

Klimatologiya

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KOSTIN, S.I., doktor geograficheskikh nauk, professor.

Some remarks on the problem of the development of climatology.

Meteor. i gidrol. no. 5:9-12 My '53.

(MLBA 8:9)

1. Gosudarstvennyy universitet, Voronezh.
(Climatology)

KOSTIN, S.I.

Climatic regions of the central Chernozem provinces. Vop.geog.
no.32:189-193 '53. (MIRA 10:11)
(Central Black Earth Region--Climate)

KOSTIN, S. I.

The Committee on Stalin Prizes (of the Council of Ministers USSR) in the fields of science and inventions announces that the following scientific works, popular scientific books, and textbooks have been submitted for competition for Stalin Prizes for the years 1952 and 1953. (Sovetskaya Kultura, Moscow, No. 22-40, 20 Feb - 3 Apr 1954)

<u>Name</u>	<u>Title of Work</u>	<u>Nominated by</u>
Kostin, S. I. Podrovskaya, T. V.	"Climatology"	Main Geophysics Observatory imeni A. I. Voyeykov

80: W-30604, 7 July 1954

KOSTIN, S. I.

"The Influence of Forest on Climate".
Trudy in-ta Lesa AN SSSR, 22, pp 129-134, 1954.

The influence of forest on climate is determined first of all by its action upon the microclimatic conditions under the forest cover, which mainly depends upon the geographical location of the forest mass, composition, growth, completeness of planting, etc. The author considers the influence of continuous forest masses upon the climatic conditions in a forest steppe region (Voronezhskaya Oblast) in accordance with data of observations (with use of automatic recorders) in an oak forest and at a distance of 500 meters from the outskirts. On bright summer days the forest canopy admits 10-12% of the radiation. The forest floor litter possesses the least heat conduction, less than the bare soil; therefore during the day it is heated more and during the night it is cooled more. (RZhGeol, No 11, 1955)

SO: Sum No 884, 9 Apr 1956

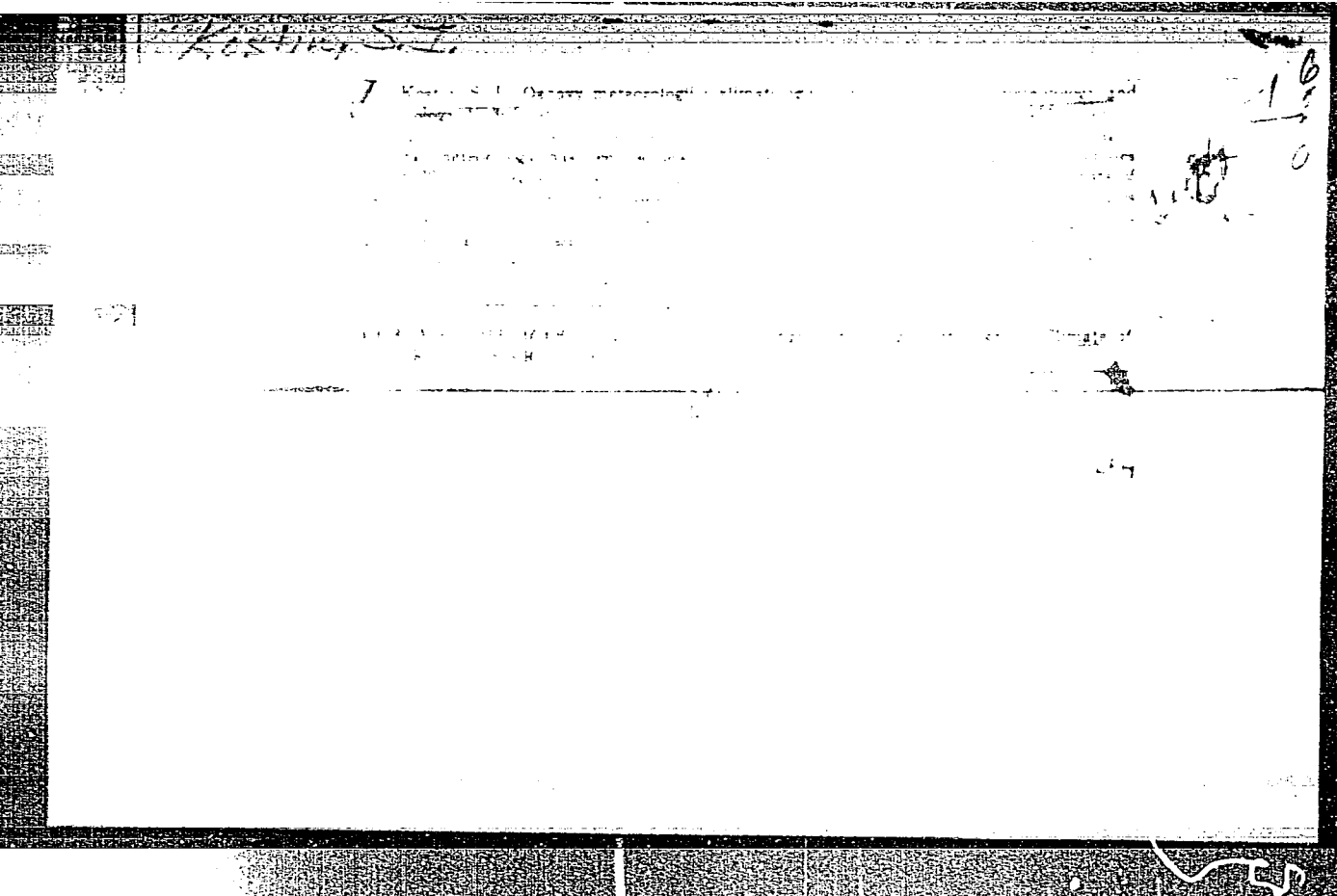
KOSTIN, S. I.

"Climatic Zones and Regions of the Central and Eastern Forest Steppe and of the Russian Lowland Steppe".

Trudy Voronezhsk. un-ta, 30, pp 3-21, 1954.

A climatic regional classification of the central and eastern regions of the southern half of the European part of the USSR is given. At the basis of a distinction of four climatic zones lies the following index of local moisture: $K = P/E$, where P is the amount of precipitation during a given period and E is the conditional index of evaporability for the same period, as calculated from the formula $E = (dn/4) (1 + 0.004t)^2$ (d is the mean deficit of moisture for the period, n is the number of days in the period, and t is the mean temperature for this same period). Indexes of moisture which were computed by this method for the period May-September were represented by isolines on a map. (RZhGeol, No 7, 1955)

SO: Sum No 884, 9 Apr 1956



PHASE I BOOK EXPLOITATION 997

Kostin, Sergey Iosifovich, Doctor of Geographical Sciences

Osnovy meteorologii i klimatologii (Principles of Meteorology and Climatology)
4th ed., rev. and enl. Leningrad, Gidrometeoizdat, 1958. 403 p.
10,000 copies printed.

Ed.: Yasnogorodskaya, M.M.; Tech. Ed.: Soloveychik, A.A.

PURPOSE: This is a textbook for students of universities, especially of forestry institutes. The book may be of interest to practical meteorologists and climatologists and to technical personnel employed in agriculture and silviculture.

COVERAGE: The book describes the principal physical phenomena of the atmosphere, particularly the processes effecting the growth of vegetation. In addition, the book analyzes the impact of vegetation on meteorological factors and climatic conditions, discusses plant protection from frosts and dry winds, and finally outlines the main climatic zones of the world in general, and of the USSR in particular. The author acknowledges the assistance of A.I. Rudenko, Yu.D. Yanishevskiy, and the Chinese scientists Van-Pen-Fei who translates the book into Chinese. There are 29 Soviet references, 143 figures (including maps), and 9 tables in the appendix.

Card 1/12

KOSTIN, S.I.

Solar activity and its influence on the growth of trees and the state of forest stands in the central steppe of the Russian Plain. Trudy GGO no.111:108-117 '61. (MIRA 15:1)
(Voronezh Province--Forests and forestry)
(Meteorology, Agricultural) (Growth (Plants))

BR

PHASE I BOOK EXPLOITATION

SOV/5941

Kostin, Sergey Iosifovich, and Taisiya Vasil'yevna Pokrovskaya

Klimatologiya (Climatology) 2d ed., rev. and enl. Leningrad, Gidrometeoizdat, 1961. 485 p. Errata slip inserted. 5000 copies printed.

Resp. Ed.: O.A. Drozdov; Ed.: L.P. Zhdanova; Tech. Eds.: A.A. Soloveychik and M.I. Braynina.

PURPOSE: This book is intended for students of climatology, meteorology, and hydrometeorology.

COVERAGE: The present volume is a revised edition of a handbook on climatology which first appeared in 1953. Part I (except for Ch. III and sections 35 and 39) was revised by its author, S.I. Kostin; and Part II and Ch. III and sections 35 and 39 of Part I were revised by T.V. Pokrovskaya, the original writer of Part II. The following are discussed: principles, scope, and application of climatology and climatography; general atmospheric circulation; the role of radiation and circulation in the formation of climate; effect of underlying surfaces (land, water, relief, snow, vegetation) and human activity on microclimate; individual climatic elements (temperature, humidity, precipitation, cloud cover, moisture

Card 1/4

Climatology

SOV/5941

and heat balance, etc.); climatic changes and variations; principles and methods of processing climatological observations and data; and the application of climatology to agriculture, industry, commerce, transportation, communications, construction, and city planning. A large portion of the book is devoted to a detailed and thorough description and analysis of climate in different parts of the world, with particular attention given to the climate of the USSR. Several systems of climate classification, of which L.S. Berg's and B.P. Alisov's are considered the most important, are discussed. The historical development of the science of climatology in the USSR is briefly outlined. The following participated in reviewing the present volume: V. Yu. Milevskiy, V.M. Mikhel', S.A. Sapozhnikova, I.A. Gol'tsberg, R.F. Usmanov, T.G. Beryland, L.I. Zubenok, and K.M. Kommatnaya. There are 25 references, all Soviet.

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Card 2/4	

KOSTIN, S.I., prof.

Does Brikner's cycle exist? Priroda 54 no.2:78-79 F '65.

(MIRA 18:10)

1. Voronezhskiy lesotekhnicheskij institut.

KOSTIN, S.I.

Climatic fluctuation on the East European Plain in the historical
epoch. Trudy GGO no. 181:56-74 '65.

(MIRA 18:10)

KOSTIN, S.M., inzh.- kapitan 2-go ranga; MIRIN, J.M., kapt. voyenno-morskikh nauk, kapitan 2-go ranga

Statistical study of the efficiency of tactical maneuvers. Mor.
sbor. 48 no.12:15-20 D '64. (MIRA 18:2)

S/169/62/000/009/075/120
D228/D307

AUTHOR: Kostin, S. N.

TITLE: Radiation and heat balance on the territory of the central black-earth regions

PERIODICAL: Referativnyy zhurnal, Geofizika, no. 9, 1962, 20, abstract 9B121 (Nauchn. Zap. Voronezhsk. lesotekhn. in-ta, 24, 1961, 57-65)

TEXT: The components of the underlying surface's radiation and heat balance were calculated for the territory of the central black-earth regions. The summary radiation $(Q + q)_n$ was computed from the empirical formula: $(Q + q)_n = (Q + q)_0 [1 - (1 - k)_n]$. Here $(Q + q)_0$ is the summary radiation when the sky is clear; n is the average cloudiness in parts of a unit; and k is the factor for the territory under consideration, taken as being equal to 0.36 - 0.37. The values of n were determined from the data of observations over

Card 1/3

Radiation and heat ...

