

FOLWARCZNY, Czeslaw, mgr inż.; SOBOTKOWSKI, Witold, mgr inż.; SUCHORAB, Antoni, inż.

Reliability and safety testing of thermal installations in power stations. Pt.1. Energetyka Pol 18 no.3:Supplement: Energopomiar 10 no.2:14-16 Mr'64

1. Pion Ciepłny, Zakład Badan i Pomiarow, Warszawa.

FOLWARCZNY, G.

FOLWARCZNY, G. How a machine-tractor station can considerably increase the yields per  
hectare. p. 244.

Vol. 6, No. 13, July 1956.

MECHANISACE ZEMEDLSTVI.

AGRICULTURE

Praha, Czechoslovakia

So: East European Accession, Vol. 6, No. 3, March 1957

FOLWARCZNY, Czeslaw, mgr inz.; SOBOTKOWSKI, Witold, mgr inz.;  
SUCHORAB, Antoni, inz.

Methods of performing acceptance tests of thermal equipment in  
power plants. Pt. 2. Energetyka Pol 18 no. 1 [i.e.5]:Suppl. Bl 1  
nauk techn energopomiar 10 no. 3:17-23 My '64.

FOLMARČNY, G.

Making work with contracts easier. p. 450. (MECHANISACE ZEMĚDELSTVI,  
Vol. 6, No. 23, Dec 1956, Praha, Czechoslovakia)

SO: Monthly List of East European Accessions (EEAL) LC, Vol. 6, No. 12, Dec 1957. Uncl.

COUNTRY : Poland ii-2  
CATEGORY : Chemical Technology. Chemical Products and Their  
Applications--Chemical engineering.  
ABS. JOUR. : RZKhim., No. 5 1960, No. 18054  
AUTHOR : Ochoduzsko, S. and Folwarczny, J.  
INST. : Not given  
TITLE : Adiabatic Flow of Gases in Pipes  
  
ORIG. PUB. : Zesz Nauk Politechn Slaskiej, No 19, 3-12 (1959)  
ABSTRACT : Using the first law of thermodynamics and the con-  
tinuity equation, the authors have derived an equa-  
tion which corresponds to the Fanno [Fannot?]  
curve. On the basis of the equation derived, the  
work required for the overcoming of friction and  
the increase in kinetic energy are discussed in  
P-V and T-S coordinates. V. Sokol'skiy

CARD: 1/1

BARTEL'S, A.V.; LEBEDEVA, M.A.; GRASHCHENKOVA, Z.P.; FOMCHENKO, I.V.

Use of staphylococcal anatoxin in the treatment of mastitis.  
Akush. i gin. 40 no.1:17-21 Ja-F '64. (MIRA 17:8)

1. Nauchno-issledovatel'skiy institut akusherstva i ginekologii (dir. - prof. O.V. Makeyeva) Ministerstva zdravookhraneniya SSSR, Moskva.

24.5200

P/032/62/009/002/001/001  
D265/D308

AUTHOR: Folwarczny, Józef, Gliwice

TITLE: Thermodynamic similarity in the case of laminar flow through a straight circular tube

PERIODICAL: Archiwum budowy maszyn, v. 9, no. 2, 1962, 191-238

TEXT: The laminar flow of heat is analyzed under conditions of hydrodynamic and thermodynamic stability i.e. for constant values of Nusselt number. Taking axial heat conduction into consideration, the differential equation in dimensionless form is derived from first principles

$$\frac{d^2 \varepsilon}{d\sigma^2} + \frac{1}{\sigma} \frac{d\varepsilon}{d\sigma} - \varepsilon \left[ K(1 - \sigma^2) - \left( \frac{K}{Pe} \right)^2 \right] = k \left[ K(1 - \sigma^2) - \left( \frac{K}{Pe} \right)^2 \right], \quad (23)$$

for which the solution shows the relationship  $Nu = f(K, Pe)$  where  $K$  - appears as a new dimensionless quantity defined by

Card 1/2

Thermodynamic stability in the case ...

P/032/62/009/002/001/001  
D265/D308

$$K = - Pe \frac{R}{L} . \quad (21)$$

where R - tube radius, L - constant length of subtangent to a curve of the temperature difference as a function of tube length, Pe - Peclet number. Neglecting the axial heat conduction, the values Nu = 3.65 for K = 7.313 and Nu = 4.36 for K = 0, are shown on Nu - K system graph. Considering the external heat flow through an annular duct around the inner tube the heat exchange conditions are analyzed which depend on the coefficient of heat conductivity, the wall thickness, thermal capacity of the outer flow and the cross-section of the outer duct. Particular cases of heat flow for which K = 0 and Nu = 4.36 and K ≠ 0 and Pe ≈ ∞ are discussed in detail, and heat exchange conditions are determined for several values of Nu - number of the inner tube. There are 11 figures and 5 tables. ✓c

ASSOCIATION: Katedra teorii maszyn cieplnych politechniki śląskiej  
(Department of the Theory of Heat Engines, Silesian Polytechnic

SUBMITTED: December, 1961  
Card 2/2



FOLWARCZNY, J., dr. ing. (Poland)

Control of complex combustion processes. Ipari energia 5 no.3:  
55 M '64.

WILIMARCZYNY, Jozef (Gliwice)

Control of complex combustion processes. Archiw bud masz 11  
no. 1:59-72 '64.

FOLYADOV, V. and TISHCHENKO, D.

"New Type of Terpene Transformations. VIII. Synthesis of Terpenylacetoacetic and Terpenylmalonic Esters and Their Cleavage Products," Zhur. Ob. Khim., 22, No.6, 1952

TISHCHENKO, D.; FOLYADOV, V.

New type of terpene transformations. X. Synthesis of homoterpenes. Zhur.  
Obshchey Khim. 22, 1591-3 '52. (MIRA 5:9)  
(CA 47 no.18:9306 '53)

TISHCHENKO, D.; PERSIANTSEVA, N.; FOLYADOV, V.

New type of terpene transformations. XII. Synthesis of nitriles of the homo-  
terpene series. Zhur. Obshchey Khim. 22, 1829-32 '52. (MLRA 5:11)  
(CA 47 no.18:9307 '53)

FOM, P. Ye.

"Special Tongs for Sealing Diamonds in a Mandrel by means of a Voltaic Arc",  
Stanki I Instrument, 14, No. 11-12, 1943.

BR-52059019

MOYCHO, W.; GUBANSKI, M.; FOMAIDIS, B.; LEMANSKA, M.; WAJSBARD, E.

The occurrence of tobacco mosaic virus in tomatoes in Lodz and its  
neighborhood. Postepy nauk roln 7 no.1:79-82 Ja/F '60. (EEAI 9:10)  
(Poland--Tomatoes)  
(Mosaic disease)  
(Viruses)

FOMCHENKO, A., podpolkovnik militsii

Our common cause. Za rul. 17 no.1:8-9 Ja '59. (MIRA 12:3)

1. Predsedatel' ekzamenatsionnoy komissii Gosudarstvennoy avtomobil'noy  
inspeksii UVD Orenburgskogo oblispolkona.  
(Orenburg Province--Automobile drivers)



FOMCHENKO, G.K., general-mayor meditsinskoy sluzhby; KOVALEV, Ye.I.,  
polkovnik meditsinskoy sluzhby; ASKEROV, A.A.

Electrocardioscopic and electrospygmographic study of the functional  
state of the cardiovascular system. Voen.-med.zhur. no.10:31-35  
0 '59. (MIRA 13:3)

(ELECTROCARDIOGRAPHY)

PHASE I BOOK EXPLOITATION SOV/5304

Soveshaniye po teorii liternykh professorov. 5th, 1959  
Technost' otivok; trudy sovetskaniya (Accuracy of Castings; Transactions of the Fifth Conference on the Theory of Founding Processes) Moscow, Mashgiz, 1960. 206 p. 3,500 copies printed.  
Sponsoring Agency: Akademiya nauk SSSR. Institut mashinovedeniya. Tomskiya po tekhnologii mashinostroyeniya.

Ed. (title page): B. B. Gulyayev, Doctor of Technical Sciences, Professor; Ed. of Publishing House: G. M. Soboleva; Tech. Ed.: A. P. Uvarova; Managing Ed. for Literature on Hot-Processed Metals: S. Ya. Golovin, Engineer.

PURPOSE: This book is intended for scientific and technical personnel at scientific research institutes, factories, and schools of higher education.

COVER: The book contains 19 reports read at a conference on the accuracy of castings. The conference was organized by the Committee on Processing in Machine Building and sponsored by the Institut mashinovedeniya AN SSSR (Institute of the Science of Machines of the Academy of Sciences USSR). The reports, presented by leading specialists, science workers, and production personnel, discuss the present state of the problem of the accuracy of castings and methods of solving the problems involved. There are 50 references, mostly Soviet.

Kumarov, L. Ye. (Engineer). Distortion of Sand Molds

Zhukovskiy, S. S. (Engineer), and V. Ts'yan-chen (Engineer). Dimensional Errors of Castings Caused by Patterns and Flasks 125

Dubrovskiy, A. M. (Engineer). Effect of Thermal Distortion of the Moldlin Mixtures on the Accuracy of Castings  
The work of investigating the distortions and thermal stress in the molding mixtures was carried out under the supervision of P. P. Berg. 131

Fomchenko, S. I. (Engineer), and B. B. Gulyayev. Production of Precision Castings in Shell Molds Filled From a Water-Glass Mixture 146

Kolchin, I. P. (Engineer), and V. V. Krzhenkov (Engineer). Production of Large Precision Steel Castings By Using Chemically Hardening Mixtures 153

Rubtsov, M. N. (Doctor of Technical Sciences, Professor), and I. L. Zaitkov (Engineer). Dimensional Accuracy of Investment Castings 160

Goryunov, I. I. (Candidate of Technical Sciences). Dimensional Accuracy and Surface Roughness of Castings Obtained by Various Methods  
C. A. Kantor, A. Ye. Danilov, A. I. Belyayev, and Engineer V. B. Shul'man participated in making castings. 180

Iskakov, M. E. (Engineer), and B. B. Gulyayev. Formation of the Contours of Castings in Die Casting 193

Kolesnichenko, A. G. (Engineer). Accuracy of Castings Obtained in Metal Molds 203

Card 6/7

11

YEVSTAF'YEV, I.N., inzh.; BOROVSKIY, Yu.P., kand. tekhn. nauk; FOMCHENKO,  
S.I., kand. tekhn. nauk; GULYAYEV, B.B., doktor tekhn. nauk

Compacting molding mixtures by vibration squeezing. Lit. prosv.  
no.9:4-6 S '65. (MIRA 18:10)

По м.ч.е.н.к.о.в.н.

