

87336

S/078/60/005/011/023/025/XX  
B004/B060

5.4110 2209, 1043, 1273

AUTHORS: Dombrovskaya, N. S., Alekseyeva, Ye. A.

TITLE: Methods of Decomposing Diagrams of the Composition of Multicomponent Systems According to the Indices of the Peaks of Prisms of the First Kind

PERIODICAL: Zhurnal neorganicheskoy khimii, 1960, Vol.5, No. 11, pp. 2612-2620

TEXT: The following definition is given: A prism of the  $r$ th kind is an  $n$ -dimensional polytope, formed by the parallel shift of an  $n-r$  dimensional polytope in independent directions in an  $n$ -dimensional space. The authors were concerned with the problem of studying such multicomponent systems as frequently arise in chemical technology. They proceeded from papers by N. S. Kurnakov (Ref. 1), A. G. Bergman, V. P. Radishchev (Refs. 2-5), which had dealt with the triangulation of diagrams of the chemical equilibrium and the search for singular stars. The singular stars of the chemical diagram constitute the geometrical representation of the chemical interaction between the components of the system. This is illustrated by Card 1/6

Methods of Decomposing Diagrams of the Composition of Multicomponent Systems According to the Indices of the Peaks of Prisms of the First Kind

B7336  
S/078/60/005/011/023/025/XX  
B004/B060

Fig. 1, a diagram of the reciprocal ternary system  $A, B \parallel X, Y$ . This is a cut through the tetrahedron of the quaternary system  $A - B - X - Y$ , brought about by the formation of the binary compounds  $AX, BX, AY, BY$ . The square formed by a plane cut is divided by the stable diagonal  $AY - BX$  into two stable cells, i.e., two triangles each of which represents a ternary system in the case of irreversibility of reaction. The exchange reaction is denoted in the square by the point  $O$  of conversion where the stable and the unstable diagonal intersect in conformity with the reaction  $AX + BY \rightarrow AY + BX$ . The stability of the diagonal is determined from the thermal data or the character of the liquidus surface. After a thorough description of the conventional method of decomposing complicated diagrams, which requires practice in spatial representation, the authors introduce their simplified method, as recommended in Ref. 10. Decomposition is performed on the basis of the peak indices. An index table is utilized for reciprocal systems of the  $2/n$  type. The first row is characterized by the natural series  $1, 2, 3, \dots, n$ , and the second row by the inverse series  $n, \dots, 3, 2, 1$ . Table 2 holds for the singular star in the system  $Na, K \parallel F, Cl, Br, I$  (Fig. 4):

Card 2/6

87336

Methods of Decomposing Diagrams of the Composition of Multicomponent Systems According to the Indices of the Peaks of Prisms of the First Kind

Indices of peaks

S	I	Br	Cl	F	$\Sigma$
Na	0	1	2	3	6
K	3	2	1	0	6
$\Sigma$	3	3	3	3	12

The pentatope which contains the zero peak NaI is cut off by means of the tetrahedron with the peak indices 1,2,3 of the first row and index 3 of the second row. This constitutes the tetrahedron NaF - NaCl - NaBr - KI. Pentatope NaF - NaCl - NaBr - NaI - KI is obtained. The tetrahedron for cutting off the pentatope with

the zero peak KF has the index 3 of the first row and the indices 3,2,1 of the second row. The common edge of the two tetrahedra has the largest indices 3 - 3, i.e., NaF - KI. The third tetrahedron, finally, has the edge 3 - 3 and two peaks with the remaining largest indices 2 - 2. The following stable diagonal tetrahedra are thus formed: 1) 123 - 3; 2) 23 - 32; 3) 3 - 321. The stable pentatope cells have the indices 1) 0123 - 3; 2) 123 - 32; 3) 23 - 321; 4) 3 - 3210. The stable cells are established by a nondiagonal transition from the largest index of the 1st row to the largest index of the 2nd row, as shown in Table 3. The same procedure is illustrated by the Li,Na,K,Rb,Cs,Cl,I system (Fig. 7). Table 6 gives

Card 3/6

Methods of Decomposing Diagrams of the Composition of Multicomponent Systems According to the Indices of the Peaks of Prisms of the First Kind

87336  
S/078/60/005/011/023/025/XX  
B004/B060

the indices, while Table 8 shows how to find the stable cells. There are 7 figures, 7 tables, and 11 references: 10 Soviet and 1 German.

SUBMITTED: October 9, 1959

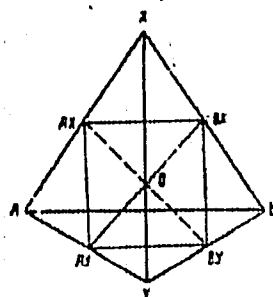


Fig. 1

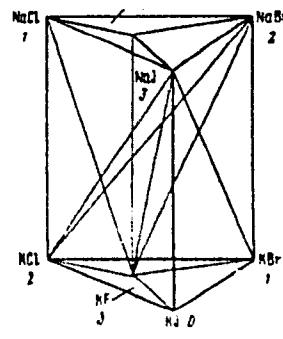


Fig. 4

Card 4/6

5.4110

68610

(3)

S/020/60/130/05/020/061

AUTHORS:

Dombrovskaya, N. S., Alekseyeva, Ye. A. B01/B005  
Khokhlova, N. V., Posypayko, V. I.

TITLE:

The Basal Tetrahedron  $1/2 Li_2SO_4 - NaCl - RbNO_3 - TlBr$  in the  
7-Component Reciprocal System  $Li, Na, Rb, Tl || Br, Cl, NO_3, SO_4$

PERIODICAL:

Doklady Akademii nauk SSSR, 1960, Vol 130, Nr 5, pp 1027-1029  
(USSR)

ABSTRACT:

The singular point of the reciprocal system of 16 salts  
 $Li, Na, Rb, Tl || Br, Cl, NO_3, SO_4$  (Ref 1) determining the  
direction of the exchange reactions is described. The position  
of the most stable basal tetrahedron  $1/2 Li_2SO_4 - NaCl - RbNO_3 -$   
 $TlBr$  was determined in the center of the cube orienting the  
singular point. Only 4 of its diagonals are fully stable:  
 $TlBr - RbNO_3$ ;  $TlBr - 1/2 Li_2SO_4$ ;  $RbNO_3 - NaCl$  and  $NaCl - 1/2 Li_2SO_4$ .  
The stability of the diagonal  $TlBr - NaCl$  is less certain since  
the solid solutions  $Tl(Br, Cl)$  and  $Na(Br, Cl)$  occur in the system  
 $Na, Tl || Br, Cl$ . M. N. Zakhvalinskiy (Ref 2) found the presence

Card 1/4

65610

The Basal Tetrahedron  $1/2 \text{Li}_2\text{SO}_4 - \text{NaCl} - \text{RbNO}_3 - \text{TlBr}$  S/O20/60/130/05/020/061  
in the 7-Component Reciprocal System Li, Na, Rb, BO11/BO05  
Tl||Br, Cl,  $\text{NO}_3$ ,  $\text{SO}_4$

of 2 complex compounds<sup>1</sup> on the diagonal  $\text{RbNO}_3 - 1/2\text{Li}_2\text{SO}_4$  in lithium- and rubidium salts. They are presumably:  $\text{Li}_2\text{SO}_4 \cdot \text{Rb}_2\text{SO}_4$  (1:1) and  $4\text{Li}_2\text{SO}_4 \cdot \text{Rb}_2\text{SO}_4$  (4:1). The base of the tetrahedron is formed by the ternary system  $1/2 \text{Li}_2\text{SO}_4 - \text{NaCl} - \text{RbNO}_3$ . Besides the 3 crystallization fields of the components, this system contains 2 additional fields which correspond to the binary compounds mentioned. Besides the 4 crystallization volumes of the components, the investigated part of the tetrahedron contains 2 relatively small volumes of the complex compounds of lithium- and rubidium sulfate (1:1 and 4:1). Rubidium sulfate is the exchange product between  $\text{Li}_2\text{SO}_4$  and  $\text{RbNO}_3$ . The 6 crystallization volumes meet in 2 quaternary points: the eutectic and the transition point lying in the "rubidium" corner of the diagram. Table 1 shows temperatures and compositions of the multiple points of the

Card 2/4

68610

The Basal Tetrahedron  $1/2 \text{Li}_2\text{SO}_4 - \text{NaCl} - \text{RbNO}_3 - \text{TlBr}$  S/020/60/130/05/070/061  
in the 7-Component Reciprocal System Li, Na, Rb,  $\text{NO}_3$ ,  $\text{SO}_4$ , Br, Cl,  $\text{NO}_3$ ,  $\text{SO}_4$  B011/B005

ternary systems and of the quaternary system. Figure 1 shows an evolvment, figure 2 a perspective representation of the tetrahedron. The composition of the ternary and quaternary eutectic and transition points was determined by graphic constructions; the temperatures were determined by recording the heating curves on the recording pyrometer of N.S.Kurnukov. In conclusion, the following can be said about the type of the 7-component system of 16 salts: the tetrahedron investigated determines the reaction direction in a way similar to the "basal" triangle in a quinary reciprocal system of 9 salts (Ref 3), and also similar to the stable diagonal triangles in a quaternary reciprocal system of 6 salts (Ref 4), and finally similar to the stable diagonal of the square of a ternary reciprocal system of 4 salts. By means of an experimental determination of the fusibility of the system  $1/2 \text{Li}_2\text{SO}_4 -$

Card 3/4

$-\text{NaCl} - \text{RbNO}_3 - \text{TlBr}$ , it was ascertained that the reciprocal

65610

The Basal Tetrahedron  $1/2 \text{Li}_2\text{SO}_4 - \text{NaCl} - \text{RbNO}_3 - \text{TlBr}$  S/020/60/130/05/020/061  
in the 7-Component Reciprocal System Li, Na, Rb, BO11/BO05  
TlBr, Cl,  $\text{FO}_3$ ,  $\text{SO}_4$

7-component system Li, Na, Rb, TlBr, Cl,  $\text{NO}_3$ ,  $\text{SO}_4$  may be ✓  
assigned to the class of reversible-reciprocal systems. There  
are 2 figures, 1 table, and 5 Soviet references.

ASSOCIATION: Vsesoyuznyy nauchno-issledovatel'skiy i konstruktorskiy  
institut khimicheskogo mashinostroyeniya (All-Union Scientific  
Research and Design Institute of Chemical Machine Construction)

PRESENTED: October 15, 1959, by I. I. Chernyayev, Academician

SUBMITTED: October 12, 1959

Card 4/4



83559

also 2308

18.1285

S/020/60/134/001/011/021  
B004/B060

AUTHORS: Andreyeva, V. V., Alekseyeva, Ye. A.

TITLE: Thin Oxide Films<sup>1</sup> on Titanium<sup>2,1</sup>, Zirconium<sup>2,1</sup>, Molybdenum<sup>2,1</sup>, and Titanium Alloys

PERIODICAL: Doklady Akademii nauk SSSR, 1960, Vol. 134, No. 1, pp. 106 - 109

TEXT: The authors wanted to study the kinetics of the oxidation of Ti, Zr, Mo, and the alloys Ti+10% Mo and Ti+3% Al+5% Cr in the temperature range 50-400°C. The thickness of the thin, invisible oxide films was measured by means of a previously described (Refs. 1,4,5) optical polarization method which is based on the ellipticity of reflected light. The oxidation took place in a tubular furnace in an oxygen stream. The thickness of the oxide films was measured every 60 min. Figs. 1,2 show the results of measurement; Table 1 lists the film thicknesses attained after 6 h. Zirconium is noticeably oxidized already at 50°C, titanium and the alloy Ti+10% Mo at 100°C, molybdenum and the alloy Ti+3% Al+5% Cr at 150°C. The oxidation is fast at the beginning, and gradually stops

Card 1/3

83559

Thin Oxide Films on Titanium, Zirconium,  
Molybdenum, and Titanium Alloys

S/O20/60/134/001/011/021  
B004/B060

as soon as the oxide film attains a certain thickness. The increase in film thickness,  $L$ , can be represented by equation (1):  $L = k \log(k_1 t + k_2)$ .

