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PLASMA LEVELS OF PROTEINS OF THE ALTERNATIVE COMPLEMENT PATHWAY IN INBRED MICE THAT DIFFER IN RESISTANCE TO *TRYPANOSOMA CONGOLENSE* INFECTIONS

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Abstract

Inbred BALB/c, A/J, and CS7BII6J mice were infected with Trypanosoma congo/ense (Trans Mara strain). clone TC 13, and monitored for parasitemia, survival times, and plasma levels of complement components C3, CS, factor B, and factor H. Parasitemia was highest in BALB/c, intermediate in NJ, and lowest in CS7BV 6J mice. The mean survival times were II.S \pm 0.9, 23:8 \pm 2.3, and 119 \pm 26 days for BALB/c, NJ, and CS7BV6J mice, respectively. Preinfection levels of factor H were significantly correlated with survival times (r = 0.7722, P < 0.00I). Marked differences were observed between the plasma levels of C3, factor B, and factor H in the 3 mouse strains following infection. Complement CS levels showed the fewest changes. In the initial postinfection period, BALB/c mice had highest increases in the levels of the 4 complement proteins but also had the greatest declines toward the end of the infection. Factor H levels showed a biphasic increase in BALB/c and C57B1I6J, but not in NJ mice, with peaks at days 3 and 9. Complement C3 levels declined in all mice toward the terminal stage of the disease. In the late stages of infection, factor B levels markedly decreased in BALB/c but significantly increased in CS7BII6J mice. Factor B levels measured at the terminal stages in BALB/c, NJ, and CS7BII6J were correlated positively with their respective survival times (r = 0.714, P < 0.01).

The results suggest that genetic differences in the alternative complement pathway might affect the resistance to *T. congolense* infections.