

# Credit Distribution through Data Provenance in Relational Scientific Databases

Dennis Dosso<sup>a</sup>, Susan B. Davidson<sup>b</sup>, Gianmaria Silvello<sup>a</sup>

<sup>a</sup>*Department of Information Engineering, University of Padua, Italy*

<sup>b</sup>*Department of Computer and Information Science, University of Pennsylvania, United States*

---

## Abstract

In the current world of research data is a fundamental method to disseminate scientific knowledge, to determine scholarship, and to provide credit and recognition to the authors of research endeavors. However, issues like data citation, handling and counting the credit generated by such citations are still open research questions.

In this context, data credit has recently emerged as a new measure of value, defined and built on top of the data citation theory. Data credit is a real value that represents the importance of data cited by a paper, or by another research entity. As such, credit can be used to annotate data contained in curated scientific databases, and it can be considered as a measure for their importance and impact in the research world. As such, it is a new method that, together with traditional citations, helps to recognize the value of data and its creators in a world more and more dependent on data.

In this paper we explore the problem of Data Credit Distribution, the process by which credit is divided and assigned to the data in a database that are responsible for the production of data being cited by a research entity.

We adopt as use case the IUPHAR/BPS Guide to Pharmacology (GtoPdb), a curated and well-known scientific relational database. We define two new distribution strategies, functions that perform this task, based on two form of data provenance, why-provenance, and how-provenance.

Using different distribution strategies, we show how credit can highlight areas of a database that are frequently used, and how it can work as a new bibliometric measure for data and their corresponding curators. Credit in particular rewards data and authors based on their research impact, and not

merely on the number of citations. Also, we show how different distribution strategies, based on different types of data provenance, can be more sensible to the role of an input tuple in the generation of the output, and thus rewarding it differently.

*Keywords:* Data Citation, Data Credit

---

## 1 Introduction

Citations are an essential component of scientific research, enabling research products to be found as well as the relationships between research products to be understood. They form a basis on which to give credit to authors, papers, and venues [54, 19, 20]. Citations are used, among other things, to decide on tenure, promotion, hiring, and funding of grants for researchers [41, 21, 32, 38].

Nowadays, science and research are increasingly digital. There are numerous curated databases that are at the core of scientific research efforts [12]. It is therefore generally accepted that data must be cited and citable [39, 15], and that data citations should contribute to the scientific reputation of researchers, scientists, data curators, and creators [4, 50]. It is also accepted that data citations should be counted alongside of traditional citations, and contribute to bibliometrics indicators [7, 44].

A central problem in data citation is how to attribute credit to data creators and curators [11]. How to handle and count the credit generated by data citation, and how it contributes to traditional and new bibliometrics, are long-standing research issues Garfield [28], Borgman [9]. However, even when correctly applied, data citations and the bibliometric computed using them do not always correctly reward the creators of data used in a database. Data, in fact, is often cited at the “database level” or the “webpage level”. In the first case, the whole database is cited and therefore all credit goes to the key personnel of the database. In the second case, the database has a website with webpages that can be individually cited. The webpages use data extracted from the database, which is aggregated by topic and built to resemble a traditional research paper. Often the creators and curators of the webpage’s data are not credited or only marginally credited for their work [3].

Recently, the concepts of *data credit* and *Data Credit Distribution* (DCD) [26, 36, 53] have emerged, built on top of methodologies for data citation. Data

credit is a value that is computed based on the importance of the data being cited in a paper, and represents the impact of the data on the citing paper. The Data Credit Distribution problem consists of distributing this credit to elements in the databases in the citation graph that are responsible for the generation of the data being cited. The goal of DCD is to improve and expand the reach of data citation, rather than being an alternative to it. This means that to employ DCD techniques, we need data citations in some form.

[37] defined credit as a “quantity” that describes the importance of a research entity, such as papers or data mentioned in a citation, and proposed the idea of a *distribution* of credit from research entities, such as papers or data, to other research entities through citations. This can be done by exploiting the structure of the *citation graph*, a directed graph whose nodes are publications and edges are citations. This graph is the model at the core of systems such as Google Scholar and the Web of Science. Zeng et al. [53] and Fang [26] further explored this concept by defining frameworks for the computation and distribution of credit between papers, authors, and data used by papers in the citation graph.

In this paper, we consider data credit as a data value measure in a (curated) scientific database; credit can be assigned to data of any kind and at any level of granularity. Therefore the concept of “data” is left intentionally vague, although in this paper we focus on relational databases. Credit is a positive *real* value, acting as a proxy for the value of data based on the measure of citations, accesses, clicks, downloads, or other surrogates for data use. We call Data Credit Distribution the process, method, or algorithm used to assign credit to a given datum or dataset.

The DCD problem differs from the traditional citation setting since:

1. In a traditional setting, when a paper cites another paper, a +1 “credit” is given to the cited paper (and to its authors). It does not matter why or how paper  $p_1$  cites paper  $p_2$ <sup>1</sup>, the result is always +1 from  $p_1$  to  $p_2$  and thus a +1 to the citation count of the authors of  $p_2$ . With a different credit distribution strategy, the “value” given to the cited entity can be *proportional* to the role played in the citing entity. Hence, we can weigh the importance of the cited entities and assign credit according to their role.

---

<sup>1</sup>Note that there is vast research on this topic and many alternative proposals, but none of them currently work at a large scale.



Figure 1: Overview of the credit distribution pipeline.

2. Traditional citations are considered to be *atomic*. A citation from  $p_1$  to  $p_2$  can never be broken into pieces and assigned in part to  $p_2$  and in part to other papers or data that contributed to  $p_2$ . This is due to the intrinsic difficulty in grasping the role and “weight” of the other papers and data, and in automating the credit assignment process. In contrast, we consider data credit to be a *non-atomic* real value, which can be divided and distributed to multiple components of a database.
3. Credit can be *transitive*, that is, it can be propagated through one cited entity to other entities cited by it that contributed to its content.

We study the DCD problem in the context of relational databases (RDBs) since they are widely used<sup>2</sup> and are the main focus of current work in data citation methods [14, 12, 45]. RDBs are also frequently a test-bed for new methods that can be adapted to other databases, e.g., graphs or document databases. Furthermore, the “portions” of data in an RDB that can be credited can be defined at different levels of granularity, in particular: (i) the whole database, (ii) tables, and (iii) tuples.

The DCD process is summarized in Figure 1:

<sup>2</sup>The “relational database market alone has revenue upwards of \$50B” [1].

- 82 **Step 1** Scientists and experts contribute the curated information contained  
83 in a scientific database. These are called the “Data Curators”.
- 84 **Step 2** Other researchers use the data in their research, and when possible,  
85 cite them.
- 86 **Step 3** The citation to the data generates credit, that can be used as a  
87 proxy for the impact of the data on the citing paper. This credit is  
88 represented as a real value  $k \in \mathbb{R}_{>0}$ .
- 89 **Step 4** Given the database instance  $I$  and the query  $Q$ , it is possible to  
90 compute the *data provenance* of  $Q(I)$ . The provenance of  $Q(I)$  is a  
91 form of metadata that describes the generation process undertaken by  
92  $Q$ , and the data used in  $I$  to generate the output [17]. Many different  
93 notions of provenance have been proposed in the literature for data in  
94 database management systems [22, 13, 30], describing different kinds  
95 of relationships between data in the input and the output of a query.  
96 As reported in [17], these provenances have been used in several appli-  
97 cations beyond giving information on how queries work, for example,  
98 annotation propagation and the view update problem. In this paper,  
99 we consider three types of provenance: lineage, why-provenance, and  
100 how-provenance.
- 101 **Step 5** Provenance is input to the CDC problem, whose aim is to compute  
102 the *Credit Distribution Strategy* (CDS, also referred only as Distribu-  
103 tion Strategy, DS). The CDS is a function that distributes  $k$  to the data  
104 in the input database  $I$ , and is defined on the basis of citation policies  
105 decided at the database administration level or at the domain commu-  
106 nity level. In this paper, since we base CDS on data provenance, we  
107 describe three CDS, each one based on a different form of provenance.
- 108 **Step 6** Once the CDS is computed, it is used to distribute the given credit  
109  $k$  to the parts of the database that are responsible for the generation  
110 of  $Q(I)$ . Transitively, this credit is also divided and given to the corre-  
111 sponding authors of those data.

112 This paper expands our recent work in [24], which addressed the problem  
113 of how to reward data and data curators who are typically overlooked in  
114 current citation systems. In that work, we first defined the problem of DCD

115 in relational databases, and proposed a viable Distribution Strategy (DS)  
 116 based on *lineage*, which is the simplest form of *data provenance*. The lineage  
 117 of a tuple  $t$  in the output  $Q(I)$  is defined as the set of all and only the tuples  
 118 in the database instance  $I$  that are “relevant” to the production of  $t$ , that  
 119 is the tuple that are used by  $Q$  in the production of  $t$ . The lineage-based  
 120 strategy equally redistributes the credit  $k$  to the tuples in the lineage set,  
 121 thus each tuple receives credit  $k/|L_t|$ , where  $L_t$  is the lineage set of  $t$ .

122 One may argue that this DS is too simplistic, since lineage only tells  
 123 the relevant tuple used to produce the output, and does not convey any  
 124 information about their role or importance in the query. Therefore, one may  
 125 desire to give more credit to the tuples that are more relevant or *essential*  
 126 to the production of the output, i.e. those tuples that, if removed, would  
 127 prevent the output tuple from appearing in the final result, or those tuples  
 128 used more than once by the query.

129 Therefore, in this paper, we expand the ideas in [24] by proposing two  
 130 new DSs based on other forms of data provenance: why-provenance [13]  
 131 and how-provenance [30]. We compare them with the lineage-based solu-  
 132 tion, and discuss why one may be preferred to another depending on the  
 133 application and its goals. In particular, we show that why-provenance and  
 134 how-provenance are more sensitive to the *role* of a tuple in a query, i.e. how  
 135 many times the tuple is used and how it is used. The DS based on why-  
 136 provenance give more reward to tuples that are essential to the production  
 137 of the result set, whereas the DS based on how-provenance also takes into  
 138 consideration the different ways that a tuple is used.

139 For evaluation, we use a well-known curated database, the IUPHAR/BPS<sup>3</sup>  
 140 Guide to Pharmacology [31], also known as GtoPdb<sup>4</sup>, which contains ex-  
 141 pertly curated information about diseases, drugs, cellular drug targets, and  
 142 their mechanisms of action. We chose GtoPdb for two main reasons: (i) it  
 143 is a widely-used and valuable curated relational database, (ii) many papers  
 144 in the literature use, and cite its data (i.e., families, ligands, and receptors).  
 145 Real queries used in papers can therefore be seen as data citations which, in  
 146 turn, can be used to assign data credit.

147 We perform three sets of experiments. In the first one, real queries are ex-

---

<sup>3</sup>International Union of Basic and Clinical Pharmacology/British Pharmacology Soci-  
 ety

<sup>4</sup><https://www.guidetopharmacology.org/>

148 tracted from papers published in the British Journal of Pharmacology (BJP),  
149 that represent data citations to GtoPdb, and are used to distribute credit  
150 in the database using the three different provenance-based DSs. In the sec-  
151 ond and third experiment we analyse the behaviour of the different DS when  
152 complex citation queries are employed.

153 **Contributions.** Contributions of this work include:

- 154 • The definition of new distribution strategies for the problem of Data  
155 Credit Distribution, based on why-provenance and how-provenance;
- 156 • An in-depth analysis of the effects of credit distribution on real-world  
157 curated data and of the differences between the three proposed Distri-  
158 bution Strategies.

159 **Outline.** The rest of the paper is organized as follows: Section 2 presents the  
160 background and related work. Section 3 describes the use case we adopted.  
161 Section 4 briefly presents the forms of provenance used in the paper. Section  
162 5 describes the problem of DCD and the proposed DS. In Section 6 we present  
163 the experimental evaluation. Finally, Section 7 draws some conclusions and  
164 outlines future work.

## 165 2. Background

166 *Data in Research.* As described by Jim Gray in his last talk [33], the world of  
167 research is rapidly transitioning towards the *fourth paradigm of science*, that  
168 is, data-intensive scientific discovery, where data are important for scientific  
169 advances as well as for traditional publications [6].

170 The scientific community is promoting an *open research culture* [43],  
171 founded on methods and tools to share, discover, and access experimental  
172 data. The community has identified the FAIR principles (Findable, Acces-  
173 sible, Interoperable, and Reusable) [51], that should be enforced by every  
174 database. In particular, data should be accessible from the articles, journals,  
175 and papers that cite or use them [19]. Aspects such as the need for the *repro-*  
176 *ducibility* of experiments through the used data; the *availability* of scientific  
177 data; the *connections* between data and the scientific results are all needed  
178 aspects for the fourth paradigm, and are all relevant to the domain of *data*  
179 *citation* [34].

180 *Data Citation: Principles and Motivations.* Data Citation principles were  
 181 first described in detail in [18], and later summarized and endorsed by the  
 182 Joint Declaration of Data Citation Principles (JDDCP) [40]. The principles  
 183 are divided into two groups [48]. The first one contains principles concerning  
 184 the role of data citation in scholarly and research activities such as the (i)  
 185 *importance* of data (why data citation is important and why data should be  
 186 considered as first-class citizens); (ii) *credit* and *attribution* to the creators  
 187 and curators of the data; (iii) *evidence*; (iv) *verifiability*; and *interoperability*,  
 188 with these last three requiring data citation methods to be flexible enough to  
 189 operate through different communities. The second group defines the main  
 190 guidelines to establish a data citation systems, and contains principles such  
 191 as the (i) *unique identification* of the data being cited; (ii) (*open*) *access* to  
 192 data; (iii) guarantee of *persistence* and *availability* of citations even after the  
 193 lifespan of the cited entity; the (iv) *specificity* of a citation, i.e. it must lead  
 194 to the data set originally cited.

195 It is possible to outline six main motivations for data citation [48]:

- 196 • *Data attribution*: identify the individuals that should be credited for  
 197 data with variable granularity.
- 198 • *Data connection*: connect papers to the data being used.
- 199 • *Data Discovery*: citations helps to find data records and subsets that  
 200 would be otherwise not findable via search engines.
- 201 • *Data Sharing*: share data obtained by researchers within the whole  
 202 community.
- 203 • *Data Impact*: highlight the results obtained in writing papers using  
 204 specific data, the frequency and modality data were used.
- 205 • *Reproducibility*: data citation greatly impacts the reproducibility of  
 206 science [5]. Many authoritative journals ask to share data and provide  
 207 valid methodologies to reproduce experiments.

## 208 2.1. Data Citation in Relational Databases

209 In this paper, we develop our methods and experiments on relational  
 210 databases. RDBs have been the main target of data citation methods since  
 211 the surge of the data-centric research paradigm. The RDA “Working Group



212 on Data Citation: Making Dynamic Data Citable”<sup>5</sup> [46] has been working in  
213 the last years on large, dynamic, and changing datasets. The working group  
214 has finished the development of its guidelines and has now moved on into an  
215 adoption phase. The datasets considered by the WG are often relational.

216 In one of its most recent sessions [47], the Working Group (WG) on  
217 Data Citation reported that there are various implementations of its guide-  
218 lines for Data Citation on MySQL/Postgres relational databases. Some of  
219 these databases are: DEXHELPP<sup>6</sup> (Social Security Records); NERC (ARGO  
220 Global Array); EODC (Earth Observation Data Centre) [29]; LNEC (River  
221 dam monitoring); MDS (Million Song Database) [8]; CBMI<sup>7</sup> (Center for  
222 Biomedical Informatics); VMC (Vermont Monitoring Cooperative); CCA<sup>8</sup>  
223 (Climate Change Center Austria); VAMDC (Virtual Atomic and Molecular  
224 Data Center) [25, 55].

225 More examples of work on data citation in relational databases are [12,  
226 52, 2, 23]. The website <https://fairsharing.org/> keeps a long updated  
227 list of curated and scientific databases (many of which are relational or graph-  
228 based) following FAIR guidelines. These databases are citable since they are  
229 compliant with the most recent guidelines, and they are in the vast majority  
230 of cases accessible via dynamically created Webpages. In all these databases  
231 is, therefore, possible to implement DCD on top of the existing infrastructures  
232 for citing data.

233 Data citation techniques are primarily applied to relational databases  
234 because of their diffusion and also because the portions of data that are to  
235 be cited are easily identified: the whole database, a relation, a tuple, or  
236 even an attribute. Many papers [10, 12, 2] consider more complex citable  
237 units, recognizing that often the *views* of a database are the ones to be cited.  
238 Generally, a *view* is a query on the database. To this end, [52] suggested  
239 decomposing the database in a set of views, where each view is associated  
240 with its citation.

241 At present, the most common practices to cite databases include:

- 242 1. A database cited as a whole, even though only parts of the databases  
243 are used in the papers or datasets. Alternatively, the so-called “data pa-

---

<sup>5</sup><https://www.rd-alliance.org/groups/data-citation-wg.html>

<sup>6</sup><http://www.dexhelpp.at/>

<sup>7</sup>[https://medicine.missouri.edu/centers-institutes-labs/  
center-for-biomedical-informatics](https://medicine.missouri.edu/centers-institutes-labs/center-for-biomedical-informatics)

<sup>8</sup><https://ccca.ac.at/startseite>

- pers” can be cited, being traditional papers that describe a database [16]. In this case, all the credit from the citations goes to the database administrators or to the authors of the data papers.
2. Subsets of data, obtained by issuing queries to a database, are individually cited. This is the solution adopted by the *Resource Data Alliance* (RDA) working group on Data Citation [46]. In this case, the credit generated from citations can be distributed among the contributors of the portions of data being cited, and/or to the database administrators.
  3. The database is accessible via a series of Webpages that arrange the content of the database by topic or theme. Examples in the life science domain include the Reactome Pathway database [35], the GtoPdb [31], and the VAMDC [55]. Every single Webpage is unequivocally identifiable and can be individually cited.

Despite all the research efforts dedicated to the study and promotion of data citation, none of the largest citation-based systems, such as Elsevier Scopus, Web of Science, Microsoft Academia, or Google Scholar, consider scientific datasets as citable objects in academic work. Clarivate Analytics Data Citation Index (DCI) [27] is an exception, since its infrastructure tracks data usage in scientific domains and provides the technical means to connect datasets and repositories to scientific papers. However, DCI considers only citations to (previously registered and approved) databases as a whole and does not count citations to database portions such as views, tables, or tuples.

## 2.2. Data Credit

Data credit is related to data citation: they both aim to recognize the work of data creators and curators. Data credit can therefore also be seen as a by-product of data citation, since credit attribution is impossible without the presence of data citations.

Katz [36] suggests the need for a *modified citation system* that includes the idea of *transient* and *fractional credit*, to be used by developers of research products as software and data. In the paper two considerations are made: (i) research objects such as data and software are currently not formally rewarded or recognized by the community; (ii) even in traditional papers, the contribution of each author to the work is hard to understand, unless explicitly specified in the paper. This is even more true for data, where different groups of people work on the same database.

In [36] credit is defined as a “quantity” that describes the importance of a research entity, such as papers, software, or data, mentioned in a citation. We

add that the concept of credit can be built on top of the existing infrastructure handling traditional and data citations. Katz [36] further explores the idea of a *distribution* of credit from research entities (i.e., papers and data) to other research entities through citations that connect them. Thanks to traditional citations and now also to data citations, this distribution is finally possible, at least between papers and data. Some problems related to traditional citations can thus be solved by citations:

1. Credit rewards research entities that to date are not (formally) recognized (a goal shared with data citation).
2. Credit can reward authors *proportionally* to their role in generating the entity. The more an author contributes to a paper, the more credit is given to him. Zou and Peterson [54] work on something similar with their zp-index, which includes in its formulation the position (and thus the role) of a publication author to represent its impact in the work itself.
3. Credit can be *transitively* channeled through a chain of papers citing each other, thus enabling the rewarding of older papers that are no more cited, since other papers summarize or report their content but are nevertheless crucial in a research area for the influence of their content.

Fang [26] presents a framework to distribute the credit generated by a paper to its authors and to the papers in its reference list in a transitive way. Let us consider the *citation graph* as the graph where the nodes are papers and the links are the citations among them. In this graph, every paper is a source of credit, which is then transferred to the neighboring nodes. The quantity of credit received by each cited paper depends on its impact/role in the citing paper. So far, this theoretical framework is limited to papers, but it can be easily extended to a citation graph including both papers and data.

Zeng et al. [53] proposes the first method to compute credit within a network of papers citing data. Adopting a network flow algorithm, they simulate a random walker to estimate a score for each dataset, leveraging real-world usage data to compute the credit. This is the first step towards an automatic credit computation procedure. This proposal is, however, limited to assigning credit to whole datasets, and it does not deal with the granularity of data. It does not work to assign credit to a single research entity within a dataset.

317 Differently from Zeng et al. [53], we do not treat the credit computation  
 318 process, but we focus on the distribution process.

### 319 2.3. Data Provenance

320 To distribute credit, we base our methods on *data provenance*. Data  
 321 provenance is information that describes the origin and the process of cre-  
 322 ation of data. It can also be seen as metadata pertaining to the derivation  
 323 history of the data. It is particularly useful to help users to understand  
 324 where data are coming from, and the process they went through. Data ci-  
 325 tation and data provenance are closely linked [3] since both are forms of  
 326 annotations on data retrieved through queries. Data provenance has been  
 327 widely studied in different areas of data management. In this paper, we fo-  
 328 cus on provenance for database management systems (DBMS). For further  
 329 details on data provenance, please refer to surveys like [17] and [49].

330 Cheney et al. [17] presents four main types of data citation for DBMS: *lin-*  
 331 *age* [22], *why-provenance* [13], *how-provenance* [30] and *where-provenance* [13].

332 Let us start with the first three provenances. Given a database instance  
 333  $I$ , a query  $Q$ , and the result  $Q(D)$ , consider one tuple  $t$  of the output. Its  
 334 provenance is information about its generation through the tuples of the  
 335 input that are used by  $Q$ . Different types of provenance convey different  
 336 levels of information. Since these three provenances are computed for each  
 337 tuple of the output, they are also referred to as *tuple-based*.

338 Lineage is somehow the simplest among the forms of provenance. It has  
 339 been defined in different ways [17], but it can be thought of as the set of all  
 340 the tuples that are used in some way by the query to produce the output  
 341 tuple, the ones that are somehow *relevant* to its generation.

342 The definition of why-provenance is based on the notion of *witness set*.  
 343 A witness is a set of relevant tuples that guarantees the existence of  $t$  in  
 344  $Q(D)$ . The lineage is therefore an example of a witness. The why-provenance  
 345 of a tuple  $t$  is a peculiar set of witnesses – described in [13] – that are  
 346 computed from the query, called *witness basis*. A witness basis may be  
 347 composed of more than one witness. Therefore, the why-provenance contains  
 348 more information than the lineage, since it describes *alternative* ways in  
 349 which the same output may be generated.

350 The how-provenance takes the form of a polynomial, called *provenance*  
 351 *polynomial*, where the variables are taken from the set of identifiers of the  
 352 tuples (provided that each tuple in  $I$  has an identifier) and the coefficients are  
 353 taken from  $\mathbb{N}$ . This provenance also contains information on *how* the input

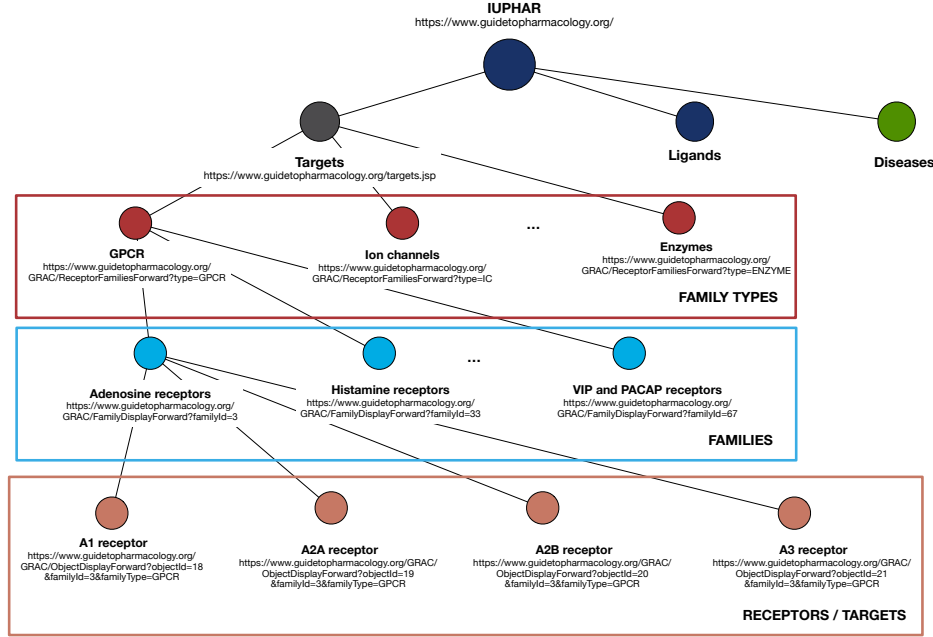


Figure 2: Partial map of the GtoPdb hierarchical structure grouping the targets into families and family types.

354 tuples are used. For example, when two tuples are combined by a join, they  
 355 are also combined in the polynomial by the  $\cdot$  operator. When two or more  
 356 tuples become equivalent due to a union or a projection, the corresponding  
 357 monomials are combined by the  $+$  operator.

358 It has been shown in [17] that the how-provenance is the more general  
 359 and informative of the three, containing the other two.

360 Where-provenance, differently from the other three, is *attribute-based*, so  
 361 we do not take it into account in this work since we consider the tuple as the  
 362 finest citable unit.

### 363 3. Use Case: GtoPdb

364 As use case we refer to the IUPHAR/BPS Guide to Pharmacology [31]  
 365 or GtoPdb<sup>9</sup>. GtoPdb is a well-known and well structured scientific relational  
 366 database that contains expertly curated information about diseases, drugs

<sup>9</sup><https://www.guidetopharmacology.org/>

367 in clinical use, their cellular targets, and the mechanisms of action on the  
368 human body. It is curated and maintained by the GtoPdb Committee, and  
369 by 96 subcommittees, comprising 512 scientists collaborating with in-house  
370 curators who draw the information contained in the database from high-  
371 quality pharmacological and medicinal chemistry literature. Roughly 1000  
372 researchers from all over the world have contributed to the database, and the  
373 curators wanted to give recognition to these contributors. This led to some  
374 early work on data citation [10].

375 GtoPdb is relational, but its logical structure is hierarchical as shown  
376 in Figure 2. The information contained in the database is also organized  
377 into webpages focused on specific diseases, targets or ligands, and families  
378 for easier access by users. As depicted in Figure 2, the database can be  
379 thought of as a tree where the root is the database; the first level consists  
380 of all targets, ligands, and diseases; and the lower levels consists of specific  
381 targets, ligands and diseases. In this paper, we focus on targets; thus at the  
382 third level in the figure we show examples of family types, at the fourth level  
383 we show specific families of targets (a finer level of granularity), and finally,  
384 at the last level, the single targets (also known as receptors).

385 GtoPdb provides access to the webpages corresponding to all these nodes  
386 through URLs. The webpages corresponding to target families all present a  
387 similar structure, as shown in Figure 3 for the “Adenosine receptors” family.  
388 Each page has an *Overview*, a brief text describing the content of the page;  
389 a list of *Receptors* comprising the family; a section of *comments* about the  
390 family; the *References*, a list of the papers consulted by the curators of the  
391 page, similar to a reference list of a paper; the *further reading* list, reporting  
392 papers that an interested reader may want to consult to obtain more insight  
393 on the family; and a final section called *How to cite this family page*, con-  
394 taining text snippets useful to cite the specific page or the whole database.  
395 Figure 3 shows the SQL code that retrieves the information used to build the  
396 corresponding sections (apart from the References section). Therefore, each  
397 family page can be considered a full-fledged traditional publication, consist-  
398 ing of title, authors, abstract (the overview), content, and references.

399 In practice, many papers in the literature only reference GtoPdb (the  
400 root) without including a reference to the specific page being cited. That is,  
401 they only cite a paper describing GtoPdb as a whole (e.g., [31]) and refer  
402 to targets, ligands, diseases, etc. only by name. Thus, citations to specific  
403 families are *de-facto* “hidden” to citation systems such as Google Scholar,  
404 and useless for the computation of bibliometrics.

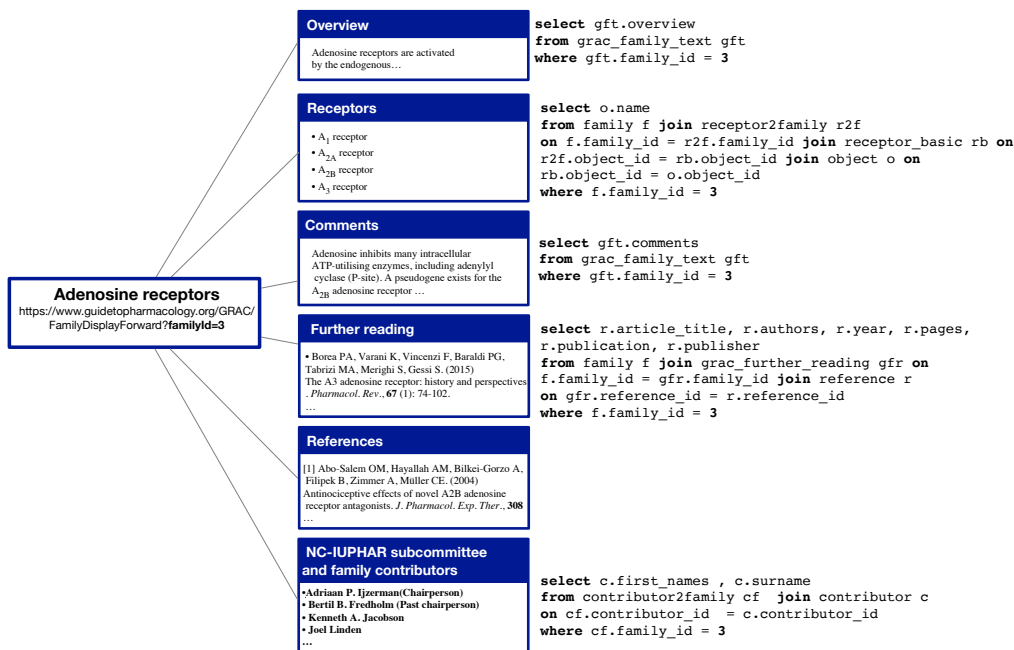


Figure 3: Basic web-page structure of “Adenosine receptors” family (ID 3), with queries used to retrieve the information contained in every section, except references.

In certain “lucky” cases, as with papers available in PDF and published in the British Journal of Clinical Pharmacology<sup>10</sup> (BJCP), when a family, ligand, receptor name, etc. are used, they have a hyperlink pointing to the corresponding webpage in GtoPdb. Therefore, the citations to the families can be detected and counted using the URLs reported in the papers. However, these citations to GtoPdb webpages are not counted as such by citation systems, so they are not converted into credit for curators and collaborators.

For our running example, consider Table 1. This simplified version of GtoPdb illustrates three tables: **family**, **contributor** and **contributor2family**. The first table, **family**, has tuples representing families with three attributes: the id of the family, its name, and type. Table **contributor** consists of people who have helped generate the data of the database. The third table, **contributor2family**, serves as a link between the families and the people who contributed to them. For instance, “John Smith” ( $c_1$ ) contributed to

<sup>10</sup><https://bpspubs.onlinelibrary.wiley.com/journal/13652125>

family			contributor2family		
id	name	type	id	family_id	contributor_id
$f_1$	Dopamine Receptors	gpcr	$c2f_1$	$f_1$	$c_1$
$f_2$	Bile Acid Receptor	gpcr	$c2f_2$	$f_1$	$c_2$
$f_3$	FAK Family	enzyme	$c2f_3$	$f_2$	$c_3$
$f_4$	YANK Family	enzyme	$c2f_4$	$f_4$	$c_1$

contributor		
id	Name	Country
$c_1$	John Smith	UK
$c_2$	Jim Doe	UK
$c_3$	Hans Zimmerman	Germany
$c_4$	Roberta Rossi	Italy

Table 1: Example of a database consisting of three tables. **family** includes some receptor families in the database; **contributor** contains the name and country of contributors; **contributor2family** connects contributors to the families they contributed to.

419 “Dopamine Receptors” ( $f_1$ ) as well as to the “YANK Family” ( $f_4$ ). We use  
420 this example throughout the rest of the paper. In particular, we are using  
421 the **id** attribute of the tables as *provenance token* of its corresponding tu-  
422 ples, that is, as a symbol that serves to identify a tuple when talking about  
423 provenance.

## 424 4. Data Provenances

425 In this section, we present the three types of provenance used in this  
426 paper: lineage, why-provenance, and how-provenance.

### 427 4.1. Lineage

428 Lineage was first introduced by Cui et al. [22]. Given a database instance  
429  $I$  and query  $Q$ , lineage associates with each tuple  $o \in Q(I)$  the set of tuples  
430 in the input that helped “produce” it [17]. As an example, consider the  
431 following SQL query **Q1**, applied to the database described in Table 1, that  
432 asks for the names of families curated by researchers based in the United  
433 Kingdom (UK):

```

434 Q1: SELECT DISTINCT f.name
435 FROM family AS f JOIN contributor2family AS c2f
436 ON f.id = c2f.family_id

```



437 JOIN contributor AS c ON c2f.contributor\_id = c.id  
 438 WHERE c.country = 'UK'

id	name	lineage
$o_1$	Dopamine Receptors	$\{f_1, c2f_1, c_1, c2f_2, c_2\}$
$o_2$	YANK Family	$\{f_4, c2f_4, c_1\}$

Table 2: Result of an SQL query applied to the database instance in Table 1, which asks for the names of families curated by a researcher based in the UK. Attribute `id` is not part of the output and was added to succinctly identify each tuple as provenance token. Each tuple is also annotated with its lineage.

439 Table 2 shows the query result, which consists of two tuples. We add  
 440 an extra attribute `id` so that we can easily refer to each result tuple. The  
 441 lineage for tuple  $o_1$  is the set  $\{f_1, c2f_1, c_1, c2f_2, c_2\}$ , since the tuple  $f_1$  was  
 442 joined with  $c2f_1$  and then with  $c_1$ , and was also joined with  $c2f_2$  and  $c_2$ . No  
 443 other tuple is used in the database to produce  $o_1$ . For tuple  $o_2$  the lineage is  
 444  $\{f_4, c2f_4, c_1\}$ . Lineage is defined for each tuple of the output, and can differ  
 445 between tuples.

#### 446 4.2. Why-Provenance

447 Why-Provenance was first defined in terms of a deterministic semistruc-  
 448 tured data model and query language [13]. While why-provenance can be  
 449 defined in many ways, we refer to [17], where it is expressed in terms of the  
 450 relational model using the relational algebra.

451 In particular, while lineage aims to find all and only the tuples in the  
 452 input relevant to the production of an output tuple, why-provenance aims to  
 453 find sub-instances of the input that “witness” a part of the output. Given a  
 454 tuple  $t$  in the query’s output, a *witness* is any sub-instance of the database  
 455 that produces  $t$ . In particular, the whole database and the lineage of  $t$  are  
 456 both witnesses of  $t$ . Since the definition of witness allows for the presence  
 457 of “irrelevant” tuples, the set of all witnesses is finite (since the database  
 458 instance  $I$  is finite), but it is potentially exponentially large [17].

459 Buneman et al. [13] defined the why-provenance of an output tuple  $t$  in  
 460 the result  $Q(I)$  as a special *subset* of the set of witnesses called the *witness*  
 461 *basis*. The witnesses of the basis depend on  $Q$ ; thus, each basis’s size is  
 462 bounded by the size of  $Q$ . The witnesses of the basis exclude tuples that  
 463 are irrelevant to  $t$  being produced by  $Q$ , and thus the basis tends to be very  
 464 small compared to the set of all possible witnesses [17]. The witnesses are

also *minimal*, in the sense that if one tuple is removed from one of these witnesses, it cannot produce the output.

id	name	why-provenance
$o_1$	Dopamine Receptors	$\{\{f_1, c2f_1, c_1\}, \{f_1, c2f_2, c_2\}\}$
$o_2$	YANK Family	$\{\{f_4, c2f_4, c_1\}\}$

Table 3: Result of a SQL query applied on the database of Table 1 with the why-provenance of the corresponding results.

In a sense, each witness in the witness basis captures one possible way in which the query can generate the output. To better understand this, consider the example in Table 3, where each tuple in the result of query Q1 is annotated with its why-provenance.

The why-provenance of output tuple  $o_2$  has only one witness, which coincides with its lineage. This happens because there is only one way this output tuple can be produced, i.e., for tuple  $f_4$  to be joined with  $c2f_4$  and  $c_1$ . On the other hand,  $o_1$  has a witness basis with of two witnesses, since there are two possible ways in which the query can generate  $o_1$ . One possibility is that  $f_1$  is joined with  $c2f_1$  and  $c_1$  (the first witness), and the second possibility is that  $f_1$  is joined with  $c2f_2$  and  $c_2$  (the second witness). This means that to generate  $o_1$ , it is sufficient that only one of the two witnesses is present in the input database.

#### 4.3. How-Provenance

While why-provenance describes the source tuples that witness an output tuple in the result of the query, it leaves out information about how the source tuples are used. How-provenance was therefore defined in [30] to capture this information using a *semiring* algebraic structure, and is a form of provenance that takes the form of a *polynomial*.

The key idea in Green et al. [30] is to use the two operators  $+$  and  $\cdot$  to represent two basic transformations that source tuples undergo as a result of applying a relational query to a database [17]. Two tuples may either be joined together, as an effect of a join (represented with the  $\cdot$  operator) or merged via union or projection (represented with the  $+$  operator).

Table 4 shows a simple example in which the two output tuples of our running example are annotated with their respective how-provenances. Tuple  $o_2$  was produced through the join among the input tuples  $f_4, c2f_4$ , and  $c_1$ . The three provenance tokens are, therefore “multiplied” together. The case of

id	name	how-provenance
$o_1$	Dopamine Receptors	$f_1 \cdot c2f_1 \cdot c_1 + f_1 \cdot c2f_2 \cdot c_2$
$o_2$	YANK Family	$f_4 \cdot c2f_4 \cdot c_1$

Table 4: Result of the example SQL query **Q1** with the corresponding how-provenances of the output tuples annotated.

$o_1$  is slightly more complex. This tuple, as already discussed, can be obtained through two different joins. The two monomials composing the polynomial represent these two alternatives. They correspond, in a way, to the witnesses of the why-provenance of  $o_1$ . The  $+$  operator represents the fact that the two monomials describe alternative derivations. The output tuple is the result of a merge of two distinct tuples after the projection on the attribute **name**. This merge is due to the fact that the result of a relational algebra expression is always a *set* of tuples, which corresponds to the presence of the **DISTINCT** operator in an SQL query. This simple example gives the basic idea behind how-provenance and how it allows us to track the operations that produced an output tuple.

