

Assessment of prenatal exposure using human biomonitoring: *the project component 3* 2015-2017

Irina Zastenskaya

WHO European Centre for Environment and Health



Is a global mercury monitoring feasible?

Yes

- Capacities for mercury HBM for reliable assessment of exposure exist or can be built in countries with different level of economic development
- Scientific/Technical/Methodological basis is created
- Applicable for general population and hot spots
- Provide data allowing identification of population groups at risk globally

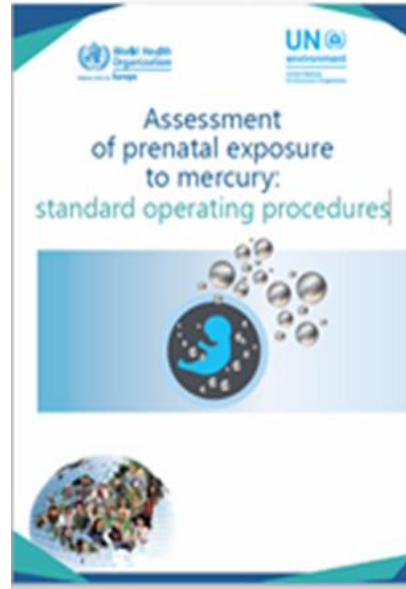
Selection of biological matrix

Biological matrix

Selection of sample matrices and biomarkers for human biomonitoring surveys of prenatal exposure to mercury

Report prepared by Prof. Milena Horvat and Dr. Janja Snoj Tratnik, Jožef Stefan Institute, Ljubljana, Slovenia in collaboration with WHO Europe.

Justification for the selection of biomarkers of prenatal exposure to organic and inorganic mercury



- Scalp hair
- Cord blood
- Urine

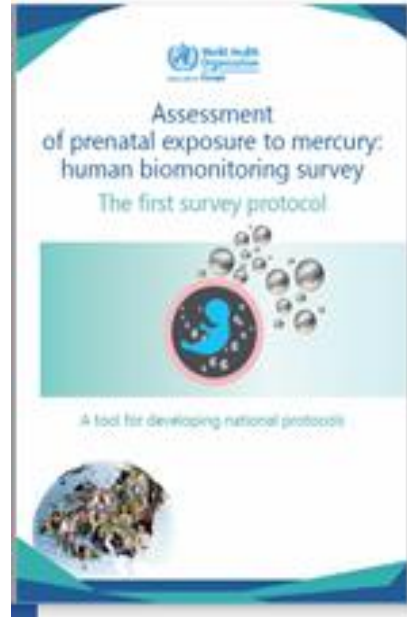
All forms of mercury
All non-invasive

Survey design

Prenatal exposure

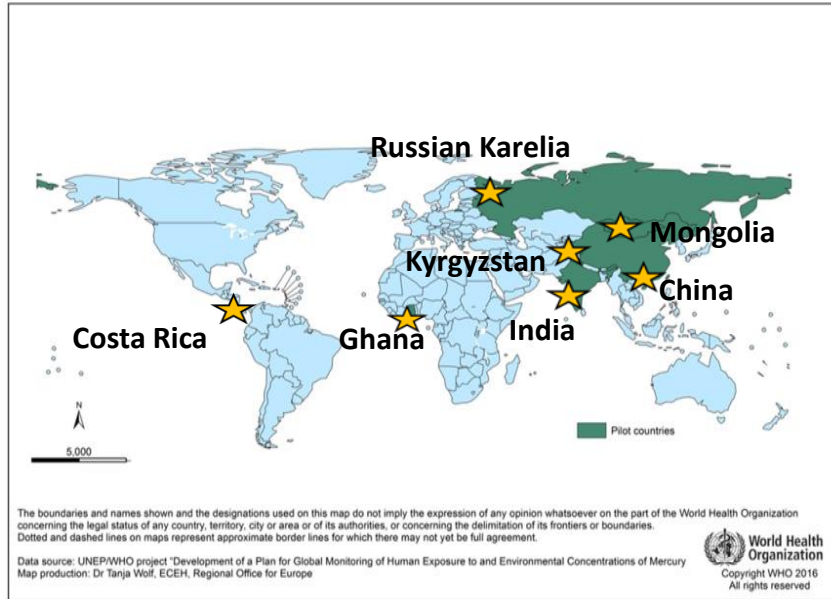
250 women just delivered a child

Maternity hospitals



Harmonized approach to biomonitoring of human exposure to mercury:
International technical experts workshop
UNEP/WHO project "Development of a Plan for Global Monitoring of Human
Exposure to and Environmental Concentrations of Mercury"
Bonn, Germany, 26 June 2015

