

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: NHI:

Brentuximab

INITIATION – relapsed/refractory Hodgkin lymphoma

Re-assessment required after 6 months

Prerequisites (tick boxes where appropriate)

and 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

and 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

CONTINUATION – relapsed/refractory Hodgkin lymphoma

Re-assessment required after 9 months

Prerequisites (tick boxes where appropriate)

and 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

INITIATION – anaplastic large cell lymphoma

Re-assessment required after 9 months

Prerequisites (tick boxes where appropriate)

and 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

I confirm that the above details are correct:

Signed: Date:

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PREScriBER

Name:

Ward: NHI:

Brentuximab - *continued*

CONTINUATION – anaplastic large cell lymphoma

Re-assessment required after 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

I confirm that the above details are correct:

Signed: Date: