

TOLSTOPYATOVA, A.A.; YUY TSI-TSYUAN' [Yu Ch'i-ch'uan]; DULITSKAYA, K.A.

Catalytic properties of neodymium oxide in the reactions of
dehydrogenation of tetralin. Izv. AN SSSR. Ser. khim. no.12:
2095-2100 D '63. (MIRA 17:1)

1. Institut organicheskoy khimii im. N.D. Zelinskogo AN
SSSR.

KONENKO, I.R.; TOLSTOPYATOVA, A.A.; BALANDIN, A.A.

Scandium oxide as a catalyst of dehydrogenation and dehydration.
Izv. AN SSSR. Ser. khim. no.11:1899-1905 N '63. (MIRA 17:1)

1. Institut organicheskoy khimii imeni N.D. Zelinskogo AN SSSR.

BALANDIN, A. A.; TOLSTOPYATOVA, A. A.

"Selectivity of catalysis and bond energies."

report submitted to 3rd Intl Cong on Catalysis, Amaterdam, 20-25 Jul 64.

Inst or Organic Chemistry im Zelinskiy, AS USSR, Moscow.

TOLSTOPYATOVA, A.A.; BALANDIN, A.A.; YUY TSI-TSYUAN' [Yü Ch'i-ch'üan]

Kinetics of dehydrogenation and dehydration of isopropyl alcohol and of the dehydrogenation of tetralin on lanthanum oxide. Zhur.fiz.khim. 37 no.10:2220-2227 0 '63. (MIRA 17:2)

1. Institut organicheskoy khimii AN SSSR.

BALANDIN, A.A.; KONENKO, I.R.; TOLSTOPYATOVA, A.A.

Kinetics of dehydrogenation and dehydration of ethyl and isopropyl alcohols on yttrium oxide. Kin.i kat. 2 no.6:900-906 N-D '61.
(MIRA 14:12)

1. Institut organicheskoy khimii imeni N.D. Zelinskogo AN SSSR.
(Ethyl alcohol) (Isopropyl alcohol)
(Dehydrogenation) (Dehydration (Chemistry))

BALANDIN, A.A., akademik; TOLSTOPYATOVA, A.A.; NAUMOV, V.A.

Bond energies of interaction between organogens and the surface
of oxide catalysts. Dokl. AN SSSR 148 no. 4: 825-828 F '63.

(MIRA 16:4)

1. Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosova.
(Chemical bonds) (Chemical elements) (Catalysts)

FERAPONTOV, V.A.; BALANDIN, A.A.; TOLSTOPYATOVA, A.A.

Catalytic dehydrogenation of ethylbenzene to styrene on
cadmium oxide in the presence of water vapors. Izv. AN SSSR. Otd.
khim. nauk no. 3:414-423 Mr '63. (MIRA 16:4)

1. Institut organicheskoy khimii im. N.D. Zelinskogo AN SSSR.
(Benzene) (Styrene)

TOLSTOPYATOVA, A.A.; BALANDIN, A.A.; NAUMOV, V.A.

Kinetic method used in the determination of bond energies of the reacting atoms of organic molecules having a blue molybdenum oxide surface. *Izv. AN SSSR. Otd. khim. nauk* no. 3:423-429 Apr '63.
(MIRA 16:4)

1. Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosova.
(Chemical bonds) (Molybdenum oxides) (Organic compounds)

TOLSTOPYATOVA, A.A.; BALANDIN, A.A.; PYN BI-SYAN [P'ing Pi-hsiang]

Kinetics of the dehydrogenation and dehydration of isopropyl alcohol
and of the dehydrogenation of tetralin on thulium oxide. *Izv.AN SSSR.*
Otd.khim.nauk no.9:1524-1533 S '62. (MIRA 15:10)

1. Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova.
(Isopropyl alcohol) (Dehydrogenation) (Dehydration (Chemistry))
(Naphthalene)

BALANDIN, A.A.; TOLSTOPYATOVA, A.A.; PYN BI-SYAN [P'ong Pi-hsiang]

Catalytic properties of dysprosium oxide with respect to the reaction of dehydrogenation and dehydration of alcohols and dehydrogenation of tetralin. Izv.AN SSSR.Otd.khim.nauk no.6: 974-980 '62. (MIRA 15:8)

1. Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova. (Dysprosium oxide) (Catalysis) (Dehydrogenation)

TOLSTOPYATOVA, A.A.; BALANDIN, A.A. PYN BI-SYAN [P'êng Pi-hsiang]

Kinetics of dehydrogenation and dehydration of isopropyl alcohol
and dehydrogenation of tetralin of dysprosium oxide. Izv.AN SSSR.
Otd.khim.nauk no.7:1154-1163 J1 '62. (MIRA 15:7)

1. Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova.
(Chemical reaction, Rate of) (Catalysis) (Dysprosium oxide)

TOLSTOPYATOVA, A.A.; PYN BU-SYAN [P'ing Pi-hsiang]; BALANDIN, A.A.

Kinetics of dehydrogenation and dehydration of isopropyl alcohol and of dehydrogenation of tetralin on ytterbium oxide. Izv.AN SSSR.Otd.khim.nauk no.8:1322-1329 Ag '62. (MIRA 15:8)

1. Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova.
(Isopropyl alcohol) (Naphthalene) (Dehydrogenation)

BALANDIN, A.A.; TOLSTOPYATOVA, A.A.; NAUMOV, V.A.

Determination of bond energies of the reacting atoms of organic molecules with the MoO₂ catalyst surface using a kinetic method.
Izv.AN SSSR.Otd.khim.nauk no.7:1150-1154 JI '62. (MIRA 15:7)

1. Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova.
(Chemical bonds) (Molybdenum oxide) (Chemical reaction, Rate of)

BALANDIN, A.A.; FYN BI-SYAN [P'ing Pi-hsiang]; TOLSTOPYATOVA, A.A.

Kinetics of dehydrogenation and dehydration of isopropyl alcohol
and of dehydrogenation of tetralin on gadolinium oxide. Izv.AN
SSSR.Otd.khim.nauk no.8:1330-1336 Ag '62. (MIRA 15:8)

1. Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova.
(Isopropyl alcohol) (Naphthalene) (Dehydrogenation)

TOLSTOPYATOVA, A.A.; BALANDIN, A.A.; MATYUSHENKO, V.Kh.

Determination of bond energies of atoms of organic molecules
reacting with the surface of the MnO catalyst. Izv.AN SSSR Otd.
khim.nauk no.8:1333-1336 Ag '60. (MIRA 15:5)

1. Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova i
Institut organicheskoy khimii im. N.D.Zelinskogo AN SSSR.
(Chemical bonds) (Chemical reaction, Rate of)

S/062/62/000/008/002/016
B101/B180

AUTHORS: Tolstopyatova, A. A., Ping Pi-hsiang, and Balandin, A. A.

TITLE: Kinetics of the dehydrogenation and dehydration of isopropanol and the dehydration of tetralin on ytterbium oxide

PERIODICAL: Akademiya nauk SSSR. Izvestiya. Otdeleniye khimicheskikh nauk, no. 8, 1962, 1322-1329

TEXT: Data was obtained on the catalytic dehydration of i-propanol and the dehydrogenation of i-propanol and tetralin on Yb_2O_3 , using methods described earlier (Dokl. AN SSSR, 138, 1365 (1961); Izv. AN SSSR, Otd. khim. n., 1962, 974, 1154). The low specific surface area of the catalyst ($9.1 \text{ m}^2/\text{g}$) precluded capillary condensation and complications due to diffusion. When the catalyst was mixed with quartz powder in a 1:1 ratio, the reaction took place isothermally. α , the relative adsorption coefficient of the reaction products (H_2O , C_3H_8 , acetone, H_2 , naphthalene), and the thermodynamic functions ΔH (kcal/mole), ΔF (kcal/mole), ΔS

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S/062/62/000/008/002/016
B101/B180

Kinetics of the dehydrogenation ...

(kcal/mole·deg) were calculated:

dehydration of i-propanol

°C	z_{H_2O}	ΔH	ΔF	ΔS
325	7.23	-18.7	-2.39	-10.4
365	3.48	"	-1.57	-10.3

dehydrogenation of i-propanol

°C	z_{acet}	ΔH	ΔF	ΔS
345	2.56	-21.9	-1.13	-33.5
365	1.54	"	-0.54	"

$z_{C_3H_6}$

329	1.80	-23.8	-0.69	-38.3
360	0.61	"	0.40	"

z_{H_2}

325	0.15	34.7	2.24	54.4
345	0.40	"	1.13	54.5

dehydrogenation of tetralin

z_{napht}

500	1.37	26.5	-0.49	34.9
524.5	2.32	"	-1.35	"

$z_{H_2}^{\dagger}$

500	1.40	-48.4	-0.49	-62.0
530	0.42	"	1.39	-62.1

The real and apparent activation energies and the energy of adsorption of the C,H and O atoms onto the surface of the Yb_2O_3 were calculated from the Card 2/3

Kinetics of the dehydrogenation ...

S/062/62/000/008/002/016
B101/B180

above data (the first figure being the apparent value, the second the true value, in kcal/mole): dehydration of i-propanol ϵ_{H_2O} 25.1, 30.6;

dehydrogenation of i-propanol ϵ_{H_2} 20.6, 22.9; dehydrogenation of tetralin

ϵ_{H_2} 34.7, 36.6; Q_{C-cat} 22.1, 19.4; Q_{H-cat} 53.5; 55.1; Q_{O-cat} 52.5, 49.7.

There are 3 figures and 13 tables.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova
(Moscow State University imeni M. V. Lomonosov)

SUBMITTED: February 13, 1962

Card 3/3

TOLSTOPYATOVA, A.A.; KONENKO, I.R.; BALANDIN, A.A.