Provenance polynomials may also have monomials whose exponents and/or coefficients are greater than one, for example,  $3f_1 \cdot c2f_1 \cdot c_1 + f_1 \cdot c2f_2^3 \cdot c_2^3$ . This is a polynomial of a tuple produced by a query where the result of the join between the tuples  $f_1$ ,  $c2f_1$ , and  $c_1$  is produced three times and then merged (e.g. as the result of a union), and the tuples  $c2f_2$  and  $c_2$  are used three times in the operation described by the second monomial (e.g., with nested queries).

## 5. Credit Distribution and Distribution Strategies

We now give formal definitions of data credit and Data Credit Distribution (DCD), and present three different Distribution Strategies (DSs) based on the forms of provenance discussed earlier: Lineage-based DS, Why-Provenance-based DS, and How-Provenance-based DS. We also show how these strategies distribute credit in the IUPHAR example discussed earlier.

### 5.1. Data Credit and Data Credit Distribution

Given a database instance  $I$ , a *recipient of credit* is a unit of information within  $I$ . In the case of relational databases, recipients may be (i) the whole database; (ii) a table; (iii) a tuple; or (iv) an attribute.

523 *Data credit* is a value  $k \in \mathbb{R}_{>0}$ . Every recipient in a database is annotated  
 524 with a quantity of credit as a proxy for its importance. In this paper, we  
 525 focus on *tuples* as recipients of credit.

526 Given a *distribution strategy* (DS), *Data Credit Distribution* (DCD) takes  
 527 a database instance  $I$ , quantity of credit  $k$ , and query  $Q$  over  $I$ , and splits  $k$   
 528 among the recipients of credit in  $I$ .

529 In the following, we use the notation in Cheney et al. [17]: Given an  
 530 instance  $I$ , a *tuple location*  $(R, t)$  is a tuple  $t$  in relation  $R$ . With reference to  
 531 the running example,  $(\text{family}, \langle f_1, \text{Dopamine Receptors}, \text{gpcr} \rangle)$  is the  
 532 tuple location of the first tuple in the `family` relation. The set of all tuple  
 533 locations in  $I$  is called *TupleLoc*. We use this to formally define DCD at the  
 534 *tuple level*.

535 **Definition 5.1. Tuple Level Data Credit Distribution (DCD) [24]**  
 536 *Given a query  $Q$  over  $I$  and  $k \in \mathbb{R}_{>0}$ , DCD is defined by the function  $f_{I,Q} :$   
 537  $\text{TupleLoc} \times \mathbb{R}_{>0} \rightarrow \mathbb{R}_{\geq 0}$  such that  $f_{I,Q}(t, k) = h$  where  $0 \leq h \leq k$  and  
 538  $\sum_{t \in \text{TupleLoc}} f_{I,Q}(t, k) = k$ . The function  $f_{I,Q}$  is the distribution strategy (DS).*

539 As we can see, the DS is a function that annotates each tuple in the  
 540 database with a real value, which is a fraction of the given quantity  $k$ . The  
 541 only constraint is that the sum of the credit annotations on tuples must be  
 542  $k$ , i.e. that no credit is generated or destroyed during the distribution. Given  
 543  $I$  and  $Q$ , many different DSs may be defined as long as they sum up to  $k$ .

544 In what follows, we use information provided by data provenance to de-  
 545 fine distribution functions. For simplicity, we assume that the credit  $k$  is  
 546 distributed equally across the set of output tuples (i.e. the result of a query),  
 547 and discuss how the credit of one output tuple  $o$ ,  $k_o$ , is distributed across the  
 548 instance  $I$ .

## 549 5.2. A Lineage-based Distribution Strategy

550 In the lineage-based distribution strategy, each tuple in the output of  
 551 a query distributes credit equally to each input tuple that appears in its  
 552 lineage. More formally:

**Definition 5.2. Lineage-based Distribution Strategy [24]**

*Let  $I$  be a database instance,  $Q$  a query over  $I$ ,  $o \in Q(I)$  an output tuple and  
 $k_o$  the credit associated to  $o$ . Let  $L$  be the lineage of  $o$  and  $t$  be a tuple in  $I$ ,*

then  $t$  receives credit equal to:

$$f_{I,Q}(t, k_o) = \begin{cases} 0 & \text{if } t \notin L \\ \frac{k_o}{|L|} & \text{if } t \in L \end{cases}$$

553 Note that lineage-based DS distributes credit only to input tuples that  
 554 have a role in creating  $o$  by the query  $Q$ , and that each receives an equal  
 555 share of credit via  $o$ . Thus, the more tuples in a lineage set, the less credit  
 556 each tuple receives.

557 As an example, consider the output tuples of Table 2, and assume that  
 558 each output tuple has credit  $k_o = 1$ . The lineage of the first tuple,  $o_1$ , is  
 559 the set  $\{f_1, c2f_1, c_1, c2f_2, c_2\}$ . Therefore, each tuple in this set receives credit  
 560  $1/5$ . The other tuples of the database receive zero credit. The lineage of the  
 561 second output tuple is  $\{f_4, c2f_4, c_1\}$ , therefore each of these tuples receives  
 562 credit  $1/3$ .

563 At the end of the process, tuples  $f_1$ ,  $c2f_2$  and  $c_2$  each receive credit  $1/5$ ,  
 564 tuples  $f_4$  and  $c2f_4$  receive  $1/3$ , while tuple  $c_1$  receives  $8/15$ . Note that if a  
 565 tuple appears in more than one lineage set, then it will accumulate credit  
 566 from the distribution associated with each one of these sets, implying that  
 567 it has a more significant role in the context  $Q$ , as is the case with  $c_1$  in this  
 568 example.

569 Not all of the tuples in the lineage of an output tuple are necessary to be  
 570 present at the same time for the output tuple to appear in the query results.  
 571 For example, if the database only had the set of tuples  $\{f_1, c2f_1, c_1\}$  or the set  
 572  $\{f_1, c2f_2, c_2\}$ , the existence of  $o_1$  would still be guaranteed. In other words,  
 573 while  $f_1$  is always needed for  $o_1$  to appear in the output, only one of the sets  
 574 of tuples  $\{c2f_1, c_1\}$  and  $\{c2f_2, c_2\}$  is required. One could therefore argue that  
 575 it would be more fair for  $f_1$  to receive more credit than the other four tuples,  
 576 given its role in producing  $o_1$ .

577 This highlights one limitation of the lineage-based DS: while able to find  
 578 all and only the relevant tuples of the output, it does not distinguish the  
 579 *importance* of tuples in the query computations. We therefore present two  
 580 other, more sophisticated, forms of distribution strategies based on why- and  
 581 how-provenance.

### 582 5.3. A Why-Provenance-Based Distribution Strategy

583 The distribution strategy based on why-provenance first equally distributes  
 584 the credit  $k_o$  among the witnesses of the witness basis for  $o$ , and then equally

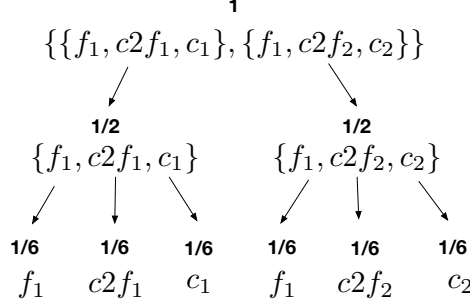


Figure 4: Distribution of credit using why-provenance-based DS for tuple  $o_1$ .

divides the credit of a witness among the tuples in the witness. Since a tuple may appear in more than one witness, it will receive more than one portion of credit from the same distribution. More formally:

**Definition 5.3.** *Why-Provenance-based Distribution Strategy*

Let  $I$  be a database instance,  $Q$  a query over  $I$ ,  $o \in Q(I)$  an output tuple and  $k_o$  the total credit associated to  $o$ . Let  $\mathcal{W} = \text{Why}(Q, I, o)$  be the witness basis of  $o$  according to  $Q$  and  $I$ , and  $W \in \mathcal{W}$  be a witness.

Then tuple  $t$  in  $I$  receives credit equal to:

$$f_{I,Q}(t, k_o) = \frac{k_o}{|\mathcal{W}|} \sum_{W \in \gamma(\mathcal{W}, t)} \frac{1}{|W|}$$

where  $\gamma$  is a function which returns all witnesses  $W$  in which  $t$  appears:

$$\gamma(\mathcal{W}, t) = \{W \in \mathcal{W} : t \in W\}$$

Figure 4 shows the distribution of credit with why-provenance-based DS for tuple  $o_1$ . The credit is first equally divided between the two witnesses, so that both receive credit  $1/2$ . The credit is then further divided among the tuples in each witness. Since each witness has three tuples, each tuple in a witness receives  $1/6$  of credit. At the end of the distribution,  $f_1$  receives a total credit of  $1/3$ , and the other tuples receive  $1/6$  each. This distribution better reflects the role of  $f_1$  in the generation of  $o_1$  since, as discussed earlier, it is the only mandatory tuple for  $o_1$  to appear in the output; only one of the two other pairs of tuples are necessary for  $o_1$  to appear in the result.

This example illustrates that why-provenance can better reward input tuples depending on their role. Tuples that appear in more than one witness are rewarded more than others.

$$\begin{aligned}
\mathcal{H} &= \underbrace{3f_1 \cdot c2f_1 \cdot c_1}_{M_1} + \underbrace{f_1 \cdot c2f_2^3 \cdot c_2^3}_{M_2} \\
c(\mathcal{H}) &= 4 & c(M_2) &= 7 \\
mc(M_1) &= 3 & mc(M_2) &= 1 \\
e(c_2, M_2) &= 3 & \gamma(c_1, \mathcal{H}) &= \{M_1\} \\
\gamma(f_1, \mathcal{H}) &= \{M_1, M_2\}
\end{aligned}$$

Figure 5: Illustration of notation used to define the how-provenance based DS in Definition 5.4.

#### 5.4. A How-Provenance Based Distribution Strategy

How-provenance conveys more information than why-provenance since it not only captures what tuples are relevant to the output and in which combination, but also how they are used. The “how” is captured through the provenance polynomials.

The how-provenance-based DS therefore first distributes the credit to the monomials of the polynomial accordingly to the weight represented by their coefficients, then to the tuples of each monomial accordingly to the weights represented by their exponents.

To define the DS more formally, we introduce some notation and illustrate it using the provenance polynomial  $\mathcal{H}$  shown in Figure 5.

We call  $c$  the function that, given a polynomial, returns the sum of the coefficients of the polynomial; thus  $c(\mathcal{H}) = 3 + 1 = 4$ . We use the same name for the function that, given a monomial, returns the sum of its exponents; thus  $c(M_2) = 1 + 3 + 3 = 7$ .  $mc$  is the function that takes as input a monomial and returns its coefficient.  $e$  is a function that takes as input a tuple and a monomial, and returns the exponent of the tuple in the monomial, if present; thus  $e(c_2, M_2) = 3$ .  $\gamma$  takes as input a tuple and the whole polynomial, and returns a set containing the monomials containing that tuple, if present in the polynomial; thus  $\gamma(f_1, \mathcal{H}) = \{M_1, M_2\}$ .

#### Definition 5.4. How-Provenance-Based Distribution Strategy

Let  $I$  be a database instance,  $Q$  a query over  $I$ ,  $o \in Q(I)$  an output tuple,  $\mathcal{H}$  be the provenance polynomial for  $o$ , and  $k_o$  the credit given to  $o$ . The credit given to tuple  $t$  in  $I$  is:

$$f_{I,Q}(t, k_o) = \frac{k_o}{c(\mathcal{H})} \sum_{M \in \gamma(t, \mathcal{H})} mc(M) \frac{e(t, M)}{c(M)}$$

id	name
$oxs_1$	Dopamine Receptors

lineage	why-provenance	how-provenance
$\{f_1, c2f_1, c_1, c2f_2, c_2\}$	$\{\{f_1, c2f_1, c_1\}, \{f_1, c2f_2, c_2\}\}$	$f_1^2 c2f_1 c_1 + f_1^2 c2f_2 c_2$

Table 5: Result of query Q2 applied on the database of Table 1 and its different provenances. The reported numbers are the credit distributed through the process.

628        Going back to the example of Table 4, consider  $o_1$  with provenance poly-  
629        nomial  $f_1 c2f_1 c_1 + f_1 c2f_2 c_2$ . The how-provenance-based DS firstly divides  
630        the credit between the two monomials. Since the coefficients of each mono-  
631        mial are 1, the credit is split in half. If they were, for example, 1 and 2  
632        respectively, 1/3 of the credit would go to the first monomial, and 2/3 to  
633        the second. Since in our example each variable has exponent 1, the credit  
634        is further divided equally among the three variables. Thus, at the end of  
635        the computation,  $f_1$  receives 1/3, and the other tuples receive 1/6. If, for  
636        example, the first monomial was  $f_1^2 c2f_1 c_1$ , then the portion of credit of this  
637        monomial would be divided in this way: 1/2 to  $f_1$  and 1/4 to each of the  
638        other two tuples.

639        In this specific example, the how-provenance-based DS has the same out-  
640        come as the one based on why-provenance. We therefore consider another  
641        query over GtoPdb, Q2, that asks for the families of type **gpcr** that have as  
642        contributor a researcher located in the UK:

```

643        Q2: SELECT DISTINCT F.name
644        FROM family as F JOIN
645        (SELECT DISTINCT f.name AS name
646        FROM family AS f JOIN contributor2family AS c2f ON f.id = c2f.family_id
647        JOIN contributor AS c ON c2f.contributor_id = c.id
648        WHERE c.country = "UK") AS R ON F.name = R.name
649        WHERE F.type = "gpcr"

```

650        The result of Q2 is shown in Table 5, and consists of one tuple, anno-  
651        tated with each of the three provenances. As can be seen, lineage and why-  
652        provenance are identical to those of the tuple  $o_1$  in the previous example.  
653        The how-provenance, however, is different since tuple  $f_1$  is used twice: first  
654        in the join of the inner query, and second in the join of the outer query. This  
655        information is lost in the first two forms of provenances since they are sets,  
656        but it is captured in how-provenance through the use of the operator ‘.’.



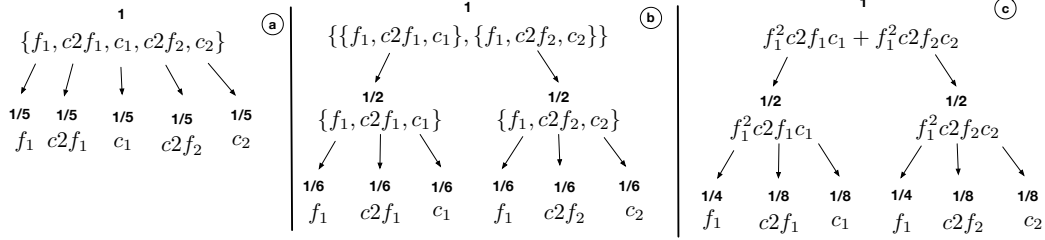


Figure 6: Comparison of different distributions strategies for tuple  $o_1$  produced by query Q2.

Figure 6 shows the differences between the three DS for the tuple  $o_1$  of Table 5. Subfigure 5.a uses lineage, sub-figure 5.b uses why-provenance, and sub-figure 5.c uses how-provenance. The DS based on the provenance polynomial gives credit  $1/2$  to  $f_1$ , and  $1/8$  to the other tuples. This is reasonable since Q2 relies on  $f_1$  even more than Q1 does. The distribution based on how-provenance can reward  $f_1$  more, showing that how-provenance is even more sensitive to the tuples' role in a query than why-provenance. This is a direct consequence of the fact that, as proven in [30], how-provenance is more general than why-provenance and lineage, in the sense that it contains more information.

## 6. Experimental Evaluation

To understand the trade-offs between these Distribution Strategies (DSs), we perform four sets of experiments using queries over target families presented on the GtoPdb website. The first set of experiments use real queries extracted from citations to GtoPdb published in the British Journal of Pharmacology. The second set uses synthetically produced provenance polynomials, corresponding to more complex queries, in order to highlight the differences between the DSs. The third set of experiments considers the accrual of credit over time by the three strategies, again using synthetic queries. The fourth set of experiments shows how the DSs compare to traditional citations in giving credit to data curators using both real and synthetic queries. We close by discussing relative execution times of the three strategies.

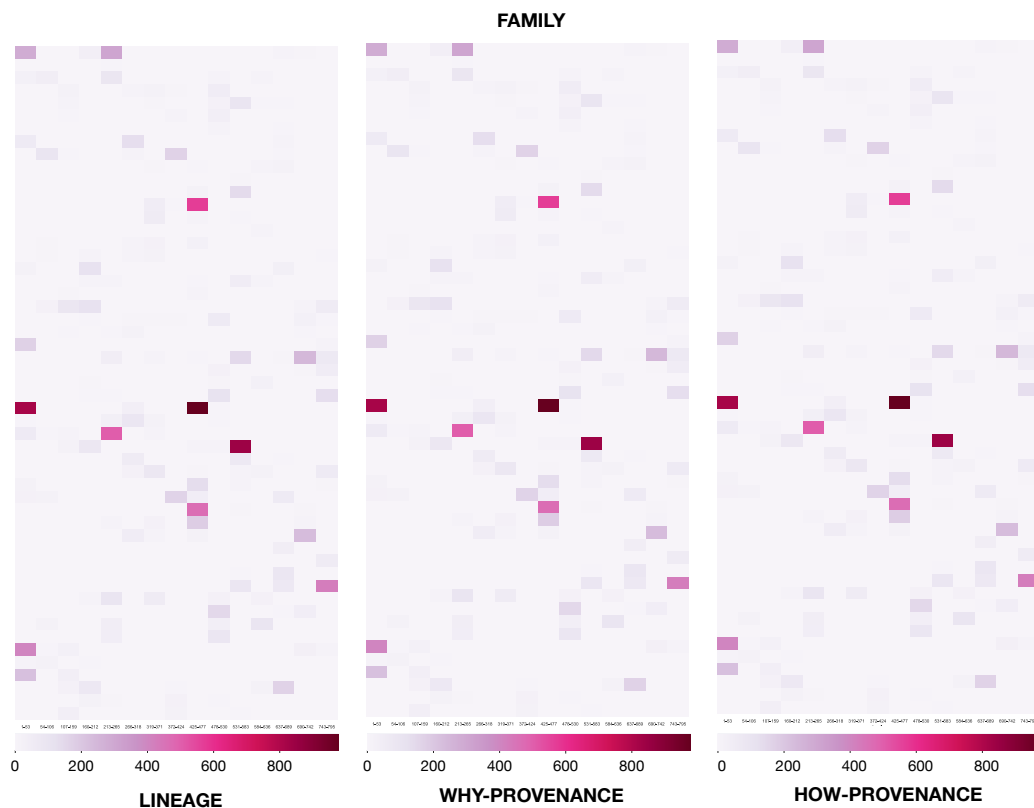


Figure 7: Comparison of three DS on the same table `family` using the distribution given by the queries retrieved from papers.

### 6.1. Real-world queries

Examples of real queries are drawn from papers published in the British Journal of Pharmacology (BJP) <sup>11</sup>. Each time a paper in this journal cites a webpage from GtoPdb, it reports the URL of the page. From this URL, the query used to obtain the webpage data can be determined. We considered all 889 papers in BJCP citing the IUPHAR/BPS Guide to pharmacology [31] as of October 2020, and extracted all webpage URLs to GtoPdb contained within the paper<sup>12</sup>.

<sup>11</sup><https://bpspubs.onlinelibrary.wiley.com>

<sup>12</sup>The IUPHAR/BPS Guide is a journal that describes the structure and evolution of GtoPdb. At the time of writing, it had received more than 1200 citations on Google Scholar.

687 There are eight target family types presented on the GtoPdb website:  
688 *GPCR, Ion channels, NHRs, Kinases, Catalytic receptors, Transporters, En-*  
689 *zymes and Other protein targets.*

690 The queries that we inferred are those used to build target family web-  
691 pages. An example was given in Figure 3, where we show how the structure of  
692 the “Adenosine receptors” family can be mapped into queries over the under-  
693 lying database. In GtoPdb, all target family pages share a similar structure;  
694 the only difference is that individual sections, such as “contributors” or “fur-  
695 ther readings”, may be absent. Therefore, the same queries can be used to  
696 build all of the target family pages by simply changing the family id used in  
697 the query (in Figure 3, it is 3). Note that the queries are fairly simple SQL  
698 queries, and fall into a class called “select-project-join” or “SPJ” queries. A  
699 total of more than 12K different queries were built in this way.<sup>13</sup> Without  
700 loss of generality, we give each tuple in the output of a query a credit of 1.

701 *Results.* Figure 7 shows the heat-maps obtained by the distribution of credit  
702 according to the three different DS on one of the tables in the underlying  
703 database, **family**, which is often joined with other tables in the database to  
704 build the webpages. It can be seen that the result of credit distribution over  
705 **family** is the same for all three strategies. The same result is also obtained  
706 with the other tables of the database used by the queries shown in Figure 3.

707 The reason why credit distribution is the same for all three strategies  
708 is that the queries are all simple SPJ queries, which use each table only  
709 once and do joins on key attributes. Under these conditions, each tuple of  
710 the output presents: (i) a how-provenance that is a single monomial with  
711 coefficient 1 and exponent 1 in each variable; (ii) a why-provenance with  
712 only one witness; and (iii) a lineage that coincides with the witness in the  
713 basis. Hence, for these queries, the three DSs behave in the same way: credit  
714 is uniformly distributed among the tuples present in each provenance.

715 To illustrate this, consider one of the queries in Figure 3 which is used to  
716 build the output webpage:

```
717 Q3: SELECT c.first_names, c.surname
718 FROM contributor2family AS cf JOIN contributor AS c ON
719 cf.contributor_id = c.contributor_id
```

---

<sup>13</sup>For reproducibility purposes, the code we used for our experiments and all queries are available here: [https://bitbucket.org/dennis\\_dosso/credit\\_distribution\\_project](https://bitbucket.org/dennis_dosso/credit_distribution_project).

**How-provenance:**  $3f_1^3c_1^2f_1^2c_1^2 + 2f_1c_2f_2^3c_2^3 + 4f_5c_2f_{17}^4c_{18}^3$

**Credit distribution:**

$$f_1 = \frac{59}{315}, f_5 = \frac{1}{18}, c_2f_1 = \frac{2}{21}, c_2f_2 = \frac{2}{15}, c_2f_{17} = \frac{2}{9}, c_1 = \frac{2}{21}, c_2 = \frac{2}{15}, c_{18} = \frac{1}{6}$$

**Why-provenance:**  $\{\{f_1, c_2f_1, c_1\}, \{f_1, c_2f_2, c_2\}, \{f_5, c_2f_{17}, c_{18}\}\}$

**Credit distribution:**

$$f_1 = \frac{2}{9}, f_5 = \frac{1}{9}, c_2f_1 = \frac{1}{9}, c_2f_2 = \frac{1}{9}, c_2f_{17} = \frac{1}{9}, c_1 = \frac{1}{9}, c_2 = \frac{1}{9}, c_{18} = \frac{1}{9}$$

**Lineage:**  $\{f_1, f_5, c_2f_1, c_1, c_2f_1, c_2f_2, c_2f_{17}, c_1, c_2, c_{18}\}$

**Credit distribution:**

$$f_1 = \frac{1}{8}, f_5 = \frac{1}{8}, c_2f_1 = \frac{1}{8}, c_2f_2 = \frac{1}{8}, c_2f_{17} = \frac{1}{8}, c_1 = \frac{1}{8}, c_2 = \frac{1}{8}, c_{18} = \frac{1}{8}$$

Figure 8: Sample synthetic provenance polynomial (how-provenance) and corresponding why-provenance and lineage expressions with deriving credit distributions.

720 WHERE f.family\_id = 3

721 Q3 returned 10 tuples from the version of GtoPdb used. The first tu-  
 722 ple, <Bertil B., Fredholm>, has  $c_{939} \cdot c_2f_{496}$  as its provenance polynomial.  
 723  $c_{939}$  represents the provenance token of a tuple in `contributor`, and  $c_2f_{496}$   
 724 the provenance token of a tuple in table `contributor2family`. The why-  
 725 provenance of this tuple is  $\{\{c_{939}, c_2f_{496}\}\}$  and its lineage is  $\{c_{939}, c_2f_{496}\}$ .  
 726 Therefore, the credit assigned to these tuples is 1/2 using all three DS. This  
 727 happens for all the tuples in the output of each query of GtoPdb, thus making  
 728 the distributions equivalent over all outputs.

729 However, this is not the case with more complex queries. As we showed  
 730 in the previous section, when two or more tuples are merged as a result of  
 731 a projection or union, the credit distributions will differ between the three  
 732 strategies.

## 733 6.2. Synthetic queries

734 To simulate synthetic queries, we randomly generated provenance poly-  
 735 nomials in which the coefficients and exponents could be greater than 1 over  
 736 three GtoPdb tables: `family`, `contributor2family`, and `contributor`. An  
 737 example can be found in Figure 8, which shows a sample synthetic provenance  
 738 polynomial (the how-provenance) and the corresponding why-provenance and  
 739 lineage expressions. The resulting credit distribution for each DS is shown  
 740 after the provenance expression.

