ACCESSION NR: AT4033995

the feasibility of trans- and cis-configurations. Orig. art. has: 2 tables and numerous chemical formulas.

ASSOCIATION: Moskovskiy gosudarstvennyky universitet im. M. V. Lomonosova

(Moscow State University)

SUBMITTED: 31Ju162

DATE ACQ: 30Apr64

ENCL: 00

SUB CODE: OC

NO REF SOV: 005

OTHER: 001

Card 2/2

APPROVED FOR RELEASE: 06/14/2000

CIA-RDP86-00513R001134820011-0"

ACCESSION NRt		ASD PO-14 RM/MAY S/0190/63/005/006	0837/08/1	63
AUTHOR: Terent	yev. A. P.; Rukhadze	, Iq. G.; Mochaline, I.		
	on high-moleculer no	ridine derivatives. I.		
Source: Vy*soko	molekulyarny*ye soyed	lineniya, v. 5, no. 6, 3	963, 837-841	
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ABSTRACT: To ac	hieve the synthesis o	of desired polyamides a	solution of the d	12
olycondensation	(with an alkaling on	acid in benzene was re	acted by interfac	ial
enediamine, ben	zidine. o-tolidine. /	dylenediamine, tetramet	hylenediamine, o-	pheny-
lightly colored	Substances, almost i	ngoluble in angel	yamides were white	or
35-2400 and sil	chain lowered the me	solubility and wise poly	emides from 298-3(OC to
rd 1/2	u tue macromolecules	rendered them harder and	i more thermoresis	itant
The first territory of the contract of a single contract of the contract of th	riprose de capación a propio de la company d			

T 13520-63 ACCESSION NR1 AP3001154			
resistance to concentrate	ity. The obtained polyamides we depreciate the hydrochloric acid, sodium hyd ination revealed their amorphouses.	roxide, browine, and	Maro-
ASSOCIATION: Moskovskiy State University)	gosudarstvenny*y universitet im	. M. V. Lomonosova (M	oscow
SUBMITTED: 20Nov61	DATE ACQ: OlJul63	ENCL: OO	
SUB CODE: CO	NO REF SOV: 010	OTHER: 002	
Cord 2/2		. 1996년 - Bartis - 1997년 1월 1일 등학 등 1921년 - 1921년 1921년 - 1921년 - 1921년 1921년 1921년 1921년 1921년 1921년 1921년 1	Cir.

UTHOR: Terent'yev, A. F.; Rukhadse, Ye. G.; Panova. G. V.; Mochalina, I. G. ITIE: Studies on high-molecular pyridine derivatives. 2. Polyamides and poly- hiosmides on the basis of certain alkylpyridines 7 SOURCE: Vy*sokomolekulyarny*ye soyedineniya, v. 5, no. 6, 1963, 842-845 FOPIC TAGS: pyridine, polyamide, polythioamide, alkylpyridine, tautomerism ABSTRACT: The present investigation is a continuation of the earlier work by the authors; on polyamides on the basis of 2,6-lutidine. Since the objective was the synthesis of thioamides and polythioamides, this study also included alpha-picoline. These compounds were obtained by prolonged heating of the corresponding alkylpyri- line and of diamines (bensidine, o-tolidine, and o-dianizidine) with an excess of sulfur at 160-200C. Where 2,6-lutidine constituted the base, it was necessary to sulfur at 160-200C. Where 2,6-lutidine constituted the base, it was necessary to obtain polythioamides, while at a 4:1:10 ratio only one methyl group of 2,6- lutidine entered into the reaction. These synthesized polythioamides are colored lutidine entered into the reaction. These synthesized polythioamides are colored powdery substances, some of which show an amorphous structure on a Debyegram, while infrared spectra revealed the presence of a thioureide group. It is concluded that	. 13521-63 CCESSION WR:	EMP(j)/EMT(m)/BDS AP3001155	ASDPc=4RM S/0190/63/005/0		62
TILE: Studies on high-molecular pyridine derivatives. 2. Polyamides and poly- hicamides on the basis of certain alkylpyridines 7 SOURCE: Vy*sokomolekulyarny*ye soyedineniya, v. 5, no. 6, 1963, 842-845 FOPIC TAGS: pyridine, polyamide, polythicamide, alkylpyridine, tautomerism ABSTRACT: The present investigation is a continuation of the earlier work by the authors on polyamides on the basis of 2,6-lutidine. Since the objective was the synthesis of thicamides and polythicamides, this study also included alpha-picoline. These compounds were obtained by prolonged heating of the corresponding alkylpyridine and of diamines (bensidine, o-tolidine, and o-diamizidine) with an excess of sulfur at 160-200C. Where 2,6-lutidine constituted the base, it was necessary to sulfur at 160-200C. Where 2,6-lutidine and a tenfold quantity of sulfur in order react it with a double amount of diamine and a tenfold quantity of sulfur in order to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6- to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6- lutidine entered into the reaction. These synthesized polythicamides are colored lutidine entered into the reaction.	UTHOR: Teren	yev, A. P.; Rukhadse, Y	e. C.; Panova, C. V	.; Mochalina, I. G.	
COURCE: Vy*sokomolekulyarny*ye soyedineniya, v. 5, no. 6, 1963, 842-845 EOPIC TAGS: pyridine, polyamide, polythicamide, alkylpyridine, tautomerism ABSTRACT: The present investigation is a continuation of the earlier work by the authors on polyamides on the basis of 2,6-lutidine. Since the objective was the authors of thicamides and polythicamides, this study also included alpha-picoline. Synthesis of thicamides and polythicamides, this study also included alpha-picoline. These compounds were obtained by prolonged heating of the corresponding alkylpyriline and of diamines (benzidine, o-tolidine, and o-dianizidine) with an excess of sulfur at 160-200C. Where 2,6-lutidine constituted the base, it was necessary to sulfur at 160-200C. Where 2,6-lutidine and a tenfold quantity of sulfur in order react it with a double amount of diamine and a tenfold quantity of sulfur in order to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6- to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6- to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6- to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6- to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6- to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6- to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6-	TMT 0	on high-molecular pyrid	ine derivatives. 2.	Polyamides and poly	
FOPIC TAGS: pyridine, polyamide, polythicamide, alkylpyridine, tautomerism and the present investigation is a continuation of the earlier work by the authors on polyamides on the basis of 2,6-lutidine. Since the objective was the authors of thicamides and polythicamides, this study also included alpha-picoline. Synthesis of thicamides and polythicamides, this study also included alpha-picoline. These compounds were obtained by prolonged heating of the corresponding alkylpyriline and of diamines (benzidine, o-tolidine, and o-dianizidine) with an excess of dine and of diamines (benzidine, o-tolidine, and o-dianizidine) with an excess of sulfur at 160-200C. Where 2,6-lutidine constituted the base, it was necessary to react it with a double amount of diamine and a tenfold quantity of sulfur in order to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythicamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythicamides are colored lutidine on a polythicamides are colored lutidine on a polythicamide and a tenfold quantity of sulfur and a tenfold	OURCE: Vy*so	komolekulyarny*ye soyedin	eniya, v. 5, no. 6,	1963, 842-845	
ABSTRACT: The present investigation is a continuation of the earlier work by the authors on polyamides on the basis of 2,6-lutidine. Since the objective was the authors of thioamides and polythioamides, this study also included alpha-picoline. Synthesis of thioamides and polythioamides, this study also included alpha-picoline. Synthesis of thioamides and polythioamides, and o-dianizidine) with an excess of dine and of diamines (benzidine, o-tolidine, and o-dianizidine) with an excess of sulfur at 160-200C. Where 2,6-lutidine constituted the base, it was necessary to sulfur at 160-200C. Where 2,6-lutidine and a tenfold quantity of sulfur in order react it with a double amount of diamine and a tenfold quantity of sulfur in order to obtain polythioamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythioamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythioamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythioamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythioamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythioamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythioamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythioamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythioamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythioamides, while at a 4:1:10 ratio only one methyl group of 2,6-to obtain polythioamides are colored lutidine.					
	and the second synthesis of the second of displaying the second of displaying the sulfur at 160-react it with to obtain poly lutiding enter	present investigation is yamides on the basis of 2 hioamides and polythioamis were obtained by prolonamines (benzidine, o-toli 2000. Where 2,6-lutidines double amount of diaminthioamides, while at a 4 ed into the reaction. The	a continuation of 2,6-lutidine. Since ides, this study also read heating of the iden, and o-dianize constituted the bene and a tenfold que:1:10 ratio only on hese synthesized po	the earlier work by the objective was to included alpha-pi corresponding alkyl (dine) with an excessive, it was necessariantity of sulfur in methyl group of 2, lythicamides are collars on a Debyegram,	coline. pyri— s of y to order 6— ored while

ACCESSION NRf AP3001155			
the thicamides and polythi has: 4 formulas.	cemides may exist in two taut	omeric forms. Orig	art.
ASSOCIATION: Moskovskiy g State University)	osudarstvemy*y universitet i	m. M. V. Lomonosova	(Koscovi
SUBMITTED: 20Nov61	DATE ACQ: Oljul63	ENCL: OL	
SUB CODE: OO	No ref sove co6	OTHER: 002	
Cord 2/80			

ACCESSION NR: AP4042188

\$/0190/64/006/007/1267/1271

AUTHOR: Terent'yev, A. P.; Mochalina, I. G.; Rukhadze, Ye. G.; Povolotskaya, Ye. H.

