Kimura's disease associated with IgA nephropathy: A case report

WEI ZHANG¹, ANCHARAZ PREEATUM² and CHUNYAN LIU¹

¹Department of Nephrology, Second Affiliated Hospital of Dalian Medical University, Dalian, Liaoning 116000, P.R. China; ²Nephrology Unit, Internal Medicine Department, Flacq Regional Hospital, Central Flacq 40606, Mauritius

Received April 14, 2018; Accepted November 11, 2018

DOI: 10.3892/etm.2018.7103

Abstract. Kimura's disease (KD) is a rare chronic inflammatory disorder of unknown etiology that manifests as painless subcutaneous lesions in the head and neck. It primarily affects young Asian males and is characterized by peripheral eosinophilia and elevated levels of serum Immunoglobulin (Ig) E. Histologically, a variety of renal lesions have been observed in patients with KD, although IgA nephropathy is rarely reported. In the current study, a case of KD with an atypical manifestation accompanied with IgA nephropathy was reported in a middle-aged Chinese man. The subcutaneous lesion was located on the elbow of the patient. Although the recurrence rate of KD-associated nephrotic syndrome is high, the patient has not experienced relapse following 6 months of pulse treatment with methylprednisone and cyclophosphamide. In conclusion, methylprednisone combined with cyclophosphamide is effective in the treatment of KD-associated IgA nephropathy, and long-term administration of methylprednisone may achieve the effect of preventing KD recurrence.

Introduction

Kimura's disease (KD) was first described by Kimm and Szeto in 1937 (1) and the definitive histological description of the disease was subsequently reported by Kimura in 1948 (2). KD is a rare chronic inflammatory disorder that may lead to lymphadenopathy and subcutaneous lesions. It is characterized by plasma and tissue eosinophilia, raised levels of immunoglobulin (Ig)E and frequent complications of nephropathy (3). KD occurs mostly in Asian patients and has been considered limited to the Far East (4). It has previously been reported that 12-60% of patients with KD also exhibit coexisting renal disease (5). A number of histopathological renal lesions have been reported in this disease, particularly minimal change disease, membranous nephropathy and mesangioproliferative disease (6). However, to the best of our knowledge, IgA nephropathy in patients with KD has rarely been reported. In the current study, a case of steroid-responsive IgA nephropathy associated with KD is reported.

Case study

A 45-year-old Chinese man was diagnosed with eosinophilic hyperplasia in May 2011. In June 2012, during a routine health examination, the patient's urine was analyzed and the results revealed the presence of grade 3+ proteinuria. Approximately 6 months later, the patient went to another hospital (The First Affiliated Hospital Of Jinzhou Medical University (Jinzhou, China) and was diagnosed with proteinuria and hematuria. The patient was then admitted to Second Affiliated Hospital of Dalian Medical University (Dalian, China) due to gross hematuria and proteinuria in March 2017.

The patient also exhibited an oval-shaped subcutaneous swelling in his right elbow that had been present for 2 months. The swelling was 2.5 cm in diameter and presented a clear edge. In addition, the mobile swelling was neither tender nor indurated.

Upon physical examination, the blood pressure and heart rate of the patient were 135/80 mmHg and 72 beats/min, respectively. The patient was also determined to be febrile.

The patient had no history of hypertension or diabetes. Laboratory tests revealed proteinuria 3+ (normal range, negative) and 2 consecutive 24 h urine analyses determined a protein content of 3.9 and 3.8 g/d, respectively. Normal values were provided previously (7). The urinary red blood cell (RBC) count of the patient was RBC:20/HP (normal range: 0-2/HP). Upon morphological examination (magnification, x40), the shape of the urine RBCs were dysmorphic, indicating glomerular hematuria. A full blood count revealed: White blood cell count, 13.4x10⁹/l (normal range: 3.50-9.50x10⁹/l); hemoglobin, 155 g/l (normal range: 130-175 g/l); and eosinophilia, 3.53x10⁹/l (normal range, 0.02-0.52x10⁹/l). Furthermore, a renal function test revealed the following: Urea, 6.10 mmol/l (normal range: 3.1-8.0 mmol/l) and serum creatinine, 115 µmol/l (normal range, 57-97 µmol/l). Serum total IgE levels were high, at 9,790.00 IU/ml (normal range 0-100 IU/ml). N Latex IgE mono kit (Siemens AG, Munich, Germany) was used for IgE assay. Hypersensitive c-reactive protein [analysed using an ELISA kit from Monobind, Inc. (Lake Forest, CA, USA); cat. no. 3125-000A] was 77.13 mg/l (normal range: 0-5 mg/l) and the erythrocyte sedimentation

