

# UCSF

## UC San Francisco Previously Published Works

### Title

Fifth Joint Meeting of J-CaP and CaPSURE: Advancing the Global Understanding of Prostate Cancer and Its Management

### Permalink

<https://escholarship.org/uc/item/6gd2s0fj>

### Journal

Japanese Journal of Clinical Oncology, 42(3)

### ISSN

0368-2811

### Authors

Akaza, Hideyuki  
Carroll, Peter  
Cooperberg, Matthew R  
et al.

### Publication Date

2012-03-01

### DOI

10.1093/jjco/hyr194

Peer reviewed

Meeting Report

## Fifth Joint Meeting of J-CaP and CaPSURE: Advancing the Global Understanding of Prostate Cancer and Its Management

Hideyuki Akaza<sup>1,\*</sup>, Peter Carroll<sup>2</sup>, Matthew R. Cooperberg<sup>3</sup> and Shiro Hinotsu<sup>4</sup>

<sup>1</sup>Department of Strategic Investigation on Comprehensive Cancer Network, Research Center for Advanced Science and Technology, The University of Tokyo, Tokyo, Japan, <sup>2</sup>Department of Urology, Helen Diller Family Comprehensive Cancer Center, University of California, San Francisco, <sup>3</sup>Urology, Urologic Oncology, UCSF Helen Diller Family Comprehensive Cancer Center, San Francisco, CA, USA and <sup>4</sup>Department of Pharmacoepidemiology, Graduate School of Medicine and Public Health, Kyoto University, Kyoto, Japan

\*For reprints and all correspondence: Hideyuki Akaza, Department of Strategic Investigation on Comprehensive Cancer Network, Research Center for Advanced Science and Technology, The University of Tokyo, 4-6-1, Komaba, Meguro-ku, Tokyo 1530041, Japan. E-mail: akazah@med.rcast.u-tokyo.ac.jp

Received October 25, 2011; accepted December 7, 2011

This report summarizes the presentations and discussions that took place at the Fifth Joint Meeting of J-CaP and CaPSURE held in Tokyo, Japan, in July 2011. The J-CaP and CaPSURE Joint Initiative was established in 2007 with the objective of analyzing, reviewing, comparing and contrasting data on prostate cancer patients from Japan and the USA within the two important large-scale, longitudinal, observational databases—J-CaP and CaPSURE. Since its inception, the initiative has reviewed a wide range of topics and generated valuable data on the patterns of prostate cancer treatment and patient outcomes in the two geographical regions. The objectives of this 5th Joint Meeting were to provide an update on the current status of the J-CaP and CaPSURE databases, and also to discuss perspectives from a range of other Asian countries—Japan, China, Indonesia and Korea—on the use of androgen deprivation therapy for prostate cancer. The collaborators recognize that large databases, such as J-CaP and CaPSURE, provide valuable ‘real-world’ information, to complement data from clinical trials, which can help to advance the clinical management of prostate cancer patients worldwide. It is anticipated that in the near future, the Joint Initiative will expand globally to include patient registries from other countries so that best practice can be shared and regional differences in patients, treatments and outcomes can be explored.

*Key words: prostate cancer – androgen deprivation therapy – overall survival – risk stratification – J-CAPRA*

### OVERVIEW

The J-CaP and CaPSURE Joint Initiative was established in 2007 with the overall objective of analyzing, reviewing, comparing and contrasting data on prostate cancer patients from Japan and the USA within the two important large-scale, longitudinal, observational databases—J-CaP and CaPSURE. The aim of this ongoing collaboration is to identify trends within patient characteristics, treatment

approaches and outcomes (clinical, economic and health-related quality of life) that might assist physicians, patients and others when selecting treatment options at different prostate cancer disease stages. The collaborators recognize that patient registries are an important source of data on therapeutic efficacy and patient outcomes in the ‘real-world’ situation and are an invaluable complement to data obtained from randomized clinical trials.

This report summarizes the presentations and discussions that took place at the Fifth Joint Meeting of J-CaP and CaPSURE held in Tokyo, Japan, in July 2011. At the four previous meetings of the Joint Initiative, a wide range of topics have been addressed and valuable data have been generated on the patterns of prostate cancer treatment and patient outcomes in the two geographical regions. One important tool that has been developed by the group is a modified CAPRA risk stratification method, named 'J-CAPRA'. This novel risk instrument is the first to be developed and validated for patients undergoing primary androgen deprivation therapy (PADT) and is applicable to those with both localized and advanced disease. It performs well in diverse populations more relevant than existing instruments to the high-risk patients that form a large proportion of the J-CaP database and (5).

In Japan, the J-CaP database was established in 2001 when the Japan Study Group of Prostate Cancer (J-CaP Study Group) supported by the Japan Kidney Foundation and authorized by the Japanese Urological Association commenced a study to gather information about hormone therapy administered to Japanese patients and to analyze the outcomes of treatment. This was prompted by the extensive use of the different forms PADT, including combined androgen blockade (CAB) therapy, for the treatment of prostate cancer within Japan. The result was the J-CaP registry, a large, multicenter, population-based database of men newly starting PADT for prostate cancer. The rationale for its development and an interim analysis of the registration status of the patients and their background variables was reported in 2003 (1), and more recently, treatment patterns with PADT have been reported along with an interim analysis of prognosis (12). As of 2005, J-CaP included data for 26 272 patients from 406 institutions comprising 77 University Hospitals (67% of those in Japan), 267 General Hospitals and 62 Private Hospitals. Around 50% of new prostate cancer patients treated with hormone therapy in Japan were registered with J-CaP at that time.

In the USA, the CaPSURE database was founded in 1995 and currently contains data on around 14 000 prostate cancer patients treated with all forms of therapy. The objectives of the CaPSURE initiative are to collect longitudinal data on a large cohort of patients with prostate cancer, to record existing and developing patterns in these data and in patient outcomes and to identify any variables that might predict outcome. Data on socio-demographics, clinical characteristics, resource use and health-related quality of life are collected prospectively from patients, community-based urologists and hospital records at centers across the USA.

The objectives of this Fifth Joint Meeting of J-CaP and CaPSURE were to provide an update on the current status of the J-CaP and CaPSURE databases and to discuss perspectives from a range of Asian countries—Japan, China, Indonesia and Korea—on the use of androgen deprivation therapy (ADT) for prostate cancer. The latest comparative data from the J-CaP and CaPSURE databases were also

reviewed. The meeting was co-chaired by Professor Hideyuki Akaza (The University of Tokyo, Japan) and Professor Peter Carroll (University of California, San Francisco, CA, USA).

### **PRESENTATION 1: WHAT CONCLUSIONS CAN BE DRAWN USING INFORMATION FROM LARGE-SCALE DATABASES? REGISTRY DATA VERSUS RANDOMIZED CONTROLLED TRIALS**

S.H. (Kyoto University, Kyoto, Japan) discussed the advantages and disadvantages of the types of information that could be drawn from large-scale databases, such as J-CaP and CaPSURE, in contrast to that which could be concluded from randomized, controlled trials (RCTs).

A range of different designs of clinical study can contribute to the different levels of evidence specified in treatment guidelines, such as those of the European Association of Urology (11). These include meta-analysis, randomized, case-controlled, observational, cohort, cross-sectional, plus retrospective medical record reviews. When comparing between RCTs and observational studies, each has its own particular advantages: RCTs eliminate confounding and minimize bias while observational studies have the advantage of greater generalizability, studies can be conducted more rapidly, expense is minimized and they address a broader range of questions.

Confounding factors are recognized as an inherent problem in observational studies—these are factors associated with the risk factor being observed and are causally related to the outcome. Methods to eliminate confounding include randomization (as in RCTs), matching (as in case-controlled trials) and exclusion of certain patients. In addition, in observational studies, the statistical methodology employs stratification and multivariate analyses.

Generalizability of study data refers to the ability to apply the results of a study to patient populations other than the study sample. In general, the results of RCTs only apply to a selected patient population as identified in the study protocol, and many patients are specifically excluded from a trial. As an example, the results of a study performed in patients with good renal function might not be generalized to patients with renal dysfunction. In observational studies, no such selection is performed and as a result their generalizability is increased and they are more likely to reflect treatment effects in the real world.

Results from RCTs and observational studies, including data from large-scale databases such as J-CaP and CaPSURE, should therefore be considered as complementary: information from both is of value and should be considered to help build a full clinical picture and inform clinical decision making.

S.H. cited the example of the results of an RCT evaluating CAB with bicalutamide for advanced prostate cancer (2).

The study reported survival outcomes after long-term follow-up of a Phase III RCT. Five-year survival with CAB was 75.3% and with luteinizing hormone-releasing hormone (LHRH) alone was 63.4% ( $P = 0.043$ ). These findings are supported by results from the J-CaP database of overall survival in high-risk patients (J-CAPRA score 8+) which also found a significant difference in favor of CAB compared with non-CAB hormonal therapy ( $P < 0.0001$ ).

### **PRESENTATION 2: CONFOUNDING FACTORS: AN INHERENT PROBLEM WHEN COMPARING DIFFERENT DATABASES**

P.C. [University of California, San Francisco (UCSF), CA, USA] provided the meeting participants with some insight into an inherent problem often encountered when analyzing and comparing information from observational databases—confounding factors. A confounder can affect the relationship between the predictor and outcome leading to bias, and it is therefore important to control for confounders in order to get closer to the true relationship between the predictor and outcome. In observational research, confounding is commonly addressed by adjusting or controlling for these potentially confounding factors. However, databases differ in the demographic and clinical factors they capture and the outcomes they assess and therefore some confounding factors might not be recorded in the database. As a result, these cannot be adjusted for, leading to potential bias.

P.C. went on to describe how a range of studies investigating an association between ADT and cardiovascular (CV) mortality had shown mixed results in observational studies, while RCTs had failed to show any such association. An analysis was initially undertaken on the CaPSURE database to assess mortality outcomes by treatment type with a particular focus on the effects of ADT and CV mortality. The results suggested that there was an increase in CV mortality in some men receiving ADT (27). Because of the conflicting data in the literature, this subject was addressed in the CaPSURE database more recently. The results showed an almost 2-fold increase in CV mortality in the ADT group compared with those receiving local treatment only. However, there was also a greater than 2-fold increased CV mortality in the watchful waiting/active surveillance group compared with local treatment alone. It was therefore considered that perhaps factors affecting treatment selection, e.g. age or co-morbidity, might be contributing to CV risk. These data were also analyzed using propensity scoring—a measure of the patient's probability of receiving treatment (ADT), conditional on important clinical and socio-demographic covariates (21). No differences in CV mortality were noted between those who did and did not receive ADT inferring that unmeasured variables—which influence treatment decision-making but are not recorded in CaPSURE—might have confounded the association between ADT and CV mortality in many previous studies.

He concluded that it could be challenging to properly adjust risk for outcomes like CV mortality. Although CaPSURE assessed the presence or absence of important comorbidities like cardiac disease and diabetes, it did not record HbA1c levels, heart failure risk classifications and other important risk factors for cardiac death. Therefore, it was possible that these variables might be confounding the association between ADT use and CV mortality. Matching patients on their propensity to receive ADT was an effective method of attempting to control for confounding in observational studies, although it could not eliminate it completely.

### **PRESENTATION 3: OBSERVED DIFFERENCES BETWEEN THE J-CaP AND CaPSURE DATABASES IN THE SURVIVAL OF PATIENTS RECEIVING PADT**

H.A. (Research Center for Advanced Science and Technology, The University of Tokyo, Tokyo, Japan) reminded participants that any hormonal therapy for prostate cancer should take into account not only testosterone of testicular origin but also the locally produced androgens: testosterone and dihydrotestosterone derived from dehydroepiandrosterone of adrenal origin. CAB—either medical or surgical castration plus an antiandrogen—is often the therapy of choice when the tumor is localized; however, resistance to the disease can develop if the tumor becomes metastatic. This is thought to be due to either elevated levels of the androgen receptor or to local biosynthesis of androgens.

New agents for the management of prostate cancer, in particular more potent antiandrogens, are continually being sought and H.A reviewed some recent examples that have shown promising results in patients with castration-resistant prostate cancer (CRPC).

Abiraterone acetate, an inhibitor of androgen biosynthesis, has recently received approval from the FDA for the treatment of metastatic CRPC. The results of a Phase III RCT have been published (10) which demonstrated that abiraterone acetate was able to prolong the overall survival among patients with metastatic CRPC who had previously received chemotherapy.

MDV3100 is an androgen receptor antagonist which also induces tumor cell apoptosis. Results of a Phase I–II trial have shown that MDV3100 has promising antitumor activity in patients with CRPC and also provides evidence that sustained androgen receptor signaling is a key driver in this disease (22).

XL184 (cabozantinib) is an inhibitor of tumor growth, metastasis and angiogenesis, which simultaneously targets MET and VEGFR2, key kinases involved in the development and progression of many cancers. Interim results of a Phase II clinical trial of cabozantinib for the treatment of men with metastatic CRPC have shown encouraging clinical activity in these patients (24).

The initial concept of the benefit of CAB therapy over LHRH monotherapy was proposed in 2002 (14); however, several other studies going back over 20 years have also demonstrated the benefit of this regimen (15): Crawford *et al.* (9) concluded that in patients with advanced prostate cancer, CAB treatment with leuprolide plus flutamide was superior to treatment with leuprolide alone, and more recently, Akaza *et al.* (2) showed that CAB with bicalutamide 80 mg offered a significant overall survival benefit compared with LHRH agonist monotherapy without reducing tolerability in patients with locally advanced or metastatic prostate cancer.

Despite such evidence, the National Comprehensive Cancer Network (17) Clinical Practice Guidelines in Oncology do not currently recommend primary hormone therapy for localized prostate cancer. However, the NCCN Clinical Practice Guidelines in Oncology: Asia Consensus Statement (18) focuses on the specific situation for Asian patients and takes into account the differences between Asian and Caucasian patients in their response to therapy. Primary ADT is recommended in these guidelines as an additional option in low-, intermediate- or high-risk Asian patients with a life expectancy of <10 years.

Published data on the benefits of CAB from RCTs are now supported by observational data from the J-CaP database, which although not randomized does include a large number of patients. H.A. presented data showing that for low-risk patients (J-CAPRA score 0–2), there is no observed difference in overall survival between those receiving CAB therapy or non-CAB therapy and by 10 years overall survival does not drop below 50% in either group. Overall survival of intermediate-risk (J-CAPRA score 3–7) and high-risk (J-CAPRA score 8+) patients in the J-CaP database is significantly higher for those receiving CAB therapy compared with non-CAB therapy ( $P < 0.0001$ ). In the intermediate-risk group, overall survival only drops below 50% after 8 years and for high-risk patients after 4 years in each treatment group (Fig. 1a).

He then compared the J-CaP data with those analyzed by the J-CAPRA score from the CaPSURE database (Fig. 1b). Interestingly, in the low-risk group in the CaPSURE cohort, 50% overall survival is reached by 6 years, much shorter than that seen in the low-risk J-CaP cohort. Notably, in the CaPSURE database, the overall survival of low- and intermediate-risk patients is better for non-CAB treated patients than for CAB-treated patients—the opposite of that seen in the J-CaP database. Only high-risk patients in the CaPSURE database show improved survival with CAB compared with non-CAB therapy.

He concluded by saying that to explain these observed differences between the two databases, it is important to consider carefully the influence of possible confounding factors. It is possible that even within a single category, such as the intermediate risk group with J-CAPRA scores of 3–7, there were patients who responded differently to therapy.

## PRESENTATION 4: ASIAN PERSPECTIVES ON THE USE OF ADT FOR PROSTATE CANCER

### KOREA

B.-H.C. (Yonsei University College of Medicine, Seoul, Korea) reported that data from the National Statistics Office in Korea showed that the number of patients diagnosed with prostate cancer had quadrupled between 2002 and 2008 (16,19). A survey of 50 urologists demonstrated an increase in the average number of prostate cancer patients seen each month, which had risen to an average of 70 per month in 2009 in the outpatient setting (unpublished data). These patients have predominantly localized disease (37%) with 23% having locally advanced disease, 29% advanced (metastatic) disease and 12% hormone-refractory disease. There appears to be a stage migration to a greater proportion of patients with localized disease and a reduction in the numbers with advanced disease.

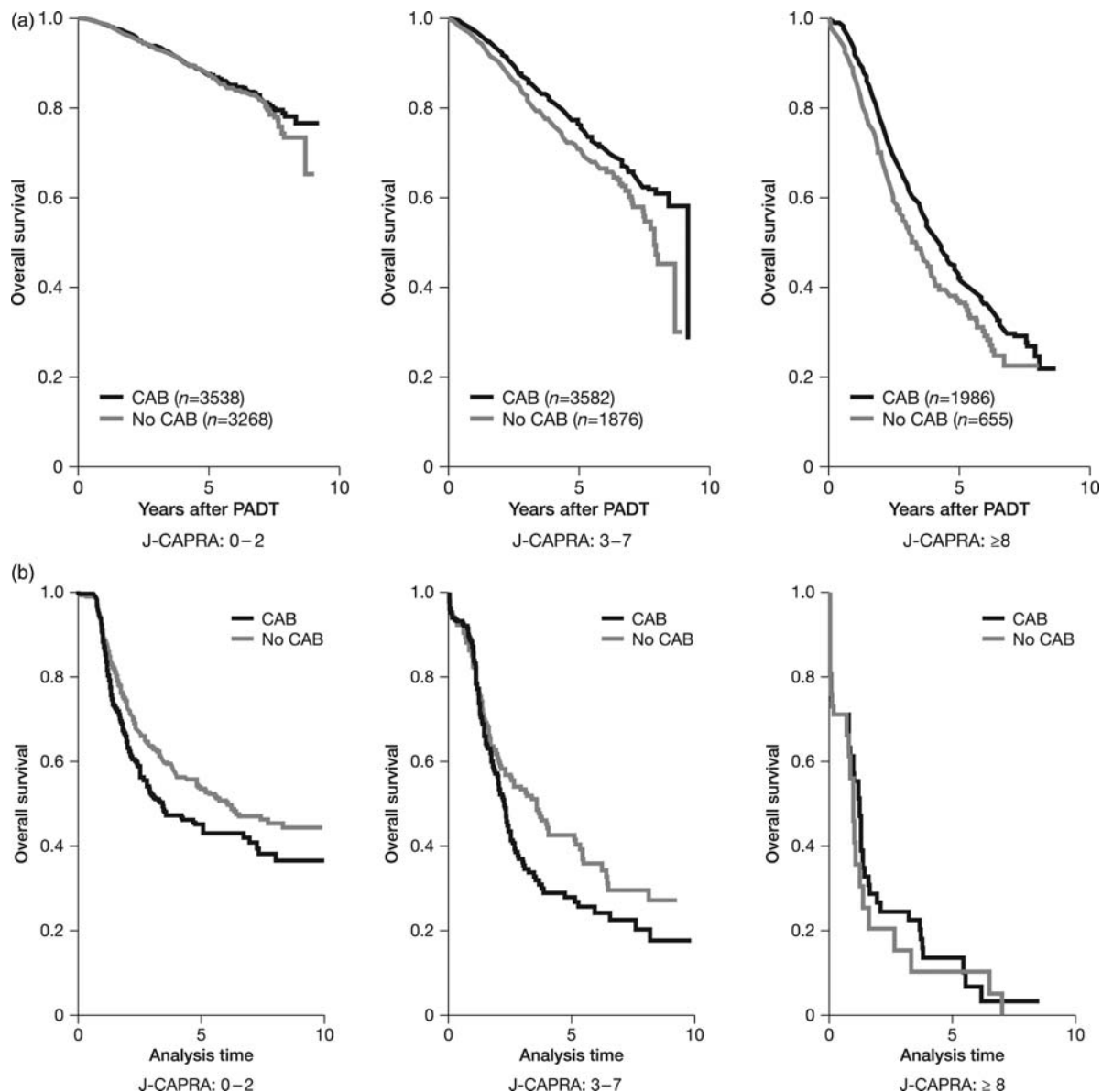
In terms of treatment modalities used in Korea, hormone therapy alone is administered to 13% of the patients with localized disease, 17% with locally advanced disease, 88% with advanced disease and 37% with hormone-refractory disease. In the case of localized or locally advanced disease, often it is used in combination with radical prostatectomy or radiotherapy. He went on to compare these survey data with those from his own institution, the Gangnam Severance Hospital, where a total of 157 patients were newly diagnosed with prostate cancer between 2008 and 2010. Hormone therapy alone is administered as a primary therapy to 12% of the patients with localized disease, 25% of those with locally advanced disease and 76% of those with advanced disease.

