# Treatment of Localized and Locally Advanced, **High-Risk Prostate Cancer: A Report From the** First Prostate Cancer Consensus Conference for **Developing Countries**

Raja Khauli, MD1; Robson Ferrigno, MD2; Gustavo Guimarães, MD2; Muhammad Bulbulan, MD3; Pedro Luiz Serrano Uson Junior, MD4; Bernardo Salvajoli, MD5,6; Daniel Moore Freitas Palhares, MD7; Douglas Racy, MD2; Erlon Gil, MD2; Fernando Freire de Arruda, MD7; Gustavo Caserta Lemos, MD4; Gustavo Franco Carvalhal, MD8; Icaro Thiago de Carvalho, MD4; Igor Austin Fernandes Martins, MD2; Ivan Frederico Pinto Gimpel, MD9; João Victor Salvajoli, MD5,6; José Luis Chambo, MD2; José Pontes Jr, MD5; Leopoldo Alves Ribeiro Filho, MD7; Lucas Nogueira, MD10; Marcelo Roberto Pereira Freitas, MD11; Marcelo Wroclawski, MD4; Marco Antonio Arap, MD7; Marcus Vinicius Sadi, MD2; Rafael Coelho, MD5; Rafael Gadia, MD7; Rodrigo Antonio Ledezma Roja, MD12; Rodrigo de Moraes Hanriot, MD13; Ronaldo Baroni, MD4; Stenio Zequi, MD14; William Carlos Nahas, MD5; Wladimir Alfer Jr, MD4; and Fernando Cotait Maluf, MD<sup>2,4,15</sup>

PURPOSE To generate and present survey results on important issues relevant to treatment and follow-up of localized and locally advanced, high-risk prostate cancer (PCa) focusing on developing countries.

METHODS A panel of 99 PCa experts developed more than 300 survey questions of which 67 questions concern the main areas of interest of this article: treatment and follow-up of localized and locally advanced, high-risk PCa in developing countries. A larger panel of 99 international multidisciplinary cancer experts voted on these questions to create the recommendations for treatment and follow-up of localized and locally advanced, highrisk PCa in areas of limited resources discussed in this article.

**RESULTS** The panel voted publicly but anonymously on the predefined questions. Each question was deemed consensus if 75% or more of the full panel had selected a particular answer. These answers are based on panelist opinion and not on a literature review or meta-analysis. For questions that refer to an area of limited resources, the recommendations considered cost-effectiveness as well as the possible therapies with easier and greater access. Each question had five to seven relevant answers including two nonanswers. Results were tabulated in real time.

**CONCLUSION** The voting results and recommendations presented in this article can guide physicians managing localized and locally advanced, high-risk PCa in areas of limited resources. Individual clinical decision making should be supported by available data; however, as guidelines for treatment of localized and locally advanced, high-risk PCa in developing countries have not been defined, this article will serve as a point of reference when confronted with this disease.

JCO Global Oncol 7:530-537. © 2021 by American Society of Clinical Oncology

Licensed under the Creative Commons Attribution 4.0 License (c)



#### **ASSOCIATED** CONTENT

#### **Data Supplement**

Author affiliations and support information (if applicable) appear at the end of this

Accepted on February 22, 2021 and published at ascopubs.org/journal/ go on April 15, 2021: DOI https://doi.org/10. 1200/G0.20.00421

#### INTRODUCTION

High-risk prostate cancer (PCa) accounts for 15% of cancer diagnoses. This percentage may be higher in developing countries, especially where multidisciplinary care is limited to few academic medical centers and there are major access barriers from screening to confirmation diagnosis. This article will summarize the recommendations of a large panel of physicians from developing countries, specializing in PCa, regarding the treatment of patients presenting with high-risk PCa both with and without contemplating the restrictions of limited resources in the decision-making process, with the objective of providing guidance in clinical practice and policy development and modification. The

complete methodology of the Prostate Cancer Consensus Conference for Developing Countries including the elaboration process of the questionnaires to guide the panelists, the design of voting sessions, and consensus criteria were presented for the editorial and are valid for all the manuscripts (Data Supplement).

## TREATMENT: LOCALIZED AND LOCALLY ADVANCED, **HIGH-RISK PROSTATE CANCER**

For patients with life expectancy of > 10-15 years with the diagnosis of localized high-risk PCa, with Gleason score 8-10 and/or prostate-specific antigen (PSA) > 20 ng/mL, most panelists (61.15%) recommended radical prostatectomy plus lymph node dissection for





#### **CONTEXT**

#### **Key Objective**

Generate a consensus on critical issues relevant to treatment of localized and locally advanced, high-risk prostate cancer (PCa) focusing on developing countries.

