Refractive Accommodative Esotropia

Gina M. Rogers, MD and Susannah Q. Longmuir, MD

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Chief Complaint: Eye crossing

History of Present Illness: The patient is a 4-year-old boy who originally presented to an outside ophthalmologist at 2 years of age after his parents noted that his eyes were crossing. The crossing was first noted a few weeks prior to that visit. He was evaluated and placed in glasses, which resulted in partial improvement of the crossing. Over the ensuing 2 years his prescription was updated as needed by the outside provider. His mother seeks a second opinion to ensure that his condition is being managed appropriately. The mother reports that the patient wears his glasses well. She sees occasional crossing with the glasses on. He has never been patched. He is otherwise healthy, growing and developing normally.

Past Medical History:
He was born full term by normal spontaneous vaginal delivery without complications.

Past Surgical History:
None

Family History:
No known strabismus or inherited eye disease

OCULAR EXAMINATION: AGE 2

Visual acuity without correction:
- OD: 20/80
- OS: 20/80

Pupils: Equally round and briskly reactive. No relative afferent pupillary defect.

Stereopsis testing: Unable to test

Strabismus exam: Alternate Cover Testing
- Motility: Full OU
- Esotropia (ET): 30 prism diopters in the distance
- Esotropia (ET): 25 prism diopters at near

Cycloplegic refraction:
- OD: +3.75 D sphere
- OS: +3.50 D + 0.50 D x 090

Slit lamp exam: Within normal limits OU

Dilated fundus exam: Within normal limits OU
OCULAR EXAMINATION: AGE 4

Visual acuity with correction: HOTV

- OD: 20/20
- OS: 20/20

Pupils: Equally round and briskly reactive. No relative afferent pupillary defect.

Stereopsis testing:

- Titmus Fly: stereopsis present
- Animals: 2/3 identified
- Circles: 2/9 identified

Wearing spectacle prescription:

- OD: +4.00 D sphere
- OS: +3.25 D + 0.75 D x 100

Cycloplegic over-refraction:

- OD: +0.50 D sphere
- OS: +0.75 D sphere

Strabismus exam: Alternate Prism Cover Test

Without correction:

<table>
<thead>
<tr>
<th>Right Gaze</th>
<th>Motility</th>
<th>Upgaze: ET 30</th>
<th>Motility</th>
<th>Left Gaze</th>
<th>ET 30</th>
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</thead>
<tbody>
<tr>
<td>ET 30</td>
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<td></td>
<td>ET 30</td>
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<tr>
<td>Downgaze:</td>
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<td>0 0 0</td>
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<tr>
<td>Near:</td>
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<td>0 0 0</td>
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</table>

With correction:

<table>
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<tr>
<th>Right Gaze</th>
<th>Motility</th>
<th>Distance:</th>
<th>Orthotropic</th>
<th>Left Gaze</th>
<th>Orthotropic</th>
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</thead>
<tbody>
<tr>
<td>ET 30</td>
<td>0 0 0</td>
<td>0 0 0</td>
<td>0 0 0</td>
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<tr>
<td>Distance:</td>
<td>0 0 0</td>
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<td>Orthotropic</td>
<td>0 0 0</td>
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</table>

1 Please see http://webeye.ophth.uiowa.edu/eyeforum/tutorials/Bhola-BinocularVision.htm for a more in depth discussion about binocular vision.
**General:** Normal mood and behavior; no nystagmus or abnormal head position

**Slit lamp exam:** Normal OU

**Dilated fundus exam:** Normal macula, vessels and periphery OU. Cup to disc ratio 0.2 OU.

**DIAGNOSIS:** Refractive Accommodative Esotropia

**DISCUSSION:**
Accommodative esotropia is defined as the convergent deviation of the eyes associated with activation of the accommodative reflex. It is classically divided into three categories:

1. Refractive accommodative esotropia (low accommodative convergence/accommodation or AC/A ratio of less than 5),
2. Nonrefractive accommodative esotropia (high AC/A ratio), and
3. Partially accommodative esotropia.
Accommodative esotropia is the most common cause of childhood esotropia.

These esodeviations share common characteristics. All accommodative esodeviations are acquired with onset generally between 6 months and 7 years of age, with average onset at 2.5 years of age. Rare cases have been reported from age 3 months to 11 years old. At the onset, the deviation is usually intermittent, but usually becomes constant in the following weeks to months. It is not uncommon to obtain a history of illness, trauma or fatigue as a precipitating factor for the onset of the deviation. There often is a family history of strabismus. Studies suggest that approximately 77% of patients with accommodative esotropia have a first- or second-degree relative with the same condition. Among first-degree relatives alone, the prevalence of accommodative esotropia is 23%; therefore, it is important to have siblings examined. These esodeviations are frequently associated with amblyopia (Hutchison 2004).