Bond energies of carbon, hydrogen, and oxygen atoms of organic molecules with yttrium oxide. Kin.i kat. 3 no.1:103-106 '62.
(MIRA 15:3)

1. Institut organicheskoy khimii imeni N.D.Zelinskogo AN SSSR.
(Chemical bonds) (Yttrium oxide) (Catalysis) (Organic compounds)

Tolstop'yatova, A. A.

60000

Zirconia as a dehydrogenation and dehydration catalyst. A. A. Tolstop'yatova, and V. A. Ferafontov (Dokl. Akad. Nauk SSSR, 1955, 103, 611-614). The following reactions are catalysed by ZrO_2 : cyclohexane \rightarrow benzene + $3H_2$ (480-522°; $\epsilon_{cyc} = 33.4$); $Pr^iOH \rightarrow COMe + 2H_2$ (297-333°; $\epsilon_{cyc} = 30.9$); $Pr^iOH \rightarrow C_2H_4 + H_2O$ (297-333°; $\epsilon_{cyc} = 44.3$); $EtOH \rightarrow MeCHO + H_2$ (316-348°; $\epsilon_{cyc} = 29.1$); $EtOH \rightarrow C_2H_4 + H_2O$ (316-348°; $\epsilon_{cyc} = 38.5$ kg.-cal./mole). The dehydrating activity of ZrO_2 exceeds its dehydrogenating activity. In accordance with theory, the activation energies ϵ_a are very similar for all the reactions, but the ϵ_{cyc} differ considerably. The binding energies of C and H with the catalyst, as calculated from the kinetic data, are very close to those found for Ni and Cr_2O_3 catalysts, whilst the catalyst-O bond-energy is much lower.

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BALANDIN, A.A.; TOLSTOPYATOVA, A.A.; DUDZIK, Z.

Catalytic properties of thorium dioxide in the dehydrogenation and dehydration of alcohols, and in the dehydrogenation of cyclic hydrocarbons. *Kin.i kat.* 2 no.2:273-284, Mr-Ap '61.

(MIRA 14:6)

1. Moskovskiy gosudarstvennyy universitet, kafedra organicheskogo kataliza.

(Thorium oxide)
(Dehydrogenation)
(Dehydration (Chemistry))

TOLSTOPYATOVA, A.A.; BALANDIN, A.A., akademik

Regularities in the catalytic properties of rare earths. Dokl.AN
SSSR 138 no.6:1365-1368 Je '61. (MIRA 14:6)

1. Institut organicheskoy khimii im. N.D.Zelinskogo AN SSSR i
Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova.
(Rare earths) (Catalysis)

BALANDIN, A.A.; STSHIZHEVSKIY, V.; TOLSTOPYATOVA, A.A.

Thermal effect of the reaction of alcohol dehydration affecting
the results of kinetic measurements. Vest.Mosk. un. Ser.2:khim.
17 no.1:30-33 Ja-F '62. (MIRA 15:1)

1. Moskovskiy gosudarstvennyy universitet, kafedra organicheskogo
kataliza.

(Alcohols)

(Heat of denervation)

(Catalysis)

BALANDIN, A.A.; TOLSTOPYATOVA, A.A.; DUDZIK, Z.

Catalytic properties of thorium dioxide in the dehydrogenations and dehydration of alcohols, and in the dehydrogenation of cyclic hydrocarbons. *Kin.i kat.* 2 no.2:273-284 Mr-Ap 1961. (MIRA 14:6)

1. Moskovskiy gosudartsvennyy universitet, kafedra organicheskogo kataliza.

(Thorium oxide) (Dehydrogenation)
(Dehydration (Chemistry))

TOLSTOPYATOVA, A.A.; BALANDIN, A.A.; MATYUSHENKO, V.Kh.; PETROV, Yu.I.

Kinetics of the dehydrogenation and dehydration of alcohols, and of the dehydrogenation of hydrocarbons over WS_2 and MoS_2 catalysts. Izv. AN SSSR Otd.khim.nauk no.4:583-590 Ap '61. (MIRA 14:4)

1. Institut organicheskoy khimii im. N.D.Zelinskogo AN SSSR.
(Dehydrogenation) (Dehydration (Chemistry))
(Molybdenum sulfide) (Tungsten sulfide)

TOLSTOPYATOVA, A.A.; KONEKO, I.R.; BALANDIN, A.A.

Catalytic properties of yttrium oxide. Conversions of alcohols and hydrocarbons. *Kin. i kat.* 2 no.1:135-143 Ja-F '61. (MIRA 14:3)

1. Institut organicheskoy khimii imeni N.D. Zelinskogo AN SSSR.
(Yttrium oxide)
(Alcohols)
(Hydrocarbons)

BALANDIE, A. A.; TOLSTOPIATOVA, A. A.; KONENKO, I. R.

Study of the catalytic conversions of isopropyl alcohol and cyclic hydrocarbons on titanium dioxide (anatase) by means of a differential thermocouple. Izv. AN SSSR. Otd. khim. nauk no. 12:2096-2102 D '60.
(MIRA 13:12)

1. Institut organicheskoy khimii im. N. D. Zelinskogo AN SSSR.
(Isopropyl alcohol) (Hydrocarbons)
(Anatase)

TOLSTOPYATOVA, A.A.; KONEIKO, I.R.; BALANDIN, A.A.

Kinetics of dehydrogenation and dehydration of isopropyl alcohol over titanium dioxide (Anatase). Izv. AN SSSR. Otd. khim. nauk no. 1:38-44 Ja '61. (MIRA 14:2)

1. Institut organicheskoy khimii im. N.D. Zelinskogo AN SSSR. (Anatase) (Isopropyl alcohol)

BALANDIN, A.A.; KOLENKO, I.P.; TOLSTOPELOVA, A.A.

Effect of the preparation method on the catalytic properties of titanium dioxide in reactions of ethyl and isopropyl alcohol and cyclohexane. Izv. AN SSSR. Otd. khim. nauk no. 1:45-50 Ja '61. (MIRA. 14:2)

1. Institut organicheskoy khimii im. M.D. Zelinskogo AN SSSR.
(Titanium oxide) (Ethanol) (Isopropyl alcohols)
(Cyclohexane)

TOLSTOPYATOVA, A.A.; BALANDIN, A.A.; KONENKO, I.R.

Energies of the bonds between reacting organic compounds and the catalytic active centers of titanium dioxide. Izv. AN SSSR. Otd. khim. nauk no.2:214-217 F '61. (MIRA 14:2)

1. Institut organicheskoy khimii im.N.D.Zelinskogo AN SSSR.
(Titanium oxide) (Chemical bonds)

S/062/61/000/001/004/016
B101/B220AUTHORS: Tolstopyatova, A. A., Konenko, I. R., and Balandin, A. A.

TITLE: Kinetics of dehydrogenation and dehydration of isopropyl alcohol on titanium dioxide (anatase)

PERIODICAL: Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk, no. 1, 1961, 38-44

TEXT: By way of introduction the authors offer a survey of reports dealing with the catalytic efficacy of TiO_2 , and then report on the dehydrogenation and dehydration of isopropyl alcohol on anatase. Regarding the production of the catalyst they refer to another report published by them. For the verification of the kinetics of monomolecular heterogeneous-catalytic reactions they proceed from the equation $k = A_1(Z_2 - Z_3) \ln A_1 / (A_1 - m) - (Z_2 + Z_3 - 1)m$ (1). A_1 is the volume of alcohol converted per minute, which is introduced at the rate v ; m is the volume of propylene (or hydrogen) formed per minute; Z_2, Z_3 are the relative adsorption coefficients of the reaction products (hydrogen and acetone)

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Kinetics of dehydrogenation and dehydration...

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in the case of dehydrogenation; propylene and water in the case of dehydration); Z_2 and Z_3 were calculated from $Z = (m_0/m-1)/(100/p-1)$ (2), where m_0 , m are the amounts of the reaction products resulting on the passage of pure alcohol (m_0) and on the passage of a p molar mixture with the reaction product. Since Z is the equilibrium constant of the displacement of the isopropyl alcohol from the catalytic centers by the reaction products, it was possible to calculate also ΔF of the free energy, ΔS of entropy, and ΔH of the heat content. The adsorption coefficient Z_{ac} of acetone was found to be highly dependent on temperature and much less so on the degree to which the catalyst was covered with carbon. The following is indicated for not carbonized TiO_2 : $Z_{ac} = 2.4$ at $282^\circ C$; 1.7 at $294^\circ C$; 1.2 at $310^\circ C$. For carbonized TiO_2 these values amounted to 2.7 , 1.8 and 0.9 . The adsorption coefficient Z_{H_2} of the hydrogen remains constantly 0.9 in the range of $276-306^\circ C$. Moreover, it was found that $Z_{H_2O} = 1.9$ at $282^\circ C$, 1.4 at $294^\circ C$, and 0.9 at $310^\circ C$, whereas $Z_{propylene}$ remains a constant 0.2 in the temperature range of $300-330^\circ C$. A. M. Rubinshteyn and S. G. Kulikov are mentioned.

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Kinetics of dehydrogenation and dehydration... S/062/61/000/001/004/016
B101/B220

There are 1 figure, 8 tables, and 24 references: 8 Soviet-bloc and 13 non-Soviet-bloc.

ASSOCIATION: Institut organicheskoy khimii im. N. D. Zelinskogo Akademii nauk SSSR (Institute of Organic Chemistry imeni N. D. Zelinskiy, Academy of Sciences USSR)

SUBMITTED: June 29, 1959

Card 3/3

S/062/61/000/001/005/016
B101/B220

AUTHORS: Balandin, A. A., Konenko, I. R., and Tolstopyatova, A. A.

TITLE: Effect of the method of production on the catalytic properties of titanium dioxide in the reaction with ethyl alcohol, isopropyl alcohol, and cyclohexane

PERIODICAL: Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk, no. 1, 1961, 45-50

TEXT: The authors were concerned with the investigation of the catalytic properties of titanium dioxide. Here, they studied the dependence of those properties on the method of TiO_2 synthesis. Four specimens were used.

Catalyst 1 (anatase), whose production is described in Ref. 1, a previous report of the authors; catalyst 2, obtained by hydrolysis of $TiCl_4$ by means of ammonia solution in a weakly acid medium at room temperature; catalyst 3, obtained by precipitation of $Ti(OH)_3$ from $TiCl_3$ by means of ammonia at room temperature and conversion to $Ti(OH)_4$ which occurred automatically

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Effect of the method of production....

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while washing the precipitate; catalyst 4, obtained by hydrolysis of ethyl orthotitanate by means of distilled water at room temperature. The further treatment of the hydroxides for the purpose of obtaining TiO_2 was performed according to Ref. 1. Ethanol reacted with these catalysts according to three reactions: a) dehydrogenation, b) dehydration, c) hydrogenation of the resulting ethylene. Moreover, up to 2% ethyl acetate was formed on all four catalysts. The reaction of isopropanol corresponded to data mentioned in Ref. 1. Due to the growing accumulation of carbon on the catalyst, the rate of dehydration increases, whereas that of dehydrogenation decreases. Cyclohexane is dehydrogenated on all four catalysts. The differences between the catalysts regarding the activation energy ϵ and the factor k_0 of the Arrhenius equation are listed in Table 8. The linear function $\log k_0 = a\epsilon + b$ ($a, b = \text{constants}$) is conserved for all reactions. A. M. Rubinshteyn and S. G. Kulikov are mentioned. There are 2 figures, 8 tables, and 6 references: 4 Soviet-bloc and 2 non-Soviet-bloc.

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Effect of the method of production...

S/062/61/000/001/005/016
B101/B220

ASSOCIATION: Institut organicheskoy khimii im. N. D. Zelinskogo Akademii
nauk SSSR (Institute of Organic Chemistry imeni N. D.
Zelinskiy, Academy of Sciences USSR)

SUBMITTED: July 9, 1959

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Effect of the method of production...

S/062/61/000/001/005/016
B101/B220

Влияние способа приготовления катализатора TiO₂ на величины энергии активации

1 Состояние поверхности катализатора *	2 Реакции	3 ккал/М катализатор				4 ккал/М катализатор			
		1	2	3	4	1	2	3	4
		4 ч. о.	7 Дегидрогенизация этилового спирта	22,8	20,4	0,4	12,4	1,0·10 ⁸	1,9·10 ⁷
4 ч. о.	8 Дегидратация этилового спирта	25,0	25,4	0,9	12,5	1,5·10 ⁸	5,8·10 ⁸	8,7·10 ⁸	4,4·10 ⁴
5 п. о.	9 Дегидратация изопропилового спирта	22,9	9,1	28,2	17,2	3,7·10 ⁸	9,1·10 ⁸	1,0·10 ¹⁰	2,8·10 ⁷
4 ч. о.	10 Гидрогенизация этилена	21,2	24,2	13,0	18,8	1,8·10 ⁷	1,0·10 ⁸	1,3·10 ⁴	2,7·10 ⁸
4 ч. о.	11 Дегидрогенизация циклогексана	19,0	14,7	21,0	15,0	8,7·10 ⁸	5,1·10 ⁴	1,3·10 ⁸	3,5·10 ⁴
4 ч. о.	12 Дегидрогенизация изопропилового спирта	9,1	10,4	—	—	1,0·10 ⁴	2,2·10 ⁴	—	—
4 ч. о.	13 Дегидратация изопропилового спирта	24,4	15,3	—	—	4,5·10 ⁸	8,9·10 ⁸	—	—
6 н. о.	14 Дегидрогенизация изопропилового спирта	11,9	—	—	—	1,9·10 ³	—	—	—
5 п. о.	15 Дегидрогенизация циклогексана	25,2	—	—	—	4,9·10 ⁷	—	—	—
4 ч. о.	16 Дегидрогенизация пропилового спирта	19,4	—	—	—	2,3·10 ⁸	—	—	—
4 ч. о.	17 Дегидратация н.пропилового спирта	12,1	—	—	—	4,5·10 ⁴	—	—	—

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Effect of the method of

ε/lg R ₂ +10 ⁴			
3 КИТАЛИЗАТОР			
1	2	3	4
2,84	2,81	3,01	3,05
2,73	2,90	2,70	2,70
2,39	2,28	2,54	2,31
2,91	3,01	3,01	3,10
3,20	3,12	3,53	3,30
2,24	2,40	—	—
2,53	2,02	—	—
2,25	—	—	—
3,27	—	—	—
2,71	—	—	—
3,05	2,73	2,96	2,89
2,80	—	—	—

S/062/61/000/001/005/016
B101/B220

Legend to Table 8: Effect of the method of preparing the TiO₂ catalyst on the amounts of activation energies. 1) Condition of catalyst surface; 2) reaction; 3) catalyst; 4) partially carbonized; 5) completely carbonized; 6) not carbonized; 7) dehydrogenation of ethanol; 8) dehydration of ethanol; 9) dehydration of isopropanol; 10) hydrogenation of ethylene; 11) dehydrogenation of cyclohexane; 12) dehydrogenation of isopropanol; 13) dehydrogenation of propanol; 14) dehydration of n-propanol.

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TOLSTOPYATOVA, A.A.

A.A.Balandin's work in the field of heterogeneous catalysis.
Vest.Mosk.un.Ser.mat., mekh., astron., fiz., khim. 14 no.3:
159-169 '59. (MIRA 13:5)
(Catalysis) (Balandin, Aleksei Aleksandrovich, 1898)

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S/020/60/133/01/36/070
B011/B003

5.1190

AUTHORS: Tolstopyatova, A. A., Balandin, A. A., Academician,
Konenko, I. K.

TITLE: Catalytic Transformations of Alcohols and Cyclic Hydro-
carbons on Titanium Dioxide

PERIODICAL: Doklady Akademii nauk SSSR, 1960, Vol. 133, No.1,
pp. 130 - 133

TEXT: Although titanium dioxide is easily available, it belongs to the little investigated catalysts. The authors wanted to study its catalytic properties with various modes of preparation, in reactions with ethyl-, isopropyl, and n-propyl alcohol as well as with cyclohexane, cyclohexene, and 1,4-cyclohexadiene. Moreover, they wanted to investigate the kinetics of these reactions and the energies of the bonds of C-, H-, and O-atoms with the TiO_2 surface. The method of the differential thermocouple is used for the study of the catalytic reactions. Fig. 1 shows the position of the catalyst in relation to the thermocouple. The electromotive force (emf) was uninterruptedly recorded on

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Catalytic Transformations of Alcohols and
Cyclic Hydrocarbons on Titanium Dioxide

S/020/60/133/01/36/070
B011/B003

a potentiometer of the type ЭПП-09¹⁸ (EPP-09). Apparatus and methods are described in Ref. 4. By combining the above method with analytical methods the authors found that in isopropyl alcohol the surface of the catalyst is coated by carbon-containing sediments. In this way, the relation between dehydrogenation and dehydration of the alcohol is considerably influenced. Within the first 2 - 5 min it is only dehydrogenated. Subsequently, the two processes mentioned take place, and after 20-25 min there occurs dehydration only. Meanwhile, the total amount of the gas liberated per unit of time remains unchanged. All this proves that the centers that are active for dehydrogenation, are poisoned after decarbonization with respect to this reaction, but at the same time are capable of dehydrating the alcohol at the same rate (Table 1). Also the regeneration conditions of the catalyst were determined with the same apparatus: The carbon film was removed after each experiment by means of a strong air current, with the catalyst being previously brought to a temperature exceeding the one of the experiment by 40-50°C. Regeneration usually took 20-30 min. The authors studied the influence of the mode of preparation of TiO₂ on the catalytic properties on four samples: No. 1 - by precipitation of the hydroxide from TiCl₄ with

Card 2/3

Catalytic Transformations of Alcohols and
Cyclic Hydrocarbons on Titanium Dioxide

81725
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B011/B003

water, No. 2 - the same with ammonia. No. 3 by oxidation of freshly prepared $Ti(OH)_3$ (precipitated from $TiCl_3$ with ammonia) with air, and No. 4 by hydrolysis of orthoethyl titanate. It was established by X-ray structural analysis that anatase resulted in No. 1. Table 2 shows the reactions studied in certain temperature ranges, the activation energies E as well as K_0 of the Arrhenius equation; furthermore, the degree of decarbonization of each sample surface. It may be seen from these data and Table 4 that the mode of preparation exerts a great influence on the above-mentioned values and the binding energy. Table 3 shows the dependence of the adsorption coefficient z on ΔH^0 (heat content), the entropy ΔS^0 , and the change in free energy ΔF^0 . The binding energy values were calculated for the first time (Table 4). There are 1 figure, 4 tables, and 11 references: 10 Soviet. 44

ASSOCIATION: Institut organicheskoy khimii im. N. D. Zelinskogo
Akademii nauk SSSR (Institute of Organic Chemistry imeni
N. D. Zelinskiy of the Academy of Sciences, USSR)

SUBMITTED: April 2, 1960

Card 3/3

00300

5.4300

S/195/60/001/004/008/015
B017/B055

AUTHORS: Tolstopyatova, A. A., Balandin, A. A., Stshizhevskiy, V.
TITLE: The Kinetics of Alcohol Dehydration on Tungsten Oxide and
the Energy of Carbon, Hydrogen, and Oxygen Bonds With
Catalysts
PERIODICAL: Kinetika i kataliz, 1960, Vol. 1, No. 4, pp. 558-565

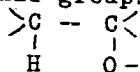
TEXT: The kinetics of the dehydration of ethyl, isopropyl, n-butyl and
tert-butyl alcohol and cyclohexanol on W_2O_5 were investigated under
isothermal conditions. This catalyst was also used to study the de-
hydrogenation of methyl alcohol and 1,2,3,4-tetrahydro naphthalene. The
catalyst was prepared by subjecting the yellow tungsten oxide WO_3 to heat
treatment in air at $350-450^{\circ}C$ for 5-6h, and subsequent treatment with
alcohol vapors at $200-390^{\circ}C$. Thermal effects in dehydration reactions of
n-butyl alcohol are shown in Table 1. Table 2 gives the apparent activa-
tion energies of alcohol dehydration on the tungsten-oxide catalyst. A
relation was found to exist between the apparent activation energy and
Card 1/4