### **Knowledge Generated**

Definitive treatment for high-risk PCa is achieved either surgically or with radiotherapy and androgen deprivation therapy. Androgen deprivation therapy alone is inadequate. Several factors such as life expectancy, tumor stage, prostate-specific antigen, lymph node involvement, and regional metastasis may affect the decision-making process between surgery and radiation. In this article, we developed discussions on the multiple questions voted on at the first Prostate Cancer Consensus Conference for Developing Countries for high-risk PCa.

#### Relevance

The voting results presented in this article can be used to support the treatment of localized and locally advanced, high-risk PCa in areas of limited resources lacking specific guidelines.

treatment, whereas a little more than one-quarter (27.50%) selected the combination of hormonal therapy and external beam radiation therapy (EBRT). For the same patient in an area of limited resources, panelists reached consensus (74.84%) in recommending radical prostatectomy plus lymph node dissection (where Robot Platform is not available), whereas less than one-quarter (23.46%) recommended the combination of hormonal therapy and EBRT (intensity-modulated radiation therapy [IMRT] not available).

For patients with life expectancy of > 10-15 years with the diagnosis of clinical T3/T4 and/or clinical N+, high-risk PCa, the panelists reached consensus in recommending the combination of hormonal therapy and EBRT (IMRT preferred) with or without brachytherapy (76.54%) as well as in recommending the combination of hormonal therapy and EBRT (IMRT not available) with or without brachytherapy (78.75%) for treatment in an area of limited resources.

For patients with life expectancy of < 10-15 years with the diagnosis of localized high-risk PCa, with Gleason score 8-10 and/or PSA > 20 ng/mL, the panelists almost reached consensus (71.76%) in recommending the combination of hormonal therapy and EBRT (IMRT preferred) as treatment, with some (17.65%) choosing the combination of hormonal therapy, EBRT (IMRT preferred), and brachytherapy. For the same patient in an area of limited resources, the panel reached consensus (77.38%) in recommending the combination of hormonal therapy and EBRT (IMRT not available).

The treatment for patients with life expectancy of < 10-15 years with the diagnosis of high-risk PCa with clinical T3/T4 and/or clinical N+, the panel reached consensus (79.76%) in recommending the combination of hormonal therapy and EBRT (IMRT preferred), including in an area of limited resources (85.54%) (IMRT not available).

The panel reached consensus (79.01%) in recommending the combination of hormonal therapy and conformal EBRT

for patients with the diagnosis of high-risk PCa with clinical T3/T4 and/or clinical N+ in institutions where there is no availability of IMRT technique.

In institutions where there is no availability of IMRT technique and conformal EBRT, the panel also reached consensus (82.35%) in recommending radical prostatectomy plus lymph node dissection for patients with the diagnosis of high-risk PCa with Gleason score 8-10 and/or PSA > 20 ng/mL and disease confined to the prostate.

In institutions where there is only conventional radiation therapy (RT) technique, for patients with high-risk, organ-confined PCa, the panel reached consensus (78.31%) in recommending external RT. In institutions where there is only cobalt RT technique, the panel reached strong consensus (89.95%) in recommending that patients with high-risk, disease-confined PCa cannot be treated with external RT.

In the case where some form of radiation is an option for treatment of a high-risk patient with PCa with life expectancy of > 10-15 years, the panel agreed, and in most cases reached consensus, on a recommendation for the association of hormonal treatment under varying circumstances. In the case where radical prostatectomy is an option for treatment for the same patient, the panel recommended extended lymph node dissection, in all circumstances (Tables 1 and 2).

In the case where radical prostatectomy is an option for treatment of a patient with life expectancy of > 10-15 with a diagnosis of high-risk PCa, the panel reached consensus in recommending a surgical approach under varying circumstances. The panel reached strong consensus in recommending an open radical surgery in areas of limited resources (Table 3).

In the case where exclusive hormonal therapy is an option for treatment of high-risk PCa, the panel had differing recommendations for their preferred treatment under

**TABLE 1.** Association of Hormonal Treatment to Radiation in High-Risk Patients With PCa With Life Expectancy > 10-15 years

Recommended Approach Regarding the Association of Hormonal Treatment (%)

85.00

In the Case Where Some Form of Radiation Is a Treatment Option		Mid-Term ADT (18 Months)	Long-Term ADT (24-36 Months)	
Gleason score 8-10	PSA > 20 ng/mL, organ-confined	Standard of care		80.72
and/or		Limited resources	28.92	68.67
With clinical T3/T/Lan	d/or clinical N±	Standard of care		77 78

Limited resources

Abbreviations: ADT, androgen deprivation therapy; PCa, prostate cancer; PSA, prostate-specific antigen.

varying circumstances and reached consensus in only one instance, the application of orchiectomy alone as a means of androgen deprivation therapy (ADT) for patients with clinical T3/T4 and/or clinical N+ in areas of limited resources (Table 4).

