Prostate and Irregular Afferent-Triggered Reflex Responses

Voiding Dysfunction in Old Male Rats Associated With Enlarged Prostate and Irregular Afferent-Triggered Reflex Responses

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Purpose: This study was conducted to evaluate the hypothesis that an enlarged prostate in old rats may lead to complications associated with voiding dysfunction involving ionotropic P2X2/3-type purinergic receptors.

Methods: Intact animals were divided into male young (MYR; 8–10 weeks old) and male old (MOR; 20 months old) rats. The animals underwent simultaneous detrusor electromyography (EMG) and suprapubic cystometry (CMG) under urethane anesthesia. Immunofluorescence techniques were used to evaluate prostatic autonomic innervation and P2X3R expression in bladder urothelial cells. The functional role of P2X3R was characterized by intramuscular application of AF-353, a selective P2X2/3R antagonist.

Results: The prostate index significantly increased in MOR, suggestive of an enlarged prostate affecting micturition patterns. Significant EMG and CMG differences were found between MYR and MOR. Higher immunoreactivity for P2X2/3R in the urothelial layer and for prostatic neurofilaments was seen in MOR. Systemic inhibition of P2X2/3R had minimal effects on MYR responsiveness, but improved voiding function in MOR with a marked decrease of intravesical pressure and bladder contractile responses.

Conclusions: The data support the hypothesis that an enlarged prostate in MOR may contribute to voiding dysfunction involving activation of P2X2/3R, which enhances a prostate-bladder reflex. This reflex may increase bladder afferent transmission and activation of increased prostate innervation, leading to voiding dysfunction.

Keywords: Prostate; Urinary bladder; Urothelium; Receptors, Purinergic P2X3; Micturition

*Research Ethics: All experimental procedures were approved by the Institutional Animal Care and Use Committee of Houston Methodist Research Institute (IACUC; ID: AUP-0615-0044/IS00001252; responsible investigator: Alvaro Munoz) and performed in accordance with the Guidelines from the National Institutes of Health on the Care and Use of Laboratory Animals.

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HIGHLIGHTS
- Immunoreexpression of both P2X3 receptors in bladder-urothelium and neurofilaments in prostate tissue from aged male rats is higher than in young animals.
- Experiments performing simultaneous electromyography of the BLADDER DOME with bladder cystometry in anesthetized rats, uncovered clear dysfunctional voiding-patterns in aged male animals, which are improved by systemic inhibition of P2X3 receptors.
- Aging-related changes in the lower urinary tract and prostate from aged male-rats may contribute to voiding dysfunction, involving a hyperexcitable prostate-bladder reflex via activation of urothelial P2X3 receptors.
INTRODUCTION

Lower urinary tract symptoms generate elevated financial burdens and healthcare costs in older men [1,2]. One of the most common symptoms is difficulty urinating due to narrowing of the urethral lumen associated with an enlarged prostate, otherwise known as benign prostatic hyperplasia. This condition is not associated with prostate cancer, but generates similar symptoms of bladder outlet obstruction [2].

Although the pathophysiology of voiding dysfunction associated with prostatic hyperplasia is not clearly understood, several pharmacological treatments are available to modestly improve the quality of life in patients with benign prostatic hyperplasia. Common therapies include the use of phosphodiesterase inhibitors, alpha-adrenergic blockers, or 5-alpha reductase inhibitors, but these treatments do not dramatically improve urinary symptoms [3]. Meanwhile, few basic research studies have correlated histological findings with functional evaluations, making it even more difficult to interpret pharmacological effects and targets.

Consequently, more comprehensive preclinical investigations using animal models to evaluate the effects of aging on pathological alterations of the lower urinary tract are necessary. Delineating a mechanistic understanding of voiding dysfunction in the older may be helpful for making clinical diagnoses and selecting better therapeutic interventions.

Several animal models have been used to evaluate bladder outlet obstruction in rats [4]. Some of these involve the mechanically induced partial occlusion of the urethra in rats to simulate similar conditions as those observed in patients with prostate-induced urethral obstructions [5]. Despite the efforts made to evaluate the impact of aging on micturition, the reported results have varied greatly, which may be explained by differences in species, sex, the application of anesthetics, and the experimental procedures used [6].

