

# Burden of Prominent Anhedonia in Major Depressive Disorder Reflected in Polypharmacy, Healthcare Use, and Humanistic Outcomes

Hrishikesh Kale,<sup>1</sup> Michael L. Ganz,<sup>2</sup> Rajrupa Ghosh,<sup>3</sup> Cynthia Saiontz-Martinez,<sup>3</sup> Tiina Drissen<sup>1</sup>, Andrew J. Cutler<sup>4</sup>

<sup>1</sup>Janssen Scientific Affairs, LLC, a Johnson & Johnson Company, Titusville, NJ, USA; <sup>2</sup>Formerly Evidera, Inc.; <sup>3</sup>Evidera, Inc., Waltham, MA, USA; <sup>4</sup>SUNY Upstate Medical University, Lakewood Ranch, FL, USA

## Background

- Major depressive disorder (MDD) is a serious mental health condition that imposes vast economic, medical, and personal burdens.<sup>1-5</sup>
- About 21 million adults in the US had at least one depressive episode in 2020; the 12-month prevalence was approximately 9.2%.<sup>6,7</sup>
- Anhedonia, a key symptom and diagnostic criterion of MDD, is characterized by deficits in pleasure and/or interest<sup>8</sup> and has been linked to worse prognosis, lower rate of remission, and higher functional impairments and suicidality in individuals with MDD.<sup>9-12</sup>
- Although 40%–70% of individuals with MDD exhibit symptoms of anhedonia,<sup>13,14</sup> little is known about the clinical and humanistic burden associated with prominent anhedonia in individuals with MDD.

## Objectives

Assess the clinical and humanistic burden associated with prominent anhedonia in patients with MDD.

## Methods

### Study Design and Data Source

- Pooled cross-sectional study using data from the Medical Expenditure Panel Survey (MEPS) (2016 through 2019).

### Study Cohorts and Measures

- Respondents with MDD were identified as those with at least one record in the MEPS medical conditions file containing an International Classification of Diseases, 10th Revision, Clinical Modification (ICD-10-CM) code of F32\* (depressive episode) or F33\* (major depressive disorder, recurrent).
- The presence and degree of anhedonia were assessed using the first item of the Patient Health Questionnaire-2 (PHQ-2), administered during the second and fourth interview rounds of each MEPS panel ("During the past two weeks, bothered by having little interest or pleasure in doing things").
- We classified respondents with MDD into two groups, based on their MDD and anhedonia status:
  - Respondents with MDD who reported more than half the days (2) or nearly every day (3) were classified as having MDD with prominent anhedonia (MDD-ANH).
  - Respondents with MDD who reported not at all (0) or several days (1) were classified as having MDD with no/low anhedonia (other-MDD).

### Sample Selection

#### Included

- Respondents who were at least 18 years of age during the first interview round of each year.

#### Excluded

- Respondents with bipolar disorder, dementia, Alzheimer's disease, and other neurological conditions during the calendar year.

### Use of Healthcare Services

- Use of pharmacologic treatments (medications) was measured by the proportion of respondents who used specific medications or medication classes and by the number of unique medications respondents used.
- Medications were classified as psychotropic or non-psychotropic; psychotropic medications were further classified into MDD or non-MDD (other) treatment.
- Polypharmacy was defined as the concomitant use of two or more psychotropic medications during the same calendar year.
- The numbers of visits and hospitalizations (per 100 persons) were assessed for office-based/outpatient visits, emergency department (ED) visits, inpatient hospitalizations, and home health services.
- These measures were also defined for mental health and non-mental health-related services.

- The Physical Component Summary (PCS) score.
- The Mental Component Summary (MCS) score.
- The PCS and MCS scores range from 0 to 100, with 0 representing the worst and 100 representing the best health status; 50 represents the national average.

- HRQoL was assessed using the 12-item Short Form Health Survey Version 2 (SF-12v2) instrument.
- The Physical Component Summary (PCS) score.
- The Mental Component Summary (MCS) score.
- The PCS and MCS scores range from 0 to 100, with 0 representing the worst and 100 representing the best health status; 50 represents the national average.

