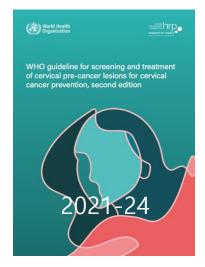
WHO Cervical Cancer Screening and Treatment Recommendations to Prevent Cervical Cancer

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Cancer Team
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2021 Cervical cancer screening and treatment guidelines HPV testing – (triage) - ablative treatment



WHO guidelines for the use of thermal ablation for cervical pre-cancer lesions

2019

GENERAL FEMALE POPULATION

Screen-and-Treat or Screen, Triage & Treat

Primary screening test

- High-performance HPV DNA Test
 - On provider- or self-collected samples
 - Starting at age 30
 - Every 5 to 10 years
- High-performance mRNA Test
 - Only on provider-collected samples
 - Every 5 years

<u>Triage</u> with HPV16/18, VIA, cytology, colposcopy or dual-stain cytology

Treatment

- Ablative treatment if eligible
- Referral for excision or other

WOMEN LIVING WITH HIV Screen, Triage & Treat

Primary screening test

- High-performance HPV DNA test
 - On provider- or self-collected samples
 - Starting at age 25
 - Every 3 to 5 years

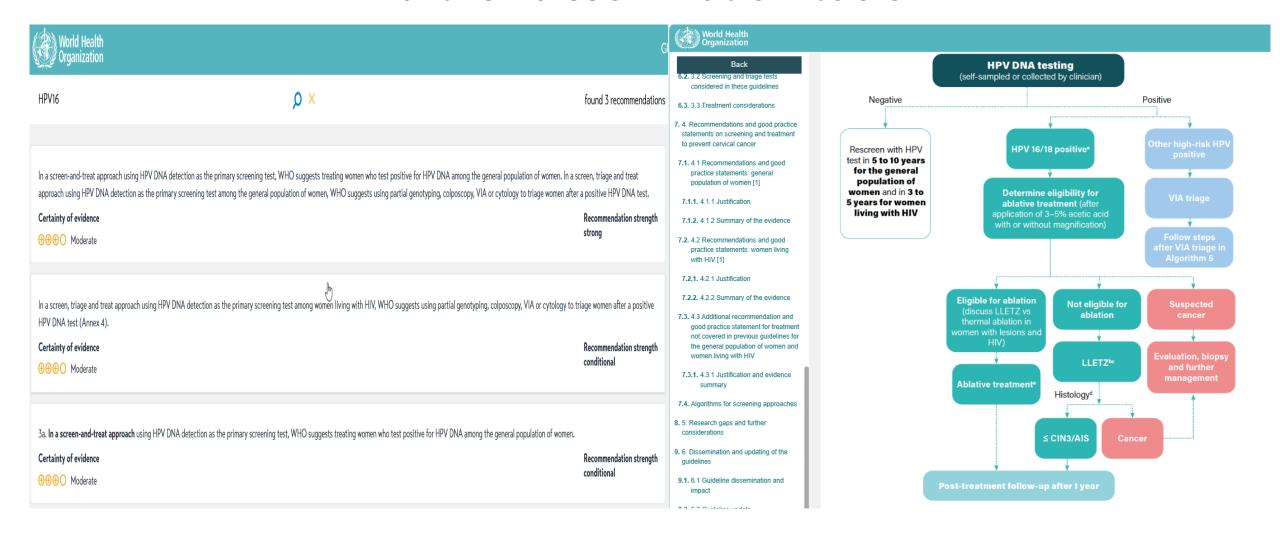
<u>Triage</u> with HPV16/18, VIA, cytology or colposcopy

Treatment

- Ablative treatment if eligible
- Referral for excision or other



WHO Cervical Cancer Screening and Treatment Guidelines Further dissemination tools



Updating recommendations - Systematic reviews

Emerging considerations when carrying recommendations & systematic reviews process

When is there sufficient evidence to consider formulating or updating a recommendation?

