Autoimmune pulmonary alveolar proteinosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autoimmune pulmonary alveolar proteinosis. ORPHA:747

Pulmonary alveolar proteinosis (PAP) is a rare lung disease characterized by the accumulation of a lipoproteinaceous substance in the distal air spaces which positively stains with periodic acid-Schiff (PAS).