The available ADT for prostate cancer in Korea includes leuprolide acetate, leuprorelin acetate, goserelin acetate, bicalutamide (50 mg and 150 mg), cyproterone acetate (50 mg) and flutamide (250 mg).

The Korean Urological Oncology Society issued guidelines in 2008 for the use of ADT for prostate cancer. In contrast to the situation in Japan where ADT is used as a primary therapy in around 80% of low-risk patients, primary ADT is not commonly used in Korea for this patient group. Again, in contrast to the recommendations for high-risk patients within the NCCN Clinical Practice Guidelines in Oncology: Asia Consensus Statement, CAB with bicalutamide (80 mg daily) is not used in Korea as this dose was not available; bicalutamide 50 or 150 mg daily was used instead, according to efficacy and the patient's quality of life. B.-H.C. concluded that Korean treatment patterns for prostate cancer patients are similar to those seen in the USA and Europe, rather than in Japan, and primary ADT is not the preferred choice for T1c–T3N0M0 patients (26).

### CHINA

X.Z. (Peking University, China) reported that the overall incidence of prostate cancer in Asia is relatively low at around



**Figure 1.** Overall survival by type of hormone therapy according to the J-CAPRA score in (a) the J-CaP database and (b) the CaPSURE database.

90 per 100 000 men, but there are variations in the incidence across different Asian regions. In the city of Shanghai, China, the incidence is 2–3 per 100 000 men. Ten-year data from 1990 to 2000 show a sharp 3-fold increase from a standardized incidence of 2.7 to 7.7.

When data for the incidence of different forms of cancer in various cities in China from the period 1993–97 are compared with that for the period 1998–2002, it can be seen that prostate cancer was the 10th most common form of cancer in Shanghai in the period 1993–97 but has risen to become the eighth most common in the period from 1998–2002.

Hormonal therapy, particularly CAB treatment, is widely used in China, based on guidelines for prostate cancer treatment issued by the Chinese Urological Association. X.Z. advised that the types of patients who are selected for hormone therapy generally fall into four categories:

metastatic prostate cancer (clinical stage M), those given it prior to radical prostatectomy (clinical stage T3a), those given it following radical prostatectomy (pathological stage T1–T3) or those receiving radiation therapy (RT) (clinical stage T3–T4, M0). The usual treatment modality is CAB in the majority of patients, while antiandrogen alone, medical castration alone or surgical castration alone are only administered to around 10% of patients each.

He concluded by noting that the incidence of prostate cancer in China was likely to increase significantly in future years, particularly in major cities.

#### INDONESIA

R.U. (University of Indonesia, Depok, Indonesia) reported that prostate cancer had accounted for 40% of all the

urological malignancies diagnosed between 1995 and 2010 at two hospitals in Jakarta, Indonesia—RSCM and RSKD. The incidence of prostate cancer at these centers had increased around 3-fold from the period 1995–99 (135 cases) to 2005–09 (404 cases). In many cases, patients presented with advanced, metastatic disease; only about 38% had non-metastatic disease.

He went on to describe the treatment options administered to patients with T1–T4,N0,M0 prostate cancer from 1995 to 2010 ( $n = 262$  patients) at these two institutions. Hormonal therapy had been given to 22% of the patients with T1 disease, 29% with T2, 57% with T3 and 94% with T4. The type of hormonal therapy given was either orchiectomy (about one-third of the patients), LHRH monotherapy or intermittent androgen blockade (IAB) using CAB [6 months on treatment and 6 months off, depending on prostate specific antigen (PSA) levels]. R.U. had compared the patients treated with IAB and orchiectomy at these institutions but found no statistically significant difference in survival between the two treatment groups.

Data collected for the period 2004–08 from four cities in Indonesia (Jakarta, Bandung, Surabaya, Yogyakarta;  $n = 851$  cases) confirmed that ADT is the most commonly used prostate cancer treatment (around 60% of the cases) in Indonesia.

#### JAPAN

Y.H. (Nara Medical University, Kashihara City, Nara, Japan) reported results of a study evaluating the distribution of primary therapy and clinical characteristics of over 2000 prostate cancer patients at Nara Medical University and its 23 affiliated hospitals (25). The study had also investigated the differences between the preferential primary therapy conceived by the primary doctors and the primary therapy actually administered for prostate cancer patients.

Data from 2004 to 2006 showed that of the 2371 patients included in the analysis, 30% had undergone radical prostatectomy (RP), 13.8% RT and 50.1% PADT. However, when Nara Medical University (NMU) was compared with its affiliated hospitals (non-NMU), differences were observed in the treatment modalities used. Non-NMU hospitals were found to treat a much higher proportion of patients with PADT (55.7%) compared with NMU (24.3%) and a lower proportion with RT (7.4%) compared with NMU (42.8%). The proportion of patients treated with RP was similar between NMU and non-NMU (around 30%). The likely reason is that RT (IM-RT and brachytherapy) is more easily available at the NMU than at affiliate hospitals.

Data from NMU comparing the two different time periods (2004–06 and 2007–09) showed that the trends in patients' ages, PSA levels, Gleason scores, clinical stage and risk classification of prostate cancer patients diagnosed between 2007 and 2009 were significantly lower than those diagnosed between 2004 and 2006 ( $P < 0.001$ ). In addition, the proportion of patients treated with PADT from 2007 to 2009 was

significantly lower than that seen from 2004 to 2006 (40 vs. 50%). The proportion undergoing RP was similar in both time periods (30%); however, the proportion receiving radiotherapy from 2007 to 2009 was significantly higher than in 2004–06 (24 vs. 14%,  $P < 0.001$ ).

The trend in primary therapy was significantly different between university hospital and affiliated hospitals in both time periods (2004–06 and 2007–09). Within NMU, over the two time periods, there was a trend to increasing use of RT with a decrease in RP and the use of PADT. Non-NMU hospitals did not show much change in the use of primary therapy over this time other than a slight decrease in the use of PADT.

