

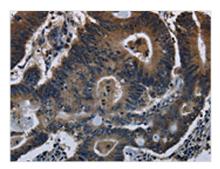
Image





DHCR7 Antibody

CSB-PA110763
Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Q9UBM7
Fusion protein of Human DHCR7
Rabbit
Human, Mouse
ELISA,IHC;ELISA:1:1000-1:2000,IHC:1:25-1:100
This gene encodes an enzyme that removes the C(7-8) double bond in the B ring of sterols and catalyzes the conversion of 7-dehydrocholesterol to cholesterol. This gene is ubiquitously expressed and its transmembrane protein localizes to the endoplasmic reticulum membrane and nuclear outer membrane. Mutations in this gene cause Smith-Lemli-Opitz syndrome (SLOS); a syndrome that is metabolically characterized by reduced serum cholesterol levels and elevated serum 7-dehydrocholesterol levels and phenotypically characterized by mental retardation, facial dysmorphism, syndactyly of second and third toes, and holoprosencephaly in severe cases to minimal physical abnormalities and near-normal intelligence in mild cases. Alternative splicing results in multiple transcript variants that encode the same protein.
Liquid
Non-conjugated
-20°C, pH7.4 PBS, 0.05% NaN3, 40% Glycerol
Antigen affinity purification
IgG
Homo sapiens (Human)
DHCR7



The image on the left is immunohistochemistry of paraffin-embedded Human colon cancer tissue using CSB-PA110763(DHCR7 Antibody) at dilution 1/20, on the right is treated with fusion protein. (Original magnification: ×200)