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	PATIENT:
Name:	
Ward:	NHI:
Gemtuzuma	b ozogamicin
INITIATION	
Prerequisites	(tick boxes where appropriate)
and	Patient has not received prior chemotherapy for this condition
and	Patient has de novo CD33-positive acute myeloid leukaemia
and	Patient does not have acute promyelocytic leukaemia
and _	Gemtuzumab ozogamicin will be used in combination with standard anthracycline and cytarabine (AraC)
O	Patient is being treated with curative intent
and and	Patient's disease risk has been assessed by cytogenetic testing to be good or intermediate
0	Patient must be considered eligible for standard intensive remission induction chemotherapy with standard anthracycline and cytarabine (AraC)
and	Gemtuzumab ozogamicin to be funded for one course only (one dose at 3 mg per m² body surface area or up to 2 vials of 5 mg as separate doses)

Note: Acute myeloid leukaemia excludes acute promyelocytic leukaemia and acute myeloid leukaemia that is secondary to another haematological disorder (eg myelodysplasia or myeloproliferative disorder).

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