Y.H. then compared data for NMU (2004–06 and 2007–09) with that from the CaPSURE database (1989–2002) according to risk: T-stage, Gleason score, PSA and D'Amico risk score. The results showed that Japan appeared to be 15–20 years behind the USA in terms of the pattern of prostate cancer risk, having a greater proportion of high-risk patients. When comparing primary treatment used at NMU and recorded in CaPSURE, substantial differences were observed. In NMU, a much smaller proportion of patients underwent watchful waiting (around 3%) compared with CaPSURE (around 23%) while a much higher proportion were treated with PADT (40%) compared with CaPSURE (17%).

He summarized by noting that while the number of high-risk patients at NMU had decreased significantly from 2004–06 to 2007–09, as had the proportion of patients treated with PADT, the proportion of high-risk patients was still greater than that seen in the CaPSURE database.

### PRESENTATION 5: EVALUATION OF THE OUTCOMES OF PATIENTS RECEIVING CAB THERAPY USING THE J-CAPRA RISK SCORE

#### RESULTS FROM THE J-CAP DATABASE

S.H. (Kyoto University, Kyoto, Japan) noted that a major difference between the J-CaP and CaPSURE databases related to age distribution: in the J-CaP database, almost 50% of the patients were aged 75 years or over and around 90% were aged 66 years or over. In contrast, in CaPSURE only around 15% were aged 75 years or over and almost 50% were 65 years of age or younger. Analysis of age distribution and risk according to the J-CAPRA score showed that a higher proportion of younger patients had high-risk disease compared with older patients; this was the opposite of the trend seen in CaPSURE. When the trend in the type of PADT employed is analyzed for the period 2001–03, it is apparent that there were no significant changes in treatment trends over this time with the majority of patients (about 60%) receiving PADT.

An important factor potentially affecting treatment outcomes in any patient is the presence of co-morbidities. Within the J-CaP database, a range of co-morbidity factors is

recorded; however, it is not possible to specify the severity of the co-morbidity or assign any importance or degree of risk. S.H. described how using Charlson's comorbidity index co-morbid conditions can be 'weighted' depending on the risk of dying associated with this condition to give a score that predicts mortality. Unfortunately, this could not be applied accurately to patients in the J-CaP database since certain data were lacking—the database does not record all conditions, e.g. dementia, leukemia, lymphoma, AIDS, plus it did not record the severity of liver or renal disease. In view of this, an alternative method of analysis was employed: patients were assigned as 'cardiovascular disease (CVD)+' if they had either heart disease or stroke. Then, the combination of number of different co-morbidities plus the presence or absence of CVD was assigned to four categories: no co-morbidities (6248 patients),  $\leq 2$  co-morbidities but CVD- (7341 patients),  $\leq 2$  co-morbidities but CVD+ (2990 patients) and  $> 2$  co-morbidities (2458 patients). An analysis was then undertaken of co-morbidity distribution according to the J-CAPRA score (low, intermediate and high), age range ( $\leq 65$ , 66–75 and  $> 75$  years) and the type of hormonal therapy selected (CAB or non-CAB). The results showed that the distribution of the number of co-morbidities was comparable between CAB and non-CAB treatment regimens.

S.H. went on to describe the results of the survival analysis of J-CaP patients. Both cause-specific and overall survival showed similar separations of the curves for patients with low, intermediate and high J-CAPRA scores. In terms of overall survival, for low-risk patients, there was no difference in survival between CAB and non-CAB therapy; however, in both intermediate- and high-risk patients, the prognosis was better with CAB than with non-CAB therapy.

In order to evaluate these differences, PADT treatment of patients in the J-CaP database was reclassified according to the following three categories: (i) orchiectomy, (ii) LHRH monotherapy ( $n = 2991$ ) or LHRH plus short antiandrogen ( $n = 1657$ ) or (iii) CAB ( $n = 10\,369$ ). In terms of cause-specific survival, CAB and LHRH monotherapy resulted in better prognosis than orchiectomy in patients with an intermediate J-CAPRA score (3–7). CAB therapy resulted in a better prognosis than LHRH monotherapy or orchiectomy in patients with a high J-CAPRA score ( $> 8$ ). No significant differences were observed between CAB and LHRH monotherapy according to the J-CAPRA score in terms of background factors, such as age, PSA at diagnosis and Gleason score at diagnosis.

#### RESULTS FROM THE CAPSURE DATABASE

M.C. (University of California, San Francisco, CA, USA) reported that the prostate cancer mortality rate in the USA has fallen by 40% in the PSA screening era at a time when men were generally living longer and should in fact be at greater risk from prostate cancer mortality (13). However, he

noted that the observed treatment patterns in the USA give rise to concerns about both overtreatment and undertreatment of many patients. Few men with low-risk disease undergo watchful waiting/active surveillance and, the improved mortality rates have come at the cost of many other men being treated unnecessarily with surgery or radiation for tumors that would ultimately have not been fatal.

He noted that PADT is commonly used in the USA, particularly for higher risk patients (6). This raises the issue of undertreatment of men with high-risk disease who might benefit from local therapy. The reason for this appears to in part reflect an age bias, as reported in a recent publication from the CaPSURE team (4). The results had shown that older patients were more likely to have high-risk prostate cancer at diagnosis but less likely to receive local therapy and that underuse of potentially curative local therapy among older men with high-risk disease might in part explain observed differences in cancer-specific survival across age groups.

When comparing risk groupings and age between J-CaP and CaPSURE, he noted that the patterns were quite different: in J-CaP the oldest patients were more likely to be diagnosed with high-risk disease, and the opposite was true for CaPSURE (5), possibly due to the different screening practices in Japan and the USA. Interestingly, patients in CaPSURE treated with PADT showed similar relationship between age and risk as the J-CaP cohort (4).

Thus, in CaPSURE at least, age appears to be a stronger driver of clinical decision-making than disease risk. Younger patients tend to receive surgical therapy regardless of their disease risk, whereas in the older age group ( $> 75$  years), a much smaller proportion receive local therapy and many high-risk patients only receive ADT monotherapy. Primary treatment trends in CaPSURE from 1999 to 2007 showed that the use of ADT has been relatively consistent over this time period. Two other large databases—Medicare and i3—show similar trends to CaPSURE across all forms of therapy (7). In 2005, however, reimbursement arrangements for ADT in the USA changed significantly and there is now less incentive to prescribe ADT, which might influence treatment patterns.

Results of a study into factors that drive selection of ADT for prostate cancer treatment suggest that the particular urologist a patient sees might be more important in determining whether they receive ADT than tumor or patient characteristics (23). Treatment patterns in CaPSURE also vary markedly across clinical sites, and this variation is not explained by case-mix variability or known patient factors (6).

