PERSPECTIVE

Classification and diversity of amblyopia

DAPHNE MAURER[1] AND SUZANNE P. McKEE[2]

[1]Department of Psychology, Neuroscience and Behavior, McMaster University, Hamilton, Ontario, Canada
[2]Smith-Kettlewell Eye Research Institute, San Francisco, California

Abstract

Amblyopia is a developmental disorder that affects the spatial vision of one or both eyes in the absence of an obvious organic cause; it is associated with a history of abnormal visual experience during childhood. Subtypes have been defined based on the purported etiology, namely, strabismus (misaligned eyes) and/or anisometropia (unequal refractive error). Here we consider the usefulness of these subclassifications.

Keywords: Strabismus, Anisometropia, Acuity, Contrast sensitivity, Binocularity

The generally accepted definition of amblyopia is reduced visual acuity, despite best optical correction, when measured with an optotype chart, such as the LogMAR chart. Typically, a two-line difference between the eyes is taken as evidence of unilateral amblyopia. Special pediatric charts with single letters or pictures are used to test preschool children. Estimates of the prevalence of unilateral amblyopia in preschool children range between one and 3%, with the proportion depending somewhat on the ethnic composition of the study sample (MEPEDS, 2008; Friedman et al., 2009; Ying et al., 2014). It is the most common cause of monocular visual impairment in children (Webber & Wood, 2005; Gunton, 2013). A similar proportion of adults suffer from unilateral amblyopia (Attebo et al., 1998). Bilateral amblyopia is much less common (0.1-0.45%; Roboaei et al., 2005; MEPEDS, 2008) and is generally associated with a history visual deprivation produced by cataracts, ptosis, or high refractive errors in both eyes.

Amblyopia has been subdivided traditionally based on the condition thought to be its cause, primarily strabismus (misaligned eyes) or anisometropia (difference in refractive error between the two eyes) or some combination of both, yielding the common labels of strabismic amblyopia, anisometropic amblyopia, and strabismic-anisometropia or mixed amblyopia. Amblyopia in patients with the rare history of deprivation is labeled deprivation amblyopia.

Does this classification, based on purported etiology, produce different functional types of amblyopia, or are there different functional types that would arise from some other classification scheme, or is amblyopia, however caused, essentially the same abnormality? This answer to these questions is especially important in determining the treatment because the etiology may not be fully apparent at the time that amblyopia is diagnosed, and different forms of amblyopia may require different management.

Classification based on visual function

Numerous behavioral studies, made almost exclusively on adults with amblyopia, have suggested strongly that strabismus and anisometropia do indeed lead to different patterns of visual loss (Levi & Klein, 1982a,b, 1985; Hess & Pointer, 1985). In the largest study to date, McKee et al. (2003) measured the acuity and contrast sensitivity of 427 individuals either with amblyopia or with risk factors for amblyopia, e.g., strabismus or anisometropia, plus 68 normal control observers. Participants were assigned to one of eleven groups, based on their history and a detailed clinical examination of their oculomotor and refractive characteristics. Participants were not classified on the basis of their optotype (LogMAR) visual acuity; instead the study explored the relationships between optotype acuity and four other measures (grating acuity, vernier acuity, edge contrast sensitivity, and contrast sensitivity measured with Pelli-Robson chart). Statistical analysis of this large data set revealed that two factors accounted for about 80% of the variance separating the 11 clinical groups: an acuity factor and a sensitivity factor. In Fig. 1, the average sensitivity factor of the nonpreferred eyes (the eyes with the poorer acuity) is plotted *versus* the average acuity factor of the nonpreferred eyes for each of these clinically defined groups. Note that these groups include both amblyopic and nonamblyopic observers, e.g., all patients with pure strabismus whether amblyopic or not.

Figure 1 represents a kind of 'map' of amblyogenic conditions with the normal control group (black symbol) on the right showing the best acuity and the strabismic-anisometropic group (blue symbol) on the left showing the poorest acuity. Clearly, the strabismic groups (red symbols) and the anisometropic group (green symbol)

Address correspondence to: Dr. Daphne Maurer, Psychology, Neuroscience & Behaviour, McMaster University, Hamilton, ON L8S 4K1 Canada. E-mail: mcmaster@maurer.ca
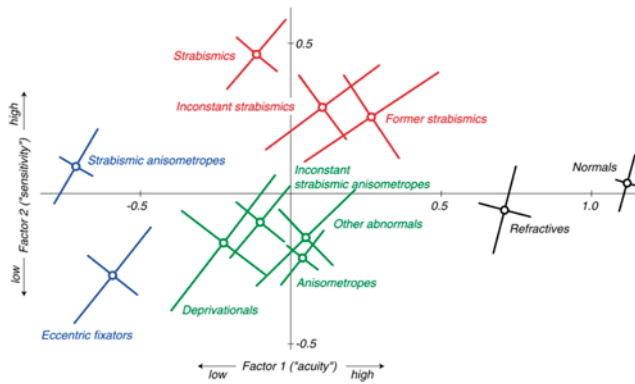
Fig. 1. McKee et al. (2003). The pattern of visual deficits in amblyopia. *Journal of Vision* 3, 380–405 (used with permission).

fall in different sectors of this map, suggesting that different patterns of visual loss are associated with these different conditions. The difference between the red and green groups is largely dependent on a significant difference in contrast sensitivity. The strabismic groups are more sensitive than the anisometropes; indeed, they are significantly more sensitive than the normal controls, a curious hypersensitivity that may arise from the loss of binocularity. McKee et al. (2003) speculated that the reorganization produced by the loss of binocularly-driven neurons would lead to a greater visual input to monocularly driven neurons, which would thus enhance their thresholds at lower contrast. Thus, they predicted that monocular contrast sensitivity in both eyes of the strabismic groups would be slightly better than normal, which is indeed what their results showed. Another item of interest in this map is that the deprivation group falls near the anisometropic groups, which may arise from the degraded retinal image produced by both anisometropia and deprivation.

What is missing from this map is an important functional difference between the strabismic groups and the anisometropic group—namely the quality of their binocular performance. In addition to their measures of contrast and acuity, McKee et al. (2003) made two measurements of the binocular function. The first was a widely used clinical test of stereopsis—the Randot "Circles" test. Observers viewed the test circles with the best optical correction, but without prisms. The second test, binocular motion integration (BMI), was an experimental measure of the cortical capacity to integrate or fuse diverse images into a coherent motion. This test used the dichoptic quadrature motion stimulus devised by Shadlen and Carney (l986). Briefly, each eye viewed a horizontal sinusoidal grating whose contrast was modulated sinusoidally at 2 Hz. The stimuli presented to the two eyes were spatially and temporally 90 deg out of phase; the direction of the phase shift determined whether the gratings appeared to move up or down. Observers were asked to judge the direction of motion for 30 trials. Performance on each of these two tests was graded as 'pass–fail'. If an observer had any measureable stereopsis based on the Circles test, he or she was given a pass. In the BMI test, an observer was given a pass if he or she correctly judged the direction of motion on more than 70% of trials. About 80% of participants either passed both tests or failed both; all normal observers passed both. Most members of the strabismic groups (~90%; red and blue symbols) failed both measures, but a significant proportion of anisometropes (~64%) passed both. In short, based on these tests, most of the strabismic participants had difficulty combining the information binocularly, while the majority of anisometropic participants retained this capability.

Although the classification scheme shown by the 'map' is based on rigorous psychophysical measurements made on a large sample of the participants, it can be criticized on many grounds. First, the psychophysical measurements tested only acuity, contrast sensitivity, and binocular functioning. A different configuration might emerge if more functions had been tested. Some types of motion judgments (Simmers et al., 2003; Ho & Giaschi, 2006; Thompson et al., 2011), subitizing (counting number of items seen during a brief presentation; Sharma et al., 2000), and attentional control (Popple & Levi, 2008; Farzin & Norcia, 2011) have been shown to be abnormal in amblyopia. The inclusion of oculomotor measurements might also alter the classification scheme since they appear to vary with amblyogenic conditions. Many studies have found that strabismic patients with amblyopia show poorer oculomotor control than anisometropic patients with amblyopia; their fixation is much more unsteady, and the latency of their saccades is also longer (Schor & Hallmark, 1978; Cuiffreda et al., 1979; Zhang et al., 2008; Niechwiej-Szwedo et al., 2010; Gonzalez et al., 2012; Niechwiej-Szwedo et al., 2012; Chung et al., 2015; McKee et al., 2016). It is also possible to break down the clinical groups shown in the map into further subgroups. For example, there is considerable evidence that the early onset (infantile) strabismus produces different functional deficits than those of the late onset (refractive) strabismus (Schor et al., 1997; Brosnahan et al., 1998; Sloper, 2016).

Another problem with the 'map' is the categorical designation of the clinical groups. These designations were based on measurements made by ophthalmologists and optometrists who had been trained on an extensive clinical protocol. Strabismus and anisometropia were defined by the conventional criteria at the time of the study: a difference between the eyes of 1 diopter in refractive error at the maximum anisometropic meridian for anisometropia; eye movement seen under unilateral and alternating cover tests at near (0.3 m) and distance (6 m) for constant strabismus. The dichotomy between these two conditions might have been less obvious if more precise measurements of ocular alignment had been used (Hunter et al., 1999; Gramatikov et al., 2007). Were the two eyes of the amblyopic members of the anisometropic group really aligned within the normal range? About half of the patients with anisometropic amblyopia passed both of our binocular tests, so we do know that their binocular functioning was better than that of the amblyopic members of the designated strabismic groups. However, it is intriguing that other functional characteristics (vernier acuity, saccadic latency) of the patients with anisometropia who failed both of our binocular tests were somewhat similar to strabismic patients (McKee, 1998; McKee et al., 2016). Generally, the nonbinocular anisometropic patients were those with the poorest acuity in their nonpreferred eye—individuals suffering from severe unilateral amblyopia. If they had no capacity for binocular integration, what mechanism was keeping their eyes aligned? A more sensitive measure of alignment might have revealed that these nonbinocular anisometropic patients were microstrabismic.

The fluidity of these categories probably reflects the complex interaction between refractive error (spherical, astigmatic and anisometropic), spatial visual performance, and eye alignment during development. Apparently similar infants can develop along quite different paths (Babinsky & Candy, 2013; Barrett et al., 2013). Thus a patient with strabismic amblyopia may present with quite different forms (ranging from intermittent exotropia, or microtropia to constant alternating or unilateral esotropia) and different forms of neural adaptation or consequence (from suppression to diplopia). We are yet to fully understand the factors that predict the developmental path different infants will take and that define the range of untreated visual function that presents for an examination in a clinic.

In conclusion, it may be more useful to describe prototypes that differ but have unclear boundaries between them, rather than to assign amblyopic individuals or individuals at risk for amblyopia to distinct clinical categories.

Does classification matter?

Despite the vast amount of basic research demonstrating that different patterns of functional loss are associated with different amblyogenic conditions, the presenting condition does not predict treatment outcome. Following patching or penalization, which force use of the amblyopic eye by depriving the fellow eye of vision, patients classified as suffering from strabismic and anisometropic amblyopia, treated before age 7, both show the same amount of improvement in acuity (PEDIG, 2003*a,b,c,d*). One could argue either that the patients were misclassified due to insensitive measures of misalignment or that this analysis was too narrowly focused on acuity and that more extensive testing including measures of contrast sensitivity and binocularity (Bosworth & Birch, 2003; Birch, 2013) would show that the associated condition does matter to the outcome. Using a battery of tests to classify the patients might lead to a better prediction of the outcome, although obtaining reliable behavioral measures in preschoolers, the age at which amblyopia or an amblyogenic condition is diagnosed, is challenging. However, a test battery might be prognostic in older patients with amblyopia in whom a later intervention is being contemplated and provide insight on treatment of young patients with amblyopia if more reliable behavioral measures in preschoolers become available.

