A comprehensive framework to capture the arcana of neuroimaging analysis

Thomas G. $\operatorname{Close}^{1,2*} \cdot \operatorname{Phillip} G. D.$ $\operatorname{Ward}^{1,3,4} \cdot \operatorname{Francesco} \operatorname{Sforazzini}^1 \cdot \operatorname{Wojtek} \operatorname{Goscinski}^6 \cdot \operatorname{Zhaolin} \operatorname{Chen}^{1,5} \cdot \operatorname{Gary} F. \operatorname{Egan}^{1,3,4}$

the date of receipt and acceptance should be inserted later

Abstract Mastering the "arcana of neuroimaging analysis", the obscure knowledge required to apply an appropriate combination of software tools and parameters to analyse a given neuroimaging dataset, is a time consuming process. Therefore, it is not typically feasible to invest the additional effort required generalise workflow implementations to accommodate for the various acquisition parameters, data storage conventions and computing environments in use at different research sites, limiting the reusability of published workflows.

We present a novel software framework, Abstraction of Repository-Centric ANAlysis (Arcana), which enables the development of complex, "end-to-end" workflows that are adaptable to new analyses and portable to a wide range of computing infrastructures. Analysis templates for specific image types (e.g. MRI contrast) are implemented as Python classes, which specify a range of potential derivatives and analysis methods. Arcana retrieves data from imaging repositories, which can be BIDS datasets, XNAT instances or plain directories, and stores selected derivatives back into a repository with provenance data for reuse by subsequent analyses. Workflows are constructed using Nipype and can be executed on local workstations or in high performance computing environments without alteration. Generic analysis methods can be consolidated within a common base classes to facilitate code-reuse and collaborative development,

Thomas G. Close

E-mail: tom.close@monash.edu

¹Monash Biomedical Imaging, Monash University, Melbourne, Australia

² Australian National Imaging Facility, Australia

 $^{^3}$ Australian Research Council Centre of Excellence for integrative Brain Function, Melbourne, Australia

 $^{^4\}mathrm{Monash}$ Institute of Cognitive and Clinical Neurosciences, Monash University, Melbourne, Australia

 $^{^5\}mathrm{Department}$ of Electrical and Computer Systems Engineering, Monash University, Melbourne, Australia

 $^{^6\}mathrm{Monash}$ e
Research Centre, Monash University, Melbourne, Australia

which can be specialised for study-specific requirements via class inheritance. Therefore, Arcana provides a platform to develop unified neuroimaging workflows that can be reused across many research sites and studies.

Keywords Neuroimaging \cdot Workflows \cdot Repository \cdot Reproducibility \cdot Reusability \cdot Large-scale \cdot Python

Introduction

Despite the availability of well-established neuroimaging analysis packages (Cox, 1996; Smith et al., 2004; Friston, 2007; Tournier et al., 2012), the arcana of neuroimaging analysis is substantial due to the range of available tools, tuneable parameters, and imaging sequences involved (Cusack et al., 2015). The distribution of complete "end-to-end" workflows, from acquired data to publication results, is necessary for routine reproduction because of the effort required to accurately exactly reimplement such analyses (Kennedy, 2018). It is also difficult to adapt existing workflows to new studies without detailed knowledge of their design. Therefore, flexible, portable and complete workflows are important to promote reproduction and code reuse in neuroimaging research.

A barrier to designing portable and complete workflows is the heterogeneity of data storage conventions (Marcus et al., 2007; Das et al., 2012; Gorgolewski et al., 2016). To address this, the emerging Brain Imaging Data Standard (BIDS) (Gorgolewski et al., 2016) provides a way to standardise the storage of neuroimaging data in file system directories. BIDS specifies strict file and directory naming conventions, which facilitate the design of portable BIDS Apps (bids-apps.neuroimaging.io). However, for research groups with sufficient informatics support, software-managed repositories (Marcus et al., 2007; Das et al., 2012) can provide additional features, such as flexible access-control and automated pipelines. Therefore, for published workflows, the choice of repository should be transparent in order to maximise their audience.

While neuroimaging analyses are generally amenable to standardisation (Kennedy, 2018), minor modifications are often required to accommodate idiosyncrasies of the acquisition protocols in use at different sites (Esteban et al., 2018). Therefore, workflows may require conditional logic in construction or execution to be portable. Nipype is a flexible Python framework for neuroimaging analysis in which workflows are constructed programmatically in Python (Gorgolewski et al., 2011). Programmatic construction allows for rich control-flow logic that is not readily available in alternative workflow frameworks (Cusack et al., 2015; Achterberg et al., 2016; Amstutz et al., 2016), and has been used to implement workflows that are robust to differences in fMRI protocols across a large number of sites (Esteban et al., 2018).

The trend towards large multi-site and multi-contrast datasets collected over a number of years (Van Essen et al., 2012; Thompson et al., 2014; Sudlow et al., 2015) presents additional challenges to workflow design. Analysis packages are constantly being developed and improved, so the state-of-the-art

workflow for a particular analysis can change over time. Therefore, it is challenging to ensure workflows are applied consistently over the course of long studies (Cusack et al., 2015).

While analysis workflows for different contrasts and modalities are typically implemented independently, they can share common processing steps (e.g. non-linear registration to standard space, surface parcellation) and their outputs may need to be integrated to produce publication results. For large scale studies, which are typically processed on the cloud or high-performance computing (HPC) clusters, rerunning common segments can lead to significant increases in computation time and therefore project cost. In addition, duplication of processing segments increases time for manual quality control (QC), making the reuse of intermediate derivatives a practical requirement for some large studies (Schreiber et al., 2018).

To maximise the reusability of neuroimaging workflows and avoid frequent reimplementation of standard analyses, workflow implementations should be flexible, extensible and applicable to a wide range of storage systems. In addition, in order to promote routine reproduction of neuroimaging studies, published workflow implementations should include the complete procedure, from acquired data to publication results. However, ensuring workflow implementations are flexible, portable and complete adds a high degree of complexity and effort to the design process.

Our objective was to extract common elements of repository-centric work-flow design into an abstract framework to make it practical to implement flexible, portable and complete workflows for a wide range of neuroimaging analyses. Abstraction of Repository-Centric ANAlysis (Arcana) (arcana. readthedocs.io) is a Python framework for designing complex workflows in which modular Nipype pipelines operate on data stored in various repositories. Intermediate derivatives are derived on demand, checking against stored provenance for required updates. Analyses can be applied to XNAT, BIDS and plain-directory repositories, and using Nipype's execution plugins, run on workstations or be submitted as batch jobs to HPC schedulers. Arcana's architecture, with programmatic workflow construction—yet clear delineation between analysis design and application—facilitates the implementation of complex workflows that are portable and complete.

The utility of the Arcana framework is demonstrated by the implementation of analysis suites for T1, T2* and diffusion weighted MRI (dMRI) data and the application of dMRI tractogram (Tournier et al., 2012) and composite vein image (Ward et al., 2018) workflows to data collected from a healthy subject.

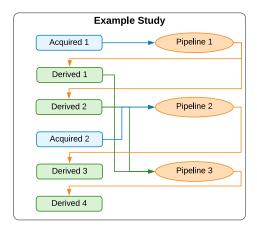


Fig. 1 Example study. Blue boxes represent acquired (input) data (filesets or fields) stored in a repository and green derivatives from that data stored alongside the original data. Orange ovals are pipelines that operate on data in the repository to derive the derivatives. Arrows represent data flows, i.e., inputs and outputs to pipelines

Methods

Framework overview

The separation of analysis design and application in the Arcana framework follows the conceptual divide between classes and objects in Object-Oriented (OO) software design. *Study* classes encapsulate types of data, such as scans of a specific imaging contrast or modality, with the suite of analysis methods that can be performed on them. Study objects apply the analysis suite defined in the Study class to a specific dataset.

The set of acquired data, the derivatives that can be derived from it, and methods that construct pipelines to derive the derivatives, are linked together by the data specification table of the Study class (Figure 1). Likewise, free parameters used in pipeline construction are defined in the class' parameter specification table. Class inheritance can be used to specialise analysis suites by overriding entries in the specification tables or pipeline constructor methods. Analysis suites for multi-modal data can be implemented by combining Study classes within MultiStudy classes.

Analysis methods defined by a Study class are applied to a specific dataset by instantiating an object of the class and requesting a derivative listed in the class' data specification table. At initialisation, a Study object is passed references to a *Repository*, a *Processor*, and an *Environment*, which define where and how data is stored and processed. When a derivative is requested, a Study object queries the Repository for intermediate derivatives that can be reused before constructing a workflow to produce the requested derivative. The manner in which the workflows are executed (i.e. single/multi-process or via SLURM scheduler) is specified by the Processor and software modules

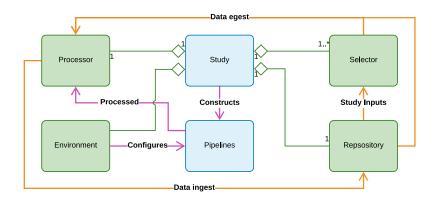


Fig. 2 Unified Modelling Language (UML) diagram of information flow in the Arcana framework. Boxes: Python classes, blue=analysis-design, green=analysis-application. Arrows: orange=data, magenta=workflow description, diamond=aggregated-in. Study classes construct analysis pipelines, which are sent to the *Processor* to be processed. Input data is selected by *Selector* objects and pulled to the compute environment to be processed along with existing intermediate derivatives. After the derivatives are pushed back to the repository.

required by the analysis are loaded by the Environment. Selected workflow products are stored back in the repository for reuse by subsequent analyses (Figure 2). Input data to a study are selected from repositories using criteria defined in *Selector* objects passed to the Study object at initialisation and matched against entries in the class' data specification table.

