

Kinetics and Mechanism of the Decomposition S/153/60/003/004/04/04C/XX  
Reaction of the Hexacyanoferrate Ion in the B020/B054  
Presence of Silver Compounds

relation between the light absorption of the solutions and the logarithm of the silver salt concentration for a certain time in a concentration range of silver ions between  $3 \cdot 10^{-7}$  and  $2 \cdot 10^{-6}$  moles/l, and in the presence of thiourea. The catalytic reaction is much slowed down by addition of a NaCN solution to a weakly acid hexacyanoferrate solution containing  $\text{AgNO}_3$  and thiourea (Fig. 8). The authors revealed the reaction mechanism according to which the slowest reaction is the decomposition of the intermediate complex formed from positively charged catalyst ions ( $\text{Ag}^+$  or  $\text{Ag}(\text{CSN}_2\text{H}_4)_2^+$ ) and the cyanide complex of bivalent iron. They also derived the kinetic equation for this reaction, and calculated the rate constants (Table 1). At a pH of 3.7, and at  $40^\circ\text{C}$ , the rate constant of the reaction is equal to  $0.016 \pm 0.003 \text{ min}^{-1}$  (Table 2). The authors determined the equilibrium constant of the conversion of the cyanide complex of silver into a thiourea complex in weakly acid solution. Silver can be quantitatively determined in concentrations of the order of

Card 3/4

Kinetics and Mechanism of the Decomposition S/153/60/003/004/014/040/XX  
Reaction of the Hexacyanoferrate Ion in the Presence of Silver Compounds B020/B054

magnitude between  $10^{-7}$  and  $10^{-6}$  moles/l on the basis of light absorption measurements in solutions containing potassium hexacyanoferrate, an acetate buffer, a silver salt, and thiourea. There are 8 figures, 2 tables, and 14 references: 3 Soviet, 1 US, 4 British, 2 Yugo-Slav, 2 French, 1 Austrian, and 1 Dutch.

ASSOCIATION: Ivanovskiy khimiko-tekhnologicheskii institut, kafedra analiticheskoy khimii (Ivanovo Institute of Chemical Technology, Department of Analytical Chemistry)

SUBMITTED: December 9, 1958

Card 4/4

**YATSIMIRSKIY, K.B.; ORLOVA, M.N.**

Kinetics and mechanism of the conversion of the hexacyanoferrate(II) ion in the presence of a gold-thiourea complex. Zhur. neorg. khim. 5 no.10:2184-2189 0 '60. (MIRA 13:10)

1. Ivanovskiy khimiko-tekhnologicheskoy institut, Kafedra analiticheskoy khimii.

(Gold compounds)

(Iron compounds)

GRIGVA, M. N.

Cand Chem. Sci - (russ) "Kinetics and mechanism of transformation of hexacyanoferrate ion in the presence of gold and silver compounds." Ivanovo, 1961. 15 pp; (Ministry of Higher and Secondary Specialist Education RSFSR, Ivanovo Chemical Technology Inst.; 150 copies; price not given; (KI, 5-61 sup, 179)

PIRYKHIN, B.V.; MOPACHENSKIY, V.G.; OLSOVA, M.N.

Evaporation rate of droplets of aqueous solutions of surface-active agents. Probl. fiz. atm. no.2:142-150 '63.

(MFA 17:5)

LIKHTENSHEYN, Ye.A., starshiy nauchnyy sotrudnik; ORLOVA, M.P., nauchnyy sotrudnik

X-ray diagnosis of metastases and growth of malignant tumors in the mandible. Stomatologiya 39 no.1:31-36 Ja-F '60. (MIRA 14:11)

1. Iz rentgenodiagnosticheskogo otdeleniya (zav. - prof. Ye.E. Abarbanel') Gosudarstvennogo onkologicheskogo instituta imeni P.A.Gertsena (dir. - prof. A.N.Novikov, nauchnyy rukovoditel' - prof. A.I.Savitskiy).

(JAWS--CANCER)

(JAWS--RADIOGRAPHY)

A. C. S. ORLOVA, M. P.

*Glass*

Improving the light transmission of Fournet glass. I. B. SHAPRO AND M. P. ORLOVA. *Soviet Phys. J. Techn. Phys.*, 1966, No. 17, pp. 4-8. -- To increase the transparency of glass and to diminish its coloration, it is necessary to keep the Fe content in the glass as low as possible and to transform the little that must be included into  $FeO$ . The latter is favored by (a) maintaining oxidizing conditions during melting, (b) maintaining as low a temperature as possible during melting, and (c) shortening the time during which the melt is exposed to high temperatures. These conditions are readily satisfied when the melt is carried out in pot furnaces operating periodically. They are harder to realize in tank furnaces, because of the longer period the melt spends in the furnace and the higher operating temperatures. The continuous operation of the furnace also limits the variety of compositions that can be used in the batch. The investigation was concerned with selecting a batch composition for a continuous furnace that would yield a glass at least as transparent as that obtained in pot furnaces. It was required to obtain in a Fournet furnace a transparency of 80% at a sheet thickness of 20 mm. There were two objectives: (1) to obtain a glass with a minimum of  $Fe$  oxides, and (2) to create such conditions that the equilibrium  $Fe_2O_3 \rightleftharpoons 2FeO + O_2$ , within the melting mass would be shifted to the left. To reduce the Fe content, the sand was cleaned on a Willey table. This alone reduced the Fe oxide content by 50% or more. Furthermore, the iron drums used for drying the washed sand were replaced by a hot-flow furnace, which eliminated the contact of the sand with iron. The dolomite and limestone used in the batch were carefully graded and, after being ground twice, were passed through a magnetic separator to remove any iron picked up in the grinding. The combined effect of

these operations reduced the  $Fe_2O_3$  content in the glass from 0.12 to 0.05%. Several experimental melts were made in pots to establish the most propitious oxidizing conditions. The following agents were used to increase the O pressure within the melt and thus prevent the decoloration of  $Fe_2O_3$ : (1) sulfite, (2) sulfite +  $KNO_3$ , (3) sulfite +  $KNO_3$  +  $CaF_2$ , (4) sulfite +  $As_2O_3$  +  $KNO_3$ , (5) sulfite +  $K_2CO_3$ , and (6) sulfite +  $As_2O_3$  +  $K_2CO_3$ . The experimental melts show that glasses can contain a high percentage of  $K_2O$  introduced as  $KNO_3$ , clarify easier and therefore melt more easily in the furnace. An addition of  $KNO_3$  in quantities up to 5% and an addition of  $KNO_3$  with 0.25 to 0.5% of  $As_2O_3$  hasten clarification. Of all the batches tested, the highest light transmission and the lowest coloring were found with the batch containing 5% of  $K_2O$  added as  $KNO_3$  and 0.25 to 0.5% of  $As_2O_3$ . The new composition of the glass was chosen as  $SiO_2$  71.5,  $B_2O_3$  0.8,  $CaO$  0.0,  $MgO$  3.0,  $Na_2O$  15.0,  $K_2O$  1.0, and  $As_2O_3$  0.25. The glass was made from a batch of sand 88, limestone 12.0, dolomite 0.4, soda 74.5, sulfite 0.77, calcifer 0.8, and  $As_2O_3$  1.0 kg. Within the furnace, temperatures were maintained as follows: at the first pair of burners  $1450^\circ \pm 10^\circ$ ,  $10^\circ$ , beyond the second pair of burners  $1400^\circ \pm 10^\circ$ , and in the cooling zone  $1340^\circ \pm 10^\circ$ . The batch fed into the degasser melted within 20 min. Beyond the second pair of burners the glass was free of bubbles and strain. After 4 days of leaching this batch, the glass pulled from the working end and made into 20-mm. sheets had a transmission of 87% and a very satisfactory weak yellowish color. M.M.

ORLOVA, M. P. Engineer

" Method for Determining the Homogeneity of Certain Physical Properties of Glasses."  
Thesis for degree of Cand. Technical Sci Sub 30 May 50, All-Union Sci Res Inst of  
Glass, Ministry of the Construction Materials Industry USSR

Summary 71, 4 Sep 52, Dissertations Presented for Degrees in Science and Engineering  
in Moscow in 1950. From Vechernyaya Moskva. Jan-Dec 1950



ORLOVA, M.P., kandidat tekhnicheskikh nauk; SESOROVA, V.N., kandidat  
tekhnicheskikh nauk; TYKACHINSKIY, I.D., kandidat tekhnicheskikh  
nauk.

Investigating the performance of the VVS machines at high speeds in  
the Bytoshevskiy and Chagodoshchenskiy glass works. Trudy VNI  
Stekla no.36:82-94 '56. (MLRA 9:11)  
(Glass manufacture) (Furnaces)

*ERLOVA M P*

ORLOVA, M.P.; POLLYAK, V.V.; TYKACHINSKIY, I.D.

Speeding up the melting process is a powerful means for increasing  
the productivity of glass furnaces. Stek. i ker. 14 no.9:1-4 S '57.  
(MIRA 10:10)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut stekla.  
(Glass manufacture)

ORLOVA, M.P.

**AUTHORS:** Engver, Ye.A., Chief Engineer of the  
"Proletariy" Works, Katayeva, G.V., Orlova, M.P.,  
Collaborators of the Institute for Glass

72-2-3/20

**TITLE:** The Practical Application of Ammonium-Sulfate for the Acceleration  
of the Process of Glass Smelting (Praktika primeneniya sul'fata  
ammoniya kak uskoritelya varki stekla).

**PERIODICAL:** Steklo i Keramika, 1958, Nr 2, pp. 6-7 (USSR)

**ABSTRACT:** The staff of the "Proletariy" works, together with the working group  
of the Institute for Glass, carried out a practical test with the  
continuous glass-smelting furnace Nr 2 having a total surface of  
141.6 m. A.L. Nikanorova participated in this work. The authors  
further describe the temperature conditions of the furnace, the chem-  
ical composition of the glass, and the composition of layers. The  
correlation of the Na<sub>2</sub>O-quantities, which were introduced by soda  
and sulfate into the layer, was 90:10, the moisture content of the  
layer 0.5%. 20-25% of scrap was added. Before the use of ammonium-  
sulfate the layer contained 0.15% F', which exercises no noticeable  
influence on the acceleration of glass smelting. This quantity was,  
however, left in the layer also further. After the introduction of

Card 1/2

The Practical Application of Ammonium-Sulfate for the  
Acceleration of the Process of Glass Smelting

72-2-3/20

3% ammonium-sulfate, smelting and refining of the glass mass improved considerably. The entire heat conditions of the furnace as well as the technological process remained unchanged, only the means of reduction (coal) was increased from 8 to 11% of the weight of the ammonium-sulfate introduced. The characteristic values of work before and after introduction of the ammonium-sulfate are shown in a table. There are 1 table and 2 Slavic references.

ASSOCIATION: Zavod "Proletariy" (Proletariy Works), Institut stekla (Institute for Glass)

AVAILABLE: Library of Congress

Card 2/2

SOV 2-58-8-2000

AUTHORS: Kerzhinskaya, I. V., Shlova, E. I., Selezneva, N. N.,  
Smirnov, E. I., Shubin, V. P.

TITLE: **Industrial Experiment in Replacing Sodiumsulphate by Astrachanite**  
in the Melting of Glass (Promyshlennyy opyt zameny sulfata  
natriya astrakhanitim pri varke stekla)

PERIODICAL: Steklo i keramika 1958, Nr 8, pp. 3 - 5 (UDC 661.7)

ABSTRACT: The possibilities of using astrachanite in the melting of  
glass were investigated at the Institute of Glass (Institut  
stekla) by N. Ia. Raf in 1940 - 1953, as well as at the  
Belorussian Polytechnical Institute (Belorusskiy politekhn-  
cheskiy institut) by A. S. G-zburg in 1941. Besides, the  
All-Union Institute of Metallurgy (Vsesoyuznyy institut galyur-  
gii) carried out investigations on the working up of astrachanite  
from 1947 to 1954. The great attention which was  
attracted by this mineral may be explained by the fact that  
huge deposits may be found in the area of the Aral and Caspian  
Seas (Aral'skoye i Kaspiyskoye morya), the lower Volga  
(Nizhnnyaya Volga) and at a number of other places. The fol-

Card 1/3

SOV/72-58-8-2, 17

**Industrial Experiment in Replacing Sodiumsulphate by Astrachanite in the Melting of Glass**

Following formula holds for the composition of astrachanite:  
 $A = 278x / (100 + B)$ , where x denotes the percentage of  $MgCO_3$   
and B the percentage of  $H_2O$ . Earlier papers showed that  
astrachanite may be used only after its homogeneity had been  
improved (Ref 1). At the end of 1954 a working team of the  
Institute of Glass together with the collective of the kras-  
nousol'sk glass factory carried out a continuous experiment  
of glass melting in a tank furnace with astrachanite. More  
than 400 t of this mineral were used. Its chemical composi-  
tion and the sample taking are given and described. Its work-  
ing up was carried out according to scheme (Fig), and this  
process is then described in detail. By the introduction of  
astrachanite into the charge the properties of glass melting  
are not changed. The comparative data concerning work may  
be seen from Table 2. I. G. Druzhinin (Ref 2) showed in his  
paper that astrachanite melts at a temperature of  $670^\circ$ .

**Conclusions:**

- 1) Astrachanite may be used to replace sodiumsulfate.
- 2) This increases a little the costs of the charge.
- 3) To use this material successfully a respective preparation must be organized at its place of finding.

Card 2/3

New Tasks and a New Orientation of Our Periodical

SOV/72-58-3-1/17

glass and ceramics. Finally it is stated that the reorganization and improvement of the periodical cannot be solved by the editors alone. It needs the active participation of collaborators in the glass and ceramic industry.

1. Glass industry--USSR
2. Ceramic materials--USSR
3. Periodicals

Card 3/3

~~LOPUK, I.G.; ORLOVA, M.P.~~

Use of petrolatum in the drying and impregnation of materials for  
the manufacture of containers. Trudy Military no.2:71-80 '58.

(MIRA 13:12)

(Petrolatum)

(Wood--Preservation)

(Lumber--Drying)



ORLOVA, M.P.

Standardization of containers made of plywood. Trudy NIL  
Tary no.4:44-49 '60. (MIRA 14:12)  
(Containers--Standards)  
(Plywood)

ORLOVA, M.P.

Plywood drums. Standartizatsiya no.7:38-39 J1 '60.

(MIRA 13:7)

(Drums (Containers))

L 39557-66 EWT(1)/EWT(a)/EWP(t)/EPI IJP(c) JD/C  
ACC NR: AP6008780 SOURCE CODE: UR/0115/66/000/001/0057/000

AUTHOR: Orlova, M. P.; Kats, G. A.; Astrov, D. N.; Belyanskiy, L. B.;  
Shibayeva, O. A.; Shubin, V. E.

ORG: none

TITLE: Alloyed germanium for low-temperature thermometry

SOURCE: Izmeritel'naya tekhnika no. 1, 1976, 57-61

TOPIC TAGS: thermometry, germanium alloy, thermometer

ABSTRACT: The results are reported of an experimental investigation of the galvanomagnetic properties of Ge doped with various amounts of Sb, As, In, Ga. The Ge properties were studied in a range of temperatures from room to liquid helium in order to find out the best impurity and its concentration suitable for low-temperature thermometers. Most measurements were made with Sb-doped Ge.

Card 1/2

UDC: 546.289.001.5:036.071

L 39557-66  
ACC NR: AP6008780

whose Nd was  $4.6 \times 10^{16} < Nd < 1 \times 10^{17}$  per  $cm^3$ ; the resistivity was found to be  $0.00042-0.00046$  ohm·m at 20-4.2K; acceptor-impurity concentration  $N_a < 0.1 Nd$ . A few thermometers were made from Sb-doped Ge (Nd = 5%,  $N_a = 0.1$  per  $cm^3$ , K = 6%) for the 40-4.2K range; their resistivity was  $0.025-0.027$  ohm·m at boiling-helium temperature. The relation  $lg \rho (1/T)$  was satisfactory for the thermometers only under 7K. A relatively high value of magnetoresistance in doped Ge is noted. Orig. art. has: 4 figures, 4 formulas, and 4 tables.

SUB CODE: 20, 09 / SUBM DATE: none / ORIG REF: 003 / CTR REF:

Card 2/2 HS

ORLOVA, M.P.; OKZHDESTVINSKIY, Yu.P.; BARANOVA, Ye.N.

Mineralogy of the rare-metal carbonatites of the Salla-Latvinskii  
Massif (northern Karelia). Trudy VSEGEI 96:3-20 '63.  
(MIA 17:9)

ORLOVA, M.P.

Some problems of the petrochemistry and petrology of the  
Caledonian complex of alkali-ultrabasic rocks of the Kola  
Peninsula. Trudy VSEGEI 96:65-103 '63. (MIRA 1':9)

ORLOVA, M.P.

Alkali basaltoid rocks in the Daubaba Valley (Talas Ala-Tau).  
Inform.sbor. VSEGEI no.16:87-95 '59. (MIRA 15:3)  
(Daubaba Valley--Basalt)

ORIOVA, M.P.

Alkaline gabbroid intrusions in the northwestern spurs of the  
Talas Ala-Tau. Uch. zap. IGU no.291:91-121 '60.

(MIRA 13:7)

(Talas Ala-Tau--Gabbro)



ORLOVA, M.P.; KUKHARENKO, A.A.

Melilite from alkali-ultrabasic massifs of the Kola Peninsula. Uch.zap.  
LGU no.312:173-189 '62. (MIRA 15:6)  
(Kola Peninsula--Melilites)

ORLOVA, M. I.

Dissertation: "Investigation of the Possibility of Reproduction of Temperature Phase Conversions in Solid Oxygen." Cand Tech Sci, All-Union Sci Res Inst of Metrology, Leningrad, 1954. Referativnyy Zhurnal--Khimiya, Moscow, No 7, Apr 54.

SO: SUM 284, 26 Nov 1954

ORLOVA, M. P.

SECRET

336.511 : 536.48

4662. Apparatus for reproducing the boiling point of hydrogen. A. S. BOROVIK-ROMANOV, M. P. ORLOVA AND P. G. STRELKOV. Zh. tekhn. Fiz., 24, No. 7, 1219-21 (1954) In Russian.

The method described was used in the measurement of the boiling point of H<sub>2</sub> (Abstr. 7233 (1952)). The reproducibility of the datum-point temperature (equilibrium between the liquid and vapour phase) can only be attained if the effect of the ortho-H/para-H conversion is either eliminated or allowed for. The apparatus includes 2 condensation thermometers in one of which the composition of H<sub>2</sub> is maintained by means of a catalyst in a state of equilibrium corresponding to room temperature, whereas in the other the equilibrium corresponding to ~20°K is established. The pressure differences between these thermometers indicate the degree of ortho-para conversion. Comparison with the results of the National Bureau of Standards and the Laboratory of Leyden shows that the systematic temperature difference is of the order of 0.001-0.002 deg. Systematic measurements of the boiling point with the aid of a reliable Pt resistance thermometer showed that, with a probability of 90%, the measurement of the boiling point of H<sub>2</sub> carried out by means of the apparatus described, is accurate within ±0.003%.