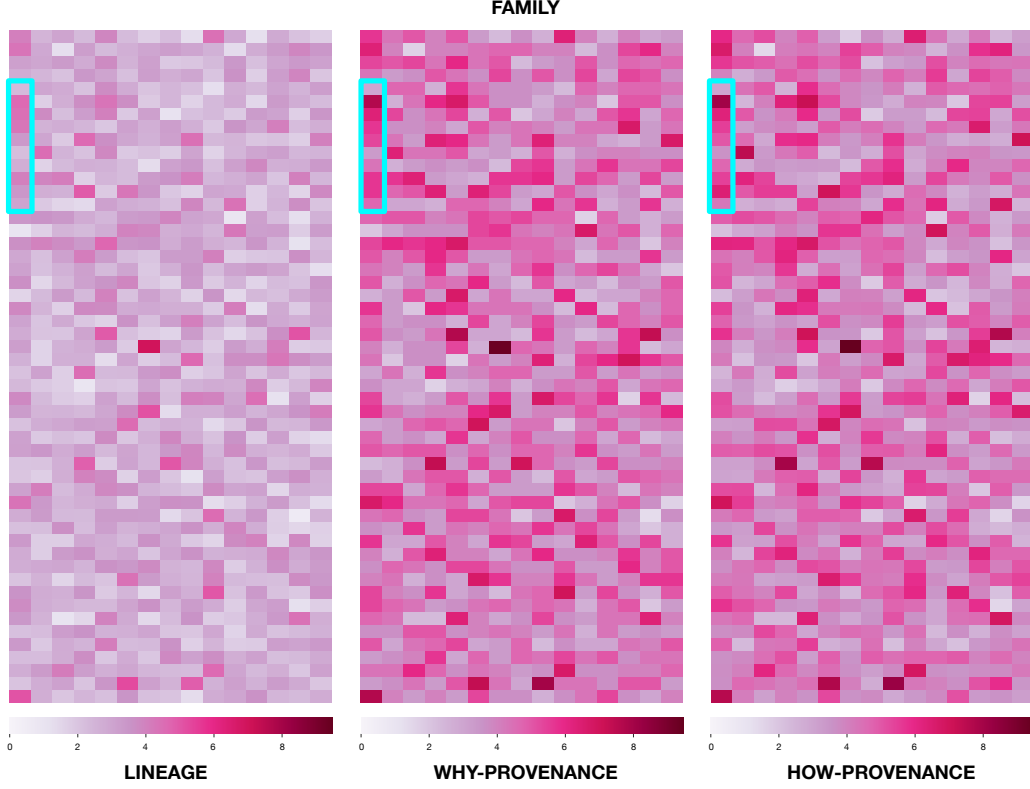


Figure 9: Comparison of three DS on the same table `family` after the distribution computed using 10K synthetic and randomly generated provenance polynomials. The tuples in the blue rectangles are used as example in the discussion connected to Figure 10.

As an example of how the distribution strategies behave with these synthetic queries, consider tuple  $f_5$  in Figure 8. This tuple receives the highest quantity of credit using lineage-based distribution, and less credit using why- and how-provenance because more information is available about the role of the tuple in the overall computation. Generally speaking, the more complex the distribution (the most complex being how-provenance), the more credit is given to tuples which are more frequently used, and thus have a higher impact in producing the output tuple.

Although synthetic, these provenance polynomials represent realistic queries. The polynomials can be obtained by any nested query with join and union operations that use the same tuple multiple times (in which case the exponents are bigger than 1), and the same combination of operations more than

753 once (in which case the coefficients of monomials are bigger than 1).

754 *Results.* The results of credit distribution on the **family** table using 10K  
755 randomly generated synthetic provenance polynomials are shown in Figure  
756 9. We set the maximum value in the heat maps to the highest value reached  
757 by a tuple in all three distributions (i.e., 9.4).

758 As can be seen, the three strategies generate significantly different credit  
759 distributions indicated by the varying hues. We note that, however, there is  
760 a consistency in how credit is distributed between the tuples, i.e. tuples that  
761 are highly rewarded by one strategy are also highly rewarded by the others,  
762 and vice-versa. This shows that the three DS consistently reward certain  
763 tuples more than others.

764 In particular, lineage-based DS gives the least credit to tuples in the  
765 **family** table, indicated by an overall lighter hue. This is because the DS  
766 equally distributes credit to all tuples appearing in the lineage. Since these  
767 queries also use two other tables, credit is distributed to tuples in those  
768 tables.

769 Moving to why-provenance based DS, we see that more credit is given to  
770 tuples in the **family** table than with the previous strategy. This is because  
771 the DS considers the different ways that a tuple is used, e.g. in joins with  
772 other tuples. If the same tuple is present in more than one witness, it will  
773 draw more credit and take it from other tuples in the witness basis. In this  
774 case, tuples in **family** drew more credit, taking it from tuples in the other  
775 two tables, due to the role that **family** tuples played in the queries that were  
776 executed.

777 Finally, consider the how-provenance-based DS heat-map. As with why-  
778 provenance, more credit is typically given to tuples in **family** compared to  
779 lineage-based DS since it recognizes the role of these tuples in the queries,  
780 and the overall hue is deeper. The two distributions appear similar, although,  
781 to a closer inspection, it is possible to note some smaller differences between  
782 the two distributions. This is because this DS also considers the frequency by  
783 which tuples are used, not only the ways in which they are used. Therefore,  
784 although the overall distribution is similar, there are small differences due  
785 to the presence of exponents and coefficients in the provenance polynomials,  
786 influencing the distribution of credit.

787 To better show this difference, let us consider the ten tuples within each  
788 large blue rectangle (each small rectangle within the large blue rectangle is  
789 a tuple). We will number them from 1 (top) to ten (bottom), and we use



Figure 10: Comparison of the distribution of credit performed by the three DSs on a subset of 10 tuples taken from the `family` table, simulating the passing of time. The number at the top of each group of heat-maps represents the number of queries.

790 them in the discussion in the next section.

### 791 6.3. Credit accrual over time

792 Since credit accrues over time, we simulate the passage of time by varying  
 793 the number of queries executed, and look at the “snapshots” of credit for each  
 794 of the strategies using synthetic queries. The results are shown in Figure 10.

795 In this figure, four groups of heat-maps are shown. Each group represents  
 796 a “snapshot” taken after 1K, 2K, 5K and 10K provenance polynomials have  
 797 been considered for credit distribution. The ten tuples in each heat-map are  
 798 from the `family` table, and are the ones highlighted previously in the blue  
 799 boxes of Figure 9.

800 The queries used are the same as the experiment reported in the previous  
801 section. The range of credit in each map goes from 0 (no credit) to 8 (the  
802 maximum quantity of credit reached on one of the tuples of the considered  
803 window at the “snapshot” with 10K queries). The color hue of the legend,  
804 as can be seen, still ranges from 0 to 9.5.

805 Focusing on the 1K and 2K groups, we see that credit distribution by the  
806 three DS are very similar. Still, there are small differences. We note that,  
807 in the 1K group, tuple 4 is the one with the highest value of credit with all  
808 3 strategies. Moving to 2K group, it is still the one with the highest value  
809 of credit, although it presents the highest value with the why-provenance  
810 DS. The other two strategies rewarded it less. Also, why-provenance and  
811 how-provenance rewarded more tuples 2 and 3 than the strategy based on  
812 lineage. Tuple 5 appears to be more rewarded by why-provenance, and less by  
813 how-provenance and even less by lineage. This shows, even with these lower  
814 values of polynomials, that the strategies may differ and reward certain tuples  
815 more than others, and vice versa. We see the tendency of the lineage DS to  
816 reward the tuples in this table less than the other strategies, since it does  
817 not take into consideration their importance. Instead, the DS based on why-  
818 provenance rewards more tuples like 4 and 5 (values 2 in both cases). The  
819 same can be said of the strategy based on how-provenance. However, in this  
820 case, tuples 4 and 5 are rewarded a little less (with credit values of 1.9 and  
821 1.5 respectively). This is due to the fact that how-provenance contains more  
822 information. Thus, this DS rewarded more other tuples in the other used  
823 tables. Viceversa, tuples 2 and 3 are rewarded more by how-provenance DS  
824 (values 1.5 and 1.6) than the why-provenance DS (value 1.3 in both cases),  
825 due to the fact that their roles in the polynomials are more important.

826 Moving to the 5K group, we see how credit was accumulated on the  
827 tuples. Now tuple 2 is the one with the biggest quantity of credit in this  
828 window. This shows how credit is able to track how the importance of tuples  
829 changes over time. In this group we see of it is more evident the difference  
830 between the distribution based on lineage and the other two strategies. The  
831 why-provenance and how-provenance based DSs appear to work similarly,  
832 that is to give similar values of credit to the same tuples. We can still see  
833 differences, for example on tuples like 8 and 6, that are more rewarded by  
834 the DS based on lineage.

835 Similar observations can be seen for the 10K group. We see how tuple  
836 2 is still highly rewarded by all three provenances. In the case of lineage,  
837 however, it is at the same level with tuples 3, 4, and 8, while the other two



838 strategies reward it the most. Once again we see the DSs based on why-  
839 and how-provenance operate similar distributions (we still note differences of  
840 few decimals between the values assigned to the tuples). However, it is still  
841 possible to see how tuples like 9 are more rewarded by one DS, in this case  
842 the how-provenance one, than the other. This shows how the last two DS  
843 operate in a similar way. The differences between the credit assigned to the  
844 same tuple is of few decimals between the two strategies in most of the cases.  
845 However, there are certain situations when the role of a tuple is particularly  
846 critical in a query. This information is captured by the provenance polyno-  
847 mial, and this is why in certain cases the differences in the credit assigned to  
848 one tuple is notably different between the two strategies.

849 To sum up, the DS based on lineage is sufficient when a user only wants  
850 to highlight the tuples of the database used by a query. This strategy equally  
851 distributes the credit to the tuples of the lineages. The resulting distribution  
852 rewards more the tuples that are used by more queries, but does not reward  
853 how the tuples are used in the same query.

854 For this reason, a user may want (depending on the nature of the queries)  
855 to use DS based on why-provenance and how-provenance. Using the why-  
856 provenance and how-provenance DS, it is possible to change the distribution  
857 of credit to the tuple, rewarding more the tuples that have a more critical  
858 role in generating the output. The distribution based on how-provenance is  
859 most of the times similar to the one based on why-provenance. However, in  
860 certain instances, it differ from the other one due to the specific role of a  
861 tuple, an information that is present in the provenance polynomial but lost  
862 in the witness basis. These two DSs can be preferred when the user aims to  
863 find “hotspots” in the database based on the tuples’ role, and the DS based  
864 on how-provenance can be chosen when the user wants the higher sensibility  
865 to the role of a tuple.

#### 866 6.4. *Credit vs Citations*

867 In the last set of experiments, we compare traditional citations to the  
868 proposed credit distribution strategies to see the difference in reward for  
869 authors, including data curators.

870 Each target family page in GtoPdb has a list of curators, that represents  
871 all the people who worked on the page and that can be considered as co-  
872 authors of that page. It is possible to obtain the list of one page authors  
873 using the last query shown in Figure 3. As set of authors, we consider all  
874 the curators of one target family in GtoPdb. Each time data are cited, we

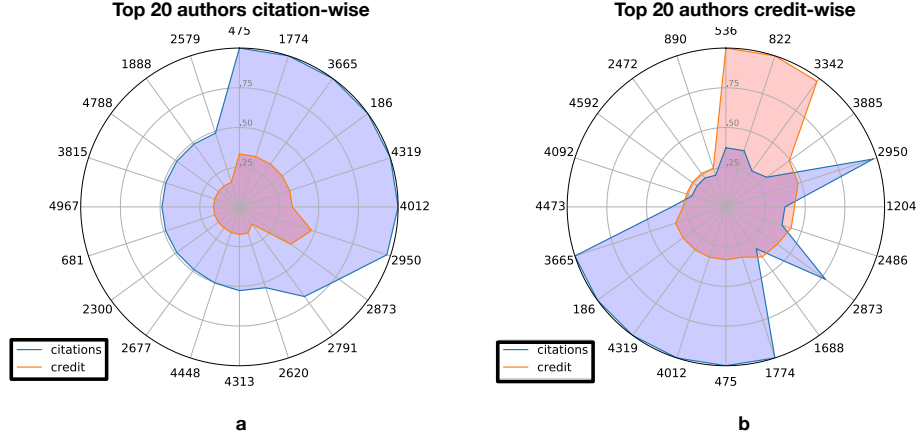


Figure 11: Radars presenting the top 20 authors citation-wise and credit wise, together with their (normalized between 0 and 1) values of citations and credit.

875 assign one citation to each author associated to that data. Each time some  
876 data, e.g. a tuple, receives credit, we equally divide and transitively assign  
877 that quantity of credit to the authors of that data.

878 *Results: Real-world queries.* As described in Section 6.1, we consider real-  
879 world queries, taken from papers published in the BJCP which reference  
880 webpages in GtoPdb. Since for these queries there is no difference in the  
881 distribution of credit between the three DS, only one value for credit is given  
882 in Figure 11. As we said, each time a webpage is cited, the authors of that  
883 webpage receive one citation, and also receive a quantity of credit that is  
884 equal to the credit assigned to the data and generated from the citation,  
885 equally divided among them.

886 The results are shown in the radar plots of Figure 11, in which each  
887 number on the outer circle (e.g. 475, 1774 and 3665) represents an author (id)  
888 and the blue (pink) line represents the normalized value of credit generated  
889 by citations (credit), respectively. The first radar plot, Figure 11.a, shows the  
890 top 20 authors, ordered in a clockwise direction based on citations, whereas  
891 Figure 11.b, orders the authors with the highest value of credit. As we  
892 see from Figure 11.a, the top 20 authors that present the highest values of  
893 citations do not also have the highest values of credit. Vice-versa, as seen in  
894 Figure 11.b, the authors with the highest values of credit do not necessarily  
895 also have the highest values of citations, i.e. the two sets of authors are not

896 equal (although there is partial overlapping between the two, e.g. authors  
897 1774, 475, 4012, etc.).

