

Product datasheet for TA500304M

OriGene Technologies, Inc.

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Neurogenin 1 (NEUROG1) Mouse Monoclonal Antibody [Clone ID: OTI5H10]

Product data:

Product Type: Primary Antibodies

Clone Name: OTI5H10

Applications: IF, IHC, IP, WB

Recommended Dilution: WB 1:2000, IHC 1:50, IF 1:100; IP 2-4ug/mg

Reactivity: Human, Mouse, Rat

Host: Mouse Isotype: IgG1

Clonality: Monoclonal

Immunogen: Full-length protein expressed in 293T cell transfected with human NEUROG1 expression

vector

Formulation: PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide

Concentration: 0.65 mg/ml

Purification: Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography

(protein A/G)

Conjugation: Unconjugated

Storage: Store at -20°C as received.

Stability: Stable for 12 months from date of receipt.

Predicted Protein Size: 25.7 kDa

Gene Name: neurogenin 1

Database Link: NP 006152

Entrez Gene 18014 MouseEntrez Gene 29410 RatEntrez Gene 4762 Human

O92886

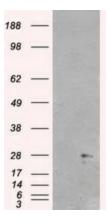
Synonyms: AKA; bHLHa6; Math4C; NEUROD3; ngn1

Protein Families: Druggable Genome, ES Cell Differentiation/IPS, Transcription Factors

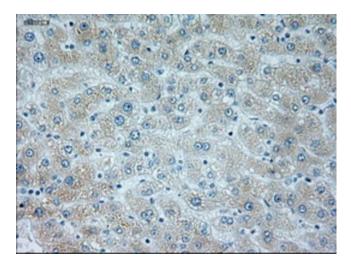




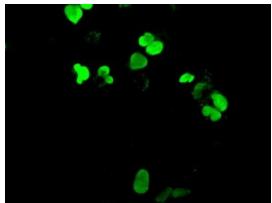
Product images:



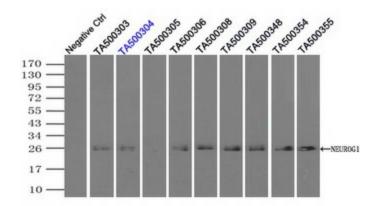
HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY NEUROG1 ([RC207029], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-NEUROG1. Positive lysates [LY401855] (100ug) and [LC401855] (20ug) can be purchased separately from OriGene.



Immunohistochemical staining of paraffinembedded liver tissue within the normal limits using anti-NEUROG1mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.



Anti-NEUROG1 mouse monoclonal antibody ([TA500304]) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY NEUROG1 ([RC207029]).



Immunoprecipitation of NEUROG1 by using TrueMab monoclonal anti-NEUROG1 antibody (Negative control: IP without adding anti-NEUROG1 antibody). For each experiment, 500ul of DDK tagged NEUROG1 overexpression lysates (at 1:5 dilution with HEK293T lysate), 2ug of anti-NEUROG1 antibody and 20ul (0.1mg) of goat antimouse conjugated magnetic beads were mixed and incubated overnight. After extensive wash to remove any non-specific binding, the immuno-precipitated products were analyzed with rabbit anti-DDK polyclonal antibody.