M.C. advised that there were currently 13 893 patients in the CaPSURE database. Follow-up as of September 2010 showed an all-cause mortality (ACM) of 3752 patients (27.0%) at a median of 7.0 years and a prostate cancer-specific mortality (PCM) of 714 patients at a median of 6.2 years—this represented 5.1% of overall cohort and 19% of all deaths. The overall median follow-up was 4.6 years.

Both ACM and PCM by treatment type in CaPSURE show similar trends, with surgery patients having a better



prognosis than those treated with radiation, followed by watchful waiting/active surveillance and ADT; however, it should be noted that these are unadjusted figures and reflect the particular case mix.

Looking at treatment according to the J-CAPRA risk group over time, it is apparent that in recent years, there has been less use of ADT monotherapy for low-risk patients and an increase in its use for higher risk patients (J-CAPRA score  $\geq 8$ ). For patients treated with ADT, PCM according to the J-CAPRA score shows a similar separation of the curves (low-, intermediate- and high-risk patients) as seen in J-CaP except that they are shifted to the left—i.e. for a given J-CAPRA category, survival is shorter in CaPSURE.

With the accumulation of data now in CaPSURE due to longer follow-up and a greater number of events, cancer-specific mortality can also be analyzed by the individual J-CAPRA score and shows good distinction between the curves with a discriminatory accuracy (*C*-index) of 0.81. Curves for ACM show a similar distribution pattern but slightly more overlap between the curves and a *C*-index of 0.61.

Of the CaPSURE cohort, 2030 men received ADT monotherapy, predominantly either LHRH monotherapy (916 patients) or CAB (864 patients). In general, the use of CAB seems to correlate with risk, with higher risk patients being more likely to receive it. However, if figures are adjusted according to the J-CAPRA score, this divergence disappears and there is no difference between LHRH monotherapy and CAB. A multivariate analysis (Cox's proportional hazards model) also confirms no increased mortality risk with CAB [hazard ratio (HR): 0.92] but an increased risk with orchiectomy (HR: 1.72) compared with LHRH monotherapy. Similarly, predicted 10-year survival curves using this model show no difference between LHRH monotherapy and CAB. A competing risks analysis (Fine–Gray) also supports these results.

M.C. summarized by saying that while US guidelines for prostate cancer, such as NCCN(17) and the American Urological Association (3), do not currently endorse the use of PADT, it is commonly used in practice. J-CAPRA scoring provides excellent risk stratification for prediction of mortality for US men receiving PADT and reveals that CAB does not appear to result in improved survival over LHRH monotherapy. These observations are consistent with a meta-analysis undertaken by the Prostate Cancer Trialists' Collaborative Group (PCTCG) of 27 trials per 8275 patients (20). The underlying reasons for the different responses to hormonal therapy across populations remain to be elucidated.

#### **PRESENTATION 6: SIGNIFICANCE OF J-CAPRA SCORE ON THE NEW PROGNOSTIC GROUPING WITHIN THE TNM CLASSIFICATION (7TH EDITION)**

S.H. (Kyoto University, Kyoto, Japan) described how the American Joint Committee on Cancer (AJCC) Cancer

Staging Manual 7th edition had introduced new prognostic groupings for prostate cancer. He reviewed these new groupings in terms of the specified T-stage, N and M category, PSA level and Gleason score for Group I–IV disease and contrasted them with the J-CAPRA score which was calculated from a sum of points assigned to different variables (T, N, M, PSA and Gleason score) from 0–12 score, where low risk scored 0–2, intermediate risk 3–7 and high risk 8 or greater (5).

When TNM prognostic groupings were calculated using data from J-CaP, the results showed that about one-third of the patients fell into Group IV (advanced disease). He went on to examine the relationship between J-CAPRA risk and TNM prognostic grouping for the J-CaP data: all patients identified as high risk by J-CAPRA score ( $>8$ ) were TNM Group IV, those identified as intermediate-risk by J-CAPRA score (3–7) comprised a mixture of Group IIA, II and IV, and those in the low-risk J-CAPRA group comprised a mixture of TNM groups from I to IV.

S.H. concluded by saying that all groups within the TNM prognostic grouping classification are represented within the J-CAPRA low-risk patient group in J-CaP, while about one-third of the patients are TNM Group IV. Analysis of progression-free and overall survival shows that patients in TNM Group IV have a poor prognosis. He therefore suggested that to achieve a balanced proportion and prognosis between TNM prognostic groupings, some modifications should be considered for patients in J-CaP database, for example, creating a subcategory in Group IV, which currently comprises the majority of the patients.

#### **DISCUSSION**

Participants agreed that due to the possibility of confounding factors, it was important to exercise caution when reporting and interpreting data from observational databases such as J-CaP and CaPSURE. The possible confounding factors that might contribute to the differences in the results seen in the two databases were discussed. M.C. commented that it depended whether the endpoint being considered was overall mortality or cancer-specific mortality. For cancer-specific mortality, it was important to confirm that the risk profiles were the same in the two arms, or whether the patients selected for CAB were actually higher risk patients in the first place, as was probably true of the CaPSURE cohort. In the case of overall mortality, it was important to consider what the effects of treatment might be, for example, on CV risk. Patients might have different severities of comorbidities, such as diabetes, and these subtle differences in risk profile were not captured within the database information. P.C. considered that propensity analysis was one way to help address this.

M.N. noted that aside from any confounding factors, there were racial differences in lifetime CV risk profile between Asian and Caucasian patients, so it was difficult

to compare the effect of ADT on CV risk in these two populations. P.C. agreed and commented that conclusions about prostate cancer therapy were often based on the US experience; however, it was a worldwide disease and what was applicable in one population might not be applicable in another, due to environmental or dietary influences, for example. A larger, global registry, as proposed by H.A., would be very valuable in evaluating these differences.

It was noted that the effect of ADT on CV risk had also been analyzed in the J-CaP database and no adverse effects had been found, in line with the CaPSURE data presented by P.C.

P.C. considered that it would be interesting to evaluate the effectiveness and outcomes of different therapeutic modalities in the different populations. It was known from work at UCSF that lifestyle differences after radical prostatectomy could influence treatment outcomes.

It was recognized that patients were now being diagnosed with prostate cancer at a younger age and differences in the treatment of younger versus older patients were discussed. Baseline J-CaP data had been collected over 10 years ago and now new data were being collected on treatment patterns which would allow an analysis of the change in treatment modalities over time.

While the incidence of prostate cancer was increasing in Asia, the stage distribution seemed to be different from that in the USA. It was queried whether routine screening was performed in the countries represented at the meeting. It was noted that in Korea, screening was not done routinely; ~20% are screened but it varies between rural and urban areas. China does not have a routine PSA screening system; it is normally done if the patient presents with symptoms. The underlying reason for the rise in the incidence of prostate cancer in Asian countries was discussed. It was suggested that this could be due to an increase in PSA screening and education about testing, and also to a rise in living standards in certain areas which resulted in better access to health care.

