# Autosomal dominant hyper-IgE syndrome

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant hyper-IgE syndrome. ORPHA:2314*

Autosomal dominant hyper-IgE syndrome (AD-HIES) is a very rare primary immunodeficiency disorder characterized by the clinical triad of high serum IgE (>2000 IU/ml), recurring staphylococcal skin abscesses, and recurrent pneumonia with formation of pneumatoceles.