SECOND SEMI-ANNUAL TECHNICAL REPORT

ELECTRICAL and OPTICAL PROPERTIES of AMORPHOUS MATERIALS*

by

A Solid-State Group of the Physics Department Northern Illinois University DeKalb, Illinois 60115

> *Sponsored by Advanced Research Projects Agency ARPA Order No. <u>1562</u>

The views and conclusions contained in this document are those of the authors and should not be interpreted as necessarily representing the official policies, either expressed or implied, of the Advanced Research Projects Agency or the U. S. Government.



GENERAL INFORMATION

ARPA Order Number	1562
Program Code Number	0010
Name of Contractor	U. S. Army Research Office
Effective Date of Contract	June 1, 1970
Contract Expiration Date	May 31, 1971
Amount of Contract	\$45,868.00
Contract Number	DA-ARO-D-31-124-70-G77
Principal Investigator Phone Number	Dr. Charles Wood (815) 753-1773

No.

CONTENTS

THEORET	ICAL RESEARCH		
Ι.	The Kramers-Kronig (Hilbert) Transform	1	
II.	Electronic Structure	2	
MATERIA	MATERIALS RESEARCH		
Ι.	Amorphous Thin Films	4	
II.	Crystalline Thin Films	4	
ш.	Single Crystals	5	
IV.	Differential Thermal Analysis	5	
STRUCTURAL INVESTIGATIONS			
Ι.	Mossbauer Studies	6	
п.	Electron Microscopy	7	
OPTICAL	PROPERTIES		
Ι.	Transmission and Reflectivity Measurements	9	
п.	Modulated Reflectivity	10	
III.	Photoconductivity	11	
IV.	Photoemission	12	
TRANSPORT PROPERTIES			
Ι.	Resistivity vs. Temperature	14	
REFERENCES 15			
FISCAL STATUS 17			
APPENDICES 18		18	

Summary

A continuing effort is being made to maintain a closely knit experimental and theoretical program. Theoretical work on the use of Hermite functions to compute Hilbert (Kramer Kronig) transforms is completed and has been successfully tested on several systems and applied to the analysis of optical data on single crystalline and amorphous thin films of Sb₂Se₃.

A satisfactory method of preparing thin amorphous films of controlled composition by coevaporation has been developed and improved upon in the past six months. Differential thermal analysis has yielded important information on the compositional dependence of the glass to crystalline transition in Sb_xSe_y amorphous films.

> Mossbauer studies have yielded important information on the short range order of amorphous films and on the structural chemistry of mixed chalcogenide systems. This technique, where applicable, appears to fulfill a need for analysis on a microscale where more conventional analytical tools such as X-ray and electron microscopy are inadequate. However, initial studies of Sb_XSe_y amorphous films by electron microscopy are yielding very interesting data but a full interpretation of the data awaits compositional analysis of the films.

In the optical studies of thin amorphous films, one of the more significant findings has been the close correspondence between the energy band gaps in the crystalline and amorphous form of Sb_2Se_3 and the apparent lack of band tailing. Further optical measurements on single crystal Sb_2Se_3 in the neighborhood of the band edge as a function of crystal orientation

have supported the initial findings of a direct transition. We are currently pressing ahead with resistivity versus temperature measurements on both the amorphous and single crystal forms to obtain thermal activation energies for comparison with the optical values.

4

Several different approaches to modulated reflectivity (and modulated absorption) are under development. This data should prove invaluable in the analysis of the electronic structure of the antimony chalcogenides. Along the same lines, the setting-up of appartus for the very powerful technique of photoemission is at a final stage of completion and is already yielding results on the density of states.

As evidenced by the large number of references to Bull. Am. Phys. Soc. much of the work is still in the initial stages. In keeping with past practice we have included preprints of work currently being submitted for publication to elaborate, for the interested reader, on details not suitable for this general progress report. A summary of the more significant findings of this program to date will be presented at the forthcoming International Symposium on Amorphous and Liquid Semiconductors and an abstract is appended to this report. Reprints of papers, published since the last six-monthly report, are also appended.

THEORETICAL RESEARCH

I. The Kramers-Kronig (Hilbert) Transform

During the last six months the work initiated earlier on the use of Hermite functions to compute Hilbert (Kramers-Kronig) Transforms has been completed. A paper outlining the general mathematical technique has been submitted for publication and a preprint was included in the previous technical report. This paper concentrates upon the application of the method to functions of compact support such as densities of states. Hilbert Transforms of such functions are useful, e.g., in the computation of optical functions from the results of electronic structure calculations. Briefly, a convergent, matrix representation of the Hilbert Transform kernel has been employed to compute Hilbert Transforms of two densities of states, 1) the classic simple cubic s-band (triple cosine) density of states and 2) the electronic density of states of fcc palladium¹. The results indicate that critical point structure does not shift in frequency under a proper Hilbert Transform. Details of the computat tional method and its analytical development are found in the paper.

The application of the above technique to optical data analysis, i.e., to the extraction of optical functions from experimental mear-normal incidence reflectivity measurements has also come to fruition. A preprint of a paper which is now in its final stages of preparation is attached to this report as Appendix A. A preliminary report of the results of the technique as applied to Sb_2Se_3 crystalline samples was presented at the March Meeting of the American Physical Society in Cleveland ²(Appendix B). Further application of the technique has been made to thin amorphous films

of the composition Sb_2Se_3 and this work was reported at the April Meeting of the American Physical Society in Washington³(Appendix C). There are indications of substantial differences in the structural details of the optical functions of crystalline and amorphous Sb_2Se_3 . Further investigations in this area are underway. Basically the method employs the reflectivity of systems of simple charged oscillators to extrapolate the experimental data outside the measured data range and to convert the function whose Hilbert Transform needs to be computed to one of compact support. The methods employed in the previous work described above are then applicable. As a check upon the power of the technique it has been applied to reflectivity data⁴ on BaF_2 in the far ultraviolet. This data has a wealth of sharp structure arising, most likely, from deep excitons and provides a test of the resolution of the computational technique.

II. Electronic Structure

Some progress has been made on the computation of the electronic spectra of amorphous materials. The method we have explored is similar in spirit to the work of Maschke and Thomas⁵ for electronic structure in amorphous solids with short-range order. We are planning to apply our method to specific amorphous solids. A computer program to compute electronic energies of a model of amorphous Ge and the associated density of states, is under development. The computational problems are severe both as to complexity and as to computational time required. The problem is, however, tractable and with continuing work will be completed.

In another vein work has been initiated to attempt to make use of the results recently obtained by Romeris⁶ to connect the electronic structure of amorphous solids to that of crystals of similar composition.

2

ß

Romeris has shown that critical point structure behaves in a particular manner as a "crystal" potential varies from being truly periodic to being <u>almost periodic</u>. Since there is evidence³ of critical point structure in the optical functions found for amorphous films of Sb_2Se_3 it may be feasible to correlate this structure in a definite manner with that found in crystalline Sb_2Se_3 . We have, therefore begun to explore methods for tracking the behavior of critical point structure with disorder.

MATERIALS RESEARCH

I. Amorphous Thin Films

A large number of amorphous films of various compositions in the Sb-Se system have now been prepared by coevaporation. The compositions were determined by micro-probe analysis in conjunction with the Geophysical Laboratory of the University of Chicago. A fair degree of reproducibility has been obtained but, because of the critical dependence of optical properties on composition⁷ (Appendix B) the deposition controllers (Sloan Omni IIA) have had to be upgraded. The most significant improvements are: newly-developed circuit boards have been installed, gold-plated quartz crystals of circular geometry are now employed and, most importantly, the frequency and deposition rate of each source is continuously monitored externally by chart recorders throughout deposition. Despite these improvements control of stochiometry to within 1 atomic percent cannot be guaranteed.

In an attempt to obtain even closer control of composition two further approaches will be tried: 1) sputtering, under equilibrium conditions: 2) coevaporation of a graded composition across an extended substrate. Both systems are under construction, the former being nearly completed.

II. Crystalline Thin Films

The three-temperature technique⁸ was successfully applied to the deposition of a stochiometric, polycrystalline film of Sb_2Se_3 . There were difficulties in achieving adequate deposition rates which are determined by both substrate temperature and melt temperature. Too fast

4

a deposition rate produced a film with small, irregular crystal grains. With too slow a rate there is virtually no deposition. It was found that a fused-quartz substrate temperature of 450° C and a melt temperature of 630° C produced a satisfactory film for optical studies.

III. Single Crystals

A new single crystal growth apparatus is under development using a tin-oxide coated quartz tube for both the background heating elements and the zone heating element. These tubes have the advantage that they will develop temperatures well above 600° C while remaining very transparent to allow good visibility of the molten zone and hence good control of the growth process.

IV. Differential Thermal Analysis

DTA measurements have been carried out on various compositions of Sb-Se films in the temperature range of 30° C to 300° C⁹ (Appendix C). The glass to crystalline phase transitions of the samples were investigated. This transition was found to be endothermic and located ar $\sim 190^{\circ}$ C for Sb₂Se₃. Those films that were on the Se-rich side of Sb₂Se₃ showed amorphous-crystalline transition for Se, which was located at $\sim 120^{\circ}$ C, as well as the Sb₂Se₃ transition. Plans have been made to incorporate a more sensitive arrangement of thermo-couples more suited to a thin film geometry.

STRUCTURAL INVESTIGATIONS

I. Mossbauer Studies

Work performed in collaboration with Dr. S. Ruby of Argonne National Laboratory produced the very significant finding that all Sb atoms in Sb_xSe_y (including Sb_2Se_3) amorphous solids sit in closely similar environments in contrast to the Sb_2Se_3 crystal where there are two distinct sites¹⁰ (Appendix B). This is of particular significance in that it lends support to the view that the evaporated films are in reality amorphous and not "micro-crystalline", since we now know that the short-rangeorder differs in the two forms. This work will be submitted for publication and a preprint is attached (Appendix D).

In order to investigate cross-linking effects of Group IV elements in amorphous Group V chalcogenides, Mossbauer studies using ¹¹⁹Sn in a BaSnO₃ source, have been carried out on $Sn_x(As_2Se_3)_{1-x}$ glasses with varying concentrations of tin, i.e. (x = 2.5, 5.0, 8.0, 10.0 At. %) and on the two crystalline samples of $SnSe_2$ at liquid nitrogen and liquid He temperatures¹¹ (Appendix B). $Sn_x(As_2Se_3)_{1-x}$ glasses samples were prepared by quenching the melts into a brine bath at 200K.

The limit of tin solubility in the quenched glassy state is tentatively set by Kolomiets et al¹² at 7 At. %. Thus 8.0 and 10.0 At. % Sn, glass samples may have a crystalline component. X-ray patterns for all the samples, however, were similar and characteristic of amorphous systems. The major phase present in a deliberately crystallized $As_2Se_3 + 10$ At. % Sn sample is SnSe and the corresponding amorphous sample also contains a

6

10

1 . www. allowid. or hall Ser

small amount of SnSe. A critical Mossbauer spectra for 10 At. % Sn amorphous sample at 4.2° K shows an increase of absorption by about 15% compared to that at liquid nitrogen temperature. The overall character of the spectra is similar to that obtained by Borisova et. al.¹³, the major difference occurred in the intensities of the spectra corresponding to SnSe₂ and SnSe. The average shift in all systems corresponding to SnSe₂ is 1.68 mm/sec which is slightly less than that of α -Sn. On the basis of Ruby's¹⁴ atomic calculations, the electronic configuration is 0.7s, 3.3p. The average shift corresponding to SnSe is 3.45 mm/sec and the electronic configuration turns out as 1.6s, 2.4p.

