Research Article

P2Y Receptor Modulation of ATP Release in the Urothelium

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Received 14 February 2014; Accepted 28 March 2014; Published 14 April 2014

Academic Editor: Rok Romih

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The release of ATP from the urothelium in response to stretch during filling demonstrates the importance of the purinergic system for the physiological functioning of the bladder. This study examined the effect of P2 receptor agonists on ATP release from two urothelial cell lines (RT4 and UROtsa cells). Hypotonic Krebs was used as a stretch stimulus. Incubation of urothelial cells with high concentrations of the P2Y agonist ADP induced ATP release to a level that was 40-fold greater than hypotonic-stimulated ATP release ($P < 0.001$, ADP EC₅₀ 1.8 μ M). Similarly, an increase in ATP release was also observed with the P2Y agonist, UTP, up to a maximum of 70% of the hypotonic response (EC₅₀ 0.62 μ M). Selective P2 receptor agonists, $\alpha\beta$ -methylene-ATP, ATP- γ -S, and 2-methylthio-ADP had minimal effects on ATP release. ADP-stimulated ATP release was significantly inhibited by suramin (100 μ M, $P = 0.002$). RT4 urothelial cells break down nucleotides (100 μ M) including ATP, ADP, and UTP to liberate phosphate. Phosphate liberation was also demonstrated from endogenous nucleotides with approximately 10% of the released ATP broken down during the incubation. These studies demonstrate a role for P2Y receptor activation in stimulation of ATP release and emphasize the complexity of urothelial P2 receptor signalling.

1. Introduction

The first evidence for P2 receptor signalling from the urothelium was provided in 1997 with the work of Ferguson et al., who demonstrated in tissue strips that ATP is released from the urothelium in response to stretch [1]. Our understanding of the sensory role of the urothelium has since greatly increased, with stretch-induced ATP release from the urothelium also identified in cultured cells [2] and *in vivo* preparations [3].

ATP binds to two families of P2 receptors, P2X ligand-gated ion channels and G protein-coupled P2Y receptors. To date seven P2X receptors have been identified [4], with P2X₄ and P2X₂ receptors [5] expressed on the urothelium and P2X₃ expressed on the urothelium [5, 6] as well as on suburothelial afferent nerves and myofibroblasts [7]. It is hypothesised that ATP released from the urothelium interacts with P2 receptors located on both afferent nerves (P2X₃) [8] and myofibroblasts to signal bladder fullness. A role for ATP in bladder dysfunction has been postulated with increased ATP release associated with sensory disorders such as interstitial

cystitis [9] and painful bladder syndrome [10]. Furthermore, ATP has been shown to play a role in bladder sensation, with ATP concentration in the intravesical fluid correlating with the first desire to void in patients with both overactive bladder [3] and painful bladder [11] but not in control patients.

P2X receptors are activated preferentially by ATP; thus urothelial derived ATP may have an autocrine action on urothelial P2X receptors. However the urothelium also expresses ectoATPase enzymes [12] and so is able to hydrolyse ATP (and other nucleotides) to breakdown products such as ADP, AMP, and adenosine. The perceived function of these enzymes is to limit the availability of ATP. Recently, the urothelium has been shown to release nucleotides other than ATP with twelve nucleotides quantified by HPLC in patient urine specimens [13]. The levels of ADP, UTP, UDP, and GTP were found to be more than 10 times higher than the level of ATP [13]. Other epithelial cells have also been shown to release nucleotides such as ATP, ADP, AMP, adenosine, UTP, and UDP, in response to hypotonic stimulus [14]. The role of these nucleotides in P2 receptor signalling in the bladder is yet to be determined. It is known that in addition

to ATP, other nucleotides, in particular ADP and UTP, can also be substrates for P2Y receptors; however, their effects on urothelial P2Y receptors have not been explored. Eight P2Y receptors have been identified [15] with P2Y₁ [5, 16], P2Y₂ [5, 16, 17], P2Y₄ [16, 17], and P2Y₁₁ [5] expressed on urothelial cells. P2Y₆ receptors are expressed on suburothelial myofibroblasts [18] but their expression on urothelial cells remains controversial [5, 19]. The function of these receptors on the urothelium is currently undetermined. In addition, adenosine formed from the breakdown of ATP binds to P1 receptors which are also expressed on the urothelium [20].

