

New Methods for the Synthesis of Organosiloxanes SOV/20-125-4-35/74

and leads to higher yields. A further advantage of this method is that it is possible to use also arylsilanes. This is impossible in the case of H_2SO_4 because of the disaryllating effect of this acid (Ref 15). It is easy to control the course of the reactions (1) and (4) according to the amount of hydrogen separated. Its rate can be easily controlled by the amount of the catalyst and the temperature of the medium. The mentioned reactions take place with solvents (benzene, toluene, petroleum ether etc) and without solvents (Ref 16). There are 1 table and 16 references, 3 of which are Soviet.

ASSOCIATION: Institut khimii silikatov Akademii nauk SSSR (Institute of Silicate Chemistry of the Academy of Sciences, USSR)

PRESENTED: December 7, 1958, by A. N. Nesmeyanov, Academician

SUBMITTED: November 4, 1958

Card 3/3

ORLOV, N. F., Cand Chem Sci -- (diss) "Synthesis of triorganosiloxy-derivatives of elements of the III, IV and V Groups from triorganosilanol and hexaalkylsiloxy-siloxanes," Leningrad, 1960, 18 pp (Leningrad State Univ im A. A. Zhdanov)

(KL, 36-60, 113)

ORLIV N F

Stekloobraznye sostoyaniya: trudy Tret'ego vostochno-sovetskaniya Leningrad, 1969.

16-20 noyabr 1969 (Vitreous State; Transactions of the Third All-Union Conference on the Vitreous State, held in Leningrad on November 16-20, 1969) Moscow, Izdatvo AN SSSR, 1969. 528 p. Errata slip inserted. 3,200 copies printed. (Series: Iss. Trudy)

Sponsoring Agencies: Institut khimii silikatov Akademi nauk SSSR. Vostochnyye opticheskiye obshchestvo Iseini D.I. Mendeleeva and Gosudarstvennyy opticheskiy Institut Iseini S.I. Vavilova.

Editorial Board: A.I. Argutinskiy, V.F. Barzakovskiy, M.A. Bezborodov, O.K. Borzishin, V.V. Vargin, A.G. Vlasov, K.S. Yevashovskiy, A.A. Lebedev, M.A. Matkovskiy, V.S. Molchanov, R.L. Myller, Ye.A. Ponomarevskiy, Chairman, N.A. Tolopon, V.A. Florinskaya, A.K. Yabinskiy; Ed. of Publishing House: I.V. Surorov, Tech. Ed.: V.T. Kochever.

PURPOSE: This book is intended for researchers in the science and technology of glasses.

COVERAGE: The book contains the reports and discussions of the Third All-Union Conference on the Vitreous State, held in Leningrad on November 16-20, 1969. They deal with the methods and results of studying the structure of glasses, the relation between the structure and properties of glasses, the nature of the chemical bond and glass structure, and the crystallochemistry of glasses. Pressed silica, mechanics of vitrification, optical properties and glass structure, and the electrical properties of glasses are also discussed. A number of the reports deal with the dependence of glass properties on composition, the tinting of glasses and radiation effects, mechanical, technical, and chemical properties of glasses. Other papers treat glass semiconductors and solar boron-doped glasses. The Conference was attended by more than 300 delegates from Soviet and East German scientific organizations. Among the participants the discussion were R.V. Solomin, Ye. Kuzminskiy, Yu.A. Gastev, V.P. Priglasnikov, Ya. Ya. Oortlib, O.P. Kuchelov, P. Kuznetsov, G.P. Mikhaylov, S.M. Petrov, A.N. Lazarev, D.I. Levin, A.V. Shatilov, M.T. Plonchinskii, A.Ya. Kuznetsov, K.K. Keller, Ya.A. Byrganovskaya, A.A. Salanov, M.M. Skorniyakov, P.F. Bolshakov, O.S. Molchanova, Kuznetsov, V.P. Podkov, R.S. Shevelovich, Z.G. Plinar, and O.S. Molchanova. The final session of the Conference was addressed by Professor I.I. Kitaygorodskiy, Honored Scientist and Engineer, Doctor of Technical Sciences. The following institutes were cited for their contribution to the development of glass science and technology: Gosudarstvennyy opticheskiy Institut (State Optical Institute), Institut khimii silikatov AN SSSR (Institute of Silicate Chemistry), Fizicheskii Institut AN SSSR (Physics Institute AS USSR), Fiziko-khimiya Institut AN SSSR (Physicochemical Institute AS USSR), Institut fiziki AS USSR, Minsk (Institute of Physics, Academy of Sciences, Belorussian SSR, Minsk), Laboratory of Physical Chemistry of Silicates of the Institut obschey i neorganicheskiy khimii AN SSSR (Institute of General and Inorganic Chemistry, Academy of Sciences, Belorussian SSR, Minsk), Institut vysokomolekulyarnykh soedineniy AN SSSR (Institute of High Molecular Compounds, AS USSR), Gosudarstvennyy Institut stekla (State Institute for Glass), Gosudarstvennyy Institut elektrotekhnicheskogo stekla (State Institute for Electrical, Glass), Sibirskiy fiziko-khimiya Institut (State Institute for Physical, Chemical Institute, Tomsk), Leningradskiy tekhnicheskii universitet (Leningrad State University), Mookovskiy Khimiko-tekhnologicheskii Institut im. Lencovetsa (Leningrad Chemical Technological Institute named after Lencovets), Belorusskiy politekhnicheskii Institut (Belorussian Polytechnical Institute, Minsk), Novosibirskiy politekhnicheskii Institut (Novosibirsk Polytechnical Institute), and Sverdlovskiy politekhnicheskii Institut (Sverdlovsk Polytechnical Institute). The Conference was sponsored by the Institut of Silicate Chemistry AS USSR (Acting Director - A.P. Orlikh), the Vostochnyye Khimicheskiye Obshchestvo im. D.I. Mendeleeva (All-Union Chemical Society named D.I. Mendeleev), and the Gosudarstvennyy opticheskiy Institut Iseini S.I. Vavilova. D.I. Vavilova (State Optical Institute named S.I. Vavilova) acted as the Chief for the purpose of coordinating their research on glass, to publish a book of the proceedings of the Conference in the series "Vostochnyye Opticheskiye Obshchestva" under the title "Fiziko-khimiya stekla". The Conference took place in Leningrad, and was attended by the members of the A.A. Lebedev, Academician, Professor of Physics and was attended by the members of the Organizational Committee, and R.L. Myller, Doctor of Chemical Sciences, Member of the Organizational Committee. The editorial board was M. V. Burdakov, M.V. Vavilovskiy, L.I. Pashkin, D.P. Podkov, S.K. Daryus, V.A. Izoff, and P.T. Florinskaya. References accompany individual reports.

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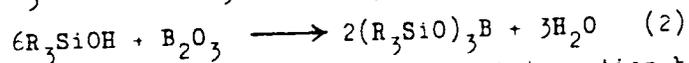
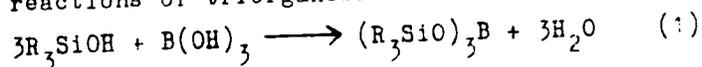
87122

S/062/60/000/009/000/000
B023/B064

15-2220

AUTHORS: Orlov, N. F., Dolgov, B. N., and Voronkov, M. G.
 TITLE: Tris(triorganosilyl)borate
 PERIODICAL: Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh
 nauk, 1960, No. 9, pp. 1607-1609

TEXT: In their development of new methods of synthesizing silicon elemental-organic compounds, the authors aimed at completing the previous methods of producing triorganosilyl esters of boric acid - tris(triorganosilyl)borate by new methods. Three new methods were developed. They distinguish themselves from the older ones by high yields (80-95%). The first two are based on the reactions of triorganosilanes with boric anhydride or boric acid:

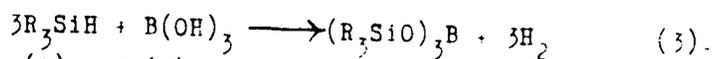


The third method of synthesis is based on the interaction between triorganosilanes and boric acid in the presence of colloidal nickel:

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87122

Tris(triorganosilyl)borate

S/062/60/000/009/009/001
B023/B064

The reactions (1) and (2) are based on an uninterrupted azeotropic distilling off of water from the mixture of triorganosilanoles with boric acid or with boric anhydride, containing an inert solvent (benzene). Tris(triorganosilyl)borate was produced according to scheme (3) by heating mixtures of triorganosilane and boric acid in the presence of colloidal nickel, forming in the interaction of nickel chloride with triorganosilane. 6 tris(triorganosilyl)borates were obtained by the three methods of synthesis. 4 of the compounds have hitherto been unknown. There are 1 table and 21 references: 8 Soviet, 6 US, 3 British, 1 German, and 3 Japanese.

