

APPLIED PHYSICS REVIEWS

Nanoscale thermal transport. II. 2003–2012

David G. Cahill,^{1,a)} Paul V. Braun,¹ Gang Chen,² David R. Clarke,³ Shanhui Fan,⁴ Kenneth E. Goodson,⁵ Pawel Keblinski,⁶ William P. King,⁷ Gerald D. Mahan,⁸ Arun Majumdar,⁹ Humphrey J. Maris,¹⁰ Simon R. Phillpot,¹¹ Eric Pop,¹² and Li Shi¹³

¹Department of Materials Science and Engineering and the Frederick Seitz Materials Research Laboratory, University of Illinois, Urbana, Illinois 61801, USA

²Department of Mechanical Engineering, MIT, Cambridge, Massachusetts 02139, USA

³School of Engineering and Applied Sciences, Harvard University, Cambridge, Massachusetts 02138, USA

⁴Department of Electrical Engineering, Stanford University, Stanford, California 94305, USA

⁵Department of Mechanical Engineering, Stanford University, Stanford, California 94305, USA

⁶Department of Materials Science and Engineering, Rensselaer Polytechnic Institute, Troy, New York 12180, USA

⁷Department of Mechanical Sciences and Engineering, University of Illinois, Urbana, Illinois 61801, USA

⁸Department of Physics, Penn State University, University Park, Pennsylvania 16802, USA

⁹Department of Mechanical Engineering, University of California, Berkeley, California 94720, USA

¹⁰Department of Physics, Brown University, Providence, Rhode Island 02912, USA

¹¹Department of Materials Science and Engineering, University of Florida, Gainesville, Florida 32611, USA

¹²Department of Electrical and Computer Engineering, University of Illinois, Urbana, Illinois 61801, USA

¹³Department of Mechanical Engineering, University of Texas, Austin, Texas 78712, USA

(Received 6 April 2013; accepted 7 August 2013; published online 14 January 2014)

A diverse spectrum of technology drivers such as improved thermal barriers, higher efficiency thermoelectric energy conversion, phase-change memory, heat-assisted magnetic recording, thermal management of nanoscale electronics, and nanoparticles for thermal medical therapies are motivating studies of the applied physics of thermal transport at the nanoscale. This review emphasizes developments in experiment, theory, and computation in the past ten years and summarizes the present status of the field. Interfaces become increasingly important on small length scales. Research during the past decade has extended studies of interfaces between simple metals and inorganic crystals to interfaces with molecular materials and liquids with systematic control of interface chemistry and physics. At separations on the order of ~ 1 nm, the science of radiative transport through nanoscale gaps overlaps with thermal conduction by the coupling of electronic and vibrational excitations across weakly bonded or rough interfaces between materials. Major advances in the physics of phonons include first principles calculation of the phonon lifetimes of simple crystals and application of the predicted scattering rates in parameter-free calculations of the thermal conductivity. Progress in the control of thermal transport at the nanoscale is critical to continued advances in the density of information that can be stored in phase change memory devices and new generations of magnetic storage that will use highly localized heat sources to reduce the coercivity of magnetic media. Ultralow thermal conductivity—thermal conductivity below the conventionally predicted minimum thermal conductivity—has been observed in nanolaminates and disordered crystals with strong anisotropy. Advances in metrology by time-domain thermoreflectance have made measurements of the thermal conductivity of a thin layer with micron-scale spatial resolution relatively routine. Scanning thermal microscopy and thermal analysis using proximal probes has achieved spatial resolution of 10 nm, temperature precision of 50 mK, sensitivity to heat flows of 10 pW, and the capability for thermal analysis of sub-femtogram samples. © 2014 Author(s). All article content, except where otherwise noted, is licensed under a Creative Commons Attribution 3.0 Unported License. [<http://dx.doi.org/10.1063/1.4832615>]

TABLE OF CONTENTS

I. INTRODUCTION	2
II. THERMAL CONDUCTANCE OF INTERFACES.....	3

A. Vibrational thermal energy transport at interfaces.....	3
1. Overview.....	3
2. Interfacial thermal transport via molecular dynamics techniques.....	3
3. Role of interfacial bonding and structure.....	4
4. Interfaces in composite materials and nanoparticle suspensions.....	5

^{a)}Electronic mail: d-cahill@illinois.edu

5. Heated nanoparticles	6
6. Experiment	6
B. Near-field radiation transport across a gap ..	8
1. Overview	8
2. Experiment	8
3. Theoretical formalism	10
4. Exact treatment of near-field thermal transfer in non-planar geometries	11
5. Active modulation and control of near field thermal transfer	12
6. Image charge theory	12
III. THERMAL TRANSPORT IN NANOSCALE STRUCTURES AND DEVICES	12
A. Physics of phonon transport	12
B. Nanowires and graphene: Experiment and theory	17
1. Particle transport description of phonon-boundary scattering	17
2. Wave interference models of phonon transport in silicon nanostructures	19
3. Phonon scattering in graphene and nanotubes	20
C. Applications to information technology and high-power density electronics	21
1. Data storage	21
2. Nanoscale CMOS and interconnects	23
3. Composite substrates for power electronics	26
IV. THERMAL CONDUCTIVITY OF NANOSTRUCTURED MATERIALS	27
A. Doping and point defects	28
B. Grain boundaries	28
C. Anisotropic crystals and natural superlattices	30
V. METROLOGY AND PROCESSING TOOLS	31
A. Time-domain thermorefectance	31
1. Advances	31
2. Limitations	31
3. Implementation	32
4. Signal analysis and sensitivities	32
5. Forefront issues	34
B. Scanning thermal microscopy	34
C. Nanometer-scale thermal analysis and thermal nano-manufacturing	35
VI. OUTLOOK	36

I. INTRODUCTION

The relentless decrease in the size of devices and structures, the increase in their operating speeds and frequencies, and the ever more aggressive thermal conditions imposed upon them requires sophisticated understanding and control of thermal transport at the nanoscale. While thermal performance itself is a key metric for applications such as thermal barrier coatings, nuclear fuels, and materials for thermoelectric energy conversion, there are many other technologies for which the ability to manipulate heat is vital

even though the primary goal of the structure is non-thermal. Some examples of technology drivers for the science of nanoscale thermal transport are phase change memory devices, heat assisted magnetic recording, thermal management of wide variety of high power and nanoscale electronic and optoelectronic devices, and the proposed use of intensely heated nanoparticles in medical therapies.

The objective of this review is to provide perspectives on new developments in the applied physics of nanoscale thermal transport that have resulted from advances in experiment, theory, and simulation over the past decade; to identify gaps in current state-of-knowledge; and discuss directions for future research. This review is an update to an earlier review,¹ co-authored by some of the same authors, which surveyed progress in the field through 2002. The topics we have selected emphasize the importance of interfacial phenomena in nanoscale thermal transport and avoid topics that have been extensively reviewed by others in recent years, e.g., nanostructured thermoelectric materials^{2–7} and the transport properties of isolated graphene and carbon nanotubes.^{8–10} We limit our discussion of thermoelectric materials to the lattice thermal conductivity of Si nanowires. Our discussion of carbon allotropes focuses on the effects of interfaces on thermal transport. We also omit discussion of non-linear effects and transport in low-dimensional systems as these topics were recently reviewed by Li and co-workers.¹¹ Experimental methods for measuring the transport properties of nanowires were recently reviewed by Weathers and Shi.¹² Thorough development of the fundamentals of nanoscale thermal transport are available in recent textbooks.^{13–16}

For the purposes of this review, the term nanoscale is interpreted liberally, embracing characteristic dimensions ranging from the atomic up to sub-micron. Another way of envisaging the definition of nanoscale is the regime in which sub-continuum effects are important; that is, the system cannot be adequately described by bulk transport properties and continuum heat transport equations that do not explicitly consider interfaces. For example, while it has long been recognized that grain boundaries and interphase boundaries provide significant thermal resistance, there continues to be significant efforts to understand the transport of thermal energy across interfaces. One of the major advances in the past decade has been the extension of experiment and theory to interfaces that involve molecular materials and liquids. Systematic control of the strength of interface bonding has revealed the importance of interface chemistry and physics on heat conduction.

We include discussion of radiative transport by thermally excited electromagnetic modes in this review because we believe that there is an increasing intersection between the science of near-field radiative transport through nanoscale gaps and the thermal conductance of interfaces G between materials when G is small because of weak bonding, interface roughness, or a large mismatch in vibrational density of states. The coupling of electronic and vibrational modes at an interface is usually thought to be limited to atomic scales but recent theory suggests that longer-range coupling across the interface can be significant.

Heterostructures provide both an approach to reducing thermal conductivity in the cross-plane direction and a laboratory for studies of highly anisotropic thermal transport. Anisotropy in engineered superlattices has now been extended to natural superlattice materials, such as those in layered perovskite structures. Anisotropy is most extreme in two-dimensional or near two-dimensional materials, of which carbon-based materials (nanotubes, and most recently graphene) are the most prominent; however, considerable work is now being performed on layered metal chalcogenide systems, such as MoS₂ and WSe₂.

Historically, theory and simulation have been able to provide predictions with spatial and time resolutions for which corresponding experimental techniques did not exist or were not fully developed. In the last decade, there have been significant improvements in the spatial and temporal resolution of key techniques, and a much improved ability to determine temperatures and heat flow. This is now allowing theory and experiment to be confronted with experimental results at an unprecedented level. Advances in density functional theory and high performance computing have reached the point where parameter-free predictions of the thermal conductivities of simple crystals are now possible.

We begin our discussion with a review of theory and experiment on the transport of heat at an interface between two materials. In Sec. III, we focus on thermal transport in nanostructures and nanoscale electronic devices. The related topic of low thermal conductivity in nanostructured materials, including anisotropic crystals and natural superlattices, is addressed in Sec. IV. We explore issues associated with thermal measurement science and thermal processing at the nanoscale in Sec. V.

II. THERMAL CONDUCTANCE OF INTERFACES

A. Vibrational thermal energy transport at interfaces

1. Overview

A thermal boundary resistance is found for the heat flow between two materials in contact along a planar interface. The heat flow is perpendicular to the planar interface

$$J_Q = G\Delta T, \quad (1)$$

where ΔT is the small temperature difference between the two sides of the interface, and G is the boundary conductance; G has the units of power per area per degree.

A common interface in thermal transport is between two insulators. Then, the heat is carried only by phonons. The thermal boundary resistance G^{-1} from phonons was first calculated by Khalatnikov.^{17,18} He considered only longitudinal phonons. His calculation is called the *acoustic mismatch theory* (AMT). It is found to be a good approximation at low temperatures, where $G \propto T^3$. At room temperatures, it is a poor approximation. Young and Maris¹⁹ provided the first theory of G , which included all phonons on both sides of the junction. They derived only the special case that both sides of the interface had face-centered-cubic structures, with the same lattice constant. Pettersson and Mahan²⁰ extended the

Young-Maris theory to include any kind of lattices on the two sides of the interface, but assumed they were in intimate contact. These theories are based on phonon scattering from the interface atoms. Persson and Ueba²¹ recently derived the phonon-phonon case using surface Green's functions. That allowed them to consider rough surfaces.

Superlattices are periodic arrays of alternating materials. One layer in the array will have n -atomic layers of a material, while the alternating layers have m atomic layers. Thermal conductivity can be measured parallel to the layers, or perpendicular to them. For a crystalline superlattice, the thermal conductivity parallel to the layers is typically comparable to the average of the two bulk materials, as long as the interfaces between the layers do not have a large density of defects. The interesting measurement is for heat flow perpendicular to the layers, which is called the "cross-plane direction." Hyldgaard and Mahan²² originally predicted that the thermal conductivity of a superlattice in the cross-plane direction could be ten times smaller than the values of the bulk materials that makeup the superlattice. This reduction is generally observed. The thermal conductivity is sometimes lower in value than a random alloy of the same material. The large reduction in heat flow is due to the numerous interfaces of the superlattice. The thermal boundary resistance is a major component of the thermal resistance. Later Simkin and Mahan²³ suggested that a superlattice would have a minimum thermal conductivity as a function of the number of atomic layers within one superlattice layer. Several experiments have tested these ideas.^{24–28} (A summary of experimental studies of superlattices prior to 2002 is given in Ref. 1.)

Until recently all theory and experiments on superlattices were for insulating layers. In this case only phonons carry heat, and the boundary resistance is due to phonon scattering. Recently, there has been experimental work on metal-metal interfaces²⁹ and multilayers,³⁰ where the electrons are the important carriers of heat. Some work has been done also on metal-insulator superlattices.^{27,31}

2. Interfacial thermal transport via molecular dynamics techniques

Modeling and simulation of interfacial heat transport has been the focus of numerous computational efforts in the past decade. The most prominent approach is to use molecular dynamics (MD) simulations where atoms and molecules follow classical dynamics based on the numerical solution of the Newton's second law of motion. The forces between atoms and molecules are derived from interatomic potentials.³² In the most popular scheme, equal power, P , heat source, and sink are introduced on both sides of the interface, and when the steady state is established, the interfacial thermal conductance, G , is evaluated from the temperature drop at the interface, ΔT , via $G = P/(A\Delta T)$, where A is the interfacial area. While the heat source-sink method allows determination of the value of the conductance, detailed understanding of the thermal coupling across the interface can be obtained from the equilibrium MD simulations.^{33,34} Specifically, a Green-Kubo relationship is employed, which

relates the fluctuating cross interfacial heat power, $P(t)$, to the interfacial conductance via³⁵

$$G = \frac{1}{Ak_B T^2} \int_0^\infty \langle P(t)P(0) \rangle dt, \quad (2)$$

where $\langle P(t)P(0) \rangle$ is the interfacial heat power autocorrelation function. The interfacial heat power can be easily obtained by monitoring the total energy of the material on the one side of the interface, and then taking its time derivative. Also, the upper limit of the integral given by Eq. (2) is in practice finite as the autocorrelation function decays to zero. In addition to the value of G , the full functional form of $\langle P(t)P(0) \rangle$ provides dynamical information about coupling between molecular vibrations in the two phases forming the interface.³⁶

Equilibrium and non-equilibrium MD thermal conductivity techniques characterize either macroscopic or interfacial behavior “integrated” over all vibrations. To explicitly provide information on underlying mechanism of heat flow in terms of individual vibrations, recently developed phonon dynamics MD based techniques have proved to be useful. The basic idea of phonon dynamics simulations³⁷ is to generate wave packets of lattice vibrations of a narrow range of frequencies and well-defined polarization, using the normal modes of the perfect crystal structure, and then to allow them to propagate through the system. Following the scattering process, one can decompose the transmitted and reflected energy into phonons and thus can identify the nature of the scattering processes and to calculate the transmission coefficient for each phonon.³⁸ For example, scattering of high frequency phonons at interfaces involves a mixture of “acoustic scattering” where the reflected phonon has the same character as the incident phonon and “diffusive scattering” where the correlation with the incident phonon wave packet is lost.³⁸

The phonon dynamics technique relates to Green’s functions methods.³⁹ Both methods enable calculations of the phonon transmission coefficients. However, due to computational limitations and difficulties with the correct description of anharmonic effects, Green’s functions calculations were mostly performed on simplified interfacial models, such as junctions,⁴⁰ or nanowires.⁴¹ The transmission coefficients for longitudinal acoustic phonons determined by phonon dynamics simulations are similar to those obtained from GF calculations for a simple one dimensional chain model of masses connected by springs.^{37,42} Green’s function calculations on fully 3-dimensional structures, on the other hand, lump together all phonons at a given frequency, whereas phonon dynamics simulations can discriminate among different directions and polarizations.

3. Role of interfacial bonding and structure

The acoustic mismatch model (AMM) and diffusive mismatch model (DMM) have traditionally provided a basis for understanding interfacial thermal resistance. These models provide prescriptions for calculating the transmission coefficient of a phonon with which the interfacial conductance can be obtained by appropriate summations over all

phonons.^{19,43} While both AMM and DMM provide values for transmission coefficients, they calculate them purely based on the bulk properties of the two solids forming the interface; thus they do not account explicitly for the strength of the interfacial bonding. Consequently, MD simulations were used to model the effect of interfacial bonding on the interfacial conductance. In particular, a study of a self-assembled monolayer (SAM) water interface (Fig. 1) exposed the relationship between interfacial bonding and interfacial conductance, suggesting a more or less linear relationship between the two and saturation on the conductance at high interfacial bonding.⁴⁴ Interestingly, in this case, the MD results are in quantitative agreement with the experiment.⁴⁵ A similar relationship between the interfacial conductance and the interfacial bonding strength was observed in a number of MD simulation on interfacial systems, including hard solid-polymer,⁴² solid-solid,⁴⁶ solid-SAM-solid,^{47,48} carbon nanotube on silica substrate,⁴⁹ and for gas-solid interfaces.⁵⁰

Within the theoretical treatment, the effect of interfacial bonding on the phonon transmission coefficients and consequently on the interfacial thermal conductance can be demonstrated via an analysis of a simple model of semi-infinite one-dimensional chains of masses connected by springs.⁵¹ A calculation based on this model shows that the interfacial phonon transmission coefficient for a typical phonon representing those carrying the majority of the heat strongly increases with increasing interfacial stiffness.⁴⁶ Furthermore, the transmission coefficient saturates at strong bonding at the value coinciding which by the prediction of the AMM. A similar conclusion was reached in the case of a fully 3 dimensional model, which developed an analytical extension of the AMM that includes the effects of weak interface bond spring constants.⁵² Analysis of the one dimensional chain junctions and of the extended AMM model shows that in the limit of very weak interfacial bonding the conductance is

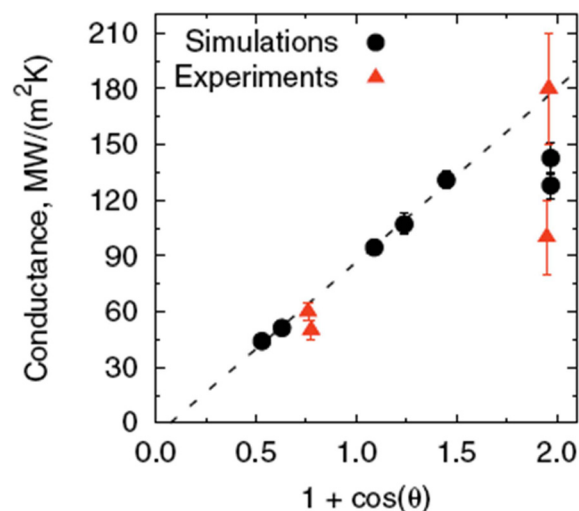


FIG. 1. The interfacial thermal conductance of a range of SAM-water surfaces as a function of $[1 + \cos(\theta)]$, which measures interfacial adhesion strength, where θ is the wetting angle. The experimental data are for interfaces between water and surfactant coated surfaces, see Ref. 45. Reproduced with permission from Phys. Rev. Lett. **102**, 156101 (2009). Copyright 2009 American Physical Society.

proportional to the square of the bonding strength. However, this scaling has little relevance to real interfaces as it holds only for unrealistically weak bonding. At larger interfacial bonding strength corresponding to van der Waals bonding, the relationship between bonding and the conductance is more-or-less linear, and at even larger bonding strength the conductance saturates. This is also consistent with observations made for 3 dimensional lattice models.¹⁹ All of the above findings suggest that AMM can be, at least empirically, extended to account explicitly for the strength of the interfacial interactions.

The above considerations indicate that when the diffuse phonon scattering is not important, the AMM model prediction is an upper limit of the possible conductance associated with strong interfacial bonding. When the diffuse scattering is dominant, the DMM predicts that the interfacial transport is maximized when the vibrational spectra on both sites of the interface are closely matching. This motivated MD simulations to explore the potential of enhancing and tuning interfacial thermal transport beyond the limits set by the inherent mismatch between bulk properties the two materials forming the interface.⁵³ In particular, it was demonstrated that an interface between vibrationally mismatched solids can be enhanced by the insertion of an interfacial film with mediating vibrational properties, or by intermixing of the two solid at the interfacial region.⁵⁴ Another example of such a mediating layer is an organic SAM used in the bonding of solids, or solids and liquids. In particular, MD simulations demonstrated that a confined water layer between quartz solids could lead to very good thermal transport properties.⁵⁵ Possible effects of surface nano-patterning^{56,57} on thermal transport were also explored, generally indicating benefits for thermal conduction due to increased surface area and better vibrational matching at the interface.

4. Interfaces in composite materials and nanoparticle suspensions

Interfaces generally present a resistance to heat flow in addition to the resistance of bulk materials forming the interface. This, due to high density of interfaces, can be particularly limiting to heat flow in nanocomposite materials. This effect manifests itself most spectacularly in the case of carbon nanotube (CNT) and graphene composites. Despite the fact that the reported value of thermal conductivity of CNTs and graphene is 2000 to 6000 W m⁻¹ K⁻¹ along axial and in-plane direction,^{58,59} while most of the polymers have thermal conductivity of less than 0.5 W m⁻¹ K⁻¹, an addition of nanocarbon fillers leads to rather limited thermal conductivity enhancements. MD simulations combined with transient laser heating measurement indicated that the interfacial conductance is very low, of the order of 20 MW m⁻² K⁻¹.^{60,61} The associated resistance is equivalent to the resistance of ten of nanometers the matrix materials. Analysis of the MD results shows that such high resistance is due to weak bonding between carbon nanofiller and the matrix and very strong bonding within the nanocarbon filler.

In the context of the composite thermal transport, MD simulation exposed differences in value of the interfacial

thermal conductance, depending on the mode of the heat transfer across the interface. In particular, it was found that the graphene-organic matrix cross-plane interfacial thermal resistance determined via thermal relaxation method is about 20 MW m⁻² K⁻¹ (Ref. 62) and consistent with experimental laser based pump-probe measurements on CNTs water.⁶⁰ Such consistency lies in the fact that the relaxation method mimics pump-probe measurements. In fact, in both experiment and standard MD simulations, the energy is pump predominantly to high frequency modes. As demonstrated by simulations, before dissipating to the matrix, first the energy flows from high to low frequency modes within graphene. Consequently, the interfacial thermal resistance in such thermal relaxation can be understood in terms of a sum of internal to the graphene resistance and the “external” resistance associated with the heat flow across the interface.

In the heat source-sink simulations where the heat is added and removed in the matrix away from the interface the interfacial conductance for the heat flow perpendicular to the graphene matrix interface was determined to be about 170 MW m⁻² K⁻¹,⁶² i.e., almost order of magnitude higher than that obtained from the relaxation simulations. This high conductance can be related to the “external” resistance alone, and heat can flow from the matrix via graphene to the matrix on the other side without employing high frequency modes in graphene.

Yet another value of the effective interfacial thermal conductance of about 30 MW m⁻² K⁻¹ was calculated for the case of heat entering a high-aspect ratio carbon nanofiber in order to propagate along the fiber.⁶³ In these cases, the heat enters the fiber predominantly via low frequency modes involving vibrations normal to the interface. However, rapid transport along the carbon fiber is facilitated by acoustic modes with no or little normal the interface displacement. Consequently the heat flow in this configuration involves energy redistribution over modes within the fiber and is thus affected by associated internal to the fiber thermal resistance.

All the cases described above show that interfacial thermal transport can involve a complicated process in which the phonons carrying most of the heat across the interface are not necessarily the phonons carrying the heat in the bulk or along the fiber, which in turn makes phonon-phonon coupling within each material forming the interface an important factor.

In the context of thermal transport in nanofluids (colloidal suspensions of solid nanoparticles in heat transfer fluids), it was postulated that at the interface an ordered liquid layer might have higher thermal conductivity than bulk liquid.⁶⁴ Molecular dynamics simulation on simple liquids concluded that such an interfacial layer does not contribute significantly to heat transfer.⁶⁵ For strong solid-liquid interactions, typical of those in nanofluids with metallic nanoparticles, a percolating network of amorphous-like fluid structures can emerge. This can facilitate additional thermal conduction paths.⁶⁶ However, a discernible increase in thermal conductivity is possible only for exceedingly small colloidal particles, limited to a few tens of atoms. Furthermore, interpretation of the cooling rates of Au nanoparticles suspended in water and organic solvents does not appear to require any unusual

thermophysical properties of the surrounding liquid to explain the experimental results.⁶⁷

5. Heated nanoparticles

A potential for using metallic nanoparticles and electromagnetic radiation induced nanoscopic heat sources is perhaps the ultimate frontier of the nanoscale thermal transport. In particular, when large laser heating is applied, the local temperature gradients can be as high at 100 K per nanometer, creating a unique opportunity of studying highly non-equilibrium systems. In this case, molecular dynamics simulations demonstrated the fluid next to the small nanoparticle can be heated well above its boiling point without a phase change, while on flat surface, a vapor formed readily.⁶⁸ These differences in heat transfer were explained by the curvature-induced pressure close to the nanoparticle, which inhibits boiling. When the nanoparticle temperature is much larger than the critical fluid temperature, a very large temperature gradient develops, resulting in close to ambient temperature just a radius away from the particle surface. The behavior reported allows us to interpret recent experiments where nanoparticles can be subjected to transient heating up to and above 1000 K, without observing boiling of the surrounding liquid.^{69,70} In cases of such extreme heating and associated high thermal gradients, a significant temperature drop occurs the interface. Moreover, nature of a supercritical interfacial liquid layer can contribute to the interfacial resistance and well to the stability. These issues are highly intriguing and remain largely unexplored.

6. Experiment

Interfaces between elements or simple inorganic crystals represent the easiest system to fabricate and model and have attracted considerable attention. Care must be taken however to ensure that the interface is atomically abrupt and clean; otherwise, interpretation of the data and rigorous modeling is difficult. Reactive sputter deposition of TiN on MgO substrates⁷¹ at high temperatures produced the highest thermal conductance observed to-date for individual interfaces at room temperature, $G \approx 700 \text{ MW m}^{-2} \text{ K}^{-1}$. Nearly ideal interfaces for fundamental studies of thermal transport were prepared by Horn-von Hoegen and colleagues using deposition of Bi on atomically clean Si under UHV conditions.^{72,73} The thermal conductance was measured *in-situ* by heating of the Bi layer with a laser pulse and measurement of the Bi temperature by ultrafast electron diffraction (UED). G for the Bi-Si interface was determined to be as low as low as $13 \text{ MW m}^{-2} \text{ K}^{-1}$, comparable to G for Bi deposited on diamond.⁷⁴ Interestingly, G for Bi-Si was nearly the same for (111) and (001) oriented Si substrates, indicating that details of the crystallographic structure of the interface have a minimal effect on G . G was also independent of thickness even for Bi layers only 2.5 nm thick.⁷³ Going forward, UED can probe interfaces and structures, e.g., non-planar Ge-Si interfaces⁷⁵ that are not accessible in more conventional approaches based on pump-probe optical methods.

Although single interfaces are often easier to fabricate, a key advantage of a multilayer systems is that although the

effect of each interface may be small, in a system which consists of a “solid of interfaces,” the effect of interface conductance on thermal transport can be dramatic. In one multilayer system, atomic layer deposition and magnetron sputter deposition were used to synthesize thin-film multilayers of W/Al₂O₃. The high interface density produced a strong impediment to heat transfer, giving rise to a thermal conductivity as low as $\kappa \approx 0.6 \text{ W m}^{-1} \text{ K}^{-1}$ as the interface density approached 0.5 nm^{-1} .⁷⁶ Even lower conductivity, $\kappa < 0.5 \text{ W m}^{-1} \text{ K}^{-1}$, was observed in Ta/TaO_x multilayers.⁷⁷ At an interface density of 0.14 nm^{-1} , Mo/Si multilayers⁷⁸ of the type used for mirrors in the extreme-ultraviolet have a thermal conductivity of $\approx 1 \text{ W m}^{-1} \text{ K}^{-1}$. A multilayer approach has also been applied to the study of organic/metal interfaces, namely, copper phthalocyanine interfaces with Ag and Al.^{79,80}

Common to all four studies^{76–79} is the use of a metallic layer as a component in the multilayer. The thermal conductivity of ultrathin metallic layers in a metal/dielectric multilayer depends on the strength of electron-phonon coupling.^{77,78,81,82} The electronic contribution to thermal conductivity κ_{el} is suppressed when the metal layer thickness is smaller than $\sqrt{\kappa_l/g}$, where g is the electron-phonon coupling parameter⁸³ and κ_l is the lattice contribution to the thermal conductivity. Similarly, the interface conductance of a metal-dielectric interface has an additional series conductance⁸¹ of $\sqrt{\kappa_l g}$. Au has weakest electron-phonon coupling and lowest lattice thermal conductivity among the metal elements: at room temperature, $g \approx 3 \times 10^{16} \text{ W m}^{-3} \text{ K}^{-1}$ and $\kappa_l \approx 3 \text{ W m}^{-1} \text{ K}^{-1}$.⁸⁴ Therefore, for an interface with Au, the additional series conductance is $\sqrt{\kappa_l g} \approx 300 \text{ MW m}^{-2} \text{ K}^{-1}$. For all other metals, $\sqrt{\kappa_l g}$ is significantly larger.

The lowest thermal conductivity observed in a fully dense solid at room temperature is $\kappa \approx 0.05 \text{ W m}^{-1} \text{ K}^{-1}$ for the cross-plane thermal conductivity of thin films of WSe₂ grown from vacuum deposition of alternating W and Se layers.⁸⁵ This conductivity is 30 times smaller than the *c*-axis conductivity of WSe₂ single crystal but more significantly is a factor of ≈ 5 times lower than minimum thermal conductivity⁸⁶ calculated from the atomic density and the speeds of sound. The origin of this “ultralow” conductivity not yet fully understood—for example, it is not clear if the transport can be described as a the series sum of the thermal conductance of closely spaced interfaces acting independently—but the strong anisotropy of the elastic constants that reduces the density of states of phonon modes that propagate in the cross-plane direction^{87,88} is thought to be an important factor.

The thermal conductivity of amorphous soft materials (e.g., polymers) is inherently low ($\kappa \approx 0.2 \text{ W m}^{-1} \text{ K}^{-1}$), and thus the effect of interfaces is less dramatic. There are specific cases however where interface conductance is still important, e.g., surfactant stabilized nanoparticles. Most inorganic nanoparticles are made dispersible in a solvent by coating their surface with a thin layer of organic ligands, and thus are best described as a system containing two interfaces. The first is the interface between the inorganic nanoparticle and the surrounding ligand, and the second is the interface between the ligand and the solvent phase. A motivation for

understanding heat flow surrounding nanoparticles comes from the broad interest in nanoparticle photothermal therapies.⁸⁹ In nanoparticle photothermal therapies, nanoparticles, which are designed to target various tumors, are injected into the body. Then, near IR radiation is used to heat the nanoparticles and locally kill tissue. Simple calculations show, however, that for any reasonable combination of optical power, nanoparticle loading, and background optical absorption, it is very challenging to create a nanometer-scale local temperature excursion using a cw laser source.^{90,91} Excitation with pulsed lasers with low duty-cycle can localize temperature excursion in the immediate vicinity of a nanoparticle.⁹⁰ At short time-scales, interface thermal conductance plays an important role in the heat transfer from nanoparticle to surroundings.⁶⁷

Model systems consisting of nanoparticles dispersed in water and other solvents have been studied to investigate the effect of the organic stabilizing group on the thermal conductance G of the particle/fluid interface.^{67,92,93} In a series of experiments on heat conduction from ligand-stabilized nanoparticles to solvents, G ranged from $5 \text{ MW m}^{-2} \text{ K}^{-1}$ for alkanethiol stabilized AuPd alloy nanoparticles in toluene to $100\text{--}300 \text{ MW m}^{-2} \text{ K}^{-1}$ for various metal nanoparticles in water.^{67,92} The similar values of G for all nanoparticles in water indicates that the thermal coupling between nanoparticles and water is strong regardless of the self-assembled stabilizing ligand. This was true for tiopronin (anionic), ethylene glycol (nonionic), and cetyltrimethylammonium bromide (cationic bilayer) ligands on particles ranging from 3 to 24 nm.

In principle, the measured value of G of a nanoparticle-surfactant-solvent system could include at least four contributions: (i) the flow of heat from the nanoparticle into the stabilizing group; (ii) the flow of vibrational energy along the length of the stabilizing group or across the molecular layer; (iii) the transport of vibrational energy from the stabilizing group to the surrounding fluid; and (iv) the possibility of the direct transfer of heat from the nanoparticle to fluid molecules that penetrate the self-assembled molecular layer of the organic stabilizing group. The large difference in the values of G for toluene and water suspensions suggests that either the third or fourth mechanism listed above is playing a critical role since we do not have any reason to believe that the first two mechanisms would have a strong dependence on the solvent. The large values for the thermal conductance G of nanoparticles in aqueous solution may either indicate that the thermal coupling between hydrophilic molecular layers and water is strong, while the thermal coupling between alkanethiol monolayers and toluene is much weaker, resulting in a smaller G , or that water penetrates deeply within the molecular layer and interacts directly with the nanoparticle. This second explanation appears unlikely in light of the cetyltrimethylammonium bromide results since this molecule forms a bilayer on the surface of the nanoparticle with a hydrophobic core, and the water content of the interior of the bilayer is small.

With nanoparticle systems, it is not possible to explore heat transfer between hydrophobic surfaces and water, or highly polar surfaces and a non-polar solvent. These systems

are not stable and aggregate rapidly. However, by moving to a planar system, such experiments are possible. Using time-domain thermorefectance (TDTR), Ge and co-workers⁴⁵ measured the transport of thermally excited vibrational energy across planar interfaces between water and solids that have been chemically functionalized with both hydrophobic and hydrophilic self-assembled monolayers. In these systems, it is useful to think of the interfacial conductance in the form of a Kapitza length—i.e., the thermal conductivity of water divided by G . The Kapitza length at hydrophobic interfaces (10–12 nm) is a factor of ~ 3 larger than the Kapitza length at hydrophilic interfaces (36 nm),⁴⁵ indicating a much stronger thermal coupling between hydrophilic surfaces and water than hydrophobic surfaces and water.

Hard-soft-hard interfaces (e.g., two inorganic layers sandwiching an organic layer) are interesting both because they can be experimentally realized in systems with high degree of structural and chemical control, and for the technological reason that they may provide a route to dramatically alter thermal conductances in composite materials with attractive physical properties. A key question in hard-soft-hard systems is the relationship between interfacial bonding character and thermal conductance. MD simulations^{42,44,48} as well as analytical models⁵² predict interfacial bond stiffness can cause G to vary by an order of magnitude. This has been probed by sandwiching a SAM between two surfaces. In one example, AuSAMGaAs junctions were made using alkanedithiol SAMs and assembled by transfer printing.⁹⁴ Thermal conductance was measured using the 3ω technique, and no thermal conductance dependence on alkane chain length was observed over a relatively narrow chain length (8–10 carbons). The thermal conductances ranged from 25 to $28 \text{ MW m}^{-2} \text{ K}^{-1}$.⁹⁴ Because the character of the bonding does not change, this experiment provides only one data point for the chemical effects on G .

To explicitly study the effect of interfacial bonding character, a gold film was transfer-printed to a self-assembled monolayer with systematically varied termination chemistries. Using a combination of ultrafast pump-probe techniques (time-domain thermorefectance and picosecond acoustics) and laser spallation experiments, changes in bonding strength and heat flow at the gold-SAM interface were independently measured and correlated.⁹⁵ It was shown that varying the density of covalent bonds within this single bonding layer modulates both interfacial stiffness and interfacial thermal conductance. Heat transport at Au/SAM/Qz interfaces using SAMs of the same length and different ω -end-groups: (1) Methyl ($\text{CH}_3\text{-C}_{11}\text{-Si} \equiv$) and (2) thiol ($\text{SH-C}_{11}\text{-Si} \equiv$) were studied. The thiol functionality is expected to form a strong covalent-like bond to the gold film^{96,97} while only a weak van der Waals attraction should attach the gold film to the methyl-terminated surface. $G = 68 \text{ MW m}^{-2} \text{ K}^{-1}$ for the Au/SH- $\text{C}_{11}\text{-Si} \equiv$ Qz interface and $G = 36 \text{ MW m}^{-2} \text{ K}^{-1}$ for the Au/ $\text{CH}_3\text{-C}_{11}\text{-Si} \equiv$ Qz interface, confirming that interfacial bonding directly impacts thermal conductance. Particularly striking is the similarity between G measured for Au/SH- $\text{C}_{11}\text{-Si} \equiv$ Qz ($G_{\text{avg}} = 65 \text{ MW m}^{-2} \text{ K}^{-1}$) and the recent calculations by Keblinski for a Au/SAM/Si interface with strong bonding at both SAM end-groups ($G = 60 \text{ MW m}^{-2} \text{ K}^{-1}$).⁴²

Hsieh and co-workers continuously varied the strength of interfacial bonding using high hydrostatic pressure in a SiC anvil cell.⁹⁸ An Al-graphene-SiC interface was formed on the face of the SiC anvil using transfer printing of a graphene followed by sputter deposition of Al. As the pressure increases from ambient to 10 GPa, the value of G increases by nearly an order of magnitude from $G \approx 30$ to $G \approx 200 \text{ m}^{-2} \text{ K}^{-1}$. At high pressures, G for Al-graphene-SiC is the same as G for Al deposited directly on SiC cleaned at high temperatures.

Hard-soft-hard interfaces also offer the potential for switchable nanoscale thermal conduction. Recently, it was shown that the thermal conductivity of a boron nanoribbon bundle can be modulated between high and low thermal conductivity states by wetting the interface between the nanoribbons comprising the bundle.⁹⁹ When the ribbons are welded together cleanly by the solvent (ethanol/water mixture), phonons transmit across the interface, leading an increase in the thermal conductivity. Once this solvent is removed and replaced with isopropyl alcohol (IPA), the thermal conductivity decreases. This process is reversible. IPA leaves residues after evaporation, and it is thought these residues separate the nanoribbons, making them effectively non-interacting.

B. Near-field radiation transport across a gap

1. Overview

Photons can transfer energy between two distant bodies that are at different temperatures. The formula for the thermal conductance contains the factor σT^3 , where σ is the Stefan-Boltzmann constant. This energy transfer is called a *far-field* transfer. There is also a *near-field* energy transfer that is electro-magnetic in origin. It operates on a distance scale less than ten micron, i.e., less than the wavelength of thermally excited infrared radiation. This near-field energy transfer has been much studied.^{100–116} A review of experiment and theory of near-field thermal transport is given in Secs. II B 2 and II B 3.

Polder and van Hove¹⁰¹ wrote out the basic equations in terms of local dielectric functions. The dielectric function for a homogeneous material is a function of wave vector and frequency $\epsilon(q, \omega)$. The dielectric formulations typically neglect the wave vector dependence and just use $\epsilon(0, \omega)$ to match the electromagnetic fields at the surface. For insulating solids, that is a good approximation since the wave vector dependence of the dielectric response is small, but for metallic samples, we expect deviations when the gap becomes comparable to atomic length scales.

2. Experiment

In the far field, radiation heat transfer between macroscopic bodies is limited to below that given by the Planck blackbody radiation law for the spectral-based flux or the Stefan-Boltzmann law for the total flux, both of which will be referred to as the blackbody law limit. When two objects are brought close to each other, radiation can *exceed* the blackbody limit because of the tunneling of evanescent

waves. There are two types of evanescent waves.^{14,117,118} One decays exponentially from one-side of the interface but is a propagating wave on the other side of the interface, and the other decays exponentially on both sides of the interface. The former happens when electromagnetic waves inside a material with a high and positive dielectric constant propagate across the interface with a material of a lower but positive dielectric constant at an angle of incidence larger than the critical angle as determined by the Snell law, creating an exponentially decaying electromagnetic field, i.e., an evanescent wave, on the other side. The latter is called a surface wave, which happens when the dielectric constants on both sides of an interface are opposite signs, and the absolute value of the negative dielectric constant is larger than that of the positive dielectric constant. A negative dielectric constant occurs when the frequency of photons falls in between the transverse and longitudinal optical phonon frequencies for dielectrics or below the plasma frequency for metals, when there are resonances between material waves (e.g., electrons of optical phonons) and the electromagnetic waves.

Examples of surface waves are surface plasmon-polaritons or surface phonon-polaritons, which are resonance modes of the electrons and electromagnetic waves, and the optical phonons and electromagnetic waves, respectively. Evanescent waves normally do not carry a net energy flux and decay rapidly away from the interface over a distance of the order of the wavelength, and hence do not contribute to radiation heat transfer between objects separated at macroscopic distances. However, when the two surfaces are brought close by, evanescent waves can reach the other medium and enhance heat transfer.

Measurement of near-field radiation is challenging because (i) thermal radiation heat transfer rate is small, and (ii) near-field effects start to be important only when the separation between the two surfaces are comparable to the dominant wavelength. Using Wiens displacement law $\lambda T = 2898 \text{ } \mu\text{m K}$ as a guideline, at room temperatures, blackbody radiation peaks at the wavelength of $\approx \lambda = 10 \text{ } \mu\text{m}$. Measuring small heat transfer at small separations of two surfaces is difficult. Tien and co-workers^{119,120} reported nearfield radiative heat transfer at cryogenic temperatures between two copper disks with a diameter of 8.5 cm, whose cold and the hot sides were at 4.1 K and 15.1 K, respectively. Due to the low temperatures, the near-field effect can be observed at large gaps. They observed increasing radiative heat transfer as the size of the gap decreases from 1 mm to 50 μm , consistent with theoretical expectations. Their experiments were stimulated by cryogenic multilayer insulations. Due to the low emittance of metallic surfaces, the maximum heat transfer observed is $\approx 1/29$ of the Planck theory prediction. Hargreaves¹²¹ carried out near-field radiation heat transfer measurements between two chromium surfaces with a diameter of 4.0 cm,^{118,121} to gaps down to 1 μm at room temperature, demonstrating an enhancement from ≈ 1.75 in the farfield to $\approx 2.95 \text{ W m}^{-2} \text{ K}^{-1}$ at a 1 μm gap, which is $\sim 50\%$ of blackbody radiation. More recently, Kralik *et al.*¹²² carried out near-field measurements between tungsten surfaces, each 35 mm in diameter, with a separation distance between $\sim 1 \text{ } \mu\text{m}$ to 1 mm. The hot-side surface was maintained between 10 and 100 K and cold side maintained at 5 K. They

reported a three orders of magnitude increase in the emittance between the two surfaces.

Metallic surfaces were used in these experiments because capacitive sensing can be used to control the separation and parallelism between the surfaces. Reducing the separation between surfaces less than microns, however, is challenging because of the difficulties in maintaining the surfaces parallel and the presence of dust particles and other imperfections. Xu *et al.*¹²³ took the strategy of reducing the surface area of two metallic surfaces using a scanning tunneling microscope (STM) configuration. One side of the surface is a thin-film thermocouple, and the other side of the surface is an indium tip flattened by pressing it against a glass flat. Their experiment, however, was not conclusive due to the significantly reduced heat transfer with smaller surface area and the low sensitivity of the thermocouple heat flux sensor. Kittel *et al.*¹⁰⁵ used an improved STM configuration to measure heat transfer of a gold tip with a gold substrate and with a GaN substrate for tip-substrate separation in the range of 1–100 nm. They observed saturation of heat transfer when the tip-substrate separation is less than 10 nm.

To study near-field heat transfer between non-metallic surfaces, especially between two surfaces supporting surface phonon-polaritons, Hu *et al.*¹²⁴ used two quartz surfaces with a diameter of 1.27 cm separated by polystyrene beads with a nominal diameter of 1 μm . They reported near-field heat transfer between the quartz surfaces exceeding that of the blackbody radiation. Recently, Ottens *et al.*¹¹⁴ reported measurements of near-field radiation between two large sapphire surfaces, $5 \times 5 \text{ cm}^2$ in area, for distance between 2–100 μm near room temperature. They maintained the surface separation and parallelism using small capacitive sensors deposited on the surfaces. Their experiment observed increasing heat transfer with decreasing separations between two surfaces starting from 10 μm , and heat transfer coefficient exceeding that between blackbodies.

The Chen group at MIT developed a sphere-plate experimental configuration to investigate near-field heat transfer.^{125,126} This experimental configuration overcomes the difficulties of maintaining parallelism in the two parallel plates and unknown tip geometry in the STM experiment. Their experiment took advantage of the proven extreme thermal sensitivity of bi-layer cantilevers used for atomic force microscopes,^{127–129} which compensates for a smaller area involved in radiative transfer due to the curvature of the sphere compared to the parallel plate configuration. They reported experimental results between a glass sphere and different substrates including glass, silicon, and gold, for sphere and plate separation down to 30 nm. Such experiments were extended to smaller separations and between metallic surfaces.^{130,131} Rousseau and co-workers used a slightly modified configuration and reported near-field heat transfer between a glass sphere and a glass substrates.¹³² A subtle but significant difference between the two experiments is that in the MIT experiment, the cantilever is heated while the surrounding is maintained at ambient temperature, while in the experiment of Rousseau *et al.*, the substrate is heated. In the former case, only near-field heat transfer is measured, since the far-field contribution is included in the initial bending of the

cantilever. In the latter case, the influence of the heated substrate to the sphere, and cantilever is modeled and used as a fitting parameter. Theoretical simulation by Otey and Fan¹¹⁵ shows that the far-field component creates significant ambiguity in the interpretation of the data.

Figure 2 summarizes some of the reported near-field experimental data. Experimental data in different papers were presented in different forms: power, conductance, effective emittance, and heat transfer coefficients. These data are converted into heat transfer coefficients. For sphere-plate experiments, the conversion of conductance and heat transfer coefficient between two parallel plates relies on the proximity-force approximation, also called the Derjaguin approximation,¹²⁸ which treats interactions between curved surfaces as locally parallel surfaces. Exact solutions of the problems have shown that this approximation is valid as long as the far-field contributions are properly separated.^{129,133} Figure 2 does not include the conductance data from Refs. 120 and 126, as the former has undefined geometry and the latter is complicated by the large far-field contributions.

Although the existing experiments have clearly established the phenomena of enhanced radiation heat transfer in

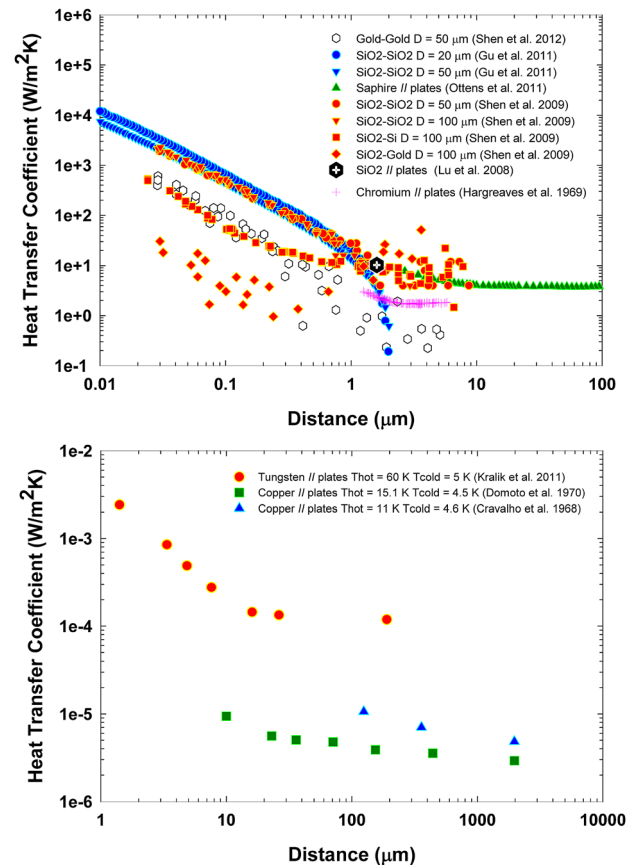


FIG. 2. Summary of experimental data on near field thermal radiation heat transfer between different materials, (a) at or near room temperature and (b) at cryogenic temperatures, expressed as heat transfer coefficients. All heat transfer coefficients in (a) between a sphere-and-plate were extracted based on measured near-field radiation heat transfer conductance^{126,130,131} using the proximity approximation and do not include the far-field contributions, while experimental data between parallel plates in both (a) (Refs. 114, 121, and 124) and (b) (Refs. 119, 120, and 122) include both far-field and near-field contributions.

the near-field, there are still many unanswered questions that require further development of experimental techniques and theoretical understanding. In the sphere-plate experiments, as the minimal separation between the sphere and the plate enters into a few to tens of nanometers regimes, surface roughness of the sphere and the plate becomes important. The impact of this roughness on experimental results and data interpretation are not clear. Experimentally, determining accurately the separation between the two surfaces, especially between nonconducting surfaces, is a nontrivial problem that has not yet been solved. In addition to thermal signal, the force interactions between the sphere and the plate can also complicate the experiments. The convolution between the force and thermal signals becomes more severe at small separations and better experiments or new designs of cantilevers are needed to separate the force and thermal signals.

At nanometer separations, the dielectric constant used to describe averaged response of a medium may lose its validity. A few theoretical studies consider the nonlocal description of dielectric constant,^{134,135} although the nonlocal theory on the dielectric function is not well-established. In the limit when two surfaces are in contact with each other, heat transfer is described by the heat conduction picture as discussed in Sec. II A. A complete description of the transition between radiation when the surfaces are nanometers apart to heat conduction when the surfaces are in contact has yet to be accomplished.

To close this section, we discuss the connection between the near-field radiation heat transfer and the Casimir force, a generalization of the van der Waals force between surfaces.¹³⁶ As Casimir first pointed out, the van der Waals force between two objects can also be interpreted as the change in the zero point energy stored in the space between the two objects. As the separation between the objects shrinks, the density of states of the electromagnetic wave changes and hence the energy stored in the space changes, leading to a net force between the surfaces. A generalized theory for the Casimir force was developed by Lifshitz and co-workers,¹³⁷ based on the fluctuational electrodynamics theory of Rytov.¹³⁸ As we discuss in more detail further, the fluctuational electrodynamics theory is also the foundation for near-field heat transfer. The Casimir force is generated by the zero-point fluctuations in the two interacting media and described by the Maxwell tensor, while the near-field radiation heat transfer is generated by the thermal fluctuation and described by the Poynting vector. A large amount of theoretical and experimental work has been done in the context of the Casimir force,^{139,140} and the two areas can benefit from each other through increased interactions.

3. Theoretical formalism

The theoretical treatment of near-field electromagnetic thermal transfer aims to calculate the heat flux between two bodies separated by a vacuum gap. (Fig. 3) These bodies are maintained at temperatures T_1 and T_2 , respectively. The term “near-field” refers to the regime where the characteristic

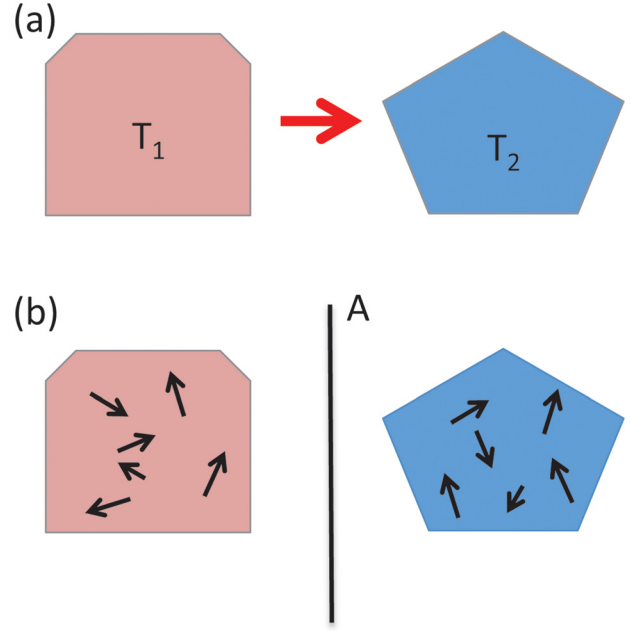


FIG. 3. (a) Near-field thermal transfer between two arbitrarily shaped bodies maintained at different temperatures. (b) Schematic of the computational set up. The heat transfer is modeled by placing fluctuating current sources inside the bodies, and by calculating the resulting electromagnetic flux at a surface in vacuum, (labelled as “A”), between the bodies.

width of the vacuum gap is below the thermal wavelength at these temperatures.

At a fundamental level, near-field thermal transfer can be treated using the formalism of fluctuation electrodynamics.¹⁴¹ In this formalism, the geometries are modeled by a dielectric constant distribution $\epsilon(r, \omega) = \epsilon'(r, \omega) + i\epsilon''(r, \omega)$, where r represents spatial location and ω is the frequency. The imaginary part of the dielectric constants, $\epsilon''(r, \omega)$, represents material loss. By the fluctuation-dissipation theorem, such material loss is directly connected to the presence of thermally induced current sources $j_\alpha(r, \omega)$ with an ensemble-averaged fluctuation of

$$\langle j_\alpha^*(r, \omega) j_\beta(r', \omega') \rangle = \frac{4}{\pi} \omega \Theta(\omega, T) \delta(r - r') \delta(\omega - \omega') \epsilon''(r, \omega) \delta_{\alpha\beta}, \quad (3)$$

where

$$\Theta(\omega, T) = \frac{\hbar \omega}{\exp(\hbar \omega / k_B T) - 1} \quad (4)$$

is the mean thermal energy for a given mode at a frequency ω , α , and β label different spatial directions. With the source defined, the spectral density of the heat flux is then calculated by integrating the Poynting vector as generated by these fluctuating currents on a plane in the vacuum region (labelled as ‘A’ in Fig. 3) between the two bodies

$$S(\omega) = \frac{1}{2} \Re \int_A dn \cdot \int d\omega' \langle E^*(r, \omega) \times H(r, \omega') \rangle. \quad (5)$$

Here, $\Re$ refers to the real part. The total heat flux is determined as

$$S = \int_0^\infty \frac{d\omega}{2\pi} S(\omega). \quad (6)$$

As outlined above, the computation of near-field electromagnetic thermal transfer is reduced, in principle, to a sequence of calculations on the electromagnetic radiations from various sources. This process of calculating thermal flux assumes that the material properties are accurately characterized by the dielectric constant of the material. Moreover, the dielectric constant is typically assumed to be local, i.e., the dielectric constant depends on the frequency but not the wave vector of the excitation. One expects that such a local dielectric function should be applicable unless one is in the deep near-field regime where the size of the vacuum gap is reduced to a few nanometer.^{105,142}

The implementation of fluctuational electrodynamic treatment of near-field electromagnetic thermal transfer involves integration over large number of sources and frequencies and thus in general is very demanding computationally. While the basic formalism of fluctuational electrodynamics was developed in the 1950s,¹⁴¹ for several decades afterwards, exact calculations of near-field electromagnetic thermal transfer were restricted to planar geometries.^{101,102}

For non-planar geometries, a number of approximate methods have been developed.^{103,132,133,143,144} The dipole approximation was developed to treat the thermal transfer between a deep sub-wavelength nanoparticle and a planar substrate.^{143,144} The proximity approximation, which was also extensively applied, approximate the two bodies in terms of a series of parallel planes.¹³² One should note however that the proximity approximation is an uncontrolled approximation in the sense that there is no systematic process through which one can improve the approximation towards an exact result. Also, the proximity approximation does not reproduce the correct far-field results when the two bodies are far apart from each other.¹³³

4. Exact treatment of near-field thermal transfer in non-planar geometries

One of the most significant developments in the past decade for near-field thermal transfer is the development of techniques that implement fluctuational thermodynamics in a numerically exact fashion in non-planar geometries.

Narayanaswamy and Chen solved the thermal near-field radiative transfer between two spheres.¹⁴⁵ The sphere-plate geometry (Fig. 4), which directly connects to some of the recent experiments, has been solved independently by Krüger *et al.*,¹⁴⁶ and by Otey and Fan.¹¹⁵ Direct comparison between numerically exact calculations and the experiment as well as several previously used approximations has been done in Ref. 115. The exact calculations did agree with the experiment of Ref. 132, if one were to modify some of the fitting parameters that were used to interpret the experiment. The exact calculations also correctly reproduced the far-field limit when the sphere-plate separation is significantly larger than the thermal wavelength (Fig. 4), as well as the results of the dipole approximation when the sphere is small. There is, however, a significant discrepancy between the exact calculations and the proximity approximation (Fig. 4).

The works of Refs. 115, 145, and 146 all involve the expansion of the fields outside the bodies in terms of either spherical or cylindrical harmonics that are solutions of the individual bodies. Therefore, these approaches are applicable only for relatively simple geometries such as spheres, cylinders, and plates. As an alternative to these approaches, finite-difference time-domain (FDTD) simulations of near-field radiative heat transfer between photonic crystal slabs has been performed in Ref. 147. In this approach, the current-current correlation of Eq. (3) is implemented directly in the simulation by introducing randomly fluctuating currents with the fluctuation of these currents simulated by a random number generator with a correlation determined from the fluctuation dissipation theory. By calculating the flux as emitted from these random sources, and by performing a time average over a long simulation time, as well as an ensemble average over different random source configurations, the thermal flux can be calculated. Boundary element approach has been implemented in Ref. 148, which may significantly reduce the computational cost compared with the FDTD approach, since in the boundary elements one only stores a current distribution on the surface of the body, thus only a two-dimensional discretization is required for a three-dimensional object. This is in contrast with the FDTD methods where storage of fields in full three dimensions are required.¹⁴⁷

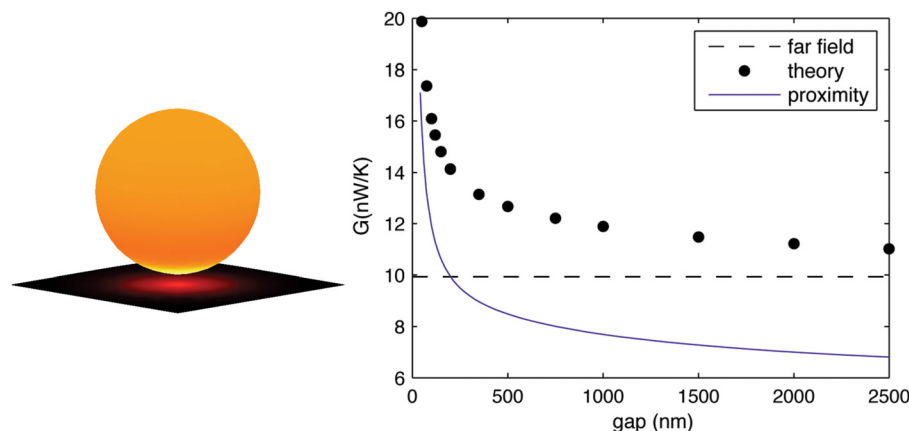


FIG. 4. Left panel: The sphere-plate geometry. Right panel: Thermal conductance between a 20 μm radius SiO_2 sphere and a SiO_2 substrate. The sphere is at a temperature of 321 K, the plate is at 300 K. The dots are exact numerical results taken from Ref. 115. The solid line is from proximity approximation. The dashed line is the far-field limit.

5. Active modulation and control of near field thermal transfer

An emerging recent direction is the theoretical exploration of active control of near-field thermal transfer. Otey *et al.* have theoretically shown that significant thermal rectification can be achieved in near-field thermal transfer between two planar bodies of SiC.¹¹⁰ In the near field of a thermal body, due to electromagnetic surface excitation, the entire thermal electromagnetic field can have a spectrum that consists of a single resonant peak.¹⁴⁹ Otey *et al.* showed that a strong rectification behavior can be achieved by exploiting the temperature dependency of the resonance frequency of such resonant peaks.¹¹⁰ Predictions about thermal rectifications using doped silicon,¹⁵⁰ as well as the possibilities of negative differential thermal conductance in near-field thermal transfer,¹⁵¹ and significant modulation of near-field thermal transfer,^{152,153} have also been made. Further developments of theory will need to take into account the variation of the phonon-polariton line width as function of temperature, which can be significant in a number of polar materials.¹⁵⁴ These works have indicated that many of the thermal circuit concepts, which have been originally explored in phonon thermal transport,¹¹ may be useful in controlling photon-based thermal transport as well.

6. Image charge theory

Stoner and Maris¹⁵⁵ and Lyeo and Cahill⁷⁴ measured the conductance G between various metals and insulators. Some of their values at room temperature are summarized in Table I.

Mahan was able to explain these results by calculating the heat flow between a metal and an insulator when their parallel surfaces are in contact.¹⁰⁷ He invoked image charge theory, which is a type of near-field effect. His theory did not account for dielectric matching but instead evaluated the effects of the electric fields on the conduction electrons in the surface layer. His theory used large wave vectors in the metals, small frequencies in both materials, and is not given by dielectric theory.

When the ions in the insulator vibrate, they create local electric fields. At the metal surface, the fields are given by classical image charge theory. These fields interact with the conduction electrons at the surface of the metal. The theory involved only parameters of the insulator such as dielectric constant, ion mass, ion Sziget charge, and lattice constants. In the metal, the only parameter is electron density, which determines the Fermi wave vector. He calculated the boundary conductance G for several alkali halide crystals, where

the metal is aluminum, since all parameters are known. He found room temperature values of G in the range of 40–100 MW m⁻² K⁻¹. These fall in the same range as the values in Table I.

Mahan's theory does not apply to the case where the insulator is diamond. Diamond is not infrared active, and its vibrations do not create oscillating electric fields. There is, as yet, no quantitative explanation of the experimental results in Table I for diamond as the insulator although three-phonon scattering at the interface^{74,156} may be particularly important at interfaces between materials with highly dissimilar phonon density of states such as Pb/diamond and Au/diamond.

III. THERMAL TRANSPORT IN NANOSCALE STRUCTURES AND DEVICES

A. Physics of phonon transport

In this section, we give a brief review of the basics of phonon propagation and scattering. An understanding of these two topics is essential for the development of a theory of the heat flow in a small structure.

To develop a quantitative understanding of the heat flow in a small structure, there are at the present time two rather different approaches. The first is to use some version of molecular dynamics. (For a review of some of the variations of this technique, see Ref. 34.) This involves, as a first step, the construction of a model for the interatomic potential; this can be based on a fit to measured quantities such as the phonon dispersion relation or it can be calculated from first principles. The interatomic potential has to include terms up to at least third order in the atomic displacements so that anharmonic phonon-phonon interactions can be correctly modeled. Various methods can be used to calculate heat flow.³⁴ The most straightforward is to connect each end of the nanostructure to heat baths with different temperatures and then calculate the steady state flow of heat from one bath to the other. As an alternative, one can calculate the fluctuating thermal currents in a structure and relate these to the thermal conductivity through some form of the fluctuation-dissipation theorem. (For early work, see Ref. 33).

As a tool for calculating heat flow, molecular dynamics has two limitations. The first problem is that for some nanostructures the number of atoms that need to be included in the simulation may be rather large. As an example, suppose that it is desired to calculate how the thermal conductivity of a nanowire varies with diameter in the range of diameters where there is a crossover from ballistic propagation to diffusive propagation. If the phonon mean free path is 50 nm, it is necessary to consider wires of diameter up to at least 100 nm and lengths of these wires of perhaps 200 nm. In such a volume, they are on the order of 100 million atoms, making a simulation challenging. The second problem is that the calculation is classical. As a result, in the simulation, the average energy of each phonon mode is equal to $k_B T$, whereas the correct energy distribution is given by the Planck distribution. For modes with frequency such that $\hbar\omega < k_B T$ the error is small, but the error becomes large for any solid in which the phonon spectrum extends up to frequencies such

TABLE I. Thermal boundary conductances (G) between some metals (in the top row) and insulators (in the left column) at 300 K. Units of G are MW/(m² K). The first three columns are data from Ref. 155. The last column is data from Ref. 74.

	Al	Au	Pb	Pb (Ref. 74)
BaF ₂	100	40	62	—
Al ₂ O ₃	105	45	55	28
Diamond	46	40	31	19

that $\hbar\omega > k_B T$. It is hard to judge the effect of this error. The fact that the simulation overestimates the specific heat will lead to an overestimate of the conductivity. But it is also true that the error in the average energy of the high frequency modes will result in an overestimate of the rate of phonon-phonon scattering; this leads to an underestimate of the heat flow. For materials such as silicon with a Debye temperature of 600 K, the error may not be too large for simulations at room temperature, but for materials such as diamond ($\Theta_D \sim 2000$ K), carbon nanotubes and graphene, most of the phonon modes have such a high frequency that they are poorly described by classical mechanics. For a recent discussion of attempts to correct molecular dynamics for quantum effects, see Ref. 157.

Molecular dynamics has been used to successfully model the heat flow across interfaces (see Sec. II A) and heat flow in superlattices.^{158–161} For these applications, the method has the great advantage that it is easy to include in the model different aspects of the microscopic structure of an interface, such as the roughness^{48,57,162–164} and mixing.¹⁶⁵

The second approach is based on the phonon Boltzmann equation¹⁶⁶

$$\frac{\partial n(\vec{k}j)}{\partial t} = \left. \frac{\partial n(\vec{k}j)}{\partial t} \right|_{coll} - \vec{v}(\vec{k}j) \cdot \nabla n(\vec{k}j), \quad (7)$$

where $\partial n(\vec{k}j)/\partial t|_{coll}$ is the rate at which the phonon distribution $n(\vec{k}j)$ changes due to collisions, $\vec{k}$ is the wave vector, j is an index indicating the polarization, and $\vec{v}(\vec{k}j)$ is the group velocity. Since we are primarily interested in heat flow as a result of small and constant temperature gradients (gradients such that the temperature changes by a small amount over a distance of one phonon mean free path, and over a distance equal to the transverse dimensions of the sample), we can use the time-independent and linearized version of this equation

$$\frac{\hbar\omega(\vec{k}j)}{k_B T^2} \bar{n}(\vec{k}j) (\bar{n}(\vec{k}j) + 1) \vec{v}(\vec{k}j) \cdot \nabla T = \left. \frac{\partial n(\vec{k}j)}{\partial t} \right|_{coll}, \quad (8)$$

where $\bar{n}(\vec{k}j)$ is the equilibrium number of phonons of wave vector $\vec{k}$ and polarization j frequency at temperature T . This equation has to be supplemented by appropriate boundary conditions at the surface of the sample. These conditions correspond to diffuse scattering or specular scattering, etc.

To solve the Boltzmann equation one therefore needs to know the phonon dispersion relation, the group velocity, and the rate at which collisions occur. The phonon dispersion relation is known from neutron scattering for most materials of interest. Frequencies at special points in the Brillouin zone can be measured by light scattering. Recently, inelastic X-ray scattering has also become useful for measuring phonon frequencies.¹⁶⁷ This has the important advantage that X-rays from a synchrotron source can be focused into a very small volume, thereby enabling small samples to be used. The phonon group velocity can be obtained from the dispersion relation by

$$\vec{v}(\vec{k}j) = \frac{\partial \omega(\vec{k}j)}{\partial \vec{k}}. \quad (9)$$

The variation of group velocity with wave vector is complicated; there are focusing directions in which the group velocity is stationary with respect to variations in $\vec{k}$ and so there are a large number of phonons which propagate in nearly the same direction.^{168–170} But to evaluate the collision term is much harder. It is important to note that to construct the collision term of the Boltzmann equation it is not sufficient to calculate just the lifetime $\tau(\vec{k}j)$ of each phonon $\vec{k}j$. A collision of a phonon $\vec{k}j$ with another phonon inevitably destroys not only the phonon $\vec{k}j$ but also another phonon. In addition, a new phonon is produced. Thus, one needs to know not just the lifetime but also the collision kernel C defined such that

$$\left. \frac{\partial n(\vec{k}j)}{\partial t} \right|_{coll} = -\frac{n(\vec{k}j)}{\tau(\vec{k}j)} + \sum_{\vec{k}'j'} C(\vec{k}j; \vec{k}'j') n(\vec{k}'j') d^3 \vec{k}'. \quad (10)$$

The kernel is not symmetric with respect to the interchange of $\vec{k}j$ with $\vec{k}'j'$; instead,¹⁷¹ it can be shown that

$$C(\vec{k}'j'; \vec{k}j) = C(\vec{k}j; \vec{k}'j') \frac{n(\vec{k}'j')[n(\vec{k}'j') + 1]}{n(\vec{k}j)[n(\vec{k}j) + 1]}. \quad (11)$$

To calculate the rate of anharmonic processes involves the following steps:

1. A model for the harmonic part of the interatomic potential has to be constructed. This can be done using first-principles theory or more simply by choosing a model for the interatomic forces, which gives reasonably good agreement with the measured dispersion relation.
2. One then needs a model for the anharmonic part of the potential. For the purpose of calculating three-phonon processes, this needs to contain terms that are cubic in the atomic displacements. Again, this part of the potential can be obtained from first-principles¹⁷² or by choosing a model that gives agreement with experimentally measured parameters. These parameters might include, for example, the third order elastic constants (known accurately for many materials), and the thermal expansion coefficient and its variation with temperature. In addition, for many materials, experimental values exist for the variation with applied elastic strain of the frequency of zone center optical phonons. (See, for example, the early work on Si and GaP by Weinstein and Piermarini.¹⁷³)
3. From these two models, the matrix elements and the rate of three-phonon scattering can be calculated. These matrix elements involve not just the anharmonic potential and phonon wave vectors but also the polarization vectors of the interacting phonons. The calculation of the matrix elements and the collision kernel C is straightforward but complicated.^{174–176} Because C involves the polarization vectors (quantities which are not directly measurable), it is not easy to estimate the accuracy of the calculation.

The general formula for the phonon lifetime is given in a classic paper by Maradudin and Fein.¹⁷⁷ A phonon k_{1j1} can

be scattered by a phonon $\vec{k}_{2j_2}$ to produce another phonon $\vec{k}_{3j_3}$. In this process, energy and crystal momentum are conserved so that

$$\begin{aligned}\hbar\omega(\vec{k}_{1j_1}) + \hbar\omega(\vec{k}_{2j_2}) &= \hbar\omega(\vec{k}_{3j_3}) \\ \vec{k}_1 + \vec{k}_2 &= \vec{k}_3 + \vec{G},\end{aligned}\quad (12)$$

where $\vec{G}$ is a reciprocal lattice vector. The phonon $\vec{k}_{1j_1}$ can also decay into phonons $\vec{k}_{2j_2}$ and $\vec{k}_{3j_3}$ in which case

$$\begin{aligned}\hbar\omega(\vec{k}_{1j_1}) &= \hbar\omega(\vec{k}_{2j_2}) + \hbar\omega(\vec{k}_{3j_3}) \\ \vec{k}_1 &= \vec{k}_2 + \vec{k}_3 + \vec{G}.\end{aligned}\quad (13)$$

Processes in which $\vec{G} = 0$ are called Normal (N) processes, and those for which $\vec{G} \neq 0$ are Umklapp (U) processes. Umklapp processes are of special importance because these collisions directly reduce the heat flux. Early work on phonon lifetimes was summarized in review articles by Klemens¹⁷⁸ and Carruthers,¹⁷⁹ and in a book by Ziman.¹⁸⁰ Much of this work focused on determining approximate expressions for the phonon lifetime in the limit of low or high frequency, or when the temperature was much less than or greater than the Debye temperature.

Some important results are the following:

1. When the temperature is zero, the phonon lifetime is always limited by boundary scattering. It is also limited by the spontaneous anharmonic decay process governed by the conservation laws (Eqs. (12) and (13)). For longitudinal and fast transverse acoustic phonons, the rate τ^{-1} of this process varies as ω^5 ;¹⁸¹ this assumes that the wave number is small compared to the wave number at the Brillouin zone boundary. A rough estimate of the magnitude of this rate for a number of crystals is given in Ref. 182, and more detailed calculations have been performed by Tamura¹⁸³ and Berke *et al.*¹⁸⁴ For slow transverse phonons, the rate varies as ω^5 when the wave vector lies in some directions, but in other directions, the decay rate is zero because there is no solution to the conditions of conservation of energy and momentum.^{185–188}
2. A discussion of the lifetime of optical phonons at zero temperature has been given by Klemens.¹⁸⁹
3. For temperatures much less than the Debye temperature Θ_D and provided that ω is much less than the Debye frequency, the phonons have wavelength which is much larger than the lattice parameter. The calculation of the phonon lifetime is then simplified because the anharmonic coupling can be expressed in terms of the third order elastic constants. These constants have been measured for many crystals of current technical interest. The phonon lifetime varies with frequency ω and temperature T according to the law

$$\tau^{-1} \propto T^5 f(\hbar\omega/k_B T). \quad (14)$$

For transverse phonons with $\omega \ll k_B T/\hbar$, τ^{-1} is proportional to ωT^4 . Herring¹⁹⁰ has shown that for longitudinal phonons, again with $\omega \ll k_B T/\hbar$, $\tau^{-1} \propto \omega^a T^{5-a}$, where a is an integer, which is larger than 1 and dependent on

the symmetry of the crystal. The lifetime is determined by the collisions of the longitudinal phonon with a transverse phonon and producing a transverse phonon of different polarization. These collisions are called Herring processes. For cubic crystals, the inverse lifetime can be calculated in closed form as¹⁹¹

$$\begin{aligned}\tau^{-1} &= \frac{3\xi(3)\hbar k^2}{4\pi^2\rho^2 c_T^5} \left(\frac{k_B T}{\hbar}\right)^3 \frac{C_{11} + C_{12}}{C_{11} + 2C_{12} + C_{44}} \frac{1}{|C_{11} - C_{12} - 2C_{44}|} \\ &\times \{ (1 - 3G)C_1^2[E(R) - (1 - R)K(R)] \\ &+ 8GC_2^2[K(R) - E(R)] \},\end{aligned}\quad (15)$$

where $\xi(3)$ is the Riemann zeta function, C_{ij} are second order elastic constants, c_T is the sound velocity of transverse waves in the (100) direction, and C_1 and C_2 are related to the second and third order elastic constants by

$$C_1 = C_{155} - C_{144} + 2C_{44}, \quad C_2 = C_{456} + C_{44}. \quad (16)$$

$K(R)$ and $E(R)$ are complete elliptic integrals of the first and second kind with

$$R = (C_{11} - C_{44})(C_{11} + 2C_{12} + C_{44})/(C_{11} + C_{12})^2, \quad (17)$$

and $G = l^2 m^2 + m^2 n^2 + n^2 l^2$ where l, m, n are the direction cosines of $\vec{k}$.

4. For temperatures greater than Θ_D the inverse lifetime for all phonons is proportional to temperature. This result holds provided that three-phonon processes are the most important; at sufficiently high temperatures (for example, near the melting point), there may be a significant contribution from four-phonon processes.
5. In addition to these general results, phonon lifetimes have been calculated for the nearest-neighbor central force model of a fcc crystal with one atom per unit cell. This model has the advantage that the harmonic and anharmonic forces are each described by only one parameter. These parameters can be related to the Debye temperature and the Grüneisen constant γ . The phonon lifetime based on this model was first considered by Maradudin *et al.*,¹⁹² and later investigated in more detail.¹⁹³ The inverse lifetime can be expressed as

$$\frac{1}{\tau(\vec{k}j)} = \frac{\hbar\gamma^2\rho^{2/3}}{M^{5/3}} G_{total}, \quad (18)$$

where ρ is the mass density, M is the mass of each atom, and G_{total} is a dimensionless function of wave vector, temperature, and phonon polarization. As an example of the results obtained, Fig. 5 shows G_{total} for longitudinal phonons propagating along the [100] direction for five different values of the wave number. In the low temperature limit such that $\hbar\omega \gg k_B T$, the inverse lifetime is determined by spontaneous quantum decay; if the frequency is much less than the Debye frequency but still satisfying $\hbar\omega \gg k_B T$, the rate is proportional to ω^5 . At high temperatures, the decay rate is proportional to temperature and increases monotonically with increasing wave number. The results in Fig. 5 are for the total

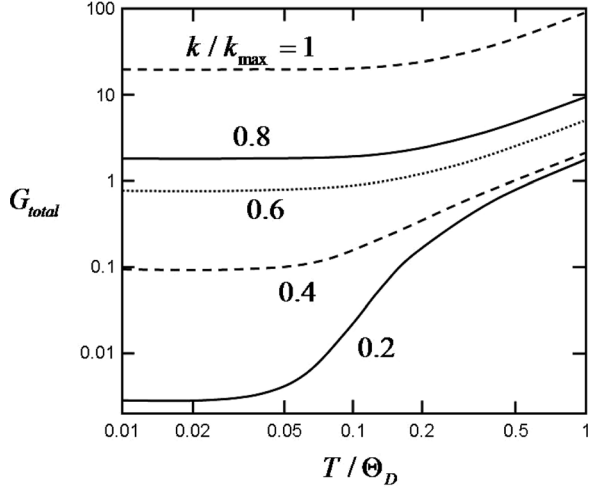


FIG. 5. Calculated value of the parameter G_{total} giving the lifetime of longitudinal phonons propagating in the [100] direction as a function of temperature using the results of Ref. 194. These results are for a simplified model of a fcc crystal. k_{max} is the phonon wave number at the zone boundary.

decay rate, i.e., they include both Umklapp and normal processes. It turns out that the fraction f_U of the scattering events, which are U-processes, varies with wave number in a surprisingly complex way (Fig. 6). At the zone boundary ($k = k_{max}$), there is no difference between a U and an N-process so $f_U = 0.5$. U-processes can occur when $k = 0.8k_{max}$ or $0.2k_{max}$, but there is a range of k between these values in which U-processes cannot occur. The reason for this is discussed in Ref. 193. Because of these complexities, it is difficult to judge the validity of calculations based on highly simplified expressions for the lifetime of phonons such as Klemens' result Eq. (32) that is discussed in Sec. IV. Certainly, that formula does not agree with the exact results for an fcc crystal shown in Fig. 5. It seems likely

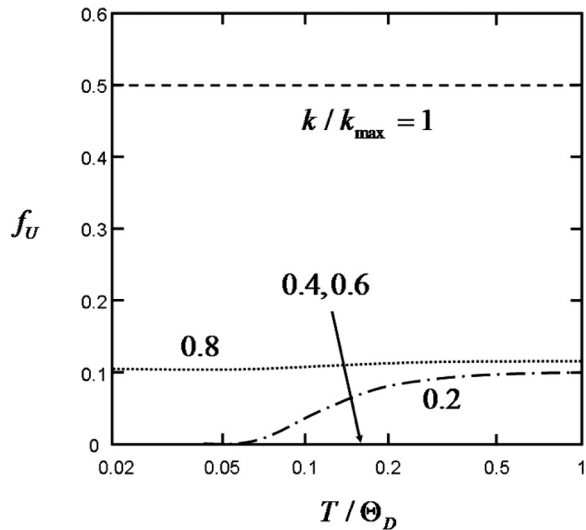


FIG. 6. Calculated fraction f_U of phonon interactions which are U-processes as a function of temperature. Results are for longitudinal phonons propagating in the [100] direction from the results of Ref. 194. These results are for a simplified model of a fcc crystal. k_{max} is the phonon wave number at the zone boundary.

that within the next few years, as the number of first-principles calculations based on the Boltzmann equation increases, we will have a better understanding of the usefulness of the approximate formulas.

6. It is important to note that all of the results discussed so far assume implicitly that we are interested in the lifetime of a phonon of frequency ω such that $\omega\bar{\tau} \gg 1$, where $\bar{\tau}$ is the average lifetime of the thermal phonons. To understand this, first note that for a very low frequency phonon, the lifetime is dominated by collisions with thermal phonons (see Eq. (12)); decay processes (Eq. (13)) are negligible. The finite lifetime of phonons k_{2j2} and k_{3j3} makes their energy uncertain by amounts of $\hbar/\tau(k_{2j2})$ and $\hbar/\tau(k_{3j3})$, respectively. If the energy uncertainty is comparable to or larger than the energy of phonon k_{1j1} the conservation laws break down. One can show that under these conditions, the inverse lifetime varies as $\tau^{-1} \propto \omega^2\bar{\tau}$; this is referred to as the Akhiezer regime.¹⁹⁵ In this regime, one can consider that the inverse lifetime τ^{-1} is the sum of two processes, both of which can be viewed in macroscopic terms.¹⁹⁶ The first is the so-called thermoelastic damping. As the low frequency phonon propagates, it expands and contracts the lattice resulting in temperature fluctuations. The flow of heat between the heated and cooled regions results in entropy production and a damping of the phonon at a rate

$$\tau_{thermo}^{-1} = \frac{[e_\alpha(\vec{k}j)\kappa_\beta C_{\alpha\beta\gamma\delta}\alpha_{\gamma\delta}]^2 T \kappa_{\lambda\mu} \kappa_{\lambda\mu}}{\rho \omega(\vec{k}j)^2 C^2}, \quad (19)$$

where e_α are the elements of the phonon polarization vector, $\kappa_{\lambda\mu}$ are the elements of the thermal conductivity tensor, $C_{\alpha\beta\gamma\delta}$ are the second order elastic constants in tensor notation, and C is the specific heat per unit volume. The second mechanism is by phonon viscosity, also called the Akhiezer process. The fluctuating strain associated with the low frequency phonon raises the frequency of some of the high frequency ($\omega \sim k_B T/\hbar$) phonons and lowers the frequency of others. This creates a non-equilibrium phonon distribution which produces entropy as it relaxes. The damping rate by this process is

$$\tau_{vis}^{-1} = \frac{e_\alpha(\vec{k}j)\kappa_\beta e_\gamma(\vec{k}j)\kappa_\delta v_{\alpha\beta\gamma\delta}}{\rho}, \quad (20)$$

where $v_{\alpha\beta\gamma\delta}$ are the elements of the phonon viscosity tensor. Both of these processes give a damping proportional to $\omega^2\bar{\tau}$, but generally the viscosity contribution is larger than the thermoelastic term. The damping rate in the transition regime where $\omega\bar{\tau}$ goes from less than one to greater than one is complicated, and is discussed in Refs. 197 and 198. It is important to note that these are two distinct regimes in which separate formulas apply. Thus, in the $\omega\bar{\tau} > 1$ regime, one should calculate the damping rate using the phonon collision approach; a contribution from the thermoelastic and Akhiezer mechanisms should not be added in.

The other important scattering process in crystals which are free of defects is isotope scattering. In a cubic crystal with a single atom per unit cell, it is easy to show that the phonon lifetime is given by¹⁷⁹

$$\frac{1}{\tau(\vec{k}j)} = \frac{\pi}{6} V_0 g_2 \omega^2(\vec{k}j) D(\omega(\vec{k}j)), \quad (21)$$

where V_0 is the volume per atom, and g_2 is given by

$$g_2 = \sum_i f_i \left(\frac{\delta m_i}{\langle m \rangle} \right)^2, \quad (22)$$

where i labels the different isotopes, f_i is the fraction of isotope i , δm_i is the deviation of the mass of isotope i from the average mass $\langle m \rangle$. D is the phonon density of states per unit volume. It can be shown that the same formula also holds for silicon and germanium.¹⁹⁴ Isotope scattering has some similarities to the Barkla effect for X-rays. The scattering from state $\vec{k}j$ to $\vec{k}'j'$ is proportional to $[\vec{e}(\vec{k}j) \cdot \vec{e}(\vec{k}'j')]^2$. Thus, for example, for scattering at an angle of $\pi/2$ a longitudinal phonon $\vec{k}$ is most likely to be scattered into a transverse phonon with polarization vector in roughly the same direction as $\vec{k}$.

Isotope scattering is another example of a process for which it would be helpful to have a better understanding of the usefulness of simplified formulas. If we compare Eq. (21) with the Klemens' formula Eq. (33) discussed in Sec. IV, one can see immediately that the difference between Eqs. (21) and (33) is simply that the phonon density of states has been approximated by the Debye model, i.e., has been taken to be proportional to frequency squared. In fact, of course, the density of states $D(\omega)$ goes as ω^2 only at low frequencies. At larger ω , $D(\omega)$ usually contains strong peaks, which come from regions of the Brillouin zone where the phonon group velocity is small. In the frequency range of these peaks, the true value of D is much larger than the Debye estimate. Thus, one can see that Eq. (33) underestimates the scattering rate of the phonons with low group velocity and probably overestimates the scattering of phonons with less dispersion.

Once C is calculated, the final step is to solve Eq. (7) to find the phonon distribution. The heat current $\vec{j}$ can then be obtained from

$$\vec{j} = \sum_{\vec{k}j} n(\vec{k}j) \vec{v}(\vec{k}j) \hbar \omega(\vec{k}j). \quad (23)$$

From the discussion above, it is clear that even for a bulk crystal the calculation of the thermal conductivity is challenging. One way to simplify the problem is to ignore the collision kernel and write

$$\left. \frac{\partial n(\vec{k}j)}{\partial t} \right|_{\text{coll}} = -\frac{n(\vec{k}j)}{\tau(\vec{k}j)}. \quad (24)$$

This leads to the “relaxation time approximation” for the thermal conductivity tensor

$$\kappa_{\alpha\beta} = \sum_{\vec{k}j} c_{\vec{k}j} v_{\alpha}(\vec{k}j) v_{\beta}(\vec{k}j) \tau_{\vec{k}j}, \quad (25)$$

where $c(\vec{k}j)$ is the contribution to the specific heat from the phonon mode $\vec{k}j$. For a cubic crystal, the conductivity simplifies to become

$$\kappa = \frac{1}{3} \sum_{\vec{k}j} c_{\vec{k}j} v^2(\vec{k}j) \tau_{\vec{k}j}. \quad (26)$$

There are several well known difficulties with the use of this formula; an old but still very useful discussion of some of the problems is given in Ziman's book.¹⁸⁰ The major difficulty with this formula is that it is not clear whether every type of scattering process should be included in the calculation of $\tau_{\vec{k}j}$. Anharmonic processes of the N type do change the direction of the scattered phonon but leave the total flux of energy in a given direction essentially unchanged. Thus, one can argue that these processes do not contribute to the thermal resistance and so should be ignored and only U processes considered. But suppose that there is a particular group of phonons which cannot be scattered in a U process. Then, if N processes are ignored, the thermal conductivity would be infinite, which is clearly impossible. To treat this situation correctly, it is necessary to go beyond the relaxation time approximation and include the effects of both N and U processes in an appropriate way.¹⁸⁰

To overcome these difficulties, quantitative calculations using the full Boltzmann equation have been made. In 1996, Omini and Sparavigna¹⁷⁴ reported calculations for Ar and Kr using an accurately calculated collision kernel together with a solution of the Boltzmann equation. For these materials, the interatomic potential is relatively simple. More recently, Broido and coworkers^{176,199,200} have published a series of papers in which they report calculations of this type for silicon, germanium, and diamond. These calculations are considerably more difficult because of the more complicated form of the interatomic potential. The results are in good agreement with experiment.

Of course, in the numerical calculations of both Sparavigna and of Broido *et al.*, the information about the lifetime and the collision kernel is contained within the program. It would be very valuable to extract this information and in this way learn more about the detailed phonon processes that take place when heat flows. Which phonons carry most of the heat? How does the phonon lifetime vary with wave vector and polarization? What is the ratio of normal processes to Umklapp processes and which phonons are involved in U-processes? Sparavigna does not provide a plot of this information, but Ward and Broido²⁰¹ do present a plot of the lifetime of longitudinal phonons at 300 K averaged over all directions (Fig. 7). They also divide the contributions to the lifetime into U and N-processes. Lindsay and Broido²⁰² have found correlations between the region of phase space available for three-phonon processes and the thermal conductivity. Study along these lines has the potential to give an indication about the validity of some of the approximations which are often made in theories of the bulk thermal conductivity. For example, it is usually assumed that although the optic modes may affect the heat flow by contributing to the scattering rate of the acoustic phonons, the actual heat carried by these modes is unimportant. It would be good to know how reliable an approximation this is.

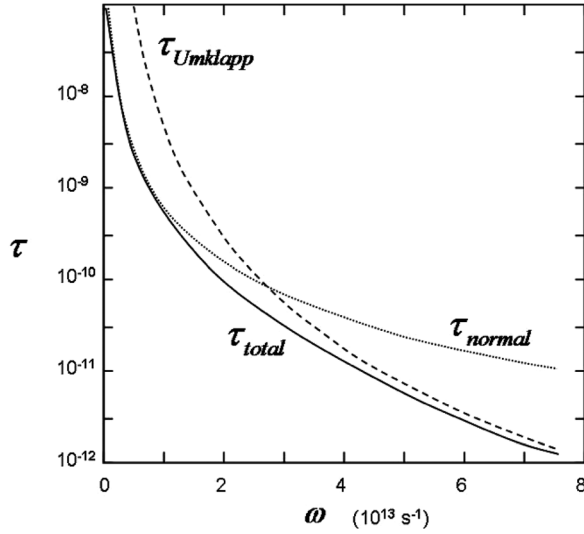


FIG. 7. Calculated lifetime of longitudinal phonons in silicon at 300 K as a function of frequency from Ref. 201.

To calculate the heat flow in a simple nanostructure, such as a nanowire composed of a single material, it is sufficient to use the same phonon lifetime and collision kernel as for a bulk material. This approach will break down only when the lateral dimensions of the structure become comparable to the phonon wavelength. However, the calculation of heat flow is much more difficult than for the bulk because now the phonon distribution function $n(\vec{k}_j)$ varies across the transverse dimensions of the sample. In a cylindrical nanowire, for example, the phonon distribution varies with distance r from the axis, and the solution for the phonon distribution function has to satisfy the Boltzmann equation at each value of r . Calculations for a diamond nanowire have recently been reported;²⁰³ it was necessary to make some approximations in order to reduce the computer time that was needed. In a single-walled carbon nanotube (SWCNT), the situation is simpler, since the phonons only exist at one value of r . Calculations of the thermal conductivity in a SWCNT²⁰⁴ and in graphene²⁰⁵ have been performed by Lindsay *et al.* They do not give details of the phonon lifetimes but do discuss the relative importance of N and U processes.

Of course, if the mean free path is very large compared to all sample dimensions (the Casimir regime) the problem is tractable. It is then sufficient to simply add up the contribution from each of the phonon modes which are thermally excited. For each mode, one can calculate the distance $\Lambda_{\vec{k}_j}$ that a phonon leaving a point on the surface of the structure travels in the direction of heat flow before being scattered at another point on the surface; this distance depends on the particular point from which the phonon starts. For a uniform rod of arbitrary cross-section, one can show²⁰⁶ that the conductivity is given by the formula

$$\kappa = \frac{\hbar^2}{16\pi^3 k T^2 A} \oint dl \sum_j \int \int \int \vec{v}_{\vec{k}_j} \cdot \hat{e}(l) n_{\vec{k}_j} \times (n_{\vec{k}_j} + 1) \Lambda_{\vec{k}_j}^2 \hbar \omega_{\vec{k}_j} k^2 dk d\cos\theta d\phi, \quad (27)$$

where A is the cross-sectional area, the line integral runs around the boundary of the rod, and $\hat{e}(l)$ is a vector normal to the surface at the position l . Note that the contribution from a mode depends, amongst other things, on the square of $\Lambda_{\vec{k}_j}$.²⁰⁶ A change in modes for which the group velocity vector lies in a direction close to the direction of heat flow clearly make a large contribution. In fact, one can show that this contribution is so large that for heat flow in the in-plane x -direction of a plate of thickness in the y -direction D and width in z of nD , the thermal conductivity diverges logarithmically as n goes to infinity.²⁰⁷ Quantitative calculations of the Casimir regime heat flow in cylindrical nanowires and plates of Si and GaAs allowing for the details of the phonon dispersion relation and the variation of the phonon group velocity with wave vector and polarization have been presented in Ref. 206.

For a nanostructure with a complex geometry and composed of more than one material, new issues arise. As an example, consider a superlattice with repeat distance d . If the phonon mean free path in the bulk materials is larger than d , then it is necessary to first calculate the phonon dispersion relation in the superlattice and then calculate the phonon lifetime and collision kernel. Ward and Broido²⁰¹ have calculated the thermal conductivity of a series of Si/Ge and GaAs/AlAs superlattices. For the GaAs/AlAs structures, they find a theoretical conductivity substantially higher than the experimentally measured thermal conductivity. They attribute this to the effect of roughness of the interfaces.

B. Nanowires and graphene: Experiment and theory

Over the past decade, the establishment of several nano-scale thermal transport measurement techniques has made it possible to use nanostructures as model systems to test the phonon transport theories reviewed in Sec. III A. Compared to bulk crystals, phonon transport in nanostructures is unique in the importance of phonon-boundary scattering as well as confinement of phonons in low-dimensional structures such as nanotubes and graphene. In the following, the assumptions behind the particle transport theories of phonon-boundary scattering are examined, followed by a discussion of a number of phonon wave interference models that have emerged over the past decade for explaining the thermal conductivity of some nanostructures. Besides the central issue regarding whether wave interference effects are observable in the thermal conductivity measurements, the size dependence and interface scattering of phonon transport in one-dimensional (1D) nanotubes and two-dimensional (2D) graphene are reviewed.

1. Particle transport description of phonon-boundary scattering

Casimir²⁰⁸ employed a radiation analogue and the Debye approximation to analyze phonon transport in a rod in the low temperature limit where internal scattering of phonons was negligible. The rod surface was treated as a black-body that absorbed all the incident phonons and re-emitted phonons diffusely according to the local equilibrium distribution. Under these assumptions, the average phonon-

boundary scattering mean free path Λ_B is the diameter d for a circular rod, and $1.12W$ for a rod with a square cross section of width W .

Building upon Casimir's result²⁰⁸ and several other studies,^{209–211} Ziman¹⁸⁰ developed a solution of the phonon Boltzmann transport equation (BTE) in a rod under the relaxation time approximation. (Although the Debye approximation was mentioned in Ziman's derivation,¹⁸⁰ the assumption of an isotropic crystal is sufficient to arrive at his result without the need of assuming a frequency-independent phonon group velocity.) For diffuse boundary condition at the rod surface and absence of internal scattering, the boundary mean free path given by the BTE solution¹⁸⁰ is the same as Casimir's result ($\Lambda_{B,C}$), and is essentially an average over all orientations and positions of the distance traveled by phonon quasi-particles originated from the cross section and intercepted by the rod surface. For a partially specular and partially diffuse surface, the specular parameter p is used to describe the probability for specular scattering at the surface. Under the isotropic assumption, the phonon BTE can be solved to obtain the boundary scattering mean free path as $\Lambda_B = \Lambda_{B,C}(1+p)/(1-p)$.¹⁸⁰

Because the phase velocity is not collinear with the group velocity in an anisotropic crystals, McCurdy *et al.*²⁰⁷ calculated that the Λ_B of a Si rod of diffuse boundary condition ranges between $1.66\Lambda_{B,C}$ and $0.97\Lambda_{B,C}$ when the rod axis is aligned along the $\langle 100 \rangle$ and $\langle 111 \rangle$ direction, respectively, and obtained low-temperature thermal conductivity data that revealed this phonon focusing effect.

With internal scattering due to phonon-phonon processes or impurities accounted for, the BTE can be solved under the isotropic assumption to yield that the total mean free path (Λ) differs only slightly from that given by the Matthiessen's rule, $1/\Lambda = 1/\Lambda_B + 1/\Lambda_i$, where Λ_i is the mean free path due to internal scattering processes.^{180,210} Wang and Mingo²¹² recently presented BTE solutions of Λ for nanowires of different cross sections and 2D nanoribbons with two diffuse edges, and verified the accuracy of the Matthiessen's rule only for the nanowire case, because diffuse boundary scattering alone does not lead to a finite mean free path in 2D nanoribbons, similar to the case of the logarithmic divergence for an infinite thin film discussed at the end of Sec. III A.

These particle transport theories have been used to analyze the thermal conductivity data of nanostructures measured by a number of methods that have been established over the past decade^{8,9,12} In addition to the axial thermal conductivity of different nanowire structures, the thermal conductivity of thin films²¹³ and periodic nanoporous membrane structures^{214,215} along the in-plane direction have been obtained with the use of suspended micro-devices with built-in resistance thermometers.²¹⁶ TDTR techniques have also been employed to measure the cross-plane thermal conductivity of nanowire arrays.^{217,218} The theoretical framework of Casimir and Ziman can explain well the suppressed thermal conductivity found in InAs nanowires,^{217,219} Bi₂Te₃ nanowires,²²⁰ SnO₂ nanobelts,²²¹ and Si nanowires²²² grown by a vapor liquid solid (VLS) method with a diameter larger than about 30 nm, as well as Bi nanowires,²²³ InSb

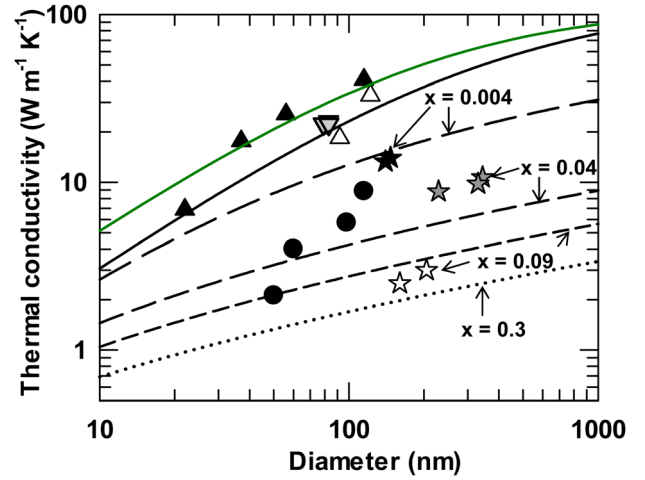


FIG. 8. Thermal conductivity as a function of diameter for Si and Si_{1-x}Ge_x nanowires. Except for those labeled in the figure, the data are for Si nanowires or $x = 0$. The green line is the calculation result in Ref. 231 for the case of Si with 10^{19} cm^{-3} boron doping. The black lines are the calculation results in Ref. 228. The symbols are experimental data reported in Ref. 222 for VLS Si nanowires (black triangles), in Ref. 232 for RIE patterned Si nanowires (gray down triangles), in Ref. 230 for electroless etched Si nanowires (black circles), and in Ref. 225 for Si nanowires (unfilled triangles) and Si_{1-x}Ge_x nanowires.

nanowires,²²⁴ and SiGe nanowires²²⁵ when additional impurity and defect scattering were considered.

Figure 8 includes a comparison between the measured thermal conductivity of some Si and SiGe nanowires and two calculated thermal conductivity results based on Callaway-type models, Mathessens rule, bulk phonon dispersion, and diffuse phonon-surface scattering. Because the thermal conductivity of bulk Si is dominated by phonons with mean free paths longer than 100 nm, the size effect on the thermal conductivity can be readily observed in Si nanowires. Because the relative thermal conductivity contribution of low-frequency phonons with long mean free paths is decreased by diffuse boundary scattering, the relative contribution from the high-frequency or short-wavelength zone boundary phonons increases in a nanowire. Consequently, the calculated thermal conductivity becomes more sensitive to the choice of the cutoff frequency in the Callaway-type model for nanowires than for bulk crystals.²²⁶ As shown in Fig. 8, the use of the same bulk cutoff frequency given by Morelli *et al.*²²⁷ combined with diffuse phonon-boundary scattering yields approximately the experimental thermal conductivity values reported by VLS Si nanowires with a diameter larger than 30 nm.²²²

Using a cutoff frequency adjusted to fit the data for one nanowire, Wang and Mingo²²⁸ obtained somewhat lower thermal conductivity values for the diffuse boundary condition. Based on the results of Wang and Mingo,²²⁸ phonons scattered by the surface of the VLS Si nanowires of diameter larger than 30 nm should contain a small specular component together with a large diffuse component. In addition, the low-temperature thermal conductivity data measured by Heron *et al.*²²⁹ for patterned Si nanobeams were explained by an increased specular component with increasing phonon wavelength or decreasing temperature. By accounting for alloy

scattering of short-wavelength phonons and diffuse boundary scattering of all phonons, the calculation by Wang and Mingo²²⁸ was used to fit the experimental thermal conductivity values reported for $\text{Si}_{1-x}\text{Ge}_x$ nanowires,²²⁵ which suggested that alloy scattering instead of surface scattering should be the dominant scattering mechanism. However, without considering scattering by high-concentration defects, impurities or their clusters, these Callaway-type models suggest that the Casimir limit of diffuse surface scattering alone is insufficient to suppress the thermal conductivity to the experimental values reported for electroless etched rough Si nanowires,²³⁰ as shown in Fig. 8.

Other types of scattering phase function at the surface can in principle lead to shorter Λ_B than the diffuse boundary condition. For example, the Monte Carlo (MC) phonon particle transport simulation result of Moore *et al.*²³³ suggested that adding periodic sawtooth features of a sufficiently large height to width ratio to a Si nanowire of $22 \times 22 \text{ nm}^2$ cross section can suppress the thermal conductance despite the increase in the solid volume of the nanowire. This effect has been attributed to the presence of a large backscattering component of the scattered phonons relative to the phonons incident on the sawtooth features. However, the room-temperature thermal conductivity obtained for the $22 \times 22 \text{ nm}^2$ nanowire with 4-nm-high, 2-nm-wide sawtooth features is still four times larger than the reported experimental value shown in Fig. 8 for the $\sim 50 \text{ nm}$ diameter rough Si nanowire. In addition, a scattering phase function with a large component normal to the nanowire axis can also reduce the average v_z^2 to be smaller than the $|v_g|^2/3$ value for the diffuse scattering case in an isotropic crystal, where v_g and v_z are the group velocity and its component along the nanowire axis. However, within the particle transport description used in the relaxation time approximation or MC solution of BTE, no mechanisms have been found to yield a backscattering or normal scattering component that is sufficiently large to produce the experimental thermal conductivity values reported for rough Si nanowires.

2. Wave interference models of phonon transport in silicon nanostructures

The seeming discrepancy between the experimental data and the phonon particle transport theories has stimulated a number of investigations of different wave interference effects that are not accounted for by the particle description of phonon transport. For example, it has been suggested that phonon interference can result in phononic band gaps and reduce the phonon group velocity, which has been used to explain the experimental values of the in-plane thermal conductivity of Si nanomesh.²¹⁴ The reported room-temperature upper bound value of $2 \text{ W m}^{-1} \text{ K}^{-1}$ for the Si nanomeshes is about a factor of two lower than the already surprisingly small thermal conductivity of $\sim 3.5 \text{ W m}^{-1} \text{ K}^{-1}$ for patterned Si nanowires of a comparable critical dimension of $\approx 20 \text{ nm}$.²¹⁴ If the measurement result of the Si nanomesh of periodic holes was analyzed with the use of effective medium theories^{234,235} that do not account for any size effects on the solid thermal conductivity, the obtained room-temperature solid

thermal conductivity would be as low as about $1.4 \text{ W m}^{-1} \text{ K}^{-1}$, comparable to those reported for amorphous Si and SiO_2 .²³⁶ It is worth noting that the reported thermal conductivity of the nanomesh increases with temperature at near room temperature. However, if reduced phonon group velocity was the only reason for the very low thermal conductivity of the nanomesh, the room-temperature thermal conductivity would be expected to decrease with increasing T because of increasing umklapp scattering.

Moreover, confinement of phonon waves along the radial direction in a small-diameter nanowire can in principle modify the phonon dispersion because of interference of the waves reflected by the nanowire surface. Some earlier theoretical works suggested that the phonon group velocity component along the nanowire axis of a 20-nm-diameter Si nanowire is reduced to be about half of the v_g value in the bulk,²³⁷ suppressing the thermal conductivity. However, the axial component of the phonon group velocity in a nanowire is equivalent to the bulk v_z instead of the bulk v_g . For Si nanowires of a diffuse surface and a diameter of 20 nm and larger, Mingo²²⁶ calculated that the phonon dispersion is rather close to the bulk dispersion and the room-temperature thermal conductivity calculated based on the actual dispersion are similar to those calculated based on the bulk phonon dispersion, because the energy separation between 1D phonon sub-bands in the nanowire is much smaller than $k_B T$ for temperature $T > 40 \text{ K}$ in a 20 nm Si nanowire.

In addition, variation of the cross section due to the surface roughness of a nanowire can in principle result in a perturbation in the allowed wave vector states (q_{mn}) perpendicular to the transport direction in the nanowire, because $|q_{mn}|$ scales with $\pi/d(x)$ with $d(x)$ being the local nanowire diameter. This modulation is proportional to σ/D , where σ is the root mean square (rms) height of the surface roughness and D is the average nanowire diameter. The effect is equivalent to a perturbation of the local phonon dispersion $\omega(q)$ along the nanowire. Martin *et al.*²³⁸ employed first order perturbation theory, also referred as the Born approximation, to calculate the volumetric scattering rate caused by this perturbation. Their results²³⁸ suggest that the thermal conductivity in the surface roughness scattering-dominant regime is proportional to $(D/\sigma)^2$.

Sadhu and Sinha²³⁹ employed a different Bourret approximation and the Green's function approach to calculate the phonon transmission coefficient in 50-nm-diameter Si nanowires of different σ and L_c values. Besides the σ/D ratio, their result suggested that the σ/L_c ratio is another critical parameter. By setting $\sigma = 2.5 \text{ nm}$ and $L_c = 30 \text{ nm}$ to increase the σ/L_c ratio, Sadhu and Sinha²³⁹ obtained a transmission coefficient that decreases with increasing frequency to be well below the Casimir limit, and used the result to match the experimental data for rough Si nanowires with $\sim 50 \text{ nm}$ diameter.

A different atomistic Green's function calculation by Carrete *et al.*²⁴⁰ suggested that the transmission coefficient can be suppressed to be one order of magnitude smaller than that given by Casimir's limit only for a very large σ/D value exceeding 20%. These authors²⁴⁰ emphasized that the Born approximation is invalid when the surface roughness is

on the order of wavelength and can underestimate the transmission coefficient by one order of magnitude.

It is worth noting that the surface amorphous layer on a nanowire was shown to considerably reduce the thermal conductivity of a 15 nm diameter Si nanowire in a molecular dynamics simulation²⁴¹ and in an experiment with Si crystals.²⁴² The presence of an amorphous native oxide on the surface of Si nanostructures can destroy the phase coherence of phonons transmitted into the amorphous shell and back into the crystalline nanowire core.

A recent MD simulation²⁴¹ has suggested that interior defects such as grain boundaries are needed to obtain the abnormally low thermal conductivity in the experimental Si nanowires. While it is unclear how grain boundaries could have formed in the nanowires etched from single crystal silicon, it has been found that the electroless etching process can create nanopores in nanowires etched from highly doped Si wafers.²⁴³ In addition, RIE was shown to cause damage in a 30-nm-thick surface layer of Si.²⁴⁴ Although nanopores were not observed by transmission electron microscopy (TEM) in nanowires etched from low-doped wafers,²³⁰ TEM is effective in detecting extensive defects such as grain boundaries or pores but not local defects such as point defects, impurities and their clusters.

Hence, further experimental and theoretical studies are needed to clarify the cause of the unusually low experimental thermal conductivity of Si nanostructures. The recent combined structure-thermal transport measurement of Si nanowires has represented an important progress toward this direction.²⁴⁵ It is found that the thermal conductivity of the rough Si nanowires shows a much clearer correlation with σ/L_c than D , σ , and σ/D . The dependence of the thermal conductivity on the σ/L_c ratio has not only been suggested by the coherent scattering model of Sadhu and Sinha,²³⁹ but also revealed by the MC phonon particle transport simulation of Moore *et al.*²³³

3. Phonon scattering in graphene and nanotubes

Graphene and carbon nanotubes are unique in their atomically smooth surfaces and confinement of phonons to the atomic scale along one or two dimensions. These distinct features can also be found in the hexagonal boron nitride counterparts. The phonon dispersion of suspended single-layer graphene (SLG) is different from that of graphite in the absence of the cross-plane modes. Appreciable difference also appears in the basal-plane modes between graphene and highly oriented pyrolytic graphites (HOPG), also referred to as Bernal or 3D graphite.²⁴⁶ In addition to the approximately linear longitudinal acoustic (LA) and in-plane transverse acoustic (TA), the out-of-plane polarized, in-plane propagating transverse acoustic modes, the so-called flexural or ZA modes, show a quadratic dispersion in SLG. Such quadratic ZA dispersion can also be found in graphite except in the long-wavelength limit, where interlayer interaction breaks the rotational symmetry and the energy cost for rotation of the graphene sheet results in a linear term in the ZA dispersion that dominates over the quadratic term.²⁴⁷ Although the acoustic branches of graphite are similar to those in SLG,

and the low-frequency longitudinal optical (LO) and transverse optical (TO) branches of HOPG are nearly degenerate with the LA and TA branches, the additional low-frequency out-of-plane optical (ZO') branch in HOPG lies appreciably above the ZA branch.^{248,249} Because the cross-plane modes and ZO' modes can scatter the basal-plane acoustic modes, the absence of these scattering channels should result in higher basal-plane thermal conductivity of suspended flat SLG than that of graphite.

Moreover, because of the reflection symmetry in flat suspended SLG, three-phonon scattering processes involving the ZA branch must involve an even number of ZA phonons.²⁰⁵ The selection rule restricts the phase space for ZA phonon scattering. Because of the reduced ZA scattering rate and the large density of states of the ZA phonons, it was suggested that the ZA contribution dominates the thermal conductivity of flat suspended SLG,²⁰⁵ contradicting earlier calculations²⁵⁰ that did not use this selection rule or calculate the actual phase space for phonon-phonon scattering. Together with scattering by cross-plane and ZO modes, it was suggested that breaking of the reflection symmetry in few-layer graphene (FLG) relaxes the selection rule, decreasing the basal plane thermal conductivity to approach the graphite limit when the FLG layer thickness is increased to be above five layers.²⁵¹

When a graphene sheet is rolled up and joined into a single-walled CNT, the curvature can also break the reflection symmetry, and increase the ZA phonon scattering rate. It has been reported that this effect results in a decreasing thermal conductivity when the SWCNT diameter is decreased to a critical value, 1.5 nm for a 3- μ m-long long SWCNT.²⁵² For diameters below this critical value, increasing confinement of the phonon waves along the circumference of the SWCNT increases the energy separation between 1D phonon subbands and pushes the optical modes to higher frequencies. Consequently, scattering between the optical modes and the acoustic modes is suppressed in small-diameter SWCNTs, increasing the thermal conductivity with decreasing diameter. Besides the FLG thickness and SWCNT diameter dependence, it has also been suggested that the thermal conductivity of SLG and SWCNTs increases with the lateral dimension,^{253,254} because first-order three-phonon processes are insufficient to prevent the divergence of the thermal conductivity contribution from low-frequency acoustic modes that play a larger role in lower dimensions. The predicted size dependence is for the case of clean, suspended defect-free graphene and nanotubes, because supporting or embedding graphene and nanotubes on a substrate or in a medium may cause interface phonon-scattering as well as breaking of the reflection symmetry. In addition, higher-order phonon-phonon processes and scattering by defects, isotope impurities, and other structure disorders can also remove the predicted length dependence.²⁵⁴

Over the past decade, the establishment of a number of nanoscale thermal transport measurement methods has enabled the experimental investigation of phonon transport in nanotubes and graphene, as discussed in several recent reviews.^{8,9,231} Results from a micro-Raman spectroscopy measurement have been used to support the theoretical

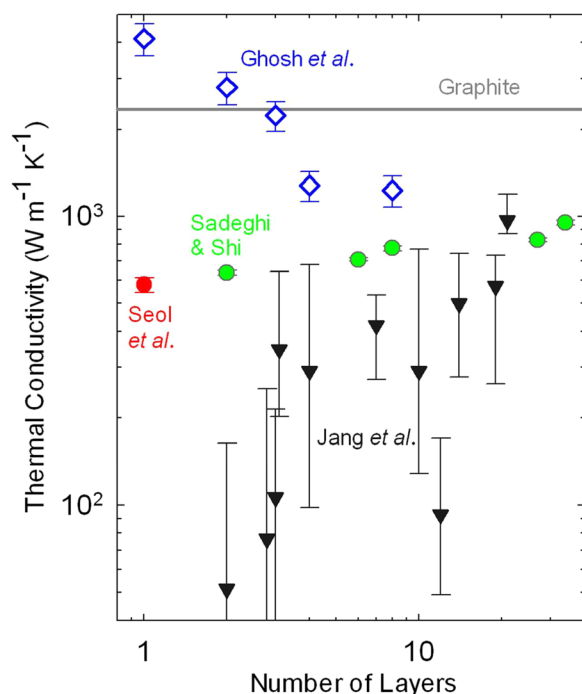


FIG. 9. Reported thermal conductivity near room temperature as a function of number of layers for suspended graphene reported by Ghosh *et al.*,²⁵⁵ graphene supported on amorphous SiO₂ reported by Seol *et al.*,²⁶⁰ and Sadeghi *et al.*,⁹ and graphene encased in amorphous SiO₂ reported by Jang *et al.*²⁶¹

prediction of increasing basal plane thermal conductivity with decreasing FLG layer thickness and suggest that the thermal conductivity approaches the graphite value when the FLG layer thickness increases to about eight layer,²⁵⁵ as shown in Fig. 9. Other Raman measurements^{256–259} yielded lower optical absorbance and thermal conductivity values for suspended SLG than the initial work,⁵⁹ as shown in Fig. 10. In addition to large uncertainties in the optical absorbance, the limited temperature sensitivity of the Raman thermometry method has prevented experimental verification of the lateral size dependence of the thermal conductivity of suspended graphene.²⁵⁸ It is also necessary to verify the reported thickness dependence with other experimental techniques.

Different designs of micro-devices with built-in resistance thermometers have been employed to observe decreasing thermal conductivity of FLG supported on^{260,262} or embedded in²⁶¹ SiO₂ with decreasing thickness, as shown in Fig. 9. Raman spectra obtained on the supported SLG²⁶⁰ and FLG²⁶² samples do not contain D peaks associated with defects, with a large electron mobility found in the supported SLG sample.²⁶⁰ In addition, the experimentally determined acoustic phonon dispersions of a graphene monolayer weakly interacting with a substrate^{264,265} are similar to those of suspended SLG except for ZA phonons near the zone center. Hence, the thermal conductivity suppression found in the supported graphene has been attributed to phonon scattering by the graphene-SiO₂ interface. Compared to the results reported for suspended FLG, the opposite trend is found for supported and embedded FLG and indicates much stronger interface scattering than interlayer scattering within the FLG. The results from the supported FLG further suggests

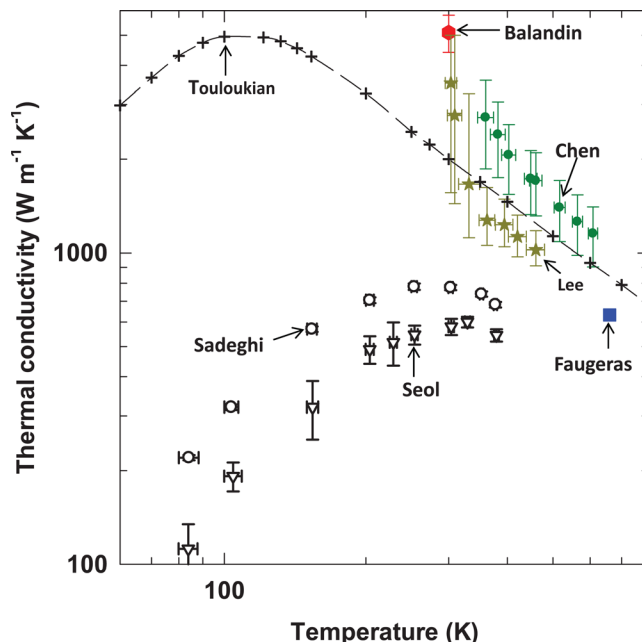


FIG. 10. Experimental basal plane thermal conductivity of suspended SLG reported by Balandin *et al.*,⁵⁹ Faugeras *et al.*,²⁵⁶ Chen *et al.*,²⁵⁸ and Lee *et al.*,²⁵⁹ SLG supported on SiO₂ by Seol *et al.*,²⁶⁰ and eight-layer graphene supported on SiO₂ by Sadeghi *et al.*²⁶² Shown for comparison is the value recommended for high-quality graphite by Touloukian.²⁶³

that the interface scattering can impact the transport of phonons in graphene sheets more than tens of layers away from the interface. A key question is whether the strong interface scattering is simply caused by the roughness of the SiO₂ surface, or largely due to transmission or leaking of the scattered phonons across the interface.²⁵³

C. Applications to information technology and high-power density electronics

Modern semiconductor devices and their constituent materials and interfaces present some of the most promising and challenging opportunities for the study of nanoscale heat conduction. Example technologies and devices include highly scaled planar or 3-dimensional (3-D) field-effect transistors,^{266,267} phase change memory devices based on chalcogenides,^{268,269} photonic crystal lasers and waveguides for optical interconnects,^{270,271} and composite substrates for power semiconductor devices.^{272–274} The temperature distributions in these structures, especially near interfaces, consume an increasing fraction of the available “thermal budget” for electronic, optoelectronic, and amplifier circuitry. While a sub-continuum analysis is not always essential for the accurate simulation of these structures, this situation is gradually changing with continued dimensional scaling, heterogeneous materials integration, and geometries targeting electron and photon band manipulation. Section III C summarizes three areas of special relevance and proposes future nanoscale experiments and simulations.

1. Data storage

While heat generation and transport in logic^{266,275} and interconnects^{276,277} have received much attention, thermal

phenomena in data storage technologies are also very relevant due to leakage in volatile memory and to the sheer number of memory elements employed. In this context, phase change materials (PCMs) have attracted wide interest for memory and reprogrammable circuits with low voltage operation, fast switching, and high endurance.^{268,269} PCMs are chalcogenides like $\text{Ge}_2\text{Sb}_2\text{Te}_5$ (GST) which have amorphous and crystalline phases with a large ratio ($>100\times$) of electrical resistivity. The data in electrically programmable PCMs are stored as changes in bit resistance, which can be reversibly switched with short voltage pulses and localized Joule heating. However, a drawback of PCMs has been their high programming current ($>0.1\text{ mA}$), as Joule heat must be delivered to a finite bit volume.

Figure 11 shows the distribution of amorphous and polycrystalline material in a phase change random access memory (PCRAM) cell that has been subjected to varying levels of programming current. The key metrics for contemporary devices include the switching energy and time, which are governed by heat conduction and crystallization kinetics in regions of GST alloys with dimensions below 10 nm. For exploratory phase change devices, such as those developed for brain-inspired computing,²⁷⁸ nanoscale heat conduction governs the staged transition from the amorphous to the crystalline phase during pulsed electrical heating. The temporal stability of the amorphous region is another challenging research topic,^{279,280} which is critical for advanced cell designs including those targeting multibit operation.^{281,282}

Thermal transport studies in PCMs examined the impact of interface properties, phase purity, and the relative contributions of electrons and phonons to transport.²⁸⁴ The temperature-dependent thermal conductivities of the amorphous and polycrystalline face centered cubic (fcc) and hexagonally close packed (hcp) phases of GST films and several stoichiometric variants were measured using time-domain

thermoreflectance²⁸⁵ and nanosecond transient thermoreflectance (TTR).²⁸⁶ Electrons contribute $\sim 1\text{ Wm}^{-1}\text{K}^{-1}$ to the hcp crystalline phase and help to nearly quadruple the thermal conductivity relative to the amorphous and fcc phases. By exploring stoichiometric variations including pure GeSb, the electron contribution can be varied²⁸⁷ by nearly a factor of 5. Electrothermal measurements using suspended nanostructures and the 3ω method identified strong anisotropy in mixed phase films, which was attributed to the alignment of amorphous material along vertically oriented grain boundaries.²⁸⁸ Thermal property data remain urgently needed for the melted condition.

Thermal boundary resistances (TBRs) of PCMs with surrounding passivation and electrode materials (e.g., SiO_2 , SiN_x , and TiN) are also key for the energy-efficient operation of a memory cell.²⁸⁹ The TBRs of $\text{Ge}_2\text{Sb}_2\text{Te}_5$ are difficult to predict owing to the comparable contributions of both electrons and phonons to heat flow in the hcp phase and the large mismatch of acoustic properties with respect to common metals and dielectrics. Early work identified a strong reduction of TBR with disordered $\text{ZnS}:\text{SiO}_2$ films due to transition to the crystalline phase.²⁹⁰ More recent work measured the relatively large TBR with respect to device electrode materials, and simulated their large impact on the writing current for a vertical cell geometry.^{291,292}

Thermoelectric transport can have a large impact on temperature fields in PCM nanodevices and can strongly influence the write current.²⁹³ For the reset process to the amorphous phase, Peltier heating at the electrode interface and Thomson heating or cooling in the crystalline phase of the chalcogenide material can govern the size of the amorphous region resulting from a given current pulse. Nanoscale electrothermal simulations are complicated by size and temperature dependence of thermoelectric properties of $\text{Ge}_2\text{Sb}_2\text{Te}_5$. Experimental data exhibit differing magnitudes

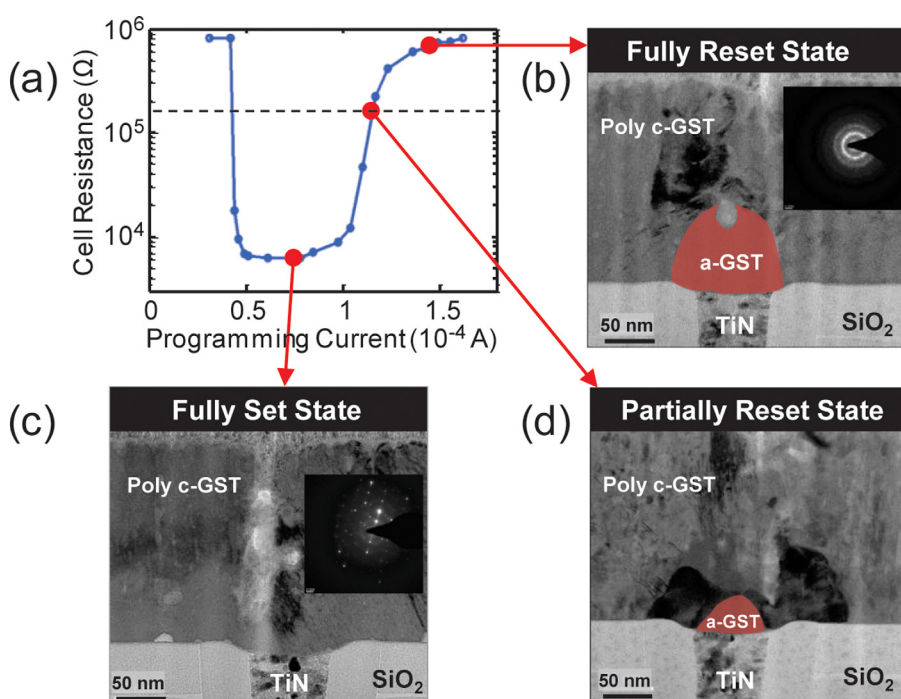


FIG. 11. Transmission electron micrographs, electron diffraction data, and multiphysics prediction of the electrical resistance as a function of programming current for a phase change memory nanocell.^{278,283} (a) A high current generates a peak temperature rise exceeding 600°C and induces melting of a dome-shaped region (b), which is preserved in the amorphous condition through rapid cooling when the current stops. Lower currents recrystallize the amorphous region either fully (c) or partially (d). The evolution of the phase distributions due to an electrical current is governed by nanoscale heat conduction, thermoelectric transport, and crystallization kinetics during timescales near 10 ns.²⁶⁸ Reproduced with permission from Nano Lett. 12, 2179 (2012). Copyright 2012 American Chemical Society.

and temperature derivatives for the Seebeck coefficient as the material is converted from the amorphous to the crystalline phase through annealing.^{280,294} Furthermore, when the material is predominantly crystalline, a strong thickness dependence of the Seebeck coefficient may be linked to the variation of the phase purity with film thickness for a given annealing history

Recent work has also explored the fundamental scaling limits of PCRAM using electrodes made from carbon nanotubes rather than metals.^{295–297} CNTs can have a higher electrical conductivity and can be more reliable than metals down to diameters of only ~ 2 nm; thus, they can form electrodes contacting PCM bit volumes of just tens of cubic nanometers. Figure 12 shows such a device,²⁹⁶ where the CNT is “cut” in the middle, forming a small nanogap (~ 10 to 200 nm). Sputtering ~ 10 nm GST²⁹⁵ fills the nanogap, forming the extremely small PCM bit with two CNT electrodes. Such devices switch with programming currents of ~ 1 μ A, two orders of magnitude lower than the previous state-of-the-art,^{268,296} and programming energy < 100 fJ per bit. Testing hundreds of such structures²⁹⁶ has uncovered trends suggesting that PCRAM could be scaled down to achieve sub-fJ per bit operation in sub-5-nm bits. Thus, the fundamental operating limits of PCMs have not yet been reached, and these limits are set by phase transformations and heating at single nanometer dimensions.

At these scales, the influence of even subtle atomistic changes (e.g., grain boundaries, and stoichiometry) on the thermophysical properties of such materials are unknown, and their contact resistances (electrical and thermal) with the surrounding materials play a very important role. Moreover, the ultimate packing density of such bits in memory arrays is also limited by thermal disturb between adjacent bits,^{298,299} motivating further research into heat flow on sub-10 nm length scales in PCMs, CNTs, and their interfaces and contacts.

Another memory technology for which thermal transport plays a central role is heat-assisted magnetic recording (HAMR), which involves near-field radiation, ballistic thin-film conduction, interface resistance, and nanoscale lubricant transport.³⁰⁰ Near-field radiative heating, intensified and localized through plasmon excitation, must heat a ~ 10 nm spot in the recording medium above its Curie temperature (generally above ~ 500 °C) in ~ 1 ns. The recording medium consists of metal and dielectric nanolayers of thickness 10–40 nm along with the functional magnetic film. The effective thermal conductivities of this stack are highly anisotropic and governed by electron and phonon scattering and energy conversion. One challenge is to optimize the complex multilayer system, considering both photon absorption and transient, anisotropic thermal conduction, such that the lateral dimensions of the hotspot are minimized. Recent HAMR progress includes detailed electromagnetic simulations of the near-field transducer, studies of the interacting thermal and magnetic properties of the recording medium,³⁰¹ approximate analyses of the combined optical and thermal transport in the medium,³⁰² and even studies of the heated lubricant flow in the gap below the head.³⁰³ A systematic characterization of HAMR materials for optical and thermal performance remains important for future work.

2. Nanoscale CMOS and interconnects

Heat conduction in electronics begins with the energy dissipated in nanoscale transistors and interconnects.²⁷⁵ Complementary metal-oxide-semiconductor (CMOS) field-effect transistors (FETs) are the building blocks of modern electronics, typically based on Si technology with features down to ~ 28 nm.³⁰⁴ However, scaling devices to sub-10 nm dimensions will likely involve new materials from groups III-V such as GaAs, InAs, or InSb,³⁰⁵ or group IV such as Ge or C,^{306,307} which have electronic properties (e.g., mobility)

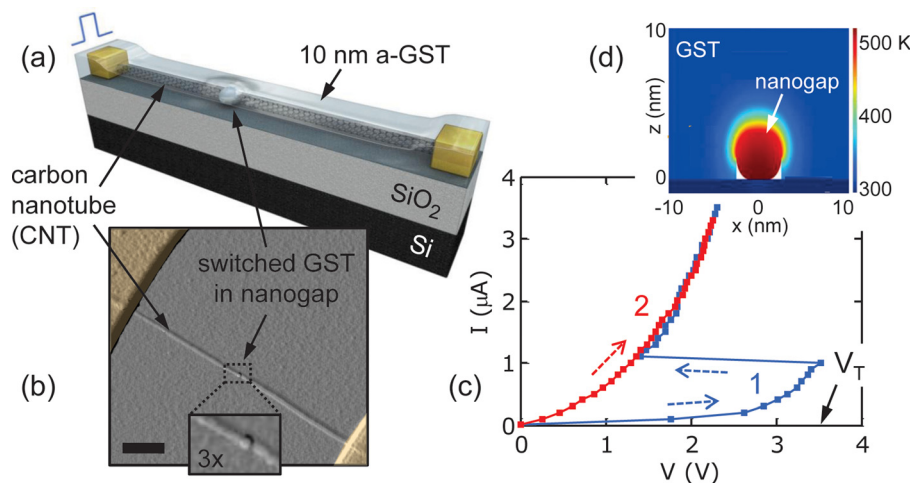


FIG. 12. (a) CNT electrodes have been used to contact the smallest and most energy-efficient phase-change material memory devices to date, see Refs. 295–297. The CNT spans two Pd contacts on an oxidized Si wafer. A nanogap is formed electrically or mechanically with the tip of an atomic force microscope. Amorphous Ge₂Sb₂Te₅ (a-GST) is sputter-deposited to fill the nanogap, and voltage pulses are used to reversibly switch its state. (b) AFM image of experimental device (scale bar is 500 nm); zoom-in shows the top surface of the PCM bit. (c) Experimental current-voltage (I - V) curves reveal switching at ~ 1 μ A, two orders of magnitude lower than previous state-of-the-art PCM devices.²⁹⁷ V_T marks the threshold voltage of the amorphous state. (d) Simulation of nanogap bit cross-section confirms that small dimensions are essential for achieving low-power operation.²⁹⁷ TBR, Seebeck effects, and non-equilibrium transport of both electrons and phonons are crucial for energy-efficient operation at sub-10 nm dimensions and sub-10 ns time scales.

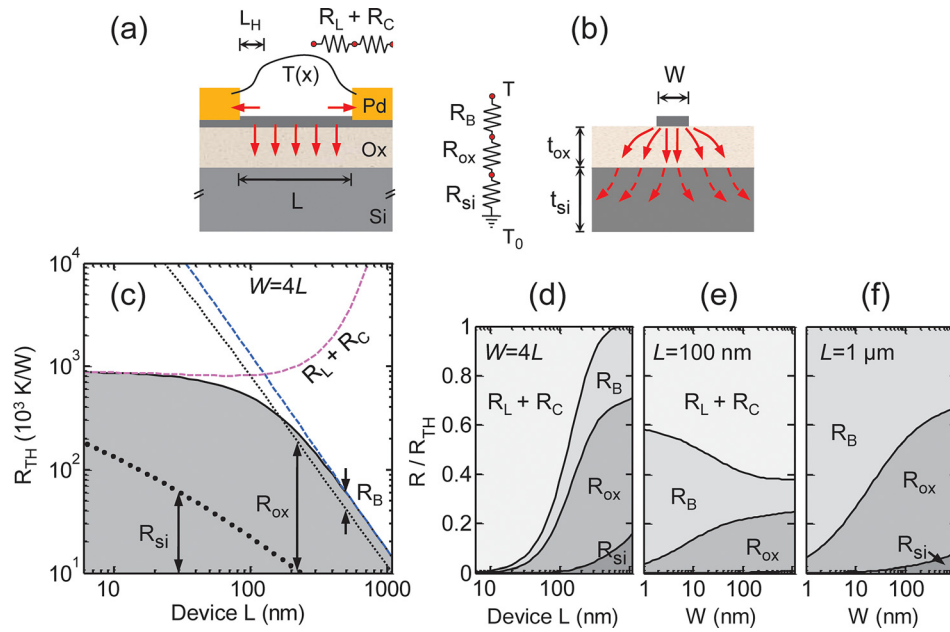


FIG. 13. Competing heat conduction pathways in nanoscale devices. (a) Cross-section along length and (b) along width of a device, here for the case of a single GNR.³¹⁸ Red arrows show heat flow pathways; thermal resistances and other dimensions are as defined in the text. (c) Calculated thermal resistance R_{TH} vs. device length L following models introduced in Refs. 275, 318, and 319. The gray area tracks R_{TH} , while the lines track individual thermal resistance components, as labeled. Trends are consistent with experimental data summarized in Ref. 275. (d) Fractional contribution of individual thermal resistances vs. device length. Assumed width $W = 4L$ and $t_{ox} = 50$ nm. (e) and (f) Fractional contribution of thermal resistances vs. device width for a “short” ($L = 100$ nm) and a “long” device ($1 \mu\text{m}$), respectively. Extrinsic device properties (such as TBR R_B and contact thermal resistance R_C) dominate over intrinsic properties (such as thermal conductivity of channel and substrate) for devices that are shorter and narrower than ~ 100 nm.

superior to Si. Further scaling will also involve new geometries such as multi-gate (FinFET) devices,³⁰⁸ nanowires, or nanotubes.³⁰⁶

In all cases, power dissipation begins with electrons or holes that are accelerated by high electric fields, of the order ~ 10 V/ μm . These “hot” carriers scatter with the lattice, emitting acoustic and optical phonons (OPs).^{309,310} Modern devices operate at voltages near ~ 1 V, strongly emitting OPs which have energies ~ 35 meV in Ge or GaAs, 60 meV in Si, 180 meV in CNTs and graphene.^{309,310} Typical OP lifetimes are of the order ~ 1 ps;^{311,312} however, the thermal time constants of CMOS devices are of the order ~ 10 ns due to the larger volumetric heat capacity.²⁷⁵ It is worth noting that the cooling time of a single carbon nanotube is ~ 0.1 ns,³¹³ likely representing the smallest thermal time constant achievable in a CMOS device. The thermal time constant of a nanoscale device represents how quickly it cools down or heats up after an electrical transient, and such temperature excursions determine both the device leakage and long-term reliability.^{275,314}

Once a nanoscale device reaches thermal steady state, the pathway of heat conduction to the environment is key in determining its temperature rise and ultimately its reliability.³¹⁴ The temperature rise of a modern device can reach $\Delta T > 100$ K above room temperature, being determined by its power input P and the thermal resistance R_{TH} , as $\Delta T = PR_{TH}$. For instance, the thermal resistance of a single device on the surface of a Si wafer is approximately $R_{si} \approx 1/(2\kappa_{Si}D)$, where $D \approx (LW)^{1/2}$, L and W are the length and width.²⁷⁵ However, as devices are scaled down, four additional heat flow considerations must be taken into account. First, heat conduction into contacts or the gate

electrode could play a greater role. Second, novel devices (e.g., FinFET, nanowire, and CNT) contain many interfaces between the semiconductor body where dissipation takes place, and the external heat sink. Thus, the associated thermal boundary resistances must be carefully taken into account. Third, nanoscale devices are smaller than the bulk phonon mean free path, $\Lambda \approx 100$ nm in Si and comparable in most semiconductors. Thus, the thermal conductivity is lower than well-known bulk values, for instance, that of a ~ 20 nm thin silicon-on-insulator (SOI) film is $\kappa_{SOI} \approx 25$ W m⁻¹ K⁻¹ compared to the thermal conductivity of bulk Si, $\kappa_{Si} \approx 150$ W m⁻¹ K⁻¹.³¹⁵ Fourth, modern devices have electrical switching transients of the order 10–100 ps, ranging between those of carrier scattering events (0.1–1 ps) and the thermal relaxation of the lattice (~ 10 ns). Thus, nanoscale devices operate in a complex temporal fashion, where non-equilibrium situations can arise between the electrons, the OPs, and the lattice whose heat capacity is primarily determined by acoustic phonons.³¹⁶

Figure 13 illustrates the various heat conduction mechanisms in transistor-like or interconnect-like nanoscale devices. For simplicity, the figure ignores heat loss through a top gate, which is discussed later in Fig. 14. The key questions answered here concern whether heat dissipation from a modern nanoscale device takes place along the channel, through the contacts, or via the substrate. The various heat flow paths are denoted by thermal resistances R_L along the device, R_C through the contacts, R_{ox} for the supporting oxide (of an SOI-like device), R_{si} for the silicon substrate, and R_B for the channel-oxide thermal boundary resistance. The figure also clarifies the *relative* roles played by intrinsic thermal properties (such as thermal conductivity of the channel and

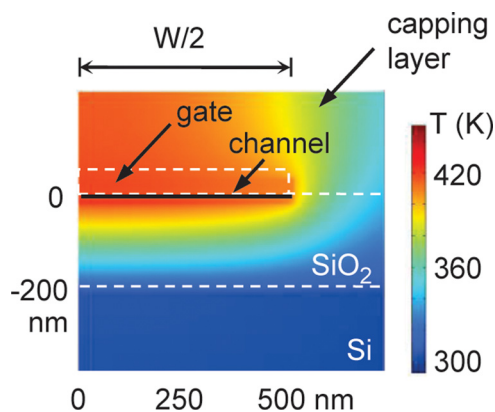


FIG. 14. Finite element simulation of a graphene transistor including its top gate.³²¹ The cross-section along the device width only shows one-half ($W/2$) by symmetry considerations. The simulation finds that heat sinking into the top gate is minimal, in part due to TBR with the two surfaces of the gate-channel dielectric, and in part due to much larger heat spreading to the substrate and contacts. The top gate simply assumes the same temperature as the device channel. However, top gates and any additional capping layers provide thermal capacitance, which significantly extends the thermal time constants of such devices.³²¹ Reproduced with permission from IEEE Electron Device Lett. **34**, 166–168 (2013). Copyright 2013 IEEE.

substrate) versus extrinsic properties (such as the contacts and TBR), and how these change as the device length (L) and width (W) are scaled toward sub-10 nm dimensions.

Numerical calculations in Fig. 13 are based on the models in Refs. 317–319, and correspond to a graphene device on SiO_2 with thickness $t_{ox} = 50$ nm, graphene- SiO_2 boundary resistance $R_B = 2 \times 10^{-8} \text{ m}^2 \text{ K W}^{-1}$, and graphene thermal conductivity $\kappa_g = 150 \text{ W m}^{-1} \text{ K}^{-1}$, consistent with graphene nanoribbons³²⁰ or encased graphene.²⁶¹ The numerical analysis is similar for an ultra-thin SOI device with thickness $t_{SOI} = 5$ nm and thermal conductivity $\kappa_{SOI} = 10 \text{ W m}^{-1} \text{ K}^{-1}$ (same $t\kappa$ product).³¹⁵ Thus, the trends observed in the figures are also typical for other modern and future nanoscale transistors, including ultra-thin body SOI, multi-gate (FinFET), CNT, or nanowire FETs.

Figure 13(a) shows a cross-section of the device along its length, and Fig. 13(b) shows the cross-section across its width. The heat conduction pathways can be both “vertical” into the substrate and “lateral” along the device and into the contacts, as shown by red arrows. Figure 13(c) displays the total thermal resistance R_{TH} , in units of temperature rise per unit input power, of a typical device as a function of channel length L ; in this example, for a device width $W = 4L$. The thermal resistance of “longer” devices is dominated by the vertical heat flow path which includes the TBR (R_B), the thermal resistance of the oxide (R_{ox}), and, to a smaller degree, the Si substrate (R_{Si}). However, the thermal resistance of “shorter” devices (particularly with $L < 100$ nm) becomes limited by heat flow along the channel and into the contacts ($R_L + R_C$). The cross-over of these two regimes occurs at a device length that is approximately three times the thermal healing length (L_H) along the channel.^{318,321} The results of Fig. 13(c) are consistent with experimental data summarized in Fig. 5(a) of Ref. 275.

The fractional contribution of each thermal pathway to the total thermal resistance is shown in Fig. 13(d) as a

function of channel length L . For long-channel devices, the thermal resistance is approximately 30% due to TBR (R_B) and the rest dominated by the thermal resistance of the underlying oxide (R_{ox}) for the combination of parameters used in these estimates. (The relative importance of TBR would be larger if thinner underlying oxides are used.) The contact thermal resistance (R_C) becomes increasingly important in sub-100 nm devices, almost regardless of the thermal properties of the device or substrate.

Figures 13(e) and 13(f) show the fractional contributions of various thermal resistances as a function of device width W , for “short” ($L = 100$ nm) and “long” devices ($L = 1 \mu\text{m}$), respectively. In both cases, we see the increasing relative importance of the TBR between the device and the substrate as device widths are narrowed. However, for longer devices, this TBR component can almost entirely dominate as device widths approach $W \sim 1$ nm, which has been experimentally confirmed for the case of CNTs.^{322,323}

We also examine the role of top gates in heat sinking from transistors in Fig. 14. Finite-element simulations³²¹ suggest that top gates are not significant heat sinks, in part due to the additional TBRs associated with the channel-dielectric and dielectric-gate interfaces, and in part because the main heat conduction pathway remains into the substrate and towards the back-side heat sink. This can be seen as the top gate assumes nearly the same temperature as the channel in Fig. 14. However, top gates and the dielectric capping layers covering them add thermal capacitance near the device channel, thus lengthening the device thermal time constants. For instance, the thermal time constant of the graphene device is approximately 50 ns without a capping layer, and >100 ns with the capping layer shown (here taken as 200 nm of SiO_2).³²¹ These results illustrate that the exact heat dissipation picture is complex and must often be evaluated for the particular, 3-D device designs.^{324–326}

Unlike transistors, nanoscale interconnects are typically metals like Cu with liners or interfaces of TaN, TiN, or W.^{327,328} In these materials, both electrons and phonons can contribute to heat conduction, while the interfaces, grain boundaries (GBs), and line edge roughness (LER) play an increasingly important role as dimensions are reduced. Importantly, both the magnitude of thermal conductivity (κ) and the relative contributions of electrons and phonons ($\kappa = \kappa_e + \kappa_p$) are modified by GB and LER scattering. For instance, the electrical conductivity (σ) of Cu nanowires decreases by $\sim 5\times$ from the bulk value at linewidths of ~ 10 nm.³²⁸

However, data for Pt films and nanowires reveal that σ and κ do not always scale proportionally, as would be predicted by the Wiedemann-Franz Law (WFL) in metals. Platinum films with thickness down to 7.3 nm fabricated using atomic layer deposition (ALD) show a stronger reduction of σ (by $4.5\times$) than of κ ($3\times$) with respect to bulk values.³²⁹ The authors suggested that the excess thermal conductivity in the Pt films, above that predicted by the WFL, may be due to phonon transport. However, electrochemically deposited Pt nanowires with diameter ~ 100 nm reveal a weaker decrease of σ (by $2.5\times$) than of κ ($3.4\times$),³³⁰ which suggests that the WFL overpredicts the electron contribution

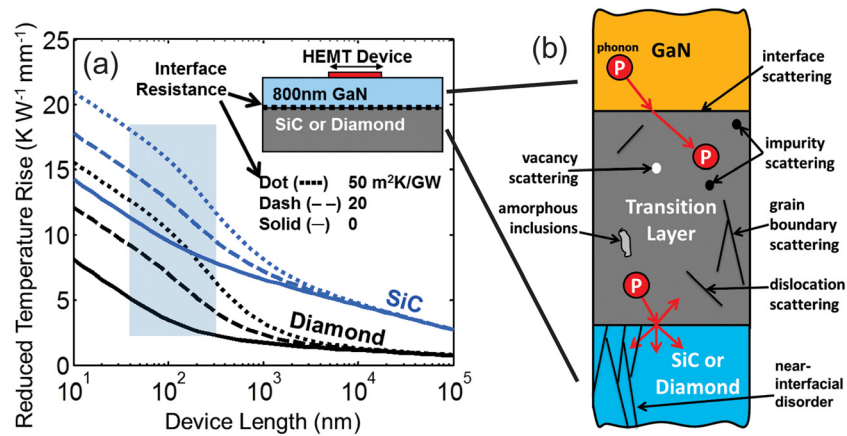


FIG. 15. Physical mechanisms, simulated impact, and experimental data for the room temperature interface thermal resistance in composite substrates involving diamond and GaN, which is a leading material combination for radar amplifiers. (a) The blue box indicates the range of simulations corresponding to contemporary gate lengths for HEMTs and shows that an effective interface resistance of $50 \text{ m}^2 \text{K/GW}$ can fully eliminate the benefits of using diamond relative to silicon carbide. The SiC or diamond substrate is assumed semi-infinite, which yields little error for heater widths less than $10 \mu\text{m}$. (b) Phonon scattering on crystal imperfections depicted at top right can be particularly important in the AlN transition layers common for GaN grown using MOCVD on SiC. For GaN-diamond composites, the best transition layers are a subject of vigorous research.

to the thermal conductivity. This result is surprising because the WFL is generally assumed to work well for boundary scattering owing to the elastic nature of this process. These studies suggest that electron transport at nanoscales in metals is governed by subtle differences in sample and interface characteristics. Similar observations have been made by recent thermal measurements of graphene nanoribbon (GNR) interconnects,³²⁰ where simulations have also pointed out that LER scattering is likely to affect phonons more strongly than electron conduction in narrow GNRs with zigzag edges.³³¹ Much research remains to be carried out to understand the relative contributions to heat conduction of electrons and phonons at sub-10 nm dimensions.

3. Composite substrates for power electronics

Nanoscale heat transfer can be particularly important in semiconductor devices that deliver or modulate power. Examples include semiconductor lasers, power field-effect switches, and radar amplifiers, which can have power densities exceeding 1 kW/cm^2 . Improvements have been particularly rapid for quantum cascade lasers (QCL), which achieve efficient, tunable mid-infrared wavelength emission through quantum confinement in GaInAs/AlInAs multilayers.^{332,333}

Composite substrates aim to bring a higher thermal conductivity heat spreader, such as SiC or diamond, within micrometers of active semiconducting regions. This is generally achieved by bonding or directly growing the functional semiconductor onto a high thermal conductivity substrate, such as CVD diamond. Eutectic bonding of a QCL with a diamond submount yielded high-quality diamond about $5 \mu\text{m}$ from the active epitaxial region.³³⁴ The lower bound thermal resistance can be approached, however, only by thinning or eliminating the attachment film.³³⁵ Progress on such composite substrates has been particularly rapid for high-electron-mobility transistors (HEMT), for which the substrate thermal resistance was reduced through GaN/AlGaIn heterostructure growth on SiC³³⁶ or bonding to diamond.^{274,337}

Figure 15 and Table II summarize the impact of the thermal resistances in GaN composite substrates on their relative heat spreading capabilities, as well as the phonon scattering mechanisms that contribute to the thermal resistance. To significantly outperform GaN-SiC, the GaN-diamond interface resistance must approach the DMM predictions under the Debye approximation, which is near $1 \text{ m}^2 \text{K GW}^{-1}$. While the Debye approach yields some error due to phonon dispersion, the challenge for materials science is clear: strongly reduce defect concentrations within nanometers of a complex interface.

The challenge for the nanoscale thermal transport community is to measure and identify the specific microstructural causes for resistances at interfaces roughly a micrometer below the surface. The effective interface resistance sums the components at the discrete material interfaces with the effects of defect scattering in the two primary substrate materials and the transition film. Diamond composite substrate fabrication procedures include direct growth of the semiconductor on diamond, in some cases with a transitional layer, and growth of the diamond on the semiconductor or a very thin adhesion film. There has been some progress on relating the concentrations of point defects and dislocation segments in the transition region to metal-organic chemical vapor deposition (MOCVD) growth conditions of AlN/GaN on SiC.³³⁸ Future work for diamond composites should relate concentrations of

TABLE II. Summary of the minimum thermal resistances observed in several studies of composite substrates and comparisons to the prediction of the DMM.

	Lowest data ($\text{m}^2 \text{K/GW}$)	DMM prediction
GaN-diamond	<10 (Ref. 274) 15–30 (Ref. 337)	3.0
GaN-SiC	3–5 (Ref. 336)	0.9
Diamond-Si	20–30 (Ref. 340)	0.2

specific types of defects and impurities to the thermal resistance. The optimal fabrication procedures for GaN-diamond, i.e., those that yield the lowest defect concentrations in the semiconductor and the lowest thermal resistance at its interface with diamond, remain undetermined. For the case of diamond growth on the functional semiconductor the transition film, defects in the diamond can impair heat spreading. There has been progress on measuring and modeling the impact of phonon scattering on defects near the deposition interface with silicon^{339,340} and optimization of nucleation parameters to minimize the effective interface thermal resistance.³⁴¹

IV. THERMAL CONDUCTIVITY OF NANOSTRUCTURED MATERIALS

In this section, we review progress in nanostructuring materials for low thermal conductivity applications, and materials having low thermal conductivity as a result of their intrinsic atomic structure and interatomic bonding. The former includes nano-grain sized materials as well as those doped to create point defect scattering and second phases by thermal treatment. The latter includes crystal structures that constitute natural superlattices with superlattice periodicities of the order of nanometers.

In addition to their intrinsic scientific interest, materials having low thermal conductivity are important for several technologies, and for understanding larger scientific issues. For instance, there is a world-wide search for oxides with lower thermal conductivity at high temperatures for use as thermal barriers for gas turbines;³⁴² the thermal conductivity of uranium dioxide nuclear fuel is crucial in reactor performance, and it evolves continuously in the reactor as the chemistry and microstructure evolves.³⁴³ In identifying oxide materials with unusually low thermal conductivities at high temperatures as well as predicting the lowest attainable thermal conductivity of a compound under extreme burn-up, guidance comes from two closely related models based on classic kinetic theory. The first, proposed by Kittel in 1949 (Ref. 344), pertains to amorphous solids and argues that the length scale associated with the atomic disorder determines the mean free path for phonon scattering. The other, the Cahill and Pohl model,⁸⁶ describes the minimum thermal conductivity of a crystal when the mean-free path approaches the inter-atomic spacing and is usually described in terms of the longitudinal and transverse sound velocities, and, respectively in the high temperature limit

$$\kappa_{\min} = 0.40k_B \left(\frac{N}{\Omega_0} \right)^{2/3} (v_{L,s} + 2v_{T,s}). \quad (28)$$

Here, Ω_0 is the unit cell volume and N is the number of atoms per unit cell. At the minimum thermal conductivity, one can view heat transport in the spirit of an Einstein oscillator model as a process involving the successive hopping diffusion of energy from one atom to a nearest neighbor. This limiting behavior can occur at high temperatures even in crystalline materials as well as in glasses.

For crystalline materials below temperatures at which the minimum thermal conductivity, $\kappa_{\min}$, is reached, the general

principles underlying the effects of many microstructural effects on phonon-mediated thermal transport were largely worked out in the 1950s and 1960s by Klemens,^{345,346} Callaway,^{347,348} and others.¹⁸⁰ These analyses led to expressions for the thermal conductivity in terms of the phonon relaxation times for different phonon scattering processes integrated over the phonon energy distribution. Assuming a Debye density of states distribution, the thermal conductivity described by Eq. (26) can be written as

$$\kappa = \frac{1}{2\pi^2 v_D} \int_0^{\omega_D} C_v(\omega) \tau(\omega) \omega^2 d\omega, \quad (29)$$

where ω_D is the Debye frequency, v_D is the Debye velocity of sound, and $C_v(\omega)$ is the frequency dependent heat capacity. In practice, the mean-free-path for phonon scattering cannot be smaller than approximately half the phonon wavelength and, in turn, smaller than the inter-atomic spacing, as pointed out by Roufosse and Klemens.^{349,350} This requires the integral to be split into two parts

$$\kappa = \frac{1}{2\pi^2 v} \int_0^{\omega_i} C_v(\omega) \tau(\omega) \omega^2 d\omega + \frac{1}{2\pi^2 v} \int_{\omega_i}^{\omega_D} C_v(\omega) \tau(\omega) \omega^2 d\omega, \quad (30)$$

where the second term is, in essence, the same temperature-independent minimum conductivity described in greater detail by Cahill and Pohl.⁸⁶ In materials in which multiple phonon scattering processes occur, the contributions from each process are added inversely through Matthiessen's rule^{345,351,352}

$$\frac{1}{\tau} = \frac{1}{\tau_u} + \frac{1}{\tau_d} + \frac{1}{\tau_b} + \dots \quad (31)$$

Klemens³⁴⁵ derived specific scattering times associated with phonon-phonon Umklapp scattering τ_u , phonon scattering from point defects τ_d , dislocation strain fields τ_s , and grain boundaries τ_{GB}

$$\frac{1}{\tau_u} = A\omega^2 T^\xi \exp(-B/T), \quad (32)$$

$$\frac{1}{\tau_d} = V_{ac} \left(\frac{m_i - m_{ave}}{m_{ave}} \right)^2 \frac{\omega^4}{4\pi v^3}, \quad (33)$$

$$\frac{1}{\tau_s} = \frac{2^{3/2}}{3^{3/2}} N_d b^2 \gamma^2, \quad (34)$$

$$\frac{1}{\tau_{GB}} = \frac{d}{v_s}. \quad (35)$$

Equation (32) is a commonly used approximation of the anharmonic scattering rates that are discussed in detail in Sec. III A, in particular, see Eq. (18) and Fig. 5. Equation (6) is an approximation of Eq. (21) that assumes a Debye density of states.

There are several important results pertinent to nanostructured materials that follow directly from the Klemens-Callaway model³⁴⁸ in Eq. (30) and the form of the relaxation

times. First, in the absence of any defects and at intermediate temperatures, below 2 to 5 times the Debye temperature, T_D , the thermal conductivity decreases approximately as $1/T$ and asymptotes to a temperature independent value given by $\kappa_{\min}$. Microstructural defects, including point defects, anti-site defects, and grain boundaries, all decrease the scattering time with contributions that are independent of temperature. As the thermal conductivity due to anharmonic phonon scattering decreases with increasing temperature, all forms of microstructural defects have a greater apparent effect in lowering thermal conductivity at lower temperatures than at the higher temperatures, decreasing the observable temperature dependence. In the extreme, when the minimum conductivity is reached, they can result in a temperature-independent conductivity over a very large temperature range. We will return to this point later in discussing grain size effects and high-defect concentrations.

