

Review

Hematopoietic Cell and Renal Transplantation in Plasma Cell Dyscrasia Patients

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Gammopathies, multiple myeloma, and amyloidosis are plasma dyscrasias characterized by clonal proliferation and immunoglobulin overproduction. Renal impairment is the most common and serious complication with an incidence of 20–30% patients at the diagnosis. Kidney transplant has not been considered feasible in the presence of plasma dyscrasias because the immunosuppressive therapy may increase the risk of neoplasia progression, and paraproteins may affect the graft. However, recent advances in clinical management of multiple myeloma and other gammopathies allow considering kidney transplant as a possible alternative to dialysis. Numerous evidence indicates the direct relationship between hematological remission and renal function restoring. The combination of kidney and hematopoietic cell transplant has been reported as a promising approach to reestablish end-organ function and effectively treat the underlying disease. This review describes current protocols used to perform kidney transplantation in patients with plasma dyscrasias.

Key words: Plasma dyscrasia; Multiple myeloma (MM); Amyloidosis (AL); Hematopoietic cell transplantation; Kidney transplant

INTRODUCTION

Gammopathies, multiple myeloma (MM), and amyloidosis (AL) are plasma cell dyscrasias (PCD) characterized by clonal proliferation of plasma cells, monoclonal immunoglobulin secretion, and often renal impairment (20–30% of patients at the diagnosis) (2,14). In the past decades, the diagnosis of hematological dyscrasias discouraged kidney transplant for many reasons; the immunosuppressive therapy required for transplantation may enhance neoplasia progression and increase the risk of sepsis and infection, and paraprotein overexpression may affect the graft (25). Recently, it has been reported that novel chemotherapy combined with autologous stem cell transplantation may improve the outcome of kidney transplant in PCD patients (25). The aims of this review are to describe the current protocols for kidney transplant specifically applied in patients with MM and AL and to identify patients who may benefit from this therapeutic option.

MEDLINE SEARCH PARAMETERS

We performed a MEDLINE search to identify published articles on kidney transplantation and hematopoietic cell transplantation (HCT) in multiple myeloma or amyloidosis patients. Search terms were “multiple myeloma,” “monoclonal gammopathy,” “stem cell transplantation,” “kidney transplantation.” We chose the English language and clinical trial and case reports as the types of manuscripts. Reviews were excluded. A further selection was done by using “kidney transplantation” and “stem cell transplantation” and “multiple myeloma” or “amyloidosis.” We selected 16 studies that described renal transplantation before, after, or concomitant with hematopoietic cell transplantation (Fig. 1).

PLASMA CELL DYSCRASIAS

Monoclonal Gammopathy of Undetermined Significance

Monoclonal gammopathy of undetermined significance (MGUS) is one of the most common premalignancy

