



Gene Wiki: Making data FAIR on Wikidata

Andra Waagmeester,

Micelio, Antwerp, Belgium | Email: andra@micelio.be, Twitter: @andrawaag

Micelio, Ekeren, Antwerp, Belgium



Structure

- Introduction
 - Wikidata
 - Wikibase
 - Gene Wiki
- Bits and Bolts
 - How to get your data into Wikidata/Wikibase
 - Using Wikidata/Wikibase
- Is Wikidata FAIR?
 - Where is it located in the research life cycle
 - How FAIR is Wikidata/Wikibase/Genewiki
 - interoperate/collaborate/engage
 - one key change to successfully implement FAIR data infrastructure

Wikidata and its sisters

Wikidata is to data as Wikipedia is to text

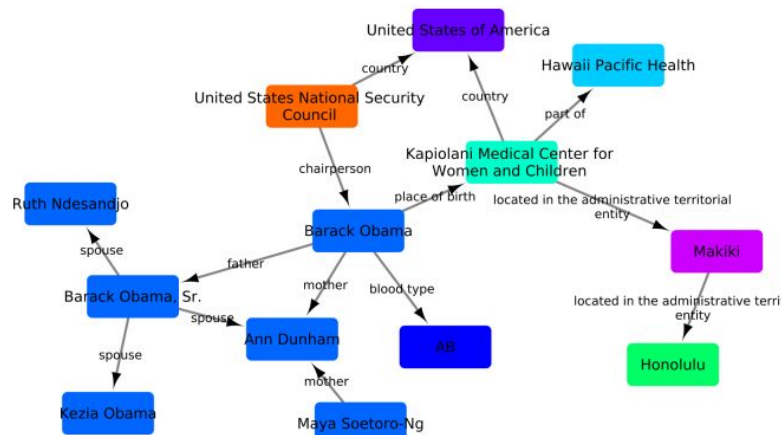
Wikidata is a collaboratively edited knowledge base operated by the Wikimedia Foundation



- Completely free, even for commercial usage (CC0)
- Anybody can contribute
- Covers all domains of knowledge
- Extensive item history, talk pages, projects, users
- Integration with the semantic web
- High performance query engine (SPARQL)

- Stable! Long term support not dictated by funding cycles
- Actively developed
- Already has large number of active users, editors, contributors!

A giant graph of knowledge!



FAIR on Wikidata The Wikimedia infrastructure

Infrastructure



Resource



Content

Data

Where?

<https://www.wikidata.org>

<https://www.wikibase.org>



Text

<https://<lang>.wikipedia.org>

<https://releases.wikimedia.org/mediawiki/>



Under development

Schema

<https://shex.io>

<https://wikidata-shex.wmflabs.org/>

FAIR on Wikidata: Providing identifiers

Wikidata as an intuitive resource towards semantic data modeling in data FAIRification

Annika Jacobsen¹[0000-0003-4818-2360], Andra
Waagmeester²[0000-0001-9773-4008], Rajaram
Kaliyaperumal¹[0000-0002-1215-167X], Gregory S. Stupp³[0000-0002-0644-7212],
Lynn M. Schriml⁴[0000-0001-8910-9851], Mark Thompson¹[0000-0002-7633-1442],
Andrew I. Su³[0000-0002-9859-4104], and Marco Roos¹[0000-0002-8691-772X]

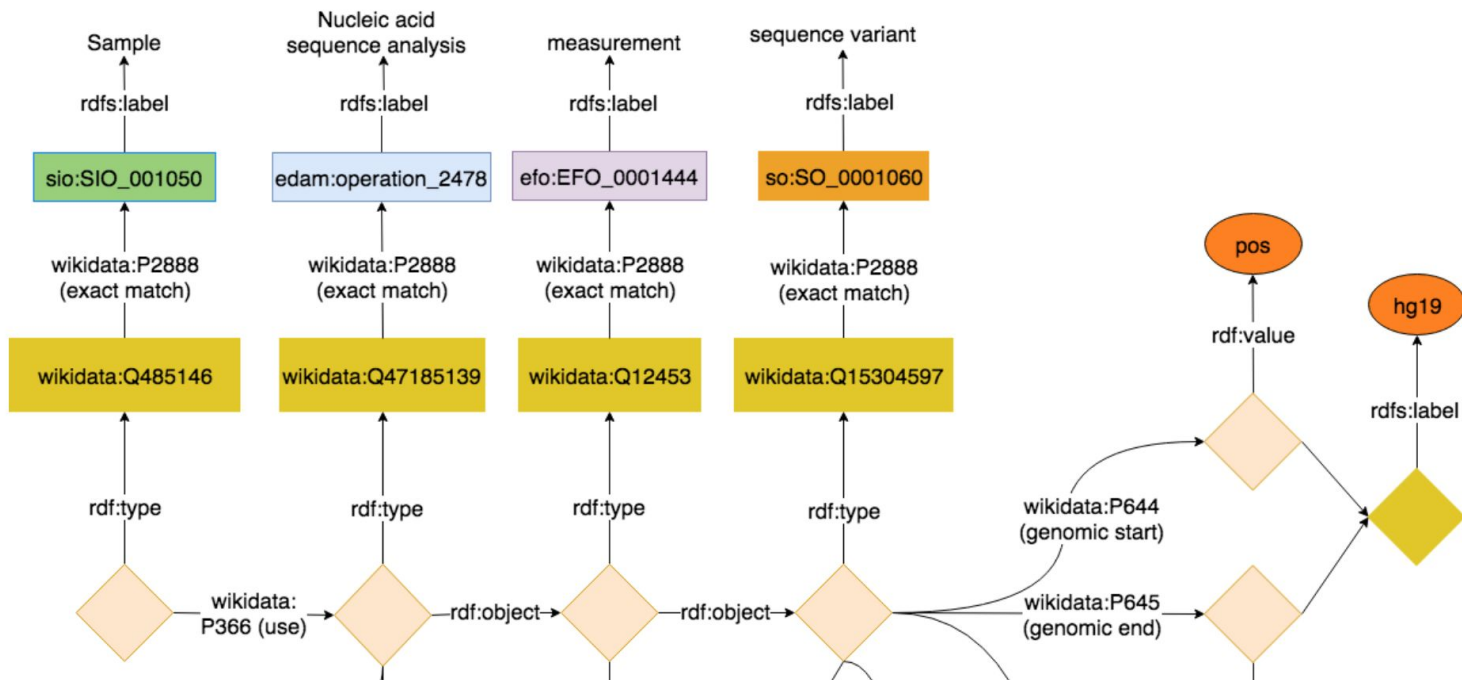
¹ Department of Human Genetics, Leiden University Medical Center, Leiden, The
Netherlands

