

Thermal Conductivity of MgO, Al₂O₃, MgAl₂O₄, and Fe₃O₄ Crystals from 3° to 300°K*

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The thermal conductivity from 3° to 300°K of single crystals of the two spinels, diamagnetic MgAl₂O₄ and paramagnetic Fe₃O₄, has been measured and analyzed. The heat is transported by phonons in both cases. The phonon scattering processes in the two crystals are quite similar except that the phonons in Fe₃O₄ are also scattered by the magnetic lattice in its disordered state. The thermal conductivity of Fe₃O₄ falls abruptly with increasing temperature at the 121±2°K transition.

Comparison of results on the double oxide MgAl₂O₄ with those on the single oxides MgO and α-Al₂O₃ indicates the double oxide has an intermediate thermal conductivity. Measurements on crystals of MgO·3.5Al₂O₃ and Mg_xFe_yAl_zO₄ from 3° to 300°K indicate that vacancies and certain impurities can noticeably lower the thermal conductivity of MgAl₂O₄. The effect of vacancies and impurities on Fe₃O₄ is also noticeable, particularly below 30°K.

INTRODUCTION

THE thermal conductivity of transition metal compounds which are electrically nonconducting or are semiconducting has not been studied very extensively in the past. The main interest at present is to study compounds in which most if not all of the thermal energy is transported by phonons, and possibly by magnons. If the thermal conductivity is measured over a wide range of temperatures, particularly through regions of magnetic transitions, or phase transitions, it may be possible to gain some insight into the interaction of phonons with the magnetic moments in the solid.

In this paper a brief summary is given of the thermal conductivity measurements available in the literature for paramagnetic transition metal compounds of the first transition metal series in which anomalous behavior has been seen. A few cases can be found where a magnetic or crystallographic change is associated with a change in thermal conductivity. In the next section an understanding of the thermal conductivity as a function of temperature for pure and impure spinel, MgAl₂O₄, is developed by comparison of this double oxide with the single constituent oxides MgO and α-Al₂O₃. Then these results on diamagnetic MgAl₂O₄ are employed to help understand the thermal conductivity of Fe₃O₄, which has the same spinel crystal structure. The observation is that in Fe₃O₄ and related ferrites there is an additional phonon scattering caused by interaction of the phonons with the magnetic moments.

PREVIOUS WORK

The present measurements on Fe₃O₄ single crystals provide a system in which the interaction of phonons and magnetic moments is particularly pronounced. Before considering these results it is instructive to consider the previous work that has been done on the thermal conductivity of some transition metal compounds. Probably some of the earliest such measure-

ments were reported in 1905 by Kruckenberg¹ on natural crystals of Fe₃O₄ and Fe₂O₃ at 343°K. In 1911 Koenigsberger and Weiss² also reported measurements on a natural crystal of Fe₂O₃ at 303°K. Some average results on ceramic ferrites at 300°K have recently been reported by Smit and Wijn.³

None of these measurements have been made as a function of temperature, and it is the temperature dependence of the thermal conductivity that provides the most useful information about the phonon interactions. In 1917 Bidwell⁴ reported what were probably the first measurements on a transition metal compound as a function of temperature. He studied a ceramic sample of Fe₂O₃ from 430° to 1320°K. Even though his absolute values of the thermal conductivity K are unreasonably low, and his temperature dependence seems wrong for a low porosity sample, he does observe an inflection in the K versus T curve at 980°K. This is quite close to the reported⁵ Curie temperature T_c for α-Fe₂O₃ at 950°K. It seems reasonable to attribute this inflection to the change in phonon scattering with temperature as the sample changes from the ferrimagnetic to the paramagnetic state. His measurements do not extend to temperatures low enough to cover the transition in α-Fe₂O₃ at 260°K.⁶ Measurements of the K of single crystals of Mn and Co ferrite have recently been made between 1.5° and 25°K by Douthett and Friedberg.⁷ In this temperature range none of the samples possess a magnetic or crystallographic transition, so that no anomalies in K versus T were observed. They did, however, observe a change in the phonon scattering as a function of applied magnetic field. Ceramic samples of NiO have been studied

¹ I. Kruckenberg, *Arkiv Matematik, Astoromi, Fysik* **2**, 1 (1905).

² J. Koenigsberger and J. Weiss, *Ann. Physik* **35**, 1 (1911).

³ J. Smit and H. P. J. Wijn, *Advances in Electronics and Electron Physics*, edited by L. Marton (Academic Press Inc., New York, 1954), Vol. 6, pp. 69, 133; *Ferrites* (John Wiley & Sons, Inc., New York, 1959), p. 225.

⁴ C. C. Bidwell, *Phys. Rev.* **10**, 756 (1917).

⁵ K. Endo, *Sci. Repts. Tôhoku Univ.* **25**, 879 (1937).

⁶ L. M. Corliss, J. M. Hastings, and J. E. Goldman, *Phys. Rev.* **93**, 893 (1954).

⁷ D. Douthett and S. A. Friedberg, *Phys. Rev.* **121**, 1662 (1961).

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by Kingery *et al.*⁸ from 420° to 1220°K. These measurements cover the T_N at 525°K,⁹ but they show no anomaly to within the scatter of the data. Single crystals of MnO, and NiO have been measured by Slack and Newman,¹⁰ and single crystals of CoO by Slack¹¹ from 3° to 300°K. Anomalies have been seen at the T_N values of 118°K for MnO,¹² and 290°K for CoO.¹³ Anomalous behavior of the K of single crystals of MnF₂ and CoF₂, measured from 3° to 300°K, has been observed by Slack¹⁴ at their respective T_N values¹⁵ of 67° and 38°K.

PROCEDURE

There are two main problems. The first is to measure the thermal conductivity K of the diamagnetic spinel MgAl₂O₄ versus temperature, and to explain its absolute magnitude and temperature dependence. Since MgAl₂O₄ is a double oxide, the results can be compared with the K measurements on single crystal samples of the constituent oxides, MgO and α -Al₂O₃. The question raised here is whether or not the double oxide has a K significantly different from that of the single constituent oxides.

MgAl₂O₄ was chosen in order to have a comparison for Fe₃O₄. Actually there are a number of diamagnetic spinels to choose from. For example, MgAl₂O₄ and ZnGa₂O₄ both have the spinel structure, and the atomic weights of the two cations in each one are not very different ($\leq 10\%$). MgAl₂O₄ is a normal rather than inverse spinel¹⁶ in that the Al³⁺ ions are on the octahedral sites, and the Mg²⁺ ions are on the tetrahedral ones. It is known¹⁷ that CdGa₂O₄ is a normal spinel, and from the calculations of Miller¹⁸ it can be concluded that ZnGa₂O₄ is also a normal spinel. However, the only single crystals that were available were those of MgAl₂O₄. Natural crystals of MgAl₂O₄ are quite common,¹⁹ and can be obtained through gem and jewelry sources in sizes large enough to enable useful measurements to be made. The technical problems of making large synthetic single crystals of MgAl₂O₄ are quite

TABLE I. Description of the samples studied.^a

Crystal	Run No.	S (cm)	d (cm)	a_0 (Å)
MgO	14	1.24	0.41	4.213 ^b
MgO	38	1.11	0.28	4.213 ^b
MgAl ₂ O ₄	42	0.29	0.18	8.0866 ^c
MgAl ₂ O ₄	54	0.45	0.25	8.0868 ^c
MgO·3.5Al ₂ O ₃	53	1.12	0.40	7.979 ^c
Mg _{0.75} Fe _{0.41} Al _{1.85} O ₄	56	1.12	0.34	8.128 ^c
Mg _{0.73} Fe _{0.33} Al _{1.93} O ₄	62	1.00	0.24	8.117 ^c
Fe ₃ O ₄	57	1.23	0.24	8.398 ^c
Fe ₃ O ₄	37	1.26	0.54	8.397 ^c
Co _{0.07} Fe _{2.93} O ₄	63	0.80	0.24	8.394 ^c

^a S = over-all length of the sample in the direction of heat flow. d = average diameter = $(4A/\pi)^{1/2}$, where A is the cross-sectional area perpendicular to the heat flow. a_0 = lattice constant.

^b See reference 32.

^c Explicitly measured at 300°K for the samples as listed.

complex.²⁰ From the phase diagram studies²¹ it appears to be easier to make nonstoichiometric Mg-Al spinels which are rich in Al₂O₃. Thus single crystals approximating the composition MgO·3.5 Al₂O₃ are available commercially.²² These nonstoichiometric Mg-Al spinels still possess the spinel crystal structure,²⁰ but the metal-ion-to-oxygen-ion ratio is less than 0.75. Hence there are vacancies in the metal ion lattice which noticeably affect the thermal conductivity.

Therefore, the natural, stoichiometric MgAl₂O₄ crystals serve as the most useful comparison for Fe₃O₄. The nonstoichiometric, artificial Mg-Al-spinels provide us with information regarding the effect of large vacancy concentrations and partial inversion of the normal spinel structure on K .

The second problem is to explain the effect of phonon-magnetic moment interactions on the K of Fe₃O₄ by using the results of MgAl₂O₄ as a basis. Fe₃O₄ magnetite, is ferrimagnetic, and is the starting point for many other ferrite materials.³ It has a ferrimagnetic Curie temperature T_c of 848°K,²³ and an internal ordering transition at a transition temperature T_x of 119°K.^{24,25} At all temperatures above T_x , samples of Fe₃O₄ have the spinel crystal structure. In this structure there are both octahedral and tetrahedral sites for the metal ions, where there are 8×1 tetrahedral sites, 8×2 octahedral sites, and 8×4 oxygen sites in a unit cell of 8 molecules. In stoichiometric Fe₃O₄ the tetrahedral sites are occupied by Fe³⁺ ions, while half of the octahedral sites are occupied by the remaining Fe³⁺ ions, and the other half by the Fe²⁺ ions. Thus Fe₃O₄ is an inverse spinel, as pointed out by Verwey and Haayman,²⁶ and con-

²⁰ F. Rinne, Neues Jahrb. Mineral. Geol., beilage band 58A, 43 (1928).

²¹ D. M. Roy, R. Roy, and E. F. Osborn, Am. J. Sci. 251, 337 (1953).

²² Linde Air Products Company, New York, New York.

²³ D. O. Smith, Phys. Rev. 102, 959 (1956).

²⁴ R. W. Millar, J. Am. Chem. Soc. 51, 215 (1929).

²⁵ B. A. Calhoun, Phys. Rev. 94, 1577 (1954).

²⁶ E. J. W. Verwey and P. W. Haayman, Physica 8, 979 (1941).

⁸ W. D. Kingery, J. Francl, R. L. Coble, and T. Vasilos, J. Am. Ceram. Soc. 37(II), 107 (1954).

⁹ J. R. Tomlinson, L. Domash, R. C. Hay, and C. W. Montgomery, J. Am. Chem. Soc. 77, 909 (1955).

¹⁰ G. A. Slack and R. Newman, Phys. Rev. Letters 1, 359 (1958).

¹¹ G. A. Slack, Bull. Am. Phys. Soc. 4, 37 (1958); G. A. Slack, "Proceedings of the International Conference on Semiconductor Physics" (Czechoslovakia Academy of Science, Prague, 1961), p. 630.

¹² S. S. Todd and K. R. Bonnickson, J. Am. Chem. Soc. 73, 3894 (1951).

¹³ G. Assayag and H. Bizette, Compt. rend. 239, 238 (1954).

¹⁴ G. A. Slack, Phys. Rev. 122, 1451 (1961).

¹⁵ J. W. Stout and E. Catalano, J. Chem. Phys. 23, 2013 (1955).

¹⁶ G. E. Bacon, Acta Cryst. 5, 684 (1952).

¹⁷ W. Rudorf and B. Reuter, Z. anorg. u. allgem. Chem. 253, 194 (1947).

¹⁸ A. Miller, Suppl. J. Appl. Phys. 30, 24S (1959).

¹⁹ C. Palache, H. Berman, and C. Frondel, Dana's System of Mineralogy (John Wiley & Sons, Inc., New York, 1944), 7th ed., Vol. 1, p. 689.

