



# Correlation Between Bladder Sonographic Parameters and Uroflowmetry Patterns in Children with Urinary Dysfunction: A Case-Control Study

Reza Nafisi Moghaddam<sup>1</sup>, Mohammad Reza Dashti<sup>2</sup>, Ahmad Shajari<sup>2,\*</sup>

<sup>1</sup> Department of Radiology, Shahid Sadooghi University of Medical Sciences, Yazd, Iran

<sup>2</sup> Ali-Ebne-Abitaleb School of Medicine, Yazd Branch, Islamic Azad University, Yazd, Iran

\*Corresponding Author: Ali-Ebne-Abitaleb School of Medicine, Yazd Branch, Islamic Azad University, Yazd, Iran. Email: a\_shajari@yahoo.com

Received: 15 June, 2025; Revised: 23 September, 2025; Accepted: 31 December, 2025

## Abstract

**Background:** Ultrasound and uroflowmetry are widely used, noninvasive tests for pediatric lower urinary tract dysfunction (LUTD), yet the extent to which structural sonographic parameters mirror uroflowmetry patterns in individual children remains uncertain.

**Objectives:** The objective of this study is to compare bladder sonographic measures and uroflowmetry findings between symptomatic children and asymptomatic controls, and to examine whether specific uroflowmetry flow patterns correlate with bladder volume and wall thickness after adjustment for age and sex.

**Methods:** In this retrospective, two center case-control study (Shohadaye Kargar and Shahid Sadoughi hospitals, Yazd, Iran; January 2022 - December 2023), 240 children aged 5 - 14 years were enrolled with individual 1:1 matching on age ( $\pm 6$  months) and sex (120 cases; 120 controls). Cases had LUTD symptoms; controls were asymptomatic by standardized questionnaire. All children underwent transabdominal bladder ultrasound [volume, posterior wall thickness, postvoid residual (PVR)] and uroflowmetry (flow pattern, Qmax, Qavg, voided volume, flow time, TQmax). Analyses used *t*/Mann-Whitney and  $\chi^2$ /Fisher's tests as appropriate and analysis of covariance (ANCOVA) to adjust for age and sex. Sample size was calculated a priori (G\*Power v3.1) to detect a 1.0 mm difference in bladder wall thickness ( $\alpha = 0.05$ , power = 0.80).

**Results:** Bladder wall thickness was greater in cases than controls ( $3.65 \pm 1.02$  vs.  $2.48 \pm 0.27$  mm;  $P < 0.001$ ), and PVR was markedly higher ( $41.19 \pm 19.25$  vs.  $9.80 \pm 4.77$  mL;  $P < 0.001$ ). Bell shaped curves predominated in controls (87.5%) but were uncommon in cases (14.2%;  $P < 0.001$ ). Cases had lower Qmax and Qavg ( $8.88 \pm 3.34$  vs.  $13.82 \pm 3.45$  mL/s and  $5.76 \pm 2.23$  vs.  $8.82 \pm 2.38$  mL/s; both  $P < 0.001$ ). After adjustment for age and sex, uroflowmetry flow patterns were not significantly associated with bladder volume or wall thickness in either group (all  $P > 0.05$ ).

**Conclusions:** Children with LUTD demonstrate increased bladder wall thickness, higher PVR, and abnormal uroflowmetry profiles compared with asymptomatic peers; however, specific flow patterns do not map reliably onto structural ultrasound measures at the individual level. These findings support a multimodal diagnostic approach in which ultrasound and uroflowmetry are interpreted together rather than in isolation.

**Keywords:** Ultrasonography, Pediatrics, Lower Urinary Tract Symptoms, Uroflowmetry, Urinary Incontinence, Bladder Wall Thickening, Urinary Bladder Diseases, Dysfunctional Voiding

## 1. Background

Pediatric lower urinary tract dysfunction (LUTD) includes a heterogeneous spectrum of storage and voiding abnormalities that impair quality of life and, if

unrecognized, increase the risks of recurrent urinary tract infection, vesicoureteral reflux progression, and renal scarring. Noninvasive tests, particularly transabdominal bladder ultrasound and uroflowmetry, now anchor routine evaluation because they

characterize complementary domains of LUTD. Ultrasound assesses bladder morphology, wall characteristics, and postvoid residual (PVR). Uroflowmetry quantifies voiding dynamics and flow curve morphology. Although both modalities are well established, it remains uncertain how closely structural sonographic parameters map onto specific uroflowmetry patterns in individual children, and how best to integrate these data for diagnosis and management.