Are other technical products needed to support the guideline process and public health needs?

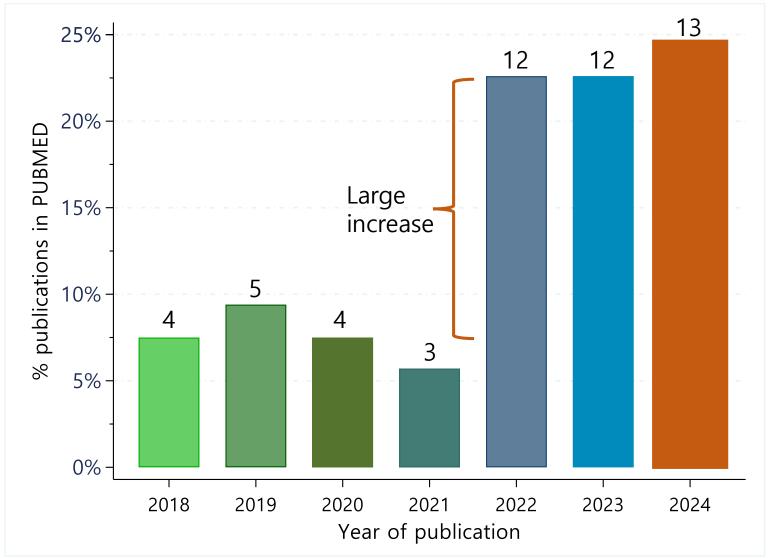
Do we know the metrics to use in the performance evaluation of novel technologies?

1. When is there sufficient evidence to consider formulating or updating a recommendation?

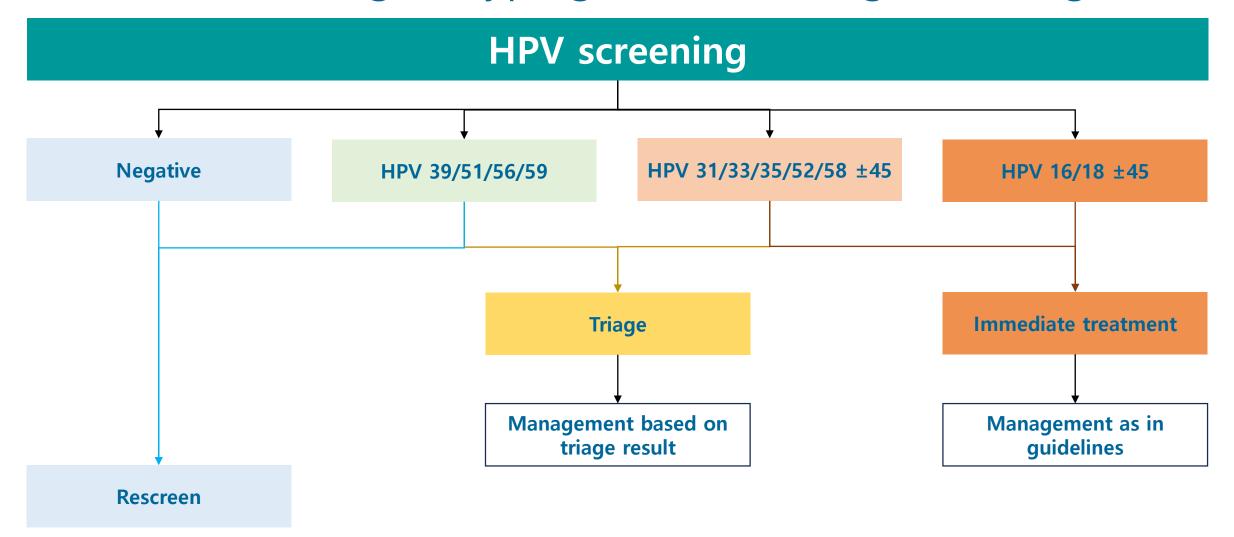
 Monitoring through regular systematic reviews



PUBMED publications "referring" HPV extended genotyping 2018-24



Possible extended genotyping-based management algorithms



Do other technical products need to be developed to support the living process and public health needs?

264 HPV tests but availability of affordable high-performance HPV tests remains limited!

Number of HPV molecular tests in the market



79% without clinical validation

WHO Target Product Profiles for HPV screening tests to detect cervical pre-cancer and cancer

- WHO TPPs guide and coordinate development of new health products with clear product characteristics, considering populations, access and equity from the outset
- HPV TPPs Technical Development Group (TDG) composed of 39 members
 - Multiple expertise, stakeholders and women's representatives
 - ✓ Representation balanced by WHO region

- WHO TPPs for HPV screening tests aim to direct tests developers & manufacturers to prioritize technologies that can contribute to countries' efforts to reach 70% screening coverage elimination target
- TPPs outline desired profile of a product, with two characteristics per parameter:
 - ✓ <u>minimal</u> (lowest acceptable)
 - ✓ <u>preferred</u> (ideal)



Target Product Profiles for HPV screening tests to detect cervical pre-cancer and cancer



Step 1. Identify need Step 3. Convene group Step 5. Consult Step 7. Publish TPP • Post draft 0.1 for 28 days Upload to WHO website Literature review Convene steering group Consult experts Convene TPP Upload to TPP Directory **Development Group** Landscape analysis

standard procedure

Second edition



Step 2. Scope

- Prepare scope and purpose
- Seek initial clearance within WHO
- Determine there is an external audience

Step 4. Develop

- Prepare draft v0
- Prepare v0.1 with TPP **Development Group**

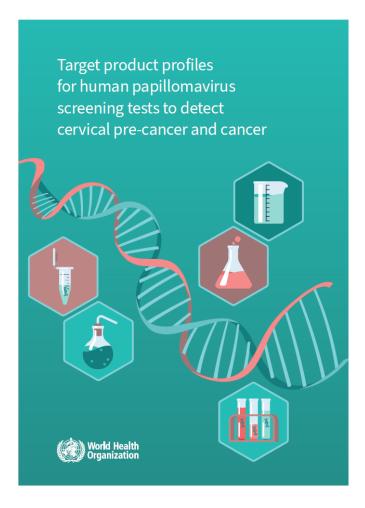
Step 6. Finalize

- Prepare v1.0
- Refine with TPP **Development Group**
- Clearance within WHO

Step 8. Disseminate

- Using networks
- Peer-reviewed journals
- Retire after 5 years

WHO Target Product Profiles for HPV screening tests



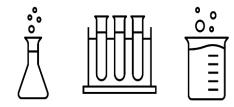
In person meeting in Costa Rica November 2023



https://www.who.int/publications/i/item/9789240100275

2 Target Product Profiles for HPV screening tests

✓ For laboratory use



✓ For point-of-care use



We need to scale up!

Minimal and Preferred Characteristics 41 Parameters across Eight Domains

Scope

Performance

Conditions

Connectivity

Technical specifications

Design and operation

Quality and standards

Cost



Genotype Spectrum and result output

LABORATORY

✓ Minimal:

8 cHPV, Groups 1a, 1b and 1c at least TWO signals:

- 16 individual or grouped with 18/45
- o 31, 33, 35, 52, 58 pooled

✓ Preferred:

12 cHPV, Groups 1a, 1b, 1c and 1d at least FOUR signals:

- 0 16
- o 18, 45 individual or pooled
- o 31, 33, 35, 52, 58 pooled
- o 39, 51, 56, 59 pooled

POINT-OF-CARE

SubgroupCarcinogenic HPV types includedGroup 1aHPV16Group 1bHPV18, HPV45Group 1cHPV31, HPV33, HPV35, HPV52 and HPV58Group 1dHPV39, HPV51, HPV56 and HPV59

✓ Minimal:

8 cHPV, Groups 1a, 1b and 1c ONE pooled signal (P/N):

