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The free Everest Ultimate Edition 5.50 (Barre) Serial.Variety of regulatory T cells in rheumatoid arthritis and their in vitro suppressive function on autoreactive T cells. To clarify whether the distinct population of CD4(+)CD25(+)FOXP3(+) T cells in rheumatoid arthritis (RA) is functionally distinct from the circulating CD4(+)CD25(-)FOXP3(-) cells or whether both cell populations are functionally similar. The in vitro suppressive function of CD4(+)CD25(+)FOXP3(+) T cells from RA synovial fluid or peripheral blood was compared with that of circulating CD4(+)CD25(-)FOXP3(-) T cells. Expression of CD152 (CTLA-4) and CD25 by RA CD4(+) T cells from synovial fluid or peripheral blood was analyzed by flow cytometry. Allergic and autoantigen-specific T cell proliferation was assessed after CD4(+)CD25(-) and CD4(+)CD25(+) T cells were stimulated with autologous dendritic cells in the presence of autoantigen. Suppression of T cell proliferation by CD4(+)CD25(+) T cells was determined by quantitating [3H]thymidine incorporation of the proliferating CD4(+)CD25(-) cells. These CD4(+)CD25(+)FOXP3(+) T cells inhibit effector T cell proliferation. In addition, some CD4(+)CD25(+)FOXP3(+) T cells express CTLA-4 and/or CD25 on the surface and exert suppressive function on T cell proliferation in the presence of autoantigen. The majority of CD4(+)CD25(+) T cells in RA was identified as CD4(+)CD25(+)FOXP3(+) cells, and they suppressed effector T cell proliferation in the presence of autoantigen. Their suppression of T cell proliferation was not diminished when the cells were stimulated with autoantigen in the presence of anti

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