Normative work has clarified expectations for uroflow parameters and typical bell-shaped curves in healthy school age children. Sonographic protocols increasingly emphasize standardized filling and posterior wall measurements to improve reproducibility. However, robust age and sex stratified reference ranges for bladder wall thickness in children remain limited, and prior studies have reported mixed results regarding the strength of structure-function associations. Clinically, this uncertainty can lead to overreliance on a single modality or to inconsistent phenotyping of LUTD subtypes.

To address these gaps, we conducted a two center, age and sex matched, retrospective case-control study of children aged 5 to 14 years. We compared bladder wall thickness, bladder volume, and PVR with uroflowmetry patterns and quantitative flow measures. We hypothesized that, while cases would differ markedly from controls on both structural and functional indices, specific uroflowmetry flow patterns would show at most weak associations with ultrasound parameters after accounting for age and sex.

## 2. Objectives

Our objectives were threefold: (1) To quantify between group differences in sonographic and uroflowmetry measures; (2) to describe the distribution of flow curve phenotypes in symptomatic versus asymptomatic children; and (3) to examine the adjusted relationships between flow patterns and bladder volume and wall thickness. These data aim to inform a multimodal, evidence based diagnostic approach that interprets ultrasound and uroflowmetry together, rather than in isolation, for optimized assessment of pediatric LUTD.

## 3. Methods

### 3.1. Study Design and Setting

This retrospective, two center case-control study was conducted at Shohadaye Kargar and Shahid Sadoughi hospitals (Yazd, Iran) from January 2022 to December 2023. The protocol was approved by the Ethics Committee of Shahid Sadoughi University of Medical Sciences (IR.IAU.KHUISF.REC.1404.408) and adhered to the Declaration of Helsinki. We enrolled 240 children aged 5 - 14 years and allocated them equally to case and control groups using individual 1:1 matching.

### 3.2. Participants and Sample Size Calculation

Cases were children presenting with urinary dysfunction symptoms, including daytime incontinence, urgency, frequency > 8 voids/day, dysuria, nocturnal enuresis ( $\geq 2$  episodes/week in children > 5 years), or subjective voiding difficulty. Controls were recruited from routine health visits or non-urological evaluations and were confirmed asymptomatic via standardized questionnaires completed by parents and children. Individual 1:1 matching was performed on sex (identical) and age (within  $\pm 6$  months) using computer generated random selection from eligible controls.

Sample size was calculated a priori with G\*Power v3.1 assuming a between group mean difference of 1.0 mm in bladder wall thickness (SD = 1.2 mm),  $\alpha = 0.05$ , and power = 0.80, yielding 118 participants per group. To accommodate potential missingness, we included 120 per group (total n = 240). Exclusion criteria were incomplete diagnostic data, neurological disease affecting bladder function, congenital urinary tract abnormalities, active urinary tract infection at assessment, prior urological surgery, or lack of parental consent.

### 3.3. Sonographic Assessment

Transabdominal bladder ultrasound was performed on a Voluson™ E8 system (GE Healthcare, Chicago, IL) with a 2 - 8 MHz convex transducer by radiologists with  $\geq 5$  years' pediatric experience, following standardized protocols. Children were scanned with moderate bladder filling {~50 - 75% of expected capacity; [age (years) + 1]  $\times$  30 mL if < 12 years and 350 - 400 mL if  $\geq 12$  years}.

Bladder volume was calculated with the ellipsoid formula (length  $\times$  width  $\times$  height  $\times$  0.52) using sagittal and transverse measurements. Bladder wall thickness was measured at the posterior wall in the midsagittal plane during moderate filling, avoiding trabeculation and artifacts; the average of three measurements was

recorded. The PVR was measured within 5 minutes of complete voiding.

For reliability, a subset of 30 examinations underwent duplicate measurements by two independent operators, yielding intraclass correlation coefficients (ICC) > 0.90 for all parameters.

### 3.4. Uroflowmetry Assessment

Uroflowmetry was performed with the MMS UD 2000 system (Medical Measurement Systems, Enschede, Netherlands) according to International Children's Continence Society guidelines. Children received brief verbal instructions and voided in a private setting in their natural position (standing for boys, sitting for girls) after achieving comfortable bladder fullness. Only voids  $\geq 50$  mL were analyzed; testing was repeated when initial volume was insufficient.

Recorded parameters included maximum flow rate (Qmax), average flow rate (Qavg), voided volume, flow time, and time to maximum flow (TQmax). Flow curves were classified as bell shaped (normal), plateau, tower shaped, staccato, or intermittent by two independent observers, with discrepancies resolved by consensus.

### 3.5. Data Collection and Statistical Analysis

Data extracted from medical records included age, sex, weight, height, BMI, presenting symptoms, family history of urinary problems, prior urinary tract infections, comorbid conditions (particularly constipation), and all sonographic and uroflowmetry measurements. Data were entered on standardized forms with double data entry by two independent researchers; discrepancies were resolved by record review. De identified datasets were stored in password protected databases.

Analyses were conducted in SPSS v26 (IBM, Armonk, NY). Distributions were assessed with the Shapiro-Wilk test and visual inspection. Continuous variables are reported as mean  $\pm$  SD or median (IQR); categorical variables as No. (%). Between group comparisons used independent *t*-tests or Mann-Whitney U tests (continuous) and chi square or Fisher's exact tests (categorical). Analysis of covariance (ANCOVA) adjusted for age and sex. Correlations were assessed with Spearman's rank correlation (continuous) and contingency analysis (categorical). All tests were two tailed, with statistical significance at  $P < 0.05$ . Effect sizes are reported as Cohen's *d* (pooled SD) for

continuous outcomes and Cramer's V for categorical associations.

## 4. Results

### 4.1. Participant Characteristics

A total of 240 children were enrolled (120 cases; 120 controls). Sex distribution did not differ between groups [147 males (61.3%), 93 females (38.8%);  $\chi^2 = 0.235$ ,  $P = 0.629$ ]. Mean age was comparable (controls:  $10.05 \pm 2.77$  years; cases:  $9.92 \pm 2.79$  years;  $t = 0.348$ ,  $P = 0.728$ ). Anthropometric measures were similar (controls vs. cases):

- Weight:  $32.8 \pm 11.2$  vs.  $31.9 \pm 10.8$  kg.
- Height:  $135.4 \pm 16.3$  vs.  $134.2 \pm 15.8$  cm.
- BMI:  $17.6 \pm 3.2$  vs.  $17.3 \pm 3.1$  kg/m<sup>2</sup> (all  $P > 0.05$ ).

### 4.2. Clinical Features

In the case group, the most frequent presenting symptoms were incontinence (27/120, 22.5%), urinary frequency (26/120, 21.7%), dysuria (22/120, 18.3%), and nocturnal enuresis (21/120, 17.5%). Controls were confirmed asymptomatic via standardized parent- and child-reported questionnaires. Family history of urinary problems was more prevalent in cases than controls (38.3% vs. 10.8%;  $\chi^2 = 23.1$ ,  $P < 0.001$ ). Prior urinary tract infection was also more common in cases (42.5% vs. 9.2%;  $\chi^2 = 34.2$ ,  $P < 0.001$ ). Constipation occurred in 35.0% of cases vs. 10.0% of controls ( $\chi^2 = 20.8$ ,  $P < 0.001$ ). Comorbidities beyond constipation were not systematically assessed.

### 4.3. Sonographic Parameters

Bladder volume did not differ significantly between groups (controls:  $217.92 \pm 88.12$  mL; cases:  $205.67 \pm 81.90$  mL;  $t = 1.098$ ,  $P = 0.625$ ). In contrast, bladder wall thickness was greater in cases ( $3.65 \pm 1.02$  mm) than controls ( $2.48 \pm 0.27$  mm;  $t = 11.2$ ,  $P < 0.001$ ; Cohen's *d* = 1.45). The PVR volume was markedly higher in cases ( $41.19 \pm 19.25$  mL) vs. controls ( $9.80 \pm 4.77$  mL; Mann-Whitney U = 1,847,  $P < 0.001$ ; Cohen's *d* = 2.31). Abnormal bladder morphology was more prevalent in cases (28.3% vs. 4.2%;  $\chi^2 = 24.1$ ,  $P < 0.001$ ; Cramer's V = 0.32), as was trabeculation (30.8% vs. 5.0%;  $\chi^2 = 26.3$ ,  $P < 0.001$ ; Cramer's V = 0.33). In controls only, males had larger mean bladder volumes than females ( $238.5 \pm 91.2$  vs.  $201.3 \pm 82.4$  mL;  $t = 2.07$ ,  $P = 0.042$ ); no sex difference was

**Table 1.** Sonographic Parameters in Cases and Controls <sup>a</sup>

Parameters	Controls (N = 120)	Cases (N = 120)	P-Value	Effect Size <sup>b</sup>
Bladder volume (mL)	217.92 ± 88.12	205.67 ± 81.90	0.625	0.15
Wall thickness (mm)	2.48 ± 0.27	3.65 ± 1.02	< 0.001	1.45
Post-void residual (mL)	9.80 ± 4.77	41.19 ± 19.25	< 0.001	2.31
Normal morphology	115 (95.8)	86 (71.7)	< 0.001	0.32
Trabeculation present	6 (5.0)	37 (30.8)	< 0.001	0.33

<sup>a</sup> Values are expressed as mean ± SD or No. (%).

<sup>b</sup> Effect size denotes Cohen's d for continuous variables and Cramer's V for categorical variables.

observed in cases ( $P = 0.187$ ). Sonographic parameters are summarized in [Table 1](#).

#### 4.4. Uroflowmetry Patterns and Parameters

Normal bell-shaped curves predominated in controls (105/120, 87.5%) but were uncommon in cases (17/120, 14.2%;  $\chi^2 = 128.7$ ,  $P < 0.001$ ; Cramer's  $V = 0.73$ ). Among cases, abnormal patterns included intermittent (30/120, 25.0%), staccato (30/120, 25.0%), plateau (25/120, 20.8%), and tower-shaped (18/120, 15.0%); tower-shaped and staccato were observed exclusively in cases.

Maximum flow rate ( $Q_{max}$ ) was lower in cases ( $8.88 \pm 3.34$  mL/s) than controls ( $13.82 \pm 3.45$  mL/s;  $t = 11.1$ ,  $P < 0.001$ ; Cohen's  $d = 1.45$ ). Average flow rate ( $Q_{avg}$ ) was likewise reduced ( $5.76 \pm 2.23$  vs.  $8.82 \pm 2.38$  mL/s;  $t = 10.0$ ,  $P < 0.001$ ; Cohen's  $d = 1.32$ ). Voided volume ( $215.43 \pm 92.79$  vs.  $209.54 \pm 87.86$  mL;  $P = 0.614$ ), flow time ( $19.63 \pm 4.81$  vs.  $19.86 \pm 4.64$  s;  $P = 0.703$ ), and time to maximum flow ( $TQ_{max}$ :  $8.42 \pm 2.89$  vs.  $8.15 \pm 2.74$  s;  $P = 0.482$ ) did not differ significantly. Detailed uroflowmetry parameters and flow pattern distributions are presented in [Table 2](#).

#### 4.5. Structure-Function Correlations and Covariate Effects

No significant associations were detected between flow pattern categories and bladder volume or wall thickness (controls: Volume:  $F = 1.10$ ,  $P = 0.335$ ; wall thickness:  $F = 0.18$ ,  $P = 0.835$ ; cases: Volume:  $F = 1.17$ ,  $P = 0.326$ ; wall thickness:  $F = 0.75$ ,  $P = 0.558$ ). In ANCOVA models adjusting for age and sex, these findings remained non-significant. Age correlated positively with bladder volume in both groups (controls:  $R = 0.73$ ,  $P < 0.001$ ; cases:  $R = 0.69$ ,  $P < 0.001$ ). A sex effect on bladder volume was observed only among controls ( $F = 4.2$ ,  $P = 0.042$ ). No significant interactions between flow pattern and sex were identified (all  $P > 0.10$ ). Comparisons of bladder sonographic parameters by flow pattern are shown in [Table 3](#).

## 5. Discussion

In these two centers, age and sex matched case-control study, children with urinary dysfunction exhibited significantly thicker bladder walls, markedly higher PVR volumes, and substantially more abnormal uroflowmetry patterns than asymptomatic peers, reinforcing the clinical value of a multimodal, noninvasive evaluation pathway advocated by the International Children's Continence Society and related guidance (1-3). The magnitude of the between group differences for bladder wall thickness and PVR, together with the predominance of non-bell shaped curves in the case group, mirrors prior pediatric reports linking structural bladder changes and inefficient emptying with dysfunctional voiding phenotypes (4-11). At the same time, our analyses confirmed that, after adjustment for age and sex, specific uroflowmetry flow patterns did not correlate strongly with ultrasound parameters at the individual level, a finding that is consistent with the emerging view that these modalities provide complementary rather than redundant information (1, 12).

The absence of strong pattern specific correlations likely reflects the multifactorial pathophysiology of pediatric LUTD. Similar flow morphologies can arise from different underlying mechanisms; staccato curves, for example, may reflect pelvic floor discoordination, behavioral withholding, or pain, while increases in bladder wall thickness may result from chronic detrusor overactivity, reduced compliance, or compensatory responses to outlet resistance (6-8, 13-17). Sonographic measures also remain state dependent: Despite our standardization to moderate filling, residual variability in filling level and regional wall thickness can attenuate linear associations with concurrent flow morphology (5, 8, 18). Moreover, both ultrasound and uroflowmetry in routine practice are often obtained as single occasion

**Table 2.** Uroflowmetry Parameters and Flow Patterns<sup>a</sup>

Parameters	Controls (N = 120)	Cases (N = 120)	P-Value	Effect Size
Qmax (mL/s)	13.82 ± 3.45	8.88 ± 3.34	< 0.001	1.45
Qavg (mL/s)	8.82 ± 2.38	5.76 ± 2.23	< 0.001	1.32
Voided volume (mL)	209.54 ± 87.86	215.43 ± 92.79	0.614	0.07
Flow time (s)	19.86 ± 4.64	19.63 ± 4.81	0.703	0.05
Time to Qmax (s)	8.15 ± 2.74	8.42 ± 2.89	0.482	0.10
<b>Flow pattern</b>			< 0.001	0.73 <sup>b</sup>
Bell-shaped (normal)	105 (87.5)	17 (14.2)		
Plateau	10 (8.3)	25 (20.8)		
Tower-shaped	0 (0)	18 (15.0)		
Staccato	0 (0)	30 (25.0)		
Intermittent	5 (4.2)	30 (25.0)		

<sup>a</sup> Values are expressed as mean ± SD or No. (%).

<sup>b</sup> Effect size for flow pattern is Cramer's V.

tests; within child variability in hydration, stool burden, anxiety, and cooperation can affect volumes, PVR, and flow curves, blurring structure–function coupling on any one day (1, 11, 19, 20). Finally, relationships between parameters may be non-linear or threshold based, e.g., risk increases above certain PVR levels, so average differences across broad pattern categories may not capture clinically meaningful inflection points.

Clinically, these observations argue against inferring pathophysiology from either modality in isolation. Instead, ultrasound and uroflowmetry should be interpreted together and in clinical context, including symptom profiles, voiding diaries, and constipation assessment. In practice, clearly increased bladder wall thickness and consistently elevated PVR, particularly when accompanied by staccato or intermittent flow patterns and reduced flow rates, should prompt structured urotherapy (timed voiding, fluid optimization, constipation treatment), consideration of pelvic floor biofeedback for suspected dysfunctional voiding, and escalation to urodynamic testing when abnormalities persist or when findings are discordant. Where local protocols apply numeric triggers, thresholds such as bladder wall thickness above approximately 3.0 mm for this age range or repeated PVR values above 20 mL may justify earlier referral, especially in the setting of recurrent urinary tract infections or refractory symptoms.

Our results fit well within the literature. The predominance of bell-shaped curves among controls and the lower Qmax and Qavg in cases align with normative uroflow data for school age children (5) and

with studies reporting reduced flow in dysfunctional voiding syndromes (21-25). The larger bladder wall thickness in symptomatic children and the markedly higher PVR echo prior work that associates structural bladder changes with impaired emptying and greater risk for infectious or reflux related sequelae (5-11, 26, 27). At the same time, the weak individual level structure–function coupling we observed has been noted in multimodal pediatric series, supporting a diagnostic strategy that synthesizes rather than substitutes across modalities (1, 12).

This study has several strengths. We used individual 1:1 matching on age and sex, standardized acquisition protocols for both ultrasound and uroflowmetry, and demonstrated excellent inter operator reliability for sonographic measurements (ICC > 0.90). We captured both curve morphology and quantitative flow parameters, including time to maximum flow, and we adjusted analyses for age and sex using ANCOVA, acknowledging known developmental effects on bladder capacity and voiding function. The sample size afforded robust power for the primary between group comparisons.

Limitations must also be acknowledged. As a retrospective, hospital based, two center study, our design limits causal inference and may not fully represent community populations. Although ultrasounds were performed at moderate filling, bladder wall thickness remains filling dependent, and single occasion imaging and uroflowmetry cannot account for day-to-day variability. Flow pattern classification, while based on established criteria, is

**Table 3.** Sonographic Parameters by Flow Pattern <sup>a,b</sup>

Variables	Bladder Volume (mL)	Wall Thickness (mm)
<b>Controls</b>		
Bell-shaped	222.30 ± 87.98	2.49 ± 0.28
Plateau	184.90 ± 93.45	2.46 ± 0.26
Intermittent	192.20 ± 77.63	2.42 ± 0.08
<b>Cases</b>		
Bell-shaped	189.88 ± 79.71	3.67 ± 1.13
Plateau	193.88 ± 80.07	3.42 ± 0.92
Tower-shaped	214.06 ± 77.38	3.56 ± 1.07
Staccato	230.57 ± 84.97	3.88 ± 1.01
Intermittent	194.50 ± 83.10	3.66 ± 1.01

<sup>a</sup> Values are expressed as mean ± SD.

<sup>b</sup> One-way analysis of covariance (ANCOVA, adjusted for age and sex) showed no significant association between flow pattern and bladder volume or wall thickness in either group (controls:  $F = 1.10, P = 0.335$ ;  $F = 0.18, P = 0.835$ ; cases:  $F = 1.17, P = 0.326$ ;  $F = 0.75, P = 0.558$ ).

susceptible to residual inter observer differences; similarly, we concentrated comorbidity ascertainment on constipation and did not systematically quantify other modifiers such as medication use, psychological factors, or stool burden by imaging. Finally, we did not incorporate invasive urodynamics or surface electromyography, which could have validated noninvasive phenotypes in discordant cases and refined mechanistic interpretation.

Future research should move beyond cross sectional snapshots to prospective, longitudinal, multicenter designs with repeated same day ultrasound and uroflowmetry under standardized filling targets, permitting within child modeling of variability and response to urotherapy. Parallel efforts should generate age and sex stratified normative nomograms for pediatric bladder wall thickness and PVR, ideally reported as Z scores to facilitate interpretation across the 5 - 14-year range (2, 3, 5, 19, 20, 28, 29). Analytically, pre specified multivariable models and modern methods such as penalized regression or machine learning classifiers could integrate quantitative flow features, curve shape descriptors, PVR, and sonographic metrics to improve discrimination and calibration for clinically relevant outcomes. Mechanistic substudies incorporating selective urodynamics and, where feasible, surface EMG would help link noninvasive signatures to detrusor pressure and sphincter activity. Comprehensive comorbidity assessment, including standardized constipation scoring, stool burden quantification, hydration tracking, and patient reported outcomes, should be embedded to contextualize structure-function relationships and tailor therapy.

Finally, implementation studies should test the cost effectiveness and clinical impact of “ultrasound plus uroflowmetry for all” versus risk stratified pathways in real world pediatric urology clinics.

### 5.1. Conclusions

Children with urinary dysfunction have greater bladder wall thickness, higher PVR, and abnormal uroflowmetry compared with asymptomatic peers, yet individual level correlations between ultrasound parameters and specific flow morphologies are weak. These findings support a multimodal diagnostic approach in which ultrasound and uroflowmetry are interpreted together and against the clinical backdrop, rather than relied upon singly. Prospective, age and sex stratified, and analytically advanced studies are needed to establish normative sonographic values, validate noninvasive phenotypes against mechanistic measures, and determine the pathway level impact of integrated testing on outcomes and resource use.

### Footnotes

**AI Use Disclosure:** The authors declare that no generative AI tools were used in the creation of this article.

**Authors' Contribution:** Study concept and design: Reza Nafisi Moghaddam and Mohammad Reza Dashti; Acquisition of data: Ahmad Shajari and Mohammad Reza Dashti; Analysis and interpretation of data: Ahmad Shajari and Reza Nafisi Moghaddam; Drafting of the

manuscript: Ahmad Shajari; Critical revision of the manuscript for important intellectual content: Reza Nafisi Moghaddam; Statistical analysis: Ahmad Shajari; Administrative, technical, and material support: Mohammad Reza Dashti; Study supervision: Reza Nafisi Moghaddam.

**Conflict of Interests Statement:** All authors declare no conflict of interest.

**Data Availability:** The dataset presented in the study is available on request from the corresponding author during submission or after publication. The data are not publicly available due to ethical and privacy restrictions regarding pediatric patient information.

**Ethical Approval:** This study was approved by the Ethics Committee of Shahid Sadoughi University of Medical Sciences, Yazd, Iran, under the ethical approval code [IR.IAU.KHUISF.REC.1404.408](https://doi.org/10.1016/j.jpuro.2014.10.002).

**Funding/Support:** This study received no external funding and was conducted using institutional resources.

**Informed Consent:** Written informed consent was obtained from the parents or legal guardians of all participating children prior to inclusion in the study.

## References

1. Van Batavia JP, Combs AJ. The Role of Non-invasive Testing in Evaluation and Diagnosis of Pediatric Lower Urinary Tract Dysfunction. *Curr Urol Rep*. 2018;**19**(5):34. [PubMed ID: [29623450](https://pubmed.ncbi.nlm.nih.gov/29623450/)]. <https://doi.org/10.1007/s11934-018-0784-1>.
2. Austin PF, Bauer SB, Bower W, Chase J, Franco I, Hoebeke P, et al. The standardization of terminology of lower urinary tract function in children and adolescents: Update report from the standardization committee of the International Children's Continence Society. *Neurourol Urodyn*. 2016;**35**(4):471-81. [PubMed ID: [25772695](https://pubmed.ncbi.nlm.nih.gov/25772695/)]. <https://doi.org/10.1002/nau.22751>.
3. Tekgül S, Dogan HS, Hoebeke P, et al. *EAU Guidelines on Paediatric Urology*. European Association of Urology; 2020. Available from: <https://uroweb.org/guidelines/paediatric-urology>.
4. Ozturk YD, Elmas AT, Tabel Y. Uroflowmetry parameters in healthy children between 5 and 15 years old. *Low Urin Tract Symptoms*. 2023;**15**(6):231-7. [PubMed ID: [37614060](https://pubmed.ncbi.nlm.nih.gov/37614060/)]. <https://doi.org/10.1111/luts.12499>.
5. Marzuillo P, Guarino S, Capalbo D, Acierno S, Menale F, Prisco A, et al. Interrater reliability of bladder ultrasound measurements in children. *J Pediatr Urol*. 2020;**16**(2):219 e1-7. [PubMed ID: [31980386](https://pubmed.ncbi.nlm.nih.gov/31980386/)]. <https://doi.org/10.1016/j.jpuro.2019.12.015>.
6. Taskinen S, Heikkilä J, Rintala R. Ultrasonographic bladder wall thickness measurement in the evaluation of neurogenic bladder dysfunction in children and adolescents. *J Urol*. 2007;**178**(6):2571-5.
7. Brownrigg N, Hane B, Pemberton J; et al. The bladder wall thickness in normal paediatric populations. *J Pediatr Urol*. 2009;**5**(5):395-8.
8. Yeung CK, Chiu HN, Sit FKY. Bladder dysfunction in children with refractory monosymptomatic primary nocturnal enuresis. *J Urol*. 1999;**162**(3 Part 2):1049-54.
9. Nguyen HT, Benson CB, Bromley B, Campbell JB, Chow J, Coleman B, et al. Multidisciplinary consensus on the classification of prenatal and postnatal urinary tract dilation (UTD classification system). *J Pediatr Urol*. 2014;**10**(6):982-98. [PubMed ID: [25435247](https://pubmed.ncbi.nlm.nih.gov/25435247/)]. <https://doi.org/10.1016/j.jpuro.2014.10.002>.
10. Koff SA, Wagner TT, Jayanthi VR. The relationship among dysfunctional elimination syndromes, primary vesicoureteral reflux and urinary tract infections in children. *J Urol*. 1998;**160**(3 Pt 2):1019-22. [PubMed ID: [9719268](https://pubmed.ncbi.nlm.nih.gov/9719268/)]. <https://doi.org/10.1097/00005392-199809020-00014>.
11. Sillen U. Bladder dysfunction in children: pitfalls and challenges for the clinician. *Pediatr Nephrol*. 2001;**16**(7):518-23.
12. Palmer LS, Zebold K, Firlit CF, Kaplan WE. Complications of intravesical oxybutynin chloride therapy in the pediatric myelomeningocele population. *J Urol*. 1997;**157**(2):638-40. [PubMed ID: [8996387](https://pubmed.ncbi.nlm.nih.gov/8996387/)].
13. Franco I. Functional bladder problems in children: pathophysiology, diagnosis, and treatment. *Pediatr Clin North Am*. 2012;**59**(4):783-817. [PubMed ID: [22857829](https://pubmed.ncbi.nlm.nih.gov/22857829/)]. <https://doi.org/10.1016/j.pcl.2012.05.007>.
14. Bauer SB. Pediatric uroflowmetry. *Urol Clin North Am*. 1996;**23**(2):273-7.
15. Neveus T. Pathogenesis of enuresis: Towards a new understanding. *Int J Urol*. 2017;**24**(3):174-82. [PubMed ID: [28208214](https://pubmed.ncbi.nlm.nih.gov/28208214/)]. <https://doi.org/10.1111/iju.13310>.
16. de Jong TP, Chrzan R, Klijn AJ, Dik P. Treatment of the neurogenic bladder in spina bifida. *Pediatr Nephrol*. 2008;**23**(6):889-96. [PubMed ID: [18350321](https://pubmed.ncbi.nlm.nih.gov/18350321/)]. [PubMed Central ID: [PMC2335291](https://pubmed.ncbi.nlm.nih.gov/PMC2335291/)]. <https://doi.org/10.1007/s00467-008-0780-7>.
17. Mahant S, Friedman J, MacArthur C. Renal ultrasound findings and vesicoureteral reflux in children hospitalised with urinary tract infection. *Arch Dis Child*. 2002;**86**(6):419-20. [PubMed ID: [12023172](https://pubmed.ncbi.nlm.nih.gov/12023172/)]. [PubMed Central ID: [PMC1762998](https://pubmed.ncbi.nlm.nih.gov/PMC1762998/)]. <https://doi.org/10.1136/adc.86.6.419>.
18. Berger RM, Maizels M, Moran GC, Conway JJ, Firlit CF. Bladder capacity (ounces) equals age (years) plus 2 predicts normal bladder capacity and aids in diagnosis of abnormal voiding patterns. *J Urol*. 1983;**129**(2):347-9. [PubMed ID: [6834505](https://pubmed.ncbi.nlm.nih.gov/6834505/)]. [https://doi.org/10.1016/s0022-5347\(17\)52091-1](https://doi.org/10.1016/s0022-5347(17)52091-1).
19. von Gontard A, Equit M. Comorbidity of ADHD and incontinence in children. *Eur Child Adolesc Psychiatry*. 2015;**24**(2):127-40. [PubMed ID: [24980793](https://pubmed.ncbi.nlm.nih.gov/24980793/)]. <https://doi.org/10.1007/s00787-014-0577-0>.
20. Averbek MA, Madersbacher H. Constipation and LUTS - how do they affect each other? *Int Braz J Urol*. 2011;**37**(1):16-28. [PubMed ID: [21385476](https://pubmed.ncbi.nlm.nih.gov/21385476/)]. <https://doi.org/10.1590/s1677-5538201000100003>.
21. Shaikh N, Morone NE, Bost JE, Farrell MH. Prevalence of urinary tract infection in childhood: a meta-analysis. *Pediatr Infect Dis J*. 2008;**27**(4):302-8. [PubMed ID: [18316994](https://pubmed.ncbi.nlm.nih.gov/18316994/)]. <https://doi.org/10.1097/INF.0b013e31815e4122>.
22. Bauer SB, Retik AB, Colodny AH, Hallett M, Khoshbin S, Dyro FM. The unstable bladder in childhood. *Urol Clin North Am*. 1980;**7**(2):321-36. [PubMed ID: [7404871](https://pubmed.ncbi.nlm.nih.gov/7404871/)].
23. Glassberg KI, Combs AJ, Horowitz M. Nonneurogenic bladder dysfunction in children and adolescents: current concepts in diagnosis and management. *Urol Clin North Am*. 1995;**22**(1):133-50.