A. Doping and point defects

As Eq. (33) shows, the effect of doping depends on the square of the normalized mass mismatch of the dopant and the host lattice ions. This effect has been widely investigated for low concentrations of dopants for many years and is clearly manifested in a number of materials systems. The thermal conductivity of natural diamond (containing 1.1% of ^{13}C) at room temperature is $\approx 2000 \text{ W m}^{-1} \text{ K}^{-1}$. This isotopic concentration, and high normalized mass mismatch, has a large impact on the thermal conductivity: diamond, isotopically enhanced such that it contains only 0.001% ^{13}C has a considerably higher thermal conductivity.³⁵³ Isotopic effects have little effect for higher mass systems. For example, natural uranium is an isotopic mixture of 99.3% ^{238}U and 0.7% ^{235}U ; atomic-level simulations of isotopic effects in uranium dioxide, UO_2 , fuel show that this leads to only $\approx 4\%$ decrease in the thermal conductivity.³⁵⁴ Turning to off-stoichiometry, UO_2 can easily accommodate oxygen interstitials compensated by U^{5+} ions. As Fig. 16 shows, such off-stoichiometry can result in a significant reduction in the thermal conductivity. Very similar quantitative behavior is observed in the $\text{Gd}_{8+x}\text{Ca}_{2+y}(\text{SiO}_4)_6\text{O}_{2+3x/2+y}$ apatite system where ranges of solid solution with off-stoichiometry accommodated by either oxygen vacancies, oxygen interstitials or a combination can be produced by varying the Gd:Ca ratio.³⁵⁵

At higher doping and solute concentrations, the details of phonon scattering are more complex. A case in point is the decrease in thermal conductivity of yttria-stabilized zirconia (YSZ) with doping. Without any yttria, zirconia exhibits the typical $1/T$ dependence of many other oxides; however, as the yttria doping concentration is increased, its conductivity decreases and the temperature dependence becomes weaker. The yttrium ions substituting on the Zr cation sub-lattice do not themselves introduce appreciable point defect scattering because their atomic mass is similar to that of Zr; however, the oxygen vacancies they create do, just as they do in UO_2 . At 7–8 wt. % yttria, 3YSZ, which is the prototypical material for thermal barrier applications, has a tetragonal crystal structure and its conductivity is almost temperature independent. For fully dense 3YSZ at 1373 K, a

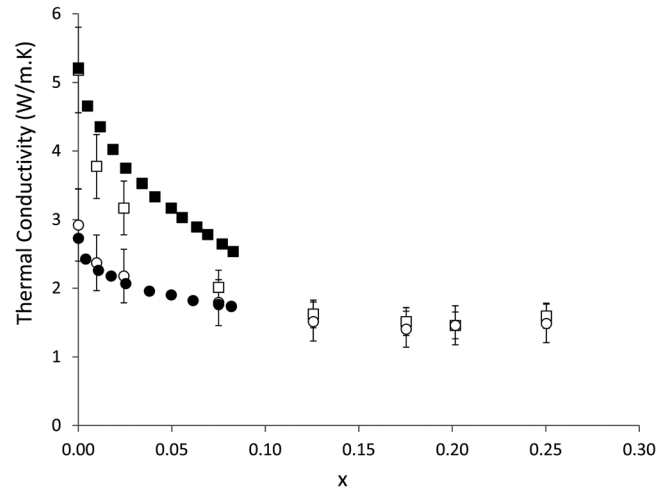


FIG. 16. Thermal conductivity of UO_{2+x} from simulation at 800 K (open circles) and 1600 K (open squares)³⁵⁴ compared to experiment trends³⁶² at 773 K (solid circles) and 1673 K (solid squares). Reproduced with permission from J. Am. Ceram. Soc. **92**, 805–856 (2009). Copyright 2009 John Wiley and Sons.

typical thermal barrier coating (TBC) operating temperature, the thermal conductivity is $2.5 \text{ W m}^{-1} \text{ K}^{-1}$.³⁵⁶ Deliberately introduced porosity further reduces this to $1.5\text{--}2.0 \text{ W m}^{-1} \text{ K}^{-1}$, depending on the deposition method used.^{357,358} Also, stabilization of the tetragonal phase by oxygen vacancies has the highly desirable effect of imparting high fracture toughness as a result of stress-induced ferroelasticity.³⁵⁹ Analysis of the phonon properties on both UO_2 (Ref. 354) and YSZ (Ref. 360) showed that chemical disordering through the presence of high densities of point defects means that for most vibrational modes, it is not possible to clearly define a polarization or a wave-vector, and hence consider simple Rayleigh scattering. Rather, from a heat transport perspective, the thermal-transport properties systems behave in a similar manner to structurally amorphous materials.³⁶¹

B. Grain boundaries

For many materials, the simplest approach to nanostructuring is to reduce grain size. Grain boundaries act as obstacles to the movement of phonons through a material by scattering phonons (Eq. (35)), and lower the thermal conductivity by decreasing the time between phonon scattering events. A simple two-state model for the overall thermal conductivity of a polycrystal in terms of the thermal conductivity of the bulk and of a grain-boundary region leads to

$$\kappa(T) = \frac{d\kappa_i(T)}{d + R_K\kappa_i(T)}, \quad (36)$$

where R_K is the interfacial (Kapitza) resistance, κ_i is the thermal conductivity of the bulk and d is the grain size. Figure 17 shows the effects of grain size for SrTiO_3 . There is a substantial reduction in the thermal conductivity as the grain size decreases, particularly at lower temperatures, at which the anharmonic phonon-phonon scattering effects are less significant, consistent with the expectations of the Debye model discussed above.³⁶⁶ Also, as expected, the difference

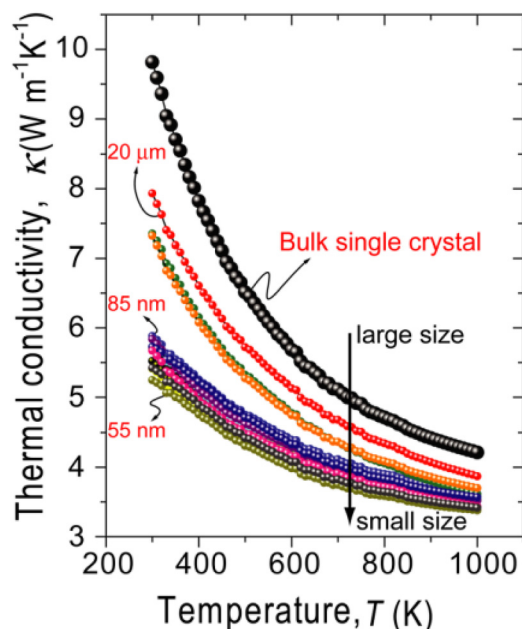


FIG. 17. Temperature dependence of the thermal conductivity of SrTiO_3 from bulk single crystal down to 55 nm grain sizes shows a strong reduction due to increased contribution from grain boundary scattering. Reproduced with permission from Appl. Phys. Express 3, 03101 (2010). Copyright 2010 The Japan Society of Applied Physics.

between the conductivities at high temperatures becomes less marked for larger grain sizes. Similar results have been obtained with yttria-stabilized zirconia.³⁶⁸ This is an important result since it means that grain size manipulation for grain sizes in excess of $\sim 1\mu\text{m}$ is a much less effective strategy for decreasing thermal conductivity for high temperature applications than for low-temperature applications.

While grain growth at high temperatures is familiar, decreases in grain size can also occur during the operational lifetime of a material. For example, microstructural evolution of UO_2 during in-reactor burn-up involves the development of very small grain sizes (averaging $\sim 130\text{nm}$, but with a significant fraction less than 100nm)³⁶³ and high porosity in the outer regions of the fuel pellet, i.e., the “rim effect.”^{343,364} The combination of high GB area, segregation of fission gases, and high porosity in this region leads to a significant decrease in the thermal conductivity.³⁶⁵

The interfacial thermal (Kapitza) conductance can be determined from data on the grain size and temperature dependence of thermal conductivity. Figure 18 is an example obtained from low temperature measurements from cubic zirconia³⁵⁶ and high temperature data for tetragonal 7YSZ zirconia.³⁶⁷ Up to about the Debye temperature (475 K), the Kapitza conductance increases with temperature, whereas above the Debye temperature, it is temperature independent. This temperature independence is largely due to the mean free path of the phonons approaching inter-atomic dimensions near the Debye temperature, with the result that the grain boundaries can only scatter the long wave-length phonons.

The interfacial thermal resistance is the result of the interactions of an ensemble of phonons with the GB. At the simplest level, the effect of a grain boundary can be

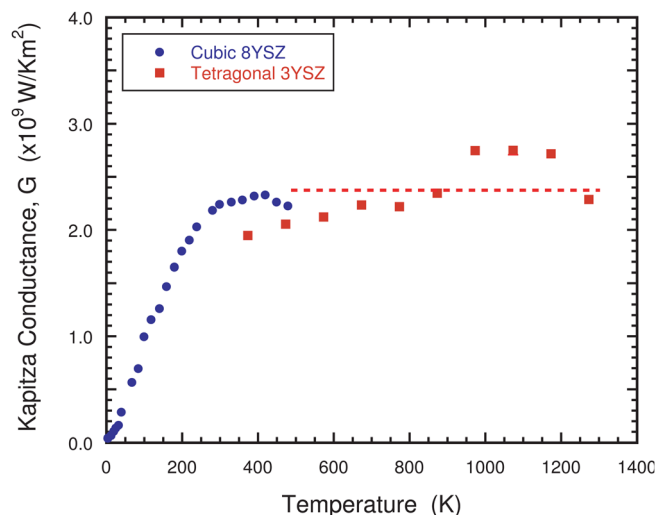


FIG. 18. The temperature dependence of the Kapitza conductance in two closely related yttria-stabilized zirconia materials.^{356,367} The error bars of the data from the tetragonal zirconia are considerably larger because of the small variation in thermal conductivity above the Debye temperature.

characterized in terms of the phonon transmittance, t , the fraction of phonon energy that passes through the interface. One model for this transmittance based on phonon hopping is that it is related to the Kapitza conductance by³⁶⁸

$$G_K = \frac{1}{R_K} = \frac{0.735k_B}{4(\Omega_0/N)}vt. \quad (37)$$

Here, v is the phonon velocity, t is the transmittance, and the other symbols have the same meaning as in Eq. (28). The transmittance for YSZ grain boundaries deduced from Fig. 18 is 0.89; this is of course, averaged over all of the grain boundaries in the polycrystal and over all of the phonon branches and frequencies.

To dissect into individual phonon-grain boundary interactions, it has proved instructive to characterize the interactions of packets of phonons from a single branch and a narrow frequency range with specific interfaces. As is well-known, the standard models in which to interpret the phonon/grain boundary interactions are the DMM and the AMM.⁴³ In the DMM, it is assumed that all of the energy in a phonon is trapped by the grain boundary, and then transmitted or reflected with probability proportional to the available density of states on the two sides of the grain boundary. In the AMM, the transmission coefficient is determined from the acoustic impedances, $Z = \rho v$, of the crystals on the two sides of the GB. Thus, for example, for a twist GB in electronic insulators and semiconductors (e.g., Si) the DMM predicts a transmission coefficient of 0.5, while the AMM predicts a transmission coefficient of 1.0. Detailed analysis of the transmission coefficient of transverse acoustic phonons through a $\Sigma 29$ twist GB in silicon showed that the AMM is valid for long wavelength phonons, whereas the DMM is valid for short wavelength phonons; the transmission coefficient shows a more complex behavior at intermediate wavelengths.³⁸

There thus remains considerable need for a more comprehensive understanding of phonon transmission through

interfaces and their relationship to the crystallography and orientation relationships. This is especially so for non-planar boundaries with high average curvature as they dominate in nano-crystalline materials. Furthermore, more comprehensive modeling is required to gain a more complete understanding of the thermal resistance across interfaces between dissimilar phases having different crystal structures and different mean atomic masses. This would be extremely useful in guiding the selection of two and three phase materials for the next-generation thermal barrier coatings.

C. Anisotropic crystals and natural superlattices

The engineering of artificial superlattices allows significant reduction in the thermal conductivity. It also introduces significant anisotropy between the cross- and parallel-plane thermal conductivities of such structures. It thus seems reasonable to expect that crystal structures having a strong structural anisotropy might also display anisotropic thermal transport. This has been found to be the case for several natural superlattices and is well illustrated by perovskite-based compounds. Natural superlattices are compounds whose unit cell consists of alternating crystallographic blocks having different atomic structure. For instance, bismuth titanate (BiT , $\text{Bi}_4\text{Ti}_3\text{O}_{12}$), the prototypical Aurivillius phase, consists of $(\text{Bi}_2\text{Ti}_3\text{O}_9)^{2-}$ perovskite blocks separated by $(\text{Bi}_2\text{O}_2)^{2+}$ layers.³⁶⁹

Similarly, the prototypical Ruddlesden-Popper (RP) phase consists of one or more SrTiO_3 perovskite blocks separated by a SrO rocksalt layer.³⁷⁰ Simulations³⁷¹ for various values of n , the number of perovskite layers between each SrO layer, show that for large values of n , the system responds as a classical interfacial system and the thermal conductivity normal to the modulation directions decreases linearly with density of the interfaces. For the lowest values of n , the system behaves as a monolithic single crystal with a larger unit cell. The thermal conductivity of rocksalt SrO is higher than that of perovskite-structured SrTiO_3 . Thus, for these low values of n , the thermal conductivity decreases slightly with increasing n . As a result, there is a minimum in the thermal conductivity at a critical value.

Figure 19 summarizes the conductivity of a range of recently discovered low thermal conductivity crystalline oxides. For comparison, the conductivity of fully dense 8YSZ, the current material of choice for thermal barrier coatings, is shown. In addition to having low thermal conductivities, it is notable that with the exception of LaPO_4 , all of the conductivities are almost temperature-independent above room temperature. This is particularly striking for $\text{Bi}_4\text{Ti}_3\text{O}_{12}$ where the conductivity perpendicular to the superlattice plane is almost constant from 150 K up to 1273 K. The figure includes materials, such as (Tm, Eu) co-doped YSZ, $(\text{Zr,Hf})\text{Y}_{0.13}\text{O}_2$,³⁷² and $(\text{Zr}_{0.6}\text{Y}_{0.2}\text{Ta}_{0.2})\text{O}_2$,³⁷³ whose conductivity is attributed to reductions in thermal conductivity from the fluorite-derived 8YSZ due to doping. The case of $(\text{Zr}_{0.6}\text{Y}_{0.2}\text{Ta}_{0.2})\text{O}_2$ is particularly instructive, since by doping with equal concentrations of a trivalent and pentavalent cations, the crystal structure does not contain oxygen vacancies. (As discussed above, the low thermal conductivity of YSZ

had been attributed to the presence of oxygen vacancies). LaPO_4 and $\beta\text{-La}_2\text{Mo}_2\text{O}_9$ are examples of crystal structures with internal distortions. The compound $\beta\text{-La}_2\text{Mo}_2\text{O}_9$ is unusual because it has an exceptionally low elastic modulus, half that of fused silica and almost one-tenth of sapphire has unusual seven-fold coordination for the Mo ion and is also a fast ion oxygen conductor.

The data in Fig. 19 illustrate that materials can manifest low intrinsic thermal conductivity for a variety of reasons and that the well-established method of introducing a high concentration of point defects, by alloying or altering stoichiometry, or by refining the grain size, can also be used to decrease thermal conductivity. The strategy of selecting a refractory, high mean atomic mass compound and then alloying with cations of different atomic mass to increase phonon scattering has proven to be a viable approach to lowering thermal conductivity in many oxides.^{374,375} Solute alloying on the A and B cation sites of the pyrochlore oxides has been explored in considerable detail.^{374,376–380} Of these alloyed oxide systems, $\text{Sm}_2\text{Zr}_2\text{O}_7$ with Yb substitution is especially interesting as there is a drop in conductivity associated with possible lattice softening at the composition where the ordered pyrochlore transitions to the disordered fluorite structure.³⁸¹ There is also evidence in the Yb^{3+} alloyed $\text{La}_2\text{Zr}_2\text{O}_7$ solid solution that the low thermal conductivity can be attributed to the Yb^{3+} behaving as a “rattler.”³⁷⁷

Evidently, defect-free oxide crystal structures that are natural superlattices can have low thermal conductivity perpendicular to the stacking plane as shown by both experiment and simulation. The origin of the lower conductivity in terms of the relative contribution from phonon scattering and zone folding as well as from possibly weaker bonding perpendicular to the stacking plane represented by a different Gruneisen parameter remains to be established. In addition, the increased complexity and number of atoms in the unit cell of some of the oxides, such as $\text{Gd}_{8+x}\text{Ca}_{2+y}(\text{SiO}_4)_6\text{O}_{2+3x/2+y}$ apatites³⁷³ and $\beta\text{-La}_2\text{Mo}_2\text{O}_9$,³⁷² for instance, are consistent

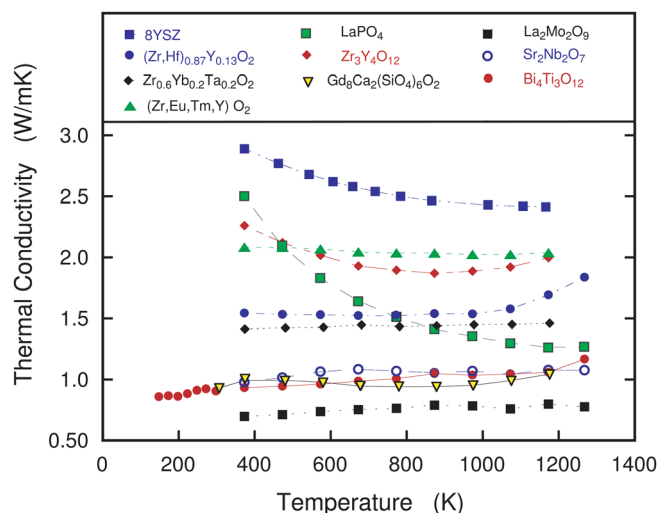


FIG. 19. The thermal conductivity of a variety of complex oxides, all exhibiting little or no variation with temperature. Although all crystalline, the reasons for their low thermal conductivity vary as discussed in the text. Each material was fully dense, so there is no contribution from porosity. Data compiled from Refs. 355, 372, 373, 382, and 383.

with the expectation that more of the thermal energy is carried by optical modes and contribute less to the thermal conductivity. To make further headway in understanding thermal conductivity in complex oxides such as these, more simulations and detailed inelastic neutron scattering studies of the phonon dispersion curves are required.

V. METROLOGY AND PROCESSING TOOLS

A. Time-domain thermoreflectance

1. Advances

During the past decade, TDTR has emerged as a powerful, general-purpose tool for the measurement of the thermal transport properties of materials.^{384,385} TDTR is applicable to materials with a wide range of properties and sample geometries. TDTR has been applied across the entire range of thermal conductivity from diamond and high thermal conductivity metals^{386,387} to the ultralow thermal conductivity of disordered layered crystals⁸⁵ and fullerene derivatives.³⁸⁸ Essentially the same method can be applied to bulk materials, thin layers, and individual interfaces with both high (e.g., TiN/MgO and Al/Cu) and low (e.g., Bi/H-terminated diamond) thermal conductance.^{29,71,74} High-throughput mapping³⁸⁹ of thermal conductivity as a function of position is enabled by the micron-size focus of the laser spots and high signal-to-noise. Applications of thermal conductivity mapping include the analysis of metallurgical phase diagrams,³⁹⁰ and characterization of thermal barriers³⁹¹ and coatings for nuclear fuels.³⁹² Since TDTR is an optical, non-contact method, TDTR can be directly applied to samples in optical cryostats or high temperature microscope stages; and samples exposed to other extreme conditions such as the high pressure environment of a diamond anvil cell.^{98,393–395}

More than 25 years ago, the research groups of Maris³⁹⁶ and Eesley³⁹⁷ independently demonstrated ultrafast optical measurements of thermal transport properties using thermoreflectance as the detection scheme. The approach of Maris and co-workers—the deposition of a metal film transducer on the sample of interest, heating of the transducer by a pump optical pulse, and measurement of the changes in reflectivity of the transducer for thermometry—is essentially the same as what we use today. The key advance in the past decade has been in a method for data analysis that starts from an analytical solution to the diffusion equation in cylindrical coordinates³⁸⁴ and includes the effects of so-called “pulse accumulation,” i.e., the fact that the signals are measured in a TDTR are not only a function of the pump-probe delay time but are also a function of the modulation frequency and the laser repetition rate.^{384,398} The use of single objective lens and integration of the pump and probe beams into an optical imaging system have made the alignment and focusing of the pump and probe beams fast and convenient.^{1,385}

The determination of thermal transport properties by TDTR is typically accomplished by adjusting free parameters in a thermal model³⁸⁴ to obtain the best fit between the predicted and measured thermal response of the sample. The free parameters are the unknown thermal transport

properties, e.g., the thermal conductivity of a thin layer or the thermal conductance of an interface. Thus, TDTR data analysis proceeds in much the same manner as the analysis of ellipsometry data where reflectivity data as a function of wavelength and polarization is used to measure optical constants and layer thicknesses; or the manner in which x-ray reflectivity data as a function of scattering angle are analyzed to determine mass densities and layer thicknesses.

Consider a typical measurement of the thermal conductivity of a material. The sample is coated with an 80 nm film of Al. The unknowns in the thermal model are the thermal conductivity of the sample κ and the thermal conductance G of the Al/sample interface G . The other parameters in the thermal model are independently measured or taken from literature values. The thickness of the Al film h is measured by picosecond acoustics; the heat capacity C of the Al is known from literature values; and the thermal conductivity of the Al film can be estimated based on the in-plane electrical conductivity and the Wiedemann-Franz law. The rms value of the pump and probe beam radius is measured by a spatial correlating the pump and probe or by measuring the transmission as a knife edge is translated through the focused pump and probe. The heat capacity of the sample is usually known from literature values or can be estimated based on data for related materials.

2. Limitations

TDTR requires that the surface of the sample is smooth enough so that any undesired modulation of diffuse scattering by thermoelastic effects does not produce a significant contribution to the desired signal coming from the change in specular reflectivity produced by the thermoreflectance of the metal film transducer. The in-plane length-scale or correlation length of the surface morphology also plays a role in these effects but an approximate rule-of-thumb is that the rms surface roughness should be <15 nm. Measurement errors created by modulation of diffuse light scattering are clearly observed in experiments on rough samples but our understanding of the mechanisms are qualitative at best and have not been systematically studied.

TDTR is mostly sensitivity to the thermal conductivity in the through-thickness direction, i.e., heat transport in the direction anti-parallel to the surface normal. For thermally anisotropic materials such as superlattices, textured polycrystalline films, and anisotropic crystals, knowledge of the in-plane thermal conductivity is also desired. In special cases, e.g., highly anisotropic materials³⁹⁸ or a relatively high thermal conductivity thin layer on a low thermal conductivity substrate, the in-plane thermal conductivity can be accurately measured using tightly focused laser beams and an analysis of how the TDTR signal varies as the pump beam is translated relative to the probe.²¹⁸ The development of methods to accurately determine the in-plane thermal conductivity of thin layers is an active area of research.^{218,399}

Applicability of TDTR at low temperatures, <30 K is limited because the small heat capacity of the metal film transducer and sample requires a greatly reduced average laser power to keep the temperature excursions created by

the pump optical pulses below 10% the absolute temperature. At low temperatures, this constraint means that the maximum laser power that can be used in the experiment scales at the $\propto TC$, where T is absolute temperature and C is the heat capacity per unit volume of the metal film transducer. This scaling places a severe constraint on measurements at low temperatures. (High temperature measurements, on the other hand, do not present any fundamental difficulties. The technical challenges in high temperature measurements are the chemical and physical stabilities of the metal film transducer, measurement of the sample temperature, and mitigating heating of the microscope objective by air convection and thermal radiation from the hot sample environment.)

A key concept in understanding a TDTR measurement is the penetration depth δ of thermal waves in the sample at the modulation frequency f , $\delta = \sqrt{D/(\pi f)}$, where D is the thermal diffusivity, $D = \kappa/C$, where C is the heat capacity per unit volume.⁴⁰⁰ For the measurement of a thin film of thickness h_f deposited on a substrate, we describe the layer as “thermally thin” if $h_f < \delta$ and “thermally thick” if $h_f > \delta$. For a thermally-thin sample, a possible limitation in the measurement is the fact that thermal conductance of the film/substrate interface cannot usually be separated⁴⁰⁰ from the thermal conductance of the film, κ/h_f . For a thermally thick film, the measurement is equally sensitive to both κ and C so any uncertainty in the estimation of C propagates linearly⁴⁰⁰ into an uncertainty in κ .

3. Implementation

The most common implementation of TDTR uses a Ti:sapphire laser oscillator as the light source. The laser oscillator produces a train of sub-picosecond optical pulses at a high repetition rate, typically 80 MHz. The output of the laser is split into two optical paths, a “pump” path and a “probe” path, and the pump and probe are focused on the sample by a single microscope objective.^{1,385} The pump path includes an optical delay line and electro-optic modulator capable of chopping the pump beam at frequencies beyond 10 MHz. The probe path typically includes a audio-frequency mechanical chopper to facilitate the removal of coherent background signals. The term “thermoreflectance” refers to the fact that the probe measures changes in the temperature of the sample through the dependence of the optical reflectivity R of a metal film transducer on temperature, T ; dR/dT is the coefficient of thermoreflectance.^{401,402} An Al thin film has an unusually large thermoreflectance at probe wavelengths generated by Ti:sapphire lasers, $dR/dT \approx 10^{-4} \text{ K}^{-1}$ near room temperature at wavelengths near 800 nm. This fact, combined with the strong sensitivity of Al reflectivity to strain, makes Al ideally suited for both TDTR and picosecond acoustics.

A typical temperature excursion from the metal film is on the order of 1 K and therefore the signal (the change in reflectivity of the probe) is $\Delta R \sim 10^{-4}$. This signal is well above the noise floor of laser oscillators (the noise of Ti:sapphire laser oscillators is typically $<10^{-6}/\sqrt{\text{Hz}}$ at frequencies $>1 \text{ MHz}$), but care has to be taken to suppress light from the modulated pump beam from reaching the detector.

The polarizing beam splitter that combines the pump and probe beam suppresses the reflected pump beam by a factor of ~ 100 as long as optical elements between the beam splitter and the sample have small birefringence. (This is usually, but not always, the case; for example, in diamond anvil cells, inhomogeneous stress fields can produce significant birefringence in the diamond anvil.) Additional suppression of the reflected pump beam is provided by one of two approaches that can be combined: (i) spatially separating the pump and probe and passing only the specularly reflected probe beam with an aperture; or (ii) spectrally separating the pump and probe and blocking the reflected pump light with an optical filter. (If an aperture is used, care must be taken that the aperture does not partially block the probe; if the probe is partially blocked, thermal lensing and small deflections created by misalignment of the pump and probe beams can create artifacts in the signals and errors in the measurement.) At Illinois, we choose, in most cases, to spectrally separate the pump and probe by a small amount, $\approx 7 \text{ nm}$ using what we call a “two-tint” approach.³⁸⁵ Others groups have chosen to frequency double either the pump or probe.^{403,404} Double-modulation, i.e., adding a mechanical chopper to the probe path and audio-frequency lock-in detection of the output of the rf-lockin, eliminates any remaining background from the pump beam and also removes coherent pick-up by the rf-lockin.

Fiber laser oscillators operating at $1.55 \mu\text{m}$ (Er: fiber) or $1.05 \mu\text{m}$ (Yb: fiber) have also been used for TDTR measurements.^{401,402,405} The advantage of fiber lasers over Ti:sapphire lasers is the smaller footprint and lower cost. Also, Si is transparent at $1.55 \mu\text{m}$, and therefore, an Er: fiber source can be used in experiments where it is desired to pass the pump and probe beam through the backside of a Si wafer.^{405,406} (At $1.55 \mu\text{m}$, Si has the useful property that the transient absorption near zero delay time can be used as optical thermometer with high bandwidth and high spatial resolution.⁴⁰⁷) At Illinois, we have not been fully satisfied these fiber lasers for general-purpose TDTR experiments because of problems in reproducing low-noise mode-lock conditions and the lack of suitable metal film transducers with the desired properties of high thermoreflectance, strong picosecond acoustics signals, high thermal conductivity, and small optical absorption depth that are provided by Al at Ti:sapphire wavelengths. While some metals have large thermoreflectance, the picosecond acoustic signals are typically weak and therefore the determination of the metal film thickness (the most important parameter in the data analysis) has to be accomplished by some other method such as x-ray reflectivity or Rutherford backscattering spectrometry. Since uncertainty in the thickness of the metal film transducer is usually the dominant source of uncertainty in the measurement,^{400,408} the inability to measure the transducer thickness by picosecond acoustics is a significant disadvantage.

4. Signal analysis and sensitivities

Signal analysis in TDTR is complicated by the presence of multiple time and frequency scales associate with the repetition rate of the laser and the modulation of the pump

beam.^{384,398} Because the temperature excursions are a small fraction of absolute temperature, the system response is linear in both the pump and probe powers, and therefore, a frequency domain and time-domain descriptions of the signals are equivalent. In the frequency domain, the output of the laser oscillator is a comb of delta functions separated by 80 MHz. Modulation of the pump beam at $f = 10$ MHz produces side bands at 10, 70, 90, 150, 170 MHz, etc. The action of the lockin amplifier is to evaluate the convolution of the frequency comb of the probe with the thermal response of the sample at these side bands. The time delay between pump and probe is equivalent to a phase shift in the frequency domain. The in-phase signal V_{in} is thus given by a sum of the real part of the frequency response at these side bands. When the modulation frequency is $\approx 1/8$ of the repetition rate of the laser oscillator, pulse accumulation effects for V_{in} are minimized and V_{in} approximates the thermal response of the sample to a single pump optical pulse. The out-of-phase signal V_{out} is mostly dominated by the imaginary part of the frequency response at the lowest frequency component (10 MHz) with much smaller contributions to V_{out} coming from the differences between side bands at higher frequencies, e.g., the difference between the imaginary parts of frequency response at 90 and 70 MHz.

In most cases, the greatest sensitivity to the thermal properties of the material under study comes from V_{out} , not the time-evolution of V_{in} as was conventionally analyzed prior to 2004.^{76,389} V_{in} is mostly dependent on the heat capacity per unit area of the metal film transducer hC_t , which is known from independent measurements, and the thermal conductance of the transducer/sample interface G . The sensitivity of V_{in} to the thermal conductivity of the sample κ is usually small because only a minority fraction of the heat deposited by the pump optical pulse has moved from the metal film transducer to the sample at large delay times. V_{out} is mostly dependent on κ , and to a lesser extent, G and hC_t . The out-of-phase signal does not vary significantly with delay time. Analysis of the ratio V_{in}/V_{out} is essentially a method of normalizing V_{out} by the known value of hC_t while at the same time providing a measurement of G and eliminating artifacts created by changes in the rms spot size or small variations in the alignment of the pump and probe as a function of delay time.

It is useful to consider the relative magnitudes of an effective thermal conductance per unit area that is characteristic of the metal film transducer, the interface, and the sample. For the metal film transducer, we define $G_t = hC\omega$, where ω is the angular frequency of the modulation of the pump beam. For typical parameters, $G_t \approx 10 \text{ MW m}^{-2} \text{ K}^{-1}$. A common interface conductance is $G = 200 \text{ MW m}^{-2} \text{ K}^{-1}$. For κ ranging from 0.1 to $1000 \text{ W m}^{-1} \text{ K}^{-1}$, the effective conductance of the sample $G_s = \sqrt{\omega\kappa C}$ varies between $G_s \approx 3$ and $G_s \approx 300 \text{ MW m}^{-2} \text{ K}^{-1}$. Sensitivity to κ is typically largest when $G_t < G_s < G$. (Similarly, TDTR measurements are most sensitive to the interface conductance G , when $G_t < G < G_s$.) Thus, TDTR measurements have reduced sensitivity when κ is very small $G_s < G_t$ or when κ is very high, $G_s > G$.

We can make the term “sensitivity” quantitative by defining a sensitivity coefficient that is the logarithmic derivative of the ratio signal $\phi = -V_{in}/V_{out}$ with respect to the parameter of interest. For example, the sensitivity coefficient for thermal conductivity is $S_\kappa = d\ln(\phi)/d\ln(\kappa)$. Figure 20 shows how S_κ and S_G vary with κ for two values of G that bracket common values for G encountered in TDTR measurements, and two values of the delay time.

Pump-probe measurements that use two laser oscillators with slightly different repetition rates to rapidly scan the delay time between pump and probe are often referred to as asynchronous optical sampling (ASOPS). While ASOPS has clear advantages in accessing long delay times over the more traditional form of modulated pump-probe metrology where the delay time is limited by the length of a delay stage, the lack of a frequency component in the measurement less than the repetition rate reduces the sensitivity in many thermal measurements. In other words, a typical ASOPS experiment can only access the frequency response of the sample at multiples of the laser repetition rate while a conventional TDTR experiment accesses the frequency response at the modulation frequency in addition to the side bands that straddle each multiple of the repetition rate. This extra frequency

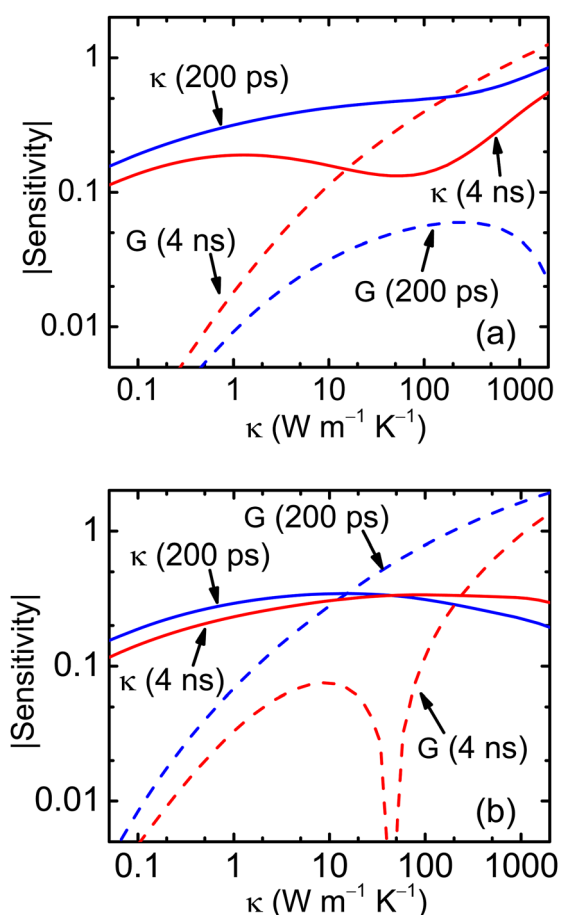


FIG. 20. Absolute value of the sensitivity coefficients for thermal conductivity S_κ (solid lines) and interface thermal conductance S_G (dashed lines) for a prototypical TDTR measurement of a thermally thick sample of thermal conductivity κ . The Al film thickness is 80 nm; the $1/e^2$ radius of the laser spot is $10 \mu\text{m}$; and the heat capacity of the sample is $2 \text{ J cm}^{-3} \text{ K}^{-1}$. In panel (a), $G = 200 \text{ MW m}^{-2} \text{ K}^{-1}$; in panel (b), $G = 40 \text{ MW m}^{-2} \text{ K}^{-1}$.

component is critical in enhancing the sensitivity to the thermal conductivity of most samples. (Dilhare and co-workers have recently described an approach that combines ASOPS with high frequency modulation of the pump beam.⁴⁰⁹) Furthermore, ASOPS evenly weights all delay times in the data acquisition, while the TDTR data acquisition can be weighted to optimize the signal-to-noise in the regions of delay time that are most important for the measurement. For example, we typically acquire data in 1 ps intervals between -20 and 40 ps to accurately determine the relative phase of the rf lockin reference channel and measure the film thickness by picosecond acoustic signals, and then acquire data from 40 ps to 4 ns with ≈ 50 data points that are spaced uniformly in $\log(t)$.

Schmidt *et al.*⁴¹⁰ analyzed the sensitivities of frequency-domain thermoreflectance (FDTR), i.e., experiments performed with a modulated cw laser as the pump and a second cw laser as the probe. The maximum sensitivity to the thermal conductivity κ of a bulk sample in an FDTR experiment is ≈ 0.15 ;⁴¹⁰ for comparison, the maximum sensitivity to κ in a TDTR experiment is ≈ 0.5 , see Fig. 20. FDTR has recently been extended to frequencies as high as 200 MHz using heterodyne detection to improve the signal-to-noise, an approach the authors refer to as broad-band frequency-domain thermoreflectance (BB-FDTR).^{411,412} Measurements of the thermal response of a sample between 200 kHz and 200 MHz are essentially equivalent to accessing time-scales for thermal transport between ~ 1 μ s and ~ 1 ns. FDTR cannot provide measurements of the picosecond time-scale acoustics that are available in TDTR but has advantages in the simplicity of the laser sources and the ability to continuously scan frequency over a wide range. Information about the high frequency response is also embedded in the delay-time dependence of the in-phase TDTR signal; analysis of TDTR data is, however, more complex because it involves summations of the thermal response over many frequencies.³⁸⁴

5. Forefront issues

All analysis of TDTR data to-date has depended on the assumption that (i) the diffusion equations is an adequate description of the temperature fields and heat fluxes; and (ii) boundaries between materials can be described by a radiative boundary condition that relates a temperature drop at an interface ΔT to the heat flux crossing the interface J_Q : $J_Q = G\Delta T$. Neither assumption can be rigorously justified unless the mean-free-paths of all of the significant heat carriers are small compared to the thermal penetration depth and couplings between various heat carriers is sufficiently strong that the distribution of heat carriers can be described by a single temperature.

When the thermal conductivity κ extracted experiment varies in an unexpected way, e.g., when κ varies with frequency^{413,414} or the size of the laser spot,⁴¹⁵ the approach has been to interpret the result as a value that is modified by the exclusion of phonons that have mean-free-paths larger than a spatial scale in the experiment. This is, of course, not a completely satisfying situation because the modified value of κ is extracted from the data using an analysis based on the

diffusion equation while simultaneously concluding that the diffusion equation cannot accurately describe the experiment. While semiconductor alloys⁴¹³ and specific type of amorphous Si⁴¹⁴ show pronounced frequency dependence in TDTR measurements, pure single crystals generally do not. It is possible that the frequency dependence is present but too weak to observe clearly. The spot-size effects at low temperature are large.⁴¹⁵ Frequency domain measurements at very high frequencies⁴¹¹ may provide new insights.

The assumption that an interface in a TDTR experiment can be described as a simple radiative boundary condition on the diffusion equation has received little critical attention. This assumption ignores the true complexity of the problem—and misses a great deal of interesting science—because it assumes that the temperature is well-defined at each side of the interface; in other words, this description assumes that all excitations of each material are in equilibrium with each other at all distances from the interface. Using current methods and materials, we cannot usually resolve these effects. One exception was an experiment that combined TDTR with time-resolved x-ray diffraction where a significant fraction of the heat leaving an Al/GaAs interface was interpreted as evidence of phonons that did not equilibrate in a near-surface layer, because the mean-free-paths of these heat-carrying phonons were larger than the thickness of the near-surface layer.⁴¹⁶ The effects of ballistic carriers on heat transport near a surface has continued to attract a great deal of attention in both experiment and theory,^{415,417–421} but we do not yet have a self-consistent method of incorporating these ideas into the analysis of TDTR data.

B. Scanning thermal microscopy

In scanning thermal microscopy (SThM), a nanometer-scale probe tip measures the temperature of a surface with nanometer-scale spatial resolution. The temperature measurement has spatial and temperature resolution governed by the size of the tip, size, and shape of the substrate features, and the heat transfer between tip and substrate. Over the last decade, there have been significant improvements in spatial resolution to ≈ 10 nm, temperature precision to ≈ 50 mK, and measurements of nanometer-scale heat flows to ≈ 10 pW.

The temperature can be measured using either a temperature sensor in the cantilever tip or by measuring local thermomechanical strains in either the sample or the cantilever.^{422,423} The temperature sensor integrated into a scanning probe tip can be either a thermocouple⁴²⁴ or a thermistor. Noncontact SThM near-field optical probes can measure temperature with spatial resolution as small as 10 nm.⁴²⁵ These near-field optical measurements do not rely on thermal conduction between tip and sample, but instead are governed by near-field electromagnetic interactions between the tip and the sample, which are still not fully understood, see Sec. IIB.