None given  
 A Conference on the Accuracy of Machine Building Castings  
 Investiya Akademiya nauk SSSR, Otdelnyye tekhnicheskikh  
 nauk, Metallurgiya i toplivo, 1959, Nr 4, pp 255-256 (USSR)  
 A conference on the above subject took place in the  
 Institute of Machine Building of the Academy of Sciences  
 of the USSR on 23-24 April 1959. About 300 representatives  
 of scientific-research institutes, laboratories,  
 universities and large enterprises from 31 towns participated  
 in the conference. The following papers were read:  
 B.B. Gulvayev "The present state of knowledge of the  
 accuracy of castings"; P.N. Akhmetov "Methods of  
 investigations of the dependence of the accuracy of  
 castings on technological factors"; N.F. Berz "Methods of  
 analytical evaluation of dimensions of castings";  
 V.A. Babitskiy "Theoretical and experimental investigations  
 of the accuracy of castings"; I.F. Iosorokhuy - "The  
 system of allowances for mechanical working of castings";  
 Ye.G. Kononovich "Methods for the determination of  
 tolerances for dimensions of cast parts"; S.A. Lazrenko  
 "Tolerances for non-ferrous cast parts"; S.A. Lazrenko  
 "Methods of castings"; G.N. Nikolayev "Method of  
 controlling the cleanliness of the surface of castings";  
 I.S. Konstantinov "The influence of stresses formed during  
 casting on the accuracy of castings"; I.Ye. Korotok "The  
 process of packing moulds as a factor determining the  
 accuracy of castings"; S.S. Zhukovskiy and  
 M. Zhukovskiy "Sources of errors in the dimensions  
 of castings caused by specific features of operation of  
 the pattern-mould boxes equipment"; A.Ye. Borozdyuk  
 "Conditions of the casting of accurate castings in sand moulds";  
 M.K. Jozanov "The influence of the chemical composition of  
 iron on the accuracy of dimensions of castings";  
 S.N. Kuznetsov and B.N. Zhurav "Improvement of the  
 accuracy of castings made in air-cast moulds";  
 V.Y. Nerzhov "Experience in increasing the cleanliness  
 and accuracy of large castings"; V.N. Radlov and  
 I.I. Zhelikov "On the accuracy of castings made by the  
 lost wax method"; I.Ye. Zhurav "An investigation of the  
 accuracy and surface cleanliness of castings made under  
 pressure"; V.Ye. Zhurav "The accuracy of castings made  
 during casting under pressure"; I.Ye. Zhurav, A.I. Zhurav  
 and S.S. Zhukovskiy "An investigation of the accuracy of  
 castings made under pressure by form surface quality  
 of the pressure moulds". It was established that studies on  
 the subject of the accuracy of castings are developing  
 too slowly mainly due to lack of coordination in the  
 research work and insufficient numbers of specialists in  
 the field of mathematics, physics and electronics. In  
 order to develop methods for overall calculations of the  
 accuracy, productivity and economics of casting processes  
 the authors recommended organizing a scientific center  
 and a new (at present) research institute in Moscow, involving  
 universities and scientific institutions of various  
 specialists, mathematicians, physicists and economists.

Card 1/3

Card 2/3

FORCHENKOV, G. A.

Dispensary Storage Table PriP. Voenno-Meditsinskiy Zhurnal, No 1, p. 76, 1955.

VOROB'YEV, V., inzh.; ZHUNIN, A., inzh.; SPIROV, V., inzh.;  
FOMCHENKOV, I., inzh.

Building made of light alloys. Na stroi. Ros. no.10:37-38  
0 '61. (MIRA 14:11)  
(Moscow--Aluminum, Structural)

FOMCHENKO, Oleg[Fomchanka, Aleh] (Brest)

People praise her work. Rab. i sial. 39 no.2:15-16 F '63.  
(MIRA 16:4)

(Juvenile delinquency)

GAL'PERIN, Ye.I.; GORDONOV, A.Yu.; POMCHENKOV, V.M.

Designing trigger circuits for point contact crystal triodes with  
consideration of interchangeability. Poluprov. prib. i ikh prim.  
no.2:340-352 '57. (MIRA 11:6)  
(Transistors) (Electronic circuits)



MOLCHANOV, A.A. (Novosibirsk); SIGORSKIY, V.P. (Novosibirsk);  
FOMEL', B.M. (Novosibirsk)

Study of the dynamics of multistable elements based on a  
simplified model. Izv. AN SSSR. Tekh. kib. no.5:156-162  
S-0 '65. (MIRA 18:11)

ACC NR: AR7004318

SOURCE CODE: UR/0271/66/000/011/B016/B017

AUTHOR: Fomel', B. M.

TITLE: Investigation of subharmonic oscillations in computer elements with one degree of freedom

SOURCE: Ref. zh. Avtomat. telemekh. i vychisl. tekhn., Abs. 11B120

REF SOURCE: Sb. Vychisl. sistemy. Vyp. 21. Novosibirsk, 1966, 33-45

TOPIC TAGS: computer design, computer component, computer theory, *analog computer, subharmonic oscillation, oscillation / IPT-5 analog computer*

ABSTRACT: Elements that use parametric subharmonic oscillations have found wide usage at the present time; the phase of the subharmonic oscillation serves as a dynamic state indicant in these elements. As nonlinear systems have no principal difference between an external periodic disturbance and a parametric effect (because every periodic process, in such a system, is accompanied by a parametric modulation of its nonlinear elements), it is suggested that the oscillations of any shape whose frequency is submultiple of the external applied frequency be considered as subharmonic oscillations. A class of dynamic computer elements is considered which is describable by second-order equations containing, in addition to a strongly nonlinear function, a periodic function of this form:  $\ddot{x} + b\dot{x} + \sigma(x) = f(t)$  where  $b$  - positive coefficient,  $\sigma(x)$  - nonlinear function,  $f(t)$  - periodic function. Qualitative methods are used in an examination of such an equation which

Cord1/2

UDC: 681.142.32.001

ACC NR: AR7004318

permits finding all possible motions in the system. The results were verified on an IPT-5 analog computer equipped with a set of nonlinear units. The results were readout from an oscilloscope screen calibrated in  $(x, y)$ -coordinates of the state plane and having time markings. The above qualitative method of studying subharmonic oscillations has the advantage over other methods in that it is noncritical to the form of the nonlinear function contained in the initial differential equation and in that it permits obtaining the entire set of possible motions simultaneously. In addition to presentation of the general method, a case of single-domain model of a ferromagnetic film is considered in detail; the model is constructed after the Landau-Livshits-Gilbert equation which describes the magnetic moment in a ferromagnetic:  $n^2\ddot{x} + \beta n\dot{x} + \frac{1}{2} \sin 2x + f(t) \sin(x - \theta_0) = 0$ , where  $x$  - magnetic-moment angle in the external field,  $n$ ,  $\beta$ ,  $\theta_0$  - film parameters,  $f(t)$  - external field varying on the same law as the function  $f(t)$  in the initial equation. Eight figures. Bibliography of 3 titles. G. V. [Translation of abstract]

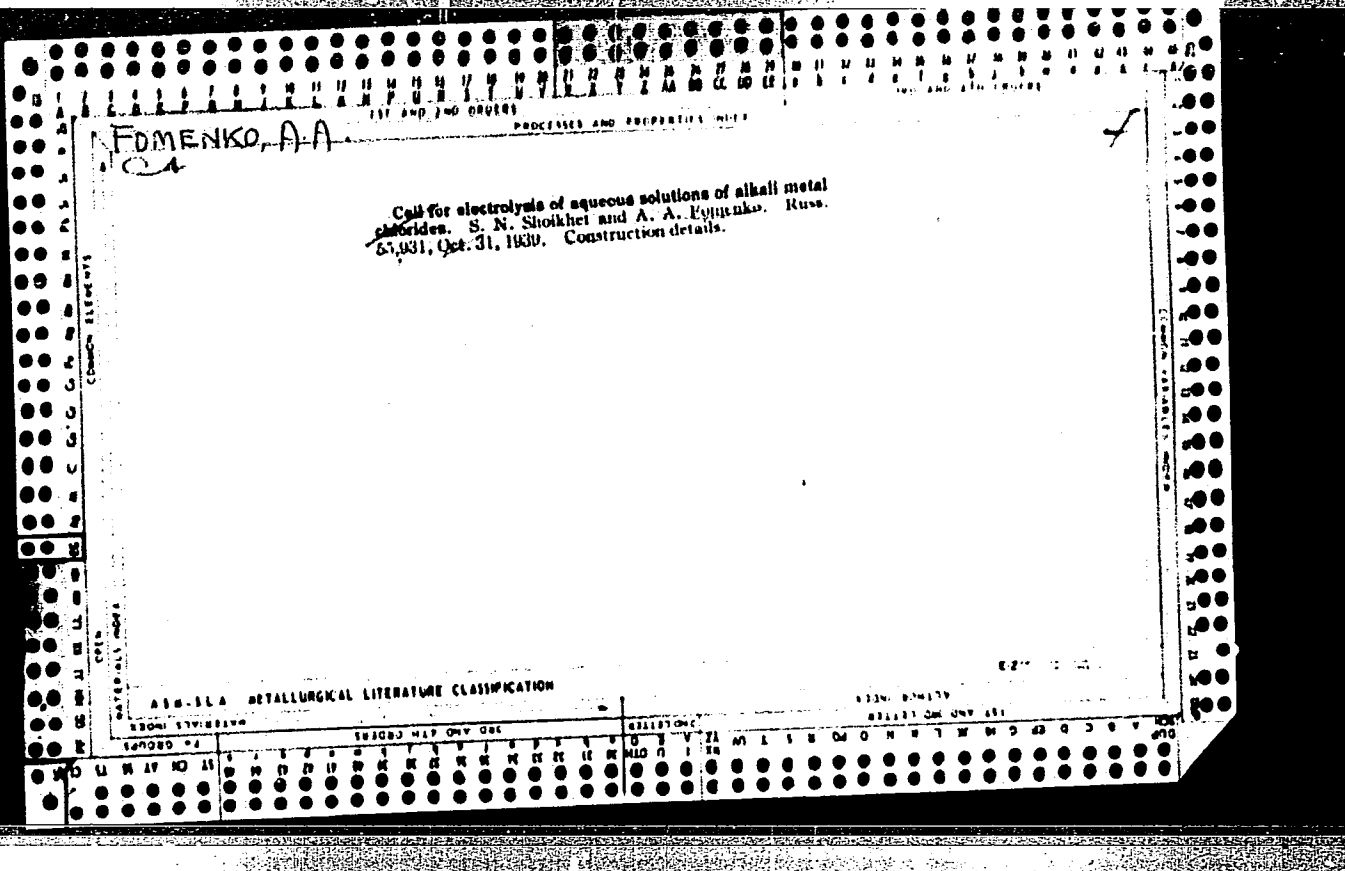
SUB CODE: 09

Card 2/2

BARKHIN, G.S.; FOMEN, V.T.

Bending of surfaces of positive curvature under certain  
boundary conditions. Dokl. AN SSSR 140 no.5:987-989 0  
'61. (MIRA 15:2)

1. Rostovskiy-na-Donu gosudarstvennyy universitet. Predstavleno  
akademikom P.S.Aleksandrovym. <sup>7</sup>  
(Surfaces of ~~constant~~ curvature)  
(Boundary value problems)



FOMENKO, A.A.

Radiofrequency magnetic and electric spectra of permeability of the  
body and substance of MnZn-ferrites. Zhur. tekhn. fiz. 28 (MIRA 11:4)  
no.3:506-510 Mr '58.  
(Ferrites) (Magnesium-zinc alloys--Electric properties)

RUSSIA, A. G.

