



SDRF (Scientific Data Repository Framework)

Improving Data Sets' Machine Readability and Interoperability with Published Research

A New Protocol for Sharing Scientific Data

Linguistic Technology Systems (LTS) is developing **SDRF** to encompass data models, publishing guidelines, and code libraries for deploying open-access research data sets associated with scientific publications. Nowadays there are many general-purpose and domain-specific portals hosting scientific data; there are also several available formats for describing and encoding scientific data, such as Research Objects, schema.org/Dataset, Digital Curation Center (**DCC**), **SciDATA**, **BIOCODER**, and **MIBBI** (Minimum Information for Biological and Biomedical Investigations). The purpose of **SDRF** is to merge these different data-set formats into a unified, overarching standard which can be adapted to different publishing houses and pipelines.

Shaping Protocols to Conform to Current Specifications in Publishing

In order to conform to current specifications — such as FAIRsharing (Findable, Accessible, Interoperable, Reusable) or the Bill and Melinda Gates Foundation guidelines for authors (<https://gatesopenresearch.org/for-authors/data-guidelines>) — proper protocols must be implemented at several stages of the publishing process, in particular (1) publications should provide clear descriptions of accompanying data sets and where that data is hosted; (2) publication repositories should make data-set links and metadata clearly visible on web pages where documents are read or previewed; (3) data sets themselves need to include metadata and supporting files which help researchers properly access, visualize, and reuse the data; and (4) data sets need to be connected with software which has the correct features to load and display the relevant raw data files. **SDRF** will include technology applicable to each of these four facets of the publishing and data-sharing pipeline in order to conform to current standards.

It is important to emphasize that data sets are only truly valuable if they are machine-readable and seamlessly integrated into domain-specific software ecosystems. Scientists who examine and reuse published data sets are generally researchers doing technical work in a field closely related to that of the original authors; in many cases there are specialized software applications, computational methods, algorithms, and research protocols which are endemic to the relevant subject areas. When sharing research data, accordingly, publishers/authors should make it as easy as possible for scientists to examine the data within the digital ecosystem that they utilize for their own research. This often implies that document viewers — e.g., **PDF** viewers and/or **HTML** pages on publisher portals — should be ideally interconnected with scientific applications so that scientists, when reading books/articles, can seamlessly launch domain-specific software and visualize/examine associated data sets. Unfortunately, most scientific software does not incorporate code libraries to parse metadata (summarizing file types, download instructions, etc.) describing open-access data sets. This can be addressed by providing plugins or extensions that add data-set-accession capabilities to existing scientific software, so that publisher repositories and data-hosting repositories can be made truly interoperable with scientific applications.

To demonstrate how such plugins can work, as well as other facets of the data-publication process facilitated via **SDRF**, this paper will review two case-studies addressing research in the academic literature, each involving papers that have been linked with multiple data sets (data which was either reused or newly created during the course of the research described).

First Case Study: “Parkinson’s Disease Diagnosis: The Effect of Autoencoders on Extracting Features from Vocal Characteristics,” by Ashena Gorgan Mohammadi, Pouya Mehralian, Amir Naseri, and Hedieh Sajedi (International Journal of Speech Technology, pending review)

This case study demonstrates a scenario where an article reuses multiple pre-existing data sets. The article examines Parkinson’s Disease symptomology from multiple perspectives, including gait (loss of motor function), speech impairment, and bioimaging (**MRIs**). The authors apply Machine Learning to data sets focused on these three different diagnostic areas, in an effort to advance research to refine Parkinson’s predictors and diagnoses. The overall information can be summarized as follows:

- 1. Gait Data:** this data was primarily drawn from a PhysioNet data set (<https://physionet.org/content/gaitpdb/1.0.0/>) obtained via sensors attached to subject’s feet as they walked (the study includes both Parkinson’s patients and healthy controls). This sensor data is provided as a collection of text files, each file corresponding to one patient (or control subject), with each line in a file representing a single time snapshot. The lines are divided into space-separated columns, each representing force exerted on a single sensor, plus two additional columns calculating total force on the left-foot and right-foot sensors respectively. This data set also includes demographic and clinical information for each patient in a spreadsheet format. The authors also use an additional source of gait data derived from a more recent (2019) study whose data is available only upon request (see <https://www.nature.com/articles/s41598-019-53656-7#MOESM1>).
- 2. Speech Data:** this data was primarily drawn from a data set hosted by the University of California Irvine Machine Learning Repository (<https://archive.ics.uci.edu/ml/datasets/Parkinsons>). The central information is a **CSV** file, where each line represents a single voice recording from one of 31 subjects (consisting of 23 Parkinson’s patients and 8 healthy controls). Each subject made multiple recordings. The individual lines in the **CSV** file present a quantitative model of subjects’ speech via a collection of acoustic features/attributes. A similar “telemonitoring” data set from the same archive was used to study how the analysis of Parkinson’s-related speech data may be applicable to samples obtained via devices such as smartphones.
- 3. Radiological Data:** this data is not immediately available for reuse, but requires special (non-commercial) authorization from the Parkinson’s Progressive Markers Initiative (<https://www.ppmi-info.org/>) or making a request of corresponding authors of referenced papers introducing the relevant data (<https://www.frontiersin.org/articles/10.3389/fnins.2019.00874/full#h6>).¹ Although this data is derived from bioimaging (so that the underlying raw data files are radiological) the authors of the IJST paper under review utilize the data in a more structured form, building off of image-feature extraction already performed when the data sets were first published by the prior authors. However, the republished unified data set itself (to appear in conjunction with the IJST submission) might include code to allow researchers to reproduce the original analytic workflow if desired: for instance, among other things, we can enable the implementations published via <https://biomedica.doc.ic.ac.uk/software/malp-em/> to be embedded as a **CAPTk** module (see in particular https://cbica.github.io/CaPTk/tr_integration.html#tr_cppIntegration).
- 4. Python Code Repository:** the authors also provide Python code, hosted on GitHub, which they used to analyze these various data sources.

Unifying the Data Sets into a Single Package

In this article pending review, the authors summarize the data sets in a table within the main text (see Figure 1 here) and, in their bibliography, they cite these data sets either directly or by referencing publications where the relevant data sets are described (see Figure 2 here). Doing so properly credits authorship to the researchers who curated the data sets, and it gives readers a means to locate the raw data. However, accessing and working with the raw data is inconvenient from a reader’s point of view without most or all of the data sets being repackaged into a *single* archive that could be hosted and downloaded as one unit. Obtaining raw data from the resources identified in the bibliography

¹When the unified data set is published, we will request permission to include a copy of the original data set to spare future readers from having to request this data on their own.



3.2. Data preprocessing and Feature extraction

Since the data is extracted using different signal processing methods, it ranges diversely. This contributes to inadequate learning procedures. Consequently, to get started with the task, we apply rescaling or in a more common term, min-max normalization. Using this method, the data is scaled in a specified range, and here we scale the features to the [0, 1] range.

Table 1: A summary of reviewed datasets.

Data type	Description	Study
Brain MRI	In this retrospective study, we enrolled 56 patients and 28 healthy control subjects.	[9]
GAIT	This database contains measures of gait from 93 patients with idiopathic PD (mean age: 66.3 years; 63% men), and 73 healthy controls (mean age: 66.3 years; 55% men).	[10]
GAIT	303 subjects were recruited from the "Incidence of Cognitive Impairment in Cohorts with Longitudinal Evaluation-GAIT" (ICICLE-GAIT) study.	[11]
Vocal Features	UCI Parkinson's Disease Classification.	[13]
Vocal Features	The dataset range of biomedical voice measurements from 31 people, where 23 people are showing Parkinson's disease.	[14]
Vocal Features	-UCI Parkinson Speech Dataset with Multiple Types of Sound Recordings Data Set	[15], [17]
Vocal Features	The database consisted of 23 columns and 197 rows. The dataset was created by Mark Little of the University of Oxford, in collaboration with the National Centre for Voice and Speech, Denver, Colorado, who recorded the speech signals. This dataset is composed of a range of biomedical voice measurements from 31 people, 23 with Parkinson's disease (PD). Each column in the table is a particular voice measure	

The authors provide citations for data sets used in their analyses ...

Figure 1: Table Listing Analyzed Data Sets in the Parkinson’s Article

requires several steps — for instance, the PhysioNet sensor data can be downloaded as a zipped folder, while the demographic data attached to it has to be downloaded separately. Furthermore, some of the information obtained from **MRI** and speech analysis (reported in papers cited as data sources for the IJST submission) is provided as supplemental materials within the secondary papers; this requires readers to browse one at a time through each of the relevant articles so as to find downloadable links (see Figure 4). In short, piecing all the source data together puts the onus on readers to manually inspect multiple web resources and to manually interconnect files once they are downloaded.

Another complicating factor is that certain information present in the data sets is implicit within how the data sets are organized, requiring extra effort to extract this information in a machine-readable manner. For instance, the PhysioNet sensor data uses a file-naming convention which encodes several pieces of information in the file names, such as whether the file presents a Parkinson’s patient or a control subject (see Figure 3). Though by examining file names it would be possible to construct a table with additional information providing context for the file contents, such information is not directly included within the PhysioNet data set; it needs to be extracted by computer code.

Constructing Machine-Readable Supplemental Archives

This collection of data sets serves as an example of how technologies such as **SDRF** can fill the gap between publication/data repositories and scientific computing, making scientific data more “**FAIR**” (Findable, Accessible, Interoperable, Reusable). Upon publication of this submission in IJST (pending peer review) the disparate open-access data from the article’s secondary sources would be provided as a single **SDRF** archive. This archive would provide *machine-readable* access to information spread across multiple sources, translated into a common file format. In general, **SDRF** encourages and implements features to help data sets conform to **FAIR** and related standards, such as (1) bundling multiple data sets into a single archive; (2) migrating data to general-purpose representations wher-



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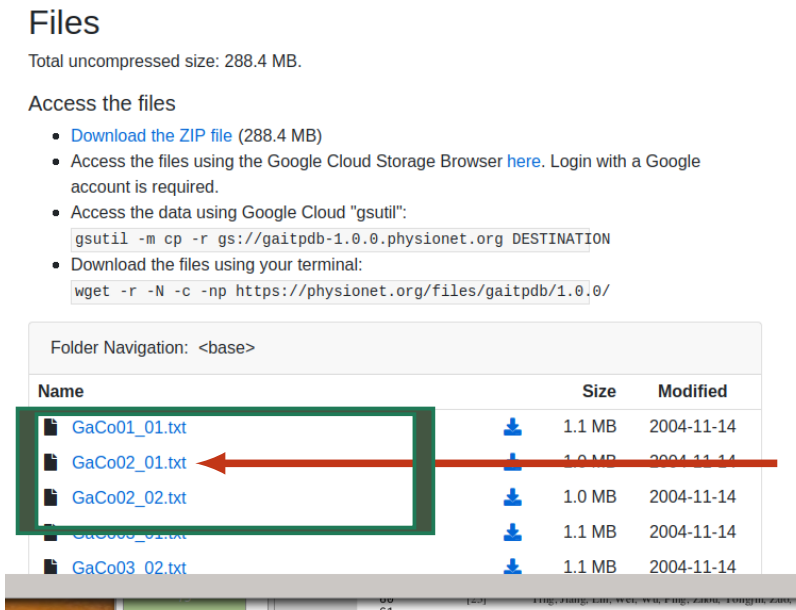
Some data sets are directly available through the bibliography; others have to be located by reading the cited articles.

Figure 2: Bibliography (With Data Set Hyperrefs) in the Parkinson's Article

ever possible — formats such as **XML**, **HDF5**, **ARFF**, or **DICOM**; (3) providing meta-data in multiple formats (**DCC**, **schema.org/Dataset**, **MIBBI**, **BIOCORDER**, etc.) to be compatible with different organizations' platforms; (4) identifying one or more "preferred applications" for examining/reusing the published data; (5) explicitly representing information encoded via file-names; (6) bundling raw data, meta-data, and (where possible) machine-readable article text into a single resource, which **SDRF** calls a "Supplemental Archive;" and (7) annotating the data sets to support microcitations that granularly link the publication to its associated Supplemental Archive.

Once a Supplemental Archive has been downloaded, an important question for any **SDRF** archive is how researchers will productively access the data. Unlike the Flow Cytometry use-case discussed below, the Parkinson's archive spans several scientific disciplines; as such, there is no obvious application which could be preferred by default for examining the data files. As a fallback option, **SDRF** is designed to present data sets via **Qt Creator**, a **C++** Integrated Development Environment associated with the **Qt** application-development framework. **SDRF** includes code libraries to represent research meta-data as **C++** objects; these libraries can be opened as **Qt** projects. These may be supplemented with separate libraries extracting and managing information specific to individual data sets. In particular, the Supplemental Archive for the Parkinson's article under peer review would provide **C++** classes encapsulating spreadsheet-like data (whether originally in **.xls**, **CSV**, or space-delimited formats) republished by the Journal in the unified data set.

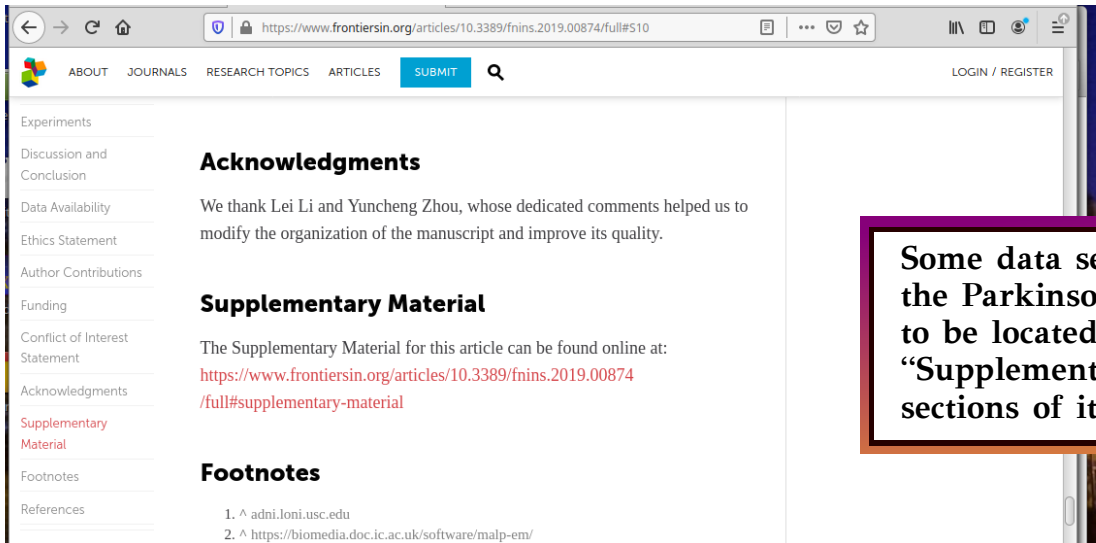
An additional concern for **SDRF** archives is how to properly annotate publications and data sets side-by-side. In the Parkinson's article, individual **C++** classes encapsulating tabular data serve as convenient microcitation targets: annotations within the relevant **C++** code represent anchors through which the data set may be referenced (on a more precise scale than merely citing the Supplemental Archive as a whole). In some places, individual class attributes can also be linked to lines in the authors' Python source code. On the text side, certain paragraphs within the Parkinson's article can be linked to the corresponding **C++** code annotations. This illustrates **SDRF**'s recommended



Some information in the PhysioNet is encoded in file names, where the initial two letter-pairs and following two number-pairs all provide information about the patient and data source. Unfortunately, encoding data in this manner requires extra computer code when reusing the data base, because the file names need to be analyzed so as to parse the information expressed via the naming conventions.

Figure 3: Extracting Information Encoded in File Names

annotation/microcitation system, where segments in publication texts (identified for instance via $\text{L}^{\text{A}}\text{T}_{\text{E}}\text{X}$ `phantomsection` commands or `JATS statement` tags) are linked to annotations or comments in code and/or raw data files in the Supplemental Archive.



Some data sets analyzed in the Parkinson’s article have to be located by browsing “Supplementary Material” sections of its cited papers.

Figure 4: Indirectly Locating Data Sets from Cited Papers

Second Case Study: “Marked T cell activation, senescence, exhaustion and skewing towards TH17 in patients with COVID-19 pneumonia” from nature.com, 2020

This article presents a use-case with some noteworthy contrasts to the Parkinson’s publication described in the previous section. This Covid-19 paper (<https://www.nature.com/articles/s41467-020-17292-4>) was published along with two data sets comprising Flow Cytometry Standard (FCS) files hosted via the Flow Repository (<http://flowrepository.org/id/FR-FCM-Z2N4> and <http://flowrepository.org/id/FR-FCM-Z2N5>). Links to the data sets (via flowrepository.org pages) are explicitly provided in the publication’s “Data Availability” section. However, researchers still need to perform several steps to manually download the full set of relevant FCS and meta-data files.

One feature of this second use-case is that the technical information in the data sets belong to a single scientific area (Flow Cytometry) and are mostly encoded in a single format (FCS). As such, it is straightforward to identify the kind of software which researchers need to use to visualize the raw data — basically, any application that can parse FCS files. There are a variety of commercial as well as open-source Flow Cytometry (FCM) applications which can be used to access FCS data. Once readers have downloaded the Flow Repository archives, they may individually load the .fcs files to study data which, in the original article, is summarized via figure illustrations.

This workflow nonetheless requires researchers to perform several manual steps before being able



to use the published data. The core problem is that existing Flow Cytometry software does not intrinsically have capabilities to read **PDF** files, locate **FCS** data sets, and interoperate with hosting platforms such as Flow Repository. Employing this use-case as an example with which to illustrate proper alignment between document viewers, publication/data repositories, and scientific software, We are developing a new Flow Cytometry application which *can* interoperate with **PDF** viewers and **SDRF** archives. This application is designed so that, when authors are reading a **PDF** file associated with an **FCS** data set, the **PDF** viewer can automatically launch and signal to the **FCM** application when a reader wishes to download and visualize **FCS** files. In short, the **FCM** application — having received data from a **PDF** viewer which implements an **SDRF** inter-application protocol — will automatically execute download and extraction steps that scientists otherwise would have to perform manually.

Note also that, although most of the relevant data for this Covid-19 article is in **FCS** form, there is, in addition, supplemental clinical information provided in other formats (...). For cases such as these, the LTS **FCM** software includes code libraries allowing researchers to parse non-**FCS** data in standard formats such as **XML**, **HDF5**, or **ARFF**.

This use-case illustrates a general principle: that research data is most convenient for scientists when it is deployed within an infrastructure where portals, document viewers, and scientific applications can seamlessly interoperate. Wherever possible, when researchers are reading published books or articles, they should be able to *automatically* launch the proper scientific application, download data sets, and examine raw data files in the preferred software with only one or two clicks. These steps would be performed automatically as much as is feasible, instead of readers having to waste time with manually finding, downloading, merging/extracting, and then opening data files.