Correspondence to: Dr Chunyan Liu, Department of Nephrology, Second Affiliated Hospital of Dalian Medical University, 467 Zhongshan Road, Dalian, Liaoning 116000, P.R. China
E-mail: liu_chunyan031@sina.com

Key words: Kimura's disease, immunoglobulin A nephropathy, methylprednisone
rate (analysed using an automatic blood sedimentation dynamic analyzer) was 22.00 mm/h (normal range: 0-15 mm/h). The hepatitis profile of the patient was normal. The results for antineutrophilic cytoplasmic antibody and antibodies associated with autoimmune hepatitis (including surface antibodies, E antibodies and core antibodies) were determined to be negative. Ultrasoundography of the subcutaneous mass presented two swollen lymph nodes (2.0 and 1.3 cm, respectively), and the hilus of these nodes indicated that these swollen lymph nodes had a blood supply. Bone marrow aspiration revealed eosinophilia hyperplasia only and excluded the possibility of other hematological diseases. A chest X-ray did not indicate any abnormalities. Ultrasoundography results of the liver and renal artery were normal.

After informed consent was obtained from the patient, a renal biopsy was performed. The tissue was prefixed with ethanol:formaldehyde:PBS (7:3:1) at 4°C for 10 min. Following prefixing, the tissues were fixed with Bouine stationary liquid (25 ml 80% ethanol saturated picric acid solution + 120 ml formaldehyde + 220 ml anhydrous ethanol + 60 ml glacial acetic acid + 50 ml water) at 4°C for 6 h. Then the tissues were dehydrated at different ethanol concentrations (75% ethanol, 85% ethanol, 95% ethanol, absolute ethyl alcohol) for 20 min at each concentration. The environmental friendly transparent dewaxing liquid was on the tissue for 30 min, then specimens were embedded in paraffin at 65°C for 40 min.

The specimens were embedded in paraffin and sliced to: i) 3-µm thickness and for immunohistochemistry and Congo red staining, ii) 2-µm sections for hematoxylin and eosin (H&E), periodic acid-Schiff and Masson staining, iii) 1.5-µm sections for periodic acid silver methenamine (PASM) staining, and iv) 1-µm sections for PASM + HE staining.

For H&E staining, the nucleus was stained with hematoxylin at 25°C for 10 min, then differentiated with 1% hydrochloric acid for 1 sec, and counterstained with Scott's bluing reagent (0.35 g sodium bicarbonate + 2 g magnesium sulfate + 100 ml water) at 25°C for 10 sec. The cytoplasm was then stained with 5% eosin at 25°C for 5 min.

A total of 34 glomeruli were detected via light microscopy (magnification, x100): Five glomeruli exhibited global sclerosis, one glomerulus exhibited segmental sclerosis, and two glomeruli presented with a cellular segmental crescent (Fig. 1). Renal tubules exhibited localized atrophy and tubule cells presented granular degeneration. A small quantity of protein casts were identified in the lumen and inflammatory cell infiltration was observed in the interstitium. Most infiltrating cells were eosinophils and several regions of the interstitium exhibited marked pathological changes to granulomas (0.35 g sodium bicarbonate + 2 g magnesium sulfate + 100 ml water) at 25°C for 10 sec. The cytoplasm was then stained with 5% eosin at 25°C for 5 min.

Based on the results, the patient was diagnosed with IgA nephropathy and interstitial nephritis.

The patient was administered methylprednisone pulse therapy (intravenous administration, 500 mg/day for 3 days), followed by oral 24 mg/day methylprednisone and 0.8 mg/kg/day cyclophosphamide (the body weight of the patient was 72 kg) for 4 weeks. Following 10 days of treatment, the swelling situated on the elbow disappeared while the peripheral eosinophil count returned to normal (0.06x10⁹/l). However, the levels of total serum IgE (9,030.00 IU/ml) and proteinuria (2.8 g/day) did not show any marked reduction. Following 3 months, the level of serum creatinine dropped to 91 µmol/l, while the level of IgE dropped to 5,653 IU/ml. The patient was discharged in March 2017 following 20 days of hospitalization. During the follow-up visit conducted 6 months following discharge, IgE levels dropped to 2,810 IU/ml and 24 h proteinuria reduced to 0.4 g/day with no evidence of hematuria. In April 2018, the dose of methylprednisone was reduced to 8 mg/day and there was no evidence of relapse, although the patient is still being followed-up.

**Discussion**

KD was first described by Kimm and Szeto (1) in 1937 and the definitive histological description of the disease was reported by Kimura et al (2) in 1948 (2). The majority of KD cases in Asian adults are associated with a protracted course (4,8). KD is characterized by obvious peripheral blood eosinophilia and an elevated level of IgE, with subcutaneous swelling around the head and neck, particularly in the periauricular region (9). In addition, epitrochlear lymph nodes, inguinal lymph nodes and submandibular salivary glands are sites that are commonly affected (10). However, in cases of relapse, the disease may present over the entirety of the patient’s body surface. Among previously reported cases, 1% of swellings were observed on the elbow, >50% of patients exhibited a single lesion and 46.6% of patients exhibited multiple lesions (11). In the majority of cases, a definitive diagnosis can only be made following biopsy of the subcutaneous swelling (6). Although the patient in the current study refused to undergo a biopsy of subcutaneous swelling, his demographics and symptoms were typical of KD.

The association between KD and renal diseases including minimal change disease, membranous nephropathy and mesangiproliferative disease is well recognized (10). Minimal change disease and membranous nephropathy are the primary symptoms observed in patients from Asia, the majority of whom exhibit nephrotic syndromes (12). However, IgA nephropathy in patients with KD is rarely reported. To the best of our knowledge, the association between KD and IgA nephropathy was first reported in 1998 (13). Since then, only one more article (9) reported a case of KD associated IgA nephropathy. In each case, the patients also exhibited nephrotic syndrome. The renal histology results of the patient in the current study revealed granular and mesangial deposits of IgA and complement as detected by immunofluorescence. Therefore, the manifestation of nephritic syndrome in this patient strongly indicated the presence of IgA nephropathy.

KD is an allergic and inflammatory disorder with unknown cause. It is believed that KD is caused by a type I hypersensitivity reaction or via a T cell-associated immune disorder (3). Ohta et al (14) demonstrated that Th2 cells (a subset of T helper...
cells), and Tc1 cells (a subset of CD8+ T cells), may contribute to the pathogenesis of KD. Furthermore, Yamazaki et al (15) demonstrated that an increased level of Th2 cells may serve important roles in the pathogenesis of KD by increasing the synthesis of certain Th2-type cytokines, including interleukin (IL)-4 and IL-5, which consequently increase eosinophilic infiltration and IgE synthesis. In addition, several previous studies have indicated that T cell-mediated tissue damage serves an important role in the immunopathogenesis of IgA nephropathy (16). The involvement of various novel subsets of T cells in the immune responses of IgA nephropathy, including Th17 and Th1 cells, have also been confirmed (17). Previous studies have also determined that, compared with IgA nephropathy (IgAN) patients without proteinuria, a higher percentage of Th22 cells were present in IgAN patients with proteinuria (18). In addition, Th22 cells were revealed to be involved in the immune responses of IgAN (19). During the patient's hospitalization the current study, he presented massive and relatively infrequent proteinuria, which we hypothesize may have been caused by an abnormality in T cell subsets. Taken together, the changes of T cells in patients with concurrent IgA nephropathy and KD remain unclear and should be further investigated.

There is no consensus regarding the optimal treatment for KD. However, surgical excision and systemic steroid therapy are the preferred treatments. Recurrence of KD is frequent after the cessation of treatment (20). A previous study has demonstrated that KD with renal recurrence is affected by sex, age and history of hypertension, and it usually presents a good prognosis (12). Nephrotic syndrome in patients with KD is usually treated with the oral administration of steroid hormones, such as methylprednisolone, but patient response to the treatment varies (21). The occurrence of relapse was also revealed to depend on the results of renal histology (22, 23). In the case of the present study, the patient responded well to methylprednisolone treatment and there was no recurrence of either elbow lesions or nephritis syndrome during follow-up.

A rare case of KD associated with IgA nephropathy was described in the current study. The case also exhibited hematuria and massive proteinuria. Short-term, the patient was successfully treated with methylprednisolone combined with cyclophosphamide, while the prevention of KD relapse was achieved via the long-term administration of methylprednisolone at a maintenance dose (8 mg/d). However, it cannot be concluded whether the long-term administration of methylprednisolone may be used as an effective strategy to prevent KD relapse and further studies are required for confirmation.

Acknowledgments

Not applicable.

Funding

The present study was supported by The technology Plan Projects for Liaoning Province, China (grant no. 20170540279).

Availability of data and materials

All data generated or analyzed during this study are included in this published article.
Authors' contributions

WZ put forward the concepts of the present study, was responsible for designing and data analysis, and was a major contributor in writing the manuscript. CL provided the histological examination of the kidney, and contributed to the manuscript preparation and editing. AP contributed to the manuscript review and editing, and performed the pathological and immunohistochemical interpretation of the renal tissue.

Ethics approval and consent to participate

The present study was approved by the Ethics Committee of Second Affiliated Hospital of Dalian Medical University.

Patient consent for publication

Concent for publication has been provided.

Competing interests

The authors declare that they have no competing interests.

References