² Micelio, Antwerp - Ekeren, Belgium

³ The Scripps Research Institute, San Diego CA 92037, USA

⁴ University of Maryland School of Medicine, Baltimore, MD, USA

FAIR on Wikidata: Providing identifiers



W Reelin - Wikipedia

Secure | https://en.wikipedia.org/wiki/Reelin

☆ ... P U B Y < > G A S :

Andrawaag 21 99+ Talk Sandbox Preferences Beta Watchlist Contributions Log out

Read Edit View history More

Search Wikipedia



WIKIPEDIA
The Free Encyclopedia

Main page

Contents

Featured content

Current events

Random article

Donate to Wikipedia

Wikipedia store

Interaction

Help

About Wikipedia

Community portal

Recent changes

Contact page

Toots

What links here

Related changes

Upload file

Special pages

Permanent link

Page information

Wikidata item

Cite this page

Print/export

Create a book

Download as PDF

Printable version

Article

Tabl

Reelin

From Wikipedia, the free encyclopedia

Reelin (RELN)^[5] is a large secreted **extracellular matrix glycoprotein** that helps regulate processes of **neuronal migration** and positioning in the developing brain by controlling **cell-cell interactions**. Besides this important role in early **development**, reelin continues to work in the adult brain. It modulates **synaptic plasticity** by enhancing the induction and maintenance of **long-term potentiation**.^{[6][7]} It also stimulates dendrite^[8] and **dendritic spine**^[9] development and regulates the continuing migration of **neuroblasts** generated in **adult neurogenesis** sites like **subventricular** and **subgranular zones**. It is found not only in the **brain**, but also in the **spinal cord**, **blood**, and other body organs and tissues. ^[*citation needed*]

Reelin has been suggested to be implicated in pathogenesis of several brain diseases. The expression of the protein has been found to be significantly lower in **schizophrenia** and psychotic **bipolar disorder**,^[10] but the cause of this observation remains uncertain as studies show that **psychotropic medication itself affects reelin expression**. Moreover, epigenetic hypotheses aimed at explaining the changed levels of reelin expression^[11] are controversial.^{[12][13]} Total lack of reelin causes a form of **lissencephaly**. Reelin may also play a role in **Alzheimer's disease**, **temporal lobe epilepsy** and **autism**.^[*citation needed*]

Reelin's name comes from the abnormal reeling **gait** of **reeler** mice,^[14] which were later found to have a deficiency of this brain **protein** and were **homozygous** for mutation of the RELN gene. The primary phenotype associated with loss of reelin function is a failure of neuronal positioning throughout the developing **central nervous system** (CNS). The mice **heterozygous** for the reelin gene, while having little neuroanatomical defects, display the **endophenotypic** traits linked to psychotic disorders.^[15]

Contents [hide]

1 Discovery

2 Tissue distribution and secretion

3 Structure

4 Function

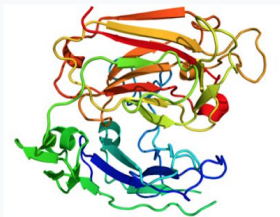
4.1 During development

4.2 In adults

5 Evolutionary significance

6 Mechanism of action

RELN



Available structures

PDB

Ortholog search: PDBe RCSB

List of PDB id codes

[show]

Identifiers

Aliases

RELN, LIS2, PRO1598, RL, reelin, ETL7

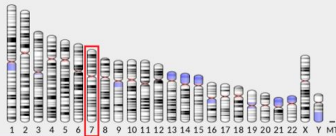
External IDs

OMIM: 600514 MGI: 103022 HomoloGene: 3699

GeneCards: RELN

Gene location (Human)

[hide]



Chr.

Chromosome 7 (human)^[1]

https://en.wikipedia.org/wiki/Reelin#Psychotropic_medication



Read

[View history](#)

More ▾

Search Wikidata



mammalian protein found in Homo sapiens

RELN | reelin | uniprot:P78509

► In more languages

instance of

protein

► 1 reference

+ add value

subclass of

protein



► 1 reference

Reelin



► 1 reference

+ add value

image

2DDU.png

 edit

► 1 reference

- [Main page](#)
- [Community portal](#)
- [Project chat](#)
- [Create a new item](#)
- [Recent changes](#)
- [Random item](#)
- [Query Service](#)
- [Nearby](#)
- [Help](#)
- [Donate](#)

Tools

- What links here
- Related changes
- Special pages
- Permanent link
- Page information
- Concept URI
- Cite this page
- Import interwiki

Retinoic acid receptor alpha (Q254943)

mammalian protein found in Homo sapiens
Nuclear receptor subfamily 1 group B member 1 | RARA

Statements

molecular function

molecular function (P680)
represents gene ontology function annotations

Wikipedia (7 entries)

ar

مستقبل حمض الريتينويك ألفا

en

Retinoic acid receptor alpha

es

Receptor de ácido retinoico alfa

sh

Receptor retinoinske kiseline alfa

sr

Receptor retinoinske kiseline alfa

uk

RARA

zh

视黄酸受体α

retinoic acid binding (Q14901431)
Interacting selectively and non-covalently with retinoic acid, 3,7
GO:0001972

Statements

subclass of

retinoid binding

1 reference

retinoic acid binding

determination method

1 reference

retrieved

stated in

3 January 2017

A human retinoic acid receptor which belongs to the family of nuclear receptors

UniProt-GOA

British Heart Foundation

http://www.ebi.ac.uk/QuickGO/GA
nnotation?protein=P10276

IDA

transcription corepressor activity

determination method

IDA

1 reference

IDA (Q23174122)
Gene Ontology evidence code
Inferred from Direct Assay

Statements

instance of

Gene Ontology Evidence code

manual assertion

A human retinoic acid receptor which belongs to the family of nuclear receptors (Q24339631)

Statements

instance of

scientific article

Identifiers

PubMed ID

2825025

British Heart Foundation (Q4970039)

Statements

instance of

organization

official website

http://www.bhf.org.uk/

Identifiers

GRID ID

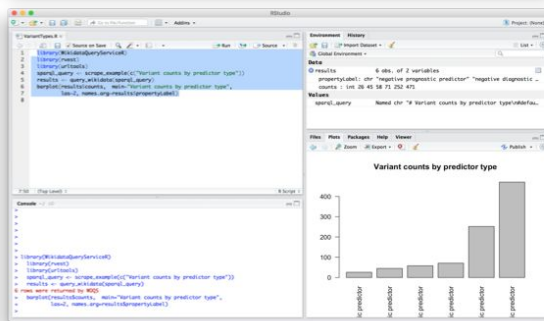
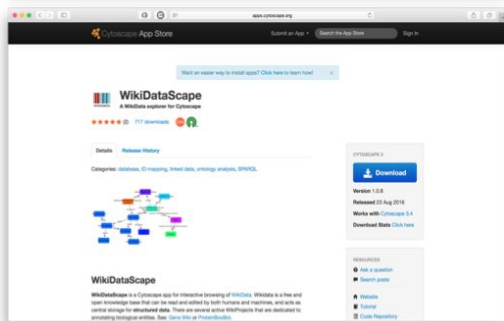
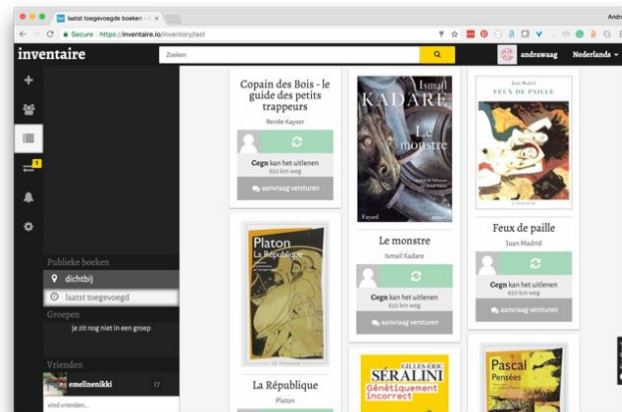
grid.452924.c

Tools using Wikidata

<http://www.wikigenomes.org> Wikipedia and Wikidata
google plugin



<http://inventaire.io>



Cytoscape

R plugins

Wikidata and its sisters

The Gene Wiki project, circa 2008

Summarized
knowledge via
crowdsourcing

IL2-inducible T-cell kinase

Function

This gene encodes an intracellular tyrosine kinase expressed in T-cells. The protein is thought to play a role in T-cell proliferation and differentiation.^{[2][3]}

Structure

This protein contains the following domains, which are often found in intracellular kinases:^[4]

- N-terminus – PH (pleckstrin homology domain)
- ITK – Bruton's tyrosine kinase Cys-rich motif
- SH3 – (Src homology 3)
- SH2 – (Src homology 2)
- C-terminus – tyrosine kinase, catalytic domain

Interactions

ITK (gene) has been shown to interact with FYN,^{[5][6]} Wiskott-Aldrich syndrome protein,^{[7][8]} KHDRBS1,^{[8][9][10]} PLCG1,^{[10][11]} Lymphocyte cytosolic protein 2,^{[11][12]} Linker of activated T cells,^{[12][13]} Karyophormin alpha 2,^[14] Grb2^{[5][9]} and Peptidylprolyl isomerase A.^[15]

References

- Gibson S, Leung B, Squire JA, Hill M, Arima N, Goss P, Hogg D, Mills GB (September 1993). "Identification, cloning, and characterization of a novel human T-cell-specific tyrosine kinase located at the hematopoietin complex on chromosome 5q". *Blood* **82** (5): 1561–72. PMID 8354205.
- Kosaka Y, Felices M, Berg LJ (October 2006). "Itk and Th2 responses: action but no reaction". *Trends Immunol.* **27** (10): 453–60. doi:10.1016/j.it.2006.08.008. PMID 16931156.
- "Entrez Gene: ITK (IL2-inducible T-cell kinase)".
- Hawkins J, Marcy A (July 2001). "Characterization of Itk tyrosine kinase: contribution of noncatalytic domains to enzymatic activity". *Protein Expr. Purif.* **22** (2): 211–9. doi:10.1006/prep.2001.1447. PMID 11437596.
- Bunnell, S.C.; Diehn M; Yaffe M B; Findell P R; Cantley L C; Berg L J (Jan. 2000). "Biochemical interactions integrating Itk with the T cell receptor-initiated signaling cascade". *J. Biol. Chem. (UNITED STATES)* **275** (3): 2219–30. ISSN 0021-9258. PMID 10639929.
- Intramolecular association in a tyrosine kinase of the Tec family. *Nature (ENGLAND)* **385** (6611): 93–7. doi:10.1038/385093a0. ISSN 0028-0836. PMID 8985255.
- Perez-Villar, J. J; Kanner S B (Dec. 1999). "Regulated association between the tyrosine kinase Emh1/Tsk and phospholipase-C gamma 1 in human T lymphocytes". *J. Immunol. (UNITED STATES)* **163** (12): 6435–41. ISSN 0022-1767. PMID 10580033.
- Shim, Eun Kyung; Moon Chang Suk; Lee Gi Yeon; Ha Yun Jung; Chae Suhm-Ke; Lee Jong Ran (Sep. 2004). "Association of the Src homology 2 domain-containing leukocyte phosphoprotein of 76 kD (SLP-76) with the p85 subunit of phosphoinositide 3-kinase". *FEBS Lett. (Netherlands)* **575** (1–3): 35–40. doi:10.1016/j.febslet.2004.07.090. ISSN 0161-6759. PMID 15388330.
- Shan, X; Wang R L (Oct. 1999). "Itk/Emt/Task activation in response to CD3 cross-linking in Jurkat T cells requires ZAP-70 and Lat and is independent of membrane recruitment". *J. Biol. Chem. (UNITED STATES)* **274** (41): 29323–30. ISSN 0021-9258. PMID 10501192.
- Perez-Villar, Juan J; Whitehead-Boake, Simon

IL2-inducible T-cell kinase

Available structures

1ku, 1lu, 1um, 1un, 1sn2, 1snu, 1snx, 2eltz, 2evd

Identifiers

symbols ITK; PSCTK2; EMT; LYK; MGC128257; MGC128258

external IDs OMIM: 169873; MGI: 96621; HomoloGene: 4051

GeneCards: ITK Gene

number 2.7.10.2

Gene ontology [show]

RNA expression pattern

21199_z_at

Protein domains

Orthologs

species	Human	Mouse
entrez	3702	16428
ensembl	ENSG000000113263	ENSMUSG000000202095
UniProt	Q08881	A1A560
RefSeq	NM_005546	NM_010583
RefSeq	NP_065537	NP_034713

Data imported
from structured
databases

Reelin

From Wikipedia, the free encyclopedia

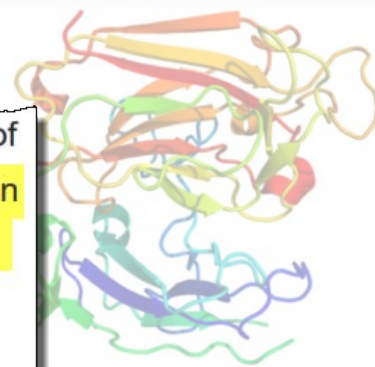
Reelin is a large secreted [extracellular matrix glycoprotein](#) that helps regulate processes of [neuronal migration](#) and positioning in the developing brain by controlling [cell–cell interactions](#).

Besides this important role in early [development](#), reelin continues to work in the adult brain. It modulates [synaptic plasticity](#) by ^{[2][3]} It also stimulates dendrite^[4] migration of [neuroblasts](#) generating [zones](#). It is found not only in the [tissues](#).

Reelin has been suggested to be implicated in pathogenesis of several brain diseases. The expression of the protein has been found to be significantly lower in schizophrenia and psychotic bipolar disorder, but the cause of this observation remains uncertain as studies show that [psychotropic medication itself affects reelin expression](#). Moreover, epigenetic hypotheses aimed at explaining the changed levels of reelin expression^[6] are controversial.^{[7][8]} Total lack of reelin causes a form of [lissencephaly](#). Reelin may also play a role in [Alzheimer's disease](#), [temporal lobe epilepsy](#) and [autism](#).

Reelin's name comes from the abnormal reeling [gait](#) of *reeler* mice,^[9] which were later found to have a deficiency of this brain [protein](#) and were [homozygous](#) for mutation of the RELN gene. The

Reelin



3D ribbon structure of the third reelin repeat domain.^[1]

Available structures

PDB Ortholog search: [PDBe](#) [RCSB](#)

List of PDB id codes [\[show\]](#)

Identifiers

Symbols [RELN](#) ; [LIS2](#); [PRO1598](#); [RL](#)

External [OMIM: 600514](#) [MGI: 103022](#)

Chlambase.org and Wikigenomes.org

 WikiGenomes

Organism Search:

[Authorize to Edit Wikidata](#)

Helicobacter pylori 26695

Organism

Annotation Landscape

Proteins
1422

Genes
1415

NCBI Taxonomy ID:
85962

Wikidata Item ID:
Q21065231

0100,000200,000300,000400,000500,000600,000700,000800,000900,0001,0001,100,0001,200,0001,300,0001,400,0001,500,0001,600,000

NC_000915.1NC_000915.1:67020..77817 (10.8 Kb)Go

70,00072,50075,00077,500

Select tracks

hypothetical protein HP0062

protein HP0061

HP0063hypothetical protein HP0063

HP0064ATP-binding protein HP0066

hypothetical protein HP0064

HP0065hypothetical protein HP0065

HP0067urease accessory protein UreH HP0067

HP0068urease accessory protein UreG HP0068

HP0069urease accessory protein UreF HP0069

HP0070urease accessory protein UreE HP0070

HP0071acid-activated urea channel HP0071

HP0072urease subunit beta HP0072

HP0073urease subunit alpha HP0073

Operons

Gene Search:

Gene

Protein

Gene Name:
urease accessory protein UreH HP0067

Entrez ID:
899986

Wikidata ID:
Q21629448

NCBI Locus Tag:
HP0067

Molecular Function (2)Biological Process (2)Cellular Component (1)InterPro Domains (0)Enzyme Activity (0)Operon (0)Genomic Position

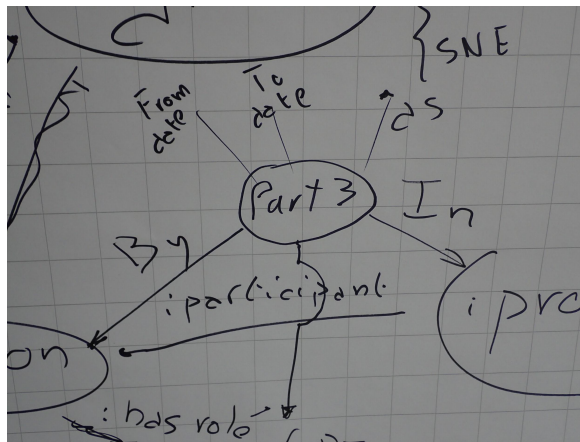
Molecular Function	Gene Ontology ID	Evidence
protein binding	GO:0005515	IPI
nickel cation binding	GO:0016151	IEA



FAIRification

Community engagement and model discussion

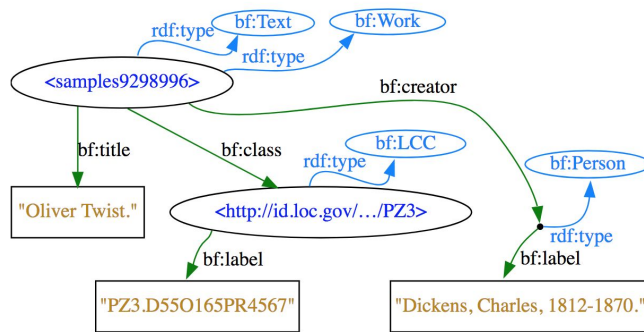
Data
Integration
Extension for
Grants
Ontology



Formally capture and describe model and community consensus

Model development

- Legacy review – develop punch lists for existing data issues that needs fixing
- Documentation – terse, human-readable representation helping contributors and maintainers quickly grok the model
- Client pre-submission – submitters test their data before submission to make sure they're saying what they want to say and that the receiving schema can accommodate all of their data
- Server pre-ingestion – submission process checks data as it comes in and either rejects or warns about non-conformant data

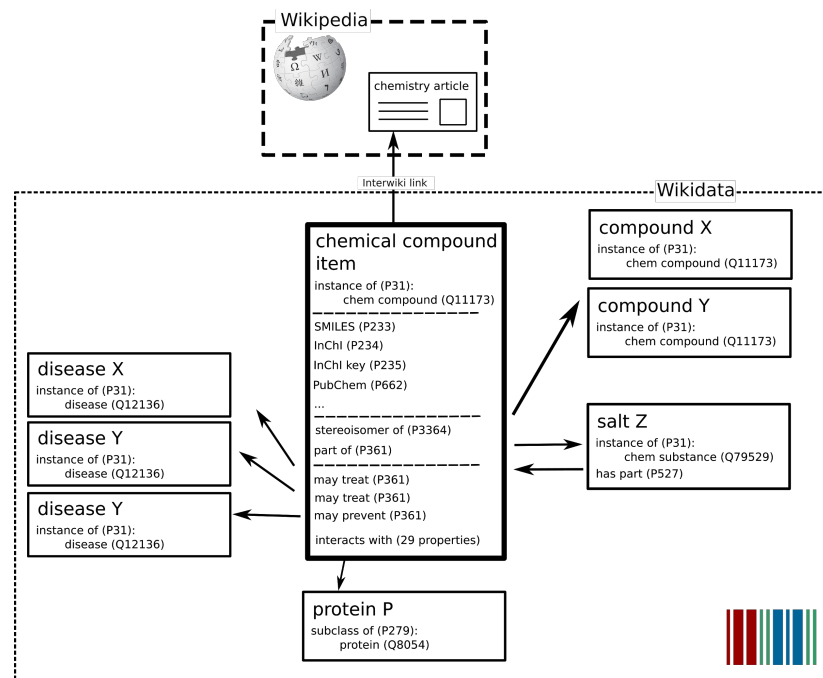


```
Data (Turtle)
<samples9298996>
  rdf:type bf:Text ;
  rdf:type bf:Work ;
  bf:title "Oliver Twist." ;
  bf:class <id.loc.gov/.../PZ3> ;
  bf:creator [
    rdf:type bf:Person ;
    bf:label "Dickens, Charles, 1812-1870." ;
  ] .

<id.loc.gov/.../PZ3>
  rdf:type bf:LCC ;
  bf:label "PZ3.D55O165PR4567" .
```

Seeding with data

- Model structure of items (genes, drugs, diseases, .. etc) & relationships between items
- Import data from many sources and ontologies
- Linked to many identifiers from external databases
- Architecture for maintaining data from external sources



Platforms for (semi-) automatic data ingestion

- Quickstatements: <https://tools.wmflabs.org/quickstatements/>
- Open Refine: <https://www.wikidata.org/wiki/Wikidata:Tools/OpenRefine>
- Wikidata Integrator: <https://github.com/SuLab/WikidataIntegrator>

A Wikidata Python module integrating the MediaWiki API and the Wikidata SPARQL endpoint

397 commits

2 branches

1 release

7 contributors

MIT

Branch: master

New pull request

Find file

Clone or download

 sebotic fixed an omission where new items don't get created when domain not s... Latest commit 2f5d2fd 22 hours ago

 doc Wikidata to Wikipedia mapping prototype for diseases added. 2 years ago

 wikidataintegrator fixed an omission where new items don't get created when domain not s... 22 hours ago

 Jenkins

Jenkins > Running >

- New Item
- People
- Build History
- Edit View
- Delete View
- Manage Jenkins
- My Views
- Credentials
- Build Queue

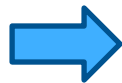
Running Bots

AllRunning+

S	Name	Last Success ↑	Last Failure
	ProteinBot_homo_sapiens	1 day 21 hr - #12	N/A
	GOBot_bigmern	2 days 15 hr - #15	9 days 15 hr - #14
	GeneBot_Homo_sapiens	2 days 19 hr - #25	2 days 20 hr - #24
	Disease_Ontology	2 days 23 hr - #11	4 days 13 hr - #8
	GeneDiseaseBot	2 days 23 hr - #9	1 mo 6 days - #2

Simple data retrieval

“Retrieve genes with
GWAS association
with asthma”



39 genes

gene	geneLabel		gene	geneLabel		gene	geneLabel		gene	geneLabel
Q5013317	COL22A1		Q18027370	IGSF3		Q18053559	CDHR3		Q14903974	SMAD3
Q14912759	SLC22A5		Q18045382	HPSE2		Q18045669	ATG3		Q18033889	IL1RL1
Q14914243	PSAP		Q18048437	IL33		Q18035037	RAD50		Q17917202	ERBB4
Q14907990	SLC30A8		Q18051900	PYHIN1		Q18036984	FBXL7		Q18027836	IL6R
Q18025002	GAB1		Q17709208	ACO1		Q18033919	XPR1		Q18030185	NOTCH4
Q18035589	C6orf10		Q18027822	IL2RB		Q15326496	RORA		Q18030409	PDE4D
Q18054256	GSDMA		Q18030364	PBX2		Q18042132	GSDMB		Q18045645	IKZF4
Q18058487	C5orf56		Q18037773	ABI3BP		Q18029145	MKLN1		Q18039979	KLHL5
Q18030785	PRKG1		Q18039623	CTNNA3		Q18036729	RAP1GAP2		Q18026947	HLA-DQA1
Q18033424	IL18R1		Q18046350	ZNF665		Q14878303	IL13			

```

1 SELECT DISTINCT ?gene ?geneLabel where {
2   ?gene wdt:P2293 wd:Q35869 .    # gene has genetic association to "asthma"
3   ?gene wdt:P31 wd:Q7187 .       # gene is subclass of "gene"
4   SERVICE wikibase:label { bd:serviceParam wikibase:language "en". }
5 }
```

Data integration

“Retrieve genes with
GWAS association
with asthma and gene
product is localized to
membrane”



22 genes

gene	geneLabel	gene	geneLabel	gene	geneLabel	gene	geneLabel
Q14912759	SLC22A5	Q18027370	IGSF3	Q18035037	RAD50	Q18027836	IL6R
Q14914243	PSAP	Q18033424	IL18R1	Q18033919	XPR1	Q18030409	PDE4D
Q14907990	SLC30A8	Q18045382	HPSE2	Q18042132	GSDMB	Q18030185	NOTCH4
Q18035589	C6orf10	Q18027822	IL2RB	Q18036729	RAP1GAP2	Q18026947	HLA-DQA1

