The incidence and diagnostic factors of polydipsia and polyuria: a single center survey in Japan

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Introduction and aim

Polydipsia and polyuria are one of the common chief complaints in the field of pediatric endocrinology. The differential diagnosis of polydipsia and polyuria are various diseases including diabetes mellitus (DM), central diabetes insipidus (CDI), and primary polydipsia (PP). DM is not difficult to diagnose, however, between DI and PP is sometimes difficult to diagnose. Although many examinations including a water deprivation test or hypertonic saline infusion test are needed to distinguish between CDI and PP, they are not convenient.

The objective of our study is to reveal the incidence of CDI and PP, and to investigate predictive factors for differentiation between CDI and PP.

Methods

This study was a retrospective chart review, performed from January 2014 to December 2018 in Department of pediatrics, Kurume University Hospital in Japan. The chief complaints of the patients, whose age was from 0 to 15 years old, were polydipsia and polyuria.

Variables analyzed including age, sex, symptom duration, height, BMI, drinking volume, urine volume and several biochemical tests.

Results Subjects From January 2014 to December 2018 First visit patients in the department of endocrinology of pediatrics, Kurume University Hospital n=1611**Exclusion:** 1) A chief complaint is not polydipsia and polyuria. 2) Patients diagnosed with diseases causing of polydipsia and polyuria previously. 3) Patients treated for polydipsia and polyuria. n=1584Patients with polydipsia, polyuria and both n = 27Exclusion: Hyperglycemia, Urine sugar positive Type 1 diabetes mellitus (T1DM) n = 16In our study, patients with hypercalcemia, hypercalciuria, hypokalemia Central diabetes insipidus (CDI) Primary polydipsia (PP) Noctural enuresis and nephrogenic diabetes insipidus were not present n=5n=5n=1

Biochemical tests*

	CDI (n=5)	PP (n=5)	P
Serum Na (mEq/L)	142.4 ± 4.15	140.0 ± 1.58	0.26
Hb (g/dL)	12.3 ± 1.7	13.9 ± 0.4	0.10
Alb (g/dL)	4.27 ± 0.44	4.69 \pm 0.34	
BUN (mg/dL)	11.82 ± 3.72	9.88 ± 1.14	0.31
Cr (mg/dL)	0.42 ± 0.23	0.35 ± 0.09	0.55
Serum osmolality (mOsm/kg)	296.2 ± 17.29	301.8 ± 45.4	0.80
Urine osmolality (mOsm/kg)	135.5 ± 35.0	430.4 ± 306.6	0.09
AVP (pg/ml)	0.50 ± 0.14	2.62 ± 2.31	0.10
Urine specific gravity (early morning)**	1.005 ± 0.002	1.015 ± 0.004	0.004

Date of first visit examination except for Urine specific gravity Without water deprivation

Significant at *p*<0.05

Patient Clinical Characteristics

	CDI (n=5)	PP (n=5)	P
Age (years)	7.8 ± 5.08	9.51 ± 3.90	0.54
Sex	M=2,F=3	M=3,F=2	1.00
Symptom duration (months)	26.4 ± 35.2	23.8 ± 33.75	0.9
Height (SD)	-1.46 ± 1.01	-0.39 ± 0.52	0.06
BMI (SD)	-0.75 ± 1.18	1.01 ± 0.27	0.02
Urination and drinking during bedtime	4*	3	_

 \times One patient is difficult to evaluate for infant . Significant at p<0.05

Significant at *p*<0.05

Drinking and Urine volume for 24 hours

	CDI (n=9)	PP (n=7)	p
Drinking (ml/m ²)	3628 ± 850	1900 ± 529	0.0006
Urine (ml/m²)	3707 ± 1278	1364 ± 881	0.001

MRI

	Age (years)	Posterior pituitary bright spot (T1WI)	Other MRI findings	Diagnosis
PP1	4	Present		
PP2	8	Present		
PP3	10	Present		
PP4	11	Not implemented		
PP5	12	Absent		
CDI1	0.3	Absent		Aftereffect of Neonatal meningitis
CDI2	5	Absent	Pituitary Cyst	Xanthogranuloma of the sellar region
CDI3	8	Absent	Pituitary gland enlargement, stalk thickening	germinoma
CDI4	11	Absent	Pituitary stalk thickening	Idiopathic?
CDI5	12	Absent	Pituitary stalk thickening	IgG4-related hypophysitis

Conclusion

- ✓ 11 patients (0.6% of first visit patients) with a chief complaint of polydipsia and polyuria in the absence of hyperglycemia, 5 CDI patients (0.4% of first visit patients) were identified. Moreover, in our medical region (Chikugo region), the incidence of T1DM was 2.2 per 100,000 population, and the incidence of CDI was 0.7 per 100,000 population of children. Hence, the incidence of CDI is low. To our knowledge, this study is the first to investigate the incidence of CDI in Japan.
- ✓ BMI and Urine Specific gravity were significantly lower in CDI than PP. Urine osmolality tended to be lower in CDI than in PP, however, not significantly.

 It may be due to small sample sizes. Moreover, the amount of drinking and urine volume was significantly higher in CDI than in PP. In MRI, posterior pituitary bright spot showed absence with all 5 CDI. These characteristics can help to distinguish between CDI and PP.

Reference

1) Haddad NG, et al. Endocr Pract. 2016 2) Dabrowski E, et al. Best Pract Res Clin Endocrinol Metab. 2016 3) Babiker AM, et al. J Trop Pediatr. 2015







