

Technical Data Sheet

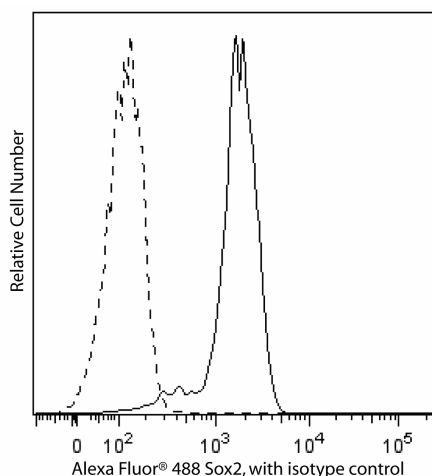
Alexa Fluor® 488 Mouse anti-Sox2

Product Information

Material Number:	560301
Size:	50 tests
Vol. per Test:	5 µl
Clone:	245610
Immunogen:	Human SOX2 Recombinant Protein
Isotype:	Mouse IgG2a
Reactivity:	QC Testing: Human Reported: Mouse
Storage Buffer:	Aqueous buffered solution containing BSA and ≤0.09% sodium azide.

Description

The monoclonal antibody 245610 recognizes the Sox2 transcription factor. Sox2 [SRY (sex determining region Y)-box 2] is a member of the SRY-related HMG-box (SOX) family of transcription factors. Sox2 is required for the maintenance of the undifferentiated state of pluripotent stem cells. Complexes of Sox2 with the homeobox transcription factors Oct3/4 and/or Nanog bind to the promoters of a network of genes that are involved in the maintenance of pluripotency and self renewal in stem cells. The 245610 antibody recognizes both human and mouse proteins. Sox2 is also a marker of neural stem cells during embryonic development and in the adult brain.



Flow cytometric analysis of Alexa Fluor® 488 anti-Sox2 in H9 cells. H9 human embryonic stem (ES) cells (WiCell, Madison, WI) were fixed (BD Cytotix™ buffer, Cat. No. 554655) for 10 minutes at 37°C, permeabilized with BD™ Phosflow Perm Buffer III (Cat. No. 558050) on ice for 30 minutes, and then stained with either Alexa Fluor® 488 Mouse anti-Sox2 (solid line) or Alexa Fluor® 488 Mouse IgG2a Isotype Control (Cat. No. 558055, dashed line). Flow cytometry was performed on a BD FACSCanto™ II flow cytometry system.

Preparation and Storage

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

The antibody was conjugated to Alexa Fluor® 488 under optimum conditions, and unreacted Alexa Fluor® 488 was removed.

Store undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

Application Notes

Application

Intracellular staining (flow cytometry)	Routinely Tested
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Recommended Assay Procedure:

Any of the three BD™ Phosflow permeabilization buffers may be used with this antibody: BD™ Phosflow Perm/Wash Buffer I (Cat. No. 557885), BD™ Phosflow Perm Buffer II (Cat. No. 558052), or BD™ Phosflow Perm Buffer III (Cat. No. 558050).

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Suggested Companion Products

<u>Catalog Number</u>	<u>Name</u>	<u>Size</u>	<u>Clone</u>
554655	Fixation Buffer	100 ml	(none)
558050	Perm Buffer III	125 ml	(none)
558055	Alexa Fluor®488 Mouse IgG2a, κ Isotype control	50 tests	MOPC-173

Product Notices

1. This reagent has been pre-diluted for use at the recommended Volume per Test. We typically use 1×10^6 cells in a 100- μ l experimental sample (a test).
2. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
3. The Alexa Fluor®, Pacific Blue™, and Cascade Blue® dye antibody conjugates in this product are sold under license from Molecular Probes, Inc. for research use only, excluding use in combination with microarrays, or as analyte specific reagents. The Alexa Fluor® dyes (except for Alexa Fluor® 430), Pacific Blue™ dye, and Cascade Blue® dye are covered by pending and issued patents.
4. Alexa Fluor® 488 fluorochrome emission is collected at the same instrument settings as for fluorescein isothiocyanate (FITC).
5. For fluorochrome spectra and suitable instrument settings, please refer to our Fluorochrome Web Page at www.bdbiosciences.com/colors.
6. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
7. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
8. Alexa Fluor® is a registered trademark of Molecular Probes, Inc., Eugene, OR.

References

- Boyer LA, Lee TI, Cole MF, et al. Core transcriptional regulatory circuitry in human embryonic stem cells. *Cell*. 2005; 122:947-956. (Biology)
- Pan G, Thomson JA. Nanog and transcriptional networks in embryonic stem cell pluripotency. *Cell Res*. 2007; 17:42-49. (Biology)
- Takahashi K, Yamanaka S. Induction of pluripotent stem cells from mouse embryonic and adult fibroblast cultures by defined factors. *Cell*. 2006; 126:633-676. (Biology)