

# THE EXPRESSION OF E6 HPV, p53 AND p16INK4A AT WELL, MODERATELY, AND POORLY DIFFERENTIATED CERVICAL ADENOCARCINOMA

*by Gondo Mastutik*

---

**Submission date:** 28-Oct-2022 04:13PM (UTC+0800)

**Submission ID:** 1937677914

**File name:** ODERATELY,\_AND\_POORLY\_DIFFERENTIATED\_CERVICAL\_ADENOCARCINOMA.pdf (295.63K)

**Word count:** 3824

**Character count:** 19879

## THE EXPRESSION OF E6 HPV, p53 AND p16INK4A AT WELL, MODERATELY, AND POORLY DIFFERENTIATED CERVICAL ADENOCARCINOMA

Gondo Mastutik<sup>1</sup>, Alphania Rahniayu<sup>1,2</sup>, Nila Kurniasari<sup>1,2</sup>, Anny Setijo Rahaju<sup>1,2</sup>, Rahmi Alia<sup>3</sup>, Sjahjenny Mustokoweni<sup>1,2</sup>

<sup>1</sup>Department of Anatomic Pathology, Faculty of Medicine, Universitas Airlangga, Surabaya, <sup>2</sup>Dr. Soetomo General Academic Hospital, Surabaya, <sup>3</sup>Prof Dr Soekandar Hospital, Mojokerto, Indonesia

### ABSTRACT

25

The objective of this study is to analyze the expression of E6 Humanpapilloma virus (HPV), p53, and p16INK4A in cervical adenocarcinoma grade well differentiated (WD), moderately differentiated (MD), and poorly differentiated (PD). A cross sectional study conducted at Department of Anatomic Pathology, Dr. Soetomo General Academic Hospital Surabaya Indonesia using formalin fix paraffin embedded (FFPE) from cervical normal and cervical adenocarcinoma grade WD, MD, and PD. The expression of E6 HPV, p53, and p16INK4A was performed by immunohistochemistry (IHC) staining. Data were analyzed with Kruskal-Wallis and continued with Mann-Withney test. The expression of E6 HPV in the cervical adenocarcinoma showed 35.9% specimens represented negative and 64.1% specimens represented positive. There was no significant difference in the expression of E6 HPV and p53 in cervical adenocarcinoma between grade WD, MD, and PD. The p16INK4A was overexpressed, shown as diffuse appearance in 89.7% of the specimens. There was a significant difference in the expression of p16INK4A between grade WD and MD with PD. In conclusion, some of cervical adenocarcinoma were not caused by infection of HPV type 16 or 18 and the expression of p16INK4A might take a role in the developing of malignancy that caused by infection of HPV.

**Keywords:** Grading of cervical adenocarcinoma; E6 HPV; p53; p16INK4A

### ABSTRAK

Tujuan penelitian ini adalah menganalisa ekspresi E6 Humanpapilloma virus (HPV), p53, dan p16INK4A pada adenocarcinoma servik grade well differentiated (WD), moderately differentiated (MD), dan poorly differentiated (PD). Penelitian cross sectional dilaksanakan di Departemen Patologi Anatomi, Rumah Sakit Umum Pendidikan Dr. Soetomo Surabaya Indonesia menggunakan blok paraffin (BP) dari jaringan serviks normal dan adenocarcinoma servik grade WD, MD, dan PD. Evaluasi ekspresi E6 HPV, p53, dan p16INK4A dilakukan dengan pewarnaan imunohistochemistry (IHC). Data dianalisa dengan Kruskal-Wallis kemudian dilanjutkan dengan Mann-Withney test. Ekspresi E6 HPV pada adenocarcinoma servik menunjukkan 35.9% specimen adalah negatif dan 64.1% specimen adalah positif. Tidak terdapat perbedaan yang nyata ekspresi of E6 HPV dan p53 pada adenocarcinoma servik antara grade WD, MD, dan PD. Terjadi overekspresi p16INK4A, yang menunjukkan sebaran merata pada 89.7% dari spesimen. Terdapat perbedaan yang nyata pada ekspresi p16INK4A antara grade WD dan MD dengan PD. Kesimpulan, beberapa specimen adenocarcinoma serviks tidak disebabkan oleh infeksi HPV 16 atau HPV 18 dan ekspresi p16INK4A mungkin berperan dalam perkembangan keganasan servik yang disebabkan oleh infeksi HPV.

