

Commentary

## The Role of Plus Disease Detection in Pediatric Ophthalmology

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## **DESCRIPTION**

Retinopathy of Prematurity (ROP) is a potentially blinding disease that affects premature infants, particularly those born before 31 weeks of gestation or with a birth weight of less than 1500 grams. Among the various stages and manifestations of ROP, identifying and managing plus disease is important in preventing vision loss and ensuring optimal visual outcomes for affected infants.

Plus disease refers to the vascular dilation and tortuosity observed in the posterior pole of the eye in infants with ROP. It is a critical clinical sign indicative of severe ROP requiring urgent intervention to prevent retinal detachment and subsequent blindness. Plus disease is diagnosed based on the examination of the retina using indirect ophthalmoscopy or wide-field retinal imaging, often supplemented with digital imaging technologies. The accurate detection of plus disease is essential for timely intervention and appropriate management strategies. Early identification allows for prompt treatment, typically through laser therapy or anti-VEGF injections, to prevent progression to advanced stages of ROP and associated complications such as retinal detachment and permanent vision loss. However, the diagnosis of plus disease can be challenging and subjective, requiring skilled ophthalmologists experienced in pediatric retinal examinations.

The caliber of the blood vessels in the posterior pole of the retina is compared to that of the adjacent optic nerve. Dilated vessels that are larger than the optic nerve are indicative of plus disease. The degree of tortuosity or twisting of the retinal blood vessels is evaluated. Increased tortuosity, particularly in the presence of vascular dilation, suggests the presence of plus disease.

Serial examinations are essential to monitor the progression of ROP and detect the development of plus disease. Changes in vascular appearance over time can indicate worsening disease and the need for intervention.

Plus disease primarily affects the posterior pole of the retina, but it can extend into the mid-periphery in severe cases. The extent and location of vascular abnormalities help assess the severity of ROP and guide treatment decisions. To enhance the accuracy and consistency of plus disease detection, ophthalmologists may utilize imaging modalities such as wide-field fundus photography, fluorescein angiography, and Optical Coherence Tomography (OCT). These technologies provide detailed visualization of the retinal vasculature and aid in documenting disease progression over time.

Despite advances in imaging and diagnostic techniques, the subjective nature of plus disease assessment remains a challenge. Interobserver variability among clinicians can lead to discrepancies in diagnosis and treatment decisions. Standardized protocols and training programs aimed at improving consistency in ROP evaluation are significant for reducing diagnostic errors and optimizing patient outcomes.

Moreover, the interpretation of plus disease in the context of other clinical findings, such as stage of ROP, zone of disease, and presence of extraretinal fibrovascular proliferation, is essential for comprehensive management planning. Collaborative approaches involving neonatologists, pediatricians, and ophthalmologists are necessary to provide multidisciplinary care and address the complex needs of infants with ROP.

In addition to accurate diagnosis, ongoing monitoring and follow-up are essential components of ROP management. Infants diagnosed with plus disease require frequent evaluations to assess treatment response and monitor for disease recurrence. Close coordination between healthcare providers ensures timely adjustments to treatment plans and facilitates continuity of care.

Research efforts aimed at elucidating the pathophysiology of ROP and identifying novel biomarkers for disease progression hold potential for improving diagnostic accuracy and prognostication. Biomolecular markers present in the vitreous or peripheral blood may offer insights into the underlying mechanisms driving ROP and help predict disease severity and treatment response.

Furthermore, advancements in Artificial Intelligence (AI) and machine learning present opportunities for automated image analysis and decision support in ROP diagnosis. AI algorithms

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trained on large datasets of retinal images can assist clinicians in detecting plus disease and predicting disease progression with high sensitivity and specificity. Integrating AI-driven tools into

clinical practice has the potential to enhance diagnostic precision and streamline patient care.