Diagrams explanatory of the chemical decompositions of the London Pharmacopæia, and of the various processes used in medical chemistry necessary to be known by students preparing for examination at Apothecaries Hall. / By Thomas Harper Whitaker.

Contributors

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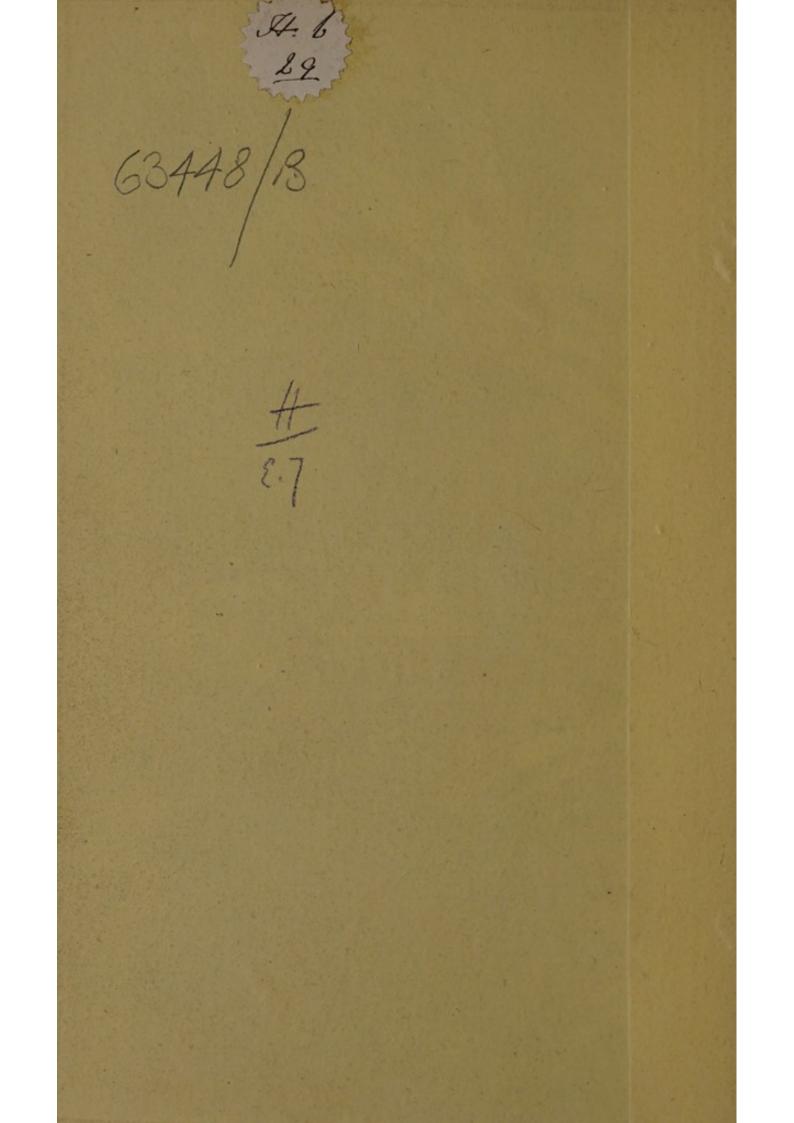
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CHEMICAL DIAGRAMS OF THE LONDON PHARMACOPCEIA

WHITAKER.



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WHITAKER, T.H.

AMSA H.n.22 DIAGRA EXPLANATORY OF THE Daublou MOL CHEMICAL DECOMPOSITIONS Loudow

OF THE

LONDON PHARMACOPCEIA,

AND OF

THE VARIOUS PROCESSES USED IN MEDICAL CHEMISTRY

NECESSARY TO BE KNOWN BY STUDENTS

PREPARING FOR EXAMINATION AT APOTHECARIES' HALL.

By THOMAS HARPER WHITAKER,

MEMBER OF THE ROYAL COLLEGE OF SURGEONS IN LONDON; LICENTIATE OF THE WORSHIPFUL COMPANY OF APOTHECARIES; PRIVATE TEACHER OF MEDICINE, SURGERY, ETC.

> LONDON: JOSEPH BUTLER, MEDICAL BOOKSELLER AND PUBLISHER, 4, ST. THOMAS'S STREET, SOUTHWARK.

> > MDCCCXXXIX.

G. WOODFALL, ANGEL COURT SKINNER STREET LONDON.

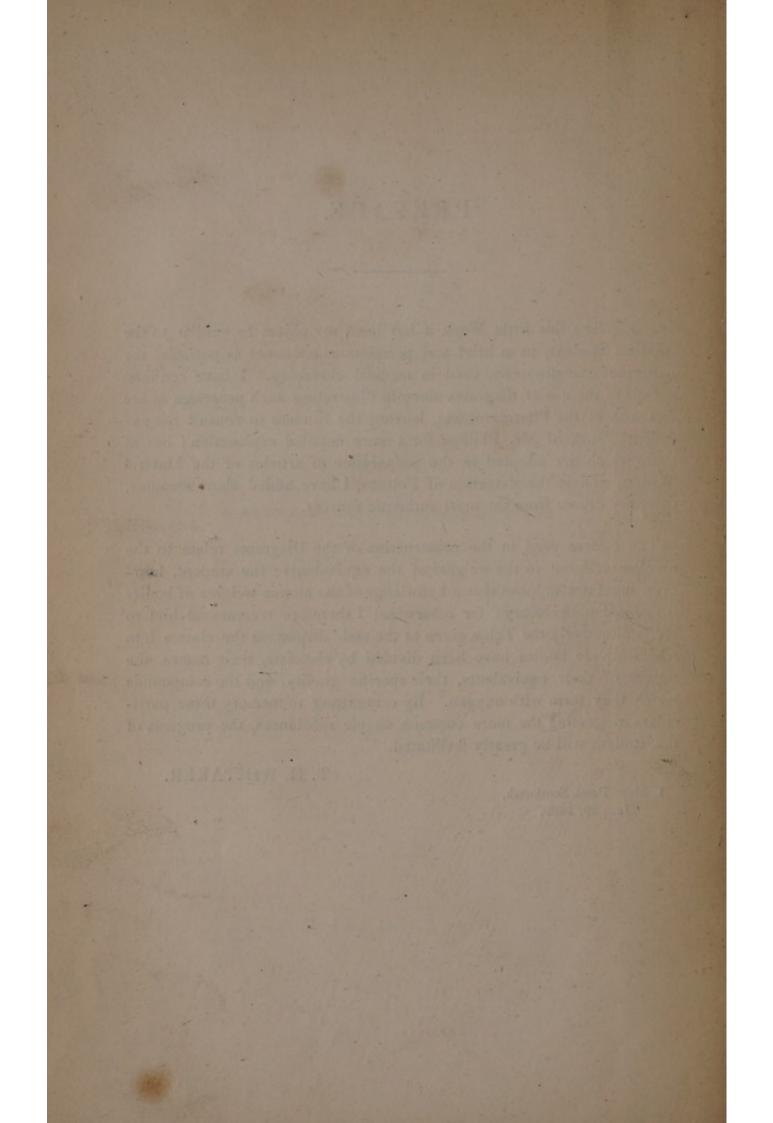
PREFACE.

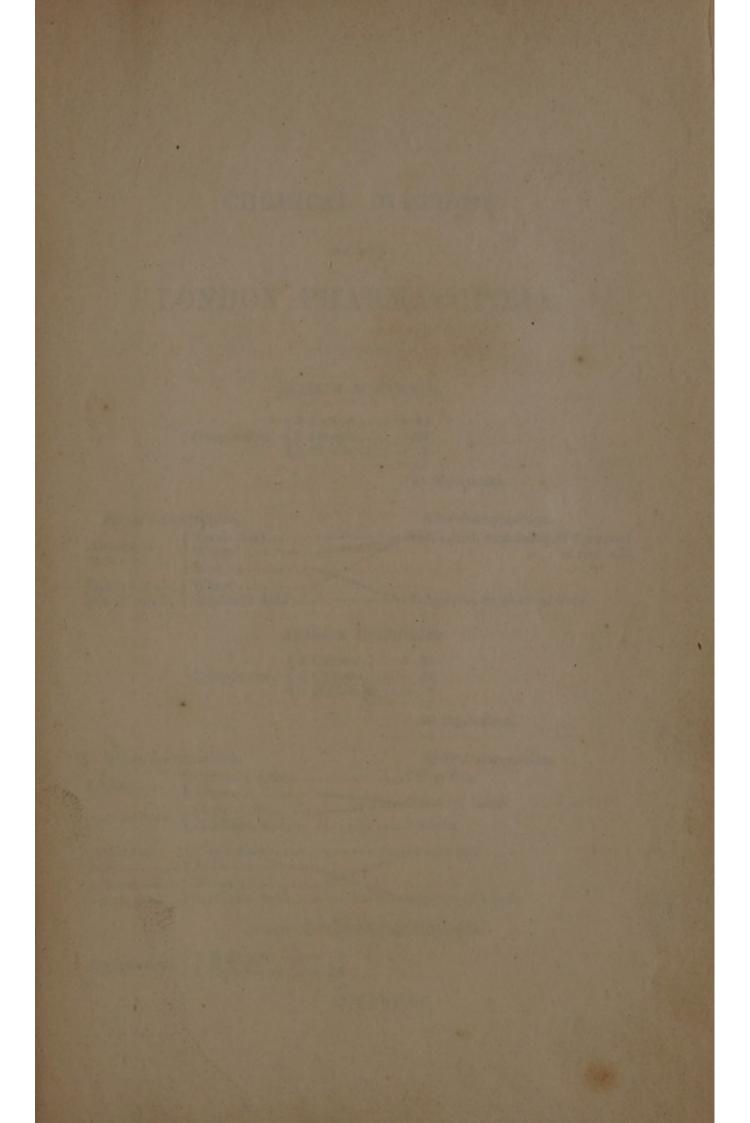
In compiling this little Work it has been my object to exhibit to the Medical Student, in as brief and perspicuous a manner as possible, the nature of the processes used in medical chemistry. I have confined myself to the use of diagrams alone in illustrating such processes as are met with in the Pharmacopœia, leaving the Student to consult the excellent Work of Mr. Phillips for a more detailed explanation; but of those which are adopted in the preparation of articles of the Materia Medica, and in the detection of Poisons, I have added short accounts, carefully drawn from the most authentic sources.

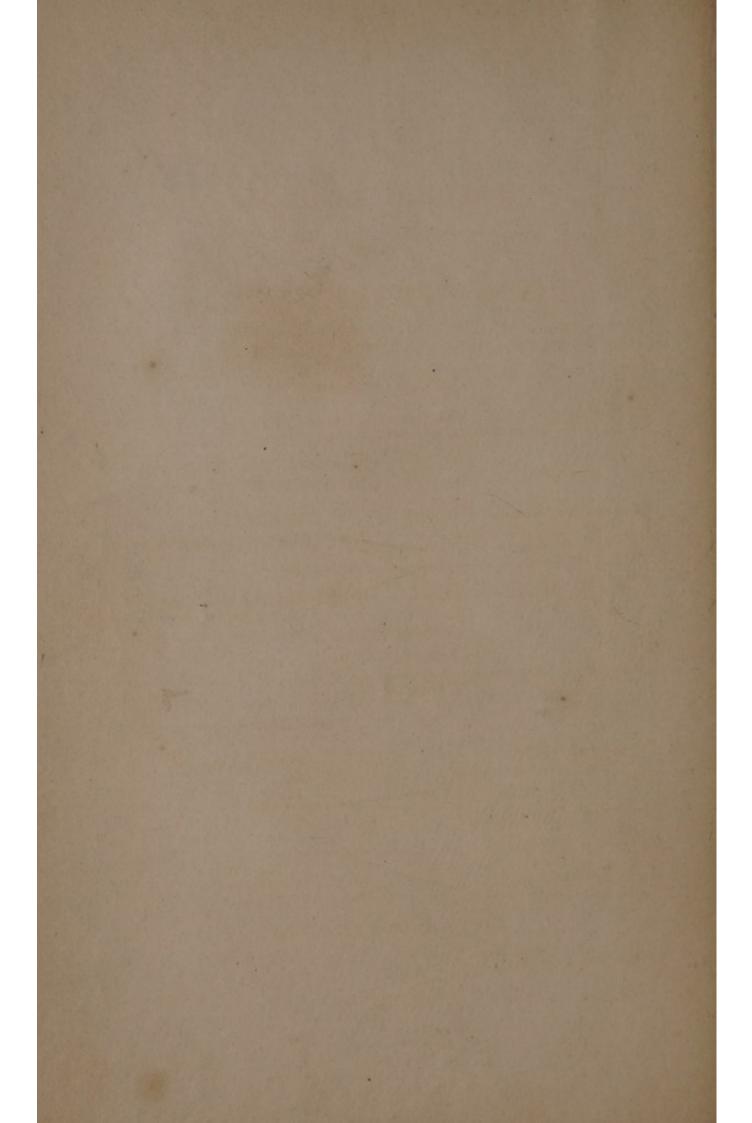
The figures used in the construction of the Diagrams relate to the number and not to the weights of the equivalents; the student, however, must not suppose that a knowledge of the atomic weights of bodies is deemed unnecessary; far otherwise; I therefore recommend him to study attentively the Table given at the end, displaying the classes into which simple bodies have been divided by chemists, their names, the weight of their equivalents, their specific gravity, and the compounds which they form with oxygen. By committing to memory these particulars respecting the more common simple substances, the progress of the Student will be greatly facilitated.

T. H. WHITAKER.

1, Maze Pond, Southwark, June 29, 1839.







OF THE

LONDON PHARMACOPŒIA.

ACIDUM ACETICUM.

Composition.

 $\begin{cases} 4 \text{ Carbon } \dots = 24 \\ 3 \text{ Oxygen} \dots = 24 \\ 3 \text{ Hydrogen} \dots = 3 \end{cases}$

51 Equivalent.

 Before decomposition.
 After decomposition.

 Acetate of Soda.....
 Acetic Acid....
 Acetic Acid, containing 30.8 per cent. of real acid.

 Diluted Sul-phuric Acid.
 Water
 Anhydrous Sulphate of Soda.

ACIDUM CITRICUM.

Composition. $\begin{cases} 4 \text{ Carbon } \dots = 24 \\ 4 \text{ Oxygen } \dots = 32 \\ 2 \text{ Hydrogen } \dots = 2 \end{cases}$

58 Equivalent.

1.

B

Before decor	nposition.	After decompositio
	Carbonic Acid	Off as Gas.
Lemon juice.	{ Citric Acid	Citrate of Lime.
Citrate of Lime	{ Citric Acid	In solution.
Diluted Sul- phuric Acid.	Water	Sulphate of Lime.

ACIDUM HYDROCHLORICUM.

Composition. $\begin{cases} 1 \text{ Hydrogen} \dots = 1 \\ 1 \text{ Chlorine} \dots = 36 \end{cases}$

37 Equivalent.

2

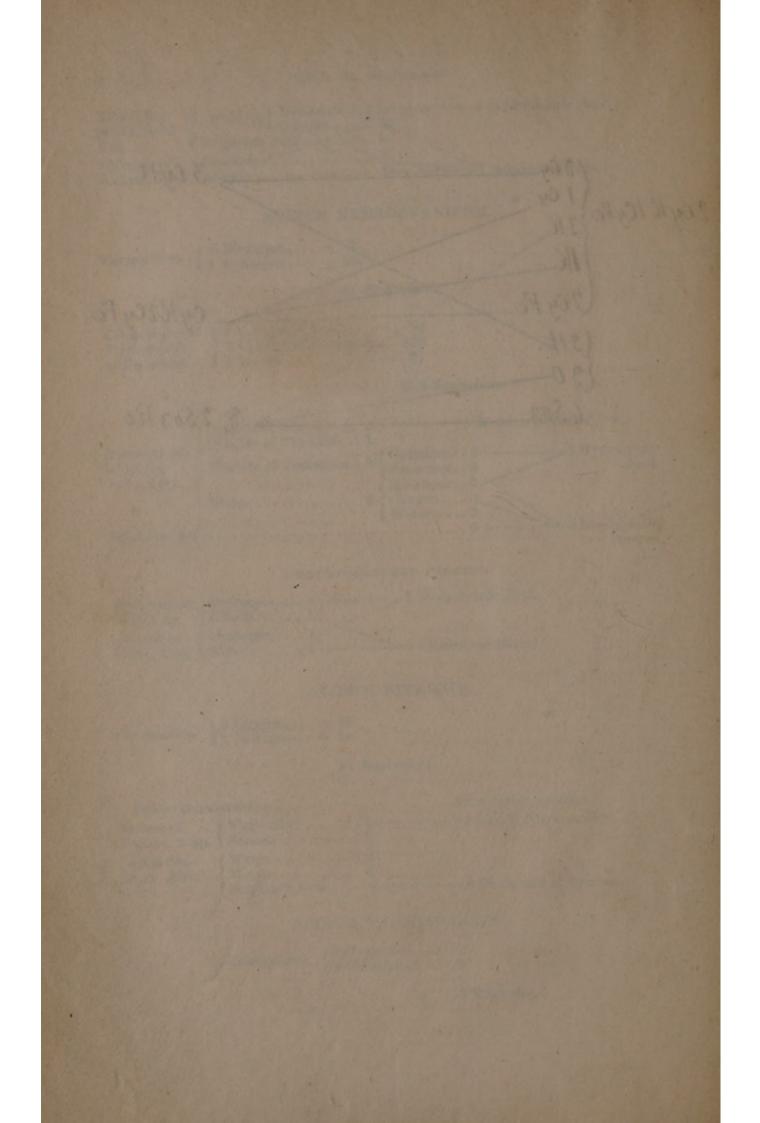
Liquid Sul- phuric Acid, 1 eq. Water 1 Hydrogen1 — 1 Hydrochloric Acid. Sulphuric Acid.
1 eq. (Sulphuric Acid Acid Chloride of Chloride of Chlorine Sodium, 1 eq. (Sodium, 1 Sulphate of Soda.
ACIDUM HYDROCYANICUM.
nonbeni intenteeni.
Composition. $\begin{cases} 1 & \text{Hydrogen} \dots = 1 \\ 1 & \text{Cyanogen} \dots = 26 \end{cases}$
27 Equivalent.
Composition of Ferrocyanide of Potassium 132 3 Water 27
213 Equivalent.
Ferrocyanide Cyanide of Iron
of Potas- { Cyanide of Potassium. 3 Potassium. 3
sium, 2 eqs. Hydrogen 3
Water 6 Oxygen 3 Water 3
Sulphuric Acid
Sulphuric Acid 6
Sulphuric Acid
Sulphuric Acid 6 3 Bisulphate of Potassa. EXTEMPORANEOUS PROCESS. Hydrochloric { Hydrogen 1 1 Hydrocyanic Acid. Acid, 1 eq. { Chlorine 1 1 Hydrocyanic Acid. Acid, 1 eq. { Chlorine 1 1 Hydrocyanic Acid. Cyanide of { Cyanogen 1 1 Chloride of Silver. Silver, 1 eq. { Silver 1 1 Chloride of Silver. ACIDUM NITRICUM. Composition. { 5 Oxygen
Sulphuric Acid 6 3 Bisulphate of Potassa. EXTEMPORANEOUS PROCESS. Hydrochloric { Hydrogen 1 1 Hydrocyanic Acid. Acid, 1 eq. { Chlorine 1 1 Hydrocyanic Acid. Cyanide of { Cyanogen 1 1 Chloride of Silver. Silver, 1 eq. { Silver 1 1 Chloride of Silver. ACIDUM NITRICUM. 6 6 Composition. { 5 Oxygen 14 54 Equivalent. 54 Equivalent. Before decomposition. After decomposition. After decomposition. Nitrate of { Nitric Acid 2 2 2 Potassa, 2 eqs. { Potassa 2 2 2

Composition. $\begin{cases} 5 \text{ Oxygen} \dots = 40 \\ 2 \text{ Phosphorus} = 32 \end{cases}$

72 Equivalent.

.

2. 2 Cy K ICy Fe 3 K 1 Cy 3 K 2 Cy Fe 3 H 3 O 6 Si 3 Cytt CyK2CyFe 6503 3, 2 503 120



OF THE LONDON PHARMACOPŒIA.

Nitric Acid. {Nitric Oxide.....Off as Nitrous Acid Gas. Oxygen......Phosphorus.....Phosphoric Acid.

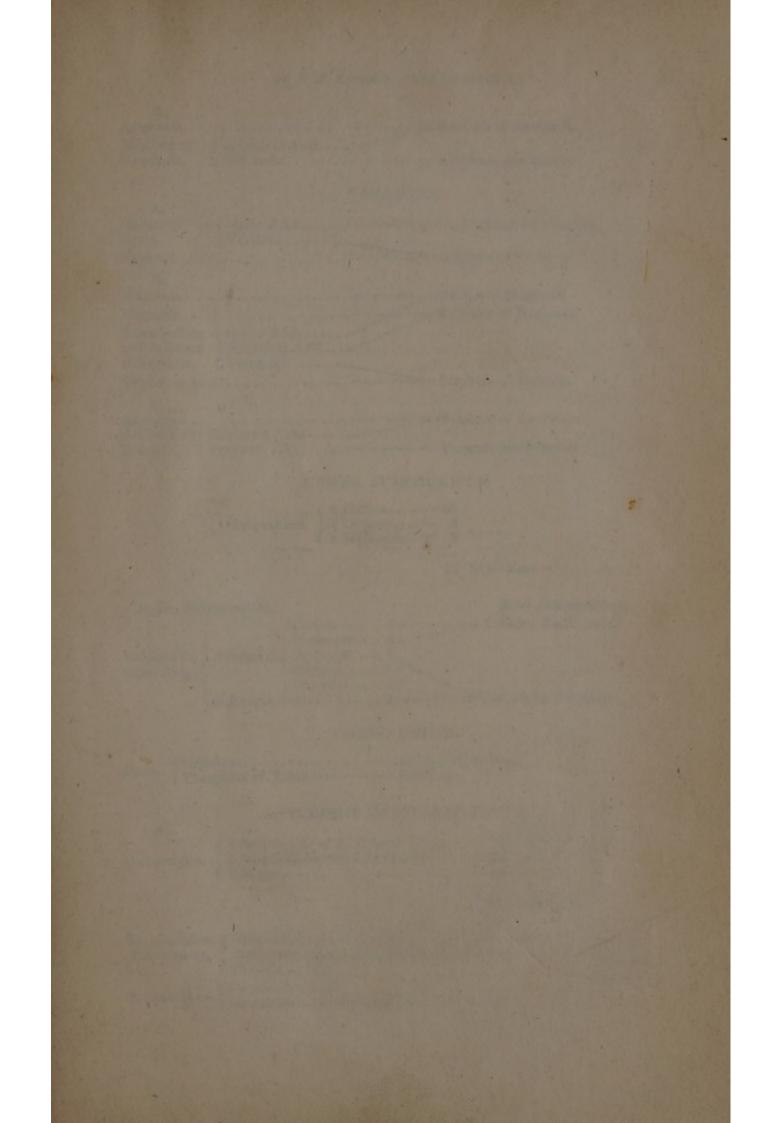
ACIDUM TARTARICUM.

