# Combined Mechanism Glaucoma

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## **Definition:**

Eves that have an occludable angle but fewer signs of angle closure not corroborating with the degree of raised intraocular pressure (IOP) and glaucomatous optic neuropathy are diagnosed as having 'combined mechanism' glaucoma (CMG). In other words, CMG is defined as a combination of both the primary types of

glaucoma, the primary open angle glaucoma (POAG) and the primary angle closure glaucoma (PACG).1,2

## **Clinical Features:**

The patients may present with non-specific symptoms, and it is more often detected incidentally, or as a part of screening programs.

- Age group: Commonly between 50-60 years of age
- Laterality: Usually bilateral.
- Baseline IOP: May vary from 23-28 mmHg.
- Anterior chamber depth: Usually normal centrally or slightly shallow, but may have narrow recess (Figure 1).

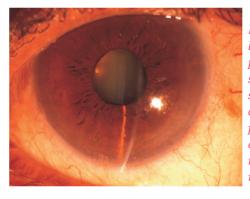


Figure 1 : Slitlamp diffuse photographshowing a mildly shallow anterior chamber, patchy pupillary ruff atrophy and immature senile nuclear sclerosis.

Gonioscopy: There would be an occludable angle (posterior trabecular meshwork not visible in at least 180°) which on manipulative/indentation gonioscopy show up limited signs of angle closure, i.e, goniosynechaie<90° and sparse blotchy pigmentation (Figure 2 & 3).



Figure 2: Goniophotograph showing occludable angle

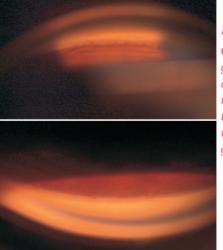


Figure-3: On manipulation gonioscopy, the angle opens up to reveal the posterior trabecular meshwork and few goniosynechaie.

- Pupillary ruff: May show patchy ruff atrophy
- Iris pattern: May be decreased.
- Optic disc: Features consistent with glaucomatous optic neuropathy (Figure 4).



Figure 4: Stereoscopic fundus photograph showing CDR 0.8:1

The sparse goniosynechiae and blotchy pigmentation imply that the patient has had fewer episodes of angle closure attacks in the past than of that which would be expected in a PACG eye with equal amount of optic neuropathy. Therefore, with the anterior segment picture not corroborating with the degree of optic neuropathy to either label a POAG or PACG, this conditionpresents as a combination of both of the primary types, and therefore referred to as the combined mechanism glaucoma'.

## **Investigations**

- Central corneal thickness (Pachymetry)
- Biometry (axial length, anterior chamber depth, lens thickness, white-to-white)
- Angle AS-OCT (angle opening distance, trabecular iris space area, angle recess area, trabecular iris angle, iridotrabecular contact)
- Visual field analyser
- Retinal nerve fibre layer OCT

Sihota et al' in their study on CMG patients showed that their mean corneal diameter was  $12.11 \pm 0.54$  mm, axial length  $23.48 \pm 0.95$  mm, anterior chamber depth was  $3.06 \pm 0.26$  mm and lens thickness  $4.44 \pm 0.29$  mm. On angle anterior segment optical coherence tomography (AS-OCT), the mean angle opening distance (AOD 500) was 0.32 mm and trabecular iris space area (TISA 500) was 0.13 mm2. The mean circumferential iridotrabecular contact (ITC) in CMG eyes was 15% as against PACG whose mean ITC was 87% and POAG o%. All these parameters fell in the mid-rangebetween those of POAG and PACG.

## **Differential Diagnosis:**

The closest differential diagnosis to CMG is primary open angle glaucoma. A CMG could be missed if gonioscopy were not done in presumed POAG. It is important to differentiate CMG from the latter because the pathogenesis and management strategy differfor both. While POAG is primarily due to age related trabecular meshwork changes, CMG has anoverlap of the angle closure component.

Secondary glaucomas such as pseudophakic glaucoma, aphakic glaucoma, post-uveitic glaucoma and pseudoexfoliation glaucoma may have a normal central chamber depth and blotchy pigmentations and goniosynechaieon the angles. Hence, secondary glaucomas must be ruled out before making a diagnosis of CMG.

A confounder by name is the 'mixed mechanism glaucoma' which can be confused for CMG.<sup>3,4</sup> While CMG is a combination of the two primary glaucomas (POAG+PACG), mixed mechanism glaucoma is referred to when there is a

combination of a primary and a secondary glaucoma (or)> 1 secondary glaucomas (eg, POAG + steroid induced glaucoma; post-uveitic + steroid induced glaucoma respectively).

The treatment of secondary and mixed glaucomasdiffer from primary glaucomas in that, they focus not only on the management of intraocular pressures, but also on the control of the inciting factor.

#### **Treatment:**

The management of combined mechanism glaucoma is distinctly different from that of POAG. In contrast to the latter, CMG eyes require a peripheral iridotomy in order to circumvent the existing relative pupillary block owing to their occludable angle status. This should be followed by medical management and measurement of diurnal variation of IOP four weeks later. Based on the highest readingrecorded, a target IOP should be set and treatment titrated accordingly.

Selective laser trabeculoplasty can be attempted in areas of visible trabecular meshwork, however results are not known at present.

In the event of failure to achieve target IOP or disease progressiondespite maximal therapy, filtering surgeries may be proceeded with.

## Take home messages:

- Combined mechanism glaucoma isa distinct entity which is a combination of both the primary type of glaucomas, the POAG and the PACG.
- Eyes which have an occludable angle but goniosynechaie<90° not consistent with PACG fall into this category.
- Management includes a peripheral iridotomy followed by the medical/laser/surgical procedures as per the severity.
- Mixed mechanism glaucoma differs from CMG in being a combination of a primary and a secondary glaucoma (or)>1 secondary glaucomas.

## **References:**

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