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**Tolvaptan for patients with acute neurological injuries.**

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**Study results**

Seventy-nine patients were screened, of whom 25 were enrolled; 24 of these patients had an admission diagnosis of subarachnoid hemorrhage, one had an intracranial hemorrhage. The average dose of tolvapan given was 29.4mg. Sodium levels were 130.2±1.7 before tolvaptan, 132.3±3.4 at 12hrs, 135.4±3.1 at 24hrs, 137.2±3.5 at 48hrs, 136.1±3.4 at 72hrs, 137.8±2.9 at 96hrs, and 137.4±3.9 at 120hrs; the median treatment duration was 9 days. Enrollment was discontinued when 2 patients developed polyuria, persisting after tolvaptan discontinuation and likely unrelated to the drug; however, we deemed our screening process insufficient in excluding CSW (the likely cause of polyuria) and decided to halt the current study and develop a new protocol with better screening for CSW. No other adverse events occurred.