

Determination of the Volume and Weight of Living Animals

WHEN accurate measurements of the weight of aquatic animals are necessary, most authors remove the adherent water by towelling the body with filter paper and weigh quickly and repeat until a constant weight is obtained. The method is defective since, apart from possible damage to the animals, the degree of dryness reached may not be the same each time. These difficulties increase when the animals are small and several have to be used at a time. Similarly, determinations of volume necessitate an elaborate procedure or the use of complicated apparatus.

For small marine Crustacea I find the following procedure better than the volumometer which I described¹. A basket is suspended by a nylon thread from one arm of a balance and immersed in a definite volume of sea-water in a beaker. The apparent weight of the basket and the thread is determined. The animals are then transferred to the basket and weighed. Next, a fourth of the sea water in the beaker is removed and distilled water is added, stirred and the animals weighed again. The animals are removed and the empty basket is weighed in the 75 per cent sea-water. The difference in apparent weights of the animals in sea-water of two densities will thus help to calculate the volume of sea-water displaced and also the weight in air.

It has been found that determinations of apparent weight are not vitiated by osmotic transport, since most Crustacea take more than 10 min. to respond to changes in media. This method is particularly suitable for making continuous observations on the rate of water transport in animals left in anisotonic media.

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¹Gnanamuthu, C. P., *Nature*, **170**, 587 (1952).

β -Galactosidase Changes in the Developing Intestinal Tract of the Rat

IN some micro-organisms β -galactosidase (lactase) is an adaptive enzyme, that is, its biosynthesis is induced by external agents such as lactose or lactose-related substances¹. Apparently this is not the case for the intestinal β -galactosidase of the rat, since a lactose-rich diet does not modify the intestinal specific activity². In the work reported here, the behaviour of this enzyme has been investigated in the albino rat, starting from the last third of the intra-uterine life until adult age.

Homogenates of the small intestine from the duodenal flexure to the ileo-caecal valve were used. The enzyme was determined by the *ortho*-nitrophenol released from the *ortho*-nitrophenyl- β -galactoside, at 38° C. and pH 3.5⁴. The age of foetuses was estimated according to the histo-differentiation shown in the duodenal, ileo-caecal and umbilical regions³, independently of the biochemical values. This technique allows a very close correlation between histological and biochemical data.

The changes in β -galactosidase activity during the end of pregnancy, new-born period and adult age of the rat are plotted in Fig. 1.

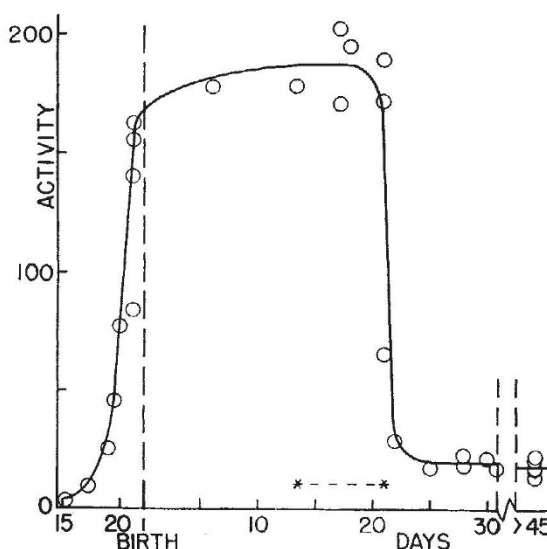


Fig. 1. Intestinal β -galactosidase activity expressed as μ moles of *ortho*-nitrophenol released per mgm. of tissue (wet weight) in 1 hr. *---* Transition from lactation to adult diet; gestation, 22 days

The enzyme activity was also examined in different physiological and dietetic conditions: in adults on a lactose-rich diet, in pregnant rats, in nursing mothers, and in young rats kept after weaning on a complete diet with lactose (25–30 per cent) as the only carbohydrate (lactose-rich diet). The results are recorded in Table 1 together with the values for the normal adult and suckling rats.

Table 1. INTESTINAL β -GALACTOSIDASE ACTIVITY EXPRESSED AS μ MOLES *ortho*-NITROPHENOL RELEASED PER MG. OF TISSUE (WET WEIGHT) PER 1 HR.

Diet	Condition	No. of animals	β -galactosidase activity	Increase
Normal	Adult	5	18 \pm 2.9	$\times 1.00$
Normal	Pregnant	4	18.6 \pm 1.0	$\times 1.03$
Normal	Nursing	3	18.4 \pm 2.4	$\times 1.02$
Mother's milk	1–20 days old	14	183 \pm 8.8	$\times 10.0$
Lactose*	Adult	3	19.3 \pm 2.1	$\times 1.07$
Lactose	25–27 days old	3	19.8 \pm 3.0	$\times 1.10$

* 'Lactose' means a complete diet with lactose 25–30 per cent as the only carbohydrate.

The features which became apparent from these results are: (1) The specific activity does not change on the diet rich in lactose (no induction in a strict sense), confirming discoveries of other authors². (2) There are no significant changes either in the pregnancy or in the nursing period. (3) Activity increases rapidly at the end of foetal life until birth, parallel to the very fast histo-differentiation of the digestive tract³. The high level remains rather constant in the new-born. (4) At the time of weaning (21–22 days) the activity decreases abruptly to the low adult-value. (5) This net decrease in the activity also occurs when the new-born is kept on the lactose-rich diet.

Several authors have shown that lactose hydrolysis is necessarily previous to its intestinal absorption; the somewhat low β -galactosidase adult activity makes its action a limiting step in lactose assimilation, and therefore laxation and diarrhoea occur². An activity ten times higher in the new-born, as compared with the adult animal, is in agreement with the expected great efficiency as regards milk lactose assimilation in the sucklings. The 'physiological