

MIT Open Access Articles

*Generalized Two-Temperature Model
for Coupled Phonon-Magnon Diffusion*

The MIT Faculty has made this article openly available. **Please share** how this access benefits you. Your story matters.

Citation: Liao, Bolin, Jiawei Zhou, and Gang Chen. "Generalized Two-Temperature Model for Coupled Phonon-Magnon Diffusion." *Physical Review Letters* 113, no. 2 (July 2014).

As Published: <http://dx.doi.org/10.1103/PhysRevLett.113.025902>

Publisher: American Physical Society

Persistent URL: <http://hdl.handle.net/1721.1/88472>

Version: Author's final manuscript: final author's manuscript post peer review, without publisher's formatting or copy editing

Terms of use: Creative Commons Attribution-Noncommercial-Share Alike



Generalized two-temperature model for coupled phonon-magnon diffusion

Bolin Liao, Jiawei Zhou and Gang Chen*

Department of Mechanical Engineering, Massachusetts Institute of Technology,
Cambridge, Massachusetts, 02139, USA

Abstract

We generalize the two-temperature model [Sanders and Walton, Phys. Rev. B, **15**, 1489 (1977)] for coupled phonon-magnon diffusion to include the effect of the concurrent magnetization flow, with a particular emphasis on the thermal consequence of the magnon flow driven by a non-uniform magnetic field. Working within the framework of Boltzmann transport equation, we derive the constitutive equations for coupled phonon-magnon transport driven by gradients of both temperature and external magnetic fields, and the corresponding conservation laws. Our equations reduce to the original Sanders-Walton two-temperature model under a uniform external field, but predict a new magnon cooling effect driven by a non-uniform magnetic field in a homogeneous single-domain ferromagnet. We estimate the magnitude of the cooling effect in yttrium iron garnet, and show it is within current experimental reach. With properly optimized materials, the predicted cooling effect can potentially supplement the conventional magnetocaloric effect in cryogenic applications in the future.

Spin caloritronics [1,2] is a nascent field of study that looks into the interaction between heat and spin. In addition to providing ways of thermally manipulating magnetization and magnetic domain walls [3–5] as supplements to conventional spintronics, it also holds

* To whom correspondence should be addressed: gchen2@mit.edu.

promise of novel energy harvesting and cooling applications owing to the recent discovery of the spin Seebeck effect (SSE) [6–10] and its reciprocal spin Peltier effect (SPE) [11,12]. Despite existing debates on details, it has been widely recognized that the aforementioned spin caloritronic effects are consequences of the interactions between phonons, electrons and spins [13–16]. From this perspective, spin caloritronics is a natural extension of both thermoelectrics and spintronics. Phonons are responsible for heat conduction in most solids; in metallic and semiconducting materials, electrons are carriers of charge, heat and spin; in magnetic materials, magnons [17] – the collective excitations of spins – also participate in transporting spin [18] and heat [19]. Coupled to these carriers are thermodynamic forces that drive their flows [20]: the gradients of temperature, electrochemical potential and non-equilibrium magnetization [21]. For conditions close to equilibrium, it is particularly convenient to treat the coupled transport phenomena within the phenomenological framework of irreversible thermodynamics [20], where the Onsager reciprocity relation serves as the link between concurrent flows. Routinely used in studying the coupled transport of electrons and phonons [20–22], the method of irreversible thermodynamics has also been utilized in analyzing the coupled transport of heat and charge with spins [21,23–26].

In this paper we limit our discussion to ferromagnetic insulators without free conducting electrons. Further steps to understanding the spin caloritronic effects require microscopic models that provide quantitative information of the transport processes, for example the kinetic coefficients [20] that connect the driving forces to the corresponding fluxes. For studying thermoelectrics, the coupled transport processes are typically treated within the framework of Boltzmann transport equation (BTE) [22], which in the diffusion regime

