Spravato® (esketamine) C III
28 mg nasal spray

An Established Therapy for Two Subsets of Challenging to Treat Major Depressive Disorder (MDD) in Adults

PRESENTED BY

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Consultant is a paid speaker for Janssen Pharmaceuticals, Inc.

Thursday, February 02, 2023 at 6:30 PM

LOCATION

Koestler Prime
1000 Highland Colony Parkway
Suite 6001, Ridgeland, MS (601) 957-3753

REGISTRATION INFORMATION

Please RSVP to Kimberly Morgan, (228) 547-3205, before Saturday, January 28, 2023, or visit http://www.medforcereg.net/SOMP/180091

To attend this program, I will need to attest that:

- I understand that Coronavirus/COVID-19 is very contagious, that COVID-19 is believed to spread mainly from person-to-person contact, and that it is not possible to eliminate any and all risks related to the potential spread of COVID-19
- I am participating in this event voluntarily and at my own risk
- I am not experiencing any symptom of illness such as cough, shortness of breath or difficulty breathing, fever, chills, repeated shaking with chills, muscle pain, headache, sore throat, or new loss of taste or smell.

I acknowledge that Janssen cannot guarantee that I will not become infected with COVID-19 (coronavirus) by virtue of attendance at this event and that Janssen has not made any representations or undertaken any obligations with respect to the venue of this event or the health of any staff or other participants. The number of attendees and the use of face coverings for this event will be determined based on local government guidance and a local assessment of risk.

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Please note that the company prohibits the offering of gifts, gratuities, or meals to federal government employees/officials. Thank you for your cooperation.

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INDICATIONS

SPRAVATO® (esketamine) CIII Nasal Spray is indicated, in conjunction with an oral antidepressant, for the treatment of:

- Treatment-resistant depression (TRD) in adults.
- Depressive symptoms in adults with major depressive disorder (MDD) with acute suicidal ideation or behavior.

LIMITATIONS OF USE:

- The effectiveness of SPRAVATO® in preventing suicide or in reducing suicidal ideation or behavior has not been demonstrated. Use of SPRAVATO® does not preclude the need for hospitalization if clinically warranted, even if patients experience improvement after an initial dose of SPRAVATO®.
- SPRAVATO® is not approved as an anesthetic agent. The safety and effectiveness of SPRAVATO® as an anesthetic agent have not been established.

IMPORTANT SAFETY INFORMATION

WARNING: SEDATION, DISSOCIATION; ABUSE AND MISUSE; and SUICIDAL THOUGHTS AND BEHAVIORS

See full prescribing information for complete boxed warning

- **Risk for sedation and dissociation after administration. Monitor patients for at least two hours after administration (5.1, 5.2).**
- **Potential for abuse and misuse. Consider the risks and benefits of using SPRAVATO® prior to use in patients at higher risk of abuse. Monitor for signs and symptoms of abuse and misuse (5.3).**
- **SPRAVATO® is only available through a restricted program called the SPRAVATO® REMS (5.4).**
- **Increased risk of suicidal thoughts and behaviors in pediatric and young adult patients taking antidepressants. Closely monitor all antidepressant-treated patients for clinical worsening and emergence of suicidal thoughts and behaviors. SPRAVATO® is not approved for use in pediatric patients (5.5).**

Please see additional Important Safety Information continued on next page.

Please see accompanying full Prescribing Information, including Boxed WARNINGS, and Medication Guide for SPRAVATO®.