
$\overline{\text { Applications }}$
$\overline{\text { Background Information }}$

Tested Applications:
WB, ELISA
Cited Applications:
WB, IF
Species Specificity:
human, mouse, rat
Cited Species:
human

Positive Controls:
WB : mouse liver tissue, rat liver tissue

Background Information
FAH is also named as FAA(fumarylacetoacetase), the last enzyme in tyrosine degradation, functionally important in the liver and kidney(PMID:8473520). FAH is a metabolic enzyme catalyzing the last step of tyrosine and phenylalanine catabolism: the hydrolysis of fumarylacetoacetate into acetoacetate and fumarate. In humans, deficiency of this activity is associated with the metabolic disease hereditary tyrosinaemia type I, which is also known as hepatorenal tyrosinaemia(PMID:17768357).

## Notable Publications

## Storage

| Author | Pubmed ID | Journal | Application |
| :--- | :--- | :--- | :--- |
| Jinghai Hu | 27626805 | PLoS One |  |
| Guodong Lian | 29115543 | Oncol Rep |  |
| Peng-Ming Yu | 32532183 | Cartilage | WB |

Storage:
Store at $-20^{\circ} \mathrm{C}$. Stable for one year after shipment.

Storage Buffer:
PBS with $0.02 \%$ sodium azide and $50 \%$ glycerol pH 7.3 .
Aliquoting is unnecessary for $-20^{\circ} \mathrm{C}$ storage

For technical support and original validation data for this product please contact:
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Selected Validation Data


Various lysates were subjected to SDS PAGE
followed by western blot with 14928-1-AP (FAH antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours.