- Insurance coverage during the year, %
  - Any private coverage: 69.5
  - Only public coverage: 26.5
  - Uninsured: 3.5
- Charlson comorbidity index, mean: 0.8

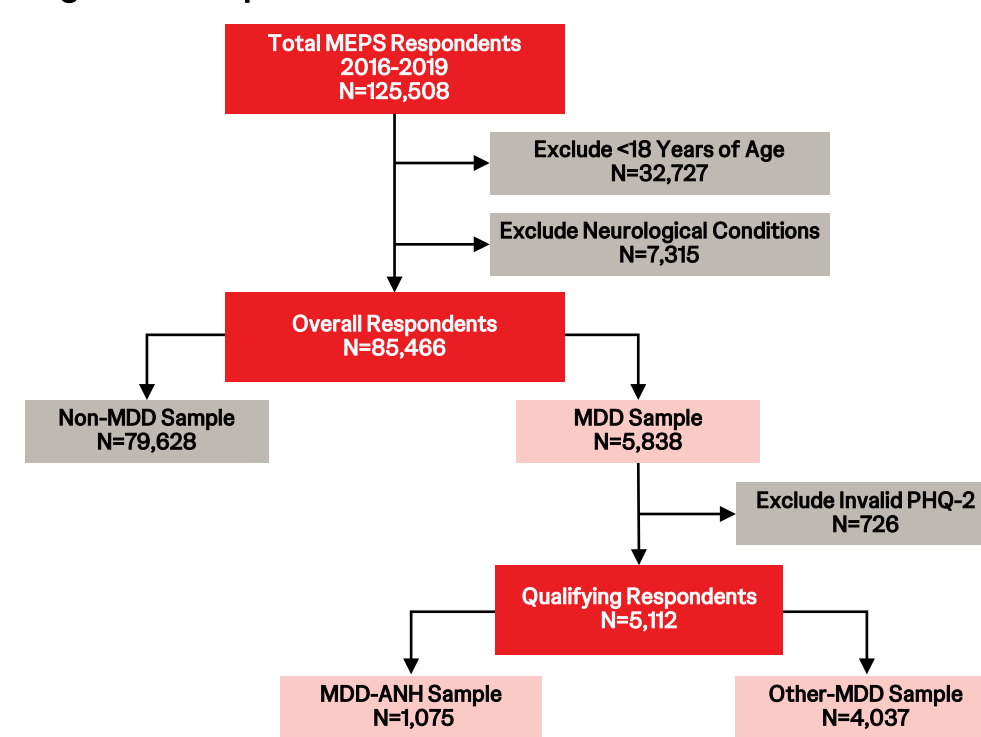
- HRQoL was assessed using the 12-item Short Form Health Survey Version 2 (SF-12v2) instrument.
- The Physical Component Summary (PCS) score.
- The Mental Component Summary (MCS) score.
- The PCS and MCS scores range from 0 to 100, with 0 representing the worst and 100 representing the best health status; 50 represents the national average.

- Weighting accounted for the non-MDD group as well; findings presented in this poster focus on respondents with MDD-ANH and other-MDD, only.

### Statistical Analyses

- The MDD-ANH and other-MDD groups were compared using propensity score-based inverse probability weighting to minimize the effect of confounding and selection bias on outcomes.
- All outcomes were summarized by their weighted means (continuous variables) or weighted proportions (categorical variables) and their corresponding 95% confidence intervals (CIs).
- Weighting accounted for the non-MDD group as well; findings presented in this poster focus on respondents with MDD-ANH and other-MDD, only.

Figure 1. Sample Selection Flowchart



## Results

### Respondent Characteristics (Tables 1–2; Figure 1)

- We identified 5,838 (7.2%) respondents with MDD, of whom 5,112 respondents had a valid response for presence and degree of anhedonia in the first item of PHQ-2.
- 1,075 individuals with MDD had MDD-ANH.
- 4,037 individuals with MDD had other-MDD.
- Selected respondents were, on average (mean), 50 years of age, 70% were female, >87% were White, and about 66% were covered by private health insurance plans.
- Respondents with MDD-ANH were mostly Black (9.0% vs. 5.9%), previously married (36.6% vs. 28.5%), had low to negative income (44.8% vs. 27.9%), and had more comorbid conditions as measured by the Charlson comorbidity index (0.8 vs. 0.6) than those in the other-MDD group.

