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:
Gemtuzumab 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)
and	Patient is being treated with curative intent
and	Patient's disease risk has been assessed by cytogenetic testing to be good or intermediate
	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).