

DOROKHOV, V.N.; RUBAKHIN, V.N.; BIL'GIL'DEYEV, A.S.; DOMANOVSKIY, A.V.

Use of synthetic oils and fatty acids for oil-coating of rabbit
pelts. Kosh.-obuv.prom. 2 no.5:15-17 My '60. (MIRA 13:9)
(Hides and skins)

SOLOV'YEV, D.A.; DOROKHOV, V.N.

International Fall Fair in Leipzig. Kozh.-otuv.prom. 3 no.6:
30-31 Ja '61. (MIRA 14:8)

1. Direktor pushno-mekhovoy kontory Vsesoyuznogo ob'yedineniya "Soyuzpushnina" (for Solov'yev) 2. Glavnyy inzhener Leningradskoy mekhovoy fabriki (for Dorokhov).
(Germany, East--Fur industry) (Leipzig--Exhibitions)

DOROKHOV, V.N.; CHERNOMORDIK, L.M.

New developments in the equipment of Leningrad Fur Factory No.1.
Kozh.-obuv.prom. 4 no.2:17-20 F '62. (MIRA 15:4)
(Leningrad--Fur industry)

DOROKHOV, V.V.

USSR/Nuclear Physics - Structure and Properties of Nuclei.

C-4

Abs Jour : Ref Zhur - Fizika, No 4, 1957, 8706

Author : Demirkhanov, R.A., Gutkin, T.I., Dorokhov, V.V.,
Rudenko, A.D.

Inst :

Title : Masses of Isotopes H, D, He⁴ and C¹²

Orig Pub : Atom. energiya, 1956, No 2, 21-27

Abstract : A new exact measurement of the masses of the atoms H, D, He⁴ and C¹² has been made. The measurements were carried out with a mass spectrograph developed by Ardennet with the participation of Eger and the authors of this work. The apparatus has double focusing by means of electric and magnetic fields, effected respectively by cylindrical capacitor and a sector magnet. The ion beam is created by a plasma source with single contraction of the discharge. Recording of the mass spectrum is photographic; "Schumann" plates are used. There

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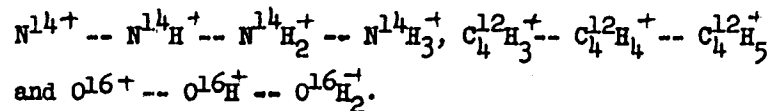
USSR/Nuclear Physics - Structure and Properties of Nuclei.

C-4

Abs Jour : Ref Zhur - Fizika, No 4, 1957, 8706

is the possibility of visually observing the spectrum with the aid of an ion-optical converter.

The maximum resolution of the instrument is 100,000 -- 120,000, and the dispersion (roughly calculated) is not less than 2.34 cm percent of relative mass difference. The calibration of the scale of the masses of the instrument is made using photographs of certain basic lines, namely those of the groups



The masses of the atoms H, D, He⁴ and C¹² were found from photographs of the doublets H₂ -- D, D₂ -- H⁴, D₃ -- 1/2

C¹²H₄ -- O¹⁶. The results are:

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USSR/Nuclear Physics - Structure and Properties of Nuclei.

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Abs Jour : Ref Zhur - Fizika, No 4, 1957, 8706

H -- 1.008142 ± 10^{-6} , He₄ -- $4.003872 \pm 4 \times 10^{-6}$,

D -- $2.014736 \pm 2 \times 10^{-6}$, and C¹² -- $12.003620 \pm 5 \times 10^{-6}$.

The data obtained are in agreement with the values determined from the energy balance of the nuclear reactions.

Card 3/3

MASS OF THE H, D, He, AND C ISOTOPES. E. A. Dimirkhanov, I. I. Gukin, V. V. Dorokhey, and A. D. Widenko. *Soviet Atomic Energy*, No. 1, 103-9 (1956).

Results are given of the measurements of the masses of the H, D, He, and C isotopes carried out by means of a mass spectrometer with a resolving power of 70,000 to 100,000. The data obtained agree well with the corresponding mass values obtained from the energy balance of nuclear reactions. (auth)

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DOROKHOV, V.V.

USSR/Nuclear Physics - Structure and Properties of Nuclei.

C-4

Abs Jour : Ref Zhur - Fizika, No 1, 1958, 452

Author : Demirkhanov, R.A., Gutkin, I.I., Dorokhov, V.V.

Inst : -

Title : Mass of the Isotope He³.

Orig Pub : Atomn. energiya, 1957, 2, No 5, 469-470

Abstract : A mass-spectroscopic determination was made of the mass of the isotope He³, in a mixture of helium isotopes enriched with He³ to 99.5%, using a setup previously described (Referat Zhur Fizika, 1957, No 4, 8706). The mass was measured in the doublets H³ -- He³ and HD³ -- He³. The results of the measurements were checked against the HD -- H³ doublet. The mass scale was calibrated against the spectrum N¹⁴ H -- N¹⁴ H₂ -- N¹⁴ H₃. The value obtained for the mass of He³ is 3.016970 ± 2 atomic units of mass. The data of this

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DOROKHOV, V.V.

USSR/Nuclear Physics - Structure and Properties of Nuclei

C-4

Abs Jour : Ref Zhur - Fizika, No 1, 1958, 453
Author : Demirkhanov, R.A., Gutkin, I.N., Dorokhov, V.V.
Inst : -
Title : Masses of the Isotopes C¹³, N¹⁴, and N¹⁵.
Orig Pub : Atomm. energiya, 1957, 2, No 6, 544-551

Abstract : Results are reported on new mass-spectrographic measurements of the masses of C¹³, N¹⁴, and N¹⁵. It is shown that there exists "an internal agreement" for the values of the masses of these isotopes, obtained from various systems of doublets. The measurements were performed under conditions that exclude the systematic errors. A procedure is given for a precision adjustment of the ion-optical system. For the masses of C¹³, N¹⁴, and N¹⁵, the values obtained were $13.007491 \pm 3 \times 10^{-6}$, $14.007527 \pm 4 \times 10^{-6}$ and $15.004890 \pm 5 \times 10^{-6}$ atomic units of mass respectively, which is in good agreement with the values

Card 1/2

24(5)

AUTHORS:

Demirkhanov, R. A., Gutkin, T. I.,
Dorokhov, V. V.

SOV/56-35-4-13/52

TITLE:

Nuclear Bond Energy in the Region of the 82 Proton and 126
Neutron Magic Numbers (Energiya svyazi yader
v oblasti magichekikh chisel po protonam 82 i neytronam
126)

PERIODICAL:

Zhurnal eksperimental'noy i teoreticheskoy fiziki, 1958, v. 35, No. 4
pp 917 - 925 (USSR)

ABSTRACT:

The authors of the present paper report on rich ex-
perimental material which is arranged in a clear manner
in tables. By means of a mass spectrograph (resolving
power 60000-80000, description in references 4,5)
the masses of the following isotopes were measured:
Lead: Pb 204, 206, 207 and 208 (Table 2)
Mercury: Hg 198, 199, 200, 201, 202 and 204 (Table 3)
Thallium: Tl 203 and 205 (Table 4)
Bismuth: Bi 209.
Determination of masses was carried out by direct
comparison with the masses of the corresponding

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Nuclear Bond Energy in the Region of the 82 Proton and
126 Neutron Magic Numbers

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organic compounds, as e.g. for
Pb 204 - $C_{16}H_{12}$, Pb 208 - $C_{14}H_8O_2$, Hg 199 - $C_{13}H_{11}O$,
Hg 204 - $C_{16}H_{12}$, Tl 203 - $C_{16}H_{11}$, Tl 205 - $C_{16}H_{13}$,
Bi 209 - $C_{14}H_{13}N_2$. The masses for the various isotopes
are given up to 6 decimals, data given are accurate
up to 3-4 decimals. From mass measurements the nucleon
bond energies in the nucleus are calculated. The results
obtained indicate a shell structure of the nucleus
with a well-filled shell of 82 protons and 126 neutrons.
The difference of the nuclear bond energy for an even
and odd number of nucleons in the nucleus and its
smoothing out as the shell is filled up can distinctly
be seen. After the shell is filled up with $Z=82$
and $N=126$, the bond energy of the next neutron is
higher than that of the next proton. The energy of
two bound neutrons (which yields the Hg 204 nucleus)
is greater than the energy of attachment of two
protons in the formation of the Pb 204 nucleus. The authors
thank Ye.Ye.Baroni, T.N.Lebsadze, K. A.

Card 2/3

Nuclear Bond Energy in the Region of the 82 Proton and 126 Neutron Magic Numbers SOV/56-35-4-13/52

Kovyrzina and V.M.Shoniya for placing the metallographic compounds and the heavy hydrogen at their disposal, and they also express their gratitude to P.S.Brostyuk, M.I. Dzkuya and G.A.Dorokhova for their practical help. There are 2 figures, 9 tables, and 10 references, 4 of which are Soviet.

SUBMITTED: May 17, 1958

Card 3/3

AUTHORS: Demirkhanov, R. A., Cutkin, T. I., 20-118-6-14/43
Dorokhov, V. V.

TITLE: Masses of Lead Isotopes (Massy izotopov svintsa)

PERIODICAL: Doklady Akademii Nauk SSSR, 1958, Vol. 118, Nr 6,
pp. 1103-1104 (USSR)

ABSTRACT: The present paper reports on the results of the measuring of the masses of lead isotopes Pb^{204} , Pb^{206} , Pb^{207} and Pb^{209} . These measurements were carried out in connection with the determination of the binding energy of the nucleons in a nucleus in the range of the magic numbers 82 and 126 with respect to the protons and with respect to the neutrons, respectively. All this is connected with the necessary exact definition of the mass of the isotope Pb^{208} which is used as base value for the computation of the masses of heavy isotopes with

$z \leq 82$

from the data of the nuclear reactions. The measurements were carried out by means of a device described already earlier by the same authors (ref 1). The dissolving power

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Masses of Lead Isotopes

20-118-6-14/43

of this device amounted in this region to 60,000 - 80,000. In order to increase the accuracy of the measurements the masses of the lead isotopes were determined by immediate comparison with the corresponding mass of hydrocarbons which contain the isotopes H^1 , C^{12} and O^{16} . The values obtained here were controlled by the determination of the mass of the lead isotope from various doublets and by the production of lead ions from various compounds. Each value ΔM of the doublet was determined by treatment of 18-20 mass spectrograms (which were photographed on different plates). The results of the measurements are given in a table. Following is shown by the data of this table: Within the measuring error limits a satisfying "inner" connection exists between the mass values detected from various doublets. The results found here confirm the absence of systematic measuring faults and the reliability of the data obtained here. Finally the differences between the present measurements and the earlier ones are pointed out in short.

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Masses of Lead Isotopes

20-118 -6-14/43

There are 3 tables and 4 references, 2 of which are Soviet

PRESENTED: September 26, 1957, by L. A. Artsimovich, Member of the
Academy of Sciences, USSR

SUBMITTED: July 5, 1957

Card 3/3

21(7), 21(1)

AUTHORS:

Demirkhanov, R. A., Gutkin, T. I.,
Dorokhov, V. V.

SOV/56-36-5-62/76

TITLE:

The Mass of the Isotope Pu²³⁹ (Massa izotopa Pu²³⁹)

PERIODICAL:

Zhurnal eksperimental'noy i teoreticheskoy fiziki, 1959,
Vol 36, Nr 5, pp 1595-1596 (USSR)

ABSTRACT:

Already in a number of previous papers the authors reported on the mass determinations of lead and uranium isotopes, and they also described the mass-spectrometric device used for these measurements (Refs 1, 3, 4). In the present "Letter to the Editor" they give a report on measurements carried out with Pu²³⁹ by means of this spectrometer, which has a resolving power of 60,000 - 80,000. For mass determination doublets of various organic compounds were used, which consisted of the already exactly known elements H, C¹² and O¹⁶, viz. alizarin (C₁₄H₈O₄, M = 240) and perilen (C₂₀H₁₂, M = 252). Ion formation occurred in an arc discharge in helium, the pairs Pu²³⁹ - organic compound

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The Mass of the Isotope Pu²³⁹

SOV/56-36-5-62/76

were introduced into the discharge by evaporation from special crucibles. The mass differences, ΔM of the doublet and the corresponding mass values of Pu²³⁹ are:

doublet	M [mME]	mass of Pu ²³⁹ , [ME]
Pu ²³⁹ - C ₁₄ H ₇ O ₄	18.448±0.082	239.128922±92
C ₁₉ H ₁₁ - Pu ²³⁹	33.447±0.067	239.128695±74

Mean value: 239.128784±165

The mass of Pu²³⁹ calculated from nuclear reactions gives 239.128025±155 if a correction of the more accurately

known value of Pb²⁰⁸ is taken into account, and 239.126999±150 if this correction is not taken into account. It is found that the difference of the masses of

Pu²³⁹ and U²³⁸ calculated according to the authors' data, when compared with the data obtained from nuclear reactions, amounts to only 0.166±0.250 mME, i. e. that it is still within the limits of errors. It is therefore assumed that the error of ~1 mME is due to an inaccurate Q-value.

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The Mass of the Isotope Pu²³⁹

SOV/56-36-5-62/76

There are 1 table and 4 references, 3 of which are Soviet.

SUBMITTED: February 2, 1959

Card 3/3

21(8)

AUTHORS:

Demirkhanov, R. A., Gutkin, T. I.,
Dorokhov, V. V.

SOV/20-124-2-16/71

TITLE:

The Masses of the Isotopes Th²³², U²³⁴, U²³⁵ and U²³⁸
(Massy izotopov Th²³², U²³⁴, U²³⁵ i U²³⁸)

PERIODICAL:

Doklady Akademii nauk SSSR, 1959, Vol. 124, Nr 2, pp 301-303
(USSR)

ABSTRACT:

Measurement of the masses of Th²³², U²³⁴, U²³⁵ and U²³⁸ made it possible to determine the binding energy of nucleons in the nuclei not only of these isotopes but also of many radioactive isotopes connected with them by the naturally-radioactive series 4n, 4n + 2 and 4n + 3. The exact masses of these isotopes have hitherto not been determined by direct measurements. The authors determined the masses of these isotopes by means of an already previously (Ref 3) described mass-spectrographical device having a resolving power of the order of 60000 - 70000. The masses of the isotopes were determined by direct comparison with the corresponding mass of organic compounds. These organic compounds contained H¹, C¹² and N¹⁴, the masses of which are known.

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The Masses of the Isotopes
Th²³², U²³⁴, U²³⁵ and U²³⁸

SOV/20-124-2-16/71

The substances used for measurements are given; control was carried out by determining the mass of U²³⁸ from the doublets C¹²H₁₀-U²³⁸ and C¹²C¹³H₉ - U²³⁸. Short reference is made to a second control method. Each doublet was determined by the evaluation of 18-20 mass spectrograms (which had been photographed on different plates). Results of measurements are given by a table. The masses of the isotope U²³⁸, which were determined from 2 different doublets, agree well with one another within the limits of measuring errors. The "mean value" calculated by taking account of weight amounts to $M_{U^{238}} = 238.127284 \pm 35.10 \cdot 10^{-6}$ mass units. The mass values determined by the present paper are lower than the corresponding values determined by nuclear reactions. Also these differences remain within the limits of permissible deviations, an exception being formed only by uranium.

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The Masses of the Isotopes
Th²³², U²³⁴, U²³⁵ and U²³⁸

SOV/20-124-2-16/71

The authors thank Ye. Ye. Baroni and K. A. Kovyrzina for placing heavy hydrogen at their disposal, and they also thank M. I. Dzkuya, G. A. Dorokhova and P. S. Brostyuk for their active help. There are 3 tables and 11 references, 6 of which are Soviet.

PRESENTED: September 26, 1958, by L. A. Artsimovich, Academician

SUBMITTED: August 29, 1958

Card 3/3

89257

S/048/61/025/001/023/031
B029/B063

24.6510

AUTHORS:

Demirkhanov, R. A., Gutkin, T. I., Dorokhov, V. V.

TITLE:

Masses of heavy atoms and binding energies of nuclei in the range of $174 \leq M \leq 239$

PERIODICAL:

Izvestiya Akademii nauk SSSR. Seriya fizicheskaya, v. 25, no. 1, 1961, 124-129

TEXT: The results of mass-spectrographic measurements of nuclear masses in the range of $174 \leq M \leq 239$, carried out for a large number of isotopes with an accuracy of 10^{-7} to $5 \cdot 10^{-7}$, are presented. The mass spectrograph with double focusing used for the purpose had a resolution of 50,000-80,000. The masses of heavy nuclei were measured by the doublet method and with the use of the organic compounds $C_n H_m$, $C_n C^{13} H_m$, $C_n N_m H_k$, and $C_n O_m N_k H_p$ as standard masses. The question as to whether there is a fine structure in the curve of binding energy in the mass range with $A \sim 200$ can only be answered if the accuracy of measurement is improved by one

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89257

Masses of heavy atoms and binding ...