**Kata kunci:** Grade adenocarcinoma servik; E6 HPV; p53; p16INK4A

**Correspondence:** Gondo Mastutik, Department of Anatomic Pathology, Faculty of Medicine, Universitas Airlangga, Jl. Prof Dr Moestopo 47, Surabaya 60132, Indonesia, E-mail: gondomastutik@fk.unair.ac.id, gondomastutik@gmail.com

pISSN:2355-8393 • eISSN: 2599-056x • doi: <http://dx.doi.org/10.20473/fmi.v55i4.17327>  
 • Fol Med Indones. 2019;55:295-300 • Received 24 May 2019 • Accepted 28 Nov 2019  
 • Open access under CC-BY-NC-SA license • Available at <https://e-journal.unair.ac.id/FMI/>

### INTRODUCTION

Cervical cancer is the fourth most common cancer diagnosed in women worldwide. In Indonesia, it is the second most common cancer diagnosed in women after breast cancer, with an estimated 20,928 new cases and 4,097 deaths in 2012 (Ferlay et al 2013). Based on the histological classification of tumours, the most common

type of cervical cancer is squamous cell carcinoma (Wells et al 2003) and cervical adenocarcinoma comprises 15% of cervical cancer cases (Siriaunkgul et al 2013). The incidence rates of cervical adenocarcinoma tend to increase in Korean women (Oh et al 2013), in women in the United States (Adegoke et al 2012), and in the Netherlands where it is found predominantly in women of 25-39 years old (van der

Horst et al 2017). This trend is particularly evident among females aged <40 years and has occurred despite extensive cytology-based screening programs (Tornesello et al 2014). Adenocarcinoma originates from glandular precursor lesions of the endocervical mucosa that are difficult to reach by pap smear examination. Therefore, although screening programs have led to a substantial decrease in the incidence of squamous cell carcinoma (Andersson et al 2013), it is not protective for precursor cancer whose cells cannot be reached by screening programs, including cervical adenocarcinoma.

Humanpapilloma virus (HPV) has already been established as the cause of cervical cancer and it is related to other anogenital cancers (anus, vulva, vagina and penis) as well as head and neck cancers. HPV types 16 and 18 are responsible for about 70% of all cervical cancer cases worldwide (Bruni et al 2017). The E6 and E7 of the high risk HPV have associated with oncogenic transformation. E6 interacts with the product of tumour suppressor genes p53. It targets p53 for degradation via ubiquitin pathway, resulting in loss of G2/M checkpoint regulation. In addition, degradation of p53 by E6 HPV leads to irreversible DNA damage and the cells enter the cell cycle and cease to apoptose, resulting in the accumulation of genomic instability and genetic alteration. E7 interacts with retinoblastoma (RB) family member RB1, RBL1, and RBL2, then targets them for degradation, resulting in nuclear translocation of E2F and promotion of S-phase transition. The result of pRb downregulation of the loss of feedback inhibition and overexpression of p16INK4A (Scheffner et al 1990, Hietanen et al 2000, Crosbie et al 2013).

During the progression of cervical adenocarcinoma, cells differentiate into well-differentiated (WD), moderately-differentiated (MD), or poorly-differentiated (PD). However, the expression of p53 and p16INK4A in cervical adenocarcinoma grade WD, MD, PD is still not clear. The objective of this study was to analyze the expression of E6, p53, and p16INK4A in cervical adenocarcinoma grade WD, MD, and PD.

## 29 MATERIALS AND METHODS

### Specimens

This was a cross sectional study conducted at the Department of Anatomic Pathology, Dr. Soetomo General Academic Hospital Surabaya Indonesia. The specimens were cervixes from the uterine prolapse cases and adenocarcinoma cases that had been diagnosed by pathologist. There were 10 formalin fix paraffin embedded (FFPE) of the uterine prolapse as control group and 39 FFPE of cervical adenocarcinoma that

consisted of 14 tissues were grade WD, 12 tissues were grade MD, and 13 tissues were grade PD.

### Immunohistochemistry staining

The expressions of E6 HPV, p53, and p16INK4A are detected by immunohistochemistry (IHC) staining using the HPV16 E6/18 E6 Antibody (C1P5): sc-460 (Santa Cruz Biotechnology), monoclonal antibody p53 clone Y5 (Biocare), and Anti-CDKN2A/ p16INK4A Antibody (clone 1E12E10) IHC-plus™ LS-B5261 (LS Bio).

The scoring of the positive interpretation for E6 HPV is performed when the nucleus and cytoplasm were stained clearly and scored using an arbitrary semiquantitative scale. Score 0 is given when there are no cells stained representing negative staining, score 1 when 5-25% of the cells stained representing mild positive staining, score 2 when 25-50% of the cells stained representing moderate positive staining, and score 3 when >50% of the cells stained representing extensive immunostaining (Chaudhary et al 2013).

Expression of p53 is evaluated when the nucleus is stained clearly and scored semiquantitatively. The score is 0 if up to 5% of the cells are positive, score 1 if 5-25% of the cells are positive, score 2 if 26-50% of the cells are positive, score 3 if 51-75%, and score 4 if more than 75% of the cells are positive (Baalbergen et al 2013).

16 Expression of p16INK4A is evaluated as positive when nuclear or cytoplasmic immunostaining is clearly demonstrated. Scoring of p16INK4A is performed using the criteria of Klaes et al. (2001). In this scoring, score 0 is given if there is no staining or <1% of the cells are positively representing negative, score 1 if 1-5% of the cells positively representing sporadic, score 2 if 5-25% of the cells are positively representing focal, and score 3 if >25% of the cells are positively representing diffuse (Kazlouskaya et al 2013).

### Statistical analysis

To evaluate different expression of E6 HPV, p53 and p16INK4A at normal cervix tissue and cervical adenocarcinoma tissue based on differentiated grading, the analysis was performed by using The Kruskal-Wallis ( $p < 0.005$ ) and continued with Mann Withney test ( $p < 0.005$ ).

## RESULTS

This study was performed to cervix specimen of 49 women with ages 30-81 years old and average 51.78 ±

9.395 (mean + SD). There were 20.4% FFPE from cervical normal and 79.6% from cervical adenocarcinoma that consisted of 35.9%, 30.8%, and 33.3% FFPE from WD, MD, and PD tissues, respectively.

The expression of E6 HPV, p53, and p16INK4A in cervical adenocarcinoma grade WD, MD, and PD showed in IHC staining (Figure 1). The expression of E6 HPV in the cervical adenocarcinoma showed that those with score 0 were 14 (35.9%) specimens, score 1 were 11 (28.2%), score 2 were 11 (28.2%), and score 3 were 3 (7.7%). The expression of E6 HPV in cervical adenocarcinoma in grade WD, MD, and PD have the same "a" subscript (Table 1). It means that there was no significant difference in the expression of E6 HPV between cervical adenocarcinoma in grade WD, MD, and PD.

Expression of p53 in the cells of cervical adenocarcinoma WD, MD, and PD showed that those with score 0 were 12 (30.8%), score 1 were 15 (38.5%), score 2 were 3 (7.7%), score 3 were 4 (10.2%), and score 4 were 5 (12.8%). The result showed that there was no significant difference in the expression of p53 between cervical adenocarcinoma grade WD, MD, and PD ( $p > 0.05$ ) (Table 1). Expression of p16INK4A in cervical adenocarcinoma grade WD, MD, and PD showed that almost all of the specimens were positively diffuse (score 3) and there was no negative (Fig. 1). The result showed

that there was a significant difference in the expression of p16INK4A between cervical adenocarcinoma in grade WD and MD with PD (Table 1).

## DISCUSSION

Chronic infection of HPV is correlated with the development of cervical cancer (Wells et al 2003). HPV 16, 18 are the most common genotypes of HPV that are presenting in cervical adenocarcinoma (An et al 2005, Tornesello et al 2011, Siriaunkgul et al 2013). In Korean women, the prevalence of HPV infection in cervical adenocarcinoma was 90%. The infection of HPV 16 and/or HPV 18 accounted for 78% of HPV-positive adenocarcinomas. Multiple HPV types were found in 13% of the cases. Mostly HPV 16 and HPV 18 are highly associated with most of cervical adenocarcinomas (An et al 2005). In Thailand, predominant genotype is HPV 18, being twice as common as HPV 16 (Siriaunkgul et al 2013). In Uppsala Sweden, 95% were HR HPV positive; HPV18/45 predominated (77%), followed by HPV16 (27%) (Andersson et al 2013). In the Netherlands, HPV1 8 is mainly the risk factor for the development of adenocarcinoma, while HPV16 is associated with both SCC and adenocarcinoma (Bulk et al 2006). Because of that in this study we used the E6 of HPV 16 and HPV 18.

**Table 1.** Different expression of E6 HPV, p53, and p16INK4A at cervical normal and well differentiated, moderately differentiated, and poorly differentiated adenocarcinoma

E6	AWD <sup>a</sup>	AMD <sup>a</sup>	APD <sup>a</sup>	CN <sup>b</sup>	Total	p=0.001
0	4 (28.6%)	5 (41.7%)	5 (38.5%)	10 (100%)	24 (49%)	
1	6 (42.9%)	2 (16.7%)	3 (23.1%)	0 (0%)	11 (22.4%)	
2	2 (14.3%)	5 (41.7%)	4 (30.8%)	0 (0%)	11 (22.4%)	
3	2 (14.3%)	0 (0%)	1 (7.7%)	0 (0%)	3 (6.1%)	
p53	AWD	AMD	APD	CN		p=0.179
0	4 (28.6%)	3 (25%)	5 (38.5%)	6 (60%)	18 (36.7%)	
1	6 (42.9%)	4 (33.3%)	5 (38.5%)	3 (30%)	18 (36.7%)	
2	1 (7.1%)	1 (8.3%)	1 (7.7%)	1 (10%)	4 (8.2%)	
3	1 (7.1%)	1 (8.3%)	2 (15.4%)	0 (0%)	4 (8.2%)	
4	2 (14.3%)	3 (25%)	0 (0%)	0 (0%)	5 (10.2%)	
p16 <sup>INK4A</sup>	AWD <sup>a</sup>	AMD <sup>a</sup>	APD <sup>b</sup>	CN <sup>c</sup>		p= <0.0001
0	0 (0%)	0 (0%)	0 (0%)	9 (90%)	9 (18.4%)	
1	0 (0%)	0 (0%)	1 (7.7%)	1 (10%)	2 (4.1%)	
2	0 (0%)	0 (0%)	3 (23.1%)	0 (0%)	3 (6.1%)	
3	14 (100%)	12 (100%)	9 (69.2%)	0 (0%)	35 (71.4%)	