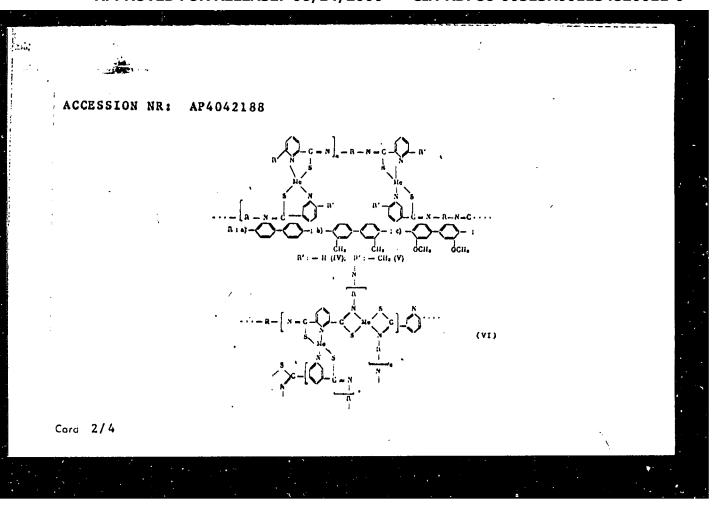
TITLE: Study in the series of polymeric chelates. X. Some physicochemical studies of polymeric chelates based on thio amide and poly(thio amide) derivatives of pyridine

SOURCE: Vy*sokomolekulyarny*ye soyedineniya, v. 6, no. 7, 1964, 1267-1271

TOPIC TAGS: polymeric chelate, thiamide pyridine derivative, polythiamide pyridine derivative, Cu, Ni, Co, Zn, polymeric chelate property, chelate group structure

ABSTRACT: The properties of the following Cu-, Ni-, Co-, or Zn-containing polymeric chelates have been studied.

Card 1/4



ACCESSION NR: AP4042188

The polymers are fine, colored amorphous powders which are soluble only with difficulty. They dissolve most readily in dimethyl-formamide, chloroform, and benzene; the best solubility is exhibited by polymers with metoxy groups in the backbone. The polymers are attacked by acids but are resistant to alkalis. They lose 5-15% of their weight on heating to 200C and 20-80% on heating to 400C. Comparative studies showed that polymers with a higher solubility have a lower chemical and thermal stability. The density of polymeric chelates varies from 1.27 to 1.60. They are dielectrics at room temperature; at 383K the highest electrical conductivity (10-10 to 10-11 ohm-1 cm-1) is exhibited by Cu-containing polymers. Catalytic. properties are inherent only in Cu-, Ni-, and Co-containing polymers. Their catalytic activity surpasses by two orders of magnitude that of inorganic Cu semiconductors. The nature of the band in the chelate group, as determined from x-ray absorption spectra, is =C-S-Me. Orig. art. has: 3 tables.

ASSOCIATION: Moskovskiy gosudarstvenny*y universitet im M. V. Lomonosova (Moscow State University)

Card 3/4

ACCESSION NR: AP4042188

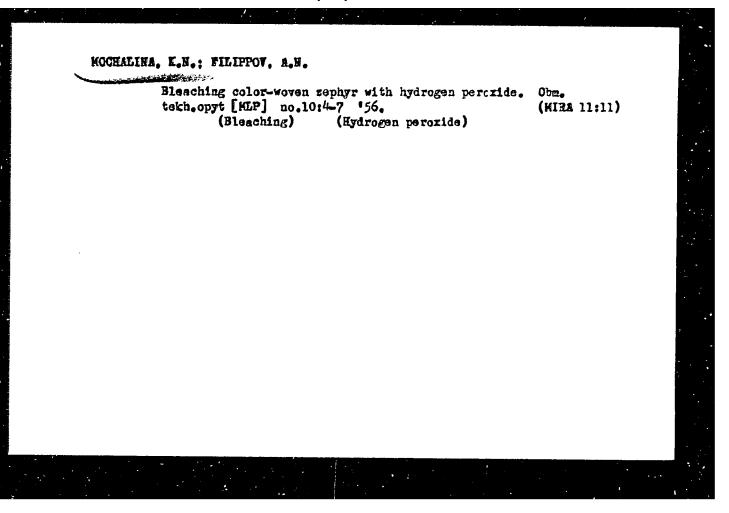
SUBMITTED: 02Aug63 ATD PRESS: 3073

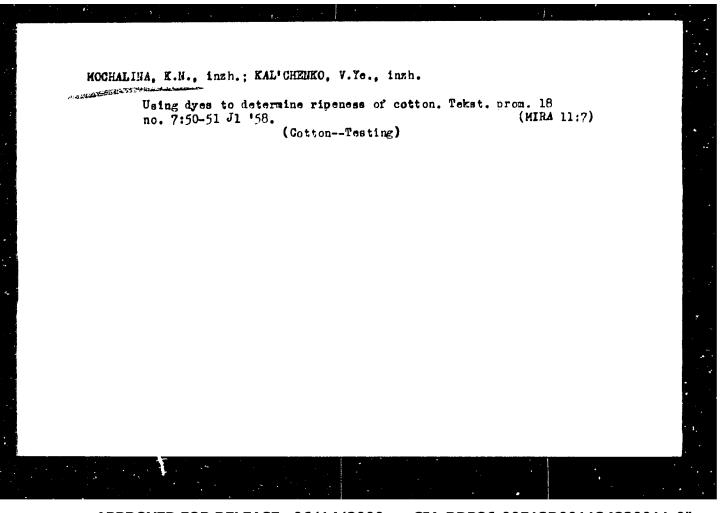
ENCL: 00

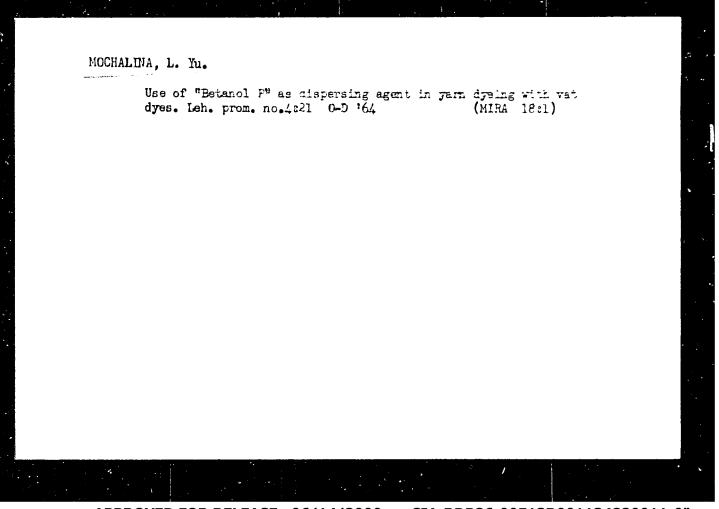
SUB CODE: OC, GC NO REF SOV: 006

OTHER: 002

Card 4/4







APPROVED FOR RELEASE: 06/14/2000 CIA-RDP86-00513R001134820011-0"

82041 s/062/60/000/02/04/012 B003/B066

5.3600

Knunyants, I. L., Dyatkin, B. L., German, L. S., AUTHORS:

Mochalina, Ye. P.

TITLE:

Reactions of Flucro-olefins. 12th Report. Interactions of

Polyfluoro-chloro Butenes With Alcohols

PERIODICAL: Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk,

1960, No. 2, pp. 231 - 236

TEXT: The authors investigate the action of sodium methylate and ethylate on linear dimers of 1,2-difluoro-1,2-dichloro ethylene and trifluoro-chloro ethylene. The experiment is described in detail in the experimental part of the paper. The structure was clarified by means of infrared spectrography. The investigations revealed that the reaction of 1,2,3,4-tetrafluoro-1,3,4,4-tetrachloro butene-1 with the alcoholates mentioned yields 1,1,1-trialkoxy-2,3,4-trifluoro-4,4-dichloro butene-2. When treating the linear dimer of trifluoro-chloro ethylene with the alcoholates, 3-alkoxy-4-chloro-perfluoro butene-1 results. The linear dimer of trifluoro-chloro ethylene was identified to be a mixture of

Card 1/2

CIA-RDP86-00513R001134820011-0" **APPROVED FOR RELEASE: 06/14/2000**

Reactions of Fluoro-olefins. 12th Report. Interactions of Polyfluoro-chloro Butenes With Alcohols \$/062/60/000/02/04/012 B003/B066

3,4-dichloro-perfluoro butene-1 and 1,4-dichloro-perfluoro butene-2 (with the latter being predominant). There are 17 references: 4 Soviet, 10 American, 1 Belgian, and 1 German.

ASSOCIATION: Institut elementoorganicheskikh soyedineniy Akademii nauk SSSR (Institute of Elemental-organic Compounds of the

Academy of Sciences USSR)

SUBMITTED: July 4, 1958 (initially)

July 31, 1959 (after revision)

Card 2/2

KNUNYANTS, I.L.; GERMAN, L.S.; DYATKIN, B.L.; MOCHALINA, Ye.P.

