

PIRYATINSKII, A. L.

1. Presence of α -carene in the turpentine of the common spruce (*Picea excelsa*). T. T. Hardy, V. L. Piryatinskii, K. V. Bardysheva, and O. A. Chetayeva. Applied Chem. USSR 23, 805 (1950) (Engl. translation's Russian Ed., 817, 523; cf. J. A. 44, 10347/6). The properties and composition of two samples of spruce turpentine were determined. Turpentine distilled from spruce gum contains in the portion distilling up to 200°, 48% 1,4-pinen (1-nitrosochloride, m. 102.3°), 17% 1,8-pinen (converted to naphthalic acid, m. 120°), 1% d- Δ^4 -carene (nitrosoate, m. 147°), 18% of a mixture of dipentene (tetrabromide, m. 125.6°), and limonene, and higher-boiling constituents. Turpentine obtained from relatively fresh spruce gum contains I-10, 1,4-pinen 35, d- Δ^4 -carene (m.p. 170-170.7°), nitrosoate m. 147°) 10% of a mixture of dipentene and 1-limonene, and higher-boiling constituents. The optical activity of I in the first sample was much lower than that of I in the second, which is a relatively fresh sample.
Richard I. Akawle

26

2. Dependence of viscosity of linseed oil on its oxidation. II. V. G. Georgievskii and B. N. Shakhnel'dyan (Moscow)

PIRYATINSKIY, A.L.

BUGLAY, B.M., kandidat tekhnicheskikh nauk; PIRYATINSKIY, A.L., kandidat
tekhnicheskikh nauk; KORSHUN, L.L., inzhener.

Terpene-collodion lacquers for finishing furniture. Der.i lesokhim.
(MLRA 7:2)
prom. 3 no.1:3-5 Ja '54.

1. TSentral'nyy nauchno-issledovatel'skiy institut mekhanicheskoy
obrabotki drevesiny (for Buglay). 2. TeNIIKhI (for Piryatinskiy and
Korshun).
(Lacquer and lacquering)

PIHYATSIKY, A. L.; BUGLAY, B. M.; KORSHUN, L. L.

New polishing and softening agents for the refining of nitro lacquer coatings. Sbor. trud. TSMILKHI no.13:115-118 '59.
(MIRA 13:10)
(Lacquer and lacquering)

BUGLAY, B.M., doktor tekhn.nauk; PIRYATINSKIY, A.L., kand.khim.nauk; SHUBINA,
I.I., inzh.; KORSHUN, L.L., Inzh.

New materials used for finishing furniture. Der.prom. 7 no.9:1-5
(MIRA 11:11)
S '58.
(Wood finishing)

KORSHUN, L.L.; TRIMONOVA, T.V.; PIRYATINSKIY, A.L.; BUGLAY, R.M.; SHUBINA, I.I.

Fungicidal nitro varnishes based on oxyterpene resins. Der.prom.
7 no.11:1-2 N '58. (MIRA 11:11)
(Varnish and varnishing) (Fungicides)

KALANTAROV, Pavel Lazarevich; TSEYTLIN, Lev Aleksandrovich; PIRYATINSKIY,
A.Z., redaktor; ZABRODINA, A.A., tekhnicheskij redaktor

[Inductance calculations; reference book] Raschet induktivnosti:
spravochnaja kniga. Moskva, Gos. energ. izd-vo 1955. 367 p. (MLRA 8:3)
(Inductance)

1. PIRYATINSKIY, A. Z.
 2. USSR (60)
 4. Ionization
 7. Electric puncture of technical dielectrics.
Zhur. tekh. fiz., 22 No. 10, 1952.
-
9. Monthly List of Russian Accessions, Library of Congress, February 1953. Unclassified.

1. VYATINOV, N. A.
2. NSCR (60)
4. Dielectrics
7. Electric nature of technical materials. Indus. Techn. Phys., 1958, No. 1.
9. Monthly List of Russian Acquisitions, Library of Congress, February 1958. "In the USSR".

PA 236T17

PIRYATINSKIY, A. Z.

USSR/Electricity - Dielectrics, Breakdown Oct 52

"Problem of Electric Breakdown of Technical Dielectrics," A. Z. Piryatinskiy

"Zhur Tekh Fiz" Vol 22, No 10, pp 1556-1564

Discusses mechanism of breakdown of ceramics at high frequencies. Assumption of "thermitization" character of breakdown of porous dielectric makes it possible to establish a connection between physicomechanical properties and electric strength. Indebted to N. P. Bogoroditskiy. Received 17 Jan 1942 [sic].

236T17

PIRYATINSKIY, B.O.

Two species of Upper Jurassic Trigoniidae from western Turkmenia.
Vest. LGU no. 24:146-149 '62. (MIRA 16:2)
(Turkmenistan—Lamellibranchiata, Fossil)

"APPROVED FOR RELEASE: 07/13/2001

CIA-RDP86-00513R001341020006-4

OPERATION KITE, B-6.

Components of the system developed by the Central and Western
Armaments' Vertical Joint Venture of the USSR - VZKFA

APPROVED FOR RELEASE: 07/13/2001

CIA-RDP86-00513R001341020006-4"

PROZOROVSKAYA, Ye.L.; PIRYATINSKIY, B.G.

Some characteristics of upper Callovian sediments in the Tuar-Kyr
region. Trudy VSEGEI 46:101-105 '61. (MIR 14:11)
(Tuar Kyr region--Geology, Stratigraphic)

AMANNIYAZOV, K.; PROZOROVSKAYA, Ye.L.; PIRYATINSKIY, B.G.

Upper Jurassic sediments in the Kyzylkyr boundary (Tuar-kyr region).
Trudy VSEGEI 46:106-107 '61. (MIRA 14:11)
(Tuar-kyr region--Geology, Stratigraphic)

The Production of Vinyl Phenols by the Catalytic
Cracking of Some Dioxydiarylalkanes

S-41-

S/020/60/132/G2, 72
B011/B002

the catalysts almost always three fractions developed: I phenol; II p-phenol admixtures of ethyl phenol and p-vinyl phenol; III p-vinyl phenol with slight high concentration of p-vinyl phenol. Under the condition of selective cracking and of a III in the form of paleish green lamina. The yield in fraction III and the conversion of dioxydiphenylethane into light products increased with a higher volume velocity of the dioxydiphenylethane solution. The authors describe some of the most successful experiments. After several processes of recrystallization obtained Crude crystals dissolved easily in benzene, alcohol, and ether, and not so well in water. After left standing in the vacuum desiccator, for a short time the solubility was reduced due to polymerization. The crystals dissolved in lye turned the solution brown. An admixture of p-vinyl phenol to concentrated H₂SO₄ gave it a vividly red color. An admixture of a ferric chloride solution to the aqueous solution of p-vinyl phenol gave it a brownish green color. In the dark, p-vinyl phenol rapidly polymerizes into an insoluble white resin. In a protective gas however, it keeps up to 50 hours and more. Cracking dioxydimethylethane (ethylidene-di-o-cresol) in acetone and benzene the following substances were obtained: o-cresol, 4-ethyl-o-cresol and o-vinyl

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The Production of Vinyl Phenols by the Catalytic
Cracking of Some Dioxydiarylalkanes

S/C2C/6C, 112/
B011/B002

cresol. The latter is a white, crystalline substance with a melting point of 74°. It is soluble in ordinary solvents, and under the action of air it forms into a sticky resin from which after treatment with benzene the p-isopropenyl derivative of 4-vinyl-o-cresol precipitates in the form of an indissoluble white powder. Dicyanodiphenylpropane (diphenylolpropane) was obtained from a commercial product supplied by GIPI-4 (Gosudarstvennyy nauchno-issledovatel'skiy i proyektnyy institut-4, State Design and Planning Scientific Research Institute) by distillation and recrystallization. Cracking was the same as above, but was conducted in acetone-benzene. White, scale-like crystals of p-isopropenyl phenol with a melting point of 80.5° was obtained from the catalyst. Exposed to air they transformed into a red resin difficultly soluble in organic solvents. There are 1 figure and 7 references, 2 of which are Soviet.

ASSOCIATION: Institut neftekhimicheskoy i gазовой промышленности им. И. М. Губкина (Institute of Petroleum-chemical and Gas Industry, I. M. Gutkin)

PRESENTED: November 5, 1954, by A. V. Topchiyev, Academician

SUBMITTED: November 5, 1959
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S/080/61/074 06/10/87
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AUTHORS: Vayser, V.L., Ryabov, V.D., and, Piryatinskiy, P.V.

TITLE: The condensation of acetylene and phenol in the presence of cation exchange resin KU-2

PERIODICAL: Zhurnal prikladnoy khimii, v. 34, no. 6, 1961,
1381 - 1381

TEXT: The aim was to discover more effective methods of synthesizing 4,4'-dioxydiphenylethane (diphenol) using catalysts containing a mercury salt. Cationite KU-2 was chosen. Diphenol which is of great use in the synthesis of high molecular compounds is formed from the condensation of acetylene and phenol in aqueous and alkoholic solution in the presence of various acidic catalysts and mercuric oxide. The best catalyst was $H_3Pb_4 \cdot BF_4$. Commercial cationite was treated with hydrochloric acid, washed with water, treated with an alkoholic solution of mercury salt, and dried. 1-2 % by weight of mercury salt was adsorbed on the surface of the

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catalyst. The experiments were carried out in a three-neck flask provided with a stirring rod, a reflux condenser, mercury seal and glass funnel for the addition of acetylene. Calcium carbide, phenol (30 g.) were placed in the flask and, at a temperature of 170° , acetylene was run in for 4 hours at the rate of 5 liters an hour. When the reaction was over the flask contents were vacuum-filtered to separate the catalyst, the latter washed with a small quantity of phenol and used again. The reaction products were distilled under pressure, the fraction of 4.4'dioxydiphenylethane collected at $210-220^{\circ}$ and 8 mm Hg. A series of tests was done to study the variation in catalyst activity with time. Acetylene and phenol were condensed also in aqueous solution at 90° , other conditions remaining constant. 4.4'dioxydiphenylethane was obtained and it was shown that in this case acetaldehyde was formed at an intermediate stage. The advantages of KM-2, activated by mercury salts, as a catalyst in this reaction, are as follows: It avoids neutralization of the reaction product, it is active for a long time and easily separable, though the yield of diphenol is considerably lower than

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The condensation of ...

when using $H_3PO_4 \cdot BF_3$. Conclusions: Acetylene condenses with phenol in the presence of attionite KU-2, activated by mercury salts, at $50-150^\circ$ forming 4,4'-oxydiphenylethane (yield at 4%); the yield is at a time of more than 4 hours, its activity rising to constant level; in the presence of water, acetaldehyde is an intermediate product. There are 2 figure and 3 Soviet-bis references.

SUBMITTED: April 16, 1960

Card

PIKYATINSKIY, I. L.

27152. PIKYATINSKIY, I. L. SHVARTSMAN, I. SH. Proizvodstvo i sluzhba stalerazivochnykh probok na zavodakh urala i vostoka. Ogneurny, 1949 No. 8, s. 340-45.

So: Letopis' Zhurnal'nykh Statey, Vol. 36, 1949

MIRYATINSKII, I. L.

C
(S) 5

Manufacture and service of steel-pouring plugs in the Urals and
OSS EAST XI. L. MIRYATINSKII AND I. L. MIRYATINSKII. Orenburg, 14 (21-340 IA (1648). Details are given on the production
of plugs in various refractory and metallurgical works in the
Urals and in the eastern part of the Soviet Union and on their
service in (V. A. 13) and (V. A. 14). Results show that the
spherical portion of the plug is subject to greatest wear and defor-
mation. W.Z.K.