ASSOCIATION: Institut khimii silikatov Akademii nauk SSSR (Institute of Silicate Chemistry of the Academy of Sciences USSR)

SUBMITTED: May 4, 1959

Card 2/2

82296

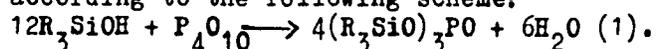
S/079/60/030/007/010/020
B001/B063

5.3700B

AUTHORS: Orlov, N. F., Voronkov, M. G.TITLE: Tris(triorganosilyl)phosphates 9

PERIODICAL: Zhurnal obshchey khimii, 1960, Vol. 30, No. 7, pp. 2223-2229

TEXT: The present paper describes new syntheses of tris(triorganosilyl) phosphates on the basis of triorganosilanols, hexaorganodisiloxanes, or triorganosilanes. It was found that these phosphates are formed with a yield of 50% by a reaction of phosphoric anhydride with triorganosilanols according to the following scheme:



A side reaction, $6R_3SiOSiR_3 + P_4O_{10} \longrightarrow 4(R_3SiO)_3PO$ (2), is to be assumed because of the comparatively high tendency of triorganosilanols to intermolecular condensation with water separation, which is accompanied by the formation of hexaalkyldisiloxanes, with P_4O_{10} acting as a catalyst (Ref. 18). The mechanism of this reaction is illustrated by Scheme (3).

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Tris(triorganosilyl)phosphates

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S/079/60/030/007/010/020
B001/B063

Another synthesis of tris(triorganosilyl)phosphates (yield of 90%), worked out by the authors, is based on a reaction of triorganosilanols with orthophosphoric acid: $3R_3SiOH + H_3PO_4 \rightleftharpoons (R_3SiO)_3PO + 3H_2O$ (4) in which the water must be distilled off (Ref. 22). A mechanism of this reaction is suggested in Scheme (5). An attempt to synthesize tris(triethylsilyl)phosphate according to Scheme (7) was unsuccessful, whereas orthophosphoric acid could be reacted with hexaalkyl disiloxanes according to Scheme (8). The mechanism of this reaction is shown in Scheme (9). A new interesting synthesis of tris(trialkylsilyl)phosphates with colloidal nickel acting as a catalyst is shown in reaction (10) in which a side reaction occurs, so that a yield of only 60% is obtained. All tris(triorganosilyl)phosphates obtained are only stable in dry air, whereas moist air splits the SiOP bond. The synthesized products are tabulated. A mechanism of the catalytic action of colloidal nickel is also suggested. There are 1 table and 26 references: 13 Soviet, 7 US, 1 Italian, 3 German, and 2 British.

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82296

Tris(triorganosilyl)phosphates

S/079/60/030/007/010/020
B001/B063

ASSOCIATION: Institut khimii silikatov Akademii nauk SSSR (Institute of
Silicate Chemistry of the Academy of Sciences USSR)

SUBMITTED: July 11, 1959

X

Card 3/3

S/661/61/OJO/006/026/021
D205/D302

AUTHORS: Orlov, N. F., Dolgov, B. N. and Voronkov, M. G.

TITLE: Synthesis of triorganosyloxy-derivatives of boron, titanium, phosphorous and vanadium, from triorgano-silanolis and hexaalkyl disiloxanes. Reaction of hexaalkyl disiloxanes with aluminum and titanium halides

SOURCE: Khimiya i prakticheskoye primeneniye kremneorganicheskikh soyedineniy; trudy konferentsii, no. 6, Doklady, diskussii resheniye. II Vses. Konfer. po khimii i prakt. prim. kremneorg. soyed., Len., 1958. Leningrad, Izd-vo AN SSSR, 1961, 123-126

TEXT: The field of elemental-silico-organic compounds is of practical interest because of the possibility of synthesis of compounds which are similar to inorganic compounds in their thermal stability while being soluble in organic solvents. The syntheses proposed give better yields than those previously described in literature. As to the application of the synthesized compounds, the silico-

Card 1/2

Card

ORLOV, N. F.

SOV/6034

PHASE I BOOK EXPLOITATION

Konferentsiya po khimii i primeneniyu fosfororganicheskikh soyedineniy. 2d. Kazan', 1959.

Khimiya i primeneniye fosfororganicheskikh soyedineniy; trudy (Chemistry and Use of Organophosphorus Compounds; Conference Transactions) Moscow, Izd-vo AN SSSR, 1962. 630 p. Errata slip inserted. 2800 copies printed.

Sponsoring Agency: Akademiya nauk SSSR. Kazanskiy filial.

Resp. Ed.: A. Ye. Arbuzov, Academician; Ed. of Publishing House: L. S. Povarov; Tech. Ed.: S. G. Tikhomirova.

PURPOSE: This collection of conference transactions is intended for chemists, process engineers, physiologists, pharmacists, physicians, veterinarians, and agricultural scientists.

COVERAGE: The transactions include the full texts of most of the scientific papers presented at the Second Conference on the Chemistry and Use of

Card 1/14

SECTION

Scientific Research Institute of Plastics,
for the Industrial Use of Organophosphorus

Card 2/14

Chemistry and the Use of Organophosphorus (Cont.)

SOV/6034

The preparation and uses of organophosphorus compounds are discussed, and prospects for obtaining new high-molecular organophosphorus compounds are outlined. Gefer recommends 1) the preparation of new organophosphorus polymers based on oxides of unsaturated tertiary phosphines, 2) the utilization of thermal stability and good mechanical properties of known organophosphorus compounds as a basis for creating new heterochain phosphorus-containing compounds, and 3) application of the latest methods currently used in the chemistry of high-molecular compounds to organophosphorus polymers.

Orlov, N. F., and M. G. Voronkov [Institut khimii silikatov (Institute of Silicate Chemistry, Leningrad)]. New Methods of Synthesis of Triorganosilyl Esters of Orthophosphoric and Alkylphosphonic Acids

212

New methods have been developed for the synthesis of tris-(triorganosilyl)-phosphates $(R_3SiO)_3PO$, bis-(triorganosilyl)methylphosphonates $CH_3(R_3SiO)_2PO$, and organophosphorussilicon polymers based on H_3PO_4 and $CH_3PO(OH)_2$. One method is based on the interaction of

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S/051/62/012/002/016/020
E202/E192

AUTHORS: Byurganovskaya, G.V., and Orlov, N.F.

TITLE: Formation of colour centres in quartz and sodium silicate glasses under the action of gamma radiation

PERIODICAL: Optika i spektroskopiya, v.12, no.2, 1962, 278-284

TEXT: The relations between the absorption bands of quartz glasses and sodium silicate glasses, and the amount of gamma irradiation, were studied together with the effect of traces of cerium, antimony, arsenic, manganese, iron and lead on the properties of these bands. A large number of industrial and experimental glasses were used in the studies of colour centres. The glasses were melted preferably in an oxy-fuel burner. The irradiation was started at ambient temperature. Co^{60} , at 9×10^3 roentgen/hour was used as a gamma source. The spectral absorption of the glasses was measured on a CΦ-4 (SF-4) spectrophotometer and expressed as optical density. Quartz glasses showed three characteristic spectral absorption curves. The first, exhibited by glasses melted in reducing conditions or containing large admixtures of elements of Group I and III, had three groups of
Card 1/2

Formation of colour centres in ...

S/051/62/012/002/016/020
E202/E192

strong bands with centroids at 540, 300 and 200-210 m μ . The second, represented by optical quality quartz glasses, had a weak absorption band with a maximum in the visible spectrum, discernible only after a large dose of irradiation ($1 \times 10^7 - 1 \times 10^8$ roentgen). The third, characteristic of high purity quartz glasses, had only one shortwave UV absorption band with a maximum at 200-210 m μ . The formation of colour centres for all quartz glasses was the same, i.e. requiring doses within the ranges of $(4-8) \times 10^4$ and $(2-4) \times 10^6$ roentgen. The silica glasses studied were based on $\text{Na}_2\text{O} \cdot 4\text{SiO}_2$ and $\text{Na}_2\text{O} \cdot 3\text{SiO}_2$, with the additives of oxides of the above listed elements within 0.1-0.2% w/w. The rate of formation and quantity of colour centres in these glasses were found to be influenced by the additives. All colour centres due to irradiation were of one type only. The anomalies in the relations between the optical absorption and the irradiating dose in glasses containing Fe_2O_3 were not fully explained, but were thought to be due to the traces of contaminants or due to the conditions of melting.

There are 8 figures. SUBMITTED: January 21, 1961
Card 2/2

352

S/079/62/032/002/008/011
D204/D303

5. 3700

AUTHORS: Orlov, N.F. and Voronkov, M.G.

TITLE: Silico-organic esters of methyl phosphinic acid (MPA)

PERIODICAL: Zhurnal obshchey khimii, v.32, no. 2, 1962, 608-612

TEXT: This is a description of the methods of synthesis of the hitherto unknown bis (triorganosilyl) methyl phosphinates, based on interaction of $\text{CH}_3\text{P}(\text{O})(\text{OH})_2$ with: (1) Triorganosilanol $\text{CH}_3\text{P}(\text{O})[\text{OSiEt}_3]_2$ was prepared in 68% yield from Et_3SiOH (2 moles) and MPA (1 mole), by heating for 2 hrs, removing continuously the water formed. (2) Triorganosilanes $\text{CH}_3\text{P}(\text{O})[\text{OSiEt}_2\text{Me}]_2$, $\text{CH}_3\text{P}(\text{O})[\text{OSiEt}_3]_2$, $\text{CH}_3\text{P}(\text{O})[\text{OSiMe}(\text{n-Pr})_2]_2$, $\text{CH}_3\text{P}(\text{O})[\text{OSiMe}(\text{Ph})_2]_2$ and $\text{CH}_3\text{P}(\text{O})[\text{OSiEt}(\text{Ph})_2]_2$ were synthesized from the corresponding trisubstituted silanes and MPA in the molar ratio of 2:4:1, in presence of colloidal Ni, in 70-90% yields, by heating for 2-3 hr at 130°C . (3) Triorganoalkoxysilanes $\text{CH}_3\text{P}(\text{O})[\text{OSiEt}_3]_2$ and $\text{CH}_3\text{P}(\text{O})$

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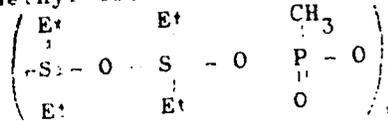
3973

S/079/62/032/002/008/011

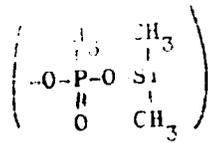
D204/D303

Silico-organic esters of ...

$[\text{OSi}(\text{Ph})_3]_2$ were obtained from Et_3SiOMe and $(\text{Ph})_3\text{SiOEt}$ taken in the molar ratio of silane MPA equal to 2.4:1 and 2:1 respectively, in 60 and 70% yields. The reaction of MPA and tetraethyl disiloxane (1:1 ratio) was catalyzed by colloidal Ni and yielded



whilst another polymer



was obtained from an equimolar

lar mixture of MPA and dimethyl diethoxysilane. All the phosphates with the exception of crystalline $\text{CH}_3\text{P}(\text{O})[\text{OSi}(\text{Ph})_3]_2$ were colorless, high-boiling liquids, readily hydrolyzed by water. The polymers were pale yellow oils. Experimental details are given and physical constants of

Card 2/3

35923
S/079/62/032/002/008/011
D204/D303

Silico-organic esters of ...

the products are tabulated. There are 1 table and 17 references: 11 Soviet-bloc and 6 non-Soviet-bloc. The 4 most recent references to the English-language publications read as follows: R.O. Sauer, J. Am. Chem. Soc., 66, 1707, (1944); W.H. Keeber and H.W. Post, J. Org. Ch., 21, 509, (1958); Canadian Pat. 496,623 (1953); Ch. A., 47, 4128, (1953).

ASSOCIATION Institut khimii silikatov Akademii nauk SSSR (Institute of Silicate Chemistry of the Academy of Sciences, USSR)

SUBMITTED March 2, 1961

Card 3/3

S/079/62/032/002/009/011
D204/D303

AUTHORS: Orlov, N.F., Bogatkin, R.A., Sergeyeva, Z.I., and Veronkov,
M.G.

TITLE: Interaction of triorganosilanes with carboxylic acids in
the presence of colloidal nickel

PERIODICAL: Zhurnal obshchey khimii, v. 32, no. 2, 1962, 650-651

TEXT: A short note on the reactions of triethyl silane with carboxylic acid, using colloidal Ni as a catalyst. Monocarboxylic acids reacted giving the corresponding triorganosilyl esters, in 50-85% yield. Esters of general formula $\text{Et}_3\text{SiOCO}(\text{CH}_2)_n\text{OCOSiEt}_3$ were synthesized in 60-80% yields from simple dicarboxylic acids. Colloidal Ni promoted hydrogenation as well as dehydrocondensation, as was shown by the reactions of Et_3SiH with halogenated and unsaturated acids. Monochloroacetic acid yielded either $\text{Et}_3\text{SiOCOCH}_2\text{Cl}$ or $(\text{Et}_3\text{SiOCOCH}_3 + \text{Et}_3\text{SiCl})$, depending on the molar ratio of the reagents. Unsaturated acids yield hydrogenated

Card 1/2

ORLOV, N.F.; BOGATKIN, R.A.; SERGEYEVA, Z.I.; VORONKOV, M.G.

Interaction of hydroxysilanes with saturated acids in the
presence of colloidal nickel. Zhur.ob.khim. 32 no.8:2561-2566
Ag '62. (MIRA 15:9)
(Silane) (Acids, Organic)

ORLOV, N.F.; BOGATKIN, R.A.; SERGEYEVA, Z.I.; VORONKOV, M.G.

Nickel catalyst in the reactions of organosilane hydrides with
organic acids. Zhur.ob.khim. 33 no.6:1934-1938 Je '63.
(MIRA 16:7)

1. Leningradskiy gosudarstvennyy universitet i Institut
khimii silikatov AN SSSR.
(Silane) (Acids, Organic) (Nickel catalysts)

I 34860-65 EPA(s)-2/EWT(m)/EPF(c)/EPR/EWP(j)/I/EWA(c) Pc-4/Pt-4/Ps-4/Pt-10 WH/RM
 ACCESSION NR: AP5008143 S/0286/65/000/005/0022/0022

AUTHOR: Minsker, Ye. I.; Solovey, G. G.; Borisov, M. F.; Orlov, N. E.

TITLE: A method of preparing polyaluminodiorganosiloxanes, Class 12, No. 168689

SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 5, 1965, 22

TOPIC TAGS: elastomer, polymer, aluminum containing polymer, polysiloxane

ABSTRACT: This Author Certificate introduces an improved preparative method for aluminum-containing polysiloxanes utilizing dihydroxypolydiorganosiloxanes and methylpropylsiloxy-bis-(β -chloroalkoxy)-aluminum and triphenylsiloxydiisopropoxy-aluminum as starting materials. This results in a product with improved thermal stability. [VS]

ASSOCIATION: none

SUBMITTED: 29Dec63

ENCL: 00

SUB CODE: 00,00

NO REF SOV: 000

OTHER: 000

ATD PRESS: 3211

Card 1/1

I 35556-65 EWT(m)/EPF(c)/EWP(j) Pc-4/Pr-4 RM 8/0286/65/000/005/0023/0023
ACCESSION NR: AP5008146

AUTHORS: Orlov, N. P.; Milashkevich, V. P.; Gefter, Ye. L.

TITLE: A method for obtaining triorganosilanol esters of oximethylphosphinic acid.
Class 12, No. 168695

SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 5, 1965, 23

TOPIC TAGS: ester, triorganosilanol, hydroxymethylphosphinic acid,
triorganoalkoxysilane

ABSTRACT: This Author Certificate presents a method for obtaining triorganosilanol esters of hydroxymethylphosphinic acid. The latter is heated to 100-230C and is subjected to interaction with triorganoalkoxysilanes.

ASSOCIATION: none

SUBMITTED: 13Jan64

ENCL: 00

SUB CODE: 00

NO REF SOV: 000

OTHER: 000

Card 1/1

L 65102-65 EWT(m)/EPP(c)/EWP(j)/T RM

ACCESSION NR: AP5021973

UR/0286/65/000/014/0024/0024
547.419.1.5.07

AUTHOR: Orlov, N. F.⁶⁶; Mileshevich, V. P.⁵¹; Vaynburg, V. M.⁵²

TITLE: A method for producing organosilicon esters of chlorinated α - or β -organophosphonic acids, Class 12, No. 172788¹⁵

SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 14, 1965, 24¹¹

TOPIC TAGS: chlorinated organic compound, organic phosphorus compound, silane esterification, phosphonic acid 24
B

ABSTRACT: This Author's Certificate introduces a method for producing organosilicon esters of chlorinated α - or β -organophosphonic acids containing a radical of the $\text{RCHCl}(\text{CH}_2)$ type at the phosphorus atom. Organophosphonyl chlorides are reacted

L 65101-65 SWP(m)/EPF(c)/EWP(j)/T RM

ACCESSION NR: AP5021972

UR/0286/65/000/014/0024/0024
547.419.1.5.07

AUTHOR: Orlov, N. F.; Volodina, L. N.

7.65
22
B

TITLE: A method for producing bis-(triorganosilyl)-phosphinates. Class 12,
No. 172787

SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 14, 1965, 24

TOPIC TAGS: organosilicon compound, organic phosphorus compound, silane, phosphorus acid

ABSTRACT: This Author's Certificate introduces a method for producing bis-(triorganosilyl)-phosphinates by interacting monofunctional silanes with phosphorous acid in the presence of heat. A wider selection of raw materials is provided by using triorganoacetoxysilanes.

