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ADA2 Antibody Blocking Peptide

Catalog Number: bs-9005P

Activity: Not tested

Purification: HPLC

Storage: Shipped at 4°C. Stored at -20°C for one year. Avoid repeated freeze/thaw cycles.

Background: CECR1 is a member of the adenosine and AMP deaminases family. It may act as a growth

factor and have adenosine deaminase activity. It is a candidate gene for cat eye syndrome.

Two transcript variants encoding distinct isoforms have been identified for this gene.

Adenosine deaminase is an enzyme that is present in most tissues and exists predominantly

as a monomer, although in some tissues it is associated with adenosine deaminase-binding $% \left(1\right) =\left(1\right) \left(1\right)$

protein. Adenosine deaminase degrades extracellular adenosine, which is toxic for

lymphocytes. A novel family of growth factors that share sequence similarity to adenosine deaminase has been identified. The cat eye syndrome critical region protein (CECR) family

includes CECR1, CECR2, CECR3, CECR4, CECR5, CECR6, CECR7, CECR8 and CECR9. The genes

encoding CECR proteins are candidates for Cat Eye Syndrome (CES), a developmental

disorder associated with the duplication of a 2 Mb region of 22q11.2. CES is characterized by

the combination of coloboma of the iris and anal atresia with fistula, downslanting

palpebral fissures, preauricular tags and/or pits, frequent occurrence of heart and renal

malformations, and normal or near-normal mental development. CECR family members are

widely expressed. Specifically, CECR1 has the highest expression in adult heart, lung,

lymphoblasts and placenta. CECR2 is also involved in neurulation and chromatin

remodeling. Mutations in the CECR2 gene result in neural tube defects.