TABLE II. Concentration of major impurities in the crystals by quantitative spectrographic analysis.

Crystal	MgO	MgO	MgAl ₂ O ₄	MgAl ₂ O ₄	MgO·3.5 Al ₂ O ₃	Mg _z Fe _y Al ₂ O ₄	Fe ₃ O ₄
Run No.	14	38	42	54	53	56	37, 57
Impurity ^a							
Al	18.4	18.9	<i>m</i>	<i>m</i>	<i>m</i>	<i>m</i>	18.1
Be	18.4	18.4	<18.1	<18.1	<18.1	n.d.	<18.2
Ca	18.3	18.7	17.7	<17.7	17.4	n.d.	<17.9
Cr	17.6	17.6	19.2	19.3	<17.6	low	17.8
Fe	17.6	17.6	18.9	19.2	18.8	<i>m</i>	<i>m</i>
K	18.4	<18.4	17.7	18.3	<17.7	n.d.	<17.9
Li	17.5	17.5	18.2	18.2	18.0	n.d.	<18.1
Mg	<i>m</i>	<i>m</i>	<i>m</i>	<i>m</i>	<i>m</i>	<i>m</i>	17.6
Mn	<17.3	<17.3	18.1	17.9	<17.6	low	17.1
Na	18.7	18.7	18.3	18.3	18.0	n.d.	18.1
Ni	<17.2	<17.2	<17.6	<17.6	<17.6	low	17.6
Si	18.7	17.9	18.2	18.2	19.1	trace	<18.0
Ti	<17.6	<17.6	18.6	18.3	18.5	medium	<17.8
V	<17.3	<17.3	18.8	19.4	<17.6	low	<17.8
Zn	<17.8	<17.8	20.1	19.5	<17.8	low	<18.0
Zr	18.1	<17.7	<17.6	<17.4	<17.4	n.d.	<17.5

^a The concentration of the *j*th type of impurity is ϵ_{pj} in atoms/cm³. The entries in the table are $\log_{10}\epsilon_{pj}$. The letter *m* denotes a major constituent. Only qualitative values are given for the Mg_zFe_yAl₂O₄ spinel because it is very impure. Here n.d. = not detected. In these crystals, the total number of atoms/cm³ is about 1.0×10^{23} . The metal impurities not listed may be present in concentrations less than 10^{18} cm⁻³, except for R-56 for which this limit is about 10^{19} cm⁻³.

firmed by the neutron scattering results of Shull, Wollan, and Koehler.²⁷

SAMPLES

The single-crystal samples that were studied came from a variety of sources. The MgO, Fe₃O₄, and MgO·3.5 Al₂O₃ were synthetic crystals. The MgAl₂O₄ and Mg_zFe_yAl₂O₄ were natural crystals. These are listed in Table I. In addition, Table I contains the x-ray lattice constant at 300°K, the sample size, and the particular run number (e.g., R-) in which the thermal conductivity was measured.

The MgO crystals were supplied by Hansler,²⁸ and were grown from the melt in an arc furnace using carbon electrodes and a self-crucible technique.²⁹ They were transparent, colorless, well-formed, and free of visible defects. The Fe₃O₄ crystals were grown by Horn³⁰ from the melt using the Czochralski technique. These were opaque, possessed a black luster, and contained no second phase inclusions of FeO or Fe₂O₃. The MgO·3.5 Al₂O₃ crystals were obtained from Linde,²² and were made by the Verneuil³¹ process. These had the same outward appearance as the MgO crystals. Two different natural crystals of MgAl₂O₄ both came from Burma. They were transparent and light pink in color, and were in the shape of small, imperfectly formed, single crystal octahedra before cutting to size.

²⁷ C. G. Shull, E. O. Wollan, and W. C. Koehler, Phys. Rev. **84**, 912 (1951).

²⁸ R. L. Hansler, Lamp Division, General Electric Company, Cleveland, Ohio.

²⁹ H. Moissan, Ann. chim. et phys. **4**, 136 (1895).

³⁰ F. H. Horn, J. Appl. Phys. **32**, 900 (1961).

³¹ A. Verneuil, Ann. chim. et phys. **3**, 20 (1904).

The two Mg_zFe_yAl₂O₄ crystals were opaque, water worn, crystals of pleonaste spinel from Queensland, Australia. The chemical analyses indicated a formula of (Ti_wMg_zFe_yAl₂)O₄ for both crystals, where for R-56 $w=0.02$, $x=0.75$, $y=0.41$, $z=1.85$, and for R-62 $w=0.01$, $x=0.73$, $y=0.33$, $z=1.93$. Note that $w+x+y+z=3.0$. The Ti concentration in both is small enough to be neglected. Traces of other minor constituents are listed in Table II. Both pleonaste spinels were single crystals, as verified by x-ray studies, and were virtually free of second phase inclusions. Thin polished sections of the crystals 40 μ thick were transparent and yellow-brown in color. These sections indicated about 0.1% by volume of some unidentified second phase inclusions or pores.

The lattice constants of the natural MgAl₂O₄ crystals are 0.08% larger than the value³² of 8.0800 Å at 297°K for pure MgAl₂O₄. This discrepancy is probably not caused by a deviation from the stoichiometric MgO to Al₂O₃ ratio of 1:1, but rather by the traces of other impurities which, from Table II, occupy about 0.2% of the metal ion lattice sites. The lattice constant of 7.979 Å for the synthetic spinel from Linde is considerably less than that for pure MgAl₂O₄. This difference is caused by the large excess of Al₂O₃ compared to MgO. The Al₂O₃ to MgO ratio was measured by chemical analysis for the present sample as 3.7 ± 0.3 , and thus agrees with the manufacturers value of 3.5. Using the value of $a_0=7.979$ Å and the variation of a_0 of MgO-Al₂O₃ solid solutions found by Roy, Roy, and Osborn²¹ the computed Al₂O₃ to MgO ratio is 3.3. This

³² H. E. Swanson and E. Tagte, National Bureau of Standards Circular No. 539, 1953 (unpublished), Vol. 1, p. 38, 39.

is also in good agreement with the 3.5 value, which will be used hereafter.

The measured a_0 of 8.398 Å for stoichiometric Fe_3O_4 (R-57) agrees quite well with Hagg,³³ who found 8.397 Å, and with Abrahams and Calhoun,³⁴ who found 8.394 Å at 295°K. The iron deficiency in R-37 and the Co content of R-63 account for the slightly smaller a_0 values. The lattice constant of the mixed Mg-Fe-Al spinels lie between the values for MgAl_2O_4 and FeAl_2O_4 , as might be expected. These limiting values³⁵ are $a_0=8.135$ Å for FeAl_2O_4 and 8.080 Å for MgAl_2O_4 . These pleonaste spinels are essentially MgAl_2O_4 with 25% of the Mg^{2+} and about 10% of the Al^{3+} replaced by Fe^{2+} . If the Ti is Ti^{4+} , the charge balance is maintained with practically all of the Fe in the Fe^{2+} state.

Since the effects of chemical impurities play a large part in determining thermal conductivity, a quantitative spectrochemical analysis was made for most of the samples. The impurity concentrations are given in Table II, and are probably accurate to within a factor of 2. For ease in tabulating, the \log_{10} of the impurity concentration is given. This method also allows a quick assessment of the order of magnitude of the impurity concentration. Note that all samples except Fe_3O_4 have an aggregate impurity concentration of at least 10^{19} cm^{-3} , and some are much higher. In the Fe_3O_4 the total concentration is 4×10^{18} and could probably be reduced to around 10^{18} cm^{-3} with improvements in the growing technique. These Fe_3O_4 crystals were grown from iridium crucibles by the Czochralski technique,³⁰ the iridium content is less than 3×10^{17} cm^{-3} . An analysis of R-63 is not given in Table II because it has the same purity as R-37 and R-57 except for the added cobalt.

In addition to foreign chemical impurities variable valence crystals like magnetite can possess large concentrations of vacancies. " Fe_3O_4 " can deviate substantially from an iron-to-oxygen ratio of exactly 3:4. For illustration, the fcc oxygen lattice and the spinel structure is maintained even for $\gamma\text{-Fe}_2\text{O}_3$. This crystal is essentially magnetite in which there are no Fe^{2+} ions,^{33,36} and $\frac{1}{6}$ of the octahedral lattice sites are vacant.³⁷ This corresponds to a fractional concentration of iron vacancies p , of $\frac{1}{6}$. That is, 11.1% of all the possible iron ion sites in the spinel lattice are unoccupied. Thus in magnetite crystals the limits on p are $0 \leq p \leq 0.111$. From the data of Greig *et al.*³⁸ and Darken and Gurry³⁹ the maximum value of p for magnetite at high temperatures in equilibrium under an oxygen partial pres-

sure P_{O_2} of 1 atmosphere is $p=0.034$ for a temperature of 1725°K. In an $\text{O}_2/\text{CO}/\text{CO}_2$ environment at a total pressure of 1 atm produced by introducing pure CO_2 at room temperature into the growing furnace, the maximum value is $p=0.010$ at 1375°K. The "as-grown" magnetite crystal measured in R-37 was analyzed by standard wet chemistry methods in order to determine the $\text{Fe}^{2+}:\text{Fe}^{3+}$ ratio. The fractional vacancy concentration was $p=0.006$, which corresponds to an equilibration temperature of about 1420°K in the $\text{O}_2/\text{CO}/\text{CO}_2$ atmosphere of the growing furnace. This atmosphere corresponds to a partial pressure of oxygen of 1×10^{-4} atm. A section of the R-37 crystal, which is designated here as R-57, was subjected to an annealing of 1500°K in a CO/CO_2 atmosphere with a ratio of 1 part CO to 20 parts CO_2 by volume. This mixture produced an equilibrium partial pressure of oxygen of 1×10^{-8} atm at 1500°K. After this treatment the vacancy concentration is too low to measure by standard chemical techniques. However, an estimate of p can be made using the approach of Flood and Hill.⁴⁰ The value of p , computed from the data of Greig *et al.*³⁸ and making the assumption that the equilibrium constant⁴⁰ $K_1 \ll 10^{-11}$, is $p=5 \times 10^{-6}$. This heat treated crystal was measured in R-57. From Fig. 6 it can be seen that this change in vacancy concentration from 2.6×10^{20} cm^{-3} in R-37 to 2×10^{17} cm^{-3} in R-57 increases the thermal conductivity below 60°K. Crystal R-63 was given a similar anneal at 1620°K in an oxygen pressure of 2×10^{-6} atm. Its vacancy concentration, based on the same calculation as used for pure Fe_3O_4 , is about 8×10^{18} cm^{-3} .

In view of the high content of chemical impurities and iron vacancies, the effect of the naturally occurring isotopes on the thermal conductivity⁴¹ will be quite small in most of the crystals studied. However the Γ values for the pure crystals containing the normally occurring isotopes⁴² are given in Table III. Here Γ for the single element A is defined by

$$\Gamma(A) = \frac{1}{12} \sum_j f_j \left(\frac{m_j - m_A}{m_A} \right)^2, \quad (1)$$

$$m_A = \sum_j f_j m_j,$$

where f_j is the fractional concentration of the j th isotope of mass m_j . Note that this definition of Γ is the $B\Gamma$ of reference 41 with $B = \frac{1}{12}$. This new definition has also been used in reference 14. The Γ values computed for the individual elements⁴² are $\Gamma(\text{Al})=0$, $\Gamma(\text{oxygen})=0.28 \times 10^{-5}$, $\Gamma(\text{Fe})=0.69 \times 10^{-5}$, $\Gamma(\text{Mg})=6.28 \times 10^{-5}$. For a compound⁴³ $A_x B_y C_z$ the composite

³³ G. Hagg, Z. physik. Chem. **B29**, 95 (1935).

³⁴ S. C. Abrahams and B. A. Calhoun, Acta Cryst. **6**, 105 (1953).

³⁵ R. W. G. Wyckoff, *Crystal Structures* (Interscience Publishers, Inc., New York, 1948-1960).

³⁶ E. J. W. Verwey, Z. Krist. Mineral. Petrog. **91**, 65 (1935).