All children with new-onset esotropia require a thorough history, review of systems, and comprehensive examination. The examination must include assessment of vision, ocular motility, strabismus measurements at distance and near and cycloplegic refraction in addition to a general ocular examination. Distance and near measurements are essential to determine the accommodative convergence to accommodation (AC/A) ratio. The AC/A ratio should be investigated if there is good alignment in the distance and greater than 10 prism diopters of increased esodeviation at near in full correction. The next step is to relax accommodation and determine the remaining amount of convergence. We typically hold +3.00 diopter lenses over both eyes and remeasure the esodeviation at near. The esodeviation while looking through +3.00 D lenses is subtracted from the original esodeviation at near without the +3.00 D lenses and the difference is divided by +3.00. If the value is greater than 5, then the patient has a high AC/A ratio. The treatment in this setting may include bifocals if they allow the patient to fuse.

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\text{AC/A ratio} = \frac{\text{Change in deviation}}{\text{Change in lens power}} = \frac{(\text{Esodeviation without additional lens} - \text{esodeviation with lens})}{\text{Additional lens power}}
\]

This case is an example of refractive accommodative esotropia with normal AC/A ratio (in proper correction, the patient is orthophoric in the distance and at near). The mechanism of this type of esodeviation involves 3 factors:

1. Uncorrected hyperopia,
2. Accommodative convergence, and
3. Insufficient fusional divergence

The uncorrected hyperopia forces the patient to exert excessive accommodation to focus images on the retina, thus evoking increased convergence. If the patient’s fusional divergence mechanism is insufficient to compensate the increased convergence, esotropia results. Generally, in refractive accommodative esotropia, the deviation is between 20 and 30 prism diopters, the deviation is
similar in the distance and at near, and the average amount of hyperopia is +4 diopters (range +3.00 to +10.00 diopters) (BCSC Pediatrics and Strabismus, 2007).

The goals of treating this condition are to restore normal ocular alignment, maintain good visual acuity in each eye, and promote good binocular function. Treatment consists of prescribing spectacle correction with the full amount of hyperopic correction as determined by cycloplegic retinoscopy. Significant delay in the initial treatment following the onset of esotropia increases the likelihood that a nonaccommodative component (partially accommodative esotropia) may develop (Mulvihill et al 2000). Of course, if there is any concurrent amblyopia, treatment should be initiated to address this issue.

To the ophthalmologist, the clinical diagnosis and treatment seems straightforward, but to the parent or caretaker the condition may not be as easily understood. Many parents often request surgery, thinking it will be a “quick fix.” For the typical refractive accommodative esotropia patient, the treatment is glasses, not surgery. Strabismus surgery is indicated if optical correction is ineffective in restoring normal ocular alignment. It is important for the eye care provider to take the time to discuss the mechanism of the condition so that the child’s caretaker can comprehend the situation and fully cooperate with the treatment. The importance of full-time glasses wear and amblyopia treatment, if applicable, cannot be over-emphasized. To prevent frustration or misunderstanding, one should mention that the eyes will cross when the child is not wearing glasses because glasses control the deviation and do not cure it. It is also important to convey to the caretaker that this is a long-term condition that requires routine monitoring for amblyopia and that changes in spectacle prescription may be required not only to maintain vision but also alignment. Parents are also frequently concerned over their child’s long-term prognosis. Although the child’s course cannot be predicted at the initial visit or even within years of initiating treatment, it is often helpful to mention that some children can be weaned out of their glasses, while others will require optical correction to maintain alignment throughout adulthood. Contact lenses and refractive surgical procedures may be options as the child nears adulthood.

**Differential diagnosis:**
- Cranial Nerve 6 palsy
- Duane’s syndrome
- Basic, acute, and cyclic esotropia
- Divergence insufficiency
- Spasm of the near synkinetic reflex
### SUMMARY
- Most common type of childhood strabismus
- Prompt recognition and treatment provides the most successful outcomes
- Onset between 6 months and 7 years, average 2.5 years
- Deviation typically measures 20-30 prism diopters
- Similar amount of esodeviation at distance and near (normal AC/A ratio)
- Average hyperopia of +4 diopters

### SIGNS
- Inward deviation of eye
- Full motility
- Improvement of ocular alignment in proper glasses correction

### SYMPTOMS
- Onset in childhood, crossing noted by caretaker
- Diplopia may occur in older children, but will usually disappear as a suppression scotoma develops in the deviating eye

### TREATMENT
- Glasses with full amount of hyperopic treatment as determined by cycloplegic refraction

### References:

### Suggested Citation Format: