

PROCESSES AND PROPERTIES

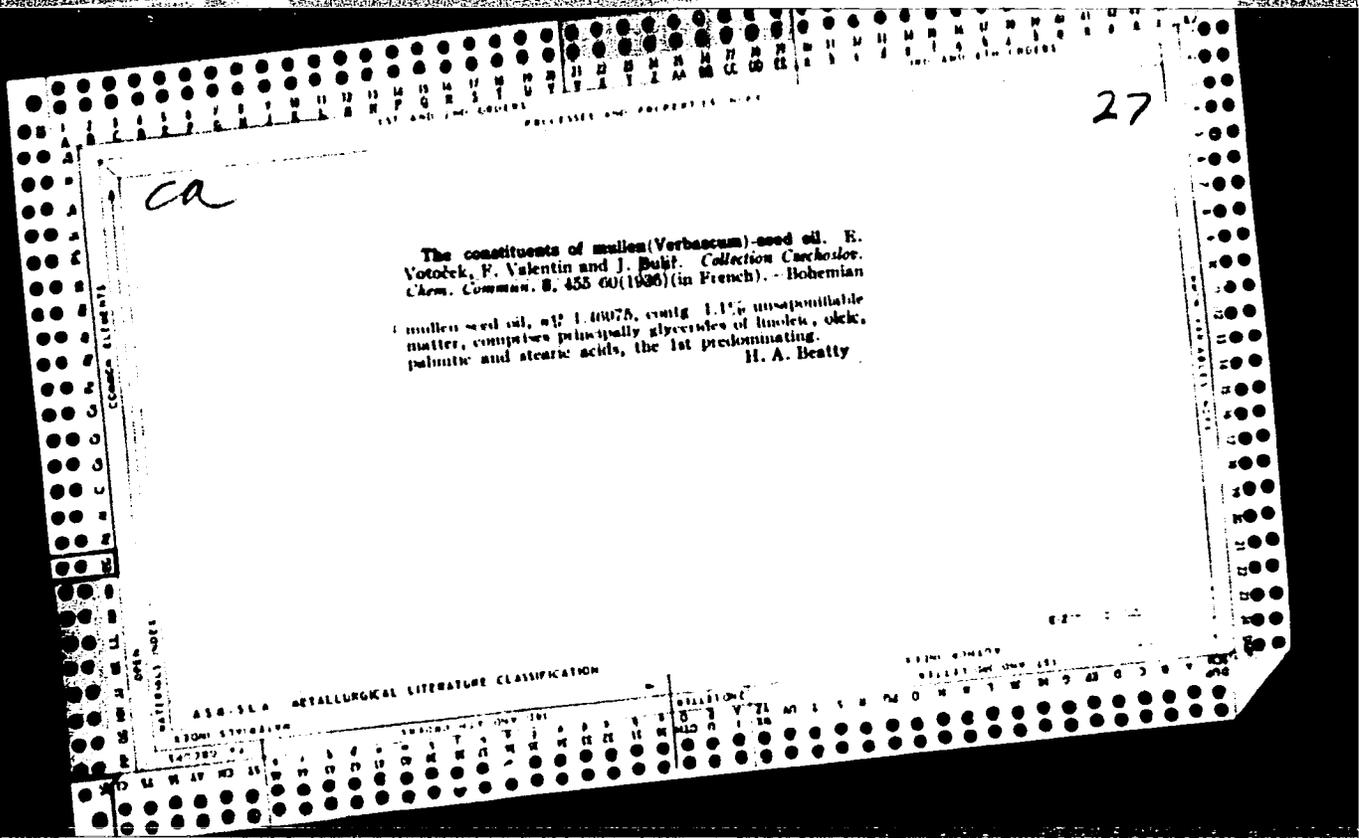
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ca

The rotatory power of certain sugar hydrazones with relation to the stereochemical structure of the α -carbon. R. VOTOUK, R. VALENTIN AND O. LEMINGER. *Collection Czechoslov. Chem. Comm.* 3, 250-64 (1931). The authors prepd. hydrazones of various sugars with 1,1-benzylphenyl, *p*-chlorobenzylphenyl and dibenzylhydrazines, resp. They conclude that the benzyl group of these hydrazones has such a marked effect on the rotation of the α -C atom that the configuration of the rest of the sugar mol. has no effect on the rotation of the hydrazone. Of the hydrazones studied only 2 have not been previously prepd.; the *benzylphenylhydrazones* of *D*-arabinose, m. 173°, and of *D*-thiambohexose, m. 183-84°. *p*-Chlorobenzylphenylhydrazine, m. 41°, was prepd from *p*-ClC₆H₄CH₂Cl and PhNHNH₂. J. L. WILKS

ASB-SLA METALLURGICAL LITERATURE CLASSIFICATION

12



1ST AND 2ND LAYERS
3RD AND 4TH LAYERS

PROCESSES AND PROPERTIES INDEX

7

BC

New form of two enantiomorphous rhamsites.
 F. VALENTIN (Ost. Chem. Gesell. Comm., 1900: 2, 600-602).—Reduction of rhamsite with 2% sodium amalgam and crystallization of the product from water. *d*-rhamsite: m.p. 80°, [α]_D +0.45° in water; *l*-rhamsite: m.p. 80°, [α]_D -0.45° in water; both soluble in water in a vacuum to 1.5% in water; *d*-rhamsite: m.p. 117°, *l*-rhamsite: m.p. 117°. *d*-rhamsite trihydrate has m. p. 117° in water, whilst *l*-rhamsite, prepared by crystallizing equal quantities of the enantiomorphs from water, has m. p. 117° and is anhydrous. Crystallographic data for the *d*- and *l*-rhamsite trihydrates are given; the crystals exhibit enantiomorphous facets. O. W. SMORAN.

ASS.-SLA METALLURGICAL LITERATURE CLASSIFICATION

ROOM SYMBOL

ALIST ONE ONE 111

COMPOUND ELEMENTS

NATURAL SYMBOLS

10000 11000 12000 13000 14000 15000 16000 17000 18000 19000 20000 21000 22000 23000 24000 25000 26000 27000 28000 29000 30000 31000 32000 33000 34000 35000 36000 37000 38000 39000 40000 41000 42000 43000 44000 45000 46000 47000 48000 49000 50000 51000 52000 53000 54000 55000 56000 57000 58000 59000 60000 61000 62000 63000 64000 65000 66000 67000 68000 69000 70000 71000 72000 73000 74000 75000 76000 77000 78000 79000 80000 81000 82000 83000 84000 85000 86000 87000 88000 89000 90000 91000 92000 93000 94000 95000 96000 97000 98000 99000

