

# Connexin 31 Antibody (C-term)

Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP1553c

#### **Product Information**

**Application** WB, E **Primary Accession** 075712 Reactivity Human Host Rabbit Clonality Polyclonal Isotype Rabbit IgG **Clone Names** RB0443 **Calculated MW** 30818 **Antigen Region** 204-233

### **Additional Information**

**Gene ID** 2707

Other Names Gap junction beta-3 protein, Connexin-31, Cx31, GJB3, CX31

**Target/Specificity** This Connexin 31 Antibody antibody is generated from rabbits immunized

with a KLH conjugated synthetic peptide between 204~233 amino acids from

the C-terminal region of human GJB3.

**Dilution** WB~~1:1000 E~~Use at an assay dependent concentration.

**Format** Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide.

This antibody is prepared by Saturated Ammonium Sulfate (SAS) precipitation

followed by dialysis against PBS.

**Storage** Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions** Connexin 31 Antibody (C-term) is for research use only and not for use in

diagnostic or therapeutic procedures.

## **Protein Information**

Name GJB3

Synonyms CX31

**Function** One gap junction consists of a cluster of closely packed pairs of

transmembrane channels, the connexons, through which materials of low

MW diffuse from one cell to a neighboring cell.

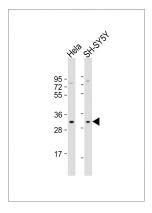
# **Background**

Gap junctions are conduits that allow the direct cell-to-cell passage of small cytoplasmic molecules, including ions, metabolic intermediates, and second messengers, and thereby mediate intercellular metabolic and electrical communication. Gap junction channels consist of connexin protein subunits, which are encoded by a multigene family. GJBs (gap-junction proteins or connexins) play crucial functional roles associated with these channels. Defects in GJB3 have been linked to erythrokeratodermia variabilis (EKV) is an autosomal dominant genodermatosis characterized by transient figurate red patches or hyperkeratosis. Mutations in GJB2 have also been associated with genetically derived hearing impairments, including autosomal recessive nonsyndromic deafness.

#### References

Plantard, L., et al., Hum. Mol. Genet. 12(24):3287-3294 (2003). Rouan, F., et al., Exp. Dermatol. 12(2):191-197 (2003). Mhatre, A.N., et al., Clin. Genet. 63(2):154-159 (2003). Diestel, S., et al., Biochem. Biophys. Res. Commun. 296(3):721-728 (2002). Di, W.L., et al., Hum. Mol. Genet. 11(17):2005-2014 (2002).

## **Images**



All lanes: Anti-GJB3 Antibody (C211) at 1:1000 dilution Lane 1: Hela whole cell lysate Lane 2: SH-SY5Y whole cell lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size: 31 kDa Blocking/Dilution buffer: 5% NFDM/TBST.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.