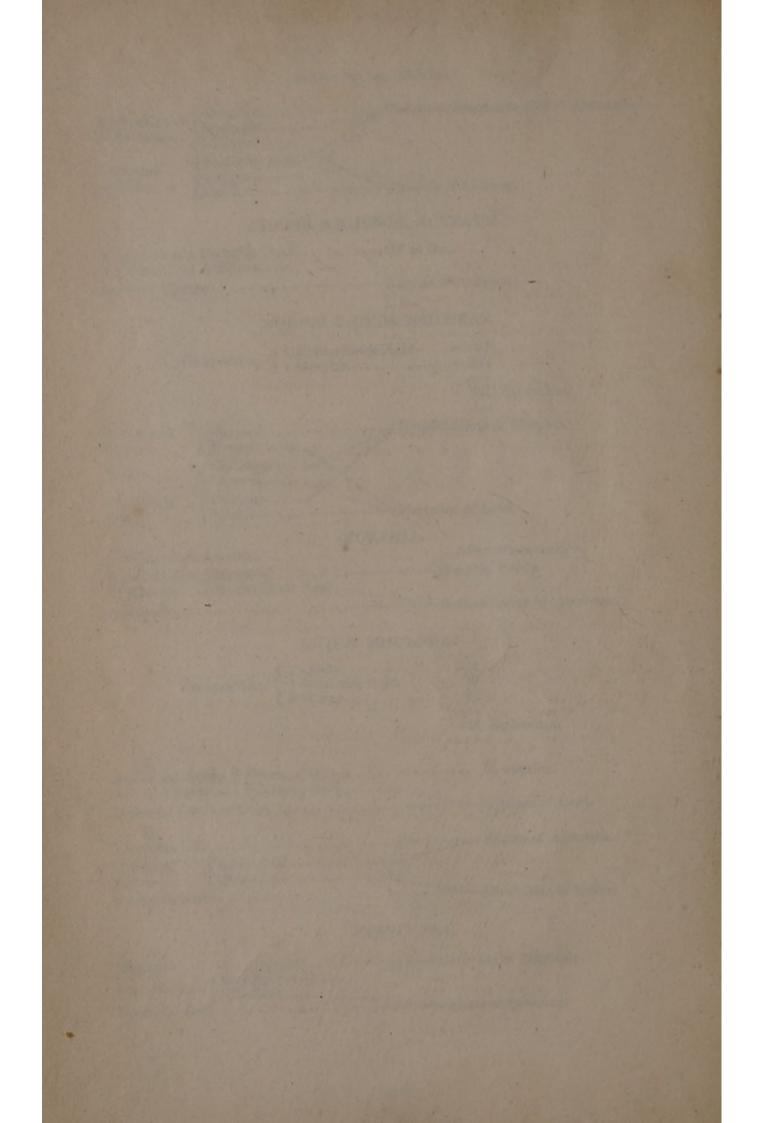
Composition. $\begin{cases} 4 \text{ Carbon } \dots = 24 \\ 5 \text{ Oxygen} \dots = 40 \\ 2 \text{ Hydrogen} \dots = 2 \end{cases}$

66 Equivalent.

Before decomposition. After decomposition. A. [Tartrate of Potassa 1--- In solution. Bitartrate of Potassa, 1 eq.) Tartaric Acid 1_ **B**. $\begin{cases} Carbonic Acid.....1 \dots 1 Off as gas \\ Lime \dots 1 \\ Calcium \dots 1 \end{cases}$ Carbonate of Lime, 1 eq. {Hydrogen..... 1 Chlorine 1 Chloride of Calcium. Hydrochloric Acid, 1 eq ... C. _____ Chloride of Potassium. Chloride of (Chlorine 1-Calcium, 1 eq. | Calcium Tartrate of S Potassa 1 { Potassium ... 1 Oxygen..... 1 Potassa, 1 eq.) Tartaric Acid 1->1 Tartrate of Lime. D. -In solution. Tartrate of Tartaric Acid..... Lime..... Lime. Water Diluted Sul-Sulphate of Lime. Sulphuric Acid phuric Acid. LIQUOR AMMONIÆ. Composition of Ammoniacal Gas. $\begin{cases} 1 \text{ Nitrogen} \dots = 14 \\ 3 \text{ Hydrogen} \dots = 3 \end{cases}$ 17 Equivalent. Hydrochlorate Ammonia.... -Liquor Ammoniæ. Hydrogen of Ammonia. Chlorine Oxygen..... Lime. - Chloride of Calcium. Calcium....-AMMONIÆ SESQUICARBONAS. 11 Carbonic Acid.... = 33 1 Ammonia = 17 Composition. 1 Water.... = 959 Equivalent. B 2

Hydrogen of Ammonia. Carbonate Carbonate Of Lime. Carbonic Acid Carbonic Acid Oxygen Calcium Chloride of Calcium. LIQUOR AMMONLÆ ACETATIS.
of Lime. Calcium Chloride of Calcium. LIQUOR AMMONIÆ ACETATIS.
LIQUOR AMMONIÆ ACETATIS.
Securicant of Carbonia Arid Office Car
Sesquicarb. of y Carbonic AcidOff as Gas.
Ammonia. (Ammonia Distilled VinegarAcetate of Ammonia.
MORPHIÆ HYDROCHLORAS.
Composition. $\begin{cases} 1 \text{ Hydrochloric Acid} = 37 \\ 1 \text{ Morphia} = 284 \end{cases}$
321 Equivalent.
Solution of Morphia Hydrochlorate of Morphia.
Water J Hydrogen
Chloride of (Chlorine
Lead. Lead Meconate of Lead.
' MORPHIA.
Before decomposition. After decomposition.
Hydrochlorate Morphia Morphia Precip.
Ammonia
QUINÆ DISULPHAS.
(2 Quina = 324
Composition. $\begin{cases} 1 \text{ Sulphuric Acid} = 40 \\ 8 \text{ Water} = 72 \end{cases}$
And a second s
436 Equivalent.
A. Acidulated Decoc- J Kinate of Quina In solution.
tion of Cinchona. Sulphuric Acid Sulphate of Lead.
B. Ammonia Kinate of Ammonia.
Kinate of Kinic Acid
Quina. Quina
Sulpharte Acia
STRYCHNIA.
A. Magnesia Strychnate of Magnesia.
Nux Vomica. {Strychnic Acid
Sulphuric AcidSulphate of Strychnia.





OF THE LONDON PHARMACOPCEIA.

B. Ammonia		Sulphate of Ammonia.
Sulphate of Strychnia.	Sulphuric Acid	
Su yemma.	(Suyennia	Strychnia precipitated.

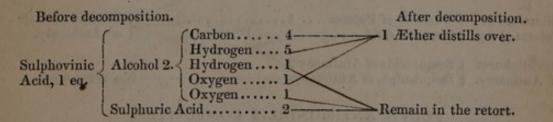
VERATRIA.

Gallate of Ve- { Gallic Acid	Super gallate of Veratria.
Sulphuric Acid	Sulphate of Veratria.
B. Magnesia Magnesia Super gallate { Gallic Acid Suphate { Sulphuric Acid	
of Veratria. (Veratria	Sulphate of Veratria.
C. Ammonia Sulphate of { Sulphuric Acid Veratria. { Veratria	Sulphate of Ammonia.

ÆTHER SULPHURICUS.

Composition.
$$\begin{cases} 4 \text{ Carbon} \dots = 24 \\ 1 \text{ Oxygen} \dots = 3 \\ 5 \text{ Hydrogen} \dots = 5 \end{cases}$$

37 Equivalent. .



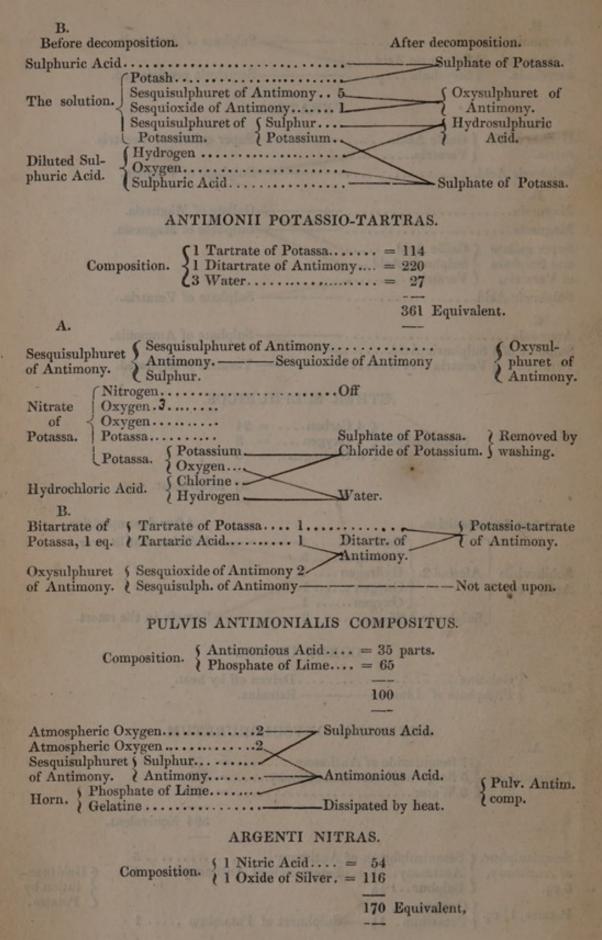
CORNU USTUM.

ANTIMONII OXYSULPHURETUM.

Composition.	3 5	Sesquioxide of Antimony Sesquisulphuret of Antimony	=	445
composition		Water		

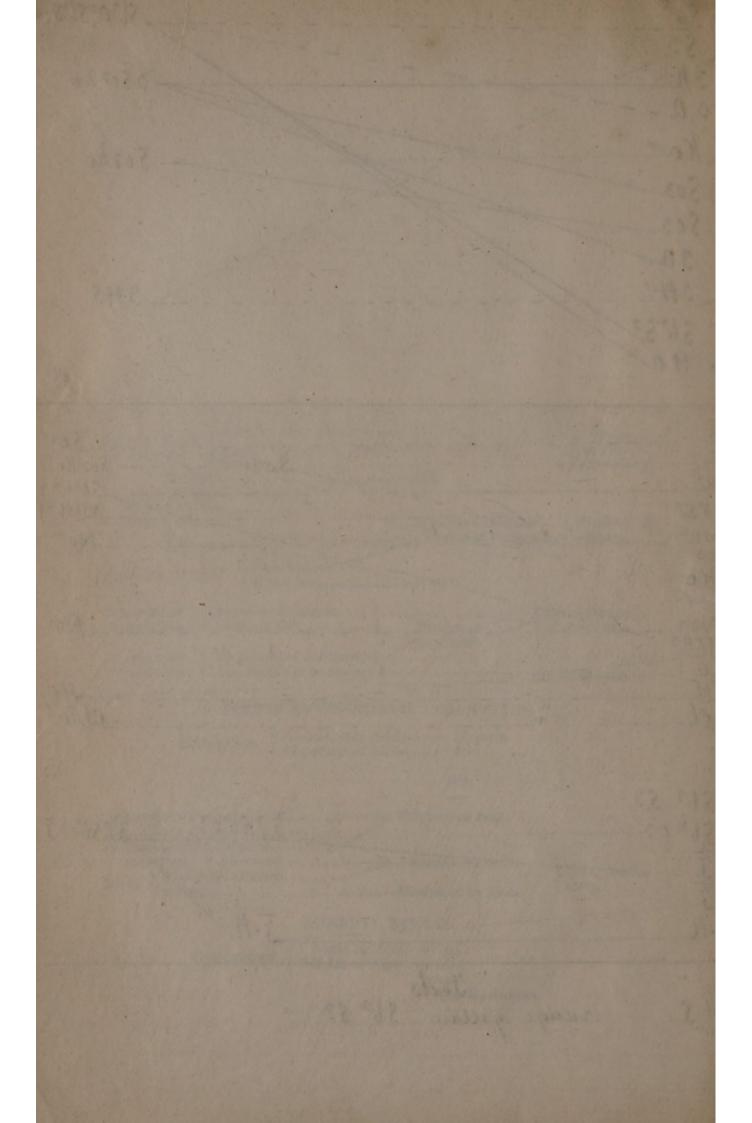
594 Equivalent.

Sesquisulphur.	Sesquisulphuret of Antimony 5	
of Antimony,	Antimony 1Sesquioxide of Antimony 1	(Heldinso-
6 eq.	Sulphur 11	} lution by
Botasen Il or	Oxygen 11 Potassium. 11 Sulphuret of Potassium 1	C Potassa.
rocassa, 12 eq. 3	Potassium. 11-Sulphuret of Potassium 1	



6

oxyenephoner of antimony 5 56253 SHO 5625 562035,56253,840 53 35K. 3K 3503 Ko 30 S03 30 3H 3#5 25. - Soz 503 Ko 503 512 Sl203. 551562 5 5653 NO2-Noz Ko. No2 Noz 2 0-TH_ -HO CUK 1cl 50562 53 525603 2562 03-32.560 × 5/10 ĩ Ī 7.12 1.K Jests HS trange yellan Sbr S3



Atmospheric	Oxygen	
Nitric Acid,	Nitric Acid 1 Binoxide of Nitrogen 1 Oxygen	Simularity Constant These
Silver		

ARGENTI CYANIDUM.

Composition. $\begin{cases} 1 \text{ Cyanogen} = 26 \\ 1 \text{ Silver} \dots = 108 \end{cases}$

134 Equivalent.

Before dec	omposition.	After decompositio
Nitrate of silver, 1 eq.	Silver 1	Free.
Hydrocyanic Acid, 1 eq.	§ Hydrogen 1 Cyanogen 1	Cyanide of Silver.

LIQUOR POTASSÆ ARSENITIS.

Composition of Arsenious Acid. $\begin{array}{c} 2 & \text{Arsenicum} = 76 \\ 3 & \text{Oxygen} \\ \end{array}$

100 Equivalent.

n.

BARII CHLORIDUM.

Composition. $\begin{cases} 1 \text{ Chlorine} = 36 \\ 1 \text{ Barium} = 68 \end{cases}$

104 Equivalent.

Carbonate of Baryta, 1 eq. Carbonic Acid.. 1.....Off as gas. Oxygen Hydrochloric Acid, 1 eq. Carbonic Acid.. 1....Off as gas. Water. Water. Chlorine....... Chloride of Barium.

A.

BISMUTHI TRISNITRAS.

Composition. $\begin{cases} 1 \text{ Nitric Acid } \dots = 54 \\ 3 \text{ Oxide of Bismuth} = 240 \end{cases}$

294 Equivalent.

Before dec	omposition.		After dec	composition.	
Nitric Acid,	Nitric Acid 1	Binoxide of Nitrogen Oxygen	3	1 Nitrous Acid Gas.	1
Bismuth			. 3	3 Nitrate of Bis- muth.	

. 7

D.				
Nitrate of	Oxide of Bismuth	3	-Trisnitrate of Bismuth (precipitate	ed).
Bismuth,	Nitric Acid	1	and the second se	
3 eqs.	Nitric Acid	2		
Water	The second se	The second	-Unite	

CALX.

Composition.
$$\begin{cases} 1 \text{ Oxygen } \dots = 8\\ 1 \text{ Calcium } \dots = 20 \end{cases}$$

28 Equivalent.

Chalk {Carbonic Acid 1.....Driven off by heat. Lime 1————Remains.

CALCII CHLORIDUM.

Composition. $\begin{cases} 1 \text{ Chlorine } = 36\\ 1 \text{ Calcium } = 20 \end{cases}$

56 Equivalent.

Before decomposition.

Co

Sulphate of (O

After decomposition.

Carbonate of Lime, 1 eq.	Carbonic Acid 11 Off as gas. Oxygen1 Calcium1	
Hydrochloric Acid, 1 eq.	Hydrogen 1 Chlorine 11 Chloride of Cal	cium.

CALX CHLORINATA.

Composition. $\begin{cases} 1 \text{ Chlorine } \dots = 36 \\ 2 \text{ Hydrate of Lime} = 74 \end{cases}$

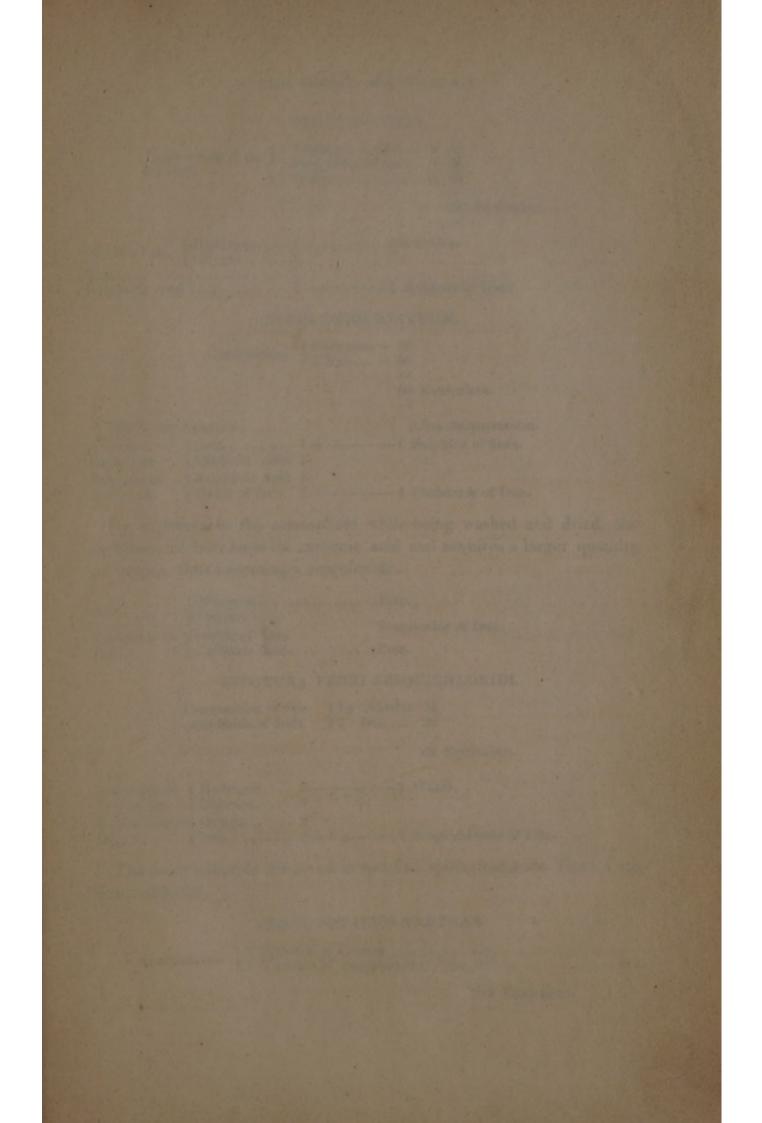
110 Equivalent.

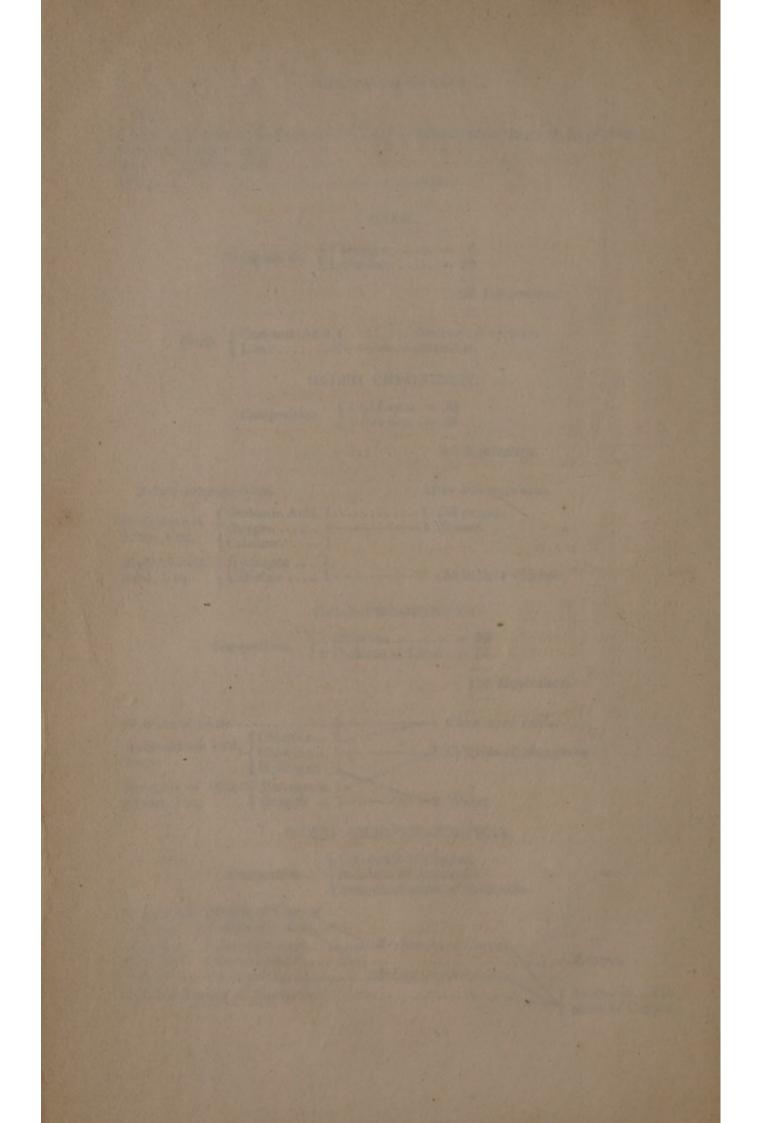
CUPRI AMMONIO-SULPHAS.

	Carbonate of Copper. Sulphate of Ammonia. Sesquicarbonate of Ammonia.
of Connor	

Surburnes or 1	our of copper
Copper. 2	Sulphuric Acid
Sesquicar-	Carbonic AcidCarbonate of Copper
bonate of -	Carbonic Acid Escapes.
Ammonia.	AmmoniaSulphate of Ammonia
Sesquicarbon	ate of Ammonia S Ammonio Sul-
Tanta	phate of Copper.

8





OF THE LONDON PHARMACOPCEIA.

FERRI SULPHAS.

Composition of the Crystals.	Sulphuric Acid Protoxide of Iron Water		
	water	-	00

139 Equivalent.

Water, 1 eq. {Hydrogen 1 Oxygen 1	Off as Gas.
Iron 1 Sulphuric Acid 1	

FERRI SESQUIOXYDUM.

Composition. $\begin{cases} 3 \text{ Oxygen} = 24\\ 2 \text{ Iron} \dots = 56 \end{cases}$

80 Equivalent.

Before decon	nposition.	After decomposition.
	Soda Carbonic Acid	Sulphate of Soda.
	Sulphuric Acid Oxide of Iron	Carbonate of Iron.

By exposure to the atmosphere while being washed and dried, the carbonate of iron loses its carbonic acid and acquires a larger quantity of oxygen, thus becoming a sesquioxide.

Air.	NitrogenFree.	
	Oxygen Sesquioxide of Iron.	
Iron.	Carbonic AcidFree.	

TINCTURA FERRI SESQUICHLORIDI.

Composition of Sesquichloride of Iron. {11/2 Chlorine 54 11 Iron.... 28

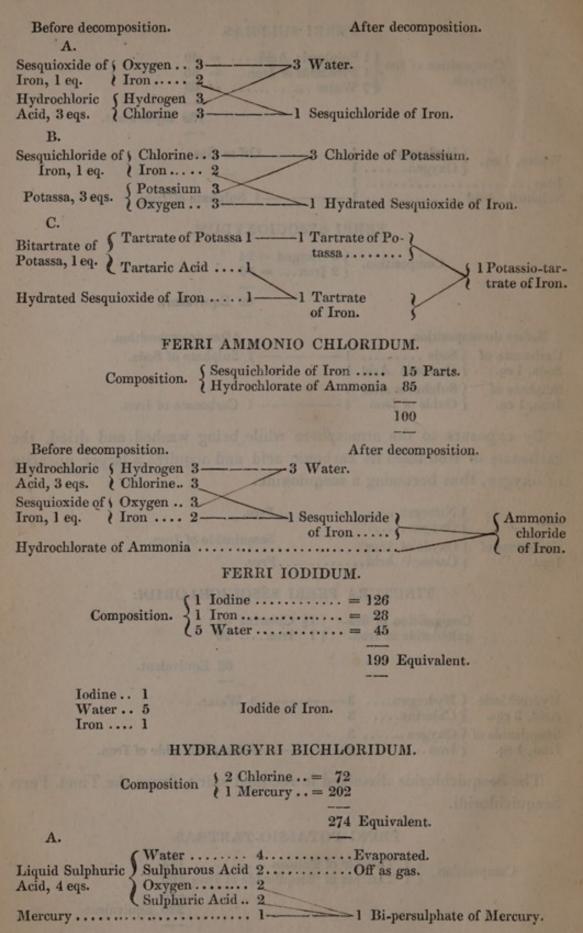
82 Equivalent.

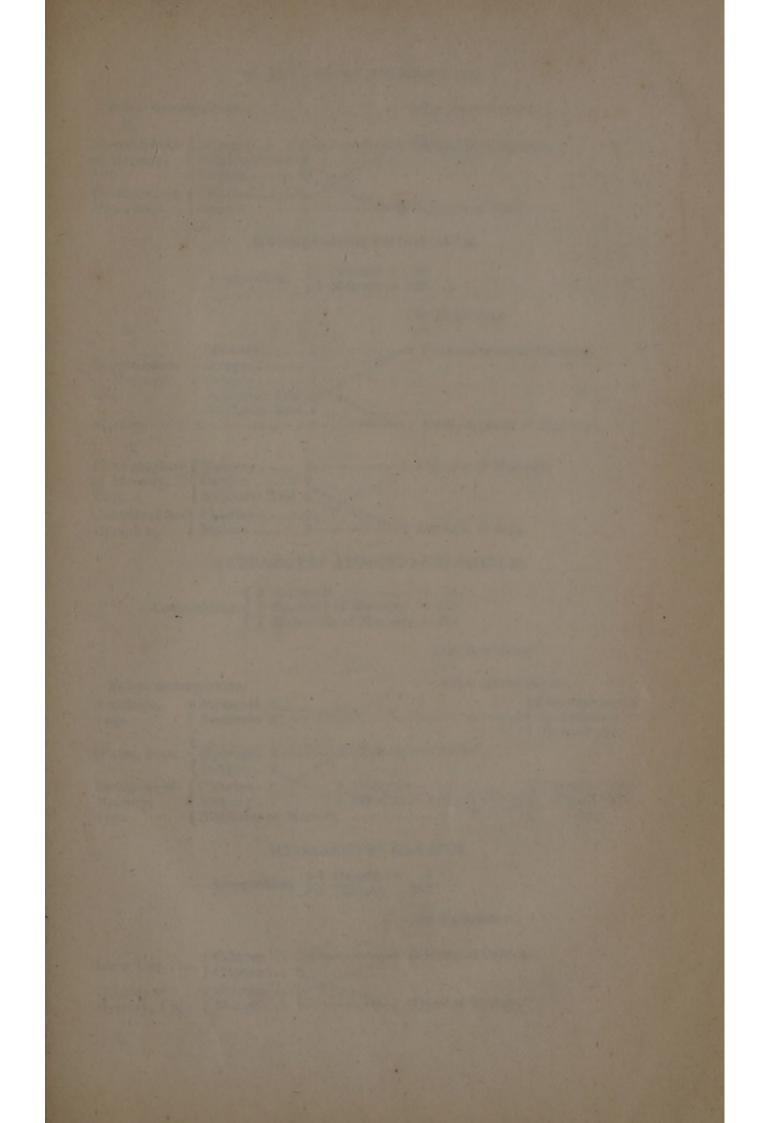
The Sesquichloride dissolved in rectified spirit, forms the Tinct. Ferri Sesquichloridi.