BC
 [3:6]-Anhydrogalactose. F. VALNTIN (Coll. Chem. (Chem. Comm., 1232, 4, 364-378).— α -Methylgalactoside (modified prep.), when treated in C_6H_5N with CPh_3Cl and then with Ac_2O , gives 2:3:4-

triacetyl-6-triphenylmethyl- α -methylgalactoside, m.p. 179-181°, $[\alpha]_D^{20} +86^\circ$ in C_6H_6 , which, when treated with PHr , in ethylene dichloride and then with NH_3-MeOH , affords α -methylgalactosidyl 6-bromide, decomp. 163°, $[\alpha]_D^{20} +157^\circ$ in H_2O . This with $Be(OH)_2$ yields 3:6-anhydro- α -methylgalactoside, m.p. 141-143°, $[\alpha]_D^{20} +82.4^\circ$ in H_2O , hydrolysed by 1% H_2SO_4 to 3:6-anhydrogalactose (I), amorphous, $[\alpha]_D^{20}$



in $H_2O +37.6^\circ$, changing to $+27.2^\circ$ (phenylglucosazone, m.p. 215° (decomp.), $[\alpha]_D^{20} +48.3^\circ$ in $MeOH$). The possibility that (I) contains an ethylenic linking is excluded by its stability to halogens, and the constitution of the 3:6-ring follows from stereochemical reasons and the formation of the osazone. (I) decolorizes Schiff's reagent in 3-4 sec. and is considered to be an anhydro-aldehyde. Mutarotation is due to opening of the 1:5-ring, and not the more stable 3:6-ring, and equilibrium is between (I) and the aldehyde. For stereochemical reasons the pyranose ring can re-form only in one direction. R. S. C.

110-11A METALLURGICAL LITERATURE CLASSIFICATION

12-3

Hydrazones and osazones of sugars. E. VORO-
 ON and F. VALENTIN (Ark. Hronju, 1931, 8, 165—
 1932).—The sugar of which a given hydrazone is a
 derivative can be identified by distilling the hydr-
 azone with 12% HCl, when the production of furfur-
 aldehyde indicates the presence of pentose, and of
 methylfurfuraldehyde methylpentose, whilst neither
 is evolved by hexoses. This reaction can be applied
 to determine whether the pentose or methylpentose
 component of a disaccharide is responsible for its
 reducing action. Lactosephenylosazone exhibits
 mutarotation in MeOH, whilst the rotation of the
 phenylosazone of anhyd. lactose is constant. *p*-
 Nitrobenzaldehyde can conveniently be substituted
 for PhCHO in the regeneration of sugars from their
 hydrazones, in view of the greater insolubility of its
 hydrazone. Fructosephenylmethylsazone yields the
 corresponding phenylosazone on heating with excess
 NHPH-NH₂, whilst with *p*-bromophenylmethylhydr-
 azone a mixed osazone is obtained. The muta-
 rotation of osazones is not due to tautomerism
 between the dihydrazone and azo forms, as fructose-
 phenylmethylsazone, which does not possess a
 labile H atom, exhibits mutarotation.
 R. TRUSKOWSKI.

ASB-51A METALLURGICAL LITERATURE CLASSIFICATION

FROM SYNONYMS

SYNONYMS

RELATIONS

ALPHABETICALLY

NUMERICALLY

INTERNAL INDEX

OPEN

CLOSED

EXTERNAL INDEX

PROCESSES AND PROPERTIES INDEX

A-3

BC

β-Anhydromannonic- γ -lactone. Y. VALENTIN
 (Coll. Czech. Chem. Comm., 1957, 0, 315-326).—
 3:6-Anhydromannose treated with Br in H₂O for
 several days gives non-cryst. *β*-anhydromannonic
 acid (I) (amorphous as salt), which yields a phenyl-
 hydrazide, m.p. 180-5° (decomp.), [α]_D +16.7° in
 MeOH. This with KOH-H₂O-PbCHO at the b.p.
 gives the γ -lactone, m.p. 113°, [α]_D +120.5° in H₂O,
 falling slowly to +115.5° after 250 hr. K₂ selenate
 reacts violently with AcCl-H₂SO₄ giving *α*-
 dioxolactone- γ -*γ*-dilactone, m.p. 160-162°, [α]_D
 +155° in Ac₂O. It is concluded that the two rings of
 these and of other sugar compounds containing the
 bicyclic system  have the same optical
 character, [α] then being augmented, and that the
 effect increases with the no. of CO groups.
 E. W. W.

ASD SIA ORYTHOLOGICAL LITERATURE CLASSIFICATION

PROCESSED AND REPRODUCED FROM THE ORIGINAL SOURCE

10

CK

Anhydromannose, a new sugar anhydride. K. Yalen-
 tin. *Collection Czechoslov. Chem. Communications* 6,
 354-70 (1934).—Mannose, treated with MeOH and HCl,
 gave α -*M*s mannopyranoside (I) in 75% yield. By treat-
 ment of I in C_6H_5N with Ph_3CCl and acetylation, *6-trityl-2,3,4-triacetyl- α -methylmannoside* (II), $C_{41}H_{54}O_{10}$, was ob-
 tained, m. 131-2° (from MeOH or EtOH and ligroin),
 $[\alpha]_D^{25} 58.0^\circ$; yield 80%. Treated with PBr_3 in CCl_4 , II
 gave the corresponding *6-Br deriv.* (III), $C_{41}H_{52}O_{10}Br$, m.
 78-81° (from C_6H_6 and ligroin), $[\alpha]_D^{25} 57.8^\circ$; yield 45%.
 By action of NH_3 in MeOH, III was transformed into the
6-bromo- α -methylmannoside, m. 97-98°, $[\alpha]_D^{25} 62^\circ$; analy-
 sis showed this material to be slightly impure. By treat-
 ment with $Ba(OH)_2$, *3,6-anhydro- α -methylmannopyrano-
 side* (IV), $C_{12}H_{18}O_5$, was obtained, m. 130-2° (from EtOAc),
 $[\alpha]_D^{25} 97.1^\circ$. Crystals of IV showed $a:b:c = 0.996:1:$
 1.057 . The β m by Becke's method with H_2SO_4 , IV gave a
 β m. 1.522, γ m. 1.526. Hydrolyzed with H_2SO_4 , IV gave a
 sirup which crystd., m. 102-3°, $[\alpha]_D^{25} 95.92^\circ$. This was
3,6-anhydromannose (mannofuranoside) (V). It showed
 no mutarotation and required 10-15 min. to develop color
 with fuchsin reagent. It formed a *benzylphenylhydrazone*,
 $C_{21}H_{24}O_2N_2$, m. 144-5° (from C_6H_6), $[\alpha]_D^{25} 43.59^\circ$; a phenyl-
 hydrasone insol. in H_2O ; and an *osazone*, $C_{12}H_{12}O_4N_4$, m.
 188-90° (40% alc.), $[\alpha]_D^{25} -110.14^\circ$. V, treated with HCl
 in dry MeOH, gave a sirup which became cryst., $C_{12}H_{18}O_5$,
 m. 85° (from AcOEt and ligroin), $[\alpha]_D^{25} 157^\circ$; this was
3,6-anhydro- α -methylmannofuranoside. V's reasons for
 believing the structures assigned are correct are given in
 detail. M. P. Benoy

ASB-SLA METALLURGICAL LITERATURE CLASSIFICATION

MATERIALS INDEX												GENERAL INDEX											
GROUP												SUBJECT											
A B C D E F G H I J K L M N O P Q R S T U V W X Y Z												A B C D E F G H I J K L M N O P Q R S T U V W X Y Z											

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PROCESSES AND PROPERTIES INDEX

CA

Chemistry of sugar hydrazones and osazones. E. VOTUCEK AND F. VALENTIN. *Archiv. Hem. Farm.* 5, 155 61(101-2 French)(1931). -The sugar components can be recognized by boiling the hydrazones with 12% HCl, the hydrazones of a pentose methylpentose yield furfural or methylfurfural and hexose yields neither. The test can be effected microchemically with a few cg. of the sugar hydrazone. In osazone derivs. of disaccharides it can be seen whether a pentose or methylpentose component does or does not constitute the reducing part of the sugar mol. V. and V. measured the rotatory power of the phenylosazone of lactose and its anhydride in MeOH. The first diminishes with time, the second (often higher on account of the anhydride ring) remains const. Due to the smaller soly. of its hydrazone, β -O₂NC₆H₅-CHO renders possible an integral solution even where it is incomplete with BrH as in the case of rhodose methylphenylhydrazones. The hydrazone radicals can be replaced by other hydrazone residues, not only in hydrazones, but also often in osazones with the formation of a new osazone either simple or mixed. Thus, fructose methylphenylosazone, treated in the cold with an excess of PhNHNH₂·AcOH, gives the corresponding phenylosazone, e. g., the β -bromophenylhydrazone would furnish the mixed osazone, HOCH₂(CH(OH))₂C(:NNMePh)CH:NNHC₆H₅·Br. The theory of Zerner and Waltuch, according to which the inconstancy of rotatory power of osazones is due to the tautomeric change of a dihydrazone form into an azoic form, cannot be maintained, since fructose methylphenylosazone, which does not possess any H susceptible of migration, shows also the phenomenon of mutarotation. J. Krcma

COMMON ELEMENTS

MATERIALS INDEX

ASSOCIATED METALLURGICAL LITERATURE CLASSIFICATION

RESEARCH AND DEVELOPMENT

GROUPS

INDICES

GROUPS

GROUPS

10

ca

A new anhydrogalactose, *E. VALMIAN*. *Collection Czechoslov. Chem. Communica-*
tions 4, 364-75 (1952). *3,6-Anhydrogalactose* (I) has been prepd. and differs from those
 known in the stability of its anhydride ring which even decreases the stability of the
 original galactose ring. V. compares I to Fischer's 3,6-anhydroglucose (C. A. 6, 1295).
 On the basis of reducing power, osazone formation, the original pyran ring in galactose.
 and stereochem. reasons V. assigns to the anhydride ring the 3,6-positions. He notes
 that models show this configuration to be the only one which would weaken the pyran
 ring of galactose. He concludes that the ring is thus weakened since I reacts with Schiff's
 reagent and also shows a change of rotation. A stable 2-ring system should not show
 mutarotation (Fischer's 3,6-anhydroglucose does not) and V. believes that the new
 rotation is due to an equil. mixt. of bicyclic and aldehydic forms of I. V. also cites
 reasons to show that the pyran ring did not change to a furan ring. Anhyd. α -methyl
 galactoside in dry pyridine was treated with $PbCl_2$, then with Ac_2O , giving 78% 2,3,4-
 triacetyl-6-triisopropyl- α -methylgalactoside (II), m. 170-81°, $[\alpha]_D^{25}$ 6°. By treating II with
 $PbCl_2$ in $(CH_2Cl)_2$ he obtained 70% of the bromohydrin (III), brown and decmpa. 167-
 163°, $[\alpha]_D^{25}$ 18°. After refluxing a mixt. of III and $Ba(OH)_2$ for 2 hrs., 3,6-anhydro- α -
 methylgalactoside (IV) was isolated in almost theoretical yield, recrystd. from $AcOEt$, it
 m. 141-2°, $[\alpha]_D^{25}$ 24° (in H_2O). Hydrolysis of IV with 1% H_2SO_4 gave I which has
 been obtained only in the amorphous state. A water soln. of I was examd. in the
 polarimeter and the rate of change of rotation was found to follow the equation for a
 monomol. reaction. Extrapolation of the curve to zero time gave $[\alpha]_D^{25}$ 38.6°. The
 osazone m. 215°, $[\alpha]_D^{25}$ 48.2° (Me-OH).
 ANNIE E. WHITE

ABB. 51A 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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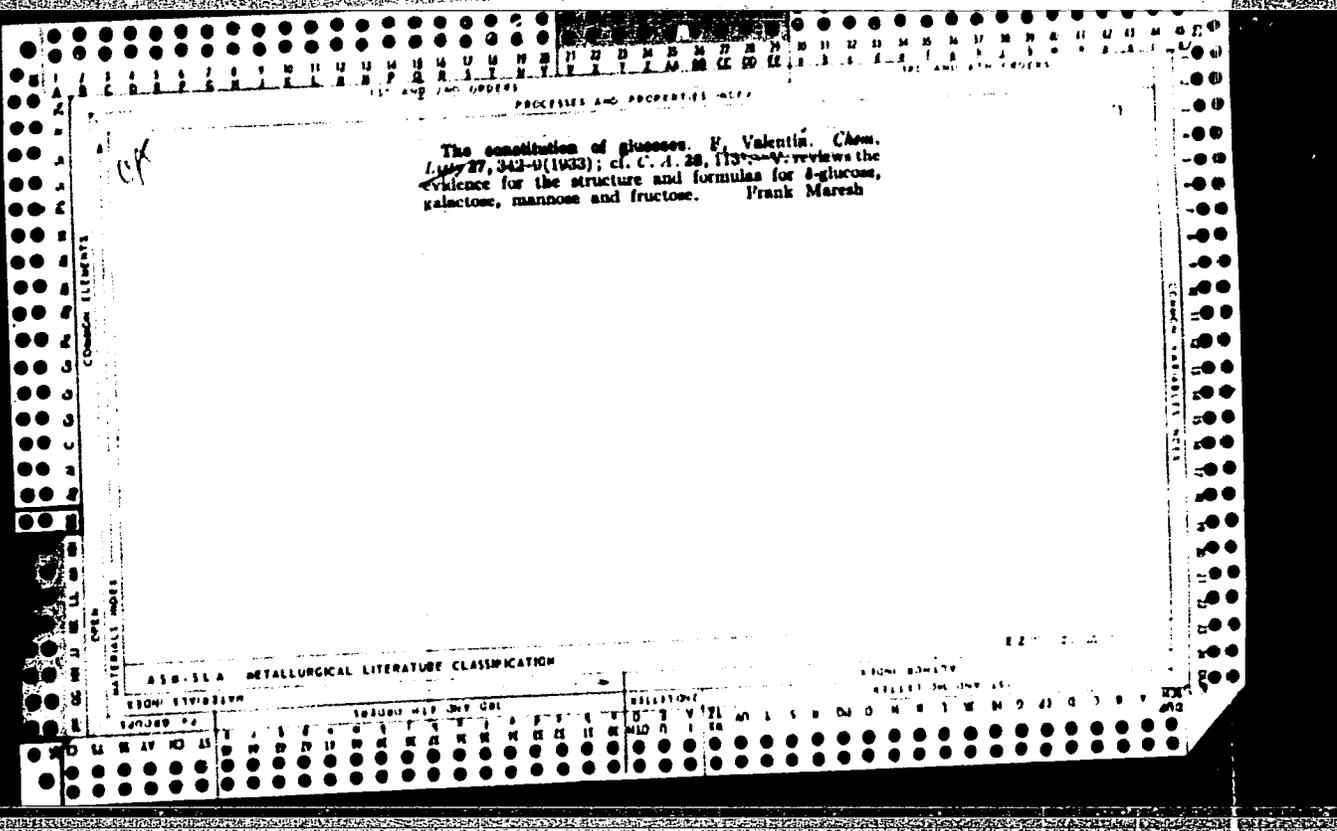
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ca