Condensation of 1,2-difluoro-1,2-dichloroethylene with formaldehyde. Zhur.VKHO 6 no.1:114 '61. (MRA 14:3)

1. Institut elementoorganicheskikh soyedineniy Akademii nauk SSSR. (Ethylene) (Formaldehyde)

25481 \$/020/61/139/001/013/018 B103/B226

AUTHORS:

Dyatkin, B. L., Mochalina, Ye. P., and Knunyants, I. L.,

Academician

TITLE:

Condensation of formaldehyde with perfluoro olefines - tetrafluoro ethylene, hexafluoro propylene, and trifluoro

chloroethylene

PERIODICAL:

Akademiya nauk SSSR. Doklady, v. 139, no. 1, 1761, 106-109

TEXT: The authors continued the investigation of chlorosulfonic acid as a catalyst of H. Prins' reaction (Ref. 1: Rec. trav. chim., 51, 469 (1952)) and endeavored to use this acid when extending the Prins reaction to perfluoro olefines: tetrafluoro ethylene, hexafluoro propylene, and trifluoro chloroethylene, in their interaction with formaldehyde. As is known, the Prins reaction belongs to the typical reactions of hydrocarbon olefines with electrophile reagents; especially, in this case, the condensation with formaldehyde in the presence of strong acids is meant. Such reactions are very difficult and, therefore, little investigated. The authors demonstrated by means of 1,2-difluoro-1,2-dichloro ethylene that Card 1/5

25481 S/020/61/139/001/013/018 B103/B226

Condensation of formaldehyde...

both chlorosulfonic and fluorosulfonic acid are in this case highly effective catalysts of the Prins reaction, while $\rm H_2SO_A$ of various

concentrations could not release this reaction (I. L. Knunyants et al. Ref. 5: Zhurn. Vsesoyuzn. khim. obshch. im. Mendeleyeva. 6, 114 (1961)). It has been proved that a mixture of tetrafluoro ethylene CF₂ = CFC1.

paraformaldehyde, and chlorosulfonic acid, heated up to 100° C, is subject to a condensation according to the general scheme of the Prins reaction, and yields α,α -diffuoro hydracrylic acid which is isolated as its ethyl ester. As the yield of this ester was 6°6%, the authors were of the opinion that chlorosulfonic acid is much more active than $\frac{1}{2}$ SO₄. The

condensation of paraformaldehyde with hexafluoro propylere leads in the presence of chlorosulforic acid at 130 - 150°C to a 41-% yield of

Card 2/5

25481 S/020/61/139/001/013/018 B103/B226

Condensation of formaldehyde...

of fluoroacetic acid are produced. Thus, the condensation of formaldehyde with trifluoro chloroethylene could be expected to lead to α -fluoro- α -chloro

hydracrylic acid: $EOCH_2^+ + CFC1 = CF_2 \longrightarrow \left[HOCH_2CFC1CF_2\right]^{\frac{1}{2}O} \rightarrow HOCH_2CFC1COOH.$

This acid has actually been isolated as its ethyl ester. The yield amounted to 19.8 % of the theoretical one. However, beside this yield, another 30.5 % of ethyl ester of α, α -difluoro hydracrylic acid have been produced. Its formation can be only explained as a result of the electrophile attack to the CF₂ group, i.e., to a negatively polarized carbon atom being more weak

than is the case with the C atom in the CFC1 group of trifluoro chloroethylene. I. L. Knunyants, V. V. Shokina and Li Chih-yüan (Ref. 9: DAN, Card 3/5

25\tau81 \$/020/61/139/001/013/018 B103/B226

Condensation of formaldehyde ...

136, 611 (1961)) observed two types of orientation in the addition of todine chloride to trifluoro chloroethylene: $CF_2 = CFC1 + IC1 \longrightarrow CF_2I \longrightarrow CFC1_2 + CF_2C1 \longrightarrow CFCII$.

It could not be found out whether this reaction is released by the ionic or by the radical mechanism. However, in the condensation with formaldehyde, an orientation being opposed to polarity appears in a pronounced ionic process, viz., that of the electrophile addition to the double tond. The authors try to explain this phenomenon by the competition of the polar and steric factors. The effective radius of F is 1.25 Å, that of Cl 1.58 Å. Thus, the orientation of reaction which corresponds to polarity meets a great steric hindrance. The steric and polar factors, however, agree as to their effect in the nucleophile addition to trifluoro chioroethylene. Due to this fact, orientation in these reactions has to be a rigorously unambiguous one. There are 9 references: 2 Soviet-bloc and 7 non-Soviet-bloc. The three references to English-language publications read as follows: M. S. Raasch. (Ref. 2: Am. pat. 2452791); D. D. Coffman et al. (Ref. 3: J. Org. Chem. 14, 747 (1949)); E. T. McBee et al. (Ref. 4: J. Am. Chem. Soc., 74, 444 (1952)).

Card 4/5

S/020/61/139/001/013/018 Condensation of formaldehyde... B103/B226

ASSOCIATION: Institut elementoorganicheskikh soyedineniy Akademii nauk

SSSR (Institute of Elemental-organic Compounds of the

Academy of Sciences USSR)

SUBMITTED: March 23, 1961

Card 5/5

39794 \$/062/62/000/008/014/016 B101/B180

11.7714

AUTHORS:

Knunyants, I. L., Dyatkin, B. L., and Mochalina, Ye. P.

TITLE:

Anionotropic rearrangement in reactions of perfluorobutadiene

PERIODICAL:

Akademiya nauk SSSR. Izvestiya. Otdeleniye khimicheskikh

nauk, no. 8, 1962, 1483-1484

TEXT: The addition of nucleophilic reagents (e.g. alcohol) to hexafluorobutadiene-1,3 in the presence of triethylamine was investigated at 80° C. As oxidation of the addition compound with permanganate gives trifluoroacetic acid, while hydrolysis with $\rm H_2SO_4$ gives trifluoroacetone,

Card 1/2

Anionotropic rearrangement in ...

S/062/62/000/008/014/016 B101/B180

fluoroacetic acid in quantitative yield, and its hydrolysis with ${\rm H_2SO_4}$ gave trifluoroacetone. Since neither anhydrous HF nor triethylaminefluorohydrate cause a rearrangement of the dialkylamine of (I) into the corresponding compound of (II), the triethylamine is assumed to have a

 $CF_2 = CF - CFHCONR_2 + R_3N \longrightarrow CF_2 = CF + CH + \Theta NR_3$ → CF₃CF=CH-CONR₂ + R₃N.

The formation of a carben as an intermediate is also possible:

ASSOCIATION:

Institut elementoorganicheskikh soyedineniy Akademii nauk SSSR (Institute of Elemental Organic Compounds of the

Academy of Sciences USSR)

SUBMITTED:

February 19, 1962

Card 2/2

KNUNYANTS, 1.L.; DYATKIN, B.L.; GERMAN, L.S.; MECHALINA, Ye.P.

Condensation of formaldehyde with trifluoroethylene. Izv.AN SSSR.0td.
khim.nauk no.9:1674 1677 S '62. (MIRA 15:10)

1. Institut elementoorganicheskikh soyedineniy AN SSSR.,
(Formaldehyde) (Ethylene) (Chlorine compounds)

DYATKIN, B.L.; MOCHALINA, Yo.P.

Fluoroaliphatic diazo compounds. Report No.1: Some new reactions of trifluorodiazoethane. Izv. AN SSSR Ser. khim. no.7:1225-1229 J1 '64. (MIRA 17:8)

1. Institut elementoorganicheskikh soyedineniy AN SSSR.

MOCHALINA, Ye.P.; DYATKIN, B.L.

Synthesis of 2-diazoperfluoropropane (bistrifluoromethyldiazomethane. Izv. AN SSSR. Ser. khim. no.5:926-928 '65. (MIRA 18:5)

1. Institut elementoorganicheskikh soyedineniy AN SSSR.

DYATKIN, B.L.; MOCHALINA, Ye.P.

Fluoroaliphatic diazo compounds. Report No.2: ∞-Diazoperfluoro ketones. Izv. AN SSSR. Ser. khim. no.6:1035-1039 '65.

1. Institut elementoorganicheskikh soyedineniy AN SSSR.

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1. Institut elemen objgantorepolika byje ilmenty (1912).	

DYATKIN, B.L.; MOCHALINA, Ye.P.; KNUNYANTS, I.L.

Oxidation of hexafluoroacetone oxime in anhydrous hydrogen fluoride. Izv. AN SSSR. Ser. khim. no.9:1715-1716 '65.

(MIRA 18:9)

1. Institut elementoorganicheskikh soyedineniy AN SSSR.

DYATKIN, B.L.; MOCHALINA, Ye.P.; LANTSEVA, I.T.; KNUNYANTS, I.L.

Hexafluoroisobutyric acid in the Boredin-Hunsdiecker reaction. Zhur.VKHC 10 no.4:469-470 '65.

(MIRA 18:11)

1. Institut elementoorganicheskikh soyedineniy AN SSSR.

L 31885-66 ENT(m)/ENP(j)/T WW/JW/JWD/RM

ACC NR: AP6012539

SOURCE CODE: UR/0062/66/000/003/0585/0585

AUTHOR: Cyatkin, B. L.; Mochalina, Ye. P.; Bekker, R. A.; Knunyants, I. L.

