

Access to official microdata and accreditation of researchers in Europe

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University of Bucharest, 23 January 2012



Outline

- 1 Background
- 2 Accreditation
 - Eligibility
 - Applications
 - Decision-making
- 3 Similarities, differences, and a way forward
- 4 DwB and today's workshop

European research and data access

- Official microdata as a major resource for science-based policy-making;
- Growing demand for both highly anonymised and detailed datasets;
- Needs of scientific community now often recognised in legal frameworks;
- Ongoing negotiations in the ESS for access to European data;
- Recent national-level improvements in several countries.

However...

- Access arrangements for *national* data still highly heterogeneous;
- Trans-national access difficult or (at best) burdensome;
- Comparative cross-country research particularly penalised.

	1. Public Use Files	2. Scientific Use Files	3. Extracts (subsets)	4. Standard tabulations	5. Bespoke tabulations	6. Secure remote access / execution	7. Safe data labs
a. Census	7-9 countries	10-12 countries	7-9 countries	10-12 countries	10-12 countries	7-9 countries	7-9 countries
b. Social Survey	7-9 countries	10-12 countries	7-9 countries	10-12 countries	10-12 countries	7-9 countries	7-9 countries
c. Person Register	no country	4-6 countries	4-6 countries	7-9 countries	7-9 countries	4-6 countries	1-3 countries
d. Business Survey	1-3 countries	10-12 countries	7-9 countries	10-12 countries	10-12 countries	7-9 countries	10-12 countries
e. Business Register	1-3 countries	4-6 countries	4-6 countries	7-9 countries	7-9 countries	1-3 countries	4-6 countries
f. Other	no country	4-6 countries	4-6 countries	1-3 countries	1-3 countries	1-3 countries	1-3 countries

Table 1. Modes of access in European countries (2008).

no country
 1-3 countries
 4-6 countries
 7-9 countries
 10-12 countries
 ≥ 13 countries

Researcher accreditation

- Accreditation contributes to enabling safe research access to official data;
- It is the process of:
 - defining **eligibility** criteria (who is a researcher, what is a research);
 - establishing **application** procedures (how to request access);
 - designing rules for **decision-making** (who decides, on what basis) and monitoring.

How researcher accreditation contributes to risk management

- A “fit and proper” person;
- Comparable to official statistics staff;
- Safe data = **safe person** / safe project / safe place.

Who is a (safe) researcher?

Different answers in different countries:

- qualifications / experience?
- institutional backing?
- employer / employment status?
- country of birth / residence?
- threat of prosecution / assurance that they will bind themselves to a contract?
- and what about...
 - students?
 - foreign researchers?

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How to apply for accreditation?

Various aspects and options:

- Who submits an application —PI, team, institution?
- Whose signatures are needed —PI, team, institution?
- What forms to use —and are they available in different languages?
- What evidence to provide?
- ...

Yet commonalities exist!

- Much of the variation is in practices and processes rather than principles;
- In fact, most of the existing procedures attempt to capture the same information!
- There are similarities in key criteria and application contents, though they often go under different names.

⇒ Is there scope for improvement?

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Our task within DwB

- Map current accreditation criteria, rules, procedures and practices across Europe;
- Identify commonalities that may help to design a better integrated future system;
- Also, identify major obstacles to the process, if any.

Today's workshop

- A forum to overview accreditation rules, practices and procedures in Eastern Europe;
- Understand current state, planned changes, and challenges ahead;
- Identify opportunities and threats, and the needs and expectations of all countries represented;
- Discuss how DwB may contribute to improving access under safe conditions in Eastern Europe.

Thank you!

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