



Connexin 43 (phospho Tyr265) rabbit pAb

Cat#: orb768396 (Manual)

For research use only. Not intended for diagnostic use.

Product Name Connexin 43 (phospho Tyr265) rabbit pAb

Host species Rabbit

Applications WB;ELISA

Species Cross-Reactivity Human; Mouse; Rat

Recommended dilutions Western Blot: 1/500 - 1/2000. ELISA: 1/10000. Not yet tested in other

applications.

Immunogen Synthesized phospho-peptide around the phosphorylation site of human

Connexin 43 (phospho Ser265)

Specificity Phospho-Connexin 43 (S265) Polyclonal Antibody detects endogenous

levels of Connexin 43 protein only when phosphorylated at S265.

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium

azide..

Storage Store at -20°C. Avoid repeated freeze-thaw cycles.

Protein Name Gap junction alpha-1 protein

Gene Name GJA1

Cellular localization Cell membrane; Multi-pass membrane protein. Cell junction, gap junction.

Endoplasmic reticulum. Localizes at the intercalated disk (ICD) in cardiomyocytes and the proper localization at ICD is dependent on

TMEM65...

Purification The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.





Clonality Polyclonal

Concentration 1 mg/ml

Observed band 43kD

2697 **Human Gene ID**

Human Swiss-Prot Number P17302

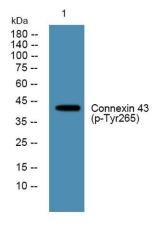
Alternative Names GJA1; GJAL; Gap junction alpha-1 protein; Connexin-43; Cx43; Gap

junction 43 kDa heart protein

Background This gene is a member of the connexin gene family. The encoded protein is a

component of gap junctions, which are composed of arrays of intercellular channels that provide a route for the diffusion of low molecular weight materials from cell to cell. The encoded protein is the major protein of gap junctions in the heart that are thought to have a crucial role in the synchronized contraction of the heart and in embryonic development. A related intronless pseudogene has been mapped to chromosome 5. Mutations in this gene have been associated with oculodentodigital dysplasia,

autosomal recessive craniometaphyseal dysplasia and heart malformations. [provided by RefSeq, May 2014],



Western blot analysis of lysates from K562 cells, primary antibody was diluted at 1:1000, 4° over night