44

ORG: Institute of Elemental Organic Compounds, Academy of Sciences SSSR (Institut elementarnoorganicheskikh soyedineniy Akademii nauk SSSR)

TITLE: Mechanism of addition of nitrosyl fluoride to fluoroolefins

SOURCE: AN SSSR. Izvestiya. Seriya khimicheskaya, no. 3, 1966, 585

TOPIC TAGS: organic synthesis, fluorine compound

ABSTRACT: The authors obtained experimental proof of the nucleophilic mechanism of addition of FNO to higher fluoroolefins. It was shown that alkali metal fluorides (especially in strongly ionizing media) are effective catalysts of this reaction

Perfluoroethylene reacts with FNO only at 120-150°C and the reaction of perfluoroethylene (15 g), FNO (5.5 g), CsF (2 g) and KF (4 g) produces 2-nitrosoperfluoropro-

Card 1/2

UDC: 541.124 + 546.16

L 31885-66

ACC NR: AP6012539

pane (14.3 g). Perfluorocyclobutene according to our observations does not react with FNO even during heating to 120°C; however, shaking of 8 g of perfluorocyclobutene, 4.5 g of FNO, 1 g CSF and 2 kg KF with 5 ml of tetramethylene sulfone at 35° for 3 hrs produces nitrosoperfluorocyclobutane with 70% yield.

SUB CODE: 07/ SUBM DATE: 30Dec65/ ORIG REF: 002/ OTH REF: 001

(Card 2/2

ACC NR: AP7010719

SOURCE CODE: UR/0062/66/000/012/2247/2248

AUTHOR: Mochalina, Ye. P.; Dyatkin, B. L.; Knunyants, I. L.

ORG: Institute of heteroorganic compounds, Academy of Sciences USSR (Institut elementoorganicheskikh soyodiniy AN SSSR)

TITLE: Fluorine-containing phosphazo-compounds

SOURCE: AN SSSR. Izvestiya. Seriya khimicheskaya, no. 12, 1966, 2247-2248

TOPIC TAGS: organic nitrogen compound, organic phosphorus compound, fluorinated organic compound

SUB CODE: 07

ABSTRACT: It was found that fluorine-containing phosphazo-compounds may be prepared by reaction triphenylphosphine and triethylphosphite with tertiary perfluoronitrosoalkanes, in 75 and 68% yield, respectively. The reaction products were characterized. The first case of the formation of phosphazo compounds from perfluoronitrosoalkanes and derivatives of trivalent phosphorus was reported in 1965 at the laboratory of the authors. Orig. art. has: 2 formulas. [JPRS: 40,351]

Card 1/1

UDC: 547.221 + 661.718

OMETILIN, J. A.; COUNLETT, J. Ya. - Largest a a ricelter of Institute "Fundamentian of Mounds" Vetering riya, 31, No. 7, Andrews, prof-50 of County and the county and County a

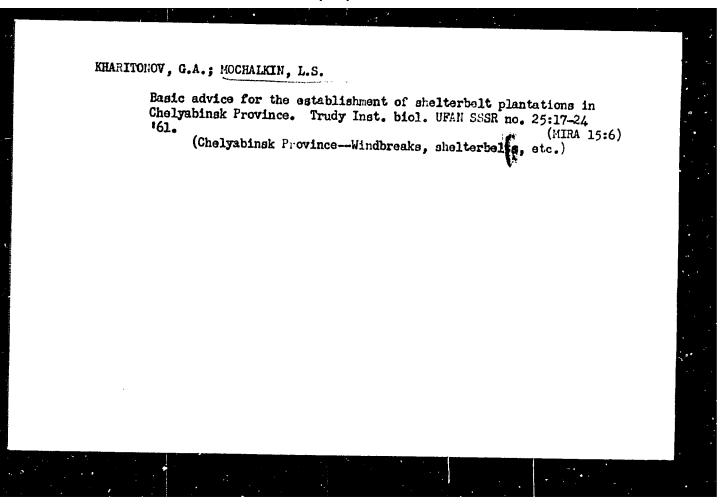
KURHATOV, L.N.; KABANOV, A.N.; SIGRIYANSKIY, V.V.; MASHCHENKO, V.Ye.; MOCHALKIN, N.N.; SHARIK, A.I.; SOROHO-NOVITSKIY, N.V.

Generation of coherent radiation in specimens of gallium arsenide following electronic excitation. Dokl. AN SSSR 165 no.2:303-304 N *65. (MIRA 18:11)

1. Submitted March 15, 1965.

	10241-66 EWT(d)/FBD/EWT(1)/EEC(k)-2/EPF(n)-2/T/EWP(k)/EWA(m)-2/EVIA(h) SCTR/LIP(c) CC NR. AP5028275 HG/WW/AT SOURCE CODE: UR/0020/65/165/002/0303/0304	
PK	UTHOR: Kurbatov, L. N.; Kabanov, A. N.; Sigriyanskiy, V. V.; Mashchenko, V. Ye.;	
OI	RG: name	
T	ITLE: Generation of coherent radiation in GaAs samples excited by electrons	
S	OURCE: AN SSSR. Doklady, v. 165, no. 2, 1965, 303-304	
	OPIC TAGS: laser, semiconductor laser, electron beam, gallium arsenide,	
Α.	BSTRACT: Laser action at 77K and at room temperature is reported in both n- and p- ype GaAs excited with a beam of electrons. The Fabry-Perot cavity was prepared by	
	leaving in the (110) plane. The resonator mirror surfaces were separated by a dis-	
6	ance of 50-60 μ . An electron beam device supplied electrons with energies up to 0 kev. The repetition rate and the pulse duration were 50-200 pulses per second	
8	nd 9 x 10 ⁻⁸ sec, respectively. The maximum beam current at a beam diameter of 0-70 u was 17 mamp. The electron beam was normal to the polished surface of the	
8	ample. The light was emitted from the faces normal to the polished faces. The	- 7
7	hreshold current densities were different for different samples and varied between 0 and 150 amp/cm ² . Since the effective mass of the electron and the width of the orbidden gap in GaAs are larger than in InSb and InAs (two of the other semiconduc-	
ŧ	or lasers) and the lifetime of the electrons is very short, population inversion in	
Co	urd 1/2 UDC: 537.311.33	
		47

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temperatures	s betwee	n 77-300K t	he thresho	ld current	should	depend we	akly o	n the tem-	
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ACCESSION NR: AP3003731

5/0109/63/008/007/1279/1280

AUTHOR: Kuzimin, V. A.; Mochalkina, O. R.

TITLE: Method for reducing the cutoff time of p-n-p-n semiconductor devices

SOURCE: Radiotekhnika i elektronika, v. 8, no. 7, 1963, 1279-1280

TOPIC TACS: p-n-p-n semiconductor device, D235 semiconductor device

ABSTRACT: Practical 4-layer semiconductor structures include one wide (100-200-micron) n-type base. The authors cut the switching time of the D235 device to one-fifth by reducing the lifetime of minority carriers in the wide base. The latter was alloyed with Au atoms. Other characteristics of the device were not impaired. Orig. art. has: I figure, 3 formulas, and I table.

ASSOCIATION: none

SUBMITTED: 28Dec62

SUB CODE: CE

DATE ACQ: 02Aug63

ENCL: 00

NO REF SOV: GOL

OTHER: OOL

Card 1/4

ACCESSION NR: AT4040780

8/2657/64/000/011/0121/0129

AUTHOR: Mochalkina, O. R.

TITLE: Current of an open controllable diode in the switching mode

SOURCE: Poluprovodníkovy*ye prihory* i ikh primeneniye; sborník statey, no. 11, 1964,

121-129

TOPIC TAGS: diode, controllable diode, diode current, switch, diode switch, semiconductor device, rectifier, diode power

ABSTRACT: The author considers the operation of the high-power controllable diode D238 in the switching mode. The maximum current of this diode in the open state is on the order of 10 amperes. The condition of thermal stability, an essential factor in limiting the power given off in a semiconductor device, is defined and explained. The author determines the maximum permissible power and current which may be liberated in the diode in the open condition. It is shown on the basis of a typical characteristic curve of a controllable diode that the current is an almost linear function of voltage over a rather large voltage range. Exactly as in the case of rectifier diodes, temperature instability in a controllable diode may occur in the event that the power dissipated by the diode increases as the junction temperatures rises. For a controllable diode operating

Car.4/8

ACCESSION NR: AT4040780

in the switching mode, instability involves the possibility of an avalanche-type current build-up in the reverse direction as the temperature increases. From this fact, there follows the possibility of an avalanche-like rise in the power given off in the reverse voltage. The power given off by the diode in the open state may be considered as independent of the junction temperature. In fact, this power will decrease somewhat in view of the drop in residual voltage as the temperature of the junction rises. This power cannot rise in an avalanche form, heats the diode only to a certain temperature and cannot lead to thermal instability and the failure of the diode, provided this temperature does not cause melting of the materials present in the device. There exists a critical power, dissipated in the reverse direction, after the attainment of which the diode will always be unstable regardless of the magnitude of the power dissipated in the open state. Critical power is thus determined only by the thermal factor of the device and by the temperature coefficient. In addition, the author shows that the maximum permissible mean current of the open state is a function of the cutoff angle. This function is related to a rise in the residual voltage in the diode as the amplitude of the current flowing through it increases. Formulas are given which determine the maximum permissible value of the mean current of the open diode for several typical cutoff angle

ACCESSION NR: AT4040780

values. The calculation given in the article permits an approximate estimate of the value of the open diode for several typical cutoff angle values. The calculation given in the article permits an approximate estimate of the value of the maximum permissible power of a high-power controllable diode, for specific heat emission, depending on the ambient temperature. Finally, the author demonstrates the considerable dependence of the maximum mean current of the open diode on the cutoff angle, which fixes the duration of time the diode is in the open state. It is important that this dependence be borne in mind when operating a controllable diode in various pulse systems. Orig. art. has: 6 figures and 7 formulas.