#### **RADIOTHERAPY**

Over the past few decades, radiation techniques have improved, allowing better coverage of tumor volumes with improved sparing of adjacent normal structures. <sup>2,3</sup> EBRT is the most used type of radiation treatment, in particular IMRT. The enhanced conformity of IMRT allows for dose escalation to the prostate while reducing the dose to the bladder and rectum, and trials have demonstrated reduced toxicity with IMRT; however, reimbursements for IMRT were higher, leading to increased costs in the overall care for PCa. <sup>4</sup> When comparing different radiotherapy strategies, including three-dimensional conformal radiation therapy (3DCRT) with IMRT, IMRT was found to be cost effective in one Australian study, demonstrating an approximate 1.1 million dollars in savings per 1,000 patients. <sup>5</sup>

#### ANDROGEN DEPRIVATION THERAPY

RT combined with ADT was the preferred option for highrisk patients with clinical T3/T4 and/or clinical N+ if IMRT or conformal EBRT is available. Despite IMRT being preferred over conformal RT because of reduced late toxicity, the efficacy of both techniques is comparable; therefore, the latter is a reasonable option in the treatment of localized and locally advanced disease, particularly in areas of limited resources where IMRT is not available. A study evaluating two prospective cohorts of men treated for localized PCa investigated the hypothesis of reductions in toxicity and showed that the 5-year cumulative incidence of grade  $\geq 2$  GI toxicity was 24.9% for IGRT and IMRT and 37.6% with 3DCRT (P = .005), with significant reductions in proctitis (P = .047) and increased stool frequency (P < .001). On the other hand, genitourinary grade  $\geq 2$  toxicity levels at 5 years were comparable.<sup>6</sup>

The panel favored radical prostatectomy over radiation when neither IMRT nor conformal radiation was available. Cobalt RT in this consensus was defined as using the external beam from a <sup>60</sup>Cobalt unit, with parallel opposed, three-field or four-field box technique, with anatomic bony landmarks on plain radiographs being used to define shape and location of the prostate. This technique presents target inaccuracies, higher volumes of irradiation to normal tissue, and lower energy used (average beam energy of 1.25 MeV) when compared with modern linear accelerators. Consequently, the panel considered this treatment to have inferior efficacy because of the lower doses toward the tumor and pelvic drainage as well more toxicity.<sup>7</sup>

ADT remains a cornerstone in the treatment of high-risk localized and advanced PCa. In localized and locally advanced disease, ADT is typically combined with RT. The duration of ADT is usually 2-3 years, based on clinical trials that established that long-term ADT (28-36 months) is more effective than short-term ADT (4-6 months). On the other hand, ADT alone is inferior to ADT combined with radiation in high-risk and/or locally advanced PCa.<sup>8</sup>

Recommended Extension of Lymph Node

TABLE 2. Lymph Node Dissection in Addition to Radical Prostatectomy in High-Risk Patients With PCa With Life Expectancy > 10-15 Years

Treatment Option			Dissection (%)
High-risk PCa with Gleason score 8-10 and/or	PSA > 20 ng/mL, organ- confined	Standard of care	Extended pelvic lymph node dissection 100
		Limited resources	Extended pelvic lymph node dissection 100
High-risk PCa with clinical T3/T4 and/or clinical N+		Standard of care	Extended pelvic lymph node dissection 98.78
		Limited resources	Extended pelvic lymph node dissection 97.53

Abbreviations: PCa, prostate cancer; PSA, prostate-specific antigen.

In the Case Where Radical Prostatectomy Is a

TABLE 3. Recommended Surgical Approach for Treatment With Radical Prostatectomy in High-Risk Patients With PCa With Life Expectancy > 10-15 Years

Recommended Surgical Approach (%)

In the Case Where Radical Prostatectomy Is a Treatment Option	Open Surgery	Robotic Surgery	
Standard of care			
With Gleason score 8-10 and/or PSA > 20 ng/mL, organ-confined		84.34	
With clinical T3/T4 and/or clinical N+		79.01	
Area of limited resources			
With Gleason score 8-10 and/or PSA > 20 ng/mL, organ confined	93.83		
With clinical T3/T4 and/or clinical N+	93.83		

Abbreviations: PCa, prostate cancer; PSA, prostate-specific antigen.

ADT alone should be considered in patients with a contraindication to RT or in areas of limited resources with no access to RT. Bilateral orchiectomy is a surgical option for metastatic disease and for selected elderly patients with locally advanced disease. It is well known that orchiectomy is less expensive than any other ADT therapy. It is estimated that the cost of luteinizing hormone-releasing hormone (LHRH) agonist treatment for androgen suppression is 10-13 times and that for combined androgen blockade is 17-20 times higher than the cost of bilateral orchiectomy. 10

### RADICAL PROSTATECTOMY

Radical prostatectomy is a second treatment option for patients with high-risk PCa, as noted in the NCCN, American Urological Association, and European Association of Urology guidelines. <sup>11,12</sup> In addition, in this setting, pelvic lymph node dissection is considered the surgical standard for PCa staging. There are no randomized clinical trials completed comparing surgery versus radiation. Therefore, there is no consensus regarding what the best approach

is focusing on efficacy, toxicity, and quality of life. A retrospective German analysis including 910 men with localized disease treated with surgery, 292 with radiation, and 124 with active surveillance concluded that surgery was associated with more life-years gained. However, because of the high inpatient costs of the initial surgery, radical prostatectomy had €11,000 in Euros higher total per capita costs than radiation or active surveillance.¹³ The length of RT compared with surgery may also present an issue in patients with limited transportation and/or access to the health centers.

## ADJUVANT TREATMENT AFTER RADICAL PROSTATECTOMY: LOCALIZED AND LOCALLY ADVANCED, HIGH-RISK PROSTATE CANCER

When considering patients with life expectancy of < 10-15 years or > 10-15 years with adverse factors post-prostatectomy (positive margins and/or extracapsular disease and/or seminal vesicle involvement) but no pathologic lymph node involvement (pNO) and undetectable post-operative PSA, the panel differed in their recommendations for adjuvant radiation and hormonal therapy, including in

TABLE 4. Preferred Hormonal Therapy for Treatment of High-Risk PCa

In the Case Where Exclusive Hormonal Therapy Is a Treatment Option	Preferred Treatment	% of Votes
Standard of care		
With Gleason score 8-10 and/or PSA $>$ 20 ng/mL, disease-confined	ADT by LHRH agonist alone (with or without first-generation AR antagonist)	51.22
	ADT by LHRH antagonist alone (with or without first-generation AR antagonist)	21.95
	Any form of intermittent ADT	17.07
With clinical T3/T4 and/or clinical N+	ADT by LHRH agonist alone (with or without first-generation AR antagonist)	65.43
	ADT plus abiraterone	18.52
Area of limited resources		
With Gleason score 8-10 and/or PSA > 20 ng/mL, disease-	ADT by orchiectomy alone	63.75
confined	ADT by LHRH agonist alone (with or without first-generation AR antagonist)	18.75
With clinical T3/T4 and/or clinical N+	ADT by orchiectomy alone	81.93

Abbreviations: ADT, androgen deprivation therapy; AR, androgen receptor; LHRH, luteinizing hormone-releasing hormone; PCa, prostate cancer; PSA, prostate-specific antigen.

areas of limited resources. Consensus was not reached on any recommendation for these patients (Table 5).

In line with recent recommendations on the basis of randomized controlled trials, when considering hormonal therapy as an option as part of adjuvant therapy in association with RT because of positive margins and/or extracapsular disease and/or seminal vesicle involvement, although not reaching consensus, most of the panel (72.29%) preferred ADT by LHRH agonist alone (with or without first-generation androgen receptor [AR] antagonist), whereas less than one-fourth (21.69%) recommended no ADT. If adjuvant hormonal therapy alone is recommended for a patient with positive margins and/or extracapsular disease and/or seminal vesicle involvement, slightly more than half (51.76%) of the panel voted to abstain from the answer. Some of the panel members (21.18%) selected long-term ADT (24-36 months) and others (18.82%) chose short-term ADT (6 months).

For the same patient, in an area of limited resources, the panel was almost equally divided in their preference, with one-third (34.18%) recommending ADT by LHRH agonist alone (with or without first-generation AR antagonist), another third (34.18%) indicating no ADT would be recommended, and just less than one-quarter (22.78%) abstaining. If the panelists recommend adjuvant hormonal therapy alone for this patient, as to the preferred duration of hormonal therapy, most panelists (68.67%) abstained, whereas few selected short-term ADT (6 months) and long-term ADT (24-36 months) in combination with radiation.

In institutions where only conventional conformal EBRT is available (no IMRT), the panelists were divided in recommending adjuvant radiation therapy (ART) in men with life expectancy of > 10-15 years who recovered urinary continence, with adverse factors postprostatectomy (positive margins and/or extracapsular disease and/or seminal vesicle involvement but no pathologic lymph

node involvement [pN0]) and undetectable postoperative PSA. More than half (48.10%) recommended ART in the majority of the cases. For the same patient, in institutions where only cobalt RT is available, the panel reached consensus (93.83%) in not recommending ART.

For men with life expectancy of > 10-15 years who recovered urinary continence with pathologic lymph node involvement (pN+) and undetectable postoperative PSA, the majority of the panel (54.55%) recommended ART in the majority of cases, whereas almost one-third (31.17%) recommended it only in a few cases.

For men with pathologic lymph node involvement (pN+) and undetectable postoperative PSA, most of the panelists (68.35%) recommended hormonal therapy as part of adjuvant therapy associated with adjuvant radiation in the majority of cases, some (16.48%) did not recommend it, and others (15.19%) recommended it only in a few cases. This poses a trend toward a consensus in that the majority of the panelists (84.83%) recommended hormonal therapy for either all or some cases.

For men with pathologic lymph node involvement (pN+) and undetectable postoperative PSA, most panelists (50.62%) do not recommend isolated hormonal therapy as adjuvant therapy, whereas almost one-third (31.10%) recommend it only in a few cases.

When hormonal therapy is an option as part of adjuvant therapy in association with RT because of pathologic lymph node involvement (pN+), the panelists reached consensus (90.12%) in preferring ADT by LHRH agonist alone (with or without first-generation AR antagonist).

If the panelists recommended adjuvant hormonal therapy because of pathologic lymph node involvement (pN+), most (70.51%) selected long-term ADT, of which 55.13% would give it for 24-36 months and 15.38% recommended ADT indefinitely; 17.95% abstained.

**TABLE 5.** Adjuvant Radiation and Hormonal Therapy for Patients With Adverse Factor Postprostatectomy Defined by Positive Margins and/or Extracapsular Disease and/or Seminal Vesicle Involvement but No Pathologic Lymph Node Involvement (pN0) and Undetectable Postoperative PSA

	Recommendation for ART (%)			Recommendation for Hormonal Therapy As Part of Adjuvant Therapy Associated With Adjuvant Radiation (%)		
Patient Characteristic	Yes, in the majority of cases	Yes, only in a few cases	No	Yes, in the majority of cases	Yes, only in a few cases	No
Life expectancy < 10-15 years						
Who recovered urinary continence	28.75	28.75	42.50			
In an area of limited resources				17.07	18.29	64.63
Life expectancy > 10-15 years						
Who recovered urinary continence	48.10	34.18			34.62ª	47.44
In an area of limited resources				16.25		68.75

Abbreviations: ART, adjuvant radiation therapy; PSA, prostate-specific antigen.

<sup>&</sup>lt;sup>a</sup>In the case of two or more adverse factors.

In areas of limited resources, the panelists' opinions differed for recommendations of ART, hormonal therapy as part of adjuvant therapy associated with adjuvant radiation, and isolated hormonal therapy as adjuvant therapy for men under varying circumstances, as noted in Table 6.

In institutions where there is only conventional conformal EBRT technique (no IMRT), for men with life expectancy of > 10-15 years who recovered urinary continence, with pathologic lymph node involvement (pN+) and undetectable postoperative PSA, half the panel (50.52%) recommend ART in the majority of cases, whereas one-quarter (26.25%) recommend it in a few cases and one-fifth (20.21%) do not recommend it. For the same patient, in institutions where there is only cobalt RT technique, the panel reached consensus and 95% of the voters would not recommend ART.

In this situation, slightly more than half of the panelists (54.32%) recommended hormonal therapy as part of adjuvant therapy associated with radiation, either conformal or cobalt (no IMRT), for men with pathologic lymph node involvement (pN+) and undetectable postoperative PSA in the majority of cases, whereas almost one-quarter (23.26%) did not.

The panelists were divided when asked about the extension of the RT field in case of ART in men with high-risk PCa (positive margins and/or extracapsular extension and/or seminal vesicle involvement) and pN0 postprostatectomy. Almost half of the panel (45.12%) recommended prostatic bed only and the same percentage (45.12%) recommended prostatic bed plus whole pelvis.

In institutions where there is only conventional conformal EBRT available (no IMRT) and ART is recommended for men with high-risk PCa (positive margins and/or extracapsular extension and/or seminal vesicle involvement) and

pNO postprostatectomy, most panelists (62.96%) recommended to irradiate the prostatic bed only. For the same patient and circumstance, in institutions where there is only cobalt RT technique, the panelists reached consensus (93.83%) in not recommending RT. This recommendation contrasts with recent literature recommending whole-pelvis radiation in addition to prostatic fossa radiation.

When treating with ART in men with pathologic lymph node involvement (pN+) and undetectable postoperative PSA, the panel reached consensus (89.02%) in recommending prostatic bed and whole pelvis as the RT volumes. In institutions where there is only conventional conformal EBRT technique (no IMRT) and ART is recommended in men with pathologic lymph node involvement (pN+) and undetectable postoperative PSA, the panel reached consensus (85.71%) in recommending RT to the prostatic bed plus whole pelvis. For the same patient and circumstance, in institutions where there is only cobalt RT technique, the panel reached almost complete consensus (98.73%) in not recommending RT in this scenario.

For patients with high-risk PCa, most panelists (60%) recommended external iliac, internal iliac, obturator, and common iliac as the pelvic lymph node dissection template, although some (22.35%) recommended external iliac, internal iliac, obturator, common iliac, and presacral. For the same patient in an area of limited resources, the panel made a similar recommendation with most panelists (62.03%) recommending external iliac, internal iliac, obturator, and common iliac.

