

SOV/51-7-3-12/21

Plastic Scintillators with Additions of Aryl Derivatives of 1,3,4-Oxadiazole

prepared by cyclization of the corresponding dihydrazides by heating with phosphorus oxychloride (Ref 3). All compounds were purified by re-crystallization and chromatographic treatment. The scintillators were in the form of polystyrene discs (with the appropriate 1,3,4-oxadiazole derivative added to them) of 20 mm diameter and 12 mm height; they were prepared by high temperature polymerization in an atmosphere of nitrogen. The absorption spectra were recorded by means of a spectrophotometer SF-4. The luminescence spectra were obtained by means of the same instrument used as a monochromator; they were recorded photoelectrically. The scintillation efficiency was deduced from the current of a FEU-19 photomultiplier. A sample of Ag^{110} of 0.1 μ curie intensity was used as the source of excitation. The absorption spectra of the eight oxadiazoles are shown in Figs 1 (curves 1-4) and 2 (curves 1-4). The luminescence spectrum of polystyrene is shown as curve 5 in both figures. The greatest amount of overlapping of the absorption spectrum with the luminescence spectrum of polystyrene was exhibited by the compounds with 1-naphthyl radical, that is the compounds ~~ONPD~~, ~~ONPD~~ and ~~ONPD~~. Figs 3 and 4 show the photoluminescence spectra (excited with 253 and 313 m μ mercury lines). Here again the oxadiazoles with 1-naphthyl radical show the greatest amount of overlap with the maximum of the FEU-19 sensitivity. The dependence of the scintillation

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efficiency on the concentration of the oxadiazoles (Fig 7) shows that the compounds ~~ANPD~~, ~~ANPD~~, ~~ANPD~~ and ~~ANPD~~ and ~~ANPD~~ are the most efficient. In a table on p 369 the authors list the absorption and luminescence maxima (cols 3 and 4), the concentration oxadiazole in polystyrene (col 5) and the scintillation efficiency (col 6) of the eight oxadiazole derivatives listed above and eight other 1,3,4-oxadiazole derivatives studied earlier. The authors found that the scintillation efficiency of organic compounds in plastics is determined primarily by their absorption and luminescence spectra and their luminescence yield. The scintillation efficiency may be measured in relative units by Swank and Buck's method (Ref 8), allowing for the overlapping of the luminescence spectrum of the base (polystyrene) and the absorption spectrum of the additive (oxadiazole derivative), the photoluminescence yield of the additive and the efficiency of recording of the emission by the additive. The best scintillation property among the diaryl derivatives of oxadiazole were found in the compounds with 1-naphthyl and biphenyl radicals. Among the sixteen compounds listed in the table on p 369 the following were found to be most efficient in

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plastic scintillators:

- 2,5-di-(4-biphenyl)-1,3,4-oxadiazole (BED);
- 2,5-di-(1-naphthyl)-1,3,4-oxadiazole (dNED);
- 2-phenyl-5-(4-methoxyphenyl)-1,3,4-oxadiazole (MTPD);
- 2-(4-biphenyl)-5-(2-naphthyl)-1,3,4-oxadiazole (dNED);
- 2-phenyl-5-(1-naphthyl)-1,3,4-oxadiazole (dNPD).

There are 7 figures, 1 table and 9 references, 3 of which are Soviet, 4 English, 1 German and 1 translation into Russian.

DATE ISSUED: December 26, 1958

Card 1/4

5 3610

5 4500

AUTHORS:

Bezuglyy, V.D., and Shimanskaya, N.P.

TITLE:

Polarographic study of some oxozoles

PERIODICAL:

Zhurnal obshchey khimii, v. 31, no. 10, 1961,
3160-3177

TEXT: The investigations were concerned with phenyl-, diphenyl-, and naphthyl substituted oxozoles, used as luminophores in a scintillator to establish the relation between their polarographic and optical properties. The measurements were conducted with the Geyrovsky-Shikal-polarograph using a saturated, high specific resistance solution of $N(C_2H_5)_4 I$ in 92% methanol. The solutions of oxozoles used in the investigation were prepared using 60:40 methanol:dioxane mixtures. The experiments showed that phenyl-, naphthyl-, and biphenyl- substituted oxozoles undergo reduction at the cathode giving unique polarographic waves for different substituents. 2-Methyl-5-phenyl-oxazole did not reduce under the experimental conditions used, but 2,5-diphenyl oxazole gave two waves on the polarograph, whose half wave potentials corresponded to

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D227/D304

Polarographic study ...

-2.10 and -2.33V, referred to the standard calomel electrode. Substitution of the 2-phenyl radical with 2 α -naphthyl displaces the first half-wave potential to -1.85V; β -naphthyl substitution displaces $E_{1/2}$ of the first wave to -2.02V. $E_{1/2}$ of the second wave for 1-naphthyl derivative is -2.23 and for 2-naphthyl derivative -2.23, i.e. very near the value for the second wave of 2,5-diphenyloxazole. A greater effect may be obtained by introducing into position 2 of the diphenyloxazole radical, when $E_{1/2}$ of the first wave = -1.77 with $E_{1/2}$ for the second wave -2.18V and $E_{1/2}$ for the third wave -2.28V. This compound may be considered composed of two 5-phenyloxazole groups joined by a phenyl radical forming a bridge between the electron interaction of the two groups. If the bridge is provided by -CH=CH-, a group containing π electrons, $E_{1/2}$ is displaced towards less negative values and is equal to -1.36V. Introduction of a -CH₂-CH₂- bridging group causes the loss of the polarographic activity of 1,2 - di X

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Polarographic study ...

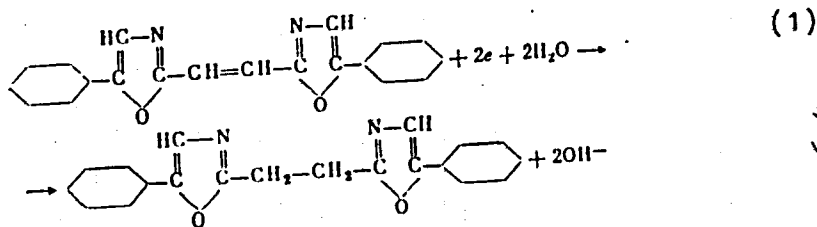
(5 phenyloxazole)-ethone. To investigate the effect of 5-position substituted derivatives, 2,5-di-(4-biphenyl) oxazole and 2-(4-biphenyl)-5-(1-naphthyl)- had $E_{1/2}^1 = -2.00V$, $E_{1/2}^2 = 2,22V$ and $E_{1/2}^1 = -1.97V$, and $E_{1/2}^2 = 2,22V$ respectively. From the experimental results it follows that the 5-membered oxazole ring undergoes reduction, under specific conditions, at the mercury dropping cathode and the ease of reduction depends on the substituent, and its position in the ring. Substitution of phenyl instead of methyl group in 2-position gives rise to unique polarographic waves. Substituents in position-5 have a smaller effect and the introduction of 1-naphthyl in place of biphenyl in the 5-position (substituents in position-2 remaining the same) changes the half-wave potential very little. It follows then that the most readily reducible is the C=N- bond and only after its reduction can the link between position-2 substituents and $>C=C<$ bond (between 4th and 5th C) be broken; the substituent has practically no effect on the $E_{1/2}$ of the second wave. The electro-negative effect of

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Polarographic study ...

aromatic radicals, no reduction occurring with alkyl substituents joined to >C=N- group, is presumably due to the lengthening of the chain of the conjugated double bonds. It must be mentioned that contrary to other investigated oxazole derivatives, compound no. underwent reduction along the -CH=CH- bond and its number of electrons taking part in the reduction of one molecule of the compound was equal to 2. It follows from research that the electrochemical reaction on the mercury dropping cathode for 1,2-di(5-phenyloxazolyl)-ethylene may be represented by first wave Eq.



