



The Health Information Portal: a one-stop shop for health information

Miriam Saso, Sciensano European Public Health Conference, Dublin

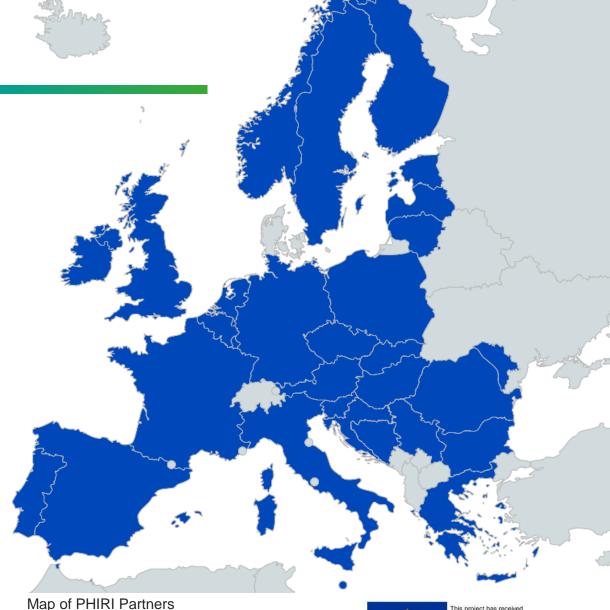




PHIRI

The Population Health Information Research Infrastructure:

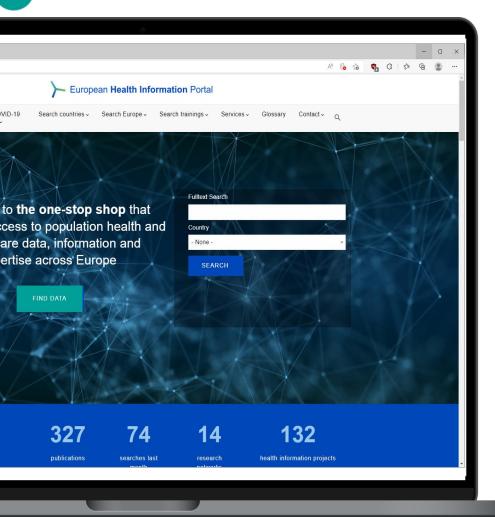
- a European mechanism, that aims to
- facilitate and support data-driven population health research
- and exchange of best practices
- to support decision making





The Health Information Portal





www.healthinformationportal.eu

A one-stop shop that facilitates access to population health and health care data, information and expertise across Europe.



Data sources



Publications



Countries and national nodes



Trainings in all areas of population health



Research infrastructures, Research networks



COVID-19 Policy measures



Health information projects



COVID-19 Rapid Exchange Forum

Data sources metadata catalogue





OVERVIEW	DEMOGRAPHICS	DATASET	CONTACT
Acronym	MNCR		
Type of	Registry data		
information			
URL of the	data source		
https://deput	yprimeminister.gov.i	mt/en/dhir/Pa	ges/Registries/can
cers.aspx			
Description			
The aims of	this register are 1)Te	o collect data	, keep a register
and follow u	p all residents of the	Maltese Isla	nds who are

diagnosed with cancer, and 2)To provide a surveillance function

It provides structured information on

- Population registries
- Health survey data
- Surveillance data
- Hospitalizations (in- and out-patient)
- Healthcare resources, costs and activities
- Etc.

The metadata template is mapped to **Schema.org and to DCAT metadata standards**, further increasing the findability of the data on the web





Sources on the Health Information Portal



CATEGORY

- □ Data / Hi Source (20)
- □ Output & Dissemination (8)
- ☐ European Research Network (5)
- □ Capacity Building Activity (1)
- □ National Node (1)

Fulltext Search

health interview

You can use boolean operators like AND, OR and NOT (in capital letters) to narrow your search.

Country

International

SEARCH

RESET

SEARCH RESULTS LEGEND



National Node



European Research Network



Data / Hi source



Output & Dissemination



HEALTH INTERVIEW SURVEY Topics Determinants of health; Health status Free keywords household; general population; home visit; lifestyle Type of information Survey/interview data Website https://his.wiv-isp.be/SitePages/Home.aspx Country(ies):









- Type of Information
- URL of the data source
- Description
- Governance & legal framework
- Keywords & topics

OVERVIEW	DEMOCDADHICS	DATASET	CONTACT
OVERVIEW	DEMOGRAPHICS	DATASET	CONTACT

HIS Acronym

Type of information Survey/interview data

URL of the data source

https://www.sciensano.be/en/projects/health-interview-survey

Description

A national Health Interview Survey (HIS) provides an overall picture of the health status of the population and allows to identify the main health problems, as well as the social and behavioral factors that influence them. This information enables health authorities to pursue a proactive health policy aimed at improving public health and addressing the peeds of groups at rick. Desceptors, expects and the processor the recults of the LUC.





OVERVIEW DEMOGRAPHICS DATASET CONTACT

Country(ies) Belgium

GEO coverage Nuts 1

Target Population General population

Sex Both





- Data collection period
- Languages
- Personal identifier
- Level of aggregation
- Data access

OVERVIEW

DEMOGRAPHICS

DATASET

CONTACT

Data Collection Period

Language(s) Dutch, English, French

Personal Identifier Participant number

Level of aggregation Individual

Terms of data access - URL https://www.sciensano.be/en/node/55737/health-interview-survey-microdata-request-procedure

Terms of data access

The procedure for obtaining the HIS data is initially done by completing the electronic application form.





