



REVIEW

Under-correction or full correction of myopia? A meta-analysis



Negareh Yazdani^{a,b}, Ramin Sadeghi^c, Asieh Ehsaei^{a,b}, Ali Taghipour^d,
Samira Hasanzadeh^{a,b}, Leili Zarifmahmoudi^c, Javad Heravian Shandiz^{a,b,*}

^a Refractive Errors Research Center, School of Paramedical Sciences, Mashhad University of Medical Sciences, Mashhad, Iran

^b Department of Optometry, Mashhad University of Medical Sciences, Mashhad, Iran

^c Nuclear Medicine Research Center, Mashhad University of Medical Sciences, Mashhad, Iran

^d Health Sciences Research Centre, Department of Biostatistics and Epidemiology, School of Health, Mashhad University of Medical Sciences, Mashhad, Iran

KEYWORDS

Myopia;
Nearsightedness;
Under-correction;
Full correction

Abstract

Purpose: To compare the effect of full-correction versus under-correction on myopia progression.

Methods: A literature search was performed in PubMed, Scopus, Science Direct, Ovid, Web of Science and Cochrane library. Methodological quality assessment of the literature was evaluated according to the Critical Appraisal Skills Program. Statistical analysis was performed using Comprehensive Meta-Analysis (version 2, Biostat Inc., USA).

Results: The present meta-analysis included six studies (two randomized controlled trials [RCTs] and four non-RCTs) with 695 subjects (full-correction group, n = 371; under-correction group, n = 324) aged 6 to 33 years. Using cycloplegic refraction, the pooled difference in mean of myopia progression was -0.179 D [lower and higher limits: -0.383 , 0.025], which was higher but not in full correction group as compared to under correction group ($p = 0.085$). Regarding studies using non-cycloplegic subjective refraction according to maximum plus for maximum visual acuity, the pooled difference in myopia progression was 0.128 D [lower and higher limits: -0.057 , 0.312] higher in under-correction group compared with full-correction group ($p = 0.175$). Although, difference in myopia progression did not reach significant level in either cycloplegic or non-cycloplegic refraction.

Conclusions: Our findings suggest that, myopic eyes which are fully corrected with non-cycloplegic refraction with maximum plus sphere, are less prone to myopia progression, in comparison to those which were under corrected. However, regarding cycloplegic refraction, further studies are needed to better understand these trends.

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* Corresponding author at: Department of Optometry, School of Paramedical Science, Mashhad University of Medical Science, Azadi Square, Mashhad, Iran.

E-mail address: heravianSJ@mums.ac.ir (J. Heravian Shandiz).

Introduction

Myopia has already become the more prevalent refractive error worldwide, and its incidence is increasing particularly in young adults and school-aged children.^{1,2} For decades, researchers had typically focused on the genetic basis of the myopia. While, genetic factor alone cannot explain this dramatic increase. Increasing evidence suggests a role of environmental factors in etiology of the myopia or even myopia progression, such as outdoor activities, physical activities,³ socioeconomic factors and near work activities such as studying, reading, watching television, and using computer.⁴

There are some available options for myopia correction including spectacle, contact lens and refractive surgery. Spectacles are the most acceptable method in comparison with contact lens and refractive surgery, which is less invasive approach for correction.⁵ It has been approved that while myopia occurs early in life, it has a tendency toward progression due to related possible factors.⁶ Different approaches have been used in order to slow or reduce the progression of myopia, such as cycloplegic drugs (Atropine),⁷⁻⁹ specially designed contact lenses such as orthokeratology contact lenses,⁸⁻¹⁰ multifocal contact lenses,^{11,12} specially designed spectacles such as bifocal or multifocals^{13,14} and under-correction of the refractive error.^{6,9}

While there is no evidence for FDA (United State Food and Drug Association) approved pharmaceutical agents which slow the myopia progression, there are some studies concern the possible effect of some drugs on controlling myopia progression such as atropine.^{15,16} Contributed studies show that atropine is an effective method in slowing the myopia progression.¹⁷⁻¹⁹ However, in a study which considered the different concentrations of atropine, side effects were cautioned with 1% concentrate of atropine and also with long term usage.¹⁹ Chia et al. also compared three different concentrations of atropine and reported that 0.01% concentration has less side effects compared with 0.50 and 0.1%.²⁰ Although it has been shown that, lower dosages of atropine were found to have less rebound myopic progression.²¹

Special design of contact lenses such as orthokeratology, rigid contact lenses, and soft multifocal contact lenses,^{11,12} are alternative methods to control the progression of myopia.²² Orthokeratology contact lenses control myopia through reshaping and flattening the cornea during sleep, these lenses provide the clear vision in all day. It has been proved that, providing the successful fit, it will slower the progression of the myopia, particularly in low-to-moderate myopic patients.²³⁻²⁶ It has been shown that multifocal soft contact lenses reduce myopia progression through providing less peripheral refractive error.²⁷ Similar to contact lenses, bifocal or multifocal spectacles also control myopia progression through reducing accommodative effort at near vision.¹³

