



CS/BSE&NSE/PR/2022-2023
November 16, 2022

To
The Manager
Department of Corporate Services
BSE Limited
25th Floor, P. J. Towers,
Dalal Street, Mumbai - 400 001

To
The Manager
Listing Department
National Stock Exchange of India Limited
Exchange Plaza, Bandra Kurla Complex
Bandra (E), Mumbai – 400 051

Scrip Code: 530239

Scrip Symbol: SUVEN

Dear Sir/Madam,

Sub: News Release

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With reference to above subject, please find enclosed News Release of our company titled **“Suven Life Sciences Limited Announces Randomization of First Patient in Phase-3 Global Clinical Trial of Masupirdine (SUVN-502) for the Treatment of Agitation in Patients with Dementia of the Alzheimer's Type”**

This is for your information and record.

Thanking You,
Yours faithfully,
For **Suven Life Sciences Limited**

Shrenik Soni
Company Secretary

Suven Life Sciences Limited

Registered Office: 8-2-334 | SDE Serene Chambers | 6th Floor Road No.5 | Avenue 7
Banjara Hills | Hyderabad – 500 034 | Telangana | India | CIN: L24110TG1989PLC009713
Tel: 91 40 2354 1142/ 3311/ 3315 Fax: 91 40 2354 1152 Email: info@suven.com website: www.suven.com

News Release

Suven Life Sciences Limited Announces Randomization of First Patient in Phase-3 Global Clinical Trial of Masupirdine (SUVN-502) for the Treatment of Agitation in Patients with Dementia of the Alzheimer's Type

HYDERABAD, INDIA (November 16, 2022) Suven Life Sciences Limited announces the randomization of first patient in the Phase-3 global clinical trial of Masupirdine for the treatment of agitation in patients with dementia of the Alzheimer's type.

The trial is a multicenter, randomized, double-blind, placebo-controlled study planned across 50 sites in the North America and Europe. The study will enroll approximately 375 patients who will be randomized in a 1:1:1 ratio to receive masupirdine either 50 mg QD or 100 mg QD or placebo QD for 12 weeks. The primary outcome measure is Cohen-Mansfield Agitation Inventory (CMAI) - Change in the CMAI items score aligning to the International Psychogeriatric Association (IPA) agitation criteria domains from baseline to Week 12. Key secondary outcome measure is modified Alzheimer's Disease Cooperative Study-Clinical Global Impression of Change (mADCS-CGI-C) scale as related to the agitation. Topline data from the trial is expected to be available by early 2025. Additional information on the trial can be found at ClinicalTrials.gov (NCT05397639).

About Masupirdine:

Masupirdine, a serotonin-6 (5-HT₆) receptor antagonist is being developed for the treatment of agitation in patients with dementia of the Alzheimer's type. In animal models, masupirdine showed significant reduction in agitation like behavior and modulated the neurotransmitters implicated in mood and behavior. Post-hoc analyses of the Phase 2 study (NCT02580305) evaluating masupirdine for the treatment of cognitive deficits in patients with moderate Alzheimer's disease (AD) suggested potential beneficial effects on agitation/aggression.

About Suven Life Sciences Limited ("Suven"):

Suven is a bio-pharmaceutical company, focused on discovering and developing novel pharmaceutical products, for central nervous system ("CNS") disorders using G Protein-Coupled Receptor targets. Our focus has been on discovery and development of innovative molecules targeting diseases and areas, which has undiscovered medical treatment opportunities. Our Company singularly focuses on development of "New Chemical Entities" ("NCEs") molecules for CNS diseases such as Alzheimer's, various forms of Dementia, Narcolepsy, Major Depressive Disorder ("MDD"), Attention Deficient Hyperactivity Disorder ("ADHD"), Huntington's disease, Parkinson, Bipolar disorder and different forms of neuropsychiatry disorders, gastro and pain. Suven has 7 clinical stage compounds, including

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this Phase 3 Masupirdine (SUVN-502) for treatment of agitation in patients with dementia of the Alzheimer's type, ongoing Phase 2 Samelisant (SUVN-G3031) for sleep disorders (Narcolepsy), Phase 2 ready Ropanicant (SUVN-911) for Major Depressive Disorder (MDD) and Phase 2 ready Usmarapride (SUVN-D4010) for cognitive disorders. In addition to clinical candidates, Suven has 8 molecules in development pipeline.

For more information please visit our website at <http://www.suven.com>

Risk Statement:

Except for historical information, all the statements, expectations, and assumptions, including expectations and assumptions, contained in this news release may be forward-looking statements that involve several risks and uncertainties. Although Suven attempts to be accurate in making these forward-looking statements, it is possible that future circumstances might differ from the assumptions on which such statements are based. Other important factors which could cause results to differ materially including outsourcing trends, economic conditions, dependence on collaborative partnership programs, retention of key personnel, technological advances, and continued success in growth of sales that may make our products/services offerings less competitive; Suven may not undertake to update any forward-looking statements that may be made from time to time.

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