Preliminary Schedule ESHG 2012

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**	**S09. Primary microcephaly** **S10. Cancer genetics** **S11. De novo mutations: A common cause of common disease?** **S12. Molecular basis of Lymphedema**	09.00	**PL3. Targeted pharmacological therapies in genetic disorder**
			ES4. Applying family dynamics/ therapy in genetic counselling	**ES6. Trinucleotid repeat disorders**		
		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**	13.15	**Concurrent Sessions C01 - C06** from submitted abstracts	**Concurrent Sessions C07- C12** from submitted abstracts	12.30	*Lunch Break*
	WS01. Cascade screening: what about relatives' right (not) to know?				13.30	**Plenary Session 4** **Mendel Lecture**
	Corporate Satellites	14.45	*Vitamin Break*	*Vitamin Break*	14.15	**Plenary Session 5** **EJHG Nature Awards** **Young Investigator Awards** **Poster Awards** **ESHG Award Lecture** **Closing**
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**	**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**		
15.45	**Welcoming Addresses**					
16.30	**Plenary Session 1**		**Corporate Satellites**	**Corporate Satellites**		
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**	**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**	15.45	*End of Meeting*
			ES5. Array CGH & Next generation sequencing	**ES8. Nucleotid repeat disorders**		
		18.45	**Corporate Satellites**	**Corporate Satellites**		
		19.00	*ESHG Membership Meeting*			
20.00	*Welcome Reception*	20.00		*Congress Party*		

Plenary Sessions | Concurrent Symposia | Concurrent Sessions | Educational Sessions | Workshops

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Abstract submission deadline: February 17, 2012