24. Ergun R, Ozturk NI, Sekerci CA. The effect of duration between sessions on biofeedback treatment in children with dysfunctional voiding. *Low Urin Tract Symptoms*. 2022;**14**(5):387-92. [PubMed ID: 35808986]. <https://doi.org/10.1111/luts.12456>.
25. Hoebeke P, Van Laecke E, Van Camp C, Raes A, Van De Walle J. One thousand video-urodynamic studies in children with non-neurogenic bladder sphincter dysfunction. *BJU Int*. 2001;**87**(6):575-80. [PubMed ID: 11298061]. <https://doi.org/10.1046/j.1464-410x.2001.00083.x>.
26. Chang SJ, Chiang IN, Hsieh CH, Lin CD, Yang SS. Age- and gender-specific nomograms for single and dual post-void residual urine in healthy children. *Neurourol Urodyn*. 2013;**32**(7):1014-8. [PubMed ID: 23595887]. <https://doi.org/10.1002/nau.22342>.
27. Herndon CD, Decambre M, McKenna PH. Interactive computer games for treatment of pelvic floor dysfunction. *J Urol*. 2001;**166**(5):1893-8. [PubMed ID: 11586256].
28. Bael A, Lax H, de Jong TP, Hoebeke P, Nijman RJ, Sixt R, et al. The relevance of urodynamic studies for Urge syndrome and dysfunctional voiding: a multicenter controlled trial in children. *J Urol*. 2008;**180**(4):1486-93. discussion 1494-5. [PubMed ID: 18710726]. <https://doi.org/10.1016/j.juro.2008.06.054>.
29. Kaefer M, Zurakowski D, Bauer SB, Retik AB, Peters CA, Atala A, et al. Estimating normal bladder capacity in children. *J Urol*. 1997;**158**(6):2261-4. [PubMed ID: 9366371]. [https://doi.org/10.1016/s0022-5347\(01\)68230-2](https://doi.org/10.1016/s0022-5347(01)68230-2).