Ollok liere to decess the guidelines	MIOL digorithm				
N&WICS - TA693 - Olaparib plus	bevacizumab - Maintenan	ce treatment of advanced cancer	l ovarian, fallopia	an tube or primar	y peritoneal
Before providing patient identifiable of parent/legal guardian/carer) has give CSU for processing this funding requirements.	en appropriate explicit consent	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, leas	ole. This has taken into cons st expensive, will be chosen	sideration therapeu	itic need and wheth	ner 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:	Compliant —	e □NHS care □Hospital	Sub-Type:	I/A 🔽	
For support regarding IFRs, please  For support regarding the criteria list	contact: norfolkicd@nhs.net	·	i runding request (i	ifk).	
Please indicate whether patier	nt meets the following NICE	criteria:		Please tick	
Olaparib plus bevacizumab is r maintenance treatment of advance and 4) high-grade epithelial ovaria	ed (International Federation o	of Gynecology and Obstetrics	s [FIGO] stages 3		
there has been a complete or partial response after first-line platinum-based chemotherapy plus bevacizumab, and				□Yes □No	
the cancer is associated with ho conditions in the managed access			nded only if the		
As per NICE guidance, treatment the dose required, price per dose			ug (considering		
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.					
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:		