NOTES ON THE ESTIMATION OF CAFFEIN.

BY W. A. PUCKNER.

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SOME time ago Gomberg published a method for the estimation of caffein, by means of Wagner's reagent,¹ wherein appear certain statements from which is to be inferred the superiority of this method over such where the caffein is shaken out of an aqueous solution by means of chloroform, and which, if true, would show that most methods now in use, give low results since but an imperfect separation of caffein is attained. Thus Spencer² is said to have demonstrated the difficulty with which the alkaloid is abstracted from watery solutions, he directing that at *least* seven portions of chloroform be used for this purpose, but offering no proof of the necessity for this departure from the usual direction of shaking out the liquid with three or four portions of the solvent. Spencer is at variance with Allen,³ who investigated this matter and found that from a solution, slightly acidulated with sulphuric acid, one treatment with chloroform removed seventy to eighty-five per cent. of the amount present, while four usually effected complete extraction, especially if toward the end the solution is rendered faintly alkaline.

This agrees well with the results of my own experiments, where anhydrous caffein, in quantities from one-tenth to four-tenths gram, dissolved in fifty cc. one per cent. sulphuric acid, was shaken successively with twenty-five, ten and ten cc. chloroform, the united chloroform solution evaporated at a gentle heat and the residue dried over sulphuric acid to constant weight. In each case the solution was shaken with a further quantity of ten cc. chloroform and the weight of the caffein so extracted ascertained as before.

Caffein taken. Gram.	Residue from first, second and third extraction. Gram.	Residue from fourth extraction. Gram.	Total per cent. recovered.
0.1285	0.1277	0.0004	99.69
0.1852	0.1820	0.0026	99.67
0.1988	0.1980	0.0002	99.69
0.2011	0.1977	0.0025	99.55
0.2559	0.2552	0.0005	99.92
0.4416	0.4355	0.0043	99.58

¹ This Journal, 18, 331.

² J. Anal. Chem., 4, 390.

³ Com. Org. Anal., 3, Part II, 485.

This shows that the extraction of caffein from an aqueous solution presents no difficulties since, even when the solution is quite acid, practically the entire amount is obtained when four portions of chloroform are used ; while, even if the fourth be omitted the results will be sufficiently correct for most purposes.

In the article referred to we are also told, although it is usually stated caffein may be shaken out of an acid solution, since its salts are broken up by water, that this is but relatively true; as a proof thereof the following is offered :

“ 1.0085 grams of caffein were dissolved in sixty cc. of sulphuric acid (1.10), and this solution was repeatedly shaken with chloroform, twenty-five cc. at a time :

Ten consecutive portions of chloroform gave a total of	0.3514	gram	caffein.
Three additional “ “ “ made “ “	0.4859	“	“
Three more “ “ “ “ “ “	0.5034	“	“

Since the degree of dissociation of caffein salts is inversely proportional to the acid strength of the solution, it is to be expected that it will be extremely difficult to shake out the alkalioid from a solution containing so great a quantity of free acid ; but while at times it may be advantageous to extract caffein from a solution having an acid reaction, in no instance would there seem need of a sufficient amount to render the method inapplicable ; further, according to Knox and Prescott¹ Gombert's method becomes uncertain under similar conditions.

In the experiments just quoted ten extractions with chloroform yielded but 34.85 per cent. of the total caffein, or on an average each treatment removed only 3.485 per cent., while the three subsequent treatments removed an additional 13.33 per cent. of the whole, or 4.44 per cent. for each extraction, *i. e.*, although the total substance in solution had been decreased by more than one third the average amount given up to chloroform increased in the 11th, 12th and 13th treatment ; while in the 14th, 15th and 16th but 1.735 per cent., or on an average of 0.578 per cent. for each shaking was obtained.

Although the writer had never attempted a caffein determination under the conditions mentioned, he was, from theoretical considerations, inclined to question the figures given, and accordingly made the following experiments.

¹ Proceedings Am. Pharm. Ass., 1896.

