

by rinsing the beaker with 5 to 10 cc. of glacial acetic acid. The acid is transferred to another test-tube for comparison with the first. By holding a sheet of white paper back of the 2 test-tubes the difference in color is made sufficiently pronounced to be seen by a large audience. To preclude the possibility that the color of the second rinsing is merely accentuated by the presence of acetic acid, some of the acid may be added to the water in the first test-tube.

The experiment illustrates not only the fact of adsorption itself, but the effect of the solvent upon the equilibrium, and the rapidity with which adsorption takes place.

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**Qualitative Analysis of the Iron Group.**—The writer has found it feasible to modify the generally accepted analysis of the iron group and finds that the resultant method both shortens the procedure and increases the accuracy of the work of his students.

The following table of results shows that, with proportions rarely met, all separations are quite sharp. The numbers represent milligrams of the ion named.

Zn	Cr	Mn	Fe	Co	Ni
...	..	9	9	1.8	9
...	..	3.6	3.6	3.6	18
...	..	1.3	13.3	0.67	13.3
...	..	2	100	1	100
5	..	..	250	..	..
5	..	250	..	..	..
5	..	250	250	..	..
5	..	500	500	..	..
5	..	500	500	2	3
5	8.5	500	500	2	3
0.5	..	1	..	0.5	5
0.5	..	5	..	5	5
0.5	17	18	20	10	5
2	..	500	500	1	10

The method of W. A. Noyes<sup>1</sup> is used for the separation of aluminum, chromium and zinc, but his method for separating these metals from those of the iron division is modified by the use of a large excess of sodium hydroxide. As the table shows, this gives a sharp enough separation of the zinc to make it unnecessary to search for it in the iron division. No difficulty has been met because of the solution of the iron division with the aluminum division.

<sup>1</sup> Noyes, "Qualitative Analysis," Henry Holt and Co., 1911.

The iron division precipitate is treated directly with nitric acid and potassium chlorate to remove the manganese. (At this point phosphates may be removed if present.) Iron is removed by adding a large excess of ammonium hydroxide. The filtrate from the iron precipitate is evaporated until only a faint odor of ammonia remains and nickel is tested for by adding dimethyl glyoxime directly to  $\frac{1}{8}$  of this solution, no adjustment of hydrogen ion being necessary.

The remainder of the solution is further evaporated to a sirup and transferred to a test-tube; enough water is then added to dissolve the precipitated salts, and cobalt is tested for by a modified Vogel's test; 3 cc. of 10% potassium thiocyanate solution is added, followed by 3 cc. of a mixture of amyl alcohol and ether (1 vol. : 3 vol.) and the tube gently shaken. Cobalt causes the ether layer to become blue-green. If the color is not distinct, the careful addition of a few drops of 12 *N* hydrochloric acid to the ether layer will give the blue-green color if cobalt is present, or a straw-colored or colorless layer, if it is absent. This test is extremely sensitive and can be applied to the original solution if iron is absent.

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## EFFECTS OF THE METHOD OF DESICCATION ON THE CARBOHYDRATES OF PLANT TISSUE<sup>1</sup>

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### Introduction

In determining the composition of plant tissue it is essential that the material undergo a minimum of possible chemical changes from the time the plant is harvested to the time actual analysis is begun. When a large number of samples are involved simultaneously, various mechanical difficulties encountered in extraction render it almost impossible to analyse the fresh tissue directly. Therefore, it is practically necessary to resort to some method of preserving the sample for subsequent analysis.

Two methods have been used to preserve such tissues, namely: (1) desiccation by heating at 85° to 105°, and (2) immersion in conc. alcohol.

<sup>1</sup> Published with the permission of the Director of the Wisconsin Agricultural Experiment Station.