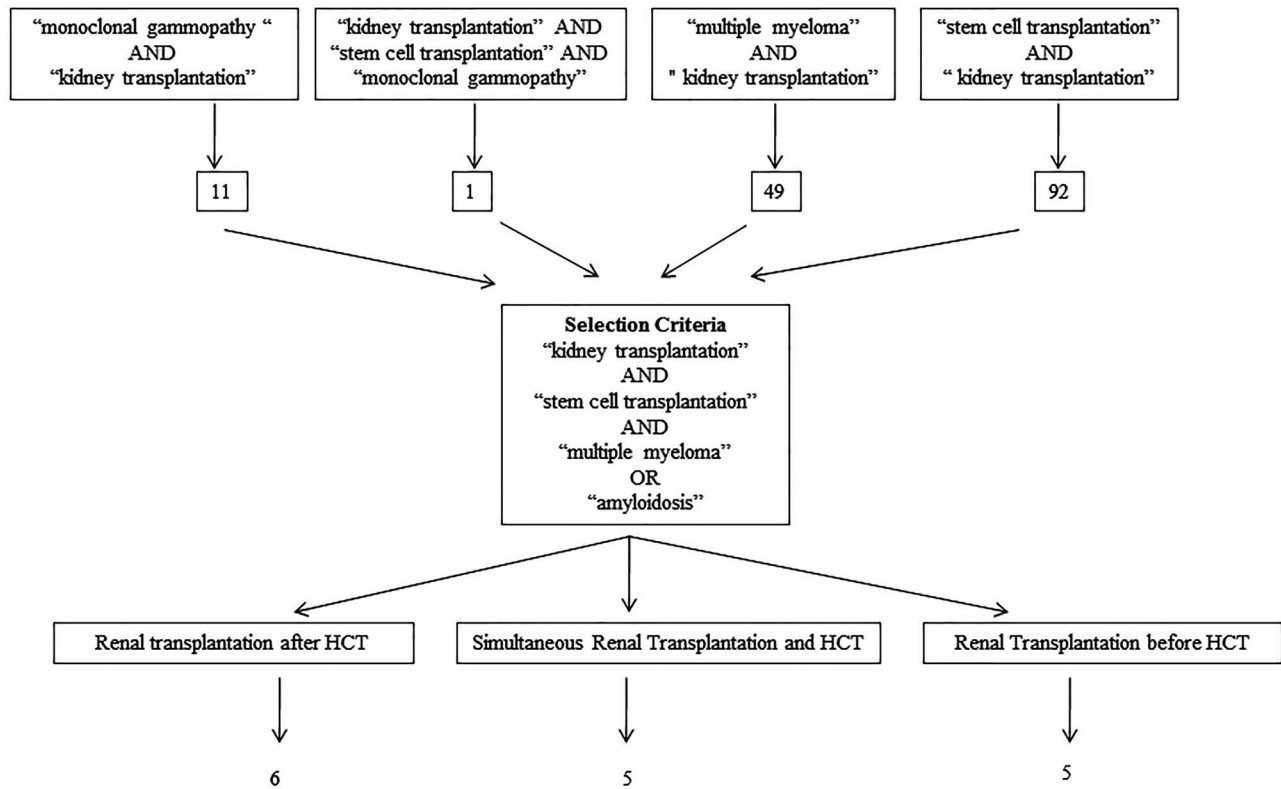


Figure 1. Terms and procedure of literature evaluation.

plasma cell dyscrasia in the Western countries (prevalence 3.2%). Immunoglobulin (M protein) concentration less than 3 g/dl and plasma cells in the bone marrow less than 10% are specific MGUS features, as well as the absence of hypercalcemia, renal impairment, anemia, lytic bone lesions, and any other clinical symptoms of multiple myeloma or lymphoproliferative disorders (16). MGUS is considered a prestage of multiple myeloma, and the increased ratio of M protein concentration and free light chains (FLCs) are predictive of a progression toward

malignancy (17). The rate of malignant transformation of MGUS is 1% per year (1). Considering the high prevalence of MGUS in the general population, recipients for kidney transplantation with MGUS may be common, and therefore it is important to evaluate the risk profile in terms of rejection and MM progression for these patients. Furthermore, MGUS is often associated to renal impairment, even when the hematological disorders do not meet clinical parameters for smoldering multiple myeloma or MM in terms of malignancy. Leung et al. proposed

Table 1. Kidney Disease in Multiple Myeloma

Renal Disease	Molecular Mechanism	Cause and Related Syndromes
Tubulointerstitial lesions cast nephropathy (40–63%)	Glomerular filtration and deposition of free light chains leading to distal obstruction	Dehydration, hypercalcemia, contrast medium usage, nephrotoxic medications (NSAIDs, diuretics, angiotensin-converting enzyme inhibitors, angiotensin II receptor blockers); acute kidney failure
Monoclonal immunoglobulin deposition disease (19–26%)	Deposition of light and/or heavy chains in the basement membranes of the glomeruli and tubules	Nephrotic syndrome and progressive chronic kidney damage
AL amyloidosis (7–30%)	Proteinaceous material deposition in the mesangium, glomerular basement membranes, blood vessels	Nephrotic syndrome and progressive chronic kidney disease, often associated with a systemic involvement

NSAIDs, nonsteroidal anti-inflammatory drugs.

