

Cervical Cancer Control Current Practices

From Research to Practice: Training Course in
Sexual and Reproductive Health Research
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Situational Analysis: Cancer Control in Developing Countries

- 75% of the new cancer cases worldwide
- 5% of the world cancer resources
- 80-85% cases diagnosed at late incurable stages - if at all
- majority not covered with cancer care
- 5% women screened for Cx Ca compared to 40% in industrialised countries

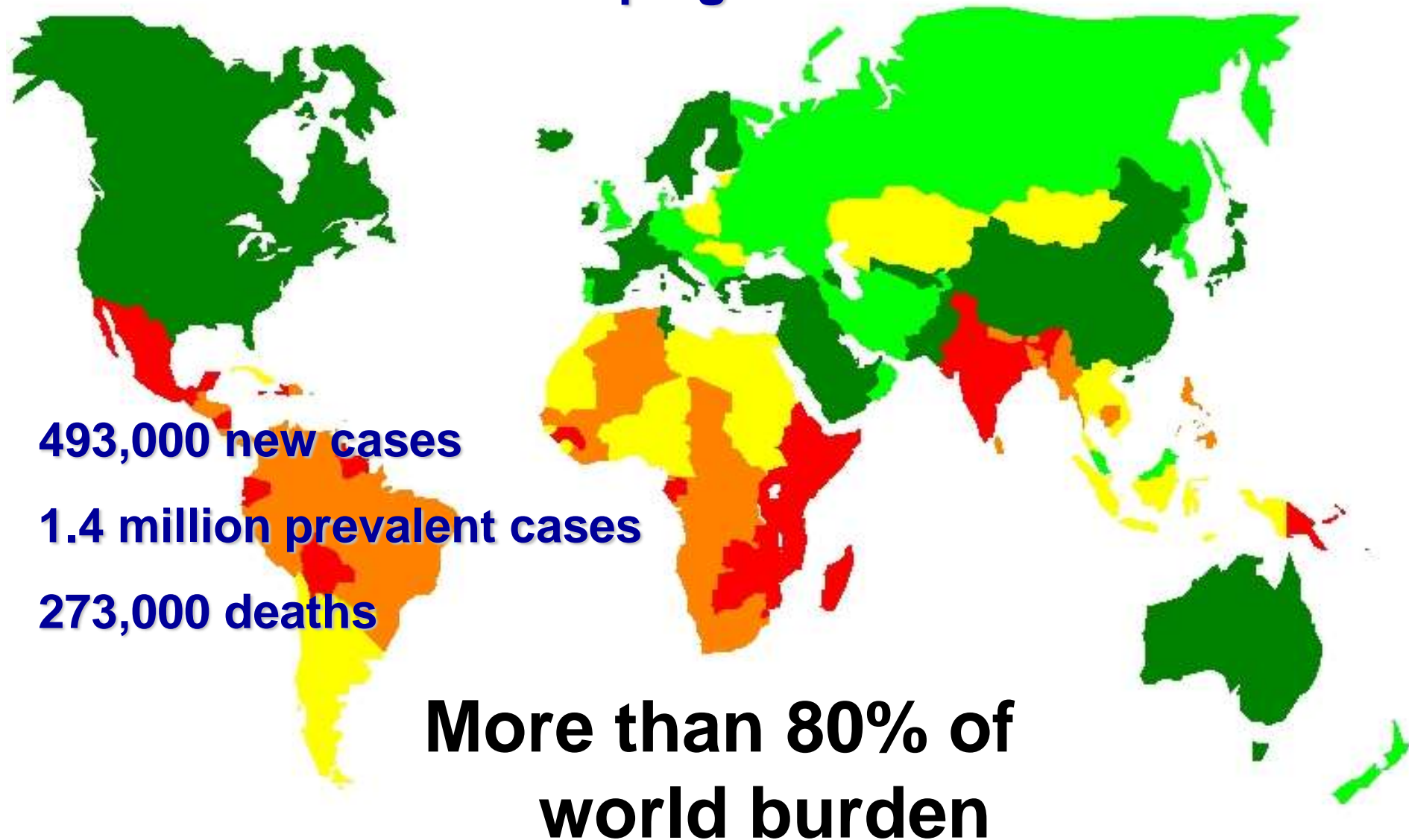
Priorities and strategies for the eight most common cancers worldwide

WHO, 1995

Site of cancer	Primary prevention	Early diagnosis	Curative therapy	Pain relief, palliative care
Lung	++	-	-	++
Stomach	+	-	-	++
Breast	+	++	++	++
Colorectum	+	++	++	++
Cervix	++	++	++	++
Mouth pharynx	++	+	++	++
Oesophagus	+	-	-	++
Liver	++	-	-	++