0 16, 18, 45, 33, 58, 31, 52, 35

✓ Preferred:

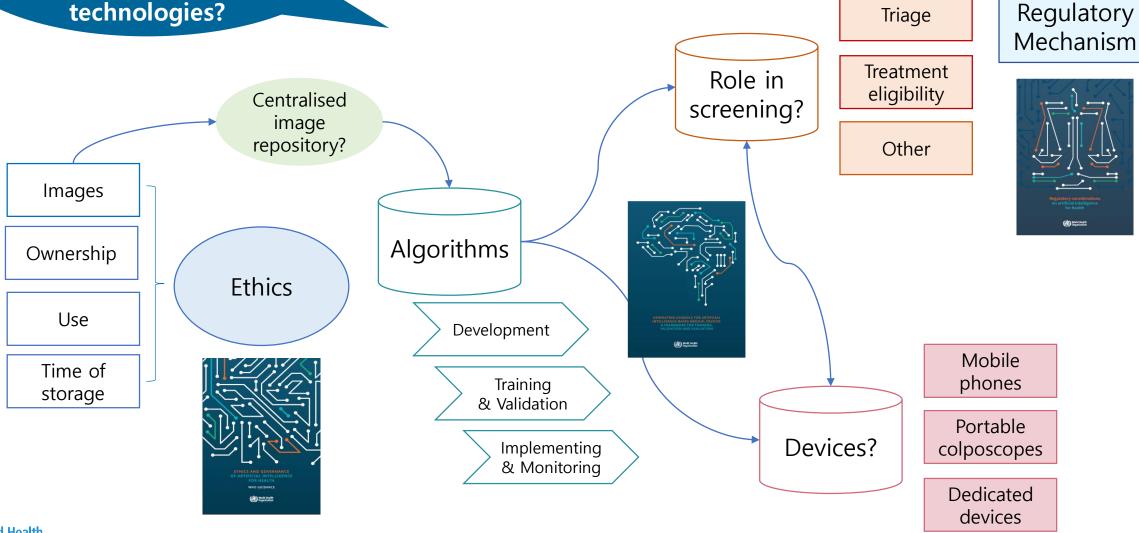
12 cHPV, Groups 1a, 1b, 1c and 1d at least TWO signals:

- 16 individual or grouped with 18/45
- o 31, 33, 35, 52, 58, 39, 51, 56, 59 pooled



3. Do we know the metrics to use in the performance evaluation of novel technologies?

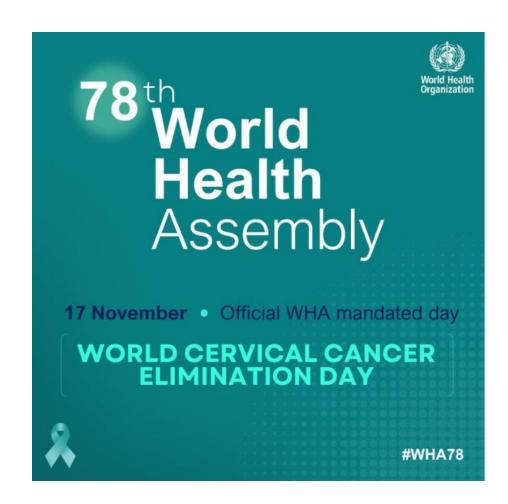
Artificial intelligence for cervical cancer screening and treatment





In summary

- Technology is evolving fast, a process for updating recommendations is essential
- Recommendations should be based on evidence on performance and feasibility to facilitate countries to make informed decision when adoption emerging technologies
- Other workstreams will require attention, such as the WHO Target Product Profiles for HPV screening tests, having dialogues with the private sector and WHO Prequalification IVDs
- Validation criteria appropriate for some novel technologies such as AI needs to be carefully defined and agreed upon by experts and stakeholders





make to all women and girls – to spare millions hardmyner Kanustofta preventable cancer

No one left behind

Many thanks to the GDG, WHO Secretariat, stakeholders and multiple collaborators