V. LACHMAN

ORLOVA, M. P.

Magnetic and thermal properties of the three modifica-  
 tions of solid oxygen. A. S. Borovik-Romanov, M. P.  
 Orlova, and P. G. Sirel'kov (State Inst. of Measures and  
 Measuring Apparatus, Moscow). *Doklady Akad. Nauk*  
 S.S.S.R. 99, 698-702 (1954); cf. C.A. 46, 10725b. — Solid O  
 exists in 3 modifications:  $\alpha$  (below 23.58°K.,  $\beta$  (23.58-  
 43.80°K.), and  $\gamma$  (from 43.80° to the triple point, 54.37°K.).  
 The heat capacity was measured in the region of the transi-  
 tion points. The data show that the  $\beta \rightarrow \gamma$  transition is of  
 the first order and the  $\alpha \rightarrow \beta$  transition is of the second order.  
 The magnetic permeability  $\chi$  of solid O was measured be-  
 tween 26.5 and 77.5°K. For the  $\gamma$ -modification (as for  
 liquid O)  $\chi$  obeys the Curie-Weiss law. Upon transition to  
 the  $\beta$ -modification,  $\chi$  decreases sharply and continues to de-  
 crease with decreasing temp. This is characteristic of  
 antiferromagnetic substances. The large heat effect of the  
 $\beta \rightarrow \gamma$  transition is attributed to the change to the antiferro-  
 magnetic state. J. Roytar Lashin

62

2

ORLOVA, M.P.

USSR/Physics - Measuring Instruments

Card 1/1 Pub. 147 - 22/27

Authors : Strelkov, P.G.; Borovik-Romanov, A.S.; and Orlova, M.P.

Title : Thermodynamic investigations at low temperatures. Part 1.-Measurement of temperatures between 12 and 300° K.

Periodical : Zhur. fis. khim. 28/2, 345-352, Feb 1954

Abstract : A technique was developed for the manufacture of thermometers with international scale graduation. The technique of calibrating thermometers, at a temperature corresponding to the boiling point of hydrogen, is described. A simple way of fixing the scale of a platinum resistance thermometer, by reducing it to the standard table, is explained. The technique described can also be applied in measuring the temperatures between 12 and 300° K with deviations from the thermodynamic scale of about 0.03 - 0.04°. Fifteen references: 8-USSR; 3-USA; 2-German and 2-English (1929-1952). Tables; drawings.

Institution : State Institute of Measures and Measuring Instruments, The S.I. Vavilov Institute of Physical Problems, Moscow

Submitted : June 8, 1953

ORLOVA, M.P. and BOROVIK-ROMANOV, A.S.

"Magnetic Properties of Co and Mu Carbonates and MU-Oxydes"  
Moscow

Conference on Physics of Magnetic Phenomena,  
May 1956, Sverdlovsk, USSR

ORLOVA, M. P.

SUBJECT USSR / PHYSICS CARD 1 / 2 PA - 1780  
AUTHOR BOROVIK-ROMANOV, A.S., ORLOVA, M.P.  
TITLE The Magnetic Properties of the Carbonates of Cobalt and Manganese.  
PERIODICAL Žurn. eksp. i teor. fis., 31, fasc. 4, 579-582 (1956)  
Issued: 1 / 1957

Here the temperature dependence of the magnetic susceptibility of  $MnCO_3$  and of a waterless  $CoCO_3$  preparation is investigated within the temperature range of from 14 to  $300^\circ$ . Three different  $MnCO_3$  samples were investigated: an undried industrial preparation, and the same preparation dried at  $160^\circ C$ ; the third preparation was made by the authors themselves by means of the heating for 20 hours (at  $160^\circ C$ ) of a saturated solution of  $MnCl_2$  with  $CaCO_3$  in a sealed test glass. The values obtained in the case of the first and third sample were always lower than in the case of the second. This is due to the presence of water in the first sample and of  $CaCO_3$ -remains in the second sample. After the necessary corrections the susceptibilities of all three samples were in agreement within the limits of measuring errors throughout the entire temperature range.

Further, two samples of waterless  $CoCO_3$  were examined, which had been produced, like the  $MnCO_3$ , by the heating of  $CoCl_2$  and  $CaCO_3$  in a sealed ampule. The results obtained for both samples were in agreement within the limits of measuring errors. Susceptibilities were measured by FARADAY'S method. The temperature dependence of the magnetic susceptibility of both carbonates at higher temperatures satisfies

Zurn.eksp.i teor.fis, 31, fasc.4, 579-582 (1956) CARD 2 / 2

PA - 1780

the law by CURIE-WEISS  $\chi_m = C_M(T + \Theta)$ . The values of the constants  $C_M$  and  $\Theta$  are shown in a table. The same table contains the values of  $\mu_{\text{eff}}$  computed from the here measured values of  $C_M$  and the theoretical values of  $\mu_{\text{eff}}$  computed on the assumption of the total "freezing in" of the orbital moments. Below a certain critical temperature  $T_c$  permeability increases sharply and depends considerably on field strength. Also a slight hysteresis is found. At temperatures of less than  $T_c$  and at field strengths of more than 600 Oersted the dependence of the magnetic moment  $M$  on the field strength  $H$  can be represented as the sum of two terms  $M = M_0 + \chi'H$ . Similar isotherms were obtained also for  $\text{CoCO}_3$ . The temperature dependence of  $M_0$  in the case of  $\text{MnCO}_3$  and  $\text{CoCO}_3$  has the shape which is characteristic of the "settling curve". However, at  $T \rightarrow 0$  the value of  $M_0$  tends towards a considerably lower value than might be expected in the case of ferromagnetic saturation: For  $\text{MnCO}_3$  :  $M_0 = 68$ ,  $M_{\text{ferr}} = 32000$ , for  $\text{CoCO}_3$  :  $M_0 = 400$  to  $1000$ ,  $M_{\text{ferr}} = 27200$ . The results obtained can be explained qualitatively by assuming that an antiferromagnetic process occurs in manganese- and cobalt manganese below  $T_c$ .

INSTITUTION: All-Soviet Institution for Physical, Technical, and Radiotechnical Measuring.



ORLOVA, M.P.

**AUTHOR:** BOROVIK-ROMANOV, A.S., ORLOVA, M.P. 56-5-49/55  
**TITLE:** The Magnetic Properties of Manganese Oxides at Temperatures of 20 - 300° K. (Magn'nyye svoystva kislorodnykh soedineniy manganitsa pri temperaturakh ot 20 do 300° K, Russian)  
**PERIODICAL:** Zhurnal Eksperim. i Teoret. Fiziki, 1957, Vol 32, Nr 5, pp 1255- 1256 (U.S.S.R.)  
**ABSTRACT:** In connection with the anomalous magnetic behavior of manganese carbonate below 31° K becoming known the following manganese oxides were investigated with a temperature range of 20 - 300° K, on which occasion the following was found with respect to their magnetic behavior.  
1.)  $Mn_3O_4$  has ferromagnetic properties below 42,5° K.  
2.)  $Mn_2O_3$  remains a paramagnetic substance within the entire investigated temperature domain of 20-300° K.  
**ASSOCIATION:** All-Union Scientific Research Institute for Physical-Technical and Radiotechnical Measurements  
**PRESENTED BY:**  
**SUBMITTED:**  
**AVAILABLE:** Library of Congress  
Card 1/1



SOV/115-59-5-14/27

24(8)

**AUTHOR:**

Orlova, M.P.

**TITLE:**

How to Attain the Boiling Temperature of Oxygen

**PERIODICAL:**

Izmeritel'naya Tekhnika, 1959, Nr 5, pp 23-25 (USSR)

**ABSTRACT:**

The reference point at the bottom of the international temperature scale is the temperature prevailing at the equilibrium of fluid oxygen and saturated steam, which is determined by the formula (1). In the course of the development it became desirable to attain this point with the greatest possible accuracy. In the years between 1948-1949, the laboratories of the MGIMIP developed the following method. Oxygen condensating thermometer: (cf. Fig.1) In order to determine the purity of oxygen of different fillings in regard to the difference in the solidity of the vapors, two oxygen condensating thermometers a) and b), which are protected by vacuum covers, were used. The glass of the thermometer contained platinum foils. A description follows. It was found out that a possible impurity causes an error of less than  $2 \cdot 10^{-3}$  degrees. The thermal process in the oxygen tank was investigated. Good results were

Card 1/2

SOV/115-59-5-14/27

How to Attain the Boiling Temperature of Oxygen

gained, when heat exchange with the walls was avoided. The following part of the article describes the principle of measurement in connection with the instrument in fig.1. Fig.2 shows two diagrams of a tank under certain test conditions. Fig.3, as fig.2, shows the difference of temperature of the oxygen and the atmosphere outside, i.e. the balance temperature which corresponds to the atmospheric pressure. The graphs show results of tests with an error calculation. There are 1 diagram, 5 graphs, 1 table, and 1 Soviet reference.

Card 2/2

24(5), 28(2)

SOV/115-59-8-14/33

AUTHOR:

Astrov, D. N., Orlova, M. P., Strelkov, F. G., and Sharevskaya, D. I.

