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This fax machine is located in a secure location as required by HIPAA regulations. Fax complete signed and dated forms to CVS/Caremark at . Please contact CVS/Caremark at 1-888-413-2723 with questions regarding the prior authorization process. When conditions are met, we will authorize the coverage of the medication.

Patient Name: Patient ID: Patient Group No: Physician Office Address: Drug Name (specify drug) Quantity: Route of Administration: Diagnosis:		Frequency:	Expected Length of Therap	ength:
Cor				
Plea		te answer for each applica Lupron Depot being prescri ate	•	
	Yes - Lupron Depot			
	No			
2.	Ovulation induction (e	fallopian transfer [GIFT], zy m injection [ICSI])		
3.	What is the intent of the Inhibition of premature	rapy? e luteinizing hormone (LH) s	surge	
	Trigger of oocyte maturation and ovulation			
	Other, please specify			
4.	Leuprolide acetate or Luinstructions.	upron Depot does not meet	guidelines. See comments for	
	Review more infertility	y medications		
	Infertility review comp	olete		
	the drug name field. F Clinician. Open one n	Pend initial PA to clinician by	es, remove the approved drug(s) from the selecting Team Infertility Pend to next did meet guidelines, attach any ions.	
5.	Leuprolide acetate will b	be approved for 12 months.		
	Review more infertility	y medications		
	Infertility review comp	lete		

6.	s cetrorelix acetate, Cetrotide, Fyremadel and/or Ganirelix being prescribed for this nember?			N	
7					
7.	What is the type of procedure the patient will be undergoing? Ovulation induction (e.g., intrauterine insemination [IUI])		П		
	Assisted reproductive technology (e.g., in vitro fertilization [IVF], frozen embryo transfer, gamete intrafallopian transfer [GIFT], zygote intrafallopian transfer [ZIFT], intracytoplasmic sperm injection [ICSI])		Ц		
	Other, please specify.				
8.	What is the intent of therapy?				
	Inhibition of premature luteinizing hormone (LH) surge				
	Other, please specify.				
9.	Cetrorelix acetate, Cetrotide, Fyremadel, and/or Ganirelix will be approved for 12 months. Review more infertility medications.				
	Infertility review complete				
10.	Cetrorelix acetate, Cetrotide, Fyremadel, and/or Ganirelix does not meet guidelines. See comments for instructions.				
	Review more infertility medications.				
	Infertility review complete				
	If one or more medications did not meet guidelines, remove the approved drug(s) from the drug name field. Pend initial PA to clinician by selecting Team Infertility Pend to Clinician. Open one new PA for all medications that did meet guidelines, attach any relevant documents, approve and load authorizations.				
11.	Are hCG, Novarel, Ovidrel and/or Pregnyl being prescribed for this member?	Y		N	
12.	What is the patient's diagnosis or the type of procedure the patient will be undergoing?				
	Ovulation induction (e.g., intrauterine insemination [IUI])				
	Assisted reproductive technology (e.g., in vitro fertilization [IVF], frozen embryo transfer, gamete intrafallopian transfer [GIFT], zygote intrafallopian transfer [ZIFT], Intracytoplasmic sperm injection [ICSI])				
	Prepubertal cryptorchidism				
	Hypogonadotropic hypogonadism		П		
	Other, please specify.		_		
	Cutof, ploado opcomy.				
13.	Does the patient have a low pretreatment testosterone level? ACTION REQUIRED: If Yes, attach laboratory results of testosterone level.	Y		N	
14.	Does the patient have low or low to normal levels of follicle stimulating hormone (FSH) or luteinizing hormone (LH)? ACTION REQUIRED: Attach laboratory results of FSH or LH levels.				
	Yes - Follicle stimulating hormone (FSH) level				
	Yes - Luteinizing hormone (LH) level				
	No				
15.	Novarel, Pregnyl, hCG and/or Ovidrel will be approved for 6 months.				
	Review more infertility medications				
	Infertility review complete				