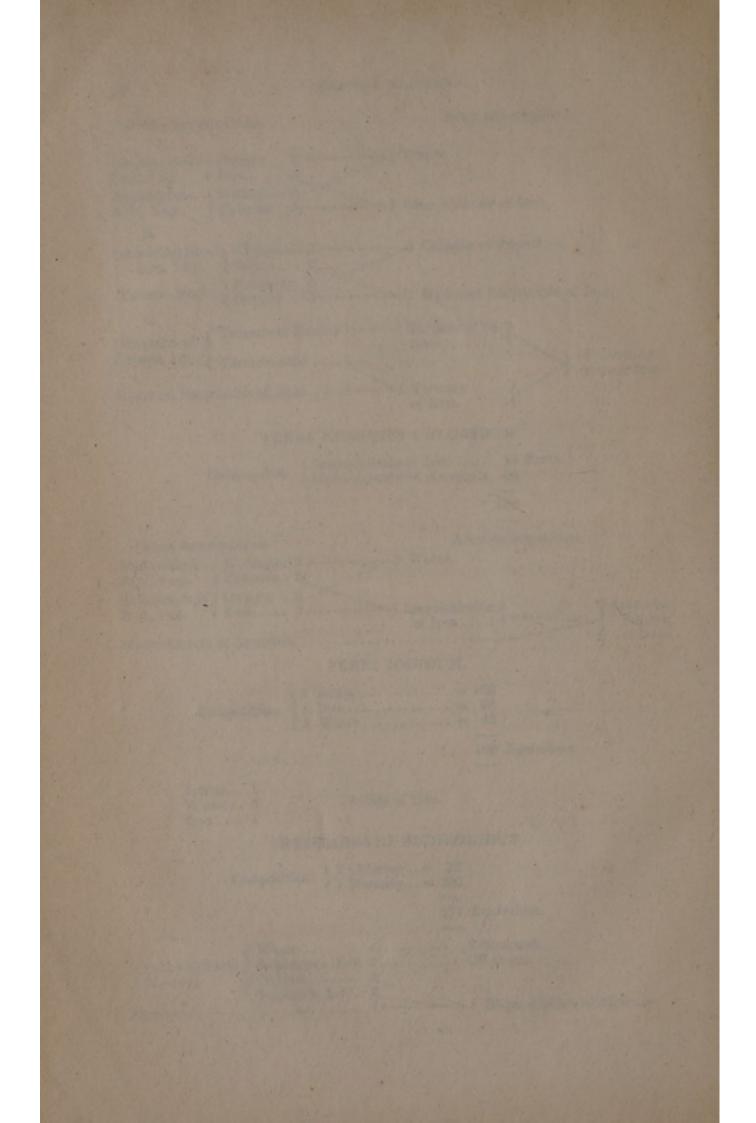
FERRI POTASSIO-TARTRAS.

Composition. { 1 Tartrate of Potassa 114 1 Tartrate of Sesquioxide of Jron 106

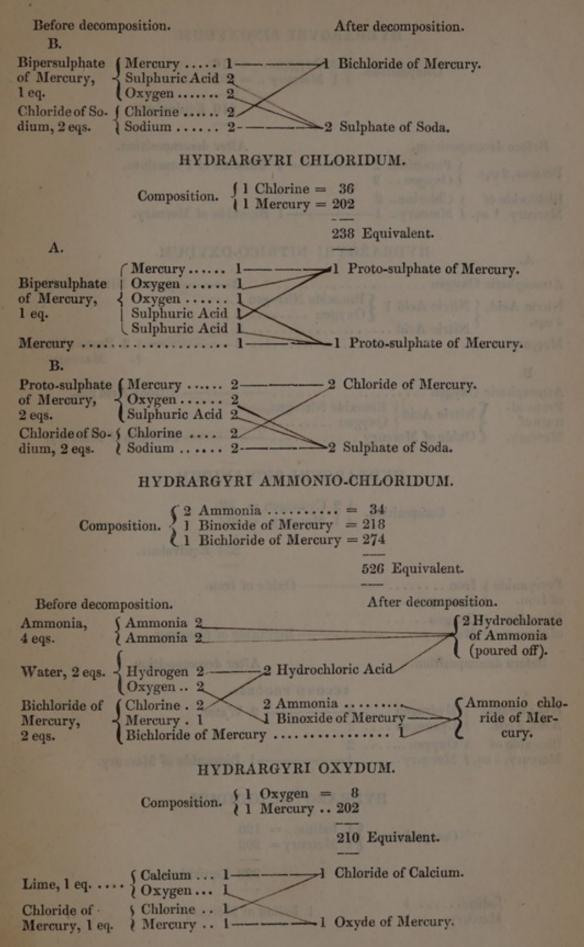
220 Equivalent.







OF THE LONDON PHARMACOPCEIA.



11

HYDRARGYRI BINOXYDUM.

Composition. $\begin{cases} 2 \text{ Oxygen..} = 16 \\ 1 \text{ Mercury.} = 202 \end{cases}$

218 Equivalent.

Before decom	position.		After decomposition.
Potassa, 2 eqs.	{ Potassium Oxygen	2- <u></u> 2 2	Chloride of Potassium.
Bichloride of Mercury, 1 eq.	Chlorine Mercury		Binoxide of Mercury.

HYDRARGYRI NITRICO-OXYDUM.

Nitric Acid, §	Dxygen 2 Nitric Acid 1 Binoxide Nitrogen 1 Oxygen 3 Nitric Acid 3	the representation of the second second
Mercury		3 Proto-nitrate of Mercury.
B. Atmospheric (Oxygen	
Proto-ni- trate of	Nitric Acid Binoxide Nitrogen	alan (Angelenger Sain

Mercury. (Oxide of Mercury Binoxide of Mercury.

HYDRARGYRI BICYANIDUM.

Composition. § 2 Cyanogen = 52 (1 Mercury. = 202

254 Equivalent.

Iron Cyanogen	Oxide of Iron.
Oxygen Mercury	Bicyanide of Mercury.

Before decomposition.

After decomposition.

SECOND PROCESS.

	Hydrogen		Water.
Acid, 2 eqs.	Cyanogen	2	I start have been the
Binoxide of	Oxygen	2	
Mercury, 1 eq.	Mercury	11	Bicyanide of Mercury.

HYDRARGYRI IODIDUM.

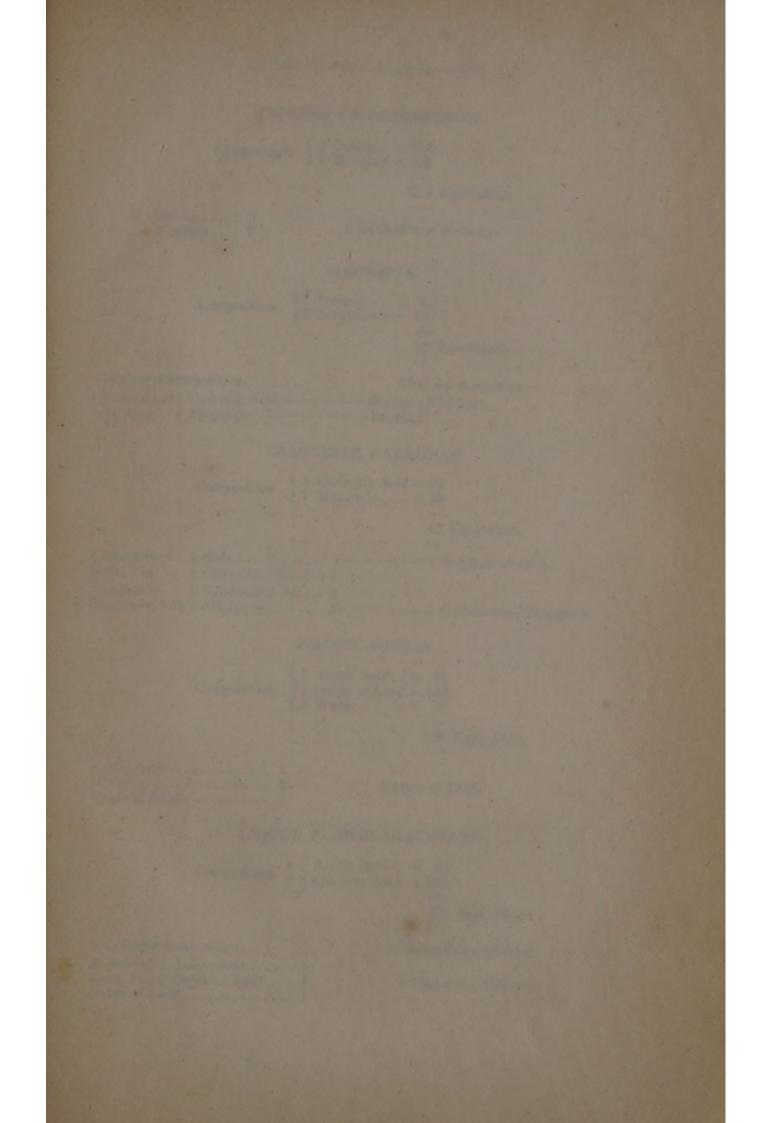
Composition. $\begin{cases} 1 \text{ Iodine..} = 126 \\ 1 \text{ Mercury} = 202 \end{cases}$

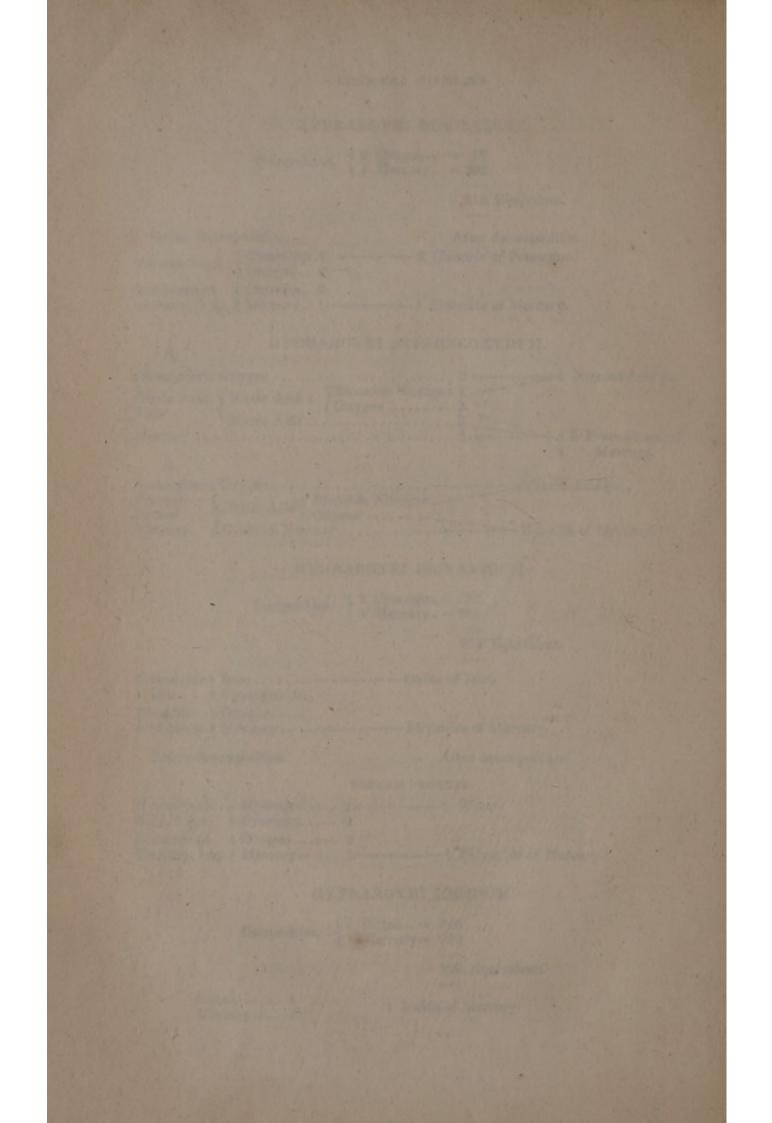
328 Equivalent.

 Iodine..... 1
 1 Iodide of Mercury.

 Mercury.... 1
 1

A.





HYDRARGYRI BINIODIDUM.

Composition.

§ 2 Iodine.. = 252 1 Mercury = 202

454 Equivalent.

Iodine 2 Mercury.... 1 *

1 Biniodide of Mercury.

MAGNESIA.

1 Oxygen ... = 8

Composition.

1 Magnesium = 1220 Equivalent.

Before decomposition. After decomposition. _____Driven off by heat. _____Remains. Carbonate of) Carbonic Acid-Magnesia A Magnesia-

MAGNESIÆ CARBONAS.

1 Carbonic Acid = 22 Composition. 1 Magnesia = 20

42 Equivalent.

Carbonate of	§ Soda	1
Soda, 1 eq.	(Carbonic Acid	La sharmont + Westmannet
Sulphate of	Sulphuric Acid	Inter sector of the sector of the sector of
Magnesia, 1 eq.	A Magnesia	1

PLUMBI ACETAS.

Composition.	(1	Acetic Acid	=	51	
	31	Oxide of Lead	=	112	
	63	Water	-	27	

190 Equivalent.

Acetic Acid 1	
Water 3	Acetate
Oxide of Lead 1	

e of Lead.

LIQUOR PLUMBI DIACETATIS.

1 Acetic Acid ... = 51Composition. 2 Oxide of Lead = 224

275 Equivalent.

Before decomposition. After decomposition. Acetate of & Acetic Acid 1 Lead, 1 eq. / Oxide of Lead 1 1 Diacetate of Lead. Oxide of Lead 1

PLUMBI CHLORIDUM.

Composition. $\begin{cases} 1 \text{ Chlorine} = 36 \\ 1 \text{ Lead} \dots = 104 \end{cases}$

140 Equivalent.

Chloride of Sodium, 1 eq. Sodium...... 1 — 1 Acetate of Soda. Chlorine 1 Acetate of Lead, 1 eq. Acetate of Lead. Lead. 1 eq. Lead 1

PLUMBI IODIDUM.

Composition. $\begin{cases} 1 \text{ Iodine} = 126 \\ 1 \text{ Lead.} = 104 \end{cases}$

230 Equivalent.

Before decomp	osition.	Af	ter decomposition.
	Potassium		Acetate of Potassa.
Lend Leg	Acetic Acid Oxygen Lead	1 amount I	Iodide of Lead.

PLUMBI OXYDUM HYDRATUM.

Composition. Protoxide of Lead. Water, (proportion unknown).

and the second	Potassa Acetate of Potassa. Water
	Acetic Acid Oxide of Lead — Hydrated oxide of Lead.

POTASSÆ CARBONAS.

Composition. $\begin{cases} 1 & \text{Carbonic Acid } 22 \\ 1 & \text{Potassa} \dots 48 \\ 1\frac{1}{2} & \text{Water} \dots 12 \end{cases}$

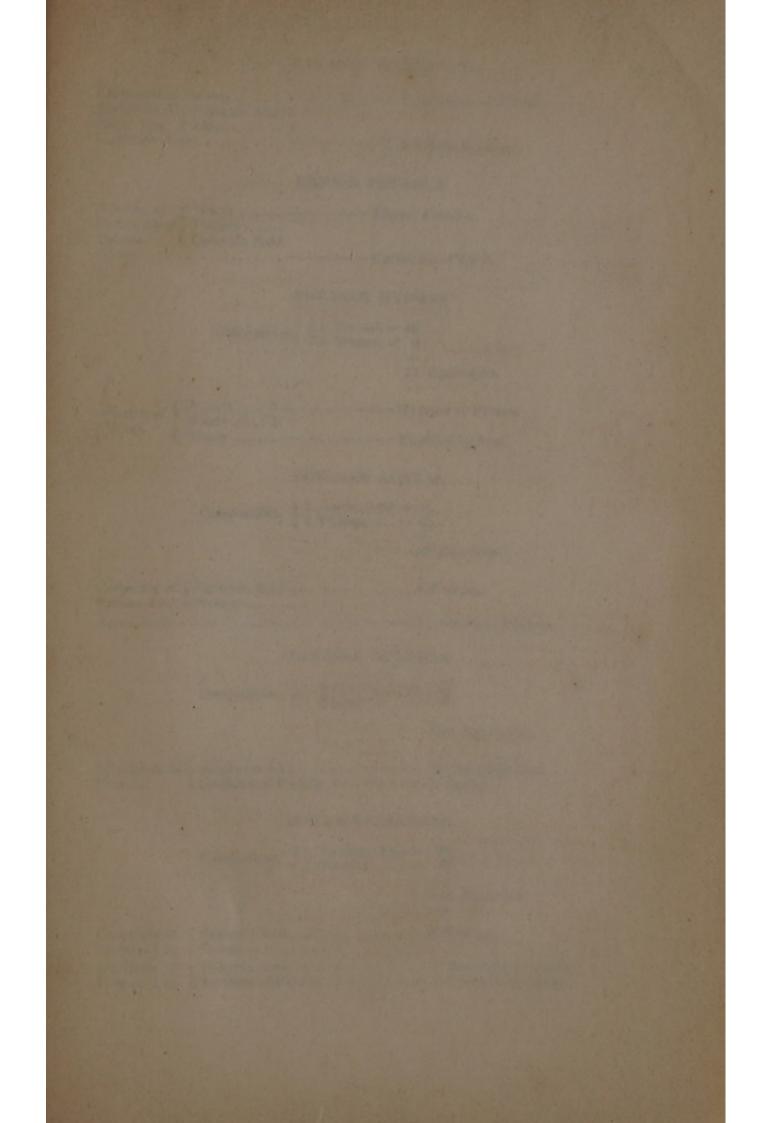
82 Equivalent.

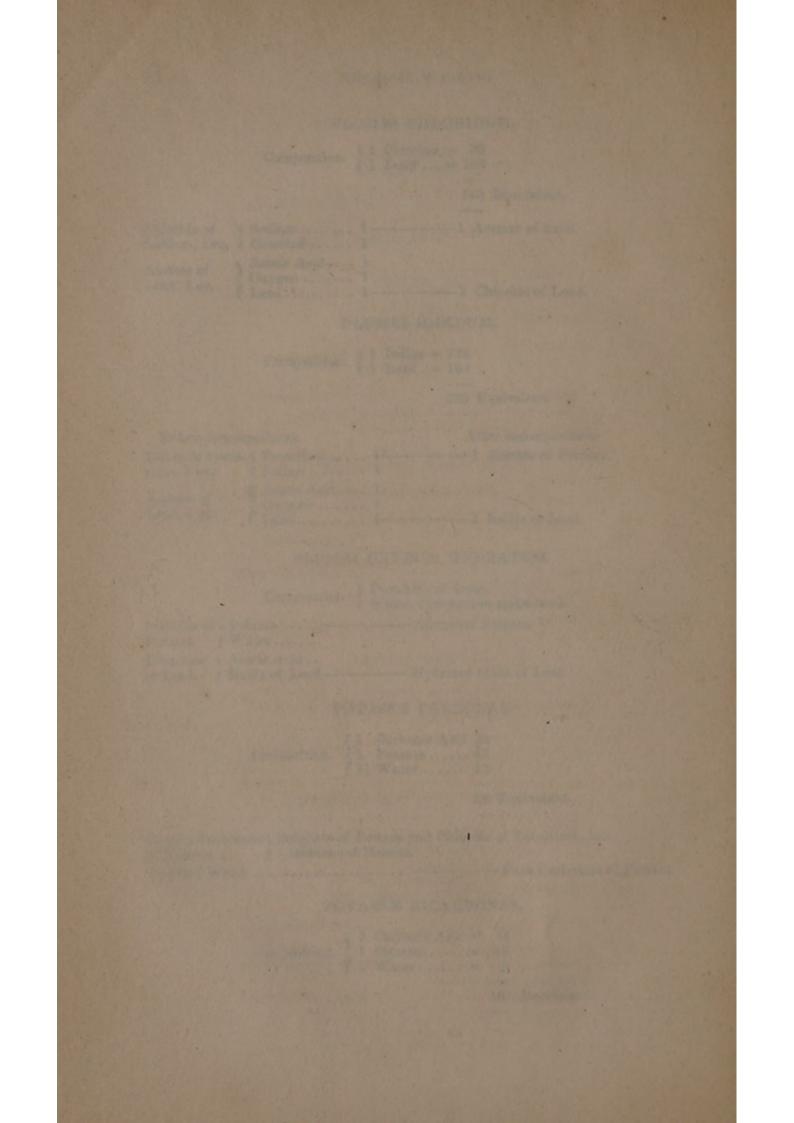
Impure Carbonate | Sulphate of Potassa and Chloride of Potassium, left. of Potassa. | Carbonate of Potassa. Distilled WaterPure Carbonate of Potassa.

POTASSÆ BICARBONAS.

Composition. $\begin{cases} 2 \text{ Carbonic Acid} = 44\\ 1 \text{ Potassa} \dots = 48\\ 1 \text{ Water} \dots = 9 \end{cases}$

101 Equivalent.





Carbonate of) Carbonic Acid	1. MILMORAL 1	Bicarbonate of Potassa.
Lime, 1 eq. ? Lime Sulphuric Acid		Sulphate of Lime.

LIQUOR POTASSÆ.

Solution of Carbonate of Potassa. LimeCarbonic Acid LimeCarbonate of Lime.

POTASSÆ HYDRAS."

Composition. $\begin{cases} 1 & Potassa = 48 \\ 1 & Water = 9 \\ 57 & Equivalent. \end{cases}$

Solution of Potassa.... 1 — Hydrate of Potassa. Water 1 — Expelled by heat.

POTASSÆ ACETAS.

Composition. $\begin{cases} 1 & \text{Acetic Acid} = 51 \\ 1 & \text{Potassa} \dots = 48 \end{cases}$

99 Equivalent.

POTASSÆ SULPHAS.

Composition. $\begin{cases} 1 & \text{Sulphuric Acid} = 40 \\ 1 & \text{Potassa} \dots = 48 \end{cases}$

88 Equivalent.

Bi-sulphate of Sulphuric Acid 1 — Driven off by heat. Potassa. Sulphate of Potassa 1 — Remains.

POTASSÆ TARTRAS.

Composition. $\begin{cases} 1 & \text{Tartaric Acid} = 66 \\ 1 & \text{Potassa} \dots = 48 \end{cases}$

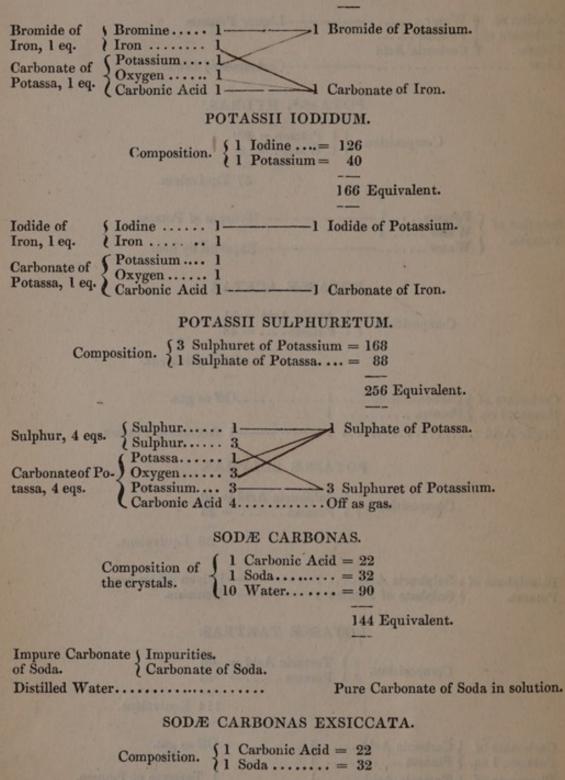
114 Equivalent.

Carbonate of	Carbonic Acid	1Off as gas.
	? Potassa	
Bitartrate of	[Tartaric Acid	1 1 Tartrate of Potass
Potassa, 1 eq.	Tartrate of Potassa	11 Tartrate of Potass

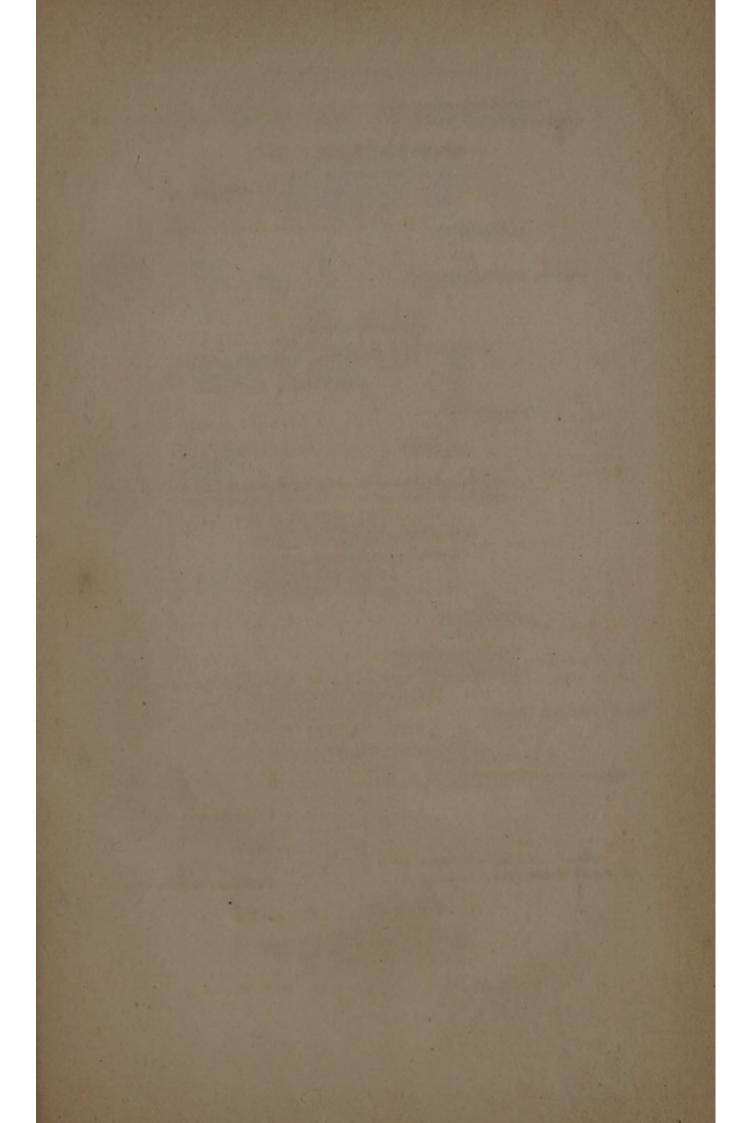
POTASSII BROMIDUM.

