

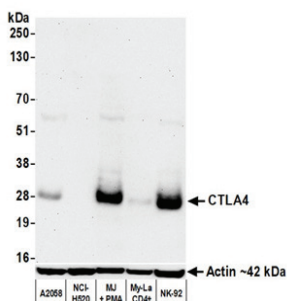
# CTLA4

**Product:** CTLA4 (A700-257)  
**Reactivity:** Human  
**Validated Applications:** IHC, WB  
**Full Name:** Cytotoxic T-lymphocyte protein 4  
**Gene ID:** 1493  
**Uniprot ID:** P16410  
**Alternative Names:** CTLA-4, CD152, GSE

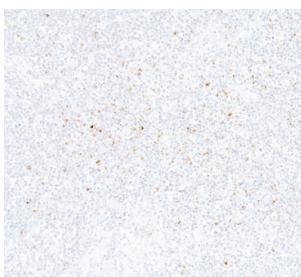
## Background Information

CTLA4 plays a critical role in regulating peripheral immune responses and has become a target for cancer immunotherapy. CTLA4 is expressed on activated T cells and regulatory T cells, where it functions primarily to inhibit T cell responses through interactions with CD80 and CD86<sup>1,2</sup>. CTLA4 is part of the T cell checkpoint pathways that dampen inappropriate or sustained immune activation. Tumors and pathogens can exploit this pathway as a way to evade the immune system. In 2011, CTLA4 became the first immune checkpoint receptor to have a targeted immunotherapy approved by the FDA. CTLA4 blockade immunotherapy has improved the survival rates of patients with melanoma, renal cell carcinoma, head and neck squamous cell cancer, and non-small cell lung cancer.<sup>2-4</sup>

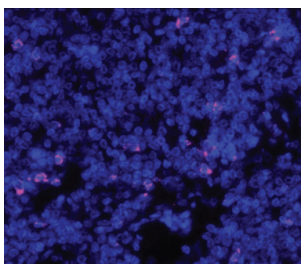
## Featured Applications



**Detection of human CTLA4 by western blot. Antibody:** A700-257 used at 1:1000.



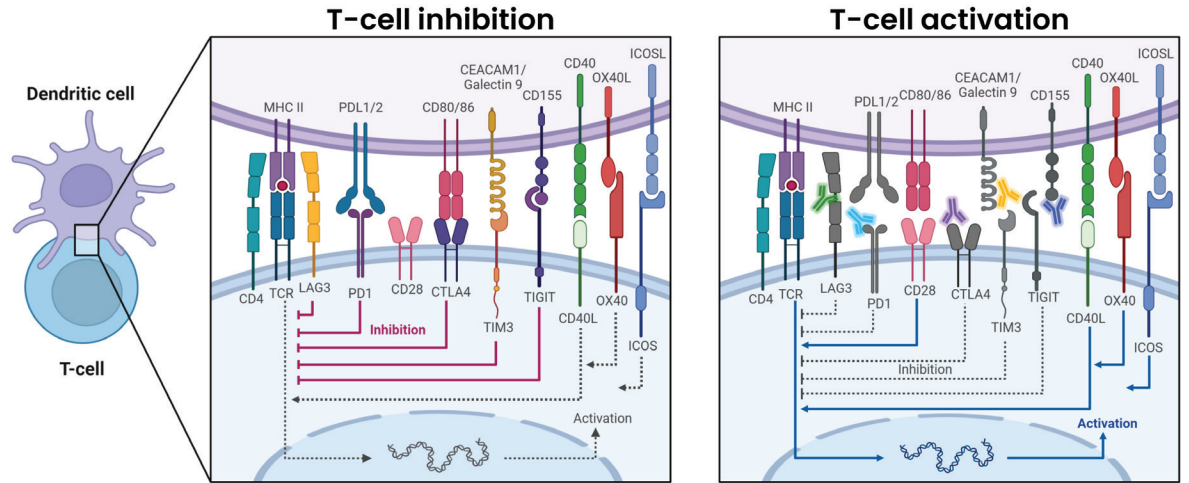
**Detection of human CTLA4 by immunohistochemistry.** Sample: FFPE section of human tonsil. Antibody: A700-257.



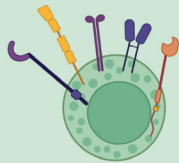
**Detection of human CTLA4 (magenta) by immunofluorescence.** Sample: FFPE section of metastatic lymph node. Antibody: A700-257 used at 1:20. Counterstain: DAPI (blue).

**References:**

1. Bateman A, Martin MJ, Orchard S, et al. UniProt: the Universal Protein Knowledgebase in 2023. *Nucleic Acids Res.* 2023;51(D1):D523-D531. doi:10.1093/nar/gkac1052
2. Masteller EL, Chuang E, Mullen AC, Reiner SL, Thompson CB. Structural Analysis of CTLA-4 Function in Vivo. *The Journal of Immunology.* 2000;164(10):5319-5327. doi:10.4049/jimmunol.164.10.5319
3. Atkins MB, Clark JI, Quinn DI. Immune checkpoint inhibitors in advanced renal cell carcinoma: experience to date and future directions. *Annals of Oncology.* 2017;28(7):1484-1494. doi:10.1093/annonc/mdx151
4. Sobhani N, Tardiel-Cyril DR, Davtyan A, Generali D, Roudi R, Li Y. CTLA-4 in Regulatory T Cells for Cancer Immunotherapy. *Cancers (Basel).* 2021;13(6):1440. doi:10.3390/cancers13061440

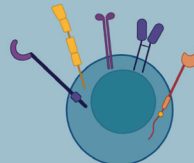


**Cytotoxic T Cells**



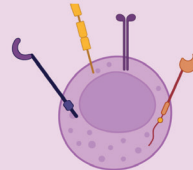
Engagement of immune checkpoints like CTLA4, LAG3, PD1, TIM3, and TIGIT on cytotoxic T cells can result in decreased cell killing, cytotoxic T cell exhaustion, and a pro-tumorigenic environment.

**Regulatory T Cells**



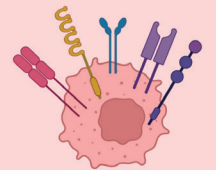
Engagement of immune checkpoints like CTLA4, LAG3, PD1, TIM3, and TIGIT on regulatory T cells can lead to increased suppressive activity, blocked APC maturation, dampened inflammation, and a pro-tumorigenic environment.

**Natural Killer Cells**



Engagement of immune checkpoints like LAG3, PD1, TIM3, and TIGIT on natural killer cells can lead to decreased NK cell activation, cell killing, and a pro-tumorigenic environment.

**Cancer Cells**



Engagement of immune checkpoints like CTLA4, LAG3, PD1, TIM3, and TIGIT through ligand expression on cancer cells can lead to decreased cell killing and increased immune suppression culminating in a pro-tumorigenic environment.