TITLE:

Comparing Low-Temperature Scales of Platinum Resistance Thermometers

PERIODICAL:

Izmeritel'naya tekhnika, 1959, Nr 8, p 29 (USSR)

ABSTRACT:

At the 1958 session of the Konsul'tativnyy komitet po termometrii (Advisory Committee of Thermometry), a comparison of platinum resistance thermometers at temperatures below 90°K was recommended. Complying with this recommendation, the Vsesoyuznyy nauchno-issledovatel'skiy institut fiziko-tekhnicheskikh i radiotekhnicheskikh izmereniy (All-Union Scientific Research Institute of Physical Engineering and Radio Engineering Measurements) and the National Physics Laboratory compared their platinum thermometers. The British platinum thermometer was sent to the USSR, where the authors performed this comparison at 35 temperature points ranging from 10 to 90°K. The comparison was performed in an adiabatic cryostat with a temperature change of  $1 \cdot 10^{-4}$  degree/minute. The experimental characteristics of the

Card 1/3

SOV/115-59-8-14/33

## Comparing Low-Temperature Scales of Platinum Resistance Thermometers

British thermometer with the calibration of the scale of the National Physics Laboratory was compared to the IKh-6 scale. The scale of the British thermometer was obtained by calculations using the "Z-function" tables of the US National Bureau of Standards [Ref 1] and their corrections [Ref 2]. This method is fully satisfactory for the given types of platinum. Although it decreases the range of platinum brands which are applicable in this temperature range. For example, the Soviet industrial platinum "Pobeda" is about equal in purity to the British platinum ( $R_{100^{\circ}\text{C}}/R_{0^{\circ}\text{C}} = 1.39243$  for the "Pobeda" and 1.39250 for the British platinum), and does not satisfy the additional criterion. For this reason, individual calibrations of such platinum thermometers cannot be calculated by the method suggested by the National Physics Laboratory. In addition, the aforementioned method was developed for temperatures of  $90-20^{\circ}\text{K}$ , while presently a scale is

Card 2/3

SOV/115-59-8-14/33  
Comparing Low-Temperature Scales of Platinum Resistance Thermometers

required reaching below  $20^{\circ}\text{K}$ . The deviation between the practical scale IKh-6 and the calibration of the thermometer of the National Physics Laboratory in the range of  $90$  and  $20^{\circ}\text{K}$  is about  $0.01^{\circ}$  according to the authors' data. For completing the comparison of temperature scales below  $90^{\circ}\text{K}$ , direct comparisons of the scales of the National Bureau of Standards and the Soviet scale are required, since these two scales are based on primary measurements with gas thermometers. There are 1 table and 2 references, 1 of which is American and 1 Soviet.

Card 3/3

SOV/115-60-1-16/28

AUTHOR: Borovik-Romanov, A.S., Orlova, M.P. and Strelkov,  
P.G.

TITLE: Establishing a Practical Temperature Scale for the  
10-90° K range. Deviations of the International Tem-  
perature Scale From the VNIIFTRI Group Standard  
Scale and the Thermodynamic Scale Near the Oxygen  
Point.

PERIODICAL: Izmeritel'naya tekhnika, 1960, Nr 1, pp 34-35 (USSR)

ABSTRACT: The VNIIFTRI temperature scale for the 10-95° K range coincides with the International Scale ("MShT") at the boiling-point of oxygen (-182.97° C) except for a discrepancy of 0.01° in the 90-95° K range, which means that the interpolation formula is only suitable for temperatures near 90° K. Former comparisons made by Heuse and Otto /Ref. 37, Keesom and Damers /Ref. 47, and Bricwedde and Höge /Ref. 57

Card 1/3



SOV/115-60-1-16/28

Establishing a Practical Temperature Scale for the 10-90° K Range.  
Deviations of the International Temperature Scale From the VNIIFTRI  
Group Standard Scale and the Thermodynamic Scale Near the Oxygen  
Point

appear to be insufficiently accurate. The authors suggest a better practical scale for the 90-273° K range. Use of the interpolation power formulas is not advised and recommendations are made to establish a scale according to the principle suggested by Strelkov and Sharevskaya /Ref. 67. The VNIIFTRI group standard thermometers were compared at the boiling-point of "natural composition" hydrogen, which was determined as

$$T = 20.39 \pm 0.003$$

This value can differ from the thermodynamic temperature of boiling hydrogen by the value

$$20.39 \left[ \frac{T_{O_2}}{90.19} - 1 \right] \pm 0.006^\circ \text{ K}$$

Card 2/3

SCV/115-60-1-16/28

Establishing a Practical Temperature Scale for the 10-90° K Range.  
Deviations of the International Temperature Scale From the VNIIFTRI  
Group Standard Scale and the Thermodynamic Scale Near the Oxygen  
Point

The article includes a temperature table of the boiling-point of "natural composition hydrogen", measured by different authors, after the phenomenon of ortho-para conversion became known. There are 3 graphs, 1 table and 12 references, of which 5 are Soviet, 4 German, 1 Dutch and 2 unidentified.

Card 3/3

ORLOVA, M.P.

Reproducing temperatures of phase transitions in solid oxygen.  
Izm.tekh. no.2:21-23 F '61. (MIRA 14:2)  
(Oxygen)

ORLOVA, M. P.; ASTROV, D. N.

"Le thermometre a tension de vapeur de l'helium de condensation  
pour la realisation de l'echelle T<sub>58</sub> "  
Report presented at the 6th Session of the Advisory Committee  
on Thermometry to the International Committee on Weights and  
Measures, Sevres, France, 25-27 Sep 62

Institut "ational des Recherches Scientifiques pour les Mesures  
Physiques et Radiotechniques (U. R. S. S.)

U

ASTROV, D. M.; REVA, M. P.; CHAYNOVA, E. I.

"Extension de l'échelle Internationale Pratique de Température au-dessous de  $-152,9^{\circ}\text{C}$  ( $10,1^{\circ}\text{R}$ )"  
Report presented at the 7th Session of the Advisory Committee on Thermometry to the International Commission on Weights and Measures, Sevres, France, 25-27 Sep 67

Institut National des Recherches Scientifiques pour les Mesures Physiques et Radiotechniques (U. R. S. S.)

CHAYVSKAYA, D. I.; ATRHOV, D. N.; BUKHARINOV, A. S.;  
SILINA, M. P.; STASLAV, P. G.

"Qualification de l'échelle pratique de température dans le  
domaine de 10 à 90°K,"  
Report presented at the 7th Session of the Advisory Committee  
on Thermometry to the International Committee on Weights and  
Measures, Sevres, France, 25-27 Sep 1962.

Institut des Mesures Physicotechniques (U. R. S. S.)

ORLOVA, M. P.; ASTROV, D. N.

L'echelle de temperature dans le domaine de  $4,2$  a  $10^0$  K  
Report presented at the 6th Session of the Advisory Committee  
on Thermometry to the International Committee on Weights and  
Measures, Sevres, France, 25-27 Sep 62

Institut national des Recherches Scientifiques pour les  
Mesures Physiques et Radiotechniques (U. R. S. S.)

S/181/62/004/004/035/042  
B102/B104

AUTHORS: Astrov, D. N., Kytin, G. A., and Orlova, M. P.

TITLE: Shift of Curie point of manganese and cobalt carbonates in uniform compression

PERIODICAL: Fizika tverdogo tela, v. 4, no. 4, 1962, 1055-1057

TEXT: The Curie points of  $MnCO_3$  and  $CoCO_3$ , which are weak ferromagnetics, were determined from gravimetrical measurements of the magnetic susceptibilities (Faraday method). The pressure ( $1900 \pm 100 \text{ kg/cm}^2$ ) was exerted according to the Lazarev method (ZhETF, 14, 470, 1944) in a water-filled autoclave of beryllium bronze. The specimens were pressed from powder and coated with a waterproof film of polymerized LF-4 (BF-4). The measurements were carried out in fields of  $\sim 1800$  oe. Correction was made for the susceptibilities of the autoclave and the water. The measuring error was less than  $\pm 1.5\%$ . From the  $\chi_m(T)$  curves plotted with and without compression applied to the specimen, the Curie point shift

Card 1/2



СЕРГЕЕВ И И П

37952

24.5700

3/181/62/104/005/000/001  
B162/B108

Authors: Piterman, N. Sh., Krol', L. Ia., Medvedev, V. A.,  
Sergeev, S. S., No. 1830, 7. 5.

Title: Impurity Hall conductivity in n-type GaAs

Source: Radiotekhnika, v. 4, no. 5, 1968, 1353-1361

Main results are given of measurement of the resistivity  $\rho$ , the Hall coefficient  $R_H$  and the kinetic resistance  $\frac{\rho^2}{R_H}$  on single crystals of n-type GaAs with impurity concentration of  $10^{16} - 10^{17} \text{ cm}^{-3}$ , at which impurities between the impurities and formation of an impurity band not far from the conduction band can be expected. The specimens were prepared by zone melting in a horizontal boat of an ingot of chemically pure GaAs. Analysis of the data show that the single crystal specimens at temperatures below  $50^\circ\text{K}$  display conductivity in the impurity band. This effect is absent in the more contaminated single-crystal and polycrystalline specimens. The Hall mobility in the conduction band

Impurity band conductivity in ...

U/81/62, 004/100, 000/001  
B162/B108

is three to four times greater than in the impurity band. The magnetic resistances of the single-crystal specimens measured in a field of 100 G become negative at temperatures below 10°K, and for a polycrystal  $\frac{1}{\rho}$  over the wide range of 1.7° - 300°K. The conductivity in the impurity band in n-type GaAs does not lead to a change in the sign of the Hall effect at low temperatures, as might have been expected for holes in the impurity band.

f

ADRESA: Institut Fiziko-tekhnicheskikh i radiotekhnicheskikh izmereniy (Institute of Physicotechnical and Radiotechnical Measurements), Moscow

DATA: November 10, 1961 (initially)  
February 14, 1962 (after revision)

Card 2/2

ORLOVA, M.P.; ASTROV, D.N.

Measuring temperatures below  $10^0$  K. Izv. tekhn. no. 8:37-38  
Ag '62. (MIRA 16:4)  
(Thermometry)

ACCESSION NR: AP4018401

S/0120/64/000/001/0230/0232

AUTHOR: Orlova, M. P.; Astrov, D. N.; Medvedeva, L. A.