1.0137 gram caffein, rendered anhydrous by keeping in a desiccator over sulphuric acid until its weight remained constant, was dissolved in sixty cc. ten per cent. sulphuric acid and shaken successively with nine portions of chloroform, twenty-five cc. each; the chloroform solutions evaporated at a gentle heat and the residue dried over sulphuric acid to constant weight.

1st portion of twenty-five cc. yielded a residue of	0.5525	gram.
2nd " " " " " " " "	0.2514	"
3rd " " " " " " " "	0.1155	"
4th " " " " " " " "	0.0535	"
5th " " " " " " " "	0.0237	"
6th " " " " " " " "	0.0114	"
7th " " " " " " " "	0.0058	"
8th " " " " " " " "	0.0029	"
9th " " " " " " " "	0.0015	"
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	1.0182 ¹	"

In the second experiment 1.0001 gram anhydrous caffein in sixty cc. ten per cent. sulphuric acid, extracted as before, with chloroform in proportions of twenty-five cc. each :

1st, 2d and 3rd portions gave a total residue of	0.9086	gram.
4th, 5th and 6th " " " " " "	0.0854	"
7th, 8th and 9th " " " " " "	0.0134	"
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	1.0074 ¹	"

The sulphuric acid used in Gomberg's experiments was designated as "(1:10)" by which it is presumed an acid containing ten per cent. by weight of sulphuric acid was meant; since, however, it was possible that sulphuric acid 1:10 *by volume* was the strength of the acid used, a determination was made with an acid with such concentration, *i. e.*, ten cc. concentrated sulphuric acid mixed with water enough to make when cold, 100 cc. In sixty cc. of this were dissolved 0.9790 gram caffein and extracted with chloroform in portions of twenty-five cc. each as before.

1st, 2nd and 3rd portion yielded a total residue of	0.6484	gram.
4th, 5th and 6th " " " " " "	0.2222	"
7th, 8th and 9th " " " " " "	0.0756	"
10th, 11th, 12th, 13th, 14th, 15th and 16th "	0.0379	"
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	0.9841 ¹	"

¹ No explanation is offered to account for the plus error in the above. Contamination with sulphuric acid was suspected, but disproved.

As was to be expected, this confirms in a general way, the statement relative the difficulty with which caffen is shaken out of solutions containing a large proportion of sulphuric acid; in no way, however, does it agree with the data given by Gomberg, who by ten successive treatments with chloroform removed only 34.85 per cent., while my figures show that when a ten per cent. sulphuric acid was used, with but three extractions, fully ninety per cent. was recovered, and even with a still stronger acid (1 + 9 by volume), three portions of chloroform removed about sixty-five per cent.

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CONTRIBUTION TO THE KNOWLEDGE OF THE RUTHENOCYANIDES.

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POTASSIUM ruthenocyanide was described by Claus, in 1854, in his "Beiträge zur Chemie der Platinmetalle." The salt was formed by fusing ammonium rutheninitrosochloride¹ (tetrachloride of Claus) with potassium cyanide. The attempt was also made to form it by fusing potassium ferrocyanide with ruthenium, but it was found impossible to separate the ferrocyanide and ruthenocyanide. It is probable that some of Claus' experiments were carried out with a ruthenocyanide contaminated with ferrocyanide, from the fact that he describes copper ruthenocyanide as brown, whereas, when free from the ferrocyanide, it is pale green. Potassium ruthenocyanide in reactions and crystallization resembles very closely the ferrocyanide, except that when pure it is white. Its crystallography as well as that of the isomorphous ferrocyanide and osmocyanaide are described by A. Dufet.²

Preparation of potassium ruthenocyanide for the purpose of carrying out experiments upon it not yet completed, gave occasion to the work recorded in this paper.

In the Claus method of preparation, a large proportion of the ammonium rutheninitrosochloride is decomposed with separation of metallic ruthenium, and while a part of the ruthenocya-

¹ Joly; *Compt. rend.*, 108, 854, 1889; Howe: *J. Am. Chem. Soc.*, 16, 388, 1894.

² *Compt. rend.*, (1895), 120, 377.