Analysis design: Study classes

Study classes encapsulate a study dataset (i.e. data collected across multiple subjects using the same acquisition protocol) with the suite of analytical methods that can be applied to the dataset. The hierarchy of a study dataset is assumed to have two levels, *subjects* and *sessions*, with each session for each subject corresponding to a specific *visit*, e.g. timepoint in longitudinal study. Derivatives can be created at any point in this hierarchy: per-session, per-subject, per-visit and per-study. Iteration over subjects and visits is handled implicitly by the framework. All Study classes must inherit from the *Study* base class and be created by the *StudyMetaClass* metaclass or subclasses thereof.

$Data\ and\ parameter\ specification\ tables$

At the heart of each Study class is the data specification table, which specifies both the input and output data of the analysis, and all the stages in between. There is a one-to-one relationship between entries in the data specification table and data that is stored in the repository or will be stored if and when it is derived. Which intermediate derivatives to include in the data specification table, and therefore store in the repository, is left to the discretion of the researcher designing the analysis. However, as a general rule, derivatives that require manual QC or are likely to be reused between different branches of analysis should be included in the table.

Data specified in the data specification table can be of either a *fileset* or *field* type. Filesets represent single files, or sets of related files typically contained within a directory (e.g. a multi-volume DICOM dataset). Fields represent integer, floating point or character string values. By default, a Field represents a single value but if the *array* flag is set a Field represents a list of values. Each Fileset references a *FileFormat* object, which specifies the formats of the files in the set. File formats are explicitly registered by the researcher at design time using the *FileFormat.register(format)* class method to avoid conflicts where the same extension is used for different formats in different contexts.

Fileset and field specifications are passed to the data specification of a Study class via the add_data_specs class attribute as a list of named Fileset-Spec and FieldSpec objects (Figure 3). Specifications for acquired data (i.e. input data to the study) are distinguished from derived data by using the AcquiredFilesetSpec and AcquiredFieldSpec subtypes. However, the distinction is fluid, with derived specifications able to be overridden by acquired specifications in subclasses or MultiStudy classes, and vice-versa, or passed inputs when the class is instantiated.

All data specifications have a *frequency* attribute which specifies where the data sits in the hierarchy of the dataset and can take the values 'per_session', 'per_subject, 'per_visit' or 'per_study'. In addition, derived specifications are passed the name of a method in the class that constructs the pipeline to derive them. Therefore, while a pipeline can have multiple outputs, each derivative is derived by only one pipeline.

For fileset specifications that correspond to a known type in the BIDS standard, a *BidsSelector* or *BidsAssociatedSelector* can be provided to the *bids* keyword arg of the spec. BidsSelector specifies a primary scan in the BIDS standard using its type, modality and optional run number. BidsAssociatedSelector is used to select associated files, such as field maps and diffusion encoding matrices. When passing a BidsSelector to a FilesetSpec the *run* keyword argument of the selector is typically not set, so it can be read from the *bids_run* attribute of the Study. This allows the Study class to be applicable to any BIDS run.

Similar to data specifications, parameter specifications are included in the Study class by providing a list of ParameterSpec objects to the add_param_specs class attribute. ParameterSpec objects are initialised with a name and default value. Special parameters that specify a qualitative change in the analysis, for example using ANTs registration (Avants et al., 2011) instead of FSL registration (Smith et al., 2004), are specified by the SwitchSpec subtype. SwitchSpecs take a name, default value and a list of accepted values at initialisation.

```
from arcana import (
     Study, StudyMetaClass, AcquiredFilesetSpec, FilesetSpec,
      Acquired Field Spec\;,\;\; Field Spec\;,\;\; Parameter Spec\;,\;\; Switch Spec\;,\;\;
      FilesetSelector, XnatRepository)
from banana.file_format import (
      {\tt nifti\_gz\_format} \;,\;\; {\tt dicom\_format} \;,\;\; {\tt nifti\_format} \;,\;\; {\tt analyze\_format} \;,
      text_format , text_mat_format)
STD\_IMAGE\_FORMATS = (dicom\_format \,, \, nifti\_format \,, \, nifti\_gz\_format \,, \,
                             analyze_format)
# Select a Fileset collection to use as a default for template1
template_repo = XnatRepository(
     server='http://central.xnat.org', project_id='TEMPLATES'
     cache=os.path.expanduser(os.path.join('~', 'xnat-cache')))
template_selector = FilesetSelector(
     name='template1_default', format=nifti_gz_format,
      pattern='MNI152_T1', frequency='per_study')
template_collectn = template_selector.match(template_repo.tree())
class ExampleStudy(Study, metaclass=StudyMetaClass):
      add_data_specs = [
           # Acquired file sets
           AcquiredFilesetSpec('acquired_file1', text_format),
AcquiredFilesetSpec('acquired_file2', STD_IMAGE_FORMATS),
           # Acquired fields
           AcquiredFieldSpec('acquired_field1', int, array=True, frequency='per_subject'),
           AcquiredFieldSpec('acquired_field2', float, optional=True),
           # "Acquired" file set with default value. Useful for
           # standard templates
           AcquiredFilesetSpec('template1', STD_IMAGE_FORMATS,
                                      frequency='per_study',
                                      default=template_collectn),
           # Derived file sets
           FilesetSpec('derived_file1', text_format, 'pipeline1'),
FilesetSpec('derived_file2', nifti_gz_format, 'pipeline1'),
FilesetSpec('derived_file3', text_mat_format, 'pipeline2'),
FilesetSpec('derived_file4', dicom_format, 'pipeline3'),
FilesetSpec('derived_file5', nifti_gz_format, 'pipeline3',
                           frequency='per_subject'),
           FilesetSpec('derived_file6', analyze_format, 'pipeline2',
                           frequency='per_visit'),
           # Derived fields
           FieldSpec('derived_field1', float, 'pipeline2'),
FieldSpec('derived_field2', int, 'pipeline4',
                         frequency='per_study')]
     add_param_specs = [
           # Standard parameters
          ParameterSpec('parameter1', 10),
ParameterSpec('parameter2', 25.8),
# "Switch" parameters that specify a qualitative change
           # in the analysis
           SwitchSpec('node1_option', False), # Boolean switch
SwitchSpec('pipeline2_tool', 'toolA', ('toolA', 'toolB'))]
```

Fig. 3 Example data and parameter specifications. The data specification specifies two "acquired" file sets, 'one' and 'ten' and ten derived file sets that can be derived from them, at least indirectly. Each derived data spec, specifies the name of the pipeline constructor that creates the pipeline that derives them. Parameter specifications specify a name and default value for free parameters of the Study class

Workflow design: pipeline constructor methods

Workflows are implemented in Arcana as a series of pipelines, which each perform a unit of the analysis (e.g. registration, brain extraction, quantitative susceptibility mapping). Pipelines are represented by *Pipeline* objects, which are thin wrappers around Nipype workflows to handle input and output connections, and namespace management. Each pipeline consists of a (typically small) graph of Nipype nodes, with each node wrapping a stand alone tool (e.g. FSL's FLIRT) or analysis package function (e.g. SPM's *coreg* tool). Pipelines source their inputs from, and sink their outputs to, entries in the data specification table, thereby linking the table together.

Pipelines are constructed dynamically by "pipeline constructor" methods of the study. Pipeline constructor methods are referenced by name in the pipeline_name argument of the FilesetSpec and FieldSpec objects of data specifications they derive. Pipeline constructor methods should only receive wild-card keyword arguments (e.g. my_pipeline(self, **name_maps)), and these arguments should be passed as a dictionary directly to the name_maps argument of the pipeline initialisation to enable inputs and outputs of the pipeline to be rerouted to alternative data specifications in modified constructor methods, typically in subclasses and multi-studies (see Extension and specialisation by class inheritance and Implementing multi-modal studies).

The syntax for pipeline construction is inspired by a proposed form for Nipype v2.0 (github.com/nipy/nipype/issues/2539). Within a pipeline constructor method, a Pipeline object is constructed by the *Study.pipeline* method (Figure 4). At initialisation, each pipeline is given a name, which must be unique amongst the pipelines constructed by the Study. The methods implemented by the pipeline can be characterised by providing a the list of scientific references and a text description to the *references* and *desc* keyword arguments, respectively.

The add(name, interface) method is used to add a node to a pipeline and takes a unique name for the node (within the pipeline) and a Nipype Interface object, and returns a reference to the newly added node. For clarity, is recommended to put all static inputs (i.e. parameters) of the interface as keyword arguments of the interface constructor (Figure 4). However, if an input conditionally depends on a parameter of the study it can be set via the inputs attribute of the node (e.g. my_node.inputs.my_param = 1.0).

Interface traits are connected to inputs and outputs of the pipeline by providing *inputs* and *outputs* keyword arguments to the *add* method. Both arguments take a dictionary, where the keys of the dictionary are the names of traits of the interface. The values of the dictionary are 2-tuples each consisting of the name of a data specification and the format the pipeline either expects or produces the data in (i.e. a FileFormat for Fileset specifications or Python datatype for Field specifications). If the expected format does not match that of the corresponding data specification or study input then a conversion node is implicitly connected to the pipeline. Inputs and outputs that are conditional

```
def pipeline2(self, **name_maps):
    pipeline = self.pipeline(
    name='pipeline2',
         name_maps=name_maps
         desc="Description of the pipeline",
         references = [methods_paper_cite])
    node1 = pipeline.add(
         'node1'.
         Interface1 (
             param1=3.5,
             param2=self.parameter('parameter1')),
         inputs={
             'in_file1': ('acquired_file1', text_format),
'in_file2': ('acquired_file2', analyze_format),
'in_field': ('acquired_field1', int)},
              'out_field': ('derived_field1', int)}
         wall_time=25, requirements=[software_req1])
    if self.branch('node1_option'):
         node1.inputs.an_option = 'set-extra-option'
    if self.branch('pipeline2_tool', 'toolA'):
         pipeline.add(
              'node2',
             Interface2 (
                  param1=self.parameter('parameter2')),
                  'template': ('template1', nifti_gz_format)},
             connect={
                  'in_file': (node1, 'out_file')},
             outputs={
                  'out_file': ('derived_file3',
                                text_mat_format)},
             wall_time=10, requirements=[software_req2])
         self.connect_output('derived_file6', node1, 'out',
                                nifti_format)
    elif self.branch('pipeline2_tool', 'toolB'):
         pipeline.add(
              'node2',
             Interface3(),
             inputs={
                  'template': ('template1', nifti_gz_format)},
             connect={
                  'in_file': (node1, 'out_file')},
             outputs={
                  'out_file': ('derived_file3'
                                 text_mat_format)},
             wall_time=30, requirements=[matlab_req,
                                             toolbox1_req])
    else:
         self.unhandled_branch('pipeline2_tool')
    return pipeline
```

Fig. 4 Example pipeline constructor method. Pipelines are created using the *pipeline* method of the Study class. Pipeline objects are thin wrappers around Nipype Workflow objects to in order manage the namespaces of the workflow's inputs, outputs and nodes. Every pipeline constructor method should allow wildcard keyword arguments, which are passed to the *name_maps* argument of the pipeline initialisation. This allows pipeline constructors in sub and multi classes to map the inputs and outputs of the pipeline onto different data specifications.

on parameters or inputs provided to the study can be set outside of the *add* method using the *connect_input* or *connect_output* methods, respectively.

Connections between nodes are specified by a dictionary passed to the *connect* keyword argument of the *add* method that creates the receiving node. The dictionary keys are the names of the receiving traits and the values are 2-tuples consisting of the sending node and the name of the sending trait. Alternatively, conditional connections can be specified via the *connect* method of the pipeline, which simply calls the method of the same name in the underlying Nipype workflow.

Any external software packages required by a node should be referenced in the *requirements* keyword argument as a list of *Requirement* objects when the node added to the pipeline. Similarly, the expected memory requirements in MB and wall time for the node execution should be provided to the keyword arguments *memory*, and *wall_time*.

Iteration over subjects and visits is handled implicitly by Arcana and depends on the frequency of the pipeline's inputs and outputs. To create a summary derivative (i.e. frequency != 'per_session') from more frequent data, Study.SUBJECT_ID or Study.VISIT_ID should be passed to the joinsource keyword of the add method to join over subjects or visits, respectively. In this case, a JoinNode will be created instead of standard Node, which should be passed the additional keyword argument joinfield to specify the list of input traits to convert into lists to receive the joined input. Similarly, if the name of an input trait is provided (or list thereof) to the the iterfield keyword argument, then a MapNode will be created and the interface will be applied to all items of the list connected to that input (Gorgolewski et al., 2011). Additionally, the values of the subject and visit IDs are directly accessible as input fields of the pipeline named Study.SUBJECT_ID and Study.VISIT_ID, respectively.

Study parameters can be accessed during pipeline construction with the Study.parameter(name) method. If conditional logic is included in the workflow construction that alters the pipeline inputs, outputs or parameters then it should be controlled by a switch instead of a parameter. The analysis branch designated by a switch value should be tested with Study.branch(name) in the case of boolean switches and $Study.branch(name, ref_value)$ in the case of string switches.

Extension and specialisation by class inheritance

Because Arcana analyses are implemented as Python classes, class inheritance can be used to specialise existing analyses.

Instead of being set directly, the data and parameter specifications are set by the metaclass of the Study (i.e. StudyMetaClass), in order to combine them with corresponding specifications in the class' bases. The combined data and parameter specifications are constructed by visiting the class' bases in reverse method resolution order (MRO) and adding specifications from their

add_data_specs, add_param_specs attributes, overriding previously added specifications with matching names. Note that in this scheme, specifications can only be appended or overridden but not removed by Study subclasses so as not to break workflows inherited from base classes.

Pipeline constructor methods can be overridden in subclasses like any Python method. Often the overriding method will call the superclass method to construct a pipeline, apply modifications and return the modified pipeline. In this scenario, references to the data specification in the superclass method can be mapped onto different entries in the table of the subclass by providing the <code>input_map</code> and <code>output_map</code> keyword arguments when calling the superclass method. Additionally the name of the pipeline can be altered by providing the <code>name</code> argument, which is useful when creating multiple constructor methods from the one base method. In these scenarios it is important to also pass the the wildcard keyword arguments of the overriding method to the <code>name_maps</code> keyword argument of the superclass method, to allow the overriding method to be overridden in turn.

Implementing multi-modal studies

While basic Study classes are typically associated with single image modality or contrast, the analysis suites implemented by them can be integrated into multi-modal analysis by aggregating multiple Study classes (sub-studies) in a *MultiStudy* class. Analysis suites are integrated by joining the data specification tables of the sub-studies of a MultiStudy class. Entries in specification tables of sub-studies are joined by mapping them to a common entry in the specification table of the MultiStudy (Figure 5). This enables derivatives from one sub-study (e.g. brain extracted T1-weighted anatomical) to be referenced by workflows of other sub-studies (e.g. anatomically constrained dMRI tractography).

All MultiStudy classes must inherit from the MultiStudy base class and be created by the MultiStudyMetaClass metaclass or subclasses thereof. As in the case of subclassing the standard Study class, additional data and parameter specifications can be added to the class via add_data_specs and add_param_specs respectively for additional analysis not included in the sub-studies.

Sub-studies are aggregated in the *sub-study specification table* of a Multi-Study class via a list of *SubStudySpec* objects in the *add_sub_study_specs* class attribute, in the manner of data and parameter specifications. A SubStudySpec consists of a name, a Study class, and a *name map* dictionary. The name map dictionary maps data and parameter specification names from the namespace of the MultiStudy class to the namespace of the Study class.

Entries in the specification tables of sub-study classes that are not referenced in the sub study's name map are implicitly mapped to the Multi-Study namespace in the construction of the MultiStudy class by the metaclass using the name of the sub-study as a prefix. If the spec is a derivative then the pipeline constructor used to derive it is also mapped into the Multi-

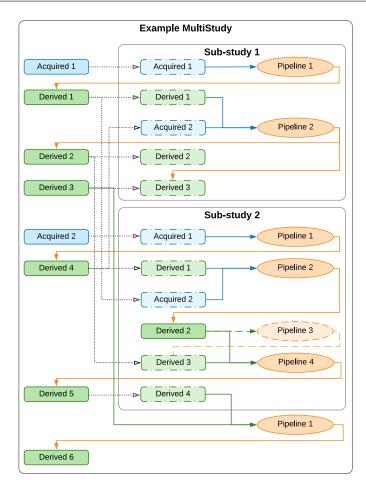


Fig. 5 Example MultiStudy. Blue boxes represent acquired (input) data (filesets or fields) and green derivatives. Orange ovals are pipelines. Blue and green arrows: acquired and derived inputs to pipelines, respectively. Orange arrows: outputs of pipelines. Dashed boxes represent data specifications in a sub-study that are present in the global namespace and mapped into the sub-study space, and dotted arrows the mappings. Sub-studies are linked by mapping the same data spec in the global space onto data specifications the multiple sub-study namespaces (e.g. Derived 1, 2 and 3). There are no restrictions between mapping acquired and derivative specifications: both acquired and derivative specifications can be mapped onto acquired data or derivative specifications in sub-studies. If a spec in the global spaced is mapped onto a derivative spec in the sub-study space, then the pipeline that generates that derivative in the sub-study will not run unless it generates other required derivatives (e.g. Pipeline 3 in Sub-Study 2)

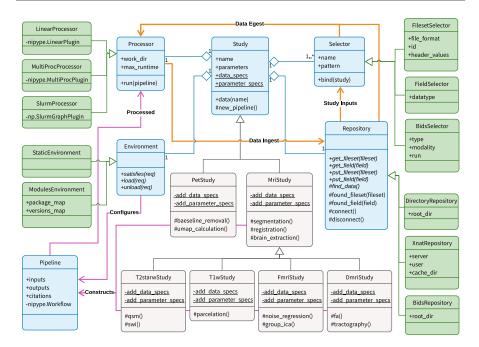


Fig. 6 Detailed Unified Modelling Language (UML) diagram of information flow in the Arcana framework. Boxes: Python classes (blue=core, green=interchangeable modules, grey=example specialisations). Arrows: orange=data, magenta=workflow description, diamond=aggregated-in, triangle=subclass-of. Calling data(name) on a Study subclass constructs the requisite pipelines (as specified in data_specs) to produce the requested data, and sends them to the Processor to be processed. Data is selected by Selector objects, pulled to the compute environment to be processed, and then the derivatives are pushed back to the repository. Repositories can be of plain directories, or BIDS or XNAT repositories

Study namespace. For example, derived1 in sub_study2 would be mapped to sub_study2_derived1 along with the method sub_study2_pipeline1.

Analysis application: Study instances

To apply the analysis blueprint specified in a Study class to a specific dataset, an instance of the Study class is created with details of where the data is stored (*Repository* module) the computing resources available to process it (*Processor* module) and the software installed in the environment it will be processed in (*Environment* module). A Study object controls the construction and execution of analysis workflows, and the flow of data to and from the repository (Figure 6).

Each Study instance is assigned a name, which is used to differentiate its results from alternative analyses on the same dataset (e.g. with different parameterisations). Parameters are set on initialisation of the Study object along with the range of subject and visit IDs to be included in the analysis (if they are not provided then all IDs found in the repository are included). The remaining arguments passed to the Study object initialisation are the Repository, Processor and Environment modules to use and a list of *Selector* objects to select input data from the repository and match it with the Study's data specification table. (Figure 7).

Repository modules

In Arcana, repository access is encapsulated within modular *Repository* objects to enable switching between different repositories and repository types at analysis "application time" (Figure 2). There are currently three supported repository types, XNAT (Marcus et al., 2007), BIDS (Gorgolewski et al., 2016) and a simple "directory" format, which are encapsulated by *XnatRepository*, *BidsRepository*, and *DirectoryRepository* classes respectively.

In its most basic form, a directory repository is just a file system directory with separate subdirectories for each subject in the study, with subject IDs taken from the subdirectory names. If the study has more than one visit then nested subdirectories for each visit are required in the subject subdirectories. Note that the directory repository is similar to the BIDS format, however, there are no naming conventions in the directory repository, which enables prototyping and testing of analyses on loosely structured data.

Derivatives are stored by their specification name in a study-specific namespaces to avoid clashes with separate analyses. In directory repositories this namespace is a subdirectory named after the study nested within each visit subdirectory if present or subject subdirectory otherwise. In BIDS repositories, the namespace is a subdirectory of the *derivatives* directory, again named after the study. In XNAT repositories, derivatives for each session are stored in separate *MrSession* objects alongside the primary session underneath its *Subject*, and are named *primary-session-name>_<study-name>* (Table 1).

Derived filesets are stored with the format specified in the study's data specification. In plain-directory and BIDS repositories, fields are stored in a single JSON file named '_fields__.json' in each derived session, and on XNAT they are stored in custom fields of the derived session. Provenance is stored in a '_prov__' sub-directory (dataset on XNAT) of the derivatives directory (MrSession on XNAT) in separate JSON files for each pipeline named after the pipeline (Table 1).

Summary data (i.e. with per_subject, per_visit, and per_study frequencies) are stored in specially named subjects and visits (e.g. 'all'), the names for which are specified when the repository is initialised. For example, given in plain-directory repository using all as the summary name for both subjects and visits, per_subject data for 'subj1' would be stored at root/subj1/all, per_visit data for 'visit1' in root/all/visit1, and per_study data in root/all/all (Table 1).

Each study can only have one repository in which derivatives are stored. However, a study can draw data from multiple auxiliary repositories, which are specified in the inputs passed to the study. When using multiple input

```
#!/usr/bin/env python3
import os.path as op
from arcana import (
    FilesetSelector\;,\;FieldSelector\;,\;XnatRepository\;,\\SlurmProcessor\;,\;ModulesEnvironment)
from .example_study import ExampleStudy
from banana.file_format import dicom_format
# Create study object that accesses an XNAT repository
# and submits jobs to a SLURM scheduler
study = ExampleStudy(
    # Give a name to this analysis to
    # differentiate it from other analyses
    # performed on the same data
    name='example',
    # Set up connection to XNAT repository
    repository=XnatRepository(
         project_id='SAMPLE_PROJECT',
         server='https://central.xnat.org',
cache_dir=op.expanduser('~/xnat-cache')),
    # Specify the use the SLURM scheduler to submit
    # nodes as jobs using Nipype's SlurmGraphPlugin
    processor=SlurmProcessor(
         work_dir=op.expanduser('~/work')),
    # Specify the use of environment modules to
    # satisfy software requirements. Non-standard
    # package names explicitly mapped to approp. reqs.
    environment=ModulesEnvironment (
         packages_map={
              'Package1_Parallel': package1_req }),
    # Link files and fields in the repository
    # to entries in the data specification
    inputs={
         'acquired1_file': FilesetSelector(
          '.*mprage.*', dicom_format, is_regex=True),
'acquired_file2': FilesetSelector(
         'SWI_Images', dicom_format),
'acquired_field1': FieldSelector(
              'YOB', int, frequency='per_subject'),
         'acquired_field2': FieldSelector(
               'weight', float)},
    # Specify parameters specific to this
    # analysis
    parameters = \{\, {}^{\backprime}parameter1 \,\, {}^{\backprime} \colon \  \, 55.0 \,\, ,
                    pipeline_tool': 'toolB'})
# Generate whole brain tracks and return path to
# a cached dataset
derived5 = study.data('whole_brain_tracks')
print("The \"derived5\" fileset for the second visit "
       "of subject 'PILOT1' were produced at:\n{}"
       .format(derived5.path(subject_id='PILOT1',
                                 visit_id=1))
```

Fig. 7 Example application of Study class to a dataset stored in an XNAT repository. Once the Study object has been initialised potential derivatives of the Study can be requested, and will be generated and stored in the repository if already present.

Table 1 Storage locations of derived data for each repository type. Derivatives are stored in separate namespaces for each Study instance to enable multiple analyses on the same datasets with different parameterisations. Where '...' is the location of the directory or MrSession that holds the derivatives, subj = subject ID, vis = visit ID, study = name of the Study instance, and pl = name of pipeline.

Datatype	Plain-directory	BIDS	XNAT
Derivatives	/subj/vis/study	$\label{lem:condition} $$ \derivatives/study/subj/vis $$/fieldsjson $$/prov/pl.json $$$	/subj/vis_study
Fields	/fieldsjson		MrSession XML
Provenance	/prov/pl.json		/_prov/pl.json

Table 2 Abstract methods in the base Repository class that need to be implemented by platform-specific sub-classes.

Method name	Function
find_data() get_fileset(Fileset) get_field(Field) put_fileset(Fileset) put_field(Field) connect() disconnect()	Queries repository to find all existing filesets and fields Caches fileset (if necessary) and returns the path to file(s). Retrieves and returns the value of the field from the repository Inserts fileset into the repository and updates cache. Inserts the value of the Field into the repository Opens a connection to the repository (optional). Closes connection to repository (optional)

repositories, subject and visit IDs will often need to be mapped from their values in the auxiliary repositories to the "ID space" of the study, which can be done by passing either by a dictionary or callable object (e.g. function) to the <code>subject_id_map</code> or <code>visit_id_map</code> keyword arguments during initialisation of a repository.

New repository modules for additional repository types can be implemented by extending the Repository abstract base class and implementing five abstract methods, find_data, get_fileset, get_field, put_fileset and put_field (Table 2).

$Study\ inputs$

While derivatives generated by a Study object are named in accordance with the data specification of the Study class, arbitrary naming conventions can be used for input datasets and fields to allow for heterogeneity in acquisition procedures. A selection stage is therefore necessary to match input data to appropriate entries in the data specification table of the Study class. The criteria for this selection is passed to the Study object at instantiation and is required to match exactly one fileset or field in every session included in the study.

Selection criteria are specified by list of FilesetSelector and FieldSelector objects provided to the inputs argument of the Study initialisation method. Matching is typically performed on file names (dataset labels for XNAT repositories), and field names. If the names are inconsistent across the study then regular expressions can be used instead of exact matches by passing is_regex=True

when initialising the Selector. Additional criteria can be used to distinguish cases where multiple filesets in the session match the pattern provided, such as DICOM header values, or the order and ID of the dataset.

Inputs can be drawn from auxiliary repositories by providing alternative Repository instances to the *repository* keyword of the Selector. Care should be taken to ensure that the subject and visit ID schemes will map correctly to that of the primary repository (see *Repository modules*). If the primary repository is empty (i.e. all inputs come from auxiliary repositories) then explicit *subject_ids*, *visit_ids* need to be provided and the *fill_tree* flag set when initialising the Study.

If the data to select has been derived from an alternative Arcana Study instance, then the name of the alternative study can be passed to the *from_study* keyword argument provided to the selector. There are no restrictions on selecting any data, derived or otherwise, to match acquired or derived specifications. For example, it is possible to deliberately skip analysis steps by selecting an output of an early step in a previous run as a match for a later derived spec although this is not recommended as standard practice.

Specific files and fields that are not stored within a repository can be passed as inputs in *FilesetCollection* and *FieldCollection* objects. Collection objects reference an entry in the data specification table and contain a single Fileset or Field for every session (or subject/visit/study depending on the frequency of the corresponding data spec). Collection objects can be used to pass reference atlases and templates as inputs to analyses. They can also be set as the "default input" for a data specification via the *default* keyword argument. However, for the sake of portability, default inputs should be restricted to data in publically accessible repositories or those included in standard software packages (e.g. FSL).