X

88360

The Kinetics of Alcohol Dehydration on Tungsten Oxide and the Energy of Carbon, Hydrogen, and Oxygen Bonds With Catalysts S/195/60/001/004/008/015 B017/B055

the structure of the alcohol. For ethyl and n-butyl alcohol, the apparent activation energy of dehydration is 30 kcal/mole, for isopropyl alcohol, 24 kcal/mole and for tert-butyl alcohol, 18 kcal/mole. The true activation energies of alcohol dehydration (isopropyl and butyl alcohols) were determined at 26.1 kcal/mole for isopropyl alcohol and 33.4 kcal/mole for n-butyl alcohol by means of the kinetic equation by Balandin (Ref. 20). The relative adsorption coefficients of the dehydration products from n-C₄H₉OH and iso-C₄H₉OH as a function of temperature are given in Table 3. From this it may be seen that the relative adsorption coefficients are independent of temperature. Results obtained in the kinetic determination of the dehydration of n-butyl and isopropyl alcohol are shown in Table 5. The true activation energy is 3 kcal/mole higher than the apparent activation energy. The alcohols to be dehydrated are oriented with their functional groups towards the surface of the catalyst:



The kinetic method was used to determine the energies of the bonds of the
Card 2/4

00300

The Kinetics of Alcohol Dehydration on Tungsten Oxide and the Energy of Carbon, Hydrogen, and Oxygen Bonds With Catalysts S/195/60/001/004/008/015 B017/B055

carbon, hydrogen, and oxygen atoms with the active centers of the W_2O_5 surface. The bond energies were found to be a function of the structure of the alcohol. The energies of the bonds of reacting atoms and molecules with the surface of the W_2O_5 catalyst are given in Table 6.

Table 6

Alcohol	Activation Energy of Dehydration	Bond Energies		
		Q_{H-Cat}	Q_{C-Cat}	Q_{O-Cat}
n-Butyl	29.9	56.7	24.1	39.4
Ethyl	29.4	56.4	24.5	39.7
Isopropyl	23.7	52.6	27.6	43.5
Cyclohexanol	21.9	51.5	29.5	44.6
Tert-butyl	17.8	48.7	32.2	47.5

I. Ye. Adadurov and P. Ya. Krayniy are mentioned. There are 6 tables and 25 references: 24 Soviet and 1 German.

Card 3/4

88360
The Kinetics of Alcohol Dehydration on Tungsten S/195/60/001/004/008/015
Oxide and the Energy of Carbon, Hydrogen, and BO17/BO55
Oxygen Bonds With Catalysts

ASSOCIATION: Moskovskiy gosudarstvennyy universitet (Moscow State
University)

SUBMITTED: May 13, 1960

Card 4/4

TOLSTOPYATOVA, A.A.; BALANDIN, A.A.; STSHIZHEVSKIY, V.

Kinetics of dehydration of alcohols on tungsten oxide, and energies of the bonds between the catalyst and carbon, hydrogen, and oxygen. Kin. i kat. 1 no. 4:558-565 N-D '60.

(MIRA 13:12)

1. Moskovskiy gosudarstvennyy universitet,

(Dehydration (Chemistry))

(Tungsten oxide)

(Chemical bonds)

S/062/60/000/008/014/033/XX
B013/B055

AUTHORS: Tolstopyatova, A. A., Balandin, A. A., and Matyushenko,
V. Kh.

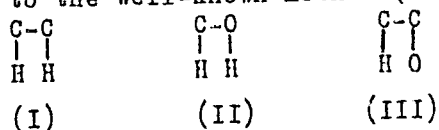
TITLE: Determination of the Bonding Energies of the Atoms of
Organic Molecules Reacting With the MnO Catalyst Surface

PERIODICAL: Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk,
1960, No. 8, pp. 1333-1336

TEXT: The present paper is a continuation of the investigation into the transformation of alcohols on manganous oxide begun in Refs. 1 and 2. The experimental methods and catalyst preparation have been described previously (Ref. 2). Dehydration was carried out between 330° and 380° C. The gaseous products consisted of unsaturated hydrocarbons only. The first series of experiments was performed to determine the apparent activation energy (Table 1, Fig. 1). The activation energy calculated by means of the Arrhenius equation from the relation $\log m = f(1/T)$ was 24.1 kcal/mol. Relative adsorption coefficients of isobutylene (Table 2) and water (Table 3) were determined. The determination of the relative adsorption coefficient
Card 1/3

Determination of the Bonding Energies of the S/062/60/000/008/014/033/XX
 Atoms of Organic Molecules Reacting With the B013/B055
 MnO Catalyst Surface

clients and their temperature dependence permitted calculation by the method described in Ref. 5 of the changes in free energy, enthalpy and entropy produced by displacement of trimethylcarbinol from the active centers of the catalyst by isobutylene and water (Table 4). The true activation energy for the dehydration of trimethylcarbinol, calculated from the relation $\log k = f(1/T)$ ($E = 31.0$ kcal/mol) was found to be 6.9 kcal higher than the apparent activation energy ($E' = 24.1$ kcal/mol). On the strength of this investigation, the authors were able to determine the bonding energies of the atoms reacting with the catalyst surface according to the well-known method (Ref. 6). These reactions are:



(I) dehydrogenation of hydrocarbons; (II) dehydrogenation of alcohols; (III) dehydration of alcohols. The following bonding energies were found:
 $Q_{\text{H-k}} = 50.75$, $Q_{\text{C-k}} = 26.7$ and $Q_{\text{O-k}} = 38.8$. V. N. Kondrat'yev is mentioned.

Card 2/3

Determination of the Bonding Energies of the S/062/60/000/008/014/033/XX
Atoms of Organic Molecules Reacting With the B013/B055
MnO Catalyst Surface

There are 1 figure, 4 tables, and 8 Soviet references.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova
(Moscow State University imeni M. V. Lomonosov).
Institut organicheskoy khimii im. N. D. Zelinskogo Akademii
nauk SSSR (Institute of Organic Chemistry imeni N. D.
Zelinskiy of the Academy of Sciences USSR) ✓

SUBMITTED: January 14, 1959

Card 3/3

S/081/60/000/021/005/018
AC05/AC01

Translation from: Referativnyy zhurnal, Khimiya, 1960, No. 21, p. 50, # 83983

AUTHORS: Tolstopyatova, A. A., Balandin, A. A.

TITLE: The Determination of the Energy of Bonding With Oxide Catalysts by
the Kinetic Method for Developing a Theory of Catalyst Selection

PERIODICAL: Probl. kinetiki i kataliza, 1960, Vol. 10, pp. 351-355

TEXT: The authors determined by the kinetic method the bonding energies of hydrogen, carbon, and oxygen with various oxides and show that, in accordance with the theory, the molecular structure, the catalyst's nature, and the mode of the catalyst's preparation affect the values mentioned.

Summary of the authors

Translator's note: This is the full translation of the original Russian abstract.

Card 1/1

S/062/60/000/010/019/031/XX
B002/B060

AUTHORS: Balandin, A. A., Ferapontov, V. A., and Tolstopyatova, A.A.

TITLE: Ability of Cadmium Oxide to Dehydrogenate Hydrocarbons by Catalysis

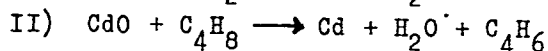
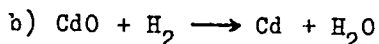
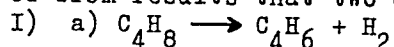
PERIODICAL: Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk, 1960, No. 10, pp. 1751-1758

TEXT: The authors based on experimental work made on the dehydrogenation and dehydration of alcohols by means of oxides to make a theoretical study of the ability of beryllium oxide, magnesium oxide, zinc oxide, and cadmium oxide to dehydrogenate hydrocarbons. For this purpose, the activation energy was calculated on the basis of the multiplet theory (Table 1). The average value found for the adsorption potential of hydrocarbon dehydrogenation was 51.3 for beryllium oxide, 70.7 for magnesium oxide, 116.2 for zinc oxide, and 132.3 kcal/mole for cadmium oxide. The catalytic properties of cadmium oxide were studied experimentally. Cadmium oxide was synthesized by precipitating a cadmium nitrate solution with ammonia, and causing hydroxide to glow at 500°C in air free from CO₂, and was examined by X-rays. Dehydrogenation of cyclohexane was studied between 458° and 540° C (Table 3A). Card 1/2

Ability of Cadmium Oxide to Dehydrogenate
Hydrocarbons by Catalysis

S/062/60/000/010/019/031/XX
B002/B060

The reaction starts at 487°C; at 522°C the catalyst is completely reduced, and dehydrogenation stops. Dehydrogenation of cyclohexene to benzene takes place between 470°C and 520°C (Table 3 B). No cyclohexadiene is formed in this connection. Conversion of butylene to butadiene in nitrogen atmosphere takes place between 575°C and 605°C (Table 4). The degree of conversion was 5.6% at most. Conversion of piperidine to pyridine was examined between 465°C and 520°C (Table 5), and is believed to take place over piperidine. Methane does not reduce cadmium oxide between 500°C and 595°C. It was concluded from results that two different types of reactions take place:



O. V. Krylov, S. Z. Roginskiy, and Ye. A. Fokina are mentioned. There are 1 figure, 5 tables, and 26 references: 24 Soviet, 1 US, and 1 British.