TITLE: Germanium resistance thermometer for low temperatures

SOURCE: Pribery\* i tekhnika eksperimenta, no. 1, 1964, 230-232

TOPIC TAGS: thermometer, germanium thermometer, low temperature thermometer, Sb alloyed Ge thermometer

ABSTRACT: The thermometer was prepared from Ge alloyed with Sb; the measured carrier concentration was from  $5 \times 10^{16}$  to  $1 \times 10^{17}$   $\text{cm}^{-3}$ . Au-Sb alloy was used for contacts. For 4 months, six thermometers were tested for stability by comparing them with the reference Pt resistance thermometers and with an H condensation-type thermometer at the H-boiling temperature. Nonreproducibility of indication was  $\pm 0.001$  to  $0.002\text{K}$ . The developed Ge thermometer is recommended for a temperature range of 1-35K; it has a small size, good accuracy.

Card. 1/2

ACCESSION NR: AP4018401

high sensitivity, and sufficient ruggedness. However, it is sensitive to magnetic fields, difficult to manufacture with identical characteristics, and there is, as yet, no formula which would establish the resistance-temperature relation. "The authors are thankful to G. A. Kats and O. I. Shibayeva who developed the method and grew Sb-alloyed Ge single crystals, and also to V. I. Petrov for his part in preparing the thermometers." Orig. art. has: 4 figures.

ASSOCIATION: Vsesoyuznyy nauchno-issledovatel'skiy institut fiziko-tekhnicheskikh i radiotekhnicheskikh izmereniy (All-Union Scientific-Research Institute of Physico-Technical and Radio-Technical Measurements)

SUBMITTED: 21Dec62

DATE ACQ: 18Mar64

ENCL: 00

SUB CODE: PH

NO REF SOV: 002

OTHER: 003

Card 2/2

ORLOVA, M.P.

Session of working groups of the Consultative Committee on Thermometry.  
Izm.tekh. no.2:50-51 F '64. (MIRA 17:4)

I 11393-63

BDS

S/120/63/000/002/034/041

50

**AUTHOR:** Orlova, M. P., Astrov, D. N., and Medvedeva, L. A.

**TITLE:** An indium resistance thermometer for 3.4-300°K temperatures

**PERIODICAL:** Priory i tekhnika eksperimenta, March-April 1963, v. 8, no. 2, 160-163

**TEXT:** The article describes resistance thermometers using extremely pure metallic indium; these instruments have a higher resistance (10-25 ohms at 0°C) and are less cumbersome than earlier instruments. The measurement range of these thermometers is 3.4-300°K and their stability is at least 0.0015°K. The authors give suggestions for calibrating the thermometers without comparison with primary instruments over a small temperature interval. There are eight figures.

**ASSOCIATION:** Vsesoyuznyy nauchno-issledovatel'skiy institut fiziko-tekhnicheskikh i radiotekhnicheskikh izmereniy (All-Union Scientific Research Institute for Physicotechnical and Radiotechnical Measurements)

**SUBMITTED:** April 12, 1962

Card 1/1 Jm/ck

ORLOVA, M.P.

Development of the research in the field of low-temperature measurements in the U.S.S.R. and abroad. Izv.tekh. no.6:22-29 Je '64.

(MIRA 17:12)



L 34868-66 EWT(d)/EWP(v)/EWP(k)/EWP(h)/EWP(l)

AEC NR: AI'6014518

SOURCE CODE: UR/0115/65/000/011/0008/0010

AUTHOR: Orlova, M. P.; Konoplev, V. A.; Sharevskaya, D. I.; Astrov, D. N.;  
Al'shin, B. I.; Medvedeva, L. A.

ORG: none

TITLE: New commercial resistance thermometer

SOURCE: Izmeritel'naya tekhnika, no. 11, 1965, 8-10

TOPIC TAGS: resistance thermometer, temperature measurement, low temperature research / PTS-100 resistance thermometer

ABSTRACT: As the PTS-100<sup>†</sup> standard platinum resistance thermometer<sup>ρ</sup> (10-300K, ± 0.01K) is suitable only for operating under laboratory conditions, two new high-accuracy designs have been developed by the authors for industrial uses. In the first design (see Figure 1), coil 1 is fastened by the glass coating of straight platinum wire 2. Four such vitrified coils constitute the sensor of the thermometer. Platinum supporting wires are used as lead-ins 4 in envelope 3 filled by

Card 1/2

UDC: 536.531

L 34868-66

ACC NR: AP6014518

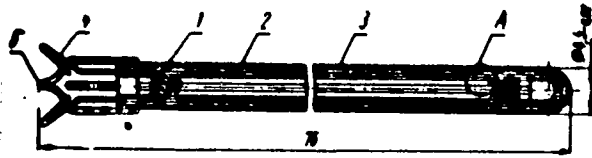


Fig 1



Fig 2



helium through throat 5. In the second design (see Figure 2), a straight 0.1-mm glass thread 2 is placed inside platinum coil 1. The latter is mounted in glass capillary 3; envelope 4 and platinum lead-ins 5 are conventional. The above designs were tested for vibration (50-3000 cps) and temperature stability (250, 100, 0C; H boiling and

triple points). Their thermal inertia was 5-8 sec. Orig. art. has: 2 figures and 1 table.

SUB CODE: 20, /4 / SUBM DATE: none / ORIG REF: 001 / OTH REF: 002

Card 2/2 vmb

ORLOVA, M.P.; KONOPLEV, V.A.; SHAREVSKAYA, D.I.; ASTROV, D.N.;  
AL'SHIN, B.I.; MEDVEDEVA, L.A.

New industrial resistance thermometers. Izv. tekhn. no. 11:  
8-10 N '65. (MIRA 18:12)

ORLOVA, M.T.

Accessory minerals of the ancient meta strata of the western slope  
of the Southern Urals. Mat.VSEGEI.Ob.ser. no.28:31-43 '60.  
(MIRA 14:6)

(Ural Mountains—Mineralogy)

MOSKALEVA, S.V.; ORLOVA, M.T.

Genesis of garnetiferous pyroxenite of the Krak massif in the  
Southern Urals. Mat.VSEGEI.Ob.ser. no.28:143-147 '60.

(MIRA 14:6)

(Ural Mountains--Garnet) (Ural Mountains--Pyroxenite)

ORLOVA, M V

NIKRA SOV, V.P.; ORLOVA, M.V. (Moskva)

Family epidemic of acute pancreatitis. Klin.med. 36 no.2:124-127  
P '57. (MIRA 11:4)

1. Iz Glavnogo voyennogo gosпитalya imeni akad. N.N.Burdenko  
(nach. H.M.Nevskiy)  
(PANCREATITIS, case reports  
acute, familial (Rus))

ASHMARIN, Yu.R., kand.med.nauk; ORLOVA, M.V. (Moskva)

Case of combined herpes zoster and chickenpox. Klin.med. no.3:  
143-146 '62. (MIRA 15:3)  
(CHICKEN POX) (HERPES ZOSTER)

ORLOVA, M. Ye.

Orlova, M. Ye. "Experience in using purified gonococcus antigens in diagnosing female gonorrhoea" (From a candidate's dissertation,) Sbornik nauč. tr. (Rec. obl. nauch.-issled. arushers o-grebol. in-t), Issue 2, 1943, p. 11-14.

So: U-3261, 10 April 1963 (Letopis 'Zhurnal 'nykh Statey, No. 12, 1943).



ORLOVA, M. Ye.

Zaleskaya, M. A. and Orlova, M. Ye. "Cytobacterioscopy of cervical canal discharges in gonorrhoea", Sbornik nauch. trudov (Rost. obl. nauch.-issled. akushersko-ginekol. in-t), Issue 8, 1948, p. 56-60.

So: U- 261, 10 April 1949 (Letopis 'Zhurnal 'nykh Statey, No. 10, 1949).

Orlova, M. Ye.

Orlova, M. Ye. "The degeneration of the gonococcus in discharges from the female sexual organs", Sbornik nauch. trudov (Rest. o l. nauch.-issled. akusersko-ginekol. in-t), Issue 8, 1948, . 72-76.

So: U-3201, 10 April 1953 (Letopis Zhurnal Inykh St. tey, No. 12, 1949).

ORLOVA, M.Ye.

