

Introduction

Dry eye disease is defined as a multifactorial disease of the ocular surface characterised by a loss of homeostasis of the tear film and accompanied by ocular symptoms in which tear film instability, hyperosmolarity, ocular surface inflammation, and neurosensory abnormalities play etiological roles.



- DED can influence individual's ability to perform daily tasks and can negatively affect their quality of life via many ways relating to decreased quality of vision and the development of psychological issues such as anxiety and depression.
- Prevalence of DED is higher in older age and in females as compared to males.

Patients & methods

This prospective comparative study was carried out at Al-zahraa University Hospital, Cairo, Egypt and included 120 eyes of 60 patients with severe dry eye disease, age 23 - 67 years



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Exclusion Criteria:

- Patients with positive serology for hepatitis B ,C, or HIV.
- Patients with hemolyzed or lipaemic serum.
- Contact lens wearers.
- Eyelid abnormality interfere with blinking
- Past history of herpetic keratitis

Our patients were divided into two groups according to treatment protocol:

- Group I: 30 patients (60 eyes) treated with autologous serum eye drops.
- **Group II:** 30 patients (60 eyes) treated with autologous platelet rich plasma eye drops.



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All patients underwent a comprehensive ophthalmic examination, including:

- Medical history, initial evaluation of the dry eye-related symptoms.
- BCVA.
- Slit-lamp examination (TBUT, corneal fluorescein staining, level of conjunctival hyperaemia).
- Schirmer's test (1).
- Conjunctival impression cytology (CIC).

The same tests as in the baseline visit were performed at each visit (2, 4, 6 weeks), except CIC which was only performed on the last visit.

Preparation Of Autologous Serum Eye Drops:

- The blood was first drawn from the recipient and allowed to clot in the absence of an anticoagulant.
- After clotting, the sample was centrifuged by the SW-12 centrifuge at 4000 RPM for 10 minutes at room temperature [20-40] C to separate serum from basal components with no haemolysis.
- After centrifugation, the serum was transferred into a sterile tube and diluted with a sterile saline solution to a 20% concentration, the final preparation was divided into 5-mL bottles.



Preparation Of Autologous Platelet Rich Plasma Eye Drops:

- The blood was first drawn from the recipient, placed in five vacutainer tubes [2-mL] containing anticoagulant, citrate dextrose solution, and centrifuged by the Hermle Z326K High-Speed Centrifuge [HERMLE Labortech-nik GmbH] at 1200 RPM for 10 minutes.
- The upper two layers, the plasma and buffy coat layer, were separated and diluted to 20% with a sterile saline solution, the final preparation was divided into 5-mL bottles.

- The patients in both groups were instructed to store these bottles at -20°C until use, a maximum for 3 months.
- The bottles being used were maintained under refrigerated conditions at 4°C, a maximum for 7 days.
- The patients were instructed to use the eye drops 4-6 times daily.





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Speed Centrifuge, sterile vacutainer tube



 Table (1): Comparison between Group 1 and Group 2 according to Age and sex.

		Group (1)	Group (2)	P-value
Age		50.53 ± 12.81	46.27 ± 12.93	0.204*
C	Male	e 9 (30.0%) 8 (26.7%)	0 77 4**	
Sex	Female	21 (70.0%)	22 (73.3%)	0.774**
* t-test. ** Chi-Sq	uare test.			E052023

Table (2): Mean Change 6 Weeks After Treatment In Group1 And Group 2

Variabla	Group (1)	Group (2)	p- value
variable	Mean± SD	Mean± SD	(Student's t-test)
BCVA (log MAR)			
Baseline	$0.15\pm\!\!0.22$	0.15±0.25	0.969
Change after Treatment	0.00	-0.05*	0.242
p- value (paired t-test)	1.00	0.001	
Break up time test			
Baseline	3.73 ± 0.86	4.02 ± 0.65	0.044
Change after Treatment	1.4*	1.69*	0.001
p- value (paired t-test)	<0.001	<0.001	

Variable	Group (1) Mean± SD	Group (2) Mean± SD	p- value (Student's t-test)
Schirmer's test			
Baseline	3.45±1.00	4.00±1.15	0.006
Change after Treatm	nent 1.23*	2.33*	<0.001
p- value (paired t-te	st) <0.001	<0.001	
Corneal fluorescein Stain	ing		
Baseline	2.23 ± 0.46	1.97 ± 0.45	0.002
Change after Treatm	-0.21*	-1.32*	<0.001
p- value (paired t-te	st) < 0.001	<0.001	
Level of hyperemia			
Baseline	1.82 ± 0.62	2.13 ± 0.5	0.002
Change after Treatm	nent -1.29*	-1.7*	0.275
p- value (paired t-te	st) < 0.001	<0.001	





CIC	Group (1) No, %	Group (2) No, %	p- value (Linear-by-linear association test)
aseline			
Grade 0	0 (0.0%)	0 (0.0%)	
Grade 1	9 (15%)	6 (10%)	1.00
Grade 2	24 (40%)	30 (50%)	1.00
Grade 3	27 (45%)	24 (40%)	
hange after treatment			
Grade 0	1 (1.7%)	2 (3.3)	
Grade 1	30 (50%)	40 (66.7)	0.04
Grade 2	29 (48.3)	18 (30%)	0.04
Grade 3	0 (0.0%)	0 (0.0%)	
P-value farginal homogeneity <u>test)</u>	<0.001*	<0.001*	











