Stem cell and renal transplantation in patients with plasma cell dyscrasias and end-stage renal disease

Beatriz Redondo, MD¹, Candela Moliz, MD¹, Maria Molina, MD¹, Enrique Morales, PhD¹, Esther Gonzalez, PhD¹, Jimena Cabrera, MD² and Amado Andres, PhD¹.

¹Nephrology Department, Hospital Universitario 12 de Octubre, Madrid, Madrid, Spain and ²Nephrology Department, Hospital Evangélico y Hospital Militar, Montevideo, Uruguay.



Introduction

- Plasma cell dyscrasias (PCD) are one of the etiologies of end-stage renal disease (ESRD).
- Traditionally, renal transplantation (RT) has been avoided in these patients due to the poor patient survival, the risk of recurrence after





RT and the high incidence of life-threatening infective complications.

However, the good results of stem cell transplantation (SCT) combined with the new drug therapies in PCD patients with ESRD have encouraged their inclusion as KT candidates.

Aims

 To describe the results of our experience of combined therapy with SCT and RT in patients with PCD and ESRD.

Methods

TABLE 2. Results during and at the end of follow-up.

Median follow-up (mo)	36
Median serum creatinine (mg/dl) after 1 year	1.6 [1.1-1.9]
Median serum creatinina (mg/dl) after 3 years	1.3 [1.1-1.9]
Acute rejection rate	1 case (16.7%) 10 months past RT
Overall renal graft survival (%)	83%
Recurrence of PCD after KT (%)	50%
	3 patients who developed 5 infection cases who required in-patient

- We performed a retrospective study that included all patients with PCD who had received both SCT and RT in our center.
- We reviewed renal, haematological, rate of infection evolution and recipient survival after 3 years of follow-up.

Severe infection rate (no)

management 2 invasive aspergillosis 2 viral infections 1 urinary infection.

Results

 TABLE 1. Demographic and clinical features of our cohort

Mean age Years (range)	55 (49-57)
Males n (%)	4 (67%)
ESRD etiology	2 myeloma cast nephropathy (33%) 2 light-chain deposition disease (33%) 1 primary amiloidosis (16.7%) 1 FSGS (16.7%)
Type of PCD	5 multiple myeloma (83%) 1 primary amiloydosis
Timing of SCT	4 previous to RT (67%) 2 after RT (33%)
Induction therapy in RT	4 basiliximab 1 timoglobuline

Sequential SCT and RT could be a suitable option for patients with PCD and ESRD.

Conclusions

- Both patient and renal graft survival are conditioned to the relapse of hematological disease and infective complications.
- The high incidence of fungal infection will require special prophylaxis measure.
- These results highlight the importance of declaring more number of patients in this situation and with longer follow-up to elucidate the best management of PCD with ESRD.

No relevant financial with respect to my research

amado.andres@salud.madrid.org





Renal transplantation - Treatment & immunosuppre

Beatriz Redondo

DOI: 10.3252/pso.eu.54ERA.2017