"APPROVED FOR RELEASE: 07/13/2001

CIA-RDP86-00513R001341020006-4

1. APPROVAL:

2. APPROVAL DATE:

3. APPROVAL SIGNATURE:

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CIA-RDP86-00513R001341020006-4"

"L.YATINSKIY, I. L. and R.

"The type of break in "new articles"

Onnepory, No. 5, 1975

PIRY/TINCKIY, I. L. M.R.

"Production and service of steel structures
in the plants of the oil and gas industry"

Obninskoy, No. 1, 1970

PLATINUM, I. L. Eng.

"Productive service to our country, being it a silent one until now".

Ottoman, N.Y., 1944

*Reed Abo**57-69-Subas 24-1*

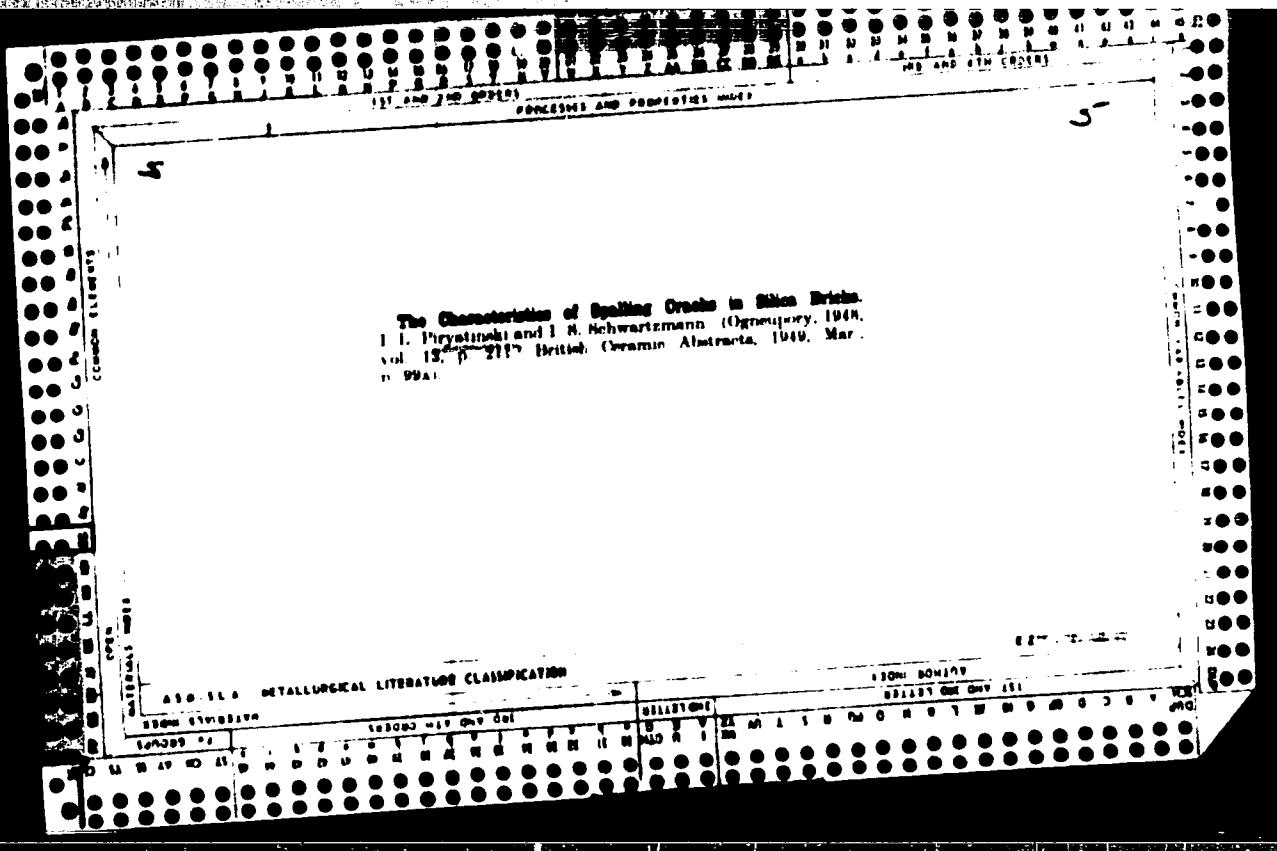
Characteristics of spalling cracks in silica bricks. I. L. Pyrzakowski and J. S. Schurman (Ogonyk, 1966, 23, 311; Brd. techn. Akad., 1966, 5(2)).—Straight spalling cracks do not always appear on the surface and may penetrate into the body of the brick; cracks in the form of a network penetrate less deeply (5–10 mm.). The depth of a crack increases with its width, but depth and width are not directly related. An increase in width from 0.1 to 0.8 mm. corresponds with an increase of 50% in the ap. area of the spalled fragments; a proportionately greater increase in area of the latter results from an increase in length of crack. Comparison of cracks in various types of refractories showed that the greatest no. of straight spalling cracks occurs in SiO_2 bricks for electric and open-hearth furnaces and the smallest no. in chrome-magnesite bricks. SiO_2 bricks for electric furnaces have the highest proportion of cracks in network form; such cracks are almost absent in magnesite bricks. Cracks in SiO_2 and fireclay were approximately of the same length but much longer than those in magnesite and chrome-magnesite ware. The no. and width of cracks of the straight and network types in SiO_2 brick increase as the d of the brick decreases. The method of determining the depth of flaws and spalling cracks by cutting a cross-section through a brick is applicable to SiO_2 bricks and in some cases to magnesite, but not to fireclay and chrome-magnesite products.

Brd. techn. Akad. (USSR)

5441. CHARACTERISTICS OF SPALLING CRACKS IN SILICA BRICKS.
Piryatinski, I. L. and Schwartmann, I. S. (Ogneupory, 1948, vol.
13, 211).

"APPROVED FOR RELEASE: 07/13/2001

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The composition of turpentine from the exudations of the
Siberian cedar *Pinus cembra*. I. I. Bardyshov, A. V. Prok-
torskii, K. V. Bardyshova, and O. Chernaveva. *Voprosy
lesovedeniya i selskogo hospodarstva SSSR*, No. 23, 1963. (English translation: See
C.I. 44, 10474) H.M.S.

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Composition of native turpentines from the common pine. I. I. Bardynshev, A. L. Piryatinskii, K. V. Bardynsheva, and O. I. Chernyavova. *Zhur. Prilobl. Khim.* (J. Applied Chem.) 23, 203-10 (1980).—The terpene portion of com. samples of turpentines: retort, furnace, extra, and sulfate types, was qualitatively analogous to the oleoresin turpentine and contained: α -pinene, β -pinene, d-terpine, δ -carenne, limonene, dipentene, and terpinolene. The results secured by extensive fractionations showed only minor variations of the ants. of components, principally as follows: limonene-dipentene mixt. 13.17% in retort turpentine, 10% in furnace variety, 7% in extra variety, and 3% in sulfate variety. The alc. content, calcd. as $C_{10}H_{16}OH$, was: 6.7, 30, 5.5, and 3.7%, resp.
O. M. Konchalovskii

1 The composition of active turpenines from the common
pine L. I. Radyshov, A. N. Sviridov, K. V. Rady-
shov, and O. I. Chernyavskaya. *Applied Chem. USSR*
23, 209-16 (1950) (Engl. translation). See C 4 04 103474
B. L. M.

6 A

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Properties and composition of turpentine from the resin
of *Picea abies* L. L. Hardyshew, A. I. Pyatnitskii,
and K. V. Bardysheva. Dobledy Akad. Nauk SSSR
3, 75 7(1950); cf. C.A. 45, 1781g. -Resin of 100 speci-
mens of the turpentine showed variations as follows:
 α_{D}^{20} = +31.40' to +31.46'; α_{D}^{20} 0.0500 to 0.0511; MW 1, 17.22.
All contained α -pinene (+31.47%) while β -pinene varied
from 0 to 6%; δ -terpine from 1 to 11%; γ -terpine from 0 to 5;
31%. High α -pinene went along with β -pinene and low
 δ -terpine.

CA

Composition of a representative commercial sample of dry-distillation furnace turpentine I. I. Bardyshov,
A. L. Piryatinskii, K. V. Bardyshova, and O. I. Cherny.
6666 "Zhur. Khim. Khim. (J. Applied Chem.) 23
552-6(1980).—Fractional dist. and Raman analysis of
the fractions of a sample representatively taken from com-
mercial turpentine gave the following compn.: 50%
d- α -pinene, 7.0% l- β -pinene, 6.6% d-terpine, 16.4%
d- β -cavene, 9.7% mixed dipinenes and l-isopinene, 1.6%
terpenodiene and 6.6% high-boiling substances, principally
terpenoals. The top fraction consists largely of products
of dry dist. of wood; no α -pyrone was found in it.
The turpentine is qualitatively identical with turpentines
made by other procedures from the same starting ma-
terial (common pine) G. M. Kosolapoff

"APPROVED FOR RELEASE: 07/13/2001

CIA-RDP86-00513R001341020006-4

A rapid analytical method for turpentine N. V.
Tukhovitskii, A. I. Bryatinskai and G. D. Atamanchuk
kor. Zembla. Izvem. S. No. 2, 68 (1960). Description
of a combined extr. distn. lab. app. A. A. P.

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CIA-RDP86-00513R001341020006-4"

Increase of α -pinene in turpentine oils from collation
I. I. Barabashev, A. I. Pugatsinskii, A. V. Karts-
sheva and O. I. Chernovskaya, *Zhur. Praktich. Khim.* 11,
Applied Chem., 1, 20, 1338, 1947 - Fractionation
of 6 kg. samples of turpentine from the Barnaul tur-
pentine plant showed that a typical material contain-
 α -pinene 62.6%, β -pinene 8.3%, δ -3-carene 21.8%
 α -limonene and dipentene 3.73, δ -caryophyllene
2.71, and higher boiling components 2.7%. The
pinene isolated is not uniform in properties. Typical
starting material had n_D^{20} 1.4991, d_4^{20} 0.8894, $\eta_{D,4}^{20}$ 14.1
 $\lambda_{\text{max}}^{\text{vis}} 300$. The distill curves and tabulations of frac-
tions are given. The properties of the pure substances
obtained were: α -pinene, n_D^{20} 1.5455, d_4^{20} 0.8957
 d_4^{20} 0.8984, $\eta_{D,4}^{20}$ 14.81, $\lambda_{\text{max}}^{\text{vis}} 29.0$; β -pinene, n_D^{20} 1.5154, d_4^{20}
 d_4^{20} 0.8984, $\eta_{D,4}^{20}$ 14.71; β -pinene, n_D^{20} 1.5154, d_4^{20}
 d_4^{20} 0.8984, $\eta_{D,4}^{20}$ 14.71; δ -3-carene, n_D^{20} 1.5154
 d_4^{20} 0.8984, $\eta_{D,4}^{20}$ 14.71; α -limonene, n_D^{20} 1.5154
 d_4^{20} 0.8984, $\eta_{D,4}^{20}$ 14.71; δ -caryophyllene, n_D^{20} 1.5154
 d_4^{20} 0.8984, $\eta_{D,4}^{20}$ 14.71. The δ -caryophyllene component
is not uniform. The first caryophyllene sample had
 n_D^{20} 1.5154, d_4^{20} 0.8984, $\eta_{D,4}^{20}$ 14.722, d_4^{20}
 d_4^{20} 0.8984, $\eta_{D,4}^{20}$ 14.71, $\lambda_{\text{max}}^{\text{vis}} 29.0$ (contrast to
 n_D^{20} 1.5154, d_4^{20} 0.8984, $\eta_{D,4}^{20}$ 14.71, $\lambda_{\text{max}}^{\text{vis}} 29.0$).
The δ -caryophyllene component (see above), n_D^{20} 1.5154
 d_4^{20} 0.8984, $\eta_{D,4}^{20}$ 14.71, $\lambda_{\text{max}}^{\text{vis}} 29.0$, $\lambda_{\text{max}}^{\text{vis}} 30.2$.
Its nature is not known. The compn. of the sample
after fractionation was determined by the dispersion method of
Dorsoot. It is believed that the failure of the earlier
Russian workers to isolate δ -pinene was caused by the use
of insufficiently effective fractionating apparatus.
- (A. Kondapoff)

Resinous substances from spruce waste. A. I. Pyryev,
Lunskil and N. V. Tukhovitskai. Russ. 42,281, March 31,
1965. The resinous substances are extracted by wood-tar
from products obtained from deciduous trees.

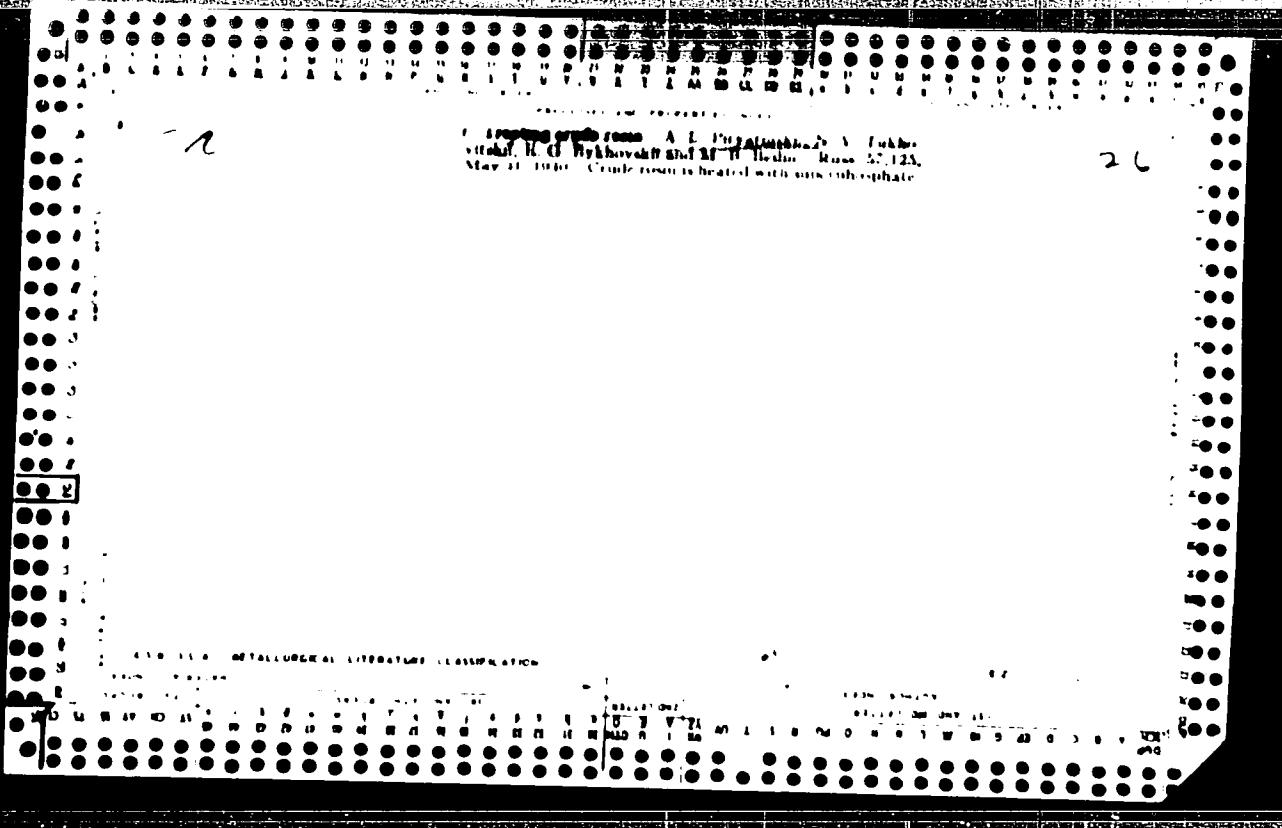
(H)

Presence of α -carenne in the turpentine of the common spruce, *Picea sylvatica*. A. L. Kwolek
Soviet. V. Rastsvetova, et al., U.S. Patent 3,487,321, filed 1964, English translation Received 1967-02-01, C.I. 144.00x47/A. The properties and composition of two samples of spruce turpentine were determined. Turpentine distilled from spruce gum contains, in the portion distilling up to 200°, 40% β -pinene (Γ -nitroacetylchloride m.p. 11°, 17% β -pinene converted to pinonic acid, m.p. 52°), 40% α -pinene (nitrosoate m.p. 147°, 18% of a mixt. of dipentene, tetrabromide m.p. 125.6°, and limonene, and higher-boiling constituents). Turpentine obtained from relatively fresh spruce galley contains 14% β -pinene, 33% α -pinene, 16% 170-170.7° nitrosoate m.p. 147°, 10% of a mixt. of dipentene and 7 limonene, and higher-boiling constituents. The optical activity of Γ in the first sample was much lower than that of Γ in the second, which is a relatively fresh sample.

Richard F. Akowic

Characteristics of spruce resin. A. I. Bykovskii
Zavod chern. No 2, 17-19 (1957). Resin contains
no less unsaturated substances 11.4; substances in-
soluble in ether 40.1%; softening point 70.0; ash
content 0.1%; melting point 126.0°C. It is dark red.
Source references: A. A. Podgornyy

Varnish resins. A. I. Pavlyuchenko, L. I. Vaynshteyn
and V. V. Tikhonov. U.S.S.R. 897816. Oct. 1967.
The resins are obtained by esterification of adipophenol
with phenyl phenyl ether. In the first step, adipophenol is
reacted with phenyl phenyl ether in a quantity of 1.5-
less than is required for complete esterification. In the
next step, the esterification is finished with benzyl alcohol taken
in a quantity equal to the surplus pentavinyloxitol.
Resins thus obtained have a higher m.p. and are more
moisture-resistant than the resins obtained when only
phenyl phenyl ether is used.



—Treating crude resin. A. L. Duganashvili, N. V. Tukhovskii and E. G. Bykbovskii. Russ. 57,125. May 11, 1955. Adds to Russ. 57,125 (preceding abstract). The method of Russ. 57,125 is modified in that a solid salt of copper sulphate is used.

Application of the Dermois method to analysis of native turpentine. I. Constancy of Biot's law. J. J. Wardy
Revista de la Universidad Católica de Chile, Vol. 11, No. 1
April-June, 21-12-1918. In view of the results
obtained by Dr. D. L. B. Dermois, showing the constancy of Biot's law
in the case of native turpentine, it was decided to
make a similar investigation. As the results of Dr. Dermois
are given in figures and tables, the differences observed in our
work, which were the result of the different methods used,
will be very apparent. On the other hand, the
method used here is the same as that of Dr. Dermois.
Hence, the results of the two methods can be
compared with great interest. The results show that the
physical constants of native turpentine, which are
given in the tables, are correct.

Crude turpentine obtained from stamps A 1, Pyatigorsk and B. A. Seregin *Zembla* from Z, No. 2, 47 (1937). The crude turpentine is characterized by the presence of heavy ends (pine oil), composed mainly of terpene ales, which are used as flotation agents, solvents and disinfectants. It has the following characteristics: sp. gr. 0.8732, Br no. 172, vis. 1.4730, b.p. 153-160° residue 180°, 190°, vis. 0.915°C (read as $\text{C}_6\text{H}_5(\text{O})_2$, esters 0.44% (read as $\text{C}_6\text{H}_5(\text{OCOC}_6\text{H}_5)_2$). In a steam distill. in the lab., 16 fractions were used. The distillates can be classified into 2 groups. The first group is characterized by its lower sp. gr. (0.8817-0.8922), low ale content and high content of terpene hydrocarbons. The 2nd group is a straw-yellow liquid of a higher sp. gr. (0.8912-0.8967) and has a high content of ales (up to 40%) and a considerably higher n. The first 6 fractions constitute a high grade turpentine with a high content of phenic fraction. The 7-11 fractions can be classified as a 2nd grade turpentine which can be used as a solvent in the varnish industry. The intermediate (12th) fraction can be recycled together with a new charge, while the higher fractions (about 13.8%) can be used as the stock in the prepn. of high-grade pine oils. The residue about 4.5% is a viscous, heavy substance that may find application in the rubber industry. A A B

ABSTRACTS OF METALLURGICAL LITERATURE CLASSIFICATION

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(A)

22

Crude turpentine obtained from stumps. A. I. Pecherskaya and V. A. Sogolov. *Vestn. Akad. Nauk SSSR*, No. 4, p. 43 (1954). The crude turpentine is characterized by the presence of heavy endo (pinole oil), composed mainly of terpenes, which are used as flotation agents, solvents and distillates. It has the following characteristics: sp. gr. 0.842; Boiling point 120°C.; density at 18°C., residue 0.41%; calculated as 30.41% C. The stock distillate in the latter 10 fractions were separated. The distillates can be classified into 2 groups. The first group is characterized by the lower sp. gr. (0.821-0.822), low olefin content and high content of terpine hydrocarbons. The 2nd group is a straw yellow liquid of a higher sp. gr. (0.822-0.826) and has a high content of aldehydes (6-10%) and a considerably higher %... The first 10 fractions constitute high grade turpentine with a high content of pinene fraction. The 11-14 fractions can be classified as a 2nd grade turpentine which can be used as a solvent or the varnish industry. The intermediate 12th fraction can be recycled together with a new charge while the higher fractions (about 14.5%) can be used as the stock in the prep of high grade pine oils. The residue (about 4.5%) is a viscous, heavy substance that may find application in the rubber industry. A. A. B.

CA

Creosote from tar obtained as residue in the rectification of "black" acetic acid in the Vilensov experimental extraction plant. A. I. Pavlinskii and I. P. Savenkov *Izvestia Akademii Nauk SSSR*, Chem. No. 3, 4, 21, 31 (1932). A dil. AcOH solution (from 1, No. 3, 4, 21, 31) was steam distilled and the distillate contg. AcOH as well as resins and higher boiling fractions was investigated. The fraction b 110-200° obtained from the residue by distn. (64% of the residue) had a sp. gr. of 1.07 and acidity of 14.70% (calcd. at Ac(III)). They were used in the prepn. of creosote. This distillate was redistilled into 2 fractions, the first fraction having an acidity of 17.80%, and a b.p. of 180-180° while the second fraction had a b.p. of 180-240°. The latter was treated with a 10% soln. of calcined soda for the removal of free acids and the oil obtained was treated with a 8-10% soln. of caustic soda (an excess is used). The phenolates formed from a small layer of oil were heated with open steam for the removal of substances which did not combine with the alkali. The caustic soln. of phenolates was acidified with a 30% H₂SO₄ to a clear acid reaction and the floating oil (phenols) was sep'd from the H₂O layer contg. water and phenols (which were wld. by salting out and were added to the basic portion). The phenols were neutralized with a 5% calcined soda soln. and washed with water. The pure phenols were distd. into 2 fractions and a residue boiling above 240°. The yield of the fraction b 180-240° amounted to 9.60% of the resins and produced a clear

soln. when 1 cc. was mixed with 2.5 cc. of a 15% NaOH soln. which could be diluted to 50cc. before the appearance of turbidity. The heavy fraction b 200-230°, conforming with the requirements of the U. S. S. R. Pharm. and obtained in the distn. of the purified phenols, was treated with a 10% soln. of NaOH (with slight excess) and blown with steam till transparent and treated in the same manner as the low boiling fraction. The liquid was then distd. into 4 fractions and the combined first 3 fractions b 98-220° were treated with chromic acid mixt., left stand overnight and redistd. The second fraction b 200-230°, which amounted to 3.80% of the resins, was an oily substance with a high s. a slightly yellow color and a sp. gr. of 1.090, yielding 96.3% with a b.p. of 200-205° on redistn. by Engle's Distilometer. It passed all tests prescribed by the U. S. S. R. Pharm. except that in the presence of cold bar phenols.

22

AIA 520 METALLURGICAL LITERATURE CLASSIFICATION

CP

Crocosm from tar obtained as residue in the rectification of "black" acetic acid in the Vilenov experimental extraction plant. A. I. Bryantsev and L. F. Samokov. *Zhurnal Prom. 1, No. 3-4, 21, 6 (1932).* A dil. AcOH and cat. Nasakim, C. I. 23, 4077 was steam distilled and the distillate contg. AcOH as well as resins and higher boiling fractions was investigated. The fractions b. 110-200° obtained from the residue by distn. 54% of the residue) had a sp. gr. of 1.07 and acidity of 14.70% (calcd. as AcOH). They were used in the prep. of esterate. This distillate was redistilled into 2 fractions, the first fraction having an acidity of 17.81%, and a b.p. of 100-110° while the second fraction had a b.p. of 180-240°. The latter was treated with a 10% soln. of calcined soda for the removal of free acids and the oil obtained was treated with a 8-10% soln. of caustic soda (an excess is used). The phenolates septd. from a small layer of oil were heated with open steam for the removal of substances which did not combine with the alkali. The cooled soln. of phenolates was acidified with a 30% H₂SO₄ to a clear acid reaction and the floating oil (phenols) was septd. from the H₂O layer contg. water-sol. phenols which were septd. by salting out and were added to the basic portion. The phenols were neutralized with a

72

5% calmed. soda soln. and washed with water. The pure phenols were distd. into 2 fractions and a residue boiling above 240°. The yield of the fraction b. 180-240° amounted to 9.36% of the resins and produced a clear soln. when 1cc. was mixed with 2.5cc. of a 15% NaOH soln. which could be diluted to 50cc. before the appearance of turbidity. The heavy fraction b. 200-220°, conforming with the requirements of the U. S. S. R. Pharm. and obtained in the distn. of the purified phenols, was treated with a 10% soln. of NaOH (with slight excess) and blown with steam till transparent and treated in the same manner as the low boiling fraction. The liquid was then distd. into 3 fractions and the combined first 2 fractions b. 108-220° were treated with chrome and mixt., left stand overnight and redistd. The second fraction b. 100-200°, which amounted to 4.86% of the resins, was a oily substance with a high w. a slightly yellow color, a sp.gr. of 1.090 yielding 54% with a b.p. of 200-240°. It passed all the tests prescribed by the U. S. S. R. Pharm. except that for the presence of coal-tar phenols. A. A. R.

CA

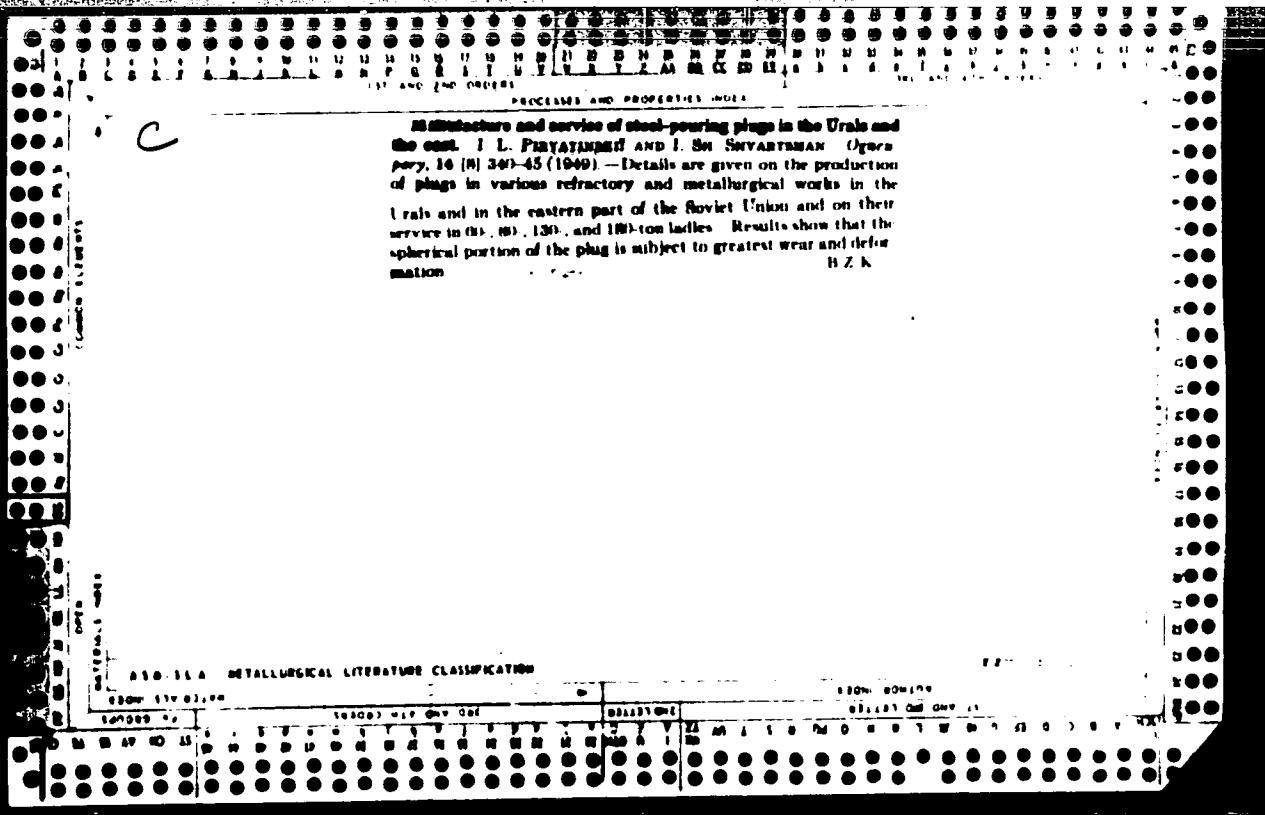
26

Some methods for improvement of rosin and for conversion to other products A. L. Finsen, Jr., U.S. Pat. No. 2,139,518, 1939, N.Y.

122. A review of new developments in the oxidation and hydrolytic reactions of rosin, the origin of condensation and polymerization products, and of resins. Hydrogenated rosin improves its properties, mainly the stability. Condensation products are formed with two molecules. Rosin can be polymerized by means of acids, bases, or heat. Resins are formed by thermal decomposition of rosin with or without catalysts. W. R. Henn

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Recovery of resins from crude resins. A. I. Piryatinskij
and N. V. Tikhovetsky. Russ. Akad. Nauk, Dec. 11, 1940.
The crude resin is exal. with neutral tar acids, oxidized with
air, and the product of oxidation is exal. with petroleum
fractions to sep. unoxidized resin.



CA

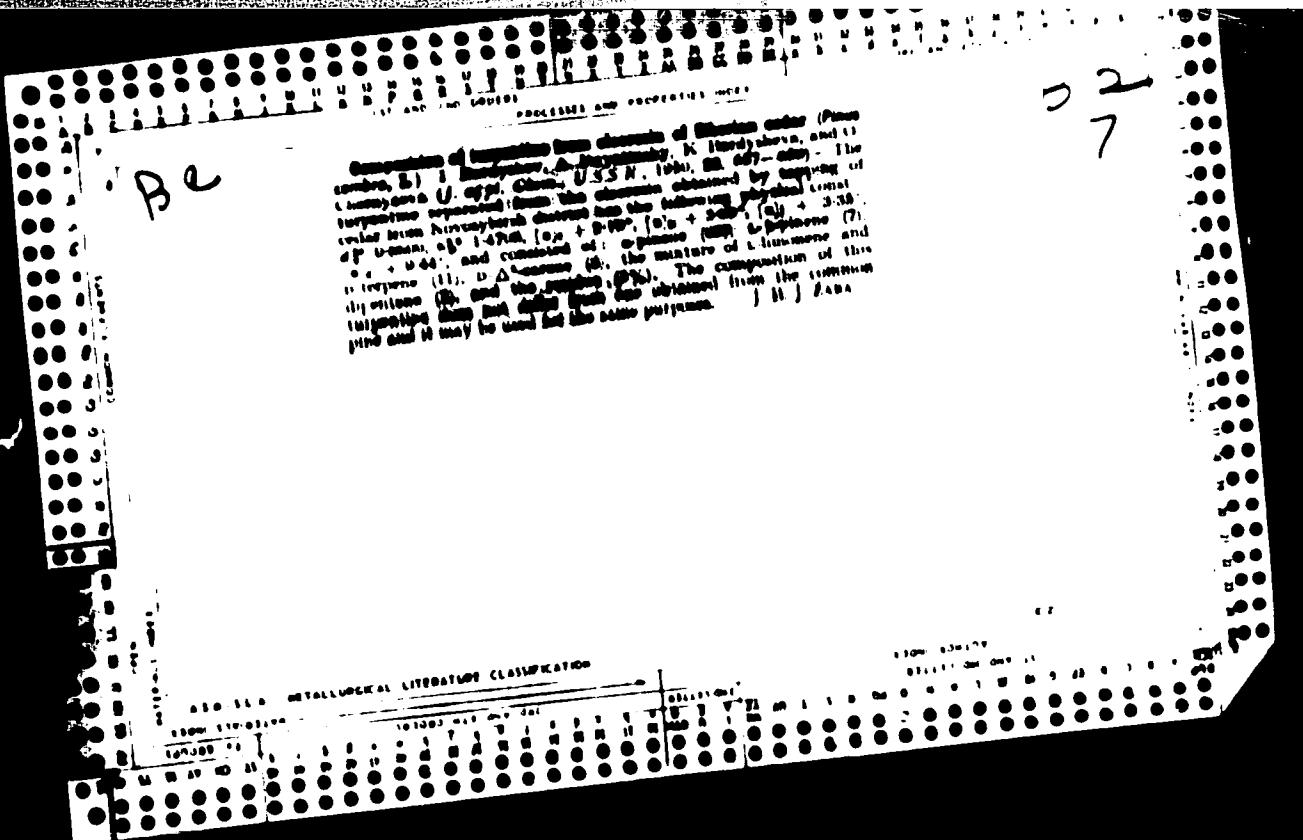
The character of the surface cracks on Dinas products
I. L. Plyatinskii and I. Sh. Shvartunian (*Ogneupory 13,*
211-16(1948); *Chem. Ztbl. (Rusinen Zess. Ed.)* 1949, I,
1155).—Straight cracks may penetrate deeply. Network
cracks, however, are usually less deep. While depth in-
creases with width there is no direct relation. Electrodinas
and open-hearth Dinas showed the greatest no. of straight
cracks; chrome-magnesite brick showed less. Electrodinas
showed the greatest network crazing; magnesite products
showed almost none. The length of the cracks was almost
the same in Dinas and greg products; the cracks were
shorter in magnesite and chrome-magnesite products. With
decreasing d. of the Dinas products, the no. of both straight
and network cracks and their width increased
M. G. Moore