S/048/61/025/OC1/023/031
B029/B063

X

order of magnitude, i.e., to $\Delta M/M \sim 3 \cdot 10^{-7}$, and a discontinuity of ~ 3 Mev in the binding energy can be established with an accuracy of $\sim 20\%$ if $\Delta M/M \sim 3 \cdot 10^{-6}$. The high degree of accuracy with which the dispersion coefficient can now be measured, and the method developed by the authors make it possible to increase the accuracy of measurement in the respective mass range by a factor of 10-50. In many cases, the mass of the isotope was determined from various doublets, i.e., the "inner agreement" was taken into account. Table 1 contains the masses of the Re, W, Ta, and Hf isotopes and, for comparison, the masses obtained by the mass-spectroscopic method and nuclear reactions. The masses of Re^{185} , Hf^{179} , Hf^{177} , and Hf^{174} were measured for the first time. The mass values of the majority of isotopes measured by the authors are higher than those obtained in Refs. 8 and 9. This is obviously due to the fact that a defective standard mass had been used. A comparison of the present data with similar values obtained by other methods is of particular interest. The results of the present paper are compared in Table 2 with those of other papers. They agree with those published by W. H. Johnson and V. B. Bhaht

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Masses of heavy atoms and binding ...

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S/048/61/025/001/023/031
B029/B063

(Ref. 17) within the limits of error, but differ from the results obtained from nuclear reactions by about the double error. According to what has been said above, the Q values for the reactions

$\text{Hf}^{177}(\gamma, n)\text{Hf}^{176}$ and $\text{Hf}^{179}(\gamma, n)\text{Hf}^{178}$ are probably erroneous, or the limits of error in the determination of the Q values of these reactions must be increased two or three times. The nuclear masses of 42 stable isotopes measured by the authors were then used to determine E/A as a function of A (per nucleon) within the range $174 \leq M \leq 210$ (cf. Fig.). In addition, the binding energies of 66 radioactive nuclei were calculated. Table 3 contains the binding energies B_n of the last neutron and B_p of the last proton, and also the pairing energies P_n and P_p of the neutrons and protons, respectively, for the Hf, Ta, W, and Re isotopes. On the strength of these measurements it is possible to establish some rules concerning nuclear energies. The nucleus has a shell structure, and the shell is completely filled at $Z = 82$ and $N = 126$. In the case of nuclei with odd A, the binding energy is always lower than in the case of nuclei with even A. At equal values of Z, the shell structure may be derived also

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Masses of heavy atoms and binding ...

S/048/61/025/001/023/031
H029/B063

X

from the mutual behavior of the $(Z+1)$ th proton and the $(N+1)$ th neutron ($Z = 82, N = 126$). The binding energy of the last neutron or proton satisfies the law of conservation of parity. The authors thank Ye. Ye. Baroni and his co-workers K. A. Kovyrina and V. M. Soyfer for several preparations, as well as M. I. Dzkuya and G. A. Dorokhova for assistance. This is the reproduction of a lecture read at the Tenth All-Union Conference on Nuclear Spectroscopy, Moscow, January 19-27, 1960. There are 1 figure, 3 tables, and 21 references: 8 Soviet-bloc and 13 non-Soviet-bloc.

Card 4/10

DEMIRKHANOV, R.A.; DORCKHOV, V.V.

Mass of the isotope Pu^{240} . Zhur. eksp. i teor. fiz. 40 no.4:1033-1034
Ap '61. (MIRA 14:7)

(Plutonium--Mass)

DEMIRKHANOV, R.A.; DOROKHOV, V.V.; DZKUYA, M.I.

Isotope masses and binding energies of nuclei in the region from
strontium to ruthenium. Zhur. eksp. i teor. fiz. 40 no.6:1572-
1582 Je '61. (MIRA 14:8)

(Nuclei, Atomic)
(Isotopes--Mass)

S/020/62/146/001/008/016
B108/B102

AUTHORS: Demirkhanov, R. A., Dorokhov, V. V., Dzukaya, M. I.

TITLE: The isotope masses of lutecium, ytterbium and thulium

PERIODICAL: Akademiya nauk SSSR. Doklady, v. 146, no. 1, 1962, 72 - 74 .

ABSTRACT: The isotope masses were measured with a double-focusing mass spectrograph. For reference, doublets formed by ions of the element to be measured and by organic compounds consisting of H¹, C¹², C¹³, N¹⁴, O¹⁶ were used. The spectrograph was able to resolve the masses of C¹² and C¹³. The mass of each isotope was calculated from the mass of the doublet. Results show good agreement with those by V. B. Shmat et al. (Phys. Rev., 129, no. 1, 235, 1960). There are 1 figure and 1 table. ✓

PRESENTED: May 3, 1962 by L. A. Artsimovich, Academician

SUBMITTED: March 26, 1962

Card 1/2

The isotope masses of ...

Table 1.

Legend: m. e. = mass units. (1) Doublet; (2) mass of isotope, (3) average mass (for O^{16} , $M = 16$); the error refers to the last figures.

3/020/32/146/01/0 0/015
B100/B102

Z	11	12	13
169	$C_{12}H_{11}N - Tu^{169}$ $Hg^{169} - Tu^{169}Cl^{169}$	151,972 ± 0,061 70,416 ± 0,029	168,987336 ± 42 168,987011 ± 160
168	$C_{12}H_{11}N - Ib^{168}$ $Ib^{168} - Ib^{168}$	147,055 ± 0,103 2001,543 ± 0,290	167,987707 ± 103 167,987635 ± 202
170	$C_{12}H_{11}N - Ib^{170}$ $C_{12}H_{11}ON - Ib^{170}$ $C_{12}C^{13}H_{11}N - Ib^{170}$	161,882 ± 0,043 125,406 ± 0,150 157,374 ± 0,210	169,990171 ± 43 169,990250 ± 150 169,989297 ± 210
171	$C_{12}H_{11}ON_2 - Ib^{171}$ $C_{12}C^{13}H_{11}N - Ib^{171}$ $Ib^{171} - Ib^{171}$	149,676 ± 0,273 164,195 ± 0,080 1004,531 ± 0,060	170,990516 ± 270 170,990533 ± 29 170,990711 ± 63
172	$C_{12}H_{11}O_2N - Ib^{172}$ $Ib^{172} - Ib^{172}$ $Ib^{172} - Ib^{172}$	103,531 ± 0,660 1500,275 ± 0,226 1082,223 ± 0,118	171,990840 ± 60 171,990802 ± 226 171,990917 ± 127
173	$C_{12}H_9 - Ib^{173}$ $C_{12}H_9O_2N - Ib^{173}$ $Ib^{173} - Ib^{173}$	101,662 ± 0,067 102,812 ± 0,063 1001,022 ± 0,050	172,990976 ± 67 172,990948 ± 63 172,990922 ± 51
174	$C_{12}H_9 - Ib^{174}$ $Ib^{174} - Ib^{174}$	108,342 ± 0,038 2601,634 ± 0,050	173,990942 ± 38 173,990916 ± 100
176	$C_{12}H_9 - Ib^{176}$ $C_{12}H_9N - Ib^{176}$	120,018 ± 0,046 167,223 ± 0,109	175,990557 ± 46 175,990772 ± 109
175	$C_{12}H_9 - Lu^{175}$ $C_{12}C^{13}H_9 - Lu^{175}$	114,157 ± 0,077 104,795 ± 0,036	174,990272 ± 37 174,990150 ± 30
170	$C_{12}H_9 - Lu^{170}$ $Lu^{170} - Lu^{170}$	120,000 ± 0,049 1002,301 ± 0,020	175,990575 ± 20 175,990519 ± 70

Card 2/2

DEMIRKHANOV, R.A.; DOROKHOV, V.V.; DZKUYA, M.I.

Isotope masses and nucleon binding energies in the rare-
earths region ($150 \leq A \leq 176$, $63 \leq Z \leq 71$). Izv. AN SSSR.
Ser. fiz. 27 no.10:1338-1356 0 '63. (MIRA 16:10)

DEMIRKHANOV, P.A.; DORONKOV, V.V.; SOLOV'YEV, V.G.

Asymptotic binding energy of the last two neutrons in the region
N = 86 and 82. Izv. Akad. Fiz. 2 no.1:10-13 Jr '65.

(MIRA 18:8)

1. Ob'yedinennyy Institut yadernykh issledovaniy.

DEMIRKHANOV, R.A.; DOROKHOV, V.V.; DZKUYA, M.I.

Masses of stable isotopes of neodymium, praseodymium, cerium, and lanthanum. Izv. AN SSSR.Ser. fiz. 29 no.5:857-861 My '65. (MIRA 18:5)

IYERUSALIMSKIY, N.D.; ANDREYEVA, Ye.A.; GRISHANKOVA, Ye.L.; GOLOVLEV, Ye.L.;
DOROSHOV, V.V.; ZHUKOVA, L.N.

Study of microflora of refinery waste waters. Prikl. biokhim.
1 mikrobiol. 1 no.2:163-166 Mr-Apr '65.

(MIRA 18:11)

1. Institut mikrobiologii AN SSSR, Moskva.

L 36092-66 EWT(m)/T WE

ACC NR: AP6015206

(A)

SOURCE CODE: UR/0411/65/001/002/0163/0166

AUTHORS: Iyerusalimskiy, N. D.; Andreyeva, Ye. A.; Grishankova, Ye. L.; Golovlev, Ye. L.; Dorokhov, V. V.; Zhukova, L. N.