S/169/62/000/009/075/120
D228/D307

a 20-year period. The calculated values of $(Q + q)_n$ were compared with the actual ones for Voronezh in 1936-1938. The divergences between the calculated and observed values constituted 3 - 12% for separate seasons and an average of 0.4% for the year. It is hence concluded that the adopted method of calculation is sufficiently precise for determining the yearly magnitudes of $(Q + q)_n$. The va-

lues of the underlying surface's albedo were established with allowance for the fact that it changes during the year. The magnitudes of absorbed radiation were derived. The effective radiation was ascertained from the formula: $I = I_0(1 - cn^2)$; where I_0 is the effective radiation when the sky is clear, n is the cloudiness index, and c is a coefficient. I_0 was found from the table in M. Ye.

Berlyand's work (Predskazaniye i regulirovaniye teplogo rezhima prizemnogo sloya atmosfery (Predicting and controlling the heat regime of the atmosphere's near-ground layer), Gidrometeoizdat, 1956); n was obtained from the data of observations over a 20-year period. A value of 0.72 - 0.73 was taken for c . Evaporation from

Card 2/3

APPROVED FOR RELEASE: 06/14/2000

CIA-RDP86-00513R000825220006-6

Radiation and heat ...

S/169/62/000/009/075/120
D228/D307

the ground surface was determined from V. B. Polyakov's graphs (Gidrologicheskiy analiz i raschety (Hydrologic analysis and calculations), Gidrometeoizdat, 1946). The calculation's results were compared with the data of 3-year measurements by a ГИИ-500 (GII-500) evaporator, set up at Voronezh, under conditions of vegetationless ground. The divergences between the calculated and observed data amount to 6% for the period May - September. The turbulent flow of heat for the year was determined as a residual term of the heat balance, the ground heat flow in this case being taken as equal to zero. A chart of the annual values of absorbed radiation, the radiation balance, and the turbulent heat flow is presented for the territory under consideration. The results obtained are discussed.

Abstracter's note: Complete translation. 7

Card 3/3

KOSTIN, V.

American military satellites. Av. i kosm. 45 no. 10:94-95 '62.
(MIRA 15:10)

(United States—Artificial satellites)

EKONOMOV, Lev Arkad'yevich; KOZMIN, V., nauchn. red.;
FEDCHENKO, V., red.

[Commanders of fiery arrows; word about rocket engineers and rockets] Poveliteli ognennykh strel; slovo o raketnikakh i raketakh. Moskva, Molodaia gvardiia, 1964. 318 p. (FIRA 18:1)

KOSTIN, V.

How to sell clothes. Sov. tog. 35 no.6:19-22 Je '62. (MIRA 15:7)
(Clothing industry)

KOSTIN, V. SHLEMIN, I.

Equipment for transporting ready-made clothing. Sov.torg. no.6:
59-60 Je '57. (MLRA 10:8)
(Clothing and dress--Transportation)

~~KOSTIN, V.~~

Permanent exhibition in Ivanovo. Pozh.delo 3 no.3:26 Mr '57.,

1. Zamestitel' nachal'nika upravleniya posharnoy okhrany Ivanovskoy oblasti. Pozh.delo 3 no.3:26 Mr '57. (MIRA 10:4)

(Ivanovo Province--Fire prevention--Exhibitions)

YAMPOL'SKIY, Boris Borisovich; KOSTIN, V., red.; KLIMOVA, T., tekhn.
red.

[Time is our ally] Nash drug - vremia. Moskva, Gos. izd-vo polit.
lit-ry, 1961. 53 p. (MIRA 14:10)
(Russia--Economic conditions) (United States--Economic conditions)

97 - 1 - 10/10

AUTHOR: Kostin, V.A., Engineer

TITLE: Joining Secondary Beams with Main Beams in Floor Constructions.
(Sopryazheniye vtorstepennykh balok s glavnoy v rebristom perekrytii).

PERIODICAL: Beton i zhelezobeton, 1957, No. 1, pp. 32-33. (U.S.S.R.)

ABSTRACT: A type of joint between a secondary and the main reinforced concrete beam as illustrated in the article is evaluated. This joint is made with the aid of steel stirrups and plates concreted to the main beam and welded together in the form of an anchor. The end-reinforcement of the secondary precast beam is welded to a steel collar. The top reinforcement of the secondary beam passes through the collar and is welded to the opposite number across the main beam by butt welding. Finally, all the steel components are welded together and the connections are grouted with expanding cement. The steel parts on the surface are covered with an expanded mesh and rendered with cement grout. The reinforced concrete slab has a thickness of 4 - 7 cm. The UHMNC recommends the above floor construction but advocates savings in steel to

Card 1/2

TITLE: Joining Secondary Beams with Main Beams in Floor Constructions.
(Sopryazheniye vtorstepennykh balok s glavnoy v rebristom perekrytii).

be achieved and also the positioning of the secondary beam at least 5 cm above the main beam.

There is 1 drawing.

ASSOCIATION: ---

PRESENTED BY: ---

SUBMITTED: ---

AVAILABLE: Library of Congress

Card 2/2

KOSTIN, V.

"The Soviet worker and automation" by A.Osipov, I. Kovalenko,
E.Petrov. Reviewed by V.Kostin. NTO 2 no.9:55 S '60.

(MIRA 13:9)

(Osipov, A.) (Automation) (Kovalenko, I.) (Petrov, E.)

KOSTIN, V.

New developments in Sverdlovsk trade. Sov. terg. 36
no. 1:25-28 Ja '63. (MIRA 1612)
(Sverdlovsk--Retail trade)

FROLOV, S.; KOSTIN, V.; VARVARIN, N.

Production organization and the establishing technical
standards. Sots. trud 8 no.1:89-91 Ja '63. (MIRA 16:2)
(Machinery industry--Production standards)

DVOYCHENKOVA, Yu.; KOSTIN, V.

Conversion of the "Krasnoe Sormovo" plant to the seven-hour
working day. *Bul.nauch.inform.; trud i zar.plata no.5:28-31*
'59. (MIRA 12:6)

(Gorkiy--Metallurgical plants)
(Industrial efficiency).

KOSTIN, Y.A., inzh.; VARVARIN, N.M., inzh.; RKKUSHIN, A.N., inzh.

Reduction of labor necessary for shipbuilding at the "Krasnoe
Sormovo" Shipyard. Sudostroenie 25 no.1:69-71 Ja '59.(MIRA 12:3)
(Gorkiy Province--Shipbuilding)

KOSTIN, V.B., kand.tekhn.nauk

Performance of elastic nitrile rubber coatings. Tekst.prom. 25
no.1:31-33 Ja '65. (MIRA 18:4)

1. Glavnyy inzh. Tashkentskogo tekstil'nogo kombinata.

VETROV, V.D.; KOSTIN, V., red.; DANILINA, A., tekhn. red.

[When you love your work....] Kogda liubish' svoe delo. Mo-
skva, Gospolitizdat, 1962. 94 p. (MIRA 15:6)
(Labor and laboring classes)

REYFMAN, V.G.; KOSTIN, V.D.

Concentration of virus X in potato leaves during the growing
period. Soob.DVFAN SSSR no.13:65-72 '60. (MIRA 14:3)

1. Dal'nevostochnyy filial im. V.L.Komarova Sibirskogo otdeleniya
AN SSSR.
(Potatoes—Diseases and pests) (Virus diseases of plants)

KOSTIN, V.D.; KRYLOV, A.V.; IVLIYEV, L.A.

Leaf beetle *Galeruca tanaceti* L., a potential potato pest in
Kamchatka. Soob. DVFAN SSSR no.17:97-99 '63.

(MIRA 17:9)

1. Dal'nevostochnyy filial im. V.L. Komarova Sibirskogo otdeleniya
AN SSSR.

TETERUK, G.I.; ZAVYAZKIN, P.G.; ALIYEV, T.M.; ALIYEV, A.G.; MELIK-SHAKHNAZAROV,
A.M.; ARULIS, B.K.; BARTENEV, G.M.; YEL'KIN, A.I.; KOSTIN, V.I.;
KHARKHARDIN, S.I.; SERGEYEV, A.I.; VARTANOV, S.Kh.; PRIMANCHUK, L.I.;
MOLODTSOV, A.A.; SHMELEV, N.V.; ROVINSKIY, M.I.; ABRAMOV, H.N.;
YEROFEYEV, L.V.; RYAKHIN, V.A.; ZELENIN, A.N.; BERKMAN, I.I.

Patent certificates for Soviet inventions. Stroi. truboprov. 9 no.5:
35-36 My '64. (MIRA 17:9)

KOSTIN, V. I.

Mathematical Reviews
Vol. 14 No. 11
December, 1953
Geometry.

↳ Kostin, V. I. Some questions of the metric duality of the geometry of Riemann. *Trudy Sem. Vektor. Tenzor. Analizu* 6, 444-446 (1948). (Russian)

This is an abstract of a paper that failed to be included in the fifth volume of the above publication. The paper is concerned with the gonometric (angular) invariants of an m -parameter family of hyper-surfaces of an n -dimensional Riemannian space. If $u(x^i, \xi^k) = 0$, $i = 1, 2, \dots, n$; $k = 1, 2, \dots, m$, is the equation of the family and $ds^2 = g_{\alpha\beta} dx^\alpha dx^\beta$ is the metric, then the angular metric is $d\theta^2 = \Omega_{\gamma\delta}(x, \xi) d\xi^\gamma d\xi^\delta$ where Ω is expressible in terms of $g^{\alpha\beta}$ and the partial derivatives of u with respect to x are ξ .

For two nearby hypersurfaces one must have $u = 0$ and $(\partial u / \partial \xi^\alpha) d\xi^\alpha = 0$. If, because of these two equations, $\Omega_{\gamma\delta}$ reduce to functions of ξ alone, the angular metric is then Riemannian and is in a sense the dual of ds^2 . Writing $u = -x^n + f(x^1, \dots, x^{n-1}, \xi) = 0$, the necessary and sufficient conditions that f must satisfy in order to define a gonometric family is a set of first-order partial differential equations of the second degree, each involving only one pair of parameters. The concluding remarks deal with some integral invariants of the family in case of n -, $(n-1)$ - and 1-parameter families. *M. S. Knebelman (Pullman, Wash.)*

KOSTIN, V. I.

USSR/Minerals
Lignites
Coal Gas

Apr 49

"Experimental Gasification of Karaganda Lignites"; V. I. Yerminev, V. I. Kostin,
Engineers, 6 pp

"Za Ekonomiyu Topliva" Vol VI, No 4

Karaganda lignite is a satisfactory gas-generating fuel. During prolonged storage, Karaganda lignite rapidly breaks up into bits of unsuitable for gasification. Higher indexes were obtained in gasification of Karaganda lignite than in gasification of Chelyabinsk coal. One deficiency of Karaganda coal, as a gas-generating fuel, is the small difference between the temperature of the start of ash deformation and the melting point (20°C).

PA 42/49T77

KOSTIN, V.I., tekhnik

Rewinding of electromagnets to operate on different current.
Energetik 11 no.10:37-38 0 '63. (MIRA 16:11)

KOSTIN, V.I.

LIOZNYANSKAYA, S.G.; KOSTIN, V.I.; VORONIN, P.V.

Accelerated cooling of the glass ribbon in the stack of a vertical
drawing machine. Stek.l ker.l3 no.11:9-10 N '56. (MIRA 10:1)
(Glass manufacture)

TAYTS, N.Yu., doktor tekhn.nauk; KOLESNIK, B.P., kand.tekhn.nauk;
YANKOVSKIY, V.M., kand.tekhn.nauk; KADINCOVA, A.S., inzh.;
KAUFMAN, M.M., inzh.; Prinsipalni uchastiyu: POLYAKOVA, N.K.,
inzh.; VOVSINA, A.D., inzh.; SHANINA, A.S., inzh.; KOSTIN, V.I., inzh.

Rapid heat treatment of drill pipes. Stal' 22 no.1:57-60 Ja '62.
(MIRA 14:12)

1. Ukrainskiy nauchno-issledovatel'skiy trubnyy institut (for
Polyakova).

(Pipe, Steel)
(Steel—Heat treatment)

15(2)

AUTHORS: Bronshteyn, Z. I., Kostin, V. I.

SOV/72-59-5-4/23

TITLE: High-efficiency Gluing Method of Curved Windshields (Vysoko-proizvoditel'nyy sposob skleyki gnutykh avtomobil'nykh stekol)

PERIODICAL: Steklo i keramika, 1959, Nr 5, pp 11 - 13 (USSR)

ABSTRACT: The Institut stekla (Glass Institute) worked out the technical method mentioned in the title by tests with the moderate panorama windshields of the car of the type "Volga". The glass packets glued together are rolled on a special apparatus (Fig 1). Figure 2 shows the various rolling cylinders. The tubes ZS-3 with a capacity of 500 w and a mirror reflector are used for heating the glass packets. Figure 3 shows the heat curves. This method makes it possible to reduce the duration of gluing and pressing the glasses to a small amount of the production time as far necessary. There are 3 figures.

Card 1/1

KOSTIN, V. I.

O gipertolicheskoy geometrii kelikleyna. Gor'kiy, Uchen. Zap. un-ta, 12 (1939), 183-195,

SO: Mathematics in the USSR, 1917-1947

Edited by Kurosh, A. G.,

Markusevich, A. I.

Rashevskiy, P. K.

Moscow-Leningrad, 1948

ACC NR: AT6036568

SOURCE CODE: UR/0000/66/000/000/0177/0178

AUTHOR: Zinov'yeva, I. P.; Rubashkina, L. A.; Kostin, V. K. 29

ORG: none

TITLE: Blood transaminase as an index of human tolerance to landing impact accelerations [Paper presented at the Conference on Problems of Space Medicine held in Moscow from 24 to 27 May 1966]

SOURCE: Konferentsiya po problemam kosmicheskoy meditsiny, 1966. Problemy kosmicheskoy meditsiny. (Problems of space medicine); materialy konferentsii, Moscow, 1966, 177-178

TOPIC TAGS: space medicine, space physiology, blood chemistry, impact acceleration, biologic acceleration effect, human physiology

ABSTRACT: A "tolerance criterion" of impact acceleration is blood transaminase activity. The literature shows that the activity of these enzymes increases as a result of injury to tissues.

In this study, a change in the activity of glutamic-alanine, and glutamic-aspartic transaminase was studied. Human subjects were exposed to landing accelerations with various buildup rates (2500-6000 G/sec and higher) acting along the longitudinal or transverse axes of the body in 32 experiments.

It was found that large accelerations (32-42 G) with a buildup rate of
Card 1/2

I 10949-67 -

ACC NR: AT6036568

APPROVED FOR RELEASE: 06/14/2000

CIA-RDP86-00513R000825220006-6

2500 G/sec (chest-back) and 22-25 G accelerations (head-pelvis) with a buildup rate of 2500-6000 G/sec (7.5 m/sec landing velocity) increased transaminase activity by a factor of 1.5-2.0. The activity of these enzymes was occasionally accompanied by fresh erythrocytes in the urine. At lower magnitudes, enzyme activity was not altered. The highest indices of transaminase activity occurred 24 hr after exposure to acceleration. After 4-5 days, these indices returned to normal. In the majority of cases, the increase in transaminase activity, which reflected disrupted tissue integrity, preceded other functional changes and indicated that acceleration which could be well tolerated. These tests showed that the activity of transaminase enzymes during impact accelerations is a sufficiently sensitive index by which to judge the reaction of the organism to an applied stress. [W.A. No. 22; ATD Report 66-116]

SUB CODE: 06 / SUBM DATE: 00May66

Card 2/2 b/p

ACSTIN, V. M.

AID P - 3975

Subject : USSR/Mining
Card 1/1 Pub. 78 - 20/27
Author : Kostin, V. M., Drilling Foreman of the Chapayev
Drilling Bureau of the United Kuybyshev Petroleum
Industries (Kuybyshevneft')
Title : For a high rate of drilling speed in complex con-
ditions.
Periodical : Neft. khoz., v. 33, #12, 85-87, D 1955
Abstract : Report on the highly satisfactory results of drilling
works for the first 9 months of 1955.
Institution : None
Submitted : No date

KOSTIN, V.M.

Hydraulic resistance of a reflux packing in a uniflow motion
of gas and liquid. Khim. prom. 40 no.10:731-733 0 '64.
(MIRA 18,3)

S/076/60/034/007/019/042/XX
B004/B068

AUTHORS: Skripov, V. P. and Kostin, V. M.
TITLE: Specific Heat of Solutions of Triethyl Amine Forming
Separate Layers in Light and Heavy Water
PERIODICAL: Zhurnal fizicheskoy khimii, 1960, Vol. 34, No. 7,
pp. 1503-1507

TEXT: Starting from a theory developed by V. K. Semenchko (Ref. 1) on critical phenomena, the predictions of a specific-heat maximum in the neighborhood of the critical concentration in separate-layer-forming solutions made according to this theory were examined. For this purpose, the specific heats of triethyl-amine (I) solutions in heavy water (II) and in light water (III) were measured with an adiabatic rocking calorimeter which contained a thermistor and was heated electrically. Five solutions of I and II containing 19.3; 25.6; 30.5; 33.0; and 43.0 % by weight of I, and four solutions of I and III containing 20.1; 25.6; 28.8; and 31.5 % by weight of I were examined. Fig. 2 shows C_p , expressed in cal/mole·degree, as a function of temperature (in °C), and Fig. 4 shows C_p as a function of
Card 1/3

Specific Heat of Solutions of Triethyl Amine S/076/60/034/007/019/042/XX
Forming Separate Layers in Light and Heavy Water B004/B068

the percent by weight of I. For 6.76 mole% of I, a shift of the critical cal separation temperature equal to 3.8°C is observed if H_2O is used instead of D_2O . The highest C_p values are near the critical temperature at which separation of the solution into two layers occurs. This is explained by the fluctuation which precedes the macroscopic layer separation and leads to microscopic latent separation dependent on concentration and temperature. A visual method developed by Alekseyev is mentioned but not described. There are 4 figures and 8 references: 7 Soviet and 1 US.

ASSOCIATION: Ural'skiy politekhnicheskiy institut im. S. M. Kirova,
Sverdlovsk (Ural Polytechnic Institute imeni S. M. Kirov,
Sverdlovsk)

SUBMITTED: September 23, 1958

Text to Fig. 2: 1: Triethyl Amine and Heavy Water; 2: Triethyl Amine and Light Water. Molar Concentration of Triethyl Amine in Both Experiments: 6.76%; a) cal/mole·degree.

Card 2/3

KOSTIN, V. N.

6
①

~~Y~~ ~~Synthesis of hydrocarbons. New reaction of tetrahydro-~~
~~phthalic anhydrides (products of diene synthesis) with phos-~~
~~phorus pentoxide. R. Ya. Levina, V. R. Skvarchenko,~~
~~V. N. Kostin and N. S. Kataeva. Doklady Akad. Nauk~~
~~S.S.S.R. 91-06-8 (1980).—Tetrahydrophthalic anhydrides~~
heated with P_2O_5 split off 2 moles CO and H_2O , forming aro-
matic hydrocarbons. Thus, 29 g. 3,6-dimethyltetrahydro-
phthalic anhydride (I) treated in the molten state with 23 g.
 P_2O_5 and heated carefully to 110–20° reacted vigorously
with heat evolution and distn. of the product (completed by
heating to 230°); washing and distn. gave 80% *p*-xylene,
identified by oxidation to $p-C_6H_4(COOH)_2$. Similarly the
3,5-di-Me isomer of I gave 60% pure *m*-xylene, and the 4,5-
isomer (II) gave 40% pure *o*-xylene. Heating II with 10%
KOH, evapn., and acidification gave 4,5-dimethyltetrahydro-
phthalic acid (III), decomp. 135°, which heated similarly
with P_2O_5 gave 80% *o*-xylene. II treated with dry HCl in
abs. EtOH gave 72% di-Et ester of III, b. 153°, n_D^{20} 1.4679,
 d_4^{20} 1.0450, which heated with P_2O_5 similarly gave 31% *o*-
xylene, with CO and C_2H_4 . G. M. Kosolapoff

MA

ROSTIN, V.N.

7

Synthesis of hydrocarbons. XLIII. Partial reduction of diene hydrocarbons with conjugated system of double bonds by sodium in liquid ammonia. R. Ya. Levina, V. R. Skvarenko, V. N. Kostin, E. G. Urshchikov, and A. S. Okunevich (Moscow State Univ.). *Sbornik Statei Oshchek Khim. Akad. Nauk S.S.S.R.* 1, 355-61(1953); cf. *C.A.* 47, 12210d.—Reduction by Na-NH₃ of 2,3-dimethyl-1,3-butadiene results in addn. of H mainly (98-0%) in the 1,4-positions, and to a minor extent in the 1,2-positions. With the unsym. diene, 2-methyl-2,4-pentadiene, 1,4-addn. predominates also (93-4%). To 87 g. Na in 500 ml. liquid NH₃ was added with stirring an Et₂O soln. of 30 g. 2,3-dimethyl-1,3-butadiene; after 2 hrs. the mixture was dild. with Et₂O, evapd. at room temp., washed and dried, yielding 53% total reduction products, which were sepd. into 3 g. crude material, *b₁₀* 50-72.1°, and 19 g. 2,3-dimethyl-2-butene, *b₁₀* 72.1-72.3°, *n_D²⁰* 1.4134, *d₄* 0.7077; the Raman spectrum of the product had a 1074 cm.⁻¹ line characteristic of the above structure. Raman spectrum of the forerun showed the presence of 2,3-dimethyl-1-butene. Similar reduction of 25 g. 2,4-dimethyl-1,3-pentadiene with Na-NH₃ gave 72% reaction products composed of material, *b₁₀* 81.9°, *n_D²⁰* 1.4033, *d₄* 0.6917, whose Raman spectrum

showed the absence of any contaminant in the 2,4-dimethyl-2-pentene produced (for comparison a specimen was prepd. by pyrolysis of diisopropylcarbinol acetate; the product, *b₁₀* 82.1-82.3°, *n_D²⁰* 1.4073, *d₄* 0.6927, Raman spectrum given. Similar Na-NH₃ reduction of 2-methyl-2,4-pentadiene (*b₁₀* 70.3°, *n_D²⁰* 1.4520, *d₄* 0.7200) gave 2 fractions: 60% 2-methyl-2-pentene, *b₁₀* 57.5-8.5°, *n_D²⁰* 1.3905, *d₄* 0.6663, and about 8% material, *b₁₀* 58.5-67°, which also contained some 2-methyl-3-pentene, confirmed by Raman spectrum of this mixt. in comparison with that of a pure synthetic specimen. Satn. of MeCH:CHCH:CH₃ (25 g.) with 30 g. dry HBr with cooling, followed by addn. of the crude resulting bromide to MeMgBr (from 39 g. MeBr) and stirring 1.5 hrs. in the cold and 6 hrs. at reflux, gave 40% pure 2-methyl-3-pentene, *b₁₀* 57.5-8.5°, *n_D²⁰* 1.3905, *d₄* 0.6665, whose Raman spectrum was not given in the paper; that of the specimen prepd. by Na-NH₃ reduction is given. Similarly, 24 g.

