191

Shim K S¹, Bae J H¹, Lee J G¹, Lee K C², Kim H J³, Park H J⁴, Kwon C H⁴, Moon D G¹
1. Korea University Medical Center, 2. Inje University Hospital, 3. Dankook University Hospital, 4. Sungkyunkwan University Hospital

THE SAFETY AND EFFECTS OF DESMOPRESSIN IN THE TREATMENT OF ELDERLY PATIENTS WITH NOCTURIA FOR 1-YEAR FOLLOW-UP.

Hypothesis / aims of study

The purpose of this study was to investigate safety and effects of oral desmopressin (DDAVP) in elderly patients with nocturia for 1-year follow-up.

Study design, materials and methods

We selected 12 male patients over 60 years old who visited our outpatient clinic on complains of nocturia or BPH symptoms. They had more than three times of nocturia per night based on a 7-day voiding diary. They took one 0.2mg DDAVP tablet at bedtime every day for 1 year and urine and blood sampling were performed every 3month. In each serum, levels of BUN/ Creatinine, urea, nitrogen, glucose, Na, K, Cl, Ca, P, Mg, uric acid were measured. For timed urine study, urinary creatinine, urea, nitrogen, glucose, Na, K, Cl, Ca, P, Mg, uric acid, osmolality, volume of urine were measured.

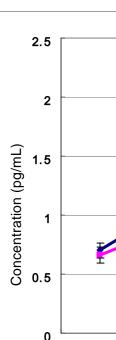
Results

Table Significant variables before and every 3 months and 1 month washout after 1 year medication with DDAVP *: stastistically significant (before vs 12mon)

†: stastistically significant (12mon vs 13mon) ‡: 1 month after stop medication

Mean±SD	Before	3mon.	6mon.	9mon.	12mon.	13mon. ‡
Total urine output (mL)	1719±521	1644±222	1768±775	1496±588	1495±394	1724±613
Noctunal urii output (mL)	^{1e} 641±265	498±268	481±304	430±510	441±308*	751±415†
No. of nocturia	4.85±0.93	1.11±1.36	1.89±0.93	1.88±0.83	1.43±0.79*	3.14±1.07†
Sleeping time (min) 400.0±36.9		394.3±66.3	421.4±47.0	439.4±59.6	427.2±55.7	414.7±48.6
1st voiding tin (min)	^{ne} 125.8±13.7	265.8±35.1	291.1±45.7	322.8±64.3	312.5±58.7*	155.5±47.1†
QoL (0-6)	5.5±0.52	2.1±0.97	1.7±0.58	1.9±0.65	1.5±0.82*	5.3±0.43†
Serum Na (mmol/L) 142±1.94		140±1.70	139±2.04	139±2.54	136±4.27*	139±1.05†
Urine Na (mmol/L) 186±81.1	203±49.4	170±64.5	190±82.6	209±63.7	157±65†
Urine osmolal (mOsm/Kg)	^{ity} 496±119	505±165	434±138	516±158	513±157	390±79.3†





11pm

Interpretation of results

Patient group of 12, average age of 69.2 (range 60 - 84) year-old had a total urine volume of 1719.08±5 nocturia volume of 641.17±265.17ml and nocturnal polyuria index of 0.37±0.13. There was no significant differential urine output before and after DDAVP administration, but nocturnal urine output was significantly decided 440.9±308.3 ml (p<0.01). After administration of DDAVP, sodium level was significantly increased from 186.8 to 209.4 mmol/L (p<0.05) in timed urinary electrolytes. Urinary chloride level was significantly increased from 180.3 a more deliced and provided the provided from 180.3 and 100.3 to 100.3 before administration.

228.3 mmol/L (p<0.05). Urine osmolality was significantly increased form 496.3 to 513.3 after administration of DDAVP

(p<0.05). The levels of ADH before medication were not significantly different from those of 1 month after stop the 1-year medication. No significant systemic complication was found during medication.

Concluding message

These results suggest that DDAVP induced natriuresis, and thus increased urine osmolality, condensing the urine, while decreasing nocturia. Desmopressin was well tolerated for I year in elderly patients with nocturia and long term administration of DDAVP did not affect the basal secretion of ADH, but the results suggest that serum sodium should be measured after treatment carefully.

References

- 1. Evaluation of the etiology of nocturia in men: the nocturia and nocturnal bladder capacity indices. Neurourol Urodyn 1999; 18: 559-65.
- 2. Abnormal diurnal rhythm of plasma vasopressin and urinary output in patients with enuresis. Am J Physiol 1989; 256: F664-71.
- 3. Efficacy of desmopressin in treatment of refractory nocturia in patients older than 65 years. Urology. 2002;59:485-9.

FUNDING: NONE DISCLOSURES: NONE

HUMAN SUBJECTS: This study was approved by the Korea University College of Medicine IRB and followed the Declaration of Helsinki Informed consent was obtained from the patients.