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## Antidepressants Are Not Overprescribed for Mild Depression

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### ABSTRACT

**Objective:** To evaluate overprescribing of antidepressant medication for minimal or mild depression.

**Method:** Electronic records data from 4 large health care systems identified outpatients aged 18 years or older starting a new episode of antidepressant treatment in 2011 with an ICD-9 diagnosis of depressive disorder (296.2, 296.3, 311, or 300.4). Patient Health Questionnaire-9 (PHQ-9) depression severity scores at time of treatment initiation were used to examine the distribution of baseline severity and the association between baseline severity and patients' demographic and clinical characteristics.

**Results:** Of 19,751 adults beginning treatment in 2011, baseline PHQ-9 scores were available for 7,051. In those with a baseline score, 85% reported moderate or severe symptoms (PHQ-9 score of 10 or more), 12% reported mild symptoms (PHQ-9 score of 5 to 9), and 3% reported minimal symptoms (PHQ-9 score of less than 5). The proportion reporting minimal or mild symptoms when starting treatment increased with age, ranging from 11% in those under age 65 years to 26% in those aged 65 and older. The proportion with minimal or mild symptoms was also moderately higher among patients living in wealthier neighborhoods and those treated by psychiatrists. Nevertheless, across all subgroups defined by sex, race/ethnicity, prescriber specialty, and treatment history, the proportions with minimal or mild symptoms did not exceed 18%. Secondary analyses, including weighting and subgroup analyses, found no evidence that estimates of baseline severity were biased by missing PHQ-9 scores.

**Conclusions:** In these health systems, prescribing of antidepressant medication for minimal or mild depression is much less common than suggested by previous reports. Given that this practice may sometimes be clinically appropriate, our findings indicate that overprescribing of antidepressants for mild depression is not a significant public health concern.

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Use of antidepressants has increased dramatically over the past 20 years in the United States and in other higher-income countries.<sup>1–5</sup> Approximately 10% of US adults now fill one or more antidepressant prescriptions in any calendar year.<sup>3</sup> Antidepressants prescribed by primary care physicians account for the majority of this increase.<sup>2,3</sup>

Increasing rates of antidepressant treatment have raised concerns about overprescribing to patients with less severe depression.<sup>6,7</sup> Community surveys suggest that the rates of antidepressant use may now exceed the prevalence of depression, especially among older adults.<sup>8</sup> In the 2003 Collaborative Psychiatric Epidemiologic Surveys, 26% of recent antidepressant users did not meet diagnostic criteria for any lifetime psychiatric diagnosis according to a structured research interview.<sup>9</sup> In the 2010 National Survey on Drug Use and Health, only 44% of respondents taking antidepressants reported experiencing a major depressive episode during the past year.<sup>10</sup> These findings were interpreted as evidence for substantial diagnostic inflation<sup>11</sup> and attracted significant public attention.<sup>12,13</sup>

Cross-sectional community surveys, however, may not accurately assess indications for antidepressant treatment. While recall of depression severity is reasonably accurate over several weeks,<sup>14</sup> more remote episodes of depression are often underreported.<sup>15–18</sup> If current antidepressant users were asked about past depression, failure to recall prior episodes would overestimate the proportion without a clear history of significant depression.

Here, we use data from 4 large health care systems to examine severity of depression at initiation of antidepressant treatment. We take advantage of the increasing use of standard depression severity measures to examine how often outpatients starting antidepressant treatment reported only minimal or mild symptoms at the time of the initial prescription. These ratings should be less subject to error or bias than would previous studies<sup>9,10</sup> relying on long-term recall.

### METHOD

Data were drawn from the Mental Health Research Network (MHRN), a consortium of public-domain research centers affiliated with 11 large not-for-profit integrated health care systems. Each of these systems provides comprehensive care (including general medical and specialty mental health care) to a defined population of members or patients. Across these systems, electronic medical records, insurance claims, and other administrative data systems have been organized in a Virtual Data Warehouse to facilitate population-based research.<sup>20</sup> Protected health information remains at each member health system, but common data definitions and formats facilitate sharing of de-identified data for research. Institutional review boards and privacy boards at each health system approved all study procedures and granted waivers of consent for this research use of de-identified records data.