Evolution of seg chromatin in the neutrophils of the blood;  
preliminary report. Probl.gemat.i perel.krovi no.7:24-28  
'62. (MIRA 15:9)

1. Iz Rostovskogo-na-Donu respublikanskogo nauchno-issledc-  
vatel'skogo instituta akusherstva i pediatrii (dir. F.S. Kara-  
novskaya, nauchnyy rukovoditel' - prof. T.V. Loverdo) Mini-  
sterstva zdravookhraneniya RSFSR.  
(LEUCOCYTES) (CHROMATIN)

ORIOVA, N., kand.med.nauk

Beware of grippe! Okhr. truda i sots. strakh. 6 no.12:19-22 D  
'63. (MIRA 17:2)

1. Zaveduyushchaya laboratoriyey profilaktiki zabolevaniy dy-  
khatel'nykh putey Instituta virusologii AMN SSSR.

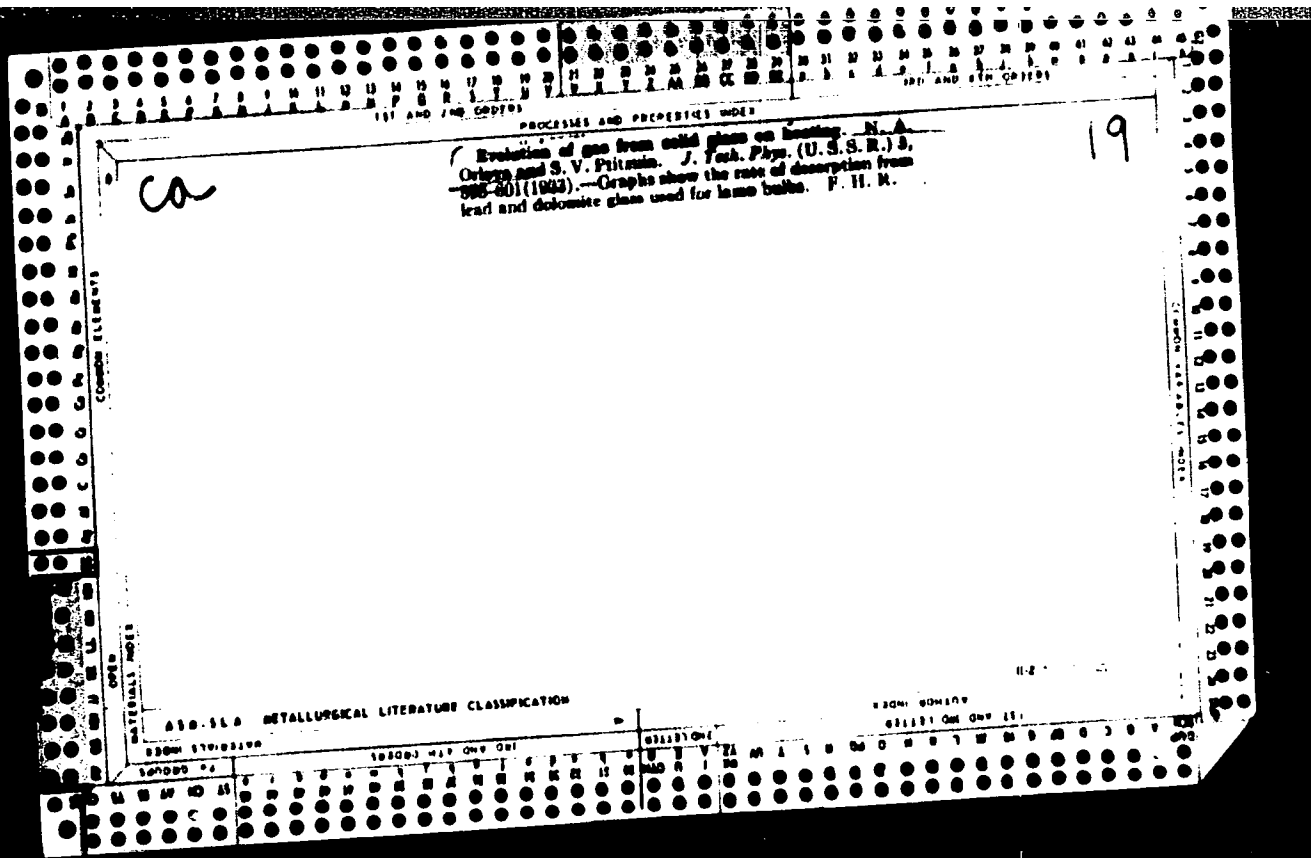
GAN, P.A.; DZIANAYEVA, V.M.; KARAFI-KORBUT, I.G.; KRIVOSHEYEVA, L.S.;  
KUNICHENKO, A.I.; ORLOVA, N.A.; PROTOPOPOV, G.F.; PRUTENSKIY,  
D.I.; TKACHENKO, V.I.; SOROKHAYEVA, H.V., red. izd-va; POPOVA,  
M.G., tekhn. red.

[Trees and shrubs of Kirghizia]Derev'ia i kustarniki Kirgizii.  
Frunze, Izd-vo Ak. Kirgizekoi SSR. No.2. [Families: Liliaceae-  
Moraceae].Semeistva lileinye-tutovye. 1961. 211 p.

(MIRA 15:10)

1. Akade: iya nauk Kirgizskoy SSR, Frunze. Institut botaniki.  
Sektor lesa.

(Kirghizistan--Angiosperms)



ORLOVA, N.

Microdetermination of nitrogen in petroleum and its products. S. Dalbaeva and N. Orlova. *Novosti Neftyanol Tekh., Neftepereabotka* 1955, No. 23-24. — The Kjeldahl method was adapted for the detn. of N in petroleum fractions, especially heterocyclic N compds. A 20-200-mg. sample in a boat was placed in the Kjeldahl flask to which were added 4.6% H<sub>2</sub>SO<sub>4</sub> (d. 1.84), glucose 200, CuSO<sub>4</sub> 40, K<sub>2</sub>SO<sub>4</sub> 200, and Se 40 mg., and the mixt. was heated 1.5 hrs. at 400-20°. After cooling two 2-ml. portions of 0.1M KMnO<sub>4</sub> were added. After each addn. the mixt. was heated for 30 min. The contents of the flask were then transferred into the Parnas-Wagner distn. app. (cf. C.A. 44, 480f), treated with 10 ml. NaOH solu. (3%), and NH<sub>3</sub> was detd. by the usual procedure. The reproducibility of this method was 0.01% for a N concn. in petroleum from 0.01% to 0.1%.

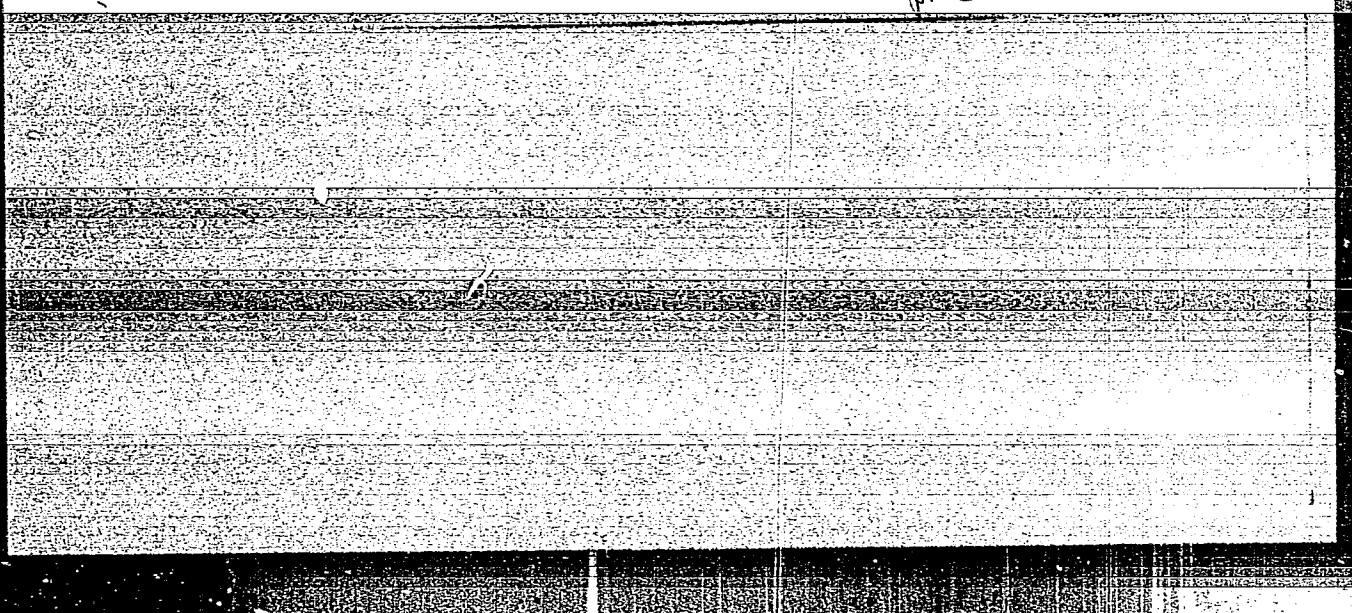
File

12

PM

ORLOVA, N. A. Cand Tech Sci -- (diss) "Salification as <sup>a</sup> ~~the~~ means of antifiltration  
for ~~minor~~ <sup>small</sup> irrigation canals of the south <sup>ern</sup> ~~of the~~ Ukraine." Kiev, 1956. 14 pp  
20 cm. (Min of Agr USSR. All-Union Sci Res Inst of Hydraulic Engineering and  
Improvement VNIIGIM), 100 copies. (KL, 13-57, 99)





ORLOVA, N. A.

DSSR/ Chemistry - Catalysis

Card 1/2 Pub. 22 - 17/43

Authors : Bashkirov, A. N.; Khotinskaya, M. I.; and Orlova, N. A.

Title : Oxygen-containing compounds formed during the synthesis of hydrocarbons from carbon monoxide and hydrogen over an iron catalyst

Periodical : Dok. AN SSSR 106/1, 65-68, Jan 1, 1956

Abstract : Experiments were conducted to separate and identify oxygenous compounds contained in the hydrocarbon part of a product formed during the synthesis from CO and H over an iron catalyst at an average pressure and 1 : 1 ratio of the CO to H in the initial gas. The acid, hydroxyl, carbonyl and ether numbers of the hydrocarbon fraction investigated are listed. Neutral

Institution : Acad. of Sc., Institute of Petroleum

Presented by: Academician A. V. Topchiyev, May 27, 1955

Card 2/2 Pub. 22 - 17/43

Periodical : Dok. AN SSSR 106/1, 65-68, Jan 1, 1956

Abstract : oxycoumpounds were separated from the investigated product by means of chromatographic adsorption on a ASM-silica gel contact with the grain dimension of 60 - 100 mesh and activity of 17.1. The ketones contained in the product were separated through semicarbazones. Four references: 2 USSR and 2 Eng. (1949-1954). Tables.

ORLOVA, N.A.

