

Assess Transcriptionally Active HPV Biomarkers in Head and Neck Cancer Biopsies

Detect HPV E6/E7 mRNA expression with tissue morphological context

- High Sensitivity and Specificity—single molecule detection with single-target specificity by proprietary probe design and detection technology
- Broadest Spectrum—with flexibility of individual genotyping, pooled high-risk and low-risk panels (HPV-HR7, HPV-HR18, HPV-LR6) and custom pooled panels
- Robust and Easy Methodology—scalable with automation for routine analysis

RNA SCOPE[®] DELIVERS
MD+MC
IN A SINGLE ASSAY

Molecular Detection visualizes and quantifies expressed genes at single-cell resolution.

Morphological Context localizes expressed genes and reveals tissue architecture.

Detection of Gold-standard E6/E7 Oncogene Transcripts Using RNAscope[®] ISH

Evidence for transcriptional active of the viral oncogenes E6/E7 is regarded as the gold standard for presence of clinically relevant high-risk human papillomavirus (HPV), but detection of E6/E7 mRNA can be challenging using conventional techniques (Bishop *et al.*, 2013). As a causal agent in head and neck squamous cell carcinoma (HNSCC), it is critical that the detection method enable pathologist review of tissue morphology and be of the highest specificity and sensitivity for accurate assessment of within the tissue microenvironment of FFPE specimens (Figure 1, 2). RNAscope HPV Biomarker Detection Reagents and its proprietary “double Z” oligonucleotide probes specific for each subtype E6/E7 mRNA enable high specificity detection of viral transcripts in routine FFPE tumor biopsies.

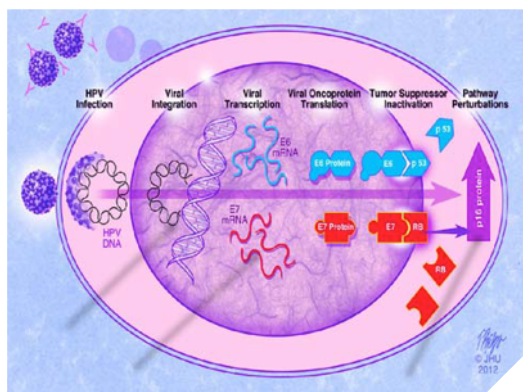


FIGURE 1. Schematic illustration of the Biology of HPV infection provides points of HPV Detection (Bishop *et al.*). Testing E6/E7 transcripts by RNA ISH is desirable because it indicates the presence of transcriptionally active virus and enables visualization of viral transcripts directly in tumor cells in tissue sections, unlike RT-PCR (Upko *et al.*, 2011).

Highest Sensitivity and Specificity

Current methodologies for HPV testing include PCR-based amplification and DNA *in situ* Hybridization (ISH). PCR amplification of HPV DNA is more sensitive, but it is less specific than DNA ISH. Published studies (Bishop *et al.*, 2013, Upko *et al.*, 2011, and Schache *et al.*, 2013) indicate that RNAscope-based ISH assay is more sensitive than DNA ISH in detecting HPV in Oropharyngeal Squamous Cell Carcinoma (OSCC) and results correlate well with p16 immunohistochemistry (IHC) staining (Table 1).

RNAscope ISH technology is an ideal platform giving the high sensitivity and specificity needed for detection of HPV biomarkers in Head and Neck Squamous Cell Carcinoma (HNSCC) samples.

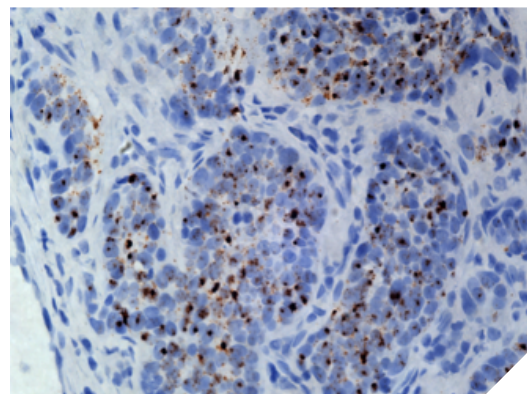


FIGURE 2. High risk HPV E6/E7 mRNA expression in Head and Neck Squamous Cell Carcinoma (HNSCC). FFPE section of HNSCC hybridized with a pool of HPV 16, 18, 31, 33, 35, 52 and 58 genotype probes, showing cytoplasmic punctate dots only in the tumor cells (40X magnification).

Assay Methods	Sensitivity	Specificity	PPV	NPV
P16 IHC	97%	82%	80%	97%
HR-HPV DNA ISH	94%	91%	89%	95%
Combined p16/HR-HPV DNA ISH	94%	91%	89%	95%
DNA qPCR	91%	87%	83%	93%
Combined p16/DNA qPCR	91%	93%	91%	93%
RNAscope HR-HPV	97%	93%	91%	98%

TABLE 1. Using the current “gold-standard” qPCR as the reference method, RNAscope® Technology demonstrated the best sensitivity and specificity for HPV status determination than existing methods (Schache *et al.*, 2013). Abbreviations: IHC=immunohistochemistry; ISH=in situ hybridization; NPV=negative predictive value; PPV=positive predictive value; qPCR=quantitative PCR

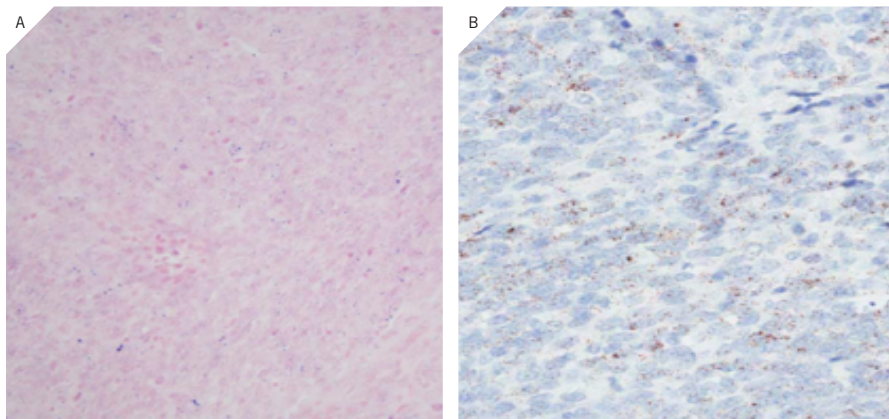


FIGURE 3. High Concordance with DNA and RNA *E6/E7* ISH for HR HPV (Upko *et al.*, 2011). Panel A shows positive DNA ISH with the Ventana III assay with punctate, blue, nuclear staining (magnification, 400X). Panel B shows positive RNA ISH with the same with diffuse, finely granular, cytoplasmic staining (magnification, 400X).

High Concordance between p16-IHC and RNAscope ISH

Case Study: Upko et al., 2011

Here the authors studied 211 oropharyngeal squamous cell carcinoma using tissue microarrays (TMAs) and observed high concordance between RNA ISH and p16 IHC, as well as superior sensitivity of RNA ISH to HPV DNA ISH (Figure 3).

Overall RNA ISH showed a consistently strong, diffuse granular (and or punctate) staining pattern in the tumor cells. The results are more readily visualized than those from DNA ISH

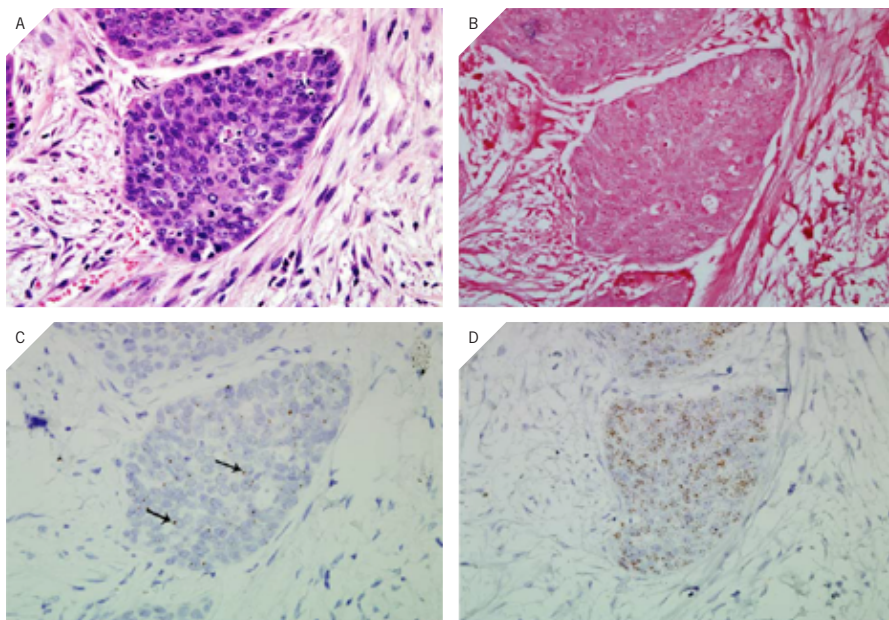


FIGURE 4. HPV-related oropharyngeal squamous cell carcinoma detection (Bishop *et al.*, 2013). Panel A is stained with hematoxylin & eosin stain. Panel B indicates that HPV was not detected using the Ventana Inform HPV III Family 16 probe. Panel C shows the presence of HPV at low viral copy numbers using the type 16-specific probe (arrows point to small hybridization signals within nuclei of tumor cells). Panel D clearly shows mRNA transcripts as numerous granular signals.

High Sensitivity of RNAscope ISH Independent of Viral Load

Case Study: Bishop et al., 2013

In this study, the authors showed that the RNAscope® ISH Assay was successful in the detection of HPV biomarkers—even in samples with very low viral load.

HPV-related Oropharyngeal squamous cell carcinoma HPV was not detected using Ventana DNA ISH, but low copy numbers were observed by using HPV 16-specific probe with the Dako GenPoint™ assay (Figure 4, Panel B and C).

Using the RNAscope assay, *E6/E7* mRNA transcripts are clearly detected as numerous granular signals (Figure 4, Panel D). Here, the authors confirmed that most oropharyngeal squamous cell carcinomas harbor transcriptionally active HPV.

HPV Quantitative Molecular Detection Coupled with Morphological Context

Broadest Spectrum HPV Quantitative Molecular Detection

HPV tumor status is the most powerful prognostic indicator for patients with head and neck cancer. Research has indicated that with the same treatment HPV positive cancer cases had a better overall survival: 80 to 85% versus approximately 35 to 38% for HPV negative cancer cases (CAP Today article, 2014). Clinical assessment of HNSCC disease basis requires accurate and comprehensive detection of active HPV subtypes. While > 90% of HPV cases are type 16, an assay assessing all high risk (HR) subtypes is recommended for labs assessing HPV status in HNSCC.

RNAscope assays offer a broad spectrum of probes targeting 18 High Risk subtypes as well as 6 Low Risk subtypes. The HR 18 HPV types were selected based on thought leader interviews and are in accordance with published literature and the International Agency for Research on Cancer (IARC). Users are able to easily customize probe pools to evaluate specific HPV genotypes.

Routinely used with FFPE and Fresh Frozen Tissues

Extensive assay development was done to ensure that the RNAscope assay will work with a wide range of tissue types—both FFPE and fresh frozen are routinely used.

Tissue Morphological Context

In many labs today, oropharyngeal cancer cases are routinely tested using a surrogate marker, p16-IHC, which may be followed by an extraction assay using RT-PCR to determine HPV involvement. A major concern with an extract assay of HPV is the loss of morphological context, which is essential to establish that active HPV is present in tumor cells. An advantage of RNAscope ISH is the specific assessment of HPV is in the context of intact tissue morphology. The RNAscope ISH signal is identified as strong, clear punctate chromogenic dots present in the nucleus and/or cytoplasm. The recommended assay format includes the use of assay control probes, bacterial gene Dap B (negative control) and housekeeping gene UBC (positive control to verify sample RNA integrity) run on serial sections alongside HPV probe. For positive identification of HPV, the staining should have granular cytoplasmic and/or nuclear brown staining that is measurably higher than the negative Dap B control slide (Figure 5).

Robust and Easy Methodology

The RNAscope ISH workflow is similar to IHC and can be automated. Thousands of off-the-shelf catalog probes and several probe pools are already available. Custom probes and probe pools are available in less than 3 weeks.

REFERENCES

1. Bishop JA, Ma XJ, Wang H, Luo Y, Illei PB, Begum S, Taube JM, Koch WM, Westra WH (2012). Detection of Transcriptionally Active High-risk HPV in Patients With Head and Neck Squamous Cell Carcinoma as Visualized by a Novel E6/E7 mRNA *In Situ* Hybridization Method. *American Journal of Surgical Pathology*, 36 (12):1874–1882. doi: 10.1097/PAS.0b013e318265fb2b.
2. Schache AG, Liloglou T, Risk JM, Jones TM, Ma XJ, Wang H, Bui S, Luo Y, Sloan P, Shaw RJ, Robinson M (2013). Validation of a novel diagnostic standard in HPV-positive oropharyngeal squamous cell carcinoma. *British Journal of Cancer*, 108(6):1332–1339. doi: 10.1038/bjc.2013.63.
3. Ukpo OC, Flanagan JJ, Ma XJ, Luo Y, Thorstad WL, Lewis JS Jr (2011). High-Risk Human Papillomavirus E6/E7 mRNA Detection by a Novel *In Situ* Hybridization Assay Strongly Correlates With p16 Expression and Patient Outcomes in Oropharyngeal Squamous Cell Carcinoma. *American J of Surgical Pathology*, 35(9):1343–1350. doi: 10.1097/PAS.0b013e318220e59d.
4. CAP TODAY, Dec 2013 Newsletter.

