The effect of dutasteride on the detection of prostate cancer: A set of meta-analyses

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Abstract

Background: Dutasteride has been shown to significantly improve symptoms of benign prostatic hyperplasia (BPH) and reduce clinical progression. Recent data from studies evaluating 5-alpha reductase inhibitors (5-ARIs) for the prevention of prostate cancer, however, suggest 5ARIs, including dutasteride, may be associated with increased incidence of Gleason 8-10 prostate tumours. This meta-analysis was undertaken to quantify the effect of dutasteride on detection of prostate cancer and high-grade prostate cancer.

Methods: Our meta-analysis includes data from GlaxoSmithKline-sponsored phase III randomized clinical trials (with a study duration of ≥2 years) evaluating the effect of dutasteride, alone or in combination with tamsulosin, to treat BPH or to reduce the risk of prostate cancer. The incidence of prostate cancer, including Gleason 7-10 and Gleason 8-10, for patients taking either dutasteride, dutasteride plus tamsulosin, tamsulosin alone, or placebo, were evaluated using the Mantel-Haenszel Risk Ratio (MHRR) method of conducting meta-analyses.

Results: The meta-analysis demonstrated that in a population with symptomatic BPH and/or at increased risk of prostate cancer, a statistically significant lower number of detectable prostate cancers was found in men taking dutasteride compared to control groups (MHRR: 0.66, 95% CI 0.52-0.85). In our analysis, there was no increased risk for Gleason 7-10 (MHRR: 0.83, 95% CI 0.56-1.21) or Gleason 8-10 prostate cancers (MHRR: 0.99, 95% CI 0.39-2.53) in men taking dutasteride over control groups. There were several limitations that need to be considered when interpreting these results.

Conclusion: These data provide support for the continued use of dutasteride in the treatment of symptomatic BPH patients.

Introduction

Prostate conditions, such as benign prostatic hyperplasia (BPH), remain a significant concern in men, especially for those over age 50. It is estimated that 40% of men over 50 and 90% over 80 have BPH.1,2 In Canada, the number of men 50 years and over is projected to grow by over 37% to 6.5 million by 2018, and the number of men with moderate-to-severe lower urinary tract symptoms (LUTS) is expected to increase by 41% to 2.6 million.3 BPH has a significant economic burden on society with direct annual costs of $1.1 billion in the year 2000 in the United States alone, exclusive of outpatient medication.4

Medical treatments for BPH include 5-alpha reductase inhibitors (5-ARIs), which block the conversion of testosterone to dihydrotestosterone (DHT) via 5-alpha reductase (5-AR).5 Dutasteride inhibits both isoforms of 5-AR and reduces serum DHT by more than 90%, subsequently reducing prostate volume and improving BPH symptoms.5,6 Levels of both isoforms of 5-AR are slightly increased in BPH compared to the normal prostate, while mainly type 1 5-AR levels are increased in prostate cancer tissue.7 Theoretically, suppression of prostatic DHT may also affect the development and progression of prostate neoplasia.

Prostate cancer incidence was reported in several Phase III studies that evaluated the efficacy of dutasteride to treat symptomatic BPH. In a pooled analysis of three Phase III BPH monotherapy studies (ARIA3001, ARIA3002, ARIB3003), the incidence of detectable prostate cancer reported as an adverse event, determined by for-cause biopsies, was almost 50% less in the dutasteride group compared to the placebo group at 27 months (1.2 and 2.5%, p = 0.002).8 In a post-hoc analysis of the Combination of Avodart and Tamsulosin (CombAT) study in BPH patients, dutasteride, alone or in combination with tamsulosin, showed a relative reduction of 40% in the risk of detectable prostate cancers compared to tamsulosin monotherapy, as well as a 40% reduction in biopsies, over the 4-year study duration.9 There were numerically fewer Gleason score 7-10 and Gleason score 8-10 tumours in the dutasteride groups combined, compared with the tamsulosin group, and a similar reduction of 40% was observed for low- and high-grade Gleason score cancers as for all tumours.9

