

# Kidney Transplantation

## Kaposi Sarcoma in Kidney Transplanted Patients

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### ABSTRACT

**Purpose:** Newly developed malignancies in kidney transplanted patients are one of the complications attributed to immunosuppression. Kaposi sarcoma is an unusual malignancy in general population, but may develop in kidney transplanted patients with highly varying prevalence. Our aim is to evaluate the prevalence, clinical manifestations, and outcome of Kaposi sarcoma in kidney transplanted patients.

**Materials and Method:** Five hundred and eighty cases (330 male, 250 female) with a mean age of 38.2 were followed for 36 months (range 9 months to 10 years), visiting every two months. History taking and physical examination with emphasis on skin and mucosa were taken. Biopsy of suspicious skin, mucosal, and visceral lesions assigned by other paraclinical methods was performed. Except 7 cases which were HLA identical to donors, all patients were managed with cyclosporine, Azathioprine and Prednisolone.

**Results:** Fourteen patients (2.2%) developed Kaposi sarcoma (biopsy documented) which constituted 60% of all post-transplantation malignancies. They were 11 males and 3 females with a mean age of 41 years. Sarcoma developed 8 to 31 months after transplantation with an average of 18 months. Of these patients, 13 had skin involvement that one of them had pulmonary involvement too. Another patient had only abdominal involvement. Azathioprine was discontinued in all patients, and cyclosporine was reduced in skin affected patients. In patients with visceral involvement cyclosporine was discontinued and then chemotherapy was initiated. All 3 patients with visceral involvement didn't respond to chemotherapy and expired after 6 months. Of 11 patients with skin involvement, one had complete and 2 had incomplete remission of whom, one expired due to acute rejection. Renal function in 8 patients was acceptable, but 2 had impaired renal function, yet didn't need dialysis.

**Conclusion:** Prevalence of Kaposi sarcoma in our patients is more than western countries. Visceral involvement is uncommon, but has poor prognosis. Reducing immunosuppression with discontinuation of Azathioprine and significant reducing cyclosporine dosage can cease skin involvement, with preserving renal function in most of the patients.

**KEY WORDS:** Kaposi sarcoma, malignancy, kidney transplantation

### Introduction

New onset malignancies as the subsequent complication of immunosuppressive therapy in kidney transplant recipients are now well known<sup>(1)</sup> and the overall incidence of malignancies is 100 folds

more than general population.<sup>(2)</sup> A series of mechanisms play a role in increasing the risk, each have its own importance in different types of cancers.<sup>(1)</sup> These include compromised immune system, direct carcinogenic effects of drugs, carcino-

genic viruses, chronic antigenic stimulation, uremia, genetic susceptibility, recipient-donor interactions, and environmental factors.<sup>(3-8)</sup>

Some studies shown a 2-8% incidence of non-cutaneous tumors in kidney recipients, made the researchers believe in a low risk of post-transplant malignancies.<sup>(9)</sup> The sampling method of transplant patients underestimates the risk, because in such samples there are always a considerable number of recently transplanted patients. Latest reports indicate 30-50% incidence 20 years after transplantation.<sup>(10-12)</sup> Common cancers in general population (pulmonary, breast, prostate,...) don't have any increase in transplant patients.<sup>(13)</sup> Following non-melanoma skin malignancies and cervix carcinoma in situ, which are the most common ones, a series of cancers uncommon in general population are more prevalent in this group of patients consisting of lymphoma (24% vs. 6%), lip malignancy (6.2% vs. 0.3%), Kaposi sarcoma (5.7% vs. non-significant incidence), renal carcinoma (5.1% vs. 2.4%), etc.<sup>(13)</sup> Kaposi sarcoma has a 400 to 500 times more risk of developing in transplant recipients than in general population.<sup>(13)</sup> However, it is an uncommon cancer which constitutes a varying proportion of new onset malignancies after transplantation.<sup>(1)</sup> Kidney recipients of African, Arabian, Italian, Jewish, Greek, and Turkish ethnics are more susceptible to Kaposi sarcoma, as an association exists between its incidence and the ratio of Mediterranean people living in a community.<sup>(4)</sup> Kaposi sarcoma develops in 0.25% of organ recipients in western countries, that is 2-3% of the whole malignancies in this group of patients. Whereas, 5% of the recipients will have Kaposi sarcoma in Saudi Arabia, which constitutes 40-70% of the tumors.<sup>(15,16)</sup> On the other hand, it is rare in Japan.<sup>(17)</sup> Male to female ratio of the disease is 3:1.<sup>(14)</sup> Interaction of the risk factors such as Herpes Simplex virus and compromised immune system is apparent.<sup>(18)</sup> Complete improvement is seen in some cases by discontinuing immunosuppressants.<sup>(14)</sup>

Kaposi sarcoma is defined as tumors with endothelium covered vascular areas, spindle shaped cells, extravasation of red blood cells, and groups of inflammatory cells.<sup>(1)</sup> Sixty percent of patients suffer from skin or oropharyngeal mucosa involvement<sup>(19)</sup> presents as lesions such as purple maculae with defined margins or refractory granuloma. The rest of patients have visceral involvement, particularly of gastrointestinal

and respiratory systems and lymph nodes.<sup>(19)</sup>

Although visceral involvement may ensue resistant to treatment, complete or relative remission can be achieved by decreasing or ceasing immunosuppressive drugs in 40% of patients with non-visceral lesions.<sup>(14)</sup> In addition, drug discontinuation ensues graft rejection.<sup>(14)</sup> The aim of this study was to depict the incidence, clinical manifestations, and outcome of Kaposi sarcoma in kidney transplant recipients at Golestan hospital, Ahwaz.