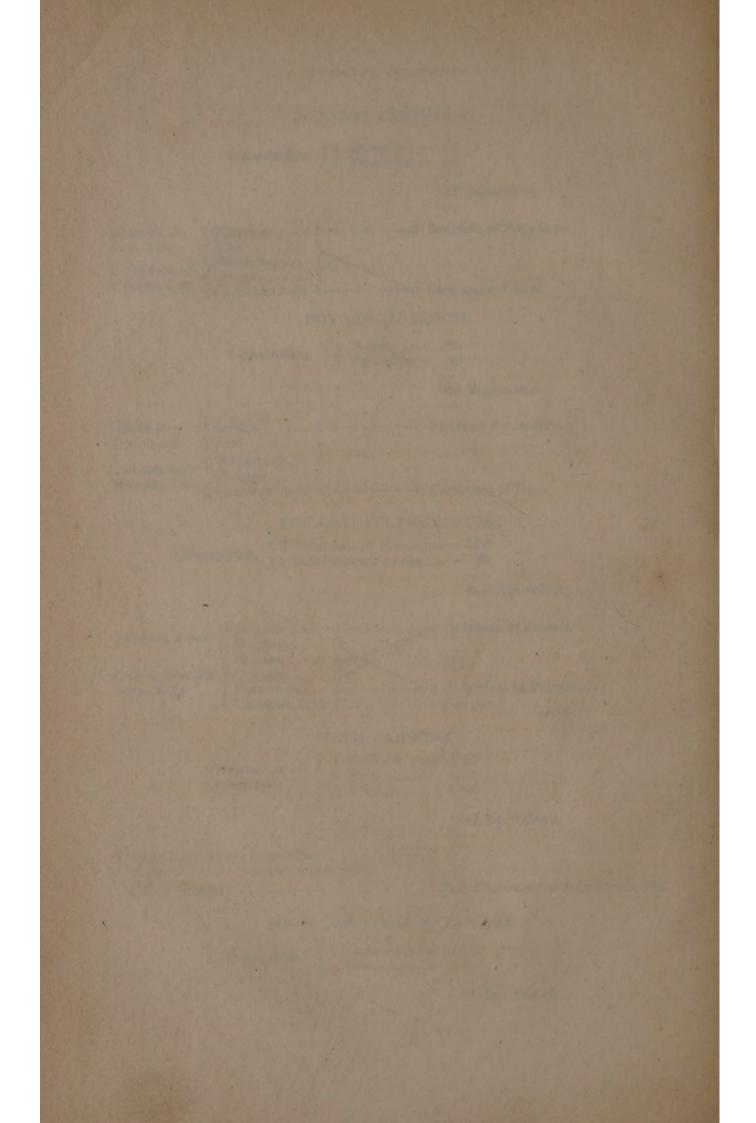
Composition. $\begin{cases} 1 \text{ Bromine } . = 78 \\ 1 \text{ Potassium } = 40 \end{cases}$

118 Equivalent.



54 Equivalents.





Carbonate of Soda. {Water-Expelled by heat. Carbonate of Soda ______ Dry Carbonate of Soda.

SODÆ SESQUICARBONAS.

Composition. $\begin{cases} 1\frac{1}{2} & \text{Carbonic Acid} = 33\\ 1 & \text{Soda} \dots = 32\\ 2 & \text{Water} \dots = 18 \end{cases}$

83 Equivalent.

Carbonate of {Soda..... 1 Soda, 1 eq. {Carbonic Acid.... 1 Carbonic Acid..... 1

1 Sesquicarbonate of Soda.

SODÆ SULPHAS.

Composition of	(1	Sulphuric Acid =	40
	31	Soda =	32
the Crystals.	(10	Water =	90

162 Equivalent.

	Carbonic Acid ———————————————————————————————————
Super-sulphate	Sulphuric AcidSulphate of Soda. Sulphate of SodaSulphate of Soda.

SODÆ POTASSIO-TARTRAS.

Composition. $\begin{cases} 1 \text{ Tartrate of Potassa} = 114 \\ 1 \text{ Tartrate of Soda} \dots = 98 \\ 8 \text{ Water} \dots \dots = 72 \end{cases}$

284 Equivalent.

Before decomposit	ion.	After decomposition.
	Carbonic Acid	1Off as gas.
Bitartrate of Po-	Tartaric Acid Tartrate of Potassa.	1 1 Potassio-Tartrate of Soda.

LIQUOR SODÆ CHLORINATÆ.

Solution of Carbon	ate of Soda	L	iquor Sodæ Chlorinatæ,
Chloride of So-	Chlorine	L	
dium, 1 eq.	Sodium	L	
Binoxide of Man- ganese, 1 eq.	Oxygen Protoxide Manganese.	1	
Sulphuric Acid		11	Sulphate of Soda.
Sulphuric Acid		1	Sulphate of Manganese.

ZINCI SULPHAS.

Composition. $\begin{cases} 1 & \text{Sulphuric Acid} = 40 \\ 1 & \text{Oxide of Zinc.} = 40 \\ 7 & \text{Water.} = 63 \end{cases}$

143 Equivalent.

C

17

Before decomposition.	After decomposition.
Water, 1 eq. {Hydrogen Oxygen	1Off as gas.
Zinc	

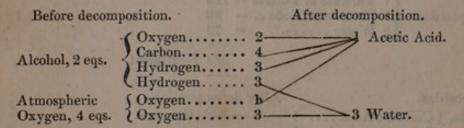
ZINCI OXYDUM.

Composition. $\begin{cases} 1 & \text{Zinc...} & 32 \\ 1 & \text{Oxygen} & 8 \\ \hline 40 & \text{Equivalent.} \end{cases}$

A		
	f Ammonia	
of Ammonia.	Carbonic Acid	
Sulphate of	Sulphuric Acid	an ord a summary but
Zinc.	Oxide of Zinc	Carbonate of Zinc.
В.		
and the second se	Carbonic Acid	Expelled by heat.
	Oxide of Zinc	

ACETOUS FERMENTATION.

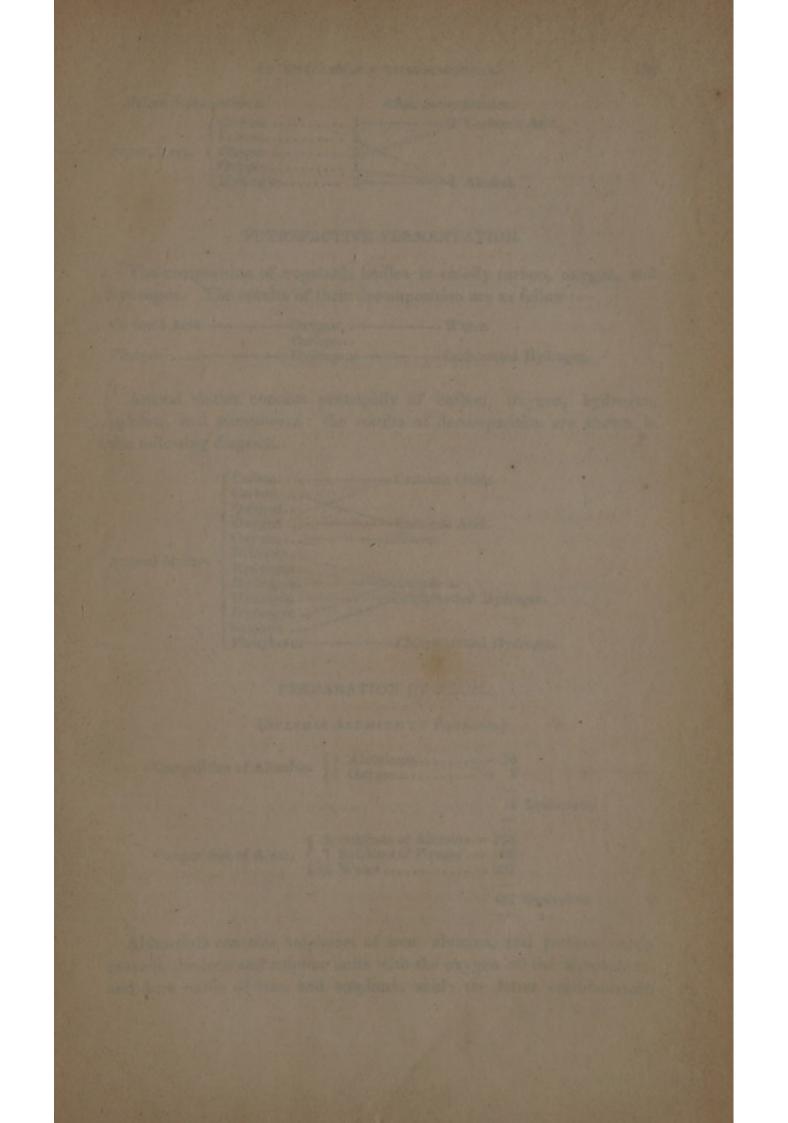
When alcohol, mixed with yeast, is exposed to the air in a warm temperature, the following changes take place.

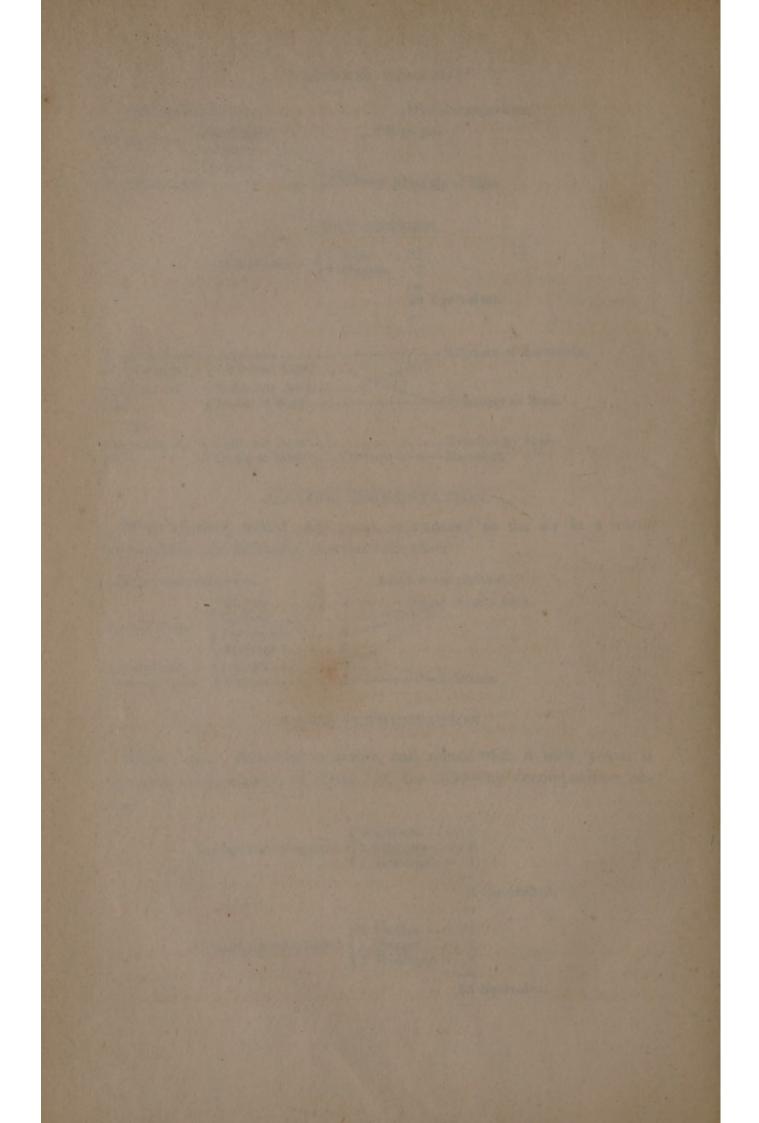


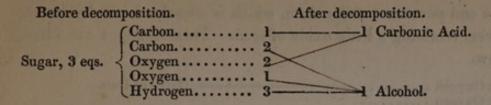
VINOUS FERMENTATION.

When sugar, dissolved in water, and mixed with a little yeast, is placed in a temperature of about 70°, the following decomposition occurs.

Composition of Sugar.
$$\begin{cases} 1 \text{ Carbon. ..} = 6\\ 1 \text{ Oxygen..} = 8\\ 1 \text{ Hydrogen} = 1\\ 15 \text{ Equivalent.} \end{cases}$$
Composition of Alcohol.
$$\begin{cases} 2 \text{ Carbon. ..} = 12\\ 1 \text{ Oxygen..} = 8\\ 3 \text{ Hydrogen} = 3\\ 23 \text{ Equivalent.} \end{cases}$$



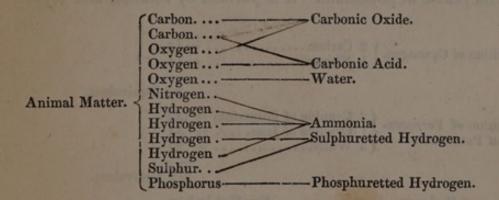




PUTREFACTIVE FERMENTATION.

Carbonic Acid	Oxygen	
	Carbon	
Vinegar		Carburetted Hydrogen.

Animal matter consists principally of carbon, oxygen, hydrogen, sulphur, and phosphorus: the results of decomposition are shown in the following diagram.



PREPARATION OF ALUM.

(SULPHAS ALUMINÆ ET POTASSÆ.)

Composition of Alumina. $\begin{cases} 1 & \text{Aluminum} \dots = 10 \\ 1 & \text{Oxygen} \dots = 8 \end{cases}$

18 Equivalent.

Composition of Alum. $\begin{cases} 3 \text{ Sulphate of Alumina} = 174 \\ 1 \text{ Sulphate of Potassa} . = 88 \\ 25 \text{ Water} ... = 225 \\ \hline 487 \text{ Equivalent.} \end{cases}$

Alum-slate contains sulphuret of iron, alumina, and potassa; when roasted, the iron and sulphur unite with the oxygen of the atmosphere, and form oxide of iron and sulphuric acid; the latter combines with

19

c 2

the alumina and potassa to form alum, which is dissolved out by water, still, however, retaining a little oxide of iron. These changes are illustrated below.

Atmospheric Oxygen	Oxide of Iron.
Alum-slate. Sulphuret { Iron of Iron. { Sulphur Alumina Potassa	

Frequently the alum-slate does not contain sufficient potassa; in this case a salt of potassa, either chloride of potassium, or sulphate of potassa, is added.

PREPARATION OF FERROCYANIDE OF POTASSIUM.

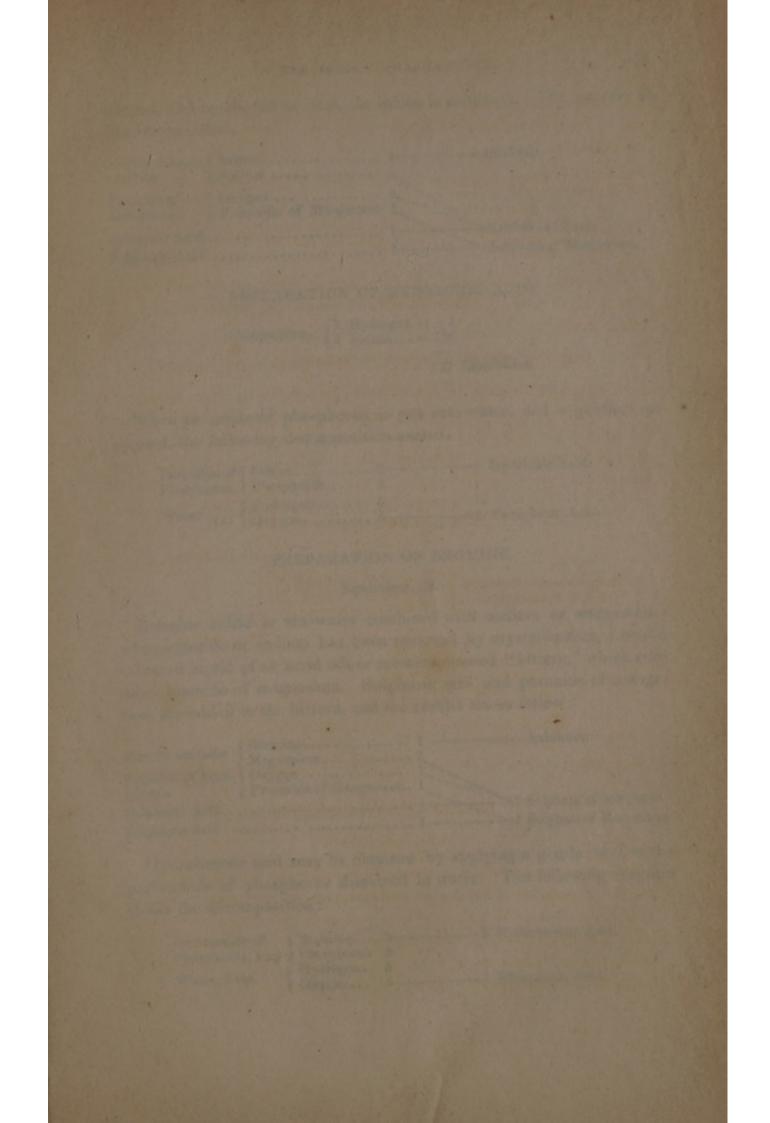
When animal matter is heated with potassa in an iron retort, the carbon and nitrogen of the former unite to form cyanogen, which, combining with the potassium of the potassa and the iron of the retort, constitutes ferrocyanide of potassium. It is purified by solution and crystallization.

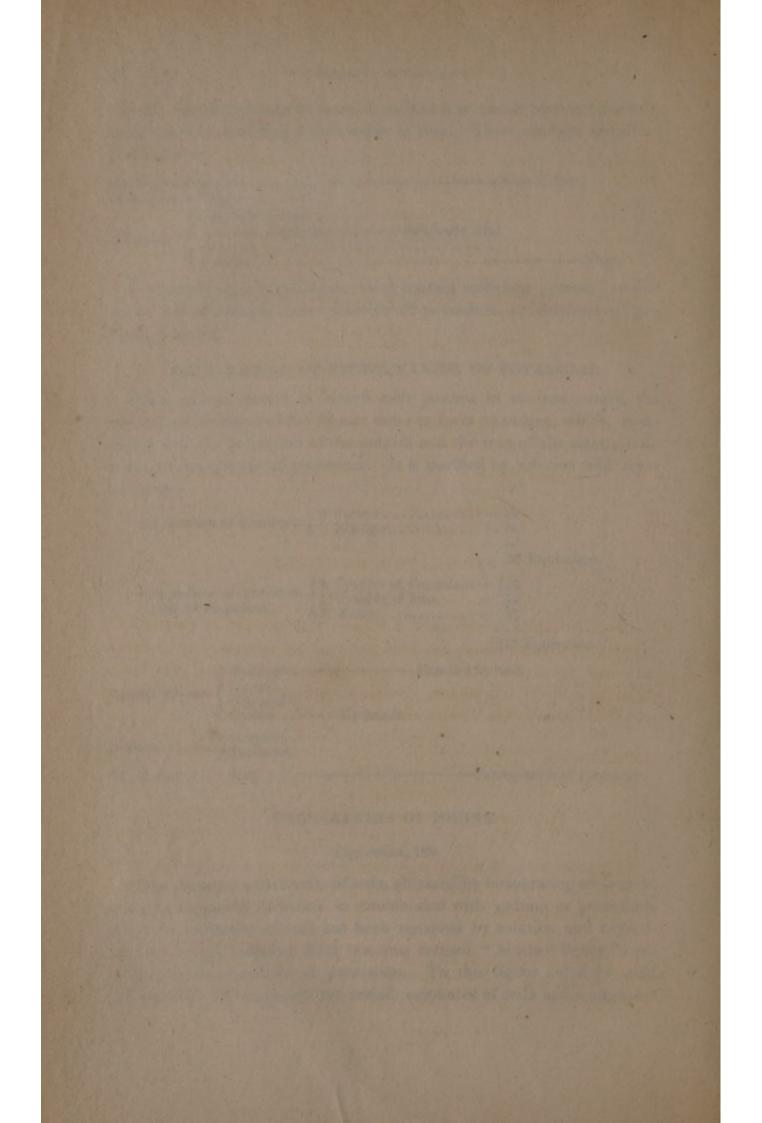
Composition of Cyanogen. 2 Carbon = 12 1 Nitrogen..... = 14 26 Equivalent. Composition of Ferrocya-nide of Potassium. $\begin{cases} 2 & \text{Cyanide of Potassium} = 132\\ 1 & \text{Cyanide of Iron.} \\ 3 & \text{Water.} \\ = 27 \end{cases}$ 213 Equivalent. - Expelled by heat. Hydrogen. Oxygen . Animal Matter. Nitrogen . Carbon .. Cyanogen Oxygen .. Potassa.... Potassium The Retort Ferrocyanide of Potassium. Iron. .

PREPARATION OF IODINE.

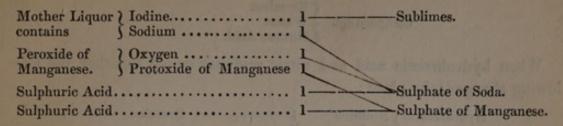
Equivalent, 126.

Kelp, the impure carbonate of soda, obtained by incinerating sea-weeds, contains a quantity of iodine, in combination with sodium or potassium. After the carbonate of soda has been removed by solution and crystallization, a dark coloured fluid remains, termed "Mother liquor," containing iodide of sodium or potassium. To this liquor sulphuric acid and peroxide of manganese are added, sulphates of soda and manganese





formed, and by the aid of heat, the iodine is sublimed. The process is illustrated below.



PREPARATION OF HYDRIODIC ACID.

Composition. $\begin{cases} 1 & \text{Hydrogen} = 1 \\ 1 & \text{Iodine...} = 126 \end{cases}$

127 Equivalent.

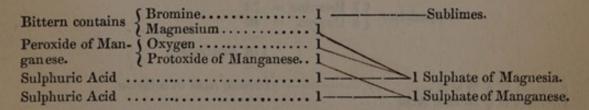
When periodide of phosphorus is put into water, and a gentle heat applied, the following decomposition ensues.

	Iodine Phosphorus		Hydriodic Acid.
Water }	Hydrogen Oxygen	5 51	Phosphoric Acid.

PREPARATION OF BROMINE.

Equivalent, 78.

Bromine exists in sea-water combined with sodium or magnesium; when chloride of sodium has been removed by crystallization, a brown coloured liquid of an acrid odour remains, named "bittern," which contains bromide of magnesium. Sulphuric acid and peroxide of manganese are added to the bittern, and the results are as follow:



Hydrobromic acid may be obtained by applying a gentle heat to the perbromide of phosphorus dissolved in water. The following diagram shows the decomposition :

Perbromide of	Bromine	5	Hydrobromic Acid.
Phosphorus, 1 eq.	Phosphorus	2	a ni benanice a
Water 5 ore	Hydrogen	5	dan bin hall
water, o eqs.	Oxygen	51	Phosphoric Acid.

PREPARATION OF NITRO-HYDROBROMIC ACID.

Composition. Bromine. Water. Nitrous Acid.

When hydrobromic acid and nitric acid are mixed together the following changes take place :

 Hydrobromic { Bromine
 1
 1
 Bromine.

 Acid, 1 eq.
 { Hydrogen ... 1
 1
 1

 Nitric Acid, { Oxygen 1
 1
 1
 Water.

 1 eq.
 { Nitrous Acid 1
 1
 1

The nitro-hydrobromic acid which is formed possesses the power of dissolving gold on account of the free bromine.

PREPARATION OF BROMATE OF BARYTA.

Composition. $\begin{cases} 1 \text{ Bromic Acid} = 118 \\ 1 \text{ Baryta} \dots = \frac{76}{76} \end{cases}$

194 Equivalent.

When bromine is added to a solution of any of the alkalies or alkaline earths the following decomposition takes place :

Baryta,	Barium	5	Bromide of Barium.
	Oxygen	5	
6 eqs.	Baryta	1 Mailavinade	
Bromine,	Bromine	5	
	Bromine		Bromate of Baryta.

The bromate can be easily separated from the bromide by evaporation and crystallization.

