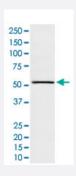


# NR0B1 monoclonal antibody, clone AECC-14

Catalog # MAB22101 Size 100 uL

## **Applications**



#### Western Blot (Cell lysate)

Western Blot (cell lysate) analysis of A-431 cell lysate.

Specification	
Product Description	Rabbit monoclonal antibody raised against synthetic protein of human NR0B1.
Immunogen	A synthetic peptide corresponding to human NR0B1.
Host	Rabbit
Reactivity	Human
Specificity	This anyibody reacts with human NR0B1, in native form and recombinant. Superfamily members of N R0B1 are not reactive to antibody.
Form	Liquid
Purification	Affinity purification
Isotype	lgG
Recommend Usage	Flow Cytometry (1:50) Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) (1:100-500) Immunofluorescence (1:50-200) Immunocytochemistry (1:50-200) Western Blot (1:500-1000) The optimal working dilution should be determined by the end user.



#### **Product Information**

Storage Buffer	In PBS, 150 mM NaCl, pH 7.4 (50% glycerol, 0.02% sodium azide).
Storage Instruction	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

### **Applications**

Western Blot (Cell lysate)

Western Blot (cell lysate) analysis of A-431 cell lysate.

- Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)
- Immunocytochemistry
- Immunofluorescence
- Flow Cytometry

Gene Info — NR0B1	
Entrez GenelD	<u>190</u>
Protein Accession#	<u>P51843</u>
Gene Name	NR0B1
Gene Alias	AHC, AHCH, AHX, DAX-1, DAX1, DSS, GTD, HHG, NROB1
Gene Description	nuclear receptor subfamily 0, group B, member 1
Omim ID	300018 300200 300473
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes a protein that contains a DNA-binding domain. The encoded protein acts as a dominant-negative regulator of transcription which is mediated by the retinoic acid receptor. This protein also functions as an anti-testis gene by acting antagonistically to Sry. Mutations in this gene result in both X-linked congenital adrenal hypoplasia and hypogonadotropic hypogonadism. [provided by RefSeq
Other Designations	OTTHUMP00000023102 gonadotropin deficiency nuclear hormone receptor