

SOV/124-57-9-10348

On the Formation of Whirlpools in Front of Water Gates in Canals

the whirlpools. In the investigations conducted the intensity of a whirlpool formation is characterized by the degree of the transporting capacity of the whirlpool, namely, the number of uniform objects passing through the whirlpool in a specific period of time. According to the test results, an increase in the intensity of the whirlpool formation results in a decrease both of the velocity coefficient  $\phi$  and of the discharge capacity of the gate aperture. As a result of the elimination of whirlpools by means of baffles preventing back currents at the gate the coefficient  $\phi$  can be increased by 1.5 - 3%. The formation of whirlpools occurs in both unsubmerged and submerged outflows. The authors also repeated the tests made by V. S. Fokeyev (Gidrotekhn. str-vo, 1951, Nr 5; Gidrotekhn. i melioratsiya, 1951, Nr 12). The eddy-stimulator water gate recommended by him has contributed toward an increase in the intensity of the whirlpools. A paper by S. M. Isaakyan (RZhMekh, 1957, abstract 533) is devoted to a question similar to that under review.

N. A. Pritvits

Card 2/2

SUBJECT USSR / PHYSICS CARD 1 / 2 PA - 1202  
AUTHOR NIKITOV, A.I.  
TITLE On the Charge Distribution of the Mesons on the Occasion of  
Nucleon-Antinucleon Annihilation.  
PERIODICAL Žurn. eksp. i teor. fis, 30, 1149-1150 (1956)  
Publ. 6 / 1956 reviewed 8 / 1956

S.Z.BELEN'KIJ and I.L.ROZENTAL' (Žurn. eksp. i teor. fis, 30, 595 (1956)) investigated the production of stars on the occasion of the annihilation of antinucleons and computed the probability of the processes with different plurality according to the statistical theory of the plural production of particles. Here the charge distribution computed on the basis of the conservation of isotopic spin is given. Denotations: p - proton, n - neutron,  $\bar{p}$  - antiproton,  $\bar{n}$  - antineutron. The products of annihilation (pions) are characterized by the signs of their charges. The charge distribution for  $p\bar{n}$  is obtained from the distribution for  $p\bar{p}$  (misprint in the original text?) by reversing the signs of the meson charges. A table shows the distribution of the process with given plurality among the charge states. If, e.g. the cross section of the annihilation of  $p\bar{p}$  with production of two mesons has the value  $\sigma_2$ , than it may be seen from this table that 0,167 of the cross section is due to the process

NIKITOV A1

CARD 1 / 2

PA - 1740

SUBJECT USSR / PHYSICS  
 AUTHOR MAKSIMENKO, V.M., NIKITOV, A.I.  
 TITLE The Multiple Production of Particles on the Occasion of  
 Nucleon-Nucleon Collisions at 5,3 BeV.  
 PERIODICAL Žurn.eksp.i teor.fis, 31, fasc.4, 727-729 (1956)  
 Issued: 1 / 1957

The authors theoretically computed the distribution of nucleon-nucleon collisions at 5,3 BeV over the number of secondary particles. These computations were carried out in accordance with the statistical theory of the multiple production of particles with and without consideration of isobaric states. For these computations the method suggested by V.M. MAKSIMENKO, I.L. ROSENTAL', Žurn.eksp.i teor.fis (in print) was used, which makes the exact computation of statistical weights possible.

The statistical weight of the various processes (in %) are shown in form of a table. Two further tables illustrate the further distribution of p - p and n - p - collisions obtained from the postulate for the conservation of isotopic spin. On the occasion of a p - p - collision the process  $NN 2\pi$  (its statistical weight is given by the table) thus leads to the charge state (pp+-) with the probability 0,300, and to the charge state (pp00) with the probability 0,100, etc.

From the aforementioned data it is easy to obtain the distribution of non-elastic collisions over the number of charged particles (rays) which, on the

... I.N. DEBEDEV" of the Academy of Science in  
 the USSR.

NIKITOVA, A.M., Dotsent

Reparative and regenerative processes in focal and fibro-focal pulmonary tuberculosis. Probl.tub. no.2:39-46 Mr-Ap '55.(MLRA 8:6)

1. Iz kafedry patologicheskoy anatomii (zav. -prof. P.P.Erofeyev) Ivanovskogo meditsinskogo instituta.

(TUBERCULOSIS, PULMONARY, physiology, regen. processes in fibrous & focal forms)

BLAGOVESHCHENSKIY, M.A., prof. (Ivanovo, 2-ya Plekhanova ul., d.10)  
NIKITOVA, A.N., dots.

Epifascial progressive gangrene. Vest.khir. 81 no.10:132-136 0 '58  
(MIRA 11:11)

1. Iz fakul'tetskoy khirurgicheskoy kliniki (zav. - prof. M.A.  
Blagoveshchenskiy) i kafedry patologicheskoy anatomii (zav. - prof  
P.P. Yerofeyev) Ivanovskogo meditsinskogo instituta.  
(FACE, ulcer  
phagedenic (Rus))

NIKITOVA, A. N. Doc Med Sci -- "Pathomorphology of the development and self-healing of pulmonary tuberculosis." Mos, 1960 (Acad Med Sci USSR). (KL, 1-61, 205)

-348-

NIKITOVA, A.N., dotsent

Reparative processes in the lungs in primary tuberculosis in children treated with antibacterial preparations. *Pediatria* no.12:3-8 '61. (MIRA 15:1)

1. Iz kafedry patologicheskoy anatomii (zav. - prof. P.P. Yero-feyev) Ivanovskogo meditsinskogo instituta (dir. - dotsent Ya.M. Romanov).  
(TUBERCULOSIS) (STREPTOMYCIN) (CHEMOTHERAPY)

*Nikitsin, P.*  
NIKITSIN, P.

Rural medical station. Rab. i sial. 34 no.1:8 Ja '58. (MIRA 11:1)  
(POGOST (MINSK PROVINCE)—MEDICINE, RURAL)



**NIKITSINA, Alena.**

Exploit. Rab. 1 sial. 33 no.8:16-17 Ag '57.

(MLRA 10:8)

(White Russia--World War, 1939-1945--Underground movements)

NIKITSINA, L.M.

LYUBOSHYTS, I.L., kand.tekhn.nauk; NIKITSINA, L.M., kand.tekhn.nauk.

Drying and heating corn in pneumatic gas dryers before sowing.  
Vestsi AN BSSR. Ser. fiz.-tekhn. nav. no.2:137-143 '57.(MIRA 11:1)  
(Corn (Maize)--Drying)

USSR/Cultivated Plants - Fruits. Berries.

M

Abs Jour : Ref Zhur Biol., No 12, 1958, 53764

Author : Cholyadinova, A.I., Nikitskaya, K.I.

Inst : -

Title : Biological Control of the Development and Growth of the Buds of Fruit and Berry Plants.

Orig Pub : Nauka i perolov, opyt s. kh., 1957, No 7, 48-49

Abstract : Studies of the morphological structure of the fruit buds and also of the degree of their differentiation before winter quiescence and the subsequent development of blossoms in spring were conducted at Moscow University on the following: Siberian crabapple, Vladimir cherry (*Prunus cerasus austera*), and black and golden currants. The blossoms of the Vladimir cherry have - before retiring for the winter - fully formed outer covering organs (calyx and corolla), a developed pistil, and only incipient stamen protuberances. The pollens, the stamen

Card 1/2

*NIKITSEAYA, K. I.*

CHELYADINOVA, A., kand. biol. nauk; NIKITSEAYA, K. I., nauchnyy sotrudnik.

Biological investigation of the growth and development of flower buds of woody plants. Nauka i pered. op. v sel'khoz. 18 no.2:47-49 P '58. (MIRA 11:3)

1. Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosova.  
(Flowers) (Buds)

NIKITSKAYA, V. A.

24

✓ Fracture Structure of Wide-Strip Steel M180. V. A. Nikit-  
skaya. (Stal', 1953, (10), 937-941). [In Russian]. The casting  
and rolling results are considered. Detailed metallographic  
investigation of test-pieces whose fractures indicated defects  
showed that this was often misleading. Laminations and  
other fracture defects were found to be due to external test  
conditions such as rate of loading, type of notch and orienta-  
tion of the test-piece with respect to the direction of rolling.  
Best conditions were maximal rates of dynamic fracture of  
the test-piece and sharp notches, the results also indicating  
tendency to cold brittleness. Transverse test-pieces are  
preferable. — S. Z.

*of g/m*

*N. K. Kraya, K. A.*

✓ 3347. Rapid analysis of Bessemer steel by the method of measurement of thermo-e.m.f. V. M. Yufarov, M. P. Kuznetsov, V. A. Nikitskaya, A. I. Novoschok and I. I. Shargol'skiy (Leningradskiy Metallurgical Inst.). *Zavod. Lab.*, 1958, 29 (4), 397-401.—The use of a thermo-e.m.f. method for determining C in steels under works conditions is discussed.

G. S. SMITH

*Approved*  
*5*  
*no file*

AUTHORS: Nikitskaya, V. A. and Karpunin, A.M., Engineers.  
(Dzerzhinskiy Works).

368

TITLE: Longitudinal cracks on flanges of railway rails.  
(Prodol'nye treshchiny na flantsakh zheleznodorozhnykh  
rel'sov).

PERIODICAL: "Stal'" (Steel), 1957, No.4, pp.347-351 (U.S.S.R.)

ABSTRACT: An investigation of the causes of surface defects on the base and head of rails in the form of cracks and fissures stretched along the rolling direction, usually associated with surface bubbles, was investigated. It was found that the observed defects originate from longitudinal cracks often present on the bottom part of the ingots of rail steel. The appearance of the above cracks on ingots was co-related with the fluidity of steel and the rate of casting of ingots. With bottom pouring the optimum temperature is limited to a narrow temperature range 1470-1475, therefore, the real solution for the problem is top pouring. From top poured ingots the yield of the quality rails increases to 90-92%. There are two tables, 9 figures and 2 Russian references.

NIKITSKAYA, V

PLATE 1 BOOK EXTRACTS 807/1380

Special Issue: Metallurgy, Descriptive  
"Rolling" for the 25th anniversary program (Metallography in the Field for  
Technical Progress) (Moscow) Izdatel'stvo Mashinostroyeniya 1979 96 p. 3,200 copies  
printed.