Pilot survey: Selection of countries



Target populations in the pilot surveys

Seafood consumers –Ghana, India

Freshwaters fish consumers – the Russian Federation

Consumers of locally produced rise in contaminated areas - China

ASGM - Mongolia

Mercury primary mining - Kyrgyzstan

Industrial activities - India

Training of the national survey laboratory analyst (Slovenia, JSI, Feb 2016)



Capacity building is critical for GMM

Pilot surveys

Preparation

- Proficiency test (freeze-drying samples of cord blood)
- Procurement of consumables
- Revision of SOPs, survey protocol
- Subcontracting countries

Pilot surveys:

- ethical approval (local/national),
- Training (for field staff),
- adaptation of WHO documents,
- Sampling (250 samples; three matrices; the epidemiological questionnaire)
- Mercury analysis
- Data analysis
- Database creation

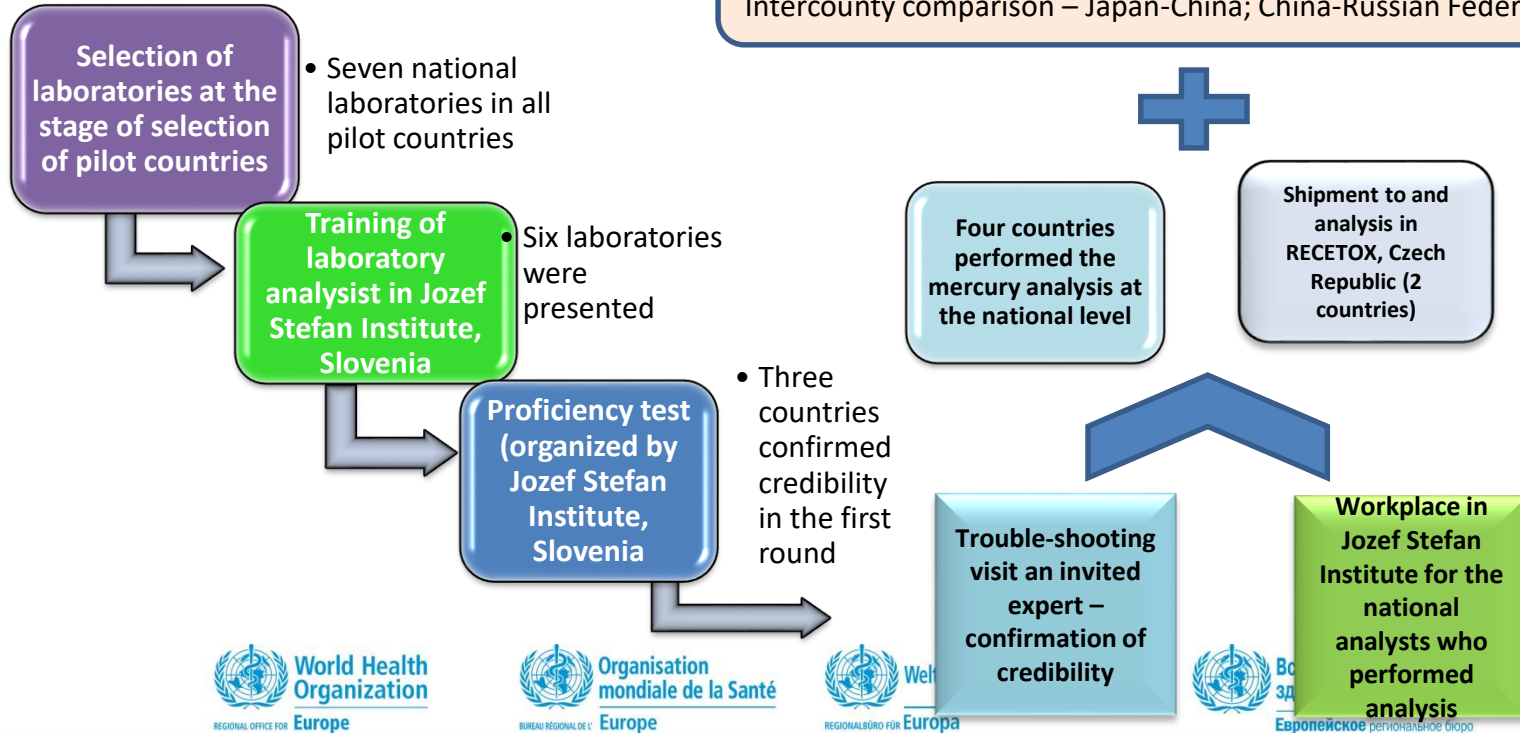
Follow up

- Communication of results



Challenges: Laboratory performance evaluation

Mirror analysis (20 samples of each matrix) from China, India, Kyrgyzstan, Mongolia – National Center for Minamata Disease, Japan;
Intercounty comparison – Japan-China; China-Russian Federation



