Lactose Absorption in the Premature Infant*

EUGENE C. JARRETT and GERALD H. HOLMAN
From the Department of Pediatrics, Medical College of Georgia, Augusta, Georgia, U.S.A.

In 1921, while addressing the American Pediatric Society, John Howland described quite vividly the syndromes now recognized as congenital and acquired small intestinal disaccharidase deficiency. In his opening paragraph he stated that he hoped, 'to stimulate an interest to inquire more deeply into a subject on which our present knowledge is disappointingly scanty'. Yet it was not until 37 years later that attention was again brought to bear on this subject after Durand (1958) described a case of lactosuria and chronic diarrhoea, and Holzel, Schwarz, and Sutcliffe (1959) described two cases of suspected lactase deficiency. The only study to be published in this 37-year period is by Fox (1950) who reported sucrose absorption problems in sprue. Since 1959 however, several case reports and other studies related to the intestinal disaccharidases have been published (Weijers, van de Kamer, Dicke, and Ijsseling, 1961; Holzel, Mereu, and Thomson, 1962; Weijers, and van de Kamer, 1962; Auricchio, Dahlqvist, Mürset, and Prader, 1963; Anderson, Messer, Townley, and Freeman, 1963; Sunshine and Kretchmer, 1963; Cozzetto, 1963; Burgess, Levin, Mahalanabis, and Tonge, 1964; Sunshine and Kretchmer, 1964; Lifshitz and Holman, 1964; Davidson, Sobel, Kugler, and Prader, 1964; Auricchio, Rubino, Prader, Rey, Jos, Frézal, and Davidson, 1965b; Lifshitz, Klotz, and Holman, 1965; Burke, Kerry, and Anderson, 1965; Townley, Khaw, and Schwachman, 1965).

The purpose of this study was to determine if the premature infant, at 2 weeks of age, possessed normal intestinal disaccharidase activity as measured by oral tolerance tests. In addition, the question of whether or not there is substrate induction of disaccharidase activity in humans, as there appears to be in some animals (Fischer, 1957), was investigated.

Material and Methods

Twenty premature infants were divided at random into two groups at the time of their birth. One group was given a formula containing lactose as the only disaccharide*. The other group was given a formula containing sucrose as the principal disaccharide† (Bernstein, 1965) until after the disaccharide tolerance tests were performed, and then this group was given the lactose-containing formula. Included in the study were two and a half sets of twins and one set of triplets. All infants were cared for in the premature intensive care unit, and any infant who had any significant difficulty, such as pneumonia or the respiratory distress syndrome, was excluded from the study.

Lactose, sucrose, and maltose tolerance tests were performed between 13 and 17 days of age on each infant by giving an oral load of 1·75 g./kg. of the sugar in question after a 4-hour fast. True blood glucose was determined while fasting just before ingestion of the sugar and at 30, 60, and 90 minutes after ingestion, using an ultramicro method (Keston and Teller, 1965) on heel prick blood.

Results

Twenty infants were studied, 10 in each group. There were 10 females and 10 males with both sexes equally divided between the two groups. In the lactose formula group, there were 7 Negro infants and 3 Caucasians, and this group had a birthweight range of 1250 to 1940 g. with a mean birthweight of 1571·3 g. (SD 82·6). In the sucrose formula group, there were 9 Negro infants and one Caucasian, and this group had a birthweight range of 1170 to 2000 g. with a mean birthweight of 1633·5 g. (SD 85·2)‡. There is no significant difference between these two means.

The mean change in blood glucose with each tolerance test is illustrated in Fig. 1-3. As can be seen, there is very little difference between the groups during the maltose tolerance. Even though the group that was fed the lactose-free formula had a smaller mean blood glucose rise during the lactose and sucrose tolerance tests, the rise is significant and is within limits of normal using the criteria of Durand (1964). There is no statistically significant

Received February 4, 1966.
* Supported in Part by Grant HD-01483-02.

* Lactose formula—SMA S26 (R) (Wyeth).
† Sucrose formula—Nutramigen (R) (Mead Johnson).
‡ Tabulated details of individual subjects are available on request to the authors.
difference between the values obtained at comparative times between the two formula groups for any of the tolerance tests.

Comment

It has been shown quite clearly that the small intestinal disaccharidases are intracellular enzymes, and actually are located in the brush border portion of the small intestinal mucosal cells (Dahlqvist, Auricchio, Semenza, and Prader, 1963; Doell and Kretchmer, 1962; Miller and Crane, 1961a, b). Auricchio, Rubino, and Mürset (1965a) have shown that the $\alpha$-glycosidases, maltase, sucrase, and others, can be found in human intestinal mucosal cells as early as the third foetal month, and that their activity increases thereafter until adult levels are reached between the sixth and eighth foetal months. The $\beta$-glycosidases, lactase and cellobiase, develop slightly later in intrauterine time and do not reach maximal activity until the end of normal gestation.

The rat, which has been used in much experimental work in this area, is born with a high level of lactase activity: however, this activity virtually disappears when the rat is weaned. Fischer (1957) has shown that if the rat is given a lactose load orally after the disappearance of his lactase activity, fermentative diarrhoea develops. If this lactose loading is continued, however, the rat will again develop sufficient lactase activity to hydrolyse the administered lactose. It was then postulated by some (Herzenberg and Herzenberg, 1959) that the intestinal lactase activity of the human infant might be substrate induced by the lactose present in the diet, or at least the presence of the lactose might increase the lactase activity.

The data presented in this paper illustrate that intestinal lactase, sucrase, and maltase activity is normal in the premature infant by 14 days of age, as measured by oral disaccharide tolerance tests. There is no indication of induction of lactase activity, since the infants who had relatively
lactose-free diets had blood glucose rises equivalent to those of the infants who had been fed lactose. The same findings were noted in regard to sucrase absorption in those infants with and without sucrase in the diet. These observations are supported by the findings of Barnett and Beard (1965), reported in a recent study on the ability of the newborn to utilize maltose. It may be possible that the small quantity of lactose in the low-lactose formula was sufficient to induce enzymatic activity, and further studies with formulas entirely free of lactose will be necessary to clarify this problem.

**Summary**

Oral lactose, sucrase, and maltose tolerance tests were performed on 20 premature infants at approximately 14 days of age. 10 infants had been on diets which contained lactose and 10 infants had been on relatively lactose-free diets containing sucrose as the principal disaccharide. The groups were the same regarding sex distribution, and there was no significant difference between the mean birth weights of the two groups.

Data are presented illustrating that lactase, sucrase, and maltase activities are normal in the premature infant at 14 days of age, as measured by oral disaccharide tolerance tests. It is also shown that the presence of the substrate sugar in the diet preceding the test did not appear to induce or enhance the absorption activity of lactase or sucrase as measured by oral tolerance tests.

**References**


— (1964). Disorders Due to Intestinal Defective Carbohydrate Digestion and Absorption, p. 71. II Pensiero Scientifico, Rome.


—, and —— (1961b). The digestive function of the epithelium of the small intestine. II. Localization of the disaccharide hydrolysis in the isolated brush border portion of intestinal epithelial cells. *ibid.*, 52, 293.


