

This international conference (now in its 43rd year) is a forum for all workers in human and medical genetics to review advances and develop research collaborations. The conference has become one of the premier events in the field of human genetics with over 2.000 delegates, more than 100 oral presentations, 14 workshops, and 8 educational sessions. The ESHG conference is where the latest developments in human genetics are discussed, and where professionals from all parts of human genetics meet.

Programme

- Invited Plenary lectures and Symposia
- *ESHG Award* and *Mendel* lectures
- "Educational Track" throughout the meeting
- Workshops
- Concurrent Sessions of submitted abstracts
- Poster presentations of submitted abstracts
- Young Scientist and Poster Awards
- Conference Fellowships for young researchers from central and eastern Europe, as well as Fellowships for National Societies are available
- Corporate Satellites
- Over 100 exhibitors from all over the world

Further details

The website www.eshg.org/eshg2012 is now open.

Abstract submission

Online abstract submission via www.eshg.org/eshg2012
Closing date: February 17, 2012

Scientific & Administrative Conference Secretariat

ESHG 2012 c/o Vienna Medical Academy
 Alser Strasse 4, 1090, Vienna, Austria
 Tel: +43 1 405 13 83 16
 Email: conference@eshg.org

Exhibition, Sponsoring, Corporate Satellites

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 P.O.Box 93260, 2509 AG The Hague
 The Netherlands
 Tel: +31 70 383 8901
 Email: eshg@rose-international.com

Further information on programme, registration and abstract submission:

www.eshg.org/eshg2012

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as per date of printing

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The European Society of Human Genetics promotes research in basic and applied human and medical genetics and facilitates contact between all persons who share these aims.

ESHG 2012

EMPAG



Nürnberg, Germany, June 23 – 26, 2012



www.eshg.org

European Human Genetics Conference
 in conjunction with the
European Meeting on Psychosocial Aspects of Genetics
 and the
German Society of Human Genetics (GfH)

Dear Colleagues,

The next ESHG meeting will take place in Nuremberg, Germany. On behalf of the German and the European Societies of Human Genetics I would like to invite you to join us in June 2012. This meeting will continue in the successful tradition of excellent conferences that cover the latest developments in the field of human genetics that are of interest for both clinicians and research scientists.

Nuremberg is a lively city with a medieval feel situated in Southern Germany half-way between Frankfurt and Munich. It is well known for its Christkindlesmarkt, its trade fairs, its world famous beer and sausages, and of course its historical sites and museums.

The meeting will be filled with exciting and up-to-date scientific sessions, educational lectures and distinguished speakers, hopefully making the conference a success both from the scientific as well as from the social point of view.

I hope that you will seize the opportunity to join us in Nuremberg and look forward to seeing you in June 2012.

With best regards,

André Reis

Local Host ESHG 2012
 Chairman of the
 German Society of Human Genetics

The preliminary programme is subject to alteration. Please consult the website as of autumn 2011.

Time	Saturday, June 23	Time	Sunday, June 24	Monday, June 25	Time	Tuesday, June 26
09.00	Registration opens	08.30	S01. The molecular basis of facial malformations S02. Mechanisms & consequences of chromosomal/genetic mosaicism S03 Epigenetics S04. Statistical analysis of sequence data in complex disease ES4. Applying family dynamics/ therapy in genetic counselling	S09. Primary microcephaly S10. Cancer genetics S11. De novo mutations: A common cause of common disease? S12. Molecular basis of Lymphedema ES6. Trinucleotid repeat disorders	09.00	PL3. Targeted pharmacological therapies in genetic disorder
		10.00	Coffee Break	Coffee Break	10.30	Coffee Break
		10.30	Poster Viewing with Authors	Poster Viewing with Authors	11.00	Concurrent Sessions C13 - C18 from submitted abstracts
11.45	Corporate Satellites	11.40	Corporate Satellites WS02. UCSC Genome Browser - Introductory WS03. Array CGH	Corporate Satellites ES7. Next generation sequencing goes diagnostics: First experiences		
14.00	ES1. Complex Diseases ES2. Skin Diseases ES3. How to get published in the EJHG WS01. Cascade screening: what about relatives' right (not) to know? Corporate Satellites	13.15	Concurrent Sessions C01 - C06 from submitted abstracts	Concurrent Sessions C07- C12 from submitted abstracts	12.30	Lunch Break
		14.45	Vitamin Break	Vitamin Break	13.30	Plenary Session 4 Mendel Lecture
15.30	Break	15.15	WS04. Dysmorphology 1 WS05. Community genetics WS06. Genetic Education WS07. Quality assurance WS08. Cellular models of human biology & genetics WS09. Plans & Strategies for Rare diseases in Europe WS10. PGD Workshop Corporate Satellites	WS11. Dysmorphology 2 WS12. Legal regulation of genetic issues WS13. Huntington's disease: how long should we wait for a cure? WS14. Genome research translation & translational research WS15. Progress in understanding of our genome through sharing of data - Debate WS 16. UCSC Genome Browser - Intermediate Tools Corporate Satellites	14.15	Plenary Session 5 EJHG Nature Awards Young Investigator Awards Poster Awards ESHG Award Lecture Closing
15.45	Welcoming Addresses					
16.30	Plenary Session 1					
18.00	Coffee Break	16.45	Coffee Break	Coffee Break		
18.30	Plenary Session 2 What's new? from submitted abstracts	17.15	S05. Advances in Neuropsychiatric Disorders S06. Prenatal diagnosis S07. Genomics and drug response S08. 25 Years Psychosocial research in Clinical Genetics: Making up the balance ES5. Array CGH & Next generation sequencing	S13. Bone diseases & therapeutic perspectives S14. Intellectual disability: genes, proteins & model organisms S15. From DNA sequence to genome function S16. Debate on Informed Consent in New Technologies ES8. Nucleotid repeat disorders	15.45	End of Meeting
		18.45	Corporate Satellites	Corporate Satellites		
		19.00	ESHG Membership Meeting			
20.00	Welcome Reception	20.00		Congress Party		



Abstract submission deadline:
 February 17, 2012