At high temperatures, however, the relation (2)  $L^2 = k_3 t$  (Fig. 3) holds.

For the metals and alloys examined, Table 1 supplies the temperature ranges in which the logarithmic or the parabolic law holds. Up to 800°C, the oxide film on titanium consists of  $TiO_2$  (rutile).  $TiO$  layers are formed above 800°C (immediately on the metal),  $Ti_2O_3$  (middle layer) and  $TiO_2$  (external layer). On zirconium,  $ZrO_2$  of a cubic structure forms up to 250°C, and  $ZrO_2$  of a monoclinic structure above 250°C. In the case of molybdenum, the oxide layer consists of  $MoO_3$  up to 300°C. Above 300°C,  $MoO_2$  is formed additionally due to dissociation. Fig. 3 further shows oxidation curves for chromium and aluminum, which are taken from the paper by T. N. Krylova (Ref. 9). These metals form only  $Cr_2O_3$  and  $Al_2O_3$ , respectively. The activation energy was calculated from the slope of the linear function  $\log k = f(1/T)$  (Table 1). The mechanism underlying

Card 2/3

8359

Thin Oxide Films on Titanium, Zirconium,  
Molybdenum, and Titanium Alloys

S/020/60/134/001/011/021  
B004/B060

the oxidation of zirconium and molybdenum differs from that of the oxidation of titanium and the alloy  $Ti+10\%Mo$ . A mixed oxide,  $TiO_2 + xMoO_3$ , is formed in the latter case. The resistivity to heat is thus improved. The high resistivity of the alloy  $Ti+3\%Al+5\%Cr$  to oxidation is explained by the formation of a mixed oxide of the spinel type. Fig. 4 shows the oxidation of titanium at 50-250°C in dry and moist, fully saturated air. In dry air, the oxide layer becomes twice as thick as in moist air. In the latter, however, the oxide layer is more compact and contains fewer defects. At 250°C, the water binding is loosened, and the thickness of the oxide layer attains the same value as in dry air. There are 4 figures, 1 table, and 12 references: 9 Soviet, 2 US, and 1 German.

ASSOCIATION: Institut fizicheskoy khimii Akademii nauk SSSR (Institute  
of Physical Chemistry of the Academy of Sciences USSR)

PRESENTED: April 28, 1960, by A. N. Frumkin, Academician

SUBMITTED: April 28, 1960

Card 3/3

DOMBROVSKAYA, N.S.; ALEKSEYEVA, Ye.A.

Methods of cleaving phase diagrams of multicomponent, reciprocal  
anhydrous salt systems for prisms of the 2nd kind, 3/3. Zhur.  
neorg. khim. 6 no.3:702-711 Mr '61. (MIRA 14:3)  
(Phase rule and equilibrium)  
(Systems(Chemistry))

POSTIPAYKO, V.I.; KHAKHLOVA, N.V.; ALEKSEYEVA, Ye.A.; DOMBROVSKAYA, N.S.

Singular decomposition of the polytope of the quintary reciprocal  
system consisting of nine salts: Na, K, Tl || Cl, Br, NO<sub>3</sub>.  
Zhur.neorg.khim. 6 no.6:1401-1407 Je '61. (MIRA 14:11)  
(Salts) (Systems (Chemistry))

15.6600

25503

S/065/61/000/007/004/005  
E030/E435

AUTHORS: Vinogradova, I.E., Alekseyeva, Ye.A.

TITLE: Thermographic investigation of E.P. (entrance pressure) additives in oils

PERIODICAL: Khimiya i tekhnologiya topliv i masel, 1961, <sup>6</sup>No.7, pp.56-61

TEXT: A differential thermocouple system has been applied to study the physical and chemical reactions of E.P. additives on heating, by themselves and in admixture with iron powder, to throw light on the reasons for their effectiveness. Of the thermocouples, one was placed in a beaker containing the additive or additive plus pure iron powder, and the other was placed in a beaker containing calcined magnesium oxide, noted for its absence of heating effects in the range investigated; the cold thermocouple junctions were in a Dewar flask of water at 18°C and the direct reading gave the absolute specimen temperatures. The oil used in testing was AC-14 (DS-14) and the additives were ХлорэФ-40 (Khloref-40) (butyl ether of trichlormethylphosphonic acid  $\text{CCl}_3\text{PO}(\text{OC}_4\text{H}_9)_2$ ), ГХС (GKHS) (hexachlorsulphide  $[\text{CCl}_3(\text{CH}_2-\text{CH}_2)_2]_2\text{S}$ ).

Card 1/2

25503

S/065/61/000/007/004/005  
E030/E435

Thermographic investigation of ...

J13-6/9 (L3-6/9) (ethylene dibutylxanthate ( $C_4H_9OCS_2-CH_2)_2$ ) and chlorinated paraffins (mixtures from  $C_{25}H_{51}Cl$  to  $C_{25}H_{40}Cl_{12}$ ). All the additives gave endothermic effects on boiling and, where relevant, on melting, either by themselves or on addition of powdered iron (1:2.5 by weight). The butyl ether also gave an exothermic effect on decomposition at  $240^\circ C$ ; one at  $135^\circ C$  with iron powder was verified by repeat experiments at lower iron concentrations to be reaction with the iron. Similarly, the hexachlorosulphide reacted with iron at  $153^\circ C$  and the dibutylxanthate at  $224$  to  $238^\circ C$ . Chlorinated paraffins scarcely react with iron but the iron catalyses their decomposition, reducing the decomposition from  $325$  to  $285^\circ C$ . All these reaction temperatures are below those generated during boundary friction accompanied by wear, thus confirming the anti-friction properties of the additives. A strong correlation is claimed to exist between the degree of wear reduction and the magnitude of the exothermic effect on reaction with iron powder. There are 6 figures, 2 tables and 2 Soviet references.

ASSOCIATION: VNII NP

Card 2/2

ALIKSEYEVA, Ye.A.; DOMBROVSKAYA, N.S.

Dividing the composition diagram of the septenary reciprocal  
system consisting of 20 salts A, B, C, D || X, Y, Z, T of the 16 G  
type. Zhur.neorg.khim. / no.9:2158-2162 S '61. (MIRA 14:9)  
(Salts) (Systems (Chemistry))



DOMBROVSKAYA, N.S.; KHAKHLOVA, N.V.; ALEKSEYEVA, Ye.A.

Intersection between a stable and a nonequilibrium tetrahedron in the  
septenary reciprocal system Li, Na, Rb, Tl Br, Cl, NO<sub>3</sub>, SO<sub>4</sub>. Dokl.  
AN SSSR 137 no.6:1361-1363 Ap '61. (MIRA 14:4)

1. Vsesoyuznyy nauchno-issledovatel'skiy i konstruktorskiy institut  
khimicheskogo mashinostroyeniya. Predstavleno akademikom I.V.  
Tamanayevym.

(Systems (Chemistry))

S/883/62/000/000/016/020  
E194/E155

AUTHORS: Vinogradova, I.E., Alekseyeva, Ye.A., and Kulagina, S.S.

TITLE: Temperature methods of assessing the properties of  
E.P. oil

SOURCE: Metody ispytaniya na iznashivaniye; trudy soveshchaniya,  
sostoyavshegosya 7-10 dek. 1960. Ed. by .  
N.M. Khrushchov. Moscow, Izd-vo AN SSSR, 1962. 164-175

TEXT: Point-contact friction machine tests are simple and  
sensitive to the effects of E.P. additives, although information  
is generally not available about the actual temperatures on the  
friction surfaces, except in four-ball machine type KT -2 (KT-2),  
where the rubbing speeds are low and the oil is assessed by the  
critical temperature at which the oil film breaks down. In  
conventional four-ball machines the effects are more complicated  
and it is recommended to assess the contact surface temperature by  
study of structural changes in the surface layers of the metal.  
A study was made of the microhardness distribution near the wear  
scar of sectioned balls from the four-ball machine. The temperature  
distribution was estimated by interpolation of microhardness  
Card 1/2

Temperature methods of assessing...

S/883/62/000/000/016/020  
E194/E155

results on the tempering curve of the steel in question. The differing effects of chlorine- and sulphur-containing additives on the load/temperature characteristics at seizure were determined in this way. Most sulphur additives reduce the temperature of the friction surfaces, whilst chlorine additives prevent welding. Thermographic analysis is a most sensitive procedure for studying physical and chemical processes but has been little used in studying E.P. oil. It was accordingly used to judge of changes in the aggregate state from inflection points on the heating or cooling curves, which correspond to endothermic or exothermic reactions. The results were compared with those obtained in four-ball machines. Test results are quoted for a number of sulphur- and chlorine-containing additives in oils, both with and without iron powder. It is, of course, necessary to separate the reactions between additives and iron from those corresponding to evaporation or thermal decomposition of the additive. It is desirable to check the reaction between additives and iron up to temperatures above the highest bulk oil temperature and below the seizure temperature, i.e. in the range 150 to 250 °C.

Card 2/2      There are 8 figures and 2 tables.

ALEKSEYEVA, Ye.A.; DOMBROVSKAYA, N.S.

Interaction of salts in the five-component reciprocal system Li, Rb,  
Ti||Br, NO<sub>3</sub>, SO<sub>4</sub>. Zhur.neorg.khim. 7 no.7:1659-1665 J1 '62.  
(MIRA 16:3)

(Systems (Chemistry))

(Salts)

DOMEROVSKAYA, N.S.; ALEKSEYEVA, Ye.A.

Singular star of the binary reciprocal system consisting of  
12 salts: Li, Rb, Tl, Br, Cl, NO<sub>3</sub>, SO<sub>4</sub>. Zhur. neorg. khim. 7  
no.8:2002-2012 Ag '62. (MIRA 16:6)

(Systems(Chemistry))  
(Fused salts)

DOMBROVSKAYA, N.S.; ALEKSEYEVA, Ye.A.

Completeness of interaction and thermochemical relations in  
the quinary reciprocal system  $Li, Rb, Tl || Br, NO_3, SO_4$ .  
Zhur.neorg.khim. 7 no.12:2801-2805 D '62. (MIRA 16:2)  
(Systems (Chemistry)) (Thermochemistry)

156600

2209

32397  
S/080/62/035/001/009/013  
D245/D304

AUTHORS: Vinogradova, I. E., and Alekseyeva, Ye. A.

TITLE: Study of the stability and reactivity of derivatives  
of chlorophosphinic acids used as anti-wear additives  
in oils

PERIODICAL: Zhurnal prikladnoy khimii, v.35, no.1, 1962, 176-182

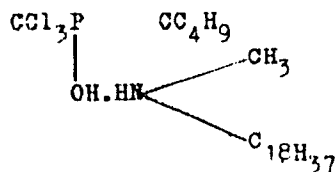
TEXT: The authors used thermographic analysis to study the behavior of chlorophosphinic acid derivatives used as anti-wear additives to gear box oil. Samples were heated to high temperatures slowly with and without addition of powdered Fe. Wear tests were carried out in accordance with GOST 9490-60. It is shown that additives which impart high anti-wear properties to oil undergo a marked exothermic reaction with Fe when heated in this way. The additive "chlorefamin" showed the most satisfactory anti-wear properties at the temperatures studied and had no corrosive effect. It consists of the methyl-octadecylamine salt of butoxytrichloromethyl-phosphinic acid: X

Card 1/2

Study of the stability ...