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The 4 MHRN health care systems contributing data to this study include Group Health Cooperative, HealthPartners, Kaiser Permanente Colorado, and Kaiser Permanente Hawaii. These 4 systems serve a combined population of approximately 2 million members/patients in the states of Washington, Idaho, Minnesota, Colorado, and Hawaii. Members are enrolled through employer-sponsored insurance, individual insurance plans, and capitated Medicare and Medicaid programs and are generally representative of each system's regional population. In 2011, 10.2% of all adult members of these health care systems filled one or more antidepressant prescriptions, similar to national rates.<sup>3</sup> While Group Health and HealthPartners are mixed-model health care systems (providing care through both internal or group-model providers and external or network-model providers), this sample was limited to patients receiving prescriptions from internal providers to ensure availability of complete electronic medical records.

The study sample included all adult members filling a new outpatient antidepressant prescription from an internal health care system provider between January 1, 2011, and December 31, 2011. Eligible antidepressant medications included all drugs approved by the US Food and Drug Administration for treatment of major depression, excluding trazodone (more often prescribed for insomnia). A list of included medications and corresponding National Drug Codes is available at [www.mhresearchnetwork.org](http://www.mhresearchnetwork.org). A new episode of antidepressant treatment was defined by an interval of at least 270 days since the last filled antidepressant prescription. While this interval is longer than that used to define new prescriptions in National Committee for Quality Assurance (NCQA)/Healthcare Effectiveness Data Set (HEDIS) measures<sup>20</sup> and in some of our previous research,<sup>21,22</sup> it is based on 2 findings in records data from MHRN health systems. First, analyses of the frequency distributions of intervals between filled antidepressant prescriptions across health systems found that the rate of prescription fills remained elevated above the background rate for approximately 270 days. Second, review of full text medical records for visits between antidepressant fills found that 50% of visit notes reported continued medication use 180 days after the prior fill, with this rate dropping to 25% by 270 days. Details of both of these analyses are available as an online appendix (see eAppendix 1 at [PSYCHIATRIST.COM](http://PSYCHIATRIST.COM)). The study sample was limited to those with a recorded diagnosis of any depressive disorder (ICD-9 diagnosis 296.2, 296.3, 300.4, or 311) in the interval starting 90 days before the index prescription and ending 15 days after. In these health care systems, approximately 60% of adults receiving antidepressant treatment have a recorded diagnosis of depressive disorder, with most of the remainder having recorded diagnoses of anxiety disorders or attention-deficit disorders.<sup>23</sup> Patients with any diagnosis of bipolar disorder or psychotic disorder prior to the index prescription were excluded. To ensure availability of records data to assess inclusion and exclusion criteria, the sample was limited to those continuously enrolled in the participating health care

- Community antidepressant prescribing is usually consistent with guideline recommendations, with only a small proportion of patients who start treatment having minimal or mild symptoms.
- Psychiatrists may be more likely than primary care physicians to start or restart antidepressants when symptoms of depression are mild.

systems for at least 270 days prior to the index prescription. These criteria are illustrated in Figure 1.

During the study period, participating health care systems all recommended use of the Patient Health Questionnaire-9 (PHQ-9)<sup>24-26</sup> for initial assessment of depression severity and at all follow-up visits for depression care. Procedures for use of the PHQ-9 varied between health systems and between clinics within health systems. In general medical clinics, use of the PHQ-9 depended on practice teams' implementation of this recommended practice. Questionnaires could be administered prior to the visit by a nurse or medical assistant (if depression was identified in advance) or by the physician during the visit. Procedures also varied across mental health specialty clinics, with some clinics routinely administering the PHQ-9 prior to every outpatient visit and some relying on providers to administer it during visits as clinically indicated. Actual adherence to recommended use of the PHQ-9 was variable.

Electronic medical records and health care system administrative databases were used to identify the specialty of the prescribing physician and the following patient characteristics: sex, age at initiation of antidepressant treatment, race/ethnicity, and neighborhood income.

