- Heusser, H., Engel, C., Herzig, P. T. and Plattner, P. A. Helv. Chim. Acta 23 (1950) 2229.
- Engel, C. J. Am. Chem. Soc. 78 (1956) 4727.
- Venus-Danilowa, E. D. J. Gen. Chem. USSR 11 (1941) 847; Chem. Abstr. 36 (1942) 4094.
- Aston, J. G. and Greenburg, R. B. J. Am. Chem. Soc. 62 (1940) 2590.
- Morrell, R. S. and Bellars, A. E. J. Chem. Soc. 85 (1904) 345.
- 18. Rappe, C. Arkiv Kemi 21 (1963) 503.

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Crystallographic Data on N,N,N',N', Tetrakis (2-aminoethyl) ethylenediamine Pentahydrochloride = [Penten,5HCl], Hydroxy-Penten-Cobalt (III) Iodide and Hydroxy-Penten-Cobalt (III) Perchlorate

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Gauss, Moser and Schwarzenbach<sup>1</sup> prepared "penten", 5HCl in connection with investigations of metal complexes with polyamines. Crystallographic data on this compound (1) and on the cobalt(III) complexes  $[Co(OH)penten]I_2$  (2) and  $[Co(OH)penten](ClO_4)_2$  (3) were obtained from rotation and Weissenberg photographs recorded with CuKa radiation and from precession photographs recorded with

CuKa and MoKa radiation. The densities of the crystals were measured by the flotation method.

The crystals of all three compounds are monoclinic; from systematic absences (1) and (2) must have the space group  $P2_1/c$  (No. 14,  $C_{2k}^{5}$ ), although (3) has space group P2 (No. 3,  $C_2^{1}$ ) or Pm (No. 6,  $C_s^{1}$ ) the reflections h0l with l=2n+1 and 0k0 with k=2n+1 are very weak and so (2) and (3) are nearly isomorphous.

Detailed studies of these three com-

pounds are in progress.

The crystals were kindly supplied by professor G. Schwarzenbach, Zürich.

 Gauss, W., Moser, P. and Schwarzenbach, G. Helv. Chim. Acta. 35 (1952) 2359.

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## A Conversion of D-Glucose to D-Mannose under Acidic Conditions

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In an attempt to prepare D-manno-furanose pentaacetate, 2,3:5,6-di-O-iso-propylidene-D-mannose was treated with a mixture of acetic acid, acetic anhydride and sulphuric acid at room temperature. When the deacetylated product was investigated by paper chromatography, however, it gave two spots, which corresponded to D-glucose and D-mannose. By fractionation of the mixture these two sugars were

Table 1.

	(1)	(2)	(3)
Formula	$\mathrm{C_{10}H_{33}N_6Cl_5}$	$\mathrm{CoC_{10}H_{29}N_6OI_2}$	$\mathrm{CoC_{10}H_{29}N_6O_9Cl_2}$
Formula wt.	414.7	562.2	507.2
$D_{m}$ (g·ml <sup>-1</sup> )	1.34	2.07	1.71
$D_x$ (g·ml <sup>-1</sup> )	1.35	2.07	1.71
Space group	$P2_{_1}/c$	$P2_{1}/c$	
Z (calc.)	4	4	4
a (Å)	10.19	10.52	10.57
b (Å)	23.18	15.31	15.99
c (Å)	9.96	11.31	11.76
β (°)	120.0	98.4	98.4
V (Å <sup>3</sup> )	2037	1802	1966

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