

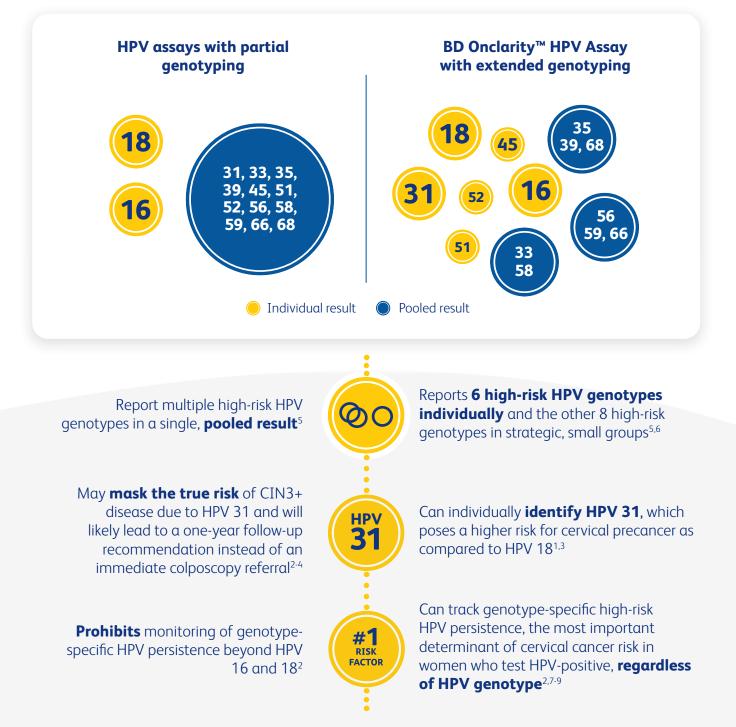
Shape the future of cervical cancer screening: Identify HPV 31

The BD Onclarity[™] HPV Assay with extended genotyping is the only FDA-approved HPV test that individually identifies HPV 31, which poses a higher risk for cervical precancer as compared to HPV 18 and should be managed similarly.¹⁻⁴



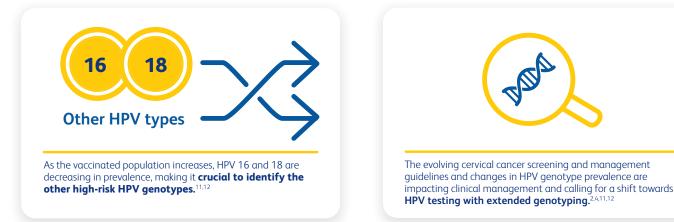
Extended genotyping brings value to clinical decision-making and patient care

Get specific, actionable insights on an extended set of HPV genotypes



The **BD Onclarity[™] HPV Assay with extended genotyping** allows for a **more precise, accurate** way to measure a woman's risk for developing cervical precancer compared to an assay with partial genotyping.^{2,3,7-10}

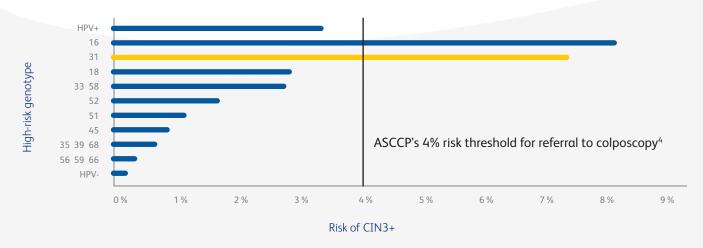
Adapt to the evolving landscape of cervical cancer screening



HPV 31 identification matters. Extended genotyping is critical.

Following the American Society for Colposcopy and Cervical Pathology (ASCCP) principle of **"similar management for similar risk"**, women with an immediate risk for CIN3+ disease above 4% should be referred to colposcopy.⁴

In the BD Onclarity[™] HPV Assay FDA trial, women 25 years and older with HPV 31 and normal cytology had an immediate risk for CIN3+ similar to HPV 16 that exceeds the colposcopy referral threshold of 4% recommended by ASCCP management guidelines.^{3,4}



Risk of CIN3+ by HPV type in women ≥ 25 years with normal cytology

Created from information provided in Stoler MH et al. Gynecol Oncol. 2019,153(1):26-33.

Only an HPV assay with extended genotyping can individually identify high-risk HPV genotypes beyond HPV 16 and 18, including HPV 31²

Shape the future of cervical cancer screening with the BD Onclarity™ HPV Assay



Provides an individual result for HPV 31, which poses a higher risk for cervical precancer as compared to HPV 18^{1,3}



Individually identifies 6 high-risk HPV genotypes to allow for genotype-specific HPV persistence tracking, the most important determinant of cervical cancer risk in women who test HPV-positive^{2,5-9}



The only FDA-approved HPV assay that offers extended genotyping^{5,6}



FDA approved for the three most-common cervical cancer screening paradigms offering the flexibility you need to adopt to changing screening guidelines^{5,6}



Consensus guidelines favor a personalized risk-based management of cervical cancer screening results with HPV testing as the foundation for risk-estimation⁴

Improve your cervical cancer screening: Identify HPV 31

Let's shape the future of women's health. Together and now.

For more information about BD Onclarity™ HPV Assay, please visit <u>womens-health-solutions.bd.com</u>

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