

Congenital and Acquired Malposition of Inferior Oblique



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ABSTRACT

Purpose: To report and characterize malposition of the inferior oblique (IO) muscle observed during orbital surgeries for congenital and acquired pathologies.

Study Design: Descriptive case series.

Place and Duration of Study: Al- Shifa Trust Eye Hospital, Rawalpindi from June 2004 to June 2024.

Methods: In this study, all the patients who underwent orbital surgery during the 20 years, by one experienced orbital surgeon, were reviewed. Out of 1200 cases of orbital surgeries four cases were identified for the study. In all four cases, IO malposition was directly observed during orbital surgery. Pre-operative orbital imaging studies could not find the malposition. All surgeries were performed by the same surgeon under general anesthesia. In addition to the planned surgical procedure, an attempt was also made to re-position the abnormal-origin IO.

Results: Out of the four patients with IO malposition, there were two females and two males. Two patients presenting with congenital dermoid cysts had a normal birth history with no history of any surgery or trauma. The other two patients with acquired malposition had a history of trauma and previous surgery.

Conclusion: IO malposition is a rare entity. It is difficult to predict preoperatively so while performing orbital or strabismus surgeries, one should be careful and watchful to prevent damage to ectopic muscles.

Keywords: Inferior Oblique muscle, Congenital, Orbit, Extra ocular muscle.

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INTRODUCTION

IO muscle takes its origin near the anterior margin of the floor of the orbit posterior other im from the orbital surface of the maxilla lateral to the nasolacrimal groove. It inserts on to the globe posteriorly near the inferior border of the lateral rectus muscle. It makes an angle of 51 degrees with the globe in the primary position. The IO muscle externally rotates, abducts, and elevates the eye and is innervated by the oculomotor nerve.¹ Among extraocular muscles, the IO most frequently shows anatomic

variations, but that is also very rare.² In this article, we will discuss malposition of the IO diagnosed during orbital surgeries. The indications for surgeries were both congenital and acquired pathologies.

METHODS

In this study, all the patients who underwent orbital surgery during the last 20 years, by orbital surgeon, were included and their charts were reviewed. The study was approved by the ethical review board of the institution (ERC-22/AST-25). Out of 1200 cases of orbital surgeries, four cases were identified to have IO malposition. In all four cases, the malposition was found during orbital surgery by direct observation. Their medical records were reviewed which included age, gender, history, complete systemic and ophthalmic examination, pre-operative investigations including orbital imaging studies. All surgeries were

Table 1: Four cases with Abnormal origin of the inferior oblique.

Case	Diagnosis	Site of origin of Inferior oblique	
01	5-year-old, male	Congenital medial orbital dermoid	Medial orbital wall around 12 mm behind the orbital rim
02	4-year-old female	Congenital inferomedial orbital dermoid	Arising from the medial lobe of the multilobed dermoid cyst
03	35-year-old male	Traumatic Orbital floor fracture	Middle of the orbital floor from the inferior orbital margin
04	45-year-old male	Traumatic Orbital floor fracture	Middle of the orbital floor from the inferior orbital margin
Normal Position of origin of inferior oblique		From the medial part of the orbital floor near the inferior orbital margin lateral to the lacrimal groove	

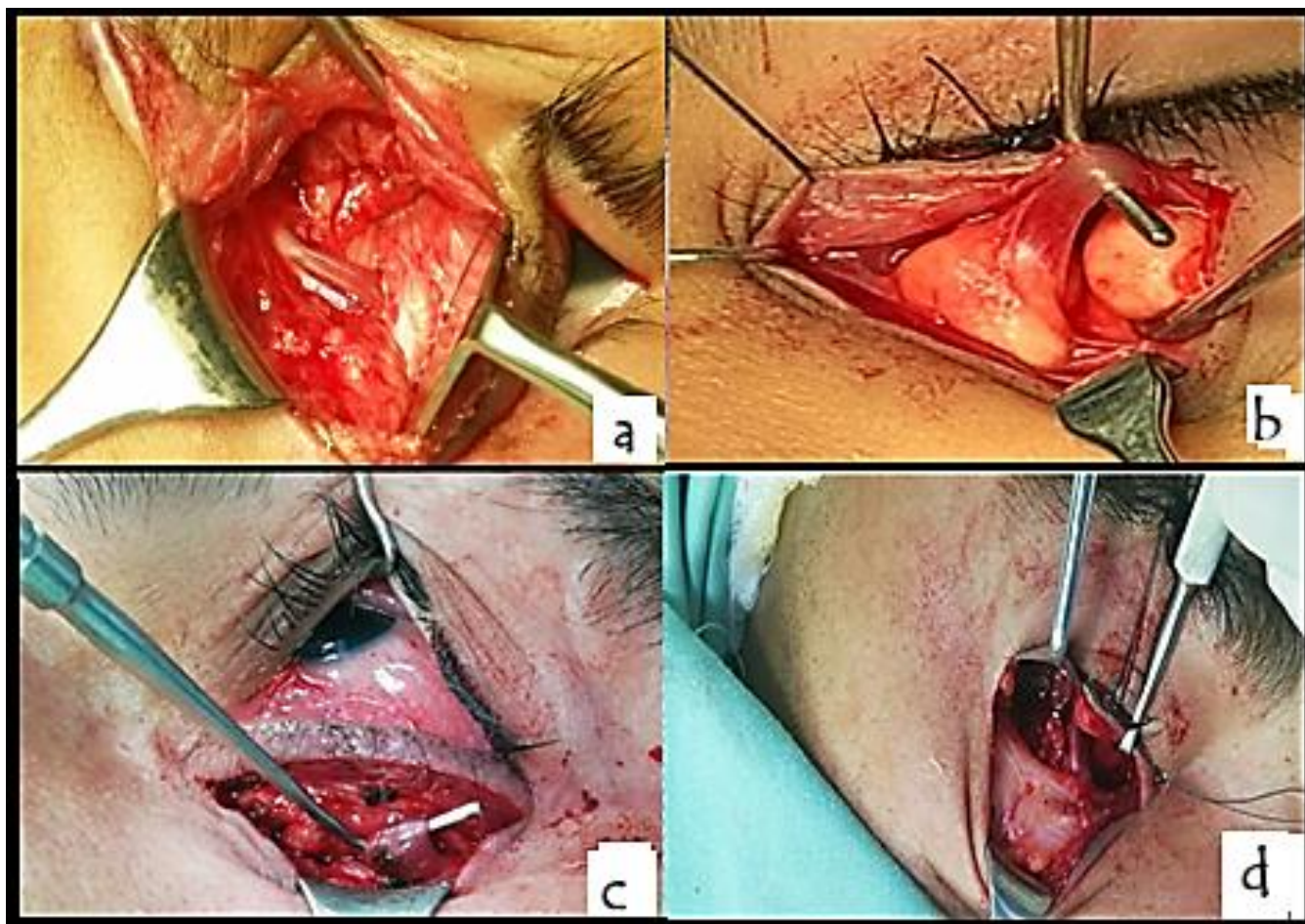


Figure 1(a): Case 1; abnormal origin of inferior oblique from the medial wall. **(b):** Case 2; inferior oblique arising from one of the lobes of the dermoid cyst. **(c) (d):** Case 3 and 4; inferior oblique taking its 'abnormal origin' from the middle of inferior orbital margin.

performed by the same surgeon under general anesthesia. In addition to the planned surgical procedure, an attempt was also made to re-position the abnormal-origin IO.

A five-year-old child presented with a congenital cystic lesion covering the left eyeball (Figure 1a) which was excised. In this case, the IO muscle was seen to be arising from the medial wall of the orbit. The muscle was found to originate 10-12mm behind the medial orbital margin, and going laterally, posteriorly, and downward. The muscle was making

an angle of around 20° to its site of origin. The insertion of the muscle was at its normal site. In another case, during the excision of a trilobed dermoid cyst along the inferomedial wall of the orbit, the IO was seen to be arising from one of the lobes of the orbital dermoid (Figure 1b). The IO was dissected away from the dermoid cyst, and it was re-inserted to its normal anatomical position after excision of the dermoid cyst. In these two cases, congenital IO malposition was seen due to structural developmental anomalies.

Two patients presented to us with a history of trauma followed by orbital floor repair by a maxillofacial surgeon. Both complained of diplopia. On examination, in both cases, the eyeball was rotated upward and medially. We planned the patients for exploration surgery. During surgery, we found identical abnormal anatomy in both cases. The proximal part of the IO was seen to be glued to the inferior orbital margin up to its middle till it left the orbital margin moving towards its insertion to the globe. Although the IO had its normal anatomical origin, the abnormal adhesion of its proximal part to the orbital margin made the IO take its functional origin from the middle of the inferior orbital margin, (figure 1c, 1d) causing the unwanted deviation of the eyeball. The abnormally adherent muscle belly was detached from the orbital margin thus restoring the normal anatomical origin.

RESULTS

Four patients of IO malposition were identified, two females and two males. In all patients, the abnormal origin of the IO was the incidental finding during the surgery, while preoperative imaging studies did not provide any clue to these anatomical variations. Two patients presenting with congenital dermoid cysts had a normal birth history with no history of any surgery or trauma. The other two patients with acquired malposition had a history of trauma and previous surgery. The abnormal positions of the IO in all four cases are outlined in Table 1.

DISCUSSION

IO dysfunction and its clinical manifestations are well-known but anatomical variations are rarely reported. This study retrospectively analyzed the anatomical abnormalities of extraocular muscles in cases undergoing orbital surgeries. Out of four cases with IO malposition, two cases were operated for congenital dermoid with no history of trauma or surgery, while two cases had a history of trauma and previous surgery.

Literature shows various anatomical variations of the IO muscle. Some authors reported double bellies.^{3,4} In another case, a muscular bridge was found between the IO and inferior rectus.⁵ Yet another case report describes IO to bear arising from the orbital apex and inserted near the inferior rectus.⁶

In clinical practice, anatomical variation of the IO is a rare entity or may be under-reported. Accurate preoperative diagnosis and proper repositioning can result in good surgical outcomes. Wang Xiaojun reported a case of IO malposition which was corrected surgically.⁷ Lin in 2012 reported three cases of IO malposition. In one case, the IO tendon was anteriorly displaced, while in the other two cases, the IO bundle was split and fused with the lateral rectus.⁸ Qureshi and Watson reported a unique case of absence of the IO.⁹

In the first two cases, we believe structural pathologies (congenital dermoid) interfered with the extraocular muscle development altering their normal anatomical origin. The abnormal origin was identified during surgical exploration, and the muscle was reinserted to their normal anatomical location. In acquired cases of IO malposition, either the trauma or previous surgery caused the abnormal adhesion resulting in malposition of the IO muscle. In the literature, there are reported cases mostly of abnormal ectopic IO insertion. In this article, we discussed the malposition of IO origin caused by developmental abnormalities and acquired causes.

The strengths of this study include long study duration (20 years) with a large pool of orbital surgeries (n=1200) reviewed by a single, experienced orbital surgeon, ensuring consistency in surgical technique and observation. Direct intraoperative visualization of inferior oblique (IO) malposition, provides reliable documentation of a rare finding and inclusion of both congenital and acquired cases, broadens the clinical spectrum of IO malposition. Moreover, detailed operative management, including attempts at re-positioning, adds practical clinical insight.

Although this condition is rare but small sample size of only 4 cases limit generalizability of the findings. The study was retrospective which can result in selection bias and reliance on operative notes. There was lack of standardized preoperative imaging correlation or postoperative functional outcomes (e.g., motility, diplopia, or long-term results).

CONCLUSION

IO malposition is a rare entity. It is difficult to predict preoperatively so while performing orbital or strabismus surgeries, one should be very careful and watchful to prevent damage to ectopic muscles. Muscle malposition should be vigilantly identified

and ectopic muscles should be repositioned to its normal anatomical location for best postoperative results.

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Conflict of Interest: Authors declared no conflict of interest.

Ethical Approval: The study was approved by the Institutional review board/Ethical review board (ERC-22/AST-25).

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