It is very clear that the complexity of urothelial P2 receptor signalling and the feedback of ATP breakdown and release have not been adequately explored to date. Therefore the aims of this study were to examine the effect of purine and pyrimidine nucleotides on urothelial cell ATP release. We hypothesise that purines and pyrimidines selective for P2Y receptors will modulate ATP release from the urothelium.

2. Materials and Methods

2.1. Cell Culture. Human urothelial RT4 cells were obtained from the ECACC. Cells were grown at 37°C with 5% CO₂ in McCoy's 5A culture medium supplemented with 10% foetal bovine serum, 100 units/mL of penicillin, 100 µg/mL of streptomycin, and 0.25 µg/mL of fungizone. When confluent, RT4 cells were passaged with 0.05% Trypsin-EDTA for 5 minutes and then plated onto T75 flasks for continuous passage or onto 24 well plates for use in experiments when confluent (approximately 3 to 5 days after passage).

Human urothelial UTOsa cells were a gift from Dr. Scott Garrett from the University of North Dakota. Cells were grown at 37°C with 5% CO₂ in Dulbecco's modified eagle's medium supplemented with 5% foetal bovine serum, 100 units/mL of penicillin, 100 µg/mL of streptomycin, 0.25 µg/mL of fungizone, and 1% glucose. UTOsa cells were passaged in the manner described above for RT4 cells.

2.2. ATP Release Studies. ATP release was determined as we have described previously [2]. Urothelial cells were washed (three times) with carbogenated Krebs-Henseleit solution (containing mM: NaCl 118, KCl 4.7, NaHCO₃ 25, KH₂PO₄ 1.2, MgSO₄ 1.2, CaCl₂ 2.5, and D-glucose 11.7). The basal level of ATP release was then determined by a 10-minute incubation in 500 µL Krebs-Henseleit solution. These media were collected and followed with the experimental treatments. Cells were exposed to normal Krebs-Henseleit (control) or the indicated concentration of nucleotide. Cells were exposed to hypotonic Krebs-Henseleit (1:2 dilution of Krebs-Henseleit in distilled water), a stimulus commonly used to mimic stretch in cultured cells [2]. Cells were treated for 10 min before the supernatant (200 µL) was collected and used for ATP determinations.

ATP concentration in the supernatant was measured using the bioluminescence assay according to the manufacturer instructions. Equal volumes of the cellular supernatant or ATP standard solutions (10⁻⁶ to 10⁻¹⁰ M) were mixed

with the bioluminescence assay mix and the luminescence generated was measured immediately using a plate reader (BMG labtech Polarstar). The standard concentrations fell within the upper and lower limits of sensitivity of the ATP bioluminescence assay. The ATP concentration in the cell supernatant was calculated relative to the standard curve. Luminescence was also measured for all concentrations of the nucleotides used. The luminescence determined for each concentration of nucleotide was used as a blank for that treatment and was subtracted from the corresponding cellular luminescence before the ATP concentration was calculated. Treatments were carried out in triplicate and the mean ATP concentration (in nM) per treatment determined.

2.3. EctoATPase Studies. Confluent urothelial cells in 24 well plates were washed (three times) in phosphate free media (containing mM: NaCl 120, KCl 5, CaCl₂ 2, HEPES 20, and D-glucose 10) [12]. After washing cells were incubated with 100 µM nucleotide. After 30 minutes the supernatant was collected and the phosphate concentration determined. In additional experiments confluent cells in 12 well plates were incubated in control or hypotonic (50% dilution in water) phosphate free media for 10 or 30 minutes and both ATP concentration and phosphate concentration determined.

Phosphate liberated from nucleotides was determined as previously described [21]. Equal volumes of cell supernatant and colour reagent (containing 1% ammonium molybdate, 0.3 M H₂SO₄, 4% FeSO₄) were mixed. After 15 minutes absorbance at 750 nm was measured in a plate reader (BMG labtech Polarstar). Phosphate concentration in the cell supernatant was determined relative to a standard curve (KH₂PO₄, 10 to 150 nM). Treatments were carried out in triplicate and the mean phosphate concentration (in nM) per treatment determined.

2.4. Statistics. Results are nonnormally distributed and as such are expressed as median with interquartile range (IQR). Two different treatments were compared using a Mann-Whitney *t*-test. Concentration response relationships were examined using a sigmoidal concentration response curve. All statistics were performed using Graphpad Prims (version 6) (San Diego, CA).

2.5. Materials. All cell culture reagents were purchased from Invitrogen (Mount Waverley, Australia). Bioluminescence ATP Assay kit and nucleotides were from Sigma-Aldrich (Sydney, Australia). All other reagents were of high analytical grade.

3. Results

3.1. Effect of Nucleotides on Urothelial Cell ATP Release. Hypotonic Krebs was used as a stretch stimulus and was seen to induce an approximate threefold increase in ATP release (*P* = 0.0006). Incubation of RT4 cells with ADP (100 µM for 10 minutes) induced ATP release to a level that was 40-fold higher compared to the hypotonic stimulus (Table 1, *P* < 0.0011). In the presence of AMP and adenosine, the level of

TABLE 1: ATP release in RT4 urothelial cells stimulated by 10-minute incubation with nucleotides (100 μ M).

	ATP release in RT4 cells
Control	20.28 (13.24–40.22) nM ($n = 33$)
Hypotonic	64.63 (43.75–92.95) nM ($n = 33$)
ADP	2641 (2006–3598) nM ($n = 13$)
AMP	8.56 (4.74–189.2) nM ($n = 17$)
Adenosine	5.58 (4.52–65.85) nM ($n = 4$)
CTP	51.12 (37.4–99.96) nM ($n = 11$)
GTP	55.33 (34.42–173.4) nM ($n = 15$)
UTP	60.57 (36.51–126.1) nM ($n = 11$)
α,β -Methylene-ATP	26.6 (22.5–77.5) nM ($n = 9$)
ATP- γ -S	2294 (429–2538) nM ($n = 16$)

TABLE 2: ATP release in UROtsa urothelial cells stimulated by 10-minute incubation with nucleotides (100 μ M).

	ATP release in UROtsa cells
Control	11.08 (4.3–30.2) nM ($n = 6$)
Hypotonic	68.3 (35.6–133.4) nM ($n = 6$)
ADP	3377 (752–6361) nM ($n = 6$)
UTP	61.2 (26.6–115.7) nM ($n = 6$)
2-Methylthio-ADP	17.7 (13.9–26.9) nM ($n = 6$)

ATP release was not significantly different to the control level (Table 1). The response to CTP and GTP was not significantly different to the ATP release induced by hypotonic media (Table 1).

When the concentration response effect of ADP induced ATP release was determined it was seen to only occur at high concentrations of ADP (Figure 1(a)) with an EC₅₀ of 1.8 μ M determined ($n = 5$). A concentration dependent inhibition of ATP release was seen with AMP (Figure 1(b)) with an EC₅₀ of 0.33 μ M ($n = 7$). A concentration dependent increase in ATP release was observed with UTP, up to a maximum of approximately 70% of the hypotonic response (Figure 1(c)). An EC₅₀ of 0.62 μ M was determined ($n = 7$).

In RT4 cells, concentration response relationships were also determined for P2 receptor agonists and antagonists. The P2X agonist, $\alpha\beta$ -methylene-ATP, and P2Y agonist, ATP- γ -S, had no effect on ATP release (Figures 2(a) and 2(b), $n = 9$ and 15, resp.). The more selective P2Y₁ agonist, 2-methylthio-ADP, inhibited ATP release in a concentration dependent manner (Figure 2(c), EC₅₀ 0.16 μ M, $n = 8$).

