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.

PRES	CRIBER		PATIENT:
Name	e:		Name:
Ward	:		NHI:
Terip	paratide		
Re-a		nt required after 18 months (tick boxes where appropriate)	
	O	The patient has severe, established osteoporosis	
	and	The patient has a documented T-score less than or equal to -3	8.0 (see Notes)
	and	The patient has had two or more fractures due to minimal trau	ma
	and	The patient has experienced at least one symptomatic new fra	acture after at least 12 months' continuous therapy with a funded

Note:

- a) The bone mineral density (BMD) measurement used to derive the T-score must be made using dual-energy x-ray absorptiometry (DXA). Quantitative ultrasound and quantitative computed tomography (QCT) are not acceptable
- b) Antiresorptive agents and their adequate doses for the purposes of this restriction are defined as: alendronate sodium tab 70 mg or tab 70 mg with colecalciferol 5,600 iu once weekly; raloxifene hydrochloride tab 60 mg once daily; zoledronic acid 5 mg per year. If an intolerance of a severity necessitating permanent treatment withdrawal develops during the use of one antiresorptive agent, an alternate antiresorptive agent must be trialled so that the patient achieves the minimum requirement of 12 months' continuous therapy.
- c) A vertebral fracture is defined as a 20% or greater reduction in height of the anterior or mid portion of a vertebral body relative to the posterior height of that body, or a 20% or greater reduction in any of these heights compared to the vertebral body above or below the affected vertebral body.

I confirm that the above details are correct:	
Signed:	. Date: