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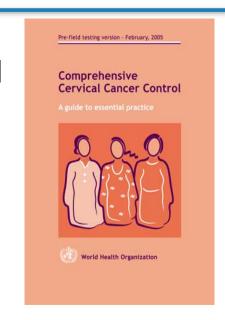
New guidelines for cervical cancer prevention and control



PAHO Women's Cancer Initiative Washington DC, 5-7 February, 2013

WHO Mandate to Develop Norms and Guidelines

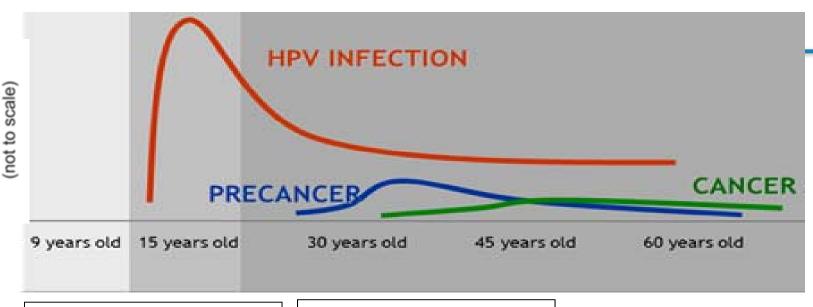
- WHO Member States rely on WHO for expertise and guidance with respect to cervical cancer control through the development of international norms and guidelines and promoting their implementation
- The Guidelines Review Committee (GRC) was established in 2007 to ensure that WHO guidelines are developed in ways consistent with best practice



 The second edition of the guidelines are comprehensively updated to take account of developments in screening, diagnosis, and treatment of cervical cancer



Comprehensive approach: Programmatic interventions over the life course to prevent HPV infection and cervical cancer



PRIMARY PREVENTION



Girls 9-13 years

HPV vaccination

Population prevalence

From 10 years old and onward

Health education and services, for example:

- •Sexual health education tailored to the age group
- •Providing contraceptive counseling and services including condoms
- Prevent tobacco use and support cessation*

SECONDARY PREVENTION



Women > 30 years of age Screening and treatment

- "screen and treat" with low cost technology VIA followed by cryotherapy • HPV testing for high risk HPV types (e.g.
- types 16, 18 and others)

