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cracked by superrw. A simple and reliable mechanism of nuclease action. The nuclease used for the removal of nonspecific phosphate groups from polynucleotides could be a single enzyme from *Escherichia coli* extract or a number of enzymes present in the extract, which has been purified to homogeneity. The activity of the latter is stabilized by reducing the concentration of EDTA to 1 mM and by addition of  $Mn^{2+}$ . Activity of the nuclease is linear with time up to 20 min and with the total concentration of DNA up to 20 micrograms/ml. At equilibrium, the enzyme splits  $10^{-10}$  mol of phosphate from a pyrimidine duplex per nanogram of protein. The mechanism of the hydrolysis of a thymidine dimer on the 5' side in 1 mM  $MnCl_2$  is proposed to proceed via the formation of a 3'-OH terminus at the dimer site followed by attack of the 5' end of the chain. At 2 mM  $MnCl_2$ , the nuclease splits the dimer at the site of the phosphorus atom;