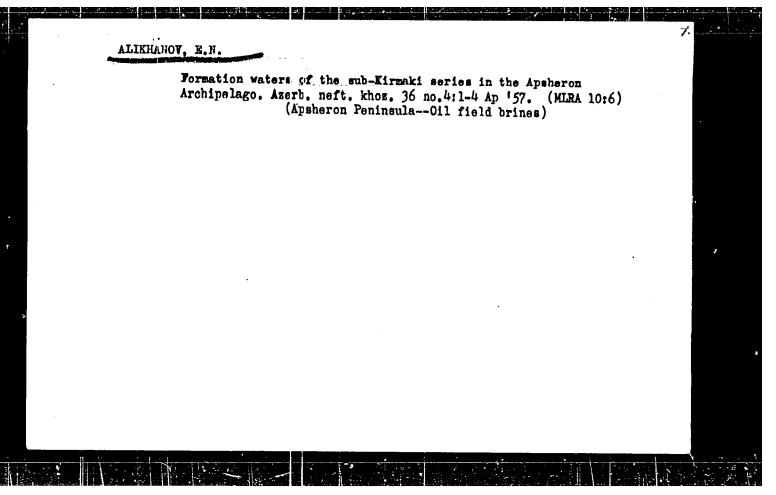
ALIVHANOV. Rever Nagara Ashard RIN, V.A., professor, redaktor; GONCHAROV,

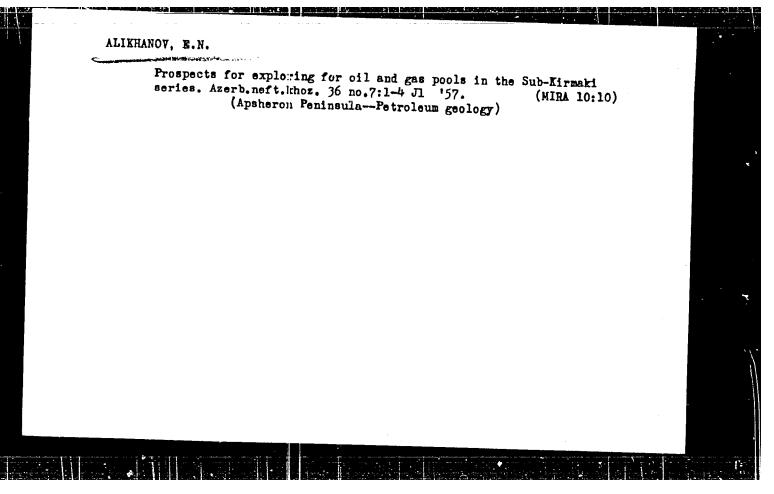
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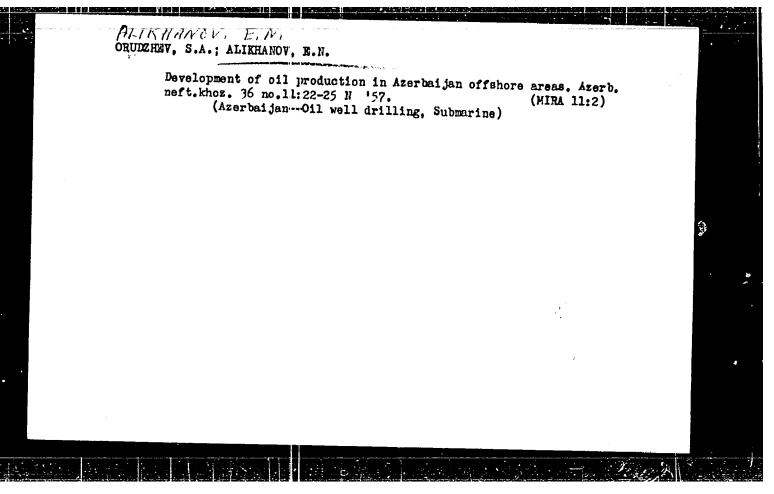
[Sub-kirmaki series of the eastern part of Apsheron Province and its oil bearing possibilities] Podkirmakinskais svita vostochnoi chasti Apsheronskci oblasti i ee neftenosnost'. Baku. Azerbaidzhanskoe gos.izd-vo neft. i nauchno-tekhn. lit-ry, 1957. 215 p.

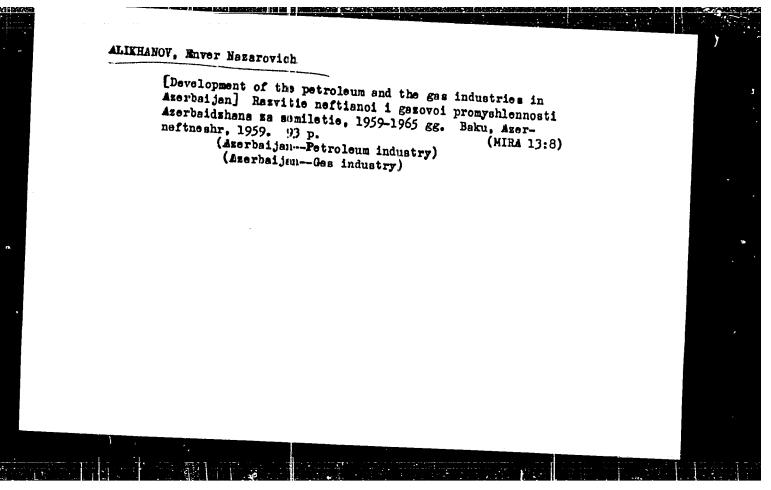
(MIRA 10:9)

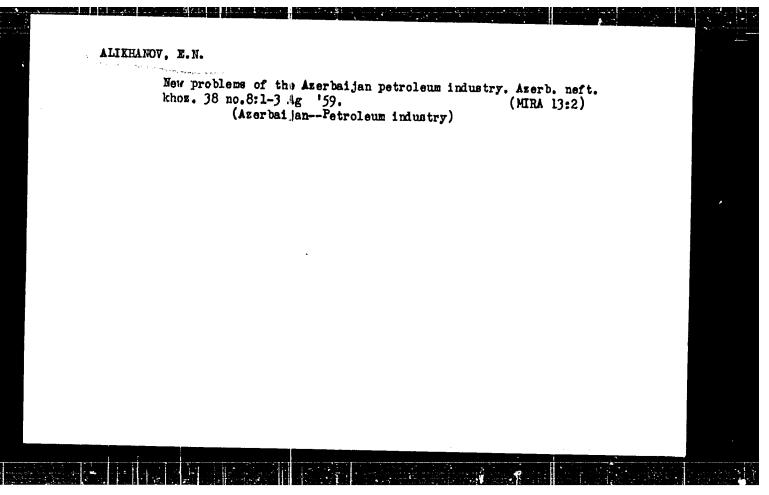
(Apsheron Frovince--Petroleum geology)











ALIERANOV, E.N.; KAUFMAN, V.P.

Improvement of the wage system in drilling departments is an urgent problem, Azerb.meft.khoz. 38 no.11:45-48 H 59. (MIRA 13:5)

(O11 well drilling) (Weges)

ALIKHAMOV, E.N.; KULIYEV, I.P.; SAMEDOV, F.I.

Characteristics and principles of the efficient development of offshore petroleum fields. Sov.geol. 4 no.10:100-107 0 '61.

(MIRA 14:11)

1. Gosudarstvennyy nauchno-issledovatel'skiy i proyektnyy institut morskey nefti.

(Amerbaijan--Oil well drilling, Submarine)

ALIKHANOV, F.M.; ARUSHANOV, N.A.; AKHUNDOV, V.Yu.; ALIZADE, M.A.; AZIZBEŁOV, Sh.A.; FAGIROV, M.A.; VEZIROV, S.A.; VOLOBUYEV, V.R.; EŁKILOV, F.M.; GADZHIYEV, N.M.; CUSEYNOV, D.M.; GUSEYNOV, I.A.; DADASHEV, K.K.; DADASHZADE, M.A.; DALIN, M.A.; ISKENDEROV, M.A.; KAZIYEV, M.A.; KARAYEV, A.I.; KASHKAY, M.S.; KEL'DYSH, M.V.; KERIMOV, A.G.; LEMBERANGKIY, A.D.; MAMEDOV, G.K.; MEKHTIYEV, M.R.; KIRZOYEV, S.A.; NAGIYEV, M.F.; NESRULLAYEV, N.I.; ORUDZHEV, A.K.; RADZHAŁOV, R.A.; RUDNEV, K.N.; SADYKHOV, R.N.; SEMENOV, N.N.; TOFCHIYEV, A.V.; TOPCHIBASHEV, M.A.; TAIROVA, T.A.; KHALILOV, Z.I.; EFENDIYEV, G.Kh.; SHUKYUROVA, Z.Z.

IUsif Geidarovich Mamedaliev; obituary. Dokl. AN Azerb. SSR 17 no.12:1123-1126 '61. (MIRA 15:2) (Mamedaliev, Iusif Geidarovich, 1905-1961)

(Azerba: janIndustries)	[Forty years of Azerbaijani industry; 1920-1960]Promys nost' Azerbaidzhana za 40 let; 1920-1960 gg. Baku, Azzhanskoe gos. izd-vo neftianoi i nauchno-tekhn. lit-ry 133 p. (Azerbaijan, Industriae)	hlen- erbaid- , 1960.		
	(wserig: JanIndustries)			,
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ALIKHANOV, E.N.; ARUSHANOV, N.A.; AKHUNDOV, V.Yu.: ALIZADE, M.A.; AZIZBEKOV, Sh.A.; BAGIROV, M.A.; VEZIROV, S.A.; VOLOBUYEV, V.R.; VEKILOV, F.M.; GADZHIYEV, N.M.; GUSEYNOV, D.M.; GUSEYNOV, I.A.; DADASHEV, K.K.; DADASHZADE, M.A.; DALIN, M.A.; ISKENDEROV, M.A.; KAZIYEV, M.A.; KARAYEV, A.I.; KASHKAY, M.S.; KEL'DYSH, M.V.; KERIMOV, A.G.; LEMBERANSKIY, A.F.; MAMEDOV, G.K.; MEKHTIYEV, M.R.; MIRZOYEV, S.A.; NAGIYEV, M.F.; NASRULLAYEV, N.I.; OGUDZHEV, A.K.; RADZHABOV, R.A.; RUDNEV, K.N.; SAECKHOV, R.N.; SEMENOV, N.N.; TOPCHIYEV, A.V.; TOPCHIBASHEV, M.A.; TAIROVA, T.A.; KHALILOV, Z.I.; EFENDIYEV, G.Kh.; SHUKYUROVA, Z.Z.

ALIKHANOV, E.N.; MIRCHINK, Y.F., red.; AKHMEDOV, G.A., red.

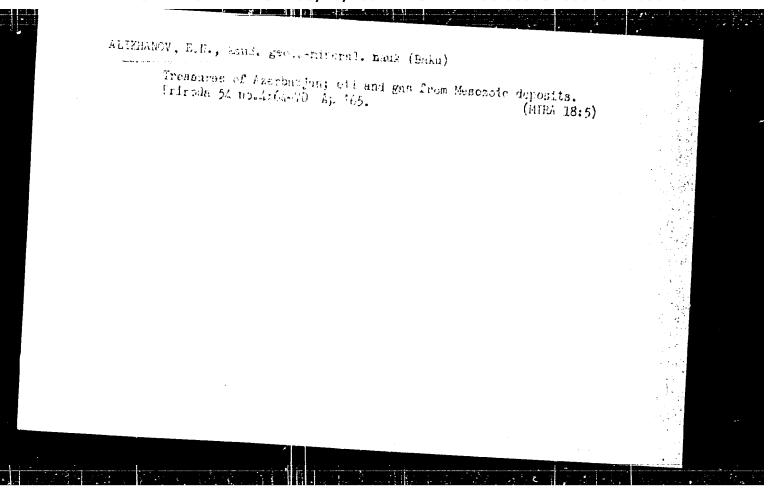
[Oll and gas fields of the Caspian Sea] Neftiarye i gazovye mestorozhdenia Kaspiiskogo moria. Baku, Azerneshr, 1964. 382 p. (MIRA 17:10)

ALIKHANOV, E.N.; ASAN-NURI, A.O.; KULIYEV, I.P.; MAMEDOV, B.M.;

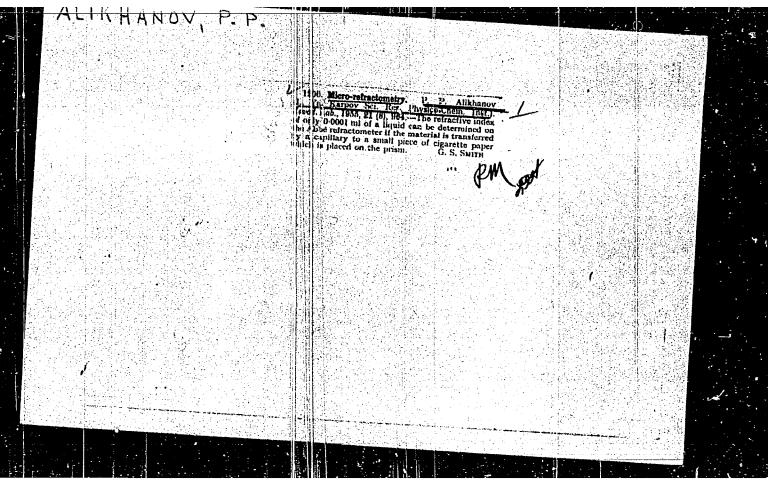
ORDZHEV, S.A.; TIMDIEYEV, N.S.

