

NIVOLUMAB: CLINICAL EXPERIENCE IN A TERTIARY HOSPITAL

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Background

4CPS-099

Nivolumab is approved in Spain for the treatment of patients with melanoma, metastatic non-small cell lung cancer (NSCLC) and renal cell carcinoma (RCC), and has been included in our hospital's formulary since 2016.

Purpose

To evaluate the effectiveness and safety of patients treated with nivolumab in our hospital.

Material and methods

To evaluate the effectiveness and safety of patients treated with nivolumab in our hospital. Retrospective observational study of all patients treated with nivolumab **from February 2016 to June 2017**.

Data collected Clinical history

Age	ECOG	Prior lines treatment	% death
Sex	Treatment duration	PFS and OS (Kaplan- Meier)	% adverse effects
Diagmosis	Number of cycles	% patients continuing treatment	

Results





64,4 (39-99) years

Median age

	Melanoma	NSCLC	RCC
Nº patients	20	20	1
Median disease stage	4	4	4
Median ECOG	1	1	0
Mean treatment duration (days)	118	129	70
Median nº (cycles)	8,4	9,2	5
Line: nº patients	First line: 12 Second-third line: 8	Second-third line: 20	Second line: 1

	Melanoma	NSCLC	RCC
(%) patients continuing treatment	20	35	0
(%) deaths	40	45	0
Median PFS (95% CI) (days)	74 (38-86)	76 (41-87)	No progression
Median OS (95% CI) (days)	96 (45-120)	99 (67-126)	No death

63,4% patients → Toxicity grade I-II

- 24% asthenia
- 14,6% pruritus and dermathologycal reactions
- 9,7% artralgia or myalgia
- 15,1% others

No patient required hospitalisation, however, 1 patient discontinued treatment for renal toxicity. Grade 3–4 reactions were not detected.

Conclusions

The effectiveness in terms of PFS and OS was more reduced than clinical essays, although we should consider that there were patients with ECOG ≥2. In most cases, nivolumab was safe and well tolerated.

To evaluate efficacy and long term safety, a longer monitoring period is required.