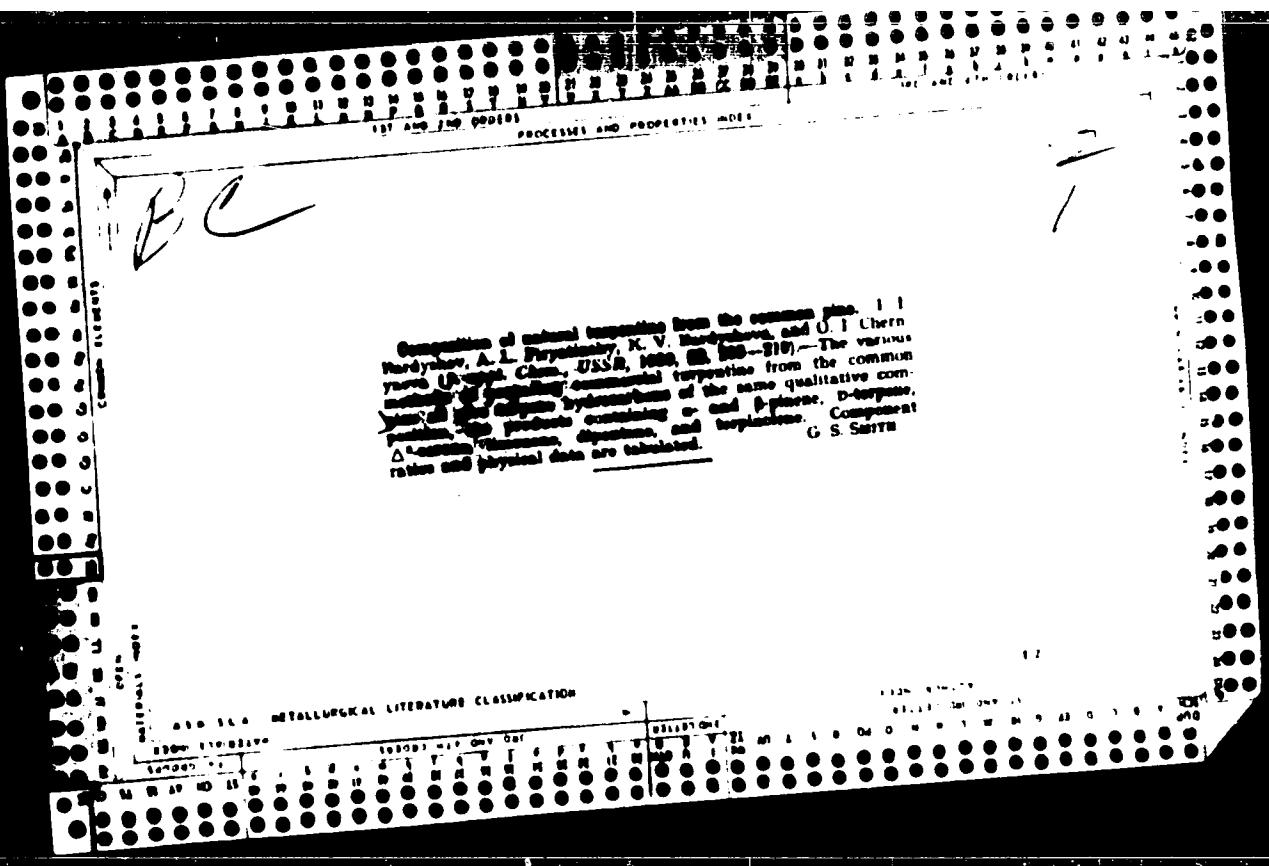
Turpentine, resin and high melting resin from pine or fir waste. V. I. Pyrstunskii and N. G. Lukhovitskii Russ. 52,498,720-737, 1987. The waste is extracted with light petroleum fractions and then with light and neutral tar oils.

ALB 110 METALLURGICAL LITERATURE CLASSIFICATION

1300-111-01000

100000-00





26

2. Tigray. Dura + system

2585 Application of the Darrone method to analysis of *Borod* impregnations. I. Consistency of the ratio $\left(\frac{\text{D}}{\text{L}}\right)_{\text{B}} : \left(\frac{\text{D}}{\text{L}}\right)_{\text{M}}$
I. I. Hardybov and A. L. Tigranyan *J. Appl. Chem. USSR*, 1948, 81, 1174-1179 - Darrone formula,
$$\left(\frac{\text{D}}{\text{L}}\right)_{\text{B}} = \left(\frac{\alpha_{\text{D}} + \alpha_{\text{L}}}{\alpha_{\text{B}}}\right) \cdot \left(\frac{\alpha_{\text{B}}}{\alpha_{\text{M}}}\right)^{\frac{1}{2}},$$
 where α is the percentage content of one constituent of a binary terpene mixture, and α_{D} , α_{L} and α_{B} are the op rotations of the individual constituents and of the mixture, respectively. It is applicable to binary, but not to ternary, mixtures of Δ^3 -carene and D- α - and L- β -pinene. R. Tausz

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"APPROVED FOR RELEASE: 07/13/2001

CIA-RDP86-00513R001341020006-4

REPLACEMENT. 1. L. SWAFFORD, JR., SM.

CC: [redacted]

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SAC: [redacted] IS: [redacted] D: [redacted]

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ALL
1-C

Presence of Δ -caryene in the turpentine from common spruce (Picea sylvestris, L.) I. I. Shandarov, A. Pugatchy, K. Bardysheva, and O. Cherysova (J. appl. Chem., USSR, 1959, 32, 847-852) — Two samples of turpentine were investigated. The first (I) was obtained (1.7 wt.-%) from spruce resin by steam-distillation. The second (II) was obtained (6.3%) from resin which stayed on the wood for a year. In I the fractions distilling up to 215° contain L- α -pinene (~48), L- β -pinene, (17) D- Δ -caryene (4), a mixture of dypnone and humulene (10%), and some substances with high b.p. whose nature has not been investigated. II consists of L- α -pinene (60), L- β -pinene (30), D- Δ -caryene (10), a mixture of dypnone and L-humulene (10%), and some substances with high b.p. whose nature has not been established. Therefore the ratio of components in the turpentine obtained from spruce resin changes with the time it remains on the tree or in storage, but the qual. composition of hydrocarbons does not change. I. B. J. ZABA.

PIRYATINSKIY, L.B.

Modifications of general immunological reactivity under arctic conditions at various seasons. Zhur.mikrobiol.epid. i immun. 28 no.3:65-66 Mr '56. (MLRA 10:6)

(CLIMATE,

immunol. reactions in arctic cond. during various seasons (Rus))

(IMMUNOLOGY,
same)

"The Problem of Changes in General Immunological Reactivity in Polar Regions at Different Periods of the Year," by L. B. Piryatinskiy, Zhurnal Mikrobiologii, Epidemiology, i Immuno-biologii, No 3, Mar 57, pp 75-61.

The author reports on experiments on 321 healthy young persons regarding the resistance of the human organism to disease in the polar regions where, according to a number of physicians who have worked in northern latitudes, certain diseases have specific characteristics. The young people were divided into three groups, one of which was tested during the polar night, the other during the polar day, and the third during both the polar night and the polar day. The tests consisted of injecting intracutaneously one ml of a 1:100 dilution of "so-called antihuman serum" furnished by the Microbiological Department of the Institute of Experimental Medicine, Academy of Medical Sciences USSR. Normal rabbit serum was used as control.

Two tables are included in the article showing the following: for group 1, during the polar night immunological reactivity was lowered positive reactivity was twice that observed during the polar day (97.3 percent against 72.0 percent); for group 2, during the polar day the percentage of positive reactions was not as high as that observed by Ioffe and co-workers, who had found 95 percent positive reactions; and for group 3, during the polar night increased reactivity was found in 55 of the 80 people studied, no change was apparent in 22, and reactivity was lowered in 3 cases. (U)

5000

PIRYAZEV, D.I.

Investigating contact friction forces in the center of deformation
in rolling. Izv. vys. ucheb. zav.; chern. met. 7 no.1:100-106 '54.
1. Ukrainskiy nauchno-issledovatel'skiy institut metallov.
(MIRA 17:2)

СИМЕОНОВ, А.М., АРКАЗЕВ, С.Л.; БУЧЕНКЕВ, А.Р.

Report on working conditions in timbering mills at the "Sibtim" plant, Sverdlovsk district, 1975-1976. (M)

DOLZHENKOV, F.Ye.; KRIVONOSOV, Yu.I.; PIRYAZEV, D.I., VOLCHEK, F.R.;
BAT', Yu.I.

Production of bimetals by the vacuum rolling method. Met.
i gornorud. prom. no.3:34-35 My-Je '64. (MIRA 17:10)

ACC NR AI6009856

SOURCE CODE: UR/0437/65/000/012/D/000145

AUTHOR: Dorzhenkov, F. Ye., Krivonosov, Yu. I., Piryazev, D. I., Bat', Yu. I., Vol'nyak, F. R.

TITLE: Production of bimetal compounds by vacuum rolling

SOURCE: Ref. zh. Metallurgiya, Abs. 12D75

REF SOURCE: Sb. tr. Ukr. n.-i. inst-metall, vyp. 1, 1965, 183-196

TOPIC TAGS: bimetal, metal rolling, titanium, low carbon steel

ABSTRACT: The optimal temperature for commencing the vacuum rolling (R) of Ti on a bimetal is 1000°C. At higher temperatures liquid phase may form. It is preferable to roll at 850°C, since a decrease in temperature leads to a sharp rise in specific pressure, as well as to the occurrence of considerable internal stresses in the bimetal layers. A high C content of steel adversely affects the cohesion to Ti, and hence it is desirable to use a steel with a lower C content as the base-layer (M). Reduction in R temperature and increase in the reduction of area contribute to the decrease of the transition zone of the steel-Ti bimetal during R of two-layer and sandwich packs with the P-plates positioned outermost, the difference

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UDC: 621.771.001

REF ID: A6009956

ACC NR: AR6009956

the deformation of layers increases with increase in reduction of area. As the thickness of the Ti layer decreases, its deformation resistance changes, and this leads to a change in the nonuniformity factor of the plastic deformation of the pack. The broadening of the contact interface of the pack is insignificant, reaching its maximum at the interface. The relation of specific pressure and torque to reduction in area, temperature, thickness ratio and other factors is investigated. 9 illustrations, 1 table. Bibliography of 6 titles. L. Kochenova. [Translation of abstract]

SUB CODE: 13, 11

Cord 2/2

S/137/62/000/001/079/23
A060/A1C1

AUTHORS: Piryazev, I. I., Golubov, M. M., Dabagyan, I. P., Timofeyev, D. I.,
Meleshko, A. M., Kovynev, M. V.

TITLE: The roll separating force of the metal and the loading of the main
motors in the course of rolling on the thick sheet mill 2800

PERIODICAL: Referativnyy zhurnal, Metallurgiya, no. 1, 1962, 4 - 5, abstract ID2:
("St. tr. Ukr. n.-i. in-t metallov", 1961, no. 7, 165 - 177)

TEXT: The authors studied the power conditions for rolling at the thick-
sheet mill 2800 of the Plant imeni Voroshilov. The mill is designed for rolling
sheets with thickness 6 - 50 mm, width 2,500 - 2,600 mm. It consists of a stand
with vertical rolls, a roughing two-high stand with working rolls 1,150 mm dia,
a universal finishing four-high stand 800/1400. The stands are arranged in a
sequence. The roll separating force of the metal in the roughing and the finish-
ing stands was measured by means of force meters with wire tensometers. The
force meters were welded to the pedestals of the working stands on the side of
drive. The pulses from the tensometers were recorded by a magnetoelectric os-
cilloscope ПОБ-14 (POB-14). A calculation of the forces from the torque was

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The roll separating force of...

