

RADCHENKO, S.I.; PETROV, A.A.

Alkylthioenynes and their analogs. Part 7: Synthesis and properties of alkylthioisopropenylacetylenes and their analogs. Zhur. org. khim. 1 no. 12:2115-2118 D '65 (MIRA 19:1)

1. Leningradskiy tekhnologicheskiy institut im. I. Lenooveta.
Submitted November 21, 1964.

SECRET, 1971

Classification of this document is "Secret" because it contains information the disclosure of which would be injurious to the national defense.

PELTON, G. J.; KEMM, V. L.; BERRY, S. R. V. J. Polym. Sci. A-1

Diene-styrene copolymerization. I. Kinetics and mechanism of the

radical copolymerization of styrene and diene monomers. Submitted May 1968.

"APPROVED FOR RELEASE: 07/19/2001

CIA-RDP86-00513R001240410015-3

APPROVED FOR RELEASE: 07/19/2001

CIA-RDP86-00513R001240410015-3"

PETROV, A.A.; RALL', K.B.; VIL'DAVSKAYA, A.I.

Synthesis and properties of 1,3-nitroalkadienes. Zhur.org.khim. 1
no.2:240-243 P 165. (MIRA 18:4)

1. Leningradskiy tekhnologicheskii institut imeni Lensoвета.

CHISTOKLETOV, V.N.; VAGINA, L.K.; PETROV, A.A.

Bipolar addition to unsaturated compounds. Part 10: Addition
of α, β -diphenylnitrene to vinylacetylene compounds. Zhur.org.
khim. 1 no.2:369-375 F '65. (MIRA 18:4)

1. Leningradskiy tekhnologicheskii institut imeni Lensovs'ya.

M. J. ... K. ... M. J. ...
... ..
... ..

PETROV, A. A.

SECRET

L 16648-65 EWT(d)/EWT(m)/EWP(w)/EWP(v)/EWP(k)/EWA(h) Pf-4/Peb
AEDC(a)/ASD(a)-5 EM

ACCESSION NR: AP4045712 S/0208/64/004/005/0880/0895 7

AUTHOR: Patrov, A. A. (Moscow); Popov, Yu. P. (Moscow); Pukhnachev, Yu. V. (Moscow)

TITLE: Calculation of natural oscillations of liquid in fixed containers by the variational method 26

SOURCE: Zhurnal vyshisitel'noy matematiki i matematicheskoy fiziki, v. 4, no. 5, 1964, 880-895

TOPIC TAGS: natural oscillations, Ritz method, eigenvalue problem, boundary value problem, Laplace equation, liquid oscillation, coordinate function

ABSTRACT: A method is presented for calculating the natural oscillations of an ideal fluid in fixed containers for a wide class of domains τ . (τ is the volume of the liquid in equilibrium.) The solution of this problem is reduced to the solution of the variational problem which consists in determining the function ϕ minimizing a certain functional $F(\phi)$. The use of Ritz method to

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ACCESSION NR: AP4045712

solve this variational problem makes it possible to determine the minimizing function ϕ in the form

$$\phi = \sum_{n=1}^N a_n f_n$$

where the coefficients a_n are determined from a certain homogeneous system of equations providing that the system of coordinate functions (f_n) in the domain τ is known. The method of constructing (f_n) is presented based on simple domains (parallelepiped, right cylinder, and others) enveloping the domain τ for which solutions of the problem are known. After the simple enveloping domain with a form closest to the form of the domain τ is chosen, the problem of determining the natural oscillations of the liquid is reduced to the evaluation of certain integrals and the solution of the system of homogeneous equations. The evaluations of the integrals and the solution of the system were carried out on electronic computers. The numerical solution of the problem on natural oscillations of liquids in containers is presented in detail for container in the shape of a cylinder with a horizontal generatrix, of a right cylinder with a spherical bottom and a spherical upper end cover, and of a torus. Orig. art. has: 19 formulas and 15 figures.

Card 2/3

L 16648-65

ACCESSION NR: AP6045712

ASSOCIATION: none

SUBMITTED: 15Oct63

ENCL: 00

SUB CODE: MA

NO REF SOV: 005

OTHER: 001

Fuel tanks

Card 3

1. The first part of the document is a list of names and titles of the members of the committee.

2. The second part of the document is a list of the names of the members of the committee.

3. The third part of the document is a list of the names of the members of the committee.

CHEKASOVA, I.A.; BAIYAN, Y.I.; ...

Reaction of ...
Telomerization of olefins with piperylene ...
ob. khim. 34 no.9:2912-2915, 1961.

Reactions of unsaturated ...
Addition of piperylene ...
tylene. 111:1242-43

Leningradskiy ...

SECRET

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SECRET

CONFIDENTIAL

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RAZIMOVA, N.A.; PETROV, A.A.

Preparation - containing catecholamines. Part of: *Preparation of*
catechol; osiferous and chloride with plane of *Preparation of*
ob.khim. 33 n. 12: 3862-3860 D 163.

1. Leningradskiy nauchno issledovatel'skiy institut khimicheskoy fiziki.

VALIACEV, H.G.; PETROV, A.A.

Testing the desalting process of Badayevskaya heavy crude. Study
Giprovostoknefti no.4:111-136 Vol. (MIRA 1:18)
(Kuybyshev Province - Petroleum desalting)

PETROV, A.A.; VERETENNIKOVA, I.V.

Reagents-demulsifiers for preparing petroleum for field of
Kashpir shale tar. Trudy Siprovoostoknefti no.4:137-160 '61.
(MIRA 16:8)
(Sodium sulfates)

PETROV, A.A.; SABO, L.V.

Theory and laboratory investigations of petroleum desalting
processes involving the use of rotary-disc contactors. Trudy
Giprovoztoknefti no.4:7'-9' '61. (MIRA 1:3)
(Volga-Ural region--Petroleum desalting)

PETROV, A.A.; PORFIR'YEVA, Yu.I.; SOBOL'EV, L.B.

Course of the reactions in which electrophilic and nucleophilic reagents are added to unsymmetrical monomers of diacetylene. Dokl. AN SSSR 161 n. 4:19, 1970. 2p. (MIRA 1971)

1. Leningradskiy tekhnologicheskii institut im. Lensoвета.
Predstavlen akademikom B.A. Astuzovym.

LETROY, A.A. ; 10307, U.S.S.R. ; 1964, U.S.S.R. (Moscow)

"An analysis of free oscillations of a diatomic molecule in a potential well by the variational method"

Report presented at the 2nd All-Union Conference on Theoretical and Applied Mechanics, Moscow 21 Jan - 1 Feb 64.

MOISEYEV, N.N. (Moscow); MYSHKIN, A.I. (Khar'kov); PERELI, A.A. (Moscow)

"On some new problems of the theory of motion of a body with liquid inside".

report presented at the 2nd All-Union Scientific Conference on Applied Mechanics, Moscow, 29 Jan - 1 Feb 61.

SABO, L.V.; PETROV, A.A.; KATYEV, V.M.

Testing the experimental petroleum desalting pilot plant combined with the rotary-disc contactor in the Novokuybyshev Petroleum Refinery. Trudy Siprovestoknefti no.4:100-110 '61. (MIRA 1:3)
(Volga-Ural region--Petroleum desalting)

BAKAYEV, M.T.; PETROV, A.A.

Seismic effect of underground blasting in Dzhezkazgan mines.