Table 2. Selected Respondent Characteristics (Unweighted)

Characteristic	MDD-ANH (N=1,075)	Other-MDD (N=4,037)
Age in years, mean	50.6	49.5
Sex, %		
Female	68.1	70.5
Male	31.9	29.5
Race, %		
White only	84.7	87.8
Black only	9.0	5.9
Other race	6.2	6.2
Marital status, %		
Never married	25.4	25.7
Married	38.0	45.9
Divorced, separated, widowed	36.6	28.5
Income level, %		
Low to negative (<200% FPL)	44.8	27.9
Middle (200% to <400% FPL)	27.4	29.8
High (≥400% FPL)	27.8	42.3
Insurance coverage during the year, %		
Any private coverage	53.1	69.5
Only public coverage	43.5	26.5
Uninsured	3.5	4.0
Charlson comorbidity index, mean	0.8	0.6

### Use of Healthcare Services (Table 3; Figures 2–3)

- Respondents with MDD-ANH had more healthcare visits than those with other-MDD:
  - Office/outpatient visits: 1,305.2 per 100 persons vs. 896.9 per 100 persons
  - ED visits: 33.1 per 100 persons vs. 22.6 per 100 persons
  - Inpatient hospitalizations: 14.3 per 100 persons vs. 9.9 per 100 persons
  - Home health visits: 38.1 per 100 persons vs. 18.9 per 100 persons

### Use of Antidepressants and Psychotropic Medications (Table 4; Figure 4)

- Proportionately more respondents with MDD-ANH used serotonin modulators, tricyclic antidepressants, bupropion + serotonin modulators, and bupropion + selective serotonin reuptake inhibitors (SSRIs) than respondents with other-MDD.
- Respondents with MDD-ANH were also more likely to have used the following medications than those with other-MDD:
  - Any psychotropic medications: 75.7% vs. 71.7%
  - Non-MDD psychotropic medications: 42.8% vs. 25.3%
  - Any non-psychotropic medications: 75.0% vs. 72.2%
- Polypharmacy was slightly more prevalent among respondents with MDD-ANH than other-MDD (43.2% vs. 27.8%); respondents with MDD-ANH were about twice as likely as those with other-MDD to have used medications from other classes in addition to antidepressants:
  - Mood stabilizers: 13.0% vs 7.1%
  - Antipsychotics: 6.2% vs 2.9%
  - Anxiolytics: 6.5% vs 3.2%
  - Attention-deficit hyperactivity disorder (ADHD) medications: 7.1% vs 4.3%

### HRQoL (Figure 5)

- Mental HRQoL, as measured by the SF-12 MCS, was substantially lower for respondents with MDD-ANH (30.9) than for those with other-MDD (47.1).
- Physical HRQoL, as measured by the SF-12 PCS, was also lower for respondents with MDD-ANH (46.2) than for those with other-MDD (50.3).

