

Product datasheet

Anti-B3GAT3 antibody ab68026

1 图像

概述

|       |   |
|-------|---|
| 产品名称  | Anti-B3GAT3抗体   |
| 描述    | 小鼠多克隆抗体to B3GAT3  |
| 宿主    | Mouse   |
| 经测试应用 | 适用于: WB   |
| 种属反应性 | 与反应: Human  |
| 免疫原   | Full length protein, corresponding to amino acids 1-335 of Human B3GAT3 |
| 阳性对照  | Transfected 293T cell line  |

性能

|      |   |
|------|---|
| 形式   | Liquid  |
| 存放说明 | Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw cycles. |
| 存储溶液 | pH: 7.4<br>Constituent: PBS   |
| 纯度   | Protein A purified  |
| 克隆   | 多克隆   |
| 同种型  | IgG   |

应用

Our [Abpromise guarantee](#) covers the use of **ab68026** in the following tested applications.

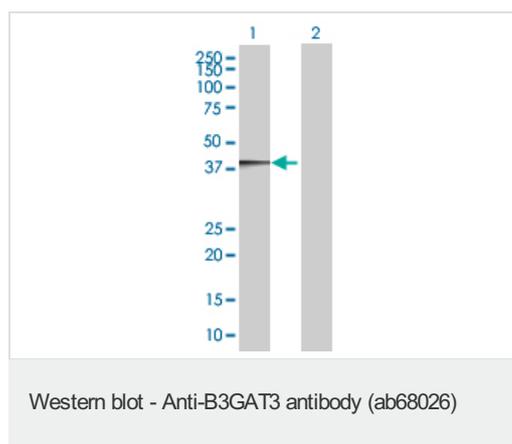
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

| 应用 | Ab评论 | 说明   |
|----|------|--|
| WB |      | 1/500 - 1/1000. Detects a band of approximately 37 kDa (predicted molecular weight: 37 kDa). |

靶标

|              |  |
|--------------|--|
| <b>功能</b>    | Glycosaminoglycans biosynthesis. Involved in forming the linkage tetrasaccharide present in heparan sulfate and chondroitin sulfate. Transfers a glucuronic acid moiety from the uridine diphosphate-glucuronic acid (UDP-GlcUA) to the common linkage region trisaccharide Gal-beta-1,3-Gal-beta-1,4-Xyl covalently bound to a Ser residue at the glycosaminoglycan attachment site of proteoglycans. Can also play a role in the biosynthesis of I2/HNK-1 carbohydrate epitope on glycoproteins. Shows strict specificity for Gal-beta-1,3-Gal-beta-1,4-Xyl, exhibiting negligible incorporation into other galactoside substrates including Galbeta1-3Gal beta1-O-benzyl, Galbeta1-4GlcNAc and Galbeta1-4Glc. |
| <b>组织特异性</b> | Ubiquitous (but weakly expressed in all tissues examined).   |
| <b>通路</b>    | Protein modification; protein glycosylation.   |
| <b>疾病相关</b>  | Defects in B3GAT3 are the cause of multiple joint dislocations short stature craniofacial dysmorphism and congenital heart defects (JDSSDHD) [MIM:245600]. An autosomal recessive disease characterized by dysmorphic facies, bilateral dislocations of the elbows, hips, and knees, clubfeet, and short stature, as well as cardiovascular defects.   |
| <b>序列相似性</b> | Belongs to the glycosyltransferase 43 family.  |
| <b>翻译后修饰</b> | N-glycosylated.  |
| <b>细胞定位</b>  | Golgi apparatus membrane. Golgi apparatus > cis-Golgi network.   |

## 图片



**All lanes :** Anti-B3GAT3 antibody (ab68026)  
at 1/500 dilution

**Lane 1 :** B3GAT3 transfected 293T cell lysate

**Lane 2 :** Non transfected 293T cell lysate

Lysates/proteins at 25 µg per lane.

### Secondary

**All lanes :** Goat Anti-Mouse IgG (H&L)-HRP

Conjugate secondary  
antibody at 1/2500 dilution

Developed using the ECL technique.

**Predicted band size:** 37 kDa

**Observed band size:** 37 kDa

**Please note:** All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE"

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- Response to your inquiry within 24 hours
  
- We provide support in Chinese, English, French, German, Japanese and Spanish
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