Table 2. Renal Transplantation After HCT in MM Patient

Reference	Sex (Age)	Nephropathy	Prior Therapies	Conditioning Regimen		BMT Donor	Time to KT	Kidney Donor	Complications	Immunosuppression	Follow-up	Outcome
				GVHD Prophylaxis	HDM							
Hassoun et al. 2008 (12)	M (42)	LCDD	Thal Dex Mel Dox	HDM	ASCT	ASCT	14.1 months	Living related donor	None	N/A	23.6 months	CR; stable graft function
Khoriaty et al. 2006 (15)	F (51)	LCDD	Thal Dex Mel Dox	HDM	ASCT	ASCT	45.7 months	Living related donor	None	N/A	23.6 months	CR; stable graft function
Beitinjaneh et al. 2010 (7)	M (56)	ESRD	N/A	N/A	ASCT	ASCT	2.9 years	Living unrelated donor	Herpes zoster	Mycophenolate Mofetil, Prednisone	6	CR
Barracough et al. 2007 (6)	M (58)	LCDD	N/A	Mel; Thal Dex Mel	Autologous PBSC	Autologous PBSC	1	HLA-identical living donor	Severe neutropenic; LCDD recurrence at 26 months after KT	N/A	26 months	Stable renal function
Sanchez Quintana et al. 2003 (25)	M (38)	LCDD	Dex	Lenalidomide	ASCT	ASCT	4	Cadaveric Donor	Acute kidney rejection at 1 month	Steroids Tacrolimus Mycophenolate Mofetil	4	CR; no allograft rejection; no infections
	F (44)	MM	VAD Thal	Lenalidomide	ASCT	ASCT	4	Cadaveric Donor	None	Steroids Tacrolimus Lenalidomide	3	VGPR no allograft rejection; no infections

GVHD, graft-versus-host disease; KT, kidney transplantation; BMT, bone marrow transplantation; MM, multiple myeloma; LCDD, light-chain deposition disease; ASCT, autologous stem cell transplantation; PBSC, autologous peripheral stem cell transplantation; Thal, thalidomide; Dex, dexamethasone; Dox, doxorubicin; Mel, melphalan; HDM, high-dose melphalan; VAD, vincristine-adriamycin-dexamethasone; CR, complete remission; VGPR, very good partial response; N/A not available.

a novel definition “monoclonal gammopathy with renal significance” to discriminate these pathological features from true MGUS (18).

Multiple Myeloma

Kidney disease in MM is mainly caused by conditions summarized in Table 1. The light chain cast, nephropathy, is the most common kidney disease and affects 32–48% of patients who died from MM (21). Histologically, myeloma-dependent cast nephropathy (MCN) shows intratubular light chain casts in distal tubules and collecting ducts. Casts are aggregates of monoclonal FLCs and Tamm Horsfall proteins (THPs). In normal conditions, FLCs are reabsorbed in proximal tubules via receptor-mediated endocytosis after glomerular filtration. In MM, the excess of FLCs overwhelms the capacity of proximal tubules to endocytose them, and monoclonal FLCs pass into the loop of Henle and interact with THPs. The FLC CD3 region binds THP carbohydrate chains and forms aggregates. Several factors such as urinary concentration of light chains, pH, urine flow rate, sodium chloride, and calcium concentrations may favor FLC–THP aggregation (21). Tubule obstructions induce strong inflammation, nuclear factor κ -light chain enhancer of activated B cell (NF- κ B) activation and production of monocyte chemoattractant protein 1 (MCP-1) and interleukin-6 (IL-6) (21).

A giant cell reaction is frequently observed around the casts where mononuclear cells are recruited, and this inflammatory response damages tubules (21). Bortezomib or thalidomide treatment and/or use of a high cutoff dialyzer are suggested in these patients to drastically reduce serum FLC levels. The combination of bortezomib and serial plasma exchange resulted in complete or partial renal response, remarkably reversing the acute renal failure secondary to cast nephropathy in MM newly diagnosed and relapsed patients (10). MCN becomes a medical emergency when patients are forced to undergo chronic dialysis. Although renal transplant is not recommended in the presence of MM because of poor prognosis, there are some cases of patients who underwent renal transplant and were in remission reported in the literature.

