In Touch with your Feminine Side: How Oestrogen Metabolism Impacts Prostate Cancer

Rahman, Habib-Ur; Hofland, Johannes; Foster, Paul

DOI:

[10.1530/ERC-16-0118](https://doi.org/10.1530/ERC-16-0118)

License:

None: All rights reserved

Document Version

Peer reviewed version

Citation for published version (Harvard):

Rahman, H-U, Hofland, J & Foster, P 2016, 'In Touch with your Feminine Side: How Oestrogen Metabolism Impacts Prostate Cancer', *Endocrine-related cancer*. <https://doi.org/10.1530/ERC-16-0118>

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

Publisher Rights Statement:

Disclaimer: this is not the definitive version of record of this article. This manuscript has been accepted for publication in Endocrine-Related Cancer, but the version presented here has not yet been copy-edited, formatted or proofed. Consequently, Bioscientifica accepts no responsibility for any errors or omissions it may contain. The definitive version is now freely available at doi: 10.1530/ERC-16-0118

Checked May 2016

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 providing details and we will remove access to the work immediately and investigate.

1 **In Touch with your Feminine Side: How Oestrogen Metabolism Impacts Prostate**
2 **Cancer**

3 ¹Habibur P. Rahman, ²Johannes Hofland, ^{1,3}Paul A. Foster

4

5 ¹ Institute of Metabolism and Systems Research, University of Birmingham, Birmingham,
6 B15 2TT, UK.

7 ² Department of Internal Medicine, Erasmus Medical Center, 3000 CA Rotterdam, The
8 Netherlands.

9 ³ Centre for Endocrinology, Diabetes and Metabolism, Birmingham Healthcare Partners,
10 Birmingham, B15 2TH, UK.

11 Tel: +44 (0)121 414 3776

12 Fax: +44 (0)121 415 8712

13

14 Short title: Oestrogens and Prostate Cancer

15

16 Proofs and Correspondence to:

17 Dr. P.A. Foster

18 Institute of Metabolism and Systems Research,

19 College of Medical and Dental School,

20 University of Birmingham,

21 Birmingham,

22 B15 2TT.

23 UK.

24

25 **Declaration of Interest**

26 The authors have nothing to declare.

27 **Funding**

28 No specific funding was received for this review.

29 **Abstract**

30 Prostate cancer is the primary male cancer with increasing global incidence rates making this
31 malignancy a significant healthcare burden. Androgens promote normal prostate maturity but
32 also influence the development and progression of prostate cancer. Intriguingly, evidence
33 now suggests endogenous and exogenous oestrogens, in the form of phytoestrogens, may be
34 equally as relevant as androgens in prostate cancer growth. The prostate gland has the
35 molecular mechanisms, catalysed by steroid sulphatase (STS), to unconjugate and utilise
36 circulating oestrogens. Furthermore, prostate tissue also expresses enzymes essential for local
37 oestrogen metabolism, including aromatase (CYP19A1) and 3 β - and 17 β -hydroxysteroid
38 dehydrogenases. Increased expression of these enzymes in malignant prostate tissue
39 compared to normal prostate indicates oestrogen synthesis is favoured in malignancy and thus
40 may influence tumour progression. In contrast to previous reviews, here we comprehensively
41 explore the epidemiological and scientific evidence on how oestrogens impact prostate
42 cancer, particularly focusing on pre-receptor oestrogen metabolism and subsequent molecular
43 action. We analyse how molecular mechanisms and metabolic pathways involved in
44 androgen and oestrogen synthesis intertwine to alter prostate tissue. Furthermore, we
45 speculate on whether oestrogen receptor status in the prostate affects progression of this
46 malignancy.

47

48

49

50

51

52

53

54 **Introduction**

55 In the UK prostate cancer is the number one male malignancy accounting for 25% of all new
56 cancer diagnoses in men (Siegel, et al. 2012). In 2011, there were almost 42,000 new cases
57 with an age-standardised incidence rate of 104.7 per 100,000. Prostate cancer is the second
58 leading cancer killer in UK men and 4th most common cause of cancer death in the general
59 population. Similarly, in Europe prostate cancer is the most common cancer in males and
60 third most common cancer overall (Jacob and Henrik 2006) . It is the third most common
61 cause of cancer deaths in men and sixth overall. Currently, prostate cancer is the second most
62 common cancer in males worldwide after lung cancer. However, it is predicted that prostate
63 cancer will become the most common cancer in men globally (Parkin, et al. 2001).

64 Survival statistics from prostate cancer have improved dramatically over the last four decades
65 which may be attributed to earlier detection and treatment granted by prostate specific
66 antigen (PSA) testing and transurethral resection of the prostate (TURP). The UK 10-year
67 survival has improved from 25% when diagnosed in 1970 to 84% in 2010 (Quaresma, et al.
68 2015). Prostate cancer primarily affects the elderly with 99.9% of patients diagnosed over the
69 age of 50 and the mean age at diagnosis being 73 (Parkin, et al. 1997). Furthermore, from
70 autopsy studies of non-cancer-related deaths, there is histological evidence of prostate
71 neoplasms in more than 50% of men in their 50s (Sakr, et al. 1993). As average male life
72 expectancy gradually increases, it is foreseeable that men will live longer with the disease
73 and may experience a poorer quality of life.

74 There are significant geographical variations between prostate cancer incidences around the
75 world with up to a 24-fold difference between the regions with the highest rates (in Australia,
76 North America and Western Europe) and the lowest rates (in India, Japan and China) (Center,
77 et al. 2012). While some of the discrepancies might be explained by disparities in healthcare

78 access, diagnostic methods, screening programmes and reporting systems; environment and
79 lifestyle remain considerable factors. Studies comparing the incidence of prostate cancer in
80 first and second generation Asian immigrants to USA with age-matched controls in their
81 native countries have found that migrants travelling from low risk countries to high risk
82 countries adopt the higher risk (Cook, et al. 1999). This advocates that environmental risk
83 factors may have a higher precedence than genetic associations in determining risk of
84 prostate cancer. Furthermore, environmental and lifestyle factors, diet in particular,
85 fundamentally alter endogenous hormones including sex steroids (Barazani, et al. 2014).
86 Indeed, factors such as smoking, increased physical exercise and a vegetarian diet increased
87 serum androgen concentrations in British men while obesity, high fat diet and sedentary
88 occupation reduced serum androgen concentrations (Allen et al. 2002). Such hormonal
89 changes have the propensity to subsequently affect tumour initiation and progression
90 (Kolonel, et al. 2004).

91

92 **Sex Steroids and Prostate Cancer**

93 Both males and females produce sex steroid hormones; the predominant androgens are
94 testosterone and the more biologically active dihydrotestosterone (DHT) and the predominant
95 oestrogens are oestrone (E_1) and the more biologically active oestradiol (E_2). However, the
96 ratio of the two hormones differs between the sexes significantly. In the prostate, androgens
97 are required for normal development and function. However, the role of oestrogens in normal
98 prostate development is ill defined as biochemical mechanisms are still under investigation;
99 the current dogma being that oestrogens are involved in the differentiation of epithelial tissue
100 (Chen, et al. 2012; Francis, et al. 2013) and regulation of prostatic angiogenesis (Montico, et
101 al. 2013).

102

103 Androgens have been implicated in prostate carcinogenesis since 1941 when Huggins
104 published his Nobel winning study showing testosterone injections exacerbate prostate cancer
105 in patients with late-stage disease and androgen deprivation alleviated the disease (Huggins
106 and Hodges 1941), this suggested prostate cancer as an androgen-dependent malignancy. The
107 primary source of androgens in males is testosterone secreted by the testicles, however, the
108 adrenal glands secrete 100-500 times greater amounts of dehydroepiandrosterone sulphate
109 (DHEAS), a testosterone precursor which can be converted peripherally in the prostate into
110 testosterone and DHT (Labrie, et al. 2005). Androgen ablation therapy is initially successful
111 in the vast majority of prostate cancers but relapse is common as tumours become castration
112 resistant; they still however continue to express androgen receptors which respond to very
113 low concentrations (as low as 10 pM) of peripherally synthesised testosterone and DHT
114 (Chen, et al. 2004; Mohler, et al. 2004). Using microarray experiments on LNCaP and
115 LAPC4 cell lines, Chen et al. (2004) showed an increase in androgen receptor mRNA and
116 protein expression *in vitro* and *in vivo* in castrated xenograft murine models which correlated
117 with tumour growth. Increased expression of androgen receptors amplified signals from low
118 levels of androgen ligands to confer castration resistance. Mohler et al (2004) demonstrated
119 using immunostaining and radioimmunoassays that activation of androgen receptors occur
120 even in human prostate cancer samples retrieved from chemically castrated patients. This
121 explains why surgical or medical castration is not 100% effective.

122 Previously, second-line hormonal therapy has proven to improve survival in patients with
123 castration-resistant disease, both before and after docetaxel chemotherapy. Both inhibition of
124 steroidogenic enzyme CYP17A1 using abiraterone and androgen receptor antagonism by
125 enzalutamide have successfully ablated continued androgen receptor activation and prostate
126 cancer growth (Beer , et al. 2014; de Bono , et al. 2011; Ryan , et al. 2013; Scher , et al.

2012). However, as with other androgen ablation therapy, resistance to abiraterone and enzalutamide inevitably develops.

Even though molecular mechanisms were not elucidated, oestrogens were traditionally considered to protect against prostate cancer. Therapeutic use of oestrogens was based on their anti-androgenic effects. Huggins reported exogenous oestrogens had protective properties mediated by a negative feedback effect on the hypothalamic-pituitary-gonadal (HPG) axis which reduced stimulation for androgen secretion from the testes (Huggins and Hodges 1941). Diethylstilbestrol (DES), a synthetic non-metabolised oestrogen is still used in certain clinics as a non-first line therapy to chemically castrate patients with metastatic prostate cancer (Bosset, et al. 2012; Clemons, et al. 2013). DES negatively feedbacks on the pituitary gland to reduce secretion of luteinizing hormone which reduces the stimulus for the testes to synthesise sex hormones. In addition to the effects oestrogens have on the HPG axis, demonstrated by quantitative PCR, DES inhibits androgen-stimulated telomerase activity and gene expression and induces apoptosis in LNCaP and PC3 prostate cancer cell lines in both the presence and absence of androgens (Geier, et al. 2010). On the contrary, while DES is still licensed in the UK for treatment of prostate cancer it is infrequently used as secondary treatment due to the accompanied high rates of cardiovascular toxicity (Malkowicz 2001).

Importantly, the interactions of oestrogens on androgen receptors should be considered. For example, E₂ can activate both wildtype and, with greater efficacy, mutated (T877A) androgen receptors in LNCaP cells (Susa et al J Cell Physiol 2015; Yeh et al. 1998; Veldscholte et al J Steroid Biochem Mol Biol. 1992). Mutations of the androgen receptor are uncommon in the early stages of prostate cancer but are much more frequent in late-stage disease. In one study, out of 99 patients diagnosed with early stage prostate cancer none were found to have

151 mutations in the androgen receptor. On the contrary, eight tumours out of 38 patients with
152 advanced prostate cancer were found to harbour androgen receptor mutations (Marcelli, et al.
153 2000; Brooke and Bevan 2009). There is, however, mounting evidence that oestrogens may
154 be involved in the initiation and progression of prostate cancer, although compelling evidence
155 confirming oestrogen binding affinity to AR is lacking.

156
157 Impact of Endogenous Oestrogens in Prostate Cancer

158 Males are exposed to a high oestrogen/androgen (E/T) ratio twice in their lifetime. The first is
159 as a foetus, during the third trimester when the maternal E₂ levels increase and foetal
160 androgen levels decrease. Raised E₂ levels stimulate the developing epithelial cells of the
161 prostate to proliferate but also cause morphological changes. For example, the prostate glands
162 of neonatal rats and mice show abnormal proliferation and cell structure when the pregnant
163 mother is injected with E₂. (Wernert, et al. 1990). This early exposure may imprint
164 intracellular changes by modulating expression pathways of steroid enzymes and receptors
165 as shown in rat models where the response to endogenous androgens and oestrogens becomes
166 abnormal, thus predisposing the animal to prostate cancer after sexual maturation (Rajfer and
167 Coffey 1978). Moreover, studies in mice show that when exposed to high levels of oestrogens
168 in utero, foetal prostate tissue develops abnormalities including intraepithelial neoplasia and
169 predisposition to carcinogenesis in adult life (Prins, et al. 2006). This hypothesis is supported
170 by epidemiological evidence obtained from African-American men having twice as high a
171 risk of developing prostate cancer than comparable Caucasian men which correlates with
172 African-American women having a higher serum oestrogen level during pregnancy compared
173 to Caucasian women (Henderson, et al. 1988).