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carried out to verify the values determined by the force meters. The mean pressures were calculated from the total forces obtained experimentally. Simultaneously with the measurement of the forces, the operation of the main drive motors was oscillographed. The oscillograms recorded the current, voltage, and the number of revolutions of the motors. The investigations have demonstrated that: 1) the separating force of the metal on the rolls of the four-high stand is, in all the cases investigated below the admissible; 2) the closest agreement with the experimental data is given by the values of the mean pressures as calculated by the Golovin-Tyagunov method; 3) the main motors of the mill 2800 are not utilized to full capacity.

G. Grigoryan

[Abstracter's note: Complete translation]

Card 2/2

PAVLOV, I.M.; PIRYAZEV, D.I.

Axial stresses in the cold rolling of pipe. Trudy Inst. met.
no.4:134-140 '60. (MIRA 14:5)
(Rolling (Metalwork))
(Pipe mills)

L 29809-66 EWT(m)/EMP(t), ETI/EMP(k) IJP(c) JD/HW
ACC NM AP6020871 SOURCE CODE: UR/0383/66/000/001/0032/0034

AUTHOR: Piryazov, D. I. (Candidate of technical sciences); Khoroshilov, N. M.;
Krivonosov, Yu. I.; Timofeyev, D. I.; Shul'ga, Ye. A.; Syts'ko, A. A.

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b6
b

ORG: none

TITLE: Variations in the thickness of clad sheet

SOURCE: Metallurgicheskaya i gornorudnaya promyshlennost', no. 1, 1966, 32-34

TOPIC TAGS: metal cladding, sheet metal, metal rolling, metallurgical furnace, thermal conduction, stool/OKh13 stool, Kh17N13M2T stool

ABSTRACT: The authors discuss the variations in thickness of two-layer steel caused by a combination of variations and nonuniformities in the thickness of the individual slabs which make up the pack. These variations may reach $\pm 20\%$ of the nominal value in individual cases. Variations in the thickness was determined for mass produced sheets with a cladding layer of Kh18N10T, Kh17N13M2T and OKh13 steel. The variations in thickness and deviations from nominal value were studied during rolling of bimetal sheet from packs weighing less than 5 tons (small packs) and from packs weighing 10-12 tons (large packs). Sheet rolled from large packs shows less variation in thickness than that rolled from small packets. This is because the large slabs were hot when they were fed into the continuous furnaces and were therefore heated more uniformly. However, completely uniform heating was impossible even in three-zone continuous furnaces. The following furnace conditions are recommended

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ACC NR: AP6020871

for reducing variations in the thickness of plates rolled on the 2800 mill. Temperature of upper and lower sections in the joining zone should be identical: 1300-1310°C; temperature of the soaking zone should be 1260-1270°C. Total heating time should be divided into 40% for preheat, 30% for joining and 30% for soaking.

Experiments showed that planing the slabs on both sides reduced variations in thickness up to approximately 20%. The lubricating interlayer has a low thermal conductivity and impedes heat exchange between the upper and lower parts of the packet during heating which prevents temperature equalization. This causes variations in the thickness of the finished sheet. It was found that the absolute variation in thickness increases with the thickness of the sheets. The relative variations in thickness are approximately the same for sheets of all thicknesses with the exception of 16 mm sheets for which variations are somewhat lower. In 80% of the cases, deviations from the nominal thickness vary within limits from -10 to +12%. The following recommendations are given for reducing deviations from the nominal thickness using existing equipment: bending or by planing on both sides; increasing thickness of the upper slab in the pack by 7% as compared with the lower slab; heating the packets in continuous furnaces with equal temperatures for the upper and lower sections in the joining zone, a temperature of 1260°C in the soaking zone and holding in this zone for 30% of the total heating time. Taking part in the work of the article were TsvNIChM specialists L. V. Moandrov, V. A. Ustimenko, A. V. Tkachev and Komunarskyy Metalurgical Plant specialists S. R. Sarkisyan and A. N. Nesmachnyy. Orig. art. has: 4 figures.

7
SUB CODE: 13, 11 / SUB DATE: nom
Cord 2/2 FV

PAVLOV, I.M.; PIRYATEV, D.I.

Investigating complete pressure on rolls during the cold rolling
of pipe. Trudy Inst. met no. 4:141-149 '60. (MIRA 14:5)
(Rolling (Metalwork)
(Pipe mills)

2025 RELEASE UNDER E.O. 14176
11937.183

AUTHOR: Savlov, I. ... and Artyazev, D. I.
TITLE: Investigation of the total roll pressure during cold
Rolling (cold reducing) of Tubes
PERIODICAL: Akademiya nauk SSSR, Institut metallurgii,
Trudy, No. 4, 1960. Metallurgiya, metallovedeniye,
fiziko-khimicheskie metody i ledovaniya, pp. 141-149
TEXT: The object of the present investigation was to study
the effect of various parameters of the rolling process on the
pressure exerted on the rolls during cold reducing of tubes made
of aluminium alloys D-1 (D-1) and AMg (AMg), brasses F-62
(L-62) and F-68 (L-68), German silver and copper. Rolls
X77-1 (Kh77-1), X77-125 (Kh77-125), X77-52 (Kh77-52) and
X77-75 (Kh77-75) were used in the experiments, and the measure-
ments were carried out with the aid of carbon pressure gauges
accommodated in the housing of the rolls the electrical pulses
generated by the gauges being recorded by a 14-loop oscillograph
TKF-14 (POB-14). The long-term object of the investigation was
to gather data that could be utilized for improvement of the roll
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Investigation of the total roll pressure during cold rolling
(Cold Reducing) of Tubes

pass design developed at katedra prokatki instituta stali
(Mechanical Rolling Department of the Steel Institute). To this
end, the passes in the rolls used in the present investigation were
calculated from the formulae due to I. G. Savlov et al. (ref. 3).

$$t_x = \frac{t_2}{\frac{n_1 - 1}{1 - e^{-n_1}} + 1 - e^{-n_1} \frac{x}{l}} \quad (1)$$

and

$$t_x = \frac{t}{\frac{n_2 - 1}{1 - n_2} + 1 + n_2 \frac{x}{l} + 1} \quad (2)$$

where: t_x = wall thickness (mm) at the given point of the pass;
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Investigation of the Total Roll Pressure During Cold Rolling
(Cold Reducing) of Tubes

t_z - wall thickness (mm) of the stock; $\mu_e = t_z/t_{tp}$ - total reduction in the wall thickness; ℓ - length (mm) of the reducing portion of the pass; x - the coordinate (distance from the wide end) of the given point of the pass (mm); n_1 and n_2 - constants ($n_1 = 0.64$, $n_2 = \cancel{0.1}$). Formula (1) was used to design the roll passes for mills KhPT-2½" and KhPT-75, formula (2) having been used for the two other mills. Some of the results obtained during rolling of alloy AMG (mill KhPT-32) through a tapered pass 34 x 3 - 23 x 1.0 mm (elongation $\mu_0 = 4.32$, feed $m = 8.0$ mm), are reproduced in Fig.1, where the roll pressure P_Σ (kg, left-hand scale, lower curve) and the decrease Δt_x (mm, right-hand scale, upper curve) in the wall thickness are plotted against the distance x (mm), from the leading end of the pass. In Fig.2, P_Σ (kg) is plotted against the distance ℓ_p (mm) from the leading end of the pass, curves 1 and 2 relating respectively to the forward and reverse movement of the rolls of the mill KhPT-75, used for rolling alloy D-16 through a 4-zone pass 54 x 4 -
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Investigation of the Total Roll Pressure During Cold Rolling
(Cold Reducing) of Tubes

35 x 1.75 mm ($m = 10$ mm). In Fig.3, P_{Σ} (kg) during the forward movement of the rolls (mill KhPT-14" used for rolling copper through a pass 40 x 2 - 27 x 0.8 mm) is plotted against feed m (mm), curves 1, 2 and 3 relating to rolling to attain elongation μ_o of 3.0, 3.9 and 5.6 respectively; the variation of P_{Σ} during the reverse movement under the same conditions is similarly illustrated in Fig.4. The effect of elongation, μ_o , is illustrated in Fig.5, where P_{Σ} during the forward movement of the rolls is plotted against μ_o , graphs (a) and (b) relating respectively to points at a distance of 99 and 140 mm from the leading end of the pass: the graphs were constructed for alloy D-1, rolled on mill KhPT-32 through a pass 34 x 3 - 23 x 1 mm ($m = 7.9$ mm). Fig.6 shows P_{Σ} (at $x = 177$ and 53 mm) as a function of the absolute deformation Δt (mm), the data having been obtained during rolling of alloy D-1 on mill KhPT-32 ($\mu_o = 4.13$). Fig.7 shows P_{Σ} (at $x = 201.5$ and 59.5) as a function of the relative deformation $\Delta t/t \times 100$. the curves

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Investigation of the Total Roll Pressure During Cold Rolling
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having been constructed for copper rolled through a pass $32 \times 3 - 20 \times 1$ mm ($\mu_0 = 4.65$). In Fig. 8, P_{Σ} at $x = 94.7$ mm (curve 1) and $x = 235.7$ mm (curve 2) is plotted against the wall thickness t_z (mm) of the stock, this graph relates to brass L-62 rolled through a pass $38 \times 3 - 25 \times 1$ mm (forward movement). The results reproduced in Fig. 9, where P_{Σ} is plotted against the rolling speed n (reciprocal revs/min), relate to alloy AMG, rolled on mill KhPT-32, through a pass $29 \times 3 - 18 \times 0.8$ mm ($m = 7.8$ mm). Finally, the results of lubricating tests are reproduced in Fig. 10, where P_{Σ} is plotted against various types of lubricants used in the rolling of brass L-68 on mill KhPT- F_2 " through a pass $36 \times 3 - 24 \times 1$ mm ($\mu_0 = 4.65$, $m = 8.3$ mm), curves I and II relating to the forward and reverse movement respectively. The type of lubricant is shown as follows: open circles - oil/graphite mixture, full circles - solidol; full triangle - emulsol, full circle (on the extreme left) - mineral oil. The following conclusions were reached.

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Investigation of the Total Roll Pressure During Cold Rolling
(Cold Reducing) of Tubes

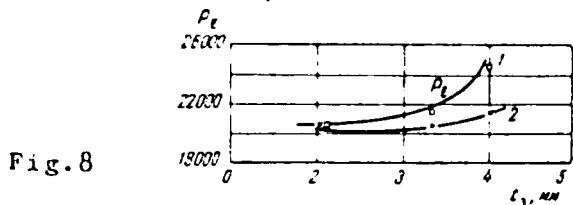
- (1) Irrespective of the size of the mill and type of alloy rolled, more favourable distribution of the roll pressure along the pass is obtained if instead of a 4-zone pass a tapered pass calculated from the formulae (1) and (2) is used. Since the maximum roll pressure in a tapered pass is 1.5 times lower than that in a 4-zone pass, the introduction of the former in industrial practice should increase the output of the mill and improve the quality of the product. (2) A two-fold increase in the feed increases the roll pressure by a factor of 1.3-1.5. (3) In rolling tubes to the final wall thickness > 1.3 mm, the increase in the roll pressure due to increased feed is approximately the same as that due to increased elongation, when the final wall thickness is below 1.3 mm the effect of elongation becomes more pronounced. (4) Doubling the wall thickness of the stock increases the roll pressure by a factor of 1.2 during the forward movement, and by a factor of 1.3 during the reverse movement of the rolls.
- (5) Within the range of the rolling speeds studied

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(10-80 reciprocal revs/min), the roll pressure remains practically constant. (6) Best results (lowest roll pressure) are obtained when an oil/graphite mixture is used for lubrication. However, this lubricant is difficult to remove from the finished product, and the application of emulsol or solidol is recommended instead. There are 10 figures, 1 table and 4 Soviet references.