"Use of preparations FB-1 and FB-2."  
SO: Veterinariya 28(2), 1951, p. 46.

S/263/62/000/020/002/006  
E194/E135

AUTHORS: Agarkov, A.P., Fomenko, A.I., and Kureyko, S.M.  
TITLE: The preparation and calibration of heat resistant pick-ups  
PERIODICAL: Referativnyy zhurnal, otdel'nyy vypusk, Izmeritel'naya tekhnika, no.20, 1962, 11, abstract 32.30.91. (Tr. Novocherk. politekhn. in-ta, 127, 1961(1962), 57-61).

TEXT: Heat-resistant pick-ups based on potassium glass are at present widely used to measure static stresses that are set up in components which operate in a high temperature zone. Cements for heat resistant pick-ups based on potassium liquid glass are of good technological properties, their sole disadvantage being their poor electrical insulating properties at high temperature. This disadvantage is overcome by introducing lead oxide PbO which reduces the ionic conductivity of the cement because of the influence of heavy ions. The procedures for making and calibrating the heat-resistant pick-ups are described. 3 figures.

Card 1/1 [Abstractor's note: Complete translation.]



1. FOMENKO, A.N., INOZEMTSEV, Ye K.
2. USSR (600)
4. Lignite - Kagul County
7. Geological report on the results of the prospecting carried out by the Izmail and Kagul lignite parties in 1940-1941. (Abstract) Izv.Glav.upr.geol.fon. no.2 1947
  
9. Monthly List of Russian Accessions. Library of Congress. March 1953. Unclassified.

A-1

BC

**Kinetics of the electrode process at an ammonia gas electrode. A. S. FOMENKO (J. Phys. Chem. Russ., 1938, 11, 878-884).—The potential change  $\Delta\epsilon$  of platinum Pt in eq.  $\text{NH}_4\text{NO}_3$  saturated with  $\text{NH}_3$  is connected with the e.d. of the polarizing current by the equation  $\Delta\epsilon = a - b \log I$ ,  $a$  and  $b$  being constants. Between 0° and 30°  $b$  rises from 0.034 to 0.039; its val. suggests that  $\text{NH}_3 + 3\text{OH}' \rightarrow \text{N} + 3\text{H}_2\text{O} + 3\ominus$  is the slowest reaction at the electrode. J. J. B.**

ASB-3LA METALLURGICAL LITERATURE CLASSIFICATION

GROUP	CLASS	SECTION	RELATIONS
1	2	3	4
5	6	7	8
9	10	11	12
13	14	15	16
17	18	19	20
21	22	23	24
25	26	27	28
29	30	31	32
33	34	35	36
37	38	39	40
41	42	43	44
45	46	47	48
49	50	51	52
53	54	55	56
57	58	59	60
61	62	63	64
65	66	67	68
69	70	71	72
73	74	75	76
77	78	79	80
81	82	83	84
85	86	87	88
89	90	91	92
93	94	95	96
97	98	99	100

9

*Ca*

**Retarding effect of inhibitors in the corrosion of metals.**  
 A. S. Fomenko and O. K. Skarre. *J. Phys. Chem. (U. S. S. R.)* 19; 1297-1303 (1939).—The hydrogen overvoltage on an Fe electrode in H<sub>2</sub>SO<sub>4</sub> soln. is increased by the corrosion inhibitor sulfurous anthracene oil, in a conc. up to 0.2%. The anode polarization curves of Fe in FeSO<sub>4</sub> or H<sub>2</sub>SO<sub>4</sub> soln. are shifted by in the direction of a positive potential of the iron electrode by about 70 mv. The current strength of the Fe-Pt pair for short contacts decreases from 0.0 ma. to 0.7 ma. for 0.3% of the inhibitor and is unaffected by further additions. F. H. Rathmann

*Instit. Phys. Chem. im Pishchevskiy, 45 Ukr. SSR, Dnepropetrovsk*

A.S.M.S.A. METALLURGICAL LITERATURE CLASSIFICATION

RECORDS SECTION

SEARCHED INDEXED SERIALIZED FILED

APR 1960

FOMENKO, A. S.

Stability of hydrogen atoms in free trichloromethyl-  
A. S. Fomenko and E. A. Sadovnikova (Acad. Sci.  
Ukrain. S.S.R., Kiev). J. Phys. Chem. (U.S.S.R.) 22,  
423-5(1948)(in Russian).--It was expected that the high  
electron charge  $d$ , in the  $\alpha$ - and  $\beta$ -positions of the benzene  
nuclei of  $(C_6H_5)_3C$  would render the H atoms there more  
mobile. Solns. of 20 g. of hexaphenylethane in 100 cc.  
 $C_6H_6$  were heated with heavy water at  $100^\circ$  for 20-150  
hrs. The fact that hexaphenylethane recovered (as per-  
oxide) contained only 2.5% of deuterium showed that the  
expectation was not fulfilled. Expts. using heavy acetone  
yielded similar results. J. J. Bikerman

102 AND 21st CROSS

107 AND 2ND CROSS

PROCESSES AND PROPERTIES INDEX

BC

A 2  
3

Microkinetics of the Claisen rearrangement. A. S. Fompahe, G. P. Mikheev, and L. A. Izrael'skaya (C. R. Acad. Sci., U.R.S.S., 1948, 1949). The Claisen rearrangement of allyl vinyl ether (OPE) to gamma-delta unsaturated ether (OPE) was studied. It was found that the reaction is first order in OPE and zero order in the catalyst. The reaction is an intramolecular process involving dissociation of the ether moiety. E. Truscov.

ASS-SLA METALLURGICAL LITERATURE CLASSIFICATION

107 AND 2ND CROSS

102 AND 21st CROSS

107 AND 2ND CROSS

102 AND 21st CROSS

FOMENKO, A. S.

Ashinazi, M. G., Kutsaya, B. F., and Fomenko, A. S. - "A simplified method of determining the contrast factor", (Of photographic plates), Ukr. khim. zhurnal, Vol. XIV, Issue 2, 1949, 47-52

SO: U-7392, 19 August '53, (Letopis 'Zhurnal 'nykh Statey, No. 21, 1949).

CA

10

**Mechanism of the Claisen rearrangement.** A. S. Ponomarev and E. A. Salovnikova (L. V. Pisarshev Inst. Phys. Chem., Acad. Sci. Ukr. S.S.R.). *Zhur. Obshchei Khim.* (J. Gen. Chem.) 20, 1898-1903(1950); cf. *C.A.* 43, 602i.—The existing data on investigations of the reaction are critically reviewed and it is pointed out that at this time no conclusive evidence exists for a final decision as to an intra- or intermol. mechanism of the primary step. The ortho-rearrangement of  $\text{CH}_2=\text{CHCH}_2\text{OPh}$  in the presence of phenol-3,5- $D_2$  by refluxing in a N atm. 6 hrs. gave 65% *o*-allylphenol, isolated either by extrn. with 20% NaOH, acidification, extrn., and fractionation, or by pptn. of allylphenol with  $\text{HgCl}_2$  as 2-(chloromercurimethyl)-2,3-dihydrobenzofuran; combustion of the product showed the absence of *o*-allylphenol-3,5- $D_2$ . A similar result was obtained on refluxing  $\text{PhCH}=\text{CHCH}_2\text{OPh}$  in the presence of allyl phenyl-3,5- $D_2$  ether 4 hrs. in a  $\text{CO}_2$  atm. Thus, the ortho rearrangement in these cases takes place initially by an intramol. route. The phenol-3,5- $D_2$  was prepd. according to Hrodski, *et al.* (*C.A.* 44, 8882e); its allyl ether, prepd. from allyl bromide as usual (cf. Houben, *Die Methoden der org. Chemie*, vol. 111, *C.A.* 24, 1389), bp 104-6°. *o*-(1-Phenylallyl)phenol bp 178.5-7.5°. G. M. K.

APPROVED FOR RELEASE



*FOMENKO, H. S.*  
ROZANOV, Ivan Grigor'yevich, starshiy nauchnyy sotrudnik; ZAVITAYEV, Petr  
Aleksseyevich, starshiy nauchnyy sotrudnik; SKATKIN, M.M., redaktor;  
FOMENKO, A.S., redaktor; DZHATIYEV, S.G., tekhnichaskiy redaktor

[Handicraft lessons for the fourth grade] Uroki ruchnogo truda v  
chetvertom klasse. Pod red. Skatkina. Izd. 2-oe, dop. i perer.  
Moskva, Gos.uchebno-pedagog. izd-vo M-va prosv. RSFSR, 1956. 231 p.  
(MIRA 10:11)

1. Institut teorii i istorii pedagogiki (for Rozanov). 2. Institut  
metodov obucheniya Akademii pedagogicheskikh nauk RSFSR (for  
Zavitayev). 3. Chlen-korrespondent Akademii pedagogicheskikh nauk  
RSFSR (for (Skatkin)  
(Handicraft)

from entrained H<sub>2</sub> and consequently the  
the process repeated. The tests were run with iron filings

FOMENKO, A.

73-2-21/22

AUTHOR: Reported by A. Fomenko.

TITLE: Second Ukrainian Conference on Physical Chemistry (Symposium). Vtoroye Ukrainskoye Respublikanskoye Soveshchaniye po Fizicheskoy Khimii).

PERIODICAL: "Ukrainskiy Khimicheskiy Zhurnal" (Ukrainian Journal of Chemistry), Vol.23, No.2, March-April, 1957, pp.272-275 (USSR).

ABSTRACT: Held December 14-18, 1956 in Kiev by the Institute of Physical Chemistry im. L.V.Pisarzhhevskii (Institut Fizicheskoi Khimii im.L.V.Pisarzhhevskogo, AN USSR). Short summaries are given of the following papers:

Theory and Practice of Catalytic and Adsorption Phenomena.

Roiter, V.A., Korneichuk, G.P., Rzayev, P.B. and Stukanovskaya, N.A. (Kiev):

Results on investigations on the kinetics of oxidation of sulphurous anhydride to sulphuric anhydride by industrial vanadium catalysts by the diaphragm method.

Card 1/20 Values for the velocity constants and heat of activation were computed. They were not affected by the influence of macrofactors.

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Theory and Practice of Catalytic and Adsorption Phenomena.

(Cont.)  
Arnol'dov, E.M., Lyubiteleva, A.Z. and Vol'fson, V.Ya. (Rubzhnye).  
Investigations on a composite charge of a vanadium oxide  
catalyst for the oxidation of naphthalene. This was pre-  
sented by IFX AN USSR.

Roiter, V.A., Stukanovskaya, N.A., Yuza, V.A., Korneichuk, G.P. and  
Ushakova, V.P. (Kiev).

These authors observed that many processes are catalysed by  
impure vanadium oxide. Oxygen interchange is revealed only  
above 500 C, in spite of the fact that catalytic oxidation  
occurs between 300 - 400 C. This is contradictory to the  
oxidation-reduction mechanism of the catalysis.

Rusov, M.T. and Strel'tsov, O.A. (Kiev).

Data obtained by investigating the macrokinetics of the  
synthesis of ammonia in a long catalyst bed at high  
pressures.