Trudy Inst. gor. dela AN Kazakh. SSSR 10:137-147 '67.

(MIRA 16:8)

(Dzhezkazgan District--Blasting) (Seismometry)

PETROV, A.A.; YAKOVLEVA, T.V.; KUPIN, B.S.

2,3,5-Trimethyl-1-hexen-3-yne. Zhur. ob. khim. 33 no.5:1701-
1702 My '63. (MIRA 16:6)

1. Leningradskiy tekhnologicheskii institut imeni Lensoвета.
(Hexenyne)

BOGOLYUBOV, G.M.; RAZUMOVA, N.A.; PETROV, A.A.

Synthesis of phospholine and phospholine, phosphorus-containing heterocycles. *Zhur.Obkhit.* 33 no.7:2419-2420 1968. (MIRA 16:8)

1. Leningradskiy tekhnologicheskij institut imeni Leningeta.
(Phospholine)

PETROV, A.A.

Hydraulic transmission system of a 4,000 hp. diesel locomotive.
Elek. i tepl. tiaga no.11:46-43 N 152. (MIRA 1971)
(Diesel locomotives--Transmission devices)

PETROV, A.A.

3/079/63/033/003/002/005
A066/A126

AUTHORS: Razumova, N.A., Petrov, A.A.

TITLE: Investigations in the field of conjugate systems. CLXIX.
Addition of dialkyl phosphorous acid and chlorides to diene hydrocarbons

PERIODICAL: Zhurnal obshchey khimii, v. 33, no. 3, 1963, 783 - 789

TEXT: The authors investigated the addition of chlorides of ethylene glycol phosphorous acid and propylene glycol phosphorous acid to divinyl, isoprene, piperylene, and chloroprene. The reactions were carried out by heating in sealed tubes. The resultant addition compounds could be sublimated at 1 - 2°C. A new kind of regrouping was obtained by the formation of five-membered phosphorous-containing heterocycles with a structure similar to that of the heterocycles formed by the reaction of diene hydrocarbons with alkyl dichlorophosphines. A number of derivatives of 3-phospholine-1 oxide is described for the first time. There are 2 figures and 2 tables.

Card 1/2

Investigations in the field of conjugate...

S/079/63/033/003/002/005
A066/A126

ASSOCIATION: Leningradskiy tekhnologicheskii institut imeni Lensoveta
(Leningrad Technological Institute im. Lensovet)

SUBMITTED: February 26, 1962

Card 2/2

PETROV, A.A.; PORFIR'YEVA, Yu.I.

Conjugated systems. Part 168: Course of the addition of
hydrogen halides to diene hydrocarbons - vinylallylacetylene
and its homologs. Zhur.ob.khim. 33 no.2:419-427 F '63.
(MIRA 16:2)

1. Leningradskiy tekhnologicheskij institut imeni Leningra.
(Hydrogen halides) (Butene) (Conjugation (Chemistry))

S/054/63/004/001/007/022
B102/B186

AUTHORS: Zaydel', A. N., Lazeyeva, G. S., Petrov, A. A.

TITLE: On the possibility of spectral-isotope determination of oxygen in metals

PERIODICAL: Leningrad. Universitet. Vestnik. Seriya fiziki i khimii, no. 1, 1963, 55-58

TEXT: The method of isotope spectral analysis for determining oxygen in metals is based on the spectroscopic determination of the isotope concentration ratio by a comparison of the band edges of the molecules $C^{12}O^{16}$ and $C^{12}O^{18}$. The 4123.6 \AA band of the CO molecule with an isotopic shift of $+8.5 \text{ \AA}$ is chosen as analytical band. The spectral width of the instrument slit was 1.7 \AA so that the band was not resolved. The relation $I^{18}/I^{16} = f(C^{18}/C^{16})$ is represented by a linear 45° calibration curve which does not intersect with the origin but is somewhat shifted to lower intensity ratios. The curve was plotted for the O^{18} concentration range 5 - 63%. The accuracy in oxygen determination was 2-3% in the case

Card 1/2

S/054/63/004/001/007/022
B102/B186

On the possibility of ...

of standard analytical conditions. The applicability of the method was demonstrated on the analysis of specially prepared powders of Armco iron and cobalt. The reproducibility of the results was 5%. The sensitivity of the method is limited by the correction of idle experiment which has to be reduced by a factor of 5-10 for an accuracy of $10^{-4}\%$. There are 3 figures and 2 tables.

SUBMITTED: July 23, 1962

Card 2/2

BUNINA-KRIVORUKOVA, L.I.; PETROV, A.A.

Interaction of bivinyl and isoprene hydrochlorides with sodium acetoacetic ester. Zhur.ob.khim. 33 no.2:427-432 F '63.

(MIRA 16:2)

1. Leningradskiy tekhnologicheskij institut imeni Lensoveta.
(Butadiene) (Isoprene) (Acetoacetic acid)

PETROV, A.A.; LEEDEEV, V.B.; PORFIR'YEVA, Yu.I.

Conjugated systems. Part 167: Nuclear magnetic resonance spectra
and structure of dienyne hydrocarbons. Zhur.ob.khim. 33
no.2:416-418 F '63. (MIRA 1642)

1. Leningradskiy tekhnologicheskii institut imeni Lensoveta.
(Hydrocarbons—Spectra)
(Nuclear magnetic resonance and relaxation)

BALAYEV, G.A.; PETROV, A.A.

Chemistry of organic oxides. Part 22: Addition of hydroxylamine
to α -oxides. Izv.vys.uch.zav.; khim.i khim.tekh. 5 no.4:608-611
'62. (MIRA 15:12)

1. Leningradskiy tekhnologicheskiy institut imeni Lensoveta,
kafedra organicheskoy khimii.
(Hydroxylamine)
(Oxides)

IONIN, B.I.; PETROV, A.A.

Prototropic isomerization of esters of alkenylphosphinic acids.
Zhur.ob.khim. 33 no.2:432-437 F '63. (MIRA 16:2)

1. Leningradskiy tekhnologicheskij institut imeni Lensoveta.
(Phosphinic acid) (Isomerization)

RAZUMOVA, N.A.; PETROV, A.A.

Conjugated systems. Part 169: Addition of dialkyl phosphoryl
chlorides to diene hydrocarbons. Zhur.ob.khim. 33 no.3:783-789
Mr '63. (MIRA 1643)

1. Leningradskiy tekhnologicheskii institut imeni Lensoveta.
(Phosphoryl chloride)
(Hydrocarbons)

CHISTOKLETOV, V.N.; TROSHCHENKO, A.T.; PETROV, A.A.

Addition of benzonitrile oxide to unsaturated compounds.
Part 3: Interaction of benzonitrile oxide with vinylacetylenes
substituted in a vinyl group in β -position. Zhur.ob.khim.
33 no.3:789-793 Mr '63. (MIRA 16:3)

1. Leningradskiy tekhnologicheskii institut imeni Lensovetu.
(Benzonitrile) (Butenyne)

PETROV, A. A.; YELSAKOV, N. V.