The glucosylalkylamines (and other aldose alkylamines). E. Votoček and E. Valentin. *Collection Czechoslov. Chem. Communications* 6, 77-90(1934).— These compds. are formed by the condensation of the aldoses with 1 mol. of the primary amine. They crystallize either in the anhyd. condition, with 1 or 0.5 mol. of H₂O or MeOH depending upon the solvent used. They show mutarotation and must therefore have the lactolic structure. The substituted hydrazines react with the alkylamines to give the hydrazone of the sugar and the corresponding amine. The following alkyl derivs. were prepd. by dissolving approx. 2 g. of the corresponding sugar in an aq. or MeOH soln. of the amine. The alkylamine deriv. crystallizes on standing. *Methylamines:* rhamnosyl, C₁₁H₁₉O₄N. 2H₂O, m. 101°; fucosyl, C₁₁H₁₉O₄N. N H₂O, m. 125°; rhamnosyl, C₁₁H₁₉O₄N. MeOH, m. 126-7°; glucosyl, C₁₁H₁₉O₄N. H₂O, m. 78-80° (decomp. 115°). *Rhamnoseethylamine,* C₁₁H₁₉O₄N. 0.5H₂O, m. 141-2°. *Propylamines:* rhamnosyl, C₁₁H₁₉O₄N, m. 145°; fucosyl, C₁₁H₁₉O₄N, m. 124-5°; galactosyl, C₁₁H₁₉O₄N, m. 127-8°. *Butylamines:* xylosyl, C₁₁H₁₉O₄N. H₂O, m. 81-2°; rhamnosyl, C₁₁H₁₉O₄N. MeOH, m. 136-7°; fucosyl, C₁₁H₁₉O₄N. 0.5H₂O, m. 88-9°; glucosyl, C₁₁H₁₉O₄N. H₂O, m. 97-8°; galactosyl, C₁₁H₁₉O₄N, m. 82-3°; mannosyl, C₁₁H₁₉O₄N. H₂O, m. 71-2°. *Amylamines:* rhamnosyl, C₁₁H₁₉O₄N. H₂O, m. 139-40°; fucosyl, C₁₁H₁₉O₄N. H₂O, m. 90-7°; galactosyl, C₁₁H₁₉O₄N. H₂O, m. 110°; mannosyl, C₁₁H₁₉O₄N. 0.5H₂O, m. 70-1°. *Hexylamines:* xylosyl, C₁₁H₁₉O₄N. H₂O, m. 87°; rhamnosyl, C₁₁H₁₉O₄N, m. 132-3°; fucosyl, C₁₁H₁₉O₄N, m. 100-10°; glucosyl, C₁₁H₁₉O₄N. H₂O, m. 79-80°; mannosyl, C₁₁H₁₉O₄N. 0.5 H₂O, m. 75°. *Heptylamines:* rhamnosyl, C₁₁H₁₉O₄N. m. 138°; fucosyl, C₁₁H₁₉O₄N. 0.5H₂O, m. 84-5°; glucosyl, C₁₁H₁₉O₄N. H₂O, m. 97°; galactosyl, C₁₁H₁₉O₄N, m. 99-100°, softens 80°. W. A. Moore

E 2

METALLURGICAL LITERATURE CLASSIFICATION



1ST AND 2ND GROUPS

PROCESSES AND PROPERTIES INDEX

ca

The constitution of glucose. *J. P. Valentin, Chem. Listy 27, 275-8, 301-3, 319-25(1933).*—The constitution of sugars as polyhydroxy aldehydes, butylene oxides and amyleno oxides is reviewed. The stable and unstable forms of their deriva. are discussed in connection with the semiacetal ring. Frank Marsh

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COMMON ELEMENTS

WATER AND INDEX

ASB-51A METALLURGICAL LITERATURE CLASSIFICATION

ALPHABETIC INDEX

1ST AND 2ND LETTERS

3RD AND 4TH LETTERS

5TH AND 6TH LETTERS

7TH AND 8TH LETTERS

9TH AND 10TH LETTERS

11TH AND 12TH LETTERS

13TH AND 14TH LETTERS

15TH AND 16TH LETTERS

17TH AND 18TH LETTERS

19TH AND 20TH LETTERS

21ST AND 22ND LETTERS

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87TH AND 88TH LETTERS

89TH AND 90TH LETTERS

91ST AND 92ND LETTERS

93RD AND 94TH LETTERS

95TH AND 96TH LETTERS

97TH AND 98TH LETTERS

99TH AND 100TH LETTERS

LIST AND NO CROSS PROCESSES AND PROPERTIES INDEX

A

Phytochemical notes. I. The glucosides from the gum of *Viscaria vulgaris* (Lycalis *Viscaria* L.). E. VOROČEK AND E. VALENTIN. *Collection Czechoslov. Chem. Commun.* 4, 282-4(1962).—The principal part of the gum of *Viscaria vulgaris* consists of a polyose or polyoses with a base of d-glucose and d-xylose. R. C. R.

ADD-51A METALLURGICAL LITERATURE CLASSIFICATION

#	1	2	3	4	5	6	7	8	9	0	A	B	C	D	E	F	G	H	I	J	K	L	M	N	O	P	Q	R	S	T	U	V	W	X	Y	Z
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PROCESSES AND PROPERTIES INDEX

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ca

The glucosylketimines, tertiary compounds of sugar, ammonia and β -diketones. R. Votreck and F. Valentin. *Collection Czechoslov. Chem. Communications* 7, 201-208 (1955).—Rhamnose, 0.4 g., is dissolved in MeOH and 2.1 g. acetylacetone and 11 cc. MeOH containing 0.1105 g. NH_3 per cc. is added. *Rhamnosyliminomethylcrotonate*, $\text{O} \cdot \text{CHMe} \cdot [\text{CH}(\text{OH})]_4 \cdot \text{CHNHCMc} : \text{CHCO}_2\text{Me}$, crystd. in 2 days; recrystd., m. 20° (decompn.), $[\alpha]_D -172.5^\circ$. Ten g. rhamnose is dissolved in MeOH and NH_3 passed in until the soln. is acid. Upon adding 12.8 g. CH_3Ac and cooling, *Me rhamnosyliminocrotonate*, $\text{O} \cdot \text{CHMe} \cdot [\text{CH}(\text{OH})]_4 \cdot \text{CHNHCMc} : \text{CHCO}_2\text{Me}$ is formed, recrystd., m. $190-2^\circ$, $[\alpha]_D -125.5^\circ$. *Et rhamnosyliminocrotonate*, similarly prepd. from rhamnose, NH_3 and $\text{Ac} \cdot \text{CH} : \text{COEt}$ in EtOH, m. 185° , $[\alpha]_D -121.0^\circ$. *Et rhamnosyliminomethylcrotonate*, m. 182° , $[\alpha]_D -112.5^\circ$, prepd. from AcCHMeCOEt . *Me rhamnosyliminomethylcrotonate*, from AcCHMeCOEt , m. $192.5-3^\circ$, $[\alpha]_D -111^\circ$, prepd. from CH_3EtAc . *Et rhamnosyliminomethylcrotonate*, m. $191.5-2.5^\circ$, $[\alpha]_D -107.5^\circ$, prepd. from AcCH_2COEt . *Et mannosyliminocrotonate*, m. 179° , $[\alpha]_D 159^\circ$, prepd. from mannose, NH_3 and AcCH_2COEt .
P. H. Moser

