



ORIGINAL ARTICLE

Active surveillance in low-risk prostate cancer. Patient acceptance and results[☆]

V. Hernández^{a,*}, C. Blázquez^a, E. de la Peña^a, E. Pérez-Fernández^b, F.J. Díaz^a, C. Llorente^a

^a Servicio de Urología, Hospital Universitario Fundación de Alcorcón, Madrid, Spain

^b Unidad de Investigación, Hospital Universitario Fundación de Alcorcón, Madrid, Spain

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KEYWORDS

Prostate cancer;
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Abstract

Objectives: To evaluate the acceptance of active monitoring by patients treated in our health-care community and to report the clinical results of an active surveillance program in patients with low-risk prostate cancer.

Materials and methods: Prospective study of patients enrolled in an active surveillance program at our center between 2004 and 2012. The inclusion criteria were PSA < 10 ng/ml, Gleason score ≤ 6, clinical stage T1c/T2a, ≤ 2 positive cores, and no more than 50% of the core being affected. Curative treatment was proposed when faced with pathological progression over the course of the monitoring.

Results: In 2011, only 17% of the total number of potential candidate patients rejected their inclusion in a surveillance program and were treated actively. We analyzed a series of 144 patients included in our active surveillance protocol. The mean follow-up time was 3.22 years (SD 2.08). A total of 110 patients (76.3%) remained under active monitoring, with an estimated median treatment-free survival after diagnosis of 6.9 years (95% CI: 6.2–7.6). The percentage of patients who remained free of treatment at 2 and 5 years was 96.3% (95% CI: 92.8–99.8%) and 70.9% (95% CI: 59.3–85.5%), respectively. Thirty-four patients (23.6%) required curative treatment. The mean time to treatment was 4.6 years (SD 2.3).

Conclusions: Active surveillance of highly selected patients with low-risk prostate cancer is a valid alternative therapy that is accepted by patients in our community.

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* Corresponding author.

E-mail address: vhernandez@fhalcorcon.es (V. Hernández).

PALABRAS CLAVE

Cáncer de próstata;
Vigilancia activa;
Resultados del
tratamiento

Vigilancia activa en cáncer de próstata de bajo riesgo. Aceptación por el paciente y resultados

Resumen

Objetivos: Evaluar la aceptación del seguimiento activo por los pacientes en nuestro entorno asistencial y describir los resultados clínicos de un programa de vigilancia activa en pacientes con cáncer de próstata de bajo riesgo.

Material y métodos: Estudio prospectivo de pacientes incluidos en programa de vigilancia activa en nuestro centro entre 2004 y 2012. Los criterios de inclusión fueron: PSA < 10 ng/ml, Gleason \leq 6, estadio clínico T1c/T2a, \leq 2 cilindros positivos, con una afectación máxima del cilindro del 50%. Se propuso tratamiento curativo ante la progresión anatomopatológica a lo largo del seguimiento.

Resultados: En el año 2011, del total de pacientes potenciales candidatos, tan solo un 17% de los mismos rechazó la inclusión en un programa de vigilancia y fue tratado de forma activa. Analizamos una serie de 144 pacientes incluidos en nuestro protocolo de vigilancia activa. La media de seguimiento fue de 3,22 años (DE: 2,08). Ciento diez pacientes (76,3%) permanecen en seguimiento activo, con una mediana estimada de supervivencia libre de tratamiento tras el diagnóstico de 6,9 años (IC 95%: 6,2-7,6%). El porcentaje de pacientes que permanecen libres de tratamiento a 2 y 5 años fue de 96,3% (IC 95%: 92,8-99,8%) y 70,9% (IC 95%: 59,3-85,5%) respectivamente. Treinta y cuatro pacientes (23,6%) precisaron tratamiento curativo. La media de tiempo hasta el tratamiento fue de 4,6 años (DE: 2,3).

Conclusiones: La vigilancia activa en pacientes altamente seleccionados con cáncer de próstata de bajo riesgo es una alternativa terapéutica válida de tratamiento y aceptada por los pacientes de nuestro entorno.

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Introduction

In the last years, active surveillance is a well-consolidated option in urological practice for management of low-risk prostate cancer, and it has been included as a treatment modality in clinical guidelines of the European and American urological associations and NCCN.¹⁻³ However, there is still no consensus on the criteria for inclusion in these programs, optimal monitoring protocol and indication criteria for active treatment. Moreover, in our health-care community it is considered as treatment option not well accepted by patients and, for this reason, difficult to apply in clinical practice (Fig. 1).

Objective

The objective of our study is to evaluate the acceptance of active surveillance by patients treated in our health-care community and to report the clinical results of an active surveillance program in patients with low-risk prostate cancer, and to assess the dwell time on this program, trying to increase the scientific evidence published to date.

Materials and methods

Prospective study of patients enrolled in an active surveillance program at our center between 2004 and 2012 were conducted.

Inclusion criteria

Active surveillance was proposed to those patients diagnosed of low-risk prostate adenocarcinoma as therapeutic option. The inclusion criteria were: PSA < 10 ng/ml, Gleason score < 6, clinical stage T1c/T2a, \leq 2 positive prostate biopsy cylinders with no more than 50% of the core being affected. Curative treatment was proposed when faced with pathological progression over the course of the surveillance, both in

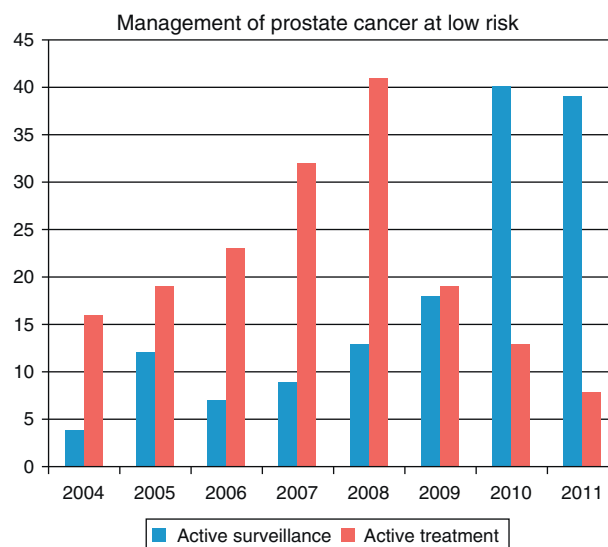


Figure 1 Evolution in the management of low-risk prostate cancer.

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