# ARAX: a graph-based modular reasoning tool for translational biomedicine

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## ABSTRACT

**Motivation:** With the rapidly growing volume of knowledge and data in biomedical databases, improved methods for knowledge-graph-based computational reasoning are needed in order to answer translational questions. Previous efforts to solve such challenging computational reasoning problems have contributed tools and approaches, but progress has been hindered by the lack of an expressive analysis workflow language for translational reasoning and by the lack of a reasoning engine—supporting that language—that federates semantically integrated knowledge-bases.
**Results:** We introduce ARAX, a new reasoning system for translational biomedicine that provides a web browser user interface and an application programming interface. ARAX enables users to encode translational biomedical questions and to integrate knowledge across sources to answer the user's query and facilitate exploration of results. For ARAX, we developed new approaches to query planning, knowledge-gathering, reasoning, and result ranking and dynamically integrate knowledge providers for answering biomedical questions. To illustrate ARAX's application and utility in specific disease contexts, we present several use-case examples.
**Availability and Implementation:** The source code and technical documentation for building the ARAX server-side software and its built-in knowledge database are freely available online (https://github.com/RTXteam/RTX). We provide a hosted ARAX service with a web browser interface at arax.rtx.ai and a web application programming interface (API) endpoint at arax.rtx.ai/api/arax/v1.2/ui/.
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## 1 Introduction

Structured biomedical knowledge is rapidly accumulating in primary databases such as ChEMBL [1], DrugBank [2], Reactome [3], OBO Foundry [4], KEGG [5], Semantic MEDLINE Database (SemMedDB) [6], UniProtKB [7], and the Unified Medical Language System (UMLS) [8]. Using knowledge-based computational reasoning to answer translational questions such as *What drugs might be repurposed to treat Adams-Oliver Syndrome?* requires integrating

facts spanning a variety of concept types including drugs, targets, pathways, genetic variants, phenotypes, and diseases. To facilitate such computational reasoning, there have been numerous efforts to integrate knowledge from various biomedical databases using knowledge graph (KG) abstractions [9, 10, 11] consisting of a labeled multigraph in which each node represents a concept and each edge represents a concept-concept relationship, i.e., a "triple". Several such biomedical KGs have been described [12, 13, 14, 15, 16, 17, 18], including RTX-KG2, which we developed and described previously [19] and which integrates all of the aforementioned primary databases with a semantic layer that is described by the open-standard Biolink model [20]. Given such a comprehensive KG with a unified semantic layer, a biomedical question can be transformed into a *query graph* [21] (which represents a search pattern, analogous to a `select` statement in the SPARQL language [22]) and/or a *graph analysis workflow* in order to generate a list of "answers", each corresponding to a *subgraph* of the KG. To facilitate that process for translational applications, an open-source, web-based, and standards-based tool is needed for designing, expressing, executing, and refining KG analysis workflows.

Efforts to develop a comprehensive platform for expressing, executing and refining KG analysis workflows have used new query languages, web application programming interface frameworks, computational reasoning paradigms, graph topology heuristics, and machine-learning methods. BIOZON [23] provides a SQL-based query and search function for its KG; Hetionet [14] integrates multiple databases into a single graph for drug repositioning and leverages graph path-finding; BioGraph [24] uses the Gremlin graph query language; SPOKE [25] provides capabilities for finding a concept of interest and visualizing its "node neighborhood"; BioThings Explorer [26] builds a biomedical knowledge graph by querying SmartAPI-registered APIs; ROBOKOP [16] constructs a query-specific knowledge graph which it analyzes to identify candidate answers; and mediKarnren [27] uses the miniKanren logic programming system for drug repositioning applications. Despite much progress, two key problems remain: (1) how to provide control and transparency of the graph analysis workflow that is executed to answer a user's question and (2) how to rank potential answers. The control/transparency problem entails a trade-off: on one extreme, a general-purpose graph query language such as Neo4j Cypher[1] provides complete control, but expressing, modifying, and extending a state-of-the-art drug repositioning analysis workflow is extremely complex and demanding for many users. On the other extreme, canned analysis workflow approaches in practice have limited customizability. In addition, many tools have a limited number of knowledge sources and/or lack a standardized semantic layer with sufficient expressiveness.

### 1.1 ARAX overview and key advantages

Here we describe ARAX (arax.rtx.ai), a new computational reasoning tool for querying and exploring biomedical knowledge and data. ARAX provides three key advantages versus previous systems:

1. ARAXi, ARAX's intuitive language for specifying a workflow for analyzing a knowledge graph (Sec. 2.1);
2. access to 15 knowledge providers (which themselves access over 100 underlying knowledge sources) from a single reasoning tool, using a standardized interface and semantic layer (Sec. 2.2); and
3. a versatile method for scoring search result subgraphs (Sec. 2.1.5).

Together, these advantages are designed to facilitate and improve reasoning-based answering of translational questions [28] such as *What genetic conditions protect against asthma? What drugs target proteins associated with the cyclooxygenase pathway?* and *How can expression of KCNMA1 be pharmacologically inhibited?*

ARAX is one of six reasoning engines in the Biomedical Data Translator system (henceforth, "Translator") [29, 30], (see "Funding"), a distributed computational system for accelerating translational science. Translator has a layered architecture; queries are interpreted by a coordinating service, which sends the query to the reasoning engines which, in turn, consult registered Knowledge Provider (KP) services to answer the query. All Translator services exchange messages using a standard web interface, the Translator Reasoner API (TRAPI),[2] whose technical specification is collaboratively maintained. For its semantic layer, TRAPI uses the Biolink Model[3] to specify types of biological entities and relationships. ARAX can be directly queried via either its TRAPI API or its web browser user interface; these provide various ways to formulate queries and explore results (Sec. 2.3).

ARAX's architecture (Figure 1) is modular, with five different analysis modules that can be run using ARAXi commands (Sec. 2.1). ARAX is mainly built on top of RTX-KG2 [19], our large-scale biomedical knowledge graph that has a TRAPI-compliant query interface and is registered as a Translator KP; but ARAX also queries 14 other KPs with specific areas of emphasis (Sec. 2.2.1), including genome-wide association study data, electronic health record data from medical centers, and molecular data from high-throughput cellular assays.

[1] `neo4j.com/developer/cypher`
[2] `github.com/NCATSTranslator/ReasonerAPI`
[3] `github.com/biolink/biolink-model`