- Data Owner
- Contact details

OVERVIEW DEMOGRAPHICS DATASET CONTACT

Data Owner(s) Sciensano

Contact name HIS team Sciensano

Contact e-mail his@sciensano.be

Contact info (address)



Sciensano

Rue Juliette Wytsmanstraat 14

1050 Brussels

Relaium

Supporting data-driven research





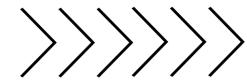
What type of data are these?

What age groups do they cover?

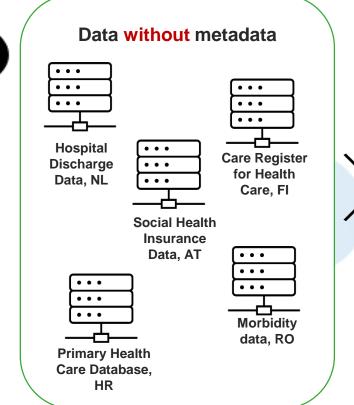
Who owns these data?

 How do I get access to these data sources?

For my research I need data on prevalence of NCD's in European countries. Where to find it?



Through my networks and internet search I was able to identify the following potential data sources





Why metadata?





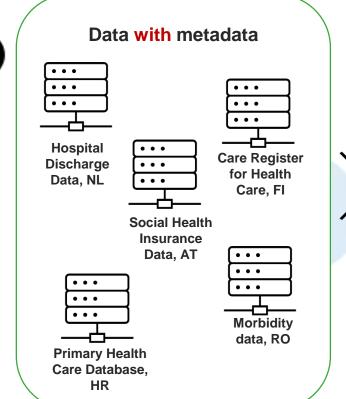
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Through the Health
Information Portal I was
able to identify the following
potential data sources



In structured format

- Data on hospitalizations
- Covers all 18+
- National coverage
- Contact details for access requests

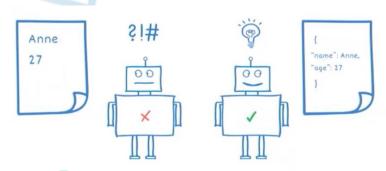


Metadata on the Health Information Portal



Development process of the metadata elements:

- 1) Metadata elements developed by population health researchers
 - In order to accomodate all metadata relevant for population health information
- 2) Mapping to schema.org
 - Schema.org is a metadata standard for the indexation of web pages
 - Improves machine to machine communication
 - Increases discoverability





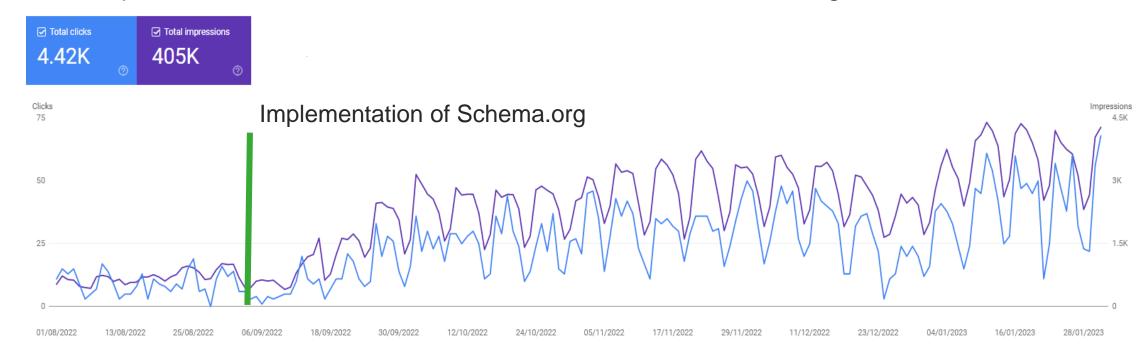


Effects of Schema.org



The Health Information Portal - Google search analytics:

- Clicks: number of times a user clicked on a link provided by Google to the HIP.
- Impressions: how often someone saw a link of the HIP on Google.







Metadata on the Health Information Portal



Development process of the metadata elements:

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- 2) Mapping to schema.org
 - Schema.org is a metadata standard for the indexation of web pages
- 3) Mapping to DCAT
 - DCAT is a vocabulary for publishing data catalogs on the web
 - Facilitates sharing of data & interoperability between data catalogs
 - Increases the discoverability of datasets
 - Allows federated search for datasets across catalogs in multiple sites





Metadata in the European landscape



Proposal for a regulation - The European Health Data Space published on May 3rd by the European Commission.

Section 5 – Article 55.1

"The health data access bodies shall inform the data users about the available datasets and their characteristics through a **metadata catalogue**. Each dataset shall include information concerning the source, the scope, the main characteristics, nature of electronic health data and conditions for making electronic health data available"

- The Health Information Portal has these metadata available in a standardized format
- →The Health Information Portal can provide a starting point for the national authorities to create a catalogue
- →Through an API these metadata are available for everyone







Thank you!

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