32397

S/080/62/035/001/009/013  
D245/D304



X

This additive does not react with Fe until a temperature of 178°C is reached. (In a gear box, the oil usually reached a maximum temperature of 150°C.) There are 4 figures, 2 tables and 5 Soviet-bloc references.

SUBMITTED: February 17, 1961

Card 2/2



DOMEROVSKAYA, N.S.; ALEKSEYEVA, Ye.A.

New types of the singular stars of senary reciprocal systems.  
Izv. neorg. khim. 9 no.5:1266-1271 My '64.

KOSTENKO, Mikhail Poliyevktovich; PIOTROVSKIY, Lyudvik  
Marianovich; ANEMPODISTOV, V.P., nauchn. red.; ALEKSEYEVA,  
Ye.A., red.

[Electrical machines] Elektricheskie mashiny. Moskva,  
Energia. Pt.2. 1965. 703 p. (MIRA 18:11)

L 15281-66		ACC NR. AP5028963		SOURCE CODE: UR/0119/64/009/009/0025/0027	
AUTHOR: Alekseyeva, Ye. A. (Engineer); Grudov, A. P. (Engineer); Il'in, Ye. P. (Engineer); Konovalova, I. K. (Engineer); Maksimova, O. V. (Engineer); Sharenko, M. A. (Engineer)					
ORG: none					
TITLE: Effect of temperature on elastic properties of thin-sheet spring alloys					
SOURCE: Priborostroyeniye, no. 9, 1964, 25-27					
TOPIC TAGS: spring, measuring instrument, industrial instrument					
ABSTRACT: The results are reported of measurements of the elastic limit $\sigma$ (with residual strains of 0.01 and 0.005%) and elasticity modulus E in bending of 85-120-micron thick specimens (10x100 mm) of BrOF6, 5-0, 15; BrKMTs 3-1; BrBz, BrBNT 1, 9 bronzes, 60SZ, H1814 steels, and N36KhTYuM8 alloy at temperatures that ranged from -70C to +150 or +500C. Also, the ultimate strength $\sigma_u$ and the yield point $\sigma_s$ of 0.1x10-mm 57-mm long specimens were determined. All specimens were thermally treated according to specifications normally used in the					
Card 1/2		UDC: 620.172.22:62-415:536.49			

L 15284-55

ACC NR. AP5028963

instrument-making industry. The numerical findings are reported in the form of curves. It is noted that many specimens suffered brittle fractures partly due to their thickness nonuniformity and high width-to-thickness ratio. Orig. art. has: 7 figures and 1 table.

SUB CODE: 11, 13 / SUBM DATE: none / ORIG REF: 004

Card 2/2

7/1 Q.S

DOMBROVSKAYA, N.S.; POSYPAYKO, V.I.; ALEKSEYEVA, Ye.A.; KHACHLOVA, N.V.

Stable elements of hepta-component reciprocal systems. Dokl.  
AN SSSR 165 no.5:1081-1084 D '65.

(MIRA 19:1)

1. Submitted May 13, 1965.

ALEKSEYEVA, Ye. F.; KIRILLOV, V. V.; LYATKOVSKAYA, N. M.; MALYSHEVA, T. D.;  
CHESLOV, V. M.; STEPANOV, A. S.; KHROPOVA, P. M.; CHERNENKO, M. I.;  
GRAMMANOV, A. G., prof., red.; SMIRNOV, P. S., tekhn. red.

[Manual on exercises in physics] Posobie k uprazhneniam po fizike.  
Leningrad, Leningr. elektrotekhn. in-t im. V. I. Ul'ianova (Lenina).  
Part. 1. [Mechanics. Molecular physics] Mekhanika, Molekuliarnaya  
fizika. Sost. E. F. Alekseeva i dr. 1960. 75 p. (MIRA 14:10)  
(Physics—Problems, exercises, etc.)

ALEKSEYEV, Ye. I.

АЛЕКСЕЕВ, Е. И.

Free transplant of bones into infected tissues. Vest.khir.  
74 no.1:45-51 Ja-P '54. (MIRA 7:2)

1. Iz nauchno-issledovatel'skogo instituta khirurgicheskogo  
tuberkuleza (nauchnyy rukovoditel' - professor P.G.Kornev).  
(Bones--Wounds and injuries)

ALEKSEYEVA, Ye. I.

ALEKSEYEVA, Ye. I.: "The Free Transplantation of Bone into Infected Tissue in Order to Treat and Avoid 'False Arthroses' From Firearm Wounds." Leningrad State Order of Lenin Inst for the Advanced Training of Physicians imeni S. M. Kirov. Leningrad, 1956. (Dissertation for the Degree of Candidate in Medical Science)

So: Knizhnyy letopis', No. 18, 1956.



~~ALEKSEYEV~~, Yeliseveta Ivanovna; BOGDANOVA, Mariya Semenovna; LIL'KHO,  
S.I., redaktor; KUDREVA, M.S., tekhnicheskii redaktor.

[Work therapy in hospitals; knitting] Trudoterapiia v lechebnykh  
uchrezhdeniyakh : viazanie na spitsakh. [Leningrad] Gos.izd-vo  
med.lit-ry, Leningr.otsk-mis, 1956. 44 p. (MIRA 10:6)  
(Knitting)



[illegible]

ALERSHAYNA, Ye. L.

"Anatomical Structure of Green Grafts of Non-Valley L. in the Root-Joining Process," Dokl. AN SSSR, 62, No. 3, 1962.

All-Union Sci. Res. Inst. Essential Oil Cultures.

1. D'TACHEN, I. I., ALIKSEYEV, YE. I.
2. USSR (600)
4. Nicotine
7. Problem of the synthesis of nicotine in the tobacco plant. Tabak.  
15 no. 5, 1952

9. Monthly List of Russian Accessions, Library of Congress, January 1953, Unclassified.

ALEKSEYEVA, Ye.I., kand. sel'khoz. nauk; BUZINOV, P.A., kand. sel'khoz. nauk; VODOLAGIN, V.D.; VOLKHOVSKAYA, U.V.; GLUSHCHENKO, N.N., kand. biol. nauk; GURVICH, N.L., doktor biol. nauk; ZHELEZNOV, P.A., kand. sel'khoz. nauk; KSENDZ, A.T.; LESHCHUK, T.Ya.; LUK'YANOV, I.A., kand. sel'khoz. nauk; MAYCHENKO, Z.G., kand. sel'khoz. nauk; TANASIYENKO, F.S., kand. khim. nauk; ZNAMENSKIY, M.P.; PERSIDSKAYA, K.G.; PODLESHOVA, A.F.; ROGOCHIIY, I.Ya.; REZNIKOV, A.R.; SHUL'GIN, G.T.; KHOTIN, A.A., doktor sel'khoz. nauk; LAPSHINA, O.V., red.; KINENKOVA, V.R., red.; MAKINOVA, N.N., tekhn. red.; BALLOD, A.I., tekhn. red.

[Aromatic plants] Efiromaslichnye kul'tury. Moskva, Sel'-khozizdat, 1963. 358 p. (MIRA 16:12)  
(Ukraine--Aromatic plants)

PUSTOVOYT, V.S., akademik, red.; SUSLOV, V.M., kand. ekon. nauk, otv. red.; ALEKSEYEVA, Ye.I., kand. sel'khoz. nauk, red.; BUZINOV, P.A., red.; VASIL'YEV, D.S., kand. sel'khoz. nauk, red.; VOSKRESENSKAYA, G.S., red.; GUNDAYEV, A.I., red.; IGNAT'YEV, B.K., kand. sel'khoz. nauk, red.; MAKSHOVA, A.Ya., red.; MOSKALENKO, V.I., red.; PANCHENKO, A.Ya., red.; TIKHONOV, O.I., red.; SHPOTA, V.I., kand. sel'khoz. nauk, red.; MONOVA, Ye.S., red.; LAPSHINA, O.V., red.

[Oilseed and aromatic crops; transactions for 1912-1926]  
Maslichnye i efiromaslichnye kul'tury; trudy za 1912-1962 gg. Pod obshchey red. V.S.Pustovoita. Moskva, Sel'khozizdat, 1963. 575 p. (MIRA 17:4)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut maslichnykh i efiromaslichnykh kul'tur. 2. Vsesoyuznaya akademiya sel'skokhozyaystvennykh nauk imeni V.I.Lenina (for Pustovoyt). 3. Direktor Vsesoyuznogo nauchno-issledovatel'skogo instituta maslichnykh i efiromaslichnykh kul'tur (for Suslov).

GLUSHCHENKO, N.M., kand. sel'khoz. nauk; ALERSEYEVA, Ye.I., kand.  
sel'khoz. nauk; VOLOB'YEVA, G.V.; IUSINA, L.V., kand.  
biol. nauk; PACHENKO, E.G., CHIKALOV, B.M., kand. sel'-  
khoz. nauk; KRYLATOVA, S.A., red.

[Recommendations for the production of aromatic plant seeds]  
Rekomendatsii po semenovodstvu efiroaslichnykh kul'tur. Mo-  
skva, Sel'khozizdat, 1963. 27 p. (MIRA 17:6)

1. Russia (1973- U.S.S.R.) Ministerstvo sel'skogo khozyay-  
stva. Upravleniye nauki, propagandy i vnedreniya peredovogo  
opyta. 2. Nauchnyye sotrudniki Vsesoyuznogo nauchno-  
issledovatel'skogo instituta maslichnykh i efiroaslichnykh  
kul'tur. (for all except Krylatova).



S/080/62/035/006/002/013  
D204/D307

AUTHORS: Andreyeva, V. V. and Alkeseyeva, Ye. L.

TITLE: The oxidation of titanium, zirconium, molybdenum  
and certain alloys.

PERIODICAL: Zhurnal prikladnoy khimii, v. 35, no. 6, 1962,  
1175-1183

TEXT: The rates of oxidation of massive Ti, Mo, Zr, 90Ti10Mo and 92Ti15Cr3Al were studied in oxygen and in air, between 20 and 400°C. Progress of oxidation was assessed by the thickness of oxide scale formed, as measured by an optical polarization method. At room temperature, in air, Ti oxidized logarithmically, the scale reaching a thickness of 80 - 90 A.V. after 545 days. Between 50 and 250°C Ti oxidized faster in dry than in humid air. The oxidation of Zr was already appreciable at 50°C, of Ti and 90Ti10Mo alloy at 100°C and of Mo and 92Ti15Cr3Al at 150°C. At 150°C the scales on Ti, Mo and Zr reached 30 - 50 A.V. after 6 hours and were 250 - 400 A.V. thick after 6 hours at 250 - 350°C. Zirconium

Card 1/2

The oxidation of titanium ...

S/080/62/035/006/002/013  
D204/D307

oxidized more rapidly than Mo at 250°C or than Ti at 350°C. This was associated with the growth of monoclinic  $ZrO_2$ . Mo oxidized rapidly above 300°C but an admixture of 10% Mo and Ti made this alloy better than Ti alone at the same temperature. The TiCrAl alloy was particularly resistant to oxidation. In general the metals oxidized logarithmically at lower and parabolically at higher temperatures. The rate of oxidation is thought to be governed initially by the rate of formation of a thin scale as the temperature is increased. The scales were found to be rutile for Ti, cubic  $ZrO_2$  up to 250°C and monoclinic  $ZrO_2$  above 250°C for Zr,  $MoO_3$  up to 300°C and  $MoO_3 + MoO_2$  above 300°C for Mo, solid solution of  $TiO_2$  and  $MoO_3$  for 90Ti10Mo and (probably) a solid solution of Ti, Cr, Al oxides for the ternary alloy. Energies of activation for the oxidation processes ranged from 10 to 21.2 kcal/mole between 200 and 350°C. The results are discussed. There are 8 figures and 2 tables.