Thus, to better understand the neurophysiological pathways involved in voiding dysfunction associated with an enlarged prostate, we performed simultaneous detrusor electromyographic and cystometric recordings in aged male rats (20 months old), and compared the responses to those in young animals (8-10 weeks old). This approach was intended to establish naturally occurring benign prostatic hyperplasia in old rats to yield further insights into the origin of the complications associated with voiding dysfunction.

Ionotropic P2X3 and P2X2/3 purinergic receptors (P2X3R) are hetero- or homo-trimer protein complexes that function as nonselective ion channels activated by adenosine triphosphate (ATP), and participate in afferent transmission of sensory stimuli in different tissues, including the lower urinary tract [7,8]. Due to the relevance of these receptors for nociception, various antagonists have been designed for pharmacological use in diseases related to pain, including AF-353, an antagonist able to block P2X3 and P2X2/3 receptors in the periphery and the central nervous system [9]. Because the bladder sensory symptoms associated with prostate hyperplasia are carried by afferent signals [10], some of them mediated by P2X2/3R [11-13], we pharmacologically tested the role of ionotropic P2X3R [9] in both groups of rats.

MATERIALS AND METHODS

All experimental procedures were approved by our Institutional Animal Care and Use Committee, and performed in accordance with the Guidelines from the National Institutes of Health on the Care and Use of Laboratory Animals. Rats were housed in a pathogen-free environment, with 12-hour light/dark cycles, a controlled room temperature of 25°C, and had ad libitum access to food and water in plastic cages containing corn cob bedding. Efforts were made to minimize the number of animals used, while maintaining the ability to make significant comparisons.

Animals and Surgical Preparation

Long-Evans rats were divided into groups of male young rats (MYR; 2–2.5 months old; N = 6) or male old rats (MOR; 20 months old; N = 6). The MOR animals were donated through the Resources and Tissue-Sharing Initiative at our institution, while maintaining the ability to make significant comparisons.
out the surgical procedures and experiments.

Cystometry and Electromyography
The status of both the electrode and suprapubic catheter was verified with a stereomicroscope before placing animals in a prone position. Voided urine was collected with a plastic cup mounted on top of a pressure sensor for monitoring effective volumes. Saline infusion was performed using a syringe pump through the suprapubic catheter, at a rate of 0.1 mL/min. Before the start of the experiment, continuous saline infusion was performed for approximately 10-15 minutes until stable voiding events, indicated by real-time cystometry (CMG), were observed. After 6 consecutive voiding events, 10 mg/kg AF-353 (diluted at a final dimethyl sulfoxide concentration of 0.01% in saline solution) was injected intramuscularly into the right quadriceps muscle. This dose was selected based on previous studies in which the P2X3R antagonist was administered intravenously [14]. Based on the biological characterization of AF-353, for which a time to peak plasma concentration of approximately 30 minutes was determined when orally administered [11], we waited for a period of 20 minutes before 6 additional consecutive voiding events were recorded. Electromyography (EMG) signals were amplified with a DP-304 differential amplifier (Warner Instruments, Hamden, CT, USA), acquired at a sampling rate of 2 kHz, and digitized using the WinDaq data acquisition software (DataQ Inst. Akron, OH, USA).

Cystometric Analysis
In contrast to typical cystometric responses, the Long-Evans male rats used in this study showed a far more complicated voiding pattern that involved the presence of nonvoiding contractions (NVCs), small volume voiding events (SVVEs), and effective contractions (ECs) during sustained high intravesical pressures before returning to baseline bladder pressure, particularly in MOR animals (;). Nonetheless, for CMG characterization, only the last 3 voiding events, indicated by visible intraluminal pressure high-frequency oscillations (IPHFOs) at each stage (intact and following intramuscular AF-353 injection) were used. NVCs were considered to occur when a change in bladder pressure higher than 5 cm H$_2$O was observed, accompanied by no evident release of saline, either at baseline or at high intravesical pressures. The frequency was calculated as the number of NVCs over the duration of the recording condition and normalized by minute (i.e., NVC/min). Additionally, the peak pressure during bladder contractions (cm H$_2$O), intercontraction interval (sec), duration of IPHFOs (sec), and voided volume (mL) for ECs were determined using MATLAB R2017a (MathWorks, Natick, MA, USA).