Monoclonal Immunoglobulin Deposition Disease

Monoclonal immunoglobulin deposition diseases (MIDDs) include light chain deposition disease (LCDD), light–heavy chain deposition disease (LHCDD), and heavy chain deposition disease (HCDD). MIDDs are characterized by amorphous to granular deposition of monoclonal immunoglobulin or its components. When the β edges in the complementary determining region 2 (CDR2) of light chains are exposed, light chains spontaneously aggregate into oligomers and fibrils and contribute to deposition.

Table 3. Renal Transplantation After HCT in Amyloidosis Patients

Reference	Sex (Age)	Prior Therapies	Conditioning Regimen GVHD Prophylaxis	BMT Donor	Kidney Donor	Complications	Immunosuppression	Follow-up (Months)	Outcome
Herrmann et al. 2011 (13)	F (62)	None	Mel	ASCT	Living related donor	None	Thymo tacrolimus, mycophenolate mofetil (MMF) corticosteroids	53	CR; KTx survival 42.4 months
	F (55)	None	Mel	ASCT	Living related donor	None	Basiliximab, tacrolimus, mycophenolate mofetil (MMF) corticosteroids	41	CR; KTx survival 41.4 months; cause of death: stroke
	M (57)	None	Mel	ASCT	Cadaveric donor	Cellular KTx	Daclizumab, tacrolimus, mycophenolate mofetil (MMF) corticosteroids	39	CR; KTx survival 33.9 months
	M (37)	None	Mel	ASCT	Living related donor	Borderline KTx	Basiliximab, tacrolimus, mycophenolate mofetil (MMF) corticosteroids	39	CR; KTx survival 60.5 months
	F (35)	None	Mel	ASCT	Living related donor	None	Basiliximab, tacrolimus, mycophenolate mofetil (MMF) corticosteroids	75	PR; KTx survival 23.1 months
	M (53)	None	Mel	ASCT	Living related donor	None	Thymo tacrolimus, mycophenolate mofetil (MMF) corticosteroids	64	CR; KTx survival 10.4 months

GVHD, graft-versus-host disease; BMT, bone marrow transplantation; KT, kidney transplantation; Mel, melphalan; ASCT, autologous stem cell transplantation; CR, complete remission; PR, partial remission. Time to kidney transplantation was not available.

The main histological lesion observed in two thirds of MIDD cases is a nodular mesangial sclerosis; mesangial sclerosis without nodules and with membranoproliferative features has also been described.

AL Amyloidosis

Amyloid light chain (AL) amyloidosis is a systemic disease characterized by extracellular deposition of Congoophilic fibrils in soft tissue. Amyloid is constituted by immunoglobulin or its components and appears as amorphous periodic acid Schiff-negative and silver-negative deposits in kidney biopsy. These deposits can be found mainly in vessel walls, in the glomeruli, and in tubular interstitium (22).

HEMATOPOIETIC CELL TRANSPLANT (HCT)

The gold standard treatment of plasma cell dyscrasias is high-dose melphalan followed by autologous stem cell transplantation (ASCT), even during renal failure (27). According to the National Comprehensive Cancer Network (NCCN) guidelines, the primary therapeutic options for transplant candidates include bortezomib/dexamethasone, bortezomib/doxorubicin/dexamethasone, bortezomib/thalidomide/dexamethasone, and lenalidomide/dexamethasone (3). These approaches are followed by hematopoietic cell transplantation (HCT) and addressed to patients younger than 65 years without renal dysfunction (24). For older patients with chronic nephropathy, reduced-intensity conditioning regimens and ASCT should be recommended. The use of bortezomib or lenalidomide, before or after autologous transplantation, significantly increases complete remission rate and progression-free survival (24).

Another possible approach may be the combination of high-dose chemotherapy and ASCT followed by reduced-intensity conditioning allo-SCT. This procedure has decreased treatment-related mortality to 15% or less (8). Evidence about higher safety of tandem transplantation compared to single hematopoietic cell transplant is lacking, and the risk of death and adverse events is still high (7).

