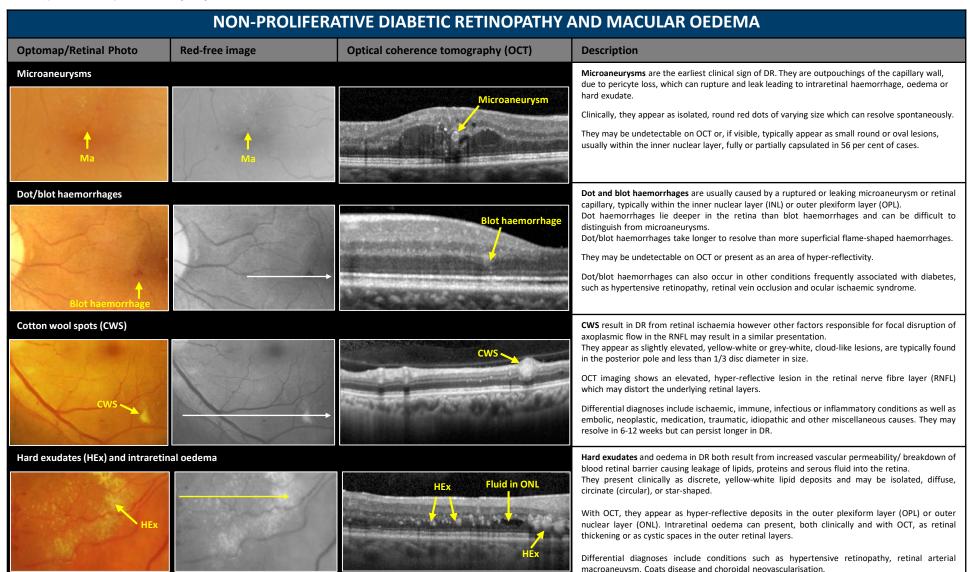


©CFEH

Diabetic retinopathy (DR) is a retinal microvascular disease which occurs in an individual with diabetes. Typical retinal microvascular lesions display a characteristic evolution and progression(see table below). Risk factors include duration of diabetes, hyperglycaemia, systemic hypertension, renal disease, hyperlipidaemia, sudden lowering of glycaemic levels, pregnancy, ethnicity and genetic factors. The International Clinical Diabetic Retinopathy and Diabetic Macular Edema Disease Severity Scales (overleaf) provide a clinically useful scale for grading the level of DR and macular oedema.



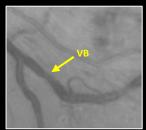


OTHER NON-PROLIFERATIVE DR LESIONS

Venous beading (VB) - colour and red-free photography

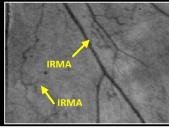
Intraretinal microvascular abnormalities (IRMA) – colour and red-free photography





Venous Beading is a venous calibre irregularity which occurs in areas of severe retinal hypoxia. A sausage-link appearance occurs in severe cases. Other calibre changes include dilation, reduplication and loops.





Intraretinal microvascular abnormalities (IRMA) are abnormal intraretinal shunts which appear as branching or dilation of capillaries within the retina in areas of poor retinal perfusion. They have a similar appearance to NV but with slightly larger vessel calibre. They are a precursor to NV which may form in close proximity.

PROLIFERATIVE DIABETIC RETINOPATHY

Optomap/Retinal Photo

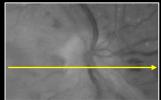
Red-free image

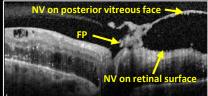
Optical coherence tomography (OCT)

Description

Neovascularisation (NV) and fibrous proliferation (FP)







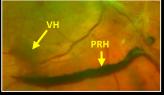
Neovascularisation (NV) appears as new vessels which loop back around or form a disorganised net, distinguishing them from normal capillaries. They are on the surface of the internal limiting membrane (ILM) or posterior hyaloid face of the vitreous and occur at the border between healthy retina and areas of capillary non-perfusion (retinal ischaemia).

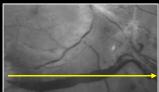
They are prone to bleeding, resulting in pre-retinal (PRH) or vitreous haemorrhage (VH).

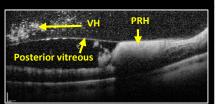
Dynamic interaction between NV and the vitreous can lead to an inflammatory response and subsequent fibrous proliferation (FP).

Any pre-retinal or vitreous haemorrhage should be considered as NV until proven otherwise. NV of the disc (NVD) describes new vessels on or within 1 disc diameter of the disc as opposed to NV elsewhere (NVE).

Pre-retinal haemorrhage (PRH) and vitreous haemorrhage (VH)





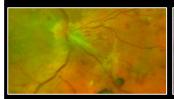


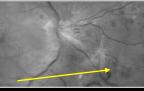
PRH or VH can occur when new vessels bleed. This may occur when the new vessels proliferate along the posterior surface of the relatively mobile vitreous, causing traction on the new vessels, particularly where there is a strong adherence between the vitreous and the retina at the area of NV or FP.

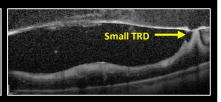
PRH may present as a D-shaped or boat-shaped haemorrhage trapped between the ILM and the posterior hyaloid face of the vitreous, although they may appear linear, blot-like or arcuate.

VH will appear as a reddish or greyish area of haze obscuring the underlying retinal detail. OCT assists in identifying the location of the haemorrhage (which appears hyper-reflective).

Tractional retinal detachment (TRD)







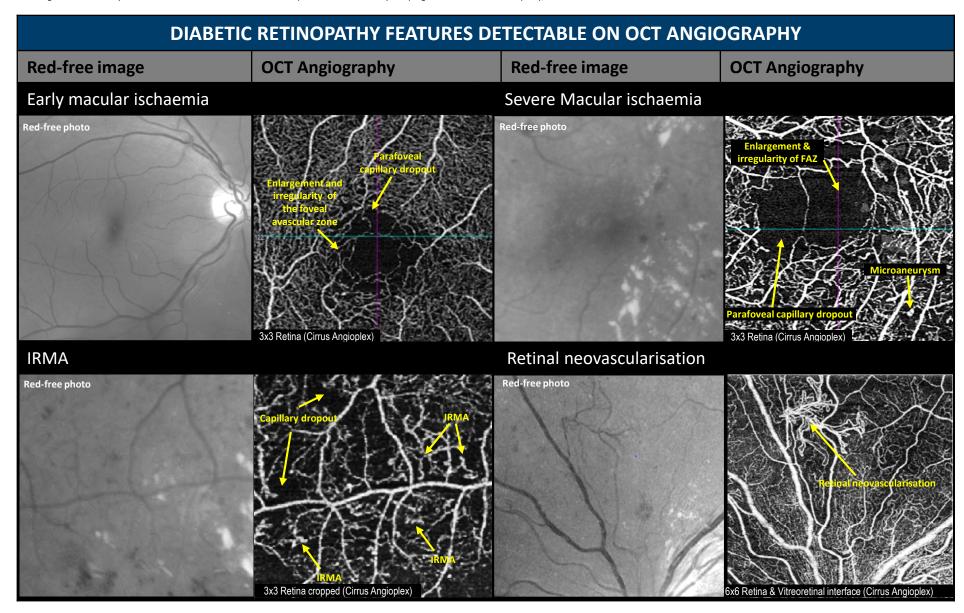
Retinal folds or tractional retinal detachment (TRD) can occur if the vitreous is adherent to the retina in an area of fibrovascular scar formation. These changes are more likely to occur along the major vascular arcades.

TRDs are concave and usually progress slowly, however a hole can form in the detached retina leading to a combined TRD and rhegmatogenous retinal detachment.

Clinically, TRD will be associated with NV and FP and appear elevated.



OCT angiography is a non-invasive technology that detects the movement of red blood cells through the retinal and vessels to produce an image of the vasculature without the injection of dye. In diabetes, OCT angiography can show features of macular or retinal ischaemia including foveal avascular zone (FAZ) enlargement, parafoveal capillary dropout, IRMA and neovascularisation. Microaneurysms can be detected in some cases, but not others where there is poor blood flow through the microaneurysm. An increased FAZ has been shown to be a predictor of the severity and progression of diabetic retinopathy,.





International Clinical Diabetic Retinopathy Disease Severity Scale	
DIABETIC RETINOPATHY STAGE	OPHTHALMOSCOPY FINDINGS
No apparent retinopathy	No abnormalities
Mild NPDR	Microaneurysms only
Moderate NPDR	More than just microaneurysms but less than severe NPDR
Severe NPDR	 Any one of the following (and NO signs of PDR): More than 20 intraretinal haemorrhages in each of 4 quadrants Definite VB in 2+ quadrants Prominent IRMA in 1+ quadrant
Proliferative DR	One of the following: Neovascularisation, vitreous/pre-retinal haemorrhage

International Clinical Diabetic Macular Edema Disease Severity Scale	
MACULAR OEDEMA STAGE	OPHTHALMOSCOPY FINDINGS
Absent	No retinal thickening or hard exudates in the posterior pole
Mild (non-centre involving*) Can occur at any level of DR	Some retinal thickening or hard exudates in posterior pole but distant from the macula
Moderate (centre approaching*) Can occur at any level of DR	Retinal thickening or hard exudates approaching the centre of the macula but not involving the centre
Severe (centre involving*) Can occur at any level of DR	Retinal thickening or hard exudates involving centre of the macula

*Modified by CFEH to reflect International Council of Ophthalmology guidelines (2017) which define centre-involved macular oedema as thickening within the central 1000µm of the macula