



News Release

HYDERABAD, INDIA (29 Oct 2024) -- SUVEN Life Sciences Limited ("Suven") today announced unaudited financial results for the quarter & half year ended 30 September 2024. The unaudited financial results were reviewed by the audit committee and approved by the Board of Directors in their meeting held on 29 October 2024 at Hyderabad.

CONSOLIDATED STATEMENT OF OPERATIONS

INR Million, except EPS

	Quarter ended			Period ended		Year ended
	30-Sep-24	30-Jun-24	30-Sep-23	30-Sep-24	30-Sep-23	31-Mar-23
Revenue	66.09	50.14	82.01	116.23	176.38	328.23
R&D and Operational expenses	547.82	315.45	265.70	863.26	583.82	1,396.98
Depreciation and Amortisation	14.55	14.86	17.26	29.42	33.91	65.02
Finance cost	0.16	0.23	0.42	0.39	0.93	1.58
Total expenses	562.52	330.54	283.38	893.06	618.66	1,463.58
Exceptional items (insurance claim received)	-	-	74.57	-	74.57	74.57
Tax	-	-	-	-	-	(10.02)
Profit/(Loss) After Tax for the period/year	(496.44)	(280.40)	(126.80)	(776.84)	(367.71)	(1,050.76)
Other comprehensive income	(0.05)	(0.05)	0.48	(0.09)	0.95	(0.60)
Total comprehensive income	(496.48)	(280.44)	(126.33)	(776.93)	(366.76)	(1,051.36)
Paid up equity capital	218.07	218.07	218.07	218.07	218.07	218.07
Earnings per share of Rs.1 each (EPS)	(2.27)	(1.29)	(0.58)	(3.56)	(1.69)	(4.82)

- (a) Suven, a Biopharmaceutical company, engaged in Drug Discovery and Development of New Chemical Entities (NCEs) in Central Nervous System (CNS) disorders targeting unmet medical needs, globally.
- (b) The statement of operations includes financial of Suven Neurosciences, Inc., a Delaware Company, wholly owned subsidiary (WOS) of Suven, involved in clinical development programs of the Company.
- (c) Clinical development pipeline:
- SUVN-502 (Masupirdine) – Ongoing phase 3 study for Agitation and Aggression in Alzheimer's type dementias in North America and Europe; Enrolling patients in sites in US and Europe. Expected completion by end of calendar year 2026.
 - SUVN-G3031 (Samelisant) – Announced positive proof-of-concept results from its Phase 2 clinical trial assessing the safety and efficacy of Samelisant for the treatment of excessive daytime sleepiness (EDS) in adult narcolepsy patients with and without cataplexy. Planning to start Phase 3 registration clinical study for treatment of Narcolepsy in Q1-FY26.
 - SUVN-911 (Ropanicant) – Phase 2 open label study for Moderate to Severe Major Depressive Disorder in USA. Completed the study and outcome successful. Planning to start Phase 2b clinical study in Q1-FY26.
 - SUVN-D4010 (Usmarapride) – Planning for Phase 2 double blind study for the treatment of Cognition in depression in Q2-FY26.
 - SUVN-I6107 – Phase 1 study initiated during Q2-FY25 for establishing safety and pharmacokinetics of the molecule.
- (d) Since last reporting period, the Company has been granted 6 patents for its innovative drug discovery covering, Brazil, China, Hong Kong, Singapore and South Africa.

For more information on Suven please visit our Web site 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 that involve a number of 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 research and clinical development outcome, outsourcing trends, economic conditions, dependence on collaborative programs, retention of key personnel, technological advances and continued success in growth of revenue that may make our products/services offerings less competitive.

CIN: L24110TG1989PLC009713

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