In patients with PCD-related chronic nephropathy, the severity of renal dysfunction does not affect the response to chemotherapy. However, autologous stem cell transplant induces complete remission and longer survival than chemotherapy (23). Numerous retrospective studies describe the efficiency of hematopoietic cell transplant even in advanced renal failure that requires kidney transplantation. Regression of renal damage after 4–5 months from ASCT has been also reported (23).

RENAL TRANSPLANTATION AFTER HCT

Kidney transplant after HCT may be a favorable therapeutic approach, especially in dialytic younger patients with a good performance status. Considerable evidence indicates that HCT improves the quality of life and offers

higher survival rate than dialysis in patients with myeloma-related end-stage renal disease (ESRD) (25).

Five studies (6,7,12,15,25) describe seven patients undergoing renal transplantation after HCT (Table 2). Five patients received kidneys for transplantation from matched living related donors and two from cadaveric donors. Time between HCT and renal transplantation ranged from 14.1 months to 4 years. Follow-up ranged from 14.3 months to 6 years. In four patients chronic immunosuppression therapy was based on prednisone, tacrolimus, mycophenolate, and in two cases on lenalidomide. Acute kidney rejection at 1 month from kidney transplantation was reported in one patient efficiently treated with steroids. Recurrence of light chain deposition 26 months after renal transplant was also described. During follow-up, complete remission and renal function restoration were observed. Noteworthy, the graft rejection rate did not increase when the kidney donor was different from the HCT donor. Similar results were described in patients with amyloidosis-related nephropathy (13) (Table 3).

SIMULTANEOUS HCT AND RENAL TRANSPLANT

Table 4 (9,11,23,26) shows results from 16 PCD patients treated with nonmyeloablative conditioning and combined kidney and bone marrow transplantation from human leukocyte antigen (HLA)-haploidentical-related donors. Fifteen patients received a conditioning regimen based on cyclophosphamide, anti-thymocyte globulin and thymic irradiation; one patient received cyclophosphamide and melphalan. Follow-up ranged from 2.4 months to 12.1 years. Ten patients showed chimerism; in seven patients a mixed lymph-hematopoietic chimerism (donor cells were >1% of recipient's cell population) was observed, and three patients achieved full donor chimerism (100% blood cells of recipients derived from donor). Two patients died; one from basal disease progression and one from therapy-related acute myeloid leukemia. Nine out of 16 patients achieved durable complete remission (CR), three a prolonged partial remission (PR), and four patients experienced disease progression. Six patients developed graft-versus-host disease (GVHD), thus requiring immunosuppressive therapy; one had acute cellular rejection of the kidney transplant. Ten patients maintained a stable tolerance to the kidney graft without immunosuppression. This result was partially attributable to mixed lymphohematopoietic chimerism that guaranteed better outcomes than full chimerism. It is remarkable that transplantation tolerance was achieved in patients who had only transient donor hematopoietic chimerism.

Therefore, nonmyeloablative conditioning followed by combined kidney–bone marrow transplantation from HLA-haploidentical-related donors may be considered as a potentially long-term beneficial option. A conditioning