Under-correction is one of the interventions to reduce or slow myopia progression. One hypothesis about under-correction is that it may decrease the accommodative effort and also the lag of accommodation and subsequently slow or reduce the myopia progression.^{6,28,29} Some studies have already compared under-correction versus full correction in myopic population.^{6,30-33} Whereas, there is contradiction in

study's findings with regards to the under-correction or full correction of myopia, it seems valuable to have a definite conclusion and provide clinically an appropriate and effective method of correction. Therefore, we aimed to conduct a meta-analysis to investigate whether the under-correction is the method of choice for myopic patients or the full correction is the better way.

Materials and methods

Current meta-analysis was performed based on PRISMA statement, the Preferred Reporting Items for Systematic Reviews and Meta-Analysis. The PRISMA Statement consists of a 27-item checklist and a four-phase flow diagram. The aim of the PRISMA Statement is to help authors improve the reporting of systematic reviews and meta-analyses. PRISMA is used as a basis for reporting systematic reviews, particularly evaluations of interventions.³⁴

Search strategy

Complete literature searches of the PubMed, Science direct, Ovid, Web of Science, Scopus and Cochrane library databases was done to find relevant published articles on comparing under-correction with full-correction of myopia. We used a search algorithm that was based on a combination of the terms: "Under-correction, Full-correction, Myopia and Nearsightedness". The Boolean operators "AND" and "OR" were used in order to specify the search. No beginning date limit or language restriction was used; the search was updated until July 2019. To expand our search, references of the retrieved articles were also screened by two independent authors (NY and SH) for more related studies.

Study selection

Studies concerning the difference between under-correction and full-correction of myopia were considered for inclusion. The exclusion criteria were shown in Flow chart. (Fig. 1) Two researchers (NY and SH) independently reviewed the titles and abstracts of the retrieved articles, applying mentioned inclusion and exclusion criteria and also reviewed the full-text version of the remaining articles to verify their suitability for inclusion.

Data extraction

For each study, information was gathered regarding basic study data (authors, journals, year of publication, country of origin, and study design), patient characteristics (mean age, gender, number of patients) and technical aspects (amount of under-correction, type of refraction and applied device). (Table 1).

Quality assessment

Critical Appraisal Skills Program (CASP), was used to assess the methodological quality of included studies based on study design. This tool presents a number of questions that

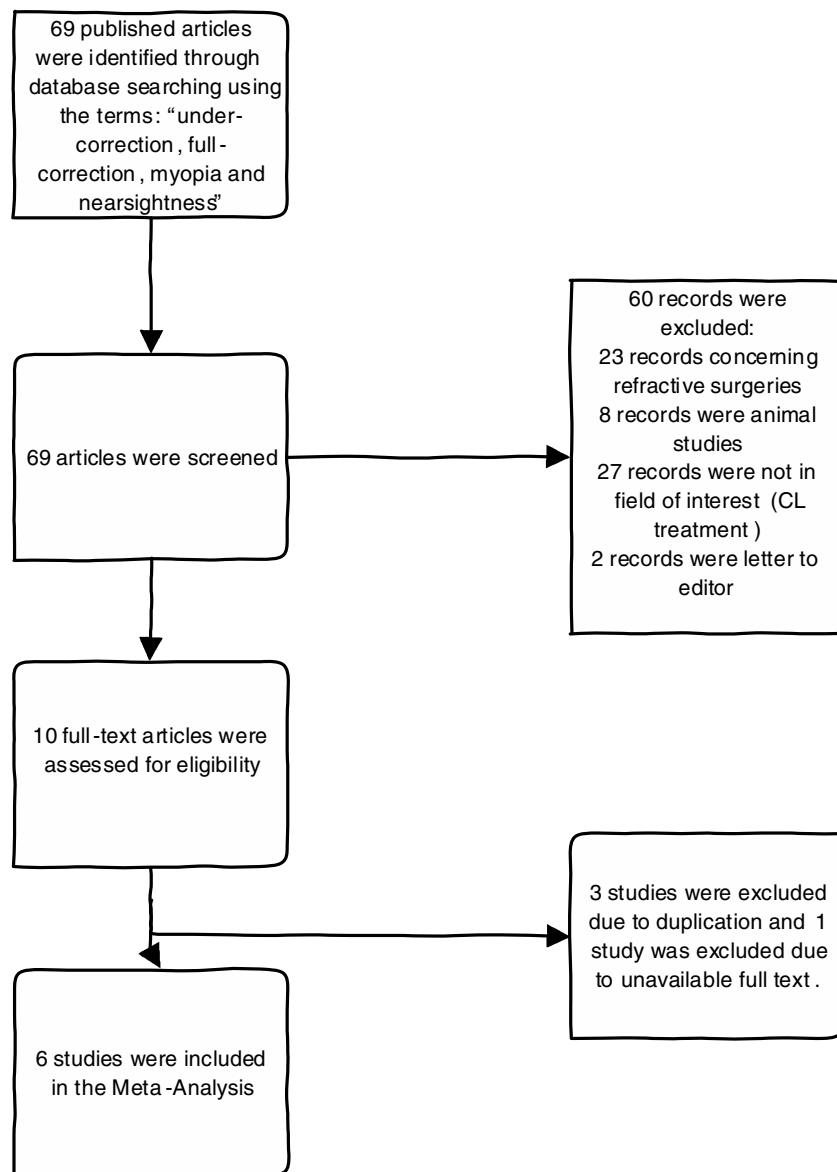


Figure 1 Flow chart of included studies.

deal very broadly with some of the principles or assumptions that characterize the qualitative aspect of a research. Three broad issues are considered when appraising a qualitative study: 1. Are the results of the study valid? 2. What are the results? 3. Will the results help locally? You are asked to record a "yes", "no" or "can't tell" to most of the questions. (CASP; <http://www.casp-uk.net/>) CASP cross-sectional tool assess quality in five aspects: selection bias which assess whether is the sample representative of a defined population or is everybody included who should have been included, measurement bias which assess if measurements truly reflect what you want, data collection clarifies how data were collected, sample size to assess whether study have enough participants to minimize role of chance and results presentation which assess the main result and if it could be applied to local population. CASP randomized control trials tool assess quality in six aspects: sample allocation which assess whether participants were appropriately

allocated to intervention and control group, blinding which assess whether participants, staff and study personnel were blind to participants, informed about the study which assess whether all of the participants who entered the trial were accounted for at its conclusion, similarity in method which assess if all participants were examined in the same way, sample size which clarify whether enough participants were enrolled to minimize role of chance and results presentation assess how the results were presented and what is main result. CASP cohort tool assess quality in four aspects: selection bias, which assess whether cohort recruited in acceptable way and if appropriate sample were included, measurement bias which assess whether exposure was accurately measured to minimize bias, considering confounding factors which assess whether authors identified all confounding factors and acceptable follow up which assess whether follow up was complete enough. (Table 2).

Table 1 Characteristics of included studies.

Author	Title	Sample size	Age	Refraction	Device	Amount of U	Result
Daniel Adler (2006) ⁶	The possible effect of under-correction on myopic progression in children.	48 (F:23, U:25)	6–15 y	Non-cycloplegic	Retinoscopy	–0.50 D	Under-correction produced a slight but not statistically significant increase in myopic progression compared to full correction.
Kahmeng Chung (2002) ³⁰	Under-correction of myopia enhances rather than inhibits myopia progression.	94 (F:47, U:47)	9–14 y	Non-cycloplegic	Retinoscopy	–0.75 D	Under-correction produced more rapid myopia progression.
Si yuan Li (2015) ³¹	Effect of under-correction on myopia progression in 12 year old children.	253 (F:133,U:120)	12 y	Cycloplegic	Autorefractometer	–0.50 ≤ D	Under-correction or full-correction of myopia by wearing spectacles did not show any differences in myopia progression or axial elongation.
Balamurali Vasudevan (2014) ³³	Under-correction of human myopia- is it myopigenic?	79 (F:35, U:12)	11–33 y	Non-cycloplegic	Autorefractometer	–0.50 D	Under-correction of myopia produced a small but progressively greater degree of myopic progression than did full correction.
Yun-Yun Sun (2017) ³⁷	Effect of under-correction versus full-correction on myopia progression in 12 year-old children.	121 (F:56, U:65)	12.7 y	Cycloplegic	Autorefractometer	–0.50 D	Myopia progression decreased with under-correction.
Yao-Hua Chen (2014) ³⁸	Clinical observation of the development of juvenile myopia wearing glasses with full-correction and under-correction.	132 (F:77, U:55)	12–18 y	Non-cycloplegic	Retinoscopy	–0.50 D	The progression of myopia is slow if patients wear glasses with full correction.

F: Full correction, U: Under-correction. UK: United Kingdom, USA: United State of America, NA: Not available, Y: Years.

Table 2 Quality assessment of included studies. (CASP).

