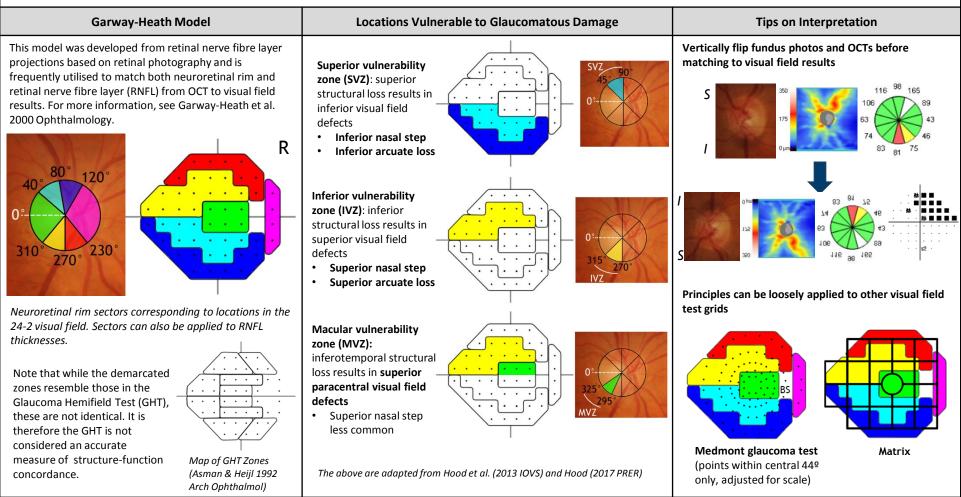


CHAIR-SIDE REFERENCE: STRUCTURE-FUNCTION RELATIONSHIP IN GLAUCOMA

Glaucoma is an optic neuropathy presenting with characteristic loss of the neuroretinal rim, retinal nerve fibre and retinal ganglion cells. These structural findings are frequently observed with corresponding defects on visual field testing, and therefore in conjunction with other clinical signs and examination findings, structure-function concordance should herald high suspicion of glaucoma. This reference provides considerations for evaluating the presence of structure-function concordance in clinical settings. While the focus of this chairside reference is glaucoma, principles may be applied in other optic nerve and inner retinal pathologies resulting in visual field defects.

STRUCTURE-FUNCTION RELATIONSHIP USING THE 24-2 VISUAL FIELD GRID

The 24-2 visual field test grid evaluates the central 54 degrees of the visual field, extending to 30 degrees from fixation nasally, and is the mainstay technique for assessing the visual field in glaucoma.





CHAIR-SIDE REFERENCE: STRUCTURE-FUNCTION RELATIONSHIP IN GLAUCOMA

FACTORS CONTRIBUTING TO VARIATION IN THE STRUCTURE-FUNCTION RELATIONSHIP **Inappropriate Sampling Density of Visual Field Test Points** b. Slit RNFL Defects a. Macula Macular OCT analyses e.g. ganglion cell analysis (GCA) and ganglion cell complex (GCC) and central visual fields testing e.g. Very thin or slit RNFL 10-2 are helpful in determining concordance between structure and function centrally. Paracentral visual field defects do not defects may fall between always appear to match structural loss with other tests due to: visual field test locations. Excessive spacing between test locations Structure and function may • Difficulty visualising associated RNFL loss due to the small MVZ not match in these circumstances, but this 24-2 10-2: Note adjustment of 10-2 locations* Due to the shift in ganglion cells secondary does not mean there is no to Henle's fibres forming the foveal pit, corresponding functional central visual field locations need to be loss. adjusted to match corresponding structural locations.* Note that peripheral 10-2 In cases of slit RNFL defects locations are not covered by macular OCT visible within the macular analyses due to differences in testing area. region, central visual fields testing (e.g. 10-2) may be In this example, macular ganglion cell loss helpful to confirm whether appears as a borderline visual field defect on there is associated Unadiusted 10-2 locations 24-2 and a corresponding superior visual Displaced 10-2 locations functional damage. field defect on 10-2. * Adjustment of visual field locations only applies for inner retinal diseases

ANATOMICAL VARIATIONS RESULTING IN ALTERED RNFL TRAJECTORY

Atypical disc insertions, locations and sizes, frequently observed in high myopia, often present with alterations in RNFL trajectory which will affect the corresponding visual field location. Examples include temporal shift of RNFL bundles and rotation of RNFL trajectory in relation to the foveal location.

Normal Eye (Comparison)	High Myopia	Atypical Optic Disc Insertion		Large Optic Disc to Fovea Angle	
		Torted	Tilted/Obliquely Inserted	High Optic Disc	Low Optic Disc
~7°					
350 175 0μm	350 175 0 μm	³⁵⁰ 175 0 µm	350 175 0 um	350 175 0 µm	350 175 0 μm