ASSOCIATION: Institut organicheskoy khimii im. N. D. Zelinskogo Akademii nauk SSSR (Institute of Organic Chemistry imeni N. D. Zelinskiy of the Academy of Sciences USSR)

SUBMITTED:
Card 2/2

June 15, 1959

S/062/60/000/012/002/020
B013/B055

AUTHORS: Balandin, A. A., Tolstopyatova, A. A., Konenko, I. R.
TITLE: Investigation of Catalytic Transformations of Isopropyl Alcohol and Cyclic Hydrocarbons on Titanium Dioxide (Anatase) Using a Differential Thermocouple
PERIODICAL: Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk, 1960, No. 12, pp. 2096-2102

TEXT: The authors of the present paper investigated the dehydrogenation and dehydration of isopropyl alcohol (Tables 1-4) and the dehydrogenation and irreversible catalysis of hydrocarbons, cyclohexane (Table 5), cyclohexene (Tables 6 and 7) and 1,4-cyclohexadiene (Table 8) - on a modification of titanium dioxide (anatase) using a differential thermocouple (Chromel - cupro nickel). The latter had 10 junctions each at both sides of the mica sheet to which it was attached. All the 20 junctions were on the one half of the sheet. The distribution of the catalyst in the quartz tube containing the thermocouple is shown in Fig. 1. A continuous reaction vessel was used for the kinetic experiments (Refs. 4 and 5). The temperature was maintained with
Card 1/3

Investigation of Catalytic Transformations of S/062/60/000/012/002/020
Isopropyl Alcohol and Cyclic Hydrocarbons B013/B055
on Titanium Dioxide (Anatase) Using a
Differential Thermocouple

an accuracy of $\pm 5^{\circ}$. The evolution rate and quantity of gaseous products were measured and recorded by a ГСП-10 (GSP-10) gas meter (Fig. 2). X-ray analysis of the catalyst prepared from titanium tetrachloride showed it to be anatase. The transformation of isopropyl alcohol on TiO_2 was accompanied by a marked change in thermal effect during the first 15-20 min (Table 1, Fig. 3a). The analysis of the gaseous products formed in the reaction showed that the alcohol is practically only dehydrogenated during the first 7-8 min. After this period dehydrogenation and dehydration take place simultaneously and after 15-20 min dehydration occurs alone. From a comparison of the thermocouple data with the analytical data of the reaction products and the degree of carbon deposition on the catalyst surface it can be seen that the rates of dehydrogenation and dehydration, and the carbon content of the surface run parallel. The change in selectivity at otherwise unchanged general activity of TiO_2 is probably due to the accumulation of carbon on the surface. Deposition of carbon on the surface of Cr_2O_3 (Fig. 3b), however, was found to have no effect on its activity and selectivity. Summarily, the

Card 2/3

Investigation of Catalytic Transformations of Isopropyl Alcohol and Cyclic Hydrocarbons on Titanium Dioxide (Anatase) Using a Differential Thermocouple

S/062/60/000/012/002/020
B013/B055

authors conclude that a gradual accumulation of carbon on the anatase surface almost entirely suppresses the dehydrogenation of the alcohol and promotes its dehydration, but does not affect the total activity of the catalyst. The catalyzed irreversible transformation of cyclohexene and cyclohexadiene is suppressed by the accumulation of carbon on TiO_2 , whereas the dehydrogenation of cyclohexene, cyclohexane, and cyclohexadiene is promoted by this process. There are 3 figures, 7 tables, and 12 Soviet references. ✓

ASSOCIATION: Institut organicheskoy khimii im. N. D. Zelinskogo Akademii nauk SSSR
(Institute of Organic Chemistry imeni N. D. Zelinskiy of the Academy of Sciences USSR)

SUBMITTED: June 26, 1959

Card 3/3

S/020/60/134/003/030/033/XX
B004/B064AUTHORS: Balandin, A. A., Academician, Tolstopyatova, A. A., and
Stshizhevskiy, V.TITLE: The Catalytic Activity of Tungsten Pentoxide 2]PERIODICAL: Doklady Akademii nauk SSSR, 1960, Vol. 134, No. 3,
pp. 625 - 628

TEXT: The authors investigated the dehydration of ethanol, isopropanol, t-butanol, cyclohexanol, methanol, and Tetralin with W_2O_5 under isothermal conditions. Blue W_2O_5 was obtained from tungstic acid at 350 - 400°C in an air current. First, WO_3 was formed and then reduced to W_2O_5 during the reaction with the alcohols at 200 - 300°C. The reaction apparatus operated continuously, and the liquid substances were automatically added. The gaseous products were collected in an automatic Patrikeyev gasometer, and analyzed with a BTM (VTI) apparatus or chromatographically. In the liquid product of catalysis, the amount of

Card 1/4

The Catalytic Activity of Tungsten
Pentoxide

S/O20/60/134/C03/C30/C33/XX
B004/B064

unsaturated hydrocarbons was determined by the method of Kaufman-Gal'pern. Since the endothermic effect of reaction affected the results of measurement, the catalyst was diluted in a ratio of 2 : 3 with quartz of the same grain size, the alcohol with water or with the corresponding unsaturated hydrocarbon. Under these conditions, the process was isothermal. By determining the apparent activation energy (Table 1) it was found that the primary alcohols were dehydrated with the same energies (approximately 30 kcal/mole), that the activation energy of the secondary alcohols was about 6 kcal/mole lower than that of the primary ones, and that the activation energy of the tertiary alcohol was approximately 6 kcal/mole lower than that of the secondary ones. The reaction constant and the relative adsorption coefficients z_2 and z_3 of water and the unsaturated hydrocarbon were computed (Table 2) by Balandin's method (Ref. 3) with reference to the adsorption coefficient of alcohol. Checking by introduction of the experimental data into Balandin's equation confirmed the validity of this equation (Table 3). Table 4 gives the actual activation coefficients for isopropanol and n-butanol, which are approximately 3 kcal/mole higher than the apparent ones. For the binding

Card 2/4

The Catalytic Activity of Tungsten
Pentoxide

S/020/60/134/003/C30/C33/XX
B004/B064

energies Q_{HC} , Q_{CC} , Q_{OC} of the H, C, and O atoms reacting with the surface of W_2O_5 , the following was computed from Balandin's kinetic equation: Table 5:

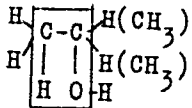
Alcohol	Activation energy of dehydration [kcal/mole]	Binding energy		
		Q_{HC}	Q_{CC}	Q_{OC}
n-butanol	29.9	56.7	15.9	39.2
ethanol	29.4	56.4	16.3	39.5
iso-propanol	23.7	52.3	19.4	43.3
cyclohexanol	21.9	51.5	21.3	44.4
t-butanol	17.8	48.7	24.0	47.3

Although the hydrogen atoms and the radicals substituting them do not directly take part in the reaction: ✓

Card 3/4

The Catalytic Activity of Tungsten
Pentoxide

S/O20/60/134/003/C30/C33/XX
B004/B064



, they affect the binding energy between the C, H, and O atoms and the catalyst. The authors mention a paper by I. Ye. Adadurov and P. Ya. Krayniy. There are 5 tables and 8 Soviet references.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosova
(Moscow State University imeni M.V. Lomonosov)

SUBMITTED: May 17, 1960



Card 4/4

TOLSTOPYATOVA, A.A.; BALANDIN, A.A., akademik; KONENKO, I.R.

Catalytic conversions of alcohols and cyclic hydrocarbons on
titanium dioxide. Dokl.AN SSSR 133 no.1:130-133 J1 '60.
(MIRA 13:7)

1. Institut organicheskoy khimii imeni N.D.Zelinskogo Akademii
nauk SSSR. (Alcohols) (Hydrocarbons) (Titanium oxide)

TOLSTOPYATOVA, A.A.; BALANDIN, A.A.; MATYUSHENKO, V.Kh.

Dehydrogenation of alcohols and cyclohexene on MnO . Izv. AN
SSSR Otd.khim.nauk no.5:787-793 My '60. (MIRA 13:6)

1. Moskovskiy gosudarstvennyy universitet imeni M.V.Lomonosova i
Institut organicheskoy khimii imeni N.D.Zelinakogo Akademii
nauk SSSR.

(Cyclohexene) (Cyclohexanol) (Isopropyl alcohol)

TOLSTOPYATOVA, P.A.

PART I BOOK REFERRATIONS SET/1164

Вещногорский завод химического машиностроения по сплавам титана и алюминия. Лт., Москва, 1957
Материалы 1-го съезда... (Rare Metals and Alloys) Transactions of the
First All-Union Conference on Rare-Metal Alloys Moscow, Metallurgizdat, 1960.
Part 1, 1960 copies printed.

Специальная литература: Атомиздат, Москва, 1958. Индустриальный титан: СССР
Индустриальный титан: перевод с немецкого языка. Москва, 1958.
Книжка по редким металлам при машиностроении. О.К. Казарян, Техн. М.:
П.О. Издательство.

М.И. И.И. Соперников, К.И. Издательство Дома О.К. Казарян, Техн. М.:
П.О. Издательство.
PURPOSE: This collection of articles is intended for metallurgical engineers,
physicists, and workers in the machine-building and radio-engineering industries.
It may also be used by students of schools of higher education.

CONTENTS: The collection contains technical papers which were presented and dis-
cussed at the First All-Union Conference on Rare-Metal Alloys held in the In-
stitute of Metallurgy, Academy of Sciences USSR in November 1957. Results of
investigations of rare-metal alloys, titanium-based alloys with ad-
ditions of rare metals are presented and discussed along with investigations of
titanium, vanadium, niobium and their alloys. The effect of rare-earth metals
on properties of magnesium alloys and steels is analyzed. The use of titanium
as a dehydrating catalyst, electropolymerizing material, and material suitable for
making plugs for automobile electrical systems are discussed. Also, the ef-
fect of the addition of certain elements on the properties of heat-resistant
steel is examined and alloys with special properties (particularly
superconductive alloys) are discussed. Some of the articles
and non-serial references mentioned in the articles.