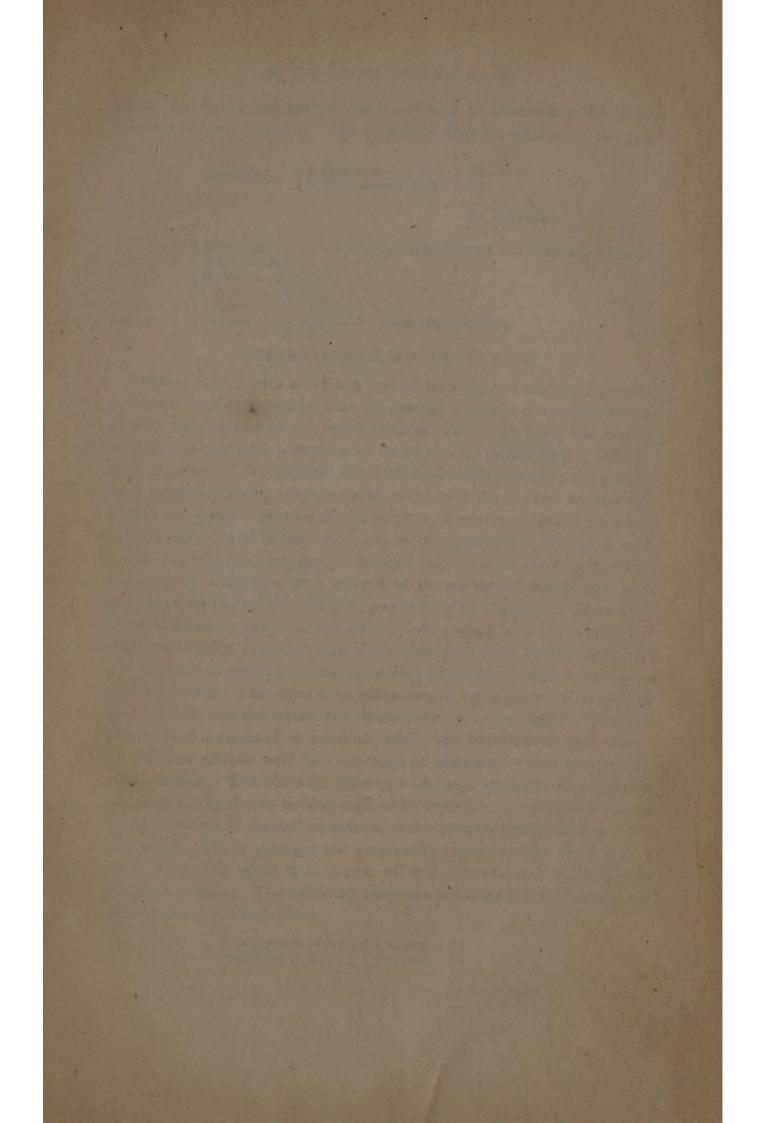
PREPARATION OF BROMIC ACID.

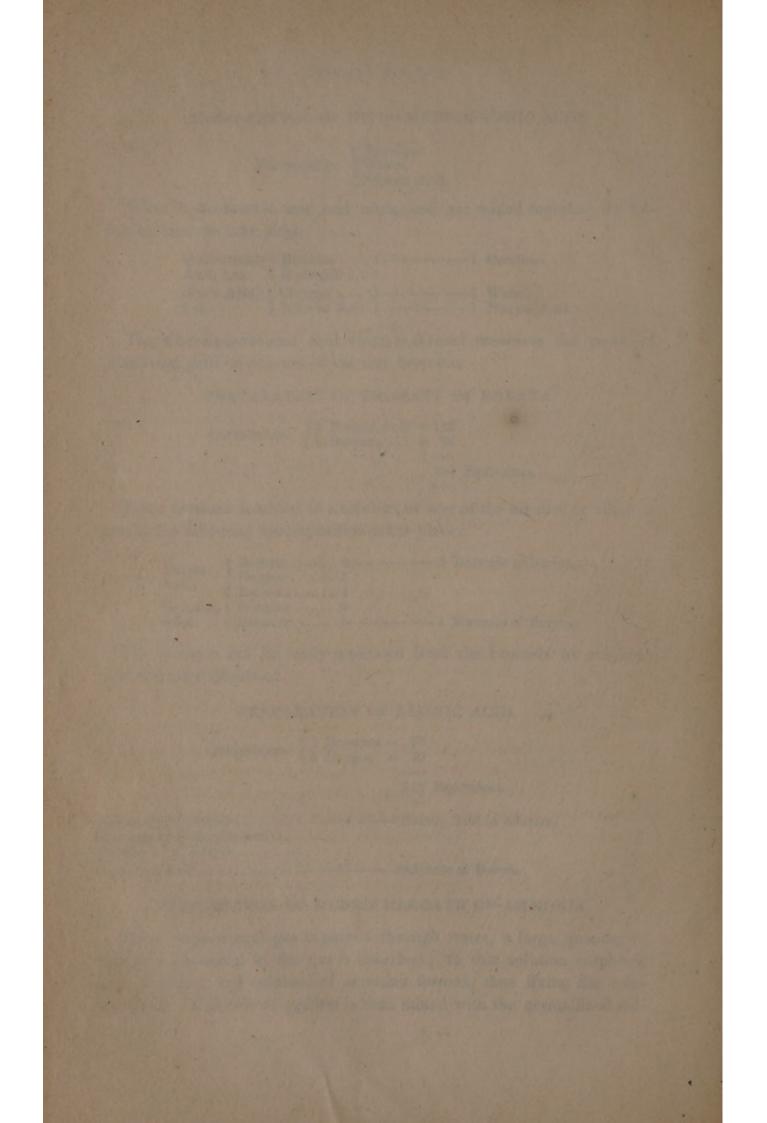
Composition. $\begin{cases} 1 & \text{Bromine} = 78 \\ 5 & \text{Oxygen} = 40 \end{cases}$

118 Equivalent.

PREPARATION OF HYDROCHLORATE OF AMMONIA.

When impure coal gas is passed through water, a large quantity of ammonia contained in the gas is absorbed; to this solution sulphuric acid is added, and sulphate of ammonia formed, thus fixing the volatile alkali. Chloride of sodium is then mixed with the crystallized sul-





phate, and heat being applied, hydrochlorate of ammonia sublimes and sulphate of soda remains. The following diagram illustrates the process:

Composition. $\begin{cases} 1 \text{ Ammonia } \dots = 17 \\ 1 \text{ Hydrochloric Acid.} = 37 \end{cases}$

54 Equivalent.

Sulphate of	Ammonia Hydrochlorate of Ammonia (sublimed).
Ammonia.	Sulphuric Acid
	Hydrogen
Sector Man	Oxygen
Chloride of	Chlorine
Sodium.	SodiumSulphate of Soda.

PREPARATION OF SULPHURIC ACID.

Sulphur, mixed with an eighth part of its weight of nitrate of potassa, is burned in such a position that the gaseous products of combustion are carried by a current of air into a chamber lined with lead, and having its floor covered with a stratum of water several inches deep. The nitrate of potassa is decomposed, the nitric acid yields part of its oxygen to a little sulphur, forming sulphuric acid, which with the potassa forms a sulphate; the greater portion of the sulphur, however, unites with atmospheric oxygen and is converted into sulphurous acid. The nitric acid, having lost part of its oxygen, is changed into binoxide of nitrogen, which immediately combining with oxygen of the air forms nitrous acid, and, together with the sulphurous acid, passes into the chamber. The watery vapour present in the chamber, coming into contact with the nitrous and sulphurous acids, causes an immediate change in their composition, and a crystalline compound results, consisting of sulphuric acid, hyponitrous acid, and water. This crystalline substance, in consequence of its specific gravity, falls into the water, and decomposition again ensues. The sulphuric acid is retained in solution, while the hyponitrous acid is converted into nitrous acid and binoxide of nitrogen, which escape with effervescence. The binoxide meeting with more atmospheric oxygen, is instantly changed into nitrous acid, and enters into the composition of a further portion of crystalline matter, which is again precipitated and decomposed. These changes are continually repeated until the water is made very acid, when it is drawn off and concentrated by boiling in platinum vessels. The following diagrams are intended to illustrate the several changes which occur :

40 Equivalent.

Composition of Aqueous $\begin{cases} 1 & \text{Sulphuric Acid} = 40 \\ 1 & \text{Water} = 9 \end{cases}$

49 Equivalent.

CHANGES WHICH TAKE PLACE WITHOUT THE CHAMBER.

Atmospheric	c Oxygen	Nitrous Acid.
Nitrate of	SBinoxide of Nitrogen	to a blink strait store ?
Potassa.	(Potassa	Sulphate of Potassa.
Sulphur		and a survey of the second
Sulphur		
Atmospherie	c Oxygen	Sulphurous Acid.

CHANGES WHICH TAKE PLACE WITHIN THE CHAMBER.

	Crystalline compound.
Water Water.	The Revenue

Crystalline J	Hyponitrous Acid, 2 eqs.	Oxygen.	1 Cagain forms Nitrous Acid. 2
compound.	Sulphuric Acid		4 ——— Nitrous acid. 2
	-Water		Retained by the water on the floor

PREPARATION OF PHOSPHORUS.

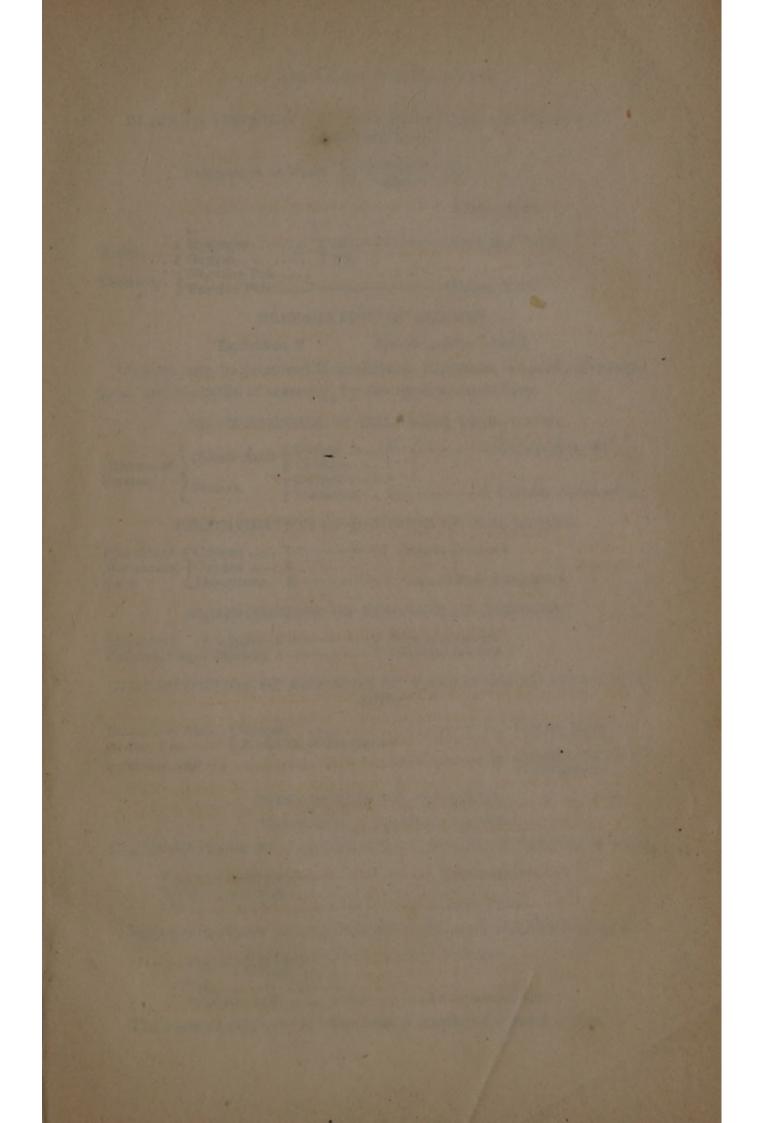
Equivalent, 16.

Sulphuric acid is added to the well burnt ashes of bone, and sulphate and superphosphate of lime are formed. The latter being the more soluble salt, is dissolved out by water, and the solution having been evaporated to the consistence of syrup, is mixed with charcoal. By the assistance of heat the oxygen is removed from the excess of phosphoric acid in the form of carbonic oxide and carbonic acid, phosphorus distils, and phosphate of lime remains in the retort.

A.	
Bone ashes. { Phosphoric Acid	Superphosphate of Lime.
Sulphuric Acid	Sulphate of Lime.
В.	
Carbon	
Superphosphate Science	usDistils over.

24

R



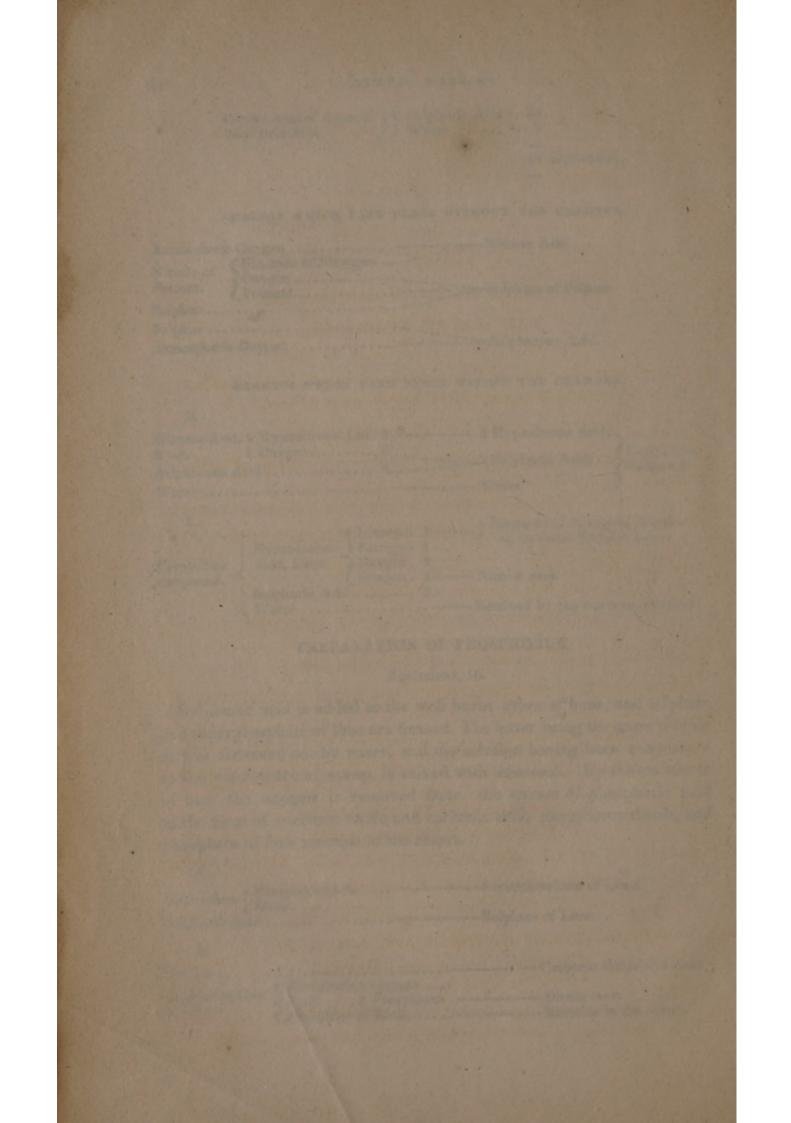


DIAGRAM SHOWING THE DECOMPOSITION OF WATER BY ELECTRICITY.

Composition of Water. $\begin{cases} 1 & \text{Hydrogen} = 1 \\ 1 & \text{Oxygen} \\ \dots = 8 \end{cases}$

9 Equivalent.

	Hydrogen 2 vols. ————————————————————————————————————	
Electricity.	Positive Pole Oxygen 1 vol.	

PREPARATION OF OXYGEN.

Equivalent 8. Specific gravity 1.1026.

Oxygen may be procured from chlorate of potassa, binoxide of manganese, and binoxide of mercury, by the application of heat.

DECOMPOSITION OF CHLORATE OF POTASSA

Chlorate of Potassa.	oric Acid. { Oxygen 56 Oxygen given off. Chlorine 1 oxygen 1 Potassium 11 Chloride of Potassium.
DECO	MPOSITION OF BINOXIDE OF MANGANESE.
Binoxide of Manganese, 2 eqs.	ygen 1 — 1 Oxygen given off. ygen 3 nganese 2- — 1 Sesquioxide of Manganese.

DECOMPOSITION OF BINOXIDE OF MERCURY.

DECOMPOSITION OF BINOXIDE OF MANGANESE BY SULPHURIC ACID.

Binoxide of Man-ganese, 1 eq. {Oxygen 1—1 Oxygen given off. Protoxide of Manganese 1 § 1 Sulphate of Protoxide of Manganese. Sulphuric Acid 1-

PREPARATION OF HYDROGEN.

Equivalent 1. Specific gravity '069.

DECOMPOSITION OF THE VAPOUR OF WATER BY RED HOT IRON.

Vapour of { Hydrogen... 1 — 1 Hydrogen given off. Water. Oxygen.... 1 ------ 1 Oxide of Iron. Iron

DECOMPOSITION OF WATER BY ZINC AND SULPHURIC ACID.

The same changes occur when iron is employed instead of zinc.

PREPARATION OF NITROGEN.

Equivalent 14. Specific gravity 0.9727.

Nitrogen is prepared by burning phosphorus in a jar containing atmospheric air inverted over water, thus—

 Atmospheric
 Nitrogen
 Nitrogen, set free, containing carbonic Acid

 Air.
 Oxygen.
 derived from the Air.

 Phosphorus
 Metaphosphoric Acid absorbed by the water.

PREPARATION OF PROTOXIDE OF NITROGEN.

Composition. $\begin{cases} 1 \text{ Nitrogen} = 14 \\ 1 \text{ Oxygen} = 8 \end{cases}$

22 Equivalent.

DECOMPOSITION OF NITRATE OF AMMONIA BY HEAT.

PREPARATION OF BINOXIDE OF NITROGEN.

Nitric Acid, Nitric Acid, Nitrogen... 1 4 eqs. Nitric Acid, Nitrogen... 2 Nitric Acid, Nitrogen... 2 Oxygen ... 2 Oxygen ... 3 Solution of Nitrogen... 2 Oxygen ... 3 Solution of Nitrogen... 2 Solution of Nitrogen... 3 Solution o

PREPARATION OF NITROUS ACID.

Composition. $\begin{cases} 1 \text{ Nitrogen} = 14 \\ 4 \text{ Oxygen} = 32 \end{cases}$

46 Equivalent.

Atmospheric Oxygen..... 2 Binoxide of Nitrogen..... 1

1 Nitrous Acid.

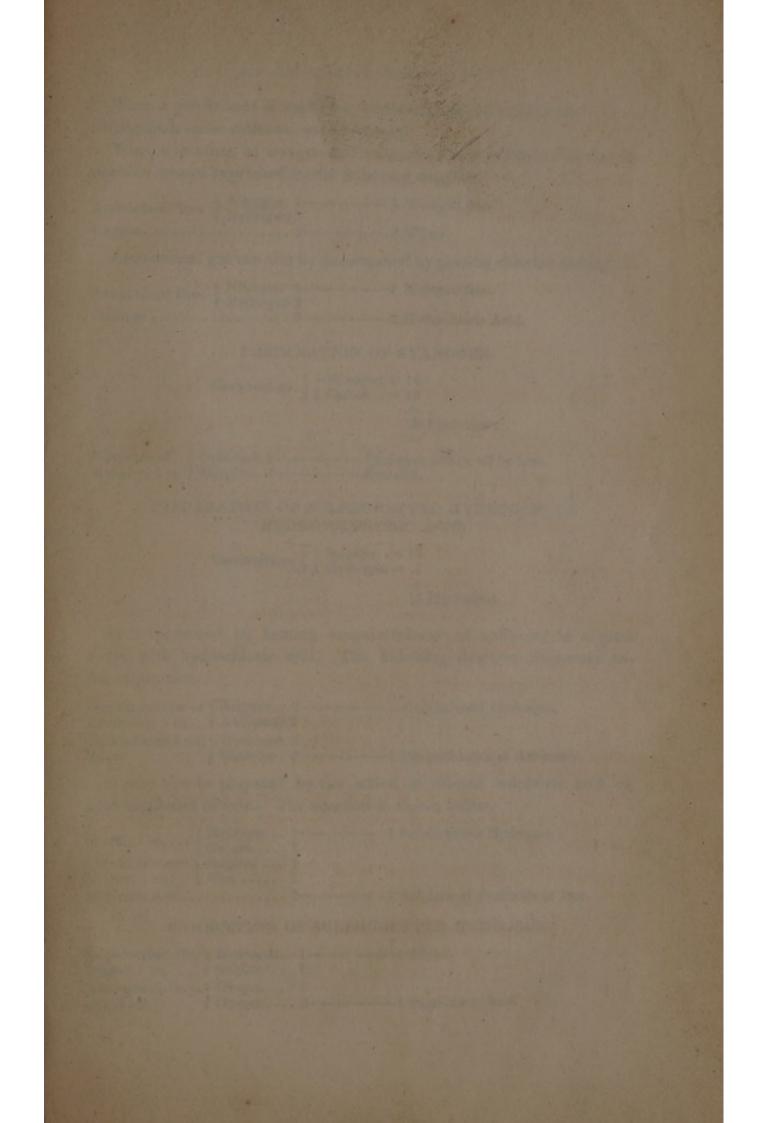
This acid may also be obtained in the liquid form by applying heat to the dry nitrate of lead, and condensing the vapours by a freezing mixture.

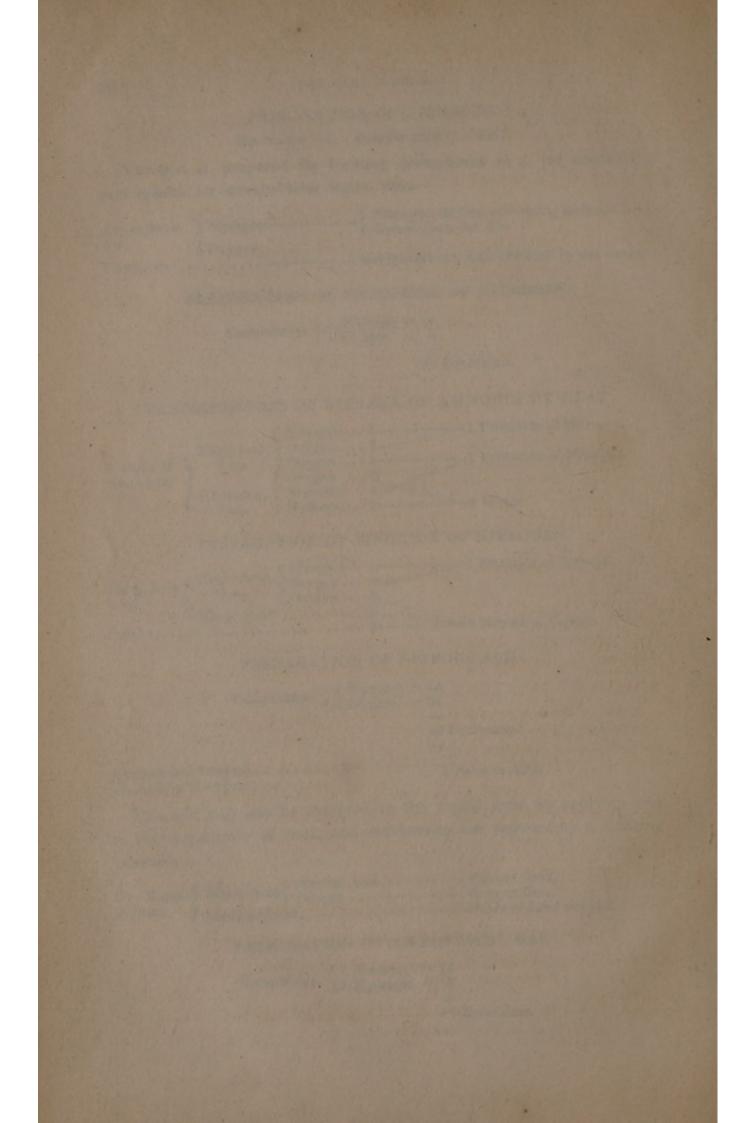
Dry Nitrate Nitric Acid Nitrous Acid. of Lead. Oxide of Lead.....Oxygen free. Oxide of Lead.....Oxide of Lead remains.

PREPARATION OF AMMONIACAL GAS.

Composition. $\begin{cases} 1 \text{ Nitrogen} = 14 \\ 3 \text{ Hydrogen} = 3 \\ -- \end{cases}$

17 Equivalent.





When a gentle heat is applied to liquor ammoniæ, ammoniacal gas is disengaged, to be collected over mercury.

When a mixture of oxygen and ammoniacal gas is fired, the decomposition ensues expressed in the following diagram.

Ammoniacal Gas.	Nitrogen Hydrogen	11	Nitrogen free.
Oxygen			Water.

Ammoniacal gas can also be decomposed by passing chlorine through it.

Ammoniacal Gas.	Nitrogen Hydrogen	$\frac{1}{3}$	Nitrogen free.
Chlorine			Hydrochloric Ac

PREPARATION OF CYANOGEN.

Composition. $\begin{cases} 1 \text{ Nitrogen} = 14 \\ 2 \text{ Carbon.} = 12 \end{cases}$

26 Equivalent.

id.

Bicyanide of { Cyanogen 2 _____Cyanogen driven off by heat. Mercury, 1 eq. { Mercury 1 _____Remains.

PREPARATION OF SULPHURETTED HYDROGEN, OR HYDROSULPHURIC ACID.

Composition. $\begin{cases} 1 \text{ Sulphur.} = 16 \\ 1 \text{ Hydrogen} = 1 \end{cases}$

17 Equivalent.

It is procured by heating sesquisulphuret of antimony in a glass vessel with hydrochloric acid. The following diagram illustrates the decomposition.

It may also be prepared by the action of diluted sulphuric acid on protosulphuret of iron. The reaction is shewn below.

Water, 1 eq }	Hydrogen	11 1	Sulphuretted Hydrogen.
Protosulphuret y	Sulphur	1	
of Iron, 1 eq. (Sulphuric Acid			Sulphate of Protoxide of Iron.

