

MSS - MAIN SCIENTIFIC SESSIONS

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.

Lunch Break 11.30 • 13.30

New guidelines for validation of HPV tests

MSS 02 for cervical cancer screening

Room 0B

13.30 • 15.00

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

Coffee Break 15.00 • 15.30

MSS 03

Monitoring HPV vaccine impact in males (including in MSM programmes - data in these are picking up)

Room 0B

15.30 • 17.00

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

Gender neural 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.



MSS - MAIN SCIENTIFIC SESSIONS

Thursday, February 9

MSS 04

Scientific approaches towards improving the cervical cancer elimination strategies

Room 0B

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

8.00 • 9.30

M. Lehtinen (Finland)

MSS 05

Debate on controversial topics

Room 0B

Chair: 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 2022 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.

MSS 06

Self-sampling in routine practice, operational

Room 0B

and technical issues

14.00 • 15.30

Chair: J. Bonde (Denmark) • C. Cocuzza (Italy)

Coffee Break

15.30 • 16.00

MSS 07

Prevention of cervical cancer among people

Room 0B

living with HIV

16.00 • 17.30

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.

MSS 08

HPV and host methylation as triage tools for cervical cancer screening in different clinical contexts

Room 0B

Chair: M. Clarke (USA) • C. Meijer (Netherlands)

17.30 • 19.00



MSS - MAIN SCIENTIFIC SESSIONS

Friday, February 10

MSS 09

VALGENT / VALHUDES

Room 0B

Chair: M. Arbyn (Belgium) • C. Cocuzza (Italy)

8.00 • 9.30

Coffee Break

9.30 • 10.00

MSS 10

Impact of HPV vaccine on cancer

Room 0B

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

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.

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 Covid19 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 were issues solved, and present evidence on the performance of self-sampling in different settings.

Coffee Break 16.00 • 16.30

MSS 12

Extended HPV genotyping in screening Chair: J. Bonde (Denmark) • E. Franco (Canada)

Room 0B

16.30 • 18.00

MSS 13

Next generation HPV based sequencing:

Room 0B

What have we learned and what's next?

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.



SS - SCIENTIFIC SESSIONS

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.

Criteria for HPV tests validation protocols for SS 02 ASCUS triage and for post-treatment monitoring

Room 0D

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

10.00 • 11.30

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. 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. Second part of the session will review clinical rationale for the use of HPV testing for ASCUS triage indication followed by description of proposed criteria for HPV tests validation protocols for this important indication. 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 description of proposed criteria for HPV tests validation protocols for post-treatment monitoring.

SS 03

Transmission: role of the male partner

Room 0D

Chair: E. Franco (Canada) • M. Goodman (USA)

13.30 • 15.00

SS - SCIENTIFIC SESSIONS

Thursday, February 9

SS 04	New screening algorithms in European countries Chair: M. Arbyn (Belgium) • P. Hillemanns (Germany)	Room 0E 8.00 • 9.30
SS 05	Molecular triage of HPV-positive women Chair: M. Arbyn (Belgium) • C. Cocuzza (Italy)	Room 0E 9.30 • 11.00



SS - SCIENTIFIC SESSIONS

Thursday, February 9

SS 06

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

Room 0E

14.00 • 15.30

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

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. On-going efforts to accumulate additional information on the efficacy and durability of different HPV vaccine options will also be highlighted.

Coffee Break 15.30 • 16.00

SS 07

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

Room 0E

16.00 • 17.30

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

The implementation and evolution of cervical screening programmes 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.

SS 08

Digital interventions to increase HPV vaccination

Room 0E

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

17.30 • 19.00





SS - SCIENTIFIC SESSIONS

Thursday, February 9

Tumour HPV status and implications for survival SS 09 outcomes in cervical and non-cervical disease

Room 0D

14.00 • 15.30

Chair: 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.

Coffee Break 15.30 • 16.00

SS 10

Microbiome

Chair: A. Lev Sagie (Israel) • A. B. Moscicki (USA)

Room 0D

16.00 • 17.30

SS 11

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

Room 0D

17.30 • 19.00

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.



PRELIMINARY PROGRAM

SS - SCIENTIFIC SESSIONS

Friday, February 10

SS 12 Benefits and harms of HPV screening

Room 0E

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

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.

Coffee Break 9.30 • 10.00

New triage methods

Room 0E

Chair: F. Carozzi (Italy) • J. Cuzick (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.

SS 14

SS 13

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 the Europe, both in term 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 presentation on HPV vaccine implementation in Hungary – a country with the highest HPV coverage rate in the region.

Lunch Break 11.30 • 13.15



PRELIMINARY PROGRAM

SS - SCIENTIFIC SESSIONS

Friday, February 10

The role of obesity in cervical cancer screening SS 15 and management

Room 0B

Chair: M. Clarke (USA)

13.15 • 14.15

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.

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 options for these clinicians who want to detect an anal malignancy as early as possible?

Coffee Break 16.00 • 16.30



SS - SCIENTIFIC SESSIONS

Friday, February 10

 SS 17
 HPV latency
 Room 0E

 Chair: A. B. Moscicki (USA)
 16.30 • 18.00

SS 18 Risk stratification in the follow up
Chair: F. Carozzi (Italy) • N. Wentzensen (USA)
Room 0E
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.

The utility of urine

Room 0A

for improved cervical cancer prevention

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

16.30 • 18.00

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.



CS - CLINICAL SESSIONS

Wednesday, February 8

CS 01	Management of CIN Chair: E. Paraskevaidis (Greece)	Room 0E 15.30 • 16.45
CS 02	Test of cure: strategies for the follow up of women treated for cervical intraepithelial neoplasia Chair: C. Cocuzza (Italy) • K. Cuschieri (UK)	Room 0E 16.45 • 18.15

CS - CLINICAL SESSIONS

Thursday, February 9

CS 03

Colposcopy in HPV era

Room 0D

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

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.

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
Coffee Break		9.30 • 10.00
CS 05	Vulvar Intraepithelial Neoplasia (VuIN) update Chair: M. Bleeker (Netherlands) • J. Bornstein (Israel)	Room 0C 10.00 • 11.30



FC - FREE COMMUNICATIONS

Wednesday, February 8

FC 01	Genotyping Chair: J. Andrews (USA)	Room 0A 10.00 • 11.30
FC 02	Anal neoplasia Chair: J. Palfesky (USA)	Room 0D 12.00 • 13.30
FC 03	Self-sampling 1 Chair: F. Inturrisi (Netherlands) • P. Sasieni (UK)	Room 0D 13.30 • 15.00
FC 04	Epidemiology and natural history 1 Chair: M. Goodman (USA)	Room 0A 12.00 • 13.30
FC 05	Vaccines 1 Chair: P. Bonanni (Italy)	Room 0A 13.30 • 15.00
FC 06	Methylation 1 Chair: S. Van Keer (Belgium) • B. Wisman (Netherlands)	Room 0A 15.30 • 17.00
FC 07	New technologies / Artificial intelligence Chair: P. Nieminen (Finland)	Room 0C 13.30 • 15.00
FC 08	Colposcopy / Management Chair: M. Hampl (Germany) • E. Siegler (Israel)	Room 0C 15.30 • 17.00



FC - FREE COMMUNICATIONS

Thursday, February 9

Chair: O. Gassama (Senegal) • J. Smith (USA) 8.00 • 9 Low income countries 2	FC 09	Self-sampling 2 Chair: C. Cocuzza (Italy) • G. Ogilvie (Canada)	Room 0D 8.00 • 9.30
Chair: O. Gassama (Senegal) • J. Smith (USA) Low income countries 2 Roon			
Low income countries 2	FC 10		Room 0A 8.00 • 9.30
Low income countries 2			
Chair: O. Gassama (Senegal) • J. Smith (USA) 9.30 • 11	FC 11		Room 0A 9.30 • 11.00

FC - FREE COMMUNICATIONS

Friday, February 10

FC 12	Vaccines 2 Chair: E. Joura (Austria)	Room 0D 8.00 • 10.00
FC 13	Screening 1 Chair: J. Bogaards (Netherlands) • S. Van Dijk (Netherlands)	Room 0C 14.30 • 16.00
FC 14	Epidemiology and natural history 2 Chair: S. Franceschi (Italy) • S. Nygård (Norway)	Room 0C 16.30 • 18.00
FC 15	Economics and modelling Chair: M. Brisson (Canada) • M. Drolet (Canada)	Room 0A 18.00 • 19.30



FC - FREE COMMUNICATIONS

Saturday, February 11

FC 16	Health education – Advocacy – Public Health Chair: N. Osazuwa-Peters (USA) • J. Paavonen (Finland)	Room 0C 8.00 • 9.30
FC 17	Epidemiology and natural history 3 Chair: A. Del Mistro (Italy) • E. Yilmaz (Sweden)	Room 0E 8.00 • 9.30
FC 18	Microbiome Chair: T. Iftner (Germany) • V. Pimenoff (Sweden)	Room 0E 9.30 • 11.00
FC 19	Methylation 2 Chair: B. Nedjai (UK) • H. Pedersen (Denmark)	Room 0E 11.00 • 12.30
FC 20	Screening methods and self-sampling Chair: H. Ikenberg (Germany) • R. Steenbergen (Netherlands)	Room 0D 8.00 • 9.30
FC 21	Triage of HPV positive women Chair: C. Depuydt (Belgium) • P. Giorgi Rossi (Italy)	Room 0D 9.30 • 11.00
FC 22	Screening 2 Chair: C. Eklund (Sweden) • A. Tropé (Norway)	Room 0D 11.00 • 12.30
FC 23	Molecular markers Chair: S. Regauer (Austria)	Room 0C 9.30 • 11.00
FC 24	Serology and immunotherapy Chair: O. Forslund (Sweden) • A. Kaufmann (Germany)	Room 0C 11.00 • 12.30



PC 01 - YOUNG SCIENTISTS PITCH CONTEST

Chair: J. Bonde (Denmark)

Room 0B

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

JURY			
M. Elfström (Sweden) Center for Cervical Cancer Prevention, Karolinska University Hospital	A. Oštrbenk Valenčak (Slovenia) University of Ljubljana Faculty of Medicine	D. Hawkes (Australia) Australian Centre for the Prevention of Cervical Cancer	
P. E. Castle (USA) Director, Division of Cancer Prevention, US National Cancer Institute	H. Botha (South Africa) 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.

^{*} Data, conclusions, perspectives of research



PC 01 - YOUNG SCIENTISTS PITCH CONTEST

Chair: J. Bonde (Denmark)

Room 0B

AGENDA

1. Welcome

Welcome by the session presenter(s), rules & process

2. Presentations

5 min segment presentation by the 6 selected presenters, including shifts

- Methylation analysis of anal swabs: the future of anal cancer screening?
- Diagnostic accuracy of human and Human Papillomavirus DNA methylation testing in cervical cancer: A systematic review and meta-analysis
- Profiling HPV antibody responses 6 years following 1, 2 or 3 doses of quadrivalent HPV vaccine
- Impact of mobile game fightHPV on cervical cancer screening attendance: retrospective cohort study
- A prospective study on association of high-risk HPV in oral cancer among Indian cases
- Oral Human Papillomavirus prevalence and risk factors among healthy populations attending routine dental care in the United States: Results from the PROGRESS (PRevalence of Oral HPV infection, a Global aSSessment) study
- Health and economic effects of introducing single-dose human papillomavirus vaccination in India

K. Rozemeijer (Netherlands)

L. Ellis (UK)

C. Quang (Australia)

M. Orumaa (Norway)

P. Tanwar (India)

M. Felsher (USA)

T. De Carvalho (Netherlands)

3. Deliberation

Deliberation by the Jury

4. Winner Announcement

Presentation of the Winner



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.

11515 VC1545 E	reflection daining inquite biopay for the viole and veinance.	
	Thursday, February 9	
HN 01	Screening for HPV-OPC Chair: K. Lang Kuhs (USA) • T. Waterboer (Germany)	14.00 • 15.30
• Updates c	n 2 US-based HPV-OPC screening studies	K. Lang Kuhs (USA)
• Prevalence	e of HPV biomarkers among screened populations	A. D'Souza (USA)
• Biomarke	r-based screening trial updates	T. Waterboer (Germany)
	f prediagnostic HPV16 E6 antibodies with oropharyngeal cancer ast 40 years	M. Lehtinen (Finland)
• Prediagno	stic liquid biopsy	D. Faden (USA)
PANEL: Pros vs. Cons of screening for HPV-OPC		E. Sturgis (USA)
		vs. K. Lang Kuhs (USA)
Coffee Bre	ak	15.30 • 16.00
HN 02	Basic science Chair: S. Virani (France)	16.00 • 17.30
• Recurrent	Respiratory Papillomatosis: Updates	S. Pransky (USA)
HPV therapeutic vaccine trials		A. Kejner (USA)
Cell plasticity in HPV-driven tumors		J. Hess (Germany)
Single cell sequencing in HPV-OPC		S. Puram (USA)
 Granular stratification of OPC patients using standard H&E staining 		S. Klein (Germany)
• Evaluation	n of HPV genome integration	E. J. Speel (Netherlands)
 Germline Genetics for risk of HPV-driven head and neck cancers 		S. Virani (France)



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

Room 0A

Thursd	lay, Feb	ruary 9
--------	----------	---------

	, ,	
HN 03	Submitted papers Chair: K. Van Able (USA)	17.30 • 19.00
men from	dependently associated with oncogenic oral-HPV infection among Brazil, Mexico and USA participating in the Human Papillomavirus n Men (HIM) Study	R. S. Dube Mandishora (USA)
• Laryngeal	papilloma and presence of bacterial species	O. Forslund (Sweden)
• Impact of	circulating cell free tumour tissue modified viral-HPV DNA testing	C. Fundakowski (USA)
on post-tre	eatment imaging surveillance protocol in oropharyngeal carcinoma	
 Feasibility 	study ONCSALIVA - non-invasive specimen for the detection of	A. B. Hums (Germany)
head and i	neck cancer via epigenetic biomarkers	
• HPV (genit	al warts in pregnancy)-diagnosis and a new therapeutic approach	I. Jeremic (USA)
the cause	of laryngeal polyps in children	
• The role of	f HPV in determining treatment, survival, and prognosis of head	C. M. Martin (Ireland)
and neck s	quamous cell carcinoma	
• Prevalence	e of HPV infection in oropharyngeal cancer in Sardinian region	N. Muresu (Italy)
 Evaluation 	of the attributable fraction and burden of HPV-related	A. Psyrri (Greece)
oropharyr	ngeal cancers in Greece - The ORPHEAS study	
 Quantifica 	tion of HPV16 cell-free DNA in liquid biopsies for early detection of	F. Rosing (Germany)
HPV-drive	n oropharyngeal cancer	



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

Room 0A

Friday, February 10		
HN 04 Epidemiolog Chair: E. Rettig	y and prevention of HPV-OPC (USA)	8.00 • 9.30
 HPV-OPC incidence trends Oral HPV infection and HIV Impact of tonsillectomy on risk of HPV-OPC HPV vaccination for prevention of oral HPV infection What should be communicated to women with oncogenic genital HPV and their partners regarding the risk of oral viral transmission 		A. Deshmukh (USA) N. Osazuwa-Peters (USA) J. D. Combes (France) A. Giuliano (USA) G. Barbara (Italy)
Patient priorities and concerns		M. Windon (USA)
Coffee Break		9.30 • 10.00
HN 05 Managemer Chair: H. Mirgho		10.00 • 11.30
• ORATOR trials	•	H. Starmer (USA) D. Routman (USA) M. Patel (USA) M. Lechner (UK) H. Mirghani (France) C. Simon (Switzerland) vs. R. Haddad (USA)
Lunch Break		11.30 • 13.15



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

Room 0A

g		
Friday, February 10		
HN 06	Submitted papers Chair: J. Hess (Germany)	13.15 • 14.30
 Risk-adapted therapy in HPV+ oropharyngeal cancer tumor-tissue modified virus (TTMV) - HPV DNA profile The react study 		J. Schoenfeld (USA)
 Detection of antibodies against HPV16 E6 oncoprotein by ELISA: validation of a new and promising biomarker for diagnosis of HPV-driven oropharyngeal cancer 		L. Alemany (Spain)
 Survival and recurrence outcomes in HPV+ oropharyngeal squamous cell carcinoma patients treated with tors: a systematic review and meta-analysis 		F. Durrant (USA)
Effective HPV vaccination education for oral health professionals: outcomes of a qualitative evaluation		J. Oliphant (USA)
HN 07	Molecular diagnosis and surveillance Chair: J. P. Klussmann (Germany)	14.30 • 16.00
· ·	implication of p16/HPV discordance	L. Alemany (Spain)
Cell-free HPV DNA in urine		C. Brenner (USA)
TTMV-HPV DNA for surveillance in the clinic		E. Rettig (USA)
Liquid biopsy techniques		N. Würdemann (Germany)
• Randomized controlled trial of standard vs. liquid biopsy-based surveillance		H. Mirghani (France)
• PANEL: Pro	s vs. Cons of liquid biopsy for surveillance	L. Mady (USA)
		vs. G. Hanna (USA)



WS - SPECIALIZED WORKSHOP

Wednesday, February 8

WS 01 - Colposcopy Course

Room 0C 8.30 • 13.30

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

WS 01-1 Colpocopy Course - Part 1	8.30 • 10.55
Opening J. Bornstein (Israel)	8.30 • 8.40
The normal cervix and the colposcopy examination A. Singer (UK)	8.40 • 9.25
Update in pathology and cytology for colposcopists S. Regauer (Austria)	9.25 • 10.10
Colposcopy of «abnormal» cervix, colposcopy terminology J. Bornstein (Israel)	10.10 • 10.55
WS 01-2 Colpocopy Course - Part 2	10.55 • 13.30
Management protocols of abnormal screening findings and the value of biomarkers J. Bonde (Denmark)	10.55 • 11.40
	V 40 10 0F
Treatment of cervical precancer and treatment's complications J. Bornstein (Israel)	11.40 • 12.25
·	12.25 • 13.15



WS - SPECIALIZED WORKSHOP

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.

Introduction

P. Sasieni (UK)

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.

HPV-based-screening PRO

H. Bogaards (Netherlands)

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.

HPV-based-screening CON

U. Ivanus (Slovenia)

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.

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.



WS - SPECIALIZED WORKSHOP

Wednesday, February 8

Room 0D

15.30 • 18.00

WS 02 - Cervical cancer screening workshop

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

Self-sampling PRO

M. Elfström (Sweden)

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 COVID19-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.

Self-sampling CON

A. Sargent (UK)

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.

Discussion

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

Methylation PRO

C. Meijer (Netherlands)

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.

Methylation CON

S. Van Dijk (Netherlands)

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).

Discussion

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



LW - PROGRAMA WORKSHOP AEPCC

Viernes, 10 de Febrero

Comité científico: A. Alba • J. De La Fuente • S. De Sanjosé
M. Del Pino • M. Ramírez • A. Torné

