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# Efficacy of amblyopia therapy initiated after 9 years of age

#### Abstract

Aims/purpose To evaluate the efficacy of occlusion therapy initiated after 9 years of age. Methods A total of 16 amblyopes of 9 years or older (range, 9.0-14.5 years; mean, 10.5 years) with a difference in visual acuity of over two lines between the eyes alter 4 weeks of first full-time spectacle wear were included. None of the children had undergone a previous ocular examination, had ever worn spectacles, received occlusion therapy, or had strabismus surgery. Initial worst visual acuity after 4 weeks of full-time spectacle wear was 20/100 in three patients, between 20/80 and 20/40 in 11 patients, and 20/30 in two patients. Full-time occlusion was performed in 14 patients and part-time occlusion in two patients. Results The final visual acuity of 15 out of 16 patients (94%) improved at least two lines. The final visual acuities ranged from 20/30 to 20/20 in 14 patients, 20/40 in one patient, and 20/50 in one remaining patient who began amblyopia therapy at 14.5 years of age, with the poorest compliance among the patients. Conclusions Occlusion therapy for anisometropic and strabismic amblyopia can

be successful even if initiated after the age of 9 years.

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*Keywords:* amblyopia; 9 years or older; occlusion; efficacy

## Introduction

Amblyopia is a reduced corrected visual acuity without any evidence of organic eye disease. It is caused by an abnormal visual experience during a sensitive period of visual development in early childhood most commonly from either a strabismus, an anisometropia, or both. It has been treated by correcting the refractive errors followed by occlusion of the fellow eye.<sup>1</sup> Many factors are related to the response to amblyopia therapy. These are the initial visual acuity of the amblyopia,<sup>2–5</sup> the age of the initial therapy,<sup>3–9</sup> the type of amblyogenic stimulus (anisometropia, strabismus, or deprivation),<sup>2–5,9</sup> the duration of amblyopia therapy,<sup>5</sup> the method of amblyopia therapy,<sup>10,11</sup> and the level of compliance.<sup>4–6,12</sup> Among them, the single factor that was most clearly related to a successful outcome was the age at commencement of therapy.<sup>3</sup> Regarding this age factor, there have been several reports with conflicting results.<sup>3-9</sup> The purpose of this study was to evaluate the efficacy of amblyopia therapy initiated after 9 years of age, one of the oldest patient groups ever reported.

## Patients and methods

The study was conducted from January 1995 to January 2000. All patients with amblyopia 9 years of age or older who were eligible and whose parents agreed to enrolment were included. All of the amblyopes with a difference in visual acuity of over two lines between the eyes after 4 weeks of full-time spectacle wear were included in this study. None of the children had undergone a previous ocular examination. Therefore, none had ever worn spectacles, received amblyopia therapy, or had strabismus surgery. Patients with previous histories of retinal or optic nerve disease, glaucoma, cataract, nystagmus, media opacities, ocular trauma, eye surgery, neurologic disease, cerebral palsy, and mental retardation were excluded. Informed consent was obtained from each patient.

A detailed ophthalmic examination, including an uncorrected visual acuity, bestcorrected visual acuity, slit-lamp examination, cycloplegic refraction, prism cover test, alternate prism cover and uncover test, examination of ductions and versions, pupillary examination, and fundus examination was Department of Ophthalmology Seoul National University Bundang Hospital College of Medicine Seoul National University Sungnam, Korea

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The authors have no proprietary interest in any of the materials used in this study This study was supported by a grant from the Seoul National University Hospital Research Fund. undertaken. Visual acuity of the amblyopic and fellow eyes was measured by means of linear optotypes of Snellen type by a masked tester. The compliance was recorded at each clinic visit. For statistical analysis, visual acuities have been converted to logMAR equivalents. Pre- and postocclusion visual acuities have been compared using paired *t*-tests. Analysis was also performed using the nonparametric method of Wilcoxon's signed rank test.