PART II. TITANIUM AND COPPER-BASED
ALLOYS WITH RARE-METAL ADDITIVES

Данилов, Г.П., Л.П. Даниловский, and В.Т. Жуковская. Investigations of Alloys of the Titanium-Copper-Aluminum and Titanium-Copper-Aluminum Systems	24
Найзер, М.В., Г.П. Данилов, and Л.А. Сидорова. Effect of Rare Metals on the Stability of Titanium and of Some Titanium Alloys	42
Найзер, М.В., and Л.А. Сидорова. Investigation of Titanium-Aluminum- Vanadium Ternary Alloy Systems	52
Данилов, Г.П., Г.С. Сидорова, Л.А. Сидорова, Л.А. Сидорова, and Л.А. Май- серов. High-Strength and Heat-Resistant Alloys of the Copper-Cobalt-Titanium Alloy System	63

Rare Metals (Cont.) SET/1164

PART III. ZIRCONIUM, TANTALUM, NIOBIUM, ERBIDIUM AND ALLOYS BASED ON THEM	
Данилов, Г.П., Л.П. Даниловский, and Л.А. Сидорова. Zirconium as a Dehydrating Catalyst	72
Гуляев, М.А., and В.М. Сердюков. Zirconium Alloys	80
Сидорова, Л.А., Л.А. Сидорова, Л.А. Сидорова, and Л.А. Сидорова. Electro- plating with Zirconium	111
Данилов, Г.П., and М.Д. Пустовалов. Electrical Contacts Made of Zirconium Sintered with Vanadium	122
Данилов, Г.П., and М.Д. Пустовалов. The Possibility of Using Alloys on Tungsten with Zirconium for Making Contacts for Automobile Electrical Equipment	133
Данилов, Г.П., and М.Д. Пустовалов. Properties of Zirconium, Niobium and of Alloys Based on Them	136

Card 4/8

Tolstopyatova, A.A.

Abstracts and Index. Institute of Chemistry, USSR Academy of Sciences. 1960. 300 pages.

Problematic literature in Russian. [3] 10. Problems of Heterogeneous Catalysis (Problems of Heterogeneous Catalysis. [vol. 1] 10. Problems and Physical Chemistry of Catalysis) Moscow, Institute of Chemistry, USSR Academy of Sciences, 1960. 461 p. 300 copies printed. 2,600 copies printed.

Editor: S.M. Bogdanov, Corresponding Member of the Academy of Sciences USSR, and O.Y. Egorov, Candidate of Chemistry, Ed. of Publishing House: A.I. Kharin, Moscow, 1960.

PURPOSE: This collection of articles is addressed to physicists and chemists and to the community of scientists in general interested in recent research on the physics and physical chemistry of catalysis.

CONTENTS: The articles in this collection were read at the conference on the Physics and Physical Chemistry of Catalysis organized by the USSR Academy of Sciences (Section of Chemical Sciences, Academy of Sciences USSR) and by the Academic Council on the problem of "the scientific bases for the selection of catalysis". The conference was held at the Institute of Chemistry USSR Academy of Sciences (Institute of Physical Chemistry of the USSR) in Moscow, March 20-23, 1960. Of the great volume of material presented at the conference, only papers not published elsewhere were included in this collection.

I. PROBLEMS OF STRUCTURAL AND INDEX CORRELATIONS IN CATALYSIS

Malozemov, A.D. [Institute of Organic Chemistry of the USSR] Structural and Index Factors in the Elementary Stages of Catalysis 304

Malozemov, A.D., and P. Tyndal. The Role of σ -Complexes in Catalysis. Kinetics of Hydrogenation of Isopropyl Alcohol in the Presence of Copper, Cobalt, Nickel, Iron, Silver, Platinum and Palladium 309

Klimov, S.M., and Malozemov, A.D. [Institute of Organic Chemistry of the USSR] Bond Energy of Nickel, Iron, Platinum and Palladium Catalysts with the Elements of Organic Compounds 314

Polozhinov, A.A., and Malozemov, A.D. [Institute of Organic Chemistry of the USSR] Kinetic Study of Catalysis by Dehydrogenation of Ethanol by the Kinetic Method with a View to Working Out a Theory for the Selection of Catalysts 321

Yastrebina, V.P., and Malozemov, A.D. [Institute of Organic Chemistry of the USSR] Investigation of the Elementary Stages of the Alcohol Dehydrogenation Reaction over Al_2O_3 Catalysts 325

Malozemov, A.D., and K.P. Sazonov [Institute of Organic Chemistry of the USSR]. Catalytic Properties of Sodium-Naphthalene Anions as Directive Structures 329

II. RADICAL AND CHAIN CONTRIBUTIONS IN CATALYSIS

Tolstopyatova, A.A. [Institute of Chemical Physics of the USSR]. Chain Contribution in Heterogeneous Catalysis 329

Amos, A.V. [Institute of Inorganic Chemistry of the USSR]. Kinetics of the Reaction of Ethylene with Ethylene Oxide Catalysts and the Mechanism of Certain Catalytic Reactions 331

Yastrebina, V.P., and Tolstopyatova, A.A. [Institute of Chemical Physics of the USSR] Kinetic Study of the Chain Characteristic of Surface Reactions 332

Yastrebina, V.P., and Tolstopyatova, A.A. [Institute of Chemical Physics of the USSR] On the Problem of the Possibility of Chain Mechanisms During Catalysis Over Metals 333

Rylov, V.S., and S.I. Kurbatov [Institute of Organic Chemistry, Soviet Academy of Sciences]. Role of Free Radicals in the Mechanism of Radicalyzed Oxidation of Alcohols Under the Action of Small Quantities of Carbon Monoxide in the Presence of Platinum 334

TOLSTOPYATOVA, N.S. (Leningrad, P-49, ul. Lizy Chaykinoy, d.18, kv.1)

Occupational epicondylitis of the arm. Vest. rent. i rad. 36 no.5:
46-51 S-0 '61. (MIRA 15:1)

1. Iz rentgenologicheskogo otdeleniya (rukovoditel' - prof. A.V. Grinberg) Leningradskogo nauchno-issledovatel'skogo instituta gigiyeny truda i professional'nykh zabolevaniy (dir. - prof. Z.E.Grigor'yev).

(ELBOW_DISEASES)

(OCCUPATIONAL DISEASES)

TOLSTOPYATOVA, N. V.

~~Work experience with parafin in the preparation of orthopedic apparatus. Med. sestra, Moskva no. 9:29-30 1951 (GIML 21:1)~~

1. Author is a plaster-of-paris cast technician belonging to the Bone Division (Head -- A.P. Sokol'skaya) of Ivanovo Oblast Tuberculosis Sanatorium No 2 (Head Physician -- Honored Physician RSFSR A. L. Gal'perin).

5.3200
5.1190

67260

5(3)

SOV/20-129-4-22/68

AUTHORS: Balandin, A. A., Academician, Karpeyskaya, Ye. I., Tolstopyatova,
A. A.TITLE: On the Irreversible Catalysis in the Presence of Metallic Rhenium ✓PERIODICAL: Doklady Akademii nauk SSSR, 1959, Vol 129, Nr 4, pp 795-798
(USSR)

ABSTRACT: Metallic rhenium, applied to active carbon dehydrogenates the cyclic 6-membered hydrocarbons. On an ammonium perrhenate (Re-1) catalyst small quantities of cyclohexene and also benzene are produced from cyclohexane (Ref 3). This is not the case on a catalyst of a dioxane complex of rhenium-acid-anhydride (Re-2). The authors investigated the transformation of cyclohexene on Re-1 and Re-2 to find the reasons for cyclohexene formation on Re-1. Moreover, the capability of rhenium to bring about irreversible catalysis (expression by N. D. Zelinskiy) was to be investigated. The gaseous reaction products were collected in the gasometer by V. V. Patrikeyev. They consisted of hydrogen with 6-8% methane. The experiments were made between 215° and 465°C. For both catalysts a temperature range was distinctly observable in which no gas is formed and the process takes place according to equation ✓

Card 1/3 $3C_6H_{10} = C_6H_6 + 2C_6H_{12}$. At higher temperatures hydrogen is separated.

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On the Irreversible Catalysis in the Presence of
Metallic Rhenium

SOV/20-129-4-22/68

Catalysates of several experiments with Re-1 and Re-2 were poured together. After the removal of the incompletely reacted cyclohexene benzene was chromatographically separated from cyclohexane. Figures 1 and 2 show the results of various experiments which were well reproducible. It may be seen from a comparison of the kinetic data on dehydrogenation of cyclohexene and the disproportionation of hydrogen on Re-1 and Re-2 that the activation energies of the processes are in agreement on both catalysts. Apparently both processes stop at the same stage of reaction. Table 3 shows that on Re-2 benzene is formed from cyclohexene twice as rapidly as from cyclohexane. According to M. Ya. Kagan and N. A. Shcheglova (Ref 6) cyclohexane is dehydrogenated on Pt by 4500 times more slowly than is the transformation rate of cyclohexene. On the basis of these data and according to M. Ya. Kagan and R. M. Flid (Ref 7) the mentioned researchers draw the conclusion that the dehydrogenation of the cyclic hydrocarbons takes place step-wise (see Scheme). The results obtained by the authors are in contradiction with this conclusion: cyclohexane is dehydrogenated on rhenium not according to a step-mechanism but according to a sextet mechanism. Benzene was formed on Re-1 from cyclohexene about twice as rapidly (Table 4).

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SOV/20-129-4-22/68

On the Irreversible Catalysis in the Presence of
Metallic Rhenium

It may hardly be assumed that the dehydrogenation of cyclohexane on two catalysts of the same chemical properties and on the same carrier should proceed according to two different mechanisms. Cyclohexene is no intermediate (in contradiction to the scheme of references 6, 7) since it was not obtained in the reaction on Re-2. Cyclohexene thus results from a side-reaction which takes place to a small degree besides the sextet dehydrogenation. There are 2 figures, 4 tables, and 8 references, 7 of which are Soviet.