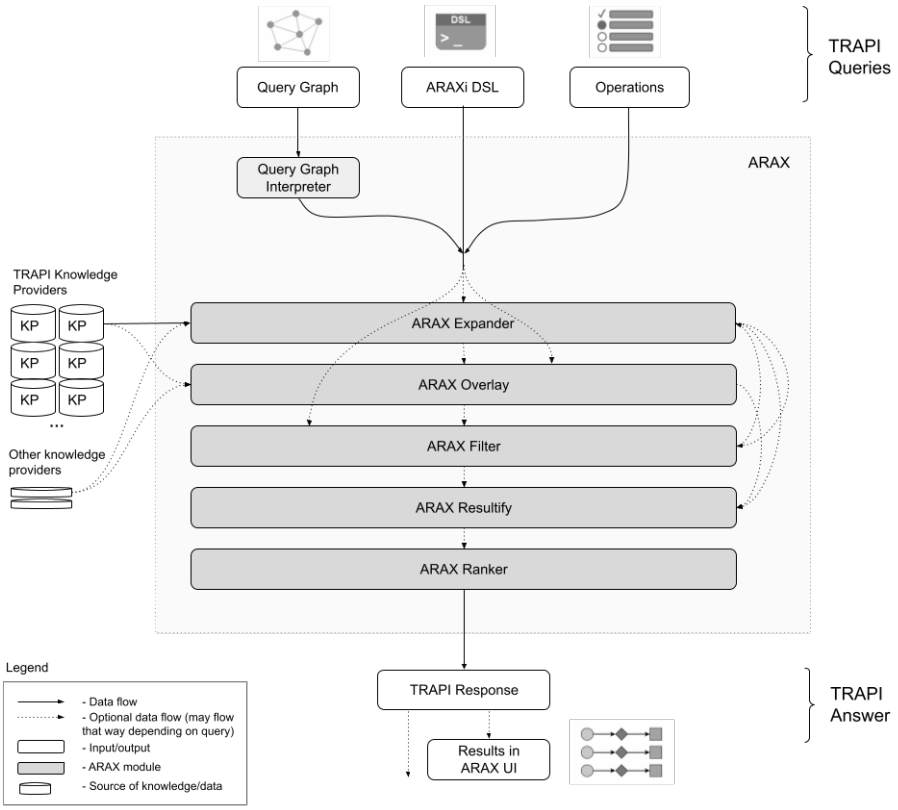


Figure 1: **ARAX's architecture is centered around five modules**. ARAX accepts queries in three different forms, all of which are based on the Translator Reasoner API (TRAPI) data model: a query graph, the ARAXi graph analysis workflow definition language, or TRAPI Operations. The ARAX system translates the two non-ARAXi input forms into an ARAXi workflow that specifies the series of ARAX modules that will be run to answer the query. The five core ARAX modules (Expander, Overlay, Filter, Resultify, and Ranker) operate independently and may be combined in various orders depending on the query. When the input is in the form of a query graph, the Query Graph Interpreter layer selects an appropriate series of ARAXi commands to answer the question at hand; this selection of ARAXi commands is done using a template-based system in which templates were manually curated by examining common query graph structures or archetypes. Both the Expander and Overlay modules pull in knowledge from external sources, either via the source's TRAPI API or a local copy of its dataset. Expander, Overlay, and Filter together build and prune a knowledge graph that combines individual knowledge providers' answers to the query. Resultify then finds all subgraphs in that answer knowledge graph that satisfy the input query and Ranker assigns each result a score. The final knowledge graph and results are returned in a TRAPI JavaScript Object Notation (JSON) response either directly to the requesting software program or to the ARAX UI for browser rendering (Sec. 2.3.1).

As illustrated in Figure 1, ARAX accepts input queries in any of three representations: a TRAPI query graph, ARAXi, or TRAPI Operations. A query graph is a graph-based template representing the user's question; it defines the structure that answers to the question must have. TRAPI query graphs consist of query nodes, which may have one or more concept identifiers (e.g., `CHEMBL.COMPOUND:CHEMBL112`) or any number of Biolink categories (e.g., `biolink:Protein`) specified, and query edges, which may have one or more Biolink predicates specified (e.g., `biolink:interacts_with`). A visual representation of an example query graph is shown in Fig. 2. The ARAXi language, which can be used to express a custom graph analysis workflow including overlaying of annotations and/or filtering of edges to reduce the result-set size, is described in Section 2.1. TRAPI Operations[4], the third way of specifying a graph analysis workflow, is an ARAXi-inspired, Translator-standardized vocabulary; it is translated into ARAXi by the ARAX interpreter.

## 2 Methods

### 2.1 ARAXi knowledge graph analysis language

To address the need for an intuitive language for expressing KG analysis workflows, we developed ARAXi, a procedural language that allows a user or software tool to concisely express biomedical questions. The five main ARAXi modules (see Secs. 2.1.1–2.1.5) can be used individually or combined in a graph analysis workflow (Fig. 2). Each ARAXi command corresponds to an analysis module; this modular design simplifies both construction and reuse of knowledge graph analysis workflows.

All input queries—whether in the form of a query graph, ARAXi, or TRAPI Operations—are ultimately translated into a series of ARAXi commands, which together define the knowledge-retrieval and analysis workflow for answering the query. In particular, two core ARAXi commands, `add_qnode` and `add_qedge`, are used to incrementally construct a query graph, which can then be expanded by the ARAX_expander module (Sec. 2.1.1) into a query-specific knowledge graph (KG); the query-specific KG can then be refined or have knowledge overlaid using the four other ARAX modules (Secs. 2.1.2–2.1.5).

#### 2.1.1 Expand a knowledge graph: ARAX_expander

The ARAX_expander module queries various knowledge providers (KPs) (see Sec. 2.2.1) to find all subgraphs that satisfy the input query graph. For each query edge in the query graph, ARAX_expander determines which KPs are capable of fulfilling that query edge (based on metadata provided dynamically by the KPs' APIs) and then queries those KPs concurrently. KPs are queried via their TRAPI-compliant APIs, where available. When querying each KP, ARAX_expander converts any node identifiers in the query to semantically equivalent identifiers from controlled vocabularies that the KP prefers. This is accomplished by using the ARAX Node Synonymizer service (Section 2.2.2). ARAX combines answers returned from the KPs into a single answer knowledge graph that is "canonicalized," meaning that it does not contain semantically redundant nodes.