5-ARIs have also been examined in prostate cancer chemoprevention. The Prostate Cancer Prevention Trial (PCPT) study showed a reduction in the incidence of for-cause prostate cancer of 24.8% in normal men treated with finasteride after 7 years.10 The ReDuction by DUtasteride of
Dutasteride treatment in patients taking finasteride. Dutasteride treatment in men administered dutasteride or finasteride over placebo, there was an increased incidence of Gleason grade 8-10 prostate cancer. The occurrence of prostate cancer in the following risk population. While these studies showed a decrease in incidence of overall prostate cancers in men administered dutasteride or finasteride over placebo, there was an increased incidence of Gleason grade 8-10 prostate cancers compared to placebo, using the classical Gleason scoring system, and 0.5% using the modified Gleason scores. In PCPT, a statistically significant higher incidence of grade 8-10 cancers was seen in the finasteride group over the placebo overall (0.96% vs. 0.56%; relative risk 1.70 [95% CI 1.22-2.39]).

Taken together, these results suggest that patients taking 5-ARIs may be at an increased risk for developing high-grade prostate cancer, a finding that has led to labelling changes in Canada, Europe and the U.S. This has subsequently increased concern among patients and physicians, even though these labelling changes also indicate a favourable benefit:risk profile for 5-ARIs in the treatment of BPH.

In the present study, we undertook a meta-analysis of all GlaxoSmithKline (GSK)-sponsored Phase III studies in which dutasteride was evaluated for the treatment of BPH or for risk reduction of prostate cancer prior to 2012, in an effort to quantify the effect of dutasteride on the detection of prostate cancer and high-grade prostate cancer.

### Methods

#### Selection of studies

All GSK-sponsored Phase III randomized, double-blind, controlled trials (≥2 years’ duration) with dutasteride compared to an active comparator/placebo in the treatment of BPH or risk reduction of prostate cancer were eligible for inclusion. We excluded prior studies that did not report prostate cancer. The occurrence of prostate cancer in the following studies was used for the analysis:

- **BPH monotherapy studies** (ARIA3001, ARIA3002, ARIB3003): 2-year studies, comprising 4325 men, to evaluate efficacy and safety of dutasteride 0.5 mg once daily to treat symptomatic men. Biopsies were conducted “for-cause” only, based on the investigator’s judgment, and Gleason scores were not collected prospectively. The intent-to-treat population was used in the analyses.
- **CombAT**: A 4-year study in 4844 men, in which dutasteride in combination with tamsulosin was compared to dutasteride or tamsulosin monotherapy, for improving BPH symptoms, and to reduce the risk of acute urinary retention (AUR) and BPH-surgery. Biopsies were conducted for-cause only, based on the investigator’s judgment. The intent-to-treat population was used for the analyses.

#### Results

We tallied study characteristics for the randomized controlled trials used in the meta-analysis (Table 1). While the patient population differs between the BPH and prostate cancer risk reduction studies, the meta-analysis allows for a broader look at the effect of dutasteride on the detection of prostate cancers in patients who have a wide range of prostate-specific antigen (PSA) values, prostate volume and symptom scores, as defined by the International Prostate Symptom Score (IPSS).

#### Any prostate cancers

Any prostate cancers were defined as a total of all types of prostate cancer detected in patients, regardless of Gleason...
For-cause prostate cancers

For-causes cancers were defined as cancers detected during a biopsy required by the investigator as a result of an abnormal finding, such as elevated PSA or an abnormal digital rectal exam (DRE). Figure 2 describes the effect of dutasteride (0.5 mg daily) or dutasteride (0.5 mg daily) + tamsulosin (0.4 mg daily) versus placebo or tamsulosin (0.4 mg daily) on prostate cancer risk reduction on for-cause prostate cancers. Since there were no scheduled biopsies in the Phase III BPH monotherapy or the CombAT studies, the for-cause cancer results were identical to any-prostate-cancer results for these studies as described above. The REDUCE trial did not show a statistically significant decrease in risk for patients using dutasteride for detectable prostate cancers (MHRR: 0.74, 95% CI 0.53-1.02), although the risk reduction (26%) was similar to risk reduction for any-prostate-cancer results (23%). The test for heterogeneity illustrated low heterogeneity ($I^2 = 10.24$) and was not statistically significant ($Q = 2.23$, $p = 0.33$), and therefore was reasonable to pool results. The meta-analysis found a significant risk reduction (37%) for detectable for-cause cancer in patients using dutasteride compared to placebo or tamsulosin (MHRR: 0.63, 95% CI 0.52-0.78).