For both thermocouple- and thermistor-based SThM temperature measurements, the key technical challenge is to understand how the measured temperature signal relates to the substrate temperature. Several groups have investigated heat transport between a nanometer-scale tip and a

surface.^{426,427} The key thermal resistances are thermal contact resistance between the nanometer-scale tip and the surface, as well as thermal resistance within the tip. In some cases, water condenses at the tip-sample interface and thus decreases the tip-sample thermal resistance. In the absence of water and for a substrate having large thermal conductivity, typical values are 10^7 K W^{-1} for the tip-sample thermal resistance and 10^6 K W^{-1} for the internal tip resistance. It is also important to understand the heat flow between the cantilever and the substrate, which can induce artifacts into the measurement. Such artifacts can be accounted for using detailed modeling or by subtracting the background thermal signal from the local tip thermal signal.^{428–430} SThM measurements in vacuum show that by operating the tip without air or water in the environment can avoid such artifacts.⁴³¹

Scanning thermal microscopy can also measure thermoelectric properties at the nanometer scale. In these measurements, the tip is held at a different temperature than the substrate, such that there is a temperature gradient near the tip-sample interface. The temperature gradient in the tip or near the tip can be as high at 10 K nm^{-1} . The tip measures the temperature-dependent voltage at the tip-sample contact to extract the local Seebeck coefficient. The measurement can be significantly faster when the tip and/or sample temperature is locally controlled, for example, using a microfabricated heater. SThM-based thermoelectric measurements have measured thermoelectricity in organic molecules,⁴³² Seebeck measurements on doped semiconductor surfaces,⁴³³ and thermal conductance at a nanometer-scale point contacts.^{434,435}

Nanometer-scale temperature distributions can be measured from nanometer-scale thermomechanical expansions.^{422,423,436,437} In Scanning Joule Expansion microscopy (SJEM), a scanning probe tip measures local thermomechanical distributions on a periodically heated substrate. For simple substrate geometries, the measured temperature distribution can be directly converted to temperature. For more complex structures, models of heat flow and thermomechanical deformations can help to resolve the temperature distribution. When the SThM-measured temperature distribution is known with high spatial resolution and precision, it is possible to calculate local heat flows and substrate thermophysical properties.^{422,423,438} The nanometer-scale temperature distributions that can be observed using SThM offer deep insight into device operation that cannot be achieved using non-local thermometry such as optical techniques and microfabricated thermometers. SThM measurements can reveal ballistic electron transport and the impact of boundary effects in carbon nanotubes⁴²⁸ and graphene devices.⁴²²

C. Nanometer-scale thermal analysis and thermal nano-manufacturing

Advancements in thermal scanning probe tools has enabled new capabilities in nanometer-scale materials characterization and nano-manufacturing. When the heated tip of an atomic force microscope (AFM) cantilever tip is in contact with a solid substrate, the tip-substrate interface forms a hotspot that is as small as a few nanometers in diameter.⁴²⁶

The temperature of this hotspot can be accurately controlled over a large temperature range. The ability to control the nanometer-scale hotspot at the end of a scanning probe tip has enabled fundamental measurements of nanometer-scale heat flow as well as new applications, such as nanometer-scale thermophysical property measurements and nanometer-scale thermal manufacturing.

Nano-thermal analysis (nano-TA) uses a heatable AFM cantilever to measure the phase change temperature of a material with nanometer-scale spatial resolution.^{439–441} In nano-TA, the AFM system holds the heated tip in contact with the surface while raising the cantilever tip temperature. As the material in contact with the tip softens, the tip penetrates into the surface and this tip motion is detected by the AFM. Nano-TA can be combined with other AFM-based measurements of substrate topography, adhesion, or stiffness. The combination of these techniques allows spatially resolved measurements of phase change temperature and temperature-dependent mechanical stiffness and loss modulus. The amount of material that can be interrogated using Nano-TA is on the order of attograms to femtograms, which is much smaller than state of the art calorimetry, thermogravimetry, or rheometry all of which require milligrams to gram scale quantities. Unlike these macroscopic techniques, Nano-TA can be performed on thin film samples and as-prepared material and can be used for failure analysis or the study of multi-component systems such as multi-layer film stacks or nanocomposites. Nano-TA measurement techniques have been aided by rigorous electro-thermal calibration and modeling,^{442,443} as well as by improvements in cantilever design, fabrication, and instrumentation.^{444–451}

The nanometer-scale hot spot at the end of a heated AFM tip can be used for nanometer-scale thermal manufacturing.^{452–459} Thermal nano-manufacturing can form nanostructures with speed faster than 1 mm s^{-1} , which is close to that of commercial electron beam lithography tools. The high speed of thermal nano-manufacturing is due to the high temperatures that can be achieved to increase mass diffusivity or chemical conversion speed. The size of the features that can be written are governed by the size and shape of the temperature field near the tip, and in some cases, the mechanical properties of the substrate being processed.^{453,455}

In thermal dip-pen nanolithography (tDPN), a heated AFM tip can be used to deposit solid meltable ink from a tip onto a substrate with sub-100 nm feature size.^{453,455,457} The flow of the ink from the tip to the surface is controlled via the tip temperature. Thus, when the cantilever tip is cold, no ink flows and the tip can be used to perform nanometrology without contaminating the surface. When the tip is hot, ink flows from the tip to the surface due to the surface forces at the tip-polymer-substrate interface.⁴⁵⁵ Materials that have been written using thermal dip-pen nanolithography include many polymers, low melting temperature metals, and nanoparticle-polymer composites.

In thermochemical nanolithography (TCNL), a heated tip is brought into contact with a material that undergoes a temperature-dependent chemical reaction.^{452,454,456,458,459} For example, a polymer having a thermally removable functional group can be made reactive when its temperature is

raised. The reactive region can be smaller than 15 nm in size, owing to the very small heat affected zone near the heated AFM tip. Chemicals presented to the reactive size can form a chemical bond only in the thermally processed region. Examples of nanopatterned materials include nanoparticles and biological molecules. Another example of TCNL uses the heated tip to locally decompose a molecular glass to form three-dimensional nanopatterns.⁴⁵⁹

Finally, the heated tip can locally convert the electronic function of a material. For example, graphene oxide can be converted to graphene or an insulating organic layer can be locally converted into a semiconducting organic.⁴⁵⁶ With thermal nano-manufacturing, it is possible to control high temperature materials processing even on low temperature substrates, since the heat load on the substrate is small. For example, one study synthesized electroceramics directly on plastic substrates.⁴⁵⁸

VI. OUTLOOK

We discuss specific examples of outstanding scientific questions and recommendations for future work throughout this review but provide a brief summary of some selected examples here.

Research in the past decade has demonstrated the importance of interface physics and chemistry for thermal transport, but advances in experiment are needed to provide improved control and characterization of interface structure. Non-local contributions to the conductance—e.g., coupling of the electronic excitations of a material on one side of the interface to the vibrational excitations of a material on the other side of the interface—are not fully understood in theory and have not yet been systematically studied in experiment. Non-local contributions may be particularly important for weakly bonded interfaces, interfaces containing sub-nm gaps, and interfaces between materials with highly dissimilar lattice vibrations. Computer simulations of these channels for thermal transport will require extensions of molecular dynamics that include the interactions between quantum mechanical electronic states and vibrational states.

Extension of experiment and theory of near-field radiation to gaps on the order of 1 nm is challenging. Extremely perfect surface morphologies and more accurate methods of measuring the gap spacing are needed. Current theory is based on the long wavelength limit of the dielectric susceptibility $\epsilon(\omega)$. As the gap size decreases, the spatial frequency dependence, $\epsilon(q, \omega)$, becomes important.

Metal nanoparticles can be heated rapidly to extremely high temperatures using lasers with short pulse durations. In many cases of interest, the temperature excursion exceeds the boiling point or even the critical point of the solvent. Phase transformations under conditions of fast transient heating and steep temperature gradients are only beginning to be explored.

First principles theory can now predict the thermal conductivity of simple crystals. A challenge for the future is to extend this work to complex crystals that are of interest for thermoelectric energy conversion. Rigorous treatment of disorder and boundaries will require further advances in theory

and computational methods. It will be useful to make detailed comparisons between the results of first principles calculations and the phenomenological scattering rates and integral equations that are typically used by experimentalists to model their data.

Thermal conductivities far below the value expected for boundary scattering have been observed in Si nanowires prepared by metal-assisted chemical etching of Si wafers. The mechanism of the suppression is controversial and may be explained by the physics of phonon waves interacting with rough surface morphologies. Metal-assisted etching process is, however, a complex process and the condition of the near-surface region of etched Si nanowires requires further study.

Carbon nanotubes and graphene have extremely high thermal conductivity when fully isolated, but the conductivity is strongly suppressed when supported by a substrate, embedded in a matrix or otherwise encased in surrounding materials. Advances in theoretical understanding of the mechanisms that produce this suppression and fabrication methods that mitigate the effects of surrounding materials will be important for thermal management of nanoscale electronics and the synthesis of nanocomposites with high thermal conductivity.

Ultralow thermal conductivity has been observed in materials that combine disorder, high interface density, and strong anisotropy of elastic constants. The conductivity of disordered WSe₂ is far below the conventional prediction of the minimum thermal conductivity of an amorphous material. Thus, the lower limit of the conductivity of a fully dense solid is not known. The materials that combine ultralow thermal conductivity with high temperature stability are needed for improved thermal barriers.

TDTR has emerged as a powerful technique for measuring the thermal conductivity of thin layers and bulk materials with high spatial resolution, but analysis of TDTR data is currently based on a solution of the diffusion equation with interfaces treated as a radiative boundary condition on heat flux and temperature. Rigorous methods for analyzing TDTR data that incorporate models of phonon scattering rates and non-equilibrium between various heat carriers at interfaces are needed to better understand the results of TDTR experiments.

ACKNOWLEDGMENTS

D.G.C., P.V.B., G.C., S.F., P.K., and W.P.K. were supported by Air Force Office of Scientific Research MURI FA9550-08-1-0407. D.R.C. acknowledges support from the Office of Naval Research (N000-14-121-0993).

K.E.G. appreciates input from Mehdi Asheghi, Elah Bozorg-Grayeli, Jungwan Cho, Jaeho Lee, and Zijian Li. K.E.G. also thanks Duygu Kuzum, Rakesh Jeyasingh, and H. S. Philip Wong of the Stanford University Electrical Engineering Department for sharing unpublished microscopy and resistance-current data for a phase change nanocell.

The work of S.R.P. was supported as a subcontractor (AC) of the U.S. Government under DOE Contract No. DE-AC07-05ID14517, under the Energy Frontier Research

Center (Office of Science, Office of Basic Energy Science, FWP 1356). Accordingly, the U.S. Government retains and the publisher (by accepting the article for publication) acknowledges that the U.S. Government retains a nonexclusive, paid-up, irrevocable, world-wide license to publish or reproduce the published form of this manuscript, or allow others to do so, for U.S. Government purposes.

E.P. acknowledges input from Andrey Serov, Feng Xiong, Albert Liao, and financial support from a Presidential Early Career Award for Scientists and Engineers (PECASE).

L.S. acknowledges research support by NSF Thermal Transport Processes Program, Office of Naval Research, and Department of Energy, and conference support from NSF (CBET-1125957 and CBET-1152129) and Office of Naval Research (N00014-11-1-0467).

- ¹D. G. Cahill, W. K. Ford, K. E. Goodson, G. D. Mahan, A. Majumdar, H. J. Maris, R. Merlin, and S. R. Phillpot, "Nanoscale thermal transport," *J. Appl. Phys.* **93**, 793–818 (2003).
- ²A. Shakouri, "Recent developments in semiconductor thermoelectric physics and materials," *Annu. Rev. Mater. Res.* **41**, 399–431 (2011).
- ³T. M. Tritt, "Thermoelectric phenomena, materials, and applications," *Annu. Rev. Mater. Res.* **41**, 433–448 (2011).
- ⁴D. L. Medlin and G. J. Snyder, "Interfaces in bulk thermoelectric materials," *Curr. Opin. Colloid Interface Sci.* **14**, 226–235 (2009).
- ⁵C. J. Vineis, A. Shakouri, A. Majumdar, and M. G. Kanatzidis, "Nanostructured thermoelectrics: Big efficiency gains from small features," *Adv. Mater.* **22**, 3970–3980 (2010).
- ⁶M. G. Kanatzidis, "Nanostructured thermoelectrics: The new paradigm?" *Chem. Mater.* **22**, 648–659 (2010).
- ⁷E. S. Toberer, L. L. Baranowski, and C. Dames, "Advances in thermal conductivity," *Annu. Rev. Mater. Res.* **42**, 179–209 (2012).
- ⁸A. A. Balandin, "Thermal properties of graphene and nanostructured carbon materials," *Nature Mater.* **10**, 569–581 (2011).
- ⁹M. M. Sadeghi, M. T. Pettes, and L. Shi, "Thermal transport in graphene," *Solid State Commun.* **152**, 1321–1330 (2012).
- ¹⁰A. M. Marconnet, M. A. Panzer, and K. E. Goodson, "Thermal conduction phenomena in carbon nanotubes and related nanostructured materials," *Rev. Mod. Phys.* **1**, 1295–1326 (2013).
- ¹¹N. Li, J. Ren, L. Wang, G. Zhang, P. Hänggi, and B. Li, "Colloquium: Phonons: Manipulating heat flow with electronic analogs and beyond," *Rev. Mod. Phys.* **84**, 1045–1066 (2012).
- ¹²A. Weathers and L. Shi, "Thermal transport measurement techniques for nanowires and nanotubes," *Annu. Rev. Heat Transfer* **15**, 101–134 (2013).
- ¹³G. Chen, *Nanoscale Energy Transport and Conversion: A Parallel Treatment of Electrons, Molecules, Phonons, and Photons* (Oxford University Press, 2005).
- ¹⁴Z. Zhang, *Nano/Microscale Heat Transfer* (McGraw-Hill Professional, 2007).
- ¹⁵S. Volz, R. Carminati, P. Chantrenne, S. Dilhaire, S. Gomez, N. Trannoy, and G. Tessier, *Microscale and Nanoscale Heat Transfer* (Springer, 2007).
- ¹⁶C. Sobhan and G. Peterson, *Microscale and Nanoscale Heat Transfer: Fundamentals and Engineering Applications* (CRC Press, 2008).
- ¹⁷I. M. Khalatnikov, "Teploobmen Mezhdru Tverdym Telom I Gelium II," *Sov. Phys. JETP* **22**, 687–704 (1952).
- ¹⁸I. Khalatnikov, *An Introduction to the Theory of Superfluidity* (Benjamin, New York, 1965).
- ¹⁹D. Young and H. Maris, "Lattice-dynamical calculation of the Kapitza resistance between fcc lattices," *Phys. Rev. B* **40**, 3685 (1989).
- ²⁰S. Pettersson and G. D. Mahan, "Theory of the thermal boundary resistance between dissimilar lattices," *Phys. Rev. B* **42**, 7386–7390 (1990).
- ²¹B. N. Persson and H. Ueba, "Heat transfer between graphene and amorphous Si O₂," *J. Phys. Condens. Matter* **22**, 462201 (2010).
- ²²P. Hyldgaard and G. D. Mahan, "Phonon superlattice transport," *Phys. Rev. B* **56**, 10754 (1997).
- ²³M. V. Simkin and G. D. Mahan, "Minimum thermal conductivity of superlattices," *Phys. Rev. Lett.* **84**, 927 (2000).
- ²⁴S. M. Lee, D. G. Cahill, and R. Venkatasubramanian, "Thermal conductivity of Si-Ge superlattices," *Appl. Phys. Lett.* **70**, 2957 (1997).
- ²⁵S. T. Huxtable, A. R. Abramson, C. L. Tien, A. Majumdar, C. LaBounty, X. F. Fan, G. Zeng, J. E. Bowers, A. Shakouri, and E. T. Croke, "Thermal conductivity of Si/SiGe and SiGe/SiGe superlattices," *Appl. Phys. Lett.* **80**, 1737 (2002).
- ²⁶M. L. Lee and R. Venkatasubramanian, "Effect of nanodot areal density and period on thermal conductivity in SiGe/Si nanodot superlattice," *Appl. Phys. Lett.* **92**, 053112 (2008).
- ²⁷V. Rawat, Y. K. Koh, D. G. Cahill, and T. D. Sands, "Thermal conductivity of (Zr,W)/N/ScN metal/semiconductor multilayers and superlattices," *J. Appl. Phys.* **105**, 024909 (2009).
- ²⁸Y. K. Koh, Y. L. Cao, D. G. Cahill, and D. Jena, "Heat-transport mechanisms in superlattices," *Adv. Funct. Mater.* **19**, 610–615 (2009).
- ²⁹B. C. Gundrum, D. G. Cahill, and R. S. Averback, "Thermal conductance of metal-metal interfaces," *Phys. Rev. B* **72**, 245426 (2005).
- ³⁰R. B. Wilson and D. G. Cahill, "Experimental validation of the interfacial form of the Wiedemann-Franz law," *Phys. Rev. Lett.* **108**, 255901 (2012).
- ³¹G. D. Mahan, "Thermal transport in AB superlattices," *Phys. Rev. B* **83**, 125313 (2011).
- ³²M. P. Allen and D. J. Tildesley, *Computer Simulation of Liquids* (Oxford Science Publications, 1986).
- ³³A. J. C. Ladd, B. Moran, and W. G. Hoover, "Lattice thermal conductivity: A comparison of molecular dynamics and anharmonic lattice dynamics," *Phys. Rev. B* **34**, 5058 (1986).
- ³⁴P. K. Schelling, S. R. Phillpot, and P. Keblinski, "Comparison of atomic-level simulation methods for computing thermal conductivity," *Phys. Rev. B* **65**, 144306 (2002).
- ³⁵J.-L. Barrat and F. Chiaruttini, "Kapitza resistance at the liquid-solid interface," *Mol. Phys.* **101**, 1605–1610 (2003).
- ³⁶S. Shenogin, P. Keblinski, D. Bedrov, and G. Smith, "Thermal relaxation mechanism and role of chemical functionalization in fullerene solutions," *J. Chem. Phys.* **124**, 014702 (2006).
- ³⁷P. Schelling, S. Phillpot, and P. Keblinski, "Phonon wave-packet dynamics at semiconductor interfaces by molecular-dynamics simulation," *Appl. Phys. Lett.* **80**, 2484 (2002).
- ³⁸P. K. Schelling, S. R. Phillpot, and P. Keblinski, "Kapitza conductance and phonon scattering at grain boundaries by simulation," *J. Appl. Phys.* **95**, 6082 (2004).
- ³⁹W. Zhang, T. Fisher, and N. Mingo, "The atomistic Green's function method: An efficient simulation approach for nanoscale phonon transport," *Numer. Heat Transfer, Part B* **51**, 333–349 (2007).
- ⁴⁰P. Hopkins, P. Norris, M. Tsegaye, and A. Ghosh, "Extracting phonon thermal conductance across atomic junctions: Nonequilibrium Green's function approach compared to semiclassical methods," *J. Appl. Phys.* **106**, 063503 (2009).
- ⁴¹N. Mingo and L. Yang, "Phonon transport in nanowires coated with an amorphous material: An atomistic Green's function approach," *Phys. Rev. B* **68**, 245406 (2003).
- ⁴²M. Hu, P. Keblinski, and P. Schelling, "Kapitza conductance of silicon-amorphous polyethylene interfaces by molecular dynamics simulations," *Phys. Rev. B* **79**, 104305 (2009).
- ⁴³E. Swartz and R. Pohl, "Thermal boundary resistance," *Rev. Mod. Phys.* **61**, 605 (1989).
- ⁴⁴N. Shenogina, R. Godawat, P. Keblinski, and S. Garde, "How wetting and adhesion affect thermal conductance of a range of hydrophobic to hydrophilic aqueous interfaces," *Phys. Rev. Lett.* **102**, 156101 (2009).
- ⁴⁵Z. B. Ge, D. G. Cahill, and P. V. Braun, "Thermal conductance of hydrophilic and hydrophobic interfaces," *Phys. Rev. Lett.* **96**, 186101 (2006).
- ⁴⁶M. Shen, W. Evans, D. Cahill, and P. Keblinski, "Bonding and pressure-tunable interfacial thermal conductance," *Phys. Rev. B* **84**, 195432 (2011).
- ⁴⁷T. Luo and J. Lloyd, "Non-equilibrium molecular dynamics study of thermal energy transport in Au-SAM-Au junctions," *Int. J. Heat Mass Transfer* **53**, 1–11 (2010).
- ⁴⁸L. Hu, L. Zhang, M. Hu, J.-S. Wang, B. Li, and P. Keblinski, "Phonon interference at self-assembled monolayer interfaces: Molecular dynamics simulations," *Phys. Rev. B* **81**, 235427 (2010).
- ⁴⁹Z. Ong and E. Pop, "Frequency and polarization dependence of thermal coupling between carbon nanotubes and SiO₂," *J. Appl. Phys.* **108**, 103502 (2010).
- ⁵⁰G. S. Hwang and M. Kaviani, "Molecular dynamics simulation of effective thermal conductivity of vapor-filled nanogap and nanocavity," *J. Appl. Phys.* **106**, 024317 (2009).

- ⁵¹C. Caroli, R. Combescot, P. Nozieres, and D. Saint-James, "Direct calculation of the tunneling current," *J. Phys. C* **4**, 916 (1971).
- ⁵²R. Prasher, "Acoustic mismatch model for thermal contact resistance of van der Waals contacts," *Appl. Phys. Lett.* **94**, 041905 (2009).
- ⁵³T. English, J. Duda, J. Smoyer, D. Jordan, P. Norris, and L. Zhigilei, "Enhancing and tuning phonon transport at vibrationally mismatched solid-solid interfaces," *Phys. Rev. B* **85**, 035438 (2012).
- ⁵⁴R. Stevens, L. Zhigilei, and P. Norris, "Effects of temperature and disorder on thermal boundary conductance at solid-solid interfaces: Nonequilibrium molecular dynamics simulations," *Int. J. Heat Mass Transfer* **50**, 3977–3989 (2007).
- ⁵⁵M. Hu, J. Goicochea, B. Michel, and D. Poulikakos, "Water nanoconfinement induced thermal enhancement at hydrophilic quartz interfaces," *Nano Lett.* **10**, 279–285 (2010).
- ⁵⁶M. Hu, X. Zhang, D. Poulikakos, and C. Grigoropoulos, "Large near junction thermal resistance reduction in electronics by interface nano-engineering," *Int. J. Heat Mass Transfer* **54**, 5183–5191 (2011).
- ⁵⁷K. Termentzidis, P. Chantrenne, and P. Keblinski, "Nonequilibrium molecular dynamics simulation of the in-plane thermal conductivity of superlattices with rough interfaces," *Phys. Rev. B* **79**, 214307 (2009).
- ⁵⁸P. Kim, L. Shi, A. Majumdar, and P. McEuen, "Thermal transport measurements of individual multiwalled nanotubes," *Phys. Rev. Lett.* **87**, 215502 (2001).
- ⁵⁹A. A. Balandin, S. Ghosh, W. Bao, I. Calizo, D. Teweldebrhan, F. Miao, and C. N. Lau, "Superior thermal conductivity of single-layer graphene," *Nano Lett.* **8**, 902–907 (2008).
- ⁶⁰S. Huxtable, D. Cahill, S. Shenogin, L. Xue, R. Ozisik, P. Barone, M. Usrey, M. Strano, G. Siddons, M. Shim *et al.*, "Interfacial heat flow in carbon nanotube suspensions," *Nature Mater.* **2**, 731–734 (2003).
- ⁶¹S. Shenogin, L. Xue, R. Ozisik, P. Keblinski, and D. G. Cahill, "Role of thermal boundary resistance on the heat flow in carbon-nanotube composites," *J. Appl. Phys.* **95**, 8136 (2004).
- ⁶²T. G. D. L. Hu and P. Keblinski, "Thermal transport in graphene-based nanocomposite," *Phys. Rev. B* **83**, 195423 (2011).
- ⁶³L. Hu, T. Desai, and P. Keblinski, "Thermal transport in graphene-based nanocomposite," *J. Appl. Phys.* **110**, 033517 (2011).
- ⁶⁴P. Keblinski, S. Phillpot, S. Choi, and J. Eastman, "Mechanisms of heat flow in suspensions of nano-sized particles (nanofluids)," *Int. J. Heat Mass Transfer* **45**, 855–863 (2002).
- ⁶⁵L. Xue, P. Keblinski, S. Phillpot, S.-S. Choi, and J. Eastman, "Effect of liquid layering at the liquidsolid interface on thermal transport," *Int. J. Heat Mass Transfer* **47**, 4277–4284 (2004).
- ⁶⁶J. Eapen, J. Li, and S. Yip, "Mechanism of thermal transport in dilute nanocolloids," *Phys. Rev. Lett.* **98**, 28302 (2007).
- ⁶⁷O. M. Wilson, X. Y. Hu, D. G. Cahill, and P. V. Braun, "Colloidal metal particles as probes of nanoscale thermal transport in fluids," *Phys. Rev. B* **66**, 224301 (2002).
- ⁶⁸S. Merabia, P. Keblinski, L. Joly, L. Lewis, and J. Barrat, "Critical heat flux around strongly heated nanoparticles," *Phys. Rev. E* **79**, 021404 (2009).
- ⁶⁹M. Hu, H. Petrova, and G. Hartland, "Investigation of the properties of gold nanoparticles in aqueous solution at extremely high lattice temperatures," *Chem. Phys. Lett.* **391**, 220–225 (2004).
- ⁷⁰A. Plech, V. Kotaidis, S. Gresillon, C. Dahmen, and G. Von Plessen, "Laser-induced heating and melting of gold nanoparticles studied by time-resolved x-ray scattering," *Phys. Rev. B* **70**, 195423 (2004).
- ⁷¹R. Costescu, M. Wall, and D. Cahill, "Thermal conductance of epitaxial interfaces," *Phys. Rev. B* **67**, 054302 (2003).
- ⁷²A. Hanisch, B. Krenzer, T. Pelka, S. Mollenbeck, and M. Horn-von Hoegen, "Thermal response of epitaxial thin Bi films on Si (001) upon femtosecond laser excitation studied by ultrafast electron diffraction," *Phys. Rev. B* **77**, 125410 (2008).
- ⁷³B. Krenzer, A. Hanisch-Blicharski, P. Schneider, T. Payer, S. Mollenbeck, O. Osmani, M. Kammler, R. Meyer, and M. Horn-von Hoegen, "Phonon confinement effects in ultrathin epitaxial bismuth films on silicon studied by time-resolved electron diffraction," *Phys. Rev. B* **80**, 024307 (2009).
- ⁷⁴H. K. Lyo and D. G. Cahill, "Thermal conductance of interfaces between highly dissimilar materials," *Phys. Rev. B* **73**, 144301 (2006).
- ⁷⁵T. Frigge, A. Kalus, F. Klasing, M. Kammler, A. Hanisch-Blicharski, and M. H. von Hoegen, "Nanoscale Heat Transport in Self-Organized Ge Clusters on Si(001)," in *Nanoscale Thermoelectrics 2012-Materials and Transport Phenomena* (Mater. Res. Soc. Symp. Proc., 2013), Vol. 1456, pp. mrss12–1456–jj05–08.
- ⁷⁶R. M. Costescu, D. G. Cahill, F. H. Fabreguette, Z. A. Sechrist, and S. M. George, "Ultra-low thermal conductivity in W/Al₂O₃ nanolaminates," *Science* **303**, 989–990 (2004).
- ⁷⁷Y. S. Ju, M.-T. Hung, and T. Usui, "Nanoscale heat conduction across metal-dielectric interfaces," *J. Heat Transfer* **128**, 919–925 (2006).
- ⁷⁸Z. J. Li, S. Tan, E. Bozorg-Grayeli, T. Kodama, M. Asheghi, G. Delgado, M. Panzer, A. Pokrovsky, D. Wack, and K. E. Goodson, "Phonon dominated heat conduction normal to Mo/Si multilayers with period below 10 nm," *Nano Lett.* **12**, 3121–3126 (2012).
- ⁷⁹Y. Jin, A. Yadav, K. Sun, H. Sun, K. P. Pipe, and M. Shtein, "Thermal boundary resistance of copper phthalocyanine-metal interface," *Appl. Phys. Lett.* **98**, 093305 (2011).
- ⁸⁰Y. Jin, C. Shao, J. Kieffer, K. P. Pipe, and M. Shtein, "Origins of thermal boundary conductance of interfaces involving organic semiconductors," *J. Appl. Phys.* **112**, 093503 (2012).
- ⁸¹A. Majumdar and P. Reddy, "Role of electronphonon coupling in thermal conductance of metal-nonmetal interfaces," *Appl. Phys. Lett.* **84**, 4768 (2004).
- ⁸²J. Ordóñez-Miranda, J. J. Alvarado-Gil, and R. Yang, "The effect of the electron-phonon coupling on the effective thermal conductivity of metal-nonmetal multilayers," *J. Appl. Phys.* **109**, 094310 (2011).
- ⁸³W. Wang and D. G. Cahill, "Limits to thermal transport in nanoscale metal bilayers due to weak electron-phonon coupling in Au and Cu," *Phys. Rev. Lett.* **109**, 175503 (2012).
- ⁸⁴A. Birch, W. R. G. Kemp, P. G. Klemens, and R. J. Tainsh, "The lattice thermal conductivity of some gold alloys," *Aust. J. Phys.* **12**, 455 (1959).
- ⁸⁵C. Chiriac, D. G. Cahill, N. Nguyen, D. Johnson, A. Bodapati, P. Keblinski, and P. Zschack, "Ultralow thermal conductivity in disordered, layered WSe₂ crystals," *Science* **315**, 351–353 (2007).
- ⁸⁶D. G. Cahill, S. K. Watson, and R. O. Pohl, "Lower limit to the thermal conductivity of disordered crystals," *Phys. Rev. B* **46**, 6131–6140 (1992).
- ⁸⁷Z. Wei, Y. Chen, and C. Dames, "Negative correlation between in-plane bonding strength and cross-plane thermal conductivity in a model layered material," *Appl. Phys. Lett.* **102**, 011901 (2013).
- ⁸⁸Z. Chen, Z. Wei, Y. Chen, and C. Dames, "Anisotropic Debye model for the thermal boundary conductance," *Phys. Rev. B* **87**, 125426 (2013).
- ⁸⁹P. K. Jain, X. H. Huang, I. H. El-Sayed, and M. A. El-Sayed, "Noble metals on the nanoscale: Optical and photothermal properties and some applications in imaging, sensing, biology, and medicine," *Acc. Chem. Res.* **41**, 1578–1586 (2008).
- ⁹⁰P. Keblinski, D. G. Cahill, A. Bodapati, C. R. Sullivan, and T. A. Taton, "Limits of localized heating by electromagnetically excited nanoparticles," *J. Appl. Phys.* **100**, 054305 (2006).
- ⁹¹C. T. Avedisian, R. E. Cavicchi, P. L. McEuen, and X. Zhou, "Nanoparticles for cancer treatment: Role of heat transfer," *Ann. N.Y. Acad. Sci.* **1161**, 62–73 (2009).
- ⁹²Z. B. Ge, D. G. Cahill, and P. V. Braun, "AuPd metal nanoparticles as probes of nanoscale thermal transport in aqueous solution," *J. Phys. Chem. B* **108**, 18870–18875 (2004).
- ⁹³G. V. Hartland, "Optical studies of dynamics in noble metal nanostructures," *Chem. Rev.* **111**, 3858–3887 (2011).
- ⁹⁴R. Y. Wang, R. A. Segalman, and A. Majumdar, "Room temperature thermal conductance of alkanedithiol self-assembled monolayers," *Appl. Phys. Lett.* **89**, 173113 (2006).
- ⁹⁵M. D. Losego, M. E. Grady, N. R. Sottos, D. G. Cahill, and P. V. Braun, "Effects of chemical bonding on heat transport across interfaces," *Nature Mater.* **11**, 502–506 (2012).
- ⁹⁶C. D. Bain, E. B. Troughton, Y. T. Tao, J. Evall, G. M. Whitesides, and R. G. Nuzzo, "Formation of monolayer films by the spontaneous assembly of organic thiols from solution onto gold," *J. Am. Chem. Soc.* **111**, 321–335 (1989).
- ⁹⁷Y. Yourdshahyan and A. M. Rappe, "Structure and energetics of alkanethiol adsorption on the Au(111) surface," *J. Chem. Phys.* **117**, 825–833 (2002).
- ⁹⁸W.-P. Hsieh, A. S. Lyons, E. Pop, P. Keblinski, and D. G. Cahill, "Pressure tuning of the thermal conductance of weak interfaces," *Phys. Rev. B* **84**, 184107 (2011).
- ⁹⁹J. K. Yang, Y. Yang, S. W. Waltermire, X. X. Wu, H. T. Zhang, T. Gutu, Y. F. Jiang, Y. F. Chen, A. A. Zinn, R. Prasher, T. T. Xu, and D. Y. Li, "Enhanced and switchable nanoscale thermal conduction due to van der Waals interfaces," *Nat. Nanotechnol.* **7**, 91–95 (2012).
- ¹⁰⁰C. M. Hargreaves, "Anomalous radiative transfer between closely-spaced bodies," *Phys. Lett. A* **30**, 491–492 (1969).

- ¹⁰¹D. Polder and M. Van Hove, "Theory of radiative heat transfer between closely spaced bodies," *Phys. Rev. B* **4**, 3303–3314 (1971).
- ¹⁰²J. J. Loomis and H. J. Maris, "Theory of heat transfer by evanescent electromagnetic waves," *Phys. Rev. B* **50**, 18517–18524 (1994).
- ¹⁰³A. I. Volokitin and B. N. Persson, "Radiative heat transfer between nanostructures," *Phys. Rev. B* **63**, 205404 (2001).
- ¹⁰⁴A. I. Volokitin and B. N. Persson, "Resonant photon tunneling enhancement of the radiative heat transfer," *Phys. Rev. B* **69**, 045417 (2004).
- ¹⁰⁵A. Kittel, W. Müller-Hirsch, J. Parisi, S.-A. Biehs, D. Reddig, and M. Holthaus, "Near-field heat transfer in a scanning thermal microscope," *Phys. Rev. Lett.* **95**, 224301 (2005).
- ¹⁰⁶K. Zhang, "Carbon nanotube thermal interface material for high-brightness light-emitting-diode cooling," *Nanotechnology* **19**, 215706 (2008).
- ¹⁰⁷G. D. Mahan, "Kapitza resistance between a metal and a nonmetal," *Phys. Rev. B* **79**, 075408 (2009).
- ¹⁰⁸P. Ben-Abdallah and K. Joulain, "Noncontact heat transfer between two metamaterials," *Phys. Rev. B* **82**, 121419 (2010).
- ¹⁰⁹S. A. Biehs, E. Rousseau, and J. J. Greffet, "A mesoscopic description of radiative heat transfer at the nanoscale," *Phys. Rev. Lett.* **105**, 234301 (2010).
- ¹¹⁰C. R. Otey, W. T. Lau, and S. H. Fan, "Thermal rectification through vacuum," *Phys. Rev. Lett.* **104**, 154301 (2010).
- ¹¹¹G. D. Mahan, "The tunneling of heat," *Appl. Phys. Lett.* **98**, 132106 (2011).
- ¹¹²A. I. Volokitin and B. N. Persson, "Near-field radiative heat transfer between closely spaced graphene and amorphous SiO₂," *Phys. Rev. B* **83**, 241407 (2011).
- ¹¹³B. N. Persson, A. I. Volokitin, and H. Ueba, "Phononic heat transfer across an interface: Thermal boundary resistance," *J. Phys. Condens. Matter* **23**, 045009 (2011).
- ¹¹⁴R. S. Ottens, V. Quetschke, S. Wise, A. A. Alemi, R. Lundock, G. Mueller, D. H. Reitze, D. B. Tanner, and B. F. Whiting, "Near-field radiative heat transfer between macroscopic planar surfaces," *Phys. Rev. Lett.* **107**, 014301 (2011).
- ¹¹⁵C. Otey and S. Fan, "Numerically exact calculation of electromagnetic heat transfer between a dielectric sphere and plate," *Phys. Rev. B* **84**, 245431 (2011).
- ¹¹⁶P. M. Norris, J. L. Smoyer, J. C. Duda, and P. E. Hopkins, "Prediction and measurement of thermal transport across interfaces between isotropic solids and graphitic materials," *J. Heat Transfer* **134**, 020910 (2012).
- ¹¹⁷K. Joulain, J.-P. Mulet, F. Marquier, R. Carminati, and J.-J. Greffet, "Surface electromagnetic waves thermally excited: Radiative heat transfer, coherence properties and Casimir forces revisited in the near field," *Surf. Sci. Rep.* **57**, 59–112 (2005).
- ¹¹⁸G. C. Arvind Narayanaswamy, "Direct computation of thermal emission from nanostructures," *Annu. Rev. Heat Transfer* **14**, 169–195 (2005).
- ¹¹⁹E. G. Cravalho, G. A. Domoto, and C. L. Tien, *Progress in Astronautics and Aeronautics Measurement of Thermal Radiation of Solids at Liquid Helium Temperatures* (Academic Press, New York, 1969), Vol. 21.
- ¹²⁰G. A. Domoto, R. F. Boehm, and C. L. Tien, "Experimental investigation of radiative transfer between metallic surfaces at cryogenic temperatures," *J. Heat Transfer* **92**, 412–417 (1970).
- ¹²¹C. Hargreaves, "Radiative transfer between closely spaced bodies," *Philips Res. Rep. Suppl.* **5**, 1–80 (1973).
- ¹²²T. Kralik, P. Hanzelka, V. Musilova, A. Srnka, and M. Zbac, "Cryogenic apparatus for study of near-field heat transfer," *Rev. Sci. Instrum.* **82**, 055106 (2011).
- ¹²³J. Xu, K. Lauger, R. Moller, K. Dransfeld, and I. H. Wilson, "Heat transfer between two metallic surfaces at small distances," *J. Appl. Phys.* **76**, 7209–7216 (1994).
- ¹²⁴L. Hu, A. Narayanaswamy, X. Chen, and G. Chen, "Near-field thermal radiation between two closely spaced glass plates exceeding Planck's blackbody radiation law," *Appl. Phys. Lett.* **92**, 133106 (2008).
- ¹²⁵A. Narayanaswamy, S. Shen, and G. Chen, "Near-field radiative heat transfer between a sphere and a substrate," *Phys. Rev. B* **78**, 115303 (2008).
- ¹²⁶S. Shen, A. Narayanaswamy, and G. Chen, "Surface phonon polaritons mediated energy transfer between nanoscale gaps," *Nano Lett.* **9**, 2909–2913 (2009).
- ¹²⁷J. R. Barnes, R. J. Stephenson, M. E. Welland, C. H. Gerber, and J. K. Gimzewski, "Photothermal spectroscopy with femtojoule sensitivity using a micromechanical device," *Nature* **372**, 79–81 (1994).
- ¹²⁸J. Varesi, J. Lai, T. Perazzo, Z. Shi, and A. Majumdar, "Photothermal measurements at picowatt resolution using uncooled micro-optomechanical sensors," *Appl. Phys. Lett.* **71**, 306–308 (1997).
- ¹²⁹S. Shen, A. Narayanaswamy, S. Goh, and G. Chen, "Thermal conductance of bimaterial microcantilevers," *Appl. Phys. Lett.* **92**, 063509 (2008).
- ¹³⁰N. Gu, K. Sasithithlu, and A. Narayanaswamy, "Near field radiative heat transfer measurement," in *Renewable Energy and the Environment* (Optical Society of America, 2011), p. JWE13.
- ¹³¹P. S. S. Shen, A. Mavrokefalos and G. Chen, "Nanoscale thermal radiation between two gold surfaces," *Appl. Phys. Lett.* **100**, 233114 (2012).
- ¹³²E. Rousseau, A. Siria, G. Jourdan, S. Volz, F. Comin, J. Chevrier, and J.-J. Greffet, "Radiative heat transfer at the nanoscale," *Nature Photon.* **3**, 514–517 (2009).
- ¹³³K. Sasithithlu and A. Narayanaswamy, "Proximity effects in radiative heat transfer," *Phys. Rev. B* **83**, 161406 (2011).
- ¹³⁴I. Dorofeyev, "Rate of heat transfer between a probing body and a sample due to electromagnetic fluctuations," *Phys. Lett. A* **372**, 1341–1347 (2008).
- ¹³⁵C. Henkel and K. Joulain, "Electromagnetic field correlations near a surface with a nonlocal optical response," *Appl. Phys. B: Lasers Opt.* **84**, 61–68 (2006).
- ¹³⁶H. B. G. Casimir, "On the attraction between two perfectly conducting plates," *Proc. K. Ned. Akad. Wet. B* **51**, 793 (1948).
- ¹³⁷I. E. Dzyaloshinskii, E. M. Lifshitz, and L. P. Pitaevskii, "General theory Of Van Der Waals' forces," *Sov. Phys. Usp.* **4**, 153 (1961).
- ¹³⁸S. Rytov, *Theory of Electric Fluctuations and Thermal Radiation* (AFRCR-TR-59-162, Electronics Research Directorate, Air Force Cambridge Research Center, Air Research and Development Command, U.S. Air Force, Bedford, MA, 1959).
- ¹³⁹U. Mohideen and A. Roy, "Precision measurement of the Casimir force from 0.1 to 0.9 μm ," *Phys. Rev. Lett.* **81**, 4549–4552 (1998).
- ¹⁴⁰K. Milton, *The Casimir Effect: Physical Manifestations of Zero-Point Energy* (World Scientific, Singapore, 2001).
- ¹⁴¹S. M. Rytov, Y. A. Kravtsov, and V. I. Tatarskii, *Principles of Statistical Radiophysics* (Springer-Verlag, Berlin, 1989).
- ¹⁴²P.-O. Chapuis, S. Volz, C. Henkel, K. Joulain, and J.-J. Greffet, "Effects of spatial dispersion in near-field radiative heat transfer between two parallel metallic surfaces," *Phys. Rev. B* **77**, 035431 (2008).
- ¹⁴³J. B. Pendry, "Radiative exchange of heat between nanostructures," *J. Phys.: Condens. Matter* **11**, 6621–6633 (1999).
- ¹⁴⁴J.-P. Mulet, K. Joulain, R. Carminati, and J.-J. Greffet, "Nanoscale radiative heat transfer between a small particle and a plane surface," *Appl. Phys. Lett.* **78**, 2931–2933 (2001).
- ¹⁴⁵A. Narayanaswamy and G. Chen, "Thermal near-field radiative transfer between two spheres," *Phys. Rev. B* **77**, 075125 (2008).
- ¹⁴⁶M. Kruger, T. Emig, and M. Kardar, "Nonequilibrium electromagnetic fluctuations: Heat transfer and interactions," *Phys. Rev. Lett.* **106**, 210404 (2011).
- ¹⁴⁷A. W. Rodriguez, O. Ilic, P. Bermel, I. Celanovic, J. D. Joannopoulos, M. Soljačić, and S. G. Johnson, "Frequency-selective near-field radiative heat transfer between photonic crystal slabs: A computational approach for arbitrary geometries and materials," *Phys. Rev. Lett.* **107**, 114302 (2011).
- ¹⁴⁸A. P. McCauley, M. T. H. Reid, M. Krüger, and S. G. Johnson, "Modeling near-field radiative heat transfer from sharp objects using a general 3d numerical scattering technique," *Phys. Rev. B* **85**, 165104 (2012).
- ¹⁴⁹A. V. Shchegrov, K. Joulain, R. Carminati, and J.-J. Greffet, "Near-field spectral effects due to electromagnetic surface excitations," *Phys. Rev. Lett.* **85**, 1548–1551 (2000).
- ¹⁵⁰S. Basu and M. Francoeur, "Near-field radiative transfer based thermal rectification using doped silicon," *Appl. Phys. Lett.* **98**, 113106 (2011).
- ¹⁵¹L. Zhu, C. R. Otey, and S. Fan, "Negative differential thermal conductance through vacuum," *Appl. Phys. Lett.* **100**, 044104 (2012).
- ¹⁵²S.-A. Biehs, F. S. S. Rosa, and P. Ben-Abdallah, "Modulation of near-field heat transfer between two gratings," *Appl. Phys. Lett.* **98**, 243102 (2011).
- ¹⁵³P. J. van Zwol, K. Joulain, P. Ben Abdallah, J. J. Greffet, and J. Chevrier, "Fast nanoscale heat-flux modulation with phase-change materials," *Phys. Rev. B* **83**, 201404 (2011).
- ¹⁵⁴A. K. Hafeli, E. Rephaeli, S. Fan, D. G. Cahill, and T. E. Tiwald, "Temperature dependence of surface phonon polaritons from a quartz grating," *J. Appl. Phys.* **110**, 043517 (2011).
- ¹⁵⁵R. J. Stoner and H. J. Maris, "Kapitza conductance and heat flow between solids at temperatures from 50 to 300 K," *Phys. Rev. B* **48**, 16373 (1993).

