

Wednesday, February 8

Prognostic biomarkers for cervical HSIL progression **MSS 01** 

Room 0B

Chair: K. Louvanto (Finland) • B. Nedjai (UK)

10.00 • 11.30

This session will highlight strategies that could identify women with a progressive precancer cervical lesion to immediate referral to colposcopy compared to follow-up testing of women with intermediate risk. These strategies are needed to maximize the benefits of cervical cancer screening and follow-up testing while decreasing the potential harm as cost of unnecessary colposcopy referrals and treatments.

 Introduction B. Nedjai (UK)

- Risk factors and natural history for HSIL progression
- Role of HPV genotypes and methylation in predicting progression
- · Host methylation biomarkers in predicting regression
- · Host and viral methylation panel in predicting progression
- Multi omics approach to inform cervical lesion progression
- Discussion and Q&A

**MSS 02** 

A. B. Moscicki (USA)

**B. Wisman (Netherlands)** 

H. Berkhof (Netherlands)

K. Louvanto (Finland)

B. Nedjai (UK)

K. Louvanto (Finland)

& B. Nedjai (UK)

Lunch Break 11.30 • 13.30

New guidelines for validation of HPV tests	Doom OD
	Room 0B

Chair: M. Arbyn (Belgium) • M. Poljak (Slovenia)

for cervical cancer screening

13.30 • 15.00

- M. Arbyn (Belgium) Introduction
  - & M. Poljak (Slovenia)
- Clinical significance of viral load or signal strength J. Bonde (Denmark)
- Analytical criteria for validation of tests with HPV genotyping capacity
- Criteria for sample adequacy assessment
- Validation of HPV point-of-care tests
- Validation of HPV tests on self-samples
- New clinical validation guidelines
- Discussion and Q&A

J. Dillner (Sweden)

K. Cuschieri (UK)

M. Almonte (France)

P. E. Castle (USA)

M. Arbyn (Belgium)

M. Arbyn (Belgium)

& M. Poljak (Slovenia)

Coffee Break 15.00 • 15.30



Wednesday, February 8

MSS 03

Monitoring HPV vaccine impact in males (including in MSM programs)

Chair: A. Giuliano (USA) • J. Palefsky (USA)

Room 0B **15.30 • 17.00** 

Gender neutral vaccination is routinely recommended in many countries but programs to vaccinate boys and men were generally initiated several years after initiation of programs to vaccinate girls and women. Consequently, less is known at this time of the impact of vaccination of boys and men than among girls and women. This session will review current knowledge of the serologic responses to HPV vaccination in boys and men; duration of response; impact of vaccination on special populations of men including men who have sex with men and those living with HIV; and status of vaccination programs.

• Introduction	A. Giuliano (USA) & J. Palefsky (USA)
• Duration of serologic response and clinical protection after HPV vaccination in men	A. Giuliano (USA)
<ul> <li>One-dose and two-dose vaccination in men - what do we know?</li> </ul>	A. Luxembourg (USA)
• HPV vaccination in heterosexual men and MSM: impact of sexual orientation	J. Palefsky (USA)
• HPV vaccination in people living with HIV - what do we know and what are	G. Ellsworth (USA)
the key questions going forward	
• HPV vaccination of men in the United States: recommendations, coverage	L. Markowitz (USA)
and vaccine effectiveness - vaccine coverage and effectiveness of male	
vaccination among men in the US	
Discussion and Q&A	A. Giuliano (USA)
·	& J. Palefsky (USA)
	, , ,



Thursday, February 9

MSS 04

# Scientific approaches towards improving the cervical cancer elimination strategies

Chair: J. Dillner (Sweden) • E. Franco (Canada)

M. Lehtinen (Finland)

Room 0B

8.00 • 9.30

• Introduction	J. Dillner (Sweden)
	- E. Franco (Canada)
	& M. Lehtinen (Finland)
Worldwide HPV vaccination and screening coverage	L. Bruni (Spain)
• Eliminating cervical cancer via vaccination & screening of young adult	M. Elfström (Sweden)
women	
<ul> <li>How to protect unvaccinated, marginalized women?</li> </ul>	P. Gray (Finland)
<ul> <li>Ethical and legal aspects of girls-only vs. gender-neutral vaccination</li> </ul>	M. Logel (Canada)
<ul> <li>Sustainability of immune protection gained from HPV vaccination</li> </ul>	F. Mariz (Germany)
<ul> <li>Sustainability and resilience of HPV vaccination programs</li> </ul>	🖵 I. Baussano (France)
<ul> <li>Cost-effectiveness of HPV vaccination and screening programs</li> </ul>	S. Vänskä (Finland)
Discussion and Q&A	J. Dillner (Sweden)
	- E. Franco (Canada)

**☐** Video presentation

& M. Lehtinen (Finland)



Thursday, February 9

**MSS 05** 

#### Debate on controversial topics

Room 0B

Chair: K. Cuschieri (UK) • E. Franco (Canada) • T. J. Palmer (UK)

9.30 • 11.00

Debate sessions have been a popular offering in EUROGIN congresses since the 1990s. Pairs of leaders in the field capture the arguments on opposing sides of controversial or hot topics in HPV science and its practical aspects, such as vaccination, cervical cancer screening, and disease etiology. They present their arguments and then debate each other. The session in 2023 will showcase debates between camps on five key areas: (i) clinical utility of HPV genotyping, (ii) adequacy of single-dose vaccination, (iii) VIA vs. HPV for cervical cancer screening in low resource settings, (iv) value of biomarkers for primary screening, and (v) feasibility of ending cervical cancer screening.

• Introduction E. Franco (Canada)

 DEBATE #1 - Do we need information on HPV types other than 16/18 in cervical cancer screening?

» Yes/Pro side J. Bonde (Denmark)

» No/Con side H. Berkhof (Netherlands)

» Debate

 DEBATE #2 - Is a single dose of HPV vaccine adequate for the long-term elimination goal?

» Yes/Pro side P. Bloem (Switzerland)

» No/Con side J. Dillner (Sweden)

» Debate

• DEBATE #3 - Should VIA be replaced by HPV screening in low resource

settings?

» Yes/Pro side S. De Sanjosé (Spain)

» No/Con side

» Debate

DEBATE #4 - Are biomarkers suitable for primary screening?

» Yes/Pro side

» No/Con side

» Debate

M. Poljak (Slovenia)

• DEBATE #5 - Can we stop cervical screening in high-income countries in

the next decade or two?

» Yes/Pro side E. Franco (Canada)

» No/Con side T. J. Palmer (UK)

» Debate



Thursday, February 9

MSS 06	Self-sampling in routine practice, operational and technical issues Chair: J. Bonde (Denmark) • C. Cocuzza (Italy)	Room 0B <b>14.00 • 15.30</b>
• Introductio	n: long term stability of self-sampling devices and media for HPV	J. Bonde (Denmark)
testing		& C. Cocuzza (Italy)
· ·	and learnings from implementation of self-sampling in low - and ome countries	R. Luckett (USA)
	ncer screening based on first-void urine self-sampling to reach eened women: ScreenUrSelf trial	S. Van Keer (Belgium)
	nd operational considerations for HPV self-sampling - lesson ouScreen trial	P. Sasieni (UK)
• Technical b	ottlenecks and mitigations of self-sampling	J. Bonde (Denmark)
• Appropriate	e media and improved QC for self- sampling	C. Cocuzza (Italy)
<ul> <li>Discussion</li> </ul>	and Q&A	J. Bonde (Denmark)
		& C. Cocuzza (Italy)

Coffee Break 15.30 • 16.00

# Prevention of cervical cancer among people Room 0B Room 0B Iiving with HIV Chair: A. Giuliano (USA)

Women living with HIV are at elevated risk of acquiring oncogenic HPV infections, having those infections persist, and progress to cervical pre-cancer and cancer. Unfortunately, the performance of tests utilized among women without HIV to screen for cervical cancer and to treat pre-cancer lesions perform sub-optimally among women living with HIV. Current research is focused on tailoring these interventions for women living with HIV in a variety of different geographic regions. Also needed are interventions to improve CIN treatment outcomes among women living with HIV.

• Introduction	A. Giuliano (USA)
Burden of cervical pre-cancer and cancer among PLWH - High level overview	S. Sudenga (USA)
of challenges with screening and treatment	
ULACNet trial 101 rationale and design	A. P. Ortiz (Puerto Rico)
ULACNet trial 202 rationale and design	T. Wilkin (USA)
• ULACNet trial 302 cervical cancer screening among WLWH in the Dominican	M. Madeleine (USA)
Republic: rationale and study design	
Challenges in CIN and cervical cancer treatment of PLWH	H. Botha (South Africa)
Research to improve CIN treatment among PLWH	E. Chiao (USA)
Discussion and Q&A	A. Giuliano (USA)



Thursday, February 9

MSS 08	HPV and host methylation as triage tools for cervical cancer screening in different clinical contexts  Chair: M. Clarke (USA) • C. Meijer (Netherlands)	Room 0B <b>17.30 • 19.00</b>
• Introductio	n	M. Clarke (USA) & C. Meijer (Netherlands)
• DNA methy	of S5 classifier performance in different populations rlation performance in women living with HIV of an HPV methylation and genotyping assay in different clinical	B. Nedjai (UK) C. Meijer (Netherlands) M. Clarke (USA)
Performance     Application	of methylation in self-collected urine samples of host cell methylation and/or HPV genotyping in women with ytology (ASCUS and HSIL)	S. Van Keer (Belgium) H. Berkhof (Netherlands)
	of methylation markers in the management of vulvar	M. Bleeker (Netherlands)
<ul><li>Monitoring</li><li>Discussion</li></ul>	of posttreatment CIN3 by methylation markers and Q&A	S. Dick (Netherlands) M. Clarke (USA)
		& C. Meijer (Netherlands)

#### **MSS - MAIN SCIENTIFIC SESSIONS**

Friday, February 10

MSS 09	VALGENT / VALHUDES Chair: M. Arbyn (Belgium) • C. Cocuzza (Italy)	Room 0B <b>8.00 • 9.30</b>
• Introduction	า	M. Arbyn (Belgium) & C. Cocuzza (Italy)
<ul><li>Performand</li><li>VALHUDES</li></ul>	e of HPV tests on vaginal self-samples, results of the Belgian	A. Latsuzbaia (Luxembourg)
<ul><li>Performand</li><li>VALHUDES</li></ul>	e of HPV tests on urine self-samples, results of the Belgian	S. Van Keer (Belgium)
	e of the OncoPrect HPV assay (SCR and QT assays) evaluated in an VALHUDES	C. Cocuzza (Italy)
<ul> <li>Findings fro</li> </ul>	m the Australian VALHUDES study	M. Saville (Australia)
• The VALGEN	NT V study design	S. Dhillon (Belgium)
• Evaluation o	of the RIATOL qPCR in the VALGENT and VALHUDES studies	D. Vanden Broeck (Belgium)
• Discussion a	and Q&A	M. Arbyn (Belgium)
		& C. Cocuzza (Italy)



Friday, February 10

**MSS 10** 

#### Impact of HPV vaccine on cancer

Chair: J. Lei (Sweden) • P. Sasieni (UK)

Room 0B

10.00 • 11.30

HPV vaccination has been approved to be highly effective. In this session, we will gather researchers from six countries who will present the latest updates regarding the impact of HPV vaccination against cancer. We will address the evidence on the already observed impact from trials, routine vaccination programmes as well as modelling results focusing on the potential scale of the impact in terms of the current strategies.

#### Keynote lecture

#### Status, progress and challenges on global elimination of cervical cancer

A broad picture from the IARC/WHO on the progress towards elimination of cervical cancer across the globe, with emphasis on successes and challenges

P. Basu (France)

- Effectiveness of HPV vaccination: a Swedish perspective
- Head-to-head comparison of two HPV vaccines for sustainable immunogenicity and efficacy against CIN3+
- Cervical cancer elimination in Scotland effect of HPV immunisation with Cervarix on incidence of invasive cervical cancer
- Scientific basis for the WHO recommendation of a single dose of HPV vaccine
- What can modelling tell us about the expected timing of the effect of HPV vaccination on rates of invasive cervical cancer?
- Impact of HPV vaccination on inequalities in cervical cancer incidence in England
- Discussion and Q&A

J. Lei (Sweden)

M. Lehtinen (Finland)

T. J. Palmer (UK)

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P. Basu (France)

M. Saville (Australia)

P. Sasieni (UK)

J. Lei (Sweden)

& P. Sasieni (UK)

Coffee Break 9.30 • 10.00



Friday, February 10

Primary self-sampling strategies:

MSS 11 Experience and evidence

Room 0B 14.30 • 16.00

Chair: H. Berkhof (Netherlands) • J. Dillner (Sweden)

There is a growing interest in HPV testing on self-collected samples (HPV self-sampling). Studies have shown that women find HPV self-sampling more convenient, less embarrassing, less uncomfortable, and less painful than clinician-based sampling. Besides, HPV self-sampling is expected to increase the coverage of screening and can be maintained during the COVID-19 pandemic. Several countries have already implemented HPV self-sampling as a primary screening option and other countries are considering a more prominent role for self-sampling in their programs. In this session, we share experiences with (primary) self-sampling, how issues were solved, and present evidence on the performance of self-sampling in different settings.

Introduction: primary self-sampling in different settings	H. Berkhof (Netherlands) & J. Dillner (Sweden)
Self-sampling in Sweden	M. Elfström (Sweden)
Self-sampling in Denmark	J. Bonde (Denmark)
<ul> <li>Self-sampling in the Netherlands</li> </ul>	F. Inturrisi (Netherlands)
Self-sampling in the UK	K. Denton (UK)
Self-sampling in Australia	M. Saville (Australia)
Self-sampling in low-resource countries	S. De Sanjosé (Spain)
Discussion and Q&A	H. Berkhof (Netherlands) & J. Dillner (Sweden)

Coffee Break 16.00 • 16.30

Cottee Break		16.00 • 16.30
MSS 12	Extended HPV genotyping in screening Chair: J. Bonde (Denmark) • E. Franco (Canada)	Room 0B <b>16.30 • 18.00</b>
• Introductio	n	J. Bonde (Denmark) & E. Franco (Canada)
The FOCAL study: can HPV genotyping be part of HPV screening in a		G. Ogilvie (Canada)
Canadian setting? • Extended genotyping in a national screening program: an example from Sweden		J. Dillner (Sweden)
Genotyping in the AUS screening program: experiences and considerations		M. Saville (Australia)
• Extended g	enotyping and cytology as combined triage in HPV screening - experience	J. Bonde (Denmark)
• Discussion	and Q&A	J. Bonde (Denmark)
		& E. Franco (Canada)



Friday, February 10

**MSS 13** 

Next generation HPV based sequencing: what have we learned and what's next?

Room 0B 18.00 • 19.30

Chair: K. Cuschieri (UK) • L. Mirabello (USA)

HPV detection using rapid next generation sequencing (NGS) technologies can provide insight into the molecular biology, epidemiology and clinical trajectory of HPV infection. In this session we reflect on recent developments and applications of NGS and how this has furthered our understanding of HPV in the context of anogenital and head and neck disease.

• Introduction - general overview of the technology developments and pace of change including COVID legacy and general intro to session	K. Cuschieri (UK) & L. Mirabello (USA)
NGS for precise determination of viral integration in cervical cancer, what	S. Baulande (France)
have we learned?	
<ul> <li>Variation of HPV 16 in oropharyngeal cancer and implications for survival</li> </ul>	L. Mirabello (USA)
& prognosis	
<ul> <li>Cervical carcinogenesis through amplification of HPV episomes</li> </ul>	M. Dean (USA)
<ul> <li>HPV 16 variant distribution in anal samples in males: influence of underlying</li> </ul>	K. Cuschieri (UK)
disease status	
Discussion and Q&A	K. Cuschieri (UK)
	& L. Mirabello (USA)



#### Wednesday, February 8

SS 01

### **HPV** and pregnancy

Room 0E

Chair: K. Louvanto (Finland) • E. Siegler (Israel)

10.00 • 11.30

This session will focus on the most common concerns that women in childbearing age have on HPV infection and its consequences. The session will cover the current knowledge on the HPV role in reproductive life, during pregnancy and in children. We will also have a round table discussion on HSIL treatment options for pregnant women.

• Introduction	K. Louvanto (Finland)
HPV infection's role in children	K. Louvanto (Finland)
HPV role in infertility	G. Oron (Israel)
<ul> <li>HPV during pregnancy and risk of birth complications</li> </ul>	H. Trottier (Canada)
<ul> <li>Vertical transmission of HPV during perinatal period</li> </ul>	N. Suominen (Finland)
ROUND TABLE - HSIL management during pregnancy	
» LLETZ Treatment of CIN 2-3 in first 15 weeks of pregnancy	E. Siegler (Israel)
» Observation only management for CIN 2-3 during pregnancy	G. Haran (Israel)
» Discussion and Q&A	E. Siegler (Israel)



Wednesday, February 8

SS 02

# Criteria for HPV tests validation protocols for ASCUS triage and for post-treatment monitoring

Room 0D 10.00 • 11.30

Chair: K. Cuschieri (UK) • M. Poljak (Slovenia)

Protocols for clinical validation of HPV tests for primary cervical cancer screening indication are well established and widely accepted in HPV community, in contrast to other indications for HPV testing. The first part of the session will review past, present and future challenges of validation protocols for primary cervical cancer screening indication using clinician-taken cervical specimens, self-samples and urine samples. The second part of the session will review clinical rationale for the use of HPV testing for ASCUS triage indication followed by a description of proposed criteria for HPV tests validation protocols for this important indication. The third part of the session will review clinical rationale for use of HPV testing for detection of residual disease after local surgical treatment ("test of cure") followed by a description of proposed criteria for HPV tests validation protocols for post-treatment monitoring.

• Introduction	K. Cuschieri (UK)
	& M. Poljak (Slovenia)
Validation protocols for HPV tests for primary cervical cancer screening	M. Arbyn (Belgium)
using clinician-taken specimens: past, present, and future challenges	
<ul> <li>Validation protocols for collection devices and HPV tests for self-samples</li> </ul>	A. Vorsters (Belgium)
and urine samples	
<ul> <li>Rationale for use of HPV testing for ASCUS triage</li> </ul>	G. Valasoulis (Greece)
<ul> <li>Criteria for HPV tests validation protocols for ASCUS triage</li> </ul>	H. Pedersen (Denmark)
• Rationale for use of HPV testing for detection of residual disease after local	M. Kyrgiou (UK)
surgical treatment	& E. Paraskevaidis (Greece)
<ul> <li>Criteria for HPV tests validation protocols for post-treatment monitoring</li> </ul>	T. J. Palmer (UK)
Discussion and Q&A	K. Cuschieri (UK)
Discussion and Quit	& M. Poljak (Slovenia)
	& IVI. Fuljak (Sluvellia)

SS 03	Transmission: role of the male partner Chair: E. Franco (Canada) • M. Goodman (USA)	Room 0E <b>13.30 • 15.00</b>
Introduction		E. Franco (Canada) & M. Goodman (USA)
• Anal HPV inf	ection	J. Palefsky (USA)
• Heterosexua	al transmission of genital HPV infection: insights from	E. Franco (Canada)
epidemiolog	ical studies	
• Male partne	r	A. Giuliano (USA)
• The role of H	ILA polymorphisms in HPV transmission	K. Louvanto (Finland)
• Discussion a	nd Q&A	E. Franco (Canada)
		& M. Goodman (USA)



# Thursday, February 9

& C. Cocuzza (Italy)

SS 04	New screening algorithms in European countries Chair: M. Arbyn (Belgium) • P. Hillemanns (Germany)	Room 0E <b>8.00 • 9.30</b>
• Introduction	า	M. Arbyn (Belgium) & P. Hillemanns (Germany)
<ul><li>Results from</li><li>Policies for</li><li>Scenarios for</li><li>Self-sampling</li><li>Longitudina</li></ul>	ons of self-sampling In HaSCO trial Hannover Germany It cervical cancer screening and triage in Europe It or offering self-sampling: a meta-analysis of RCTs Ing as the primary approach in Italy It safety of HPV testing on self-samples	K. Vos (Netherlands) M. Jentschke (Germany) M. Arbyn (Belgium) S. Costa (Netherlands) P. Giorgi Rossi (Italy) G. Stanczuk (UK)
• Discussion a	and Q&A	M. Arbyn (Belgium) & P. Hillemanns (Germany)
SS 05	Molecular triage of HPV-positive women Chair: M. Arbyn (Belgium) • C. Cocuzza (Italy)	Room 0E <b>9.30 • 11.00</b>
• Introduction	า	M. Arbyn (Belgium) & C. Cocuzza (Italy)
screening p • High-risk Hi	PV genotyping in the risk stratification of HPV-positive women:	G. Stanczuk (UK) A. Del Mistro (Italy)
<ul> <li>Methylation</li> </ul>	xperience te of methylation analysis in the triage of HPV-positive women to biomarkers: from research to clinical practice dation of molecular triage biomarkers: preliminary results of the	B. Nedjai (UK) R. Steenbergen (Netherlands) L. De Marco (Italy)
VALTRIHP st		M. Arbyn (Belgium)



Thursday, February 9

SS 06 - fro

Evolution of HPV vaccination schedules - from evidence to public health impact

Chair: S. Franceschi (Italy) • M. Jit (UK)

Room 0E 14.00 • 15.30

The pace of introduction of HPV vaccine is stagnating, the coverage low in many countries and the WHO 2030 target of 90% coverage in danger. The present session will provide an overview of the evidence and considerations which led WHO to endorse the off-label use of a single-dose in girls included in HPV vaccination for routine and multi age cohort (MAC) catch-up. Ongoing efforts to accumulate additional information on the efficacy and durability of different HPV vaccine options will also be highlighted.

• Introduction S. Franceschi (Italy)

& M. Jit (UK)

• Trials of one dose schedules N. Mugo (Kenya)

• Observational study evidence about one dose schedules and its use in L. Markowitz (USA)

policy

Modelling of different dose schedules
 M. Brisson (Canada)

• Interpreting data from vaccination programs P. Sasieni (UK)

• Global recommendations, and their early impact P. Bloem (Switzerland)

• Discussion and Q&A S. Franceschi (Italy)

& M. Jit (UK)

Coffee Break 15.30 • 16.00



Thursday, February 9

SS 07

# Quality assurance and validation of primary, triage testing and collection methods

Chair: H. Berkhof (Netherlands) • K. Cuschieri (UK)

Room 0E 16.00 • 17.30

The implementation and evolution of cervical screening programs based on molecular HPV testing continues apace. To ensure tests and processes are fit for purpose before implementation - and in perpetuity - validation and quality monitoring of the end-to-end process is essential. This session will benefit from perspectives from laboratory experts, cancer epidemiologists and screening teams. Country specific approaches to quality processes will be discussed as will the perspectives of laboratory network(s). The challenges of validation given increased use of self-sampling and the greater emergence of immunised populations in screening will also be covered.

• Introduction	H. Berkhof (Netherlands) & K. Cuschieri (UK)
<ul> <li>European Ref Lab network update on quality guidance for HPV testing</li> <li>Pre-analytical optimisation of self-taken samples for HPV testing: recent</li> </ul>	M. Correa (Argentina) K. Vos (Netherlands)
developments and considerations for best practice  • Quality processes associated with HPV testing in Australia, a view from the laboratory	D. Hawkes (Australia)
<ul> <li>Validation of HPV self-sampling tests in real-world screening programs</li> <li>Longitudinal performance monitoring of HPV primary screening through</li> </ul>	H. Berkhof (Netherlands) C. Lagheden (Sweden)
<ul><li>audit</li><li>How to evaluate screening and triage tests in an era of vaccination</li></ul>	M. Rebolj (UK)
Discussion and Q&A	H. Berkhof (Netherlands) & K. Cuschieri (UK)



Thursday, February 9

**SS 08** 

#### Digital interventions to increase HPV vaccination

Room 0E

Chair: G. Woodall (USA) • G. Zimet (USA)

17.30 • 19.00

Introduction

G. Woodall & G. Zimet (USA)

#### Randomized trials of HPV vaccine uptake improvement: web apps for parents and young adolescent girls and boys

This presentation will discuss the results of two randomized trials of parent-focused web apps to improve HPV vaccine uptake for young adolescents (ages 11-14). For the first trial, the web app was tailored to parents and young adolescent girls, and in the second trial, the web app was tailored to parents and young adolescent boys. Results of both trials indicated significant web app impact on HPV vaccine uptake for adolescent girls and boys, as well as other vaccine uptake related variables. Discussion will include a consideration of web app content and tailoring to determine HPV vaccine uptake improvement.

G. Woodall

(USA)

#### Successful technology-based rural patient HPV vaccination reminder intervention and social media assessment of strategies to reduce HPV vaccine misinformation

This presentation will describe a multi-level and multi-component intervention that included healthcare team training activities and technology-based HPV vaccination reminders. Missed opportunities for HPV vaccination declined significantly from the pre-intervention to the post-intervention period. Participants who recalled receipt of an electronically delivered vaccination reminder had higher unadjusted odds of scheduling a visit compared with those who did not recall receiving a reminder. Social media-delivered misinformation related to HPV vaccination is pervasive. We will also discuss new strategies to evaluate and reduce the impact of HPV vaccine misinformation in rural settings.

D. Kepka

(USA)

#### Promoting HPV vaccination to emerging adults in rural communities in a multi-risk factor cancer prevention social media intervention

This presentation will describe an innovative social media campaign targeting six cancer risk factors, including HPV vaccination. It is being developed for the diverse population of adults aged 18-26 in rural counties in the Mountain West region of the U.S. Emerging adults obtain health information online far more than information from health care providers and other media. A framework for social media message development will be presented based on social cognitive, self-determination, and diffusion of innovation theory. Misinformation, especially on vaccination, will be combatted by instructing emerging adults in digital and media and by using an epidemiological model of monitoring and quickly responding to correct misinformation. The campaign will be tested with a sample of 1000 emerging adults in a stepped-wedge quasi-experimental design.

D. Buller (USA)

#### U.S. national digital point of care communication to improve uptake of HPV and other adolescent vaccines in clinic settings

This presentation will report on a study that involved digital targeted adolescent vaccination infographics and videos widely disseminated to clinical practices throughout the U.S. Over 11,000 clinicians whose practices received these digital interventions (exposed condition) were matched to an equal number of non-exposed comparison practices matched on multiple practice characteristics. The outcomes of interest were the number of vaccine doses (Tetanus-diphtheria-pertussis booster or Tdap, HPV, MenACWY, and MenB) administered to patients 11-18 years of age. The exposed clinics showed significant increases in administration of adolescent vaccines, including HPV vaccine, compared to the non-exposed clinics.

J. Klein

(USA)

#### • Discussion - points will include:

1. What digital intervention approaches are being used to promote cervical cancer screening?

G. Woodall & G. Zimet (USA)

2. What kind of digital interventions are being developed and evaluated in other countries, including low- and middle-income countries?



#### Thursday, February 9

SS 09

# Tumour HPV status and implications for survival outcomes in cervical and non-cervical disease

Room 0D 14.00 • 15.30

Chair: L. S. Arroyo Mühr (Sweden) • K. Cuschieri (UK)

While it is well established that HPV status is strongly associated with clinical outcomes in oropharyngeal cancer, increasing evidence suggests that this phenomenon is consistent in other tumour types, this is reflected in the recent WHO update on gynae cancer classification that advises cervical tumours should be referred to as "HPV independent" or "HPV associated". In this session, the basic molecular characteristics of HPV associated and HPV independent neoplasms will be discussed in addition to optimal and evolving methods to determine HPV status as well as the implications and challenges of using HPV status for clinical management and treatment decisions.

• Introduction	L. S. Arroyo Mühr (Sweden) & K. Cuschieri (UK)
The mutational landscape of HPV associated vs HPV negative cancer	I. Morgan (USA)
• Implications of HPV status on survival outcomes in cervical cancer	L. S. Arroyo Mühr (Sweden)
• Implications of HPV status on survival outcomes in anal cancer	K. Cuschieri (UK)
• Treatment de-intensification of oropharyngeal disease - navigating the data	P. Lassen (Denmark)
from trials	
<ul> <li>Methods for defining HPV status of tumours – do we need site-specific</li> </ul>	Q. Lepiller (France)
algorithms?	
Discussion and Q&A	L. S. Arroyo Mühr (Sweden)
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	a n. cuscineri (on)

Coffee Break 15.30 • 16.00



#### Thursday, February 9

SS 10	<b>Microbiome</b> Chair: A. Lev Sagie (Israel) • A. B. Moscicki (USA)	Room 0D <b>16.00 • 17.30</b>
• Introduction	on	A. Lev Sagie (Israel) & A. B. Moscicki (USA)
• The role of	the vaginal/cervical microbiome and HPV progression	A. B. Moscicki (USA)
• HPV-assoc	ated disease and anal microbiome	J. Palefsky (USA)
• Penile micr	robiome	A. Lev Sagie (Israel)
<ul> <li>Vaginal mid</li> </ul>	crobial transplantation	A. Lev Sagie (Israel)
• High-resol	ution microbiome profiling	M. Molina (Netherlands)
• Discussion	and Q&A	A. Lev Sagie (Israel)
		& A. B. Moscicki (USA)

SS 11

# HPV prevention and screening in vulnerable (immunosuppressed and transgender) populations

17.30 • 19.00

Room 0D

Chair: A. D'Souza (USA) • A. B. Moscicki (USA)

This session discusses HPV vaccination and screening for HPV-related cancers among vulnerable populations including transgender individuals and non-HIV immunosuppressed populations.

• Introduction	A. D'Souza (USA) & A. B. Moscicki (USA)
HPV vaccination of non-HIV immunocompromised patients	M. Goodman (USA)
Non-HIV immunosuppression: organ transplant and autoimmune disorders:	A. B. Moscicki (USA)
cancer risk and screening	
Cervical cancer prevention in transgender men	M. Kyrgiou (UK)
Gender neutral HPV testing in transgender population	E. Joura (Austria)
Discussion and Q&A	A. D'Souza (USA)
	& A. B. Moscicki (USA)



Friday, February 10

SS 12

Benefits and harms of HPV screening

Chair: C. Bouchard (Canada) • T. Malagon (Canada)

Room 0E

8.00 • 9.30

All screening programs entail harms as well as benefits. A core principle of modern screening programs and guidelines is that the overall benefit of the screening program should outweigh its potential harms. However, the estimation and the communication of the balance of harms and benefits can be in practice complex. In this session we shall review harms and benefits of cervical cancer screening in the context of populations highly HPV vaccinated, examine the balance of harms and benefits for future anal cancer screening, and finish with related issues arising when implementing HPV-based screening, including methods for estimating the balance of benefits and harms of screening algorithms, and developing communication tools on the benefits and harms of screening for shared decision making.

Introduction

C. Bouchard & T. Malagon

(Canada)

PART 1

Cervical cancer screening in the era of vaccinated populations

» Harms» Benefits

E. Franco (Canada)

N. Wentzensen (USA)

• PART 2

Anal cancer screening: is it time to initiate recommendations?

» Benefits» Harms

J. Palefsky (USA)

M. Clarke (USA)

• PART 3

Designing screening programs with benefits and harms in mind

» Estimating benefits and harms of screening algorithms

M. Saville (Australia)

» Communicating benefits and harms of screening to the public

S. Van Dijk (Netherlands)

Discussion and Q&A

C. Bouchard & T. Malagon

(Canada)

Coffee Break 9.30 • 10.00



Friday, February 10

SS 13 New triage methods

Room 0E

Chair: F. Carozzi (Italy) • B. Nedjai (UK)

10.00 • 11.30

Advanced molecular methods now allow better characterization of HPV infections detected by screening and enable optimization of cervical screening algorithms to differentiate women carrying HPV infection at lower or higher risk of precancerosis and cervical cancer.

In this session, we will discuss and explore issues related to their introduction into screening programs to enable increasingly accurate risk stratification.

• Introduction	F. Carozzi (Italy)
	& B. Nedjai (UK)
<ul> <li>Host and viral methylations as triage tool in HPV screening programs</li> </ul>	R. Adcock (USA)
• Genotyping	J. Dillner (Sweden)
<ul> <li>Comparing genotyping vs methylation as triage methods</li> </ul>	C. Meijer (Netherlands)
<ul> <li>DNA methylation testing on clinician and self -collected samples</li> </ul>	H. Berkhof (Netherlands)
<ul> <li>Comparison of triage strategies combining p16/ki67 or cytology with</li> </ul>	M. Benevolo (Italy)
extended genotyping in the Italian NTCC2 study	
<ul> <li>Dual stain and extended genotyping for triage of HPV-positive screening</li> </ul>	N. Wentzensen (USA)
results	
• Italian guidelines for the use of biomarkers as triage in HPV screening	P. Giorgi Rossi (Italy)
Discussion and Q&A	F. Carozzi (Italy)
	& B. Nedjai (UK)



Friday, February 10

SS 14

# Screening and vaccination implementation in Eastern and Central Europe - a part of Europe with the highest burden of cervical cancer

Room 0D

10.00 • 11.30

Chair: H. Berkhof (Netherlands) • M. Poljak (Slovenia)

Burden of cervical cancer in Eastern and Central Europe is unproportionally higher than in other parts of Europe, both in terms of incidence and mortality. In the session latest available epidemiological data concerning burden of cervical cancer in the region will be presented, followed by reviews of cervical cancer screening practices and implementation, as well as HPV vaccine implementation in Eastern and Central Europe. In addition, implementation status of national organized HPV-based cervical cancer screening of two pioneering countries of HPV-based cervical cancer screening in the region (Albania and Montenegro) will be presented, followed by a presentation on HPV vaccine implementation in Hungary – a country with the highest HPV coverage rate in the region.

• Introduction	H. Berkhof (Netherlands) & M. Poljak (Slovenia)
<ul> <li>Burden of cervical cancer in Eastern and Central Europe</li> <li>Cervical cancer screening practices and implementation in Eastern and Central Europe</li> </ul>	D. Singh (France) B. Serrano (Spain)
<ul> <li>HPV vaccine implementation in Eastern and Central Europe</li> <li>Implementation status of national organized HPV-based cervical cancer screening in Albania</li> </ul>	L. Bruni (Spain) K. Filipi (Albania)
Implementation status of national organized HPV-based cervical cancer screening in Montenegro	I. Samardžić (Montenegro)
HPV vaccine implementation in Hungary	A. Molnár (Hungary)
Discussion and Q&A	H. Berkhof (Netherlands) & M. Poljak (Slovenia)

Lunch Break 11.30 • 13.15

SS 15

# The role of obesity in cervical cancer screening and management

Room 0B

13.15 • 14.15

Chair: M. Clarke (USA)

Obesity is a worldwide public health challenge, increasing the risk for several cancers including cervical cancer. Prior research has shown that individuals with obesity have a higher risk of cervical cancer, but a lower risk of precancer compared to those with normal BMI, which is thought to be at least in part explained by missed detection of precancers. The current session addresses the impact of obesity on the effectiveness of cervical

cancer screening and management.IntroductionM. Clarke (USA)

• The role of obesity in cervical cancer screening and cancer: results of a nationwide Danish cohort study

A. Urbute (Denmark)

• The impact of obesity on management of cervical precancers

M. Clarke (USA)

Assessing the impact of obesity on risk-based management guidelines

D. Egemen (USA)

Discussion and Q&A

M. Clarke (USA)



Friday, February 10

SS 16

Who should be referred for high-resolution anoscopy (HRA)?
And what if HRA is not available?

Room 0E

14.30 • 16.00

Chair: L. Abramowitz (France) • A. Nyitray (USA)

Biomarkers are under development to support decision-making about who should be referred to HRA. This session will first provide a state of the science overview about biomarkers to support anal precancer screening programs. However, there will be jurisdictions that will not have enough skilled anoscopists for many years, if ever. Clinicians in these areas cannot use biomarkers for precancers if positive results cannot be followed up with HRA. What are screening the options for these clinicians who want to detect an anal malignancy as early as possible?

• Introduction	L. Abramowitz (France)
	& A. Nyitray (USA)
Global burden of anal cancer according to sex and HIV status	A. Deshmukh (USA)
Biomarker state of the science	M. Clarke (USA)
<ul> <li>Anal smears methylation as a risk factor for progression to SCCA</li> </ul>	M. V. Ferré (France)
<ul> <li>Support for clinical decision-making in the absence of HRA</li> </ul>	L. Abramowitz (France)
<ul> <li>ANCHOR data on the risk of anal cancer after benign anal disease</li> </ul>	J. Palefsky (USA)
Discussion and Q&A	L. Abramowitz (France)
	& A. Nyitray (USA)

Coffee Break 16.00 • 16.30

SS 17	HPV latency Chair: A. B. Moscicki (USA)	Room 0E <b>16.30 • 18.00</b>
• Introduction	า	A. B. Moscicki (USA)
Latency: basic science perspective		J. Doorbar (UK)
• Latency of anal HPV: does it exist?		A. Nyitray (USA)
• Latency in v	vomen: here today, gone (for sure) tomorrow?	A. B. Moscicki (USA)
• Latency in t	he oral cavity: what we don't know and what we need to know	A. D'Souza (USA)
• Latency in t	he genitals of men: it must be there but where?	A. Giuliano (USA)
• Discussion	and Q&A	A. B. Moscicki (USA)



Friday, February 10

# Risk stratification in the follow up

Room 0E

Chair: F. Carozzi (Italy) • N. Wentzensen (USA)

18.00 • 19.30

Women treated for high grade lesions and women HPV positive/colposcopy negative represent an at-risk population compared to the routinely screened population. So, a combination of well-known risk factors associated to new molecular markers could allow a better risk stratification of these women. Balancing the benefits of robust follow-up with the harms of over-scrutiny is particularly challenging. In this session we will review the international experience from real-life program and reflecting on prevailing knowledge "gaps". We will consider modalities that may improve risk stratification and thereby optimize management in the future.

<ul> <li>An overview of tests for postcolposcopy and posttreatment surveillance</li> <li>Accuracy of the margin status of excised tissue vs post-therapeutic HPV testing to predict failure of excisional treatment of cervical pre-cancer</li> <li>HPV testing as a test of cure: experiences from applied research and service provision in Scotland</li> <li>Utility and value of HPV vaccine in the colposcopy population and after treatment</li> <li>Risk of cancer after conization</li> <li>Women with a positive high-risk human papillomavirus (HPV) test remain at increased risk of HPV infection and cervical precancer ≥15 years later</li> <li>Classification of high-grade cervical intraepithelial neoplasia by p16ink4a, Ki-67, HPV E4 and FAM19A4/miR124-2 methylation status demonstrates</li> </ul>
<ul> <li>Accuracy of the margin status of excised tissue vs post-therapeutic HPV testing to predict failure of excisional treatment of cervical pre-cancer</li> <li>HPV testing as a test of cure: experiences from applied research and service provision in Scotland</li> <li>Utility and value of HPV vaccine in the colposcopy population and after treatment</li> <li>Risk of cancer after conization</li> <li>Women with a positive high-risk human papillomavirus (HPV) test remain at increased risk of HPV infection and cervical precancer ≥15 years later</li> <li>Classification of high-grade cervical intraepithelial neoplasia by p16ink4a,</li> <li>C. Meijer (Netherlands)</li> </ul>
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<ul> <li>HPV testing as a test of cure: experiences from applied research and service provision in Scotland</li> <li>Utility and value of HPV vaccine in the colposcopy population and after treatment</li> <li>Risk of cancer after conization</li> <li>Women with a positive high-risk human papillomavirus (HPV) test remain at increased risk of HPV infection and cervical precancer ≥15 years later</li> <li>Classification of high-grade cervical intraepithelial neoplasia by p16ink4a,</li> <li>C. Meijer (Netherlands)</li> </ul>
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treatment  • Risk of cancer after conization  • Women with a positive high-risk human papillomavirus (HPV) test remain at increased risk of HPV infection and cervical precancer ≥15 years later  • Classification of high-grade cervical intraepithelial neoplasia by p16ink4a,  C. Meijer (Netherlands)
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• Classification of high-grade cervical intraepithelial neoplasia by p16ink4a, C. Meijer (Netherlands)
Ki-67, HPV E4 and FAM19A4/miR124-2 methylation status demonstrates
considerable heterogeneity with potential consequences for management
• Optimal management of older women treated for CIN: balancing the risks J. Wang (Sweden)
and benefits
• Effect of condom use after CIN treatment on cervical HPV biomarkers E. Paraskevaidis (Greece)
positivity: prolonged follow up study
• Discussion and Q&A F. Carozzi (Italy)
& N. Wentzensen (USA)



Friday, February 10

# The utility of urine for improved cervical cancer SS 19 prevention

Room 0A 16.30 • 18.00

Chair: R. Steenbergen (Netherlands) • S. Van Keer (Belgium)

Urine samples offer several advantages over conventional cervical scrapes and self-collected cervicovaginal samples for cervical cancer prevention. One of the most important advantages being the ease of collection and the wide acceptance by women. The number of studies supporting the use of urine for HPV testing is rising rapidly. Studies on testing for other biomarkers in urine are just evolving. This session will discuss current developments on the analysis of HPV and other biomarkers (including methylation) for the detection of cervical lesions in urine, and the potential of vaccine monitoring using urine as liquid biopsy to detect HPV induced antibodies.

• Introduction	R. Steenbergen (Netherlands) & S. Van Keer (Belgium)
• Urine sampling for cervical cancer screening – results from the German	M. Jentschke (Germany)
CoCoss and HaSCo trials	
<ul> <li>Urine biomarkers in cancer detection: a systematic review</li> </ul>	S. Jordaens (Belgium)
<ul> <li>Follow-up of cervical cancer patients using urine biomarkers</li> </ul>	N. Van Trommel (Netherlands)
• Endometrial cancer detection in patient-friendly sample types by DNA	R. Steenbergen (Netherlands)
methylation analysis	
• A novel cervical cancer screen-triage-treat demonstration project with HPV	L. Chinula (Malawi)
self-testing and thermal ablation for women in Malawi: protocol for a single-	
arm prospective trial	
<ul> <li>Urine for the monitoring of vaccinated women: 3.5-year persistence of</li> </ul>	A. Vorsters (Belgium)
immune responses in paired urine and serum samples	
Discussion and Q&A	R. Steenbergen (Netherlands)
	& S. Van Keer (Belgium)



# **CS - CLINICAL SESSIONS**

Room 0E <b>15.30 • 16.45</b>	Management of CIN Chair: E. Paraskevaidis (Greece)	CS 01
E. Paraskevaidis (Greece)	on	• Introductio
A. Athanasiou (UK)	tive and oncological outcomes after treatment for CIN: ons between different methods	· ·
P. Martin-Hirsch (UK)	cal treatments, when, to whom and associated morbidity	• Repeat loca
B. Strander (Sweden) E. Paraskevaidis (Greece)	risk of cervical cancer post treatment and possible explanations red management of women with cervical pathology incorporating gorithms	
E. Paraskevaidis (Greece)	n and Q&A	• Discussion
Room 0E <b>16.45 • 18.15</b>	Test of cure: strategies for the follow up of women treated for cervical intraepithelial neoplasia Chair: C. Cocuzza (Italy) • K. Cuschieri (UK)	CS 02
C. Cocuzza (Italy)	on	• Introductio
& K. Cuschieri (UK)		marodactio
& K. Cuschieri (UK)  A. Heinonen (Finland)	the key issues facing colposcopists in the post-treatment ent of women - where are the research gaps?	• What are th
	the key issues facing colposcopists in the post-treatment ent of women - where are the research gaps?  Sysis of the accuracy of markers to predict failure or cure after of cervical precancer	<ul><li>What are the manageme</li><li>Meta-analy</li></ul>
A. Heinonen (Finland)	ent of women - where are the research gaps?  ysis of the accuracy of markers to predict failure or cure after	<ul> <li>What are the manageme</li> <li>Meta-analy treatment of Longitudina</li> </ul>
A. Heinonen (Finland)  M. Arbyn (Belgium)	ent of women - where are the research gaps?  lysis of the accuracy of markers to predict failure or cure after  of cervical precancer  nal performance of test of cure in 40,000 women in Scotland;	<ul> <li>What are the manageme</li> <li>Meta-analy treatment of Longitudinal key learning</li> </ul>
A. Heinonen (Finland)  M. Arbyn (Belgium)  T. J. Palmer (UK)	ent of women - where are the research gaps?  lysis of the accuracy of markers to predict failure or cure after  of cervical precancer  nal performance of test of cure in 40,000 women in Scotland;  ng points and future developments	<ul> <li>What are the manageme</li> <li>Meta-analy treatment of Longitudinal key learning</li> <li>Use of HPV patients</li> <li>Circulating</li> </ul>
A. Heinonen (Finland)  M. Arbyn (Belgium)  T. J. Palmer (UK)  M. Martinelli (Italy)	ent of women - where are the research gaps?  Lysis of the accuracy of markers to predict failure or cure after  to of cervical precancer  hal performance of test of cure in 40,000 women in Scotland;  hag points and future developments  Vigenotyping and viral load for the risk stratification of test of cure  B DNA: a predictor of survival and outcomes after treatment for  ve cervical disease and cancer?	<ul> <li>What are the manageme</li> <li>Meta-analy treatment of Longitudinal key learning</li> <li>Use of HPV patients</li> <li>Circulating</li> </ul>



#### **CS - CLINICAL SESSIONS**

#### Thursday, February 9

**CS 03** 

#### Colposcopy in the HPV era

Chair: C. Bouchard (Canada) • P. Hillemanns (Germany)

Room 0D

9.30 • 11.00

Colposcopy is viewed as the triage gold standard in cervical cancer screening. However, it is challenged by easy-to-perform biomarkers and new AI technology amidst declining CIN prevalence in HPV-vaccinated populations.

- Introduction
- Colposcopy in the era of triage markers
- Colposcopy a challenge for AI
- Pitfalls of vulvar colposcopy
- Future of colposcopy in populations highly HPV vaccinated
- New challenges in colposcopy teaching and maintenance of competency in the world of comprehensive HPV guidelines
- Discussion and Q&A

C. Bouchard (Canada)

& P. Hillemanns (Germany)

C. Bergeron (France)

M. S. Kalteis (Germany)

M. Jentschke (Germany)

M. Brisson (Canada)

P. Nieminen (Finland)

C. Bouchard (Canada)

& P. Hillemanns (Germany)



# **CS - CLINICAL SESSIONS**

Friday, February 10

CS 04	Vaginal Intraepithelial Neoplasia (VaIN) management Chair: M. Hampl (Germany) • M. Preti (Italy)	Room 0C 8.00 • 9.30
• Introduction	on	M. Hampl (Germany) & M. Preti (Italy)
<ul><li> Vaginal mid</li><li> Vaginal col</li><li> Risk factors</li></ul>	ogy and HPV typing of Vaginal Intraepithelial Neoplasia crobiota and vaginal HPV infection poscopic assessment: are scores and classifications useful? s analysis of persistence, progression, and recurrence in Vaginal lial Neoplasia	L. Bruni (Spain) P. Vieira Baptista (Portugal) J. Bornstein (Israel) M. Bradbury (Spain)
Treatment     Discussion	of Vaginal Intraepithelial Neoplasia and Q&A	M. Hampl (Germany) M. Hampl (Germany) & M. Preti (Italy)
Coffee Bred	ak	9.30 • 10.00
CS 05	Vulvar Intraepithelial Neoplasia (VulN) update Chair: M. Bleeker (Netherlands) • J. Bornstein (Israel)	Room 0C 10.00 • 11.30
• Introduction	on	M. Bleeker (Netherlands) & J. Bornstein (Israel)
<ul><li>The termin</li><li>The signific</li><li>Lichen scle and vulvar</li><li>Treatment</li></ul>	of VIN – imiquimod or surgery	J. Bornstein (Israel) S. Regauer (Austria) M. Bleeker (Netherlands) M. Preti (Italy) G. Trutnovsky (Austria)
Discussion	and Q&A	M. Bleeker (Netherlands) & J. Bornstein (Israel)



FC 01	Genotyping Chair: J. Andrews (USA)	Room 0A 10.00 • 11.30
	ping in biopsies of HSIL and invasive cervical cancers in women IIV: a cohort- and a nested-case control study	C. Gilles (Belgium)
after HPV p	int HPV 33 / HPV 58 probe on referral rates and clinical outcome ositive screening sample using Onclarity™ HPV assay	N. Trine Dahl (Denmark)
	evidence using the Onclarity™ extended HPV genotyping assay in	L. Vaughan (USA)
II HPV28 as:	n between the novel ALLPLEXTM II HPV28 assay, the ANYPLEXTM say and INNO-LIPA HPV genotyping extra II assay for HPV and genotyping	M. Bell (Belgium)
	enotyping, cytology, and self-sampling: risk-based illustrations	J. Andrews (USA)
• Evaluation o	of the Onclarity™ HPV assay for HPV-DNA detection in cervico- ples from the Italian NTCC2 study	L. De Marco (Italy)
•	n of Anyplex™ II HPV28 detection and linear array genotyping for estimation in epidemiological studies	F. Moreno (Brazil)
•	PV mRNA types 16, 18, 45 detection on the risk of CIN3+ in young n normal cervical cytology	S. W. Sorbye (Norway)
• Validation o HPV genoty	f intra- and inter-laboratory reproducibility of the RIATOL qPCR ping assay	D. Vanden Broeck (Belgium)



FC 02	Anal neoplasia Chair: L. Abramowitz (France) • J. Palefsky (USA)	Room 0D 12.00 • 13.30
• Perianal wa	orts in HIV-infected MSM: harbingers of precancer and cancer	M. Gaisa (USA)
-	gy and human papillomavirus genotyping of a population at isk of developing anal dysplasia and cancer	C. Sani (Italy)
• High preva	ence of HPV, other STI and anal lesions among MSM in Togo	C. Charpentier (France)
among me	of HPV genotypes in self- versus clinician-collected anal swabs n who have sex with men (MSM) in Milwaukee, Wisconsin, USA: t anal cancer self-swab study	J. Nitkowski (USA)
	IPV, mRNA HPV and p16 tests for diagnosis of precursor lesions incer: systematic review and meta-analysis	A. C. Macedo (Brazil)
is very low	Rectal Examination (DARE) utilization for anal cancer screening among those most vulnerable to anal cancer: the prevent anal swab study	A. Nyitray (USA)
	coriolus versicolor-based vaginal gel and a REISHI-based food t for the treatment of high-risk HPV associated lesions: a case	P. Sanmartín (Spain)



FC 03	Self-sampling 1 Chair: F. Inturrisi (Netherlands) • P. Sasieni (UK)	Room 0D <b>13.30 • 15.00</b>
	npling among long-term non-attenders to cervical cancer Norway: a pragmatic randomized controlled trial	A. Tropé (Norway)
• Self-samplin Sweden	g within routine cervical cancer screening in region of Skåne,	O. Forslund (Sweden)
•	f-sampling method for HPV-DNA testing in an organized cervical rogram: an Italian experience	A. Chiereghin (Italy)
• Cytological t	esting on cervical versus vaginal self-samples	A. Latsuzbaia (Luxembourg)
	7-type HPV mRNA E6/E7 testing in self-collected samples as normal cytology results: a Mexican multicentric study	C. E. Aranda Flores (Mexico)
	ation test for detection of cervical pre-cancer and cancer in self- rvicovaginal specimens	D. Scibior-Bentkowsk (UK)
• Evaluation o	of alternative suspension media for vaginal self-collected samples	C. Giubbi (Italy)
	linical validation of self-collection using Copan FLOQSwabs™ and -Brush® eluted in Copan MSwab® media	D. Hawkes (Australia)
• The potentia	al of first-void urine as a source of high-quality DNA for cervical ening and triage	E. Van Den Borst (Belgium)
• HPV self san	npling-breaking barriers in cervical cancer screening	L. Balasubramani (India)
	A sequencing of vaginal self-samples versus cervical smears	W. Leenders (Netherlands)



FC 04	<b>Epidemiology and natural history 1</b> Chair: M. Goodman (USA)	Room 0A 12.00 • 13.30
sectional cl	h cervical high-risk human papillomavirus: the ANGY cross- inical study	M. Jacot-Guillarmod (Switzerland)
incidence o	oillomavirus prevalence in the European country with a high f cervical cancer	N. Berza (Latvia)
prevalence	sk human papillomavirus infection in Eastern Ethiopia: cytological profile and associated factors	A. S. Hassen (Ethiopia)
women in t	V persistence from a large screening cohort of HPV positive he district of Florence	C. Sani (Italy)
coverage (E	e of HPV 16/18 infection in a population with high vaccination ngland): findings, issues and future priorities	A. Anderson (UK)
cervical car	recurrence risk model for tailored surveillance strategy in acer patients	L. Dostalek (Czech Republic)
cytology: a	between sexually transmitted infections and abnormal cervical prospective study based on cervical cancer screening cohort	S. Chen (China)
	of the microenvironment in HPV associated lesions of the cervix-a study of the squamous and glandular neoplasia	M. Farcas (Romania)
	behavioural and biological factors in the association between HIV and cervical cancer incidence in South Africa: a mathematical tudy	D. De Bondt (Netherlands)



FC 05	HPV prophylactic vaccines l Chair: P. Bonanni (Italy)	Room 0A <b>13.30 • 15.00</b>
• The NCI-ACI	B one dose vaccine studies in Costa Rica: a status report	R. Herrero (Costa Rica)
• • • • • • • • • • • • • • • • • • • •	••••••••••••	
_	effectiveness of the 9-valent human papillomavirus (9vHPV) candinavian countries	A. Luxembourg (USA)
• • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	
	illomavirus (HPV) immunization and screening in the Italian overview of the current status of operation	P. Bonanni (Italy)
• • • • • • • • • •	•••••••••••••	
	c genital HPV-infection rates among south African women 5-10 school-based vaccination	G. Dreyer (South Africa)
• • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	
	h impact of 2-valent, 4-valent, and 9-valent HPV vaccination us coverage scenarios in China - a simulation study	I. Diakite (USA)
• • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	
	e implications of a single dose HPV vaccine regimen in a low/ me country setting: an analysis in Indonesia	V. Daniels (USA)
• • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	
-	on of human papillomavirus vaccination and series completion mid-adolescence	E. Goodman (USA)
• • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	
	lesions after hysterectomy for high-grade cervical intraepithelial nd early-stage cervical cancer: a focus on the potential role of	G. Bogani (Italy)
• • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	
	illomavirus vaccination delivery systems within national and munization programs: a systematic literature review	M. Flesher (USA)
• • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	
• HPV Mallow	Teens project - COOLTAS comics for young generation	N. T. Ildiko (Hungary)
	ation L2-Based HPV vaccines cross-protect against cutaneous rus infection and tumor development	D. Hasche (Germany)



FC 06	Methylation 1 Chair: S. Van Keer (Belgium) • B. Wisman (Netherlands)	Room 0A <b>15.30 • 17.00</b>
	enotyping and methylation triage facilitate a purely molecular of young women 23-29 years old?	H. Pedersen (Denmark)
• ScreenYu G	iyn® - cervical cancer screening triage based on a single DNA n marker	A. Hansel (Germany)
QIAsure DI	costs and diagnostic outcomes of replacing cytology with the NA methylation assay as a triage within HPV primary cervical eening in the Netherlands	S. Huntington (UK)
	logical classification to a continuous score: using epigenetic data eterogeneity of cervical lesions	M. Bonjour (France)
-	a novel method for genome-wide DNA methylation detection, can characterize different gynaecological cancers and associated HPV	J. Boers (Netherlands)
•	of-principle of marker discovery for different gynaecological a novel method for genome-wide DNA methylation profiling	R. Boers (Netherlands)
•	n of biomarkers in Multizonal Intraepithelial Neoplasia: ding Epithelial Transformation (MINUET)	E. Sumiec (UK)
FC 07	New technologies / Artificial intelligence Chair: G. Donders (Belgium) • P. Nieminen (Finland)	Room 0C <b>13.30 • 15.00</b>
	omation, data science and artificial intelligence-based analysis to curacy and throughput of a quantitative HPV-genotyping PCR	R. Pereira (Belgium)
	development and validation of an open-source software-based or 3D reconstruction of organotypic culture models	M. S. Kalteis (Germany)
• Colposcopy of Uzbekist	y based on artificial intelligence: experience in the Republic an	N. Zakhirova (Uzbekistan)
	e of electrical impedance spectroscopy in adjunction of a prospective cohort study	P. L. Omar (Finland)



Room 0C <b>15.30 • 17.30</b>	Colposcopy / Management Chair: M. Hampl (Germany) • E. Siegler (Israel)
G. Fachetti-Machado (Brazil)	nent among colposcopists on the identification of three images more ntly seen in glandular cervical precursor neoplasia
• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •
E. Alfonzo (Sweden)	cy of colposcopy in the Swedish screening programme
• • • • • • • • • • • • • • • • • • • •	
J. Bouda (Czech Republic)	ement and stratification of patients with AGC-FN PAP smear
	• • • • • • • • • • • • • • • • • • • •
L. Bergqvist (Finland)	surveillance of CIN2 in young women - a prospective cohort study
• • • • • • • • • • • • • • • • • • • •	
M. Bradbury (Spain)	perative human papillomavirus test as an early marker of cervical rade squamous intraepithelial lesion treatment failure
• • • • • • • • • • • • • • • • • • • •	
K. Madziar (Poland)	al application of modified ASCCP 2019 algorithms in the diagnosis and etection of cervical pathology
• • • • • • • • • • • • • • • • • • • •	
N. Thuijs (Netherlands)	ehensive characterization of 751 vulvar lesions, originally diagnosed -grade vulvar intra-epithelial neoplasia
• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •
A. M. Kaufmann (Germany)	al smear evaluation by RNA-based quantigene-molecular-profiling- gy assay reports diagnosis and prognosis for lesion severity and pment
• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •
G. E. Cetera (Italy)	s on my side" disease trajectory of vulvodynia: a systematic review narrative synthesis
• • • • • • • • • • • • • • • • • • • •	
Y. Li (China)	ng the value of focused ultrasound ablation therapy for fertility tion in women of childbearing age with cervical lesions: Protocol of a ctive cohort study
V. Boero (Italy)	erm efficacy of fat grafting in vulvar Lichen Sclerosus: an ational retrospective study
V. Boero (Italy)	erm efficacy of fat grafting in vulvar Lichen Sclerosus: an



# Thursday, February 9

FC 09	Self-sampling 2 Chair: C. Cocuzza (Italy) • G. Ogilvie (Canada)	Room 0D <b>8.00 • 9.30</b>
-	nd diagnostic performance of HPV self-samples compared to lected screening samples	B. Sonne (Denmark)
	f MSwab® medium for the elution of self-collected FLOQSwabs® papillomavirus (HPV) detection on six commercial PCR-based	S. Castriciano (Italy)
_	-collected vs cervical clinicians collected samples for cervical ening in COVID-19 era	A. lossa (Italy)
	on for cervix screening in the British Columbia organized cervix opulation-based program: preliminary program findings	G. Ogilvie (Canada)
·	roximity-incentive strategy based on vaginal self-sampling for cer screening in the French departments of Aude and Hérault	N. Boulle (France)
•	ysician-collected high-risk HPV mRNA testing to detect high- cal lesions among Thai women visiting a colposcopy clinic	N. Phoolcharoen (Thailand)
	arriers to clinic-based cervical cancer screening and motivators -sampling during the COVID-19 pandemic	J. Montealegre (USA)
• Validation o	of the clinical performance and reproducibility of the NEUMODX self-sample workflow	B. Hesselink (Netherlands)
		K. Zheng (USA)
	of new strategies of storage and extraction for the ation of self-collection in cervical screening programs	l. Sechi (ltaly)



# Thursday, February 9

FC 10	Low income countries 1 Chair: O. Gassama (Senegal) • J. Smith (USA)	Room 0A <b>8.00 • 9.30</b>
• Barriers to	cervical cancer prevention and control in Guatemala	M. Dean-Smith (USA)
• Low levels in Guatema	of lifetime PAP test receipt among vulnerable patient populations ala	K. Garcia (USA)
advanced s	area-level socioeconomic status and health care access with stage at diagnosis by race/ethnicity among women with cervical ne USA: a case for improving cervical cancer screening access to ulnerable	J. Islam (USA)
	e of conventional PAP smear as a screening tool for cervical ow socioeconomic countries	G. Gunasekaran (India)
	g HPV self-testing in 4 resource-constrained countries to enhance rvical cancer screening programs: a preliminary field evaluation	J. Beltman (Netherlands)
	king cervical cancer care at a tertiary oncology facility in Uganda ional treatment targets on cervical cancer control	M. De Fouw (Netherlands)
• The Emerg	ing Technologies in Cervical Cancer Screening 'ETiCCS' initiative	M. S. Kalteis (Germany)



Thursday, February 9

FC 11	Low income countries 2 Chair: O. Gassama (Senegal) • J. Smith (USA)	Room 0A <b>9.30 • 11.00</b>
	d facilitators to cervical cancer screening among under-screened Cuenca, Ecuador: the perspectives of women and health als	B. Verberckmoes (Belgium)
	ntred design of an HPV rapid test for cervical cancer screening in ed populations	N. Rodriguez (USA)
	by thermoablation of pre-cancerous lesions of the uterine cervix cal regions of Tambacounda and Kedougou (Senegal)	O. Gassama (Senegal)
-	s: learnings from HPV-based cervical cancer self-screening s in remote indigenous communities	L. Smith (Canada)
•	ty, feasibility, and appropriateness of integrating HPV self- or cervical cancer screening into voluntary family planning Malawi	P. Mhango (Malawi)
areas from	-sampling for the diagnosis of human papillomavirus in rural Cuenca Ecuador: acceptance, sensitivity and specificity among lling, self-sampling and clinician sampling	B. Vega (Ecuador)



FC 12	HPV prophylactic vaccines 2 Chair: E. Joura (Austria)	Room 0D <b>8.00 • 10.00</b>
_	efficacy, immunogenicity, and safety of the quadrivalent and PV vaccines: an overview of clinical trial long-term follow-up	A. Luxembourg (USA)
	utral HPV vaccination policies in the European Union and the UK: equity, the role of LGBT permissive societies and advocacy	K. Verbeke (UK)
• Safety and idiopathic a	immunogenicity of Gardasil4® in females age 9-26 with juvenile arthritis	N. Singer (USA)
	nent use of prophylactic HPV vaccines in patients with HPV-disease: review of the mechanism of action	M. Reuschenbach (Germany)
-	PV vaccination in patients treated for vulvar HSIL, a randomised ntrolled trial	R. Van de Laar (Netherlands)
Association initiation	of child's age, parental HPV vaccine hesitancy, and HPV vaccine	M. Silver (USA)
	e prescription and compliance in a cohort of women undergoing of the transformation zone	M. Santos (Portugal)
	ation changes over time in young population in Brazil: results	E. Wendland (Brazil)
• Sociodemo eligible USA	graphic barriers to human papillomavirus vaccination among A veterans	J. Zevallos (USA)
	and uptake of HPV vaccination and cervical cancer screening: v data from an Italian region	A. Acuti Martellucci (Italy)
• After-coniza	ation HPV vaccination uptake: real-world data from Puglia (Italy)	P. Berardi (Italy)
• HPV vaccina analysis	ation uptake in men who have sex with men in Europe: a meta-	A. Di Lorenzo (Italy)



FC 13	Screening / HPV testing 1 Chair: J. Bogaards (Netherlands) • S. Van Dijk (Netherlands)	Room 0C 14.30 • 16.00
	cypical glandular cells lesions in cervical cancer screening: erformances, correlation with HPV test and histology	I. Paganini (Italy)
	of patients with histological diagnoses in the follow-up after two -testing HPV and cytology in Germany	A. Xhaja (Germany)
	n of primary HPV programs in Russia as a means to improve ervical screening	I. Olkov (Russia)
• Compariso Columbia's	n of long-term colposcopy referrals by age groups in British cervix screening program among those who did or did not IPV-based screening in the HPV-FOCAL trial	G. Ogilvie (Canada)
	al colposcopy referral in HPV screening using combined genotype gy triage compared to LBC-screening with HPV triage	J. Bay (Denmark)
• HPV primai region of N	ry screening with extended genotyping in the south-east health orway	A. K. Lie (Norway)
• • • • • • • • • • • • • • • • • • • •	quality assurance program based on re-analysis of "HPV negative"	C. Lagheden (Sweden)
FC 14	Epidemiology and natural history 2	Room 0C
1014	Chair: S. Franceschi (Italy) • S. Nygård (Norway)	16.30 • 18.00
_	cervical cancer screening patterns among women living with HIV using a population-based sample: an analysis of the NIH's All of	J. Islam (USA)
	dence decrease in Norway: observational registry-based cohort 2004 to 2020	S. Nygård (Norway)
• P16 expres a retrospec	sion in invasive adenocarcinomas of Mozambican patients: tive study	L. Lovane Matias (Mozambique)
• Compariso socioecono	n of cervical and anal cancer incidence trends by race and mic status in the USA, 2006-2018	A. Mazul (USA)
	of HPV-independent cervical cancer: an increasingly important city	Z-Y. NG (Singapore)
•	n and oncogenic potential of HPV infections in cohort study s in Vojvodina, North region of Serbia	A. Mandic (Serbia)
• The impact intraepithe	of diet and sleep disorders on the degree of cervical lial lesions	B. Jugeli (Gorgia)



FC 15	Economics and modelling Chair: M. Brisson (Canada) • M. Drolet (Canada)	Room 0C 18.00 • 19.30
• Economic b literature re	urden of cervical cancer in mainland China: a systematic eview	C. Velicer (USA)
cervical can	veness of computer-assisted cytology in a primary HRHPV-based cer screening programme	E. Olthof (Netherlands)
Cost-effecti genotyping	veness of HPV-based cervical screening using HPV16/18 and cytology as triage test: a modelling study	K. Castañeda (Netherlands)
	ation and model validation for historical cervical cancer screening	C. Palmer (USA)
• Prevalence- individual ri	incidence model for the cumulative risk of CIN2+ based on sk-factors	K. Kroon (Netherlands)
	pact of one-dose HPV vaccination in Low-and-middle-income- MIC): a modeling analysis using HPV-ADVISE LMIC	M. Drolet (Canada)
	economic effects of nonavalent versus bivalent HPV vaccination erlands: a data-driven analysis	B. Sollie (Netherlands)
	for conducting rapid assessments of the impact of interventions way towards global cervical cancer elimination	R. Stuart (USA)



Room 0A 18.00 • 19.30	C 16  Health education – Advocacy – Public Health Chair: N. Osazuwa-Peters (USA) • J. Paavonen (Finland)	FC 16
K. Saxena (USA)	Attitudes towards human papillomavirus vaccination among adults in the USA	• Attitude USA
D. Kepka (USA)	Higher levels of HPV vaccine hesitancy among rural hispanic young adults in the western U.S., 2020-2021	
D. Gartner (USA)	Missed opportunities: low engagement of healthcare providers in HPV-related discussion with American Indian & Alaska natives, by U.S. region, 2011-2020	related
P. Hull (USA)	Improving HPV vaccination through quality improvement: coach-based versus web-based practice facilitation	•
N. Vielot (USA)	Clinician insights on the use of electronic medical records to improve human papillomavirus vaccine uptake	
K. Taavela (Finland)	The quality of life of frequently vs. infrequently screened HPV vaccinated women	
J. Paavonen (Finland)	Stopping HPV: a national network to drive elimination of cervical cancer and other HPV-related cancers in Finland	• •
E. Crespo (USA)	Oral health providers' human papillomavirus (HPV) health education training expands skills during COVID-19: a case study	
V. Colón-López (Puerto Rico)	Impact of the COVID-19 pandemic on HPV vaccination in Puerto Rico	



FC 17	Epidemiology and natural history 3 Chair: A. Del Mistro (Italy) • E. Yilmaz (Sweden)	Room 0E <b>8.00 • 9.30</b>
• Human pap to cervical s	illomavirus types in invasive cervical cancer in relation creening	J. Wang (Sweden)
	ical glandular cells: assessment of non-HPV16/18 specific risks	E. Yilmaz (Sweden)
• The HPV dis	tribution in CIN2, CIN3 and cervical cancer lesions in the s region of Madeira	D. Santos (Portugal)
	ation of type-specific HPV prevalence in a population of utaneous warts in Flanders, Belgium	N. Redzic (Belgium)
	tology in the follow-up of cervical cancer patients who received ring surgery	S. Schaafsma (Netherlands)
molecular e	community-based follow up study of HPV infection in Ethiopia: pidemiology, genotyping, persistence, clearance, and re- tes among rural women	B. Teka (Ethiopia)
and hyaluro	new supplementation based on EGCG, folic acid, vitamin B12 onic acid on patients with human papillomavirus (HPV) persistent and cervical lesions	R. Gambioli (Italy)
• Epidemiolog update	gy of cervical adenocarcinoma in situ in the HPV vaccine era: an	J. Gargano (USA)
	SIL and CIN2-3 prevalences among commercially insured olds screened for cervical cancer from 2007-2020, USA	R. Lewis (USA)



FC 18	Microbiome Chair: T. Iftner (Germany) • V. Pimenoff (Sweden)	Room 0E <b>9.30 • 11.00</b>
•	tors of persistency and CIN2+ in women infected with HR HPV genomic analysis of the cervical microbiome	T. lftner (Germany)
_	ed analysis vaginal microbiota dysbiosis and the relationship with ad in HPV positive women	M. Li (China)
· ·	omposition of the cervicovaginal microbiome correlates with ction outcomes: a longitudinal cohort study	M. Molina (Netherlands)
	crobiome associated to HPV vaccinated women who developed ve years post-vaccinated	V. Pimenoff (Sweden)
	ation of vaginal microbiome in cervical samples from the vical cancer screening program of the northern region of	A. Rosário (Portugal)
_	microbiota in cervical carcinogenesis: findings from the cervical npling in screening study	M. Logel (Canada)
	oriolus versicolor-based vaginal gel and a reishi-based food for the treatment of high-risk HPV associated lesions: a case	A-E. Del Villar Vázquez (Spain)
Therapeution with Humar	gut microbiome modulation with Microbial Ecosystem : 4 (MET4) in the context of definitive chemoradiation in patients n-papillomavirus-related or oropharyngeal squamous cell HPV+ OC) (ROMA2 trial)	M. Oliva (Spain)
• Willingness in Puerto Ri	to perform anal PAP self-collection among adults living with HIV	A. P. Ortiz (Puerto Rico)
pregnant w	nulti-ingredient coriolus versicolor-based vaginal gel in a HPV18+ oman with CIN 2/3 lesions	J. J. Hijona Elósegui (Spain)
• A conservat	ive treatment of CIN 2 using a coriolus versicolor-based vaginal rvational study	G. Pardina Claver (Spain)



FC 19	Methylation 2 Chair: B. Nedjai (UK) • H. Pedersen (Denmark)	Room 0E 11.00 • 12.30
·	fic methylation marker analysis to improve the triage of ive self-samples in the Dutch population-based cervical cancer ogramme	B. Wisman (Netherlands)
neoplasia w	ation markers for optimal detection of vulvar intraepithelial ith a high cancer risk	F. Voss (Netherlands)
rapid assess	mination of cervical cancer - increased use of biobanks enables sment of emerging biomarkers in screening	E. W. Stratford (Norway)
• Validation o	f Methica CC kit as triage test for cervical cancer screening	N. Van Belzen (Netherlands)
• Comparison	of the performance of the DNA methylation marker test and the CINtec Plus cytology assay	l. Zeiser (Germany)
• •	oval trial of Gong An Li (GynTect®), a DNA methylation assay omarkers for detecting cervical cancer and its precancerous	M. Schmitz (Germany)
• HPV vaccina	tions association to pregnancy and childbirth prevalence	T. Koivisto (Finland)
	ost and viral methylation panel in detecting high-grade cervical ng HPV-vaccinated women	K. Louvanto (Finland)
• Epigenome-	wide associations between methylation and progression to cervical intraepithelial neoplasia (cin2+): a prospective clinical	A. Bukowski (USA)



FC 20	Screening methods and self-sampling Chair: H. Ikenberg (Germany) • R. Steenbergen (Netherlands)	Room 0A <b>8.00 • 9.30</b>
•	of histological diagnoses before and after two years of coand cytology in Germany	H. Ikenberg (Germany)
opportunist	to recommended follow-up in opportunistic vs. non- ic cervical screening	S. F. Joergensen (Denmark)
• Cervical car the COVID-1	cer screening activity in the capital region of Denmark during 9 pandemic	B. T. Pedersen (Denmark)
• A nationwid cancer elim		L. S. Arroyo Mühr (Sweden)
• Exploring in women	vitation strategies for HPV self-sampling among underscreened	K. Bunzeluk (Canada)
• Experience cancer scree	from HPV self-sampling as part of a population based cervical ening program in the region of Stockholm, Sweden	D. Öhman (Sweden)
• Cervical car	cer screening improvements with self-sampling precipitated by 9 pandemic	E. Hultin (Sweden)
	nter-laboratory reproducibility of the ONC OREDICT HPV SCR and sing the VALGENT-2 framework	S. K. Dhillon (Belgium)



FC 21	<b>Triage of HPV positive women</b> Chair: C. Depuydt (Belgium) • P. Giorgi Rossi (Italy)	Room 0A <b>9.30 • 11.00</b>
• Multiple HP grade cervi	V type combinations among unvaccinated women with high- cal lesions	K. Vormisto (Finland)
and HPV RN	V primary screening study: 3 year follow up of HPV DNA positive IA positive women with normal cytology	C. White (Ireland)
	MRNA test in triage of HPV-DNA primary screen positive women	
	urements with quantitative HPV genotyping in liquid-based ows triage of HPV positive women	C. Depuydt (Belgium)
	ge dependent HPV clearance after HPV positive screening index alth care policy implications for recommended follow-up	E. Korsgaard Andreasen (Denmark)
	ytology help in the management of women with HPV 16/18 in cancer screening?	S. Bras (Portugal)
•	and clinical outcome at 12 months re-test after HPV+ index PV screening using extended genotyping and cytology as triage	A. Arday (Denmark)



FC 22	Screening / HPV testing 2 Chair: C. Eklund (Sweden) • A. Tropé (Norway)	Room 0A 11.00 • 12.30
	le trial of rapid elimination of HPV and cervical cancer	J. Dillner (Sweden)
	PV prevalence in the second round of HPV screening in Norway	B. Engesaeter (Norway)
_	obal HPV DNA typing and HPV screening proficiency studies	C. Eklund (Sweden)
• Diagnosis o	f cervical cancer in region Skåne, Sweden 2017-2020 after the ation of primary HPV screening: a quality assurance audit	C. Hellsten (Sweden)
	f an HR-HPV proficiency panel by the Belgian national reference IPV: concept and pilot projects	K. Kehoe (Belgium)
•	nd vaccination: results on number of vaccine doses from the y evaluating best strategies on how to screen vaccinated women	P. Armaroli (Italy)
•	ation sequencing (NGS) and COBAS HPV test to analysis of high- bes in European and Latin women: preliminary results from the dy	L. Godoy (Brazil)
	atch-up HPV test in women aged 65 and above: a population- randomized intervention study	M. Tranberg (Denmark)
• Cervical car	ncer screening by visual inspection and HPV testing in Eswatini	T. G. Ginindza (South Africa)



FC 23	Molecular markers Chair: J. Doorbar (UK) • S. Regauer (Austria)	Room 0C 8.00 • 9.30
	ve management of women with CIN2 lesions enrolled in a e multicentric study: clinical outcome and predictive biomarkers	S. Gori (Italy)
	al monitoring of HPV16 genomes in cervical infections and impact of H78Y and L83V mutations of HPV16-E6 protein	N. Di Domizio (France)
	tion of locoregional minimal residual disease after surgery for opharyngeal cancer using a surgical drain fluid assay	N. Earland (USA)
• Genetic eve	ents in HPV-induced and HPV-independent penile squamous cell	S. Regauer (Austria)
	of circulating human papillomavirus (HPV) DNA detection in omen with a recent history of cervical dysplasia	M. Martinelli (Italy)
• Are TP53 m HIV?	utations associated with HPV genotypes in women living with	T. Magwaku (Zimbabwe)
• HIV-1 prote oral epithel	ins Gp120 and tat promote invasiveness of neoplastic genital and ial cells	S. Tugizov (USA)



FC 24	Serology and immunotherapy Chair: O. Forslund (Sweden) • A. Kaufmann (Germany)	Room 0C <b>9.30 • 11.00</b>
quadrivale	total and neutralizing HPV18 L1 antibodies in bivalent and neutralizing to 12 years after 3-dose vaccination: a follow-up of two phase 3 trials	P. Gray (Sweden)
	rology standardization initiative: aims and progress to date at the ational laboratory for cancer research	L. Pinto (USA)
•	on of the laboratory workflow for improved efficiency and quality tine surveillance data	G. K. Presthus (Norway)
neonates fo	antibodies to HPV6 L1, E2, E4, E6 and E7 proteins among ollowed-up for three years	H. Suominen (Finland)
	filtrates among women with cervical pre-cancer and cancer	J. Rathwell (USA)
inhibitor te	eoxycytidine in combination with the cytidine deaminase trahydrouridine significantly improves survival in a preclinical del of HPV-induced cancer	L. Schlegel (Germany)
	ow-dose 5-AZA-2´-deoxycytidine (DAC) treatment on ion of HPV-transformed cells	O. Seidel (Germany)



# PC - OPENING CEREMONY with YOUNG SCIENTISTS PITCH CONTEST

Wednesday, February 8

Room 0B

Chair: J. Bonde (Denmark) • J. S. Smith (USA)

18.00 • 19.30

The aim of the session is to have 6 outstanding young researchers compete in a research presentation competition.

JURY		
<b>M. Elfström (Sweden)</b> Center for Cervical Cancer Prevention, Karolinska University Hospital	<b>A. Oštrbenk Valenčak (Slovenia)</b> University of Ljubljana Faculty of Medicine	<b>D. Hawkes (Australia)</b> Australian Centre for the Prevention of Cervical Cancer
P. E. Castle (USA)  Director, Division of Cancer Prevention, U.S. National Cancer Institute	<b>H. Botha (South Africa)</b> Stellenbosch University, South Africa	L. Chinula (Malawi) University of North Carolina (UNC) at Chapel Hill/Clinical Research Site Leader UNC Project Malawi

#### **EVALUATION SET-UP**

Each juror gets a pre-printed sheet for the 6 contestants.

The presentations are given points 0-10 for the following elements for a max point of 50:

Adherence	Clarity	Clarity	Format	X-factor
to time	of project aim	of presentation*	of presentation	

By conclusion of each presentation the Jury hands in the score sheet to the assistant for ranking. By the end the Jury will have 5 min to deliberate whether their assessment stands, and whether they agree to the Winner of the competition.

<sup>\*</sup> Data, conclusions, perspectives of research



# PC - OPENING CEREMONY with YOUNG SCIENTISTS PITCH CONTEST

Wednesday, February 8

Chair: J. Bonde (Denmark) • J. S. Smith (USA)

	S. Smith (USA)	
Introduction J. Bonde (Denmark) • J. S. Smith (USA)		Room 0B <b>18.00 • 18.10</b>
Welcome to the session presenter(s), rule	es & process	
Presentations		Room 0B <b>18.10 • 18.50</b>
5 min segment presentation by the 6 sele	ected presenters.	
<ul> <li>Methylation analysis of anal swabs: the future of anal cancer screening?</li> <li>Diagnostic accuracy of human and human papillomavirus DNA methylation testing in cervical cancer: a systematic review and meta-analysis</li> </ul>		K. Rozemeijer (Netherlands) L. Ellis (UK)
<ul> <li>Profiling HPV antibody responses 6 year HPV vaccine</li> </ul>	rs following 1, 2 or 3 doses of quadrivalent	C. Quang (Australia)
<ul> <li>Impact of mobile game fightHPV on cerv retrospective cohort study</li> </ul>	vical cancer screening attendance:	M. Orumaa (Norway)
Oral human papillomavirus prevalence a		M. Felsher (USA)
populations attending routine dental ca PROGRESS (PRevalence of Oral HPV infe • Health and economic effects of introduc vaccination in India	ection, a Global aSSessment) study	T. De Carvalho (Netherlands)
Congress Welcome	Deliberation by the Jury	Room 0B <b>18.50 • 19.15</b>
Congress Welcome  Tribute to Massimo Tommasino • S. Fr	· ·	
<b>Tribute to Massimo Tommasino •</b> S. Fr <b>Welcome by the Chairman of the EUR</b>	anceschi (Italy)  COGIN Scientific Commitee • Joseph Monsor	18.50 • 19.15 nego (France)
<b>Tribute to Massimo Tommasino •</b> S. Fr <b>Welcome by the Chairman of the EUR</b>	ranceschi (Italy)	18.50 • 19.15 nego (France)
<b>Tribute to Massimo Tommasino •</b> S. Fr <b>Welcome by the Chairman of the EUR</b>	anceschi (Italy)  COGIN Scientific Commitee • Joseph Monsor	18.50 • 19.15 nego (France)
Tribute to Massimo Tommasino • S. Fr Welcome by the Chairman of the EUR and by the Congress Presidents • Jenn	anceschi (Italy)  COGIN Scientific Commitee • Joseph Monsor	18.50 • 19.15 nego (France) rk)  Room 0B
Tribute to Massimo Tommasino • S. Fr Welcome by the Chairman of the EUR and by the Congress Presidents • Jenn Announcement of the Winner	anceschi (Italy)  COGIN Scientific Commitee • Joseph Monsor	18.50 • 19.15  nego (France) rk)  Room 0B 19.15 • 19.30  Exhibition Hall
Tribute to Massimo Tommasino • S. Fr Welcome by the Chairman of the EUR and by the Congress Presidents • Jenn Announcement of the Winner Welcome Reception	ranceschi (Italy)  COGIN Scientific Commitee • Joseph Monsor nifer S. Smith (USA) and Jesper Bonde (Denma	18.50 • 19.15  nego (France) rk)  Room 0B 19.15 • 19.30  Exhibition Hall 19.30 • 20.30  Guggenheim Museum



Coordinators: K. Lang Kuhs (USA) • J. P. Klussmann (Germany)
H. Mirghani (France) • E. Rettig (USA)

Room 0A

The EUROGIN HPV and Head & Neck Cancer Forum highlights recent advances and areas of active research in the field of HPV-related head and neck cancers. This year's Forum features talks on epidemiology and prevention, HPV-OPC screening studies, updates on current management, innovations in surveillance and new discoveries of the molecular landscape of HPV-OPC tumors. New for this year, the Forum will also feature several panel discussions exploring the potential promise and peril of screening, surgery versus chemoradiation therapy and risks versus benefit of using liquid biopsy for HPV-OPC surveillance.

Thursday, February 9		
HN 01	Screening for HPV-OPC Chair: K. Lang Kuhs (USA) • T. Waterboer (Germany)	14.00 • 15.30
• Introduct	ion	K. Lang Kuhs (USA) & T. Waterboer (Germany)
<ul><li>Prevalence</li><li>Biomarke</li><li>Relation cover the land</li></ul>	on 2 US-based HPV-OPC screening studies e of HPV biomarkers among screened populations r-based screening trial updates of prediagnostic HPV16 E6 antibodies with oropharyngeal cancer ast 40 years	K. Lang Kuhs (USA) A. D'Souza (USA) T. Waterboer (Germany) M. Lehtinen (Finland)
•	ostic liquid biopsy os vs. Cons of screening for HPV-OPC	D. Faden (USA) E. Sturgis (USA) vs. K. Lang Kuhs (USA)
• Discussio	n and Q&A	K. Lang Kuhs (USA) & T. Waterboer (Germany)
Coffee Bre	eak	15.30 • 16.00
HN 02	Basic science Chair: S. Virani (France)	16.00 • 17.30
• Introducti	on	S. Virani (France)
<ul><li> HPV thera</li><li> Cell plasti</li><li> Single cell</li></ul>	Respiratory Papillomatosis: updates speutic vaccine trials city in HPV-driven tumors sequencing in HPV-OPC stratification of OPC patients using standard H&E staining	S. Pransky (USA) A. Kejner (USA) J. Hess (Germany) S. Puram (USA) S. Klein (Germany)
• Evaluation	genetics for risk of HPV-driven head and neck cancers	E. J. Speel (Netherlands) S. Virani (France)
	n and Q&A	S. Virani (France)



Coordinators: K. Lang Kuhs (USA) • J. P. Klussmann (Germany)
H. Mirghani (France) • E. Rettig (USA)

Room 0A

		- ^
Thursday,	repruar\	79

HN 03

Submitted papers

Chair: K. Van Abel (USA)

17.30 • 19.00

- Factors independently associated with oncogenic oral-HPV infection among men from Brazil, Mexico and USA participating in the human papillomavirus Infection in Men (HIM) Study
- Laryngeal papilloma and presence of bacterial species
- Impact of circulating cell free tumour tissue modified viral-HPV DNA testing on post-treatment imaging surveillance protocol in oropharyngeal carcinoma
- Feasibility study ONCSALIVA non-invasive specimen for the detection of head and neck cancer via epigenetic biomarkers
- The role of HPV in determining treatment, survival, and prognosis of head and neck squamous cell carcinoma
- Prevalence of HPV infection in oropharyngeal cancer in Sardinian region

- R. S. Dube Mandishora (USA)
  - O. Forslund (Sweden)
  - C. Fundakowski (USA)
  - A. B. Hums (Germany)
  - C. M. Martin (Ireland)
    - N. Muresu (Italy)



Coordinators: K. Lang Kuhs (USA) • J. P. Klussmann (Germany) H. Mirghani (France) • E. Rettig (USA)

Room 0A

Friday, February 10	
HN 04 Epidemiology and prevention of HPV-OPC Chair: E. Rettig (USA)	8.00 • 9.30
• Introduction	E. Rettig (USA)
HPV-OPC incidence trends	A. Deshmukh (USA)
Oral HPV infection and HIV	N. Osazuwa-Peters (USA)
<ul> <li>HPV Vaccination in Otolaryngology Clinics: a study of feasibility, potential</li> </ul>	E. Rettig (USA)
impact and provider sentiment	
<ul> <li>HPV vaccination for prevention of oral HPV infection</li> </ul>	A. Giuliano (USA)
What should be communicated to women with oncogenic genital HPV and	G. Barbara (Italy)
their partners regarding the risk of oral viral transmission	NA Mindon (UGA)
Patient priorities and concerns	M. Windon (USA)
Discussion and Q&A	E. Rettig (USA)
Coffee Break	9.30 • 10.00
HN 05  Management  Chair: H. Mirghani (France)	10.00 • 11.30
• Introduction	H. Mirghani (France)
Outcomes beyond survival	H. Starmer (USA)
Aggressive adjuvant radiation reduction after surgery for HPV-OPC	D. Routman (USA)
ORATOR trials	M. Patel (USA)
<ul> <li>Overview of de-escalation strategies and potential of genomics for prediction</li> </ul>	M. Lechner (UK)
<ul> <li>Neoadjuvant immunotherapy pre-CRT in HPV-OPC</li> </ul>	H. Mirghani (France)
PANEL: surgery versus CRT for HPV-OPC	C. Simon (Switzerland)
	vs. R. Haddad (USA)
Discussion and Q&A	H. Mirghani (France)
Lunch Break	11.30 • 13.15



Coordinators: K. Lang Kuhs (USA) • J. P. Klussmann (Germany) H. Mirghani (France) • E. Rettig (USA)

Room 0A

	Friday, February 10	
HN 06	Submitted papers Chair: J. Hess (Germany)	13.15 • 14.30
	ed therapy in HPV+ oropharyngeal cancer tumor-tissue modified V) - HPV DNA profile   The react study	J. Schoenfeld (USA)
	of antibodies against HPV16 E6 oncoprotein by ELISA: validation of a romising biomarker for diagnosis of HPV-driven oropharyngeal cancer	L. Alemany (Spain)
	nd recurrence outcomes in HPV+ oropharyngeal squamous cell patients treated with tors: a systematic review and meta-analysis	F. Durrant (USA)
	PV vaccination education for oral health professionals: outcomes ative evaluation	J. Oliphant (USA)
	of the attributable fraction and burden of HPV-related geal cancers in Greece - The ORPHEAS study	A. Psyrri (Greece)
HN 07	Molecular diagnosis and surveillance Chair: J. P. Klussmann (Germany)	14.30 • 16.00
• Introduction	on	J. P. Klussmann (Germany)
• Prognostic	implication of p16/HPV discordance	L. Alemany (Spain)
	PV DNA in urine	C. Brenner (USA)
	DNA for surveillance in the clinic	E. Rettig (USA)
	osy techniques ed controlled trial of standard vs. liquid biopsy-based surveillance	N. Würdemann (Germany) H. Mirghani (France)
	os vs. Cons of liquid biopsy for surveillance	н. Mirghani (France) L. Mady (USA)
		vs. G. Hanna (USA)
• Discussion	and Q&A	J. P. Klussmann (Germany)



Wednesday, February 8

Room 0C

8.30 • 13.30

# WS 01 - Colposcopy Course

## Coordinators: J. Bornstein (Israel) • A. Singer (UK)

Welcome to the EUROGIN Colposcopy course. Taking care of cervical precancer has evolved significantly in recent years. However, the basis remains – Colposcopy. Performing colposcopy necessitates knowledge and experience. In this course, you will learn the fundamentals of the use of the colposcope and the essentials of diagnosing and treating precancerous cervical lesions.

The EUROGIN course has traditionally been led by professor Albert Singer, and we have the great pleasure of having him with us again this year, co-sharing the leadership of this course with Professor Jacob Bornstein, who headed the IFCPC Nomenclature committee that produced the contemporary colposcopy terminology.

Colposcopy is the visual examination of the epithelial cervix using either uni - or binocular vision. Specific abnormalities associated with both squamous and glandular precancer can be identified especially after the application of a 5% acetic acid solution. After this application, the abnormalities become visible as a result to changes in the epithelium and blood vessels in the stroma.

These changes occur within an area of the cervix called the transformation zone, an area bounded by the junction of the vaginal epithelium and the glandular epithelium arising from the endocervix (canal). Within this area, a change occurs in which glandular epithelium changes to squamous by a process of transformation, called metaplasia. The upper border of this metaplastic change is called the new squamocolumnar junction. The inability to see this junction means that abnormality may exist higher up in the endo cervix. A sample of any abnormality within the transformation zone can be taken by a simple punch biopsy.

Colposcopy is an essential part of the diagnosis and treatment of cervical precancer. It is indicated in the presence of abnormal cytology or in the finding of certain types of HPV and also when there are clinical symptoms and signs of early invasive cancer.

#### **Educational Objectives**

Upon completion of this educational activity, participants should be able to:

- Describe the anatomy, cytology, histology, and colposcopic findings of the normal and abnormal cervix;
- Define the pathophysiology of lower genital tract neoplasia, including the role of HPV in preinvasive and invasive diseases of the cervix;
- Define the IFCPC colposcopy terminology;
- Recognize the diagnostic characteristics of cervical abnormalities (minor-grade and major-grade cervical lesions as well as glandular lesions and cervical cancer) on the cytologic, colposcopic, and histologic examination;
- Interpret and correlate cytologic, colposcopic, and histologic results;
- Describe treatment options to include cryosurgery and large loop excision of the transformation zone (LLETZ) of the cervix;
- Provide appropriate patient education and support.



