

**DISEASE:**

**Autosomal dominant generalized dystrophic epidermolysis bullosa**

<b>NAME:</b>	Autosomal dominant generalized dystrophic epidermolysis bullosa
<b>DESCRIPTION:</b>	A rare dystrophic epidermolysis bullosa (DEB) characterized by generalized blistering, milia formation, atrophic scarring, and dystrophic nails.
<b>ORPHACODE:</b>	231568
<b>SYNOMYS:</b>	Generalized DDEB
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">ICD-10</a> <a href="#">OMIM</a>
<b>ANALYTE(S):</b>	<a href="#">COL7A1</a>
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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## RELATED CONTENT

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### Related Genetic Tests

- [Epidermolysis bullosa \(gene panel\)](#)

### Related Laboratories

- [Centrum Menselijke Erfelijkheid - KUL](#)

### Related Analytes

- [collagen type VII alpha 1 chain](#)

### Related Gene Panels

- [Epidermolysis bullosa and bladder diseases \(60 genes\) - KUL](#)

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