³⁷ K. P. Sinha and A. P. B. Sinha, J. Phys. Chem. **61**, 758 (1957).

³⁸ J. W. Greig, E. Posnjak, H. E. Merwin, and R. B. Sosman, Am. J. Sci. **30**, 239 (1935).

³⁹ L. S. Darken and R. W. Gurry, J. Am. Chem. Soc. **68**, 798 (1946).

⁴⁰ H. Flood and D. G. Hill, Z. Elektrochem. **61**, 18 (1957).

⁴¹ G. A. Slack, Phys. Rev. **105**, 829, 832 (1957).

⁴² D. Strominger, J. M. Hollander, and G. T. Seaborg, Revs. Modern Phys. **30**, 585 (1958).

⁴³ The values of $\Gamma(\text{CaF}_2)$ and $\Gamma(\text{ZnF}_2)$ in reference 14 were incorrectly computed. They should be a factor of 3 larger in order to account for the 3 atoms per molecule. The $\Gamma(\text{Ca})$ and $\Gamma(\text{Zn})$ values are correct.

TABLE III. Parameters for calculating K_I , the thermal conductivity when limited by isotope scattering.^a

Crystal	$\Gamma \times 10^5$	Λ_0/Λ_i^0	$\theta\Lambda_0/TA_i^0$	K_I (w/cm deg)	K_{obs} (w/cm deg)
MgO	4.66	5.6×10^3	8.6×10^4	15	15.
MgAl ₂ O ₄	1.38	2.0×10^3	3.6×10^4	45	1.4
Al ₂ O ₃	0.10	0.14×10^3	0.27×10^4	200	59.
Fe ₃ O ₄	0.88	1.5×10^3	1.8×10^4	22	0.95

^a Γ = isotope scattering parameter. Λ_0/Λ_i^0 = ratio of mean free paths for phonon-phonon and phonon-isotope scattering (see text). K_I = calculated thermal conductivity at 50°K when limited only by isotope scattering. K_{obs} = observed value of the thermal conductivity at 50°K.

Γ , denoted by $\Gamma(A_xB_yC_z)$ is given by

$$\Gamma(A_xB_yC_z) = \frac{x}{(x+y+z)} \left(\frac{m_A}{\bar{m}}\right)^2 \Gamma(A) + \frac{y}{(x+y+z)} \left(\frac{m_B}{\bar{m}}\right)^2 \Gamma(B) + \frac{z}{(x+y+z)} \left(\frac{m_C}{\bar{m}}\right)^2 \Gamma(C),$$

and

$$\bar{m} = (xm_A + ym_B + zm_C)/(x+y+z). \quad (2)$$

This is equivalent to Eq. (12) of Berman, Foster, and Ziman.⁴⁴ By using $(A_xB_yC_z)$ the compound can now be treated as a monatomic material with an average mass \bar{m} and an average volume per atom V_0 .

Equation (2) is only valid if the atoms A , B , C have definite positions in the lattice. This is important, especially in the spinels, since in a random spinel the atoms A and B may be distributed randomly on the tetrahedral and octahedral sites. Then the Γ from Eq. (2) would be much too small. The only samples for which such an effect might be important are the MgO·3.5Al₂O₃, the Mg_xFe_yAl₂O₄, and the Co_{0.07}Fe_{2.93}O₄. The Fe₃O₄ is an inverse spinel where all the cations have the same mass, and the MgAl₂O₄ is a normal spinel. So Eq. (2) should be valid for these latter crystal as well as for MgO and α -Al₂O₃.

EXPERIMENTAL TECHNIQUE

The thermal conductivity of the crystals was measured by a steady state heat flow method in a rod-shaped sample. The power input was measured electrically, and the temperature gradient was measured with a gold-cobalt versus manganin differential thermocouple. The exact details have previously been published.¹⁴

RESULTS

MgO

The present results from 2.9° to 300°K on two single crystals, R-14 and R-38, of MgO are shown in Fig. 1 together with previous data on single crystals from the

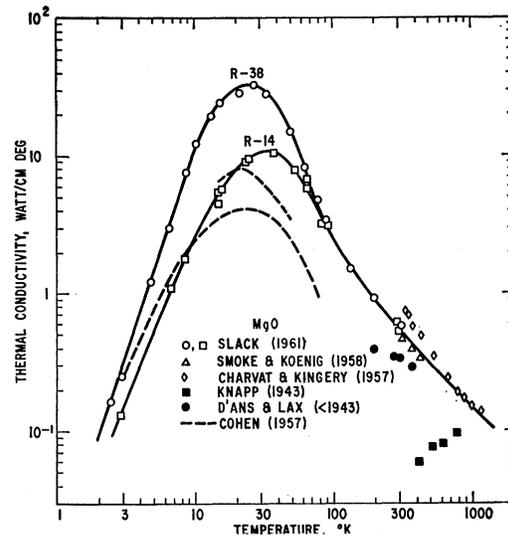


Fig. 1. The thermal conductivity vs temperature of single crystals of MgO. The data of various other authors are also shown.

literature. The agreement with Smoke and Koenig,⁴⁵ and with Charvat and Kingery,⁴⁶ who made measurements at temperatures above 300°K, is satisfactory. The agreement of the values quoted (from an unknown source) by D'Ans and Lax⁴⁷ with the present results is only fair, while Knapp's⁴⁸ values are definitely incorrect. Knapp failed to consider thermal contact resistance effects in his temperature gradient measurements. This failure has also been noted by other authors.⁴⁹ Some of Cohen's thermal conductivity results at low temperatures on single crystals of MgO from the Norton Company⁵⁰ are also shown in Fig. 1. The author wishes to thank her for permission to quote these results, which have previously been mentioned in the literature.^{51,52} The upper of the two dotted curves measured by Cohen refers to a crystal with an effective diameter d of 0.21 cm, and the lower curve to one with $d=0.51$ cm. These are similar to the d values given in Table I. The estimated impurity content of Cohen's crystals was kindly furnished, and the dominant impurities were Al, Fe, and Si, all at a concentration of about 10^{19} cm⁻³. These concentrations are somewhat higher than those given in Table II, particularly for

⁴⁵ E. J. Smoke and J. H. Koenig, College of Engineering Research Bulletin 40, Rutgers University, New Brunswick, New Jersey 1958 (unpublished).

⁴⁶ F. R. Charvat and W. D. Kingery, J. Am. Ceram. Soc. **40**, 306 (1957).

⁴⁷ J. D'Ans and E. Lax, *Taschenbuch für Chemiker und Physiker* (Springer-Verlag, Berlin, Germany, 1949), p. 1125.

⁴⁸ W. J. Knapp, J. Am. Ceram. Soc. **26**, 48 (1953).

⁴⁹ J. Francl and W. D. Kingery, J. Am. Ceram. Soc. **37**(II), 80 (1954).

⁵⁰ The Norton Company, Worcester, Massachusetts.

⁵¹ A. F. Cohen and L. C. Templeton, Bull. Am. Phys. Soc. **2**, 188 (1957).

⁵² A. F. Cohen, *Low Temperature Physics and Chemistry*, edited by J. R. Dillinger (University of Wisconsin Press, Madison, Wisconsin, 1958), p. 385.

⁴⁴ R. Berman, E. L. Foster, and J. M. Ziman, Proc. Roy. Soc. (London) **A237**, 344 (1956).

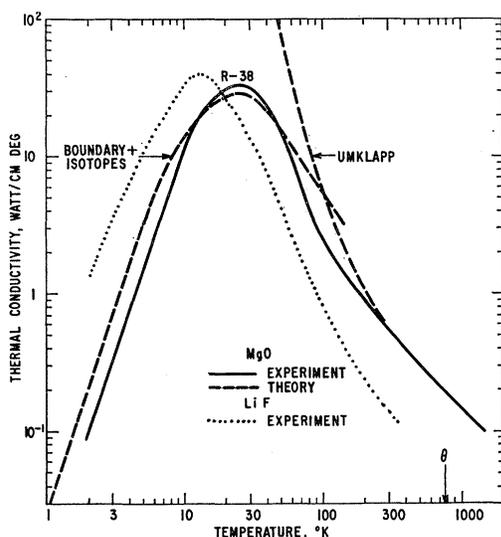


FIG. 2. The thermal conductivity vs temperature of the best MgO crystal compared to the theory. The two dashed curves are the theoretical limits imposed on the phonon heat transport by umklapp scattering, and by a combination of boundary plus isotope scattering. The Debye temperature of MgO is θ . A curve for LiF is shown for comparison.

Fe, and probably account for the lower K_{\max} values found by Cohen.

The curve in Fig. 1 above 300°K also agrees quite well with data⁸ on high purity ceramic samples corrected to zero porosity. This is understandable because the particle size of the ceramic is large compared to the average phonon mean free paths for temperatures above 300°K. Therefore the thermal conductivity in both the single crystal and ceramic samples is determined mainly by phonon-phonon scattering.

The results on the crystal with the highest thermal conductivity, R-38, can be compared with theory. This comparison is shown in Fig. 2. The theoretical calculations for MgO as well as for the other crystals will be made by calculating the thermal conductivity at $T = 10^\circ\text{K}$, 50°K , and $\theta^\circ\text{K}$. The curve-fitting-by-machine-computation method of Callaway⁵³ does not seem appropriate here because the theory of the effects of magnetic impurities and magnetic disorder is not yet well developed. At 10°K the most important phonon scatterers are boundaries or chemical impurities, at 50°K they are isotopes or other mass fluctuations, at $T = \theta$ they are umklapp processes. The ideal K vs T relation for $0^\circ\text{K} \leq T \leq 100^\circ\text{K}$ is determined on an absolute basis using well-developed theories. For $100^\circ\text{K} \leq T \leq \theta$ an empirical K vs T curve is matched to coincide with K_θ , the experimentally determined value of K at $T = \theta$. In those temperature ranges where several scattering processes are equally important approximate averages are used.

For any such calculation it is first necessary to know

⁵³ J. Callaway, Phys. Rev. **113**, 1046 (1959); *ibid.* **122**, 787 (1961).

some of the physical constants for MgO as shown in Table IV. Using these constants a theoretical value K_θ' for the thermal conductivity at the Debye temperature θ can be calculated using the theory of Leibfried and Schlömann.⁵⁴ Assuming the crystal is monatomic, and using the average values of M , θ , V_0 with their units as listed in Table IV, this theory gives

$$K_\theta' = 5.72 \times 10^{-8} (M\theta^2 V_0^{1/3} \gamma^{-2}) \text{ w/cm deg}, \quad (3)$$

where θ is the value at 300°K. The Grüneisen constant γ is taken to be equal to 2.0 for all of the crystals. The value for MgO at 300°K as calculated from the elastic constants,⁵⁵ the specific heat,^{56,57} and the thermal expansion,⁵⁸ is $\gamma = 1.54$. Since K_θ' is being used mainly for comparison of the various crystals, since the γ values for most of them are not known, and since the experimental value of γ is not necessarily the correct one⁵⁹ to use in the theory for K_θ' , $\gamma = 2.0$ seems to be a satisfactory choice. With this choice of variables, the K_θ' values for the four crystals Al_2O_3 , MgAl_2O_4 , Fe_3O_4 , and MgO all lie between 0.54 and 0.35 w/cm deg, and decrease in this order. This very nearly reflects the variation in θ . In the last column of Table IV, the K_θ values are listed. The K_θ show a greater range of values than do the K_θ' . The closest agreement between the computed K_θ' and the experimental K_θ is for MgO, which has the simple NaCl crystal structure, where $K_\theta/K_\theta' = 0.57$. This lattice has very high symmetry, as does the fcc lattice used in the theoretical computation⁵⁴ of K_θ' . The MgAl_2O_4 is also cubic, but possesses a lower symmetry than MgO. The Al_2O_3 is not cubic, but rather has a rhombohedral structure. The ratio $K_\theta/K_\theta' = 0.15$ for these two crystals. The general behavior is that crystals of low symmetry appear to have K_θ values lower than crystals of high symmetry.

