

## So much cost, such little progress

Bryan, Richard T; Kirby, Roger; O'Brien, Tim; Mostafid, Hugh

DOI:

[10.1016/j.eururo.2014.02.031](https://doi.org/10.1016/j.eururo.2014.02.031)

License:

Creative Commons: Attribution-NonCommercial-NoDerivs (CC BY-NC-ND)

*Document Version*

Peer reviewed version

*Citation for published version (Harvard):*

Bryan, RT, Kirby, R, O'Brien, T & Mostafid, H 2014, 'So much cost, such little progress', *European urology*, vol. 66, no. 2, pp. 263-264. <https://doi.org/10.1016/j.eururo.2014.02.031>

[Link to publication on Research at Birmingham portal](#)

### **Publisher Rights Statement:**

Checked October 2015

### **General rights**

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

- Users may freely distribute the URL that is used to identify this publication.
- Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.
- User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)
- Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

### **Take down policy**

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact [UBIRA@lists.bham.ac.uk](mailto:UBIRA@lists.bham.ac.uk) providing details and we will remove access to the work immediately and investigate.

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19

**Editorial: SO MUCH COST, SUCH LITTLE PROGRESS**

**Richard T Bryan, Roger Kirby, Tim O'Brien, Hugh Mostafid**

- RT Bryan PhD MRCS** School of Cancer Sciences, University of Birmingham, UK.  
Council, Section of Urology, Royal Society of Medicine.
- R Kirby MD FRCS(Urol)** The Prostate Centre, London, UK.  
Secretary, The Urology Foundation.
- T O'Brien DM FRCS(Urol)** Guy's and St. Thomas' NHS Foundation Trust, London, UK.  
Chairman, Section of Oncology, British Association of Urological Surgeons.
- H Mostafid FRCS(Urol) FEBU** Royal Berkshire NHS Foundation Trust, UK.  
Chairman, Action on Bladder Cancer.

**Word count:** 986

**Keywords:** Bladder cancer, economics, expense

**Correspondence to:** Mr RT Bryan, School of Cancer Sciences, University of Birmingham, Edgbaston, Birmingham B15 2TT, United Kingdom.  
E: [r.t.bryan@bham.ac.uk](mailto:r.t.bryan@bham.ac.uk), T: +44 121 414 7870, F: +44 121 414 2230.

20 Urothelial bladder cancer (UBC) is burdensome for patients and expensive for healthcare  
21 providers [1]. Outcomes have changed little for three decades, despite significant  
22 improvements in 5-year survival rates for prostate and kidney cancers during this period [1].  
23 Furthermore, patient pathways are complex, prolonged, and practiced in various permutations  
24 at every stage:

- 25 • The investigation of patients with suspected UBC requires multiple diagnostic  
26 procedures [2;3]; various combinations of tests are utilised [4].
- 27 • TURBT can be performed by a number of different techniques with a number of different  
28 energy sources, and utilising a variety of optical image enhancement technologies [5].
- 29 • Further treatment may be required in the form of intravesical therapy with various  
30 agents, with or without chemohyperthermia or electromotive drug administration [3].
- 31 • Long-term surveillance is the mainstay of subsequent management [2;3]; various  
32 surveillance schedules are practiced [6].
- 33 • Surveillance may or may not utilize urinary biomarkers, and treatment of recurrence  
34 may be carried out in the office or the operating theatre [6].
- 35 • For curative intent, patients who present with or progress to MIBC are treated by  
36 radiotherapy [2;7], chemoradiotherapy [8], radical cystectomy (open, laparoscopic or  
37 robot-assisted), or neoadjuvant chemotherapy followed by radical cystectomy [2;7];  
38 adjuvant chemotherapy is utilized in some units.

39  
40 In this month's issue of *European Urology*, Svatek *et al* provide a review of the costs and other  
41 considerations for these approaches, and discuss key issues regarding the wider economics of

42 bladder cancer care [6]. This represents a useful overview for the practicing urologist. In  
43 particular, the authors demonstrate that there are large gaps in our knowledge regarding the  
44 efficacy and cost-effectiveness of these approaches and a lack of sufficiently-powered  
45 randomised controlled trials (RCTs), with expensive tools having crept into everyday practice  
46 without the necessary thorough evaluations. They highlight that there is a clear and urgent  
47 need for the development of new drugs for UBC, both NMIBC and MIBC. The prevalence of  
48 NMIBC and its protracted course compared to MIBC is such that the cumulative cost of care is  
49 thought to be even more substantial than MIBC [6], so the gains to be made in preventing  
50 recurrence and progression of NMIBC could be the most significant. Furthermore, the individual  
51 physician has the greatest impact on the cost of care of NMIBC, yet variation in treatment  
52 intensity does not impact survival or the avoidance of subsequent major interventions [6].  
53 Other authors have recently highlighted these and other issues in bladder cancer care [1].  
54 However, in order to make practice-changing recommendations, robust and detailed  
55 assessments of specific elements of these complicated pathways are required, utilizing complex  
56 modeling and statistics, and measures of cost-effectiveness. Such analyses have previously  
57 been undertaken in the UK in the form of Health Technology Assessments [4;9]. Reasonably,  
58 Svatek *et al* do not venture into this complex territory, but such health services research is  
59 urgently needed alongside basic and translational research and clinical trials [1]. Furthermore,  
60 the non-medical costs associated with UBC care (that are borne by patients, their families, their  
61 employers) and the costs associated with untimely deaths due to UBC are simply staggering [6].  
62 Perhaps the treatment of UBC has far more impact on HRQoL than we have previously realised?