Off-shore oil of the U.S.S.R. Neft. khoz. 42 no.9/10:
46-51 S-0 '64.

(MIRA 17:12)



"APPROVED FOR RELEASE: 03/20/2001 CIA-RDP86-00513R000101110007-0



ALIKHANOY

USSR/Organic Chemistry. Theoretical and General Questions of Organic Chemistry.

RUE

E-1

Abs Jour

: Ref Zhur - Khimiya, No. 8, 1957, 26613.

Author

Kololev, A.; Shatenshteyn, A.; Yurygina,

Inst

Ye.; Kalinachenko, V., Alikhanov, P.

Title

Isomerization of Monodeuteronaphthalenes.

Orig Pub

Zh. obshch. khimii, 1956, 26, No. 6, 1666 -

Abstract

The question of the possibility of transposition of X-H and B-H in the naphthalene molecule was investigated by the method of deuterium interchange. It is shown that if vapors of X-deuteronaphthalene, as well of B-deuteronaphthalene (I and II) in a flow of nitrogen are passing above silica gel at 4200,

Card 1/3

USSR/Organic Chemistry. Theoretical and General Questions of Organic Chemistry.

E-1

Abs Jour

Ref Zhur - Khimiya, No. 8, 1957, 26613.

transposition of D and the formation of π mixture consisting of approximately equal amounts of the two isomers will be observed. Dueterium was not transposed in the nainthalene molecule at a heating of I and II at 40(10 for 52 hours in absence of silica gel. The method of investigation is based on the difference in speeds of hydrogen interchange between I or II with liquid Hir. The speed constants C were determined basing on experiments of interchange of I and II with HBr at 250: C in sec-1 is 2 to 3 x 10-3 for I and 5 x 10-5 for II. It is shown at the same time, that a-methylnaphthalene, as well as &-methylnaphthalene (III and IV) are converting mutually one into the

Card 2/3

AUTHORS:

Shatenshtern, A. I., Vedeneyev, A. V., Alikhanov. P. P.

SOV/79-28-10-3/60

TITLE:

Hydrogen Reaction of Phenol, Its Ethers and of the Aromatic Amines With Liquid DBr (Vodorodnyy obmen fenola, yego efirov i aromaticheskikh aminov s zhidkim DBr)

PERIODICAL:

Zhurnal obshchey khimii, 1958, Vol 28, Nr 10, pp 2638 - 2644 (USSR)

ABSTRACT:

Shatenshteyn and his collaborators had earlier found the rules governing the deutero reaction in hydrocarbons dissolved in liquid DBr (Refs 1,2). In this paper the results are given which were obtained in the hydrogen contain with liquid DBr in aromatic compounds that The free electron pairs of oxygen or nitrogen atom. Substituents are in mesomeric relation to the \pi-electrons of the aromatic nuclei, which fact causes an increase of the electron density in the ortho and para-atoms (Ref 4). The affiliation of the proton (deuteron) to the electron pair of the nitrogen or oxygen atom causes its transition to the quaternary or ternary ion

Card 1/3

Hydrogen Reaction of Phenol, Its Ethers and of the Aromatic Amines With Liquid DBr

507/79-28-10-3/60

with simple positive charge. These characteristic features of the compounds mentioned above are the decisive characteristics in their deutero reaction with acids. The hydrogen reaction on phenol and its ethers $(c_6H_5OCH_3, c_6H_5OC_6H_5)$ and on aromatic amines $(c_6H_5N(CH_3)_2,$ (C6H5)2NH, (C6H5)3N with liquid DBr as well as with DBr+AlBr, were investigated at 25°. In all compounds of the first group the ortho and para-atoms react immediately whereas in the second group this rapid reaction takes place only with $(c_6\bar{H}_5)_3\bar{N}$, with all others only slowly or not at all. AlBr3 causes the reaction of the meta-atoms in the phenol ethers and delays the reaction in $(C_6H_5)_3N$. The different behaviour of compounds containing oxygen and nitrogen in the hydrogen reaction with DBr+AlBr, depends on their different relation to the proton and on the different coordination capability of oxygen and nitrogen atoms.

Card 2/3

Hydrogen Reaction of Phenol, Its Ethers and of the Aromatic Amines With Liquid DBr

SOV/79-28-10-3/60

There are 3 tables and 17 references, 10 of which are Soviet.

ASSOCIATION:

Fiziko-khimicheskiy institut imeni L.Ya.Karpova (Physical Chemical Institute imeni L.Ya.Karpov)

SUBMITTED:

August 20, 1957

Card 3/3

66428

5(4) 5.2400(H), 5.3200

SOV/20-128-6-33/63

AUTHORS:

Alikhanov, P. P., Varshavskiy, Ya. M.

TITLE:

Equilibrium Distribution of Deuterium on Hydrogen Exchange With

Liquid Deuterium Iodide

PERIODICAL:

Doklady Akademii nauk BSSR, 1959, Vol 128, Nr 6, pp 1214-1216

(USSR)

ABSTRACT:

This paper belongs to a series of investigations of the isotope equilibria in various systems of hydrogenous compounds (Refs 1-4). An investigation is made of the exchange reaction between the aromatic C-H bond and the deuterium-containing HJ. Benzene, which would be well suited for this investigation between

would be well suited for this investigation because of the equivalence of all H atom, reacts—too slowly so that diphenyl had to be used. In the dissolution of diphenyl in liquid hydrogen iodide the 6 H-atoms in ortho- and para-position are easily exchanged while the 4 H-atoms in meta-position virtually did not react. The deuterium concentration in HJ was determined by analysis of the water obtained by decomposition of HJ in the nitrogen current over CuO at 350 - 400. The reaction with diphenyl took place in sealed glass tubes at a pressure of 4 - 15 atm. After equilibration had been reached, the HJ was evaporated, the diphenyl in the oxygen current was burnt, and

Card 1/2

66428

SOV/20-128-6-33/63 Equilibrium Distribution of Deuterium on Hydrogen Exchange With Liquid Deuterium Iodide

> the deuterium content of the water obtained was determined. The resulting distribution coefficient a was checked by the counterreaction of I-substituted diphenyl in ortho- and para-position with HJ. The experimental data on c are compared in the table with the values calculated by means of the formulas contained in reference 8. The values coincide well. As is shown in figure 1, good agreement was also obtained for the other hydrogen halides (Refs 2-4). In the present paper the following values are given for the equilibrium coefficient α : $\alpha_{00} = 4.07$,

 α_{250} = 3.47 and α_{500} = 3.13. There are 1 figure, 1 table, and 9 Soviet references.

ASSOCIATION: Nauchno-issledovatel skiy fizil o-khimicheskiy is stitut im. L. Ya. Karpova (Scientific Research Institute of Physical Chemistry imeni L. Ya. Karpov)

PRESENTED:

June 19, 1959, by S. S. Medvedev, Academician

SUBMITTED:

April 13, 1959

Card 2/2

SHATENSHTEYN, A.I.; SANNIKOV, A.P.; ALIKHANOV, I.P.

Deuterium exchange method for studying the catalytic activity of systems consisting of hydrogen acid and eprotic acid-like substance. Acetic acid stannichloride-system. Zhur. ob. khim. 35 no.3:419-425 Mr 165. (MIRA 18:4)

1. Fiziko-khimicherkiy institut imeni L Ia. Karpova.

5.2000 78298 SOV/79-30-3-52/69 AUTHORS: Shatenshteyn, A. I., Alikharov, P. P. TITLE: Concerning Catalytic Action of Iodine in the Deuterium Exchange in Liquid Hydroge: Iodide PERIODICAL: Zhurnal obshchey khimii, 1960, Vol 30, Nr 3, pp 992-999 (USSR) ABSTRACT: Rate of deuterium exchange of monodeuteriotoluene (I) and monodeuteriobiphenyl (II) with HI was studied in the presence of iodine as a catalyst. HI was synthesized in a quartz apparatus described previously (V. R. Kalinachenko and others, ZhFKh, 30, 1140, 1956). A test tube containing a thin-walled glass ampoule filled with the investigated compound and catalyst (iodine) was attached to the apparatus to fill it with the HI obtained (-45°) . After filling with HI, the test tube was detached from the apparatus and kept at $25 \pm 0.2^{\circ}$. The glass ampoule inside the test tube Card 1/8 was imploded by the HI vapor, and the exchange

Concerning Catalytic Action of Iodine in the Deuterium Exchange in Lichid Hydrogen Iodide

78298 **SOV**/79-30-3-52/69

reaction started. The reagents used were prepared by Ye. N. Yurygina. Results of the experiments are shown in Table 1. Absorption spectra of iodine in liquid HI were taken at room temperature according to a previously described method (A. I. Shatenshteyn, Ye. A. Izriilevich, ZhFKh, 26, 377, 1952). SF-4 spectrophotometer and a quartz cell (0.11 cm) were used. Specific electric conductivity of liquid HI

was measured at -44° and found to be equal to $2 \cdot 10^{-8}$ shm⁻¹cm⁻¹. The measurements were taken in a cell with nonplatinized platinum electrodes using an alternate current bridge constructed under the supervision of V. Ye. Kazakevich. There was no noticeable increase in the conductivity of the solution after the addition of iodine, mesitylene, or hexamethylbenzene. Cataly it action of iodine is due to the polarization of the bond H-I in the complex formed in the ternary system: promatic hydrocarbon-hydrogen iodide-ioding:

Card 2/8

Concerning Catalytic Action of Todine in the Deute-ium Exchange in Liquid Hydrogen Iod1de

73008 SOV/79-30-3-52/69

 $ArD \oplus HJ \oplus J_{\pi_2} \cdot ArD \oplus \dots \oplus H \oplus J_{\pi}$

Comparison of the rates of deuterium exchange is given in Table 2. The authors express their gratitude to Ye. N. Yurygina for the deuterium-containing hydro-carbons. There are 2 figures; 3 tables; and 45 references, 13 U.S., 7 U.K., 1 German, 24 Soviet. The 5 most recent U.S. and U.K. references are: G. Olah, S. Kuhn, A. Puvlath, Nature, 178, 693 (1956); E. L. Mackor, F. J. Smit, J. H. van der Waals, Trans. Farad. Soc., 53, 1309 (1957); V. Gold, D. P. Satchell, J. Chem. Soc., 1927, 3904, 3910 (1958); H. H. Hyman, M. Kilpatrick, J. J. Katz, J. Am. Chem. Soc., 79, 3668 (1957); W. G. Schneider, "Hydrogen Bonding," ed. Hadzi, Pergamon Press, N. Y., 54 (1959). L. Ya. Karpov Scientific Research Institute of Physi-

ASSOCIATION:

cal Chemistry (Nauchno-issledovatel'skiy fiziko-

khimicheskiy institut imeni L. Ya. Karpova)

SUBMITTED:

April 10, 1959

Card 3/8

78298 SOV/79-30-3-52/69

Table 1. Results of experiments on deuterium exchange with liquid HT_{\ast}

14	C _{Arly} · 10 ^a	CJ, 10*	ို့	d	e 1	k - 104	<i>h</i> ' · 1₁2	h C
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1	1.4	•	10.80	5.86	1 45	4.0		
2	1.4	•	10.80	5.90	45	4.0		_
3	1.7	-	5.50	3.05	43	4.0		
4	1.7	-	5.50	3.02	43	4.0		
5	1.6	0.6	5,60	3,43	28	5.2	8.7	270
6	1.9	1.2	5,60	2,86	22	9.2	7.7	160
7	1.8	2.0	5,60	3.16	14	12	6.0	90
8	1.8	3.5	5,50	2.88	8	24	6.9	51
9	1.7	5.0	5,50	2.70		43	7.3	29
0	1.9	6.9	5.60	3,23	3	55	8.0	28
1 2	1.8	10	5,60	3,02	2	92	9.1	18
- 1	2.2	12	5,60	2,94	1.5	130	10.8	18

Card 4/8

(Table 1 cont'd on Card 5/8)

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-	Mr	a	b	С	d	78. <u>e</u>	$\frac{1}{ r }$	<u> </u>	1 h	<u>52/69</u>	
	13 14 15 16 17 18 19 20 21	1.6 1.6 1.9 1.6 1.9 1.6 1.5 1.8	1.7 2.0 2.0 2.0 5.8 11 14 25	5.50 5.50 5.50 5.50 5.50 5.50 5.50 5.50	3,05 2,96 3,09 3,32 2,43 2,76 2,68 3,48 2,21	64 50 30 23 24 12 5 3	2.1 4.0 6.4 7.1 12 19 47 61 160	3.6 2.9 2.7 3.1 3.3 4.3 4.4 6.4	150 86 62 49 26 14 13 7		
	22 23 24 25 26	0,75 0,63 0,67 0,86 0,71	0.8 2.1 3.8 12	8,10 8,10 8,10 8,10 8,10	k 4.27 3.97 4.44 4.93 5.08	45 28 12 6 1.5	4,2 8,6 15 25 91	10.8 7.2 6.6 7.0	79 32 23 6		
Card 5/8	27 28 29 30 31 32 33	0.71 0.82 0.75 1.1 0.24 0.83 0.82	1.8 3.4 5.2 7.8 11 12 17 (Key	7.80 7.80 7.80 7.80 7.80 7.80 7.80 7.80	1 4.76 5.49 4.27 4.76 5.42 5.40 5.01 ble 1	45 23 25 12 6 5 4	3.5 5.6 7.9 14 20 23 36	2.0 1.7 1.5 1.8 1.8 1.9 2.1	39 24 14 14 2 7 5		
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Concerning Catalytic Action of Iodine in the Deuterium Exchange in Liquid Hydrogen Iodide 2. 45

78298 SOV/79-30-3-52/69

Key: (a) ${\rm C_{APD}} \cdot 10^2$ is the number of hydrocarbon moles per mole of HI; (b) ${\rm C_{J2}}$ is the iodine concentration in moles per mole of HI; (c) ${\rm C_B^O}$ is the deuterium concentration in water after the precipitation of the compound before the experiment (in atom \$\mathcal{H}\$); of the compound before the experiment; (e) \$\mathcal{T}\$ is (d) ${\rm C_{B}}$, after the end of the experiment; (f) k is constant duration of the experiment in hours; (f) k is constant of the rate of deuterium exchange; (g) k' = k/C_{J_2} is the specific rate constant in reference to the concentration of iodine in the solution; (h) C = C_{ArD}/C_{J_2}; (i) p-D-toluene; (j) o-D-toluene; (k) p-D-biphenyl; (1) o-D-biphenyl.

card 6/8

Concerning Catalytic Action of Toding in the Deuterium Exchange in Liquid Hydrogen Todine

Fig. 2. Absorption spectra of iodine solutions; (1) in CCl_h; (2) in C₆H₆; (3) in liquid HI.

Concerning Catalytic Action of Todine in the Deuterium Exchange in Liquid Hydrogen Todide 78298 **SOV/**79-30-3-52**/**69

Table 2. Comparison of the constants of deuterium exchange rate with different hydrogen halides at 25° (sec $^{-1}$).

O THE STREET AND A STREET	HI	11Br	HI,
e	2 · 10 · 6 4 · 50 · 6 1 · 10 · 6 4 · 10 · 6 (5 · 10 · 5)	5 - 10 5 2 - 10 4 3 - 10 5 1 - 10 3 23 - 10 9	> 2 \cdot 10^2

Key: (a) Hydrocarbon; (b) o-D-toluene; (c) p-D-toluene; (d) o-D-biphenyl; (e) p-D-biphenyl; (f) α -D-naph-thalene.

Card 8/8

AUTHORS:

Yurygina, Ye. N., Alikhanov, P. P., S/076/60/60/12railevich, Ye. L., Manochkina, P. N., B115/B016 8/076/60/034/03/015/038

Shatenshteyn, A. I. (Moscow)

TITLE:

The Kinetics of Dauterium Exchange of the Isomers of Monodeuterotoluene, Monodeutero-diphenyl, and Monodeutero-naphthalene With Liquid Hydrogen Bromide and a Solution of Potassium Amide in

Liquid Ammonia

PERIODICAL:

Zhurnal fizicheskoy khimii, 1960, Vol 34, Nr 3, pp 587 - 593

(USSR)

TEXT: The aim of the investigation under review was the determination of the factors of the partial rate f in the hydrogen isotopic exchange of the substances mentioned in the title with the reagents likewise mentioned in the title. The synthesis of monodeuterated hydrocarbons, the deuterium concentration in water on combustion of the hydrocarbons, and the carrying out of experiments are described. The rate constant of the deuterium exchange is calculated by an equation and, when using ammoniacal solutions, by a simplified form of this equation. The results of the measurements made with liquid HBr are given in table 1, those of the experiments with ammoniacal solutions in table 2, and the mean values of the constants of the deuterium exchange rate

Card 1/3

The Kinetics of Deuterium Exchange of the Isomers of S/076/60/034/03/015/038 Monodeutero-toluene, Monodeutero-diphenyl, and Mono- B115/B016 deutero-naphthalene With Liquid Hydrogen Bromide and a Solution of Potassium Amide in Liquid Ammonia

in table 3. The activation energy of deuterium exchange is also given. Table 4 presents the factors of the partial rate f of the deuterium exchange of isomeric monodeuterated hydrocarbons with a ${\rm KNH_2}$ solution in ${\rm NH_3}$, and with HBr.

The order of the partial rate factors in the isotopic exchange between non-equivalent deuterium atoms in the toluene and diphenyl molecule differs in reactions with acids and with lyes, which is due to the different mechanism of these reactions in which the conjugative and the inductive effect considerably manifest themselves in the mutual action of the atoms in the hydrocarbon molecule. The rules in the deuterium exchange in toluene and diphenyl indicate that the inductive effect of the methyl group has the reverse sign compared to that of the phenyl group. Finally, it can be said that the acid and the base react in the deuterium exchange with the carbon atoms of the CH-bonds and the base with the hydrogen atoms of the CH-bonds, and protonize these atoms. A method is described for obtaining isomeric monodeutero-diphenyls. It is described exactly how the authors divided this work among themselves. There are 4 tables and 19 references, 12 of which are Soviet.

Card 2/3

The Kinetics of Deuterium Exchange of the Isomers of Monodeutero-toluene, Monodeutero-diphenyl, and Mono-deutero-naphthalene With Liquid Hydrogen Bromide and a Solution of Potassium Amide in Liquid Ammonia

ASSOCIATION: Fiziko-khimicheskiy institut im. L. Ya. Karpova (Institute of Physical Chemistry imeni L. Ya. Karpov)

SUBMITTED: June 13, 1958

Card 3/3

ALIKHANOV, P. P.

Cand Chem Sci - (diss) "Investigation of reactions of deutero-exchange of some aromatic compounds with liquid hydrogen halides."

Moscow, 1961. 15 pp; (Ministry of Higher and Secondary Specialist Education RSFSR, Moscow Order of Lenin Chemical-Technological Inst imeni D. I. Mendeleyev); 150 coples; price not given; (KL, 5-61 sup, 175)

SHATENSHTEYN, A.I., prof.; VYRSKIY, Yu.P., kand. khim. nauk; PRAVIKOVA, N.A., kand. tekhn. nauk; ALIKHANOV, P.P., kand. khim. nauk; ZEDANOVA, K.I., kand. khim. nauk; IZYUMNIKOV, A.L., mlad. nauchn. sotr.; LEVINSKIY, Yu.V., red.

[Practical Maboratory manual on the determination of the molecular weights and molecular weight distribution of polymers] Prakticheskoe rukovodstvo po opredeleniu molekuliarnykh vescy i molekuliarno-vesovogo raspredelenia polimerov. [By] A.I.Shatenshtein i dr. Moskva, Khimiia, 1964. 188 p. (MIRA 18:2)

USSR/Physics - Heat of evaporation of 02

FD-3283

Author

: Alikhanov, R. A.

Title

Heat of evaporation of oxygen in the interval of temperatures 80-106°K

Periodical: Zhur. eksp. i teor. fiz., 29, No 6(12), Dec 1955, 902-903

Abstract

The author made 35 measurements of the heat of evaporation of \mathcal{O}_2 for 7 different temperatures, obtaining values different from those of other researchers; e.g. he obtained L (cal/mole) = 1628.7±1.3 at ToK = 90.19. He gives the other values of L for 6 other temperatures and the empirical expression relating L to T; e.g.

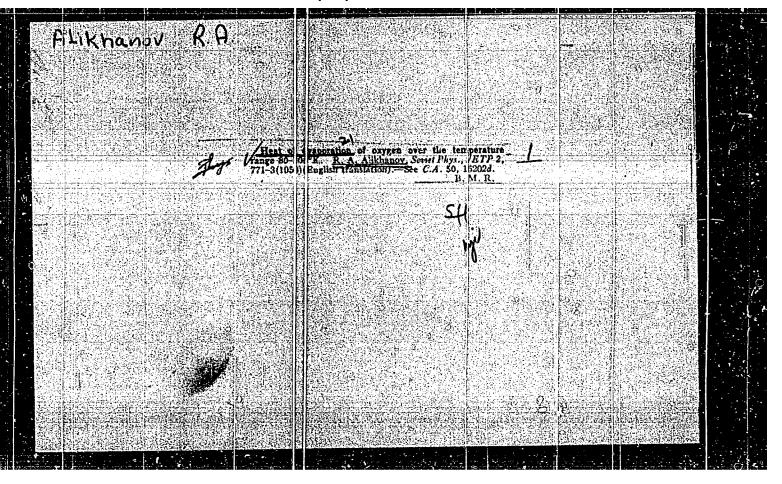
 $L^2 = 67208.68(T_{cr}-T) - 555.221(T_{cr}-T)^2 + 2.32505(T_{cr}-T)^3$

The author thanks P. G. Strelkov for his attention and interest. Five

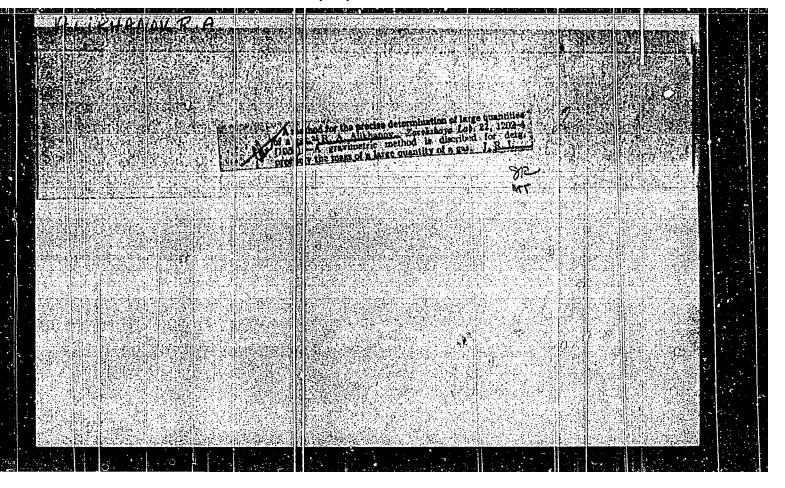
Institution: Institute of Physical Froblems, Academy of Sciences USSR

Submitted : July 21, 1955

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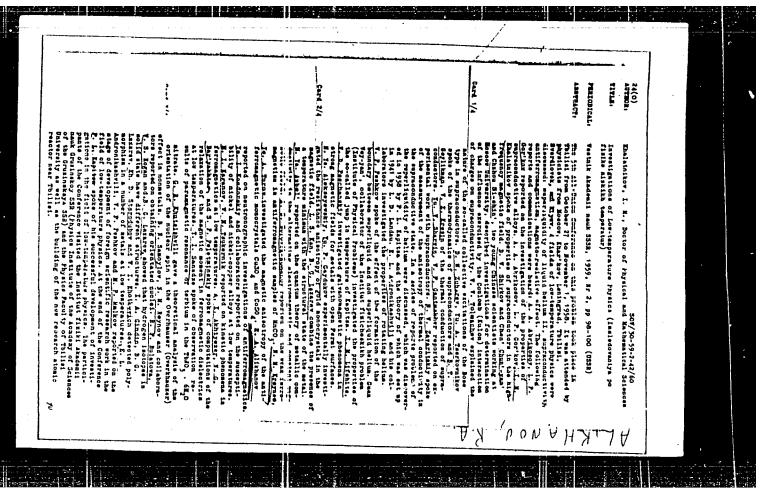


ALIKHAROV, R. A., Cand Phys-Eath Sci — (diss) "Moutronographic study of antiferropagnetics knoo3, Feco3 and NiF21."

Los, 1959. 10 pp (Acad Sci USSR. Inst of Physica Problems).

150 copies. Bibliography at end of text (10 titles)

(KL, 39-59,100)



24(3) SCV/56-36-6-10/66 AUTHOR: Alikhanov, R. A. TITLE: Neutronographic Investigation of the Antiferromagnetism of the Carbonates of Mn and Fe (Neytronograficheskoye issledovaniye antiferromagnetizma karbonatov Mn i Fe) PERIODICAL: Zhurnal eksperimental'noy i teoreticheskoy fiziki, 1959, Vol 36, Nr 6, pp 1690 - 1696 (USSR) Borovik-Romanov and Orlova (Ref 1) already inventigated the ABSTRACT: magnetic properties of polycrystalline $\mathrm{MnCO_3}$ and $\mathrm{CoCO_3}$, and I. Ye. Dzyalcshinskiy (Ref 2) published a theory on this subject. Dzyaloshinskiy pointed out the connection between weak ferromagnetism and crystal symmetry: In Ma-, Fe-, and Co-carbonates 3 forms of magnetic symmetry are distinguished: 1) the spins are in the direction of the rhombohedral axis, 2) the spins are in the basis plane and are aligned along the axis of the second order, 3) the spins are in the basis plane and in the symmetry plane. In the first-mentioned case no weak ferromagnetism can be observed, whereas in the two Card 1/3 latter cases such a weak ferromagnetism is found to exist. In

Neutronographic Investigation of the Antiferromagnetism SOV/56-36-6-10/66 of the Carbonates of Mn and Fe

the present paper the author investigates MnCO_3 and FeCO_3 by means of the neutron diagram method with respect to Dzyaloshinskiy's theory. A crystal powder was exposed to a monochromatic neutron beam (λ = 1.37 A), and the scattered neutrons were recorded by a spectrometer constructed by Yu. G. Abov (Ref 5). The neutrons originated from the heavy water reactor of the ITEF AN SSSR. Figure 1 shows the neutron diffraction pictures obtained of MnCO_3 powder at room temperature and

at hydrogen temperature. The latter is between the background curve and that recorded at room temperature, and has two additional maxima before the maximum of the room temperature neutron diffraction picture. The lattice constants were determined as amounting to a=5.84~Å and $\alpha=47^{0.15}$; further data are given by table 1. The structure of FeCO₃ was investigated

in a siderite sample found at Bakal (USSR). Figure 2 shows the neutron diffraction picture at hydrogen temperature. The nuclear scattering amplitudes of Fe⁺⁺ are double those of Mn⁺⁺. Table 2 contains the calculated and measured relative

Card 2/3

Neutronographic Investigation of the Antiferromagnetism SOV/56-36-6-10/66 of the Carbonates of Mn and Fe

in MnCO₃ the spins in the antiferromagnetic state are in the basis plane and symmetry plane; in FeCO₃ the spins are aligned along the rhombohedral axis. According to Dzyaloshinskiy this would mean that MnCO₃ has a weak ferromagnetism, whereas with FeCO₃ this is not the case. The author finally thanks Academician P. L. Kapitsa for his interest in this investigation, A. S. Borovik-Romanov for his valuable advice, I. Ye. Dzyaloshinskiy for discussions, Yu. G. Abov for his advice in connection with the neutronoscopic part of the work, and Yu. V. Sharvin for placing a special thermometer at his disposal. There are 6 figures, 2 tables, and 11 references, 5 of which are Soviet.

intensities for the first group of peaks. It was found that

ASOCIATION:

Institut fizicheskikh problem Akademii nauk SSSR (Institute for Physical Problems of the Academy of Sciences, USSR) February 11. 1959

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CIA-RDP86-00513R000101110007-0

S/056/60/C38/O3/15/O33 B006/B014

AUTHOR:

Alikhanov, R. A.

TITLE:

A Cryostat for Neutronographic Investigations //

PERIODICAL:

Zhurnal eksperimental'noy i teoreticheskoy fiziki, 1960,

Vol. 38, No. 3, pp. 806-808

TEXT: The present paper described a low-temperature (including the temperature of liquid helium) cryostat employed for investigating the scattering of thermal neutrons on polycrystalline samples. This apparatus, which offers the possibility of investigating neutron diffraction within a wide range of Bragg angles (up to about $O_{\rm B}$ = 45°, corresponding to a rotation of the de-

tector through 90°), is described in the introduction. Inside, it has a Dewar designed by P. L. Kapitas (cf. Fig. 1). As was shown by a number of neutronographic investigations, this apparatus operates satisfactorily. The background due to the aluminum casing of the samples amounts to 20% of the counter background and does not disturb the neutron diffraction picture. In this instrument samples can be kept at the temperature of liquid helium for

Card 1/2

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A Cryostat for Neutronographic Investigations

S/056/60/038/03/15/033 B006/B014

48 hours. The amount of helium required is 7.5 liters. Fig. 2 shows a typical evaporation curve of helium from the cryostat (1.31). A thermometer supplied by Yu. V. Sharvin and exhibiting a resistance which strongly depends on temperature in the low-temperature range was used for measuring the temperature of the samples. Fig. 3 shows the calibration curve of this thermometer, which is 1.2.3.5 mm large. In some cases, sample temperatures were also checked during the experiment, for which purpose a recording potential eter of the type EPP-09 was used. The author finally thanks Academician P. L. Kapitsa for his interest and Yu. V. Sharvin for the thermometer supplied. There are 3 figures and 5 references, 4 of which are Soviet.

ASSOCIATION:

Institut fizicheskikh problem Akademii nauk SSSR (Institute of

Physical Problems of the Academy of Sciences, USSR)

SUBMITTED:

October 10, 1959

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1144, 1395, 1160, 2209

s/056/60/039/005/048/051 B006/B077

AUTHOR:

Alikhanov R. A.

TITLE:

The Antiferromagnetism of CoCO,

PERIODICAL:

Zhurnal eksperimental'noy i teoreticheskoy fiziki; 1960,

 $Vol_{*} 39$, $Nc_{*} 5(11)$, $pp_{*} 1481 - 1483$

TEXT: The author reports on the investigation of hydrothermically produced finely crystalline CoCO₃ samples with the help of neutron reflec-

tion experiments at low temperatures. He received these samples from I. Yu. Ikornikova of the Institut kristallografii AN SSSR (Institute of Crystallography AS USSR). Fig. 1 shows a neutron reflection diagram taken at 4.20K. The position of the magnetic moments was computed from these results. The direction of antiferromagnetism with regard to the rhombic axis was determined from the magnetic intensity ratios and it was found that the moments and the [111] axis enclose an angle of (46.4). The magnetic reflections (111) and (100) (see diagram) and also (110) and (211) showed a decrease in intensity on transition from 20.4 to 4.20K.

Card 1/3

86929

The Antiferromagnetism of CoCO,

5/056/60/039/005/048/051 B006/B07?

The results are briefly discussed applying the thermodynamic theory by I. Ye. Dayaloshinskiy. Fig.2 illustrates the position of the magnetic moment in CoCO, as determined experimentally. Among other things it was

found that the regnetic moments of the ions fall out of the symmetry plane by the angle γ; an estimate of this angle resulted in $(15\pm5)^\circ$, and a calculation of this angle using the data from absolute measurements (Ref.8) gave $\gamma = 7^\circ$. The author thanks Academician P. L. Kapitsa for his interest, A. S. Borovik-Romanov and I. Ye. Dzyaloshinskiy for discussions, and N. Yu. Ikornikov and N. N. Mikhaylov for preparing and sorting the samples and I. P. Karpikhin for his experimental assistance. S. D. Chetverikov, P. V. Kalinin, M. G. Spiridonova, B. M. Shzakin, M. P. Orlova, and V. I. Ozhogin are mentioned. There are 2 figures and 8 references: 4 Soviet and 4 US.

ASSOCIATION: Institut fizicheskikh problem Akademii nauk SSSR (Insti-

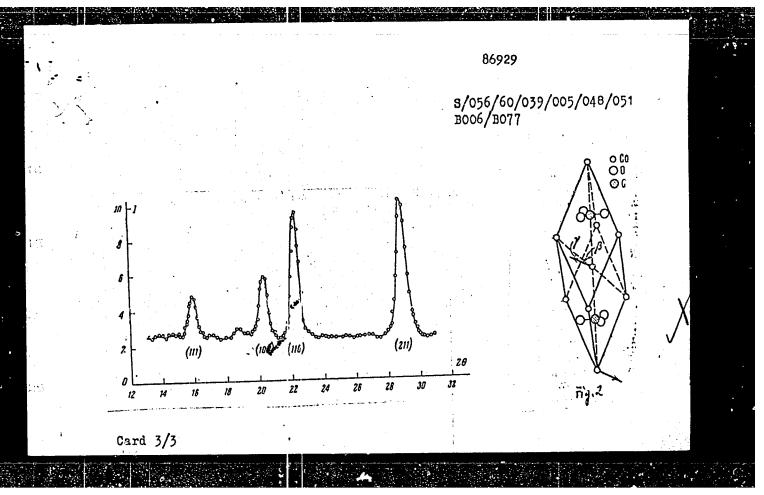
tute of Physical Problems of the Academy of Sciences

USSR)

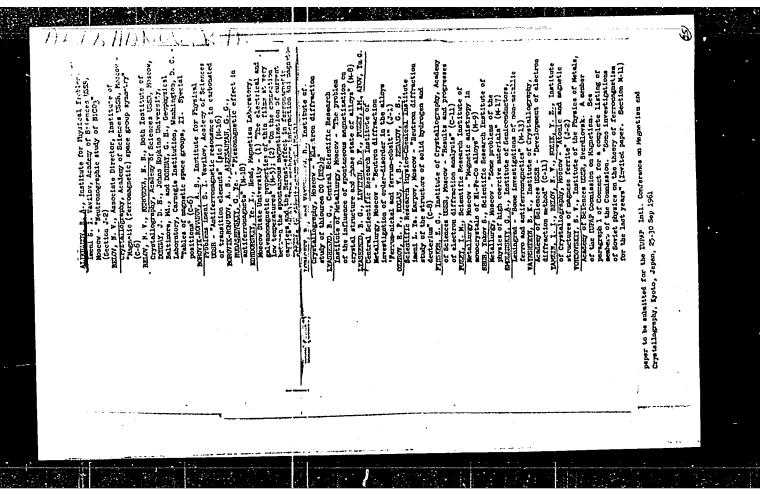
SUBMITTED: August 26, 1960

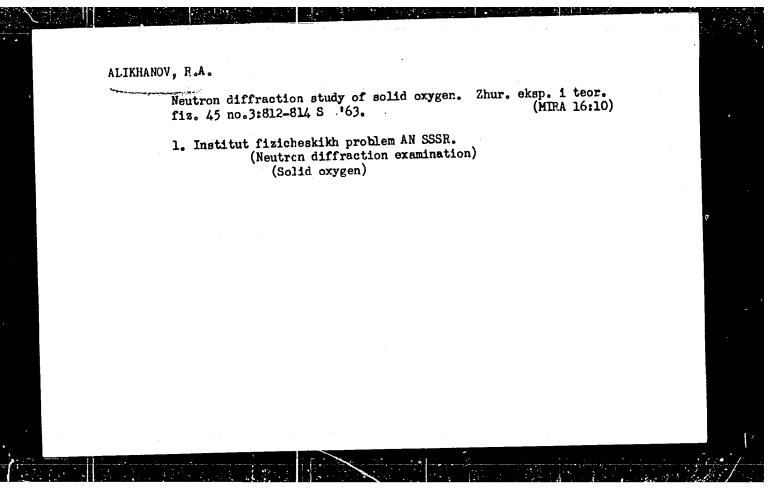
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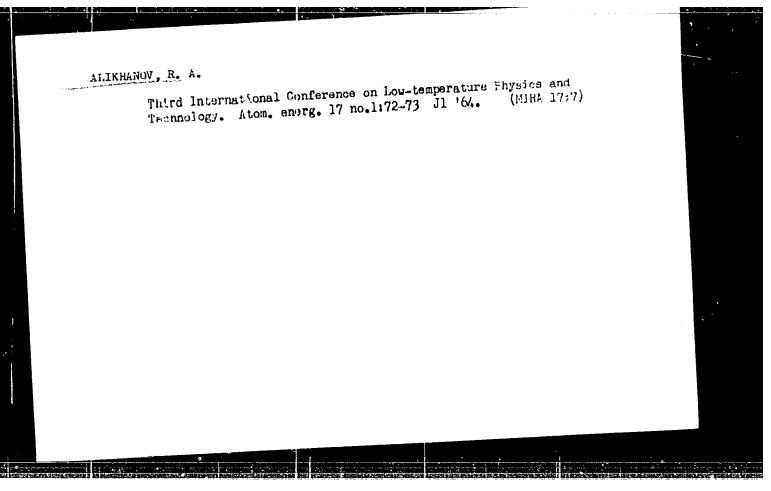
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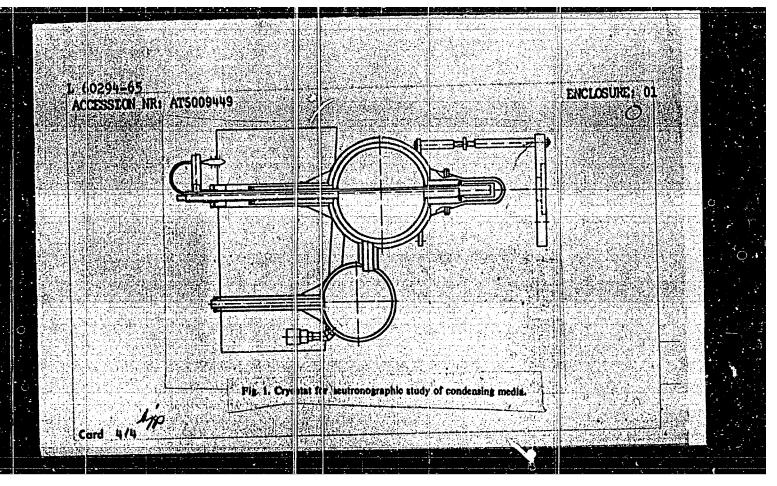
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1-602/1-65 EPF(a)/EPF(n)-2/EWG(d)/EEG(k)-2/EWA(h)/EWT(d)/EWT(1)/EWT(E)/T/ EWP(b)/EWP(t): Pf-4/Pg-4/Pu-4/Pe): IJP(c)/Z/0000/64/000/000/0127/0131 ACCESSION NR1 /AT5009449 RPL - W/JW/JD - 55	
AUTHOR: Alikhanov, R. 19 6+1 TITLE: Neutronographic investigations of solid oxygen 31	
SOURCE: Conference on Low Temps rature Physics and Techniques. 3d, Prague, 1963. Physics and techniques of low temperatures; proceedings Prague, 1963. Physics and techniques of low temperatures; proceedings Of the conference. Prague, Fubl. House of the Ozedhosl. Academy of Sciences, 1964, 127-131 TOPIC TAGS: neutron diffraction, solid oxygen, crystal structure, phase transition, low temperature research ABSTRACT: Neutron-diffraction studies of solid oxygen were made with an aim at assessing the applicability of neutron-diffraction techni- an aim at assessing the applicability of neutron-diffraction techni- ques to measurements of crystalline and magnetic properties of oxygen ques to measurements of crystalline and magnetic properties of oxygen at low temperatures. Some of the earlier studies are reviewed and the difficulties of the research are mentioned. The technique used the difficulties of the research are mentioned. The technique used was described by the author earlier (ZhETF v. 36 (1959) 1690), with	

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entailed the use also described by	method used of of a Kapitsa-I (2	working with condensed gas ewar, the application of wh hETF v. 38, 806, 1960). The Enclousre. An important	es. This nick was ne pryostat festure of	
the new cryostate came temperature of the sample as The measurements with the bath fipositions of the rhombohedral lat	as the bath, is sumes in statle of the neutror lied with liquiobtained diffitice with parar	s used to ensure that the fashion the temperature of diffraction from oxygen we describe action lines are described eters a = 4.210 and l = 46	temperature I the bath. Ere performed Im. The I well by a If Dif- similar	
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AUTHOR: Alikhanov. R. A.

ORG: none

SURCE: USSR. Gosudarstvennyy komitet po ispol'zovaniyu atomnoy energii.
Institut teoreticheskcy i eksperimental'noy fiziki. Doklady, no. 355, 1965.

Neuprugoye rasseyaniye neytronov gematitom, 1-20

TOPIC TAGS: neutron scattering, inelastic scattering, inelastic neutron scattering, hematite neutron scattering, tricrystal spectrometer, hematite spin wave, magnon moment

ABSTRACT: A three-crystal spectrometer was used to investigate the spectrum of spin in hematite (& Fe₂O₃) originating from the Shabra deposits. The experiment was conducted over the 0.64A⁰-2 to 5.3⁻³ ev range of transmitted magnon moment and energy. Two branches with energy gaps characterized by magnetic anisotropy were observed. Values of (1.3 + 0.7)10⁻³ ev and (0.4 + 0.15)10⁻³ ev were observed for the gap, and (1.3 + 0.7)10⁻⁶ cm/sec and (4.6 + 0.5)10⁵ cm/sec

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for spin wave velocity for t Kapitsa and A. I. Alikhanov	he branches.	The author	thanks A	cademicia	ns P. I.		
Kapitsa and A. I. Alikhanov Dzyaloshinskiy and L. K. F.	for their con	tinuous atteni	tion to th	is work,	Ye.		
Dzyaloshinskiy and L. K. F thanks Professor G. P. Bar	ar Kovskiy for	their helpful	l discuss	ions. The	e author		3
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AT6033191

SOURCE CODE: UR/3138/65/000/393/0001/0015

31

AUTHOR: Alikhanov, R. A., Smirrov, L. S.

ORG: none

TITLE: Neutronographic investigation of the low-temperature magnetic transition

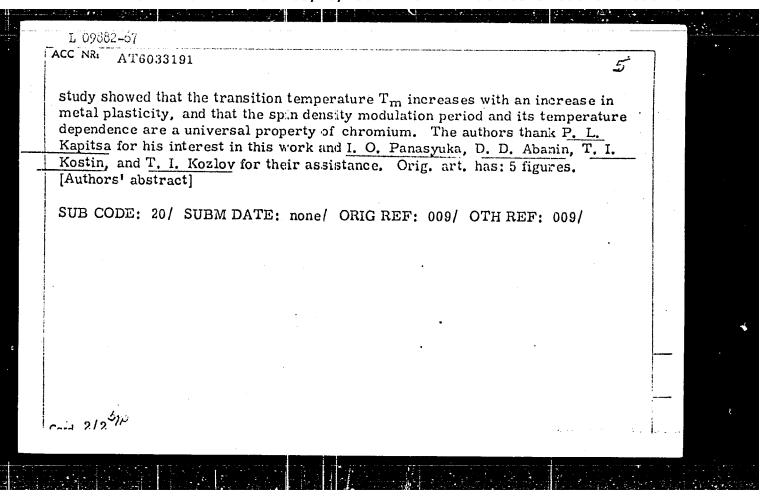
in chromium

SOURCE: USSR. Gosudarstvennyy komitet po ispol'zovaniyu atomnoy energii. Institut teoreticheskoy eksperimental'noy fiziki. Doklady, no. 393, 1965. Netronograficheskiye issledovaniya nizkotemperaturnogo magnitnogo prevrashcheniya v khrome, 1-15

TOPIC TAGS: chromium, cryostat, magnetic transition, low temperature. magnetic transition, crystal spin structure, chromium crystal, neutron scattering, magnetic scattering

ABSTRACT: A neutronographic study was made of the low-temperature magnetic transition in samples of crystalline chromium and polycrystalline chromium containing equal amounts of impurities but subjected to different thermomechanical treatment. A miniature cryostat (weight 530 g, height 200 mm) was used. The

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ALIKHANOV, S.G.

SUBJECT

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20

AUTHOR TITLE ALICHANOV, S.G.

A New Impulse Method for Measuring Ion Masses. Zurn.eksp.i teor.fis, 31, fasc. 3, 517-518 (1956)

Issued: 12 / 1956

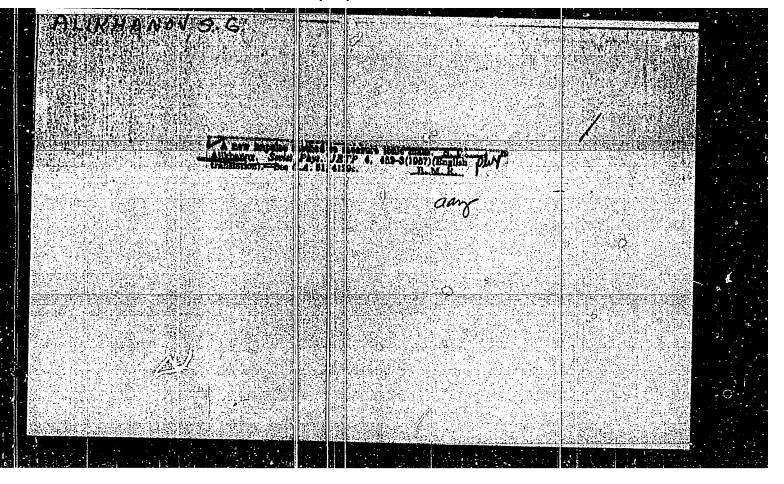
By the focussing method suggested here it is possible to increase the resolving power of the spectrometer considerably. The mass spectrometer consists of a drive tube which is limited on both sides by slowing-down fields with linear potential distribution. The whole device is in a weak longitudinal field. The condensed ions produced in an impulse-like source are "injected" into the tube, and at the same time the slowing-down field is switched on. The condensed ions inciding into the potential well begin to move from one reflector to the other, on which occasion the time gained by the fastest ions with given e/M in comparison with slow ions is compensated by the loss of velocity during motion in the slowing-down field. The ions with equal e/h and different energies then gather at a certain point in the drive tube. After a sufficient number of cycles, when the ions with approximately equal e/M separate, the shift of the time of arrival in the focus exceeds the time needed for condensation. Hereby the latter are deflected and annihilated. In order that the condensations of ions with different masses oscillate in the tube, it is necessary to connect a voltage on to the deflecting plates which is switched on only on the occasion of the passing of the ions with the masses to be measured. For the duration T of a complete cycle