Special Ed.: Ye. V. Lopatin, P. M. Korotkov, and I. B. Polubny. Ed.: S. A. Malozemov;  
Tech. Ed.: S. D. Shadrin.

NOTE: This book is intended for technical personnel interested in metallurgical  
processes.

CONTENTS: The book contains 9 articles dealing with technical improvements  
developed and implemented by workers at the Plant Lenin Dzerzhinsky,  
production of the machine-rolling mills (mechanical engineering company Dzerzhinskii  
[Dzerzhinskii] Machine-Building Plant, Dzerzhinskii). Descriptive articles  
describe technical progress in laminar rolling, continuous casting, thermal treatment  
of open-hearth processes, light rolling, and improvements in roll production.

Editor: M. Dzerzhinskii. Improving the Quality of Rails  
Ed.: Professor Szeel

Karpulin, A. [Engineer]. Best Treatment of Rails 38

Nikitina, I. [Engineer]. A New Steel for Rolling the Plate 37

Chelapov, A. [Manager of Best-Engineering Laboratory].  
Improvement in the Design of Reciprocating Rolling Pits 31

NOTE: Library of Congress (DT705.83)

108-769

82/000/000  
11-13-80



TYLKIN, M.A., kand.tekhn.nauk; NIKITSKAYA, V.A., inzh.; BURKHAN, G.N., inzh.

Efforts to avoid discards in rolled telegraph wire rods. Stal'  
21 no.5:448-451 My '61. (MIRA 14:5)

1. Dneprodzershinskiy metallurgicheskiy zavod-vtuz i zavod im.  
Dzerzhinskogo.

(Rolling (Metalwork)--Quality control)  
(Telegraph wire)

TYLKIN, M. A., kand. tekhn. nauk; GREBENIK, V. M., kand. tekhn. nauk;  
KUCHERENKO, V. F., inzh.; ALPEYEV, V. G., inzh.;  
NIKITSKAYA, V. A., inzh.

Heat treatment of crane wheels. Mashinostroenie no.5:57-60  
S-0 '62. (MIRA 16:1)

1. Dneprodzerzhinskiy metallurgicheskiy zavod-vtuz im. M. I.  
Arsenicheva (for Tykin, Grebenik, Kucherenko). 2. Metallur-  
gicheskiy zavod im. Dzerzhinskogo (for Alpeyev, Nikitskaya).

(Steel—Heat treatment)  
(Cranes, derricks, etc.)

NIKITSKAYA, V.

Research carried out at the Dzerzhinskii Metallurgical Plant.  
Stal' 22 no.12:1075, 1086-1087, 1105, 1122 D '62.

(MIRA 15:12)

(Dneprodzerzhinsk--Metallurgical research)

NIKITSKAYA, V.

Research at the Dzerzhinskii Metallurgical Plant. Stal'  
23 no.2:180 F '63. (MIRA 16:2)  
(Dneprodzerzhinsk—Blast furnaces)

W. R. RICHMOND, W. R.

Prose: The methods for underground waters. Vol. 1. (1954)  
173 (1954) (P. 1)

NIKITSKAYA, E. A.

Chemical Abst.  
Vol. 48 No. 8  
Apr. 25, 1954  
Petroleum, Lubricants, and Asphalt

②  
iNA  
A method of determining the threshold shear stress of mineral oils. P. I. Sanin and E. A. Nikitskaya. *Trudy Inst. Nefti, Akad. Nauk S.S.S.R.* 2: 47-52 (1952).—Static shear stress,  $\theta_s$ , computed from  $\theta_s = P_c d / 4l$  (where  $P_c$  is the crit. pressure at which motion of oil occurs,  $d$  = diam. of capillary in cm,  $l$  = length of column of oil in cm.) is independent, within broad limits, of rate of application of load, diam. of capillary, and length of oil column if the pressure accurately corresponding to the beginning of motion is taken as the crit. pressure. V. N. Bednarski

10-19-54  
JP

SOV/65-59-9-5/14

**AUTHORS:** Sanin, P. I; Sher, V. V. and Nikitskaya, Ye. A.

**TITLE:** Metal Dialkyl Dithiophosphates as Complex Additives to Lubricating Oils. (Dialkilditiofosfaty metallov kak kompleksnyye prisadki k smazochnym maslam).

**PERIODICAL:** Khimiya i Tekhnologiya Topliv i Masel, 1958, Nr.9. pp. 24 - 28. (USSR).

**ABSTRACT:** In early articles it was shown that metal dialkyl dithiophosphates are active complex additives (Ref.1 - 2). Dialkyl dithiophosphates of various metals have varying effect on the deterative and corrosion properties of oils. Tests were carried out on two types of oil: the oil MS-20 (from the Emba Region) and the oil MK-22 (from the Baku Region). Properties of these oils are given. From Table 1 it can be seen that these additives show varying degrees of activity. The most active additive was the barium dialkyl dithiophosphate DF-1 when added to the oil MS-20. This additive contained about 4% P, 9% S, and 8% barium, and was used in the form of a 50% solution in spindle oil AD. The action of this additive on the characteristics of various oils was investigated under laboratory conditions. Table 3: the dependence of the corrosion of oils on the concentration of DF-1. Results of this

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SOV/65-52-9-5/14

Metal Dialkyl Dithiophosphates as Complex Additives to Lubricating Oils.

Investigation indicates that the optimum concentration of the additive DF-1 is about 3%. Other tests concerned the effect of the additive on the oil MS-20 with regard to its stability to oxidation (GOST 4953-49), and its tendency to lacquer formation (GOST 6049-51) (Table 4). The acid number of the samples containing the additive, after testing in the device PZV, were considerably lower than for oils not containing the additive (Table 5). Practical experiments were carried out on the one-cylinder engine IT-2-3 (designed by VNII NP) under the supervision of V. F. Filippova. Results of these tests are given in Table 6. Table 7: the effect of the additive on the solidification point of the oils; Table 8: the effect of complex additives on some properties of the oil MS-20 (containing 3% of the additive). There are 8 Tables and 4 Soviet References.

ASSOCIATION: Institut nefti AN SSSR. (Petroleum Institute, AS USSR).

1. Lubricant additives--Effectiveness
2. Phosphates--Applications
3. Lubricating oils--Test results

Card 2/2



NIKITSKAYA, Ye. A.

5(3), 3(4)  
ARRIVED:

SOV/62-39-d-15/42  
Petrov, A. A., Sargiyenko, S. B., Zedilina, A. L.,  
Bodilya, E. A., Sasin, P. I., NIKITSKAYA, Ye. A.

TITLE: Synthesis and Properties of the Dimethyl-substituted-alkanes  
Having the Composition C<sub>12</sub>-C<sub>16</sub>

PERIODICAL: Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk,  
1959, Nr 6, pp 1421-1424 (USSR)

ABSTRACT: The present paper discusses the synthesis and properties of  
some of the compounds mentioned in the title. The properties  
of the synthesized materials are given in table 1. Nearly  
all substances crystallize at low temperatures; only 2,4-di-  
methyldecane and 3,5-dimethyldodecane vitrify at much lower  
temperatures than do their isomers or adjacent homologs.  
Besides references 3 investigations aiding at the separation  
of these phenomena have also been mentioned in the literature  
(Ref 4). It was assumed that the compounds mentioned in the title  
vitrify at low temperatures. Various investigations were carried out to prove  
this assumption (determination of viscosity as a function  
of temperature (Table 2) and determination of molecular  
weight). From the results it is seen that the influence of  
the structure on the vitrification effect cannot be limited.

Card 1/2

3

It was only possible to establish a certain dependence on  
the branching degree of the compounds. There are 2 tables  
and 3 Soviet references.

ASSOCIATION: Institut nefii khimii nauki SSSR  
(Petroleum Institute of the Academy of Sciences, USSR)

SUBMITTED: December 10, 1957

Card 2/2

NIKITSKAYA, YL. A.

S 3380 1105, 1158, 1161

2/06/60/000/010/012/018  
1019/8054

4458

ADDRESS:

Petrov, M. A., Gorkiyevko, S. B., Tselinina, A. L.,  
Krasovskiy, A. G., and Nikitskaya, Y. L.

TITLE: Synthesis and Properties of High-molecular Hydrocarbons of Elased Structures. Information 1. Synthesis of Hydrocarbons of the Compounds C<sub>24</sub>

PERIODICAL:

Izvestiya Akademi nauk SSSR. Otdeleniye Khimicheskikh  
nauk, 1960, No. 10, pp. 1648 - 1651

NOTE: The authors synthesized several hydrocarbons that, up to a certain extent, had served as models for the hydrocarbons containing in high-molecular weight compounds of the type of the hydrocarbons of the Elased and polymeric of 2) Hydrocarbons with aliphatic structures containing 24-merid ones by R. Saltselger et al. (Ref. 2), the present studies were made on a larger scale. The influence of the degree of cyclization of the hydrocarbon molecule, the effect of the relative position of some cycles in the paraffin chain of the molecules, and the effect of the  
and 1/3

degree of substitution of the aromatic of cyclohexane rings in the molecule upon the properties of the whole molecule were investigated (method of the structural changes). The hydrocarbons were synthesized by the Grignard reaction. The alcohols were dehydrated in the vapor phase by means of an aluminum catalyst used (method of the American Petroleum Institute); this was, however, done in vacuum (1-4 mm Hg). Purification was carried out by distillation and absorption. The conditions of synthesis are described in detail for 1,1-diphenyl ethane, 7-ethyl-2,2-hydrocarbons. Since a peculiar behavior of 7,8,6-trimethyl chloro benzyl was observed under the preparation conditions of the ordered fragments, the characteristics of the reaction between acetylated benzyl halides and magnesium are discussed (reaction between acetylated benzyl chloride and magnesium). The structure of the most important fragments of the 24 hydrocarbons obtained. The analyses of 1,1,2,2,4,4-hexachloro-2,3-dimethyl-2-butene and 1,1,2,2,4,4-hexachloro-2,3-dimethyl-2-butene derivatives are presented, i.e., the aromatic hydrocarbons having  
and 2/3

Several methyl groups on the ring have a much higher viscosity than the monosubstituted isomers. In the near future, the authors will publish a paper on the physico-chemical properties of the hydrocarbons described here (data on various spots). There are 7 tables and 10 references: 3 Soviet, 3 US, 1 German, and 1 British.