gives quantitative kinetic coefficients, and is capable of delineating ballistic transport [27] when solved with proper boundary conditions. It is particularly a natural way to describe thermally induced transport processes where coherent contributions are not important. On the other hand, the spintronics community often uses the Landau-Lifshitz-Gilbert (LLG) [28] equation for the dynamics of magnetization. Compared with BTE, LLG adopts more a “wave-like” point of view, where the coherent dynamics is important and the thermal relaxation acts as a damping factor. Indeed the long wavelength magnons have been shown to exhibit macroscopic coherence lengths at room temperature [29], and LLG is necessary to account for their behaviors. For the thermal transport, however, magnons with a wide range of wavelengths and coherent lengths will be excited, and LLG seems no longer a particularly convenient description. A recent work by Hoffman *et al.* [30] applied a “semi-phenomenological” stochastic LLG equation to modeling the longitudinal SSE, where the temperature effect was incorporated via a thermally fluctuating Langevin field. Since a linear phonon temperature distribution was presumed in their work, it did not fully solve the coupled phonon-magnon transport problem. An alternative approach to this problem adopts a more “particle-like” picture. The pioneering work by Sanders and Walton [31] treated the coupled phonon-magnon thermal diffusion process with a two-temperature model, where phonons and magnons were modeled as two gases of bosons, each locally in thermal equilibrium with different temperatures, and the local energy exchange rate between them is proportional to the temperature difference. This model was later used to explain the spin Seebeck effect [13], and was recently extended to take into account the boundary heat and spin transfer [32]. It also served as a modeling tool for interpreting dynamic

measurements of the thermal conductivity of spin ladder compounds [33,34] and the static direct measurement of the magnon temperature [35], and has been applied to other carrier systems such as electron-phonon [36] and acoustic-phonon-optical-phonon [37].

In their original formulation, Sanders and Walton did not consider the associated magnetization flow with the magnon heat flow. On the other hand, Meier and Loss [38] showed that the magnon flow could also be generated by a non-uniform external magnetic field, but they did not look into the thermal aspect of this transport process. More studies also discussed the similarity between field-driven magnon transport and electron transport [39–43]. In a recent work by Kosevich and Gann [44], both quantum and semiclassical dynamics of a field-driven magnon flow was thoroughly studied. In the current paper we attempt to combine the two paths, one from the thermal perspective and the other from the magnetic perspective, and give a unified description of the coupled phonon-magnon diffusive transport of both heat and magnetization, which is also applicable when the external magnetic field is non-uniform, with a special focus on the thermal effect associated with the field-driven magnon flow.

Magnons are (in most cases [34]) bosonic excitations, and in equilibrium obey the Bose-Einstein distribution:

$$f_0(\mathbf{r}, \mathbf{k}) = \frac{1}{\exp\left[\frac{\hbar\omega(\mathbf{k}) + g\mu_B B(\mathbf{r})}{k_B T_m(\mathbf{r})}\right] - 1}, \quad (1)$$

where $\hbar\omega(\mathbf{k})$ is the magnon energy [17,28] without external magnetic field, g is the Landé g -factor, μ_B is the Bohr magneton ($-g\mu_B$ combined represents the amount of magnetic moment carried by a single magnon [38]), T_m is the magnon temperature and B is the external magnetic field. Here we neglect the magnetic dipolar interaction and

magnetic anisotropy for simplicity. Although magnons can reach a quasi-equilibrium state with a finite chemical potential under parametric pumping [45], here we treat magnons with vanishing chemical potential for local equilibrium is assumed. Next we write down the steady-state Boltzmann transport equation with the relaxation time approximation (RTA):

$$-\frac{f - f_0}{\tau_m} = \mathbf{v} \cdot \nabla_{\mathbf{r}} f_0, \quad (2)$$

where $f(\mathbf{r}, \mathbf{k})$ is the non-equilibrium distribution function of magnons, $\mathbf{v}(\mathbf{k})$ is the

group velocity of magnons, $\tau_m = \left(\frac{1}{\tau_{m-m}} + \frac{1}{\tau_{m-p,ela}} + \frac{1}{\tau_{m-imp}} \right)^{-1}$ is a lumped relaxation time

of magnons including effects of magnon-magnon scattering [46], elastic magnon-phonon scattering [47] and elastic magnon-impurity scattering [48]. The inelastic magnon-phonon scattering is responsible for the local energy exchange between magnons and phonons [47], and in general cannot be written in a relaxation-time form [22]. Thus we follow Sanders and Walton [31] here and consider the energy exchange process separately in the conservation laws later. We emphasize the validity of this separation requires that phonon-magnon interactions be much weaker than magnon-magnon interactions.