ASSOCIATION: none

SUBMITTED: 19Jun64

ENCL: 00

SUB CODE: 00, C-C

NO REF SOV: 000

OTHER: 000

Card 1/1 MRP

L 65103-65 EWT(m)/EPT(c)/EWP(j)/T RM

ACCESSION NR: AP5021974

UR/0286/65/000/014/0025/0025
547.419.5.1.07

AUTHOR: Orlov, N. F.; Mileshevich, V. P.

22
B 755

TITLE: A method for producing dialkyl-(triorganosilylmethoxymethyl)-phosphinates.
Class 12, No. 172789

SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 14, 1965, 25

TOPIC TAGS: chlorinated aliphatic compound, alkylphosphine, organosodium compound, silane, organosilicon compound

ABSTRACT: This Author's Certificate introduces a method for producing dialkyl-(triorganosilylmethoxymethyl)-phosphinates. Dialkyl-(sodium hydroxymethyl)-phosphinates are interacted with chloromethyltriorganosilanes with the application of heat in an organic solvent.

ASSOCIATION: none

SUBMITTED: 24Jun64

ENCL: 00

SUB CODE: 00, 60

NO REF SOV: 000

OTHER: 000

Card 1/1
MOR

L 2948-66 EWT(m)/EPP(c)/EWP(j)/T RM

ACCESSION NR: AP5024970

UR/0286/65/000/016/0033/0033

547.419.1.07

AUTHOR: Orlov, N. P.; Milieshevich, V. P.; Andronov, Ye. S.

TITLE: Preparation of organosilicon derivatives of hydroxyalkylphosphonic acids.
Class 12, No. 173761

SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 16, 1965, 33

TOPIC TAGS: organosilicon compound, phosphonic acid, phosphonate

ABSTRACT: An Author Certificate has been issued for a preparative method for organo-
silicon hydroxyalkylphosphonic acid derivatives involving the reaction of phosphonic
acid derivatives with alkylchlorosilanes in organic solvents. The method provides
for the use of dialkyl(sodiooxymethyl)phosphonates as the phosphonic acid derivatives.

[B0]

ASSOCIATION: none

SUBMITTED: 10Jul64

ENCL: 00

SUB CODE: DC,GC

NO REF SOV: 000

OTHER: 000

ATD PRESS: 4/08

Card 1/1 DP

L 11136-66 EWT(m)/EWP(j) RM

ACC NR. AP6002513

SOURCE CODE: UR/0286/65/000/023/0018/0018

INVENTOR: ^{44 55} Orlov, N. F.; ^{44 55} Kaufman, B. L.

ORG: none

TITLE: Preparation of organosilicon esters of acetophosphonic acid.
Class 12, No. 176585, 16

SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 23, 1965, 18

TOPIC TAGS: organosilicon phosphinates, synthetic material

ABSTRACT: An Author Certificate has been issued for a preparative method for organosilicon esters of acetophosphonic acid [sic]. The method involves heating of triorganacetoxysilanes with phosphorus trichloride in an inert gas. [B0]

SUB CODE: 07, 11/ SUBM DATE: 14Oct64/ ATD PRESS: 4173

PC

UDC: 547.419.5.07
547.419.1.07

Card 1/1

Im. G. M. Karava

L 42422-65 EWT(m)/EPF(c)/BWP(j) Re-4/Pr-4 RM S/0079/65/035/003/0590/0591
ACCESSION NR: AP5008842

*27
26
D*

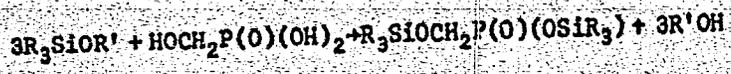
AUTHOR: Orlov, N. F.; Mileshevich, V. P.; Gefer, Ye. L.

TITLE: The reaction of hydroxymethylphosphinic acid with triorganoalkoxysilanes

SOURCE: Zhurnal obshchey khimii, v. 35, no. 3, 1965, 590-591

TOPIC TAGS: silicon organic compound, organic synthesis, methylol group, phosphinic acid

ABSTRACT: The reaction of organoalkoxysilanes with hydroxymethylphosphinic acid, HOCH₂P(O)(OH)₂ is studied. It was found experimentally that alcohol and acid group reactions with triorganoalkoxysilanes take place at comparable rates and the main reaction product is a triorganosilyl ester of hydroxymethylphosphinic acid, which is formed with 45-55% yield as follows:



The reaction is carried out by heating a mixture of triorganoalkoxysilane with hydroxymethylphosphinic acid in a molar ratio of 3:1. The alcohol formed during the reaction is removed and the reaction product is then isolated. The

L 42422-65

ACCESSION NR: AP5008842

alcohol yield is 75-100% of the theoretical yield. The reaction takes 6-8 hours.
The authors give the properties and analytical data for the synthesized compounds.

ASSOCIATION: Leningradskiy institut tekstil'noy promyshlennosti im. S. M. Kirova
(Leningrad Institute of the Textile Industry)

SUBMITTED: 09Oct64

ENCL: 00

SUB CODE: 00

NO REF SOV: 002

OTHER: 000

cc
Card 2/2

OBLOV, N.F., MILIKOVICH, V.F., GEFTEL, Ya.

Synthesis of bis (inorganically and organically) phosphinates by the reaction of organoaluminum hydrides with hydroxymethylphosphinic acid. Zhur. obshch. khim. 1965, 39, 1312-1313, 1315.

1. Leningradskiy institut tekstil'noy i legkoy promyshlennosti im. S.M. Kirova.

L 1811-66 BT(m)/EPF(c)/BWP(j)/T RM
ACCESSION NR: AP5024002/

UR/0020/65/164/002/0344/0346

AUTHOR: Orlov, N. F.; Mileshevich, V. P.

TITLE: Dialkyl triorganosilylmethoxymethylphosphonates and their stability toward hydrolysis -

SOURCE: AN SSSR. Doklady, v. 164, no. 2, 1965, 344-346

TOPIC TAGS: polymer, silicon containing polymer, phosphorus containing polymer, phosphonic acid, phosphonic ester

ABSTRACT: Previous work had shown that silicon- and phosphorus-containing organic compounds, although of promise in various practical applications, suffer from insufficient stability toward hydrolysis. The only class of compounds in which hydrolysis does not produce scission into silicon- and phosphorus-bearing fragments is Si-(CH₂)_n-P. This work deals with the synthesis of esters of trialkylsilylmethoxymethylphosphonic acids



Card 1/2

L 1814-66
ACCESSION NR: AP5024002

where R = alkyl, R' = alkyl, R'' = alkyl or alkoxy. When R'' = alkoxy, the reaction is completed in 5-7 hr. The starting reagents are taken in equimolar ratios, the yields are 50-55%. The colorless, liquid products can be distilled under vacuum without decomposition. On boiling for 20 hr with 18% hydrochloric acid, they are converted to the corresponding phosphonic acids in 85-90% yield. Compounds containing alkoxy groups at the Si atom undergo hydrolysis with appreciable breakup of the molecular skeleton and formation of polymeric materials. The results obtained suggest that the structure P-C-O-C-Si is considerably more stable toward hydrolysis than Si-O-P, Si-C-O-P, Si-C-O-P(S), and approaches Si-(CH₂)_n-P with respect to hydrolytic stability. Orig. art. has: 1 table. [VS]

ASSOCIATION: Leningradskiy institut tekstil'noy i legkoy promyshlennosti im. S. M. Kirova (Leningrad Institute of the Textile and Light Industry)

SUBMITTED: 16Mar65

ENCL: 00

441/65
SUB CODE: OC, GC

NO REF SOV: 003

OTHER: 005

ATD PRESS: 4111

Card 2/2

ORLOV, N.F.; MILESHKEVICH, V.I.; ANDRONOV, Ye S.

Synthesis of dialkylphosphonemethoxyorganosilanes and their
thermal decomposition. Zhur.ob.khim. 35 no.12:2193-2197
'65. (MIRA 1961)

1. Leningradskiy institut tekstil'noy i legkooy promyslovenosti
imeni S.M.Kirova. Submitted March 31, 1965.

I 28360-66 EPF(n)-2/EWI(1)/EWI(m)/EWP(e) GG/NH

SOURCE CODE: UR/0368/66/004/004/0323/0326

ACC NR: AF6012855

AUTHOR: Orlov, N. F.; Iakob, N. A.

