

DNA Binding Protein-7 (DBP-7) Antibody

Rabbit Polyclonal Antibody Catalog # ABV10721

Specification

DNA Binding Protein-7 (DBP-7) Antibody - Product Information

Application WB **Primary Accession** 09P2D1 Other Accession NP 060250 Reactivity Human Host **Rabbit** Clonality **Polyclonal** Isotype Rabbit IgG Calculated MW 335927

DNA Binding Protein-7 (DBP-7) Antibody - Additional Information

Gene ID 55636

Application & Usage Western blot analysis (0.5-4 μg/ml).

However, the optimal conditions should be determined individually. Recombinant DBP-7 can be used as a positive control.

Other Names

DNA Binding Protein 7, DNA Binding Protein-7

Target/Specificity

DBP7

Antibody Form

Liquid

Appearance

Colorless liquid

Formulation

 $100~\mu g$ (0.5 mg/ml) affinity purified rabbit anti-human DBP-7 polyclonal antibody in phosphate buffered saline (PBS), pH 7.2, containing 30% glycerol, 0.5% BSA, 0.01% thimerosal

Handling

The antibody solution should be gently mixed before use.

Reconstitution & Storage

-20 °C

Background Descriptions

Precautions

DNA Binding Protein-7 (DBP-7) Antibody is for research use only and not for use in diagnostic or



therapeutic procedures.

DNA Binding Protein-7 (DBP-7) Antibody - Protein Information

Name CHD7

Synonyms KIAA1416

Function

Probable transcription regulator. Maybe involved in the in 45S precursor rRNA production.

Cellular Location [Isoform 1]: Nucleus

Tissue Location

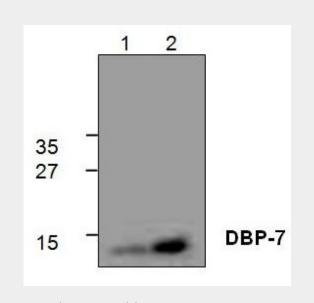
Widely expressed in fetal and adult tissues.

DNA Binding Protein-7 (DBP-7) Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

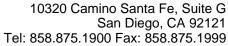
- Western Blot
- Blocking Peptides
- Dot Blot
- <u>Immunohistochemistry</u>
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

DNA Binding Protein-7 (DBP-7) Antibody - Images



Western blot analysis of DBP-7 using recombinant DBP-7. Lane 1:10 ng; Lane 2:50 ng

DNA Binding Protein-7 (DBP-7) Antibody - Background





DNA binding protein 7 (DBP-7) is also known as Chd7. Mutations in the Chd7 gene is believed to be a major implication of CHARGE syndrome. CHARGE syndrome is associated with anomalies such as malformations of the heart, inner ear, retina and cranial nerve defects.