TERTIARY PREVENTION



All women as needed

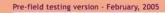
Treatment of invasive cancer at any age

- Ablative surgery
- Radiotherapy
- Chemotherapy



WHO standards for cervical cancer prevention and control

http://www.who.int/
reproductivehealth/en/

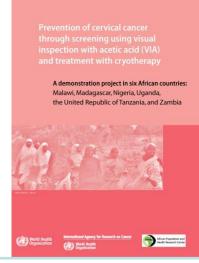


Comprehensive Cervical Cancer Control

A guide to essential practice























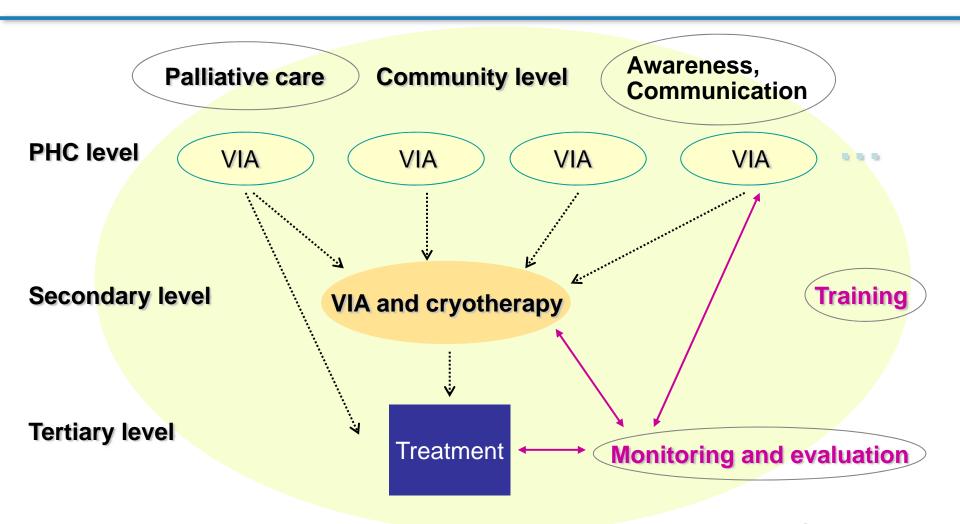






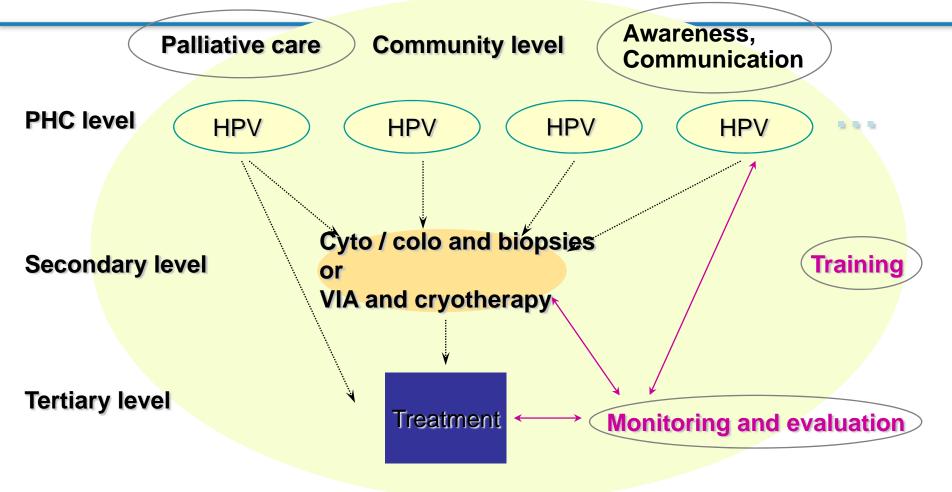


Strengthening Cervical Cancer Prevention Programme - Operational framework





Strengthening Cervical Cancer Prevention Programme – New algorithm?





Technical specifications for cryotherapy equipment

Technical specification for cyrotherapy equipment

This manual addresses key issues that will ensure the procurement and effective use of quality assured cryotherapy equipment to support the early management of precancerous cervical lesions as part of a comprehensive cervical cancer prevention programme.

Contents:

- Technical Basis Paper. Cryotherapy equipment for the treatment of pre-cancerous cervical lesions
- Generic Specification. Cryotherapy equipment for the treatment of pre-cancerous cervical lesions
- Advice and guidance, gas supplies for cryotherapy treatment of precancerous cervical lesions
- Recommendations for handling gas cylinders
- Procurement guidance.

WHO technical specifications Cryosurgical equipment for the treatment of precancerous cervical lesions and prevention of cervical cancer World Health Organization WHO guidelines Use of cryotherapy for cervical intraepithelial neoplasia



QA/QC for VIA-cryotherapy based programmes

Companion guides to (C4GEP)

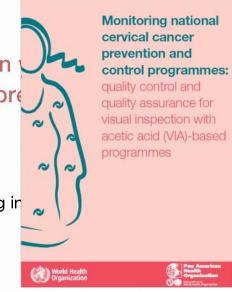
Quality control and quality assurance for visual inspection vacetic acid (VIA) and for cryotherapy for cervical cancer preand control

Intended primarily for programme managers and other stakeholders working in health programmes for cervical cancer prevention and control.

Purpose

This guide focuses on quality control and quality assurance for VIA and cryotherapy, given that both have been extensively evaluated through cross-sectional studies, prospective randomized trials and demonstration programmes.

The recommendations provided in this document need to be adapted to national policies, health systems, needs, language and culture.



Cervical cancer indicators

Performance indicators

Screening rate of the target population (women aged 30–49 years): Percentage of women aged 30–49 years who have been screened for the first time with VIA in the previous 12-month period.

Positivity rate: Percentage of screened women aged 30–49 years with a positive VIA test result in the previous 12-month period.

Treatment rate: Percentage of VIA-positive women receiving treatment in the previous 12-month period.

Result indicator

Coverage rate indicator: Percentage of women aged 30–49 years who have been screened with VIA or another screening test at least once between the ages of 30 and 49 years.

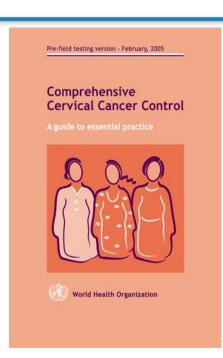
Impact indicator

Cervical cancer age-specific incidence.



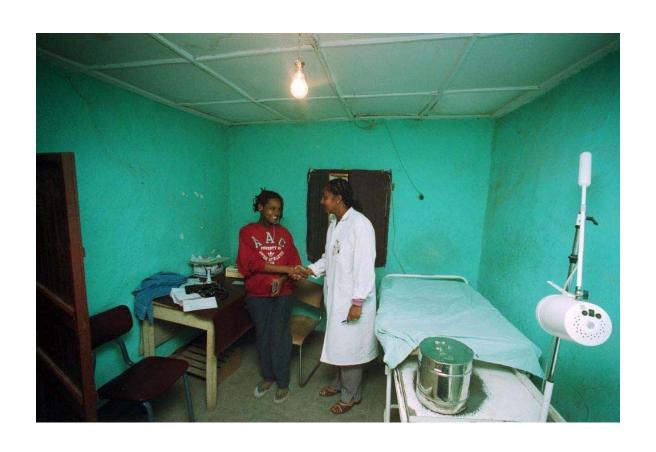
Purpose of the update

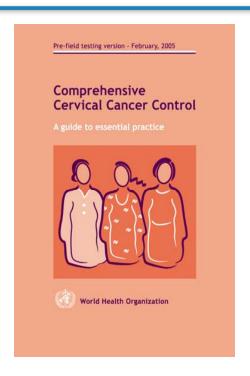
- Health education to be expanded
- HPV vaccines to be included
- New data on use of screening tests and algorithms
- New data on HIV and cervical cancer:
 - Natural history of HPV infection in HIV+ women
 - Age of first screening
 - Frequency of screening tests
 - Management of positive screening tests in HIV positive women (cryotherapy, LEEP) and follow-up, also safety issues
- HIV screening in women undergoing cervical cancer screening – how to incorporate?





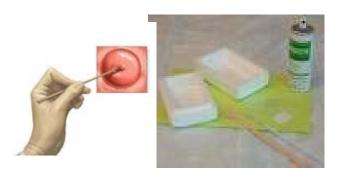
Screen and Treat





Which screening test for which population and where?

Conventional pap smear



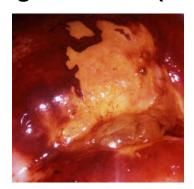
HPV DNA test



Visual inspection with acetic acid (VIA)



Visual inspection with Lugol's iodine (VILI)



