

## **Supplementary Material**

Article Title: Effect of Zuranolone on Concurrent Anxiety and Insomnia Symptoms in Women With

Postpartum Depression

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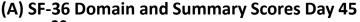
Response and Remission and NNH Estimates for Discontinuation Due to AE and Specific

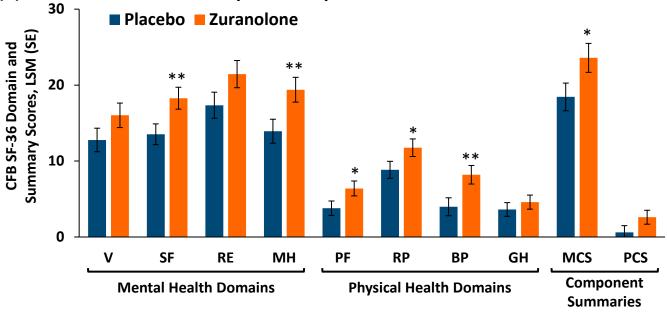
TEAEs (≥2% Incidence With Zuranolone and Greater Than That With Placebo).

## **Disclaimer**

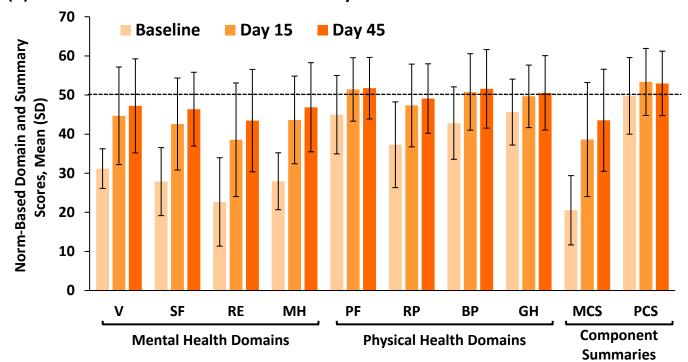
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Supplementary Figure 1. Patients' Perceptions of Their Own Functional Health and Well-being as Reported on the SF-36v2. (A) Change From Baseline in Domain and Summary Scores at Day 45. (B) Mean Domain and Component Summary Scores at Baseline, Day 15, and Day 45 (in Patients Receiving Zuranolone Only).





## (B) Norm-Based SF-36 Domain and Summary Scores



<sup>\*</sup>P<.05; \*\*P<.01 vs placebo. P values have not been adjusted for multiplicity and are nominal.

Abbreviations: BP = Bodily Pain; CFB = change from baseline; GH = General Health; LSM = least squares mean; MCS = Mental Component Score; MH = Mental Health; PCS = Physical Component Score; PF = Physical Functioning; RE = Role Emotional; RP = Role Physical; SD = standard deviation; SE = standard error; SF = Social Functioning; SF-36v2 = 36-Item Short Form Health Survey Instrument version 2; V = Vitality.

Supplementary Table 1. Summary of NNT Estimates for HDRS-17 Response, Remission, and Sustained Response and Remission and NNH Estimates for Discontinuation Due to AE and Specific TEAEs (≥2% Incidence With Zuranolone and Greater Than That With Placebo).

(A) NNT Estimates			
Outcome, n (%) <sup>a</sup>	Placebo (N = 74)	Zuranolone (N = 76)	Associated NNT (95% CI)
HDRS-17 response (day 15)	35 (47.9)	53 (71.6)	5 (3 to 13)
HDRS-17 remission (day 15)	17 (23.3)	33 (44.6)	5 (3 to 17)
Sustained HDRS-17 response (days 15/45)	27 (38.6)	44 (58.7)	5 (3 to 27)
Sustained HDRS-17 remission (days 15/45)	9 (12.7)	28 (37.3)	5 (3 to 10)
(B) NNH Estimates			
Outcome, n (%)	Placebo (N = 73)	Zuranolone (N = 78)	Associated NNH <sup>b</sup>
Treatment discontinuation due to AE	0	1 (1.3)	78
TEAE			
Somnolence	8 (11.0)	12 (15.4)	23
Dizziness	4 (5.5)	6 (7.7)	46
Sedation	0	4 (5.1)	20
Diarrhea	2 (2.7)	5 (6.4)	28
Dry mouth	0	3 (3.8)	26
Upper respiratory infection	1 (1.4)	6 (7.7)	16
Nasopharyngitis	1 (1.4)	3 (3.8)	41
Pain in extremity	1 (1.4)	2 (2.6)	84
Fatigue	1 (1.4)	3 (3.8)	41

<sup>&</sup>lt;sup>a</sup>Percentages were calculated using the numbers of patients with data evaluable at that day as denominator: N = 73 (placebo) and N = 74 (zuranolone) for HDRS-17 response and remission at day 15; N = 70 (placebo) and N = 75 (zuranolone) for sustained HDRS-17 response; and N = 71 (placebo) and N = 75 (zuranolone) for sustained HDRS-17 remission.

Abbreviations: AE = adverse events; CI = confidence interval; HDRS-17 = 17-item Hamilton Depression Rating Scale; NNT = number needed to treat; NNH = number needed to harm; TEAE = treatment-emergent adverse event.

<sup>&</sup>lt;sup>b</sup>Not statistically significant versus placebo at the p<0.05 threshold.