ORG: Institute of Microbiology, Academy of Sciences, SSSR, Moscow (Institut mikrobiologii Akademii nauk SSSR)

53

B

TITLE: A study of the microflora of sewage of petroleum refineries

SOURCE: Prikladnaya biokhimiya i mikrobiologiya, v. 1, no. 2, 1965, 163-166

TOPIC TAGS: bacteria, fuel microorganism, industrial waste, petroleum refining, yeast, aromatic hydrocarbon, diesel fuel, kerosene

ABSTRACT: The results of a study of active slime from petroleum refineries are given. Active slimes from waste phenolic water and from oil traps (purified of petroleum by six-fold extraction by benzene) were studied. Recent and old slimes from oil refinery No. 4 and a sample of slime from the trap of No. 4 were also studied. The specimens were kept in the active state in Sengen's medium at pH 7. From the slimes, 575 cultures were extracted, and 145 other cultures were extracted from similar sources. The mycobacteria were 44%, the bacteria 28%, and yeast 26%. All the bacteria were gram-negative nonspore-forming. They were represented mostly by Pseudomonas and Achromobacter. The yeasts were Candida and Torulopsis. All of the extracted microorganisms grew well in pure kerosene, pure paraffin, diesel-fuel distillate, and

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mm. 622.35+613.663

L 36092-66

ACC NR: AP6015206

paraffin-base petroleum. It was found that only certain mycobacteria and bacteria grow in aromatic hydrocarbons. Orig. art. has: 3 tables.

SUB CODE: 06/1/SUBM DATE: 18Jan65/ ORIG REF: 003/ OTH REF: 009

LS
Card 2/2

DOROKHOV, V.Ya.

Genesis of platform structures of the second and third orders.
Izv.AN SSSR.Ser.geol. 25 no.1:15-27 Ja '60. (MIRA 13:8)

1. Nizhne-Volzhskiy filial Vsesoyuznogo nauchno-issledovatel'skogo
geologo-razvedochnogo neftyanogo instituta, g.Saratov.
(Saratov Province--Geology, Structural)

ACC NR: AR6029303

SOURCE CODE: UR/0271/66/000/006/B056/B056

AUTHOR: Dorokhov, Ye. Ya.

TITLE: The optimum frequency of data readout from discrete signal transducers

SOURCE: Ref. zh. Avtomatika, telemekhanika i vychislitel'naya tekhnika, Abs. 6B426

REF SOURCE: Sb. Algoritmiz, proizv. protsessov. Seminar. Vyp. 8. Kiyev, Nauk. dumka, 1965, 16-31

TOPIC TAGS: data readout, data retrieval, discrete automation, optimal control

ABSTRACT: The problem is considered of selecting the frequency of data readout from discrete signal transducers; thereby it is assumed that a certain value of the frequency can lead to data losses. Both data losses caused by equipment failure and structural losses associated with the method and speed of processing the given type of input signals serve as the criterion of system's operation. A method for determining the optimum frequency of data readout is indicated, and the upper and lower limits of the optimum frequency of data readout are shown. Equations are derived which make it possible, for a given magnitude of permissible data losses, to select a definite value of the data processing cycle which would determine the upper limit of the optimum frequency of data readout. [Translation of abstract] 4 illustrations and bibliography of 8 titles. N. S.

SUB CODE: 09,13
Card 1/1

UDC: 681.142.621

BYKHOVSKIY, I.I. (Moskva); DOROKHOVA, A.D. (Moskva); ZARETSKIY, L.B.
(Moskva); LUKOMSKIY, S.I. (Moskva)

Some periodic movements and the structure of the phase space
of an impact-vibration system with a regularly recovered
force. Izv. AN SSSR. Mekh. i mashinostr. no. 2:161-165
Mr-Apr '64. (MIRA 17:5)

BOROVIKOV, Vasilii Aleksandrovich; KOSAREV, Vladimir Kuz'mich; KHODOT, Georgiy Aleksandrovich; SLAVIN, M.I., kand. tekhn.nauk, retsenzent; DOROKHOVA, A.I., inzh., retsenzent; GESSEN, V.Yu., doktor tekhn. nauk, red.; SOBOLEVA, Ye.M., tekhn. red.

[Electrical networks and systems] Elektricheskie seti i sistemy. Moskva, Gosenergizdat, 1963. 459 p. (MIRA 16:8)
(Electric lines--Overhead)

USSR/Diseases of Farm Animals, Non-Contagious Diseases.

R-2

Abs Jour : Ref Zhur-Biol., No 18, 1958, 83583

Author : Fedorov, B. T.; Miroljubov, I.I.; Polivanskaya, K.
D.; Dorokhova, A. K.

Institute : No institute is given

Title : Steatitis Disease in Minks.

Orig Pub : Karakulevodstvo i zverovodstvo, 1957, No 6, 54-56

Abstract : At one of the sovkhoses for animal breeding, an outbreak of polyavitaminotic steatitis ("yellow fat" disease) occurred among young minks. The disease was characterized by a general depression, by food refusal, by diarrhea with yellow or dark-green feces, sometimes by seizures accompanied by spasms or paralyzes. An autopsy of succumbed animals uncovered a well advanced degenerative adiposity. The disease was caused by continuous feedings of fish remnants containing rancid fat to the animals.

Card 1/1

DOROKHOVA, D.V.

Vasomotor lesions in focal inflammatory brain diseases in children.
Fiziol.zhur. [Ukr.] 10 no.4:551-554 J1-Ag '64.

(MIRA 18:11)

1. Detskaya psikhonevrologicheskaya klinika i laboratoriya
fiziologii vysshey nervnoy deyatel'nosti Ukrainского
psikhonevrologicheskogo instituta, Khar'kov.

DOROKHOVA, K.Ya.; SOKOLOVA, T.A.

Chemical composition of the layer covering the grains of primary
minerals in some mountain-taiga soils of eastern Transbaikalia.
Pochvovedenie no.10:34-36 0 '63. (MIRA 16:12)

1. Pochvennyy institut imeni V.V.Dokuchayeva.

FRIDLAND, V.M.; DOROKHOVA, K.Ya.; ZHITKOVA, A.I.

Nature of the structure of humid tropical soils (North Vietnam).
Dokl. AN SSSR 154 no. 3:707-709 Ja '64. (MIRA 17:5)

1. Pochvennyy institut im. V.V.Dokuchayeva i Ministerstvo sel'
skogo khozyaystva SSSR. Predstavleno akademikom I.P.Gerasimovym.

DOROKHOVA, M.A.

Following Valentina Gaganova's example. Transp.stroi. 10
no.3:7 Mr '60. (MIRA 13:6)
(Shipitsina, Anfisa Ivanovna)

CA 1020116V, M-1

Syntheses of derivatives of α -amino acids. II. New method of synthesis of amides of α -amino acids. V. F. Kurbatov and M. I. Danilova. *Zhur. Obshch. Khim.* (J. Gen. Chem.) 31, 1485-6 (1951); *J. Gen. Chem. U.S.S.R.* 31, 1821-3, 1957-30 (Eng. Translation); cf. Bergmann and Zervas, *C.A.* 29, 1672; preceding abstr.— β -Alkyl- β , γ -dihydroamides are prepared from α -(carbobenzylamino) acids with PCl_5 ; the products with NH_3 yield amides of α -amino acids. The following *N*-carbobenzylamides of α -amino acids were prepared conventionally from 10% excess of $\text{PCl}_5 \cdot \text{O}(\text{COCl})$ in alk. soln. at 0-3°: DL-valine (I), 73.8%, m. 73-3° (from petr. ether); DL-glycine (II), 71%, m. 85-7° (from CCl_4 -petr. ether); DL-leucine (III), 70%, m. 84-6°; DL-isoleucine (IV), 74%, m. 105-6° (from CCl_4 -petr. ether). To 9 g. I in dry Et_2O was added at -3 to 0° 8.3 g. powdered PCl_5 , and the mixt. stirred 15 min. at 0° and 1 hr. at room temp., filtered, and evapd. in vacuo with addn. of petr. ether, yielding 90% *N*-isopropyl- β , γ -dihydroamidine, decamp. 80-1° (from Et_2O); on standing in air or on heating it loses CO_2 and forms a high-melting polymer, $(\text{C}_7\text{H}_{11}\text{N})_n$. Similarly, II yields *N*-propyl- β , γ -dihydroamidine (V), 84.3%, decamp. 67-9° (from Et_2O), while III gives 80% *N*-isobutyl analog, m. 48-50°, and IV gives 85% *N*-isobutyl analog, m. 78-80°. Similarly, DL-*N*-(carbobenzyl)phenylalanine and PCl_5 in dry Et_2O at 0° gave 72% DL-*N*-(carbobenzyl)phenylalanyl chloride, isolated by diln. with petr. ether in the cold; refluxing in Et_2O 1 hr. yields *N*-isopropyl- β , γ -dihydroamidine, decamp. 123-5° (from CHCl_3). Addn. with cooling of 3.5 g. V to 30 ml. MeOH aqcd. with

