Open Peer Review on Qeios

Junctional epidermolysis bullosa inversa

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Junctional epidermolysis bullosa inversa. ORPHA:79405

Junctional epidermolysis bullosa inversa is a rare severe subtype of junctional epidermolysis bullosa (JEB, see this term) characterized by blistering and erosions confined to intertriginous skin sites, the esophagus, and vagina.