Wednesday, February 8

# WS 01 - Colposcopy Course

Room 0C 8.30 • 13.30

Coordinators: J. Bornstein (Israel) • A. Singer (UK)

WS 01-A Part A	8.30 • 10.55
• Opening	J. Bornstein (Israel)
The normal cervix and the colposcopy examination	A. Singer (UK)
Update in pathology and cytology for colposcopists	S. Regauer (Austria)
Colposcopy of «abnormal» cervix, colposcopy terminology	J. Bornstein (Israel)
WS 01-B Part B	10.55 • 13.30
<ul> <li>Management protocols of abnormal screening findings and the value of biomarkers</li> </ul>	J. Bonde (Denmark)
Treatment of cervical precancer and treatment's complications	J. Bornstein (Israel)
Interactive session (What is your diagnosis)	A. Singer (UK)
• Course Summary	A. Singer (UK)



Wednesday, February 8

## WS 02 - Cervical cancer screening workshop

Room 0D

15.30 • 18.00

Coordinators: M. Elfström (Sweden) • P. Giorgi Rossi (Italy)
M. Rebolj (UK) • S. Van Dijk (Netherlands)

The workfield of cervical cancer screening is in full development. Due to tremendous efforts of researchers and researchgroups worldwide, we have already been able to achieve a lot in the field of innovation. In recent years, more evidence for new screening technologies has become available from scientific research. In order to achieve the objective of the WHO, it is important that research is implemented into cervical cancer screening programs. However, this implementation is not self-evident and can sometimes take quite a long time. By means of a discussion between researchers and clinicians, we want to provide more insight into this transition from research to implementation on the basis of 3 potential new screening technologies on the roadmap to eliminate cervical cancer: HPV-screening, self-sampling and methylation.

#### WS 02-A

#### Introduction

An overview of the development of cervical cancer screening over the years, focusing on the achievements until now, in terms of absolute health gains and in terms of reduction and increase in inequities. And how further improvement can be reached by the introduction of (potential new) screening technologies.

P. Sasieni (UK)

#### WS 02-B

#### • HPV-based-screening PRO

Research has shown that primary HPV screening can increase the sensitivity of screening programs. In addition HPV-testing may be accessible to more countries, as it is cheaper and easier to perform. And it's opening up opportunities for self-sampling and HPV-vaccinated cohorts. So, it's time to implement primary HPV screening everywhere to eliminate cervical cancer.

H. Bogaards (Netherlands)

#### HPV-based-screening CON

The time may be right scientifically, but then there are several practical obstacles that make it impossible to move quickly in certain countries. The Slovenian organized programme is coping with some unique challenges. Certain conditions must be met when implementing primary HPV screening.

U. Ivanus (Slovenia)

#### Discussion

A discussion will be held with the participants of the workshop focusing on the conditions under which countries could successfully implement HPV-based screening.



Wednesday, February 8

# WS 02 - Cervical cancer screening workshop

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#### WS 02-C

#### Self-sampling PRO

Research in recent years gives strong indications that self-sampling can reach a significant part of the non-responder population of cervical cancer screening programs. Experiences from Sweden, especially since COVID-19-times have shown great opportunities for self-sampling. Self-sampling is easy and reliable. Therefore, the time is right for the primary use of self-sampling in screening programs.

M. Elfström (Sweden)

#### Self-sampling CON

We need to move carefully on self-sampling, particularly in countries that have been operating a very efficient and quality-assured screening service for a long time. The early-adopter evidence is eye-opening in this respect. We will discuss the necessary mitigations, impact of lab-processes and what kind of research work needs to be prioritized. Not for every country, the time is right to introduce self-sampling.

A. Sargent (UK)

#### Discussion

A discussion will be held with the participants of the workshop focusing on the conditions under which countries could successfully implement self-sampling.

#### WS 02-D

#### Methylation PRO

Research in the field of biomarkers indicates that methylation as a triage method could increase the specificity of primary HPV screening programs. Introduction of primary HPV screening in several countries has shown that the sensitivity of the program increases, but at the same time the specificity decreases resulting in a higher proportion of unnecessary referrals. So, methylation shows enormous opportunities and the time is right to introduce methylation in the management of HPV positive women.

C. Meijer (Netherlands)

#### Methylation CON

To implement a new technique into organized screening programs, we need more than just research. For instance within the domains of equity, feasibility, acceptability, (balance of) desirable and undesirable effects, validation, CE-approval, high-throughput machines, trained professionals and cost-effectiveness. We cannot implement methylation in screening (yet).

S. Van Dijk (Netherlands)

#### Discussion

A discussion will be held with the participants of the workshop focusing on the conditions under which countries could successfully implement methylation.



## **LW - WORKSHOP AEPCC**

Viernes, 10 de Febrero

# **Highlights Congreso AEPCC**

Room 0D **15.30 • 18.00** 

Comité científico: A. Alba • J. De La Fuente • S. De Sanjosé

M. Del Pino • M. Ramírez • A. Torné

Organiza: Asociación Española de Patología Cervical y Colposcopia



LW 01 Part 1	14.30 • 16.30
Vacunación VPH	J. De La Fuente
Cribado del cáncer de cérvix en España	M. Ramírez
Patología vulvar premaligna	Mª P. Cano
Patología anal premaligna	M. Del Pino
Debate	Modera: <b>D. Andía</b>
Charla magistral - Infección por VPH - Transmisión y latencia	E. Franco (Canadá)
Preguntas	Modera: S. De Sanjosé
Pausa Café	16.30 • 17.00
LW 02 Part 2	17.00 • 19.30
Charla magistral - Objetivo: eliminación cáncer de cérvix	Modera: L. Bruni
Preguntas	Modera: <b>S. De Sanjosé</b>
Presentación AEPCC-Guía	A. Torné
Novedades en VPH/LSIL	L. Fernández-Villarrenaga
Novedades en HSIL/AIS	J. C. Quílez
Comité Clínico	Moderan: M. Del Pino & A. Torné



## WF - WORKSHOP HPV FRANCE

Coordinateur: J. Monsonego

(Coordinator)

Jeudi 9 Février (Thursday, February 9)

14.00 • 17.00 | Salle 5H Terrasse

(Room 5H Terrace)

## Élimination du cancer du col utérin en marche

Comment la France peut parvenir à une stratégie nationale d'élimination des cancers HPV-induits ?

Dans son rapport de lutte contre le cancer pour 2020-2030, l'InCA nous indique qu'en France le taux de couverture du dépistage au cancer du col utérin est de 59,5% et celui de la vaccination HPV est de moins de 25% pour les jeunes filles de 16 ans. La feuille de route présente des éléments d'action et des objectifs ambitieux à l'horizon 2030.

Alors que l'OMS a fixé en 2019-2020 un objectif clair et détaillé, à savoir l'élimination du cancer du col dans les 10 années à venir, et que de nombreux pays – en particulier européens – ont amorcé ce tournant décisif, la France progresse peu dans ce domaine, bien que des efforts stratégiques et de communication aient été entrepris ces dernières années.

Le constat montre malheureusement qu'il nous faut aborder cette question de manière pragmatique, collaborative et dynamique et faire en sorte que les rapports qui se suivent ne soient pas que de simples idées mais de justes orientations appliquées.

Le cancer du col est totalement évitable ; nous avons à notre disposition un vaccin très efficace en termes de protection presque absolue et bien toléré, un dépistage HPV censé éviter le cancer par la détection précoce des pré-cancers, et des méthodes de diagnostic et de traitement de plus en plus performantes.

Ce meeting d'experts se fixe comme objectifs :

- · Confronter les expériences européennes avec la nôtre,
- Répertorier les problématiques, les dérives et les difficultés rencontrées,
- Proposer des pistes de veille programme de dépistage HPV disponible depuis 3 ans,
- Analyser pourquoi la politique de prévention vaccinale ne porte pas ses fruits dans notre pays alors que le vaccin HPV est disponible et remboursé depuis une quinzaine d'années.

Avec la contribution des acteurs scientifiques, professionnels, institutionnels, de la société civile et associatifs, ce workshop, organisé dans un contexte international, veut faire prendre conscience et alerter, analyser et proposer des orientations afin d'encourager et faciliter les recommandations institutionnelles.

# Elimination of cervical cancer under way

How France can succeed to set up a national strategy to eliminate HPV induced cancers?

In its report on fighting cancer for the period 2020-2030, the French National Cancer Institute InCA indicates that the coverage rate for cervical cancer screening is 59.5% and that the HPV vaccine coverage rate is less than 25% for girls aged 16. The report presents ambitious goals and elements of action for 2030.

Whereas in 2019-2020 the WHO has set a clear and detailed objective of elimination of cervical cancer in the next 10 years, many countries – in particular countries in Europe – managed to make decisive headway. In France progress is still slow despite various strategic and communication efforts undertaken in recent years.

Unfortunately, one has to concede that the problem needs to be tackled in a pragmatic, collaborative and active way to ensure that the reports do not simply reflect ideas but also give guidance on how to apply these.

Cervical cancer is fully evitable. All tools are fully available, offering a highly efficient and safe vaccine, HPV screening for optimal early detection of pre-cancers and ever better diagnostic and treatment procedures.

The objectives of this expert meeting are:

- · Confront and share European experiences,
- Take stock of implementation problems, errors and diffulties encountered,
- Efficient roll-out of the HPV screening program which has been available for 3 years,
- Understanding why the vaccination policies are not successful although HPV vaccine has been available and reimbursed for the last 15 years.

In an international context and with the contribution of representatives from science, health care providers, institutions, civil society and associations, this workshop endeavours to raise awareness, to alert, analyse and offer directions to encourage and facilitate institutional recommendations.



## WF - WORKSHOP HPV FRANCE

Jeudi 9 Février

Salle 5H Terrasse

## **Programme**

WF 01	Introduction & objectifs du Workshop HPV France J. Monsonego	14.00 • 14.10
WF 02	Élimination des cancers HPV induits au niveau international - Objectifs et recommandations P. Basu	14.10 • 14.30
WF 03	Partage d'expériences européennes (dépistage et vaccination)	14.30 • 15.30

Chaque représentant d'un pays présente 3 slides en 5 minutes, sur l'état des lieux, la couverture, la mise en œuvre et les performances. Les sujets suivants seront abordés :

- Dépistage
- Vaccination
- Difficultés

## **PAYS INVITÉS**



P. Sasieni





Allemagne

P. Hillemanns



Italie
P. Giorgi Rossi



Pays-Bas S. Van Dijk



Finlande **K. Louvanto** 



M. Elfström



🖵 B. Frey-Tirri\*

**☐** Présentation vidéo

Table ronde - Comment la France peut-elle WF 04 rattraper son retard ? •••

Modérateur : J. Monsonego

15.30 • 16.30

- Une ambition : quels objectifs ?
- Définition de notre ambition et de nos objectifs
- Est-on plus efficace ensemble? Comment se coordonner?

Comment créer une coalition des parties prenantes scientifiques, médicales, associatives et institutionnelles?

- Un pilote dans l'avion?
- Comment inviter les pouvoirs publics à prendre leur place légitime ?
- Quelles actions concrètes pour 2023?

#### **PARTICIPANTS**

L. Abramowitz	B. Julia B.	J. L.Brun
H. Baffet	O. Launay O.	C. Clavel
C. Charpentier	J. B. Lusignan	A. S. Le Duc-Banaszuk
R. Cohen	A. Menard	J. Nicolet
D. Culié	H. Mirghani	C. Marjollet
M. A. Dommergues	H. Peré	J. P. Martin
G. Dolivet	F. Vié Le Sage	P. Pautier
O. Jourdain	K. Ardaens	

WF 05 Conclusions et prochaines étapes

16.30 • 17.00



## WF - WORKSHOP HPV FRANCE

Thursday, February 9

Salle 5H Terrace

## Agenda

WF 01	Introduction & objectives of the Workshop HPV France J. Monsonego	14.00 • 14.10
WF 02	Elimination of HPV induced cancers - International objectives and recommendations P. Basu	14.10 • 14.30
WF 03	Sharing experiences from Europe (screening and vaccination)	14.30 • 15.30

Each country representative gives a very brief outline (5 minutes and 3 slides maximum) of the situation, coverage, performance and difficulties, addressing:

- Screening
- Vaccination
- Difficulties

#### **INVITED COUNTRIES**















P. Sasieni

Germany
P. Hillemanns

Italy

P. Giorgi Rossi

Netherlands S. Van Dijk

K. Louvanto

M. Elfström

B. Frey-Tirri\*

**□** Video presentation

Table ronde - Comment la France peut-elle

WF 04 rattraper son retard ? 🕩

15.30 • 16.30

Moderator: J. Monsonego

- Une ambition : quels objectifs?
- Définition de notre ambition et de nos objectifs
- Est-on plus efficace ensemble ? Comment se coordonner ? Comment créer une coalition des parties prenantes scientifiques, médicales, associatives et institutionnelles ?
- Un pilote dans l'avion?

Comment inviter les pouvoirs publics à prendre leur place légitime ?

• Quelles actions concrètes pour 2023?

#### **PARTICIPANTS**

· / · · · · · · · · · · · · · · · · · ·		
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M. A. Dommergues	H. Peré	J. P. Martin
G. Dolivet	F. Vié Le Sage	P. Pautier
O. Jourdain	K. Ardaens	

WF 05 Conclusions and next steps

16.30 • 17.30