

POZNANSKAYA, A.A., Cand Biol Sci -- (diss) "The effect of biotin deficiency upon processes ^{of formation} ~~in~~ the ~~development~~ of certain proteins in animal tissues." Mos, 1958. 15 pp (Acad Med Sci USSR Inst of Biol and Med Chemistry). 200 copies.
(KL, 12-58, 97)

-32-

POZNANSKAYA, A.A.

Effect of biotin deficiency in chicks on the synthesis of serum albumin in liver sections and on amylase synthesis in sections of the pancreas [with summary in English]. Biokhimiia 22 no.4:668-676 (MIRA 10:11)
Jl-Ag '57.

1. Laboratoriya obmena azotistykh veshchestv i Institut biologicheskoy i meditsinskoy khimii AMN SSSR, Moskva.

(PANCREAS, metabolism,

amylase in slices from biotin-defic. chicks (Rus))

(AMYLASES,

in pancreatic slices from biotin-defic. chicks (Rus))

(BIOTIN, deficiency,

eff. on liver serum albumin & pancreas amylases in slices from defic. chicks (Rus))

(LIVER, metabolism,

serum albumin in slices from biotin-defic. chicks (Rus))

(SERUM ALBUMIN, metabolism,

liver, in slices from biotin-defic. chicks (Rus))

USSR / Human and Animal Physiology (Normal and Pathological).
Metabolism.

T-3

Abs Jour : Ref Zhur - Biologiya, No 13, 1958, No. 60102

Author : Poznanskaya, A. A.

Inst : Not given

Title : Effect of Biotin Deficiency in Chicks Upon the Synthesis
of Serum Albumin in Liver Sections and Amylase in
Pancreatic Sections

Orig Pub : Biokhimiya, 1957, 22, No 4, 668-676

Abstract : Chicks were kept on a special diet producing biotin
deficiency. When the signs of avitaminosis became pro-
nounced, the animals were killed and the ability of
pancreatic (P) sections and liver sections to synthesize,
respectively, amylase (I) and serum albumin from glucose
and essential amino acids was studied. In chicks with
avitaminosis, the protein synthesis was almost completely

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USSR / Human and Animal Physiology (Normal and Pathological).
Metabolism.

T-3

Abs Jour : Ref Zhur - Biologiya, No 13, 1958, No. 60102

inhibited. The addition of α -ketoglutarate re-established the ability of the P sections to synthesize I, but did not affect the ability of the liver section to synthesize albumin; the latter process was restored by the addition of glutamine to the sections. The impairment of the capacity of normal P slices in the synthesis of I, produced in vitro by substances inhibiting the citric cycle (mesotartarate, fluoroacetate), could also be counteracted by the addition of α -ketoglutarate. --
V. I. Rozengart

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BRAUNSHTEYN, A.Ye.; GHUGHEV, N.V.; POZNANSKAYA, A.A.

Nonenzymatic reamination of δ -aminovaleric acid. Dokl. AN SSSR
152 no.5:1239-1242 0 '63. (MIRA 16:12)

1. Institut radiatsionnoy i fiziko-khimicheskoy biologii AN SSSR.
2. Chlen-korrespondent AN SSSR (for Braunshteyn).

5(3), 17(3)
AUTHORS:

SOV/20-125-4-67/74
Rozenfel'd, Ye. L., Poznanskaya, A. A., Rudakova, N. K.

TITLE:

A Study of the Composition and Properties of Zymosan (Izucheniye sostava i svoystv zimozana)

PERIODICAL:

Doklady Akademii nauk SSSR, 1959, Vol 125, Nr 4, pp 928-930 (USSR)

ABSTRACT:

The polysaccharide zymosan and the complex it forms with the newly discovered serum protein properdin (Refs 1,2), which plays an important role in natural immunity, are being more and more investigated. The authors examined a zymosan preparation (Nr 1) which is active with regard to the properdin system and which Mrs. R. A. Rutberg obtained from ordinary yeast by her modified method (Ref 5). 2 fractions were obtained from the zymosan, which were conditionally named: a) soluble and b) insoluble. It has been found that the nitrogen content of both fractions is considerably lower than in the original zymosan preparation. Table 1 shows the results of further investigations. As may be seen from it, the soluble fraction of zymosan consists of glucose and mannose, whereas the insoluble is a glucan. It is evident from figure 1 that the soluble fraction consists of 2 fractions, A and B, differing by their

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A Study of the Composition and Properties of Zymosan

electrophoretic motility. The proportion of glucose and mannose in fraction A is 1 : 1.6, in fraction B 1.6 : 1. Therefore the soluble zymosan fraction is composed of 2 glucomannans differing by their structure. In chromatography the insoluble fraction consisting of glucose alone shows after a partly acid hydrolysis (30 mins in 0.5 n HCl) 3 spots in the hydrolysate, 2 of which apparently are due to disaccharides. It is possible that in glucan there are no less than 2 types of glucoside bonds between remains of glucose, or else that fraction constitutes a mixture of 2 glucans differing by the character of bonds. Therefore zymosan is a complicated mixture of polysaccharides differing by their composition and structure, namely a glucan (or glucans), and two different glucomannans. The composition of zymosan is explained by a diagram. There are 1 figure, 1 table, and 7 references, 2 of which are Soviet.

ASSOCIATION: Tsentral'nyy institut gematologii i perelivaniya krovi
(Central Institute of Hematology and Blood Transfusion)
Card 2/3 Laboratoriya fiziologicheskoy khimii Akademii nauk SSSR

SOV/20-125-4-67/74

A Study of the Composition and Properties of Zymosan

(Laboratory of Physiological Chemistry of the Academy of
Sciences USSR)

PRESENTED: December 8, 1958, by A. I. Oparin, Academician

SUBMITTED: December 4, 1958

Card 3/3

POZNANSKAYA, A.A.; GORKIN, V.Z.

Modern concepts of the role of biotin in metabolism; participation
of biotin in carbon dioxide fixation. Vop. med. khim. 8 no.2:115-131
Mr-Apr '62. (MIRA 15:4)

1. Institut biologicheskoy i meditsinskoy khimii ANI SSSR, Moskva.
(CARBOXYLATION) (BIOTIN)

POZNANSKAYA, A.A.

Effect of biotin deficiency on induced increase in the activity of the tryptophan-peroxidase system in rat liver [with summary in English]. *Biokhimiya* 23 no.2:230-233 Mr-Apr '58 (MIRA 11:6)

1. Laboratoriya obmena azotistykh veshchestv Instituta biologicheskoy i meditsinskoy khimii AMN SSSR, Moskva.

(BIOTIN, deficiency

exper., eff. on activity of liver tryptophan-peroxidase system in rats (Rus))

(LIVER, metabolism

tryptophan-peroxidase system activity, eff. of exper. biotin defic. in rats (Rus))

(OXIDASES, metabolism

tryptophan-peroxidase system activity in liver, eff. of exper. biotin defic. in rats (Rus))

POZNANSKAYA, A.A.

YEFIMOVICHKINA, Ye.F.; POZNANSKAYA, A.A.

Biological synthesis of purine and pyrimidine substances and mononucleotides. Vop.med.khim. 3 no.4:243-254 J1-Ag '57.

(MIRA 10:11)

1. Laboratoriya obmena azotistykh soedineniy Instituta biologicheskoy i meditsinskoy khimii Akademii meditsinskikh nauk SSSR, Moskva.

(PURINES, metabolism,
biosynthesis, review (Rus))

(PYRIMIDINES, metabolism,
same)

(NUCLEOSIDES AND NUCLEOTIDES, metabolism,
mononucleotides, biosynthesis, review (Rus))

POZNANSKAYA, A.A.; ROZENFEL'D, Ye.L.

Composition and properties of different zymosan preparations. *Biokhimiia*
25 no.4:624-629 J1-Ag '60. (MIRA 13:11)

1. Institute of Hematology and Blood Transfusion, and Institute of
Biological and Medical Chemistry, Academy of Medical Sciences of
the U.S.S.R, Moscow.

(ZYMOSAN)

POZNAJSKAYA, A. A., and NEYMAN, L. A. (U.S.S.R.)

"New Routes of Synthesis of 4,5-Dioxovaleric (Glyoxalylpropionic) Acid."

Report presented at the 5th International Biochemistry Congress,
Moscow, 10-16 Aug 1961

POZNANSKA, Hanna

Behavior of fructose and inorganic phosphorus in the blood following oral loading with saccharose as the index of the fructose metabolism. Intermediately determination of the liver fructokinase activity. Pol. tyg. lek. 19 no. 44: 1675-1677 ■ 2'64

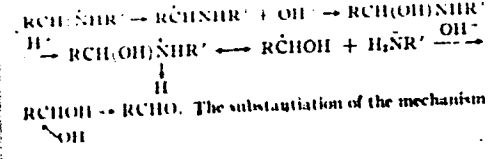
1. Z II Kliniki Chorob Zakaznych Akademii Medycznej w Warszawie (Kierownik: prof. dr. med. B. Kaszur).

Azomethines I Hydrolysis reaction of azomethines

B. A. Pocat Koshity, R. M. Poznanovskaya, V. S. Shevchenko, and I. A. Pavlova (Leningrad Technol. Inst.), *J. Gen. Chem. (U.S.S.R.)* 17, 1774-87 (1947) (in Russian).