CH₃:CMeCH:CH₃ treated with dry HBr and the resulting bromide treated with MeMgBr gave 13 g. (43%) 2-methyl-3-pentene, *b₁₀* 67°, *n_D²⁰* 1.4035, *d₄* 0.6919, Raman spectrum given. G. M. Kosolapoff

M
gpc

KOSTIN, V.N.

③

Reaction of cyclopropane hydrocarbons with salts of mercuric mercury. II. New organomercury compounds from 1,1,2,2-tetramethylcyclopropane. R. Ya. Levina and V. N. Kostin (M. V. Lomonosov State Univ., Moscow). *Zhur. Obshchei Khim.* 23, 1054-9 (1953); cf. *C.A.* 44, 8321c. — 1,1,2,2-Tetramethylcyclopropane (I), b_m 75.2°, n_D^{20} 1.4003, d_4^{20} 0.7185, (12.8 g.) and 41.6 g. Hg(OAc)₂ in 160 g. H₂O kept 6 days at -5° with occasional shaking gave 64% 3-hydroxy-2,2,3-trimethylbutylmercuric acetate (from heptane), m. 68-70°. Exchange with the corresponding K salts gave: 92% chloride, m. 118.5-17.0°; bromide, 80%, m. 123.5-4.5°; iodide, 85%, m. 102-3.5°; cyanide, 98%, m. 95.5-6.0°; thiocyanate, 93%, m. 73-3.5°. Similarly the reaction in MeOH gave 3-methoxy-2,2,3-trimethylbutylmercuric acetate, 71%, m. 31.5-2.0°, which yielded: 90% chloride, m. 103-3.5°; 91% bromide, m. 133-3.5°; 85% iodide, m. 147-7.5°; cyanide, 90%, m. 53-4°; thiocyanate, 85%, m. 91-1.5°. I in EtOH gave 56% oily 3-ethoxy-2,2,3-trimethylbutylmercuric acetate, which gave: 76% chloride, m. 58.5-0.0°; 72% bromide, m. 61-1.5°; 70% iodide, m. 60-0.5°; 60% oily cyanide; 82% thiocyanate, m. 80.5-1.0°. The acetates stirred with NaOH and SnCl₂ 1 hr. symmetrized to: bis(3-hydroxy-2,2,3-trimethylbutyl)mercury, 90%, m. 113-14°; MeO analog, 85%, m. 50-0.5°; EtO analog, 83%, m. 52-2.5°. G. M. Kosolapoff.

10-15-54

mlf

KOSTIN, V. N. (Engr.)

Dissertation: "The effect of the Material and Character of a Heating Surface on Heat Emission During Boiling." Cand. Tech. Sci., Moscow Institute of Chemical Machine Building, 1 July 54. (Vechernyaya Moskva, Moscow, 22 Jun 54)

SO: Sum. 318, 23 Dec 54

KOSTIN, V. N.

USSR

Reaction of cyclopropane hydrocarbons with salts of mercuric oxide. Organometallic compounds from 1,1,2-trimethylcyclopropane. R. Yu. Lavina and V. N. Kostin (M. V. Lomonosov State Univ., Moscow). *Doklady Akad. Nauk S.S.S.R.* 97, 1027-30 (1954); cf. *C.A.* 48, 8171c. — The reaction between 1,1,2-trimethylcyclopropane (I) (b.p. 52-2.4°, n_D^{20} 1.3940, d_4^{20} 0.6580) and $Hg(OAc)_2$ in aq. medium and in MeOH or EtOH was performed as described previously. The reaction of I with $Hg(OAc)_2$ in aq. soln. or in EtOH yields $Me_2C(OH)CHMeCH_2HgOAc$, which treated with KCl, KBr, KI, KCN, or KCNS gave the expected exchange products. The cleavage of the 3-membered ring of I occurs at 1-3 link since reduction of $Me_2C(OH)CHMeCH_2HgCl$ with Na-Hg in H_2O gave 74% iso-PrCMe₂OH. The following substances are reported: $Me_2C(OH)CHMeCH_2HgX$ (K, X, % yield, and m.p. given): H, OAc , 62, 34.5-5°; H, Cl , 60, 60.5-1°; H, Br , 58, 65.5-6°; H, I , 80, 68-3.5°; H, CN , 92, 64-3°; Me, Cl , 92, 33-4°; Me, Br , 92, 58.5-9°; Me, I , 85, 31-3°; Me, CNS , 83, 33.5-4°; Et, I , 70, 40-40.5°; Et, CNS , 80, 45-8°.

G. M. Kosolapoff

KOSTIN, V.N.

Name: KOSTIN, V. N.

Dissertation: Cyclopropane hydrocarbons in the synthesis of mercury organic compounds

Degree: Cand Chem Sci

Defended at:
Affiliation: Moscow State U imeni M. V. Lomonosov

Publication
~~Defense Date~~, Place: 1956, Moscow

Source: Knizhnaya Letopis', No 2, 1957

LEVINA, R. Ya.; KOSTIN, V.N.; TARTAKOVSKIY, V.A.

APPROVED FOR RELEASE: 06/14/2000 CIA-RDP86-00513R000825220006-

Interaction of cyclopropane hydrocarbons and mercury oxide salts.
Part 4. Effect of mercury acetate on 1,1,2-trimethyl-2-ethylcyclo-
pentylsulfide. Vest.Mosk.un.11 no.2:77-82 F '56. (MLBA 9:8)

1. Kafedra organicheskoy khimii.
(Mercury acetate) (Cyclopropane)

Kostin, V.N.

6

7
 ✓ Reaction of cyclopropane hydrocarbons with salts of
 mercuric acetate Action of mercuric acetate on
 methylcyclopropane and 1,1-dimethylcyclopropane
 Ya. Levina, V. N. Kostin and V. I. Pavlovskii
 (1960), Chem. Abstr. 54, 12484
 Addn. of 180 g. HBr to 150 g. cyclopropane
 a mixt. of monobromides which is obtained at room
 temp. is dried at 40°C. and 10 mm. Hg.
 HBr was added to the residue and the mixture
 was dried at 40°C. and 10 mm. Hg.
 The residue was dried at 40°C. and 10 mm. Hg.
 The residue was dried at 40°C. and 10 mm. Hg.
 The residue was dried at 40°C. and 10 mm. Hg.
 The residue was dried at 40°C. and 10 mm. Hg.

chem 1-4E 2C

11/12

washed with cold, dil. $KMnO_4$. Shaking 25 g I with 110 g $Hg(OAc)_2$ in 400 ml. H_2O 10-12 hrs gave, after filtration, concn., and extr. with Et_2O , 75% $HO(CH_2)_3CH_2HgOAc$ (II). The use of MeOH as solvent gave similarly 40% oily $MeO(CH_2)_3CH_2HgOAc$. These treated with the corresponding K salts gave the following $RtHgX(R, X, \text{ and } m.p. \text{ shown})$: $HO(CH_2)_3CH_2HgCl$ (I, 92.3°; $f. 119-20^*$); $MeO(CH_2)_3CH_2HgCl$ (II, 15.8°; $CNS, 37.8^*$; $CN, 15.1^*$). I and II and CN derivs. of the HO compd. were prepared in 80% yield by similar treatment of 10 g of I with 120 g $Hg(OAc)_2$ in 400 ml. H_2O for 10 hrs. The corresponding $RtHgOAc$ (III) was prepared from 10 g of I with 120 g $Hg(OAc)_2$ in 400 ml. H_2O for 10 hrs. The corresponding $RtHgOAc$ (IV) was prepared from 10 g of II with 120 g $Hg(OAc)_2$ in 400 ml. H_2O for 10 hrs. The iodide and cyanide were only 10% MeOH soluble. The $HO(CH_2)_3CH_2HgOAc$ (II) was prepared from 10 g of I with 120 g $Hg(OAc)_2$ in 400 ml. H_2O for 10 hrs. The corresponding $RtHgOAc$ (III) was prepared from 10 g of I with 120 g $Hg(OAc)_2$ in 400 ml. H_2O for 10 hrs. The corresponding $RtHgOAc$ (IV) was prepared from 10 g of II with 120 g $Hg(OAc)_2$ in 400 ml. H_2O for 10 hrs. Thus the ring is not broken by $Hg(OAc)_2$ acting between the substituted C atoms of the ring.

Kostin, V.N.

SOV/137-58-8-16597

Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 8, p 51 (USSR)

AUTHOR: ~~Kostin, V.N.~~

TITLE: [Shortcomings in Gas Cleaning in Nonferrous Metallurgy and Methods of Improving the Situation] ([O nedostatkakh v oblasti pyleulavlivaniya v tsvetnoy metallurgii i mery po uluchsheniyu yego])

PERIODICAL: Sb. materialov po pyleulavlivaniyu v tsvetn. metallurgii. Moscow, Metallurgizdat, 1957, pp 9-12

ABSTRACT: A paper delivered at the conference on problems of gas cleaning (G) in nonferrous metallurgy held in Moscow April 10-14, 1956. Improvement in the state of G at nonferrous metals plants during recent years are noted. It is noted that the rates of construction and launching of new gas-cleaning installations are found to be inadequate. Examples of shortcomings in the servicing of gas-cleaning plants are cited. Ideas are put forth on the organization of planning and scientific research in the field of G. Note is taken of the need to improve organization of the structure of G services and of the system of payment of service personnel. 1. Gases--Cleaning 2. Industrial plants G.G. --Performance

Card 1/1

KOSTIN, V.N.
LEBEDEVA, M.N.; YEFRENOVA, S.A.; KOSTIN, V.N.; LEVINA, R.Ya.

Bacteriostatic properties of ~~mercurated~~ -mercurated alcohols and their
alkyl ethers. Vest.Mosk.un.Ser.mat., mekh., astron., fiz., khim.
12 no.3:149-158 '57. (MIRA 11:3)

1. Kafedra mikrobiologii I Moskovskogo ordena Lenina meditsinskogo
instituta i Kafedra organicheskoy khimii Moskovskogo gosudarstvennogo
universiteta.

(Alcohol) (Ethers) (Bacteriostasis)

Koslov, V. N.