++ effective, + partly effective, - ineffective

Cervical cancer continues to be a major burden in most developing countries



■ < 3.9 ■ < 7.4 ■ < 11.3 ■ < 16.8 ■ < 53.5

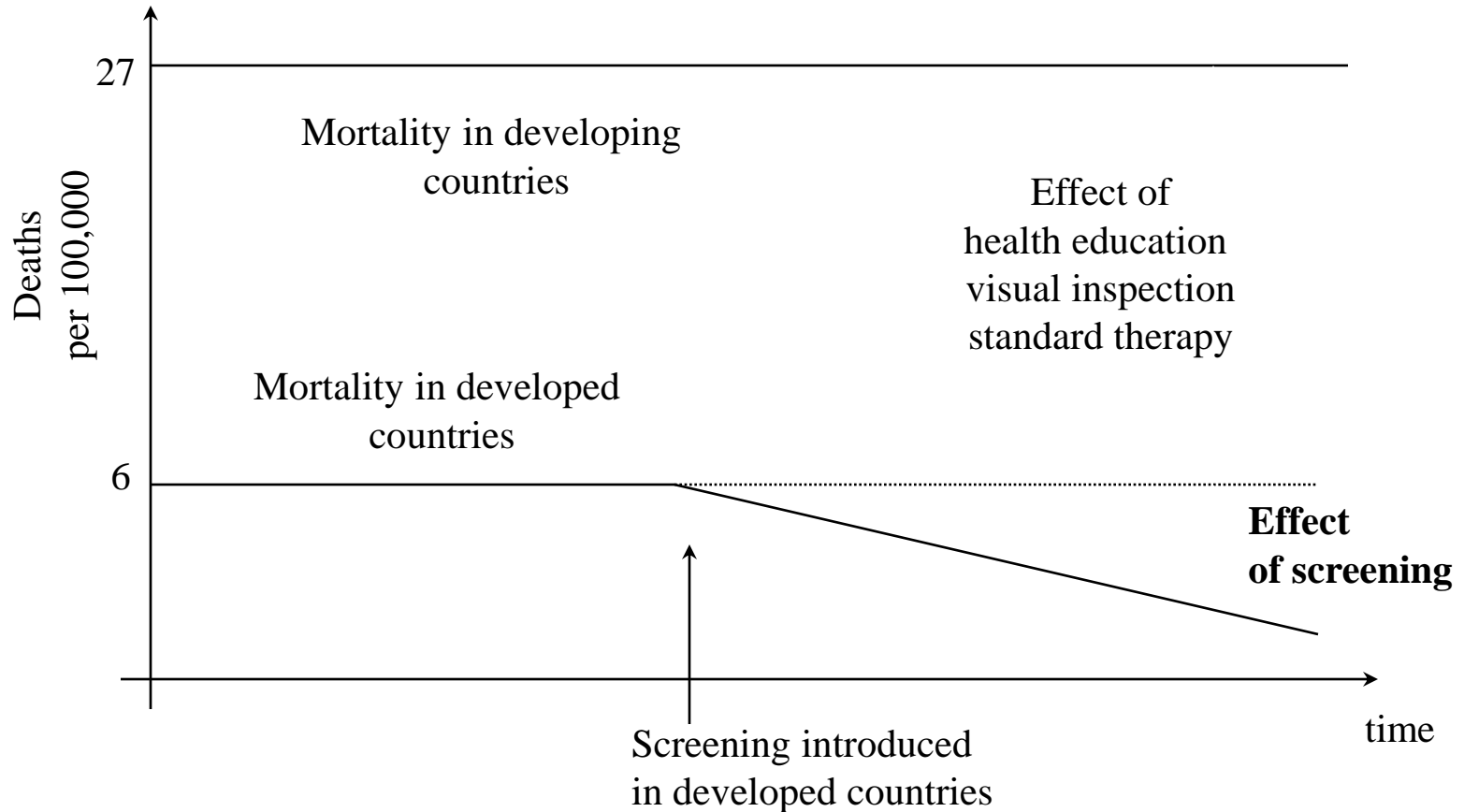
Estimated cases of Cx Ca in Regions and selected countries

