

Perinatal health monitoring in EU+3: data sources

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Aim

- An inventory of existing perinatal health data sources in 26 countries (25 EU + Norway) was done with the 2004 data.
 - Gissler M, Mohangoo A, Blondel B, Chalmers J, Macfarlane A, Gaizauskiene A, Gatt M, Lack N, Sakkeus L, Zeitlin J for the EURO-PERISTAT group: Perinatal health monitoring in Europe: results from the EURO-PERISTAT project. Informatics for Social and Health Care 35 (2): 64-79, 2010.
- The problems in comparisons of perinatal health indicators within Europe will be re-analysed.



Data collection 2010

- Update for 30 countries with 2010 data:
 - No data (yet): Bulgaria, Denmark, Iceland, Greece, Slovak Republic.
 - UK data refers to Scotland, other data pending.
 - Belgian data refers to Brussels, Flanders and Wallonia.
 - Spanish data has also some data on Catalonia and Valencia.



Data sources: 92,

mean 3.7 per country (range 1-7)

A Register-based data sources

- 1. Civil registration based on birth and death certificates, causes-of-death
- 2. Medical Birth Registers/Perinatal Databases, Quality Registers
- 3. Combined MBR and data from civil registration
- 4. Other registers: congenital anomalies, hospital discharges and DRGs, induced abortions, morbidity-specific registers, premature newborns

B Survey data

- 1. Perinatal surveys
- 2. Other routine surveys: breastfeeding and parturients
- 3. Confidential enquiries/audits: maternal deaths, stillbirths, infant deaths

C Aggregate data sources

1. Statistics on ART, induced abortions and miscarriages



Results: Data availability and problems

Questions complicating international comparisons

- 1. Registration criteria
- 2. Coverage of data collection
- 3. Data collection by other definitions than recommended
- 4. Denominators and numerators
- 5. Missing data
- 6. Random variation



92 data sources

- Coverage:
 - Country-level 92%, regional 3%, other 5%
 - Population-based 58%, hospital-based 30%, other 13%
- Data collection:
 - Obligatory 79%, voluntary 21%
 - Death cohort (deaths that occurred in 2010) 60 %,
 birth cohort deaths (babies born in 2010) 40 %,



Registration criteria

- Fetal deaths
 - 32% only gestational age criterion
 - 30% gestational age OR birthweight criterion
 - 15% only birthweight criterion
 - 4% gestational age AND birthweight criterion
 - 19% other criteria
- Livebirths
 - 68 % no
 - 7 % only gestational age criterion
 - 5 % gestational age OR birthweight criterion
 - 2 % only birthweight criterion
 - 18 % other



Discussion

- Improvement since 2004:
 - Strengths and limitations will be studied.
- Main messages:
 - Perinatal health data can be improved everywhere.
 - Indicators have to be developed further, also with other groups
 - International comparisons have to be improved.
 - What is the best way to collect perinatal data?
 - Registers, aggregated statistics, surveys



The way forward

- How to establish a routine collection of perinatal health data?
 - Commission of European Union (DG Health and Consumers or Eurostat)
 - EU Institutes (European Centre for Disease Prevention and Control ECDC)
 - launching a new organisation, such as European
 Perinatal Health Monitoring Centre.
 - Active research networks to promote more research.
 - Collaboration with EUPHA

