

**APPLICANT** (stamp or sticker acceptable)      **PATIENT NHI:** .....      **REFERRER** Reg No: .....

Reg No: .....      First Names: .....      First Names: .....

Name: .....      Surname: .....      Surname: .....

Address: .....      DOB: .....      Address: .....

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Fax Number: .....      Fax Number: .....

**Brentuximab**

**Initial application — CD30 positive systemic anaplastic large-cell lymphoma**  
Applications from any relevant practitioner. Approvals valid for 12 months.  
**Prerequisites**(tick boxes where appropriate)

Patient is currently on treatment with brentuximab vedotin and met all the following criteria prior to commencing treatment

or

Patient has CD30 positive systemic anaplastic large-cell lymphoma  
and  
 Patient must have histological confirmation of CD30 expression  
and  
 Patient must not have received prior treatment with curative intent chemotherapy for this condition  
and  
 Treatment must be in combination with cyclophosphamide, anthracycline, and steroids for a maximum of 8 cycles  
and  
 Brentuximab vedotin is to be administered at doses no greater than 1.8 mg/kg every 3 weeks

**Initial application — relapsed/refractory Hodgkin lymphoma**  
Applications from any relevant practitioner. Approvals valid for 6 months.  
**Prerequisites**(tick boxes where appropriate)

Patient has relapsed/refractory CD30-positive Hodgkin lymphoma after two or more lines of chemotherapy  
and  
 Patient is ineligible for autologous stem cell transplant

or

Patient has relapsed/refractory CD30-positive Hodgkin lymphoma  
and  
 Patient has previously undergone autologous stem cell transplant

and  
 Patient has not previously received funded brentuximab vedotin  
and  
 Response to brentuximab vedotin treatment is to be reviewed after a maximum of 6 treatment cycles  
and  
 Brentuximab vedotin to be administered at doses no greater than 1.8 mg/kg every 3 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 Health New Zealand, Private Bag 3015, Wanganui – email: [customerservice@health.govt.nz](mailto:customerservice@health.govt.nz)

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**Brentuximab - continued**

**Renewal — relapsed/refractory Hodgkin lymphoma**

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

Applications from any relevant practitioner. Approvals valid for 9 months.

**Prerequisites**(tick boxes where appropriate)

- Patient has achieved a partial or complete response to brentuximab vedotin after 6 treatment cycles
- and  Treatment remains clinically appropriate and the patient is benefitting from treatment and treatment is being tolerated
- and  Patient is to receive a maximum of 16 total cycles of brentuximab vedotin treatment

**Initial application — relapsed/refractory anaplastic large cell lymphoma**

Applications from any relevant practitioner. Approvals valid for 9 months.

**Prerequisites**(tick boxes where appropriate)

- Patient has relapsed/refractory CD30-positive systemic anaplastic large cell lymphoma
- and  Patient has an ECOG performance status of 0-1
- and  Patient has not previously received brentuximab vedotin
- and  Response to brentuximab vedotin treatment is to be reviewed after a maximum of 6 treatment cycles
- and  Brentuximab vedotin to be administered at doses no greater than 1.8 mg/kg every 3 weeks

**Renewal — relapsed/refractory anaplastic large cell lymphoma**

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

Applications from any relevant practitioner. Approvals valid for 9 months.

**Prerequisites**(tick boxes where appropriate)

- Patient has experienced a partial or complete response to brentuximab vedotin after 6 treatment cycles
- and  Treatment remains clinically appropriate and the patient is benefitting from treatment and treatment is being tolerated
- and  Patient is to receive a maximum of 16 total cycles of brentuximab vedotin treatment

**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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