Table 4. Simultaneous Bone Marrow and Kidney Transplantation

Reference	Sex (Age)	Nephropathy	Prior Therapies	Complications	Immunosuppression	Follow-up	Outcome
Wagner et al. 2013 (27)	M (45)	κ light-chain disease	B THAL Dex	Steroid-responsive acute GVHD	Tacrolimus sirolimus, methyl-PRED gradually discontinued	2.4 years	CR; full donor chimerism; good renal function
Fudaba et al. 2006 (11)	M (34) F (35)	MM ESRD due to MM	Dox + Dex Mel VAD	Chronic GVHD None	MMF, PRED for GVHD CY discontinued by day 60 PRED after 2nd PBSC for GVHD	4.3 years 3.5 years	CR; full donor chimerism; no kidney rejection PD; second PBSC at 2.1 years; mixed donor chimerism <71 day; no kidney rejection
	M (35)	ESRD due to MM	Thal + Dex followed by B/Dex; CP VAD	Acute/chronic GVHD None	PRED for GVHD	2.8 years	PD; no kidney rejection; full donor chimerism
	M (51)	ESRD due to MM	VAD	None	CY discontinued by day 76	5.3 years	PR, then PD; mixed donor chimerism <123 day; no kidney rejection
	F (55)	ESRD due to MM	Mel Dex	None	CY discontinued by day 73	7.3 years	CR; mixed donor chimerism <105 day; no kidney rejection
	M (56)	ESRD due to multiple myeloma	Thal + Dex	None	CY discontinued by 1.8 year	2 years	CR MM; mixed donor chimerism <98 day; kidney rejection at 108 day
Spitzer et al. 2011 (26)	F (55)	Cast nephropathy	MEL/PRED, DEX	None	None	12.1 years	CR; mixed donor chimerism <105 day stable renal allograft function without immunosuppressive therapy
	F (51)	ESRD due to MM	VAD	None	None	7.7 years	PR; dead 7.7 years after transplant due to progressive MM
	M (34)	LCDD	VAD, MEL	Chronic GVHD	Steroid	9.2 years	CR followed by relapse; stable renal allograft function
	F (35)	Cast nephropathy	VAD	Chronic GVHD after second SCT	CNI	8.3 years	PD (CR after second SCT)
	F (35)	Cast nephropathy	THAL/DEX, CY	Acute GVHD	Steroid	4.2 years	PD; recurrent cast nephropathy; hemodialysis; death for therapy-related acute myeloid leukemia
	M (57)	LCDD	THAL/DEX	Acute cellular rejection of KT _x	None	6.8 years	CR
	F (47)	Cast nephropathy	THAL/DEX/Dox	Chronic GVHD	None	4.0 years	CR
Buhler et al. 2002 (9)	F (55)	ESRD secondary to κ light-chain multiple myelomas	Mel Dex	None	CY discontinued on day 73 posttransplant	4 years	CR; mixed donor chimerism <105 day; no kidney rejection
	F (50)	ESRD secondary to κ light-chain multiple myelomas	VAD	None	CY discontinued on day 77 posttransplant	2 years	PR, then PD of MM; mixed donor chimerism <123 day; no kidney rejection

For all patients the conditioning regimen was CP+ thymic irradiation + equine antithymocyte globulin; all patients received BMT and KT from HLA-identical related siblings. In Wagner et al. (27) the conditioning regimen was CP+Mel, and the patient received ASCT and BMT and KT from HLA-identical living allogeneic donor. GVHD, graft-versus-host disease; KT, kidney transplantation; BMT, bone marrow transplantation; MM, multiple myeloma; ASCT, autologous stem cell transplantation; PBSC, peripheral blood stem cell transplantation; CP, cyclophosphamide; B, bortezomib; Thal, thalidomide; Dex, dexamethasone; PRED, prednisone; Dox, doxorubicin; CY, cyclosporine A; Mel, melphalan; VAD, vincristine-adriamycin-dexamethasone; CR, complete remission; PR, partial remission; PD, disease progression.

Table 5. Renal transplantation before HCT

Reference	Sex (Age)	Nephropathy	Prior Therapies	Conditioning Regimen GVHD Prophylaxis		Kidney Donor	Time to BMT	BMT Donor	Complications	Immunosuppression	Follow-up*	Outcome
				HDMEL	HDMEL							
Herrmann et al. 2011 (13)	M (52)	Amyloidosis	Tacrolimus, MMF and corticosteroids	HDMEL	HDMEL	Living related donors	N/A	ASCT	None	Tacrolimus, MMF, corticosteroids	103.9 months	CR; KT survival 103.9 months
	M (64)	Amyloidosis	Thymo, CY, MMF, corticosteroids	HDMEL	HDMEL	Living unrelated donors	N/A	ASCT	None	Thymo, CY, MMF, corticosteroids	95.1 months	CR; KT survival 95.1 months
	F (60)	Amyloidosis	Thymo, tacrolimus, MMF, corticosteroids	HDMEL	HDMEL	Living related donors	N/A	ASCT	KT rejection borderline	Thymo tacrolimus, MMF, corticosteroids	83.7 months	CR; KT survival 83.7 months
	F (64)	Amyloidosis	Thymo, tacrolimus, MMF, corticosteroids	HDMEL	HDMEL	Living related donors	N/A	ASCT	None	Thymo tacrolimus, MMF, corticosteroids	78.3 months	CR; KT survival 78.3 months
	M (50)	Amyloidosis	Thymo tacrolimus, MMF, corticosteroids	HDMEL	HDMEL	Living related donors	N/A	ASCT	Cellular KT rejection	Thymo tacrolimus, MMF, corticosteroids	72.7 months	CR; KT survival 72.7 months
	F (61)	Amyloidosis	Thymo tacrolimus, MMF, corticosteroids	HDMEL	HDMEL	Living related donors	N/A	ASCT	None	Thymo tacrolimus, MMF, corticosteroids	61.7 months	CR; KT survival 61.7 months
	M (58)	Amyloidosis	Thymo, tacrolimus, MMF, corticosteroids	HDMEL	HDMEL	Living unrelated donors	N/A	ASCT	Cellular KT rejection	Thymo tacrolimus, MMF, corticosteroids	8.8 months	CR; KT survival 8.8 months; death for acute pulmonary embolism 9 months after KTx
	M (56)	Amyloidosis	Thymo, tacrolimus, MMF, corticosteroids	HDMEL	HDMEL	Living unrelated donors	N/A	ASCT	Cellular KT rejection	Thymo tacrolimus, MMF, corticosteroids	8 months	CR; KT survival 8 months; death CMV infection and posttransplant lymphoproliferative disorder

(continued)

Table 5. (Continued)

Reference	Sex (Age)	Nephropathy	Prior Therapies	Conditioning Regimen		Kidney Donor	Time to BMT	BMT Donor	Complications	Immunosuppression	Follow-up*	Outcome
				GVHD	Prophylaxis							
Leung et al. 2009 (20)	M (58)	Amyloidosis	Thymoglobulin, tacrolimus, MMF, corticosteroids	MEL	MEL	Living unrelated donor	4 months	ASCT	Steroid-resistant cellular rejection (Banff 1A)	Thymo tacrolimus, MMF, corticosteroids	43 days	No negative impact of calcineurin inhibitor and MMF on hematopoietic SCT
Leung et al. 2005 (19)	M (65)	Amyloidosis	ATG/SRL/ MMF	MEL	MEL	Living kidney donor	379 days	ASCT	Bacteremia; no KT rejection	SRL, MMF, PRED	70 months	Alive with stable renal allograft function
	F (60)	Amyloidosis	ATG/TAC/ MMF	MEL	MEL	Living kidney donor	132 days	ASCT	CMV colitis, oral candida; No KT rejection	Thymo tacrolimus, MMF, PRED	55 months	Alive with stable renal allograft function
	F (64)	Amyloidosis	ATG/TAC	MEL	MEL	Living kidney donor	245 days	ASCT	Nausea/anorexia; no KT rejection	Thymo tacrolimus, PRED	40 months	Alive with stable renal allograft function
	M (51)	Amyloidosis	ATG/TAC/ MMF	MEL	MEL	Living kidney donor	138 days	ASCT	Nausea/vomiting; no KT rejection	Thymo tacrolimus, MMF, PRED	33 months	Alive with stable renal allograft function

F (62)	Amyloidosis	ATG/TAC/ MMF	MEL	Living kidney donor	87 days	ASCT	Nausea/vomiting; no KT rejection	Thymo tacrolimus, MMF, PRED	13 months	Alive with stable renal allograft function
Aoki et al. 2013 (4)	M (35) MM	PRED, CY, MMF after KTx; discon- tinuation of MMF, Vel, Dex Mel, CP, THAL	Fludarabine MEL; tacrolimus methotrexate	HLA- identical related siblings	5 years	tandem auto- PBSC/RIC allo-HSCT from an unrelated male donor	Primarily cytomegalo- virus (CMV) pneumonia, pneumocystis pneumonia (PCP), and disseminated candida; HBV reactivation; GVHD	Lenalidomide Dex as salvage therapy; immunosuppressive therapy was discon- tinued after MM relapsed (5 months after allo-HSCT)	8 months	Alive; without recurrence of GVHD; full donor chime- rism; stable renal function
Au et al. 2003 (5)	M (65) ESRD	CY, corticos- teroids CP, epirubicin, vincristine, PRED	Fludarabine total body irra- diation CYC, MMF	Living kidney donor	13 years	HSCT from an related donor	Monomorphic plasmacytoma- like PTLD after 13 years KT ; 4 months post- HSCT, multiple spinal cord infiltrations	CY, MMF	14 years	Cause of death: cerebral metastases