Note: Different superscript shows significant differences. AWD=Well differentiated adenocarcinoma, AMD=moderately differentiated adenocarcinoma, APD= poorly differentiated adenocarcinoma, CN=cervical normal

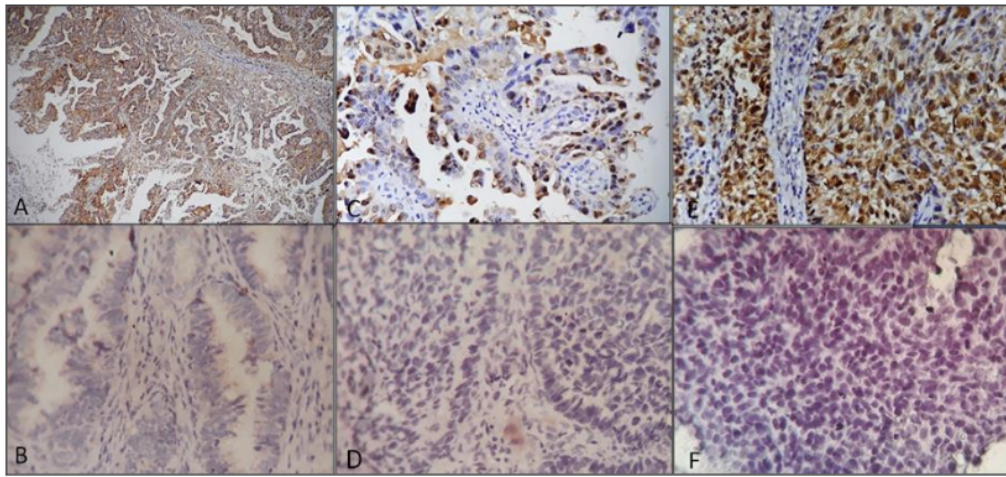


Fig. 1. Expression of E6 HPV, p53, and p16INK4A in cervical adenocarcinoma grade well differentiated (WD), moderately differentiated (MD), poorly differentiated (PD). A. Expression E6 HPV, positive in adenocarcinoma WD, 100X; B. Expression E6 HPV, negative in adenocarcinoma WD, 400X; C. Expression p53, positive in adenocarcinoma MD, 400X; D. Expression p53, negative in adenocarcinoma MD, 400X; E. Expression p16INK4A, positive in adenocarcinoma PD, 400X; F. Expression p16INK4A, negative in adenocarcinoma PD, 400X.

The histology of cervical adenocarcinoma exhibits a variety of architectural patterns and ranges from WD, MD, and PD. The WD of cervical adenocarcinoma may show exophytic papillary growth or infiltrative growth, or both. The invasive tumor may be composed of irregular cystic and tubular glands, glands with intraluminal papillary infoldings, or cribriform glands. In the MD tumors the growth is more confluent with sheets of small cribriform glands. The poorly differentiated adenocarcinomas show solid areas of undifferentiated cells that may be undistinguishable from poorly differentiated squamous cell carcinoma (Pirog, 2017). This grading change occurs along with the development of malignancy so that it will affect the expression of tumor suppressor gene, p53 and p16INK4A.

E6 HPV in this study represented the infection of HPV 16 and or 18 (HPV 16/18), showed that 35.9% and 64.1% specimens of cervical adenocarcinoma represented negative and positive for E6 HPV 16/18, respectively. This suggested that some of cervical adenocarcinoma tissues were not infected by HPV type 16 or 18, so it might be related with other HPV types or any others causes. This study was in accordance with other study that the prevalence of HPV in cervical adenocarcinoma varied from 62 to 100%, depending on geographic region and tumor subtype (Baalbergen et al 2013). The expression of E6 HPV in cervical

adenocarcinoma between grade WD, MD, and PD, there are no significant different. This data suggested that the infection of HPV type 16 or 18 did not affect the grading differentiation of cervical adenocarcinoma.