ASSOCIATION: Institut Geologii i Petrologii Gornichikh Institutov  
Akademii nauk SSSR (Institute of Geology and Geophysics  
of the Academy of Sciences of the USSR)

SUBMITTED: May 6, 1959

Card 3/3

SANIN, P.I.; PETROV, A.I.A.; SERGIYENKO, S.R.; NIKITSKAYA, Ye.A.

Viscous properties of some  $C_{24}$  cyclic hydrocarbons. Zhur.prikl.  
khim. 33 no.4:919-930 Ap '60. (MIRA 13:9)

1. Institut neftekhimicheskogo sinteza AN SSSR i Institut geologii  
i razrabotki goryuchikh iskopayemykh AN SSSR.  
(Hydrocarbons) (Viscosity)

67570

5.3300 (B)

5(3)

SOV/20-130-2-26/69

AUTHORS: Sanin, P. I., Petrov, Al. A., Sergiyenko, S. R., Academician  
AS Turkm SSR, Nikitskaya, Ye. A.

TITLE: Viscosity<sup>1</sup> Properties of Alkyl-aromatic Hydrocarbons and  
Their Hydrogenated Analogs

PERIODICAL: Doklady Akademii nauk SSSR, 1960, Vol 130, Nr 2, pp 338 - 340  
(USSR)

ABSTRACT: An investigation of the viscosity of aromatic hydrocarbons  
containing isolated benzene rings, and their hydrogenated  
analogs, showed (Table 1) that the change in viscosity on  
hydrogenation considerably depends on the structure of the  
hydrocarbons. Hydrogenation of certain structures reduces  
the viscosity extraordinarily. The aromatic hydrocarbons<sup>1</sup> 4  
(C<sub>24</sub>) investigated here may be divided into 2 groups: 1)  
without substituents on the ring (Table 1, Nrs 1-5); 2) with  
methyl groups on the ring (Nrs 6-8). Hydrogenation (or trans-  
formation of aromatic into naphthene hydrocarbons, respecti-  
vely) of the hydrocarbons of the 1st group increases the  
viscosity, and causes a higher viscosity increase with de-

Card 1/3

63570

Viscosity Properties of Alkyl-aromatic Hydrocarbons and SOV/20-130 2-26/69  
Their Hydrogenated Analogs

The data obtained by the authors make it possible to assert that the viscosity on hydrogenation of the higher-boiling petroleum fractions may also be reduced by the presence of polycyclic aromatic hydrocarbons with isolated benzene rings containing alkyl- (methyl-) groups on the ring. The cause of the viscosity change of some types of aromatic hydrocarbons on hydrogenation is unknown and must be investigated yet. There are 2 tables and 4 references, 3 of which are Soviet

ASSOCIATION: Institut neftekhimicheskogo sinteza Akademii nauk SSSR (Institute of Petroleum-chemical Synthesis of the Academy of Sciences, USSR) Institut geologii i razrabotki goryuchikh iskopayemykh Akademii nauk SSSR (Institute of Geology and the Working of Combustible Minerals of the Academy of Sciences, USSR)

SUBMITTED: September 22, 1959

Card 3/3

SANIN, P.I.; BAGRIY, Ye.I.; PETROV, Al.A.; NIKITSKAYA, Ye.A.; TSEDILINA, A.L.

Viscosity of  $C_{24}$  and  $C_{28}$  polycyclic hydrocarbons. Neftekhimiya 3  
no.6:835-844 N-D '63. (MIRA 17:3)

1. Institut neftekhimicheskogo sinteza AN SSSR im. A.V.Topchiyeva  
i Institut geologii i razrabotki goryuchikh iskopayemykh.

Nikitskaya, E. S.

"Sur la question de l'hydrolyse des chlorures de l'acétylène et de l'acétylène.  
by S. I. Lourje et collaborateurs C. J. Starobogotov et E. S. Nikitskaya. (p. 21)

SO: Journal of General Chemistry (Zhurnal Obshchey Khimii) 1941, 10, 21.

cd

**Syntheses in the amino-sulfone series.** I. *p*-Amino-phenyl dialkylaminosulfonyl sulfones. I. Kh. Feldman and E. S. Nikitskaya. *Zhur. Obshch. Khim.* (J. Gen. Chem.) 19, 134-42(1949). Refluxing 55 ml.  $PCl_5$  in 20 ml.  $(CH_2Cl)_2$  and 59.5 g.  $Et_3NCH_2CH_2OH$  in 15 ml.  $(CH_2Cl)_2$  6 hrs. gave, after concn., addn. of  $H_2O$ , filtration, and addn. of 50% NaOH, 80%  $Et_3NCH_2CH_2Cl$ . The *di Me analog* was made similarly in 85% yield. Addn. of 3 g.  $Et_3NCH_2CH_2Cl$  to 1.74 g.  $4-A-NHC_6H_4SO_2NH_2$  in 30 ml. re-

fluxing  $EtOH$ , heating 5 hrs., filtration, and washing gave  $(p-A-NHC_6H_4SO_2CH_2)_2$ , m. 265-7° (from  $AcOH$ ). Heating 21 g.  $p-CH_3NC_6H_4SH$  and 6.8 g. KOH in 150 ml.  $EtOH$  to boiling, and addn. of 13.3 g.  $Me_3NCH_2CH_2Cl$  gave, after filtration, concn., and diln., 68%  $4-O_2NC_6H_4SCH_2CH_2NMe_3$ , red oil ( $HCl$  salt, m. 215-17° (from  $EtOH$ )), reduction with H and Raney Ni in  $EtOH$  at room temp. gave 81% *aniso deriv.* (oil) ( $di-HCl$  salt, m. 222-4° (from  $EtOH$ )). (I) (15 g.) in 105 ml.  $AcOH$  treated with 21 ml. 28%  $H_2O_2$  at 70° and heated 2 hrs. at 75-80° gave  $4-O_2NC_6H_4SO_2CH_2CH_2NMe_3$ , m. 109-10° (from  $EtOH$ );  $HCl$  salt, m. 199-201°. This (2 g.) and 0.8 g.  $NH_4Cl$  in 50 ml.  $H_2O$  treated at 70° with 4.3 g.  $Fe$  powder and stirred 6 hrs. gave, upon extrn. with  $Me_2CO$ , 0.3 g. *unknown substance* (II), m. 100-71°, while  $Et_2O$  extrn. gave 41%  $4-H_1NC_6H_4SO_2CH_2CH_2NMe_3$ , m. 137-8°, also obtained in 95% yield by reduction with H and Raney Ni. A similar sequence of reactions, starting with  $Et_3NCH_2CH_2Cl$ , gave 77%  $4-O_2NC_6H_4SCH_2CH_2NEt_3$ , an oil, whose  $HCl$  salt, m. 172-4°, gave 94% of the only *4-NH\_2 analog* ( $di-HCl$  salt, m. 188-90°) by H reduction, while  $H_2O_2$  oxidation gave 72%  $4-O_2NC_6H_4SO_2CH_2CH_2NEt_3$ , m. 93-5° (from  $EtOH$ ) ( $HCl$  salt, m. 185-7°), which by  $Fe$  reduction gave some *4-NH\_2 analog*, m. 98-100° and 34% II, m. 171°. Raney Ni-H reduction gave 90% of the pure sulfone, m. 98-100°. Use of  $Et_3NCH_2CH_2Cl$  gave in turn 75% only  $4-O_2NC_6H_4SCH_2CH_2NEt_3$  ( $HCl$  salt, m. 146-8°), 85% *4-NH\_2 analog*, an oil ( $di-HCl$  salt, m. 101-3°), 79%  $4-O_2NC_6H_4SCH_2CH_2NEt_3$ , m. 57-9° ( $HCl$  salt, m. 191-3°), and 95% *4-NH\_2 analog*, m. 78-19° (from  $Et_2O$ -petr. ether). G. M. Kosolapoff



NIKITSKAYA, Ye. S.

USSR/Chemistry - Pharmaceuticals

Feb 52

"Synthesis of Aminosulfides and Aminosulfones. VIII. Synthesis of Aminoaryl- $\beta$ -Ketonosulfones and Their Derivatives," I. Kh. Fel'dman, Ye. S. Nikitskaya, All-Union Sci Res Chem-Phar Inst imeni S. Ordzhonikidze, Moscow

"Zhur Obshch Khim" Vol XXII, No 2, pp 278-285

Prepd 12 aminoaryl- $\beta$ -ketonosulfones and derivs not described in the literature. Studied acid properties of methylene group in 2 of these: p-acetylamino-phenylphenacylsulfone (I) and p-acetylamino-phenyl-p'-acetylaminophenacylsulfone (II). Found that (a) H atom of methylene group of I reacts with alkylhalides  
209725

USSR/Chemistry - Pharmaceuticals (Contd) Feb 52

to form monoalkyl compds, with aldehydes to form ethylene derivs, while H atom of methylene group of II under same conditions does not react with alkylhalides, but reacts with aldehydes to yield ethylene derivs; (b) ethylene derivs of I and II prepd by reaction with salicylic aldehyde (III) react with Br<sub>2</sub>, adding Br in nucleus of III to form dibromides; (c) hydrolysis of acetyl groups of ethylene derivs yields amino compds whose double bonds can not be hydrogenated at normal pressure and room temp or in presence of ether Raney Ni or Pt (according to Adams) catalysts.

209725

SHCHUKINA, M.M.: PERCHIN, G.M.: MAKHEVA, G.G.: SAZONOVA, YE. D.: MEYTSKAYA, YE. G.  
YAMINA, A.T.: YAKOVLEVA, A.T.

Tuberculosis

Isonicotinoylhydrazones and their antituberculous activity. 1975. ANTITUBERCULOSIS, 1:11.

Monthly List of Russian Accessions, Library of Congress, October 1, 1975, 1975, 1975

NIKITSKAYA, E.S.

Synthesis of  $\gamma$ -substituted pyridines. M. V. Rehtsov, E. S. Nikitskaya, and A. D. Yalova (S. Dvorkovskaya All-Union Chem. Res. Inst., Moscow). *Zhur. Obshchei Khim.*, 23, 1069-9 (1953); *cf. C. A.*, 48, 1632g. To 2.42 g. 4-formylpyridine (I) (prepd. by oxidation of 4-picoline) in PhMe. was added 10 ml. 50% NaHSO<sub>3</sub>, the mixt. cooled to 10°, stirred until a mush formed, the liquid was decanted off, and the solid mass rubbed with EtOH, yielding 7.3 g. Na. (1-pyridyl)hydroxyacetatesulfonate (II) mixed with NaHSO<sub>3</sub>. Crystn. from H<sub>2</sub>O gave needles of \*HN:Cl.

CH<sub>3</sub>C(Cl)(OH)(SO<sub>3</sub><sup>-</sup>)CH<sub>2</sub>CH (A), sol. in hot H<sub>2</sub>O, insol.

in org. solvents, does not melt. A (0.56 g.) treated with 2.25 ml. N NaOH followed by 50 ml. abs. EtOH gave 0.5% regenerated II, which with NaHSO<sub>3</sub> again gave A. Prolonged stirring in PhMe of 3.1 g. II with 2 Cl. g. H<sub>2</sub>N<sub>2</sub>SCH<sub>2</sub>SH in H<sub>2</sub>O at 70-80° gave a green ppt. m. 213-14° (crude), m. 213-17° (from EtOH), identical as I *flumetasarbone*, which with 10% HCl formed a yellow HCl salt, m. 238.3-2.5° (from 50% EtOH). Crude II (7.0 g.) heated 5 min. with 15 ml. H<sub>2</sub>O and 20 ml. 50% K<sub>2</sub>CO<sub>3</sub> developed a strong aldehydic odor, and after rapid extr. with CHCl<sub>3</sub>, the ext. gave 1.56 g. l. b. 125-2°, readily forming from it monohydrate, m. 53-60°; I forms a HCl salt, *hexahydrate*, m. 133-4°, which sublimes to white after loss of H<sub>2</sub>O; the *dry salt*, m. 159.5-81.5°, is again a white, converted to the monohydrate. Heating 3.1 g. IIHCl with 1.26 g. Cl<sub>2</sub>(CO)Et and 2.0 ml. AcOH 15 min. at 85-90° gave 3.8 g. 1-(2,2-dichloroethyl)pyridine-HCl (III), m. 210-21°, which with an equiv. of NaOAc yielded the free base, decamp. 253-6° (*isomeric salt*, losing H<sub>2</sub>O in vacuo at 100°). III (1 g.) hydrogenated in 1% HCl over PtO<sub>2</sub> at slight pressure gave 91% 1-(2,2-dichloroethyl)pyridine-HCl, m. 237-9°, yielding with AcONa the free base, b. comp. 235.5-6.0°. The HCl salt refluxed 4 hrs. with 4%

alc. HCl gave, after the usual treatment, 79.5% *nitro ester-HCl*, m. 122-3°. Letting 3.3 g. I.HCl stand 1 day with 3.53 g. Cl<sub>2</sub>(CO)Et, 9.6 ml. pyridine, and 3.3 ml. piperidine gave, after filtration of the piperidine HCl, evapn. of the filtrate at 50° in vacuo, dib. with hex. filtration, washing the solids with 15% Na<sub>2</sub>CO<sub>3</sub> and distn., 62% 7,7-(2,2-dichloroethyl)pyridine, b. 176-8°, alc. HCl gave the HCl salt, m. 180-2° (from EtOH). Hydrogenation gave the Et analog described previously. G. M. E.

NIKITSKAYA, YE. S.

USSR/Chemistry - Drugs

Sep 53

"Aminoalkyl Derivatives of Quinuclidine," M.V. Rubtsov, Ye.S. Nikitskaya, Ye.Ye. Mikhlina, A.D. Yanina, and V.Ya. Furshtatova, All-Union Sci-Research Chemico-Pharmaceut Inst im Ordzhonikidze

Zhur Obshch Khim, Vol 23, No 9, pp 1555-1559

A number of substituted 2-aminomethyl quinuclidines and 2-aminomethyl-3-( $\beta$ -aminoethyl)-quinuclidines were synthesized.

268133

NIKITSKAYA E.S.

Chemical Abst.  
Vol. 48 No. 8  
Apr. 25, 1957  
Organic Chemistry

Synthesis of 4-substituted derivatives of pyridine. M. V. Rubtsov, E. S. Nikitskaya, and A. D. Yudin (Institute of Organic Chemistry, Pharm. Inst., Moscow, U.S.S.R.; Dokl. Akad. Nauk S.S.S.R. 89, 91-2 (1954)).  
Direct oxidation of 4-picoline is possible in the presence of 4-picoline. 4-Formylpyridine (I) in PhMe with 60% NaHSO<sub>3</sub> yields a colorless solid mixt. of NaHSO<sub>3</sub> and Na  $\alpha$ -hydroxy-4-pyridinemethanesulfonate (II); crystn. of the mixt. from H<sub>2</sub>O yields Na-free needles of the inner salt of  $\alpha$ -hydroxy-4-pyridinemethanesulfonic acid, with evolution of Na<sub>2</sub>SO<sub>3</sub>; AcOH acts similarly. Heating II briefly with 85% K<sub>2</sub>CO<sub>3</sub> extn. with CHCl<sub>3</sub> or Et<sub>2</sub>O, and evapn. of the ext. gave an odoriferous liquid, b. 185-7°, which crystallizes in air; this is pure I; in contact with air it yields the hydrate, m. 58-60°. Much I is distd. along with the solvent (above), as can be detected by addn. of EtOH-HCl, which ppt. I.HCl, m. 132-4° (hemihydrate), m. 159.6-61.5° (anhyd. after drying by vacuum distn.). I yields a m. semicarbazone, yellow, m. 215-17° (from EtOH). I.HCl condenses with CH<sub>3</sub>(CO)H in AcOH at 85-90°, yielding 81%, 4-(2,2-dicarboxyvinyl)pyridine-HCl (III), m. 219-20°; w/ NaOAc it yields the free base, cong. 1.5H<sub>2</sub>O, m. 26-3°, which loses all the H<sub>2</sub>O *in vacuo* at 100°. I.HCl and Cl<sub>2</sub>-(CO<sub>2</sub>Et)<sub>2</sub> in pyridine and a slight excess of piperidine kept 4-5 days in the cold gave 62% 4-[2,2-bis(ethoxycarbonyl)methyl]pyridine (IV), b. 162-4°. III heated with EtOH.HCl loses CO<sub>2</sub>, yielding Et-4-pyridinacrylate. The free base of III reacts with SOCl<sub>2</sub> only at 70° with partial decarboxylation. Hydrogenation of III gave 4-(2,2-dicarboxyethyl)piperidine-HCl, m. 235-7°; free base, m. 353.5-6.0°, esterified to di-Et ester, whose HCl salt, m. 113-9°, is also obtained by hydrogenation of IV over Pt. (C. 54-6)

NIKITSKAYA - E. S.

Synthesis of  $\gamma$ -formylpyridine and isonicotinic acid. /  
M. V. Kuznetsov, E. S. Nikitskaya, and A. D. Yanina. J.  
Gen. Chem. U.S.S.R. 24, 1631-2 (1951) (Engl. translation).  
See C.A. 49, 11241h. B.M.B.

NIKITSKAYA - E.S.

CH Synthesis of substituted 2-aminoethylquinolines  
M. V. Rubtsov and E. S. Nikitskaya. *J. Gen. Chem.*  
*U.S.S.R.* 24, 1641-4 (1954) (Engl. translation).—*See C.A.*  
49, 13250c. B. M. P.

**NIKITSKAYA, E. S.**

USSR/Chemistry - Synthesis

Card 1/1 : Pub. 151 - 34/42

Authors : Rubtsov, M. V.; Nikitskaya, E. S.; and Yanina, A. D.

Title : Synthesis of gamma-formylpyridine and isonicotinic acid

Periodical : Zhur. ob. khim. 24/9, 1648-1651, Sep 1954

Abstract : The conditions favorable for the synthesis of gamma-formylpyridine by selective oxidation of gamma-picoline, with selenium dioxide in the presence of beta-picoline, are described. Two variants for the derivation of isonicotinic acid, with a yield of 75-80% of the initial gamma-picoline, were developed. The effect of selenium dioxide, on the selective oxidation of gamma-picoline, is explained. Five references: 2-USSR; 1-Swiss; 2-German and USA (1934-1953).

Institution : The S. Ordzhonikidze All-Union Scientific Research Chemical  
Pharmaceutical Institute

Submitted : April 14, 1954



**NIKITSKAYA, E. S.**

USSR/Chemistry - Synthesis

**Card** 1/1 : Pub. 151 - 36/42**Authors** : Rubtsov, M. V., and Nikitskaya, E. S.**Title** : Synthesis of substituted 2-aminomethylquinuclidine**Periodical** : Zhur. ob. khim. 24/9, 1659-1664, Sep 1954

**Abstract** : The synthesis of numerous 2-alkyl(aryl)aminomethylquinuclidines, from 2-quinuclidine carboxylic acid, is described. The derivation of 2-aminomethylquinuclidine containing the quinoline and acridine cycles through the reaction of 2-aminomethylquinuclidine with 6-methoxy-4-sulfoquinoline and 9-phenoxyacridine is reported. One USSR reference (1953).

**Institution** : The S. Ordzhonikidze All-Union Scientific Research Chemical Pharmaceutical Institute**Submitted** : January 13, 1954



NIKITSKAYA E.S.

*Chem* Synthesis of  $\beta$ -(2-quinolindinyl)propionic acid. M. V. Rubtsov, L. N. Vakhontov, and E. S. Nikitskaya. *J. Gen. Chem. U.S.S.R.* 25, 2281-3(1955)(Engl. translation). See *C.A.* 50, 9401c. B.M.R.

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NIKITSKAYA, E. S.

