

Product datasheet

Anti-Corneodesmosin/S protein antibody ab90517

Overview

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|----------------------------|---|
| Product name | Anti-Corneodesmosin/S protein antibody |
| Description | Rabbit polyclonal to Corneodesmosin/S protein |
| Host species | Rabbit |
| Tested applications | Suitable for: WB, IHC-P, IHC-Fr |
| Species reactivity | Reacts with: Human |
| Immunogen | Synthetic peptide corresponding to Human Corneodesmosin/S protein (N terminal). |
| General notes | Protein previously labeled as Corneodesmosin. |

Properties

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|-----------------------------|---|
| Form | Liquid |
| Storage instructions | Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw cycles. |
| Storage buffer | Constituent: Whole serum |
| Purity | Whole antiserum |
| Clonality | Polyclonal |
| Isotype | IgG |

Applications

Our [Abpromise guarantee](#) covers the use of **ab90517** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

| Application | Abreviews | Notes |
|-------------|-----------|---|
| WB | | 1/500 - 1/2000. Predicted molecular weight: 51 kDa. |
| IHC-P | | Use at an assay dependent concentration. |
| IHC-Fr | | Use at an assay dependent concentration. |

Target

| | |
|-------------------------------|--|
| Function | Important for the epidermal barrier integrity. |
| Tissue specificity | Exclusively expressed in skin. |
| Involvement in disease | <p>Defects in CDSN are a cause of hypotrichosis simplex of the scalp (HTSS) [MIM:146520]; also known as hypotrichosis Spanish type. HTSS is an autosomal dominant form of isolated alopecia. Affected individuals have normal hair in early childhood but experience progressive loss of scalp hair beginning in the middle of the first decade and almost complete baldness by the third decade.</p> <p>Defects in CDSN are the cause of peeling skin syndrome type B (BPSS) [MIM:270300]; also known as peeling skin syndrome or deciduous skin or keratolysis exfoliativa congenita. BPSS is a genodermatosis characterized by the continuous shedding of the outer layers of the epidermis, associated with pruritus and atopy. It is an ichthyosiform erythroderma characterized by lifelong patchy peeling of the entire skin with onset at birth or shortly thereafter. Several patients have been reported with high IgE levels.</p> |
| Cellular localization | Secreted. Found in corneodesmosomes, the intercellular structures that are involved in desquamation. |

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