Conclusion

The two case-studies considered here are similar in that each involve articles which are linked to multiple data sets. For maximum convenience, it is optimal for researchers to be able to access this data without performing manual download and merging/extracting actions. There are also some differences between the two case-studies: in particular, the Parkinson's data spans multiple disciplines, whereas the Covid-19 data is more rigorously grounded in Flow Cytometry. As such, the operational requirements for the Covid-19 data, from a reader's point of view, are more clearly delineated: effective integration between the publication and its accompanying data sets is defined by launching Flow Cytometry software while a researcher is reading the publication, allowing the researcher to view the data via software similar to that used to generate/analyze the data while the reported research was being conducted. In the case of the Parkinson's data, in contrast, there is no single application which would seamlessly display the specrum of information considered in that article; as mentioned above, in such cases, **SDRF** would default to using **QT** Creator as a fallback for loading Supplemental Archives where no other software is available.

Regardless of whether one is using **QT** Creator or a domain-specific application, it is preferable that each **SDRF** archive be associated with one or more applications that researchers can use to view data (and extract information) from the archive. Moreover, these applications would ideally be linked to document viewers and also to publisher's portals, so that readers can automatically launch preferred applications and view Supplemental Archives while reading concomitant publications. In order to achieve this, scientific applications need to be augmented with plugins to parse **SDRF** data. To address this need, we are developing an inter-application messaging protocol so that disparate applications with **SDRF** plugins may interoperate. In particular, this protocol would entail **PDF** viewers being able to interoperate with scientific applications so that publications' data sets may be automatically downloaded and visualized via the preferred software.

Our prototype example for an application utilizing such plugins, as mentioned above, is software for Flow Cytometry. We are also working on a prototypic **QT** Creator plugin so that **QT** Creator (as the "default" **SDRF** software) can participate in **SDRF** networks using the same protocol. We will then expand the scope of this protocol via plugins for software in other domains, such as image-analysis, molecular visualization, radiology, **3D** graphics, and so forth.



Future Projections: Operationalizing SDRF

This section will present a concrete outline of the steps necessary to package scientific data into an **SDRF** Supplemental Archive. Data Sets may be published with content and organization compatible with both **SDRF** and other formats, such as Research Object Bundles, so using **SDRF** does not preclude adopting other formats as well. For example, a data set which provides **SDRF** meta-data may also include the **meta-inf** files required by Research Objects. Indeed, it is recommended that authors aim for compatibility with multiple standards, not only **SDRF**. This section, however, will focus on the components of Supplemental Archives that are specific to **SDRF**.

This section will describe two different approaches to using **SDRF**. One approach employs **SDRF** to a limited extent, deferring to older technologies for such basic operations as encoding data or annotating publications. An alternative is to adopt **SDRF** more holistically, adopting experimental or under-development features in the **SDRF** libraries. When discussed in the following outline, these features will usually be characterized as “experimental” or “specific to **SDRF** code libraries.”

Formulating Scientific Data Repository Models for Individual Publishing Environments

SDRF is intended to interoperate with multiple publisher and data-hosting platforms, each of which have their own requirements and technology. Therefore, the precise contents and organization of a given **SDRF** Supplemental Archive will depend on specifications pertaining the specific repository and hosting platform where the publication and data set, respectively, are to be deposited. To facilitate Supplemental Archive preparation for different environments, **SDRF** seeks to include “Scientific Data Repository Models” (**SDRMs**) which are tailored to individual publishers. The **SDRMs** would encapsulate information pertaining to the specific platforms where the relevant publication and data set are hosted. For example, research funded by the Bill and Melinda Gates Foundation must follow certain guidelines, which are documented whenever an article is submitted to the Gates Open Research portal. Consequently, the specific **SDRF** meta-data for such papers can be designed to notate the concordant research’s adherence to these organizational guidelines. In these situations, the layout and vocabulary for **SDRF** meta-data would be determined at least in part by the **SDRM** germane to the environment where the research is published.

The purpose of an **SDRM** is also to facilitate the implementation of technologies which interoperate with publishers’ platforms — for instance, **API** clients which can request information from document/data portals (e.g., locating articles or data sets via keyword searches), or **PDF** viewers enhanced with capabilities to obtain data sets from hosting platforms (to automatically download data sets upon request from a user reading the corresponding publication). For these use-cases, individual **SDRM** would include information and/or computer code about how to access publishers’ **APIs**, how to download and extract a data set given a digital identifier, and so forth.