Electromyographic Analysis
Second-order Butterworth notch filters were applied to remove the power line interference and harmonics for all electrophysiological signals. EMG signals were further bandpass-filtered at 5–400 Hz using a second-order Butterworth filter. A template subtraction method was applied to remove the electrocardiogram contamination. Epochs of bladder EMG discharging were automatically detected using a preset amplitude threshold. The onset/offset of EMG activity was temporally referenced to the bladder pressure peak. The bladder pressure that induced the onset/offset of signal activation was also determined. For each EMG experiment, the signal turns (indicative of frequency), amplitude, and duration were calculated in triplicate, averaged, and grouped according to the age condition.

Morphometric Analysis
After performing CO$_2$-thoracothomy euthanasia, the ventral, lateral, and dorsal prostate lobes were removed from the MYR and MOR animals and wet-weighed as a whole, and the resulting weight values were recorded. The same procedure was performed for the urinary bladders including tissue from the bladder base to the bladder dome. Thereafter, the bladder and prostate (ventral + lateral + dorsal lobes) indexes were calculated respectively as the ratio of bladder or prostate weight per 100-g body weight.

Immunofluorescent Procedure in Bladder Tissue
Following systemic treatment with the P2X3R antagonist during simultaneous CMG-EMG evaluations, urinary bladders were processed and stained as previously described [15]. Briefly, equatorial bladder specimens from each animal were incubated in 4% formalin overnight and cryoprotected in 30% sucrose at 4°C. Thereafter, specimens were sectioned at 30 µm with a cryostat, and blocked overnight at 4°C (10% goat serum, 2% bovine serum albumin, 0.1% Triton X-100, and 0.05% Tween 20). After washout, bladder sections were incubated overnight with antibodies against-actin (1:400; AM1829B, ABGENT, San Diego, CA, USA) and P2X3R receptors (1:1,000; AB90905, AB-CAM, Cambridge, MA, USA). After washing, sections were incubated in the dark for 2 hours at room temperature with a mixture of donkey anti-rabbit (Alexa Fluor 647; AB150075, ABspin, San Diego, CA, USA) and donkey anti-mouse (Alexa Fluor 488; AB150752, ABspin). Slides were coverslipped with fluorescent mounting medium (DakoCytomation, Carpinteria, CA, USA) and images were acquired using a Leica confocal microscope (Wetzlar, Germany) with a 20× objective.
ABCAM) and goat anti-mouse (Alexa Fluor 488, AB150113, ABCAM) secondary antibodies. Sections were washed, covered with mounting medium containing 4′-6-diamidino-2-phenylindole (DAPI) (ab104139, ABCAM), and protected with glass coverslips.

**Immunofluorescent Procedure in Prostatic Tissue**

A section of the ventral prostate lobe was fixed in formalin and then cryoprotected in 30% sucrose. Briefly, prostatic tissue was sectioned at 30-μm thickness and incubated overnight at 4°C with a mouse anti-neurofilaments 68 and 200 kDa antibody (1:100; AB1989, Sigma-Aldrich) and β-actin (1:400). Secondary antibody incubation was performed as described for bladder tissues. Sections were covered with mounting medium containing DAPI and protected with glass coverslips. The antibody solutions were simultaneously used for all bladder and prostate slides.

**Immunofluorescent Evaluations**

A dry ×40 objective on a Leica TCS SPE confocal microscope (Leica Microsystems, Buffalo Grove, IL, USA) was used for acquiring images, which were digitized using Leica LAS-X interface software. The image analysis was performed with ImageJ software (U.S. National Institutes of Health, Bethesda, MD, USA) by determining the signal contribution (as percent) for each channel (i.e., DAPI, β-actin, neurofilaments, and P2X3R) over the full image area. Additionally, the P2X3R signals were normalized to DAPI and β-actin as immunoreactivity indicators for each bladder.

**Statistical Analysis**

Intergroup statistical comparisons between the MOR and MYR groups were performed using 1-way analysis of variance with multiple comparisons between means, and corrected with the Tukey post hoc test. Intragroup analyses were performed with the 2-tailed unpaired t-test. All values are expressed as mean ± standard error. In all cases, the level of statistical significance was set at P < 0.05. All statistical analyses were performed with GraphPad 6.0 software (Prism, GraphPad Software Inc., San Diego, CA, USA).

**RESULTS**

**Morphometric Differences Between MYR and MOR**

As expected, the average body weight of MYR (289.2 ± 7 g) was significantly lower than that of MOR (594.5 ± 7 g) (P < 0.001). Similarly, the weights of the isolated bladder and prostate tissues were significantly higher in MOR (Fig 1A, B). However, when compared to the MYR animals, a lower bladder index (Fig. 1C) and a higher prostate index were observed in MOR, indicative of an enlarged prostate (Fig. 1D).