Challenges: ethics approval

National surveys			
WHO Master Survey Protocol →	<i>Site specific Protocol, China</i>	↔	← National ERC: Institute and Hospital Local governmental???
→	<i>Site specific Protocol, Costa Rica</i>	↔	← National ERC
→	Site specific Protocol, Ghana	↔	← National ERC
→	Site specific Protocol, India	↔	← two institutional ERC: Govt. Hospital & private hospital
→	Site specific Protocol, Kyrgyzstan	↔	← Institutional ERC
→	Site specific Protocol, Mongolia	↔	← National ERC
→	Site specific Protocol, Russian Federation	↔	← Institutional ERC

Database

my cells of "INPUT SHEET". A new dataset will be generated in sheet "TOD_DATA".
Original measurements please fill in measured value. If measurement is below the LOD, please fill in value equal to half the LOD (1/2 LOD).
For each measurement whether value is above LOD (">LOD") or below LOD ("<LOD"), values that are equal to LOD are indicated as "=LOD".

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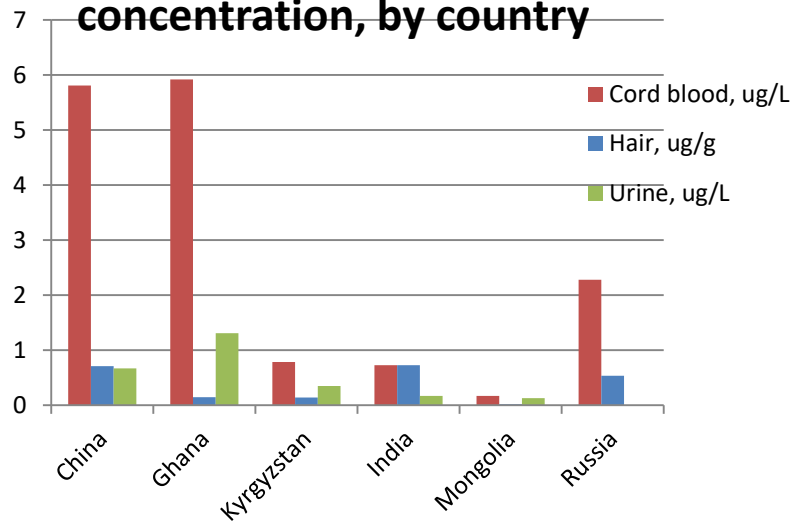
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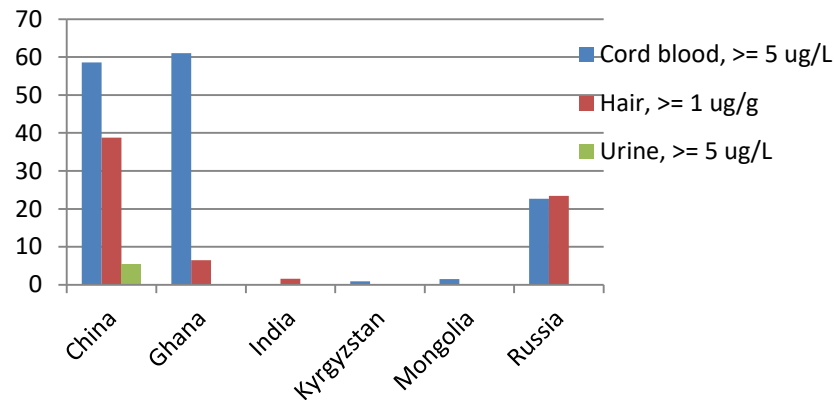
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Mercury concentration

Biomarker mercury concentration, by country



Percent of individuals above reference level, by country*



The survey national coordinators



We are very thankful to:

- The Jozef Stefan Institute, Slovenia (Milena Horvat and the team)
- The National Centre for Minamata Disease, Japan (Mineshi Sakamoto and the team)
- The National Centre for Environmental Health, Carlos III Institute of Health, Spain (Argelia Castaño and the team)
- Research Centre for Toxic Compounds, Czech Republic (Katerina Sebkova and the team)
- WHO SEARO (Lesley Onyon)

THANK YOU

WHO European Centre for
Environment and Health
Platz der Vereinten Nationen 1
53113 Bonn, Germany

