

Trifocal IOLs

Clinical Outcome at 6 months

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Financial Disclosures

Consultant to:

- Nidek
- Staar Surgical
- PhysiOL

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Clinical Evaluation of Trifocal IOLs Study Design

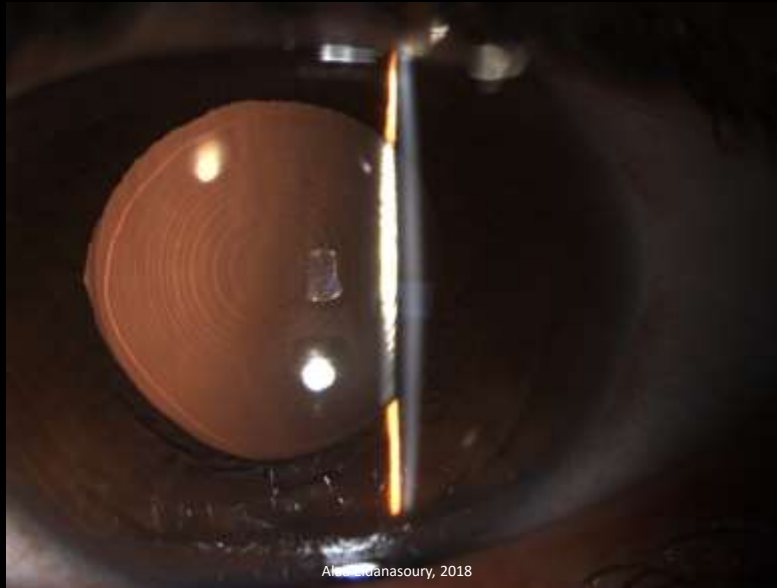
- Prospective Study on consecutive patients.
- 36 patients (72 eyes).
- Inclusion criteria:
 - Bilateral cataract.
 - Stable tear film.
 - No ocular co-morbidity.
- Exclusion criteria:
 - Patients with high corneal irregularities.
 - Significant dry eyes.
 - Conditions that might affect visual rehab. (AMD, glaucoma, amblyopia...)
- Phacoemulsification with Trifocal IOL:
 - Spheric (POD F): 37 eyes
 - Toric (POD FT): 35 eyes

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PODF (PhysIOL)



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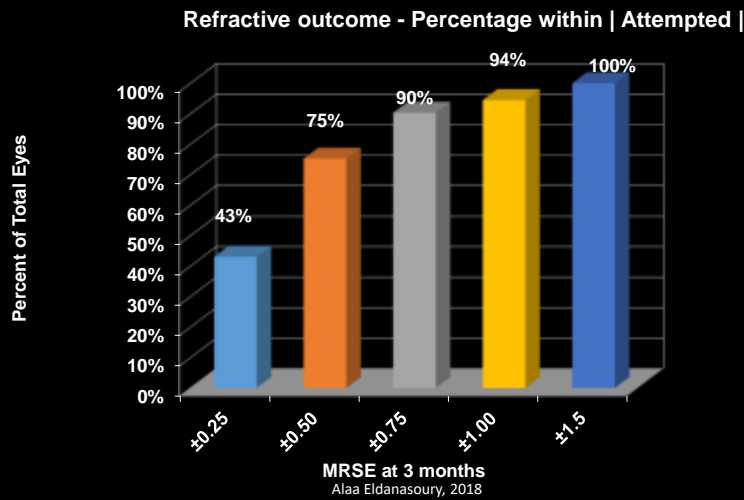
Clinical Evaluation of Trifocal IOLs:

- I. Visual Acuity: "F.I.N.e"
 - Uncorrected.
 - Distance corrected.
- II. Range of pseudoaccommodation:
 - Defocus curve.
- III. Quality of vision:
 - Scatter (OSI, HD Analyzer).
 - Entire eye HOA (OPD scan III)
 - Modulation transfer function (cut off ratio).
 - Contrast Sensitivity (HACSS, CTS).
- IV. Patients' satisfaction:
 - Subjective questionnaire.

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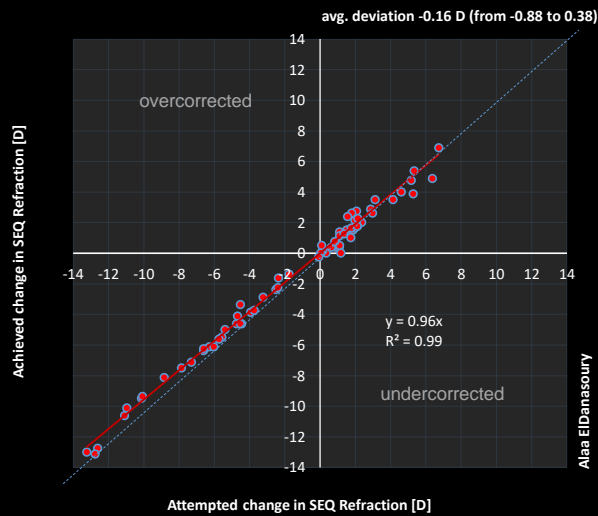
I- Evaluation of Visual Acuity Refractive Outcome at 6 months

68 Eyes – 34 patients



Refractive predictability

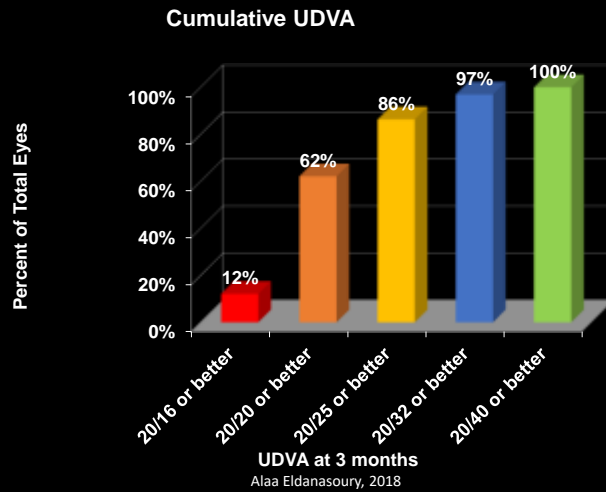
PREDICTABILITY 72 eyes - 3 Mo postOP



I- Evaluation of Visual Acuity

Cumulative UDVA at 6 months (monocular)

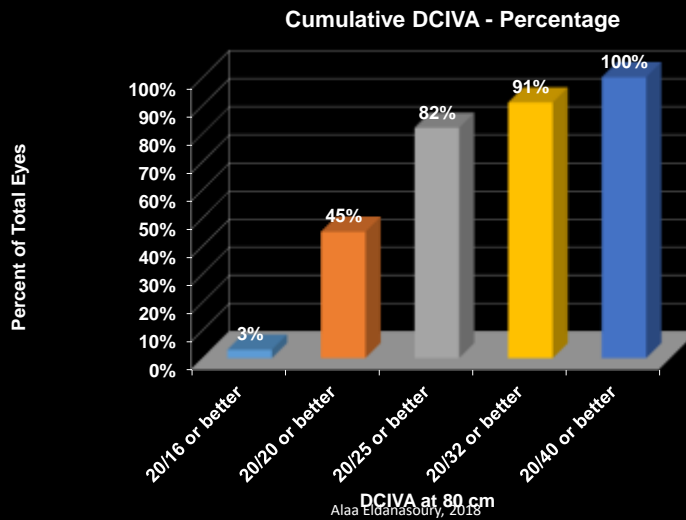
66 Eyes of 33 patients



I- Evaluation of Visual Acuity

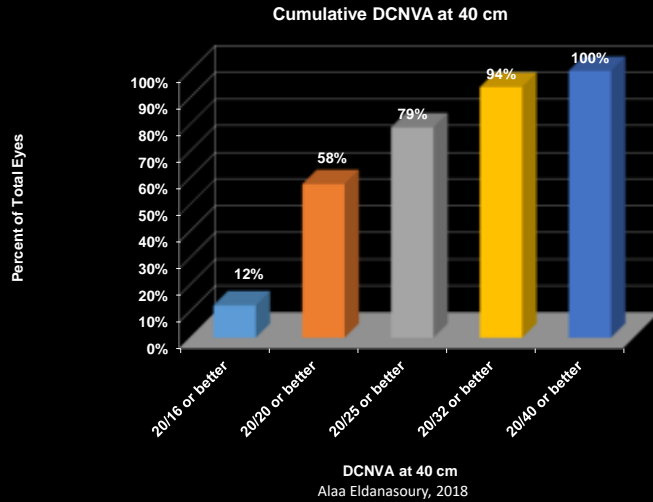
Binocular DCIVA at 80 cm

N = 33 patients

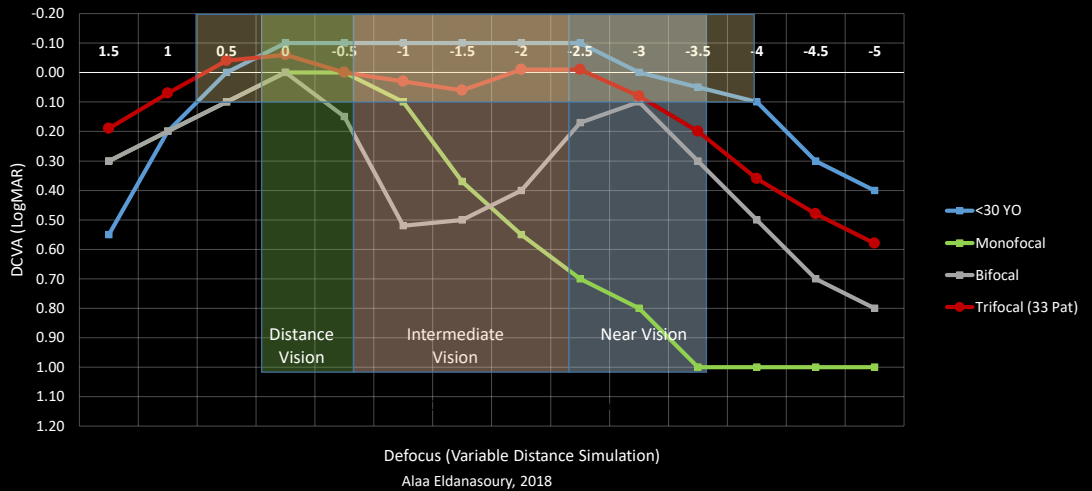


I- Evaluation of Visual Acuity Binocular DCNVA at 40 cm

N = 33 Patients



II- Evaluation of Range of "pseudoaccommodation" Defocus Curve



III- Evaluation of Quality of Vision A- Contrast Sensitivity “HACCS”

Clinical Trial Suite “CTS”:

- Conforms to ANSI* & ISO** guidelines.
- Fully automated system.

*American National Standards Institute

** International Standards Organization



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III- Evaluation of Quality of Vision – Clinical Trial Suite Mesopic Contrast Sensitivity with Glare

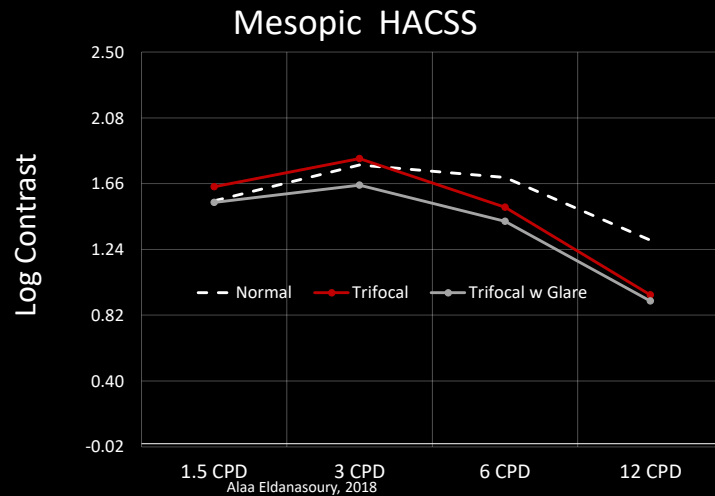


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III- Evaluation of Quality of Vision