1000

✓ Synthesis of  $\beta$ -(2-quinuclidinyl)propionic acid. M. V. Rubtsov, L. M. Yakhontov, and E. S. Nikitskaya (S. Odeskenskaya All-Union Chem. Pharm. Research Inst., Moscow). *Zhur. Obshchey Khim.* 25, 2311-13 (1955).  
 Keeping 0.7 g. 2-formylquinuclidine, 3 ml. dry pyridine, and 5 drops piperidine with 0.8 g.  $\text{CH}_2\text{CO}_2\text{Et}$  4 days gave after extr. with  $\text{Et}_2\text{O}$  86% *di-Et* 2-quinuclidinylmethyl-*enemalonate*,  $\text{C}_{12}\text{H}_{20}\text{N}_2$ , *b.p.* 142-3°, *n<sub>D</sub><sup>20</sup>* 1.4821. Refluxed with concd.  $\text{HCl}$  6 hrs. it gave 80%  $\beta$ -(2-quinuclidinyl)acrylic acid-HCl, *m.* 197-8°, which hydrogenated over Raney Ni gave  $\beta$ -(2-quinuclidinyl)propionic acid-HCl, *decomp.* 215-16°, also formed from upon hydrolysis of the ester, *b.p.* 140-50°, obtained by condensation of 2-bromoethylquinuclidine-HBr with  $\text{NaCH}(\text{CO}_2\text{Et})_2$ . Treatment of the acid HCl salt with  $\text{SOCl}_2$ , followed by refluxing the crude acyl chloride with  $\text{EtOH}$  gave *Et*  $\beta$ -(2-quinuclidinyl)propionate, isolated as the methoxide, *m.* 87-8°, in 37% yield.  
 G. M. Krasovskii

Chem

EM

NIKITSKAYA, Ye. S.

Action of nitric acid on methylol derivatives of a ribose  
pyridine / M. V. Rubtsov, E. S. Nikitskaya, and V. S.  
Usorskaya. J. Gen. Chem. U.S.S.R. 25, 2341-3 (1965)  
(English translation) - See C.A. 50, 9401b. B. M. R.

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N. Kitskaya, E.S.

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✓ Action of nitric acid on methyl derivatives of 4-ethylpyridine. M. V. Kuznetsov, E. E. Zhuravskaya, and V. S. Usotskaya (S. Gubzonnikov ~~Department of Chemistry, Pharm. Sci. Research Inst., Moscow~~). Zhur. Obshch. Khim. 25: 2450-7 (1951). Heating 5.35 g. 4-ethylpyridine at reflux 18 hrs. with 18 g. 53% formalin, removal of excess reagent with steam and evapn. of the residual soln. to 12 ml. gave an aq. soln. of mixed methyl derivs. which can be used directly for oxidations or which can be sepd. as follows: after evapn. *in vacuo*, the residue was evapd. twice with abs. EtOH *in vacuo* at 100°, yielding 83.5% dimethyl-4-ethylpyridine, m. 93-5° (from EtOH), sepd. by washing with Et<sub>2</sub>O; the Et<sub>2</sub>O washings on evapn. gave 80.6% monomethyl-4-ethylpyridine, b.p. 41°. To the mixed methyl derivs. (12 ml.) at 90-100° there was added in 20-3 min. 50 g. 25% H<sub>2</sub>O<sub>2</sub> and 50.8 g. HNO<sub>3</sub> (92.8%); after subsidence of the exothermic reaction the mixt. was heated 5 hrs. at 100-110°; after cooling and neutralization there was obtained 53.0% isonicotinic acid and 0.85 g. apparently 4-(2,5-dihydroxy)pyridine, m. 55-7°, HCl salt, m. 142-50°; *recryst.*, m. 137°. Reduction of this in EtOH over Raney Ni at room temp. gave 91.8% 4-(2-aminoethyl)pyridine, b.p. 78-80°; di-HCl salt, m. 230-7°. 4-Acetylpyridine forms an oxime, which has *two modifications*, m. 121-3° (from CCl<sub>4</sub>, sol. in hot solvent), and m. 157-9° (from sol. in hot CCl<sub>4</sub>). Hydrogenation of 4-oxoethylpyridine oxime in EtOH over Pt gave 4-(2-aminoethyl)pyridine, identical with the above specimen.

*Handwritten initials*

*Handwritten initials*

НИКИТСКАЯ, Е. С.

USSR/Organic Chemistry - Synthetic Organic Chemistry, E-2

Abst Journal: Referat Zhur - Khimiya, No 19, 1956, 61543

Author: Rubtsov, M. V., Nikitskaya, Ye. S., Usovskaya, V. S.

Institution: None

Title: Alkamino Esters of Some Heterocyclic Acids as Possible Hypotensive Remedies

Original

Periodical: Zh. obshch. khimii, 1956, 26, No 1, 130-134

Abstract: There have been synthesized the diethylaminoethyl esters of dipicolinic (I), dipipecolinic (II), N-methyl dipipecolinic (III), 6-methyl picolinic (IV), 6-methyl pipecolinic (V), 1,6-cimethyl pipecolinic (VI), and quinuclidine carboxylic-2 acid (VII). On pharmacological investigation it was found that the di-methyl iodides of VI and VII have high ganglion-blocking activity. A mixture of 3 g dipicolinic acid (VIII) and 30 ml SOCl<sub>2</sub> is boiled until completely dissolved (6 hours) heat the thus formed di-acid chloride (IX) with 30 ml diethylaminoethanol (X) for 6 hours at

Card 1/3

USSR/Organic Chemistry - Synthetic Organic Chemistry, E-2

Abst Journal: Referat Zhur - Khimiya, No 19, 1956, 61543

Abstract: 90%, BP 103-105°/0.25 mm, and the ethyl ester of 6-methyl pipercolinic acid (XV), yield 92%, BP 99-100°/13 mm; hydrochloride MP 213-215°. Mixture of 4.27 g XIV, 1.32 g CH<sub>3</sub>I and 23 ml absolute alcohol heated for 6 hours at 40-45°, evaporated in vacuum, residue extracted with dry C<sub>6</sub>H<sub>6</sub>, the insoluble hydroiodide of XIV is filtered off and from the benzene extract is recovered the diethyl ester of N-methyl dipipercolinic acid (XVI), yield 52.7%, BP 107-108°/0.2 mm. Analogously is prepared the ethyl ester of 1,6-dimethyl pipercolinic acid, yield 43.7%, BP 53-54°/0.2 mm; hydrochloride MP 198-200°. In 7 ml of X are dissolved 0.01 g Na, added with stirring 1.32 g XVI, heated 3 hours at 150° (distilling off the alcohol) excess of X is distilled off, the residue is treated with 50% solution K<sub>2</sub>CO<sub>3</sub> and extracted with ether; III is thus obtained, yield 51.2%, BP 176-178°/0.2 mm; methyl iodide and hydrochloride are oily substances. Analogously is synthesized VI, yield 44.7%, BP 106-108°/0.25 mm; dimethyl iodide MP 201-202°.

Card 3/3





NIKITSKAYA, S.

4

Preparation of N-ethylpiperidine, M. V. Nubtsov and S. S. Nikitskaya (S. Ordzhonikidze All-Union Chem. Pharm. Res. Research Inst., Moscow). *Zhur. Priklad. Khim.* 29, 1887 (1976). Heating 40 g. C<sub>11</sub>H<sub>17</sub>N, 5 g. Raney Ni, and 200 ml. ahs. EtOH in autoclave with 50 atmo. H<sub>2</sub> to 140-60° 30 hrs. gave 55-60% N-ethylpiperidine; mp. 106-8°; Cf. Jones, C.A. 45, 6186. G. M. Kosolunoff

EM

AUTHORS: Nikitskaya, Ye. S., Rubtsov, M. V.

79-11-26/36

TITLE: Synthesis of Bicyclic Systems Starting From 2,6-Lutidine.  
Synthesis of 9-Methyl-2-Oxy-9-Azabicyclo (3,3,1)-Nonanes  
(Azobicyclic )  
(Sintez bitsiklicheskih sistem, iskhodya iz 2,6-lutidina)  
(Sintez 9-Metil- 2- oksi - 9 - azobitsiklo (3,3,1) - nonana).

PERIODICAL: Zhurnal Obshchey Khimii, 1957, Vol. 27, Nr 11,  
pp. 313-316 (USSR)

ABSTRACT: The investigation of the azobicyclic compounds of the octane series (quinuclidine, tropane) showed that they are of great interest as raw products for the synthesis of remedies. Thus compounds with curative, ganglion-blocking, spasmatic, mydriatic and other properties were discovered among the tropine derivatives. It was of interest to investigate the bicyclic systems close to the tropane series. Thus the authors synthesized 9-methyl-2-oxy-9-azobicyclo- (3,3,1)-nonane by starting from the ethyl ester of  $\alpha$ -methylpicolinic acid (obtained from 2,6-lutidine). (See the process of reaction). The initial, intermediate and final products are as follows: the ethyl ester of 6-methylpicolinic acid, the product of

Card 1/2

Synthesis of Bicyclic Systems Starting From 2,6-Lutidine. 79-11-46/56  
Synthesis of 9-Methyl-2-Oxy-9-Azabicyclo (3,3,1)-Nonanes (Azobicyclic)

its condensation with chiral, 2-carboxy-6- (1-carboxyvinyl)-pyridine, 2-carbethoxy-6- (p-carbethoxyethyl)-piperidine, 9-methyl-2-keto-9-azobicyclo (3,3,1)-nonane which on reduction with aluminumhydrate of lithium is converted to 9-methyl-2-oxy-9-azobicyclo (3,3,1)-nonane.  
There are 3 references, 1 of which is Slavic.

ASSOCIATION: All-Union Scientific Research Institute for Pharmaceutical Chemistry  
imeni S. Ordzhonikidze (Vsesoyuznyy nauchno - issledovatel'skiy khimiko - farmatsevticheskiy institut im. S. Ordzhonikidze).

SUBMITTED: November 27, 1976

AVAILABLE: Library of Congress

1. Cyclic compounds - Synthesis

Card 2/2

NIKITSKAYA, YE. S.

AUTHORS:

Nikitskaya, Ye. S., Naumova, V. B., *ibid.*, 1958, No. 7, 141-142

TITLE:

Tertiary Amines of Some Heterocyclic Compounds as Possible Means For Blocking Nerve Ganglia (Tretichnyye aminy nekotorykh geterotsiklov kak vozmozhnyye gipertensivnyye sredstva)

PERIODICAL:

Zhurnal Obshchey Khimii, 1958, Vol. 28, No. 1, 141-142 (USSR).