Cross-sectional studies						
Study	Selection Bias	Measurement Bias	Data collection	Sample size	Results presentation	
Daniel Adler (2006) ⁶	Yes	Yes	Yes	Cannot tell	Yes	
Balamurali Vasudevan (2014) ³³	Yes	Yes	Yes	Cannot tell	Yes	
Randomized control trial studies						
Study	Sample allocation	Blinding	Informed about the study	Similarity in method	Sample size	Results presentation
Kahmeng Chung (2002) ³⁰	Yes	Yes	Yes	Yes	Cannot tell	Yes
Yao-Hua Chen (2014) ³⁸	Yes	Yes	Cannot tell	Yes	Yes	Yes
Cohort studies						
Study	Selection bias	Measurement Bias	Considering Confounding factors	Acceptable follow up		
Si yuan Li (2015) ³¹	Yes	Yes	Yes	Yes		
Yun-Yun Sun (2017) ³⁷	Yes	Yes	Yes	Yes		

Statistical analysis

For each study, the mean difference in myopia progression recorded in diopter notation was determined for the full-correction and under-correction groups. To pool the effect sizes across studies, a random effects model was used. The random effects model is a statistical method in which between study variability is accounted for. This method is used especially in pooling data across studies which are different in terms of design, included patients etc. In the forest plot, left column shows the identity (type of applied refraction whether it is cycloplegic or non-cycloplegic) of each included study, and is followed by name of the first author. Next, to the right, we meet some statistic data for each study which includes difference in means of myopia progression in diopter between two approaches of under or full correction, lower and upper limits of confidence interval and p -value. The right-hand column visually displays each study results. The horizontal lines through the boxes illustrate the length of the confidence interval. The longer the lines, the wider the confidential interval, the less reliable the study results. The width of the diamond serves the same purpose. The vertical line is the line of no effect. The boxes show the effect estimates from the single studies, while the diamond shows the pooled result. The larger the box, the bigger the sample size and the narrower the confidence interval (CI), the higher the percentage weight and more the influence the study has on the pooled result. If the diamond shape does not touch the line of no effect, the difference found between the two

groups was statistically significant. In that case, the p -value is usually <0.05 .

Heterogeneity was evaluated by the Cochrane Q test (the significance level was considered to be 0.05.), and I^2 index.³⁵ I^2 index is the inconsistency index and represents how much of the heterogeneity among the included studies is real and cannot be attributed to sampling error. Publication bias was evaluated graphically by funnel plots and statistically by Egger's regression intercept method.³⁶ The Funnel plot is the plot of the standard errors of the included studies on the y-axis and the effect size on the x-axis. Each dot represents a single study. The y-axis is usually the standard error of the effect estimate. Larger studies with higher power are placed toward the top. Lower powered studies are placed toward the bottom. Asymmetry of this plot may be due to publication bias. Egger's regression intercept is the mathematical counterpart of this visual assessment. Statistically significant results of this test indicate a large asymmetry in the funnel plot. All statistical analyses were performed using Comprehensive Meta-Analysis (version 2, Biostat Inc., USA).

Results

PRISMA flow chart (PRISMA 2009 flow diagram) of the study has been presented in Fig. 1. Overall six studies were included in this meta-analysis. Two studies with Randomized Controlled Trial (RCT) design, two studies with Cross-sectional and two Cohort studies. (Table 1). Among

included studies, 2 studies, used cycloplegic auto refraction (group1) and 4 studies, used non-cycloplegic refraction with retinoscopy and auto refractometer, which were followed by refined subjective refraction based on maximum plus sphere as the end point for best corrected distance visual acuity (group2).

The pooled difference in mean of myopia progression using cycloplegic refraction was -0.179 D [lower and higher limits: $-0.383, 0.025$], which was higher in the full-correction group as compared to the under-correction group ($p=0.085$). In group 2, the pooled difference in myopia progression was 0.128 D [lower and higher limits: $-0.057, 0.312$] higher in under-correction group compared with full-correction group ($p=0.175$). Fig. 2 shows the forest plot of the myopia progression. The Cochrane Q value was 6.99 ($p=0.008$) and $I^2=85.69$ for cycloplegic studies and Q value was 1.48 ($p=0.686$) and $I^2=0.000$ for non-cycloplegic studies. Fig. 3 shows the funnel plot of the included studies. Estimated Eggers regression intercept was 3.70 , $p=0.33$, which indicates a symmetrical funnel plot.

The subgroup analysis of the results of the studies in group 1 and 2 demonstrated pooled progression rates of -0.26 diopter (95% CI -0.24 to -0.29) in group 1, in comparison with 0.15 diopter (95% CI 0.10 to 0.21) in group 2. In the first group of studies, myopia progression was significantly higher in full-corrected subjects in relation to under-corrected cases. ($p<0.001$) While, in the group 2, myopia progression decreased significantly with full-correction ($p<0.001$). Although, difference in means of progression was statistically significant in both types of refraction, but the difference was not clinically significant.

