45th ICBDSR Annual Meeting
Hotel Olšanka, Prague, Czech Republic
September 30 - October 4, 2018

Sunday, September 30, 2018

9:00 - 13:00	Executive Committee meeting
14:00 - 18:00	Meeting of the World Birth Defects Day Participating Organizations *(invitation only)*
16:00 - 18:00	*Registration ICBDSR Annual Meeting*
19:30	*Welcome cocktail at Hotel Olšanka*

Monday, October 1, 2018

8:30 - 8:50	**Welcome**
	Antonin Šípek, Jr., Institute of Medical Biology and Genetics,First Faculty of Medicine Charles University
	prof. Tomáš Zima, Head Institute of Medical Biochemistry and Laboratory Medicine
	prof. Dušek, Head of the Institute of Health Information and Statistics
	Marcia Feldkamp, ICBDSR EC Chair
8:50 - 9:20	Lorenzo Botto, Director ICBD
9:20 - 10:00	J. David Erickson Lecture: Salimah Walani, PhD Vice President of Global Programs
10:00 - 10:10	ICBDSR Distinguished Service Award
10:10 - 10:50	Alessandra Lisi Award Winner Lecture
10:50 - 11:20	Keynote Lecture
11:20 - 11:40	*Coffee break - Networking*
11:40 - 12:40	Session I - Oral communications
12:40 - 14:00	*Lunch*
14:00 - 15:45	Session II - Oral communications
15:45 - 16:45	Session III - Oral communications
16:45 - 17:00	*Coffee break*
17:00 - 18:00	Poster session I
20:00	*Dinner at Hotel Ol šanka, included in the registration fee*

Tuesday, October 2, 2018

8:30 - 10:30	Annual Business Meeting - First part *(ICBDSR members only)*
10:30 -10:45	*Coffee break*
10:45 - 12:30	Annual Business Meeting - Second part *(ICBDSR members only)*
12:30 - 13:30	*Lunch*
13:30 - 15:30	Session IV - Oral communications
15:30 - 16:00	*Coffee break*
16:00 - 17:00	Session V - Oral communications
17:00 - 18:00	Poster session II
19:00	Dinner in the city centre at Municipal House (personal pay)

Wednesday, October 3, 2018

Scientific Session on Rare Defects - Moderator: Dr. Jiří Horáček and Dr. Antonín Šípek Jr

8:30 - 8:40	Introduction
8:40 - 9:20	Rare disease initiatives in the Czech Republic and European ReferenceNetworks 2018. prof. Milan Macek Jr
9:20 - 10:00	Rare Diseases: focus on the rarest and the undiagnosed. Prof Bruno Dallapiccola
10:00 - 10:45	*Coffee break - Networking*
10:45 - 11:15	Screening of Pregnancy complications: prof. Pavel Calda (Head of the Centre of Fetal Medicine, General University Hospital, Prague).
11:15 - 11:40	New methods of chromosomal analysis (microarrays, NIPT): clinical specialist, GENNET, Prague
11:40 - 12:05	Preimplantation genetic testing of monogenic diseases by karyomapping: Dr. Jakub Horak, Rerpomeda center, Brno, Czech Republic
12:05 - 12:35	Neonatal screening – The Czech experience: prof. Viktor Kožich (former head of Institute of Inherited metabolic diseases, General University Hospital, Prague)
12:35 - 14:00	*Lunch*

Classification and Surveillance

14:00 - 14:20	Classification of rare diseases – the local translation of Orphanet classification: dr. Miroslav Zvolský (Institute of health Information and Statistics of the Czech Republic, Prague).
14:20 - 14:40	Czech Registry of Congenital Anomalies: New online reporting system and the implementation of rare diseases reporting tools: Jitka Jirova, MSc. (Institute of health Information and Statistics of the Czech Republic, Prague).
14:40 - 15:10	*Coffee break - Networking*
15:10 - 15:30	Surveillance of rare diseases: UK Experience - Mary Bythell (Head of Rare Disease Registration, NCARDRS)
15:30 - 15:50	Surveillance of rare diseases - the Spanish experience: Eva Bermajo Sanchez
15:50 - 17:00	*Round Table – Discussion about registration and classificationof rare diseases*

Thursday, October 4, 2018

	Optional Day Tour to medieval mine city of Kutna Hora (personal pay)