Haim-Munk syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Haim-Munk syndrome. ORPHA:2342

Haim-Munk syndrome (HMS) is characterized by palmoplantar hyperkeratosis, severe early-onset periodontitis, onychogryposis, pes planus, arachnodactyly and acroosteolysis.