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which explains the absence of second waves. A table also gives the wave lengths in the region of maximum absorption and it may be seen that the character of λ_{\max} , in general, corresponds to the variation of $E_{1/2}$ values. The fact also confirms the dependence of polarographic results on the character of the substituent. It was also interesting to compare scintillation effectiveness of the compounds with the polarographic results which shows a certain correlation between these properties. It may be concluded that the polarographic method may be used for determining the effectiveness of a given substance as a scintillator, this effectiveness being higher for less negative $E_{1/2}$ values. There are 1 table, 10 figures and 9 references: 5 Soviet-bloc and 4 non-Soviet-bloc. The references to the English-language publications read as follows: F. Hayes, L. King, J. Am. Chem. Soc. 74, 1106 (1952); E. Hartnell, C. Bricker, J. Am. Chem. Soc. 70, 3385 (1948).

Card 5/6

Polarographic study ...

27901
S/079/61/031/010/001/010
D227/D304

ASSOCIATION: Khar'kovskiy filial vsesoyuznogo nauchno-issledovatel'skogo instituta khimicheskikh reaktivov
(Khar'kov branch of All-Union Scientific Research Institute for Chemical Reagents)

SUBMITTED: September 10, 1960

UX

Card 6/6

L 13353-63 EWP(j)/EPF(c)/EWT(m)/BDS AFFTC/ASD Pc-4/Pr-4 RM/WW
ACCESSION NR: AP3002622 8/0079/63/033/006/1726/1732

AUTHOR: Shimanskaya, N. P.; Bezugly*, V. D.

TITLE: Polarographic investigation of oxadiazole derivatives

SOURCE: Zhurnal obshchey khimii, v. 33, no. 6, 1963, 1726-1732

TOPIC TAGS: polarographic investigation, oxadiazole derivative, half-wave potential, polystyrene

ABSTRACT: The polarographic properties of oxadiazole derivatives such as phenyl, biphenyl, naphthyl and others used as luminescent additives in plastics were determined. It was found that substituents in the 2 and 5 positions influenced advantageously the reducibility of the oxadiazole ring. The polarographic data are compared with some optical properties. The half-wave potential values are tabulated. A correlation is found between the half-wave potential (diffusion and catalytic) and the scintillating effectiveness of the substances. On the basis of the polarographic data, the possible scintillating effectiveness of these substances can be determined. The polarographic properties are influenced by the effect of conjugation. Polystyrene was used as an experimental substance. Orig. art. has: 4 figures and 1 table.

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Cord 1/2

L 13353-63

ACCESSION NR: AP3002622

ASSOCIATION: Vsesoyuznyy nauchno-issledovatel'skiy institut monokristallov
i osobo chistykh khimicheskikh veshchestv, Kharkov (All-Union Scientific
Research Institute of Monocrystals and Ultrapure Chemical Substances)

SUBMITTED: 29Jun62

DATE ACQ: 20Jul63

ENCL: 00

SUB CODE: 00

NO REF SOV: 004

OTHER: 009

Card 2/2

REF ID: A25028.18
S 0081/65/000/003/S061/S061

SOURCE: Ref. zh. Khimiya, Abs. 35356

AUTHOR: Shimanskaya, N. P.; Bezuglyy, V. D.

STUDIES OF SOME PROPERTIES OF PLASTIC SCINTILLATORS

Abstract: Studies of some properties of plastic scintillators. Vyp. 3

KEYWORDS: plastic scintillator, electrochemical property, optical property, aging, polystyrene analysis

TRANSLATION: The authors evolved a procedure for the polarographic analysis of plastic scintillators. This enabled them to define the electrochemical properties of scintillators containing furfural-2,3,4 and oxazole-1,3, as well as

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L 43131-64

ACCESSION NO: AR5006436

pounds in polystyrene was carried out polarographically, making possible a study of aging in plastic scintillators. Studied was the stability of luminescent compounds under the influence of temperature and exposure to radiation or violet illumination. Experimental results are cited and evaluated. 6

EEZUGLYY, V.D.; SHIMANSKAYA, N.P.; PERESLENI, Ye.M.

Mechanism of reduction of 1,3-oxazole and 1,3,4-oxadiazole
derivatives. Zhur. ob. khim. 34 no.11:3540-3545 N '64
(MIRA 18:1)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut monokristallov,
Khar'kov.

BEYDOLYI, V.D.;

Polarographic behavior of
17-22 (M.A. 18:2)

L 02421-67 EWT(m)/EWP(j)/T WW/RM/JW

ACC NR: AP6031380

SOURCE CODE: UR/0079/66/036/009/1601/1603

AUTHOR: Shimanskaya, N. P., Malkes, L. Ya.; Bezuglyy, V. D.

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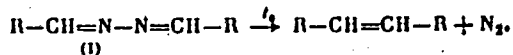
ORG: All Union Scientific Research Institute of Monocrystals, Scintillating Materials and High Purity Chemicals (Vsesoyuznyy naudno-issledovatel'skiy institut monokristallov, stsintillyatsionnykh materialov i osobo chistykh khimicheskikh veshchestv)

TITLE: Reaction rate of thermal decomposition of some azines

SOURCE: Zhurnal obshchey khimii, v. 36, no. 9, 1966, 1601-1603

TOPIC TAGS: ~~thermal decomposition~~, thermal decomposition, reaction rate ~~constant~~, ORGANIC AZINE COMPOUND

ABSTRACT: Polarographic studies were made of the effect of the structure of azines (I) on the reaction rate constants of their thermal decomposition at 300C.



The results showed that the decomposition of azines is a first-order reaction. Reaction. Reaction rate constants calculated for some azines from the experimental data are given in the table:

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UDC: 57.8524

~~SHIMANSKAYA, N.S.~~ SHIMANSKAYA, N.S.

CARD 1 / 2

PA - 1803

SUBJECT USSR / PHYSICS ✓
 AUTHOR GORŠKOV, G.V., SIMANSKAJA, N.S.
 TITLE On Calorimetric Measurements of Preparations of Naturally Radioactive Families.
 PERIODICAL Atomnaja Energija, 1, fasc.5, 86-93 (1956)
 Issued: 1 / 1957

The preparations themselves can be liquid or mixed with other non-active substances which, in some cases, are highly absorbent. The effect of absorption and of self-absorption can mostly not be estimated, for which reason the accuracy of measurements mostly does not go beyond from 2 to 3%. Even greater errors are committed on the occasion of the determination of the radioactivity of preparations by comparison with a gauging preparation of different origin.

On the occasion of calorimetric measurements of naturally radioactive preparations the authors were faced with the lack of complete data in modern tables of radioactive constants (e.g. J.HOLLANDER, J.PERLMAN, G.SEABORG, Rev.Mod.Phys. 25, 429, (1953)). The tables above all contain no detailed information concerning the energies of gamma rays and their relative and absolute intensities. Furthermore, there are no data concerning the number of conversion electrons and the average energies of β -spectra. It is for this reason that the authors carefully analyzed all existing data on energies and on the radiation yields of the elements of the three naturally radioactive families. By the critical investigation of a great number of experimental works it was possible to determine the energy E_1 for each

Atomnaja Energijs, 1, fasc. 5, 86-93 (1956) CARD 2 / 2 PA - 1803
element as well as to determine the total "thermal" energies^E corresponding to one act of decay of the preparations of Ra, MsTh, RaTh, and Ac which are in equilibrium. For the radium preparations also the corrections for the increase of the heat effect caused by the accumulation of RaE and Po²¹⁰ was computed.

