


Risk Assessment of Recurrence and Autoimmune Disorders in Kikuchi Disease

This article was published in the following Dove Press journal:
Risk Management and Healthcare Policy

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Purpose: Kikuchi disease (KD) is typically a benign disease. Recent studies reporting recurrence or serious cases suggest a possible association of KD with systemic autoimmune disorders. We performed a long-term analysis of the characteristics of KD in patients of all ages and assessed KD recurrence or progress to systemic autoimmune disorders.

Patients and Methods: Electronic medical records of patients diagnosed with KD between April 1995 and May 2017 were reviewed for clinical and laboratory manifestations.

Results: In total, 480 patients were confirmed to have KD based on histopathology findings. The mean age at KD diagnosis was 24.4 years. Recurrence occurred in 11.3% of patients; 2.7% developed autoimmune diseases after KD diagnosis. Patients who experienced recurrence had more extranodal symptoms, lymphopenia, and a longer lymphopenia-recovery duration. Patients who developed autoimmune diseases after KD were more likely to have extranodal symptoms, KD recurrence, and anti-nuclear antibody positivity.

Conclusion: KD patients with risk factors need to be followed-up for KD recurrence and the development of systemic autoimmune diseases.

Keywords: Kikuchi disease, histiocytic necrotizing lymphadenitis, recurrence, autoimmune disorders

Introduction

Kikuchi disease (KD), also known as histiocytic necrotizing lymphadenitis or Kikuchi-Fujimoto disease, is characterized by fever and lymphadenopathy of unknown etiology and usually affects young Asian women.¹⁻⁴ Recent reports describe KD as a form of lymphadenitis that can occur in people of all races and in both children and adults of both sexes.^{2,5-7} The extranodal symptoms of KD are uncommon and diverse, including skin rash, night sweats, weight loss, headache, cough, and abdominal pain.^{2,6,8,9} Although KD is generally a self-limiting and benign disease, varying rates of recurrence and fatality have been reported.^{2,6,10-14} Associations between KD and systemic autoimmune disorders, such as systemic lupus erythematosus (SLE), Sjögren's syndrome, and adult-onset Still's disease, have been reported.¹⁵⁻²⁰ Studies have also suggested that KD is more likely to develop into systemic autoimmune disorders in patients with KD recurrence, extranodal symptoms, or anti-nuclear antibody (ANA) positivity.^{9-12,21} However, these studies had limitations such as short study durations and small sample sizes. In this study, we aimed to analyze the clinical characteristics of patients with KD of all ages over a period of more than 20 years and to evaluate the risk factors for KD recurrence or progression to systemic autoimmune disorders.

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Patients and Methods

We reviewed the electronic medical records of patients who had been diagnosed with KD between April 1995 and May 2017 at Ajou University Hospital, a tertiary university hospital. KD was diagnosed when all of the following criteria were met: (1) systemic symptoms or physical findings compatible with those for KD; (2) exclusion of any other causes of lymphadenopathy; (3) histological examination of a lymph node biopsy specimen showing typical findings compatible with those for KD. The protocol of this study was approved by the Institutional Review Board of Ajou University Hospital (AJIRB-MED-MDB-20-058). Informed consent from study participants was waived due to the retrospective data collection and patient data de-identification prior to study analyses. The data collected included those of age, sex, presenting symptoms, laboratory test results, histological findings, treatment, and clinical outcome. Recurrence was defined as additional episodes of febrile lymphadenopathy before or after the pathological diagnosis, and these episodes were determined from the patients' medical records, regardless of whether repeated pathological confirmation had been performed. We excluded cases involving recurrent episodes of lymphadenopathy in which other definite causes were identified. Between-group differences were evaluated using Student's *t*- or Pearson χ^2 -test, as appropriate. Two-sided *p*-values of less than 0.05 were considered statistically significant. All statistical analyses were performed using the PASW Statistics 13 (SPSS Inc., Chicago, IL, USA) software package.