ABSTRACT:

The quaternary ammonia salts with their quaternary nitrogen were formerly considered the most important source of remedies for blocking ganglia. But the most recent investigations showed that this may also be the case with secondary and tertiary amines (reference 2). Thus the authors had already earlier found that e.g. the pertinent 2-diethylaminoethylaminomethylquinuclidine (formula (a)) possesses a high activity in the above-mentioned sense. As compounds of this type of activity are of great importance for treating hypertension it was expedient to synthesize simpler compounds of a similar type, namely that of the pyridine and piperidine series. By the conversion of the hydrochlorides or esters of picolinic and 6-methylpicolinic acid with different amines it was possible to produce the amides (I and II). In spite of

Card 1/2

Tertiary Amines of Some Heterocyclic Compounds as Possible  
Means For Blocking Nerve Ganglia.

79-1 31/57

indications in publications that no amines can be obtained from the amides of pyridinecarboxylic acids with the aid of the aluminum hydride of lithium the authors succeeded in converting most of the obtained amides to the amines (III) although the yield on that occasion was small and by-product occurred. The reduction of the amides of piperidinecarboxylic acids took place much better with good yields and easy isolation (IV). The pharmacological investigation of the pyridine and piperidine derivatives which was performed by I. M. Sharapov showed that 1,6-dimethyl-2-( $\beta$ -diethylaminoethyl-aminomethyl)-piperidine (IV d) possesses a high activity in the above-mentioned sense that it even ten times surpasses that of tetraethylammoniumiodide. There are 1 table and 6 references, 5 of which are Slavic.

ASSOCIATION. **All Union** Scientific Chemical-Pharmaceutical Institute imeni S. Ordzhonikidze (Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevticheskiy institut imeni S. Ordzhonikidze)

SUBMITTED: January 7, 1957

AVAILABLE: Library of Congress

Card 2/2 1. Chemistry 2. Cyclic compounds 3. Amides

AUTHORS: Nikitskaya, Ye. S., Mikhlina, Ye. Ye., SOV/79-28-10-32/60  
Yakhontov, L. N., Furshtatova, V. Ya.

TITLE: Synthesis of the Hydrazines and Hydrazones of Some Hetero-  
cyclic and Aromatic Acids (Sintez gidrazidov i gidrazonov  
nekotorykh geterotsiklicheskikh i aromaticsikh kislot)

PERIODICAL: Zhurnal obshchey khimii, 1968, Vol 28, Nr 10,  
pp 2786 - 2790 (USSR)

ABSTRACT: In earlier investigations (Ref 1) it was shown that  
the hydrazone of isonicotinic acid and its hydrazones  
develop an antitubercular activity. It was, therefore,  
of interest to the authors to synthesize the hydrazides  
and their derivatives of the pyridyl-4-acetic acid,  
 $\beta$ -(pyridyl-4)acrylic and  $\beta$ -(pyridic-4)-propionic acid,  
as these differ from the isonicotinoyl hydrazone only by the  
presence of one and more methyl groups between the  
pyridine nucleus and the hydrazone radical. Therefore  
it was desired to obtain hydrazides and hydrazones  
from acids of the piperidine and quinoline series  
in order to explain the effect of the mentioned cycles  
on the biological effect of these compounds and to

Card 1/3

Synthesis of the Hydrazines and Hydrazones of Some  
Heterocyclic and Aromatic Acids

SOV, 79-28-10-11, 1979

compare them in this respect with the similar compounds of the pyridine series. To this end the hydrazides of the following acids were synthesized: isonipecotinic, pyridyl-4-acetic-, piperidyl-4-acetic-,  $\beta$ -(pyridyl-4)propionic-,  $\beta$ -(piperidyl-4)propionic-,  $\beta$ -(pyridyl-4)-acrylic-, 6-methyl picolic- and  $\alpha$ -quinuclidine carboxylic acid. As the p-nitro-benzoic acid is closely related to the isonicotinic acid, its hydrazide and hydrazones were also synthesized to explain its structure and activity. The synthesis of the hydrazides was carried out by the reaction of the ethyl esters of the acids with hydrazine hydrate in alcohol solution (Refs 5,6) already earlier synthesized by the authors. The subsequent reaction of the hydrazides with various aldehydes lead to the hydrazones. The constants of the obtained products, analyses and yields are given in tables 1-4. The biological investigation of the antitubercular activity showed that the synthesized products are much less effective than the corresponding derivatives of isonicotinic acid. The data are in tables

Card 2, 3



Synthesis of the H<sub>2</sub> Irizines and Hybrids of Some  
Heterocyclic and Aromatic Acids

SV, 01-10-1967, 1

and 6 references, 3 of which are Soviet.

ASSOCIATION: Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevticheskiy institut imeni S.Ordzhonikidze (All-Union Scientific Chemopharmaceutical Research Institute imeni S.Ordzhonikidze)

SUBMITTED: September 28, 1967

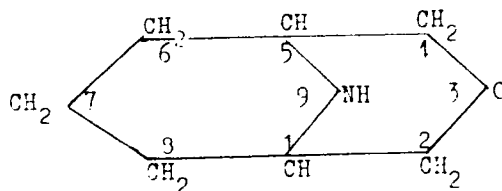
Card 3, 3

AUTHORS: Nikit'skaya, Ye. S., Usovskaya, V. S., SOV/79-29-1-28/74  
Kubtaov, M. V.

TITLE: Bicyclic Systems Derived From 2,6-Lutidine (Bitsiklicheskiye  
 sistemy na baze 2,6-lutidina)  
 II. Synthesis of the 3,9-Oxazabicyclo-[3,3,1]-Nonane and Its  
 N-Derivatives (II. Sintez 3,9-oksazabitsiklo-[3,3,1]-nonana  
 i yego N-proizvodnykh)

PERIODICAL: Zhurnal obshchey khimii, 1959, Vol 29, Nr 1, pp 124-129 (USSR)

ABSTRACT: In continuing work<sup>on</sup> the synthesis of the bicyclic systems derived  
 from 2,6-lutidine the authors obtained a new compound, the  
 3,9-oxazabicyclo-(3,3,1)-nonane



Card 1/3

The diethyl ester of the dipipecolinic acid, obtained from

Bicyclic Systems Derived From 2,6-lutidine.

SCV/79-29-1-28/74

II. Synthesis of the 3,9-Oxazabicyclo-[3,3,1]-Nonane and Its N-Derivatives

2,6-lutidine, was used as initial product (Ref 1). By the reduction of the ethyl ester of this acid with aluminum-lithium hydride in ether solution compound (I) was obtained which yielded (II) by methylation. By the action of thionyl chloride in the hydrochlorides of (I) and (II), (III) and (IV) were formed. On longer boiling of (I) with sulfuric acid (V) resulted, a slightly volatile, crystalline and salt-forming product (on nitrogen), from which some of its N-substituted derivatives were obtained. From compound (I) the nonane (VI) was formed by formic acid and formaldehyde. The sulfurization yielded the N-sulfo acid which was separated in the form of potassium salt (VII). By the reaction of (I) with the chloric acid anhydride of  $\beta$ -chloro propionic acid in alkaline medium with subsequent boiling of the resulting amide of this acid with piperidine and diethylamine the compounds (VIII) and (IX) were formed. By reduction of the amides obtained with aluminum-lithium hydride (X) and (XI) were synthesized. The reaction of an excess of (I) with dichloric acid anhydride of glutaric and adipic acid the diamides (XII) and (XIII) were obtained. The latter were transferred by reduction with aluminum-lithium

Card 2/3

Bicyclic Systems Derived From 2,6-Lutidine.

SCV/79-29-1-28/71

II. Synthesis of the 3,9-Oxazabicyclo-[3,3,1]-Nonane and Its N-Derivatives

hydride and subsequent treatment of the resulting amines with methyl iodide into the compounds (XIV) and (XV). Compounds (V) and (VI) show a nicotine-like activity, whereas compounds (VIII-XI) exert a lower activity. There are 2 references, 1 of which is Soviet.

ASSOCIATION: Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevticheskiy institut imeni S. Ordzhonikidze (All-Union Chemical-pharmaceutical Scientific Research Institute imeni S. Ordzhonikidze)

SUBMITTED: November 30, 1957

Card 3/3

SOV, '79-29-2-2

AUTHORS: Nikitskaya, Ye. S., Usovskaya, V. S., Rubtsov, M. V.

TITLE: Piperidine Derivatives as Possible Hypotensive Agents (Proizvodnyye piperidina kak vozmozhnyye gipotensivnyye sredstva)

PERIODICAL: Zhurnal obshchey khimii, 1959, Vol 29, Nr 2, pp 472-476 (USSR).

ABSTRACT: According to the sec tertiary amines of the quinolidine and piperidine series, which develop a high ganglion-blocking activity, the authors synthesized some N-substituted piperidine derivatives, in order to examine further tertiary amines (Ftivazid), served as initial product. The reaction of 2,6-lupetidine, a waste product in the preparation of piperidine (obtained from 2,6-lutidine) with the chloride anhydride of  $\beta$ -chloropropionic acid and subsequent boiling of the reaction product in ethyl alcohol with piperidine and diethylamine gave the compounds (I) and (II). By reduction, the latter correspondingly passed over to compounds (III) and (IV). Some out the synthesis, beginning from 2,6-lupetidine, of the sec quaternary salts by the aid of dichloric anhydride of glutaric and adipic acid, namely, compounds (V) and (VI). These

Card 1/2

SOV/79-29 2 25/71

Piperidine Derivatives as Possible Hypotensive Agents

piperidides of both acids could, correspondingly, be converted by reduction into 1,5-bis(2',6'-dimethyl piperidine-1'-yl)hexane (VII) and 1,6-bis(2',6'-dimethyl piperidine-1'-yl)hexane (VIII). Sec quaternary salts (Scheme 2) easily result from these compounds. By reaction of ethyl ester of 6-methyl piperidine-2-carboxylic acid with anhydride of 3-chloro propionic acid and by subsequent treatment of the reaction product with piperidine or diethyl amine, piperidines (IX and X) were obtained, which in their turn changed over to piperidones (XI and XIII) by reduction (Scheme 3). The constants of the compounds synthesized will be given in a following paper. There is a Soviet reference.

