Poster 26

Real-World Experience With Treatment and Expectations by Satisfaction Status of Persons With Major Depressive Disorder

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Key Objective

To examine the relationship between self-reported treatment satisfaction and treatment response, experiences, and expectations from the perspective of people with major depressive disorder (MDD)

Introduction

- MDD is a highly prevalent and often persistent psychiatric disorder characterized by sadness or loss of interest in
- It is often accompanied by changes in weight, sleep habits, and energy levels, as well as a decreased quality of life^{3,4} Individuals with MDD additionally have an increased risk of suicidal ideation^{4,5}
- People who take medications for MDD exhibit varying levels of satisfaction and differing expectations regarding
- Despite many approved treatments, patients often have trouble achieving desired outcomes due to delayed or inadequate therapeutic effects and side effects⁴⁻⁶
- Individuals' treatment expectations and experiences can differ across satisfaction levels with MDD treatment⁶⁻⁸

Methods

- A survey was designed in collaboration with mental health experts and patients through the Depression and Bipolar Support Alliance (DBSA) to assess MDD treatment expectations and experiences
- The survey was conducted online from December 2021 to January 2022 among adults with self-reported MDD in the United States

Outcomes

- Sociodemographic and clinical characteristics, including co-occurring non-psychiatric and psychiatric conditions
- Current MDD treatment, including line of therapy and time on treatment
- MDD severity and current treatment experiences, including confidence in their current treatment, belief in their treatment being the best available, functioning, and side effects, as well as treatment expectations and goals
- MDD severity was assessed using the validated 16-item Quick Inventory of Depression Symptomology (QIDS-SR-16) and then was categorized based on total scores: none/mild (0-10), moderate (11-15), or severe/very severe (16-27)9

- Among respondents reporting current MDD treatment, 3 satisfaction groups were constructed: dissatisfied (somewhat, very, or extremely dissatisfied), neither (neither satisfied nor dissatisfied), and satisfied (somewhat, very, or extremely satisfied)
- Descriptive analyses were conducted with means and standard deviations (SDs) for continuous variables and counts and percentages for categorical variables
- Between-group differences on sociodemographic and clinical characteristics, co-occurring medical conditions, and current MDD treatment were examined via regression analysis with a significance level set at 0.05
- Treatment experience outcomes were analyzed using logistic regression adjusting for age, gender, race (White vs. non-White), education (bachelor's degree or greater vs. less than bachelor's degree), insurance (private vs. nonprivate), arthritis, diabetes, obesity, chronic fatigue syndrome, autoimmune disorder, hypertension, chronic pain syndrome, migraine, anxiety disorder, and obsessive-compulsive disorder; predicted values between treatment satisfaction groups were tested with a significance level set at 0.05, using a two-tailed approach and adjusting for multiple comparisons

Acknowledgments

- Pan Z, et al. CNS Spectr. 2019;24(1):22-9.
- Tolentino JC, Schmidt SL. Front Psychiatry. 2018 (9):450 4. Karrouri R, et al. World J Clin Cases. 2021;9(31):9350-67. 5. McIntyre RS, et al. World Psychiatry. 2023;22(3):394-412.
- Baune BT, et al. Neuropsychiatr Dis Treat. 2021;17:2995-3006. Demyttenaere K, et al. Prim Care Companion CNS Disord. 2011;13(4): 8. IsHak WW, et al. Int Neuropsychiatr Dis J. 2016;7(2):INDJ.26203. 9. Rush AJ, et al. Biol Psychiatry. 2003;54(5):573-83.

