A pilot study of bladder voiding with real-time MRI and computational fluid dynamics

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Abstract

Lower urinary tract symptoms (LUTS) affect many older adults. Multi-channel urodynamic studies provide information about bladder pressure and urinary flow but offer little insight into changes in bladder anatomy and detrusor muscle function. Here we present a novel method for real time MRI during bladder voiding. This was performed in a small cohort of healthy men and men with benign prostatic hyperplasia and lower urinary tract symptoms (BPH/LUTS) to demonstrate proof of principle; The MRI urodynamic protocol was successfully implemented, and bladder wall displacement and urine flow dynamics were calculated. Displacement analysis on healthy controls showed the greatest bladder wall displacement in the dome of the bladder while men with BPH/LUTS exhibited decreased and asymmetric bladder wall motion. Computational fluid dynamics of voiding showed men with BPH/LUTS had larger recirculation regions in the bladder. This study demonstrates the feasibility of performing MRI voiding studies and their potential to provide new insight into lower urinary tract function in health and disease.

Introduction

Lower urinary track symptoms (LUTS) and changes in bladder function occur frequently as individuals age [1–3]. Studies have evaluated the anatomical and functional changes of the bladder in patients with LUTS [4, 5]; however, the biomechanical characteristics of the lower urogenital tract, and how these are altered in patients with LUTS, are not fully understood. There is a compelling need to better delineate the anatomic and functional changes of the lower urinary tract in individual patients to improve diagnostic precision and allow for individualized treatment.

Lower urinary tract function is commonly assessed through multi-channel urodynamic studies that determine bladder pressure and flow during voiding. These studies can be performed in combination with fluoroscopic imaging to visualize the urine flow during voiding. However, these studies are invasive and provide little insight into the changes in bladder
anatomy and detrusor muscle function that occur with aging and lower urinary tract obstruction [6]. Use of non-invasive methods for the study of lower urinary tract anatomy and function has been limited. Image based patient specific computational models have been extensively used for cardiovascular evaluation and personalized treatment planning [7–10]. We envision a comparable approach for the evaluation of patients with LUTS. In this pilot study, we describe a magnetic resonance imaging (MRI) urodynamics method, as well as a patient specific MRI-based computational fluid dynamics (CFD) simulation of bladder voiding. These methods were applied to a small cohort of healthy men and men with benign prostatic hyperplasia and lower urinary tract symptoms (BPH/LUTS) to demonstrate proof of principle.

Materials and methods

This study was approved by the University of Wisconsin Health Sciences IRB (approval number 2017-1373-CP006). Three men with BPH/LUTS were recruited from the University of Wisconsin Urology clinic (ages 73, 71, and 54). The inclusion criteria were adult men recently diagnosed with BPH. Three control subjects were recruited from a database of healthy controls maintained by the MR research group at UW-Madison (ages 66, 42, and 44). Inclusion criteria were healthy adult men not experiencing any symptoms consistent with BPH. Exclusion criteria for both groups were contraindication to MRI (e.g. pacemaker, contraindicated metallic implants, claustrophobia, etc) and patients who have undergone prostatectomy. MRI was performed on a clinical 3T scanner (Premier, GE Healthcare, Waukesha, WI) using a high-density flexible surface coil array (AIR Coil, GE Healthcare). An MRI urodynamics protocol was implemented which involved a fluid challenge and voiding during MRI. Three-dimensional ‘Fast-spin echo’ (FSE) T2-weighted acquisitions were performed immediately before and after voiding. A sagittal plane 2D spoiled gradient echo (SGRE) dynamic real-time imaging (RTI) acquisition was performed during voiding. The MRI protocol timeline is shown in Fig 1. The bladder was segmented from pre- and post-voiding 3D images of the bladder, while bladder cross-sectional area change during voiding was calculated from the 2D RTI. Measurements from both the 3D and 2D images were incorporated in a patient specific simulation of bladder voiding. Further details on the fluid challenge, 3D FSE MRI, 2D RTI MRI, post-processing and computational fluid dynamics are in the subsequent sub-sections.

Fluid challenge

The subject was instructed to fast for 4 hrs. prior to arrival to the MRI session. Upon arrival, the subject was asked to empty his bladder and to drink approximately 1 L of a water and
electrolyte beverage (Gatorade G2, The Gatorade Company, Chicago, IL) 30 minutes before MRI scanning, and an additional 100 ml every 10 minutes until scanning (total of approximately 1.3 L). Before the imaging session started, the subject was equipped with a condom catheter system that allowed him to void supine in the scanner. The condom catheter was connected to a container through a flexible plastic tube.

3D MRI of the lower urinary tract

2D fast-spin echo (FSE) T2-weighted axial acquisitions are currently used for clinical evaluation of patients with bladder or prostate-related diseases [11, 12]. Even though these images provide relevant information for clinical diagnosis, they are limited by their slice thickness (~7 mm), demanding additional assumptions when performing volumetric measurements. Anatomical changes of the bladder can be spatially heterogeneous, requiring a volumetric (three-dimensional) evaluation. As an alternative, and in order to improve the spatial resolution for the current study, a single-slab 3D FSE T2-weighted imaging sequence was used [13]. This methodology utilizes parallel imaging–simultaneous acceleration in two directions–allowing the acquisition of additional and thinner slices, producing voxels with the same size in the slice and in-plane directions (isotropic resolution), contrary to conventional axial FSE acquisitions (anisotropic resolution, i.e.: slice thickness larger than in-plane pixel size).

Dynamic imaging of the bladder

Advanced MRI sequence development has allowed for dynamic real-time acquisition of data for anatomical and functional assessment. Dynamic anatomical MRI has been used extensively for heart function assessment, where balanced steady-state-free precession (bSSFP) images have been shown to provide sufficient contrast to separate the myocardium and the blood pool [14]. bSSFP has previously been used to determine the bladder motion during voiding in healthy volunteers [15]. We endeavored to reproduce this protocol and obtain bSSFP images in a healthy volunteer during voiding. However, the volunteer expressed significant abdominal discomfort due to peripheral nerve stimulation during the real time MR acquisition. This made this approach untenable for use in our study. To circumvent this road block, we used an MRI sequence that consists of a series of parallel sagittal 2D spoiled gradient echo (SGRE) dynamic real-time images capturing the bladder neck and urethra as well as other regions of the bladder. In this protocol, images are constantly acquired for four minutes allowing complete capture of voiding mechanics during bladder voiding without inducing any discomfort. Dynamic images were segmented to measure the relative displacement of the bladder wall during voiding. Only one control subject underwent the real-time imaging protocol. The other two control subjects were imaged prior to the real-time imaging being added to the IRB protocol and instead were asked to step out of the scanner and void in the bathroom after the 3D MRI was performed. Immediately after voiding the subject was asked to return to the MRI scanner and the 3D MRI was repeated.

MRI post-processing

Using 3D FSE T2-weighted images the bladder was segmented (Mimics, Materialise, Leuven, Belgium) for both pre and post voiding. The pre and post voiding bladder volumes were then exported as stereolithography (STL) files. From 2D RTI data during voiding, the area of a sagittal plane through the bladder was measured over time.

To estimate the motion of the bladder wall during voiding a spherical coordinate system was defined for the bladder, similar to previous motion estimation algorithms used in cardiac chambers [16, 17]. The coordinate system origin was set to be the center of the post voiding
bladder volume. In general, the bladder wall displacement \( \mathbf{d} \) is a three dimensional vector that has spatial and time dependence.

\[
\mathbf{d} = f(\theta, \phi, t)
\]  

(1)

Based on prior work in cardiac chambers, we simplify the complete description of bladder wall motion by assuming the bladder wall only moves radially \((d_\theta = d_\phi = 0)\) and the spatial and time dependence of the wall motion can be separated as

\[
d_r = d_y(\theta, \phi)\alpha(t)
\]  

(2)

where \(d_y(\theta, \phi)\) is the total displacement from the pre to post void anatomies and \(\alpha(t)\) is the time dependence function that varies from 0 at the start of voiding to 1 at the end of voiding. For wall displacement analysis, the bladder wall was divided into anterior-posterior, dome-base, and left-right regions (Fig 2) and an asymmetry ratio was calculated based on the difference between the median displacement of the left and right bladder wall regions.

