

Programme

Time	Topic	Speaker
08:30 - 09:00	Welcome & Registration	
09:00 - 09:10	Introduction	<i>Maria Luisa Brandi</i>
	Understanding lifetime impact of rare disorders in:	Chair: Bernard Cortet
09:10	Skeletal dysplasias – a lifelong transition process	<i>Lothar Seefried</i>
09:30	Achondroplasia	<i>Jörg Semler</i>
09:50	Fibrodysplasia Ossificans Progressiva	<i>Richard Keen</i>
10:10	GACI / ARHR 2	<i>Frank Rutsch</i>
10:30 - 11:00	Coffee Break	
	Understanding genotype-phenotype complexity in:	Chair: Lothar Seefried
11:00	Osteogenesis Imperfecta	<i>Luca Sangiorgi</i>
11:20	X-linked Hypophosphatemia	<i>Peter Kamenicky</i>
11:40	Hypophosphatasia	<i>Lothar Seefried</i>
12:00	Osteopetrosis	<i>Uwe Kornak</i>
12:20 - 13:20	Lunch Break	
	Understanding manifestations and management in:	Chair: Peter Ebeling
13:20	Pregnancy + Lactation associated Osteoporosis	<i>Valentina Degennaro</i>
13:40	Hypoparathyroidism	<i>Maria Luisa Brandi</i>
14:00	Tumor-induced Osteomalacia	<i>Suzanne Jan De Beur</i>
14:20	Early-onset Osteoporosis vs. rare disorders	<i>Ralf Oheim</i>
14:40 - 15:10	Coffee Break	
	Understanding guidelines in:	Chair: Manju Chandran and Radmila Matijevic
15:10	Fibrous Dysplasia / McCune Albright Syndrome	<i>Roland Chapurlat</i>
15:30	X-linked Hypophosphatemia	<i>Aliya Khan</i>
15:50	HPP Diagnostic Criteria	<i>Aliya Khan</i>
16:10	Conclusion & Wrap Up	<i>Lothar Seefried</i>