Role of screening in cancer control

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Purpose of screening

- reduce mortality
- sometimes reduce incidence
- improve quality of life

Means of screening

- early diagnosis
- finding preinvasive lesions

WHO criteria

- 1. Important health problem
- 2. Acceptable treatment
- 3. Resources available
- 4. Preclinical phase
- 5. Screening test
- 6. Acceptable test
- 7. Natural history known
- 8. Treatment policy agreed
- 9. Cost acceptable

Sufficient criteria

- Objective is defined (in terms of health)
- Objective can be made quantitative
- There is scientific evidence on effect of screening on health

Health services activity e.g. routine screening

- There is an objective
- There is a chain of actions
- There is evidence on effect

Change in outocome is the objective.

Outcome in cancer screening is

- Length of life, mortality
- Quality of life, mastectomy

Infrastructure for Screening Programme

- 1. Population
- 2. Individuals
- 3. Coverage and attendance
- 4. Field facilities
- 5. Laboratory facilities
- 6. Quality control of 4 and 5
- 7. Facilities for confirmation
- 8. Facilities for treatment
- 9. Referral system
- 10. Evaluating and monitoring

Infrastructure provides the essential elements or the chain of actions of a screening programme

- It is causal chain
- Cf treatment chain
- Cf path of patient

The causal chain

- Effect assumes that each round and link works
- Diagnostics only is not sufficient
- The theory of screening is derived from the theory of health services research

Evidence is

- The effect of program on outcome
- Outcome is health
- Empirical research with scientific method

What is the scientific method that provides evidence for public health policy

Direct evidence, conclusive

- randomised allocation of screening within the routine
 Indirect evidence, inconclusive
- time trends
- geographical differences
- screen detected cases
- test validity
- survival difference between localised and nonlocalised disease

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Evidence on effect

- Cervix cancer
- Breast cancer
- Colorectal cancer
- Prostate cancer

Evidence on effect

- Pap-test
- HPV-test
- Visual inspection
- Mammography
- FOBT
- PSA

Evidence on harm

- Overdiagnosis etc.
- Cost
- Lead time
- QoL

Finnish health services

- Responsibility of municipality
- More than 400 in number
- Size from 200 to 550 000
- Expenses subsidised by governement
- Guided and regulated less and less by governement

Screening as public health policy

- For cervical cancer since 1963
- For breast cancer since 1987
- For colorectal cancer since 2004

Smaller and smaller effects

1960's	cervical cancer	80%
1980's	breast cancer	30%
2000's	colorectal cancer	20%
2020's	prostate cancer	?%

What new in cervix cancer

Implementing new tests by randomised design

- automation, Papnet
- HPV-test, Hybrid capture II

Improvement in sensitivity or increase in overdiagnosis

What new in breast cancer

digital mammography

Danger of uncontrolled implementation

What new in colorectal cancer

- First organized programme as public health policy
- Individual level randomised election of invitees

What new in prostate cancer

- Randomised screening trial of 80 000
- Test sensitivity high, episode sensitivity low
- Effect on mortality small
- Overdiagnosis large
- QoL unknown

What new in health policy

- Less regulation by government
- Less guidelines by government
- More competition between providers
- More freedom to elect by municipalities

Consequences of health policy

- More emphasis on cost
- Less emphasis on health
- Loss in effectiveness

Main characteristics of screening in Finland

- Organised programmes with high effectiveness and low cost
- Early implementation of public health policy before opportunistic screening is common
- An advanced infrastructure that allows active design and unbiased evaluation for outcome
- Implementation of the routine screening by experimental design with randomisation
- Change of technology (test) in the routine by experimental design with randomisation

Role of screening in cancer control

Deaths from cancer in the Nordic Countries in 2010-2015

Primary site	Screening		Prevented	
	No	Yes	No	%
Present				
Breast	5300	4300	1000	18
Cervix	1600	130	1500	91
Future				
Colorectum	8300	6900	1500	18
Prostate	5000	4100	900	18
All sites	69000	64000	4900	7

Life years gained per year in the Nordic Countries

Deaths avoided 4900

Life years gained 27000

Good Q LYG 21000

To screen or not to screen I

- Effect 7 per cent of all cancer deaths
- Effect one year prolongation in 1000 years of life
- Effect small but tried and tested
- Almost all interventions have at most a small effect in medicine

To screen or not to screen II

Harm is important

Benefit vs Harm (e.g. GQLYG)

- Evidence (data) is limited
- Theory is limited
- Values i.e. weights are biased
- Cost is not a primary issue

Organised programme

- More effective than the spontaneous one
- Produces less harm than the spontaneous one
- Can be evaluated (by randomised design)
- Can be stopped

If routine screening is started

- Organise a programme
- Find the evidence on effectiveness i.e. observation on reduced mortality
- Do not trust on indirect evidence
- Do not trust on nonexperimental evidence