Date:

N&WICS - TA684 - Nivolumab - Adjuvant treatment of completely resected melanoma with lymph node involvement or metastatic disease								
Before providing patient identifiable data on this form, please confirm that the patient (or in the case of a minor or vulnerable adult with the parent/legal guardian/carer) has given appropriate explicit consent for sensitive personal information on this form to be passed to the CCG and/or CSU for processing this funding request and validating subsequent invoices. Consent given:								
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, leas	ole. This has taken into cor st expensive, will be chose	nsideration therapeut	tic need and wheth	er or not the			
	I preference is stated in the	1A5	l					
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 🔽				
commissioning statement. Any required For support regarding IFRs, please For support regarding the criteria list	contact: norfolkicd@nhs.net	·	al funding request (II	FR).				
Please indicate whether patier	nt meets the following NICE	criteria:		Please tick				
1. Nivolumab is recommended, within its marketing authorisation, as an option for the adjuvant treatment of completely resected melanoma in adults with lymph node involvement or metastatic disease. It is recommended only if the company provides nivolumab according to the commercial arrangement.								
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.								
3. The product is being used as described by local commissioning position.				□Yes □No				
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		<u> </u>						
Name:								