HPV rapid **DNA** test



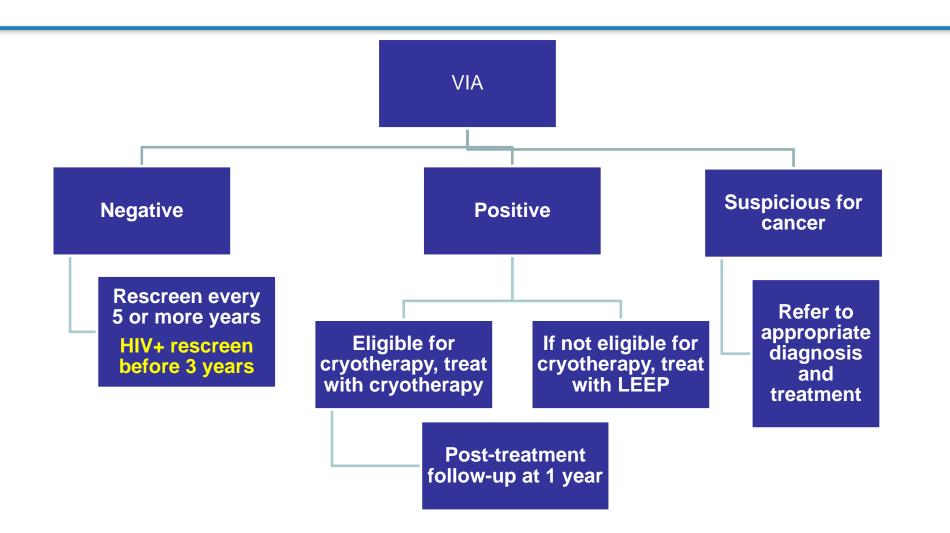


Characterictics of screening tests for secondary prevention

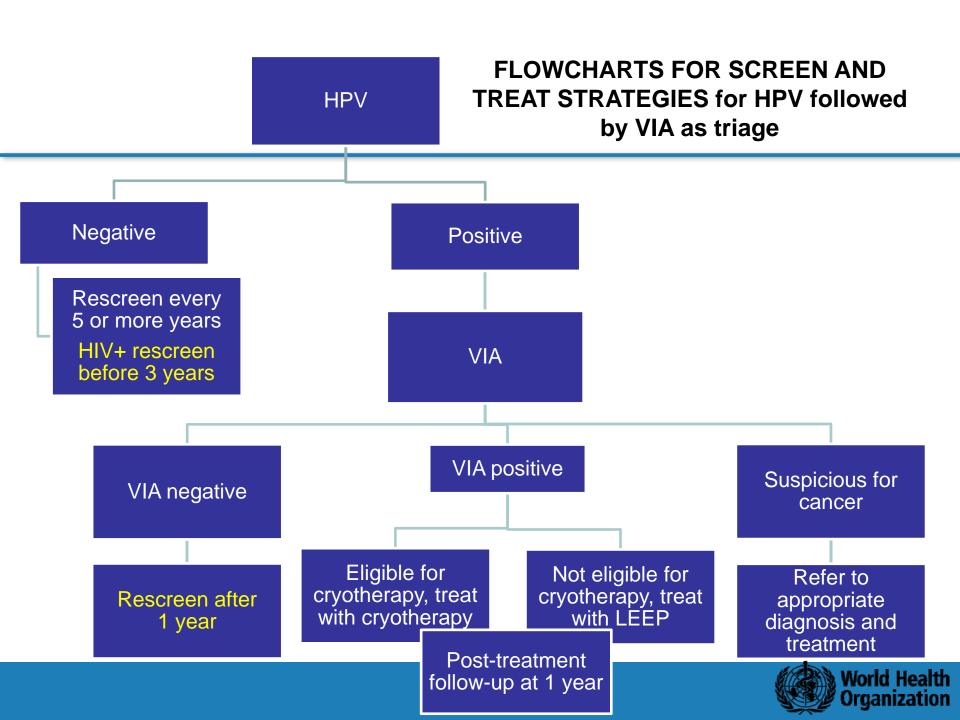
| Characteristics | Conventional cytology | HPV DNA tests | Visual inspection tests | |
|---|--|---|---|---|
| | | | VIA | VILI |
| Sensitivity | 47-62% | 82-100% | 67-79% | 78-98% |
| Specificity (for high-grade | 60-95% | 75-96% | 49-86% | 73-91% |
| lesions and invasive cancer) Comments | Assessed over the last 50 years in a wide range of | Assessed over the last decade in many settings in | Assessed over the last decade in many settings in | Assessed by IARC over the last four years in India and 3 countries in Africa. |
| | settings in developed and developing countries | developed and relatively few in developing countries | developing countries | Need further evaluation for reproducibility |
| Number of visits required for screening and treatment | 2 or more visits | 2 or more visits | Can be used in single-visit or 'see and treat' approach where outpatient treatment is available | |



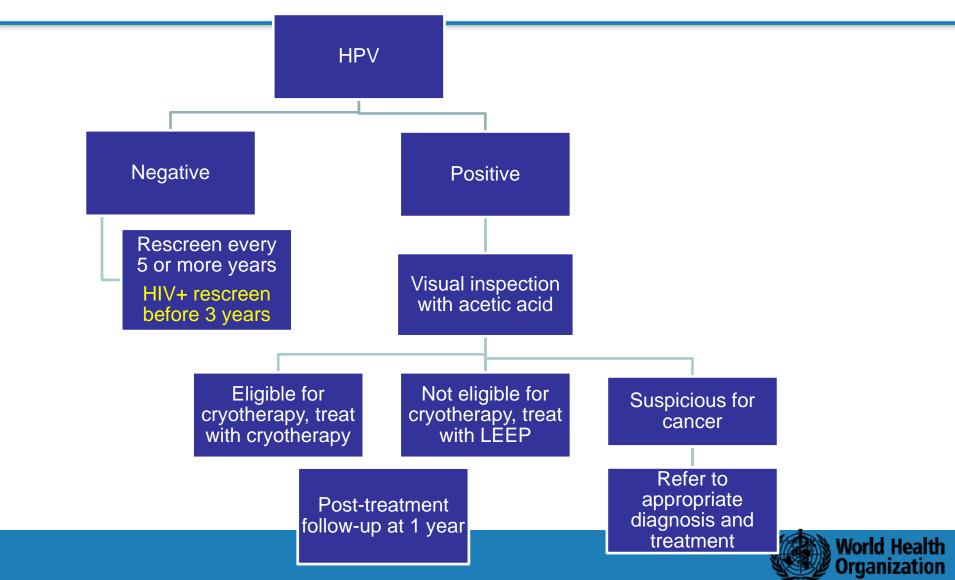
FLOWCHARTS FOR SCREEN AND TREAT STRATEGIES WITH VIA



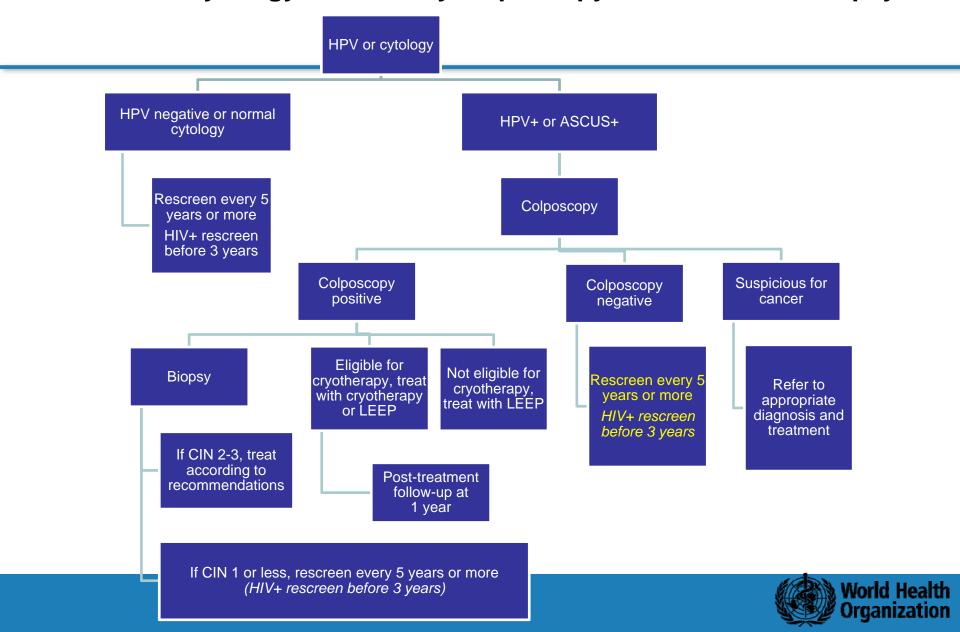




FLOWCHARTS FOR SCREEN AND TREAT STRATEGIES WITH HPV alone – VIA used to determine eligibility for cryotherapy



FLOWCHARTS FOR SCREEN AND TREAT STRATEGIES WITH HPV or cytology followed by colposcopy with or without biopsy



The issue



Overcoming the transfer and application of knowledge gap



To take evidence into practice



Scope of IR/OR

Any research producing practically usable knowledge (evidence, findings, information, etc) which can improve programme implementation (e.g., effectiveness, efficiency, quality, access, scale-up, sustainability) regardless of the type of research (design, methodology, approach) falls within the boundaries of operational research

Jane Kengeya-Kayondo



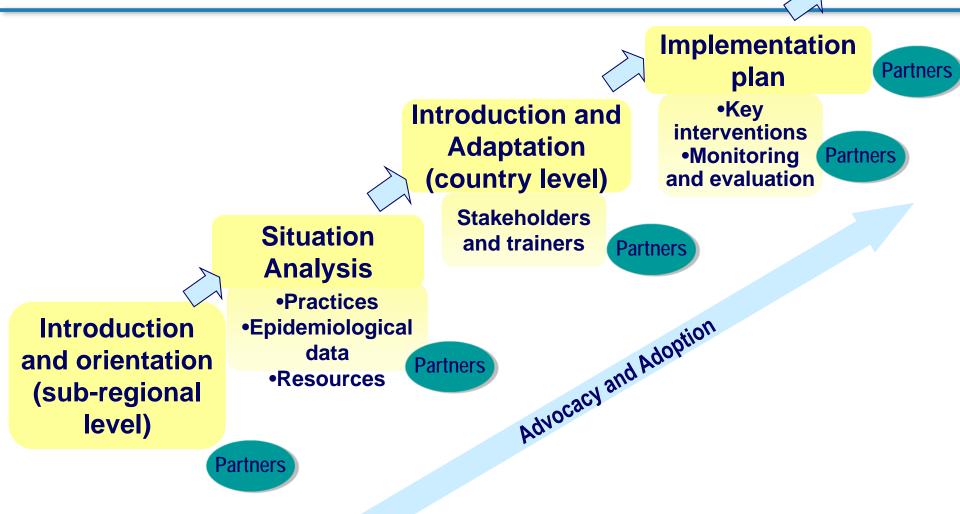
IR/OR Can Accomplish the Following:

- Identify and solve programme problems in a timely manner
- Help policy-makers and programme managers make evidence-based programme decisions
- Improve programme quality and performance using scientifically valid methods
- Help programme managers and staff understand how their programmes work



A process to introduce and adapt guidelines and tools

Scalingup





Example of programmatic linkages: sexual and reproductive health services

