



Product Specifications

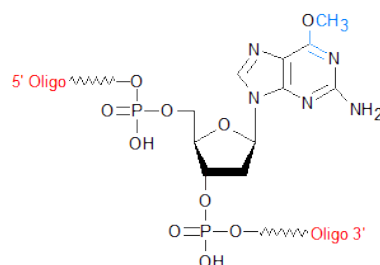
Custom Oligo Synthesis, antisense oligos, RNA oligos, chimeric oligos, Fluorescent dyes, Affinity Ligands, Spacers & Linkers, Duplex Stabilizers, Minor bases, labeled oligos, Molecular Beacons, siRNA, phosphonates Locked Nucleic Acids (LNA); 2'-5' linked Oligos

Oligo Modifications

For research use only. Not for use in diagnostic procedures for clinical purposes.

O6 Methyl dG

Category	Minor Bases
Modification Code	O6-Me-dG
Reference Catalog Number	26-6409
5 Prime	Y
3 Prime	Y
Internal	Y
Molecular Weight(mw)	343.24



O6 Methyl deoxyguanosine dG
[26-6409-XX]

O6-Methyl-deoxyguanosine (O6-Me-dG) is classified as an O-alkyl purine, and O6-Me-dG-modified oligonucleotides are primarily used in studies of the role of DNA alkylating agents in mutagenesis and carcinogenesis, and in studies into possible enzymatic mechanisms involved in repair of DNA alkylation damage. Both in vitro and in vivo, O6-Me-dG DNA lesions are formed by reaction with N-nitrosamines (known carcinogens), particularly those found in tobacco smoke (for example, 4-(methylnitrosamine)-1-(3-pyridyl)-1-butanone (NNK)) (1,2). O6-Me-dG is highly mutagenic due to its strongly mis-pairing characteristics (DNA polymerases preferentially insert thymine opposite O6-Me-dG, leading to G-to-A transitions) (3). Generation of O6-Me-dG lesions in the K-ras proto-oncogene or p53 gene results in activation of the proto-oncogene or inactivation of p53, leading to loss of control over cell growth and cancer (4). O6-Me-dG lesions are repaired by the specialized repair protein O6-alkylguanine DNA alkyltransferase (AGT), which removes the methyl group (as well as other alkyl groups) from O6 via a "suicide" inactivation mechanism, thereby restoring the natural guanine base (5). **References**

1. Hecht, S.S. Tobacco smoke carcinogens and lung cancer. *J. Natl. Cancer. Inst.* (1999), **91**: 1194-1210.
2. Hoffmann, D., Brunnemann, K.D., Prokopczyk, B., Djordjevic, M.V. Tobacco-specific N-nitrosamines and Areca-derived N-nitrosamines: chemistry, biochemistry, carcinogenicity, and relevance to humans. *J. Toxicol. Environ. Health* (1994), **41**: 1-52.
3. Warren, J.J., Forsberg, L.J., Beese, L.S. The structural basis for the mutagenicity of O6-methyl-guanine lesions. *Proc. Natl. Acad. Sci. USA* (2006), **103**: 19701-19706.
4. Zochbauer-Muller, S., Gazdar, A.F., Minna, J.D.. Molecular pathogenesis of lung cancer. *Annual Review of Physiology* (2002), **64**: 681-708.
5. Zang, H., Fang, Q., Pegg, A.E., Guengerich, F.P. Kinetic analysis of steps in the repair of damaged DNA by human O6-alkylguanine-DNA alkyltransferase. *J. Biol. Chem.* (2005), **280**: 30873-30881.