- SUBMITTED: May 12, 1961

Card 2/2

DOC NO:	AP-013787	DOC CASE:	DA/0000/65/000/000/0043/0058
AUTHOR:	Andreyev, V. V.; Kazarin, V. I.; Alekseyeva, Ye. L.; Glasunov, S. G.; Solovina, O. P.; Kikilova, V. P.		
ORIG:	USSR		
TITLE:	Study of the corrosion resistance and electrochemical and mechanical properties of alloys of the titanium-niobium system		
SOURCE:	Korroziya metallov i spлавov (Corrosion of metals and alloys), no. 2 Moscow, Izd-vo Metallurgiya, 1963, 43-58		
TOPIC TAGS:	corrosion resistance, electrochemistry, titanium containing alloy, niobium containing alloy, acid, metal heat treatment		
ABSTRACT:	This is a continuation of a previous investigation (this issue, pp 29-42) with the difference that it deals with alloys of the Ti-Nb system containing up to 50% wt. Nb. Both metals in unalloyed state have a high corrosion resistance, but in certain solutions, e.g. sulfuric and hydrochloric acid solutions, Ti dissolves at a sufficiently fast rate whereas Nb remains corrosion-resistant. Hence, the addition of Nb to Ti should increase the corrosion resistance of Ti. Mechanical tests of these alloys show that as the Nb content increases (up to 5%) the ultimate strength of the alloy increases from 57 kg/mm <sup>2</sup> to 92 kg/mm <sup>2</sup> ; as the Nb content is further		
Card	1/2		

1. 28.06-66

ACC NR. AT6013787

increased above 51, however, ultimate strength decreases; a similar pattern of variation with Nb content is displayed by plasticity and hardness. In 10, 40, 60, 75 and 94% solutions of  $H_2SO_4$ , the alloys at 40°C, whether in hot-forged state or after heat treatment (heating at 920-950°C for 1 hr, water quenching, aging at 450°C for 10 hr with cooling in air), display a general increase in corrosion resistance with increase in Nb content. A similar pattern, on the whole, is observed when the alloys are placed in  $HPO_3$ ,  $HCl$ ,  $HNO_3$ , and oxalic acid. For the alloys containing upward of 30% Nb, however, aging leads to decomposition of the  $\beta$ -phase, which deteriorates their corrosion resistance. Plotting of the curves of variation in current density as a function of the specified potentials (starting with -0.8 v and ending with +2.2 v) showed that the maximum corrosion rate corresponds to a potential of -0.25 v. As the Nb content of the alloys increases, the critical density of the passivation current decreases and the normal hydrogen potential shifts in the direction of more positive values. The addition of Nb to Ti enhances the corrosion resistance of Ti in solutions of non-oxidizing acids and does not affect the high corrosion resistance of Ti in oxidizing solutions such as 57%  $HNO_3$  or a mixture of  $HNO_3$  and  $HCl$  in the ratio of 1:1 or 2:1 at 100°C. Orig. art. has: 9 figures, 5 tables.

SUB CODE:

07.12

SUBJ DATE: 19JUL65/

ORIG REF: 003/

Card 2/2

47367-66 EWT(m)/EMP( )/I/EMP( )/ETI( )/E(c) ID/IG/WB  
ACC NR AR60284 SOURCE CODE: UR/0137/66/000/005/1085/1085

AUTHOR: Andreyeva, V. V.; Kazarin, V. I.; Alekseyeva, Ye. L.; Glazunov, S. G.; Nikulova, V. F.; Solonina, O. P.

TITLE: Investigation of the corrosion resistance and electrochemical and mechanical properties of alloys of the system niobium and titanium

SOURCE: Ref. zh. Metallurgiya, Abs. 51590

REF SOURCE: Sb. Korroziya met. i splavov. No. 2, M., Metallurgiya, 1965, 43-58

TOPIC TAGS: niobium titanium alloy, corrosion resistance/Ti20Nb alloy

ABSTRACT: Titanium alloys with 2--50% niobium have been investigated. Alloying of titanium with niobium considerably increases  $\sigma_s$  and  $H_B$  of Ti. Thus, after hot forging the Ti-20Nb alloy has  $\sigma_s$  of  $\sim 104 \text{ Mn/m}^2$  (Ti  $60 \text{ mn/m}^2$ ),  $\delta \sim 11\%$  (Ti  $\sim 18\%$ ). The corrosion resistance of alloys in solutions of unoxidative acids is considerably higher than that of titanium. In such acids as  $\text{HNO}_3$ , the resistance of titanium and titanium-niobium is identical. The critical density of passivating current decreases with an increase of niobium content in

Card 1/2

UDC: 669.295.5

L 47367-55

ACC NR: AR6028442

alloys, and the stationary potential shifts to more positive values. [Translation  
of abstract] [NT]

SUB CODE: 11/

Card 2/2 afs

"APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6

APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6"

ALIKSEYEV, Y. N.

In memory of the Academician Vladimir Mikhailovich Rodionov. Soob.o  
nach.rub.chl.VKHO no.4:1-4 '54. (MIRA 10:10)  
(Rodionov, Vladimir Mikhailovich, 1878-1954)



TIKRONOV, G.A., dots., red.; ~~ALEXSEYEV~~ ALEXSEYEV, Ya.N., red.; VORONTSOVA,  
Z.Z., tekhn. red.

[Automatic metering and control devices] Avtomaticheskie  
ustroystva ucheta i kontrolya; sbornik statei. Izhevsk,  
Udmurtskoe knizhnoe izd-vo, 1963. 43 p. (MIRA 17:3)

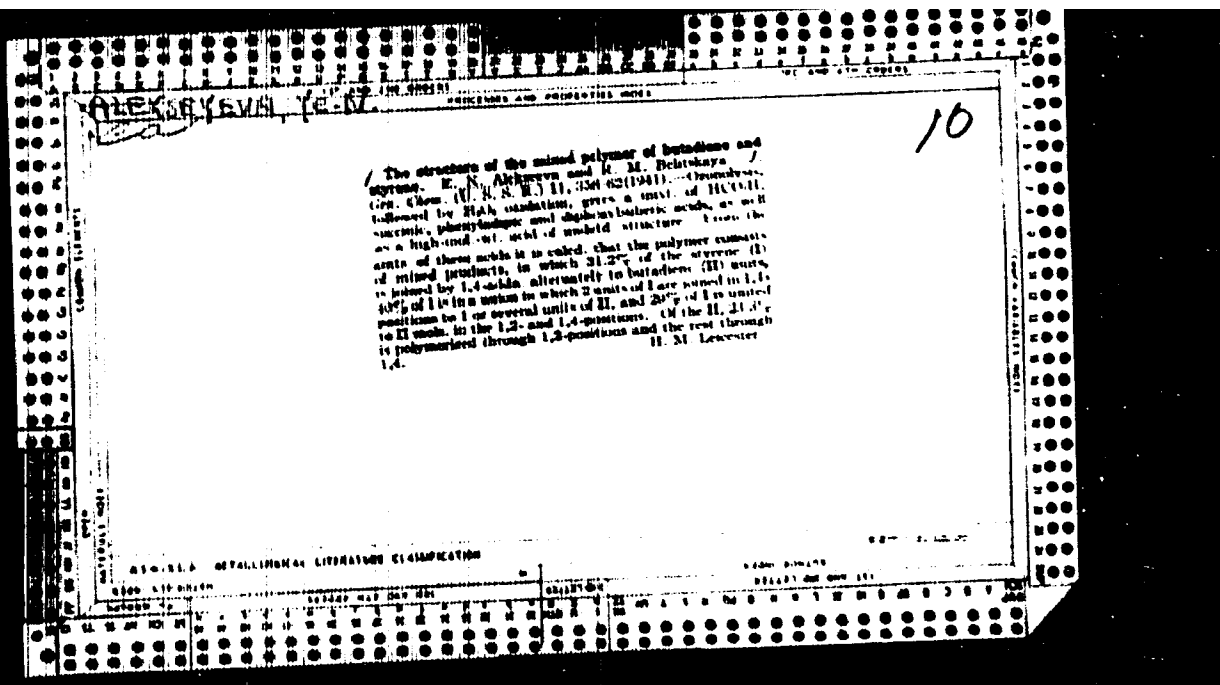
10

1A

STRUCTURE OF THE MIXED POLYMER OF BUTADIENE AND  
ACRYLONITRILE. E. N. Akhmedov. *Izv. Akad. Nauk SSSR Khim. Nauk* 1960, 1420-30 (1960). An equimolar mix of butadiene  
(I) and acrylonitrile (II) contg. 1% HCl, II was poly-  
merized at 60° for 116 hrs. and the product was vacuum-  
distilled at 60° for 116 hrs. and the polymer was dist. by the  
Kjeldahl method. The polymer was oxidized and oxidized  
with H<sub>2</sub>O<sub>2</sub> by the method of Putnam et al. (J. Am. Chem. Soc.  
70, 1948 (1948)). The resulting carboxylic acids were methylated  
with MeOH contg. 10% HCl and the distn. fractions  
analysed, giving succinic, butanetricarboxylic, hexane-  
tetracarboxylic and dodecanetricarboxylic acids in  
definite proportions, indicating that about one half of the  
polymerization product is formed by the alternate co-  
polymerization of I and II in the mix. About one third of the  
polymer consists of the units in which I and II corresponds  
to 2 or 3 mols. of II.  
Chas. Blum

ASAC-11.4 METALLURGICAL LITERATURE CLASSIFICATION

1000 1100 1200 1300 1400 1500 1600 1700 1800 1900 2000 2100 2200 2300 2400 2500 2600 2700 2800 2900 3000 3100 3200 3300 3400 3500 3600 3700 3800 3900 4000 4100 4200 4300 4400 4500 4600 4700 4800 4900 5000 5100 5200 5300 5400 5500 5600 5700 5800 5900 6000 6100 6200 6300 6400 6500 6600 6700 6800 6900 7000 7100 7200 7300 7400 7500 7600 7700 7800 7900 8000 8100 8200 8300 8400 8500 8600 8700 8800 8900 9000 9100 9200 9300 9400 9500 9600 9700 9800 9900



29

АЛЕКСЕЕВА, Е.Н.

718. The Structure of a Mixed Polymer of Butadiene and Acrylonitrile. E. N. Alekseeva. *Rubber Chemistry and Technology*, v. 20, Oct. 1947, p. 687-692. Translated from *Journal of General Chemistry (U.S.S.R.)*, v. 15, no. 1948, p. 1426-1430.

Gives results of an experimental polymer as determined by ozonolysis and investigation of the products.

ASR-51.4 DETAILING LITERATURE CLASSIFICATION

ASR-51.4 DETAILING LITERATURE CLASSIFICATION

ASR-51.4 DETAILING LITERATURE CLASSIFICATION

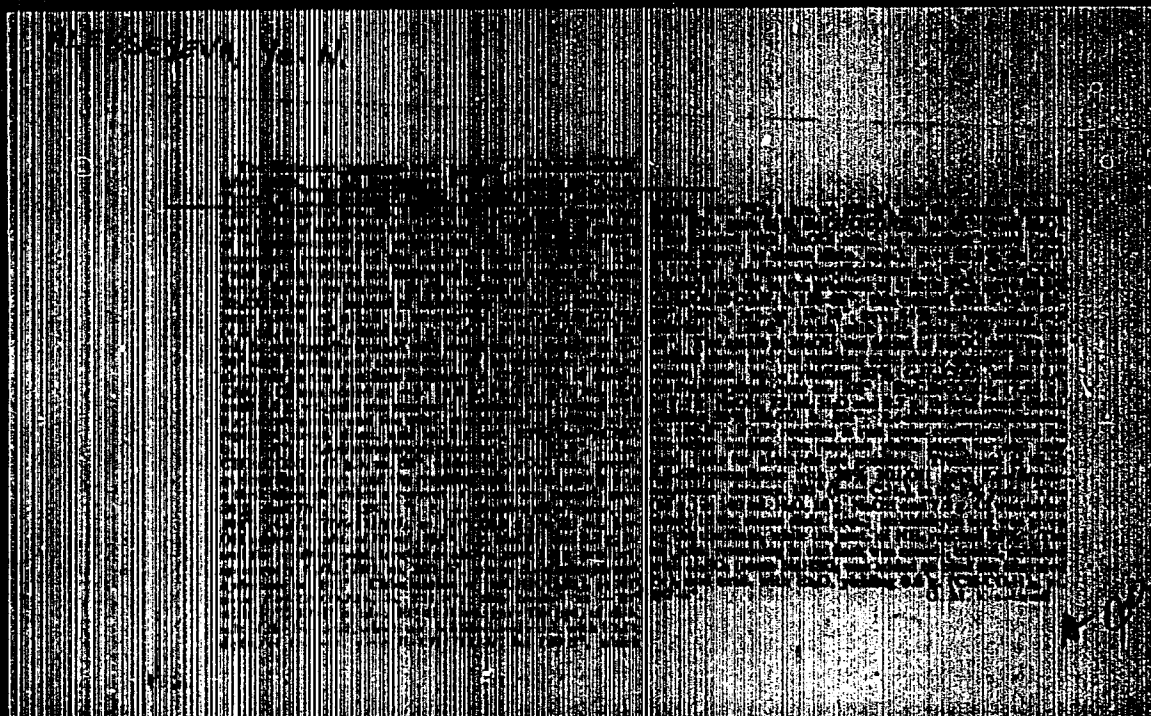
ALEKSEYEV, E. N.

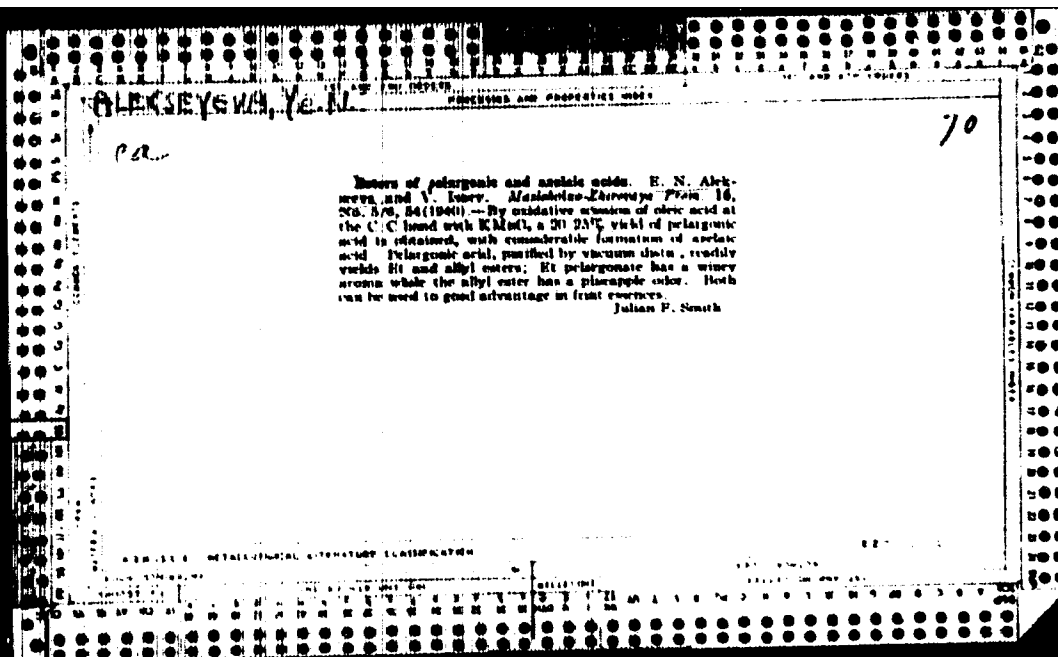
Chemical Abst.  
Vol. 48 No. 8  
Apr. 25, 1954  
Organic Chemistry

7  
(2) 1176  
The structure of butadiene and acrylonitrile copolymers.  
Synthesis of butadiene-triacetylenic acids. E. N. Alekseyev  
and V. D. Moshov. J. Gen. Chem. (U.S.S.R.) 23, 1851-4  
(1953) (Engl. translation). - See C.A. 47, 5357d.

H. L. H.

11-17-54





AUTHORS: Alekseyeva, Ye. E., Vavov, V. A. SOV/156-58-3-37/52

TITLE: The Deposition of Phthalimide on the Esters of  $\alpha$ -,  $\beta$ -Unsaturated Acids (Prisoyedineniye ftalimida k estiram  $\alpha$ -,  $\beta$ -nenasyshchennykh kislot)

PERIODICAL: Nauchnyye doklady vysshey shkoly, Khimiya i khimicheskaya tekhnologiya, 1958, Nr 3, pp. 545 - 549 (USSR)

ABSTRACT: The possibility of the use of the catalyst of Rabinov was investigated in the deposition of phthalimide on the double bonds of the alkylidenemalononic and alkylidenecetonic esters, the esters of cinnamic acid and benzene acetone. The reaction takes place effectively for the ethylidene-, propylidene-, isopropylidene-, and benzylidene-malononic esters. The acid hydrolysis of the phthalimide derivatives of the alkylidene malonic esters leads to the corresponding  $\beta$ -amino acids. By employing this method  $\beta$ -aminoisocaproic acid,  $\beta$ -aminobutyric acid,  $\beta$ -aminovaleric acid, and  $\beta$ -amino- $\beta$ -phenylpropionic acid were synthesized for the first time. There are 2 tables and 6 references, 1 of which is Soviet.

Card 1/2



The Deposition of Phthalimide on the Esters of  $\alpha$ -,  
 $\beta$ -Unsaturated Acids

SOV, 156-50-3-37/52

ASSOCIATION:

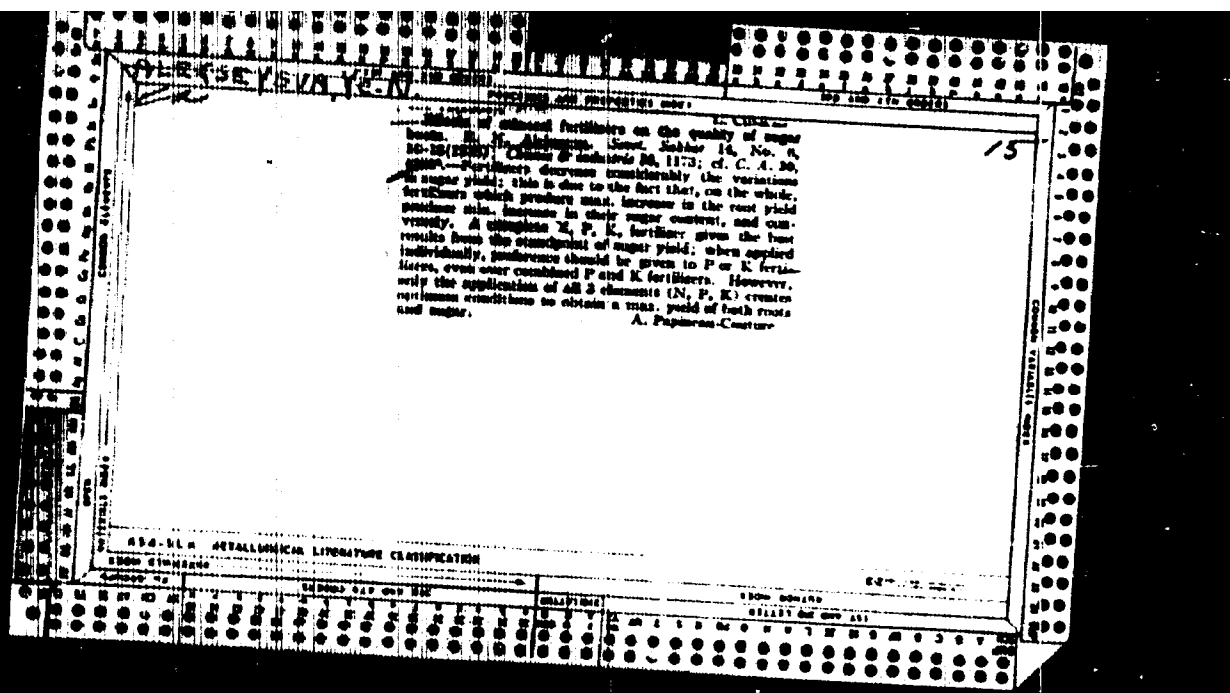
**Kafedra** organicheskoy khimii Mo Kovskogo khimiko-  
tehnologicheskogo instituta im.D.I.Mendeleyeva (Chair of In-  
organic Chemistry at the Moscow Chemical and Technological  
Institute imeni D.I.Mendeleyev)

SUBMITTED: November 19, 1957

Card 2/2

POKROV, Sergey Pavlovich; GIKH, V. M., ed.

[Semiconductor rectifiers for charging storage bat-  
teries] Poluprovodnikovye vypryamiteli dlia zaryadki  
akkumulatorov. Izd. 2., ispr. i dop. Izhevsk,  
Udmurtskoe knizhnoe izd-vo, 1962. 55 p. (MIRA 17.6)





1. ALEKSEYEV, YE.N.

2. USSR (600)

4. Fertilizers and Manures

7. Action of fertilizers in grassland crop rotation on deep moderately leached chernozem, Sov.Agron. 11 no. 4, 1953.

9. Monthly List of Russian Accessions, Library of Congress, APRIL 1953, Uncl.

ALMESKYEVA, Ye. H.

Foliar nutrition of sugar beets. Sakh.prom.29 no.5:29-42 '55.  
(MIRA 8:11)

1. Ramenskaya opytno-selektcionnaya stantsiya  
(Sugar beets) (Fertilizers and manures)

SHIROBOKOV, Stepan Ivanovich; ALEKSEYEVA, Ye.N., red.; VORONTSOVA, Z.Z.,  
tekhn. red.

[The Udmurt A.S.S.R.; economic and geographical features] Udmurt-  
skaya ASSR; ekonomiko-geograficheskaya kharakteristika. Izhevsk,  
Udmurtskoe knizhnoe izd-vl, 1961. 214 p. (MIRA 15:1)  
(Udmurt A.S.S.R.—Economic geography)

BURKSER, E.S. [Burksar, IE.S.]; ALEKSEYEVA, Ye.N. [Alekseieva, K.M.];  
VETSHTEYN, V.Ye.; GOL'DENFELD, I.V.; DAVYDYUK, L.A. [Davydyuk, L.O.];  
DEMIDENKO, S.G. [Demydenko, S.H.]; YELISEYEVA, G.D. [Eliseieva, H.D.];  
LECHEKHLEB, V.R. [Lechakhlid, V.R.]; SHCHERBAK, M.P.

Accurate determination of the absolute age of rocks by the lead  
method. Geol.sbur. 21 no.5:48-57 '61. (MIRA 14:10)

1. Institut geologicheskikh nauk AN USSR.  
(Geological time) (Mineralogy)



BICHUKIN, Filipp Demidovich; KAZACHENOK, Vladimir Isidorovich;  
RABANEYEV, F.Sh., inzh., retsennent; ALEKSEYEVA, Ye.N., red.

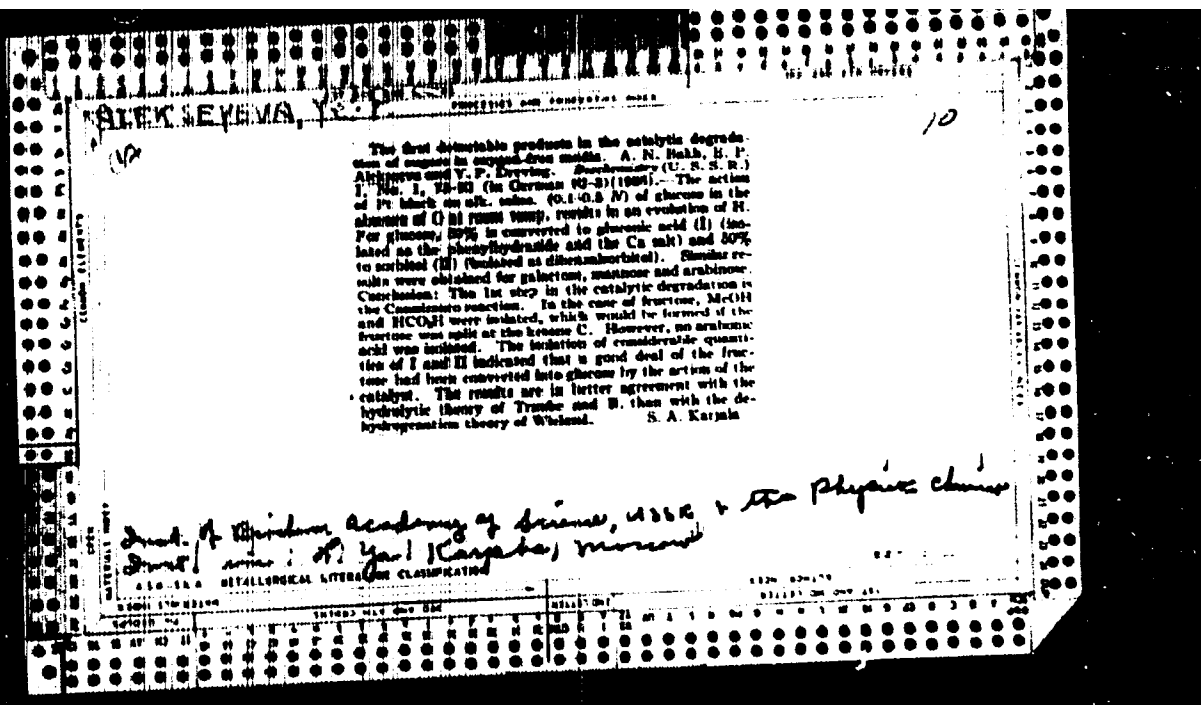
[Low-waste precision forging] Malootkhodnaia i tochnaia shtam-  
povka. Izhevsk, Udmurtskoe knizhnoe izd-vo, 1961. 92 p.  
(MIRA 17:4)

GORBOV, Mikhail Ivanovich; ALEKSANDROV, A.A., kand. ist. nauk,  
red.; ALEKSEYEVA, Ye.N., red.

[Gunsmiths of Izhevsk] Izhevskie oruzheiniki. Izhevsk,  
Udmurtskoe knizhnoe izd-vo, 1963. 102 p. (MIRA 17:4)

VERSHININ, V.V.; ALEKSEYEVA, Ye.P.

Improving methods of a commercial sampling of ores. Gor.shur. no.2:  
35-36 P'55. (MIRA 8:7)  
(Ores--Sampling and estimation)



ALENSEYEVA, Ye.P.; SHOSHINA, O.Yu.

Comparing interferometers of the institutes of the committee. Trudy  
inst. Kon. stand., ser 1 izm. prib no. 47:63-79 '61. (MIRA 15:12)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut metrologii im.  
D.I. Mandeleeva.

(Interferometers—Testing)

ALEKSHYEVA, Ye.P.; SHOSHINA, O.Yu.

Comparing interferometers of the institutes of the committee. Trudy  
inst. K<sup>o</sup> stand. i. prib. no. 63-79 '61. (MIRA 15:12)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut metrologii im.  
D.I. Mendeleeva.

(Interferometers--Testing)

22194

S/048/61/025/004/043/048  
B117/B209

24.3500

AUTHOR:

Alekseyeva, Ye. P.

TITLE:

Luminescence of LiF crystals activated with uranyl nitrate

PERIODICAL:

Investiya Akademii nauk SSSR. Seriya fizicheskaya,  
v. 25, no. 4, 1961, 545-547

TEXT: The present paper has been read at the 9th Conference on Luminescence (Crystal Phosphors). The author has studied the absorption and luminescence spectra as well as the thermoluminescence of LiF crystals activated with uranium salts. The crystals were excited by X-rays from an X-ray tube ( $V = 50 \text{ kV}$ ,  $I = 20 \text{ mA}$ ) with a tungsten cathode and beryllium windows. The absorption spectra were studied by means of an  $\text{CQ-4}$  (SF-4) spectrometer. The absorption bands of the activator have maxima at  $\lambda_1 = 2500 \text{ \AA}$  and  $\lambda_2 < 2100 \text{ \AA}$ . An increase in activator concentration causes the maximum at  $\lambda_1 = 2500 \text{ \AA}$  to shift to  $2600 \text{ \AA}$ , and a new band with its maximum at  $\lambda_3 = 3150 \text{ \AA}$  appears. The fluorescence spectra of activated LiF were taken

Card 1/4

Luminescence of LiF crystals ...

22194  
S/048/61/025/004/043/048  
B117/B209

with an ИСП-53 (ISP-53) spectrograph with an ФЭП-1 (FEP-1) recorder. The phosphor was excited by light from a ПРК-2 (PRK-2) lamp with a УФС-2 (UPS-2) filter. At room temperature, the fluorescence spectra showed a number of individual bands in the green region. When the activator concentration is varied, the shape of the curves remains in general the same. The individual spectra differ as to the relative intensity of the individual bands. The comparison of present results with those obtained by P. P. Feofilov (Ref. 2: Materialy V Soveshchaniya po lyuminestsentsii, Izd. AN EstSSR, Tartu, 1957) for the luminescence spectra of  $\text{CaF}_2\text{-U}^{6+}$  shows a fairly good agreement of the characteristic wavelengths of these spectra. For this reason, the luminescence of a phosphor activated with uranyl salts may also be considered a radiation of the uranium ion which has been transformed through the crystal lattice. The author found that the excitation of LiF phosphors with uranyl admixtures by X-rays entails the formation of the same color centers as the action of an ionizing radiation upon pure LiF crystals. When the concentration of the activator (of the order of 0.2%) is sufficiently high, a new absorption band with its maximum at  $\lambda \approx 5700 \text{ \AA}$  appears, giving the crystal a violet color. This

Card 2/4



22194

S/048/61/025/004/043/048  
B117/B209

Luminescence of LiF crystals ...

band is stable and is conserved for several months when the samples are stored at room temperature. It is undoubtedly related to the addition of the activator, as it does not appear in pure crystals, not even in the case of very long exposure. The thermoluminescence of pure and activated crystals was investigated, too. The luminescence of the activator was isolated by means of a special filter. The thermoluminescence curve characteristic of a non-activated crystal was found to exhibit a number of maxima, corresponding to the following temperatures: 110°, 163°, 215°, 300°, and 360°C (for  $\beta = 0.7 \text{ deg. sec}^{-1}$ ). Moreover, the activation of crystals was found to change the interrelations between these maxima, without, however, giving rise to any new maxima. This leads to the conclusion that the light sums in this case are accumulated on the trapping levels of the base substance, just like in the case of other alkali halide crystals. Finally, the author thanks L. M. Belyayev and Z. P. Perekalina for having supplied the samples. In the discussion to this paper, Z. P. Perekalina stated that luminescent LiF crystals activated with uranyl salts were grown in air according to the method of Kyropoulos at the Institut kristallografii (Institute of Crystallography). Crystals grown in vacuo did not exhibit any luminescence. This is proof

Card 3/4

22194

S/048/51/025/004/043/048  
B117/B209

Luminescence of LiF crystals ...

that  $U^{6+}$  has to be contained in activated LiF to produce luminescence centers. The luminescence and absorption centers of LiF-U crystals were studied for various uranium concentrations. At concentrations of 0.01 ÷ 0.02% by weight in the melt, intensity is redistributed in the bands of the luminescence spectra: Bluish-green luminescence changes over into yellowish-green luminescence. At the same time, the absorption spectrum is changed, too: the absorption bands shift into the long-wave region. This paper was read at the Conference on Alkali-halide Compounds at Tartu, July, 1959. There are 1 figure and 8 references: 4 Soviet-bloc and 4 non-Soviet-bloc. The four references to English-language publications read as follows: C. Delbeck, P. Pringsheim, J. Chem. Phys., 21, 5 (1953); F. Morehead, F. Daniels, J. Chem. Phys., 27, 6 (1957); J. Sharma, Phys. Rev., 78, 3, 535 (1952); D. F. Saunders, F. Morehead, F. Daniels, J. Amer. Chem. Soc., 75, 3096 (1953).

ASSOCIATION: Irkutskiy universitet (Irkutsk University)

Card 4/4

S/076/62/036/007/008/010  
B101/B138

AUTHORS: Yemel'yanenko, G. A., and Alekseyeva, Ye. P.

TITLE: Effect of temperature on the electrodeposition of some metals

PERIODICAL: Zhurnal fizicheskoy khimii, v. 36, no. 7, 1962, 1532 - 1536

NOTE: To find the optimum temperatures for the electrodeposition of Cu, Zn, and Ag from various electrolytes, the curves  $\log i$  versus  $1/T$  were plotted. The following electrolytes were used: 12.5 g/l  $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$  + 2.5 g/l  $\text{H}_2\text{SO}_4$  (I); 200 g/l  $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$  + 50 g/l  $\text{H}_2\text{SO}_4$  (II); 450 g/l  $\text{SnSO}_4 \cdot 7\text{H}_2\text{O}$  + 50 g/l  $\text{Al}_2(\text{SO}_4)_3 \cdot 18\text{H}_2\text{O}$  (III); 17 g/l  $\text{AgNO}_3$  (IV), and 40 g/l  $\text{AgCl}$  + 200 g/l  $\text{K}_4[\text{Fe}(\text{CN})_6] \cdot 3\text{H}_2\text{O}$  + 20 g/l  $\text{K}_2\text{CO}_3$  (V). A deviation from linearity was observed for the function  $\log i = f(1/T)$ . It occurred at certain polarizations and temperatures and was more distinct at high polarization. It lies at 40 mv, 40-50°C, for electrolyte I; at 50 mv, 50-60°C, or 100 mv, 40°C, for II; at 50 mv, 30-40°C, for III; at 10 mv, Card 1/2

Effect of temperature ...