Effect of solvents on the nuclear magnetic resonance spectra
of diacetylene and vinylacetylene. Zhur. ob. khim. 33 no. 11
319-320 1963. (MIRA 16.1)

1. Leningradskiy tekhnologicheskij institut imeni Lensoveta.

(Butadiyne--Spectra) (Butenyne--Spectra)
(Solvents)

L 11060-63

EWP(j)/EFF(c)/EWT(m)/BDS--Pc-4/Pr-4--RM/WW/JFW

ACCESSION NR: AP3000482

S/0153/63/006/001/0170/0171

AUTHOR: Kheruze, Yu. I.; Petrov, A. A.TITLE: Addition of triphenylmethyl radicals to divinylacetylene and its homologs

SOURCE: Izv. VUZ: Khimiya i khim. tekhnologiya, v. 6, no. 1, 1963, 170-171

TOPIC TAGS: triphenylmethyl radicals, divinylacetylene, vinylpropenylacetylene, vinylisopropenylacetylene, addition reactions

ABSTRACT: Crystalline adducts were formed between the triphenylmethyl radical and divinylacetylene (1), vinylpropenylacetylene (2), and vinylisopropenylacetylene (3) in benzene solution. Addition occurred chiefly in the 1,4-positions to form 7, 7, 7-triphenyl-3-triphenyl-methyl-heptatriene, 8, 8, 8-triphenyl-3-triphenylmethyl-heptatriene-1,3,4. Adducts were characterized by their IR spectra. Weak absorption occurs. Adducts were recrystallized from propylacetate and formed molecular compounds with acetone. Orig. art. has: 1 table.

ASSOCIATION: Kafedra organicheskoy khimii, Leningradskiy tekhnologicheskii institut im. Lensoveta (Department of Organic Chemistry, Leningrad Technological Institute)

Card 1/1

Perlov, A.A.

10

S/169/63/000/003/006/042
D263/D307

AUTHORS: Alekseyev, P.P., Besyadovskiy, Ye..., Biryukova, L.A.,
Golyshev, G.I., Ivanovskiy, A.I., Izakov, I.I.,
Kokin, G.A., Kurilova, Yu.V., Livshits, L.S., ~~Perlov,~~
~~A.A.~~ Rozhdestvenskiy, B.G., Solov'yev, N.V., Speran-
skiy, K.Ye., Khvostikov, I.A., Shvidkovskiy, Ye.G.
and Shcherba, I.A.

TITLE: Study of the upper layers of the atmosphere with the
aid of meteorological rockets

PERIODICAL: Referativnyy zhurnal, Geofizika, no. 3, 1963, 28,
abstract 3.1166 (Tr. Vses. nauchn. meteorol. sovesh-
chaniya. T.I.L., Gidrometeoizdat, 1962, 91-103)

TEXT: In the present review-type article the authors give
the results of studies carried out at Tsentralnaya aerologicheskaya
observatoriya (Central Aerological Observatory) on atmospheric sound-
ing with meteorological rockets. Measuring methods are described and
the main points are given for obtaining such atmospheric character-

Card 1/2

S/169/63/000/003/006/042
D263/D307

Study of the upper layers ...

istics as pressure, temperature, and wind. Certain results are given: data of seasonal temperature variations at heights up to 50 km in the middle latitudes of the USSR and in polar regions, cases of sudden warming up, characterization of temperature distribution curves, a table characterizing the temperature inversion below the stratopause under the conditions of polar night, and data regarding the circulation in the upper atmospheric layers. Information is given on the constructed meridional sections of temperature fields and on the zonal component of the gradient wind. (25 references).
[Abstracter's note: Complete translation]

Card 2/2

ORLOVA, N.M., 1970, A.S.

Hydrogen determination of the amount of the ^{137}Cs in the soil by the
balancing method. Dokl. Akad. Nauk SSSR, 1970, 203, 104. (MIRA, 1970)

YELBAKOV, S.V.; BERTOV, A.A.

Studying H-bonds, formed by acetylene hydrocarbons, with the aid of
nuclear magnetic resonance spectroscopy. Opt. i spekt. 16: 201
148-151 Jan 1964. MIRA 1963

1A 197710

USSR/Chemistry - Petroleum, Isomer-
ization Nov/Dec 51

"Isomerizations of Olefinic Hydrocarbons Over
Aluminosilicates," A. A. Petrov, A. V. Frost, M. I.
Babuyev, Petroleum Inst, Acad Sci USSR

"Iz Ak Nauk SSSR, Otdel Khim Nauk" No 6, pp 745-
752

Investigated catalytic isomerization on aluminosilicate catalyst of a number of olefinic hydrocarbons under conditions excluding cracking. Proposes mechanism of isomerization involving splitting off of olefin from catalyst under formation

197710

USSR/Chemistry .. Petroleum, Isomer-
ization (Contd) Nov/Dec 51

of intermediate alkyl cyclopropane product. These cyclopropane derivs then undergo ring opening through fission of the most highly asym bond. In the unsatd products, the double bond is usually next to the side alkyl group.

197710

... A. A.

USSR/Chemistry - Liquid Fuels

11 Jun 51

"Conversion of 1-Methylcyclopentane-1 Over an Active Aluminum Sulfate Catalyst" A. A. Petrov, V. V. Sheberin, Inst. Petroleum, Acad. Sci. USSR

"Dok Ak Nauk SSSR" Vol. LXXVIII, No. 1, pp. 41-915

Monomer fractions of product obtained by conversion at 2500 do not contain 6-membered napthalenes. Polymer product resulting from conversion of 1-methylcyclopentane-1 is cyclohexylmethylcyclopentane, substance identical with that formed from cyclohexane under the same conditions. Apparently ring expansion of simple pentamethylenes proceeds only

184715

USSR/Chemistry - Liquid Fuels
(Contd)

11 Jun 51

simultaneously with polymerization. Dehydrogenation over this catalyst of 6-membered rings into aromatics also seems to occur together with polymerization only.

184715

PETROV, A. A.

USSR.

4

✓Catalytic transformation of olefins over aluminum silicates. A. A. PETROV. *Trudy Vsesoyuz. Sovetskoye na Khim. i Tsellyul. Inst. (Sept. 18-23, 1951)* (Izdatel. Akad. Nauk Azerbaidzhan, S.S.R., Baku) 1953, 141-50; *Referat. Zhur., Khim.* 1954, No. 20650.—A discussion of catalytic transformations of olefins which practically preclude cracking. M. Hosen

AD per

Chemical Abst.
Vol. 48 No. 8
Apr. 25, 1954
Organic Chemistry

2
Redistribution of hydrogen in olefinic hydrocarbons on an
aluminosilicate catalyst A. A. Petrov and A. V. Ergot
Zh. Fiz. Khim. U.S.S.R. 22, 1813 (1948); Eng. Transla-
tion: See C. I. 47 8336 H. I. H.

10-5-54
JHP

PETROV, A. A.