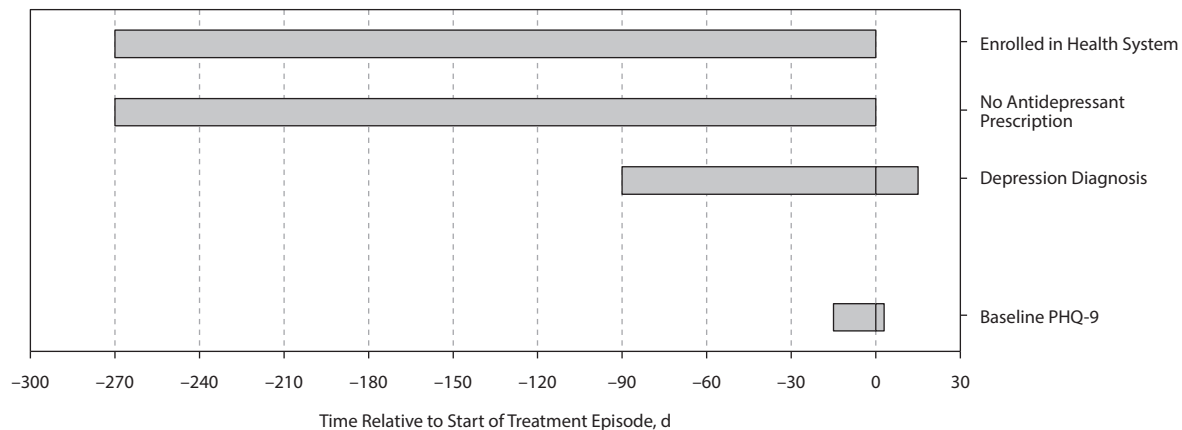
For these analyses, we defined a baseline PHQ-9 score as any measurement in the interval between 15 days prior to and 3 days after the index prescription. For patients with more than one PHQ-9 record in this interval, the PHQ-9 score recorded prior to and closest to the index prescription date was considered the baseline value.

Descriptive analyses examined the availability of baseline PHQ-9 scores and distributions of scores across health care systems and patient subgroups. Severity of depression according to PHQ-9 score was categorized as 0 to 4 (minimal depression), 5 to 9 (mild depression), 10 to 14 (moderate depression), 15 to 19 (moderately severe depression), and 20 or more (severe depression).<sup>26</sup> Predictors that a baseline PHQ-9 score would be recorded in the medical record and predictors that that baseline PHQ-9 score would be less than 10 were examined using logistic regression. To account for possible selection bias due to missing baseline PHQ-9 scores, weighted distributions of baseline scores were calculated using inverse probability weights<sup>27,28</sup> based on all covariate information listed in Table 1.

## RESULTS

Health care system records identified 19,751 patients aged 18 years and over with an eligible episode of antidepressant

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Figure 1. Criteria for Defining a New Antidepressant Treatment Episode<sup>a</sup>

<sup>a</sup>Criteria were as follows: at least 270 days of enrollment in the health system, at least 270 days since the most recent antidepressant prescription fill, and a recorded depression diagnosis in the interval from 90 days before to 15 days after the index prescription. The inclusion window for baseline Patient Health Questionnaire-9 (PHQ-9) severity measures extended from 15 days before to 3 days after the index prescription.

treatment in 2011. These treatments included 7,141 (36%) initial prescriptions for citalopram, 4,427 (22%) for fluoxetine, 2,838 (14%) for sertraline, and 2,361 (12%) for bupropion. No other single medication accounted for more than 5% of initial prescriptions. Seven thousand fifty-one (36%) episodes had a baseline PHQ-9 score recorded in the electronic medical record. As shown in Table 1, baseline PHQ-9 scores were recorded slightly more often among men than women. The proportion with a recorded baseline depression severity score declined progressively with age, decreasing from 44% in younger patients to only 28% in those aged 65 and older. Recording of a baseline severity score varied moderately across racial/ethnic groups (higher among African Americans and Native Americans, lower among Hispanics, Asians, and Native Hawaiians). Baseline scores were recorded much less often for those with no recorded race or ethnicity. Presence of a baseline PHQ-9 score was moderately higher among patients residing in economically disadvantaged neighborhoods and lower among patients treated by psychiatrists. The strongest predictor of having a recorded baseline severity score was the prescribing provider's number of patients starting antidepressant treatment for depression during the study year. The proportion with a baseline score increased from approximately 15% in those treating 5 or fewer patients to over 40% in those treating 11 or more patients.