S/076/62/036/007/008/010  
B101/B138

40°C, or 20 mv, 35°C, for IV; and at 30 mv, 35°C or 100 mv, 45°C, for V. The activation energy  $W$  for electrolytes I-IV is much lower above than below the bend in the curve, whereas V shows the opposite behavior to  $W$ . The deviation from linearity and the drop in  $W$  for I-IV are attributed to transition from chemical to concentration polarization. The deviation of V is attributed to the fact that  $Ag^+$  ions are discharged at low polarization, but complex silver ions at high polarization. Conclusion: This transition from chemical to concentration polarization must be allowed for when deciding the optimum temperature. Optimum temperatures for a rapid and irreversible process with prevailing chemical polarization are: for I (no data); II 60 mv, up to 40°C; III 50 mv, 18-25°C; IV (no data); V 250 mv, 60-80°C. There are 5 figures and 2 tables.

ASSOCIATION: Dnepropetrovskiy gosudarstvennyy universitet (Dnepropetrovsk State University)

SUBMITTED: May 29, 1961

Card 2/2

"APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6

APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6"

ACCESSION NR: AP4025093

S/0139/63/000/006/0110/0116

AUTHOR: Alekseyeva, Ye. P.

TITLE: Luminescence of activated LiF-crystals

SOURCE: IVUZ. Fizika, no. 6, 1963, 110-116

TOPIC TAGS: lithium fluoride, single-crystal luminescence, uranyl salt, absorption spectra, spectrophotometer, activator concentration, M-center absorption, thermo-luminescence

ABSTRACT: A detailed investigation has been made of lithium fluoride single-crystal luminescence activated with uranyl salt, copper, magnesium, and iron. Absorption spectra were obtained by means of the SF-4 spectrophotometer in wave lengths from 2050 to 8000 Å. The specimen was a plane parallel plate 1 mm thick. The absorption spectra of the activated crystal show three maximum absorption bands. Increasing the activator concentration shifts the maxima towards long wave lengths. Auxiliary absorption of LiF is discussed, and the M-center absorption is noted under continuous and intermittent x-ray irradiation at 120°C. The effect of ionizing radiation on activator centers is analyzed in the LiF-U crystal phosphor. It is

Cord 1/2

ACCESSION NR: APL025093

found that green fluorescence brightness in LiF-U decreases under x-ray irradiation. The thermoluminescence of LiF-Hg shows two maxima at 80 and 110C temperatures, whereas in LiF-Cu the thermoluminescence maximum occurs above 215C. In LiF-Fe the maximum moves up to 300C, and LiF-U maxima occur both at high and low temperatures. "The author is grateful to Professor I. A. Parfianovich under whose guidance this work was done." Orig. art. has: 6 figures.

ASSOCIATION: Irkutskiy gosuniversitet imeni A. A. Zhdanova (Irkutsk State University)

SUBMITTED: 30Jun62

DATE ACQ: 14Feb64

ENCL: 00

SUB CODE: RH

NO REF SOV: 008

OTHER: 013

Card 2/2

ACC NR: AP6030719

SOURCE CODE: UR/0368/66/005/002/0216/0220

AUTHOR: Alakseyeva, Ye. P.

ORG: none

TITLE: Dependence of thermoluminescence curves of LiF phosphors on the nature of impurity, preliminary thermal treatment, and x-ray irradiation dose 17

SOURCE: Zhurnal prikladnoy spektroskopii, v. 5, no. 2, 1966, 216-220

TOPIC TAGS: lithium fluoride, crystal phosphor, thermoluminescence, radiation dosimeter, crystal impurity, x ray irradiation

ABSTRACT: This study was made because LiF phosphors might be used as radiation dosimeters. The thermoluminescence of doped and undoped LiF crystals irradiated by x-rays was studied at room temperature. The experimental results show that the magnitude of the sum of light, accumulated at the individual capture levels, is substantially affected by the irradiation time of the crystal, by the annealing of the crystal prior to irradiation, and by the presence of impurities in the crystal. This paper was read at the Twelfth All-Union Conference on Luminescence of Crystal Phosphors, held in Lvov in January 1964. The author thanks Prof. I. A. Parfianovich for setting up the problem and directing the work. Orig. art. has: 3 figures.

SUB CODE: 20/ SUBM DATE: 25Jan65/ ORIG REF: 009/ OTH REF: 005

Card 1/1

UDC: 535.377



ACC NR: AP7004986

SOURCE CODE: UR/0048/66/030/009/1506/1508

AUTHOR: Alekseyeva, Ye. P.

ORG: none

TITLE: M band stimulation of luminescence in x-irradiated LiF crystals /Report, Fourteenth All-Union Conference on Luminescence (Crystal Phosphors) held at Riga, 16-23 Sept. 1965/

SOURCE: AN SSSR, Izvestiya. Seriya fizicheskaya, v.30, no.9, 1966, 1506-1508

TOPIC TAGS: luminescent crystal, lithium fluoride, luminescence spectrum, excitation spectrum, luminescence center

ABSTRACT: The author has recorded the excitation spectrum of the red luminescence of LiF crystals colored by x irradiation. The resulting spectrum was similar to that obtained by C.Klick (Phys. Rev., 112, 1620 (1950)): the red luminescence was stimulated not only by irradiation in the M band (at 4450 Å), but also by irradiation at certain other wavelengths, namely, at 2200, 2500, 3200, 3800 and 4050 Å. Three of these stimulation peaks are close to the F, R<sub>1</sub>, and R<sub>2</sub> induced absorption peaks at 2450, 3150 and 3800 Å, respectively, and it has been suggested that the red luminescence is stimulated at these wavelengths by transfer of energy from F, R<sub>1</sub>, and R<sub>2</sub> centers to M centers. The author proposes an alternative explanation, based on the calculations of A.Mayer and R.F.Wood (Phys.Rev., 133, A1436 (1964)) of the energy

Card 1/2

ACC NR: AP7004986

levels of an M center in a LiF crystal. Differences between the energies of some of these levels correspond to wavelengths of 2220, 4050, and 2520 Å, and it is suggested that the M centers are excited by direct absorption of photons of the corresponding energies. The temperature dependences of the stimulation of the red luminescence by 4450, 3800, 2220 and 2520 Å radiations were found to be very similar, indicating that in all these cases the stimulation is effected by transitions in the same centers. By identifying the 4450 Å "ordinary" M absorption peak with a transition involving one state whose energy had not been calculated, the author was able to account also for the 3800 Å excitation peak. On the basis of this identification, however, one would expect the 3200 Å excitation peak to occur at 3380 Å; this discrepancy cannot be explained at present. The author thanks I.A.Parfianovich for his guidance of and interest in the work. Orig. art. has: 2 figures.

SUB CODE: 20 : SUBM DATE: none

ORIG. REF: 005

OTH REF: 004

Card 2/2

ALEXANDER, Ye. S.

ALEXANDER, Ye. S.: "Local varieties of *L. grandis* for soil's tilth and measures to improve them." In: *Plodovodstvo Ukrainy*, No. 1, 1956, Order of Lenin: USSR Agricultural Academy. Kiev, 1956. (Dissertation for the Degree of Candidate in Agricultural Sciences)

So: Russkaya Letopis' No 3, 1951. Moscow

ALEXANDER, Ye.S.

Selecting parental types for developing hybrids of Hungarian  
millet. Agrobiologiya no. 3:383-385 My-Je '60. (MIRA 13:12)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut kornov  
im. V.R. Vil'yamsa.

(Millet breeding)

ALEKSEYEVA, Ye. S., Cand Agr Sci -- "Creation of hybrid starting material for  
the cultivation of new <sup>varieties</sup> ~~species~~ of Italian millet in northern regions." Nos. 1960  
(All-Union Sci Res Inst of Fodders in V.R. Vilyams). (RL 1-01, 200)

-261-

S/020/61/139/005/009/021  
B104/B201

AUTHORS: Berestnev, V. A., Razikov, K. Kh., Alekseyeva, Ye. S., and  
Kargin, V. A., Academician

TITLE: Structure of oriented polymers

PERIODICAL: Akademiya nauk SSSR. Doklady. v. 139, no. 5, 1961, 1093-  
1094

TEXT: The structure of polymers includes a so-called supermolecular formation which reaches beyond the molecular dimensions. The authors used an electron microscope to examine this supermolecular formation on thread sections cut by a microtome. Longitudinal and cross sections were prepared, and special care was taken to save the structure from damages in the cutting process. In this process, all threads were found to split into small fibers with diameters ranging between 1 and 10 $\mu$ . The fibers had partly a nonoriented, coarse spherulitic structure, and fine globular formations; others had nonoriented macroformations. The authors base on these results to assume that the structure of chemical threads consists of anisodiametric elements. the fibers. The fibers do not contain any

Card 1/2

Structure of oriented polymers

S/020/61/139/005/009/021  
B104/B201

oriented microstructural elements. The anisotropy of thread properties is evidently caused by the fiber shape. From a micropicture of a fiber (not reproducible) in which there is a hole the authors infer from the fact that this hole is neither circular nor elliptic that there are stress inhomogeneities in the fibers. The results yielded by examinations of fibers by a polarization microscope (400 - 600 - fold magnification) cause the authors to conclude that some fibers lack an orientation of the molecular structure; an orientation of the surface layer of the fiber was established in a number of cases. Thus, the structure of chemical threads appears to consist of fibers having a nonoriented regular macroformation. The anisotropy of a number of properties in these materials is caused by the fibrous structure. There are 4 figures and 3 Soviet references.

ASSOCIATION: Fiziko-khimicheskiy institut im. L. Ya. Karpova  
(Physicochemical Institute imeni L. Ya. Karlov)

SUBMITTED: April 18, 1961

Card 2/2

BERESTNEV, V.A.; ALEKSEYEVA, Ye.S.; NOSOV, M.P.

Measuring birefringence by the thickness of the fiber. Khim.volok.  
no.2:40-43 '63. (MIRA 16:5)

1. Nauchno-issledovatel'skiy institut shinnoy promyshlennosti (for  
Berestnev, Alekseyeva). 2. Kiyevskiy filial Vsesoyuznogo nauchno-  
issledovatel'skogo instituta iskusstvennogo volokna (for Nosov).  
(Textile fibers--Optical properties)



DUBOVA, L.S.; BERESTNEV, V.A.; NAGLASEVA, I.P.; Prinimali uchastiye:  
ALEKSEYeva, Ye.S.; PRYAMIKOVA, T.S.

Studying the double refraction of some polyamide fibers.  
Khim.yolok. no.5:52-55 '64. (MIRA 17:10)

1. Nauchno-issledovatel'skiy institut shveytnoy promyshlennosti.

BERESTNEV, V.A.; LUBOVA, L.S.; ALEKSEYEVA, Ye.S.

Structure of polyamide fibers studied by the etching test. Khim.  
volok. no.5:29-31 '63. (MIRA 16:10)

1. Nauchno-issledovatel'skiy institut shinnoy promyshlennosti.

1. ALEKSEYEVA, YE.V.
2. USSR (600)
4. Dairy Cattle - Feeding and Feeding Stuffs
7. Useful pamphlet ("Practice of leaders in widespread feeding for milk production."  
N. P. Semenov. Reviewed by YE. V. Alekseyeva), Sots.zhiv. 15 no. 4, 1953.
9. Monthly List of Russian Accessions, Library of Congress, APRIL 1953, Uncl.

FALINA, N.N.; MASLOVA, R.A.; YAKIMOV, P.A.; ANDREYEV, S.M.; ALEKSEYEV, Ye.V.