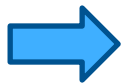
```

1 SELECT DISTINCT ?gene ?geneLabel where {
2   ?gene wdt:P2293 wd:Q35869 . # gene has genetic association to "asthma"
3
4   ?gene wdt:P31 wd:Q7187 .      # gene is subclass of "gene"
5
6   ?gene wdt:P688 ?protein .      # gene encodes a protein
7   ?protein wdt:P681 ?cc .        # protein has a cellular component
8   ?cc wdt:P279*|wdt:P361* wd:Q14349455 . # cell component is 'part of' or 'subclass of' membrane
9
10  SERVICE wikibase:label { bd:serviceParam wikibase:language "en". }
11 }
```

http://bit.ly/bosc2017_wikidata

Leveraging the Disease Ontology structure

“Retrieve genes with GWAS association with any respiratory disease and gene product is localized to membrane (non-IEA)”



31 genes / 8 diseases

diseaseGALabel	gene_counts	geneList
asthma	15	SMAD3, RAP1GAP2, IL18R1, HPSE2, SLC30A8, SLC22A5, PSAP, ERBB4, HLA-DQA1, IGSF3, IL2RB, IL6R, NOTCH4, PDE4D, RAD50
chronic obstructive pulmonary disease	5	HLA-C, SFTPD, ANXA5, ANXA11, ATP2C2
lung cancer	3	TGM5, VTI1A, PHACTR2
interstitial lung disease	2	DSP, ATP11A
non-small-cell lung carcinoma	2	NALCN, DLST
nasopharynx carcinoma	2	ITGA9, TNFRSF19
adenocarcinoma of the lung	1	BTNL2
pulmonary emphysema	1	BICD1

```

1 SELECT ?diseaseGALabel (count (DISTINCT ?geneLabel) as ?gene_counts)
2 (group_concat(DISTINCT ?geneLabel; separator="," as ?geneList)
3   ?gene wdt:P2293 ?diseaseGA . # gene is subclass of "gene"
4   ?diseaseGA wdt:P279* wd:Q3286546 . # to a type of respiratory system disease
5
6   ?gene wdt:P31 wd:Q7187 ; wdt:P688 ?protein ; # gene is subclass of "gene" and encodes protein
7   rdfs:label ?geneLabel .
8   FILTER (lang(?geneLabel) = "en")
9   ?protein p:P681 ?s . # protein's cell component statement
10  ?s ps:P681 ?cp . # get statement value
11  FILTER NOT EXISTS (

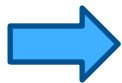
```

http://bit.ly/bosc2017_wikidata

... and show associated pathways

16 genes / 59 pathways

“Retrieve genes with GWAS association with any respiratory disease and gene product is localized to membrane (non-IEA), show causative chemical hazards and **show pathways where they have a role.**”



gene	pathway
SMAD3	Androgen receptor signaling pathway
SMAD3	TGF-beta Receptor Signaling
SMAD3	mechlorethamine exposure
HLA-C	Allograft Rejection
SFTPD	Regulation of toll-like receptor signaling pathway
....	

```

11  .cp wdt:P279 | wdt:P301 wd:Q4915012 .
12
13  ?pathway wdt:P31 wd:Q4915012 ;           # instance of a biological pathway
14      wdt:P527 ?gene .
15
16  SERVICE wikibase:label { bd:serviceParam wikibase:language "en". }
17  }
```

http://bit.ly/bosc2017_wikidata

PREFIX wp: <http://vocabularies.wikipathways.org/wp#>

PREFIX dcterms: <http://purl.org/dc/terms/>

PREFIX dc: <http://purl.org/dc/elements/1.1/>

SELECT DISTINCT ?metabolite1Label ?metabolite2Label ?mass1 ?mass2 WITH {

```
SELECT ?metabolite1 ?metabolite2 WHERE {  
  ?pathwayItem wdt:P2410 "WP706";  
  wdt:P2888 ?pwIri.
```

Wikidata

```
SERVICE <http://sparql.wikipathways.org/> {  
  ?pathway dc:identifier ?pwIri.  
  ?interaction rdf:type wp:Interaction;  
    wp:participants ?wpmb1, ?wpmb2;  
    dcterms:isPartOf ?pathway.  
  
  FILTER (?wpmb1 != ?wpmb2)  
  ?wpmb1 wp:bdbWikidata ?metabolite1.  
  ?wpmb2 wp:bdbWikidata ?metabolite2.  
}
```

Wikipathways

} AS %metabolites WHERE {

```
INCLUDE %metabolites.
```

```
?metabolite1 wdt:P2067 ?mass1.
```

```
?metabolite2 wdt:P2067 ?mass2.
```

```
SERVICE wikibase:label { bd:serviceParam wikibase:language "[AUTO_LANGUAGE],en". }
```

Wikidata

[Try me....](#)

From a remote SPARQL endpoint to Wikidata



SPARQL Downloads

Documentation/Help Contact

Your query

Add common prefixes

```
20 SELECT DISTINCT ?wd_item ?physically_interacts_with ?interactswithLabel ?type ?iri ?uniprot ?text WHERE {
21   {SELECT * WHERE { ?iri a up:Protein ;
22     up:organism taxon:9606 ;
23     up:annotation ?annotation .
24     ?annotation a up:Natural_Variant_Annotation ;
25     rdfs:comment ?text .
26     FILTER (CONTAINS(?text, 'loss of function'))
27   }}
28   SERVICE <https://query.wikidata.org/bigdata/namespace/wdq/sparql> {
29     VALUES ?use {wd:Q427492}
30     ?wd_item wdt:P352 ?uniprot ;
31       wdt:P129 ?physically_interacts_with ;
32       wdt:P2888 ?iri ;
33       wdt:P703 wd:Q15978631 .
34     ?wd_item p:P129 ?phys_interacts_with_node .
35     ?phys_interacts_with_node ps:P129 ?physically_interacts_with ;
36       pq:P366 ?use .
37     ?physically_interacts_with wdt:P31 ?type ;
38       rdfs:label ?interactswithLabel .
39     FILTER (lang(?interactswithLabel) = "en")
40   }}
```

UniProt

Wikidata

Submit Query

Cancel



Cellosaurus

@Cellosaurus

Following



ITS DONE!! All 107'576 [#Cellosaurus](#)
[#celllines](#) are now in [#Wikidata](#). There is still
some work to be carried out in the next
weeks in term of disease mappings but its a
major step. Thanks again to Lelia Debornes
who wrote the bot and to [@andrawaag](#) who
was of tremendous help



6:09 PM - 16 Jun 2018

Is Wikidata FAIR?

