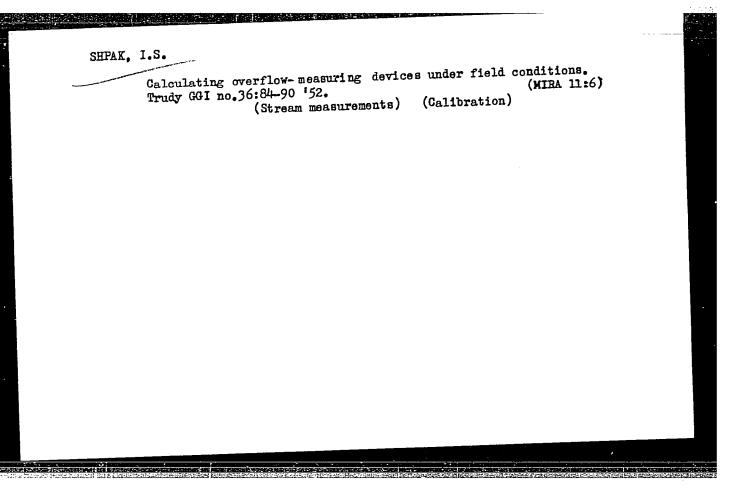
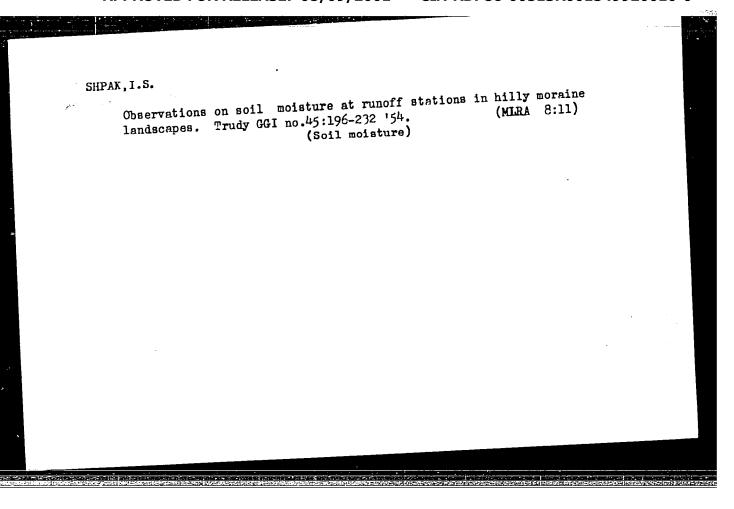
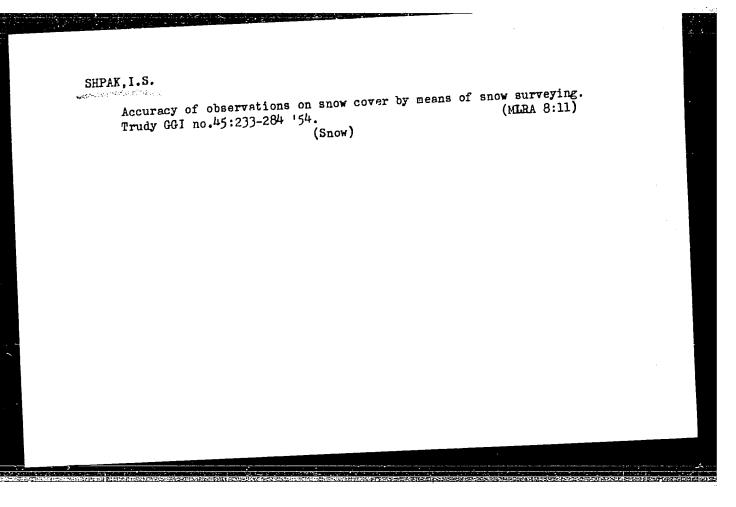


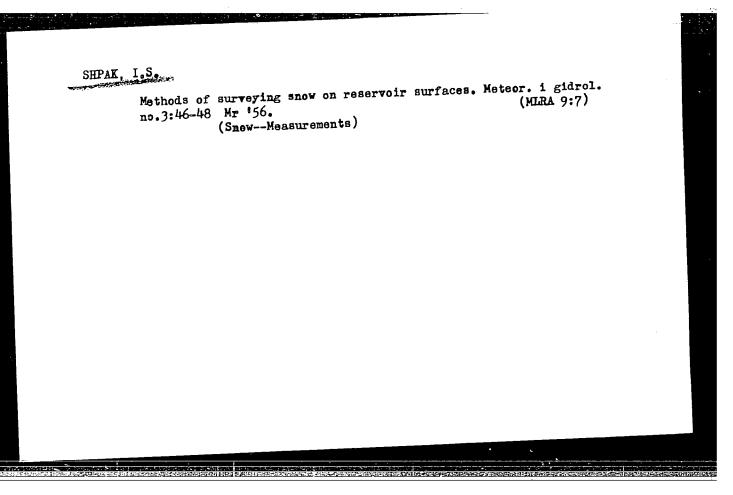
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Squall in the region of the Tsimlyansk Reservoir, June 25, 1956.

Meteor.i gidrol. no.4:28-30 Ap '57. (MLRA 10:5)

(Tsimlyansk Reservoir--Storm, 1956)

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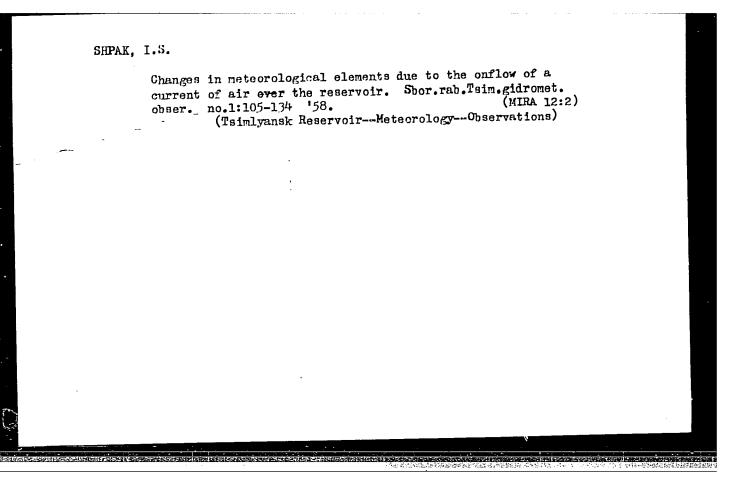
sov/1945

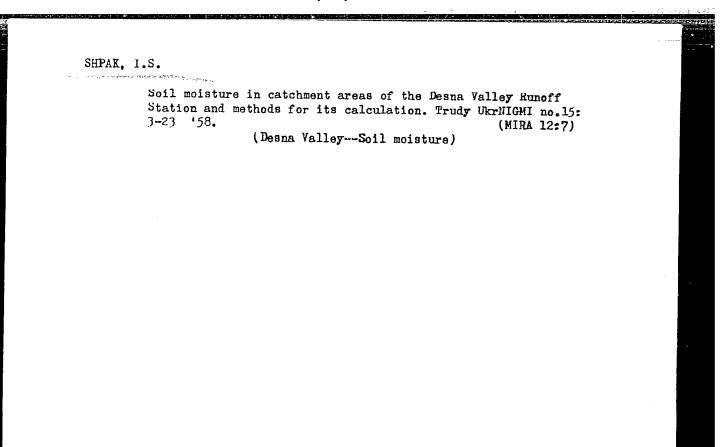
Tsimlyanskaya gidrometeorologicheskaya observatoriya

- Sbornik rabot...Vyp. 1. (Collected Papers of the Tsimlyansk Hydrometeorological Observatory Nr. 1) Leningrad, Gidrometeoizdat, 1958. 159 p. 460 copies printed.
- Additional sponsoring agency: USSR. Glavnoye upravleniye gidrometeorologicheskoy sluzhby.
- Ed. (Title page): P.P. Kokoulin; Ed. (Inside book): Z.I. Mironenko; Tech. Ed.: M.Ya. Flaum.
- PURPOSE: This publication is intended for all specialists concerned with the study and exploitation of water reservoirs and large lakes.
- COVERAGE: This collection of articles is concerned with a study of the following factors as they concern the Tsemlyanskoye Water Reservoir: wind produced agitation in the lake, the formation of

card 1/3

sov/1945 Hydrometeorological Observatory (Cont.) shorelines, changes in the meteorological conditions induced by the flow of air currents onto the reservoir surface, surface evaporation, and the gaseous regime of the lake. The studies are based on data obtained from observations. This information is shown in tables and graphs. Each article is accompanied by diagrams, tables, and bibliographic references. TABLE OF CONTENTS: 3 Foreword 5 Kokoulin, P.P. Tsimlyanskoye Reservoir Kokoulin, P.P., and L.V. Kokoulina. Data Obtained in Carrying Out Observations on the Formation of the Tsimlyanskoye 13 Reservoir Shoreline Kokoulin, P.P., and Ye.F. Semenov. Methods and Results Obtained in Observing Wave Patterns on the Tsimlyanskoye Reservoir 65 During the Years 1953-1955 Card 2/3



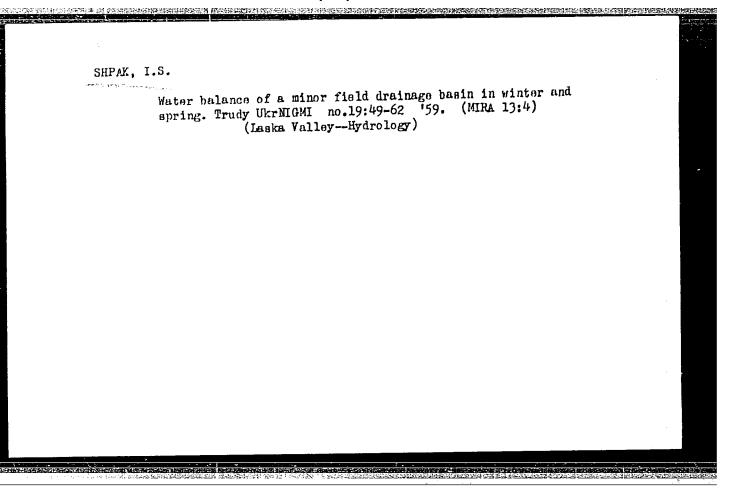


SHPAK, I.S.

Fluctuations of the water temperature in Tsimlyansk Reservoir.

Sbor. rab. po gidrol. no.1:62-66 59. (MIRA 15:2)

1. TSimlyanskaya nauchno-issledovateliskaya gidrometeoobservatoriya. (TSimlyansk Reservoir--Temperature)

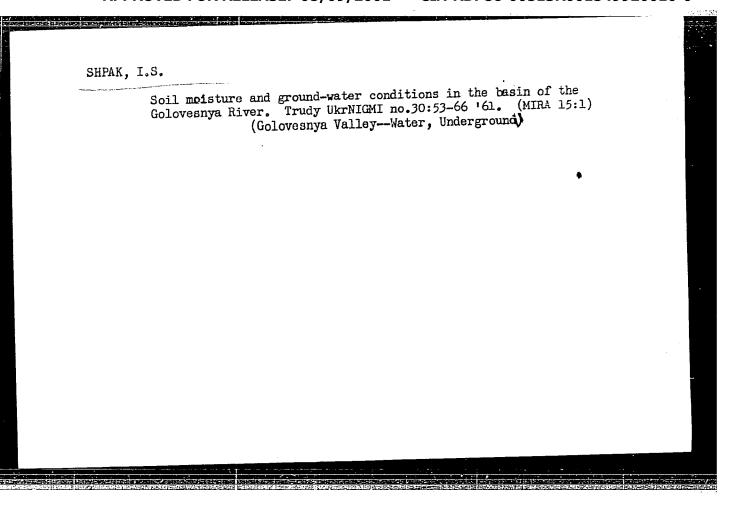


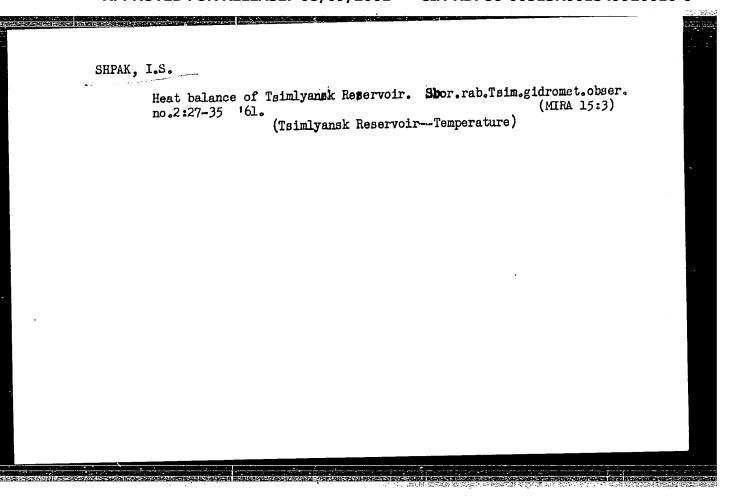
SHPAK, I.S.; KOSTYUCHENKO-PAVLOVA, M.M.

Accuracy of the single sample system in determining soil moisture by the gravimetric method. Pochvovedenie no.6: 107-111 Je '61. (MIRA 14:6)

1. Severo-ukrainskaya opornaya gidrogeologicheskaya stantsiya. (Soil moisture)

Water balance of a small field watershed. Trudy UkrNICHI
no.30:41-52 161. (MIRA 15:1)
(Desna Valley—Hydrology)





SHPAK, I.S.

Moisture variations in steppe soils during summer. Trudy UkrNICMI no.34:
39-44 162. (MIRA 15:7)

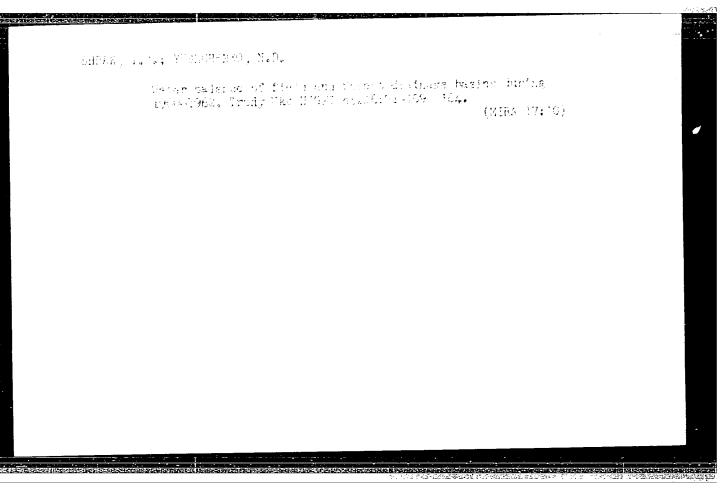
(Soil moisture) (Steppes)

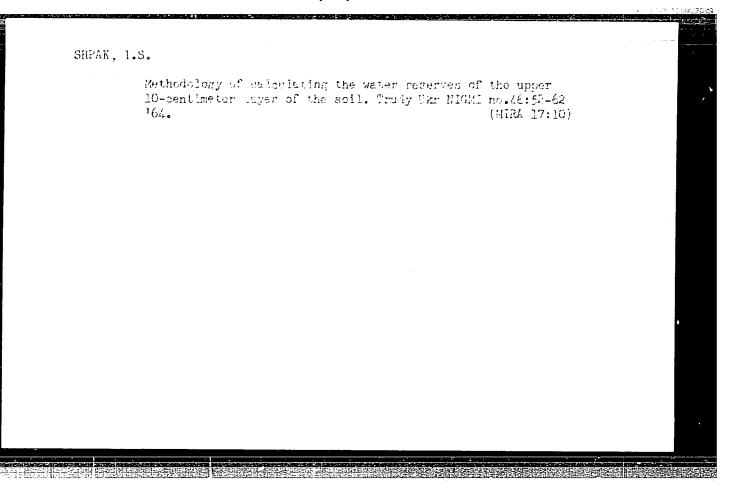
SHPAK, I.S.

Water balance of a small field and a forest drainage basin for the hydrological year 1959-60. Trudy UkrNIGMI no.34:24-38 *62.

(MIRA 15:7)

(Hydrology)





Serak, 1.°.

Methods for calculating the moisture supply in a lo-dentimeter soil layer. Formwowelshie mo.9:50-53 5 ltg.

(MIRA 18:10)

l. Parainskiy nauchno-destedows bullakiy gidrometeomologicheskly institut.

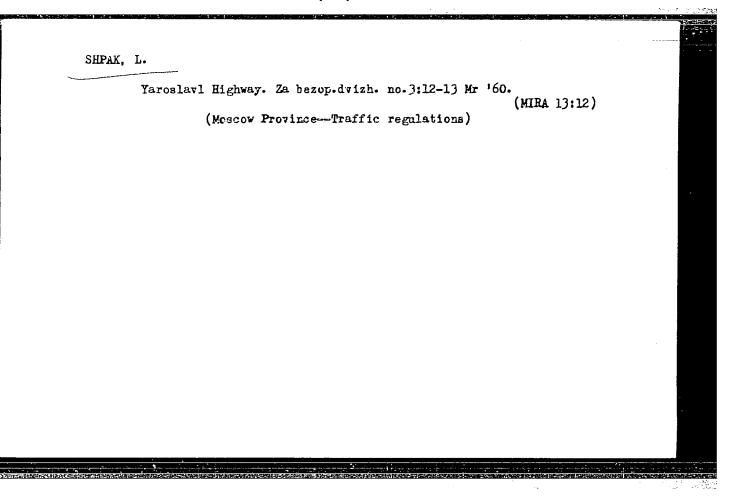
SHPAK, I.S.

Natural replenishment of groundwater resources. Trudy UkrNIGMI no.50:29-39 '65.

Closed depressions (saucer-shaped sinkholes, hollow depressions, etc.) in the Ukraine and their hydrologic role. Ibid.:60-69 (MIRA 18:11)

SHPAK, I.S.; ZAPOL'SKIY, I.A.

Water balance of the drained bottomland of the Trubezh River, 1960-1963. Trudy UkrNIGMI no.50:40-49 '65. (MIRA 18:11)



ZHIGAYLO, Ya.V.; SHPAK, L.I.

Effect of potassium addition on the rate of regeneration of a zinc-chromium catalyst. Khim. prom. no.8:591-593 Ag '63. (MIRA 16:12)

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18(5)

S07/128-59-9-20/25

AUTHOR:

Shpak L.V. and Gladkiy S.I., Engineers

TITLE:

Using Tall Oil in Producing Foundry Cores

PERIODICAL:

Liteynoye proizvodstvo, 1959, Mr 9, p 45 (USSR)

ABSTRACT:

Foundry cores used in serial production should be neither hygroscopic nor brittle. To impart them the required properties, the Zaporozh'ye Plant Kemmunar' has introduced a process of making the foundry cores out of a mixture of the fixing agent P with tall oil. The latter, a liquid of a dark-brown color, is a waste material of cellulose industry. Its moisture content is 0.6%; viscosity at 50°C - 17.7 (according to Englerl); specific gravity - 0.97; value of saponification - 206. For preparation of the mixture the tall oil, in quantity of 0.6%, is charged together with the fixing agent P into the runners, where it is mixed during 45-50 minutes. The volume of mixture is 250 litres; moisture - not over 3%; strength of cores - 0.1 to 0.25 kg/cm. The cores are dried at a temperature of 280° - 300°C in the course of 60-80 minutes. Tensile strength of dry cores is not under 10 kg/cm². The cores made on the basis of the fixing

Card 1/2

SOV/128-59-9-20/25

Using Tall Oil in Producing Foundry Cores

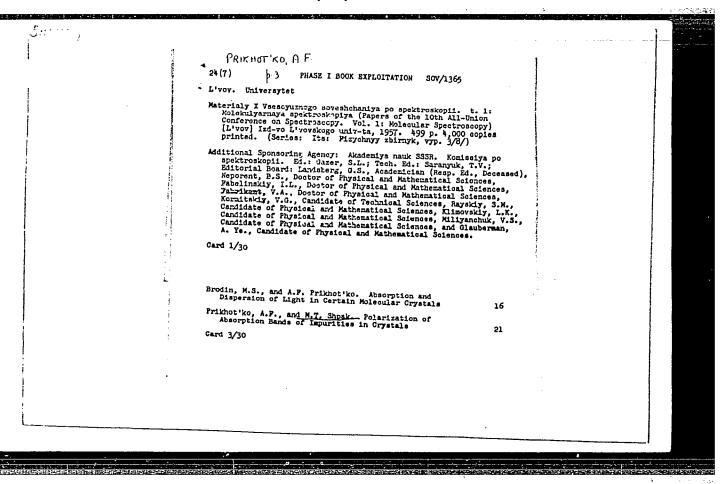
agent P with the addition of tall oil are "ar less hygroscopic and brittle than those prepared without oil. Addition of tall oil does not practically raise the cost of the mixture.

Card 2/2

SHPAK, M. T.

SHPAK, M. T. "Spectral Investigations in a Series of Polycyclic Hydro-carbons." Acad of Sci Ukrainian SSR. Inst of Physics. Kiev, 1956. (Dissertations for Degree of Candidate in Physicomathematical Science)

So: Knizhnaya Letopis', No. 17, 1956.



Sherrum analysis of the simplest polycyclic hydrocarbon solutions in a wide temperature range [with summary in English]. Ukr. fiz. zhur. 2 no.4:347-357 O-D '57. (KIRA 11:3)

1. Institut fiziki AN URSR. (Hydrocarbons--Spectra)

PRIKHOT'KO, A.F.; SHPAK, M.T.

Polarization of impurity absorption bands in crystals. Fiz. sbor. no.3:21-24 *57. (MIRA 11:8)

1. Institut fiziki AN USSR.
(Luminescence) (Absorption spectra)

51-4-1-3/26

AUTHORS: Prikhot'ko, A. F., and Shpak, M. T.

TIPLE: Absorption and Tuminescence of Stilbene Impurity in

Dibenzyl Crystals at 20.40K. (Pogloshcheniye i lyumin-

estsentsiya stil'bena kak primesi v kristallakh

dibenzila pri 20.4°K.)

PERIODICAL: Optika i Spektroskopiya, 1958, Vol.IV, Nr.1,

pp.17-29. (USSR)

ABSTRACT: The absorption bands of strongly absorbing crystals

(e.g. stilbene and tolane) are still comparatively

wide at liquid-hydrogen temperatures. When the absorb-

ing molecules are separated by molecules of a foreign

lattice (e.g. stilbene as an impurity in a dibenzyl

crystal) the absorption bands of the impurity are

sufficiently narrow to be called lines. Crystals with such impurities usually luminesce. The present paper

deals with absorption and luminescence of stilbene in

Card 1/7 dibenzyl crystals. The results were obtained using a

51-4 -1-3/26 Absorption and Luminescence of Stilbene Impurity in Dibenzyl Crystals at 20.40K.

> Hilger quartz spectrograph of high dispersive power. The absorption spectra were photographed in polarized The cell with the crystal was cooled to the temperature of liquid hydrogen, A hydrogen or a krypton lamp was used as a source, and an are spectrum between iron electrodes was used for calibration. crystal thicknesses in absorption studies were from several hundredths of a micron to 160 μ . The stilbene content was from about 0.001% to 3-5% by weight. To study luminescent spectra crystalline plates of 1 mm thickness were used. These spectra were obtained from the same surface of the crystal onto which the exciting Luminescence was excited by means of a light fell.

Card 2/7 mercury lamp. Crystals of dibenzyl possess intrinsic

51-.4-1-3/26 Absorption and Luminescence of Stilbene Impurity in Dibenzyl Crystals at 20.40K.

absorption in the region of 37000 cm^{-1} which consists of a large number of narrow bands and lines, some of which are strongly polarized. The stilbene impurity begins to absorb light from about 29500 cm⁻¹. shows the absorption spectra of dibenzyl monocrystals 160 μ thick with 0.04% of stilbene (Fig.la), 5 μ thick with 5% of stilbene (Fig.1b) and 2 μ thick with 5% stilbene (Fig.lv). All results in this figure and subsequentfigures and tables are quoted in Table 1 shows the results of measurements on a 160 μ thick dibenzyl crystal with about 0.04% of stilbene (Fig.la). The first column gives relative Card 3/7 intensities obtained visually. The second column

Absorption and Luminescence of Stilbene Impurity in Dibenzyl Crystals at 20,40K.

gives frequencies in cm⁻¹ measured with 1-2 cm⁻¹

precision for narrow lines, 3-5 cm⁻¹ for narrow
bands and 10-15 cm⁻¹ for wide bands. The third

column gives the wave-number differences with respect
to the 29904 cm⁻¹ line and similar information. The
fourth column contains remarks. Fig.2 shows the
luminescence spectrum of dibenzyl with 3-4% of stilbene
(5% according to the caption under Fig.2). All

measured luminescence lines of a crystal with 0.015%
of stilbene (Fig.3a) are given in Table 2. The
first column gives the relative intensities, the second the frequencies in cm⁻¹, the third - the frequency
differences with respect to the 29902 cm⁻¹ line, the
Card 4/7 fourth - similar frequency differences calculated with

Absorption and Luminescence of Stilbene Inpurity in Dibenzyl Crystals at 20.4 $^{\circ}$ K.

respect to other lines. The fifth column contains remarks. Fig.3 gives the luminescence spectra for dibenzyl crystals with 0.015% of stilbene (Fig.3a) and 5% of stilbene (Fig.3b). Analysis of the spectra yields the main absorption series due to intramolecular vibrations with frequencies: 203, 297, 402, 590, 745, 847, 978, 1249, 1333, 1427, 1538 and 1593 cm⁻¹. Each of these frequencies, except 402 and 590 cm⁻¹, is itself a beginning of a further series. In addition to the series of strong lines beginning at 29904 cm⁻¹, there is a second, weaker, series, which begins at 29840 cm⁻¹ (Fig.1b). Fig.1v gives the absorption spectrum of dibenzyl 2 m thick with 5% card 5/7 stilbene. Compared with Fig.1b, the absorption bands

Absorption and Luminescence of Stilbene Impurity in Dibenzyl Crystals at 20.40K.

in Fig.1v are slightly displaced, broadened, weakened and immersed in a strong background. The reason for such a change may lie in disorder in the orientation of stilbene molecules and formation of disordered aggregates on rapid crystallization. Comparison of the absorption and luminescence spectra shows that certain lines are either identical or very close to one another in both spectra. Table 3 shows that in a frequency interval of 200 cm⁻¹ no less than nine lines coincide. It is suggested that these lines and the short series occurring with them are due to nolecules of stilbene which are deformed in the dibenzyl lattice. There are 3 figures, 3 tables and 6 references, of which 5 are

Card 6/7 Russian and 1 German.

Absorption and Luminescence of Stilbene Impurity in Dibenzyl Crystals at 20.40K.

ASSOCIATION: Institute of Physics, Academy of Sciences of the Ukrainian SSR. (Institut fiziki, AN USSR.)

SUBLITTED: March 25, 1957.

AVAILABLE: Library of Congress.

1. Stilbene crystals-Impurities 2. Stilbene crystals-Absorption

Card 7/7

AUTHORS: Prikhot'ko, A. F., and Shpak, H. T.

TITIM: Absorption and Tuminescence of Stilbene Impurity in Tolane Crystals at 20.40K. (Pogloshcheniye i

lyuminestsentsiya stil'bena kak primesi v kristallakh

tolana pri 20.40K.)

PERIODICAL: Optika i Spektroskopiya, 1958, Vol. IV, Nr.1, pp.30-39. (USSR).

ABSTRACT: Absorption by stilbene molecules in the tolane crystal lattice differs in certain respects from abscrption by the same molecules in the dibenzyl lattice (see the preceding paper). The most important difference is in the polarization of certain absorption bands of stilbene in tolane which are rather similar to the absorption bands of tolane itself (Ref.2). Theory of this effect was developed by E.I. Rashba (Ref.3). To study this effect in detail, the technique used in

Card 1/5 Ref.2 was followed. A Hilger quartz spectrograph of

51-4-1-4/26

Absorption and Luminescence of Stilbene Impurity in Tolane Crystals at 20.40 K.

high dispersion was used. Fig.1 shows the absorption spectra of 160 μ thick tolane monocrystals with 0.4% Table 1 gives the absorption bands and of stilbene. lines of stilbene in tolane at 20.40K. The first four columns give the intensities of strong and weak lines shown in Fig.1. The fifth column gives the frequency in cm⁻¹, the sixth column gives the various frequency differences and the seventh column contains remarks. Table 2 gives some frequencies and intensities of the absorption spectra of tolane crystals with various amounts of stilbene. The impurity lines are found to be strongly polarized; each new transition to a Card 2/5 deformed or differently oriented molecule could depress

Absorption and lumine scence of Stilbene Impurity in Tolane Crystals at 20.40 K. 51-4 -1-4/26

Card 3/5

the remaining transitions and become the beginning for all series. The absorption bands remain narrow and sharp, although they are somewhat broadened on increase of stilbene concentration. From these results the fundamental frequencies of intramolecular vibrations of stilbene are determined. The general nature of luminescence of stilbene (1%) in tolane is shown in The spectrum begins with a pair of very weak Fig.2. lines of frequencies 29672 and 29620 cm⁻¹. of the luminescence spectrum is given in Fig. 3. Table 3 gives the luminescence of stilbene in tolane at 20.4 K. The first column gives a rough intensity value; the second column gives the frequency; third, fourth and fifth columns give various frequency

51-14-1-4/26 Absorption and Luminescence of Stilbene Impurity in Tolane Crystals at 20.40K.

> differences. If the three series, which begin at 29672, 29528 and 29468 cm^{-1} , are separated out from the luminescence spectrum, then the frequencies of intramolecular vibrations of stilbene in tolane are found to be close to the corresponding frequencies obtained from the Raman spectrum and from luminescence of stilbene in dibenzyl. Stilbene and tolane form a continuous series of solid solutions (Ref.6), since their lattices are practically identical. Introduction of stilbene molecules into the lattice sites produces "defects" which become absorption centres. accounts for the spectral peculiarities of tolane crystals with stilbene impurity which were described

Card 4/5 above. The authors note that tolane itself does not

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Absorption and Iuminescence of Stilbene Impurity in Tolane Crystals at 20.40K.

luminesce, but luminescence occurs only from the impurity levels. In the absorption and luminescence spectra of cadmium sulphide (Ref.5) the same peculiarities are observed as in tolane with stilbene. The difference between CdS and stilbene-tolane lies in the fact that the "defects" in the former are due to an excess of sulphur or cadmium, while in the latter the "defects" are stilbene molecules. There are 3 tables, 3 figures and 7 references, of which 6 are Russian and 1 French.

ASSOCIATION: Institute of Physics, Academy of Sciences of the Ukrainian SSR. (Institut fiziki AN USSR.)

SUBMITTED: March 25, 1957.

AVAILABLE: Library of Congress.

Card 5/5 1. Tolane crystals-Stilbene Absorption 2. Tolane crystals-

Stilbene-luminescence

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SOV/51-6-1-26/30

AUTHORS:

Prikhot'ko, A.F. and Shpak, M.T.

TITLE:

On the Origin of Certain Electron Transitions in Naphthalene Crystals (O proiskhozhdemii nekotorykh elektronnykh perekhodov v kristallakh

naftalina)

PERIODICAL: Optika i Spektroekopiya, 1959, Vol 6, Nr 1, pp 119-120 (USSR)

ABSTRACT &

Because of doubts about the purity of crystals in earlier work (Refs 2, 3) the authors re-investigated the spectra of naphthalene in order to check the origin of electron transitions in crystals. The authors used samples of naphthalene of various origins and purified by various methods: chemical, recrystallization from solution, sublimation and roze melting. It was found that the intensities of the absorption lines at 29941 and 31050 cm-1 varied considerably from sample to sample (the samples were monocrystalline, of 1 to 5 mm thickness). If the method of zone melting is used it is possible to obtain a rod of naphthalene from which layers of various purity can be cut. Such a rod was presented to the authors by V.I. Startsev and Yu.V. Naboykin. It was found that the 31060 cm line was virtually absent in sample cut from the region of the rod with the highest purity. On the other hand samples of low purity, also cut from this rod, had a strong polarized absorption hand at the latter, Frequency and an extension white

Card 1/2

APPROVED FOR RELEASE: 08/09/2001 CIA-RDP86-00513R001549920010-0"

SOV/51-6-1-26/30

On the Origin of Certain Electron Transitions in Naphthalene Crystals

only the absorption but also luminescence spectra were found to depend on the sample purity. The observed effects suggested presence of impurities in the crystals. This was confirmed by adding \$\beta\$-naphthol which increased the intensity of the 29941 cm⁻¹ line. Addition of \$\beta\$-methylaschthalens strengthened the 31060 cm⁻¹ line. Irradiation of naphthalene crystals with a beam of neutrons showed the presence of sulphur compounds in it. This irradiation and verification of the presence of sulphur compounds was carried out together with V.A. Batalia in the laboratory of M.V. Pasechnik. It is possible that one of these sulphur compounds in responsible for the 33441 cm⁻¹ line and all the other related weakly polarized absorption lines. There are 6 references, 4 of which are Soviet, 1 English and 1 mixed (Soviet and English).

SUBMITTED: May 29, 1958

Card 2/2

24,3500

68313

AUTHORS:

Shpak, M.T. and Sheka, Ye.F.

SOV/51-8-1-12/40

TITLE :

On the Intrinsic Luminescence of Crystalline Naphthalene

PERIODICAL: Optika i spektroskopiya, 1960, Vol 8, Nr 1, pp 66-72 (USSR)

ABSTRACT:

The authors investigated the luminescence spectrum of crystalline naphthalene lying in the region of fundamental absorption. The luminescence and absorption spectra were obtained at 20.4, 77, 200-210, 293 and 330-340°K. Medium-dispersion quartz spectrographs ISP-22 and Hilger E-2 were used. For the absorption spectra a krypton lamp GSVD-120 was used; luminescence was excited with 2530-2400 Å light from a mercury lamp SVDSh-1000. A Glan-Thomson prism was used as an analyser. Very pure samples of naphthalene were obtained by repeated zone purification (45 runs). Purity of the samples was judged by the absence of β -methyl naphthalene bands in the absorption spectrum. Fig 1 gives the luminescence spectra of naphthalene crystals at 20.40K. Fig 1 shows a luminescence band at 31480 ± 3 cm-1 weakly polarized along the a-direction (this is called an A-band). The A-band practically coincides with a strongly polarized Al-band at 31476 cm⁻¹ in the absorption spectrum (Ref 11). The A-band is more intense in purer samples. On increase of temperature to 770K

Card 1/3

APPROVED FOR RELEASE: 08/09/2001 CIA-RDP86-00513R001549920010-0"

68313

On the Intrinsic Luminescence of Crystalline Naphthalene

507/51-8-1-12/40

another band appears in the luminescence spectrum; it lies at 31623 ± 3 cm-1 (the B-band) and it is polarized in the b-direction At 200-210°K (Fig 3) the A- and B-bands are broader and partly depolarized (they coincide with the Al-and Bl-bands in the absorption spectrum at the same temperature). At 200-210°K a very weak luminescence band (the I-band) appears at 31900 cm-1. On increase of temperature to 293°K the A- and B-bands are depolarized still further (the spectrum is practically unaffected by a 900 rotation of the Glan-Thomson prism) and the intensity of the I-band rises (Fig 4a). The I-band is also completely depolarized. At 330-340°K the A-, Band I-hands become stronger and another band appears in the spectrum (the II-band), which lies in the region of the A2 and B2 absorption bands (Ref 11). Rise of temperature depolarizes also the absorption cands which are strongly polarized at 20.40% (Fig 5). At 770% a new band (the III-band) is observed in the absorption spectra of thick maphthalene crystals; this band lies at 31100 cm-1 and its intensity

Card 2/3

"APPROVED FOR RELEASE: 08/09/2001

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68313

On the Intrinsic Luminescence of Crystalline Naphthalene

SCV/51-8-1-12/40

rises on increase of temperature. At 330-340°K no further changes are observed in the absorption spectrum. The authors show that the results point to exciton nature of luminescence of very pure naphthalene crystals in the region of fundamental absorption. Acknowledgment is made to A.F. Prikhot'ko for his advice. There are 5 figures and 14 references, 9 of which are Soviet, 3 English and 2 German.

SUBMITTED:

June 8, 1959

Card 3/3

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AUTHORS :

TITLE 8

Shpak, M.T. and Sheka, Ye.F.

The Effect of Impurities on Luminescence of Crystalline Naphthalene

PERIODICAL: Optika i spektroskopiya, 1960, Vol 9, Nr 1, pp 57-63 (USSR)

ABSTRACT 8

Earlier Prikhot'ko and Shpak (Ref 1) showed that luminescence of crystalline naphthalene (Ref 2) is due to a β -methylnaphthalene impurity. It was also reported that crystalline naphthalene contains small amounts of d-naphthol, \$-naphthol, thionaphthene and other The present paper reports an investigation of the substances. effect of temperature and solute (impurity) concentration on the luminescence and absorption spectra of solid solutions of A-methylnaphthalene, a-methylnaphthalene, A-naphthol and a-naphthol in crystalline naphthalene. The experimental technique was the same The experiments were carried out as described earlier (Ref 3). at 20.4, 77 and 200°K. The results (Figs 1-5 and Tables 1-4) showed

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The Effect of Impurities on Luminescence of Crystalline Naphthalene

that solute molecules produced two types of luminescence centres in the naphthalane lattice. Acknowledgments are made to A.F. Prikhot'ko for his advice and to a graduate of the Kiyev State University, Ye.G. Ruseva. There are 5 figures, 4 tables and 11 references, 8 of which are Soviet, 2 English and 1 German.

SUEMITTED: September 18, 1959

Card 2/2

SHPAK, M.T.; SHEKA, Ye.F.

Luminescence of crystalline naphthalene containing minor admixtures. Izv.AN SSSR.Ser.fiz. 24 no.5:553-555 by 160.
(MIRA 13:5)

1. Institut fiziki AN USSR.
(Naphthalene--Optical properties)

9,4160 (also 1137, 1395)

S/048/61/025/001/002/031 B029/B067

AUTHOR:

Shpak, M. T.

TITLE:

The role of impurities in the luminescence of molecular

crystals at low temperatures

PERIODICAL:

Izvestiya Akademii nauk SSSR. Seriya fizicheskaya, v. 25,

no. 1, 1961, 13-19

TEXT: This is a review on studies made at the Institut fiziki AN USSR (Institute of Physics of the AS UkrSSR) which covered the following subjects: a) the simplest aromatic compounds (diphenyl, terphenyl, quaterphenyl, etc.); b) compounds with benzene nuclei which are bound by carbon atoms not contained in the ring (dibenzyl, stilbene, etc); compounds with condensed benzene nuclei (naphthalene, anthracene, phenantrene, etc.); d) derivatives and deuterosubstitutes of some of these substances. Substances of the same class of hydrocarbons were used as impurities. The studies were made with monocrystals. Following are the most important results obtained: 1) General properties of the spectra: The absorption bands of strongly absorbing substances (stilbene, tolane, anthracene, etc.)

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are relatively broad even at hydrogen temperatures (50-100 $\rm cm^{-1}$). On the other hand, the impurity spectra of such substances consist of very narrow, weakly polarized bands. One and the same molecule has similar frequencies in different solvents. 2) Nature of impurity centers. Usually, the impurity crystals have multiplet-type spectra. The spectra of the solutions of monosubstitutes of napthalene in naphthalene consist of two equal series of bands which are mutually shifted by constant quantities $\Delta \, {f v}.$ The phenomena predicted by the quantum-mechanical theory of absorption and luminescence of molecular crystals should be studied experimentally. 3) The role of defects. In the spectra of several crystals near the longwave edge of self-absorption, very weak bands of largely varying intensity and position lie in an interval of sometimes several hundred cm⁻¹. These bands are in resonance with the shortest-wave bands of the corresponding luminescence spectra. These characteristic features of spectra are related to defects in real molecular crystals. 4) Mirror symmetry. The bands in the impurity absorption and luminescence spectra are almost always surrounded by groups of satellite bands of different intensity. These satellite bands are mirrorsymmetrically arranged with respect to the band of a pure electron transition. Hence, the satellite bands are assumed to belong to the same type of Card 2/3

89236

The role of impurities in the

S/048/61/025/001/002/031 B029/B067

discrete impurity centers and to be related to the interaction between intermolecular transitions and vibrations of the solvent lattice. The fifth section deals with the polarization of bands in absorption spectra. First, strongly polarized bands were observed in the absorption spectrum of stilbene in tolane. With increased impurity concentration, the bands in their absorption spectra are usually blurred and widened. The problem of the origin of strongly polarized bands in the absorption spectra of impurities in organic crystals has as yet not been definitely solved. The introduction of a second impurity into a molecular crystal strongly changes its spectrum considerably. The material discussed in this paper needs further examination. This is the reproduction of a lecture read at the Ninth Conference on Luminescence (Crystal Phosphors), Kiyev, June 20-25, 1960. There are 34 references: 30 Soviet-bloc and 4 non-Soviet-bloc.

ASSOCIATION: Institut fiziki Akademii nauk USSR (Institute of Physics of the Arademy of Sciences UkrSSR)

Card 3/3

.7

SHPAK, M.T.; SOLOV'YEV, A.V. [Solovyov, A.V.]; SHEREMET, N.I.;

DMITHERKO, I.P. [Dmytenko, O.P.]

Spectra investigation of chemical transformations in crystalline triphenylmethane. Ukr.fiz.zhur. 7 no.4:422-429 Ap '62.

(MIRA 15:8)

1. Institut fiziki AN UkrSSR, g. Kiyev.

(Methane) (Chemical reactions)

SHPAK, M.T.; SOLOV'YEV, A.V.; SHEREMET, N.I.

Nature of the luminescence spectra of crystalline benzene at low temperatures. Opt.i spektr. 13 no.52694-700 N '62.

(Benzene crystals—Spectra)

(Benzene crystals—Spectra)

"APPROVED FOR RELEASE: 08/09/2001

CIA-RDP86-00513R001549920010-0

L 11169-63 EPF(c)/EPR/EWP(j)/EWT(1)/EWT(m)/BDS/EEC(b)-2--AFFTC/ASD/SSD--Pr-h/Ps-h/Pc-h--IJP(C)/RM/WW

ACCESSION NR: AP3002792

\$/0051/63/014/006/0816/0819

AUTHOR: Shpak, M. T.; Sheremet, N. I.

80

TITLE: On the nature of the luminescence of crystalline anthracene at low

temperatures

SOURCE: Optika i spektroskopiya, v. 14, no. 6, 1963, 816-819

TOPIC TAGS: anthracene, luminescence

ABSTRACT: Although there have been many investigations of the luminescence of anthracene, the nature of its luminescence is still obscure. It has been shown that the luminescence at low temperatures is not due to radiative annihilation of excitons in the main lattice. On the other hand, it has also recently been shown that the luminescence spectra of naphthalene and benzene have certain characteristics that may be associated with exciton effects and that should be exhibited by other molecular crystals. The paper gives the results of investigation of the luminescence of zone refined (40 zone) crystal anthracene in polarized light at 20.4, 77 and 290°K. Excitation was realized by the light from a DRSh-250 mercury discharge tube opassing through a Woods filter; the spectra were observed by ISP-22°C and DFS-130 spectrographs of The spectra along two crystal axes are reproduced and

Card 1/2

L 11169-63

ACCESSION NR: AP3002792

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described. Analysis of the luminescence data in conjunction with the literature data on the absorption spectrum shows that the first bands of both components partially overlap which indicates that they correspond to pure electronic transitions from exciton bands in the crystal. This is substantiated by the temperature dependence of the intensity and width of these bands. Three other groups of bands were distinguished in the luminescence spectrum at 20.4°K: one is attributed to an impurity; the second is associated with transitions in distorted host molecules located near lattice defects; the third is tentatively attributed to transitions from the lowest exciton band to a set of ground-state electronic-vibrational levels. Orig. art. has: 1 figure.

ASSOCIATION: none

SUBMITTED: 10Dec62

DATE ACOD: 15Jul63

ENCL: 00

SUB CODE: 00

NO REF SOV: Oll

OTHER: 005

16/wm

Card 2/2

MALYKHINA, N.N.; SHPAK, M.T.

Spectroscopic study of solutions of stilbene in n-octane in polarized light at 20.4°K. Opt. i spektr. 14 no.6:829-831 Je '63. (MIRA 16:8)

(Stilbene-Spectra)

L 10164-63 EFF(c)/EWT(1)/EWT(m)/HDS-AFFTC/ASD/SSD--Pr-4-RM/WW/MAY

ACCESSION NR: AP3000308

s/0048/63/027/005/0596/0605

AUTHOR: Broude, V. L.; Sheka, Ye. F.; Shpak, M. T.

65 63

TITLE: Exciton luminescence of molecular crystals [Report: Eleventh Conference on Luminescence held in Minsk 10-15 Sept. 1962]

SOURCE: Izvestiya AN SSR. Seriya fizicheskaya, v. 27, no. 5, 1963, 596-605

TOPIC TAGS: molecular crystals, naphthalene, excitons, energy bands, luminescence

ABSTRACT: Observation, interpretation and indentification of exciton luminescence of molecular crystals is often difficult. Analysis of luminescence data for molecular crystals shows that in many cases the observed luminescence is not due to excitons, but to impurities or defects. The specific case of low temperature luminescence of naphthalene crystals is examined and attributions are made for the different luminescence bands. Exciton series are identified. The general energy structure of naphthalene crystals can be explained in the framework of a molecular diagram, but certain distinctive features of the

Card1/2

L 10164-63

ACCESSION NR: AP3000308

naphthalene luminescence spectrum are specifically associated with the exciton character of its crystal structure. It is hoped that the approach to analysis of these features may help elucidate the structure of the exciton bands (zones) in this crystal and can be extended to the study of other molecular crystals. Orig. art. has: 6 figures and 1 table.

ASSOCIATION: Institut fiziki Akademii nauk USSR (Institute of Physics, Academy

of Sciences, USSR)

SUBMITTED: CO

DATE ACQ: 12Jun63

ENCL: 00

SUB CODE: PH

NR REF SOV: 013

OTHER: 007

Card 2/2

"APPROVED FOR RELEASE: 08/09/2001

CIA-RDP86-00513R001549920010-0

IJP(c) GG/RM EWI(1)/EWI(m)/EWP(j)/IL 34000-66 UR/0058/65/000/012/D042/D042 SOURCE CODE: ACC NR: AR6017247 61 AUTHORS: Malykhina, N. N.; Shpak, M. T. \mathcal{B} TITLE: Low-temperature spectral investigations of the stilbene-tolar impurity crystals SOURCE: Ref. zh. Fizika, Abs. 12D355 REF SOURCE: Tr. Komis. po spektroskopii. AN SSSR, t. 3, vyp. 1, 1964, 173-477 [probably should be 473-477] TOPIC TAGS: crystal impurity, absorption spectrum, luminescence spectrum, light polarization, impurity center, light excitation ABSTRACT: The absorption and luminescence spectra of single crystal tolan with stilbehe molecules enter into the tolan lattice in several ways, forming impurity centers of different natures. The impurity concentration influences the intensity distribution in the luminescence spectra of different centers. The sharp difference in the polarization properties of the bands corresponding to the pure-electronic and electronic vibrational transitions in one and the same impurity center is qualitatively explained by means of a theory which takes into account the delocalization of the excitations of the impurity molecules. [Translation of abstract]. SUB CODE: 20 Card 1/1

MALYKHINA, N.N. [Malykhina, N.N.] ; SHPAK, M.T.