Some results of studying Basidiomycetes as a source for obtaining  
feed proteins and diet-deficient amino acids. Rest. res. 1 no.1:  
122-127 '65. (MIRA 18:6)

1. Botanicheskiy institut im. V.L. Komarova AN SSSR, Leningrad.

SOV/137-58-7-14118

Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 7, p 20 (USSR)

AUTHORS: Levenshteyn, S. A. , Alekseyeva, Ye. V.

TITLE: Possibilities for the Utilization of "Biscuit" Clays in the Binder Component of Fireclay (O vozmozhnosti primeneniya sukharnykh glin v svyazuyushchey chasti shamotnykh mass)

PERIODICAL: Tr. Leningr. tekhnol. in-ta im. Lensovet, 1957, Nr 43, pp 99-105

ABSTRACT: Studies performed establish that it is possible to introduce up to 15% nonplastic, "biscuit"-type Borovichi clays into the mix of fireclay refractories without impairing the quality of the latter, thus making possible economies in plastic binder clays.

1. Refractory materials--Preparation 2. Refractory S. G.  
materials--Binders 3. Clays--Applications

Card 1/1

ALEXSEYEVA, E. V., BARANTSEV, R. G. (Leningrad)

"A Circular Plate in Almost Free-Molecule Flows."

report presented at the First All-Union Congress on Theoretical and Applied Mechanics, Moscow, 27 Jan - 3 Feb 1960.

BARANTSHV, R.G.; ALEKSEYEVA, Ye.V.

Highly rarefied monatomic gas flow past bodies. Aerodin. razresh.  
gaz. no.1:183-194 '63.

Highly rarefied gas flow around a mirror-image reflecting sphere.  
Ibid.:195-211 (MIRA 17:3)

"APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6

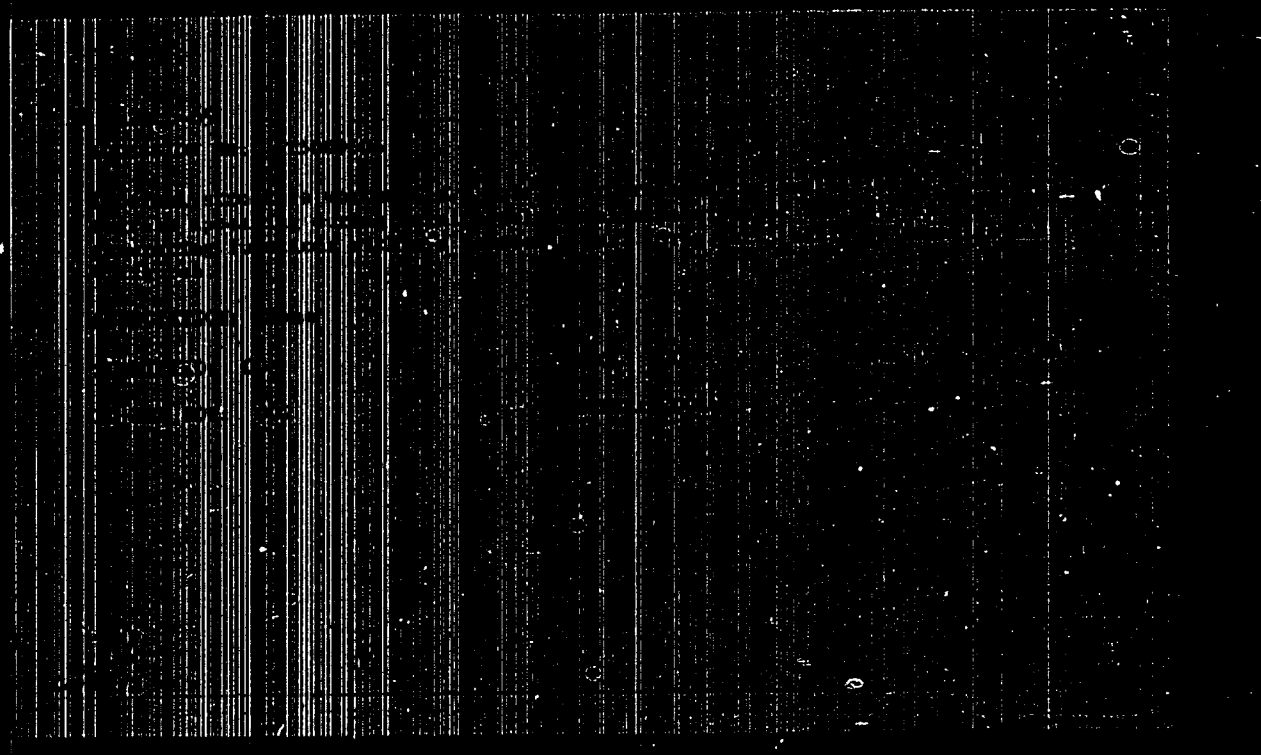
APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6"



"APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6

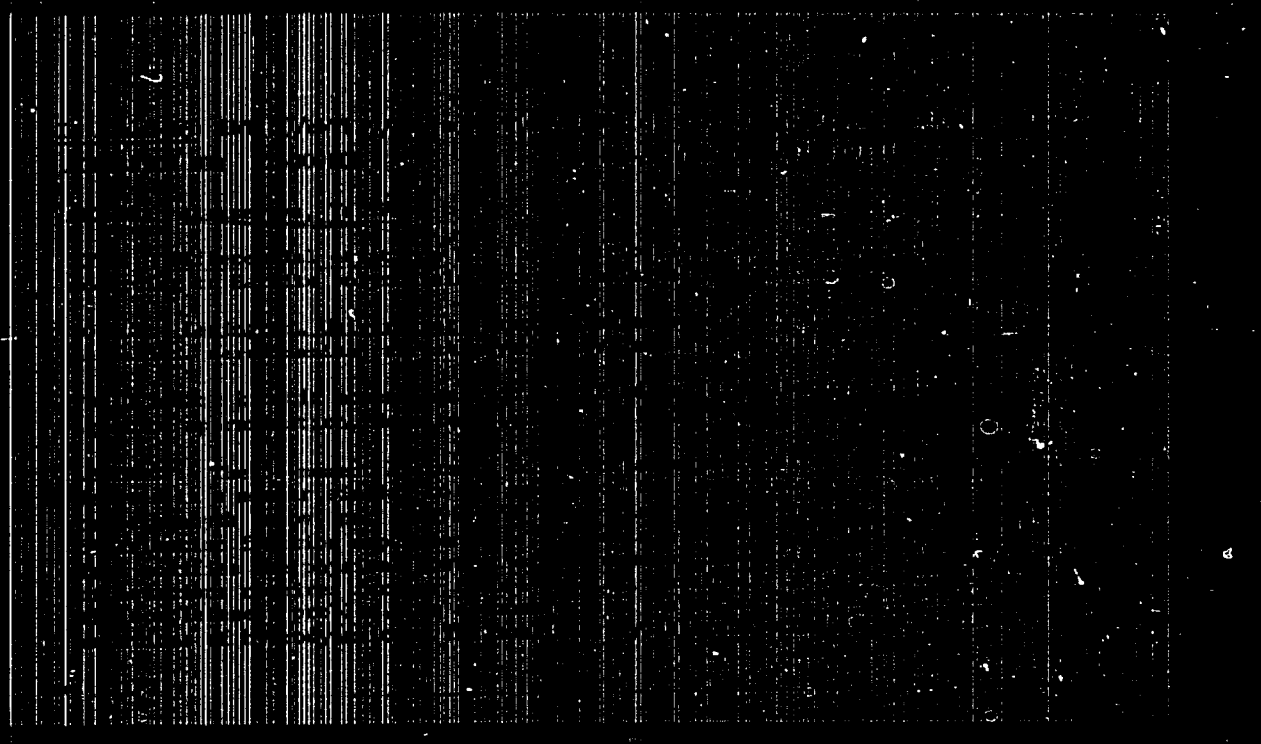


APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6"

"APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6

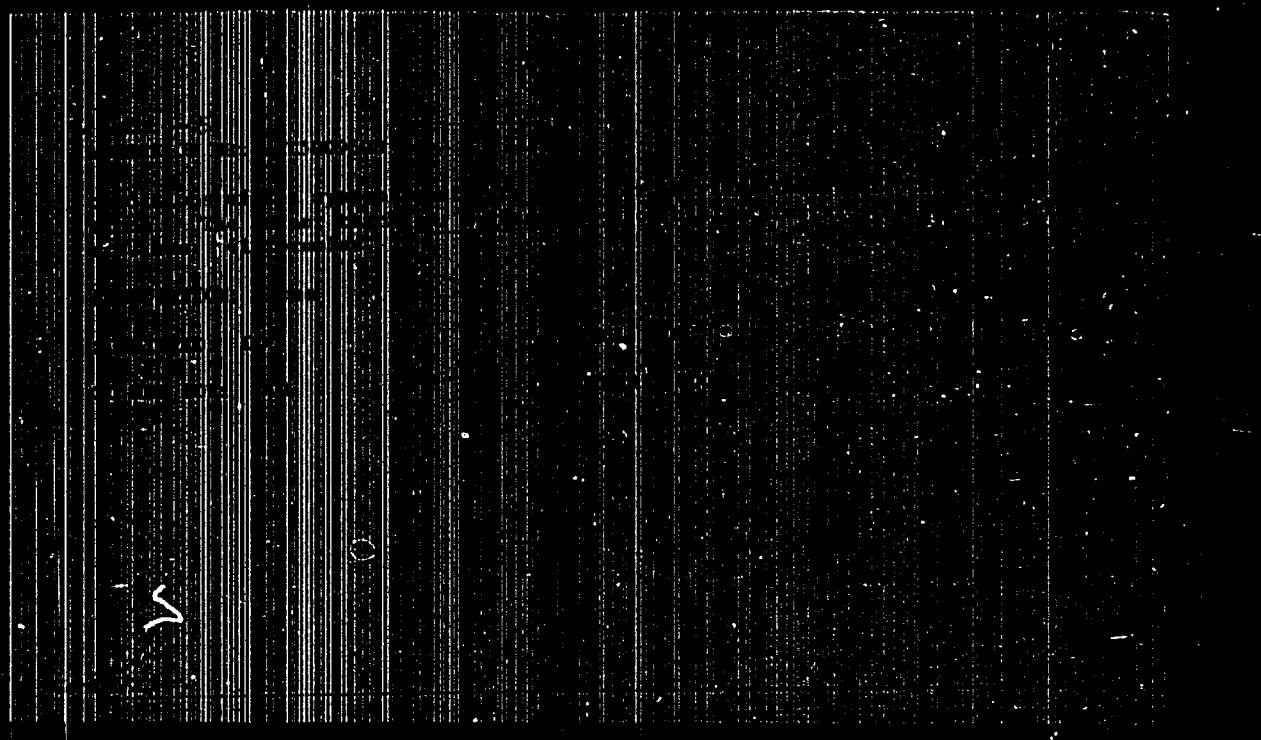


APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6"

"APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6



APPROVED FOR RELEASE: 09/24/2001

CIA-RDP86-00513R000101010011-6"

RYABCHUK, A.F.; YUR'YEV, A.M.; PLETNIKOV, K.V., redaktor; ALEKSEYEVA, Ye.Ye.,  
redaktor; MATISSEN, Z.M., tekhnicheskiiy redaktor

[Some 16mm sound motion-picture apparatus] Uskoplennochnye zvukovye  
kinoustanovki. Pod obshchei red. K.V.Pletnikova. Isd. 2-oe, perer.  
i dop. Moskva, Gos. izd-vo "Iskusstvo," 1956. 193 p. 96 illus. 1.  
(Motion-picture projectors) (MLRA 10:1)