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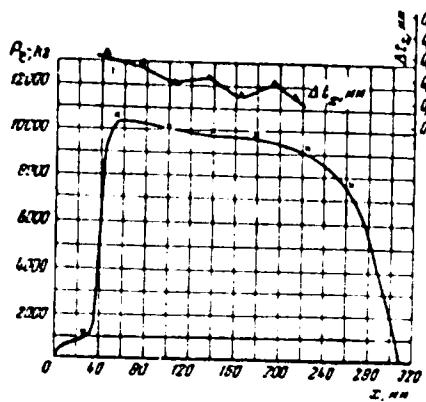


Fig. 1

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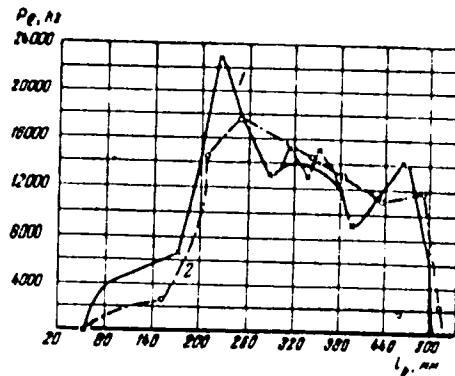
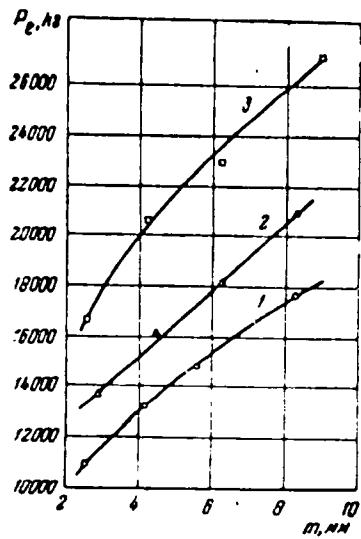


Fig. 2

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Fig. 3

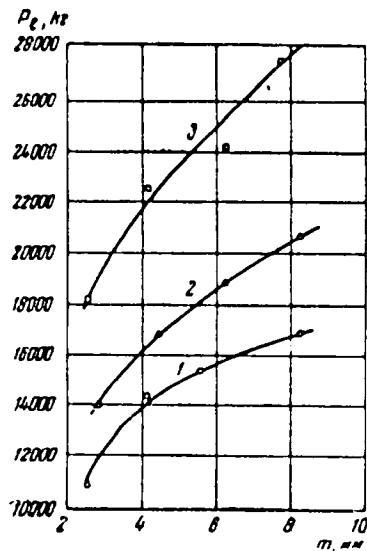


Fig. 4

Investigation of the Total Roll... S/509/60/000/004/011/024
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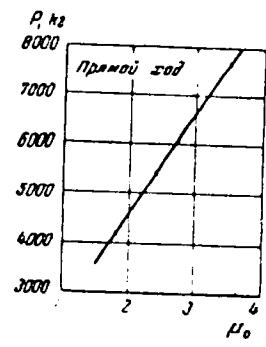
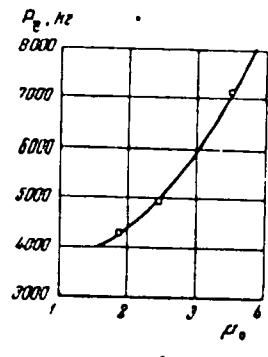


Fig. 5

a



b

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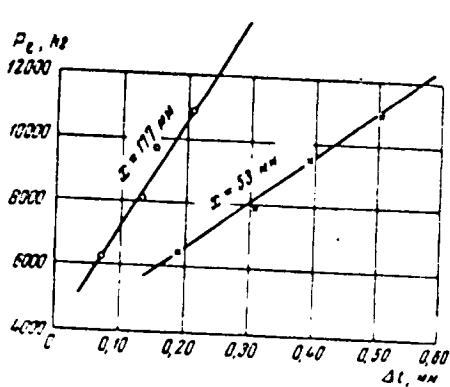


Рис. 6 Зависимость P_t от Δt

Fig. 6

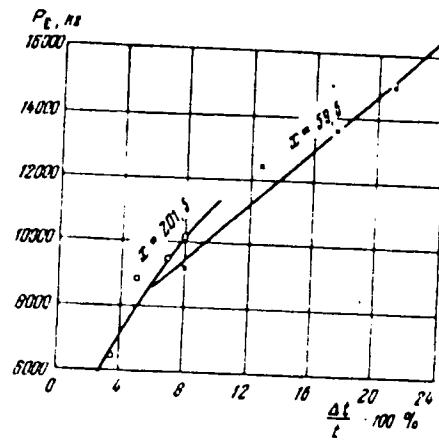


Рис. 7 Зависимость P_t от $\Delta t \cdot 100$ пр

Fig. 7

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Investigation of the Total Roll....

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Fig. 9

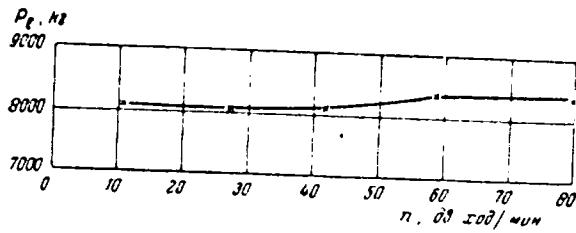


Fig. 10

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PURPOSE, I.I.

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"APPROVED FOR RELEASE: 07/13/2001

CIA-RDP86-00513R001341020006-4

PAVLOV, I.M.; PIRYAZEV, D.I.

Unit pressure in the cold rolling of tubes. Trudy Inst. met.
no.4:123-134 '60. (MIRA 14:^c)
(Pipe mills)

APPROVED FOR RELEASE: 07/13/2001

CIA-RDP86-00513R001341020006-4"

S 276.000 OCT 1973 042
AUG 1973

AUTHOR: Rizayer I.

TITLE: Investigating metal pressure on rolled and the main thickness
during rolling in a 3000 tonne mill

PERIODICAL: Referativnyy zhurnal Metalurgiya i metalloobrabotka
"Tr. Konferencii Tekhnicheskogo komiteta po izuchenii
vysokikh metallicheskikh struktur"

TEXT: Metal pressure in rolling was made at 1000 tonne rolling mills on 100 mm
sheets of various steel grades. Data given in tables show that the pressure in
the two-high stand is distributed non-uniformly over the passes. Maximum force
here is in the middle of the passes received in the last passes (1.14 kg/mm²)
in the two-high stand and 0.75 kg/mm² in the first stand. The admissible
pressure of these rolling mills is 1.0 kg/mm² in the first two passes (1.0 kg/mm²)
and 0.75 kg/mm² in the last passes. In a four-high stand the pressure
is distributed uniformly. Metal pressure in rolling was also tested by the formula
 $P = \frac{F}{t^2}$. The pressure in rolling was 1.0 kg/mm². The exact metal
thickness was 100 mm. The pressure in rolling was 1.0 kg/mm². The thickness of the metal
and the calculated data are in a satisfactory agreement. The author also points

that in

Investigating the following:

1. 17 SEP 1971
2. 17 SEP 1971

Turned up in the course of investigation that the location of RSP-1000 was 'West', or 'West', in the area square number 10 RSP-1000 is the location of the current location of the data center. It is also present in the file that the location of the data center is located in the same area.

3. 17 SEP 1971

A set of recommendations for further investigation.

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PIRYAZEV, D.I.; ALEKSANDROV, P.A.

Unit pressures in hot rolling and the analysis of formulas and
methods for their determination. Trudy Ukr. nauch.-issl. inst.
(MIRA 14:3)
met. no.6:157-170 '60.
(Rolling (Metalwork))

PIRYAZEV, D.I., kand.tekhn.nauk; GOLUHOV, M.M., inzh.; DABAGYAN, I.P., inzh.;
TIMOFEEV, D.I., inzh.; MELESHKO, A.M., inzh.; KOVYNEV, M.V., inzh.;
Prinimali uchastiye: VOLCHEK, F.R.; SOKOLOV, B.A.; KRIVONOSOV, Yu.I.

Metal pressure on rolls and loading of the main motors during the
operation of 2800 plate rolling mills. Trudy Ukr. nauch.-issl.
inst. met. no.7:165-176 '71. (MIRA 1+11)
(Rolling mills--Electric driving)

PIRYAEV, D.I.

Investigation of slipping, contact friction and forces in the
deformation center. Trudy Ukr. nauch.-issl.inst. met. no.6:171-
179 '60. (MIRA L4:3)
(Rolling mills)(Deformations(Mechanics))

S/509/60/000/004/010/024
E193/E183

AUTHORS: Pavlov, I.M., and Piryazev, D.I.

TITLE: Axial Loads in Cold Rolling (Cold Reducing) of Tubes

PERIODICAL: Akademiya nauk SSSR. Institut metallurgii.
Trudy, No. 4, 1960. Metallurgiya, metallovedeniye,
fiziko-khimicheskiye metody issledovaniya, pp.135-140.

TEXT: Many of the mechanical failures, encountered in the cold-reducing process (seizure of the stock, bending of the rod supporting the mandrel, excessive wear of various parts of the feeding mechanism) are caused by axial loads which, in addition, constitute a factor limiting the protective capacity of the mill. It was for these reasons that the present investigation, concerned with axial loads in rolling non-ferrous metals and alloys, was undertaken. The measurements were carried out on cold-reducing mills XPT-1 $\frac{1}{2}$ " (KhPT-1 $\frac{1}{2}$ ") and XPT-2 $\frac{1}{2}$ " (KhPT-2 $\frac{1}{2}$ "), used for rolling copper and brass tubes. The axial loads, acting directly on stock, were measured with the aid of carbon pressure gauges, mounted in a special device attached to the end of the stock. In the case of mill KhPT-1 $\frac{1}{2}$, only the Card 1/9

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Axial Loads in Cold Rolling (Cold Reducing) of Tubes

compressive loads were measured; the device used during rolling on mill KhPT- $2\frac{1}{2}$ " was designed to measure both compressive and tensile loads. A general view of this device is reproduced in Fig.1, which shows a cylinder (1) to which the stock (2) was rigidly attached, and flanges (3) and (4); the compressive loads were measured with the aid of three carbon gauges (5), similar gauges of the membrane type having been used to measure the tensile loads. The electric pulses generated by the gauges were recorded with the aid of a magneto-electric oscillograph №6-14 (POB-14). In addition to the axial loads, the roll pressure was also determined. In the case of mill KhPT- $1\frac{1}{2}$ ", the measurements were carried out during rolling of copper and brass tubes through six different passes. Mill KhPT- $2\frac{1}{2}$ " was used to study the variation of axial loads during rolling of brass tubes through a tapered pass (61 x 6 - 36 x 3 mm) and through a 4-zone pass (61 x 6 - 38 x 3 mm). Some of the typical results are reproduced graphically. In Fig.2, the roll pressure, P_{Σ} (kg, left-hand scale) is plotted against the distance, x (mm) from the leading