Over the whole temperature range studied in MgO phonons are by far the dominant carriers of thermal energy. As can be seen from Fig. 2 the experimental results on the highest conductivity crystal, R-38, can be explained by invoking only three main scattering processes. These are boundary scattering,⁶⁰ isotope scattering,⁴¹ and umklapp scattering.^{54,61} So far the magnitude of the umklapp scattering has been used to calculate K_θ' which agrees reasonably well with K_θ . The theory of umklapp scattering has not been developed to the point where the variation in K with temperature for $T < \theta$ can be calculated. Leibfried and Schlömann⁵⁴ have suggested that K_u/K_θ , where K_u is

⁵⁴ G. Leibfried and E. Schlömann, Nachr. Akad. Wiss. Göttingen, Math.-physik. Kl. **4**, 71 (1954).

⁵⁵ M. A. Durand, Phys. Rev. **50**, 449 (1936).

⁵⁶ W. F. Giauque and R. C. Archibald, J. Am. Chem. Soc. **59**, 561 (1937).

⁵⁷ T. H. K. Barron, W. T. Berg, and J. A. Morrison, Proc. Roy. Soc. (London) **A250**, 70 (1959).

⁵⁸ S. S. Sharma, Proc. Indian Acad. Sci. **A32**, 268 (1950).

⁵⁹ T. H. K. Barron, Nature **178**, 871 (1956); Ann. Phys. **1**, 77 (1957).

⁶⁰ H. B. Casimir, Physica **5**, 495 (1938).

⁶¹ R. E. Peierls, Ann. Physik **3**, 1055 (1929).

TABLE IV. Physical properties of some oxide crystals and values of K_θ and K_θ' .^a

Crystal	V_0 (Å ³)	M (g)	ρ (g/cm ³)	\bar{v}_θ (10 ⁵ cm/sec)	θ_0 (°K)	θ (°K)	K_θ' (w/cm deg)	K_θ w/cm deg
MgO	9.35	20.16	3.58	7.0 ^b	949 ^{b,e}	760 ^h	0.35	0.20
MgAl ₂ O ₄	9.42	20.33	3.58	(7.0)	(900)	900 ⁱ	0.50	0.080
Al ₂ O ₃	8.47	20.39	3.99	6.6 ^c	1010 ^f	950 ^f	0.54	0.080
Fe ₃ O ₄	10.58	33.08	5.19	2.5 ^d	660 ^g	600 ^j	0.37	(0.06)

^a V_0 = average volume per atom, M = average atomic weight per atom, ρ = x-ray density of crystal, \bar{v}_θ = average sound velocity, θ_0 = Debye temperature at 0°K, θ = Debye temperature at 300°K, K_θ' = value of thermal conductivity at $T = \theta$ calculated from Eq. (3), K_θ = experimental value of thermal conductivity at $T = \theta$, () = an approximate value.

^b See references 55, 57.

^c J. B. Wachtman, W. E. Teft, D. G. Lam, and R. P. Stinchfield, J. Research Natl. Bur. Standards 64A, 213 (1960).

^d M. S. Doraiswamy, Proc. Indian Acad. Sci. A25, 413 (1947); M. E. Fine and N. T. Kenney, Phys. Rev. 94, 1573 (1954); H. J. McSkimin, A. J. Williams, R. M. Bozorth, Phys. Rev. 95, 616 (1954); see also reference 93.

^e J. deLaunay, J. Chem. Phys. 24, 1071 (1956).

^f G. T. Furukawa, T. B. Douglas, R. E. McCoskey, and D. C. Ginnings, J. Research Natl. Bur. Standards 57, 67 (1956).

^g J. S. Kouvel, Phys. Rev. 102, 1489 (1956).

^h See reference 57.

ⁱ E. G. King, J. Phys. Chem. 59, 218 (1955).

^j See reference 17; G. S. Parks and K. K. Kelley, J. Phys. Chem. 30, 47 (1926).

the thermal conductivity determined only by umklapp scattering, should be a "universal function" of reduced temperature T/θ . An empirical function of this nature has been constructed by Slack¹⁴ from a combination of the thermal conductivity data on isotopically clean (or nearly so) α -Al₂O₃, H₂, He⁴, and He³. This "universal function" has been matched to the MgO curve in Fig. 2 at $T = \theta$, and is shown as the dashed umklapp curve. From the comparison of the umklapp curve \bar{K}_u with the experimental one in Fig. 2 it can be seen that umklapp scattering is dominant for $100^\circ\text{K} \leq T \leq 1400^\circ\text{K}$, is important for $50^\circ\text{K} \leq T \leq 100^\circ\text{K}$, and is negligible for $T \leq 50^\circ\text{K}$. The difference between the umklapp and experimental curves for $50^\circ\text{K} \leq T \leq 300^\circ\text{K}$ is attributable mainly to isotope scattering.

The theoretical calculation of isotope scattering has been studied by Klemens,^{62,63} Callaway and von Baeyer,^{63,64} and Berman *et al.*⁶⁵ Of these the variational method of Ziman in Berman *et al.* is favored here. It agrees⁶⁴ with the more involved computation by Callaway. Provided that the dimensionless parameter $\theta\Lambda_0/T\Lambda_i^0$ lies in the range 10^3 to 10^7 , the thermal conductivity for $T/\theta \ll 1$ is given^{14,65} approximately by

$$K_I = 0.055 k^{3/2} M^{1/2} h^{-2} \gamma^{-1} N_A^{-1/2} (\Gamma^{-1/2} T^{-3/2}), \quad (4)$$

where k = Boltzmann's constant, h = Planck's constant, N_A = Avogadro's number, and where boundary and umklapp scattering can be neglected. For a temperature of $T = 50^\circ\text{K}$ the calculated value for K_I for MgO as well as of the other crystals is given in Table III. The parameter Λ_0/Λ_i^0 has been calculated from

$$\Lambda_0/\Lambda_i^0 = 7.08 \times 10^3 \theta M \Gamma V_0^{3/2} \gamma^{-2}. \quad (5)$$

Γ is given in Table III, and θ , M , and V_0 with their units in Table IV, and $\gamma = 2.0$. The parameter Λ_0/Λ_i^0

⁶² P. G. Klemens, Proc. Phys. Soc. (London) A68, 1113 (1955).

⁶³ P. G. Klemens, Proc. Phys. Soc. (London) A70, 833 (1957).

⁶⁴ J. Callaway and H. C. von Baeyer, Phys. Rev. 120, 1149 (1960).

⁶⁵ R. Berman, P. T. Nettle, F. W. Sheard, A. N. Spencer, R. W. H. Stevenson, and J. M. Ziman, Proc. Roy. Soc. (London) A253, 403 (1959).

is equal to

$$\frac{\Lambda_0}{\Lambda_i^0} = 2.5 \times 10^4 \left(\frac{\text{phonon m.f.p. for umklapp scattering}}{\text{phonon m.f.p. for isotope scattering}} \right)$$

evaluated at $T = \theta$ for phonons of frequency $\nu = (k\theta/h)$. The ratio Λ_u^0/Λ_i^0 given by Ziman in his book⁶⁶ is just the above factor in the parentheses (). He has changed the definitions of both Λ 's so as to eliminate the factor of 2.5×10^4 . From Table III it can be seen that $\theta\Lambda_0/T\Lambda_i^0$ is in the appropriate range at 50°K so that Eq. (4) is valid for all of the crystals. The calculated value of K_I at 50°K for MgO is in good agreement with the observed value, as can be seen from Table III and also from Fig. 2. This very good agreement is somewhat fortuitous because crystal R-38 still has an over-all impurity concentration of about $2 \times 10^{19}/\text{cm}^{-3}$, as shown in Table II. If these impurities were eliminated, it is estimated that the maximum thermal conductivity at 25°K might increase by a factor of 2.

From Fig. 2 it can be seen that the curve for R-38 comes close to the thermal conductivity limit imposed by boundary scattering for $T < 5^\circ\text{K}$. The dashed curve labeled "boundary + isotopes" was computed by graphically combining (a) the low temperature effects of boundary and isotope scattering calculated using the method of Slack and Glassbrenner⁶⁷ with (b) the isotope scattering calculated from Eq. (4). It was assumed that the pure boundary scattering for $T < 5^\circ\text{K}$ was for a perfectly diffuse scattering of phonons from the crystal surface. Hence the mean free path is equal to the sample diameter d in Table I. The fact that the thermal conductivity is lower than the boundary limit is attributed to the effects of the residual chemical impurities. This rather pronounced effect of impurities at temperatures below the maximum in the \bar{K} vs T curve has been noticed before in other crystals by

⁶⁶ J. M. Ziman, *Electrons and Phonons* (Oxford University Press, New York, 1960), p. 311.

⁶⁷ G. A. Slack and C. Glassbrenner, Phys. Rev. 120, 782 (1960).

Slack,^{10,14,41} Pohl,⁶⁸ Rosenberg and Sujak,⁶⁹ Keyes,⁷⁰ and Klein.⁷¹ This effect of impurities can be seen in Fig. 1 for crystal R-14 where the maximum K is lower than in R-38, and where the K at 3°K is a factor of 6 below the boundary scattering limit. Crystal R-14 is less "pure" than R-38. The major difference between the impurities in R-14 and R-38 that would cause this K difference would seem to be the Si and Zr listed in Table II. These might be expected to fit less well into the MgO lattice than the Ca and Al impurities which, incidentally, are more prevalent in R-38 than in R-14.

The general result for MgO is that crystal R-38 is sufficiently free of chemical impurities so that its thermal conductivity approaches the theoretical limits. A comparison of the results for MgO in R-38 with data for the alkali halide LiF is instructive. A composite curve of K versus T for a single crystal of LiF with a diameter of 0.79 cm and with the Li and F isotopes present in their natural abundances⁴² is shown dotted in Fig. 2. This curve has been constructed using the data available in the literature.^{44,65,72-74} The two curves have very nearly the same shape. The thermal conductivity ratio at some chosen temperature is denoted by $[K(\text{MgO})/K(\text{LiF})]$. Experimentally this ratio is 0.1 from 2° to 10°K, and from 30° to 300°K is nearly constant at a value of 3 to 4. MgO and LiF both have the cubic NaCl structure, and have similar phonon spectra.⁵⁷ Their main differences in K should be caused by M , θ , V_0 , Γ , and d , and perhaps by the fact that MgO is a divalent crystal while LiF is monovalent. From 2° to 10°K the thermal conductivity K_B is determined by boundary scattering for both crystals. For LiF $d = 0.79$ cm, $V_0 = 8.06 \text{ \AA}^3$, $\theta_0 = 743^\circ\text{K}$.⁷⁵ Thus from 2° to 10°K the calculated ratio is $[K_B(\text{MgO})/K_B(\text{LiF})] = 0.2$ compared to the observed 0.1.

From 30° to 60°K the thermal conductivity is determined by isotope scattering. For LiF $M = 12.97$ g, and Γ has a value⁴² of $\Gamma = 1.70 \times 10^{-5}$. This yields a calculated value of $[K_I(\text{MgO})/K_I(\text{LiF})] = 1.6$. For $T > 200^\circ\text{K}$ the conductivity K_u is determined by umklapp scattering, and at temperatures near and above $T = \theta$ one has $K_u = (\theta/T)K_\theta$. This means that for $T > 200^\circ\text{K}$ the calculation gives $[K(\text{MgO})/K(\text{LiF})] = 3.4$, which is only twice the value computed at lower temperatures. For both K_I and K_u a value of $\gamma = 2$ has been assumed for LiF and MgO. The general agreement of the computed ratio from 30° to 300°K with the observed one of 3 to 4 indicates that there is no appreciable difference

in the application of the theory to monovalent or divalent crystals. The conclusion is that in the application of the same theory to Al_2O_3 , MgAl_2O_4 , and Fe_3O_4 no explicit account of the various valences need be taken.