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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CA

10

Acetone derivatives of xylitol and their constitution. Frantiek Valentin and Dezider Tomkuljak (Slovak Tech. Univ., Bratislava, Czech.). *Chem. Zvesti* 3, 146-64 (1949). --Diacetonexylitol was prepd. and its constitution detd. On treatment with Me_2CO and concd. H_2SO_4 , xylitol (I) binds first 1 mol. of Me_2CO in a 6-membered heterocycle at positions 3 and 5 of the I mol., forming the amorphous moniacetonexylitol, $b.p.$ 145-7°, n_D^{20} 1.4778. Prolonging the reaction time and increasing the amt. of the condensation agent gives diacetonexylitol, in which the 2nd mol. of Me_2C is bound at positions 1 and 2 of the sugar mol., forming a 5-membered heterocycle; the amorphous 1,2,3,5-diacetonexylitol $b.p.$ 99-101°, n_D^{20} 1.4534. The following derivs. of I were prepd. by oxidation with $\text{Pb}(\text{OAc})_2$: 1,2,3,5-diacetone-4-tolylsulfonyl, *cryst.*, *m.* 71-3°; 3,5-acetone-1,2,4-tribenzoyl, *cryst.*, *m.* 103-4°; 3,5-acetone-1,2,4-trimethyl, amorphous, $b.p.$ 69-71°, n_D^{20} 1.4324; 1,2,4-trimethyl, amorphous, $b.p.$ 97-9°, n_D^{20} 1.4610; 1,2,3,5-diacetone-4-methyl, amorphous, $b.p.$ 78-80°, n_D^{20} 1.4383; 3,5-acetone-4-methyl, amorphous, $b.p.$ 109-11°, n_D^{20} 1.4603; 4-methyl, amorphous, $b.p.$ 167-9°; 1,2,3,5-tetrabenzoyl-4-methyl, *cryst.*, *m.* 121-3°.

Jan Micka

CA

Vitamin C in fresh and canned vegetables. František
Valentia and Danica Žuffová (Research Inst. Food Ind.,
Bratislava, Czech.). *Chem. Zvesti* 4, 309-12(1950).—
Twenty-one various kinds of vegetables in Bratislava region
was tested by Tillman's method for vitamin C content when
fresh and canned vitamin C was very much lower in the
canned vegetables. Jan Miska

CA

Vitamin C in various fruits. Frantíček, Valentin and
Dana Zúřová (Food Research Inst., Bratislava, Czech.).
Chem. Zvesti 4, 8-13(1950).— Twenty varieties of fruit
from the Bratislava region were tested for their vitamin C
by the Tillmans method and compared with the finished
product. Jan Muka

A

Quantitative separation of magnesium cation from sodium and potassium cations. Prantlček Václav and Magda Suchánová-Toufková (Bratislava, Czech). *Chem. J. Zvesti 6, 68-69 (1950)*. The salts of Na⁺, K⁺, and Mg²⁺ is evaporated to dryness with H₂SO₄ and ignited. After the sulfates are dissolved, freshly prepd. AgOH or Ag₂O (H₂O) is added. Mg(OH)₂ is adsorbed on the dispersed particles of AgOH.

Jan Miska

12

CA

Vitamin C content of tomatoes. Frantisek Valentin
and Danica Zuffová (Food Research Inst., Bratislava,
Czech.). *Chem. Zvesti* 3, 340-53(1949).—The effect of
vegetation period and climatic conditions on vitamin C in
tomatoes was studied. There were losses of 61.4-85.2%
in the finished product (catsup) as compared with the
original tomatoes contg. 15.0-38.00 mg.% of vitamin C
as detd. by Tillmans method. Jan Micka

Valentin, F.

Vitamins in food research in Slovakia. F. Valentin, D. Žuffová, P. Hanula, M. Čunderliková, and I. Stein (Výsk. ústav potravinárského priemyslu, Bratislava, Czech.). *Průmysl Potravin* 4, 20-4(1953).--In 72 different kinds of vegetables, fruits, oils, fats, and food products the contents of the following vitamins (I) were detd.: A, B₁, B₂, E, niacin, and folic acid. More than 450 tabulated values from approx. 5000 tests. are presented. Preventive measures against losses of I in manufg. processes are suggested. L. J. Ucháček

VALENTIN, F.

ry

C Z E C H

Vitamins A (axerophthal) and carotene in raw materials of Slovakian food industry. F. Valentín, D. Žaffová, and M. Čunderlíková (Výzkumný ústav potravního průmyslu, Bratislava, Czech.). *Chem. Zvesti* 8, 267-71(1954); cf. C.A. 49, 625c.—Structural formulas and occurrence of β -carotene in Slovakian fruits and vegetables are discussed. J. M.

VALENTIN, F.

"Vitamin A (axerophthol) and carotenes in basic raw materials of the Slovak food industry."
Chemicke Zvesti, Bratislava, Vol. 8, no. 5, May 1954, p. 167.

SO: Eastern European Accessions List, Vol. 3, No. 11, Nov. 1954, L.C.

VALENTIN, F.

"Chemistry of Penicillin as an Antibiotic of Today", P. 218, (TECHNICKA
FPACA, Vol. 6, No. 4, April 1964, Bratislava, Czechoslovakia)

SO: Monthly List of East European Accessions, (EEAL), IC, Vol. 4,
No. 1, Jan. 1955, Uncl.

VALENTIN, FRANT

Czechoslovakia/Chemical Technology. Chemical Products and Their Application --
Food industry, I-28

Abst Journal: Referat Zhur - Khimiya, No 2, 1957, 6769

Author: Valentin, Frant

Institution: None

Title: Capsaicin -- The Substance that Imparts the Burning Taste to Red
Pepper

Original
Publication: Prumysl potraviny, 1955, 6, No 8, 383-387

Abstract: Data concerning the structure of capsaicin, its occurrence in nature,
its physiological and pharmacological action and methods for its de-
termination. Bibliography, 8 references.

Card 1/1

VALENTIN, F.

Coloring matter of paprika. p. 638. CHEMICKE ZVESTI. Bratislava.
Vol. 9, no. 10, Dec. 1955.

SOURCE: East European Accessions List (EEAL), LC, Vol. 5, no. 3, March 1956.

VALENTIN, F. ; HANULA, F; ARSAL, J.

Effect of raw materials on the biological quality of corn extracts. p.55.
CHEMICKA ZVEST. (Slovenska akademija vied a Spolok chemicov na Slovensku)
Bratislava. Vol 10, no. 1, January 1956.

SOURCE: East European Accessions List, (EEAL), Library of Congress
Vol. 5, no. 12, December 1956.