After obtaining the non-equilibrium distribution function $f(\mathbf{r}, \mathbf{k})$, we can calculate the local magnetization and heat flows carried by magnons. The magnetization flow is

$$\mathbf{J}_m = -g\mu_B \int \frac{d^3\mathbf{k}}{(2\pi)^3} f \mathbf{v},$$

where the minus sign accounts for the fact that the excitation of magnons reduces the total magnetization [49]: $\mathbf{M}(\mathbf{r}) = \mathbf{M}_s - g\mu_B n_m$, where \mathbf{M}_s is the

saturation magnetization, and n_m is the number density of magnons. To calculate the magnon heat flow, we start with the thermodynamic relation of a magnet [20]: $dE = dQ + BdM = dQ - Bg\mu_B dn_m$, where E is the total energy of the magnet and the interaction energy (BM) between the magnet and the magnetic field, and thus field-independent [50], corresponding to $\hbar\omega(\mathbf{k})$ microscopically (in contrast $\hbar\omega(\mathbf{k}) + g\mu_B B$ corresponds to the field-dependent “spectroscopic energy” [50]). Differentiating the above relation with respect to time, we get the magnon heat flux $\mathbf{J}_{qm} = \mathbf{J}_e - B\mathbf{J}_m = \int \frac{d^3\mathbf{k}}{(2\pi)^3} (\hbar\omega + g\mu_B B) f \mathbf{v}$, where \mathbf{J}_{qm} is the magnon heat flux, \mathbf{J}_e is the magnon energy flux. The term $B\mathbf{J}_m$ describes the transport of the magnetic interaction energy associated with the magnetization flow, analogous to $\phi\mathbf{J}_e$ in the case of electrons, where ϕ is the electro-static potential and \mathbf{J}_e is the electrical charge flux. Combining the above expression with Eqs. (1) and (2), we arrive at the constitutive equations for the magnon transport:

$$-\mathbf{J}_m = L_{11}\nabla B + L_{12}(-\nabla T_m), \quad (3)$$

$$\mathbf{J}_{qm} = L_{12}T_m\nabla B + L_{22}(-\nabla T_m), \quad (4)$$

with the kinetic coefficients given by (assuming an isotropic magnon dispersion):

$$L_{11} = -\frac{(g\mu_B)^2}{3} \int_{\omega} \tau_m v^2 \frac{\partial f_0}{\partial(\hbar\omega)} D(\omega) d\omega, \quad (5)$$

$$L_{12} = -\frac{g\mu_B}{3T_m} \int_{\omega} (\hbar\omega + g\mu_B B) \tau_m v^2 \frac{\partial f_0}{\partial(\hbar\omega)} D(\omega) d\omega, \quad (6)$$

$$L_{22} = -\frac{1}{3T_m} \int_{\omega} (\hbar\omega + g\mu_B B)^2 \tau_m v^2 \frac{\partial f_0}{\partial(\hbar\omega)} D(\omega) d\omega, \quad (7)$$

where $D(\omega)$ is the magnon density of states. We can interpret L_{11} as the isothermal magneto-conductivity σ_m and L_{22} as the uniform-field magnon thermal conductivity κ_m , and define L_{12} as a magneto-thermal coupling coefficient ζ_m . Note that the Onsager reciprocity relation manifests itself explicitly in Eqs. (3) and (4). It can be shown using Cauchy-Schwartz inequality that $L_{11}L_{22} \geq T_m L_{12}^2$ (in the case of electron transport, this inequality implies a positive zero-current thermal conductivity [22]), which guarantees the net entropy generation in this system is non-negative [51]. Eqs. (3)-(7) are reminiscent of electron transport, and the external field B seems to play a similar role as the electrochemical potential of electrons. We need to point out here, however, a critical difference between electrons and magnons. The number of electrons is conserved, thus the electrochemical potential includes the contribution of a finite chemical potential that can be “self-adjusted” during the transport process, whereas the number of magnons is not conserved, and the B field does not contain a similar contribution as the chemical potential of electrons (given the magnetic dipolar interaction is negligible).