ASSOCIATION: none

SUBMITTED: 00

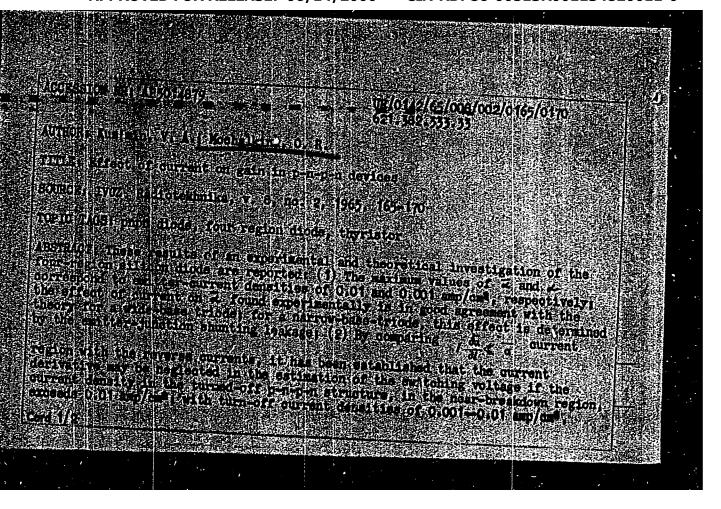
ENCL: 00

SUB CODE: EC

NO REF SOV: 001

OTHER: 003

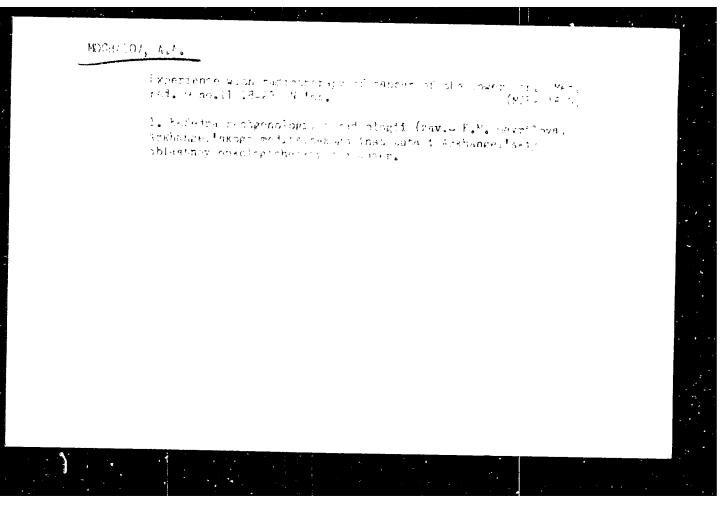
Cara 3/3



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	L 13292-66 EWT(m)/EWP(j) RM	سهد بارد	
	ACC NR: AP6000325 (A) SOURCE CODE: UR/0286/65/000/021/0012/0012		
	INVENTOR: Volkova t. T. Zatena A. W.		
	Nazarova, L. Yu.; Nazarov, V. I.; Pryakhina, M. S.; Petrov, V. N.; Rachkovskiy, E. E.; Savel yev, A. P.; Syrova, A. A.; Tikhanovskaya, S. G.		
	ORG: none		
	Class 12. No. 175920 Farrage and butanol by synthesis from ethyl alcohol.		
_	Petroleum Refining (Bashkirskiy nauchno-issledovatel'skiy institut po pererabotke	-	
	SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 21, 1965, 12		5
	TOPIC TAGS: catalysis, butanol, ethyl alcohol		
	ABSTRACT: This Author's Certificate introduces: 1. A method for producing normal butanol by synthesis from ethyl alcohol on a catalyst. The process is done in a single stage by using a catalyst consisting of aluminum oxide, magnesium oxide, silicon oxide and a salt or oxide of an alkali retail.		
	silicon oxide and a salt or oxide of an alkali metal. 2. A modification of this		
	Card 1/2		
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& magne	in which sium oxi ukali me	de, fr	atalys	t conta o 50 %	nins f silic	rom 5 on oxid	to 80 % de and f	aluminum rom O to	oxide,	from 95 a salt	i to 10 or oxide	
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KOCHALOV, A.

Equipment for making silicate wall blocks. Stroi.mat.itdel. i konstr. 1 no.9:16-19 S'55. (MLRA 9:1)

1. Glavny, inshener Krasnopresnenskogo kombinata stroitel'nykh materialov.
(Building blocks)

MOCHALOV A. SHINKAREV. S.

Two thousand kilograms of lime from one cubic-meter furnace.

Stroi. mat., izdel. i konstr. 2 no.8:21 Ag '56. (MLRA 9:10)

1. Glavnyy inzhener Krasnopresnenskogo kombinata (for Mochalov). (Lime kilns)

MOCHALOV, A.; SHINKAREV, A.

Chromium magnesite bricks used for lining limekilms. Stroi.mat. 3 no.11:20-21 N '57. (MIRA 10:12)

1.Glavnyy inzhener Krasnopresnenskogo kombinata stroitel'nykh materialov (for Mochalov). 2.Nachal'nik tsekha Krasnopresnenkogo kimbinata stroitel'nyk materialov (for Shinkarev). (Magnesite) (Firebrick) (Limekilns)

WCCHALOUA / VOLZHENSKIY, A.V., prof.; MOCHALOV, A.I., inzh.; BUROV, Yu.S., kand. tekhn.nauk; SILAYENKOV, Ye.S., inzh. Autoclaved concrete made with metallurgical slag and ash binders. Bet. i zhel. -bet. no.8:322-325 Ag '57. (MIRA 10:10) 1. Deystvitel'nyy chlen Akademii stroitel'stva i arkhitektury (for

(Concrete)

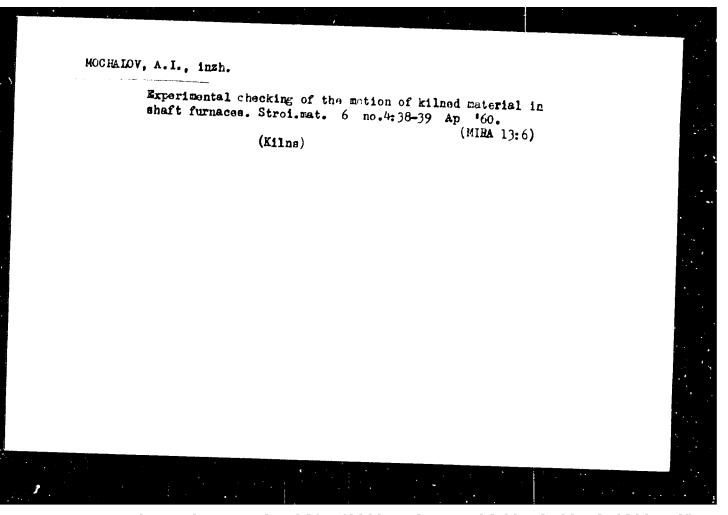
Volzhenskiy)

MOCHALOV, A.; SHAPOVAL, I.; TOMASHPOL'SKIY, L., tekhnolog.

Improving equipment for making lightweight blocks. Stroi. mat. 4 no.3:29-31 Mr '58. (MIRA 11:3)

1. Glavnyy inzhener Krasnopresnenskogo kombinata stroitel'nykh materialov (for Kochalov). 2. Nachal'nik tekhnicheskogo otdela Krasnopresnenskogo kombinata stroitel'nykh materialov (for Shapoval).

(Lightweight concrete)

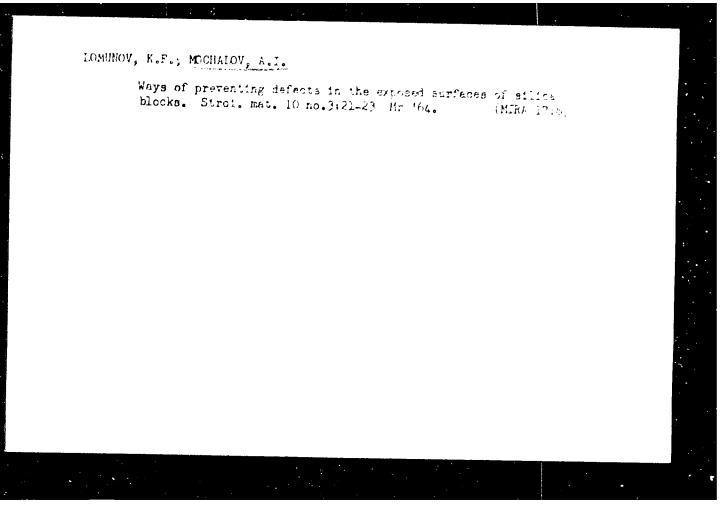


APPROVED FOR RELEASE: 06/14/2000 CIA-RDP86-00513R001134820011-0"

MOCHALOV, A.I., inch.

