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Studies on Antibodies Accompanying Homograft Rejection

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The immunological nature of homograft rejection has been established beyond any reasonable doubt. The role of cell mediated and humoral immunological mechanisms has been extensively investigated without a final answer as to which of these immunological responses should be primarily blamed for the homograft rejection. Whether the humoral antibodies are by themselves deleterious to the homograft or not, their detection still brings most valuable information.

Several serological procedures have been used in order to detect humoral antibodies accompanying homograft rejection, including hemagglutination [7, 8], cytotoxicity [9], leukoagglutination [1], and cutaneous anaphylaxis [12, 18]. In the present study, application of mixed agglutination and antiglobulin consumption tests has been explored.

Mixed agglutination should be defined as a procedure in which cells of morphologically distinguishable types are brought together by antibodies. The technique was originally described by TOPLEY *et al.* [16] and then it was used by a few other investigators [6, 11, 17].

COOMBS (for references see 4) should be given credit for elaborating this procedure to the point where it is a valuable, almost routine, laboratory tool. HÖGMAN [10] and FAGRÆUS and ESPMARK [5] applied the mixed agglutination technique to studies on tissue cultures. This method has also been used extensively in our laboratory. Reactions between cell cultures infected by measles virus and anti-viral antibodies were studied [2]. In testing cell cultures of various species origin against heteroimmune sera to organ suspensions, a new type of species-specific antigens which are thermolabile and saline non-extractible was described [13]. Recently, we found that this procedure may be successfully employed to study antibodies accompanying homograft rejection.

Briefly, the principle of the procedure employed was as follows. Monolayer cell cultures were incubated with transplantation sera. The binding of