Future directions

- It would be useful to agree on a common set of sensitive tests that can be used clinically and in research to classify patients with amblyopia. That list should include measures of crowded acuity (acuity measured with symbols spaced as in words), contrast sensitivity, and binocular functions (stereopsis and binocular interactions). Models of binocular interactions are necessary to understand patterns of empirically observed binocular suppression in different behavioral paradigms (Huang et al., 2011). Normative data for agreed-upon measures are needed at different ages. Although some of the measures may not be possible with very young children, the patterns found in older children can inform the understanding of etiology and possibly effective treatments. There is an urgent need for both more efficient and precise behavioral measurements of the functional vision (e.g., Lesmes et al., 2010) as well as theoretical models that link core visual deficits in amblyopia to the observed poor visual performance of patients with amblyopia.
- One of the major problems in amblyopia is regression—a return to amblyopic acuity levels following successful treatment. Clinical classification, based on amblyogenic condition, may be useful in determining who is likely to regress, and thus, who may benefit from additional treatment or alternative treatments during elementary school years.
- Currently, there are a variety of new experimental treatments, involving perceptual learning, video game play, and dichoptic presentation (Zhou et al., 2006; Li et al., 2011; Hess et al., 2012). These new treatments might be especially useful for improving acuity in older children and young adults who have residual amblyopia following treatment as children. Here again, a new classification system may predict who will benefit from these new treatments.
- As part of a new classification system, it would be worthwhile to pursue research on genetic markers and epigenetic factors, which may predict which patients with amblyopia are likely to resist treatment or to regress to poor acuity following treatment. Special treatment protocols could be developed for those individuals whose risk profiles, based on a new classification, will make them more likely to suffer from persistent amblyopia.

References

ATTEBO, K., MITCHELL, P., CUMMING, R., SMITH, W., JOLLY, N. & SPARKES, R. (1998). Prevalence and causes of amblyopia in an adult population. *Ophthalmology* **105**, 154–159.

BABINSKY, E. & CANDY, T.R. (2013). Why do only some hyperopes become strabismic? *Investigative Ophthalmology & Visual Science* **54**, 4941–4955.

BARRETT, B.T., BRADLEY, A. & CANDY, T.R. (2013). The relationship between anisometropia and amblyopia. *Progress in Retinal and Eye Research* **36**, 120–158.

BIRCH, E.E. (2013). Amblyopia and binocular vision. *Progress in Retinal and Eye Research* **54**, 1998–2003.

BOSWORTH, R.G. & BIRCH, E.E. (2003). Binocular function and optotype-grating acuity discrepancies in amblyopic children. *Investigative Ophthalmology & Visual Science* **44**, e3183.

BROSNAHAN, D., NORCIA, A.M., SCHOR, C.M. & TAYLOR, D.G. (1998). OKN, perceptual and VEP direction biases in strabismus. *Vision Research* **38**, 2833–2840.

CHUNG, S.T.L., KUMAR, G., LI, R.W. & LEVI, D.M. (2015). Characteristics of fixational eye movements in amblyopia: Limitations on fixation stability and acuity? *Vision Research* **114**, 87–99.

CUIFFREDA, K.J., KENYON, R.V. & STARK, L. (1979). Fixational eye movements in amblyopia and strabismus. *Journal of the American Optometric Association* **50**, 1251–1258.

FARZIN, F. & NORCIA, A.M. (2011). Impaired visual decision-making in individuals with amblyopia. *Journal of Vision* **14**, 1–10.

FRIEDMAN, D.S., REPKA, M.X., KATZ, J., GIORDANO, L., IBIRONKE, J., HAWSE, P. & TIELSCH, J.M. (2009). Prevalence of amblyopia and strabismus in white and African American children aged 6 through 71 months the baltimore pediatric eye disease study. *Ophthalmology* **116**, 2128–2134.

GONZALEZ, E.G., WONG, A.M.F., NIECHWIEJ-SZWEDO, E., TARITA-NISTOR, L. & STEINBACH, M.J. (2012). Eye position stability in amblyopia and in normal binocular vision. *Investigative Ophthalmology & Visual Science* **53**, 5386–5394.