Reaction of cyclopropane hydrocarbons with mercuric salts. VI. Organomercury compounds from phenylcyclopropane. R. Ya. Levina, V. N. Kostin, and V. A. Tarakovskii (Sov. Chem. Soc.), *Zh. Obshch. Khim.* 27, 231-2 (1957); cf. C.A. 51, 8658a. Shaking 12 g. Hg(OAc)₂ with 45 g. phenylcyclopropane (I) in 40 ml. H₂O 12 hrs. gave after extn. with Et₂O and distn. of the ext. 71% PhCH(OH)CH₂CH₂HgOAc, m. 65-6°; this in ice cooled aq. soln. was treated with KBr, KI, KCNS, and KCN, but only PhCH(OH)CH₂CH₂HgCNS, m. 61°, was isolated. Treatment of the acetate with Na stannite gave Hg[CH₂CH(OH)Ph]₂, m. 77-8° (aq. Me₂CO). Shaking 37 g. Hg(OAc)₂ and 10 g. I in dry MeOH 12 hrs. gave 82% PhCH(OMe)CH₂CH₂HgOAc, oil. This with appropriate salts gave: PhCH(OMe)CH₂CH₂HgCl, m. 82°; PhCH(OMe)CH₂CH₂HgBr, m. 87°; PhCH(OMe)CH₂CH₂HgI, m. 79°; PhCH(OMe)CH₂CH₂HgCN, m. 84°. To EtMgBr from 3 g. Mg was added dropwise 30 g. PhCHOHCH₂CH₂HgCl in Et₂O; after refluxing 1 hr. and decompn. with aq. AcOH there was obtained EtHgCH₂CH₂CHPhOH, an oil, which heated in vacuo symmetrized and yielded 51% Et₂Hg and 47.5% Hg[CH₂CH₂CH(OH)Ph]₂. When 3 moles EtMgBr was used/mole Hg deriv. and the mixt. refluxed 12 hrs. there were obtained Et₂Hg and 44% EtPhCHOH, identified as the 3,5-dinitrobenzoate, m. 67-8°. G. M. Kosolapoff //

Distr: hEh j/hE3d/hE2c(j)

KOSTIN, V.N.

7

~~* Reaction of cyclopropane hydrocarbons with salts of mercuric mercury. VII. Thermal decomposition of products of symmetrization of γ -mercurated alcohols. R. Ya. Levina, V. N. Kostin, and V. A. Tartakovsky (State Univ. Moscow). *Zhur. Obshch. Khim.* 27, 2049-52 (1957); cf. *C.A.* 52, 3713i. Heating $(PhCH(OH)CH_2CH_2)_2Hg$ to 120-30° at 15 mm. gave a distillate of Hg and a mixt. of $EtPhCHOH$ and $EtCOPh$. Similar heating to 100-10° of $(Me_2C(OH)CF_2CH_2)_2Hg$ gave *tert*-AmOH and Me_2CO and Hg, with apparently 1-hexene. The 1st substance apparently yields free radicals the HO group of which is the source of H for the final products. The 2nd compd. in its decompn. yields tertiary free radicals which are cleaved at the central C atom to yield the above products with probably intermediacy of Me_2CO , C_2H_4 , and H.~~

G. M. Kosolapoff

Bistr: 4E4j/4E2c(j)/
4E3d

7
2m up
3
1/1

[Handwritten signature]

KOSTIN, V. N.

18(5,6); 25(1) P. 2

PHASE I BOOK EXPLOITATION

SOV/3134

Kommunisticheskaya partiya Sovetskogo Soyuz. Vysshaya partiynaya shkola

Dostizheniya nauki i tekhniki i peredovoy opyt v promyshlennosti i stroitel'stve. Vyp. 2, chernaya i tsvetnaya metallurgiya (Progress in Science and Technology and Advanced Methods Applied in Industry and Construction. No. 2, Ferrous and Nonferrous Metallurgy) Moscow, Izd-vo VPSH i AON pri TsK KPSS, 1958. 157 p. 22,000 copies printed.

Ed. (Title page): G. I. Pogodin-Alekseyev, Doctor of Technical Sciences, Professor;
Ed. (Inside book): G. V. Popova; Tech. Ed.: K. M. Naumov.

PURPOSE: This book is intended for the informed reader and should also be of interest to metallurgists.

COVERAGE: This is a collection of lectures, presumably delivered at the Vysshaya partiynaya shkola (Higher Party School) of the Communist Party, USSR, describing recent advances in the field of metallurgy. The approach is basically nontechnical, though a number of processes are briefly described. Specific ore deposits and metallurgical plants are referred to.

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Progress in Science (Cont.)

Some statistics are given. No personalities are mentioned. There are no references.

TABLE OF CONTENTS:

Myrtsyomov, A. F., Candidate of Technical Sciences. Technical Progress in Ferrous Metallurgy. Ways of Increasing the Productivity of the Blast-furnace Process

Claims of actual achievements described in this article include the highest blast-furnace operating efficiency in the world in 1955, when the Soviet yield of pig iron per cubic meter of useful blast furnace is stated to have been 11 percent higher than in the United States. The 1956 Soviet yield of open-hearth steel per square meter of hearth is given as 7 metric tons per 24-hr. period, as compared with 5.7 for the United States. Current objectives in Soviet ferrous metallurgy include the following: 1) greater application of open-pit mining; increased mechanization of mining operations, prospecting for additional deposits; accelerating concentration operations, construction of several large ore-beneficiation combines in the Krivoy Rog Basin, and new concentration plants in other areas;

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Card 2/7

Progress in Science (Cont.)

SOV/3134

process; vacuum melting and teeming; high-temperature refractories; and direct reduction of iron from ore.

Tselikov, A. I., Corresponding Member, Academy of Sciences, USSR.
Progressive Methods and Basic Trends in the Development of Metal Forming 43
 According to the author, improvements in metal-forming techniques are currently being achieved with two basic objectives, 1) reduction of metal waste, and 2) automation of processes. Attempts are being made to attain the first objective by designing lighter finished products and by using pre-sized blanks and slugs to reduce allowances. As regards the increased-automation program, the main line of attack is the conversion of noncontinuous to continuous forming processes, especially in rolling, drawing, and extruding. In the case of hot and cold upsetting continuity of auxiliary operations is the aim, while in stamping operations, continuous feed is considered important. Particular reference is made to the Soviet method of helical cross rolling, by means of which cavities may be developed in originally solid bodies. The method is frequently used in the rolling of hollow bodies of rotation.

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Progress in Science (Cont.)

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Charikhov, L. A. **Automation of Production Processes in Ferrous Metallurgy** 54
 An account is given of recent Soviet advances in the automation of blast-furnace, steelmaking, and rolling processes.

Kornev, Yu. V., Candidate of Physical and Mathematical Sciences. **Radioactive Isotopes in Metallurgy** 80

Basic information on both natural and artificial radioactive isotopes is given. Production of artificial isotopes is briefly discussed. Such applications of radioactive isotopes in metallurgy as the following are described: use of cobalt 60 in the continuous casting of steel to measure the level of liquid metal in the crystallizing mold; flaw detection; measurement of metal thicknesses; controlling composition of alloys; determining rate of wear of blast-furnace well lining; tracing the motion of charge materials in the blast furnace; and determining the weight of steel in the teeming ladle.

Kostin, V. N. **Nonferrous Metallurgy and Basic Trends in Its Development** 104
 Nonferrous ore deposits and their locations are described. It is stated that 63 nonferrous metals are currently being extracted from ores in the Soviet Union, 45 in metallic form, the remainder as

Card 5/7

KOSTIN, V.N.

136-1-2/20

AUTHOR: Kostin, V.N.TITLE: Electrothermics Should be Adopted in Lead and Zinc Metallurgy
(Dorogu elektrotermii v metallurgii svintsa i tsinka)

PERIODICAL: Tsvetnyye Metally, 1958, No.1, pp. 3 - 10 (USSR).

ABSTRACT: The author states that electrothermics represents a potentially valuable technical development for lead and zinc production in the near future. He lists possible fields of application and, without going into process details, examines the main peculiarities of those developments and touches on reasons for comparative slowness of the adoption of electrothermics. The author deals with work carried out since 1953 by the VNIItsvetmet institute and the Leninogorsk Combine (Leninogorskiy kombinat) on the electric smelting of lead sinter on a scale up to that of a 2 500 kVA furnace. He summarises results obtained in twenty days of operation in the autumn of 1957 with a fine sinter containing 38.2% lead with the furnace operating at 2 000 kVA, 84 V, and a daily production rate equivalent to 3.6 tons of sinter per m² of furnace bottom. The power consumption was 600 - 650 kWh per ton sinter, and the author recommends the adoption of this procedure at existing plants. For new plants he advocates the adoption of a process so far only based on laboratory-scale experiments by the VNIItsvetmet in

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136-1-2/20

Electrothermics Should be Adopted in Lead and Zinc Metallurgy

which a granular lead concentrate, roasted in a fluidised bed, is smelted. The author lists and discusses the advantages of both the variants over shaft smelting. He states that the power cost of 6-8 kopeks per kWh prevailing in several regions of the USSR makes the process competitive with shaft smelting. He makes suggestions on a programme for the near future, which includes the construction at the Leninogorsk Combine of a two-furnace electric shop in 1958 and of new electric shop at the Ust'-Kamenogorsk Works (Ust'-Kamenogorskiy zavod) in 1959-1960 from designs to be worked out by the Giprovetsmet organisation; an extensive research programme on the second variant in 1958 by VNIItsvetmet; the adoption of electric smelting at the "Elektrotsink" Works after appropriate estimates by the works staff, together with the Kavgiprotsvetmet organisation; the adoption of electric smelting at the Chimkentskiy Works, plans being drawn up in 1958. The author urges the VNIItsvetmet and Giprotsvetmet organisations to take a more positive attitude towards the wider adoption of the process and touches on the functions of a number of research and design organisations, including these and Gintsvetmet and Giredmet. He advocates the creation of large, suitably supported planning offices (SKB).

Card2/4

AUTHORS: Levina, R. Ya., Kostin, V. N. SOV/79-28-12-33/41

TITLE: Reaction of Cyclopropane Hydrocarbons With Salts of Mercury Oxide (Vzaimodeystviye tsiklopropanovykh uglevodorodov s solyami okisi rtuti) VIII. Reaction of γ -Mercurized Alcohols With Zinc Dust (VIII. Reaktsiya γ -merkurirovannykh spirtov s tsinkovoy pyl'yu)

PERIODICAL: Zhurnal obshchey khimii, 1959, Vol 28, Nr 12, pp 3307-3311 (USSR)

ABSTRACT: The structure of the γ -mercurized alcohols (I) obtained by the authors by a synthesis earlier carried out by them, where the product was synthesized by the reaction of alkyl and aryl cyclopropanes with mercury salts dissolved in water (Refs 1-6), had to be proved by reduction (i.e. by means of substituting the HgX-group by hydrogen). The first reduction experiments were too complicated (Scheme 1). This time the authors attempted to realize the symmetrization of the γ -mercurized alcohols (I) and the thermal decomposition of the products obtained in this symmetrization in one stage by heating the γ -mercurized alcohols with zinc dust. The previous symmetrization stage was thus avoided to explain its

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Reaction of Cyclopropane Hydrocarbons With Salts
of Mercury Oxide, VIII. Reaction of γ -Mercurized
Alcohols With Zinc Dust

SOV/79-28-12-33/41

structure. Thus, the structural proof was made easier. A secondary alcohol and a ketone with the carbon skeleton of the organo-mercury compound used as initial product are formed from the γ -mercurized alcohols containing a secondary alcohol group. The γ -mercurized alcohols with a tertiary alcohol group form tertiary alcohols with the carbon skeleton of mercurized alcohol, whereas the formation of the ketones is connected with a cleavage of the C-C bond. On heating the β -mercurized alcohol obtained from triptene by the action of mercury acetate a separation of the mercury salt and the formation of the initial ethylene hydrocarbon, triptene, took place. The reaction of the thermal decomposition in the presence of zinc dust was also attempted with 3-oxy-2,3-dimethyl-butyl mercury acetate (III). From this γ -mercurized tertiary alcohol the dimethyl isopropyl carbinol, acetone and propylene were obtained (Scheme 4). There are 14 references, 10 of which are Soviet.

Card 2/3

Reaction of Cyclopropane Hydrocarbons With Salts
of Mercury Oxide. VIII. Reaction of γ -Mercurized
Alcohols With Zinc Dust

SOV/79-28-12-33/41

ASSOCIATION: Moskovskiy gosudarstvennyy universitet (Moscow State University)

SUBMITTED: November 12, 1957

Card 3/3

AUTHORS:

Levina, R. Ya., Kostin, V. N.,
Shanazarov, K. S.

SOV/79-29-1-9/74

TITLE:

Reaction of Cyclopropane Hydrocarbons With Salts of Mercury Oxide (Vzaimodeystviye tsiklopropanovykh ulevodorodov s solyami okisi rtuti) IX. γ -Mercurized Alcohols and Their Ethers From Aryl Cyclopropanes (IX. γ -Merkurirovannyye spirty i ikh efiry iz ariltsiklopropanov)

PERIODICAL:

Zhurnal obshchey khimii, 1959, Vol 29, Nr 1, pp 40-44 (USSR)

ABSTRACT:

In an earlier paper the authors described (Ref 1) the decomposition of the three-membered cycle in phenyl cyclopropane under the action of mercury acetate. The present paper deals with the reaction of this acetate in an aqueous and methanol solution with aryl cyclopropanes which in the benzene ring contain a methyl- and methoxy group in para-position. It was found that the reaction takes place under the same conditions as in the case of phenyl cyclopropane, i.e. when the reaction mass is shaken for hours at low temperature. The addition of mercury acetate to the two aryl cyclopropanes (p-anisyl- and p-tolyl cyclopropane) leads to the formation of 3-cxy-3-aryl propyl mercury acetates and

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Reaction of Cyclopropane Hydrocarbons With Salts of
Mercury Oxide. IX. γ -Mercurized Alcohols and Their
Ethers From Aryl Cyclopropanes

SOV/79-29-1-9/74

β -methoxy- β -aryl propyl mercury acetates respectively. Under the action of potassium salts upon them the corresponding crystalline salts of γ -mercurized secondary alcohols and their methyl ethers (Scheme 1) are formed. The structure of the γ -mercurized secondary alcohols was determined according to the method which had already been previously worked out by the authors (Ref 2). The synthesis of the crystalline organo-mercury compounds can serve for the identification of p-anisyl cyclo- and p-tolyl cyclopropane. A new description was given of the following compounds: β -oxy- β -p-anisyl propyl mercury acetate and its chloride, bromide, iodide, cyanide and thiocyanate; β -oxy- β -p-tolyl propyl mercury acetate; β -methoxy- β -p-tolyl propyl mercury chloride and its bromide. There are 2 tables and 6 references, 3 of which are Soviet.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet (Moscow State
University)

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Reaction of Cyclopropane Hydrocarbons With Salts of
Mercury Oxide. IX. γ -Mercurized Alcohols and Their
Ethers From Aryl Cyclopropanes

SO7/79-29-1-9/74

SUBMITTED: November 18, 1957

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5(3)

SOV/79-29-6-41/72

AUTHORS:

Levina, R. Ya., Kostin, V. N., Kim Dyay Gir, Ustynyuk, T. K.

TITLE:

Reaction of Cyclopropane Hydrocarbons With Mercuric Salts
(Vzaimodeystviye tsiklopropanovykh uglevodorodov s solyami
okisi rtuti). X. Action of Mercury Acetate on (0,1,3)-Bicyclo-
hexanes (X. Deystviye atsetata rtuti na (0,1,3)-bitsiklogeksany)

PERIODICAL:

Zhurnal obshchey khimii, 1959, Vol 29, Nr 6,
pp 1956 - 1960 (USSR)

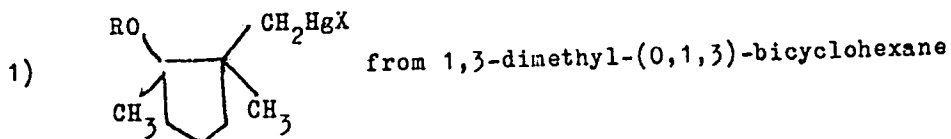
ABSTRACT:

The products of the reactions described earlier (Refs 1-4) which are characteristic of the above hydrocarbons and which take place under the opening of the three-membered cycle i.e. the γ - alcohols and their ethers were the products of the reaction with mercury acetate in aqueous and alcoholic solutions. Their investigation revealed the cleavage of the three-membered cycle between the hydrocarbon atoms which were the most strongly and the most weakly substituted (Scheme 1). The forming crystalline organomercury compounds may be successfully for the identification of the cyclopropane hydrocarbons. In the present paper it was found that the above reaction is also characteristic of the "bridge bicyclo hydrocarbons" in the system of which

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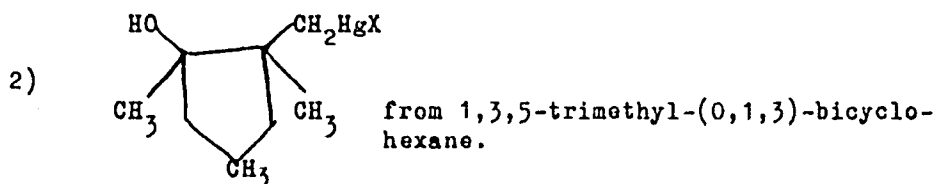
Reaction of Cyclopropane Hydrocarbons With Mercuric Salts. X. Action of Mercury Acetate on (0,1,3)-Bicyclohexanes SOV/79-29-6-41/72

the three-membered cycle is integrated. It was shown (Scheme 2) that the opening of the three-membered cycle in the homologues of (0,1,3)-bicyclohexane in the action of mercury acetate on these homologues in aqueous or methanol solution does not take place at the bridge C-C-bond. The products of this reaction are cyclopentanes γ -mercurized in the side chain or their methyl ethers. The synthesis of the crystalline γ -mercurized alcohols and their ethers may thus serve for the determination of 1,3-dimethyl-1,3,5-trimethyl and 1-methyl-3-ethyl-(0,1,3)-bicyclohexane. The structure of the γ -mercurized alcohols and their ethers is the following:



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Reaction of Cyclopropane Hydrocarbons With Mercuric Salts. X. Action of Mercury Acetate on (0,1,3)-Bicyclohexanes SOV/79-29-6-41/72



There are 3 tables and 7 Soviet references.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet (Moscow State University)

SUBMITTED: May 25, 1958

Card 3/3

5 (3)

AUTHORS:

Levina, R. Ya., Kostin, V. N.,
Gembitskiy, P. A.

SOV/79-29-7-80/83

TITLE:

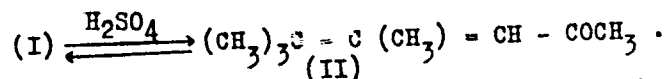
Letter to the Editor (Pis'mo v redaktsiyu). On the Photochemical
Isomerization of Vinyl Ketones Into Allyl Ketones (O foto-
khimicheskoy izomerizatsii vinilketonov v allilketony)

PERIODICAL:

Zhurnal obshchey khimii, 1959, Vol 29, Nr 7, pp 2456-2458 (USSR)

ABSTRACT:

The acylation of triptene (2,2,3-trimethyl butene-3) with acetic anhydride in the presence of phosphoric acid yielded the unsaturated ketone, with a β, γ -position of the double bond with respect to the carbonyl group, the 2,2-dimethyl-3-methylene hexanone-5 (I) (Scheme 1). On standing for some months the ketone (I) isomerized to form the 2,2,3-trimethyl-hexen-3-one-5 (II), under rearrangement of the double bond. An equilibrated mixture of the ketone (I) and its isomerization product (II) is there formed



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If this resulting equilibrated mixture of the isomeric unsaturated ketones is irradiated with ultraviolet light in

Letter to the Editor. On the Photochemical Isomerization SOV/79-29-7-80/83
of Vinyl Ketones Into Allyl Ketones

quartz, its α,β -form, the vinyl ketone (II), isomerizes completely to produce the initial β,γ -form, the allyl ketone (I): [(II) \longrightarrow (I)]. In this way a pure β,γ -unsaturated ketone, the 2,2-dimethyl-3-methylene hexanone-5 is formed from the mixture of the α,β - and β,γ -unsaturated ketones. In order to clarify whether this photochemical isomerization of the vinyl into the allyl ketones is of general nature, 1-acetyl cyclohexene-1 was likewise irradiated; about the half of this compound was found to be isomerized into the 1-acetyl-cyclohexene-2 (Scheme 3). Thus the vinyl ketones isomerize by a photochemical process completely or partially to give the allyl ketones. There are 2 Soviet references.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet (Moscow State University)

SUBMITTED: March 24, 1958

Card 2/2

83353

S/139/60/000/004/010/033
E032/E514

9.6150
26.2420
AUTHORS:

Kostin, V.N. and Zabara, M.Ya.

TITLE: Control of the Spectral Characteristics of Composite Photocathodes for the Ultraviolet Part of the Spectrum

PERIODICAL: Izvestiya vysshikh uchebnykh zavedeniy, Fizika, 1960, No.4, pp.110-116

TEXT: The control of the spectral characteristics of composite photocathodes is particularly important in the ultraviolet region. In the dosimetry of ultraviolet radiation it is desirable for the spectral sensitivity of the dosimeter to be roughly the same as the spectral sensitivity of the corresponding biological or photo-chemical process. The most widely used method for obtaining this correspondence is based on the use of light filters in combination with photo-electric devices. The aim of the present paper was to obtain an approximate matching by controlling the spectral characteristics of the photocathode used in the dosimeter. The control of the spectral characteristics of a composite photocathode can be obtained as follows. In Fig.1 the curves marked 1 and 2 represent plots of sensitivity versus wavelength for two different

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S/139/60/000/004/010/033
E032/E514Control of the Spectral Characteristics of Composite Photocathodes
for the Ultraviolet Part of the Spectrum

materials. If the surface of the photocathode made of material 1 is covered by fine and uniformly distributed spots of metal 2 and the area covered by the latter metal is gradually increased, then the characteristic of the composite photocathode will be displaced from position AB to position CD, i.e. it will lie between the characteristics of metals 1 and 2. If the metals 1 and 2 have widely differing characteristics, then a composite photocathode made up of these two metals can be varied within relatively wide limits. In the present work the cathode 1 (Fig.2) was prepared from aluminium or red copper and was in the form of a hollow cylinder (25 mm long, internal diameter 18 mm) and cut along a plane containing its axis. The cathode was baked in a high vacuum and then deposits of cadmium or silver or silver spots on cadmium were evaporated onto the cathode. It was shown that by using cadmium in combination with silver it is possible to obtain a spectral characteristic which is similar to the erythematic effect curve in the region between 240 and 280 m μ . An investigation was also made of the stability of such photocathodes and the ion

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bombardment treatment ensuring constant photo-sensitivity. It was found that a photon counter filled with pure hydrogen and having a composite cadmium-silver cathode had stable characteristics after intense ion bombardment. Fig.7 shows a typical characteristic obtained with a cadmium-silver composite photocathode as well as the separate characteristics for cadmium and silver. Acknowledgments are made to Academician K. D. Sinel'nikov and Docent V. K. Tkach for valuable advice and assistance. There are 7 figures and 3 Soviet references.