This study showed that the score 0 for the expression of p53 in the cells of cervical adenocarcinoma grade WD, MD, and PD was 30.8%, and score 1, 2, 3, and 4 were 69.2%. It suggests that some specimens have not and have shown the expression of p53. The wild type p53 has half-life of about 5-20 minutes (Dowell et al 1994) and then degraded, whereas the E6 HPV 16 and 18 binds to p53 and also stimulate degradation (Crosbie et al 2013), so it would be negative for IHC staining. The ubiquitin form of p53 as result of degradation by E6 HPV and mutant type p53 should be positive by IHC staining. Other studies showed that the mutation of p53 gene in cervical adenocarcinoma is more frequent than that in squamous cell carcinoma or cervical intraepithelial neoplasia. There is a single nucleotide substitutions in p53 gene detected in 36% cervical adenocarcinoma cases and this mutation is independent from HPV infection status (Tornesello et al 2014). In cervical adenocarcinoma, the positivity of p53 is associated with poor survival. The survival rate in cervical adenocarcinoma is 67%, and it is not influenced by estrogen receptor, progesterone receptor, MIB-1, or bcl-2 with strongly positive staining, but significantly influenced by the expression of p53. The p53 staining is

strongly positive. The survival is significantly worse than that in tumors scored as negative or weak positive (Baalbergen et al 2013).

This study found that the expression of p16INK4A was overexpressed in cervical adenocarcinoma grade WD, MD, and PD. This was demonstrated by the diffuse appearance in 89.7% (35/39) of the specimens. It was also showed that a significant difference in the expression of p16INK4A in cervical adenocarcinoma between grade WD and MD with PD. The expression of p16INK4A in WD and MD was diffusely positive. This result was the same with the finding in other study that p16INK4A was expressed in diffusely or strongly expression in cervical intraepithelial neoplasia I, II, III, squamous cell carcinoma, endocervical glandular dysplasia, adenocarcinoma in situ, and invasive adenocarcinoma (Tringler B et al 2004); as well as in endocervical carcinoma (Liang et al 2016). The finding in this study was in accordance with the results of other studies. The overexpression of p16INK4A in uterine cervix adenocarcinoma is common and significantly associated with HPV infection (mainly high-risk HPV types). It suggests that p16INK4A is a putative molecular biomarker that consistently discriminates uterine cervix adenocarcinomas from benign lesions and from endometrioid adenocarcinomas of the uterine corpus (Missaoui et al 2006). The overexpression of the p16INK4A was typical for dysplastic and neoplastic epithelia of the uterine cervix. The expression of p16 increases from normal to invasive squamous carcinoma in the uterine cervix emphasizing that it might be a useful marker for predicting risk of developing cervical cancer in women (Izadi-Mood et al 2012).

## 21 CONCLUSION

There was no significant difference in the expression of E6 HPV and p53 in cervical adenocarcinoma between grade WD, MD, and PD, but there was a significant difference in the expression of p16INK4A between grade WD and MD with PD. This finding suggest that the expression of p16INK4A might take a role in the developing of malignancy that caused by infection HPV.

## 6 ACKNOWLEDGMENT

This study was supported by the Ministry of Research, Technology, and Higher Education of the Republic of Indonesia by the university operational funding fiscal years at 2017. Thank you for the Indonesian Government and the Faculty of Medicine Airlangga University.

## REFERENCES

- Adegoke O, Kulasingam S, Virnig B (2012). Cervical cancer trends in the United States: a 35-year population-based analysis. *J Womens Health (Larchmt)* 21, 1031-7
- An HJ, Kim KR, Kim IS, et al (2005). Prevalence of human papillomavirus DNA in various histological subtypes of cervical adenocarcinoma: a population-based study. *Mod Pathol* 18, 528-34
- Andersson S, Mints M, Wilander E (2013). Results of cytology and high-risk human papillomavirus testing in females with cervical adenocarcinoma in situ. *Oncol Lett* 6, 215-9
- Baalbergen A, Smedts F, Ewing P, et al (2013). HPV-type has no impact on survival of patients with adenocarcinoma of the uterine cervix. *Gynecol Oncol* 128, 530-4
- Bruni L, Barrionuevo-Rosas L, Albero G, et al 2017. Human Papillomavirus and Related Diseases in Indonesia. Summary Report 27 July 2017 [Online]. ICO/IARC Information Centre on HPV and Cancer (HPV Information Centre) 2017. Accessed April 12, 2018
- Bulk S, Berkhof J, Bulkman NW, et al (2006). Preferential risk of HPV16 for squamous cell carcinoma and of HPV18 for adenocarcinoma of the cervix compared to women with normal cytology in The Netherlands. *Br J Cancer* 94, 171-5
- Chaudhary A, Pandya S, Singh M, et al (2013). Identification of high-risk human papillomavirus-16 and -18 infections by multiplex PCR and their expression in oral submucous fibrosis and oral squamous cell carcinoma. *Head and Neck Oncology* 5, 1-10
- Crosbie EJ, Einstein MH, Franceschi S, et al (2013). Human papillomavirus and cervical cancer. *The Lancet* 382, 889-99
- Dowell SP, Wilson PO, Derias NW, et al (1994). Clinical utility of the immunocytochemical detection of p53 protein in cytological specimens. *Cancer Res* 54, 2914-8
- Hietanen S, Lain S, Krausz E, et al (2000). Activation of p53 in cervical carcinoma cells by small molecules. *Proc Natl Acad Sci U S A*, 97, 8501-6
- Izadi-Mood N, Asadi K, Shojaei H, et al (2012). Potential diagnostic value of P16 expression in premalignant and malignant cervical lesions. *J Res Med Sci* 17, 428-33
- Kazlouskaya V, Shustef E, Allam SH, et al (2013). Expression of p16 protein in lesional and perilesional condyloma acuminata and bowenoid papulosis: clinical significance and diagnostic implications. *J Am Acad Dermatol* 69, 444-9
- Klaes R, Friedrich T, Spitkovsky D, Ridder R, Rudy W, Petry U, et al. (2001). Overexpression of p16INK4A