The extensive use of PADT in Japan was possibly due to older urologists who were accustomed to using it to treat the high number of advanced cases seen in previous years and who continued to use it for lower-risk patients. It was considered that younger urologists might be more likely to choose other, newer treatment modalities.

It was noted that intermittent hormonal therapy was popular in some Asian countries. P.C. confirmed that it was also popular in the USA in patients with low-risk disease but cautioned that it may not be optimal for patients with higher risk disease. M.C. added that there were also many inconsistencies between intermittent therapy regimens in terms of PSA trigger, time on and off therapy etc., which made comparison between patients and studies difficult.

M.N. queried why the cause-specific survival of patients in J-CaP who underwent orchiectomy was inferior to that seen with LHRH monotherapy, and also why CAB had

not been found to be superior to LHRH monotherapy. In the case of orchiectomy, he proposed it might be that patients did not receive a secondary therapy if orchiectomy failed. P.C. suggested that there might be a confounding factor and noted that the sample size in the case of orchiectomy was small. S.H. also commented that it could be due to the age distribution of the of the orchiectomy cohort: patients who received orchiectomy were older than other treatment groups, so the number at risk was smaller at an earlier time point in orchiectomy group than in the LHRH group, meaning that a single event would cause a large decrease in survival. It might also reflect the inferiority of the cause-specific survival of patients who underwent orchiectomy. M.C. added that there could be differences in biological treatment effects between LHRH and orchiectomy—*in vitro* studies had suggested a possible direct cytotoxic effect of LHRH therapy.

Secondary treatment effects were considered to be an important issue. S.H. confirmed that in the J-CaP database, only data on initial treatment were recorded routinely. The first 90 days of treatment were recorded as the initial therapy and survival data were calculated for this initial treatment.

The four PSA categories within the J-CAPRA risk scoring system were queried: 0–20 ng/ml = 0, 20–100 ng/ml = 1, 100–500 ng/ml = 2 and >500 ng/ml = 3. These differed from the levels used in some countries, e.g. 0–10, 10–20 and >20 ng/ml. S.H. explained that registration of patients to the J-CaP database had been undertaken for 3 years with 7-year follow-up. Ten years ago, the PSA distribution had been higher so the cut-points had been set relatively high. M.C. confirmed that the J-CAPRA scoring system had been developed for patients who were relatively high risk. He advised that another risk scoring system, CAPRA-S, which incorporated pathological data, had been developed for post-surgical patients (8).

K.N. queried how T-category within CaPSURE was assessed. M.C. confirmed that they recorded both digital rectal examination and ultrasound imaging results and then calculated the stage; few patients underwent magnetic resonance imaging for tumor staging. Stage did not appear to be a meaningful predictor of outcomes until the patient reached T-stage 3.

H.A. commented on the poor prognosis of patients undergoing PADT within CaPSURE compared with those who received local therapy and queried whether there were any underlying reasons for this, e.g. socio-economic factors. M.C. agreed that the survival benefit of ADT in US patients was indeed limited, but the reason for this was still unclear. Within CaPSURE, there was also a 2-fold benefit in survival of surgery over radiation and a 70% benefit of radiation over ADT, although mostly in higher risk patients. The durability of response to a particular therapy may be different in the US and Japanese populations, due to, for example, lifestyle or genetic factors, and these remained to be elucidated.

## CONCLUSION AND FUTURE DIRECTION OF THE J-CaP and CaPSURE JOINT INITIATIVE

H.A. (Research Center for Advanced Science and Technology, The University of Tokyo, Tokyo, Japan) acknowledged that the discussions at the Fifth Joint Meeting of J-CaP and CaPSURE had been extremely thought provoking; however, there were still many unanswered questions. He was convinced that large databases, such as J-CaP and CaPSURE, provided valuable ‘real-world’ information that would help advance clinical management of prostate cancer patients in the future.

P.C. (University of California, San Francisco, CA, USA) stressed the importance of the relationship between the J-CaP and CaPSURE teams and noted the contribution that the initiative had made over the last 5 years to the understanding of prostate cancer in these two distinct regions—the USA and Japan. He was pleased to see that the initiative had now expanded to include input from other Asian countries and hoped to see that the collaboration further expand globally as there was a lot more information about prostate cancer and its management to learn and share between countries.

It was noted that the Korean group was now planning to establish ‘K-CaP’ which would be a complementary database to J-CaP and CaPSURE. In addition, the Korean group had held the 1st Congress of the Asian Pacific Prostate Society (APPS) in March 2011. The President of the APPS was Professor Kim, the Vice-President was Professor Chung and the Secretary General was Professor Lee, all participants at this meeting. Future APPS meetings were already planned: the 2nd APPS Meeting would be held in Seoul, Korea, in 2012 and the 3rd APPS Meeting was to be held in Australia in 2013.

In closing the meeting, the Co-Chairmen thanked all the participants for attending and for their valuable contributions to the discussions, and also acknowledged Takeda Pharmaceutical Company Ltd for their continued support of these joint meetings.

### Conflict of interest statement

None declared.