GVHD, graft-versus-host disease; KT, kidney transplantation; BMT, bone marrow transplantation; ASCT, autologous stem cell transplantation; HSCT, hematopoietic stem cell transplantation; auto-PBSC, autologous peripheral blood stem cell; RIC, reduced-intensity conditioning; MMF, mycophenolate mofetil; PRED, prednisone; THAL, thalidomide; BXM, basiliximab; CY, cyclosporine; CP, cyclophosphamide; ATG, anti-thymocyte globulin; SRL, sirolimus; HDMEF, high-dose mephalan; MEF, mephalan; CR, complete remission; CMV, cytomegalovirus.
*Follow-up was considered after kidney transplant.

regimen with reduced toxicity may improve renal allograft tolerance and diminish the need of long-term immunosuppression (11).

RENAL TRANSPLANTATION BEFORE HCT

In five studies kidney transplantation from living donor before autologous ASCT has been described (4,5,13,19,20) (Table 5). Out of 16 patients, 14 had amyloidosis-related ESRD, one MM-related ESRD, and one primary ESRD. Follow-up ranged from 43 days to 103.9 months. Time between renal transplantation and HCT ranged from 87 days to 13 years. In patients with amyloidosis, immunosuppressive therapy after transplant was calcineurin inhibitors + mycophenolate mofetil + steroids. Five patients had acute cellular rejection with a higher tendency for graft rejection in patients who received renal transplant after HCT. All patients had complete remission from amyloidosis and stable renal function. Out of 16 patients, three died of cardiovascular complications (one acute pulmonary embolism 9 months after kidney transplantation) and two of neoplasia (one posttransplant lymphoproliferative disorder and one brain metastases). Therefore, renal transplant followed by a high dose of melphalan and ASCT represents an effective treatment for AL amyloidosis with a good hematological response and an improved median survival (13,21). In MM patients, the effectiveness of tandem autoperipheral blood cell transplantation/ reduced-intensity conditioning (PBSC/RIC) allo-HCT after renal transplantation remains unclear.

CONCLUSION

Renal impairment is detrimental both in multiple myeloma and in gammopathies and is often associated with morbidity and mortality. Several studies have highlighted that renal and hematopoietic transplantation from the same donor could restore organ function and efficiently treat the underlying disease. In particular, the transient mixed chimerism after combined kidney and hematopoietic cell transplantation allows the withdrawal from all immunosuppressive therapy, thus favoring normal graft functioning. Sequential living donor kidney transplant followed by autologous ASCT after conditioning with melphalan seems to be the most effective treatment for amyloidosis in ESRD patients.

Therefore, the combination of chemotherapy/ASCT and kidney transplant is pivotal to pursue renal restoration and improve the overall survival rate. The introduction of high cutoff dialyzers might further facilitate FLC removal and partially overcome renal impairment. Understanding the molecular mechanism of each renal pathological pattern is critical to choose the best strategy to manage kidney transplant in these patients. General health conditions and type of hematological dyscrasia have to be considered to select those patients who may

benefit most from kidney transplant (i.e., young patients who can better tolerate a high dose of melphalan). The need of kidney transplant in multiple myeloma patients remains high, especially in those patients who present some degree of persistent chronic kidney disease after all therapeutic options described above have been exhausted. Future strategies will optimize the protocol to reduce immunoglobulin production and preserve renal function as early as possible after the diagnosis of gammopathy in order to prevent further renal damage. Further large prospective studies to identify novel strategies to manage renal and hematopoietic cell transplantation in ESRD patients should be encouraged.

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