IgE and IgG4 Antibodies in Subjects Reinfected with *Schistosoma mansoni* in an Endemic Area of Northeast Brazil

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Analysis of the immune response of resistant and susceptible subjects in endemic areas indicated that resistance was associated with enhanced anti-parasite-IgE levels and that reinfection occurred when patients were producing high levels of antibodies that could compete with IgE (AE Butterworth et al. 1985 *Trans R Soc Trop Med Hyg* 79: 393-408, P Hagan et al. 1987 *Trans R Soc Trop Med Hyg* 81: 938-946, P Rihet et al. 1991 *Eur J Immunol* 21: 2679-2686, DW Dunne et al. 1992 *Eur J Immunol* 22: 1483-1494). The results of these reinfection studies suggest that acquired immunity develops slowly with age and that, although other factors can not be excluded, IgE specific antibodies play an important role in anti-schistosome resistance.

Recently, the role of different factors involved in *Schistosoma mansoni* infection (including the nutritional status of the population) was evaluated by E Coutinho et al. (1997 *Mem Inst Oswaldo Cruz* 92: 710-715) in a population living in two contiguous endemic villages (Itapinassu and SãJoaoquim), in northeast Brazil. The patients were identified by stool examinations (WA Hoffman et al. 1934 *Puerto Rico J Publ Hlth Trop Med* 9: 626-653, N Katz et al. 1972 *Rev Inst Med Trop São Paulo* 14: 397-400) and the intensity of infection was classified as light (<100 epg), moderate (101-400) and severe (>400 epg). All patients positive for *S. mansoni* were treated with oxamniquine in a single dose (15 mg/kg for adults and 20 mg/kg for patients under 15 years old). A previous therapy against other helminth infections was carried out with mebendazole and/or thiabendazole (Coutinho et al. *loc. cit.*) before starting the study.

In the present communication we evaluated the influence of the IgE and IgG4 levels on the resistance and susceptibility to infection by *S. mansoni* of 141 patients living in the area mentioned above.

Blood was obtained by venipuncture, six months after treatment, and serum was stored at -20°C until use. Soluble worm antigen preparation (SWAP) and soluble egg antigen (SEA) were prepared using standard procedures (DG Colley et al. 1977 *Int Arch Allergy Appl Immunol* 53: 420-443, G Gazzinelli et al. 1983 *J Immunol* 130: 2891-2895). Antigen preparations were dialysed against distilled water and the protein concentration determined by the method of OH Lowry et al. (1951 *J Biol Chem* 193: 263-275). IgE and IgG4 specific antibody response to SWAP and SEA was evaluated by ELISA (P Hagan 1991 *Nature* 349: 243-245). Each antigen was diluted in 0.05 M Na₂CO₃ buffer, pH 9.6, onto flat-bottomed microtitre plates at the optimum concentration determined by chequerboard titration using pooled positive and negative control sera. The levels of specific IgE and IgG4 bound by these antigens was determined using mouse monoclonal anti-human IgE and IgG4 (Fc fragments). Assays were developed using a horseradish peroxidase-conjugated rabbit anti-mouse IgG. Optimum concentrations of all reagents were determined by titration. Statistical analysis was performed using the software EPIINFO V 6.03. Analysis of variance was employed to determine the differences between frequencies; p<0.05 was considered to be statistically significant.

In the present note only the results concerning reinfection after treatment are mentioned. Reinfection after treatment was moderate and severe in young adults while in subjects older than 35 years, the intensities were lighter (data not shown). An association between age and levels of IgE schistosome-specific antibodies was found. IgE levels to SWAP antigen were maximal in the 35+ age group (p<0.05) and the age-IgE profile follows that expected for an antibody involved in resistance to infection. The IgE levels to SEA antigen was also presented. The statistical analysis revealed that the levels of specific IgE and IgG4 antibodies to SWAP and SEA were significantly higher in the 35+ age group compared to the 15-35 age group. The results suggest that the acquisition of resistance to *S. mansoni* infection is associated with the development of specific IgE and IgG4 antibodies, and that the levels of these antibodies are influenced by age.

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slightly more elevated in the 35+ age group although this was not significant (Fig. 1). Similar results were found by Hagan (1991 *loc. cit.*). The stratified analysis of the IgG4 levels to SWAP and SEA was not well defined and conclusive. IgG4 levels were more elevated to SEA and SWAP in the young adults and in the 35+ age group although this was not statistically significant (Fig. 2). Taken into account the pattern of age related intensity of infection (high in young adults, low in adults), the present results suggest that the levels of IgE antibodies to SWAP may be used as a marker of resistance against schistosomiasis, supporting results from other groups (Hagan 1991 *loc. cit.*, DW Dunne et al. 1997 *Parasite Immunol* 19: 79-89).

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