Kilning fine lime in a staft kiln. Strof. mat. 10 no.1:
37-38 Ja'b4.

(MIRA 17:5:



BUROV, Yu.S., kand. tekhn. nauk; MOCHALOV. A.I., laureat Leninskoy promit KORABLINOV, A.M., inzh.; PEREVOZOV, V.S., inzh.; SEMCHENKOVA T.S., inzh.

Large products made of autoclaved concrete from carbonacects sands. Stroi. mat. 10 no.6:38-40 Je *64. (MIRA 17.10, l. Direktor Krasnopresnenskogo kombinata stroitel*nykh materialov (for Mochalov).

MOCHALOV, A.M.

Preparation of high-purity cadmium by the method of zone melting. Trudy Alt. GMNII AN Kazakh. SSR 9:233-237 '60. (MIRA 14:6)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut tsvetnykh metallov.

(Cadmium-Metallurgy)

(Zone metling)

Mochanical crusher for laboratory samples. Masl.-zhir. prom.
24 no.4:43-44 '58. (MIRA 11:5)

1.Millerovskiy maslockstraktsionnyy zavod.
(Laboratories--Equipment and supplies)
(Crushing machinery)

SOV/137-58-10-20866

, Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 10, p 74 (USSR)

Smirnov, V.S., Pavlov, N.N., Mochalov, A.Ya. AUTHORS:

Grooving for Steel Angles by the Equivalent-strip Method (Kali-TITLE:

brovka uglovov stali po metodu sootvetstvennov polosy)

Prokatnoye i trubnoye proizvodstvo (prilozheniye k zhurnalu PERIODICAL:

"Stal"), Metallurgizdat, 1958, pp 147-167

A communication is presented on the results of the develop-ABSTRACT: ment of the rolling of steel angles when the rolls are grooved by the method of an equivalent strip. A presentation is made of the overall course of the calculation of grooving for angle sections by this method. Prior to the calculation, the relationship of the spread-ratio correction coefficient to the axial ratio is determined, since a change in this relationship induces a change in the effect of the pass walls on the spread. The ratio of the index of spread to reduction for the rolling conditions on the given mill (a 470 mill at one of the Leningrad plants) is determined. It is found that the fluctuating value of the spread ratio correction coefficient renders impossible any further correc-

tion of pass dimensions if it is found that they do not properly Card 1/2

CIA-RDP86-00513R001134820011-0" APPROVED FOR RELEASE: 06/14/2000

SOV/137-58-10-20866

Grooving for Steel Angles by the Equivalent-strip Method

fill with metal. The pass-grooving calculation is adduced. Since 1952 grooving for several angle sections calculated by the method of the equivalent strip has been successfully employed at that plant. An investigation for the purpose of determining roll-separating pressure was conducted during the rolling of angle steel Nr 5 on the 470 mill, and the nature of the distribution of the pressure through the grooves was determined.

B.Ts.

1. Angle bars---Production 2. Rolling mills---Design 3. Rolling mills---Operation

Card 2/2

MCCHALOV, B. D.

Cutting Fachines

"Improvement of bearings for spindles of metal-cutting machines." Pokshipnik no. ϵ , 1952.

Monthly List of Russian Accessions, library of Congress, October 1 52. UNCLASSIFIED.

HOCHILOY. B.M.

Bronchography in a district hospital. Vest. rent. 1 rad. 33 no.6: 82 N-0 '58.

1. Iz Serpukhovskoy bol'nitsy "Krasnyy tekstil'shchik" (glavnyy vrach Ya. G. Surkin) Moskovskoy oblasti. (BHONGHI, radiography in district hosp. (Rus))

MOCHALOV, Boris Kikhaylovich

[Production and consumption in the period of the extensive building of communism] Proizvodstvo i potreblenie v period razvernutogo stroitel stva kommunisma. Hoskva. Znanie. 1960.
45 p. v (MIRA 13:12)

(Russia -- Economic conditions)

MOCHALOV, Boris Mikhaylovich; SUVOROVA, M.I., otv.red.; PAL'CHUN, I.F.,

[Trade under socialism; a lecture on the course in the political economy of socialism] Torgovlie pri sotsialisme; lektalia po kursu politicheskoi ekonomii sotsialisma. Otv.red. M.I.Suvorove. Hoskva, Izd-vo Kosk.univ., 1961. 50 p.

(HIRA 14:4)

(Russia--Commerce)

MOCHALOV, Boris Mikhaylovich; PAL°CHUN, I.F., red.; YERMAKOV, M.S., tekhm.

[Commodity production, the law of value, and money under socialism; lacture on the economics of socialism] Tovarnoe proizvodstvo, zakon stoimosti i den'gi pri sotsializme; lektsiia po kursu politicheskoi ekonomii sotsializma. Moskva, Izd-vo Mosk. univ., 1961. 64 p.

(MIRA 14:8)

(Economics)

MOCHALOV, B. V.

Sep 52

USSR/Electricity - Measuring Instruments

"Review of P. N. Pigina, and N. N. Shumilovskiy's Book, 'Electric Meters'" E. L. Rymar', V. A. Kochan, V. A. Fetrovskiy, B. V. Mochalov, Members of Chair of Automatic and Measuring Devices, L'vov Polytech Inst.

"Elektrichestvo" No 9, pp 95, 96

Favorable review of subject book on elec meters/watt-hour meters/. States that no Soviet tooks had been written on this subject since the works of F. N. Sedov, N. N. Shumilovskiy, and others published in 1933-1935

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(1)

5/129/60/000/011/011/016 E073/E535

AUTHORS:

Mitrofanov, A.A., Candidate of Technical Sciences,

Volkova, M.A/, Letchford, N.I., Mochalov, G.N.,

Engineers

TITLE:

Application of Converter Steel in the Automobile

Industry

PERIODICAL:

Metallovedeniye i termicheskaya obrabotka metallov,

1960, No.11, p.46.

Data are given on industrial tests relating to the TEXT: use of converter steel (0.17% C, 0.46% Mn, 0.032% S, 0.038% P) in the motor car industry. From 8 ton ingots of three commercial melts, strip was rolled which was used for producing rims of truck wheels. It was found that the chemical composition and the mechanical properties are the same as for open hearth steel. the Gor'kiy Automobile Works 2900 such rims were produced and the performance of 1684 of them was closely observed. The number of rejects due to cracking along the weld seam during stretching of the rim was 0.87% for the experimental batch as compared to 0.71% for the batch made of open hearth steel of a similar composition. With these rims disc wheels were made which were fitted on 264 trucks. Card 1/2

S/129/60/000/011/011/016 E073/E535

Application of Converter Steel in the Automobile Industry

So far, these trucks have run over 50 000 km. It is concluded from the results that the investigated converter steel is as good as open hearth steel, particularly for hot rolled and cold rolled sheets which are to be used for deep drawing. There is 1 table.

ASSOCIATIONS: TSNIICHM, GAZ and ZIL

Card 2/2

ARIYA, S.M.; YEROFETEVA, M.S.; MOCHALOV, G.P.

Magnetic susceptibility of strontium subnitride. Zhur.ob.khim.
27 no.7:1740-1743 J1 '57. (MIRA 10:10)

1. Leningradskiy gosudarstvennyy universitet.

(Strontium nitride--Magnetic properties)

MOCHALOV, I.P.

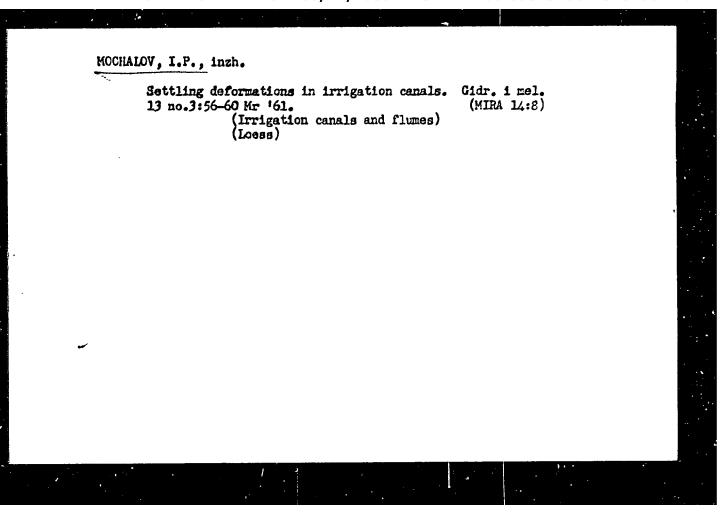
Filtration characteristics of soils in the Golodnaya Steppe. Mat. po proizv. sil. Uzb. no.15:107-112 '60. (MIRA 14:8)

1. Moskovskiy institut inzhenerov vodnogo khozyaystva im.V.R. $Vil^{\dagger}yamsa$.

(Golodnaya Steppe—Soil percolation)

MOCHALOV, I. P.

