A REAL-WORLD STUDY COMPARING ATROPINE MONOTHERAPY TO THE SYNERGISTIC EFFECTS OF COMBINATION TREATMENT: DEFOCUS INCORPORATED MULTIPLE SEGMENTS SPECTACLE LENSES AND LOW DOSE ATROPINE

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Background

- The increasing of global prevalence of myopia has bee concern. In myopia management there are many types of used as monotherapy or combination treatment.
- The effectiveness of Defocus Incorporated Multiple Seg spectacle lenses (as an optical intervention) has been demo as a stand-alone treatment and when combined (pharmacological treatment) in diverse populations.^{1,2}
- There is no literature on the effectiveness of combination to DIMS spectacle lenses and low dose atropine (LDA) in the S population.

Purpose

The objective of this study was to evaluate whether the comb and DIMS spectacle lenses was beneficial in South-Ame undergoing myopia control treatment.

Retrospective data for 51 patients attending a private January – September 2020.

- Inclusion criteria:
- Age 8-13y \bullet
- Myopia with SER between -5.00 to -1.00 D
- Myopia progression \geq -0.50 D/year in the previous 12 months
- Attended for 6-month, one- and two- year follow-up visits
- Phase 1 Environmental control: Following initial myo participants were advised to spend 2 hours a day in outdoors
- Phase 2 Monotherapy: Participants whose axial length (AL ≥0.15mm in 6 months were prescribed LDA for the next 12-m
- Phase 3 Combination treatment: If at the 12-month visit, th by \geq 0.17 mm/year, combination treatment (LDA + DIMS sp was prescribed for the next 12 months period.
- To observe changes, only data from the right eye was accepte
- Treatment effectiveness was calculated based on the myop across time periods.
- This study was approved by ethics committee of Centro L Várzea Grande, Brazil, under number 2127639 (December, 8^{TI}

We thank PhD. Gilmar Jorge de Oliveira junior for all the statistical support.

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lean age: 10.16 ± 1.63 years		Gender: Males 49%, Females 51%	
Table 1: General data			
Outcome	Phase	Mean ± SD	
Spherical Equivalent Refraction (D)	Baseline	-3.01 ± 1.22	
	Phase 1	-3.33 ± 1.22	
	Phase 2	-3.40 ± 1.21	
	Phase 3	-3.46 ± 1.23	
Axial Length (mm)	Baseline	24.60 ± 1.03	
	Phase 1	24.79 ± 1.03	
	Phase 2	24.99 ± 1.02	
	Phase 3	25.12 ± 1.03	
Keratometry (D)	Baseline	43.13 ± 1.19	
	Phase 1	43.12 ± 1.21	
	Phase 2	43.13 ± 1.24	
	Phase 3	43.17 ± 1.22	

Graph 1: Boxplot with the distribution in each phase of spherical equivalent refraction (A), axial length (B), variation axial length (C), and keratometry (D).







Discussion

- 0.01% ³.

- Attributed to participant selection

- Limitations:
 - Retrospective study
 - Small sample size
 - Short follow-up duration

Conclusions

- or environmental control in a Brazilian population.
- progression.

References

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LDA is the most widely used pharmaceutical intervention in clinical settings

• Several studies that showed weak effectiveness in long-term follow-up, particularly when AL elongation was the outcome of interest.^{4,5}

• Moreover, 0.05% atropine has been suggested as the most effective LDA tested in the young Asian population.⁶ In the western population, there were reports of frequent side effects when using this low-concentration.⁷

• In the present study, AL elongation was 0.13 ± 0.05 mm/year with combination treatment, compared to 0.21 \pm 0.03 mm/year from LDA alone, confirming the synergistic effect between LDA and DIMS.

 Although 0.025% atropine was used in combination with DIMS spectacle lenses in the present study, the results were indifferent to combination of 0.01% atropine + DIMS spectacle lenses in a European population.²

European study: participants selected from progressive myopes

Present study: sub-group of participants selected from progressive myopes in whom monotherapy (0.025% atropine) was not effective.

Combination treatment (DIMS spectacle lenses and 0.025% atropine) resulted in the most significant reductions in myopia progression based on AL elongation when compared to the use of 0.025% atropine monotherapy

Further randomized, double-blind clinical trials with longer follow-up may elucidate the true impact of this combination therapy on myopia

