

[Open Peer Review on Qeios](#)

Mu-heavy chain disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Mu-heavy chain disease. ORPHA:100024

Mu-heavy chain disease (mu-HCD) is a type of HCD (see this term) characterized by the production of incomplete monoclonal mu-heavy chains without associated light chains. The clinical presentation resembles that of patients with chronic lymphocytic leukemia/small lymphocytic lymphoma (CLL/SLL; see this term).