Average Contrast Sensitivity Curve

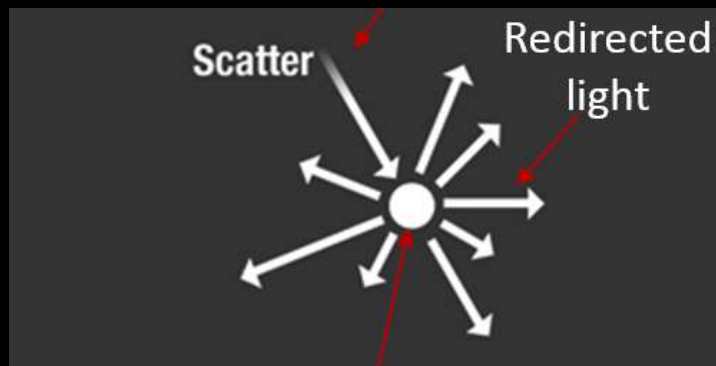
33 Patients @ 6 months



III- Evaluation of Quality of Vision

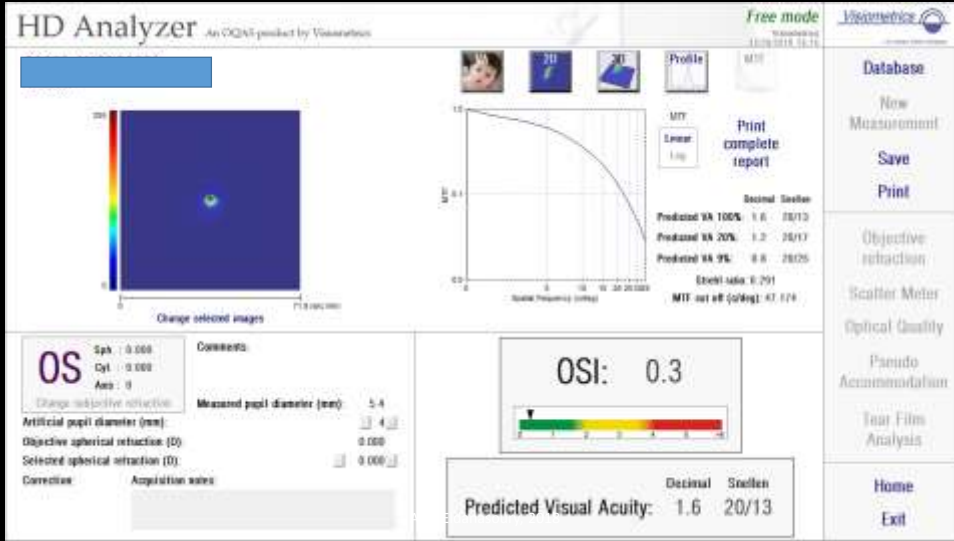
B- Scatter Measurement

- Scatter cannot be measured by aberrometers!



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Objective Scatter Index



IV- Evaluation of Patients Satisfaction Subjective Questionnaire at 6 months

- All patients are very satisfied or satisfied with FINE vision
- 12.5% patients reported halos at night.
- 87.5% patients had non-disturbing photic phenomena when asked.
- No patients were dissatisfied with vision at any distance.
- No patients is using spectacles at any distance.
- All patients would have the same lens again.

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What did we learn:

- Trifocal IOLs give satisfactory far, Intermediate and near vision.
- Night vision phenomena remain but the trade-in is very satisfactory.
- Essentials to achieve good results:
 - Well informed patient.
 - Objective measure of preop quality of vision (OSI, CST)
 - Tear film assessment.
 - Accurate IOL power calculation.
 - Correction of corneal astigmatism.

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Acknowledgment

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Thank You!