NH_3 at 0°, gives 3.2 g. crude III, salt of DL-*N*-(carbobenzyl)phenylalanineamide, decamp. 110-1°, which, boiled until dissolved in CHCl_3 , loses CO_2 and NH_3 , yielding 60% DL-phenylalanineamide, m. 67-9°. Similarly were obtained: DL-isoleucineamide, 61.5%, m. 77-9°; DL-isovalineamide, 68%, m. 80-1° (NH_3 salt of α -(*iso*-leucylamino)isovalineamide, decamp. 108-9°); DL-leucineamide, 60%, m. 104-5°; DL-phenylalanineamide, 61%, m. 127-9°. III. Synthesis of decarboxylated peptides. *Ibid.* 1491-4.—Extension of the reaction described in paper II of the series showed that amides may be substituted for NH_2 in the reactions with the cyanidides. Thus, addn. to 3-6 moles of the amine in dry MeOH at about 0° of the β -alkyl- β , γ -dihydroamidine, followed by 0.5 hr. at 0°, yields on evapn. in vacuo, the amine salt of the amide of the corresponding α -(carbobenzylamino) acid, which, boiled 1 hr. in MeOH , then allowed to stand 1 hr., yields the desired decarboxylated peptide in 80-85% yields. Such a treatment of *N*-isopropyl- β , γ -dihydroamidine (I) (3.3 g.) and 30 ml. 20% MeNH_2 in MeOH gave the MeNH_2 salt of DL-*N*-(carbobenzylamino)-*N*-methylisoleucineamide, decamp. 96-102°.

which yielded 83% *N*-*tert*-butylcarbamoyllysine (DL- α -amino-*N*-methylisoleucine), bp 100-10°, n_D^{20} 1.4712, d_4^{20} 0.9040. Similarly, the 3-Pr analog of I gave 86.3% DL- α -amino-*N*-methylisoleucine, bp 110-11°, n_D^{20} 1.4703, d_4^{20} 0.9054. The 3-*iso*-Bu analog of I gave the *MeNH* salt of DL- α -(carbamoyl-*tert*-butyl)-*N*-methylisoleucine, decamp. 98-104°, which gave 80% DL- α -amino-*N*-methylisoleucine, bp 131°, n_D^{20} 1.4670, d_4^{20} 0.9060. The 3-Bu analog of I gave 60% DL- α -amino-*N*-methylisoleucine, bp 120-1°, n_D^{20} 1.4665, d_4^{20} 0.9074, which crystallizes on prolonged standing. The 3-benzyl analog of I gave 45% DL- α -amino-*N*-methyl- β -phenylpropionamide, bp 165-7°, low-melting solid. Similarly, *iso*-AmNH₂ gave with the corresponding oxazolidinone: DL- α -amino-*N*-isomethylisoleucine, 80%, bp 140-0°, n_D^{20} 1.4620, d_4^{20} 0.9273; *isocaproamide* analog, 84.3%, bp 170-1°, n_D^{20} 1.4608, d_4^{20} 0.9161. $\text{FeCl}_3 \cdot \text{CH}_2\text{Cl}_2$ salt, similarly gave: DL- α -amino-*N*-phenethylisoleucine, 80%, bp 190-1°, n_D^{20} 1.5275, d_4^{20} 1.0376; *isocaproamide* analog, 85%, bp 170-1°, n_D^{20} 1.5213, d_4^{20} 1.0175 (HCl salt, m. 242-3°), and *caproamide* analog, 82%, bp 175-6°, crys. on cooling.

(C. M. Kosolapoff

DOROKHOVA, M. I.

USSR/Chemistry - Pharmaceuticals

11 Feb 53

"The Synthesis of Quinuclidinecarboxylic Acid-(2),"
M. V. Rubtsov, M. I. Dorokhova; All-Union Sci-Res
Chemicopharmaceutical Inst im S. Ordzhonikidze

DAN SSSR, Vol 88, No 5, pp 843, 844

Worked out a simple 5-step synthesis for quinu-
clidinecarboxylic acid-(2) starting with γ -picoline
and mesoxalic ester. Presented by Acad V. M.
Rodionov 28 Nov 52.

264T26

Dorokhova, M. I.

Syntomycin analogs. A. I. Franov, A. P. Arsendoruk, V. P. Klavko, V. A. Mikhailov, T. V. Protopenova, A. P. Skoldinov, and N. B. Smolina. U.S.S.R. 102,626, Apr. 30, 1956. Salts of the corresponding α -aminocrotonophenones are acylated in an aprotic solvent with acyl halides of the corresponding ω -acids in the presence of aq. NaOAc. The resulting α -acylaminoacetoephonones are condensed with CH_2O and the oxo group reduced by known means. M. Hoesel. ///

Dorobh... M...

Distri: 4E43/4E3d/4E2c(3) 7

(Nitr-o-acetamide-4-hydroxyphenolacetate, A. P. ...
Kondraty, M. L. Dorobh... V. A. Biktay, G. ...
Ieva, A. P. Goidinov, G. M. Socolin, and N. B. ...
W O B P 107 945 May 25 1964 The title compound is ...
formed by the interaction of formaldehyde with ...
a-aminocretin-phenone in the presence of ...
agents such as trimethylamine

SM 2 ✓

Dora Khatun, #11

37 7 6
~~Precipitation of aluminum compounds in alcoholic or aqueous-alcoholic media. Y. A. Mikheyev, M. I. Durovaya, A. M. Zhelezovskaya, and N. E. Melnik. U.S.S.R. 102,751, May 25, 1956. To the alc. or aq.-alc. medium is added an alkali sulfate and H₂SO₄ in a quantity less than is required by the equation: 2Al(OH)₃ + 14SO₄ + 3H₂SO₄ = 2MAl(SO₄)₃ + 6H₂O, where M is an alkali metal. The pptn. is best carried out at 45-65°. Thus pptd. Al filters well and the yield of the end product is increased.~~

~~M. Kosob~~

fm

Dorekines, M.

7
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 11

p-Nitro- α -(dichloroacetyl)aminoacetophenone. A. P. Ar-
 eudovskiy, A. I. Ivanov, M. I. Duzakhtova, Y. A. Mikhalev,
 A. P. Skoldinov, D. D. Smolin, and N. R. Smolin. U.S.-
 S.R. 103,915, June 25, 1957. Addn. to U.S.S.R. 69,838.
 1-Hydroxy-1-(*p*-nitrophenyl)ethanamine is treated with
 Cl_2CHCO_2Me and the resulting 1-hydroxy-1-(*p*-nitrophenyl)-
 2-dichloroacetylaminethane is oxidized to *p*-nitro- α -(dichloro-
 acetyl)aminoacetophenone. M. Hoach

Darokhova, M. I.

~~1,1'-Diphenyl-2,2'-diethyl ether, and their di-~~
 nitro derivatives M. I. Darokhova, V. A. Mikhailov, and
 N. P. Sholite. U.S.S.R. 104,154, Nov. 25, 1955. The
 corresponding styrene hydrocarbons are dehydrated by heat-
 ing with Fe, FeCl₃, Fe salts, or other dehydrating agents,
 such as H₂SO₄ or sulfonic acids, and the resulting 1,1'-di-
 phenyl-2,2'-diethyl ethers are nitrated with concd.
 HNO₃ in the usual manner. M. Hirsch

