EDTA-C2-dT is a deoxymyidine conjugated with the triethanoic (that is, triacetic) acid derivative of EDTA at the C-5 position of the thymine base. Oligonucleotides containing EDTA-C2-dT are typically used as artificial nucleases for cleaving both single and double-stranded DNA (1,2). In such cases, the modified oligo serves as a molecular recognition moiety, while the EDTA complexed to a metal ion (such as Fe(II)) acts as the actual cleavage agent. This analytical approach has been used to investigate the tertiary structure of tRNA (3).

Note that, because addition of Fe(II) and dithiothreotol (DTT) to form the Fe-EDTA complex in the oligo results in fairly rapid autocleavage of the oligo’s own phosphodiester backbone (thereby rendering the oligo useless as an artificial nuclease), substitution of methylphosphonate into this backbone can be done to confer resistance to such autocleavage (4).

Besides being used in the synthesis of oligo-based artificial nucleases, another possible use for EDTA-C2-dT is to introduce a metal complex (for example, a Cu-64-EDTA complex) into an oligonucleotide probe for use in experiments requiring in vivo imaging with positron emission tomography (PET).

References