ARAX_expander processes the query graph in a breadth-first fashion; that is, it retrieves all answers from KPs for a given query edge before moving on to the next query edge. It avoids combinatorial explosion by pruning down the answer set that it receives for each query edge in an adaptive and predictive manner: it uses the ARAX_overlay module's Fisher Exact Test function (Sec. 2.1.2) combined with ARAX_ranker (Sec. 2.1.5) to decide which triples to retain for the given query edge. By default, the number of triples to retain for each query edge (the "pruning threshold") is dynamically determined according to the projected degree of combinatorial explosion the given query edge will produce (based on heuristics pertaining to the depth of categories and predicates in the Biolink hierarchy). Alternatively, a user may specify a per-query pruning threshold that overrides the dynamically-determined default pruning.

Because users have different preferences in terms of the amount of time they are willing to wait for ARAX to provide answers to their query, ARAX_expander also provides a KP timeout parameter, which specifies how long ARAX_expander should wait for a response from each KP. Users who are more interested in getting answers quickly can specify a short (e.g., 30 seconds) timeout, while a user whose use-case prioritizes comprehensiveness over expediency can use a longer KP timeout.

#### 2.1.2 Overlay/annotate contextual information: ARAX_overlay

The ARAX_overlay module enhances the ARAX_expander-built KG by overlaying "virtual edges" (which are virtual in the sense that they are not present in the original query graph) to denote structural similarity (i.e., a statistically significant number of shared neighbors) of two nodes in the KG or to denote statistically significant co-association

[4] `github.com/NCATSTranslator/OperationsAndWorkflows`

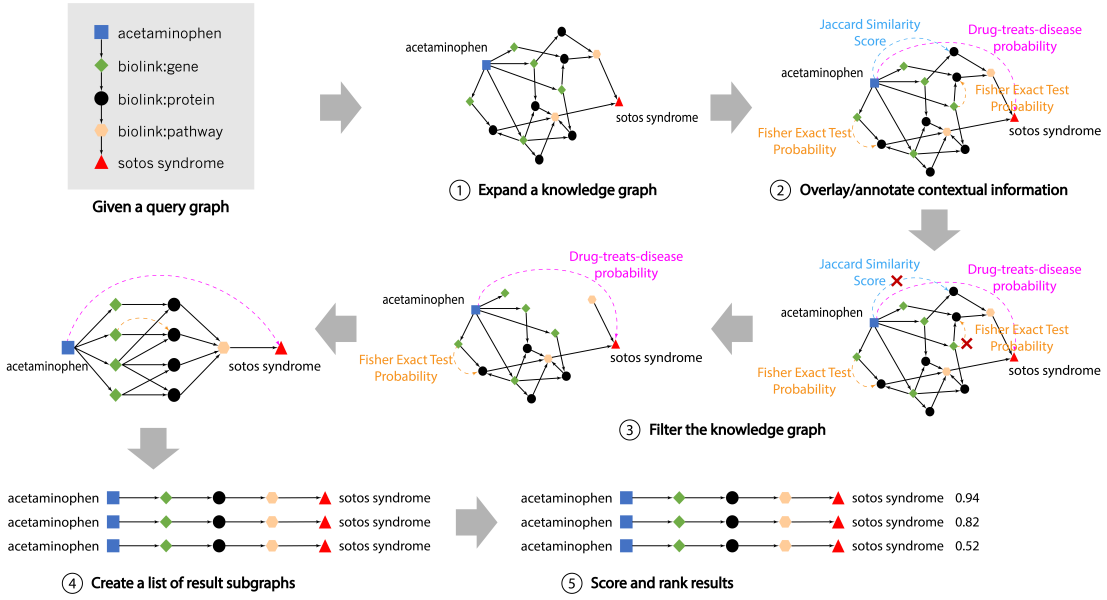


Figure 2: An abstract representation of a typical ARAXi workflow utilizing the five primary ARAXi modules: 1) Expand the query graph to an answer knowledge graph by consulting the registered Knowledge Providers; 2) Overlay/annotate the answer KG with contextual quantitative information in "virtual" edges (dashed lines); 3) Filter nodes and edges from the answer KG based on the annotated statistical information in virtual edges; 4) Create a list of result subgraphs in the answer KG fulfilling the query graph; 5) Calculate a score for each result subgraph indicating the degree of overall epistemic support for that answer.