**Higher Expression of Urothelial P2X3R in MOR**

Both MYR and MOR showed similar levels of DAPI signals (indicative of a comparable numbers of cells), and urothelial immunoreactivity for β-actin and P2X3R (Fig. 2A–D, MYR; Fig. 2E–H, MOR). In an analysis of the total contribution of DAPI, β-actin, and P2X3R to the total signal area, higher values were found for P2X3R in MOR than in MYR (Fig. 2I; P < 0.01). When the P2X3R signal contribution was normalized to either DAPI (P < 0.05, MOR vs. MYR) or β-actin (P < 0.01, MOR vs. MYR), a greater index remained for urothelial P2X3R in MOR than in MYR (Fig. 2J).

**Higher Expression of Neurofilaments in Ventral Prostate Lobes From MOR**

Immunoreactivity for β-actin and neurofilaments, as well as DAPI staining, was observed in the prostatic ventral lobes from both MYR (Fig. 3A–D) and MOR (Fig. 3E–H). A group analysis of these 3 markers showed a significantly higher expression of neurofilaments in MOR prostatic tissues (Fig. 3I) (P < 0.05 vs. MYR).
Micturition in MYR Before and After Systemic Inhibition of P2X3R

Simultaneous CMG/EMG demonstrated typical micturition patterns in MYR during intravesical saline infusion. Fig. 4A shows multiple micturitions where EMG was correlated with voiding events. A closer view at one of these events corroborates muscle activation (Fig. 4B) and maximal activity during IPHFO (Fig. 4C). The intramuscular application of AF-353 (10 mg/kg) to the same rats increased the number of micturitions in the same period of recording time (Fig. 4D), but decreased the maximal voiding pressure with a parallel reduction in EMG activity (Fig. 4E). The correlation of EMG with IPHFOs remained the same despite a decrease in IPHFO after P2X3R inhibition. No evident NVCs were observed.

Micturition in MOR Before and After Systemic Inhibition of P2X3R

In contrast to MYR, the simultaneous EMG/CMG recordings in MOR showed significantly altered voiding patterns (Fig. 5A, B). These were characterized by a mixture of multiple NVCs (indicated with * in Fig. 5C), SVVE, and ECs (Fig. 5C), all of them accompanied by some degree of EMG activity; these abnormal activities were observed during high intravesical pressures, but were more evident during IPHFOs (Fig. 5A-C). The systemic inhibition of P2X3R in the same MOR animals substantially affected both EMG and CMG responses. First, an in-
crease in the intercontractile interval was observed (Fig. 5D). Second, the frequencies of NVCs and SVVE were reduced (Fig. 5E). Finally, a more typical voiding event was observed in MOR following AF-353 application (Fig. 5F).

**CMG and EMG Group Analysis in MYR and MOR**

A group analysis showed a higher frequency of EMG spikes in MOR than in MYR (P < 0.001 vs. MYR), and a reduction after P2X3R inhibition (P < 0.01 vs. MOR) (Fig. 6A). The amplitude of the spikes was also reduced in MOR (P < 0.01 vs. MYR) and not altered by AF-353 application (Fig. 6B). The duration of the EMG signal was significantly higher in MOR (P < 0.001 vs. MYR) and only affected by the purinergic antagonist in MOR (P < 0.05) (Fig. 6C). Although a trend was observed towards reduced peak bladder pressure, no significant differences were found between MYR or MOR before or after P2X3R inhibition (Fig. 6D). The intercontractile interval was lower in MOR (P < 0.05 vs. MYR); it decreased in response to AF-353 in MYR (P < 0.05) but increased in MOR (P < 0.01) (Fig. 6E). The duration of IPHFOs was significantly lower in MOR than in MYR (P < 0.05), and no effects were induced by the P2X3R antagonist in either group (Fig. 6F). Similar responses between groups were determined for voided volume (Fig. 6G). Finally, the frequency of NVCs was significantly higher in MOR (P < 0.05), and no effects were induced by the P2X3R antagonist in either group (Fig. 6H).

**Fig. 3.** Neurofilament expression was higher in male old rats. Representative images of immunofluorescent signals in the prostate from 4’-6-diamidino-2-phenylindole (DAPI), β-actin (β-ACT), neurofilaments (NFs), and their merged signals from MYR (A–D) or MOR (E–H). (I) Group analysis of the contributions of DAPI (left), β-actin (middle), and neurofilaments (right) to the total signal area in prostate sections from MYR and MOR. Prostates were dissected after systemic treatment with AF-353 following CMG-EMG characterization. The scale bar in panel A indicates 50 µm, and applies to all images. *P < 0.05 vs. the MYR group (t-test). MYR, male young rats; MOR, male old rats; CMG, cystometry; EMG, electromyography.
DISCUSSION

The present study showed a significantly higher prostate-to-body weight index in aged rats, suggesting that age-related conditions may induce a partial urethral obstruction affecting urination due to complicated voiding patterns in MOR. This observation suggests that it is possible to avoid chemically induced changes in prostate size [16,17]. The immunofluorescence results also revealed an increased presence of neurofilaments in MOR, supporting the possibility that pathological conditions stimulate prostatic innervation and consequently the overgrowth of the gland [18]. Additionally, the expression of P2X3R involved in the amplification of purinergic sensory signals from the bladder was also increased in the urothelium from MOR, suggesting an increased afferent tone mediated by the release of ATP [19]. As both MYR and MOR animals were treated in the same way with AF-353, we consider that this difference in the immunoreactivity of the receptor and neurofilaments did not result from exposure to the antagonist.

The current MOR model may be considered as analogous to benign prostatic hyperplasia in older humans, which may alter the normal micturition process. Prostatic hyperplasia is closely related to bladder outlet obstruction and urinary infection in males [6], although its etiology has not yet been well established.

Fig. 4. Systemic inhibition of P2X3R had mild effects on cystometric and electromyographic bladder responses from young male rats. (A) Representative multiple voiding events during simultaneous cystometry and bladder electromyography in control conditions. (B) Single voiding event as indicated in panel A. (C) Example of IPHFOs during the single voiding event indicated in panel B. (D) Representative pattern of the response during simultaneous cystometry and bladder electromyography after intramuscular application of 10-mg/kg AF-353 in the same rat. (E) Single voiding event as indicated in panel D. (F) Characteristic IPHFOs during the single voiding event indicated in panel E after AF-353 injection. IPHFO, intraluminal pressure high-frequency oscillation; EMG, electromyography.
Although direct urethral urine flow rate was not evaluated in this study, the CMG responses suggest a dysfunctional micturition process in older animals, which may be associated with obstructive uropathy caused by a large prostate, increased prostate innervation, and enhanced afferent signals from the bladder.

The voiding patterns observed in MOR differ from those of previous murine studies in female rats under anesthesia [21], but are somewhat consistent with cystometric findings in conscious male mice [22]. This may be explained by differences in animal models and sex [6]. Nonetheless, our results suggest that a correlation between an enlarged prostate and increased expression of urothelial P2X3R may explain cystometric differences observed in MOR, and that this condition may be associated with afferent hyperexcitability. Supporting this suggestion, it has been proposed that a prostate-to-bladder reflex, which is neurally mediated, may participate in regulating the micturition process [23]. Therefore, a reciprocal reflex involving the prostate and the bladder may be responsible for the impaired voiding function in MOR, and may be further related to the increased expression of P2X3R-mediated receptors in the bladder, promoting a higher expression of neurofilaments in the prostate. Our results support this hypothesis by showing the coexist-
tence of prostate enlargement, increased prostate innervation, and bladder hyperactivity in MOR. Indeed, in a recent study using a formalin-induced prostatic inflammation model in rats, hyperexcitability of afferent neurons combined with bladder overactivity was observed [16], supporting the idea of more complex interactions among the prostate, the urinary bladder,
and afferent innervation and the possibility that a localized reflex may control both organs.

In this regard, ionotropic P2X3R have been implicated as key mediators of sensory transmission in the lower urinary tract [24]. Previous reports on lower urinary tract dysfunction have found increased immunoreactivity and expression of bladder P2X3R [12,25]. Therefore, the observed bladder dysfunction in MOR may hypothetically also be associated with the increased expression of purinergic P2X3R in the bladder urothelium, as these receptors mainly function as mediators of bladder afferent activity following ATP release [11]. The increased urothelial expression of P2X3R in old rats may directly lead to an increased sensory tone, which is closely related to bladder overactivity. Furthermore, the current results also coincide with previous findings in male mice [12], where increased P2X3R expression, along with concurrent transmitter overflow of ATP and acetylcholine, was found in the bladder lumen. Our findings suggest that these changes in parasympathetic neurotransmitter release may also occur in MOR, and may be associated with the prostate-to-bladder reflex in a hyperexcitable mode.

As P2X3R in bladder sensory fibers and urothelial cells trigger the activation of afferent pathways [26], which may be a factor contributing to the voiding dysfunction observed in MOR, P2X2/3 receptors were pharmacologically evaluated by systemic application of AF-353, a selective antagonist [9]. Although we did not measure nerve electrical activity, our results suggest that intramuscular injections of AF-353 in MOR resulted in marked improvements in voiding function, with significantly decreased detrusor EMG frequency and duration. Although the impact of the purinergic antagonist on EMG findings in MYR was negligible, its stronger impact on MOR can be well explained by the attenuation of enhanced purinergic signaling mediated by urothelial P2X3R, triggering the urothelial release of ATP [27] to enhance the activation of bladder afferent innervation in MOR animals.

These results suggest that pharmacological inhibition of P2X3R may be a promising therapeutic alternative for treating age-related bladder dysfunction. Although neither EMG nor CMG properties were restored to those observed in MYR, the overall number of nonvoiding events, indicative of bladder overactivity, improved in MOR. One reason for these differences may be that the intramuscular injections of AF-353 affected the activation of P2X3R in sensory neurons from dorsal root ganglia or spinal cord sensory neurons, making micturition effects less evident than is the case for intravenous injections or direct intravesical application, as observed in female rats with neurogenic bladder following semichronic spinal cord injury [8,14]. Nonetheless, it is important to keep in mind that P2X3R are also required for the activity of primary and sensory neurons involved in the micturition process [28]. Thus, the EMG/CMG effects observed in MOR after AF-353 injection may be related to a negative effect on the activity of sensory neurons, with a concentrated effect on urothelial P2X3R at the lower urinary tract level.

In congruence with the prostate-bladder reflex, where sensory neurons in the lumbosacral spinal cord region may be responsible for their regulation [23], the application of the P2X3R antagonist could attenuate the corresponding ATP-mediated neuronal activity. For example, an intrathecal application of a similar P2X3 antagonist (AF-792) reduced the expression of phosphorylated extracellular signal-related kinase in lumbosacral dorsal horn neurons from rats with bladder overactivity induced with acetic acid, resulting in improved voiding function in a bladder dysfunction model, and supporting the relevance of additional cell targets for purinergic modulation and voiding control [29,30].

It is important to keep in mind that the sources for the EMG recordings from the bladder dome may include a combination of detrusor, innervation, and mechanical artifacts that may alter electrophysiological recordings. Although the source of the acquired EMG from the bladder dome remains unknown, the signals from the lower urinary tract are highly reproducible and persistently reflect voiding activity [8,13]. In this study, EMG responses were affected by AF-353 in MOR, which can be attributed to the fact that P2X3R play a fundamental role in the activity of sensory neurons [7], in turn attenuating the hyperexcited putative prostate-bladder reflex [23].

We took first steps towards evaluating the impact of an aging prostate on voiding dysfunction in Long-Evans rats, as a potential naturally occurring hyperplasia model. Bladder dysfunction in these aged rats may be strongly related to prostate enlargement, exacerbating a complex prostate-bladder reflex where P2X3R in urothelial cells and sensory neurons play a significant role, leading to voiding dysfunction. The results obtained after systemic application of the P2X3R antagonist AF-353 suggest that targeting these receptors can be a promising clinical alternative for treating age-related lower urinary tract symptoms. However, special care should be taken regarding the effects on other cell types expressing P2X3R, as inhibition may alter normal sensory function not related to the lower urinary tract [7].
A detailed comparison within the lifespan of the male rat should be performed to consolidate the current observations on micturition. Our findings bring about new perspectives for understanding voiding dysfunction and further evaluating uroinary function in an animal model with potential translational impact.

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AUTHOR CONTRIBUTION STATEMENT

- Conceptualization: AM
- Data curation: CZ, AM
- Formal analysis: CZ, YC, YZ, AM
- Funding acquisition: TB, YZ, AM
- Methodology: CZ, X Li, AM
- Project administration: AM
- Visualization: T Boone, YC, YZ, AM
- Writing-original draft: CZ, X Li, AM
- Writing-review & editing: CZ, X Li, TB, YC, YZ, AM

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