ASSOCIATION: Institut organicheskoy khimii im. N. D. Zelinskogo Akademii nauk SSSR (Institute of Organic Chemistry imeni N. D. Zelinskiy of the Academy of Sciences, USSR) ✓

SUBMITTED: July 24, 1959

Card 3/3

VIKSNE, A.; VIKSNE, J.; DENISOVA, U. [translator]; KASPARSONA, G.
[translator]; LEGZDINA, Zh. [Legzdina, Z.] [translator];
POISHA, Ya. [Poisa, J.] [tranalator]; TOLSTOPYATOVA, R.
[translator]; ALKSNE, B., red.; BERZINA, K., red.; SILINS, V.,
tekhn. red.

[Riga Zoological Garden] Rizhskii zoologicheskii sad. Riga,
Latvijas Valsts izdevnieciba, 1957. 1 v. (chiefly illus).
(MIRA 14:12)

(Riga--Zoological gardens)

TOLSTOSHEV, A.V., student V kursa OMF

Methods of testing central photographic shutters. Trudy MIIGAIK
no.36:117-122 '59. (MIRA 13:4)

1. Studencheskoye nauchnoye obshchestvo Moskovskogo instituta
inzhenerov geodezii, aerofo os"yemki i kartografii.
(Shutter, Photographic--Testing)

TOLSTOSHEY, A.N., assistant

Analytical method for determining the basic norms for switching operations. Nauch.trudy KHIIT no.55:39-50 '62.

Device for recording the parameters of the movement of switcher locomotives (electric switcher recorder). 81-87 (MIRA 16:10)

KON'KOV, P.S., , kand. tekhn.nauk, dots.; DONTSOV, A.Ya., inzh.;
YURCHENKO, I.F., inzh.; ANGELEYKO, V.I., retsenzent;
BABENKO, V.I., retsenzent; ZAPREVSKIY, G.S., retsenzent;
KRIMNUS, G.Kh., retsenzent; MANIN, I.I., retsenzent;
NAUMOV, G.K., retsenzent; TOLSTOSHEY, A.N., retsenzent;
TUCHKEVICH, T.M., retsenzent; FEDORETS, V.M., retsenzent;
FEL'DMAN, M.F., retsenzent; FRANKOV, N.Ya., retsenzent;
USENKO, L.A., tekhn. red.

[Establishing work norms in railroad transportation] Tekh-
nicheskoe normirovanie truda na zheleznodorozhnom transporte.
Moskva, Transzheldorizdat, 1963. 366 p. (MIRA 16:9)
(Railroads—Production standards)

LEBEDEVA, A.P.; TOLSTOSHEY, O.N.

Contamination of soil with hexachloran from its use in agriculture.
Gig. i san. 26 no.11:15-18 N '61. (MIRA 14:11)

1. Iz Ukrainского nauchno-issledovatel'skogo instituta kommunal'noy
gigiyeny. (CYCLOHEXANE) (SOIL POLLUTION)

TOLSTOSHEV, O.N.

Enlarged conference on sanitary protection of the sea near the
shoreline. Gig. 1 san. 24 no.1:88-89 Ja '59. (MIRA 12:2)
(SEA WATER--POLLUTION)

BARAN, N.A.; TOLSTOSHEY, O.N.

Hygienic problems of sewage and waste disposal in towns of the
Donets Basin [with summary in English]. Gig. i san. 24 no.2:22-26
F '59. (MIRA 12:3)

1. Iz Ukrainskogo nauchno-issledovatel'skogo instituta kommunal'noy
gigiyeny.

(SANITATION

hyg. problems of sewage & waste disposal of towns
of Donbas (Rus))

TOISTOSHEY, O.N.

Hygienic features of garbage disposal in Kiev. Gig. i san. 23 no.12:60-
62 D '58. (MIRA 12:1)

1. Iz Ukrainskogo nauchno-issledovatel'skogo instituta kommunal'noy
gigiveny.

(SANITATION
garbage disposal in Russia (Rus))

TOLSTOSHEY, O.N. (Kiyev)

Significance of a regular trash pick-up service for the public health in cities. Vrach.delo no.6:621-624 Je '58 (MIRA 11:7)

1. Ukrainskiy Institut kommunal'noy gigiyeny.
(KIEV--REFUSE AND REFUSE DISPOSAL)

Tolstauklov, A.S.

NOV 8-39-6-21/22

3(2),3(4)
ATTACHED
TITLE
PERIODICAL
ABSTRACT

None Given
Chronicle (Khronika)

Geodesiya i kartografiya, 1959, Nr 6, pp 74-75 (USSR)

At the Moskovskiy Institut inzhenerov geodesii, aerofotogrammetrii i kartografii (Moscow Institute of Geodesic, Aerial Survey and Cartographic Engineers), the Ordinary Scientific Conference took place on April 22-24. A. I. Kozlov, Docent, Candidate of Philosophic Sciences, spoke on the subject "The Role of the Geodesic Materialistic Philosophy". A. M. Zhukovskiy, Chief of the Glavnoye upravleniye geodesii i kartografii, spoke "On the Seven-year Plan for the Academy and the Scientific Progress".

Following reports were delivered in the geodesic section:

Ac. M. Pavlov, Professor, "Some Integrals of the Surface of the Earth"; A. V. Kondrakhin, Docent, "Radio-electronics and Geodesy"; G. V. Krasnitskiy, Docent, "Accuracy in the Solution of Inverse Position Computations by the Coordinates of Different Geodesic Systems"; P. S. Shukhin, Docent, "Gravity in the Present Stage of Development"; Yu. V. Kozlov, Assistant, reported on the influence of rounding errors on the accuracy of solution of linear equation systems; I. M. Kuznetsov, Candidate of Technical Sciences, spoke on the subject "The Rules of Distribution of Errors in the Solution of the Problem in Surveys"; I. M. Zhelezovskiy, Post-graduate student, reported on the solution of the problem of the adjustment of geodesic observations by M. Kopylovskiy, Docent, demonstrated an apparatus designed by him for parallaxless traversing with a short constant vertical baseline. The following reports were delivered in the aerophotogrammetric section: A. S. Valuyev, Docent, reported on a parallaxless method; an additional device to the stereocomparator; N. M. Veselovskiy, Docent, spoke on the possibility of generalizing the formulas for the air survey of outcrops and altitudes; B. M. Medvedev and E. P. Zakharov, Docents, reported on a band-shaped optical shutter for aerial cameras; M. G. Gromov and Engineer V. I. Terkhov on the scheme of the automatic device for the automatic entry of the survey data into the route for air surveys; K. P. Khramov reported some simplifications for the completion of constant of aerial cameras; Yu. K. Kuznetsov, Post-graduate student, spoke on the use of rapid film recording for the investigation of aerial camera shutters; V. I. Grishin, Engineer of the Geodesic Aerial Camera KRYM, spoke "On Some Results and Plans in the Execution of Large-scale Photogrammetric Surveys". The following reports were delivered in the cartographic section: Professor V. I. Sukhov spoke on the content of the new map on a scale of 1:2,500,000; Professor A. I. Prokhorov spoke on "Mineral Resources of the USSR and Their Representation on Economic Maps"; S. S. Sidorov, Assistant, reported on the method of geographic field research during the preparatory editorial work at the subject of cartography; A. S. Kuznetsov, Assistant, reported on the improvement of relief representation of 1:10,000; Ya. S. Bilich, the topographic map on a scale of 1:10,000; In. S. Bilich, Assistant, reported on maps of the section of building of apparatus; I. V. Yakovlev, Assistant, reported on the life of the Academy; I. G. Sidorov, Assistant, reported on the accuracy in determining physical heights; Engineer V. M. Kuznetsov on vertical aerial systems for highly accurate optical theodolites; K. S. Durov, Assistant, on sighting with telescopes with some plates; E. P. Zakharov, Assistant, on the automation of the evaluation of image couples.

Card 2/4

Card 3/4

TOLSTOUKHOV, A.S.

Practice of conducting preliminary topographic and geographic investigations. Geod. 1 kart. no. 2:49-53, F. 163. (MIRA 16:3)
(Topographic maps)

AUTHOR: Tolstoukhov, A. S., Graduate Student SOT/154-58-1-16/22

TITLE: Some Peculiarities in Representing the Relief of Plains on Topographic Maps on a Scale of 1 : 10 000 - Shown in the Case of the Terek Delta (Nekotoryye osobennosti izobrazheniya ravninnogo rel'yefa na topograficheskikh kartakh masshtaba 1 : 10 000 - na primere del'ty reki Tereka)

PERIODICAL: Izvestiya vysshikh uchebnykh zavedeniy. Geodeziya i aéro-fotos"yemka, 1958, Nr 1, pp 123-131 (USSR)

ABSTRACT: Topographic maps are used to a large extent in studying the characteristics of various geographical areas in the USSR. This is true especially of maps on a scale of 1 : 10 000. These maps in use today do, however, no longer meet all requirements. Above all, the representation of the relief with the micro-forms in the plains on topographic maps (large scale) has not yet been sufficiently developed. The author and his collaborators therefore selected the delta of the Terek River for their investigations. One of the peculiarities amongst others of the delta is the elevated position of the central river bed in the delta (up to four meters). On both sides of this river bed the terrain gradually falls.

Card 1/3

SOV/154-58 1-16/22

Some Peculiarities in Representing the Relief of Plains on Topographic Maps on a Scale of 1 : 10 000 - Shown in the Case of the Terek Delta

The numerous other branches of the Terek therefore are situated lower than the central river bed. Innumerable small elevations, dome-shaped and overgrown, are characteristic of this delta region, as well as the amorphous salt soil. The author discusses in detail the methods of relief representation developed by him. The combination of the micro relief (the ground with small dome-shaped elevations), of the flora on the one hand and of the semi-arid area on the other hand, requires a comprehensive and uniform representation. The author proposes new topographic symbols as marks of the small dome-shaped elevations. With respect to the flora the author suggests the method already developed by K. A. Borodina. The elevation marks and figures should be printed on topographic maps in a way that they follow the turns of the contours. The present density of topographic marks within one square decimeter has been insufficient and should therefore be increased considerably. There are 11 figures and 5 references, 5 of which are Soviet.