Knowledge of all these quantities made calorimetric measuring of the absolute activity of many different preparations of naturally radioactive families possible. For this purpose double calorimeters of the static type were used. A table shows the measuring results of some Ra-, RaTh- and Ac-preparations. The following measurements were furthermore carried out by the calorimetric method: The relations between the milligram equivalent and the data in millicurie for the RaTh- and Ac-isotopes which are in equilibrium; these relations are of importance in the practice of ionization measuring. With the conditions usual in the USSR for ionization measuring (lead filter of 5 mm thickness and standard chamber SGM-1) the following results are obtained: 1 mg-equ RaTh = $1,29 \pm 0,02$ millicurie RaTh, 1 mg-equ. Ac = $10,0 \pm 0,5$ millicurie Ac; the content of radioactive substances on neutron sources of the type ($\alpha + \text{Be}$), the content of Ra and MsTh in radium-mesothorium preparations. The calorimetric method is not suited for radium-mesothorium preparations with unknown time of production ("age"), but it is well suited for the determination of the composition of "young" preparations.

INSTITUTION:

SIMIANSKAYA, N.S.

CARD 1 / 2

PA - 1759

SUBJECT USSR / PHYSICS
 AUTHOR SIMIANSKAJA, N.S., JASUGINA, E.A.
 TITLE Determination of the Half-Life of Ac^{227} by the calorimetric
 Method.
 PERIODICAL Atomnaja Energija, 1, fasc.5, 133-133 (1956)
 Issued: 1 / 1957

As the authors had a weighable quantity of pure Ac^{227} available, they attempted to measure its half-life calorimetrically. The calorimetric measuring of half-life of long-lived isotopes is known to be reduced to the determination of the heat generated in the calorimeter by a known quantity of the radioactive isotope. The formula for the computation of the half-life is written down. The preparation used here for investigations was first chemically purified. The preparation ($Ac_2^{227}O_3$) had a weight of $2,01 \pm 0,02$ milligrams. The impurities in the preparation are quantitatively mentioned. The preparation contained no radioactive impurities. On the occasion of the purification of the preparation also its isotope, $RdAc$, a daughter-product of Ac^{227} , was deposited together with Th. Therefore all basic calorimetric measurements were carried out after 6 months, i.e. after establishment of radioactive equilibrium in the preparation. Measurements were carried out in a double static calorimeter, which is being used in the Radium Institute of the Academy of Science of the USSR for the calorimetric measuring of radioactive substances. The thermal efficiency of the preparation was 23,7 milliwatts ($\pm 0,5\%$). When determining its activity

SIMANSKAYA, N.S.

CARD 1 / 2

PA - 1532

SUBJECT USSR / PHYSICS
 AUTHOR SIMANSKAJA, N.S.
 TITLE The Determination of the Bifurcation Ratio in the Decay Scheme of Po²¹⁰.
 PERIODICAL Zurn.eksp.i teor.fis, 31, fasc.2, 174-177 (1956)
 Issued: 5.10.1956

At first several previous works dealing with the same subject are discussed. When this bifurcation ratio is determined the number of γ -quanta emitted by the Po²¹⁰ source per time unit must be estimated, and absolute activity must be measured with sufficient accuracy. Attention is drawn to the frequently occurring grave errors committed when measuring α -intensity with the help of ionization chambers, impulse chambers and luminescence counters.

The authors used a pure Po²¹⁰ preparation. The number of γ -quanta emitted per second was determined by comparison with a Co⁶⁰ source of known intensity ($N_0 = 19.3 \pm 0.8$). 10^3 acts of decay per second. These measurements were carried out by means of a standard- β -counter with an additional 2 mm aluminium filter, and the results of two measuring series are shown in a diagram. If the ratio k of the intensity of the γ -radiation of the gauged Co⁶⁰ source and of the Po²¹⁰ preparation to be measured as well as their absolute activities N_0 and N_a are known, the number n_γ of the 800 keV- γ -quanta emitted by the Po²¹⁰ per act of decay can be determined. The correction coefficients yet to be intro-

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SHIMANSKAYA, N.S.

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1. 10/10/54 N.S.
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Shimanskaya, N.S.
SHIMANSKAYA, N.S.

In memory of Nikolai Ivanovich Dobronravov. Trudy Radiev.inst.
AN SSSR 6:20-23 '57. (MIRA 11:2)
(Dobronravov, Nikolai Ivanovich, 1891-1949)

SOV/120-53-2-23/37

AUTHOR: Shimanskaya, N. S.

TITLE: The Determination of Half-Lives of Radioactive Isotopes Using a Double Static Calorimeter (Opredeleniye periodov poluraspada radioaktivnykh izotopov s pomoshch'yu dvuyhogo staticheskogo kalorimetra)

PERIODICAL: Priroda i Tekhnika Eksperimenta, 1958, Nr 2, pp 95-97 (USSR)

ABSTRACT: The calorimeter is shown in Fig.1 and consists of two identical internal calorimetric cylinders A and B placed inside the screen C whose temperature T_0 is kept constant. If the radioactive sample is placed, for example, in A then after a certain interval of time a temperature difference is established between A and B. This temperature difference is a function of the heating power of the radioactive sample only, for a given calorimeter system. The calorimeter is calibrated in a preliminary experiment by a special heater placed in the same cylinder A. A particularly useful variant of this apparatus is the case when the radioactive sample is placed in the cylinder A and the heater in the other

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The Determination of Half-Lives of Radioactive Isotopes Using a Double Static Calorimeter.

cylinder B . The heater is adjusted so that the temperature difference is reduced to zero. The device was used to measure the half-life of Rn^{222} . The half-life was found to be $T = 3.83 \pm 0.03$ days, which is in good agreement with the accepted value which is $T = 3.825 \pm 0.005$ days. There are 2 figures, no tables and 8 references, 4 of which are English, 1 French and 3 Soviet.

ASSOCIATION: Radiyevyy institut AN SSSR (Radium Institute of the Academy of Sciences of the USSR)

SUBMITTED: July 2, 1956.

Card 2/2

1. Radioisotopes--Half life
2. Half life--Measurement
3. Calorimeters--Applications

SOV/48-22-7-19/26

Trigor'ev, G. I., Kuznetsov, B. S., Shiganskaya, N. I.,
Molodtsov, I. A.

Determination of the Ratio L/K in ^{159}Gd and ^{165}Ho and the
estimation of the Transmutation Energies of $^{159}\text{Gd} \rightarrow \text{Tb}$
and $^{165}\text{Ho} \rightarrow \text{Ho}^{165}$ (Opredeleniye otnosheniya L/K dlya
 ^{159}Gd i ^{165}Ho i otsenka energii perekhodov $^{159}\text{Gd} \rightarrow \text{Tb}$
i $^{165}\text{Ho} \rightarrow \text{Ho}^{165}$)

IZVESTIYA AKADEMII NAUK SSSR, Seriya fizicheskaya, 1959,
Vol. 22, Nr 7, pp. 850-860 (USSR)

ABSTRACT: The decay energy ϵ_0 of radioactive isotopes, which are sub-
jected to an electron capture can be determined by 5 different
methods. They are described. From the evidence given it is
concluded, that the 5th method, that utilizing the ratio L/K
is very convenient in the determination of small transmutation
energies (< 200 keV) in isotopes with a relatively simple
decay scheme, which do not exhibit a considerable converting
cascade γ -radiation. The application of this method is limit-
ed by the imperfections still inherent in the modern theory

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SOV/43-22-7-1)/26

Determination of the Ratio L/K in Dy^{159} and Er^{165} and an Estimation of the Transmutation Energies of $Dy^{159} \rightarrow Tb^{159}$ and $Er^{165} \rightarrow Ho^{165}$

of K-capture and by the incomplete knowledge of the qualitative and quantitative rules governing the processes of the re-arrangement of the electron shell of the atom. L/K was determined for two isotopes of rare earths, Dy^{159} and Er^{165} , both having a neutron deficit. Proceeding from the results the transmutation energies of the processes $Dy^{159} \rightarrow Tb^{159}$ and $Er^{165} \rightarrow Ho^{165}$ were estimated. A γ -spectrometer combined with a proportional counter was used for measuring the energies and the intensities of an X-ray K- and L-radiation. The proportional counter (Ref 20) permitted to measure the γ - and X-ray radiation of small energies, which is quite impossible with other methods. The proportional counter with a cylindrical aluminum cathode and its circuit diagram is described. The recording power of the counter for γ - and X-ray-radiation of varying energy is computed according to the known absorption coefficients for this radiation in argon and beryllium (Ref 22), taking into account the geometry of the experimental arrangement. The electronic circuit diagram

