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leukemias) and one T-cell leukemia line. IL-7Ralpha expression was not detectable by Western analysis in seven of the 22 cell lines and ranged from low to intermediate in 16 cell lines. Reverse transcriptase-polymerase chain reaction (RT-PCR) analysis detected IL-7Ralpha mRNA transcripts in all of the cell lines except for two types of Burkitt lymphomas (PDL and Daudi). The IL-7Ralpha mRNA transcripts were variably expressed, ranging from a few to many copies per cell. Southern analysis of IL-7Ralpha revealed one restriction fragment in six cell lines (ie, all lymphoma cell lines except for the two types of Burkitt lymphomas) and two restriction fragments in the T-cell leukemia line. IL-7Ralpha was not detectable in the two types of Burkitt lymphomas either by Western or Southern analysis. Cell surface IL-7Ralpha expression was examined by immunofluorescence flow cytometry in eight cell lines. While five cell lines expressed IL-7Ralpha on the cell surface, two cell lines expressed no detectable amounts of IL-7Ralpha by flow cytometry. Northern blot analysis indicated that the IL-7Ralpha mRNA