ASSOCIATION: Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsiyaevskiy institut imeni S. Ordzhonikidze (All-Union Scientific Chemical-pharmaceutical Research Institute imeni S. Ordzhonikidze).

SUBMITTED: January 3, 1958

Card 2/2

RUBTSOV, M.V.; NIKITSKAYA, Ye.S.; LANINA, A.D.; USOVSKAYA, V.S.

New ganglion blocking preparations. Khim. i med. no.15:16-22 '60.  
(MIrA 15:1)

1. Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevticheskiy  
institut imeni S. Ordzhonikidze.  
(AUTONOMIC DRUGS)

5.3610

77375  
SOV/79-30-1-35/70

AUTHORS: Nikitskaya, Ye. S., Usovskaya, V. S., Rabtsov,  
M. V.

TITLE: Bicyclic Systems Based on 2,6 Lutidine. III.  
N-Derivatives of 3-Oxa-9-azabicyclo-(3,3,1)-Nonane

PERIODICAL: Zhurnal obshchey khimii, 1960, Vol 30, Nr 1, pp  
171-182 (USSR)

ABSTRACT: Acyl and alkyl derivatives of 3-oxa-9-azabicyclo-  
(3,3,1)-nonane (I) were synthesized. Acid chlorides of  
acetic, propionic, and benzoic acids were reacted with I  
in anhydrous benzene with cooling and 9-acetyl-  
(IIa), 9-propionyl- (IIb), and benzoyl-3-oxa-9-aza-  
bicyclo-(3,3,1)- nonanes (IIc) were obtained. The ob-  
tained products, on reduction with lithium aluminum  
hydride, were converted into corresponding amines.  
Morpholine and dimethylamine in anhydrous alcohol,  
phenothiazine in anhydrous benzene, and the sodium  
salt of quinoxaline-4 in anhydrous alcohol were

Card 1/10

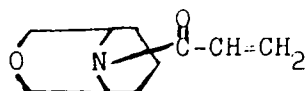


Bicyclic Systems Based on 2,6-Lutidine. III

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SOV/79-30-1-36/78

reacted with 9-( $\beta$ -chloropropionyl)-3-oxa-9-azabicyclo-(3,3,1)-nonane and corresponding  $\beta$ -substituted derivatives of 9-propionyl-3-oxa-9-azabicyclo-(3,3,1)-nonanes (IIId, IIe, IIIf, IIg) were obtained. The above reaction with phenothiazine and quinoxalone takes place with formation of a sideproduct, 9-acryloyl-3-oxa-9-azabicyclo-(3,3,1)-nonane.

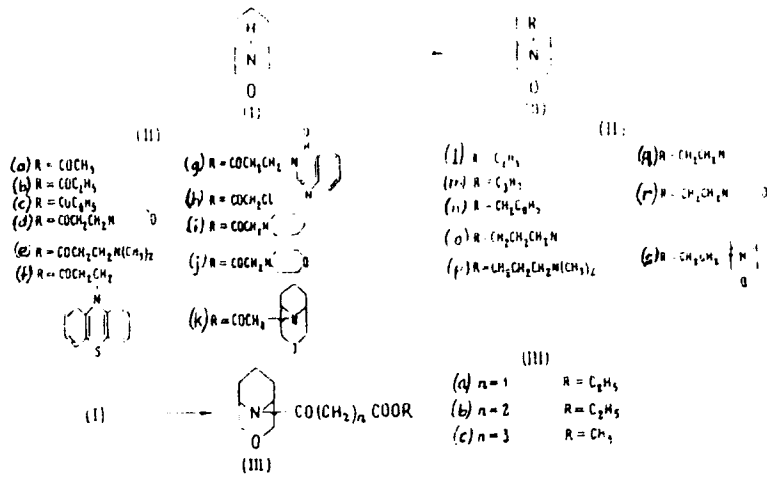


Acetyl chloride reacts with I, in aqueous alkali, forming as main product 9- [3'-oxa-9'-azabicyclo-3', 3', 1'-nonano-9'] -acetyl-3-oxa-9-azabicyclo-(3,3,1)-nonane (IIj).

Card 2/10

Biogel: Systems Based on 2,6-Pyridinedione. III

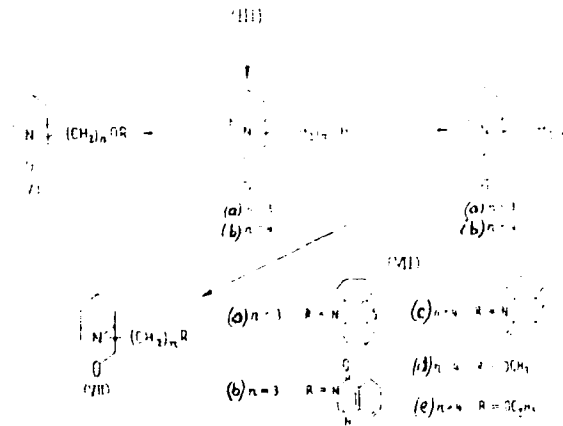
11/1/77  
SOV/77-3-1-1-1/77



Card 3/10

Biopolymers: Systems Based on  $\alpha, \omega$ -Dialkylamines. III

7757  
301/100-1-2-11



Card 4/10

Bicyclic Systems Based on 2,6-Lutidine. III

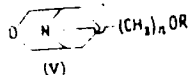
7737:  
SCV/77-30-1-33/75

The corresponding amines (IIo, IIp, IIr, IIs, IIt) were obtained on reduction of IID, IIe, III, IIJ, IIK, with lithium aluminum hydride. Attempts to reduce compounds IIf and IIg were unsuccessful. The desired amines were prepared as follows: I was reacted with carbethoxyacetyl chloride. The obtained IIIa was reduced to IVa; the latter with thionyl chloride gave VIa. Phenothiazine and quinoxalin-4-one were reacted with VIa; corresponding VIIa and VIIb were obtained. IIIb and IIIc were obtained similarly from  $\beta$ -carbomethoxypropionyl chloride and  $\beta$ -carbomethoxypropionyl chloride, forming on reduction IVb. Thionyl chloride was reacted with IVb and a corresponding hydrochloride (VIb) was obtained. Phenothiazine reacts with VIb, forming VIIc (yield 34%). Alkoxides react with VIb, forming corresponding ethers. VIId and VIIe were obtained by the above reaction.

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Bicyclic Systems Based on 2,6-Lutidine. III

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SOV/79-30-1-36/78



n	R	REACTION TIME (HR)	REACTION TEMPERATURE	YIELD (%)	BOILING POINT (PRESSURE IN MM)	MELTING POINT OF HYDROCHLORIDE
3	COCH <sub>3</sub>	4	On boiling	67	—	200—202°
3	COC <sub>2</sub> H <sub>5</sub>	4	On boiling	58	—	170—172
3	COC <sub>6</sub> H <sub>5</sub>	4	On boiling	80	—	189—191
3*		3	60—70°	59	183.5° (0.9)	179—181
3**		1	45—50	72	183 (1)	150—152

(Continuation, and explanation of asterisks, on next card)



Card 6/10

Bicyclic Systems Based on 2,6-Lutidine. III

77375

30/7/77-53-1-50/77

(table cont'd)

n	R	REACTION TIME (HR)	REACTION TEMPERATURE	YIELD (%)	BOILING POINT (PRESSURE IN MM)	MELTING POINT OF HYDROCHLORIDE
4	COCH <sub>3</sub>	4	On boiling	95		201-202
4	COC <sub>2</sub> H <sub>5</sub>	4	On boiling	~100		191-196
4	COC <sub>6</sub> H <sub>5</sub>	4	On boiling	87		194-195.5
4*		2	60	67	200-201 (0.8)	137-139
4**		2	60	60	184 (0.9)	152-151

\* Was isolated in the form of dihydrochloride.

\*\* Was isolated in the form of dihydrochloride monohydrate.

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Bicyclic Systems Based on 2,6-Lutidine. III

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SOV/79-30-1-36/78

The yields and properties of compounds are given below:

Compound	Yield (%)	bp (°C) (Pressure in mm)	mp (°C)
IIa	70	106-109/1	74-75
IIb	60	113-114/0.6	-
IIc	81	162-163/0.7	78-80
IIc	72	183-185/0.2	-
IIe	75	140/0.8	68-70
II f (1st fraction)	~30	101-103	-
II f (2nd fraction)	56	260	-
IIg	27	-	138-139
IIh	78	124-126/0.5	77-79
IIi	83	157-159/0.55	97-99
IIj	90	148-150/0.4	100-102
IIk	43	-	140-142
III	81	67-67.5/3	-
IIl	64	55-56/0.8	-
II n	93	119-121/0.7	38-40

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Bicyclic Systems Based on 2,6-Lutidine. III 77375  
SOV/79-30-1-36/78

(Continued from Card 8/10.)

The yield and properties of compounds are given below:

Compound	Yield (%)	bp (°C) (Pressure in mm)	mp (°C)
IIo	72	140-142/0.6	-
IIp	62	98-100/0.6	-
IIq	79	108/0.35	-
IIr	70	118-120/0.3	-
IIs	84	-	113-115
IIIa	77	157-159/0.7	-
IIIb	55	151-152/0.5	-
IIIc	77	171-172/1	63-65
IVa	65	107-109/0.5	-
IVb	70	135-137/1	-
VIa	75	217-219 (dec) .	-
VIIb	80	-	173-175
VIIa	41	-	234-236 (alc)
VIIb	52	215/0.4	-
VIIc	34	-	194-196
VIIId	-	-	163-165
Card 9/10			



Bicyclic Systems Based on 2,6-Lutidine. III

77375

SOV/79-30-1-36/78

VIIe

(Continued from card 9/10.)

64

176-177

There is 1 table; and 1 Soviet reference.

ASSOCIATION: Ordzhonikidze All-State Scientific Research Chemical-  
Pharmaceutical Institute (Vsesoyuznyy nauchno-  
issledovatel'skiy khimiko-farmatsevticheskiy institut  
imeni S. Ordzhonikidze)

SUBMITTED: January 21, 1959

Card 10/10

NIKITSKAYA, Ye.S.; USOVSKAYA, V.S.; RUBTSOV, M.V.