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SOV/48-22-7-1)/26

Determination of the Ratio L/K in Dy¹⁵⁹ and Er¹⁶⁵ and an Estimation of the Transmutation Energies of Dy¹⁵⁹ → Tb¹⁵⁹ and Er¹⁶⁵ → Ho¹⁶⁵

and the calibration of the device is described. The Dy¹⁵⁹ source was obtained from a tantalum target, which was irradiated in the synchrocyclotron of the "United Institute of Nuclear Research" with 660 MeV protons. The ratio L/K was computed according to formula (3). It is shown that the transition Dy¹⁵⁹ → Tb¹⁵⁹ must be classified as being superforbidden. Marshak's formula was used, giving an energy value of 79^{+10}_{-5} keV for this transition. The lowest level of Tb¹⁵⁹ at 57 keV is apparently not excited in the decay of Dy¹⁵⁹. An estimation of the quantity ft on the basis of the decay energy of 79 keV and a half-life of 156 days furnishes a value for lg ft of about 6,2. According to the classification of King (Ref 32) this value agrees with the assumption, that this transmutation is a superforbidden one. The Er¹⁶⁵-sources were also obtained from tantalum irradiated with fast protons ($\epsilon_0 = 660$ MeV). The X-ray radiation

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Determination of the Ratio L/K in Dy¹⁵⁹ and Er¹⁶⁵ and an Estimation of the Transmutation Energies of Dy¹⁵⁹ → Nb¹⁵⁹ and Er¹⁶⁵ → Ho¹⁶⁵

of a series of tantalum targets irradiated for different periods was measured. The ratio I_L/I_K (for the intensities of these radiations) was equal to 0,40. From this value for L/K a result of $1,2 \pm 0,4$ was obtained. Using Marghak's formula and the experimentally found value of L/K (Er¹⁶⁵) 82_{-5}^{+10} keV were found for the transmutation energy of the process Er¹⁶⁵ → Ho¹⁶⁵. The value of lg ft was 3,1 with a half-life of 10,5 hours, which is in agreement with the permitted character of the transmutation. There are 9 figures, 1 table, and 35 references, 3 of which are Soviet.

ASSOCIATION: Radiyevyy institut im. V. G. Khlopina Akademii nauk SSSR
(Radium Institute imeni V. G. Khlopin, AS USSR)

Card 4/4

AUTHORS: Gorshkov, G. V. , Gritchenko, Z. G. , SOV/56-34.3-39/55
Shimanskaya, N. S.
TITLE: The Calorimetric Determination of the Half-Life of Ra²²⁶
(Kalorimetricheskoye opredeleniye perioda poluraspada Ra²²⁶)
PERIODICAL: Zhurnal Eksperimental'noy i Teoreticheskoy Fiziki, 1958,
Vol. 34, Nr 3, pp. 756 - 757 (USSR)

ABSTRACT: First, brief reference is made to some previous works dealing with the same subject. The authors of the present report carried out careful calorimetric measurements on 3 equilibrated radium preparations which were liberated from possible contaminations by means of additional crystallization. The purity of these preparations was controlled by means of the spectroscopic method. The results of the immediate weighing of the radium preparations prior to their sealing, their radium-content and the results of the calorimetric measurements carried out by means of a double static calorimeter, are contained in a table. The last column of the table contains the values found here for $Q_{\alpha+\beta}/p$ - the

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The Calorimetric Determination of the Half-Life of Ra²²⁶

SOV/56-34 -3-39/55

thermal effect of the α - and β -radiation of 1 g radium. Calculating these values, the absorption of the γ -rays in the preparation itself (self-absorption), in the protective container, in the glass of the ampules and within the walls of the calorimetric cylinder, were taken into consideration. Also the increase of the thermal effect due to the accumulation of Po²¹⁰ and RaE in the preparations was taken into account. ϵ (the energy liberated in the calorimeter in a process of decay) was calculated on the basis of the last experimental data on the α - and β -spectra of the elements of the radium-series for an equilibrated preparation of Ra²²⁶. This energy amounted to 25.335 MeV ($\pm 0.3\%$). Utilizing this value, the authors found the value $T = 1577 \pm 9$ years for the half life of Ra²²⁶. Hence results the value $z = 3.71 \pm 0.02 \cdot 10^{10}$ decay-processes/sec.g. for the specific activity. Further measurements of these important values z and T for Ra²²⁶ with the methods discussed here and also by other methods, would be desirable. There are 1 table and 10 references, 4 of which are Soviet.

Card 2/3

The Calorimetric Determination of the Half-Life of Ra²²⁶

SOV/56-34-3-39/55

ASSOCIATION: Radiyevyy institut Akademii nauk SSSR
(Radium Institute AS USSR)

SUBMITTED: December 6, 1957

Card 3/3

ШИМАНСКАЯ, В.И.

1950-1951 (Proceedings of the Institute of Atomic Energy, Academy of Sciences USSR, No. 1, 9, Moscow, 1950, 1951, 187 p. Extracts being inserted, 1,750 copies printed.)
Ed. I. K.M. Perfilov, Doctor of Physical and Mathematical Sciences, Ed. of Publishing House G.V. Arnold Tech. Ed. I. A.V. Shirova.

FOREWORD: The volume is intended for physicists.
COMMENT: The book represents volume 9 of the Proceedings of the Institute of Atomic Energy and contains the results of studies conducted at the Institute during the period 1945 to 1950. There is a number of articles dealing with the study of nuclear reactions occurring with particles of different energies ranging from a few MeV up to hundreds of MeV. Others treat different problems in the energy district between 100 and 1000 MeV. Some articles deal with the theory of neutron interaction with matter, and other problems connected with the theory of neutron interaction with matter are presented. The majority of the articles are concerned with problems of method. The authors provide complete descriptions of the construction of equipment and of the details of tests performed under laboratory conditions. In particular, the personalities are mentioned. References accompany individual articles.

Shimansky, V.I. Strahlung fission due to high excitation energy	42
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21(8)
AUTHORS: Gorshkov, G.V., Karavayev, F.M., and Shimanskaya, N.S. SOV/115-59-3-24/29

TITLE: The Determination of the Radium Content in Radium Compounds (Ob opredelenii soderzhaniya radiya v radiyevykh preparatakh)

PERIODICAL: Izmeritel'naya tekhnika, 1959, Nr 3, pp 52-53 (USSR)

ABSTRACT: The radium content of radium compounds is mainly determined by the ionization method, or more exactly, its gamma equivalent is determined. The ionization effect of the radiation of the compound under investigation is compared to that of a standard with a known radium content. At VNIIM, two state standards, X and XI, are used, whose radium content was set equal (for 1957) to 29.37 and 14.27 mg radium elements. The self-absorption of the gamma radiation within the radiation source itself is not considered sufficiently. Although lead filters are used, which are 2 cm thick at VNIIM, whereby the soft gamma radiation is eliminated, the error can attain a consider-

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SOV/115-59-3-24/29

The Determination of the Radium Content in Radium Compounds

able magnitude, if the differences of self-absorption are not taken into consideration. The authors determined the accuracy of contemporary ionization methods used for determining the radium content. For this purpose, three pure radium compounds were available which were to be used for the calorimetric determination of the radium half decay period (Ra^{226}). The results of these investigations and measurement results of VNIIM and the Radiyevyy institut AN SSSR -RIAN- (Radium Institute AS USSR) are shown in one table. The calculations performed by the authors show that the difference of the self-absorption of the gamma radiation of radium in 15 mg $RaCl_2$ and 150 mg $RaBr_2$ is of a considerable magnitude. The effective self-absorption in standard XI was found to be 0.9% while it was 1.7% in 150 mg $RaBr_2$, whereby the difference was 0.8%. The authors recommend to establish new standards in the USSR with a radium content of 1, 5, 10, 25, 100, 200, 500 mg, whereby the error

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SOV/115-59-3-24/29

The Determination of the Radium Content in Radium Compounds

caused by the different self-absorption were reduced to a greater extent. In addition they recommend the application of lead filters with thicknesses of not less than 1-1.5 cm. Until new state standards are created the authors recommend the application of a formula for obtaining an accuracy of 0.3-0.5%

$$p = I (1.006 + 3.6 \cdot 10^{-3} \sqrt[3]{I})$$

where I is the milligram-equivalent of the compound under investigation. A footnote says that the standards X and XI are regarded also as secondary international standards. There are: 1 table and 6 references, 3 of which are Soviet and 3 English.