--The behavior of Schiff bases from aromatic compds. on acid and alk. hydrolysis was investigated. In the majority of cases there is a direct relation between the rate of acid hydrolysis and the basicity of the compd.; the latter most frequently depends on the basicity of the amine used. Theoretical examn. of the mechanism indi-

cates that the major role in acid hydrolysis is played by the protons. An approx. calcul. of the magnitudes of the resonance effect and the alternate charge effect and of the field effect of the substituted groups showed that for the NO₂ group the resonance effect is 4.5 times greater than the field effect, which is 1.5 times greater than the alternate effect. The acid hydrolysis is represented as follows:



is seen in the dependence of the rate of hydrolysis on the basicity of the base used. In alk. soln. the primary reaction is addn. of OH⁻, followed by: RCH(OH)NHR' → RCHOH + NHR' → RNH₂ + RCHOH → RCHO + OH⁻

where the addn. of OH⁻ is simultaneous with polarization of the CN bond; support for this is seen in more difficult hydrolysis in alk. soln. than in acid soln. if such groups are introduced which are not acidic enough to counteract the basic properties of the azomethine. Hydrolyses in 0.1 N HCl were followed at room temp. and 50° by analysis of free amine and aldehyde. The following values were obtained (the figures are in the order: hydrolysis rate consts. at room temp. and 50°, activation energy rate consts. at room temp. and 50°, N-benzylideneamine, 1.3 × 10⁻⁴, 2.0 × 10⁻⁴, 4800, 5.3 × 10⁻¹⁰; N-(p-nitrobenzylidene)aniline, 1.7 × 10⁻⁴, 1.0 × 10⁻⁴, 15,510, 5.3 × 10⁻¹⁰; N-(m-nitrobenzylidene)aniline, 2.2 × 10⁻⁴, 5.7 × 10⁻⁴, 4448, 5.3 × 10⁻¹⁰; N-benzylidene-p-nitroaniline, 7.0 × 10⁻⁴, 1.24 × 10⁻¹⁰; N-benzylidene-m-nitroaniline, 1.2 × 10⁻⁴, 5.07 × 10⁻¹⁰; N-(p-nitrobenzylidene)-m-nitroaniline, 1 × 10⁻⁴, 1 × 10⁻¹⁰; N-(p-nitrobenzylidene)-p-nitroaniline, 0, 1.24 × 10⁻¹⁰; p-benzylideneaminobenzoic acid, 1.2 × 10⁻⁴, 8.15 × 10⁻¹⁰; (O,N,N), 1.0 × 10⁻⁴, m-benzylideneaminobenzoic acid, (over)

ASH-31A METALLURGICAL LITERATURE CLASSIFICATION

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

Also dyes from 5-amino-1-naphthol and some of its derivatives II. Determination of the structures of monoazo dyes from 5-amino-1-naphthol. A. I. Poral-Koshits, B. A. Poral-Koshits, and V. V. Perekalin (Leningrad Chem. Technol. Inst.). *J. Gen. Chem. (U.S.S.R.)* 17, 1768-67 (1947) (in Russian); cf. *C.I.* 40, 4885.

In Part I it was shown that coupling of 5,1-H₂NC₁₀H₇OH with PhN₂Cl and *p*-O₂NC₆H₄N₂Cl yielded 3 isomers for each reagent. These were given the provisional assignment of 2-, 4-, and 6-isomers. The establishment of some of the structures has now been made on the basis of the following observations. *N*-1-Naphthylphthalimide, *N*-1-naphthylphthalamic acid, and *N*-(3-nitrobenzylidene)-1-naphthylamine are incapable of azo coupling. Dyes obtained by coupling *N*-(5-hydroxy-1-naphthyl)phthalamic acid with PhN₂Cl are identical with dyes prepd. by phthaloylation of 6-phenylazo-5-amino-1-naphthol and 8-phenylazo-5-amino-1-naphthol. Dyes from PhN₂Cl and the 5-(3-nitrobenzylideneamino)-1-naphthol are identical with the products of the reaction of 3-O₂NC₆H₄CHO with 6-phenylazo-5-amino-1-naphthol and 8-phenylazo-5-amino-1-naphthol. *N*-(6-Nitroso-5-hydroxy-1-naphthyl)phthalamic acid does not react with diazo compounds. Replacement of the H atoms of the NH₂ group in the new dyes by acyl or benzylidene groups causes a hypsochromic shift; the color becomes deeper. When 5-amino-1-naphthol (1.50 g.) was fused with 1.48 g. phthalic anhydride, the reaction was complete in 1-2 min and the product, pptd. from 10% NaOH by HCl, yielded 100% *N*-(5-hydroxy-1-naphthyl)phthalamic acid (1), needles, m. 210° (from EtOH). 1 (3.00 g.) in 12 ml. 10% NaOH dild. to 100 ml., coupled with 100 ml. 0.1 *N* PhN₂Cl, and acidified with 10% HCl gave 80% crude dye; this was extd. with 100 ml. hot Me₂CO, followed by 50 ml. Me₂CO and the residue was repptd. from 10% NaOH by 10% HCl, while the dye in Me₂CO was purified in a similar manner. The product, m. 172-4°, was identical with that obtained by heating 2.63 g. of the dye m. 181° (see part I) with 1.48 g. phthalic anhydride 1-2 min. until a melt is achieved; this establishes the structure *N*-(6-phenylazo-5-hydroxy-1-naphthyl)phthalamic acid. The other product, violet, m. 215°, was identical with the product of similar phthaloylation of the dye m. 195° (see part I); this established the structure *N*-(8-phenylazo-5-hydroxy-1-naphthyl)phthalamic acid. Attempted coupling of the above reagents in HCl failed to take place. *m*-O₂NC₆H₄CHO (8.4 g.) in 10 ml. warm AcOH was added to 6 g. 1,5 HOC₆H₄NH₂ HCl in 30 ml. 50% AcOH, giving on cooling 100% *o*-nitrobenzylideneamino-1-naphthol, green plates, m. 108° (from alc.). This (2.0 g.), in 8 ml. 10% NaOH and 20 ml. EtOH dild. to 100 ml. with water, was mixed with 100 ml. 0.1 *N* PhN₂Cl, and the pptd. dye taken up in 100 ml. 10% alc. NaOH and pptd. by 10% AcOH, giving 70% crude dye, which was extd. with hot EtOH and isolated as a dark powder, m. 108-70°. It was identical with the product obtained by heating 3.96 g. of the dye m. 195° (see part I) with 1.5 g. *m*-O₂NC₆H₄CHO in AcOH. only

A.S.M.-I.I.A. METALLURGICAL LITERATURE CLASSIFICATION

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The dye m. 195^o, thus established as being *p*-phenylazo-5-amino-1-naphthol, was, in addn., decimated by treatment of its diazonium deriv. in H₂SO₄ with abs. EtOH at 100^o, yielding a red solid, purified by pptn. from 10% NaOH by HCl, identified as *p*-phenylazo-1-naphthol, decomp. 205^o; in addn., replacement of the NH₂ by a OH group by the conventional diazonium reaction gave *p*-phenylazo-1,5-dihydroxynaphthalene, decomp. 210^o (the OH group in the conventional diazonium reaction gave *p*-phenylazo-1,5-dihydroxynaphthalene, decomp. 210^o). Similar to diazo soln. was heated with 10% H₂SO₄. Similar to placement of the NH₂ group in the dye m. 184^o (see part I), i.e. *p*-phenylazo-5-amino-1-naphthol, gave 2-phenylazo-1-naphthol, m. 138^o. The remaining dye isomer (see part I) was shown to be 2-phenylazo-3-amino-1-naphthol by the following: reduction of the dye with hydrosulfite gave 50% 1,2-diamino-3-naphthol, decomp. 206-7^o, which (7.63 g.) was heated to 50^o in 20% AcOH with 2 (8 g. phenanthrenequinone, giving 5-hydroxy-1,2-naphthophenanthraquinone, yellow-brown, m. 322-2.5^o (from PhCl). III. Some anomalous cases of reaction of aminonaphtholsulfonic acids with diazo compounds V. V. Perekahin. *Ibid.* 1784 (1930) (in Russian). Some aminonaphtholsulfonates, e.g., M acid (5-amino-1-naphthol-3-sulfonic acid) (I), 7-acid (7-amino-1-naphthol-3-sulfonic acid) (II), and 8-amino-1-naphthol-3-sulfonic acid (III), after coupling with 1,3,6-trisulfonic acid (P.P.-acid), do not couple mole of diazo deriv. in acid soln. although there are no apparent steric factors to explain such a behavior. It may be explained by resonance phenomena and by formation of a H bond between the sulfo group and a *peri*-H atom. By summation of the resonant structures formed individually by each substituent it was possible to arrive at the following table by comparative activities of C atoms in aminonaphtholsulfonic acids and in their mono-*o*-aminazo derivs.: I: C activity in the parent substance, *o*-OH -1, *p*-OH 0; in monoazo deriv.: *o*-OH 0, *p*-OH +1; II: -1, 0, 0, +1; the formation observed. II: -1, 0, 0, +1; 8-Amino-1-naphthol-2-sulfonic

acid: -2, -1, -2, -1; disazo dyes form. 8-Amino-1-naphthol 3,6-sulfonic acid: -1, 0, -1, 0; disazo dyes form. Prognostic extension of such summation gave the following theoretical results: 1, 5-amino-1-naphthol-2-sulfonic acid, and 5-amino-1-naphthol-8-sulfonic acid should not form disazo dyes, while 5-amino-1-naphthol-2-sulfonic acid and 5-amino-1-naphthol-4-sulfonic acid should readily couple twice; similarly, 7-amino-1-naphthol-4-sulfonic acid and 3-sulfonic acid and 8-amino-2-naphthol-6-sulfonic acid and 8-amino-2-naphthol-4-sulfonic acid should not give disazo dyes, and 6-amino-2-naphthol-4-sulfonic acid and 8-amino-2-naphthol-6-sulfonic acid should form disazo dyes. The mechanism of action of the substituents was assumed to be the normal, differing only in the algebraic sign and only the normal valence bonds were assumed to be operative. Similar to 1,5- and 2,8-naphtholsulfonic acids the distance between the O of SO₃H and the adjacent ring H bond exceed 1.7 Å. (*peri*-position), the formation of a H bond is likely. In order to investigate the possibilities named above, the following expts. showed that the cause for failure of 2nd coupling by I does not lie in the specific orientation of the substituents but in the presence of the SO₃H group. (No data on products or exptl. data are given.) 2-Phenylazo-5-amino-1-naphthol with Ph₂N₂Cl in alk. soln. (pH 10.5-11.5) gave 2,6-bis(phenylazo)-5-amino-1-naphthol, while an identical product, 2,4-bis(phenylazo)-5-amino-1-naphthol, was produced by similar treatment of either 2-phenylazo-5-amino-1-naphthol or 4-phenylazo-5-amino-1-naphthol; the last 2 substances did not couple with PhN₂Cl in acid medium. From Ph₂N₂Cl at pH 4.5-5.5 with 5-amino-1-naphthol-3-, -4-, -8-, -2-, and -4-sulfonic acids the corresponding monoazoderivs. were obtained and characterized by spectrum analysis and potentiometric titrations (no data given); the resulting 4-phenylazo-5-amino-1-naphthol-3-sulfonic acid, 8-phenylazo-5-amino-1-naphthol-8-sulfonic acid, and 6-phenylazo-5-amino-1-naphthol-8-sulfonic acid with PhN₂Cl in alk. soln. failed to yield any disazo products; similar failure took place with the isomers in which the monoazo

(cont)