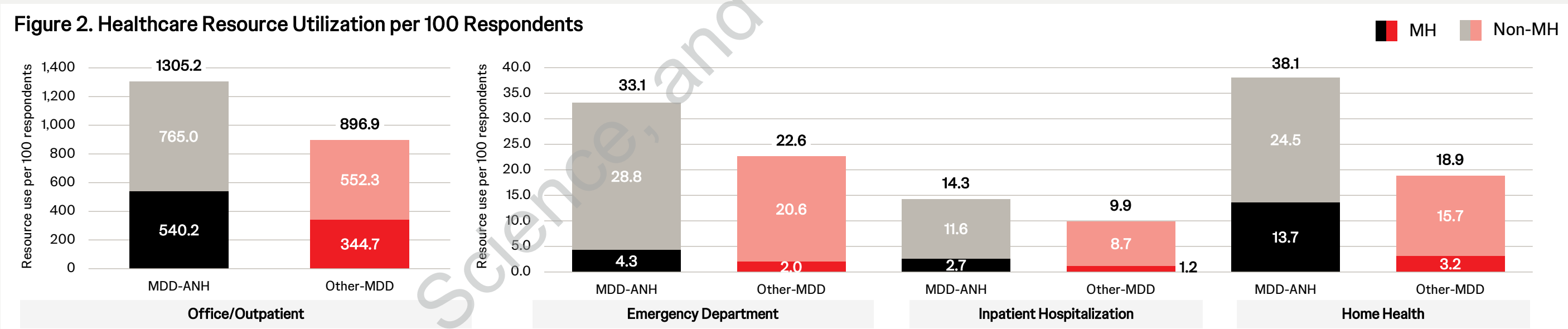


Table 3. Use of Healthcare Services

Use Measure	MDD-ANH	Other-MDD
All medication types, any use, % (95% CI)	90.2 (86.4, 93.9)*	85.7 (83.7, 87.8)*
Psychotropic, any use, % (95% CI)	75.7 (70.7, 80.6)*	71.0 (68.4, 73.6)*
MDD treatment, %	67.8 (62.6, 73.1)	66.0 (63.2, 68.7)
NDRI	16.2 (11.6, 20.8)	12.7 (11.0, 14.4)
Serotonin modulator	7.0 (3.8, 10.2)*	2.7 (2.0, 3.3)*
SNRI	14.0 (10.2, 17.9)	11.2 (9.7, 12.6)
SSRI	47.8 (42.3, 53.3)	46.5 (43.6, 49.3)
Tetracyclic antidepressant	2.1 (0.7, 3.6)	1.3 (0.8, 1.8)
Tricyclic antidepressant	3.6 (1.5, 5.7)*	1.6 (1.2, 2.0)*
Bupropion + serotonin modulator	2.5 (0.03, 4.9)*	0.5 (0.2, 0.8)*
Bupropion + SSRI	1.9 (0.6, 3.2)	1.2 (0.7, 1.7)
Bupropion + SSRI	9.1 (5.5, 12.8)*	4.4 (3.2, 5.5)*
Other (non-MDD) treatment, %	42.8 (37.3, 48.3)*	25.3 (23.3, 27.4)*
No. of unique psychotropic medications, mean (95% CI)	6.8 (6.5, 7.1)*	6.3 (6.2, 6.5)*
Non-psychotropic medication, any use, % (95% CI)	75.0 (69.6, 80.5)*	72.2 (69.7, 74.7)*

Figure 3. Rate Ratios Comparing Healthcare Resource Utilization between Study Cohorts