3

Dependence between the amount of liberated hydrogen bromide and structure of olefins undergoing bromination. Al. A. Petrov (Petroleum Inst., Acad. Sci. U.S.S.R., Moscow). *Zhur. Obshchei Khim.*, 23, 1806-0 (1953).—Tests with 49 olefins with different structures are reported in the reaction of these with 1% Br soln. in MeOH satd. with NaBr. The hydrocarbons were treated with excess reagent and the excess Br was back-titrated with $\text{Na}_2\text{S}_2\text{O}_3$ after addn. of KI. Usually 1 min. contact was sufficient for the reaction. The colorless soln. was treated with 6% NaOI and the liberated iodine was again titrated as usual. It was shown that the amount of liberated HBr depends on the structure of the olefin. Linear olefins and those with a side chain that is not proximate to the double bond eliminate some 14-16 mole-% HBr. Olefins with branching at the double bond give 27-39% HBr, while those with branching at the double bond and in addn. having bulky groups at the double bond (iso-Pr, Me₂C) give large amts. of HBr (41-75%). The course of the reaction may be represented by the following scheme: the MeOH-Br₂ complex yields MeOH and the olefin dibromide (straight chain olefins), or a carbonium ion after addn. of pos. Br ion, which is then attacked by MeOHBr⁻, yielding HBr and a methoxy-bromoalkane (for branched olefins). Both courses are affected by steric factors. The reaction of 2,5-dimethyl-2-hexene gave a colored product, n_D^{20} 1.4860, d_4^{20} 1.3300, which contained 6% MeO and 46.6% Br, thus confirming the above hypothesis. G. M. Kosolapoff

MA
gaw

The order of addition of bromine to homologs of vinylacetylene. A. A. Petrov and Yu. I. Porfir'eva. *Doklady Akad. Nauk S.S.S.R.* 89, 873 (1953); *cf. CA* 44, 7751c. To 4.3 g. MeC CCH=CH₂ in 200 ml. CHCl₃ at -8° was added 53 g. Br in 100 ml. CHCl₃, distn gave a no. of fractions up to 145°/10 mm. of which the most abundant were those b.p. 70.5-80.5° (17.5 g.) and b.p. 132.6° (13 g.). The former, d₄ 1.403, n_D 1.5588, was identified as MeC CCHBrCH₂Br (I) on the basis of the following reactions, and the latter, d₄ 1.403, n_D 1.5588, was identified as MeC CCHBrCH₂Br (II) on the basis of the following reactions, and the Raman spectrum of the frequency 2240 cm⁻¹ which is that of a triple link. I (7.3 g.) at room temp. with 2.7 g. KOH in 25 ml. EtOH gave in 4 hrs. 60% bromide ion, while the soln. yielded 60% bromopentenyne. Heating I with concd. HBr-Cu₂Br₂ in Et₂O 4 hrs. gave some 80% stable dibromide, MeCBr:CBBr:CH:CH₂, b.p. 57-7.5°, d₄ 1.8305, n_D 1.5768, which in alc. KOH lost but 5% of its Br. under the above conditions. Ozonolysis of I and treatment with H₂O₂ gave AcOH and BrCH₂CHBrCO₂H, m. 64-6°. With KMnO₄, I gave the same products. The high-boiling fraction (b.p. 132-6°) was impure tetrabromide, d₄ 2.3406, n_D 1.6160. Similar reaction of 40 g. EtC:CCH=CH₂ with Br in CHCl₃ gave 52 g. mixed di- and polybromides which distd. up to 190°/10 mm. From these was isolated 24 g. material, b.p. 86-91°, which after redistn. gave pure EtC CCHBrCH₂Br, b.p. 87.7.5°, d₄ 1.6920, n_D 1.5470, whose chem. properties were identical with those of the Me analog. Along with this dibromide was obtained 10 g. crude tetrabromide, b.p. 140-60°, d₄ 2.2035, n_D 1.6020. Similar bromination of 65 g. HC:CCMe:CH₂ in CHCl₃ over 9 hrs. gave a range of fractions of which the principal one (40 g.), b.p. 44-53°, was redistd., yielding pure HC:CCMeBr:CH₂Br (II), b.p. 50.1.5°, d₄ 1.7581, n_D 1.5320, Raman frequency 2118 cm⁻¹. II with ammoniacal AgNO₃ gave a white ppt. of the Ag salt, but this could not be reversed by ahdn. of HCl. With alc. KOH II gave 60% 4-bromo-3-methyl-3-buten-1-yne, b.p. 35.

9°, d₄ 1.3690, n_D 1.5070. II heated with HBr-Cu₂Br₂ gave CHBr:CB:CMc:CH₂, b.p. 48.5-9.0°, d₄ 1.7740, n_D 1.5430, which was quite stable to alc. KOH, losing but 4% of its Br after 4 hrs. at room temp. In addn. to II, dibromide the original reaction mixt. gave 2.3 g. product, b.p. 70-80°, probably CHBr:C:CMcCH₂Br, which, heated with HBr-Cu₂Br₂, gave the above dienic dibromide, b.p. 48.9°. An unstated amt. of impure tetrabromide, b.p. 140-45°, was also obtained, but this decomp. during distn. MeC CCBBrCH₂, b.p. 52.3°, d₄ 1.4010, n_D 1.5255. EtC CCBBrCH₂, b.p. 1.3531, n_D 1.5156. G. M. Kosolapoff

PETROV, A.A.; ARBUZOV, A.E., akademik.; PORFIR'YEVA, Yu.I.

Order of addition of alkylhypohalogenides to vinylacetylene homologs.
Dokl.AN SSSR 90 no.4:561-564 Je '53. (MLRA 6:5)

1. Akademiya Nauk SSSR (for Arbuzov). (Halogenides) (Vinylacetylene)

PETROV, A. A.

USSR/Chemistry - Fuels

11 Sep 53

"Isomerization of Unsaturated Hydrocarbons Having
the Composition $C_{12}-C_{16}$." A. A. Petrov, Inst of
Petroleum, Acad Sci USSR

DAN SSSR, Vol 92, No 2, pp 329-33.

Studied the isomerization of dodecene-1, tridecene-1,
and hexadecene-1 using Al_2O_3 and HCl as catalyst.
Aluminosilicate catalyst was used for the isomeriza-
tion of hexadecene-1. Did not detm the structure of
the isomers. Presented by Acad A. V. Topchiyev
14 Jul 53.

269T17

PEKROV, AL. A.

U.S.S.R.

Isomeric transformations of olefin hydrocarbons over
 aluminum silicate. II. Al. A. Pekrov, *Izv. Akad. Nauk
 S.S.R., Otdel. Khim. Nauk* 1954, 124-32; *Bull. Acad.
 Sci. U.S.S.R., Div. Chem. Sci.* 1954, 101-7 (English trans-
 lation); *et. C.A.* 46, 4776c. Olefin and cycloolefins under-
 go considerable changes over aluminum silicate catalyst at
 about 260°. The most characteristic changes are: (1)
 isomerization of *n*-alkenes to monoalkylalkenes with an
 alkyl side chain at the site of the double bond; (2) forma-
 tion of secondary C atom groupings, except for cases in which
 the double bond is proximate to a tertiary C atom at a double bond;
 (3) isomerization of cycloolefins of the C₇ type and higher
 (cycloheptenes); (4) shift of the double bond to form unsym-
 metrical olefins; (5) decrease of the no. of C atoms in side
 alkyl chains. Thus, the following changes were observed:
 1-methyl-1-hexene gave Me₂CHPr and (CHMe)₂h; 6-
 methyl-1-hexene gave Me₂CHBu, CH₂(CHMe)₂, a small
 amt. of BiCHMeCHMe; 6-methyl-1-heptene gave Am-
 diethylhexanes; 1-heptene gave BuCHMe₂, PrCHMeEt,
 and a small amt. of CH₂(CHMe)₂; Me₂CCH₂CH₂CH₂ gave