Room 0D **14.30 • 19.30**

Organiza: Asociación Española de Patología Cervical y Colposcopia



LW 01-1 Programa Workshop AEPCC - Part 1	14.30 • 16.30
Vacunación VPH J. De La Fuente	14.30 • 14.45
Cribado del cáncer de cérvix en España M. Ramírez	14.45 • 15.00
Patología vulvar premaligna Mª P. Cano	15.00 • 15.15
Patología anal premaligna M. Del Pino	15.15 • 15.30
Debate Modera: D. Andía	15.30 • 15.45
Charla magistral - Infección por VPH - Transmisión y latencia E. Franco (Canadá)	15.45 • 16.15
Preguntas	10.15 10.00
Modera: S. De Sanjosé	16.15 • 16.30
	16.30 • 17.00
Modera: S. De Sanjosé	
Modera: S. De Sanjosé Pausa Café	16.30 • 17.00
Pausa Café LW 01-2 Programa Workshop AEPCC - Part 2 Charla magistral - Objetivo: eliminación cáncer de cérvix	16.30 • 17.00 17.00 • 19.30
Pausa Café LW 01-2 Programa Workshop AEPCC - Part 2 Charla magistral - Objetivo: eliminación cáncer de cérvix Modera: L. Bruni Preguntas	16.30 • 17.00 17.00 • 19.30 17.00 • 17.30
Pausa Café LW 01-2 Programa Workshop AEPCC - Part 2 Charla magistral - Objetivo: eliminación cáncer de cérvix Modera: L. Bruni Preguntas Modera: S. De Sanjosé Presentación AEPCC-Guía	16.30 • 17.00 17.00 • 19.30 17.00 • 17.30 17.30 • 17.45
Pausa Café LW 01-2 Programa Workshop AEPCC - Part 2 Charla magistral - Objetivo: eliminación cáncer de cérvix Modera: L. Bruni Preguntas Modera: S. De Sanjosé Presentación AEPCC-Guía A. Torné Novedades en VPH/LSIL	16.30 • 17.00 17.00 • 19.30 17.00 • 17.30 17.30 • 17.45 17.45 • 18.00



WF - WORKSHOP HPV FRANCE

Coordinateur: J. Monsonego

(Coordinator)

Jeudi 9 Février (Thursday, February 9)

14.00 • 17.30 | Salle 3F

(Room 3F)

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,
- Déployer efficacement le 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 available: 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

Programme

WF 01	Introduction et 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 (IARC / OMS)	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

P. Giorgi Rossi

S. Van Dijk

K. Louvanto

M. Elfström

B. Frey-Tirri*

WF 04

Table ronde

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

H. Baffet (SFCPCV) • M. A. Dommergues & F. Vié Le Sage (Infovac) • G. Dolivet (SFCO) B. Julia (URPS Pharmaciens) • A. S. Le Duc-Banaszuk (CRDC Pays de la Loire) C. Marjollet (IMAGYN) • J. P. Martin (Ligue Contre le Cancer)

L. Abramowitz • J. L. Brun • C. Clavel • R. Cohen • D. Culié • O. Jourdain • O. Launay • J. Levêque J. B. Lusignan • A. Mesnard • H. Mirghani • J. Nicolet • H. Peré

^{*} Swiss HPV Alliance Initiative



WF - WORKSHOP HPV FRANCE

Thursday, February 9

Agenda

WF 01	Introduction and objectives of the Workshop HPV France J. Monsonego	14.00 • 14.10
WF 02	Elimination of HPV induced cancers - International objectives and recommendations () (IARC / WHO)	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 min. and 3 slides maximum) of the situation, coverage, performance and difficulties, addressing:

- Screening
- Vaccination
- Difficulties

INVITED COUNTRIES















P. Sasieni

P. Hillemanns

Italy P. Giorgi Rossi

S. Van Dijk

K. Louvanto

M. Elfström

B. Frey-Tirri*

WF 04

Round table ()

Moderator: 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

H. Baffet (SFCPCV) • M. A. Dommergues & F. Vié Le Sage (Infovac) • G. Dolivet (SFCO) B. Julia (URPS Pharmaciens) • A. S. Le Duc-Banaszuk (CRDC Pays de la Loire) C. Marjollet (IMAGYN) • J. P. Martin (Ligue Contre le Cancer)

L. Abramowitz • J. L. Brun • C. Clavel • R. Cohen • D. Culié • O. Jourdain • O. Launay • J. Levêque J. B. Lusignan • A. Mesnard • H. Mirghani • J. Nicolet • H. Peré

^{*} Swiss HPV Alliance Initiative