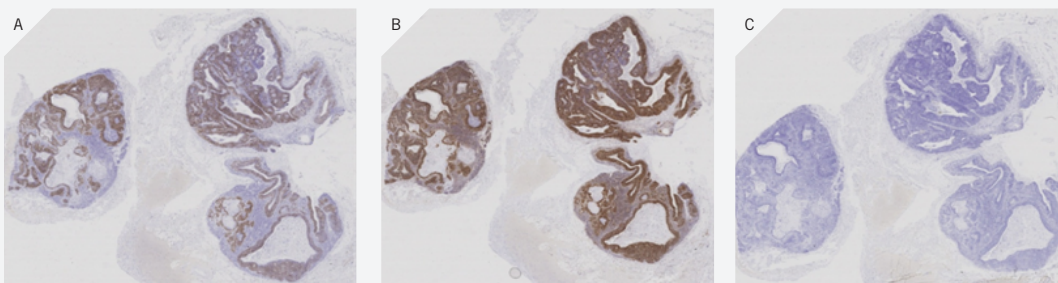


FIGURE 5. High expressing HR-HPV case study using RNAscope® assay with a positive and a negative control slide. Panel A. HPV-HR18. Panel B. UBC Positive Control. Panel C. DAP B Negative Control.

Ordering Information

RNAscope® assay is available in both manual and automated formats, to suit individual lab demand. The manual assay workflow is similar to IHC and only requires a Hybridization oven (HybEZ™ System) to help control the temperature and humidity during hybridization. The RNAscope automated assay is available on Ventana® Discovery® XT and Discovery Ultra systems.

RNAscope Reagent Kits

CAT #	Product Name	CAT #	Product Name
310035	RNAscope 2.0 HD Reagent Kit - BROWN (20 slides/reagent kit)	320600	RNAscope VS Reagent Kit - BROWN (60 slides/reagent kit)

HPV Catalog Pooled Probes

A broad spectrum of probe selection is available for HPV genotyping. Note each probe vial is sufficient for 20 slides.

CAT #	Probe Name	Description
312351	Probe - HPV HR7	HPV High Risk 7 Pool: 16,18, 31, 33, 35, 52 and 58
312591	Probe - HPV HR18	HPV High risk 18 Pool: 16,18, 26, 31, 33, 35, 39, 45, 51, 52, 53, 56, 58, 59, 66, 68, 73 and 82
407601	Probe - HPV LR6	HPV Low risk 6 Pool: 6, 11, 40, 42, 43 and 44
314551	Probe - HPV LR10	HPV Low risk 10 Pool: 6, 11, 40, 43, 44, 54, 70, 69, 71 and 74
311121	Probe - HPV16/18	HPV Pool: 16 and 18

HPV Catalog Pooled Probes

A broad spectrum of probe selection is available for HPV genotyping. Note each probe vial is sufficient for 20 slides

CAT #	Probe Name	CAT #	Product Name	CAT #	Probe Name
407501	Probe - HPV 6	407511	Probe - HPV 40	311651	Probe - HPV 59
316431	Probe - HPV10	407521	Probe - HPV 42	311661	Probe - HPV 66
311471	Probe - HPV 11	407531	Probe - HPV 43	404091	Probe - HPV 67
311521	Probe - HPV 16	311591	Probe - HPV 45	311671	Probe - HPV 68
311531	Probe - HPV 18	311601	Probe - HPV 51	407561	Probe - HPV 69
311541	Probe - HPV 26	311611	Probe - HPV 52	407571	Probe - HPV 70
311551	Probe - HPV 31	311621	Probe - HPV 53	407581	Probe - HPV 71
311561	Probe - HPV 33	407551	Probe - HPV 54	311681	Probe - HPV 73
311571	Probe - HPV 35	311631	Probe - HPV 56	407591	Probe - HPV 74
311581	Probe - HPV 39	311641	Probe - HPV 58	311691	Probe - HPV 82

ACD offers an ever-growing selection of RNA biomarker probes. Don't see your gene of interest? Custom probes designed within 2 weeks for virtually ANY gene from ANY species in ANY tissue.

Experience unprecedented molecular specificity and morphological data in one sensitive assay at acdbio.com/hpv

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