ORG: none

TITLE: Absorption centers produced at low temperatures in certain glasses of simple composition

SOURCE: Zhurnal prikladnoy spektroskopii, v. 4, no. 4, 1966, 323-326

TOPIC TAGS: borate glass, silicate glass, glass property, gamma radiation, color center, radioluminescence, absorption spectrum

ABSTRACT: To obtain more information on the nature of the color centers and radioluminescence in irradiated glass, the authors investigated the effect of γ rays from Co^{60} at liquid-nitrogen temperature on various glasses. The experiment was carried out in a dismantlable glass Dewar with side windows of quartz glass, immune to discoloring by γ rays. The γ -ray dose intensity was 4100 r/hr. The induced absorption spectra of the glasses were measured at the same temperature by determining the difference between the optical densities of the irradiated and non-irradiated glasses. The results show that the spectra of both glasses change noticeably at low temperatures. In the sodium-silicate and non-alkali sodium-borosilicate glass, a strong increase was observed in the band near 640 nm. In the case of potassium-silicate glass, a new band was observed at 820 nm. In the case of sodium-borate glass and alkali sodium-borosilicate glass the intensity of the 240 nm band increased. All these

UDC: 661.11: 639.16

Cont 1/2

53
52
B

L 32200-66 EWT(m)/T/EWP(1) IJP(c) WW/RM

ACC NR: AP6007119

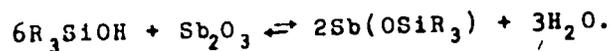
SOURCE CODE: UR/0079/66/036/002/0347/0350

AUTHOR: Orlov, N. F.; Voronekov, M. G.45
44
BORG: Leningrad Institute of Textiles and Light Industry im. S. M. Kirov (Leningradskiy institut tekstil'noy i legkoy promyshlennosti); Institute of Organic Synthesis, AN LatSSR (Institut organicheskogo sinteza AN LatSSR)TITLE: Tris(triorganylsilyl) antimonites ↑

SOURCE: Zhurnal obshchey khimii, v. 36, no. 2, 1966, 347-350

TOPIC TAGS: antimony compound, organic synthetic process, silicon compound

ABSTRACT: A study by N. F. Orlov and M. G. Voronkov contains data on two new methods for the synthesis of tris(trialkylsilyl) antimonites. The first method (A), which is simple and economical, consists of reacting a trialkylsilanol with antimony trioxide on heating:



The water formed is removed by azeotropic distillation with an inert solvent, e. g., benzene, toluene, etc. The yield reaches 70% of the theoretical. Previously, the authors used a similar method to prepare tri-

Card 1/3

INC. 516.287+516.863

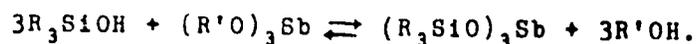
L 32200-66

ACC NR: AP6007119

silanol derivatives of some other trivalent elements or groups: boron, phosphorus, or vanadyl. 0

The method is similar to that used in the preparation of trialkyl antimonites, $Sb(OR)_3$, by reacting Sb_2O_3 with alcohols. The higher reactivity of trialkylsilanols, as compared with alcohols, can be explained by their higher acidity caused by the $p_\pi - d_\pi$ interactions between silicon and oxygen atoms.

The second method (B) suggested is the silanolysis of trialkyl antimonites:



On heating of the mixture the alcohol formed is stripped off. It is advisable to add the silanols in small quantities, e. g., dropwise, especially in benzene solution, in order to prevent silanol condensation, thereby increasing the yield of the desired product.

The presence of sodium in catalytic amounts accelerates the reaction; however, the reaction proceeds readily even without any catalyst.

Tris(trialkylsilyl) antimonites are either colorless liquids or crystals with a weak silanol odor. They are stable on storage in dry air, but

Card 2/3

L 32200-66

ACC NR: AP6007119

hydrolyze in humid air. Hydrolytic stability of tris(trialkylsilyl) antimonites increases with increasing length of the organic radical, and is higher than that of the trialkyl antimonites.

The properties of the five compounds synthesized (four of them for the first time) are given in Table 1.

Table 1. Tris(trialkylsilyl) antimonites

Compound	Preparative method	Yield %	bp °C (p mm Hg)	d ₄ ²⁰	n _D ²⁰	MR _D		Found %		Formula	Calculated	
						Found	Calculated	Si	Si		Si	Si
(CH ₃) ₃ SiO ₂ Sb	B	07.0	80° (3)	1.1448	1.4374	89.17	89.13	31.25,	21.52,	C ₉ H ₂₇ O ₂ SbSi ₃	31.27	21.84
(CH ₃) ₂ C ₁₁ SiO ₂ Sb	B	00.0	132 (6)	1.1318	1.4508	102.59	102.66	31.71	21.29	C ₁₉ H ₃₃ O ₂ SbSi ₃	28.22	—
(CH ₃) ₂ (C ₂ H ₅) ₂ SiO ₂ Sb	A	61.5	150 (3.5)	1.1132	1.4588	116.23	116.19	28.25,	—	C ₁₃ H ₂₉ O ₂ SbSi ₃	25.72	17.80
(C ₂ H ₅) ₃ SiO ₂ Sb	A	83.0	170 (3)	1.1041	1.4675	129.69	129.72	25.75,	17.60,	C ₁₈ H ₃₅ O ₂ SbSi ₃	23.81	16.34
(C ₂ H ₅) ₂ (C ₄ H ₉) ₂ SiO ₂ Sb	B	90.5	203 (30)	1.1038	1.4681	129.04	129.72	25.64,	17.44,	C ₂₁ H ₄₃ O ₂ SbSi ₃	23.81	16.34
	B		mp 214° (benzene)			—	—	23.83,	18.01,	C ₂₁ H ₄₃ O ₂ SbSi ₃	12.84	8.89
								23.89	18.01			
								—	16.08,			
								12.68,	18.14			
								12.18	8.39			

The refraction increment for the Sb-O bond, absent in the literature according to the authors, was computed to be equal to $5.25 \pm 0.05 \text{ cm}^3$. Orig. art. has: 1 table. [FSB: v.2, no. 5]

SUB CODE: 07 / SUBM DATE: 05Jan65 / ORIG REF: 010 / OTH REF: 002

Card 3/3

1-32718-66 FMP(j)/EWT(m) RM
ACC NR: AP6021414

SOURCE CODE: UR/0413/66/000/011/0019/0019

INVENTOR: Orlov, N. F.; Kaufman, B. L.

11
B

ORG: none

TITLE: Preparative method for organic organosilicon esters of acetylphosphonic acid. Class 12, No. 182146 (announced by Leningrad Institute of the Textile and Light Industry im. S. M. Kirov (Leningradskiy institut tekstil'noy i legkov promyshlennosti))

SOURCE: Isobreteniya, promyshlennyye obrabotki, tovarnyye znaki, no. 11, 1966, 19

TOPIC TAGS: organosilicon ester, acetylphosphonic acid

ABSTRACT: An Author Certificate has been issued for a preparative method of organic organosilicon esters of acetylphosphonic acid. The method involves heating of alkyl(aryl)dichlorophosphites with triorganoacetoxysilanes. [B0]

SUB CODE: 07 / SUBM DATE: 03Apr65/ ATD PRESS: 5025

Cord 1/1 JS

UDC: 547.419.5'419.1.07

L 31797-66 ENT(m)/ENP(j) RM

ACC NR: AP6021688

SOURCE CODE: UR/0079/66/036/003/0518/0521

AUTHOR: Orlov, N. F.; Mlleshkevich, V. P.

50

B

ORG: Leningrad Institute of the Textile and Light Industry im. S. M. Kirov
(Leningradskiy institut tekstil'noy i legkoy promyshlennosti)TITLE: Synthesis and certain properties of organosilicon esters of hydroxymethylphosphinic acid

SOURCE: Zhurnal obshchey khimii, v. 36, no. 3, 1966, 518-521

TOPIC TAGS: chemical synthesis, phosphinic acid, ester, organosilicon compound, silane, hydroxyl group, chemical bonding, heterogeneous catalysis, acetic anhydride, chemical reaction, reaction mechanism

ABSTRACT: A method for producing bis(triorganosilyl)hydroxymethylphosphinates not previously described in the literature was developed. The reaction of triorganoalkoxysilanes with hydroxymethylphosphinic acid in a 2:1 ratio is conducted under mild conditions, distilling off the alcohol under vacuum while heating the reaction mixture to a temperature not higher than 50-60°. The formation of such products is direct evidence that the hydroxyl bonded to the phosphorus through the carbon atom in the hydroxymethylphosphinic acid molecule is less reactive than the acid hydroxyls toward alkoxysilanes. The reaction of bis(trialkylsilyl)hydroxymethylphosphinates with trialkylsilane

Card 1/2

UDC: 547.281.1+546.185

L 31797-66

ACC NR: AP6021688

hydrides in the presence of a colloidal nickel catalyst and with acetic anhydride in the presence of catalytic amounts of sulfuric acid results in the formation of esters of trialkylsiloxymethylphosphinic or acetoxymethylphosphinic acids, respectively. Orig. art. has: 1 table. [JPRS]

SUB CODE: 07 / SUBM DATE: 02Mar65 / ORIG REF: 006 / OTH REF: 005

Card 2/2

1 14641-66

ACC NR: AP6004093

quaternary ammonium catalysts. It was shown that acidic catalysts such as boron trifluoride etherate promote polymerization of the obtained glycidylmethylphosphonates; this is accompanied by cleavage of the epoxide ring. Orig. art. has: 1 table
[VS]

SUB CODE: 11.071 SUBM DATE: 28Apr65/ ORIG REF: 006/ OTH REF: 001/ ATD PRESS: 4198

Card 212 AC

I 40967-66 EWP(e)/EWT(m) WH

ACC NR: AP6023365

SOURCE CODE: UR/0237/66/000/007/0006/0011

AUTHOR: Orlov, N. F.