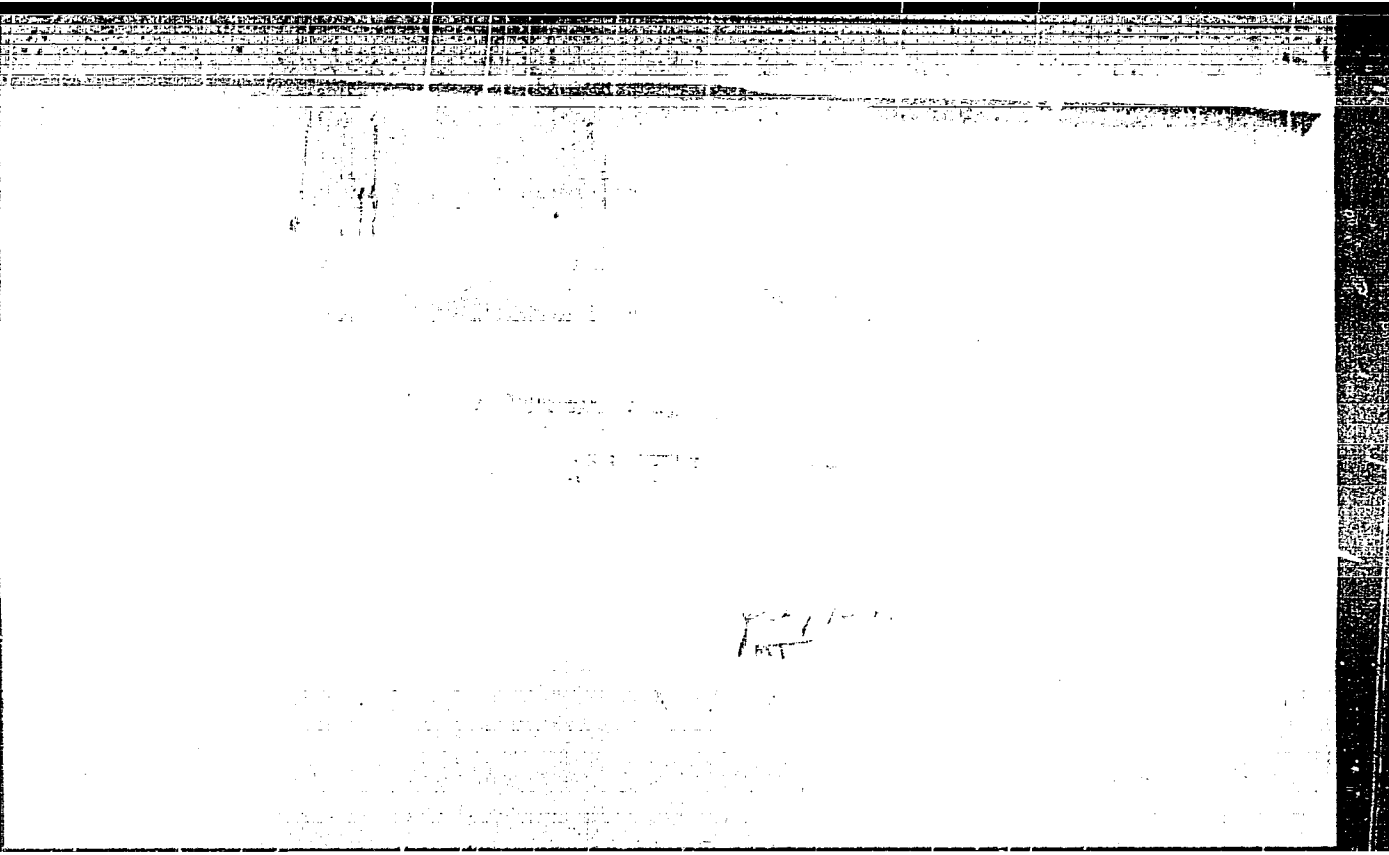
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DOROKHOVA, M. I.

Dorokhova, M. I. -- "Investigation of Certain Transformations of Simple Ethers of Styrol Halohydrins." All-Union Sci Res Chemicopharmaceutical Inst imeni S. Ordzhonikidze. Moscow, 1956. (Dissertation For the Degree of Candidate in Chemical Sciences.)

So: KNIZHNAYA Letopis', No. 11, 1956, pp 103-111



DOROKHOVA, M. I.

Diary 4243

~~Phenothiazine series. II. Synthesis of 10-(N-methyl-
 piperidinyl)phenothiazines. S. V. Zhuravlev, A. N.
 Gritsenko, and M. I. Dorokhova. Zhur. Obshch. Khim.
 27, 1828-1831 (1957), Cl. C.A. 31, 6021a; Nieschulz, et al.
 C.A. 49, 7761d. — Refluxing 10.6 g. pic acid with 18.9
 g. HSO₃Me and 25 ml. dry Me₂CO 2-2.5 hrs. gave after
 addn. of Et₂O 81.5% quaternary salt, m. 73-4°, which be-
 dry-crystallized in EtOH over Pt at atm. pressure to 70.5%. Fr
 V. *N*-methyl-3-piperidinylpicinate, bp., 92-5°, n_D^{20} 1.4499. This
 (91.2 g.) to 100 g. iso-AmOH and 163 ml MePh was added
 to 12.5 g. powd. Na in MePh at reflux yielding after aq.
 treatment 36.9% *N*-methyl-3-piperidinylcarbamid, bp. 189-11°,
 n_D^{20} 1.4769. This treated with dry HCl in CHCl₃ followed
 by SOCl₂ and heating 2 hrs. on a steam bath gave 61.4%
N-methyl-3-piperidinylmethyl chloride HCl salt, m. 168-9°,
 which with NaOH gave the free chloride (I), bp. 58-60°,
 n_D^{20} 1.4729. A stream of dry CO-free air was passed 1 hr.
 through 0.3 g. Na in 200 ml. liquid NH₃ and 0.1 g. Fe-
 (NO)₃; the mixt. was then treated with 1.4 g. Na and
 stirred 1 hr. and the resulting suspension of NaNH₂ treated
 with 12 g. phenothiazine; when a red soln. formed, 9 g. I
 was added and after 2 hrs. NH₃ was distd. and the residue
 isolated 4 hrs. to 130-40°; after refluxing 1 hr. with MePh
 the mixt. was treated with H₂O and dil. HCl, yielding *N*-*N*-
 dimethyl-3-piperidinylmethylphenothiazine HCl salt, m. 228-9°,
 12.54 g. yield; free base, bp., 150-3°. Similarly, 2-chloro-
 phenothiazine gave *N*-*N*-methyl-3-piperidinylmethyl-2-chloro-
 phenothiazine, m. 56-8°; HCl salt, m. 210-11°.~~

G. M. Kosolapoff

DM 1/

MIKHALEV, V.A.; DOROKHOVA, M.I.; SMOLINA, N.Ye.; ZHELOKHOVTSEVA, A.M.; IVANOV, A.I.; ARENDARUK, A.P.; GALCHENKO, M.I.; SKORODUMOV, V.A.; SMOLIN, D.D.

Styrene as raw material for the production of synthomycin and levomycetin. Part 1: Synthesis of p-nitro- α -acylaminoacetophenones. Antibiotiki, 4 no.2:21-24 Mr-Apr '59. (MIRA 12:7)

1. Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevticheskiy institut imeni S. Ordzhonikidze (for Mikhalev, Dorokhova, Smolina, Zhelokhovtseva). 2. Institut farmakologii i khimioterapii AMN SSSR (for Skoldinov, Ivanov, Arendaruk, Galchenko, Skorodumov, Smolin).

(CHLORAMPHENICOL, prep. of.

synthesis from styrene through p-nitro- α -acylaminoacetophenones (Rus))

(VINYL COMPOUNDS

styrene, use in chloramphenicol synthesis through p-nitro- α -acylaminoacetophenones (Rus))

(KETONES

p-nitro- α -acylaminoacetophenones, intermediate in chloramphenicol synthesis from styrene (Rus))

MIKHALEV, V.A.; DOROKHOVA, M.I.; SMOLINA, N.Ye.; ZHELOKHOVTSEVA, A.M.;
TIKHONOVA, O.Ya.; SKOLDINOV, A.P.; ARENDARUK, A.P.; SMOLIN, D.D.;
GOLOVKINA, T.V.; SLONOVA, L.A.

Styrene as an initial product for synthomycetin and levomycetin
production. Part 2: Synthesis of p-nitroacetophenone and
p-nitro- α -bromacetophenone. Antibiotiki 4 no.4:21-24 J1-Ag
"59. (MIRA 12:11)

1. Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevticheskiy
Institut imeni S.Ordzhonikidze (for Mikhaev, Dorokhova, Smolina,
Zhelokhovtseva, Tikhonova). 2. Institut farmakologii i khimio-
terapii AMN SSSR (for Skoldinov, Arendaruk, Smolin, Golovkina,
Slonova).

(CHLORAMPHENICOL chem)
(KETONES chem)

MIKHALEV, V.A.; DOROKHOVA, M.I.; SMOLINA, N.Ye.

Mechanism of conversions of α -acylamino- β -oxypropiofenones into the corresponding benzoyl acetyls. Part 2: Synthesis and cleavage of α -benzenesulfamidoacrylophenones. Zhur. ob. khim. 30 no.11:3714-3718 N'60. (MIRA 13:11)

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

MIKHALEV, V.A.; DOROKHOVA, M.I.; SMOLINA, N.Ye.; TIKHONOVA, O.Ya.

β -Haloalkyl amines and products of their transformations.

Part 1: Reaction of bis(β -chloroethyl)amine with α -oxides.
Zhur. ob. khim. 34 no.11:3716-3719 N '64 (MIRA 18:1)

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

L 41024-65

ACCESSION NR: AP5008582

S/0286/65/000/006/0130/0130

AUTHORS: Mikhailev, V. A.; Vlasov, A. S.; Dorokhova, M. I.; Moskalik, Ya. K.;
Smolina, N. Ye.; Tikhonova, O. Ya.; Shagalov, L. B.

