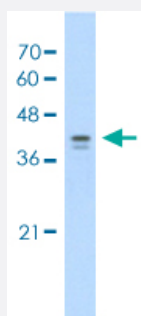


# LOR polyclonal antibody

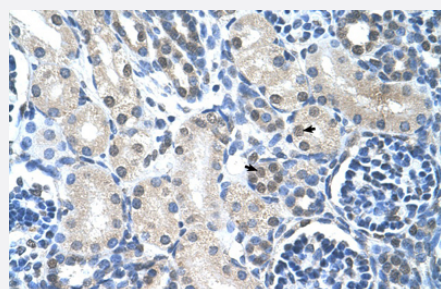
Catalog # PAB29914      Size 100 uL

## Applications



### Western Blot (Cell lysate)

Western blot analysis of HepG2 cell lysate with LOR polyclonal antibody (Cat # PAB29914).



### Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining (Formalin-fixed paraffin-embedded sections) of human kidney with LOR polyclonal antibody (Cat # PAB29914).

## Specification

Product Description	Rabbit polyclonal antibody raised against partial synthetic protein of human LOR.
Immunogen	A synthetic peptide corresponding to amino acids 58-107 of human LOR.
Sequence	GYSGGGCGGGSSGGGGGGGIGGCGGGSGGSVKYSGGGGSSGGSGCFSSG
Host	Rabbit
Theoretical MW (kDa)	26
Reactivity	Human
Form	Liquid

<b>Purification</b>	Protein A purification
<b>Isotype</b>	IgG
<b>Recommend Usage</b>	Immunohistochemistry (1:250) Western Blot (1:1000) The optimal working dilution should be determined by the end user.
<b>Storage Buffer</b>	In 1X PBS , pH 7.4 (2% sucrose, 0.09% sodium azide).
<b>Storage Instruction</b>	Store at 4°C for up to one week. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
<b>Note</b>	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

## Applications

- Western Blot (Cell lysate)

Western blot analysis of HepG2 cell lysate with LOR polyclonal antibody (Cat # PAB29914).

- Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining (Formalin-fixed paraffin-embedded sections) of human kidney with LOR polyclonal antibody (Cat # PAB29914).

## Gene Info — LOR

<b>Entrez GeneID</b>	<a href="#">4014</a>
<b>Protein Accession#</b>	<a href="#">P23490</a>
<b>Gene Name</b>	LOR
<b>Gene Alias</b>	MGC111513
<b>Gene Description</b>	loricrin
<b>Omim ID</b>	<a href="#">152445 602036 604117</a>
<b>Gene Ontology</b>	<a href="#">Hyperlink</a>
<b>Gene Summary</b>	This gene encodes loricrin, a major protein component of the cornified cell envelope found in terminally differentiated epidermal cells. Mutations in this gene are associated with Vohwinkel's syndrome and progressive symmetric erythrokeratoderma, both inherited skin diseases. [provided by RefSeq]

Other Designations

OTTHUMP00000015823

## Disease

- [Cleft Lip](#)
- [Cleft Palate](#)
- [Dermatitis](#)
- [Genetic Predisposition to Disease](#)