ASSOCIATION: Khar'kovskiy gosuniversitet imeni A. M. Gor'kogo
(Khar'kov State University imeni A. M. Gor'kiy)

SUBMITTED: July 3, 1959

Card 3/3

5.3300

77852
SOV/79-30-2-3/78

AUTHORS: Levina, R. Ya., Kostin, V. N., Ustuyuk, T. K.
TITLE: Reaction of Cyclopropanes With Mercuric Salts. XI. Cleavage of the Three-Membered Ring in (0, 1, N)-Bicycloalkanes (N = 2,3, and 4)

PERIODICAL: Zhurnal obshchey khimii, 1960, Vol 30, Nr 2, pp 359-363 (USSR)

ABSTRACT: The action of mercuric acetate on (0,1,3)-bicyclohexane (I, northujane), (0,1,4)-bicycloheptane (IV, norcarane), and (0,1,2)-bicyclopentane (VI), as well as other (0,1,N)-bicycloalkanes, can be used to identify these substances by the final reaction products -mercurated alcohols and their derivatives. The reactions for compounds I, IV, and VI are represented below:

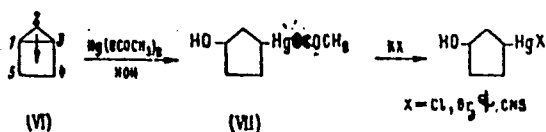
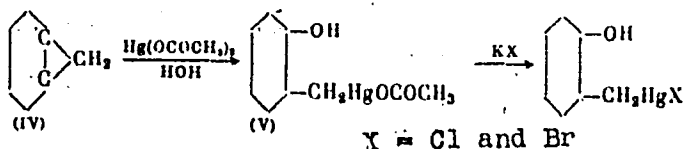
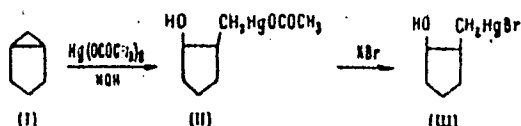
Card 1/5

APPROVED FOR RELEASE: 06/14/2000

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Reaction of Cyclopropanes With Mercuric Salts. XI

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SOV/79-30-2-3/78



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Reaction of Cyclopropanes With Mercuric Salts.
 XI

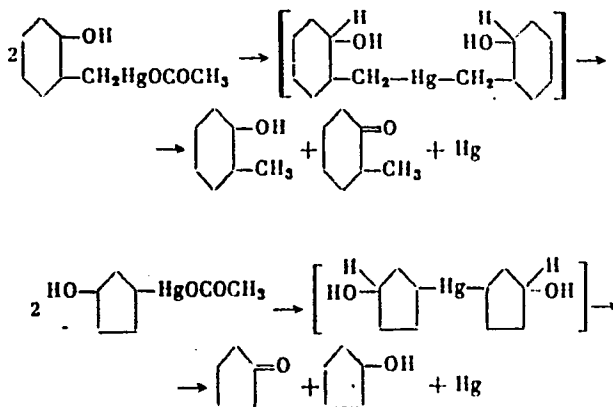
77852
 SOV/79-30-2-3/78

The reactions were performed by mixing the respective hydrocarbon with mercuric acetate (at room temperature for compounds I and IV and at 0° for VI) and by subsequent addition of potassium bromide to the product (the mercurated alcohols II, V, and VII) to obtain the halomercuri-derivatives. The cleavage of the three-membered ring in (0,1,2)-bicyclopentane takes place at the carbon-carbon "bridge" bond, while in the case of (0,1,3)-bicyclohexane and (0,1,4)-bicycloheptane the bond between the least and the most substituted carbon atoms is broken, leading to the formation of mercurated alcohols of the cycloheptane and cyclohexane series, respectively. The site of the ring cleavage was determined by results of thermal decomposition of mercurated alcohols. The reactions and final products for the pyrolysis of V and VII are shown below:

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Reaction of Cyclopropanes With Mercuric Salts.
 XI

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The following derivatives of the mercurated alcohols were isolated and analyzed: 2-hydroxy-1-bromo-

Reaction of Cyclopropanes With Mercuric Salts.
XI

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mercurimethylcyclopentane (III), 151-152°; 2-hydroxy-1-acetoxymethylcyclohexane (V), mp 129-130°; 3-hydroxy-1-chloromercuricyclopentane, mp 97-98°; 3-hydroxy-1-bromomercuricyclopentane, mp 129-130°; 3-hydroxy-1-thiocyanomercuricyclopentane, mp 110-111°; 2-hydroxy-1-chloromercurimethylcyclohexane; mp 122-123°; 2-hydroxy-1-bromomercurimethylcyclohexane, mp 114-115°. There are 13 references, 4 Soviet, 5 German, 1 Belgian, 1 Swiss, 2 U.S. The U.S. references are: W. E. Doering, A. K. Hoffman, J. Am. Chem. Soc., 76, 6162 (1954); C. R. Noller, R. Adams, J. Am. Chem. Soc., 48, 1084 (1926).

ASSOCIATION: Moscow State University (Moskovskiy gosudarstvennyy universitet)

SUBMITTED: March 2, 1959

Card 5/5

5.3500

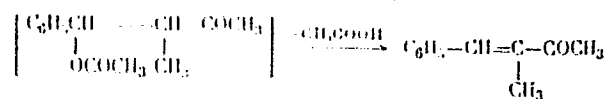
73775
SOV/79-30-3-27/69

AUTHORS: Levina, R. Ya., Kostin, V. N., Gembitskiy, P. A.,
Treshchova, Ye. G.

TITLE: Cyclopropanes and Cyclobutanes. X. Acylation of
Cyclopropanes

PERIODICAL: Zhurnal obshchey khimii, 1960, Vol 30, Nr 3,
pp 868-875 (USSR)

ABSTRACT: The behavior of the cyclopropane ring in phenyl-
cyclopropane, an acylation with acetic anhydride
in the presence of phosphoric acid, was studied.
The reaction proceeds through isomerization of
cyclopropanes into alkenes, which on further acylation
yield an unsaturated ketone, 2-benzylidenbutanone
(yield 36.5%), mp 30°. $C_6H_5-CH_2-CH_2-CH_2 \xrightarrow{(CH_3CO)_2O, (H_3PO_4)}$

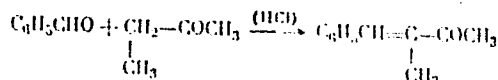


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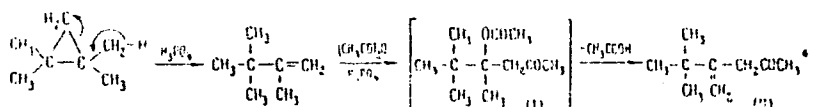
Cyclopropanes and Cyclobutanes. X.
Acylation of Cyclopropanes

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SOV/79-30-3-27/69

All other cyclopropanes behave similarly. The structure of the prepared 2-benzylidenbutanone was confirmed by parallel synthesis.



Isomerization of phenylcyclopropane and 1,1,2,2-tetramethylcyclopropane in acetic anhydride in the presence of phosphoric acid yields propenylbenzene and 2,2,3-trimethylbutene, bp 162° (745 mm), n_D²⁰ 1.4530, respectively. This constitutes a heterolytic cleavage of the cyclopropane ring.

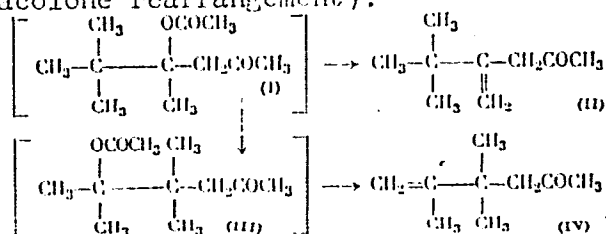


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Cyclopropanes and Cyclobutanes. X.
 Acylation of Cyclopropanes

1959
 SOV/79-30-3-27/69

Acylation of 1,1,2,2-tetramethylcyclopropane or 2,2,3-trimethylbutene yields a β, γ -unsaturated ketone, 2,2-dimethyl-3-methylenehexan-5-one, which is formed from the intermediate acetoxyketone by elimination of an acetic acid molecule (without retro-pinacolone rearrangement).



The structure of unsaturated ketones formed on acylation of phenylcyclopropane and 1,1,2,2-tetramethylcyclopropane was confirmed by absorption spectra of their 2,4-dinitrophenylhydrazones, and by comparison of their properties with those of authentic samples. There are 12 references, 6

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Cyclopropanes and Cyclobutanes. X.
 Acylation of Cyclopropanes

1959
 SOV/79-30-3-27/69

Soviet, 2 U.S., 2 U.K., 1 French, 1 German. The U.S. and U.K. references are: Hart, H., Curtis, O. E., Jr., J. Am. Chem. Soc., 79, 931 (1957); Sukh, Dev, Chem. and Ind., 1071 (1954); Hartough, H., Kosak, A., J. Am. Chem. Soc., 69, 3093 (1947); Perkin, W. H., J. Chem. Soc., 69, 1028 (1896).

ASSOCIATION: Moscow State University (Moskovskiy gosudarstvennyy universitet)

SUBMITTED: March 31, 1959

Card 4/4

LEVINA, R.Ya.; KOSTIN, V.N.; GEMBITSKIY, P.A.; SHOSTAKOVSKIY, S.M.;
TRESHCHOVA, Ya.G.

Cyclopropylmesitylene and *p*-cyclopropylcumene. Zhur.ob.
khim. 30 no.7:2435-2436 J1 '60. (MIRA 13:7)

1. Moskovskiy gosudarstvennyy universitet.
(Mesitylene) (Cumene)

84886

5.3300 only 2209, 1236

S/079/60/030/010/029/030
B001/B066AUTHORS: Levina, R. Ya., Kostin, V. N., Gembitskiy, P. A., and Shostakovskiy, S. M.TITLE: New Hydrocarbons of the Cyclopropane SeriesPERIODICAL: Zhurnal obshchey khimii, 1960, Vol. 30, No. 10,
pp. 3502 - 3503

TEXT: The authors continued their investigations on cyclopropanes (Refs. 1 and 2) by synthesizing some new compounds of this series. By partial reduction of phenyl- and p-tolyl cyclopropane with sodium (in liquid ammonia) and methyl alcohol, 1-cyclopropyl-cyclohexadiene-1,4 and 1-methyl-4-cyclopropyl-cyclohexadiene-1,4 were synthesized. The subsequent catalytic hydrogenation of the double bonds in 1-cyclopropyl-cyclohexadiene-1,4 (on a copper-chromium catalyst at a pressure of 100 atm) at 95° and 125° gave cyclopropyl-cyclohexene-1 and, later, cyclopropyl-cyclohexane; the latter was also obtained by methylenation (Ref.3) of vinyl cyclohexane. p-cyclopropyl styrene and p-cyclopropyl-isopropenyl benzene were, accordingly, synthesized by dehydration of

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New Hydrocarbons of the Cyclopropane Series S/079/60/030/010/029/030
B001/B066

methyl- and dimethyl-p-cyclopropyl-phenyl carbinols (both carbinols were obtained from p-cyclopropyl-acetophenone). p-dicyclopropyl benzene resulted both from p-cyclopropyl acetophenone, according to the stepwise reactions of Mannich and Kizhner (Ref.4), and from p-cyclopropyl styrene by methylenation (Ref.3). The constants of the resultant hydrocarbons are tabulated. There are 1 table and 4 references: 3 Soviet and 1 US.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet (Moscow State University)

SUBMITTED: June 13, 1960

Card 2/2

LEVINA, R.Ya.; KOSTIN, V.N.; GEMBITSKIY, P.A.; VINOGRADOV, A.D.

