

- tral baroreceptor pathways, *J. Physiol.* 184: 535, 1966.
7. Millar, R. A., and Biscoe, T. J.: Pre-ganglionic sympathetic activity and the effects of anaesthetics, *Brit. J. Anaesth.* 37: 804, 1965.
 8. Bartlestone, H. J., Katz, R. L., and Ngai, S. H.: Effect of cyclopropane on reflexly induced circulatory responses in the dog, *ANESTHESIOLOGY* 27: 756, 1966.
 9. Markee, S., Wang, H. H., and Wang, S. C.: Effects of cyclopropane on the central vasomotor mechanism of the dog, *ANESTHESIOLOGY* 27: 742, 1966.
 10. Ngai, S. H., and Bolme, P.: Effects of anesthetics on circulatory regulatory mechanisms in the dog, *J. Pharmacol. Exp. Ther.* 153: 495, 1966.
 11. Smith, T. C., Colton, E. T., and Dripps, R. D.: Mixed venous-arterial carbon dioxide difference in anesthetized man, *Fed. Proc.* 6: 333, 1967.
 12. Linde, H. W., and Price, H. L.: Gas analyzer for rapid estimation of cyclopropane, *ANESTHESIOLOGY* 19: 757, 1958.
 13. Kahn, N., and Mills, E.: Centrally evoked sympathetic discharge: A functional study of medullary vasomotor areas, *J. Physiol.* 131: 339, 1967.
 14. Ranson, S. W., and Billingsley, P. R.: Affected spinal paths and the vasomotor reflexes. Studies in vasomotor reflex arcs, VI. *Amer. J. Physiol.* 42: 16, 1916.
 15. Price, H. L., King, B. D., Elder, J. D., Libben, B. H., and Dripps, R. D.: Circulatory effects of raised airway pressure during cyclopropane anesthesia in man, *J. Clin. Invest.* 30: 1243, 1951.
 16. Price, H. L., and Helrich, M.: The effects of cyclopropane, diethyl ether, nitrous oxide, thiopental and hydrogen ion concentration on the myocardial function of the dog heart-lung preparation, *J. Pharmacol. Exp. Ther.* 115: 206, 1955.
 17. Price, H. L., Jones, R. E., Deutsch, S., and Linde, H. W.: Ventricular function and autonomic nervous activity during cyclopropane anesthesia in man, *J. Clin. Invest.* 41: 604, 1962.
 18. Moe, G. K., Malton, S. D., Rennick, B. E., and Freyburger, W. A.: The role of arterial pressure in the induction of idioventricular rhythms under cyclopropane anesthesia, *J. Pharmacol. Exp. Ther.* 94: 319, 1948.

Drugs

MAO INHIBITORS Monoamine oxidase inhibitors potentiate the effects of dietary and injected tyramine and other sympathomimetic amines. Studies in various rat and cat preparations have indicated that MAO inhibitors act by retarding the binding and/or breakdown of these sympathomimetic amines by liver microsomal enzyme systems, allowing higher blood levels of these amines. This leads to the release of more endogenous catecholamines from storage sites. Other enzymes (diamine oxidase, choline oxidase) are also inhibited, thus potentiating the action of barbiturates, amphetamine, and morphine and its congeners. Since destruction by catechol-o-methyltransferase and uptake into storage sites terminate catecholamine action, MAO inhibitors do not potentiate the action of exogenous adrenalin or noradrenalin. (Rand, M. J., and Trinker, F. R.: *The Mechanism of the Augmentation of Responses to Indirectly-acting Sympathomimetic Amines by Monoamine Oxidase Inhibitors*, *Brit. J. Pharmacol.* 33: 287 (June) 1968.)