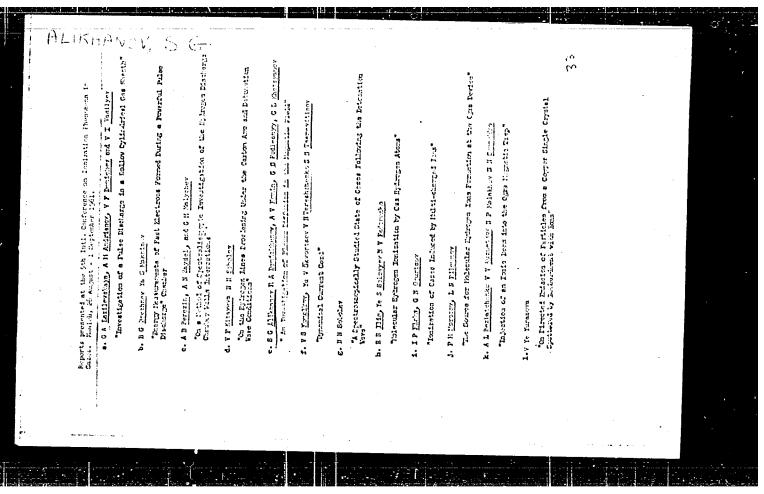
Žurn.eksp.i teor.fis, 31, fasc.3,517-518 (1956) CARD 2 / 2 PA - 1656 $T = 1\sqrt{2}$ M/UZe + $2((1/E_1)+(1/E_2))\sqrt{2}$ is found with the slowing-down fields E_1 and E_2 . Here M denotes the mass, Ze = the charge, and U = the energy (in volts) of the ions, and l = the length of the drive tube. The condition $\partial T/\partial U$ results in $(1/E_1) + (1/E_2) = 1/2$ U₀. If the suitably selected quantities U₀, 1, E₁ and E₂ satisfy the latter equation, a grouping in time and in space (focussing) of the first order of ions of different energies is attained. However, the focussing condition of the second order is not satisfied and the washing out in time of the cycle for ions with the same mass is

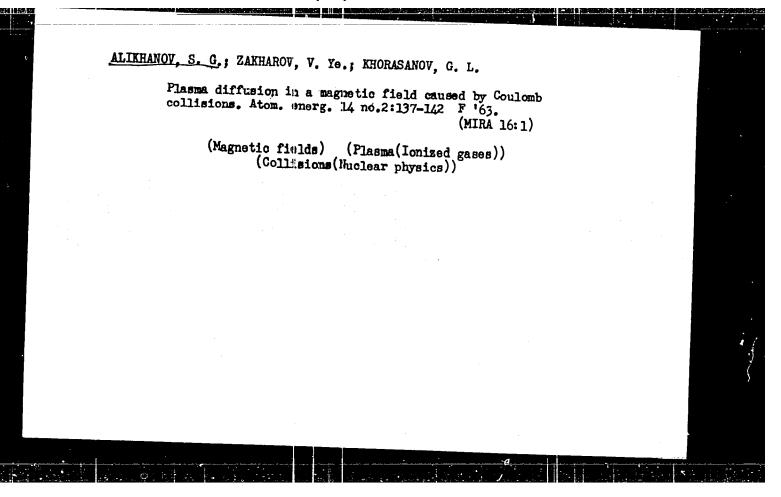
 \triangle T_U ~ $(1/2)\sqrt{M/2ZeU^5}$ $(\triangle$ U)². The limiting value of the resolving power is \triangle M/M = $(1/4)(\triangle$ U/U)². The accuracy of measurements depends on the duration and the linear dimensions of ion condensation. The ratio (duration of ion condensation / time needed for passage) must be smaller than the limiting value of the resolving power. In order to improve accuracy it is necessary either to increase the length of the tube or the ions must perform the necessary number of cycles. The field strength of the longitudinal magnetic field is selected in such a manner that the resolving power of the device is not diminished by an increase of light intensity.

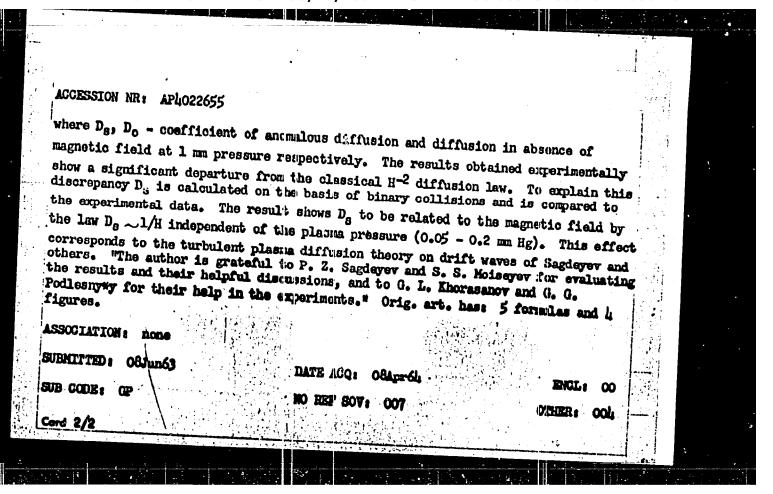
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ACCRSSION NRI APBO07307	8/0057/63/035/001/0557/0551
AUTHOR: Alikhanov, S.G. Ivaniya	63
TIME: Temperature and density di	ribut on in a hot stationary are
SCURCE; Zhurnal tekhnicheskoy fizil	, v.3t, no.3, 1965, 557-561
TOPIC 1ACS: plasma plasma arc, plasma arc, plasma	ma corfinement, hot plasma confinement, deu-
ABSTRACT: In order to explore one of A.Alfven and E.Smars (Nature, 188 insulated from a surrounding dense tions for an axially symmetric composally. The results are presented gray insulating magnetic field war as in a galeous shell). Both electron equations, the electron and ion tem loss by radiation was neglected. The T5/p, where T and p are the temperature of the composal of th	If the possibilities inherent in the suggestion 801,1960) that a hot plasma might be thermally asse by a magnetic field, the steady state equationized deuterium are were so wed numeriphically and are discussed briefly. The thermal- umed to be that due to the arc current (pinch heat conduction were included in the gratures were assumed to be equal, and energy steady state equations involve the parameter ure and pressure on the axis of the arc. Solu-
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of somewhat similar calculations of Whereas Falthammar finds the pinch plasma, the present calculations sho layer at the wall of the confining not represent a solution of his equal to some concluded that the solution of transport coefficients provides no might be achieved without employing one may expect to be able to isolational wall of the chamber. "In conclusion and support of the work. N.K. Fare for	is parameter ranging from 1017 to 1014 (0.05 calculations are not in agreement with those C.G.Fs thammar (Phys.Fluids 4,1145,1961). o be surrounded by a thick dense shell of cold within the density rises sharply only in a thin essel. The curve in Falthammar's figure 6 does tions (19)-(21), and is accordingly erroneous. the steady state equations with the classical asis for hope that a very hot stable plasma for the special means for its stabilization, but that the key plasma from impurities coming from the the suthers thank G.L.Budkek for his interest a valuable consultations, and L.P.Kovachevich lons. Orig.art.has: 23 formulas, 8 ligures	
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ALIKHANOVA, KH. M.

Category: USSR/ Farm Animal Diseases Caused by Bacteria and Fungi V-2

Abs Jour: Refer. Zhur-Biologiya, No 16, 1957, 72308

Author : Gryslov V. P. Alikhanova Kh. M.

: Not given Title

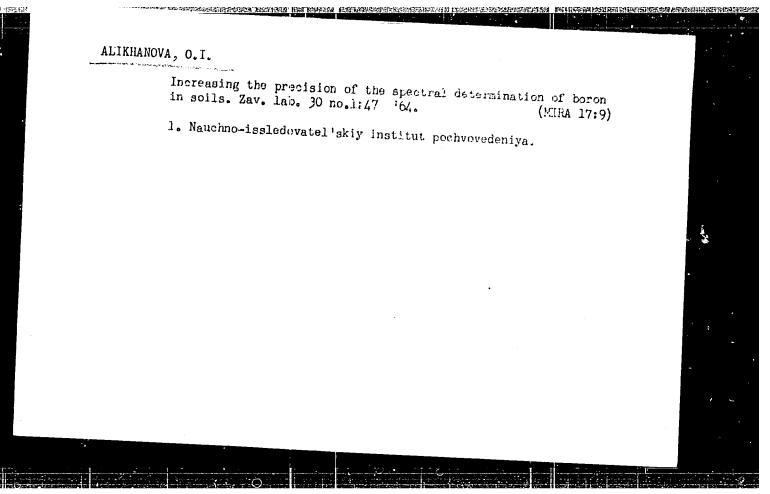
: A Case of Actinomycosis of the Breat-Bone of a Cow

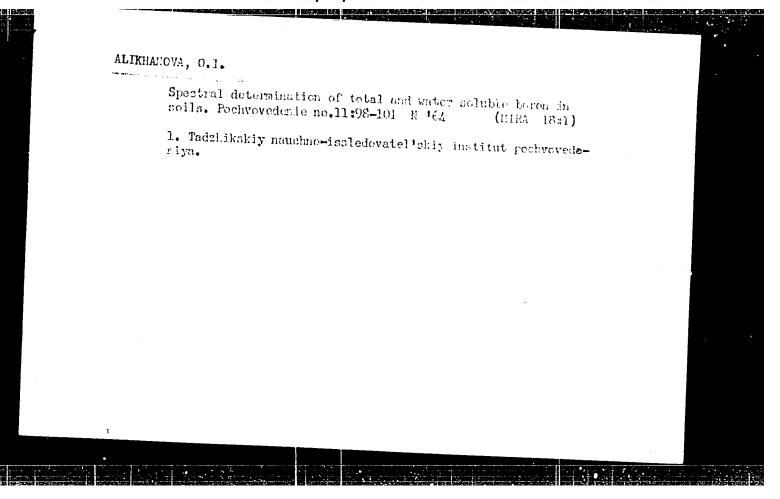
Orig Pub: Tr. Dagestansk. S.-Kh. In-ta, 1956, 8, 193-194

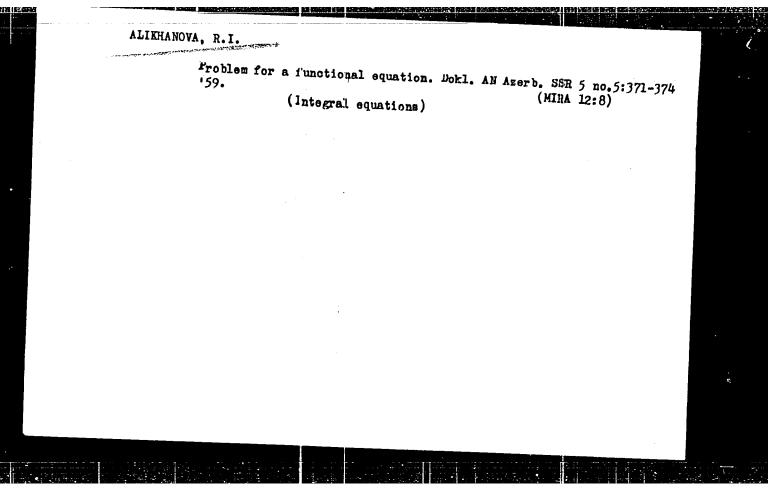
In the region of 6-7 segments of the breast-bone and 7-9 rib carti-Abstract: lage a painless swelling was found, which had a pus secreting fistula in the center. The study of of the pus demonstrated the presence of actinomyces. On the basis of the findings, upon opening the authors suspect that the actinomycosis localized in the breast bone occurred due to a trauma (piercing of the bone) by a wire covered with Actinomyces, which penetrated into the breast region.