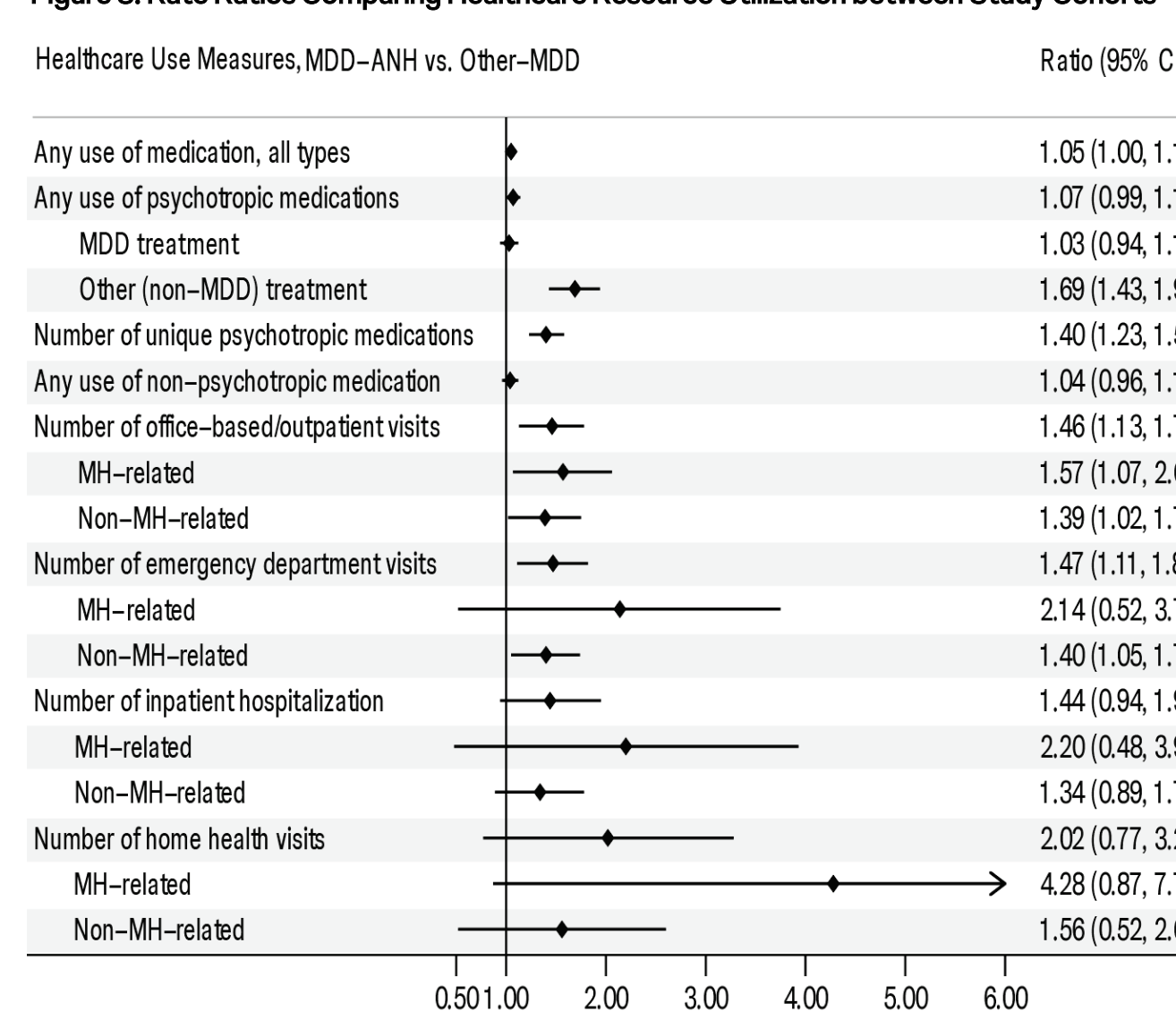
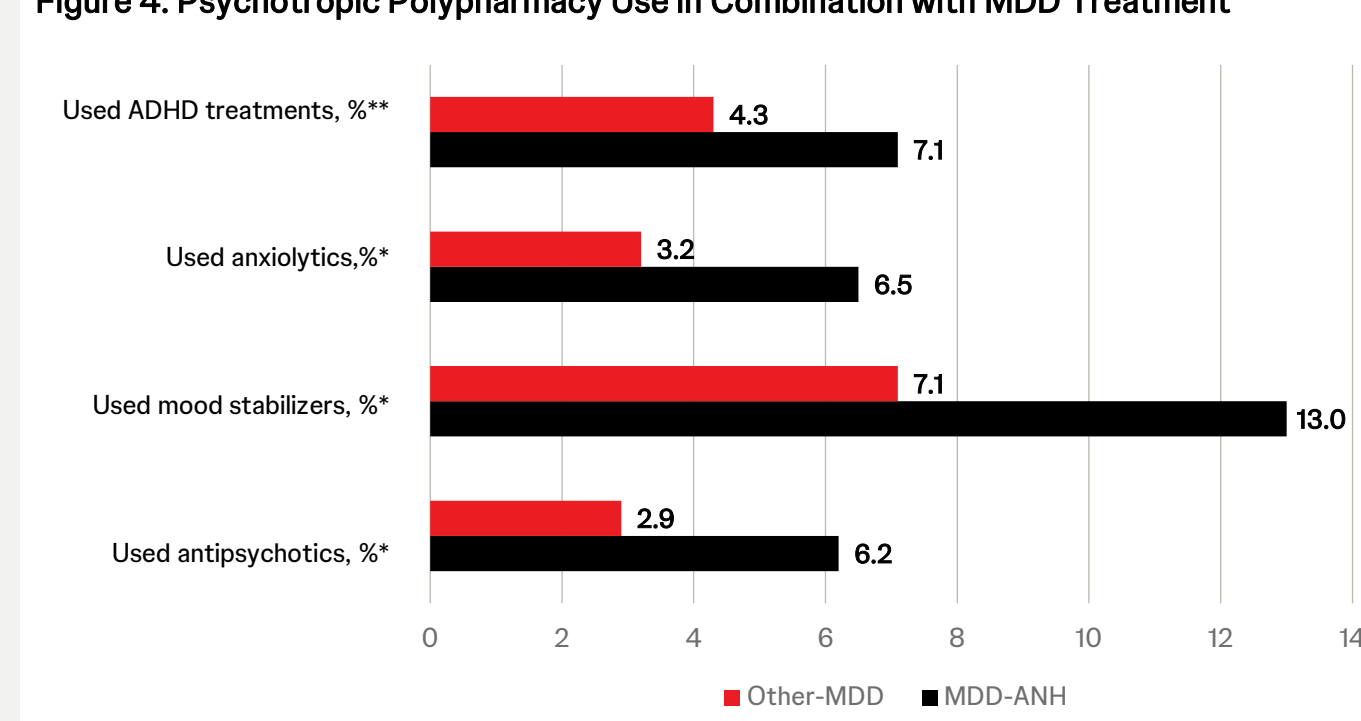


