
The impact of hodgkins disease on the immune system

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Sixty children below the age of 14 years with confirmed diagnosis of H.D both clinically and laboratory were studied. They presented at the Pediatric Clinic of National Cancer Institute, Cairo University. They were divided into three groups: Group 1. Twenty children with HD seen at diagnosis and before starting treatment. Twenty children with HD under treatment. Twenty children Studied in complete remission and after ending therapy. Group 4. Twenty children were healthy control. Lymphadenopathy was the most prominent clinical finding in 88.4%. It was more prevalent in the cervical group of lymph nodes (68.33%). B-symptoms was present in 40% of all patients. The height was severely reduced in patients under relapse therapy compared to the control. The percentile of height was significantly decreased in all studied group compared to the controls. The mean percentile of weight was significantly diminished in all studied groups compared to controls. This decrease was more apparent in relapsed patients. Hepatomegaly was found in 26.66% with a mean size of 1.275 ± 1.832 cm with a maximum of 5 cm below the costal margin. It was decreased by treatment, However this decrease did not reach significance. Splenomegaly was found in 20% at presentation while only one case showed splenomegaly at sample due to splenectomy. Combined therapy reduced the enlarged lymph nodes significantly from a mean 4.3 ± 2.623 cm at presentation to 1.25 ± 1.034 cm at sample. Infectious complication of therapy including viral, fungal, bacterial infections was present in 15%, 15%, 55% respectively. Anergy to tuberculin skin test was present in 15/20 (75%) of pretreatment children, 9/10 (90%) and 10/10 (100%) of relapsing children 70% in remission group. This high frequency of anergy was statistically significant compared to controls. The Anergy was present in 7/8 (87.5%) of patients in stage III & IV and 7/12 (58.3%) in stage I & II .. This higher percentage in stage III & IV does not reach statistical significance. The mean indurated area to tuberculin test was significantly reduced in all patients compared to controls. The mean Hemoglobin percent was significantly decreased in pretreatment and relapsing compared to controls patients revealing hypochromic microcytic anaemia in those children. Leukopenia was found in all patients compared to controls. It was more severe in patients under treatment. Relative lymphocytopenia was found in 13/60 (21.6%) while relative lymphocytosis in 20/60 (20%). The lymphocytopenia was statistically significant in relation to controls. Relative eosinophilia was present in patients before treatment, in relapse and in remission group. It was

statistically significant. Significant monocytosis was found in the same groups of eosinophilia. Absolute lymphocytopenia was significant in all studied groups compared to controls. It was more prevalent in patients during treatment. The E-R active (T-cell) was significantly reduced in all studied groups compared with the control group. At diagnosis, the mean ER was significantly reduced in relation to controls, but it was significantly higher than patients under treatment. It also was significantly lower than in remission group. The reduction in E-R active in relapsed patients was the worst of all groups in relation to controls.¹³² The mean percentage of lymphoblastic transformation in response to PHA was statistically significantly reduced in relation to control; this reduction was prevalent in relapsed groups than other groups. It was also more prevalent in patient with advanced disease (stage III and IV) than in patients with early disease (stage I & II). The Mean E-A (B-cell) active percentage was normal in untreated patients compared to controls. It was severely depressed in relapsing patients due to repeated therapy. The mean E-A active was little higher in untreated children compared to patients under treatment whereas, it was nearly the same as remission level. The absolute B-cells count (active) was significantly reduced in all studied children with HD compared to controls. The reduction of the absolute B-cell count was more prevalent in children under treatment. Serum alkaline phosphatase was significantly elevated in all patients studied in relation to the controls. Relapse group showed the highest level compared to the controls. SGPT was significantly elevated in all groups studied compared to the controls. It was strikingly elevated in patients during treatment. The mean ESR showed a significant elevation in all groups of HD during the first & Second hours in relation to controls. The untreated children had the highest ESR compared to the controls. In conclusion HD causes reduced cellular immunity in -all affected children manifested by anergy to tuberculin skin test, diminished T Cell functions (decreased E-R and decreased lymphoblastic transformation in response PHA) in the different phases of H.D; at diagnosis, during treatment, in relapse and in remission with preserved B-cell functions. Therapy causes further impairment of the cellular immune defects. All deficient functions of T-cells persisted during remission except the delayed skin reactivity which shows improvement.