



# Go-ChIP-Grade™ Purified anti-NFATc1 Antibody

**Catalog# / Size** 649607 / 25 μg

649608 / 100 µg

Clone 7A6

Regulatory Status RUO

Other Names Nuclear factor of activated T-cells, cytoplasmic 1, NFAT transcription complex cytosolic

component, NFAT2, NFATc

**Isotype** Mouse IgG1, κ

**Description** The product of this gene is a component of the nuclear factor of activated T cells DNA-binding

transcription complex. The protein complex consists of NFAT1, NFAT2 (NFATc1 or NFATc), NFAT3, and NFAT4. All members of this family are transcription factors with a Rel homology domain and regulate gene transcription in concert with AP-1 (Jun/Fos) to orchestrate an effective immune response. NFAT proteins are predominantly expressed in cells of the immune system but are also expressed in skeletal muscle, keratinocytes and adipocytes, regulating cell

differentiation programs in these cells. In resting cells, NFAT proteins are heavily phosphorylated and localized in the cytoplasm. Increased intracellular calcium concentrations

activate the calcium/calmodulin-dependent serine phosphatase calcineurin, which

dephosphorylates NFAT proteins, resulting in their subsequent translocation to the nucleus. Proteins belonging to this family of transcription factors play a central role in inducible gene transcription during immune response. The product of this gene is an inducible nuclear component. It functions as a major molecular target for the immunosuppressive drugs such as cyclosporin A. Five transcript variants encoding distinct isoforms have been identified for this gene. Different isoforms of this protein may regulate inducible expression of different cytokine

aenes.

## **Product Details**

Verified Reactivity Human, Mouse, Rat

Antibody Type Monoclonal

Host Species Mouse

Immunogen Recombinant protein of human NFATc1 amino acids 197-304.

**Formulation** Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.

**Preparation** The antibody was purified by affinity chromatography.

Concentration 0.5 mg/ml

Storage & Handling The antibody solution should be stored undiluted between 2°C and 8°C.

Application <u>ChIP - Quality tested</u>

WB - Verified

ICC, ICFC - Reported in the literature, not verified in house

Recommended Usage Each lot of this antibody is quality control tested by ChIP Assay. The suggested dilution for ChIP

application is 1:100-1:2000 by volume. For Western blotting, the suggested use of this reagent is 1.0 µg per ml. It is recommended that the reagent to be titrated for optimal performance before

each experiment.

**Application Notes** 25 μg, 100 μg of Go-ChIP-Grade™ Purified Antibody can be used for 4-50, 16-200

immunoprecipitations, respectively, at the recommended dilution.

7A6 antibody detects endogenous human NFATC1 in Western blot. There are 10 isoforms with a predicted MW of 39KD, 74kD, 76kD, 77kD (3), 88kD (2), 100kD, 101kD. With this antibody, observed MW bands range from 70 - 120 kD. The optimal concentration should be determined by

titration for each individual assay of interest.

Application References

1. Timmerman LA, et al. 1997. J. Immunol. 159:2735. (IF)

(PubMed link indicates BioLegend citation)

2. Brandt C, et al. 2010. Cytometry A. 77:607. (FC)

3. Fan W, et al. 2012. Arthritis Rheum. 64:3715. PubMed

**Product Citations** 

1. Zhong Y, et al. 2022. Nat Immunol. 23:122. PubMed

**RRID** AB\_2721595 (BioLegend Cat. No. 649607)

AB 2721596 (BioLegend Cat. No. 649608)

## **Antigen Details**

Structure 943 amino acids with predicted molecular weight of 101kD. There are 10 isoforms with a

predicted MW of 39kD, 74kD, 76kD, 77kD (3), 88kD (2), 100kD, 101kD.

Distribution Cytoplasm, Nucleus. NFATc1 is cytoplasmic in the phosphorylated form and nuclear after

calcineurin-mediated dephosphorylation controlled activation.

Biology Area Cell Biology, Immunology, Neuroscience, Neuroscience Cell Markers, Signal Transduction,

Transcription Factors

Molecular Family Nuclear Markers

Antigen References 1. Zhao Q, et al. 2010. Int. J. Biochem. Cell Biol. 42:576.

2. Hoey T, et al. 1995. Immunity 2:461.

Northrop JP, et al. 1993. J. Biol. Chem. 268:2917.
 Hogan PG, et al. 2003. Genes Dev. 17:2205.

5. Shaw KT, *et al.* 1995. *P. Natl. Acad. Sci. USA* 92:11205.

6. Serfling E, et al. 2007. Sci. STKE 138:pe42.

Gene ID <u>4772</u>

#### **Related Protocols**

BioLegend's Tools for Chromatin Immunoprecipitation (ChIP) Assays - Video

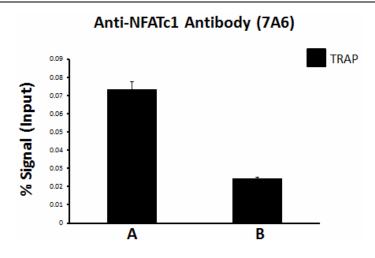
Chromatin Immunoprecipitation (ChIP) Assay Protocol

Western Blotting Protocol

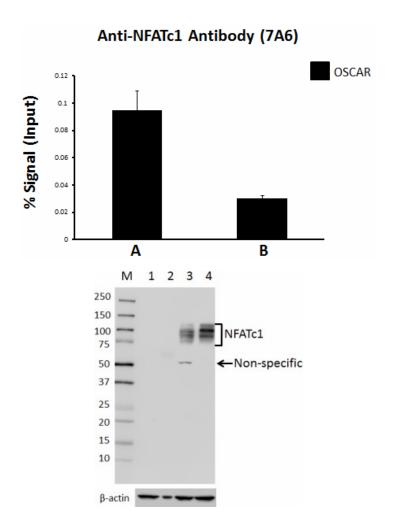
## **Other Formats**

PE anti-NFATc1, Purified anti-NFATc1, Alexa Fluor® 488 anti-NFATc1, Go-ChIP-Grade™ Purified anti-NFATc1

### **Product Data**



Chromatin Immunoprecipitation (ChIP) was performed using Go-ChIP-Grade make the contract of the co Protein G Enzymatic kit by loading 3 µg of cross-linked chromatin samples from Raw264.7 cells treated with RANKL with either A) 1:500 dilution of Go-ChIP-Grade™ Purified anti-NFATc1 (Clone 7A6), or B) equal amount of Purified Mouse IgG1 Isotype Control Antibody. The enriched DNA was purified and quantified by real-time qPCR using primers targeting mouse TRAP gene region. The amount of immunoprecipitated DNA in each sample is represented as signal relative to the 5% of total amount of input chromatin.



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Total lysates (15 µg protein) from EL4 (lane 1), HeLa (lane 2), Jurkat (lane 3) and Raw264.7 (lane 4) cells were resolved by electrophoresis (4-20% Tris-Glycine gel), transferred to nitrocellulose, and probed with 1:500 diluted (1 µg/mL) Purified anti-NFATc1 Antibody, clone 7A6 (upper). Proteins were visualized by chemiluminescence detection using a 1:3000 diluted goat anti-mouse-IgG secondary antibody conjugated to HRP for the anti-NFATc1 Antibody. 7A6 antibody detects different endogenous human NFATc1 spicing variants with an observed molecular weight from 70 - 120 kD. 1:5000 diluted Direct-Blot HRP anti-β-Actin Antibody (2F1-1) was used as a loading control (lower). Lane M: Molecular weight ladder.

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