Click here to access the guidelines/NICE algorithm						
N&WICS - TA836 - Palbociclib with fulvestrant - Hormone receptor-positive, HER2-negative advanced breast cancer after endocrine therapy						
Before providing patient identifiable parent/legal guardian/carer) has giv CSU for processing this funding rec	en appropriate explicit conser	nt for sensitive personal infe	ormation on this form			
CSO for processing this funding rec	quest and validating subseque	nt invoices. Consent given	. 🗀			
If there is more than one NICE-appr about the advantages and disadvar patient is likely to adhere to treatme	tages of the treatments availa	able. This has taken into co	onsideration therapeu	itic need and wheth	ner or not the	
and price per dose) unless an orde	r of preference is stated in the	TAs.				
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.n	et account ONLY)	Contact name & number:			
Start date	Provider:	te NHS				
requested treatment:	Supplier: Home	ecare Hospital	Sub-Type:	I/A 💙		
commissioning statement. Any requirements for support regarding IFRs, please For support regarding the criteria list	contact: norfolkicd@nhs.net	·	ual funding request (l	FR).		
Please indicate whether patie	nt meets the following NIC	E criteria:		Please tick		
Palbociclib plus fulvestrant is recommended as an option for treating hormone receptor-positive, HER2-negative locally advanced or metastatic breast cancer in adults who have had endocrine therapy only if:						
exemestane plus everolimus is the most appropriate alternative to a cyclin-dependent kinase 4 and 6 (CDK4/6) inhibitor and			kinase 4 and 6	☐Yes ☐No		
the company provides it according to the commercial arrangement.						
2. If patients and their clinicians consider palbociclib plus fulvestrant and abemaciclib plus fulvestrant or ribociclib plus fulvestrant to be suitable options, use the least expensive treatment. Take account of the monitoring and adverse event costs, dosage, price per dose and commercial arrangements.				☐Yes ☐No		
As per NICE guidance, treatment the dose required, price per dose.			drug (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.				□Yes □No		
Where an alternative lower cost that the clinical rationale for presauditing.		, ,				
I confirm that the patient meet	ts the criteria for treatment					
Name of person completing:		Contact Details:				
Designation of person completing	g:	Date:				

Trust Authorising Pharmacist	
Name:	
Date:	