VALENTIN, N.

The circulation speed of current assets in industrial enterprises. p. 70.

METALURGIA SI CONSTRUCTIA DE MASINI

Vol. 8, no. 3, Mar. 1956

Rumania

Source: EAST EUROPEAN LISTS Vol. 5, no. 10 Oct. 1956

BAKATIS, V.; VALENTINOVICH, A.

Investigating the increase in the use of
heat-insulating radi-peat boards. *Izvestiya*,
B no.2:253-264 '62.

1. Institut stroitel'stva i arkhitektury

VALENTINCIC, Joza, ing. (Beograd, Mladena Stojanovica 4)

Housing construction in the Scandinavian countries and in Yugoslavia.
(To be contd.) Tehnika Jug 17 no.1:28-31 Ja '62.

(Scandinavia—Housing)
(Yugoslavia— Housing)

VALENTINCIC, Joze, ing. (Beograd, Mladena Stojanovica 4)

Construction of housing in the Scandinavian countries and in
Yugoslavia. Tehnika Jug 17 no.2:245-253 F '62.

1. Deputy Secretary for Industry of the Federal Executive Council,
Beograd.

(Scandinavia--Housing)
(Yugoslavia--Housing)

VALENTINCIC, M.

Intestinal parasites in Slovenia. Zdrav. vest., Ljubljana 23 no.
11-12:312-316 1954.

1. Mikrobioloski institut medicinske visoke sole v Ljubljani-
predstojnik prof. dr. Milica Valentincic.

(PARASITES,
intestinal, in Slovenia, in child., determ. technic (Slov))

VALENTINCIC, M.; STROPNIK, Z.

Histoplasma capsulatum. Zdrav. vest., Ljubljana 24 no.7-8:273-277
1955.

1. Mikrobioloski institut medicinske fakultete v Ljubljani-
predstojnik prof. dr. Milica Valentincic.

(HISTOPLASMA
capsulatum (S1))
(HISTOPLASMOSIS, diag.
(S1))

VALENTINCIC, M.; LEBEZ, D.; VOZELJ, M.

Jakobstahl complement fixation test with cardiolipin antigen.
Acta med. iugosl. 10 no.1:50-58 1956.

1. Department of Serology, Institute of Microbiology, Medical
Faculty, University of Ljubljana.

(COMPLEMENT

fixation with cardiolipin antigen in diag. of syphilis,
Jacobsthal technic.)

(CARDIOLIPIN

antigen complement fixation in diag. of syphilis,
Jacobsthal technic)

(SYPHILIS, diag.

complement fixation test with cardiolipin antigen,
Jacobsthal technic)

(ANTIGENS

cardiolipin antigen complement fixation in diag. of syphilis,
Jacobsthal technic)

VALENTINCO, M.

CZECHOSLOVAKIA / Zooparasitology. Parasitic Protozoa. G-2

Abs Jour: Ref Zhur-Biol., No 20, 1958, 91046

Author : ~~Valenticio, M.~~, Kozak, M.

Inst : Not given

Title : The Dysentery Ameba and Environmental Temperature

Orig Pub: Zdravstv. vestn., 1957, 26, No 26, No 11, 456-457 (Slovenian)

Abstract: No abstract

Card 1/1

EXCERPTA MEDICA Sec 12 Vol 13/6 Ophthalmology June, 59...

969. PYRAZOLIDINE TREATMENT OF ENDOGENIC IRIDOCYCLITIS (Russian text) - Valentinene A. B. - VESTN. OPTALM. 1959, 7³ (31-36)
Pyrazolidine, an analogue of butazolidine (pyrazolon derivative), was the drug used by the author in treatment of 50 patients with endogenic iridocyclitis of various aetiology. It is most effective in relieving pain and decreasing inflammation. Acute clinical forms with increased exudation are especially susceptible to treatment by pyrazolidine. Investigation of the permeability of the blood vessels of the anterior portion of the eye in 15 patients demonstrated that this drug promotes normalization of vessel permeability. However, this takes place only after the disappearance of inflammation. Oral administration of the drug in the dose of 0.15 g. t.i.d. gives the greatest effect on the 5th-6th day. However, it is recommended that the treatment be continued up to the 7th-8th day for prophylaxis of early recurrences. At the same time the main endogenic cause of iridocyclitis should be treated. Gastro-duodenal ulcers, diseases of the haemopoietic organs, liver and kidney, insufficiency of the cardiac valves and pronounced arteriosclerosis are contraindications to the use of this drug. There were no complications when the drug was used in the above doses. Pyrazolidine may be employed in out-patient departments, with control of blood and urine data.

VALENTINENE, A. B., Cand Med Sci -- (diss) "Pyrazolidine in the treatment of of iridocyclitis." Moscow, 1960. 16 pp; (Second State Moscow Medical Inst im N. I. Pirogov); 250 copies; free; (KL, 51-60, 120)

VELKEY, Laszlo, dr.; TOTH, Anna, dr.; VALENTINI, Jozsef, dr.

Methemoglobinemia in infants caused by drinking water. Orv.
hetil. 105 no.5:201-203 2F '64.

1. Borsod megyei Sennelweis Korhaz, I. Gyermekosztaly.

*

NAZAROV, M.I.; PATRUSHEV, M.F., inz., retsenzent; LEGOSTAYEV, A.M., retsenzent;
TALMAZA, V.F., retsenzent; VALENTINI, L.A., kand.tekhn.nauk, retsenz-
sent; KABAKOV, M.M., red.; ANOKHINA, M.G., tekhn.red.

[Paved canals] Moshchenye kanaly. Frunze, Akad.nauk Kirgizskoi
SSR, 1958. 104 p. (MIRA 12:3)
(Irrigation canals and flumes)

VALENTINI, L.A., kand.tekhn.nauk; TYAN, V.K., inzh.

Investigating the regimen of bed load flow in small
mountain rivers. Trudy SANIIRI no.95:3-15 '58.

(MIRA 13:6)

(Sukuluk River--Hydraulics)

VALENTINI, L.A., kand.tekhn.nauk

Construction of earth dams by depositing earth into water.

Trudy SANIIRI no. 98:101-104 '59.

(MIRA 14:1)

(Dams) (Hydraulic engineering)

VALENTINI, L.A., kand.tekhn.nauk

Headworks on mountain and piedmont sections of small rivers.
Trudy SANIIRI no. 104:3-14 '59. (MIRA 14:1)
(Hydraulic engineering)

VALENTINI, I.A., kand. tekhn. nauk; DERLYATKA, T.I., inzh.; NAUMENKO, Yu.G.
inzh.; SHISHORINA, G.I., inzh.

Destruction of the Kugart Dam and its analysis. Gidr. i mel. 13
no.9:54-61 S '61. (MIRA 14:9)

(Kugart River--Dams)

ARG024060

(N)

SOURCE CODE: UR/0124/66/000/004/B071/B071

AUTHOR: Valentini, L. A.; Derlyatka, T. I.