Davtyan, O.K. and Ovchinnikova, E.N. (Odessa).

Oxidation of sulphurous anhydride on the surface of a

Card 2/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Theory and Practice of Catalytic and Adsorption Phenomena.  
(Cont.)

series of solid catalysts and on activated coal proceeding  
at normal temperatures.

Stepko, I.I. and Goncharenko, G.F. (Kiev).

They observed a decreased discharge of electrons from  
germanium during adsorption of CO and CO<sub>2</sub> and formic acid  
vapours. As in the case of CuO, the catalytic oxidation  
of CO on germanium causes a decrease of electron emission.

Glikman, T.S. and Podlinyaeva, M.E. (Kiev).

Card 3/20 The authors proved that sensitised photo-reduction (by  
zinc oxide) of methylene blue and thionine can be accom-  
plished at the expense of oxidation of OH-ions from zinc  
hydroxide. Thus the oxidation of water can be achieved  
with dyes by the effect of small light quanta in the  
presence of ZnO.

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Theory and Practice of Catalytic and Adsorption Phenomena.

Kornienko, T.P. and Polyakov, M.V. (Kiev).

Oxidation reactions of methane and methanol to formaldehyde.

The oxidation reactions of methane were proved to start on walls of the reaction vessel.

Shalya, V.V. and Polyakov, M.V. (Kiev).

The authors discussed results obtained during the investigation of the kinetics of oxidation of butane-propane fractions and of methanol by differential calorimetric measurement. A heterogeneous-homogeneous radical-chain mechanism of the process was established.

Tovbin, M.V. and Savinova, Ye.V. (Kiev).

They investigated the kinetics of the non-stationary evaporation of water from the surface of a vibrating string which is in contact with the gas. The mechanism of the process consists in the formation of transient self-adsorptive layer and of the desorption of substances from this layer. In a further discussion they investigated the mechanism of self-adsorption on the liquid-gas boundary.

Card 4/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Theory and Practice of Catalytic and Adsorption Phenomena.  
(Cont.)

Rastrenenko, A.I. and Neymark, I.Ye. (Kiev).

The production of titanium silicagel, titanium-gels and alumino-silica gel with different pore structure was investigated as well as the influence of the macro-structure of these sorbents on their catalytic properties.

I.E.Neymark's theory on the formation of the structure of hydrophilic sorbents can be applied.

Sheynfayn, P.Yu. and Neymark, I.Ye. (Kiev).

Results of investigations on the adsorption of various substances on fluorinated silicagels.

Adsorption is linked with the formation of molecular complexes and proceeds on the basis of the acceptor-donor mechanism.

Tarkovskaya, I.A., Chervyatsova, L.L. and Strazhesko, D.N. (Kiev),

presented a paper on the mechanism of adsorption of electrolytes with the aid of radioactive isotopes. In aqueous as well as in aqueous-organic media the selective absorption of salt cations represents a typical ion-exchange process.

Card 5/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Theory and Practice of Catalytic and Adsorption Phenomena.  
(Cont.)

Davydov, A.I. (Kharkov).

The mechanism of the poly-anion exchange.

Shostenko, Yu. V. (Kharkov).

Results of investigations of the dynamics of adsorption  
and desorption of organic substances.

Strazhesko, D.N. (Kiev).

Results of investigations in the field of electrochemical  
adsorption of ions on carbon from aqueous and non-aqueous  
solutions.

The mechanism of the influence of non-electrolytes on the  
adsorption of electrolytes by active carbon was formulated.

This amplifies the theory of electrochemical adsorption as  
stated by A.N.Frumkin.

Theory of Solutions of Electrolytes and Non-Electrolytes.

Esel'son, B.N. (Kharkov).

Card 6/20 The author discussed the vapour-liquid diagrams of  $\text{He}^3$  in  $\text{He}^4$   
near to the transition point  $\text{HeI} \rightarrow \text{HeII}$ .



73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Theory of Solutions of Electrolytes and Non-Electrolytes.  
(Cont.)

Lifshits, I.M. (Kharkov).

The lecturer elucidated the quantum theory of liquid solutions and the theory of phase transitions of type II. by application to the transformation of He I in He II. The theory is proved by investigations on the separation of the solutions into 2 phases, He<sup>3</sup> and He<sup>4</sup>. The quantum theory of solutions explains the discrepancy of the solutions He<sup>3</sup> in liquid He<sup>4</sup> from ideal compositions.

Izmaylov, N.A. (Kharkov).

The influence of solvents on the strength of electrolytes. The dissociation of any electrolytes is considered to proceed in a number of successive stages: the interaction of the electrolyte with the solvent with the formation of addition products, its dissociation into ions as a result of further solvation and association of ions into pairs of ions. The influence of the solvents on the properties of electrolytes is determined quantitatively by an equation

Card 7/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Theory of Solutions of Electrolytes and Non-Electrolytes.  
(Cont.)

put forward by the author.

Mikulin, G.I. (Donsoda).

Critical review of the contemporary theories on electrolytes.

The author's thermodynamic theory on hydration compares the theory of electrolytes with Mendeleev's hydration theory. On the basis of experimental data it was stated that the molar energy of the interaction of particles of concentrated electrolytes is in a linear proportion to  $\sqrt{c}$ . The chemical composition and the constants of instability of liquid hydrates has to be defined by the quantitative theory of solutions as well as the physical theory elucidating the linear dependence of the energy on  $\sqrt{c}$ .

Golik, A.Z. (Kiev).

Investigations on the viscosity, density, heats of evaporation and other properties of molecular-mixed solutions the components of which have similar structures.

Card 8/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Theory of Solutions of Electrolytes and Non-Electrolytes.  
(Cont.)

Roshchinaya, G.P. (Kiev).

The author investigated the molecular light diffusion in solutions. Results are in good agreement with those obtained by A.Z.Golik during the investigations of the physical properties of liquid solutions.

Urazovskiy, S.S. and Kuris'ko, A.I. (Kharkov).

They discussed the anomalies of the thermal relation of the refractive index for menthol and benzophenone in points corresponding to the melting points of their stable modifications. In the case of cis-decahydronaphthalene the melting point is considerably higher. These anomalies can be explained by the change of the coordination number in the structure of the neighbouring order during the transition from the liquid to the supercooled state. The anomaly in the thermal dependence of the refractive index for cis-decahydronaphthalene can be explained by the reverse isomery which characterises the cyclic compounds of the saturated series.

Card 9/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Theory of Solutions of Electrolytes and Non-Electrolytes.  
(Cont.)

Izmaylov, N.A. and Franke, A.K. (Kharkov).

Investigations on the reactions between butyl alcohol and carboxylic acids in benzene, and calculation of the yield of complex products of the reaction.

The compounds have the composition  $AB_2$  (where A = acid and B = alcohol). The constants of non-stability and reaction energies were calculated from data of their yield. Acids with acetone, acetonitrile and nitrobenzene give compounds corresponding to the formula AB.

Izmaylov, N.A., Aleksandrov, V.V., and Ivanova, E.F. (Kharkov).  
Influence of the chemical and physical nature of the solvent and of the concentration of the electrolyte on the state of energy of ions in solutions.

Markov, B.F. and Shimina, L.A. (Kiev).

It was shown that the electric conductivity of binary dissolved salt mixtures, the components of which do not form compounds, does not appear to be an additive property.

Card 10/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Theory of Solutions of Electrolytes and Non-Electrolytes.  
(Cont.)

This digression is more marked during the transition from continuous solid solutions to congruently melting compounds.

Gorenbeyn, Ye.N. (Kiev).

The influence of cations and anions on the molecular composition of electrolytes in solvents with low dielectric constants.

Rapshtynskaya, Ye.A. and Skarre, O.K. (Dnepropetrovsk).

Dielectric constant of pure liquids at various temperatures. The authors found that the dielectric constant is a function of their energy of orientated interaction. Assuming that this applies also to mixtures the authors calculated the energy of orientated interaction for 10 solutions at various temperatures.

Vasenko, Ye.N. and Blank, M.G. (L'vov).

Investigation on the mutual solubility in ternary systems: formamide-acetone in benzene, toluene and carbon tetrachloride.

Card 11/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Theory of Solutions of Electrolytes and Non-Electrolytes.  
(Cont.)

Ravikovich, S.D. (Kiev).

Relations between the molecular structure and density of  
organic liquids.

Photochemistry.

Dain, B.Ya. and Ashkinazi, M.S. (Kiev).

Results of investigations on the photochemistry of hemin-  
resembling derivatives of chlorophyll.

With the aid of these pigments it was possible to show one  
of the stages of photosynthesis outside the plant cell:  
the photodecomposition of water by small quanta of light.  
By observing the spectra of hemin-resembling analogues of  
chlorophyll it was possible to clarify the long-wave dis-  
placement of the spectral region of the photodecomposition  
of water by basing it on the principle of Frank-Kondon.

Mechanism of Chemical Processes and the Structure of the  
Molecule.

Tsimbal, R.N., Burmistrov, S.I. and Loshkarev, M.A. (Dnepropetrovsk).

Card 12/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

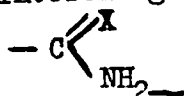
Mechanism of Chemical Processes and the Structure of the  
Molecule. (Cont.)

Reaction kinetics of nitrosation of azo-dyes with a second-  
ary amino-group.

It was found that the reaction velocity changes with the  
concentration  $\text{Cl}^-$  and  $\text{Br}^-$ . Isobaric potentials and ther-  
mal effects of the nitrosation reaction were calculated  
from the equilibrium constants and from the functions of  
temperature.

Brodskiy, A.I. and Strizhak, L.L. (Kiev).

Nitrogen-exchange of amino-groups between liquid ammonia  
and organic nitrogen-containing compounds at 100 - 200 C.  
Nitrogen does not enter into the interchange. In com-  
pounds containing amino-groups attached directly to the  
aromatic nucleus, the interchange occurs only in the  
presence of strongly negative substituents. Considerable  
interchange is revealed in the groups



, where X = O, S, N-(amides of acids, urea and

Card 13/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Mechanism of Chemical Processes and the Structure of the  
Molecule. (Cont.)

its analogues). The velocity of interchange depends on the negative substituents and on the interlinking. The lecturers explained this by the mechanism of bimolecular nucleophilic substitution.

Geller, B.A. (Kiev).

Investigations on the mechanism of some reactions with heavy nitrogen.

During the reaction between an azo-dye and diazonium hydrate diazo-compounds migrate together with their N-atoms. During the condensation of nitroso-naphthosultam with o-phenylenediamine the nitroso-group is split off in the form of ammonia.

Morozov, V.P. (Dnepropetrovsk).

Calculation of basic frequencies of oscillation of isotope molecules.

1). Method of intervals, allowing the determination of limits of possible displacement of basic frequencies of oscillation of molecules during complete isotope

Card 14/20



73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Mechanism of Chemical Processes and the Structure of the  
Molecule. (Cont.)

substitution.

- 2). Method of partial frequencies. Approximate relations were obtained which permit the definition of frequencies of basic oscillations of the molecules  $XY_2^1$  and  $XY^1$  when the frequencies of oscillation of the molecule  $XY_2$  are known.

