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

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: