## NLN Antibody

| Product Code | CSB-PA863161LA01HU |
| :---: | :---: |
| Storage | Upon receipt, store at $-20^{\circ} \mathrm{C}$ or $-80^{\circ} \mathrm{C}$. Avoid repeated freeze. |
| Uniprot No. | Q9BYT8 |
| Immunogen | Recombinant Human Neurolysin, mitochondrial protein (109-241AA) |
| Raised In | Rabbit |
| Species Reactivity | Human |
| Tested Applications | ELISA, WB, IHC, IF; Recommended dilution: WB:1:500-1:5000, <br> IHC:1:200-1:500, IF:1:50-1:200 |
| Form | Liquid |
| Conjugate | Non-conjugated |
| Storage Buffer | Preservative: 0.03\% Proclin 300 <br> Constituents: $50 \%$ Glycerol, 0.01M PBS, pH 7.4 |
| Purification Method | >95\%, Protein G purified |
| Isotype | $\lg G$ |
| Clonality | Polyclonal |
| Alias | Neurolysin, mitochondrial (EC 3.4.24.16) (Angiotensin-binding protein) (Microsomal endopeptidase) (MEP) (Mitochondrial oligopeptidase M) (Neurotensin endopeptidase), NLN, AGTBP KIAA1226 |
| Immunogen Species | Homo sapiens (Human) |
| Research Area | Tags \& Cell Markers |
| Target Names | NLN |
| Image | $120 \mathrm{KD} \rightarrow$$90 \mathrm{KD} \rightarrow$$50 \mathrm{KD} \rightarrow$$35 \mathrm{KD} \rightarrow$$25 \mathrm{KD} \rightarrow$$\quad$Western Blot <br> Positive WB detected in: 293 whole cell lysate, <br> PC-3 whole cell lysate, HepG2 whole cell lysate, <br> LO2 whole cell lysate, A549 whole cell lysate <br> All lanes: NLN antibody at $3.2 \mu \mathrm{~g} / \mathrm{ml}$ <br> Secondary <br> Goat polyclonal to rabbit IgG at $1 / 50000$ dilution <br> Predicted band size: 81 kDa <br> Observed band size: 81 kDa |



IHC image of CSB-PA863161LA01HU diluted at 1:300 and staining in paraffin-embedded human pancreatic cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with $10 \%$ normal goat serum 30 min at RT. Then primary antibody ( $1 \%$ BSA) was incubated at $4^{\circ} \mathrm{C}$ overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.


Immunofluorescence staining of HepG2 cells with CSB-PA863161LA01HU at 1:100, counterstained with DAPI. The cells were fixed in $4 \%$ formaldehyde, permeabilized using 0.2\% Triton X-100 and blocked in $10 \%$ normal Goat Serum. The cells were then incubated with the antibody overnight at $4^{\circ} \mathrm{C}$. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit $\lg G(\mathrm{H}+\mathrm{L})$.