- as a specific marker for dysplastic and neoplastic epithelial cells of the cervix uteri *Int J Cancer* 2001 92, 276-84
- Liang L, Zheng W, Liu J, et al (2016). Assessment of the Utility of PAX8 Immunohistochemical Stain in Diagnosing Endocervical Glandular Lesions. *Arch Pathol Lab Med*, 140, 148-52
- Missaoui N, Hmissa S, Frappart L, et al (2006). p16INK4A overexpression and HPV infection in uterine cervix adenocarcinoma. *Virchows Arch* 448, 597-603
- Oh CM, Jung KW, Won YJ, et al (2013). Trends in the incidence of in situ and invasive cervical cancer by age group and histological type in Korea from 1993 to 2009. *PLoS One* 8, e72012
- Pirog EC (2017). Cervical Adenocarcinoma: Diagnosis of Human Papillomavirus-Positive and Human Papillomavirus-Negative Tumors. *Arch Pathol Lab Med* 141, 1653-67
- Scheffner M, Werness BA, Huibregtse JM, et al (1990). The E6 oncoprotein encoded by human papillomavirus types 16 and 18 promotes the degradation of p53. *Cell* 63, 1129-36
- Siriaunkgul S, Utaipat U, Suthipintawong C, et al (2013). HPV genotyping in adenocarcinoma of the uterine cervix in Thailand. *Int J Gynaecol Obstet* 123, 226-30
- Tornesello ML, Annunziata C, Buonaguro L, et al (2014). TP53 and PIK3CA gene mutations in adenocarcinoma, squamous cell carcinoma and high-grade intraepithelial neoplasia of the cervix. *J Transl Med*, 12, 255
- Tornesello ML, Losito S, Benincasa G, et al (2011). Human papillomavirus (HPV) genotypes and HPV16 variants and risk of adenocarcinoma and squamous cell carcinoma of the cervix. *Gynecol Oncol*, 121, 32-42
- van der Horst J, Siebers AG, Bulten J, et al (2017). Increasing incidence of invasive and in situ cervical adenocarcinoma in the Netherlands during 2004-2013. *Cancer Med*, 6, 416-23
- Wells M, Östör AG, Crum CP, et al (2003). Tumours of the uterine cervix, epithelial tumours. In 'Pathology and Genetics of Tumours of the Breast and Female Genital Organs', Eds IARC Press, Lyon, 259-72

# THE EXPRESSION OF E6 HPV, p53 AND p16INK4A AT WELL, MODERATELY, AND POORLY DIFFERENTIATED CERVICAL ADENOCARCINOMA

## ORIGINALITY REPORT

17%

SIMILARITY INDEX

11%

INTERNET SOURCES

17%

PUBLICATIONS

1%

STUDENT PAPERS

## PRIMARY SOURCES

- 1 Ashesh B. Jani. "Prostate Cancer Modality Time Trend Analyses From 1973 to 2004 : A Surveillance, Epidemiology, and End Results Registry Analysis", American Journal of Clinical Oncology, 09/2009  
Publication 1%
- 2 [www.semanticscholar.org](http://www.semanticscholar.org)  
Internet Source 1%
- 3 ALBERTO RAMÍREZ-TORRES, JEOVANIS GIL, SANDRA CONTRERAS, GRACIELA RAMÍREZ et al. "Quantitative Proteomic Analysis of Cervical Cancer Tissues Identifies Proteins Associated With Cancer Progression", Cancer Genomics - Proteomics, 2022  
Publication 1%
- 4 Submitted to Universitas Airlangga  
Student Paper 1%
- 5 Yong Cui, Qi Wang, JunYu Wang, Yan Dong, Chun Luo, Guohan Hu, Yicheng Lu. 1%