What makes a service FAIR?



WikiCite
@Wikicite

Following

Schema.org 3.5 has been released. It includes a vocabulary to represent grants/funding sources for research projects and the outputs they produce. Developed jointly for Schema and Wikidata.

[blog.schema.org/2019/04/schema ...](https://blog.schema.org/2019/04/schema...)

- **Issue #383**: Added initial vocabulary towards improved support description of project-based funding, for example to use with **Dataset** citations. See also **DINGO draft**, another outcome from the same **Wikidata / Wikibase workshop**. New types for **Grant**, **MonetaryGrant**, **Project**, **FundingAgency**, **FundingScheme**, and **ResearchProject**. Developed in collaboration with the **Wikidata/Wikibase** community.

3:22 AM - 3 Apr 2019

16 Retweets 30 Likes



WikiCite @Wikicite · Apr 3

More on the DINGO project ("Data Integration and Extension for Grant Ontology") github.com/dcodings/DINGO... [#linkeddata](#)

**projects and grants,
and specialisations for
Wikidata and
Schema.org .**

Editors:

- Diego Chialva
- Alexis-Michel Mugabushaka
- Andra Waagmeester
- Thomas Baker
- Dan Brickley
- Eric Prud'hommeaux

You, Wikidata, Peter Murray-Rust and WikiDigi



2



5



Recognition



Final Report and Action Plan
from the European
Commission Expert Group
on FAIR Data

TURNING
FAIR INTO
REALITY

Rec. 7: Support semantic technologies

Semantic technologies are essential for interoperability and need to be developed, expanded and applied both within and across disciplines.

Wikidata as a cross-disciplinary FAIR data platform

Wikidata (<https://www.wikidata.org>) is a multilingual collaborative database collecting, reusing and providing structured Open data. The platform hosts information across all areas of knowledge and is tightly integrated with all Wikipedia sites. About 18,000 people contribute in a typical month. The human contributors are aided by hundreds of automated or semi-automated tools that perform similar tasks at scale, based on community-agreed standards.

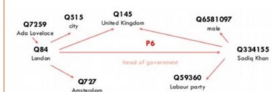


An identifier-first architecture

Each entity in Wikidata (e.g. an 'item' or a 'lexeme') has a globally unique and persistent identifier that can be used by humans and machines to retrieve information on the topic. Entities can be described using an increasingly rich metadata vocabulary that consists of several thousand uniquely identifiable 'properties'. Some of these express relationships between Wikidata entities, while others can be used to link concepts with concrete values, e.g. the height of a mountain or the pseudonym of a writer.



In contrast to classical Subject Predicate Object triples, Wikidata's data model includes optional qualifiers to make statements more specific, as well as references to highlight the provenance of a specific piece of information. Every entity is linked to multiple different assertions.



The identifier-first architecture has many benefits. It enables Wikidata to support hundreds of languages and allows editors from all over the globe to review, refine, expand, correct or otherwise build on each other's contributions in a FAIR manner.

Wikidata is a FAIR data platform:

- » It can be searched and queried in multiple ways, including via SPARQL, the query language of the Semantic Web
- » Wikidata is accessible via open, free, and universally implementable protocols, with authentication and authorization where necessary
- » Metadata provided by automated tools are usually associated with detailed provenance
- » Except for specific circumstances, metadata about deleted data remains available.
- » The data and metadata are published under CCO, which allows for reuse without restrictions
- » The software for the site and for most of the user-generated tools is openly licensed, which allows an ecosystem of federated FAIR databases to grow around Wikidata.

By acting as an identifier hub, Wikidata helps other resources across and beyond the research landscape – e.g. including the cultural heritage sector – increase their FAIRness.

Image credit: CC-BY Elena Simperl and Alessandro Piscopo, slides 6 & 7 www.slideshare.net/elenasimperl/quality-and-collaboration-in-wikidata

