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## **ARX Polyclonal Antibody**

| Catalog No         | YP-Ab-15747   |
|--------------------|---|
| lsotype            | lgG   |
| Reactivity         | Human;Mouse;Rat   |
| Applications       | WB;ELISA  |
| Gene Name          | ARX   |
| Protein Name       | Homeobox protein ARX  |
| Immunogen          | Synthesized peptide derived from ARX . at AA range: 250-330   |
| Specificity        | ARX Polyclonal Antibody detects endogenous levels of ARX protein.   |
| Formulation        | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.   |
| Source             | Polyclonal, Rabbit,IgG  |
| Purification       | The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.   |
| Dilution           | Western Blot: 1/500 - 1/2000. ELISA: 1/5000. Not yet tested in other applications.  |
| Concentration      | 1 mg/ml   |
| Purity             | ≥90%  |
| Storage Stability  | -20°C/1 year  |
| Synonyms           | ARX; Homeobox protein ARX; Aristaless-related homeobox  |
| Observed Band      | 55kD  |
| Cell Pathway       | Nucleus .   |
| Tissue Specificity | Expressed predominantly in fetal and adult brain and skeletal muscle. Expression is specific to the telencephalon and ventral thalamus. There is an absence of expression in the cerebellum throughout development and also in adult.   |
| Function           | disease:Defects in ARX are a cause of Partington syndrome (PRTS)<br>[MIM:309510]; also known as X-linked syndromic mental retardation 1 (MRXS1).<br>PRTS is characterized by mental retardation, episodic dystonic hand movements,<br>and dysarthria.,disease:Defects in ARX are the cause of agenesis of corpus<br>callosum with abnormal genitalia (ACC with abnormal genitalia) [MIM:300004].<br>ACC with abnormal genitalia consists of a brain and genital malformations<br>syndrome.,disease:Defects in ARX are the cause of epileptic encephalopathy<br>early infantile type 1 (EIEE1) [MIM:308350]; also known as myoclonic epilepsy<br>X-linked with intellectual disability and spasticity, X-linked West syndrome or<br>X-linked infantile spasm syndrome (ISSX). EIEE1 is a severe form of epilepsy<br>characterized by frequent tonic seizures or spasms beginning in infancy with a<br>specific EEG finding of suppression-burst patterns, characterize |
| Background         | This gene is a homeobox-containing gene expressed during development. The expressed protein contains two conserved domains, a C-peptide (or aristaless  |



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domain) and the prd-like class homeobox domain. It is a member of the group-II<br/>aristaless-related protein family whose members are expressed primarily in the<br/>central and/or peripheral nervous system. This gene is thought to be involved in<br/>CNS development. Expansion of a polyalanine tract and other mutations in this<br/>gene cause X-linked mental retardation and epilepsy. [provided by RefSeq, Jul<br/>2016],matters needing<br/>attentionAvoid repeated freezing and thawing!Usage suggestionsThis product can be used in immunological reaction related experiments. For<br/>more information, please consult technical personnel.

## **Products Images**