For each point on the bladder surface \(d_0\) the distance between the pre and post voiding anatomies was calculated using a fast, minimum storage ray-triangle intersection algorithm [18] implemented in MATLAB. The time dependence function \(\alpha(t)\) can be calculated from real time measurements of bladder area during voiding,

\[
\alpha(t) = \left( \frac{A(t) - A(t_0)}{A(t_{end}) - A(t_0)} \right)^{1/2}
\]  

(3)

where \(A(t)\) is the bladder area, \(t_0\) is the time at the start of voiding and \(t_{end}\) is the time at the end of voiding. Bladder area measurements from the real time sagittal MR images showed a sigmoidal behavior (Fig 3). Based on that behavior, \(\alpha(t)\) was chosen to be a square root of cosine function.

**Computational fluid dynamics**

The patient-specific bladder anatomies were imported into the computational fluid dynamics (CFD) software CONVERGE v2.4 (Convergent Science Inc, Madison, WI). Bladder wall motion was estimated as described above and imposed with a user-defined function to virtually drive voiding. As only one control subject had real-time MRI the same \(\alpha(t)\) function was used for all three control subjects. The urethra wall was assumed to be rigid and the urethra outlet was set to atmospheric pressure. CFD simulations with large boundary motion are typically time-consuming, but this study employed two complementary strategies to enhance simulation speed. First, a cut-cell Cartesian mesh was used as these meshes have advantages in biomedical flows with moving boundaries [19]. Compared to typical boundary fitted meshes, cut-cell grids rapidly re-mesh to handle the moving bladder wall while simultaneously minimizing numerical diffusion from mesh motion. Second, the efficient re-meshing capabilities of cut-cell Cartesian meshes were used by adapting the mesh to the instantaneous flow field, which minimizes simulation time while maintaining accuracy. Results figures were generated with Tecplot 360 (Tecplot, Bellevue, WA). Vorticity, the curl of the velocity field, was averaged over the bladder volume (urethra not included). Dimensionless vorticity was then calculated based on average urethra flow rates and the prostatic urethra diameter for each subject.

**Results**

The MRI urodynamics protocol was successfully completed in three healthy volunteers and three patients with BPH/LUTS. 2D planes of the bladder during voiding are shown in Fig 3,
along with the bladder cross-section area over time. The rate of area change is slowest at the beginning and end of voiding and greatest during the middle of voiding. Three-dimensional plots of the estimated bladder displacement throughout voiding at various time points are shown in Fig 3. The estimated displacement maps show that greatest displacement occurs at the dome of the bladder.

MRI urodynamics was performed on two additional healthy volunteers and three men with BPH/LUTS. The pre and post-voiding anatomies, estimated displacement maps and box plots showing the regional displacement behavior for each subject are shown in Fig 4. The control subjects had large displacements at the bladder dome with little observed asymmetry. The men with BPH/LUTS had smaller displacements and unlike the controls did not exhibit a consistent displacement pattern. These qualitative observations from the displacement maps were confirmed looking at probability functions of the displacement and the left-right asymmetry (Fig 5). Overall the bladder walls of men with BPH/LUTS moved only 25%-50% as much as the

Fig 2. Schematic of the bladder wall divided into anterior-posterior, dome-base, and left-right regions for regional displacement and asymmetry analysis. Different colors represent different regions (anterior–aqua, posterior–yellow, dome–navy blue, base–royal blue, left–red, right–not shown).