When using BIDS repositories, the selection stage is typically already included in the data specification (see *Data and parameter specifications*) so inputs do not need to be provided to the initialisation of the Study. However, BidsSelector and BidsAssociatedSelector objects can be provided to override the default selections is required.

Processor modules

Processor modules control how pipelines generated by a Study are executed. There currently three Processor modules implemented in Arcana: LinearProcessor, MultiProcProcessor, SlurmProcessor, which wrap the correspondingly named Nipype execution plugins. The main task performed by the processor, as separate from the Nipype execution plugin it wraps, is to determine which pipelines need to be run and link them into a single workflow. Since this logic is implemented in the Processor abstract base class, wrapping additional Nipype plugins as required is trivial.

A Processor is used internally by a Study instance to execute pipelines to derive derivatives requested from the data specification by the data(name[,name,...]) method (Figure 2). The first step in this procedure is to query the

Table 3 Effect of the *reprocess* flag value on the behaviour of the study when existing derivatives are found that were derived with mismatching inputs and parameters.

Reprocess value	Behaviour on provenance mismatch
False (default)	Raises an exception
True	Mismatching derivatives will be reprocessed
'ignore'	Mismatches are ignored
'ignore_versions'	Ignore versions, otherwise raise exception
'ignore_versions_true'	Ignore versions, otherwise the derivative will be reprocessed

repository tree for all data and provenance associated with the study. Sessions for which the requested outputs of the pipeline are already present in the repository, and the stored provenance matches the current parameters of the study, excluded from the list to process. For the remaining sessions to process, inputs of the pipeline that are derivatives themselves are added to the stack of requested derivatives. This procedure is repeated recursively until there are no sessions to process or all inputs to the pipeline are study inputs at a given depth.

When a pipeline is processed it is connected to source and sink nodes, which get and put the pipeline inputs and outputs from and to a repository, respectively. Separate source and sink nodes are used for each data frequency (i.e. per-session, per-subject, per-visit, per-study). If implicit file format conversion is required (i.e. the input or output format differs from the data specification) then additional format converter nodes are inserted after the source nodes or before the sink nodes. Iterator nodes that iterate over the required subjects and visits are connected to the sources, and "deiterator" nodes that join over subjects and visits are connected to the sink nodes. Final nodes of upstream pipelines are connected to downstream iterator nodes in order to create a single workflow, which is then executed using the Nipype execution plugin.

Provenance is stored for each pipeline run alongside the generated derivatives and consists of: parameter values used by the pipeline, software versions used by the pipeline, a graph representation of the underlying Nipype workflow, checksums of inputs, version of Arcana used, version of Nipype used, subject and visit IDs used by study.

For subsequent analyses, changes w.r.t. Any of the stored values, with the exception of subject and visit IDs, will be flagged as a mismatch. Subject or visit IDs only flag a mismatch if any of the inputs, or inputs of upstream pipelines, are per-visit or per-subject, respectively, or per-study. How provenance mismatches are handled by the study is determined by the *reprocess* flag, which is passed to the Study on initialisation (Table 3).

Additionally, a time limit can be passed to the *reprocess_time_limit*, and if the estimated time to reprocess all the required sessions is less than the time limit they will be reprocessed and otherwise an exception will be raised.

Environment modules

The software packages installed on the system that are required by a study's workflows (e.g. FSL, SPM) are detected and managed by the Environment object passed to the study at initialisation. There are currently two types of Environment class implemented in Arcana: *StaticEnvironment* and *Module-sEnvironment*.

Modules Environment objects can be used if environment modules (Furlani, 1991) are installed on the system (typical in many HPC systems). In this case, environment modules are loaded before a workflow node is run, based on the requirements specified for the node during construction of the pipeline (see Pipeline constructors), and then unloaded afterwards. Mappings from non-standard module names installed on the system to those expected by the Study can be can be passed as a dictionary to the packages_map keyword argument at initialisation. Likewise non-standard versions for packages can be mapped onto requirements using the versions_map keyword argument.

StaticEnvironment objects don't actively manage the environment, and instead only check the current environment for appropriate software versions before running requested workflows.

Acquisition of test dataset

A healthy volunteer was scanned using a 3T Siemens Skyra with a 32-channel head and neck coil to demonstrate the application of analyses implemented in Arcana. The protocol was a T1-weighted MPRAGE (1mm contiguous, matrix size 256x240x192, FOV 256x240x192, TE = 2.13ms, TR, = 2300ms, TI = 900ms, bandwidth = 230Hz/pixel), GRE (1.8 mm contiguous, matrix size 256x232x72, FOV 230x208x130, TE = 20ms, TR, = 30ms, bandwidth = 120Hz/pixel), and diffusion MRI (1.2 mm contiguous, matrix size 110x100x60, FOV 256x240x192, TE = 95ms, TR, = 8200ms, 33 diffusion directions with b = 1500 mm2/s and 3 b=0, bandwidth = 781Hz/pixel).

Results

The Arcana framework is distributed as a publicly available software package via GitHub (github.com/MonashBI/arcana) and the Python Package Index (PyPI) (pypi.org/project/arcana/). Study classes for T1, T2* and diffusion weighted MRI data have been implemented as part of the Biomedical imAgiNg ANAlysis (Banana) package (github.com/MonashBI/banana; pypi.org./project/banana). All three classes, T1Study, T2starStudy and DmriStudy, inherit generic image analysis methods, such as registration and brain extraction, from the base class MriStudy.

The DmriStudy class implements the extraction of diffusion tensor metrics, FA and ADC, as well as whole-brain tractography using streamlines tracking from the MRtrix toolbox (Tournier et al., 2010, 2012) (Figure 8).

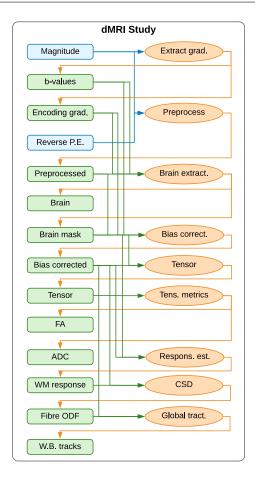


Fig. 8 Example diffusion MRI (dMRI) study, which can derive tensor metrics, fractional anisotropy (FA) and apparent diffusion coefficient (ADC) as well as streamlines fibre tracking. Blue boxes: acquired (input) data (filesets or fields). Green boxes: derivatives. Orange ovals: pipelines. Blue and green arrows: acquired and derived inputs to pipelines, respectively. Orange arrows: outputs of pipelines. The dMRI magnitude image is preprocessed for motion correction and EPI distortions masked and bias corrected. From the bias corrected image two branches of analysis can be performed using the same intermediate derivatives: FA and ADC and/or streamlines fibre tracking.

The T2starStudy class implements an algorithm to generate $composite\ vein\ images$ (Ward et al., 2018) and vein masks (Ward et al., 2017) from the combination of vein atlases derived from manual tracings with QSM and SWI contrasts derived from the T2*-weighted acquisition (Figure 9).

The T1Study, T2starStudy and DmriStudy classes are aggregated into a single MultiStudy class, *ArcanaPaper* (Supplementary material), which is specialised to produce the figures in the Results section of this manuscript. In order to warp the vein atlases to the subject space for comparison with the SWI and QSM images, nonlinear registration to Montreal Neurological In-

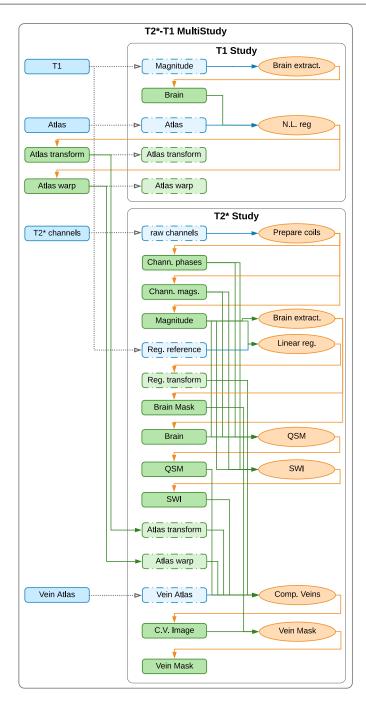


Fig. 9 Combined T2*-weighted/T1-weighted multi-study, which can derive vein masks by combining Quantitative Susceptibility Mapping (QSM) and Susceptible Weighted Imaging (SWI) contrasts with a manual atlas. Blue boxes: acquired (input) data (filesets or fields). Green boxes: derivatives. Orange ovals: pipelines. Blue and green arrows: acquired and derived inputs to pipelines, respectively. Orange arrows: outputs of pipelines. Dashed boxes represent data specifications in a sub-study that are present in the global namespace and mapped into the sub-study space, and dotted arrows the mappings. The acquired T1-weighted image is mapped to both the magnitude spec of the T1-weighted sub-study and the registration reference spec of the T2*-weighted sub-study. The nonlinear transformation from subject to atlas space are mapped from the T1-weighted sub-study and combined with the linear registration between T1-weighted and T2*-weighted images, QSM and SWI images mapped from the T2*-weighted sub-study to produce the composite-vein image. The brain mask from the T1-weighted sub-study is combined with the composite vein image to produce the vein mask.

stitute templates (Grabner et al., 2006) is performed in the T1Study. The transform and warp field from this registration are then mapped onto the 'coreg_to_atlas_mat' and 'coreg_to_atlas_warp' specifications in the T2starStudy, in order to improve the registration. These transforms were used to produce the composite vein image in conjuction with the linear transform from the brain-extracted T2*-weighted magnitude image to the brain extracted T1-weighted image, which required the 'brain' specification in the T1Study to be mapped to the 'coreg_ref_brain' specification in the T2starStudy.