PROCESSES AND PROPERTIES

coupling was governed by the OH group. 6-Phenylazo-5-amino-1-naphthol-2-sulfonic acid and 6-phenylazo-5-amino-1-naphthol-4-sulfonic acid, however, gave diazo derivs. which showed a displacement of the absorption max. toward longer wave lengths in comparison with the initial materials. Since the pH of the medium does not affect the location of the absorption max. in o-hydroxy azo dyes and since the OH group in them can be titrated with difficulty, the azoid structure (Auwers, C.A. 20, 1219) appears to be disproved for these compds. The "quinone-hydrazone" structure is also disproved by the absence of characteristic —NH— infrared absorption. Thus, the H of the OH is attached neither to O nor to N, and the OH group H bonds to —N=N—. This is confirmed by the higher pH titration requirement for o-OH-azo dyes than for the corresponding p-HO or o-amino derivs. IV. Nitrosation of 5-acetamido-1-naphthol. L. S. Efros, A. B. Poral-Koshits, and B. A. Poral-Koshits (Leningrad Technol. Inst.). *Ibid.* 1801-6(1947) (in Russian). —Nitrosation of 5,1-AcNHCO₂H₂OH gives 2-nitroso-5-acetamido-1-naphthol (I), whose structure was confirmed by formation of 1-acetamido-5,6-naphthophenazine (II). To 10 g. 5,1-H₂NC₂H₄OH suspended in 100 ml. H₂O was added 25 ml. Ac₂O and the mixt. was warmed until soln. occurred; on cooling, 8.5-9 g. 5-acetamido-1-naphthol, m. 174-5° (from water), sepd. This (4 g.) in 50 ml. cold 70% AcOH with 1.4 g. NaNO₂ in 6 ml. H₂O, gave 4 g. I, m. indefinitely with darkening at 200°, sol. in alkalis, pptd. by acids. I dyes wool in aq. suspension to a brown shade, with Fe mordant to grass-green shades. I (1 g.) in 5 ml. EtOH suspension was warmed with 1.5 ml. PhNHNH₂; after subsidence of the reaction and cooling, there was obtained 0.9 g. 2-amino-5-acetamido-1-naphthol (III), sandy plates, m. indefinitely, darkening at 142°, decomp. 185-7°, sol. in aq. acids and alkalis, and capable of being diazotized and coupled. I (2 g.) suspended in 75 ml. AcOH was warmed with 2.5 ml. PhNHNH₂; after subsidence, the mixt. was dil. with H₂O, giving 1 g. 5,6'-diacetamido-1,1'-dihydroxy-2,2'-azobenzophenone, red needles, m. 248-50° (some decompn. from alc.), which is almost insol. in aq. alkalis. I (1.5 g.) and 1.5 g. o-(H₂N)₂C₆H₄ were warmed in 4.5 ml. AcOH; after subsidence, the product was ground with EtOH, followed by cold 10% NaOH and washing with H₂O, giving 1.5 g. II, m. 311-13° (from AcOH). The same product was obtained by treating a cold suspension of 0.5 g. III in 10 ml. 70% AcOH with 0.25 g. Cr oxide in 5 ml. AcOH, dilg. with 100 cc. H₂O, extg. with CHCl₃, evapg. the ext., and warming the residue with 0.2 g. o-(H₂N)₂C₆H₄ in 3 ml. AcOH. III (1 g.) and 20 ml. H₂O heated 15 min. with 10 ml. AcOH gave on cooling 0.5 g. 2,5-diacetamido-1-naphthol, m. 215° (from dil. AcOH). This (5 g.), treated in 100 ml. cold AcOH with 1.5 g. NaNO₂ in 5 ml. H₂O and the ppt. recrystd. from AcOH, gave 0.8 g. red needles, m. 208°, while diln. of the mother liquor gave brown-yellow needles, m. 302-4°; both products are insol. in alkali; their structure is unknown.

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D.H.

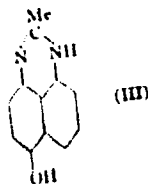
AS 50-514 METALLURGICAL LITERATURE CLASSIFICATION

1300 1310 1320

1300	1310	1320	1330	1340	1350	1360	1370	1380	1390	1400	1410	1420	1430	1440	1450	1460	1470	1480	1490	1500

V. Coupling of 5-amino-1-naphthol with diazotized II acid. *Ibid.* 1807-15 (in Russian).—Coupling of 5,1-AcNHCl₂H₂O (I) and 5,1-H₂NCO₂H (II) with diazotized II acid at pH over 4 gives *p*-HO dyes, while at pH below 4 I does not couple. II gives a *p*-aminoazo dye, which hitherto has been considered to be an *o*-amino deriv. (Ger. Pat. 95,100), or an *o*-HO deriv. The structure was established by reduction, followed by oxidation to juglone. I (2 g.) in 10 ml. 10% KOH and 20 ml. H₂O was treated with the diazonium deriv. from 3.2 g. II acid in 30 ml. H₂O, and the blue soln. acidified with AcOH, giving 4.7 g. of a metallic-looking powder, forming a violet soln. in H₂O, blue in dil. alkalis, red in stronger alkalis, absorption max. 520 mμ, dyes wool violet. Titration gave breaks at pH 9 and 4.7, the mol. wt. found being 550; the same product was obtained when coupling was run at pH 4. The product is probably 8-(8-acetamido-4-hydroxy-1-naphthylazo)-1-naphthol-3,6-disulfonic acid, since 5 g. of it in 50 ml. H₂O and a little NaOH, treated with Zn dust, followed by 10 ml. concd. HCl and boiled, gave the HCl salt of methyl-5-hydroxy-1,8-naph-

thimidazole (III), yellow needles without a definite m.p. When II was coupled, as above, in alkali and at pH 5, the products obtained were identical: metallic-looking powders, with color behavior similar to that of the Ac deriv. described above; the absorption max. was 510 mμ. The same product was obtained by deacetylation of the Ac compd. by hot 1:4 HCl. Reduction by Zn in HCl gave II acid and 4,5-diamino-1-naphthol-HCl (by addn. of HCl to the filtrate), which on heating with AcOH and 1:3 HCl gave the same imidazole deriv. as described above. If the coupling of II is conducted in AcOH-HCl or at pH 3, again identical products are obtained: a crystal powder, giving a violet soln. in water, blue in acids, and red in alkalis, with pptn. of some of the dye, heating in alkalis yields NH₃ (detected by color) and a soln., absorption max. 540 mμ (pH 3). Reduction of this product with Zn-HCl gave II acid and a soln. of 5,8-diamino-1-naphthol, which with 10% FeCl₃ *in situ* gave juglone, red-orange needles, m. 117° (from CHCl₃). This confirms the structure of the dye as 8-(8-hydroxy-3,6-disulfo-1-naphthylazo)-4-amino-1-naphthol. The *o*-aminoazo deriv. (1 g.) from PhN₂Cl and II (P.-K. and Perevalin, C. 1. 40, 1885) in 20 ml. H₂O and 10 ml. 10% KOH was reduced by Na hydrosulfite and acidified; after removal of S the filtered soln. was neutralized by NaOAc, treated with 0.8 g. phenanthrenequinone in 200 ml. H₂O with NaHSO₃, and boiled 5-10 min. to give 0.75 g. of the corresponding azine, yellow needles, m. 307-10° (from PhBr). Similar treatment of the above dyes failed to give the azine, thus confirming the absence of the *o*-amino structures. G. M. Kosolapoff



POZNANSKA, Hanna

Determination of basic phosphatase level in blood serum as a test for differentiation of parenchymal and mechanical jaundice. Polskie arch.med.wewn. 25 no.3a:587-590 '55.

1. II Klinika Chorob Wewnętrznych AM w Łodzi Kierownik: prof.dr med. J. Jakubowski Zakład Chemii Fizjologicznej AM w Łodzi. Kierownik: prof. dr med. B. Filipowicz.

(JAUNDICE

parenchymal & mechanical, differ.diag.,determ. of basic phosphatase level in blood serum)

(PHOSPHATASE, in blood

determ. of level in differ. diag. of parenchymal & mechanical jaundice)

(BLOOD

phosphatase level determ. in differ. diag. of parenchymal & mechanical jaundice)

POZNANSKA, I.

Parnas, J., Lorkiewicz, Zb., Poznanska, I., Nowak, B.: "Ze studiow nad wloskowcami rozycy" (Studies on Erysipelothrix Rhusiopathiae Suic), Medecyna Weterynaryjna, No. 7, p. 530, 1951.

POZNANSKA, I.

Parnas, J., Lorkiewicz, Zb., Poznanska, I.: "Badania nad uzjadliwieniem szczepu Stauba oraz hemaglutynacja z wloskowcami rozycey" (Studies on the reversion of virulence of an avirulent Staub's strain and hemagglutination with *Erysipelothrix suis*), Annales U.M.C.S., DD. IX.3.41, 1954.

PARNAS, J.; LORKIEWICZ, Z.; NOWAK, B.; POZHANSKA, I.

Studies on *Erysipelothrix rhusiopathiae*; studies on Stamb hem-
agglutinating strains; allergic phenomenon. Med. dosw. mikrob.,
Warsz. 4 no. 3:336-337 1952. (GLML 23:3)

1. Summary of work progress presented at 11th Congress of Polish
Microbiologists held in Krakow May 1951. 2. Lublin.

L 34451-66 T JK

(A)

SOURCE CODE: PO/0071/65/000/009/0546/0548

ACC NR: AP6026216

AUTHOR: Dziekonski, Jozef--Dzekon'ski, Y. (Doctor; Bydgoszcz); Drozdzyński, Witold--
Drozdzyński, V.; Poznanska, W.--Poznan'ska, V.

ORG: Regional Institute of Veterinary Hygiene/headed by Veterinarian J. Borowiecki,
Bydgoszcz (Wojewodzkie Zaklad Higieny Weterynaryjnej)

TITLE: Pasteurella hemolytica epizootic in lambs

SOURCE: Medycyna weterynaryjna, no. 9, 1965, 546-548

TOPIC TAGS: epidemiology, animal disease, commercial animal

ABSTRACT: Report of field outbreak on a large (764 sheep, 529 lambs) ovine farm; detailed description of clinical, biochemical and other laboratory observations; several mice-pathogenic strains of Pasteurella hemolytica were isolated from sick animals. [JPRS: 33,500]

SUB CODE: 06, 02 / SUBM DATE: none / ORIG REF: 002 / OTH REF: 002

Card 1/1

0976 1964

ACC NR: AP6021775

SOURCE CODE: UR/0413/66/000/012/0035/0035

INVENTOR: Adamovich, A. I.; Poznanskaya, E. M.; Fel'dman, R. M.; Sarenko, A. S.;
Mikhaylova, N. P.; Tsirlina, S. S.