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Axial Loads in Cold Rolling (Cold Reducing) of Tubes

end of the pass, curves 1 and 2 relating to the forward and reverse movements of the rolls respectively. Similarly, curves 3 (forward movement) and 4 (reverse movement) show the variation of the axial load, Q_{Σ} (kg, right-hand scale). The results, reproduced in Fig.2, relate to copper tubes rolled on mill KhPT-1 $\frac{1}{2}$ " through a pass 40 x 3 - 27 x 0.8 mm, the other rolling parameters being μ_0 (elongation) = 3.9 and m (feed) = 8.3 mm. The results for brass П-68 (L-68) rolled on mill KhPT-2 $\frac{1}{2}$ " through a 4-zone pass 61 x 6 - 38 x 3.0 mm (μ_0 = 2.9, m = 4 mm) are reproduced in the same manner in Fig.3, except that in this case P_{Σ} is given in tons. In Fig.4, the axial load Q_{Σ} (kg) is plotted against the distance x (mm) from the leading end of the pass, curves 1 and 2 relating respectively to the forward and reverse movement during rolling of brass L-68 through a tapered pass 61 x 6 - 36 x 3 mm (μ_0 = 3.5, m = 4.0 mm). The combined effect of the variation of feed, m , and elongation, μ_0 , on Q_{Σ} (kg) during rolling of copper (reverse movement) on mill KhPT-1 $\frac{1}{2}$ ", through a pass 40 x 2 - 27 x 0.8 mm, is plotted against m (mm), curves 1, 2 and Card 3/9

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Axial Loads in Cold Rolling (Cold Reducing) of Tubes

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3 relating to $\mu_0 = 3.0, 3.9 and 5.6 respectively, see Fig.5). In Fig.6, Q_Σ (kg) during rolling of brass L-68 on mill KhPT-1 $\frac{1}{2}$ " through a pass 36 x 3 - 24 x 1 mm ($\mu_0 = 3.9$) is plotted against m (mm), curves 1 and 2 relating respectively to points at a distance of 154.7 mm from the leading end of the pass (forward movement) and 126.7 mm (reverse movement). In the final experiments, the effect of various lubricants on Q_Σ was studied. The results, obtained during rolling of brass L-68 on mill KhPT-1 $\frac{1}{2}$ " through a tapered pass 36 x 3 - 24 x 1 mm ($\mu_0 = 3.9, m = 8.3$ mm), are reproduced in Fig.7, showing the variation of Q_Σ due to change of the lubricant, curves 1 and 2 having been constructed for the forward and reverse movement of the rolls, and the experimental points relating to an oil/graphite mixture (open circles), solidol (full circles), emulsol (full triangles), and mineral oil (full squares). The main conclusions reached by the present authors can be summarised as follows. (1) In analogy to the roll pressure, the axial loads during cold reducing of tubes vary along the pass. The axial loads during the reverse movement are considerably higher$

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Axial Loads in Cold Rolling (Cold Reducing) of Tubes

than those during the forward movement rolls, constituting 8-10% of the roll pressure in the former, and only 2.5-6% in the latter case. If, therefore, seizure of the stock occurs, it probably takes place during the reverse movement of the rolls.

(2) Two-fold increase in the feed increases the axial loads 1.5-1.8 times; a similar increase in the wall thickness of the stock increases the axial loads by a factor of 2.3.

(3) Minimum axial loads are ensured by using an oil/graphite mixture for lubrication; mineral oil, used for this purpose, raises the magnitude of the axial loads to its maximum. There are 7 figures, 2 tables and 2 references: 1 Soviet and 1 German.

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Axial Loads in Cold Rolling....

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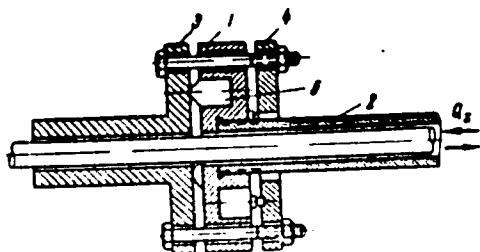


Fig. 1

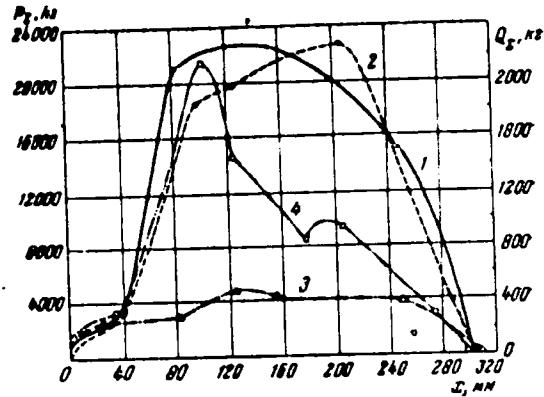
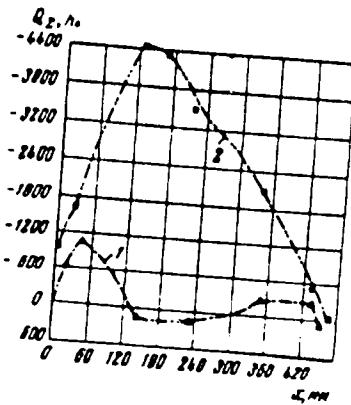
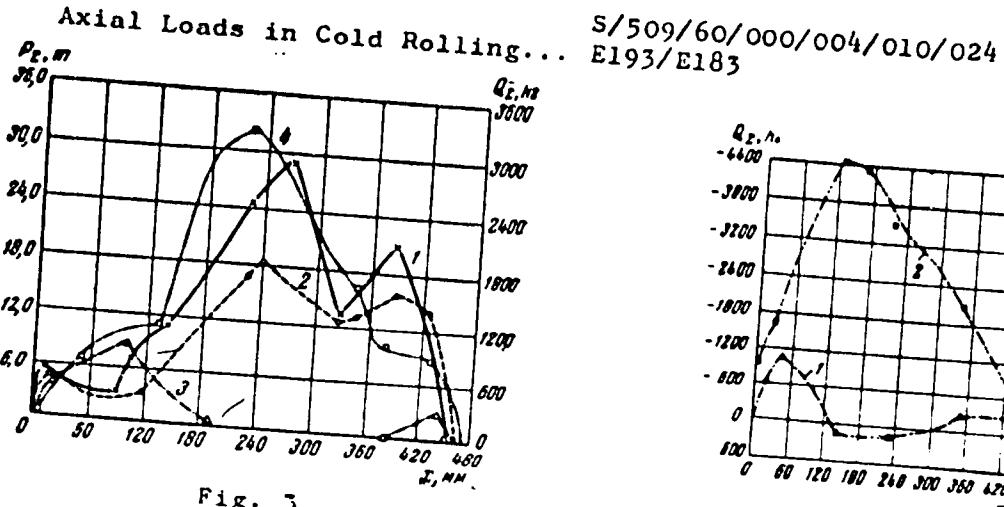


Fig. 2

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Axial Loads in Cold Rolling... S/509/60/000/004/010/024
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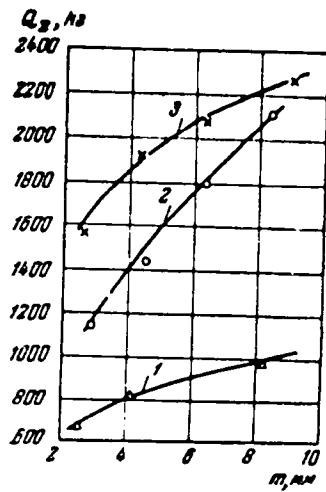


Fig. 5

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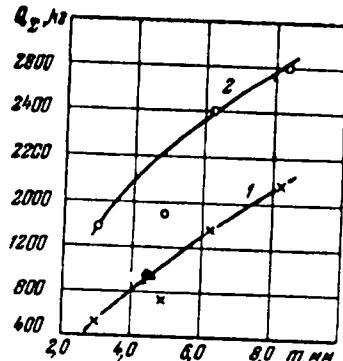


Fig. 6

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Axial Loads in Cold Rolling (Cold Reducing) of Tubes

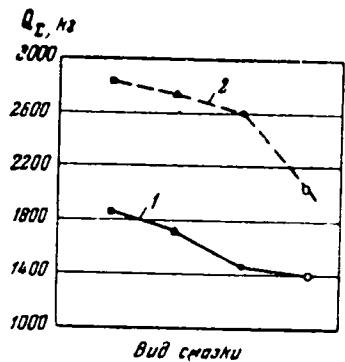


Fig. 7

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2195 2185

AUTHORS: Pavlov, I.N., and Piryazev D.I.

TITLE: Specific Pressure in Cold Rolling (Cold Reducing) of Tubes

PERIODICAL: Akademiya nauk SSSR. Institut metallurgii
Trudy, No.4, 1960 Metallurgiya metallovedeniye,
fiziko-khimicheskiye metody issledovaniya, pp 123-134

TEXT: Problems such as the determination of the roll pressure in tube rolling, roll pass design, and assessment of the degree of wear of various parts of the rolling mill, become easier to deal with if data on the magnitude and distribution of specific pressure are available and if it is known how these parameters are affected by other variables of the process. Since the only experimental data on this subject are those due to Yu.F. Shevakin (Ref.5) the investigation described in the present paper was undertaken in order to study the effect of feed, elongation, and the magnitude of absolute and relative deformation on the specific pressure and its distribution along both the deformation region (contact zone) and the roll pass (reducing

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Specific Pressure in Cold Rolling (Cold Reducing) of Tubes
groove). In addition, the average magnitude of specific pressure was determined, and an attempt was made analytically to solve the problem of distribution of pressure in the deformation region. The measurements of the specific pressure were carried out under industrial conditions on a cold-reducing mill XMT-32 (KhPT-32). Specially designed rolls (300 mm in diameter) permitted direct determination of the pressure at six points of the pass with the aid of six carbon pressure gauges of the membrane type constructed by TsNIITMASH. Fig.1 shows the expanded pass with the location of the pressure gauges indicated by dots and their distance from the wide end of the pass given in mm. Each of the two semi-circular rolls accommodated three of these gauges in the manner shown in Fig.2. All gauges were located in the plane of the crown of the pass the problem of distribution of pressure across the groove being outside the scope of this investigation. The electrical pulses generated by the pressure gauges were recorded on a photographic film with the aid of a magneto-electric oscillograph N^S-14 (POB-14). The groove and the mandrel were

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Specific Pressure in Cold Rolling (Cold Reducing) of Tubes

designed to give a pass which tapered from 34 x 3.0 to 23 x 1.0 mm. The pressure measurements were carried out during rolling of tubes of aluminium alloys A-1 (AMG), D-1 (D-1) and D-16 (D-16). The stock (33.2 outside diameter 3.0-3.2 mm wall thickness) was rolled to the following final sizes: 23 x 0.75, 23 x 0.83, 23 x 1.0, 23 x 1.1, 23 x 1.5, and 23 x 1.75 mm. Both the roll grooves and inside walls of the tubes were lubricated with mineral oil. The magnitude of feed was determined from the number of reversals per 100 mm of the length of the stock rolled. Owing to the difficulties encountered in measuring the pressure at normal rolling speeds a speed of 10-12 reciprocal revs/min was used in the experiments. In addition to the specific pressure, the total roll pressure was measured with the aid of a gauge accommodated in the roll housing. Preliminary to experiments proper, a formula was derived for the critical angle, β , in the plane of the groove crown, and the values of this angle and of the contact angle θ_0 , were calculated for various feeds, m . It was shown that at small m (e.g. $m = 1.5$ mm) $\beta < \theta_0$ for the entire length of the

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