ART for patients with higher risk of relapse after radical prostatectomy has been studied in several randomized trials with questionable results regarding an overall survival benefit of this strategy. <sup>14-16</sup> One of the most important questions in this space refers to the optimal timing of the

**TABLE 6.** Hormonal Therapy as Part of Adjuvant Therapy Associated With Adjuvant Radiation and Isolated Hormonal Therapy As Adjuvant Therapy for Men With High-Risk Prostate Cancer in Areas of Limited Resources

In Areas of Limited Resources	Recommendation of ART	Recommendation of Hormonal Therapy As Part of Adjuvant Therapy Associated With Adjuvant Radiation	Recommendation of Isolated Hormonal Therapy As Adjuvant Therapy
In men with life expectancy > 10-15 years who recovered urinary continence with pathologic lymph node involvement (pN+) and undetectable postoperative PSA	No (42.50%) Yes, in the majority of cases (35.00%) Yes, but only in a few cases (22.50%)		
For men with pathologic lymph node involvement (pN+) and undetectable postoperative PSA		Yes, in the majority of cases (63.38%) No (23.08%)	No (51.28%) Yes, in the majority of cases (32.05%)
In case the option for hormonal therapy is made as part of adjuvant therapy in association with RT because of pathologic lymph node involvement (pN+)	Long-term ADT (24-36 months; 50.52%) Short-term ADT (6 months; 18.75%)	ADT by LHRH agonist alone (with or without first-generation AR antagonist; 70.73%) ADT by orchiectomy alone (14.63%)	

Abbreviations: ADT, androgen deprivation therapy; AR, androgen receptor; ART, adjuvant radiation therapy; LHRH, luteinizing hormone-releasing hormone; PSA, prostate-specific antigen; RT, radiation therapy.

postoperative RT. To answer this question, one randomized trial (RADICALS-RT) and one meta-analysis did not show that ART improved event-free survival when compared with early salvage radiation.<sup>17</sup>

Although the use of ADT concurrent with ART was selected by many panelists as a treatment for patients with high risk of recurrence after radical prostatectomy, this approach is still controversial. Until recently, there were no phase III trials favoring the use of concurrent ADT in the adjuvant setting.

## FOLLOW-UP: LOCALIZED AND LOCALLY ADVANCED, HIGH-RISK PROSTATE CANCER

In a patient with high-risk and/or locally advanced PCa, who underwent surgery with a curative intent, most of the panelists (60.76%) recommended following the majority of patients by anamnesis, physical examination including digital rectal examination, and PSA every 3-6 months for 5 years then every year. For the same patient, in an area of limited resources, the panel reached consensus (77.22%) in recommending follow-up by anamnesis, physical examination including digital rectal examination, and PSA every 3-6 months for 5 years then every year.

After RT (any form) with curative intent (with or without ADT) in a patient with high-risk and/or locally advanced PCa, the panel reached consensus (76.62%) in recommending following the majority of patients by anamnesis, physical examination including digital rectal examination, and PSA every 3-6 months for 5 years then every year. Consensus (82.05%) was also reached to follow up patients the same way in areas of limited resources (Data Supplement).

After definitive therapy with curative intent for high-risk PCa, most of the panel (64.02%) indicated they would never order imaging as follow-up for the majority of patients. For the same patient, in an area of limited resources, the panel reached consensus (85%) indicating that they would not order imaging at follow-up.

In conclusion, definitive treatment of the primary tumor for high-risk PCa is achieved either surgically or with RT and ADT. ADT alone is inadequate. Several factors such as life expectancy, tumor stage, lymph node involvement, and regional metastasis may affect the decision-making process between surgery and radiation. In scarce-resource settings, given the low availability of IMRT, 3DCRT, or image-guided radiation therapy combination of hormonal therapy and EBRT if conformal RT is available, particularly conventional conformal external beam, as the minimum accepted technique or radical prostatectomy in expert hands may be the best available option for the treatment and management of these patients. Nonetheless, even in areas of limited resources, when the only available RT is a cobalt machine, this treatment method is not recommended, and alternate surgical or medical options should be sought to manage the high-risk patient with disease confined or not confined to the prostate. Regarding longterm ADT, it was determined that orchiectomy may be an alternative for areas of resource limitations. Radical prostatectomy could translate into a more cost-effective strategy in areas of scarce resources, with limited access to RT. Open surgery is deemed to be adequate.

