49th Scientific Session of the International Clearinghouse for Birth Defects Surveillance and Research (ICBDSR) Draft SCIENTIFIC SESSION PROGRAM

	Monday, November 6 th , 2023 Meeting Room Babylon
8:30-9:00	Registration
9:00-9:15	Welcome – Walter Busuttil, Chief Medical Officer, Malta and Boris Groisman, Chair ICBDSR
9:15-9:30	ICBDSR Executive Committee Talk Boris Groisman, ICBDSR Chair and Lorenzo D Botto, ICBD Director
9:30-10:10	J. David Erickson Lecture – Pablo Duran, Pan American Health Organization (PAHO)
10:10-10:30	Keynote Lecture Surveillance research and NTDs prevalence estimates around the world: the importance of the WHA67.19 Resolution Sylvia Roozen, Secretary General, International Federation for Spina Bifida and Hydrocephalus
10:30-10:45	EUROCAT Updates David Tucker, President EUROCAT Association, CARIS, Swansea University, Wales, UK NBDPN Updates–Wendy A Nembhard, Chair of NBDPN, University of Arkansas for Medical
10:45-11:00	Sciences, USA
11:00-11:15	Coffee break
11:15-11:30	From ConcePTION project to sustainable solution for medicines safety assessment during pregnancy and breastfeeding in Europe and beyond Miriam Sturkenboom, Department of Data Science & Biostastistics, University Medical Center Utrecht, Netherlands
11:30-13:00	Session I – Oral Presentations Birth defects surveillance – Methodology I
	Global burden of birth defects estimation: methods, progress and selected results Kathleen L Strong
	Estimating the contribution of different disease groups to the global rare diseases burden using data from the Veneto region RD registry, 2002 through 2019 <i>Monica Mazzucato</i>
	MALFOVAR Registry / Repository Update <i>Tidhar Steiner</i>
	Using population-level, real world data to establish cytogenetic testing patterns and evaluate genotype-phenotype correlations in congenital anomalies in England <i>Corinne R Mallinson</i>
	Current activities of the JRC-EUROCAT Central Registry Agniezska Kinsner-Ovaskainen
	Validating routinely collected data to improve ascertainment of congenital anomalies to support national registration in England Jennifer M Broughan
	Lunch

	Session II – Oral Presentations
	Birth defects surveillance - Descriptive epidemiology
	Prenatal findings in craniofacial microsomia using the Alberta Congenital Anomalies Surveillance System, 1997-2019 <i>Mary Ann Thomas</i>
	Prevalence of skeletal dysplasia in Emilia Romagna, Italy <i>Elisa Ballardini</i>
14:15-15:45	Monitoring an emerging maternal and infant health threat: expanding a birth defects surveillance program to include infants with congenital cytomegalovirus infections <i>Paul A Romitti</i>
	Diaphragmatic hernia in the Czech Republic. Incidence, prenatal diagnosis and sex ratio Antonin Šípek, Jr
	Population-based study on epidemiology and outcomes of Congenital Tracheo-Oesophageal Anomalies in South Wales over a 21-year period <i>Margery Morgan</i>
	Years of potential life lost for children and adults with Congenital Heart Defects, United States, 2007-2017 Wendy N Nembhard
15:45-16:00	Coffee break
10.00.10.15	
16:00-16:45	Poster session (meeting room Olympia)
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	Tuesday, November 7 th , 2023
	Scientific session: Surveillance, management, care and transition to adulthood
	Meeting room Babylon
8:00-8:30	Registration
8.30–10:30	Session I - Surveillance and management Moderators: Neville Calleja, Directorate for Health Information & Research, Ministry for Health, Malta; Charmaine Cordina, Directorate for Health Information & Research, Ministry for Health, Malta
	Surveillance of Congenital Anomalies in Malta - Setting the scene <i>Miriam Gatt</i>
	Neonatal mortality due to congenital anomalies in Malta and beyond <i>Merle Wilhelm</i>
	Evolution of Screening in Malta Mark Cordina
	The accuracy of antenatal ultrasound screening in Malta: a population-based study Jeremy Borg Myatt
	Fetal Cardiac Screening in Malta Sarah Darmanin
	Prenatal Orofacial screening in Malta Maria Christina Tabone
	Orofacial Clefts in Malta: Prevalence, Impact & Public Health Implications – A 20-Year Analysis Terence Micallef
	The epidemiology of congenital heart disease in Malta Victor Grech
	Maternal and Gestational Risk Factors of Congenital Diaphragmatic Hernia Sophie Hackenbruch
10:30-11:00	Coffee break
11:00–13:15	Session II - Care, transition to adulthood and personal experiences Moderators: Victor Grech, Paediatric Department, Mater Dei Hospital, Tal-Qroqq, Malta Miriam Gatt, Directorate for Health Information & Research, Ministry for Health, Malta
	Breaking the news Ryan Farrugia
	Several decades later - a look at long-term outcomes of congenital heart defects in Malta <i>Maryanne Caruana</i>
	The European Reference Networks - Raising the bar for patients with rare diseases and their healthcare providers across Europe <i>James Clark</i>
	Rare Diseases in Paediatrics : Outsourcing the specialised paediatric services into the Community Resource Centres <i>Chris Sciberras</i>
	Adult Down Syndrome Clinic Maria Callus
	Patient / Carer experiences Closing – Yves Muscat Baron – Clinical Chairperson, Department of Obstetrics and Gynaecology
13:15-14:15	Lunch

Posters

(Alphabetical order by corresponding author's last name)

The posters will be shown on Monday 6th November Meeting room "Olympia"

Using national congenital anomaly registration to evaluate the roll out of the non-invasive prenatal testing (NIPT) in England Jennifer M Broughan

Infants with Congenital Anomalies of the Kidneys and Urinary Tract (CAKUT) in Malta *Gabrielle de Gray*

Congenital anomalies as rare diseases: the importance of genomic linkage for congenital anomaly registries *Kathryn Johnson*

The unexplored effects of social and health inequality on adverse outcomes to children born with Spina Bifida: 2010-2020 *Kathryn Johnson*

Sibling birth defects recurrence in Florida, 2000 to 2019 *Russell S Kirby*

Prenatal detection of birth defects: pathway of care in obstetric ultrasound services in Bogotá, Colombia Maria Camila León-Sanabria

Echocardiography evaluation of congenital heart disease with its outcome in a Medical University Hospital of Dhaka, Bangladesh *M Abdul Mannan*

Prevention of neural tube defects in Chile: more than two decades of folic acid fortification *Cecilia Mellado*

WHO's Resolution calls for micronutrient fortification to prevent spina bifida and other Neural Tube Defects *Rosa Pardo*

Epidemiology and risk ractors concerning transposition of the great vessels: A population-based case-control study in Bogotá and Cali, Colombia *Pablo Pineda-Sanabria*

Characterization of risk factors for Clubfoot in Bogotá and Cali, between 2002-2020 *Esteban Portilla*

Polydactyly risk factors in Bogotá and Cali, Colombia between 2002-2020: A case-control study *Lina María Ramírez*

Severe congenital protein C deficiency: epidemiology, diagnosis and long-term management *Csaba Siffel*

Sex ratio and congenital anomalies in children born after assisted reproduction. *Antonin Šípek Sr*

Application of Photon counting CT in the diagnostics of Post mortem fetal malformations for the assessment of skeletal , external skin surface and vasculature anomalies *Tidhar Steiner*

Characterization of environmental exposures in the French ATENA registry *Julire Thomas-Chabaneix*

Multidisciplinary consultation in fetuses with malformations: prenatal exome experience in a tertiary public hospital *Eduardo Tizzano*

Prevalence of Congenital Heart Defects Among Boys with Hypospadias in England *Lee T Watson*