- ¹⁵⁶E. S. Landry and A. J. H. McGaughey, "Thermal boundary resistance predictions from molecular dynamics simulations and theoretical calculations," *Phys. Rev. B* **80**, 165304 (2009).
- ¹⁵⁷J. E. Turney, A. J. H. McGaughey, and C. H. Amon, "Assessing the applicability of quantum corrections to classical thermal conductivity predictions," *Phys. Rev. B* **79**, 224305 (2009).
- ¹⁵⁸B. Daly, H. Maris, S. Tamura, and K. Imamura, "Molecular dynamics calculation of the thermal conductivity of superlattices," *Phys. Rev. B* **66**, 024301 (2002).
- ¹⁵⁹B. Daly, H. Maris, Y. Tanaka, and S. Tamura, "Molecular dynamics calculation of the in-plane thermal conductivity of superlattices," *Phys. Rev. B* **67**, 033308 (2003).
- ¹⁶⁰Y. Chen, D. Li, J. Lukes, Z. Ni, and M. Chen, "Minimum superlattice thermal conductivity from molecular dynamics," *Phys. Rev. B* **72**, 174302 (2005).
- ¹⁶¹T. Kawamura, Y. Kangawa, and K. Kaimoto, "An investigation of thermal conductivity of nitride-semiconductor nanostructures by molecular dynamics simulation," *J. Cryst. Growth* **298**, 251–253 (2007).
- ¹⁶²S.-F. Ren, W. Cheng, and G. Chen, "Lattice dynamics investigations of phonon thermal conductivity of Si/Ge superlattices with rough interfaces," *J. Appl. Phys.* **100**, 103505 (2006).
- ¹⁶³K. Termentzidis, P. Chantrenne, J.-Y. Duquesne, and A. Saci, "Thermal conductivity of GaAs/AlAs superlattices and the puzzle of interfaces," *J. Phys. Condens. Matter* **22**, 475001 (2010).
- ¹⁶⁴K. Termentzidis, S. Merabia, P. Chantrenne, and P. Koblinski, "Cross-plane thermal conductivity of superlattices with rough interfaces using equilibrium and non-equilibrium molecular dynamics," *International J. Heat and Mass Transfer* **54**, 2014–2020 (2011).
- ¹⁶⁵E. S. Landry and A. J. McGaughey, "Effect of interfacial species mixing on phonon transport in semiconductor superlattices," *Phys. Rev. B* **79**, 075316 (2009).
- ¹⁶⁶J. E. Turney, E. S. Landry, A. J. H. McGaughey, and C. H. Amon, "Predicting phonon properties and thermal conductivity from anharmonic lattice dynamics calculations and molecular dynamics simulations," *Phys. Rev. B* **79**, 064301 (2009).
- ¹⁶⁷M. Mohr, J. Maultzsch, E. Dobardzic, S. Reich, I. Milosevic, M. Damnjanovic, A. Bosak, M. Krisch, and C. Thomsen, "The phonon dispersion of graphite by inelastic X-ray scattering," *Phys. Rev. B* **76**, 035439 (2007).
- ¹⁶⁸B. Taylor, H. J. Maris, and C. Elbaum, "Focusing of phonons in crystalline solids due to elastic anisotropy," *Phys. Rev. B* **3**, 1462–1472 (1971).
- ¹⁶⁹H. J. Maris, "Enhancement of heat pulses in crystals due to elastic anisotropy," *J. Acoust. Soc. Am.* **50**, 812–818 (1971).
- ¹⁷⁰P. Wolfe, *Imaging Phonons* (Cambridge University Press, Cambridge, 1998).
- ¹⁷¹G. Leibfried, *Encyclopedia of Physics*, edited by S. Flugge (Springer-Verlag, Berlin, 1955), Vol. 7, p. 309.
- ¹⁷²G. Deinzer, G. Birner, and D. Strauch, "Ab initio calculation of the linewidth of various phonon modes in germanium and silicon," *Phys. Rev. B* **67**, 144304 (2003).
- ¹⁷³B. A. Weinstein and G. J. Piermarini, "Raman scattering and phonon dispersion in Si and GaP at very high pressure," *Phys. Rev. B* **12**, 1172–1186 (1975).
- ¹⁷⁴M. Omini and A. Sparavigna, "Beyond the isotropic-model approximation in the theory of thermal conductivity," *Phys. Rev. B* **53**, 9064 (1996).
- ¹⁷⁵A. Sparavigna, "Role of nonpairwise interactions on phonon thermal transport," *Phys. Rev. B* **67**, 144305 (2003).
- ¹⁷⁶D. A. Broido, A. Ward, and N. Mingo, "Lattice thermal conductivity of silicon from empirical interatomic potentials," *Phys. Rev. B* **72**, 014308 (2005).
- ¹⁷⁷A. A. Maradudin and A. E. Fein, "Scattering of neutrons by an anharmonic crystal," *Phys. Rev.* **128**, 2589 (1962).
- ¹⁷⁸P. Klemens, *Solid State Physics* (Academic, New York, 1958), Vol. 7.
- ¹⁷⁹P. Carruthers, "Theory of thermal conductivity of solids at low temperatures," *Rev. Mod. Phys.* **33**, 92–138 (1961).
- ¹⁸⁰J. M. Ziman, *Electrons and Phonons: The Theory of Transport Phenomena in Solids* (Clarendon Press, 1960).
- ¹⁸¹P. G. Klemens, "Decay of high-frequency longitudinal phonons," *J. Appl. Phys.* **38**, 4573 (1967).
- ¹⁸²H. J. Maris and S. Tamura, "Anharmonic decay and the propagation of phonons in an isotopically pure crystal at low temperatures: Application to dark-matter detection," *Phys. Rev. B* **47**, 727–739 (1993).
- ¹⁸³S. Tamura, "Spontaneous decay rates of LA phonons in quasi-isotropic solids," *Phys. Rev. B* **31**, 2574 (1985).
- ¹⁸⁴A. Berke, A. Mayer, and R. Wehner, "Spontaneous decay of acoustic phonons in calcium fluoride and silicon," *Solid State Commun.* **54**, 395–397 (1985).
- ¹⁸⁵R. Orbach and L. A. Vredevoe, "The Attenuation of high frequency phonons at low temperatures," *Phys.* **1**, 91 (1964).
- ¹⁸⁶H. J. Maris, "In Elastic Scattering Of Neutrons By An Anharmonic Crystal At Low Temperatures," *Phys. Lett.* **17**, 228–230 (1964).
- ¹⁸⁷M. Lax, P. Hu, and V. Narayanamurti, "Spontaneous phonon decay selection rule: N and U processes," *Phys. Rev. B* **23**, 3095–3097 (1981).
- ¹⁸⁸A. Berke, A. P. Mayer, and R. K. Wehner, "Spontaneous decay of long-wavelength acoustic phonons," *J. Phys. C* **21**, 2305 (1988).
- ¹⁸⁹P. G. Klemens, "Anharmonic decay of optical phonons," *Phys. Rev.* **148**, 845–848 (1966).
- ¹⁹⁰C. Herring, "Role of low-energy phonons in thermal conduction," *Phys. Rev.* **95**, 954 (1954).
- ¹⁹¹S. Simons, "The absorption of very high frequency sound in dielectric solids," *Proc. Cambridge Philos. Soc.* **53**, 702 (1957).
- ¹⁹²A. A. Maradudin, A. E. Fein, and G. H. Vineyard, "Thermal expansion and phonon frequency shifts," *Phys. Status Solidi* **2**, 1493 (1962).
- ¹⁹³S. Tamura and H. J. Maris, "Temperature dependence of phonon lifetime in dielectric crystals," *Phys. Rev. B* **51**, 2857–2863 (1995).
- ¹⁹⁴S. Tamura, "Isotope scattering of dispersive phonons in Ge," *Phys. Rev. B* **27**, 858 (1983).
- ¹⁹⁵A. Akhiezer, "On the absorption of sound in solids," *J. Phys. (USSR)* **1**, 277 (1939).
- ¹⁹⁶H. J. Maris, "Phonon viscosity," *Phys. Rev.* **188**, 1303–1307 (1969).
- ¹⁹⁷H. J. Maris, *Physical Acoustics* (Academic, New York, 1971), Vol. 8, p. 279.
- ¹⁹⁸T. O. Woodruff and H. Ehrenreich, "Absorption of sound in insulators," *Phys. Rev.* **123**, 1553–1559 (1961).
- ¹⁹⁹D. A. Broido, M. Malorny, G. Birner, N. Mingo, and D. A. Stewart, "Intrinsic lattice thermal conductivity of semiconductors from first principles," *Appl. Phys. Lett.* **91**, 231922 (2007).
- ²⁰⁰A. Ward, D. S. Broido, D. A. Stewart, and G. Deinzer, "Ab-initio theory of the lattice thermal conductivity of diamond," *Phys. Rev. B* **80**, 125203 (2009).
- ²⁰¹A. Ward and D. A. Broido, "Intrinsic phonon relaxation times from first-principles studies of the thermal conductivities of Si and Ge," *Phys. Rev. B* **81**, 085205 (2010).
- ²⁰²L. Lindsay and D. A. Broido, "Three-phonon phase space and lattice thermal conductivity in semiconductors," *J. Phys. Condens. Matter* **20**, 165209 (2008).
- ²⁰³W. Li, N. Mingo, L. Lindsay, D. A. Broido, D. A. Stewart, and N. A. Katcho, "Thermal conductivity of diamond nanowires from first principles," *Phys. Rev. B* **85**, 195436 (2012).
- ²⁰⁴L. Lindsay, D. A. Broido, and N. Mingo, "Lattice thermal conductivity of single-walled carbon nanotubes: Beyond the relaxation time approximation and phonon-phonon scattering selection rules," *Phys. Rev. B* **80**, 125407 (2009).
- ²⁰⁵L. Lindsay, D. A. Broido, and N. Mingo, "Flexural phonons and thermal transport in graphene," *Phys. Rev. B* **82**, 115427 (2010).
- ²⁰⁶H. J. Maris and S. Tamura, "Heat flow in nanostructures in the Casimir regime," *Phys. Rev. B* **85**, 054304 (2012).
- ²⁰⁷A. K. McCurdy, H. J. Maris, and C. Elbaum, "Anisotropic heat conduction in cubic crystals in the boundary scattering regime," *Phys. Rev. B* **2**, 4077–4083 (1970).
- ²⁰⁸H. B. G. Casimir, "Note on the conduction of heat in crystals," *Physica* **5**, 495–500 (1938).
- ²⁰⁹K. Fuchs, "The conductivity of thin metallic films according to the electron theory of metals," *Math. Proc. Cambridge Philos. Soc.* **34**, 100–108 (1938).
- ²¹⁰R. B. Dingle, "The electrical conductivity of thin wires," *Proc. R. Soc. London, Ser. A* **201**, 545–560 (1950).
- ²¹¹R. Berman, E. L. Foster, and J. M. Ziman, "Thermal conduction in artificial sapphire crystals at low temperatures. I. Nearly perfect crystals," *Proc. R. Soc. London, Ser. A* **231**, 130–144 (1955).
- ²¹²Z. Wang and N. Mingo, "Absence of Casimir regime in two-dimensional nanoribbon phonon conduction," *Appl. Phys. Lett.* **99**, 101903 (2011).
- ²¹³A. Mavrokefalos, M. T. Pettes, F. Zhou, and L. Shi, "Four-probe measurements of the in-plane thermoelectric properties of nanofilms," *Rev. Sci. Instrum.* **78**, 034901 (2007).
- ²¹⁴J.-K. Yu, S. Mitrovic, D. Tham, J. Varghese, and J. R. Heath, "Reduction of thermal conductivity in phononic nanomesh structures," *Nat. Nanotechnol.* **5**, 718–721 (2010).

- ²¹⁵J. Tang, H.-T. Wang, D. H. Lee, M. Fardy, Z. Huo, T. P. Russell, and P. Yang, "Holey silicon as an efficient thermoelectric material," *Nano Lett.* **10**, 4279–4283 (2010).
- ²¹⁶L. Shi, D. Li, C. Yu, W. Jang, D. Kim, Z. Yao, P. Kim, and A. Majumdar, "Measuring thermal and thermoelectric properties of one-dimensional nanostructures using a microfabricated device," *J. Heat Transfer* **125**, 881–888 (2003).
- ²¹⁷A. I. Persson, Y. K. Koh, D. G. Cahill, L. Samuelson, and H. Linke, "Thermal conductance of InAs nanowire composites," *Nano Lett.* **9**, 4484–4488 (2009).
- ²¹⁸J. P. Feser and D. G. Cahill, "Probing anisotropic heat transport using time-domain thermoreflectance with offset laser spots," *Rev. Sci. Instrum.* **83**, 104901 (2012).
- ²¹⁹F. Zhou, A. L. Moore, J. Bolinsson, A. Persson, L. Froberg, M. T. Pettes, H. Kong, L. Rabenberg, P. Caroff, D. A. Stewart, N. Mingo, K. A. Dick, L. Samuelson, H. Linke, and L. Shi, "Thermal conductivity of indium arsenide nanowires with wurtzite and zinc blende phases," *Phys. Rev. B* **83**, 205416 (2011).
- ²²⁰A. Mavrokefalos, A. L. Moore, M. T. Pettes, L. Shi, W. Wang, and X. Li, "Nanowires, thermoelectric and structural characterizations of individual electrodeposited bismuth telluride," *J. Appl. Phys.* **105**, 104318 (2009).
- ²²¹L. Shi, Q. Hao, C. Yu, N. Mingo, X. Kong, and Z. Wang, "Thermal conductivities of individual tin dioxide nanobelts," *Appl. Phys. Lett.* **84**, 2638–2640 (2004).
- ²²²D. Li, Y. Wu, P. Kim, L. Shi, P. Yang, and A. Majumdar, "Thermal conductivity of individual silicon nanowires," *Appl. Phys. Lett.* **83**, 2934–2936 (2003).
- ²²³A. L. Moore, M. T. Pettes, F. Zhou, and L. Shi, "Thermal conductivity suppression in bismuth nanowires," *J. Appl. Phys.* **106**, 034310 (2009).
- ²²⁴F. Zhou, A. L. Moore, M. T. Pettes, Y. Lee, J. H. Seol, Q. L. Ye, L. Rabenberg, and L. Shi, "Effect of growth base pressure on the thermoelectric properties of indium antimonide nanowires," *J. Phys. D: Appl. Phys.* **43**, 025406 (2010).
- ²²⁵H. Kim, I. Kim, H.-j. Choi, and W. Kim, "Thermal conductivities of Si_{1-x}Ge_x nanowires with different germanium concentrations and diameters," *Appl. Phys. Lett.* **96**, 233106 (2010).
- ²²⁶N. Mingo, "Calculation of Si nanowire thermal conductivity using complete phonon dispersion relations," *Phys. Rev. B* **68**, 113308 (2003).
- ²²⁷D. Morelli, J. Heremans, and G. Slack, "Estimation of the isotope effect on the lattice thermal conductivity of group IV and group III–V semiconductors," *Phys. Rev. B* **66**, 195304 (2002).
- ²²⁸Z. Wang and N. Mingo, "Diameter dependence of SiGe nanowire thermal conductivity," *Appl. Phys. Lett.* **97**, 101903 (2010).
- ²²⁹J. S. Heron, T. Fournier, N. Mingo, and O. Bourgeois, "Mesoscopic size effects on the thermal conductance of silicon nanowire," *Nano Lett.* **9**, 1861–1865 (2009).
- ²³⁰A. I. Hochbaum, R. Chen, R. D. Delgado, W. Liang, E. C. Garnett, M. Najarian, A. Majumdar, and P. Yang, "Enhanced thermoelectric performance of rough silicon nanowires," *Nature* **451**, 163–167 (2008).
- ²³¹L. Shi, "Thermal and thermoelectric transport in nanostructures and low-dimensional systems," *Nanoscale Microscale Thermophys. Eng.* **16**, 79–116 (2012).
- ²³²K. Hippalgaonkar, B. Huang, R. Chen, K. Sawyer, P. Ercius, and A. Majumdar, "Fabrication of microdevices with integrated nanowires for investigating low-dimensional phonon transport," *Nano Lett.* **10**, 4341–4348 (2010).
- ²³³A. L. Moore, S. K. Saha, R. S. Prasher, and L. Shi, "Phonon backscattering and thermal conductivity suppression in sawtooth nanowires," *Appl. Phys. Lett.* **93**, 083112 (2008).
- ²³⁴A. Eucken, *Forsch. Geb. Ingenieurwes.* **353**, 16 (1932).
- ²³⁵S. Hyun and S. Torquato, "Optimal and manufacturable two-dimensional, Kagome-like cellular solids," *J. Mater. Res.* **17**, 137 (2002).
- ²³⁶D. G. Cahill, M. Katiyar, and J. R. Abelson, "Thermal conductivity of a-Si:H thin films," *Phys. Rev. B* **50**, 6077–6081 (1994).
- ²³⁷J. Zou and A. Balandin, "Phonon heat conduction in a semiconductor nanowire," *J. Appl. Phys.* **89**, 2932–2938 (2001).
- ²³⁸P. Martin, Z. Aksamija, E. Pop, and U. Ravaioli, "Impact of phonon-surface roughness scattering on thermal conductivity of thin Si nanowires," *Phys. Rev. Lett.* **102**, 125503 (2009).
- ²³⁹J. Sadhu and S. Sinha, "Room-temperature phonon boundary scattering below the Casimir limit," *Phys. Rev. B* **84**, 115450 (2011).
- ²⁴⁰J. Carrete, L. J. Gallego, L. M. Varela, and N. Mingo, "Surface roughness and thermal conductivity of semiconductor nanowires: Going below the Casimir limit," *Phys. Rev. B* **84**, 075403 (2011).
- ²⁴¹Y. He and G. Galli, "Microscopic origin of the reduced thermal conductivity of silicon nanowires," *Phys. Rev. Lett.* **108**, 215901 (2012).
- ²⁴²T. Klitsner and R. O. Pohl, "Phonon scattering at silicon crystal surfaces," *Phys. Rev. B* **36**, 6551–6565 (1987).
- ²⁴³A. I. Hochbaum, D. Gargas, Y. J. Hwang, and P. Yang, "Single crystalline mesoporous silicon nanowires," *Nano Lett.* **9**, 3550–3554 (2009).
- ²⁴⁴G. S. Oehrlein, "Dry etching damage of silicon: A review," *Mater. Sci. Eng., B* **4**, 441–450 (1989).
- ²⁴⁵J. Lim, K. Hippalgaonkar, S. C. Andrews, A. Majumdar, and P. Yang, "Quantifying surface roughness effects on phonon transport in silicon nanowires," *Nano Lett.* **12**, 2475–2482 (2012).
- ²⁴⁶J. D. Bernal, "The structure of graphite," *Proc. R. Soc. London, Ser. A* **106**, 749–773 (1924).
- ²⁴⁷A. H. Castro Neto, F. Guinea, N. M. R. Peres, K. S. Novoselov, and A. K. Geim, "The electronic properties of graphene," *Rev. Mod. Phys.* **81**, 109–161 (2009).
- ²⁴⁸R. Nicklow, N. Wakabayashi, and H. G. Smith, "Lattice dynamics of pyrolytic graphite," *Phys. Rev. B* **5**, 4951–4962 (1972).
- ²⁴⁹R. Al-Jishi and G. Dresselhaus, "Lattice-dynamical model for graphite," *Phys. Rev. B* **26**, 4514–4522 (1982).
- ²⁵⁰D. L. Nika, E. P. Pokatilov, A. S. Askerov, and A. A. Balandin, "Phonon thermal conduction in graphene: Role of umklapp and edge roughness scattering," *Phys. Rev. B* **79**, 155413 (2009).
- ²⁵¹L. Lindsay, D. A. Broido, and N. Mingo, "Flexural phonons and thermal transport in multilayer graphene and graphite," *Phys. Rev. B* **83**, 235428 (2011).
- ²⁵²L. Lindsay, D. A. Broido, and N. Mingo, "Diameter dependence of carbon nanotube thermal conductivity and extension to the graphene limit," *Phys. Rev. B* **82**, 161402 (2010).
- ²⁵³P. G. Klemens, "Theory of thermal conduction in thin ceramic films," *Int. J. Thermophys.* **22**, 265–275 (2001).
- ²⁵⁴N. Mingo and D. Broido, "Length dependence of carbon nanotube thermal conductivity and the 'Problem of long waves,'" *Nano Lett.* **5**, 1221–1225 (2005).
- ²⁵⁵S. Ghosh, W. Bao, D. L. Nika, S. Subrina, E. P. Pokatilov, C. N. Lau, and A. A. Balandin, "Dimensional crossover of thermal transport in few-layer graphene," *Nature Mater.* **9**, 555–558 (2010).
- ²⁵⁶C. Faugeras, B. Faugeras, M. Orlita, M. Potemski, R. R. Nair, and A. K. Geim, "Thermal conductivity of graphene in Corbino Membrane Geometry," *ACS Nano* **4**, 1889–1892 (2010).
- ²⁵⁷W. Cai, A. L. Moore, Y. Zhu, X. Li, S. Chen, L. Shi, and R. S. Ruoff, "Thermal transport in suspended and supported monolayer graphene grown by chemical vapor deposition," *Nano Lett.* **10**, 1645–1651 (2010).
- ²⁵⁸S. Chen, A. L. Moore, W. Cai, J. W. Suk, J. An, C. Mishra, C. Amos, C. W. Magnuson, J. Kang, L. Shi, and R. S. Ruoff, "Raman measurements of thermal transport in suspended monolayer graphene of variable sizes in vacuum and gaseous environments," *ACS Nano* **5**, 321–328 (2011).
- ²⁵⁹J.-U. Lee, D. Yoon, H. Kim, S. W. Lee, and H. Cheong, "Thermal conductivity of suspended pristine graphene measured by Raman spectroscopy," *Phys. Rev. B* **83**, 081419 (2011).
- ²⁶⁰J. H. Seol, I. Jo, A. L. Moore, L. Lindsay, Z. H. Aitken, M. T. Pettes, X. Li, Z. Yao, R. Huang, D. Broido, N. Mingo, R. S. Ruoff, and L. Shi, "Two-dimensional phonon transport in supported graphene," *Science* **328**, 213–216 (2010).
- ²⁶¹W. Jang, Z. Chen, W. Bao, C. N. Lau, and C. Dames, "Thickness-dependent thermal conductivity of encased graphene and ultrathin graphite," *Nano Lett.* **10**, 3909–3913 (2010).
- ²⁶²M. M. Sadeghi, I. Jo, and L. Shi, "Phonon-Interface Scattering in Multilayered Graphene on an Amorphous Support," *Proceedings of National Academy of Sciences* **110**, 16321–16326 (2013).
- ²⁶³Y. S. Touloukian and E. H. Buyco, *Thermophysical Properties of Matter* (IFI/Plenum, New York, 1970).
- ²⁶⁴T. Aizawa, R. Souda, S. Otani, Y. Ishizawa, and C. Oshima, "Bond softening in monolayer graphite formed on transition-metal carbide surfaces," *Phys. Rev. B* **42**, 11469–11478 (1990).
- ²⁶⁵D. Farias, K. Rieder, A. Shikin, V. Adamchuk, T. Tanaka, and C. Oshima, "Modification of the surface phonon dispersion of a graphite monolayer adsorbed on Ni (111) caused by intercalation of Yb, Cu and Ag," *Surf. Sci.* **454**, 437–441 (2000).
- ²⁶⁶T. N. Theis and P. M. Solomon, "In quest of the 'next switch': Prospects for greatly reduced power dissipation in a successor to the silicon field-effect transistor," *Proc. IEEE* **98**, 2005–2014 (2010).

- ²⁶⁷E. Pop, S. Sinha, and K. E. Goodson, "Heat generation and transport in nanometer-scale transistors," *Proc. IEEE* **94**, 1587–1601 (2006).
- ²⁶⁸H.-S. P. Wong, S. Raoux, S. Kim, J. L. Liang, J. P. Reifenberg, B. Rajendran, M. Asheghi, and K. E. Goodson, "Phase change memory," *Proc. IEEE* **98**, 2201–2227 (2010).
- ²⁶⁹G. W. Burr, M. J. Breitwisch, M. Franceschini, D. Garetto, K. Gopalakrishnan, B. Jackson, B. Kurdi, C. Lam, L. A. Lastras, A. Padilla, B. Rajendran, S. Raoux, and R. S. Shenoy, "Phase change memory technology," *J. Vac. Sci. Technol. B* **28**, 223–262 (2010).
- ²⁷⁰H. Altug, D. Eglund, and J. Vuckovic, "Ultrafast photonic crystal nanocavity laser," *Nat. Phys.* **2**, 484 (2006).
- ²⁷¹B. Concoran, C. Monat, C. Grillet, D. Moss, B. Eggleton, T. White, L. O'Faolain, and T. Krauss, "Green light emission in silicon through slow-light enhanced third-harmonic generation in photonic-crystal waveguides," *Nat. Photonics* **3**, 206–210 (2009).
- ²⁷²A. Sarua, H. Ji, K. Hilton, D. Wallis, M. Uren, T. Martin, and M. Kuball, "Thermal boundary resistance between GaN and substrate in AlGaIn/GaN electronic devices," *IEEE Trans. Electron Devices* **54**, 3152–3158 (2007).
- ²⁷³M. Alomari, A. Dussaigne, D. Martin, N. Grandjean, C. Gaquiere, and E. Kohn, "AlGaIn/GaN HEMT on (111) single crystalline diamond," *Electron. Lett.* **46**, 299–301 (2010).
- ²⁷⁴J. Kuzmik, S. Bychikhin, D. Pogany, E. Pichonat, O. Lancry, C. Gaquiere, G. Tsiakouras, G. Deligeorgis, and A. Georgakilas, "Thermal characterization of MBE-grown GaN/AlGaIn/GaN device on single crystalline diamond," *J. Appl. Phys.* **109**, 086106 (2011).
- ²⁷⁵E. Pop, "Energy dissipation and transport in nanoscale devices," *Nano Res.* **3**, 147–169 (2010).
- ²⁷⁶M. Pedram and S. Nazarian, "Thermal modeling, analysis, and management in VLSI circuits: Principles and methods," *Proc. IEEE* **94**, 1487–1501 (2006).
- ²⁷⁷D. A. B. Miller, "Device requirements for optical interconnects to silicon chips," *Proc. IEEE* **97**, 1166–1185 (2009).
- ²⁷⁸D. Kuzum, R. G. D. Jeyasingh, B. Lee, and H.-S. P. Wong, "Nanoelectronic programmable synapses based on phase change materials for brain-inspired computing," *Nano Lett.* **12**, 2179 (2012).
- ²⁷⁹C. Ahn, B. Lee, R. G. D. Jeyasingh, M. Asheghi, G. A. M. Hurkx, K. E. Goodson, and H.-S. P. Wong, "Crystallization properties and their drift dependence in phase-change memory studied with a micro-thermal stage," *J. Appl. Phys.* **110**, 114520 (2011).
- ²⁸⁰J. Lee, T. Kodama, Y. Won, M. Asheghi, and K. E. Goodson, "Phase purity and thermoelectric properties of $\text{Ge}_2\text{Sb}_2\text{Te}_5$ films down to 25 nm thickness," *J. Appl. Phys.* **112**, 014902 (2012).
- ²⁸¹Y. Zhang, J. Feng, Y. Zhang, Z. Zhang, Y. Lin, T. Tang, B. Cai, and B. Chen, "Multi-bit storage in reset process of phase change access memory (PRAM)," *Phys. Status Solidi (RRL)* **1**, R28–R30 (2007).
- ²⁸²G. Oh, Y. Park, J. Lee, D. Im, J. Bae, D. Kim, D. Ahn, H. Horii, S. Park, H. Yoon, I. Park, Y. Ko, U.-I. Chung, and J. Moon, "Parallel multi-confined (PMC) cell technology for high density MLC PRAM," *Dig. Tech. Pap. - Symp. VLSI Technol.* **2009**, 220–221.
- ²⁸³D. Kuzum, R. G. D. Jeyasingh, and H.-S. P. Wong, "Energy efficient programming of nanoelectronic synaptic devices for large-scale implementation of associative and temporal sequence learning," *Tech. Dig. - Int. Electron Devices Meet.* **2011**, 30.3.1–30.3.4.
- ²⁸⁴E. Bozorg-Grayeli, J. P. Reifenberg, M. Asheghi, H.-S. P. Wong, and K. E. Goodson, "Thermal transport in phase change memory materials," *Annu. Rev. Heat Transfer* **15**, 397–428 (2012).
- ²⁸⁵H.-K. Lyoo, D. G. Cahill, B.-S. Lee, J. R. Abelson, M.-H. Kwon, K.-B. Kim, S. G. Bishop, and B. Ki Cheong, "Thermal conductivity of phase-change material $\text{Ge}_2\text{Sb}_2\text{Te}_5$," *Appl. Phys. Lett.* **89**, 151904 (2006).
- ²⁸⁶J. P. Reifenberg, M. A. Panzer, S. Kim, A. M. Gibby, Y. Zhang, S. Wong, H.-S. P. Wong, E. Pop, and K. E. Goodson, "Thickness and stoichiometry dependence of the thermal conductivity of GeSbTe films," *Appl. Phys. Lett.* **91**, 111904 (2007).
- ²⁸⁷W. P. Risk, C. T. Rettner, and S. Raoux, "Thermal conductivities and phase transition temperatures of various phase-change materials measured by the 3 omega method," *Appl. Phys. Lett.* **94**, 101906 (2009).
- ²⁸⁸J. Lee, Z. Li, J. P. Reifenberg, S. Lee, R. Sinclair, M. Asheghi, and K. E. Goodson, "Thermal conductivity anisotropy and grain structure in $\text{Ge}_2\text{Sb}_2\text{Te}_5$ films," *J. Appl. Phys.* **109**, 084902 (2011).
- ²⁸⁹I. R. Chen and E. Pop, "Compact thermal model for vertical nanowire phase-change memory cells," *IEEE Trans. Electron Devices* **56**, 1523–1528 (2009).
- ²⁹⁰E.-K. Kim, S.-I. Kwun, S.-M. Lee, H. Seo, and J.-G. Yoon, "Thermal boundary resistance at $\text{Ge}_2\text{Sb}_2\text{Te}_5/\text{ZnS}:\text{SiO}_2$ interface," *Appl. Phys. Lett.* **76**, 3864–3866 (2000).
- ²⁹¹J. Reifenberg, D. Kencke, and K. Goodson, "The impact of thermal boundary resistance in phase-change memory devices," *IEEE Electron Device Letters* **29**, 1112–1114 (2008).
- ²⁹²J. Reifenberg, K.-W. Chang, M. Panzer, S. Kim, J. Rowlette, M. Asheghi, H.-S. P. Wong, and K. Goodson, "Thermal boundary resistance measurements for phase-change memory devices," *IEEE Electron Device Letters* **31**, 56–58 (2010).
- ²⁹³J. Lee, M. Asheghi, and K. Goodson, "Impact of thermoelectric phenomena on phase-change memory performance metrics and scaling," *Nanotechnology* **23**, 205201 (2012).
- ²⁹⁴T. Kato and K. Tanaka, "Electronic properties of amorphous and crystalline $\text{Ge}_2\text{Sb}_2\text{Te}_5$ Films," *Jpn. J. Appl. Phys. Part 1* **44**, 7340–7344 (2005).
- ²⁹⁵F. Xiong, A. Liao, and E. Pop, "Inducing chalcogenide phase change with ultra-narrow carbon nanotube heaters," *Appl. Phys. Lett.* **95**, 243103 (2009).
- ²⁹⁶F. Xiong, A. D. Liao, D. Estrada, and E. Pop, "Low-power switching of phase-change materials with carbon nanotube electrodes," *Science* **332**, 568–570 (2011).
- ²⁹⁷J. Liang, R. G. D. Jeyasingh, H. Y. Chen, and H.-S. P. Wong, "An ultra-low reset current cross-point phase change memory with carbon nanotube electrodes," *IEEE Trans. Electron Devices* **59**, 1155–1163 (2012).
- ²⁹⁸U. Russo, D. Ielmini, A. Redaelli, and A. L. Lacaita, "Modeling of programming and read performance in phase-change memories—part II: Program disturb and mixed-scaling approach," *IEEE Trans. Electron Devices* **55**, 515–522 (2008).
- ²⁹⁹S. Kim, B. Lee, M. Asheghi, G. A. M. Hurkx, J. Reifenberg, K. Goodson, and H.-S. P. Wong, "Thermal disturbance and its impact on reliability of phase-change memory studied by the micro-thermal stage," *IEEE Int. Reliab. Phys. Symp. Proc.* **2010**, 99–103.
- ³⁰⁰L. Pan and D. B. Bogy, "Data storage: Heat-assisted magnetic recording," *Nat. Photonics* **3**, 189–190 (2009).
- ³⁰¹R. Fernandez, D. Teweldebrhan, C. Zhang, A. Balandin, and S. Khizroev, "A comparative analysis of Ag and Cu heat sink layers in $\text{L}_{10}\text{-FePt}$ films for heat-assisted magnetic recording," *J. Appl. Phys.* **109**, 07B763 (2011).
- ³⁰²D.-S. Lim, M.-H. Shin, H.-S. Oh, and Y.-J. Kim, "Opto-thermal analysis of novel heat assisted magnetic recording media based on surface plasmon enhancement," *IEEE Trans. Magn.* **45**, 3844–3847 (2009).
- ³⁰³Y. Ma, L. Gonzaga, C. An, and B. Liu, "Effect of laser heating duration on lubricant depletion in heat assisted magnetic recording," *IEEE Trans. Magn.* **47**, 3445–3448 (2011).
- ³⁰⁴R. Chau, B. Doyle, S. Datta, J. Kavalieros, and K. Zhang, "Integrated nanoelectronics for the future," *Nature Mater.* **6**, 810–812 (2007).
- ³⁰⁵J. A. del Alamo, "Nanometre-scale electronics with III-V compound semiconductors," *Nature* **479**, 317–323 (2011).
- ³⁰⁶P. Avouris and R. Martel, "Progress in carbon nanotube electronics and photonics," *MRS Bull.* **35**, 306–313 (2010).
- ³⁰⁷K. Saraswat, C. O. Chui, T. Krishnamohan, D. Kim, A. Nayfeh, and A. Pethe, "High performance germanium MOSFETs," *Mater. Sci. Eng., B* **135**, 242–249 (2006).
- ³⁰⁸N. Collaert, A. De Keersgieter, A. Dixit, I. Ferain, L. S. Lai, D. Lenoble, A. Mercha, A. Nackaerts, B. J. Pawlak, R. Rooyackers, T. Schulz, K. T. San, N. J. Son, M. J. H. Van Dal, P. Verheyen, K. von Armin, L. Witters, K. De Meyer, S. Biesemans, and M. Jurczak, "Multi-gate devices for the 32 nm technology node and beyond," *Solid-State Electron.* **52**, 1291–1296 (2008).
- ³⁰⁹E. Pop, R. W. Dutton, and K. E. Goodson, "Monte Carlo simulation of Joule heating in bulk and strained silicon," *Appl. Phys. Lett.* **86**, 082101-3 (2005).
- ³¹⁰V. Perebeinos and P. Avouris, "Inelastic scattering and current saturation in graphene," *Phys. Rev. B* **81**, 195442 (2010).
- ³¹¹K. Kang, D. Abdula, D. G. Cahill, and M. Shim, "Lifetimes of optical phonons in graphene and graphite by time-resolved incoherent anti-Stokes Raman scattering," *Phys. Rev. B* **81**, 165405 (2010).
- ³¹²Z.-Y. Ong, E. Pop, and J. Shiomi, "Reduction of phonon lifetimes and thermal conductivity of a carbon nanotube on amorphous silica," *Phys. Rev. B* **84**, 165418 (2011).
- ³¹³Z.-Y. Ong and E. Pop, "Molecular dynamics simulation of thermal boundary conductance between carbon nanotubes and SiO_2 ," *Phys. Rev. B* **81**, 155408 (2010).