ORG: None

TITLE: A method for producing diethylaminoethyl ester of diphenylacetic acid (base
of adiphenine). Class 12, No. 182715

SOURCE: Izobreteniya, promyshlennyye obraztzy, tovarnyye znaki, no. 12, 1966, 35

TOPIC TAGS: drug, ester

ABSTRACT: This Author's Certificate introduces a method for producing diethylamino-
ethyl ester of diphenylacetic acid (base of adiphenine). The technological process
is simplified by interacting diethylaminoethyl chloride in an aqueous solution with
an alkali metal salt of diphenylacetic acid.

SUB CODE: 07, 11/ SUBM DATE: 15Jul64

Card 1/1

UDC; 66,095.132;615.717

POZNANSKAYA, L.

Hans Christian Andersen. Nauka i zhizn' 22 no.4:57-58
Ap '55. (MLBA 8:6)

(Andersen, Hans Christian, 1805-1875)

POZNANSKAYA, L.

Henrik Ibsen; on the 50th anniversary of his death. Nauka i zhizn'
23 no.5:58-59 '56. (MLRA 9:8)
(Ibsen, Henrik, 1828-1906)

VALUYKO, G.G.; GODIN, K.G.; POZNANSKAYA, M.N.

Systems of the thermal processing of grapes. Trudy VNIIViV

"Magarach" 13:44-56 '64.

(MIRA 17:12)

POZNANSKAYA, N.

"The Selective Ionic Permeability of the Human Skin," Zhur.Fiz., Vol.28,
No. 4, pp 323-29, 1940

Dept. of Biological Physico-Chemistry (Head: Prof. D.L.Rubenstein), VIEM

CA

IIA

IONIC PERMEABILITY AND PROPERTIES OF THE SKIN

Ionic permeability of the human skin II. Topographic peculiarities of the electric resistance and ionic permeability of the skin N. B. Ponomareva, *Bull. biol. med. exp. 1. R. S. S. G.* 194 8(1968), *Chem. Zentr.* 1940, II, 1170-1; cf. *C. A.* 34, 5101; 36, 788. - A number of people had various regions of their skin tested for elec. resistance to potentials 100 mv. 8 v. The same places were tested for ionic permeability with AlCl₃, NaCl, Na acetate and Na salicylate solns. No correlation was found between the two phenomena. M. Hosh

AS 4.51.4 METALLURGICAL LITERATURE CLASSIFICATION

8204 577-02174

147040 74

147040 74

147040 74

CP

11A

PROCESSES AND PROPERTIES INDEX

The ionic permeability of human skin. I. N. S. Poznanskaya. *Bull. Acad. Sci. Div. Chem. Sci. USSR* 1954(10:17) (in English). The cation coeff. of skin permeability, K , which is the ratio J_e/J_i , where J_e denotes the permeability of the skin to the cations of the external soln. and the anions of the internal soln. (cell and tissue fluids) and J_i denotes the permeability to external anions and internal cations, was detd. for $N NaCl$, KCl , $LiCl$, $CaCl_2$, $MgCl_2$, and $AlCl_3$. In undamaged human skin the internal soln. remains unchanged so J_e and J_i represent the skin permeability to external cations and external anions (Cl^-), resp. K shows the greatest permeability, Na and Li give approx. similar values slightly lower than K , Ca permeability in most cases is of the same order as Na , but in 3 cases was found higher than Na , Mg shows slightly less permeability than Na and Al shows the least permeability. K increases the permeability to Cl^- considerably, Li and Al increase it slightly, while Ca and Mg have a strong inhibiting action on Cl^- permeability. Lowering of the pH of $AlCl_3$ solns. in $NaCl$ from 5 to 3 with glycine + HCl buffer causes little increase in Al permeability. In expts. on $N NaCl$, NaI , $NaNO_3$, Na_2SO_4 , and Na citrate, I^- , Cl^- and NO_3^- had the highest permeability, followed by sulfate and citrate ion. I^- increases cation permeability considerably, but the other anions have little effect on cation permeability.

S. A. Karjala

A 58-11.4 METALLURGICAL LITERATURE CLASSIFICATION

SUBJECT INDEX		AUTHOR INDEX	
GROUP	NO.	GROUP	NO.
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100	100	100	100

CA

117

The phenomena of physical electrotonus in solutions of different electrolytes. N. R. Ponomareva. *J. Physiol.* U.S.S.R. 20, 50 (1966) (English transl. *J. Physiol.* 34, 510). By expts. in which NaCl, KCl, AlCl₃, Na citrate, Na salicylate, and dil. HCl and H₂SO₄ were the electrolytes, it was shown that in cutaneous electrotonus, the polarization induced by a cation is the stronger and the anelectrotonus is the more pronounced the less readily the cation of the external soln. penetrates through the skin. The H⁺ ion penetrates the skin very quickly but causes an inversion of the anelectrotonus, possibly as a result of damage to the cutaneous membrane. Thus the role of cation is very important. Catelectrotonus is a result of penetration of anions through the skin. Therefore catelectrotonus is the weaker, the less penetrating is the anion of the external soln. Human skin has a selective permeability to anions. Hence the presence or absence of changes in the skin permeability, as result of cathodic polarization, is detd. largely by the choice of the anion of the external soln. It is probable that the cathodic inversion of current strength is due to alterations produced in the skin by the anions. Cl⁻ ion shows great penetrating power, while citrate and salicylate ions can barely pass through the skin. The skin permeability to H⁺ ion is evident from the fact that both catelectrotonus and anelectrotonus with acid solns. give much pain to the subject; other cations do not.

C. S. Shapiro

ASME-51A METALLURGICAL LITERATURE CLASSIFICATION

USSR / Human and Animal Morphology (Normal and Patho- 3-4
logical). Nervous System.

Abs Jour: Ref Zhur-Biol., No 17 1958, 79082.

Author : Magrupov, A. I., Semenova, Ye. N., Patrusheva,
T. M., Poznanskaya, Sh. L., Abdukhalinkov, F.,
Surkova, L. F.

Inst : Not given.

Title : Pathomorphology of the Internal Organs During
Toxic Encephalitis.

Orig Pub: Sb. nauchn. tr. Samarkands k. med. in-ta, 1955,
10, 145-153.

Abstract: No abstract.

Card 1/1

POZNANSKAYA, V. S.

Ca

21

Loke from anthracite, etc. V. S. Poznanskaya. Russ. 58,266, Oct. 31, 1940. Anthracite or other lean coal is powdered and mixed with finely divided pitch, asphalt or similar bituminous substance m. 65° or higher, and the mixt. is coked.

ASB-SLA METALLURGICAL LITERATURE CLASSIFICATION

GROUP	SECTION	SUBSECTION	CLASSIFICATION
A	1	1	
B	2	2	
C	3	3	
D	4	4	
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POZNANSKAYA, V.S.

USSR:

✓ Brown coal as coking material. I. F. Pakhalok and V. S. Poznanskaya. *Ugol'* 30, No. 3, 32-5 (1955).—The recommended process of coking brown coal involves drying to 10% moisture, crushing to 0.5-1.0 mm. size, briquetting at 1000-2000 kg./sq. cm. pressure into 50-g. briquets, slowly raising the temp. to 350° and coking at 1000-1100°. The coke strength was in line with that of coke produced at the Far Eastern by-product coking plants. W. M. Sternberg

Smelting ferro-silicon from roasted pyrite. S. I. Khitrik and A. E. Ponomarek. *Doklady Akad. Nauk SSSR*, 1934, No. 2, 3, 57-58. In a study of the application of roasted pyrite to the production of ferro-silicon, the authors had in view the effect of the starting material, roasted pyrite, iron filings or Fe ore, on caking of the charge and on energy consumption. The charge consisted of the Fe carrier, quartz and coke. The expts. were carried out in a 40-kw. elec. furnace. Ferro-Si alloys contg. the following amts. of Si were prepd.: 75, 65 and 45% from roasted pyrite; 75% from Fe filings; 75 and 60% from Fe ore. The roasted pyrite was in powder form. In the prepn. of 45% Si alloy from roasted pyrite (where the pyrite represented 25% of the charge), no difficulties were encountered and the charge descended regularly. In the charge of alloys with a higher Si content caking of the charge took place. In this respect pyrite was superior to Fe filings, but inferior to the ore. In general, the performance of the furnace depended not so much on the nature of the Fe carrier as on its amt. in the charge. As to energy consumption, it required, per ton of alloy, 46,000, 42,500 and 31,000 kw. hrs. for the 75, 65 and 45% alloys from pyrite, resp.; 43,000 and 38,000 kw. hrs. for the 75% alloy from filings and for the 60% alloy from ore, resp. A large amt. of pyrite was lost as flue dust, on account of its fineness. Its high S and C contents did not interfere with obtaining a high-grade ferro-Si.

S. L. Malovsky

1ST AND 2ND ORDERS PROCESSES AND PROPERTIES INDEX

B-I-5

BC

Smelting ferroalloys from roasted pyrites. S. I. KNITLIK and A. E. POMANSKI (Dokl. 1934, No. 2-3, 37-45).—Fe-Si alloys (45-75% Si) were prepared from powdered roasted pyrites. In alloys with > 45% Si caking of the charge took place. In this respect pyrites was superior to Fe silage, but inferior to Fe ore. The energy consumption in electrical smelting is discussed. Cu. Ans. (e)

ASB-SLA METALLURGICAL LITERATURE CLASSIFICATION

1ST AND 2ND ORDERS 1ST AND 2ND ORDERS

COMMON ELEMENTS COMMON ELEMENTS

COMMON VARIABLES INDEX

1ST AND 2ND ORDERS

1ST AND 2ND ORDERS

PROCESSES AND PROPERTIES INDEX

11 D

ca

Raciborski's nitrite and diazo reaction for plant cell membranes. **E. POZYŃSKI**. *Bull intern acad Polonaise 1929B*, 219-36 (in German). -- Raciborski's diazo reaction (*Bull intern acad Polonaise 1908*) for identification of aromatic incrustations in cell membranes is not specific for lignin, as Wiseligh (Die Zellmembran, Berlin, 1925) assumes, nor for suberin. The nitrite reaction is better and quicker. Micro sections are successively soaked in NaNO_2 (10%), H_2SO_4 (10%) and Na_2CO_3 (15% soln.) and washed. Coloration appears in the carbonate soln. Although this reaction does not indicate albuminous substances as was primarily expected, it is a characteristic means of identification of certain systematic units, since species of the same family with no exception gave the same shade and coloration. **J. WIENIĘLAK**

ASB-SLA METALLURGICAL LITERATURE CLASSIFICATION

SIGNATURE

SIGNATURE

17

CPX

Remarks on the methods of determination of alkaloids in the products and the raw material of Solanaceae. ADAM JURKOWSKI AND FRANCISZEK POJANAKI. *Wiadomości Farm.* 60, 133-5, 147-8(1935).—Three methods of detg. alkaloids are compared: (1) weighing a pptd. salt (picrolonate, picrate, etc.), (2) weighing the base and (3) treating the free base with an acid and titrating the excess acid. All the methods, although in use, are not easily carried out. Even Gadamer's modifications (*Deutsches Arzneibuch* 6th ed., 1926) are insufficient. The best method seems to be: Bata. of the substance with Et₂O, concn. of the ext., addn. of dil. HCl and H₂O, filtration and titration of the filtrate in the presence of methyl red.

J. WISNIEWSKI

ATM-31A METALLURGICAL LITERATURE CLASSIFICATION

GROUP	CLASS	SUBCLASS	SECTION	INDEX
1	2	3	4	5
6	7	8	9	10
11	12	13	14	15
16	17	18	19	20
21	22	23	24	25
26	27	28	29	30
31	32	33	34	35
36	37	38	39	40
41	42	43	44	45
46	47	48	49	50
51	52	53	54	55
56	57	58	59	60
61	62	63	64	65
66	67	68	69	70
71	72	73	74	75
76	77	78	79	80
81	82	83	84	85
86	87	88	89	90
91	92	93	94	95
96	97	98	99	100

JOZEF POZNANSKI

JOZEF POZNANSKI: an obituary. p. 319. Vol. 6, no. 12, Dec. 1956. TECHNIKA
I GOSPODARKA MORSKA. Gdansk Poland.

SOURCE: East European Accessions List (EEAL) Vol. 6, No. 4--April 1957

POZNANSKI, MARCELI, ed.

Kto miluje ksiegi; antologia tekstow o ksiazce.

(Wyd. 1. Warszawa), Poland Stowarzyszenie Bibliotekarzy Polskich (1968) 288 p.

Monthly List of East European Accessions, (EEAI) LC, Vol. 9, No 1, Jan. 1960
Uncl.

POWIAZANIE, P.

Remarks on a 1-level indicator for credit tools. p. 612.

PRACOWNIAKOWIE ZEM. (Stowarzyszenie Inżynierów i Techników Rolniczych
Polskich) Warszawa, Poland. Vol. 19, no. 19/20, Oct. 1956.

Monthly List of East European Acquisitions (CBI) 19, Vol. 9, no. 2, 1956.
Encl.

SUWALSKI, Ludomir (Warszawa); POZNANSKI, Tomasz (Warszawa)

Preliminary studies on the acceleration of the maturing process of concrete by high frequency currents. Archiw inz lad 7 no.3:403-414 '61.

POLAND

KORNIEWICZ, Adolf, KOTLIŃSKI, Jerzy, and POLIŃSKI, Wiesław, Experimental Research Office (Zakład Doświadczalny) of the Zootechnical Institute (Instytut Zootechniki) in Czołuch and the Department of Hog Breeding (Zakład Hodowli Trzody Chlewniej) of the WSR [Wyższa Szkoła Rolnicza, Higher School of Agriculture] in Wrocław (Director: Dr. Jerzy KOTLIŃSKI)

"Impasit, Mineral Mixture and Forest Soil as Media Preventing Anaemia in Sucking Piglets, Used in Indoor and Outdoor Systems of Management."

Warsaw-Lublin, Nadwojna weterynaryjna, Vol 14, No 11, Nov 62, pp 689-694.

Abstract: [Authors' English summary modified] Authors report materials, procedure, and tabulated results of their investigation, on the basis of which they conclude that piglets are particularly prone to anaemia in the absence of outdoor management and forest soil, that a supplementary supply of forest soil, mineral mixture, and Impasit are effective prevention in the order listed. One English, two Polish, and one Soviet references.

1/1

R-11-5

PROCESSES AND PROPERTIES INDEX

Cause of formation of "milky spots" in viscous silk. A. FUKUDA (Furuyal Chem., 1936, 19, 46-47).
—Loss of lustre after drying is due to evaporation of CO_2 bubbles in the fibres, with their replacement by air; the dimensions and no. of these bubbles diminish with increasing maturity of the viscose, and vary according to the conditions of manufacture of the latter. E. T.

METALLURGICAL LITERATURE CLASSIFICATION

1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100
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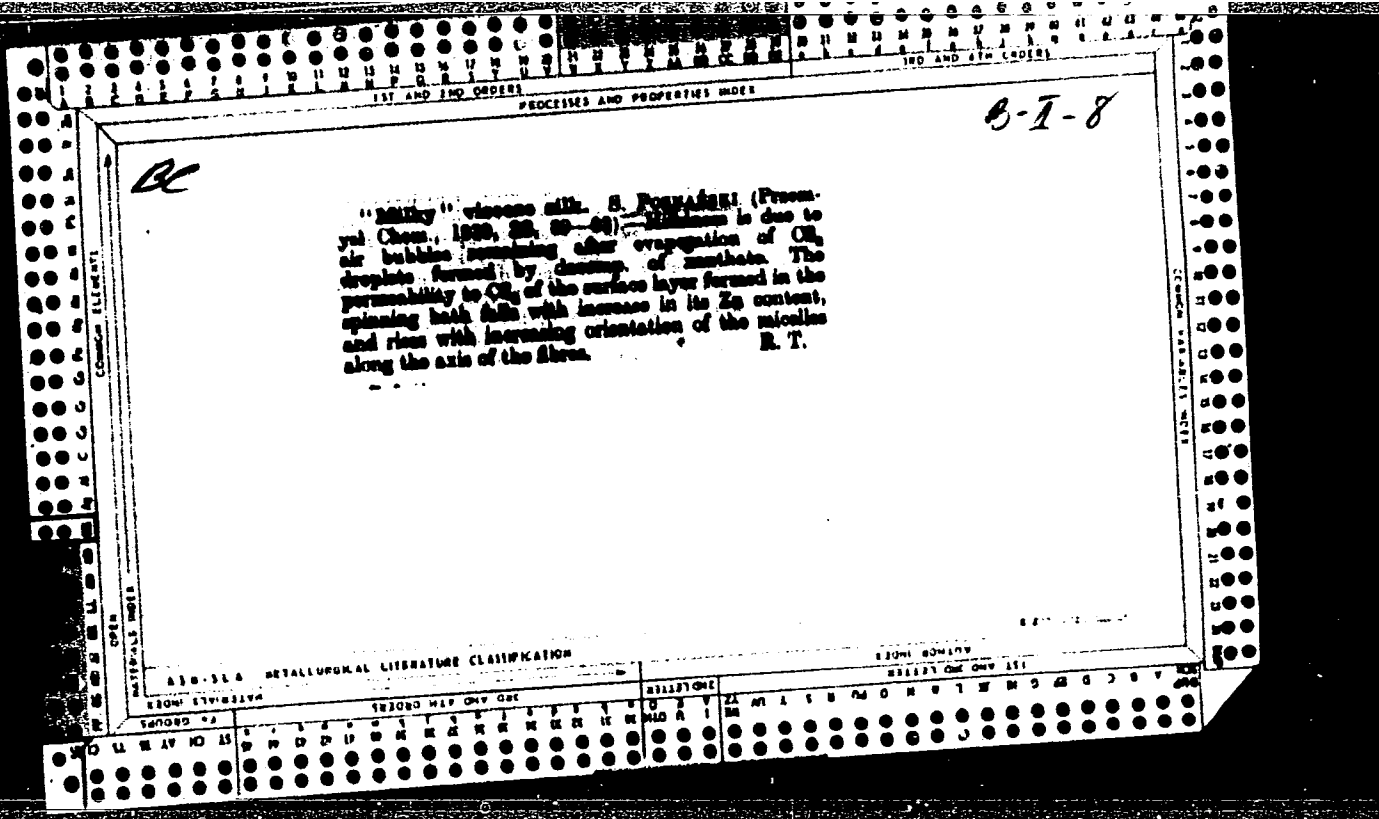
B-7-8

BC

Gravimetric determination of dihydrocarbonic acid residues of cellulose nanthate in viscose.
S. FERNANDEZ (Farmacia Chem., 1930, 23, 22-26).—
 Films of viscose (3-4 g.) are shaken with 80 c.c. of saturated aq. NaCl and 10 c.c. of 0.1N-I in KI. After 1 hr. 3-4 c.c. of 0.1N-Na₂S₂O₄ are added, to decolorize the films, and excess of Na₂S₂O₄ is titrated with I in KI. The C₂H₄ content as obtained agrees closely with that given by the gravimetric method, or by that of D'Ann and Jager (B., 1936, 587). Slightly higher results are given by titration of films of Zn cellulose nanthate, and considerably higher ones by the method of Bari and Dillencus (B., 1933, 222).
 R. T.

ASS-SLA METALLURGICAL LITERATURE CLASSIFICATION

1930 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100



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

BC *B-Z-8*

FUNCTIONS OF CELLULOSE NITRATE, AND THEIR TECHNOLOGICAL IMPORTANCE. S. FORTA (Frascati Chem., 1938, 22, 463-470).—In aq. ZnSO₄, part of the Na of viscose is substituted by Zn, to yield a mixed Na-Zn salt, which is less readily decomposed by acids than is the Na salt. Formation and decomp. of the Zn cellulose nitrate layer in spinning baths proceeds from the surface of the thread to its centre. The final product contains a certain amount of Zn combined directly with cellulose.
R. T.