The second time men are exposed to a high E/T ratio is during old age when serum testosterone decreases, partly due to a dampened HPG axis and partly due to reduced Leydig cell function in the testes. In addition to this, sex hormone-binding globulin (SHBG), which has a higher affinity to testosterone than E_2 (Knochenhauer, et al. 1998), also increases with age which further decreases free serum testosterone relative to free serum E_2 (Samaras, et al. 2012). Furthermore, there is evidence that E_1 and E_2 not only remain at the same level, but in fact increase with age even when accounted for BMI and other metabolic diseases (Jasuja, et al. 2013). While the evidence for an association between serum oestrogen concentration and risk of prostate cancer is unclear and inconsistent, increased serum oestrogen concentrations may stimulate the prostate stroma and epithelia to proliferate and subsequently become neoplastic. Indeed a higher oestrogen:androgen ratio stimulates proliferation of normal prostate stromal (PrSC) and normal epithelial (PrEC) cell lines in vitro (King, et al. 2006).

Another interesting population which is exposed to a high E/T ratio are transsexual male to female individuals. Often in this group of former males, individuals are orchiectomised and then supplemented with anti-androgens to relinquish masculine secondary sex characteristics. They are also supplemented with oestrogens to acquire and enhance feminine characteristics. Their prostates, however, remain unadulterated. A study observing such a cohort of transsexual persons for over 30 years has not identified any increase in risk for prostate cancer (Gooren and Morgentaler 2014). However the study has suggested that when presenting these patients are more likely to be diagnosed with a later stage disease. One limitation admitted by the authors is that the majority of the cohort has not reached the mean age at which prostate cancer is typically diagnosed (Gooren and Morgentaler 2014). Observations made to this cohort over the next two or three decades will be most

198 enlightening in ascertaining whether oestrogens have any significant effects in the
199 development of prostate cancer.

200

201 **Oestrogen Metabolism in Adipose and Prostate Cancer**

202

203 While in pre-menopausal females the primary source of oestrogens are the ovaries, in males
204 there is no central organ which produces substantial quantities of E₂. Instead, peripheral
205 conversion of oestrogen precursors is the main source of oestrogen in men. Local synthesis of
206 E₁ and E₂ is regulated by a plethora of enzymes. DHEA secreted from the zona reticularis of
207 the adrenal glands, and stored in the blood as a reservoir as DHEAS, is the ultimate
208 precursor. Adipose tissue is another notable source of oestrogen synthesis (Cui, et al. 2013).
209 White adipose tissues (the predominant type in obesity) express significant quantities of
210 cytochrome P450 aromatase enzyme (*CYP19A1*) in the abdominal adipose fat of male human
211 samples, which is the final catalyst in the conversion of androgens to oestrogens (Polari, et al.
212 2015; Wang, et al. 2013). There is also a positive correlation between the amount of visceral
213 adipose tissue and serum E₂ levels as shown in a study of 229 man with a mean age of 53.6
214 years where visceral fat was measured using magnetic resonance imaging (Gautier, et al.
215 2013).

216

217 There have been conflicting reports as to whether obesity is a risk factor for prostate cancer
218 as some suggest it decreases risk while others have found the opposite. Allott *et al.* have
219 summarised the findings published between 1991 to 2012 in their review and conclude
220 obesity is associated with aggressive prostate cancer (Allott, et al. 2013). There is further

robust evidence that obese patients are more likely to present with aggressive high-grade prostate cancer (De Nunzio, et al. 2013; Vidal, et al. 2014). It is possible that the risk associated with obesity may in fact be due to elevated circulating oestrogen levels secondary to increased adipose deposition. If this is the case, it would parallel the effects of oestrogen that have been observed in colorectal cancer where oestrogen exposure in the form of hormone replacement therapy or oral contraceptives are initially protective against colorectal cancer but when patients present, they present with a later stage disease (Foster 2013). The intra- and extracellular handling and metabolism of oestrogens within the prostate gland may clarify what effects oestrogens have on tumours. However, studies are lacking regarding the exact intra-tumoural metabolism of oestrogens in prostate cancer cells and human prostate cancer tissue.

Impact of Exogenous Oestrogen on Prostate Cancer

Exogenous oestrogen intake and subsequent availability to the prostate should be considered when determining whether oestrogens affect the development and progression of prostate cancer. A Western diet comprising of high meat, saturated fat, and dairy products has been associated with increased risk of prostate cancer as highlighted by numerous epidemiological studies (Grönberg 2003; Howell 1974; Whittemore, et al. 1995). Additionally, it has been observed that such a Western diet is more likely to cause men diagnosed with prostate cancer to die from the disease when compared to a diet rich in fruits, vegetables, and whole grain cereals (Yang, et al. 2014). Supporting this, it has been widely speculated that dietary oestrogenic compounds from plant sources, termed phytoestrogens, are protective against prostate cancer and are the reason behind lower incidence rates in East Asia where per capita consumption of phytoestrogen-rich foods, such as soya beans, are considerably higher than

245 the Western world (Adlercreutz, et al. 2000; Goetzl, et al. 2007; Strom, et al. 1999). It is
246 possible that phytoestrogens reduce the risk of prostate cancer through multiple mechanisms.
247 In rodent models phytoestrogens can upregulate SHBG synthesis in the liver leading to a
248 higher circulating concentration (Pilšáková, et al. 2010). Increased SHBG is anti-androgenic
249 as it binds to free testosterone with a higher affinity than oestrogens (Knochenhauer et al.
250 1998) implementing a net reduction of testosterone relative to E₂ (Ronde, et al. 2005). This
251 reduction in androgen is thought to be important in the reduction of risk. In addition to
252 chelation of free testosterone via SHBG, phytoestrogens have a negative feedback effect on
253 the HPG axis directly leading to reduced secretion of luteinising hormone and consequently
254 reduced stimulation of androgen and oestrogen synthesis (Goetzl et al. 2007).

255

256 Phytoestrogen compounds are similar enough to endogenous oestrogens to be able to bind to
257 oestrogen receptors (ER) and evoke ligand-specific intracellular responses (Usui 2006).
258 Preference for different types of nuclear ER varies between phytoestrogens (see section on
259 oestrogen receptors). Isoflavones and coumestans are two main categories of phytoestrogens
260 and are structurally similar to E₂ (Figure 1). The prostate cancer cell lines LNCaP and DU145
261 are more sensitive to apoptotic factors when treated with isoflavones *in vitro*. A dose-
262 response relationship between concentration of biochanin A and apoptosis was observed
263 using cytotoxicity and lactate dehydrogenase release assays, flow cytometry and fluorescence
264 microscopy (Szliszka, et al. 2013). Coumestans are able to induce caspase-dependent
265 apoptosis in LNCaP, DU145 and PC3 cells. When treated with wedelolactone, a plant derived
266 coumestan, there was dose-dependent apoptosis in androgen-sensitive cell lines (LNCaP) and
267 androgen-independent cell lines (DU145 and PC3). However, normal non-cancerous PrEC
268 prostate epithelial cells were not affected as harshly showing 90% cell viability compared to
269 circa 20% in cancerous cell lines at concentrations of 30µM. (Sarveswaran, et al. 2012).

While in vitro evidence argues that phytoestrogens are protective against prostate cancer, clinical trials looking at the relationship between consumption of dietary phytoestrogens and progression of prostate cancer have been inconclusive (Goetzl et al. 2007). One double blind randomised control trial in which 81 healthy men were either given a soy protein drink with high isoflavone concentration (83mg/day) or a drink with low isoflavone concentration (3mg/day) showed no significant difference in PSA over 12 months (Adams, et al. 2004). Another trial offering men with confirmed prostate cancer who had either failed medical/surgical therapy or had chosen active surveillance a high dose (450mg/day) oral isoflavone supplement for 6 months showed only a clinically insignificant improvement in PSA in the active surveillance group with no difference in the failed therapy group (deVere White, et al. 2004). Furthermore, a study following up 3628 men with diagnosed prostate cancer for a median duration of 11.5 years showed an increased risk of advanced prostate cancer (HR: 1.62) but a reduced risk of non-advanced prostate cancer (HR: 0.88) in the higher dietary intake of isoflavones group. Dietary intake of phytoestrogens was measured using a validated food frequency questionnaire and so exact doses of phytoestrogens are subject to variation (Reger et al. 2015). This preliminary evidence could infer that dietary phytoestrogens might protect against initiation of prostate cancer, however may promote the progression of advanced prostate cancer.

Steroid metabolism in the prostate

Androgens

The metabolism of oestrogens and oestrogen precursors is important for availability of biologically active E_2 to prostate cancer cells. Oestrogens are synthesised from androgens which themselves are synthesised from progestogens (Khurana 2008). In addition to circulating androgens secreted from the testes, normal prostate tissues have the potential to produce androgens from circulating C19 steroids DHEA and androstenedione (Figure 2). There have been conflicting reports on the possibility of prostate cancer to synthesize androgens *de novo* through the conversion of progestogens via cytochrome P450 17A1 (17-hydroxylase and 17, 20 lyase enzyme [*CYP17A1*]). In prostate cancer, the expression of cytochrome P450 17A1 was reportedly increased in LNCaP and LuCaP cells and human prostate tissue samples ascertained by PCR and immunoblotting (Locke, et al. 2008; Montgomery, et al. 2008); however not all studies support this (Ellem and Risbridger 2009; Hofland, et al. 2010). Although DHT formation from cholesterol was detected using mass spectrometry in castration-resistant prostate cancer (CRPC) models in one study (Locke et al. 2008) these steroid fluxes have not been confirmed quantitatively to date in either *in vitro* or *in vivo* models.

Another key enzyme in the synthesis of biologically active androgens and oestrogens is 3- β -hydroxysteroid dehydrogenase (3 β -HSD) which converts dehydroepiandrosterone and androstenediol to androstenedione and testosterone, respectively (White, et al. 2013). 3 β -HSD is expressed in the normal human prostate, with immunoblotting revealing that the highest concentrations are found in basal epithelial cells (Luu-The, et al. 2008). Certainly, in mouse xenograft studies using the CRPC LAPC4 cell line, expression of 3 β -HSD is increased within the tumour in addition to AKR1C3 and 17 β -HSD3 (Chang, et al. 2011), although its mRNA expression almost completely mutually excludes that of CYP17A1 (Hofland et al. 2010).

318

319 Inhibitors of 3 β -HSD have been explored as an androgen deprivation technique as they are
320 effective in decreasing proliferation in androgen sensitive LNCaP or CRPC cell lines 22Rv1,
321 VCaP and PC346C *in vitro* (Evaul, et al. 2010; Kumagai, et al. 2013). Furthermore,
322 abiraterone was found to inhibit 3 β -HSD activity in addition to CYP17A1 in prostate cancer
323 cell lines and isolated yeast microsomes (Li, et al. 2012). This mechanism might rely on
324 abiraterone being converted to the more active $\Delta(4)$ -abiraterone (D4A) within the prostate
325 gland by 3 β -HSD itself (Li, et al. 2015b). Further research into 3 β -HSD inhibition are
326 currently being pursued, however alternative pathways which bypass androstenedione
327 synthesis exist and so 3 β -HSD function is not strictly necessary.

328

329

330 An alternative pathway has been demonstrated by which synthesis of DHT within the
331 prostate may bypass testosterone and instead be synthesised by reduction of androstenedione
332 by 5 α -reductase SRD5A1 to 5 α -androstenedione which is converted to DHT by 17 β -HSD5.
333 Mass spectrometry has shown that even in patients on anti-androgen therapy with very low
334 serum testosterone levels, intratumoral DHT concentrations remain at the pre-treatment level
335 (Chang et al. 2011; Sharifi and Auchus 2012). 17 β -HSD-5, also known as AKR1C3, appears
336 to be the key enzyme responsible for intratumoural androgen production in CRPC. Its
337 expression in LNCaP, DU145 and PC3 cells are potently stimulated by androgen deprivation
338 *in vitro* and in humans *in vivo* (Ellem and Risbridger 2009; Ellem, et al. 2004) and this
339 secures continued production of testosterone and DHT from circulating adrenal androgens.
340 Local growth factor activin A was shown to be a key intermediate in the castration-induced
341 rise of *AKR1C3* expression levels and intratumoural testosterone production as observed in

342 LNCaP, VCaP and PC3 cells. The concentration of activin A and testosterone were also
343 shown to be increased in the cultured supernatants, as measured by ELISA and mass
344 spectrometry (Hofland, et al. 2011). 17 β -HSD-5 has also been implicated in enzalutamide
345 resistance to anti-androgen therapy. Knockdown of 17 β -HSD-5 using shRNA or inhibition
346 with indomethacin has shown to resensitise enzalutamide-resistant cells *in vitro* and *in vivo*
347 (Liu, et al. 2015).

348 ***Peripheral Oestrogen Metabolism in Prostate Cancer***

349 As mentioned previously, aromatase is a key enzyme required for oestrogen synthesis from
350 androgen precursors. Aromatase converts androstenedione and testosterone to E₁ and E₂,
351 respectively (White et al. 2013). The local synthesis of E₂ within the prostate has previously
352 been debated as not all experiments have identified aromatase expression in normal prostate
353 tissue (Ellem et al. 2004). However, it has been demonstrated in human samples by substrate
354 conversion assays and mass spectrometry that E₂ synthesis does occur in prostate cancer cells
355 (and benign prostatic hyperplasia) via aromatisation (Ellem and Risbridger 2009; Härkönen
356 and Mäkelä 2004). In normal prostate, aromatase is expressed by the stromal tissue but not
357 the epithelial cells, however once malignant, epithelial cells also express aromatase (Ellem
358 and Risbridger 2007). Aberrant expression and activity of aromatase is crucial in the
359 pathophysiology of endometrial and breast cancers where an imbalance of oestrogen is a key
360 factor in tumour growth (Chen 1998; Cunha 1994). As with the developmental similarities
361 between breast and prostate tissues (Ellem and Risbridger 2010), abnormal aromatase activity
362 also plays a major role in breast and prostate tumourigenesis (Ellem and Risbridger 2010).
363 Tumourigenic growth factors including epidermal growth factor and transforming growth
364 factor-1 can modulate aromatase activity in androgen-sensitive LNCaP cells lines leading to
365 decreased oestrogen synthesis (Block, et al. 1996). Furthermore, the expression of aromatase
366 is up to 30-fold greater in metastatic prostate cancer compared to primary tumours

(Miftakhova, et al. 2016). In addition, overexpression of aromatase increased the progression of bony metastasis in xenograft experiments where nude mice were injected with PC3 cell lines transfected to overexpress aromatase (Miftakhova, et al. 2016). Consequently, the use of aromatase inhibitors for the treatment of prostate cancer has been investigated many times in patient cohorts. The first generation aromatase inhibitor aminoglutethimide is non-selective and showed poor objective responses including serum PSA levels and disease stability in some studies while showing a significant increase in survival in others (Santen, et al. 1997). One study treated 58 castrated men with advanced prostate cancer resistant to conventional therapy with 500-750mg daily aminoglutethimide; 11 men showed an objective response with a mean remission of 10 months and a further two showed disease stabilisation for a mean seven months (Murray and Pitt 1985). The second generation aromatase inhibitor, 4-hydroxyandrostenedione showed good subjective responses in 18 out of 25 patients with advanced CRPC, particularly alleviation of bone pain in prostate metastases. However the objective responses were still poor with a reduction in tumour volume seen in only three patients and all patients progressed to have skeletal metastasis. (Davies, et al. 1992). A Phase II clinical study looking at the effects of oral letrozole, a third-generation aromatase inhibitor more commonly used in the treatment of hormone-dependent breast cancer, in 43 men with CRPC showed no significant disease regression with serum PSA decreasing by more than 50% in only one patient and decreasing by less than 50% in one further patient (Smith, et al. 2002). A very similar conclusion was drawn from clinical studies looking at anastrozole, another third generation aromatase inhibitor, where out of 14 patients with CRPC none showed a decrease in serum PSA and mild bone pain relief was reported by only two patients (Santen, et al. 2001). While aromatase is of utmost importance in local oestrogen synthesis, it appears as though therapeutic approaches targeting aromatase may be futile in treating prostate cancer. An alternative possibility is that E₂ is not synthesised from androgens within

the prostate but instead is converted from systemic sulphated E_1 within the prostate via steroid sulphotase (STS).

STS is widely expressed in almost all peripheral tissues and is responsible for hydrolysing sulphate moieties off of circulating sulphate-conjugated steroids in order to make them biologically active (Mueller, et al. 2015). Oestrone sulphate (E_1S) is the most abundant circulating oestrogen in adult humans (Muir, et al. 2004) with plasma levels between 2-4nmol/L in men (Mueller et al. 2015) and while oestradiol sulphate also exists, plasma levels are very low. Furthermore, serum E_1S levels have been correlated with increased risk of prostate cancer. In a cohort study of 5995 men aged over 65 where the mean serum E_1S levels in the 275 patients who developed prostate cancer was significantly higher than those who did not develop prostate cancer (Daniels, et al. 2010).

Before sulphated oestrogens can be unconjugated by intracellular STS, transport of sulphated oestrogens into cells requires the expression of organic anion transporter peptides (OATP) (Raftogianis, et al. 2000) and indeed several different OATPs involved in the transport of oestrone sulphate are expressed in prostate cancers (Buxhofer-Ausch, et al. 2013; Giton, et al. 2015; Wright, et al. 2011). STS has been shown to be expressed in normal human prostate tissue (Reed, et al. 2005), prostate cancer cell lines LNCaP, DU-145 and PC3 (Nakamura, et al. 2006) and in primary prostate homogenates (Klein, et al. 1989). Furthermore, one study found that STS is expressed in the majority of localised prostate cancers showing higher expression in malignant tissues compared to benign (Nakamura, et al. 2006). The activity of STS has been proven within the human prostate for the desulphation of dehydroepiandrosterone sulphate (DHEAS) into DHEA, an androgen precursor (Farnsworth

1973). Moreover, E_1 synthesis from desulphation of E_1S within the prostate is putatively 10-fold greater than synthesis via aromatase (Nakamura et al. 2006). The relevance of STS in cancer has been more extensively studied in breast cancer where there is significantly higher expression of STS than in normal breast (Utsumi, et al. 2000). Consequently, several STS inhibitors have been developed for the treatment of breast cancer, some of which have shown early promise (Stanway, et al. 2006). Moreover, first and second generation STS inhibitors have been effective pre-clinically against breast cancer (Foster et al. 2006; Foster et al. 2008; Purohit and Foster 2012). Meanwhile, investigations into the efficacy of STS inhibitors in prostate cancer have been undertaken. It has been observed that middle-aged rats treated with oral STS inhibitor, STX64 decreased conversion of E_1S to E_1 (Giton et al. 2015; Roy, et al. 2013). Neither study presented evidence of STS inhibition affecting any proliferative markers of proliferation, however the latter study did demonstrate that STS inhibition in middle-aged rats prevented increase of prostate mass when treated with E_1S + STX64 vs E_1S alone where prostate mass increased (Giton et al. 2015). An alternative conjugate of circulating oestrogens is glucuronide (Raftogianis et al. 2000), however, research into oestrogen glucuronide transport into prostate cells and evidence of glucuronidase enzymes within the prostate is lacking.

Conversion of E_1 to E_2 (and androstenedione to testosterone) requires 17-beta-hydroxysteroid dehydrogenase (17β -HSD) enzymes (White et al. 2013). 17β -HSDs enzymes are alcohol oxidoreductases which catalyse reduction (E_1 to E_2) and oxidation (E_2 to E_1) at carbon atom 17. There are over 14 different isozymes of 17β -HSDs (17β -HSD 1-14) and certain 17β -HSDs have a higher propensity to catalyse the reaction in a certain direction, for example 17β -HSD-1 favours reduction whereas 17β -HSD-2 favours oxidation (Lukacik, et al. 2006; Oduwale, et al. 2003). 17β -HSDs play an important role in hormone sensitive cancers.

441 Increased expression of 17 β -HSD-1 in breast cancers of post-menopausal women helps
442 maintain high intratumoural E₂ levels (Lukacik et al. 2006). Moreover, expression of 17 β -
443 HSD-2 and 17 β -HSD-3 mRNA is significantly higher in malignant prostatic tissues
444 compared to normal prostate tissues (Day, et al. 2013) with one study reporting prostate
445 cancer biopsies showing 30-fold higher mRNA expression than normal. In addition to
446 converting androstenedione to testosterone, 17 β -HSD 5 can convert E₁ to E₂. Inhibitors of
447 17 β -HSD 5 have been explored in castration-resistant prostate cancer and breast cancer, in
448 the latter where androgens are not considered to play an important role (Adeniji, et al. 2013).
449 The study found no appreciable decrease in E₂ synthesis in breast cancer cell lines when
450 treated with a 17 β -HSD 5 inhibitor and only a moderate decrease in E₂ synthesis in some
451 subpopulations of prostate cancer cell lines. Interestingly, inflammation associated with
452 tumours modulates the expression of 17 β -HSD-2 and 17 β -HSD-5 (and also 3 β -HSD).
453 Treatment of prostate cancer stromal cell lines PrSC with TGF β 1 showed a marked down-
454 regulation in mRNA expression of 17 β -HSD-2 and 17 β -HSD-5 in a dose-dependent manner
455 (Piao, et al. 2013). The counterintuitive action of TGF β 1 again demonstrates how little is
456 understood about oestrogenic pathways in prostate cancer. Regardless of the mechanisms by
457 which oestrogens become available within the prostate gland, tumour-promoting or tumour-
458 suppressing effects must be mediated by activation of oestrogen receptors (ER).

459

460 **Oestrogen receptors (ER) in the prostate**

461 The effects of oestrogens on tissues are mediated via activation of oestrogen receptors (ER).
462 There are two well studied ERs; ER alpha (ER α) and ER beta (ER β) encoded by two separate
463 genes *ESR1* and *ESR2*, respectively. ER α and ER β are members of the nuclear receptor
464 superfamily (Robinson-Rechavi, et al. 2003). When bound and activated, ERs interact

directly with the genome acting as transcription factors (or activating transcription factors) which act directly on oestrogen response elements (Deblois and Giguere 2013). As well as E_2 , ERs can be stimulated by phytoestrogens, and different classes of phytoestrogens have selected preferences for each type of ER. In general, phytoestrogens show agonistic activity towards $ER\beta$ at lower concentrations than towards $ER\alpha$ using hamster uterine cells (Takeuchi, et al. 2009). When human cells are examined, the relative binding affinity (RBA) of genistein to $ER\beta$ is approximately 20-30 times greater than for $ER\alpha$ as shown in MCF-7 breast cancer cell lines (Pilšáková et al. 2010). The affinity of phytoestrogens for ER widely varies with most molecules having an RBA to $ER\beta$ 1000-fold lower than E_2 . However, molecules such as genistein and coumesterol have an RBA 100-fold lower than E_2 . Genistein and coumesterol are able to activate transcriptional activities of $ER\alpha$ and $ER\beta$ at concentrations of 1-10nM compared to physiological E_2 concentrations of 20-40pM in males (Kuiper, et al. 1998; Mueller et al. 2015). Of course, the ability of phytoestrogens to bind to ER also depends on the existing levels of E_1 and E_2 as these molecules are direct competitors with phytoestrogens.

ERs have been studied more extensively in the context of breast cancers, a neoplasm that has been likened as the sister disease to prostate cancer, especially in regards to their hormonal responses and sensitivities (Risbridger, et al. 2010). In breast cancer, activation of $ER\alpha$ promotes tumour growth as it initiates anti-apoptotic (Chaudhri, et al. 2014; Razandi, et al. 2000) and mitogenic effects (Bhatt, et al. 2012; Yamnik and Holz 2010). This anti-apoptotic effect of $ER\alpha$ makes $ER\alpha$ positive breast cancers more likely to metastasise (Ross-Innes, et al. 2012). In fact, a review of ERs in breast and ovarian cancers has found $ER\alpha$ expression correlates with worse prognosis whereas $ER\beta$ expression correlates with better clinical

489 outcomes (Burns and Korach 2012). Generally, ER α activation promotes proliferative
 490 pathways whereas ER β activation leads to apoptotic pathways (Acconcia, et al. 2005).

