

# AP3B1 Polyklonaler Antikörper

Katalog-Nr.: 13384-1-AP

Vorgestelltes Produkt

14 Publikationen

## Allgemeine Informationen

Katalog-Nr.:	GenBank-Zugangsnummer:	Reinigungsmethode:
13384-1-AP	BC038444	Antigen-Affinitätsreinigung
<b>Größe:</b>	<b>GenID (NCBI):</b>	<b>Empfohlene Verdünnungen:</b>
150ul, Konzentration: 600 µg/ml von Nanodrop;	8546	WB 1:500-1:3000 IP 0.5-4.0 ug für IP und 1:500-1:1000 für WB IHC 1:50-1:500
<b>Wirz:</b> Kaninchen	<b>Vollständiger Name:</b> adaptor-related protein complex 3, beta 1 subunit	
<b>Isotyp:</b> IgG	<b>Berechneté Masse:</b> 1094 aa, 121 kDa	
<b>Immunogen Katalognummer:</b> AG4225	<b>Beobachteté Masse:</b> 140 kDa	

## Anwendungen

<b>Geprüfte Anwendungen:</b> IHC, IP, WB, ELISA	<b>Positivkontrollen:</b> WB: A431-Zellen, COLO 320-Zellen, HeLa-Zellen, HepG2-Zellen, Maus-Thymusgewebe, SKOV-3-Zellen
<b>In Publikationen genannte Anwendungen:</b> IF, WB	<b>IP :</b> COLO 320-Zellen,
<b>Getestete Reaktivität:</b> Human, Maus, Ratte	<b>IHC :</b> Rattenhirngewebe,
<b>Zitierte Arten:</b> Human, Maus	
<b>Hinweis-IHC:</b> Antigendemaskierung mit TE-Puffer pH 9,0 empfohlen. (*) Wahlweise kann die Antigendemaskierung auch mit Citratpuffer pH 6,0 erfolgen.	

## Hintergrundinformationen

AP3B1 is the 140-kDa β3A subunit of the adaptor-related protein complex-3 (AP-3), a ubiquitous heterotetrameric complex that is localized to the trans-Golgi network and endosomes and is involved in protein trafficking to lysosomes or specialized endosomal-lysosomal organelles (PMID: 9182526; 9545220). This complex is composed of two larger subunits (δ and β3A or β3B), a medium subunit (μ3A or μ3B), and a small subunit (ο3A or ο3B). The absence of the β3A subunit (AP3B1) results in the loss of stability of AP3 and leads to degradation of μ3A, to which β3A is directly bound, while the other subunits are variably affected (PMID: 16507770). AP3B1 contains three main domains: the N-terminal head domain, the hinge, and the C-terminal ear domain. It has been reported as a target of IP(7)-mediated pyrophosphorylation (PMID: 19934039). Defects in AP3B1 are the cause of Hermansky-Pudlak syndrome type 2 (HPS2) (PMID: 10024875; 16507770).

## Bemerkenswerte Veröffentlichungen

Verfasser	Pubmed ID	Journal	Anwendung
Weina Sun	25210190	J Virol	WB, IF
Joshi Stephen	28296950	PLoS One	WB
Maria B Bagh	28266544	Nat Commun	WB

## Lagerung

**Lagerungsbedingungen:**  
Bei -20°C lagern. Nach dem Versand ein Jahr lang stabil  
**Lagerungspuffer:**  
PBS mit 0.02% Natriumazid und 50% Glycerin pH 7.3.  
**Aliquotieren ist nicht notwendig bei -20°C Lagerung**

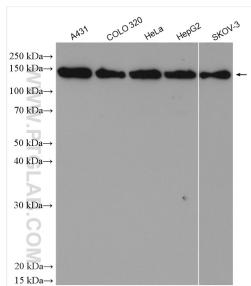
\*\*\* 20ul-Größen enthalten 0.1% BSA

For technical support and original validation data for this product please contact:  
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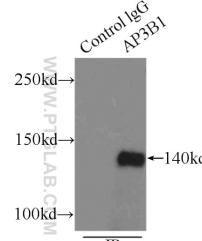
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W: [ptglab.com](http://ptglab.com)

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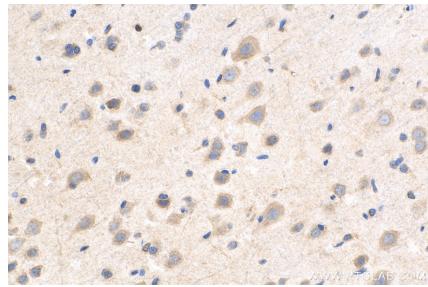
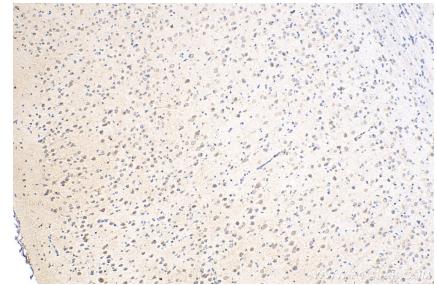
## Ausgewählte Validierungsdaten



Various lysates were subjected to SDS PAGE followed by western blot with 13384-1-AP (AP3B1 antibody) at dilution of 1:1500 incubated at room temperature for 1.5 hours.



IP Result of anti-AP3B1 (IP:13384-1-AP, 3ug; Detection:13384-1-AP 1:500) with COLO 320 cells lysate 2500ug.



Immunohistochemical analysis of paraffin-embedded rat brain tissue slide using 13384-1-AP (AP3B1 antibody) at dilution of 1:200 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).