A total of 16 subjects consisting of eight boys and eight girls were enrolled in this study. All of them were 9 years or older (range, 9.0–14.5 years; mean, 10.5 years) at the commencement of amblyopia therapy for anisometropic (nos. 1-11; mean, 10.4 years; range, 9-14.5 years), strabismic (nos. 12, 13; mean, 11.5 years; range, 10.5–12.3 years), or anisometropic and strabismic (nos. 14-16; mean, 10.3 years; range, 9.2-11.2 years) amblyopia (Table 1). Anisometropic amblyopes were defined as amblyopia with spherical difference between the two eyes of +1.00 diopters (D) or more of hypermetropia or -2.00 D or more of myopia, and with no heterotropia in the alternate cover test. Strabismic amblyopes had exotropia of at least 15 prism diopter (PD) in the alternate cover test and a spherical difference between the two eyes of less than 0.75 D. Anisometropic and strabismic amblyopes had both exotropia of at least 10 PD in the alternate cover test and a spherical difference between the two eyes of -2.00 D or more of myopia. The average age at the commencement of amblyopia therapy for the 16 patients was 10.5 years (range, 9.0-14.5 years).

The initial therapy consisted of full optical spectacle correction for the amblyopic eye for 4 weeks and then

followed by a full-time occlusion with adhesive patch in 14 patients and a part-time occlusion in two patients who showed a difference in visual acuity of over two lines between the eyes. A full spectacle correction was prescribed in all cases on the basis of a cycloplegic refraction with cyclopentolate hydrochloride 1% for 20-40 min. Following the initial period with occlusion along with spectacles, the changes in visual acuity were compared at 3-, 6-, 12-, and 24-month intervals. In patients with a low compliance, both the patient and their parents were asked to visit the clinic once a month and were given a great deal of encouragement and education as to the significance of amblyopia therapy. The patients and parents were asked to keep a diary of the occlusion time of each day. The parents were asked to bring a letter to school so that the patients could occlude their eyes even there. This therapy was continued until there was no further improvement in the visual acuity of the amblyopic eye for at least 3 months. In the 14 patients with full-time occlusion, 11 patients had part-time occlusion with a patch and three patients were given occlusive Min's glasses<sup>®</sup> (P & M Inc., Daejon, Korea)<sup>13</sup> so as to maintain the visual acuity after they were considered to have reached a plateau. Final visual acuity was defined as the visual acuity at the end of the study. After the maintenance therapy was discontinued, the visual acuity was retested 6-12 months later. If there was any decrease in visual acuity, the maintenance therapy was reinstituted.

The compliance was categorized as good if occlusion was performed >70% of the time, fair if occlusion was performed for 30–70% of the time, and poor if occlusion was performed <30% of the time.

Table 1 Clinical characteristics and outcome of amblyopia therapy in 16 patients

Patient/sex	Initial age (years)	Refraction		Anisome-	Strabismus	Final age	Occlusion	Initial VA		Final VA		Compliance
		Amblyopic eye	Fellow eye	tropia (D)		(years)	full/part-time (months)	Snellen	logmar	Snellen	logMAR	(%)
1/F	10.3	-4.75	-0.25	-4.50	Ortho	11.1	1/4	20/60	0.5	20/20	0	
2/F	9.0	+3.75	+0.75	+3.00	Ortho	11.8	0/9	20/40	0.3	20/20	0	>70
3/F	9.6	-3.25	-0.50	-2.75	Ortho	11.0	8/0	20/40	0.3	20/25	0.1	>70
4/F	9.0	+5.75	-1.75	+7.50	Ortho	12.5	1/17	20/40	0.3	20/20	0	>70
5/M	9.0	-4.75	-1.75	-3.00	Ortho	10.2	$1/12^{a}$	20/100	0.7	20/30	0.2	>70
6/M	11.8	+1.25	-1.00	+2.25	Ortho	13.6	1/7	20/40	0.3	20/20	0	>70
7/M	14.5	+3.25	+0.25	+3.00	Ortho	15.5	0/10	20/60	0.5	20/50	0.4	30-70
8/M	9.0	+2.75	+1.75	+1.00	Ortho	11.8	$1/12^{a}$	20/50	0.4	20/20	0	
9/F	9.0	+4.50	+0.50	+4.00	Ortho	11.3	1/26	20/100	0.7	20/20	0	>70
10/M	9.2	-3.75	-1.75	-2.00	Ortho	10.2	5/2	20/40	0.3	20/20	0	>70
11/M	13.9	+4.50	+3.00	+1.50	Ortho	14.9	1/4	20/100	0.7	20/40	0.3	
12/M	10.5	-1.25	-0.75	-0.50	40XT	11.5	2/2	20/40	0.3	20/25	0.1	>70
13/F	12.3	-3.00	-2.25	-0.75	16X(T)	10.9	3/12	20/30	0.2	20/20	0	>70
14/M	9.2	-6.50	-2.50	-3.50	35XT	11.4	3/0	20/40	0.3	20/20	0	>70
15/F	11.2	-4.00	-2.00	-2.00	10X(T)	12.4	1/13	20/30	0.2	20/20	0	>70
16/F	10.5	-9.75	-1.00	-8.75	55XT	15.0	3/10a	20/40	0.3	20/25	0.1	>70

-: compliance unrecorded; "Part-time occlusion with occlusive Min's glasses.13

Anisometropia was present in 11 patients (nos. 1–11), strabismus in two patients (nos. 12 and 13), and both anisometropia  $\geq -2.00 \text{ D}$  and strabismus in three patients (nos. 14–16) (Table 1).

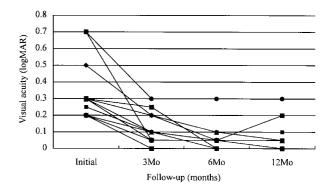
The initial worst visual acuity after 4 weeks of full-time spectacle wear was 20/100 in three patients, ranged from 20/80 to 20/40 in 11 patients, and 20/30 in two patients. The refractive errors of the amblyopic eye were myopic in seven patients, hyperopic in seven patients, and astigmatism in two patients. The average amount of anisometropia between the amblyopic eye and the fellow eye was 2.56 D. The type of strabismus was a constant exotropia of 35, 40, and 55 PD in three patients, and an intermittent exotropia of 10 and 16 PD in two patients (Table 1). Patient no. 16 showed a visual acuity of 20/401 month after he first came. The visual acuities of fellow eyes were 20/20 in 14 patients and 20/15 in the two remaining patients before patching, which did not show any significant improvement during the follow-up period.

The final visual acuity of 15 out of 16 patients (94%) at the end of the study improved by at least two lines. The final visual acuities ranged from 20/20 to 20/30 for 14 patients, 20/40 in one patient (no. 11), and 20/50 in the remaining patient (no. 7). Patient no. 11 underwent occlusion therapy from age 13.9 to 14.9 years and the visual acuity had improved from 20/100 to 20/40. Patient no. 7 underwent occlusion therapy from age 14.5 to 15.5 years and showed just a one-line improvement, with the poorest compliance among the patients (Table 1).

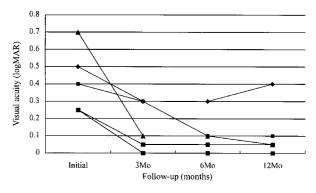
Paired *t*-tests and Wilcoxon's signed rank test showed a statistically significant difference between mean preand postocclusion visual acuities converted to logMAR equivalents (0.39375 *vs* 0.075, P < 0.001).

The duration of therapy averaged 10 months (range, 3–27 months). The compliance was >70% in 12 out of the 16 patients (75%). Improvement in the visual acuity occurred mostly in the initial 3–6 months (Figures 1 and 2). Occlusion-induced amblyopia did not occur in any patient. The average follow-up period was 24 months (range, 12–63 months).

