AH Alsuhaibani¹, TM Bosley¹, RA Goldberg² and

Entropion in children with isolated peripheral facial nerve paresis

Abstract

Purpose Adults with facial nerve paresis (FNP) generally develop ectropion, but a recent report of children with syndromatic FNPs implies that entropion may be more common in this setting than ectropion. This study evaluates eyelid position and other periorbital changes in children with isolated, non-syndromatic FNP.

Methods Charts were reviewed of 10 sequential children who presented to a major national eye referral centre with isolated FNP of variable aetiology. Severity of FNP was assessed according to the House-Brackmann scale.

Results All 10 patients (4 males and 6 females; mean age at presentation, 4 years) had unilateral, isolated FNP. Mild lowereyelid entropion was present in four patients, and severe lower-eyelid entropion required surgical correction in three patients. All patients had lower eyelid retraction (mean 2.3 mm) and lagophthalmos (mean 2.9 mm). None had enophthalmos, lower eyelid ectropion, or brow ptosis. Conclusion Unlike adults, children with isolated FNP seem prone to develop entropion rather than ectropion. Entropion reported previously in five syndromic children with FNP seems more likely related to patients' age than to their congenital syndromes.

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Keywords: entropion; ectropion; children; facial paresis; facial palsy

Introduction

Ectropion is common in adults with facial nerve paresis (FNP),^{1,2} but a recent report described entropions in a small group of

children with syndromic FNP.³ Eyelid position has not yet been carefully evaluated in children with isolated FNP,^{2,4–10} so this study reports eyelid and periorbital changes in 10 consecutive children presenting with isolated FNP.

Methods

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Medical records were reviewed of 10 consecutive children who were presented to King Abdulaziz University Hospital, Riyadh, Saudi Arabia, with isolated lower motor neuron FNP between January 2007 and January 2011. Patients who had undergone previous ophthalmic surgery and patients with obvious craniofacial disorders were excluded. The information that was recorded included the cause of FNP, the complete ophthalmologic clinical examination, and ophthalmologic surgical procedures needed to manage the consequences of FNP.

Results

Table 1 summarizes the aetiology of FNP, age at onset and presentation, lower eyelid observations, and surgeries performed. All 10 patients (4 males and 6 females; mean age 4 years at presentation) had unilateral FNP with House-Brackmann grade III or IV at presentation.¹¹ Severe lower eyelid entropion with trichiasis was present in three patients (Figures 1a and b), while mild lower eyelid entropion without trichiasis was seen in four patients when initially evaluated (Figures 2a and b). None of the patients had enophthalmos, lower eyelid ectropion, or a significant abnormality of the eyelid skin, tarsus, or conjunctiva. All patients had lower eyelid retraction (mean 2.3 mm; range 1–3 mm) and lagophthalmos (mean 2.9 mm; range 1-4 mm) (Figure 2c),

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Case	Gender	Age (years)		HBG	Aetiology	Lower lid entropion	Lower lid retraction (mm)	Lagophthalmos (mm)	Surgical procedure
		At onset	At presentation				, , , , , , , , , , , , , , , , ,		1
1	Female	Congenital	3	III	Congenital	Mild entropion	2	3	None
2	Female	4	5	III	MEI	Severe entropion	1	1	Entropion repair
3	Female	Congenital	3	IV	Congenital	Mild entropion	2	2	None
4	Male	Congenital	5	III	Congenital	No entropion	3	4	None
5	Female	2	3	IV	Traumatic	Mild entropion	2	3	None
6	Male	Congenital	4	IV	Congenital	No entropion	3	3	None
7	Female	2	3	IV	MEI	Severe entropion	2	2	Entropion repair
8	Male	3	8	III	Traumatic	Mild entropion	3	4	None
9	Female	4	4	IV	Traumatic	No entropion	3	4	None
10	Male	Congenital	2	III	Congenital	Severe entropion	2	2	Entropion repair

Table 1	Patient demogra	phics, lid	position,	and	surgical	procedures

Abbreviations: HBG, House-Brackmann grade; MEI, middle ear infection

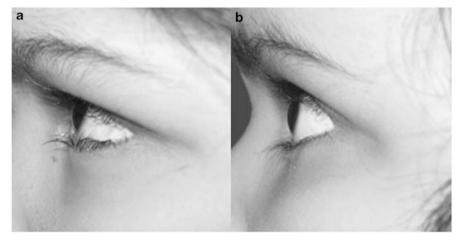


Figure 1 (a) A 4-year-old girl with left-sided facial paralysis secondary to middle ear infection presented with severe lower eyelid entropion causing marked ocular irritation due to trichiasis. (b) The same patient 6 months after entropion repair by reinsertion of the lower eyelid retractors into the lower anterior part of the tarsus. Entropion was improved, but she continued to have the same degree of lower eyelid retraction postoperatively.

and three patients had 1 mm of upper eyelid retraction on the affected side.

All patients had moderate to severe decrease in frontalis function on the affected side, but none had eyebrow ptosis. Also, none of the patients had clinically evident aberrant regeneration of the facial nerve, but five patients had mild atrophy of the mid-face on the affected side. All patients had normal ocular motility and otherwise grossly normal neurologic and general medical examinations; none had major general medical problems. The five patients with congenital onset did not have a family history of FNP.

In three patients, lower eyelid retractors were reinserted into the lower anterior part of the tarsus through a subciliary incision to correct severe lower eyelid entropion (Figures 1a and b).

Discussion

We report 10 children who presented sequentially to a national ophthalmology centre between the ages of 2 and 8 years with congenital or acquired, isolated, unilateral lower motor neuron facial nerve palsies. Four had some degree of entropion, and three had severe entropion with trichiasis requiring surgical correction. These patients had no other obvious visual, neurologic, or general medical problems. Age at onset and presentation, etiology and severity of FNP, and duration of symptomatology were not consistently associated with severity of entropion. The three operated patients were relatively young (ages 2, 3, and 5 years at the time of surgery), but their lid problems may have brought them to medical attention relatively early.

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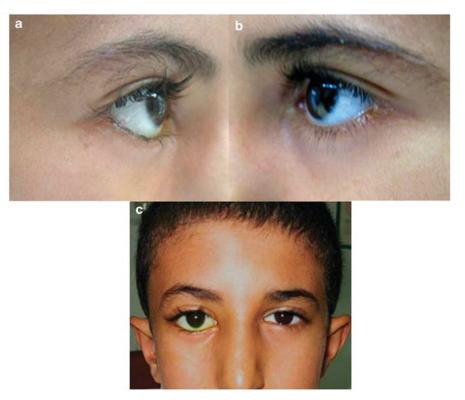


Figure 2 (a) Right eye and (b) left eye of an 8-year-old boy with right-sided facial paralysis causing a mild right lower eyelid entropion with no trichiasis. (c) He had significant right lower eyelid retraction and mild right upper lid retraction in comparison to the unaffected left eye.

Lower eyelid entropion has been described previously in a small group of children with FNP and craniofacial abnormalities.³ Our report shows that entropion is also a frequent observation in children with diverse causes of isolated, unilateral FNP without craniofacial syndromes. The mechanism of lower eyelid changes observed in this study was not clear, although the presence of tight medial and lateral canthal tendons in children might combine with a loss in the balance between the orbicularis oculi and lower eyelid retractors in patients with FNP to contribute to the development of lower eyelid entropion and lid retraction.^{3,12} Alternatively, maldevelopment or repair of the peripheral facial nerve injury might lead to subclinical hyperactivity of previously paralyzed periocular muscles.

A few additional observations are probably valuable regarding the facial appearance of these children. These children did not have eyebrow ptosis even after substantial loss of frontalis function, while adults with FNP generally exhibit clinical brow ptosis.² This may be due to a laxity of the forehead and eyebrows that develops with age.¹² Impairment of facial nerve innervation of the cheek muscles almost certainly contributed to the mild cheek atrophy observed in five of the patients.

There are a few limitations to this study. The patient group is relatively small and was studied retrospectively. Patients were seen at a national ophthalmology centre, which may reflect a referral bias for somewhat worse symptomatology than the typical child with isolated FNP. Finally, neuroimaging and electrodiagnostic studies were not performed on these patients because these were not necessary for the clinical care provided. The observations reported here should be confirmed with a larger group of patients from multiple ethnicities in the future.

Summary

What was known before

• A recent report of children with syndromatic facial nerve paresis implies that entropion may be more common in this setting than ectropion.

What this study adds

- Children with isolated facial nerve paresis seem prone to develop entropion rather than ectropion.
- Entropion reported previously in five syndromic children with facial nerve paresis seems more likely related to patients' age rather than to their congenital syndromes.



Conflict of interest

The authors declare no conflict of interest.

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