Reactions of cyclopropane hydrocarbons with mercury oxide salts.
Part 12: γ -Mercurated alcohols from 1,1-dimethyl-2-alkylcyclo-
propanes. Vest. Mosk. un. Ser. 2:Khim. 16 no.1:67-68 Ja-F '61.
(MIRA 14:4)

1. Kafedra organicheskoy khimii Moskovskogo universiteta.
(Mercury organic compounds)

LEVINA, R.Ya.; KOSTIN, V.N.; GEMBITSKIY, P.A.; SHOSTAKOVSKIY, S.M.

New hydrocarbons of the cyclopropane series. Zhur.ob.khim. 30
no.10:3502-3503 0 '61. (MIRA 14:4)

1. Moskovskiy gosudarstvennyy universitet.
(Cyclopropane)

LEVINA, R. Ya.; KOSTIN, V.N.; GEMBITSKII, P.A.; TRESHCHOVA, Ye. G.

Cyclopropanes and cyclobutanes. part 17: Reduction of arylcyclopropanes by metals and methyl alcohol in liquid ammonia.
Zhur. ob. khim. 31 no.3 (29-836 Mr '61. (MIRA 14:3)

1. Moskovskiy gosudarstvennyy universitet imeni M.V. Lomonosova.
(Cyclopropane)

LEVINA, R. Ya.; KOSTIN, V.N.; GEMBITSKIY, P.A.; SHOSTAKOVSKIY, S.M.;
TRESHCHOVA, Ye.G.

Cyclopropanes and cyclobutanes. Part 18: ρ -Cyclopropylcumene
and β -isopropenylcumene. Zhur. ob. khim. 31 no.4:1185-1190
Ap '61. (MIRA 14:4)

1. Moskovskiy gosudarstvennyy universitet.
(Benzene)

LEVINA, R.Ya.; SHANAZAROV, K.S.; KOSTIN, V.N.

Synthesis of hydrocarbons. Part 77: Stereoisomeric 1, 3, 5-trialkyl-[0, 1, 3]-bicyclohexanes. Zhur.ob.khim. 31 no.6: 1823-1827 Je '61. (MIRA 14:6)

1. Moskovskiy gosudarstvennyy universitet imeni M.V.Lomonosova.
(Bicyclohexane)

LEVINA, R.Ya.; KOSTIN, V.N.; GEMBITSKIY, P.A.; SHOSTAKOVSKIY, S.M.;
TRESHCHOVA, Ye.G.

Cyclopropanes and cyclobutanes. Part 24: Cyclopropylmesitylene.
Zhur.ob.khim. 32 no.5:1377-1382 My '62. (MIRA 15:5)

1. Moskovskiy gosudarstvennyy universitet.
(Mesitylene)

L 8547-65 EWT(1)/EWG(k)/EWT(m)/EEC(t)/EWP(q)/EWP(b) Pz-6 IJP(c)/SSD/
ASD(p)-3/ASD(a)-5/BSL/AFWL/ESD(gs)/ESD(dp)/ESD(t)/RAEM(t)/AS(mp)-2 JD/AT
ACCESSION NR: AR4044073 8/0058/63/000/011/HO45/HO46

SOURCE: Ref. zh. Fizika, Abs. 11Zh363

AUTHOR: Kostin, V. N.; Petrenko, L. A.

TITLE: The influence of ion bombardment on the photoelectric properties of copper

CITED SOURCE: Uch. zap. Khar'kovsk. un-t, v. 127, 1962, Tr. Radiofiz. fak., v. 6, 93-98

TOPIC TAGS: ion bombardment, photoelectric property, copper, counter, control cathode

TRANSLATION: Investigates the influence of ion bombardment on the photoelectric properties of Cu in nonself-quenching hydrogen-filled counters. The design of the counters permits investigation of the influence of bombardment on the working Cu-cathode and simultaneous comparison of it with an identical control cathode which is not bombarded. It is shown that the characteristics of both cathodes under ordinary circumstances change little with time, and that these changes are identical for both cathodes. However, ion bombardment (positive corona) of the working

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L 8547-65

ACCESSION NR: AR4044073

cathode with gradually increasing current values (25-300 μ a) greatly decreased the sensitivity of both cathodes to UV radiation (more sharply for the working cathode than for the control cathode); at a bombardment current of 125 μ a the working cathode ceases to be sensitive to UV radiation. With further voltage increase the coronal discharge abruptly became an intermediate discharge (voltage 1500 v, current 3 ma), and bombardment under these conditions for 2 minutes resulted in a sharp increase of cathode sensitivity to UV radiation. The sensitivity of the control cathode in this case remained as before. The threshold of spectral sensitivity of the control cathode also remained unchanged, while for the working cathode it was displaced by approximately ~ 25 m μ toward higher wavelengths and the work function decreased by 0.4 ev. Thus it was shown that bombardment of a cathode with positive ions decreases its work function.

SUB CODE: IC, NP

ENCL: 00

Card 2/2

LEVINA, R.Ya.; SHANAZAROV, K.S.; KOSTIN, V.N.; TRESHCHOVA, Ye.G.

Synthesis of hydrocarbons. Part 78: New synthesis of 1,1,3,3,4-pentamethylcyclopentane. Zhur.ob.khim. 32 no.8:2637-2640 Ag
'62. (MIRA 15:9)

1. Moskovskiy gosudarstvennyy universitet imeni M.V. Lomonosova.
(Cyclopentane)

LEVINA, R. Ya.; SHANAZAROV, K. S.; TRESHCHOVA, Ye. G.; KOSTIN, V. N.

Synthesis of hydrocarbons. Part 79: Synthesis of stereoisomeric
1,3-dimethyl-5-alkylbicyclo(0,1,3)hexanes and their Raman
spectra. Zhur. ob. khim. 32 no.12:3935-3941 D '62.
(MIRA 16:1)

1. Moskovskiy gosudarstvennyy universitet imeni M. V. Lomonosova.

(Bicyclohexane—Spectra)

LEVINA, R.Ya.; GEMBITSKIY, P.A.; KOSTIN, V.N.; TRESHCHOVA, Ye.G.

Cyclopropanes and cyclobutanes. Part 27: Cyclopropylalkylbenzenes.
Zhur.ob.khim. 33 no.2:359-365 F '63. (MIRA 16:2)

1. Moskovskiy gosudarstvennyy universitet imeni M.V.Lomonosova.
(Benzene) (Cyclopropyl group)

LEVINA, R.Ya.; GEMBITSKIY, P.A.; KOSTIN, V.N.; SHOSTAKOVSKIY, S.M.;
TRESHCHOVA, Ye.G.

Cyclopropanes and cycloputanes. Part 28: p-Acetylphenyl-
cyclopropane in the synthesis of para-substituted cyclopropyl-
benzenes. Zhur.ob.khim. 33 no.2:365-371 F '63. (MIRA 16:2)

1. Moskovskiy gosudarstvennyy universitet imeni M.V.Lomonosova.
(Acetophenone) (Benzene derivatives)

KOSTIN, V.N.

Widely expand the use of oxygen in nonferrous metallurgy. TSvet.
met. 36 no.5:1-4 My '63. (MIRA 16:10)

-KOSTIN, V.N., kand. tekhn. nauk; RYCHKOV, A.I., doktor tekhn. nauk,
prof. [deceased]

1. Heat emission of corrugated heating surfaces when boiling
water. Khim. i neft. mashinostr. no.2:30-32 Ag '64 (MIRA 18:1)

KOSTIN, V.N.; TKACHENKO, V.M.

Effect of a constant magnetic field on a stationary corona discharge. Zhur. tekhn. fiz. 34 no.5:883-887 Ny'64 (MIRA 17:8)

1. Khar'kovskiy gosudarstvennyy unicersitet imeni Gor'kogo.

KOSTIN, V.N.; TKACHENKO, V.M.

Effect of a constant magnetic field on a stationary corona
discharge. Zhur. tekhn. fiz. 34 no.7:1252-1258 J1 '64

(MIRA 17:8)

1. Khar'kovskiy gosudarstvennyy universitet imeni Gor'kogo.

S/120/62/000/001/043/061
E192/E382

7.4160
AUTHORS:

Kostin, V.N. and Tkachenko, V.M.

TITLE:

Operation of photo-cells under pulsed conditions

PERIODICAL:

Pribory i tekhnika eksperimenta, no. 1, 1962,
173 - 176

TEXT:

The operation of gas-filled photo-cells under pulsed conditions offers new possibilities in photo-electronic automation systems. The pulsed operating regime has therefore been investigated in detail by employing argon-filled photo-cells, type УГ-4 (TsG-4), with O-Cs cathodes and vacuum photo-cells, CU5-4 (STsV-4), with Sb-Cs cathodes (manufactured at Moskovskiy elektrozavod (Moscow Electrical Works)). The spectral characteristics of these cells extend over the visible range so that a 6 V battery-fed lamp was used as the illumination source. The intensity of the illumination was controlled by varying the current. The cell and the lamp were enclosed in a special envelope to eliminate the outside radiation of the cell. The cell was connected to an amplifier head (type БГС (BGS)) through

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a spark gap consisting of a plate and a point. The width of the spark gap could be varied and measured with an error of ± 0.005 mm since the point was mounted on a micrometer screw. The plate was in the form of a solid metal slab. The basic circuit for the investigation is illustrated in Fig. 2, where $\frac{1}{4}$ is the spark gap. The positive terminal of a high-voltage stabilized supply source was connected to the cell through the spark gap and a quenching resistor of $2\text{ M}\Omega$. The negative terminal of the supply source and the cathode of the photo-cell were grounded. The pulses produced in the circuit were applied to the first amplifier tube through a coupling condenser of 47 pF . The amplifier head was connected to a counter by means of a screened cable. The operation of the photo-cells was investigated by measuring their electrical and photometric characteristics. It is concluded from the investigation that:

- 1) the number-of-pulses versus voltage characteristics of the photo-cells under pulsed conditions are qualitatively comparable with the voltage-current characteristics under normal DC

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conditions;

2) the photo-cells can be used photometrically under pulsed conditions since the number of output pulses is linearly dependent on the illumination;

3) the absolute sensitivity under pulsed conditions is comparable with that of the DC operation but the threshold sensitivity is at least 20 times higher;

4) the pulsed operation has the advantage of the absence of dark currents, a high pulsed voltage, possibility of a direct telemetry transmission and a high breakdown voltage.

There are 7 figures.

ASSOCIATION: Khar'kovskiy gosudarstvennyy universitet
(Khar'kov State University)

SUBMITTED: June 30, 1961

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ACCESSION NR: AP4017599

S/0109/64/009/002/0293/0299

AUTHOR: Tkachenko, V. M.; Kostin, V. N.; Belous, V. V.

TITLE: Effect of a high-frequency electric field on the glow discharge

SOURCE: Radiotekhnika i elektronika, v. 9, no. 2, 1964, 293-299

TOPIC TAGS: glow discharge, glow discharge column, glow discharge column compression, hf glow discharge column compression, glow discharge in air, glow discharge in hydrogen, glow discharge in deuterium

ABSTRACT: A theoretico-experimental investigation of the effect of an axially-symmetric h-f field produced by a set of rings on the glow discharge for various field frequencies, discharge currents, gases, and gas pressures is reported. A formula is developed for the diameter of the compressed-discharge column which depends on the field frequency, electron temperature, discharge-tube radius, and the frequency of collisions between ions and gas molecules (i. e. , ion temperature

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