Card 3/3

BAK, M.A.; GOPSHKOV, G.V.; MATVIYENKO, V.I.; PETRZHAK, K.A.; SHIMANSKAYA, N.S.

Determination of the neutron yields of the sources Ra + Be, Ac +
Be, MsTh + Be, and P + Be. Trudy Radiev.inst.AN SSSR 9:120-125
159. (MIRA 14:6)

(Neutrons)

SHIMANSKAYA, N.S.

Determination of the correction for the thermal inertia of the
calorimeter in calorimetric measurements of radioactive preparations.
Trudy Radiov.inst.AN SSSR 9:126-130 '59. (MIRA 14:6)
(Calorimetry) (Radioactive substances)

SHIMANSKAYA, N.S.

Role of physicochemical processes in calorimetric measurements of
radioactive substances. Trudy Radiov. inst. AN SSSR 9:131-133 '59.
(MIRA 14:6)

(Radioactive substances) (Calorimetry)

SOV/89-6-4-14/27

21(8)

AUTHORS: Gorshkov, G. V., Shimanskaya, N. S.TITLE: Total Energy of the Radioactive Radiation of a Radium Preparation (Ra²²⁶) in Equilibrium (Polnaya energiya radioaktivnogo izlucheniya ravnovesnogo preparata radiya (Ra²²⁶))

PERIODICAL: Atomnaya energiya, 1959, Vol 6, Nr 4, pp 474-475 (USSR)

ABSTRACT: In 1935 I. Zlotovskiy calorimetrically measured the total energy of all radioactive radiations radiated from a radium preparation in equilibrium. This value was now checked with the help of 3 sources the exact radium content (Ref 4) of which was known. This measurement was carried out with the static γ -calorimeter (Ref 5) the tungsten walls of which absorbed ~93% of the γ -radiation of Ra-Ra(B+C). For q (total energy) the value 138.9 ± 0.7 cal/h.lg Ra was measured. It is by 0.7% lower than that obtained by Zlotovskiy. This lower value agrees well with expectations. Individual data, from which q was calculated, are shown by a table. Ye. K. Smirnova produced the radium preparations. Yu. S. Martynov took part in the measurements. There are 1 table and 6 references, 4 of which are Soviet.

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87959

87959
S/115/60/000/012/013/018
B019/B056

Legend to Table 1: 1) Day of measurement. 2) Number of years between sealing and measuring. 3) Composition of the source as calculated on the various days of measurement. 3a) Ra, millicuries. 3b) Milligram-equivalent of radium. 3c) Millicuries. 3d) Ratio between the number of decaying atoms: $T_t = (\lambda N_t)RdTh / (\lambda N_t)MsTh$, where λ is the decay constant and N_t the atom number. 3e) Millicuries. 3f) Milligram-equivalent of radium. 4) Radium- γ -equivalent, milligram-equivalent of radium. 4a) J_{calc} . 4b) J_{exp} . 4c) J/J_o calc. 4d) J/J_o exp. 5) Relative neutron number. 5a) A_n calc. 5b) A_n exp. 5c) A_n/A_{n_o} calc. 5d) A_n/A_{n_o} exp. 6) Q value, cal/h; 6a) Q_{calc} . 6b) Q_{exp} .

Legend to Fig. 3: The curves denoted by 1, 2, 3, 4, and 5 stand for the relative neutron yield and the relative radium- γ -equivalent for five different original compositions of the Ra+MsTh source with 55% Ra+45% MsTh, 60% Ra + 40% MsTh; 65% Ra + 35% MsTh; 70% Ra + 30% MsTh; 75% Ra + 25% MsTh. a) = years.

Card 2/3

33.3 | 44.0

BIRYUKOV, Ye.I.; BRIGOR'YEV, O.I.; KUZNETSOV, B.S.; SHIMANSKAYA, N.S.

Decay of Nd¹⁴⁰ and Pr¹⁴⁰. Izv.AN SSSR.Ser.fiz. 24 no.9:
1135-1144 S '60. (MIRA 13:9)
(Neodymium--Decay) (Praseodymium--Decay)

0727

S/048/61/025/001/019/07
B029/B060

24.6/20

Biryukov, Ye. I., Grigor'yev, O. I., Kuznetsov, B. S.,
Shimanskaya, N. S.

AUTHORS:

Decay of Dy¹⁵⁹

TITLE:

PERIODICAL:

Izvestiya Akademii nauk SSSR. Seriya fizicheskaya, v. 25,
No. 1, 1961, 109-110

TEXT: The authors studied the electromagnetic radiation of Dy¹⁵⁹ (T_{1/2} = 144 days) arising from the irradiation of a tantalum target by Mev protons by means of a spectrometer with proportional counter and a scintillation gamma spectrometer. The enclosed figure shows the spectrum of the electromagnetic radiation of Dy¹⁵⁹ in the range of 15 to 60 kev, taken with a filter of 130 mg cm⁻² Al. The ratio between intensities of 58-kev gamma radiation and the KX radiation of Dy (44.5; 50.4 kev) is I_{KX}/I_{γ58} = 53. The contribution of the nonconverted 58-kev gamma radiation amounts to 6.1%, which is also in agreement with the data

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89253

Decay of Dy¹⁵⁹S/048/61/025/001/019/031
B029/B060

relative to gamma decay of Gd¹⁵⁹. Apart from the γ_{58} line, a weak line with an energy of 350 keV was also observed (Ref. 2). The intensity of this line amounts to $2 \cdot 10^{-5}$ quanta per decay event. Shorter wave lines in the energy range up to 2 MeV were no more observed, or at least not any such with an intensity exceeding 10^{-4} to 10^{-5} quanta per decay event. Simultaneous measurements of the two Dy¹⁵⁹ sources in the 4π scintillation counter and in the 4π gas counter gave the following ratios between the intensities of the LX and KX radiation and the intensities of the corresponding LX - LX and KX - KX coincidences:

$$\frac{I_{KX}}{I_{KX-KX}} = 6.56 \pm 0.18, \quad \frac{I_{LX}}{I_{LX-LX}} = 48.1 \pm 4.1, \quad \frac{I_{KX-KX}}{I_{LX-LX}} = 37.1 \pm 5.8, \quad \frac{I_{LX}}{I_{KX}} = 0.21 \pm 0.01.$$

One may calculate therefrom the ratio L_1/K_1 for the transition to the first excited 58-keV level of Tb¹⁵⁹ and the amount \mathcal{X} of the bifurcation. If the value $\bar{\omega} = 0.18 \pm 0.02$ is assumed for the L fluorescence yield of Tb, one obtains $L_1/K_1 = 0.58$ and $\mathcal{X} = 0.32 \pm 0.08$. The article under consideration is the reproduction of a lecture delivered at the 10th All-Union Conference on Nuclear Spectroscopy, which took place in Moscow

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89253
S/048/61/025/001/019/031
B029/B060

Decay of Dy¹⁵⁹

from January 19 to 27, 1960. There are 1 figure and 3 non-Soviet-bloc references.