Cand Tech Sci - (diss) "Features of the utilization of irrigated lands on forest soils." Moscow, 1961. 24 pp with illustrations; (Ministry of Agriculture USSR, All-Union Order of Lenin Academy of Agricultural Sciences imeni V. I. Lenin, All-Union Scientific Research Inst of Hydraulic Engineering and Land Reclamation imeni A. N. Kostyakov); 200 copies; price not given; (KL, 6-61 sup, 221)



APPROVED FOR RELEASE: 06/14/2000 CIA-RDP86-00513R001134820011-0"

Seepage preventing facings and coverings in canals laid in losss-type soils [with summary in English]. Izv. TSEHA no.1:193-205 '62.

(MIRA 15:6)

(Irrigation canals and flumes) (Seepage) (Losss)

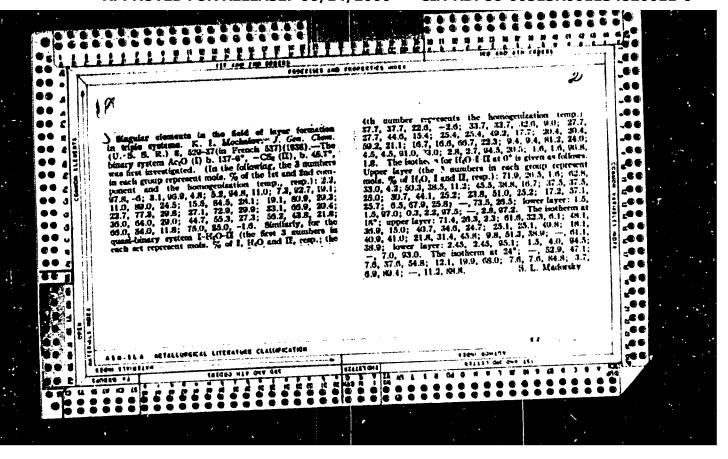
POSLAVSKIY, V.V., akademik; GIRSHKAN, S.A., kand.tekhn.nauk;
MOCHALOV, I.P., kand.tekhn.nauk

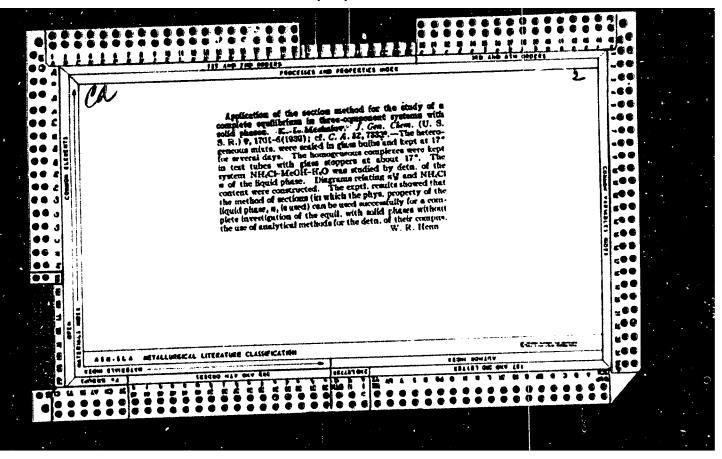
Investigating the seepage from canals. Gidr. i mel.
14 no.6:11-18 Je '62. (MIRA 15:9)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut
gidrotekhniki i melioratsii im. Kostyskova. 2. Akademiya
nauk Uzbekskoy SSR (for Poslavskiy).

(Golodnaya Steppe—Irrigation canals and flumes)

(Seepage)





MOCHALOV, K. I.

"A supplement to the Sectioning Method (metoda secheniy) for the Study of the Full Equilibrium in Tri-member Systems having Solid Phases", Zhur. Obsch. Khim., ?, No. 12, 1039. Laboratory of Inorganic Chemistry, Perm' University. Rec'd 26 March 1939.

Report U-1614, 3 Jrn 1952

S/153/60/003/003/016/036/XX B016/B038

AUTHOR:

Mochalov, K. I.

TITLE:

Equilibrium of the Liquid Phases in the Three-component

System Water - Sulfuric Acid - Ethylienzylan.line

PERIODICAL:

Izvestiya vysahikh uchebnyko zavedeniy. Khimiya . khimicheskaya tekhniligiya , 1960, Vol. 3. No. 3.

pp. 434 - 438

TEXT: The author reports on the study of the recipical solubility in the ternary liquid system water—sulfuric acid—ethylbersylantline $(C_{15}H_{17}N)$. The study was made to obtain better purification and higher yield of $C_{15}H_{17}N$. For this purp se, the author determined the limits of the range of separation into layers, by means of the Alekseyev method (not explained in the text). Moreover, he determined the squalition of the layers being in a state of equilibrium. All together, the author investigated ten cross sections of the temperature correctivation prism with constant content of H_2SO_4 of 6, 10, 15, 20, 20, 17, 35, 40, 46.

Card 1/3

Equilibrium of the Liquid Phases in the Three-component System Water Sulfurio Acid - Ethylbenzylaniline

\$/153/05/053/053/054/046/XX B016/B058

and 46.0%. From the results obtained, he determined a ber of layer formation senthermal lines in the system. Dee numerical data, which that a terize the impositions of the joints of the bic day curves at 0.15, 30, 38, and 50°C are lested in Table 1 and represented in Fig.1. The composition of the equilibrium layers was determined at 15°C (Table 2). Fig. 2 shows the isothermal lines of the terrary sys tem $\rm H_2O$ - $\rm H_2SO_4$ - $\rm C_{15}H_{17}N$. The author concludes from these data that two independent layer formation ranges exist in the system investigated. The one is closed and has an upper critical triple point of dissolution at 40 to 41°C. This point contains (in %): 45 $\rm H_2O$ 34 $\rm H_2SC_4$, and 21 $C_{15}^{\rm H}_{17}^{\rm N}_{\rm L}$ The second range juins the system $\rm H_2C=C_{15}^{\rm H}_{17}^{\rm N}_{\rm L}$ and widens with increasing temperature without exceeding the line of the quasi binary pross section. The author further established that a reversal of the layers takes place near the horizontal control (Fig. 2) in the closed layer formation range. Finally he proved that told theed range develops in consequence of the restricted mutual solubility between the compound

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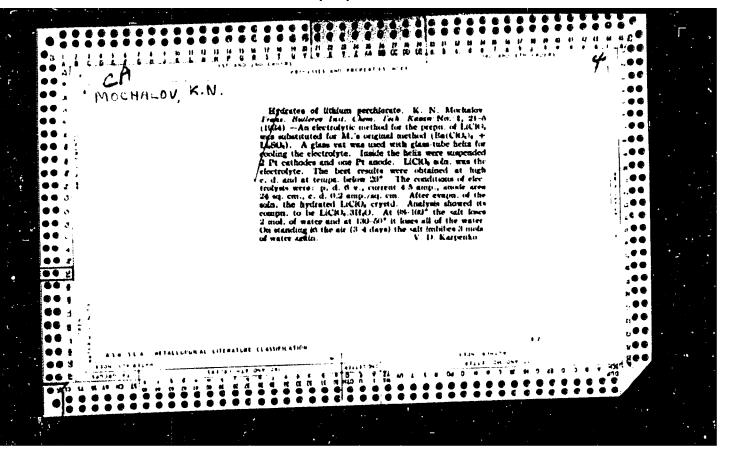
Equilibrium of the Liquid Phases in the Three-component System Water - Sulfaria B016/B058 S/153/60/003/016/036/XX

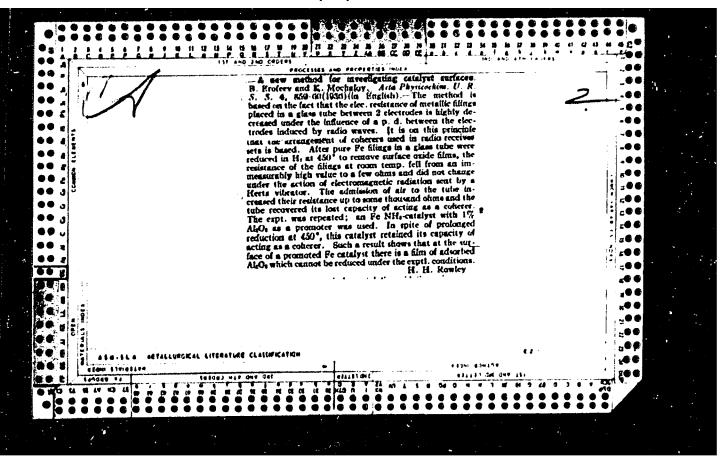
 $c_{15}H_{17}N \cdot H_2SO_4$ and the H_2SO_4 solutions with a maximum electrical fondultivity. Conversely, $c_{15}H_{17}N \cdot H_2SO_4$ is unrestrictedly miscible with water and acts itself as a homogenizer for the layer formation in the binary system $H_2O = c_{15}H_{17}N$. The author mentions papers by V. F. Ust: Kachkintsev and P. A. Khlebnikov (Ref. 4) and reports that the student Yu. L. Medvedev participated in the experimental work. There are 2 figures, 2 tables, and 5 Seviet references.