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Respondent Characteristics

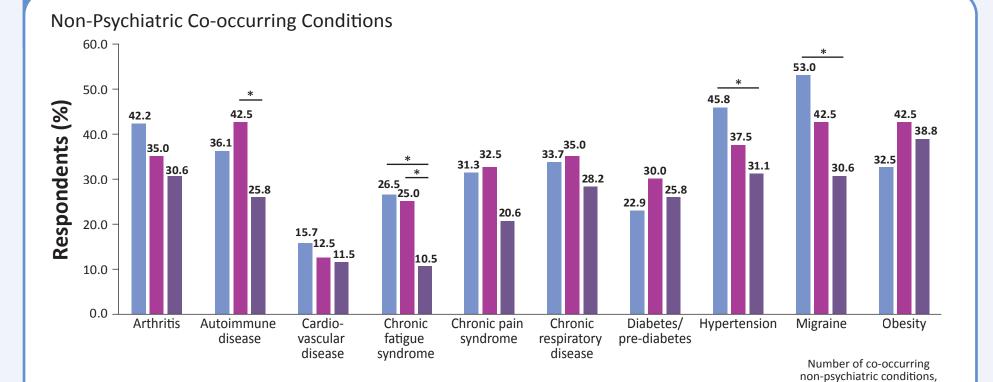
Results

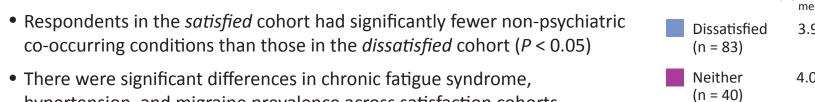
- A total of 332 individuals currently receiving pharmacologic MDD treatment completed the survey
- Self-reported satisfaction levels varied, with 62.9% of respondents being satisfied, 25.0% being dissatisfied, and 12.1% being neither satisfied nor dissatisfied with their current MDD treatment
- There were significant between-group differences on race, education, insurance type, and duration of current MDD-related treatment (all P < 0.05)

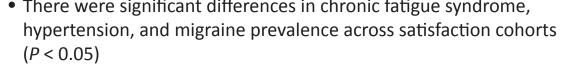
	Dissatisfied (n = 83)	Neither (n = 40)	Satisfied (n = 209)
lge, years; nean (SD)	46.1 (11.9)	46.7 (10.5)	47.6 (13.2)
Gender, n (%)			
Female	66 (79.5)	33 (82.5)	168 (80.4)
Race, n (%)*			
White	62 (74.7)	27 (67.5)	179 (85.6)
ducation, n (%)*			
Less than a bachelor's degree	53 (64.6)	24 (60.0)	100 (47.8)
Bachelor's degree or greater	29 (35.4)	16 (40.0)	109 (52.2)
nsurance type, ı (%)*			
Private	31 (39.7)	15 (39.5)	117 (57.6)
Not private ^a	47 (60.3)	23 (60.5)	86 (42.4)
ime on current reatment, n (%)*			
< 5 years	50 (60.2)	19 (47.5)	90 (43.1)
5-10 years	24 (28.9)	16 (40.0)	69 (33.0)
> 10 years	9 (10.8)	5 (12.5)	50 (23.9)
ine of therapy,			
First	17 (20.5)	9 (22.5)	43 (20.6)
Second	22 (26.5)	12 (30.0)	47 (22.5)
Third	11 (13.3)	6 (15.0)	34 (16.3)
Fourth or more	27 (32.5)	12 (30.0)	71 (34.0)
Don't know/not sure	6 (7.2)	1 (2.5)	14 (6.7)

Note: Group comparison P values were derived from a linear regression model for age; a multinomial logistic regression model for gender, race, and insurance type, a logistic regression model for education; and a proportional odds logistic regression model for time on current treatment and line of therapy ^a Includes Medicaid, Medicare, Veterans Affairs, and other

Co-occurring Conditions







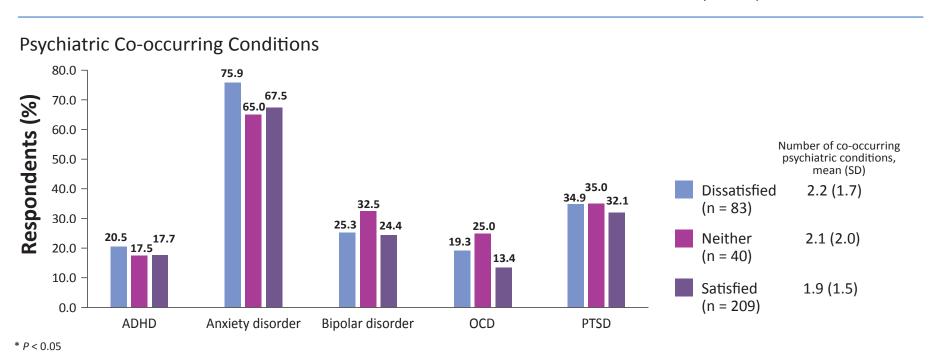
Current MDD-related Treatments

• Across cohorts, 54.2% of respondents were receiving monotherapy.

Es = esketamine; SNRI = serotonin-norepinephrine reuptake inhibitor; SSRI = selective serotonin reuptake inhibitor

Note: Group comparison P values were derived from a logistic regression model for all MDD-related treatments

SNRIs showed the largest difference between dissatisfied and satisfied groups



Satisfied

(n = 209)

3.1 (2.5)

ADHD = attention deficit hyperactivity disorder; OCD = obsessive-compulsive disorder; PTSD = post-traumatic stress disorder Notes: Other non-psychiatric comorbidities included atopic dermatitis, cancer, endometriosis, epilepsy, osteoporosis, and polycystic ovary syndrome. Other psychiatric comorbidities included eating disorder, personality disorder, postpartum depression, schizophrenia, schizoaffective disorder, and substance use disorder. Group comparison P values were derived from a logistic regression model for specific co-occurring non-psychiatric and psychiatric conditions and from a negative binomial regression model for the total number of co-occurring non-

• Common treatments included SSRIs (dissatisfied, 50.6%; neither, 42.5%; satisfied, 44.5%), SNRIs (dissatisfied, 41.0%;

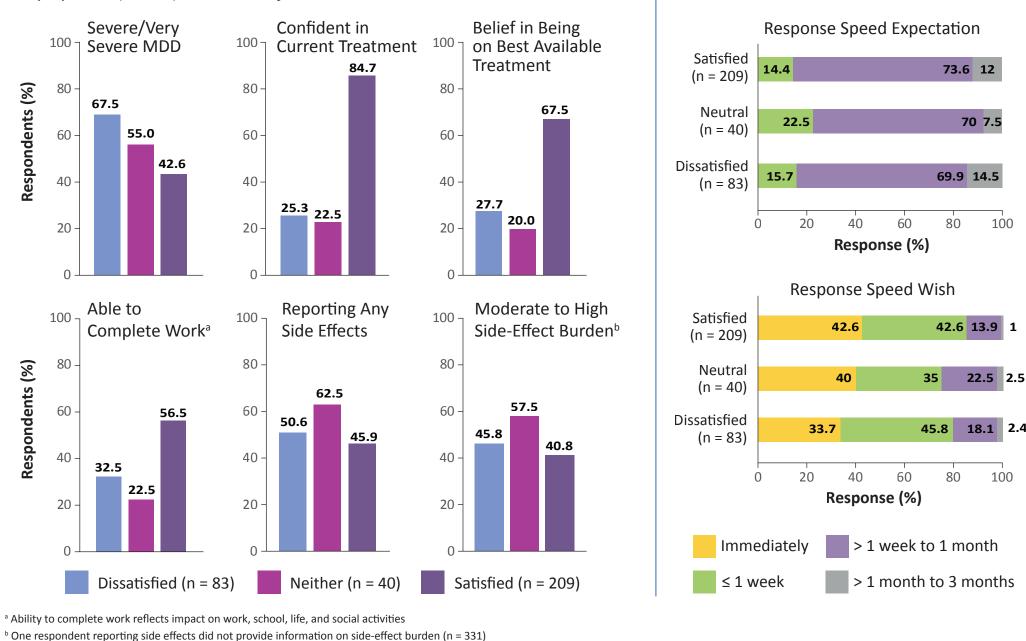
neither, 32.5%; satisfied, 27.8%), and antipsychotics (dissatisfied, 31.3%; neither, 30.0%; satisfied, 21.5%)

Es/ketamine Modified SSRI^a

Dissatisfied (n = 83) Neither (n = 40) Satisfied (n = 209)