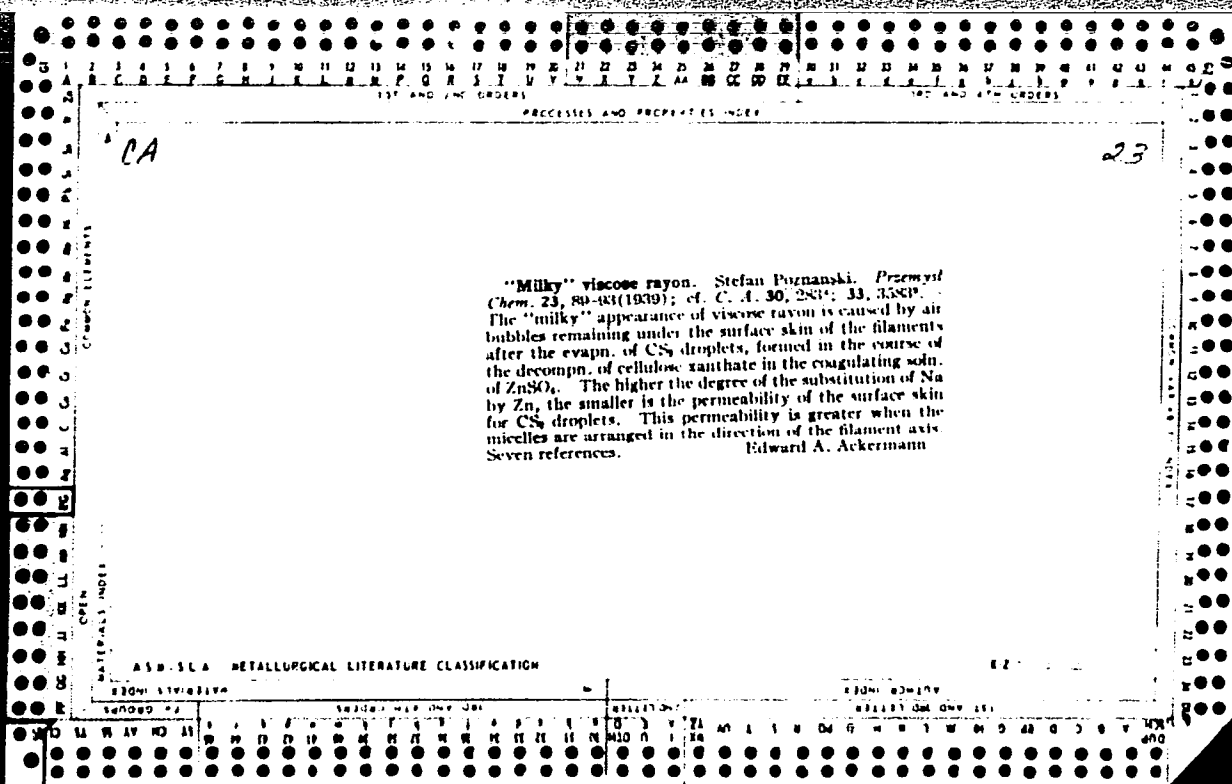
COMMON ELEMENTS
COMMON VARIABLES INDEX

ASB-31A METALLURGICAL LITERATURE CLASSIFICATION

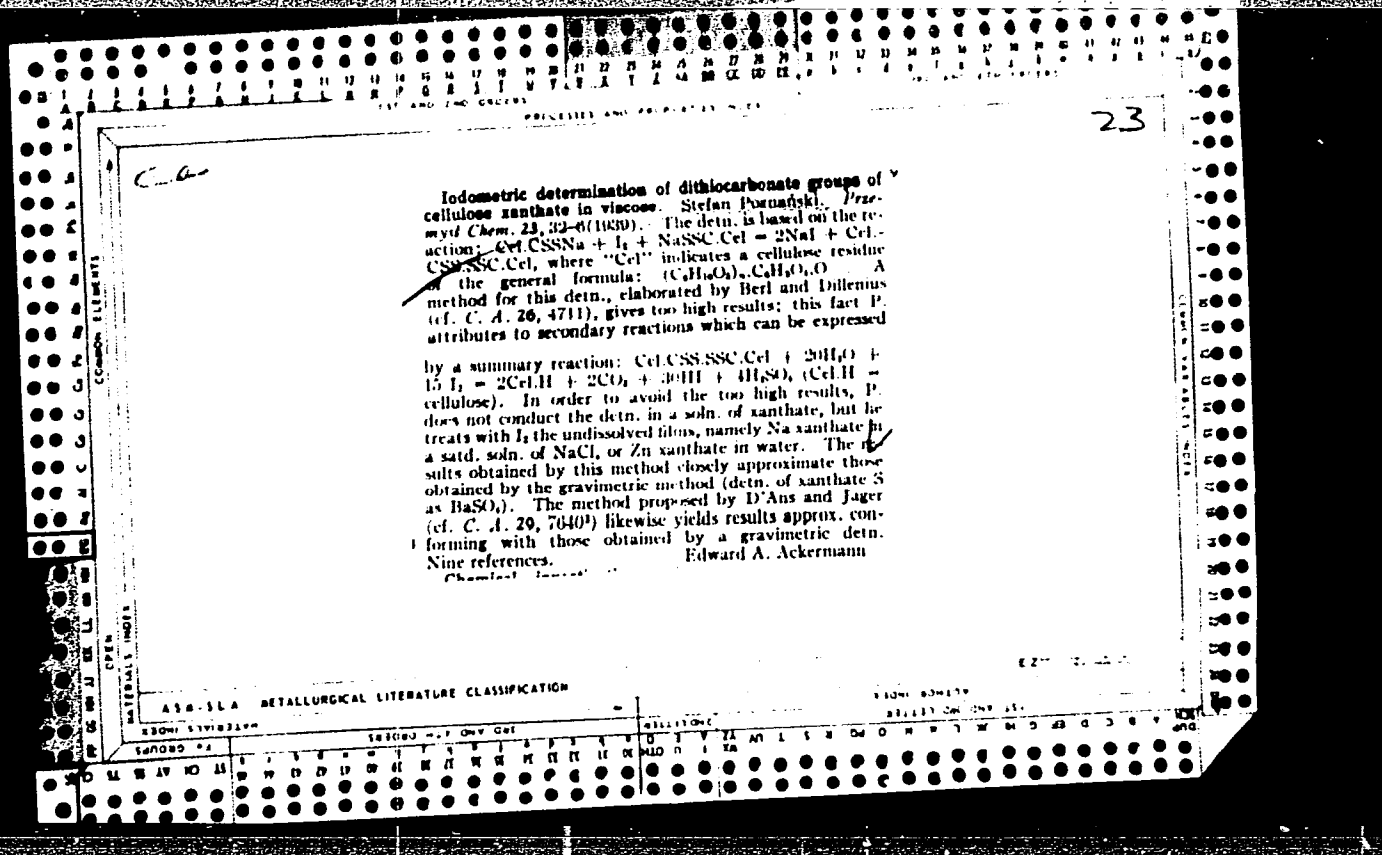
6-21-72

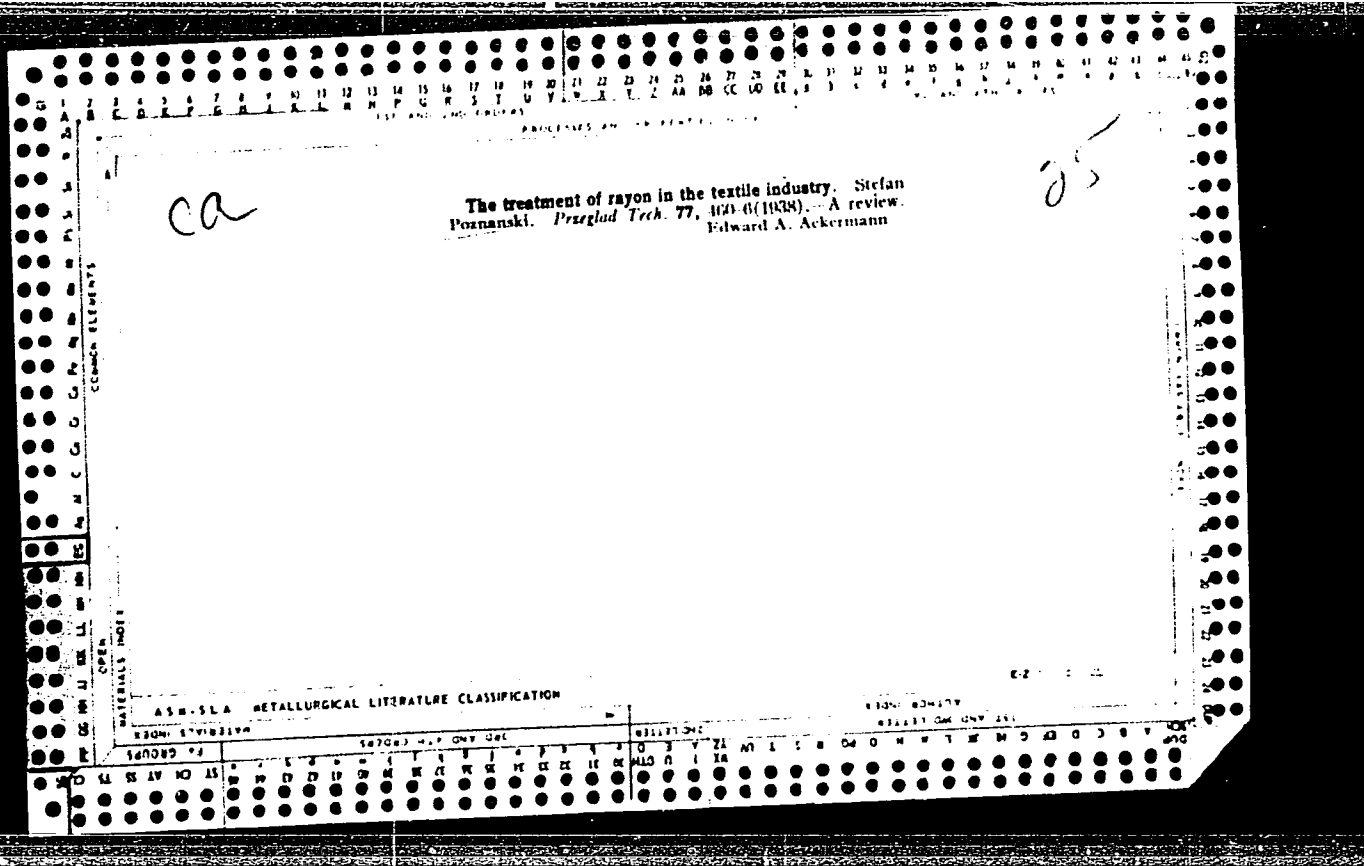
FROM SYNONYMS COLLECTION TO WHICH BELONGS

1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100



"Milky" viscose rayon. Stefan Poznanski. *Przemysl Chem.* 23, 80-83(1939); cf. *C. A.* 30, 2831; 33, 3589. The "milky" appearance of viscose rayon is caused by air bubbles remaining under the surface skin of the filaments after the evapn. of CS₂ droplets, formed in the course of the decompn. of cellulose xanthate in the coagulating soln. of ZnSO₄. The higher the degree of the substitution of Na by Zn, the smaller is the permeability of the surface skin for CS₂ droplets. This permeability is greater when the micelles are arranged in the direction of the filament axis. Seven references. Edward A. Ackermann