Spectral study of a mixed bibenzyl ~ stilbene crystal at 20.42 and 4.20%. Ukr. Fiz. zhur. 9 no.2:172-178 F'64 (MIRA 17:7)

1. Institut fiziki AN UkrSSR, Klyev.

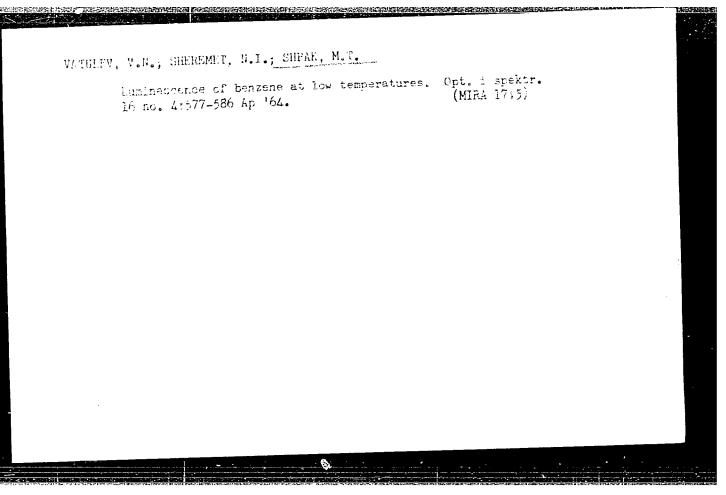
MALYKHINA, N.N. [Malykhina, N.K.]; SHFAK, M.T.

Effect of impurities on the absorption of crystalline
naphthalene and its deuterosubstituted compounds. Ukr.
naphthalene and its deuterosubstituted compounds. Ukr.

fiz. zhur. 9 no.9:991-1000 S '64.

1. Institut fiziki AN UkrSSR, Kiyev.

APPROVED FOR RELEASE: 08/09/2001 CIA-RDP86-00513R001549920010-0"



5/0051/64/017/002/0235/0243

AP4043011

AUTHORS: Maly*khina, N. N.; Shpak, M. T.

TITLE: Polarized-light investigations of the spectra of a crystalline solution of stilbene in tolane

SOURCE: Optika i spektroskopiya, v. 17, no. 2, 1964, 235-243

TOPIC TAGS: light polarization, absorption spectrum, luminescence spectrum, single crystal, thin film, impurity center, electronic transition

ABSTRACT: Results are reported of an experimental investigation of the absorption and luminescence spectra of the stilbene-tolane system, made in polarized light at temperatures 20.4 and 4.2K. The tolane used was prepared by zone melting to increase its purity. Most tests were made on single-crystal thin films prepared by sublimation. Luminescence was excited by a mercury lamp and the

1/3

ACCESSION NR:

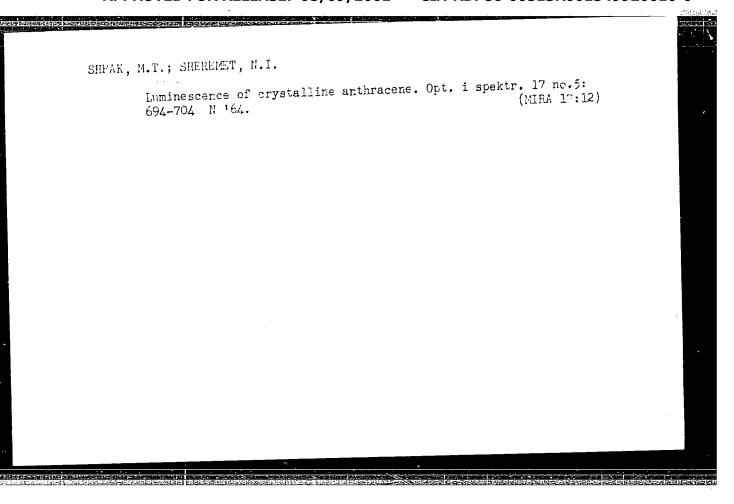
ACCESSION NR: AP4043011

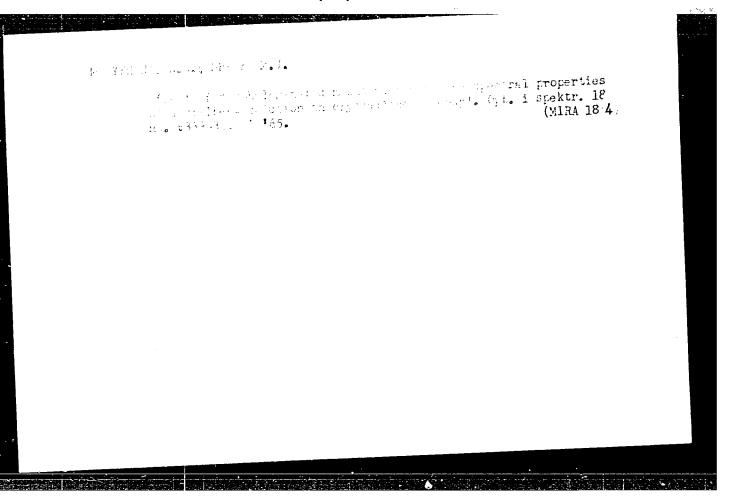
spectral sections were separated either with a monochromator or with a glass filter. A Glan-Thomson prism was used as a polarized-light analyzer, and most of the measurements were made with a DFC-13 spectrograph with resolution 4 Å/mm. The possible arrangement of the stilbene molecules, which formed various types of impurity centers in the tolane lattice, is analyzed in the approximation of the oriented-gas model. It is shown that the concentration changes in the luminescence spectra can be related to the occurrence of an interaction between the impurity centers of different types as the impurity concentration is increased. The polarization properties of the absorption and luminescence spectra of the investigated system are found to differ greatly, and contain bands that can be set in correspondence with combinations of a purely electronic transition in the stilbene molecule and the vibration frequencies of the tolane. Orig. art. has: 4 figures and 2 tables.

ASSOCIATION: None

2/3

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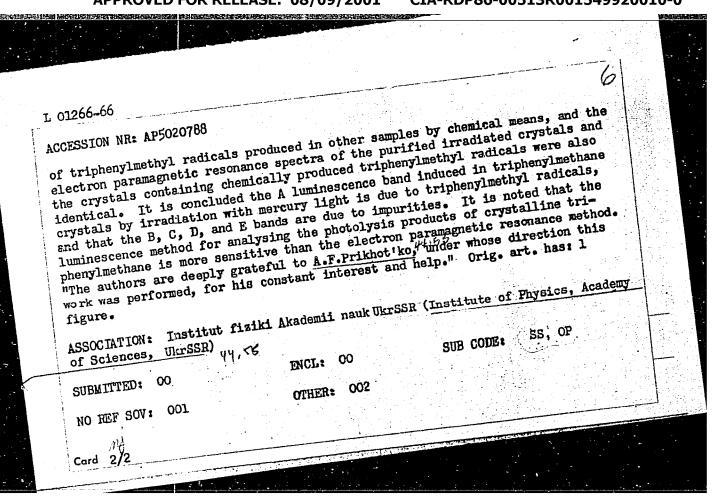
APPROVED FOR RELEASE: 08/09/2001 CIA-RDP86-00513R001549920010-0"

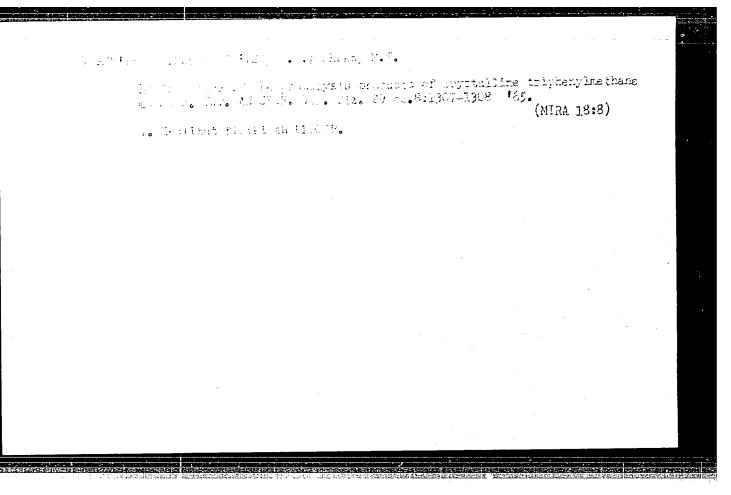
EWT(m)/EPF(c)/EWP(j) RM L 01265-66 UR/0048/65/029/008/1304/1306 ACCESSION NR: AP5020787 AUTHOR: Lisovenko, V. A.; Shpak, M. T. TITLE: Effect of imperfections of the crystal lattice on the luminescence proper ties of anthracene (Report, 13th Conference on Luminescence held in Khar'kov 41,55 25 June to 1 July 1964/ SOURCE: AN SSSR. Izvestiya. Seriya fizicheskaya, v. 29, no. 8, 1965, 1304-1306, and insert facing p. 1304 TOPIC TAGS: anthracene, absorption spectrum, luminescence spectrum, crystal deformation, crystal lattice defect, crystal impurity ABSTRACT: This paper reports a continuation of previous work (M.T. Shpak and N.I. Sheremet, Optika i spektroskopiya, 17, 694, 1964) on the effect of deformation on the exciton luminescence spectrum of anthracene. Lattice defects were introduced into the anthracene crystals by doping with molecules that are comparable in size to or larger than anthracene molecules, and by thermal deformation. The absorption and luminescence spectra were observed at 20.4°K. Doping with large molecules led to the appearance of a new series in the spectrum, similar to the room Card 1/2

