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| | 57-9-36/40 |
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| AUTHOR | Gross, Ye.F., Zakharchenya, B.P., Pavinskiy, P.P. |
| TITLE: | Diamagnetic Exiton Levels and Cyclotron Resonance |
| PERIODICAL | Thurnal Tekhn, Fiz., 1957, Vol. 27, Nr 9, pp. 21// - 2100 (000K) |
| ABSTRACT : | New phenomena are described. Nearer towards the series border, where diamagnetic displacement in the case of a lacking magnet- ic field attains the amount of the distance between the neigh- boring terms of the series, a spectrum, consisting of nearly equidistant lines, was observed at a distance between the lines of H = 29 coolersted of the order of 2 cm ⁻¹ . This striped spec- trum is continued also beyond the series boundary, where, with a lacking magnetic field, (H=O) the through-going spectrum which corresponds to exiton dissociation is located. The farther one penetrates into the shortwave range, the less distinct does the structure of the spectrum become, and the spectral lines ap- proach more closely to one another over a distance of 1,6 cm ⁻¹ . Hereafter their distribution becomes irregular. These lines are observed on the base of the through-going spectrum, where its intensity does not take a monotonous course but shows absorption maxima. The distance between the maxima is reduced as the short- |
| Card $1/2$ | maxima. The distance between the maximum is approached. Thus, the spectrum wave part of the spectrum is approached. Thus, the spectrum |
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| | na se presenta a la constante de la constante d La constante de la constante de |

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57-9-36/40 Diamagnetic Exiton Levels and Cyclotron Resonance here consists of absorption maxima upon which the aforementioned striped spectrum is impressed in form of a thin structure. The intensity of the absorption maxima becomes weaker to the extent as they shift towards the vdolet part of the spectrum, and coalesce with the limit of the continuous absorption. Investigations showed that the through-going exiton spectrum is a superposition of the absorption spectra corresponding to the exiton states at various A-values. A - is the magnetic quantum number of the exiton. There is 1 figure and 2 Slavic references. Physical-Technical Institute AN USSR, Leningrad ASSOCIATION (Fiziko-tekhnicheskiy institut AN SSSR, Leningrad) July 8, 1957 SUBMITTED: Library of Congress AVAILABE: Card 2/2

| Sciences TITLE: An International Symposium on the Hydrogen Bond in Ljubljama (Mezhdunarodnyy simpozium po vodorodnoy svyszi v Lyublyane) PERIODICAL: Vestnik AliSSSR, 1957, Vol. 27. Nr 11, pr. 157-159 (JSSR) ABSTRACT: Scientists from Yugoslavia, the Nest- and Emsteuropean countries, Australia, the USA, Canada, the USSR, Scandinavia and the state of Israel participated in the symposium held from July 29 to August 3. More than 60 speakers got a hearing. The soviet delegation read 6 papers: Ye.F.Gross talked on "The vibration spectrum of the hydrogen bond". D.N.Shigorin on "The nature of the hydrogen bond and its influence upon the vibration- and electron-spectra of the molecules", V.M.Chulanovskiy on "The spectroscopic investigation of the hydrogen bond". K.V.Vol'kentroscopic investigation of the hydrogen bond in vitrificable on the vibration theory of the hydrogen bond "A.N.Terenin and V.Filiaonov "The hydrogen bond" between edsorbed molecules and the structural OH-groups on the surface of solid bodies". Kany papers were devoted to the surface of solid bodies". Kany papers were devoted to the surface of solid bodies". | | GROS'S | Ye. F. | |
|--|---|-------------|--|--|
| TITLE: An International Symposium on the hydrogen point of the hydrogen point. PERIODICAL: Vestnik AlESSR, 1957, Vol. 27. Nr 11, p. 137-139 (JSSR) ABSTRACT: Scientists from Yugoslavia, the West- and Lasteuropean countries, Australia, the USA, Canada, the USSR, Scandinavia and the state of Israel participated in the symposium held from July 29 to August 3. More than 60 speakers got a hearing. The soviet delegation read 6 papers: Ye.F.Gross talked on "The vibration spectrum of the hydrogen bond". D.N.Shigorin on "The nature of the hydrogen bond and its influence upon the vibration- and electrometroscopic investigation of the hydrogen bond", M.V.Vol'kentroscopic investigation of the hydroge: bonds in vitrificashteyn on "The behavior of the hydroge: bonds in vitrificability of the hydrogen bond". A.N.Terenin and V.Filimonov "The hydrogen bond between edsorbed molecules and the structural OH-groups on the surface of solid bodies". Many papers were devoted to the | • | AUTHOR: | Voltkenshtern, M. V., Loctor of Physics-Asth matical 30-11-22/23 Sciences | |
| PERIODICAL: Vestnik ALSSSR, 1957, Vol. 27. Nr 11, p. 137-139 (JSSR) ABSTRACT: Scientists from Yugoslavia, the West- and Easteuropean countries, Australia, the USA, Canada, the USSR, Scandinavia and the state of Israel participated in the symposium held from July 29 to Au- gust 3. More than 60 speakers got a hearing. The soviet delega- tion read 6 papers: Ye.F.Gross talked on "The vibration spectrum of the hydrogen bond". D.N.Shigorin on "The nature of the hy- drogen bond and its influence upon the vibration- and elec- tron-spectra of the molecules", V.M.Chulanovskij on "The spec- troscopic investigation of the hydrogen bond", M.Y.Vol'ken- tion (steklovanije)", N.D.Sokolov "On the quantum theory of the hydrogen bond Between edsorbed molecules and the structural OH-groups on the surface of solid bodies". Many papers were devoted to the | | TITLE: | An International Symposium on the hjurogen Luni v Lyublyane) (Mezhdunarodnyy simpozium po vodorodnoy svyszi v Lyublyane) | |
| Australia, the USA, Canada, Chinada, Chinada, Chinada, The Symposium held from July 29 to Au- of Israel participated in the symposium held from July 29 to Au- gust 3. More than 60 speakers got a hearing. The soviet delega- tion read 6 papers: Ye.F.Gross talked on "The vibration spectrum of the hydrogen bond". D.N.Shigorin on "The nature of the hy- drogen bond and its influence upon the vibration- and elec- tron-spectra of the molecules", V.M.Chulanovskij on "The spec- tron-spectra of the molecules", V.M.Chulanovskij on "The spec- troscopic investigation of the hydrogen bond", K.Y.Vol'ken- troscopic investigation of the hydroge: bonds in vitrifica- shteyn on "The behavior of the hydroge: bonds in vitrifica- tion (steklovanije)", N.D.Sokolov "On the quantum theory of the hydrogen bond" A.N.Terenin and V.Filimonov "The hydrogen bond between edsorbed molecules and the structural OH-groups on the surface of solid bodies". Many papers were devoted to the | | PERIODICAL: | Vestnik AlbSSR, 1957, Vol. 27. Nr 11, p. 137-139 (USSR) | |
| | | | Australia, the OSA, Chinada, Chinada, Shinada, S | |
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An International Symposium on the Hjurogen Bond

in Ljubljana. 30-11-22/23

given by Dah.Pimentel (USA) on the spectral properties of the compounds at low temperatures and G.Marrinan (England) on the investig tion-results of the orgstalline modifications of cellulose by means of the method of polarized infrared spectra. E. Lippert (German Federal Republic) gave an extensive survey of the upon the electroninfluence exerted by the hydrogen bonds spectra. The session in which the participants especially dealt with the problems of the crystallography of the compounds with those of hydrogen, was opened by Dzh.Bernal, England, with an extensive report on the part played by the hydrogen bonds in solids and in liquids for which the participants showed great interest. R.Popinskiy (USA) talked on the investigation of the by means of the X-ray and neutronographic mehjdrogen bond thod. U.Shneyder (Canada) and others also dealt with this method. The following sessions mainly dealt with problems of the theory Speaker was: Ch.Koulson, England. His of the hydrogen bond. statement caused a lively discussion in which above all the American scientists participated. Although there exists no strict definition on the conception of the hydrogen bond , , all participants in the discussion agreed that the evidence of the quantum-mechanical process of the formation of a donor-acceptor (donorno-aktseptornaja svjaz') were necessary for the bond

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| AUTHOR:Gross, Ye.P.TIFLE:The Spectrum of the Excitation of Exitons in a Crystal Lattice (Spektr vozbuzhdeniya eksitonov v kristallicheskoy reshetke)PERIODICAL:Uspekhi Fiz. Nauk , 1957, Vol. 63, Er 3, pp. 575 - 611 (USSR)ABSTRACT:The present survey is arranged as follows: Introduction, investi- gation of the structure of the long-wave edge of the principal absorption of a ciprous oxide crystal at room temperature and at the temperature T = 77.3 K (liquid nitrogen). The hydrogen- like series of narrow absorption lines. The optical spectrum of the excitation of exitons in a crystal lattice. The yellow exiton series in a Cu_O-crystal. Comparison of the exiton spec- trum with that of the hydrogen atom. Pree and polarized exitons. The green exiton series in a Cu_O crystal at the temperature of liquid nitrogen (77.3 K) and at the temperature of liquid helium (T = 4.2 K). (The wave length of the newly found absorp- tion lines of the yellow and green exiton series in Cu_O are given in a table). The exiton spectrum in Cu_O at T = 1.3 K (liquid helium). The Stark effect at the lines of the exiton field).Card 1/2 | 4 ********** | Gross, Ye.F. |
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| (Spektr vozbuzhdeniya eksitonov v kristarrinkenský felder v PERIODICAL: Uspekhi Fiz. Nauk , 1957, Vol. 63, Nr 3, pp. 575 - 611 (USSR) ABSTRACT: The present survey is arranged as follows: Introduction, investigation of the structure of the long-wave edge of the principal absorption of a coprous oxide crystal at room temperature and absorption of a coprous oxide crystal at room temperature and absorption of a coprous oxide crystal at room temperature and the temperature T = 77.3 K (liquid nitrogen). The hydrogen-like series of narrow absorption lines. The optical spectrum of the excitation of exitons in a crystal lattice. The yellow exiton series in a Cu₂O-crystal. Comparison of the exiton spectrum with that of the hydrogen atom. Free and polarized exitons. The green exiton series in a Cu₂O crystal at the temperature of liquid nitrogen (77.3 K) and at the temperature of liquid helium (T = 4.2 K). (The wave length of the newly found absorption lines of the yellow and green exiton series in Cu₂O are given in a table). The exiton spectrum in Cu₂O at T = 1.3 K (liquid helium). The Stark effect at the lines of the exiton | AUTHON: | |
| ABSTRACT: The present survey is arranged as follows: Introduction, investi- gation of the structure of the long-wave edge of the principal absorption of a ciprous oxide crystal at room temperature and at the temperature $T = 77.3$ K (liquid nitrogen). The hydrogen- like series of narrow absorption lines. The optical spectrum of the excitation of exitons in a crystal lattice. The yellow exiton series in a Cu ₃ O-crystal. Comparison of the exiton spec- trum with that of the hydrogen atom. Free and polarized exitons. The green exiton series in a Cu ₂ O crystal at the temperature of liquid nitrogen (77.3) K) and at the temperature of liquid helium $(T = 4.2)$ K). (The wave length of the newly found absorp- tion lines of the yellow and green exiton series in Cu ₂ O are given in a table). The exiton spectrum in Cu ₂ O at $T = 1.3$ K (liquid helium). The Stark effect at the lines of the exiton | TI TLE: | The Spectrum of the Excitation of Exitons in a Crystal Lattice (Spektr vozbuzhdeniya eksitonov v kristallicheskoy reshetke) |
| gation of the structure of the long-wave case of comparature and absorption of a coprous oxide crystal at room temperature and at the temperature $T = 77.3$ K (liquid nitrogen). The hydrogen- like series of narrow absorption lines. The optical spectrum of the excitation of exitons in a crystal lattice. The yellow exiton series in a Cu ₂ O-crystal. Comparison of the exiton spec- trum with that of the hydrogen atom. Free and polarized exitons. The green exiton series in a Cu ₂ O crystal at the temperature of liquid nitrogen (77.3 K) and at the temperature of liquid helium ($T = 4.2$ K). (The wave length of the newly found absorp- tion lines of the yellow and green exiton series in Cu ₂ O are given in a table). The exiton spectrum in Cu ₂ O at $T = 1.3$ K (liquid helium). The Stark effect at the lines of the exiton | PERIODICAL: | |
| Card 1/2 aportrum (in a homogeneous and in an inhomogeneous error | | gation of the structure of the long-wive cupt of the provide the second temperature and absorption of a cuprous oxide crystal at room temperature and at the temperature $T = 77.3$ K (liquid nitrogen). The hydrogen- like series of narrow absorption lines. The optical spectrum of the excitation of exitons in a crystal lattice. The yellow exiton series in a Cu ₂ O-crystal. Comparison of the exiton spectrum with that of the hydrogen atom. Free and polarized exitons. The green exiton series in a Cu ₂ O crystal at the temperature of liquid nitrogen (77.3 K) and at the temperature of liquid hybrid comparison of the newly found absorption lines of the yellow and green exiton series in Cu ₂ O are given in a table). The exiton spectrum in Cu ₂ O at $T = 1.3$ K |
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53-3-5/6 The Spectrum of the Excitation of Exiton in a Crystal Lattice The Zeeman effect on the lines of the yellow exiton series. The investigation of the structure of the principal absorption in other crystals. (Experimental data are available only for some crystals). Thus, a complicated structure consisting of a group of 11 narrow and 4 broader bands was observed in the spectrum of cadmium sulphide at 4,2 K at the absorption edge. Besides, a considerable distortion of lines was observed in the case of some CdS crystals. Also in HgJ_-crystals a structure of the ab-sorption edge was observed. In PbJ_-crystals a group of 4 narrow lines was observed on the absorption edge of this crystal at T = 4,2°K. Next, some observations made in the case of other crystals are enumerated. There are 21 figures, 8 tables and 58 references, 33 of which are Slavic. Library of Congress AVAILABLE: Card 2/2

| | Ver All |
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| AUTHORS : | Gross, Ye. F. Corresponding Member AN USSA 20-4-15/60 Shullin, A. A. |
| TITLE: | The Interaction Between Intermolecular and Lattice Vibrations in Crystals According to the Data of Infrared Spectra(Vzaimodeystviye vnutrimolekulyarnykh i reshetochnykh kolebaniy kristallov po dannym infraktasnykh spektrov). |
| -PERIODICAL: | Doklady Akad.Nauk SSSR, 1957, Vol. 115, Nr 4, pp. 689-692 (USSR) |
| ABSTRACT : | At first reference is made to the present state of the problem and to some earlier papers. The problem of the interaction of phonons with the innermolecular vibrations is of great interest. Therefore the author examines the infrared absorption spectra of monocrystalline samples of $Ba(NO_3)_2$ and $Pb(NO_3)_2$. It was the pur- pose of these investigations to clear up the problem of the exi- stence and the peculiarities of the "compound" transitions (who- se intensities are markedly dependent on temperature). The prob- ability of such transitions must also depend on the type of the inner-ionic excitation and on the type of lattice vibrations. In the crystals selected here the composed NO_3^- - ions play the part of structural units which have internal degrees of freedom. The- se ions are plane equilaterial triangles having the nitrogen atom in the center. Such a system has 6 normal vibrations with frequ- |
| Card 1/3 | encies in the range 700- 1500 cm-1. The samples of lead nitrate |

The Interaction Between Intermolecular and Lattice Vibrations in 20-4-15/60 Crystals According to the Data of Infrared Spectra.

> and barium nitrate consisted of plates 0,3ox1ox2omm or 0,56x1ox x20 mm in dimension respectively. The spectra obtained at -160° and $+ 20^{\circ}$ are illustrated by diagrams. The analysis of the selection rule by the group theory leads in crystals belonging to the space group T_h^6 to the following conclusions: All inner group vibrations must be effective in the absorption and this is also observed in reality. But in addition to that, much weaker absorption bands are also noticed in the spectra. The clearly expressed absorption maxima that lie symmetrically to the absorption peak might furnish a contribution toward their explanation. These concomitant bands might be caused by compound transtions in which the quantum numbers of the inner-ionic and the lattice levels simultaneously change. That means that, when a light quantum is absorbed in these transitions, an exiton is produced and at the same time a phonon is created or disappears. Most of these weak bands can be explained by the scheme given here. There are 4 tables and 19 references, 3 of which are Slavic.

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The Interaction Between Intermolecular and Lutice Vibraticus in 2024-15/60 Crystals According to the Data of Infrared Spectra. ASSOCIATION: Leningrad State University imeni A.A.Zhdanov (Leningradskiy gosudarstvennyy universitet imeni A.A.Zhdanova) SUBMITTED: March 25, 1957 AVAILABLE: Library of Congress. Card 3/3



KENERAL SALE

| AUTHOR: | Gross, Ye. F., Corresponding Member, AS USSR |
|-------------|---|
| T1TLX: | An Important Problem of Modern Physics (Vaznnaya problema sovremennoy fiziki` Excitons in the Crystal Lattice (Eksitony v kristallicheskoy reshetke) |
| PERIODICAL: | Vestnik Akademii nauk 3358, 1958, Nr 10, pp 11-19 (USSR) |
| BSTRACT : | The concept o excitons was introduced into physics in 1931 by the Soviet scientist Ta. I. Frenkei'. Exitons are supposed to be cauable of migrating on a crystal. In theory the idea of Frenkel' was developed further by the foreign scientists - Vamier (1937) and N. F. Mott (1938) among others. According to M''., exitons are hydrogen-like quasi-atoms which move on crystals, as well as neutral forms. In the opticheskaya laboratoriya Fiziko-tekhnicheskogo instituta Akademii nauk SSUP (Uptical Laboratory of the Physico-Technical Institute. AS HSSR) some phenomena were recently discovered which prove the exist- ence of excitons in the crystal lattice (N. A. Karryyev, B. P. Zakharchen', A. A. Kaplyanskiy, B. S. Hazbirin, M. A. Yakobson, V. V. Sobolev, and others). The pattern of exciton movement (m |
| 2 ard 1/2 | the crystal lattice is represented in figure 1. Figure 2 snows |

CIA-RDP86-00513R00051703

SOV/30-58-10-2/55 An Important Problem of Modern Physics. Excitons in the Crystal Lattice

> the spectrogram of a hydrogen-like series in the crystal of cuprous oxide at a temperature of $T \sim 1.5~K.$ As the effects of Shtark and Zeeman observed in the exiton spectrum prove, excitons are huge forms in the crystal. Mention was made of the studies by A. G. Samoylovich and L. L. Korenblit (1955). Two excitons of small and large radius can exist in one crystal. In figure 3 patterns of para- and ortho excitons are shown. Recently 1. **Descriti** (Ref 2) reported on the **exciton** level in his theoretical study. 1. M. Dykman, S. I. Pekar, 1. 1. Ansel'm. L. T. Korovin. and I. P. Ipatova also work on this problem. Figure 4 gives a diagram of the origin of resonance luminescence and figure y of inducing luminescence of excitons Apart from electrons, holes and phonons, excitons play an important role in many physical phenomena in the crystal lattice. As the studies of A. V. and A. F. Ioffe (1956) show, excitons play an important part in the thermal conductivity of crystals. Research of excitons and their role in the crystal lattice is still in its beginning stages. There are 5 figures.

Card 2/2

CARLES TRANS

| AUTHCRS: | Gross, 10. F., Zakharchenya, E. P. | 57-2-2/32 |
|-------------|---|--|
| TITLE: | Ionization of Excitons in a Cu ₂ C Crystal by a siya eksitonov v kristalle Cu ₂ O elektricheski | an Electric Field (Ionizat= im polem). |
| PERIODICAL: | Zhurnal Tekhnicheskoy Fiziki, 1958, Vol. 28, | |
| ABSTRACT: | Reference is made to the great difference bet the latter proceeding from the correct assum cuproud oxide is a Motta exciton. As this di- mined, the Stark effect was again investigate first of all the test conditions were perfec- showed a good agreement with theory. First t mer tests are enumerated and it is shown that in the determinations of the field voltages appearance of the members of the exciton-ser be removed when the observations in the doma cut by measurement of the electric potential probes. The probes were put onto a small any of evaporation of gold is a vacuum. The prob- from the silver base electrodes. The Stark e | ption that the exclusion in fference was still undeter= ed in a Cu ₂ C crystal, where ted. The results obtained he deficiencies of the for= t all these sources of error in which a successive dis= ies takes place may easily in investigated are carried . tradient with the aid of stalline Cu ₂ O plate by means les had a distance of 1 ± 2 mon ffect was investigated at |
| Card $1/2$ | the exciton lines under conditions of coolin | g of the crystal to the tem |

| ASSOCIATION. Technical Physics Institute ASUSSR. Leningrad (Fiziko-tekhnicheskiy institut AN SSSR. Leningrad). SUBMITTED. August 22, 1957. AVAILABLE. Library of Congress. Card 2/2 I. Crystals-Excitation 2. Crystals-Ionization | | perature of liquid nitrogen. A spectrograph with a dispersion of log5 A/mm was used for the observation of the spectrum. The consecutive dis- appearance of the members in the yellow exciton-deries with the quantum numbers $n = 4,3,2$ due to the ionization of the exciton by the electric field was distinctly observed. It became evident that a field voltage of 2,5 kV/cm is notessary for the ionization of the exciton from the state with $n = 4$. In the case of $n = 3$, $E = 9$ kV/cm and in the case of n = 2, $E = 29$ kV/cm. The values for the field voltages are highly dif= ferent from those measured earlier and the near those obtained by Samoy= lovich and Korenblit for the Stark effect. I. 4. Polovnikova, Diplomant= ka in the State University, Leningrad, helped in the experiment. There are 5 references, 4 of which are Slavic. |
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| AVAILABLE. Library of Congress. | ASSOCIATION. | Technical Physics Institute ASUSSR. Leningrad (Fiziko-tekhnicheskiy institut AN SSSR. Leningrad). |
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| Card 2/2 1. Crystals-Excitation 2. Crystals-Ionization | AVAILABLE. | Library of Congress. |
| | Card 2/2 | 1. Crystals-Excitation 2. Crystals-Ionization |

| AUTHORS: | SK285, 718, F., Gross, Ye. F., Razbirin, B. S. | 57-2-4/32 |
|-------------|--|---|
| TITLE: | The Influence of Deformations on the Spectrum deformatsiy na spektr kristallov CdS). | n of CdS Crystals (Vliyaniye |
| PERIODICAL: | Zhurnal Tekhnicheskoy Fiziki, 1958, Vol. 28, | Nr 2, pp. 237-239 (USSR). |
| ABSTRACT: | Tests were made here to find out whether the pasting on of crystals are responsible for the absorption-spectrum of CdS-crystals. The long one end pasted to a small glass-plate. The of did not rest against the surface of the glass sible to compare the absorption-spectra of the and of the free part at the same time in one wed that at $T = h_2^{OK}$ as well the narrow line bands N h860-h800 Å, in the spectrum of the p glass are narrowed to almost half of their for by 8-9 Å toward the short-wave side (in compa- the free part of the crystal). In this connec- trum of the thin lines changes in a complica- them becomes smaller, the intensity is redis- lines is diminished and some of them are depart | the distortion of lines in the thered CdS crystals was with ther half remained free and a. In this manner it was post the part of crystal pasted on spectrogram. The tests shot es AA4889-4860 Å, as the part of crystal pasted to the ormer width and are displaced arison to their position in ction the shape of the spect ted way, the distance between tributed, the blurring of the |
| Card $1/3$ | - lines is diminished and some of them are deb | OTSLINGT, THE GOSOLPGION TH |

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57-2-1./32

The Influence of Deformations on the Spectrum of CdS-Crystals.

the lines $\lambda\lambda$ 4857 Å and 4869 Å is intensified. The part of the crystal between the two halves (the half pasted on and the free half) yields a distorted image of spectrum with a continuous transition from the spec= trum of the one half (that pasted on) to the other one (the free one). Analogous distortions were observed after putting drops of glue upon the crystal. It is shown that the quantity and the direction for the displace ment in the spectrum depend on the quantity and the sign of the force deforming the crystal. This becomes evident from a comparison concerning the behaviour of the absorption-spectra of CdS crystals which are glued on glass- and quartz-bases. The spectrum of a crystal-part glued on quartz is displaced to the long-wave-side (by 1 R) (in comparison to the crystal-part not glued on) and simultaneously with the displacement the absorption-lines become wider. This displacement to the long-wave side is connected with the expansion of the crystal by its base (on cooling quartz contracts leas than CdS). It is pointed out that the posting on of the CdS-crystals onto glass and quartz at T = 4,2% is recording to its effect equivalent to an additional cooling and heating of them. The phenomena described here show a high sensitivity of the spectrum of CdScrystals to deformation. There are 3 figures, and 4 Slavic references.

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CIA-RDP86-00513R00051703

| | 57-28-4-17/39 |
|-------------|---|
| THQR5: | Gross, Ye. F., Novikov, B. V. |
| ITLE : | On the Problem of the Structure of a Spectral Curve of the Internal Photoeffect in Cd_S-Crystals (K voprosu o strukture spektral noy krivoy vnutrennego fotoeffekta v kristallakh CdS) |
| PERIODICAL: | Zhurnal Tekhnicheskoy Fiziki, 1958-Vol.29, Nr 4, pp.782-783 (USSR) |
| ABSTRACT: | This is a letter to the editor. In connection with Ref 3 the investigation of the spectral characteristics of photo- conductivity in various CdS-moncrystals at $T = 77.3$ K was continued. These experiments showed that the shape of the spectral curves essentially depends on the orientation of the crystals with regard to the direction of the incident light beam and on the state of its polarization. The occur- rence of the absorption lines in the form of peaks or depressions in the photoeffect-curves however, is mainly not determined by these conditions (Ref 1). but depends |
| Card 1/3 | on the type of crystal. The crystals are according to the |

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57-20-4-17/39 On the Problem of the Strucutre of a Spectral Curve of the Internal Photoeffect in CdS-Crystals

nature of the spectral curves of photoconductivity divided into two groups. 1) At T = 77 3°K the peaks of the curves of photoconductivity correspond to the absorption lines λ 4870 Å and λ 4840 Å. The peak λ 4870 Å is polarized with the electric vector ELC, also the line corresponding to it. In the declineof the short wave portion of the curve of photoconductivity wide maxima are observed at λ 4795 Å and λ 4730 Å. 2) The minima at the photocurrent-curve here correspond to the same absorption lines. Here, too, the polarization of the spectral curves agrees with the polarization of the absorption curves. In the short-wave part . of the photocurrent curve a wide minimum is here observed at 7 4795 A. - A more intensive infrared illumination exerts an extinguishing influence upon the photocurrent in crystals of both groups. It especially strongly influences the longwave part of the photocurrent-curve in crystals of the second group. A detailed description is given. There are 4 references, 4 of which are Soviet.

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| AUTHORS : | 57-28 4-26/39 Gross, Ye. F., Abolin'sh, Ya. Ya., Shultin, A. A. |
|-------------|---|
| TITLE: | On the Observation of the Optico Acoustic Effect in a Liquid (O nablyudenii optiko-akusticheskogo effekta v zhidkosti) |
| PERIODICAL: | Zhurnal Tekhnicheskoy Fiziki, 1958, Vol. 28, Nr 4, pp. 832-835 (USSR) |
| ABSTRACT: | The authors here started from the idea whether it was not possible to utilize the optico-acoustic phenomenon for the determination of the duration of existing states of exci- tation in the molecules of liquids and solids and to work out a method of investigation for condensed systems on the basis of this phenomenon. From these considerations ex- periments were performed in the authors' laboratory. In these experiments the optico-acoustic phenomena in liquids and solids were observed. In the course of these experi- ments in the year 1952, which were repeated in 1957 such phenomena were observed in water, methyl alcohol and ethyl |
| Card 1/2 | alcohol. A perceptible signal was only obtained in a small |

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ŕ 57-28-4-26/39 On the Observation of the Optico-Acoustic Effect in a Liquid range of the modulation frequencies at about 200 cycles. The optimum modulation frequency at which the acoustic signal attains its maximum value depends on the geometric dimensions of the chamber and on the frequency-characteristic of the microphone whose membrane touches the liquid. The spectral dependence of the optico-acoustic signal was observed in liquid ethyl alcohol. The signal was observed in the domain of from $\lambda = 0.95 \,\mu$ to $4 \,\mu$ where the maximum amplitude was attained at $\lambda = 2 \mu \tilde{\mu}$. A comparison of the experiments with analogous ones in which a gas had been investigated shows that the acoustic signal forming in liquids is many times weaker than that forming in gases. By a certain perfection of the scheme it will be possible to use the principle of the gas analyzer by Veyngerov also for an analysis of liquids. At present the experiments for the observation of optico-acoustic phenomena in crystals are continued. There are 3 figures, and 7 references 5 of which are Soviet. Card 2/ Langrad Stale U. 2_

| 24 (6) 30790R - : | 00V-57-23-30-22:40 Gross, 'e, F., Spolin'sh, fa. fa., Shultin, A. A. |
|---------------------------|---|
| TITLS: | Optical-,coustic Effect in Crystals (Optiko-akusticheskiy eifekt v kristallakh) |
| PSRIODIO L: | Zhurnal tekhnicheskoy fiziki Nol 28, Nr 10. pp 2255-2258 (USER) AJS |
| ⁻ B* 甲式 - UT : | The experiments, of which this paper gives an account, for the determination of the optical-account effect were performed according to a scheme died in work with fluid substances (Ref 1). These experiments substantiated the existence of such an effect in crystal. It appears from the information gained that a utilization of this effect in the investigation of the solid state of a substance is dependent on whether ways and means are found of improving the experimental technique and the instrument sensitivity. This paper also covers experiments on piezoelectric properties, a Rochelle-salt crystal serving as a sample. The oscillograms obtained demonstrate that is ontical-accoustic effect in Rochelle-salt crystals is very intensive and comparable to the optic-accoustic effect in gases. It was found that |
| Ctard 1/2 | crystal, which is not clamped down. The intendity in both cases, |

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Optical-'contic mirect in Crystals

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the clamped down and the free one, is the same. In explanation of this effect is advanced and it is consumed that the effect in both cases is due to the same causes. It was also found that the intensity of the optical-acoustic effect gradually decreases after connecting the light source. This is considered to be due to a general increase in temperature of the angle crystal and the gradual approach of the upper Curie point. The opticalacountic offect in the crystal of Rochelle-salt is so intensive that it can be used for the solution of a number of problems. The experiments described in this paper are at present continued by investigating the spectral distribution of the optical-acoustic effect in Rochelle-salt crystals and in other ferroelectric substances. The experiments presented in this paper are only the first stage of investigations of the opticalacoustic effect in crystals. There are 5 figures and / references. b of which are Soviet.

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| 462++0 | S/058/62/000/005/047/119 A001/A101 |
| AUTHORS: | Gross, Ye. F., Sobolev, V. V. |
| TITLE: | Investigation of the structure of absorption, emission and photo- electric effect at the edge of CdSe crystal fundamental absorption (Theses) |
| PERIODICAL: | Referativnyy zhurnal, Fizika. no. 5, 1962, 33. abstract 5V227 (V sb. "Fotoelektr. i optich. yavleniya v poluprovodnikakh", Kiyev, AN USSR, 1959, 40-42) |
| in the spect spectra are emission spectrongly in | A fine structure is discovered at low temperatures, most complicated a absorption and emission spectra of CdSe single crystals, as well as tral distribution of internal photoeffect. Absorption and emission strongly polarized. Position of lines and bands in absorption and ectra is constant for specimens being in free state, but varies very dependence upon strains and stresses in the specimen. Conclusions in the observed bands in CdSe absorption spectra. |
| [Abstracter | 's note: Complete translation] |
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| 26 242 0 | S/ AC | /058/62/000/005/044/119 001/A101 |
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| AUTHORS: | Gross, Ye. F., Razbirin, B. S. | |
| ritle : | Investigation of the structure of main al crystals (Theses) | bsorption edge in CdS |
| PERIODICAL: | Referativnyy zhurnal, Fizika, no. 5, 1963 (V sb. "Fotoelektr. i optich. yavleniya Kiyev, AN USSR, 1959, 61-62) | 2, 32, abstract 5V223 v poluprovodnikakh", |
| 4,889 A was This variable and dependent their ascrib these lines | A strong variability of a group of weak dge of fundamental absorption in the regi discovered in the absorption spectrum of lity, independence of absorption value of dec of the lines on the state of crystal s bing to the CdS principal lattice; they in with the crystal surface. Contrary to th orter wavelength than the narrow ones, are brigin, a high absorption coefficient and | CdS crystals at 4.2 [°] K. the crystal thickness surface do not warrant dicate a connection of lese lines, strong a permanent. Their |
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| · | \$/058/62/000/004/105/160 A061/A101 | |
| AUTHORS: | Gross, Ye. F., Kaplyanskiy, A. A., Novikov, B. V. | |
| TITLE: | Structure of photoconductivity spectral curves in crystals at low temp-ratures (Theses) | |
| PERIODICAL: | Referativnyy zhurnal, Fizika, no. 4, 1962, 39, abstract 4E342 (V sb. "Fotoelektr. i optich. yavleniya v poluprovodnikakh", Kiyev, AN USSR, 1959. 66-73) | i |
| single cryst to correspon of the absor chiefly depe cal vector E cient in the | The shape of photoconductivity spectral curves in crystals with corption edge structure at 77.3 K was studied on CdS. HgI_2 , and PbI_2 cals. For these crystals, the maxima of photoconductivity were found ad to the discrete absorption lines ascribed to excitons. The shape option spectra and of the photoconductivity curves in CdS and HgI_2 ends on the mitual orientation of the crystal axis c and the electri- i of the exciting light. In HgI_2 crystals, when the absorption coeffi- e lines is large, a self-reversal of the maxima of photoconductivity which is related to the increased annihilation of excitons near the ne constant infrared illumination reduces photoconductivity; however, | |
| Card 1/2 | | |
| | | |



GROSS, Te.F.; PASTRNYAK, I. Fine structure of the spectral photosensitivity curve of cuprous oxide crystals. Fis.tver.tela 1 no.1:162-166 Ja '59. (MIRA 12:4) 1. Leningradskiy fisiko-tekhnicheskiy institut. (Copper oxide crystals) (Photoelecticity)

GROSS, Ye.F.; NOVIKOV, B.V.

Fine structure of the spectrum curves of photoconductivity in cadmium sulfide crystals, Fiz. tver. tela 1 no.3:357-362 Mr '59. (MIRA 12:5)

1. Leningradskiy gesudarstvennyy universitet, Fizicheskiy institut. (Cadmium sulfide crystals) (Photoelectricity)

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GROSS, Ye.F.; PASTRNYAK, I.

Formation of polarons in crystals during the absorption of light. Fis. twor. tola 1 no.3:518-521 Mr 159. (MIRA 12:5)

1. Leningradskiy fisike-tekhnicheskiy institut AN SSSR. (Excitens) (Crystal lattices)

GROSS, Ye.F.; PASTRNYAK, I.

Effect of surface treatment on the structure of the curve of spectral distribution of photoconductivity of cuprous oxide crystals. Fiz.tver. tela 1 no.5:837-840 My '59. (MIRA 12:4)

1. Leningradskiy fiziko-tekhnicheskiy institut AN SSSR. (Photoconductivity) (Copper oxide orystals)

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"APPROVED FOR RELEASE: Thursday, July 27, 2000

GROSS, Ye.F.; FASTRHYAK, I. Optical spectrum of the formation of polarons in cuprous oxide crystals. Fiz. tver. tela 1 no.6:973-976 Je '59. (MIRA 12:10) 1.Leningradskiy fisiko-tekhnicheskiy institut AN SSSR. (Copper oxides--Spectra)
CIA-RDP86-00513R00051703

SOV/51-6-1-24/30 AUTHORS : Grozs, St.F. and Susline, L.G. TITLE: Single figurate of ZeS soi the Spectrum of Their Absorption Edge at Low Temperatures (Monokristally ZaS 1 spektr ikh kraya pogloshcheniya pri nizkikh temperaturakh) FERIODICAL: Optika i Spektroskopiya, 1959, Vol 6, Nr 1, pp 115-117 (USSR) ABS TRACT : In single systems were protected by ablimation of very pure powder in an atmosphere of argon. This method is similar to that employed for growing of CdS monocrystals in the laboratory of S.M. Ryvkin. at the Physico-Technical Institute of the Loadeny of Sciences of the U.S.S.R. Sublimation was carried out in a quartz tube (1, in Fig 1) placed in an electric furnace (2, in Fig 1). A quartz boat 3 filled with ZnS was placed in the heuset part of the tube 1. This boat was heated to 1270°C. Before the electric furnace was switched on, a stream of argon was passed for 1 hour via a small tube 4. This was done to expel all air from the system. The electric furnace was switched on for 2 hours. During this time the vapours of ZrS produced from the powder were moved by a stream of argon to a quarts screen 5 ZoB single crystals were found to grow on this screen and on the tube walls near it. The single armir.u were in the form of colourless plates of 25 x 5 mm area Card 1/3

CIA-RDP86-00513R00051703

SOV/51-6-1-24/30 Single Gystals of ZoS and the Spectrum of Their Absorption Suge 25 Low "or paratures and their this meas varied from 0.1 mm to 0.1 µ. The size crystals were found to be uniaxial (hexagonal modification of 2nS) and the optical axis was found to lie in the plane of the crystal. Using crystals of various thicknesses the authors studied the absorption spectra in the region of 3300-5100 Å (Fig 2). Curves a b, 4, 5 in Fig 2 represent spectra obtained using crystals of 1C, 1, 1 and J.1 u thickness respectively. The structure of the long-mivelength edge of absorption was found to be clearly exhibited at the liquid-helium temperature (4.2°K), as shown in Fig 2. Heating of crystals from 4.2 to 77°K and then to 2000 displaces the absorption lines towards longer wavelength: and makes them less distinct. Positions of the absorption lines near the long-mavelength edge are given in various temperatures in a table on p 11". This table lists three absorption maxima at 4.2°K and 77°K and only two such maxima at 293°K (20°C). All these maxima lie between Card 2/3

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SOV/51-6-1-24/30 Single Grystels of ZnS and the Spectrum of Their Absorption Edge at Low Temperatures. S120 and 3200 Å. Acknowledgments are made to O.A. Katveyev and L.V. Maslova for their advice on the method of growing of single crystels. There are 2 figures, 1 table and 10 references, 6 of which are soviet, 3 English and 1 German. SUBMINTED: July 18, 1989 Card 3/3

Laborate 1

| 24(7), 24(6) AUTHORS: | SOV/51-6-4-29/29 Gross, Ye.F., Novikov, B.V., Razbirin, B.S. and Suslina, L.G. |
|--------------------------|--|
| TITLE : | Absorption Spectra of Crystals of Certain Gallium Chalcogenides (Spektry poglosncheniya kristallov nekotorykh khalkogenidov galliya) |
| PERIODICAL: | Optica i Spektroskopiya, 1959, Vol 6, Nr 4, pp 569-572 (USSR) |
| ABSTRACT: Card 1/2 | Linear structure in the long-wavelength edge of fundamental absorption was obsurved in the spectra of some semiconductors (Refs 1-10). These lines were ascribed by some authors to exiten states and by others to excess of one of the components of the semiconductor or to a foreign impurity. The present paper reports an investigation of the absorption apectra of gallium sulphide and selenide crystals (GaS and GaSe) with hexagonal laminar structure and crystals of β -Ga ₂ S ₃ and Ga ₂ Se ₃ . GaS crystals were obtained by melting together at 1000-1050°C stoichicmetric amounts of gallium and sulphur in evacuated quartz ampules. Crystals of β -Ga ₂ S ₃ were prepared similarly but at a higher temperature (1200-1250°C). Preparation of GaSe and Ga ₂ Se ₃ (cubic symmetry) was described by Goryunova et al (Ref 13). GaS and GaSe were used in the form of monocrystals of thicknesses varying from several microns to 100 p Ga ₂ S ₃ and Ga ₂ Se ₃ were 50-100 µ thick. Structure in the fundamental absorption edge was observed in the spectra of GaS and GaSe at 77°K (Figs 1a and 2a respectively). Such structure was also visible in the |
| Caro A/C | (rige is and is respectively). But is bruckers and side victors in the |

| | absorption spectrum of GaSe at room temperature. In contrast to GaS and GaSe, no structure was observed in the fundamental absorption edges of Ga2S3 and Ga2Se3 either at room temperature or at 77° K (Figs 1d and 2d). The absence of structure in the absorption spectra of ρ -Ga2S3 and Ga2Se3 is probably due to a large number of randomly distributed imperfections in these crystals. Such imperfections impede formation and migration of excitons and consequently the exciton lifetime is very short. Under such conditions the exciton structure of the absorption bands may be very diffuse or it may disappear altogether. From the absorption spectra the authors deduced the energy gaps in these semi- conductors. A table on p 571 lists the values of the energy gaps so deduced at 290°K (col 2) and 77°K (col 3). These values agree satisfactorily with those deduced from photoelectric measurements at room temperature, which are listed in col 3. Acknowledgments are made to N _{AA} . Goryunova for supply of GaSe and Ga2Se3 and for advice on preparation of GaS and Ga2S3 crystals. There are 2 figures, 1 table and 17 references, 10 of which are Soviet, 5 French and 2 German. |
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| SUBMITTED: | November 27, 1958 |
| Card 2/2 | |

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- 24(4), 24(6) AUTHORS: Bansi-Griyo, M. (Bancie-Grillot), Gross, Ye.F., Griyo, E.(Grillot) and Razbirin, B.S.
- TITLE: Studies of Linear Fluorescence and Absorption of Fure Cadmium Sulphide Crystals at the Temperature of 4.2°K (Issledovaniye lineychatoy fluorestsentsii i pogloshcheniya kristallov chistogo sernistogo kadmiya pri temperature 4.2°K)

PERIODICAL: Optika i Spektroskopiya, 1959, Vol 6, Nr 5, pp 707-710 (USSR)

ABSTRACT: The present note supplements earlier work (Refs 1, 2, 4) on fluorescence and absorption of CdS monocrystals observed at low temperatures. The monocrystals were produced by sublimation and their thicknesses were ~50 μ or less. A spectrograph of 4 Å/nm dispersion was used to obtain the spectra at 4.2°K. The fluorescence spectrum of sublimated crystals is shown schematically by the band A in the figure on p 709. This spectrum was found to contain one new line (4968.2 Å) in addition to those reported earlier (Ref 2). As before (Refs 1, 2), the fluorescence was mainly blue with very feeble emission at green wavelengths. The background between 4870 and 4942 Å and the fluorescence lines were polarized with the electric vector at right-angles to the optical axis of the monocrystal, suggesting a common origin for all of them. The absorption spectrum of the sublimated crystals is shown by the band B

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207/31-6-5-29/34 Studies of Linear Fluorescence and Absorption of Pure Jadmiu. Bulchido Crystals at the Temperature of 4.2°K

> (the figure on p 709); it did not vary from sample to sample. Three absorption lines (4853.1, 4857.2, 4869.1 Å) were observed in unpolarized light at positions very close to three fluorescence lines at 4856.6, 4861.4 and 4870.0 Å. In the region where continuous absorption was somewhat weaker, the light which had ussed through the crystal was completely polarized with the electric vector parallel to the optical axis of the crystal. The authors obtained also the fluorescence spectrum of a crystal which was not prepared by unblimition (the method of preparation is not given). This spectrum was characterized by a strong green band and a line structure in the blue region (D in the figure on p 709) which was quite different from that observed in sublimited crystals; neither the green band nor the blue lines were polarized. See also the following abstract. There are 5 references, 2 of which are Soviet, 1 Dutch, 1 French and 1 mixed (Gentan, Anglish and Rossian).

SUEMITTED: December 31, 1958

Card 5/2

| 4(4), 24(6) | .07/1-6-5-30/34 |
|-------------|--|
| AUTHORS : | Gross, Ye.F., Griyo, E. (Grillot), Zakharchenya, B.P. and Bansi-Griyo, M. (Baucie-Grillot). |
| title : | The Effect of a Magnetic Field on the Blue Fluoresten to and on the |
| | Absorption Lines of Some Pure Cadmium Sulphide Srystels st the |
| | Temperature of 4.20K (Vliyaniye magnitnogo polya na linii siney |
| | fluorestsentsii i na linii pogloshcheniya nekotorykh kristallev |
| | chistogo sernistogo kadmiya pri temperature 4.2°k) |
| | Optika i Spektroskopiya, 1959, Vol 6, Nr 5, pp 710-713 (, |
| ABS TRACT : | seports continuation of the work described in the proceeding of them |
| | (see preceding abstract). The present work was carried out at the |
| | Physico-Technical Institute of the Ac. Sc. USOR in Loningred. A 3d2 |
| | monocrystal propared by sublimation (dimensions 4 mm x 2 mm x 40-00 my |
| | was placed between the poles of an electromagnet. a diffraction |
| | spectrograph with 1.7 Å/mm dispersion was used to record the flaorescence |
| | spectrum of the crystal excited by the 3650 Å line at 4.2° n. In a |
| | magnetic field of 28 000 0e, oriented at right-angles to the optical |
| | axis of the crystal, the fluorescence lines at 4970, 4060 and 4001 a |
| | exhibited Zeeman splitting into doublets (separations of 0.5%. 0.5 and |
| | 1.2-1.3 Å respectively, cf. Fig 1). The doublet components were |
| | polarized in the same way as the original lines, i.e. with the electric |
| 1 | vector at right-angles to the optical axis of the crystel. To splitting |
| 1/2 I/2 | cas observed in magnetic field up to 28 000 0e, oriented carabel to |
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| RUMANIA/Op | tie | s - Spectroscopy. | К |
|-------------------------|-----|--|-------------|
| Abs Jour | : | Ref Zhur Fizka, No 2, 1960, 4671 | |
| Author Inst Title | :: | Gross, E.F. Important Problem of Modern Physics (Excitons Crystal Lattice) | in |
| Orig Pub | : | An. Rom/-Sov. Ser. matfiz., 1959, 13, No 2, | 128-136 |
| Abstract | : | Translated from Vestn. AN SSSR, 1958, No 10, Zhur Fizika, 1959, No 4, 9434. | See Referat |

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F. Y= This Conference both place from October 21 to Hernther 1 of Thisis is an erganized by the October 21 to Hernthemit-finities have Andersis and 1532 (Specimer, CT20-Anders) anthomatical Sciences of the Assisy of Stances, TS18). In Anderly and Gruinatory 538 (Lataby of Stances, TS18). The Anderly and Gruinatory 538 (Lataby of Stances, TS18). The Anderly and Gruinatory 538 (Lataby of Stances, TS18). The Anderly and Gruinatory 538 (Lataby of Stances, TS18). The Anderly and Gruinatory 538 (Lataby of Stances, TS18). The Anderly and Gruinatory 538 (Lataby of Stances, TS18). The Anderly and Cruinator 100 (Lataby of Science, TS18). The Anderly and Cruinator 100 (Lataby of Science, TS18). The Anderly and Cruinator 100 (Science frame Phart Stances, Edit V, Standrof V, Science 40 (Science edit), and This Passati Ander 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition and Addition are 10 aby 10 (Science edit) and Addition are 11 aby CS22. Abort 90 (Science edit) and Addition and Addition are 10 aby 10 (Science edit) and Addition and Addition are 10 aby 10 (Science edit) and Addition and Addition are 10 aby 10 (Science edit) and 10 aby 10 (Science edit) and 10 aby The of the next intersting leaves delivers it will con-former that by $[1, \dots, 1]$ there is a lister if will con-former and the ball there is a lister if will con-grandene and the intersting leaves deliver if will be delived is a lister (the intersting lister) if [1, 1] with a constraint is a lister (the intersting lister) if [1, 1] with a constraint is a lister of the outperturber if [1, 1] with a constraint is a lister of the outperturber if [1, 1] with a constraint is a lister of the outperturber if [1, 1] with a constraint is a lister of the outperturber if [1, 1] with a constraint is a lister of the outperturber if [1, 1] with a constraint is a lister of the outperturber of the outperturber is a lister is a lister of the outperturber of the lister of the lister is a lister of the lister of the lister of the lister of the lister is a lister of the lis A *** and *** and second with the involution as carried out a produced by Gurrich the involution as carried and a produced by Gurrich (1971 - and 7000 Berline) as Samh an in the involution of the a restor in the merican fraction and the involution of the and the involution of the anomynemic (1971) and the fraction of the involution (1971) apple about attract and the involution of the anomynemic (1971) apple and the involution of the involution fraction of the involution of the involution fraction of the involution of the involution fraction of the involution of the fraction of the involution of the involution of the involution fraction in the observation of the involution fraction of the involution of the involution in the involution in the involution involution fraction of the involution of the involution in the involution in the involution of the involution in the involution of the involution in the involution in the involution of the involution in the involution of the involution in the involution in the involution of the involution in the involution of the involution in the involution in the involution of the involution in the involution in the involution of the involution in the involution in the involution of the involution in the involution in the involution of the involution of the involution in the involution of the involution of the involution of the involution in the involution of the involution of the involution of the involution in the involution of the Fifth All-Dhies Canforeace on the Physics of Low Temporatures (3-ye Teensymissye soundehaiye ye fisike mistikh seeperstur) The second second second second is a second Сараћы fisiebeekith ламк. 1353. 7el fT. 3r 4. вр 743-750 (1558) Leatures delivered at 21 10 Can 2/1---19-65/2018 Chanteer, 2. PERCENCALS : LITURIT 2 (0) 17700 TTLE Caré 10/11 Card 3/11 4. đ 1 j

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GROSS, Ye.F.; KAPLYANSKIY, A.A.

Quadrupole optical excitation of the ground state of excitons in copper oxide crystals. Fiz. tver. tela 2 no.2:379-380 F '60. 1. Leningradskiy fiziko-tekhnicheskiy institut AN SSSR. (Excitons) Copper oxide crystals)

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| S/181/60/002/007/047/047/XX P006/B067 | |
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| THORS: Gross, Ye. F., Kaplyanskiy, A. A. | |
| TLE: Splitting of the Fundamental Absorption Edge of Cu ₂ O Due to the Removal of the Energy Band Degeneracy in Orientated Deformation of Crystals | |
| RIODICAL: Fizika tverdogo tela, 1960, Vol. 2, No. 7, pp. 1676-1677 | |
| XT: The authors studied the effect of deformation on the crystal spectra ich are related to 1) optical transitions between energy bands and excitation of exciton states. The effect of uniaxial compression of 20 crystals at 77°K on their spectra of the long-wave main absorption ge, the lines $\lambda = 6164$, 6085, and 6125 A inclusive, was studied. The 20 single crystals were compressed in the direction of the C ₄ , C ₅ , C ₂ mmetry axes (the direction of compression is indicated by P), and the sorption spectra were taken perpendicular to P (in the direction L). The sults are briefly discussed and shown in a figure Pictures were also ken in polarized light The main characteristics of the splitting | J |

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| Splitting of | the Fundamental Absorption | 5/181/60/00 | 02/007/047/047/XX |
| Edge of Cu ₂ O | Due to the Removal of the Degeneracy in Orientated | B006/B067 | ~~/ 00 // 04 // 04 // XX |
| that one of the contract of th | multiplicity and the relative the energy bands is single, and sme case at $\vec{k} = 0$). This exper- ical investigations (Ref. 11) is up-theoretical considerations, ith the 2p levels of 0^{2-}) is tr a levels of Cu) is single. Then Y Soviet and 4 US. | l the other i mental resul in which it w that the wal iply-degener | s triply-degenerate t is in agreement as shown, on the ency band (in ate, and that the |
| ASSOCIATION | Leningradskiy fiziko-tekhnich (Leningrad Institute of Physi AS USSR) | eskiy instit cs and Techn | ut AN SSSR ology of the |
| SUBMITTED: | May 17, 1960 | | |
| Card 2/3 | | | |

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| 9.4160(224),1003,1137) 26.24,200 AUTHORS: Gross YarF, Razbirin, B. S., and Safarov, V. I. TITLE: A Study of the Longwave Edge of Intrinsic Absorption of Polycrystalline Films of CdS and ZnSe at Low Temperatures PERIODICAL: Fizika tverdogo tela, 1960, Vol. 2, No. 11, pp. 2945-2949 TEXT: The authors prepared CdS and ZnS films by sublimating the powdery or crystalline initial substances onto glass or quartz bases; the films were between 0.04 and 1-2µ thick, the sputtering time ranging between 5 and 30 min. A total of over 100 specimens were examined which were all and 30 min. A total of over 100 specimens were examined which were all subjected to a considerable temperature gradient during sublimation (along the base the temperature varied from 600-800°C to room temperature). Fig.1 shows the spectra taken on a 0.42µ thick CdS film at 77.3°K in the range of the longwave absorption edge; while the absorption spectrum of the film has a smooth course on the "cold" part of base b, that of the film on the "warm" base (a) exhibits three distinct peaks at 4376, 4830, and 4706 A which can be assigned to the three lines with 4870, 4840, and 4720 A exhibited by the CdS monocrystal. The shift is ascribed to deforma- Card 1/3 | • | | 8644 | 9 | | |
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| AUTHORS: Gross varies, Razoffin, Drotting of entrinsic Absorption of TITLE: A Study of the Longwave Edge of Intrinsic Absorption of Polycrystalline Films of CdS and ZnSe at Low Temperatures PERIODICAL: Fizika tverdogo tela, 1960, Vol. 2, No. 11, pp. 2945-2949 TEXT: The authors prepared CdS and ZnS films by sublimating the powdery or crystalline initial substances onto glass or quartz bases; the films were between 0.04 and 1-2 μ thick, the sputtering time ranging between 5 and 30 min. A total of over 100 specimens were examined which were all subjected to a considerable temperature gradient during sublimation (along the base the temperature varied from 600-800°C to room temperature). Fig.1 shows the spectra taken on a 0.42 μ thick CdS film at 77.3°K in the range of the longwave absorption edge; while the absorption spectrum of the film has a smooth course on the "cold" part of base b, that of the film on the "warm" base (a) exhibits three distinct peaks at 4878, 4838, and 4706 A which can be assigned to the three lines with 4870, 4840, and 4720 A exhibited by the CdS monocrystal. The shift is ascribed to deforma- | 9.4160 (32 | .01,1003,1137) | | S/181/60/002 B006/B060 | /011/038/042 | |
| PERIODICAL: Fizika tverdogo tela, 1960, Vol. 2, No. 11, pp. 2945-2949 TEXT: The authors prepared CdS and ZnS films by sublimating the powdery or crystalline initial substances onto glass or quartz bases; the films were between 0.04 and 1-2 μ thick, the sputtering time ranging between 5 and 30 min. A total of over 100 specimens were examined which were all subjected to a considerable temperature gradient during sublimation (along the base the temperature varied from 600-800°C to room temperature). Fig.1 shows the spectra taken on a 0.42 μ thick CdS film at 77.3°K in the range of the longwave absorption edge; while the absorption spectrum of the film has a smooth course on the "cold" part of base b, that of the film on the "warm" base (a) exhibits three distinct peaks at 4878, 4838, and 4706 A which can be assigned to the three lines with 4870, 4840, and 4720 A exhibited by the CdS monocrystal. The shift is ascribed to deforma- | | Gross Va F., R | azbirin, B. S., a | and Safarov, V | '. I. | |
| PERIODICAL: Fizika tverdogo tela, 1960, Vol. 2, No. 11, pp. 2945-2949 TEXT: The authors prepared CdS and ZnS films by sublimating the powdery or crystalline initial substances onto glass or quartz bases; the films were between 0.04 and 1-2 μ thick, the sputtering time ranging between 5 and 30 min. A total of over 100 specimens were examined which were all subjected to a considerable temperature gradient during sublimation (along the base the temperature varied from 600-800°C to room temperature). Fig.1 shows the spectra taken on a 0.42 μ thick CdS film at 77.3°K in the range of the longwave absorption edge; while the absorption spectrum of the film has a smooth course on the "cold" part of base b, that of the film 4706 A which can be assigned to the three lines with 4870, 4840, and 4720 A exhibited by the CdS monocrystal. The shift is ascribed to deforma- | TITLE: | Polycrystalline | Films of cub and | | | |
| of the longwave absolution of the "cold" part of base b, that of the film film has a smooth course on the "cold" part of base b, that of the film on the "warm" base (a) exhibits three distinct peaks at 4878, 4838, and 4706 A which can be assigned to the three lines with 4870, 4840, and 4706 A which can be assigned to the three lines with 4870, 4840, and 4720 A exhibited by the CdS monocrystal. The shift is ascribed to deforma- | TEXT: The au or crystallin were between and 30 min. A subjected to the base the | thors prepared Cd ne initial substan 0.04 and 1-2µ thi A total of over 10 a considerable to temperature vario ectra taken on a 0 | S and ZnS films ces onto glass o ck, the sputteri O specimens were emperature gradie of from 600-800°C 0.42 thick CdS f | by sublimating r quartz base ng time rangi examined whi nt during sub to room temp ilm at 77.3 K | g the powdery s; the films ng between 5 ch were all limation (along erature). Fig.1 in the range rum of the | X |
| | of the longw film has a s cn the "warm 4706 A which 4720 A exhib | mooth course on t base (a) exhibi | he "cold" part of ts three distinct | base b, that t peaks at 487 | 8, 4838, and 4840, and | |
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A Study of the Longwave Edge of Intrinsic Absorption of Polycrystalline Films of CdS and ZnSeat Low Temperatures s/181/60/002/011/038/042 B006/B060

tion owing to the backing and the great number of defects. The first two peaks have a half-width of 10, the third one of 30 A. SnSe exhibits the same effect. The "hot" part exhibits three peaks, two distinct ones at 4430 and 4412 A, and a very weak one at 4365 A. The half-widths are the same. One of these maxima has been already observed by G. A. Zholkevich. An investigation of the films at 4.2°K did not give different results. Electron diffraction studies showed that the difference in the spectra of "cold" and "hot" films cannot be due to different crystallization forms or amorphism of the "cold" film. The differences are explained by the fact that films forming on the "hot" parts of the base consist of considerably larger and less defective crystals than those forming on the "cold" parts. This assumption has been checked experimentally. M. A. Rumsh and Y. N. Shchemelev as well as V. N. Vertsner and M. I. Rudenok are thanked for assistance in the experiments; K. V. Shalimova and N. V. Karpenko are mentioned. There are 3 figures and 7 references: 5 Soviet, 1 British, 1 French, and 1 German.

Card 2/3





CIA-RDP86-00513R00051703

86451 5/181/60/002/011/041/042 9,4160 (3201, 1003, 1137) B006/B060 26.2420 Grossever Fac and Kaplyanskiy, A. A-Study of the Effect of Oriented Deformations on the Spectrum AUTHORS of the Fundamental Absorption Edge of Cu₂O Single Crystals TITLE: Fizika tverdogo tela, 1960, Vol. 2, No. 11, pp. 2968-2981 TEXT: This is a very detailed report on studies made on monoaxially compressed Cu₂O single crystals. As a consequence of this deformation, an anisotropic polarized splitting of the longwave edge was observed to take place along with an exciton structure of the Cu_2O absorption spectrum. Multiplicity, amount of splitting, and polarization in the spectra were examined with different directions of compression The method used for the investigation was similar to the one described in Refs 13,14. The specimens were about 1% 2% 4 mm large platelets placed in a special press inside a Dewar vessel (77°K). The spectra were taken in polarized (EIP, ELP) and unpolarized light by an NCR 51 (ISP-51) spectroscope, dispersion $\sim 5A/mm$, and a $y_{\Phi}^2 - 85$ (UF-85) camera. The spectra were all taken Card 1/8

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86451 5/181/60/002/011/041/042 Study of the Effect of Oriented Deformations on the Spectrum of the Fundamental Absorption B006/B060 Edge of Cu₂O Single Crystals in the direction L which was perpendicular to the direction of compression P (L1P), with P being oriented along the axes C_4 , C_5 , and C_2 ; Figs. 1-3 show pictures of the polarized splitting of the absorption edge of Cu_2O_{π} The spectra in the figures are described in the text and certain characteristics are pointed out. The magnitude of the splitting is in all cases (L1P; $P \parallel C_4$, $P \parallel C_5$, $P \parallel C_2$) directly proportional to the compression pressure. The polarized splitting was calculated, and the theoretical and experimental splitting amounts Δ were intercompared ($\Delta = \gamma - \nu_0$, ν_0 being the position of the line in the free crystal). The agreement is satisfactory. From results obtained it was possible to infer the character of the energy levels. The results can be explained by assuming that the triple degeneracy of the valency band in deformation is removed at $\vec{K} = 0$. The final part of the paper under consideration deals with the effect of the crystal deformation upon the yellow exciton series of Cu₂O (V. T. Agekyan, a student of LGU (Leningrad State University) participated in Card 2/8

| Study of the Effect of Oriented Deformations S/181/60/002/011/041/042 Sige of Cu₂O Single Crystals Station) It was found that at P # C₄ the set is sharply shifted Sige of Cu₂O Single Crystals Station of longer waves; the same, but less markedly, applies to Section at P # C₂ a polarization effect appears in addition to this Shift. The uniform shift of the lines allows the conclusion that deformation has an effect upon the position of the band, but not upon the Section levels; a relationship was found, furthermore, to Section are are 7 figures, 2 tables, and 29 references: 17 Soviet, SUEMITTED: August 4, 1960 | <pre>budy of the Effect of Orientou Brows Brows in the Spectrum of the Fundamental Absorption B006/B060 dige of Cu₂O Single Crystals his investigation). It was found that at P C₄ the set is sharply shifted in the direction of longer waves; the same, but less markedly, applies to n the direction of longer waves; the same, but less markedly, applies to C₂, and at P C₂ a polarization effect appears in addition to this C₃, and at P C₂ a polarization effect appears in addition to this C₄, and at P C₂ a polarization of the band, but not upon the shift. The uniform shift of the lines allows the conclusion that deforma- hift. The uniform shift of the lines allows the conclusion that deforma- hift is an effect upon the position of the band, but not upon the individual exciton levels; a relationship was found, furthermore, to individual exciton levels; a relationship was found, furthermore, to exist between the hydrogen-like exciton levels with n =2 and the degenerate problems. There are 7 figures, 2 tables, and 29 references: 17 Soviet, 10 US, 1 French, 1 Japanese, and 1 Italian. ASSOCIATION: Fiziko-tekhnicheskiy institut AN SSSR Leningrad (Institute of Physics and Technology of the AS USSR, Leningrad)</pre> | | 8645 | | |
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| SUBMITTED: August 4, 1960 | | on the Spectrum Edge of Cu_2O Sithis investiga in the directi $P \parallel C_3$, and at shift. The unition has an eff individual exc exist between band. A. G. Zh problems. They 10 US, 1 Frem | m of the function ingle Crystals tion). It was found that at P [C ₂ on of longer waves; the same, but P C ₂ a polarization effect append form shift of the lines allows to ffect upon the position of the ba- citon levels; a relationship was the hydrogen-like exciton levels hilich is finally thanked for dis re are 7 figures, 2 tables, and 2 ch, 1 Japanese, and 1 Italian. Fiziko-tekhnicheskiy institut AM of Physics and Technology of the | the set is sharply shifted t less markedly, applies to ars in addition to this he conclusion that deforma- nd, but not upon the found, furthermore, to with $n \rightleftharpoons 2$ and the degenerate cussions on theoretical 29 references: 17 Soviet, | X |
| | Card 3/8 | SUBMITTED: | August 4, 1960 | | |

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Redst Signification of the absorption edge spectrum of Cu₂O on a crystal compression in the direction of the symmetry axis of 4th order; Fig. 2: the same on a compression in the direction of the symmetry axis of the symmetry axis of 2nd order. Legend to Fig. 7: spectrogram of the yellow exciton series of Cu₂O in monoaxially compressed crystals.

"APPROVED FOR RELEASE: Thursday, July 27, 2000















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| 24.350 | S/051/60/008/02/015/036 |
| AUTHORS: | Gross YesF., Razbirin, B.S. and Shekhmamet'yev, R.I. |
| TITLE: | Investigation of the Reflection and Luminescence Spectra of Copper Halides at Low Temperature |
| PERIODICAL | : Optika i spektroskopiya, 1960, Vol 8, Nr 2, pp 232 - 238 (USSR) |
| ABSTRACT: | This paper is based on the results of the diploma work of <u>B.S. Razbirin</u> and <u>R.I. Shekhmamet'yev</u> carried out at Leningradskiy gosudarstvennyy universitet im. Zhdanova (Leningrad State University im. Zhdanov) in 1955-1957. The paper reports the results of an investigation of the diffuse reflection and luminescence spectra of CuI (Figures 1, 2), CuBr (Figures 3-5) and CuCl (Figure 6) crystals at 77 K in the spectral region around the fundamental absorption edges of these three compounds (some of these results have been reported earlier, cf. Ref 8). The crystals were used in the form of sublimated layers deposited in vacuo on glass plates and in the form of fine-grained powders. Luminescence was excited with ultraviolet light from a mercury lamp SVDSh-1000: |
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68888 s/051/60/008/02/015/036 E201/E391 Investigation of the Reflection and Luminescence Spectra of Copper Halides at Low Temperature for CuI and CuBr the 3 660 Å wavelength was used and for CuCl shorter wavelengths (3 100 - 5 500 Å) were employed. The reflection spectra were obtained using a continuousspectrum source (an incandescent lamp). The reflection and luminescence spectra of the same sample were recorded by means of a quartz spectrograph Q-12 with 50 Å/mm dispersion in the 4 000 Å region. The results obtained can be summarized as follows: 1) The reflection and luminescence spectra of copper halides are similar. The positions of the long-wavelength absorption edges of CuCl, CuBr and CuI do not differ greatly. Reflection maxima identical with absorption lines are obtained for all these crystals in the region of the absorption edge. The luminescence spectra of the three crystals have groups of narrow lines, of which those lying at shorter wavelengths coincide with the appropriate absorption lines. The luminescence spectra contain also wide bands at longer wavelengths; 2) Adsorbed gases affect strongly the structure of the Card2/3 ADDRESS OF A STREET AND ADDRESS OF A STREET ADDRESS OF ADDRES

| 68888 S/051/60/008/02/015/036 E201/E391 Lides at Low Temperature |
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| <pre>luminescence and reflection spectra of CuI and CuBr crystals. After adsorbed gas is removed the luminescence and reflection spectra recover their original form (this process can be repeated many times); 3) The luminescence spectra of CuI and CuBr contain groups of equidistant lines similar to those observed in other semiconducting crystals (CdS, CdSe, ZnS, ZnO, etc); 4) The results obtained indicate that the short-wavelength weak luminescence lines of CuI, CuBr and CuCl, which coincide with absorption lines and are not greatly affected by surface treatment, are due to processes occurring in the crystal lattice. The long-wavelength strong luminescence lines, which are very sensitive to surface treatment, are due to some processes occurring at the surface. There are 6 figures and 12 references, 6 of which are Soviet, 3 English and 2 French, 1 German.</pre> |
| UBMITTED: June 3, 1959 |
| ard 3/3 |
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| | S/051/60/008/04/014/032 B201/B091 | |
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| HORS : | Gross, Ye.F., Susling, L.G. and Komarovskikh, K.F. | |
| 1.2 : | Gross, Ye.F., Susline, L.G. and Russicvillary for Sulphide Crystals Investigation of the Absorption Spectra of Zinc Sulphide Crystals | |
| IODICAL | Optika i spektroskopiya, 1960, Vol 8, Nr 4, pp 516-520 (USSR) | |
| BTRACT : | The paper deals with polarization of the line structure of the absorption edge of hexagonal ZnS monocrystals at 4.2 and 77° K and with the absorption spectra of sublimated polycrystalline ZnS films. Nonocrystals of ZnS were grown in the authors' laboratory by sublimation of ZnS powder in a neutral atmosphere (Ref 2). To sublimation of ZnS powder in a neutral atmosphere (Ref 2). The other monocrystals was found to subtract and and natural ultraviolet light using a quarts spectrograph Q-12 (25 Å/mm linear dispersion in the 3200 Å region). The line structure of the absorption edge of ZnS monocrystals was found to be structure of the absorption edge of ZnS monocrystals was completely polarized polarized (Fig 1). At 4.2°K the 3205 Å l | / |
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rsday, July 27, 2000 CIA-RDP86-00513R00051703

"APPROVED FOR RELEASE: Thursday, July 27, 2000

S/051/60/008/04/014/032 **B201/B691** Investigation of the Absorption Spectra of Zinc Sulphide Crystals oscillator. The 3160 & line was unpolarized or slightly polarized and the 3115 Å line was strongly polarised with E || c (the absorption oscillator close to a linear electric dipole). The nature of polarization of the spectrum of ZnS monogrystals is similar to polarization found in other uniaxial crystals with discrete structure of the absorption edge (Refs 4-6). Thin crystals $(d \sim 0.1 \mu)$ were found to stick to the base and the consequent deformation produced displacement of the absorption lines (Table 1), as well as broadening (the 3205 Å line). The absorption spectra of some "free" crystals were also found to be displaced by 1-2 Å due to internal stresses produced during growth of the monocrystals or due to differences in attachment The authors investigated also the absorption spectra (Fig 2) of polycrystalline ZnS films produced by sublimation in vacuo. to the bases. At 4.2°K the following absorption lines were observed (the widths are given in brackets): $3212 \stackrel{1}{\downarrow} (10 \stackrel{1}{\downarrow})$, $3190 \stackrel{1}{\downarrow} (10 \stackrel{1}{\downarrow})$, $3128 \stackrel{1}{\downarrow} (20 \stackrel{1}{\downarrow})$; all these are shown in Fig 2a. In some films a weak line at 3271 Å (20 Å width) could be seen (Fig 23). The positions of the absorption lines in polycrystalline ZnS films at 4.2 and 77°K are listed in Table 2. Comparison of the data in Tables 1 and 2 shows that the positions and Card 2/3

| Investigati | S/051/60/008/04/014/032 E201/E691 widths of the absorption lines of polycrystalline films are close |
|-------------|---|
| | to the positions and widths of ZnS monocrystals stuck to substrates. It follows that the films have hexagonal structure. The 3212, 3190 and of monocrystals are the lines of the lattice itself (exciton excitation). The 3271 Å line of ZnS films is due to lattice defects (for example excitation of "impurity" excitons in the defective regions). There are 2 figures, 2 tables and 9 references, 8 of which are Soviet and |
| SUEVITED : | August 13, 1959 |
| Card 3/3 | |
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83927 s/051/60/009/004/030/034 E201/E191 26.1512 **B**. Grillot Ye.F. Gross, Bancie-Grillot, M., AUTHORS: and Razbirin, B.S. The Effect of Temperature on Two Series of Bands in TITLE: the Green Fluorescence Spectrum of Pure Cadmium Sulphide at Low Temperatures of PERIODICAL: Optika i spektroskopiya, 1960, Vol 9, No 4, pp 542-544 Very pure cadmium sulphide, which does not luminesce at room temperature, exhibits intense green fluorescence at the TEXT: temperature of liquid air (Refs 1-3). The spectrum consists of two vibrational series whose maxima are given by $v_1 = 19450 - 300n \text{ cm}^{-1}$ and $v_2 = 19310 - 300p \text{ cm}^{-1}$, where n and p are small integers. The present paper reports further studies on the effect of temperature on the relative intensities of the two series, between 4 and 77 °K. In some crystals only the second series (N_2) was observed at 4 °K; heating of these crystals to the boiling point of liquid nitrogen destroyed gradually this series, which was (also gradually) replaced by the first series at 77 °K. If a crystal exhibited only the first series at $\frac{1}{4}$ °K, then Card 1/2

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^R3927 **S/051/60/009/004/030/034 E201/E191** O Series of Bands in the Groov

The Effect of Temperature on Two Series of Bands in the Green Fluorescence Spectrum of Pure Cadmium Sulphide at Low Temperatures

heating to 77 °K did not produce the second series. In crystals with both series at 4 °K heating to 77 °K weakened the bands of the second series so that only the first series (slightly broadened) remained at 77 °K, as shown in Fig 1. On application of an electric field (about 1 kV/cm) to a crystal immersed in liquid helium and exhibiting both series, the intensity of the first series bands was raised and the intensity of the second series was lowered, as shown in Fig 2. Further studies of the effects of electric fields are proceeding. Acknowledgement is made to N.M. Reynov for his help in work with liquid helium. There are 2 figures and 4 references: 1 Dutch, 1 French and 2 mixed (English, German, Dutch, Russian and French).

SUBMITTED: May 3, 1960

Card 2/2

"APPROVED FOR RELEASE: Thursday, July 27, 2000

8/020/60/132/01/25/064 T., Corresponding Member of the AS USSR, Kaplyanakiy, A.A. The Optical Anisotropy of Gubic Crystals Which Gauses the Effect of Spatial Dispersion. Quadrupole Exciton Absorption of Light 1 Guprous Oxide 7/ n PERIODICAL: Doklady Akademii nauk SSSR, 1960, Vol. 132, No. 1, PP. 98-101 24.7100 When examining the absorption spectrum of monoorystalline samples of $(T = 77^{\circ} \text{ K})$ the authors of the present paper detected the effect of TEXT: When examining the absorption spectrum of monocrystalline samples TEXT: (T = 77° K) the authors of the present paper detected in the case of the present paper detected in the through the set of the present paper detected in the through the set of the present paper detected the paper detect $v_{2}v_{1} = (1 - k)^{v_{1}} e^{auv_{1}v_{1}v_{1}} e^{av_{1}v_{1}v_{1}} e^{av_{1}v_{1}v_{1}} e^{av_{2}v_{2}v_{2}} u_{1} e^{av_{2}v_{2}v_{2}} u_{1} e^{av_{2}v_{2}v_{2}} e^{av_{2}v_{2}v_{2}v_{2}} e^{av_{2}v_{2}v_{2}} e^{av_{2}v_{2}v_{2}} e^$ AUTHORS: TITLE: cubic orystals. It was found that in the spectrum of light passing throu cu₂O lamina the intensity of these absorption lines perpendicular to eac arbitrarily chosen states of polarisation which are perpendicular Cu₂O lemine the intensity of these absorption lines is not equal in two arbitrarily ohosen states of polarization which are perpendicular from spe other. The degree of polarization and the integral intensity vary from arbitrarily chosen states of polarization which are perpendicular to each from speci-other. The degree of polarization and the integral intensity vary A sufficiention of the beam obanges. other. The degree of polarization and the integral intensity vary from speci-nee to specimen and also when the direction of the beam ohanges. Of intensity large Cu2O orystal was bred and used to determine of the beam in the orystal and polarization of these lines upon the direction of the beam in the set of the set o large Cu20 orystal was bred and used to determine the dependence of intensity and polarization of these lines upon the direction of the beam in the orystal Card 1/3 APPROVED FOR RELEASE: Thursday, July 27, 2000 CIA-RDP86-00513R00051703(

CIA-RDP86-00513R00051703

80053 s/020/60/132/01/25/064 The Optical Anisotropy of Cubic Crystals Which Causes B014/B014 the Effect of Spatial Dispersion. Quadrupole Exciton lattice, Thin sheets, which were differently oriented relative to the crystallo-Absorption of Light in Cuprous Oxide graphic axes, were cut out of this crystal. It was found that the intensity and the degree of polarization of the lines under consideration are fully determined by the orientation of the beam relative to the crystallographic axes. The results obtained are described by means of the spectrogram shown in Fig. 1 and the scheme shown in Fig. 2. Herefrom it may be seen that the spatial distribution of intensity and of the state of polarization has some elements of cubic symmetry. When taking account of polarization only (neglecting absorption intensity) it is possible to determine seven "optical" ares along which absorption is isotropic. The anisotropy of the absorption lines is ascribed by the authors to electric quadrupole transitions since the observable dependence of intensity and polarization upon the direction of light inside the crystal corresponds to the spatial field distribution of an electric quadrupole system. In discussing the results obtained here the authors refer to papers by S.I. Pekar et al. (Refs. 12 and 13) in which it was pointed out that optical anisotropy of cubic crystals associated with the occurrence of excitons is theoretically possible. Furthermore, a paper by V.L. Ginzburg (Ref. 14) is referred to, in which spatial light dispersion was taken into consideration. Card 2/3


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| 24.7700 AUTHORS: | Gross, Ye. F., Corresponding Member of the AS USSR, |
| T IT LE : | <u>Photoluminescence</u> Within the Edge of the Fundamental Absorp- tion of Mixed <u>CdSe - CdS</u> Crystals |
| PERIODICAL: | Doklady Akademii nauk SSSR, 1960, Vol. 133, No. 1, pp. 56-59 |
| structure of number of of absorption luminescent crystals as | cheir long introduction the authors discuss the complicated of emission and absorption arising at low temperatures in a brystals (CdS, CdSe, HgI ₂ , ZnS, and others) within the longwave edge. In the present paper, the authors study the photo- edge. In the present paper, the authors study the photo- edge of macrocrystalline CdSe-CdS solid solutions of CdSe single of macrocrystalline CdSe- and CdS layers within their absorp- and of macrocrystalline CdSe- and CdS layers within their absorp- mathematical control of the structure is discussed in greater is pointed out. The structure is discussed in greater |
| 1 | n the first chapter. The great analogy with the spectrum greater crystals is pointed out. The structure is discussed in greater d, among other things, the great differences existing between |

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Photoluminescence Within the Edge of the Fundamental Absorption of Mixed CdSe - CdS Crystals 81717 S/020/60/133/01/15/070 B014/B011

the bands of different crystals are described. The second chapter treats pure macroorystalline CdS- and CdSe layers. Agreement is found between pure macroorystalline CdS- and CdSe layers. Agreement is found between the emission and absorption lines of the CdS layers and those of the CdS the gristals. The emission lines of CdS layers at T = 4.2 K exhibit single crystals. The emission lines of CdS layers at T = 4.2 K exhibit riplet structure, whereas the single crystals have a doublet structure. According to the authors' results, the emission of CdSe layers has a friplet structure. At T = 77.3 K the emission of the CdS layers consists of structureless bands, the CdS single crystals and pure CdSe layers have of structureless bands, the CdS single crystals and pure CdSe layers have a doublet structure. The third chapter deals with the macroorystalline layers of mixed CdSe-CdS crystals. In the case of T = 4.2 K, the photolayers of all of the 20 samples under consideration has a structure, luminescence of all of the 20 samples under consideration has a structure, and the line spectrum consists of a few weak lines. On heating to 77.3 K, the emission intensity drops, the clearness of the structure and the inthe shortwave lines of the edge emission likewise drop sharply, tensity of the shortwave lines of the edge emission likewise drop sharply, while the intensities of the shortwave components of the doublet and triplet structures rise. There are 3 figures and 18 references: 8 Soviet, 2 French, 3 German, 1 British, and 4 American.

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| Photoluminesc Fundamental A CdSe - CdS Cr | | 1717 0/133/01/15/070 11 |
| ASSOCIATION: | Fiziko-tekhnicheskiy institut Akademii (<u>Institute of Physics and Technology o</u> Sciences, USSR) | nauk SSSR If the Academy of |
| SUBMITTED : | March 28, 1960 | Y |
| Card 3/3 | | |





CIA-RDP86-00513R00051703

GROSS, Yevgeniv F. and NOVIKOV, B. V.

"Fine Structure of Spectral Curves for Excitation of Photoconducttivity and Luminescence and its Connection with Exciton Absorption."

REPORT TO be submitted for the Intl. Conference on Photoconductivity, IUPAP, Cornell University, Ithaca, N. Y., 21-24 Aug 1961.

Leningrad State Univ.

CIA-RDP86-00513R00051703

89299 5/181/61/003/001/041/042 B102/B204 26.2421 **AUTHORS:** Gross, Ye. F., Zakharchenya, B. P., and Konstantinov, O. V. TITLE: Effect of the inversion of a magnetic field in the exciton absorption spectrum of a CdS crystal PERIODICAL: Fizika tverdogo tela, v. 3, no. 1, 1961, 305-308 TEXT: Studies of the effect of a magnetic field upon the absorption spectrum of CdS, on which the authors have made a report in Ref. 1, are intended to determine the exciton energy spectrum and its relation to the band structure in CdS. The experiments described here were carried out with $1 - 3\mu$ thick foils of CdS single crystals, whose optical axis \vec{A} was in the plane of the foil. H was either parallel or perpendicular to \vec{A} . (A is considered to be a vector, because the crystal has no inversion center). The crystals were cooled to 1.3°K and remained free from deformation. In the case of E II A, the exciton absorption lines with $\lambda = 4853$, 4813, and 4806 A were weak and so narrow that the effect of the H-field upon them could be easily observed. The line with $\lambda = 4813$ A, on Card 1/4

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Effect of the inversion of a magnetic ...

which the inversion effect could be best observed, had a satellite line with $\lambda = 4814$ A. At $\vec{X} \perp \vec{H}$, the 4813-line split up into a doublet, whose center of mass was shifted toward higher energies relative to the original line. The weak 4814-line, whose origin is not quite clear, is also split up into a doublet; the components are weak and not so far apart as those of the main line. In the case of inversion of the field direction, the manner of splitting is considerably changed (shift of the main doublet $\Delta \lambda = 0.4$ A; intensity change). The essential change in the spectrum occurring when the field direction is inverted, consists in a shift of the Zeeman components and in a change of their intensity; the same effect is attained if the field is left as it is, and the crystal is rotated through 180°. Also the line with 4853 A, which is not split in the field, shows no effect of inversion. The line with 4806 A shows a complex splitting, and the inversion effect may be observed only with difficulty. The change of the absorption spectrum cannot be explained within the framework of the spectroscopy of an isolated atom, above all, because the effect is in contradiction to the invariance of the Schrödinger equation with respect to time reversal. The question is now

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GROSS, Ye.F.; SHEKHMAMET'YEV, R.I.

Connection between edge luminescence and the structure of the basic absorption edge. Fiz. tver. tela 3 no. 3:889-894 Mr '61. (MIRA 14:5)

1. Leningradskiy gosudarstvennyy universitet imeni A.A. Zhdanova. (Absorption of light) (Luminescence)

CIA-RDP86-00513R00051703

GROSS, Ye.F.; ZAKHARCHENYA, B.P.; KANSKAYA, L.M.