TITLE: Theory of an oblique hydraulic jump and its practical application

SOURCE: Ref. zh. Mekhanika, Abs. 4B486

REF SOURCE: Sb. Vopr. gidrotekhniki. Vyp. 23, Tashkent, Nauka, 1965, 12-18

TOPIC TAGS: hydraulics, fluid flow, flow analysis

ABSTRACT: The authors examine the problem of conjugate depths and magnitude of the angle β between the direction of the front of a jump and the direction of a turbulent flow in an oblique hydraulic jump arising at the vertical break of the sides of the channel. The equation of the law of conservation of mass and the equation of the theorem of impulses in projections onto the normal to the front of the jump and onto the direction of the front of the jump itself are used. This makes it possible to obtain equations determining the conjugate depths and angle β . Results are shown that the angle β can be found by calculating the propagation velocity of the disturbing wave in a flow of finite depth. Graphic relations for the above-indicated jump parameters are constructed. It is pointed out that the vertical break of the sides of the channel leads to a change in the conditions of the bottom streams of the flow which begin to move in the direction of the jump front. This can be used for reducing the silt saturation of the flow by constructing a special opening for

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ACC NR: AR6024060

discharging the silt. Abstractor's comment. The results obtained by the authors correspond to one case of an oblique jump which is realized provided the front of the jump passes through the site of the break of the channel sides and provided the lengths of the channel sections parallel to the front of the oblique jump are equal on both sides of the front and therefore have a special character. The experiments used by the authors pertain, for example, to the case where the second provision is not fulfilled. [Translation of abstract] V. S. Sinel'shchikov

SUB CODE: 20

Card 2/2

VALENTINOV, A., podpolkovnik

Determination of geodetic data for long-range firing. Voent.
vest. 41 no.11:79-83 N '61. (MIRA 16:11)

KUKUYEV, Ye.M.; YEFIMOV, V.F.; FLIORIN, B.S., otv.red.; VALENTINOV,
A.M., red.; ABRAMYAN, A.A., red.; KISELEV, N.A., red.; METLIN,
V.A., red.; ANDREYEV, G., tekhn.red.

[Handbook with nomenclature and prices for materials and equipment
used in the coal industry] Nomenklaturnyi spravochnik i tseny na
materialy i oborudovanie, primenyaemye v ugol'noi promyshlennosti.
Moskva. Group 2. [Nonferrous metals] TSvetnye metally. 1950.
275 p. (MIRA 13:4)

1. Russia (1923- U.S.S.R.) Ministerstvo ugol'noy promyshlennosti.
(Nonferrous metals)
(Coal mines and mining--Equipment and supplies)

VALENTINOV, A.M.

For firmly establishing measures of economy in all branches of the
coal industry. Ugol' 29 no.1:7-11 Ja '54. (MLRA 7:1)

1. Nachal'nik Finansovogo upravleniya Ministerstva ugol'noy promyshlen-
nosti SSSR.

(Coal mines and mining)

VALENTINOV, B. (Krymskaya oblast')

Improving the system of collecting income tax from collective farms. Vop. ekon. no.3:154-156 Mr '62. (MIRA 15:3)
(Nishnegorskiy District--Collective farms--Taxation)

VALENTINOV, B.

We are helping collective farmers to organize their finances.
Fin.SSSR 37 no.4:66-68 Ap '63. (MIRA 16:4)

1. Starshiy inspektor gosudarstvennykh dokhodov Nizhnegorskogo rayonnogo finansovogo otdela Krymskoy oblasti.
(Nizhnegorskiy District--Collective farms--Finance)

VALENTINOV, G.

Propagandist of aeronautical knowledge. Kryl. rod. 16 no.2:26
F '65. (MIRA 18:3)

VALENTINOV, I.

Africa - Social Conditions

"Stories about Africa." Vokrug Sveta no. 5, '52.

2

9. Monthly List of Russian Accessions, Library of Congress, July 195~~4~~. Unclassified.

VALENTINOV, N.

USSR/Radio - Trade Organization

Dec 51

"Concerning the Work of 'Soyuzposyltorg,'"
N. Valentinov

"Radio" No 12, pp 14, 15

Finds the following defects in the work of
"Soyuzposyltorg": poor assortment of receivers
(only the Moskvich and Rodina were available in
1951), delay in filling orders (40-50 days),
high shipping costs (set of batteries for the
Rodina receiver priced at R 101.20 cost 78 rubles
to ship), and poor advertising of available radio
parts.

208T83

1. VALENTINOV, N.
2. USSR (600)
4. Radio, Short-Wave
7. Master radio amateurs, Radio No. 1, 1953.

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

VALENTINOV, N

107-5-22/54

AUTHOR: Valentinov, N.

TITLE: The Victory of Soviet Sportsmen. International Encounters
(Pobeda sovetskikh sportsmenov. Mezhdunarodnyye vstrechi)

PERIODICAL: Radio, 1956, Nr5, pp. 22-23 (USSR)

ABSTRACT: Over 500 duplex radio amateur stations and hundreds of receiving stations took part in the IV International Short-Wave Ham Contest which was organized by the League of Friends of Soldier of the Polish People's Republic. There were hams from Bulgaria, Hungary, East Germany, Poland, Romania, Czechoslovakia, and USSR. The subject of contest: telephone and telegraph duplex radio communication.

The Chief Umpire Board worked in Warsaw and consisted of the following members: A. Yeglinski SP1CM (Poland), K. Nesterov LZ2KAC (Bulgaria), T. Matusek SP6XA (Poland), P. Vasilesku YO6VG (Romania), N. Kazanskiy YA3AΦ (USSR), Kaminek, K. OK1CX (Czechoslovakia), I. Yezerski SP2SJ (Poland).

Soviet hams won the first place in the contest. Among them:

Operator of the Kiyev radioclub Y55KAA, and of the Saratov radioclub YA4K4E; L. Labutin of Moscow YA34P; operator of the Dnepropetrovsk radioclub Y55KAD; Yu. Chernov of Saratov YA44B; A. Shchennikov of Pensa YA4Φ4; operator of the Taganrog Radiotechnical Institute YA6KOD;

Card 1/2

VALENTINOV, N.

Forward in force. Radio no.9:7-8 S '56.
(Radio, Shortwave--Competitions)

(MLBA 9:11)

MADZHAROV, D.I.; VALENTINOV, H.

Electrocardiographic changes following anesthesia in children.
Vest. khir. no.10:91-93 '64. (MIRA 19:1)

1. Iz nauchno-issledovatel'skogo instituta vosstanovitel'noy
khirurgii, protezirovaniya i trudoustroystva (dir. - doktor
Iv. Iliyev), Sofiya, Bolgariya.

VALENTINOV, N., inzh.

Lighting will be cheaper. Izobr.i rats. no.4:8-11 Ap '60.
(MIRA 13:6)
(Electric lamps)

VALENTINOV, N., inzh.; NOVINSKIY, G., vrach

An invention should subsist. Izobr.i rats. no.12:16-18 D '60.

(MIRA 13:12)

(Medical instruments and apparatus--Technological innovations)

KYARDI, Ya., brigadir (g.Tallin); KAPRANOV, G. (g.Mal'chik); KNYAZEV,
Yu. (g.Mal'chik); SHAPKUN, N., inzh. (g.Krasnodar); KHOKHLOV,
Yu. (g.Ural'sk); VALENTIKOV, N., inzh.; HOVINSKIY, G., vrach

Innovations. Izobr. i rsts. no.9:12-13 S '61. (MIRA 14:8)

1. Machal'nik tekhnicheskogo otdela zavoda imeni Zemlyachki,
g. Ural'sk (for Khokhlov).

(Technological innovations)

VALENTINOV, N. (Alma-Ata)

Photographic clubs have not yet been organized in Alma-Ata.
Sov. foto 19 no.5:26-27 My '59. (MIRA 12:9)
(Alma-Ata--Photography--Societies, etc.)

