## LOR rabbit monoclonal antibody

Catalog # H00004014-K

ocification

Size 100 ug x up to 3

| Specification           |   |
|-------------------------|---|
| Product Description     | Rabbit monoclonal antibody raised against a human LOR peptide using ARM Technology.   |
| Immunogen               | A synthetic peptide of human LOR is used for rabbit immunization.<br>Customer or Abnova will decide on the preferred peptide sequence.  |
| Host                    | Rabbit  |
| Library Construction    | Non-fusion antibody library from rabbit spleen (ARM Technology).  |
| Expression              | Overexpression vector and transfection into 293H cell line.   |
| Reactivity              | Human   |
| Purification            | Protein A   |
| lsotype                 | lgG   |
| Quality Control Testing | Antibody reactive against human LOR peptide by ELISA and mammalian transfected lysate by West ern Blot.   |
| Storage Buffer          | In 1x PBS, pH 7.4   |
| Storage Instruction     | Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.  |
| Deliverable             | Up to three rabbit IgG clones of 100 ug each will be delivered to customer.   |
| Note                    | <ol> <li>Customer may provide cell or tissue lysate for antibody screening.</li> <li>Rabbit monoclonal antibody generated by ARM technology is amenable to antibody engineering in<br/>cluding F(ab)<sub>2</sub>, lgG, scFv and different Fc and non-Fc conjugates per customer request.</li> </ol> |

## Applications

• Western Blot (Transfected lysate)

Protocol Download



• ELISA

| Gene Info — LOR     |  |
|---------------------|--|
| Entrez GenelD       | <u>4014</u>  |
| GeneBank Accession# | LOR  |
| Gene Name           | LOR  |
| Gene Alias          | MGC111513  |
| Gene Description    | loricrin   |
| Omim ID             | <u>152445 602036 604117</u>  |
| Gene Ontology       | Hyperlink  |
| Gene Summary        | This gene encodes loricrin, a major protein component of the cornified cell envelope found in term inally differentiated epidermal cells. Mutations in this gene are associated with Vohwinkel's syndr ome and progressive symmetric erythrokeratoderma, both inherited skin diseases. [provided by RefSeq |
| Other Designations  | OTTHUMP0000015823  |

## Disease

- Cleft Lip
- <u>Cleft Palate</u>
- Dermatitis
- <u>Genetic Predisposition to Disease</u>