Discussion

Today myopia is the main global cause of visual impairment, especially in Asia.^{39,40} Rapid increase of myopia prevalence has become a considerable global concern for both visual science researchers and practitioners⁴¹ and assessment of approaches to slow down the myopia progression are the main topics of recent studies. Different approaches, such as single-focus spectacles, contact lenses and atropine eye drops, have been employed in an attempt to reduce myopia progression.⁴²⁻⁴⁶ Spectacles are still the most common and easiest option. The corrected amount of myopia (full-correction or under-correction) has been the topics of several studies and discussed in several literatures.^{6,30,31,33,37,38} Under-correction has been believed that can reduce the accommodative stimulus and demand at near,⁴⁷ and subsequently reduce the blur drive for accommodation, which may be a myopigenic factor.⁴⁸ Moreover, animal studies have also revealed that myopic defocus resulting from under correction or lack of correction could slow myopia.⁴⁹⁻⁵¹ A controversial hypothesis regarding under-correction is that defocused retinal image might cause myopia progression due to form deprivation myopia.⁵² On the other hand, it has been hypothesized that full-correction of myopia optically position the far point of the eye close to the infinity and thus reducing any retinal defocus-induced, blur signals at distance.⁵³ Several studies have compared the effect of under-correction and

full-correction of myopia on its progression, but there are discrepancies in findings. Some studies showed that under-correction slows down the myopia progression,^{31,37} however, other studies revealed that full-correction could be effective in reducing myopia progression.^{6,30,33,38}

This systematic review and meta-analysis showed that the rate of myopia progression significantly differs between under-correction and full-correction approaches. The difference was statistically significant but clinically did not reach the significance level. According to the findings in this meta-analysis, in non-cycloplegic sub-group (retinoscopy followed by maximum plus to the best visual acuity), under-correction caused significantly greater degree of myopia progression as compared to full-correction. The possible rational belief behind it, is that full-correction could prevent any induced defocus of retinal image⁵³ and form deprivation,⁵⁴ which is believed to be an effective factor in myopia progression or incidence. The present findings are in consistence with Adler,⁶ Chung,³⁰ Chen³⁸ and Vasuadevan³³ studies, which reported that full-correction could significantly reduce the myopia progression compared with under-correction group. Besides, eyes with under-correction are continuously stimulated by blurred image which could causes greater rate of progression.³⁰ Regarding studies with cycloplegic refraction, under-correction could significantly reduce myopia progression compared with full-correction. The possible reason is that under-correction reduce the accommodative stimulus and demand at near,⁴⁸ and thus reduce the blur drive for accommodation, which is thought to be a myopigenic factor.

The reason that inference on the basis of between sub-groups differences is that there may be another factor, aside from types of refractions, which could be possible reason that could explain the difference between cycloplegic and non-cycloplegic sub-groups. Amount of under-correction may be more than what is really applied in cycloplegic group, as it has been reported myopia progression significantly decreased with increasing amount of under-correction.³¹ Besides, it has been approved as if under-correction continues, then myopia progression shows higher degree of reduction.⁵⁵

Moreover, our findings show that, the results of two studies with cycloplegic refraction^{31,37} could not be comparable with other four non-cycloplegic studies.^{6,30,33,38} The reason may be that Li³¹ and Sun,³⁷ in their studies, have applied different amount of under-correction, and just in one study³¹ the exact rate of progression was reported based on each amount of under-correction, which was a limitation in Sun study. Sun et al.³⁷ have just reported the myopia progression for no correction group and compared it with full-correction. Another reason concerns the definition of under-correction could be the cause of error. In Li³¹ study, under-correction of myopia was defined as improved presenting visual acuity for at least 2 lines with subjective refraction. Using this criterion of under-correction of myopia may not have detected all under-corrected children because myopes may possess reduced blur sensitivity. Lastly, further randomized control trials investigations are needed using cycloplegic refraction, which in essence serve as the gold standard for precise refraction, to understand better the relation between amount of correction and myopia progression.

This study has also some limitations. Possible heterogeneity has already been reported in all meta-analysis,

