|            | N                                       | lumber        | Type                                         | Mean relative potencies |   |      |            |
|------------|-----------------------------------------|---------------|----------------------------------------------|-------------------------|---|------|------------|
|            |                                         | of<br>ecimens | of<br>response                               | Ā                       | N | PE   | ISO        |
| Vein       | Saphenous<br>circular muscl             | 3<br>e        | Contraction                                  | 4                       | 1 | 0.08 | 0.005      |
| Artery     | Popliteal circular muscl                | 1<br>e        | Contraction                                  | 4                       | 1 | 0.2  |            |
| Oesophagus | Lower third circular musc               | le            | Contraction                                  | 5                       | 1 | 0.2  |            |
| Stomach    | Pyloric antrum<br>longitudina<br>muscle | 4             | Inhibition of acetylcholine induced contract | >1                      | 1 | <1   | ≤1         |
| *1         |                                         | •             |                                              |                         |   |      | 0.005      |
| Ileum      | Longitudinal muscle                     | 3             | Relaxation                                   | 5                       | 1 | 0.2  | 0.005      |
| Colon      | Longitudinal                            | 3             | Relaxation                                   | 3                       | 1 | 0.2  | 0.005      |
|            | muscle                                  |               |                                              | 3                       | 1 |      | 0.04       |
|            | mascie                                  |               |                                              | 10                      | î |      | 0.03       |
| <b>.</b>   | T                                       | •             | TO 1                                         |                         | 1 | ~    |            |
| Rectum     | Longitudinal muscle                     | 3             | Relaxation                                   | 3                       | ı | 0.1  | 0.003      |
|            |                                         |               |                                              | 2                       | 1 | 0.1  | 0.2        |
|            |                                         |               |                                              | 4                       | ī |      | 0.1        |
| Bladder    | Dotmison                                | 1             | Contraction                                  | 2                       | 1 | 0.3  | <b>V</b> 1 |
| Diaguer    | Detrusor                                | 1             | Contraction                                  | 2                       | 1 | 0.3  |            |

TABLE 1. Potencies of adrenaline (A), isoprenaline (ISO), and phenylephrine (PE), relative to noradrenaline (N) as free bases on a number of human tissues containing smooth muscle

-, No response obtained in the concentrations used.

muscle

amechol (0·1 to 5  $\mu$ g/ml.) to induce tone. Relative potencies were determined by the method of Furchgott (1967), the results being shown in Table 1.

In all tissues studied the order of potency was adrenaline, noradrenaline, phenylephrine followed by isoprenaline. In artery and vein the response to these sympathetic amines was contraction (excitation). In gastro-intestinal, smooth muscle the response was relaxation (or inhibition) apart from oesophagus, in which contraction (excitation) occurred. The order of potency obtained in these tissues indicates a significant α-adrenoceptive receptor population (Furchgott, 1967; Ahlquist, 1948).

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## Effects of prostaglandin E<sub>2</sub> (PGE<sub>2</sub>) on the output of histamine and acid in rat gastric secretion induced by pentagastrin or histamine

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The relationship between the output of histamine and acid from the rat stomach has been studied *in vivo* during intravenous infusion of pentagastrin and histamine. The lumen of the stomach was perfused with 0.8% saline (0.2-0.5 ml./min) and samples of perfusate were collected at 15 min intervals. Acid content was estimated by titrating an aliquot of each sample with N/100 sodium hydroxide and the histamine content of the remainder was assayed on isolated superfused ileum from the guinea-pig.

In five out of six experiments with pentagastrin  $(0.05-0.2 \mu g/min intravenously)$  there was a significant correlation between the outputs of histamine and acid. The ratio of histamine (ng base) to acid ( $\mu$ -equiv) in different rats ranged from 0.22 to 1.43 with a mean of 0.57, similar to the ratio of 0.86 reported by Adam & Main (1960) during intravenous infusion of histamine.

Preliminary results suggest that inhibition of pentagastrin-induced acid secretion by PGE<sub>2</sub> ( $0.2-2~\mu g/min$  intravenously) is not accompanied by a corresponding reduction in histamine output. During histamine infusion (5–20  $\mu g/min$  intravenously) PGE<sub>2</sub> inhibited acid secretion but increased histamine output which may indicate an increase either in mucosal blood flow or in permeability.

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## A digital oscillogram analyser used in biological assay and other experimental procedures

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This instrument consists of a micrometer with a travel of 150 mm, linked through

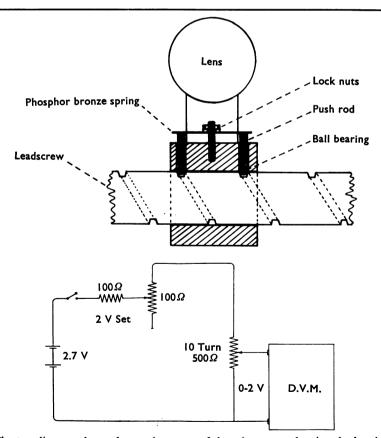


FIG. 1. The top diagram shows the moving parts of the micrometer showing the bearing between the saddle and the rotating rod. The bottom diagram shows the electrical circuit of the oscillogram analyser.