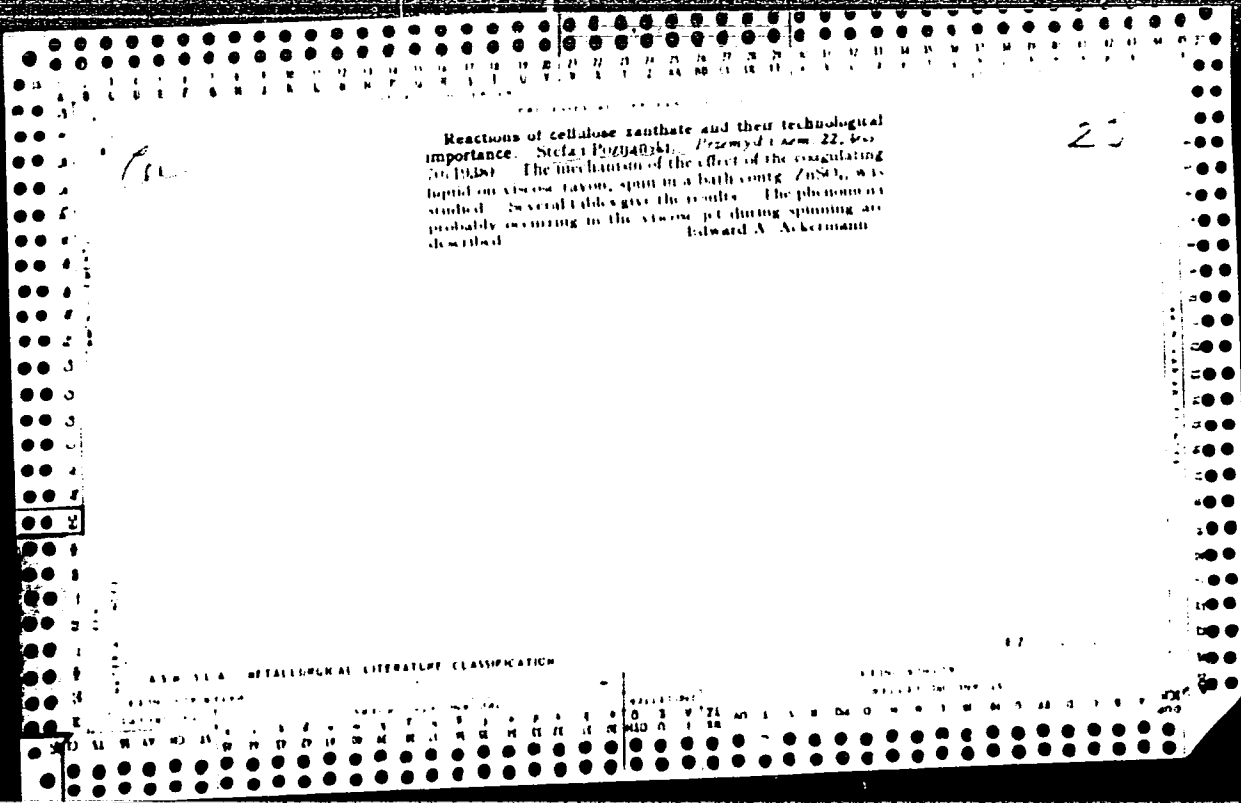
CA

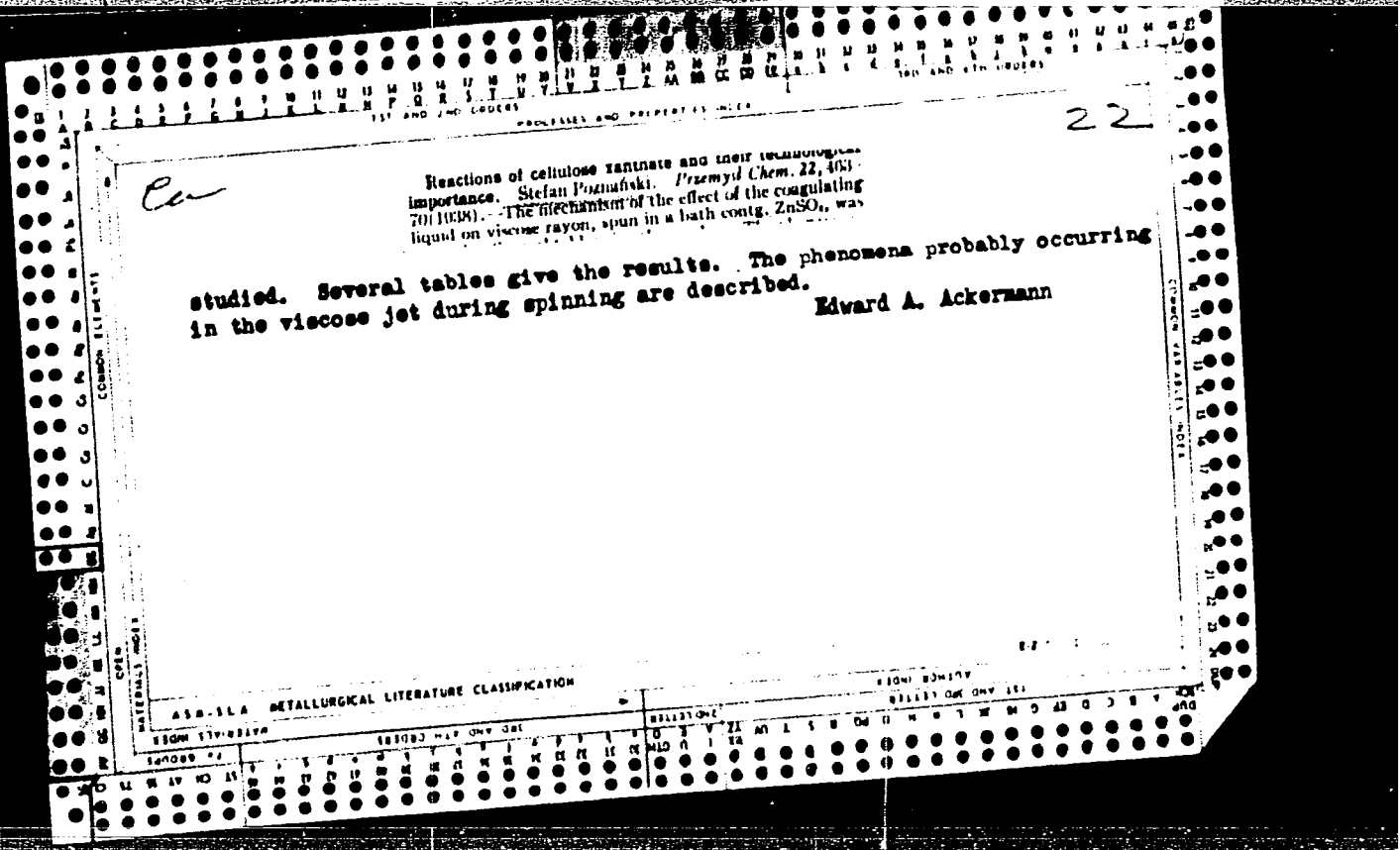
23

Cause of "milky" spots on viscose. Stefan Potapanaki, *Russia* 9, 179-83, 257, 259 (1931).—"Milky" spots in viscose are decreased by: (1) decreasing the amt. of CS₂ (even very slight differences can have a remarkable effect), (2) decreasing the viscosity of the viscose, (3) increasing the NaOH content of the viscose, (4) addn. of a small amt. of Na sulfonate to the viscose, (5) increasing the H₂SO₄ content of the coagulating bath, (6) decreasing the ZnSO₄ content of the coagulating bath, (7) lowering the titer of the elementary fibers, (8) decreasing the no. of elementary fibers in the threads. From the results of microscopic examn. of sections of fibers produced from viscose with very high CS₂ contents (50, 60 and 70%), P. considers that the voids under the surface skin which cause "milky" are not due to a difference in the structure of the skin and of the central portion of the fiber (contrary to Preston, C. A. 25, 4702-3), but to the retention of excess CS₂ within the fiber on coagulation, this CS₂ evapp. out through the surface on subsequent drying of the thread. Photomicrographs are given and discussed in support of this theory, and it is shown that the action of most of the factors enumerated above (except (6), the action of which remains as yet unexplained) agree with this hypothesis.

A. Papineau-Couture

AS B-S-L-A METALLURGICAL LITERATURE CLASSIFICATION





NIKIFOROV, I.; MAKAROV, A.; SMOLYAKOV, N.; SIPER, E.; MOGILA, V.; LARIN, M.;
FILIPPOV, K.; TOKMAKOV, V.; BARANOVSKIY, V.; CHETVERIKOV, K.;
POZNANSKIY, A.; SHUTOV, M.; ROZENFEL'D, L.; RUD', A.

Mechanization of waterproofing operations. Stroitel' 8 no.11:
15-20 N '62. (MIRA 16:1)
(Waterproofing--Equipment and supplies)

POZNANSKIY, A.S.

Relation between the pathodynamic structures of disease and the selective properties of drugs. Trudy Gos.nauch.-issl.inst.psikh. (MIRA 16:2)
35:36-43 '62.

1. Bashkirskiy meditsinskiy institut (dir. dotsent N.F. Vorob'yev)
kafedra psikhiiatrii (zav. kafedroy -- prof. A.S. Poznanskiy).
(Mescaline) (Schizophrenia)

POZNANSKIY, A. S. Doc Med Sci -- (diss) "Dynamics of schizophrenic syndroms in the process of psychopharmacological and other experimental clinical studies." Gor'kiy, 1959. 21 pp (Gor'kiy State Med Inst im S. M. Kirov), 200 copies. List of author's works pp 20-21 (18 titles) (KL, 49-59, 142)

POZNANSKIY, A. S. (Dotsent)

Kliniko-farmakodinamicheskiy metod issledovaniya patofiziologicheskikh mekhanizmov
pri lechenii shizofrenii, p. 293
V sb. Aktual'n. probl. nevropatol. i psikiatrii., Kuybyshev 1957.

POZNANSKIY, A.S.; AKOPYAN, P.O.

Our experience with the specialization of physicians in psychiatry. Zhur.nevr.1 psikh. 62 no.8:1273 Ag '62.

(MIRA 15:12)

1. Kafedra psikhiatrii (zav. - prof. A.S.Poznanskiy) Bashkirskogo meditsinskogo instituta i Bashkirskaya respublikanskaya psikhonevrologicheskaya bol'nitsa (glavnyy vrach P.O.Akopyan).
(PSYCHIATRISTS)

POZNANSKIY, A.Z. (Tallin)

Laboratory apparatus for determining the concentration of carbon monoxide
in the air. Gig. i san. 26 no. 6:65-68 Je '61. (MIRA 15:5)
(CARBON MONOXIDE) (AIR--ANALYSIS)

POZNANSKIY, A.S. [Poznans'kiy, A.S.]