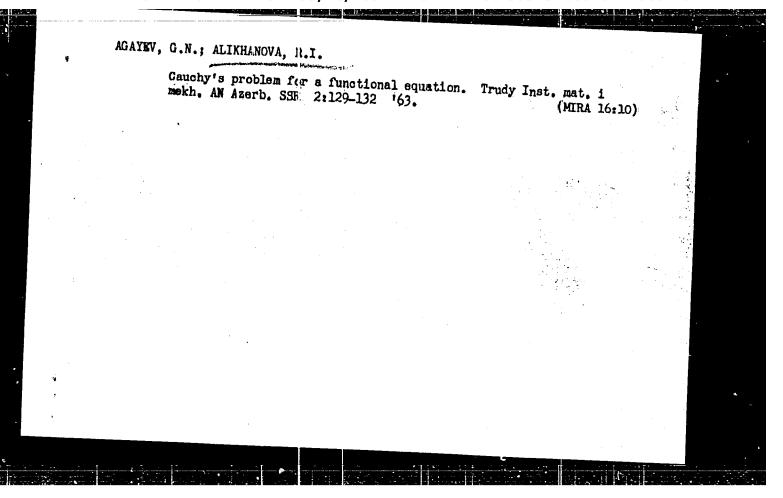
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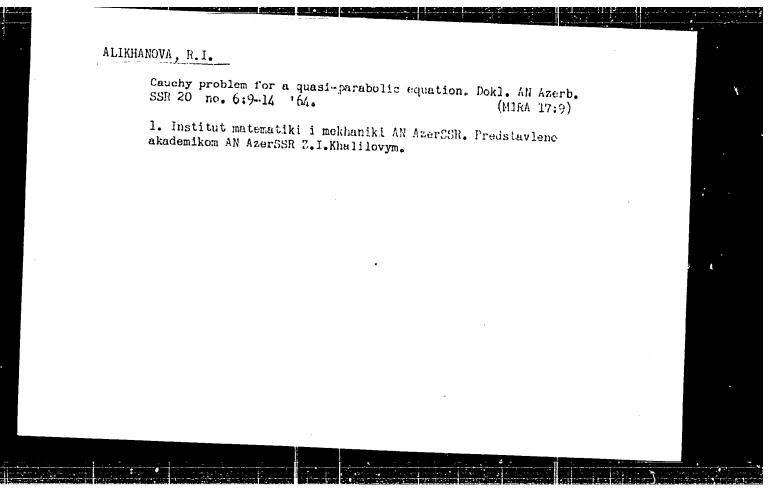
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ACCESSION NR: AP4045055 S/0249/64/020/006/0000/0014 AUTHOR: Alikhanova, R. I., Khalilov, Z. I. -TITLE: The Cauchy problem for a quasi-parabolic equation SOURCE: AN AzerbSSR. Boklady*, v. 20, no. 6, 1964, 9-14 TOPIC TAGS: Cauchy problem, parabolic equation, quasiparabolic equation ABSTRACT: In continuation of earlier work (Agayev G. N., Alikhanova R. I. Trudy* Inta matematiki i mekhaniki AN Azerb. SSR, vol. 10, 1963), the author considers the Cauchy problem for the functional integrodifferential equation (1) with the inital condition (2) $|u(t,x)|_{1=0}=u_0(x).$ where $\mathbf{R}_{\mathbf{n}}$ is an n-dimensional Euclidean space. It is assumed that (3) Card 1/4

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where Co>O is a defi	nite number, independent of t	and z. The form	•		
	$\sum_{z} \varphi^{m}(z) A_{m}(t) \rho^{m}$	•	(4)		
in the class of function order > 2r-1, and re	tonously in z with fixed t and a ons vanishing near infinity, tog duces to the solution of a dual	real vector p. The sol	ution is sought ives of the		
area that if	$v(t, p) = \int e^{t(p,z)} u(t,x) dx,$	problem. It can readily	be demonstr-		
and v(t, p) is the solut	ion of the dual anablem		(5)	Ž	
with the initial conditi	$\frac{dv(t,p)}{dt} = \sum_{ \mathbf{m} \leq 2t} v^{\mathbf{m}} [\int v^{\mathbf{k}}(t,p)dp].$	$A_m(t)\rho^{-n}(t,p)$	(6)		
	$v(t,\rho) _{t\to v_{\bullet}(\rho)}$	•	(7)		
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rd ^{2/4}				• •	
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then u(t,x) is the formal solution of the problem represented by (1) and (2). Assuming that

$$0 < v_{\bullet}(\rho) < \frac{N}{[1+|\rho|^{2}]^{r+\frac{n+1}{3}}},$$
 (8)

where N O is a definite number, it is clear that the integrals

$$\int [1+|\rho|^{1}]^{2r+\frac{n+1}{2}} v_{0}^{2}(\rho) d\rho \tag{9}$$

$$\int v_0^2(\rho)d\rho \tag{10}$$

converge. This is proven by means of four preliminary lemmas. The author then goes on to prove that the problem represented by (6) and (7) has a solution, arrived at by the methods of successive approximation and mathematical induction. In final form, the required solution of the problem represented by (1) and (2) is shown to be

$$u(t,x) = \left(\frac{1}{2\pi}\right)^n \int_{0}^{\infty} V(t,p) \dot{e}^{-i(p,x)} dp. \tag{11}$$

Card 3/4

and

APPROVED FOR RELEASE: 03/20/2001

CIA-RDP86-00513R000101110007-0"

ACCESSION NR: AP4045055

"The author expresses his appreciation to G. N. Agayev for suggesting the problem and helping in the daily work." Orig. art. has: 12 numbered equations.

ASSOCIATION: Institut matematiki i mekhaniki Akademii nauk Azerbaydzhanskoy SSR (Institute of Mathematics and Mechanics, Academy of Sciences of the Azerbaidjan SSR)

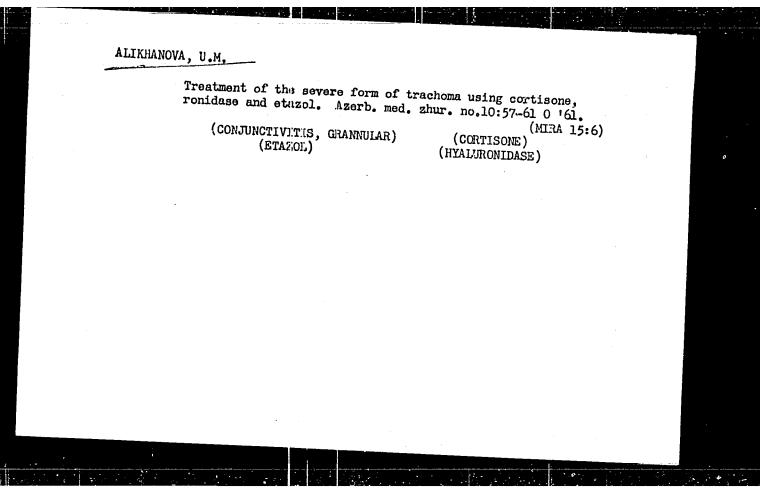
SUBMITTED: 02Jul63

ENCL: 00

SUB CODE: MA

NO REF SOV: 003

OTHER: 000



S/078/61/006/005/004/015 B121/B208

AUTHORS:

Markov, V. P., Alikhanova, Z.M.

TITLE:

Uranyl compounds with trihydroxy-glutaric acid

PERIODICAL: Zhurnal neorganicheskoy khimii, v. 6, no. 5, 1961, 1066 = 1073

TEXT: The complex formation of the uranyl ion with trihydroxy-glutaric acid COOH - CHOH - CHOH - CHOH - COOH was studied by physico-chemical methods such as potentiometric titration, determination of the specific electrical conductivity and transference number. The potentiometric titrations were made on the JNI-5 (LP-5) potentiometer by means of a glass-and a calomel electrode. The titrations were carried out with a 0,1 M allutions and trihydroxy-glutaric acid with different ratios of the components at 20°C. No hydroxide precipitates in the titration of a mixture of uranyl nitrate and trihydroxy-glutaric acid with alkali hydroxide solution. It may be seen from the potentiometric titrations and measurements of the specific electrical conductivity of isomolar solutions that tri-

Uranyl compounds with ...

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hydroxy-glutaric acid reacts with uranyl ions by forming complexes in which the ratio of the components U: C5 H807 varies from 1: 3 to 2: 1. Various uranyl compounds with trihydroxy-glutaric acid were synthesized. UO2C5H6O7.1 1/2 H2O is obtained by two methods. In the first method trihydroxy-glutaric acid is added to a nearly saturated uranyl nitrate solution up to a ratio of the components of 1: 2. After 2 - 3 days a fine--crystalline, light-yellow precipitate is obtained. This method is also applicable when using uranvl chloride, but not, when using uranyl sulfate. In the second method, trihydroxy-glutaric acid is added to an aqueous suspension of uranyl oxide in the heat up to a ratio of the components U: C5H8O7 = 1: 3. The analytical results obtained for the crystals prepared by the two methods are the same. The crystals of $00_2^{\circ}5^{\circ}6^{\circ}7^{\circ}1$ 1/2 $^{\circ}1_2^{\circ}$ 0 are insoluble in water and organic solvents. The compound (UO2)2C5H4O7.4H2O is precipitated from a solution of uranyl nitrate and trihydroxy-glutaric acid, where the ratio of the components is 2: 1, by a mixture of ethyl alcohol (2 parts) and ether (5 parts). Chemical ana-Card 2/7

Uranyl compounds with ...

S/078/61/006/005/004/015 B121/B208

lysis confirmed the given formula. The compound is soluble in water, but insoluble in organic solvents, it is decomposed in aqueous solution by lyes or acids. The compound is a nonelectrolyte, as was confirmed by measuring the molar electrical conductivity. The compound (UO2)2C5H6O7(HSO4)2.H2O

is precipitated by treating a solution of uranyl sulfate and trihydroxy-glutaric acid with alcohol and ether, The chemical analysis of the compound confirmed the above composition. The compound is very well soluble in water, but insoluble in organic solvents. The compound

NH₄(UO₂)₂C₅H₃O₇·3H₂O is precipitated from an uranyl nitrate solution with trihydroxy-glutaric acid at a pH of about 2 after 2 - 3 hr in the form of a fine-crystalline, light-yellow powder. The analysis of the compound confirmed the above formula. This compound is insoluble in water and organic solvents, but soluble in alkalies. The alkaline solutions of this compound may be destroyed by mineral acids. $K(UO_2)_2C_5H_3O_7 \cdot 3H_2O$ and $Ba[(UO_2)_2C_5H_3O_7]_2 \cdot 12H_2O$ are obtained by reaction of

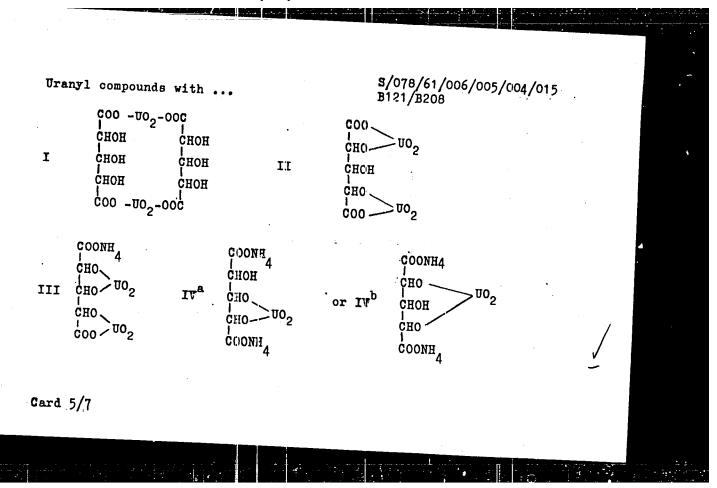
potassium- and barium hydroxide, respectively, with solutions containing uranyl nitrate and trihydroxy-glutaric acid. The above formulas are con-

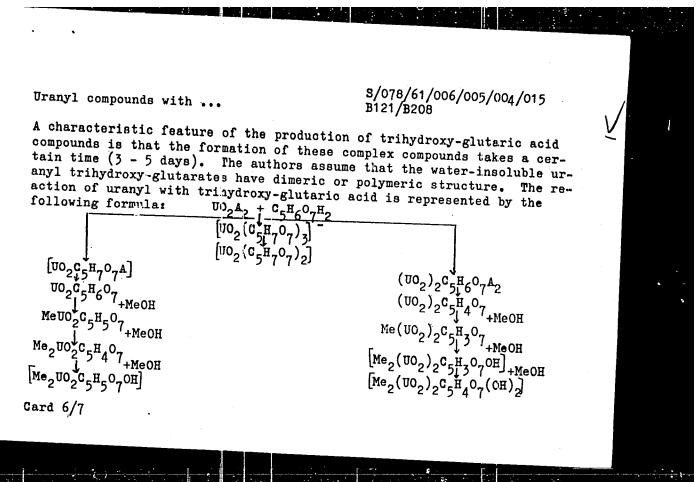
Uranyl compounds with ...