ASSOCIATION: Radiyevyy institut im. V. G. Khlopina Akademii nauk SSSR
(Radium Institute imeni V. G. Khlopin, Academy of Sciences
USSR)

Card 3/4

BIRYUKOV, Ye.I.; KUZNETSOV, B.S.; SHIMANSKAYA, N.S.

Mean energy of the β -spectrum of Y^{90} . Zhur.eksp.i teor.fiz. 41
no.1:22-23 JI '61. (MIRA 14:7)

1. Radiyevyy institut AN SSSR.
(Beta rays—Spectra) (Yttrium—Isotopes)

S/048/62/026/002/008/032
B101/B102

AUTHORS: Biryukov, Ye. I., and Shimanskaya, N. S.

TITLE: K/β^+ ratio for Pr^{140} PERIODICAL: Akademiya nauk SSSR. Izvestiya. Seriya fizicheskaya,
v. 26, no. 2, 1962, 215 - 216

TEXT: A direct measurement was made of K/β^+ of a Pr^{140} preparation obtained from the reaction $\text{Pr}^{141}(n,2n)\text{Pr}^{140}$. Pr_2O_3 was applied in a thin layer (1.3 mg/cm^2) to a polyethylene film, and bombarded with 14-Mev neutrons (intensity $10^{10} \text{ neutrons}\cdot\text{cm}^{-2}\cdot\text{sec}^{-1}$). Check tests showed that the emission from polyethylene and oxygen was negligibly small. K/β^+ was determined with a 4π scintillation gamma spectrometer with $40\cdot40 \text{ cm}$ CsI(Tl) . For measuring the annihilation radiation spectrum, the source was shielded with a lead filter ($1000 \text{ mg}\cdot\text{cm}^{-2}$) and placed into a channel (6 mm in diameter) bored through the crystal axis. The K-radiation

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K/β^+ ratio for Pr^{140}

S/048/62/026/002/008/032
B101/B102

spectrum was then measured without a filter. Two AM-100 (AI-100) multi-channel analyzers were used for the measurement. From the two spectra, the mean value of K/β^+ was found to be 0.75 ± 0.03 , which is in good agreement with the Pr^{140} decay scheme published earlier (Izv. AN SSSR, Ser. fiz., 24, 1135 (1960)). By extrapolating K/β^+ for allowed transitions one obtains 2480 keV as the limit of the Pr^{140} positron spectrum. L. Zyryanova and K. Gromov are mentioned. There are 2 figures and 5 references: 3 Soviet and 2 non-Soviet. The two references to English-language publications read as follows: Browne, C., Rasmussen, J., Surls, J., Martin, D., Phys. Rev., 85, 146 (1952); Handley, T., Olson, E., Phys. Rev., 96, 1003 (1954).

ASSOCIATION: Radiyevyy institut im. V. G. Khlopina Akademii nauk SSSR
(Radium Institute imeni V. G. Khlopin of the Academy of
Sciences USSR)

Card 2/2

BIRYUKOV, Ye.I.; SHIMANSKAYA, N.S.

Decay of ^{141}La . Izv. AN SSSR. Ser. fiz. 27 no.11:1402-1407
N '63. (MIRA 16:11)

BIRYUKOV, Ye.J.; NOVIKOV, V.T.; SHIMANSKAYA, N.S.

Decay of Pr¹³⁹. Izv. AN SSSR. Ser. fiz. 27 no.11:1408-1411
N '63. (MIRA 16:11)

BIRYUKOV, Ye. I.; NOVIKOV, V. T.; SHIMANSKAYA, N. S.

"Concerning the Decay Chain $^{134}_{58}\text{Ce} \rightarrow ^{134}_{57}\text{La} \rightarrow ^{134}_{56}\text{Ba}$."

report submitted for All-Union Conf on Nuclear Spectroscopy, Tbilisi, 14-22
Feb 64.

Radiyevyy Inst (Radium Inst)

SHILANSKAYA, N. S.; ZHIBETSKIY, E. G.

Mean energies of electron and positron beta spectra. Atoms. Energy.
17 no.1:9-17 J1 '64. (MIRA 17:7)

BIRYUKOV, Ye.I.; MARTYNOV, Yu.S.; NOV'KOV, V.T.; SHIMANSKAYA, N.S.

Mean energy of the Pr^{142} β -spectrum. Zhur.eksp.i teor.fiz 46
no.6:2242-2243 Je '64. (MIRA 17:10)

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SECRET

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... prepared by ...

"APPROVED FOR RELEASE: 08/23/2000

CIA-RDP86-00513R001549510009-7

APPROVED FOR RELEASE: 08/23/2000

CIA-RDP86-00513R001549510009-7"

SHIMANSKAYA, O.A.

Experiment demonstrating the significance of vitamins for pigeons.
Biol. v shkole no.2:66-67 Mr-Apr '59. (MIRA 12:4)

1. Kremenetskiy pedagogicheskiy institut.
(Beri-beri) (Pigeons as laboratory animals)

SHIMANSKAYA, R.I.; FLYUSHIN, V.G.; VAYSBERG, N.S.

Use of pyrolysis tar from wastes of the synthetic alcohol
manufacture. Khim.i tekhn. topl.i masel 7 no.9:34-37 S '62.
(MIRA 15:8)

1. Ural'skiy filial AN SSSR.
(Petroleum products)

SHIMANSKAYA, S. A.

Cand. Med. Sci.

Dissertation: "Vaccinotherapy of Dysentery with Heated Flexner's
Monovaccine."

2/3/50

Acad. Med. Sci. USSR

SO Vecheryaya Moskva
Sum 71

SHIMANSKAYA, S.A. (Kaunas)

Teaching epidemiology in stomatological institutes. Zhur.mikrobiol.
epid. i immun. 27 no.5:95-96 My '56. (MIRA 9:8)
(EPIDEMIOLOGY--STUDY AND TEACHING)

SHIMANSKAYA, S.A.

Clinical aspects and laboratory diagnosis of typhus. Zhur.mikrobiol.
epid. i immun. 30 no.2:108-111 F '59. (MIRA 12:3)

1. Iz Nauchno-issledovatel'skogo instituta sanitarii i gigiyeny
Vil'nyusa.

(TYPHUS,
clin. aspects & diag. (Rus))

ZAVERUKHA, B.V.; SHIMANSKAYA, V.O.

Rare medical plant. Priroda 49 no.11:109-110 N '60.(MIRA 13:11)

1. Kremenetskiy pedagogicheskiy institut.
(Euphorbia)

SIROTA, H.H.; DANIL'KEVICH, M.I.; SIROTA, A.G.; SHIMANSKAYA, V.P.

Electrets made from high polymers. Dokl. AN BSSR 2 no.10:413-
415 N '58. (MIRA 12:8)

(Electrets)

ACCESSION NR: APL011696

S/0250/61/008/001/0024/0025

AUTHORS: Sirota, N. N.; Shimanskaya, V. P.

TITLE: Lattice constant of zinc sulfide-cadmium sulfide solid solution films

SOURCE: AN BSSR. Doklady*, v. 8, no. 1, 24-25

TOPIC TAGS: zinc sulfide, cadmium sulfide, ZnS CdS solid solution, lattice constant, ZnS CdS phase composition, ZnS CdS structure, vaporized coating, x ray apparatus URS 501

ABSTRACT: Films of the binary system ZnS-CdS have been subjected to x-ray analysis in order to determine its phase composition, its structure, and its crystal lattice constant. Experiments were conducted directly after film deposition and also after a heat treatment of films which were produced by sublimating a pressed mixture of ZnS and CdS of a definite composition onto glass and quartz plates. The process was carried out in a vacuum of no less than 10^{-4} mm Hg. For CdS the temperature of the plates was held at 90C and for ZnS at 200C. After the deposition the samples were held in vacuum at 300C. Microscope inspection and x-ray analysis proved that the films were either polycrystalline or monocrystalline, the latter ranging from 0.5 to 1.6 μ in thickness. They were monophasic in type and

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ACCESSION NR: AP4014696

sphaleritic in structure. Lattice constant was calculated from x-ray photographs taken with apparatus URS-501. Figure 1 of the Enclosure shows the relation of this constant to the composition of the solid solution. Lower rates of coating in vacuum and in hydrogen sulfide tended to produce monocrystalline film, while faster rates favored the polycrystalline ones. Orig. art. has: 1 graph and 2 micro-photographs.

ASSOCIATION: Institut fiziki tverdogo tela i poluprovodnikov AN BSSR (Institute of Solid State Physics and Semiconductors AN BSSR)

SUBMITTED: 01Aug63

DATE ACQ: 26Feb64

ENCL: 01

SUB CODE: FH

NO REF SOV: 000

OTHER: 002

Card 2/32

S 0257 64 078/001.0436/0437

AUTHOR: Sirota, N. N.; Shimanskaya, V. P.

The energy gap of the energy gap according to the absorption spectra

1974, 436-437

zinc sulfide base compound, cadmium sulfide containing
zinc sulfide, cadmium sulfide, sulfide, zinc blende,
energy gap, absorption coefficient, film, spectroscopy,
resistor

In ZnS-CdS systems with zinc
has determined measurements of the absorption on the
side of the side of the... The values averaged from
changes linearly, de-
investigations
glass of quartz
of the absorption spectra

REF ID: A64044253

obtained figures close to those obtained by other researchers (Gross, G. I., *Usp. Fiz. Nauk*, 1955, 11; Khansevarov, N. Yu., *Izv. Akad. Nauk SSSR Ser. Fiz. Nauk*, 1955, 48). The study has been published in *Usp. Fiz. Nauk*, 1955, 11, and some other journals.

Institute of Solid State Physics and Semiconductors, AN BSSR

SUBMITTED: 12Feb64

ENCL: 00

REF SOV: 004

OTHER: 001

SIROTA, N.M.; SHIMENSKAYA, V.P.

Width of the forbidden zone according to absorption spectra of thin films in the system ZnS - CdS. Dokl. Akd. Nauk SSSR 3 no.7:436-437 '64.

(MIRA 17:10)

1. Institut fiziki tverdogo tela i poluprovodnikov Akd. Nauk SSSR.

SHIMANSKAYA, Ye.T. [Shymans'ka, O.T.]

Tepler method for investigating critical states of substances, Part 1:
Experimental procedure [with summary in English]. Ukr.fiz.zhur. 3 no.4:
542-551 J1-Ag '58. (MIRA 11:12)
(Optical measurements) (Specific gravity)

GOLIK, A.Z. [Holyk, O.Z.]; SHIMANSKAYA, Ye.T. [Shymans'ka, O.T.]

Investigation of the critical state of substances by Toepler's method. Part 2. Temperature dependence of the density of hexane near the critical point. Ukr.fiz.zhur. 4 no.6:769-788 II-D '59.
(MIRA 14:10)

1. Kiyevskiy gosudarstvennyy universitet im. T.G.Shevchenko.
(Hexane--Thermal properties)

SHIMANSKAYA, Ye.T. [Shymans'ka, O.T.]

Tepler method for investigating the critical state of a substance. Part 3: Dependence of the density of an octane solution in hexane on the temperature near the liquid-vapor critical point. Ukr. fiz. zhur. 5 no.4:549-559 J1-Ag '60. (MIRA 13:11)

1. Kiyevskiy gosudarstvennyy universitet.
(Critical point) (Hexane) (Octane)

СОВЕТСКОМУ, ye T.

30

PHASE I BOOK EXPLOITATION SOV/5469

Soveshchaniye po kriticheskim yavleniyam i flyuktuatsiyam v rastvorakh. Moscow, 1960.

Kriticheskiye yavleniya i flyuktuatsii v rastvorakh; trudy soveshchaniya, yanvar' 1960 g. (Critical Phenomena and Fluctuations in Solutions; Transactions of the Conference, January 1960) Moscow, Izd-vo AN SSSR, 1960. 190 p. 2,500 copies printed.

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Responsible Ed.: M. I. Shakhparonov, Doctor of Chemical Sciences, Professor; Ed. of Publishing House: E. S. Draganov; Tech. Ed.: S. G. Tikhomirova.

PURPOSE : This collection of articles is intended for scientific personnel concerned with chemistry, physics, and heat power engineering.

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Critical Phenomena and Fluctuations

SOV/5469

COVERAGE: The book contains 24 of the 26 reports read at the Conference on Critical Phenomena and Fluctuations in Solutions organized by the Chemical Division of Moscow State University, January 26-28, 1960. The reports contain results of investigations carried out in recent years by Soviet physicists, chemists, and heat power engineers. The Organizing Committee of the Conference was composed of Professor Kh. I. Amirzhanov, A. Z. Golik, I. R. Krichevskiy (Chairman), V. K. Semchenko, A. V. Storonkin, I. Z. Fisher, and M. I. Shakhparonov (Deputy Chairman). References accompany individual articles.

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S/081/61/000/021/010/094
B*02/B138

AUTHORS: Shimanskaya, Ye. T., Shimanskiy, Yu. I., Golik, A. Z.

TITLE: Investigation of the critical state of pure substances by Tepler's method

PERIODICAL: Referativnyy zhurnal. Khimiya, no. 21, 1961, 43, abstract 21B347 (Sb. "Kritich. yavleniya i flyuktuatsii v rastvorakh", M., AN SSSR, 1960, 171 - 188)

TEXT: A method has been developed, for the investigation of critical states, by means of which the density ρ of a substance can be measured in any point in a chamber (by the optical Tepler method) with long-time thermostating. The apparatus is described in detail. Heptane and hexane were examined. Density has a non-monotonic gradient with respect to the chamber height Z , and has a maximum at the meniscus. This maximum increases as the temperature approaches the point T_m at which the meniscus vanishes. With a steady temperature change (rate: 2 deg/hr) the $d\rho/dZ$ maximum is present on heating and absent on cooling (i. e. a hysteresis is observed). With irregular changes in temperature and long-time
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B102/B138

Investigation of the critical...

(15 - 20 hr) thermostating, the $d\rho/dZ$ maximum is, however, present on heating as well as cooling; the maxima are then lower than in the case of steady heating. The $\rho(Z)$ curves are found by integrating $d\rho/dZ = f(Z)$. For $T > T_m$ they are S-shaped. In the lower part of the chamber density is higher, and in the upper part lower, than critical. This is in full agreement with classical representations regarding the existence of a critical point, and not a region, when allowing for the effect of gravitational field. The critical state is realized only in a narrow layer at the point where the meniscus vanishes. Above and below this layer the substance is not in a critical state, although its temperature is critical. As the density difference throughout the chamber corresponds at the critical temperature to the equilibrium state, then it must be assumed that displacement sometimes occurs, levelling the density and removing the system from the state of equilibrium. [Abstracter's note: Complete translation.]

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SHIMANSKIY, Yu.I.; SHIMANSKAYA, Ye.T.; Primalni uchastiye: YATSYUTA, N.A.,
student; ZAVALIN, I.V., aspirant

Study of the density of benzene near the critical point. Ukr.
fiz. zhur. 7 no.8:861-868 S '62. (MIRA 16:1)

1. Kiyevskiy universitet.

(Benzene--Density) (Critical point)

ZAVALI, I.V.; SHLENKAYA, Ye.T.; SHIMENSKIY, Yu.I.; Prinsipali uchastiye:
MITYUKOVSKAYA L.M., student; KOVALENKO G.F., student; KHOMUTOVA, Z.L.
student

Behavior of the density of the solution benzene-propol alcohol near
the critical point at the liquid - vapor boundary. Ukr. Fiz. zhur.
9 no.5:491-496 My '64. (ISSA 17:9)

1. Kiyevskiy gosinzhenerstvennyy universitet.

BOGIYAKOV, G.V.; SHIMANSKIY, M.A.; SHIMANSKIY, G.V.