COMBUSTION OF SULPHURETTED HYDROGEN.

Sulphuretted Hy-5	Hydrogen		Water.	
	Sulphur			
Atmospheric Oxy-			Sulphurous	Acid
gen, 3 eqs.	Oxygen	2	Sulphurous	ration

Sulphuretted hydrogen is decomposed by chlorine, iodine, and bromine, with the formation of hydrochloric, hydriodic, and hydrobromic acids, sulphur being precipitated.

Chlorine		11	Hydrochloric Acid.
Sulphuretted Hy-	Hydrogen	1	and the second second second
drogen, 1 eq.	Sulphur	1	Sulphur precipitated.

Sulphuretted hydrogen decomposes the salts of the following metals:-

Antimony.	Columbium.	Molybdenum.	Tellurium.
Arsenicum.	Copper.	Osmium.	Tin.
Bismuth.	Copper. Gold.	Palladium.	Titanium.
Cadmium.	Iridium.	Platinum.	Tungsten.
Cerium.	Lead.	Rhodium.	Vanadium.
Chromium.	Mercury.	Silver.	Zinc.

The following diagram, showing the decomposition of acetate of lead by sulphuretted hydrogen, may serve as an example.

Acetate of Lead,	Acetic Acid 1 1 Acetic Acid free. Oxide of {Oxygen 1
Sulphuretted Hy-	Hydrogen 1 Sulphur

Salts of the metals named below are not decomposed by sulphuretted hydrogen.

Aluminum.	Iron.	Nickel.	Thorinum.
Barium.	Lithium.	Potassium.	Uranium.
Calcium.	Magnesium.	Sodium.	Yttrium.
Cobalt.	Manganese.	Strontium.	Zirconium.
Glucinum.			

But the salts of cobalt, iron, manganese, nickel, and uranium, are precipitated by sulphuret of potassium.

HYDROFLUORIC ACID

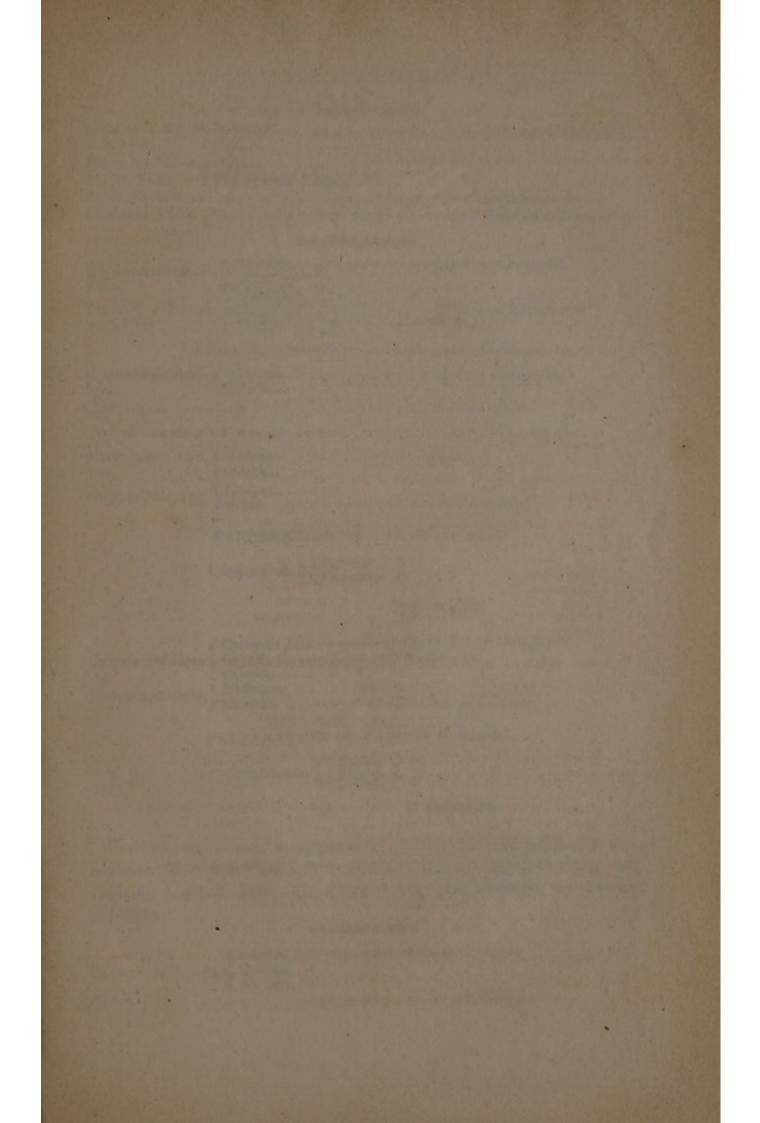
Is obtained by mixing sulphuric acid and fluoride of calcium in a leaden retort, and applying a gentle heat.

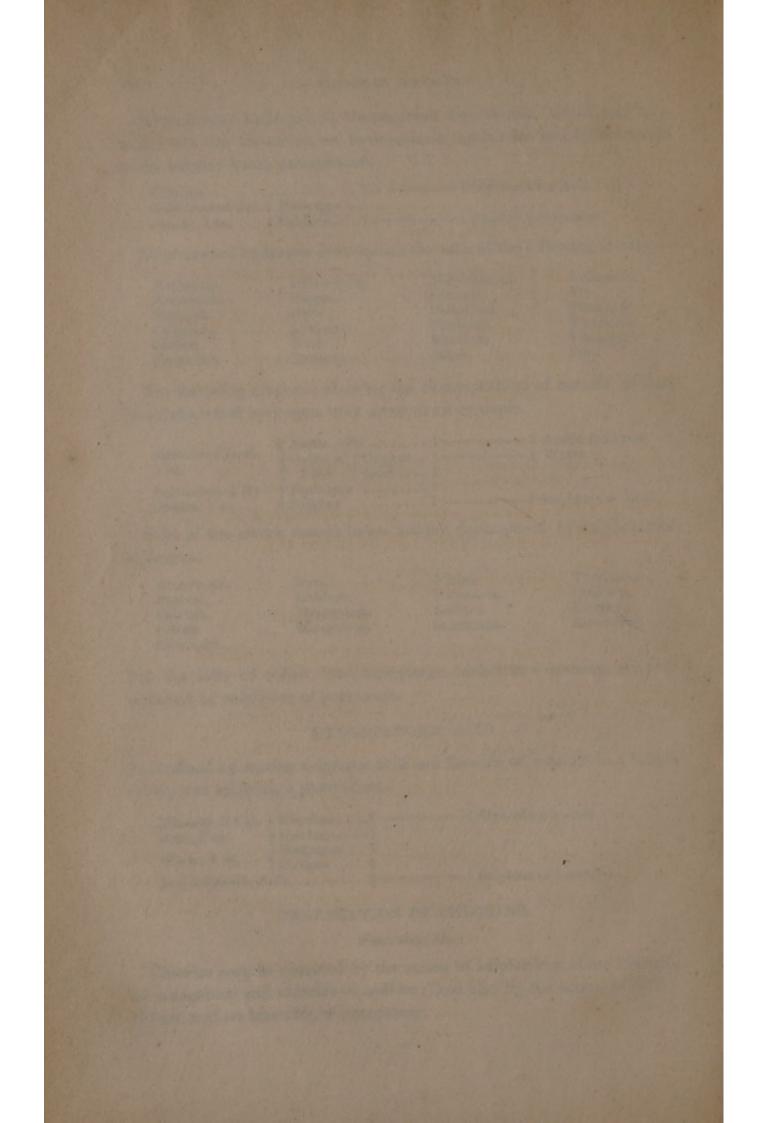
Fluoride of Cal-	Fluorine	11	Hydrofluoric Acid.	
	Calcium			
Water, 1 eq	Hydrogen	1		
Dry Sulphuric A	cid	î1	Sulphate of Lime.	

PREPARATION OF CHLORINE.

Equivalent 36.

Chlorine may be obtained by the action of sulphuric acid on binoxide of manganese and chloride of sodium; and also by the action of hydrochloric acid on binoxide of manganese.





PROCESS FIRST.

Chloride of So- { Chlorine 1 dium, 1 eq. { Sodium 1	
Binoxide of Man- 5 Oxygen 1	
ganese, 1 eq. ? Protoxide of Manganese 1	and a construction of a second second second
Sulphuric Acid 1 Sulphuric Acid 1	
PROCESS SEC	
Hydrochloric Acid, 2 eqs. Chlorine1 Hydrogen2	I Chlorine disengaged.
Binoxide of Manga- Manganese. 1-	
Binoxide of Manga- nese, 1 eq. Manganese 1 Oxygen 2	
ACTION OF HYDROCHLORIC	ACID ON A METAL.
Hydrochloric Acid, 5 Hydrogen 1 1 eq. (Chlorine 1	
Zinc 1	-1 Chloride of Zinc.
ACTION OF HYDROCHLORIC ACID	ON A METALLIC OXIDE.
Hydrochloric Acid, { Hydrogen 1	
Oxide of Zinc, 1 eq. } Oxygen 1 Zinc 1	-1 Chloride of Zinc.
PREPARATION OF CA	ARBONIC ACID.
Composition. { 1 Carbon = 2 Oxygen =	6 16
	22 Equivalent.
Carbonic Acid	- Carbonic Acid disengaged.
Carbonate of Lime. Oxygen	
Hydrochloric Acid. { Hydrogen Chlorine	-Chloride of Calcium.
PREPARATION OF CAL	RBONIC OXIDE.
Composition. { 1 Carbon 1 Oxygen	= 6 = 8
	14 Equivalent.
Carbonic oxide may be prepared by	passing carbonic acid over red
hot charcoal or iron filings, or by mixin	
applying a gentle heat; the annexed positions.	diagrams illustrate the decom-
PROCESS FIL	RST.
Carbonic Acid, 1 eq. { Carbon 1 Oxygen 1 Oxygen 1	-1 Carbonic Oxide.
Carbon	-1 Carbonic Oxide.

PROCESS SECOND.

Carbonic Acid, 1 eq.	Carbon 1 Oxygen 1 Oxygen 1	1 Carbonic Oxide. 1 Protoxide of Iron.
	PROCESS	
Oxalic Acid	Carbon 1 Oxygen 2 Oxygen 1 Water 3	1 Carbonic Oxide off.
Sulphuric Acid		Left in the retort.

The carbonic acid formed in the last process is removed by agitating the gases with lime water.

PREPARATION OF CHLORATE OF POTASSA.

Chlorine.	Chlorine	1	Chlorate of Potassa.
Potassa	Potassa Oxygen Potassium	1 5	Chloride of Potassium.

The chlorate is easily separated by evaporation and crystallization. When sulphuret of potassium is put into water, the following changes take place.

Sulphuret of Potas-	A COMPANY OF A COM	
sium, 1 eq.	Hydrogen	
Water, 1 eq	Oxygen	ll Potassa.

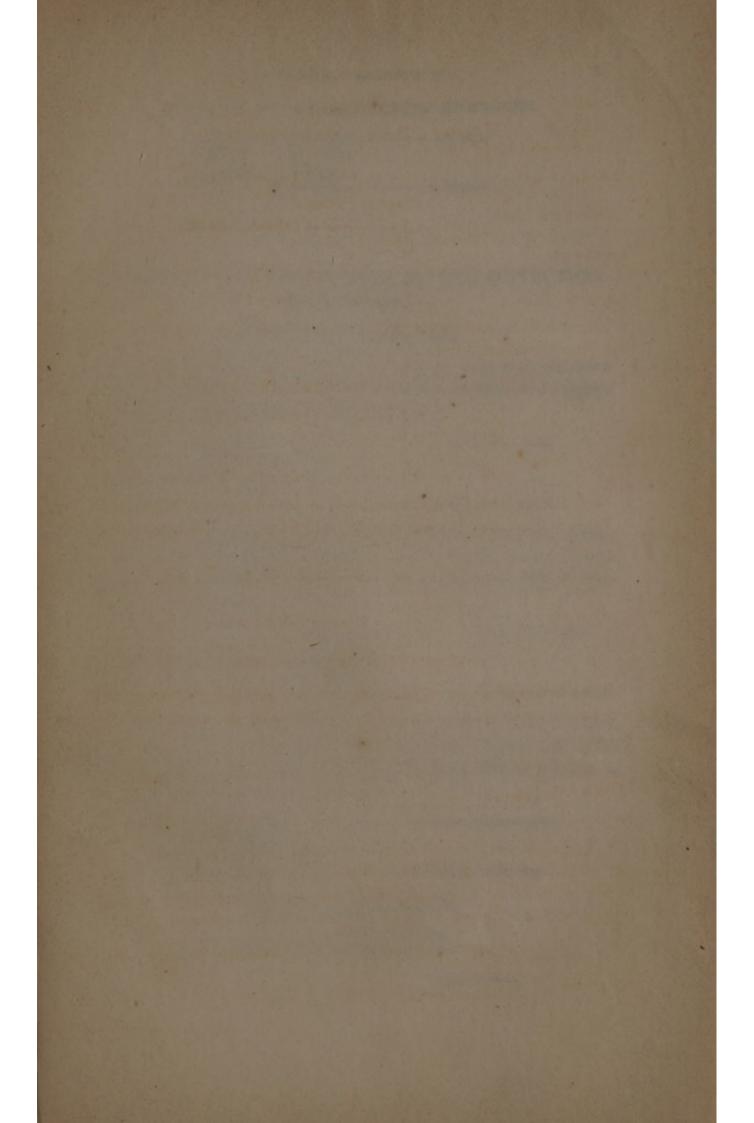
PREPARATION OF POTASSIUM.

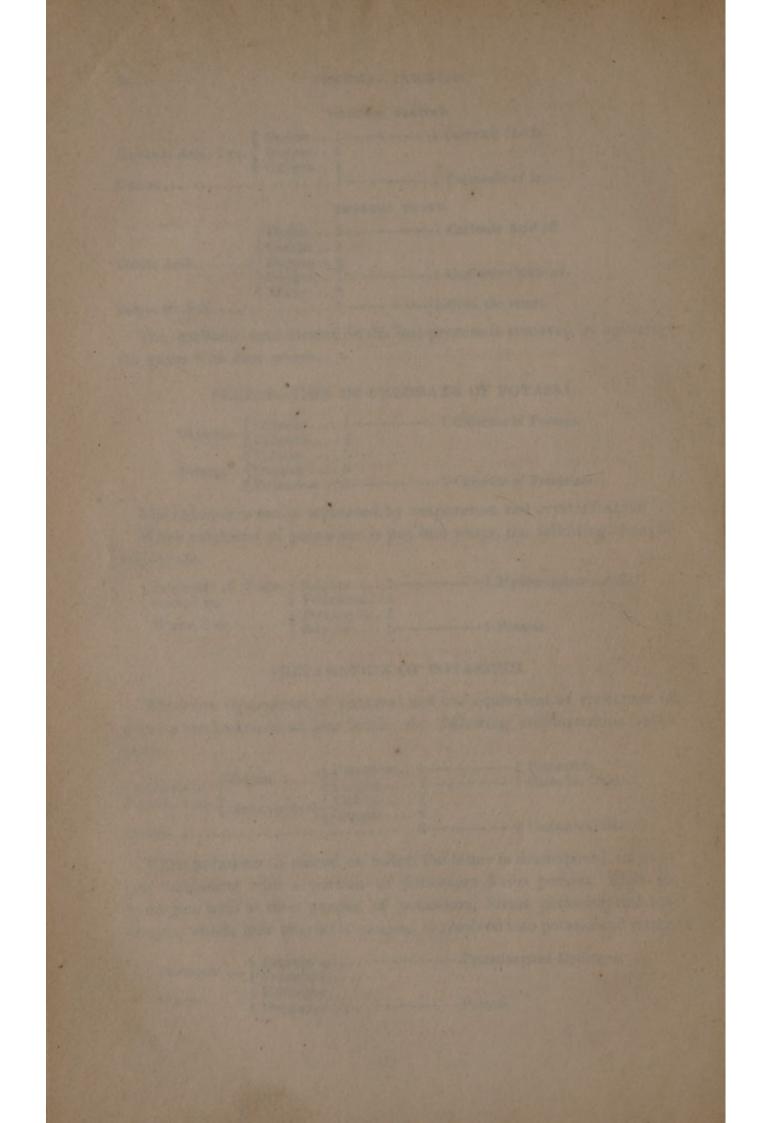
When two equivalents of charcoal and one equivalent of carbonate of potassa are heated in an iron bottle, the following decomposition takes place.

Carbonate of Potassa	Potassium	11	Potassium. Carbonic Oxide.
Potassa, 1 eq. Carbonic Acid			curbonic childer
Carbon			Carbonic Oxide.

When potassium is placed on water, the latter is decomposed, its oxygen combining with a portion of potassium forms potassa, while its hydrogen, with another portion of potassium, forms potassiuretted hydrogen, which, inflaming as it escapes, is resolved into potassa and water.

Potassium }	Potassium	Potassiuretted	Hydrogen.
	Hydrogen	Potassa.	





COMBUSTION OF POTASSIURETTED HYDROGEN.

Potassiuretted { Hydrogen.	Hydrogen ——— Potassium	Water.
Atmospheric §	Oxygen	Potassa

DIAGRAMS OF PROCESSES USED IN THE DETECTION OF POISONS.

TESTS FOR SULPHURIC ACID.

When chloride of barium is added to a solution containing sulphuric acid, water is decomposed, hydrochloric acid being formed, and sulphate of baryta precipitated, insoluble in nitric acid.

Chloride of Ba-	Chlorine	1
rium, 1 eq.	Barium	
Water, 1 eq	Hydrogen	I supply heading a supply of the
Water, 1 eq	Oxygen	1
Sulphuric Acid		1

The sulphate of baryta being heated with charcoal, the 4 eqs. of oxygen contained in the sulphate unite with 4 eqs. of carbon, to form 4 eqs. of carbonic oxide, which are expelled, while the sulphur forms a sulphuret with the barium.

Carbon		
Sulphate of Ba-	Sulphuric Acid .	Oxygen3 Sulphur. 1
		Oxygen . 1 Barium . 11 Sulphuret of Barium.

To the sulphuret of barium thus obtained hydrochloric acid is added, chloride of barium is formed, and hydrosulphuric acid given off, which may be known by its blackening paper moistened with a solution of acetate of lead and held over the fumes.

TESTS FOR OXALIC ACID.

Composition of Dry Acid. $\begin{cases} 3 \text{ Oxygen} = 24 \\ 2 \text{ Carbon} = 12 \end{cases}$

36 Equivalent.

CHEMICAL DIAGRAMS

Composition of Crystals. $\begin{cases} 1 \text{ Oxalic Acid} = 36 \\ 3 \text{ Water } \dots = 27 \\ \hline 63 \text{ Equivalent.} \end{cases}$

When chloride of calcium is added to a solution containing oxalic acid, hydrochloric acid is formed, and a heavy white oxalate of lime thrown down, soluble in nitric acid, but not in hydrochloric acid, unless in excess.

Chloride of Cal-	Chlorine Calcium	
Oxalic Acid		1

Sulphate of copper causes a greenish white precipitate of oxalate of copper, not dissolved by a small quantity of hydrochloric acid, but completely so in an excess of the acid.

Sulphate of Copper { Sulphuric Acid. _____ Free. Oxalic Acid . _____ Oxalate of Copper _____ Oxalate of Copper.

When chalk has been taken as an antidote, an insoluble oxalate of lime is already formed; this must be boiled with carbonate of potassa; oxalate of potassa and carbonate of lime are thus formed; the former is held in solution, the latter is precipitated.

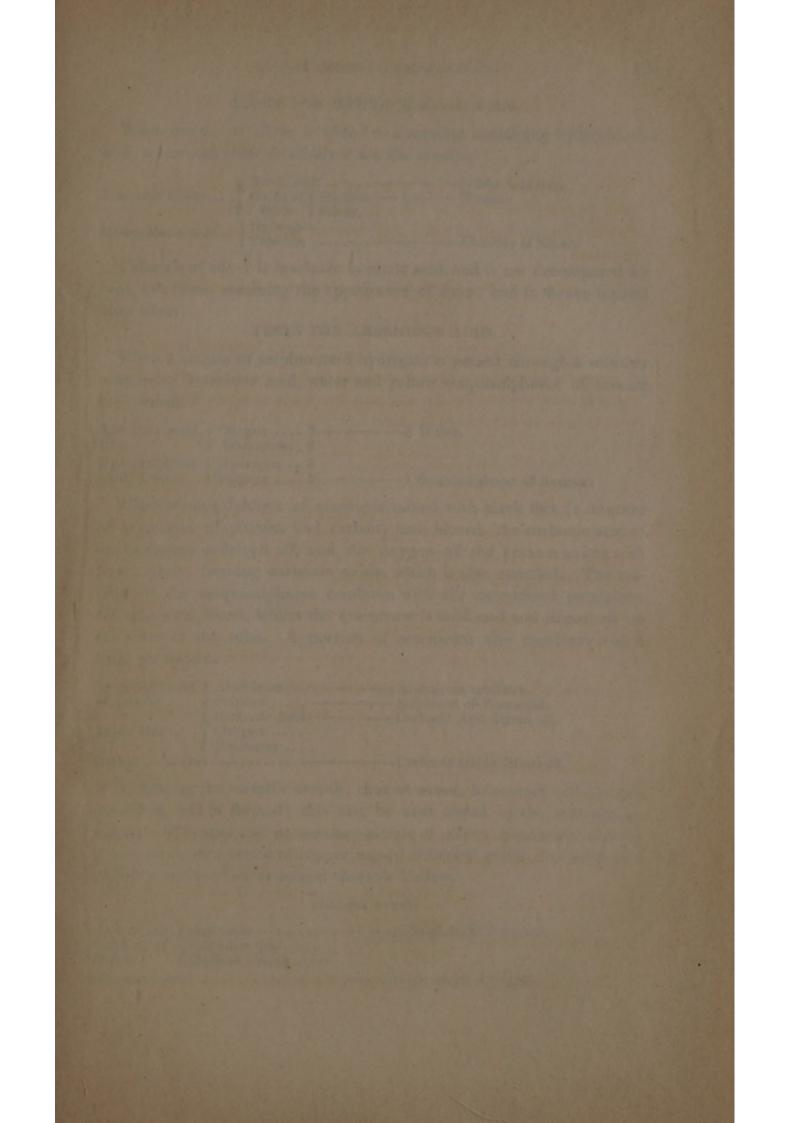
Carbonate of Potassa	PotassaOxalate of Potassa.
	0 11 4 11

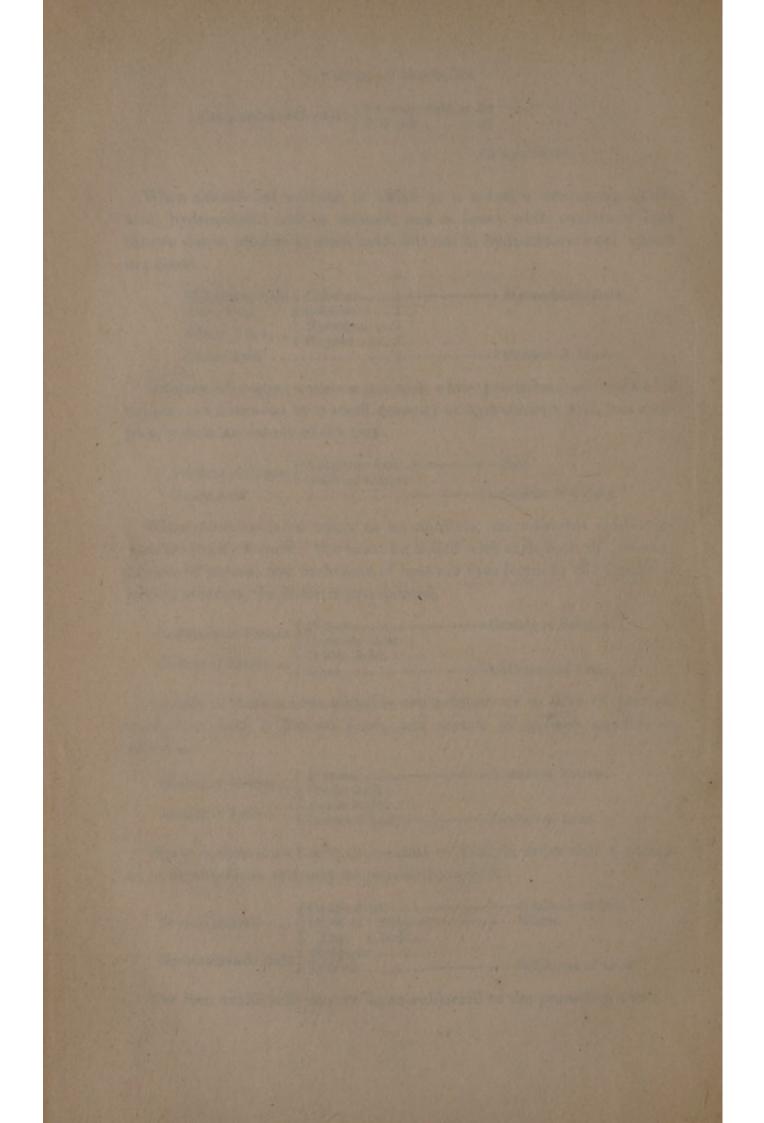
Acetate of lead is next added to the solution of oxalate of potassa, oxalate of lead is thrown down, and acetate of potassa remains in solution.