CH

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HI. H. PETROV

EtCHMeCHMe₂ and BuCHMe₂; Me₂CHCMc:CH₂ gave Me₂CHCMc₂ and a small amt. of CH₂(CHMe₂); (CH₂:CH:CH₂)₂ gave PrCHMe₂ and (CHMe₂)₂; Et₂C:CHMe gave Me₂CHBu, EtCHMePr, and CH₂(CHMe₂); PrCH:CM₂ gave BuCHMe₂, PrCHMeEt and CH₂(CHMe₂). In this work the attention was largely directed to the satd. products; the unsatd. products were shown to have the same configuration as the satd. analogs listed above. At 260° the following products formed: from 1-ethylcyclopentene, 45% 5-membered cyclanes and 55% methylcyclohexane; from 1-ethylcyclohexene, 35% 5-membered cyclanes and 65% 1,4-dimethylcyclohexane; from 1-propylcyclohexene, 72% 6-membered and 28% 5-membered cyclanes; from allylcyclohexane, 64% 6- and 36% 5-membered cyclanes; from allylbenzene, alkylbenzene. At 400° 1-ethylcyclopentene gave 27% methylcyclohexane, 8% MePh, 65% 5-membered cyclanes (largely 1,3-dimethylcyclopentane); 1-ethylcyclohexene similarly gave 42% 5-membered cyclanes, 30% aromatics (xylenes), and 38% 6-membered cyclanes (1,4- and a little 1,3-dimethylcyclohexanes). The schemes of formation of the products are discussed.

2/3

G. M. Korolapoff

PETROV, A.A.

Effect of sulfuric compounds on catalytic functions of aluminosilicates
Trudy Inst. nefti no. 6: 86-88 196. (MIRA 8:12)
(Catalysis) (Aluminum silicates) (Sulfur compounds)

Translation D-411569

PETROV, A. A.

~~Catalytic properties of activated aluminum oxide, K. G. Mischenko, O. B. Morozova, and A. A. Petrov. J. Gen. Chem. U.S.S.R. 23, 2166-8 (1966) (Engl. translation). See C.A. 50, 4611d.~~

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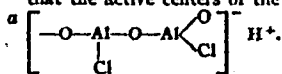
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PETROV, AL. A.

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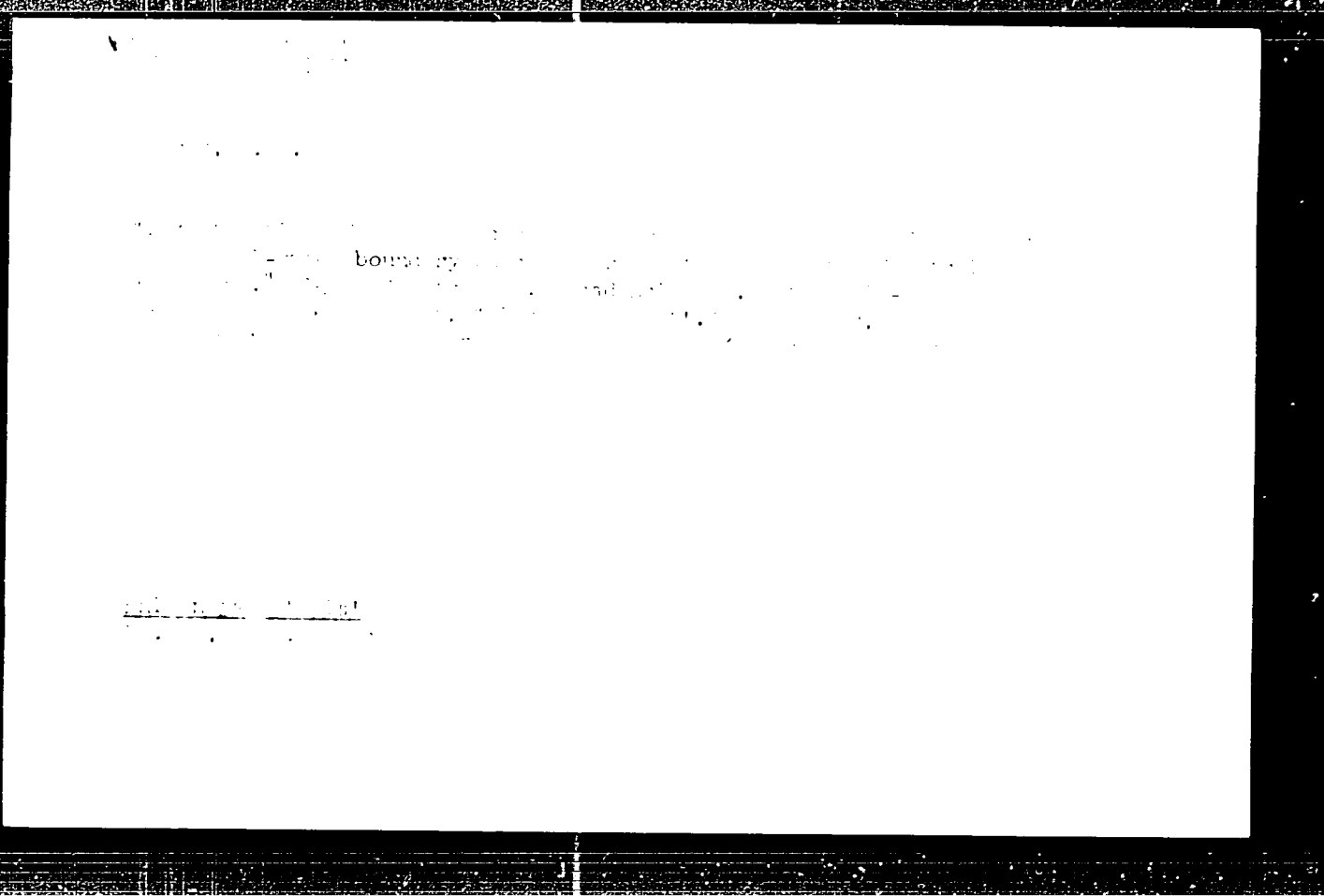
\checkmark Catalytic properties of activated aluminum oxide. K. G. Messerov, O. E. Ngorozova, and Al. A. Petrov. *Zhur. Obshchei Khim.* 25, 2204-8(1955). -- The catalytic properties of Al_2O_3 , activated with mineral acids in the isomerization of olefins were studied. The activating actions of the acids increase in the following order: H_3BO_3 , HBr , H_3PO_4 , HCl . HF , H_2SO_4 , HNO_3 , has no effect. The activity of Al_2O_3 activated with HCl is directly related to the extent to which the surface OH groups are replaced by Cl^- . It is proposed that the active centers of the catalyst have the structure



J. Rovtar Leach

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Petrov, A. A.

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✓ 923. BROMINATION OF UNSATURATED HYDROCARBONS. Petrov, A. A.
(Khim. Tekhnol. Topilva (Chem. Technol. Fuel. Moscow), July 1956, 32).
The determination of unsaturated hydrocarbons with a solution of bromine in methanol is examined in the light of results with 120 compounds. A large group of compounds, containing some olefins and five-membered cycles, gave a higher bromine number than the calculated one. This phenomenon is explained. Bromination is accompanied by separation of hydrogen bromide. A method is proposed for determining the number of branches in alkenes and cycloenes from the results of bromination. (L).

in copy

PETROV, A.I.A.; SERGIYENKO, S.R.; TSEDILINA, A.L.; YEGOROV, Yu.P.

