Splenic function in childhood coeliac disease*

G R CORAZZA†, R LAZZARI, M FRISONI, A COLLINA, and G GASBARRINI

From the Third Department of Medical Pathology and Paediatrics, Policlinico S. Orsola, Bologna University, Bologna, Italy

SUMMARY We measured splenic function using a simple, non-isotopic method in childhood coeliac disease. No patients were shown to have hyposplenism. This has important clinical and therapeutic implications.

Hyposplenism is a well-recognised feature of adult coeliac disease1,2 and it has recently been suggested that it is the cause of the autoimmune disorders frequently associated with coeliac disease.3 Although the risk of infection connected with splenectomy or impaired splenic function is particularly severe in children,4 only scanty and inadequate information on splenic function in childhood coeliac disease is available.

We measured splenic function in coeliac children by a new test,5 proved to be useful in adult coeliac patients6 and based on the observation of red cell membrane abnormalities under interference-phase microscopy.

Methods

PATIENTS

Thirty-seven patients (aged 8 months to 13 years) with flat jejunal mucosa were studied. Ten untreated patients who later showed a clear clinical remission after gluten withdrawal and eight patients with a deterioration of the mucosa during gluten challenge were regarded as active coeliac patients. Ten patients on a gluten-free diet who showed a histological improvement at the second biopsy and nine children who had met all the ESPGAN criteria for diagnosis7 were regarded as inactive coeliac patients. The controls comprised 19 healthy children and 25 patients with minor gastrointestinal disturbances matched for sex and age with the coeliac patients. Thirteen patients (aged 5 to 15 years) who had undergone an elective splenectomy during the staging procedure of Hodgkin’s disease were also studied.

SPLENIC FUNCTION MEASUREMENT

A drop of venous blood was mixed with a buffered glutaraldehyde solution; then 2000 red blood cells were examined blind under interference-phase microscopy. The number of cells showing one or more surface pits (pitted red cells) was expressed as a percentage and considered as a measure of splenic function.5

STATISTICS

The percentage of pitted red cells in coeliac children, splenectomised patients, and controls were compared using Wilcoxon’s rank sum test.

Results

No significant difference was found between coeliac patients (mean 0.74%, range 0–2.7%) and controls (mean 0.79%, range 0–1.8%) or in the coeliac group between active (mean 0.74%, range 0–2.1%) and inactive (mean 0.74%, range 0.1–2.7%) patients. Values for splenectomised patients (mean 25.25%, range 15–31%) were significantly higher than in coeliac patients or controls (p<0.001) (Figure).

Discussion

The risk of infection connected with functional or surgical asplenia4 and the finding at necropsy of splenic atrophy in two children thought to be coeliacs,8 9 prompted us to evaluate carefully splenic function in childhood coeliac disease. The study of splenic function in coeliac children has been hampered by the lack of a non-radioisotopic method. The percentage of pitted red cells has been shown to correlate with splenic function as measured by heat-damaged red cell clearance,6 and therefore this method is ideally suited for investi-
gation of splenic function in children. By counting pitted cells we have shown that splenic function in both active and inactive coeliac children was not depressed when compared with controls and splenectomised patients. These results confirm and extend previous findings by McCarthy et al.10 who, by counting Howell-Jolly bodies, were not able to show any lymphoreticular dysfunction in 29 young coeliac patients. The incidence of hyposplenism in adult treated coeliacs seems to be related to the duration of exposure to gluten.6 Thus it is possible to speculate that the immunological abnormalities of adult coeliac disease have not been present long enough in childhood coeliac disease to produce hyposplenism. Our results therefore suggest that splenic hypofunction is not a complication of childhood coeliac disease and that young coeliacs, even in a poor nutritional state, do not need any particular protection against the risks of hyposplenism.

Counting of pitted red cells is an easily repeatable test of splenic function and we intend to follow up a group of coeliac children in order to gain further information on the timing of the development of hyposplenism and its possible connection with other immunological abnormalities of coeliac disease.

References