$\alpha\text{-Al}_2\text{O}_3$

Many previous measurements of the thermal conductivity of $\alpha\text{-Al}_2\text{O}_3$ have been made, and therefore such measurements were not repeated in the present series of experiments. A K vs T curve for single crystals has been constructed from the results of Berman *et al.*⁷⁶ from 3° to 93°K, Berman⁷⁷ from 3° to 200°K, Ballard and McCarthy⁷⁸ from 300° to 350°K, Smoke and Koenig⁴⁵ from 315° to 420°K, Weeks and Seifert⁷⁹ at 340°K, Lee and Kingery⁸⁰ from 330° to 1300°K, Kingery *et al.*⁸ from 370° to 2070°K, and Charvat and Kingery⁴⁶ from 500° to 1500°K. The anisotropy in K for these rhombohedral crystals of $\alpha\text{-Al}_2\text{O}_3$ has been measured accurately by Smoke and Koenig⁴⁵ who obtained 1.08 for the ratio of K_c , measured along the c axis, to K_a , measured in the a - b plane. Jannettaz⁸¹ found $K_c/K_a = 1.18$, while the results of Ballard and McCarthy⁷⁸ give 1.11 at 300°K. Since this anisotropy is small, it has been neglected in determining the resultant curve.

The composite K vs T curve for $\alpha\text{-Al}_2\text{O}_3$ is shown in Fig. 3 together with the data for the highest conductivity samples of MgO and MgAl_2O_4 . The experimental curve for $\alpha\text{-Al}_2\text{O}_3$ is reasonably well explained by assuming that only umklapp scattering¹⁴ and boundary scattering⁷⁶ are important. The K , as limited by isotope scattering, is shown in Table III for Al_2O_3 for 50°K. Previous authors^{41,44} have pointed out that the isotope scattering in $\alpha\text{-Al}_2\text{O}_3$ can be neglected. Since many of the K measurements^{46,76,78,79} have been made on Linde²² crystals, a spectrographic analysis of a random piece of Linde $\alpha\text{-Al}_2\text{O}_3$ crystal was made. The most abundant chemical impurities found were Na and Si at about $1 \times 10^{18} \text{ cm}^{-3}$. Thirteen other common metallic impurities were checked for and, if present, were each $\leq 5 \times 10^{17} \text{ cm}^{-3}$. This low concentration of chemical impurities accounts for the fact that the expected exponential rise in K at low temperatures can be observed.

The main differences in the curves for MgO and $\alpha\text{-Al}_2\text{O}_3$ in Fig. 3 are caused by crystal symmetry effects which make $K(\text{MgO}) > K(\text{Al}_2\text{O}_3)$ for $300^\circ\text{K} \leq T \leq 1500^\circ\text{K}$, and by isotope scattering in the MgO which makes $K(\text{MgO}) < K(\text{Al}_2\text{O}_3)$ for $30^\circ\text{K} \leq T \leq 100^\circ\text{K}$.

⁶⁸ R. O. Pohl, Phys. Rev. **118**, 1499 (1960).

⁶⁹ H. M. Rosenberg and B. Sujak, Phil. Mag. **5**, 1299 (1960).

⁷⁰ R. W. Keyes, Phys. Rev. **122**, 1171 (1961).

⁷¹ M. V. Klein, Phys. Rev. **122**, 1393 (1961); *ibid.* **123**, 1977 (1961).

⁷² S. S. Ballard, K. A. McCarthy, and W. C. Davis, Rev. Sci. Instr. **21**, 905 (1950).

⁷³ R. Srinivasan, J. Indian Inst. Sci. **A37**, 200 (1955).

⁷⁴ R. L. Sproull, M. Moss, and H. Weinstock, J. Appl. Phys. **30**, 334 (1959).

⁷⁵ T. H. K. Barron and J. A. Morrison, Can. J. Phys. **35**, 799 (1957).

⁷⁶ R. Berman, E. L. Foster, and J. M. Ziman, Proc. Roy. Soc. (London) **A231**, 130 (1955).

⁷⁷ R. Berman, Proc. Phys. Soc. (London) **A65**, 1029 (1952).

⁷⁸ S. S. Ballard and K. A. McCarthy, J. Opt. Soc. Am. **41**, 1062 (1951).

⁷⁹ J. L. Weeks and R. L. Seifert, Rev. Sci. Instr. **24**, 1054 (1953).

⁸⁰ D. W. Lee and W. D. Kingery, J. Am. Ceram. Soc. **43**, 594 (1960).

⁸¹ E. Jannettaz, Compt. rend. **114**, 1352 (1892).

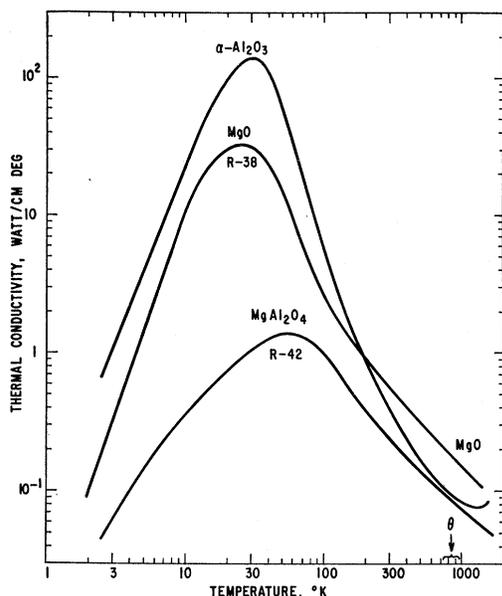


FIG. 3. A comparison of the experimental results of thermal conductivity vs temperature for the best crystals of α -Al₂O₃, MgO, and MgAl₂O₄. All three Debye temperatures, θ , fall within the range indicated.

MgAl₂O₄

Two natural crystals of MgAl₂O₄ were measured, and the results are shown in Fig. 3. These crystals, R-42 and R-54, have very nearly the same thermal conductivity from 100° to 300°K. At 300°K the value of $K=0.24$ w/cm deg. The only previous results in the literature on MgAl₂O₄ are some by Anderson⁸² on ceramic samples from 370° to 1050°K. The agreement between the ceramic samples and the single crystals is quite good, as can be seen in Fig. 4.

The analysis of the results on MgAl₂O₄ in terms of the simple theory used for the MgO is not very satisfactory because the crystals are quite impure. The total impurity content, as given in Table II, is about 0.6×10^{20} cm⁻³ for R-42 and 1×10^{20} cm⁻³ for R-54. That is, 0.2% of the metal atoms in the crystal are foreign atoms. This concentration of impurities, however, will not have much effect on the value of K_θ , which is 0.080 w/cm deg, and is listed in Table IV. Thus MgAl₂O₄ will serve as a useful guide to estimate K_θ for Fe₃O₄. It is assumed that the experimental to theoretical ratio K_θ/K'_θ for Fe₃O₄ is the same as for MgAl₂O₄ since they both have the same crystal structure and somewhat similar atomic masses. This assumption leads to K for Fe₃O₄ of 0.06 w/cm deg, which is not an implausible extrapolation of the curve in Fig. 8.

The umklapp and boundary scattering limits have been computed in the same manner as they were for MgO. These limits are shown in Fig. 4. The experi-

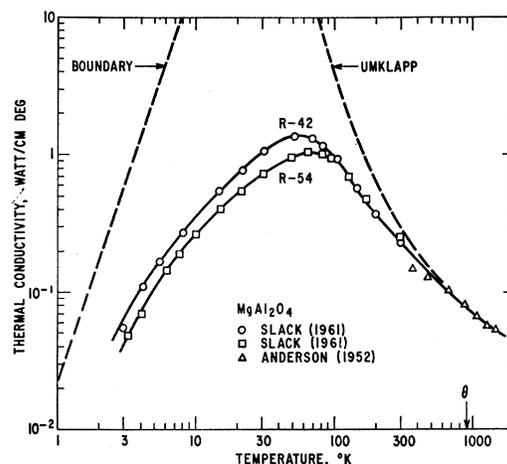


FIG. 4. The thermal conductivity vs temperature of two natural crystals of MgAl₂O₄. The data above 300°K by Anderson are for a ceramic sample. The dashed curves are the theoretical limits calculated for umklapp scattering and boundary scattering.

mental curve of K vs T begins to fall markedly below these limits for $T < 200^\circ\text{K}$. Notice that the limit on K imposed by isotope scattering in Table III is 45 w/cm deg at 50°K. This limit falls way above the graph in Fig. 4.

Impurities in MgAl₂O₄

Of the impurities listed in Table II for R-42 and R-54 the V, Ti, Cr, and Fe impurities may be the ones most responsible for the low thermal conductivity at low temperatures. It can be concluded that certain paramagnetic impurities can have a large effect on K at low temperatures from the results of Rosenberg and Sujak.⁶⁹ Since neither the charge state nor the lattice environment of these trace impurities in the present samples is known, let us consider first the total content per cm³ of the transition metal impurities that are listed as present in Table II. Denote this total concentration as C_1 . For R-42 $C_1=4 \times 10^{19}$ cm⁻³, and for R-54 $C_1=6 \times 10^{19}$ cm⁻³. The total concentration of nontransition metal impurities is denoted by C_2 . The C_2 values are 1×10^{20} cm⁻³ for R-42 and 4×10^{19} for R-54. Zinc accounts for at least 80% of these C_2 values. However, this high concentration of Zn in both crystals will not affect the thermal conductivity appreciably because the Zn enters the tetrahedral sites in place of the Mg and essentially forms a mixed crystal of MgAl₂O₄-ZnAl₂O₄. Both of these are normal spinels of nearly the same lattice parameter, so that the phonon scattering by the Zn is mainly due to the mass difference. The effect of this small amount of Zn (0.2%) on the thermal conductivity is thus^{83,84} quite small. For example $\Gamma=4 \times 10^{-4}$ due to the mass fluctuation produced by the Zn in R-42. It will be noticed later in

⁸² As quoted in reference 8.

⁸³ W. S. Williams, Phys. Rev. **119**, 1021 (1960).

⁸⁴ A. M. Toxen, Phys. Rev. **122**, 450 (1961).

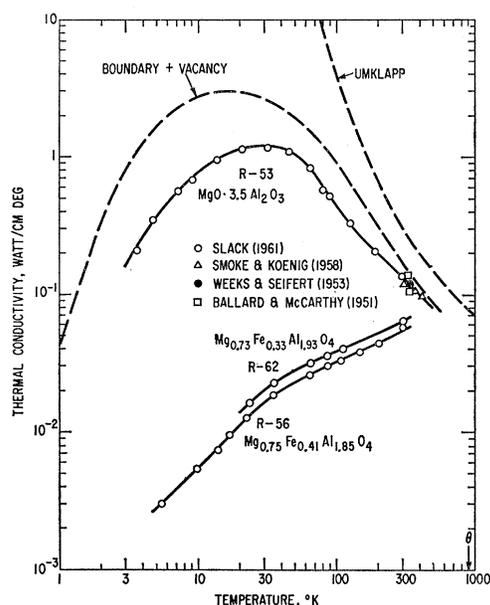


Fig. 5. The thermal conductivity vs temperature of a synthetic $\text{MgO} \cdot 3.5 \text{Al}_2\text{O}_3$ spinel crystal and two natural $\text{Mg}_x\text{Fe}_y\text{Al}_2\text{O}_4$ crystals. The literature data are for synthetic spinel. The dashed curves are the theoretical limits for the synthetic spinel computed for boundary, vacancy, and umklapp scattering.

Fig. 9 that the "best-fit" thermal resistivity versus concentration curve passes close to the transition metal concentration C_1 for both R-42 and R-54. This agrees with the present conclusion that the transition metal impurities rather than the Zn are primarily responsible for the low thermal conductivity for $T < 50^\circ\text{K}$.