Medvedev, K.P. (Kharkov).

Theory on the participation of single-electron bonds in the structure of molecules and in the electron bonding.

Lutskiy, A.Ye. (Kharkov).

Functional equations on the relation between the micro-properties of substances and their dependence on the microproperties of particles.

Carl 15/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Electrolysis and Electrode Processes.

Delimarskiy, Yu. K. (Kiev).

Polarographic investigations on dissolved salts. The kinetics of electrode processes were investigated. It was found that the limit currents due to the concentration polarisation are defined by the diffusion which is the slowest phase of the electrode process. The polarographic waves on solid electrodes conforms to the equation by Geirovskii - Il'kovich which defines the polarographic waves for dissolved salts.

Stender, V.V., Kalinovskiy, Ye.A. and Nikiforov, Ya.F. (Dnepropetrovsk).

Mechanism of electrolysis of zinc chloride on graphite electrodes.

Afanasyev, A.S. and Miroshnichenko, O.Ya. (Dnepropetrovsk).

Investigations on the electroreduction of oxygen and the separation of hydrogen on magnetite monocrystals.

Gritsan, D.N. and Shun, D.S. (Kharkov).

Influence of fatty alcohols on the spontaneous periodic

Card 16/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Electrolysis and Electrode Processes. (Cont.)

changes of cathodic polarisation and the current strength  
in the chain.

This is due to the periodic passivation and activation of  
the cathode surface during the electrolysis.

Galushko, V.P. (Dnepropetrovsk).

Theoretical analysis of periodic oscillations in electro-  
chemical systems and the author's experiments proving the  
phenomena.

Galushko, V.P. and Dorosh, T.P.

Mechanism of the electro-reduction of some sparingly  
soluble silver compounds.

Portnov, M.A. (Rubezhnoye).

New thymolquihydron electrodes for measuring the pH and  
its possible uses.

Pamfilov A.V. and Tsinman, A.I. (Chernovtsy)

Analysis of the relation of the effective activation  
energy of electrode processes and the polarisation  
potential.

Card 17/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Electrolysis and Electrode Processes. (Cont.)

In some cases of chemical polarisation the slow stage of  
the process can be defined.

Ksenzhek, O.S. (Dnepropetrovsk).

Calculation of the activation energy on simple electrodes  
from activation energy values on a smooth electrode and  
from the electroconductivity of the solutions.

Experiments were carried out on the activation energy of  
separation of chlorine on graphites.

Afanasyev, A.S. and Shevchenko, V.V. (Dnepropetrovsk).

Experimental data proving the theory of the mechanism of  
electro-reduction of oxygen on oxidised steel by inter-  
mediate formation of hydrogen peroxide.

Brodskiy, A.I., Fomenko, A.S., Abramova, T.M. and Gankina, I.L.  
(Kiev).

Results of the investigation on the corrosion mechanism  
of some metals.

Card 18/20 The electrochemical corrosion mechanism and magnesium was  
confirmed by using the heavy isotope of oxygen.

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Electrolysis and Electrode Processes. (Cont.)

Roykh, I.L. (Odessa).

Investigations on the formation of hydrogen peroxide by metals during atmospheric corrosion with photographic methods.

Phase Equilibria.

Garber, Yu. N. (Dnepropetrovsk).

Calculation of the mean buoyancies of vapours of complex mixtures from phase equilibrium curves.

Yermolayeva, Ye.V. (Kharkov).

The viscosity of solutions of tri-component systems  
( $MgO-Al_2O_3-SiO_2$ ,  $CaO-Al_2O_3-SiO_2$ ,  $MgO-CaO-Al_2O_3$  and

$MgO-CaO-SiO_2$ ). Their surface tension on the liquid-gas boundary, the wetting with solid solutions and electromotive forces arising between the solid phases and equilibrium solutions.

Card 19/20

73-2-21/22

Second Ukrainian Conference on Physical Chemistry  
(Symposium). (Cont.)

Electrolysis and Electrode Processes. (Cont.)

Gluzman, M.Kh. and Rubtsova, V.P. (Kharkov).

Quick definition of the melting point of eutectic binary, ternary and 4-component organic systems by contact melting.

Yasnitskiy, B.G. and Satanovskaya, Ts.I. (Kharkov).

The definition of the molecular weight, density and refractive index of chloroacetic aldehyde and its aqueous solutions in wide ranges of concentration.

Pamfilov, A.V. and Prodan, Ye.A. (Chernovtsy).

Physico-chemical investigations on tri-polyphosphates.

It was recommended to the Institute of Physical Chemistry im. L.V. Pisarzhevskiy AN USSR to hold the next Conference in 1958. The Department of Physical Chemistry of the Khar'kov Institute was advised to hold a meeting on the Theory of Solutions in 1957.

AVAILABLE: Library of Congress  
Card 20/20

SOV-21-58-2-15/28

AUTHORS: Abramova, T.M., Gankina, I.L. and Fomenko, A.S.

TITLE: Investigation of Cathode Reduction of Oxygen to Hydrogen Peroxide on a Coal-Nickel Electrode (Issledovaniye katodno-go vosstanovleniya kislороda do perekisi vodoroda na ugol'no-nikelevom elektrode)

PERIODICAL: Dopovidi Akademii nauk Ukrain's'koi RSR, 1958, Nr 9, pp 974 - 976 (USSR)

ABSTRACT: The process of cathode reduction of oxygen is used in technical production of hydrogen peroxide. However, the mechanism of the reaction which takes place in this process has not been explained thus far. The authors employed the heavy isotope  $O^{18}$  in order to clarify the origin of oxygen in hydrogen peroxide, which forms on a coal-nickel cathode in the oxygen reduction. As a result of this investigation it was shown that only molecular oxygen blown through the electrode plays a part in the cathode formation of hydrogen peroxide, but not the oxygen of water. These findings are in agreement with the concept of A.N. Frumkin that hydrogen peroxide

Card 1/2

SOV-21-58-9-15/28

Investigation of Cathode Reduction of Oxygen to Hydrogen Peroxide on a Coal-Nickel Electrode

formation is due to "newly"-adsorbed oxygen. There are 2 diagrams, 1 table and 10 references, 7 of which are Soviet, 2 English and 1 American.

ASSOCIATION: Institut fizicheskoy khimii imeni L.V. Pisarzhevskogo AN UkrSSR (Institute of Physical Chemistry im. L.V. Pisarzhevskiy of the AS UkrSSR)

PRESENTED: By Member of the AS UkrSSR, A.I. Brodskiy

SUBMITTED: April 21, 1958

NOTE: Russian title and Russian names of individuals and institutions appearing in this article have been used in the transliteration

1. Oxygen--Reduction
2. Hydrogen peroxide--Production
3. Electrolysis

Card 2/2



FOMENKO, A.S.

PHASE I BOOK EXPLOITATION SOV/2216

5(4)

Sveshchaniye po elektrokimii. 4th, Moscow, 1956.  
 Trudy...; [sbornik] [transactions of the Fourth Conference on Elect-  
 rochemistry; Collection of Articles] Moscow, Izd-vo AN SSSR,  
 1959. 868 p. Errata slip inside. 2,500 copies printed.  
 Sponsoring Agency: Akademiya nauk SSSR. Otdelatsiya khimicheskikh  
 nauk.  
 Editorial Board: A.M. Prumkin (Resp. Ed.), Academician, O.A. Yesin,  
 Professor, S.I. Zhdanov (Resp. Secretary), B.M. Kabanov, Prof.  
 Feodor S.I. Zhdanov (Resp. Secretary), B.M. Kabanov, Prof. P.D.  
 Ya. M. Kolotyrkin, Doctor of Chemical Sciences, Prof. P.D.  
 Lukovtsev, Professor, Z.A. Solov'yeva, V.V. Skudnov, Professor,  
 and G.M. Florianovich; Ed. of Publishing House M.G. Yegorov;  
 Tech. Ed.: T.A. Frusakova.

PURPOSE: This book is intended for chemical and electrical engi-  
 neers, physicists, metallurgists and researchers interested in  
 various aspects of electrochemistry.  
 COVERAGE: The book contains 127 of the 138 reports presented at  
 the Fourth Conference on Electrochemistry sponsored by the Depart-  
 ment of Chemical Sciences of the Institute of Physical Chemistry,  
 Academy of Sciences, USSR. The collection pertains to different  
 branches of electrochemical kinetics, double layer theories and  
 galvanic processes in metal electrodeposition and industrial elect-  
 rolysis. Abridged discussions are given at the end of each divi-  
 sion. The majority of reports not included here have been  
 published in periodical literature. No personalities are mentioned.  
 References are given at the end of the articles.

Kresal'shchikov, A.I. (Gosudarstvennyy institut azotnoy  
 promyshlennosti - State Institute of the Nitrogen Industry), 272  
 Electrochemical Reactions of Oxygen

Gerbovich, M.A. (Deceased), and R.I. Kaganovich (Moscow State  
 University). Study of the Mechanism of Some Anode Processes  
 by Combining Electrochemical and Tagged-Atom Methods 277

Shlygin, A.I., and G.A. Bogdanovskiy (Moscow State University),  
 Mechanism of the Electrochemical Oxidation of Some Compounds  
 on Platinum 282

Khomyakov, V.G., M.G. Bakhcharaevskiy, and A.P. Tomilov  
 (Moskovskiy khimiko-tekhnicheskii institut imeni  
 D.I. Mendeleeva-Moscow Institute of Chemical Technology  
 imeni D.I. Mendeleeva); Mechanism of the Electrolytic  
 Oxidation of Acetone in Alkaline Solutions 287

Rhomotov, M. Ye. (Moscow Institute of Chemical Technology imeni  
 D.I. Mendeleeva). Mechanism of Some Irreversible Elect-  
 rochemical Reactions 292

Card 12/34

Polytic-Oxidation Reactions 292  
 Fomenko, A.S., T.M. Abramova and I.F. Garkina (Institut  
 Khimicheskoy Khimii AN USSR-Institute of Physical Chemistry  
 imeni D.I. Mendeleeva, Moscow); Mechanism of the Co-  
 oxidation of Iron, Magnesium, Zinc and Aluminum With the Aid of Heavy Oxygen Isotopes 299  
 Discussion [A.M. Ginzberg, A.P. Tomilov, P.D. Lukovtsev,  
 G.A. Tedorsz and contributing authors] 302  
 309

PART IV. ELECTRODE PROCESSES IN FUSIONS

Yesin, O.A. (Ural'skiy politekhnicheskii institut Ural Poly-  
 technic Institute). Electrode Processes in Fu-ed Oxides 311  
 Montgalli, R., G. Starabala, M. Francini, and O. Montanelli  
 (Italy). Investigation of Overvoltage Phenomena in Fu-ed  
 Salts 323

Card 13/ 34

5 (4), 18 (7)

05824

AUTHORS: Fomenko, A. S., Abramova, T. M.,  
Gankina, I. L.

