Click here to access the guidelines/NICE algorithm							
N&WICS - TA323 - Erythropoiesis		in alfa, beta, theta and zei incer having chemothera		tin alfa) - Treatin	g anaemia in		
Before providing patient identifiable of parent/legal guardian/carer) has give CSU for processing this funding requ	en appropriate explicit consent	t for sensitive personal inform	mation on this form				
If there is more than one NICE-approabout the advantages and disadvant patient is likely to adhere to treatme and price per dose) unless an order	ages of the treatments availal nt. The most appropriate, lea	ble. This has taken into cons st expensive, will be chosen	sideration therapeu	tic need and wheth	er or not the		
Patient NHS No:	Trust:		Practice Name:				
Patient Hospital No:	Consultant Making Request:		Practice Postcode:				
Patient's Initials and DoB:			Practice Code:				
Notification Email Address:	(@NHS.ne	et account ONLY)	Contact name & number:				
Start date of requested treatment:	Committee	e □NHS care □Hospital	Sub-Type:	/A 🔽			
By completing this form, you confirm commissioning statement. Any requirements approximately provided the confirmation of the	ests which fall outside of this contact: norfolkicd@nhs.net	use will require an individua		•			
Please indicate whether patier		Please tick					
Erythropoiesis-stimulating agents (epoetin alfa, beta, theta and zeta, and darbepoetin alfa) are recommended, within their marketing authorisations, as options for treating anaemia in people with cancer who are having chemotherapy.							
2. If different erythropoiesis-stimulating agents are equally suitable, the product with the lowest acquisition cost for the course of treatment should be used.							
3. As per NICE guidance, treatment should normally be started with the least expensive drug (considering the dose required, price per dose and any additional administration costs). Please confirm that the choice of drug is considered the most cost-effective treatment for this individual patient, noting that alternative treatment options recommended by NICE may be lower in cost, including the use of biosimilar medications if applicable.							
Where an alternative lower cost treatment is available and has not been tried by the patient, please confirm that the clinical rationale for prescribing has been included in the patients' medical record for the purpose of auditing.							
4. The product is being used as described by local commissioning position.							
I confirm that the patient meets	s the criteria for treatment						
Name of person completing:		Contact Details:					
Designation of person completing		Date:					

Trust Authorising Pharmacist

Name:			
Date:			