ORG: none

TITLE: Radioluminescence in nonorganic glasses

SOURCE: Optiko-mekhanicheskaya promyshlennost', no. 7, 1965, 6-11

TOPIC TAGS: optic glass, radioluminescence, phosphorescence, impurity center, nuclear radiation, Cerenkov radiation, Cerenkov effect, radiation dosimetry, cerium

ABSTRACT: General questions concerning luminescence in nonorganic glasses excited by nuclear and other types of radiation are discussed. Radioluminescence of glass in high energy fields is considered in the following terms: a) luminescence of active centers and particular types of structural formations initially present in the glass; b) luminescence of impurity centers and certain types of trap centers, generated by the interaction of radiation with the material; c) luminescence due to radicals created in the process of irradiation; d) Cerenkov effect. Type a) luminescence is observed in lead and cerium glasses. Since only trivalent cerium is capable of luminescence, the brightness of luminescence in optic glasses, normally containing tetravalent cerium, is low. Due to high concentration of lead in flint glasses the extinction is very pronounced, reducing the specific intensity of luminescence. Other activators, e. g., titanium and

UDC: 666.11.01 : 535.37

Card 1/2

47
B

L 40967-66

ACC NR: AP6023365

antimony, present in glass as impurities, do not exhibit measurable amounts of luminescence. Absorption zones present in quartz glass are an example of structural formations capable of luminescence. Type b) luminescence is due to impurity elements not initially contained in the glass, but generated due to the interaction of nuclear particles with the material. The magnitude of this effect is determined by the properties of radiation and the initial concentration of appropriate elements. This effect is utilized in radiation dosimetry. Type c) luminescence results from radicals created due to the ionizing effects of radiation and the breakup of the molecular bonds. Type d) luminescence due to the Cerenkov effect is more pronounced in heavy flint glass, than in crown glass. This effect is not temperature dependent. The contribution of type b) luminescence is negligible, compared to the other three types. The effects of temperature, phosphorescence, effects of glass darkening (due to irradiation), and glass structure on radioluminescence are also discussed. The spectral output curves for luminescence of various types of glass are given. Orig. art. has: [14]
4 figures, 1 table.

SUB CODE: 20,11/

SUBM DATE: 27May65/

ORIG REF: 002 / ATD PRESS: 5056

Card 2/2 MLP

L 40809-66 EWT(m)/T/EWP(j) IJP(c) WW/RM

ACC NR: AP6025590

SOURCE CODE: UR/0413/66/000/013/0020/0020

AUTHORS: Orlov, N. F.; Mileshevich, V. P.

32
B

ORG: none

TITLE: Preparative method for silicon-containing hydroxymethylphosphonates Class 12, No. 183207 *6*

SOURCE: Izobreteniya, promyshlennyye obraztsy, tovarnyye znaki, no. 13, 1966, 20

TOPIC TAGS: ~~organosilanes~~, ~~hydroxymethylphosphonic acid~~, organosilicon compound

ABSTRACT: This Author Certificate presents a method for preparing silicon-containing (hydroxymethyl)phosphonates. (Hydroxymethyl)phosphonic acid reacted with organo(alkoxy)silanes at 40-800C. [04]

SUB CODE: 07/ SUBM DATE: 28Dec64/ ATD PRESS: 5059

Card 1/1

1964 12 28 15.07

L 05182-67 EWT(m)/EWP(j) RM

AQC NR: AP7000742

SOURCE CODE: UR/0079/66/036/005/0920/0923

ORLOV, N. F., VOLODINA, L. N., Leningrad Institute of the Textile and Light Industry S. M. Kirov (Leningradskiy institut tekstil'noy i legkoy promyshlennosti)

"Bis(triorganylsilyl)phosphites" ↑

30
B

Moscow, Zhurnal Obshchey Khimii, Vol 36, No 5, 1966, pp 920-923

Abstract: Two new methods were developed for synthesizing bis(triorganylsilyl) phosphites, based on the reaction of phosphorous acid with hexaalkyldisiloxanes and triorganylacetoxysilanes. 5-Bis(triorganylsilyl)-phosphites were synthesized; among them bis(dimethylethylsilyl) phosphite and bis(dimethylphenylsilyl) phosphite were prepared for the first time. The bis(triorganylsilyl)phosphites are colorless liquids with a weak silanol odor, stable in dry air and readily hydrolyzed by water to form the original phosphorous acid and triorganylsilanols. The latter are usually dehydrated to form hexaorganylidisiloxanes.

Orig. art. has: 2 formulas and 1 table. [JPRS: 37,023]

TOPIC TAGS: siloxane, organic phosphorus compound

SUB CODE: 07 / SUBM DATE: 21Apr65 / ORIG REF: 007 / OTH REF: 001

Card 1/1 vmb

UDC: 546.287 + 547.26.118
092.3 1912

L 04850-67 EWP(j)/EWT(m) RM

ACC NR: AP7000237

SOURCE CODE: UR/0079/66/036/004/0699/0704

AUTHOR: Orlov, N. F.; Mileshevich, V. P.

ORG: Leningrad Institute of the Textile and Light Industry im. S. M. Kirov (Leningrad-skiy institut tekstil'noy i legkoy promyshlennosti)

21
8

TITLE: Reaction of Hydroxymethylphosphinic Acid with Trialkylacetoxy-silanes. Synthesis of Esters of Acetoxymethylphosphinic Acid

Moscow, Zhurnal Obshchey Khimii, Vol 36, No 4, 1966, pp 699-704

Abstract: Two new methods of producing esters of acetoxymethylphosphinic acid were developed, based on the reaction of hydroxymethylphosphinic acid with trialkylacetoxy-silanes and acetic anhydride, as well as that of esters of trialkoxy-silyloxymethylphosphinic acids with acetic anhydride in the presence of catalytic amounts of sulfuric acid. The reactivity of various silicon-containing derivatives of hydroxymethylphosphinic acid with acetic anhydride was studied, and a scheme of the mechanism of their interaction was proposed. Under the same conditions, the Si-O-C group is quantitatively cleaved, the Si-O-P group is cleaved by only 25.35%, while the C-O-P group remains inert.

Orig. art. has: 2 formulas and 1 table. [JPRS: 37,177]

TOPIC TAGS: organic synthetic process, silane esterification, phosphinic acid
SUB CODE: 07 / SUBM DATE: 21 Mar 65 / ORIG REF: 008 / OTH REF: 004
Card 1/1 DOC: 547.281.1546.185

0923 0774

I 06502-67 EWP(j)/EWT(m) RM
 ACC NR: AP7000475

SOURCE CODE: UR/0079/66/036/006/1075/1078

ORLOV, N. F., MILESHKEVICH, V. P., VAYNBURG, V. M., Leningrad Institute of the
 Textile and Light Industry im. S. M. Kirov (Leningradskiy Institut tekstil'noy
 i legkoy promyshlennosti)

"Synthesis of Organosilicon Esters of Alpha- and Beta-Chloroalkyl(aryl)-
 phosphinic Acids. Interaction of Bis(trialkylsilyl)chloromethyl-
 phosphinates with Sodium Triethylsilanolate"

Moscow, Zhurnal Obshchey Khimii, Vol 36, No 6, 1966, pp 1075-1078

Abstract: A method was developed for producing new bis(trialkylsilyl) esters
 of chloromethyl-, beta-chloroethyl-, and alpha-chlorobenzylphosphinic acids.
 They were synthesized by the reaction of dichlorides of the corresponding
 chloroalkylphosphinic acids with trialkylacetoxysilanes under conditions of
 continuous distillation of the acetylchloride formed. In the reaction of
 sodium triethylsilanolate with bis(trialkylsilyl)chloromethylphosphinates,
 hexaalkyldisiloxanes and sodium salts of trialkylsilyl esters of chloromethyl-
 phosphinic acid were formed. The reaction mechanism deduced indicates a sub-
 stantial sensitivity of the P-O-Si group to the action of nucleophilic agents.
 Orig. art. has: 1 formula and 1 table. [JPRS: 37,023]

TOPIC TAGS: organosilicon compound, ester, organic synthetic process, phosphinic acid

SUB CODE: 07 / SUBM DATE: 06Feb65 / ORIG REF: 003 / OTH REF: 004

Card 1/1 m LE

547.419

0923

1186

L 05184-67 EWT(m)/EWP(j) RM

ACC NR: AP7000740

SOURCE CODE: UR/0079/66/036/005/0892/0895

25
3

ORLOV, N. F., MILESHKEVICH, V. P., Leningrad Institute of the Textile and Light Industry imeni S. M. Kirov (Leningradskiy institut tekstil'noy i legkoy promyshlennosti)

Interaction of Hydroxymethylphosphinic Acid⁷ with Trialkylsilane Hydrides"

Moscow, Zhurnal Obshchey Khimii, Vol 36, No 5, 1966, pp 892-895

Abstract: In a previous communication the authors reported on the synthesis of bis(triorganosilyl) triorganosiloxymethylphosphinates by the reaction of hydroxymethylphosphinic acid with triorganosilane hydrides in the presence of a colloidal nickel catalyst. In a further study of the side reactions, it was found that the reaction of hydroxymethylphosphinic acid with trialkylsilane hydrides results in the formation not only of bis(trialkylsilyl)trialkylsiloxymethylphosphinates, but also O-trialkylsilyl-O-[bis(trialkylsilyl)phosphonemethyl]trialkylsiloxymethylphosphinates, a new class of compounds. A reaction mechanism is proposed. [JPRS: 37,023]

TOPIC TAGS: silane, phosphinic acid, organic synthetic process

SUB CODE: 07 / SUBM DATE: 28Jan65 / ORIG REF: 009 / OTH REF: 001

Card 1/1 vmb

UDC: 547.245 + 547.241
0923-1909

TOPIC TAGS: phosphorous acid, silane

SUB CODE: 07 / SUBM DATE: 20Dec65 / ORIG REF: 003

APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

Card 1/1 mL

UDC: 547.245 + 547.241

ACC NR: AP6035688 (A,N) SOURCE CODE: UR/0413/66/000/019/0032/0032

INVENTOR: Orlov, N. F.; Viktorov, O. F.

ORG: none

TITLE: Preparation of dialkyl-1-(3-chloro-2-hydroxypropyl) hydroxymethyl phosphinates. Class 12, No. 186474 [announced by Leningrad Institute of the Textile and Light Industry im. S. M. Kirov (Leningradskiy institut tekstil'noy i legkoy promyshlennosti)]

SOURCE: Izobreteniya, promyshlennyye obraztsy, tovarnyye znaki, no. 19, 1966, 32

TOPIC TAGS: ^{organic} ~~organic~~ phosphorus compound, phosphinic acid, ester

ABSTRACT: In the proposed method, dialkyl-1-(3-chloro-2-hydroxypropyl)-hydroxymethyl phosphinates are obtained by the reaction of HCl with dialkylglycidylhydroxymethyl phosphinates.

[WA-50; CBE No. 14] [PS]

SUB CODE: 07/ SUEM DATE: 23Nov65

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UDC: 547.26'118.07

ACC NR: AP7002970

(A)

SOURCE CODE: UR/0413/66/000/024/0050/0050

INVENTOR: Orlov, N. F.; Vol'f, L. A.; Androsova, M. V.; Kirilenko, Yu. K.

ORG: none

TITLE: Preparative method for poly(vinyl alcohol)-based fireproof fibers, films or fabrics. Class 29, No. 189515

SOURCE: Izobreteniya, promyshlennyye obraztsy, tovarnyye znaki, no. 24, 1966, 50

TOPIC TAGS: polyvinyl alcohol, fire resistant material, polymer

ABSTRACT: An Author Certificate has been issued for a method of preparing poly(vinyl alcohol)-based fireproof fibers, films or fabrics. The method involves treatment of dehydrated fibers, films or fabrics with dialkyl hydrogen phosphites.

SUB CODE: 11/ SUBM DATE: 12Jul65/ ATD PRESS: 5112

Card 1/1

UDC: 677.494.744.72:66.093.6

ORLOV, N.G.

The BSh-25/100-type walking excavator. Biul.tekh.-ekon.inform.
no.12:37-39 '59. (MIRA 13:4)
(Excavating machinery)

ORLOV, N.G., inzh.

New development in the technology of manufacturing welded rails.
Zhel.dor.transp. 44 no.7:45-48 JI '62. (MIRA 15:8)
(Railroads--Rails)

GLAZMAN, M.G.; ORLOV, N.G., kandidat meditsinskikh nauk, glavnyy vrach.

Study of stable strains of staphylococci; author's abstract. Zmr.mikro-
biol.epid.i immun. no.2:53-54 F '53. (MLRA 6:5)

1. Bol'nitsa imeni Baumana.

(Staphylococcus)

ORLOV, N.G., kandidat meditsinskikh nauk

Work organization of district general practitioners in a consolidated city hospital. Sov.zdrev. 15 no.5:33-37 S-O '56. (MLRA 10:1)

1. Glavnyy vrach Ob'yedinennoy bol'nitsy imeni N.E.Baumana v Moskve.
(HOSPITAL ADMINISTRATION
in Russia, work organiz. of GP's in consolidated city
hospital)

ORLOV, N. G.

How functions the United Bauman's Hospital in Moscow. Czech.
sdravot. 5 no.1:1-6 Jan 57.

1. Kandidat lékařských ved, reditel Baumanovy nemocnice v
Moskve.

(HOSPITALS

United Bauman's Hosp., Moscow (Cs))

~~ORLOV, E.G.~~

Information service in municipal hospitals and polyclinics. Med.
sestra 16 no.5:7-10 May '57. (MLRA 10:7)

1. Is Gorodskoy bol'nitsy imeni Baumana, Moskva.
(HOSPITALS--MANAGEMENT AND REGULATION)

ORLOV, N.G., kand.med.nauk (Moskva)

Organizing epidemic control work in a polyclinic servicing a specific
area. Zdrav.Ros.Feder. 2 no.2:19-21 Y '58. (MIRA 11:3)
(COMMUNICABLE DISEASES)

~~ORLOV, N.G.~~

Two-stage treatment of patients in hospitals. Zdrav.Ros.Feder.
2 no.7:3-12 J1'58 (MIRA 11:7)

1. Olovnyy vrach bol'nitsy im. Baumana, Moskva.
(HOSPITALS)

ORLOV, N.G. (Moskva)

Experience in expert temporary disability evaluation in the
polyclinic. Sov. zdrav. 19 no. 4:38-42 '60. (MIRA 13:10)

1. Iz bol'nitsy imeni Baumana.
(DISABILITY EVALUATION)

ORLOV, N.G.

Daily dispensary. Zdrav. Ros. Feder. 5 no.5:3-5 My '61.

(MIRA 14:5)

1. Glavnyy vrach bol'nitsy imeni N.E. Baumana, Moskva.
(MOSCOWS--HOSPITALS--OUTPATIENT SERVICES)

ORLOV, N.G. (Moskva)

Eliminating lines in registering at the polyclinic. Sov. zdrav. 26
no.9:30-34, '61. (MIA 14:12)

1. Glavnyy vrach bol'nitsy imeni Baumana.
(MEDICAL CARE)

ORLOV, N.G. (Moskva)

Reorganization of the work of a polyclinic under new conditions. Sov. zdrav. 20 no.12:52-56 '61. (MIRA 15:6)

1. Glavnyy vrach Bol'nitsy imeni N.E. Baumana.
(HOSPITALS--ADMINISTRATION)

ORLOV, N.G.

A day clinic for peptic ulcer patients. Sov.sdrav. 21 no.8:10-13
'62. (MIRA 15:11)

1. Glavnyy vrach bol'nitsy imeni N.E.Baumana, Moskva.
(PEPTIC ULCER) (HOSPITALS—OUTPATIENT SERVICES)

URT'YEV, Viktor Petrovich; LUR'YE, Vitol'd Samar'yevich; ISAYEV,
Al'bert Semenovich; ORLOV Nikolay Il'ich; TSYPLUKHIN, Petr
Gavrilovich; SOKOLOV, A.N., red.; SHILLING, V.A., red.izd-va;
BELOGUROVA, I.A., tekhn. red.

[Vacuum arc furnace] Dugovaia vakuunnaia pech'. Leningrad, 1962.
25 p. (Leningradskii dom nauchno-tekhnicheskoi propagandy. Ob-
men peredovym opytom. Seria: Liteinoe proizvodstvo, no.5)

(MIRA 16:2)

(Electric furnaces) (Vacuum metallurgy)

Орлов, Н. И.

86-11-26/31

AUTHOR: Orlov, N. I., Col

TITLE: The Practice Bomb Should be Modified (Prakticheskuyu bombu sleduyet modernizirovat')

PERIODICAL: Vestnik Vozdushnogo Flota, 1957, Nr 11, p. 84 (USSR)

ABSTRACT: The author states that the practice bomb used at the present time for training navigators is inadequate, because with the aid of those optical devices and radar with which the bombing grounds are equipped, it is not always possible to obtain the precise fixes of the impacts of practice bombs. The author suggests that the present practice bomb be modified, or a new bomb designed. The author gives some details concerning the design of such a bomb.

AVAILABLE: Library of Congress

Card 1/1

25025

S/057/61/031/007/006/021
B108/B2099,4120AUTHORS: Golant, V. Ye., Orlov, N. I., Pakhomov, L. P.

