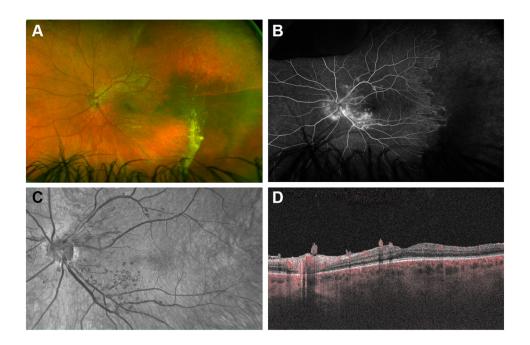
- 43. Reiter GS, Told R, Schranz M, et al. Subretinal drusenoid deposits and photoreceptor loss detecting global and local progression of geographic atrophy by SD-OCT imaging. *Invest* Ophthalmol Vis Sci. 2020;61(6):11.
- 44. Thiele S, Nadal J, Pfau M, et al. Prognostic value of intermediate age-related macular degeneration phenotypes for geographic atrophy progression. *Br J Ophthalmol*. 2021;105(2): 239–245.
- 45. Domalpally A, Danis R, Agron E, et al. Evaluation of geographic atrophy from color photographs and fundus auto-
- fluorescence images: Age-Related Eye Disease Study 2 report number 11. *Ophthalmology*. 2016;123(11):2401–2407.
- 46. Yaspan BL, Williams DF, Holz FG, et al. Targeting factor D of the alternative complement pathway reduces geographic atrophy progression secondary to age-related macular degeneration. Sci Transl Med. 2017;9(395):eaaf1443.
- Khanifar AA, Lederer DE, Ghodasra JH, et al. Comparison of color fundus photographs and fundus autofluorescence images in measuring geographic atrophy area. *Retina*. 2012;32(9): 1884–1891.

## **Pictures & Perspectives**



## Preretinal Microvascular Tufts Associated with Dyskeratosis Congenita

A 16-year-old boy with dyskeratosis congenita (DC), a very rare polymorphous inherited telomeropathy, was referred for bilateral large peripheral temporal areas of retinal nonperfusion (**A**). Absence of associated retinal neovascularization was confirmed by fluorescein angiography (FA) (**B**). Although peripheral retinal ischemia is known in DC, unexpected numerous reddish, rounded, preretinal microvascular tufts were also observed in the inferonasal macula. They were distinct on the near-infrared image (**C**) and localized above the retinal surface on OCT B-scans (**D**). Blood flow within these microvascular abnormalities was noted on FA (**B**) and OCT angiography B-scan (**D**) (Magnified version of Fig **A-D** is available online at www.aaojournal.org).

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