Investigating the Stark effect of excitons in oriented single crystals of cuprous oxide. Fiz. tver. tela 3 no. 3:972-978 Mr '61. (MIRA 14:5)

1. Fiziko-tekhnicheskiy institut AN SSSR, Leningrad. (Stark effect) (Copper oxide Spectra)

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22059 s/181/61/003/00.1/025/030 B102/B209 9,4177(1051,1482 Gross, Ye. F. and Novikov, B. V. AUTHORS: The relation between background and the fine-structure TITLE: maxima of the spectral curves of photoconductivity in CdS single crystals Fisika tverdogo tela, v. 3, no. 4, 1961, 1249-1252 PERIODICAL: TEXT: In previous papers (ZhTF, vyp. 4, 913, 1956 and DAN SSSR, 110, no. 5, 761, 1956), the authors reported on the discovery of a complex structure of the spectral photocurrent distribution at $T = 77^{\circ}K$ in the range of the known exciton absorption lines. These distribution curves may be divided into two classes: The first class contains those in which the exciton absorption lines coincide with the photocurrent maximum, while the second class encompasses such in which the exciton absorption lines coincide with the photocurrent minima. According to this classification, the crystals differ essentially in the character of the short-wave drop of photoconductivity. In a later paper, the effect of defects and surface condition of CdS crystals upon absorption lines and photocurrent curves was studied. These studies Card 1/4

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The relation ...

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have now been continued. Both classes of CdS crystals have a structureless background. Fig. 1 shows that the relative values of maxima and background differ greatly in different specimens. In the curves denoted by a, E was perpendicular to C (E - electric vector of the exciting light, C - optical axis of the crystal), whereas in those indicated by b, E was parallel to C. Like M. S. Brodin, the authors found that the background is considerably polarized in the direction of the electric vector EIC. The structure of the photoconduction curves vanishes in many cases if the surface of the crystal is subjected to a slight treatment. In this manner, curve a in Fig. 2 was obtained from as (Fig. 1) by wiping the surface of the crystal with wet cotton. The sensitivity of the specimen decreased. The structure of the curve after polishing had such a shape that the crystal had to be assigned to the second class. The authors also determined the spectral distribution curves of polycrystalline CdS films sputtered upon glass tackings. In such films which exhibited a structured absorption edge, also a structure of the photoconduction curves was found. The background was very high in this case, and the fraction of radiation used to determine the structure accounted only for some per cent. These facts speak in favor of an interrelation between the photoactive background (or part of it) and the lattice imperfections. Films

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| rent curves a Sheynkman, N. 2 figures and two reference | t 77°K. V. L. Broude, V. N. Chikovani, and M. S. 10 references: 7 Sovie s to English-language pu | n edge had completely smooth photocur- V. V. Yeremenko, V. S. Medvedev, M. K. . Brodin are mentioned. There are et-bloc and 3 non-Soviet-bloc. The ublications read as follows: D. Dutton, omas, J. J. Hopfield, Phys. Rev. <u>116</u> , | |
| ASSOCIATION: | Leningradskiy gosudars Fizicheskiy institut (1 A. A. Zhdanov, Institut | tvennyy universitet im. A. A. Zhdanova Leningrad State University imeni te of Physics) | |
| SUBMITTED: | September 26, 1960 | X | |
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24933 Effect of recoil of momentum and...

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 ΔE_{exc} . On account of the dependence of M_{exc}^{*} on the direction of propagation of light, the exciton recoil effect should also depend on the angle, and in anisotropic crystals on the direction of polarization of light. The effect should be experimentally observable in exciton spectra, e.g. from a determination of $v_{abs} - v_{emiss}$. Such a line displacement of $\sim 10^{-4}$ ev has in fact been observed in luminescence spectra (HgJ₂, CdS). However, it is not yet granted that this displacement has no other crigin. There are 6 references: 3 Soviet-bloc and 3 non-Soviet-bloc.

ASSOCIATION: Leningradskiy gosudarstvennyy universitet im. A. A. Zhdanova (Leningrad State University imeni A. A. Zhdanov)

S.B.4. 88666

SUBMITTED: February 23, 1961

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APPROVED FOR RELEASE: Thursday, July 27, 2000 CIA-RDP86-00513R00051703(

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| | 29693 S/181/61/003/010/019/036 B104/B108 |
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| 24,2100 (| Gross, Ye. F., Zakharchenya, B. P., and Razbirin, B. S. |
| TITLE: | Magneto-optical effects in the absorption spectrum of a cadmium-sulfide crystal |
| PERIODICAL | Fizika tverdogo tela, v. 3, no. 10, 1961, 3083 - 3091 |
| cadmium-sul Experiments tures of 4. | Zeeman displacement of the two groups of absorption lines of fide crystals was investigated (4889 - 4860 Å; 4860 - 4660 Å). were made in magnetic fields of up to 35,000 oe at tempera- were made in magnetic fields of up to 35,000 oe at tempera- and 1.3 K. The long-wave group was investigated with the crystals (from ~1µ up to some tens of microns). The orystals (from ~1µ up to some tens of microns). The of the diffraction-grating spectrograph used was 4 Å/mm and of the diffraction depend on the polarization and on Line splitting was found to depend on the polarization and on |
| dispersion 1.7 A/mm. the directionserved wh | Line splitting was found to depend on the permethy was on of the magnetic field. A diamagnetic line shift was high is increasing with the magnetic field strength and with the her (in the case of the hydrogen-like lines). The Zeeman her (in the case of the hydrogen-like lines). The Zeeman of the weak lines between 4889 and 4854 Å was not uniform for of the weak lines between 4889 and 4854 Å was not uniform for studied. In a discussion of these results the authors show that |

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Magneto-optical effects in the ...

an electric field acts on the exciton levels in a magnetic field. A. G. Samoylovich and L. A. Korenblit (DAN SSSR, 100, 43, 1955) studied the action of a Lorentz field on an exciton moving in a magnetic field. The results obtained here are explained as follows: The excitons of a CdS crystal have a dipole moment caused by the asymmetry of the intracrystalline field. The axia of this dipole is directed along the optical axis A of the crystal. If ANH, the electric Lorentz field is perpendicular to the dipole axis, and if ALH, it is parallel to the dipole axis. In the first case, the Stark effect obviously reaches a minimum. In the second case, a Stark effect is observed on exciton levels of greater radii. The discovered diamagnetic shift of absorption lines confirms the existence of exciton series which are related to the complex band structure in a CdS crystal. The Zeeman effect proves the complex X energy structure of an exciton in a CdS crystal. The appearance of a Lorentz field in magneto-optical exciton effects indicates the existence of a movable exciton system. There are 3 figures, 2 tables, and 12 references: 8 Soviet and 4 non-Soviet. The three most recent references to English-language publications read as follows: E. F. Gross, J. Phys. Chem. Sol., 8, 172, 1959; J. J. Jopfield and J. G. Thomas, Phys.

Card 2/3

| 24,3500 (1 | 29699 B125/B102 | |
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| AUTHORS: | Gross, Ye. F., and Snekhmanes July | |
| TITLE: | Complex structure of excitation spectra of luminescence of HgL and PbL crystals | |
| PERIODICAL | Pizika tverdogo tela, v. 3, no. 10, 1961, 3162 - 3166 | • |
| HgI ₂ crysta V. A. Arkha i spektr., an HgI ₂ cry light source authors ' la | authors determined the excitation curves for luminescence of ls and of sublimed PbI_2 layers at $T = 77^{\circ}K$. According to ngel'skays and P. P. Feofilov (DAN SSSR, <u>108</u> , 803, 1956; Opt. II, vyp. 1, 1957), the intensity of each luminescence band of stal is in different ways a function of the intensity of the the first series of HgI_2 crystals has been grown in the aboratory by K. F. Lider who employed slow crystallization from of HgI_2 in acetone. A second series of specimens was grown | |
| | aseous phase. Fig. 7 shows one random shows that $T = 77^{\circ}K$. No a function of the exciting wavelength at $T = 77^{\circ}K$. No | X |
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| Complex structure of | 29699 S/181/61/003/010/028/036 B125/B102 |
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| absorption lines correspond to the app | parent peaks at 5360 and 5280 R |
| $(\pm 20 - 30$ Å) found on the excitation citation curve for red luminescence of | curve 2. The structure of the ex- |
| be hardly influenced by surface treatm correspond to various crystal centers luminescence are closely related to th shows the excitation curves for red lu at T = 77°K. For excitation with ENC | nent. The various luminescence bands . The yellow-green and the red he exciton absorption lines. Fig. 2 uminescence of an HgI ₂ single crystal |
| on the excitation curve. For E1C the The latter is due to incomplete polar: | peaks at 5330 and 4932 X will occur. izetion of the absorption line |
| $\lambda = 4932$ Å. The authors also investig low-temperature luminescence of various | gated the excitation spectrum of the VA |
| sublimated at ~400°C upon a glass back curved, deformed, and are shifted rela of PbI ₂ single crystals toward the sh | king. The spectral lines found are tive to the absorption line $\lambda = 4948$ |
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| | | ose connection between broad lumines- ls and the structure of the self- | • |
| curves could experiments h luminescence ures and 8 re English-langu | correspond to lines win nave shown that exciton excitation of crystal efferences: 6 Soviet and tage publication reads | nima of the luminescence excitation th exciton structure. The authors' s play an essential part in the centers and defects. There are 3 fig- d 2 non-Soviet. The reference to the as follows: S. Nikitine et G. Perny, Phil. Mag., <u>4</u> , 1, 1959. | |
| ASSOCIATION: | | tvennyy universitet im. A. A. Zhdanova rsity imeni A. A. Zhdanov) | L . |
| SUBMITTED: | June 2, 1961 | | Х |
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CIA-RDP86-00513R00051703

30804 s/181/61/003/011/051/056 24,3430 (1227,1395,1163) B104/B138 Gross, Ye. F., Kalyuzhnaye, G. K., and Nedzvetskiy, D. S. AUTHORS: Complex structure of the absorption spectrum of mono-TITLE: crystalline gallium phosphide Fizika tverdogo tela, v. j, no. 11, 1961, 3543-3545 PERICDICAL: TEXT: Single crystals of GaP were investigated at nitrogen temperature. Single crystals 4 to 5 mm long, 0.3 mm to a few microns thick were obtained from the melts by crystallization (G. Wolff et al., Bull. Am. Phys. Soc., 29, 1, 1954). In transmitted light this crystals appeared orange and thick ones yellow-green. The absorption spectra were taken with an $MC\Pi-67$ (ISP-67) spectrograph with a camera of 1500 mm focal length. In the region studied the dispersion was 10.5 A/mm. The absorption edge of a GaP single crystal is shown in Fig. 1. This spectrum was taken for specimens that had Vbeen cooled slowly. Rapidly cocled specimens had only one broad line (5363.2 Å) which is shifted into the long wave range by a few angstroms. The lines can be grouped in pairs: an intense narrow and a weak narrow line, a weak and a strong broad line, and two broad lines. The distance Card 1/42 No LAS PROPERTIES

| - | 30804 S/181/61/003/011/051/056 ure of the absorption B104/B135 nes in each pair is 130 cm ⁻¹ within the limits of error. It | |
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| is concluded f consists of th to two discret action of ligh There are 2 fi The three most follows: E. O | rom this structure that the valence band of the crystal ree bands from each of which the electrons make transitions e levels below the bottom of the conduction band, under the t. A possible energy level scheme is shown in Fig. 2. gures, 1 table, and 7 references: 2 Soviet and 5 non-Soviet. recent references to English-language publications read as Kane. J. Phys. Chem. Solids. 1, 249, 1957; F. Stern, R. M. Rev., <u>108</u> , 158, 1957; R. Braunstein. J. Phys. Chem. Solids, | X |
| ASSOCIATION: | Leningradskiy gosudarstvennyy universitet im. A. A. Zhdanova (Leningrad State University imeni A. A. Zhdanov) | |
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