

# DRP1 (Phospho-Ser637) Antibody

货号: **AYP4619**

## 产品信息

|       |  |
|-------|--|
| 反应    | Human,Mouse,Rat  |
| 宿主    | Rabbit   |
| 克隆性   | Polyclonal   |
| 预测反应  |  |
| 应用    | <a href="#">WB</a> <a href="#">IHC</a> <a href="#">ELISA</a>   |
| 推荐浓度  | <b>WB:</b> 1:500 - 1:2000<br><b>IHC:</b> 1:50 - 1:200  |
| 理论分子量 | 60kDa/78kDa/79kDa/80kDa/81kDa/82kDa/83kDa  |
| 实测分子量 |  |
| 形式    | Liquid   |
| 保存条件  | Store at -20°C. Avoid freeze / thaw cycles.<br>Buffer: PBS with 0.75% BSA,50% glycerol,pH7.3.  |
| 偶联物   | Unconjugated   |
| 阳性对照  | Mouse brain,Rat brain  |
| 细胞定位  | Cytoplasm,Cytoplasmic vesicle,Endomembrane system,Golgi apparatus,Membrane,Mitochondrion outer membrane,Peripheral membrane protein,Peroxisome,clathrin-coated pit,cytosol,secretory vesicle,synaptic vesicle membrane |
| 纯化    | Affinity purification  |

## 抗原信息

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|------|---|
| 抗原信息 | Synthesized peptide derived from Human DRP1 (Phospho-Ser637). |
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## 靶点信息

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|------|--|
| 研究背景 | This gene encodes a member of the dynamin superfamily of GTPases. The encoded protein mediates mitochondrial and peroxisomal division, and is involved in developmentally regulated apoptosis and programmed necrosis. Dysfunction of this gene is implicated in several neurological disorders, including Alzheimer's disease. Mutations in this gene are associated with the autosomal dominant disorder, encephalopathy, lethal, due to defective mitochondrial and peroxisomal fission (EMPF). Alternative splicing results in multiple transcript variants encoding different isoforms. |
| 基因ID | 10059  |

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|-------|---|
| 基因名   | DNM1L   |
| Swiss | O00429  |
| 别名    | DNM1L;DLP1;DRP1;DVLP;DYMPLE;EMPF;EMPF1;HDYNIV |

产品验证

实验步骤

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