APPROVED FOR RELEASE: 08/09/2001 CIA-RDP86-00513R001549920010-0"

temperature luminescence spectrum of anthracene and shifted by 150-160 cm ⁻¹ to the long wavelength side from the beginning of the exciton luminescence. Doping anthracene with 2-methylanthracene gave rise to a spectrum coinciding with that which has been previously ascribed to the luminescence spectrum of deformed anthracene molecules (A.F.Prikhot'ko and I.Ya.Fugol', Optika i spektroskopiya, 4, 335, 1958). Mechanical stresses were introduced by depositing thin pure anthracene crystals on substrates of different thermal expansion coefficients and subsequently cooling them. This treatment led to a shift of the luminescence spectrum by 20 cm ⁻¹ toward the short wavelengths, to the appearance of an intense continuous background, and to the appearance of new "deformation bands" near 24840 and 24880 cm ⁻¹ . These phenomena were due to irreversible plastic deformation; the deformation bands could be produced by other mechanical means. The results are discussed very briefly. The intensity of the deformation bands decreased very rapidly with increasing temperature, and at 770k the bands disappeared entirely; the reason for this is not understood. Orig. Art. has: 3 figures. ASSOCIATION: none SUBMITTED: 00 ENCL: 00 SUB CODE: SS, OP	L 01265-66	•		and the second section of the section of		
temperature luminescence spectrum of anthracene and shifted by 150-160 cm ⁻¹ to the long wavelength side from the beginning of the exciton luminescence. Doping anthracene with 2-methylanthracene gave rise to a spectrum coinciding with that which has been previously ascribed to the luminescence spectrum of deformed anthracene molecules (A.F.Prikhot'ko and I.Ya.Fugol', Optika i spektroskopiya, 4, 335, 1958). Mechanical stresses were introduced by depositing thin pure anthracene crystals on substrates of different thermal expansion coefficients and subsequently cooling them. This treatment led to a shift of the luminescence spectrum by 20 cm ⁻¹ toward the short wavelengths, to the appearance of an intense continuous background, and to the appearance of new "deformation bands" near 24840 and 24880 cm ⁻¹ . These phenomena were due to irreversible plastic deformation; the deformation bands could be produced by other mechanical reams. The results are discussed very briefly. The intensity of the deformation bands decreased very rapidly with increasing temperature, and at 770K the bands disappeared entirely; the reason for this is not understood. Orig. Art. has: 3 figures. ASSOCIATION: none SUBMITTED: 00 ENCL: 00 SUE CODE: SS, OP		37			0	
SUBMITTED: 00 ENCL: 00 SUB CODE: SS, OP	emperature luminescer ong wavelength side in thracene with 2-meth which has been pravious racene molecules (A.F. 1958). Mechanical starts crystals on substrates	from the beginning of hylanthracene gave ris usly ascribed to the l .Prikhot'ko and I.Ya.F resses were introduced s of different thermal	e to a spectrum co uminescence spectrugol', Optika i sp by depositing the expansion coeffi-	pinciding with rum of deform the pektroskopiya in pure anthrocients and supercence spect	th that ned anth- i, 4, 335, racene ubsequent- crum by	
SUBMITTED: 00	20 cm ⁻¹ toward the shoackground, and to the cm ⁻¹ . These phenomen tion bands could be pased very briefly. The	ort wavelengths, to the eappearance of new "ca were due to irrevers roduced by other mecha e intensity of the definitions, and at 77°K the	le appearance of a deformation bands ⁿ vible plastic defo anical reans. The formation bands de ne bands disappear	near 24840 a reation; the results are creased very	and 24880 deforma- discus- rapidly	
	20 cm ⁻¹ toward the sheackground, and to the cm ⁻¹ . These phenomen tion bands could be part of the p	ort wavelengths, to the e appearance of new "ca were due to irrevers roduced by other mechase intensity of the deirature, and at 77°K the nderstood. Orig. Art.	deformation bands ⁿ dible plastic defo anical reans. The formation bands de ne bands disappear has: 3 figures.	near 24840 a reation; the results are creased very ed entirely;	and 24880 deforma- discus- rapidly the rea-	

L01266-66 EWT(1)/EWT(m)/EPF(c)/EWP(j) IJP(c) ACCESSION NR: AP5020788 AUTHOR: Vorob'yev, V. P ; Solov'yev, A. V.; Luminescence of the products of photolysis of crystalline triphenylmethane at 200k Report, 13th Conference on Luminescence held in Khar'kov 25 June SOURCE: AN SSSR. Izvestiya. Seriya fizicheskaya, v. 29, no. 8, 1965, 1307-1308 TOPIC TAGS: photolysis, organic crystal, free radical, luminescence analysis, ABSTRACT: Two of the authors and collaborators have previously found that irradiation of triphenylmethane crystals at 20°K with unfiltered light from a mercury arc gives rise to five new bands in the luminescence spectrum, which they have called the A, B, C, D, and E bands, in order of increasing wavelength (Ukr. fiz. zh., 7, 422, 1962). The present paper reports results of an examination of initially very pure triphenylmethane crystals which have been further purified by zone refining. After irradiation with mercury are light these crystals showed only the A band, the B, C, D, and E bands being absent. The A luminescence band of the purified irradiated crystals was identical with the luminescence spectrum





"APPROVED FOR RELEASE: 08/09/2001

CIA-RDP86-00513R001549920010-0

L 28450-66 FBD/FWT(1)/FWT(m)/EEC(k)-2/T/EWP(k)/EWP(e) LIP(c) WG/WH ACC NR. AP6016052 SOURCE CODE: UR/0185/66/011/005/0567/0569

AUTHOR: Tykhonov, Ye. O.; Shpak, M. T.

ORG: Institute of Physics, AN UkrSSR, Kiev (Instytut fizyky AN URSR)

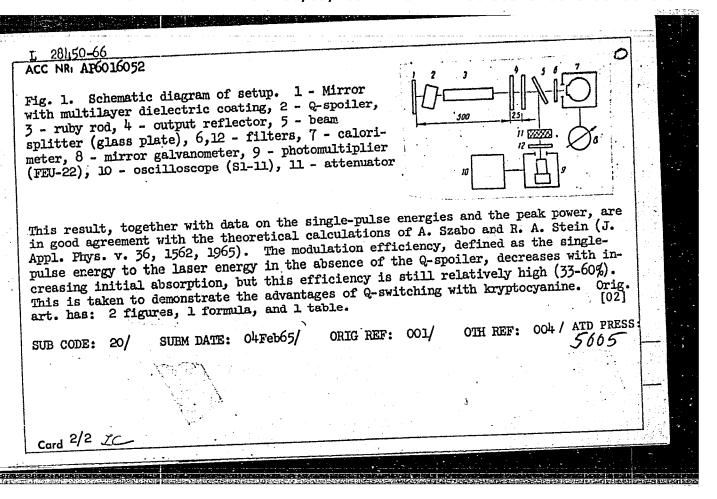
TITLE: Certain characteristics of ruby laser with passive Q switching

SOURCE: Ukrayins'kyy fizychnyy zhurnal, v. 11, no. 5, 1966, 567-569

TOPIC TAGS: ruby laser, resonator Q factor, laser modulation

ABSTRACT: The authors have experimentally determined the energy and frequency parameters of single pulses of a ruby laser with passive Q-spoiler based on a kryptocyanine solution. The solution was kept in a cell with optically finished surfaces and placed in the resonator close to the mirror with 99.5% reflectivity (Fig. 1). To prevent generation in the secondary resonator made up by the cell and the exit mirror, the cell was inclined to the resonator axis. The ruby used was 120 mm long and 13 mm in diameter, with matte surface. The energy was measured by a calorimeter method, and the system used to record the single pulses ensured a time resolution up to 4 nsec. The output reflector was a Fabry-Perot etalon with two plane-parallel quartz plates. Its energy reflection coefficient was an oscillating function of the wavelength and its maximum value was ~55%. Such a resonator yielded a single pulse with peak power ~30 MW and width ~30 nsec. A study of the oscillograms of the single pulses for different values of initial absorption showed that a considerable decrease takes place in the duration of the generated pulse with increasing initial absorption.

Card 1/2



EWT(m)/EWP(j) IJP(c) RM/JW/FDN L 38483-66 ACC NR SOURCE CODE: UR/0058/65/000/012/D042/D042 AR6017246 Vatulev, V. N.; Sheremet, N. I.; Shpak, M. T. AUTHOR: Spectral investigation of crystalline benzene at low tempera-TITLE: tures 1 SOURCE: Ref. zh. Fizika. Abs. 12D350 REF SOURCE: Tr. Komis. po spektroskopii. AN SSSR, t. 3, vyp. 1, 1964, 468-472 TOPIC TAGS: absorption spectrum, luminescence spectrum, exciton zone, crystalline benzene ABSTRACT: The luminescence and absorption spectra of crystalline benzene were investigated at 20.4 and 4.2K. The temperature relationships in luminescence spectra were evaluated. Preliminary information on the structure of benzene exciton zones, including their width and effective-mass characteristics, were obtained on the basis of a qualitative analysis of the shape of bands corresponding to transitions from the exciton zones to the basic vibration level. [Translation of [KP] abstract] SUBM DATE: none/ SUB CODE: 20/

	AN6030892	(N)		UR/9034/66/000/0		
AUTHOR:	Shpak, N. (Candidate of med	ical sciences; Od	essa)		
ORG: n	one					
TITLE:	Toxoplasmosi	s and optical pa	thology			
SOURCE:	Meditsinska	ya gazeta, 16 Se	p 66, p. 3, col.	2-5		
TOPIC T	AGS: parasit	ology, animal pa	rasite, opthalmol	.084, fissue liceau	, drug treatment	4
ABSTRAC	T: Toxoplasm	mosis has been id	lentified as one of diseases and the	of the most common cause of 36% of		
	all cases	of uveitis. Th	ne congenital for carriers (with no	m of this parasitie symptoms) to their e of inflammation	r :	
	occurs (in either the dis	seased or healtny tis or neuroretin	itis. Treatment w	ith	
	Chloridi	n and Sulfadimez:	ine reduces infla Thodies in the b	nmation and lowers lood. The frequent	t '	-
	localiza	tion of this disc	ease in the eye 1 ontibodies in acu	eous humor as com-		_
	pared wi	th blood serum.	Toxopiasmin can	be used for diagno	77 7 <u>.</u>	
	/2					

NOITHEATA, M.S. [Kome'ka, M.S.], kand, teldin, nauk; SHPAK, M.A.

132foct of coloring orides on the whiteress of kaolin,
1.ch. prom. no.4:20-22 (1-D *65. (MERA 19:1)

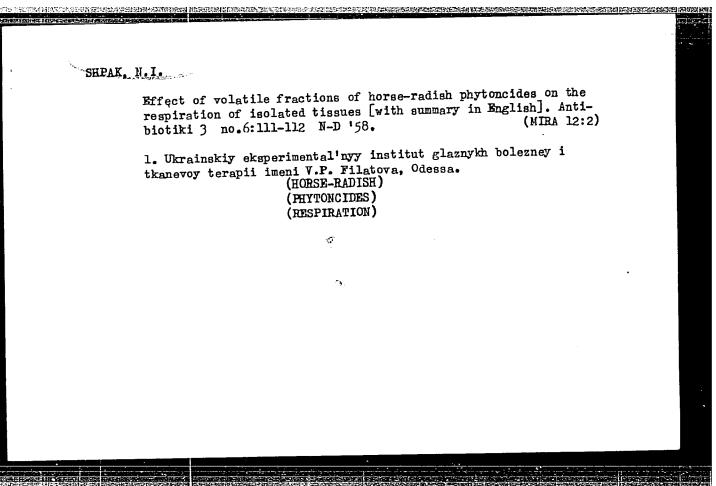
KOMSKAYA, M.S. [Koms'ka, M.S.], kand.tekhn.nauk; SHKOL'NIK, A.Ya. [Shkol'nyk, H.IA.]; SHPAK, N.A. [Shpak, N.P.]; YATSUNOVA, S.Ye. [IAtsunova, S.IU.]

Method for the regulation of the addition of electrolytes to porcelain slips. Leh.prom. no.1:63-66 Ja-Mr. 164. (MIRA 19:1)

CUPAY, N. I. -- With Conservation of Tissue for Keratoplasty and Tissue
Thorapy Using Cartain Propervatives of Biological Inthin," Classe
State Necteal Inch inend h. I. Firegov. Classa, 1995. (Dissertation
for the Tegree of Candidate in Nedical Sciences)

SC: Emishnaya letopis', %c %, 1996

SHPAK, N.I. New method of preserving soft animal tissues by using horse-radish phytoncides. Zool.zhur.35 no.3:467-468 Mr '56. (MIRA 9:7) 1.Ukrainskiy eksperimental'nyy institut glaznykh bolezney imeni V.P.Filatova. (Zoological specimens--Collection and preservation)(Phytoncides) (Horse-radish)



SHPAK, N.I., kand. med. nauk Preservation of eyes for corneal transplantations. Uch. zap. UEIGB 4: (MIRA 12:6)

91-96 '58.

1. Ukrainskiy eksperimental'nyy institut glaznykh bolezney i tkanevoy terapii imeni akademika V.P. Filatova. (EYE BANKS)

Toxoplasmosis and its manifestation in eye diseases. Oft. zhur.
16 no.2:102-106 '61.

1. Iz Ukrainskogo nauchno-issled. eksperimental 'nogo instituta
glaznykh bolezney i tkanevoy terapii imeni akademika V.P.Filatova
(direktor - prof. N.A.Puchkovskaya).

(TOXOPLASMOSIS)

(EYE—DISEASES AND DEFECTS)

SHPAK, N.I., kand.med.nauk

Diagnosis and treatment of congenital and acquired toxoplasmosis of the eye. Oft. zhur. 16 no.8:451-455 '61. (MIRA 15'4)

l. Iz Ukrainskogo nauchno-issledovatel skogo eksperimental nogo instituta glaznykh bolezney i tkanevoy terapii imeni akademika V.P.Filatova (direktor - prof. N.A.Puchkovskaya).

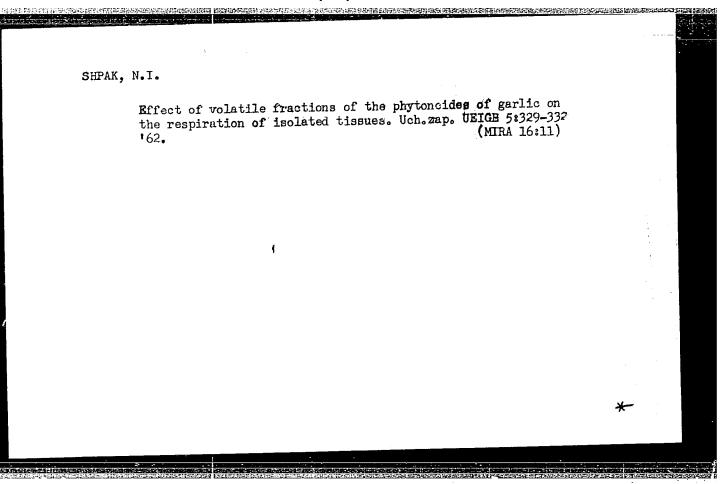
(TOXOPLASMOSIS)

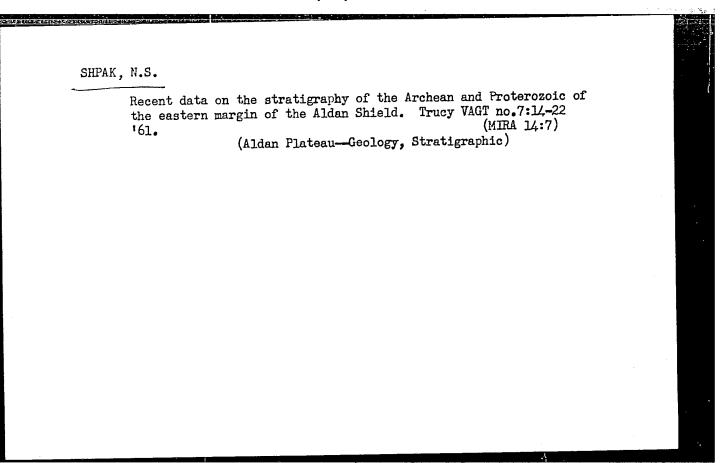
PANFILOVA, G. V., kand. med. nauk; SHPAK, N. I., kand. med. nauk

X-ray examinations in congenital toxoplasmosis. Oft. zhur. no.2: 82-87 162. (MIRA 15:4)

l. Iz Ukrainskogo nauchno-issledovatel'skogo eksperimental'nogo instituta glaznykh bolezney i tkanevoy terapii im. akad. V. P. Filatova (direktor - chlen-korrespondent AMN SSSR prof. N. A. Puchkovskaya)

(TOXOPLASMOSIS) (EYE__DISEASES AND DEFECTS)





Early Sinian subvolcanic formations of the southeastern Aldan
Early Sinian subvolcanic formations of the southeastern Aldan
Shield. Trudy VAGT no.7: 52-65 '61. (MIRA 14:7)
(Aldan Plateau—Geology, Stratigraphic)

ALEKSEYEV, V.R.; GAVRILOVA, Z.S.; KALIHULIN, S.M.; MORALEV, V.M.; HUZHNOV, S.V.; SHPAK, N.S.

Problem of the ancient rare metal placers of the eastern part of the Aldan Plateau. Dokl.AN SSSR 144 no.2:409-411 My 162. (MTRA 15:5)

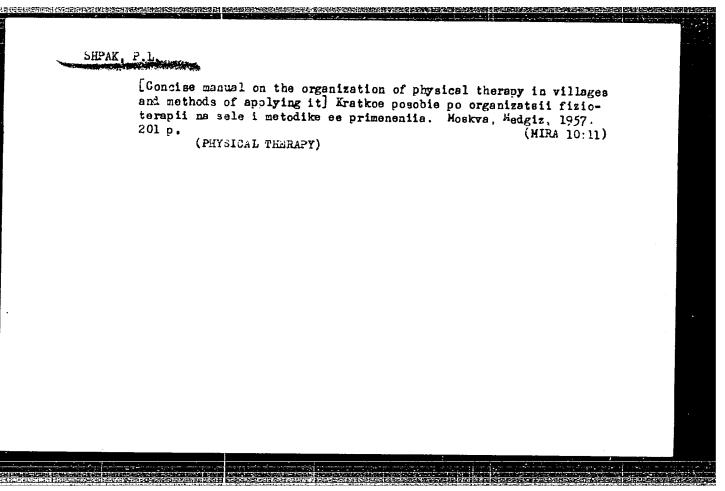
l. Aldanskaya ekspeditsiya Vsesoyuznogo aerogeologicheskogo tresta. Predstavleno akademikom N.M.Strakhovym. (Aldan Plateau—Rare earth metals) (Geological time)

20034-65 EWT(d) IJP(c) CESSION NR: AP!003637	s/0140/64/000/005/0127/0136	
Figure Shpak, G. S. (Khar'kov)	dyon values at	
ITLE: Method of obtaining emisates fiver, points OURCE: IVUZ. Matematika, no. 5, 1964;	or functions which assume given values at ;	
CPIC TAGS: function theory Astract: N.A. Libedev determined consists one to find functions that are rest points z_k , $ z_k < 1$, $k = 1, 2, \ldots$, $ z_k < 1$,	ditions which, when satisfied, engular in the circle $ z \le 1$; which n, assume given values $f(\mathcal{V})(z_k)$, $ z \le 1$ are contained in the exist functions which are subsuthor determines another property $f(z)$, $f(z)$, $f(z)$, varies contined in $f(z)$, $f(z)$	
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ACCESSION NR: AP5003637			
for which G(f) is the largest root of the struct this equation, which, estimates for various subclacarried out here, except in from that of Lebedev. Orig.	in turn, enables one to cosses of regular functions	shove how to con- \ Obtain some accurate \	
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