individually or in combinations with one another in chromatography of proteins on hydroxylapatite; often it is difficult to ascertain which buffer cation was used. Our results show the importance of knowing exactly which buffer cation is used.

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## Gas chromatographic analysis of ethylene and some fluoroethylenes

It was of interest in a recent investigation<sup>1</sup> to separate the C<sub>2</sub>-olefius ethylene, 1,1difluoroethylene and tetrafluoroethylene. Silica gel and silver nitrate-ethylene glycol packings show excellent separational properties for hydrocarbons and olefins<sup>2,3</sup>, and in the present work these materials have been used to separate the C2-olefins.

Using silica gel alone it is found that I,I-difluoroethylene is separated from the ethylene and tetrafluoroethylene peak, while silver nitrate-ethylene glycol separates ethylene from the fluoroolefins. The quantitative separation of all three olefins is achieved by using both column materials in series.

## Experimental and results

Silver nitrate in diethylene glycol on a firebrick support was purchased from the Perkin-Elmer Co., as was the silica gel. A Perkin-Elmer gas chromatograph (154-C) was used.

The silver nitrate phase was packed into a 12 ft. length of  $\frac{1}{4}$  in. O.D. aluminium tubing and coiled, while the silica gel was contained in a 3.3 ft. Pyrex glass column;

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| Column used                                       | Relention times (min) |                                 |              |
|---|-----------------------|---------------------------------|--------------|
|   | $C_2 H_4$             | CF <sub>2</sub> CH <sub>2</sub> | $C_{2}F_{4}$ |
| Silver nitrate-ethylene glycol                    | 12.5                  | 7.0                             | 7.0          |
| Silica gel  | 14.2                  | 19.0                            | 14.2         |
| Silver nitrate-ethylene glycol<br>plus silica gel | 30.8                  | 25.6                            | 21.6         |

250

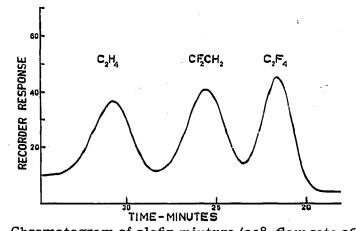


Fig. 1. Chromatogram of olefin mixture (25°, flow rate 36 ml/min).

the latter column was pretreated with nitrogen at 100° before use. Hydrogen was used as a carrier gas at a flow rate of 36 ml/min column temperature was 25°. The olefin mixtures were approximately equimolar and sample size was 0.06 ml.

The rasults of the separation are given in terms of retention times and are recorded in Table I; the recorded chromatograph using the combined columns is shown in Fig. 1.

Retention times may be decreased with some loss in separation by increasing the carrier flow and column temperature. A chromatogram at  $45^{\circ}$  and flow rate of 70 ml/min is shown in Fig. 2.

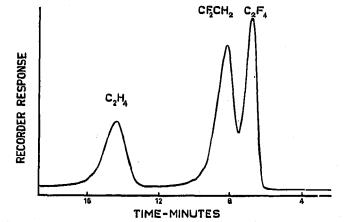


Fig. 2. Chromatogram of olefin mixture (45°, flow rate 70 ml/min).

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