63

64 The Authors could have been more prescriptive in their conclusions to send a clearer message.  
65 For example, they present data that level 1 evidence and clinical guidelines are being ignored  
66 [1;6], yet fail to recommend that such evidence and guidelines be more closely adhered to.  
67 Perhaps we don't actually need more RCTs of BCG maintenance therapy, which are both  
68 expensive and protracted? Instead, would a better use of resources be to gain a clearer  
69 understanding of BCG's mechanism of action and the immunological milieu of the bladder  
70 tumour microenvironment, potentially leading to the development of new therapeutics for all  
71 UBC patients? NMIBC is also an ideal setting in which to assess the effectiveness of novel low  
72 toxicity therapeutics and/or chemopreventive agents administered long-term (and several such  
73 RCTs are in follow-up, eg. BOXIT, SELENIB), yet such strategies are not discussed by the Authors.  
74 As for urinary biomarkers, their real utility may not actually lie in their ability to detect new or  
75 recurrent disease, but in their ability to risk stratify patients early in their pathway so that they  
76 are investigated and managed more appropriately and expeditiously [10]. There is a lot that we  
77 could do now to redesign these pathways and interventions [10], yet there is a reluctance to  
78 change and a significant lack of research funding [1].

79 And it is this lack of research funding that underlies our complex and varied pathways. We don't  
80 actually have the robust evidence base to support a lot of what we do, and where the robust  
81 evidence and high grade recommendations exist, the uptake is poor (eg. single-shot intravesical  
82 mitomycin C [3], neoadjuvant platinum-based combination chemotherapy [7]) [6].  
83 Consequently, a spectrum of alternatives is practiced by individual urologists and/or individual  
84 units, possibly accentuated in the USA by illogical reimbursement patterns [1]. The Authors'  
85 lack of decisive conclusions is therefore understandable.

86 If we are to tackle bladder cancer and improve outcomes, as we have done for prostate and  
87 kidney cancer, then we need to lobby for more funding for RCTs, translational science and  
88 health services research, and address the poor awareness of UBC among the general public and  
89 the nonurological scientific community [1]. And where are MIBC's innovative new drugs? It  
90 feels as though the pharmaceutical industry have deserted UBC in search of lower hanging fruit.  
91 These issues were specifically discussed amongst leading UK urologists and oncologists at The  
92 Royal Society of Medicine Section of Urology Annual Winter Meeting in January, and it was  
93 concluded that The Royal Society of Medicine, the British Association of Urological Surgeons,  
94 The Urology Foundation and Action on Bladder Cancer should endeavour to undertake a  
95 collaborative and concerted effort to advance the cause for bladder cancer patients. We need  
96 to make much more progress, perhaps improving cost-effectiveness along the way.

97

- 100 (1) Kaplan AL, Litwin MS, Chamie K. The future of bladder cancer care in the USA. *Nat Rev*  
101 *Urol* 2014; 11:59-62.
- 102 (2) Kaufman DS, Shipley WU, Feldman AS. Bladder cancer. *Lancet* 2009; 374:239-249.
- 103 (3) Babjuk M, Burger M, Zigeuner R, Shariat SF, van Rhijn BW, Comperat E, Sylvester RJ,  
104 Kaasinen E, Bohle A, Palou RJ, Roupret M. EAU Guidelines on Non-Muscle-invasive  
105 Urothelial Carcinoma of the Bladder: Update 2013. *Eur Urol* 2013.
- 106 (4) Rodgers M, Nixon J, Hempel S, Aho T, Kelly J, Neal D, Duffy S, Ritchie G, Kleijnen J,  
107 Westwood M. Diagnostic tests and algorithms used in the investigation of haematuria:  
108 systematic reviews and economic evaluation. *Health Technol Assess* 2006; 10:iii-259.
- 109 (5) Wilby D, Thomas K, Ray E, Chappell B, O'Brien T. Bladder cancer: new TUR techniques.  
110 *World J Urol* 2009; 27:309-312.
- 111 (6) Svatek et al. The economics of bladder cancer: costs and considerations of caring for this  
112 disease. *Eur Urol* . 2014.  
113 Ref Type: In Press
- 114 (7) Witjes JA, Comperat E, Cowan NC, De SM, Gakis G, Lebret T, Ribal MJ, Van der Heijden  
115 AG, Sherif A. EAU Guidelines on Muscle-invasive and Metastatic Bladder Cancer:  
116 Summary of the 2013 Guidelines. *Eur Urol* 2013.
- 117 (8) James ND, Hussain SA, Hall E, Jenkins P, Tremlett J, Rawlings C, Crundwell M, Sizer B,  
118 Sreenivasan T, Hendron C, Lewis R, Waters R, Huddart RA. Radiotherapy with or without  
119 chemotherapy in muscle-invasive bladder cancer. *N Engl J Med* 2012; 366:1477-1488.
- 120 (9) Mowatt G, Zhu S, Kilonzo M, Boachie C, Fraser C, Griffiths TR, N'Dow J, Nabi G, Cook J,  
121 Vale L. Systematic review of the clinical effectiveness and cost-effectiveness of  
122 photodynamic diagnosis and urine biomarkers (FISH, ImmunoCyt, NMP22) and cytology  
123 for the detection and follow-up of bladder cancer. *Health Technol Assess* 2010; 14:1-iv.
- 124 (10) Bryan RT, Shimwell NJ, Wei W, Devall AJ, Pirrie SJ, James ND, Zeegers MP, Cheng KK,  
125 Martin A, Ward DG. Urinary EpCAM in urothelial bladder cancer patients:  
126 characterisation and evaluation of biomarker potential. *Br J Cancer* 2013.  
127  
128