SOV/76-33-10-22/45

TITLE: An Investigation of Metal Corrosion With the Help of the Heavy Oxygen Isotope. II. Moist Atmospheric Corrosion of Cadmium

PERIODICAL: Zhurnal fizicheskoy khimii, 1959, Vol 33, Nr 10, pp 2249 - 2252 (USSR)

ABSTRACT: In previous articles (Refs 1,2) the corrosion of iron and magnesium was investigated by means of the heavy oxygen isotope, and it was shown that there was an electrochemical mechanism with oxygen depolarization. In this article the authors investigated the moist atmospheric corrosion of cadmium by the same method and made experiments on the oxygen exchange of the corrosion products of cadmium ( $\text{Cd}(\text{OH})_2$ ) with  $\text{H}_2\text{O}^{18}$ . The latter indicated that no exchange took place within 60 hours (Table 1). Since investigations with the help of cadmium filings of the sort KD-0 failed, experiments were made by means of cadmium plating applied to a quartz tube (inner side). 3% NaCl solution served as corrosion liquid which was poured into the tube in a definite quantity together with oxygen at atmospheric pressure. The

Card 1/3

An Investigation of Metal Corrosion With the Help of <sup>18</sup>O SOV/76-33-10-22/45  
the Heavy Oxygen Isotope. II. Moist Atmospheric Corrosion of Cadmium

content of <sup>18</sup>O in the resultant water and gaseous oxygen was determined by mass spectrometric analysis (Ref 3). Experiments were made with the aid of natural water in heavy oxygen atmosphere as well as with H<sub>2</sub><sup>18</sup>O in a common oxygen atmosphere. Results of measurement (Table 2) concerning the distribution of <sup>18</sup>O among water, gaseous oxygen and corrosion products indicate that cadmium corrodes according to two parallel mechanisms, i.e. an electrochemical mechanism with oxygen depolarization (as has already been observed by Feitknecht, Wyler (Ref 5), Ya. M. Kolotyркиn and L. A. Medvedeva (Ref 6)) and a chemical mechanism. Investigations of copper corrosion have shown that the exchange of Cu(OH)<sub>2</sub> oxygen with water is equilibrated after 48 hours. Accordingly, the afore-mentioned method cannot be applied here. In conclusion, the authors thank Academician A. I. Brodskiy for his help. There are 2 tables and 7 references, 6 of which are Soviet.

Card 2/3

05824

An Investigation of Metal Corrosion With the Help of the Heavy Oxygen Isotope. II. Moist Atmospheric Corrosion of Cadmium SOV/76-33-10-22/45

ASSOCIATION: Akademiya nauk USSR, Institut fizicheskoy khimii im. L. V. Pizarzhevskogo, Kiyev (Academy of Sciences of the UkrSSR, Institute of Physical Chemistry imeni L. V. Pizarzhevskiy, Kiyev)

SUBMITTED: March 21, 1958

Card 3/3

18-8300  
5.2600(A)

67265

SOV/20-129-4-29/68

~~5(A), 5(B)~~

AUTHORS: Abramova, T. M., Gankina, I. L., Pomenko, A. S.

TITLE: The Mechanism of Hydrogen Peroxide Formation in the Corrosion of Metals

PERIODICAL: Doklady Akademii nauk SSSR, 1959, Vol 129, Nr 4, pp 820-823 (USSR)

ABSTRACT: The authors set themselves the task of finding an answer to the following question: Is the hydrogen peroxide which is formed as an intermediate in the corrosion of metals caused by water and air formed from oxygen of the air, from that of water, or from the oxygen of both? As, according to reference 10, H<sub>2</sub>O<sub>2</sub> is formed as an intermediate in the cathodic reduction of O, and O is depolarized also in the corrosion of metals in air, the formation of H<sub>2</sub>O<sub>2</sub> from the O in air was probable. In that case its isotopic composition would have to correspond to that of gaseous O. This was checked by the authors by means of O<sup>18</sup> which was either added to the O blown through the solution corroding the metal, or was admixed to the solution as H<sub>2</sub>O<sub>2</sub><sup>18</sup>. The

Card 1/4

67265

SOV/20-129-4-29/68

## The Mechanism of Hydrogen Peroxide Formation in the Corrosion of Metals

approach of the isotopic composition of the  $H_2O_2$  formed to that of gaseous  $O$  was actually observed by the authors in the corrosion of Zn, Mg, Sn, Al, and Cd in aqueous solution of  $H_2O_2$  and during the blowing through of  $O$ . The experiments were made in the dark at room temperature and took 1 - 24 hours. The  $H_2O_2$  content of the solution was then determined by means of permanganate (in the case of Zn and Cd the insoluble peroxides were dissolved by acidification) and the isotopic composition of  $H_2O_2$  was determined by means of a mass spectrometer. Table 1 shows that in the experiments with  $H_2O_2^{18} + O_2^{16}$  the  $O^{18}$  content decreases in the  $H_2O_2$  analyzed after the experiment is ended, whereas it increases in the experiments made with  $H_2O_2^{16} + O_2^{18}$ . The  $H_2O_2$  is thus produced from air-oxygen. A rough calculation of the isotopic composition of  $H_2O_2$  to be expected, carried out, as an example, on Zn, showed a difference of 13% if compared

Card 2/4

67265

SOV/20-129-4-29/68

The Mechanism of Hydrogen Peroxide Formation in the Corrosion of Metals

with the experimental result. Also the differences found in experiments with other metals are of the same order of magnitude. The following causes are assumed to be responsible:

1) Part of the  $H_2O_2$  formed is immediately again catalytically decomposed by the metal. 2) The O liberated in this decomposition partly again enters into reaction accompanied by the formation of  $H_2O_2$ . Attempts at stabilizing the  $H_2O_2$  formed by additions of oxyquinoline, sodium pyrophosphate, sodium silicate etc. were unsuccessful. The authors refer to published data, according to which there is no exchange between the oxygen of  $H_2O_2$  and of air under the prevailing experimental conditions (Refs 14,15), which they were able to confirm by control tests. Thus, as no side-reactions occur, the results obtained by the authors prove that the  $H_2O_2$  is produced in the corrosion of metals from the oxygen in the air. It is finally mentioned that the authors thank A. I. Brodskiy, Academician of the AS UkrSSR, for supervising the investigations, and Engineer I. M. Protas for the mass-spectrometrical analyses

Card 3/4

67265

SOV/20-129-4-29/68

The Mechanism of Hydrogen Peroxide Formation in the Corrosion of Metals

carried out. There are 1 table and 16 references, 3 of which are Soviet.

ASSOCIATION: Institut fizicheskoy khimii im. L. V. Pisarzhevskogo Akademii nauk USSR (Institute of Physical Chemistry imeni L. V. Pisarzhevskiy of the Academy of Sciences, UkrSSR)

PRESENTED: July 13, 1959, by A. N. Frumkin, Academician

SUBMITTED: July 13, 1959

4

Card 4/4



FOMENKO, A.S.

~~LATYSHEV, G.D.~~

176

PHASE I BOOK EXPLOITATION SOV/5410

Tashkentskaya konferentsiya po mirnomu ispol'zovaniyu atomnoy energii, Tashkent, 1959.

Trudy (Transactions of the Tashkent Conference on the Peaceful Uses of Atomic Energy) v. 2. Tashkent, Izd-vo AN UzSSR, 1960. 449 p. Errata slip inserted. 1,500 copies printed.

Sponsoring Agency: Akademiya nauk Uzbekskoy SSR.

Responsible Ed.: S. V. Starodubtsev, Academician, Academy of Sciences Uzbek SSR. Editorial Board: A. A. Abdullayev, Candidate of Physics and Mathematics; D. M. Abdurasulov, Doctor of Medical Sciences; U. A. Arifov, Academician, Academy of Sciences Uzbek SSR; A. A. Borodulina, Candidate of Biological Sciences; V. N. Ivashev; G. S. Ikramova; A. Ye. Kiv; Ye. M. Lobanov, Candidate of Physics and Mathematics; A. I. Nikolayev, Candidate of Medical Sciences; D. Nishanov, Candidate of Chemical Sciences; A. S. Sadykov, Corresponding Member, Academy of Sciences USSR, Academician, Academy of Sciences Uzbek SSR; Yu. N. Talanin,

~~Car-1/20~~

176

Transactions of the Tashkent (Cont.)

SOV/5410

Candidate of Physics and Mathematics; Ya. Kh. Turakulov, Doctor of Biological Sciences. Ed.: R. I. Khamidov; Tech. Ed.: A. G. Babakhanova.

**PURPOSE :** The publication is intended for scientific workers and specialists employed in enterprises where radioactive isotopes and nuclear radiation are used for research in chemical, geological, and technological fields.

**COVERAGE:** This collection of 133 articles represents the second volume of the Transactions of the Tashkent Conference on the Peaceful Uses of Atomic Energy. The individual articles deal with a wide range of problems in the field of nuclear radiation, including: production and chemical analysis of radioactive isotopes; investigation of the kinetics of chemical reactions by means of isotopes; application of spectral analysis for the manufacturing of radioactive preparations; radioactive methods for determining the content of elements in the rocks; and an analysis of methods for obtaining pure substances. Certain

Card 2/20

176

Transactions of the Tashkent (Cont.)

SOV/5410

instruments used, such as automatic regulators, flowmeters, level gauges, and high-sensitivity gamma-relays, are described. No personalities are mentioned. References follow individual articles.

TABLE OF CONTENTS:

RADIOACTIVE ISOTOPES AND NUCLEAR RADIATION  
IN ENGINEERING AND GEOLOGY

Lobanov, Yp. M. [Institut yadernoy fiziki UzSSR - Institute of Nuclear Physics AS UzSSR]. Application of Radioactive Isotopes and Nuclear Radiation in Uzbekistan

7

Taksar, I. M., and V. A. Yanushkovskiy [Institut fiziki AN Latv SSR - Institute of Physics AS Latvian SSR]. Problems of the Typification of Automatic-Control Apparatus Based on the Use of Radioactive Isotopes

9

Card 3/20

14

- Brodskiy, A. I., I. P. Gragerov, I. F. Franchuk, L. V. Sulina, I. I. Kukhtenko, V. A. Lunenok, A. S. Fomenko, and A. M. Aleksankin [Institut fizicheskoy khimii AN SSSR - Institut of Physical Chemistry AS USSR]. Investigation of the Mechanism of Oxidizing Reactions by the Isotopic Method 327
- Savrukina, A. K. [Institut geokhimi i analiticheskoy khimii im. V. I. Vernadskogo AN SSSR - Institute of Geochemistry and Analytical Chemistry imeni V. I. Vernadskiy AS USSR]. Methods of Modern Radiochemistry and the Fields of Its Application 334
- Chuveleva, E. A., K. V. Chmutov, and P. P. Nazarov. [Institute of Physical Chemistry AS USSR]. Study of the Adsorption of Alkaline and Rare-Earth Elements on Black Earth by the Tracer Atom Method 341
- Novikov, A. I. [Taizhikskiy gosudarstvennyy universitet im. V. I. Lenina-Tadzhik State University imeni V. I. Lenin]. Coprecipitation of Small Quantities of Various Cations and Anions With Metal Hydroxides 349
- Ampelogova, N. I. [Radiyevyy institut im. V. G. Khlopina
- Gard 16/20

FOMENKO, A.S.