Possibility of using correlation analysis for determining
the content of some components in the iron ores of the
Angara-Ilim region. Izv. vyz. ucheb. zav.; geol. i razv.
6 no.5:97-103 My '65. (MIRA 18:10)

1. Irkutskiy politekhnicheskiy institut i Irkutskiy geologo-
razvedochnyy tekhnikum.

UCHAKIN, Yu.M.; SHIMANSKIY, A.A.; PAULLER, T.I.

Rare alkalies containing feldspars from the pegmatites of the Sayan Mountains, *Geokhimiya* no.8:673-680 '62. (MIRA 15:9)

1. East-Siberian Institute of Geology, Siberian Branch of the Academy of Sciences, U.S.S.R. Polytechnic Institute, Irkutsk.

(Sayan Mountains--Alkalies) (Sayan Mountains--Feldspar)

SHIMANSKIY, A.A.; UCHAKIN, Yu.M.

Distribution of alkalis in microclines from pegmatites
of the Eastern Sayan Mountains. Geokhimiia no.9:833-836
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1. Polytechnical Institute of East-Siberial Institute of
Geology, Siberian Branch of the Academy of Sciences,
U.S.S.R., Irkutsk.
(Sayan Mountains--Alkalies) (Sayan Mountains--Microcline)

SHIMANSKIY, A.A.

Lower Cretaceous continental deposits of western Ciscaucasia.
Dokl. AN SSSR 147 no.3:679-682 N '62. (MIRA 15:12)

1. Krasnodarskiy filial Vsesoyuznogo neftegazovogo nauchno-issledovatel'skogo instituta. Predstavleno akademikom D.V. Kalivkinym.
(Caucasus, Northern—Geology, Stratigraphic)
(Caucasus, Northern—Gas, Natural—Geology)

SHIMANSKIY, A.A.; ALADATOV, G.M.; NIKIFOROV, B.M.

Formation and characteristics of the distribution of
gas-condensate pools in the Yeysk-Berezan' District
(Krasnodar Territory). Trudy KF VNII no.10:3-18 '62.
(MIRA 15:11)
(Krasnodar Territory--Condensate oil wells)

ALADATOV, G.M.; NIKIFOROV, B.M.; SHIMANSKIY, A.A.

Distribution of Pre-Cambrian, Paleozoic, Triassic, and Jurassic
sediments in western Ciscaucasia (Yeysk-Berezan' gas-bearing
region). Trudy KF VNII no.10:136-148 '62. (MIRA 15:11)
(Krasnodar Territory—Geology)

SHIMANSKIY, A.A.

Conditions governing the formation of the multilayered gas condensate pools of western Ciscaucasia. Neftegaz, geol. i geofiz. no.11: 8-14'63 (MIRA 17:7)

1. Krasnodarskiy filial Vsesoyuznogo neftegazovogo nauchno-issledovatel'skogo instituta.

SHARDANOV, A.N.; SHIMANSKIY, A.A.

Age of igneous rocks in western Ciscaucasia. Dokl. AN SSSR. 152
no.6:1441-1444 O '63. (MIRA 16:11)

1. Krasnodarskiy filial Vsesoyuznogo neftegazovogo nauchno-
issledovatel'skogo instituta. Predstavleno akademikom D.V.
Nalivkinym.

SHIMANSKIY, A.A.

Regularities in the distribution of the oil and gas pools in the Mesozoic sediments of western Ciscaucasia. Geol. nefti i gaza 8 no. 1:33-38 Ja '64. (MIRA 17:5)

1. Krasnodarskiy filial Vsesoyuznogo neftegazovogo nauchno-issledovatel'skogo instituta.

MATALIN, L.A.; SHIMANSKIY, A.M.; CHUBAROV, S.I.; SHTRANIKH, I.V.

1024-Channel time analyzer. Prib. i tekhn. eksp. no.3:54-63
My-Je '60. (MIRA 14:10)

(Neutrons) (Nuclear counters)

SHIMANSKIY, A. M., CAND TECH SCI, "MULTICHANNEL TIME
ANALYZERS ~~WITH~~ ^{with} A SYSTEM OF RECORDING ON MAGNETIC ELEMENTS."
MOSCOW, 1961. (ACAD SCI USSR, PHYS INST IN P. N. LEBEDEV).
(KL, 2-61, 214).

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ACCESSION NR: AR4014689

S/0271/64/000/001/B028/B029

SOURCE: RZh. Avtomatika, telemekhanika i vy*chislitel'naya tekhnika, 1964, no. 1, Abs. 1B211

AUTHORS: Meshkov, N. V.; Nesterov, P. V.; Smirnov, V. I. and Shimanskiy, A. M.

TITLE: Memory unit for a multidimensional analyzer with 16,000 channels

CITED SOURCE: Tr. 5-y Nauchno-tekhn. konferentsii po yadern. radio-elektronika. T. 2., Ch. 2. M., Gosatomizdat, 1963, 62-71

TOPIC TAGS: multidimensional analyzer, memory unit, ferrite core memory, high capacity memory, multidimensional analysis, information sorter

TRANSLATION: Multidimensional analysis requires a memory of great capacity. A 16000-cell ferrite core memory has been built which can be used directly as a memory for a multidimensional analyzer or as a sorter of information from a magnetic tape. In many cases the high capacity makes it possible to rewrite all the information from one or several tapes in one run and obtain graphic picture of the spectrum. The memory is intended for use with various units. A number of auxiliary functions added to the addressing system and the arithmetic block of the

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memory converts the system to a specialized computer capable of highly simplified operations in reducing information. The memory capacity is 128 X 128 sixteen-digit bits. The dead time of the memory is 200 microsec. The ferrite cores used in the memory are type K-260, measuring 2 X 1 and 3 X 1 mm. Selection half-current is 240 ma. The circuit includes 200 type 6N3P tubes, 100 6N6P tubes, 1200 semiconductor diodes, and 200 transistors. Orig. art. has 4 figs. and 4 refs. O. B.

SUB CODE: CP

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DATE ACQ: 19Feb64

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ACCESSION NR: AR4022434

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SOURCE: RZh. Fizika, Abs. 1A268

AUTHORS: Meshkov, N. V.; Nesterov, P. V.; Smirnov, V. I.; Shiman-
skiy, A. M.

TITLE: Memory unit for multidimensional 16000 channel analyzer

CITED SOURCE: Tr. 5-y Nauchno-tekhn. konferentsii po yadern. ra-
dioelektronike. T. 2, Ch. 2. M., Gosatomizdat, 1963, 62-71

TOPIC TAGS: memory unit, ferrite core memory, multidimensional
analyzer, magnetic tape data reduction, ferrite core matrix, reading
amplifier, magnetic commutator, address current

TRANSLATION: A 16,000 address ferrite-core memory has been developed.
This memory can be used in a multidimensional analyzer or serve as
a sorting block for the reduction of information from a magnetic

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tape. The memory block is made in the form of a 128 x 128 x 16 matrix of K-260 ferrite cores measuring 2 x 1.3 x 1 mm. To increase the reliability, the matrix, the reading amplifiers, and the magnetic commutator for the address currents are placed in a container where a $(35 \pm 1)^\circ\text{C}$ temperature is maintained. The memory capacity is 128 x 128 16-digit binary numbers. The dead time is 200 microseconds. The circuit contains 300 vacuum tubes and approximately 200 transistors. The operating principle is analyzed and block diagrams of the main units of the memory are presented. Yu. Semenov.

DATE ACQ: 03Mar64

SUB CODE: CP, SD

ENCL: 00

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