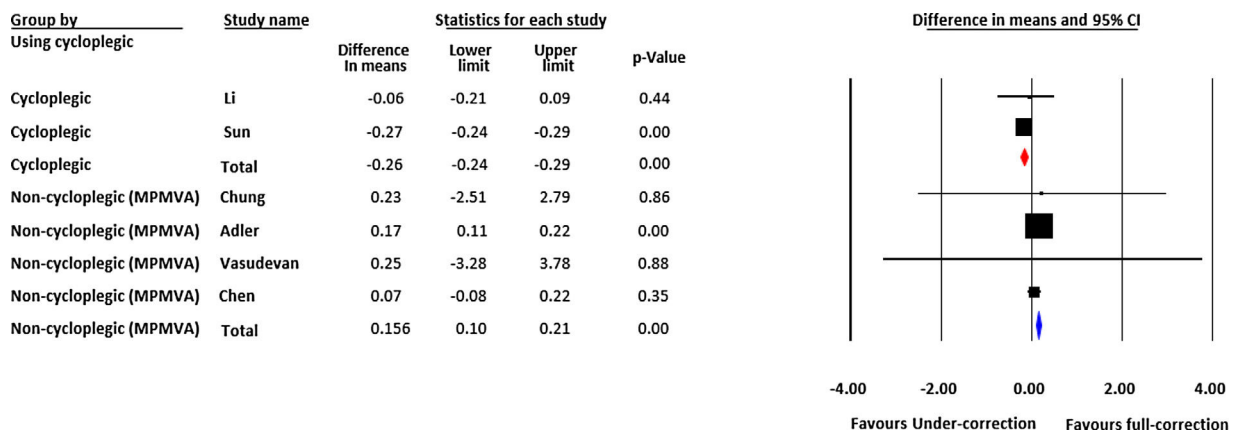


Figure 2 Forest-plot of subgroup analysis based on refraction. Left column shows the identity of each included study and name of the first author. Statistics for each study part presents difference in means of myopia progression, confidence interval and p-value. In left hand column, boxes show the effect estimates from the single studies, while the diamond shows the pooled result. MPMVA = Maximum Plus to Maximum Visual Acuity.

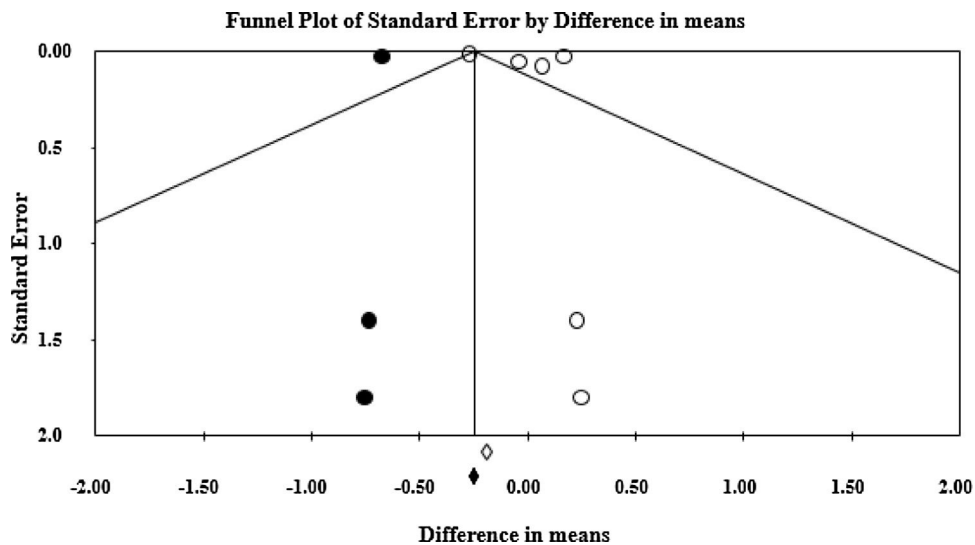


Figure 3 Funnel plot of meta-analysis. Each dot represents a single study. The y-axis is shows the standard error of the effect estimate. The x-axis shows “Difference in means”.

especially those which evaluate the pooled estimate. Our results showed the high heterogeneity which is likely to arise through differences in inclusion criteria, amount of under-correction and types of refraction. Also, the studies quality might affect the heterogeneity of the results. Therefore, any conclusion was limited by the characteristics of the studies. Otherwise, heterogeneity can be considered as an important drawback in meta-analysis to find the variables associated with variation across the primary studies can help future research on a topic. Publication bias is a major concern in all systematic reviews. The only method in order to avoid publication bias is searching a broad number of databases to locate all possible studies. We did our best to increase the sensitivity of our search strategy. However, publication bias is a major concern in all systematic reviews and cannot be avoided altogether. Citation bias is always a concern in systematic reviews. However, we only used citation analysis as an adjunct to the main search strategy.

Conclusion

The myopic eyes which are fully corrected with non-cycloplegic refraction with maximum plus sphere, are less prone to myopia progression, in comparison with those which were under corrected. Although, our results show statistically significant difference in myopia progression comparing under-correction and full-correction, in either cyclo or non-cyclo groups, this difference might not be clinically considerable. Further studies are warranted to provide reliable evidences in this regard, particularly with cycloplegic refraction.

References

1. Pan CW, Dirani M, Cheng CY, et al. The age-specific prevalence of myopia in Asia: a meta-analysis. *Optom Vis Sci.* 2015;92:258–266.