#### **AFFILIATIONS**

- <sup>1</sup>Naef K. Basile Cancer Institute, Beirut, Lebanon
- <sup>2</sup>Hospital BP, São Paulo, Brazil
- <sup>3</sup>American University of Beirut, Beirut, Lebanon
- <sup>4</sup>Hospital Israelita Albert Einstein, São Paulo, Brazil
- <sup>5</sup>Instituto do Câncer de São Paulo, São Paulo, Brazil
- <sup>6</sup>Hospital do Coração, São Paulo, Brazil
- <sup>7</sup>Hospital Sírio Libanês, São Paulo, Brazil
- <sup>8</sup>Hospital Moinho de Ventos, Porto Alegre, Brazil
- 9Clínica Santa María, Providencia, Chile
- <sup>10</sup>Universidade Federal de Minas Gerais, Belo Horizonte, Brazil
- <sup>11</sup>Centro de Pesquisas Oncológicas de Santa Catarina, Santa Catarina, Brazil
- <sup>12</sup>Universidad de Chile, Santiago, Chile
- <sup>13</sup>Hospital Alemão Oswaldo Cruz, São Paulo, Brazil
- <sup>14</sup>Hospital do Câncer AC Camargo, São Paulo, Brazil
- <sup>15</sup>Latin American Oncology Group (LACOG), Porto Alegre, Brazil

## **CORRESPONDING AUTHOR**

Fernando Cotait Maluf, MD, Avenida Albert Einstein 626/627, Jardim Leonor São Paulo, São Paulo, Brazil 05652-900; e-mail: maluffc@uol.com.br.

## **AUTHOR CONTRIBUTIONS**

Conception and design: Raja Khauli, Muhammad Bulbulan, Bernardo Salvajoli, Douglas Racy, Gustavo Caserta Lemos, Gustavo Franco

Carvalhal, Igor Austin Fernandes Martins, João Victor Salvajoli, José Luis Chambo, Rafael Gadia, William Carlos Nahas, Fernando Cotait Maluf **Administrative support**: Raja Khauli, Marcelo Wroclawski, William Carlos Nahas, Fernando Cotait Maluf

Provision of study materials or patients: Gustavo Guimarães, Muhammad Bulbulan, Douglas Racy, Fernando Freire de Arruda, Gustavo Caserta Lemos, João Victor Salvajoli

Collection and assembly of data: Robson Ferrigno, Pedro Luiz Serrano Uson Junior, Bernardo Salvajoli, Daniel Moore Freitas Palhares, Erlon Gil, Fernando Freire de Arruda, Igor Austin Fernandes Martins, José Luis Chambo, Leopoldo Alves Ribeiro Filho, Lucas Nogueira, Marcelo Roberto Pereira Freitas, Marco Antonio Arap, Marcus Vinicius Sadi, Rafael Coelho, Rodrigo Antonio Ledezma Roja, Rodrigo de Moraes Hanriot, Ronaldo Baroni, Stenio Zequi, Fernando Cotait Maluf

Data analysis and interpretation: Raja Khauli, Robson Ferrigno, Gustavo Guimarães, Bernardo Salvajoli, Icaro Thiago de Carvalho, Igor Austin Fernandes Martins, Ivan Frederico Pinto Gimpel, João Victor Salvajoli, José Luis Chambo, José Pontes Jr, Marcelo Roberto Pereira Freitas, Marcelo Wroclawski, Marcus Vinicius Sadi, Ronaldo Baroni, Stenio Zequi, Wladimir Alfer Jr, Fernando Cotait Maluf

Manuscript writing: All authors
Final approval of manuscript: All authors

Accountable for all aspects of the work: All authors

## AUTHORS' DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST

The following represents disclosure information provided by authors of this manuscript. All relationships are considered compensated unless otherwise noted. Relationships are self-held unless noted. I = Immediate Family Member, Inst = My Institution. Relationships may not relate to the subject matter of this manuscript. For more information about ASCO's conflict of interest policy, please refer to www.asco.org/rwc or ascopubs. org/go/authors/author-center.

Open Payments is a public database containing information reported by companies about payments made to US-licensed physicians (Open Payments).

#### Raja Khauli

Honoraria: Astellas Pharma, Janssen, Algorithm, Hikma, Lilly Consulting or Advisory Role: Astellas Pharma, Hikma

**Speakers' Bureau:** Algorithm, Lilly **Research Funding:** Astellas Pharma

Leopoldo Alves Ribeiro Filho Consulting or Advisory Role: Ferring Speakers' Bureau: Zodiac Pharma

Lucas Nogueira

Consulting or Advisory Role: Bayer Speakers' Bureau: Bayer, Astellas Pharma

Marcelo Roberto Pereira Freitas

Consulting or Advisory Role: Janssen Oncology, Zodiac Pharma

Speakers' Bureau: Astellas Pharma, Janssen Oncology

Rodrigo Antonio Ledezma Roja

Honoraria: Ferring, Quanta Systems

Consulting or Advisory Role: Quanta Systems, Ferring Travel, Accommodations, Expenses: Quanta Systems