VALENTINOV, Oleg

The Fifth Congress of the Trade Unions of Czechoslovakia. Vsem.
prof. dvizh. no.7/8:35-37 JI-Ag '63. (MIRA 16:10)

VALENTINOV, R.; NIKOLAYEV, I.

In primary schools, institutes of higher learning and at practical
training. Sov.foto 22 no.9:42-43 S '62. (MIRA 15:8)
(Photography)

VALENTINOV, R.; IGOREV, N.

Let's talk about your photographs. Sov. foto 23 no.4:42-44
Ap '63. (MIRA 16:5)

(Photographs)

VALENTINOV, V.

Metal workers strike. Sov.profsoiuzy 5 no.3:87-88 Mr '57.
(MLRA 10:4)

(Schleswig--Holstein--Strikes and lockouts--Metal workers)

VALENTINOV, V.

It does not suit anybody. Izobr.i rats. no.9:46-47 S '60.

(MIRA 13:10)

(Technological innovations)

VALENTINOV, V.

Callisthenics in industry. Okhr. truda i sots. strakh. 4
no. 2:50-51 F '61. (MIRA 14:2)
(Callisthenics)

30900. VALENTINOV, Ye.

Vospitaniye meditsinskoy sestry. (Shkola pri bol'nitse im. Ostroumova).
Med. sestra, 1949, No. 9, s. 30-32.

VALENTINOV, Ye. (g. Kuybyshev)

Here they repair motor vehicles. Prom.koop. 12 no.11:15
N '58. (MIRA 11:11)
(Kuybyshev-Automobiles-Maintenance and repair)

SCHWARTZ, E.; VALENTINOVA, I.; SIMKOVA, V.; DORNETZHUBER, V.

Determination of the biological effect of gibberellic acid
in guinea pigs with some biochemical methods. Bratisl. lek.
listy 44, no. 10:621-627 30 N '64

1. Krajska nemocnica tuberkulozy a chorob plucnych. (riaditel
MUDr. K. Virsik), a Ustav tuberkulozy v Bratislave (riaditel
MUDr. J. Markovic).

WENFA, Rudolf, Ing.; VALENTINOVA, Ingrid, prom. biochemik.

Spraying of sugar beets with scatox 20. Pracovni lek. 9 no.2:144-146
Apr 57.

1. Ustav hygieny prace a chorob z povolania v Bratislave, riaditel
MUDr I. Klucik.

(PARATHION,
spraying of sugar beets (Cz))

VALENTINOVA, I.

JANOK, J., RNDr. Prom biochem.; MAJEROVA, Zd., Prom biochem.; VALENTINOVA, I., Ing.;
MASEK, J., Ing.; TICHY, V., Ing.

In vitro anticholinesterase effect of coumarin dialkylphosphoric &
dialkylthiophosphoric acid esters. Pracovni lek. 9 no.6:506-512 Dec 57.

1. Ustav hygieny prace z chorob z povolania v Bratislave, prednosta MUDr.
I. Klucik Vyskumny ustav agrochemickej technologic v Bratislave. J. J.
Bratislava, Ustav hygieny prace.

(COUMARIN, eff.

dialkylphosphoric & dialkylthiophosphoric acid esters, in
vitro anticholinesterase ff. (Cz))

(CHOLINESTERASE, antag.

coumarin dialkylphosphoric & dialkylthiophosphoric acid esters
in vitro (Cz))

CZECHOSLOVAKIA/Chemical Technology - Pesticides.

H.

Abs Jour : Ref Zhur - Khimiya, No 16, 1958, 54969

Author : Tikhly, Rattay, Yanok, Valentinova

Inst : -

Title : Mixed Esters of Pyrocatechine, Phosphoric and Thiophosphoric Acid Derivatives.

Orig Pub : Chem. zvesti, 1957, 11, No 7, 398-410

Abstract : From the reaction of 1,2-phenylene chlorothiophosphate with sodium alcoholates, the following compounds of the general formula, $1,2-C_6H_4(O)_2P(S)OR$, were synthesized

(given are: R, yield of the crude material in %, m. p.

in °C., n_D^{20} , d_4^{20}) : C_6H_5 , 90.7, 71-71.5; -, -; $C_6H_4NO_2-P$, 97.3, 88, -, -; $C_6H_4NO_2-O$, 97.6, 141.5-142,

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CZECHOSLOVAKIA/Chemical Technology - Pesticides.

H.

Abs Jour : Ref Zhur - Khimiya, No 16, 1958, 54969

-, -; C_6H_4Cl-o , 99.3, 125.5, -, -; $C_6H_4OCH_3-o(1)$, 73.6,
86.5-87.5, -, -; $C_6H_3Cl_2-2',4'$, 92.0, 80.0, -, -;
4'-methyl coumarinyl-7' (II), 92.8, 146, -, -; C_2H_5 ,
70.4, -, 1.5622, 1.2954; $P(S)(OC_3H_7-n)_2$ (III), 87.8,
-, 1.5311, 1.2514.
Treating 1,2- $C_6H_4(ONa)_2$ with $(C_2H_5O)_2.PSCL$, a 94.9%
yield of 1,2-phenylene-bis-(o,o-diethyl) thiophosphate
(IV) was obtained, b. p. 110°C./0.1 mm., $n_D^{20} = 1.5110$,
 $d_4^{20} = 1.2610$. In the reaction of pyrocatechin with

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CZECHOSLOVAKIA/Chemical Technology - Pesticides.

H.

Abs Jour : Ref Zhur - Khimiya, No 16, 1958, 54969

dialkyl chlorophosphate in the presence of sodium carbonate, 1,2-phenylene-bis(dialkyl)-phosphates are formed (given are: alkyl, yield in %, b. p.

in °C./mm., n_D^{20} , d_4^{20}) : C_2H_5 , 36.0, 112-116/0.08, 1.4679,

1.2110 n- C_3H_7 (V), 28.6; 114-118/0.035 (decomposes);

1.4815, 1.1709 iso- C_3H_7 , 41.2, 112-116/0.05, 1.4715,

1.1581.

The most active insecticides for *Musca Domestica* are compounds III-V. The I_{50} for cholinesterase in the plasma (CP) and the erythrocytes (E) of human blood was determined. CP was the most sensitive to the esters obtained, and only II is active upon CP and E. I has a selective action upon CP.

Card 3/3

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cholinesterase inhib. in vitro (Cz))

(CHOLINESTERASE, antagonists,

O, O-dialkyl-S-(N,N-dialkylthiocarbamyl)-dithiophosphate,
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(RICKETS)

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PROCESSING AND PROPERTY MODES

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COMMON ELEMENTS
COMMON VARIABLES MODES
MATERIALS MODE
OPEN

450-550 METALLURGICAL LITERATURE CLASSIFICATION

FROM SOURCE
GROUP

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