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hanical vibration to al feedback by couples

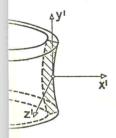
slightly modulated ne reflectance, was h quartz window (E) nt of the multiplier h setting of the more as achieved by using the photomultiplianode current of o $\Delta R/R$, the relative component was ne th by means of a phase ed on an x-y recorde checked with a phot found to be correct :

ement

the sample will be the formed by joining the r with rectangular cro s is true if the influence The cross section of the rectangular. There are tensor o and the strain the surface of the ring radius p1 and the thick d the width b is small ad the strain tensor take

$$\begin{bmatrix} e_{xx'} & & & \\ & e_{yy'} & & \\ & & e_{zz'} \end{bmatrix} . \quad (1)$$

t in our measurements the validity of Eqs. (1 measuring ρ_1 and ρ_2 0 thicknesses and widths of the surface is give $2\rho_2$. On the other hand ulated using the stress the stress tensor. Using



ing the ends of a previous ection. The stress axis is z'.

stress tensor given by (1), we found the agreement streen the measured and calculated ratio to be better $|a_{zz}| = 3\%$, provided the condition $|e_{zz}| \le 4d^2/b^2$ was filled. In the optical measurements on Cu crystals, rical numbers were $|e_{zz'}| = 4 \times 10^{-4}$, $4d^2/b^2 = 0.4$. Thus the above condition was always met.

in the actual measurements, the crystal forms a all segment of the ring shown in Fig. 2; the distorproduced by the clamps cannot always be neacted. Measurements of ρ_1 and ρ_2 of large aluminum clamped at the ends in a way similar to that bown in Fig. 1 were carried out. The difference beacen the calculated and the measured $|e_{zz}'/e_{yy}'|$ at me midpoint of the sample was again below 3%, proaded the free length of the sample (Fig. 1) was at east twice its width. Typical values for the Cu crystals sed in the optical experiments are l=10 mm, b=3 mm, e., this condition was also fulfilled.

Figure 3 shows the arrangement to determine the omponent ezz' of the strain tensor by measurement of the focal length of the cylindrical mirror, formed by the bent sample. The sample was oscillating, and the requency of the stroboscope was tuned close to the csonance frequency of the sample. The distance beween the sample and the image of the slit changed periodically with the difference frequency ω (strobo- ω (sample). The amplitude of the strain at the surface is given by

$$e_{zz}' = \Delta a d(2a_0)^{-2} \{1 + [1 + (\Delta a/a_0)^2]^{1/2}\}^{-1},$$
 (2)

where a_0 is the position of the image for zero strain and La is the difference in the position for maximum extension and compression. The accuracy of this method increases with decreasing distance between lens and sample. It was about $\pm 5\%$ for the geometry used here. The components e_{zz}' , e_{yy}' of the strain tensor are expressed in terms of the measured component e_{zz}' by means of the stress-strain relation using the form 1) of the stress tensor. The elastic constants are taken from Ref. 20. Because of the sample dimensions chosen, the errors in e_{xx}' and e_{yy}' due to deviations from (1) are smaller than 3%. During the optical mea-

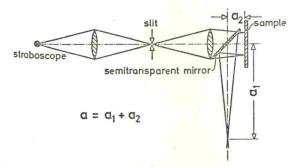


Fig. 3. The optical design which was used to determine the focal length of the cylindrical mirror formed by the bent sample.

¹⁰ American Institute of Physics Handbook (McGraw-Hill Book Co., New York, 1957), Chap. 2, p. 56.

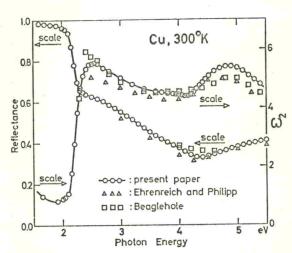


Fig. 4. The reflectance and \(\epsi_2\), the imaginary part of the dielectric constant of Cu at room temperature. The values determined by Ehrenreich and Phillipp (see Ref. 18) and by Beaglehole (see Ref. 22) are only shown if they differ by more than 2% from the values given in the present paper.

surement, the strain amplitude and phase was monitored by a pickup capacitor, consisting of one of the magnets at the ends of the lever arms (Fig. 1), moving against a fixed, insulated piece of sheet metal.

Sample Preparation

The orientation of the samples cut from a single crystal was determined to within ±1° using Laue diagrams. The surface preparation consisted of grinding, mechanically polishing, and electropolishing21 the sample. The electropolishing was terminated by quickly rinsing in deionized water and alcohol. After taking the sample from the alcohol bath, the thin film of alcohol at the surface was immediately removed by a warm stream of air. The reflectance of a freshly prepared sample, measured within 10 min after the electropolishing, is given in Fig. 4. The growth of an oxide layer at the surface of the sample is responsible for the observed decrease in the reflectance with time. This decrease is most pronounced in the ultraviolet. We observed a 1% decrease at 5.5 eV within 1 h after the electropolishing.

Although the reflectance of our samples was measured in air, it deviates less than 1% from the values determined by Beaglehole,22 which were measured in a high vacuum after reducing the oxide layer at the surface. The only exception is the region around 4.3 eV. The resolution of the vacuum monochromator used by Beaglehole was not high enough to resolve finer details of the minimum at that energy23 (see Fig. 4). Thus the oxide layer on our sample modifies the reflectance not more than 1% between 1.5 and 5.5 eV. The reflectance given by Ehrenreich and Philipp¹⁸ is slightly lower

23 D. Beaglehole (private communication).

²¹ W. J. Tegart, The Electrolytic and Chemical Polishing of Metals (Pergamon Press, Inc., New York, 1959), 2nd ed.
²² D. Beaglehole, Proc. Phys. Soc. (London) 85, 1007 (1965).