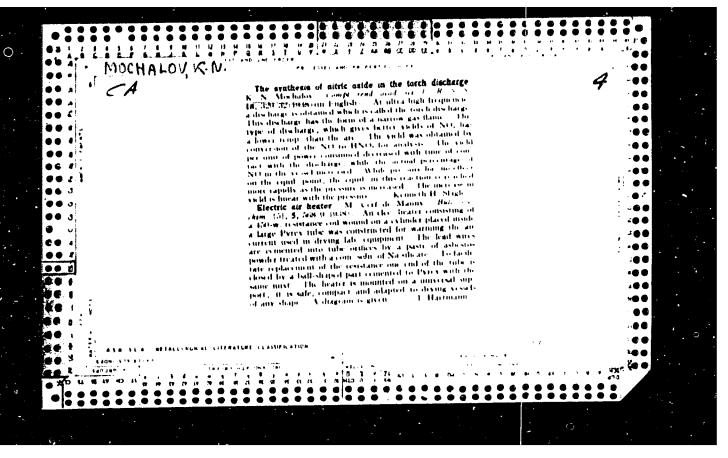
ASSOCIATION: Permskiy gosudarstvennyy universitet im. A. M. Gorikego; Kafedra neorganicheskoy khimii (Permi State University imeni A. M. Gorikiy; Chair of Inorganii Chemistry)

SUBMITTED: October 10, 1958

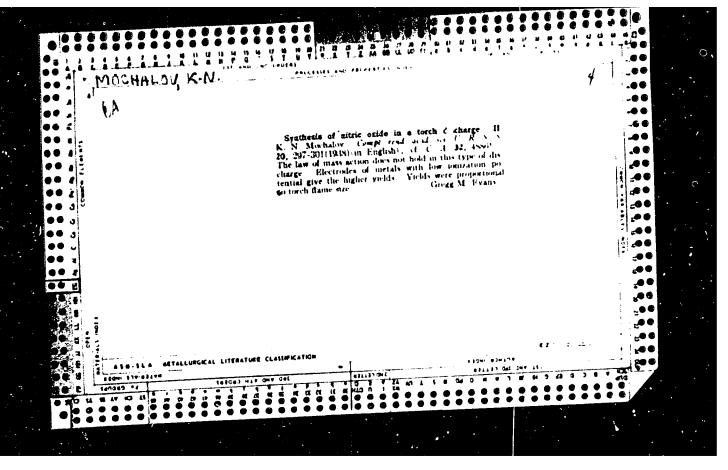
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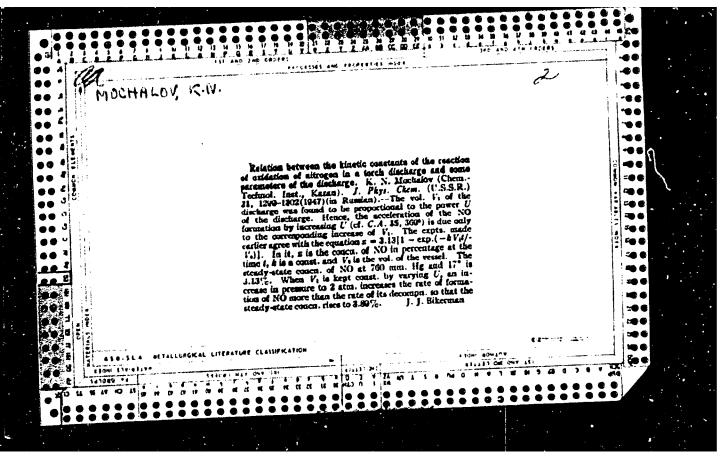


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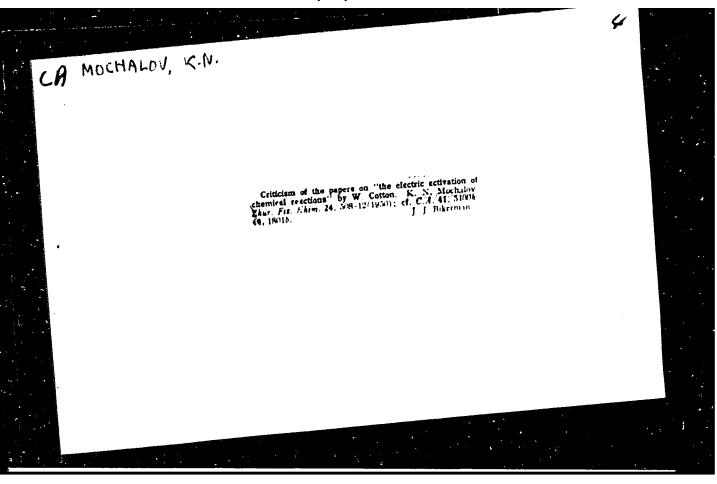
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Measured temperature of external zone of "flame" discharge for frequency of 5.107/ cycles, for various atmospheric pressures, by method of rotation of Li, Na, Tl spectral lines. At external boundary of this zone, temperature equals 2,200°K and gradually increases toward axis of "flame." Submitted 10 Nov 49.

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"Redistribution of Intensities of Spectral Lines of Elements During Discharge in Argon," K. N. Mochalov and Ye. L. Raff, Kazan Chemicotechnolog Inst im Kirov

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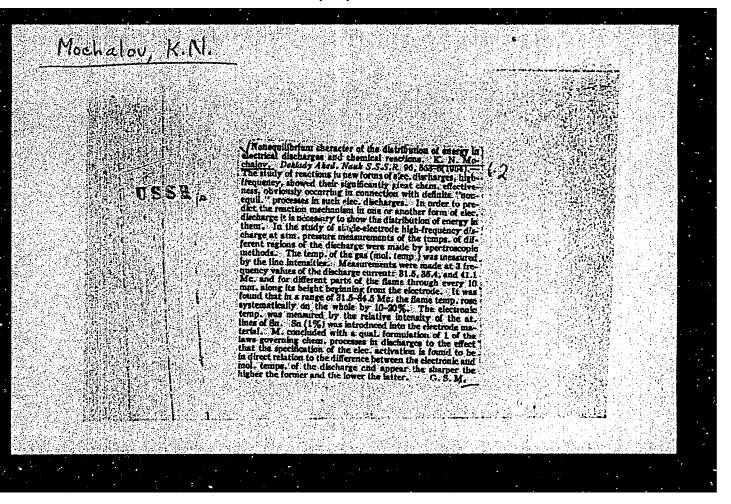
Studied spectra of metals and alloys excited in argon and compared them with spectra in air under identical conditions. Results showed that in argon spectral lines of We ions are enhanced while lines of neutral atoms are weakened. Such redistribution of intensities is also observed in spectra of many other metals. Presented by Acad A. N. Terenin 17 Jun 53.

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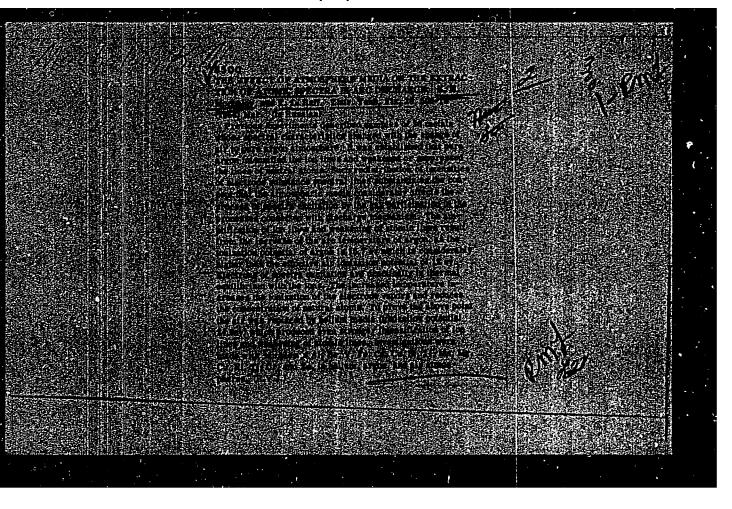
MOCHALOV, Konstantin Nikolayevich

MOCHALOV, Konstantin Nikolayevich (Kazan' Chemical-Technological Inst imeni Kirov), Academic degree of Doctor of Chemical Sciences, based on his defense, 13 April 1955, in the Council of the Inst of General and Inorganic Chemistry imeni Kurnakov, Acad Sci USSR, of his dissertation entitled: "Research in the field of chemical reactions in high-frequency electrical discharge." For the Academic Degree of Doctor of Sciences.

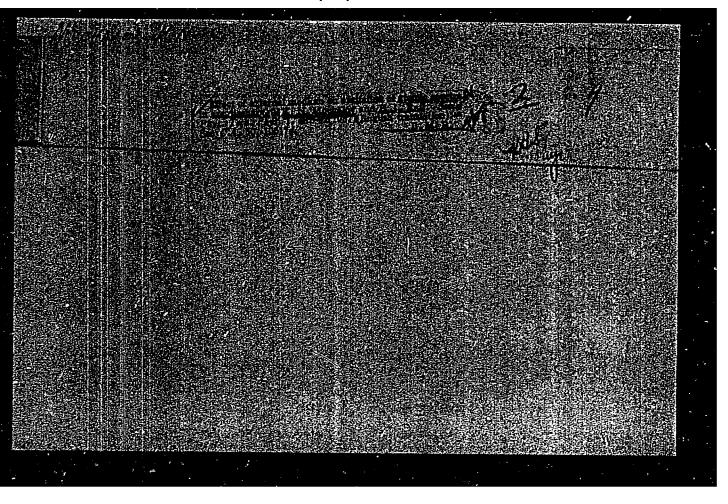
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