With the constitutive equations (3) and (4), we still need conservation laws to complete the formulation. We first look at the phonon system. At steady state, the phonon energy can either be transported by the phonon heat flux or transferred to the magnon system. Thus in the spirit of Sanders and Walton’s original work, the phonon energy conservation states:

$$\nabla \cdot \mathbf{J}_{qp} = \frac{C_m C_p}{C_m + C_p} \frac{T_m - T_p}{\tau_{mp}} \equiv g_{mp} (T_m - T_p), \quad (8)$$

where \mathbf{J}_{qp} is the phonon heat flux, T_p is the phonon temperature, C_m and C_p are the volumetric specific heat of magnons and phonons, τ_{mp} is a phenomenological time scale

characterizing the inelastic interaction between phonons and magnons, and we define g_{mp} as a lumped coefficient of phonon-magnon energy exchange. It is worth mentioning that Eq. (8) is the result of inelastic phonon-magnon scattering and in principle can be derived from a full version of BTE, similar as in the case of electron-phonon coupling [22]. Another conservation law has to do with the energy input from an external power source. The impression that a magnon flow can be generated by a non-uniform static magnetic field can be misleading because it violates the second law of thermodynamics: no energy is put into the system, while the magnon flow can potentially output work. In reality, when the magnetization of the magnet changes, an electromotive force (EMF) is induced in the electromagnet (e.g., a solenoid). To maintain the magnetic field, the current running through the electromagnet has to overcome this EMF and thus do work. It can be shown [50] that the work done by the current in this process is precisely equal to BdM . Hence the local creation and annihilation of magnons enables the energy exchange between the system (including the ferromagnet itself and its interaction with the field) and the external power supply. A local version of the above statement can be translated to $\nabla \cdot (\mathbf{J}_{qm} + B\mathbf{J}_m + \mathbf{J}_{qp}) = B\nabla \cdot \mathbf{J}_m$, or more explicitly:

$$\nabla \cdot \mathbf{J}_{qm} + \nabla B \cdot \mathbf{J}_m = g_{mp}(T_p - T_m). \quad (9)$$

Now combining Eqs. (8), (9) with (3) and (4), and the normal Fourier law for phonon heat conduction: $\mathbf{J}_{qp} = -\kappa_p \nabla T_p$ (κ_p is the phonon thermal conductivity), the governing equations for the temperature distributions of magnons and phonons read (considering 1-dimensional situations):

$$-\kappa_p \frac{\partial^2 T_p}{\partial x^2} = g_{mp} (T_m - T_p), \quad (10)$$

$$-\kappa_m \frac{\partial^2 T_m}{\partial x^2} + \left(2\zeta_m \frac{\partial B}{\partial x} \right) \frac{\partial T_m}{\partial x} + \zeta_m \frac{\partial^2 B}{\partial x^2} T_m - \sigma_m \left(\frac{\partial B}{\partial x} \right)^2 = g_{mp} (T_p - T_m). \quad (11)$$

Here we assume the applied temperature and magnetic field gradients are small and thus the transport coefficients are averaged values that do not explicitly depend on T_m or B . Eqs. (10) and (11) reduce to the original Sanders-Walton model when the external magnetic field is uniform, even though in this case the magnetization flow is present ($\mathbf{J}_m = -\zeta_m \nabla T_m$).

More interesting phenomena emerge when non-uniform external magnetic field is applied. We expect a non-uniform external field will drive magnon flow, which is associated with a magnon heat flow, and cause temperature redistribution of both magnons and phonons due to the phonon-magnon coupling. Without a concise analytic solution with the coupling terms, we turn to numerical solutions for clarity, before which we first estimate the kinetic coefficients based on information in literature on yttrium iron garnet (YIG). Since the magnetic energy scale is pretty small ($g\mu_B \approx 1.3$ K/T, for $g = 2$ in YIG), we expect the predicted effect to be more pronounced at low temperatures. Thus we use the low temperature expansion of the magnon dispersion $\hbar\omega(\mathbf{k}) = Dk^2 a^2$, where $D \approx 1.8$ meV [52], and the lattice constant $a = 12.3$ Å for YIG [53]. For a similar reason, we neglect the field dependence of the kinetic coefficients in the following discussion. Further assuming a constant relaxation time τ_m , we obtain the ratio

$$\frac{\sigma_m}{\zeta_m} = \frac{g\mu_B}{k_B} \frac{\xi(1.5)}{\xi(2.5)} = 1.304 \text{ K/T} \text{ with } \xi(t) = \int_0^{+\infty} \frac{x^t e^x}{(e^x - 1)^2} dx, \text{ which is analogous to the}$$

inverse of the Seebeck coefficient in the electron case. The value of τ_m is highly controversial [13], and here we adopt a value of $\tau_m \sim 1$ ns, which leads to the calculated uniform-field magnon thermal conductivity $\kappa_m \approx 8$ W/mK at 20K with zero field that is at least of the reasonable order of magnitude compared with the experiment [54]. With the same relaxation time, we obtain $\sigma_m \approx 0.25$ W/mT², and $\zeta_m \approx 0.19$ W/mTK. For phonons, we choose $\kappa_p \approx 50$ W/mK [54]. At 20K, the specific heat of magnons and phonons are on the same order ($\sim 10^4$ J/m³K) [55]. Different claims on the value of τ_{mp} exists, ranging from below a few nanoseconds [32,56] to longer than a few hundred nanoseconds [13,57–59] at 300K. At lower temperature, this relaxation time will be longer, and we tentatively choose $\tau_{mp} \approx 100$ ns due to the large uncertainty of available data.

Provided the above parameters, we study numerically an experimentally realizable case: a strip of YIG (100 μ m long) connected to a thermal reservoir at 20K with one end, and the other end isolated. If part of the YIG strip is covered by a magnetic shielding material with high magnetic permeability, a step-like magnetic field can be realized within YIG just by applying a uniform field. We model this step-like magnetic field as a smeared-out Fermi-Dirac function as shown in Fig. 1(a) (the length scale of the smearing is chosen to be much larger than the mean free paths of magnons to avoid the complication of ballistic transport, which in principle can still be fully captured by the BTE), and calculate the phonon temperature at the isolated end. In this case we apply adiabatic boundary conditions for magnons at both ends ($\mathbf{J}_e = \mathbf{J}_{qm} + B\mathbf{J}_m = 0$). A phonon-temperature-drop of ~ 56 mK is predicted under a step field varying from 0.5T to 1.5T, with the temperature

distribution of both phonons and magnons shown in Fig. 1(b). This temperature drop can be further amplified by increasing the field gradient as illustrated in Fig. 1(c). We would like to emphasize that the estimation here is very rough due to the lack of information, and is only intended to demonstrate a probable order of magnitude of this effect. The calculation above indicates that this magnon cooling effect may be detected under currently available experimental resolution. In passing we note that the predicted effect differs from the conventional magnetocaloric effect [60], such as adiabatic demagnetization, in that the magnetocaloric effect utilizes thermodynamic properties of the magnet (i.e. the field-dependent specific heat) in equilibrium, and a uniform field is often applied.

We provide another example where the magnon cooling effect is set up in close analogy to a thermoelectric Peltier cooling unit and calculate the coefficient of performance (COP) and effective zT . In this example the YIG strip is sandwiched between two thermal reservoirs with temperatures $T_h > T_c$, when a step field (as in Fig. 1(a)) is applied. The temperature profiles when $T_h = 20$ K, $T_h - T_c = 30$ mK and $(B_0, B_1) = (0.5$ T, 1.5 T) are plotted in Fig. 2(a), and it is clearly shown that heat is moved from the cold source to

the hot source. The COP can be calculated via
$$\text{COP} \equiv \frac{Q_c}{W} = \frac{\mathbf{J}_{qp.cold}}{-\int_0^L B \nabla \cdot \mathbf{J}_m dx}$$
, and is plotted

in Figs. 2(b) and 2(c) against varying temperature and field difference. At the fixed temperature difference of 30 mK (Fig. 2(b)), the optimal COP is around 2, corresponding to an equivalent thermoelectric module with $ZT=0.01$. From Fig. 2(c), the maximal attainable temperature difference is ~ 60 mK when $(B_0, B_1) = (0.5$ T, 1.5 T), where the COP drops to zero.

In summary, we have developed a semi-classical transport theory for coupled phonon-magnon diffusion. The merit of this work lies in the fact that we apply the techniques widely used in the field of thermoelectrics to the study of magnetization transport, utilize the analogy between field-driven electron and magnon transport, and combine the thermal effect with the field-driven magnon transport in a natural way. Our theory takes into account that magnon flow can be driven by non-uniform magnetic field, and predicts that the heat carried by magnons associated with their flow can result in a cooling effect. In real materials, non-ideal effects such as magnetic dipolar interactions and the magnetic anisotropy need to be considered as a refinement to this work. We have estimated the magnitude of the magnon cooling effect in YIG, to show it can be verified by experiments. For practical uses, however, it is necessary to search for more suitable materials (preferably with lower thermal conductivities, and strong phonon-magnon interaction), and optimize the material properties via engineering efforts. We envision this new effect could supplement the conventional magnetocaloric effect in cryogenic applications in future.