S/CS:62/000/001/004/067  
B156/B101

AUTHORS: Brodskiy, A. I., Gragerov, I. P., Franchuk, I. F., Sulima, L.V.,  
Kukhtenko, I. I., Lunenok, V. A., Fomenko, A. S.,  
Aleksankin, M. M.

TITLE: Mechanism of oxidation reactions investigated by the isotopic  
method

PERIODICAL: Referativnyy zhurnal. Khimiya, no. 1, 1962, 60, abstract  
18439 (Tr. Tashkentok. konferentsii po mirn. ispol'zovaniyu  
atomn. energii, v. 2. Tashkent, AN UzSSR, 1960, 327-334)

TEXT: A review of work done by the authors on studying the mechanism of  
certain oxidation reactions using isotopes: the oxidation of organic  
compounds with chromyl chloride, the mechanism of anthranil regrouping, the  
process of oxidation of aniline, o-anisidine and p-nitroaniline with Caro  
acid. The mechanism whereby hydrogen peroxide and certain persulfate-type  
inorganic peroxide compounds are formed and converted is examined; so also  
are the kinetics of isotopic exchange in substituted benzoic acids.

Card 1/2

Mechanism of oxidation reactions ...

3  
S/081/62/000/001/004/067  
B156/B101

benzaldehydes, alcohols, naphthalenes and nitro compounds with  $H_2O^{18}$ .  
18 references. [Abstracter's note: Complete translation.]

Card 2/2

33490

S/195/61/002/005/015/027

E111/E485

11.1310

AUTHORS: Fomenko, A.S., Gankina, I.L., Avramova, T.M.

TITLE: Study of the mechanism of the decomposition of hydrogen peroxide on activated charcoal by the isotope method

PERIODICAL: Kinetika i kataliz, v.2, no.5, 1961, 732-736

TEXT: The decomposition of hydrogen peroxide on activated charcoal was studied by many authors but their assumptions on the mechanism of this process are in contradiction. In the present work the mechanism was studied using  $O^{18}$  introduced into the charcoal oxides (with  $H_2O_2$  of the natural isotope composition) or into the  $H_2O_2$  (with ordinary oxygen charcoal). The isotope compositions of the charcoal oxides and of gaseous oxygen were determined to evaluate the mechanism. The  $O^{18}$ -containing charcoal was prepared by grinding the commercial charcoal type "BAU" (BAU), de-ashing with acid, washing, drying and activating in  $CO_2$  at  $800^\circ C$  for 6 hours and  $1000^\circ C$  for 3 hours; after this, the material was treated with hydrogen at  $1000^\circ C$  (which was then pumped off at 600 to  $650^\circ C$ ), cooled in oxygen-free nitrogen, treated at room temperature with  $O^{18}$ -enriched gaseous oxygen and  
Card 1/4

33490

S/195/61/002/005/015/027  
E111/E485

Study of the mechanism ...

stored in heavy-oxygen water vapour. The procedure for preparing and storing ordinary oxygen-charcoal was identical but ordinary oxygen and water vapour were used. Special experiments were carried out in which charcoal containing ordinary oxygen was treated with heavy-oxygen water vapour whose final isotope content was then determined, or charcoal containing  $O^{18}$  was treated with ordinary water, filtered off and its isotope composition determined by mass spectrometry on the  $CO_2$  obtained by degassing at 300 to 550°C. For  $H_2O_2$ , mass spectrometric isotope analysis was carried out on the oxygen evolved when the peroxide was decomposed with potassium permanganate: and for water, on  $CO_2$  after exchange with the water. Results showed that there is no exchange of the oxygen of the basic oxides with water. For the main  $H_2O_2$  decomposition experiments in a quartz tube containing 1 to 4 g of degassed charcoal was put in a furnace. One end of the tube was connected to a trap for freezing out the desorbed gases and to a vacuum installation; the other to a trap containing the required quantity of peroxide, freed from dissolved oxygen. The tube was periodically rotated. After the required time, the

Card 2/4

X



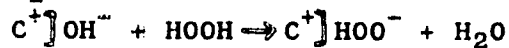
33490

S/195/61/002/005/015/027

E111/E485

Study of the mechanism ...

water and undecomposed peroxide was distilled and the charcoal degassed as before. The CO<sub>2</sub> fraction, collected at 300 to 550°C, was analysed for O<sup>18</sup>. In some experiments the isotope composition of the oxides was determined from the water obtained during their treatment with hydrogen. Mass spectrometry revealed carbon dioxide as well as oxygen in the gaseous decomposition products. In agreement with the views of G. Brinkmann (Ref. 6: Ang. Chem., v. 61, 1949, 378) the results suggest that a double decomposition type of reaction occurs between the basic OH groups in the charcoal surface and HO<sub>2</sub><sup>-</sup> ions of the peroxide



followed by  $C^+ \text{HOO}^- + \text{HOOH} \rightarrow C^+ \text{OH}^- + \text{H}_2\text{O} + \text{O}_2$ .

The origin of CO<sub>2</sub> in the decomposition of H<sub>2</sub>O<sub>2</sub> on charcoal is not clear since the simultaneous presence of H<sub>2</sub>O and CO<sub>2</sub>, in view of the exchange between them, alters the primary CO<sub>2</sub> composition. The observed change in the isotope composition of oxides on charcoal cannot be attributed to further oxidation of the coal to give oxygen compounds, differing from OH groups, whose high-Card 3/4

33490  
S/195/61/002/005/015/027  
E111/E485

Study of the mechanism ...

temperature decomposition could give a further quantity of CO<sub>2</sub> with the peroxide isotope composition. No free radicals on the charcoal surface were found by the electron paramagnetic resonance method and this is contrary to the chain mechanism proposed by various authors (e.g. Ref.8: V.A.Garten, E.Epinger, D.E.Weiss, Austr. J. Appl. Sci., v.7, 1956, 149). Academician (AS UkrSSR) A.I.Brodskiy helped in the work. There are 2 tables and 12 references: 2 Soviet-bloc and 10 non-Soviet-bloc. The four most recent references to English language publications read as follows: Ref.4: A.King, J. Chem. Soc., 1936, 1688; Ref.5: E.C.Larsen, J.H.Walton, J. Phys. Chem., v.44, 1940, 70; Ref.10: B.R.Puri, L.A.Scharma, D.D.Singh, Ind. Eng. Chem., v.50, 1958, 1075; J. Ind. Chem. Soc., v.35, 1958, 765; Ref.12: M.Cohn, H.C.Urey, J. Amer. Chem. Soc., v.60, 1958, 679.

ASSOCIATION: Institut fizicheskoy khimii im. L.V.Pisarzhevskogo  
AN UkrSSR Kiyev (Institute of Physical Chemistry  
im. L.V.Pisarzhevskiy AS UkrSSR, Kiyev)

Card 4/4

FOMENKO, A.S.; ABRAMOVA, T.M.; GANKINA, I.L.

Decomposition of hydrogen peroxide in the presence of  
potassium iodate, bromate, and chlorate. Ukr. khim. zhur.  
28 no.1:14-17 '62. (IRA 16:8)

1. Institut fizicheskoy khimii im. L.V. Pisarzhevskogo  
AN UkrSSR.

... ..  
... ..  
... ..  
... ..

... ..  
... ..

... ..

... ..  
... ..  
... ..  
... ..

... ..  
... ..

... ..  
... ..  
... ..  
... ..  
... ..  
... ..

1967-07

ATSO02665

... latter process by di-β-naphthyl-β-phenylacetamide... detected H<sub>2</sub>O and 3% CO<sub>2</sub> in the gaseous prod...

... participation in evaluating the results. ...

ASSOCIATION: Institut fizicheskoy khimii im. L. V. Plazhkevskogo AN UkrSSR (Institute of Physical Chemistry, AN UkrSSR); Kiyevskiy filial Vsesoyuznogo nauchno-issledovatel'skogo instituta iskusstvennogo volokna (Kiev Branch, All-Union Research Institute of Synthetic Fibers)

RECEIVED: 22 Jun 64

ENCL 00 SUB CODE MT, NP

NO REF SOV: 002  
Card 7/2 LL

OTHER: 003

ACCESSION NR: AP4033700

8/0073/64/030/004/0376/0384

AUTHOR: Fomenko, A. S.; Abramova, T. M.; Dar'yeva, E. P.; Galina, A. A.; Furman, Ye. G.

TITLE: Oxidative destruction of polyamides. II. Participation of free radicals in the radiolysis and radiation oxidation of polycaprolactam.

SOURCE: Ukrainskiy khimicheskij zhurnal, v. 30, no. 4, 1964, 376-384

TOPIC TAGS: polyamide, polycaprolactam, caprolactam oligomer, oxidation, free radical formation, radiolysis, radiation oxidation, EPR spectra, C N bond rupture, hydroperoxide formation, IR spectra, antioxidant, viscosity, cross linkage

ABSTRACT: The free radicals formed by irradiation of polycaprolactam with cobalt-60, their function in the radiation oxidation of polycaprolactam, and the inhibiting action of an antioxidant were investigated. The electron paramagnetic resonance spectra of polycaprolactam and caprolactam oligomers irradiated with cobalt-60, and the effects of temperature, radiation dose and presence of oxygen on the changes in these spectra are described. The gaseous products of polycaprolactam radiolysis in vacuum are hydrogen and carbon monoxide in a 3:1 ratio and about

Card

1/3

ACCESSION NR: AP4033700

3% CO<sub>2</sub>. The amount of terminal amino groups almost doubled on irradiation; with a 22 mrad dose this corresponded to the rupture of 1% of the C-N bonds in the polymer. The viscosity of the polymer also changes on irradiation--with 8 mrad irradiation the viscosity decreased during the first 30 hours, then increased, apparently due to the formation of cross-linked structures. The accumulation of hydroperoxide in polycaprolactam on gamma-irradiation in oxygen, the effect of radiation dose, the changes in terminal amino and carboxyl groups and the viscosity of the polymer were examined. H<sub>2</sub>:CO ratio in these products was 2:1; terminal NH<sub>2</sub> and COOH groups increased at doses below 15 mrad and decreased above that. These data agree with changes in the IR spectra of the irradiated polycaprolactam. It is concluded that the RO<sub>2</sub> radical formed by radiation oxidation is converted to the hydroperoxide and carbonyl-containing compounds by a parallel route. Addition of 0.5-3% antioxidant di-β-naphthyl-p-phenylenediamine to the polymer does not affect the form of the EPR spectra or concentration of free radicals formed by gamma-irradiation; but this additive significantly lowers the amount of hydroperoxide and carbonyl-containing compounds formed by radiation oxidation. "N. S. Oleynik and M. T. Kozhura took part in the experimental work."... "The authors thank AN USSR academician A. I. Brodsko for help in the work and participation in its evaluation, and also

Card 2/3

ACCESSION NR: AP4033700

coworkers in the electron paramagnetic resonance laboratory for obtaining EPR spectra and help in evaluating the spectral data." Orig. art. has: 6 figures and 1 table.

