



Registration

Payment is required in order to confirm the registration. No refunds.

Payment to:

European Society of Skin Cancer Prevention – Euroskin e.V.

IBAN: DE02 2075 0000 0090 0469 96 SWIFT-BIC: NOLADE21HAM

Mark: Euroskin Workshop 2018 and your name

Please register before march 1, 2018

Fee: €200 (registration, including 2 lunches and 1 dinner)

Location

Palais des Académies

Rue Ducale 1 1000 Bruxelles

Auditorium (April 16th): Albert II

Auditorium (April 17th): Baudouin and Prirgine



Accommodation

We have reserved rooms at the Bedford Hotel & Conference Center Rue du Midi 135 1000 BRUSSELS, Belgium

Rate : € 100,-, per night, breakfast included.

Reserve before March 1, 2018 via the registration process; use code

The local organizing committee at the Belgian cancer Foundation

Brigitte Boonen (bboonen@fondationcontrelecancer.be)
Jean-Claude Degreef (JCDegreef@fondationcontrelecancer.be)
David Ritchie (david@europeancancerleagues.org)(questions workshop 17th)

For more details on registration, location and accommodation, please visit www.euroskin.eu

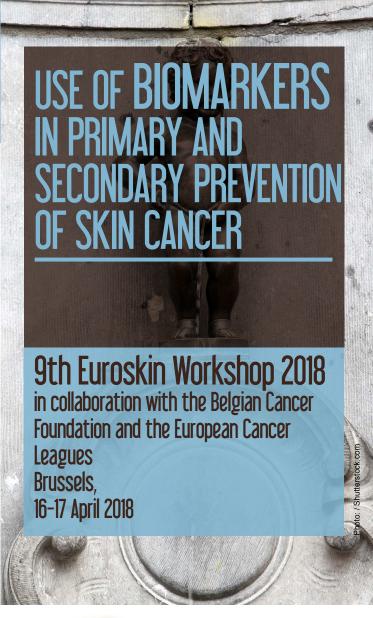
Contact

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USE OF BIOMARKERS IN PRIMARY AND SECONDARY PREVENTION OF SKIN CANCER





Skin cancer is still the most frequent cancer in the white population worldwide. Although our knowledge about risk factors (mainly UV-radiation), etiology and therapy of malignant melanoma (MM), basal cell carcinoma (BCC) and squamous cell carcinoma (SCC) has steadily increased during the last decade, there is still a need for further intensive research. However, it is widely accepted that primary and secondary prevention are powerful tools to avoid skin cancer, detecting persons at risk and/or in early stages of the disease.

For this purpose it will be important to use new knowledge about molecular markers in cells, tissues or body fluids (blood, urine, etc.) which might be indicative for certain risks to suffer from skin cancer in the future or for certain stages of skin cancer development as well as response/resistance to modern therapies. Therefore, the use of biomarkers in the field of skin cancer prevention (and therapy) will increase our possibilities to fight skin cancer.

According to a WHO definition, "a biomarker is any substance, structure or process that can be measured in the body or its products and influence or predict the incidence or outcome of a disease". This has been specified, e.g. by Institute of Medicine (IOM) to "a biomarker is an objectively measured characteristic that describes a normal or abnormal state in an organism by analyzing biomolecules such as DNA, RNA, protein, peptide, and biomolecule chemical modifications". The recent revolution in molecular biology (e.g. high-throughput sequencing, genetic or epigenetic whole genome characterization, etc.) has dramatically increased our possibilities for the search of these biomarkers. Diagnostic, predictive and prognostic biomarkers have been described, as well as biomarkers for certain risk groups and therapy response and resistance. Especially liquid biopsies are increasingly used to characterize e.g. circulating DNA/RNA, tumor cells, exosomes, miRNAs and other non-coding RNAs as biomarkers in human body fluids, which will improve our understanding of different stages of disease risk and progression. Nevertheless, it is important to mention, that biomarker research is heavily dependent on validation of its actual use in prevention and clinical settings.

Because of the tremendous increase in biomarker research and its outstanding potentials in personalized diagnostics and precision medicine, EUROSKIN has invited international experts to give us estimates, results and perspectives about the "Use of Biomarkers in Primary and Secondary Prevention on Skin Cancer". We will discuss their presentation on definitions of biomarkers, examples of biomarker use, epidemiological use of biomarkers, as well as validation of biomarkers and the communication of biomarker measurements. Results of our discussions and suggestions will be published in form of recommendations.

We are hoping that the topics of our conference attract your and your colleague's interest and we are looking forward to welcoming you in the wonderful city of Brussels.

B Boonen Dr R Greinert (President, EUROSKIN) (Secretary General, EUROSKIN)

Monday, April 16th, 2018	
Time	Description
08:00-9:00	Registration
SESSION I: INT	RODUCTION TO BIOMARKERS
9:00-09:45	Use of (epigenetic) Biomarkers (Keynote lecture)
9:45-10:30	Liquid biopsies
10:30–11:00	Coffee break
SESSION II: DIF	FERENT TYPES OF BIOMARKERS IN SKIN CANCER
11:30–12:00	Risk-biomarkers
12:00–12:30	Diagnostic, predictive and progression biomarkers
12:30-13:00	Therapy-(resistance-) biomarkers
13:00-14:00	Lunch
SESSION III: VA	ALIDATION AND EPIDEMIOLOGY
14:00–14:45	Validation of Biomarkers
14:45–15:15	Use of Biomarkers in molecular epidemiology
15:15–15:45	Coffee break
SESSION IV: H	OW TO COMMUNICATE RESULTS OF BIOMARKER USE
15:45–16:15	Differences between primary and secondary Prevention
16:15–16:45	Ethical aspects of communication of (asked and) expected (bad?) medical results
16:45–17:30	Discussion

Confirmed speakers:

- Babayan, A. (G)
- Dupuis, J. (B)
- Gandini, S. (I)
- Greinert, R. (G)
- Hemminki, K. (G)
- Leucci, E. (B)
- Nevns, B. (B)
- Page, C. (NO)
- Van Brabandt, H. (B)

Tuesday, April 17th, 2018		
Time	Description	
SESSION V: EXAMPLES OF BIOMARKER USE IN SKIN CANCER		
09:00-09:30	Liquid biopsies vs Histology	
09:30–10:00	Circulating cf-DNA/cf-RNA	
10:00-10:30	Circulating tumor cells	
10:30–11:00	Coffee break	
11:00–11:30	Methylation analysis as biomarker	
11:30–12:00	Circulating miRNA	
12:00–12:30	circRNA and IncRNA	
12:30-13:00	Formulation of recommendations	
13:00–14:00	Lunch	

Cancer Leagues Workshop Revision of ECL Sun Safety Recommendations		
Time	Description	
13:30	Welcome & Introduction	
13:45	1) What's new? Update of the scientific evidence since 2007	
14:15	2) Update of cancer leagues' activities in sun & UV safety	
14.45	Coffee break	
15.15	3) Presentation and discussion of proposed updated recommendations	
16.30	4) Adoption of the updated recommendations	
16.45	5) Concluding remarks	
17.00	Close of workshop	