TITLE: A method of preparing 3,4-bis-(n)-diethylaminoethoxy-(phenyl)-hexane.
Class 30, No. 152540

SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 6, 1965, 130

TOPIC TAGS: hexane, chloride, pharmacology

ABSTRACT: This Author Certificate presents a method of producing 3,4-bis-(n)-diethylaminoethoxy-(phenyl)-hexane by interaction between synestrol and diethylaminoethyl chloride in alcohol in the presence of alkali agents with subsequent distillation of the alcohol, addition of water, and extraction by an organic solvent such as ether. In order to increase the yield of the desired product and to suppress the by-products of the reaction, diethylaminoethyl chloride and the alkali agent are introduced gradually, in several doses, either in solid form or in alcohol solutions. Production of the pharmaceutical preparation is effected by widely accepted methods. In order to reduce danger and to facilitate

Card 1/2

L 41024-65

ACCESSION NR: AP5008582

the process, a diethylaminoethyl chloride salt is used, such as chlorhydrate. The process is also facilitated and simplified by using caustic potash or caustic soda as the alkali agent. To prevent excessive dilution of the reaction mass, the excess solvent is distilled simultaneously with introduction of the alcohol solutions of the reaction products. For all the synestrol to react, 150-170% of the theoretically computed diethylaminoethyl chloride required is used.

ASSOCIATION: none

SUBMITTED: 13Nov61

ENCL: 00

SUB CODE: CC, I&

NO REF SOV: 000

OTHER: 000

llc
Card 2/2

MIKHALEV, V.A.; DOROKHOVA, M.I.; SMOLINA, N.Ye.; TIKHONOVA, O.Ya.

β -haloalkyl amines and their transformation products. Part 2:
Derivatives of N', N"-dispirotriperazinium. Zhur.org.khim.
1 no.3:460-464 Mr '65. (MIRA 18:4)

1. Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevticheskiy
institut im. S.Ordzhonikidze.

L 61194-65 EMP(e)/EPA(s)-2/EMT(m)/EFP(c)/EAP(i)/EPA(w)-2/EMP(j)/T/EMP(b)

Fc-4/Fq-4/Pr-4/Pt-7 W/JAJ/EM/WH

ACCESSION NR: AP5019028

UR/0286/65/000/012/0063/0063
656.266.3

AUTHOR: Leko, V. K.; Dorokhova, M. L.

48
B

TITLE: Glass for glass ceramic materials. Class 32, No. 172001

SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 12, 1965, 63

TOPIC TAGS: glass, glass ceramic pyroceram, sitall, semiconductor

ABSTRACT: An Author Certificate has been issued for a glass for glass ceramics. To produce glass ceramics with an electronic conductivity of 10^{-9} to 10^{-12} ohm⁻¹.cm⁻¹, the glass is formulated to contain: 30-50% SiO₂, 5-15% TiO₂, 5-25% Al₂O₃, 30% max B₂O₃, 10-20% MgO, 15% max CaO, 3% max Na₂O, 4% max K₂O, and 2-10% Al metal. [6M]

ASSOCIATION: none

SUBMITTED: 27Mar64

ENCL: 00

SUB CODE: MT

NO REF SOV: 000

OTHER: 000

ATD PRESS: 4052

Card 1/1

ZHUKOVSKIY, V.I.; DOROKHOVA, M.P.; ZAREMBA, N.Ye.; DYKMAN, D.G.; BOYS, G.V.

Data of the thermographic investigation of barium titanate with
some admixtures. Izv. AN SSSR Ser. fiz. 24 no.10:1294-1295 0 '60.
(MIRA 13:10)

(Barium titanate)

DOROKHOVA, N. A.

PHASE I BOOK EXPLOITATION SOV/3671

Academy of Sciences USSR. Institut elektronnykh upravlyayemykh mashin Tsifrovaya tabulitsa i vychislitel'nye ustroystva; (Sbornik) (Digital Technics and Computing Devices: Collection of Articles) Moscow, Izd-vo AN SSSR, 1959. 184 p. Errata slip inserted. 4,000 copies printed.

Ed. I. N. S. Bruk, Corresponding Member, USSR Academy of Sciences; Ed. of Publishing House: O. Yu. Shchepinok; Tech. Ed.: V. V. Volkova.

PURPOSE: This collection of articles is intended for persons specializing in computer technics.

COVERAGE: Most of the work in this first issue of the Collection of Articles of the Institute of Automatic Control Machines and the Academy of Sciences, USSR, was carried out during 1958-1959, and was devoted to digital technics. The Institute conducted studies aimed at creating a high-speed measuring instrument. One of the results of this work was improvement of the accuracy by replacing its static storage device with ferrite storage devices. Other articles concern the use of computers in digital computers, stability of analog computers equipped with digital computers, and the use of the M-2 computer in solving various problems. Particular attention is given to articles which present the results of work in digital technics in mathematical investigations, and in control machines and systems of control machines mentioned on the principle of digital technics. Some particularities are mentioned in the articles.

References are given to some of the articles.
References are given to articles:
1. Dorokhova, N. A. Solving Problems of Control Machines with Direct Coupling
2. Zakharenko, A. B., and L. Ya. Chumakov. Des of Surface-Barrier Transistors
3. Zakharenko, A. B., and L. Ya. Chumakov. Des of Surface-Barrier Transistors
4. Zakharenko, A. B., and L. Ya. Chumakov. Des of Surface-Barrier Transistors
5. Zakharenko, A. B., and L. Ya. Chumakov. Des of Surface-Barrier Transistors
6. Zakharenko, A. B., and L. Ya. Chumakov. Des of Surface-Barrier Transistors
7. Zakharenko, A. B., and L. Ya. Chumakov. Des of Surface-Barrier Transistors
8. Zakharenko, A. B., and L. Ya. Chumakov. Des of Surface-Barrier Transistors
9. Zakharenko, A. B., and L. Ya. Chumakov. Des of Surface-Barrier Transistors
10. Zakharenko, A. B., and L. Ya. Chumakov. Des of Surface-Barrier Transistors

A frequency meter using a generator of standard frequency with quartz stabilization was developed at the Laboratory of Control Machines and Systems. This meter was used for measuring the frequency of signals with errors not exceeding ± 0.05 cps within a range of 50-1.5 cps. This arrangement is similar to that with PZ transistors and its power consumption is about 350 mw. It was found that the application of digital techniques permitted attainment of a high degree of stability.

Masontov, O. V. Study of the Technological Spread of Parameters in Transistors
The measurements of parametric spread made by the author demonstrate that this spread occurs independently of each individual parameter. The results of measurements are presented statistically. There are five references, all Soviet (one of these is a translation).

Masontov, O. V. Instability of Transistor Characteristics and Parameters
The author presents the results of experimental testing of the parameters and characteristics of P4, P6, P8, and P15-type transistors

Lepov, M. M. Stability of Electronic Simulation Circuits Equipped with d-c Amplifiers
The author discusses ways of obtaining stable simulation circuits for solving problems by analog computers equipped with operational d-c amplifiers with strong feedback. Such circuits are widely used for analyzing dynamics of automatic control systems. The author studies stability conditions of circuits used for the solution of problems described by differential or integral-differential equations. These problems are solved with the time-delay units of the analog computer, on which transients in variables have been instrumented in the form of time-variable does not appear as a result of the independent units. The author analyzes some typical equations, presents their block diagrams, and finds conditions for stability. There are seven references: 6 Soviet (one of which is a translation) and 1 English.

Dorokhova, N. A. Solving Problems of Control Machines with Direct Coupling

S/799/62/000/003/002/008

AUTHORS: Avaliani, Yu. Ye., Alekseyev, Yu. N., Glukhov, Yu. N., Dorokhova, N. A., Tanetov, G. I.

TITLE: The arithmetic equipment of a specialized machine.

SOURCE: Akademiya nauk SSSR: Institut elektronnykh upravlyayushchikh mashin. Tsifrovaya tekhnika i vychislitel'nyye ustroystva. no. 3. 1962, 14-23.

TEXT: The paper describes an arithmetic equipment (AE) of the parallel type, which operates with 22-digit binary numbers with a fixed decimal point and which performs addition, subtraction, multiplication, division, extraction of the square root, matching, shifting, and transposition of numbers. An acceleration in the multiplicative operations is achieved by the accumulation of the partial products without transitional carry-overs. The system of the elements and the design principles of the AE are briefly examined. The system of elements comprises a static trigger, a potential-impulse gate, and logic diode circuits. All of the elements are made up of semiconductor devices. The network of the AE is presented in skeletal form, which comprises the various equipments that serve to perform the elementary operations in each register, and the equipments that receive numbers from other partial parts of the machine. The operational algorithms of addition, subtraction,

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The arithmetic equipment of a specialized machine. S/779/62/000/003/002/008

and division, and the technical methods in the design of the logical circuits which help to realize the algorithms, are similar to those employed in some existing computers, for example, the M-2. Thus, for example, the adding equipment of the AE differs in its logic structure from that employed in the M-2 machine only by the content of cyclic carry-over circuit from the higher digit to the lower digit. While the operation of algebraic matching exhibits certain peculiarities dependent on the character of the problems to be solved, there is nothing interesting from the point of view of engineering. In this operation, the same circuits as those utilized in addition and subtraction are employed. The operation of shifting is also of no additional interest, since it employs the same shifting circuitry employed in multiplication and division. In the multiplication the partial products remain immobile, whereas the multiplicand is shifted to the right. It can be shown that to obtain, in such procedure, an accuracy of no less than a unit of the lowest digit for 22-digit initial figures, it is necessary to have 3 additional digits in the AE prior to rounding off. Extraction of the square root follows almost precisely the same method as that employed in high-school long-hand work, that is, with division of the number into pairs of digits, extraction of the square root of the highest digital pair, and all the other subsequent steps required by the 2-rectangles-cum-small-square method, until the remainder is either zero or smaller than the required accuracy residual. The duration of the extraction of the square root amounts to 112 cadences or 317 μ sec.

