

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.

Wednesday, February 08, 2023 at 6:30 PM

LOCATION

Field's Steak & Oyster Bar 119 Rue Magnolia, Biloxi, MS (228) 207-4137

REGISTRATION INFORMATION

Please RSVP to Kimberly Morgan, (228) 547-3205, before Friday, February 03, 2023, or visit http://www.medforcereg.net/SOMP/180095

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 of 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.

In adherence with PhRMA guidelines, spouses or other guests are not permitted to attend company-sponsored programs.

For all attendees, please be advised that information such as your name and the value and purpose of any educational item, meal or other items of value you receive may be publicly disclosed. If you are licensed in any state or other jurisdiction, or are an employee or contractor of any organization or governmental entity, that limits or prohibits meals from pharmaceutical companies, please identify yourself so that you (and we) are able to comply with such requirements.

Please note that the company prohibits the offering of gifts, gratuities, or meals to federal government employees/officials. Thank you for your cooperation. This promotional educational activity is not accredited. The program content is developed by Janssen Pharmaceuticals, Inc. Speakers present on behalf of the company and are required to present information in compliance with FDA requirements for communications about its medicines.

The personal information you provide may be used to contact you about your request to attend the Janssen Pharmaceuticals, Inc., educational program using your preferred method of communication as indicated by you. This information may be shared with Janssen Pharmaceuticals, Inc., its affiliates, and a third party for the sole purpose of completing your registration for this program and as required by law.

INDICATIONS

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

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

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®.