

# A STUDY ON PREVALENCE AND CORRELATES OF DRY EYE IN CONNECTIVE TISSUE DISORDERS

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## ABSTRACT

### Objective:

To determine the prevalence of dry eye disease (DED) and its severity in patients with RA, SLE, and Primary Sjogren's syndrome in a tertiary care in Department of Ophthalmology Govt General Hospital, Suryapet, Telangana, India.

### Methods:

From July 2019 to May 2022, prospective cross-sectional research of 90 individuals with RA, SLE, and Sjogren's syndrome will be conducted. Schirmer's test, Tear breakup time, and ocular surface staining was used to assess patients for DED.

### Results:

The prevalence of DED was 51.11% for RA, 31.11% for SLE, and 17.77% for Primary Sjogren's syndrome. No cases of dry eye were found in 11.11% (10 cases), 38.88% (35) patients had mild dry eye, 22.22% (20) patients had moderate dry eye, 17.77% (16) patients had severe dry eye, and 10% (9) patients had very severe dry eye.

### Conclusions:

DED has a high overall frequency in connective tissue illnesses.

**Keywords:** dry eye disease, primary Sjogren's syndrome, tear breakup time, ocular surface staining.

## Introduction:

Keratoconjunctivitis sicca or Dry eye, is a multifactorial disease of the tears and ocular surface that results in symptoms of discomfort, visual disturbance, and tear film instability with potential damage to the ocular surface. Dry eye can be caused by a number of different factors. Inflammation of the ocular surface and an increase in the osmolality of the tear film are also symptoms that accompany this condition<sup>1</sup>. Patients suffering from connective tissue disorders are more likely to have dry eye, however the ailment is often misdiagnosed. There is a prevalence of dry eye ranging from 14.5 percent to 56 percent<sup>2</sup> in systemic autoimmune illnesses such as Sjogren's syndrome (SS), rheumatoid arthritis (RA), and systemic lupus erythematosus (SLE)<sup>2</sup>. Complaints of dry eyes are one of the most common symptoms of systemic disease, but patients very often choose to ignore them since other systemic manifestations are more bothersome. And as a result, they find themselves in the outpatient ophthalmology department with severe manifestations of dry eye or complications that have led to visual morbidity or irreparable damage to the ocular surface in late stages. If the condition is not properly investigated, it is possible that dry eye is the earliest manifestation of an underlying systemic disease that will not be detected for a considerable amount of time and the treatment focuses at minimizing or alleviating signs and symptoms of dry eye, such as ocular irritation, redness, or mucous discharge; therefore maintaining or delivering an improved visual function if it is not related to any other systemic condition<sup>3</sup>. In this way, it contributes to the prevention of injury to the ocular surface. In cases when there is significant dry eye as well as other

complaints that point towards a systemic link, further evaluation may be possible. Recent developments in our understanding of the factors that contribute to dry eye disease open the door to possibilities for enhancing diagnosis and disease management, as well as the search for new treatments that are more effective in managing this condition, which is both widespread and progressive. It is possible that the diagnosis and treatment of underlying systemic immunological diseases will reduce morbidity and in certain instances may even save a patient's life.

As a result, the goal of the study was to assess the patients with rheumatoid arthritis, systemic lupus erythematosus, and primary Sjogren's syndrome for the prevalence, manifestations, and grade of dry eye disease. This was done so that we could gain a better understanding of the trend and the necessity of ophthalmic examination in such patients. In order to increase the patient's comfort and to prevent or minimize additional structural damage to the ocular surface, it is important to recognize dry eye illnesses as early as possible<sup>4,5</sup>.

#### **Objectives of the Study:**

1. To determine the prevalence of dry eye disease in patients with connective tissue disorders viz.

Rheumatoid arthritis, Systemic lupus erythematosus and Primary Sjogren's syndrome.

2. To assess the severity of dry eye in each of the above-mentioned connective tissue disorders.

#### **Inclusion Criteria:**

- Diagnosed and confirmed cases of Rheumatoid arthritis, Systemic lupus erythematosus, and Primary Sjogren's syndrome.
- Age group between 20 to 65 years.

#### **Exclusion Criteria:**

- History of previous ocular surgeries (including cataract surgery) / ocular injuries
- Smoking history
- Radiation exposure
- Age <20 and >65 years
- Contact lens users.
- Episcleritis and Scleritis.

**Materials and Methods:**

The present prospective cross-sectional study was done at outpatient Department of Ophthalmology Govt General Hospital, Suryapet, Telangana, India with following inclusion and exclusion criteria on 90 patients diagnosed with Rheumatoid arthritis, systemic lupus erythematosus and primary Sjogren's syndrome and patients diagnosed and confirmed cases of RA, SLE and primary Sjogren's over period of one year from July 2019 to May 2022 in Govt Medical College/General Hospital, Department of Ophthalmology, Suryapet, Telangana, India. In this study, Schirmer's test Types I and II, tear meniscus height, tear BUT (TBUT), fluorescein stain, and rose Bengal stain were used to diagnose and grade dry eyes. Its severity is classified into mild, moderate, severe, and very severe (level 1 to 4) according to DEWS dry eye grading system. Mild dry-eyes are diagnosed by the presence of mild irritation, dryness with variable Schirmer's, and TBUT without any other abnormalities. Moderate-dry-eyes are diagnosed by Schirmer's  $\leq 10$ , TBUT  $\leq 10$  with a variable amount of corneal and conjunctival staining and visual symptoms. Severe-dry-eyes are diagnosed by Schirmer's and TBUT of

**Results:****Table 1: Diagnosis and Sex**

			Sex		Total
			Male	Female	
Diagnosis	Rheumatoid Arthritis	Count	18	28	46
		%	20%	31.11%	51.11%
	Systemic Lupus Erythematosus	Count	9	19	28
		%	10%	21.11%	31.11%
	Primary Sjogren's	Count	4	12	16
		%	4.44%	13.33%	17.77%
	TOTAL		31	59	90
	%		34.44%	65.55%	100%

The above table describes that in study population females were more predominant (65.55%) than males (34.44%) in each of diagnosis group and the incidence of RA was 31.11% (28 cases).

**Table 2: Age group and diagnosis**

			Age range					Total	
			20-29	30-39	40-49	50-59	>60		
Diagnosis	Rheumatoid Arthritis	Count	4	18	12	6	6	46	
		%	4.44%	20%	13.33%	6.66%	2.22%	51.11%	
	Systemic Lupus Erythematositis	Count	2	12	7	6	1	28	
		%	2.22%	13.33%	7.77%	6.66%	1.11%	31.11%	
	Primary Sjogrens	Count	2	9	2	2	1	16	
		%	2.22%	10%	2.22%	2.22%	1.11%	17.77%	
	Total			8	39	21	14	8	90
			%	8.88%	43.33%	23.33%	15.55%	8.88%	100%

The above table shows the distribution of different age groups in each of the diagnosis. The present study revealed that all types of diagnosis were predominantly in the age range of 30-39 yr with 43.33% (39 cases) of these RA prevalence was about 20% (18) patients.

**Table3:DEDSeveritygradingvs.Gender**

Dryeye disease severity		Sex		Total
		Male	Female	
Nodryeye	Count	2	8	10
	%	2.22%	8.88%	11.11%
Mildeyedry	Count	11	24	35
	%	12.22%	26.66%	38.88%
Moderateeyedry	Count	9	11	20
	%	10%	12.22%	22.22%
Severedryeye	Count	6	10	16
	%	6.66%	11.11%	17.77%
Veryseveredryeye	Count	3	6	9
	%	3.33	6.66	10%
Total		31	59	90
		34.44%	63.34%	100.0%

From the above table DED was more common in females given the known epidemiology of connectivetissue disorders being more common in females. There was a significant correlation between genderdistributionandseverityof dryeye disease.

**Table4:DEDseveritygradingvs. Agegroup**

Dryeyediseaseseverity		Agegroup					Total
		20-29	30-39	40-49	50-59	>60	
Nodryeye	Count	2	3	2	2	1	10
	%	2.22%	3.33%	2.22	2.22	1.11	11.11%
Mildeyedry	Count	3	14	8	6	4	35
	%	3.33%	15.55%	8.88	6.66	4.44	38.88%
Moderateyedry	Count	1	9	6	3	1	20
	%	1.11%	10%	6.66	3.33	1.11	22.22%
Severedryeye	Count	1	9	3	2	1	16
	%	1.11%	10%	3.33	2.22	1.11	17.77%
Veryseveredryeye	Count	1	4	2	1	1	9
	%	1.11%	4.44	2.22	1.11	1.11	10%
Total		8	39	21	14	8	90
Total%		8.88%	43.33%	23.33	15.55	8.88	100.0%

There was significant correlation between presence of DED or severity grading of it in 30-39 agegroup given the varied occurrence of RA, SLE and primary Sjogrens in that particular age groups. Present study describes 43.33% of dry eye disease being prevalent in mentioned connective tissue disorders and age group.