Figure 11: Wikidata case study: a cross-disciplinary FAIR platform

Ticking the criteria

(F)indable

(A)ccessible

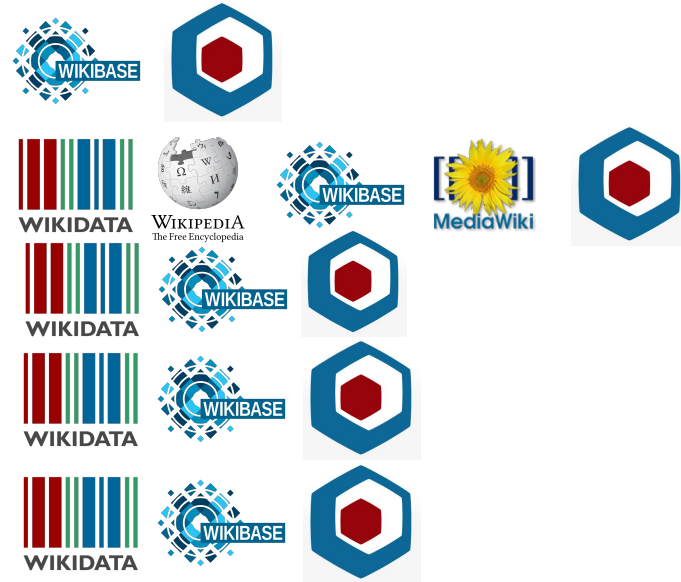
(I)nteroperable

(R)eusable



Where is Wikimedia ecosystem located in the Research Data Lifecycle

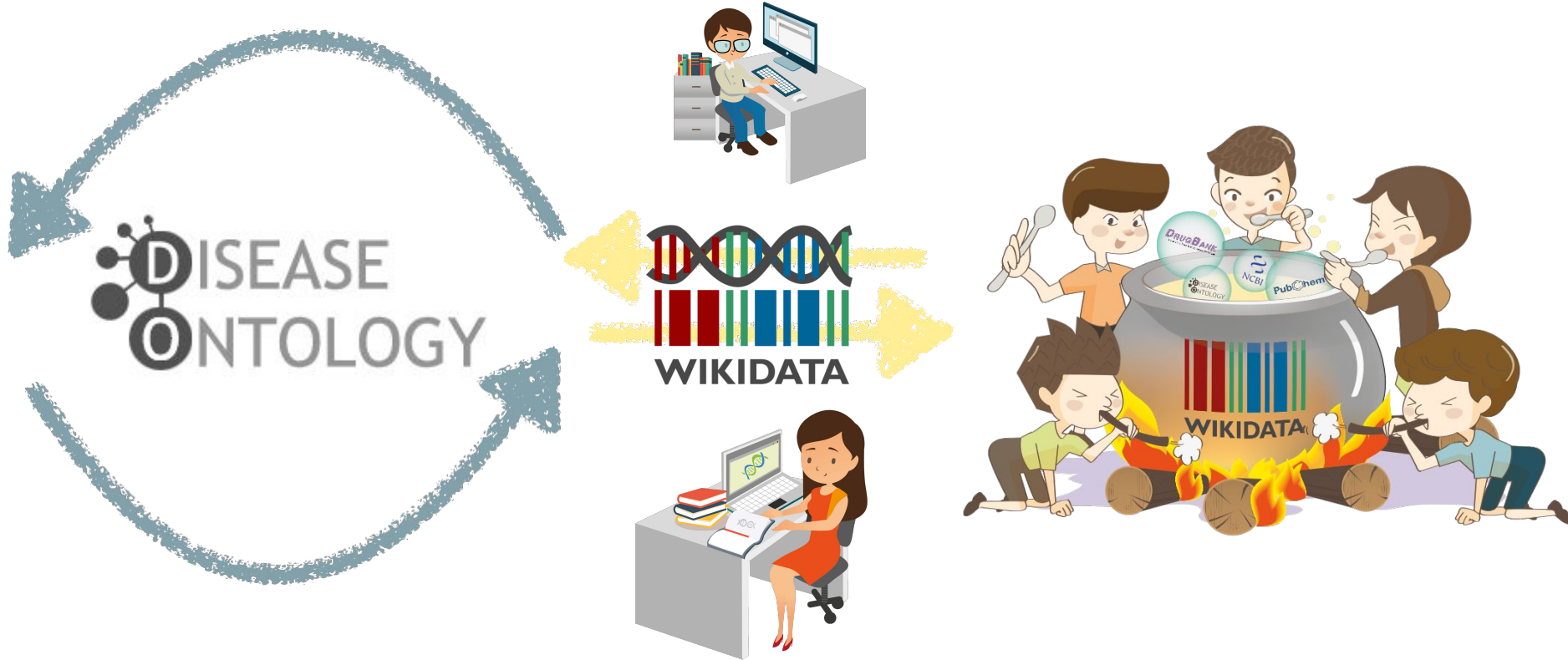
- Data creation
- Documentation
- Discovery
- Sharing
- Reuse



How do you interoperate/collaborate/engage with other services to help support FAIR data?



Updating Disease information from Disease Ontology in Wikidata



What is the one key change you think needs to happen for us to implement FAIR data and FAIR services?



Albert, 15:32

Hi Andra. Even een praktische vraag. Ken jij misschien een jr nederlandstalige data modeler / ontologist die, na een korte training voor 2019 voor 4 dagen in de week bij hjet Zorg Instituut kan worden ingezet?



Thursday, 14 March 2019



Rianne, 16:12

Ken je iemand met kennis over FAIR data / ontologiën die op zoek is naar een postdoc?



Rianne, 16:13

wij zoeken juist iemand met die kennis die een postdoc wil doen



het is blijkbaar een krappe markt, want we kunnen geen geschikte kandidaten vinden



Ruben Verborgh (UGent-imec) <Ruben.Verborgh@ugent.be>

to Eric, Andra ▾



English ▾



Dutch ▾

[Translate message](#)

I wish I knew such people, but I have trouble finding them myself :-)

Acknowledgements

Núria Queralt Rosinach



Benjamin
Good

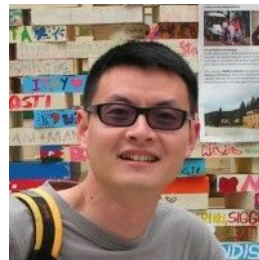


Andrew Su

Tim Putman



Sebastian
Burgstaller



Chunlei Wu



Gregory Stupp

- Elvira Mitraka
(Disease Ontology, U
Baltimore)
- Gang Fu, Evan Bolton
(NIH, PubChem)
- Wikimedia Foundation

Thousands of WikiDatans



By Helpameout - Own work, CC BY-SA 3.0,
<https://commons.wikimedia.org/w/index.php?curid=20337311>

Lynn Schriml



Funding

- National Institutes of General Medical Sciences (R01GM089820)
- NIH Common Fund programs for Big Data to Knowledge (U54GM114833)

Acknowledgements



Sebastian Burgstaller
Center for Integrative Bioinformatics,
Max Perutz Laboratories,
Vienna Biocenter.

- Elvira Mitraka
Disease Ontology, U
3altimore)
Gang Fu, Evan Bolton
NIH, PubChem)
Wikimedia Foundation

ands of WikiDatans



pameout - Own work, CC BY-SA 3.0,
commons.wikimedia.org/w/index.php?c
J337311

Núria Queralt Rosi



Lynn Schriml



Funding

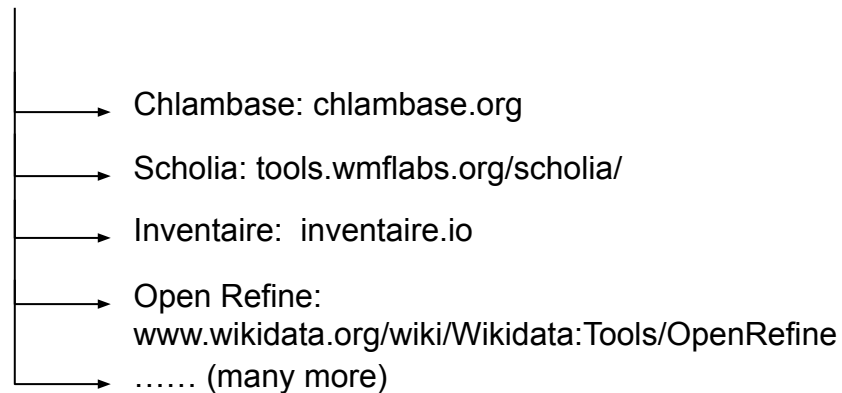
- National Inst
Sciences (R
- NIH Commo
Data to Know

Availability

Wikimedia ecosystem for FAIR data



Tools



Gene Wiki: github.com/SuLab/GeneWikiCentral