Oxalate of Potassa	Potassa Oxalic Acid	Acetate of Potassa.
Acetate of Lead		

Water is then mixed with the oxalate of lead, in order that a stream of hydrosulphuric acid may be passed through it.

The free oxalic acid may be again subjected to the preceding tests.





TESTS FOR HYDROCHLORIC ACID.

When nitrate of silver is added to a solution containing hydrochloric acid, water and chloride of silver are the results.

Nitrate of Silver	Nitric Acid Oxide of § Oxygen Silver § Silver	Nitric Acid free. Water.
Hydrochloric Acid	Hydrogen	

Chloride of silver is insoluble in nitric acid, and is not decomposed by heat, but fuses, assuming the appearance of horn, and is thence termed horn silver.

TESTS FOR ARSENIOUS ACID.

When a stream of sulphuretted hydrogen is passed through a solution containing arsenious acid, water and yellow sesquisulphuret of arsenic are formed.

Arsenious Acid,	Oxygen	3
1 eq.	Arsenicum	2
Hydrosulphuric 9		
Acid, 3 eqs.	Sulphur	3

When sesquisulphuret of arsenic is mixed with black flux (a mixture of carbonate of potassa and carbon) and heated, the carbonic acid of the carbonate is driven off, and the oxygen of the potassa unites with free carbon, forming carbonic oxide, which is also expelled. The sulphur of the sesquisulphuret combines with the deoxidized potassium, forming a sulphuret, whilst the arsenicum is sublimed and deposited on the sides of the tube. A portion of arsenicum also combines with a little potassium.

	Arsenicum	Arsenicum sublimes.
	Sulphur	Sulphuret of Potassium.
	Carbonic Acid-	Carbonic Acid driven off.
Black Flux	Oxygen	
	Potassium	
Carbon	And the second sec	Carbonic Oxide driven off.

By heating the metallic arsenic, thus obtained, in contact with the air, arsenious acid is formed; this may be next tested by the ammoniacosulphate of copper and ammoniaco-nitrate of silver, the former causing a precipitate of arsenite of copper named Scheeles' green, the latter of a yellow arsenite of silver named Queen's Yellow.

PROCESS FIRST.

Ammoniaco- Sulphate of Copper. Ammonia Sulphuric Acid Oxide of Copper	Sulphate of Ammonia.
	-Arsenite of Copper.

CHEMICAL DIAGRAMS

PROCESS SECOND.

Ammoniaco- (Ammonia	
Nitrate of Nitric Acid	
Silver. (Oxide of Silver	
Arsenious Acid	

When arsenious acid is mixed with charcoal and heated, metallic arsenic sublimes, giving off a garlic odour.

Arsenicum Arsenicum sublimes.
Carbonic Acid and Oxide.

MARSH'S TEST.

When zinc, sulphuric acid, and a solution of arsenious acid are placed in a vessel furnished with a stop-cock, the water is decomposed, the oxygen of which combines with the zinc, forming an oxide with which the sulphuric acid unites to form a sulphate. Part of the hydrogen of the decomposed water combines with the oxygen of the arsenious acid and forms water, the rest of the hydrogen unites with the metallic arsenic, forming arseniuretted hydrogen, which is allowed to escape through the stop-cock. The gas is then inflamed and the results of combustion, (namely, metallic arsenic, arsenious acid, and water,) are collected on a plate of glass held horizontally over the flame. The following diagram shows a more precise view of the changes which occur.

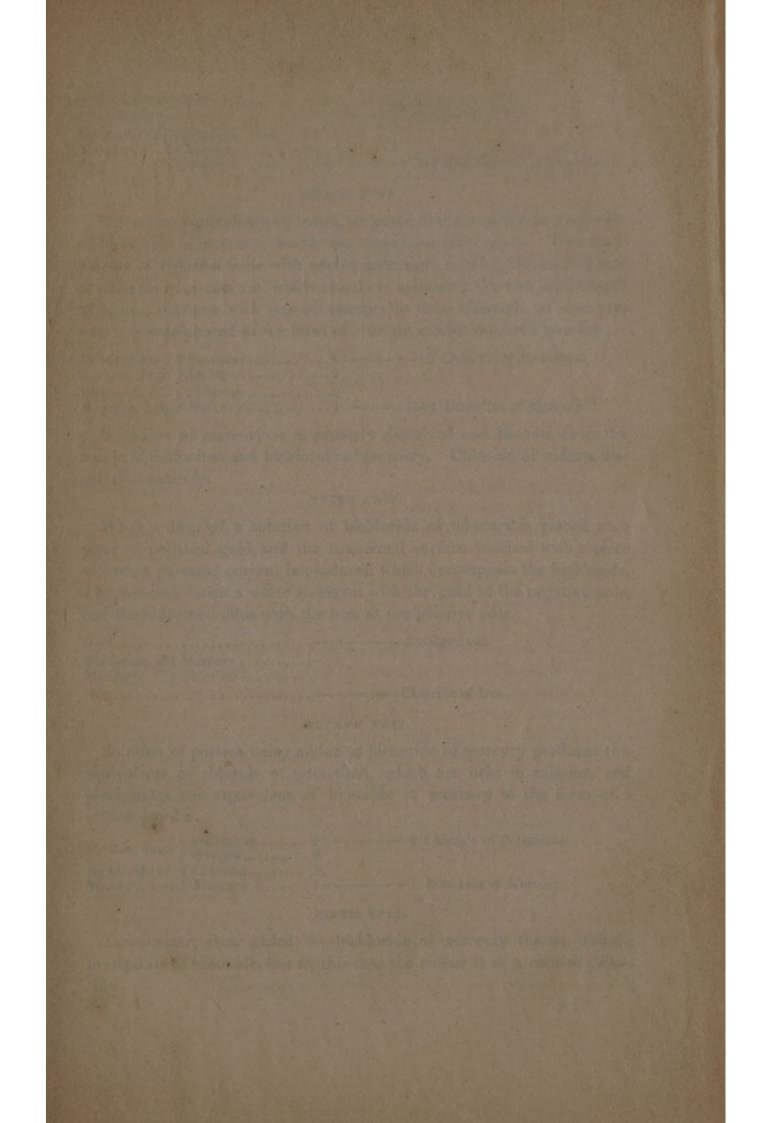
Arsenious	§ Arsenicum2	-2 Arseniuretted Hydrogen.
Acid, 1 eq.) Oxygen3	a solution while the ors
Water,	Hydrogen2	main and and and a bit of
5 000	Hydrogen3	
and the second	(Oxygen	
		5 Salabata of Ting
Sulphuric A	cid5	5 Sulphate of Zinc.

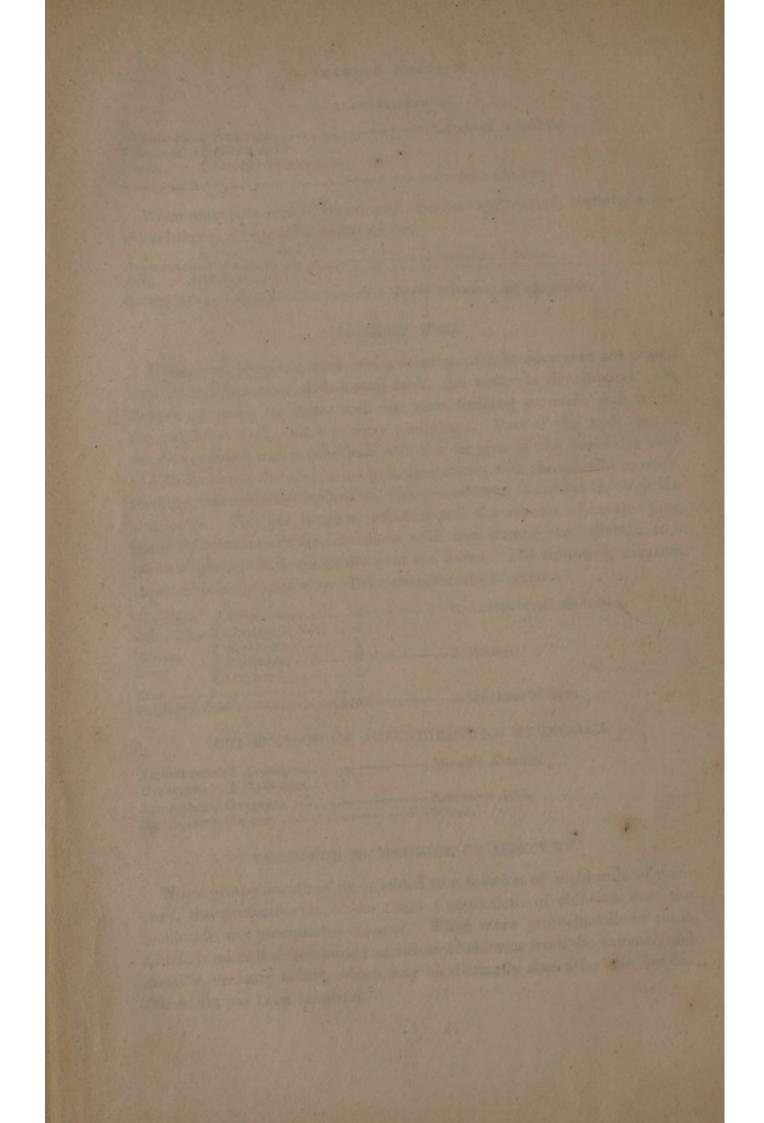
COMBUSTION OF ARSENIURETTED HYDROGEN.

	Arsenicum	Metallic Arsenic.
Hydrogen.	Hydrogen	
Atmospheric O	xygen	Arsenious Acid.
Atmospheric O	xygen	Water.

TESTS FOR BICHLORIDE OF MERCURY.

When protochloride of tin is added to a solution of bichloride of mercury, the protochloride of tin takes 1 equivalent of chlorine from the bichloride and precipitates calomel. When more protochloride of tin is added, it takes the remaining equivalent of chlorine from the calomel, and metallic mercury is left, which may be distinctly seen after the bichloride of tin has been removed.





OF THE LONDON PHARMACOPCEIA.

Protochloride	of Tin	11	Bichloride of Tin.
Protochloride	of Tin	11	Bichloride of Tin.
Bichloride of	Chlorine	idead to milit	
Mercury,	Chlorine	1 martin the state	
leq.	Mercury	11	Metallic Mercury precipitated.

SECOND TEST.

When two equivalents of iodide of potassium are added to a solution of bichloride of mercury, double decomposition takes place. Two equivalents of chlorine unite with two of potassium forming two equivalents of chloride of potassium, which remain in solution; the two equivalents of iodine combine with one of mercury to form biniodide of mercury, which is precipitated in the form of a bright scarlet coloured powder.

Iodide of Po-	§ Potassium2 -	
tassium, 2 eqs.	Iodine2	
Bichloride of	Chlorine2	

Biniodide of mercury is alternately dissolved and thrown down by iodide of potassium and bichloride of mercury. Chloride of sodium dissolves it entirely.

THIRD TEST.

When a drop of a solution of bichloride of mercury is placed on a piece of polished gold, and the moistened surface touched with a piece of iron, a galvanic current is produced which decomposes the bichloride. The mercury forms a white amalgam with the gold at the negative pole, and the chlorine unites with the iron at the positive pole.

Gold		Amalgamate.
Bichloride of	Mercury	ar and manage stress
Mercury.	Chlorine	
Iron		Chloride of Iron.

FOURTH TEST.

Solution of potassa being added to bichoride of mercury produces two equivalents of chloride of potassium, which are held in solution, and precipitates one equivalent of binoxide of mercury in the form of a yellow powder.

Potassa, 2 eqs.	Potassium	22	Chloride of Potassium.
Bichloride of	Chlorine	2 2	
	Mercury		Binoxide of Mercury.

FIFTH TEST.

Lime-water, also, added to bichloride of mercury throws down a precipitate of binoxide, but in this case the colour is of a *reddish* yellow tint.

D 2

CHEMICAL DIAGRAMS

SIXTH TEST.

Ammonia added to a solution of bichloride gives a white flocculent precipitate of ammonio-chloride of mercury, (for the decomposition see the Pharmacopœial preparation.)

TESTS FOR POTASSIO-TARTRATE OF ANTIMONY.

Potassa throws down the oxide of antimony, which is redissolved by an excess of the alkali.

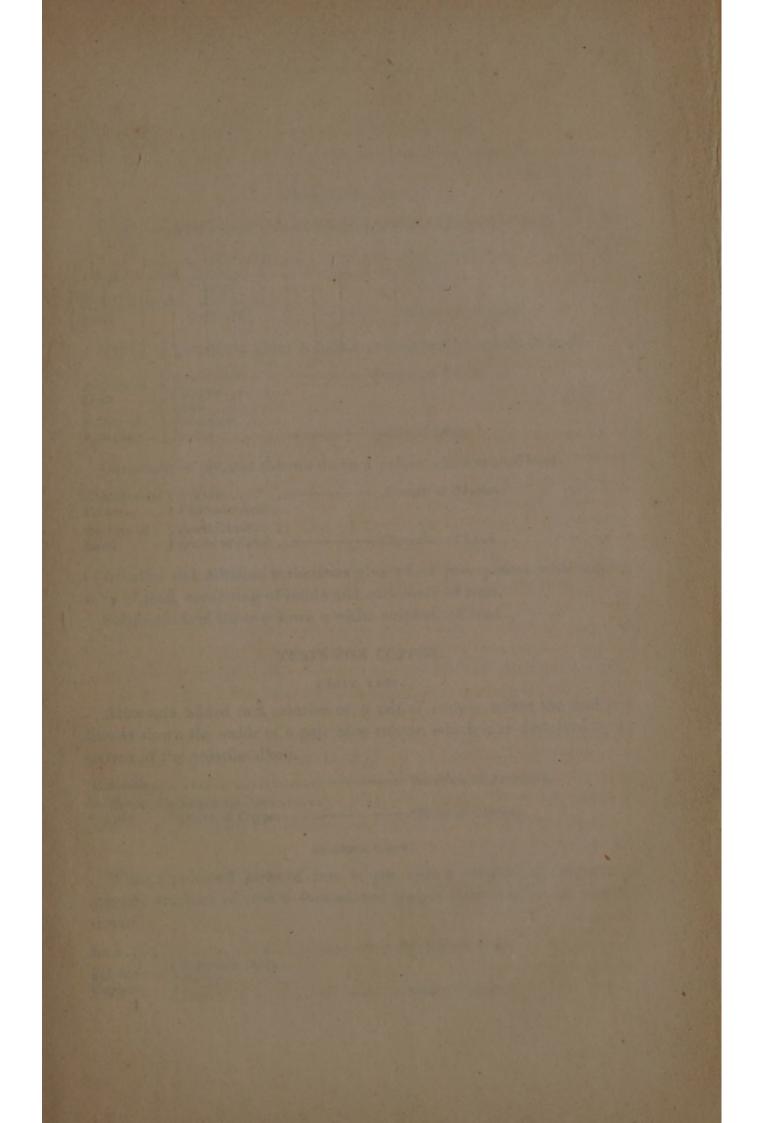
" Stripping	(Tartrate of P	otassa	Tartrate of Potassa (in solution.)
Potassio- Tartrate of - Antimony.	Ditartrate of Antimony.	Oxide of Antimony Tartaric Acid	
		Tartaric Acid	Tartrate of Potassa (in solution.)

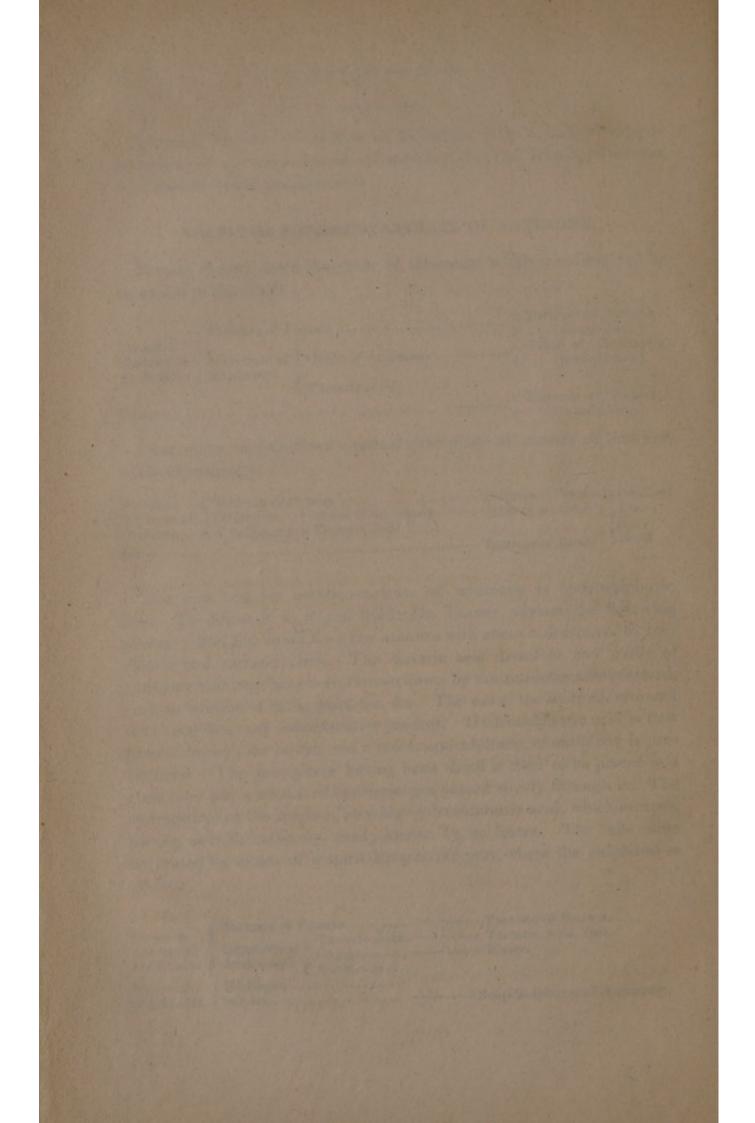
Lime water throws down a mixed precipitate of tartrate of lime and oxide of antimony.

Potassio-	Tartrate of Potassa Tartrate of Potassa in	solution.
Tartrate of	Ditartrate Oxide of Antimony Oxide of Antimony of Antimony. Tartaric Acid	Pre-
Antimony.	Tartrate of Potassa Tartrate of Potassa in Ditartrate Oxide of Antimony Oxide of Antimony of Antimony. Tartaric Acid	cipi-
	Tartrate of Lime.	tated.

The best test for potassio-tartrate of antimony is hydrosulphuric acid. To detect it in mixed fluids Dr. Turner advises the following process: Boil the liquid for a few minutes with about a drachm of hydrochloric and tartaric acids. The tartaric acid dissolves any oxide of antimony that may have been thrown down by the antidotes administered, such as infusion of bark, galls, tea, &c. The use of the hydrochloric acid is to coagulate any animal matter present. Hydrosulphuric acid is now passed through the liquid, and a red sesquisulphuret of antimony is precipitated. The precipitate having been dried is then to be placed in a glass tube and a stream of hydrogen gas passed slowly through it. The hydrogen takes the sulphur, forming hydrosulphuric acid, which escapes, leaving metallic antimony, easily known by its lustre. The tube must be heated by means of a spirit-lamp at the part where the sulphuret is placed.

A. Potassio- tartrate of Antimony.	Tartrate of Potassa Ditartrate of Tartaric Acid Antimony. Antimony	Tartrate of Potassa. Tartaric Acid, free. Water.
Hydrosul- phuric Acid.	Sulphur	Sesquisulphuret of Antimony.





B.

Sesquisulphuret		Metallic Antimony.
of Antimony. Hydrogen	(Sulphur	Hydrosulphuric Acid.

TESTS FOR LEAD.

Hydrosulphuric acid throws down a black sulphuret of lead.

Acetate of Lead.	Acetic Acid	Acetic Acid, free. Water.
Hydrosulphuric	(Lead) Hydrogen	
Acid.	! Sulphur	

Iodide of potassium gives a yellow precipitate of iodide of lead.

Acetate of Lead.	Acetic Acid Oxygen Lead	Acetate of Potassa.
Iodide of Potassium.	{ Potassium	Iodide of Lead.

Chromate of potassa throws down a yellow chromate of lead.

Chromate of	5 Potassa	Acetate of Potassa.
Potassa.	Chromic Acid	
Acetate of	Acetic Acid	doing a set in the set of the
Lead.	Oxide of Lead	Chromate of Lead.

Alkalies and alkaline carbonates give white precipitates with soluble salts of lead, consisting of oxide and carbonate of lead.

Sulphuric acid throws down a white sulphate of lead.

TESTS FOR COPPER.

FIRST TEST.

Ammonia added to a solution of a salt of copper seizes the acid and throws down the oxide of a pale blue colour, which is re-dissolved by an excess of the volatile alkali.

Ammonia Sulphate of Ammonia. Sulphate of Sulphuric Acid Copper. Oxide of Copper Oxide of Copper.

SECOND TEST.

When a polished piece of iron is put into a solution of sulphate of copper, sulphate of iron is formed and copper deposited in the metallic state.

CHEMICAL DIAGRAMS

When copper is contained in mixed fluids, hydrosulphuric acid is the best test. The resulting sulphuret having been collected and heated to redness that the organic matter may be charred, is next to be placed on a piece of porcelain and digested with nitric acid. The sulphuret is converted into a sulphate at the expense of the nitric acid, and may be submitted to the re-action of the foregoing tests.

A. Sulphate of Copper.	Sulphuric Acid Oxygen Copper	Sulphuric Acid, free. Water.
Hydrosulphur Acid.		Sulphuret of Copper.
B.		
NitricAcid.	S Nitrogen	Nitrogen.
Sulphuret of Copper.	Sulphur	

TESTS FOR MORPHIA.

Nitric acid added to morphia causes it to assume an orange-red colour soon changing to yellow.

Morphia is dissolved by sesquichloride of iron, producing a deep greenish blue solution.

When iodic acid is added to morphia, the acid is decomposed, and the iodine set free, which forms a blue colour with starch.

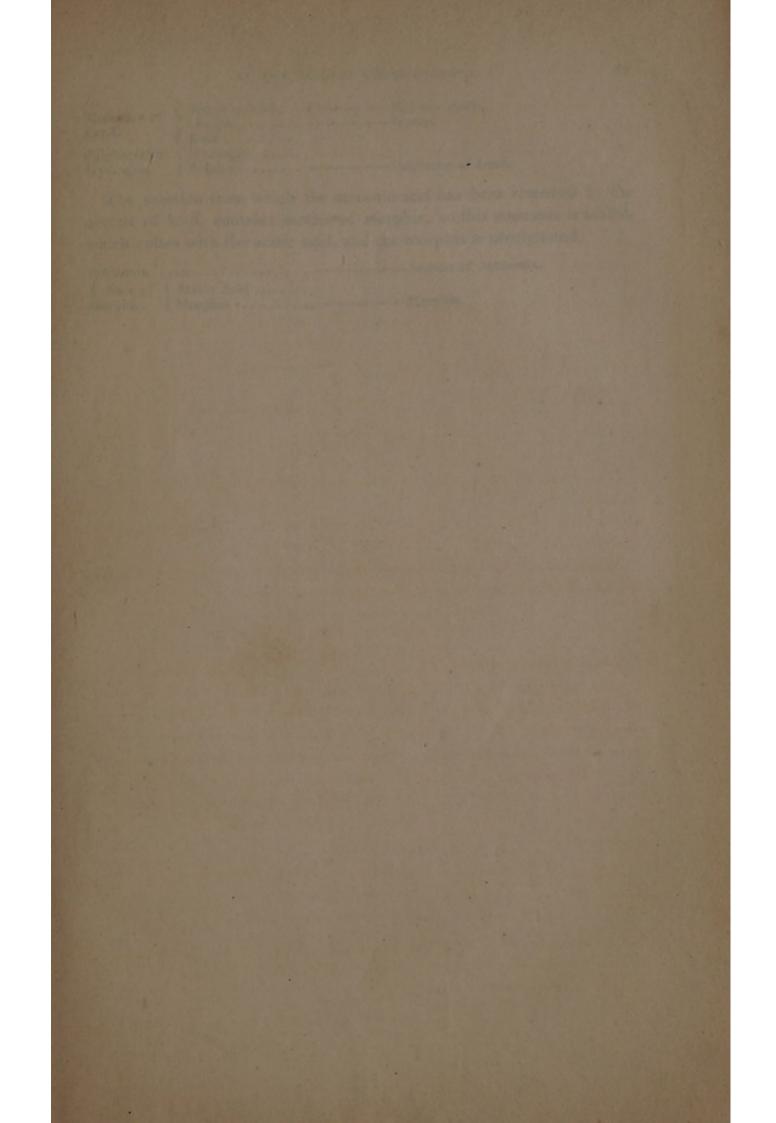
TEST FOR MECONIC ACID.