S/078/61/006/005/004/015 B121/B208

firmed by chemical analyses. Compounds of the Me₂UO₂C₅H₄O₇·nH₂O type (Me = K, Na, or NH₄) are obtained by two methods: In the first method, uranyl nitrate is added to a solution of trihydroxy-glutaric acid and alkali lye, in which the ratio of the components C₅H₈O₇: KOH = 2:3 or 2:4. The solution having a pH of 9 - 10 is then treated with ethyl alcohol and ether. The following compounds were isolated: UO₂K₂C₅H₄O₇·4H₂O and UO₂Na₂C₅H₄O₇·4H₂O. In the second method, uranyl trihydroxy-glutarate is precipitated from saturated solutions of uranyl nitrate and trihydroxy-glutaric acid, and is then dissolved in dilute ammonium hydroxide. The glutaric acid, and is then dissolved in dilute ammonium hydroxide. The yellow crystalline (NH₂)₂UO₂C₅H₄O₇·2H₂O is precipitated from the ammoniacal solution by alcohol and ether. The chemical analysis confirmed this composition. The compound is soluble in water, but insoluble in organic solvents, and is decomposed by mineral acids. For the resultant compounds the following structural formulas were suggested on the basis of the experimental

Card 4/7





Uranyl compounds with ...

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The spectrophotometric studies were carried out by Syuy Li-yuan' (Ref. 1: Dissertation at the Institut geokhimii i analiticheskoy khimii im. V. I. Vernadskogo, 1960 (Institute of Geochemistry and Analytical Chemistry imeni V. I. Vernadskiy, 1960). There are 5 figures and 10 Soviet-bloc

SUBMITTED:

December 1, 1960

Card 7/7

L5507	6-65 EWT(m)/EWP(1)/T LOW NR: AP5017999	Pg=1 Zd4			
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MARKOV, V.P.; KHARITONOV, Yr., Ya.; ALIKHANOVA, Z.M.

Structure of complex compounds of uranyl with anoins of tartaric, malic, and trihydroxyglutaric acids. Zhur.neorg.khim. 8 no.3:774-775 Mr 163.

(MIRA 16:4)

1. Institut obshcher in neorganichegkoy khimii imeni N.S. kurnakova AN SSSR.

(Uranyl compounds)

(Acids, Organic)

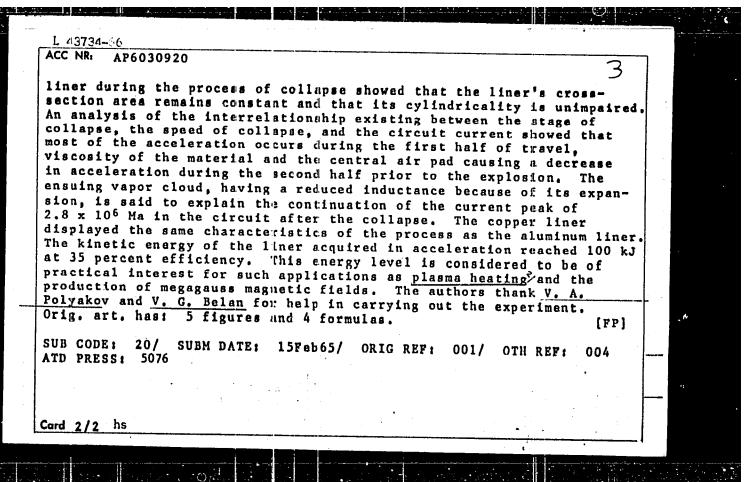
GOLOVNYA, V.A., doktor khim. nauk; ELLERT, G.V., kand. khim. nauk; SHUBOCHKIN, L.K., kand. khim. nauk; SHCHELOKOV. R.N., kand. khim. nauk; TSAPKINA, I.V., kand. khim. nauk; RARGGEMY, Ye.N., kand. khim. nauk; Markov, V.P., doktor khim. nau, [deceased];

ALKHANOVA, Z.N., FYARKINA, M.Ye., doktor khim. nauk; KIRHAYLOV, Yu.N.; TSAFKIN, V.V., kand.khim. nauk; BOLOTOVA, G.T., kand.khim. nauk; CHERNYAYEV, V.A., doktor khim.nauk; KORCHENNYA, Ye.K., red.

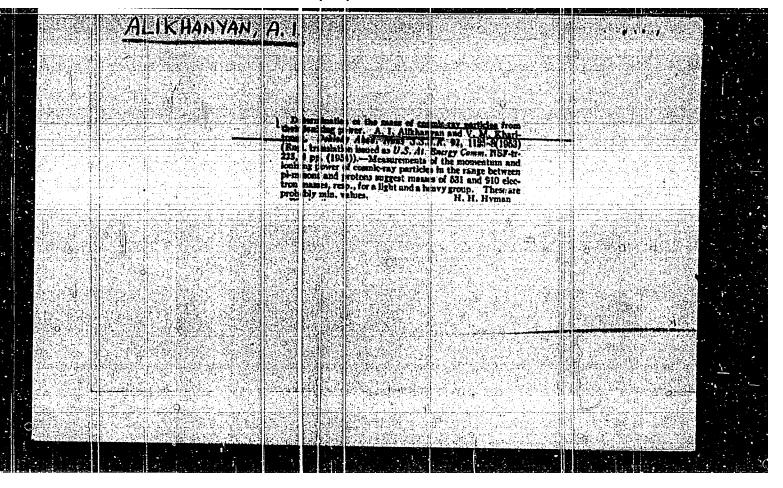
[Complex compounds of uranium] Kompleksnye soedineniia urana. Moskva, Izd-vo "Nauka," 1964. 488 p. (MIRA 17:7)

1.Akademiya nauk SSSR. Institut obshe'ey i neorganicheskoy khimii. 2. Laboratoriya khimii kompleksnykh soyedineniy aktinidov Instituta o'shchey i neorganicheskoy khimii AN SSSR (for all except Korchemmaya).

L 43734-66 E.YT(1' IJP(c) ACC NR AP6030920 SOURCE CODE: UR/0207/66/000/004/0038/0041 AUTHOR: Alikhanov, S. G. (Novosibirsk); Budker, G. I. (Novosibirsk); Kichigin, G. N. (Novosibirsk); Komin, A. V. (Novosibirsk) 50 ORG: none 47 ۶, ٧, Implosion of a metal liner by the action of a magnetic field TITLE: SOURCE: Zhurnal prikladnoy mekhaniki i tekhnicheskoy fiziki, no. 4, TOPIC TAGS: implosion, metal liner implosion, magnetic implosion, plasma heating, megagauss field, megagauss magnetic field, STRUCTURE DYNAMIC STABILITY, STRUCTURE STABILITY ABSTRACT: The experimental results of theta-pinch of metal liners by the action of a magnetic field of a single-turn solenoid are presented and compared with theoretical data on the collapse mechanics of liners. The charge of a 5 x 10^{-2} f condenser at a voltage of 4 kv was used to activate the solenoid. AD-1M (aluminum) and M-1 (copper) cylindrical liners 80 mm in outside diameter and 150 mm long were used. Wall thickness was 2.5 mm with the aluminum liner and 1 mm with the copper liner; weight was 250 g and 350 g, respectively. The circuit current and battery voltage were recorded along with other data during the experiment. A series of photographs taken from the end projection of the Card 1/2



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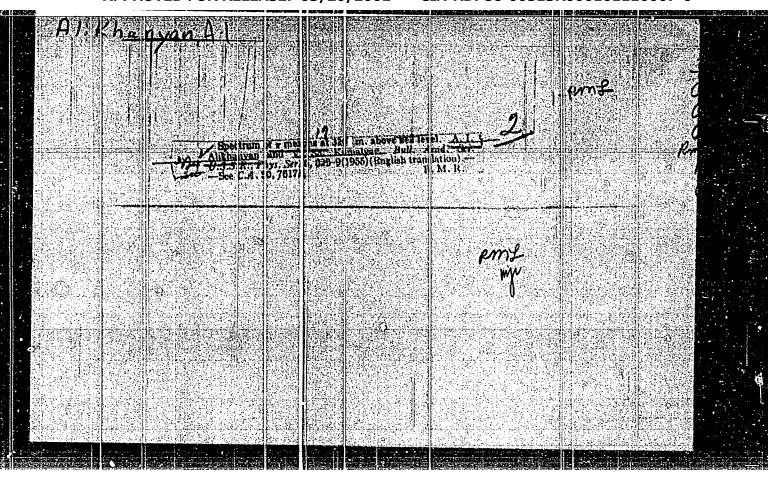


WSR/Nuclear Physics - Radioactivity Pub. 124 - 8/29 Cleared 1/1 Alikhanyan, A. I., Memb. Corresp. of Acad. of Sc. USER.; and Vaysenberg, Arithors Artificial radiolotivity Title 1 Vest. AN SSSR (, 51-61, June 1954 Periodical : Speeches held in commemoration of the 20th anniversary of the discovery, Atistract by Irene and Frederic Julio-Curie, of artificial radioactivity are presented. Various stages in the development of nuclear physics, beginning with the discovery by Marie Curie of two radioactive elements Po and Ra (1897-1898), the discovery of neutron radioactivity by Fermi and associates and including developments up to 1953, were mentioned. The direct relation between artificial radioactivity and various cosmogonic problems is explained. The speakers also predicted that by 1970 the total amount of radioactive fission products obtained from reactors will reach 100 tons per year which will correspond to a radioactive radiation energy of 12 million lw. Institution Submitted

ALIKHANYAN USSR/Physics	
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Authorn ;	Kamalyan, V. and Alikhanyan, Memb. Corres. of the Acad. of 5cs. of the
Title :	USSR About the spectrum of X-mesons at an altitude of 3200 m
Periodical :	Dokl. AN SSSR, 97, 14. 3, 425 - 428, July, 1954
Abstract :	Description is given of experiments performed at an altitude of 3200 M above sea level for the purpose of obtaining a spectrum of A -mesons formed by collisions of neutrons of cosmic radiation with the solar nuclei. The experiments were conducted with the help of a magnetic spectrometer with an additional hodoscope. Seven references. Diagrams.
Institution :	Physical Institute of the Acad. of Scs. of the Arm. SSR.
Submitted :	
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	도로바다는 이 시간 전략을 하고 있다. 그런 사람들은 사람들은 사람들은 기가 가는 시간 가는 것이 되었다. 그런 그를 보고 있다. 1985년 - 1987년 - 1985년 - 1987년 -
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ALIKHANYAN, A-1. USSE/ Physics - T-particles Card 1/1 Pub. 22 - 6/40 : Alikhonyan, A. I., member correspondent of the Acad. of Sds. of USSR;
Dayon, M.I.; Sliostakovich, N. V.; Kirillov-Ugoyumov, V. G. and Deryagin, B.N.;
Unstable charged particles heavier than protons. Authors Title Periodical : Dok. AN SSSR 99/3, 361-364, Nov 21, 1954 Abstract : Four cases of charges particles heavier than protons, observed in Wilson's camera, are described. These particles were designated T-particles and their mass, sign, durations and energy were estimated. They are considered as being particles of a decomposition process at the end of which the formation of \mathcal{J}_{l} -mesons was observed. A scheme of the decomposition process can be written as follows: $T \to n^{-1} \mathcal{J}_{l}^{\circ}$ (\mathcal{J}_{l}°) $\to Q$, where Q is energy carried away by the neutron and the meson, from the T-particle when the latter is in a state of rest. Six references; 2-USSR and 4-Foreign (1953-1954). Table; illustrations. Physical Institute of the Acad. of Scs. of the Arm SSR Physical Institute of the Acad. of Scs. of the USSR Institutions: Submitted Translation M-819, 12 Oct 17

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