Isomerization of unsaturated C₁₂-C₁₆ hydrocarbons. Khim. i tekhn.
tepl. no.1:26-32 Ja '56. (MLBA 9:7)
(Hydrocarbons)

Petrov, A. A.

1

112

Bromination of unsaturated hydrocarbons. A. A. Petrov. *Khim. i Tekhnol. Topiss* No. 7, 24-27 (1955). Based on an analysis of the data from the quant. detn. of unsat. by bromination (0.1N Br in methanol acid. with NaBr), the compds. were subdivided into the following groups: I: ethylene, propylene, 1-butene, 1-hexene, 1-heptene, 1-octene, 1-nonene, 1-dodecene, 1-tridecene, 1-tetradecene, 1-pentadecene, 1-hexadecene, 3-heptene, 4-methyl-1-pentene, 6-hexadecene, 3-methyl-1-butene, 4-methyl-1-pentene, 5-methyl-1-hexene, 6-methyl-1-heptene, 2-methyl-1-pentadecene, 3,3-dimethyl-1-butene, 4,4-dimethyl-1-pentene, 4,4-dimethyl-1-hexene, cyclohexene, 3-methyl-1-cyclohexene, allylcyclohexane, cyclopentene, allylbenzene, 4-phenyl-1-butene, 1,3-diphenylpropene, 1,6-diphenyl-1-pentene, styrene, 1,2-phenylmethylene. II: isobutylene, 2-methyl-2-butene, 2-methyl-1-pentene, 2-methyl-2-pentene, 3-methyl-2-pentene, 2,3-dimethyl-1-butene, 2,3-dimethyl-2-butene, 2-methyl-2-hexene, 3-methyl-2-hexene, 3-ethyl-2-pentene, 2,3-dimethyl-2-pentene, 2,4-dimethyl-2-pentene, 2,3,3-trimethyl-1-butene, 2-methyl-2-heptene, 3-methyl-2-heptene, 4-methyl-3-heptene, 2-ethyl-2-hexene, 3-ethyl-2-hexene, 2,3-dimethyl-3-hexene, 3,4-dimethyl-4-hexene, 4, 6-methylarene, 4-propyl-3-heptene, 2-methyl-2-undecene, 3-methyl-2-undecene, 4-methyl-3-undecene, 5-methyl-4-undecene, 6-methyl-6-undecene, 2,3-dimethyl-4-decene, 2,5-dimethyl-4-decene, 2,5,8-trimethyl-4-nonene, 6-propyl-4-nonene, 6-isopropyl-4-nonene, 2-methyl-2-dodecene, 3-methyl-2-dodecene, 4-methyl-3-dodecene, 5-methyl-4-dodecene, 6-methyl-6-dodecene, 2-methyl-2-tridecene, 3-methyl-2-tridecene, 4-methyl-3-tridecene, 6-methyl-4-tridecene, 6-methyl-6-tridecene, 7-methyl-6-tridecene, 2,3-dimethyl-3-dodecene, 4-methyl-3-tetradecene, 6-methyl-4-tetradecene, 6-methyl-6-tetradecene, 7-methyl-

PETROV, A. A.

8-tetradecene, 4-methyl-3-pentadecene, 5-methyl-4-pentadecene, 6-methyl-5-pentadecene, 7-methyl-6-pentadecene, 8-methyl-7-pentadecene, 1-methyl-1-cyclopentene, 1-ethyl-1-cyclopentene, 1,3-dimethyl-1-cyclopentene, 1-propyl-1-cyclopentene, 1-butyl-1-cyclopentene, 1-heptyl-1-cyclopentene, 1-methyl-1-cyclohexene, 1-ethyl-1-cyclohexene, 1,2-dimethyl-1-cyclohexene, 1,3-dimethyl-1-cyclohexene, 1,4-dimethyl-1-cyclohexene, 1-propyl-1-cyclohexene, 1-butyl-1-cyclohexene, 1-hexyl-1-cyclohexene, 1,1-methyl-phenylethylene, 1,1-diphenylethylene. III: 2,4-dimethyl-3-hexene, 2,4,4-trimethyl-1-pentene, 2,2,3-trimethyl-3-pentene, 2,3,4-trimethyl-2-pentene, 2,2,3-trimethyl-3-hexene, 2,4-dimethyl-3-decene, triisobutylene (mixt.), 2,4-dimethyl-3-undecene, 8-isobutyl-1-undecene, 2,4-dimethyl-3-tridecene, 3,6-dimethyl-4-dodecene, 2,4-dimethyl-3-tridecene, 2,4-dimethyl-3-tetradecene, 1,2-dimethyl-1-cyclopentene, 1-isopropyl-1-cyclopentene, 1-isobutyl-cyclopentene, 1-isopropyl-1-cyclohexene, 1-isobutyl-1-cyclohexene, 1-phenyl-1-cyclopentene. The bromination coefficient of the compds. of group I varied on the average 12-14%, that of group II, 24-27%, and of group III, 50-59%. A branching in the side-chain of cycloolefins had a stronger effect on the increase of the coefficient, than the presence of the 2nd group in the ring. In the instance of aliphatic hydrocarbons the branching of the side-chain had approximately the same effect on the value of the coefficient as the presence of two substitutions in the chain. The amt. of HBr liberated was found to be directly related to the structure of a compd. This was utilized for developing a method for the detn. of the degree of branching of unsatd. hydrocarbons from the bromination data.

A. P. Kotloby

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USSR/Physical Chemistry - Molecule. Chemical Bond.

B-4

AbsJour : Referat Zhur - Khimiya, No 6, 25 March 1957, 18159

Author : Ygorov, Yu.P. and Petrov, Al.A.

Title : Determination of the Degree of Ramification of Paraffin Hydrocarbons of C₁₂ - C₁₆ Composition by the Method of Infrared Spectroscopy.

Orig Pub : Zh. analit. chemistry, 1956, 11, No 4, 483-488

Abstract : Infrared spectra of 28 alkanes (A) of normal structure and iso structure of C₁₂ - C₁₆ composition are examined in the region of valence C-H 3.35 - 3.5 μ in a solution of CCl₄. The character of change in intensity of peaks observed allows us to show the groups where they belong: CH₃ 3.38 μ , CH₂ 3.50 and 3.42 μ and CH 3.48 μ . As the number of groups CH₃ increases with the ramification of an alkane (while the number of groups CH₂ decreases), the measurement of the ratio of intensities of peaks 3.50 and 3.42 to 3.38 μ proves to be a measure of

Card 1/2

- 26 -

Retzov, A.A.

Distr: 4E38/4E65/4E2a(5)
Determination of the degree of branching of C₆-C₆
and C₆-C₆ by infrared spectroscopy. *J. Anal. Chem. U.S.S.R.* 11,
611-18 (1964) (English translation).—See *C.A.* 51, 16291k.
R.M.R.

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Petrov, A.A.

2

Equilibrium mixtures of five- and six-membered cyclohexanes. A. Petrov. Doklady Akad. Nauk S.S.S.R. 109, 649-61 (1960). The equil. concns. of cyclohexanes resulting at 250-400° over Al₂O₃ (activated with HCl) and synthetic aluminosilicate catalysts were detd. The concns. in vol. % of hexamethylene hydrocarbons in the catalyzates from the indicated materials are shown (on aluminosilicate at 250°, 300°, 400°, on Al₂O₃ at 300° and 400°, given in order): cyclohexane, under 5%, —, —, under 5%, —; 1-methylcyclopentene, under 5%, —, —, under 5%, —; 1-methylcyclohexene, 61, 89, 21, 18, 13; 1-ethylcyclopentene 55, 34, 27, 30, —; 1,2-dimethylcyclopentene, 20, —, —, —; 1,3-dimethylcyclopentene, 15, —, —, —; 1-ethylcyclohexane, 65, 89, 88, 20, —; 1,3-dimethylcyclohexene, 64, —, —, 21, —; 1-propylcyclohexene, 72, —, —, 24, —; allylcyclohexane, 64, —, —, —; 1-butylcyclopentene, 55, —, —, —; 1-hexylcyclohexene, —, —, —, 25, —; 1-heptylcyclopentene, —, —, —, 20, —. The Δ*E*° for isomerization of cyclohexene to methylcyclopentene is estd. at 8800-4000 cal/mole which detd. the equiv. concn. of cyclohexene as 2-4%. The probable mechanism of 5-6 ring isomerization is discussed. G. M. Kosolapoff

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PETROV, A. A. Doc Chem Sci -- (diss) "Study in the field of ~~XXXXXXXXXXXX~~
of hydrocarbons." Mos, 1947. 11 pp (Acad Sci USSR. Inst of Petroleum), 1:0
copies (KL, 42-17, 91)

Петрова А.А.

USSR Chemical Technology - Chemical Products and Their Application. Treatment of Natural Gases and Petroleum. Motor and Jet Fuel. Lubricants. I-6

Abs Jour : Ref Zhur - Khimiya, No. 1, 1968, 2438
Author : Basharat'yan, Ya.S., Petrov, A.A.
Inst : -
Title : Electric Dehydration Unit at the Kalinovskiy Fields.
Orig Pub : Neftyanik, 1967, No. 12, 17-18

Abstract : The unit at the Kalinovskiy fields of Kinel'neft' is used to dehydrate all of the petroleum production of the fields which has an average water content of 65-70%. To ensure the necessary output of the dehydrators the crude petroleum is subjected to a preliminary thermo-chemical treatment which decreases the water content, on the average, to 40-50%. The consumption of NCHK reagent amounts to 20 kg per ton of oil. Heating temperature of crude petroleum is 40-50°. The electric treatment constitutes the second

Card 1 of 1

AUTHORS: Kusakov, M. M., Petrov, A. A.

20-4-30/51

TITLE: Note on the Rheologic Properties of Surface Layers at the Oil-Water Interface and Their Role in the Stability of Oil Emulsions. Reologicheskiye svoystva i povorkhnostnykh sloev na granitse razdela neft'-voia i ikh rol' v ustoychivosti neftnykh emulsiy.

PERIODICAL: Doklady AN SSSR, 1951, Vol. 117, No. 4, pp. 237-240 USSR

ABSTRACT: The authors employed an apparatus of the type of a torsion pendulum for the purpose of studying these rheologic properties. According to the composition of the hydrocarbon phase and of the water phase, of the time of formation of the layer and its temperature three types of motion of the torsion pendulum can be distinguished: a) Damped oscillations, b) An aperiodic motion, c) An elastic-viscous motion. The most exhaustive data on the character of the rheologic properties of the surface layers can be obtained from the observation of the elastic-viscous motion of the torsion pendulum. A diagram illustrates the typical curves of the temporal development of the deformation of the surface layers at the interface between

Card 1/3

Note on the Rheologic Properties of Surface Layers at the Oil-Water Interface and Their Role in the Stability of Oil Emulsions 20-4-30/51

petroleum and water at a varying total content. The deformation of these surface layers develops in a way analogous to the deformation in the adsorption layers of saponine. Such types of curves of deformation over time can be caused by three types of deformations, instantaneous elastic deformations, deformation caused by elastic after-effects and an irreversible deformation of flow. Then the quantities characterizing the rheologic quantities are enumerated. For the purpose of investigation, the influence of the nature of the petroleum on the properties of the surface layers, the rheologic properties of the surface layers on the interface were examined with a 1% solution of NaCl in H₂O with different kinds of petroleum from Eastern wells oil. The results obtained from these investigations are compiled in a table. The surface layers of the various oil types at the interface between oil and water possess high elastic and viscous properties. The surface viscosity reaches values of the order of magnitude of 10⁴ surface Poise. The occurrence of a maximum strength

Card 1/3

20-4-30/51

Note on the Rheologic Properties of Surface Layers at the Oil-Water Interface and Their Role in the Stability of Oil Emulsions

in the surface layers at the boundary of the water in the range of diluted oil solutions is apparently connected with a more intensive formation of the substances of an asphaltous and resinous type in the surface layers. The investigation of the influence of the composition of the oil phase and the water phase, of temperature, of the time of formation of the layer and of other factors on the rheologic properties of the surface layers makes it possible to select suitable methods for the dehydration and desulfurization of the petroleum and the characterization of the parameters of the petroleum and an evaluation of the efficiency of the desulfurators. There are 4 figures, 1 table and 10 Soviet references

ASSOCIATION: **Petroleum Institute** AN USSR Institute of Oil Akademi nauk SSSR
PRESENTED: April 1, 1951, by A. V. Popov, Academician
SUBMITTED: February 28, 1951
AVAILABLE: Library of Congress
Card 4/3

PETROV, A.A.

Specific conductivity of oil emulsions. Trudy Giprovoostoknefti
no.1:352-372 '58. (MIRA 13:9,
(Emulsions--Electric properties)

ATTENTION: [Illegible text]

TITLE: [Illegible text]

DESCRIPTION: [Illegible text]

ABSTRACT: [Illegible text]

REFERENCES: [Illegible text]

Isomerization of Saturated Hydrocarbons. 2. Alkanes with Isomers.
Conversions of Alkanes with $C_6 - C_8$ Structure

number of new data were determined which offer new ideas
as to the kinetic connection, the structure and the
reactivity of hydrocarbons. There are 4 tables, and 11
references, 11 of which are Soviet.

ASSOCIATION: Institut نفتي Akademiya Nauk SSSR (Petroleum Institute,
AS USSR)

SUBMITTED: November 19, 1959

AVAILABLE: Library of Congress

1. Hydrocarbons--Saturated--Isomerization 2. Alkanes
 $C_6 - C_8$ --Isomeric conversions

Card 3/3

AUTHORS:

1. IROV, A. A., 2. TSELIINA, A. L., 3. PETROVA, I. I.

TITLE:

Isomerization of saturated hydrocarbons (isomerizatsiya nasyschennykh uslevodorodnykh soedyneniy) 2. Isomerizations of the Alkanes of the C_9-C_{16} -structure (isomerizatsiya 2. Izomernyye prevrascheniya alkany sostava C_9-C_{16})

PERIODICAL:

Izvestiya Akademii Nauk SSSR, Otdel'nyye Annalen Khimii, 1958, Nr 5, pp. 575-583 (USSR)

ABSTRACT:

The use of polyfunctional catalysts when the subsequent investigation of the isomerization of saturated hydrocarbons with a boiling-point up to 150°C possible. Nevertheless, it is very difficult to achieve the isomerization of paraffins (with a boiling-point above 200°C) in the presence of heterogeneous catalysts. The purpose of this work was the investigation of the structure and of the properties of the isomerizates. In all, the isomerization of a series of alkanes (structure C_9-C_{16}) was investigated in the presence of polyfunctional catalysts. It was found that ramified hydrocarbons, mainly with methyl-secondary-groups are formed due to the isomerization:

Card 1/2

Isomerization of Saturated Hydrocarbons. Communication No. 11
2. Isomeric Conversions of the Alkanes of the $C_{12} - C_{15}$ Series

the alkanes. Moreover, a catalyst with lower activity was found in the isomerization of $C_{12} - C_{15}$. A thermodynamic calculation of the reaction equilibrium constants of the alkanes of the $C_{12} - C_{15}$ was carried out. Tables and 12 references, 8 of which are Soviet.

ASSOCIATION: Institut khimicheskoi fiziki (Soviet Academy of Sciences, AS USSR)

SUBMITTED: November 19, 1956

Original in Russian. English translation by the U.S. Atomic Energy Commission.

AUTHORS: Petrov, A. A., Serdyukov, S. R., SOV. CHEM. REV. 1958, No. 11, p. 2177.
Tsedilina, A. L., Kislitskiy, A. N., Zhukovskiy, I. I.

TITLE: The Isomerization of Saturated Hydrocarbons (Izomerizatsiya nasyschennykh uslevodorstv) Communication 3. The Isomerization and Transformation of Cyclanes (Soobshcheniye 3. Izomerizatsiya i prevrascheniya tsiklanov)

PERIODICAL: Izvestiya Akademiya Nauk SSSR, Khimicheskaya Seriya, 1958, No. 11, pp. 2177-2182 (10 p.).

ABSTRACT: In various earlier papers the isomeric transformations of alkanes are discussed, which develop in the presence of functional catalysts under hydrogen pressure (Ref. 1, 2). The main purpose of this paper is the investigation of the factors governing the isomerization of hydrocarbons and of the connection between structure on the one hand and kinetic and thermodynamic parameters on the other. The catalytic isomerization of the cycloparaffin isomers $C_7 - C_{10}$ in the presence of functional catalyst under hydrogen pressure was investigated. Furthermore, it was found that the isomerization products of the cycloparaffins C_7 and C_8 correspond with the composition to the theoretical isomers with a certain regularity.

Card 1/2

The Isomerization of Substituted Hydrocarbons
Communication 1. The Isomerization of Substituted Hydrocarbons

A mechanism for the isomerization of substituted hydrocarbons is suggested, according to which the formation of a transition state of hydrocarbons is dependent on the first stage of reaction. From the influence exerted by the reaction temperature on the direction of the reactions of cyclohexanes it is concluded that a polyfunctional catalyst was used. The results are in good agreement with the references, if they are Soviet.

ASSOCIATION: Institut Nefti Akademii N. P. S. S. S. (Petroleum Institute USSR),
SUBMITTED: November 10, 1950

1. Hydrocarbons
--chemical effects of substituents on the reaction

Card 2/2

LEACV, L.A.

ways of synthesizing higher olefins (discussion). *Lefte* 1971a
3 no. 3:430-435 My-Je '63. (MIRA 14-9)

1. Institut geologii i razrabotki goryuchikh iskopnykh sosu-
dov. Otkrytiya i razrabotka toplivnykh promyshlennosti pri Gostlane. 1971.
(Olefins)

72

The synthesis of low-molecular hydrocarbons of mixed structure

ASSEMBLY: 1. 1968, 2. 1968, 3. 1968
(Petroleum Institute, AS USSR)

DATE: January 4, 1968, by D. I. Arizov, Doctor, Academy of Sciences, USSR

SUMMARY: December 17, 1967

1. Hydrocarbons--Synthesis 2. Hydrocarbons--Spectrographic analysis

Card 3, 3

PETROV, Aleksey Aleksandrovich, kand.khim.nauk; TAKOYEV, D.A., red.;
PETROPOL'SKAYA, N.Ye., red.; TASHEN'KINA, Ye.A., tekhn.red.

[Desalting and dehydration of petroleums] Obessolivanie i
obezvozhivanie neftei. Kuibyshev, Kuibyshevskoe knizhnoe
izd-vo, 1959. 82 p. (MIRA 13:2)
(Petroleum--Refining)

5 (3)

AUTHORS:

Petrov, Al. A., Sergiyenko, S. R.,
Nechitaylo, N. A., Tsedilina, A. L.

SOV'62-59-1-00114

TITLE:

Synthesis and Properties of the Monomethyl-substituted Alkanes
of the Composition $C_{12}-C_{16}$ (Sintez i svoystva monometil-
zameshchennykh alkanov sostava $C_{12}-C_{16}$)

PERIODICAL:

Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk
1959, No. 6, pp 1091 - 1097 (USSR)

ABSTRACT:

Since the hydrocarbons of this structure are not yet sufficiently investigated, the synthesis of the monomethylated alkanes with a boiling temperature of more than 200° was investigated. With this monomethylated alkanes one may synthesize all theoretically possible isomers (there are only 29 compounds). They are furthermore of interest because they have thermodynamically stable structures and meet with all technical requirements of motorization. The hydrocarbons were synthesized according to Grignard's reaction (magnesiumbromoalkyls with acetylketones). In this reaction the purity of the compounds obtained depends in a high degree on the purity of the initial substances (alkylbromides). This fact was especially considered

Card 1/4

Synthesis and Properties of the Monomethyl-substituted
Alkanes of the Composition $C_{12}H_{26}$

in the present investigation. The methylketones were obtained by decomposition of the corresponding alkyl-acetoacetic acid. As intermediates tertiary alcohols were at first obtained which were dehydrated at 280° , redistilled, and hydrated in the autoclave on nickel at a temperature of from $150-170^{\circ}$. The temperature at which the hydrocarbons obtained crystallize was determined by plotting the heating curve by means of the photo-recording Kurnakov pyrometer. The purity degree was determined by means of a special thermographic device. N. I. Lyashkevich to whom the authors express their gratitude, carried out the measurements in the laboratory for petroleum chemistry of the institute mentioned in the Association. The purity degree of the synthesized hydrocarbons was 97-98%. The thermogram was recorded by a special aluminum block which was designed in the Institut obshchey i neorganicheskoy khimii AN SSSR (Institute of General and Inorganic Chemistry of the AS USSR). By this automatic recording of the heating curves the melting process could be fixed exactly (Fig 1). With almost all compounds obtained two stages in the melting process (-46 and -45.4° melt-

Card 2/4

Synthesis and Properties of the Monomethyl-Substituted Branched Alkanes of the Composition $C_{12}H_{26}$

ing and crystallisation temperature, were investigated. The properties of the hydrocarbons are given in a table by which it is shown that the density and the refractive index scarcely depend on the position of the outer methyl groups in the main chain, what is well in line with data by Iatovskiy (Ref. 9). The crystallisation temperature is, however, strongly influenced by these methyl groups. The change of the crystallisation temperature at the transition of one isomer to the other, and of one isomer into the other is not steady (Figs. 1, 3 with comparative data from Ref. 10). This unsteadiness is caused by the presence of either an even or an odd number of hydrocarbon atoms in the main chain. The transition from an odd to an even number of hydrocarbons exerts a parallel influence on the melting point, the reverse transition, that is an increase in the molecular weight, exerts an antiparallel influence. The laws holding for paraffins, olefins, and greases, which are well known, may thus also be applied to ramified alkanes. The crystallisation process of the latter taken place by liquid and gaseous media exhibit the same stepwise character.

Card 3/4