The vein_fig, fa_adc_fig and tractography_fig methods implemented in the ArcanaPaper class were applied to the test dataset to produce Figure 10, 11, and 12, respectively. Figure 10 displays composite vein images and vein masks for the healthy volunteer along with the SWI and QSM intermediate derivatives. The derived vein images are comparable to those generated by the original implementation (Ward et al., 2018).

Figure 11 displays the FA and ADC maps derived from the diffusion MRI acquisition. The FA map shows high intensity in known white matter tracts and low intensities in known grey matter regions. The ADC map shows high intensity in cortical spinal fluid and low intensity through the rest of the brain. Figure 12 displays the global tractography derived from the dMRI acquisition. The streamlines follow well known white matter tracts such as the corticospinal, fasciculus and corpus callosum. Intermediate derivatives derived for the FA and ADC analysis, including the preprocessed and bias-corrected dMRI image and a whole brain mask, were reused in the generation of the streamlines.

Discussion

We present Arcana, a software framework to facilitate the development of comprehensive analysis suites for neuroimaging data that implement complete workflows from repository data to publication results. The encapsulation of repository data and workflow generation in Arcana enables researchers to create robust workflows while focussing on the core logic of their analysis. Arcana's modular pipeline and OO architecture promotes code reuse between different workflows by facilitating the sharing of common segments (e.g. registration, segmentation). The clear separation of analysis design from its application leads to portable workflows, which can be applied to datasets stored in a number of storage systems. In addition, the management of intermediate derivatives, provenance and software versioning, coupled with ability to submit jobs to HPC clusters, enables workflows implemented in Arcana to scale to large datasets. Arcana thereby enables researchers to quickly prototype analysis suites on local workstations that can be deployed on enterprise-scale infrastructure without modification.

Software frameworks (Yacoub and Ammar, 2004) have been successful in improving code quality and efficiency of development in a variety of contexts (Moore et al., 2008; White, 2012; Abadi et al., 2016). By factoring out common elements, only features that are specific to the given application need to be

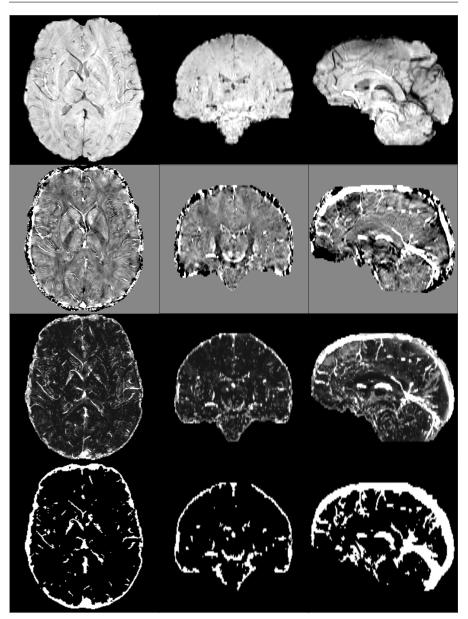


Fig. 10 Composite vein image (third row) constructed by combining susceptibility weighted imaging (SWI) (top row), quantitative susceptibility mapping (QSM) (second row) and a vein atlas from manual tracings. A vein mask was then generated (bottom row) from the composite vein image. left column: axial slices. centre column: coronal slices. right column: sagittal slices.

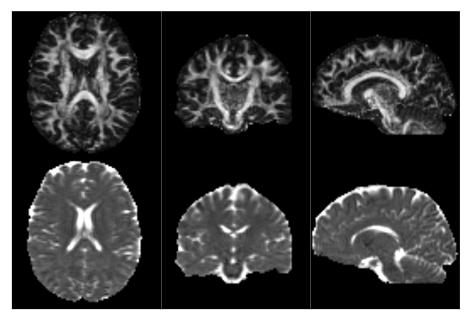


Fig. 11 Fractional Anisotropy (FA) (top row) and Apparent Diffusion Coefficient (ADC) (bottom rowI) derived from diffusion MRI data. Left column: axial midline slices. Middle column: coronal midline slices. Left column: sagittal midline slices.

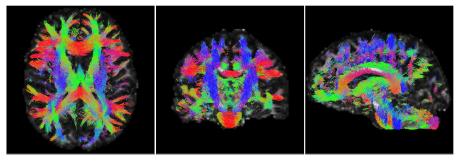


Fig. 12 Global tractography performed using the MRtrix toolbox. Probabilistic streamlines generated with the iFOD2 algorithm from fibre Orientation Distribution Function (fODF) estimated from diffusion MRI datasets using Constrained Spherical Deconvolution (CSD). Streamlines are colour-encoded by orientation: green=anterior-posterior, blue=inferior-superior, red=left-right. Left panel: axial midline slice. Middle panel: coronal midline slice. Left panel: sagittal midline slice.

implemented by the analysis designer, and the common elements become battle hardened through repeated use. Arcana handles many of the menial tasks involved with workflow implementation, such as data retrieval and storage, format conversions, and provenance, reducing the time and effort required to implement robust workflows from acquired data to publication results.

An oft-repeated mantra in the open-source software movement dubbed Linus' Law is that "given a large enough beta-tester and co-developer base, almost every problem will be characterized quickly and the fix obvious to someone" or more compactly, "given enough eyeballs, all bugs are shallow" (Raymond, 1999). Given the size of the neuroimaging research community, there are a large number of potential beta-testers and co-developers. However, it has been difficult for researchers to collaborate on the same code-base due to slight differences in acquisition protocols, storage conventions, researcher preferences, and study requirements.

The flexibility and portability of the Arcana framework increases the feasibility of community collaborations on workflow implementations. The improvement of code quality in larger community efforts, due to more eyeballs to detect and fix errors, has the potential to form a reinforcing cycle where more developers are attracted to the project. To these ends, the Banana code repository on GitHub (github.com/MonashBI/banana.git) is proposed as a code base for communal development of biomedical imaging workflows using Arcana.

A level of proficiency in Python OO design is required to design new analyses in Arcana, which may preclude inexperienced programmers. However, only a basic knowledge of Python is required to apply existing analyses to new datasets. Furthermore, a number of example Study classes have been implemented, which can guide the hand of analysis designers. Arcana imposes a consistent structure on workflows implemented within it, making the code easier to understand for developers who are familiar with the framework. In addition, class inheritance provides a manageable way to adapt and extend to existing analyses and highlights where modified analyses differ from standard procedures.

MR contrast-specific analyses are implemented in Banana via a chain of successively specialised Study sub-classes (e.g. MRI>EPI>dMRI) to enable generic processing steps (e.g. registration) to be shared between classes. While not necessary, it is recommended to create a subclass specific to the research study in question and aggregate all related analysis within it, since such classes can be applied to alternate datasets in order to reproduce the exact analysis. The *ArcanaPaper* class (Supplementary material), which contains methods to generate all figures in the Results section of this manuscript, is an example of this approach.

The abstraction of data and repositories in Arcana enables the same workflow implementation to be applied to datasets stored in BIDS format or XNAT repositories. A single code-base can therefore be containerized into BIDS apps or XNAT pipelines without adaptation, helping to form a bridge between the two communities of users and developers. Alternative data storage systems (Scott et al., 2011; Das et al., 2012; Book et al., 2013), can be integrated into Arcana by overriding a small number of methods from the Repository abstract base class. Repository modules could also be created for data portals such as DataLad (Halchenko et al., 2018) in order to take advantage of the range of platforms they support. Implementing analyses in Arcana therefore enables researchers and research groups to easily migrate their workflows between storage platforms, and not risk being locked in to a particular technology.

While Arcana was primarily developed for neuroimaging datasets, it is a general framework that could be applied to data from other fields. However, in other contexts, the subject and visit hierarchy may no longer make sense. In many cases it may be sufficient to map subjects and/or visits onto alternative concepts (e.g. for meteorological data *subjects* = *weather stations*, *visits* = *observation times*). But some cases may require a deeper data hierarchy (i.e. greater than two), which is not currently possible in Arcana.

Ensuring that the consistent versions of external tools are used throughout the analysis is important to avoid introducing biases due to algorithm updates. In systems with environment modules (Furlani, 1991) installed, Arcana can load and unload the required modules before and after each node is executed. When running Arcana within a container, environment modules and software versions can be installed inside the container giving exact control over the versions used. To these ends, a Docker container is available on Docker Hub, (hub.docker.com/r/monashbi/banana), which can be used as a base for biomedical imaging analysis containers. In future versions of Arcana, additional Environment modules could be implemented to run each pipeline node within its own container to take advantage of containers maintained by tool developers (e.g. hub.docker.com/r/vistalab/freesurfer/).

While the same tools and versions should be applied across an analysis to avoid bias, there are cases where it is desirable to rerun the same analysis with different tools substituted at various points in the workflow. In particular, when introducing new tools or upgrades to existing tools, it is important to show the effect on the final results in comparison with existing methods. Furthermore, it is typically not clear what variability between results produced by comparable tools is due to. Therefore, in the absence of *a priori* reason to favour a particular tool, perhaps the most rigorous approach is to rerun analyses with different combinations of available tools and only present results that are robust to the "analytic noise" (Maumet, 2018) they introduce. Switch parameters make it straightforward to rerun analyses in Arcana with substituted tools while controlling all other aspects of the workflow.