KAGAN, Yu.B.; BASHKIROV, A.N.; ZVEZDKINA, L.I.; ORLOVA, N.A.; KLIGER, G.A.

Influence of reduction conditions on the properties of molten iron catalysts used in alcohol synthesis from carbon monoxide and hydrogen. Trudy inst. nefti. 10:262-268 '57. (MIRA 11:4)  
(Alcohols) (Carbon monoxide) (Hydrogen)

AUTHORS: Kagan, Yu. B., Lashkirov, A. I., SOV/02-10-10-11/85  
Kryukov, Yu. B., Loktev, S. M., Orlova, N. A.

TITLE: On the Mechanism of the Catalytic Efficiency of Fused  
Iron Catalysts in the Synthesis of CO and H<sub>2</sub> (О механизме каталитического действия сплавленных железных катализаторов синтеза из CO и H<sub>2</sub>)

PERIODICAL: Izvestiya Akademii nauk SSSR. Otdel'niye khimicheskiye nauki,  
1980, No 10, pp 1274 - 1275 (USSR)

ABSTRACT: In an earlier paper the authors showed that immediately after  
the reduction (by hydrogen at 1000°) fused iron cataly-  
sts in the hydrocarbon synthesis of CO and H<sub>2</sub>  
are not active any more. Only under the working  
conditions of the synthesis when the gas mixture CO+H<sub>2</sub>  
is passed through the catalyst gradually becomes active  
(for 10-20 hours). This phenomenon may be explained  
by a number of simultaneous reactions competing with  
each other. Due to the course of these reactions com-  
peting with each other the metallic iron regenerates  
often (under the conditions of the synthesis) from its

Card 1/2

On the Mechanism of the Catalytic Efficiency of Fused Iron Catalysts in the Synthesis of CO and H<sub>2</sub> SVV/2-1-1-1/23

compounds, and at the surface of the operating catalyst the dynamic equilibrium of the surface phases of different chemical structure is obtained. As a consequence of these processes the activation of the catalyst occurs. Neither the iron itself nor compounds that might be formed from it are the reason for the activation of the catalyst surface. The hypothesis formed for the main mechanism of the catalytic efficiency of iron catalysts (according to which the synthesis of CO and H<sub>2</sub> is caused by the reactions of carbon and hydrogen monoxide with iron and its compounds on the surface of the operating catalyst) was described in detail by the authors. There are 1 table and 1 reference, which is Soviet.

ASSOCIATION: Institut nefti Akademii Nauk SSSR (Petroleum Institute AS USSR)

SUBMITTED: April 8, 1958  
Card 2/2

KAGAN, Yu.B.; BASHKIROV, A.N.; ZVEZDKINA, L.I.; ORLOVA, N.A.

Fused iron catalysts in the synthesis of higher alcohols from carbon  
monoxide and hydrogen. Trudy Inst.nefti 12:200-212 '58.  
(MIRA 12:3)

(Alcohols) (Catalysts)

KAGAN, Yu.B.; BASHKIROV, A.N.; LOKTEV, S.M.; MOROZOV, N.G.; ORLOVA, N.A.

Effect of the introduction of ferroalloys on the activity and stability  
of fused iron catalysts for synthesis based on CO and H<sub>2</sub>. *Trudy Inst.  
nefti* 12:228-239 '58.

(MIRA 12:3)

(Catalysts) (Iron alloys) (Chemistry, Organic--Synthesis)



3(5)

AUTHORS:

Chepikov, K. R., Corresponding Member *SOV/20-125-5-39/51*  
AS USSR, Yermolova, Ye. P., Orlova, N. A.

TITLE:

Epigene Minerals as Time Indicators of the Petroleum Appearance in Sandy Reservoirs Capable of Exploitation (Epigennyie mineraly kak pokazateli vremeni prikhoda nefti v peschanyie promyshlennyye kollektory)

PERIODICAL:

Doklady Akademii nauk SSSR, 1959, Vol 125, Nr 5, pp 1097-1099 (USSR)

ABSTRACT:

The problem of the time indication mentioned in the title is of great significance both practically and theoretically. In order to determine this, the authors have studied the character of the correlation between petroleum and the epigene minerals. Samples of petroleum containing rocks of the largest petroleum fields in the Volgo-Ural'skaya (Volga-Ural) region served for this purpose: Romashkino, Bavly, Tuymazy, in addition to the Yablonovyy and Zol'nyy ravines. From there came the quartz sandstones of the Pashiyskaya suite, while samples of Lower Carboniferous rocks came from Mukhanovo. The clastic material of the sand and "aleuritic" rocks is almost exclusively formed by quartz (95-98 %). Only non-clayey varieties were studied

Card 1/4

Epigene Minerals as Time Indicators of the SOV/20-125-5-39/61  
Petroleum Appearance in Sandy Reservoirs Capable of Exploitation

whose cement consists of epigene minerals. Most of the reservoirs investigated had high effective porosity values ( $P_{eff} = 17-22\%$ ). There were also, however, sandstones with a slight  $P_{eff}$  value, and even impermeable varieties. The roll played by the individual epigene minerals in the cementation of the non-clayey varieties of sandstone and "aleurite" is different. Regenerated quartz takes the first and most important place; it generally penetrates through the entire rock. Other epigene minerals occur only as local precipitations and have a limited distribution. The epigene quartz is mostly precipitated as a regeneration overgrowth of various thicknesses. It often binds only the clastic grains and only slightly fills the pore space. Carbonates (calcite and dolomite) as well as anhydrite, cement the sandstones and "aleurites" only superficially. As a rule, all of the minerals mentioned corrode the clastic and regenerated quartz. Often they replace it completely, in which case they indicate a basic cementation type. The genesis of the epigene minerals is briefly discussed. If pyrite, whose genetic conditions deviate from those of the

Card 2/4

Epigene Minerals as Time Indicators of the  
Petroleum Appearance in Sandy Reservoirs Capable of Exploitation

SOV/20-125-5-39/61

other minerals, is excluded from the observed formation sequence - quartz, pyrite, carbonate, anhydrite - the sequence of the precipitation of the remaining minerals agrees well with their increasing solubility. The regeneration process is discussed. The analysis of the formation sequence of the epigene minerals in quartz sandstones and "aleurites" has shown that the last minerals precipitated (the carbonates and anhydrites) represent new formations which originated at considerable depths. Petroleum filled all of the freely interconnecting pore channels in all samples of sandstones and "aleurites" which were already earlier cemented by epigene minerals. The form of the petroleum inclusions is determined here by the morphology of the pore space. As a rule, epigene minerals contain no petroleum inclusions; at most they have thin petroleum films on the contact of the quartz grains with the epigene carbonate and anhydrite cement in fractures and individual calcite, dolomite, and anhydrite crystals. This can be utilized as an indication that the petroleum has not filled the pore spaces until after the precipitation of the entire complex of epigene minerals. Consequently the petroleum has a younger age. By comparison

Card 3/4

Epigene Minerals as Time Indicators of the Petroleum      SOV/20-125-5-39/61  
Appearance in Sandy Reservoirs Capable of Exploitation

of the assemblage and the intensity degree of the mineral formation in water and petroleum. The same assemblage of epigene minerals is encountered again and again; additionally, their precipitation is always the same (Mukhanovo), however, the precipitation of calcite and dolomite in water bearing rocks is more intense. This is due to the fact that the formation of these minerals had come to a stand still in petroleum bearing rocks, while it continued for a time in the water bearing rocks.

SUBMITTED:      December 9, 1958

Card 4/4

CHEPIKOV, K.R.; YERMOLOVA, Ye.P.; ORLOVA, N.A.

Epigenetic minerals in arenaceous rocks of producing horizons  
and their effect on reservoir properties of rocks as revealed by  
the studies in Second Baku. Trudy Inst. geol. i razrab. gor.  
iskop. 1 '60. (MIRA 14:1)

(Second Baku--Petroleum geology)

BOGOMOLOVA, A.F. (Moskva); ORLOVA, N.A. (Moskva)

Quantitative characteristics of the structure of pore space. PMPF  
no.4:77-81 JI-Ag '61. (MIRA 14:10  
(Porous materials)

BOGOMOLOVA, A.F.; ORLOVA, N.A.

Study of the structure of a porous area. Geol. nefti i gaza  
5 no.12:46-49 D '61. (MIRA 14:11)

1. Institut geologii i razrabotki goryuchkikh iskopayemykh AN  
SSSR.

(Porosity)

CHEPIKOV, K.R.; YERMOLOVA, Ye.P.; ORLOVA, N.A.

Corrosion of quartz grains and cases of a possible effect  
of petroleum on the reservoir properties of sandy rocks.  
Dokl. AN SSSR 140 no.5:1167-1169 O '61. (MIRA 15:2)

1. Institut geologii i razrabotki goryuchikh iskopayemykh  
AN SSSR. 2. Chlen-korrespondent AN SSSR (for Chepikov).  
(Petroleum geology)



CHEPIKOV, K.R.; YERMOLOVA, Ye.P.; ORLOVA, N.A.

Variations in the porosity of sandstones with depth. Dokl. AN  
SSSR 144 no.2:435-437 My '62. (ADRA 1500)

1. Institut geologii i razrabotki goryuchikh iskopayemykh.
2. Chlen-korrespondent AN SSSR (for Chepikov).  
(Sandstone)

CHEPIKOV, K. R.; YERMOLOVA, Ye. P.; ORLOVA, N. A.

"Authigenic minerals in oil-bearing terrigenous rocks."

report submitted for 22nd Sess, Int. Geological Cong, New Delhi, India, Dec  
1964.

