Me	mber Name: {{MEMFIK51}}	{{MEMILASI}} DOB:	{{MEMB	ERDOB}} PA NI	imber: {{PANUMBER}}		
$\{\{D\}$	ANUMCODE}} olderightsize ANUMCODE}} acdescription}}						
forr {{C	s fax machine is located in a sens to {{COMPANY_NAME}} LIENT_PAG_PHONE}} with will authorize the coverage of	at {{CLIENT_PAG_FAX} questions regarding the property of the p	(}}. Please	e contact {{COM	PANY_NAME}} at		
Pati Phy	ent's Name: {{MEMFIRST}} ent's ID: {{MEMBERID}} sician's Name: {{PHYFIRST} cialty: sician Office Telephone: {{PH	Patient } {{PHYLAST}} Patient	's Date of Phone: <	Birth: {{MEME < MEMPHONE>>	>		
Phy < <f< td=""><td>sician Office Address: <<phy PHYZIP>></phy </td><td></td><td></td><td></td><td></td></f<>	sician Office Address: < <phy PHYZIP>></phy 						
	g Name: {{DRUGNAME}}						
	intity:	Frequency: Expected		_Strength:			
	te of Administration:	Expected .	Length of	Therapy:			
Dia	gnosis: < <diagnosis>> IO</diagnosis>	D Code: < <icd9>></icd9>					
1.	. What is the prescribed medication? Xeloda capecitabine Other						
2.	What is the patient's diagnosi ☐ Breast cancer	s?	□ Panc	reatic adenocarcii	noma		
	☐ Esophageal and esophagog	astric junction cancer		ric cancer			
	☐ Squamous cell skin cancer	•	☐ Amp	ullary adenocarci	noma		
	☐ Fallopian tube cancer			ary peritoneal can	icer		
	☐ Mucinous carcinoma of the	ovary		le cancer			
	☐ Anal carcinoma	1 .		noma or thymic ca			
	Gestational trophoblastic n			I bowel adenocare oendocrine and ac			
	☐ Occult primary tumor (cancer) Colorectal cancer (including cancer)						
	☐ Biliary tract cancer (including extrahepatic and intrahepatic cholangiocarcinoma and gallbladder cancer) ☐ Ovarian cancer (including epithelial ovarian cancer, carcinosarcoma [malignant mixed Müllerian tumor], clear cell carcinoma, grade 1 endometrioid carcinoma, low-grade serous carcinoma/borderline epithelial tumor)						
	☐ Head and neck cancer (incl	luding very advanced head		,			
	☐ Vulvar cancer			ical cancer			
	☐ Endometrial carcinoma		☐ Otne	r			
3.	What is the ICD-10 code?		4 1	-1:4:9			
4.	Is this a request for continuation of therapy with the requested medication? ☐ Yes ☐ No If No, skip to diagnosis section.						
5.	Is there evidence of unacceptable toxicity or disease progression while on the current regimen? ☐ Yes ☐ No No further questions.						
Con	nplete the following section bo	used on the patient's diag	nosis, if a	pplicable.			
<u>Sec</u>	tion A: Breast Cancer Will the requested medication	be given in combination	with ixab	epilone (Ixempra)	? □ Yes □ No		
7.	What is the clinical setting in which the requested medication will be used? <i>List continues on next page</i> .						
	Indicate all that apply.	requested medi	- LUION WI		z on new puge.		
	☐ Recurrent disease☐ Metastatic disease	☐ Relapsed disease☐ Locally advanced dise	ease	☐ As initial ther☐ Advanced dis			

Me	moer Name: {{MEMFIR51}} {{MEMLA51}} DOB: {{MEMBERDOB}} PA Number: {{PANOMBER}}					
	 □ Recurrent unresectable disease □ Postoperative residual disease □ The patient had no response to preoperative systemic therapy □ Other 					
8.	Does the patient have brain metastases in breast cancer?					
9.	Which of the following applies to the patient's disease? □ Early-stage HER2-negative disease □ Triple negative disease (TNBC) □ HER2-negative disease □ Other					
10.	 Will the requested be given in any of the following regimens? ☐ In combination with a HER2-inhibitor (e.g. margetuximab-cmkb [Margenza], trastuzumab [Herceptin], lapatinib [Tykerb], neratinib [Nerlynx]) ☐ In combination with trastuzumab (Herceptin) and tucatinib (Tukysa) ☐ As a single agent ☐ In combination with docetaxel ☐ Other 					
11.	 Will the requested medication be used in any of the following settings? □ As adjuvant therapy □ As maintenance therapy following adjuvant chemotherapy □ The patient has brain metastases in breast cancer □ Other 					
12.	2. Will the requested medication be used as subsequent therapy? Yes No					
13.	Will the requested medication be given as a single agent? ☐ Yes ☐ No					
	Section B: Neuroendocrine and Adrenal Tumors [44. What is the origin for the disease? The patient has neuroendocrine tumors of gastrointestinal tract, lung, or thymus (carcinoid tumors) The patient has neuroendocrine and adrenal tumors of pancreas The patient has extrapulmonary poorly differentiated disease/large or small cell disease/mixed neuroendocrine-non-neuroendocrine neoplasm The patient has well differentiated grade 3 neuroendocrine tumors Other Other					
15.	 Will the requested medication be given in any of the following regimens? Indicate all that apply. □ In combination with temozolomide (Temodar) □ With concurrent or sequential radiation □ As a component of CAPEOX (capecitabine and oxaliplatin) regimen □ Other 					
	tion C: Ovarian Cancer, Fallopian Tube Cancer, Primary Peritoneal Cancer, Mucinous Carcinoma of the Ovary Please indicate which of the following applies to the patient's disease? Carcinosarcoma (malignant mixed Mullerian tumors) Epithelial ovarian cancer Grade 1 endometrioid carcinoma Low-grade serous carcinoma/borderline epithelial tumor Other Other					
17.	What is the clinical setting in which the requested medication will be used? <i>Indicate all that apply</i> . ☐ Persistent disease ☐ Recurrent disease ☐ Platinum-sensitive (e.g., carboplatin, cisplatin) recurrence ☐ Platinum-resistant (e.g., carboplatin, cisplatin) recurrence ☐ Other					
18.	Will the requested medication be given as a single agent? ☐ Yes ☐ No					

Member Name: {{MEMFIRST}} {{MEMLAST}} DOB: {{MEMBERDOB}} PA Number: {{PANUMBER}}				
19. Will the requested medication be given in combination with oxaliplatin as adjuvant treatment? ☐ Yes ☐ No				
20. Will the requested be given in any of the following regimens? ☐ As a single agent ☐ In combination with oxaliplatin ☐ Other				
Section D: Head and Neck Cancer, Penile Cancer, Gestational Trophoblastic Neoplasia 21. Will the requested medication be given as a single agent? Yes No				
Section E: Occult Primary Tumor 22. Will the requested medication be given in any of the following regimens? ☐ As a single agent ☐ As a component of CAPEOX (capecitabine and oxaliplatin) regimen ☐ Other				
Section F: Anal Carcinoma 23. Will the requested medication be given in any of the following regimens? ☐ The requested medication with concurrent chemoradiation and in combination with mitomycin ☐ The requested medication with radiation as a single agent ☐ Other				
24. Is the requested medication being used after primary treatment of metastatic disease? ☐ Yes ☐ No				
Section G: Thymoma or Thymic Carcinoma 25. Will the requested medication be given in combination with gemcitabine? □ Yes □ No				
Section H: Squamous Cell Skin Cancer 26. What is the clinical setting in which the requested medication will be used? Indicate all that apply. Regional disease Distant Metastatic disease Uccally advanced disease Other				
27. Is the regional disease unresectable, inoperable, or incompletely resected? <i>Indicate all that apply</i> . ☐ Yes, unresectable disease ☐ Yes, inoperable disease ☐ Yes, incompletely resected disease				
28. Is the patient ineligible for immune checkpoint inhibitors and clinical trials? If Yes, skip to #31 □ Yes □ No				
29. Has the patient's disease progressed on immune checkpoint inhibitors and clinical trials? Yes No				
30. Will the requested medication be given as a single agent? ☐ Yes ☐ No				
Section I: Cervical Cancer, Vulvar Cancer 31. Will the requested medication be used as concurrent chemoradiation in combination with mitomycin? ☐ Yes ☐ No				
32. Are cisplatin and carboplatin available? ☐ Yes ☐ No				
33. Is cisplatin available? □ Yes □ No				
Section J: Endometrial Carcinoma 34. Will the requested medication be used as primary treatment as concurrent chemoradiation in combination with mitomycin? □ Yes □ No				
35. Are cisplatin and carboplatin available? ☐ Yes ☐ No				
I attest that the medication requested is medically necessary for this patient. I further attest that the information provided is accurate and true, and that the documentation supporting this information is available for review if requested by the claims processor, the health plan sponsor, or, if applicable a state or federal regulatory agency.				
Prescriber (Or Authorized) Signature and Date				