US Family Health Plan Prior Authorization Request Form for tasimelteon (Hetlioz, Hetlioz LQ)

To be completed and signed by the prescriber. To be used only for prescriptions which are to be filled through the Department of Defense (DoD) US Family Health Plan Pharmacy Program. US Family Health Plan is a TRICARE contractor for DoD.

The completed form may be faxed to 855-273-5735

OR

The patient may attach the completed form to the prescription and **mail** it to:

Attn: Pharmacy, 77 Warren St, Brighton, MA 02135

QUESTIONS? Call 1-877-880-7007

Initial and renewal prior authorization expires after 6 months. For renewal of therapy an initial USFHP prior authorization approval is required.

		4						
Step	ıΡI	ease complete patient and physician information (plo	ease print):					
.1	Pa	tient Name: Phys	ician Name:					
	Ad	dress:	Address:					
			_					
	Sponsor ID#		Phone #: _					
	Da	te of Birth: Se	ecure Fax #:					
Step	Please complete the clinical assessment:							
2	1. The provider acknowledges that Hetlioz capsules are not approved for pediatrics or adolescents and are not approved for treating Smith-Magenis Syndrome (SMS); and that Hetlioz LQ liquid is only approved for pediatrics with SMS and is not approved for Non-24 sleep wake disorder or for use in adults.		☐ Acknow ledged Proceed to question 2					
	2.	Has the patient received this medication under the	□ Y	es	□ No			
		USFHP benefit in the last 6 months? Please choose "No" if the patient did not previously have a USFHP	(subject to ve	erification)	Proceed to question 4			
		approved PA for Hetlioz.	Proceed to q	uestion 3				
	3. Has the patient been receiving Hetlioz/Hetlioz LQ for 6 months and has a documented response to therapy?		Y	es	□ No			
			Sign and date below		STOP			
					Coverage not approved			
	4. What is the requested medication?		☐ Hetlid	oz LQ	☐ Hetlioz capsules			
			Proceed to q	uestion 5	Proceed to question 6			
		Is the patient between 3 years of age and 15 years of	□ Y	es	□ No			
	age?		Proceed to q	uestion 7	STOP Coverage not approved			
	6. Is the patient greater than or equal to 18 years of ag		□ Y	es	□ No			
			Proceed to q	uestion 8	STOP Coverage not approved			

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7.	Smith-Magenis Syndrome (SMS)?	□ Yes □ No		
	Silitii-mageriis Syridi oille (SMS) !	Proceed to question 10	STOP	
	Note: Non-FDA-approved uses are not approved including jet lag disorder or other circadian rhythm disorders.		Coverage not approved	
8.	Is the patient totally blind?	☐ Yes	□ No	
		Proceed to question 9	STOP Coverage not approved	
9.	Does the patient have a documented diagnosis of non-	□ Yes	□ No	
	24 hour sleep-wake disorder?	Proceed to question 10	STOP Coverage not approved	
	Note: Non-FDA-approved uses are not approved including jet lag disorder or other circadian rhythm disorders.			
10.	.Has the patient had a trial of melatonin and either	☐ Yes	□ No	
	failed therapy or had an adverse event to therapy?	Proceed to question 11	STOP Coverage not approved	
11.	Has the patient tried and failed ramelteon?	☐ Yes	□ No	
		Proceed to question 12	STOP Coverage not approved	
12.	Is the patient taking a drug that will interact with	☐ Yes	□ No	
	Hetlioz, for example, beta blockers or strong CYP3A4 inducers? Examples of strong CYP3A4 inducers: Banzel (rufinamide), dexamethasone, Fycompa (perampanel), griseofulvin, Intelence (etravirine), modafinil (Provigil), Mycobutin (rifabutin), nafcillin, Onfi (clobazam), oxcarbazepine (Oxtellar XR, Trileptal), phenobarbital, phenytoin (Dilantin), Priftin (rifapentine), primidone (Mysoline), rifampin (Rifadin), St. John's wort, Sustiva (efavirenz), Tegretol (carbamazepine), Tracleer (bosentan), Viramune (nevirapine), Xtandi (enzalutamide), Zelboraf (vemurafenib). Examples of beta blockers: atenolol (Tenormin), betaxolol (Kerlone), bisoprolol (Zebeta), metoprolol (Lopressor, Toprol XL), nadolol (Corgard), nebivolol (Bystolic), propranolol (Inderal), sotalol (Betapace), timolol.	STOP Coverage not approved	Sign and date below	
l ce	ertify the above is true to the best of my knowle	dge. Please sign and d	ate:	
	Prescriber Signature	 Date		
	i resolibei Signature	Date	[20 April 2022	