TITLE: Production of a high-density plasma by a hot-cathode discharge in a magnetic field

PERIODICAL: Zhurnal tekhnicheskoy fiziki, v. 31, no. 7, 1961, 797-801

TEXT: The authors present the results of an investigation of a hot-cathode discharge in a magnetic field. In such a discharge with a current density of over 3 a/cm^2 in an inhomogeneous magnetic field ($H/H_c > 3 \div 5$; H - magnetic field strength in the region under examination, H_c - magnetic field strength near the cathode) a plasma concentration of over 10^{14} cm^{-3} may be attained theoretically. It was the aim of the present investigation to determine the concentration of charged particles in such a plasma. The emitting area on the loop- or ring-shaped cathode was 1.5 cm^2 , its temperature over 2000°C . The distance between the tungsten anode and cathode was 25 cm. The measurements were made in both a homogeneous and an inhomogeneous mag-

Card 1/3

ORLOV, N. I.

Magnetic ore deposits and snow cover. Priroda 52 no.1:114 '63.
(MIRA 16:1)

1. Institut geografii AN SSSR, Moskva.

(Kursk Magnetic Anomaly—Snow)

Orlov, N.I. (Moskva)

Snowstorms. Priroda 52 no.2:127-128 '63.
(Blizzards)

(MIRA 16:2)

~~ORLOV~~, N.I., glavnyy veterinarnyy vrach Bugul'minskogo rayona, Tatarskoy
SSR, KARAYSHCHIKIN, V.P.

Prophylactic effect of hemospirikin. Veterinariia 30 no.11:58 № '53.
(MLRA 6:11)

1. Glavnyy fel'dsher tsentral'nogo sovetuchastka, Bugul'minskogo
rayona (for Karayshechkin).

GROYSMAN, B.A., ORLOV, I.I.

Actinomycosis of the mammary gland. *Zdravookhranenie* 4, no. 3:
57 My-Je'61. (MIRA 16.7)

1. Iz khirurgicheskogo otdeleniya (zav. M. I. Voynova) rayonnoy
bol'nitsy g. Orgeyeva (glavnyy vrach M. A. Bagmanyant).
(ACTINOMYCOSIS) (MAMMARY GLANDS)

1ST AND 2ND COLUMNS PROCESSES AND PROPERTIES INDEX 3RD AND 4TH COLUMNS

CA *11F*

Color tests for vitamin A. N. I. Orlow. *Izv. Vsesoyuzn. Nauch.-Issledovatel. Inst. Pishchev. Khim. Prom.* Sep. 1952, No. 5, 26 pp.—The Bermanov test for vitamin A is not so. very quant. and even as a preliminary qual. test is subject to error. The H₂SO₄ test is not sensitive enough for colorimetric estn. The SbCl₅ and AsCl₃ tests, as prescribed in the Pharmacopoeia, are convenient for approx. evaluation of fish-liver oils, and may be relied on when a Lovibond tintometer is not available.

Julian F. Smith

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1ST AND 2ND ORDERS PROCESSES AND PROPERTIES INDEX 3RD AND 4TH ORDERS

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Zinc in food products and its sanitary-hygienic significance. V. P. Zolova and N. I. Golov. *Voprosy Pitaniya* 4, No. 5, 40-55(1935). Analyses are given for food products where zinc poisoning was discovered, and the source of the zinc is indicated. Zn contents in such cases varied from 400 to 7000 mg. per kg. of product. Mushrooms are especially rich in zinc. Data on about 50 species of mushrooms show variations in Zn content from 50 to 200 mg. per kg. of dry wt. of plant material.

F. H. Mathmann

COMMON ELEMENTS

OPEN

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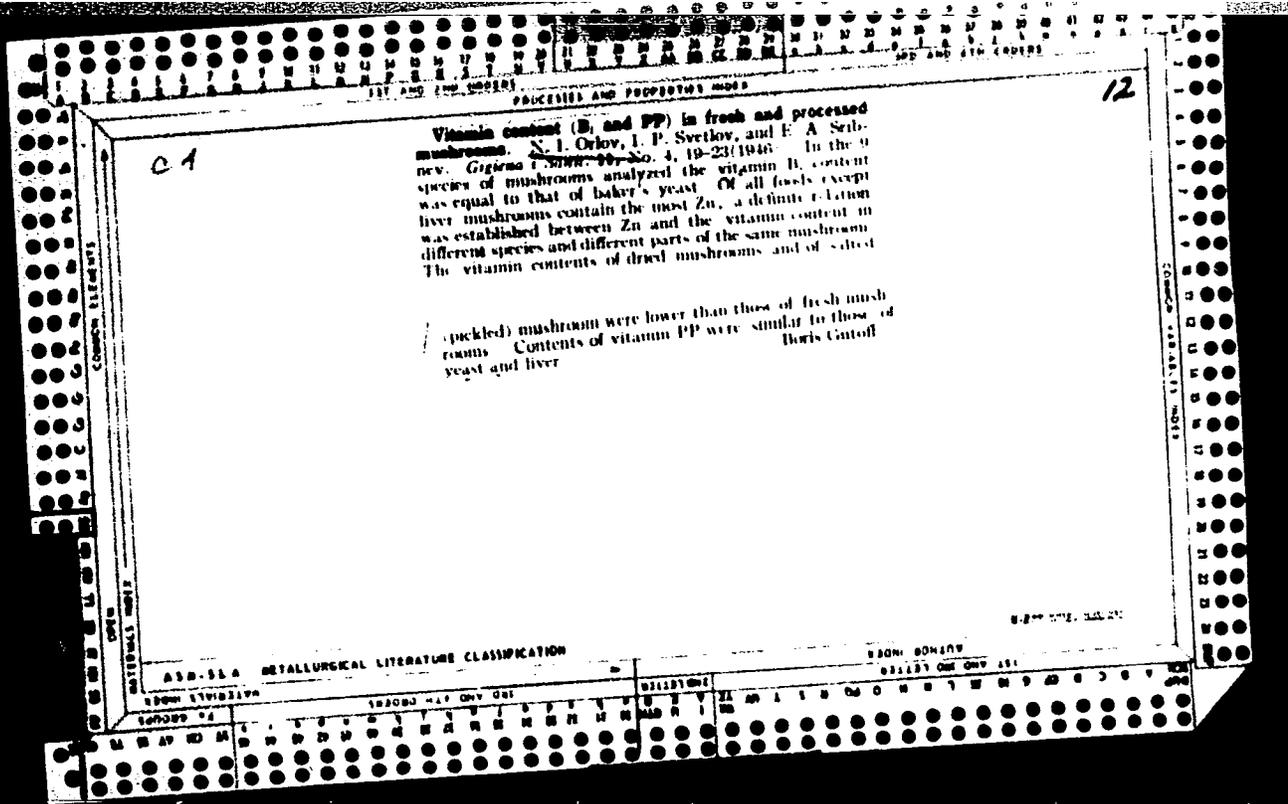
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USSR/Medicine - Food Poisoning

"Review of Monograph Food Poisoning and Its Prophylaxis," by N.I. Orlov, Library of the Practicing Physician, Medgiz, 1952, 119 pp, 25,000 copies," (Yu. A. Kozlov, reviewer)

Gig i San, No 11, pp 60, 61

Lauds the clear and concise style of the author and the up-to-date presentation of information on food poisoning of bacterial and nonbacterial origin. Objects to the brief treatment of Salmonella organisms and of their pathogenicity in producing toxic infections. Considers that too little space is allocated to a discussion of Sonne Dysentery bacilli as Originators of toxic food infections, and does not quite agree with the author's treatment of the symptomatology and pathogenesis of toxic infections caused by Sonne bacilli.

Source #264T35

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RA1242.M907

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AID P - 2168

Subject : USSR/Medicine

Card 1/2 Pub. 37 - 10/22

Authors : Belostotskaya, Ye. M., Beryushev, K. G., Kands. of Med. Sci., Orlov, N. I., Dr. of Med. Sci., Fcngauz, M. I., Kand. of Med. Sci., and Cherkinskiy, S. N., Doc. of Med. Sci.

Title : From the practical work of the Scientific Research Sanitary Institute im. Erisman in the introduction of physiological methods in investigations of hygiene

Periodical : Gig. i san., 4, 40-43, Ap 1955

Abstract : The purpose of this article is to explain the work of the Institute in the light of I. P. Pavlov's theories and his analytical approach to observed phenomena. The reactions of the organism are studied in relation to the changes in its environment, climatic, atmospheric, industrial conditions, etc. The article is illustrated by many examples, observations of human beings and tests performed on animals. 10 Russian references (1951-1954).

AID P - 2168

Gig. 1 san., 4, 40-43, Ap 1955

Card 2/2 Pub. 37 - 10/22

Institution : Scientific Research Sanitary Institute im. Erisman

Submitted : My 10, 1954

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