

Use this checklist to determine if a patient meets the restrictions for funding in the **hospital setting**. For more details, refer to [Section H](#) of the Pharmaceutical Schedule. For community funding, see the [Special Authority Criteria](#).

PRESCRIBER

Name:

Ward:

PATIENT:

Name:

NHI:

Idursulfase

INITIATION

Re-assessment required after 24 weeks

Prerequisites (tick boxes where appropriate)

☐ Prescribed by, or recommended by a metabolic physician, or in accordance with a protocol or guideline that has been endorsed by the Te Whatu Ora Hospital.

and

☐ The patient has been diagnosed with Hunter Syndrome (mucopolysaccharidosis II)

and

☐ Diagnosis confirmed by demonstration of iduronate 2-sulfatase deficiency in white blood cells by either enzyme assay in cultured skin fibroblasts

or

☐ Detection of a disease causing mutation in the iduronate 2-sulfatase gene

and

☐ Patient is going to proceed with a haematopoietic stem cell transplant (HSCT) within the next 3 months and treatment with idursulfase would be bridging treatment to transplant

and

☐ Patient has not required long-term invasive ventilation for respiratory failure prior to starting Enzyme Replacement Therapy (ERT)

and

☐ Idursulfase to be administered for a total of 24 weeks (equivalent to 12 weeks pre- and 12 weeks post-HSCT) at doses no greater than 0.5 mg/kg every week

I confirm that the above details are correct:

Signed: Date: