

62-58-1-1 -3 AUTHOR: Nechiporenko, G. N. TITLE: The Determination of Sulfate Ions by Means of the Method of Direct Titration by Lead Nitrate With Dithiazone as Indicator (Opredeleniye sullfatnykh ionov metodom pryamuje titr vaniya azotnokislym svintsom s ditizonom v kachestve irdi katora) Izvestiya Akademii Nauk SSSR, Otdeleniye Khimicheskikh Na.k PERICDICAL: 1950, Ur 3, pp. 309 - 361 (USSR) A STRACT: In the methods of the direct titration lead salts are to be preferred, for, in comparison to barrum salts they possess some advantages. Analytical reagents exist which are much more sensitive to lead ions than to barium. The solubility of PbSC can, however, be considerably reduced by addition of a  $30^4$  - 40 % alcohol or acetone. In this paper the authors suggest a method for the determination of  $SC_{\lambda}^{\frac{N}{2}}$ . Lithian re is a weak two-basic acid (see scheme). When dithiazone is Card 1/2 designated with  $\mathrm{DH}_2$  , then it would read

62-58-3-18/30

The Determination of Sulfate Ions by Means of the Method of Direct Titration by Lead Nitrate With Dithiazone as Indicator

DH  $\rightarrow$  DH  $^+$  H  $^+$   $\rightarrow$  D<sup>2-+</sup>  $\downarrow$   $^+$ . The ions of lead react with DH2 under the simultaneous formation of an inner-complex compound Pb<sup>2+</sup> + 2DH  $\rightarrow$  Pb(DH)<sub>2</sub>. (The conclusions: see equations (1) and (2), (3)). The suggested method offers the possibility to determine in the sample of 50 to 0,5 mg sulfate ions with a precision of 1 - 2 %. This method is usuable for the rapid and exact determination of sulfates in natural waters but only when the chlorides do not more than seven-fold exceed the quantity of the sulfates. There are 3 tables and 3 references, 1 of which is Soviet.

ASSOCIATION: Gidrokhimisheskiy institut Akademii nauk SSSR

(Hydrochemical Institute, AS USSR)

SUBMITTED: October 7, 1957

Card 2/2

# NECHIPORENKO, G.H. Role of organic matter in the determination of trace elements in natural waters. Gidrokhim.mat. 28:165-169 '59. (MIRA 12:9)

1. Gidrokhimicheskiy institut Akademii nauk SSSR, g. Movecherkassk. (Organic matter) (Water-Analysis) (Trace elements)

NECHIPOREMEO, G.M.; KRIVENTSOV, M.I.

Trilonometric determination of small quantities of sulfate ions in water. Gidrokhim.mat. 29:211-213 '59.

(MIRA 13:5)

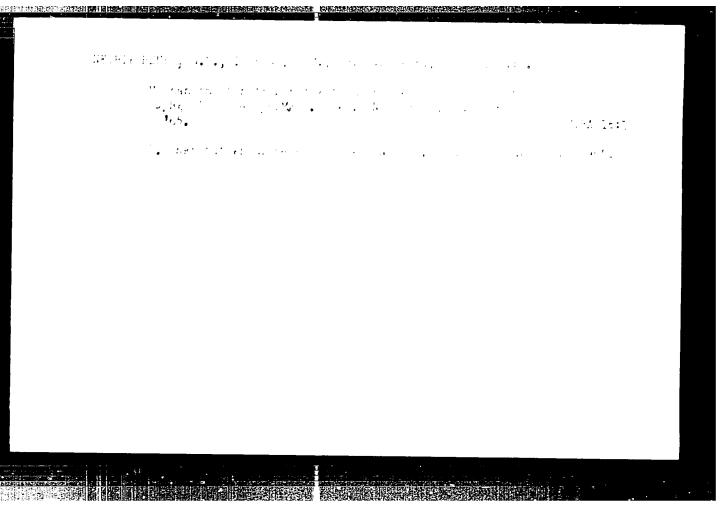
1. Gidrokhimicheskiy institut Akademii nauk SSSR, Novocherkussk.

(Mineral waters--Analysis) (Sulfates)

NECHIPORENKO, G.N.

Determining sulfate ions by direct titration with lead nitrate in the presence of diphenylcarbasone as the indicator. Gidrokhim.mat. 29:214-218 '59. (MIRA 13:5)

1. Gidrokhimicheskiy institut Akademii nauk SSSR, Novocherkassk. (Water--Analysis) (Sulfate) (Lead nitrate)



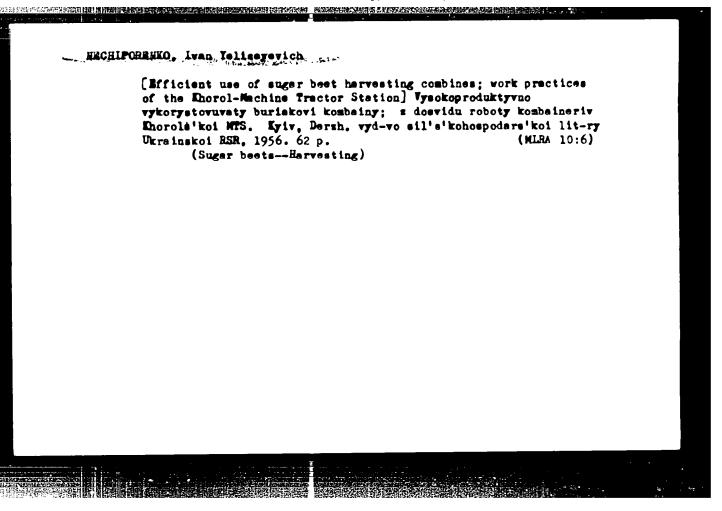
APPROVED FOR RELEASE: Wednesday, June 21, 2000 CIA-RDP86-00513R001136

NECHIPOREURO, G. V.

Horse Breeding

Results of the State stud stables in 1951, and the quota for 1952. Konevodstvo 22 No. 6, 1952

9. Monthly List of Russian Accessions, Library of Congress, September 1958, Unclassified.

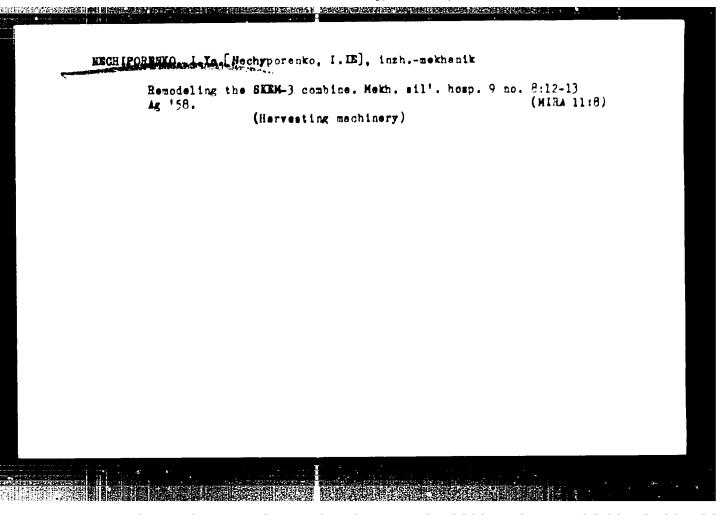


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becally or surposing blodes of the SK-2.5 combine. Mexh. sill nosp. 9 no. 7:7 J. 'Nr. (Mida 11:8)

(Maintenance and repair)

(Combines (Agricultural machinery))
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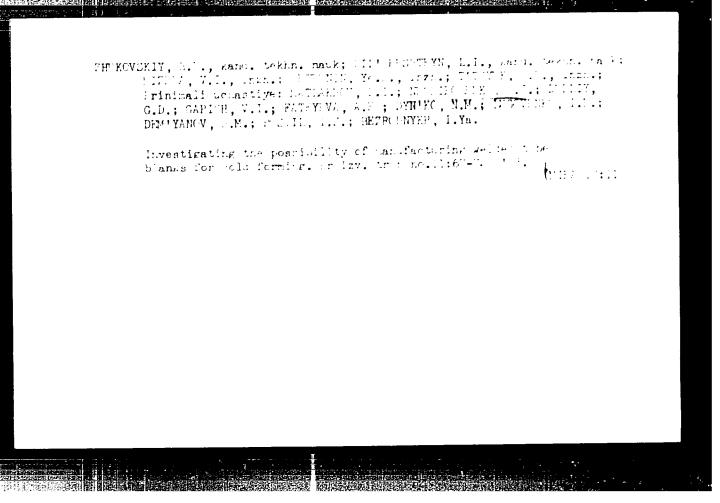
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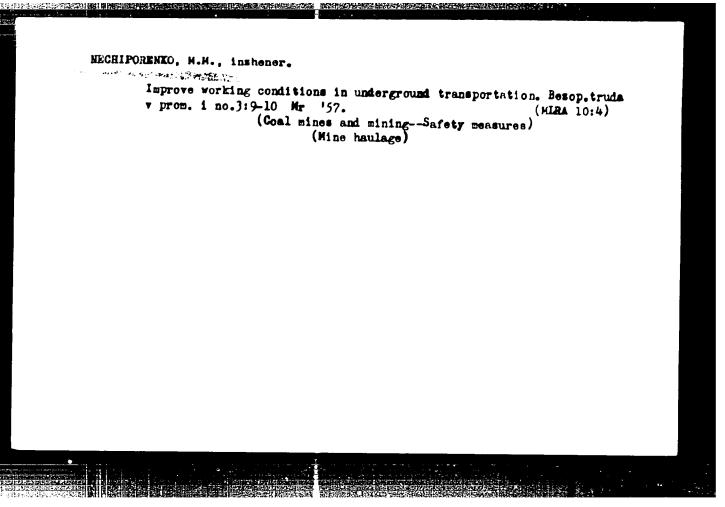
AMINOVA, M.G.; MERFINGERK, J.D., Stock t, J.D.; TRIPTMIV, M.D.

JEFELYOIN, V.V., zand. red. trank, otv. red.

Laibliography of the scientific papers of the Institute for a
1938 to 1961; Bioliografia nauchnykn recot institute za jerriod 1938-1991; g., Frunze, 1991. 77; j. 1991. A 201.

1. Kirgizskiy nauchno-issledovatel'skiy institut ejiter i
logii, mikrochilegii i riglyeny. Z. idrektor Kirgizskere
nauchno-issledovatel'skogo institute ejiteriologii, mikrobiologii i giglyeny (for Jerelygin).



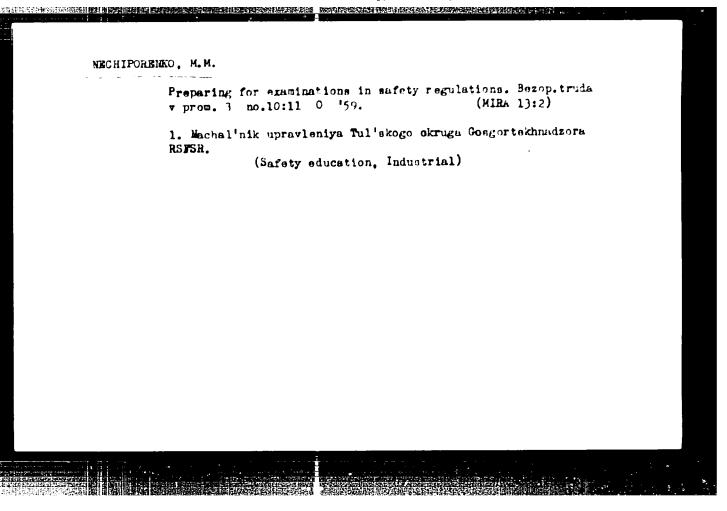


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SAVOSTIN, G.A., inzh.; TERESHCHENKO, F.P., inzh.; NECHIPORENKO, M.M.; SAMOTETEV, G.V.; DEHIRHOV, I., inzh.

