

# **Screening Libraries**

**Proteins** 

# Inhibitors

# **Product** Data Sheet

# CTLA-4 Protein, Mouse (125a.a, HEK293, Fc)

Cat. No.: HY-P70706

Synonyms: Cytotoxic T-lymphocyte protein 4; Cytotoxic T-lymphocyte-associated antigen 4; CTLA-4; CD152;

Mouse Species: Source: **HEK293** 

Accession: P09793 (A37-D161)

Gene ID: 12477 Molecular Weight: 50-60 kDa

# **PROPERTIES**

	uence

AIQVTQPSVV LASSHGVASF PCEYSPSHNT DEVRVTVLRQ TNDQMTEVCA TTFTEKNTVG FLDYPFCSGT FNESRVNLTI QGLRAVDTGL YLCKVELMYP PPYFVGMGNG TQIYVIDPEP

CPDSD

**Appearance** Lyophilized powder.

**Formulation** Lyophilized from a 0.2 μm filtered solution of PBS, pH 7.4.

**Endotoxin Level** <1 EU/µg, determined by LAL method.

Reconsititution It is not recommended to reconstitute to a concentration less than 100 μg/mL in ddH<sub>2</sub>O. For long term storage it is

recommended to add a carrier protein (0.1% BSA, 5% HSA, 10% FBS or 5% Trehalose).

Storage & Stability Stored at -20°C for 2 years. After reconstitution, it is stable at 4°C for 1 week or -20°C for longer (with carrier protein). It is

recommended to freeze aliquots at -20°C or -80°C for extended storage.

Shipping Room temperature in continental US; may vary elsewhere.

## **DESCRIPTION**

## Background

CTLA-4, a pivotal inhibitory receptor, acts as a primary negative regulator in T-cell responses, exerting its influence within the intricate network of immune modulation. This regulatory function arises from the distinct property of CTLA-4, showcasing significantly heightened affinity for its natural B7 family ligands, CD80 and CD86, in comparison to the cognate stimulatory coreceptor CD28. The homodimeric structure of CTLA-4, intricately linked by disulfide bonds, underscores its role as a molecular sentinel in immune regulation. Functionally, CTLA-4 binds avidly to CD80/B7-1 and CD86/B7.2, competitively engaging with these ligands to suppress T-cell activation and finely tune immune responses. Additionally, CTLA-4 interacts with ICOSLG, contributing to its multifaceted engagement in immune checkpoint pathways.

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