# Anti-hPD-L1-hlgG1fut

Non-fucosylated monoclonal human IgG1 antibody against human PD-L1

Catalog code: hpdl1-mab13, hpdl1-mab13-03

https://www.invivogen.com/anti-human-pdl1-igg1fut-atezolizumab

## For research use only, not for diagnostic or therapeutic use

Version 23L18-MM

## **PRODUCT INFORMATION**

Contents: Anti-hPD-L1-hlgG1fut purified monoclonal antibody (mAb) is provided azide-free and lyophilized. It is available in two quantities: hpdl1-mab13: 100 µg Anti-hPD-L1-hlgG1fut

hpdl1-mab13-03: 3 x 100 µg Anti-hPD-L1-hlgG1fut

Target: Human Programmed death-ligand 1

Species reactivity: Human and mouse

Source: CHO cells

Isotype: Human IgG1fut Light chain type: Kappa

**Clonality:** Monoclonal

**Purification:** By affinity chromatography with protein G

**Formulation:** 0.2 µm filtered solution in a sodium phosphate buffer with glycine, saccharose, and stabilizing agents

#### Storage and stability

- Product is shipped at room temperature. Upon receipt, store at -20 °C.

- Reconstituted antibody is stable for 1 month when stored at 4 °C and for 1 year when aliquoted and stored at -20 °C. Avoid repeated freeze-thaw cycles.

#### Quality control

- Binding of Anti-hPD-L1-hlgG1fut to surface expressed human PD-L1 on target cells has been validated using flow cytometry.

- ADCC has been tested using CD16-expressing Jurkat-NFAT reporter cells.

- Absence of bacterial contamination (e.g. lipoproteins and endotoxins) has been confirmed using HEK-Blue™ TLR2 and TLR4 cellular assays.

### PRODUCT DESCRIPTION

Anti-hPD-L1-hlgG1fut is a recombinant monoclonal antibody (mAb) featuring a fully sequenced variable region equivalent to Atezolizumab, that recognizes human (h)PD-L1, and a non-fucosylated constant region of the human lgG1 isotype (hlgG1fut). Anti-hPD-L1-hlgG1fut was generated by recombinant DNA technology, produced in CHO cells, and purified by affinity chromatography with protein G.

#### PD-L1 background

Programmed cell death ligand 1 (PD-L1), also known B7 homolog 1 (B7-H1) is a transmembrane protein that is constitutively expressed or induced in myeloid, lymphoid, and epithelial cells, as well as in cancer<sup>1,2</sup>. PD-L1 is the principle ligand for programmed cell death protein 1 (PD-1). This interaction is essential in the development of immune tolerance preventing excessive immune cell activity. However, PD-L1 expression is an immune evasion mechanism exploited by various malignancies<sup>3</sup>. Specifically, over-expressed PD-L1 on tumor cells and tumor infiltrating immune cells, such as macrophages, is able to bind to PD-1 on cytotoxic T cells, and ultimately inhibit the anti-tumor T cell response<sup>2</sup>. 4.

TECHNICAL SUPPORT InvivoGen USA (Toll-Free): 888-457-5873 InvivoGen USA (International): +1 (858) 457-5873 InvivoGen Europe: +33 (0) 5-62-71-69-39 InvivoGen Asia: +852 3622-34-80 E-mail: info@invivogen.com Thus, there are numerous PD-L1 inhibitors in development as promising immuno-oncology therapies. Notably, Atezolizumab (also known as MPDL3280A), a fully humanized IgG1 (N298A) mAb that blocks the interaction of PD-L1 with PD-1 and induces anti-tumor immune reactivation, has been approved by the FDA for combinational use in the treatment of lung and breast cancer<sup>2, 5</sup>.

#### IgG1fut isotype effector function

Anti-hPD-L1-hlgG1fut is a non-fucosylated antibody. The absence of the fucose residue from the N-glycans of the IgG-Fc results in an enhancement of antibody-dependent cellular cytotoxicity (ADCC) without any detectable change to complement-dependent cytotoxicity (CDC) or antigen binding capability.

1. Juneja V.R. et al. 2017. PD-L1 on tumor cells is sufficient for immune evasion in immunogenic tumors and inhibits CD8 T cell cytotoxicity. J Exp Med 214, 895-904. 2. Kythreotou A. et al. 2018. PD-L1. J Clin Pathol 71, 189-194. 3. Sun C. et al. 2018. Regulation and Function of the PD-L1 Checkpoint. Immunity 48, 434-452. 4. Lau J. et al. 2017. Tumour and host cell PD-L1 is required to mediate suppression of anti-tumour immunity in mice. Nat Commun 8, 14572. 5. Heimes A.S. & Schmidt M. 2019. Atezolizumab for the treatment of triple-negative breast cancer. Expert Opin Investig Drugs 28, 1-5.

# **METHODS**

#### Anti-hPD-L1-hlgG1fut resuspension (100 µg/ml)

Note: Ensure you see the lyophilized pellet before resuspension.

- Add 1 ml of sterile water to 100  $\mu g$  and gently pipette until completely resuspended.

- Prepare aliquots and store at -20 °C until required.

# ANTIBODY ISOTYPE COLLECTION

For your research, InvivoGen provides an Anti-hPD-L1 isotype family. This isotype family consists of mAbs comprising a variable region equivalent to Atezolizumab and differing constant regions of both natural and engineered human isotypes (*see related products*). The isotypes differ in their functional and effector functions, such as antibody-dependent cell-mediated cytotoxicity (ADCC) and complement dependent cytotoxicity (CDC).

<u>Disclaimer</u>: The terms "Atezolizumab" and "MPDL3280A" are only used as references. Anti-hPD-L1-hlgG1fut is not a pharmaceutical biosimilar of Atezolizumab. It has not been developed nor approved by Atezolizumab owner(s), and is not intended for any therapeutic or diagnostic use in human or animal.

# RELATED PRODUCTS

Product	Catalog Code
Anti-hPD-L1-hlgG1	hpdl1-mab1
Anti-hPD-L1-hlgG1 (N298A)	hpdl1-mab12