Arcana's management of intermediate derivatives and provenance guarantees that the same analysis is applied across the dataset without necessarily requiring a complete rerun of the analysis. This guarantee makes it feasible to process data as it is acquired over the course of long studies, and therefore help detect any problems that might arise with the acquisition protocol when they occur. In addition, by reusing shared intermediate derivatives between analyses, such as the preprocessed dMRI shared between tensor and fibre tracking workflows (Figure 11 and 12), processing time as well as time required for manual QC is minimised. Given analyses implemented in Arcana are also able to be processed on HPC clusters, they scale well to large studies.

Conclusion

By managing the complete flow of data from/to a repository with modular components, Arcana enables complex analyses of large-scale neuroimaging studies that are portable across a wide range of research sites. The extensibility of analyses implemented in Arcana, coupled with the flexibility afforded by programmatic constructuction of pipelines, facilitates the design of comprehensive analyses by larger communities. Larger communities of developers working on the same code-base should make it feasible to capture the arcana of neuroimaging analysis in templates that can be applied to a wide range of relevant datasets.

Acknowledgements The authors acknowledge the facilities and scientific and technical assistance of the National Imaging Facility, a National Collaborative Research Infrastructure Strategy (NCRIS) capability, at Monash Biomedical Imaging, Monash University. The "transparent repository" feature of Arcana was inspired by in-house software written by Parnesh Raniga while he was employed at Monash University prior to 2016.

References

- Abadi M, Barham P, Chen J, Chen Z, Davis A, Dean J, Devin M, Ghemawat S, Irving G, Isard M, et al. (2016) TensorFlow: A system for large-scale machine learning. Savannah, GA, USA, p 21
- Achterberg HC, Koek M, Niessen WJ (2016) Fastr: A Workflow Engine for Advanced Data Flows in Medical Image Analysis. Frontiers in ICT 3, DOI 10.3389/fict.2016.00015
- Amstutz P, Crusoe MR, Tijani N, Chapman B, Chilton J, Heuer M, Kartashov A, Leehr D, Mnager H, Nedeljkovich M, Scales M, Soiland-Reyes S, Stojanovic L (2016) Common Workflow Language, v1.0. DOI 10.6084/m9. figshare.3115156.v2
- Avants BB, Tustison NJ, Song G, Cook PA, Klein A, Gee JC (2011) A reproducible evaluation of ANTs similarity metric performance in brain image registration. NeuroImage 54(3):2033–2044, DOI 10.1016/j.neuroimage.2010. 09.025
- Book GA, Anderson BM, Stevens MC, Glahn DC, Assaf M, Pearlson GD (2013) Neuroinformatics Database (NiDB)—a modular, portable database for the storage, analysis, and sharing of neuroimaging data. Neuroinformatics 11(4):495–505, DOI 10.1007/s12021-013-9194-1
- Cox RW (1996) AFNI: software for analysis and visualization of functional magnetic resonance neuroimages. Computers and Biomedical Research, an International Journal 29(3):162–173
- Cusack R, Vicente-Grabovetsky A, Mitchell DJ, Wild CJ, Auer T, Linke AC, Peelle JE (2015) Automatic analysis (aa): efficient neuroimaging workflows and parallel processing using Matlab and XML. Frontiers in Neuroinformatics 8, DOI 10.3389/fninf.2014.00090

- Das S, Zijdenbos AP, Harlap J, Vins D, Evans AC (2012) LORIS: a web-based data management system for multi-center studies. Frontiers in Neuroinformatics 5(January):1–11, DOI 10.3389/fninf.2011.00037, iSBN: 1662-5196
- Esteban O, Markiewicz C, Blair RW, Moodie C, Isik AI, Erramuzpe Aliaga A, Kent J, Goncalves M, DuPre E, Snyder M, Oya H, Ghosh S, Wright J, Durnez J, Poldrack R, Gorgolewski KJ (2018) FMRIPrep: a robust preprocessing pipeline for functional MRI. bioRxiv DOI 10.1101/306951
- Friston K (2007) Statistical Parametric Mapping. Elsevier, DOI 10.1016/B978-0-12-372560-8.X5000-1
- Furlani JL (1991) Modules: Providing a flexible user environment. In: Proceedings of the fifth large installation systems administration conference (LISA V), pp 141–152
- Gorgolewski K, Burns CD, Madison C, Clark D, Halchenko YO, Waskom ML, Ghosh SS (2011) Nipype: a flexible, lightweight and extensible neuroimaging data processing framework in python. Frontiers in Neuroinformatics 5:13, DOI 10.3389/fninf.2011.00013
- Gorgolewski KJ, Auer T, Calhoun VD, Craddock RC, Das S, Duff EP, Flandin G, Ghosh SS, Glatard T, Halchenko YO, Handwerker DA, Hanke M, Keator D, Li X, Michael Z, Maumet C, Nichols BN, Nichols TE, Pellman J, Poline JB, Rokem A, Schaefer G, Sochat V, Triplett W, Turner JA, Varoquaux G, Poldrack RA (2016) The brain imaging data structure, a format for organizing and describing outputs of neuroimaging experiments. Scientific Data 3:160044, DOI 10.1038/sdata.2016.44
- Grabner G, Janke AL, Budge MM, Smith D, Pruessner J, Collins DL (2006) Symmetric Atlasing and Model Based Segmentation: An Application to the Hippocampus in Older Adults. In: Larsen R, Nielsen M, Sporring J (eds) Medical Image Computing and Computer-Assisted Intervention MICCAI 2006, Springer Berlin Heidelberg, Lecture Notes in Computer Science, pp 58–66
- Halchenko Y, Hanke M, Poldrack B, Meyer K, Debanjum, Alteva G, jason gors, MacFarlane D, Husler CO, Olson T, Waite A, de la Vega A, Keshavan A, bhanuprasad14, yetanothertestuser, yarikoptic private, Lau VC, tstoeter, Hardcastle N, di Oleggio Castello MV, Skytn K, Poelen J, Christian H, Ma F (2018) datalad/datalad 0.10.3.1. DOI 10.5281/zenodo.1418485
- Kennedy DN (2018) Neuroimaging Neuroinformatics: Sample Size and Other Evolutionary Topics. Neuroinformatics 16(2):149–150, DOI 10.1007/s12021-018-9379-8
- Marcus DS, Olsen TR, Ramaratnam M, Buckner RL (2007) The extensible neuroimaging archive toolkit. Neuroinformatics 5(1):11–33, DOI 10.1385/NI:5:1:11
- Maumet C (2018) Tools and standards to make neuroimaging derived data reusable. In: Neuroinformatics 2018, Montreal, Canada, URL http://www.hal.inserm.fr/inserm-01886089
- Moore D, Budd R, Wright W (2008) Professional Python Frameworks: Web 2.0 Programming with Django and Turbogears. John Wiley & Sons Raymond ES (1999) The Cathedral and the Bazaar. O'Reilly Media, p 30

- Schreiber J, Hoffstaedter F, Deepu R, Orth B, Lippert T, Amunts K, Eickhoff S, Caspers S (2018) Using a Multi-Petaflop Supercomputer for Pushing Neuroimaging Analytics to the Next Level. In: Proceedings of Organisation for Human Brain Mapping 2018, Singapore, Singapore
- Scott A, Courtney W, Wood D, de la Garza R, Lane S, King M, Wang R, Roberts J, Turner JA, Calhoun VD (2011) COINS: An Innovative Informatics and Neuroimaging Tool Suite Built for Large Heterogeneous Datasets. Frontiers in Neuroinformatics 5:33, DOI 10.3389/fninf.2011.00033
- Smith SM, Jenkinson M, Woolrich MW, Beckmann CF, Behrens TEJ, Johansen-Berg H, Bannister PR, De Luca M, Drobnjak I, Flitney DE, Niazy RK, Saunders J, Vickers J, Zhang Y, De Stefano N, Brady JM, Matthews PM (2004) Advances in functional and structural MR image analysis and implementation as FSL. NeuroImage 23 Suppl 1:S208–219, DOI 10.1016/j.neuroimage.2004.07.051
- Sudlow C, Gallacher J, Allen N, Beral V, Burton P, Danesh J, Downey P, Elliott P, Green J, Landray M, Liu B, Matthews P, Ong G, Pell J, Silman A, Young A, Sprosen T, Peakman T, Collins R (2015) UK Biobank: An Open Access Resource for Identifying the Causes of a Wide Range of Complex Diseases of Middle and Old Age. PLoS Medicine 12(3), DOI 10.1371/journal. pmed.1001779
- Thompson PM, Stein JL, Medland SE, Hibar DP, Vasquez AA, Renteria ME, Toro R, Jahanshad N, Schumann G, Franke B, et al. (2014) The ENIGMA Consortium: large-scale collaborative analyses of neuroimaging and genetic data. Brain Imaging and Behavior 8(2):153–182, DOI 10.1007/s11682-013-9269-5
- Tournier JD, Calamante F, Connelly A (2010) Improved probabilistic streamlines tractography by 2nd order integration over fibre orientation distributions. In: Proceedings of the international society for magnetic resonance in medicine, vol 18, p 1670
- Tournier JD, Calamante F, Connelly A (2012) MRtrix: Diffusion tractography in crossing fiber regions. International Journal of Imaging Systems and Technology 22(1):53–66, DOI 10.1002/ima.22005
- Van Essen DC, Ugurbil K, Auerbach E, Barch D, Behrens TEJ, Bucholz R, Chang A, Chen L, Corbetta M, Curtiss SW, Della Penna S, Feinberg D, Glasser MF, Harel N, Heath AC, Larson-Prior L, Marcus D, Michalareas G, Moeller S, Oostenveld R, Petersen SE, Prior F, Schlaggar BL, Smith SM, Snyder AZ, Xu J, Yacoub E, WU-Minn HCP Consortium (2012) The Human Connectome Project: a data acquisition perspective. NeuroImage 62(4):2222–2231, DOI 10.1016/j.neuroimage.2012.02.018
- Ward PG, Ferris NJ, Raniga P, Ng AC, Barnes DG, Dowe DL, Egan GF (2017) Vein segmentation using shape-based Markov Random Fields. In: Biomedical Imaging (ISBI 2017), 2017 IEEE 14th International Symposium on, IEEE, pp 1133–1136
- Ward PGD, Ferris NJ, Raniga P, Dowe DL, Ng ACL, Barnes DG, Egan GF (2018) Combining images and anatomical knowledge to improve automated vein segmentation in MRI. NeuroImage 165:294–305, DOI 10.1016/

```
j.neuroimage.2017.10.049
White T (2012) Hadoop: The definitive guide. O'Reilly Media, Inc.
Yacoub SM, Ammar HH (2004) Pattern-oriented Analysis and Design: Composing Patterns to Design Software Systems. Addison-Wesley Professional, google-Books-ID: dbU4ggCbqd4C
```

