# DISEASE:
# Peeling skin syndrome type B

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| **NAME:** | Peeling skin syndrome type B |
| **DESCRIPTION:** | A form of generalized peeling skin syndrome (PSS) characterized by superficial patchy peeling of the entire skin with underlying erythroderma, pruritus, and atopy. |
| **ORPHACODE:** | 263553 |
| **SYNONYMS:** | Generalized peeling skin disease type B<br>Generalized peeling skin syndrome type B<br>Inflammatory peeling skin disease<br>Inflammatory peeling skin syndrome<br>PSS type B<br>PSS1<br>Peeling skin syndrome 1 |
| **XREF(S):** | Orphanet<br>OMIM<br>ICD-10 |
| **ANALYTE(S):** | CDSN |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

# RELATED CONTENT

## Related Genetic Tests

- Epidermolysis bullosa (gene panel)
- Ichthyosis (gene panel)

## Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

## Related Analytes

- corneodesmosin

## Related Gene Panels

- Epidermolysis bullosa and bladder diseases (60 genes) - KUL
- Ichthyosis and erythroderma (98 genes) - KUL