APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Form SA2629
August 2026

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

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

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.....	.....	.....
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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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