898 From this figure we understand that it is not the case that authors with  
899 the higher number of citations also present the highest values of credit. This  
900 is due to the fact that certain citations are more “valuable” in terms of credit,  
901 i.e. an author receive more credit from his/her citations, even if fewer, than  
902 other authors. This, in turns, happens because certain citations generate  
903 more credit than others. So, for example, author 536 presents the highest  
904 value of credit, although he is not even in the top 20 authors in terms of  
905 citations. This means that he receives much more credit from his relatively  
906 few citations than author 475.

907 Given how we prepared our experiments, the citations whose query pro-  
908 duce more tuples, are also the ones that generate more credit, since we as-  
909 sumed that each output tuple carries credit 1. Thus, the authors of the data  
910 returned by these queries are the ones that will receive more credit from  
911 these citations. Also, the authors that collaborated with fewer people will  
912 also receive a biggest share from the equal subdivision of credit.

913 This shows how credit distribution actually rewards differently authors  
914 than traditional citations. An author that curated larger quantities of cited  
915 data, and together with fewer co-authors, will receive bigger quantities of  
916 credit. Thus, credit can reward him for his more relevant “contribution” to  
917 the database.

918 In more complex and sophisticated scenarios, where different strategies  
919 may be implemented to decide the generated quantity of credit to be dis-  
920 tributed, new factors beyond the only “quantity” of curated data can be  
921 factored in in rewarding data curators. The result will be a distribution of  
922 credit that represents even better the actual work and worth of data curators  
923 and their impact in the scientific community.

924 *Results: Synthetic queries.* We produced 100, 1K, and 10K synthetic poly-  
925 nomials, as described in Section 6.2, and distributed credit through them  
926 to data. Since these polynomials are composed by tuples randomly selected  
927 from three tables, they usually correspond to very large set of authors that  
928 in reality did not collaborate. To simplify the assignment of data curators to  
929 the hypothetical queries generating these polynomials, we created 20 “syn-  
930 thetic” authors, and we randomly assigned one author to each tuple in the  
931 database. The authors receive “blocks” of consecutive tuples, with each block  
932 of a size varying between 10 and 40 to simulate different quantities of “work”

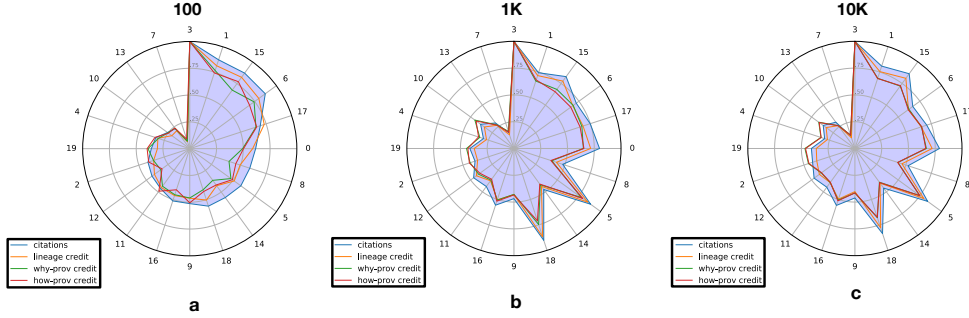


Figure 12: Radars presenting the 20 synthetic authors with corresponding citation and quantities of credit distributed through the 3 DS (all values normalized between 0 and 1) through different numbers of polynomials (respectively, 100, 1K and 10K). The order is the one defined by figure 1, i.e. descending order of citations obtained from 100 polynomials.

performed by an author. Every time an author appears as curator of one or more tuples used in a polynomial, we assigned him one citation. He also receives three kinds of credit, the ones assigned to his tuples through the three different DSs.

Figure 12 reports the three radar plots that are a consequence of the distribution of credit and citations performed and described above. Figure 12.a reports the values obtained with 100 polynomials, showing the normalized values of the citation and types of credit assigned to each author. As we see, given the synthetic nature of these queries, the correlation between the number of citations and the quantity of credit assigned to the authors appears to be a much stronger with respect to the case with the real-world queries of Figure.11 (the linear correlation between the citation number and all three types of credit is always above 0.95 with p values in the order of  $1e-11$ ). Nonetheless, it is still possible to observe how credit does not always exactly follow the citations. The credit distributed via lineage is the one that follows closer the number of citations (a linear correlation of 0.98, p value of  $6.15e-16$ ), while the other types of credit behave slightly differently (a linear correlation of around 0.95 in both cases).

Similar observations can be made for Figure 12.b and 12.c, where we kept the order of authors of Figure 12.a.

What appears from these figures is that, in certain cases, authors that do not have the highest values of citations receive more credit than others, as for example author 11 in Figure 12.a, or author 19 in Figures 12.b and 12.c, with credit distributed with how-provenance-based DS.

957 This once again shows how credit allows us to gain a different perspective  
 958 on the role of data and authors by going beyond the limitations of traditional  
 959 citations.

960 It is worth pointing out that, when scaling up to  $1K$  and  $10K$  polyno-  
 961 mials, the distributions performed via why-provenance and how-provenance  
 962 become almost equivalent. We can note that, although not exactly over-  
 963 lapping, the values of credit assigned to the authors by those DS become  
 964 quite similar with these higher quantities of polynomials, suggesting a sort  
 965 of equivalence between the two DSs in this case, at least in the task of re-  
 966 warding authors (the linear correlation for the values of Figure 12.c is more  
 967 than 0.99 with a p-value of  $1.32e-32$ ). This is consistent to what we already  
 968 pointed out in Figure 9.

#### 969 6.5. Execution time

# of polynomials	lineage	why-prov.	how-prov.
100	226.6 ms	192.0 ms	185.5 ms
200	431.2 ms	392.2 ms	403.2 ms
500	1.013 s	934.2 ms	881.8 ms
1K	2.041 s	1.934 s	1.744 s
2K	3.773 s	3.491 s	3.510 s
5K	8.992 s	8.653 s	8.889 s
10K	17.10 s	16.84 s	16.84 s
20K	34.59 s	35.30 s	39.70 s
100K	3.289 min	3.442 min	3.652 min
1M	35.91 min	34.87 min	37.91 min

Table 6: The times required to perform the three DS for different number of synthetic polynomials.

970 Table 6 shows the time required to calculate the credit distribution for  
 971 the three strategies. As can be seen, the execution time grows linearly with  
 972 the number of polynomials that are submitted to the system. With a high  
 973 number of polynomials (1M), the time required by the DS based on lineage  
 974 and why-provenance is lower than the time needed for the DS based on  
 975 how-provenance. This is due to the more significant number of operations  
 976 required to calculate the how-provenance DS and distribute the portions of  
 977 credit to be assigned to the different tuples. We note that, since we created  
 978 these polynomials on-the-fly, these values do not include the time required

979 to compute the provenances. Therefore, limited to the time required to  
980 distribute credit, the three DS are equivalent in terms of performances. The  
981 first differences can be seen only with high number of polynomials, when  
982 lineage and why-provenance may be preferred if there are no requirements  
983 to assign credit with the strategy implemented by the how-provenance-based  
984 DS.

985 All the experiments were carried out on a MacBook Pro with a 2.4 GHz  
986 processor Intel Core i5 quad-core and 8 GB of memory at 2133 MHz. Code  
987 was written in Java, supported by a PostgreSQL database.

## 988 7. Conclusions

989 This paper defines two new distribution strategies based on why- and  
990 how-provenance, and compares them against the lineage-based distribution  
991 strategy defined in [24]. The first DS, based on why-provenance, uses the  
992 concept of a witness, and gives more credit to tuples that appear in more than  
993 one witness. In this way, tuples that are more important to the query and  
994 are used in different ways are rewarded more. The second DS, based on how-  
995 provenance, considers the frequency with which a tuple or combination of  
996 tuples is used in the query through the information contained in a provenance  
997 polynomial. In this case, the distribution is even more sensitive than the first  
998 to the role and importance of tuples.

999 To show the differences between the three DSs, we performed extensive  
1000 experiments based on GtoPdb, a curated scientific relational database, using  
1001 both real and synthetic queries. In the first set of experiments, we used select-  
1002 project-join (SPJ) queries extracted from citations to webpages in GtoPdb  
1003 found in papers published in the British Journal of Pharmacology. Using  
1004 these “real” queries, we distributed credit to tuples in different tables of the  
1005 database, highlighting tuples that were more frequently used. We showed  
1006 that, with these queries, the three strategies produce the same distribution.  
1007 This is because the SPJ queries were fairly simple, and did not use self-joins.  
1008 Therefore the formulas underlying the different DSs had the same output.

1009 In the second set of experiments, we synthetically produced more complex  
1010 provenance polynomials, corresponding to more complex synthetic queries,  
1011 that resulted in exponents and coefficients in the provenance polynomials  
1012 that were greater than (or equal to) 1. These experiments highlighted the  
1013 differences between the three DSs. While the DS based on lineage rewards  
1014 all the tuples used by a query equally, the strategy based on why-provenance

gives more credit to tuples that are more critical to the query. In particular, why-provenance consider the different ways in which a tuple is used in a query. How-provenance is even more sensitive to the tuple’s role: it also considers the frequency with which a tuple or a set of tuples is used.

In the third set of experiments, we showed how the differences between the DS are compounded over time, i.e. when more and more queries are processed by the system.

In the fourth set of experiments we compared traditional citations to authors to the credit accrued to them via the DSs. We showed how, both in the real-world and synthetic scenarios, credit rewards authors who contribute/curate data that have the highest impact, and therefore receives the biggest quantity of credit, and not necessarily the data with the highest citation count. In this sense, credit appears to be an useful new measure to discover data and their corresponding curators that have a high impact in the research world, even when they are cited few times or do not appear at all in the data that are cited (i.e. the case of data used to build the output of a query but that is not visualized in the output itself).

In future work, we plan to explore different applications of credit over relational databases. One example is *data pricing*, which gives a price to a query submitted by a user who wants to buy the produced information. Currently, a commonly strategy used for data pricing is based on query rewriting: A database stores a set of views with their price. When a new query arrives, the system rewrites it using the stored views to obtain a query price, a process that can be computationally expensive. We plan to distribute credit through carefully planned and representative queries, and use credit information to define a new, faster, and potentially more flexible pricing function.

Another application is *data reduction* [42], which addresses the problem of reducing the vast – and rapidly expanding – amount of data that is being produced.

Data credit can also address this problem, by helping find “hotspots” and “coldspots” of data. A hotspot is data in a database (e.g. a tuple) with a high quantity of credit, which is therefore valuable for the set of queries that execute frequently over the data and distribute the credit. On the other hand, a coldspot is data with a low quantity of credit, which is therefore considered less important and could be deleted or moved to cheaper and/or less efficient memory.

## References

- [1] Abadi, D., Ailamaki, A., Andersen, D., Bailis, P., Balazinska, M., Bernstein, P., Boncz, P., Chaudhuri, S., Cheung, A., Doan, A., Dong, L., Franklin, M. J., Freire, J., Halevy, A., Hellerstein, J. M., Idreos, S., Kossmann, D., Kraska, T., Krishnamurthy, S., Markl, V., Melnik, S., Milo, T., Mohan, C., Neumann, T., Chin Ooi, B., Ozcan, F., Patel, J., Pavlo, A., Popa, R., Ramakrishnan, R., Ré, C., Stonebraker, M., and Suciu, D. (2020). The seattle report on database research. *SIGMOD Rec.*, 48(4):44–53.
- [2] Alawini, A., Davidson, S. B., Hu, W., and Wu, Y. (2017). Automating data citation in citedb. *PVLDB*, 10(12):1881–1884.
- [3] Alawini, A., Davidson, S. B., Silvello, G., Tannen, V., and Wu, Y. (2018). Data citation: A new provenance challenge. *IEEE Data Eng. Bull.*, 41(1):27–38.
- [4] Altman, M., Borgman, C. L., Crosas, M., and Martone, M. (2015). An Introduction to the Joint Principles for Data Citation. *Bulletin of the Association for Information Science and Technology*, 41(3):43–45.
- [5] Baggerly, K. (2010). Disclose all data in publications. *Nature*, 467(7314):401–401.
- [6] Bechhofer, S., Buchan, I. E., De Roure, D., Missier, P., Ainsworth, J. D., Bhagat, J., Couch, P. A., Cruickshank, D., Delderfield, M., Dunlop, I., Gamble, M., Michaelides, D. T., Owen, S., Newman, D. R., Sufi, S., and Goble, C. A. (2013). Why linked data is not enough for scientists. *Future Gener. Comput. Syst.*, 29(2):599–611.
- [7] Belter, C. W. (2014). Measuring the Value of Research Data: A Citation Analysis of Oceanographic Data Sets. *PLoS ONE*, 9(3):e92590.
- [8] Bertin-Mahieux, T., Ellis, D., Whitman, B., and Lamere, P. (2011). The million song dataset. In *Proceedings of the 12th International Conference on Music Information Retrieval (ISMIR 2011)*, pages 591–596.
- [9] Borgman, C. L. (2016). Data Citation as a Bibliometric Oxymoron. In Sugimoto, C. R., editor, *Theories of Informetrics and Scholarly Communication*, pages 93–116. De Gruyter Mouton.