Treatment Experience, Expectations, and Goals

- Better treatment experiences were reported by respondents in the *satisfied* cohort
- Regardless of satisfaction group, many respondents reported high side-effect burden
- Side effects were reported by 50.6%, 62.5%, and 45.9% of respondents in the dissatisfied, neither, and satisfied cohorts, respectively, with 45.8%, 57.5%, and 40.8% reporting moderate to high side-effect burden
- The most impactful side effects were cognitive impairment (26.2%), weight gain (21.4%), and sexual dysfunction (19.0%) for the dissatisfied cohort; cognitive impairment and weight gain (both 20.0%) and breakthrough symptoms and emotional blunting (both 16.0%) for the *neither* cohort; and weight gain (25.0%), sexual dysfunction (18.8%), and breakthrough symptoms (16.7%) for the satisfied cohort



Across satisfaction groups, top-reported Results from a survey of 332 individuals treatment goals included improved currently receiving MDD treatment functionality (77.7%); feelings of anxiety, agitation, and restlessness revealed that 25% of respondents were (75.9%); feelings of sadness, emptiness, dissatisfied with their current treatment

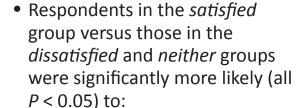
or hopelessness (74.4%); loss of interest

- or pleasure (67.5%); and tiredness or Persons with MDD—including those who lack of energy (67.2%) report overall treatment satisfaction— Among all respondents, 82.6% indicated continued to experience unmet treatment
- they desired a treatment response needs, particularly regarding symptom within 1 week control, ability to function, treatment confidence, and side effects
 - Weight gain was one of the most impactful side effects for both the satisfied and dissatisfied groups

Conclusions

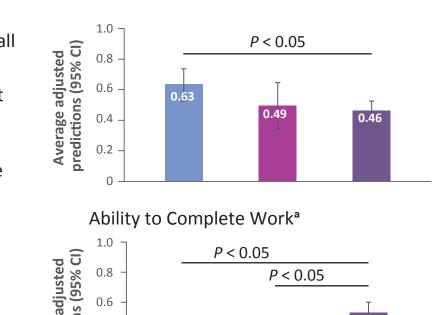
- Respondents dissatisfied with their treatment were more likely to have more severe MDD symptoms and less likely to
- Confidence in their treatment
- Belief in receiving the best available treatment
- Ability to complete work and daily activities
- Most respondents—including those satisfied with their MDD treatment indicated they desired a treatment response time of immediately or within 1 week (> 80%), although expectations for such a rapid response were low (16%), reflecting the time to response for common treatments

Regression Analysis by MDD Treatment Satisfaction

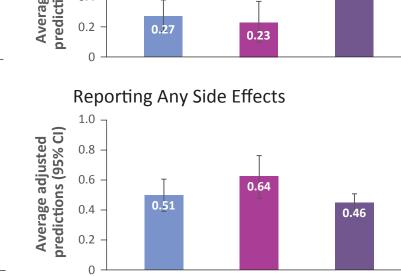


 Be confident in their current MDD medication

- Believe they were receiving the best treatment available
- Be able to complete their work or other normal daily activity
- Satisfied respondents were significantly less likely to have more severe MDD than dissatisfied respondents (P < 0.05)

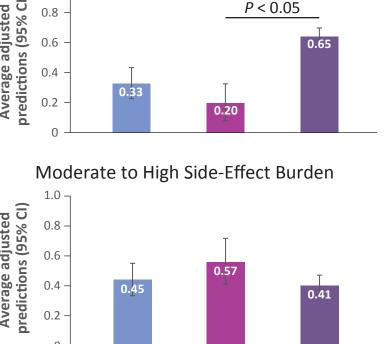


Severe/Very Severe MDD



Confidence in Current MDD Medication

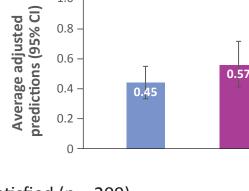
P < 0.05



Belief in Being on Best Available Treatment

P < 0.05

CI = confidence interval ^a Ability to complete work reflects impact on work, school, life, and social activities



Dissatisfied (n = 83) Neither (n = 40) Satisfied (n = 209)