# Visual Fields Glaucoma and Beyond

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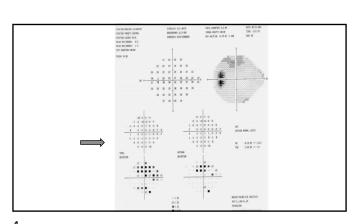
### **Disclosure**

Dr. McSoley has served as a paid consultant for Heru.

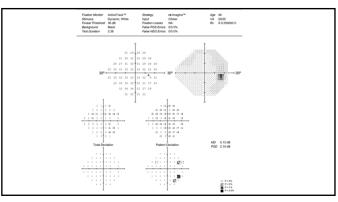
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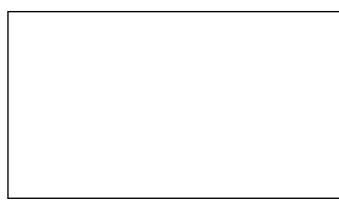
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This course discusses test selection, recognition of artifact and indicators of reliability with the goal of enhancing visual field interpretation. Attention is directed to the anatomic basis, recognition and significance of nonglaucomatous visual field defects. Examples from the Humphrey Field Analyzer and the Heru device will be discussed.



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#### **Brief intro**

Why perform visual fields? **Identify abnormality Explore clinical findings** Address symptoms Follow for change / confirm stability

#### **Brief intro**

Stimuli detected on background **Kinetic Perimetry** isopter Static Perimetry (SAP, WoW) (attenuation of maximum stimulus intensity)

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> Selecting appropriate test Test pattern and strategy Stimulus size Recognizing the quality of information provided by the test (s) Reliability Artifact **Interpreting the test** Recognizing the significance of findings

# Somethings to keep in mind

Importance of instruction and proper positioning Role of patient participation and attention Pattern and sequence of testing

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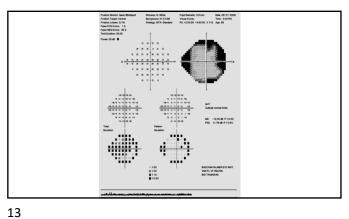
## Somethings to keep in mind

**Foveal Threshold determined** early "primary points" determined Central points determined Test "moves" peripherally More peripheral points tested later

# Somethings to keep in mind Sensitivity tends to decrease with

Longer test time **Eccentricity** Variability increases with **Eccentricity** Test duration Moderate decreased sensitivity Abnl points more variable until they are very poor

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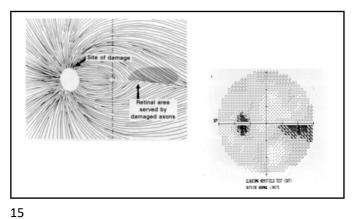


# **Reliability Parameters**

**Fixation False Positive** False negative Continuum not absolute cutoff

Clinical Picture

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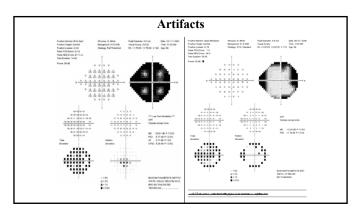


# **Clinical Correlation**

Does this make sense? May need to re-examine or repeat test Consider pattern, sequence of testing Other causes **Test time** 

Patient input

Comfort, understanding, instruction



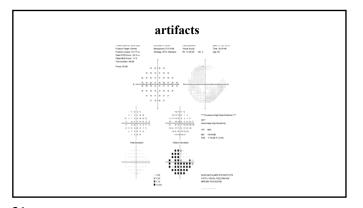
Peripheral points tested later in test "clover-leaf" pattern

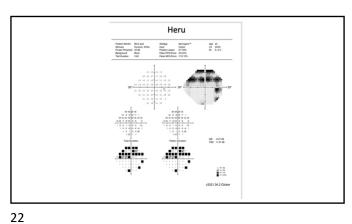
Often high FN

Sometimes longer test times

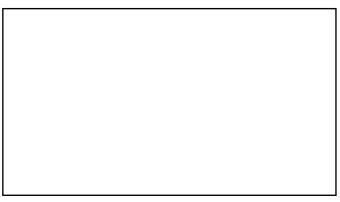
Quality of VF impacts diagnostic value
Introduction of artifact
Reliability
Patient input
Alignment
>Incorrect head or body position may not be identified
by fixation
Media fundus Refractive error
>Wrong Rx

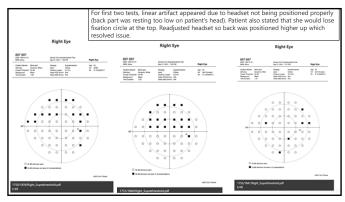
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Visual field testing is vital to management of glaucoma patients

Diagnosis

Staging of damage

In course of follow up

Role of visual fields in non glaucomatous conditions

Addressing symptoms

**Exploring clinical findings** 

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#### Review of visual pathway

Axons of ganglion cells of retina comprise retinal nerve fiber layer

Nerve fiber layer gathers and proceeds posteriorly as optic

Optic nerve proceeds posteriorly to optic chiasm with nasal fibers crossing and temporal fibers remaining ipsilateral

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#### Visual pathway and visual field

Monocular or binocular defects

Respecting vertical

Respecting horizontal (especially nasal defects)

If binocular,

Bitemporal

Homonymous

Location

congruity

#### Visual Fields - why / when

Symptoms

Subjective visual problems

Headaches in some settings

Unexplained levels of vision, acquired color vision deficit

VF defects reported or uncovered w/ confrontation techniques

Some fundus findings

Known or suspected lesions affecting the visual pathway

Drug toxicities

#### Review of visual pathway

Temporal fibers of one eye and nasal fibers of fellow eye proceed posteriorly from the optic chiasm as optic tracts

Optic tracts proceed to lateral geniculate body and synapse in the lateral geniculate nucleus

Neurons proceed posteriorly as optic radiations, inferior fibers passing through temporal lobe, superior fibers passing through parietal lobe toward occipital lobe reaching the primary visual cortex

#### Monocular

Understood clinician recognizes some VF defects will be present in binocular conditions (e.g. glaucoma, ARMD)

Conditions anterior to retina Defects may be more diffuse Media opacities, refractive error

Retinal vascular, nerve fiber layer and some optic nerve defects tend to respect horizontal, likely more apparent nasally

Some optic nerve and macular condition show central field defects Other fundus lesions (RPE / Choroid, retinoschisis) should correlate anatomically

#### Binocular

Bitemporal defects respecting the vertical localizable to crossing of nasal fibers at the optic chiasm

A temporal defect respecting the vertical in one eye and defect involving temporal crossing the vertical, often involving visual acuity in the fellow eye suggests a junctional scotoma

Defects on the same side of each eye (homonymous) suggest lesion posterior to the chiasm on the side contralateral to the VF defect

Lesions of temporal lobe affect superior field

Lesions of parietal lobe affect inferior field

The more similar in shape (congruous) the defect the more posterior Occipital lobe lesions tend to be highly congruous

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# Follow up

A patient with more than one problem

Patient with known

h/o early glaucoma

This VF represents

Non glaucomatous

Interval h/o CVA

R>L

new defect

Can have real decline within normal range

Can have progressive decline of abnormal points

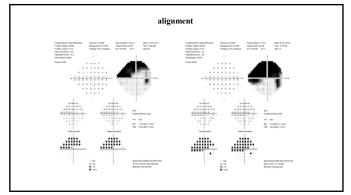
Can have artifacts that simulate change

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Will review and discuss visual field examples reflecting non

glaucomatous defects

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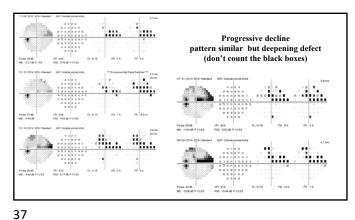


# Follow up

Once statistically abnl need to follow sensitivity values- not rely on patterns or p value symbols

Need range to detect decline Floor / pedestal

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# **Determining Progression** In general

How good your baseline is Accurately reflects status Ideally more than one test to constitute baseline How variable the patient is Can they reliably perform test Are they consistent

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# **Determining Progression In general**

How reproducible the specific test is Does it vary across the spectrum of disease? What the magnitude of change is What is the dynamic range Do you have the ability to detect change? How often the patient is tested

# **Role of Visual Field Testing**

**Exploring suspicion** Diagnosis Staging of condition Monitoring for change **Identifying progression** Confirming stability

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Thank you

All attendees

Meeting planners and sponsors

All scientists, clinicians, staff, patients and those who think and work to improve our ability to care for patients and assess visual function