Among those with a baseline PHQ-9 score, 5,988, or 85% (95% CI, 84%–86%), reported a score of 10 or more (indicating moderate or severe symptoms). Approximately 12% (95% CI, 12%–13%) had baseline scores between 5 and 9 (indicating mild symptoms of depression), and approximately 3% (95% CI, 2%–3%) had baseline scores less than 5 (indicating minimal symptoms of depression). As shown in Table 2, the distribution of baseline PHQ-9 scores did vary across the 4 health care systems more than

**Table 1. Proportion of Outpatients Starting Antidepressant Treatment With and Without a Recorded Patient Health Questionnaire-9 (PHQ-9) Depression Score at Baseline**

Variable	No Baseline PHQ-9 <sup>a</sup>	Yes Baseline PHQ-9 <sup>a</sup>	Odds Ratio <sup>b</sup> for Yes (95% CI)
Sex			
Female	8,555 (65)	4,693 (35)	Reference
Male	4,145 (64)	2,358 (36)	1.18 (1.10–1.26)
Age, y			
18–29	2,081 (56)	1,624 (44)	Reference
30–44	3,257 (62)	2,025 (38)	0.74 (0.67–0.81)
45–64	4,828 (66)	2,439 (38)	0.57 (0.52–0.62)
≥ 65	2,534 (72)	963 (28)	0.37 (0.34–0.42)
Race/ethnicity			
Non-Hispanic white	7,343 (59)	5,089 (41)	Reference
Asian	516 (63)	303 (37)	0.75 (0.64–0.87)
Non-Hispanic black	345 (51)	325 (49)	1.28 (1.09–1.51)
Hispanic	941 (65)	512 (35)	0.72 (0.64–0.82)
Native Hawaiian/Pacific Islander	135 (69)	60 (31)	0.52 (0.38–0.72)
Native American/Alaskan Native	47 (51)	45 (49)	1.35 (0.88–2.07)
Mixed race	421 (65)	230 (35)	0.65 (0.55–0.74)
Other or unknown	2,952 (86)	487 (14)	0.27 (0.25–0.31)
Neighborhood income			
≥ \$25,000	11,292 (65)	6,099 (35)	Reference
< \$25,000	1,408 (60)	952 (40)	1.36 (1.23–1.50)
Prescriber specialty			
Primary care or other	9,360 (62)	5,639 (38)	Reference
Psychiatry	3,340 (70)	1,412 (30)	0.48 (0.44–0.52)
Prescriber patient volume/y			
1–5	3,789 (85)	691 (15)	Reference
6–10	2,001 (66)	1,037 (34)	2.54 (2.26–2.84)
11–15	1,760 (54)	1,486 (46)	3.97 (3.56–4.44)
16–20	1,495 (54)	1,254 (46)	3.97 (3.54–4.46)
≥ 21	3,655 (59)	2,583 (41)	4.39 (3.95–4.87)

<sup>a</sup>Values shown as n (%).

<sup>b</sup>Odds ratios from logistic model including all covariates listed in this table.

expected by chance ( $\chi^2_{12} = 72.1$ ,  $P < .0001$ ), but the overall pattern was similar across all 4 systems. The proportion with minimal symptoms of depression ranged from 2% to 4%, and the proportion with mild symptoms of depression ranged from 11% to 21%.

Table 3 shows the proportion of patients with baseline PHQ-9 scores less than 10 for subgroups defined by



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sex, age, race/ethnicity, neighborhood income, and prescriber characteristics. The proportion with a low baseline severity score increased steadily with age—from approximately 11% among young adults to 26% among those aged 65 years or older. The proportion with minimal or mild symptoms was lower for all minority racial/ethnic groups compared to non-Hispanic whites, but odds ratios for individual racial/ethnic groups (compared to non-Hispanic whites) did not differ significantly from 1 after adjustment for other covariates. Low baseline PHQ-9 scores were less common among patients residing in economically disadvantaged neighborhoods and more common among those treated by psychiatrists. The proportion with baseline PHQ-9 score less than 10 did not differ according to the prescribing provider's number of patients treated during the study period.

Two secondary analyses examined the possibility that missing baseline PHQ-9 scores could bias estimates of baseline depression severity. First, the distribution of baseline PHQ-9 scores was recalculated after weighting each observation according to the inverse probability of PHQ-9 score availability<sup>27,28</sup> for each combination of predictors in Table 1. Second, analyses were limited to the subgroup patients treated by prescribers who recorded baseline PHQ-9 scores for at least 75% of patients starting antidepressant treatment in the study year. As shown in Table 4, the distribution of baseline depression severity scores using either of these methods was essentially identical to the unweighted results in the full sample.

## DISCUSSION

Our data do not support the prior claim<sup>10</sup> that the majority of patients treated with antidepressants have not experienced depression severe enough to warrant pharmacotherapy. In this sample, approximately 85% of adult outpatients starting antidepressants reported moderate or severe symptoms at the time of the initial prescription. This proportion was generally similar across health care systems.