16. Novarel, Pregnyl, hCG and/or Ovidrel will be approved for 12 months.

	Review more infertility medications				
	Infertility review complete				
17.	Novarel, Pregnyl, hCG, and Ovidrel do not meet guidelines. See comments for instructions.				
	Review more infertility medications				
	Infertility review complete				
	If one or more medications did not meet guidelines, remove the approved drug(s) from the drug name field. Pend initial PA to clinician by selecting Team Infertility Pend to Clinician. Open one new PA for all medications that did meet guidelines, attach any relevant documents, approve and load authorizations.				
18.	Is Follistim AQ and/or Gonal-f being prescribed for this member?	Y		N	
19.	What is the patient's diagnosis or the type of procedure the patient will be undergoing?				
	Ovulation induction (e.g., intrauterine insemination [IUI])				
	Assisted reproductive technology (e.g., in vitro fertilization [IVF], frozen embryo transfer, gamete intrafallopian transfer [GIFT], zygote intrafallopian transfer [ZIFT], Intracytoplasmic sperm injection [ICSI])				
	Hypogonadotropic hypogonadism				
	Other, please specify.				
20.	What is the patient's age?				
	Less than 37 years of age				
	Greater than or equal to 37 years of age				
21.	How many cycles of clomiphene citrate has the patient completed? Less than 3 cycles				
	Greater than or equal to 3 cycles				
22.	Does the patient have a risk factor for poor ovarian response to clomiphene?	Υ		N	
23.	Does the patient have a contraindication or exclusion to therapy with clomiphene?	Y		N	
24.	Does the patient have a low pretreatment testosterone level? ACTION REQUIRED: If Yes, attach laboratory results of testosterone level.	Υ		N	
25.	Does the patient have low or low to normal levels of follicle stimulating hormone (FSH) or luteinizing hormone (LH)? ACTION REQUIRED: Attach laboratory results of FSH or LH levels.				
	Yes - follicle stimulating hormone (FSH) level				
	Yes - luteinizing hormone (LH) level				
	No				
26.	Gonal-F and/or Follistim AQ do not meet guidelines. See comments for instructions. Review more infertility medications		П		
	Infertility review complete				
	If one or more medications did not meet guidelines, remove the approved drug(s) from the drug name field. Pend initial PA to clinician by selecting Team Infertility Pend to Clinician. Open one new PA for all medications that did meet guidelines, attach any relevant documents, approve and load authorizations.		Ш		
27.	Gonal-F and/or Follistim AQ will be approved for 12 months.				
	Review more infertility medications				

ı				
	Infertility review complete			
28.	Is Menopur being prescribed for this member?	Y	N	
29.	What is the type of procedure the patient will be undergoing?			
	Ovulation induction (e.g., intrauterine insemination [IUI])			
	Assisted reproductive technology (e.g., in vitro fertilization [IVF], frozen embryo transfer, gamete intrafallopian transfer [GIFT], zygote intrafallopian transfer [ZIFT], intracytoplasmic sperm injection [ICSI])			
	Other, please specify.			
30.	What is the patient's age?			
	Less than 37 years of age			
	Greater than or equal to 37 years of age			
31.	How many cycles of clomiphene citrate has the patient completed?			
	Less than 3 cycles			
	Greater than or equal to 3 cycles			
32.	Does the patient have a risk factor for poor ovarian response to clomiphene?	Y	N	
33.	Does the patient have a contraindication or exclusion to therapy with clomiphene?	Υ	N	
34.	Menopur does not meet guidelines. See comments for instructions.			
	Infertility review complete			
	Infertility review complete			
	If one or more medications did not meet guidelines, remove the approved drug(s) from the drug name field. Pend initial PA to clinician by selecting Team Infertility Pend to Clinician. Open one new PA for all medications that did meet guidelines, attach any relevant documents, approve and load authorizations.			
35.	Menopur will be approved for 12 months.			
	Infertility review complete			
	Infertility review complete			
36.	Have any infertility medications been approved? If yes, load authorization for the indicated period of time for the approved drugs. If one or more medications did not meet guidelines, remove the approved drug(s) from the drug name field. Pend initial PA to clinician by selecting Team Infertility Pend to Clinician. Open one new PA for all medications that did meet guidelines, attach any relevant documents, approve and load authorizations.	Y	N	

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

Now you can get responses to drug PAs immediately and securely online—without faxes, phone calls, or waiting. How? With electronic prior authorization (ePA)! For more information and to register, go to www.caremark.com/epa.