The K versus T curve for R-42 is also plotted in Fig. 3. At temperatures above 300°K the K of MgAl_2O_4 is nearly the same as that for $\alpha\text{-Al}_2\text{O}_3$, and both of these are lower than MgO , probably for symmetry reasons as previously mentioned. For temperatures $3^\circ\text{K} \leq T \leq 300^\circ\text{K}$ the high concentration of transition metal impurities in the MgAl_2O_4 is responsible for the fact that its thermal conductivity is much less than either that of MgO or $\alpha\text{-Al}_2\text{O}_3$. If the crystal boundaries and the naturally occurring isotopes in MgAl_2O_4 were the chief limitation on its thermal conductivity at low temperatures, then its K vs T curve in Fig. 3 should lie between those for MgO and $\alpha\text{-Al}_2\text{O}_3$. Therefore it is concluded that the thermal conductivity of the pure double oxide MgAl_2O_4 is not significantly different from what would have been predicted by a crude arithmetic average of the results for the single constituent oxides MgO and $\alpha\text{-Al}_2\text{O}_3$.^{84a}

Grossly Impure MgAl_2O_4

Since Fe_3O_4 crystals can possess large concentrations of chemical impurities or vacancies, it is instructive to

^{84a} Note added in proof. Recent measurements on a higher purity synthetic sample of MgAl_2O_4 gave a maximum value of K of 4.5 w/cm deg at 30°K .

see what effect such large concentrations have on the thermal conductivity of MgAl_2O_4 . For this purpose K measurements were made on crystals of $\text{Mg}_x\text{Fe}_y\text{Al}_2\text{O}_4$ and $\text{MgO} \cdot 3.5\text{Al}_2\text{O}_3$. The first contains a large number of Fe impurities, and the second metal ion vacancies, but both have essentially the same crystal structure and lattice dynamics as the parent MgAl_2O_4 . Figure 5 shows the results on $\text{Mg}_x\text{Fe}_y\text{Al}_2\text{O}_4$ (R-56 and R-62), and $\text{MgO} \cdot 3.5\text{Al}_2\text{O}_3$ (R-53). The agreement of the results on R-53 with previous data on similar single crystals by Smoke and Koenig,⁴⁵ Weeks and Seifert,⁷⁹ and Ballard and McCarthy^{78,85} is good. The analysis of these results in terms of umklapp, isotope, and boundary scattering is very nearly the same as for MgAl_2O_4 , and will not be repeated. Assuming $\theta = 900^\circ\text{K}$, the same as for MgAl_2O_4 , the umklapp limit and pure boundary scattering limit ($T \leq 2^\circ\text{K}$) are given in Fig. 5.

The most noticeable feature for R-56 (and similarly for R-62) is that its thermal conductivity at 300°K , $K = 0.055$ w/cm deg, is 0.23 of that for MgAl_2O_4 , and that K continuously decreases with decreasing temperature. These samples were quite good natural single crystals, and were free of flaws and inclusions. The low value of K can only be attributed to the large concentration of Fe^{2+} impurity. Since FeAl_2O_4 is a normal spinel⁸⁶ as is MgAl_2O_4 , the Fe^{2+} in $\text{Mg}_x\text{Fe}_y\text{Al}_2\text{O}_4$ probably occurs on the tetrahedral sites.

The distribution of the Fe impurity in the $\text{Mg}_x\text{Fe}_y\text{Al}_2\text{O}_4$ can be estimated from Miller's¹⁸ calculations. The result is for R-56 is $\text{Fe}_{0.41}\text{Mg}_{0.60}[\text{Mg}_{0.15}\text{Al}_{1.85}]_2\text{O}_4$. The mass fluctuation parameter Γ for this distribution is calculated to be $\Gamma = 5.9 \times 10^{-3}$. At temperatures above 30°K some thermal resistance is provided by this mass fluctuation. By comparison with the following analysis on $\text{MgO} \cdot 3.5\text{Al}_2\text{O}_3$ a value of $\Gamma = 5.9 \times 10^{-3}$ leads to a noticeable reduction in K only at high temperatures. At 500°K the value of K for R-56 is calculated as 0.07 w/cm deg. This value would be reasonable extrapolation of the results for R-56 in Fig. 4. At 300°K and below, the mass fluctuation scattering cannot be the explanation of the low thermal conductivity in R-56 or R-62.

The most plausible reason for the low thermal conductivity in R-56 is suggested by Orbach's⁸⁷ analysis of Fe-doped $\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}$.⁶⁹ In this latter crystal the Fe^{2+} , which has a $3d^6$ configuration, is surrounded by six H_2O molecules which produce a tetragonal environment with a slight orthorhombic distortion. Orbach assumes that there is a low lying orbital doublet which is split by the orthorhombic distortion into two levels separated by a small energy $k\Delta$, where $\Delta = 4^\circ\text{K}$. In the spinel lattice, the four oxygen ions around the tetrahedral site form an undistorted tetrahedron. The tetra-

⁸⁵ S. S. Ballard and K. A. McCarthy, *American Institute of Physics Handbook*, edited by D. E. Gray (McGraw-Hill Book Company, Inc., New York, 1957), p. 4-76.

⁸⁶ T. F. W. Barth and E. Posnjak, *Z. Krist.* **82**, 325 (1932).

⁸⁷ R. Orbach, *Phil. Mag.* **5**, 1303 (1960).

hedron remains undistorted even if the oxygen parameter u is different from 0.375. For MgAl₂O₄, $u = 0.387$,²¹ and for FeAl₂O₄, $u = 0.390$.⁸⁶ The energy level scheme of Fe²⁺ in various crystalline fields is given by McClure.⁸⁸ In the tetrahedral field, the orbital doublet is lower than the triplet. The first five electrons half fill the d shell. The sixth electron goes into the lower energy doublet state producing a twofold degeneracy. This doublet of Fe²⁺ in MgAl₂O₄ may be split by spin-orbit interactions.⁸⁹ However, the level separation Δ is not yet known for this case. Thus MgAl₂O₄ and ZnSO₄·7H₂O both appear to possess a nearly degenerate doublet ground state for Fe²⁺. Some comparison of the thermal conductivity data can be made. For a ZnSO₄·7H₂O crystal with $C_1 = 2.9 \times 10^{20}$ cm⁻³ of Fe²⁺, Rosenberg and Sujak⁶⁹ measured the thermal resistivity at 10°K, $W(10^\circ\text{K})$, 13.3 cm deg/w. W varied as $\sim T^{+0.5}$. The ratio $[W(10^\circ\text{K})/C_1] = 4.6 \times 10^{-20}$ cm⁴ deg/w. In the present case $C_1 = 6.1 \times 10^{21}$ cm⁻³ for R-56 and 4.9×10^{21} cm⁻³ for R-62. Therefore $[W(10^\circ\text{K})/C_1] = 3 \times 10^{-20}$ cm⁴ deg/w for R-56, and one obtains a similar extrapolated value for R-62 from Fig. 5. At 10°K the value of W varied as T^{-1} .

The close agreement of these two results on entirely different crystals is probably fortuitous, especially since W has a different temperature dependence in the two cases. However the agreement of the two $[W(10^\circ\text{K})/C_1]$ values to within an order of magnitude tends to indicate that the Fe²⁺ scattering process is similar in both cases.

In the temperature region between 100° and 500°K the K of MgO·3.5Al₂O₃ is noticeably lower than that of MgAl₂O₄. This difference is clearly not caused by chemical impurities since the natural MgAl₂O₄ crystals are less pure than the synthetic MgO·3.5Al₂O₃. The difference appears to be caused by vacancies. Not all of the metal ion sites in the spinel lattice are occupied by the metal ions in MgO·3.5Al₂O₃. Vacancies occur, and these are probably on the octahedral sites.⁸⁷ [Even if the vacancies are on the tetrahedral sites or are distributed at random, the numerical values in the following argument will not be appreciably altered.] Thus for each 4 oxygen O²⁻ ions there are 0.35 Mg²⁺ ions and 0.65 Al³⁺ ions in the 1 tetrahedral site, and 1.78 Al³⁺ ions in the 2 octahedral sites. The other 0.22 octahedral sites are vacant. Let C_3 denote the concentration of vacancies. Therefore $C_3 = 1.6 \times 10^{21}$ cm⁻³ for R-53, which is 6 times larger than C_3 for the R-37 sample of Fe₃O₄. If neither the Mg and Al ions on the tetrahedral sites nor the Al ions and the vacancies on the octahedral sites are ordered, their random distribution will give rise to a mass fluctuation scattering. The Γ value for this random distribution of masses is calculated to be $\Gamma = 4.4 \times 10^{-3}$, 99% of this value is due to the vacancies. The phonon scattering caused by this

mass fluctuation can be calculated for low temperatures from Eq. (4). At 50°K this yields $K_T = 2.4$ w/cm deg, and $\theta\Lambda_0/T\Lambda_i^0 = 1.1 \times 10^7$. At high temperature the theory of Ambegaokar⁹⁰ can be used. For $T = \theta$ it will be assumed that the umklapp scattering leads to the same value of K_θ as measured for MgAl₂O₄. Ambegaokar's theory applies for temperatures near $T = \theta$. The highest temperature at which comparison is possible from the available data in Figs. 4 and 5 is 500°K. Here the thermal resistivity value for MgAl₂O₄ is 7.4 cm deg/w. The additional thermal resistivity that would be caused by the mass fluctuation scattering would be

$$\Delta W = 4\pi^2 V_0 \Gamma h^{-1} \bar{v}_\theta^{-2}, \quad (6)$$

where \bar{v}_θ is 7.0×10^5 cm/sec from Table IV. Using $\Gamma = 4.4 \times 10^{-3}$ (which is $\frac{1}{2}$ of that as defined by Ambegaokar), ΔW is 4.3 cm deg/w. Thus the computed W for MgO·3.5Al₂O₃ at 500°K is $7.4 + 4.3 = 11.7$ cm deg/w, as compared to the observed value of 12.5 cm deg/w. Such agreement is considered satisfactory, especially since only the mass difference between the Al ions and the vacancies was considered in computing Γ . Any lattice distortion or changes in the elastic constants of the linkages⁶² would serve to increase Γ . The lattice distortion around a metal ion vacancy, however, is probably small because the lattice structure is determined mainly by the fcc packing of the large radius O²⁻ ions. The metal ion radius is about 40% that of the O²⁻ ions, and they fit into the "interstices" in the oxygen lattice. Thus the mass difference may be the largest part of the vacancy effect.

The dashed curve labeled "boundary+vacancy" scattering in Fig. 5 was drawn for the combination of vacancy scattering calculated from Eqs. (4) and (6) and the boundary-plus-vacancy effect calculated by the method of Slack and Glassbrenner.⁶⁷ It is apparent that this calculated limiting curve for K of MgO·3.5Al₂O₃ correctly predicts the general behavior, particularly from 50° to 500°K. Below 50°K the chemical impurities produce a K still lower than this limiting curve.

Fe₃O₄

Now that the behavior of pure and impure MgAl₂O₄ is understood in terms of the theory, it is possible to use the results for understanding Fe₃O₄. The experimental data for the Fe₃O₄ runs R-37, R-57, and R-63 are shown in Figs. 6 and 7. The fact that Fe₃O₄ has an orthorhombic crystal structure⁸⁴ at temperatures below T_x means that there are three principal thermal conductivities. Since the distortion from cubic is small, these three conductivities should be nearly equal. Therefore no effort was made to produce untwinned crystals,⁹¹ and the results in Figs. 6 and 7 represent the average K for $T < T_x$. Figure 6 also contains the data of Kruckenberg¹ on natural Fe₃O₄ crystals, Smit

⁸⁸ D. S. McClure, *Solid-State Physics*, edited by F. Seitz and D. Turnbull (Academic Press Inc., New York, 1959), Vol. 9, p. 399.

⁸⁹ W. Low and M. Weger, *Phys. Rev.* **118**, 1119 (1960).

⁹⁰ V. Ambegaokar, *Phys. Rev.* **114**, 488 (1959).

⁹¹ W. C. Hamilton, *Phys. Rev.* **110**, 1050 (1958).