Bicyclic compounds based on 2,6-lutidine. Part 4: 3-Substituted  
derivatives of 9-methyl-3,9-diazabicyclo [3.3.1]nonane. Zhur.ob.  
khim. 30 no.10:3306-3315 0 '61. (MIRA 14:4)  
(Diazabicyclononane)

NIKITSKAYA, Ye.S.; USOVSKAYA, V.S.; RUBTSOV, M.V.

Bicyclic systems based on 2, 6-lutidine. Part 5: Biquaternary salts of  $\alpha, \omega$ -bis[9-methyl-3, 9-diazabicyclo (3, 3, 1)-nonano-3]-alkanes. Zhur.ob.khim. 31 no.10:3202-3205 0 '61. (MIRA 14:10)

1. Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevticheskiy institut imeni S.Ordzhonikidze.  
(Lutidine) (Paraffins)

NIKITSKAYA, Ye.S.; USOVSKAYA, V.S.; RUBTSOV, M.V.

Bicyclic systems on the basis of 2,6-lutidine. Part 6: Synthesis of 3,9-diazabicyclo [3,3]nonane. Zhur.ob.khim. 32 no.9:2886-2888 S '62. (MIRA 15:9)

1. Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevticheskiy institut imeni S. Ordzhonikidze. (Bicyclononane)

NIKITSKAYA, Ye.S.; USOVSKAYA, V.S.; RUBTSOV, M.V.

Bicyclic systems on the basis of 2,6-lutidine.  
Part 7: Interaction of alkyl (aryl) magnesium  
halides with benzylimide of N-methyldipicolic  
acid. Zhur.ob.khim. 32 no.11:3687-3693 N '62.(MIRA 15:11)

1. Vsesoyuznyy nauchno-issledovatel'skiy ~~institut~~  
farmatsevticheskiy institut imeni S. Ordzhonikidze.  
(Pipicolic acid)  
(Magnesium organic compounds)

NIKITSKAYA, Ye.S.; LEVKUYEVA, Ye.I.; USOVSKAYA, V.S.; RUBTSOV, M.V.

Synthesis of 7-hydroxy-9-oxo-1,4-dihydrocyclo [1,3,2] s-triazine  
and some of its derivatives. Zhur. org. khim. 1 no.1:174-18. Ja  
165. (MIRA 19:6)

I. Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevticheskiy institut  
imeni S.Ordzhonikidze.

5(1) 25(5)  
AUTHORS:

06218  
SOV/64-59-6-10/28  
Atroshchenko, V. I., Doctor of Technical Sciences, Asnin, Ya. I., Candidate of Technical Sciences,  
Vilesov, G. I., Nikitskaya, Z. A., Rabin, P. S.

TITLE:

Removal of Salt From Industrial Condensates of Nitrogen  
Fertilizer Enterprises by Means of Ion Exchange Resins

PERIODICAL:

Khimicheskaya promyshlennost', 1959, Nr 6, pp 499 - 501  
(USSR)

ABSTRACT:

The vapor condensate of the evaporators used in the nitrogen fertilizer industry is contaminated with  $NH_4^+$  and  $NO_3^-$  ions and has to be purified prior to its further use (as a steam boiler feed). Experiments carried out under the supervision of B. D. Bryanskiy (deceased) showed that by means of ion exchange resins it is not only possible to remove salt from the condensate but to re-use the ammonium nitrate obtained if the cation exchanger is regenerated with nitric acid and the anion exchanger with an ammonia solution. Among the investigated cation exchangers the type KU-2 proved to be best; in this case the regeneration takes place by means of a

Card 1/2

ATROSHCHENKO, V.I., doktor tekhnicheskikh nauk; ASMIN, Ya.I., kand.tekhn.  
nauk; NIKITSKAYA, Z.A.

Investigation of the stability of KU-2 and AB-2F ion exchangers  
used in the filtration of concentrated solutions. Khim. prom.  
no. 7:551-553 O-N '60. (MIRA 13:12)  
(Ion exchange)



ATROSCHENKO, V.I., doktor tekhn.nauk; ASNIN, Ya.I., kand.tekhn.nauk;  
NIKITSKAYA, Z.A.

Investigating the possibility of the repeated use of a part of wash  
waters in desalting units. Khim.prom. no.1:66-68 Ja '61.

(MIRA 14:1)

(Saline waters—Demineralization)

*NIKOLSKY, N. I.*

PHASE I BOOK EXPLOTTATION

SOV/4488

Akademiya nauk SSSR. Energeticheskiy institut

Goreniye pri ponizhennykh davleniyakh i nekotoryye voprosy stabilizatsii plameni v odnofaznykh i dvukhfaznykh sistemakh (Combustion at Reduced Pressures and Certain Problems in the Stabilization of the Flame in Single-Phase and Two-Phase Systems) Moscow, 1960. 85 p. Errata slip inserted. 5,000 copies printed.

Sponsoring Agency: Akademiya nauk SSSR. Energeticheskiy institut imeni G. M. Krzhizhanovskogo.

Resp. Ed.: L. N. Khitrin; Ed. of Publishing House: Ye. N. Grigor'yev; Tech. Ed.: V. N. Karpov.

PURPOSE: This book is intended for scientists engaged in combustion research.

COVERAGE: The book contains five reports delivered at the Obsnchemoskovskiy seminar po goreniyu (Moscow General Seminar on Combustion) in 1958. The problems discussed in these reports concern the effect of reduced pressure on the ignition and combustion of a stream of gas-vapor mixture in turbulent flow. Each report is followed by Soviet and other references.

Card 1/6

Combustion at Reduced Pressures (Cont.)

SOV/4488

## TABLE OF CONTENTS:

## Doroshenko, V. Ye., and A. I. Nikitskiy. Study of the Effect of Mixture Parameters on Turbulent Combustion Process Characteristics 3

This study presents experimental data relating to the effect of pressure (600-60 mm Hg) and temperature (100-300°C) on the turbulent combustion process of a homogeneous gasoline-air mixture. The data lead to the following conclusions: 1) A drop in the pressure and temperature of the mixture results in considerable deterioration of combustion process characteristics (decrease in flame-propagation velocity and increase in combustion-zone width). A change in pressure substantially affects both the flame-propagation velocity and the combustion-zone width. A change in mixture temperature, however, slightly affects the flame-propagation velocity and greatly affects the combustion-zone width. These regularities are explained from the standpoint of K. I. Shchelkin's theory when turbulence loss behind grids, as well as the effect of temperature and pressure on the characteristics of turbulent flow and normal flame-propagation velocity, are taken into account. 2) Decrease in turbulence intensity and increase in turbulence rate are the main reasons for the deterioration of the characteristics of the turbulent combustion process when pressure drops.

Card 2/6

2\*323

S/124/61/000/004/022/033

A005/A126

11.7200

AUTHORS: Doroshenko, V. Ye., Nikitskiy, A. I.

TITLE: Investigation of the influence of mixture parameters on the characteristics of a turbulent burning process

PERIODICAL: Referativnyy zhurnal, Mekhanika, no. 4, 1961, 84 - 85, abstract 4 B 579 (V sb.: Goreniye pri ponizhennykh davleniyakh i nekotoryye vopr. stabilizatsii plameni v odnofazn. i dvukhfazn. sistemakh. Moscow, AN SSSR, 1960, 3 - 23)

TEXT: The authors present results of an experimental study of the effect of pressure and temperature on the propagation rate of a turbulent flame and the width of the burning zone at turbulent combustion of a homogeneous fuel-air mixture. The open steady flame in a benzene-air mixture emitted from a round nozzle was investigated. The mean flame propagation rate  $\bar{U}_T$  was determined from the correlation

$$\bar{U}_T = \frac{F}{S} U,$$

where F is the nozzle area, S is the area of inner flame cone, U is the mean mix-

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2<sup>o</sup>323

S/124/61/000/004/022/033

A005/A126

Investigation of the influence of...

ture rate. The profiles of the inner cone were determined by measuring the temperature in the flame cross sections by thermocouples. The width of the burning zone  $\delta_T$  was determined by measuring the temperature over the flame axis. The turbulence intensity of the flow was measured by an electrothermoanemometer. The turbulence intensity was varied by means of a disturbing grid. The experiments were conducted within a pressure range of from 600 to 60 mm Hg and a temperature range of from 100° to 300°C. A decrease in pressure and temperature of the mixture led to a marked deterioration of the burning characteristics (decrease of the flame propagation rate as expressed by  $U_T \sim p^{0.5}$ ;  $U_T \sim T$ ; increase of the width of the burning zone as expressed by  $\delta_T \sim p^{0.5}$ ;  $\lambda_T \sim F^{-1.6}$ ). The authors showed that a decrease in turbulence intensity and increase in turbulence rate are the main causes for the deterioration of the burning characteristics. With burning processes behind stabilizing devices, the turbulence attenuation behind the stabilizers extends along the length of the flame tongue. There are 11 references.

V. Librovich

[Abstracter's note: Complete translation.]

Card 2/2

BRODSKIY, V.B.; NIKITSKIY, A.N.; PAVSHUK, I.S.

Compensating the drift of electrical length of cables in the UKVUM  
instrument. Priborostroenie no. 5:7-8 My '57. (MLRA 10:6)  
(Electric circuits) (Measuring instruments)

NIKITSKIY, A.S., inzh.

Device for protecting tower cranes against wind loads. Mekh.  
stroil. 17 no.3:22-23 Mr '60. (MIRA 13:6)  
(Cranes, derricks, etc.) (Wind pressure)

NIKITSKIY, A. S., inzh.

Automatic grab. Mekh. stroi. 17 no.9:26-27 S '60.

(MIRA 13:9)

(Cranes, derricks, etc.--Equipment and supplies)



NIKITSKIY, Al'bert Sergeevich; KASHTANOV, F., red.; NOVIKOVA, V.,  
tekhn. red.

[Hardeners for concretes and mortars] Uskoriteli tverdenia  
betonov i rastvorov. Minsk, Gosizdat BSSR, 1962. 40 p.  
(MIRA 15:10)

(Concrete)

(Mortar)

NIKITSKIY, A. Ye., Glavgeofizika (probably Glavnoye Geofizicheskoye Upravleniye, Main Geophysical Administration)

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