Concerning the article "Increase cross sections of raulagevage"
Bezop.truda v prom. 2 no.4:22-24 Ap '59. (MIRA 11:4)

1. Institut "Krivbassproyekt" (for Savostin, Tereshchenko). 2.Upravleniye Tul'ekogo okruga Gosgortekhnadzora SSSR (for Nechiporenko, Samotayav).

(Mining engineering)
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HECHIFORENKO, M.M., ingh.; DOLOTOV, N.P., ingh.

Safety measures in mines of the Moscow Basin. Bezop.truda. v prom.
4 no.6:25-27 Je '60. (Mira 14:3)

1. Uprawleniye Tul'skogo okruga Gosgortekhnadzora RSFSR.

(Moscow Basin—Goal mines and mining—Safety measures)

APPROVED FOR RELEASE: Wednesday, June 21, 2000 CIA-RDP86-00513R001136

NECHIPORENKO, M.M.; DOLOTOV, N.P., inzh.; SUBBOTIN, A.A., Geroy Sotsialistiches-kogo truda; PERMYAKOV, P.N., laurest Leninskoy premii

Effective methods for improving work sanitation in mining. Bezop.truda v prom. 6 no.7:4-6 Jl '62. (MIRA 15:7)

ALCOHOLOGICA STATEMENT OF THE STATEMENT

1. Nachal'nik Upravleniya Tul'skogo okruga Gosudarstvennogo komiteta pri Sovete Ministrov RSFSR po nadzoru za bezopasnym vedeniyem rabot v promyshlennosti i gornomu madzoru (for Nechiporenko). 2. Nachal'nik Tul'skogo kombinata ugol'noy promyshlennosti Podmoskovskogo basseyna Ministerstva ugol'noy promyshlennosti SSSR (for Subbotin). 3. Glavnyy inzh. Tul'skogo kombinata ugol'noy promyshlennosti Podmoskovskogo basseyna Ministerstva ugol'noy promyshlennosti SSSR (for Permyakov).

(Tula Province—Coal mines and mining—Safety measures)

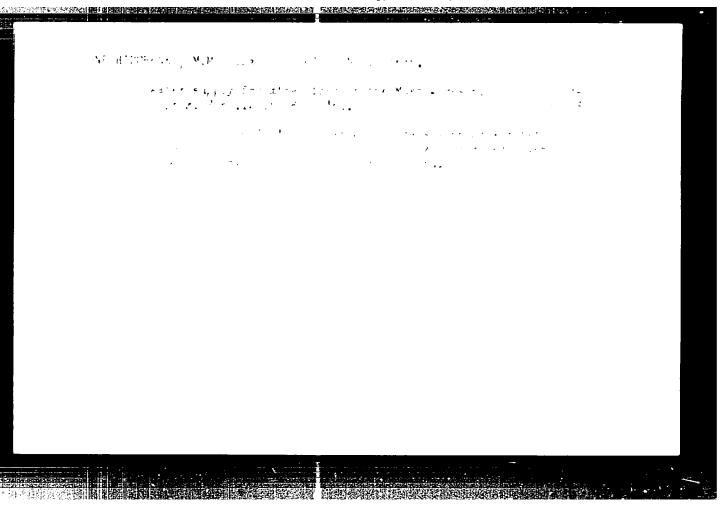
SUBBOTIN, A.A., Gerov Sotsialisticheskogo Truda; PERMYAKOV, P.W., laureat Leninskoy premii; NECHIPORENKO, M.M.; DOLOTOV, H.P.

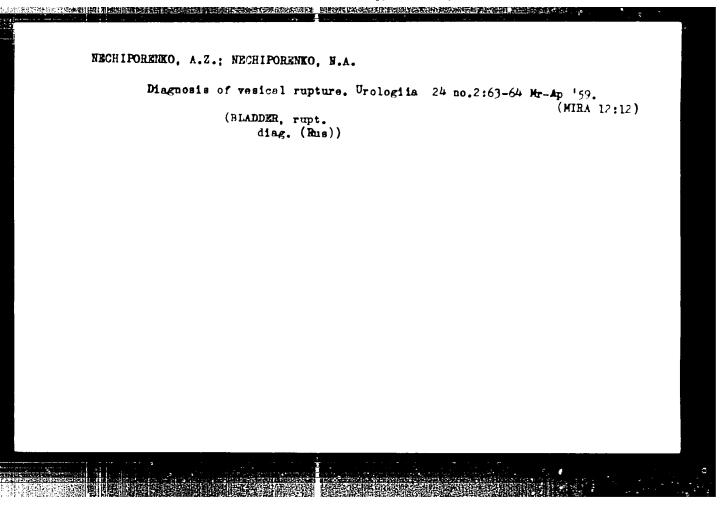
Mechanization and automation in mines of the Priokskiy Economic Council. Bezop. ruda v prom. 7 no.432-3 Ap 163.

(MIRA 16:4)

1. Machal'nik Tul'skogo kombinata ugol'noy promyshlennosti
Podmoskovnogo basseyna Ministerstva ugol'noy promyshlennosti
SSSR (for Subbotin). 2. Glavnyy inzh. Tul'skogo kombinata
ugol'noy promyshlennosti Podmoskovnogo basseyna Ministerstva
ugol'noy promyshlennosti SSSR (for Permyakov). 3. Machal'nik
Upravleniya Tul'skogo okruga Gosudarstvennogo komiteta pri
Sovete Ministrov RSFSR po nadzoru za bezopasnym vedeniyem rabot
v promyshlennosti i gornomu nadzoru (for Nechiporenko). 4. Glavnyy inzh. Upravleniya Tul'skogo okruga Gosudarstvennogo komiteta
pri Sovete Ministrov RSFSR po nadzoru za bezopasnym vedeniyem
rabot v promyshlennosti i gornomu nadzoru (for Dolotov).

(Tula Province—Coal mines and mining)
(Automation)





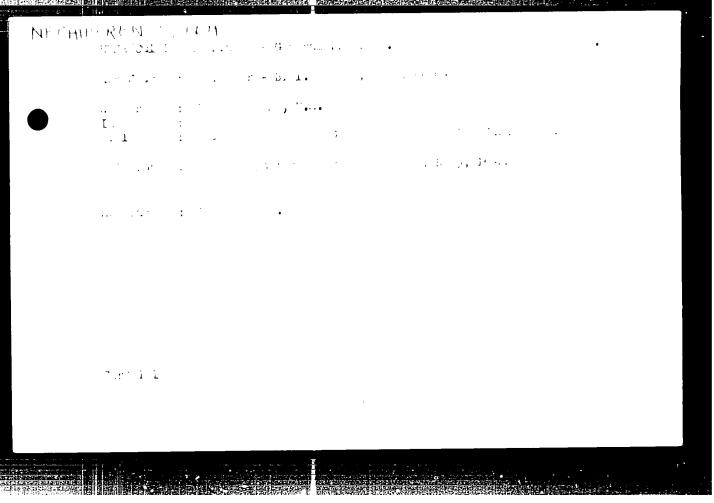
- 1. NECHIPORENKO, N. A.
- 2. USSR (600)
- 4. Siberia, Eastern-Agriculture
- Achievements of agricultural science applied on the collective farm. Dost. sel'khoz. No. 2, 1952.

9. Monthly List of Russian Accessions, Library of Congress, January 1953, Unclassified.

- 1. TSEDIK\_TOMASHEVICH, Z. F.; NECHIPORENKO, N. A.
- 2. USSR (600)
- 4. Agriculture Experimentation
- 7. Work results in scientific research institutes on agriculture for 1951.

  Dost. sel'khoz. no. 5, 1952

9. Monthly List of Russian Accessions, Library of Congress, \_\_\_\_\_\_\_\_1953, Unclassified.

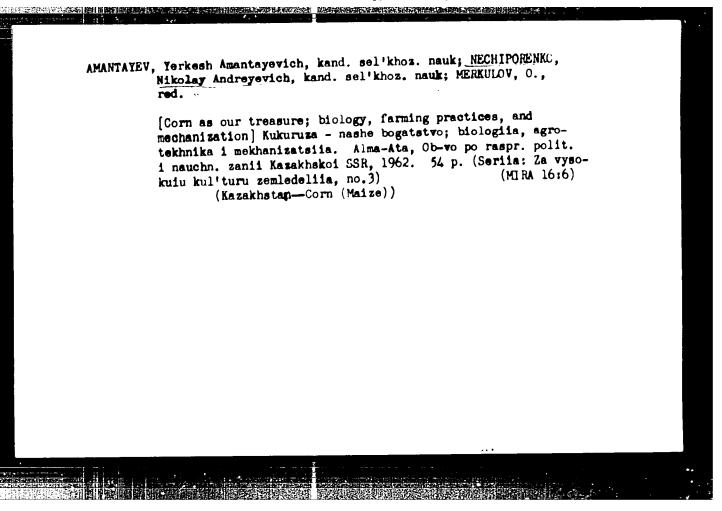


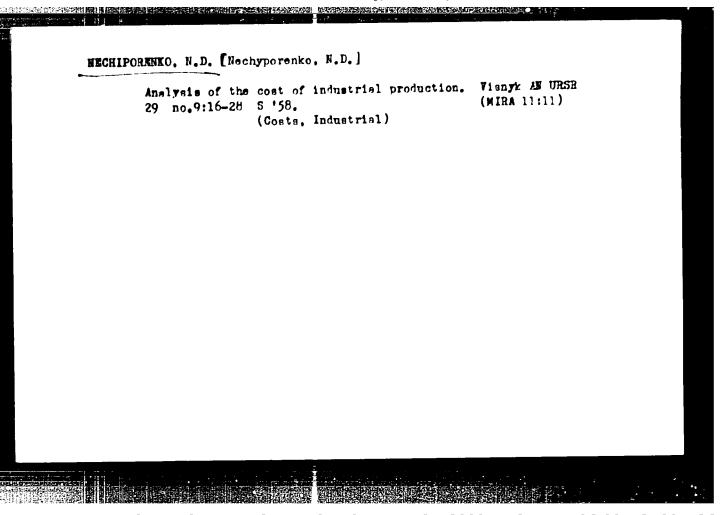
BIYASHEV, G.Z., akademik; NECHIPORENKO, N.A., FEDUROV, P.F., kand.sel'skokhozyaystvennykh nauk; AMANTAYEV, Ye.A., kand.sel'skokhozyaystvennykh nauk

Most important problems in the agriculture of southern and
southeastern Kazakhstan. Zemledelie 23 no.4:8-14 ip '61. (MPA 1/:3)

1. Kazakhskaya akademiya sel'skokhozyaystvennykh nauk (for Biyashev).
2. Chlen-korrespondent Kazakhskey akademii sel'skokhozyaystvennykh
nauk (for Nechiporenko).

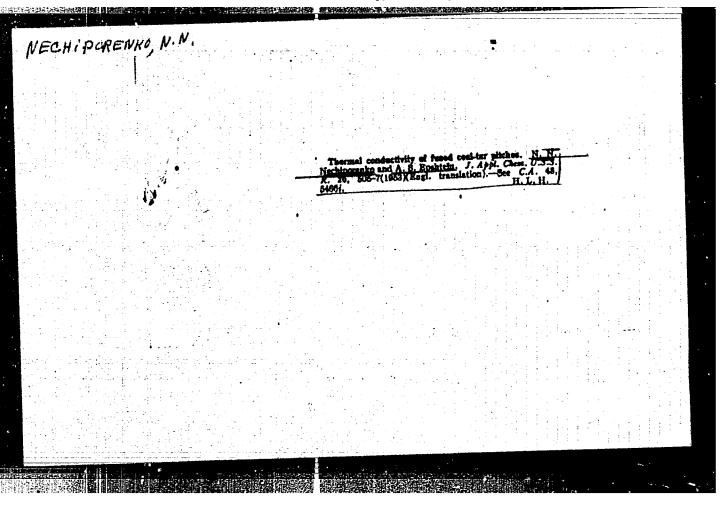
(Kazakhstan-Agriculture)

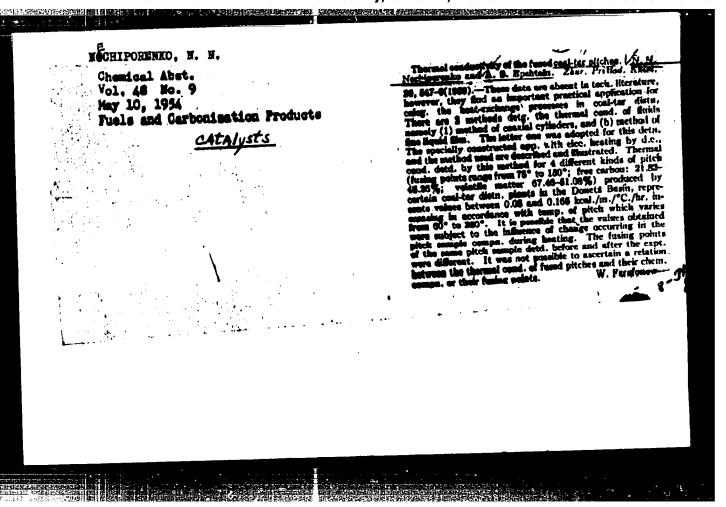




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NECHIPOTENCO, N.N.
MINENEO, V.I.; TSARIKHIN, D.A.; HECHIPOTENEO, N.N.; PUSTOVALOV, V.I.;

SPRISHEVSKIY, A.I.

Method of insulating suspension devices for galvanising parts.
Avt.trakt.pros. no.10:29 0 '54. (MIRA 7:10)

1. Ehar'kovskiy velosipednyy savod.
(Galvanising)

SOV/137-57-10-19918

Translation from Referativnyy zhurnal, Metallurgiya, 1957, Nr 10, p 208 (USSR)

AUTHORS Minenko, V.I., Nechiporenko, N.N.

TITLE

A Method of Insulating Suspension Fixtures for Nickel, Chrome, Copper, Zinc, and Other Plating Procedures (Sposob izolyatsii podvesnykh prisposobleniy dlya nikelirovaniya, khromirovaniya, medneniya, tsinkovaniya i drugikh gallvanopokrytiy detaley)

PERIODICAL Tr Khar kovsk inzh. ekon in ta 195t Vol 7 pp 135-138

ABSTRACT

A paste (P) consisting of a polychlorvinyl resin - igelite - with added plasticizers and stabilizers is applied to the surface of the suspension (S) device. The P is then polymerized by heat treatment in a drying cabinet. To improve the strength of the bond of the insulating P and the surface of the S, the latter are covered with chemically stable primer before the application of the insulating P. The raw materials used for insulation may be igelite—tech.dibutylphthalate, Pb or Zn stearate, and a chemically resistant primer KhSG-26. The insulation process proceeds via the following stages—a) Degreasing the S by the

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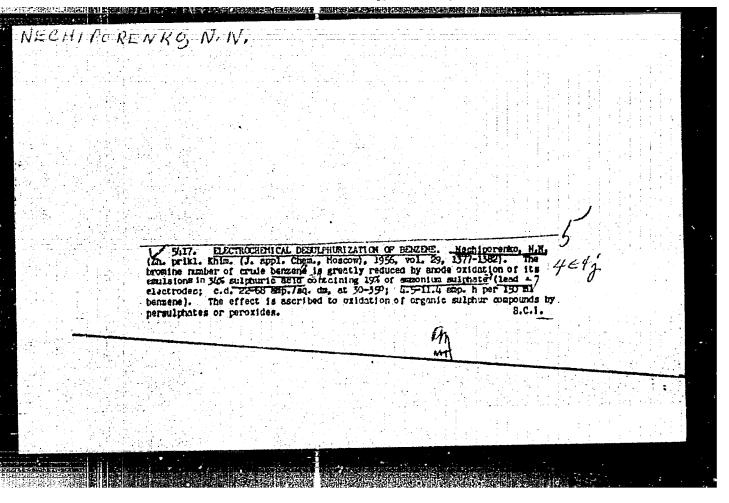
A Method of Insulating Suspension Fixtures (cont.)

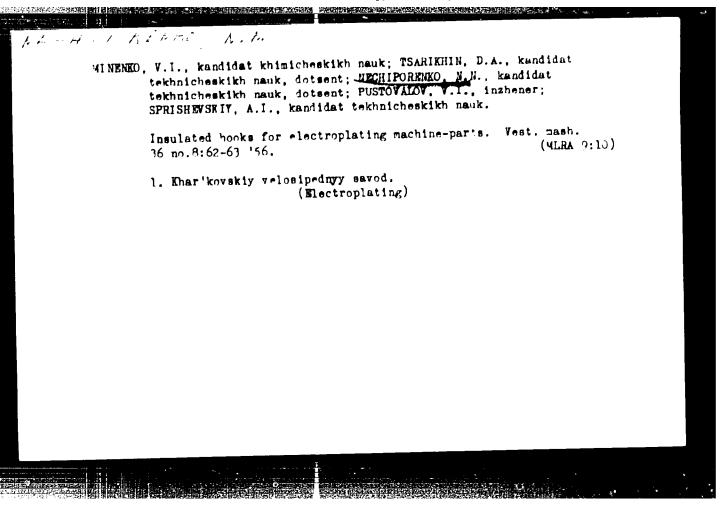
method used to prepare the parts for plating; b) coating with KhSG-26 primer, c) drying the primer in a drying cabinet, d) application of the P, e) heat treatment of the P, f) dressing the contact ends. After priming and drying, the S are immersed in a pan with the insulating P. The P is prepared as follows. The components are weighed out on a basis of 6 parts dibutylphthalate and 0.16 parts Pb or Zn stearate per weight to 10 parts igelite. The pulverized substances are thoroughly mixed, the dibutylphthalate is added, and the mixture is ground until a homogeneous P of creamy consistency is obtained. After standing for 30 to 50 minutes (to ripen) at room temperature, it is applied to the S. The insulating P is prepared in small quantities calculated for use in the next 2 or 3 hours. After the excess P has flowed off, the S, hung from racks, are placed in the drying cabinet and kept there for 20-30 min at 130-200°C.

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\$/068/60/000/003/001/003 E071/E233

AUTHORS: Nechiporenko, N. N. and Manoylenko, B. R.

TITLE: Oxidative Desulphurisation of Benzole PERIODICAL: Koks i khimiya, 1960, No. 3, pp. 37-42

TEXT: The possibility of desulphurising benzole by oxidation of sulphurous compounds to oxygen derivatives of sulphur by active or activated oxygen, obtained electrochemically or catalytically was investigated. The results of this investigation are described in the paper. The experiments were made with pure benzene to which either 0.91 or 0.50% of thiophene were added. The diagram of the apparatus for the anodic oxidation of thiophene in benzole is shown in Fig. 1. The anode and cathode were separated by a porous diaphragm made from Schott filter. The anode and cathode were made from platinum strip of a surface area of 12.6 and 15.5 cm² respectively. The anode current density was varied from 0.2 to 0.94 A/cm² and the electrolytic temperature 25-27 and 29-30°C respectively. The sulphur containing benzole was fed continuously (through the bottom of the vessel) into the anode section while the electrolyte (340 g/l of (NH4)2SO4 and 80 g/l of

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S/068/60/000/003/001/003 E071/E233

Oxidative Desulphurisation of Benzole

H2SO4) was fed from the top into the cathode section of the electrolyser. The benzole and electrolyte in the anode section were continuously stirred. The gas evolved during the electrolysis together with benzole vapours was passed through a condenser, where benzene was condensed and returned into the electrolyser. The electrolytically treated benzene was washed with a 20% solution of sodium hydroxide and redistilled. The degree of desulphurisation obtained was measured by the bromine number (Ref. 11). The experimental results are given in Table 1. The best results were obtained at a current density of 0.47 A/cm2 when up to 50% desulphurisation was obtained. A further increase in the current density (0.94 A/cm2) leads to a decrease in the effectiveness of the process. An addition of 5 g/l of sodium coloride (experiments 11-13) improved considerably the desulphurising effect. In the experiments 14 and 15 an electrolyte containing 100 g/l of  $(NH_4)_2SO_4$ , 550 g/l of  $H_2SO_4$  and 50 g/l of  $MnSO_4$  also gave good desulphurising results. However, in the latter case the formation of manganic acid was observed, so that a partial Card 2/4

3/008/60/000/003/001/003 E071/E233

Oxidative Desulphurisation of Benzole

oxidation of thiophene due to secondary reactions is possible. Oxidation of thiophene in benzole by oxidants at 85°C was also tried. The experimental procedure consisted of bubbling benzole vapours through a layer of (165 mm nigh) acid solutions of ammonium persulphate, potassium bichromate, potassium permanganate and hydrogen peroxide. The experimental results confirm that, in principle, the desulphurisation of benzole by this method is possible. Oxidising catalytic desulphurisation of benzole was tested by passing air-benzole mixture through a furnace heated to 250-380°C filled with various catalysts. As catalysts the following substances were used: 1) Pretreated activated carbon. The pretreatment consisted of extraction of silica with fluoride compounds, saturation with a solution of ferrous sulphate, precipitation of ferrous hydroxide with ammonia and ignition of the contact mass at 600°C. The product obtained contained about 1% of Fe<sub>2</sub>O<sub>3</sub>. 2) The second type of catalyst was made from Chasov-Yar clay by saturation with ferric nitrate and ferric hydroxide precipitated with ammonia, washed from alkali ions and ignited at

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Oxidative Desulphurisation of Benzole

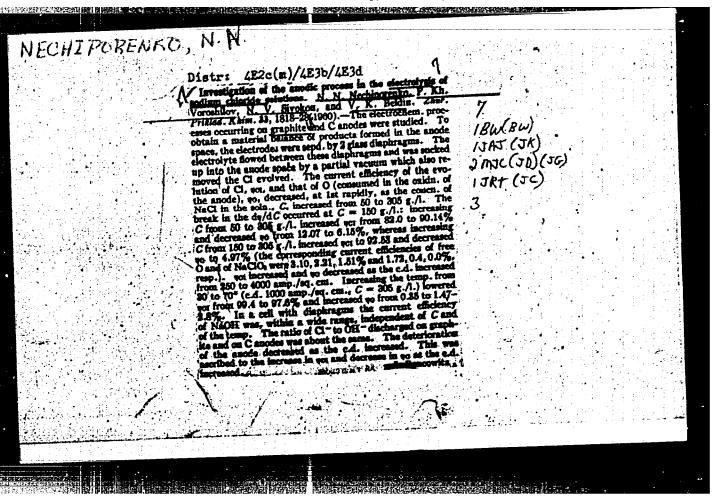
600°C. The catalyst contained about 10% of ferric oxide. This catalyst was also made in two additional modifications: a) containing phosphates and b) phosphates and a homogeneous oxygen transferring medium (not specified). In all experiments 25 ml of the catalyst was placed into the furnace; velocity of air stream - 500 ml/min, and of benzole (containing 0.5% of thio; hene) 75 ml/hr. The experimental results are given in Table 3. The best results (practically complete elimination of thiophene) were obtained with the clay catalyst activated with phosphates and containing some homogeneous oxygen transferring agents. It was also confirmed that on catalytic oxidation thiophene is completely oxidised to sulphuric acid. It is concluded that the possibility of oxidising resistant sulphurous organic compounds was proved in principle and that the catalytic method is most effective. The process however, requires further studies on a larger scale. There are 3 figures, tables and 13 references: 11 Soviet and 2 non-Soviet.

ASSOCIATION:

Khar kovskiy politekhnicheskiy institut

(Kharkov Polytechnical Institute)

Card 4/4



5/068/62/000/012/001/001 E071/E436

AUTHORS:

Nechiporenko, N.N., Kakulin, G.P., Fedorchenko, I.G.,

Manoylenko, B.R.

TITLE:

An investigation of the process of chlorination of

thiophene

PERIODICAL: Koks i khimiya, no.12, 1962, 43-45

TEXT: In view of the possibility of applying chlorine for the production of a high purity benzene, the authors investigated the process of chlorination of thiophene dissolved in benzene in order to establish the necessary amount of chlorine for a complete purification of benzole from thiophene. In addition, the influence of temperature and velocity of supply of chlorine to the reactor on the degree of purification of benzole with a given thiophene content was studied. The apparatus consisted of a reactor fitted with a mercury sealed stirrer, thermometer and inlet and outlet for chlorine. The outlet gases (air and traces of chlorine) were scrubbed in a solution of potassium iodide, crystalline sodium hydroxide (for HCl) and activated carbon (for benzene vapours). A cryoscopic benzene with an addition of 1% of thiophene was used Card 1/2

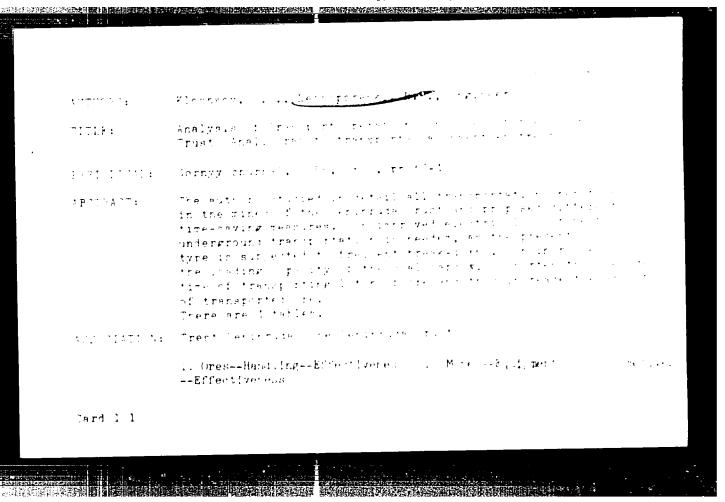
An investigation of the process ... S/C68/62/000/012/001/CC1

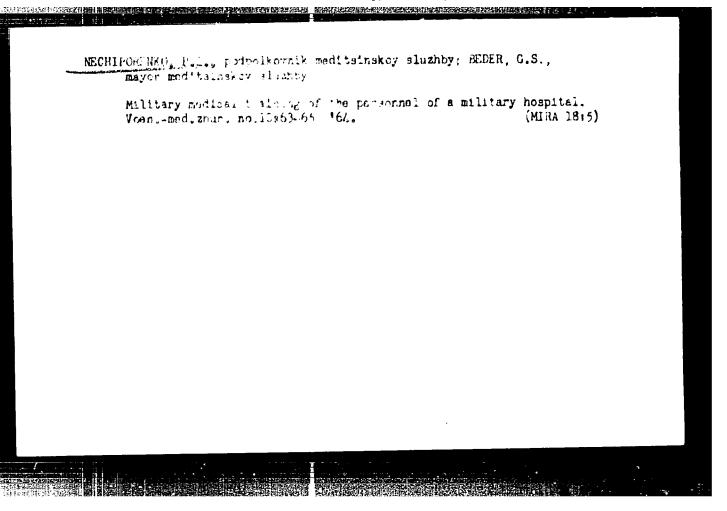
for experiments. The purification process was followed by the bromine number, determined by the bromide-bromate method. It was established that the degree of purification of benzole depends mainly on the amount of the reagent used and is practically independent of temperature (7 to 40°C) and rate of supply of chlorine. Refining with chlorine can produce a product practically free from thiophene. For a complete purification of benzole from thiophene, it is necessary to use 1.5 to 2.0 weight units of chlorine per 1 weight unit of thiophene. There are 1 figure and 3 tables.

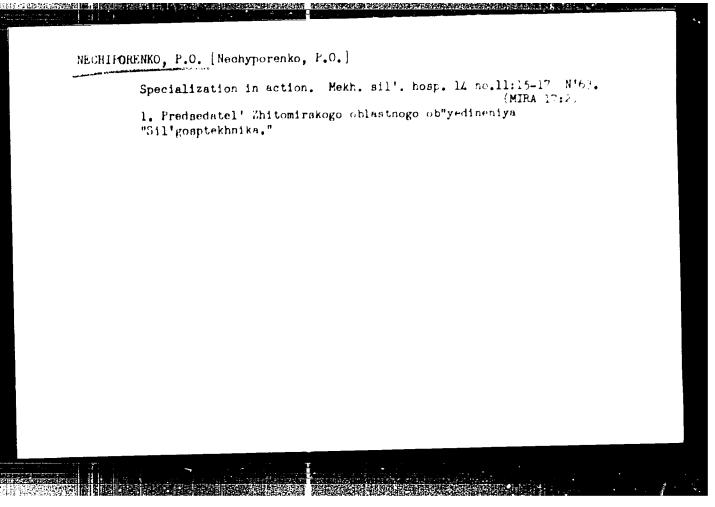
ASSOCIATION: Khar'kovskiy politekhnicheskiy institut (Khar'kov Polytechnic Institute)

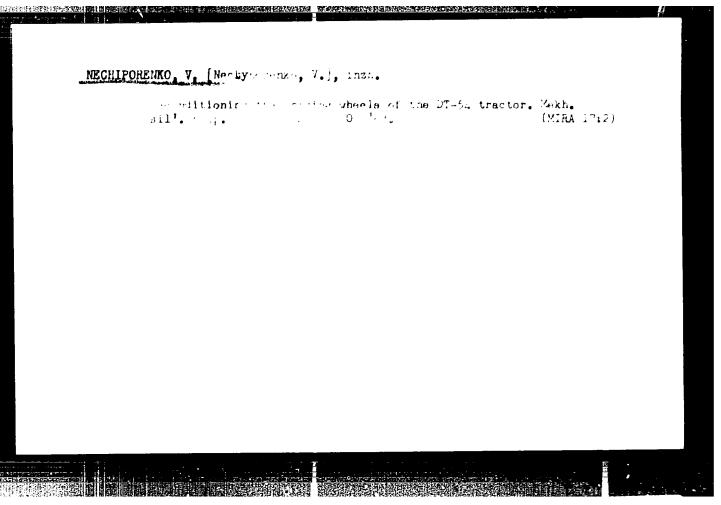
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# NECHIPORENKO, V.A.

Device for experimental study of sliding journal bearings. Trudy

[MIRA 15:2]

l. Kafedra detaley mashin i pod\*yemno-transportnykh mashin Leningradskogo korablestroitel'nogo instituta. (Marine engineering)

1413 1418 2808 2208

S/185/60/005/003/012/02C D274/D303

18 8100

26597

AUTHORS:

Gridnyev, V.N. and Mechyporenko, V.G.

TITLE:

Phase transformations during electrical heating of

manganese-vanadium steels

PERIODICAL:

Ukrayins'kyy fizychnyy zhurnal, v. 5, no. 5, 1960,

402-406

The position of the critical points was investigated as a function of the rate of heating and initial structure of low-carbon and high-carbon manganese-vanadium steels. The investigations were carried out over a range of 80 - 30,000 deg/sec; (the tiret time that such a range was investigated). The alloying constituents were chosen in such a way so as to reduce as much as possible the influence of the carbon on the phase transition process. The method of investigation is discussed. The specimens were in the form of vires 0.8 x 1.5 mm. The composition of the specimens is given in a table; the constituents are: C, V, Mn. Si, Gr, W. The steels

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Phase transformations...

were investigated in three initial states: annealed, tempered, and cold deformed. Annealing was carried out at 1050°C for 60 minutes; tempering at 1230°C for 45 min. The specimens were directly heated by alternating current of 1000 cy in a setup for the complex investigations of phase transitions as per V.N. Gridnyev and V... Cherepin (Ref. 1: Zavodskaya laboratoriya, no. 3, 1957). A figure shows the oscillograms for specimen no. 3 in the annealed state at rates of 500, 7000 and 30,000 deg/sec respectively. It is noted that the position of the Curie point can be very clearly distinguished on the dilatometric curves, and to a lesser degree - on the thermal curves. It was established by a special investigation that, with the given method of registration, reliable results in determining the position of critical points can be obtained for rates of heating which do not exceed 3000 - 4000 deg/sec. At higher rates, the readings are affected by the inertia of the system. It was found possible, however, to correct the position of the critical points at ultrahigh rates of heating by means o: the position o: the Curie point; such a correction can be carried out only if the Curie point

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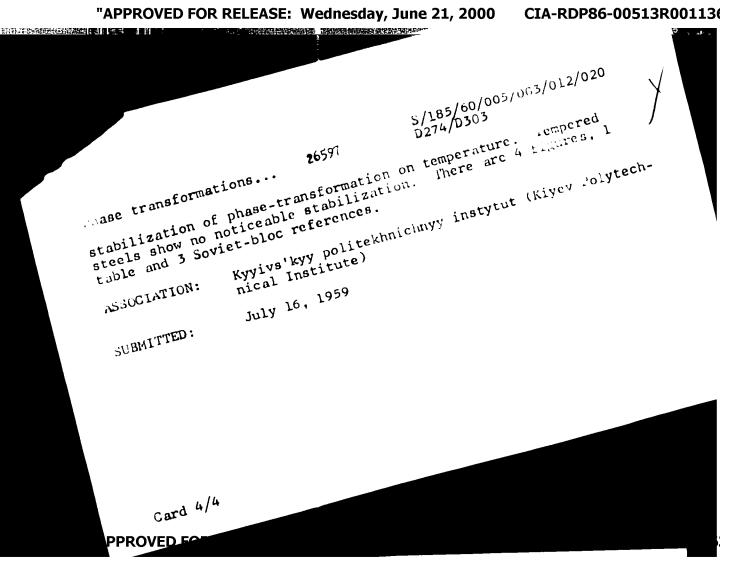
Phase transformations... 26597

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lies below the temperature of phase transition. In all cases (irrespective of initial state) the position of the critical points becomes higher with increasing rate of heating. The displacement of the critical points is especially pronounced in the case of annealed steels. some of the curves show a stabilization of phasetransformation temperature at very high rates of heating. The position of the critical points of tempered steels changed so chow unexpectedly; whereas, according to earlier results, the critical points of tempered steels lie below those of annealed steels, the present investigation showed that at relatively low rates of heating (up to 2000 - 3000 deg/sec) the critical points of temered steels lie above those of annealed steels; at high rates or heating, the present results do not disagree with the earlier results. The temperature of phase transformation of cold-deformed steels is considerably lower than that of tempered and annealed steels; this agrees with earlier results. For high-carbon steels, the temperature of phase transformations of tempered steels is considerably nigher than that of annealed steels. Cold-deformed steels show

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# "APPROVED FOR RELEASE: Wednesday, June 21, 2000



8/137/62/000/004/089/201 A052/A101

187500

AUTHOR:

Nechiporenko, V. O.

TITLE:

Magnetometric investigation of the residual austenite decomposition

at electric heating

PERIODICAL:

Referativnyy zhurnal, Metallurgiya, no. 4, 1962, 15, abstract 4193 ("Sb. nauchn. tr. aspirantov Kiyevsk. politekhn. in-ta. Kiyev",

1961, 203-211)

Wire samples of steel containing 1.19% C, 1.86% V, 1.74% Mn and 0.70% Si were water hardened at 1,230°C; a part of the samples was cold treated at -78°C. The effect of tempering at the rates of heating of 30 - 5,000 degree/ sec on the behavior of residual austenite was investigated by the magnetometric dilatometric and thermal methods. At all rates of heating volume effects corresponding to the first and third temper transformations are observed on dilatometric curves. Absolute values of these effects are considerably lower than for carbon steel. The dilatometric effect of the residual austenite decomposition is not observed. As the rate of heating increases, the first and third transformations shift into the region of higher temperatures. At heating to the

Card 1/2

UR RELEASE: Wednesday, June 21, 2000 CIA-RDP86-00513R001136

PARTICIPATION OF THE PROPERTY 5/148/61/000/006/007/013 26583 18 1500 E071/E480 ALTHORS . Gridney, V.N. and Nechiporenko, V.G. TITLE: Transformations in vanadium-manganese steel during electric annealing FARIODICAL: Izvestiya vysshikh uchernykh zavedeniy. Chernava metallurgiya, 1961, No. (. Fr. 111-118 TEXT As not many data on the behaviour of allow steels during lectric annealing are available, the authors investigated the of luence of alloying elements on the acchanism and kinetics of ocesses taking place during electric annealing of vanadiuminganese steel within a wide range of heating rates (20  $\pm c$ -000°C/sec). It was expected that the presence of the proble rming element would slow down the in emposition of marriantite, orticularly at high heating rates. The addition of many mese could have facilitated the observation of intermediate structural ates. The experimental steel (0.71% () -1.79% V, 1.40% Mm and 50% Si) was melted from Armco iron in a high frequency formace. v ingot (3 kg) was forged and drawn into wires of 0.5 and 1.5 mm meter. Wire specimens 120 mm long were hardened from +220°C in 11/4

Transformations in vanadium ... E071/E480

from high temperatures. There are 6 figures and 10 references:
9 Soviet and 1 non-Soviet. The reference to an English language
Journal of the Iron and Steel Inst., v.188, p.1, 1958.

ASSOCIATION: Kiyevskiy politekhnicheskiy institut
(Kiyev Polytechnical Institute)

SUBMITTED: May 12, 1960

Card 4/4

26583

Transformations in vanadium ...

S/148/61/000/006/007/013 E071/E480

subsequent quenching in water. After such treatment, the coercive force and Vickers hardness (P = 10 kg) of the specimens were measured. The phase composition of the annealed product was studied by X-ray analysis and by the differential magnetic method in saturation fields. It was found that: 1) Volume effects, corresponding to effects I and III of martensite decomposition were observed on the dilatometric curves at all heating rates investigated. Thus the process of martensite decomposition could not be supressed. 2) With increasing heating rate, the temperature of I and III transformations are shifted to higher With increasing heating rate, a continuous decrease of the volume effect of transformation I and an increase of the volume effect of transformation III take place. 3) During electric heating, the formation of austenite takes place in the temperature range of transformation III. On the basis of the dilatometric, magnetic and X-ray data, a partial reverse transformation of martensite into austenite was postulated. On reheating, the austenite formed decomposed at lower temperatures than the residual austenite obtained after hardening Card 3/4



Transformations in vanadium ...

S/148/61/000/006/007/013 E071/E480

from high temperatures. There are 6 figures and 10 references:
9 Soviet and 1 non-Soviet. The reference to an English language
publication reads as follows: K.Seal, R.W.K.Honeycombe,
Journal of the Iron and Steel Inst., v.188, p.1, 1958.

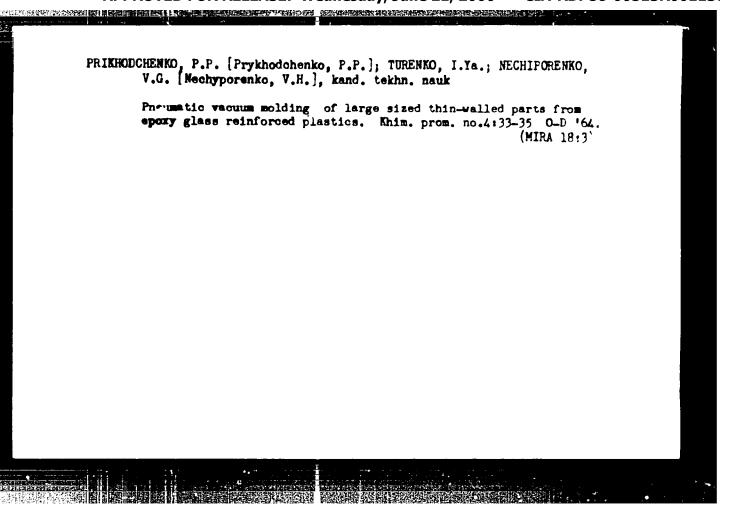
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ASSOCIATION: Kiyevskiy politekhnicheskiy institut

(Kiyev Polytechnical Institute)

SUBMITTED: May 12, 1960

Card 4/4



ACCESSION NR: AP4009590

S/0148/64/000/001/0157/0161

AUTHORS: Gridnev, V.N.; Nechiporenko, V.G.

TITLE: Electric tempering of low carbon vanadium-manganese steels

SOURCE: IVUZ. Chernaya metallurgiya, no. 1, 1964, 157-161

TOPIC TAGS: vanadium manganese steel, low carbon steel, electric tempering, tempering, coercive force, Vickers hardness, mechanical properties, rate of heating, martensite decomposition

ABSTRACT: The effects of tempering in a furnace (where desired temperature is maintained for 1 hour and sample is water cooled) and electric tempering (sample is heated at rate of 1800-2000C C./ sec. and rapidly water cooled) on the properties of the steel were compared. Vanadium (about 1.3%)-manganese (about 1.28%) steels containing different amounts of carbon were studied: allo 1, 0.054% C; alloy 2, 0.11%C; alloy 3, 0.23%C. The coercive force, Vickers hardness and mechanical properties resulting from different tempering temperatures up to 1000C were determined (fig. 1). Alloy

Card 1/82

APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001136

ACCESSION NR: AP4009590

3 was heated at rates from 80-30,000C C/soc. to determine the temperature interval of martensite decomposition (fig. 2). The study shows that in conventional furnace tempering the most favorable combination of hardness and strength is attained at about 600C. Similar or somewhat better properties are obtained with electric tempering at 700-750C than with conventional tempering at 600C. Orig. art. has: 4 figures.

ASSOCIATION: Kievskiy politekhnicheskiy institut (Kiev Polytechnical Institute)

SUBMITTED: 19May61

DATE ACQ: 14Feb64

ENCL: 03

SUB CODE: ML

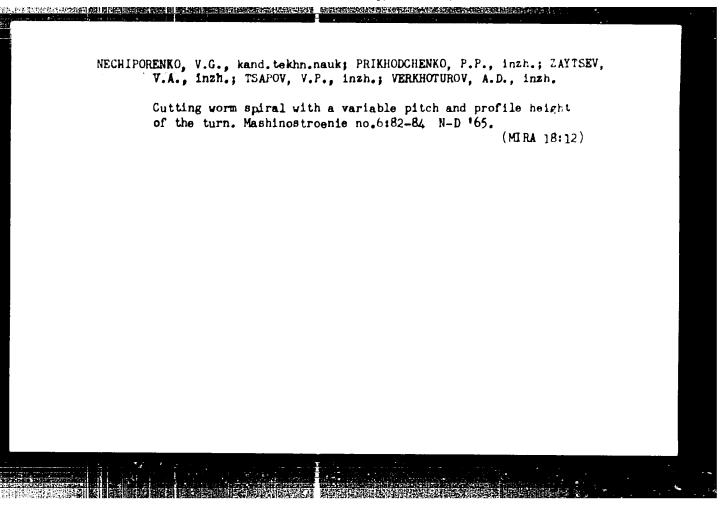
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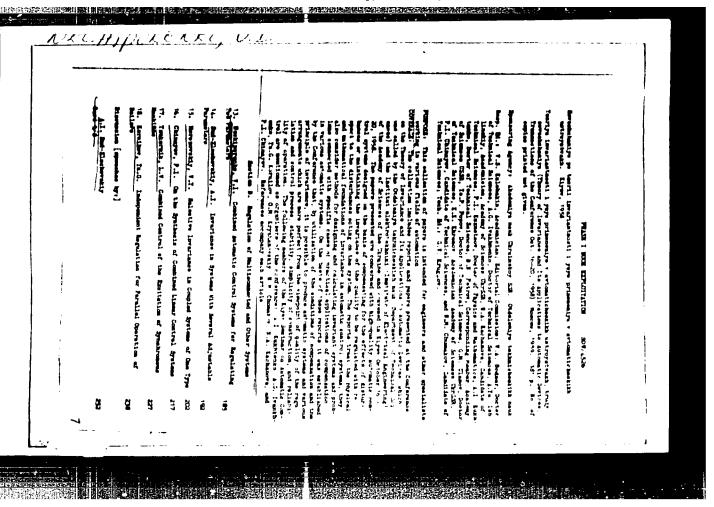
OTHER: 001

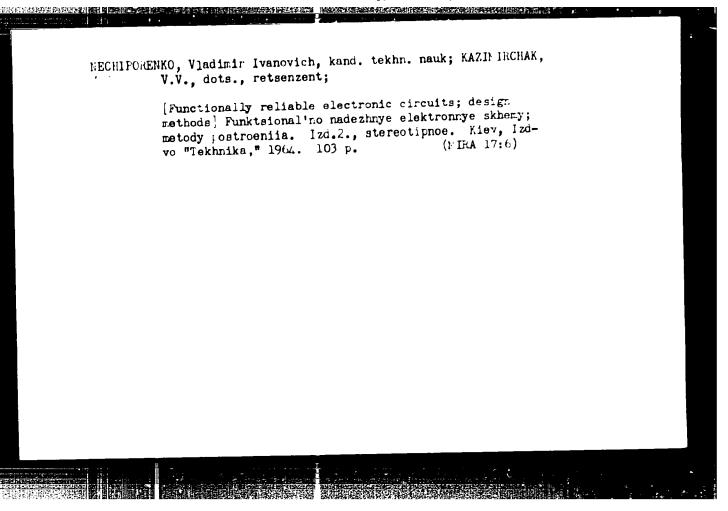
Card 2/82

APPROVED FOR RELEASE: Wednesday, June 21, 2000 CIA-RI

CIA-RDP86-00513R001136







GREBENIK, V. M.; LEONOVA, A. V.; STOROZHIK, D. A.; NECHIFORENKO, V. N.

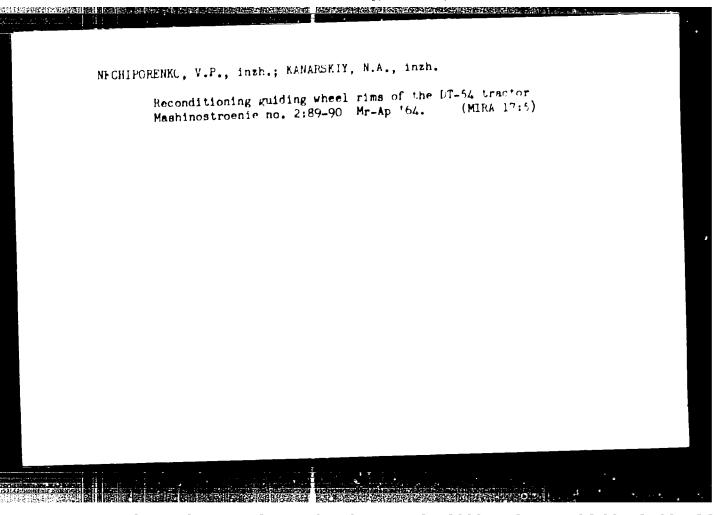
Investigating regularities of the gas flow and the wear of coupled parts in blast furnace charging arrangements. Izv.vys.ucheb.zav.; chern.met. 7 no. 4:182-185 164. (MIRA 17:5)

1. Dnepropetrovskiy metallurgicheskiy institut.

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OL'GINA, F.P., dotsent; KOSHIK, T.F.; NECHIPORENKO, V.P.

Dissecting acrtic ancuryem as a result of physical crerexertion. Vrach. delo no.ll:L21-L22 N.63 (MIRA 10:11)

1. Katedra geospital'noy terapil (zav. - prof. Ya.V.Br.m.) i patologicheskoy anatomil (zav. - prof. A.V.Sosunov) IVAI o- Frankovskogo meditsinskogo inatituta.
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APPROVED FOR RELEASE: Wednesday, June 21, 2000 CIA-RDP86-00513R001136

Cancer of the stomach accompanied with the 'rwelopment of miliary carcin matosis of the lungs in conjunction with miliary tuberculosis of the lungs. Vest. rent. i rad. 40 no.1:66-67 Ja-F '65. (Mika 18:6)

1. Hentgenovskoye otdeleniye (zav. N.Yu. Paliychuk (Irano-Frankovskoy oblastnoy klinicheskoy bol'nitsy (glavnyy vrach V.Ye. Khokhryakov) i kafedra rentgenologii i radiologii (zav. V.1. Vetoshchuk) Ivano-Frankovskogo meditsinskogo instituta.

3/137/62/000/004/049/201 A006/A101

AUTHORS:

Nechiporenko, Ye.P.; Zmiy, V.I.

TITLE:

New high-temperature heaters for electric furnaces which do not re-

quire shielding atmosphere

PERIODICAL: Referativnyy zhurnal, Metallurgiya, no. 4, 1962, 45, abstract 40300

("Poroshk. metallurgiya", 1961, no. 5, 92 - 94, English summary)

For high-temperature furnaces (1900°C), the use of Mo-rods is proposed. The rods are 6 mm in diameter, 230 mm long, with a protective MoSi2 and TEXT: refractory enamel coating. The rise of temperature in the furnace operating in air atmosphere can be brought about rapidly. The heaters were tested for 160 h at 1,750°C. A deficiency of the described heaters is the necessity of using high-power trnauformers due to the low electric resistivity of Mo.

R. Andriyevskiy

[Abstracter's note: Complete translation]

Card 1/1

CIA-RDP86-00513R001136 APPROVED FOR RELEASE: Wednesday, June 21, 2000

21224

5/126/61/011/003/012/017

11. 47. 12 5,8, 14,4 18 2,000

E021/E435

AUTHORS:

Ul'yanov, R.A., Nechiporenko, Ye.P. and Tarasov, N.D.

Vacuum Refining of Niobium TITLE:

PERIODICAL: Fizika metallov i metallovedeniye 1961 Vol.11, No.3,

pp.461-464

Results on refining experiments, the preparation of compact metal and data on the structure and mechanical properties are given. Commercially-pure miobium powder (98.7% containing 0.08% iron, 0.2% lead, 0.04% silicon and 0.18% carbon) was used. The powder also contained moisture, oxygen, nitrogen and hydrogen Hydrogen and hydrides were removed by heating in vacuo to 700°C Oxvgen and oxides were removed at 1900 to 2000 C. The powder was dried to constant weight and pressed at 5 to 6  $t/cm^2$ . Sintering was Fig.l shows carried out in vacuo at 1400°C for 4 to 6 hours. samples after this treatment. Further refining is carried out by a high temperature treatment (2300 to 2500°C) in a vacuum of 10-5 mm mercury for eight hours, in a special water cooled chamber. The samples are placed between tungsten electrodes and heated by passing a current. The appearance of the samples after treatment The purity was followed by spectrographic is shown in Fig. 2 Card 1/4

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S/126/61/011/003/012/017 E021/E435

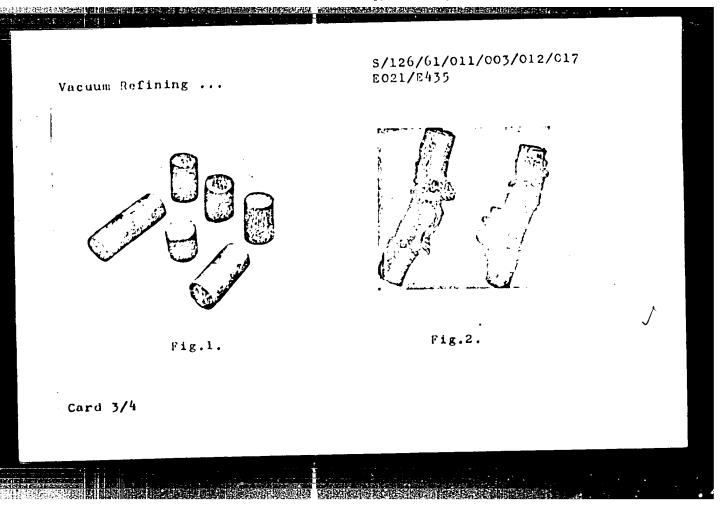
Vacuum Refining

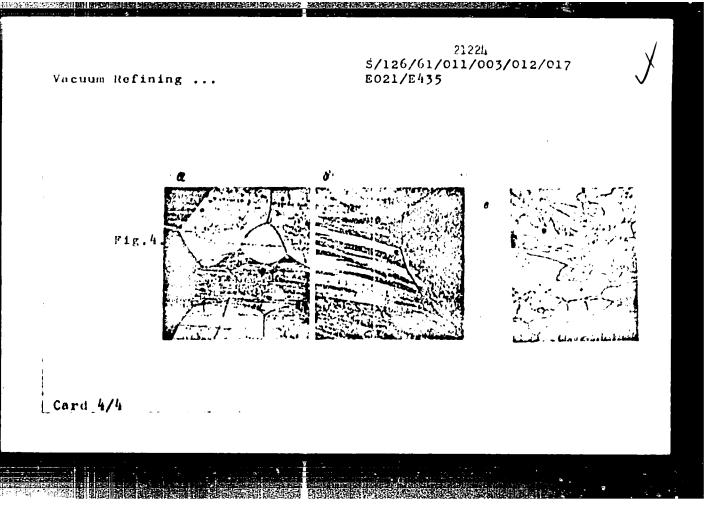
analysis, the results show how the lines corresponding to lead, silicon and iron disappear after refining. The refined metal is subjected to arc melting in an atmosphere of carefully purified argon. The ingots after melting are silver white in colour without any trace of oxidation and they have a hardness of 80 to 100 kg/mm<sup>2</sup>. The metal can be vacuum rolled at 1100 to 1200°C; the structure of the metal is shown in Fig.4 (a - as cast; b - hot rolled in vacuo at 1250°C, f - annealed at 1700°C for 10 hours). After annealing at 1700 to 1730°C in vacuo, the hardness is 80 to 90 kg/mm<sup>2</sup> (Brinell) and the tensile strength 30 to 40 kg/mm<sup>2</sup> with elongation of 30%. There are 4 figures, 1 table and 9 references: 3 Soviet and 6 non-Soviet.

ASSOCIATION: Fiziko-tekhnicheskiy institut AN UkrSSR g. Khar'kov (Physicotechnical Institute AS UkrSSR, Khar'kov)

SUBMITTED: August 2, 1960

Card 2/4





APPROVED FOR RELEASE: Wednesday, June 21, 2000 CIA-RDP86-00513R001136

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Verbinable  $n_i$  (1), ... by energy  $A_i Y_{i+1}$ , Sin(x) and  $a_i = a_i + a_i$ . 5 1 december 3 McChipprenko, Ye. C., Purachev, N. S. and Sheet, A.

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avatema.

To A0040 A : Fizika metallow a metallowed curve,  $x_1x_2^2$ ,  $x_2x_3^2$  and  $x_3x_4^2$ 

1.1 The memory of arrive means the compact molybdenna, tungsten end tyntalum her silleen en tyerman better en sere studied. Setallic samples were heated in silicon per ser on called layer, formed on the surface of the netals, a coverner by metallographic and  $N_{\tau}r_{\tau}(\mathbf{v})$  unalysis. The results shower that the saturated layer was produced, in the cain, through the vicer phase. The first stare was the fermation of lower second Afterwards, higher Silicides are formed. At 1240's, edistricted appears after 0.5, 1 and 5 hours on k, It was respectively. Once the disilicide has appeared, furt er conto occurs largely by this phase, and only after a definite fitte news has been attained is there a retardation in growth of discover e

Liffusion reac	rach states	$C(T, \zeta, z, \omega) = \sum_{i=1}^{n} \frac{1}{i} \left( \frac{1}{i} \sum_{i=1}^{n} \frac{1}{$	
snown from -1 samples during through the SI	avands a cold of the state of the cold of	and the charge of the control of the control of the charge of the control of the	
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Card 2/2			

\$/126/62/013/006/012/018 E111/E352

Glushko, P.I., Dorokhov, V.I. and Nechiporenko, Ye.P.

AUTHORS: Contribution to the kinetics of the oxidation of TITLE:

molybdenum disilicide

Fizika metallov i metallovedeniye, v. 13, no. 6, PERIODICAL: 1962, 923 - 924

TEXT: The results of a study of the kinetics of the oxidation of molybdenum disilicide in air at 900 - 1~300 C are given. Specimens were prepared by heating molybdenum plates with silicon powder at a pressure of 10 mm Hg and a temperature of 1 350 °C. After metallographic and diffraction analysis for MoSi the oxidation kinetics were studied in the interval of 900 - 1 200 °C and a duration of 6 h. The rate of oxidation per unit surface was determined from the gain in weight. The activation energy was found to be 82  $\pm$  2.5 kcal/mole and the process followed the equation:  $W = K \subset^{n} .$ 

rate constant (1.998 x  $10^{-4}$  at 900 - 2.590 x  $10^{-2}$  at 1 200 °C) Card 1/2 $\sim$  the time, K where W is the change in weight,

S/126/62/013/006/012/018 E111/E352

Contribution to ....

and n a kinetic parameter (0.72 at 900 - 0.42 at 1 200  $^{\circ}$ C). There are 3 figures and 1 table.

ASSOCIATION:

Fiziko-tekhnicheskiy institut AN UkrSSR

(Physicotechnical Institute of the AS UkrSSR)

SUBMITTED: November 28, 1961

Card 2/2

IVANOV, V. Ye.; NECHIPORENKO, Ye. P.; OSIFOV, A. D.; ZMIY, V. I.

Effect of stresses on defects in silicide layers on molytdenum.
Fiz. met. i metallowed. 14 no.4:574-577 0 '62.

(MIRA 15:10)

(Metallic films-Defects)

(Thermal stresses)

EPF(n)-2/EMP(q)/EMT(m)/BDS/T-2 AFFTC/ASD/SSD Pu-4 8/0131/63/000/007/0327/03:1 CCESSION IR: AP3004264 AUTHOR: Tvenov, V. I.; Pletenetskiy, G. Ye.; Nechiporenko, Ye. Effect of high-temperature oxide refractories on the thermal emf of tungsten, molybdenum, and tentalum in vacuum at 15000 Ogneupory+, no. 7, 1965, 527-531 TOPIC TAGE: thermocouple, high temperature, high-temperature thermocouple, insulating ceremic material, ceremic insulator, megnesia, alumina, beryllia, sirconia, tungsten, molybdomum, tentalum, tungsten wire, molybdomum wire, tentalum wire, high-temperature oxide refractory, thermal cmf, vacuum apparatus, tungstenmolybdenum thermocouple, annealing, annealed wire, vacuum furnace AMBTRACT: The stability of operation of high-temperature thermocouples made from annealed or unamealed W, No, or Ta wires after prolonged contact at 1500C with an insulating ceremic material. MgO, BcO, Al<sub>2</sub>O<sub>3</sub>, and ZrO<sub>2</sub> — has been studied in the vacuum apparatus shown in Fig. 1 of Enclosure. /, Mo, and Ts unannealed standard wires were heat-treated in contact with the pure powdered oxides for 15, 30, and 45 hr at 1500C in a vacuum (2 x 10<sup>-5</sup> mm Hg). Wires of the same netals but annealed in vacuum at 2000-2200C, were similarily treated. Temperature in Cord 1/y3

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the vacuum furnace was controlled with reference thermocomples: a VR-5/20 then couple and a platimum-platimum-rhodium thermocouple. Thermocouples were made by joining the heat-treated wire with the untreated, as a reference metal. Thermal eaf generated between the hot and cold junctions of such themsocouples was measure in the vacuum apparatus. The cold junctions of the reference thermocouple and of the thermocouples under study were maintained in vet ice. It was shown that experimental thermal cuf of the W, Mo, and Ta wires annealed and subsequently heated for 45 hr in the oxides was not significantly different from that of the unsanealed vires, except in the case of W preheated in ZrO2. Dismeter of the vires in the 0.2 to 1.0 mm range has no effect upon thermal emf stability. For each metal the changes in thermal emf due to preheating in oxides were plotted against preheating time at 1500C with each of the oxides or against temperature (in the 0-15000 range) at 45 hr of preheating. The data indicated that the thermal emf of tungsten remains stable after contact with Al,Oq, MgO, or BeO, but increases considerably with ZrO2; molybdenum thermal emf is stable after contact with Al203, MgD, or ErO2 and changes elightly after 5-hr contact with BeO; and tantalum thermal emf changes significantly after preheating in all the oxides. It was noted that small changes in the thermal cuf of W and No after contact with MgO

Card 2/N3

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ACCESSION NO: AP4009390

\$/0126/63/016/006/0931/0933

AUTHORS: Yefimenko, L. N.: Nechiporenko, Ye. P.; Pavlov, V. N.

TITLE: Oxidation of tungsten disilicide

SOURCE: Fizika metallov i metallovedeniye, v. 16, no. 6, 1963, 931-933

TOPIC TAGS: tungsten disilicide, oxidation, thermocouple, PtRh PtkRh thermocouple, oxidation curve

ABSTRACT: Oxidation of tungsten disilicide has been investigated. The process was conducted in air at a temperature range of 650-1500C. Samples 20 x 10 x 0.1 mm were produced in a vacuum of 5 x 10<sup>-5</sup> mm Hg by filling tungsten plates with powdered silicon. Nichrome elements were used to produce temperatures up to 1000C, and silicon carbide elements were used for higher temperatures. The temperatures were measured with a PtRh-PtkRh thermocouple and were kept constant. In the course of oxidation the samples were weighed with an accuracy of ±0.01 mg. Below 1000C the experiments were conducted uninterruptedly; above 1000C they were interrupted due to the formation of dense film on the surface of the plates. As can be seen from Fig. 1 of the Enclosure the rate of oxidation curves changed shape at various

Card 1/3

APPROVED FOR RELEASE: Wednesday, June 21, 2000

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ACCESSION NO: AP4009390

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temperatures. Up to 1000C the weight increase followed the formula  $W=kt^{\rm m}$ , where W is the weight change per unit area (in mg/cm²), and t is the time of oxidation (in minutes). At 1150-1250C the curves assume a descending trend because at these temperatures WO, becomes extremely volatile. A dense, glassy coating of SiO<sub>2</sub> forms at 1300C, and the process of oxidation progresses logarithmically. The formation of such a coating is described by R. Kiffer and F. Benesovsky (Symposium on Powder Metallurgy, Iron. a. Steel Inst. prep. gr., IV, 1953, 40). The logarithmic progress follows the expression  $W=k_1 \ln(k_2t+k_3)$ , where  $k_1$ ,  $k_2$ , and  $k_3$  are determined by the method described by A. Champion and T. White (J. Inst. Metals, 1949, 75, 375). Metallographic and x-ray investigation disclosed the presence of W5Siz under the glassy coating on WSi2 oxidized for a long time at high temperatures. Orig. art. has: 2 graphs, 3 formulas, and 2 tables.

ASSOCIATION: Fiziko-tekhnicheskiy institut AN UkrSSR(Institute of Physics and Technology AN UkrSSR)

SUBMITTED: 20Mar63

DATE ACQ: 03Feb64

ENCL: 01

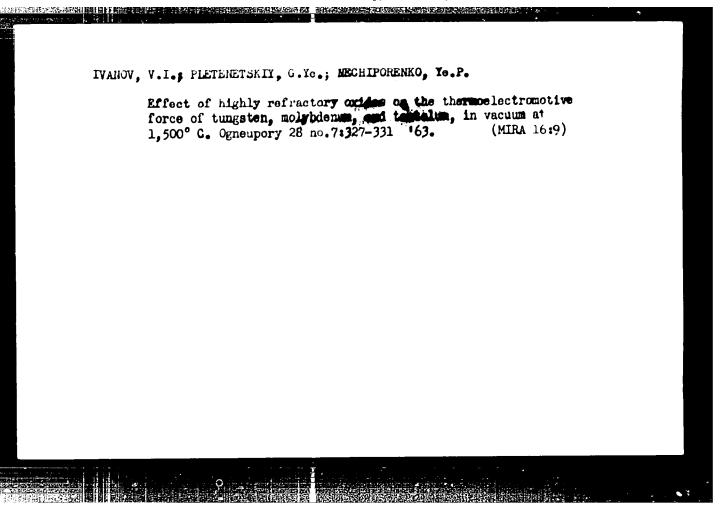
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OTHER: 003

Card 2/3

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APPROVED FOR RELEASE: Wednesday, June 21, 2000 CIA-RDP86-00513R001136

s/0126/64/017/001/0094/0099

ACCESSION MR: AP4013097

AUMICR: Ivenov, V. Ye.; Nechiporenko, Ye. P.; Zmiy, V. I.

TITIE: Study of reaction diffusion in the Mo - Si system

SCUPCE: Fizika metallov i metalloved., v. 17, no. 1, 1964, 94-99

TOPIC TAGS: metal diffusion, reaction diffusion, silicon diffusion, molybdenum silicide, molybdenum silicon system, silicide phase formation, vacuum silication

ABSTRACT: Previously published papers of the first two authors and others on various aspects of the reaction diffusion of silicon-saturated molybdenum, tungsten, and tentalum in vacuum have led to the conclusion that in the Mo - Si system the predominant role is played by diffusion of the silicon through the silicide layer; that is, the phase formation reaction takes place primarily on the internal boundary of the layer. The present article confirms this conclusion. The kinetic aspects of the vacuum silication of the molybdenum were also studied. The authors found that the growth of diffusion layers of Mossi, and Mosi, as a function of time, chays a parabolic law. From the perabolic growth of the silicide layers the authors computed the silicon diffusion factors in Mossi, and Mosi, at 12500. Used in the diffusion study were flat molybdenum samples formation millimeter in size. The

Card 1/2

ACCESSION NR: AP4013097

silicon employed in the tests was in the form of powder with a grain size of 5-7 microns (purity factor: 99.9%). The samples were located in a molybdenum bath and thoroughly sprinkled with the powder. The bath with the samples was inserted, through a precombustion chamber, into a furnace with a molybdenum heater set at the proper temperature. Orig. art. has: 6 figures, 2 formulas, and 1 table.

ASSOCIATION: Fiziko-tekhnicheskiy institut AN USSR (Physicotechnical Institute, AN UkrSSR)

SUBMITTED: 03Mar63

DATE ACQ: 26Feb64

ENCL: 00

SUBCODE: ML; PH

NO REF SOV: 009

OTHER: 000

Card 2/2

APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001136

ACCESSION IR: AP4013101

s/0126/64/017/001/01/42/01/44

**第四天中华国际的大学等于和新疆社会,并否定在这种的新疆社会的** 

AUTHOR: Ivanov, V. Ye.; Nachiporenko, Ye. P.; Zmiy, V. I.; Glushko, P. I.; Aleksandrov, O. M.; Dorokhov, V. I.

TITIE: High-temperature oxidation of molybdenum disilicide

SOURCE: Fizika metallov i metalloved., v. 17, no. 1, 1964, 142-144

TOPIC TAGS: molybdenum, silicon, molybdenum disilicide, molybdenum disilicide oxidation, molybdenum disilicide nicrohardness

ABSTRACT: Molybdenum disilicide is a metal with great promise for use in structures designed to withstand high temperatures. In the technical literature there are data on the oxidation of MoSi<sub>2</sub> achieved by various methods: hot pressing, sintering etc. The authors of this short article conducted a study of the kinetics of MoSi<sub>2</sub> oxidation in a temperature interval of 1400-17000 using a high-temperature resistance furnace. The heater was a spiral 5mm in diameter made from a molybdenum rod. For oxidation, samples of molybdenum disilicide 25x10x0.15 mm in size were used; these samples were obtained by the vacuum method. The temperature was controlled by a thermoscople (Pt - Rh 75 center: Pt-Rh 205) and an optical pyrometer, the latter placed directly on the heater. The temperature gradient between the heater

Card 1/2

APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001136

ACCESSION NR: AP4013101

and the sample was not more than 30C. A metallographic analysis of the sample was carried out with an MIM-7 microscope, with microhardness tested on a PAT-3 instrument. Oxidation time was 10 hours. It was found that with increasing time and temperature the oxidizability of MoSi<sub>2</sub> increases, the rate of oxidation obeying a parabolic law. No transition from a parabolic law of oxidation to a logarithmic one was detected in the tests. X-ray analysis in the temperature range indicated (1400-1700C) revealed an amorphous oxide film on the surface of the oxidized samples. Preliminary analysis showed that this film, in addition to SiO<sub>2</sub>, contains unknown components. These are, apparently, lower molybdic oxides, the vapor tension of which is lower than that of MoO<sub>2</sub>. The microhardness of the molybdenum disilicide, which did not change during the oxidation process, was 1200 kg/mm<sup>2</sup>. Orig. art. has: 3 figures.

ASSOCIATION: Fiziko-tekhnicheskiy institut AN USSR (Physicotechnical Institute, AN UkrSSR)

SUBMITTED: 03Mar63

DATE ACQ: 26Feb64

ENCL: 00

SUB CODE: ML

NO REF SOT: 005

OTHER: 003

Card 2/2

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IJP(c)/SSD/ EMP(e)/EMT(m)/EPF(n)-2/EPR/EMP(b)/EMP(t) Ps-4/Pu-4 L 17642-65 ASD(m)-3/ESD(t)/AEDC(b) JD/JG/AT/MH \$/0126/64/017/006/0862/0865 ACCESSION NR: AP4042042 AUTHOR: Ivanov, V. Ye.; Nechiporenko, Ye. P.; Krivoruchko, V. M.; Mitrofanov, A. S. TITLE: Some peculiarities of vacuum siliconizing refractory metals SOURCE: Pizika metallov i metallovedeniye, v. 17, no. 6, 1964, 862-865 TOPIC TAGS: Mo, W, Ta, siliconizing, vacuum siliconizing, refractory metal, silicon, vapor deposition, vacuum ABSTRACT: The authors investigate the siliconizing of W, Mo, and Ta specimens  $20 \times 20 \times 9.2$  mm in saturated silicon vapors at 1200 and 1250C and under a vacuum of 1 x 10-5 mm lig. Eliminating all contact between the Si powder and the metal, the authors observed the behavior of the vapor phase. The formation of MoSi2, W: 12, and TaSi, was identified on the surface of the specimens. The effects of time on layer thickness were plotted, and parabolas were obtained. Consequently, diffusion is a limiting factor in the process. The layer growth is defined by the equation  $h^2 = D(c_2 - c_1) \times t$ , where D is the Si diffusion coefficient in silicide, c; is Si concentration on the inner silicide layer boundary, c2 is Si concentration on the outer boundary, and t is time. W and Ta give an analogous picture. In the TaSi, phase, Ta5Si3 inclusions were found which can contribute to pinpointing Card 1/2

L 17642-65

ACCESSION NR: AP4042042

corrosion of siliconized tantalum. Siliconizing in a cell with a temperature gradient causes the rate of siliconizing to decrease as temperature of the specimen rises. The time dependence of the siliconizing at a predetermined gradient is described by a parabola. Orig. art. has: 5 figures.

ASSOCIATION: Piziko-tekhnicheskiy institut AN SSSR (Physicotechnical Institute, AN SSSR)

SUBMITTED: 15Jun63

ENCL: 00

SUB CODE: MM

NO REF SOV: 002

OTHER: 002

Card 2/2

ACCESSION NR: APHO15327

5/0032/64/030/001/0098/0099

AUTHORS: Nechiporenko, Ye. P.; Osipov, A. D.

TITLE: Apparatus for determing the modulus of elasticity of sheet materials at high temperatures

SOURCE: Zavodskaya laboratoriya, v. 30, no. 1, 1964, 98-99

TOPIC TAGS: modulus of elasticity, sheet material, high temperature apparatus, molybdenum disilicide, resonant frequency measurement

ABSTRACT: Apparatus is described for determining the modulus of elasticity of light, fragile samples by measuring the vibrational resonant frequency. The sample (in the form of a thin strip) was held at its nodal points by two metal filaments, one of which was vibrated by a solenoid. Resonance was measured by a differential capacitance device between the plates of which the sample was located. The sample and supports were enclosed by an oven. With this apparatus the modulus of elasticity at different temperatures of molybdenum disilicide was found to be  $34 \times 10^{-3} \, \text{kg/mm}^2$  at OC. It decreased linearly to  $29 \times 10^{-3} \, \text{kg/mm}^2$  at 1100C (accuracy of 5%). Orig. art. has: 1 equation and 2 diagrams.

Card 1/2

ACCESSION NR: AP4015327

ASSOCIATION: Fiziko-tekhnicheskiy institut Akademii nauk UkrSSM Institute of

Physics and Technology AN UkrGSR)

SUBMITTED: 00

DATE ACQ: 03Feb64

ENCL: 00

SUB CODE: MA

NO REP SOV: 001

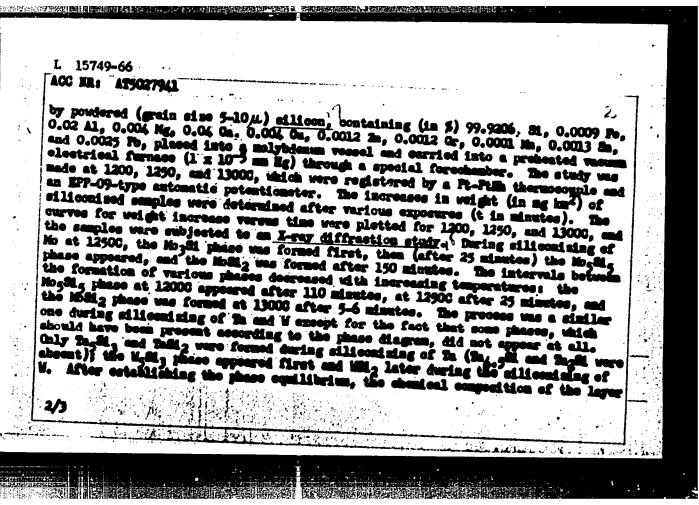
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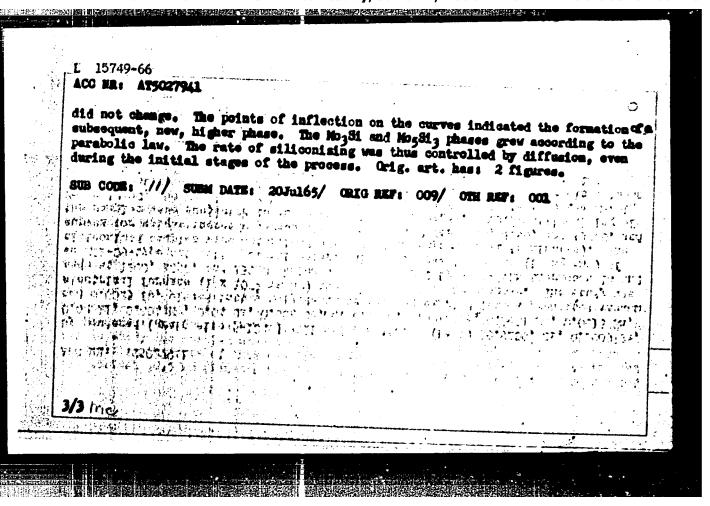
Card 2/2

APPROVED FOR RELEASE: Wednesday, June 21, 2000

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TITLE	Milloonising of ref	ractory metale	·)		<i>14</i>	
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BORISENKO, A.I., doktor tekhm. nauk, otv. red.; TOROPOV, N.A., red.; IVANOV, V.Ye., red.; APPEN, A.A., doktor khim. nauk, red.; GORBUNOV, N.S., doktor khim. nauk, red.; KLEVTSUR, S.A., doktor tekhm. nauk, red.; NECHIPORENKO, Ye.P., doktor tekhm. nauk, red.

[Heat-resistant coatings; transactions] Zharostoikie pokrytiia; trudy. Leningrad, Nauka, 1965. 233 p. (MIRA 18:9)

HATE STREET, S

- 1. Seminar po zharostoykim pokrytiyam, Leningrad, 1964.
- 2. Chlen-korrespondent AN SSSR (for Toropov, Ivanov).

L 3434-66 EMT(a)/ETC/EPF(n)-2/E	WG(m)/EWP(t)/EWP(b) _ IP(c) JD/JG/G5
ACCESSION NR: AT5024871	UR/0000/65/000/000/0045/0055
AUTHOR: Ivanov, V. Ya.; Mechiporen	ko, Ye. P.; Zniy, V. I.; Krivoruchko, V. N.
TITLE: On the vacuum siliconising	11/1/
SOURCE: AM Ukrasa. Inetitut proble	m materialovedeniya: Diffusionnyye pokrytiya
na metaliakh (Diffusion coatings for	metale). Kiev, Maukova dumka, 1965, 45-55
TOPIC TAGS: metal diffusion platin	g, silicon, refractory metal, silicide,
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ABSTRACT: The kinetics and mechani specimens measuring 40x10x1 nm vacuwith Si powder and heated at 1200-1 nation established that the formating sequence:	om of case-formation were investigated for No ium-siliconized at 1·10 <sup>-5</sup> mm Hg by being covered 350°C. Hetallographic and radiographic exami- on of molybdenum silicides occurs in the follow
ABSTRACT: The kinetics and mechani specimens measuring 40x10x1 um vacuwith Si powder and heated at 1200-1 nation established that the formating sequence:	um-siliconized at 1·10 <sup>-5</sup> mm Hg by being covered 350°C. Metallographic and radiographic exami-
ABSTRACT: The kinetics and mechani specimens measuring 40x10x1 mm vacuwith Si powder and heated at 1200-1 nation established that the formating sequence:  No + Si - Mo <sub>3</sub> Si + Si	um-siliconized at 1·10 <sup>-5</sup> mm Hg by being covered 350°C. Netallographic and radiographic examion of molybdenum silicides occurs in the follow
ABSTRACT: The kinetics and mechani specimens measuring 40x10x1 mm vacuwith Si powder and heated at 1200-1 nation established that the formating sequence:  No + Si - Mo <sub>3</sub> Si + Si	um-siliconized at 1·10 <sup>-5</sup> mm Hg by being covered 350°G. Hetallographic and radiographic examion of molybdenum silicides occurs in the follow Mo <sub>5</sub> Si <sub>3</sub> + Si MoSi <sub>2</sub>

7.50元·经验·经验公司公司的基础。 **经证明的证明的** 3 ACCESSION NR: AT5024871 lower silicides. Plotting of the curves of isothermal growth of the layers of Nossia and Nosia at 1250°C revealed that the increase in their thickness with time follows a parabolic law. This was varified by vacuum-siliconising specimens of No. Without Is in paturated Si vapore. The resulting curves also proved to follow a part rabolic law of growth in layer thickness as a function of time, thus confirming that the diffusion of \$1 is the determining factor in the rate of siliconizing. On this besis, the activation energies for the diffusion of Si in Mo<sub>5</sub>Si<sub>2</sub> and MoSi<sub>2</sub> were calculated to be  $Q_{MoSi_2} = (126,000 \pm 12,000)$  cal/mole and  $Q_{MoSi_2}$ (57,600 ± 6,000) cal/mole, respectively. Experiments to determine the effect of the presence of a temperature gradient between the box (1250°C) and the specimen (1200°C) on the growth rate of the HoSi, layer (see Fig. 1 of the Enclosure) revealed that, if the metals are siliconized in a box with a temperature gradient, the siliconising rate decreases with increase in temperature of the specimen and increases with decrease in this temperature as compared with the temperature of the box, while the growth in case-thickness follows a parabolic curve. Orig. art. has: 10 figures. ASSOCIATION:

"APPROVED FOR RELEASE: Wednesday, June 21, 2000 CIA-RDP86-00513R001136