"Knockdown of AKT2 expression by RNA interference inhibits proliferation, enhances apoptosis, and increases chemosensitivity to the anticancer drug VM-26 in U87 glioma cells", Brain Research, 2012

Publication

6

[mail.scialert.net](mailto:scialert.net)

Internet Source

1 %

7

[bmccancer.biomedcentral.com](http://bmccancer.biomedcentral.com)

Internet Source

1 %

8

"HIV/AIDS-Associated Viral Oncogenesis", Springer Science and Business Media LLC, 2019

Publication

1 %

9

"XXVII International Congress of the International Academy of Pathology October 12-17, 2008, Athens, Greece □ POSTER PRESENTATIONS", Histopathology, 10/2008

Publication

1 %

10

[www.pubmedcentral.nih.gov](http://www.pubmedcentral.nih.gov)

Internet Source

1 %

11

Chung, Tony K.H., Paul Van Hummelen, Paul K.S. Chan, Tak Hong Cheung, So Fan Yim, M. Y. Yu, Matthew D. Ducar, Aaron R. Thorner, Laura E. MacConaill, Graeme Doran, Chandra Sekhar Pedomallu, Akinyemi I. Ojesina, Raymond R.Y. Wong, Vivian W. Wang, Samuel

1 %

S. Freeman, Tat San Lau, Joseph Kwong, Loucia K.Y. Chan, Menachem Fromer, Taymaa May, Michael J. Worley, Katharine M. Esselen, Kevin M. Elias, Michael Lawrence, Gad Getz, David I. Smith, Christopher P. Crum, Matthew Meyerson, Ross S. Berkowitz, and Yick Fu Wong. "Genomic aberrations in cervical adenocarcinomas in Hong Kong Chinese women : Cervical adenocarcinoma genomics", International Journal of Cancer, 2015.

Publication

---

12 [archive.org](https://www.archive.org)  
Internet Source

1 %

---

13 Mohamed A. Youssef, Lobna Abdelsalam, Reem Abdelhameed Harfoush, Iman Mamdouh Talaat et al. "Prevalence of human papilloma virus (HPV) and its genotypes in cervical specimens of Egyptian women by linear array HPV genotyping test", Infectious Agents and Cancer, 2016

Publication

---

14 Sumalee Siriaungkul, Utaiwan Utaipat, Cheepsumon Suthipintawong, Kobkul Tungsinmunkong et al. "HPV genotyping in adenocarcinoma of the uterine cervix in Thailand", International Journal of Gynecology & Obstetrics, 2013

Publication

---

<1 %

<1 %

- |    |   |      |
|----|---|------|
| 15 | <a href="http://ir.ymlib.yonsei.ac.kr">ir.ymlib.yonsei.ac.kr</a><br>Internet Source   | <1 % |
| 16 | Filho, Adhemar. "Characterization of p16 and E6 HPV-related proteins in uterine cervix high-grade lesions of patients treated by conization with large loop excision", <i>Oncology Letters</i> , 2013.<br>Publication   | <1 % |
| 17 | Makiko Omori, Tetsuo Kondo, Hikaru Tagaya, Yumika Watanabe et al. "Utility of imaging modalities for predicting carcinogenesis in lobular endocervical glandular hyperplasia", <i>PLOS ONE</i> , 2019<br>Publication  | <1 % |
| 18 | O. M. Omran, M. AlSheeha. "Human Papilloma Virus Early Proteins E6 (HPV16/18-E6) and the Cell Cycle Marker P16 (INK4a) are Useful Prognostic Markers in Uterine Cervical Carcinomas in Qassim Region- Saudi Arabia", <i>Pathology &amp; Oncology Research</i> , 2014<br>Publication | <1 % |
| 19 | <a href="http://amsdottorato.unibo.it">amsdottorato.unibo.it</a><br>Internet Source   | <1 % |
| 20 | Jin W. Kim, Youl H. Cho, Chun G. Lee, Jae H. Kim, Heung K. Kim, Eun J. Kim, Ku T. Han, Sung E. Namkoong. "Human Papillomavirus  | <1 % |

# Infection and TP53 Gene Mutation in Primary Cervical Carcinoma", Acta Oncologica, 2009

Publication

21

S. H. Kim, J. M. Lee, H. G. Yun, U. S. Park, S. U. Hwang, J.-S. Pyo, J. H. Sohn. "Overall accuracy of cervical cytology and clinicopathological significance of LSIL cells in ASC-H cytology", *Cytopathology*, 2017

Publication

<1 %

22

Sakae Hata. "Diagnostic significance of endocervical glandular cells with "golden-yellow" mucin on pap smear", *Diagnostic Cytopathology*, 08/2002

Publication

<1 %

23

Yogi Putra Adhi Pradana, Budi Sutikno. "Olfactory neuroblastoma resection through endoscopic endonasal approach: A rare case report", *Annals of Medicine and Surgery*, 2022

Publication

<1 %

24

[academic.oup.com](http://academic.oup.com)

Internet Source

<1 %

25

[scholarlypages.org](http://scholarlypages.org)

Internet Source

<1 %

26

[www.lsbio.com](http://www.lsbio.com)

Internet Source

<1 %

27

[www.termedia.pl](http://www.termedia.pl)

Internet Source

<1 %

28

W J G Melchers. "Coexisting high-grade glandular and squamous cervical lesions and human papillomavirus infections", British Journal of Cancer, 09/01/2003

Publication

<1 %

29

medicopublication.com

Internet Source

<1 %

30

publications.ki.se

Internet Source

<1 %

31

"International Manual of Oncology Practice", Springer Science and Business Media LLC, 2015

Publication

<1 %

32

Ali Farhadi, Abbas Behzad-Behbahani, Bitra Geramizadeh, Zamberi Sekawi, Marjan Rahsaz, Sedigheh Sharifzadeh. "High-risk human papillomavirus infection in different histological subtypes of renal cell carcinoma", Journal of Medical Virology, 2014

Publication

<1 %

33

Daven N. Jackson-Humbles, John Curtis Seely, Ronald A. Herbert, David E. Malarkey et al. "Uterine Paramesonephric Cysts in Sprague-Dawley Rats from National Toxicology Program Studies", Toxicologic Pathology, 2018

Publication

<1 %

34

Galina Volgareva, Larisa Zavalishina, Yulia Andreeva, Georgy Frank et al. "Protein p16 as a marker of dysplastic and neoplastic alterations in cervical epithelial cells", BMC Cancer, 2004

Publication

---

<1 %

35

Maharani Laillyza Apriasari, Selviana Rizky Pramitha, Dewi Puspitasari, Diah Savitri Ernawati. "Anti-Inflammatory Effect of Musa acuminata Stem", European Journal of Dentistry, 2020

Publication

---

<1 %

36

Mutia Latief, Ika Agus Rini, Gita Widya Pradini, Gatot Nyarumenteng Adhipurnawan Winarno, Edhyana Sahiratmadja, Herman Susanto. "Phylogenetic Analysis of Human Papillomavirus 16 and 52 L1 Gene from Cervical Cancer in Bandung", The Indonesian Biomedical Journal, 2018

Publication

---

<1 %

37

Tornesello, M.L.. "Human papillomavirus (HPV) genotypes and HPV16 variants and risk of adenocarcinoma and squamous cell carcinoma of the cervix", Gynecologic Oncology, 201104

Publication

---

<1 %

38

Yi Guo. "Detection and identification of human papillomavirus in cervical

<1 %

adenocarcinoma", Chinese Journal of Cancer  
Research, 09/2008

Publication

39

[onlinelibrary.wiley.com](http://onlinelibrary.wiley.com)

Internet Source

<1 %

40

[publications.iarc.fr](http://publications.iarc.fr)

Internet Source

<1 %

41

[slidelegend.com](http://slidelegend.com)

Internet Source

<1 %

42

C J O'Neill. "High-grade ovarian serous carcinoma exhibits significantly higher p16 expression than low-grade serous carcinoma and serous borderline tumour",  
*Histopathology*, 5/2007

Publication

<1 %

43

Capra, G.. "HPV genotype prevalence in cytologically abnormal cervical samples from women living in south Italy", *Virus Research*, 200805

Publication

<1 %

44

S.-H. SONG. "The expression of p16 (INK4a) and Ki-67 in relation to high-risk human papilloma viral load and residual disease after conization with positive margins",  
*International Journal of Gynecological Cancer*, 7/2007

Publication

<1 %

45

Bulten, J.. "p14<sup>ARF</sup> and p16<sup>INK4A</sup>, two products of the same gene, are differently expressed in cervical intraepithelial neoplasia", Gynecologic Oncology, 200606

Publication

<1 %

46

Nemes, J.A.. "Expression of p16<sup>INK4A</sup>, p53, and Rb proteins are independent from the presence of human papillomavirus genes in oral squamous cell carcinoma", Oral Surgery, Oral Medicine, Oral Pathology, Oral Radiology and Endodontology, 200609

Publication

<1 %

Exclude quotes On

Exclude matches Off

Exclude bibliography On



# THE EXPRESSION OF E6 HPV, p53 AND p16INK4A AT WELL, MODERATELY, AND POORLY DIFFERENTIATED CERVICAL ADENOCARCINOMA

---

GRADEMARK REPORT

---

FINAL GRADE

**/100**

GENERAL COMMENTS

**Instructor**

---

PAGE 1

---

PAGE 2

---

PAGE 3

---

PAGE 4

---

PAGE 5

---

PAGE 6

---