P2 receptor antagonists were investigated for their ability to inhibit ADP-stimulated ATP release. PPADS (100 μ M) had no effect on ADP-stimulated ATP release (88.4 (65.2–113.4)% of ADP-stimulated release, $n = 8$, $P = 0.4$) while suramin (100 μ M) significantly inhibited ADP-stimulated ATP release (43.3 (27.6–41.7)% of ADP-stimulated ATP release, $P = 0.002$, $n = 4$; IC₅₀ 8.3 μ M, $n = 8$).

The results observed in RT4 urothelial cells were also confirmed in UROtsa cells (Table 2). Similar to the results observed in RT4 urothelial cells, treatment of UROtsa cells with hypotonic Krebs induced an increase in ATP release ($P = 0.0087$). Incubation of UROtsa cells with ADP (100 μ M

TABLE 3: Phosphate liberation from 30-minute incubation of RT4 urothelial cells with 100 μ M nucleotides ($n = 8$).

	Phosphate liberated (nM)
ATP	1.3 (–0.9–3.8)
ADP	7.5 (1.9–12.1)
AMP	4.1 (3.3–7.7)
CTP	3.4 (1.2–4.8)
GTP	2.6 (0.3–6.2)
UTP	7.8 (6.5–9.9)

for 10 minutes) induced an almost 50-fold increase in ATP release compared to that seen with the hypotonic stimulus ($P < 0.0022$). UTP (100 μ M) initiated ATP release similar to that seen with hypotonic stimulus while 2-methylthio-ADP (100 μ M) elicited an ATP release response similar to the control level.

3.2. EctoATPase Activity of Urothelial Cells. RT4 urothelial cells were shown to have the capacity to liberate phosphate from nucleotides (100 μ M), with the greatest amount of phosphate liberated from ADP and UTP (Table 3). In addition, a small amount of phosphate liberation was able to be demonstrated from endogenous nucleotides released during the experiment (Figure 3). When RT4 cells were incubated in either control or hypotonic phosphate free buffer for 10 or 30 minutes, significant ATP release was induced (Figure 3(a), $n = 8$). After a 10-minute incubation the level of release in the control was approximately 40 nM and in hypotonic phosphate free media was approximately 80 nM which was comparable to that seen using Krebs-Henseleit solution (Table 1). Over the 10- or 30-minute incubation some of this endogenous ATP was broken down by RT4 cells to liberate phosphate that could be detected (Figure 3(b), $n = 8$). Significantly more phosphate was detected in cells treated with hypotonic media for 10 and 30 minutes ($P = 0.0002$ and 0.0019 , resp.). After a 10-minute incubation the level of phosphate liberated indicated that approximately 10% (7 to 15%) of the endogenously released ATP was broken down during the incubations in control and hypotonic media, respectively. In the presence of the ectoATPase inhibitor ARL67156 (100 μ M) the amount of ATP detected in cells treated with hypotonic phosphate free buffer increased 5-fold after both 10- and 30-minute incubations (Figure 3(c), $n = 4$).

4. Discussion

The demonstration of ATP release from the rabbit bladder urothelium in response to stretch by Ferguson and colleagues in 1997 [1] has been pivotal to our understanding of signalling within the bladder. However, our understanding of the factors that modulate ATP release and the autocrine signalling that occurs at the urothelial cell layer is limited. The current study has demonstrated that P2 receptor agonists are able to stimulate ATP release from urothelial cells. In fact the P2Y agonist ADP was shown in two urothelial cell lines, to stimulate a level of ATP release that is far in excess of

FIGURE 1: Concentration response relationships for nucleotides ADP (a), AMP (b), and UTP (c) on RT4 urothelial cell ATP release.

that stimulated by hypotonic media. Similarly, UTP, another P2Y agonist [22], was also shown to stimulate ATP release, although to a lesser extent. In contrast, P2X agonists such as α,β -methylene-ATP were not seen to stimulate ATP release from cultured urothelial cells. These results are similar to the recently published findings by Sui and colleagues who reported that the P2Y agonist UTP stimulated ATP release in guinea pig and human bladder mucosal strips, while the P2X agonist α,β -methylene-ATP had no effect [22]. UTP stimulated ATP release has also been reported from primary cultures of rat bladder urothelial cells [17].