2. Guo L, Yang J, Mai J, et al. Prevalence and associated factors of myopia among primary and middle school-aged students: a school-based study in Guangzhou. *Eye*. 2016;30:796–804.
3. Parssinen O, Era P, AL L. Some physiological and psychological characteristics of myopic and non-myopic young men. *Acta Ophthalmol Suppl*. 1985;173:85–87.
4. Saw SM, Chua WH, Hong CY, et al. Nearwork in early-onset myopia. *Invest Ophthalmol Vis Sci*. 2002;43:332–339.
5. Nio YK, Jansonius NM, Wijdh RH, et al. Effect of methods of myopia correction on visual acuity, contrast sensitivity, and depth of focus. *J Cataract Refract Surg*. 2003;29:2082–2095.
6. Adler D, Millodot M. The possible effect of undercorrection on myopic progression in children. *Clin Exp Optom*. 2006;89:315–321.
7. Shih KC, Chan TC, Ng AL, et al. Use of atropine for prevention of childhood myopia progression in clinical practice. *Eye Contact Lens*. 2016;42:16–23.
8. Huang J, Wen D, Wang Q, et al. Efficacy comparison of 16 interventions for myopia control in children: a network meta-analysis. *Ophthalmology*. 2016;123:697–708.
9. Tay S, Farzavandi S, Tan D. Interventions to reduce myopia progression in children. *Strabismus*. 2017;25:23–32.
10. Kang P, Swarbrick H. New perspective on myopia control with orthokeratology. *Optom Vis Sci*. 2016;93:497–503.
11. Sha J, Tilia D, Diec J, et al. Visual performance of myopia control soft contact lenses in non-presbyopic myopes. *Clin Optom*. 2018;10:75–86.
12. Walline JJ, Gaume Giannoni A, Sinnott LT, et al. A randomized trial of soft multifocal contact lenses for myopia control: baseline data and methods. *Optom Vis Sci*. 2017;94:856–866.
13. Smith MJ, Walline JJ. Controlling myopia progression in children and adolescents. *Adolesc Health Med Ther*. 2015;6:133–140.
14. Cheng D, Woo GC, Drobe B, Schmid KL. Effect of bifocal and prismatic bifocal spectacles on myopia progression in children: three-year results of a randomized clinical trial. *JAMA Ophthalmol*. 2014;132:258–264.
15. Fan DS, Lam DS, Chan CK, et al. Topical atropine in retarding myopic progression and axial length growth in children with moderate to severe myopia: a pilot study. *Jpn J Ophthalmol*. 2007;51:27–33.
16. Tong L, Huang XL, Koh AL, et al. Atropine for the treatment of childhood myopia: effect on myopia progression after cessation of atropine. *Ophthalmology*. 2009;116:572–579.
17. Chia A, Lu QS, Tan D. Five-year clinical trial on atropine for the treatment of myopia 2: myopia control with atropine 0.01% eyedrops. *Ophthalmology*. 2016;123:391–399.
18. Lee JJ, Fang PC, Yang IH, et al. Prevention of myopia progression with 0.05% atropine solution. *J Ocul Pharmacol Ther*. 2006;22:41–46.
19. Shih YF, Chen CH, Chou AC, et al. Effects of different concentrations of atropine on controlling myopia in myopic children. *J Ocul Pharmacol Ther*. 1999;15:85–90.
20. Chia A, Chua WH, Cheung YB, et al. Atropine for the treatment of childhood myopia: safety and efficacy of 0.5%, 0.1%, and 0.01% doses (atropine for the treatment of myopia 2). *Ophthalmology*. 2012;119:347–354.
21. Pineles SL, Kraker RT, VanderVeen DK, et al. Atropine for the prevention of myopia progression in children: a report by the American Academy of Ophthalmology. *Ophthalmology*. 2017;124:1857–1866.
22. Cho P, Cheung SW. Retardation of myopia in Orthokeratology (ROMIO) study: a 2-year randomized clinical trial. *Invest Ophthalmol Vis Sci*. 2012;53:7077–7085.
23. Cho P, Cheung SW, Edwards M. The longitudinal orthokeratology research in children (LORIC) in Hong Kong: a pilot study on refractive changes and myopic control. *Curr Eye Res*. 2005;30:71–80.
24. Walline JJ, Jones LA, Sinnott LT. Corneal reshaping and myopia progression. *Br J Ophthalmol*. 2009;93:1181–1185.
25. Kakita T, Hiraoka T, Oshika T. Influence of overnight orthokeratology on axial elongation in childhood myopia. *Invest Ophthalmol Vis Sci*. 2011;52:2170–2174.
26. Santodomingo-Rubido J, Villa-Collar C, Gilmartin B, Gutierrez-Ortega R. Myopia control with orthokeratology contact lenses in Spain: refractive and biometric changes. *Invest Ophthalmol Vis Sci*. 2012;53:5060–5065.
27. Sankaridurg P, Holden B, Smith E, et al. Decrease in rate of myopia progression with a contact lens designed to reduce relative peripheral hyperopia: one-year results. *Invest Ophthalmol Vis Sci*. 2011;52:9362–9367.
28. Mutti DO, Mitchell GL, Hayes JR, et al. Accommodative lag before and after the onset of myopia. *Invest Ophthalmol Vis Sci*. 2006;47:837–846.
29. Nakatsuka C, Hasebe S, Nonaka F, Ohtsuki H. Accommodative lag under habitual seeing conditions: comparison between myopic and emmetropic children. *Jpn J ophthalmol*. 2005;49:189–194.
30. Chung K, Mohidin N, O’Leary DJ. Undercorrection of myopia enhances rather than inhibits myopia progression. *Vis Res*. 2002;42:2555–2559.
31. Li SY, Li SM, Zhou YH, et al. Effect of undercorrection on myopia progression in 12-year-old children. *Graefes Arch Clin Exp Ophthalmol = Albrecht von Graefes Archiv fur klinische und experimentelle Ophthalmologie*. 2015;253:1363–1368.
32. Li SM, Li SY, Liu LR, et al. Full correction and undercorrection of myopia evaluation trial: design and baseline data of a randomized, controlled, double-blind trial. *Clin Exp Ophthalmol*. 2013;41:329–338.
33. Vasudevan B, Esposito C, Peterson C, et al. Under-correction of human myopia—is it myopigenic? A retrospective analysis of clinical refraction data. *J Optom*. 2014;7:147–152.
34. Moher D, Liberati A, Tetzlaff J, Altman DG. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLoS Med*. 2009;6:e1000097.
35. Huedo-Medina TB, Sanchez-Meca J, Marin-Martinez F, Botella J. Assessing heterogeneity in meta-analysis: Q statistic or I2 index? *Psychol Methods*. 2006;11:193–206.
36. Song F, Khan KS, Dinnes J, Sutton AJ. Asymmetric funnel plots and publication bias in meta-analyses of diagnostic accuracy. *Int J Epidemiol*. 2002;31:88–95.
37. Sun YY, Li SM, Li SY, et al. Effect of uncorrection versus full correction on myopia progression in 12-year-old children. *Graefes Arch Clin Exp Ophthalmol = Albrecht von Graefes Archiv fur klinische und experimentelle Ophthalmologie*. 2017;255:189–195.
38. Chen Y-H. Clinical observation of the development of juvenile myopia wearing glasses with full correction and under-correction. *Int Eye Sci*. 2014;14:1553–1554.
39. Pan CW, Ramamurthy D, Saw SM. Worldwide prevalence and risk factors for myopia. *Ophthalmic Physiol Opt*. 2012;32:3–16.
40. Morgan IG, Ohno-Matsui K, Saw SM. Myopia. *Lancet (London, England)*. 2012;379:1739–1748.
41. Lim MC, Gazzard G, Sim EL, Tong L, Saw SM. Direct costs of myopia in Singapore. *Eye (London, England)*. 2009;23:1086–1089.
42. Walline JJ, Lindsley K, Vedula SS, et al. Interventions to slow progression of myopia in children. *Cochrane Database Syst Rev*. 2011;Cd004916.
43. Saw SM, Gazzard G, Au Eong KG, Tan DT. Myopia: attempts to arrest progression. *Br J Ophthalmol*. 2002;86:1306–1311.
44. Li SM, Ji YZ, Wu SS, et al. Multifocal versus single vision lenses intervention to slow progression of myopia in school-age children: a meta-analysis. *Surv Ophthalmol*. 2011;56:451–460.

