

AUTHORS

AUTHORS (LAST NAME, FIRST NAME): Yiu, Patrick¹; Moulton, Eric M.¹; Ploner, Stefan B.²; Lee, ByungKun¹; Husvogt, Lennart²; Maier, Andreas K.²; Spaide, Richard³; Duker, Jay S.⁴; Waheed, Nadia⁴; Fujimoto, James G.¹

INSTITUTIONS (ALL):

1. Department of Electrical Engineering and Computer Science, Massachusetts Institute of Technology, Cambridge, MA, United States.
2. Pattern Recognition Lab, Friedrich-Alexander-Universität Erlangen-Nürnberg, Erlangen, Germany.
3. Vitreous, Retina, Macula Consultants of New York, New York, NY, United States.
4. New England Eye Center at Tufts Medical Center, Boston, MA, United States.

Commercial Relationships Disclosure (Abstract): Patrick Yiu: Commercial Relationship: Code N (No Commercial Relationship) | Eric Moulton: Commercial Relationship: Code N (No Commercial Relationship) | Stefan Ploner: Commercial Relationship: Code N (No Commercial Relationship) | ByungKun Lee: Commercial Relationship: Code N (No Commercial Relationship) | Lennart Husvogt: Commercial Relationship: Code N (No Commercial Relationship) | Andreas Maier: Commercial Relationship: Code N (No Commercial Relationship) | Richard Spaide: Commercial Relationship(s); Topcon Medical Systems, Inc.: Code R (Recipient) ; Topcon Medical Systems, Inc.: Code C (Consultant) | Jay Duker: Commercial Relationship(s); Carl Zeiss Meditec, Inc.: Code F (Financial Support) ; Optovue, Inc.: Code F (Financial Support) ; Topcon Medical Systems, Inc.: Code F (Financial Support) ; Carl Zeiss Meditec, Inc.: Code C (Consultant) ; Optovue, Inc.: Code C (Consultant) ; Topcon Medical Systems, Inc.: Code C (Consultant) | Nadia Waheed: Commercial Relationship(s); MVRP: Code F (Financial Support) ; Janssen: Code C (Consultant) ; Regeneron: Code C (Consultant) ; Genentech: Code C (Consultant) ; Nidek: Code R (Recipient) ; Carl Zeiss Meditec, Inc.: Code R (Recipient) ; Ocudyne: Code C (Consultant) ; Optovue, Inc.: Code R (Recipient) | James Fujimoto: Commercial Relationship(s); Optovue, Inc.: Code I (Personal Financial Interest) ; Royalties from intellectual property owned by MIT and licensed to Carl Zeiss Meditec, Inc.: Code P (Patent) ; Royalties from intellectual property owned by MIT and licensed to Optovue, Inc.: Code P (Patent)

Study Group: (none)

ABSTRACT

TITLE: Smoothed and Resolved Thresholding (SmaRT-) Display: A New OCTA Display Technique to Resolve the Low Flow Ambiguity

ABSTRACT BODY:

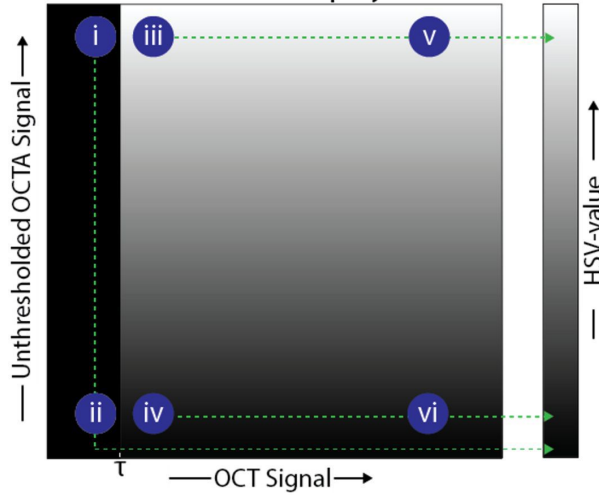
Purpose: OCT angiography (OCTA) provides clinicians with a new perspective on ocular vasculature. However, OCTA is fundamentally different from OCT, and its complex processing steps generate new artifacts that are not present in OCT. The “thresholding” step in OCTA, which removes (makes black) regions of low OCT signal, results in an ambiguity between areas of low OCT signal and areas of low blood flow. In this study we present a new OCTA display technique, SmaRT-Display OCTA, which obviates this ambiguity.

Methods: We propose a mapping scheme based on the hue-saturation-value (HSV) colorspace wherein the OCT signal is mapped to a sigmoidal curve in the value coordinate, the unthresholded OCTA signal is mapped to the saturation coordinate, and the hue coordinate is fixed at red.

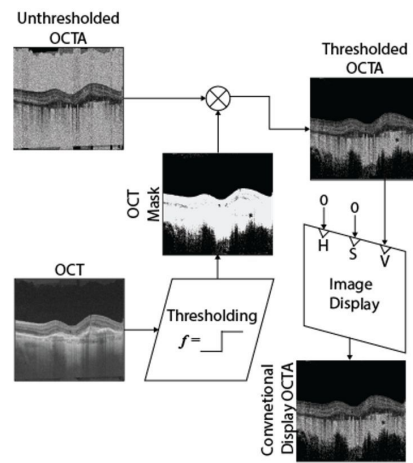
Results: SmaRT-Display removes low flow ambiguities in both cross-sectional and en face views.

Conclusions: SmaRT-Display has the potential to reduce misinterpretation of OCTA images and is a first step in expanding the standards of how OCTA data can be presented.

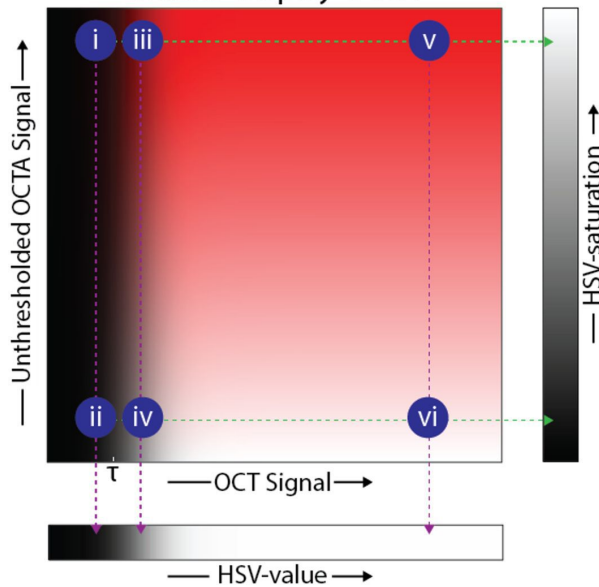
A.1 Conventional Display OCTA



A.2



B.1 Smart-Display OCTA



B.2

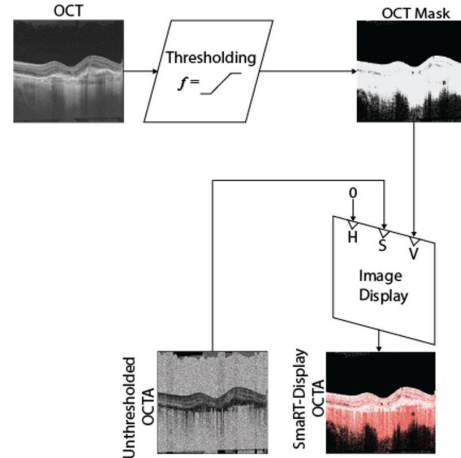


Figure 1: Conventional display vs. Smart-Display OCTA. (A.1) In the conventional display OCTA, i and ii, which are below the threshold τ , are mapped to same value; this value is essentially indistinguishable from the value to which iv and vi are mapped; finally, iii and v are mapped to the same value. These non-invertible mappings induce ambiguities. (B.1) Using Smart-Display, the mappings to the display color space are invertible. (A.2, B.2) Signal flow graphs for the conventional display and Smart-Display schemes, respectively.

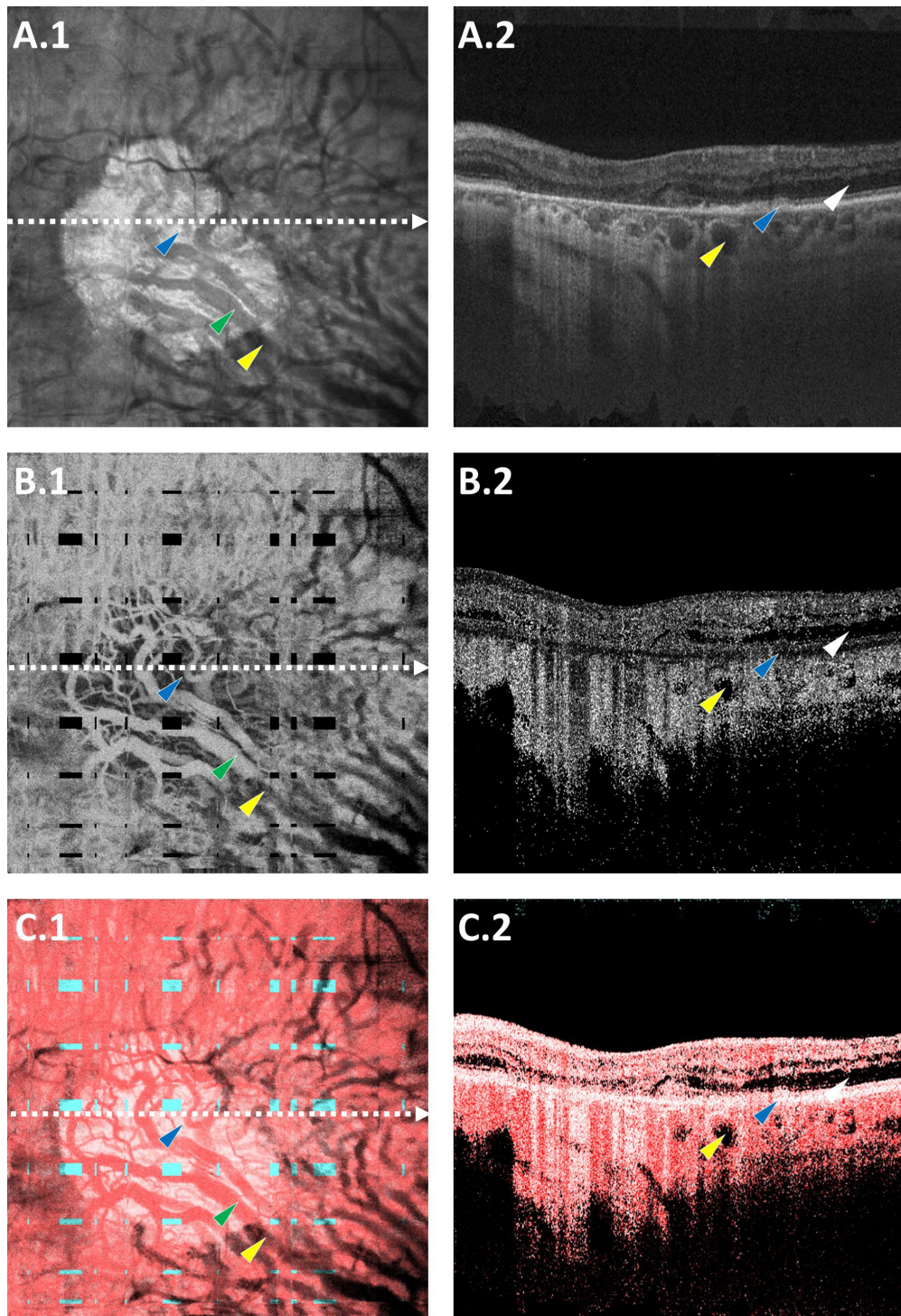


Figure 2: (A) OCT, (B) conventional display OCTA, and (C) SmART-Display OCTA from an eye with geographic atrophy. First column: en face projections of 110µm slab; the black (teal) rectangles in B.1 (C.1) are motion artifacts. Second column: B-scans from dashed white lines in first column. Blue arrows point to areas of low flow with OCT signal above the threshold (en face: intervascular stroma; B-scan: retinal pigment epithelium). Green arrows point to area of high flow with OCT signal above the threshold (en face: choroidal vessel in region of atrophy). Yellow arrows point to areas of high flow with OCT signal below the threshold (en face and B-scan: choroidal vessels outside region of atrophy). White arrows point to area of low flow with OCT signal below the threshold (B-scan: outer nuclear layer).

DETAILS

PRESENTATION TYPE: #1 Paper, #2 Poster

CURRENT REVIEWING CODE: 2510 imaging: image processing and analysis methodologies - MOI

CURRENT SECTION: Multidisciplinary Ophthalmic Imaging Cross-sectional Group

Clinical Trial Registration (Abstract): No

Other Registry Site (Abstract): (none)

Registration Number (Abstract): (none)

Date Trial was Registered (MM/DD/YYYY) (Abstract): (none)

Date Trial Began (MM/DD/YYYY) (Abstract): (none)

Grant Support (Abstract): Yes

Support Detail (Abstract): NIH: 5-R01-EY011289-28; AFOSR: FA9550-15-1-0473, FA9550-10-1-0551; MVRF

TRAVEL GRANTS and AWARDS APPLICATIONS

AWARDS:

AFFIRMATIONS

Affirmations: Affirmation of compliance with ARVO's Statement for Use of Animals.

Affirmations: Affirmation of compliance with ARVO's Statement for Use of Human Subjects and/or Declaration of Helsinki.

Affirmations: Affirmation that submission of this abstract has been approved by the Principal Investigator.

Affirmations: Affirmation to pay Annual Meeting's full registration fee.

Affirmations: Affirmation to present same work as abstract submission.

Affirmations: Affirmation that abstract data/conclusions have not been published; not redundant with other submissions from same investigators.

Affirmations: Affirmation to reveal essential structure, novel compound elements, or identify new gene compounds.

Affirmations: Affirmation of copyright transfer from each author to ARVO, or certification of public domain abstract.

Affirmations: Affirmation of compliance with ARVO policy on registering clinical trials.