491

492 Expression of ER α and ER β in the normal prostate has been determined as the role of
 493 oestrogens in prostatic development was identified (Ho 2004). Recently it has been reported
 494 that prostate progenitor stem cells, while lacking expression of androgen receptor, express ER
 495 abundantly. Indeed, the expression of ER β is putatively 6-fold greater and ER α 125-fold
 496 greater in progenitor cells compared to LNCaP mature cells (Di Zazzo, et al. 2016). Although
 497 this supports the importance of oestrogens in embryonic and neonatal development of
 498 prostate gland, it has been hypothesised that lack of androgen receptor expression could be an
 499 imprint which later predisposes to CRPC in the elderly. In non-cancerous prostate ER α is
 500 predominantly expressed in the stromal compartment and ER β is predominantly expressed in
 501 basal-epithelial cells. However in prostate cancer, ER α expression is down-regulated in
 502 stromal cells and upregulated in the cancerous epithelial cells. ER β expression is down-
 503 regulated in epithelial cells as seen by immunostaining in human prostate tissue (Yeh, et al.
 504 2014). Indeed there is evidence that down-regulation of ER β promotes activation of NF- κ B
 505 mediated by hypoxia-inducible factor 1 (HIF-1). In immortalised normal prostate epithelial
 506 cell line PNT1a, loss of ER β using shRNA showed an increase in NF- κ B mRNA expression
 507 and activity. This mirrors what is seen in high grade, late stage prostate cancer (Mak, et al.
 508 2015). Consequently, it appears that an increase in ER α expression and decrease in ER β
 509 expression is what shifts the balance between protective effects of oestrogens and
 510 proliferative effects of oestrogens as has been suggested in other cancers (Barzi, et al. 2013;
 511 Burns and Korach 2012). Figure 3 summarises the difference in ER α and ER β expression
 512 between non-cancerous and cancerous prostate tissue. Single nucleotide polymorphisms
 513 (SNP) in the ER genes have been investigated and associations have been made between

certain polymorphisms and the risk of prostate cancer (Holt, et al. 2013; Jurečková, et al. 2015). In both studies, the genomes from histologically confirmed human prostate cancer samples were analysed using polymerase chain reaction restriction fragment length polymorphism (PCR-RFLP) based analysis and compared to age-matched healthy control subjects. A meta-analysis exploring the results of 24 published studies that include Caucasian, Asian and African participants concluded that *ESR1* rs9340799 polymorphism is allied to increased risk in the general population of Caucasians and Africans whereas *ESR2* rs1256049 polymorphisms has been linked to increased risk only in Caucasians (Fu, et al. 2014).

Research into ER β has been more extensive than in ER α . McPherson et al. (2007) highlighted the potential significance of ER β manipulation when they treated prostate hyperplasia in oestrogen depleted mice with a selective ER β agonist and found it to induce apoptosis and shrink the size of the prostate. Hussain et al. (2012) carried forward this research and initial studies have found ER β agonist treatment with 8 β -VE₂ can induce apoptosis in primary human and murine prostatic basal cells, a lineage considered to be the cells of origin for prostate cancers (Taylor, et al. 2012). The mechanism behind how ER β activation induces apoptosis in prostate cancer cells lines may be via up-regulation of p53-upregulated modulator of apoptosis (PUMA) and consequent intrinsic caspase-9 mechanisms. Dey, et al. overexpressed ER β in LNCaP, PC3 and 22Rv1 prostate cancer cell lines *in vitro*, the latter which does not express ER β , and treated with E₂ and agonist 3 β -adiol. Immunofluorescence revealed that cells which expressed ER β were more likely to undergo apoptosis following expression of PUMA independent of p53 (Dey, et al. 2014). (Dey, et al. 2014). It has even been reported that ER β activation impedes on the epithelial-mesenchymal transition process thereby reducing the risk of invasion and metastasis. In human tissue samples and LNCaP

539 and PC3 cell lines, treatment with E₂ and high concentration of ERβ1 agonist 3β-adiol
540 resulted in inhibition of VEGF and destabilisation of HIF-1 in vitro thus suppressing the
541 factors that drive epithelial-mesenchymal transition necessary for metastasis. Furthermore,
542 loss of ERβ1 expression by means of shRNA transfection resulted in significant increase in
543 migration and invasion (Mak, et al. 2010). Mounting evidence also suggests that
544 pharmaceutical targeting of ERβ pathways may be effective in treating prostate cancer.
545 However, recently a 'switching roles' theory has been proposed suggesting the effects of ERβ
546 activation switches from protective to proliferative as cancer progresses (Savoy and Ghosh
547 2013). The theory is based on the observation that castration-resistant prostate cancers have
548 higher expression of ERβ compared to hormone-naïve prostate cancers. It is possible that
549 decreased levels of circulating androgens and up-regulation of androgen receptors may be
550 important in this switch however the actual mechanisms and processes are yet unknown.

551

552 Splice variants of ERβ are also important as it has been shown that at least 5 different
553 isoforms exist, many of which are expressed in the prostate (Leung, et al. 2006). Activation
554 of different isoforms may have opposing effects; for example ERβ₁ is tumour-suppressing
555 whereas ERβ₂ is tumour-promoting in LNCaP cells (Chen, et al. 2009). In a study of primary
556 prostate cancer samples from 144 patients who underwent radical prostatectomy, two
557 particular isoforms ERβ₂ and ERβ₅ have been identified to promote invasion and metastasis
558 of prostate cancer and thus correlate with worse outcomes while others continue to be studied
559 (Leung, et al. 2010; Nelson, et al. 2014). Certain ERβ isoforms, such as ERβ₂ and ERβ₃,
560 when activated interact with transcription factors which enable and promote the epithelial
561 mesenchyme transition and hence might be why advanced prostate cancers have higher
562 expression of ERβ (Leung et al. 2010). More research needs to be carried out to understand
563 the mechanisms of the complex downstream pathways of ERβ activation in prostate cancer.

564

565 The tumour promoting effects of ER α within the prostate are not as well defined. ER α is
566 expressed in significant quantities in the stromal tissue of prostate cancer where they have
567 been associated with cancer-associated fibroblasts (CAF) (Slavin, et al. 2015). Da, et al.
568 isolated CAF from adenocarcinoma of mouse prostate lentivirally transduced ER α .
569 Conditioned media from ER α + CAF promoted proliferation of LNCaP, PC3, C4-2 and
570 22Rv1 cells. Furthermore, in xenograft experiments mice co-implanted with ER α + CAF
571 showed a higher growth rate of tumour mass compared to injection of prostate cancer cell
572 lines alone (Da, et al. 2015). Activation of ER α on CAFs stimulates the release of tumour-
573 promoting factors which act on prostate epithelia in a paracrine manner. Slug (SNAI2), a
574 transcription factor with anti-apoptotic pathways can repress ER α expression by binding to
575 gene promotor regions and consequently promote epithelial-mesenchymal transition in
576 prostate cancer cells and human breast cancer samples (Li, et al. 2015a). In contrast,
577 downstream pathways of ER α activation can inhibit metastasis by down-regulating
578 expression of matrix metalloproteinase 3 and upregulating expression of thrombospondin 2
579 as seen in a range of breast cancer cell lines and LNCaP cell line, however this is not
580 evidence in primary human prostate tissue (Li et al. 2015a). This may be an effect of ER α
581 activation which diverts cell resources towards growth of prostate cancer rather than spread
582 and invasion (Hanahan and Weinberg 2011). A study investigating the role of ER α in prostate
583 cancers of PTEN-deficient mice has shown expression of ER α correlates strongly with the
584 expression of Ki67,- a proliferative marker. In addition, inhibition and knockdown of ER α
585 decreases proliferation but has no effect on cell viability thus the tumour mass remained
586 static. This further demonstrates that ER α regulates cell proliferation through PI3K and
587 MAPK signalling (Takizawa, et al. 2015).

588

Human trials in 1590 men with high grade intraepithelial neoplasia of the prostate has shown no significant decrease in risk of prostate cancer when treated with daily toremifene, a selective oestrogen receptor modulator (SERM) used for the treatment of metastatic breast cancer, compared with placebo. Of the 1467 men who underwent a biopsy during the three-year study, cancer was detected in 34.7% in the placebo group compared to 32.3% in the treatment group ($p= 0.39$) (Taneja, et al. 2013). Conversely, experimental use of toremifene, in cell lines and nude mice models have suggested that ER α antagonists can repress the tumorigenicity of prostate cancer (Hariri, et al. 2015). Intriguingly, there is recent evidence that abiraterone, used frequently in advanced prostate cancer is able to activate ER. Capper, et al. demonstrated an increase in proliferation of MCF-7 and T47D breast cancer cell lines when treated with abiraterone. The proliferative effects were diminished when the cells were treated with ER antagonist ICI 182,78 (Capper, et al. 2016). ER-mediated progression of prostate cancer might thus constitute a novel mechanism of resistance to abiraterone that warrants further investigation. The signalling mechanisms of ER α and ER β are summarised in Figure 4.

In addition to the two nuclear ERs, ER α and ER β , another relatively recently discovered ER exists. G-protein coupled oestrogen receptor (GPER), alternatively known as GPR30, is a membrane-bound receptor discovered in 1998 (O'Dowd, et al. 1998). GPER is found in 50% of breast cancers and is believed to be critically involved in how Tamoxifen (a SERM) resistance is developed (Mo, et al. 2013). Tamoxifen can bind and stimulate GPER in breast cancer (Prossnitz, et al. 2008a) activating downstream cancer promoting pathways. GPER has also been shown to be expressed in various hormone-sensitive tissues in the body including the prostate (Prins and Hu 2013; Prossnitz, et al. 2007) and has very similar affinity for E₂ as ER α and ER β with almost no interaction with androgens or glucocorticoids (Prossnitz, et al.

2008b). In addition to being activated by endogenous E₂, GPER can also be activated by phytoestrogens with similar RBA as phytoestrogens have to ER β and elicit an oestrogenic signalling pathways (Thomas and Dong 2006).

Evidence of changes in GPER expression within prostate cancer is scarce, though it has been established with immunofluorescence and immunoblotting that GPER is expressed LNCaP, DU145 and PC3 cells which have varying degrees of invasiveness (Maier, et al. 2006). In addition, expression of GPER has been identified by immunohistochemistry and immunoblotting in prostate adenocarcinomas and in pre-neoplastic lesions in 50 patients with confirmed prostate cancer of varying grades of aggressiveness and in 5 patients with benign prostatic disease (Rago, et al. 2016). Naturally, more research has been conducted in aggressive cell lines and primary tissues. In contrast to the effects of GPER activation in breast and ovarian cancers where it promotes growth, it has been identified that treatment of castration-resistant prostate cancer with a specific GPER agonist, G1, actually inhibits the growth of prostate cancer in PC-3, DU145 and LNCaP cell lines *in vitro* and *in vivo* PC3 xenografts (Chan, et al. 2010; Lam, et al. 2014). While most studies only reported tumour inhibition in castration-resistant cell lines, Lam et al. found that G1 treatment has no effect on androgen-sensitive LNCaP cells *in vitro* and *in vivo* xenograft mouse models whereas it had a significant effect on castration-resistant tumours without apparent toxicity to the host (Lam et al. 2014). Furthermore, GPER expression is significantly increased in androgen-deprived environments compared to androgen-replete milieus (Prins and Hu 2013) with increased GPER expression also evident in cells isolated from distant metastases in patients with CRPC compared to tissue from primary prostate cancers (Lam et al. 2014). Androgen receptor activation downregulates GPER expression thus explaining why expression of GPER is greater in androgen deprived environments (Lam et al. 2014). The mechanisms by

which the GPER agonist G1 has anti-tumour effects has been explored in PC3 cell line *in vitro* and *in vivo* xenograft castrated mice models and is reported to be via up-regulation of p21 and consequent cell cycle arrest at G2 phase (Chan et al. 2010). Although GPER activation inhibits growth of prostate cancer, it increases proliferation of other tissues including testicular germ cells and urothelial cells of the bladder and urinary tract (Chevalier, et al. 2011; Huang, et al. 2015). The fact that GPER activation can have opposing effects in different tissues through the same pathway illustrates the complexity of intracellular oestrogen signalling. Figure 4 grossly summarises GPER signalling pathways that have thus far been identified in prostate cancer.

Conclusion

This review has presented evidence that suggests an imbalance of circulating oestrogens and androgens may be responsible for changes to the development and progression of prostate cancer. In addition to endogenous oestrogen availability, exposure to exogenous oestrogens in the form of phytoestrogens may also have a profound effect. However, there is substantial evidence that intratumoural synthesis of oestrogens, and indeed androgens, plays a significant role as the prostate is endowed with the ability to express key enzymes required for oestrogen synthesis. There is a relationship between stage of disease and level of expression of these enzymes, as is evident from the emergence of resistance to anti-androgen therapy further supports this hypothesis.