IARC, 1999

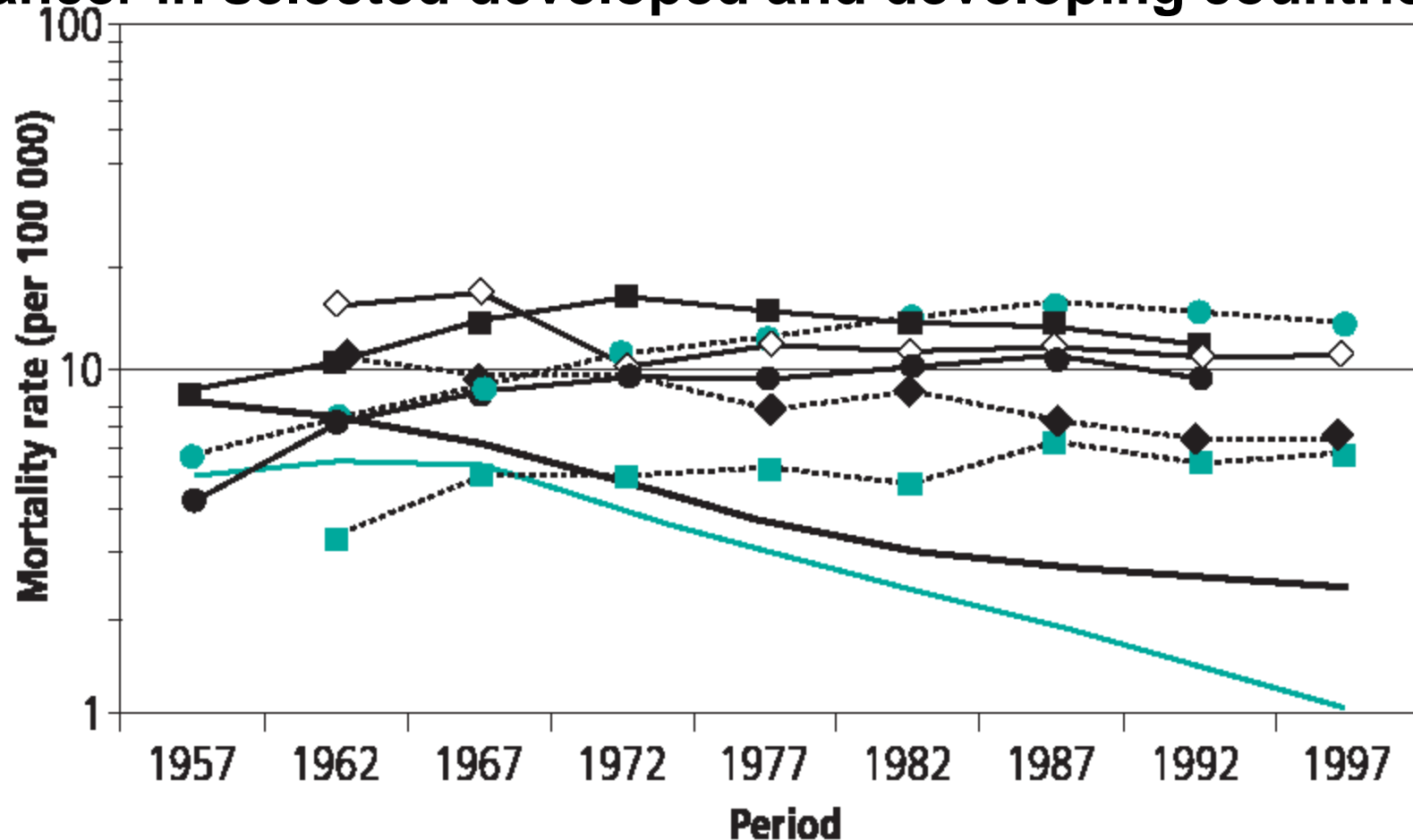
Region/Country	New cases per year
North America	15 700
Latin America	49 000
Europe	47 200
USSR	31 300
Africa	36 900
Asian excluding India & China	80 000
China	131 500
India	120 000
Australia/NZ	1 200

Cervical Cancer Control

M.I.Shafi & S.Nazeer. Colposcopy-Apractical guide; 2006



Trends in age-standardized mortality rate for cervical cancer in selected developed and developing countries



- Chile
- ◆ Costa Rica
- Colombia
- Cuba
- ◆ Singapore
- Mexico
- Finland
- USA

The **main obstacle** to a further improvement of the situation is the **high cost and labour intensive nature of all screening programmes**; for this reason, in most developing countries global preventive programs have been **rarely implemented** and almost never sustained; the usual picture is one of little financial support which entails poor quality and low coverage rates. These facts alone explain why **mortality rates in the less developed countries are twice those of the industrialized ones**

INCGC

A consortium of 32 organizations

- Local solutions
- External facilitation
- Pap smear technology revised

Akinremi TO, Nazeer S, Totsch M. (2005) Reduced alcohol use in the staining of Pap smears: a satisfactory low-cost protocol for cervical cancer screening. *Acta Cytologica* 2005; 49(2): 169-72

Cost-effective strategies

- Information systems / cancer registries
- Alternatives to Pap-smear (Education & empowerment + downstaging with simple VE; VIA; cervicography; spectroscopy; oculoscopy)
- HPV tests/vaccines

HPV VACCINES

Objective:

- To reduce Cx Ca burden globally
- To reduce costs of Cx Ca Screening Programmes in industrialised countries

RISK FACTORS FOR CERVICAL CANCER

- ❖ **Age at first sexual intercourse**
- ❖ **Multiple sexual partners**
- ❖ **OCPs**
- ❖ **Social economic status**
- ❖ **Smoking**
- ❖ **STDs**
- ❖ **HPV**

Papillomavirus types

host specifique, epitheliotropic

➤ **20** Animal types

➤ **100** Human types - **30** Infect genital tract :
Low risk / High risk (**20**
oncogenic)

Natural History of HPV

- Largely sexually transmitted
- Peak incidence: 20-24 yrs
- Incidence gradually declines upto 40-45 yrs
- May begin to increase slowly thereafter

(ref: Schiffman et al 1993; Bosch et al, 1995; Burk et al, 1996; Dillner et al, 1996; Meijer et al, 1999)

Natural History of HPV

- 80% infections transient: median range 12 mnths- no risk of CIN
- 10-20% infections persistent: high risk of CIN - only 30% of these progress if untreated
- RR of progression 40-180
- Persistence is the important factor for disease progression

(ref:Hildesheim et al,1994; Wheeler,1996; Koustky,1992)

HPV Prevalence

In cervical lesions

- Squamous carcinoma: 95% association
- HSIL/CIN II, III: 75 -95%
- LSIL/CINI: 60%
- ASCUS: 30%
- Adenocarcinoma: 12-30% association

(ref: Cuzick et al, 1992; Schiffman et al, 1993; IARC, 1995; Olsen et al, 1995)

**CERVICAL CANCER IS A RARE
LONG-TERM OUTCOME OF PERSISTENT
INFECTION WITH ONE OR MORE
OF HIGH-RISK HPV TYPES**

(16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 68, 73, 82)

HPV Vaccines Available

➤ **Polyvalent (Gardasil)**

VLPs of HPV 16; 18; 6; 11

Schedule: 3 IM injections – 0,2,6 months

➤ **Bivalent (Cervarix)**

VLPs of HPV 16; 18

Schedule: 3 IM injections – 0,1,6 months

Available HPV Vaccines

Practical Facts

- Approved by health authorities in Western countries since 2006
- Currently only approved for use in girls 10-14 years, prior to sexual debut
- To have beneficial effects the vaccination coverage should be at least 70-80% and its efficacy should last longer than 15 years

Available HPV Vaccines

Clinical Facts

- No therapeutic effect against prevalent HPV infection
- To be given before exposure – before sexual debut
- Efficient: 98-100% against CIN2+; adenocarcinoma in situ & genital lesions
- Effect on invasive Cx Ca to be proven
- Cross protection against non-vaccine types – partial at best
- Currently approved for vaccination of girls 10-14 years, with catch-up programmes for girls upto 26 years
- To date protection proven upto 7.3 years

Available HPV Vaccines

Expected Benefits

- HPV vaccines will not eradicate Cx Ca
- Effect on Cx Ca incidence will only be evident in 20-30 yrs
- Decrease (25-50%) in abnormal pap-smears
- decreased excisional treatment for high grade lesions
- Reduction in genital warts & cancers
- Beneficial in countries with and without Cx Ca screening programmes

Concluding Remarks

- Cx Ca is a public health issue
- Screening programmes – cost-effectiveness
- HPV is one of the main aetiological factors
- HPV in screening – no consensus
- HPV Vaccines – THE Plausible SOLUTION

Concluding Remarks

- HPV Vaccines – Key pending issues:
 - ? Duration of immune response – booster dose
 - ? Appropriate age of application
 - ? Cross immunisation – 30% vs 70%; geographic differences in HPV type prevalence
 - ? Monitoring methodologies
 - ? Different population groups – males; SIDA
 - ? Reduced participation in screening
 - ? Cost-benefit – continued screening

Ref.: S.Nazeer et al. European Consensus Statement on 'HPV Vaccination and Colposcopy'. *Prepared on behalf of the European Federation for Colposcopy (EFC)*, May 2010.