Thus, it is clear from the present Mossbauer investigations of the amorphous systems that the degree of local order is very high, i.e., molecules or microcrystals of small enough dimensions to escape x-ray detection are held in a disordered matrix. The microchemistry does appear to correspond to the suggested chemistry of Shkolnikov¹⁵. The Mossbauer spectra for Sn in the glasses corresponds very closely to those of crystalline SnSe and SnSe₂. However the range of chemical order is too small to be detected by x-ray diffraction.

II. <u>Electron Microscopy</u>

Electron microscopy and electron diffraction studies were made on thin (about 250° Å thick) films of Sb_xSe_y . Amorphous films appear granular with irregularly shaped particles of about 1 x 10^{-4} cm diameter. The corresponding diffraction pattern consists of three diffuse bands. One can observe crystal growth from these amorphous films as caused by the heat of the electron beam. Depending on the sample composition, different types of ordering are observed: 1) bizarre looking dendrite-like

7

needles of about 1000 \hat{A} width with a complicated fine structure, 2) fibres of about 60 \hat{A} width which are lined up parallel to one another or are spiralling around each other, 3) rounded domains of micron size with narrow subdomains of about 100 x 1000 \hat{A} which have different alignments for different domains.

The diffraction patterns of sections 50 μ in diameter of the crystallized films range from spotty powder patterns and fibre patterns to single crystal μ atterns. Typical is a one dimensional ordering resulting in irregularly spaced spots on equally spaced lines. The ordered Jimension varies somewhat around 3.24 Å. It is not known at this date if the corresponding reflections relate to the "forbidden" (100) and (010) reflections from Sb₂0₃ structure or belong to a completely new crystal type. Definitely not compatible with the Sb₂0₃ structure are recurring hexagonal patterns which cannot be attributed either to selemium or antimony.

A complete interpretation of the results will be attempted when all the sample compositions have been analyzed.

OPTICAL PROPERTIES

9

I. Transmission and Reflectivity Measurements

Several different methods have been employed to determine the optical constants n and k, from the reflectivity (R) and transmission (T) of a thin film in the spectral region near and below the absorption edge. The first technique tried was to search the n-k plane for points that would fit the observed R and T. This method had limited success because for a given R and T there are multiple solutions in the *n*-k plane and the computer program was taking excessive time. The other method tried was that of quadratic interpolation. This method is not yet fully completed, but it gives greater accuracy in less computer time.

Absorption constant (α) vs. photon energy graphs were determined by transmission and reflectivity data for a wide range of compositions of Sb-Se amorphous films⁷ (Appendix B). A close correspondence was noted between the position of the optical gap of the amorphous films and single crystal Sb₂Se₃. The α values were determined from 10^{6} cm⁻¹ to 10^{3} cm⁻¹ for selected samples using a wide variation of sample thicknesses. Thin films ($\sim 5_{\mu}$) of Sb₂Se₃ must still be made and measured inn order to determine the type of transition that occurs (direct of phonon assisted).

The reflectivity and transmission of single crystals of Sb_2Se_3 grown in a horizontal zone refiner were measured in the range 0.03 eV to 11 eV¹⁶ (Appendix B). R and T were measured as a function of polarization in the cleavage plane. The reflectivity structure was shown to

agree with Soblov et. al.¹⁷ for the two polarization directions in the cleavage plane in the energy range 1 eV to 4 eV; and for unpolarized light in the range 4-11 eV. The energy range 0.03 to 0.5 eV was covered on the Beckman IR-12 spectrophotometer; 0.5-4 eV on a Cary 14 R spectrophotometer; and 4-11 eV on a McPherson vacuum U.V. spectrophotometer at Argonne National Laboratory (kindly loaned by Dr. P. Yuster). The absorption coefficient was calculated from this data near the absorption edge at about 1 eV. Different powers of the absorption coefficient was responsible for the absorption edge. A direct allowed transition was found for both directions of polarization.

Initial low light level measurements have been made using an S.S.R. photon counter. Great lengths have been gone to in order to prevent light leaks in the system. The phototubes can be cooled down to solid CO₂ temperatures for operation in a low noise mode. This arrangement should prove to be quite valuable for the measurement of optical transmission well above the absorption edge and also for small signal modulation experiments.

II. Modulated Reflectivity

Thermally modulated reflectivity measurements were performed on single crystals of Sb_2Te_3 with the light incident on the cleavage plane $(E \perp C)^{18}$ (Appendix C). The data was taken in the visible and near infrared regions and was found to agree with critical points observed in the transmission spectrum of thin cleaved sample using a photon counter.

Electro-reflectance apparatus is now completed which will be used

10

to measure the electro-modulated reflectivity spectrum of Sb_2Se_3 single crystals, in all three orientations of the electric vector, and amorphous Sb_xSe_v films.

Also under construction is apparatus for wavelength modulated reflectivity which involves halving the optical beam from the exit slit of a monochrometer and comparing the reflectivity of the sample to light from the two halves by means of a lock-in amplifier.

Construction of apparatus for rotoreflectance is also underway. This technique utilizes a rotating polarizer to measure the changes in reflectivity as a function of electric vector orientation.

III. Photoconductivity

In the photoconductivity area most work was concentrated on study of amorphous Sb_xSe_y . The spectral dependence of photoresponse and photoconductive gain depend critically on sample composition. The photoconductivity activation energy (as determined by Moss'¹⁹ criterion) generally agrees with the optical activation energy as determined from transmission measurements⁷ (Appendix B) in case of stochiometric amorphous Sb₂Se₃ being equal to 1.1 eV, which is very close to that of single crystalline Sb₂Se₃. We feel that this point is of fundamental importance as it shows that the gap is approximately the same in amorphous and crystalline Sb₂Se₃; contrary to previously published work by Kolomiets et. al.²⁰ whose results were probably affected by deviations from stochiometry. The effect of trapping states inside the forbidden gap on photoconductive gain and on the response time was investigated using a dc light "bias" of variable intensity to fill traps to any desired degree.

In order to provide meaningful comparison of the photoconductive

11

properties of amorphous Sb_2Se_3 with single crystalline Sb_2Se_3 , a detailed investigation of the photoconductive properties of the single crystalline phase was performed²¹ (Appendix B). The onset of photoconductivity was found to depend on orientation of electrical vector E of polarized light with respect to three principal crystallographic axes in agreement with dependence of optical absorption on the above parameters.

An interesting feature of the photoconductive kinetics of Sb_2Se_3 is a rather long response time - even in the region of fundamental absorption the response time is of the order of 250-300 micro-seconds, which can be explained only by strong trapping effects which obscure the surface recombination. This long response time probably also gives rise to photosensitivity over wide spectral range (1-3 eV).

So far, we are able to make the following conclusions: The spectral response, particularly the activation energy of amorphous stochiometric Sb_2Se_3 is very similar to that of single crystal Sb_2Se_3 . The onset of photoconductivity in amorphous Sb_2Se_3 is at least that sharp as in the case of the single crystal modification, a fact which seems to be contradictory to effects of band tailing as found by Weiser and Brodsky²² for amorphous As_2Te_3 . Also the response time and photosensitivity is at least two orders of magnitude lower than that in the crystalline phase indicating, together with the previous findings, that trapping states at band edges in amorphous phase of Sb_2Se_3 seem to be electrically much less effective than in the single crystal phase.

IV. Photoemission

Construction of a ultrahigh vacuum ($10^{-10} - 10^{-11}$ torr) phototube

12

has been completed together with provision for sample cleaving in situ. Presently, an attachment is under construction for the evaporation of amorphous Sb-Se films in situ.

Also completed is the whole detection and electronic circuit employing a retarding field detector. A few preliminary runs show that the apparatus, i.e., McPherson 225 Vacuum UV Monochromator, all vacuum parts and all electronic equipment, is in good working condition and initial density of states curves have been recorded. These curves will be compared with density of states from band structure calculations for amorphous and crystalline Sb_2Se_3 . In order to obtain better resolution of transitions at critical points in Brillouen zone partially obscured by phonon broadening and by the absoptive background, first and second derivatives of the density of states vs. energy will be also recorded simultaneously. For this purpose doublers and triplers of the modulation frequency have been constructed.

13

TRANSPORT PROPERTIES

I. <u>Resistivity vs. Temperature</u>

Work has progressed in the low temperature measurement of very high resistance ($\sim 10^{11}$ ohms) thin films. These measurements have been carried out in a "Cryotip", which uses the Joule-Thompson expansion of He and H₂ to provide temperatures from ambient to 4.2^oK. A four-probe configuration of evaporated gold contacts is used to minimize contact resistance. The main problems encountered thus far are those of vacuum integrity, shielding of circuits and ground loops between circuits. These have been solved to the extent that a standard resistance of the same order of magnitude as the film has been measured in the apparatus.

18

REFERENCES

1.	F. M. Mueller, A. J. Freeman, J. O. Dimock, and A. M. Furdyna, Phys. Rev. (in press).
2.	Shaffer, J.C., and Afshar, R., Bull. Am. Phys. Soc., <u>16</u> , 374 (1971).
3.	Shaffer, J.C., Afshar, R., Wood, C., and Mueller, R., Bull. Am. Phys. Soc., <u>16</u> , 519 (1971).
4.	Rubloff, G.W., Freeouf, J., Fritzche, H., and Morase, K., Phys. Rev. Ltrs., <u>26</u> , 1317 (1971).
5.	Maschke, K. and Thomas, P., Phys. Stat. Sol. <u>39</u> , 453 (1970).
6.	Romeris, M.V., J. Math. Phys. <u>12</u> , 552 (1971).
7.	Wood, C., Mueller, R., Gilbert, L.R., Hurych, Z. and Wang, C.C., Bull. Am. Phys. Soc. <u>16</u> , 305 (1971).
8.	Gunther, K.G., Compound Semiconductors <u>1</u> , Ed. Willardson and Goering (Reinhold, 1962).
9.	Gilbert, L.R., and Wood, C., Bull. Am. Phys. Soc. <u>16</u> , 500 (1971).
10.	Ruby, S.L., Gilbert, L.R., and Wood C., Bull. Am. Phys. Soc. <u>16</u> , 303 (1971).
11.	Taneja, S.P., Gilbert, L.R., Harper, W.C., Kimball, C.W., and Wood, C., Bull. Am. Phys. Soc. <u>16</u> , 303 (1971).
12.	B.T. Kolomiets et. al. Collection:The Glassy State. Izd. Akad. Nauk. USSR, Moscow (1960) p. 456.
13.	Z.U. Borisova et. al. Sov. Phys Semiconductors <u>4</u> , 443 (1970).
14.	S.L. Ruby, Mossbauer effect Methodology, vol. 3, Ed. I.J. Gruverman, 1967 p. 203.
15.	E.V. Shkol'nikov, Solid State Chemistry ed. 7.U. Borisova. (consultants Bureau, N.Y. 1966), p. 142.
16.	Van Pelt, B., and Wood. C., Bull, Am. Phys. Soc. <u>16</u> , 306 (1971).
17.	Sobolev, V.V., Shutov, S.D., Popov, Y.V. and Shest'atskii, S.N., Phys. Stat. Sol. <u>30</u> , 349 (1968).
18.	Wood, C., Hurych, Z., and Van Pelt, B., Bull. Am. Phys. Soc. <u>16</u> , 652 (1971).

- 19. Moss, T.S., Photoconductivity in the Elements
- 20. Kolomiets, B.T., Lyubin, V.M., and Tarkin, D.V., Sov. Phys. Sol. State <u>1</u>, 819 (1959).
- 21. Hurych, Z. Wang, C.C. and Wood, C., Bull. Am. Phys. Soc. <u>16</u>, 400 (1971).
- 22. Weiser, K., and Brodsky, M.H., Phys. Rev. B1, 791 (1970).

and in the second design of the second s

FISCAL STATUS

	Amount Budgeted	Amount Expended May 31, 1971
Personne1	\$31,368.00	\$30,741.00
Supplies, Expenses & Services	12,800.00	11,004.00
Travel	2,500.00	2,268.00
Indirect Costs	11,952.00	11,952.00

APPENDIX A

APPENDIX A

A NEW METHOD FOR THE KRAMERS-KRONIG ANALYSIS OF OPTICAL DATA*

R. Afshar

Northern Illinois University, DeKalb, Illinois

F. M. Mueller

Argonne National Laboratory, Argonne, Illinois and Northern Illinois University, DeKalb, Illinois

and

J. C. Shaffer

Northern Illinois University, DeKalb, Illinois

^{*}This work was performed under the auspices of the Atomic Energy Commission and was supported by the Advanced Research Projects Agency of the Department of Defense and was monitored by the Army Research Office, Durham under Contract No. DA-ARO-D-31-124-70-G77.

ABSTRACT

Herein a new technique for performing Kramers-Kronig (Hilbert) transforms upon experimental normal incidence reflectivity data is presented. The method is based upon development of the Hilbert principal value kernel and suitably defined functions closely related to the data in the space of Hermite functions. Well defined procedures for extrapolating the data into unmeasured regions of the spectrum are employed.

The method is applied to reflectivity data taken on two solids and the results are analyzed as to their reliability as is the general utility of the technique. An important conclusion drawn from the results is the invariance of the location of critical point structure under the Kramers-Kronig transform. Some of the details of the numerical procedures and their convergence are also briefly discussed.

I. INTRODUCTION

Great progress has been made in extending the frequency region over which the optical spectroscopy of solids is regularly performed. It is presently relatively commonplace for measurements of, e.g., near-normal incidence reflectivity, to be carried out from far infra-red through vacuum ultra-violet frequencies. Reflectivity measurements on bulk single crystals over wide spectral ranges are made especially valuable by the possibility of performing a Kramers-Kronig analysis upon the data to obtain the phase of the complex reflectivity and hence all of the optical functions of a given material. This general procedure has been adopted by many workers (i-1) and several computational techniques have been developed for carrying out the requisite integral transforms. In Section II these are briefly and critically reviewed. Also in that section the method we have employed in this work is outlined. The general procedure has been separately detailed in a somewhat different $context^{(5)}$ and so the intent here is to concentrate upon those aspects of Kramers-Kronig analysis and of our method which are specific to the analysis of optical data.

In Section III the results of the application of this new method to normal incidence reflectivity Jata on $\text{Sb}_2\text{Se}_3^{(6)}$ and $\text{BaF}_2^{(7)}$ single crystals are presented. The reflectivity data in these two cases were measured over different spectral regions and, additionally, the two sets of data are quite different in structural detail. They thus provide tests of somewhat different

character (e.g., as to resolution) for the technique.

In Section IV an analysis of the accuracy, reliability, and systematic use of the method is given. Studies of the convergence of the necessary numerical expansion procedures are presented as³⁵ discussion of some general conclusions on the characteristics of the Hilbert transform as it relates to the optical properties of solids. Possible applications to other problems in solid state physics are considered.

II. THE KRAMERS-KRONIG (HILBERT) TRANSFORM APPLIED TO OPTICAL DATA

If it were possible to measure the (near-normal incidence) fractional coefficient of reflected intensity, or reflectance, $R(\omega)$, of a bulk solid sample (for a given orientation and polarization) in vacuo (air) for all frequencies then the phase of the complex (amplitude) reflection coefficient $\hat{r}(\omega) = R^{1/2}(\omega)e^{i}\varphi(\omega)$ could be determined from the Kramers-Kronig dispersion relation $\begin{pmatrix} 8 \\ \end{pmatrix}$

$$\varphi(w) = \frac{P}{\pi} \int_{-\infty}^{+\infty} \frac{\ln r(w') dw'}{w' - w}$$
(1)

where $r(w') = R^{1/2}(w')^*$, and R(w) has been extended to negative

^{*}Most often dispersion relations are derived for the real and imaginary parts of complex causal response functions which of a real variable must be of class $L^2(-\infty, \infty)$. However such relations can also be derived for the phase and logarithm of the modulus of complex functions for which $\lim_{\omega \to \infty} r(\omega) \to 0$ at least as rapidly as ω^{-n} where n > 1. Thus although the logarithm of the reflectivity is not $L^2(-\infty, +\infty)$ a dispersion relation exists between it and the

is not $L^2(-\infty, +\infty)$ a dispersion relation exists between it and the phase. For causal response functions the real and imaginary parts are each of class $L^2(-\infty, \infty)$. An excellent discussion of these points is given in Ref. (5).

frequencies as an even function. Knowledge of $R(\omega)$ and $\varphi(\omega)$ is tantamount to knowledge of all the optical functions since $\hat{n}(\omega) = n(\omega) + i_{R}(\omega)$ is related to the complex reflection coefficient by

$$\hat{n}(\omega) = (1 + \hat{r}(\omega))/(1 - \hat{r}(\omega))$$
, (2)

a simple homographic transformation. Given $\hat{H}(\omega)$; $\hat{e}(\omega)$, the complex dielectric function, and the energy loss function, $\prod_{i=1}^{\infty} [\hat{e}^{-1}(\omega)]$, are easily obtained.

There are two paramount difficulties in Eq. (1) which must be overcome in order that this data reduction be effected. In the first place data is never available for all frequencies due to the finite frequency cutoff of real spectrometers. Secondly the kernel of the Hilbert transform in Eq. (1) is highly singular and, in fact, that equation is most properly regarded as a functional and the kernel as a generalized function. A casual numerical attack upon it or any of its equivalents can be rather precarious particularly in its effect upon the critical point structure exhibited by the spectrum under consideration.

It is useful to review some of the methods which have been employed to deal with the aforementioned problems. Generally the first problem, that of finite data ranges has been handled by extrapolating the data into unmeasured regions. Above the highest frequency the data is usually extended using one or more terms of a Lavrent series. Unless great care is taken this may introduce spurious structure into the spectrum. Below the lowest frequency measured the data is normally extrapolated

frequencies as an even function. Knowledge of R(w) and $\varphi(w)$ is tantamount to knowledge of all the optical functions since $\hat{n}(w) = n(w) + i_{X}(w)$ is related to the complex reflection coefficient by

$$\hat{n}(\omega) = (1 + \hat{r}(\omega))/(1 - \hat{r}(\omega)) , \qquad (2)$$

a simple homographic transformation. Given $\Re(w)$; $\widehat{\mathfrak{c}}(w)$, the complex dielectric function, and the energy loss function, $\prod_{i=1}^{\infty} [\widehat{\mathfrak{c}}^{-1}(w)]$, are easily obtained.

There are two paramount difficulties in Eq. (1) which must be overcome in order that this data reduction be effected. In the first place data is never available for all frequencies due to the finite frequency cutoff of real spectrometers. Secondly the kernel of the Hilbert transform in Eq. (1) is highly singular and, in fact, that equation is most properly regarded as a functional and the kernel as a generalized function. A casual numerical attack upon it or any of its equivalents can be rather precarious particularly in its effect upon the critical point structure exhibited by the spectrum under consideration.

It is useful to review some of the methods which have been employed to deal with the aforementioned problems. Generally the first problem, that of finite data ranges has been handled by extrapolating the data into unmeasured regions. Above the highest frequency the data is usually extended using one or more terms of a Lavrent series. Unless great care is taken this may introduce spurious structure into the spectrum. Below the lowest frequency measured the data is normally extrapolated

using a Taylor series. Knowledge of the static dielectric constant of the material being studied can make this a most reasonable procedure. The singular kernel in (1) is usually handled by transforming (1) into the equally valid form

$$\varphi(\omega) = \frac{2\omega}{\pi} \int_{0}^{\infty} \frac{\mu_{np}(\omega') - \mu_{np}(\omega)}{\omega'^{2} - \omega^{2}} d\omega' \qquad (3)$$

The inclusion of a small additional part in the denominator of (3), η^2 , is then justified by the principal value character of (1) and if it is chosen sufficiently small the finite arithmetic character of digital computation allows the transform to carried out in an approximate manner. It is interesting to examine in more detail the methods used by some of the workers who have attacked the Kramers-Kronig transform in optical problems.

In another publication (5) (hereafter denoted I) the suitably of the Hermite functions (harmonic oscillator eigenfunctions) as a basis set for accomplishing Hilbert transforms has been demonstrated. The interest there was in the Hilbert transforms of electron (or phonon) densities of states and thus the functions to be transformed were of compact support, i.e., they vanished outside some finite range on the real line, and were bounded. In the application of Kramers-Kronig analysis to reflectivity data these characteristics are not present however as will be shown suitable treatment of the data brings the problem into a form amenable to the methods described in I.

In I it is shown that if a function $\alpha(\omega)$ have a convergent Hermite expansion

29

APPENDIX A

-7-

$$\alpha(\omega) = \sum_{n} \alpha_{n} \phi_{n}(\omega) \qquad (4)$$

where the $\phi_n(\omega)$ are the Hermite functions, then its Hilbert transform partner $\beta(\omega)$ also has a Hermite expansion

$$\beta(\omega) = \sum_{n} \beta_{n} \phi_{n}(\omega)$$
 (5)

where

$$\beta_n = \prod_{m=1}^{\Sigma} S_{nm} \alpha_m , \qquad (6)$$

 S_{nm} being a compact matrix representation of the Hilbert transform "operator" in the space of Hermite functions.* S_{nm} vanishes unless n + m is odd and, for integer h, may be written

$$S_{nm} = (-1)^{h} + 1 S_{nm}^{i}; h = (m-n-1)/2$$
 (7)

Computation of the matrix elements S_{nm}^{i} is most easily accomplished as in I; from

$$S'_{nm} = \int_{-\infty}^{+\infty} dt \ SGN(t)\phi_n(t)\phi_m(t) \ . \tag{8}$$

^{*} It can be shown (8) that causal complex response functions which are the simplest class of functions whose real and imaginary parts are Hilbert transform pairs are necessarily square integrable as are their real and imaginary parts separately. (8) However the class of square integrable functions does not exhaust all possible Hilbert transform pairs (consider $\phi(\omega)$ and $tnr(\omega)$ in (1)). The Hermite decomposition of the Hilbert transform can be directly applied only to the squareintegrable case because of the need for convergent Hermite expansions to exist.

The matrix elements are then computed by using the Hermite integration technique, as detailed in (I).

APPENDIX A

- 6 -

Direct application of this technique to (1) is obviated since as $w \to \pm \infty r(w) \to 0$, hence $\mu r(w) \to -\infty$ as $w \to \pm \infty$. The function $\mu r(w)$ is therefore not a square integrable function and has no convergent expansion in Hermite functions. A simple procedure however removes this difficulty allowing the full power of the Hermite analysis to bear upon the structure in the optical problem as well as providing a self consistent extrapolation of the data into unmeasured regions. Since the Hilbert transform (1) is a linear integral transform portions of $\mu r(w)$, the transforms of which are closed-form analytic functions, can be removed leaving a residual function whose transform must be attacked numerically. The portions removed are chosen so that they smoothly fit to $\mu r(w)$ itself near the boundaries of measurement and furthermore give reasonable extrapolations of $\mu r(w)$

$$\varphi(\omega) = \varphi_0(\omega) + \theta(\omega)$$
(9)

whe re

$$\varphi_{O}(\omega) = \frac{P}{\pi} \int_{-\infty}^{+\infty} \frac{g(\omega') d\omega'}{\omega' - \omega}$$
(10)

where $g(\omega)$ is the function which was used to fit $lnr(\omega)$ near the boundaries of measurement and to extrapolate into the unknown regions. $g(\omega)$ should be chosen from among functions with known analytic Hilbert transforms $\varphi_0(\omega)$. If we denote $lnr(\omega) - g(\omega)$

by $\Delta(\omega)$ then

$$\theta(\omega) = \frac{P}{\pi} \int_{-\infty}^{+\infty} \frac{\Delta(\omega^{\dagger}) \, d\omega^{\dagger}}{\omega^{\dagger} - \omega}$$
(11)

By construction $\Delta(\omega')$ is a compact function which is, in practice, also bounded. It is, therefore, square integrable and the Hermite analysis technique can be used to compute $\theta(\omega)$. It is then only necessary to choose appropriate $g(\omega)$. Our procedure has been as follows. The dielectric constant or index of refraction of a system of classical change oscillators with given resonant frequencies, damping constants, and oscillator strengths = iswell-known. For example, the index of refraction can be approximated (for a dilute gas of Gach oscillators) as

$$n_{0}(\omega) = 1.0 + \sum_{i} \frac{v_{i}}{(\omega_{i}^{2} - \omega^{2}) + i\Gamma_{i}\omega}$$
(12)

in which the ω_{j} are the resonant frequencies, the Γ_{j} the damping constants (linewidths) and the v_{j} essentially the oscillator "strengths". The complex amplitude reflection coefficient of a bulk sample of material with this \hat{n}_{j} is

$$p_{o}(\omega)e^{i\phi_{o}(\omega)} = \hat{p}_{o}(\omega) = (\hat{n}(\omega) - 1)/(\hat{n}(\omega) + 1)$$
(13)

By adjusting the various parameters in (12) we can find an $f_{\mu}(\omega)$ for which the resulting $\mu n p_0(\omega)$ will smoothly and closely approximate $\mu nr(\omega)$ near the data boundaries. The same function (13) provides a natureal choice of extrapolation of $r(\omega)$ and hence $\mu nr(\omega)$ into the unmeasured regions. $\varphi_O(\omega)$, the analytic portion we have extracted from the total Hilbert transform is then given by

$$\varphi_{0}(\omega) = \tan^{-1} \frac{2 \kappa(\omega)}{n_{0}^{2}(\omega) + \kappa_{0}^{2}(\omega) - 1}$$
(14)

where, of course, $n_{(\omega)}^{(\omega)}$ and $\kappa_{(\omega)}^{(\omega)}$ are the real and imaginary parts of $\hat{f}_{(\omega)}^{(\omega)}$. For the data analyzed in this contribution we have found that choosing two frequencies for classical oscillators and adjusting the other parameters (Γ_1 , Γ_2 , ν_1 , ν_2) allows us to obtain a sufficiently smooth fit to the data in the relevant regions. Note that $R(\omega) = |\mathbf{r}(\omega)|^2$ has a dominant ω^{-4} dependence at sufficiently high frequencies in the extrapolation here employed. The entire data analysis problem has now been reduced to the computation of Hermite expansion coefficients of $\Delta(\omega)$ and matrix multiplication of these to generate the expansion coefficients of $\theta(\omega)$. The next section is devoted to the details of the procedure we have described as it has been applied to data on Sb₂Se₃ and BaF₂ single crystals.

-11-

III. APPLICATION TO REAL SOLIDS

A. Optical Functions of Sb₂Se₃.

 Sb_2Se_3 , in crystalline form, has an orthorhombic lattice with a rather complex unit cell containing four units of Sb₂Se₃ with two nonequivalent sites both for the antimony and selenium atoms. The space group is D_{2h}^{16} (Pnma). The optical functions are anisotropic, however, in this work we ignore that fact as it is not pertinent to our purpose of illustrating the use of our method of Kramers-Kronig analysis. In Fig. 1 is shown the reflectivity to unpolarized light, in the frequency range 1.5eV to 11.0eV, of a cleavage plane of Sb_2Se_3 as measured by Wood⁶ et al. This data is in close agreement with the data of Shutov⁹ et al in overlapping measured regions of the spectrum. The data, as may be seen, indicates an absorption edge near 1.0eV and a series of peaks through the high reflectivity region from the absorption edge through the visible and extending into the vacuum ultra-violet region. This data, of course, must be extrapolated into the unmeasured regions by the methods we have outlined in Section II, above. The remainder of this subsection details the extrapolation and other aspects of the method.

The data were analyzed as follows. It was found that a minimum of two sets of classical charge oscillators with different resonant frequencies was required in order to extrapolate the data consistently both at high frequencies and at low frequencies using (12) and (13). The criteria for good extrapolation that were employed were (a) the residual portion, i.e., $\Delta(\omega)$ in (11) should be as small as reasonably possible at the boundaries

34

material . Sometriel

of the data, and (b) $\Delta(\omega)$ should be rather smooth function. These criteria insure the applicability of the analysis in II above as well as good convergence of the Hermite expansion of $\Delta(\omega)$. For the reflectivity spectrum in Fig. 1 it was found that a good extrapolation was obtaining by choosing $\hat{n}_{\alpha}(\omega)$ from (12) as

$$\hat{n}_{0} = 1 + \frac{20}{((2.5)^{2} - \omega^{2}) + i(1)(\omega)} + \frac{90}{((11.8)^{2} - \omega^{2}) + i(8.5)(\omega)}$$
(15)

Of course the function above is unit-free, however, the units of frequency in which it can be interpreted are electron volts. Hence we have used charged oscillator systems with resonant frequencies 2.5 and 11.8 eV. Fig. 2 shows the function $\Delta(\omega)$ which is obtaining by using for the $g(\omega)$ of section II ln $\rho_0(\omega)$ as computed from (15) and subtracting that function from ln r (ω) obtained from the data. The contribution to the phase from ln $\rho_0(\omega)$ is obtained from (14). $\Delta(\omega)$ has been expanded in a Hermite series to order 250 for which the expansion coefficients have been computed by Simpson's rule with the intrgrand evaluated at each 0.05 eV interval. Since the data extends to only 12.0 eV and the 250th Hermite function becomes essentially zero beyond the "classical turning point", ω top = $\sqrt{501}$, it is useful to scale in frequency in order to optimize the use of the higher order Hermite functions in the expansion. Thus the Hermite expansion coefficients of $\Delta(\omega)$ are given by

$$\Delta_{n}(\alpha) = \int_{-\alpha\omega top}^{\alpha\omega top} \Delta_{1}(\alpha\omega) \phi_{n}(\alpha\omega) d(\alpha\omega)$$
(16)

where $\Delta_1(\alpha\omega) = \Delta(\omega)$.

£
-13-

If $\alpha = \sim 1.77$ we have found that the mean square error between $\Delta(\omega)$ and its Hermite expansion is made minimal. The error function $\chi(\omega)$ is plotted in Fig. <u>3</u> and in that figure $\langle \chi^2(\omega) \rangle_{250}$ is given as 4.54×10^{-6} . Thus the expansion of $\Delta(\omega)$ is accurate to an r.m.s. value of order 10^{-3} . Since we know the matrix S_{nm} to be compact the convergent expansion of $\phi(\omega)$ is easily computed from (6). The logarithm of the reflectivity amplitude and its phase, $\phi(\omega)$, are shown in Fig. 4 for Sb₂Se₃.

The Kramers-Kronig analysis is now complete and the optical functions of the system can easily be computed from (2) and related equations. Fig. 5 shows the real and imaginary parts of the index of refraction which we have obtained and Fig. 6 the real and imaginary parts of the complex dielectric function. It may be seen that the frequency positions of critical points are preserved through the transform process by comparing Figs. (5) and (6) with Fig. (1).

APPENDIX A

References

1.	H.	R.	Phillip	op and	Ε.	Α.	Taft,	Phys.	Rev.	<u>113</u> ,	1002	(1959).	

- 2. H. R. Phillipp and E. A. Taft, Phys. Rev. <u>136</u>, A1445 (1964).
- 3. H. R. Phillipp and H. Ehrenreich, Phys. Rev. <u>129</u>, 1550 (1963).
- 4. M. Cardona, Chap. VI, <u>Optical Constants of Insulators</u>, Ed. (Nudelman and Mutra), Plenum Press (1969).
- 5. Afshar, R., Mueller, F. M., and Shaffer, J. C. (submitted for publication).
- 6. Van Pelt, B., and Wood, C., Bull. Am. Phys. Soc., 16, 306 (1971).
- Rubloff, G. W., Freeouf, J., Fritzche, H., and Morase, K., Phys. Phys. Rev. Ltrs., <u>26</u>, 1317 (1971).
- 8. Page, C. H., <u>Physical Mathematics</u>, Van Nostrand (1955).
- Shutov, S. D., Sobolev, V. V., Popov, Y. V., and Shestataskii, S. N., Phys. Stat. Sol. <u>31</u>, K23, (1969).











en - senten anderskalte bestämmte



ļ

.

Abstract Submitted

For the 1971 March Meeting of

The American Physical Society

<u>29 March - 1 April 1971</u> Date

Physical Review Analytic Subject Index Number <u>44.4</u> Bulletin Subject Heading in which Paper should be placed Optical Properties of Solids

The Optical Functions of Crystalline Sb_2Se_3 from a Kramers-Krönig Analysis.* J. C. SHAFFER and R. AFSHAR, Northern Illinois Univ.--In this contribution we present the results of a Kramers-Krönig analysis of normal incidence reflectivity data taken on Sb_2Se_3 single crystals. The computational method¹ employed for the data analysis which is based upon expansion in Hermite functions and suitable high frequency extrapolations is presented as are the resulting optical functions. The accuracy and computational utility of the method and the relationship of the optical functions to the electronic structure of Sb_2Se_3 are discussed.

*Supported by A.R.P.A. through A.R.O., Durham. ¹R. Afshar, F. M. Mueller, and J. C. Shaffer, (to be published).

Submitted by

Signature of APS

John C. Shaffer Same name typewritten

Department of Physics, NIU Address

DeKalb, Illinois 60115

Abstract Submitted

For the 1971 March Meeting of

The American Physical Society

Physical Review Analytic Subject Index Number <u>47.4</u> Bulletin Subject Heading in which Paper should be placed <u>Semiconductors - Optical</u> Properties

Optical Properties of Amorphous Antimony Selenides. * C. WOOD, R. MUELLER, L.R. GILBERT, Z. HURYCH and C.C. WANG, Northern Illinois Univ.--Amorphous films of antimony selenides were prepared by the standard technique of vacuum evaporation from the compound Sb₂Se₃. Micro-probe analysis showed that the films deviate considerably from the stoichiometry the exact composition depending on the source temperature.¹ Evaporation from melts of $SbSe_{0.9}$ yielded amorphous films of Sb_2Se_3 . Amorphous films of various compositions were also prepared by controlled coevaporation of Sb and Se from separate sources. Optical transmission, reflectivity and photoconductivity measurements showed large changes in the forbidden energy gap as a function of composition. In contrast to many other amorphous materials and published data on antimony selenide films² the forbidden energy gaps of amorphous and single crystal Sb₂Se₃ are approximately equal.

*Supported by A.R.P.A. through A.R.O., Durham. ¹A. Efstathion, D. M. Hoffman and E. R. Levin, J. Vacuum Sci. and Tech. <u>6</u>, 383 (1969). ²B. T. Kolomiets, V. M. Lyubin and D. V. Tarkhin, Sov. Phys. Sol. State 1, 819 (1959).

Submitted by

. M. C. Word

Signature of APS Member

Charles Wood Same Name typewritten

Department of Physics,NIU Address

DeKalb, Illinois 60115

Abstract Submitted

For the 1971 March Meeting of

The American Physical Society

29 March - 1 April 1971 Date

Physical Review Analytic Subject Index Number <u>48.3</u> Bulletin Subject Heading in which Paper should be placed Amorphous Solids



*Partially supported by A.R.P.A. through A.R.O., Durham. ¹N.W. Tideswell, F.H. Kruse, and J.D. McCullough, Acta Cryst. <u>10</u>, 99 (1957). ²L.I. Tatarinova, Kristallografiya <u>4</u>, No. 5, 678 (1959).

Submitted by

Signature of APS Member

S. L. Ruby Same name typewritten

47

Argonne National Lab. Address

Argonne, Illinois 60439

APPENDIX 8

Abstract Submitted

For the 1971 March Meeting of

The American Physical Society

29 March - 1 April 1971 Date

Physical Review Analytic Subject Index Number 48.3 Bulletin Subject Heading in which Paper should be placed Amorphous Solids

Mossbauer Investigation of Sn_x(As₂Se₃)_{1-x} Glasses.* S.P. TANEJA, L. GILBERT, W.C. HARPER, C.W. KIMBALL, and C. WOOD, Northern Illinois Univ. -- The dependence of conductivity and microhardness of glasses of the Sn-Se-As system has been attributed by Shkol'nikov to a transition from SnSe2 to SnSe with increasing tin concentration. However, Borisova et al² infer that SnSe, is found only in the glassy state and SnSe only in the crystalline state. In the present investigation of glasses of $Sn_x(As_2Se_3)_{1-x}$ (with x = 2.5,5.0,8.0,10.0 atomic percent) were prepared by quenching the melts into a brine bath at $\sim 200^{\circ}$ K. X-ray patterns show distributions characteristic of a glass; further, the patterns are qualitatively similar for all compositions. The Mossbauer study confirms the SnSe₂-SnSe transition with increasing tin concentration. The line width and concentration dependence of the isomer shift indicate a high degree of local order. The question of the presence of microcrystallinity will be discussed.

*Work supported by A.R.P.A. through A.R.O., Durham. ¹E. V. Shkol'nikov, Solid State Chemistry ed. Z. N. Borisova (Consultants Bureau, N.Y. 1966) p. 142. ²Z. U. Borisova et al, Sov. Phys. - Semiconductors <u>4</u>, 443 (1970).

48

Submitted by

Signature of APS Member

Clyde W. Kimball Same name typewritten

Physics Department, NIU Address

DeKalb, Illinois 60115

Abstract Submitted

For the 1971 March Meeting of

The American Physical Society

<u>29 March - 1 April 1971</u> Date

Physical Review Analytic Subject Index Number 47.4 Bulletin Subject Heading in which Paper should be placed Semiconductors - Optical Properties

<u>Optical Properties of Sb_2Se_3 Single Crystals.</u>* B. VAN PELT and C. WOOD, Northern Illinois Univ.--The optical absorption and transmission of single crystal Sb_2Se_3 has been measured in the region of the absorption edge with respect to the principal crystallographic axes. The optical transitions responsible for the edge will be discussed.

*Supported by A.R.P.A. through Army Research Office, Durham.

Submitted by

ila Wood

Signature of APS member

<u>Charles Wood</u> Same name typewritten

Department of Physics,NIU Address

DeKalb, Illinois 60115

Please place this paper together with and preceeding the accompanying paper Photoconductivity in Single Crystals of Sb₂Se₃ by Z. Hurych, et al.

Abstract Submitted

For the 1971 March Meeting of

The American Physical Society

<u>29 March - 1 April 1971</u> Date

Physical Review Analytic Subject Index Number 47.5 Bulletin Subject Heading in which Paper should be placed Semiconductors - Transport Properties

Photoconductivity in Single Crystals of Sb, Se. * Z. HURYCH, C.C. WANG and C. WOOD, Northern Illinois Univ.—The activation energy ΔE_a , the response time τ and the spectral distribution of photocurrent Iph were measured for p-type semiconducting orthorhombic single crystals of Sb₂Se₃ as a function of the orientation of the electric vector E₀ of the polarized light and of the drift field Ed with respect to the three crystallographic axes, a, b, and c. Values of ΔE_a were 1.09, 1.10 and 1.13 eV for $E_0 || a$, b, and c, respectively, in good agreement with corresponding optical activation energies and with published data on photoconductivity using nonpolarized light.¹ Values of τ were in the region 300-400 μ sec for $h\nu = 1.1eV$, and indicate presence of trapping levels inside the forbidden gap. The spectral distribution of τ and I_{ph} as a function of E_0 and E_d orientations will be discussed.

*Supported by A.R.P.A. through A.R.O., Durham. ¹B. T. Kolomiets and A. Kh. Zeinally, Sov. Phys. Sol. State 1, 896 (1959).

Submitted by

- Word

Signature of APS member

<u>Charles Wood</u> Same name typewritten

Department of Physics,NIU Address

DeKalb, Illinois 60115

Please place this paper together with and following the accompanying paper Optical Properties of Sb₂Se₃ Single Crystals by B. Van Pelt and C. Wood. 50 APPENDIX C

.

APPENDIX C

Abstract Submitted

For the 1971 April Meeting of

The American Physical Society

<u>26 April - 29 April 1971</u> Date

Physical Review Analytic Subject Index Number <u>44.4</u> Bulletin Subject Heading in which Paper should be placed Optical Properties of Solids

The Optical Constants of Thin Amorphous Films of Sb. Se. .* J.C. SHAFFER, R. AFSHAR, C. WOOD, and R. MUEL-LER, Northern Illinois Univ.--The optical functions of thin films are often difficult to obtain due to complications arising from the occurrence of interference in their reflection and transmission spectra. We present and discuss ar. experimental and computational method for circumventing this problem. We obtain from reflectance measurements via Kramers-Kronig analysis the optical functions of films, sufficiently thick (>5µ) that interference effects are largely suppressed. These optical functions are employed to compute exactly the reflectance and transmittance for thinner (\sim 1,000Å) film, with identical optical functions. The results are compared with experimental reflectance and transmission of thinner films of the same composition. These films are prepared simultaneously with the thick films by co-evaporation. The method is applied to amorphous films of Sb_xSe_y and comparison made with the optical functions of crystalline Sb₂Se₃.

*Supported by A.R.P.A. through Army Researh Office, Durham.

Submitted by

Signature of APS Men

John C. Shaffer Same name typewritten

Physics Department, NJU Address

DeKalb, Illinois 60115

APPENDIX C

Abstract Submitted

For the 1971 April Meeting of

The American Physical Society

<u>26 April - 29 April 1971</u> Date

Physical Review Analytic Subject Index Number 48.3 Bulletin Subject Heading in which Paper should be placed Amorphous Solids

<u>Glass-Crystalline Transitions in Amorphous</u> <u>Sb_Sey</u>.* L.R. GILBERT and C. WOOD, Northern Illinois Univ. The glass transition temperature in amorphous antimony selenides shows a strong dependence on composition. At the composition Sb₂Se₃ the transition occurs at $\sim 180-200^{\circ}$ C in agreement with the nucleation studies of Bolotov et al. The transition temperatures were determined by differential thermal analysis and resistivity versus temperature measurements and are compared.

*Supported by A.R.P.A. through A.R.O., Durham. ¹I.E. Bolotov, A.V. Kozhin, and S.B. Fisheleva, Sov. Phys.-Cryst. <u>15</u>, 461 (1970).

Submitted by

. Chah Lood

Signature of APS Member

Charles Wood Same name typewritten

Physics Department, NIU Address

DeKalb, Illinois 60115

Abstract Submitted

for the 1971 April Meeting of

The American Physical Society

26 April - 29 April 1971 Date

Physical Review Analytic Subject Index Number 47.4 Bulletin Subject Heading in which Paper should be placed Semiconductors - Optical Properties

Thermally Modulated Reflectivity of Sb₂Te₃.* C. WOOD, Z. HURYCH, and B. VAN PELT, Northern Illinois Univ.--The thermally modulated reflectivity has been determined in the visible and near IR region of the spectrum for single crystals of Sb₂Te₃ with the electric vector of the radiation in the cleavage plane (E|C). The data show the same critical points as observed in the optical transmission spectrum of very thin cleaved specimens using a photon counter. These results are compared with the low temperature reflectivity data of Sobolev et al.¹

*Supported by A.R.P.A. through A.R.O., Durham. ¹V.V. Sobolev, S.D. Shutov, Yu. V. Popov, and S.N. Shestatskii, phys. stat. sol. <u>30</u>, 349 (1968).

Submitted by

Signature of APS Member

Charles Wood Same name typewritten

Physics Department, NIU Address

DeKalb, Illinois 60115

1

55

.

MOSSBAUER STUDIES ON AMORPHOUS AND CRYSTALLINE ANTIMONY SELENIDES

S. L. Ruby Physics Division Argonne, Illinois

and

L. R. Gilbert* and C. Wood* Physics Department Northern Illinois University DeKalb, Illinois 60115

Mossbauer resonance spectra at 4° K have been determined for a range of compositions of amorphous SbSe_x films, where 0.5<x<9, and were compared with the spectrum of crystalline Sb₂Se₃. The crystal structure of Sb₂Se₃ is orthorhombic Pbnm ($D_{2h}^{16}D_8^5$) consisting of infinite chains parallel to the c-axis. There are two different equally populated Sb sites and three different Se sites within the crystal lattice. The strongest bonds are within the chains and here the Sb-Se bond distances range from 2.576Å to 2.777Å, while the 'non-bonded' Sb-Se separations range from 2.98Å to 3.74Å¹.

The Mossbauer spectrum of a powdered sample of crystalline Sb_2Se_3 is shown twice in Fig. 1. The solid line in the upper curve is the (unsuccessful) result of a computer fit assuming a single Sb lattice site. In the lower fit, the computer was allowed two different, but equally populated Sb sites. Now χ^2 shrinks to 1.2 versus 7.3 in the

^{*} This research was supported by the Advanced Research Projects Agency of the Department of Defense and was monitored by the Army Research Office, Durham, under Contract No. DA-ARO-D-31-127-70-G77.

summarize, it seems that for a best fit a range in e^2qQ and S_I are required averaging near the values given above.

The above results show that the short range order in amorphous $SbSe_{\chi}$ differs from that of the crystalline solid, and indicates a close conformity in the environment, i.e., a narrow range of coordination number and Sb-Se bond lengths, of each Sb atom in the amorphous solid. Electron diffraction studies of Tatarinova² lend support to this view-point. Intensity and radial distribution curves indicate that both the coordination number and nearest neighbor distances decrease in the transition from the crystalline to the amorphous phase of "Sb₂Se₃". (The composition may have been Se-rich from the method of preparation described in the article). In contrast, As_2Se_3 and GaAs were shown to retain in the amorphous phase the configuration inherent in the crystalline phase.

The above results resemble those from previous Mossbauer studies on glassy materials in which $Fe_{2}O_{3}$ was dissolved in both silicate and phosphate glasses³. The microenvironments of the Fe^{3+} ion in each case were found to be similar although the coordination was tetrahedral in the silicate and octahedral in the phosphate glass. Despite the glassy nature of the material in these cases, as well as in the antimony work done here, there does not seem to be a wide variety of microenvironments. However, in the amorphous Sb-Se_x films, we have a different microenvironment than in the crystalline material.

These results are of significance to questions of whether a vaporquenched solid is truly a glass or consists of micro-crystals; particularly

upper fit. The Mossbauer isomer shifts for the two peaks were 12.6 mm/sec and 15.5 mm/sec.

The amorphous thick (\sim 30 microns) films were prepared by vacuum evaporation onto room-temperature quartz substrates from large sources of various ratios of Sb and Se. The films were determined to be amorphous by x-ray diffraction. The composition as deposited (invariably different from the source material) was determined by electron microprobe and/or x-ray emission spectroscopy. For example, a source at a temperature of 720°C with x = 1.5 (i.e., Sb₂Se₃) produced a film with x \sim 9: Amorphous films of various compositions in the range 0.5<x<9 were preapred and measured and the results are shown in Table !. In all compositions of amorphous SbSe_x only a single peak was obtained which was located at an intermediate position between the double peaks in the crystalline material, as illustrated in Fig. 2.

In ¹²¹Sb Mossbauer spectroscopy, working between nuclear levels with spins of 7/2 and 5/2, a pure quadrupole interaction gives rise to an asymmetric octet. In the present case, the quadrupole interaction is not large enough compared to the minimum line width of 7.1 mm/sec to resolve these individual lines. It is found (see Fig. 1) that the data for the amorphous films can be moderately well fitted by a single site where each member of the octet has a normal width Γ =2.8 mm/sec, the value of e^2qQ is 10.5 ± 1 mm/sec and the isomer shift is -13.5 ± 0.5 mm/sec. By adding more freedom to the computer program (allowing more sites), χ^2 can be further decreased. To

in systems where the glass cannot be readily formed from a melt-quench, as in the case of Sb_2Se_3 . We can state that if the film is an assemblage of microcrystals, the crystal structure does not correspond to the equilibrium phase and the more probable conclusion is that it is amorphous. The occurrence of a softening temperature (T_g) in the films would support this conclusion.

The authors wish to thank Dr. Kimball for many helpful discussions.

REFERENCES

- N. W. Tideswell, F. H. Kruse and J. C. McCullough, Acta Cryst. <u>10</u>, 99 (1957).
- 2. L. I. Tatarinova, Kristallografiya 4, No. 5, 678 (1959).
- 3. C. R. Kurkjian and E. A. Sigety, Phys. Chem. Glasses, <u>9</u>, 3 (1968).

1

SbSe _x	Phase	x ²	SI	e ² qQ	β
x = 6.3	Amorphous	. 374	-13.97	9.64	3.39
4.4	Amorphous	.575	-14.42	9.70	2.68
2.0	Amorphous	.531	-113.38	10.85	3.26
1.5	Amorphous	1.206	-13.65	10.90	3.04
1.5	Cryst.	.253	-12.60	7.86	2.48
			-15.50	8.40	2.49
0.5	Amorphous	. 466	-13.00	11.50	2.58

TABLE 1





phys. stat. sol. 42, 871 (1970)

Subject classification: 20.1; 13.1; 22.7; 22.8

Department of Physics, Northern Illinois University, DeKalb

The Optical Properties of Sb₂Se₃-Sb₂Te₃¹)

By

W. PROCARIONE²) and C. WOOD

The optical properties of single crystal specimens of $Sb_2Tc_3-Sb_2Se_3$ solid solutions have been investigated in the photon energy range 0.04 eV to 6.5 eV. A wide range of solid solutions having hexagonal structure were found to exhibit properties closely corresponding to the isostructural compounds Bi_2Te_3 and Bi_2Se_3 . Small indirect band gaps approaching semimetallic values were obtained. The high free-carrier concentrations gave rise to plasma effects in the near infra-red. In contrast Sb_2Se_3 is orthorhombic with a direct band gap near 1 eV. The chemical bonding concepts applied to similar systems seem applicable with only slight modification.

Die optischen Eigenschaften von Sb₂Te₃-Sb₂Sc₃-Mischkristallen wurden im Bereich von 0,04 eV bis 6,5 eV untersucht. Die Festkörperlösungen haben innerhalb weiter Grenzen die gleiche hexagonale Struktur wie die isostrukturellen Verbindungen Bi₂Te₃ und Bi₃Se₃ und zeigen sehr verwandte optische Eigenschaften. Kleine indirekte Energieschwellen ähnlich denen in Halbleitern wurden beobachtet. Die große Konzentration von Seien Ladungsträgern verursachte Plasmaerscheinungen in nahen Ultrarot. Sb₂Se₃ ist orthorhombisch mit einem direkten verbotenen Energieband von 1 eV. Die chemischen Bindungskonzepte können wahrscheinlich mit nur geringen Veränderungen auf ähnliche Systeme angewendet werden.

1. Introduction

Binary compounds of the form $A_2^V B_3^{VI}$ have been studied extensively in the past. Solid solutions of these compounds, particularly the isostructural systems of $Bi_2Te_3-Bi_2Se_3$ and $Bi_2Te_3-Sb_2Te_3$, have received considerable attention in the literature. A few serious attempts have been made to study the detailed band structure of these compounds and their solid solutions; noteworthy among these being the optical work of Greenaway and Harbeke [1] on $Bi_2Te_3-Bi_2Se_3$, and Sobolev et al. [2, 3], on Bi_2Te_3 , Bi_2Se_3 , and Sb_2Te_3 .

Except for some thermoelectric data by Teramoto and Takayanagi [4], the information on the physical properties of solid solutions of Sb₂Te₃ and Sb₂Se₃ appears to be almost non-existant. However, solid solutions of these two compounds represent an interesting system to study. Although possessing different crystallographic structures, Sb₂Se₃ is orthorhombic ($D_{2h}^{1h} D_{8}^{5}$) and Sb₂Te₃ is rhombohedral ($C_{33}D_{34}^{5}$) or hexagonal on a larger unit cell, a considerable range of solid solubilities exist in this system. X-ray analysis by Teramoto and Takayanagi [4] revealed that a single-phase hexagonal structure exists between Sb₂Te₃ and Sb₂TeSe₂ with the lattice constant, *a*, decreasing approximately from 4.27 Å to 4.12 Å and the lattice constant, *c*, decreasing approximately from 30.4 Å to 29.5 Å with increasing Sb₂Se₃ content. Higher concentrations of Sb₂Se₃ resulted in an admixture of hexagonal and orthorhombie phase.

¹) Partially supported by A.R.P.A. through the Army Research Office, Durham,

²) This work partially fulfilled the requirements for the M.S. Degree in Physics.

56*

As a start toward understanding the physical properties of this system, and obtaining a knowledge of the band structure, reflection and transmission measurements on single crystals in the range 0.04 eV to 6.5 eV were investigated. Because of the marked directional preference in cleavage, (0001) for hexagonal and (010) for orthorhombic crystals, and the difficulty of preparing good reflecting surfaces perpendicular to the cleavage planes, optical measurements have been made only with the cleetric vector in the cleavage plane.

Comparison of this data with the results of Greenaway and Harbeke [1] on $Bi_3Te_3-Bi_2Se_3$, and Shutov et al. [3] on the end components, Sb_3Te_3 and Sb_3Se_3 has shown many similarities between these systems but has raised questions concerning the chemical bonding concepts formulated to explain the properties of this class of materials.

2. Experimental Procedures

2.1 Preparation of materials

All starting materials were of six 9's purity. The antimony and tellurium were obtained from COMINCO, the sclenium from Norddeutsche Raffinerie. Further purification was carried out on those elements used for preparing the end components of the system, Sb_2 Pe₃ and Sb_2Se_3 . For these, the antimony and tellurium were outgassed and scaled-off under a vacuum, and then zone-refined in a carbon crucible. The sclenium was outgassed and sublimed under vacuo.

Single crystals or large grain size ingots were grown by the Bridgman technique or by a horizontal zone-refiner with background heating, described elsewhere [5]. Specimens for optical mer. mements were cleaved from these ingots down to a thickness of a few tens of microns hy pealing a layer from the surface using Scotch-tape, or by mounting a thick specimen with epoxy onto an aluminum plate possessing a small aperture and then pulling the bulk of the material off the plate, leaving a thin layer behind.

Chemical analyses were performed on sections of these ingots to test for uniformity of composition and approximate correspondence with the nominal composition was obtained.

2.2 Optical measurements

Transmission and reflectivity measurements in the energy range 6.5 eV to 2.0 eV (wavelength 0.18 μ m to 2.6 μ m) were performed in a Cary double-beam 14R spectrophotometer. A Cary attachment, based on a V-W configuration by Strong [6] was used to measure reflectivity at near normal (< 12°) incidence, using an aluminum mirror as a standard reference. This attachment requires two reflecting samples.

A Beekman JR 12 spectrophotometer was used for transmission and reflectivity measurements in the range 2.0 eV to 0.04 eV (2.5 µm to 35 µm). A standard Beekman attachment was used to measure reflectivity at an angle of incidence of $\approx 28^{\circ}$. A long wavelength limit of 35 µm was imposed on the reflectivity measurements by the CsBr₂ lenses in the attachment.

