

Possible deferral of diagnostic and therapeutic procedures for patients with abnormal screening tests results in cervical cancer secondary prevention in current SARS-CoV-2 pandemic Interim guidelines of the Polish Society of Gynecologists and Obstetricians and the Polish Society of Colposcopy and Cervical Pathophysiology

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ABSTRACT

The Polish Society of Gynecologists and Obstetricians and Polish Society of Colposcopy and Cervical Pathophysiology Interim Guidelines goal at aiding gynecologists in providing a cervical cancer prevention care during the evolving SARS-CoV-2 pandemic. Presented guidelines were developed on a review of limited data and updated when new relevant publications were revealed. Timing for deferrals of diagnostic-therapeutic procedures were mostly covered in the guidelines. Also, a support for the existing Polish recommendations on abnormal screening results in a subject of minor and major screening abnormalities terminology were given. The guidelines are obligatory for the specified COVID-19 pandemic period only and they might be changed depending on the new available evidence.

Key words: cervical cancer prevention; abnormal screening results; HPV testing; cervical cytology; selfsampling; SARS-CoV-2 pandemic; guidelines

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The recommendations present current management that can be modified and changed in justified cases, after careful analysis of a given clinical situation, which in the future may constitute grounds for their modification and updating.

Interim guidelines are obligatory for specified period only.

INTRODUCTION

The current national epidemiological situation rapidly changed due to the SARS-CoV-2 pandemic and a public health threat has arisen [1, 2]. The clinical management and diagnostic-therapeutic procedures performed so far in the secondary cervical cancer prevention in Poland need to

be revisited to avoid the unnecessary virus spreading. The temporary guidelines for the pandemics time have been already introduced by some countries [3] for the patients' management with abnormal screening tests results. Understanding the pandemic period restrictions, Polish interim guidelines for deferral of all non-essential medical office

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Table 1. Recommended management during the SARS-CoV-2 pandemic in women with the minor cervical cancer screening abnormalities. The different screening models were given

Model of screening and detected minor screening abnormalities	Recommended management*
Primary cytology	1. Deferral of diagnostic tests up to 6-12 months or 2. Continuation of screening that do not require leaving the place of residence with the use of previously sampled liquid-based cytologic material (if applicable)
ASC-US or LSIL	
ASC-US or LSIL followed by negative HRHPV testing	
ASC-US or LSIL followed by negative p16/Ki67 test	
ASC-US or LSIL followed by positive HRHPV N16/N18 and negative p16/Ki67 testing	
Primary co-testing	
ASC-US or LSIL HRHPV-negative	
ASC-US or LSIL HRHPV N16/N18-positive	

ASC-US — atypical squamous cells of undetermined significance; LSIL — low-grade squamous intraepithelial lesion; HRHPV — high-risk human papillomavirus type

*In individual cases the management might be modified depending on a clinical- and a patient-related status and/or depending on changes in a present healthcare environment

appointments and elective procedures in the secondary cervical cancer screening have been developed, pursuant to the President of the Polish Society of Gynecologists and Obstetricians (PTGiP) statement of the 20th of March 2020, and the joint recommendations of the Polish national consultants in obstetrics and gynecology together with the national consultants in perinatology of the 19th of March 2020 [4, 5]. Guidelines for possible deferral of diagnostic and therapeutic procedures in cervical cancer prophylaxis are obligatory for the specified COVID-19 pandemic period only.

RECOMMENDATIONS

The Interim guidelines are supplementary to the existing Polish recommendations [6–8] for the secondary cervical cancer prevention in the subject of a terminology for minor and major screening abnormalities.

Minor screening abnormalities — a definition and the recommended management

Minor screening abnormalities in the pre-colposcopic stage encompass the following screening test results:

- in the primary cytology-based model the following results:
 - ASC-US,
 - LSIL,
 - ASC-US or LSIL followed by negative HRHPV test,
 - ASC-US or LSIL followed by negative p16/Ki67 test,
 - ASC-US or LSIL followed by positive HRHPV type non-16/non-18 and negative p16/Ki67 test;
- in the primary cotesting-based model the following results:
 - ASC-US or LSIL HRHPV-negative,
 - ASC-US or LSIL HRHPV types non-16/non-18-positive.

Patients with minor cervical cancer screening abnormalities may have deferred of diagnostic tests up to 6–12 months [3].

The following management in the secondary cervical cancer prevention in women with abnormal minor screen-

ing tests results during the SARS-CoV-2 pandemic period is recommended (Tab. 1).

Continuation of previously started screening that do not require leaving the place of residence is recommended, especially with the use of the liquid-based preparation systems. This concerns molecular testing for the high-risk HPV (HRHPV) and immunocytochemical p16/Ki67 testing in a case of ASC-US or LSIL cytology result [6–8].

HRHPV selfsampling at home is recommended if a primary molecular testing is required, or if a reflex testing is required in case of an ASC-US or LSIL abnormal cytology result [9].

Major Screening Abnormalities – a definition and the recommended management

Major screening abnormalities in the pre-colposcopic stage encompass the following screening test results:

- in the primary cytology-based model the following results:
 - ASC-H,
 - HSIL,
 - ASC-US or LSIL followed by positive HRHPV type 16 and/or 18,
 - ASC-US or LSIL followed by positive p16/Ki67 test;
- in the primary cotesting-based model the following results:
 - HRHPV type 16 and/or 18 positive,
 - ASC-H, HSIL and AGC cytology results, regardless the HPV status;
- in the primary HRHPV-based selfsampling model - HRHPV type 16 and/or 18 positive. For HRHPV types non-16/non-18 positive results obtained in selfsampling a management algorithm depends on clinical situation and screening history.

Patients with major cervical cancer screening abnormalities may have deferred of diagnostic tests and treatment maximum up to 3 months [3].

Table 2. Recommended management during the SARS-CoV-2 pandemic in women with the major cervical cancer screening abnormalities. The different screening models were given

Model of screening and detected major screening abnormalities	Recommended management*
Primary cytology	Deferral of diagnostic tests or treatment maximum up to 3 months
ASC-H or HSIL	
ASC-US or LSIL followed by positive HRHPV 16 and/or 18	
ASC-US or LSIL followed by positive p16/Ki67	
Primary co-testing	
Positive HRHPV 16 and/or 18	
ASC-H, HSIL, AGC regardless HRHPV status	

ASC-US — atypical squamous cells of undetermined significance; LSIL — low-grade squamous intraepithelial lesion; ASC-H — atypical squamous cells cannot exclude HSIL; HSIL — high-grade squamous intraepithelial lesion; AGC — atypical glandular cells; HRHPV — high-risk human papillomavirus type

*In individual cases the management might be modified depending on a clinical- and a patient-related status and/or depending on changes in a present healthcare environment

The following management in the secondary cervical cancer prevention in women with abnormal major screening tests results during the SARS-CoV-2 pandemic period is recommended (Tab. 2).

Histologic HSIL treatment

Specific guidelines of the gynecological societies adapted for the COVID-19 pandemic vary worldwide in the framework of histologic HSIL priority treatment [10, 11] and available data and evidence are limited. The conization due to histologic HSIL is considered by the European Society of Medical Oncology as a low-priority medical procedure while including an oncologic cervical cancer treatment of all clinical stages [10]. At the end of April, the joint American statement (of the eight gynecological and non-gynecological societies) was revealed in which authors took into account an epidemiological situation in a post-peak infectious curve when the risk of SARS-CoV-2 infection will be diminished and 'a normal new' will appear. In that document for benign gynecological indications on re-introduction of hospital and office-based procedures for the practicing gynecologists the CIN2 or CIN3 (HSIL) treatment was incorporated as an elective surgery of high acuity [12, 13]. Therefore, Polish interim guidelines suggest in the decision-making that a prioritization of patients should be established after a careful analysis of a clinical status and a patient-related situation, and also should be modified according with the dynamically changing national healthcare environment related to COVID-19.

Invasive cervical disease

In patients with suspected invasive cervical disease the recommendations for a clinical management remain unchanged.

SUMMARY

The above Polish interim guidelines for a possible deferral of diagnostic and therapeutic procedures for patients

with abnormal screening tests results in cervical cancer secondary prevention in current SARS-CoV-2 pandemic present not the final management to proceed the patients with abnormal screening test results in the secondary cervical cancer prevention. They do not replace a full clinical assessment of each individual case. The guidelines should always be considered in the context of the patient's health interest. They may change depending on new available data.

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