The agreement between experiment and the calculation of Koppe, as shown in Figs. 1 and 2, is on the whole striking, although the departure from the curve in Fig. 1 of the low temperature points for niobium and vanadium is beyond the limit of experimental error. Nevertheless, the general fit of Koppe's curve suggests a real regularity displayed by all superconductors, a regularity which may be regarded as an approximate "law of corresponding states."

\* Assisted by the ONR and Linde Air Products Company.
\* City College of New York, New York, New York.
‡ Barnard College, Columbia University, New York, New York.
<sup>1</sup> W. Heisenberg, Z. Naturforsch. IIa, 185 (1937).
<sup>2</sup> H. Koppe, Ann. phys. 1, 405 (1947).
<sup>3</sup> W. H. Keesom and P. H. van Laer, Physica 5, 193 (1938).
<sup>4</sup> W. H. Keesom and M. Désirant, Physica 8, 273 (1941).
<sup>5</sup> P. L. Bender and C. J. Gorter, Physica 18, 1 (1952). We are indebted to Professor Gorter for a copy of their paper in advance of publication.
<sup>6</sup> Brown, Zemansky, and Boorse, Phys. Rev. 86, 134 (1952).
<sup>7</sup> J. R. Clement and E. H. Quinnell, Phys. Rev. 79, 1028 (1950).
<sup>8</sup> Parkinson, Simon, and Spedding, Proc. Roy. Soc. (London) A207, 137 (1951).

<sup>6</sup> Parkinson, Simon, and Spectrug, 1 to a constraint of the second state of the low temperature Conference of the ONR General Electric Research Laboratories, Schenectady, New York, October 1952. <sup>10</sup> J. R. Clement and E. H. Quinnell, Phys. Rev. **85**, 502 (1952). ( $\gamma$  was calculated from magnetic data and taken to be 8.92 ×10<sup>-4</sup> cal/mole-deg<sup>2</sup>.) <sup>11</sup> W. H. Keesom and J. A. Kok, Physica 1, 175, 595 (1934).

## Evidence for Domain Structure in Antiferromagnetic CoO from **Elasticity Measurements**

M. E. FINE

Bell Telephone Laboratories, Murray Hill, New Jersey (Received July 15, 1952)

OUNG'S modulus in CoO on cooling undergoes a large decrease near the Néel point.<sup>1</sup> Street and Lewis<sup>1</sup> suggest that this is due to the influence of an applied stress on the preferred orientation of the spin vectors in antiferromagnetic domains. A more detailed study and explanation of the elasticity of CoO is herein reported.

A rod of CoO was prepared by pressing powdered CoO and sintering at 1130°C in  $\hat{N}_2$ . X-rays revealed only the CoO structure; the apparent density after sintering was 86 percent of the density of CoO.



Young's modulus (E) and the internal friction (1/Q) were measured from 80 to  $-196^{\circ}$ C in the pressed and sintered rod, employing both electromagnetic and piezoelectric excitation and detection of longitudinal vibrations (35 to 60 kc sec<sup>-1</sup>). On cooling from 20 to  $-20^{\circ}$ C, as shown in Fig. 1, *E* drops from 17 to  $6.3 \times 10^{11}$ dynes cm<sup>-2</sup>, the region of maximum change being 0 to  $-8^{\circ}$ C. (The Néel temperature, determined from paramagnetic susceptibilities, is  $-2^{\circ}C^{2}$ ) The effect persists to low temperature, for at

-196°C E is  $8.5 \times 10^{11}$  dynes cm<sup>-2</sup>. The internal friction (1/Q) for a constant maximum strain amplitude of  $10^{-7}$ , also shown in Fig. 1, rises on cooling from  $2 \times 10^{-4}$  (the background) to a peak value of  $91 \times 10^{-4}$  at  $-20^{\circ}$ C, then decreases to a minimum at -150°C and rises again to 8.5×10<sup>-4</sup> at 190°C. At room temperature and  $-20^{\circ}$ C, increasing the maximum strain amplitude from  $10^{-8}$  to  $10^{-5}$  has little effect on E or 1/Q, but at  $-190^{\circ}$ C E decreases 1 percent and 1/Q increases from 5 to  $25 \times 10^{-4}$ .

The general course of our modulus-temperature curve and that reported by Street and Lewis<sup>1</sup> are in substantial agreement; however, our modulus values are larger by approximately a factor of 5, and the Néel temperature appears to be  $-2^{\circ}$ C in our sample rather than 16°C. The discrepancy among modulus values and Néel points may be due to composition and porosity variations.

In considering possible explanations the following properties of CoO seem important: The formation of antiferromagnetism causes an increase in volume; the coefficient of linear expansion is high in the temperature range 5 to  $-75^{\circ}$ C, with a peak of  $22 \times 10^{-6}$ deg C<sup>-1</sup> at  $-2^{\circ}$ C. Antiferromagnetism distorts CoO from cubic (NaCl structure) to tetragonal;  $^{3,4} c/a$  becomes increasingly less than one as the temperature is decreased, being 0.9884 at  $-180^{\circ}$ C.<sup>3</sup> The antiferromagnetically aligned atomic magnetic moments are parallel to  $\langle 100 \rangle$ .<sup>5</sup> Presumably it is this direction that becomes shortened.

The large decrease in E implies that an applied stress produces strains in addition to, and in some cases even larger than, the normal elastic strain. In antiferromagnets an effect analogous to the stress-induced change in domain structure in ferromagnets is suggested.6 Neighboring domains within a crystal are postulated, differing as to which of the three possible orientations of the c axis is present. Then, for example, tension would increase the area of those domains whose c axes are nearly perpendicular to the stress axis, thereby increasing the sample length. The amount of strain from a given amount of domain boundary movement would increase on cooling since c/a decreases, but domain boundary movement becomes more difficult owing to the lattice strain produced. Thus, E from this effect would not continuously decrease on cooling. Stress-induced antiferromagnetic domain boundary movement, like domain movement in ferromagnets, is expected to be hysteretic; consequently, the proposed mechanism seems especially reasonable for low temperatures where 1/E and 1/Q increase with strain amplitude and, furthermore, well-defined domains are expected.

Near the Néel temperature, where the magnetic moments are loosely coupled, another mechanism which may yield a strain and internal friction is a stress-induced change in the degree of antiferromagnetic order. The strain would come from the resulting change in volume and tetragonality. E and 1/Q are more conceivably independent of strain amplitude by this mechanism.

This investigation has benefited from the assistance of Miss N. T. Kenney and helpful suggestions from and discussion with Mr. P. W. Anderson, Mr. W. C. Ellis, Mr. J. K. Galt, Mr. H J. McSkimin, Mr. F. J. Schnettler, and Mr. E. E. Schumacher.

<sup>1</sup> R. Street and B. Lewis, Nature 168, 1036 (1951).
<sup>2</sup> H. Bizette, Ann. phys. 1, 295 (1946).
<sup>3</sup> N. C. Tombs and H. P. Rooksby, Nature 165, 442 (1950).
<sup>4</sup> S. Greenwald and J. S. Smart, Nature 166, 523 (1950).
<sup>8</sup> Shull, Strauser, and Wollan, Phys. Rev. 83, 333 (1951).
<sup>6</sup> R. M. Bozorth, Ferromagnetism (D. Van Nostrand, Inc., New York, 1951), pp. 472, 595.

## Errata

The Symmetrical Pseudoscalar Meson Theory of Nuclear Forces, MAURICE M. LÉVY [Phys. Rev. 86, 806 (1952)]. In the last portion of Eq. (3), replace  $[(2/\pi)K_0(\mu r)]^2$  with  $(\mu/2M)[(2/\pi)K_1(\mu r)]^2.$ 

Angular Distribution of Shower Particles as a Function of Depth, L. EYGES AND S. FERNBACH [Phys. Rev. 82, 287 (1951)]. Equation (1) should read:  $\pi(E_0, E, \theta, t)2\pi\theta d\theta dE = \pi_{1ong}(E_0, E, t)dE \cdot P(x, s)\theta d\theta \cdot (E/E_s)^2$ . The present addresses