Acknowledgements

We thank Sangyeop Lee and Mehmet Onbasli for helpful discussions. This article is based upon work supported partially by S³TEC, an Energy Frontier Research Center funded by the U.S. Department of Energy, Office of Basic Energy Sciences, under Award No. DE-FG02-09ER46577 (for understanding the coupled phonon-magnon transport), and partially by the Air Force Office of Scientific Research Multidisciplinary Research

Program of the University Research Initiative (AFOSR MURI) via Ohio State University
(for studying the potential application of magnon cooling at cryogenic temperatures).

References:

- [1] G. E. W. Bauer, E. Saitoh, and B. J. van Wees, *Nat. Mater.* **11**, 391 (2012).
- [2] S. R. Boona, R. C. Myers, and J. P. Heremans, *Energy Environ. Sci.* **7**, 885 (2014).
- [3] K.-R. Jeon, B.-C. Min, S.-Y. Park, K.-D. Lee, H.-S. Song, Y.-H. Park, Y.-H. Jo, and S.-C. Shin, *Sci. Rep.* **2**, (2012).
- [4] M. Hatami, G. E. W. Bauer, Q. Zhang, and P. J. Kelly, *Phys. Rev. Lett.* **99**, 066603 (2007).
- [5] W. Jiang, P. Upadhyaya, Y. Fan, J. Zhao, M. Wang, L.-T. Chang, M. Lang, K. L. Wong, M. Lewis, Y.-T. Lin, J. Tang, S. Cherepov, X. Zhou, Y. Tserkovnyak, R. N. Schwartz, and K. L. Wang, *Phys. Rev. Lett.* **110**, 177202 (2013).
- [6] K. Uchida, S. Takahashi, K. Harii, J. Ieda, W. Koshibae, K. Ando, S. Maekawa, and E. Saitoh, *Nature* **455**, 778 (2008).
- [7] C. M. Jaworski, J. Yang, S. Mack, D. D. Awschalom, J. P. Heremans, and R. C. Myers, *Nat. Mater.* **9**, 898 (2010).
- [8] K. Uchida, J. Xiao, H. Adachi, J. Ohe, S. Takahashi, J. Ieda, T. Ota, Y. Kajiwara, H. Umezawa, H. Kawai, G. E. W. Bauer, S. Maekawa, and E. Saitoh, *Nat. Mater.* **9**, 894 (2010).
- [9] K. Uchida, H. Adachi, T. An, T. Ota, M. Toda, B. Hillebrands, S. Maekawa, and E. Saitoh, *Nat. Mater.* **10**, 737 (2011).
- [10] C. M. Jaworski, R. C. Myers, E. Johnston-Halperin, and J. P. Heremans, *Nature* **487**, 210 (2012).
- [11] J. Flipse, F. L. Bakker, A. Slachter, F. K. Dejene, and B. J. van Wees, *Nat. Nanotechnol.* **7**, 166 (2012).
- [12] J. Flipse, F. K. Dejene, D. Wagenaar, G. E. W. Bauer, J. B. Youssef, and B. J. van Wees, arXiv:1311.4772 (2013).
- [13] J. Xiao, G. E. W. Bauer, K. Uchida, E. Saitoh, and S. Maekawa, *Phys. Rev. B* **81**, 214418 (2010).
- [14] H. Adachi, J. Ohe, S. Takahashi, and S. Maekawa, *Phys. Rev. B* **83**, 094410 (2011).
- [15] H. Adachi, K. Uchida, E. Saitoh, and S. Maekawa, *Rep. Prog. Phys.* **76**, 036501 (2013).
- [16] K. S. Tikhonov, J. Sinova, and A. M. Finkel'stein, *Nat. Commun.* **4**, (2013).
- [17] C. Kittel and C. Y. Fong, *Quantum Theory of Solids* (Wiley, New York, 1987).
- [18] B. Wang, J. Wang, J. Wang, and D. Y. Xing, *Phys. Rev. B* **69**, 174403 (2004).
- [19] C. Hess, *Eur. Phys. J. Spec. Top.* **151**, 73 (2007).