## Discussion:

According to the findings of this research, the overall prevalence of dry eye in people with RA, SLE, and Primary Sjogren's disease was determined to be 51.11%, 31.11%, 17.77% respectively. The severity of each condition was also graded, in addition to its prevalence, which was documented in each of the diseases. Out of total 90 individuals 11.11% (10) were found no dry eye, 38.88% (35) patients had mild dry eye, 22.22% (20) patients had moderate dry eye, 17.77% (16) patients had severe dry eye, and 10% (9) patients had very severe dry eye. There was a prevalence of dry eye in systemic autoimmune disorders such as rheumatoid arthritis (RA), systemic lupus erythematosus, and Sjogren's syndrome (SS) that ranged from 40% to 82%, according to the study by Yogeshwari et al., which is comparable to the current study, which states that it is 51.8 percent<sup>6</sup>. In a different study conducted by Khudair Al-Bedri et al.<sup>7</sup>, ocular signs of rheumatoid arthritis were investigated in 103 patients from Iran. Keratoconjunctivitis sicca was shown to be the most prevalent ocular finding, with a prevalence rate of 28 percent to 39 percent. This might be explained by the fact that their study had a bigger sample size, as well as by the fact that the patients came from a variety of ethnic backgrounds and environmental circumstances. In their investigation of ocular symptoms in SLE, Sukhum, Silpaarcha, Joan J Lee, and Stephen Foster discovered that one third of patients had keratoconjunctivitis sicca<sup>8</sup>. This finding further supports another study done by R. R. Sivaraj et al. stating the similar prevalence. It's possible that the smaller sample size contributed to our finding of a somewhat higher prevalence of 42 percent, which we found in our research<sup>9</sup>. A study conducted by Stuart S. Kassan et al. on clinical manifestations

and early diagnosis of Primary Sjogren's syndrome also showed a prevalence of 67.5 percent in their study<sup>10</sup>

8. This establishes dry eye as the common and highly prevalent disease in primary SS, which lends further support to the findings of this current study showing a prevalence of 66 percent of dry eye in primary Sjogren's disease. The data from this study also made it possible to classify dry eye severity by using a severity grading method developed by the Dry Eye International Workshop in 2007. This scheme covered both subjective symptoms and objective findings (clinical evaluation and diagnostic tests). Even in mild to moderate cases, dry eye can have a significant influence on a person's ability to participate in day-to-day activities, and this is something that we want to underline here.

Therefore, showing the importance of making this diagnosis in patients suffering from connective tissue disorders so that a therapeutic approach can be devised, which may improve the circumstances of the ocular surface, ultimately leading to a reduction in morbidity caused by dry eye disease<sup>11,12</sup>.

## Conclusion:

In conclusion our study revealed the predominance of DED in females at 30-39 age group which is a common undiagnosed ailment in general, and especially in connective tissue disorders such as rheumatoid arthritis, systemic lupus erythematosus, and primary Sjogren's syndrome. Patients often disregard the signs of dry eye, which can include irritation, a burning feeling, grittiness, and other ocular symptoms. This is because other systemic manifestations are more unpleasant for patients. And for this reason, patients visit an ophthalmologist when they experience considerable discomfort or recurring redness of the eyes. This occurs typically in the later phases of the condition, when ocular surface damage would have already happened, making the patient visually morbid. Because of this, conducting research on the prevalence of dry eye illness and determining the degree to which it is present in each of these disorders will assist us in alerting rheumatologists and physicians to the need for an early referral. If DED is diagnosed at an early stage and treatment is begun at that time, it will help maintain the ocular surface, which in turn will provide quality of life to patients by making their daily activities less hindering due to reduced ocular discomfort. Additionally, it will eventually increase workplace productivity and provide emotional support, both of which have been reported to be hampered due to dry eye diseases.

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