Table 4. Prevalence of Psychotropic Polypharmacy Use between Study Cohorts

Polypharmacy Use	MDD-ANH	Other-MDD
Polypharmacy, any use, % (95% CI)	43.2 (37.9, 48.5)	27.8 (25.6, 30.0)
Within same class, % (95% CI)		
Used 2 unique medications	17.2 (13.1, 21.3)*	11.3 (9.8, 12.8)*
Used 3 unique medications	4.3 (2.1, 6.5)*	0.9 (0.6, 1.2)*
Used ≥4 unique medications	0.9 (0.0, 1.8)*	0.03 (0.0, 0.08)*
Across different classes, % (95% CI)		
Used 2 unique classes	26.3 (21.4, 31.1)*	17.1 (15.4, 18.8)*
Used 3 unique classes	7.7 (4.9, 10.4)*	3.2 (2.6, 3.9)*
Used ≥4 unique classes	2.4 (1.1, 3.7)*	0.9 (0.4, 1.3)*

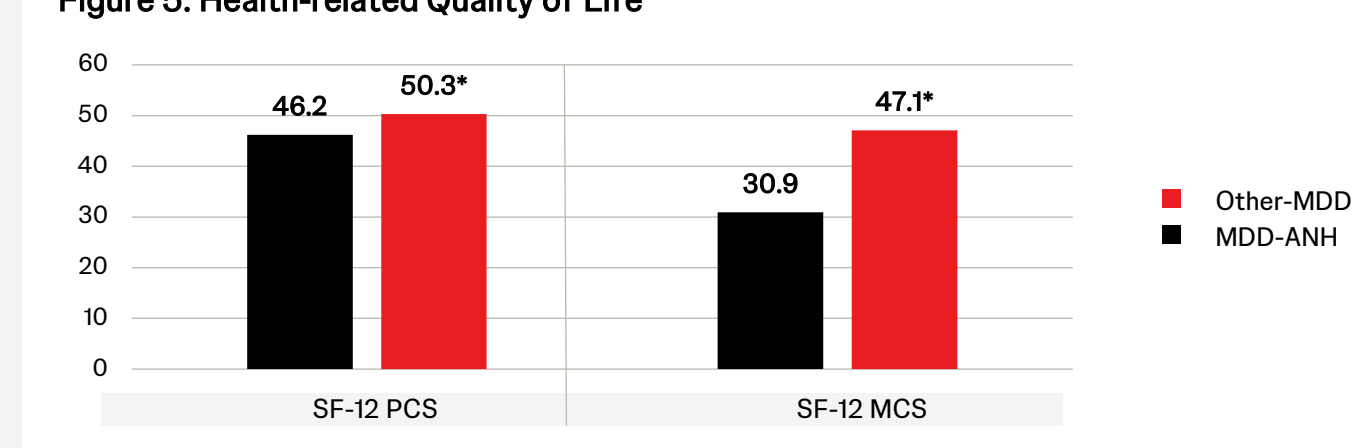
Note: Costs expressed in 2022 US dollars. \* Statistically significant differences between MDD-ANH vs other-MDD, p<0.05. Abbreviations: MDD-ANH = MDD with prominent anhedonia; other-MDD = MDD with no/low anhedonia

Figure 4. Psychotropic Polypharmacy Use in Combination with MDD Treatment



\* Statistically significant differences between MDD-ANH and other-MDD, p<0.01. \*\* Statistically significant differences between MDD-ANH and other-MDD, p<0.05. Abbreviations: ADHD = attention-deficit/hyperactivity disorder; MDD-ANH = MDD with prominent anhedonia; other-MDD = MDD with no/low anhedonia

Figure 5. Health-related Quality of Life



\* Statistically significant differences between MDD-ANH vs other-MDD, p<0.05. Abbreviations: MCS = mental component summary; MDD-ANH = MDD with prominent anhedonia; other-MDD = MDD with no/low anhedonia; PCS = physical component summary; SF-12 = 12-item Short Form Health Survey