### References

- Akaza H, Usami M, Hinotsu S, Ogawa O, Kagawa S, Kitamura T, et al. Characteristics of patients with prostate cancer who have initially been treated by hormone therapy in Japan: J-CaP surveillance. *Jpn J Clin Oncol* 2004;34:329–36.
- Akaza H, Hinotsu S, Usami M, Arai Y, Kanetake H, Naito S, Hirao Y. Combined androgen blockade with bicalutamide for advanced prostate cancer: long-term follow-up of a Phase 3, double-blind, randomized study for survival. *Cancer* 2009;115:3437–45.
- American Urological Association. *Guideline for the Management of Clinically Localized Prostate Cancer: 2007 Update*. Available at: <http://www.auanet.org/content/guidelines-and-quality-care/clinical-guidelines/main-reports/proscan07/content.pdf>.
- Bechis SK, Carroll PR, Cooperberg MR. Impact of age at diagnosis on prostate cancer treatment and survival. *J Clin Oncol* 2011;29:235–41.
- Cooperberg MR, Hinotsu S, Namiki M, Ito K, Broering J, Carroll PR, Akaza H. Risk assessment among prostate cancer patients receiving primary androgen deprivation therapy. *J Clin Oncol* 2009;27:4306–13.
- Cooperberg MR, Broering JM, Carroll PR. Time trends and local variation in primary treatment of localized prostate cancer. *J Clin Oncol* 2010;28:1117–23.
- Cooperberg MR, Smith A, Odisho AY, Carroll PR, Litwin MS, Saigal CS. Contemporary nationally-representative trends in treatment for localized prostate cancer. Abstract presented at the American Urological Association Annual Meeting, Washington DC, May 2011.
- Cooperberg MR, Hilton JF, Carroll PR. The CAPRA-S score: a straightforward tool for improved prediction of outcomes after radical prostatectomy. *Cancer* 2011 (Epub ahead of print).
- Crawford ED, Eisenberger MA, McLeod DG, Spaulding JT, Benson R, Dorr FA, et al. A controlled trial of leuprolide with and without flutamide in prostatic carcinoma. *N Engl J Med* 1989;321:419–24.
- de Bono JS, Logothetis CJ, Molina A, Fizazi K, North S, Chu L, et al. Abiraterone and increased survival in metastatic prostate cancer. *N Engl J Med* 2011;364:1995–2005.
- Heidenreich A, Bolla M, Joniau S, Mason MD, Matveev V, Mottet N, et al. *European Association of Urology: Guidelines on Prostate Cancer, 2010*. Available at: <http://www.uroweb.org/gls/pdf/Prostate%20Cancer%202010.pdf>.
- Hinotsu S, Akaza H, Usami M, Ogawa O, Kagawa S, Kitamura T, et al. Current status of endocrine therapy for prostate cancer in Japan analysis of primary androgen deprivation therapy on the basis of data collected by J-CaP. *Jpn J Clin Oncol* 2007;37:775–81.
- Jemal A, Siegel R, Xu J, Ward E. Cancer statistics, 2010. *CA Cancer J Clin* 2010;60:277–300.
- Labrie F, Candas B, Gomez JL, Cusan L. Can combined androgen blockade provide longterm control or possible cure of localized prostate cancer? *Urology* 2002;60:115–9.
- Labrie F. Blockade of testicular and adrenal androgens. *Nat Rev Urol* 2011;8:73–80.
- Ministry for Health, Welfare and Family Affairs. Annual report of cancer incidence (2006) and survival (1993–2006) in Korea; 2009.
- National Comprehensive Cancer Network (NCCN). *Clinical Practice Guidelines in Oncology—Prostate Cancer*. 2011. Available at: [http://www.nccn.org/professionals/physician\\_gls/f\\_guidelines.asp](http://www.nccn.org/professionals/physician_gls/f_guidelines.asp).
- National Comprehensive Cancer Network (NCCN). *Clinical Practice Guidelines in Oncology: Asia Consensus Statement—Prostate Cancer*. 2011. Available at: [http://www.nccn-asia.org/pdf/2010\\_Prostate\\_Cancer.pdf](http://www.nccn-asia.org/pdf/2010_Prostate_Cancer.pdf).
- National Statistical Office. 2009. Available at: <http://kosis.nso.go.kr> (14 January 2010, date last accessed).
- Prostate Cancer Trialists’ Collaborative Group. Maximum androgen blockade in advanced prostate cancer: an overview of the randomised trials. *Lancet* 2000;355:1491–8.
- Punnen S, Cooperberg MR, Sadetsky N, Carroll PR. Androgen deprivation therapy and cardiovascular risk. *J Clin Oncol* 2011 (Epub ahead of print).
- Scher HI, Beer TM, Higano CS, Anand A, Taplin ME, Efstathiou E, et al. Antitumour activity of MDV3100 in castration-resistant prostate cancer: a phase 1–2 study. *Lancet* 2010;375:1437–46.
- Shahinian VB, Kuo YF, Freeman JL, Goodwin JS. Determinants of androgen deprivation therapy use for prostate cancer: role of the urologist. *J Natl Cancer Inst* 2006;98:839–45.
- Smith DC, Smith MR, Small EJ, Sweeney C, Kurzrock R, Gordon MS, et al. Phase II study of XL184 in a cohort of patients (pts) with castration-resistant prostate cancer (CRPC) and measurable soft tissue disease. Presented at the ASCO Genitourinary Cancers Symposium, Orlando, FL, 2011. Abstract 127.
- Tanaka N, Fujimoto K, Hirayama A, Samma S, Momose H, Kaneko Y, et al. The primary therapy chosen for patients with localized prostate cancer between the university hospital and its affiliated hospitals in Nara Uro-Oncological Research Group registration. *BMC Urol* 2011;11:6.
- The Korean Urological Oncology Society. *Clinical Guidelines on Prostate Carcinoma*. 2nd edn. 2009. Available at: <http://www.kuos.or.kr>.
- Tsai HK, D’Amico AV, Sadetsky N, Chen MH, Carroll PR. Androgen deprivation therapy for localized prostate cancer and the risk of cardiovascular mortality. *J Natl Cancer Inst* 2007;99:1516–24.

## Appendix

### MEETING PARTICIPANTS



Participants at the 5th Joint Meeting of J-CaP and CaPSURE.

Byung-Ha Chung (B.-H.C.): Professor, Department of Urology, Gangnam Severance Hospital, Yonsei University College of Medicine, Seoul, Korea. Yoshihiko Hirao (Y.H.): Professor and Chairman, Department of Urology, Nara Medical University, Kashihara City, Nara, Japan. Choung Soo Kim: Professor, Department of Urology, University of Ulsan, Asan Hospital, Seoul, Korea. Tadaichi Kitamura: President, Asoka Hospital, Tokyo, Japan. Ji Youl Lee: Professor, Department of Urology, Seoul St Mary's Hospital, The Catholic University of Korea, Seoul, Korea. Jun

Miyazaki: Associate Professor, Department of Urology, Tsukuba University, Graduate School of Comprehensive Human Sciences, Tsukuba, Japan. Masaru Murai: President, International Goodwill Hospital, Yokohama, Japan. Seiji Naito: Professor and Chairman, Department of Urology, Graduate School of Medical Sciences, Kyushu University, Fukuoka, Japan. Mikio Namiki (M.N.): Professor and Chairman, Department of Integrative Cancer Therapy and Urology, Kanazawa University Graduate School of Medical Science, Ishikawa, Japan. Kazuo Nishimura (K.N.): Director, Department of Urology, Osaka Medical Center for Cancer and Cardiovascular Diseases, Osaka, Japan. Hiroyuki Nishiyama: Professor and Chairman, Urology and Andrology, Tsukuba University, Graduate School of Comprehensive Human Sciences, Tsukuba, Japan. Mototsugu Oya: Professor and Chairman, Department of Urology, Keio University School of Medicine, Tokyo, Japan. Kazuhiro Suzuki: Professor and Chairman, Department of Urology, Gunma University Graduate School of Medicine, Gunma, Japan. Taiji Tsukamoto: Professor and Chairman, Department of Urology, Sapporo Medical University, School of Medicine, Hokkaido, Japan. Rainy Umbas (R.U.): Professor, Department of Urology, Cipto Mangunkusumo Hospital, University of Indonesia, Jakarta, Indonesia. Xianghua Zhang (X.Z.): Professor and Vice Chairman, Wujieping Urology Center, Peking University, China.