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ANNUAL REPORT OF THE INTERNATIONAL COMMITTEE ON ATOMIC WEIGHTS, 1913.

Received October 12, 1912.

Since the annual report for 1912 was prepared a number of important memoirs on atomic weights have appeared. There are also one or two earlier researches which were received too late to be noticed at the proper time. These investigations may be summarized as follows:

Nitrogen.—Wourtzel,¹ by oxidizing NO to N_2O_4 , has redetermined the ratio between nitrogen and oxygen. Five concordant measurements give, in mean, N = 14.0068.

Potassium and Chlorine.—Staehler and Meyer² have made careful analyses of potassium chlorate, with especial precautions against contamination by the chloride. Their final series gives KCl = 74.555I, whence K = 39.097 and Cl = 35.458. For a discussion of their results, see also Guye,³ who concludes that the impurity above mentioned was, if not completely, at least sufficiently eliminated to be practically negligible.

Fluorine.—McAdam and Smith⁴ have published two preliminary determinations of the atomic weight of fluorine. Sodium fluoride was converted into chloride by heating in dry, gaseous hydrochloric acid, and from the ratios between the weights the atomic weight was calculated. The two values found are F = 19.0176 and 19.0133.

Phosphorus.—From analyses of phosphorus tribromide Baxter, Moore and Boylston⁵ find, in mean of three series, P = 31.027 when Ag = 107.88.

¹ Compt. rend., 154, 115.

² Z. anorg. Chem., 71, 368.

³ J. chim. phys., 10, 145.

^{*} THIS JOURNAL, 34, 592.

⁵ Proc. Amer. Acad., 47, 585; THIS JOURNAL, 34, 259.

This agrees fairly well with the former work of Baxter and Jones on silver phosphate. Further work on phosphorus trichloride is promised.

Mercury.—Easley and Brann,¹ by analyses of mercuric bromide, find Hg = 200.64. This confirms the previous work of Easley on the chloride.

Selenium.—Kuzma and Krehlik² have redetermined the atomic weight of selenium by reduction of SeO_2 with SO_2 . The mean of ten determinations is Se = 79.26.

Tellurium.--Harcourt and Baker³ have thrown doubt upon the work of Flint, who claimed to have split up the supposed element into two fractions of different atomic weight. They repeated his method of fractionation, and from the fourth fraction found Te = 127.54. This agrees with the figure found by Baker and Bennett in 1907. Similar fractionations have been carried out also by Pellini⁴ who likewise failed to find any indication of a tellurium of low atomic weight.

Radium.—Hönigschmid,⁵ by careful analyses of relatively large quantities of radium chloride, finds Ra = 225.95. On the other hand, Gray and Ramsay,⁶ using very small quantities of material, and converting the bromide into the chloride, find Ra = 226.36, in agreement with previous work by Madame Curie and Thorpe. Until the discordance between Hönigschmid's low value and the higher is explained, it is undesirable to change the figure given in the table.

Tantalum.—The determinations of this atomic weight by Chapin and Smith⁷ were made by the hydrolysis of $TaBr_5$. The mean of eight determinations gave Ta = 181.80, a figure somewhat higher than that found by Balke from similar analyses of the pentachloride.

lridium.—Hoyermann,⁸ by five reductions of $(NH_4)_2IrCl_8$ in hydrogen, finds Ir = 192.613.

Holmium.—Six determinations of the atomic weight of holmium by Holmberg⁹ gave Ho = 163.45. The well known sulfate method was employed.

There are also approximate determinations of the atomic weights of lead, zinc and copper by Pecheux,¹⁰ and of calcium by Oechsner de Coninck.¹¹ The figures obtained are not conclusive enough to justify

¹ THIS JOURNAL, 34, 137.

² Trans. Bohemian Acad. of Emperor Frances Joseph, 19, No. 13, 1910. Data furnished the committee by Professor B. Brauner.

⁸ J. Chem. Soc., 99, 1311.

⁴ Atti Accad. Lincei, 21, 218.

^b Monatsh. Chem., **33**, 253.

⁶ Proc. Roy. Soc., (A) 86, 270.

⁷ This Journal, 33, 1497.

⁸ Sitzungsb. phys. med. Soz. Erlangen, 42, 278.

[•] Z. anorg. Chem., 71, 226.

¹⁰ Compt. rend., 154, 1419.

¹¹ Ibid., 153, 1579.

their use in the table. for the methods employed were not of great accuracy.

International Atomic Weights, 1913.

	Symbol.	Atomic weight.	Symbol.	Atomic weight.
Aluminium	. A1	27.1	Molybdenum	96.0
Antimony	Sb	120.2	NeodymiumNd	144.3
Argon.	A	39.88	Neon	20.2
Arsenic	As	74.96	NickelNi	58.68
Barium	Be	137.37	Niton (radium emanation).Nt	222.4
Bismuth	Bi	208.0	NitrogenN	14.01
Boron.	B	11.0	OsmiumOs	190.9
Bromine	Br	79.92	OxygenO	16.00
Cadmium	Cd	112.40	PalladiumPd	106.7
Caesium.	Cs	132.81	PhosphorusP	31.04
Calcium	Ca	40.07	Platinum	195.2
Carbon.	C	12.00	Potassium	39.10
Cerium.	Ce	140.25	Praseodymium	140.6
Chlorine.	Cl	35.46	RadiumRa	226.4
Chromium	Cr	52.0	Rhodium	102.9
Cobalt	Co	58.97	RubidiumRb	85.45
Columbium	Сь	93.5	RutheniumRu	101.7
Copper	Cu	63.57	SamariumSa	150.4
Dysprosium	Dy	162.5	Scandium	44.I
Erbium	Er	167.7	Selenium	79.2
Europium	Eu	152.0	SiliconSi	28.3
Fluorine	F	19.0	SilverAg	107.88
Gadolinium	Gd	157.3	SodiumNa	23.00
Gallium	Ga	69.9	StrontiumSr	87.63
Germanium	Ge	72.5	SulfurS	32.07
Glucinum	Gl	9.1	Tantalum	181.5
Gold	Au	197.2	TelluriumTe	127.5
Helium	H e	3.99	TerbiumTb	159.2
Holmium	H o	163.5	ThalliumTl	204.0
Hydrogen	H	1.008	ThoriumTh	232.4
Indium	In	114.8	ThuliumTm	168.5
Iodine	. I	126.92	$Tin \dots Sn$	119.0
Iridium	Ir	193.1	Titanium	48.I
Iron	Fe	55.84	TungstenW	184.0
Krypton	Kr	82.92	UraniumU	238.5
Lanthanum	La	139.0	VanadiumV	51.0
Lead	Pb	207.10	XenonXe	130.2
Lithium	Li	6.94	Ytterbium (Neoytterbium).Yb	172.0
Lutecium	. L u	174.0	YttriumYt	89.0
Magnesium	Mg	24.32	ZincZn	65.37
Manganese	M n	54.93	ZirconiumZr	90.6
Mercury.	Hg	200 .6	-	

Only one change is recommended in the table for 1913; namely, the insertion of holmium, for which, hitherto, no good atomic weight determination has been available. Two or three other alterations of small

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importance might be made, but it seems undesirable to make changes too frequently.

(Signed) F. W. CLARKE, T. E. THORPE, W. OSTWALD, G. URBAIN.

SOME PHYSICAL CONSTANTS OF SULFUR TRIOXIDE. MELTING AND BOILING POINTS, DENSITY, COEFFICIENT OF EXPANSION AND MOLECULAR WEIGHTS.¹

BY D. M. LICHTY. Received August 30, 1912.

The existence of two forms of sulfur trioxide, distinguished as the α - and β -forms, has been generally recognized. The same method of distinguishing the two forms will be used in this paper. The former has a definit melting point, while the latter probably does not melt at all



under atmospheric pressure, but on being heated changes to vapor, whose condensation gives rise to the α -form, which may later go over into the β -form.

The sulfur trioxide used in this work was prepared by repeated distillation of it, from phosphoric anhydride, in an evacuated and sealed apparatus like that previously described.² For the purpose of taking the melting and boiling points, the receiver was given the form shown in Fig. 1. When the trioxide had been quite thoroughly dehydrated, there was distilled enough of it to fill the receiver to the line l. Dry air³ was now slowly admitted to atmospheric pressure, and then the neck of the receiver was bent through considerably more than a right angle and sealed off.

Melting Point and Boiling Point.

Fig. 1.—First made by Mr. H. P. Eastman, to whom I wish here to acknowledge my indebtedness. On being cooled to 10 or 12°, such a specimen froze to a mass of what looked like parallel lying rods. After slight

¹ Read at the Eighth International Congress of Applied Chemistry, New York, September, 1912.

² THIS JOURNAL, **30**, 1835-37. ³ Ibid., 1837.

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