## Conclusions

- Prominent anhedonia in MDD was associated with high psychotropic medication use, polypharmacy use, and healthcare resource use, as well as poor HRQoL.
- Higher psychotropic medication and polypharmacy use, along with poorer HRQoL, suggest that it is difficult to treat MDD – specifically patients with prominent anhedonia.
- Higher polypharmacy use in combination with MDD-related treatments indicates higher clinical burden and unmet needs among individuals with MDD with prominent anhedonia, compared to individuals without prominent anhedonia.
- The impact of MDD with prominent anhedonia extends beyond mental health, as reflected by higher use of non-MDD and non-mental health-related medications and healthcare services.
- These findings suggest that anhedonia severity should be considered in routine assessment of MDD; identifying targeted treatments may reduce clinical and humanistic burden associated with MDD.

## REFERENCES

- Greenberg PE, et al. *J Clin Psychiatry*. 2015;76(2):155-62.
- Greenberg PE, et al. *Pharmacoeconomics*. 2021;39(6):653-665.
- Kan K, et al. *Pharmacoeconomics*. 2021;39(6):721-730.
- Kessler RC. *Psychiatr Clin North Am*. 2012;35(1):1-14.
- Zhdanova M, et al. *J Clin Psychiatry*. 2021; 82(92).
- NIH. Major Depression. Accessed May 1, 2023. <https://www.nimh.nih.gov/health/statistics/major-depression>.
- Goodwin RD, et al. *Am J Prev Med*. 2022;63(5):726-733.
- Treadway MT, Zald DH. *Neurosci Biobehav Rev*. 2011;35(3):537-555.
- McMakin DL, et al. *J Am Acad Child Adolesc Psychiatry*. Apr 2012;51(4):404-11.
- Pelizza L, Ferrari A. *Ann Gen Psychiatry*. 2009;8:22.
- Spijker J, et al. *Acta Psychiatr Scand*. Feb 2001;103(2):122-30.
- Uher R, et al. *Depress Anxiety*. 2012;29(12):1043-9.
- Ritsner MS. *Anhedonia: A comprehensive handbook Volume I*. Springer. 2014;1:19-54.
- Cao B, et al. *Frontiers in Psychiatry*. 2019;10:17.

## Disclosures

This research was funded by Janssen Scientific Affairs, LLC. **Hrishikesh Kale** and **Tiina Drissen** are employees of Janssen Scientific Affairs, LLC. **Michael L. Ganz**, **Rajrupa Ghosh**, and **Cynthia Saiontz-Martinez** are employees of Evidera L.P.P., Inc., a contract research organization with previous and ongoing relationships with Janssen Scientific Affairs, LLC. **Andrew J. Cutler** has served in a consulting role for AbbVie, Acadia, Alfasigma, Alkermes, Axsome, Biogen, BioXcel, Boehringer Ingelheim, Bri Biosciences, Cerevel, Corium, Delpor, Evolution Research Group, Idorsia, Intra-Cellular Therapies, Ironshore Pharmaceuticals, Janssen, Jazz Pharmaceuticals, Karuna, Lundbeck, LivaNova, Luye Pharma, MapLight Therapeutics, MedAvante-ProPhase, Neumora, Neurocrine, Neuroscience Education Institute, NeuroSigma, Noven, Otsuka, Remeda, Sage Therapeutics, Sumitomo (Sunovion), Supernus, Takeda, Teva, Tris Pharma, VistaGen Therapeutics, and VivoSense; has served in a speaking role/received promotional honoraria from AbbVie, Acadia, Alfasigma, Alkermes, Axsome, BioXcel, Corium, Idorsia, Intra-Cellular Therapies, Ironshore Pharmaceuticals, Janssen, Karuna, Lundbeck, Neurocrine, Noven, Otsuka, Sumitomo (Sunovion), Supernus, Takeda, Teva, Tris Pharma, and Vanda Pharmaceuticals; and served on the Data Safety Monitoring Board for COMPASS Pathways, and Freedom Biosciences.

## Corresponding Author

Hrishikesh Kale (hkale@its.jnj.com)

## Key Contributors

Study conception and design (HK, MLG, RG, TD, AC); data analysis (CSM); data interpretation (HK, MLG, RG); draft poster preparation (RG); final approval (HK, TD, AC)

## Acknowledgements

Jason P. Swindle (Evidera, Inc.) contributed to drafting and reviewing the poster content

## Novel Pathways in Depression



Scan the QR code