Consistent with previous reports,<sup>8,10</sup> we did find that the proportion of antidepressant users reporting mild symptoms was greater in older patients: 26% of patients 65 and older compared to approximately 13% in those under age 65. Others have suggested that traditional symptoms scales or diagnostic assessments may underrepresent depression in older adults who may more often present with somatic symptoms or cognitive complaints.<sup>29</sup> Alternatively, older adults' more frequent contact with health care may result in a higher likelihood that less severe depression will be recognized and treated.

Lower baseline depression scores were more common in patients living in more economically advantaged neighborhoods. This pattern could reflect either a general tendency toward less severe depression in more advantaged patients or a tendency for more advantaged patients with mild depression to more often seek or receive treatment. Our data also suggest a higher threshold for prescribing of

**Table 2. Distribution of Baseline Patient Health Questionnaire-9 Depression Scores Among Outpatients Starting Antidepressant Treatment in 4 Health Care Systems**

Health Care System	PHQ-9 Score <sup>a</sup>				
	0–4	5–9	10–14	15–19	≥ 20
Group Health	102 (3)	434 (11)	1,139 (28)	1,354 (34)	987 (25)
HealthPartners	41 (4)	158 (15)	319 (30)	310 (30)	209 (20)
KP Colorado	29 (2)	222 (13)	517 (31)	540 (32)	385 (23)
KP Hawaii	12 (4)	65 (21)	62 (20)	98 (32)	68 (22)
Total	184 (3)	879 (12)	2,037 (29)	2,302 (33)	1,649 (23)

<sup>a</sup>Values shown as n (%).

Abbreviations: KP = Kaiser Permanente, PHQ-9 = Patient Health Questionnaire-9.

**Table 3. Proportion of Outpatients Starting Antidepressant Treatment With Baseline Patient Health Questionnaire-9 (PHQ-9) Depression Scores Above, At, or Below Threshold of 10**

Variable	Baseline PHQ-9 Score < 10 <sup>a</sup>	Baseline PHQ-9 Score ≥ 10 <sup>a</sup>	Odds Ratio <sup>b</sup> for Score < 10 (95% CI)
Sex			
Female	690 (15)	4,003 (85)	Reference
Male	373 (16)	1,985 (84)	1.03 (0.89–1.18)
Age, y			
18–29	179 (11)	1,445 (89)	Reference
30–44	274 (14)	1,751 (86)	1.27 (1.04–1.56)
45–64	360 (15)	2,079 (85)	1.39 (1.14–1.68)
≥ 65	250 (26)	713 (74)	2.76 (2.22–3.42)
Race/ethnicity			
Non-Hispanic white	804 (16)	4,825 (84)	Reference
Asian	41 (14)	262 (86)	0.86 (0.61–1.21)
African American	35 (11)	290 (89)	0.71 (0.50–1.02)
Hispanic	73 (14)	439 (86)	0.95 (0.74–1.24)
Hawaiian/Pacific Islander	4 (7)	56 (93)	0.44 (0.16–1.21)
Native American	3 (7)	42 (93)	0.39 (0.12–1.28)
More than one	33 (14)	197 (86)	1.01 (0.69–1.47)
Unknown	70 (14)	417 (86)	0.98 (0.75–1.28)
Neighborhood income			
≥ \$25,000	950 (16)	5,149 (84)	Reference
< \$25,000	113 (12)	839 (88)	0.81 (0.68–0.98)
Prescriber specialty			
Primary care or other	829 (15)	4,810 (85)	Reference
Psychiatry	234 (17)	1,178 (83)	1.16 (1.03–1.45)
Prescriber patient volume/y			
1–5	94 (14)	597 (86)	Reference
6–10	158 (15)	869 (85)	1.10 (0.84–1.46)
11–15	241 (16)	1,245 (84)	1.16 (0.90–1.51)
16–20	185 (15)	2,198 (85)	1.04 (0.79–1.36)
≥ 21	1,063 (15)	5,988 (85)	1.01 (0.79–1.30)

<sup>a</sup>Values shown as n (%).

<sup>b</sup>Odds ratios from logistic model including all covariates listed in this table.

antidepressants to members of racial/ethnic minority groups. This could reflect a bias in providers' decision processes or a difference in patients' treatment preferences. Previous research does suggest that African American and Hispanic patients are less likely than non-Hispanic whites to prefer antidepressants for treatment of depression.<sup>30</sup> Our data cannot distinguish between differences due to providers' biases and differences due to patients' preferences. It is also possible that the PHQ-9 may not accurately reflect severity of depression in some racial/ethnic groups, but previous research supports the validity of this measure across a wide range of language, culture, and race/ethnicity.<sup>24,26</sup>

Lower baseline depression scores were also more common among patients treated by psychiatrists, which may reflect a lower threshold for prescribing among psychiatrists or

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**Table 4. Secondary Analyses Examining Possible Bias Due to Missing Baseline Patient Health Questionnaire-9 (PHQ-9) Scores**

Analysis	PHQ-9 Score <sup>a</sup>				
	0–4	5–9	10–14	15–19	≥ 20
Full sample, unweighted	184 (3)	879 (12)	2,037 (29)	2,302 (33)	1,649 (23)
Full sample, weighted <sup>b</sup>	176 (3)	840 (12)	2,032 (29)	2,342 (33)	1,661 (24)
Limited to providers using PHQ-9 ≥ 75% <sup>c</sup>	66 (3)	278 (11)	762 (29)	870 (34)	615 (24)

<sup>a</sup>Values shown as n (%).<sup>b</sup>Weighted according to inverse probability of baseline PHQ-9 availability according to patients' sex, age, race/ethnicity, and neighborhood income as well as prescribing provider's specialty and number of patients treated for depression.<sup>c</sup>Limited to patients treated by prescribing providers for whom baseline PHQ-9 score was recorded for 75% or more of patients in study sample.

the fact that patients seen in specialty settings may more often present with other indications for treatment (such as comorbid psychiatric conditions or a past history of severe depression). Our data do not support concerns that increasing antidepressant prescribing by primary care physicians has led to diagnostic inflation or more frequent prescribing for mild depression.

It is possible that patterns of antidepressant prescribing in these health care systems differ from those in other practice settings. Practice guidelines in these systems all recommended routine use of the PHQ-9 for initial evaluation of depression. Guidelines in all systems did not recommend prescription of antidepressants for minimal or mild symptoms (PHQ-9 scores of 9 or less) and also advised that medication is not always indicated for moderate depression (PHQ-9 scores between 10 and 14). Those guidelines recommended either pharmacotherapy or psychotherapy as initial treatment for patients with moderate or moderately severe depression (PHQ-9 scores between 10 and 19) and combined pharmacotherapy and psychotherapy for those with more severe or chronic depression. Prescription of antidepressants for minimal or mild symptoms of depression could be more common in practice settings without these standard assessment procedures or practice guidelines. Nevertheless, the overall rate of antidepressant use in these health care systems was generally consistent with the rate in the US population. And use of standard measures, such as the PHQ-9, to guide depression treatment is increasingly common in both primary care and specialty mental health practice.<sup>31,32</sup> Furthermore, guideline recommendations regarding antidepressant treatment in these health care systems followed consensus recommendations, such as those of the Institute for Clinical Systems Improvement.<sup>33</sup>

Baseline depression severity scores were available for only 36% of patients starting antidepressant treatment, and this could bias our estimates of baseline severity. Availability of baseline severity scores was related to patient age, race/ethnicity, and neighborhood income (Table 1), but weighting for those predictors of missing baseline severity data had no meaningful effect on our estimates of baseline severity (Table 4). Availability of baseline severity scores was much more strongly related to provider characteristics (Table 1), and limiting our analyses to providers with high rates of PHQ-9 availability also had no meaningful effect on estimates of baseline severity (Table 4). Overall, we do not find evidence

that missing baseline PHQ-9 scores biased our primary finding, that only 15% of outpatients starting antidepressant treatment had only minimal or mild symptoms at the time of the initial prescription.

We believe these practice-based data more accurately assess severity of depression at initiation of treatment than do retrospective data from community surveys. As discussed above, longitudinal studies suggest that past episodes of

depression are often not recalled.<sup>15,18</sup> Furthermore, those who are not depressed at the time of interview are less likely to recall prior symptoms of depression.<sup>14</sup> Consequently, those who experience remission of depression while taking medication would be less likely to recall past symptoms of depression. Given this bias in recall, what appears to be unnecessary or inappropriate prescription of antidepressants for mild depression may actually represent successful treatment.

Several recent meta-analyses of placebo-controlled trials have attempted to identify a depression severity threshold for antidepressant prescribing—using varying patient samples and analytic methods.<sup>34–38</sup> These analyses have generally agreed regarding benefit of antidepressants for patients with severe depression and the absence of clear benefit for patients with mild depression. Conclusions have been mixed regarding a specific benefit of antidepressants for patients with moderate depression. Furthermore, the category of moderate depression is heterogeneous, including some with a high likelihood of spontaneous improvement and some for whom improvement without specific treatment is less likely.

We should acknowledge, however, that prescribing of antidepressants to patients with mild symptoms of depression may sometimes be appropriate. While every patient in the sample did receive a diagnosis of depressive disorder, it is possible that medication was prescribed primarily to address some other indication, such as co-occurring anxiety disorder. Standardized assessments, such as the PHQ-9, may not be an accurate measure of depression severity or depression-related impairment for every single patient. Some patients in our sample classified as having minimal or mild depression by the PHQ-9 could have indicated more severe symptoms to the prescribing physician during the visit. For a patient experiencing a relapse of depression following successful prior treatment with medication, reinitiating antidepressants when mild symptoms reappear (before more severe recurrence or relapse) would certainly be a reasonable practice. Our data do not allow us to identify these specific clinical decisions. Nevertheless, it is likely that these reasonable practices account for at least some of 15% of patients initiating antidepressant treatment for whom PHQ-9 scores indicated only minimal or mild depression.

The severity threshold for when to prescribe antidepressants is certainly not a bright line. Any attempt to evaluate the appropriateness of prescribing must allow

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for both the imperfection of standardized measures and the variability in individual patients' clinical histories. With those allowances, our finding that approximately 15% of outpatients starting antidepressant treatment reported mild or minimal depression does not seem particularly surprising or concerning. Data from these 4 health care systems do not indicate that overprescription of antidepressants for minimal or mild depression is a significant public health concern.

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**Supplementary material:** Available at PSYCHIATRIST.COM.

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## **Supplementary Material**

**Article Title:** Antidepressants Are Not Over-Prescribed for Mild Depression

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### **List of Supplementary Material for the article**

1. [eAppendix 1](#) Justification for 270 day interval to define a new episode of antidepressant treatment

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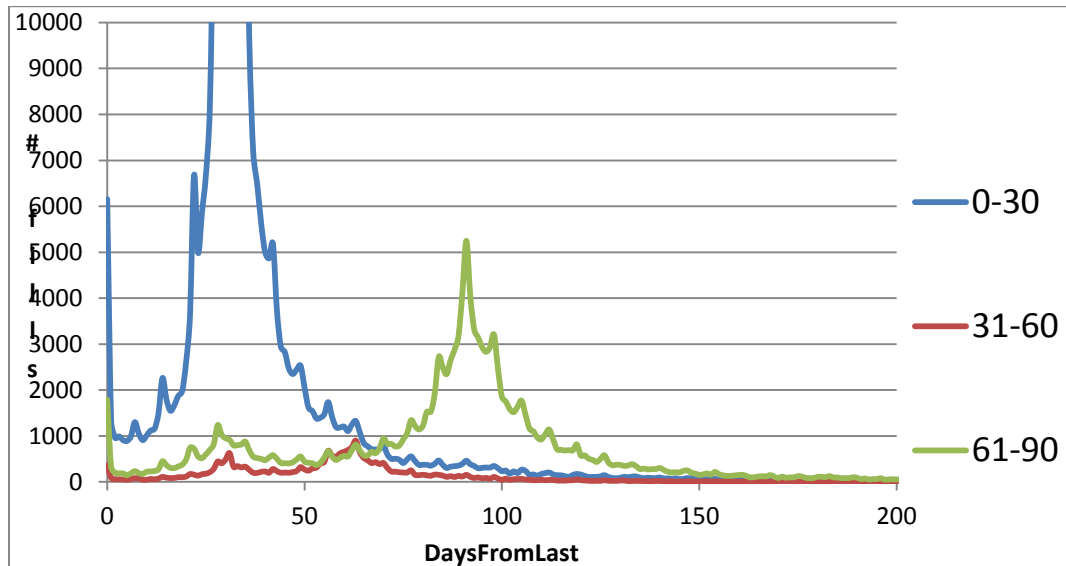
This Supplementary Material has been provided by the author(s) as an enhancement to the published article. It has been approved by peer review; however, it has undergone neither editing nor formatting by in-house editorial staff. The material is presented in the manner supplied by the author.



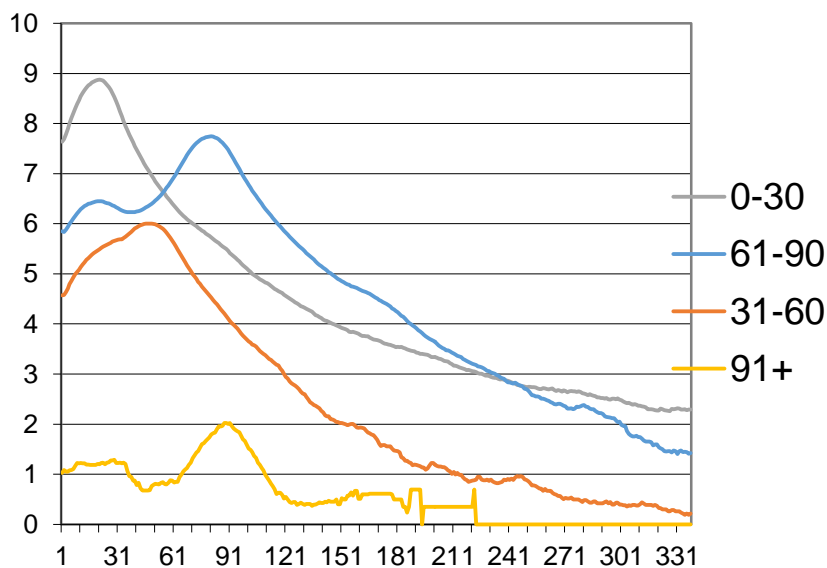
## eAppendix 1 – Justification for 270 day interval to define a new episode of antidepressant treatment

### 1) Descriptive analyses of intervals between prescription fills

Antidepressant refill data for 10 MHRN health systems in 2010 were organized to examine the frequency distribution of intervals between antidepressant fills, stratified by the days supply dispensed at the most recent prescription. Findings are shown in the figure below:



This figure suggests that the rate of refill does not decline to the background rate until the interval between refills exceeds 200 days. This is clearer when examining the signal-to-noise ratio ( $\mu/\sigma$ ) for refill rates over time, as shown in the figure below.



These data suggest that the rate of refill approaches the background rate (i.e. the rate of re-initiating treatment) after an interval of approximately 270 days.



## 2) Review of full-text medical records

Records data at two MHRN health systems were used to identify patients filling antidepressant prescriptions after intervals of varying duration since a prior antidepressant fill. Full-text medical records for a random sample of 200 patients were reviewed to evaluate how often records indicated continuation of long-term treatment vs. initiation of a new treatment episode after clear discontinuation of previous treatment. Results are shown in the table below:

Interval between fills	Continued treatment	New Episode	Missing or Unclear
180-210 days	36%	44%	20%
210-270 days	24%	54%	22%
270-300 days	18%	68%	14%

In cases where records indicated continued treatment, notes also suggested use of medication at less than the prescribed dose (so that a previous prescription intended to last 90 days might last for 180 days or longer).

Summary – We recognize that no single between-prescription interval or “washout period” will be completely accurate in distinguishing continuation of previous treatment from initiation of a new treatment episode. But these data suggest to us that using a 180 day interval to define a new episode would include an unacceptably high proportion of patients who are continuing longstanding treatment. Consequently, we choose to use a minimum interval of 270 days to define a new episode of antidepressant treatment.

### **Appendix B – Provider variability in use of PHQ9 at initiation of antidepressant treatment**

2011 data from Group Health Cooperative were used to examine variation between clinics and providers in the proportion of patients initiating antidepressant treatment with a PHQ9 score recorded during the baseline interval (14 days before to 3 days after the index prescription date).

Among clinics with at least 50 new treatment episodes, the median rate of PHQ9 completion was 49% with a range from 11% to 64%. Rates at the 25<sup>th</sup> and 75<sup>th</sup> percentile clinics were 39% and 56%, respectively.

Among providers with at least 20 new treatment episodes, the median rate of PHQ9 completion was 47% with a range from 3% to 92%.

We conclude that variation between providers and facilities in rates of PHQ9 far exceeds variation according to patients’ demographic or clinical characteristics.