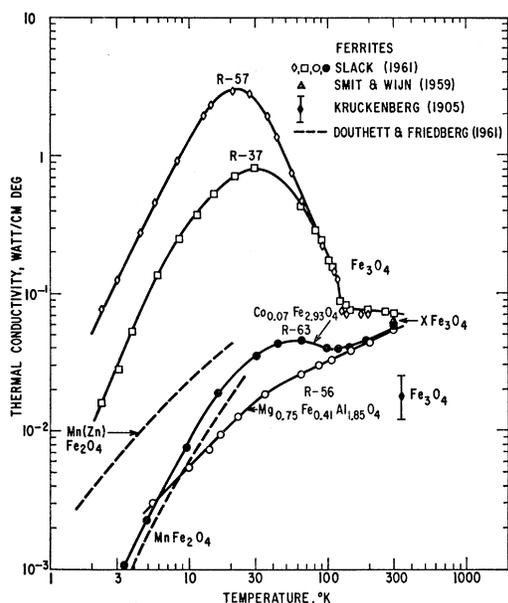


FIG. 6. The thermal conductivity vs temperature for various single-crystal ferrites. Data for a pleonaste spinel crystal, R-56, are shown for comparison.

and Wijn³ on ceramic ferrite samples, and Douthett and Friedberg⁷ on some single crystal ferrites. It is clear from the magnitude and temperature dependence of K in Fig. 6 that the explanation used for MgO , $\alpha\text{-Al}_2\text{O}_3$, and MgAl_2O_4 will have to be modified.

The most unusual feature for Fe_3O_4 is the nearly temperature independent K found for the range $119^\circ\text{K} \leq T \leq 300^\circ\text{K}$. As mentioned previously, 119°K is the transition temperature T_x below which an ordering of the Fe^{2+} and Fe^{3+} ions on the octahedral sites takes place.^{25,26,91,92} At this temperature the thermal conductivity exhibits an abrupt rise by a factor of 1.6. The details of this rise are shown more clearly in Fig. 7 where a linear plot of K vs T has been made. The transition temperature and its uncertainty from Fig. 7 is $121 \pm 2^\circ\text{K}$, which is essentially equal to the $119.4 \pm 0.3^\circ\text{K}$ found by Calhoun²⁵ from electrical measurements on synthetic crystals. As the temperature decreases below 119°K the K rises, reaches a maximum, and then decreases, a behavior similar to that of the other diamagnetic crystals studied.

The thermal conductivity results for Fe_3O_4 can be explained satisfactorily by assuming that phonons are the only important carriers of thermal energy. These phonons are assumed to be scattered by the standard processes found in the diamagnetic crystals, namely: other phonons (umklapp scattering), isotopes, vacancies, chemical impurities, and boundaries. One additional scattering mechanism then appears to be necessary. This is scattering of phonons from local disorder in the lattice formed by the magnetic moments of the

paramagnetic ions. This same model was found sufficient to explain the results for antiferromagnetic MnF_2 and CoF_2 .¹⁴ Any possible magnon heat transport¹⁴ appears to be unnoticeable in Fe_3O_4 , MnF_2 , CoF_2 , or in MnO , CoO , and NiO ,^{10,11} using the present measuring techniques.

The electrical conductivity as a function of temperature was measured for several synthetic Fe_3O_4 crystals very similar to R-37 and R-57. The results were almost the same as those reported by Calhoun.²⁵ The electrical conductivity is sufficiently low at all temperatures so that any electronic thermal conductivity contribution K_e is negligible. This lack of a significant K_e term is borne out by the agreement of the present data at 300°K with that of Smit and Wijn³ on other ferrites. At room temperature these other ferrites have very low electrical conductivities³ compared to Fe_3O_4 .

The upper limits on the conductivity of Fe_3O_4 imposed by umklapp and boundary scattering are shown in Fig. 8. The boundary limit for R-57 was calculated assuming diffuse reflection of phonons at the surface with $d=0.24$ cm. The umklapp limit was drawn using the value of $K_\theta=0.06$ w/cm deg estimated in the section under MgAl_2O_4 . The value of Γ for pure, stoichiometric Fe_3O_4 is so small that the K_I limit (see Table III) lies above the graph in Fig. 8. Therefore the results for the very nearly stoichiometric crystal, R-57, which has a vacancy concentration of about $2 \times 10^{17} \text{ cm}^{-3}$ as computed in the section entitled Samples, do not approach the simple limits found for diamagnetic crystals for $T < \theta$. Only at temperatures near $T=\theta$ where phonon-phonon scattering is dominant does the K for Fe_3O_4 appear to approach that for diamagnetic crystals.

For $T > 119^\circ\text{K}$ the Fe^{2+} and Fe^{3+} ions on the octahedral lattice sites are distributed randomly, and there is a rapid exchange of electrons from one Fe^{2+} to the

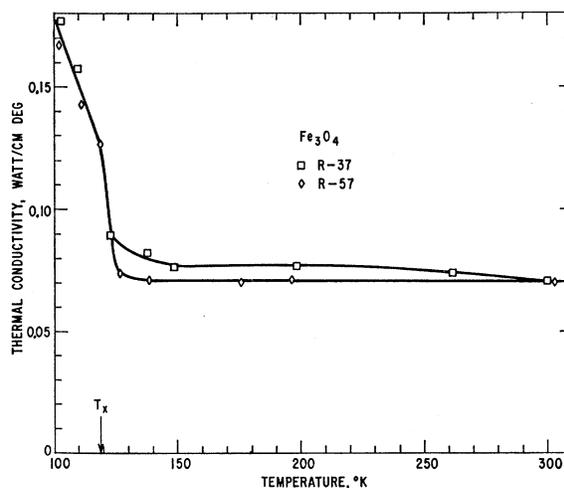


FIG. 7. The thermal conductivity vs temperature for synthetic Fe_3O_4 crystals. There is a sharp rise in conductivity at $121 \pm 2^\circ\text{K}$, the temperature at which octahedral ordering of Fe^{2+} and Fe^{3+} occurs.

⁹² R. Bauminger, S. G. Cohen, A. Marinov, S. Ofer, and E. Segal, Phys. Rev. **122**, 1447 (1961).

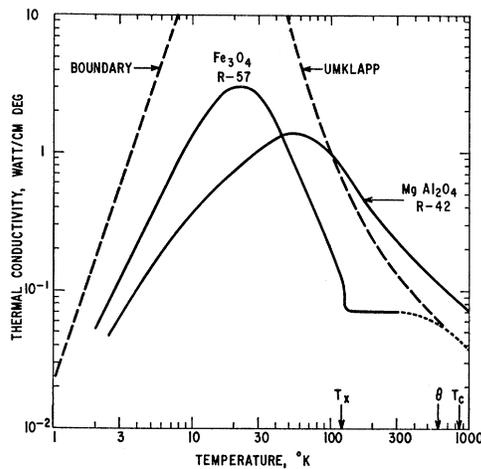


FIG. 8. A comparison of the thermal conductivity vs temperature for synthetic Fe₃O₄ and natural MgAl₂O₄ crystals. The dashed curves give the theoretical limits for Fe₃O₄ imposed by boundary and umklapp scattering. The temperatures T_x , θ , T_c are the transition, Debye, and Curie temperatures, respectively, of Fe₃O₄.

neighboring Fe³⁺. An incident compressible wave, a longitudinal phonon, can increase the overlap integrals, and, thus, may cause the octahedral ions to order. In this process of stress-induced ordering phonons will be absorbed, then these phonons will probably be reradiated incoherently, i.e., scattered. Such stress induced ordering has been observed before⁹⁸ for phonons of acoustical wavelengths in Fe₃O₄. This type of phonon scattering will occur as long as the octahedral ions are in the disordered state, i.e., $T > 119^\circ\text{K}$. From the thermal conductivity results it appears that this scattering probability varies only slowly with temperature up to temperatures at least comparable with θ . For $T > \theta$ this scattering from the magnetic disorder may still be present, but it will be masked by umklapp scattering. The predominance of the umklapp scattering also means that any anomalous behavior in K on passing through the T_c at 848°K might also be effectively masked, even though further disorder in the magnetic arrangement is present above T_c .

An average value of the phonon mean free path l at T_x can be calculated from the simple Debye formula

$$K = \frac{1}{3} l \bar{v}_0 C_v,$$

where C_v is the lattice specific heat per unit volume. C_v for the lattice can be estimated from the Debye function of $\theta = 600^\circ\text{K}$, and thus eliminate the magnetic specific heat contribution²⁴ near T_x . This computation yields an l_{qm}^0 value¹⁴ of 60 \AA for Fe₃O₄ at $T_x = 119^\circ\text{K}$. This is to be compared with very similar values of l_{qm}^0 found^{10,11} in the antiferromagnetic oxides MnO and CoO, and with the much larger values of 4000 and

2400 \AA found¹⁴ for the antiferromagnetic MnF₂ and CoF₂.

As T decreases below 119°K the octahedral ions become progressively better ordered, but there does not appear to be a convenient measure of this order, such as the sublattice magnetization that was used to interpret the data on CoF₂.¹⁴ If an average mean free path l is computed from $l = 3K\bar{v}_0^{-1}C_v^{-1}$, then l versus T from 60° to 115°K can be fitted by

$$l = [4 \times 10^{-8} \text{ cm}] \exp(U/kT),$$

where $U = 0.034 \text{ eV}$. This value of U is about $\frac{1}{3}$ of that found by Calhoun²⁵ from electrical resistivity measurements in this same temperature region. The exact significance of such an analysis of l is rather uncertain at present, and should be used with caution.

Impure Fe₃O₄

The measurements on R-63 in which 2.4% of the Fe²⁺ is replaced by Co²⁺ ($C_1 = 9.9 \times 10^{20} \text{ cm}^{-3}$ of Co) also show a break in the K curve near the T_x value for pure Fe₃O₄. The experimental value of T_x from Fig. 6 for this doped crystal is $110 \pm 5^\circ\text{K}$. The Co²⁺ enters the spinel lattice on the octahedral sites^{18,94} in place of Fe²⁺. Thus the octahedral ordering, which is responsible for the abrupt rise in K for crystals R-37 and R-57, has almost vanished in R-63. The randomly distributed Co²⁺, which has a magnetic moment different from that of Fe²⁺, cannot order at T_x because the temperature is too low for ionic diffusion to occur. The results on R-63 show that small concentrations of chemical impurities can have large effects on the K of Fe₃O₄ at low temperatures by preventing ordering. The effect of small additions of Co may rapidly saturate because the K vs T curve for R-63 is very nearly the same as the one measured by Douthett and Friedberg⁷ for a crystal approximating $(\text{Zn}_{0.5}^{2+})(\text{Fe}_{0.5}^{3+})[(\text{Co}_{0.5}^{2+})(\text{Fe}_{1.5}^{3+})]\text{O}_4$. Here, again, the Co²⁺ is on the octahedral sites. The effect of such impurities at 300°K and above is small because the intrinsic disorder is already large.

The presence of other point imperfections, such as vacancies, can also affect the K of Fe₃O₄. This is clear from the comparison of the results for R-37 and R-57 in Fig. 6 for $2^\circ\text{K} < T < 50^\circ\text{K}$. Fe₃O₄ crystal R-57 has a much lower vacancy concentration than R-37 and its thermal conductivity in this region is 4 times that of R-37. In R-37, the vacancy concentration C_3 is $2.6 \times 10^{20} \text{ cm}^{-3}$, which is much larger than the known concentration of chemical impurities. The transition metals give $C_1 = 1 \times 10^{18} \text{ cm}^{-3}$, the other metals give $C_2 = 3 \times 10^{18} \text{ cm}^{-3}$. In crystal R-57 the calculated vacancy concentration is $2 \times 10^{17} \text{ cm}^{-3}$, which is much less than the concentration of chemical impurities. Therefore the vacancies are probably the dominant phonon scatterers in R-37 at 10°K , while for R-57 the chemical impurities

⁹⁸ W. P. Mason, *Physical Acoustics and the Properties of Solids*, (D. Van Nostrand Company, Inc., Princeton, New Jersey, 1958), p. 314.

⁹⁴ E. Prince, *Phys. Rev.* **102**, 674 (1956).

can introduce local disorder into the magnetic lattice, and thus cause phonon scattering.

A concentration of $C_3 = 2.6 \times 10^{20} \text{ cm}^{-3}$ on the octahedral sites in R-37 will produce a mass fluctuation parameter of $\Gamma = 6.6 \times 10^{-4}$. Using the same arguments regarding vacancies at high temperatures ($T \geq 30^\circ\text{K}$) that were used for the $\text{MgO} \cdot 3.5\text{Al}_2\text{O}_3$, the value of K calculated from this Γ using Eq. (4) is 2.5 w/cm deg at 50°K . The observed value is only $\frac{1}{4}$ of this. Similarly the ΔW computed from Eq. (6) is quite small. Therefore the fact that the vacancies in Fe_3O_4 crystal R-37 have little effect on K for $T > 50^\circ\text{K}$ is verified by the same theory that holds for diamagnetic crystals. The small difference (10%) between crystals R-37 and R-57 evident in Fig. 7 is probably not caused by vacancies but by experimental errors.

Other Ferrites

Let us consider the K measurements on the other single crystal ferrites. The K vs T curves for some poorly defined ferrites measured by Douthett and Friedberg⁷ are shown in Fig. 6. Their K vs T curve for $\text{Co}(\text{Zn})\text{Fe}_2\text{O}_4$ is nearly the same as that for the MnFe_2O_4 curve, and is not shown. Their measurements on these Linde²² ferrite samples extend from 1.5° to 25°K , but the K vs T curves could reasonably be extrapolated to the point in Fig. 6 marked $X\text{Fe}_2\text{O}_4$. Smit and Wijn³ state that the K value given for $X\text{Fe}_2\text{O}_4$ (where X is some metallic ion) is reasonably representative of all spinel ferrites at 300°K . From the results on R-63 it appears reasonable to conclude that the ferrite curves of Douthett and Friedberg would not exhibit any pronounced rise in K , such as found in Fe_3O_4 . In fact, K may decrease monotonically as T decreases below 300°K . This is quite similar to the behavior for $\text{Mg}_x\text{Fe}_y\text{Al}_z\text{O}_4$, also shown again in Fig. 6. Some measurements of K for nickel ferrite, $\text{Ni}(\text{Fe})\text{Fe}_2\text{O}_4$, made by White and Woods⁹⁵ gave results similar to those in Fig. 6 for $\text{Mn}(\text{Zn})\text{Fe}_2\text{O}_4$.

It is known that all of these ferrites, MnFe_2O_4 , $\text{Mn}(\text{Zn})\text{Fe}_2\text{O}_4$, $\text{Co}(\text{Zn})\text{Fe}_2\text{O}_4$, and $\text{Ni}(\text{Fe})\text{Fe}_2\text{O}_4$, are either partially random spinels or inverse spinels. None are normal spinels. For instance MnFe_2O_4 has the distribution⁹⁶ $(\text{Mn}_{0.8^{2+}})(\text{Fe}_{0.2^{3+}})[(\text{Mn}_{0.2^{3+}})(\text{Fe}_{0.1^{2+}})(\text{Fe}_{1.7^{3+}})]\text{O}_4$. Here the unbracketed group is distributed randomly over the tetrahedral sites, and the [] group randomly over the octahedral sites. This is similar to the $\text{Co}_{0.07}\text{Fe}_{2.93}\text{O}_4$ in that the random distribution is frozen in during cooling, and the magnetic lattices exhibit many local fluctuations. It appears that phonons are quite effectively scattered by such a disordered magnetic lattice at low temperatures.

Douthett and Friedberg⁷ observed an increase in

the K of the MnFe_2O_4 and $\text{Co}(\text{Zn})\text{Fe}_2\text{O}_4$ in an applied magnetic field of 9400 oe. A preliminary search for a change in K with applied magnetic fields up to 5700 oe was made at 110°K in an Fe_3O_4 crystal similar to R-37. No noticeable effect, i.e. a change in K of $> 5\%$, was found. No further checks either at 4°K or at 300°K have yet been made.

The general picture of phonon scattering by disordered magnetic lattices, but not by ordered ones, indicates that thermal conductivity measurements on several different crystals might exhibit unusual behavior. For instance, ZnFe_2O_4 is a normal spinel⁹⁷ and becomes antiferromagnetic⁹⁸ below 9.5°K . Above 9.5°K it is paramagnetic. Thus an abrupt rise in the thermal conductivity of ZnFe_2O_4 might be seen at 9.5°K . A similar anomaly at low temperatures might appear in CdFe_2O_4 , which is a normal spinel⁹⁹ and is quite similar to ZnFe_2O_4 .

Point Impurity Effects for $T < 30^\circ\text{K}$

For none of the crystals studied in the present experiment was the K in the temperature range $T < 30^\circ\text{K}$ as high as the limit imposed by boundary scattering alone. Crystal R-38 of MgO came the nearest to this limit, but even its K was a factor of 2 low. The excess thermal resistivity is attributed to chemical impurities. Three different kinds of impurities have been mentioned: transition metals with a concentration C_1 , non-transition metals with a concentration C_2 , and vacancies with a concentration C_3 . In order to establish a general type of behavior the thermal resistivity at 10°K , $W(10^\circ\text{K})$, has been plotted as a function of concentration for all 10 crystals in Fig. 9. Where available, all three concentration values C_1 , C_2 , and C_3 have been entered. If the MgO and MgAl_2O_4 crystals, 7 in number, are considered, then the best correlation of $W(10^\circ\text{K})$ versus concentration is obtained by considering only C_1 and drawing the broad diagonal line shown in Fig. 9. This line indicates a relationship of the form $W(10^\circ\text{K}) \sim C_1^n$, when $n = +0.8$. An exact correlation is not expected since the C_1 values are only accurate to within a factor of 2, and since it is not really expected that all transition metal impurities will behave in the same way. The correlation of $W(10^\circ\text{K})$ with C_2 (or C_3) is considerably worse than that with C_1 . The $W(10^\circ\text{K})$ versus C_1 values for Fe^{2+} doped $\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}$ of Rosenberg and Sujak,⁶⁹ for Mn^{2+} doped NaCl of Klein,⁷¹ and for Cr^{3+} doped $\alpha\text{-Al}_2\text{O}_3$ of Berman, Foster, and Rosenberg¹⁰⁰ have also been entered in Fig. 9. The data for $\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}$ and NaCl fall on the line to

⁹⁷ J. M. Hastings and L. M. Corliss, *Revs. Modern Phys.* **25**, 114 (1953).

⁹⁸ J. M. Hastings and L. M. Corliss, *Phys. Rev.* **102**, 1460 (1956); E. F. Westrum and D. M. Grimes, *J. Phys. Chem. Solids* **3**, 44 (1957).

⁹⁹ E. J. W. Verwey and E. L. Heilmann, *J. Chem. Phys.* **15**, 174 (1947).

¹⁰⁰ R. Berman, E. L. Foster, and H. M. Rosenberg, *Defects in Crystalline Solids* (The Physical Society, London, 1955), p. 321.

⁹⁵ G. K. White and S. B. Woods, as reported in reference 7 footnote, p. 1665.

⁹⁶ F. W. Harrison, W. P. Osmond, and R. W. Teale, *Phys. Rev.* **106**, 865 (1957).

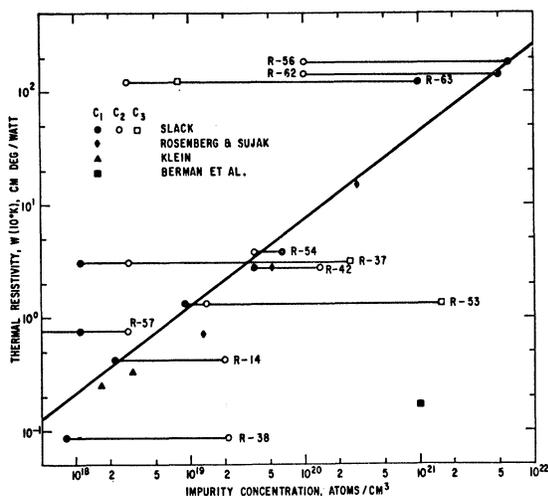
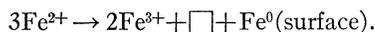


FIG. 9. A composite graph of the measured thermal resistivity at 10³K, $W(10^3\text{K})$, as a function of impurity concentration. The values C_1 , C_2 , C_3 are explained in the text. The diagonal line is the "best-fit" if $W(10^3\text{K})$ depends mainly on C_1 , the transition metal concentration.

within a factor of 2 in spite of the very different crystal structures and impurity environments. However, the data of Berman *et al.*¹⁰⁰ on $\alpha\text{-Al}_2\text{O}_3$ disagree by a factor of at least 100, thus indicating that the phonon scattering in this temperature region does depend on the particular nature of the transition metal impurity. Orbach's⁸⁷ analysis of Fe^{2+} in $\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}$ describes one such particular scattering process. In this process it appears from Fig. 9 that n in C_1^n is nearly $n=1.0$. In the theory of phonon-spin interaction given by Orbach,⁸⁷ W cannot be simply proportional to C^n . A more complex dependence involving temperature, scattering cross-section, and concentration is expected. Figure 9 simply indicates that W increases monotonically with C , and that some transition metals have roughly similar effects in various host crystals.

The results on Fe_3O_4 (crystals R-37, R-57, R-63) tend to fit the same general pattern as MgO and MgAl_2O_4 . The main difference is that both diamagnetic and paramagnetic impurities can produce disorder in the magnetic lattice of Fe_3O_4 by substituting for an Fe atom. Whether diamagnetic impurities such as Al and Na have as large an effect as the paramagnetic Co cannot be decided from Fig. 9. However it is clear from Fig. 9 that Fe vacancies (see R-37) have a scattering cross-section at 10³K only 2 to 10% of that of the chemical impurities. A vacancy \square is produced in the cation lattice according to



The change in the spin only magnetic moment in the interior of the crystal is

$$\Delta S = 2(5/2) - 3(4/2) = 1,$$

or an average $\langle \Delta S \rangle_{\text{av}}$ per lattice site of $\frac{1}{3}$. When an

Fe^{3+} ion is replaced by Al^{3+} , then $\langle \Delta S \rangle_{\text{av}} = \frac{5}{2}$. It seems plausible to assume that the scattering cross section of a defect increases with $\langle \Delta S \rangle_{\text{av}}$. Hence an Al atom should have a greater effect than a vacancy. This argument indicates why vacancies may be much less effective scatterers than substitutional diamagnetic impurities in Fe_3O_4 .

In order to reach the boundary scattering limit in Fe_3O_4 at 10³K for $d=0.24$ cm the vacancy concentration should be less than $\sim 10^{18}$ cm⁻³ and the chemical impurity content less than $\sim 10^{17}$ cm⁻³. Rather special efforts would be required in order to obtain such pure Fe_3O_4 .

CONCLUSIONS

The thermal conductivity of single crystals of MgO, MgAl_2O_4 , $\text{MgO} \cdot 3.5\text{Al}_2\text{O}_3$, $\text{Mg}_x\text{Fe}_y\text{Al}_z\text{O}_4$, and Fe_3O_4 has been measured from 3° to 300°K.

The present results for MgO and previous ones for $\alpha\text{-Al}_2\text{O}_3$ are explicable in terms of the theory of phonon scattering by umklapp processes, by isotopes, and by crystal boundaries. The thermal conductivity of pure crystals of the double oxide MgAl_2O_4 can be considered to be approximately an average between the results for MgO and $\alpha\text{-Al}_2\text{O}_3$. The thermal conductivity of the natural crystals studied is, however, considerably reduced at low temperatures by the presence of chemical impurities, particularly transition metal impurities.

The study of $\text{MgO} \cdot 3.5\text{Al}_2\text{O}_3$ and $\text{Mg}_x\text{Fe}_y\text{Al}_z\text{O}_4$ indicates that the thermal conductivity of MgAl_2O_4 can be somewhat reduced by the presence of vacancies, and greatly reduced by the presence of large quantities of Fe.

The measurements on Fe_3O_4 indicate that phonons are the dominant carriers of thermal energy, and that any magnon or electron heat transport is very small. In addition to the scattering mechanisms found in the diamagnetic crystals, the phonons in Fe_3O_4 and other ferrites can be scattered by local disorder in the magnetic lattice. Vacancies as well as chemical impurities can produce such local disorder, but vacancies are much less effective than the impurities.

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