ASSOCIATION: Institut fizicheskoy khimii im. L. V. Pisarzhevskogo AN USSR, (Institute of Physical Chemistry); Kiyevskiy filial Vsesoyuznogo NII iskusstvennogo volokna, (Kiev Branch of the All Union NII of Synthetic Fibers)

SUBMITTED: 15 May 63

ENCL: 00

SUB CODE: OC, NP.

NO REF SOV: 010

OTHER: 007

Card 3/3



ACCESSION NR: AP4040955

#/0020/64/156/005/1147/1149

‡ Corresponding member AN SSSR);  
AUTHOR: Brodskiy, A. I; Pomenko, A. S.; Abramova, T. M.; Furman, Ye. G.  
Dar'yeva, E. P.; Kukhtenko, I. I.; Galina, A. A.

TITLE: EPR spectra of radicals formed during gamma irradiation of polyamides

SOURCE: AN SSSR. Doklady\*, v. 156, no. 5, 1964, 1147-1149

TOPIC TAGS: electron paramagnetic resonance, EPR spectra,  
EPR radical spectra, polyamide, polyamide gamma  
irradiation, hexamethylene adipamide, poly-omega-undecane amide,  
deuterium, caproamide

ABSTRACT: The authors conducted this analysis because the literary data pertaining to the structure of radicals formed under the effects of irradiation are contradictory. The EPR spectra of poly-ε-caproamide were recorded. The irradiation and EPR spectra recording was taken at room temperature. The EPR spectrum of the gamma-irradiated poly-ε-caproamide is an incompletely resolved quintet 1 : 2 : 2 : 2 : 1 with an average width of 74 oersteds between the extreme maxima. The cleavage between the extreme pairs of lines 1-2 and 4-5 is 21 oersteds. This is 1.55 times less than the cleavage between the lines 2-4. This spectrum corresponds to a  $\text{-OH-CO-NH-OH-OH}$  radical in which the unpaired

card 1/3                      2                      2

ACCESSION NR: AP4040955

electron interacts with one  $\alpha$ -hydrogen and two equivalent  $\beta$ -hydrogens. The  $-\text{CO}-\text{CD}_2-\text{CH}_2-\text{CD}_2-\text{NH}-$  sample with deuterium in the two  $\text{CH}_2$  groups neighboring the carbonyl and NH groups yields a fully resolved 1 : 2 : 1 triplet with a splitting of  $a_\beta = 28$  oersteds, and with a general width of 56 oersteds between the extreme maxima. This spectrum corresponds to a  $-\text{CD}_2-\text{CO}-\text{NH}-\text{CD}-\text{CH}_2-$  radical. The spectra of irradiated polyamides containing 8 and 10  $\text{CH}_2$  groups in the monomer unit show incompletely split 1 : 3 : 3 : 1 quadruplets with identical 21 oersted cleavages. The spectrum for an irradiated completely-crystalline hexamethylene adipamide  $\text{COOH}-(\text{CH}_2)_4-\text{CO}-\text{NH}-(\text{CH}_2)_6-\text{NH}_2$  is a satisfactorily resolved 1 : 2 : 2 : 2 : 1 quintet with a general width of 84 oersteds between the extreme maxima and with  $a_\beta = 21$  oersteds and a  $\sqrt{a_\beta} = 2.0$ . It corresponds to a radical in which the hydrogen splits off from the  $\text{CH}_2$  group in the  $\alpha$ -position to the NH, just as in the poly- $\epsilon$ -caproamide radical. The irradiated  $\epsilon$ -caprolactam monomer produces a poorly resolved spectrum. When deuterium is introduced into the methylene groups of the nondeuterated and deuterated caprolactam in the NH group a sharp change in the spectrum shape can be observed. The spectrum of the  $\text{CO}-\text{CD}_2(\text{CH}_2)_3\text{CD}_2\text{ND}$  sample is not as well resolved probably on account of the participation of the NH group hydrogen in the cleavage. This spectrum can evidently also be examined as a quadruplet with intensity ratio of 1 : 1 : 1 : 1. Orig. art. has 3 figures.

Card 2/3

ACCESSION NR: AP2040955

ASSOCIATION: Institut fizicheskoy khimii im. L. V. Posarshevskogo Akademii nauk  
UkrSSR (Institute of Physical Chemistry, Academy of Sciences Ukr SSR)

SUBMITTED: 09Mar64

ENCL: 00

SUB CODE: NP, OC

NO REF SOV: 005

OTHER: 008

Card 3/3

[The following text is extremely faint and illegible, appearing as a large block of noise or a scan artifact.]

SECRET

The IP address of a PC is (see file 2 on the same)

SECRET

~~Union Scientific Research Institute of Synthetic Fibers~~

2014/05/11

ENCLOSURE 05

NO. 00000000

GELLER, B.A.; NEYMARK, I.Ye.; RUBANIK, M.Ye.; GRAGEROV, I.P.; POLYAKOV,  
M.V.; RUSOV, M.T.; DAIN, B.Ya.; REKASHEVA, A.F.; STRAZHESKO,  
D.N.; LUNENOK, V.A.; ROYTER, V.A.; SULIMA, L.V.; FOMENKO, A.S.

Aleksandr Il'ich Brodskii, 1895- ; on his seventieth birthday.  
Zhur. fiz. khim. 39 no.6:1540-1541 Je '65.

(MIRA 18:11)

L 24491-66 EPF(n)-2/EWT(m)/EWP(j)/T/EWA(h)/EWA(l) GG/RM/WW/JW

ACC NR: AP6006980

(A)

SOURCE CODE: UR/0190/66/008/002/0261/0266

AUTHORS: Fomenko, A. S.; Abramova, T. M.; Dar'yeva, E. P.; Galina, A. A.; Furman, Ye. G.

ORG: Institute of Physical Chemistry im. L. V. Pisarzhevskiy (Institut fizicheskoy khimii)

TITLE: Mechanism of action of di-β-naphthyl-p-phenylenediamine during radiation oxidation of polycaproamide

SOURCE: Vysokomolekulyarnyye soyedineniya, v. 8, no. 2, 1966, 261-266

TOPIC TAGS: polyamide, free radical, oxidation kinetics

ABSTRACT: The effect of di-β-naphthyl-p-phenylenediamine (I) upon the kinetics of accumulation of free radicals formed during the process of radiation-induced decomposition of polycaproamide (II) was investigated, and the yield of gaseous and oxygen-containing products of radiation-induced oxidation of the polymer was determined. The changes of the content of terminal NH<sub>2</sub> groups, viscosity, and IR spectra occurring in II stabilized with I during the radiation-induced oxidation were also studied. The methods, involving ESR, chromatographic, chemical, and IR

Card 1/2

UDC: 678.01:54+678.675

19  
52  
B



L 24491-66  
ACC NR: AP6006980

0  
spectroscopic studies, were previously described by A. S. Fomenko, T. M. Abramova, E. P. Dar"yeva, A. A. Galina, and Ye. G. Furman (Ukr. khimich. zh., 30, 376, 1964). It was established that I has no effect upon breaking of the C-H, C-N, and -C-CO-bonds during radiolysis of II, but does affect C-N and C-CO bonds during radiation oxidation of II. The amount of peroxy carbonyl and carboxyl compounds formed during radiation oxidation of II stabilized with I is considerably lowered as compared with the untreated II. A possible mechanism for the inhibiting action of I is offered. Orig. art. has: 1 table, 6 figures, and 4 equations.

SUB CODE: 07, 11/ SUBM DATE: 05Mar65/ ORIG REF: 007

Card 2/2 98

L 35343-66 EWT(m)/EWF(1)/T IJF(o) JWD/GG/RM

ACC NR: AP6012725

(A)

SOURCE CODE: UR/0190/66/008/004/0770/0770

AUTHOR: Fomenko, A. S.; Krasnov, Ye. P.; Abramova, T. M.; Dar'yeva, E. P.; Furman, Ye. G.; Galina, A. A.

ORG: none

25  
B

TITLE: Radiation resistance of isomeric aromatic polyamides

19

SOURCE: Vysokomolekulyarnyye soyedineniya, v. 8, no. 4, 1966, 770

TOPIC TAGS: radiation stability, aromatic polyamide, aliphatic polyamide, gamma irradiation, radiation resistance

ABSTRACT: The integral dose required for the accumulation of  $1.10^{14}$  radicals in  $\gamma$ -irradiation of aromatic polyamides is shown to be one order higher than for aliphatic polyamides. The radiation yields of hydrogen during polymer irradiation are two orders lower than for aliphatic polyamides. There were no changes in IR-spectra and thermomechanical properties of samples  $\gamma$ -irradiated in vacuo and in the presence of oxygen. This proves the high radiation stability of aromatic polyamides.  
[Based on author's abstract.]

19

[AM]

SUB CODE: 20, 11/ SUBM DATE: 22Nov65/ ORIG REF: 002

Card 1/1

ML

UDC: 678.01:54+678.675

L 40099-06 EWT(m)/EWP(j)/T IJP(c) GG/RM

ACC NR: AP6019661

(A)

SOURCE CODE: UR/0073/66/032/006/0549/0554

AUTHOR: Brodskiy, A. I.; Fomenko, A. S.; Dar'yeva, E. P.; Abramova, T. M.; Galina, A. A.; Furman, Ye. G.

ORG: Institute of Physical Chemistry im. L. V. Piszhevskiy, AN UkrSSR (Institut fizicheskoy khimii AN UkrSSR)

TITLE: Gas evolution during the radiative-oxidative degradation of poly- $\epsilon$ -caproamide

SOURCE: Ukrainskiy khimicheskiy zhurnal, v. 32, no. 6, 1966, 549-554

TOPIC TAGS: polyamide, oxidative degradation, hydrogen, carbon monoxide, gamma radiation, radiation effect

ABSTRACT: Chromatographic analysis was used to find the radiation yields of hydrogen and carbon monoxide, the main gaseous products of the radiolysis and radiative oxidation of poly- $\epsilon$ -caproamide.  $G_{H_2}$  is about 1 mole/100 eV for both processes, and does not change as the dose rate increases from 0.4 to  $5.0 \times 10^{18}$  eV/g min.  $G_{CO}$  is equal to 0.3 mole/100 eV for radiolysis and to 0.6 mole/100 eV for radiative oxidation, and rises to 0.9 mole/100 eV as the dose rate increases from 0.4 to  $5.0 \times 10^{18}$  eV/g min. It was found that the combined action of gamma radiation and increased temperature approximately doubles the values of  $G_{H_2}$  and  $G_{CO}$  in both the radiolysis and radiative oxidation of poly- $\epsilon$ -caproamide in the case of a low dose rate of gamma radiation, and that the effect of this combined action on  $G_{H_2}$  and  $G_{CO}$  diminishes with increasing

Card 1/2

UDC: 678.01:54+678.675

L 40099-66

ACC NR: AP6019661

dose rate. It is shown that the stabilization of poly- $\epsilon$ -caproamide by the addition of the antioxidant di- $\beta$ -naphthyl-p-phenylenediamine does not change  $G_{H_2}$  during radiolysis and radiative oxidation, but markedly reduces the amount of carbon monoxide formed during radiative oxidation. Orig. art. has: 6 figures and 3 tables.

SUB CODE: 07/ SUBM DATE: 31Jan64/ ORIG REF: 006

Card

2/2

*MB*