45. Li SM, Wu SS, Kang MT, et al. Atropine slows myopia progression more in Asian than white children by meta-analysis. *Optom Vis Sci.* 2014;91:342–350.
46. Li SM, Kang MT, Wu SS, et al. Efficacy, safety and acceptability of orthokeratology on slowing axial elongation in myopic children by meta-analysis. *Curr Eye Res.* 2016;41:600–608.
47. Curtin JB. *The Myopias: Basic Science and Clinical Management.* Philadelphia: Harper and Row; 1985.
48. Rosenfield M, Gilmartin B. *Myopia and Nearwork.* Boston: Butterworth-Heinemann; 1998.
49. Arumugam B, Hung LF, To CH, et al. The effects of simultaneous dual focus lenses on refractive development in infant monkeys. *Invest Ophthalmol Vis Sci.* 2014;55:7423–7432.
50. Zhu X, McBrien NA, Smith EL, et al. Eyes in various species can shorten to compensate for myopic defocus. *Invest Ophthalmol Vis Sci.* 2013;54:2634–2644.
51. Tse DY, To CH. Graded competing regional myopic and hyperopic defocus produce summated emmetropization set points in chick. *Invest Ophthalmol Vis Sci.* 2011;52:8056–8062.
52. Wallman J, Turkel J, Trachtman J. Extreme myopia produced by modest change in early visual experience. *Science (New York, NY).* 1978;201:1249–1251.
53. Ciuffreda KJ, Vasudevan B. Effect of nearwork-induced transient myopia on distance retinal defocus patterns. *Optometry (St Louis, Mo).* 2010;81:153–156.
54. Smith EL, Hung LF. Form-deprivation myopia in monkeys is a graded phenomenon. *Vision Res.* 2000;40:371–381.
55. Ong E, Grice K, Held R, Thorn F, Gwiazda J. Effects of spectacle intervention on the progression of myopia in children. *Optom Vis Sci.* 1999;76:363–369.