

















DIABETIC RETINOPATHY

Utility of WF/UWF imaging

- Live scan OCT to look for neo and determine PVD status in high risk eyes
- Determine PVD status
 - Complete PVD = lower risk for neo growth and resultant vitreoretinal traction
- Detect vitreoretinal traction & monitor for progression
- Detect retinal tears
- Montage OCTA
 - More accurate & efficient staging
 - Detection and quantification of nonperfusion to
 - determine likelihood of neo/risk for progression
 - Earlier PDR detection

































	Ocular Ischemic Syndrome (OIS)	Diabetic Retinopathy	CRVO
Symmetry	asymmetric	symmetric	asymmetric
Heme Shape	blot	blot	flame > blot
Heme Location	Midperiphery/periphery > post pole	Usually more concentrated within the post pole	Usually more concentrated within the post pole
Retinal Vessels	Veins dilated and non-tortuous, arteriolar attenuation	Veins dilated & beaded	Veins dilated and tortuous
ONH	Normal to pale	Normal	Edematous
Mac Edema	Rare	Common, usually focal (decentered) with exudate	Common (central & diffuse CME)
Oncot	Insidious	Insidious	Acute





























RETINAL VEIN OCCLUSION

RVO Zebra Work-Up

- If young patient (<50), BRVO that did not occur at a A/V crossing, no history of HTN, or bilateral RVO consider more extensive systemic workup:
- Causes of retinochoroiditis or retinal vasculitis: sarcoid, syphilis, SLE, TB, lyme, AIDS/CMV
- <u>Hyperviscosity states</u>: leukemia, Waldenström's macroglobulinemia, polycythemia, lymphoma, multiple myeloma, cryoglobulinemia .
- Hypercoagulable states and thrombophilia: factor V Leiden, hyperhomocysteinemia, protein C deficiency, antithrombin III, activated protein C resistance, sickle cell, antiphospholipid syndrome, anticardiolipin antibodies •
- Abnormal platelet function Oral contraceptive/diuretic use
- Recommend: BP, A1c, CBC with diff & plat count, PT/aPTT, ESR, INR, lipid profile, homocysteine levels, ANA, FTA-ABS/RPR, QuantiFERON TB gold or PPD, hemoglobin & serum protein electrophoresis, antiphospholipid antibodies, protein C and S levels, factor V Leiden mutation,, antithrombin III mutation, prothrombin G20210A mutation.

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С	CHRPE-like Lesions in FAP (Familial Adenomatous Polyposis)					
		Gardner's Syndrome	CHRPE			
	Appearance	Small: Identical to CHRPE Large: Oval with tail	Flat, round-oval, jet black			
	Bilaterality	Common (86% cases)	Very rare (5% cases)			
	Multiple quadrants	Common	Rare			
			p/			

Peripheral retinal pathology & Rheg RD



Utility of WF/UWF imaging

- Detect retinal breaks : Document size and extent of RDs
- . Monitor post repair

- Find holes within lattice Differentiate vitreoretinal tuft vs tear
- Differentiate retinoschisis vs RD Posterior Shafer's sign

Atrophic hole

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RETINAL DETACHMENT

- High incidence of asymptomatic retinal breaks (<u>8.4%</u>) and lattice (14.5%) in fellow eyes of RRD patients
- Fellow eye RD more common in pseudophakic and myopic patients
- Despite prophylactic barrier laser of fellow eye breaks, 5.8% of treated eyes still developed RRD



Mitry D, et al. The fellow eye in RD: findings from the Scottish RD Study. Br J Ophthalmol 2012.

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ACUTE ONSET METAMORPHOPSIA

- 38yo American Indian female
- CC: new onset floaters and slightly distorted vision OS x 3 days
- Oc Hx: LASIK OU 2010, pt reports there were "old scars" seen in the back of the eyes when she had LASIK done



- BCVA :
- OD: 20/20
- OS: 20/25+2

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THE "TAKE HOME" MESSAGE

- Eyes with predominantly peripheral findings have a greater risk for DR progression and development of PDR
- OCT useful in the detection of retinal breaks and early tractional RD
- Montage OCTA useful in detection of early PDR and quantification of peripheral nonperfusion
- OCT useful to differentiate lamellar from full thickness breaks, and to monitor SRF surrounding breaks