

NUB1 Rabbit pAb

货号: B21946

产品信息

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| 反应 | Human,Mouse,Rat |
| 宿主 | Rabbit |
| 克隆性 | Polyclonal |
| 预测反应 | |
| 应用 | WB IF/ICC |
| 推荐浓度 | WB: 1:500 - 1:2000 IF/ICC: 1:50 - 1:200 |
| 理论分子量 | 69kDa/70kDa |
| 实测分子量 | 71kDa |
| 形式 | Liquid |
| 保存条件 | Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.01% thiomersal,50% glycerol,pH7.3. |
| 偶联物 | Unconjugated |
| 阳性对照 | Mouse lung,Rat lung |
| 细胞定位 | Nucleus |
| 纯化 | Affinity purification |

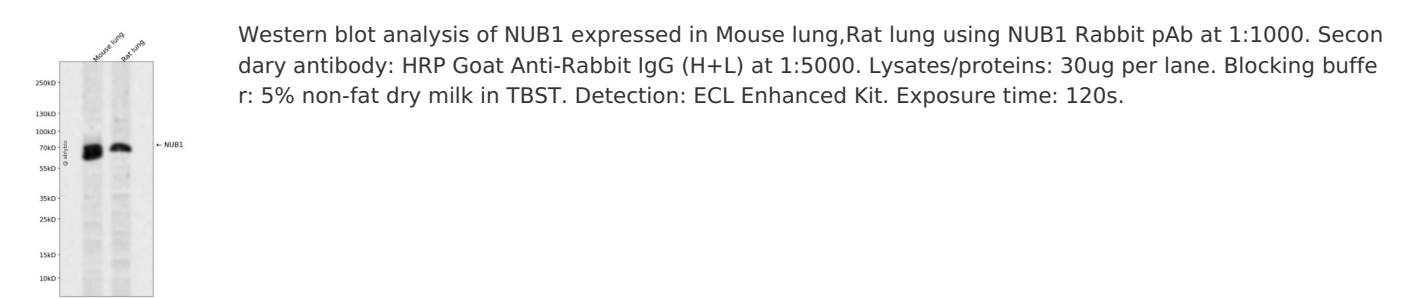
抗原信息

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|------|---|
| 抗原信息 | A synthetic peptide corresponding to a sequence within amino acids 1-100 of human IFT140 (NP_001350 458.1). |
| 序列 | MAQKKYLQAKLTQFLREDRIQLWKPPYTDENKKVGLALKDLAKQYSDRLECCENEVEKVIEEIRCKAIERGTGNDNYRTTGI ATIEVFLPPRLKKDRKNL |

靶点信息

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| 研究背景 | This gene encodes a protein that functions as a negative regulator of NEDD8, a ubiquitin-like protein that conjugates with cullin family members in order to regulate vital biological events. The protein encoded by this gene regulates the NEDD8 conjugation system post-transcriptionally by recruiting NEDD8 and its conjugates to the proteasome for degradation. This protein interacts with the product of the AIPL1 gene, which is associated with Leber congenital amaurosis, an inherited retinopathy, and mutations in that gene can abolish interaction with this protein, which may contribute to the pathogenesis. This protein is also known to accumulate in Lewy bodies in Parkinson's disease and dementia with Lewy bodies, and in glial cytoplasmic inclusions in multiple system atrophy, with this abnormal accumulation being specific to alpha-synucleinopathy lesions. Alternative splicing of this gene results in multiple transcript variants. |
| 基因ID | 51667 |
| 基因名 | NUB1 |
| Swiss | Q9Y5A7 |
| 别名 | NUB1; BS4; NUB1L; NYREN18; NEDD8 ultimate buster 1 |

产品验证



实验步骤

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