- 1083 [10] Buneman, P. (2006). How to cite curated databases and how to make  
1084 them citable. In *18th International Conference on Scientific and Statistical*  
1085 *Database Management, SSDBM*, pages 195–203. IEEE Computer Society.
- 1086 [11] Buneman, P., Christie, G., Davies, J. A., Dimitrellou, R., Harding, S. D.,  
1087 Pawson, A. J., Sharman, J. L., and Wu, Y. (2020). Why data citation isn’t  
1088 working, and what to do about it. *Database J. Biol. Databases Curation*,  
1089 2020.
- 1090 [12] Buneman, P., Davidson, S. B., and Frew, J. (2016). Why data citation  
1091 is a computational problem. *Commun. ACM*, 59(9):50–57.
- 1092 [13] Buneman, P., Khanna, S., and Tan, W. C. (2001). Why and where: A  
1093 characterization of data provenance. In *Database Theory - ICDT 2001,*  
1094 *8th International Conference*, pages 316–330.
- 1095 [14] Buneman, P. and Silvello, G. (2010). A rule-based citation system for  
1096 structured and evolving datasets. *IEEE Data Eng. Bull.*, 33(3):33–41.
- 1097 [15] Callaghan, S., Donegan, S., Pepler, S., Thorley, M., Cunningham, N.,  
1098 Kirsch, P., Ault, L., Bell, P., Bowie, R., Leadbetter, A. M., Lowry,  
1099 R. K., Moncoiffé, G., Harrison, K., Smith-Haddon, B., Weatherby, a.,  
1100 and Wright, D. (2012). Making Data a First Class Scientific Output:  
1101 Data Citation and Publication by NERC’s Environmental Data Centres.  
1102 *International Journal of Digital Curation*, 7(1):107–113.
- 1103 [16] Candela, L., Castelli, D., Manghi, P., and Tani, A. (2015). Data Jour-  
1104 nals: A Survey. *Journal of the Association for Information Science and*  
1105 *Technology*, 66(9):1747–1762.
- 1106 [17] Cheney, J., Chiticariu, L., and Tan, W. (2009). Provenance in databases:  
1107 Why, how, and where. *Foundations and Trends in Databases*, 1(4):379–  
1108 474.
- 1109 [18] CODATA-ICSTI Task Group on Data Citation Standards and Practices  
1110 (2013). *Out of Cite, Out of Mind: The Current State of Practice, Policy,*  
1111 *and Technology for the Citation of Data*, volume 12.
- 1112 [19] Cousijn, H., Feeney, P., Lowenberg, D., Presani, E., and Simons, N.  
1113 (2019). Bringing citations and usage metrics together to make data count.  
1114 *Data Science Journal*, 18(1).

- 1115 [20] Cronin, B. (1984). *The citation process. The role and significance of*  
1116 *citations in scientific communication*. London: Taylor Graham.
- 1117 [21] Cronin, B. (2001). Hyperauthorship: A postmodern perversion or evi-  
1118 dence of a structural shift in scholarly communication practices? *JASIST*,  
1119 52(7):558–569.
- 1120 [22] Cui, Y., Widom, J., and Wiener, J. L. (2000). Tracing the lineage of  
1121 view data in a warehousing environment. *ACM Trans. Database Syst.*,  
1122 25(2):179–227.
- 1123 [23] Davidson, S. B., Deutch, D., Milo, T., and Silvello, G. (2017). A model  
1124 for fine-grained data citation. In *CIDR 2017, 8th Biennial Conference on*  
1125 *Innovative Data Systems Research*. [www.cidrdb.org](http://www.cidrdb.org).
- 1126 [24] Dosso, D. and Silvello, G. (2020). Data credit distribution: A  
1127 new method to estimate databases impact. *Journal of Informetrics*,  
1128 14(4):101080.
- 1129 [25] Dubernet, M. L., Antony, B. K., Ba, Y. A., et al. (2016). The vir-  
1130 tual atomic and molecular data centre (VAMDC) consortium. *Journal of*  
1131 *Physics B: Atomic, Molecular and Optical Physics*, 49(7):074003.
- 1132 [26] Fang, H. (2018). A discussion of citations from the perspective of the  
1133 contribution of the cited paper to the citing paper. *JASIST*, 69(12):1513–  
1134 1520.
- 1135 [27] Force, M., Robinson, N., Matthews, M., Auld, D., and Boletta, M.  
1136 (2016). Research data in journals and repositories in the web of science:  
1137 Developments and recommendations. *Bulletin of IEEE Technical Com-*  
1138 *mittee on Digital Libraries, Special Issue on Data Citation*, 12(1):27–30.
- 1139 [28] Garfield, E. (1999). Journal impact factor: a brief review. *Can. Med.*  
1140 *Assoc.*, 979-980.
- 1141 [29] Gößwein, B., Miksa, T., Rauber, A., and Wagner, W. (2019). Data  
1142 identification and process monitoring for reproducible earth observation  
1143 research. In *2019 15th International Conference on eScience (eScience)*,  
1144 pages 28–38. IEEE.

- 1145 [30] Green, T. J., Karvounarakis, G., and Tannen, V. (2007). Provenance  
1146 semirings. In *Proceedings of the twenty-sixth ACM SIGMOD-SIGACT-*  
1147 *SIGART symposium on Principles of database systems*, pages 31–40. ACM.
- 1148 [31] Harding, S. D., Sharman, J. L., Faccenda, E., Southan, C., Pawson,  
1149 A. J., Ireland, S., Gray, A. J. G., Bruce, L., Alexander, S. P. H., Anderton,  
1150 S., Bryant, C., Davenport, A. P., Doerig, C., Fabbro, D., Levi-Schaffer, F.,  
1151 Spedding, M., Davies, J. A., and Nc-Iuphar (2018). The IUPHAR/BPS  
1152 guide to PHARMACOLOGY in 2018: updates and expansion to encom-  
1153 pass the new guide to IMMUNOPHARMACOLOGY. *Nucleic Acids Re-*  
1154 *search*, 46(Database-Issue):D1091–D1106.
- 1155 [32] Hartley, J. (2017). Authors and their citations: a point of view. *Scien-*  
1156 *tometrics*, 110(2):1081–1084.
- 1157 [33] Hey, T., Tansley, S., and Tolle, K. M. (2009). Jim Gray on eScience: a  
1158 transformed scientific method.
- 1159 [34] Honor, L. B., Haselgrove, C., Frazier, J. A., and Kennedy, D. N. (2016).  
1160 Data citation in neuroimaging: proposed best practices for data identifi-  
1161 cation and attribution. *Frontiers in neuroinformatics*, 10:34.
- 1162 [35] Joshi-Tope, G., Gillespie, M., Vastrik, I., D’Eustachio, P., Schmidt, E.,  
1163 de Bono, B., Jassal, B., Gopinath, G. R., Wu, G. R., Matthews, L., Lewis,  
1164 S., Birney, E., and Stein, L. (2005). Reactome: a knowledgebase of bio-  
1165 logical pathways. *Nucleic Acids Research*, 33(Database-Issue):428–432.
- 1166 [36] Katz, D. (2014). Transitive credit as a means to address social and  
1167 technological concerns stemming from citation and attribution of digital  
1168 products. *Journal of Open Research Software*, 2(1).
- 1169 [37] Katz, D. S., Hong, N., Clark, T., Fenner, M., and Martone, M. (2020).  
1170 Software and data citation. *Computing in Science & Engineering*, 22 (2):4–  
1171 7.
- 1172 [38] Kosten, J. (2016). A classification of the use of research indicators.  
1173 *Scientometrics*, 108(1):457–464.
- 1174 [39] Lawrence, B., Jones, C., Matthews, B., Pepler, S., and Callaghan, S.  
1175 (2011). Citation and Peer Review of Data: Moving Towards Formal Data  
1176 Publication. *International Journal of Digital Curation*, 6(2):4–37.

- 1177 [40] Martone, M. (2014). Joint declaration of data citation principles.  
1178 *FORCE11. San Diego CA. Data Citation Synthesis Group*. [https://www.](https://www.force11.org/datacitationprinciples)  
1179 [force11.org/datacitationprinciples](https://www.force11.org/datacitationprinciples), online September 2020.
- 1180 [41] Meho, L. I. and Yang, K. (2007). Impact of data sources on citation  
1181 counts and rankings of LIS faculty: Web of science versus scopus and  
1182 google scholar. *Journal of the american society for information science*  
1183 *and technology*, 58(13):2105–2125.
- 1184 [42] Milo, T. (2019). Getting rid of data. *Journal of Data and Information*  
1185 *Quality (JDIQ)*, 12(1):1–7.
- 1186 [43] Nosek, B. A., Alter, G., Banks, G. C., Borsboom, D., Bowman, S. D.,  
1187 Breckler, S. J., Buck, S., Chambers, C. D., Chin, G., Christensen, G.,  
1188 Contestabile, M., Dafoe, A., Eich, E., Freese, J., Glennerster, R., Goroff,  
1189 D., Green, D. P., Hesse, B., Humphreys, M., Ishiyama, J., Karlan, D.,  
1190 Kraut, A., Lupia, A., Mabry, P., Madon, T., Malhotra, N., Mayo-Wilson,  
1191 E., McNutt, M., Miguel, M., Paluck, E. L., Simonsohn, U., Soderberg, C.,  
1192 Spellman, B. A., Turitto, J., VandenBos, G., Vazire, S., Wagenmakers,  
1193 E. J., Wilson, R., and Yarkoni, T. (2015). Promoting an open research  
1194 culture. *Science*, 348(6242):1422–1425.
- 1195 [44] Peters, I., Kraker, P., Lex, E., Gumpenberger, C., and Gorraiz, J.  
1196 (2016). Research data explored: An extended analysis of citations and  
1197 altmetrics. *Scientometrics*, 107(2):723–744.
- 1198 [45] Pröll, S. and Rauber, A. (2013). Scalable data citation in dynamic, large  
1199 databases: Model and reference implementation. In *Proceedings of the*  
1200 *2013 IEEE International Conference on Big Data*, pages 307–312. IEEE.
- 1201 [46] Rauber, A., Ari, A., van Uytvanck, D., and Pröll, S. (2016). Identi-  
1202 fication of Reproducible Subsets for Data Citation, Sharing and Re-Use.  
1203 *Bulletin of IEEE Technical Committee on Digital Libraries, Special Issue*  
1204 *on Data Citation*, 12(1):6–15.
- 1205 [47] Rauber, A., Asmi, A., van Uytvanck, D., and Proell, S. (2015). Data  
1206 citation of evolving data: Recommendations of the working group on data  
1207 citation (wgdc). *Result of the RDA Data Citation WG*, 20.
- 1208 [48] Silvello, G. (2018). Theory and practice of data citation. *J. Assoc. Inf.*  
1209 *Sci. Technol.*, 69(1):6–20.

- 1210 [49] Simmhan, Y., Plale, B., and Gannon, D. (2005). A survey of data  
1211 provenance in e-science. *SIGMOD Record*, 34(3):31–36.
- 1212 [50] Spengler, S. (2012). Data Citation and Attribution: A Funder’s Per-  
1213 spective. In of Sciences’ Board on Research Data, N. A. and Information,  
1214 editors, *Report from Developing Data Attribution and Citation Practices*  
1215 *and Standards: An International Symposium and Workshop*, pages 177–  
1216 178. National Academies Press: Washington DC.
- 1217 [51] Wilkinson, M. D., Dumontier, M., Aalbersberg, I. J., Appleton, G.,  
1218 Axton, M., Baak, A., Blomberg, N., Boiten, J., da Silva Santos, L. B.,  
1219 Bourne, P. E., et al. (2016). The fair guiding principles for scientific data  
1220 management and stewardship. *Scientific data*, 3.
- 1221 [52] Wu, Y., Alawini, A., Davidson, S. B., and Silvello, G. (2018). Data  
1222 citation: Giving credit where credit is due. In *Proceedings of the 2018*  
1223 *International Conference on Management of Data, SIGMOD*, pages 99–  
1224 114.
- 1225 [53] Zeng, T., Wu, L., Bratt, S., and Acuna, D. E. (2020). Assigning credit to  
1226 scientific datasets using article citation networks. *Journal of Informetrics*,  
1227 14(2).
- 1228 [54] Zou, C. and Peterson, J. B. (2016). Quantifying the scientific output of  
1229 new researchers using the zp-index. *Scientometrics*, 106(3):901–916.
- 1230 [55] Zwölf, C. M., Moreau, N., and Dubernet, M.-L. (2016). New Model for  
1231 Datasets Citation and Extraction Reproducibility in VADMC. *Journal of*  
1232 *Molecular Spectroscopy*, 327:122–137.