Changes in the expression pattern of ER α and ER β greatly affect whether oestrogens are tumour promoting or tumour suppressing. In normal prostate and during early stages of

prostate cancer where ER β is the prominent ER, oestrogens may be beneficial as ER β activation initiates apoptotic pathways. Perhaps this is why a lifetime of increased phytoestrogen consumption can reduce the risk of prostate cancer development. In late stage prostate cancer where ER α is the dominating ER within the prostate, oestrogens are deleterious as ER α activation regulates cell proliferation through PI3K and MAPK signalling. Activation of GPER inhibits growth of prostate cancer however, GPER is not uniformly expressed in all prostate cancer and thus any GPER targeted therapy will be of benefit to a limited number of patients. Figure 5 summarises how the expression of ERs change during the progression of prostate cancer.

Before any definitive conclusions can be drawn over whether oestrogens are good or bad for prostate cancer, further research has to be conducted exploring the signalling pathways of ER within prostate tissue. In addition an understanding of the mechanisms behind abiraterone (Romanel, et al. 2015) and enzalutamide resistance (Claessens, et al. 2014), and whether this is linked to altered androgen and oestrogen metabolism, will be required before the next big step is taken towards development of endocrine therapy for prostate cancer.

References

Acconcia F, Totta P, Ogawa S, Cardillo I, Inoue S, Leone S, Trentalance A, Muramatsu M & Marino M 2005 Survival versus apoptotic 17 β -estradiol effect: Role of ER α and ER β activated non-genomic signaling. *Journal of Cellular Physiology* **203** 193-201.

Adams KF, Chen C, Newton KM, Potter JD & Lampe JW 2004 Soy Isoflavones Do Not Modulate Prostate-Specific Antigen Concentrations in Older Men in a Randomized Controlled Trial. *Cancer Epidemiology Biomarkers & Prevention* **13** 644-648.

Adeniji AO, Chen M & Penning TM 2013 AKR1C3 as a target in castrate resistant prostate cancer. *The Journal of Steroid Biochemistry and Molecular Biology* **137** 136-149.

Adlercreutz H, Mazur W, Bartels P, Elomaa V-V, Watanabe S, Wähälä K, Landström M, Lundin E, Bergh A, Damber J-E, et al. 2000 Phytoestrogens and Prostate Disease. *The Journal of Nutrition* **130** 658.

Allen NE, Appleby PN, Davey GK & Key TJ 2002 Lifestyle and nutritional determinants of bioavailable androgens and related hormones in British men. *Cancer Causes & Control* **13** 353-363.

Allott EH, Masko EM & Freedland SJ 2013 Obesity and Prostate Cancer: Weighing the Evidence. *European Urology* **63** 800-809.

Barazani Y, Katz BF, Nagler HM & Stember DS 2014 Lifestyle, Environment, and Male Reproductive Health. *Urologic Clinics of North America* **41** 55-66.

Barzi A, Lenz AM, Labonte MJ & Lenz H-J 2013 Molecular Pathways: Estrogen Pathway in Colorectal Cancer. *Clinical cancer research : an official journal of the American Association for Cancer Research* **19** 10.1158/1078-0432.CCR-1113-0325.

Beer TM, Armstrong AJ, Rathkopf DE, Loriot Y, Sternberg CN, Higano CS, Iversen P, Bhattacharya S, Carles J, Chowdhury S, et al. 2014 Enzalutamide in Metastatic Prostate Cancer before Chemotherapy. *New England Journal of Medicine* **371** 424-433.

Bhatt S, Xiao Z, Meng Z & Katzenellenbogen BS 2012 Phosphorylation by p38 Mitogen-Activated Protein Kinase Promotes Estrogen Receptor α Turnover and Functional Activity via the SCF(Skp2) Proteasomal Complex. *Molecular and Cellular Biology* **32** 1928-1943.

Block JL, Block NL & Lokeshwar BL 1996 Modulation of aromatase activity by growth factors in an androgen sensitive human prostate cancer cell line, LNCaP. *Cancer Letters* **102** 167-172.

Bosset P-O, Albiges L, Seisen T, de la Motte Rouge T, Phé V, Bitker M-O & Rouprêt M 2012 Current role of diethylstilbestrol in the management of advanced prostate cancer. *BJU International* **110** E826-E829.

Brooke GN & Bevan CL 2009 The Role of Androgen Receptor Mutations in Prostate Cancer Progression. *Current Genomics* **10** 18-25.

Bulun SE, Economos K, Miller D & Simpson ER 1994 CYP19 (aromatase cytochrome P450) gene expression in human malignant endometrial tumors. *The Journal of Clinical Endocrinology & Metabolism* **79** 1831-1834.

- 735 Burns KA & Korach KS 2012 Estrogen receptors and human disease: an update. *Archives of*
 736 *Toxicology* **86** 1491-1504.
- 737
- 738 Buxhofer-Ausch V, Secky L, Wlcek K, Svoboda M, Kounnis V, Briasoulis E, Tzakos AG, Jaeger W &
 739 Thalhammer T 2013 Tumor-specific expression of organic anion-transporting polypeptides:
 740 transporters as novel targets for cancer therapy. *Journal of drug delivery* **2013** 863539.
- 741
- 742 Capper CP, Larios JM, Sikora MJ, Johnson MD & Rae JM 2016 The CYP17A1 inhibitor abiraterone
 743 exhibits estrogen receptor agonist activity in breast cancer. *Breast Cancer Research and Treatment*
 744 1-8.
- 745
- 746 Center MM, Jemal A, Lortet-Tieulent J, Ward E, Ferlay J, Brawley O & Bray F 2012 International
 747 Variation in Prostate Cancer Incidence and Mortality Rates. *European Urology* **61** 1079-1092.
- 748
- 749 Chan QK, Lam H-M, Ng C-F, Lee AY, Chan ES, Ng H-K, Ho S-M & Lau K-M 2010 Activation of GPR30
 750 inhibits the growth of prostate cancer cells through sustained activation of Erk1/2, c-jun/c-fos-
 751 dependent upregulation of p21, and induction of G2 cell-cycle arrest. *Cell Death & Differentiation* **17**
 752 1511-1523.
- 753
- 754 Chang K-H, Li R, Papari-Zareei M, Watumull L, Zhao YD, Auchus RJ & Sharifi N 2011
 755 Dihydrotestosterone synthesis bypasses testosterone to drive castration-resistant prostate cancer.
 756 *Proceedings of the National Academy of Sciences* **108** 13728-13733.
- 757
- 758 Chen CD, Welsbie DS, Tran C, Baek SH, Chen R, Vessella R, Rosenfeld MG & Sawyers CL 2004
 759 Molecular determinants of resistance to antiandrogen therapy. *Nat Med* **10** 33-39.
- 760
- 761 Chen M, Ni J, Chang H-C, Lin C-Y, Muyan M & Yeh S 2009 CCDC62/ERAP75 functions as a coactivator
 762 to enhance estrogen receptor beta-mediated transactivation and target gene expression in prostate
 763 cancer cells. *Carcinogenesis* **30** 841-850.
- 764
- 765 Chen M, Yeh C-R, Chang H-C, Vitkus S, Wen X-Q, Bhowmick NA, Wolfe A & Yeh S 2012 Loss of
 766 epithelial oestrogen receptor α inhibits oestrogen-stimulated prostate proliferation and squamous
 767 metaplasia via in vivo tissue selective knockout models. *The Journal of Pathology* **226** 17-27.
- 768
- 769 Chen S 1998 Aromatase and breast cancer. In *Frontiers in bioscience : a journal and virtual library*, pp
 770 d922-933.
- 771
- 772 Chevalier N, Bouskine A & Fenichel P 2011 Role of GPER/GPR30 in tumoral testicular germ cells
 773 proliferation. *Cancer biology & therapy* **12** 2-3.
- 774
- 775 Christoforou P, Christopoulos PF & Koutsilieris M 2014 The Role of Estrogen Receptor β in Prostate
 776 Cancer. *Molecular Medicine* **20** 427-434.
- 777
- 778 Claessens F, Helsen C, Prekovic S, Van den Broeck T, Spans L, Van Poppel H & Joniau S 2014 Emerging
 779 mechanisms of enzalutamide resistance in prostate cancer. *Nat Rev Urol* **11** 712-716.
- 780
- 781 Clemons J, Glodé LM, Gao D & Flaig TW 2013 Low-dose diethylstilbestrol for the treatment of
 782 advanced prostate cancer. *Urologic Oncology: Seminars and Original Investigations* **31** 198-204.
- 783

- 784 Cook LS, Goldoft M, Schwartz SM & Weiss NS 1999 INCIDENCE OF ADENOCARCINOMA OF THE
785 PROSTATE IN ASIAN IMMIGRANTS TO THE UNITED STATES AND THEIR DESCENDANTS. *The Journal of*
786 *Urology* **161** 152-155.
- 787
- 788 Cui J, Shen Y & Li R 2013 Estrogen synthesis and signaling pathways during ageing: from periphery to
789 brain. *Trends in molecular medicine* **19** 197-209.
- 790
- 791 Cunha GR 1994 Role of mesenchymal-epithelial interactions in normal and abnormal development
792 of the mammary gland and prostate. *Cancer* **74** 1030-1044.
- 793
- 794 Da J, Lu M & Wang Z 2015 Estrogen Receptor Alpha (ER α)-Associated Fibroblasts Promote Cell
795 Growth in Prostate Cancer. *Cell Biochemistry and Biophysics* **73** 793-798.
- 796
- 797 Daniels NA, Nielson CM, Hoffman AR & Bauer DC 2010 SEX HORMONES AND THE RISK OF INCIDENT
798 PROSTATE CANCER. *Urology* **76** 1034-1040.
- 799
- 800 Davies JH, Dowsett M, Jacobs S, Coombes RC, Hedley A & Shearer RJ 1992 Aromatase inhibition: 4-
801 hydroxyandrostenedione (4-OHA, CGP 32349) in advanced prostatic cancer. *British Journal of Cancer*
802 **66** 139-142.
- 803
- 804 Day JM, Foster PA, Tutill HJ, Schmidlin F, Sharland CM, Hargrave JD, Vicker N, Potter BVL, Reed MJ &
805 Purohit A 2013 STX2171, a 17 β -hydroxysteroid dehydrogenase type 3 inhibitor, is efficacious in vivo
806 in a novel hormone-dependent prostate cancer model. *Endocrine-Related Cancer* **20** 53-64.
- 807
- 808 de Bono JS, Logothetis CJ, Molina A, Fizazi K, North S, Chu L, Chi KN, Jones RJ, Goodman OBJ,
809 Saad F, et al. 2011 Abiraterone and Increased Survival in Metastatic Prostate Cancer. *New England*
810 *Journal of Medicine* **364** 1995-2005.
- 811
- 812 De Nunzio C, Albisinni S, Freedland SJ, Miano L, Cindolo L, Finazzi Agrò E, Autorino R, De Sio M,
813 Schips L & Tubaro A 2013 Abdominal obesity as risk factor for prostate cancer diagnosis and high
814 grade disease: A prospective multicenter Italian cohort study. *Urologic Oncology: Seminars and*
815 *Original Investigations* **31** 997-1002.
- 816
- 817 Deblois G & Giguere V 2013 Oestrogen-related receptors in breast cancer: control of cellular
818 metabolism and beyond. *Nat Rev Cancer* **13** 27-36.
- 819
- 820 deVere White RW, Hackman RM, Soares SE, Beckett LA, Li Y & Sun B 2004 Effects of a genistein-rich
821 extract on PSA levels in men with a history of prostate cancer. *Urology* **63** 259-263.
- 822
- 823 Dey P, Strom A & Gustafsson JA 2014 Estrogen receptor [beta] upregulates FOXO3a and causes
824 induction of apoptosis through PUMA in prostate cancer. *Oncogene* **33** 4213-4225.
- 825
- 826 Di Zazzo E, Galasso G, Giovannelli P, Di Donato M, Di Santi A, Cernera G, Rossi V, Abbondanza C,
827 Moncharmont B, Sinisi AA, et al. 2016 Prostate cancer stem cells: the role of androgen and estrogen
828 receptors. *Oncotarget* **7** 193-208.
- 829
- 830 Ellem SJ & Risbridger GP 2007 Treating prostate cancer: a rationale for targeting local oestrogens.
831 *Nat Rev Cancer* **7** 621-627.
- 832
- 833 Ellem SJ & Risbridger GP 2009 The Dual, Opposing Roles of Estrogen in the Prostate. *Annals of the*
834 *New York Academy of Sciences* **1155** 174-186.

835
836 Ellem SJ & Risbridger GP 2010 Aromatase and regulating the estrogen:androgen ratio in the prostate
837 gland. *The Journal of Steroid Biochemistry and Molecular Biology* **118** 246-251.
838
839 Ellem SJ, Schmitt JF, Pedersen JS, Frydenberg M & Risbridger GP 2004 Local Aromatase Expression in
840 Human Prostate Is Altered in Malignancy. *The Journal of Clinical Endocrinology & Metabolism* **89**
841 2434-2441.
842
843 Evaul K, Li R, Papari-Zareei M, Auchus RJ & Sharifi N 2010 3 β -Hydroxysteroid Dehydrogenase Is a
844 Possible Pharmacological Target in the Treatment of Castration-Resistant Prostate Cancer.
845 *Endocrinology* **151** 3514-3520.
846
847 Farnsworth WE 1973 Human prostatic dehydroepiandrosterone sulfate sulfatase. *Steroids* **21** 647-
848 664.
849
850 Foster PA 2013 Oestrogen and colorectal cancer: mechanisms and controversies. *International*
851 *Journal of Colorectal Disease* **28** 737-749.
852
853 Foster PA, Chander SK, Parsons MF, Newman SP, Woo LW, Potter BV, Reed MJ & Purohit A 2008
854 Efficacy of three potent steroid sulfatase inhibitors: pre-clinical investigations for their use in the
855 treatment of hormone-dependent breast cancer. *Breast Cancer Res Treat* **111** 129-38.
856
857 Foster PA, Newman SP, Chander SK, Stengel C, Jhalli R, Woo LL, Potter BV, Reed MJ & Purohit A 2006
858 In vivo efficacy of STX213, a second-generation steroid sulfatase inhibitor, for hormone-dependent
859 breast cancer therapy. *Clin Cancer Res* **12** 5543-9.
860
861 Francis JC, Thomsen MK, Taketo MM & Swain A 2013 β -Catenin Is Required for Prostate
862 Development and Cooperates with *Pten* Loss to Drive Invasive Carcinoma. *PLoS*
863 *Genet* **9** e1003180.
864
865 Fu C, Dong W-Q, Wang A & Qiu G 2014 The influence of ESR1 rs9340799 and ESR2 rs1256049
866 polymorphisms on prostate cancer risk. *Tumor Biology* **35** 8319-8328.
867
868 Gautier A, Bonnet F, Dubois S, Massart C, Grosheny C, Bachelot A, Aubé C, Balkau B & Ducluzeau P-H
869 2013 Associations between visceral adipose tissue, inflammation and sex steroid concentrations in
870 men. *Clinical Endocrinology* **78** 373-378.
871
872 Geier R, Adler S, Rashid G & Klein A 2010 The synthetic estrogen diethylstilbestrol (DES) inhibits the
873 telomerase activity and gene expression of prostate cancer cells. *The Prostate* **70** 1307-1312.
874
875 Giton F, Sirab N, Franck G, Gervais M, Schmidlin F, Ali T, Allory Y, Taille Adl, Vacherot F, Loric S, et al.
876 2015 Evidence of estrone-sulfate uptake modification in young and middle-aged rat prostate. *The*
877 *Journal of Steroid Biochemistry and Molecular Biology* **152** 89-100.
878
879 Goetzl MA, VanVeldhuizen PJ & Thrasher JB 2007 Effects of soy phytoestrogens on the prostate.
880 *Prostate Cancer Prostatic Dis* **10** 216-223.
881
882 Gooren L & Morgentaler A 2014 Prostate cancer incidence in orchidectomised male-to-female
883 transsexual persons treated with oestrogens. *Andrologia* **46** 1156-1160.
884
885 Grönberg H 2003 Prostate cancer epidemiology. *The Lancet* **361** 859-864.

- 886
887 Hanahan D & Weinberg Robert A 2011 Hallmarks of Cancer: The Next Generation. *Cell* **144** 646-674.
888
- 889 Hariri W, Sudha T, Bharali DJ, Cui H & Mousa SA 2015 Nano-Targeted Delivery of Toremifene, an
890 Estrogen Receptor- α Blocker in Prostate Cancer. *Pharmaceutical Research* **32** 2764-2774.
891
- 892 Härkönen PL & Mäkelä SI 2004 Role of estrogens in development of prostate cancer. *The Journal of*
893 *Steroid Biochemistry and Molecular Biology* **92** 297-305.
894
- 895 Henderson BE, Bernstein L, Ross RK, Depue RH & Judd HL 1988 The early in utero oestrogen and
896 testosterone environment of blacks and whites: potential effects on male offspring. *British Journal of*
897 *Cancer* **57** 216-218.
898
- 899 Ho S-M 2004 Estrogens and anti-estrogens: Key mediators of prostate carcinogenesis and new
900 therapeutic candidates. *Journal of Cellular Biochemistry* **91** 491-503.
901
- 902 Hofland J, van Weerden WM, Dits NFJ, Steenbergen J, van Leenders GJLH, Jenster G, Schröder FH &
903 de Jong FH 2010 Evidence of Limited Contributions for Intratumoral Steroidogenesis in Prostate
904 Cancer. *Cancer Research* **70** 1256-1264.
905
- 906 Hofland J, Weerden WMv, Steenbergen J, Dits NFJ, Jenster G & Jong FHd 2011 Activin A regulates
907 local testosterone production and growth in prostate cancer through 17 β -hydroxysteroid
908 dehydrogenase. In *13th European Congress of Endocrinology*, p 353. Rotterdam, Netherlands:
909 Endocrine Abstracts, Bioscientifica.
910
- 911 Holt SK, Kwon EM, Fu R, Kolb S, Feng Z, Ostrander EA & Stanford JL 2013 Association of Variants in
912 Estrogen-Related Pathway Genes with Prostate Cancer Risk. *The Prostate* **73** 1-10.
913
- 914 Howell MA 1974 Factor Analysis of International Cancer Mortality Data and per capita Food
915 Consumption. *British Journal of Cancer* **29** 328-336.
916
- 917 Huang W, Chen Y, Liu Y, Zhang Q, Yu Z, Mou L, Wu H, Zhao L, Long T, Qin D, et al. 2015 Roles of ER β
918 and GPR30 in Proliferative Response of Human Bladder Cancer Cell to Estrogen. *BioMed research*
919 *international* **2015** 251780.
920
- 921 Huggins C & Hodges CV 1941 Studies on Prostatic Cancer. I. The Effect of Castration, of Estrogen and
922 of Androgen Injection on Serum Phosphatases in Metastatic Carcinoma of the Prostate. *Cancer*
923 *Research* **1** 293-297.
924
- 925 Hussain S, Lawrence MG, Taylor RA, Lo CY-W, BioResource APC, Frydenberg M, Ellem SJ, Furic L &
926 Risbridger GP 2012 Estrogen Receptor ? Activation Impairs Prostatic Regeneration by Inducing
927 Apoptosis in Murine and Human Stem/Progenitor Enriched Cell Populations. *PLoS ONE* **7** e40732.
928
- 929 Jacob M & Henrik M 2006 One, Five and Ten Year Cancer Prevalence by Cancer Network, UK, 2006.
930 In *One, Five and Ten Year Cancer Prevalence by Cancer Network, UK*. London, UK: National Cancer
931 Intelligence Network.
932
- 933 Jasuja GK, Travison TG, Davda M, Murabito JM, Basaria S, Zhang A, Kushnir MM, Rockwood AL,
934 Meikle W, Pencina MJ, et al. 2013 Age Trends in Estradiol and Estrone Levels Measured Using Liquid
935 Chromatography Tandem Mass Spectrometry in Community-Dwelling Men of the Framingham Heart
936 Study. *The Journals of Gerontology Series A: Biological Sciences and Medical Sciences* **68** 733-740.

937
 938 Jurečková J, Babušíková E, Kmeťová M, Kliment J & Dobrota D 2015 Estrogen receptor alpha
 939 polymorphisms and the risk of prostate cancer development. *Journal of Cancer Research and Clinical*
 940 *Oncology* **141** 1963-1971.
 941
 942 Khurana I 2008 *Essentials of Medical Physiology*. India: Elsevier India Pvt. Limited.
 943 Kimbro K & Simons J 2006 Hypoxia-inducible factor-1 in human breast and prostate cancer.
 944 *Endocrine-Related Cancer* **13** 739-749.
 945
 946 King KJ, Nicholson HD & Assinder SJ 2006 Effect of increasing ratio of estrogen: Androgen on
 947 proliferation of normal human prostate stromal and epithelial cells, and the malignant cell line
 948 LNCaP. *The Prostate* **66** 105-114.
 949
 950 Klein H, Molwitz T & Bartsch W 1989 Steroid sulfate sulfatase in human benign prostatic hyperplasia:
 951 Characterization and quantification of the enzyme in epithelium and stroma. *Journal of Steroid*
 952 *Biochemistry* **33** 195-200.
 953
 954 Knochenhauer ES, Boots LR, Potter HD & Azziz R 1998 Differential binding of estradiol and
 955 testosterone to SHBG. Relation to circulating estradiol levels. *The Journal of reproductive medicine*
 956 **43** 665-670.
 957
 958 Koh E, Noda T, Kanaya J & Namiki M 2002 Differential expression of 17 β -hydroxysteroid
 959 dehydrogenase isozyme genes in prostate cancer and noncancer tissues. *The Prostate* **53** 154-159.
 960
 961 Kolonel LN, Altshuler D & Henderson BE 2004 The multiethnic cohort study: exploring genes, lifestyle
 962 and cancer risk. *Nat Rev Cancer* **4** 519-527.
 963
 964 Kuiper GGJM, Lemmen JG, Carlsson B, Corton JC, Safe SH, Saag PTvd, Burg Bvd & Gustafsson J-Å
 965 1998 Interaction of Estrogenic Chemicals and Phytoestrogens with Estrogen Receptor β .
 966 *Endocrinology* **139** 4252-4263.
 967
 968 Kumagai J, Hofland J, Erkens-Schulze S, Dits NFJ, Steenbergen J, Jenster G, Homma Y, de Jong FH &
 969 van Weerden WM 2013 Intratumoral conversion of adrenal androgen precursors drives androgen
 970 receptor-activated cell growth in prostate cancer more potently than de novo steroidogenesis. *The*
 971 *Prostate* **73** 1636-1650.
 972
 973 Labrie F, Bélanger A, Luu-The V, Labrie C, Simard J, Cusan L, Gomez J & Candas B 2005 Gonadotropin-
 974 Releasing Hormone Agonists in the Treatment of Prostate Cancer. *Endocrine Reviews* **26** 361-379.
 975
 976 Lam H-M, Ouyang B, Chen J, Ying J, Wang J, Wu C-L, Jia L, Medvedovic M, Vessella RL & Ho S-M 2014
 977 Targeting GPR30 with G-1: a new therapeutic target for castration-resistant prostate cancer.
 978 *Endocrine-Related Cancer* **21** 903-914.
 979
 980 Lee J, Demissie K, Lu S & Rhoads GG 2007 Cancer incidence among Korean-American immigrants in
 981 the United States and native Koreans in South Korea. *Cancer Control* **14** 78.
 982
 983 Leung Y-K, Lam H-M, Wu S, Song D, Levin L, Cheng L, Wu C-L & Ho S-M 2010 Estrogen receptor β 2
 984 and β 5 are associated with poor prognosis in prostate cancer, and promote cancer cell migration and
 985 invasion. *Endocrine-Related Cancer* **17** 675-689.
 986

- 987 Leung Y-K, Mak P, Hassan S & Ho S-M 2006 Estrogen receptor (ER)- β isoforms: A key to
 988 understanding ER- β signaling. *Proceedings of the National Academy of Sciences of the United States*
 989 *of America* **103** 13162-13167.
 990
- 991 Li R, Evaul K, Sharma KK, Chang K-H, Yoshimoto J, Liu J, Auchus RJ & Sharifi N 2012 Abiraterone
 992 Inhibits 3 β -Hydroxysteroid Dehydrogenase: A Rationale for Increasing Drug Exposure in Castration-
 993 Resistant Prostate Cancer. *Clinical Cancer Research* **18** 3571-3579.
 994
- 995 Li Y, Wu Y, Abbatiello TC, Wu WL, Kim JR, Sarkissyan M, Sarkissyan S, Chung SS, Elshimali Y &
 996 Vadgama JV 2015a Slug contributes to cancer progression by direct regulation of ER α signaling
 997 pathway. *International Journal of Oncology* **46** 1461-1472.
 998
- 999 Li Z, Bishop AC, Alyamani M, Garcia JA, Dreicer R, Bunch D, Liu J, Upadhyay SK, Auchus RJ & Sharifi N
 1000 2015b Conversion of abiraterone to D4A drives anti-tumour activity in prostate cancer. *Nature* **523**
 1001 347-351.
 1002
- 1003 Liu C, Lou W, Zhu Y, Yang JC, Nadiminty N, Gaikwad NW, Evans CP & Gao AC 2015 Intracrine
 1004 androgens and AKR1C3 activation confer resistance to enzalutamide in prostate cancer. *Cancer*
 1005 *Research* **75** 1413-1422.
 1006
- 1007 Locke JA, Guns ES, Lubik AA, Adomat HH, Hendy SC, Wood CA, Ettinger SL, Gleave ME & Nelson CC
 1008 2008 Androgen Levels Increase by Intratumoral De novo Steroidogenesis during Progression of
 1009 Castration-Resistant Prostate Cancer. *Cancer Research* **68** 6407-6415.
 1010
- 1011 Lukacik P, Kavanagh KL & Oppermann U 2006 Structure and function of human 17 β -hydroxysteroid
 1012 dehydrogenases. *Molecular and Cellular Endocrinology* **248** 61-71.
 1013
- 1014 Luu-The V, Bélanger A & Labrie F 2008 Androgen biosynthetic pathways in the human prostate. *Best*
 1015 *Practice & Research Clinical Endocrinology & Metabolism* **22** 207-221.
 1016
- 1017 Maier KG, Brandy K, Kittur D, Kort K & LaSpina M 2006 GPR30: A novel estrogen receptor expressed
 1018 in aggressive prostate cancer. *The FASEB Journal* **20** A221.
 1019
- 1020 Mak P, Leav I, Pursell B, Bae D, Yang X, Taglienti CA, Gouvin LM, Sharma VM & Mercurio AM 2010
 1021 ER β Impedes Prostate Cancer EMT by Destabilizing HIF-1 α and Inhibiting VEGF-Mediated Snail
 1022 Nuclear Localization: Implications for Gleason Grading. *Cancer cell* **17** 319-332.
 1023
- 1024 Mak P, Li J, Samanta S & Mercurio AM 2015 ER β regulation of NF- κ B activation in prostate cancer is
 1025 mediated by HIF-1. *Oncotarget* **6** 40247-40254.
 1026
- 1027 Malkowicz SB 2001 The role of diethylstilbestrol in the treatment of prostate cancer. *Urology* **58**
 1028 108-113.
 1029
- 1030 Marcelli M, Ittmann M, Mariani S, Sutherland R, Nigam R, Murthy L, Zhao Y, DiConcini D, Puxeddu E,
 1031 Esen A, et al. 2000 Androgen Receptor Mutations in Prostate Cancer. *Cancer Research* **60** 944-949.
 1032
- 1033 McPherson SJ, Ellem SJ, Simpson ER, Patchev V, Fritzemeier K-H & Risbridger GP 2007 Essential Role
 1034 for Estrogen Receptor β in Stromal-Epithelial Regulation of Prostatic Hyperplasia. *Endocrinology* **148**
 1035 566-574.
 1036

- 1037 McTernan P, Anwar A, Eggo M, Barnett A, Stewart P & Kumar S 2000 Gender differences in the
1038 regulation of P450 aromatase expression and activity in human adipose tissue. *International journal*
1039 *of obesity* **24** 875-881.
- 1040
- 1041 Miftakhova R, Hedblom A, Semenas J, Robinson B, Simoulis A, Malm J, Rizvanov A, Heery DM,
1042 Mongan NP, Maitland NJ, et al. 2016 Cyclin A1 and P450 Aromatase Promote Metastatic Homing and
1043 Growth of Stem-like Prostate Cancer Cells in the Bone Marrow. *Cancer Research* **76** 2453-2464.
- 1044
- 1045 Mo Z, Liu M, Yang F, Luo H, Li Z, Tu G & Yang G 2013 GPR30 as an initiator of tamoxifen resistance in
1046 hormone-dependent breast cancer. *Breast Cancer Research* **15** 1-15.
- 1047
- 1048 Mohler JL, Gregory CW, Ford OH, Kim D, Weaver CM, Petrusz P, Wilson EM & French FS 2004 The
1049 Androgen Axis in Recurrent Prostate Cancer. *Clinical Cancer Research* **10** 440-448.
- 1050
- 1051 Montgomery RB, Mostaghel EA, Vessella R, Hess DL, Kalhorn TF, Higano CS, True LD & Nelson PS
1052 2008 Maintenance of Intratumoral Androgens in Metastatic Prostate Cancer: A Mechanism for
1053 Castration-Resistant Tumor Growth. *Cancer Research* **68** 4447-4454.
- 1054
- 1055 Montico F, Hetzl AC, Cândido EM & Cagnon VHA 2013 Angiogenic and Tissue Remodeling Factors in
1056 the Prostate of Elderly Rats Submitted to Hormonal Replacement. *The Anatomical Record* **296** 1758-
1057 1767.
- 1058
- 1059 Mueller JW, Gilligan LC, Idkowiak J, Arlt W & Foster PA 2015 The Regulation of Steroid Action by
1060 Sulfation and Desulfation. *Endocrine Reviews* **36** 526-563.
- 1061
- 1062 Muir M, Romalo G, Wolf L, Elger W & Schweikert H-U 2004 Estrone Sulfate Is a Major Source of Local
1063 Estrogen Formation in Human Bone. *The Journal of Clinical Endocrinology & Metabolism* **89** 4685-
1064 4692.
- 1065
- 1066 Murray R & Pitt P 1985 Treatment of advanced prostatic cancer, resistant to conventional therapy
1067 with aminoglutethimide. *European Journal of Cancer and Clinical Oncology* **21** 453-458.
- 1068
- 1069 Nakamura Y, Suzuki T, Fukuda T, Ito A, Endo M, Moriya T, Arai Y & Sasano H 2006 Steroid sulfatase
1070 and estrogen sulfotransferase in human prostate cancer. *The Prostate* **66** 1005-1012.
- 1071
- 1072 Nelson AW, Tilley WD, Neal DE & Carroll JS 2014 Estrogen receptor beta in prostate cancer: friend or
1073 foe? *Endocrine-Related Cancer* **21** T219-T234.
- 1074
- 1075 O'Dowd BF, Nguyen T, Marchese A, Cheng R, Lynch KR, Heng HHQ, Kolakowski Jr LF & George SR
1076 1998 Discovery of Three Novel G-Protein-Coupled Receptor Genes. *Genomics* **47** 310-313.
- 1077
- 1078 Oduwole OO, Mäkinen MJ, Isomaa VV, Pulkka A, Jernvall P, Karttunen TJ & Vihko PT 2003 17 β -
1079 Hydroxysteroid dehydrogenase type 2: independent prognostic significance and evidence of
1080 estrogen protection in female patients with colon cancer. *The Journal of Steroid Biochemistry and*
1081 *Molecular Biology* **87** 133-140.
- 1082
- 1083 Parkin DM, Bray FI & Devesa SS 2001 Cancer burden in the year 2000. The global picture. *European*
1084 *Journal of Cancer* **37**, **Supplement 8** 4-66.
- 1085
- 1086 Parkin DM, Whelan SL, J F, L T & DB T 1997 Cancer Incidence in Five Continents. . Lyon: International
1087 Agency for Research on Cancer Scientific Publications.

- 1088
 1089 Piao Y-s, Wiesenfeld P, Sprando R & Arnold JT 2013 TGFβ1 alters androgenic metabolites and
 1090 hydroxysteroid dehydrogenase enzyme expression in human prostate reactive stromal primary cells:
 1091 Is steroid metabolism altered by prostate reactive stromal microenvironment? *The Journal of Steroid*
 1092 *Biochemistry and Molecular Biology* **138** 10.1016/j.jsbmb.2013.1005.1016.
 1093
 1094 Pilšáková L, Riečanský I & Jagla F 2010 The physiological actions of isoflavone phytoestrogens.
 1095 *Physiological research / Academia Scientiarum Bohemoslovaca* **59** 651-664.
 1096
 1097 Polari L, Yatkin E, Martínez Chacón MG, Ahotupa M, Smeds A, Strauss L, Zhang F, Poutanen M,
 1098 Saarinen N & Mäkelä SI 2015 Weight gain and inflammation regulate aromatase expression in male
 1099 adipose tissue, as evidenced by reporter gene activity. *Molecular and Cellular Endocrinology* **412**
 1100 123-130.
 1101
 1102 Prins GS & Hu W-Y 2013 *Stem Cells and Prostate Cancer*. New York: Springer New York.
 1103
 1104 Prins GS, Huang L, Birch L & Pu Y 2006 The Role of Estrogens in Normal and Abnormal Development
 1105 of the Prostate Gland. *Annals of the New York Academy of Sciences* **1089** 1-13.
 1106
 1107 Prossnitz ER, Arterburn JB & Sklar LA 2007 GPR30: a G protein-coupled receptor for estrogen.
 1108 *Molecular and Cellular Endocrinology* **265-266** 138-142.
 1109
 1110 Prossnitz ER, Oprea TI, Sklar LA & Arterburn JB 2008a The ins and outs of GPR30: a transmembrane
 1111 estrogen receptor. *The Journal of Steroid Biochemistry and Molecular Biology* **109** 350-353.
 1112
 1113 Prossnitz ER, Sklar LA, Oprea TI & Arterburn JB 2008b GPR30: a novel therapeutic target in estrogen-
 1114 related disease. *Trends in Pharmacological Sciences* **29** 116-123.
 1115
 1116 Purohit A & Foster PA 2012 Steroid sulfatase inhibitors for estrogen- and androgen-dependent
 1117 cancers. *Journal of Endocrinology* **212** 99-110.
 1118
 1119 Quaresma M, Coleman MP & Rachet B 2015 40-year trends in an index of survival for all cancers
 1120 combined and survival adjusted for age and sex for each cancer in England and Wales,
 1121 1971–2011: a population-based study. *The Lancet* **385** 1206-1218.
 1122
 1123 Raftogianis R, Creveling C, Weinshilboum R & Weisz J 2000 Chapter 6: Estrogen Metabolism by
 1124 Conjugation. *JNCI Monographs* **2000** 113-124.
 1125
 1126 Rago V, Romeo F, Giordano F, Ferraro A & Carpino A 2016 Identification of the G protein-coupled
 1127 estrogen receptor (GPER) in human prostate: expression site of the estrogen receptor in the benign
 1128 and neoplastic gland. *Andrology* **4** 121-127.
 1129
 1130 Rajfer J & Coffey DS 1978 Sex steroid imprinting of the immature prostate. Long-term effects.
 1131 *Investigative urology* **16** 186-190.
 1132
 1133 Reed MJ, Purohit A, Woo LWL, Newman SP & Potter BVL 2005 Steroid Sulfatase: Molecular Biology,
 1134 Regulation, and Inhibition. *Endocrine Reviews* **26** 171-202.
 1135
 1136 Reger M, Zollinger T, Liu Z, Jones J & Zhang J 2015 Abstract 1884: Dietary intake of phytoestrogens
 1137 and the risk of prostate cancer in the Prostate, Lung, Colorectal, and Ovarian Cancer Screening Trial.
 1138 *Cancer Research* **75** 1884.