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The arithmetic equipment of a specialized machine.

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If the number of which the square root is to be obtained has a minus sign, then all the digits go to zero, and the operation comes to a halt. The description of the AE elements comprises the static trigger, the logical diode scheme, and the potential impulse gate, schematic circuits for all of which are shown. A block diagram is shown for a basic (k-th) digit of the AE. The AE described contains approximately 1,000 semiconductor triodes and 4,000 semiconductor diodes, all of which operate in regimes in which current intensities, voltages, and powers do not exceed the rated values. A special cooling system ensures maintenance of all semiconductor devices at room temperature. The circuits employed ensure maintenance of a stable operation of the AE under power-supply-voltage fluctuations of $\pm 10\%$ from nominal values. The electrical power supply of the AE is provided by a 400-cps rotary generator through rectifiers assembled in a 6-phase circuit. The total power requirements of the AE is approximately 0.8 kw. The AE is currently in experimental operation. There are 5 figures and 3 references (2 Russian-language Soviet and the English-language A.A. Robinson, Multiplication in the Manchester University high-speed digital computer. Electronic Engrg., v.25, no.299, 1953).

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DOROKHOVA, N.A.

Mycorrhizal nutrition of grasses. Biol. Zhurn. 1965. 73 no. 7: 119-125. (MIRA 1965)

BURNASHEV, M.S.; CHEPURNOV, V.S.; KUBRAK, I.F.; DOROKHOVA, N.I.

Materials on fishes of the Sasyk (Kunduk) Lagoon collected in the
summer of 1956. Uch.zap.Kish.un. 32:63-72 '58. (MIRA 13:6)
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IVANOV, B.I.; DOROKHOVA, N.P.; BOROZDINA, Ye.V.; KOSAREVA, Ye.A.

Dephenolizing the phenol waters of the "Slantay" Combline
with a mixture of n-butyl ether and isopropyl ether. Trudy
VNIIT no.12:266-270 '63. (MIRA 18:11)

DOHOXHOVA, V.

Main task is to improve guidance. Prof.-tekh.obr. 18 no.12:26
D '61. (MIRA 14:12)

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sovnarkhoza, Murmansk.
(Murmansk Province--Vocational education)

SLOBOV, H.H.; AKINOV, V.V.; DOROKHOVA, V.S.

Epidemiologic characteristics of tick-borne encephalitis in
Maritime Territory. Med. paraz. i paraz. bel. 33 no.2: 169-177
Mr - Ap '64 (MIRA 18:1)

1. Otdel ontologii (zav. - prof. V.N. Beklemishev [deceased])
Instituta meditsinskoy parazitologii i tropicheskoy meditsiny
imeni Ye.I. Martsinovskogo (direktor - prof. P.G. Sergiyev)
Ministerstva zdoravookhraneniya SSSR, Moskva, i Primorskaya
krayevaya sanitarno-epidemiologicheskaya stantsiya (glavnyy
vrach V.V. Akinov).

L 38466-66 INT(1)/T BK

ACC NR: AP6029183

SOURCE CODE: UR/0016/66/000/005/0008/0013

AUTHOR: Shestakov, V. I.; Mikhoyeva, A. I.; Polenova, I. N.; Dorokhova, V. S. ³³_BORG: Vladivostok Institute of Epidemiology, Microbiology and Hygiene (Vladivostokskiy institut epidemiologii, mikrobiologii i ggiyony); Regional Sanitary Epidemiological Station (Krayevaya sanitarno-epidemiologicheskaya stantsiya)TITLE: Prevention of Japanese encephalitis in Primorskiy Kray

SOURCE: Zhurnal mikrobiologii, epidemiologii i immunobiologii, no. 5, 1966, 8-13

TOPIC TAGS: encephalitis, insect control, mosquito, disease control

ABSTRACT: In Khasanskiy Rayon, where Japanese encephalitis is endemic, systematic measures have been carried out since 1960 to control the mosquito vectors of the disease (*C. tritaeniorhynchus* G., *C. bitaeniorhynchus* G., *C. pipiens* L., *A. togol* Theob., *A. escoensis* Jam.) and to protect the population from mosquito bites. The breeding places were spayed from airplanes with DDT aerosols (10% dust and 50% paste). The best results were obtained by antilarval treatment of the biotopes in the early spring. The people were protected from insect bites with dimethylphthalate, repudin, and diethyltoluamide. The latter proved to be the most effective repellent. Orig. art. has: 3 tables. [JPRS: 36,932]

SUB CODE: 06 / SUBM DATE: 22Jul65 / ORIG REF: 005 / OTH REF: 001

1/1 m, 2 UNK: 616.082.25.022.205.7.021(5.1.62)

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"Certain Methods of Dietetic Therapy of Cardiovascular Patients," from the book Theses of the Reports of the Scientific Session of the Military Medical Academy im. S. M. Kirov, Teziy Doklady Nauchnoy Sessii, 29 Oct-2 Nov, 1956, Leningrad.

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Diet therapy in hypertension. Klin.med. 38 no.12:34-38 D '60.
(MIRA 14:2)

1. Iz kafedry obshchey terapii (nach. - deystvitel'nyy ohlen
AMN SSSR prof. N.N. Savitskiy) Voyenno-meditsinskoy ordena
Lenina akademii imeni S.M. Kirova.

(HYPERTENSION) (DIET IN DISEASE)

SHAKHOVA, Z.F.; DOROKHOVA, Ye.N.

Rapid method of photometric determination of silicon in cast iron and steels. Vest. Mosk. un. Ser. 2; Khim. 20 no.2:77-78 Mr-Ap '65.
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1. Kafedra analiticheskoy khimii Moskovskogo universiteta.

SHAKHOVA, Z.F.; DOROKHOVA, Ye.N.

Formation of phosphomolybdic and germanomolybdic heteropoly acids.
Zhur. neorg. khim. 10 no.9:2060-2064, 8 '65. (MIRA 18:10)

SHAKHOVA, Z.F.; DOROKHOVA, Ye.N.

Photometric determination of chromium based on the fading-out
of silicomolybdenic acid coloration. Vest. Mosk. un. Ser. 2:
Khim. 19 no.5:77-80 S-0 '64. (MIRA 17:11)

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DOROKHOVA, E.M.

Clinical aspects and morphology of brain injuries in peace time.
Vopr. neirokhir. 17 no.61:6-15 N-D '53. (MLRA 6:12)

**1. Iz Leningradskogo nauchno-issledovatel'skogo neyrokhirurgicheskogo
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Khim. prom. [Ukr.] no.1243-44 Ja-Mr*63 (MIRA 17:7)

1. Institut prirodnogo gaza AN UkrSSR.

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tekhn. nauk

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Standardization of portable sprinkling irrigation installations.
Szabvany kozl 15 no.12:272-274 D '63.

DOBOS, Alajos, okleveles mernok, egyetemi adjunktus; DOROMBY, Iapzlo,
okleveles mernok; FEKETE, Istvan, dr., okleveles mernok;
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2. Bacs-Kiskun megyei Allami Gazdasagok Igazgatosaga (for
Doromby, Fekete and Reskovits).

DOBOS, Alajos, okleveles mernok, egyetemi adjunktus, DOROMBY, Laszlo, okleveles mernok; FEKETE, Istvan, dr., okleveles mezogazdasagi mernok; RESKOVITS, Miklos, okleveles mezogazdasagi mernok

Standardization of portable sprinkling irrigation installations.
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1.No.1 Chair of Hydraulic Engineering, Technical University
of Building and Transportation, Budapest (for Dobos).
2.Directorate of Bacs-Kiskun County State Farms (for Doromby,
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BOBOS, Alajos, okleveles mernok, egyetemi adjunktus; DOROMBI, Laszlo, okleveles mernok; FEKETE, Istvan, dr., okleveles mezogazdasagi mernok; RESKOVITS, Miklos, okleveles mezogazdasagi mernok

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1. No. 1 Chair of Hydraulic Engineering, Technical University of Building and Transportation, Budapest (for Bobos).
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