There was no patient who developed diplopia after patching. Regarding the binocular state of strabismus before and after patching, two patients (nos. 13 and 15) with intermittent exotropia showed less than 6 PD of exophoria after improvement of visual acuity with patching. Among three patients with constant exotropia, two showed less than 10 PD of exophoria or intermittent exotropia and one remaining patient (no. 14) underwent bilateral rectus recession after improvement of visual acuity with patching. Titmus stereo test in patients 12–16 showed 200–60 s of arc of stereoacuities, which did not



**Figure 1** logMAR visual acuity in anisometropic amblyopia. Improvement in the visual acuity occurred mostly in the initial 3–6 months.



**Figure 2** logMAR visual acuity in strabismic amblyopia, and in anisometropic and strabismic amblyopia. Improvement in the visual acuity occurred mostly in the initial 3–6 months.

show any significant improvement over the two-level difference before and alter improvement of visual acuity.

#### Discussion

There is general agreement that amblyopia must be treated at an early age and that treatment is more prolonged and less beneficial in older children.<sup>3-9</sup> However, the age beyond which treatment is ineffective is still controversial. Oliver *et al*<sup>5</sup> also found that children older than 8 years who complied with the treatment showed a marked improvement in their visual acuity, almost as good as that in younger children. Mintz-Hittner and Fernandez<sup>6</sup> reported significant improvements in the visual acuity of 36 compliant children between the ages of 7.0 and 10.3 years undergoing an occlusion or penalization therapy.<sup>6</sup> Hedgpeth and Sullivan<sup>7</sup> also reported that an anisometropic amblyopia could be treated successfully at least until the age of 12 years. In contrast, Rutstein and Fuhr<sup>8</sup> reported that a visual acuity of 20/40 or better could be obtained in only 27% of patients older than 8 years. Epelbaum  $et al^9$  also reported that the efficiency of strabismic amblyopia treatment decreased as a function

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of age and was nil by the time the patient had reached 12 years of age, and only children with an anisometropic amblyopia responded to therapy at the later ages. It was reported that the greater the compliance, the better the result.<sup>5,6,12</sup> The reported high failure rates in older children might be attributed to the lack of compliance rather than to age-related factors. Our study strongly reaffirms that given compliance, occlusion therapy for either anisometropic or strabismic amblyopia can be successful even if initiated after the age of 9 years.

In our amblyopes of 9.0–14.5 years of age at commencement of the occlusion, 14 out of 16 patients (88%) had improved their visual acuity by up to 20/30. If the one patient of 14.5 years of age at initiation of occlusion is excluded, all of the patients 9.0–12.3 years of age at the initiation of occlusion improved their visual acuity better than 20/40. The recorded compliance was good, with more than 70% of the occlusion in every recorded patient except the 14.5-year-old patient. This result means that amblyopia therapy for anisometropic and strabismic amblyopia can be successful even if initiated after the age of 9 years as long as there is good compliance.

There has been some debate as to how long the critical period of visual development actually lasts. The critical period for development does not necessarily follow the same time course as the plastic period during which the visual system is still amenable to successful treatment. In humans, even adults with amblyopia demonstrated residual plasticity in the visual system.<sup>4,14–18</sup> Our results also suggest that amblyopia can be treated successfully beyond the age that is usually considered to be the critical period for the development of amblyopia. Therefore, the plastic period for treating amblyopia may extend beyond the critical development period substantially, and the age of the patient should not be a limiting factor for determining whether or not to institute therapy.

In this study, most of the improvement occurred during the first 3 months of treatment regardless of the type of amblyopia (Figures 1 and 2). The same result could be found in other studies for amblyopes in other age groups.<sup>5,19,20</sup> Thus given compliance, the response of occlusion therapy for the initial 3 months could be invaluable for predicting the prognosis of the amblyopia.

In conclusion, a therapeutic trial is advisable even in older children, provided the patient is compliant. The period of visual acuity recovery for amblyopia should be extended beyond the conventional view of 7 years of age as the upper limit for treatment.

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