

Who are the BIRO Users

BIRO Collaboration



BIRO users

- Governance
- Research
- Health care
- Citizens

Governance

- European Union
 - Commission, Parliament, Directorates, Web Portal
- National and Regional Governments
 - Local Health Care Authorities (Management Clinical Networks)
 - Other local authorities
- Payers
 - Social /Private Insurance
- Non Governmental Organizations (NGOs)
 - WHO, OECD, IDF, National and local Diabetes Associations (Diabetes Atlas)

Research

- EU DG-Research, DG-INFSO
- Scientific Organizations
 - e.g. National and international scientific organizations (EASD, ESF)
- Research institutions
 - e.g. Universities, Foundations, Statistical Departments of Local Governments
- Research areas
 - e.g. Epidemiology, health policy, public health, clinical medicine

Health care

- Primary Care Societies
- Diabetes Care Units
- Health Care Professional Associations
- Quality Management Associations
- Industry

Citizens

- Consumer Organizations
- Patients organizations

Using the list of users

- Classification of users (monitoring usage)
- Customization (report template development)
- Criteria to be used to register in the web
- Users Log (find category from IP numbers)
- Relate to external stakeholders
- Put list on the web to target users

Conclusions from the list of users

- Only language: english (meta-data able to allocate translation, any initiative should be regulated according to the BIRO Consortium, rules to be defined)
- Interface for single GP/patient beyond the scope (possible, but should be planned and properly funded)
- Users should register “at some stage”
- Consider any possible abuse of information in the PIA, even beyond the expiration of the EU contract

Refining the list

- Put list on the BIRO Forum under Dissemination
- Prompt feedback from partners by the time of Graz meeting
- Use list to inform/drive report template discussion