Gleason 7-10 prostate cancers

We describe the effect of dutasteride (0.5 mg daily) or dutasteride (0.5 mg daily) + tamsulosin (0.4 mg daily) vs. placebo or tamsulosin (0.4 mg daily) on the risk of detectable Gleason 7-10 prostate cancers (Fig. 3). As Gleason scoring was not prospectively collected in the BPH monotherapy studies, these studies were excluded for this endpoint. The REDUCE trial did not show a statistically significant increased risk for patients using dutasteride for detectable...
Gleason 7-10 cancers (MHRR: 0.95, 95% CI 0.79-1.14), as did the CombAT trial (MHRR: 0.63, 95% CI 0.38-1.04). The test for heterogeneity illustrates moderate to high heterogeneity ($I^2 = 56.84$), but was not statistically significant ($Q = 2.32$, $p = 0.128$), and was therefore reasonable to pool the results. The meta-analysis did not find a statistically significant increased risk for detectable Gleason 7-10 cancers for patients using dutasteride, either as monotherapy or in combination with tamsulosin, compared to placebo or tamsulosin monotherapy (MHRR: 0.83, 95% CI 0.56-1.21).

Gleason 8-10 cancers

On risk of detectable Gleason 8-10 prostate cancers, we describe the effect of dutasteride (0.5 mg daily) or dutasteride (0.5 mg daily) + tamsulosin (0.4 mg daily) versus placebo or tamsulosin (0.4 mg daily) (Fig. 4). Gleason scoring was not prospectively collected in the BPH monotherapy studies, and therefore, was excluded for this endpoint. The REDUCE trial showed no statistically significant increased risk for patients using dutasteride for detectable Gleason 8-10 cancers (MHRR: 1.53, 95% CI 0.86-2.73), as did...
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CombAT (MHRR: 0.59, 95% CI 0.26-1.31). The test for heterogeneity illustrates high heterogeneity ($I^2 = 72.37$), but was not statistically significant at the pre-specified alpha level of $p < 0.05$ ($Q = 3.62, p = 0.06$), and represented only dutasteride randomized controlled trials that measured this endpoint. It is understood that heterogeneity statistics may lack necessary power, however, it was felt that pooling the data (as was done for Gleason 7-10 cancers) provided additional scientific insight. There was no evidence of a significant increased risk for detectable Gleason 8-10 prostate cancers in the meta-analysis in subjects using dutasteride, administered alone or in combination with tamsulosin, compared to placebo or tamsulosin monotherapy (MHRR: 0.99, 95% CI 0.39-2.53).

**Discussion**

This meta-analyses pooled data from over 9500 men and represented an exposure of 25 900 patient-years. In the dutasteride arms, the data demonstrated a statistically significant reduction in the overall risk of detectable prostate cancer by 34% to 37% (any-prostate-cancer and for-cause cancers using classical Gleason scores) in men with BPH or in men at an increased risk of prostate cancer. From the meta-analysis of CombAT and REDUCE, there was no significantly increased risk of detectable Gleason 7-10 or Gleason 8-10 prostate cancers in men taking dutasteride, either alone or in combination with tamsulosin, over placebo or tamsulosin monotherapy.

These data are consistent with published results for the treatment of BPH in symptomatic men with enlarged prostates. The BPH monotherapy studies showed a significantly reduced risk of detectable prostate cancer of 43% at 24 months and 51% at 27 months in men taking dutasteride over placebo. High-grade tumours were not reported in the BPH monotherapy trials. The CombAT study showed a 40% relative risk reduction of detectable prostate cancer compared to tamsulosin monotherapy over 4 years (combination therapy vs. tamsulosin: 43%, $p = 0.006$; dutasteride monotherapy vs. tamsulosin: 37%, $p = 0.021$). This reduction in prostate cancer with dutasteride, alone or in combination with tamsulosin, was observed across all Gleason scores. Over the 4 year study, there were numerically fewer Gleason score 7-10 tumours (combination: $n=12$; dutasteride: $n=22$; tamsulosin: $n=27$) and Gleason score 8-10 tumours (combination: $n=5$; dutasteride: $n=8$; tamsulosin: $n=11$) in the dutasteride groups combined, compared with the tamsulosin group. The reduction in risk of detectable prostate cancers is further confirmed in the recently published REDuction by Dutasteride of clinical progression Events in Expectant Management (REDEEM) trial, in which numerically fewer cases of Gleason 8 tumours in the dutasteride arm ($n=2$) compared to placebo ($n=3$) over 3 years were observed, in men who already had low-risk, low-grade cancers (T1c-T2a, Gleason score <6 and no Gleason pattern score ≥ 4). No Gleason 9 or 10 tumours were detected.