Effect of external stimulations on the dynamics of perception disorders [with summary in English]. Fiziol.zhur. Ukr. 4 no.5: 672-678 S-0 '58 (MIRA 11:11)

1. Gor'kovskiy meditsinskiy institut im. S.M. K₁rova, kafedra psikhatrii.

(HALUCINATIONS AND ILIUSIONS)

POZDNYI, S. S.

21 00 POZDNYI, S. S. Novyye etapy razvitiya gosudarstvennoy ekonomiki i upravleniya. Vrachsk. zhurn., 1986, No. 3, str. 22-27.

CC: Laboratsiya Statisticheskoy Meditsiny, No. 29, Moskva, SSSR.

ПОКАСНИЙ, С. С.

№577

Тысячелетнему Семейному Обществу На Сыле I Задчи Органов Здравоохранения Третьей Сыле, 1949, №. 9 СТБ. 825-1

SC: LETCFIS NO. 38

POZNANSKIY S.S.
GABOVICH, R.D.; POZNANSKIY, S.S.

Typical plan for practical training in the field of hygiene in
therapeutic and pediatric departments of medical institutes.

Gig. 1 san. no.9:41-45 S '54.

(MLRA 7:10)

(HYGIENE, education,

Russia, in ther. & pediatric departments of med. schools)

(EDUCATION, MEDICAL,

hyg. practical training in pediatric & ther. departments
of med. schools)

GABOVICH, R.D.; POZNANSKIY, S.S.

Aid of a medical institute to sanitary-epidemiological stations.
Sov. zdrav. 13 no.5:9-14 S-0 '54. (MLRA 7:12)

1. Iz Kiyevskogo ordena Trudovogo Krasnogo Znameni meditsinskogo
instituta imeni akad. A.A.Bogomol'tsa (dir. dotsent I.P.Alekseyenko)
(COMMUNICABLE DISEASES, prevention and control,
in Russia)

POZMANS'KIY, S.S.; DUPLENKO, K.F.

Zinovii Petrovich Solov'ev. Medych.zhur.24 no.5:85-91 '54.
(BIOGRAPHIES, (MLRA 8:10)
Solov'ev, Zinovii P.)

BATKIS, Grigoriy Abramovich [Batkin, H.A.], prof.; POZHANSKIY, S.S.
[Poznans'kiy, S.S.], red.; BOYZO, Ye.P. [Boiko, I.R.],
red.

[First steps in the organization of Soviet public health in
the Ukraine, 1918-1922] Pershi kroky budivnytstva radians'koi
okhorony zdorov'ia na Ukraini (1918-1922 rr.) Kyiv, Derzh-
medvydav Ukra, 1962. 45 p. (MLA 18:1)

GABOVICH, Rafail Davidovich, prof.; FOZNANSKIY, Semen Semonovich ,
dots.; SHAKHBAZYAN, Gayk Khachaturovich, prof.; PETROVSKIY,
K.S., red.

[Manual of hygiene] Uchebnik gigieny. Meditsina, 1964.
471 p. (MIRA 17:11)

1. Chlen-korrespondent AMN SSSR (for Shakhbazyan).

KALYUZHNYI, D.N., prof., red.; POZNANSKIY, S.S., dots., red.;
PETROV, Yu.L., red.; ZAPOL'SKAYA, L.A., tekhn. red.

[Problems in protecting the health of children and
adolescents] Voprosy okhrany zdorov'ia detei i pod-
rostkov; materialy. Pod red. D.N. Kalyuzhnogo i S.S.
Poznanskogo. Kiev, Gosmedizdat USSR, 1963. 219 p.
(MIRA 16:11)

1. Nauchnaya konferentsiya po respublikanskoy pro-
bleme "Okhrana zdorov'ya detey i podrostkov". 2. Chlen-
korrespondent AMN SSSR (for Kalyuzhnyy).
(PUBLIC HEALTH)

BARANNIK, P.I., red.; BARCHENKO, I.P., red.; GABOVICH, R.D., red.;
KAGAN, S.S., red.; KALYUZHNYI, D.N., red.; KRIVOGLAZ, B.A.,
red.; POZNAŃSKIY, S.S., red.; SUPONITSKIY, M.Ya., red.;
TRUBENBERG, I.M., red.; SHAKHBAZYAN, G.Kh., red.; SHMAL',
D.D., red.; OSETROV, V.I., red.; CHUCHUPAK, V.D., tekhn.red.

[Problems of general and specialized hygiene] Voprosy obshchei
i chastnoi gigieny. Kiev, Gosmedizdat USSR, 1963. 308 p.

(MIRA 16:10)

1. Ukraine. Ministerstvo zdravookhraneniia.
(PUBLIC HEALTH)

BRATUS', V.D., dots., red.; BARCHENKO, I.P., prof., zam. red.;
VERZHIKOVSKAYA, N.V., dots., red.; GROMASHEVSKIY, I.V.,
prof., red.; SHAKHBAZYAN, G.Kh., prof., red.; BARANIK,
P.I., prof., red.; SHMAL', D.D., dots., red.; POZNIANSKIY,
S.S., dots., red.; KALYUZHNIYY, D.N., red.; CHUCHUFAK, V.D.,
tekhn. red.

[Hygienic norms and the sanitation of the external environ-
ment]Gigienicheskie normativy i ozdorovlenie vneshnei sredy;
sbornik nauchnykh rabot. Kiev, Gosmodizdat USSR, 1961. 268 p.
(MIRA 15:11)

1. Kiev, Medychnyi instytut. 2. Deystvitel'nyy chlen Akademii meditsinskikh nauk SSSR (for Gromshevskiy). 3. Chlen-korrespondent Akademii meditsinskikh nauk SSSR (for Shakhbazyan).
4. Direktor Kiyevskogo meditsinskogo instituta (for Bratus').
5. Kafedra gigiyeny pitaniya Kiyevskogo meditsinskogo instituta im. A.A.Bogomol'tsa (for Barchenko). 6. Kafedra obshchey gigiyeny Kiyevskogo meditsinskogo instituta Kiyevskogo meditsinskogo instituta im. A.A.Bogomol'tsa (for Verzhikovskaya, Shmal').

(PUBLIC HEALTH)

BARANIK, P.I., prof.; POZNANSKIY, S.S., dotsent (Kiyev)

Sanitary and epidemiological station as a base for training physicians
of a medical institute. Sov. zdrav. 21 no.1:25-27 '62. (MIRA 15:2)

1. Iz Kiyevskogo meditsinskogo instituta.
(PUBLIC HEALTH__STUDY AND TEACHING)

KALYUZHNYIY, D.K., prof., otv.red.; GORODETSKIY, A.S., kand.med.nauk, red.;
IZDEBSKIY, A.M., kand.med.nauk, red.; KVITNITSKAYA, N.N., kand.
med.nauk, red.; KRYZHANOVSKAYA, V.V., kand.med.nauk, red.; MARTY-
NYUK, V.Z., prof., red.; PETROV, Yu.L., kand.med.nauk, red.;
POZNAVSKIY, S.S., kand.med.nauk, red.; STOVBUN, A.T., kand.med.
nauk, red.; SHMAL', D.D., kand.med.nauk, red.; POTOTSKAYA, L.A.,
tekhred.

[Hygienic study and improvement of the environment] Gigieniche-
skoe izuchenie i ozdorovlenie vneshnei sredy. Kiev, Gos.med.izd-vo
USSR, 1959. 331 p. (MIRA 13:4)

1. Ukrainskiy nauchno-issledovatel'skiy institut kommunal'noy gi-
giyeny. 2. Predsedatel' Problemnoy komissii Ministerstva zdravo-
okhraneniya USSR (for Kalyuzhnyy).
(PUBLIC HEALTH)

POZHANSKIY, V., starshiy nauchnyy sotrudnik

Dawn of Soviet inventions. Izobr.i rats. no.6:1-3 Je '59.
(MIRA 12:9)

1. Tsentral'nyy gosudarstvennyy arkhiv Kazakhskoy SSR.
(Efficiency, Industrial)

MOSKOVSKIY, A.B., otv. red.; BOKUCHAYEV, G.A., red.; POZNANSKIY,
V.S., red.; TARASOVA, N.V., red.

[Siberia during the period of the building of socialism
and transition to communism] Sibir' v period stroitel'stva
sotsializma i perekhoda k kommunizmu. Novosibirsk, Red.-
izd. otdel Sibirskogo otd-nia AN SSSR. No.3. 1964. 106 p.
(MIRA 18:9)

ISAKOV, A.I., kand.tekhn.nauk; POZNAYEV, A.P.; KORZHUK, G.K.

Quality of particle board. Bum. i der. prom. no.2:32-36 Ap-Je '63.
(MIRA 17:2)

1. Ukrainskiy nauchno-issledovatel'skiy institut mekhanicheskoy obrabotki drevesiny.

POZNAYEV, Aleksandr Petrovich; SOKOLOV, P.V., red.

[Measurement of wood moisture; electrical methods and apparatus] Izmerenie vlazhnosti drevesiny; elektricheskie metody i pribory. Moskva, Lesnaia promyshlennost', 1965. 141 p.
(MIRA 18:3)

ISAKOV, A. I., kand. tekhn. nauk; POZNAYEV, A. P., inzh.; KORZHUK,
G. K., inzh.

Automatic device for controlling and sorting panel parts and
slabs. Der. prom. 12 no.2:7-10 F '63.

(MIRA 16:4)

1. Ukrainskiy nauchno-issledovatel'skiy institut mekhanicheskoy
obrabotki drevesiny.

(Hardboard) (Sorting devices)
(Automatic control)

POZNAYEV, A.P.; KOH HUK, G.K.

Controlling the moisture of chip pulp during the production of
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SOURCE: Ref. zh. Fizika, Abs. 6B4

AUTHOR: Pozner, A. R.

TITLE: Role of methodological principles and construction of physical theories 21

CITED SOURCE: Sb. Metodol. probl. sovrem. nauki. M., Mosk. un-t. 1964, 54-83

TOPIC TAGS: general physical theory, mathematical theory, physical law, logical symbol

TRANSLATION: In the author's opinion, it is possible to separate in modern physical theories two aspects: the conceptual apparatus (physical concepts, laws, and principles) and formal calculations (mathematical equations, logical symbols and rules). It is noted that at different stages of development of physical theories, an appreciable role is played not only by the mathematical apparatus but also by the methodological principles of different degrees of generality. A. Pozner.

SUB CODE: GP, MA

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