Rodrigo de Moraes Hanriot

Speakers' Bureau: AstraZeneca, Varian Medical Systems, Bristol Myers

Squibb, Takeda, Roche

Stenio Zequi

Consulting or Advisory Role: Pfizer, Astellas Brazil

Speakers' Bureau: Pfizer, Astellas Pharma, Bayer, Janssen, Astra Zeneca

Brazil

Fernando Cotait Maluf

**Honoraria:** Janssen Oncology, Astellas Pharma, Bayer Schering Pharma, Merck Sharp & Dohme, Roche/Genentech, BMS Brazil, Ipsen, Sanofi/

Aventis, Merck Serono, AstraZeneca

Consulting or Advisory Role: Janssen Oncology, Astellas Pharma, Bayer Schering Pharma, Merck Sharp & Dohme, Roche/Genentech, BMS

Brazil, Ipsen, Merck Serono, AstraZeneca

Research Funding: Janssen-Cilag, Merck Sharp & Dohme Travel, Accommodations, Expenses: Merck Sharp & Dohme, Bayer

Schering Pharma

No other potential conflicts of interest were reported.

#### **REFERENCES**

- 1. Chang A, Autio K, Roach M, et al: "High-risk" prostate cancer: Classification and therapy. Nat Rev Clin Oncol 11:308-323, 2014
- 2. Gay HA, Michalski JM: Radiation therapy for prostate cancer. Mo Med 115:146-150, 2018
- 3. Worthington JF: Treatment for prostate cancer: External-beam radiation therapy. https://www.pcf.org/c/treatment-for-prostate-cancer-external-beam-radiation-therapy/
- 4. Fischer-Valuck BW, Rao YJ, MichalskiJM, et al: Intensity-modulated radiotherapy for prostate cancer. Transl Androl Urol 7:297-307, 2018
- 5. Konski A: Cost effectiveness of prostate cancer radiotherapy. Transl Androl Urol 7:371-377, 2018
- 6. Wortel R, Incrocci L, Pos F, et al: Late side effects after image guided intensity modulated radiation therapy compared to 3D-conformal radiation therapy for prostate cancer: Results from 2 prospective cohorts. Int J Rad Oncol Biol Phys 95:680-689, 2016
- 7. Ward M, Tendulkar R, Ciezki J, et al: Future directions from past experience: A century of prostate radiotherapy. Clin Genit Cancer 12:13-20, 2014
- 8. Mottet N, Bellmunt J, Briers E, et al: EAU—ESTRO—ESUR—SIOG guidelines on prostate cancer. https://uroweb.org/guideline/prostate-cancer/
- 9. Francesco P, Stefano D, Riccardo B, et al: Robot-assisted extended pelvic lymph nodes dissection for prostate cancer: Personal surgical technique and outcomes. Int Braz J Urol 41:1209-1219, 2015
- 10. Brandes A, Koerber F, Schwarzkopf L, et al: Cost-effectiveness of radical prostatectomy, radiation therapy and active surveillance for the treatment of localized prostate cancer—A claims data analysis. Value in Health 17:A636-A637, 2014
- Leow J, Chang S, Meyer C, et al: Robot-assisted versus open radical prostatectomy: A contemporary analysis of an all-payer discharge database. Eur Urol 70:837-845, 2016
- Coughlin G, Yaxley J, Chambers S, et al: Robot-assisted laparoscopic prostatectomy versus open radical retropubic prostatectomy: 24-Month outcomes from a randomised controlled study. Lancet Oncol 19:1051-1060, 2018
- Mottet N, Peneau M, Mazeron J, et al: Addition of radiotherapy to long-term androgen deprivation in locally advanced prostate cancer: An open randomised phase 3 trial. Eur Urol 62:213-219, 2012
- 14. Wiegel T, Bartkowiak D, Bottke D, et al: Adjuvant radiotherapy versus wait-and-see after radical prostatectomy: 10-Year follow-up of the ARO 96–02/AUO AP 09/95 trial. Eur Urol 66:243-250, 2014
- 15. Bolla M, van Poppel H, Tombal B, et al: Postoperative radiotherapy after radical prostatectomy for high-risk prostate cancer: Long-term results of a randomised controlled trial (EORTC trial 22911). Lancet 380:2018-2027, 2012
- 16. Thompson I, Tangen C, Paradelo J, et al: Adjuvant radiotherapy for pathological T3N0M0 prostate cancer significantly reduces risk of metastases and improves survival: Long-term followup of a randomized clinical trial. J Urol 181:956-962, 2009
- 17. Urotoday.com: ESMO 2019: Timing of radiotherapy after radical prostatectomy: First results from the RADICALS radiotherapy randomized controlled trial. 2020. https://www.urotoday.com/conference-highlights/esmo-2019/esmo-2019-prostate-cancer/115207-esmo-2019-timing-of-radiotherapy-after-radical-prostatectomy-first-results-from-the-radicals-radiotherapy-randomized-controlled-trial.html