3. Reflectivity

The room-temperature reflectances (R) of the Sb₂Te₃-Sb₂Se₃ solid solutions in the range 0.04 eV to 2.0 eV are shown in Fig. 1. The pronounced minima appear to be associated with the charge carrier plasma frequency; in this case,

Fig. 1. The long-wavelength reflectance spectra of Sb₂Te₃-Sb₂Se₃ solid solutions. The numbers refer to at^o₁₁ Sb₂Te₃



a hole plasma, since all materials were determined to be p-type by sign of the Seebeck coefficient. A calculation of the approximate position of ω_p , the plasma frequency minimum [7] for Sb₂Te₃ from

$$\omega_{\rm p}^2 = \frac{p \, e^2}{m^* \, \varepsilon_0 \, (n_{\rm c}^2 - 1)} = \frac{\sigma \, e}{m^* \, \mu \, \varepsilon_0 \, (n_{\rm c}^2 - 1)} \,, \tag{1}$$

where p charge earrier concentration. e electronic charge, m^* effective mass, ε_c permittivity of free space, σ conductivity, μ mobility, and n_c the refractive index in the non-dispersive region of the spectrum, using values³) of p, m^* , and n_c from Testardi et al. [8] gave a value of 7 μ m which is in reasonable agreement with experiment. Furthermore, the plasma frequency minima for Sb₂Te₃ samples, doped with various amounts of copper. shift with the square root of conductivity giving additional confirmation of this assignment.

The minima for the $Sb_2Te_3-Sb_2Se_3$ solid solutions move toward longer wavelengths with increasing amounts of Sb_2Se_3 . Again, the shift in position correlates surprisingly well with the square root of the conductivity considering no allowance was made for mobility variations.

In Fig. 2 the reflectance values at higher energies are shown and pronounced maxima are observed in the range 1 eV to 6 eV. By analogy with Bi₂Te₃, these maxima are probably due to direct interband transitions which should be associated with critical points at or between the $\Gamma(000)$ and Z(111) points of the Brillouin zone [1]. If this band model is correct, then any line structure in the curves would be due to a spin orbit split valence band. However, this model

³) Published values by various investigators of p (and σ) for undoped Sb₂Te₃ are in close agreement. The approximate value of n_c is obtained from the curve of $n^2 - k^2$ versus λ^2 in the free earrier absorption range by extrapolating back to zero wavelength.

W. PROCARIONE and C. WOOD



has been criticised by Brust [9] on the grounds that the assignment of all peaks to the high symmetry $I^{\prime}Z$ points is too restrictive. Furthermore, Sobolev et al. [2] found no evidence of a triplet structure in their reflectivity data on Bi₂Te₃, Bi₂Se₃ and Sb₂Te₃ and concluded that the reflectivity maxima have no spin-orbit nature. The band model of Borghese and Donato [10] was also too restricted to high symmetry points to satisfactorily explain their data and they suggested that assignments to the $A(D) \langle 100 \rangle$ points should not be exluded if the triplet interpretation is omitted. It appears further calculations are needed to unravel the band structure of this class of materials.

The reflectance data for Sb_sSc_s is shown in Fig. 3. The pronounced minimum in the curve is associated with the closest approach of the valence and conduction bands and corresponds to a direct transition.

4. Absorption

Transmission measurements were obtained only on those compositions containing the higher concentrations of Sb_2Se_3 and are shown in Fig. 4. For the high Sb_3Te_3 concentrations, no measureable transmission was obtained even for samples less than 10 μ m thick.

The absorption coefficient (α) and hence the extinction coefficient ($k = \alpha \lambda/4 \pi$) was calculated from the ratio of transmitted (I) to incident intensity (I_0)

$$\frac{I}{I_0} = \frac{(1-R)^2 e^{-\alpha x}}{1-R^2 e^{-2\alpha x}},$$
(2)

where x is the sample thickness, assuming no multiple interference effects. Calculations showed this equation to be valid even for those cases where inter-

The Optical Properties of Sb₂Se₃-Sb₂Te₃





Fig. 4. The transmission spectrum of Sb₃Te₃-Sb₃Se₃ solid solutions

Fig. 5. The variation of the product of absorption coefficient (a) and refractive index (*) in the free carrier absorption region with the square of the wavelength in Sb_{3}Te_{3}-Sb_{3}Se_{3} solid solutions

ference fringes were observed, leading to errors in α of less than 2% for these samples. Having obtained k, the refractive index (n) was then determined from

$$n = \frac{1+R}{1-R} + \sqrt{\frac{4R}{(1-R)^2} - k^2}.$$
 (3)

Values so obtained agreed very closely with the values of n determined from the positions of the interference fringes. The product $n \alpha$ in the free carrier absorption range is plotted against λ^2 in Fig. 5 and is shown to obey the relation [7]

$$n \alpha = \frac{\lambda^2 e^3 p}{4 \pi^2 c^3 \varepsilon_0 m_p^2 \mu_p}.$$
 (4)

The square root of the absorption coefficient was plotted against energy (Fig. 6) for each composition and found to yield two straight lines conforming to the equation [11]

$$\alpha = A \begin{bmatrix} (h \nu - E_{g} - k \theta)^{2} \\ 1 - \exp\left(-\frac{\theta}{T}\right)^{2} + \frac{(h \nu - E_{g} + k \theta)^{2}}{\exp\left(\frac{\theta}{T}\right) - 1} \end{bmatrix}$$
(5)



Fig. 6. Square root of the absorption coefficient versus incident photon energy

67

W. PROCARIONE and C. WOOD

for the simplest case of indirect allowed transitions, where $k\theta$ is the single phonon energy required to conserve momentum. By analogy with Bi_aTe₃ [1] these indirect transitions should occur somewhere along the ΓA or ΓD directions of the Brillouin zone. As is generally the case for M_0 type critical points no structure in the room-temperature reflectivity euroes corresponding to these indirect transitions was observed [12].

An arbitrary selection of an absorption coefficient value for the absorption edge [1] did not show any regular trend in energy gap with composition. Values of the energy gap (E_g) and $k \theta$ determined from the intercepts of the curves with the abscissa in Fig. 6 are listed in Table 1. The E_g -values show a decreasing trend with increasing Sb_sSe₃ content.

The exact values of these energy gaps are uncertain because of the indication of considerable charge carrier degeneracy from Fermi-level positions determined by Seebeck coefficient measurements. The possibility of the $Sb_2Te_3-Sb_2Se_3$ solid solutions having light and heavy hole valence bands as in Sb_2Te_3 [13, 14], or that both electrons and holes are contributing to the Seebeck coefficient in these low band gap materials, does not allow an accurate designation to be made of the position of the Fermi-level from Scebeck coefficient measurements. Moreover, corrections for degeneracy, as carried out by Sehr and Testardi [15] or by Greenaway and Harbeke [1], did not allow for the fact that optical transitions may take place from levels approximately 4 kT (equivalent to 0.1 eV at room-temperature) away from the Fermi-level [16], i.e. at an energy spread greater than the correction applied. Therefore, no reliable estimate of the Burstein shift [16] can be made.

For pure Sl_2Se_3 the energy gap occurs at about 1 eV and in contrast to the statement of Shutov et al. [3], appears to arise from a direct allowed transition [17]. Excellent correspondence is seen in Fig. 3 between the reflectivity minimum and the point of maximum slope of the absorption edge [7].

Sample No.	Nominal % Sb ₂ Te ₃	Resistivity (Ωem) (300 °K)	Seebeck coeff. (µV/deg) (300 °K)	Fermi level **) (eV)	E _g (eV)	ө (°К)	λ _{min} (μm)
]	100	$2.18 imes 10^{-4}$	90 ± 2	0.07	0.21 *)	400 to 1600*)	8.5
2	90	4.18×10 ⁻⁴	45 ± 3	0.14		· _ ·	10
3	78	-	75 ± 3	0,09		,	11
4	67	$4.27 imes 10^{-4}$	43 ± 4	0.17	0.16	775	13
5	1 60	4.83×10^{-4}	48 + 3	0.14	0.14	660	15
6	50	5.66×10^{-4}	97 + 2	0.06	0.09	660	ⁱ 19
7	41)	8.00×10 ⁻⁴	61 + 4	0.11	0.04	520	23
8	33	11.28×10^{-4}	66 + 2	0.10	0	500	24

Table 1

*) Values obtained from [15].

**) Fermi level values calculated from the Seebeck coefficient assuming a one-carrier single value band model.

5. Discussion

In view of the fact that Sb_2Te_3 is isostructural with Bi_2Te_3 and Bi_2Se_3 , the highest valence band and lowest conduction band would be expected to consist of six ellipsoidal isoenergetic surfaces [2] and the Brillonin zone would be expected to exhibit the same symmetry as $Bi_{3}Te_{3}$ [1]. The similarity in optical properties is therefore not surprising. $Sb_{3}Se_{3}$ differs from $Sb_{3}Te_{3}$ in its crystal structure. Thus, some differences in optical properties would be expected and are found, e.g. the smallest energy gap in $Sb_{3}Se_{3}$ is a direct one, and splitting of several $Sb_{3}Se_{3}$ reflectivity peaks (A_{4.5}) occur due to lower crystal symmetry [3].

Drabble and Goodman [18] have proposed a layer structure for the BiaTea class of compounds, each layer consisting of similar atoms stacked in a sequence Te⁽¹⁾-Bi-Te⁽²⁾-Bi-Te⁽¹⁾ perpendicular to the three-fold axis. The superscripts denote differently bonded tellurium atoms: the Te⁽²⁾ atoms being surrounded almost octohedrally by covelently bonded Bi atoms, with a weak ionic component; the Te⁽¹⁾ atoms have three nearest neighbor covalently bonded Bi atoms, with a strong ionic component, and three van der Waal bonded $Te^{(1)}$ atoms in the adjacent layer. This model was used by Drabble and Goodman to explain the results of Austin and Sheard [19] on the variation of the optical energy gap of Bi₂Te_{3-z}Se_z alloys, and was used later by Greenaway and Harbeke [1] to explain their data on similar alloys. It was proposed that if tellurium is replaced by sclenium then the more electronegative sclenium atoms will substitute preferentially for the more weakly bound Te⁽²⁾ atoms. The increased bond strength will cause an increase in the energy gap. Eventually, when all Te⁽²⁾ sites are occupied, sclenium then substitutes for Te⁽¹⁾ atoms, attracting charge away from the Bi-Se⁽²⁾ bonds, thus causing the energy gap to decrease again.

It appears more likely that selenium is substituting for the $Te^{(1)}$ atoms in the Sb_2Te_3 - Sb_3Se_3 alloys. The general trend of decreasing band gaps and the transition from hexagonal to mixed phase at the composition $Sb_2Te_1Se_2$ lend support to this view. The high hole concentration always observed in Sb_2Te_3 is thought to be due to a wrong-atom defect [20] with a stoichiometric excess of Sb (providing acceptor states) on Te sites. Although it is probable, in view of the above model, that Te vacancies would occur preferentially on $Te^{(2)}$ sites it does not necessarily follow, as Teramoto and Takayanagi [4] supposed, that it is energetically favorable for excess Sb to occupy the $Te^{(2)}$ sites rather than the $Te^{(1)}$ sites.

It appears that more definitive X-ray studies are required to resolve these problems.

References

- [1] D. L. GREENAWAY and G. HABBEKE, J. Phys. Chem. Solids 26, 1585 (1965).
- [2] V. V. SOBOLEV, S. D. SHUTOV, YU. V. POPOV, and S. N. SHESTATSKH, phys. stat. sol. 30, 349 (1968).
- [3] S. D. SHUTOV, V. V. SOBOLEV, YU. V. POPOV, and S. N. SHESTATSKII, phys. stat. sol. 81, K23 (1969).
- [4] I. TERAMOTO and S. TAKAYANAGI, J. Phys. Chem. Solids 19, 124 (1961).
- [5] W. PROCARIONE and C. WOOD, to be published.
- [6] J. STRONG, Procedures in Experimental Physics, Prentice Hall, Inc., Englewood Cliffs (N.J.) 1938 (p. 376).
- [7] T. S. Moss, Optical Properties of Semiconductors, Butterworths, London 1961.
- [8] L. R. TESTARDI, J. BIERLY, and F. DONAHOE, J. Phys. Chem. Solids 23, 1209 (1962).
- [9] D. BRUST, Phys. Rev. 184, 1337 (1964).
- [10] F. BORGHESE and E. DONATO, NUOVO Cimento 53B, 283 (1968).
- [11] G. G. MACFARLANE and V. ROBERTS, Phys. Rev. 97, 1714 (1955).

