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# Rabbit Anti-CCDC126 antibody

SL18333R

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| **Product Name:** | CCDC126 |
| **Chinese Name:** | 卷曲螺旋结构域蛋白126抗体 |
| **Alias:** | LOC90693; alpha 1,3(6) mannosylglycoprotein; beta 1,6 N acetyl glucosaminyltransferase like; CC126_HUMAN; Ccdc126; coiled coil domain containing 126; Coiled-coil domain-containing protein 126; FLJ23031; MGC104248; OTTHUMP00000158577. |
| **Organism Species:** | Rabbit |
| **Clonality:** | Polyclonal |
| **React Species:** | Human,Mouse,Rat,Dog,Pig,Cow,Horse,Rabbit,Sheep, |
| **Applications:** | ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=1:100-500IF=1:100-500（Paraffin sections need antigen repair）<br>not yet tested in other applications.<br>optimal dilutions/concentrations should be determined by the end user. |
| **Molecular weight:** | 13kDa |
| **Cellular localization:** | Secretory protein |
| **Form:** | Lyophilized or Liquid |
| **Concentration:** | 1mg/ml |
| **immunogen:** | KLH conjugated synthetic peptide derived from human CCDC126:27-100/140 |
| **Lsotype:** | IgG |
| **Purification:** | affinity purified by Protein A |
| **Storage Buffer:** | 0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol. |
| **Storage:** | Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20°C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C. |
| **PubMed:** | PubMed |
| **Product Detail:** | CCDC126 is a 140 amino acid secreted protein encoded by a gene mapping to human chromosome 7. Chromosome 7 is about 158 milllion bases long, encodes over 1000 genes and makes up about 5% of the human genome. Chromosome 7 has been linked to Osteogenesis imperfecta, Pendred syndrome, Lissencephaly, Citrullinemia and |

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| | Shwachman-Diamond syndrome. The deletion of a portion of the q arm of chromosome 7 is associated with Williams-Beuren syndrome, a condition characterized by mild mental retardation, an unusual comfort and friendliness with strangers and an elfin appearance. Deletions of portions of the q arm of chromosome 7 are also seen in a number of myeloid disorders including cases of acute myelogenous leukemia and myelodysplasia.<br><br>**Subcellular Location:**<br>Secreted.<br><br>**SWISS:**<br>Q96EE4<br><br>**Gene ID:**<br>90693<br><br>**Database links:**<br><br>Entrez Gene: 90693 Human<br><br>SwissProt: Q96EE4 Human<br><br>Unigene: 232296 Human<br><br>**Important Note:**<br>This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications. |