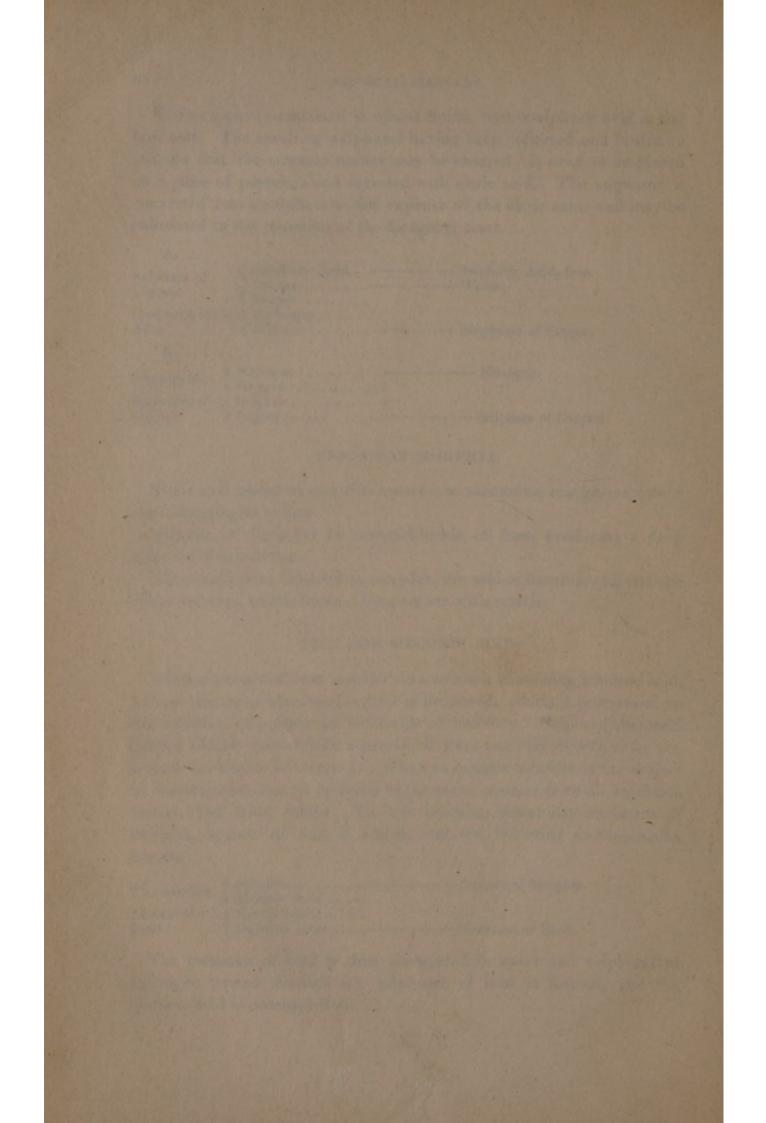
When a persalt of iron is added to a solution containing meconic acid, a deep cherry or blood-red colour is produced, which is permanent on the addition of potassa or bichloride of mercury. Sulpho-cyanic acid gives a similar colour with a persalt of iron, but it is destroyed by potassa or bichloride of mercury. When an organic mixture is the subject of investigation it is to be freed in the usual manner from all insoluble matter, and from colour. To the solution containing meconate of morphia, acetate of lead is added, and the following decomposition ensues.

The solution.	Morphia Meconic Acid	- Acetate of Morphia.
Acetate of	Acetic Acid Oxide of Lead	-Meconate of Lead.

The meconate of lead is then suspended in water and sulphuretted hydrogen passed through it; sulphuret of lead is formed, and the meconic acid is obtained free.

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OF THE LONDON PHARMACOPCEIA.

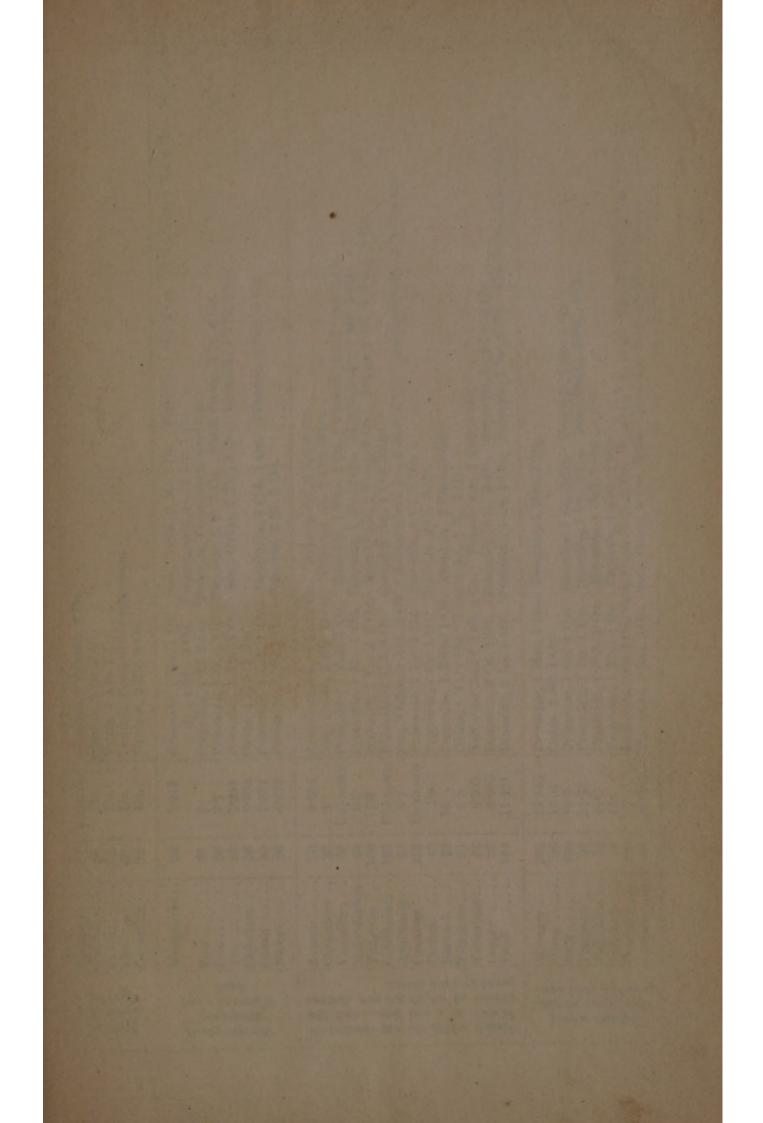
Meconate of Lead.	Meconic Acid Oxygen Lead	——Meconic Acid. ——Water.
Sulphuretted Hydrogen.) Hydrogen Sulphur	

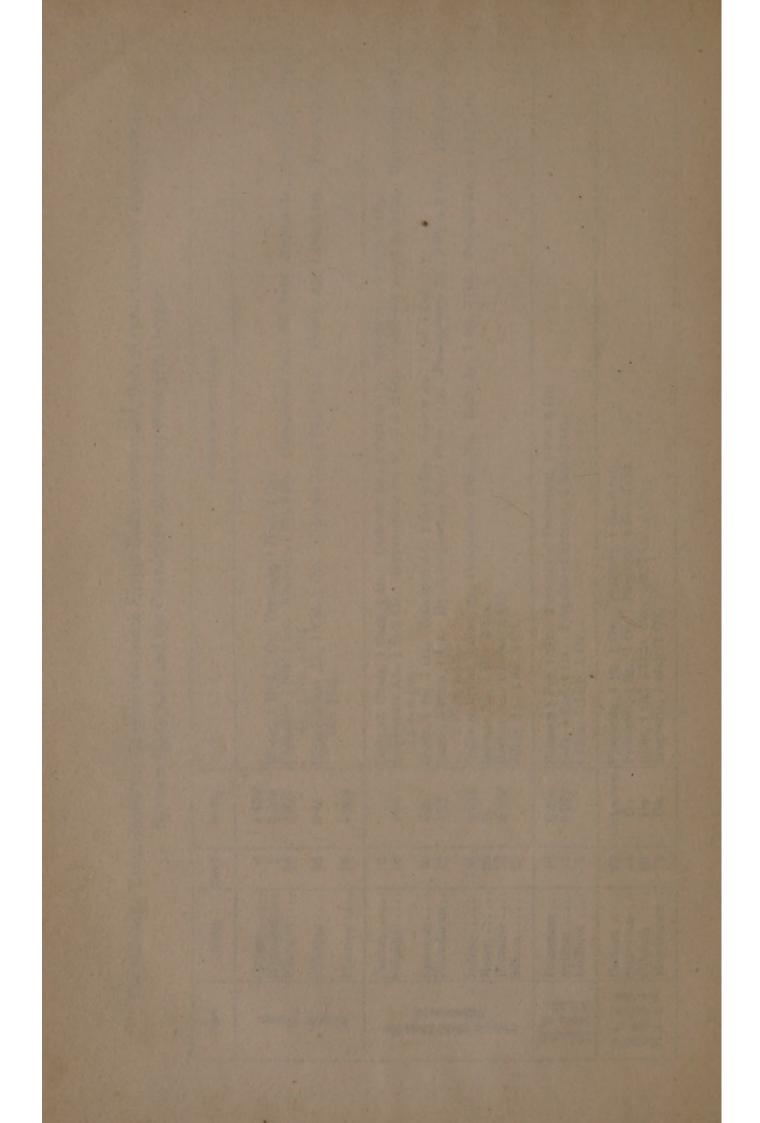
The solution from which the meconic acid has been removed by the acetate of lead, contains acetate of morphia, to this ammonia is added, which unites with the acetic acid, and the morphia is precipitated.

Ammonia	Acetate of Ammonia.
Acetate of	Acetic Acid
	MorphiaMorphia.

SPECIFIC GRAVITIES, and the COMPOUNDS WHICH they JOFIL WITH OXYGEN. Sp. Gr. Compounds with Oxygen.	Water, 1 Hyd. 1 Ox. Peroxide, 1 Hyd. 2 Ox. Protoxide, 1 Nit. 1 Ox. Binoxide, 1 Nit. 2 Ox. Hyponitrous Ac., 1 Nit. 3 Ox. Nitrous Ac., 1 Nit. 4 Ox. Nitric Ac., 1 Nit. 5 Ox. Hypochlorous Ac., 1 Chl. 1 Ox. Chlorous Ac., 1 Chl. 4 Ox. Chloric Ac., 1 Chl. 5 Ox. Perchloric Ac., 1 Chl. 7 Ox.	Carbonic Oxide, 1 Carb. 1 Ox. Carbonic Ac., 1 Carb. 2 Ox. Oxalic Ac., 2 Carb. 3 Ox. Sulphurous Ac., 1 S. 2 Ox. Hyposulphurous Ac., 2 S. 2 Ox. Sulphuric Ac., 2 S. 5 Ox. Oxide, 1 Sel. 1 Ox. Selenious Ac., 1 Sel. 2 Ox. Selenic Ac., 1 Sel. 3 Ox. Oxide, 1 Sel. 1 Ox. Selenious Ac., 1 Sel. 2 Ox. Selenic Ac., 1 Sel. 3 Ox. Oxide, 3 Phos. 1 Ox. Hypophosphorous Ac., 2 Phos. 1 Ox. Phosphorous Ac., 2 Phos. 3 Ox. Phos. 5 Ox. Oxide of Iodine, Iodous Ac., Composition unknown. Iodic Ac., 1 Iod. 5 Ox. Periodic Ac., 1 Iod. 7 Ox. Boracic Ac., 1 Bor. 3 Ox. Silicic Ac., 1 Sil. 3 Ox.	Potassa, 1 Pot. 1 Ox. Peroxide of Potassium, 1 Pot. 3 Ox. Soda, 1 Sod. 1 Ox. Peroxide of Sodium, 2 Sod. 3 Ox. Lithia, 1 Lith. 1 Ox.	Protoxide, 1 Ba. 1 Ox. Peroxide, 1 Ba. 2 Ox. Protoxide, 1 Str. 1 Ox. Peroxide, 1 Str. 2 Ox. Lime, 1 Cal. 1 Ox. Peroxide, 1 Cal. 2 Ox. Magnesia, 1 Mag. 1 Ox.
SPECIFI Sp. Gr.	1-102 0-069 0-972 2-47 1-060	1-99 4-3 1-77 4-948 2-738	0.972	Unknown Do. Do.
Equiv.	8 14 36 18	6 16 16 16 16 126 20 22	40 24 8	68 44 12 12
Name.	Oxygen Hydrogen Nitrogen Chlorine Fluorine	Carbon Sulphur Selenium Phosphorus Boron Bromine Silicon	Potassium Sodium Lithium	Barium Strontium Calcium Magnesium
Class.	Simple Gases.	Simple Non Metallic Substances.	Metallic bases of Alkalis.	Metallic bases of Alkaline Earths.

F ć are divided their Nam re into which Simple Bodies a the Cr n ala The following TAD





Aluminum10UnknownAlumina, 2 Alumin. 3 Ox.Glucinum18Do.Glucina, 1 Gl. 3 Ox.Yttrium32Do.Yttria, 1 Ytt. 1 Ox.Thorium60Do.Thorina, Composition unknown.Zirconium22Do.Zirconia, 2 Zir. 3 Ox.	Manganese288-013Protoxide, 1 Ma. 1 Ox.Sesq. Oxide, 2 Ma. 3 Ox.Peroxide, 1 Ma. 2 Ox.Red Oxide, 3 Ma. 4 Ox.ManganicZinc3277Protoxide, 1 Zi. 1 Ox.Peroxide, Composition uncertain.317 Ox.Red Oxide, 3 Ma. 4 Ox.ManganicZinc28777Protoxide, 1 Ir. 1 Ox.Peroxide, 2 Ir. 3 Ox.Black Oxide, 3 Ir. 4 Ox.Codmin.587-91Protoxide, 1 Tin, 1 Ox.Sesq. Oxide, 2 Tin, 3 Ox.Black Oxide, 3 Ir. 4 Ox.587-91Oxide, 1 Cad. 1 Ox.Sesq. Oxide, 2 Tin, 3 Ox.Binoxide, 1 Tin, 2 Ox.587:034Protoxide, 1 Cob. 1 Ox.Deutoxide, 3 Cob. 4 Ox.Peroxide, 2 Cob. 3 Ox.587:034Protoxide, 1 Ni. 1 Ox.Peroxide, 2 Ni. 3 Ox.Peroxide, 2 Cob. 3 Ox.587:034Protoxide, 1 Ni. 1 Ox.Peroxide, 2 Ni. 3 Ox.Peroxide, 2 Cob. 3 Ox.	Arsenic38 5-8845-884 5Arsenious Ac., 2 Ar. 3 Ox. Arsenio Ac., 2 Ar. 5 Ox. Vanadium5 Ox. 5 5 6Arsenious Ac., 2 Ar. 3 Ox. Chromic Ac. 1 Chr. 3 Ox. Vanadium5 Ox. 5 6 6Arsenious Ac., 2 Ar. 3 Ox. Chromic Ac. 1 Chr. 3 Ox. 7 Protoxide, 1 Yan. 1 Ox. Binoxide, 1 Yan. 2 Ox. Vanadic Ac., 1 Yan. 3 Ox. 7 Oulminum5 Ox. 7 Ox. 7 Discribe, 1 Tung, 2 Ox. Blue Oxide, 2 Tung, 3 Ox. 7 Tungstei Ac., 1 Tung, 3 Ox. 7 Tung, 3 Ox. <th>Mercury 202 13:545 Protoxide, 1 Mer. 1 Ox. Peroxide, 1 Mer. 2 Ox. Silver 108 10*51 Oxide, 1 Sil. 1 Ox. Binoxide, 1 Go. 2 Ox. Peroxide, 1 Go. 3 Ox. Platinum 200 19*3 Protoxide, 1 Fla. 1 Ox. Binoxide, 1 Pla. 2 Ox. Sesq. 0xide, 2 Pla. 3 Ox. Paladium 38 21*25 Protoxide, 1 Pall. 1 Ox. Binoxide, 1 Pla. 2 Ox. Sesq. 0xide, 2 Pla. 3 Ox. Paladium 54 11*3 Protoxide, 1 Pall. 1 Ox. Binoxide, 1 Pla. 2 Ox. Sesq. 0xide, 2 Pla. 3 Ox. Rhodium 52 11* Protoxide, 1 Rho. 1 Ox. Binoxide, 1 Pall. 2 Ox. Sesq. 0xide, 2 Pla. 3 Ox. 00 16* 10* 1 Ox. Sesq. 0xide, 2 Ox. Binoxide, 1 Os. 3 Ox. 10 7 Protoxide, 1 Rho. 1 Ox. Sesq. 0xide, 2 Ox. Binoxide, 1 Os. 3 Ox. Peroxide, 1 Os. 3 Ox. 100 7 Protoxide, 1 Ox. Sesq. 0xide, 2 Ox. Binoxide, 1 Os. 3 Ox. Peroxide, 1 Os. 3 Ox. Osmic Ac., 1 Os. 4 Ox. 100 7 Protoxide, 1 Dx. Sesq. 0xide, 2 Ox. Binoxide, 1 Os. 3 Ox. Deroxide, 1 Os.</th>	Mercury 202 13:545 Protoxide, 1 Mer. 1 Ox. Peroxide, 1 Mer. 2 Ox. Silver 108 10*51 Oxide, 1 Sil. 1 Ox. Binoxide, 1 Go. 2 Ox. Peroxide, 1 Go. 3 Ox. Platinum 200 19*3 Protoxide, 1 Fla. 1 Ox. Binoxide, 1 Pla. 2 Ox. Sesq. 0xide, 2 Pla. 3 Ox. Paladium 38 21*25 Protoxide, 1 Pall. 1 Ox. Binoxide, 1 Pla. 2 Ox. Sesq. 0xide, 2 Pla. 3 Ox. Paladium 54 11*3 Protoxide, 1 Pall. 1 Ox. Binoxide, 1 Pla. 2 Ox. Sesq. 0xide, 2 Pla. 3 Ox. Rhodium 52 11* Protoxide, 1 Rho. 1 Ox. Binoxide, 1 Pall. 2 Ox. Sesq. 0xide, 2 Pla. 3 Ox. 00 16* 10* 1 Ox. Sesq. 0xide, 2 Ox. Binoxide, 1 Os. 3 Ox. 10 7 Protoxide, 1 Rho. 1 Ox. Sesq. 0xide, 2 Ox. Binoxide, 1 Os. 3 Ox. Peroxide, 1 Os. 3 Ox. 100 7 Protoxide, 1 Ox. Sesq. 0xide, 2 Ox. Binoxide, 1 Os. 3 Ox. Peroxide, 1 Os. 3 Ox. Osmic Ac., 1 Os. 4 Ox. 100 7 Protoxide, 1 Dx. Sesq. 0xide, 2 Ox. Binoxide, 1 Os. 3 Ox. Deroxide, 1 Os.
Metallic bases of Earths.	Metals which decompose Water at a red heat.	Metals which do not decompose Water at a red heat, and the Oxides of which are not decom- posed by heat alone. Leap un Anti- Leap un de com- Leap un de com- de com- d	Metals whose Oxides are decom- posed by a red heat

TESTS FOR THE MORE COMMON SPECIES OF CALCULI.

1. Uric or lithic acid calculus is of a brownish colour and laminated texture, and has a smooth surface. Before the blowpipe it blackens, giving off an ammoniacal odour, and is gradually consumed, leaving a minute portion of white ash. It is soluble in potassa and in nitric acid, the latter solution yielding on evaporation purpurate of ammonia, known by its fine purple colour;—in water and hydrochloric acid it is sparingly soluble.

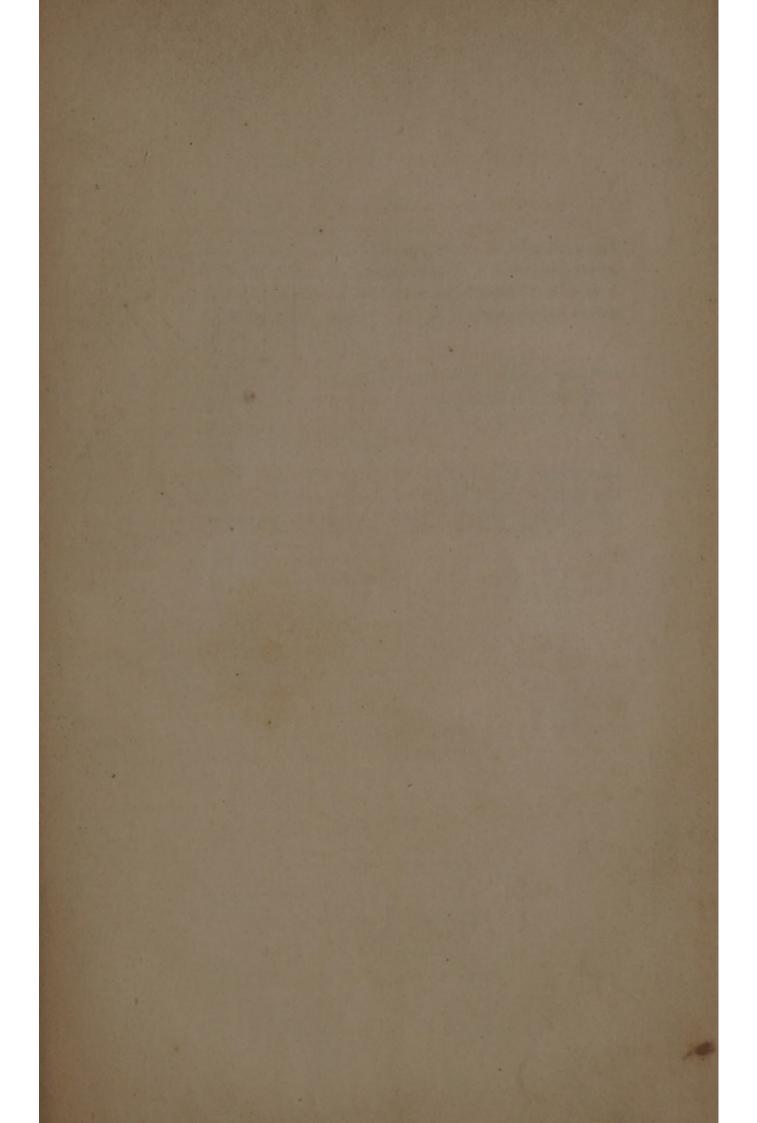
2. Phosphate of lime calculus is of a pale brown colour and loosely laminated texture; its surface is smooth. Before the blowpipe it first blackens and then assumes a white colour, remaining unchanged, unless the heat be very intense, when it fuses;—in potassa it is insoluble, but is soluble in dilute nitric and hydrochloric acids.

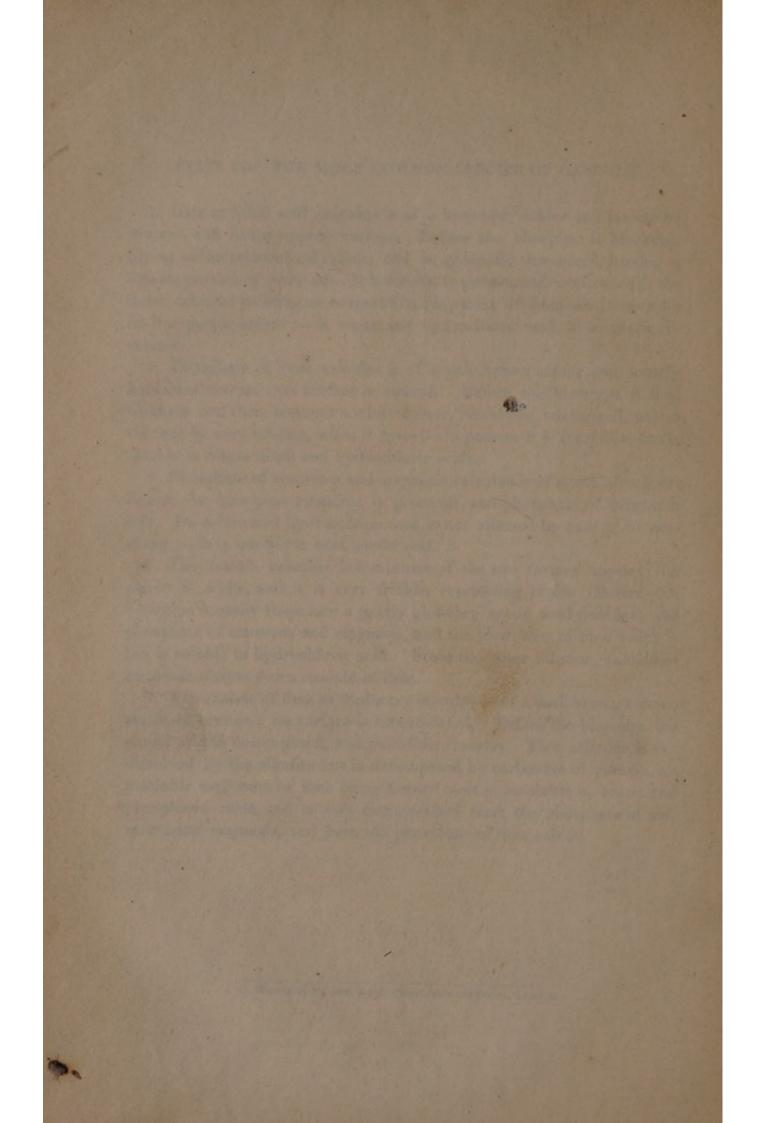
3. Phosphate of ammonia and magnesia calculus is of a whitish colour. Before the blowpipe ammonia is given off, and phosphate of magnesia left. Its solution in hydrochloric acid is not affected by oxalate of ammonia;—it is soluble in cold acetic acid.

4. The fusible calculus is a mixture of the two former species; its colour is white, and it is very friable, resembling chalk. Before the blowpipe it easily fuses into a pearly globule; acetic acid dissolves the phosphate of ammonia and magnesia, and the phosphate of lime which is left is soluble in hydrochloric acid. From the latter solution, oxalate of ammonia throws down oxalate of lime.

5. The oxalate of lime or mulberry calculus is of a dark brown colour, and hard texture; its surface is tuberculated. Before the blowpipe the oxalic acid is decomposed, and pure lime remains. This calculus is not dissolved by the alkalies but is decomposed by carbonate of potassa, an insoluble carbonate of lime being formed ;—it is insoluble in acetic and phosphoric acids, and is thus distinguished from the phosphate of ammonia and magnesia, and from the phosphate of lime calculi.

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