- ³¹⁴D. K. Schroder and J. A. Babcock, "Negative bias temperature instability: Road to cross in deep submicron silicon semiconductor manufacturing," *J. Appl. Phys.* **94**, 1–18 (2003).
- ³¹⁵W. Liu, K. Etesam-Yazdani, R. Hussin, and M. Asheghi, "Modeling and data for thermal conductivity of ultrathin single-crystal SOI layers at high temperature," *IEEE Trans. Electron Devices* **53**, 1868–1876 (2006).
- ³¹⁶S. Sinha, E. Pop, R. W. Dutton, and K. E. Goodson, "Non-equilibrium phonon distributions in sub-100 nm silicon transistors," *J. Heat Transfer* **128**, 638–647 (2006).
- ³¹⁷M.-H. Bae, S. Islam, V. E. Dorgan, and E. Pop, "Scaling of high-field transport and localized heating in graphene transistors," *ACS Nano* **5**, 7936–7944 (2011).
- ³¹⁸A. D. Liao, J. Z. Wu, X. Wang, K. Tahy, D. Jena, H. Dai, and E. Pop, "Thermally limited current carrying ability of graphene nanoribbons," *Phys. Rev. Lett.* **106**, 256801 (2011).
- ³¹⁹A. Behnam, A. S. Lyons, M.-H. Bae, E. K. Chow, S. Islam, C. M. Neumann, and E. Pop, "Transport in nanoribbon interconnects obtained from graphene grown by chemical vapor deposition," *Nano Lett.* **12**, 4424–4430 (2012).
- ³²⁰M.-H. Bae, Z. Li, Z. Aksamija, P. Martin, F. Xiong, Z.-Y. Ong, I. Knezevic, and E. Pop, "Ballistic to diffusive crossover of heat flow in graphene ribbons," *Nat. Commun.* **4**, 1734 (2013).
- ³²¹S. Islam, Z. Li, V. Dorgan, M.-H. Bae, and E. Pop, "Role of Joule heating on current saturation and transient behavior of graphene transistors," *IEEE Electron Device Lett.* **34**, 166–168 (2013).
- ³²²E. Pop, "The role of electrical and thermal contact resistance for Joule breakdown of single-wall carbon nanotubes," *Nanotechnology* **19**, 295202 (2008).
- ³²³A. Liao, R. Alizadegan, Z.-Y. Ong, S. Dutta, F. Xiong, K. J. Hsia, and E. Pop, "Thermal dissipation and variability in electrical breakdown of carbon nanotube devices," *Phys. Rev. B* **82**, 205406 (2010).
- ³²⁴D. Vasilevski, K. Raleva, and S. M. Goodnick, "Self-heating effects in nanoscale FD SOI devices: The role of the substrate, boundary conditions at various interfaces, and the dielectric material type for the box," *IEEE Trans. Electron Devices* **56**, 3064–3071 (2009).
- ³²⁵T. Takahashi, N. Beppu, K. Chen, S. Oda, and K. Uchida, "Thermal-aware device design of nanoscale bulk/SOI FinFETs: Suppression of operation temperature and its variability," *Tech. Dig. - Int. Electron Devices Meet.* **2011**, 34.6.1–34.6.4.
- ³²⁶T. Sadi, R. Kelsall, N. Pilgrim, J.-L. Thobel, and F. Dessenne, "Monte Carlo study of self-heating in nanoscale devices," *J. Comput. Electron.* **11**, 118–128 (2012).
- ³²⁷W. Steinhögl, G. Schindler, G. Steinlesberger, M. Traving, and M. Engelhardt, "Comprehensive study of the resistivity of copper wires with lateral dimensions of 100 nm and smaller," *J. Appl. Phys.* **97**, 023706-7 (2005).
- ³²⁸R. L. Graham, G. B. Alers, T. Mountsier, N. Shamma, S. Dhuey, S. Cabrini, R. H. Geiss, D. T. Read, and S. Peddetti, "Resistivity dominated by surface scattering in sub-50 nm Cu wires," *Appl. Phys. Lett.* **96**, 042116-3 (2010).
- ³²⁹S. Yoneoka, J. Lee, M. Liger, G. Yama, T. Kodama, M. Gunji, J. Provine, R. T. Howe, K. E. Goodson, and T. W. Kenny, "Electrical and thermal conduction in atomic layer deposition nanobridges down to 7 nm thickness," *Nano Lett.* **12**, 683–686 (2012).
- ³³⁰F. Völklein, H. Reith, T. W. Cornelius, M. Rauber, and R. Neumann, "The experimental investigation of thermal conductivity and the Wiedemann-Franz law for single metallic nanowires," *Nanotechnology* **20**, 325706 (2009).
- ³³¹H. Sevinçli and G. Cuniberti, "Enhanced thermoelectric figure of merit in edge-disordered zigzag graphene nanoribbons," *Phys. Rev. B* **81**, 113401 (2010).
- ³³²J. Faist, F. Capasso, D. L. Sivco, C. Sirtori, A. L. Hutchinson, and A. Y. Cho, "Quantum cascade laser," *Science* **264**, 553–556 (1994).
- ³³³B. Williams, "Terahertz quantum-cascade lasers," *Nat. Photonics* **1**, 517–525 (2007).
- ³³⁴M. Razeghi, S. Slivken, Y. Bai, B. Gokden, and S. R. Darvish, "High power quantum cascade lasers," *New J. Phys.* **11**, 125017 (2009).
- ³³⁵K. Goodson, K. Kurabayashi, and R. Pease, "Improved heat sinking for laser-diode arrays using microchannels in CVD diamond," *IEEE Trans. Compon., Packag. Manuf. Technol., Part B* **20**, 104–109 (1997).
- ³³⁶J. Cho, E. Bozorg-Grayeli, D. Altman, M. Asheghi, and K. Goodson, "Low thermal resistances at GaN-SiC interfaces for HEMT technology," *IEEE Electron Device Letters* **33**, 378–380 (2012).
- ³³⁷J. Cho, Z. Li, E. Bozorg-Grayeli, T. Kodama, D. Francis, F. Ejeckam, F. Faili, M. Asheghi, and K. E. Goodson, "Thermal characterization of GaN-on-diamond substrates for HEMT applications," *IEEE Trans. Compon. Packag. Manuf. Technol.* **3**, 79–85 (2013).
- ³³⁸N. Faleev and I. Levin, "Strain and crystal defects in thin AlN/GaN structures on (0001) SiC," *J. Appl. Phys.* **107**, 113529 (2010).
- ³³⁹K. E. Goodson, "Thermal conduction in nonhomogeneous CVD diamond layers in electronic microstructures," *J. Heat Transfer* **118**, 279–286 (1996).
- ³⁴⁰K. E. Goodson, O. W. Kading, M. Rosler, and R. Zachai, "Experimental investigation of thermal conduction normal to diamond-silicon boundaries," *J. Appl. Phys.* **77**, 1385–1392 (1995).
- ³⁴¹M. Touzelbaev and K. Goodson, "Impact of nucleation density on thermal resistance near diamond-substrate boundaries," *J. Thermophys. Heat Transfer* **11**, 506–512 (1997).
- ³⁴²D. R. Clarke and S. R. Phillpot, "Thermal barrier coating materials," *Mater. Today* **8**, 22–29 (2005).
- ³⁴³D. R. Olander, *Fundamental Aspects of Nuclear Fuel Elements* (NTIS US Department of Commerce, 1976).
- ³⁴⁴C. Kittel, "Interpretation of the thermal conductivity of glasses," *Phys. Rev.* **75**, 972–974 (1949).
- ³⁴⁵P. G. Klemens, *Solid State Physics* (Academic Press, New York, 1958), Vol. 7, Chap. Klemens55, pp. 1–98.
- ³⁴⁶P. G. Klemens, "The scattering of low-frequency lattice waves by static imperfections," *Proc. Phys. Soc., London, Sect. A* **68**, 1113–1128 (1955).
- ³⁴⁷J. Callaway, "Model for lattice thermal conductivity at low temperatures," *Phys. Rev.* **113**, 1046–1051 (1959).
- ³⁴⁸J. Callaway and H. C. von Baeyer, "Effect of point imperfections on lattice thermal conductivity," *Phys. Rev.* **120**, 1149–1154 (1960).
- ³⁴⁹M. Roufosse and P. G. Klemens, "Lattice thermal conductivity of minerals at high temperatures," *J. Geophys. Res.* **79**, 703–705, doi:10.1029/JB079i005p00703 (1974).
- ³⁵⁰M. Roufosse and P. G. Klemens, "Thermal conductivity of complex dielectric crystals," *Phys. Rev. B* **7**, 5379 (1973).
- ³⁵¹A. Mathiessen, *Ann. Phys.* **7**, 892 (1864).
- ³⁵²A. Mathiessen, *Ann. Phys.* **7**, 761 (1864).
- ³⁵³L. H. Wei, P. K. Kuo, R. L. Thomas, T. R. Anthony, and W. F. Banholzer, "Thermal-conductivity of isotopically modified single-crystal diamond," *Phys. Rev. Lett.* **70**, 3764–3767 (1993).
- ³⁵⁴T. Watanabe, S. G. Srivilliputhur, P. K. Schelling, J. S. Tulenko, S. B. Sinnott, and S. R. Phillpot, "Thermal transport in off-stoichiometric uranium dioxide by atomic level simulation," *J. Am. Ceram. Soc.* **92**, 850–856 (2009).
- ³⁵⁵Z. Qu, T. D. Sparks, W. Pan, and D. R. Clarke, "Thermal conductivity of the gadolinium calcium silicate apatites: Effect of different point defect types," *Acta Mater.* **59**, 3841 (2011).
- ³⁵⁶H.-S. Yang, G.-R. Bai, L. J. Thompson, and J. A. Eastman, "Interfacial thermal resistance in nanocrystalline yttria stabilized zirconia," *Acta Mater.* **50**, 2309–2317 (2002).
- ³⁵⁷D. Zhu and R. A. Miller, "Thermal conductivity and elastic modulus evolution of thermal barrier coatings under high heat flux conditions," *J. Therm. Spray Technol.* **9**, 175–180 (2000).
- ³⁵⁸D. Zhu, R. A. Miller, B. A. Nagaraj, and R. W. Bruce, "Thermal conductivity of EB-PVD thermal barrier coatings evaluated by a steady state laser heat flux technique," *Surf. Coat. Technol.* **138**, 1–8 (2001).
- ³⁵⁹C. Mercer, J. R. Williams, D. R. Clarke, and A. G. Evans, "On a ferroelastic mechanism governing the toughness of metastable tetragonal-prime (t) yttria-stabilized zirconia," *Proc. R. Soc. London, Ser. A* **463**, 1393 (2007).
- ³⁶⁰P. K. Schelling and S. R. Phillpot, "Mechanism of thermal transport in zirconia and yttria-stabilized zirconia by molecular-dynamics simulation," *J. Am. Ceram. Soc.* **84**, 2997–3007 (2001).
- ³⁶¹P. B. Allen, J. L. Feldman, J. Fabian, and F. Wooten, "Diffusons, locons, propagons: Character of atomic vibrations in glasses," *Philos. Mag. B* **79**, 1715–1731 (1999).
- ³⁶²P. G. Lucuta, H. Matzke, and I. J. Hastings, "A pragmatic approach to modeling thermal conductivity of irradiated UO₂ fuel: Review and recommendations," *J. Nucl. Mater.* **232**, 166–180 (1996).
- ³⁶³I. L. F. Ray and H. Matzke, "Observation of a high burnup rim-type structure in an advanced plutonium-uranium carbide fuel," *J. Nucl. Mater.* **250**, 242 (1997).
- ³⁶⁴H. Matzke, "On the rim effect in high burnup UO₂ LWR fuels," *J. Nucl. Mater.* **189**, 141 (1992).
- ³⁶⁵C. Ronchi, M. Sheindlin, D. Saticu, and M. Kinoshita, "Effect of burn-up on the thermal conductivity of uranium dioxide up to 100,000 Mwd t⁻¹," *J. Nucl. Mater.* **327**, 58–76 (2005).

- ³⁶⁶Y. Wang, K. Fujinami, R. Zhang, C. Wan, Y. Ba, and K. Koumoto, "Interfacial thermal resistance and thermal conductivity in nanograined SrTiO_3 ," *Appl. Phys. Express* **3**, 031101 (2010).
- ³⁶⁷A. Limarga and D. R. Clarke, "The grain size and temperature dependence of the thermal conductivity of polycrystalline, tetragonal yttria-stabilized zirconia," *Appl. Phys. Lett.* **98**, 211906 (2011).
- ³⁶⁸L. Braginsky, V. Shklover, H. Hofman, and P. Bowen, "High-temperature thermal conductivity of porous Al_2O_3 nanostructures," *Phys. Rev. B* **70**, 134201 (2004).
- ³⁶⁹B. Aurivillius, "Mixed bismuth oxides with layer lattices. II. Structure of $\text{Bi}_4\text{Ti}_3\text{O}_{12}$," *Ark. Kemi* **1**, 499–512 (1950).
- ³⁷⁰S. Ruddlesden and P. Popper, "The compound $\text{Sr}_3\text{Ti}_2\text{O}_7$ and its structure," *Acta Crystallogr.* **11**, 54 (1958).
- ³⁷¹A. Chernatynskiy, R. W. Grimes, M. A. Zurbuchen, D. R. Clarke, and S. R. Phillpot, "Crossover in thermal transport properties of natural, perovskite-structured superlattices," *Appl. Phys. Lett.* **95**, 161906 (2009).
- ³⁷²M. R. Winter and D. R. Clarke, "Oxide materials with low thermal conductivity," *J. Am. Ceram. Soc.* **90**, 533–540 (2007).
- ³⁷³Y. Shen, R. M. Leckie, C. G. Levi, and D. R. Clarke, "Low thermal conductivity without oxygen vacancies in equimolar $\text{YO}_{1.5}+\text{TaO}_{2.5}$ - And $\text{YbO}_{1.5}+\text{TaO}_{2.5}$ -stabilized tetragonal zirconia ceramics," *Acta Mater.* **58**, 4424–4431 (2010).
- ³⁷⁴R. Vassen, X. Q. Cao, F. Tietz, D. Basu, and D. Stover, "Zirconates as new materials for thermal barrier coatings," *J. Am. Ceram. Soc.* **83**, 2023 (2000).
- ³⁷⁵W. Ma, D. Mack, J. Malzbender, R. Vassen, and D. Stover, " Yb_2O_3 and Gd_2O_3 doped strontium zirconate for thermal barrier coatings," *J. Eur. Ceram. Soc.* **28**, 3071–3081 (2008).
- ³⁷⁶X. Q. Cao, R. Vassen, F. Tietz, and D. Stover, "New double-ceramic-layer thermal barrier coatings based on zirconia rare earth composite oxides," *J. Eur. Ceram. Soc.* **26**, 247–251 (2006).
- ³⁷⁷C. L. Wan, Z. Qu, A. B. Du, and W. Pan, "Order disorder transition and unconventional thermal conductivities of the $(\text{Sm}_{1-x}\text{Yb}_x)_2\text{Zr}_2\text{O}_7$ series," *J. Am. Ceram. Soc.* **94**, 592 (2011).
- ³⁷⁸H. S. Zhang, S. R. Liao, X. D. Dang, S. K. Guan, and Z. Zhang, "Preparation and thermal conductivities of $\text{Gd}_2\text{Ce}_2\text{O}_7$ and $(\text{Gd}_{0.9}\text{Ca}_{0.1})_2\text{Ce}_2\text{O}_{6.9}$ ceramics for thermal barrier coatings," *J. Alloys Compd.* **509**, 1226–1230 (2011).
- ³⁷⁹B. Liu, J. Y. Wang, F. Z. Li, and Y. C. Zhou, "Theoretical elastic stiffness, structural stability and thermal conductivity of $\text{La}_2\text{T}_2\text{O}_7$ ($\text{T} = \text{Ge}, \text{Ti}, \text{Sn}, \text{Zr}, \text{Hf}$) pyrochlore," *Acta Mater.* **58**, 4369–4377 (2010).
- ³⁸⁰P. K. Schelling, "Thermal conductivity of A-site doped pyrochlore oxides studied by molecular-dynamics simulation," *Comput. Mater. Sci.* **48**, 336 (2010).
- ³⁸¹C. L. Wan, W. Zhang, Y. F. Zhang, Z. Qu, A. B. Du, R. F. Wu, and W. Pan, "Glasslike thermal conductivity in ytterbium doped lanthanum zirconate pyrochlore," *Acta Mater.* **58**, 6166–6172 (2010).
- ³⁸²Y. Shen, D. R. Clarke, and P. A. Fuierer, "Anisotropic thermal conductivity of the aurivillius phase, bismuth titanate ($\text{Bi}_4\text{Ti}_3\text{O}_{12}$): A natural nanostructured superlattice," *Appl. Phys. Lett.* **93**, 102907 (2008).
- ³⁸³T. D. Sparks, P. A. Fuierer, and D. R. Clarke, "Anisotropic thermal diffusivity and conductivity of La-doped strontium niobate, $\text{Sr}_2\text{Nb}_2\text{O}_7$," *J. Am. Ceram. Soc.* **93**, 1136 (2010).
- ³⁸⁴D. G. Cahill, "Analysis of heat flow in layered structures for time-domain thermoreflectance," *Rev. Sci. Instrum.* **75**, 5119 (2004).
- ³⁸⁵K. Kang, Y. K. Koh, C. Chiritescu, X. Zheng, and D. G. Cahill, "Two-tint pump-probe measurements using a femtosecond laser oscillator and sharp-edged optical filters," *Rev. Sci. Instrum.* **79**, 114901 (2008).
- ³⁸⁶C. Monachon and L. Weber, "Thermal boundary conductance of transition metals on diamond," *Emerging Mater. Res.* **1**, 90–98 (2012).
- ³⁸⁷X. Zheng, D. G. Cahill, P. Krasnochtchekov, R. Averbach, and J.-C. Zhao, "High-throughput thermal conductivity measurements of nickel solid solutions and the applicability of the Wiedemann-Franz law," *Acta Mater.* **55**, 5177–5185 (2007).
- ³⁸⁸J. C. Duda, P. E. Hopinks, Y. Shen, and M. C. Gupta, "Exceptionally low thermal conductivities of films of the fullerene derivative PCBM," *Phys. Rev. Lett.* **110**, 015902 (2013).
- ³⁸⁹S. Huxtable, D. Cahill, V. Fauconnier, J. White, and J.-C. Zhao, "Thermal conductivity imaging at micrometre-scale resolution for combinatorial studies of materials," *Nature Mater.* **3**, 298–301 (2004).
- ³⁹⁰J.-C. Zhao, X. Zheng, and D. G. Cahill, "Thermal conductivity mapping of the NiAl system and the beta-NiAl phase in the NiAlCr system," *Scr. Mater.* **66**, 935–938 (2012).
- ³⁹¹X. Zheng, D. Cahill, and J.-C. Zhao, "Thermal conductivity imaging of thermal barrier coatings," *Adv. Eng. Mater.* **7**, 622–626 (2005).
- ³⁹²E. López-Honorato, C. Chiritescu, P. Xiao, D. G. Cahill, G. Marsh, and T. Abram, "Thermal conductivity mapping of pyrolytic carbon and silicon carbide coatings on simulated fuel particles by time-domain thermoreflectance," *J. Nucl. Mater.* **378**, 35–39 (2008).
- ³⁹³W.-P. Hsieh, B. Chen, J. Li, P. Keblinski, and D. Cahill, "Pressure tuning of the thermal conductivity of the layered muscovite crystal," *Phys. Rev. B* **80**, 180302 (2009).
- ³⁹⁴W.-P. Hsieh, M. D. Losego, P. V. Braun, S. Shenogin, P. Keblinski, and D. G. Cahill, "Testing the minimum thermal conductivity model for amorphous polymers using high pressure," *Phys. Rev. B* **83**, 174205 (2011).
- ³⁹⁵B. Chen, W.-P. Hsieh, D. G. Cahill, D. R. Trinkle, and J. Li, "Thermal conductivity of compressed H_2O to 22 GPa: A test of the Leibfried-Schlömann equation," *Phys. Rev. B* **83**, 132301 (2011).
- ³⁹⁶D. A. Young, C. Thomsen, H. T. Grahn, H. J. Maris, and J. Tauc, "Heat flow in glasses on a picosecond timescale," in *Phonon Scattering in Condensed Matter*, edited by A. C. Anderson and J. P. Wolfe (Springer, Berlin, 1986), p. 49.
- ³⁹⁷C. A. Paddock and G. L. Eesley, "Transient thermoreflectance from thin metal films," *J. Appl. Phys.* **60**, 285–290 (1986).
- ³⁹⁸A. J. Schmidt, X. Chen, and G. Chen, "Pulse accumulation, radial heat conduction, and anisotropic thermal conductivity in pump-probe transient thermoreflectance," *Rev. Sci. Instrum.* **79**, 114902 (2008).
- ³⁹⁹A. J. Schmidt, R. Cheaito, and M. Chiesa, "Characterization of thin metal films via frequency-domain thermoreflectance," *J. Appl. Phys.* **107**, 024908 (2010).
- ⁴⁰⁰Y. K. Koh, S. L. Singer, W. Kim, J. M. O. Zide, H. Lu, D. G. Cahill, A. Majumdar, and A. C. Gossard, "Comparison of the 3ω method and time-domain thermoreflectance for measurements of the cross-plane thermal conductivity of epitaxial semiconductors," *J. Appl. Phys.* **105**, 054303 (2009).
- ⁴⁰¹Y. Wang, J. Y. Park, Y. K. Koh, and D. G. Cahill, "Thermoreflectance of metal transducers for time-domain thermoreflectance," *J. Appl. Phys.* **108**, 043507 (2010).
- ⁴⁰²R. B. Wilson, B. A. Apgar, L. W. Martin, and D. G. Cahill, "Thermoreflectance of metal transducers for optical pump-probe studies of thermal properties," *Opt. Express* **20**, 28829–28838 (2012).
- ⁴⁰³A. Schmidt, M. Chiesa, X. Chen, and G. Chen, "An optical pump-probe technique for measuring the thermal conductivity of liquids," *Rev. Sci. Instrum.* **79**, 064902 (2008).
- ⁴⁰⁴M. A. Panzer, M. Shandalov, J. A. Rowlette, Y. Oshima, Y. W. Chen, P. C. McIntyre, and K. E. Goodson, "Thermal properties of ultrathin hafnium oxide gate dielectric films," *IEEE Electron Device Lett.* **30**, 1269–1271 (2009).
- ⁴⁰⁵S. H. Firoz, T. Yagi, N. Taketoshi, K. Ishikawa, and T. Baba, "Direct observation of thermal energy transfer across the thin metal lm on silicon substrates by a rear heatingfront detection thermoreflectance technique," *Meas. Sci. Technol.* **22**, 024012 (2011).
- ⁴⁰⁶J.-Y. Park, C.-K. Min, S. Granick, and D. G. Cahill, "Heat transfer and residence time when water droplets hit a scalding surface," *J. Heat Transfer* **134**, 101503 (2012).
- ⁴⁰⁷C.-K. Min, J.-Y. Park, D. G. Cahill, and S. Granick, "Fast, spatially-resolved thermometry of Si crystals by pump-probe two-photon absorption," *J. Appl. Phys.* **106**, 013102 (2009).
- ⁴⁰⁸D. G. Cahill and F. Watanabe, "Thermal conductivity of isotopically pure and Ge-doped Si epitaxial layers from 300 to 550 K," *Phys. Rev. B* **70**, 235322 (2004).
- ⁴⁰⁹S. Dilhaire, G. Pernot, G. Calbris, J. M. Rampnoux, and S. Grauby, "Heterodyne picosecond thermoreflectance applied to nanoscale thermal metrology," *J. Appl. Phys.* **110**, 114314 (2011).
- ⁴¹⁰A. J. Schmidt, R. Cheaito, and M. Chiesa, "A frequency-domain thermoreflectance method for the characterization of thermal properties," *Rev. Sci. Instrum.* **80**, 094901 (2009).
- ⁴¹¹K. T. Regner, D. P. Sellan, Z. Su, C. H. Amon, A. J. H. McGaughey, and J. A. Malen, "Broadband phonon mean free path contributions to thermal conductivity measured using frequency domain thermoreflectance," *Nat. Commun.* **4**, 1640 (2013).
- ⁴¹²K. T. Regner, S. Majumdar, and J. A. Malen, "Instrumentation of broadband frequency domain thermoreflectance for measuring thermal conductivity accumulation functions," *Rev. Sci. Instrum.* **84**, 064901 (2013).
- ⁴¹³Y. K. Koh and D. Cahill, "Frequency dependence of the thermal conductivity of semiconductor alloys," *Phys. Rev. B* **76**, 075207 (2007).
- ⁴¹⁴H.-S. Yang, D. G. Cahill, X. Liu, J. L. Feldman, R. S. Crandall, B. A. Sperling, and J. R. Abelson, "Anomalously high thermal conductivity of

- amorphous Si deposited by hot-wire chemical vapor deposition," *Phys. Rev. B* **81**, 104203 (2010).
- ⁴¹⁵A. J. Minnich, J. A. Johnson, A. J. Schmidt, K. Esfarjani, M. S. Dresselhaus, K. A. Nelson, and G. Chen, "Thermal conductivity spectroscopy technique to measure phonon mean free paths," *Phys. Rev. Lett.* **107**, 095901 (2011).
 - ⁴¹⁶M. Highland, B. C. Gundrum, K. K. Yee, R. S. Averbach, D. G. Cahill, V. C. Elarde, J. J. Coleman, D. A. Walko, and E. C. Landahl, "Ballistic-phonon heat conduction at the nanoscale as revealed by time-resolved x-ray diffraction and time-domain thermoreflectance," *Phys. Rev. B* **76**, 075337 (2007).
 - ⁴¹⁷M. E. Siemens, Q. Li, R. Yang, K. A. Nelson, E. H. Anderson, M. M. Mumane, and H. C. Kapteyn, "Quasi-ballistic thermal transport from nanoscale interfaces observed using ultrafast coherent soft X-ray beams," *Nature Mater.* **9**, 26–30 (2010).
 - ⁴¹⁸A. Minnich, G. Chen, S. Mansoor, and B. Yilbas, "Quasiballistic heat transfer studied using the frequency-dependent Boltzmann transport equation," *Phys. Rev. B* **84**, 235207 (2011).
 - ⁴¹⁹A. J. Minnich, "Determining phonon mean free paths from observations of quasiballistic thermal transport," *Phys. Rev. Lett.* **109**, 205901 (2012).
 - ⁴²⁰C. A. da Cruz, W. Li, N. A. Katcho, and N. Mingo, "Role of phonon anharmonicity in time-domain thermoreflectance measurements," *Appl. Phys. Lett.* **101**, 083108 (2012).
 - ⁴²¹J. A. Johnson, A. A. Maznev, J. Cuffe, J. K. Eliason, A. J. Minnich, T. Kehoe, C. M. S. Torres, G. Chen, and K. A. Nelson, "Direct measurement of room-temperature nondiffusive thermal transport over micron distances in a silicon membrane," *Phys. Rev. Lett.* **110**, 025901 (2013).
 - ⁴²²K. L. Grosse, M.-H. Bae, F. Lian, E. Pop, and W. P. King, "Nanoscale Joule heating, Peltier cooling and current crowding at graphene metal contacts," *Nat. Nanotechnol.* **6**, 287–290 (2011).
 - ⁴²³S. P. Gurrum, W. P. King, Y. K. Joshi, and K. Ramakrishna, "Size effect on the thermal conductivity of thin metallic films investigated by scanning Joule expansion microscopy," *ASME Trans. J. Heat Transfer* **130**, 082403 (2008).
 - ⁴²⁴P. C. Fletcher, B. Lee, and W. P. King, "Thermoelectric voltage at a nanometer-scale heated tip point contact," *Nanotechnology* **23**, 035401 (2012).
 - ⁴²⁵K. Goodson and M. Asheghi, "Near-field optical thermometry," *Microscale Thermophys. Eng.* **1**, 225–235 (1997).
 - ⁴²⁶P. O. Chapuis, J. J. Greffet, K. Joulain, and S. Volz, "Heat transfer between a nano-tip and a surface," *Nanotechnology* **17**, 2978–2981 (2006).
 - ⁴²⁷X. Chen and X. Wang, "Near-field thermal transport in a nanotip under laser irradiation," *Nanotechnology* **22**, 075204 (2011).
 - ⁴²⁸J. Chung, K. Kim, G. Hwang, O. Kwon, S. Jung, J. Lee, J. W. Lee, and G. T. Kim, "Quantitative temperature measurement of an electrically heated carbon nanotube using the null-point method," *Rev. Sci. Instrum.* **81**, 114901 (2010).
 - ⁴²⁹K. Kim, J. Chung, G. Hwang, O. Kwon, and J. S. Lee, "Quantitative measurement with scanning thermal microscope by preventing the distortion due to the heat transfer through the air," *ACS Nano* **5**, 8700–8709 (2011).
 - ⁴³⁰K. Kim, J. Chung, J. Won, O. Kwon, J. S. Lee, S. H. Park, and Y. K. Choi, "Quantitative scanning thermal microscopy using double scan technique," *Appl. Phys. Lett.* **93**, 203115 (2008).
 - ⁴³¹K. Kim, W. Jeong, W. Lee, and P. Reddy, "Ultra-high vacuum scanning thermal microscopy for nanometer resolution quantitative thermometry," *ACS Nano* **6**, 4248–4257 (2012).
 - ⁴³²P. Reddy, S.-Y. Jang, R. A. Segalman, and A. Majumdar, "Thermoelectricity in molecular junctions," *Science* **315**, 1568–1571 (2007).
 - ⁴³³H.-K. Lyeo, A. A. Khajetoorians, L. Shi, K. P. Pipe, R. J. Ram, A. Shakouri, and C. K. Shih, "Profiling the thermoelectric power of semiconductor junctions with nanometer resolution," *Science* **303**, 816–818 (2004).
 - ⁴³⁴Y. Dubi and M. Di Ventra, "Colloquium: Heat flow and thermoelectricity in atomic and molecular junctions," *Rev. Mod. Phys.* **83**, 131–155 (2011).
 - ⁴³⁵B. Gotsmann and M. A. Lantz, "Quantized thermal transport across contacts of rough surfaces," *Nature Mater.* **12**, 59–65 (2013).
 - ⁴³⁶D. Dietzel, S. Chotikaprakhan, B. K. Bein, and J. Pelzl, "Analysis of active semiconductor structures by combined SThM and SThEM," *J. Phys. IV (France)* **125**, 87–91 (2005).
 - ⁴³⁷S. Grauby, L.-D. P. Lopez, A. Salhi, E. Puyoo, J.-M. Rampnoux, W. Claeys, and S. Dilhaire, "Joule expansion imaging techniques on micro-electronic devices," *Microelectron. J.* **40**, 1367–1372 (2009).
 - ⁴³⁸O. Kwon, L. Shi, and A. Majumdar, "Scanning thermal wave microscopy (STWM)," *ASME Trans. J. Heat Transfer* **125**, 156–163 (2003).
 - ⁴³⁹B. Gotsmann and U. Durig, "Thermally activated nanowear modes of a polymer surface induced by a heated tip," *Langmuir* **20**, 1495–1500 (2004).
 - ⁴⁴⁰S. Jesse, M. P. Nikiforov, L. T. Germinario, and S. V. Kalinin, "Local thermomechanical characterization of phase transitions using band excitation atomic force acoustic microscopy with heated probe," *Appl. Phys. Lett.* **93**, 073104 (2008).
 - ⁴⁴¹W. P. King, S. Saxena, B. A. Nelson, B. L. Weeks, and R. Pitchimani, "Nanoscale thermal analysis of an energetic material," *Nano Lett.* **6**, 2145–2149 (2006).
 - ⁴⁴²E. A. Corbin and W. P. King, "Electrical noise characteristics of a doped silicon microcantilever heater-thermometer," *Appl. Phys. Lett.* **99**, 263107 (2011).
 - ⁴⁴³U. Duerig, "Fundamentals of micromechanical thermoelectric sensors," *J. Appl. Phys.* **98**, 044906 (2005).
 - ⁴⁴⁴J. H. Bae, T. Ono, and M. Esashi, "Scanning probe with an integrated diamond heater element for nanolithography," *Appl. Phys. Lett.* **82**, 814–816 (2003).
 - ⁴⁴⁵Z. Dai, W. P. King, and K. Park, "A 100 nanometer scale resistive heater-thermometer on a silicon cantilever," *Nanotechnology* **20**, 095301 (2009).
 - ⁴⁴⁶Z. T. Dai, E. A. Corbin, and W. P. King, "A microcantilever heater-thermometer with a thermal isolation layer for making thermal nanotopography measurements," *Nanotechnology* **21**, 055503 (2010).
 - ⁴⁴⁷U. Drechsler, N. Burer, M. Despont, U. Durig, B. Gotsmann, F. Robin, and P. Vettiger, "Cantilevers with nano-heaters for thermomechanical storage application," *Microelectron. Eng.* **67–68**, 397–404 (2003).
 - ⁴⁴⁸P. C. Fletcher, B. S. Bhatia, Y. Wu, M. A. Shannon, and W. P. King, "Electrothermal atomic-force microscope cantilever with integrated heater and n-p-n back-to-back diodes," *J. Microelectromech. Syst.* **20**, 644–653 (2011).
 - ⁴⁴⁹A. Jungen, M. Pfenniger, M. Tonteling, C. Stampfer, and C. Hierold, "Electrothermal effects at the microscale and their consequences on system design," *J. Micromech. Microeng.* **16**, 1633–1638 (2006).
 - ⁴⁵⁰B. Lee, C. B. Prater, and A. P. King, "Lorentz force actuation of a heated atomic force microscope cantilever," *Nanotechnology* **23**, 055709 (2012).
 - ⁴⁵¹D. W. Lee, T. Ono, T. Abe, and M. Esashi, "Microprobe array with electrical interconnection for thermal imaging and data storage," *J. Microelectromech. Syst.* **11**, 215–221 (2002).
 - ⁴⁵²A. S. Basu, S. McNamara, and Y. B. Gianchandani, "Scanning thermal lithography: Maskless, submicron thermochemical patterning of photoresist by ultracompliant probes," *J. Vac. Sci. Technol. B* **22**, 3217–3220 (2004).
 - ⁴⁵³S. Chung, J. R. Felts, D. Wang, W. P. King, and J. J. De Yoreo, "Temperature-dependence of ink transport during thermal dip-pen nanolithography," *Appl. Phys. Lett.* **99**, 193101 (2011).
 - ⁴⁵⁴J. Duvigneau, H. Schonherr, and G. J. Vancso, "Scanning thermal lithography of Tailored tert-butyl ester protected carboxylic acid functionalized (meth)acrylate polymer platforms," *ACS Appl. Mater. Interfaces* **3**, 3855–3865 (2011).
 - ⁴⁵⁵J. R. Felts, S. Somnath, R. H. Ewoldt, and W. P. King, "Nanometer-scale flow of molten polyethylene from a heated atomic force microscope tip," *Nanotechnology* **23**, 215301 (2012).
 - ⁴⁵⁶O. Fenwick, L. Bozec, D. Credgington, A. Hammiche, G. M. Lazzerini, Y. R. Silberberg, and F. Cacialli, "Thermochemical nano-patterning of organic semiconductors," *Nat. Nanotechnol.* **4**, 664–668 (2009).
 - ⁴⁵⁷P. C. Fletcher, J. R. Felts, Z. T. Dai, T. D. Jacobs, H. J. Zeng, W. Lee, P. E. Sheehan, J. A. Carlisle, R. W. Carpick, and W. P. King, "Wear-resistant diamond nanoprobe tips with integrated silicon heater for tip-based nanomanufacturing," *ACS Nano* **4**, 3338–3344 (2010).
 - ⁴⁵⁸S. Kim, Y. Bastani, H. D. Lu, W. P. King, S. Marder, K. H. Sandhage, A. Gruverman, E. Riedo, and N. Bassiri-Gharb, "Direct fabrication of arbitrary-shaped ferroelectric nanostructures on plastic, glass, and silicon substrates," *Adv. Mater.* **23**, 3786–3790 (2011).
 - ⁴⁵⁹A. W. Knoll, D. Pires, O. Coulembier, P. Dubois, J. L. Hedrick, J. Frommer, and U. Duerig, "Probe-based 3-D nanolithography using self-amplified depolymerization polymers," *Adv. Mater.* **22**, 3361–3365 (2010).