Results

General Characteristics of Patients with Kikuchi Disease

During this 22-year period, a total of 480 patients met the definition of KD. The mean age was 24.4 years (range 4–79 years), and 151 patients (31.5%) were aged <18 years (Figure 1A and B). The ratio of male patients to female patients was 1:1.32 for those under 18 years of age and 1:2.46 for those over 18 years of age (Figure 2). Table 1 shows the general characteristics of patients with KD. Of the 480 patients, 455 (94.8%) had lymphadenopathy and 381 (79.4%) had fever at the time of the initial diagnosis. Overall, 61 (12.7%) patients had systemic symptoms in addition to fever. Among systemic symptoms, rash, myalgia, fatigue, cough, rhinorrhea, abdominal pain, diarrhea, loss of appetite, and headache were

predominant. Of the 425 patients whose laboratory findings were available for analysis, 201 (47.3%) had leukopenia (white blood cell count <4000/mm³), 132 (31.1%) had lymphopenia (absolute lymphocyte count <1500/mm³), and 43 (10.1%) had neutropenia (absolute neutrophil count <1500/mm³). Among the 201 patients with leukopenia, 113 (74.8% of 151) were children (under 18 years of age) and 88 (26.7% of 329) were adults (over 18 years of age). Of the 245 patients tested for ANAs, 69 (28.2%) tested positive; of these, 32 patients were younger than 18 years of age (46.4%). Of the 69 patients with ANA positivity, there were 32 (21.2% of 151) children and 37 (11.2% of 329) adults.

Clinical Characteristics of Patients with Recurrent Kikuchi Disease

Fifty-four patients (11.3%) experienced 1–4 recurrent episodes of KD each. The initial recurrence occurred within a mean duration of 6 months (range: 1 month to 6 years). The mean age at the first pathological diagnosis in patients with recurrent KD was 28.5 years (range 8–67 years), and 20 patients (37.0% of 54) were under 18 years of age. The ratio of male patients to female patients was 1:1.57. Table 2 shows the characteristics of patients with recurrent and non-recurrent KD. Of the 54 patients with recurrent KD, 53 (98.1%) patients had lymphadenopathy; 44 (81.5%) had fever; and 18 (33.3%) had extranodal symptoms such as skin rash, myalgia, and headache. Twenty-five patients (46.3%) had leukopenia, 25 (46.3%) had lymphopenia, and 6 (11.1%) had neutropenia. Of the 45 patients with recurrent KD tested for ANAs, 17 (37.8%) tested positive; of these, 8 (47.1%) patients were younger than 18 years of age. In terms of recovery from cytopenia, the mean duration for achieving a normal neutrophil count was 7.1 days (range: 0–10 days) in recurrent cases and 8.4 days (0–32 days) in non-recurrent cases. For recovery from lymphopenia, the mean duration was 14.7 days (range: 0–58 days) in recurrent cases and 6.2 days (range: 0–14 days) in non-recurrent cases. There were no significant differences between recurrent and non-recurrent cases with respect to fever, lymphadenopathy, and ANA positivity. Patients with recurrent KD were more likely to have extranodal symptoms (*p*=0.000) and lymphopenia (*p*=0.032) and to require a longer time to achieve a normal lymphocyte count (*p*=0.002).

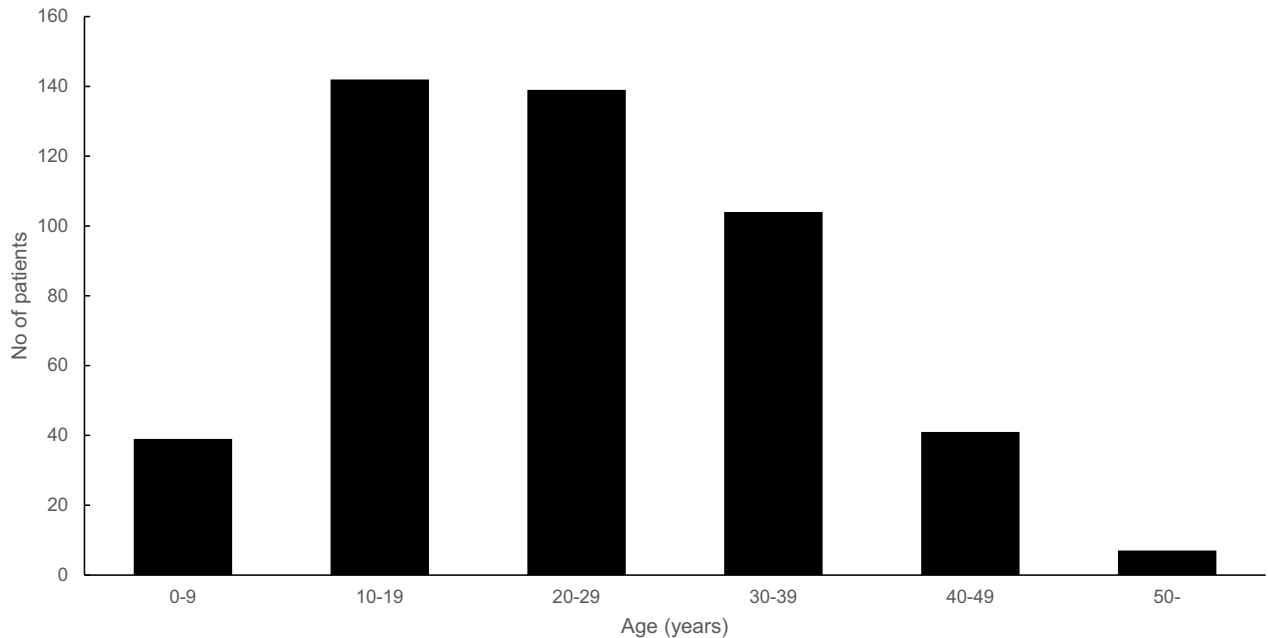
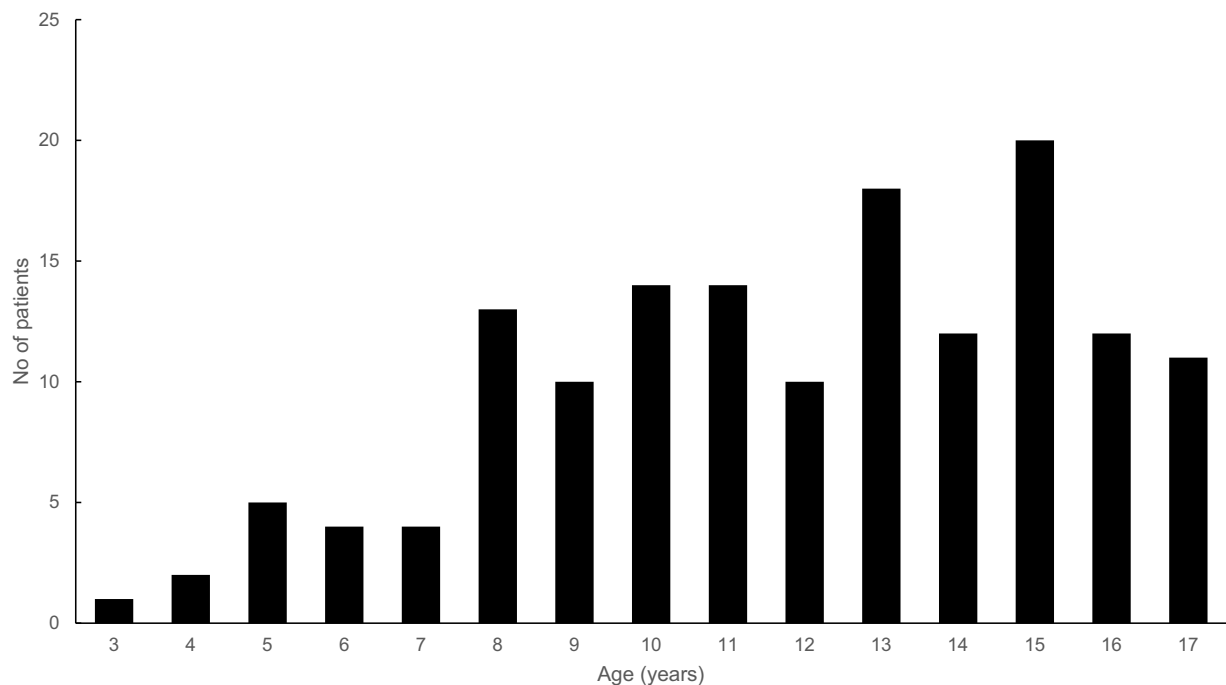
A**B**

Figure 1 Age distribution of Kikuchi disease among (A) all patients and (B) children under 18 years of age.

Clinical Characteristics of Kikuchi Disease Patients Who Developed Systemic Autoimmune Diseases

Systemic autoimmune diseases occurred in 13 (2.7% of 480) patients diagnosed with KD during the study period (Table 3). There were 10 patients with SLE, 1 with

Sjögren's syndrome, 1 with rheumatoid arthritis, and 1 with adult-onset Still's disease. The interval between KD diagnosis and the development of systemic autoimmune diseases ranged from 1 month to 5 years. The mean age at KD diagnosis was 27.9 years (range: 8–67 years). The ratio of male patients to female patients was 1:6. Of

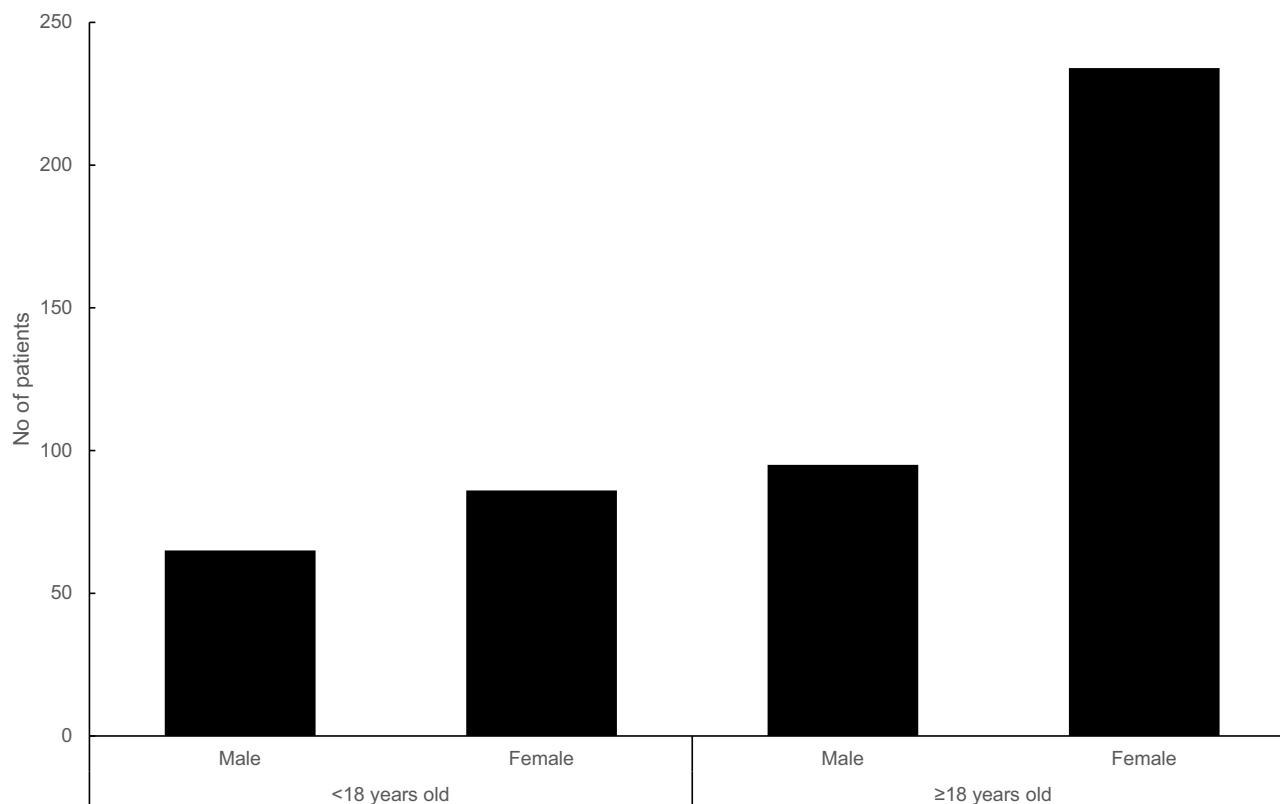


Figure 2 The ratio of male patients to female patients with Kikuchi disease.

these 13 patients, 5 (38.5%) had recurrent KD, and 8 (61.5%) showed ANA positivity at KD diagnosis. All patients in this group had lymphadenopathy, 12 (92.3%) had fever, and 4 (30.8%) had extranodal symptoms including rash, myalgia, fatigue, and abdominal pain. Except for ANA positivity, no other laboratory parameters differed significantly between the patients in this group and those who did not develop systemic autoimmune diseases.

Discussion

KD usually affects young women but can occur in any age group.^{2,5-7} The higher incidence among young women is reflected in the male: female ratio of 1:3–4 in adults; however, this ratio of approximately 1:1 in children.^{1-3,10,12,21} In our study, the mean age at KD diagnosis was 24.4 years, and 88.3% of the patients were younger than 40 years of age (Figure 1A). Interestingly, KD incidence increased significantly after 8 years of age in children (Figure 1B). The sex distribution of KD varies markedly with age. The proportions of male and female patients with KD are similar at a young age (1:1.32), but the proportion of affected female patients increases rapidly with age (1:2.46) (Figure 2). This study shows that KD occurs

mainly in women of childbearing age but can occur in both men and women of all ages. Lymphadenopathy and fever were the predominant symptoms of KD in this study, similar to those in previously reported studies.^{2,3} WBC counts in patients with KD have been reported to vary across studies. Leukopenia has been reported in many studies^{1,11,12,22} and was found in 47.3% of our patients. An interesting finding in this study was that children with KD (age <18 years) were more likely to have leukopenia (74.8% vs 26.7%) and ANA positivity (21.2% vs 11.2%) than adults with KD (Table 1).

The rate of KD recurrence has been reported to be between 3% and 40%, and KD recurrence has been reported to occur more frequently in children.^{1,11,21,22} In our study, the KD recurrence rate was 11.3% in all patients and 13.2% in children under 18 years of age; this value was not significantly different from that in adults. More extranodal symptoms, lymphopenia, and a longer duration for lymphocyte count recovery were significantly associated with KD recurrence (Table 2). Several previous studies have reported the relation between WBC counts and KD recurrence,^{9,11} but the time to recovery from cytopenia had not been previously compared between recurrence and non-recurrence groups. In this study, the

Table 1 General Characteristics of Patients with Kikuchi Disease

Characteristics	
Total patients	100% (n=480)
Mean age at the diagnosis of KD Children (age <18 yr)	24.4 yr. (4–79 yr) 31.5% (n=151)
Male: Female	1:2
Clinical features at the diagnosis of KD	
Fever	79.4% (n=381)
Lymphadenopathy	94.8% (n=455)
Extranodal symptoms	12.7% (n=61)
Recurrent KD	11.3% (n=54)
Laboratory findings (n=425)	
Leukopenia	47.3% (n=201)
Children: adults	74.8% (n=113 of 151); 26.7% (n=88 of 329)
Neutropenia	10.1% (n=43)
Lymphopenia	31.1% (n=132)
ANA (+) at the diagnosis of KD (n=245)	28.2% (n=36)
Children: adults	21.2% (n=32 of 151); 11.2% (n=37 of 329)

Abbreviations: KD, Kikuchi disease; yr, years; leukopenia, white blood cell count < 4000/mm³; neutropenia, absolute neutrophil count < 1500/mm³; lymphopenia, absolute lymphocyte count < 1500/mm³; ANA, anti-nuclear antibody.

Table 2 Clinical Characteristics of Patients with Kikuchi Disease According to Recurrence

	Recurrence (n=54)	Non-Recurrence (n=426)	p-value
Clinical symptoms			
Fever	44 (81.5%)	337 (79.1%)	0.685
Lymphadenopathy	53 (98.1%)	402 (94.4%)	0.239
Extranodal symptoms	18 (33.3%)	43 (9.9%)	0.000
Laboratory features (n=425)			
Leukopenia	25 (46.3% of 54)	176 (47.4% of 371)	0.769
Neutropenia	6 (11.1% of 54)	37 (10.0% of 371)	0.729
Lymphopenia	25 (46.3% of 54)	107 (28.8% of 371)	0.032
ANA (+) at KD diagnosis	17 (37.8% of 45)	52 (26% of 200)	0.113
Time to recovery from cytopenia			
Neutrophil	7.1 days (0–10 days)	8.4 days (0–32 days)	0.365
Lymphocyte	14.7 days (0–58 days)	6.2 days (0–14 days)	0.002

Abbreviations: leukopenia, white blood cell count < 4000/mm³; neutropenia, absolute neutrophil count < 1500/mm³; lymphopenia, absolute lymphocyte count < 1500/mm³; ANA, anti-nuclear antibody; KD, Kikuchi disease.

time to recovery from lymphopenia and lymphocyte counts were evaluated for their association with KD recurrence. KD is mostly self-limiting, but some studies have reported that steroids or hydroxychloroquine may be helpful in shortening the clinical course or reducing the severity of disease, especially in recurrent or severe cases.^{23–26} Since these studies were mostly case reports, further prospective research is needed to define the role of steroids or hydroxychloroquine in the treatment of KD.

Many studies have described the association of KD with autoimmune conditions such as SLE based on immunological and clinicopathological findings.^{15–20} Thirteen (2.7% of all patients) patients developed systemic autoimmune diseases after KD during the observation period in this study. The most common type of autoimmune condition was SLE (10 patients). One patient developed Sjögren's syndrome, one developed rheumatoid arthritis, and one developed adult-onset Still's disease. There was no difference in the mean age

Table 3 Manifestations of Kikuchi Disease in Patients Who Developed Systemic Autoimmune Diseases

Characteristics	Cases Involving the Development of SADs	All KD Cases
Total	n=13 (100%)	n=480 (100%)
Mean age at the diagnosis of KD	27.9 yr (8–67 yr)	24.4 yr (4–79 yr)
Male: Female	1: 6	1: 2
Clinical features at the diagnosis of KD		
Fever	12 (92.3%)	381 (79.4%)
Lymphadenopathy	13 (100%)	455 (94.8%)
Extranodal symptoms	4 (30.8%)	61 (12.7%)
Recurrent KD	5 (38.5%)	54 (11.3%)
Laboratory findings		
Leukopenia	6 (46.2%)	201 (47.3% of 425)
Neutropenia	2 (15.3%)	43 (10.1% of 425)
Lymphopenia	5 (38.5%)	132 (31.1% of 425)
ANA (+) at KD diagnosis	8 (61.5%)	69 (28.2% of 245)

Abbreviations: SAD, systemic autoimmune disease; KD, Kikuchi disease; yr, years; leukopenia, white blood cell count < 4000/mm³; neutropenia, absolute neutrophil count < 1500/mm³; lymphopenia, absolute lymphocyte count < 1500/mm³; ANA, anti-nuclear antibody.

at the initial diagnosis of KD between all patients and those who developed systemic autoimmune diseases. However, the incidence rate of progression to autoimmune diseases after KD diagnosis was significantly higher in female patients with KD (Table 3). Patients showing progression to autoimmune diseases were more likely to have fever (92% vs 79.4%), common extranodal symptoms (30.8% vs 12.7%), a higher recurrence rate (38.5% vs 11.3%), and a higher ANA positivity rate at KD diagnosis (61.5% vs 28.2%) than all patients with KD. However, the group of patients who developed systemic autoimmune diseases after KD diagnosis was too small to yield statistically significant results. Despite this limitation, this study suggests potential predictive risk factors of progression to systemic autoimmune disease after KD diagnosis.

Another limitation of this study was that it was performed at a single center. However, electronic medical records were available for most clinical and laboratory findings of the patients. Further, a strength of this study is the consistent observation of a large number of patients over a long study period.

Conclusion

KD is generally a self-limiting disease; however, some patients have a more severe clinical course, such as recurrence or progression to systemic autoimmune diseases. Patients with extranodal symptoms, prolonged lymphopenia, KD recurrence, or ANA positivity benefit from early diagnosis and proper management of disease progression.

Acknowledgments

This work was supported by the new faculty research fund of Ajou University School of Medicine. We would like to thank Editage for English language editing.

Disclosure

The authors report no conflicts of interest related to this work.

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