878 W. PROCABIONE and C. WOOD: The Optical Properties of Sb₂Sc₂-Sb₂Te₂

- [12] D. L. GREENAWAY and G. HABBEKE, Optical Properties and Band Structure, Pergamon Press, London 1968 (p. 41).
- [13] B. RÖNNLUND, O. BECKMAN, and H. LEVY, J. Phys. Chem. Solids 26, 1281 (1965).
 [14] I. A. SMIRNOV, A. A. ANDREEVA, and V. A. KUTASOV, Soviet Phys. Solid State 10,
- 1403 (1968).
- [15] R. SKHR and L. R. TESTABDI, J. Phys. Chem. Solids 23, 1219 (1962).
- [13] E. BUBSTEIN, Phys. Rev. 93, 632 (1954).
 [17] Z. HUBYCH and C. WOOD, to be published.
- [18] J. R. DRABBLE and C. H. GOODMAN, J. Phys. Chem. Solids 5, 142 (1958).
- [19] I. G. AUSTIN and A. SHBARD, J. Electronics and Control 3, 236 (1957).
- [20] T. C. HARMAN, B. PARIS, S. E. MILLER, and H. L. GOERING, J. Phys. Chem. Solids 2, 181 (1957).

(Received August 6, 1970)

and to alterna aller

THERMALLY MODULATED PLASMA FREQUENCY OF FREE HOLES IN THE Sb, Te 3-Se COMPOUNDS*

Z. Hurych and C. Wood,

Physics Department, Northern Illinois University, Dekalb, Illinois 60115

(Received 3 November 1970 by R.H. Silsbee)

The plasma resonance frequency ω_p of free holes in a single-crystal of the narrow-band semiconductor $Sb_2 Te_{3-x}Se_x$ was obtained using thermally-modulated reflectivity. This method gave accurate values of ω_p even in the case of broadening due to strong free-carrier and interband absorption.

SINGLE crystal solid solutions of rhombohedral Sb_2Te_3 and orthorhombic Sb_2Se_3 can be prepared of composition $Sb_2Te_{3-x}Se_x$ for values of x in the range 0-2.¹ All of these compounds possess similar physical properties: rhombohedral $(C_{33}D_{3d}^3)$ crystal structure; p-type conductivity with values of $10^{-4} - 10^{-3} (\Omega \text{ cm})^{-1}$, decreasing with x; similar reflectivity spectra with two sharp reflectivity windows, one in the u.v. region at $\lambda \sim 2,800$ Å for all samples, and the other in the i.r. region at $\lambda \sim 9 \cdot 5\mu$ for Sb_2Te_3 and at longer wavelengths with increasing x.

The long wavelength edges of these i.r. reflectivity windows were associated with the plasma resonance frequency ω_p of free holes in the valence band. For Sb₂ Te₃ this assignment of a plasma window was confirmed' from the position of the minimum reflectivity. However, absorptive processes produce a broadening of the resonance pattern as well as a non-zero reflectivity at the minimum, thus complicating the exact determination of ω_p from the reflectance spectrum. The rather

^{*} This research was supported by the Advanced Research Projects Agency of the Department of Defense and was monitored by the Army Research Office Durham, under Contract No. DA-ARO-D-31-124-70-G77.



FIG. 1a, b: The reflectivity R (dashed curve) and modulated reflectivity $\Delta R/R$ (solid curve) in the vicinity of the plasma resonance frequency for various compositions of Sb₂Te_{3-x}Se_x.

181
	Sb₂T ≈ ₃	Sb ₂ Te ₂ Se	Sb ₂ Te ₁₋₅ Se ₁₋₅	Sb ₂ TeSe ₂
$\omega_p [\text{cm}^{-1}]$				
from thermal modulation	1,040	645	435	39 5
$\omega_2 [\text{cm}^{-1}]$				
from thermal modulation	1,125	665	505	470
ϵ_L	7.3	16.7	4.0	3.70
ε _c				
from extrapolation $\lambda = 0$	50 *	28	42	<u>3</u> 9
η	0.410	0.158	0.860	0.925
$\omega_{\tau} [\mathrm{cm}^{-1}]$				
from η	426	102	375	370
ω_{p} (cm ⁻¹)				
from extrapolation $\epsilon_r = 0$	1,025*	530	490	373

* Data from reference 7.

high concentration of free carriers $(p \sim 10^{19}-10^{19}\text{cm}^{-3})$ affects not only the real part of the dielectric constant ϵ_r , but also the imaginary part ϵ_i through the free-carrier absorption. Moreover, in the case of *p*-type materials, an additional contribution to ϵ_r and to ϵ_i due to interband transition between light and heavy hole bands generally exists, 2^{23} which may be enhanced in the case of spin-orbit splitting of the valence bands. Therefore, thermomodulated reflectivity experiments were performed on the Sb₂Te_{3-x}Se_x compounds in order to obtain accurate values of ω_p and to confirm the plasma oscillation assignment.

All samples could be cleaved easily along natural cleavage planes (0001) thus allowing the preparation of thin single crystal specimens, $20-50\mu$ thick. With this sample orientation the electric vector E of the incident radiation was perpendicular to c-axis throughout our measurements. Square-wave current pulses with peak values of from 500 mA up to 2.0 A at a frequency of 11 c/s were passed through the samples. A modified Beckmann i.r.-12 Spectrophotometer was used, such that the modulated 11 cps signal from the thermopile detector was fed to a Keithley 802 Series Lock-in Amplifier and then recorded.

The thermoreflectance spectra of the samples are shown in Fig. 1 together with the reflectance spectra. All of the samples exhibit the typical feature of a thermally modulated plasma frequency,



FIG. 2. The real part of the dielectric constant, $\epsilon_r = n^2 - k^2$ and the free-carrier absorption $(n \cdot a)$ versus the square of the wavelength (λ^2) for the Sb₂Te_{3-x}Se_x compounds.

i.e. two sharp peaks of opposite sign.⁴ One of the peaks (ω_p) corresponds to the thermal modulation of the plasma frequency ω_p (in these narrow gap semi semiconductors, ω_p is modulated both by change of carrier concentration, p, as well as by change of the effective mass m_c^*), and the other peak (ω_2) corresponds to the modulation of the broadening parameter $\eta = \omega_T / \omega_p$, (where ω_T is the collision frequency which express phenomenologically the scattering of free carriers by phonons). The shorter wavelength side of the thermomodulated spectra shows additional structure which may be

and the second second

72

Table 1.

due to 'ransitions between the light and heavy hole bands.^{2,3} Light and heavy hole hands have been postulated to explain thermoelectric⁵ and Hall effect data⁶ in this class of compounds and appear to indicate a heavy to light hole concentration of at least two orders of magnitude. This large concentration ratio, even allowing for the lower effective mass of light holes, would auggest that a plasma resonance frequency due to the light holes would occur somehwere in the far i.r. region of the spectrum [see equation (2)], and that heavy holes are responsible for the plasma resonance observed.

From the two frequencies ω_p and ω_2 , the non-dispersive dielectric constant of the lattice ϵ_L and the broadening parameter η can be determined using the relations⁴

$$\epsilon_L \quad \left[1 - \left(\frac{\omega_p}{\omega_z}\right)^2\right]^{-1} \quad \text{and} \ R(\omega_2) \quad \left[\frac{\epsilon_L \omega_p^3 \eta}{4 \omega_2^3}\right]^2. \quad (1)$$

These results are shown in Table 1 together with calculated values of ω_{τ} and of the optical dielectric constant ϵ_C ; the latter was determined from the extrapolation of the spectral dependence of $n^2 - k^2$ in the free carrier absorption range to $\lambda = 0$. (Fig. 2; *n* and *k* were obtained from transmission and reflectivity measurements.¹)

Extrapolation of the same curves to $n^2 - k^2 = 0$ gave values of ω_p^* in reasonable agreement with the values of ω_p obtained from thermal modulation while values of ϵ_L , determined from equation (1) were quite different from ϵ_C , which is obviously due to large discrepancy between the wavelength at which minimum reflectivity occurs and the wavelength were $\epsilon_r = n^2 - k^2 = 1$ [ϵ_L has very strong dependence on (ω_p/ω_2)]. A similar discrepancy can be found in case of Sb₂Te₃.⁷ This discrepancy can be due either to interband transitions hetween light and heavy hole bands which contribute to the real, ϵ_r , and imaginary, ϵ_i , parts of the dielectric constant, similar to the case of p-type germanium,³ or to a high collision frequency ω_2 such that the condition $(\omega_p/\omega_r)^2 \gg 1$ is not valid. For either case, the expression:

$$\epsilon = \epsilon_C - \frac{4\pi p}{m_C^* \omega \left(\omega + i \omega_{\tau}\right)}$$
(2)

for the total dielectric constant does not apply, so equation (1) cannot be used without modification. Existence of either of the above phenomena is supported by an absorption background resulting in high values of the free carrier absorption background resulting in high values of the free carrier absorption ($\sim 10^3 \text{ cm}^{-1}$) as extrapolated to $\lambda = 0$. However, the additional structure in the thermomodulated spectra as well as the calculation of ω_{τ} from the Hall modility data⁷ indicate that the effect of interband transitions is more probable. These transitions are therefore also responsible for the discrepancy between ω_p and ω'_p .

The advantage of thermomodulating the reflection spectra is clearly indicated by the fact that despite very strong broadening due to competing absorption processes, ω_p can be determined with high accuracy.

REFERENCES

1. PROCARIONE W. and WOOD C., to be published in Phys. Status Solidi 42, No. 2, (Dec. 1970).

2. FAN H.Y., Semiconductors and Semimetals, Vol. 3, p. 406, Academic Press (1967).

3. SPITZER W.G. and FAN H.Y., Phys. Rev. 106, 882 (1957).

4. CORDONA M., Solid State Physics, Suppl. 11, p. 118, Academic Press, N.Y. (1967).

5. RONNLUND R., BECKMAN O. and LEVY H., J. Phys. Chem. Solids 26, 1281 (1965).

6. SMIRNOV I.A., ANDREEV A.A. and KUTASOV V.A., Sov. Phys. Sol. State 10, 1403 (1968).

7. SEHR R. and TESTARDI L.R., J. Phys. Chem. Solids 23, 1219 (1962).

Die Plasmørøsonanzfrequenz ω_p von freien Löchern in Einkristallen dea Schmalband-Halbleiters Sb₂ Te_{3-x} Se_x wurde mit Hilfe von thermisch modulierten Reflektivitätamessungen bestimmt. Mit dieser Methode erzielt man sogar dann genaue Ergebnisse, wenn Verschmierung durch starke Absorption der freien Träger auftritt oder wenn Interbandabsorption stattfindet.

73