SerialParaAtivarBartender100Sr4

> SerialParaAtivarBartender100Sr4 > SerialParaAtivarBartender100Sr2 Interleukin-2 treatment induces both IL-2 receptor heterodimer and IL-2 receptor p55/p70 heterotrimer expression in activated T cells from anuric rats. In the present study, we investigated whether the lack of bicarbonate-dependent reabsorption of tubular proteins, the well-known cause of chronic renal insufficiency in the X-linked hereditary renal tubular disorder of the anuric rat, could affect the expression of IL-2R on the surface of activated T cells. The expression and distribution of IL-2R on the surface of activated T cells was studied by immunofluorescence, and the expression and affinities of the receptors were analyzed by Scatchard plots. To activate T cells, anti-CD3 and anti-CD28 antibodies or phorbol 12-myristate 13-acetate (PMA) were used. Kinetics of IL-2R expression and occupancy were studied after PMA activation. The expression of IL-2R, as measured by fluorescence intensity, was reduced on activated T cells from anuric rats as compared to T cells from controls. In vitro activation of T cells with anti-CD3 and anti-CD28 antibodies increased IL-2R expression on T cells from both normal and anuric rats. The inhibitory effect of anuria on IL-2R expression was reversed by bicarbonate reabsorption in vitro. IL-2R expression increased in the presence of PMA, and this effect was detectable after as little as 5 min of activation with PMA. Bicarbonate did not alter this effect. The increase in IL-2R expression after PMA activation was accompanied by the appearance of IL-2 binding sites. A polyclonal murine anti-rat monoclonal anti-IL-2R antibody, used in Scatchard plots, detected both high- and low-affinity binding sites. Similarly, we detected 2-3 binding sites per cell with the same antibody on normal T cells stimulated with PMA. In contrast, a polyclonal rabbit anti-rat monoclonal anti-IL-2R antibody, used in Scatchard plots, detected

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