Supplementary material

```
Listing 1 ArcanaPaper class definition
```

```
#!/usr/bin/env python3
from arcana import (
   MultiStudy, MultiStudyMetaClass, SubStudySpec, ParameterSpec)
from banana.study.mri import (
   DmriStudy, T1Study, T2starStudy, MriStudy)
from banana.plot import ImageDisplayMixin
import os.path as op
class ArcanaPaper (MultiStudy, ImageDisplayMixin,
                  metaclass=MultiStudyMetaClass):
   An Study class to perform the analyses performed in the paper
    introducing Arcana. Requires a repository containing
   T1-weighted, T2*-weighted and diffusion MRI data.
    add_sub_study_specs = [
       # Include two Study classes in the overall study
        SubStudySpec(
            't1'.
           T1Study)
        SubStudySpec (
            't2star'
            T2starStudy,
           't1_coreg_to_atlas_mat': 'coreg_to_atlas_mat',
                      't1_coreg_to_atlas_warp': 'coreg_to_atlas_warp'}),
        {\bf SubStudySpec}\,(
            'dmri'
            DmriStudy),
       # Since we are using the SWI image produced from the scanner console
       # we need to brain extract it in order to display it properly, so
       # we create a basic MriStudy to use its brain extraction before
       # mapping the extracted image to the T2* study.
        SubStudySpec (
            'swi'
            MriStudy)]
    add_param_specs = [
        ParameterSpec (
            'dmri_fig_slice_offset', (6, 0, 0),
            desc=("Offset the sagittal slices of dMRI figures so they "
```

```
"intersect a cerebral hemisphere instead of the "
              "midline between the hemispheres"))]
def vein_fig(self, save_path=None, **kwargs):
    Generates an image panel containing the SWI, QSM, vein atlas,
    and vein mask
    Parameters
    save_path : str | None
        The path to save the image to. If None the image will be
        displayed instead of saved
    # Derive (when necessary) and access data SWI, QSM, vein atlas,
    # and vein mask data in the repository.
    swis = self.data('swi_brain')
qsms = self.data('t2star_qsm')
    cv_image = self.data('t2star_composite_vein_image')
    vein_masks = self.data('t2star_vein_mask')
    # Loop through all sessions
    for swi, qsm, vein_atlas, vein_mask in zip(swis, qsms, cv_image,
                                                  vein_masks):
        # Set the saturation limits for the QSM image
        row_kwargs = [\{\} for _ in range(4)]
        row_kwargs[1]['vmax_percentile'] = 95
        row_kwargs[1]['vmin_percentile'] = 5
        # Display slices from the filesets in a panel using
        # method from the ImageDisplayMixin base class.
        self. \, display\_slice\_panel \, (
            (swi, qsm, vein_atlas, vein_mask),
        row_kwargs=row_kwargs, **kwargs)
# Display or save to file the generated image.
        self.show(save_path, swi.subject_id, swi.visit_id)
def fa_adc_fig(self, save_path=None, **kwargs):
    Generates an image panel containing the derived FA and ADC
    images for each session in the study (typically only one)
    Parameters
    save_path : str | None
        The path to save the image to. If None the image will be
        displayed instead of saved
    # Derive FA and ADC if necessary and return a handle to the
    # data collections in the repository
    fas, adcs = self.data(('dmri_fa', 'dmri_adc'))
    # Loop through all sessions
    for fa, adc in zip(fas, adcs):
        # Display slices from the filesets in a panel using
        # method from the ImageDisplayMixin base class.
        self.display_slice_panel(
            (fa, adc),
            row_kwargs=({ 'vmax': 1.0}, {}),
            offset=self.parameter('dmri_fig_slice_offset'),
             **kwargs)
```

```
# Display or save to file the generated image.
            self.show(save_path, fa.subject_id, fa.visit_id)
    def tractography_fig(self, save_path=None, **kwargs):
        Generates dMRI tractography streamlines seeded from without the
        white matter.
        Is a bit more complex as it uses MRtrix's mrview to display and
        screenshot to file the high volume of streamlines created. Then
        reload, crop and combine the different slice orientations into
        a single panel.
        Parameters
        save_path : str | None
            The path of the file to save the image at. If None the
            images are displayed instead
        size : Tuple(2)[int]
            The size of the combined figure to plot
        # Generate and loop through tcks and FA maps for each
        # session in repository
        for tck, fa in zip(*self.data(('dmri_global_tracks',
                                        'dmri_fa'))):
            # Display generated streamlines with mrview from the
            # MRtrix package
            self.display_tcks_with_mrview(
                tcks=(tck,), backgrounds=(fa,),
                offset=self.parameter('dmri_fig_slice_offset'),
                **kwargs)
            # Display or save to file the generated image.
            self.show(save_path, tck.subject_id, tck.visit_id)
Listing 2 Application of ArcanaPaper class to a dataset
    import os
    from argparse import ArgumentParser
    from arcana import (
        Directory Repository, Linear Processor, Static Environment,
        FilesetSelector)
    from banana.file_format import (
        parser = ArgumentParser(
        "A script to produce figures for the Arcana manuscript")
    parser.add_argument('data_dir'
                        help="The directory containing repository")
    parser.add_argument('--work_dir', help="The work directory",
                        default=op.join(op.expanduser(',~')),
                                         'arcana-paper-work'))
    parser.add_argument('--fig_dir',
                        help="The directory to store the figures",
default=op.join(op.expanduser('~')),
                                         'arcana-paper-fig'))
    parser.add_argument('--t1', default='.*mprage.*',
                        help="Pattern to match T1-weighted scan")
    parser.add_argument('--t2star_chann', default='.*channels.*',
```

```
help="Pattern to match T1-weighted scan")
parser.add_argument('--swi', default='.*(swi|SWI).*',
                    help="Pattern to match T1-weighted scan")
parser.add_argument('--dmri', default='.*diff.*',
                     help="Pattern to match T1-weighted scan")
parser.add_argument('--distort', default='.*distortion_correction.*',
                    help="Pattern to match T1-weighted scan")
args = parser.parse_args()
# Make figure directory
os.makedirs(args.fig_dir, exist_ok=True)
# Instantiate the ArcanaPaper class in order to apply it to a
# specific dataset
paper = ArcanaPaper(
    # Name for this analysis instance
    'arcana_paper',
    # Repository is a simple directory on the local file system
    repository=DirectoryRepository(args.data_dir),
    # Use a single process on the local system to derive
    processor=LinearProcessor(args.work_dir),
    # Use the static environment (i.e. no Modules)
    environment=StaticEnvironment(),
    # Match names in the data specification to filenames used
    # in the repository
    inputs=[
        FilesetSelector (\ 't1\_magnitude\ ',\ dicom\_format\ ,\ args.t1\ ,
                         is_regex=True),
        FilesetSelector('t2star_channels', zip_format, args.t2star_chann,
                         is_regex=True),
        FilesetSelector ('t2star_header_image', dicom_format, args.swi,
        is\_regex=True)\,,\\ FilesetSelector('swi\_magnitude', dicom\_format, args.swi,
                         is_regex=True),
        FilesetSelector('dmri_magnitude', dicom_format, args.dmri,
                         is_regex=True),
        FilesetSelector('dmri_reverse_phase', dicom_format, args.distort,
                         is_regex=True)],
    parameters={
        'dmri_num_global_tracks': int(1e5),
        'dmri_global_tracks_cutoff': 0.175,
        # This is needed as the channels aren't reconstructed with the
        # correct headers
        ; t2star_force_channel_flip ': ['-x', '-y', 'z']})
# Derive required data and display them in a single step for each
# figure.
paper.vein_fig(op.join(args.fig_dir, 'veins.png'))
paper.fa_adc_fig(op.join(args.fig_dir, 'fa_adc.png'))
paper.tractography_fig(op.join(args.fig_dir, 'tractography.png'))
```