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control subjects. The control subjects had little left-right asymmetry (4%–14%) while the BPH patients had large left-right asymmetry (40%–160%).

Urodynamics results from computational fluid dynamics (CFD) are shown in Fig 6 for a sagittal plane near the center of the bladder. Control subjects had higher urine velocities in the
bladder than men with BPH/LUTS due to their greater bladder wall displacements. Near the
initiation of voiding, streamlines (showing the direction of urine flow) for all three control
subjects and two men with BPH/LUTS (P1 and P3) were directed toward the bladder neck and
prostatic urethra. Patient P2 has a small vortex just above the urethra. Towards the end of the
voiding process, control subjects C1 and C3 had small recirculation regions by the anterior
bladder wall while the streamlines for C2 were still all directed toward the bladder neck and
prostatic urethra. Near the end of voiding patient P1 had a small posterior recirculation region,
P2 had a large anterior recirculation region and P3 had large anterior and posterior recircula-
tion regions. Despite the men with BPH/LUTS having larger recirculation regions, they had
lower average vorticity (Fig 7) in the bladder due to their slower flow rates and smaller veloci-
ties. After making vorticity dimensionless (a normalization to account for differing flow rates
between patients), vorticity was similar between healthy controls and men with BPH/LUTS.

Discussion
This study demonstrates the feasibility of MRI bladder voiding studies to provide new insight
into lower urinary tract function in health and disease, and to generate patient-specific simula-
tions of voiding. Additionally, our studies have revealed a previously unsuspected aspect of
bladder voiding. It has been generally assumed that the sphere-shaped full bladder contracted
with relatively uniform displacement. Yet, the results of this study suggest that this is not the
case: Displacement analysis of voiding in healthy controls revealed a much greater displace-
ment of the bladder dome than the other regions of the bladder (Fig 3). This is a simple, but
profoundly startling, observation that may influence understanding of the contractile function
of the bladder during voiding after further study. There is increasing appreciation for the role
of impaired bladder contractility and its effects on treatment outcomes [20–24]. All of the men
with BPH/LUTS in this study exhibited decreased bladder contractility on traditional multi-
channel urodynamic evaluation (data not shown). The limitation of standard urodynamic
evaluation is that it provides only an indirect assessment of bladder contractility calculated
from voiding pressure and flow at a single point in the voiding effort (maximum flow). MRI
urodynamics, on the other hand, reveals significantly decreased and asymmetric bladder wall
motion during voiding.
Although simulations of cardiac flow have incorporated wall motion going back to the 1970s [25, 26], this is to our knowledge the first MRI urodynamic-based simulation of bladder wall motion during voiding. Prior CFD studies have set the bladder wall to be an inlet [27, 28] or used the ureters to drive voiding [29]. The CFD methodology presented here represents a significant step towards improving the physical and physiological realism of urodynamic

Fig 6. CFD results showing velocity contours and streamlines on a sagittal plane at the center of the bladder for each subject. Time points are displayed both near the initiation and termination voiding.  
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Fig 7. Bladder vorticity was lower for the men with BPH/LUTS due to their lower flowrates and urine velocities. After making vorticity dimensionless (a normalization to account for different flow rates between subjects), vorticity was similar for both groups.  
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simulations. As expected based on smaller bladder wall displacements, our simulation results showed that men with BPH/LUTS had lower urine velocities. More interestingly, the men with BPH/LUTS had larger recirculation regions in the bladder which could increase the energy demands of voiding.

In summary, this study demonstrated the feasibility of MRI bladder voiding studies to non-invasively investigate bladder function. Results from displacement analysis showed men with BPH/LUTS had decreased and asymmetric bladder wall motion compared to healthy male controls and fluid dynamic analysis of voiding showed them to have larger recirculation regions in the bladder.

Supporting information
S1 Data.
(XLSX)

Author Contributions

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References

1. Fananapazir G, Kitich A, Lamba R, Stewart SL, Corwin MT. Normal reference values for bladder wall thickness on CT in a healthy population. Abdom Radiol. 2018; 43: 2442–2445. https://doi.org/10.1007/s00261-018-1463-x PMID: 29392368
2. Güzel Ö, Aslan Y, Balci M, Tunçel A, Keten T, Erkan A, et al. Can Bladder Wall Thickness Measurement Be Used for Detecting Bladder Outlet Obstruction? Urology. 2015; 86: 439–444. https://doi.org/10.1016/j.urology.2015.06.023 PMID: 26142716
3. Wei JT, Calhoun E, Jacobsen SJ. Urologic diseases in America project: Benign prostatic hyperplasia. J Urol. 2005; 173: 1256–1261. https://doi.org/10.1097/01.ju.0000156709.37840.fe PMID: 15798764
4. Mirone V, Imbimbo C, Longo N, Fusco F. The Detrusor Muscle: An Innocent Victim of Bladder Outlet Obstruction[A figure is presented]. European Urology. Elsevier; 2007. pp. 57–66. https://doi.org/10.1016/j.eururo.2006.07.050 PMID: 16979287
5. Hou P, De EJB, Kramer LA, Westney OL. Dynamic contrast-enhanced MRI study of male pelvic perfusion at 3T: Preliminary clinical report. J Magn Reson Imaging. 2007; 25: 160–169. https://doi.org/10.1002/jmri.20813 PMID: 17173309
6. Bushman W. Etiology, Epidemiology, and Natural History. Urologic Clinics of North America. Elsevier; 2009. pp. 403–415. https://doi.org/10.1016/j.ucl.2009.07.003 PMID: 19942041
7. Marsden AL, Feinstein JA. Computational modeling and engineering in pediatric and congenital heart disease. Curr Opin Pediatr. 2015; 27: 587–596. https://doi.org/10.1097/MOP.0000000000000269 PMID: 26262579
8. Baijargeon B, Rebeco N, Fox DD, Taylor RL, Kuhl E. The living heart project: A robust and integrative simulator for human heart function. Eur J Mech A/Solids. 2014; 48: 38–47. https://doi.org/10.1016/j.euromechsol.2014.04.001 PMID: 25267880

9. Taylor CA, Steinman DA. Image-based modeling of blood flow and vessel wall dynamics: applications, methods and future directions: Sixth International Bio-Fluid Mechanics Symposium and Workshop, March 28–30, 2008 Pasadena, California. Ann Biomed Eng. 2010; 38: 1188–1203. https://doi.org/10.1007/s10439-010-9901-0 PMID: 20087775

10. Taylor CA, Figueroa CA. Patient-specific modeling of cardiovascular mechanics. Annu Rev Biomed Eng. 2009; 11: 109–134. https://doi.org/10.1146/annurev.bioeng.061807.160521 PMID: 19400706

11. Guneyli S, Ward E, Peng Y, Nehal Yousuf A, Trilisky I, Westin C, et al. MRI evaluation of benign prostatic hyperplasia: Correlation with international prostate symptom score. J Magn Reson Imaging. 2017; 45: 917–925. https://doi.org/10.1002/jmri.25418 PMID: 27487205

12. Wasserman NF, Spilseth B, Golzarian J, Metzger GJ. Use of MRI for Lobar Classification of Benign Prostatic Hyperplasia: Potential Phenotypic Biomarkers for Research on Treatment Strategies. Am J Roentgenol. 2015; 205: 564–571. https://doi.org/10.2214/AJR.14.13602 PMID: 2695642

13. Busse R. 3D FSE Reduces Scan Time, Generates Thinner Slices. 2007.

14. Hocaoglu Y, Roosen A, Herrmann K, Tritschler S, Stief C, Bauer RM. Real-time magnetic resonance imaging (MRI): anatomical changes during physiological voiding in men. BJU Int. 2012; 109: 234–239. https://doi.org/10.1111/j.1464-410X.2011.10255.x PMID: 21736694

15. Lai CQ, Lim GL, Jamil M, Mattar CNZ, Biswas A, Yap CH. Fluid mechanics of blood flow in human fetal left ventricles based on patient-specific 4D ultrasound scans. Biomech Model Mechanobiol. 2016; 15: 1159–1172. https://doi.org/10.1007/s10237-015-0750-5 PMID: 26676944

16. Vasudevan V, Low AJJ, Annamalai SP, Sampath S, Poh KK, Totman T, et al. Flow dynamics and energy efficiency of flow in the left ventricle during myocardial infarction. Biomech Model Mechanobiol. 2017; 16: 1503–1517. https://doi.org/10.1007/s10237-017-0902-x PMID: 28364199

17. Moller T, Trumbore B. Fast, Minimum Storage Ray-Triangle Intersection. J Graph Tools. 1997; 2: 21–28.

18. Marom G. Numerical Methods for Fluid–Structure Interaction Models Numerical Methods for Fluid-Structure Interaction Models of Aortic Valves. Arch Comput Methods Eng, 2015; 22: 595–620. https://doi.org/10.1007/s10237-015-0750-5 PMID: 26676944

19. Aldamanhor R, Chapple CR. Underactive bladder, detrusor underactivity, definition, symptoms, epidemiology, etiopathogenesis, and risk factors. Current Opinion in Urology. Lippincott Williams and Wilkins; 2017. pp. 293–299. https://doi.org/10.1097/MOU.0000000000000381 PMID: 28221218

20. Bosch R, Abrams P, Averbeck MA, Finazzi Agró E, Gammie A, Marcelissen T, et al. Do functional changes occur in the bladder due to bladder outlet obstruction? - ICI-RS 2018. Neurour ol Urodyn. 2019; 38: S56–S65. https://doi.org/10.1002/nau.24076 PMID: 31278801

21. Chueh YC, Plata M, Lamb LE, Chancellor MB. Underactive Bladder in Older Adults. Clinics in Geriatric Medicine. W.B. Saunders; 2015. pp. 523–533. https://doi.org/10.1016/j.cger.2015.06.002 PMID: 26476113

22. Hartigan SM, Reynolds WS, Dmochowski RR. Detrusor underactivity in women: A current understanding. Neurourology and Urodynamics. 2019; 38: 2070–2076. https://doi.org/10.1002/nau.24174 PMID: 31432566

23. Osman NI, Chapple CR, Abrams P, Dmochowski R, Haab F, Nitti V, et al. Detrusor underactivity and the underactive bladder: A new clinical entity? a review of current terminology, definitions, epidemiology, aetiology, and diagnosis. European Urology. Elsevier; 2014; 38: S6–S65. https://doi.org/10.1016/j.euro.2013.10.015 PMID: 24184024

24. Peskin CS. Flow patterns around heart valves: A numerical method. J Comput Phys. 1972; 10: 252–271. https://doi.org/10.1016/0021-9991(72)90065-4

25. Peskin CS. Numerical Analysis of Blood Flow in the Heart. J Comput Phys. 1977.

26. Zheng J, Pan J, Qin Y, Huang J, Luo Y, Gao X, et al. Role for intravesical prostatic protrusion in lower urinary tract symptom: A fluid structural interaction analysis study. BMC Urol. 2015; 15: 1503–1517. https://doi.org/10.1111/j.1464-410X.2011.10255.x PMID: 26676944

27. Altesci YZ, Senturk U, Pekedis M, Cinar M. Non-Invasive Urodynamic Analysis Using the Computational Fluid Dynamics Method Based on MR Images. Turkije Klin J Med Sci. 2011; 31: 1186–93. https://doi.org/10.5336/medsci.2010-21420

28. Jin Q, Zhang X, Li X, Wang J. Dynamics analysis of bladder-urethra system based on CFD. Front Mech Eng Chinca. 2010; 5: 336–340. https://doi.org/10.1007/s11465-010-0027-8