might be caused by a proteolytic enzyme. Accordingly, 1 per cent. solutions of pancreatin (supplied by the makers of Clarase), and of papain were used. Both preparations were ineffective, and actually caused chromosome clumping. The results were very poor when compared with the checks.

Recent studies by Greathouse, Klemme and Barker[2] on the deterioration of cellulose by fungi, suggested that certain of these organisms might be of value in the present problem. Fortunately we were able to secure cultures of *Aspergillus niger* Van Tieghem, *Chaetomium globosum* Kunze, and a species of *Metarrhizium* through the courtesy of G. A. Greathouse of this Bureau. Single flask cultures were extracted 10 to 15 days after inoculation by grinding the contents of the flask with quartz sand in a mortar containing 10 ml of a sodium acetate buffer, pH 5.0. The supernatant liquid was tested on anther sections as previously described for the enzymes. This series of treatments also included a 5 per cent. solution of Clarase in sodium acetate at pH 5.0. All treatments gave beneficial results and produced slides superior to the checks. The cell walls of most pollen mother cells were softened and the cytoplasm partially digested. As a result of these changes the cells were easily flattened and the chromosomes spread. Tests were also made in which the freshly prepared Clarase solution and fungus extracts were heated to boiling prior to use. The slides resulting from material treated in the boiled solutions were definitely inferior to those from the unheated solutions, which indicates that active enzymes were necessary to produce the beneficial effect.

The results herein briefly reported are preliminary to further work under way with other treatments and modes of application. There are many aspects of the problem needing further investigation. For instance, it is not possible as yet to measure the concentration of the fungus extracts used, or to identify the enzyme or enzyme complex that is effective. So far the method has been used on only one species other than Lilium. Dr. A. E. Clarke of this division has used Clarase on the pollen mother cells of an amphidiploid Allium with beneficial results. Some observations have also been made on smears of treated Lilium root tips. These have shown that fungus extracts affect the middle lamella so that the cells separate readily. Further investigations of pre-treated root-tip smears are under way.

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2 G. A. Greathouse, D. E. Klemme and H. D. Barker, *Ind. and Eng. Chem. Anal. Ed.*, 14: 614–620, 1942.

A NYLON BLOOD AND PLASMA FILTER

FOR the last nine months we ran laboratory tests on Nylon blood and plasma filters. Our observations and tests fully coincided with those made by Dr. S. Brandt Rose, *et al.*, as reported in SCIENCE (Vol. 98, No. 2534).

However, we found the sewing of these tiny filters rather cumbersome, and noted that small, but objectionable quantities of Nylon fibers would be entrained in the filtrate.

We took advantage of the thermo-plastic qualities of Nylon and welded the seams rather than sewing them. This method is much faster than sewing and eliminates the shedding of Nylon fibers. Furthermore, the filters can be fabricated with a cone point, and thereby can be utilized for making the drip count.

The following method was found to be satisfactory in making the filters: A double layer of finely textured Nylon cloth was placed on a flat metallic surface. A sheet metal template was made for the filters, and this was placed firmly over the Nylon cloth. The outline of the template was then traced with an electrically heated metal stylus. The stylus of an electric wood-burning set was used for this with excellent results. Flat, colorless, flexible seams were obtained after only a small amount of practice. These seams were tested carefully and were found to be safer for use in transfusion filters than sewn ones.

Prior to using the electric cutting method, experiments with flame cutting were conducted, but the results were found to be too unreliable.

The filters were welded directly to the glass delivery tube of the transfusion assembly.

Some chemical tests were made on the Nylon material. It was found to be soluble in mineral acids and was destroyed by alkalies. It withstood treatment with hydrogen peroxide and solutions of sodium citrate and sodium citrate-dextrose.

The apparatus was developed in the laboratory only, and was not used in giving transfusions to patients.

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