

APPLICANT (stamp or sticker acceptable)	PATIENT NHI:	REFERRER Reg No:
Reg No:	First Names:	First Names:
Name:	Surname:	Surname:
Address:	DOB:	Address:
.....	Address:
.....
Fax Number:	Fax Number:

Octreotide long-acting

Initial application — Malignant Bowel Obstruction

Applications from any relevant practitioner. Approvals valid for 2 months.

Prerequisites(tick boxes where appropriate)

- The patient has nausea* and vomiting* due to malignant bowel obstruction*
- and
- Treatment with antiemetics, rehydration, antimuscarinic agents, corticosteroids and analgesics for at least 48 hours has failed
- and
- Octreotide to be given at a maximum dose 1500 mcg daily for up to 4 weeks

Note: Indications marked with * are unapproved indications.

Renewal — Malignant Bowel Obstruction

Current approval Number (if known):.....

Applications from any relevant practitioner. Approvals valid for 3 months.

Prerequisites(tick box where appropriate)

- The treatment remains appropriate and the patient is benefiting from treatment

Initial application — Acromegaly

Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 3 months.

Prerequisites(tick boxes where appropriate)

- The patient has acromegaly
- and
- Treatment with surgery, radiotherapy and a dopamine agonist has failed
- or
- Treatment with octreotide is for an interim period while awaiting the effects of radiotherapy and a dopamine agonist has failed
- or
- The patient is unwilling, or unable, to undergo surgery and/or radiotherapy

Renewal — Acromegaly

Current approval Number (if known):.....

Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 2 years.

Prerequisites(tick boxes where appropriate)

- IGF1 levels have decreased since starting octreotide
- and
- The treatment remains appropriate and the patient is benefiting from treatment

Note: In patients with Acromegaly octreotide treatment should be discontinued if IGF1 levels have not decreased after 3 months treatment. In patients treated with radiotherapy octreotide treatment should be withdrawn every 2 years, for 1 month, for assessment of remission. Octreotide treatment should be stopped where there is biochemical evidence of remission (normal IGF1 levels) following octreotide treatment withdrawal for at least 4 weeks

I confirm the above details are correct and that in signing this form I understand I may be audited.

Signed: Date:

Post application to Ministry of Health, Private Bag 3015, Wanganui – email: customerservice@health.govt.nz

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Octreotide long-acting - *continued*

Initial application — Other Indications

Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 2 years.

Prerequisites(tick boxes where appropriate)

<input type="checkbox"/>	VIPomas and Glucagonomas - for patients who are seriously ill in order to improve their clinical state prior to definitive surgery
or	
<input type="checkbox"/>	Gastrinoma
and	
<input type="checkbox"/>	Patient has failed surgery
or	
<input type="checkbox"/>	Patient in metastatic disease after H2 antagonists (or proton pump inhibitors) have failed
or	
<input type="checkbox"/>	Insulinomas
and	
<input type="checkbox"/>	Surgery is contraindicated or has failed
or	
<input type="checkbox"/>	For pre-operative control of hypoglycaemia and for maintenance therapy
or	
<input type="checkbox"/>	Carcinoid syndrome (diagnosed by tissue pathology and/or urinary 5HIAA analysis)
and	
<input type="checkbox"/>	Disabling symptoms not controlled by maximal medical therapy

Note: The use of octreotide in patients with fistulae, oesophageal varices, miscellaneous diarrhoea and hypotension will not be funded as a Special Authority item

Renewal — Other Indications

Current approval Number (if known):.....

Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 2 years.

Prerequisites(tick box where appropriate)

The treatment remains appropriate and the patient is benefiting from treatment

Initial application — pre-operative acromegaly

Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 12 months.

Prerequisites(tick boxes where appropriate)

<input type="checkbox"/>	Patient has acromegaly
and	
<input type="checkbox"/>	Patient has a large pituitary tumour, greater than 10 mm at its widest
and	
<input type="checkbox"/>	Patient is scheduled to undergo pituitary surgery in the next six months

I confirm the above details are correct and that in signing this form I understand I may be audited.

Signed: Date:

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