It has been shown that P2Y₁ [5, 16], P2Y₂ [5, 16, 17], P2Y₄ [16, 17], and P2Y₁₁ [5] receptor subtypes are present in bladder urothelial cells (see introduction). While P2X receptors respond preferentially to ATP, P2Y receptors are divided on the basis of their sensitivity to adenine nucleotides (P2Y₁, P2Y₁₁, P2Y₁₂, and P2Y₁₃) or uracil nucleotides (P2Y₂, P2Y₄, P2Y₆, and P2Y₁₄) [23]. Of the P2Y receptors that respond to adenine nucleotides, P2Y₁, P2Y₁₂, and P2Y₁₃ respond to ADP while P2Y₁₁ receptors respond to ATP [23]. In the current study, ATP release was stimulated by both ADP and UTP, although the release elicited by ADP was far

FIGURE 2: P2 receptor agonist stimulated ATP release. Incubation of RT4 urothelial cells with increasing concentration of α, β -methylene-ATP (a), ATP- γ -S (b), and 2-methylthio-ADP (c).

greater than that elicited by UTP, suggesting the involvement of more than one P2Y receptor subtype. The EC₅₀ value of ADP-stimulated ATP release suggests activity at P2Y₁ or P2Y₁₁ receptors. This is in agreement with previously reported EC₅₀s at P2Y₁ of 0.9–8 μ M [24–26]. The effects of UTP were closely associated with EC₅₀ values at P2Y₂ or P2Y₄ receptors, agreeing with previously reported EC₅₀s (0.14–0.8 μ M) at the P2Y₂ receptor [24, 25] and P2Y₄ receptor (2.5 μ M) [24, 25]. In contrast, the selective P2Y₁ agonist, 2-methylthio-ADP [26], did not stimulate ATP release. The current study therefore suggests that more than one P2Y

receptor subtype is likely to be involved in nucleotide stimulated ATP release. Unfortunately, the lack of subtype selective agonists for individual P2Y receptors makes it difficult to definitively identify the receptor subtypes involved in the observed responses.

It is possible that the apparent ATP release stimulated by high concentrations of ADP was due to the conversion of ADP to ATP by cell membrane-bound adenylate kinase. While this enzyme is predominately intracellular [27], it has been identified on the cell membrane of endothelial cells [28] and other cell types [27] although there have been no

FIGURE 3: Phosphate liberation from endogenously released ATP. Incubation of RT4 urothelial cells in control or hypotonic phosphate free buffer for 10 or 30 minutes induced ATP release (a). Over the 10- or 30-minute incubation a small amount (approximately 10%) of this ATP was broken down by urothelial cells to liberate phosphate that could be detected (b). The presence of the ectoATPase inhibitor ARL67156 (100 μ M) was seen to increase the amount of ATP detected after both 10 and 30 minutes (c).

reports of cell membrane-bound adenylate cyclase in the urothelium. However, coincubation of ADP with the P2 receptor antagonist suramin inhibited ADP-stimulated ATP release, suggesting that it is indeed P2Y receptors that are involved in this response (reported IC_{50} s 3–50 μ M) [15].

The source of ADP *in vivo* is unknown; however, nucleotides have recently been isolated from human urine samples [13]. In addition, urothelial cells express ectonucleotidase enzymes capable of degrading ATP and UTP to their respective nucleotides [29]. The basal and intermediate cells of the mouse bladder urothelium express NTPDase 3 [29]. Similarly, the human urothelial cell line RT4 used in the current study expresses NTPDase 3 and 5 [12] but not NTPDase 1 [12, 30]. NTPDase 1, 2, and 3 are known to face the extracellular environment and catalyze the breakdown of

extracellular ATP [29]. It is thought that ectoATPase enzymes function to limit the exposure of P2 receptors to their ligands and to modulate the autocrine response to released ATP [29].

In the current study the activity of ectoATPase in urothelial RT4 cells was examined in three ways. Firstly, ectoATPase enzymes present on the RT4 urothelial cells were able to liberate phosphate from stretch-induced ATP release, indicating a capacity to generate mediators such as ADP from stretch-induced ATP release. Secondly, incubation of the ectoATPase inhibitor ARL67156 (100 μ M) [31] resulted in a higher level of stretch-induced ATP release being detected. At this concentration, ARL67156 has been shown to inhibit the activity of NTPDase3 (K_i 18 μ M) [32]. Finally, we demonstrated the capacity of ectoATPase enzymes associated with RT4 urothelial cells to liberate phosphate

FIGURE 4: Hypothetical schematic representation of the complexity of the effect of nucleotides and stretch on urothelial cell ATP release.

from exogenous nucleotides including ATP, ADP, AMP, CTP, GTP, and UTP. Similarly, Stella and associates showed that RT4 urothelial cells were able to liberate phosphate from exogenous nucleotides [12].

The results obtained in the current study demonstrate that P2 receptor signalling in the urothelium is a complex interaction between ATP release and breakdown. To add to the complexity, it now seems likely that products formed from breakdown of nucleotides such as ATP also have effects at P2 receptors located on the urothelium. Our understanding of this complex interaction between ATP release and breakdown has been shown schematically in Figure 4. ADP has been previously postulated to stimulate urothelial ATP release [9] and to activate intracellular calcium transients [5, 33] that were greater than those activated by hypotonic solutions [33], a commonly used stimulus for ATP release. It was proposed that the ATP-dependent calcium transients were mediated by urothelial P2Y (rather than P2X) receptors [5, 33]; however, it is known that ATP has a low affinity at P2Y receptors [23, 25]. It is therefore possible that the calcium transients induced by endogenous ATP were in fact due to the generation of ADP by membrane-bound ectoATPases. This is supported by the findings of the current study which demonstrate that ADP is capable of stimulating ATP release. Adding to the complexity of P2 receptor signalling, further breakdown of ADP to AMP and adenosine was seen to inhibit ATP release. Inhibition of ATP release by adenosine has been previously reported in rabbit bladder mucosal strips [20]. These findings indicate that while the initial breakdown of ATP to ADP may exert positive feedback for ATP release which is short lived, further breakdown of ATP to AMP and adenosine may provide negative feedback for ATP release.

Activation of urothelial P2Y receptors has been proposed to be responsible for a number of physiological functions of the urothelium. Exogenous application of P2Y receptor agonists, including ADP and UTP, has been shown to increase spontaneous activity in rat bladder sheets [34]. Similar to the results seen in this study, it was unlikely that a single P2Y receptor subtype was responsible for the reported increase in spontaneous activity [34] with the involvement of P2Y₁, P2Y₂, P2Y₄, and P2Y₆ being hypothesised. In addition,

activation of P2Y₂ receptors leads to release of the inflammatory mediators interleukins 8 and 6 from uroepithelial cells [35], indicating a role for activation of P2Y receptors in inflammatory responses in the bladder. Interestingly, expression of the P2Y₂ receptor decreased in a feline model of interstitial cystitis [16]. Intravesical instillation of a P2Y₆ selective agonist induced bladder overactivity characterised by increased voiding frequency in a rat cystometry model in addition to increased ATP release into the voided fluid [19].

Alterations in urothelial ATP release have been identified in bladder dysfunction including interstitial cystitis [9], painful bladder syndrome [10, 11], and overactive bladder [3]. The results of this study demonstrate the complexity of P2 receptor signalling in the urothelium by elucidating a role for a number of P2Y receptor subtypes in initiating ATP release. Recent literature has shown the important role of these receptors in normal bladder physiology, indicating that these receptors may represent a potential future target for the treatments of bladder dysfunction.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

Acknowledgments

The authors thank Dr. Catherine McDermott and Professor Russ Chess-Williams from Bond University for their assistance with these cell culture studies. They also thank Professor Paul Else from the University of Wollongong for assistance with establishing ectoATPase assays. This study was supported by a Small URC grant from the University of Wollongong. Thanks are also due to Tim Cowan and Nathan Ralston-Bryce for assistance with luminescence assays.

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