GRAMATIKOV, B.I., ZALLOUM, O.H., WU, Y.K., HUNTER, D.G. & GUYTON, D.L. (2007). Directional eye fixation sensor using birefringence-based foveal detection. *Applied Optics* **46**, 1809–1818.

GUNTON, K.B. (2013). Advances in amblyopia: What have we learned from PEDIG trials? *Pediatrics* **131**, 540–547.

HESS, R.F. & POINTER, J.S. (1985). Differences in the neural basis of human amblyopia: The distribution of the anomaly across the visual field. *Vision Research* **25**, 1577–1594.

HESS, R.F., THOMPSON, B., BLACK, J.M., MACHARA, G., ZHANG, P., BOBIER, W.R. & COOPERSTOCK, J. (2012). An ipod treatment of amblyopia: An updated binocular approach. *Journal of Optometry* **83**, 87–94.

HO, C.S. & GIASCHI, D.E. (2006). Deficient maximum motion displacement in amblyopia. *Vision Research* **46**, 4595–4603.

HUANG, C.B., ZHOU, J., LU, Z.L. & ZHOU, Y. (2011). Deficient binocular combination reveals mechanisms of anisometropic amblyopia: signal attenuation and interocular inhibition. *Journal of Vision*. 4. doi:10.1167/11.6.4

HUNTER, D.G., PATEL, S.N. & GUYTON, D.L. (1999). Automated detection of foveal fixation by use of retinal birefringence scanning. *Applied Optics* **38**, 1273–1279.

LESMES, L.A., LU, Z.L., BAEK, J. & ALBRIGHT, T. (2010). Bayesian adaptive estimation of the contrast sensitivity function: The quick CSF method. *Journal of Vision*, 17. doi:10.1167/10.3.17

LEVI, D.M. & KLEIN, S. (1982a). Hyperacuity and amblyopia. *Nature* **298**, 268–270.

LEVI, D.M. & KLEIN, S. (1982b). Differences in vernier discrimination for grating between strabismic and anisometropic amblyopes. *Investigative Ophthalmology & Visual Science* **23**, 398–407.

LEVI, D.M. & KLEIN, S.A. (1985). Vernier acuity, crowding and amblyopia. *Vision Research* **25**, 979–991.

LI, R.W., NGO, C., NGUYEN, J. & LEVI, D.M. (2011). Video-game play induces plasticity in the visual system of adults with amblyopia. *PLoS Biology* **9**, e1001135.

MCKEE, S.P. (1998). Binocular functioning and visual acuity in amblyopia. In *Vision Screening in the Preschool Child*, pp. 206–228. Bethesda, MD: Health Resources and Services Administration, HHS.

MCKEE, S.P., LEVI, D.M. & MOVSHON, J.A. (2003). The pattern of visual deficits in amblyopia. *Journal of Vision* **3**, 380–405.

MCKEE, S.P., LEVI, D.M., SCHOR, C.M. & MOVSHON, J.A. (2016). Saccadic latency in amblyopia. *Journal of Vision* **16**, 3.

Multi-Ethnic Pediatric Eye Disease Study Group (2008). Prevalence of amblyopia and strabismus in African American and Hispanic children ages 6 to 72 months. *Ophthalmology* **115**, 1229–1236.

NIECHWIEJ-SZWEDO, D., GOLTZ, H.C., CHANDRAKUMAR, M., HIRJI, Z.A. & WONG, A.M.F. (2010). Effects of anisometropic amblyopia on visuomotor behavior, I: Saccadic eye movements. *Investigative Ophthalmology & Visual Science* **51**, 6348–6354.

NIECHWIEJ-SZWEDO, D., CHANDRAKUMAR, M., GOLTZ, H.C. & WONG, A.M.F. (2012). Effects of strabismic amblyopia and strabismus without amblyopia on visuomotor behavior I: Saccadic eye movements. *Investigative Ophthalmology & Visual Science* **53**, 7458–7468.

POPPLE, A.V. & LEVI, D.M. (2008). The attentional blink in amblyopia. *Journal of Vision* **8**, 1–9.