- [20] H. B. Callen, *Thermodynamics and an Introduction to Thermostatistics* (Wiley, New York, 1985).
- [21] M. Johnson and R. H. Silsbee, *Phys. Rev. B* **35**, 4959 (1987).
- [22] G. Chen, *Nanoscale Energy Transport and Conversion: A Parallel Treatment of Electrons, Molecules, Phonons, and Photons* (Oxford University Press, Oxford; New York, 2005).
- [23] W. M. Saslow, *Phys. Rev. B* **76**, 184434 (2007).
- [24] W. M. Saslow and K. Rivkin, *J. Magn. Magn. Mater.* **320**, 2622 (2008).
- [25] M. R. Sears and W. M. Saslow, *Phys. Rev. B* **85**, 035446 (2012).
- [26] A. A. Kovalev and Y. Tserkovnyak, *EPL Europhys. Lett.* **97**, 67002 (2012).
- [27] G. Chen, *Phys. Rev. B* **57**, 14958 (1998).
- [28] D. D. Stancil and A. Prabhakar, *Spin Waves Theory and Applications* (Springer, New York, 2009).
- [29] Y. Kajiwara, K. Harii, S. Takahashi, J. Ohe, K. Uchida, M. Mizuguchi, H. Umezawa, H. Kawai, K. Ando, K. Takanashi, S. Maekawa, and E. Saitoh, *Nature* **464**, 262 (2010).
- [30] S. Hoffman, K. Sato, and Y. Tserkovnyak, *Phys. Rev. B* **88**, 064408 (2013).
- [31] D. J. Sanders and D. Walton, *Phys. Rev. B* **15**, 1489 (1977).
- [32] M. Schreier, A. Kamra, M. Weiler, J. Xiao, G. E. W. Bauer, R. Gross, and S. T. B. Goennenwein, *Phys. Rev. B* **88**, 094410 (2013).
- [33] M. Montagnese, M. Otter, X. Zotos, D. A. Fishman, N. Hlubek, O. Mityashkin, C. Hess, R. Saint-Martin, S. Singh, A. Revcolevschi, and P. H. M. van Loosdrecht, *Phys. Rev. Lett.* **110**, 147206 (2013).
- [34] G. T. Hohensee, R. B. Wilson, J. P. Feser, and D. G. Cahill, *Phys. Rev. B* **89**, 024422 (2014).
- [35] M. Agrawal, V. I. Vasyuchka, A. A. Serga, A. D. Karenowska, G. A. Melkov, and B. Hillebrands, *Phys. Rev. Lett.* **111**, 107204 (2013).
- [36] G. Chen, *J. Appl. Phys.* **97**, 083707 (2005).
- [37] R. B. Wilson, J. P. Feser, G. T. Hohensee, and D. G. Cahill, *Phys. Rev. B* **88**, 144305 (2013).
- [38] F. Meier and D. Loss, *Phys. Rev. Lett.* **90**, 167204 (2003).
- [39] D. Forster, *Hydrodynamic Fluctuations, Broken Symmetry, And Correlation Functions* (Westview Press, Reading, Mass., 1995).
- [40] H. Castella, X. Zotos, and P. Prelovšek, *Phys. Rev. Lett.* **74**, 972 (1995).
- [41] X. Zotos, *Phys. Rev. Lett.* **82**, 1764 (1999).
- [42] J. C. Slonczewski, *Phys. Rev. B* **39**, 6995 (1989).
- [43] J. V. Alvarez and C. Gros, *Phys. Rev. Lett.* **88**, 077203 (2002).
- [44] Y. A. Kosevich and V. V. Gann, *J. Phys. Condens. Matter* **25**, 246002 (2013).
- [45] S. O. Demokritov, V. E. Demidov, O. Dzyapko, G. A. Melkov, A. A. Serga, B. Hillebrands, and A. N. Slavin, *Nature* **443**, 430 (2006).
- [46] F. J. Dyson, *Phys. Rev.* **102**, 1217 (1956).
- [47] K. P. Sinha and U. N. Upadhyaya, *Phys. Rev.* **127**, 432 (1962).
- [48] J. Callaway and R. Boyd, *Phys. Rev.* **134**, A1655 (1964).
- [49] T. Holstein and H. Primakoff, *Phys. Rev.* **58**, 1098 (1940).
- [50] C. Kittel, *Elementary Statistical Physics* (Wiley, 1961).
- [51] See supplementary material for a detailed discussion.

- [52] A. B. Harris and H. Meyer, Phys. Rev. **127**, 101 (1962).
- [53] R. L. Douglass, Phys Rev **120**, 1612 (1960).
- [54] R. L. Douglass, Phys. Rev. **129**, 1132 (1963).
- [55] M. Guillot, F. Tch  ou, A. Marchand, P. Feldmann, and R. Lagnier, Z. F  r Phys. B Condens. Matter **44**, 53 (1981).
- [56] N. Roschewsky, M. Schreier, A. Kamra, F. Schade, K. Ganzhorn, S. Meyer, H. Huebl, S. Gepr  gs, R. Gross, and S. T. B. Goennenwein, arXiv: 1309.3986 (2013).
- [57] E. G. Spencer and R. C. LeCraw, Phys. Rev. Lett. **4**, 130 (1960).
- [58] E. G. Spencer and R. C. LeCraw, Proc. IEE - Part B Electron. Commun. Eng. **109**, 66 (1962).
- [59] C. Vittoria, P. Lubitz, P. Hansen, and W. Tolksdorf, J. Appl. Phys. **57**, 3699 (1985).
- [60] A. M. Tishin and Y. I. Spichkin, *The Magnetocaloric Effect and Its Applications* (Institute of Physics Pub., Bristol; Philadelphia, 2003).

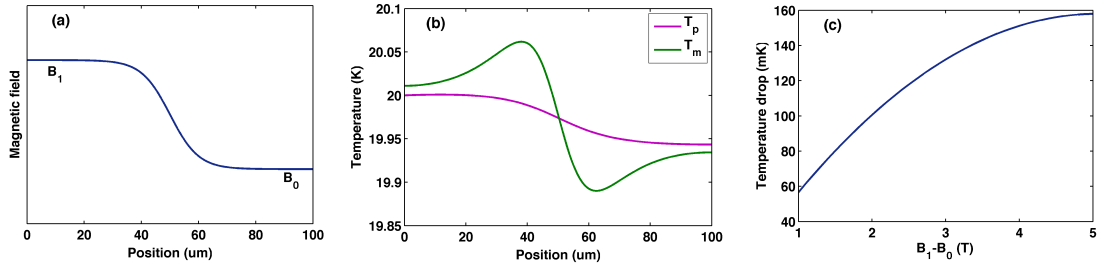


Figure 1. (a) The step-like magnetic field, smeared out as a Fermi-Dirac function. B_0 is fixed to be 0.5T in the following calculation. (b) The temperature distribution of phonons and magnons when $B_1 = 1.5$ T. One end of the sample ($x = 0$) is thermally connected to a reservoir at 20K, and the other end is isolated. (c) The dependence of the phonon temperature difference between the two ends on the difference of the magnetic field when B_0 is set to 0.5T.

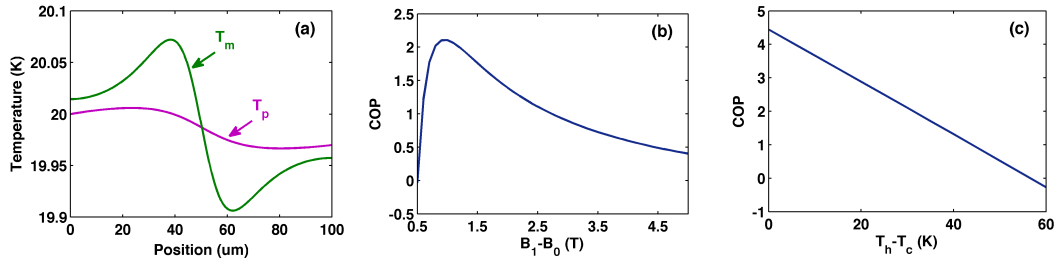


Figure 2. (a) The temperature profiles of phonons and magnons when $T_h = 20$ K , $T_h - T_c = 30$ mK and $(B_0, B_1) = (0.5$ T, 1.5 T). (b) COP versus the change of magnetic field when the temperature difference is fixed at 30mK. The hot-side temperature is fixed at 20K. (c) COP versus the temperature difference when the hot-side temperature is fixed at 20K and the magnetic field is fixed at $(B_0, B_1) = (0.5$ T, 1.5 T).