Card 2/3

SOV/154-58-1-16/22

Some Peculiarities in Representing the Relief of Plains on Topographic
Maps on a Scale of 1 : 10 000 - Shown in the Case of the Terek Delta

ASSOCIATION: Moskovskiy institut inzhenerov geodezii, aerofotos^{ny}enki i
kartografii
(Moscow Engineering Institute of Geodesy, Aerophotography
and Cartography)

Card 3/3

1625164A P. 1, 1.2.

AUTHOR: Bol'shakov, V. D., Candidate of Technical Sciences, spoke
 Scientific and Technical Conference of the NIKA i K (Nauchno-
 tekhnicheskaya konferentsiya NIKA i K) II

TITLE: Ispytaniya vyznaniya uchebnykh i avtomaticheskikh kartograficheskikh
 i topograficheskikh kart (Tests and automatic cartographic and
 topographic maps)

PERIODICAL: Izvestiya vuzovskoye nauchno-issledovatel'skoye i inzhener-
 noye otdeleniye NIKA i K (News of the Scientific and Engineering
 Department of NIKA i K), No. 2, pp 114-115 (USSR)

ABSTRACT:

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G. A. GIMBARZ, Docent, Candidate of Technical Sciences, spoke on "The Relations Between Distortions in Cartographic Projections." L. A. Bogolov, Candidate of Technical Sciences, reported on "Topographical Deciphering From the Airplane and the Use of Aerial Photographs in Cartographic Inaccessible Regions." A. S. BILICHUKHIN, Assistant, spoke on "The Relief Representation of Lines on Topographical Maps (Scale 1:10 000 000)." V. D. Kishner, Professor, Doctor of Geographical Sciences, dealt with the basic geographic structures of Antarctica and the consequent cartographical peculiarities of the region.

Engineer Ye. M. Feilitov reported on the conference held in the NIKA i K (Moscow Engineering Institute of Geodesy, topography, and Cartography) from May 6 to 10. Participants discussed various questions in relation with the design of geodetical and cartographical instruments. More than 500 delegates from many universities and scientific institutions, as well as 82 representatives of different agencies in Leningrad, Kiev, Sverdlovsk, and other cities, participated in this conference. The Deputy Head of the GUGM, M. D. Kom'zhin, read a paper on "Scientific Research in Aerial Camera Design." S. V. Felitshev, Docent, reported on "The Present State of Production of Geodetical Instruments, and Development of New Instruments of Photogrammetrical Instruments in the USSR and on developments in this field. In the different sections questions relating to the design of geodetical and photogrammetrical instruments as well as the manufacture for aerial photography were discussed. Docent S. V. Vilkovskiy, Engineer I. V. Zdobnikov reported on geodetic engineering and Engineer I. V. Zdobnikov reported on geodesic instruments. Engineer I. V. Masarov, Candidate of Technical Sciences, dealt with the new range finders of greater precision. V. A. Valchuk, Docent of Technical Sciences, on optical range finders of geodesic type. Engineer I. I. Andrianova and Ia. P. Popov, Candidates of Physical-Mathematical Sciences, on optical range finders of geodesic type. Engineer L. V. Labynovich spoke on the use of light alloys in the manufacture of geodetical instruments. Docent S. M. Muravyev and Engineer Y. K. Sayenko reported on new developments in the production of geodetical instruments. Professor D. Ya. Gal'perin dealt with the optical systems in geodetical instruments. Engineer A. M. Burago, Engineer B. A. Al'ko, Docent V. A. Kruselis, Docent I. M. Firson, and Engineer A. V. Dubrovskiy, reported on the results of the Scientific and Technical Conference held in Kiev (Planning and Production of Geodetical Instruments).

Card 2/3

Card 3/3

TOLSTOUKHOV, A. S., Ass't.

"On the Representation of Reliefs of Plane Areas on Topographic Maps"

report presented at a Scientific-Technical Conference at Moscow Inst. of Geodesy,
Aerial Photography and Cartography Engineers, 24-26 April 1958.
(Geodeziya i kartografiya, no. 6, pp. 79-80, 1958)

TOISTOUKHOV, A.S., aspirant

Certain peculiarities in representing lowland relief on topographic maps as exemplified by the Terek Delta. Izv. vys. ucheb. zav.; geod. i aerof. no. 1:123-131 '58. (MIRA 11:7)

1. Moskovskiy institut inzhenerov geodezii, aerofotos'yemki i kartografii.
(Terek Valley--Topographical surveying)

TOLSTOUKHOVA, L.I.; BYKHOVSKAYA, A.M.

I-131 treatment of postoperative recurrences of thyrotoxi-
cosis. Med.rad. 7 no.11:5-9 N^o62. (MIRA 16:9)

1. Iz kafedry rentgenologii i radiologii (zav. - prof. Ye.L.
Kevesh) Kuybyshevskogo meditsinskogo instituta.

(HYPERTHYROIDISM)
(IODINE ISOTOPES--THERAPEUTIC USE)

UZHANSKIY, Ya.G., prof., KACHANOVA, S.G., dots., TOLSTOUKHOVA, L.I., dots.

Fifth conference of the Ural branch of the All-Union Society of
Pathophysiologists, Pat.fiziol. i eksp.terap 2 no,3:60-62

My-Je '58

(MIRA 11:7)

(PHYSIOLOGY, PATHOLOGICAL)

UZHANSKIY, Ya.G., professor, predsedatel' pravleniya; TOLSTOUKHOVA, L.I., dotsent, sekretar'.

At the Ural branch of the All-Union Society of Pathophysiologists; on the organization and the first conference of the Ural branch of the All-Union Society of Pathophysiologists. Arkh.pat. 15 no.2:90-92 Mr-Ap '53.
(MLRA 6:5)

1. Ural'skiy filial Vsesoyuznogo obshchestva patofiziologov.
(Ural Mountain Region--Physiology, Pathological) (Medical societies)

BYKHOVSKAYA, A.M.; TOLSTOUKHOVA, L.I.

Treatment of thyrotoxicosis with radioactive iodine. Med. rad.
8 no.12:9-13 D '63. (MIRA 17:8)

1. Iz kafedry rentgenologii i radiologii (zav. - prof. Ye.L.
Kavesh) Kuybyshevskogo meditsinskogo instituta.

UZHANSKIY, Ya.G., professor, predsedatel'; TOLSTOUKHOVA, L.I., dotsent, sekretar'.

Second conference of the Ural Branch of the All-Union Society of Pathophysiologists. Arkh.pat. 15 no.3:88-90 My-Je '53. (MLRA 6:11)

1. Pravleniye Ural'skogo filiala Vsesoyuznogo obshchestva patofiziologov.
(Ural Mountain region--Pathology--Societies) (Societies--Pathology--
Ural Mountain region) (Ural Mountain region--Physiology--Societies)
(Societies--Physiology--Ural Mountain region)

cond
TOLSTOUKHOV, A. S.: Master Tech Sci (diss) -- "Representation of relief of a
lowland plain on topographic maps on a scale of 1:10,000 (On the example of
the delta of the river Terek)". Moscow, 1958. 15 pp (Min Higher Educ USSR,
Moscow Inst of Engineers of Geodesy, Aerial Photography, and Cartography),
150 copies (KL, No 2, 1959, 123)

TOISTOUKHOV, A.S.

Editing tasks in the representation of flatland reliefs on
1:10,000 scale topographic maps. Geod.i kart. no.9:61-67 S '57.
(MIRA 10:11)

(Relief maps) (Topographical drawing)

"APPROVED FOR RELEASE: 07/16/2001 CIA-RDP86-00513R001756120011-7

APPROVED FOR RELEASE: 07/16/2001 CIA-RDP86-00513R001756120011-7"

TOLSTOUSOV, V. P., Cand Agric Sci (diss) -- "The soils of the Yelan'-Kolemovskiy sugar-beet sovkhov, Voronezh Oblast: their genesis, properties, and ways of increasing their fertility". Voronezh, 1960. 22 pp (Min Agric RSFSR, Voronezh Agric Inst), 150 copies (KL, No 11, 1960, 136)

TOLSTOUSOV, V.P.

Ways of increasing the fertility of brown Solonetz-type spots
of the Chernozem complex. Pochvovedenie no.10:102-107 '60.
(MIRA 13:10)

1. Voronezhskiy sel'skokhozyaystvennyy institut.
(Solonetz soils) (Soil fertility)

USSR/Soil Science - Genesis and Geography of Soils. J

Abs Jour : Ref Zhur Biol., No 22, 1958, 99985

Author : Tolstousov, V.P.

Inst : Voronezh Agricultural Institute

Title : Characteristics of the Soil Cover of Yelan'-Kolenov
Sugar Beet State Farm of the Voronezhskaya Oblast' in
Connection with Topography of the Area.

Orig Pub : Zap. Voronezhsk. s.-kh. in-ta, 1957, 27, No 2, 339-342

Abstract : Yelan'-Kolinov Sugar Beet State Farm is situated in the
eastern part of the Voronezhskaya Oblast'. Here soil-
formation rocks exhibit a considerable influence on the
character of soil formation. Among the former predomi-
nate the parti-colored soft rocks, usually the salt-
containing sea deposits of the Tertiary Age. The soil
cover of the described territory, a plateau with a

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- 16 -

TOLSTOUSOVA, Z. A.

A. A. KOPEV, USSR 69,941, Dec. 31, 1947

TOLSTOUSOVA, Z. A.

23348 Tsellyuloznyy Nesmyvayemyy Appret L i Ego Primeneniye. Tekstil, Prom-St',
1949, No. 6, c. 23-25.