1139
 1140 Risbridger GP, Davis ID, Birrell SN & Tilley WD 2010 Breast and prostate cancer: more similar than
 1141 different. *Nat Rev Cancer* **10** 205-212.
 1142
 1143 Robinson-Rechavi M, Garcia HE & Laudet V 2003 The nuclear receptor superfamily. *Journal of Cell*
 1144 *Science* **116** 585-586.
 1145
 1146 Romanel A, Tandefelt DG, Conteduca V, Jayaram A, Casiraghi N, Wetterskog D, Salvi S, Amadori D,
 1147 Zafeiriou Z, Rescigno P, et al. 2015 Plasma AR and abiraterone-resistant prostate cancer. *Science*
 1148 *Translational Medicine* **7** 312re310-312re310.
 1149
 1150 Ronde Wd, Schouw YTvd, Muller M, Grobbee DE, Gooren LJG, Pols HAP & Jong FHd 2005
 1151 Associations of Sex-Hormone-Binding Globulin (SHBG) with Non-SHBG-Bound Levels of Testosterone
 1152 and Estradiol in Independently Living Men. *The Journal of Clinical Endocrinology & Metabolism* **90**
 1153 157-162.
 1154
 1155 Ross-Innes CS, Stark R, Teschendorff AE, Holmes KA, Ali HR, Dunning MJ, Brown GD, Gojis O, Ellis IO,
 1156 Green AR, et al. 2012 Differential oestrogen receptor binding is associated with clinical outcome in
 1157 breast cancer. *Nature* **481** 389-393.
 1158
 1159 Roy J, Lefebvre J, Maltais R & Poirier D 2013 Inhibition of dehydroepiandrosterone sulfate action in
 1160 androgen-sensitive tissues by EM-1913, an inhibitor of steroid sulfatase. *Molecular and Cellular*
 1161 *Endocrinology* **376** 148-155.
 1162
 1163 Ryan CJ, Smith MR, de Bono JS, Molina A, Logothetis CJ, de Souza P, Fizazi K, Mainwaring P,
 1164 Piulats JM, Ng S, et al. 2013 Abiraterone in Metastatic Prostate Cancer without Previous
 1165 Chemotherapy. *New England Journal of Medicine* **368** 138-148.
 1166
 1167 Sakr WA, Haas GP, Cassin BF, Pontes JE & Crissman JD 1993 The frequency of carcinoma and
 1168 intraepithelial neoplasia of the prostate in young male patients. *J Urol* **150** 379-385.
 1169
 1170 Samaras N, Frangos E, Forster A, Lang PO & Samaras D 2012 Andropause: A review of the definition
 1171 and treatment. *European Geriatric Medicine* **3** 368-373.
 1172
 1173 Santen RJ, Petroni GR, Fisch MJ, Myers CE, Theodorescu D & Cohen RB 2001 Use of the aromatase
 1174 inhibitor anastrozole in the treatment of patients with advanced prostate carcinoma. *Cancer* **92**
 1175 2095-2101.
 1176
 1177 Santen RJ, Thibault A, Myers C & Chung L 1997 Meeting summary: Workshop conference on
 1178 endocrine therapy of advanced prostate cancer, airle house, November 3–4, 1996. *Urologic*
 1179 *Oncology: Seminars and Original Investigations* **3** 31-38.
 1180
 1181 Sarveswaran S, Gautam SC & Ghosh J 2012 Wedelolactone, a medicinal plant-derived coumestan,
 1182 induces caspase-dependent apoptosis in prostate cancer cells via downregulation of PKC ϵ without
 1183 inhibiting Akt. *International Journal of Oncology* **41** 2191-2199.
 1184
 1185 Savoy RM & Ghosh PM 2013 The changing roles of steroid nuclear receptors with prostate cancer
 1186 progression. *Endocrine-Related Cancer* **20** C9-C11.
 1187

- 1188 Scher HI, Fizazi K, Saad F, Taplin M-E, Sternberg CN, Miller K, de Wit R, Mulders P, Chi KN, Shore
1189 ND, et al. 2012 Increased Survival with Enzalutamide in Prostate Cancer after Chemotherapy. *New*
1190 *England Journal of Medicine* **367** 1187-1197.
- 1191
- 1192 Sharifi N & Auchus RJ 2012 Steroid biosynthesis and prostate cancer. *Steroids* **77** 719-726.
- 1193
- 1194 Siegel R, Naishadham D & Jemal A 2012 Cancer statistics, 2012. *CA: A Cancer Journal for Clinicians* **62**
1195 10-29.
- 1196
- 1197 Slavin S, Yeh C-R, Da J, Yu S, Miyamoto H, Messing EM, Guancial E & Yeh S 2014 Estrogen receptor α
1198 in cancer-associated fibroblasts suppresses prostate cancer invasion via modulation of
1199 thrombospondin 2 and matrix metalloproteinase 3. *Carcinogenesis* **35** 1301-1309.
- 1200
- 1201 Smith MR, Kaufman D, George D, Oh WK, Kazanis M, Manola J & Kantoff PW 2002 Selective
1202 aromatase inhibition for patients with androgen-independent prostate carcinoma. *Cancer* **95** 1864-
1203 1868.
- 1204
- 1205 Stanway SJ, Purohit A, Woo LWL, Sufi S, Vigushin D, Ward R, Wilson RH, Stanczyk FZ, Dobbs N,
1206 Kulinskaya E, et al. 2006 Phase I Study of STX 64 (667 Coumate) in Breast Cancer Patients: The First
1207 Study of a Steroid Sulfatase Inhibitor. *Clinical Cancer Research* **12** 1585-1592.
- 1208
- 1209 Strom SS, Yamamura Y, Duphorne CM, Spitz MR, Babaian RJ, Pillow PC & Hursting SD 1999
1210 Phytoestrogen intake and prostate cancer: A case-control study using a new database. *Nutrition and*
1211 *Cancer* **33** 20-25.
- 1212
- 1213 Szliszka E, Czuba ZP, Mertas A, Paradysz A & Krol W 2013 The dietary isoflavone biochanin-A
1214 sensitizes prostate cancer cells to TRAIL-induced apoptosis. *Urologic Oncology: Seminars and*
1215 *Original Investigations* **31** 331-342.
- 1216
- 1217 Takeuchi S, Takahashi T, Sawada Y, Iida M, Matsuda T & Kojima H 2009 Comparative Study on the
1218 Nuclear Hormone Receptor Activity of Various Phytochemicals and Their Metabolites by Reporter
1219 Gene Assays Using Chinese Hamster Ovary Cells. *Biological and Pharmaceutical Bulletin* **32** 195-202.
- 1220
- 1221 Takizawa I, Lawrence MG, Balanathan P, Rebello R, Pearson HB, Garg E, Pedersen J, Pouliot N, Nadon
1222 R, Watt MJ, et al. 2015 Estrogen receptor alpha drives proliferation in PTEN-deficient prostate
1223 carcinoma by stimulating survival signaling, MYC expression and altering glucose sensitivity.
1224 *Oncotarget* **6** 604-616.
- 1225
- 1226 Taneja SS, Morton R, Barnette G, Sieber P, Hancock ML & Steiner M 2013 Prostate Cancer Diagnosis
1227 Among Men With Isolated High-Grade Intraepithelial Neoplasia Enrolled Onto a 3-Year Prospective
1228 Phase III Clinical Trial of Oral Toremifene. *Journal of Clinical Oncology* **31** 523-529.
- 1229
- 1230 Taylor RA, Toivanen R, Frydenberg M, Pedersen J, Harewood L, Australian Prostate Cancer B, Collins
1231 AT, Maitland NJ & Risbridger GP 2012 Human Epithelial Basal Cells Are Cells of Origin of Prostate
1232 Cancer, Independent of CD133 Status. *STEM CELLS* **30** 1087-1096.
- 1233
- 1234 Thomas P & Dong J 2006 Binding and activation of the seven-transmembrane estrogen receptor
1235 GPR30 by environmental estrogens: A potential novel mechanism of endocrine disruption. *The*
1236 *Journal of Steroid Biochemistry and Molecular Biology* **102** 175-179.
- 1237
- 1238 Usui T 2006 Pharmaceutical Prospects of Phytoestrogens. *Endocrine Journal* **53** 7-20.

- 1239
 1240 Utsumi T, Yoshimura N, Takeuchi S, Maruta M, Maeda K & Harada N 2000 Elevated steroid sulfatase
 1241 expression in breast cancers. *The Journal of Steroid Biochemistry and Molecular Biology* **73** 141-145.
 1242
- 1243 Vidal AC, Howard LE, Moreira DM, Castro-Santamaria R, Andriole GL & Freedland SJ 2014 Obesity
 1244 Increases the Risk for High-grade Prostate Cancer: Results from the REDUCE study. *Cancer*
 1245 *epidemiology, biomarkers & prevention : a publication of the American Association for Cancer*
 1246 *Research, cosponsored by the American Society of Preventive Oncology* **23** 2936-2942.
 1247
- 1248 Wang F, Vihma V, Soronen J, Turpeinen U, Hämäläinen E, Savolainen-Peltonen H, Mikkola TS,
 1249 Naukkarinen J, Pietiläinen KH, Jauhiainen M, et al. 2013 17 β -Estradiol and Estradiol Fatty Acyl Esters
 1250 and Estrogen-Converting Enzyme Expression in Adipose Tissue in Obese Men and Women. *The*
 1251 *Journal of Clinical Endocrinology & Metabolism* **98** 4923-4931.
 1252
- 1253 Wernert N, Kern L, Heitz P, Bonkhoff H, Goebbels R, Seitz G, Inniger R, Remberger K & Dhom G 1990
 1254 Morphological and immunohistochemical investigations of the utriculus prostaticus from the fetal
 1255 period up to adulthood. *The Prostate* **17** 19-30.
 1256
- 1257 White BA, Porterfield SP & Porterfield SP 2013 *Endocrine and reproductive physiology*. Philadelphia,
 1258 PA: Elsevier/Mosby.
 1259
- 1260 Whittemore AS, Kolonel LN, Wu AH, John EM, Gallagher RP, Howe GR, Burch JD, Hankin J, Dreon DM,
 1261 West DW, et al. 1995 Prostate Cancer in Relation to Diet, Physical Activity, and Body Size in Blacks,
 1262 Whites, and Asians in the United States and Canada. *Journal of the National Cancer Institute* **87** 652-
 1263 661.
 1264
- 1265 Wright JL, Kwon EM, Ostrander EA, Montgomery RB, Lin DW, Vessella R, Stanford JL & Mostaghel EA
 1266 2011 Expression of SLCO Transport Genes in Castration-Resistant Prostate Cancer and Impact of
 1267 Genetic Variation in SLCO1B3 and SLCO2B1 on Prostate Cancer Outcomes. *Cancer Epidemiology*
 1268 *Biomarkers & Prevention* **20** 619-627.
 1269
- 1270 Yamnik RL & Holz MK 2010 mTOR/S6K1 and MAPK/RSK signaling pathways coordinately regulate
 1271 estrogen receptor α serine 167 phosphorylation. *FEBS Letters* **584** 124-128.
 1272
- 1273 Yang M, Kenfield SA, Blarigan ELV, Kasperzyk JL, Sesso HD, Ma J, Stampfer M & Chavarro JE 2014
 1274 Abstract 1263: Dietary patterns after prostate cancer diagnosis in relation to disease-specific and
 1275 total mortality. *Cancer Research* **74** 1263.
 1276
- 1277 Yeh C-R, Da J, Song W, Fazili A & Yeh S 2014 Estrogen receptors in prostate development and cancer.
 1278 *American Journal of Clinical and Experimental Urology* **2** 161-168.
- 1279 Yeh S, Miyamoto H, Shima H & Chang C 1998 From estrogen to androgen receptor: a new pathway
 1280 for sex hormones in prostate cancer. *Proc Natl Acad Sci* **95** 5527-32.
- 1281
 1282
 1283

Figure Legends

Figure 1: Molecular similarities between phytoestrogens and E₂. E₂ contains the cyclopenta[α]phenanthrene ring structure common to all steroid molecules. Isoflavones and coumestans are two common categories of phytoestrogens and have a molecular structure similar to E₂. As a result phytoestrogens can also bind and activate the oestrogen receptors.

Figure 2: Oestrogen and Androgen synthesis pathways.

Intratumoural E₂ can be formed from desulfation and reduction of circulating oestrone-sulphate (E₁S) by steroid sulphotase (STS) and 17 β -hydroxysteroid dehydrogenase (HSD). Alternatively, oestrogens can be produced from androstenedione or testosterone by aromatase. Aromatase competes with 5 α -reductase (SRD5A1), responsible for potentiating androgens, for these substrates. DHEA, the precursor for androstenedione, is most likely derived from the large pool of circulating DHEAS by STS, as intratumoural synthesis from progestogens remains disputable.

Figure 3: The expression of ER α and ER β changes during prostate cancer progression.

During development of prostate cancer the ER β isoform is downregulated in epithelial cells. On the other hand, ER α is upregulated in tumour cells as well as the surrounding environment. The remainder of the 'normal' prostate retains its existing expression of ER α and ER β .

Figure 4: Signalling pathways in prostate cancer through ER α , ER β and GPER. ER α and ER β bind to the oestrogen response elements (ERE) of DNA and regulate transcription. Activation of ER α induces mitogenic pathways via PI3K which in turn promotes HIF-1 α which activates anti-apoptotic pathways; whereas activation of ER β induces apoptosis, cell cycle arrest and inhibits dedifferentiation pathways. GPER activation in prostate cancer is anti-tumourigenic as it upregulates p21 and induces cell cycle arrest.

Figure 5: The altered expression of ERs during prostate cancer development. Changes in ER α and ER β have been studied throughout the evolution of prostate cancer; however, expression of GPER in normal prostate and early stages of prostate cancer is currently unknown.

Fig. 1

Figure 1
18x13mm (600 x 600 DPI)

Fig. 2

Figure 2
18x13mm (600 x 600 DPI)

Figure 3
18x13mm (600 x 600 DPI)

Fig. 4

Figure 4
18x13mm (600 x 600 DPI)

Fig. 5

Figure 5
18x13mm (600 x 600 DPI)