Concern has been raised about the increased incidence of Gleason 8-10 high-grade tumours in the REDUCE and PCPT studies. When modified Gleason scores were used, an increased incidence of Gleason 8-10 cancers with finasteride (1.8%) and dutasteride (1.0%) over placebo (1.1%, 0.5%, respectively) was reported. Furthermore, no increase was seen in the detection of Gleason 7-10 cancers, and a lower incidence was observed in low-grade cancers.

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*Fig. 3. Effect of dutasteride (0.5 mg daily) or dutasteride (0.5 mg daily) + tamsulosin (0.4 mg daily) vs. placebo or tamsulosin (0.4 mg daily) on prostate cancer risk of Gleason grade 7-10 cancers.*
It is unclear whether the increased incidence of high-grade tumours was due to histopathologic artifacts such as selective reduction in overall prostate volume or low grade prostate cancer lesion caused by 5ARIs, rather than representing actual biologic activity of the tumour. However, the possibility that 5ARIs may increase the growth or the development of high grade prostate cancer cannot be ruled out. This has led to increasing concern amongst physicians and patients about the safety of 5ARIs in the management of BPH.

There are differences among the baseline patient characteristics in these clinical studies. The patient population enrolled in the BPH studies, defined by moderate to severe symptoms, prostate volume ≥30 cc, PSA ≥1.5 ng/mL and IPSS ≥12, is the more clinically relevant patient population that is prescribed a 5-ARI to manage BPH symptoms, according to national guidelines in Canada, the European Union and the U.S. In contrast, the REDUCE study included a population that was already at a higher risk of developing prostate cancer, as defined by the increased PSA values (>2.5 ng/mL) and higher prostate volumes (≤80 cc). Despite these differences, in this meta-analysis, dutasteride showed a reduction in risk of detectable prostate cancer (any-prostate-cancers and for-cause), and there was no significant increase in the detection of high-grade tumours between the CombAT and REDUCE populations (groups which ranged in baseline prostate volumes and PSA values) (Table 1).

The findings in the present study are further supported by the recent Consensus Statement from the Canadian Urological Association. The statement critically analyzed the available data on the benefit-risk of 5-ARIs. The CUA panel stated that 5-ARIs can be safely used to treat BPH in men with an enlarged prostate; however, the panel reiterated the need for a robust conversation between the physician and the patient on the risks and benefit of treatment. Regular monitoring using PSA was further reinforced.
Limitations

There were several limitations in this meta-analysis. Gleason scoring was not prospectively collected in the BPH monotherapy (ARIA3001, ARIA3002, ARIB3003) studies and this may have affected our results. Gleason scores were also not included in the analyses of Gleason 7-10 and Gleason 8-10 cancers. In CombAT and REDUCE, only for-cancer biopsies were conducted, whereas REDUCE called for both protocol-defined and for-cancer biopsies. Additionally, the Gleason scoring methodology was not consistent between the CombAT and REDUCE trials. There was no central pathology review in the dutasteride monotherapy and CombAT studies, resulting in lack of standardization and different interpretations and grading for the primary endpoint of interest in these analyses. The meta-analyses did not adjust for number of biopsies in the different trials. Finally, the analyses pooled trial results that had fundamentally different designs, population and methodology to evaluate prostate cancer status, and statistical tests for three of the four outcomes analyzed presented with moderate to high heterogeneity.

Conclusions

This is the first study to evaluate the impact of dutasteride for prostate cancer detection based on a meta-analysis of all GSK-sponsored trials. While the analysis had several limitations, the results showed that dutasteride reduced the risk of detectable prostate cancer, including low-grade prostate cancers, which are less likely to lead to death. Additionally, in our analysis, there was no increased risk of detectable high-grade prostate tumours, defined by Gleason ≥7 scores, as shown in CombAT and REDUCE. These data support the use of dutasteride in the typical clinical management of BPH, with patients appropriately monitored by their physicians.

Competing interests: All authors of this manuscript are employees of GlaxoSmithKline.

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References


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