## Materials and Methods

A total of 580 patients who had undergone kidney transplantation at Golestan hospital were followed for a mean duration of 36 months (range 9 months to 10 years). Three hundred and twenty of whom were males and 250 were females. Mean age of the patients was 38.2 (range 6 to 66) years. Outpatient visits were performed every two months and physical examination with special focus on skin and mucosa was done.

Paraclinically taken specimens of identified lesions would be sent to pathology. All except 7 patients with complete HLA matching had been under triple drug treatment of Cyclosporine (4-5 mg/kg/day), Azathioprine (2-2.5 mg/kg/day), and Prednisolone (10-12.5 mg/kg/day). Complete HLA matched patients would have undergone the treatment with Azathioprine and Prednisolone only.

## Results

An overall of 14 patients (2.2%) had biopsy confirmed Kaposi sarcoma, including 11 men and 3 women (male to female ratio of 4:1). Mean age of them was 41 (range 27 to 59) years and the mean interval between transplantation and diagnosis was 18 (range 8 to 31) months. All the cases had received triple immunosuppressant therapy and none of them were HLA identical ( $p < 0.005$ ). Two patients had developed acute rejection and subsequently treated with Methyl-Prednisolone pulse therapy (3 g) of whom one had not responded and undergone ATG (Anti-thymocyte Globulin) therapy. Skin involvement was observed in 13 cases of Kaposi sarcoma of which one had simultaneous widespread bilateral pulmonary involvement and one had simultaneous gastric and intestinal involvement. None of the patients had oral, pharyngeal, or laryngeal lesion of mucosa. Gastrointestinal involvement was presented most-

ly with hemorrhagic diarrhea, vomiting, and abdominal pain and was diagnosed by upper GI endoscopy, colonoscopy, and biopsy. The only patient with pulmonary involvement had suffered from skin lesions for a few years and had refused to lower or discontinue immunosuppressive drugs. Three years later the patient came with dyspnea and chest x-ray revealed multiple bilateral nodular lesions. Biopsy via bronchoscopy was impossible, so that open biopsy of lung was performed and Kaposi sarcoma was confirmed. Azathioprine was discontinued in all of the patients and cyclosporine dose was reduced to 50% in the patients with skin lesions, but it was discontinued in the cases of visceral involvement and interferon was initiated as well.

All the three patients with visceral involvement did not respond to drug cessation and chemotherapy and died after an average of 6 months. Renal failure ensued in one of them and led to maintenance dialysis. Complete remission was achieved in 9 out of 11 patients with skin lesions and in the other 2 ones, both refused to cease immunosuppressive drugs, skin lesions were subsided but not eliminated. At the time of this study 10 out of the respective 11 patients were still alive of whom, 8 had functional graft and 2 had impaired graft function but still not dialysis dependent and finally, one had died due to acute rejection and subsequent septicemia.

### Discussion

Kaposi sarcoma has a higher prevalence in our country than the western countries, in which Kaposi sarcoma affects 0.25% of kidney recipients and comprises 2 to 3% of post-transplant malignancies.<sup>(1)</sup> In this study Kaposi sarcoma was observed in 2.2% of the patients and comprised approximately 60% of all post-transplant malignancies (3.4%). The most prevalent malignancies of kidney recipients in western countries are skin tumors and lymphoma, respectively and Kaposi sarcoma takes the third place;<sup>(13)</sup> whereas, it is the most common one according to our study, following by skin tumors and lymphoma. Thus, the prevalence of Kaposi sarcoma in our country seems to be similar to the one in Mediterranean and Arabian countries.<sup>(15,16)</sup> So that, environmental and genetic factors should be taken into account. Solar radiation, ultraviolet radiation, some viral infections, and a series of drugs are the examples of environmental factors.<sup>(1)</sup>

Kaposi sarcoma had a male to female ratio of

4:1, almost the same as the results in other studies.<sup>(14)</sup> Skin and visceral involvement was observed in 78% and 22% of the cases, respectively, compared to 58-60% and 40-42% in other studies.<sup>(13)</sup> The difference may be due to our small sample.

The mean interval between transplantation and developing of malignancies is 61 months and it is 21 months for Kaposi sarcoma which is the earliest.<sup>(20)</sup> This interval was 18 months in our patients.

Skin lesions were recovered in 9 out of 11 patients and relatively improved in 1. Renal graft function was normal in 8 live patients and moderately impaired in 2. It can be concluded that lowering immunosuppression by Azathioprine cessation and aggressive reduction of cyclosporine dose is effective in the improvement of Kaposi sarcoma and renal function will remain normal.<sup>(13)</sup> All the three patients with visceral involvement died in 6 months; although other studies have shown improvement in 30% of such cases, 57% of whom improved merely by dose alteration of immunosuppressants,<sup>(13)</sup> our study showed a poor prognosis for this uncommon complication.

### Conclusion

Prevalence of Kaposi sarcoma in our patients is more than western countries. Visceral involvements are uncommon but have poor prognosis. Reducing immunosuppression with discontinuation of Azathioprine and significant reducing cyclosporine dosage can cease skin involvement, with preserving renal function in most of the patients. It is recommended that Kaposi sarcoma should be considered if refractory infectious granuloma, blue to red maculae or plaques in the skin, or oropharyngeal mucosa is observed and subject to confirmed diagnosis, visceral involvement is necessary to be fully evaluated.

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