

described in cats and rabbits during lidocaine infusion (ANESTHESIOLOGY 28: 155, 1967) were not observed. Although no seizure focus was evident, spike and slow-wave complexes were recorded in cortical and depth leads and were quite evident in the amygdala and caudate nuclei. (Supported in part by USPHS Grants NB-07844 and FR00169.)

The Effects of Acidosis on Glycolysis and Pentose Shunt in Skeletal Muscle. G. C. NAHAS, M.D., J. KYPSON, M.D., L. TRENER, M.D., and Y. VULLIEMOZ, M.S., *Department of Anesthesiology, College of Physicians and Surgeons, Columbia University, New York, N. Y.* Acidosis *in vivo* produces marked metabolic and functional alterations. The purpose of the present study was to investigate *in vitro* the effect of acidosis on carbohydrate metabolism. **Methods and Results:** Rat diaphragm was incubated in Krebs-Ringer phosphate buffer for one hour at pH 7.4 or 6.8. At pH 6.8 there were significant decreases in lactate (-36 per cent) and pyruvate (-22 per cent) production in the medium. Concurrently, glycogen content was significantly increased (+25 per cent). In a second series, the site of action of acid pH of muscle on glycolysis was investigated by comparing the concentrations of intermediates of the glycolytic pathway in diaphragm incubated at pH 7.4 or 6.8. At pH 6.8 glucose-6-phosphate was increased significantly (+22 per cent); fructose-6-phosphate was consistently, but not significantly, higher. Fructose-1,6-diphosphate was decreased significantly (-30 per cent), indicating a decrease in phosphofructokinase activity. Similar changes in intermediates also occurred when muscle was incubated with ouabain 10^{-4} M in medium at pH 7.4 or pH 6.8. There was no significant change in citrate concentration or O_2 uptake, which might indicate that the Krebs cycle was not affected by pH or by ouabain. Glucose-6-phosphate dehydrogenase activity was increased by 100 per cent at pH 6.8 (with or without ouabain), indicating an increase in activity of the pentose shunt. Ouabain at pH 7.4 did not change the activity of this enzyme. Decrease in phosphofructokinase activity at low pH could account for the decrease in lactate production. Maintenance

of the Krebs cycle could account, in part, for decrease in pyruvate. Marked increase in pentose shunt by acid pH, with resulting increase in TPNH, might be a compensatory mechanism in the presence of decreased glycolysis. **Summary:** Acid pH produces marked alteration in some of the key enzymes which regulate carbohydrate metabolism.

The Effects of Halothane on Mitosis. J. F. NUNN, PH.D., F.F.A.R.C.S., K. L. DIXON and J. D. LOVIS, B.Sc., PH.D., *Departments of Anesthesia and Botany, University of Leeds, Leeds, Yorkshire, England.* In cultures of the ciliate *Tetrahymena pyriformis*, halothane rapidly causes inhibition of growth, the 50 per cent effective dose being about 1.25 per cent of normal atmospheric pressure at room temperature. The rapidity of action suggested interference with mitosis, although inhibition of DNA synthesis cannot be excluded. **Methods:** Studies of mitosis were carried out on root-tips of *Vicia faba* (the broad bean) exposed for periods of two to 24 hours to concentration of halothane (prepared by volumetric dilution of saturated vapor) ranging from 0.5 to 2.0 per cent. Germinating beans were exposed to halothane when lateral roots had attained length of more than one cm. Experimental treatments were carried out in a desiccator which was continuously swept with vapor at a flow rate of about 250 ml/min. The roots were completely exposed to the vapor, their tips not being immersed in water. Controls remained exposed to room air or to air in contact with water containing crystals of thymol, some of which floated. At certain intervals after the commencement of treatment lateral roots were removed and fixed in acetic alcohol. The chromosomes were stained by treatment of the root-tips with the Feulgen reagent following controlled acid hydrolysis. Permanent squash preparations were then made and examined by light microscopy. **Results:** Four hours' exposure to 2 per cent halothane resulted in a very marked reduction in the proportion of the later mitotic stages, anaphase and telophase. Prophases were normal in appearance, but the great majority of mid-division figures showed abnormalities entirely comparable to those associated with exposure to