REPKA, M.X., BECK, R.W., HOLMES, J.M., BIRCH, E.E., CHANDLER, D.L., COTTER, S.A. et al., PEDIATRIC EYE DISEASE INVESTIGATOR GROUP (2003). A randomized trial of patching regimens for treatment of moderate amblyopia in children. *Archives of Ophthalmology* **121**, 603–611.

ROBAEI, D., ROSE, K., OJAIMI, E., KIFLEY, A., HUYNH, S. & MITCHELL, P. (2005). Visual acuity and the causes of visual loss in a population-based sample of 6-year-old Australian children. *Ophthalmology*, **112**, 1275–1282.

SCHOR, C.M., FUSARO, R.E., WILSON, N. & MCKEE, S.P. (1997). Prediction of early-onset esotropia from components of the infantile squint syndrome. *Investigative Ophthalmology & Visual Science* **38**, 719–740.

SCHOR, C.M. & HALLMARK, W. (1978). Slow control of eye position in strabismic amblyopia. *Investigative Ophthalmology & Visual Science* **17**, 577–581.

SHADLEN, M. & CARNEY, T. (1986). Mechanisms of human motion perception revealed by a new cyclopean illusion. *Science* **232**, 95–97.

SHARMA, V., LEVI, D.M. & KLEIN, S.A. (2000). Undercounting features and missing features: Evidence for a high level deficit in strabismic amblyopia. *Nature Neuroscience* **3**, 496–501.

SIMMERS, A.J., LEDGEWAY, T., HESS, R.F. & MCGRAW, P.V. (2003). Deficits to global motion processing in human amblyopia. *Vision Research* **43**, 729–738.

SLOPER, J. (2016). Knapp lecture: The other side of amblyopia. *Journal of AAPOS* **20**, 1–13.

THOMPSON, B., RICHARD, A., CHURAN, J., HESS, R.F., AAEN-STOCKDALE, C. & PACK, C.C. (2011). Impaired spatial and binocular summation for motion direction discrimination in strabismic amblyopia. *Vision Research* **51**, 577–584.

THE PEDIATRIC EYE DISEASE INVESTIGATOR GROUP (2003a). The course of moderate amblyopia treated with patching in children: Experience of the amblyopia treatment study. *American Journal of Ophthalmology* **136**, 620–629.

THE PEDIATRIC EYE DISEASE INVESTIGATOR GROUP (2003b). The course of moderate amblyopia treated with atropine in children: Experience of the amblyopia treatment study. *American Journal of Ophthalmology* **136**, 630–639.

THE PEDIATRIC EYE DISEASE INVESTIGATOR GROUP (2003c). A randomized trial of prescribed patching regimens for treatment of severe amblyopia in children. *Ophthalmology* **110**, 2075–2087.

THE PEDIATRIC EYE DISEASE INVESTIGATOR GROUP (2003d). A randomized trial of patching regimens for treatment of moderate amblyopia in children. *Archives of Ophthalmology* **121**, 603–611.

WEBBER, A.L. & WOOD, J. (2005). amblyopia: Prevalence, natural history, functional effects and treatment. *Clinical and Experimental Optometry* **88**, 365–375.

YING, G.S., MAGUIRE, M.G., CYERT, L.A., CINER, E., QUINN, G.E., KULP, M.T., OREL-BIXLER, D. & MOORE, B. (2014). Prevalence of vision disorders by racial and ethnic group among children participating in head start. *Ophthalmology* **121**, 630–636.

ZHANG, B., STEVENSON, S.S., CHENG, H., LARON, M., KUMAR, G., TONG, J. & CHINO, Y.M. (2008). Effects of fixation instability on multifocal VEP (mfVEP) responses in amblyopes. *Journal of Vision* **8**, 16.

ZHOU, Y., HUANG, C., XU, P., TAO, L., QIU, Z. & LI, X., LU, Z.L. (2006). Perceptual learning improves contrast sensitivity and visual acuity in adults with anisometropic amblyopia. *Vision Research* **46**, 739–750.