Spravato (Esketamine) continued request

Instructions: Please **thoroughly complete** all fields in the treatment request form. Missing information will delay processing as all requested clinical information is needed to determine if medical necessity is met for this treatment. "See attached" is not a sufficient response, as all information on form needs to be accurate as of the date signed by the provider.

Please submit this form through provider self-service at **HumanaMilitary.com** to ensure all necessary clinical information is included and to expedite the authorization process.

	Date Submitted.		
Beneficiary information			
Name:	_ DOB:	TRICARE ID:	
Address:			
City:			
Phone #:	_		
Rendering provider			
Provider name:	TIN/NPI:		
Address:			
City:			
Point of contact direct phone #:	Fax #:		
Provider enrolled/certified in Risk Evaluation and Mitigation Strategy (REMS):	☐ Yes ☐ No		
Who will be billing TRICARE for the drug (Spravato/ Esketamine nasal spray)?			
\square Rendering provider (buy and bill) or \square Specialty pharmacy			
If pharmacy, the following information is required:			
Pharmacy name:	TIN/NPI:		





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Current psychiatric and medical conditions Diagnosis (DSM-5/ICD-10)	Onset	Descripti	on (include symptoms, treatment, etc.)
Diagnosis (DSM-S/TCD-10)	Oliset	Descripti	on (molade symptoms, treatment, etc.)
Current medications			
Medication name	Dose	Duration	Efficacy
Spravato will be used in conjunction with an o	ral antidepressant: [] Yes □ No	
Please detail any medication changes made si	nce initial Spravato tr	eatment:	
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-			
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Have there been any suicide attempts since initial Spravato treatment? Yes No If yes, please provide further details:					
Clinical progress summary supporting ongoing Spravato treatment:					
Evidence based rating scale outcomes over course of Spravato treatment:					
Assessment:	Score:	Date:			
Assessment:	Score:	Date:			
Assessment:	Score:	Date:			
Assessment:	Score:	Date:			
Assessment:	Score:	Date:			
Assessment:	Score:	Date:			
Assessment:	Score:	Date:			
Assessment:	Score:	Date:			
Beneficiary will be enrolled in REMS while receiving Spravato treatment: Yes No Beneficiary will be monitored for at least two hours following administration of Spravato (Esketamine) nasal spray by a qualified healthcare provider: Yes No					
There are no contraindications to the continuation of treatment with Spravato: \square Yes \square No					
Signature indicates that the beneficiary is physically and intellectually capable to actively participate in all aspects of the therapeutic program and information provided is true and accurate to the best of my knowledge.					
Provider signature:	Date				

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