Because **SDRF** Supplemental Archives may differ somewhat according to the specific **SDRM** in effect, any description of these archives is provisional. To be precise, **SDRF** assumes a generic **SDRM** — dubbed “**MOSAIC**” — which can be extended or modified in consultation with individual publishers, as required. As such, the following outline will apply intrinsically to the **MOSAIC SDRM**; depending on the platform, the details for individual **SDRF** Supplemental Archives may be different than the generic model presented here.

Standard Components of SDRF Supplemental Archives

The components of an **SDRF** Supplemental Archive can be grouped into several facets, concerning meta-data, raw data, and machine-readable text, respectively.

1. SDRF Meta-Data

Meta-Data Files **SDRF** uses a vocabulary for describing the contents of data sets which merges the data models of several existing formats, such as **DCC** and schema.org/Dataset. **SDRF** recommends encoding this data in **TAGML** (“Text-as-Graph Markup Language”), which

is a very flexible, and computationally powerful representation format.² The **SDRF**-specific libraries include parsers for (an extended version of) **TAGML**.

C++ Files **SDRF** also recommends that authors provide **C++** code to initialize objects representing Supplemental Archive's meta-data. These objects may be directly constructed from the **TAGML** meta-data files, or authors may choose to customize the **C++** logic as desired. For lab-based research, authors may employ **BIOCORDER**, which uses **C++** code to notate research protocols and workflows (the **SDRF** libraries include a modified version of **BIOCORDER**). Moreover, for data sets whose "preferred application" for opening raw data files is implemented in **C++** (as is the case for many, if not most, scientific-computing applications), the Supplemental Archive may include plugins or extensions to these applications. In general, using **C++** objects specific to Supplemental Archive meta-data as a basis, authors (or programmers coding on authors' behalf) may introduce additional **C++** code demonstrating or enabling analysis, visualization, or application-integration for the Supplemental Archive's raw data.

SDRM-Specific Meta-Data Either via data files or computer code, information or capabilities specific to the platforms where papers and Supplemental Archives are hosted may be presented in conjunction with the applicable **SDRMs**. For instance, **C++** code distributed with the archive may include procedures for accessing **APIs** of the hosting service where the archive is deposited.

Qt Project Files As explained earlier, **SDRF** defaults to using **QT** Creator as an application with which to examine data sets, if there is no obvious "preferred application" based on the research topic.

- 2. Raw Data:** As mentioned above, **SDRF** recommends that raw data be encoded in general-purpose formats such as **ARFF**, **HDF5**, **XML**, or **TAGML** (as compared to formats such as **.xls**, which are associated with specific applications). An experimental "Hypergraph Exchange Format" (**HGXF**) may also be used.

An exception to the guidelines for general-purpose formats is when data should be presented in optimized formats endemic to a certain scientific field, such as **FCS** files for Flow Cytometry, or **DICOM** files for bioimaging. In these cases, it is recommended to employ such formats for most of the raw data files but also to construct summaries of the data, which may facilitate properly loading and accessing the domain-specific files, represented in a more general-purpose format.

- 3. Annotated Manuscripts** Where authors have permission to share full-text versions of their books/articles, **SDRF** recommends that a machine-readable representation of these publications, in formats such as **TAGML** or **XML**, be included in the Supplemental Archive (**SDRF** has an experimental "Hypergraph Text Encoding Protocol" which may be utilized as well). This machine-readable manuscript may then be annotated and cross-referenced with raw data and/or code files also included in the Supplemental Archive. An experimental **SDRF** **L^AT_EX** package provides an annotation framework which not only marks locations in document text, but also encodes **PDF** viewport coordinates for use by specialized **PDF** viewers which implement an **SDRF** protocol for integrating **PDF** viewers with scientific applications.

²**TAGML** is "powerful" in the sense that, with suitable mappings between their disparate syntactic expressions, **TAGML** represents a superset of **XML** and other common data-representation languages, such as **JSON**.