TYULENEV, N.A., doktor sei'khod. nauk, prof., otv. red.;  
ALIAT'YEV, S.M., kand. sei'khod. nauk, otv. red.;  
IATA, I.S., kand. sei'khod. nauk, red.; KISHIN'KO, I.Y.,  
k.i., kand. tekhn. nauk, red.; TROVAKIN, B.I., kand.  
tekhn. nauk, red.; ORLOV, V.A., kand. sei'khod. nauk, red.;  
KISHIN'KO, I.Y., kand. tekhn. nauk, red.;  
KOROTKIY, V.I., kand. tekhn. nauk, red.; GUSHKO, I.Y., red.

Materialy konferentsii "Voprosy razvitiya i  
the Field of the ... ..  
ob"edineniya ... .. -tekhnicheskoi konferentsii molodyzhi na  
ustroystva raznykh v oblasti melioratsii i gidrotekhniki.  
Kiev, Ukr. zool. ... ..

1. Ob"edineniye ... ..  
v oblasti melioratsii i gidrotekhniki ... ..  
konferentsii ... ..

KAGAN, S.Z.; KOVALEV, Yu.N.; KACAN, Yu.B.; ORLOVA, N.A.

Studying the extraction of higher alcohols from their mixtures  
with hydrocarbons. Trudy MKHTI no.40/128-133 '67.

(MIRA 18:12)

Citation:

Infrared and Raman spectra and the rotational structure of  
in liquids. Methane and perdeuterated methane. *J. Chem. Phys.*  
17:30-34, 1949. (1949)

*ORLOVA, N. D.*

LEVINA, R.Ya.; KIM DYAY GIR, SMIRNOVA, E.N.; ORLOVA, N.D.; TRESHCHOVA,  
Ye.G.

Synthesis of hydrocarbons. Part 59: (0,1,3)-bicyclohexanes with  
two quaternary carbon atoms. Zhur.ob.khim. 27 no.7:1779-1783  
Jl '57. (MIRA 10:10)

1.Moskovskiy gosudarstvennyy universitet.  
(Bicyclohexane)

ORLOVA, N. D.

Synthesis of hydrocarbons. LIX. Bicyclo[0.1.3]hexanes with two quaternary carbon atoms. R. Ya. Levina, Dyal-Gir Kim, R. N. Sviridova, N. D. Orlova, and E. G. Treshchova (State Univ., Moscow). *Zh. Obshch. Khim.* 27, 1779-83 (1957); *cf. C.A.B.* 50, 14500b; 51, 4027d. Treatment of 1-methyl-1-cyclohexene-3-one with RMgX gave the following mixts. of 1-methyl-3-alkyl-1,3-cyclohexadienes, contaminated with 1-methyl-3-alkylidencyclohexenes (alkyl group shown): 63% Et, bp 155-61°, n<sub>D</sub><sup>20</sup> 1.4872, d<sub>4</sub> 0.8472; 58% Pr, bp 173-83°, 1.4891, 0.8483; 47% Bu, bp 195-201°, 1.4900, 0.8478; 19% iso-Pr, bp 180-9°, 1.5003, 0.8099. These were hydrobrominated and treated with Zn dust in 95% EtOH yielding after distn. either completely pure bicyclo[0.1.3]hexane homologs or contg. at most 1% unsatd. isomers: 30% 1-methyl-3-ethylbicyclo[0.1.3]hexane, bp 133.5-4°, 1.4406, 0.8106 (principal Russian lines 670(0), 833(5), 702(5)); 3-Pr analog, 18%, bp 158.5-57°, 1.4435, 0.8171 (671(5), 697(3.5), 735(3.5)); 3-Bu analog, 11%, bp 179.5-9.8°, 1.4471, 0.8201 (676(0), 687(6), 734(4)); iso-Pr analog 7%, bp 150.5-50.6°, 1.4433, 0.8170 (282(0.5), 512(3.5), 664(12), 807(8), 333(4), 702(16), 853(3.5)). High-boiling mixts. were also formed in each case but pure monocyclic hydrocarbons could not be isolated from them. LX. Homologs of ethylbenzene from adducts of alkenes with methyl-ethylmaleic anhydride. V. R. Sevast'yanov, R. Ya. Levina, and M. G. Kuz'min. *Ibid.* 1781-7. Gunking AcCHEE-CO. Et 12 hrs. with 300 cc. H<sub>2</sub>O.

14  
 4E4y  
 4E3d  
 4E2c(j)  
 2 may

TWT  
 01

and treatment of the aq. phase with 50 g. NaCN for 2 hrs.  
and diln. with H<sub>2</sub>O, gave an org. layer which was refluxed 30

Levina, R. M.; Kim, Dhan-Gin; Smirnova, E. N.; Orlova, N. O;  
hrs. with 2 vols. concd. HCl and dry-distd. residue



Orlova, N. D.

USSR/ Chemistry - Synthesis

Card 1/1 Pub. 22 - 28/54

Authors : Levina, R. Ya.; Shusherina, N. P.; Lurye, N. Yu.; Orlova, N. D.

Title : Cyanethylated ketones in the synthesis of unsaturated lactones. Delta lactones with semicyclic double bond

Periodical : Dok. AN SSSR 106/2, 279-282, Jan 11, 1956

Abstract : The synthesis of delta-lactones with semicyclic double bond from basic ketones (diisopropyl- and methylisopropyl ketone) is described. The lactones obtained were found to have a definite semicyclic bond inasmuch as their formation was due mainly to the ketoenol regrouping of the keto acids, the enol forms of which are formed only as result of the isopropyl group and methyl group, respectively. The chem. properties of synthesized lactones are described. Eight references: 4 USSR, 2 USA and 2 French (1899-1955).

Institution : Moscow State University im. M. V. Lomonosov

Presented by: Academician B. A. Kazanskly, July 16, 1955

ORLOVA N.D.

**AUTHOR:** Bulanin, M.O. and Orlova, N.D.

**TITLE:** Investigation of the Changes of the Rotation-Vibration Spectrum of Certain Simple Molecules Upon Dissolution (Issledeniye izmeneniy vrashchatel'no-kolebatel'nogo spektra rektortopov prostykh molekul pri rastvorenii)

**PERIODICAL:** Optika i Spektroskopiya, 1968, Vol IV, Nr 5, pp 662-671 (USSR)

**ABSTRACT:** Transition from the vapour state to liquid or solution is normally accompanied by disappearance of the rotational structure of bands in the vibrational spectrum. This effect is ascribed to the absence of free rotation of molecules in liquids. The rotational motion is least affected on liquefaction of hydrogen halides. In liquid oxygen, nitrogen and methane the discrete rotational spectrum is absent (Ref 2) and only the form of bands in the Raman scattering indicates certain freedom of rotation of the molecules. No rotational structure was discovered in the infrared solutions. The present paper reports results of studies of the infrared absorption spectra of solutions of hydrogen halides (HCl, DCl, HBr, DBr, HF) and of water. The measurements were made using infrared spectrometers with LiF and NaCl prisms.

Card 1/3

Investigation of the Changes of the Rotation-Vibrational Spectra of Certain  
Simple Molecules Upon Dissolution

The spectrometers were calibrated using the absorption spectra of gaseous  $H_2O$ ,  $CO$ ,  $HBr$ ,  $HCl$  and  $NH_3$ . The error in determination of frequency did not exceed  $5\text{ cm}^{-1}$ . Thin layers of solutions were used ( $d = 0.05\text{--}8.0\text{ cm}$ ). The spectra of the following solutions were obtained:  $HCl$  in  $CCl_4$ ,  $SiCl_4$ ,  $TiCl_4$ ;  $DCl$  in  $CCl_4$ ,  $SiCl_4$ ,  $PCl_4$ ,  $CHCl_3$ ;  $HBr$  and  $DBr$  in  $CCl_4$ ;  $HF$  in  $C_5F_{12}$ ;  $H_2O$  in  $C_2Cl_2F_6$ ,  $C_2Cl_3F_2$ ,  $CCl_4$ ,  $C_2Cl_4$ ,  $CHCl_3$ ,  $CH_3NO_2$  and  $D_2O$  in  $C_2Cl_3F_2$ , all at room temperature, as well as  $HCl$  and  $Hbr$  in  $CCl_4$  at  $75^\circ C$  and in  $CH_3NO_2$  at  $90^\circ C$ . Some of the spectra obtained are shown in Figs 1-5. Figs 1, 2 and 3 show that the absorption bands of all hydrogen halides in solution consist of a central peak and additional maxima on both sides of the peak. The positions of the central peak and the two additional maxima, denoted by  $\nu_0$ ,  $\nu_-$  and  $\nu_+$ , are given in Table 1. In the spectra of solutions of water (Figs 4 and 5) two bands were observed, corresponding to symmetrical ( $\nu_1$ ) and antisymmetrical ( $\nu_3$ ) valence vibrations. The antisymmetrical band is more intense. It is concluded that the observed additional absorption bands in the substances studied are not due to internal molecular degrees of freedom. Comparison of the solution spectra with the spectra of gases indicates that the additional absorption bands are really not

Card 2/3

Investigation of the Changes of the Rotation-Vibration Spectrum of Certain  
Simple Molecules Upon Dissolution

of rotational branches. This indicates that almost free rotation of the solute molecules is possible in solutions. On increase of interaction of the solute with the solvent rotation is transformed INTO librational motion. The author thanks V.I. Chulanovskiy who directed this work. There are 5 figures, 2 tables and 26 references, of which 10 are American, 2 English, 5 Soviet, 3 French, 1 German and 1 Canadian.

ASSOCIATION: Fizicheskiy institut Leningradskogo gosudarstvennogo Universiteta  
(Physical Institute, Leningrad State University)

SUBMITTED: July 5, 1957

- Card 3/3
1. Molecules - Rotation - Vibration Spectra. Halides - Infrared absorption spectra
  3. Spectrometers - Applications