in a database such as a clinical database, a combined clinical-epidemiological database, or the biomedical literature. Currently, the overlay function (which is called with the `overlay` command in ARAXi) can overlay nodes or edges with the following seven kinds of contextual information:

**Fisher Exact Test** – evaluates how important/significant an association is between a subject node and an object node in a one-hop connection. Given an edge in a one-hop query, its annotated $p$-value is computed based on a $2\times2$ contingency table defined by (a) the number of nodes in the subject side of this one-hop query which are connected to the object node in this edge; (b) the number of nodes in knowledge providers with the same category as the nodes in the subject side which are connected to the object node in this edge but not in the subject side of this one-hop query; (c) the number of nodes in the subject side but not connected to the object node in this edge; and (d) the number of nodes in knowledge providers with the same category as the nodes which are not in the subject side and not connected to the object node in this edge.

**Jaccard Similarity** – measures how many shared intermediate nodes exist between a subject node and an object node. Given a user-defined start query node identifier (e.g., "n01"), a user-defined intermediate query node identifier (e.g., "n02") and a user-defined end query node identifier (e.g., "n03"), the Jaccard similarity is defined to be the number of shared intermediate nodes with the given query node identifier divided by the union number of intermediate nodes with that query node identifier between the start query node identifier and the end query node identifier.

**Drug-Treats-Disease Probability** – predicts the probability that a given drug can treat a given disease. We developed this computational drug repurposing approach based on the model framework proposed by [31]. In order to adapt this model to our core knowledge graph RTX-KG2, we modified the implementation of the model by replacing node2vec [32] with GraphSage [33] for generating the node embeddings as well as using direct concatenation instead of the Hadamard product to generate the input features for a Random Forest classifier for drug-disease pairs.

**Normalized PubMed Distance** – measures the significance of co-occurrence of pairs of terms (corresponding to node names) in article abstracts in the PubMed (i.e., MEDLINE) database, using the "Normalized Google Distance" (NGD) measure [34]

$$\mathrm{NGD}(t_1, t_2) = \frac{\max\{\log f(t_1), \log f(t_2)\} - \log f(t_1, t_2)}{\log N - \min\{\log(t_1), \log(t_2)\}}, \tag{1}$$

where $t_1$ and $t_2$ are the biomedical terms (node names) used in RTX-KG2; $f(t_1)$ and $f(t_2)$ respectively represent the total number of unique PubMed IDs associated with this term; $f(t_1, t_2)$ is the number of unique and shared PubMed IDs between $t_1$ and $t_2$; and $N$ is the number of pairs of article abstract and MeSH term [35] annotations in PubMed.

**PubMed abstracts** – obtained by mapping node identifiers to MeSH terms and obtaining the PubMed articles associated with this MeSH term.

**Columbia Open Health Data (COHD) Clinical Information** [36] – obtained from electronic health records of 1.7M patients during a five-year period. For a given pair of terms, three statistical measures are obtained from COHD: observed clinical frequencies; the log-ratio between observed and expected counts; and the $\chi^2$ test statistic.

**Exposure Data** – statistical association data for pairs of terms in health records and between clinical terms and environmental exposure variables in epidemiological databases, obtained through the Integrated Clinical and Environmental Exposures Services (ICEES) service [37], which obtains data from University of North Carolina Health; the Drug Induced Liver Injury Network; and the National Institute of Environmental Health Sciences Personalized Environment and Genes Study.

#### 2.1.3 Filter the knowledge graph: ARAX_filter

Multi-hop queries can produce such large answer KGs that it is sometimes useful to prune down the KG to remove less important nodes or edges before proceeding. The ARAX_filter module allows one to do this; it provides several options for selectively removing nodes/edges based on their contextual information (often quantitative information added by ARAX_overlay) according to user-defined thresholds. This functionality can be invoked using the `filter_kg` ARAXi command. ARAX_filter also provides various methods for limiting the number of results returned and for sorting the results created by ARAX_resultify (see Sec. 2.1.4), for example, by score, edge/node counts, or particular edge/node attributes.

#### 2.1.4 Create results: ARAX_resultify

The ARAX_resultify module finds and returns all subgraphs in the answer KG that fit the structure of the query graph; this step is typically performed after the KG has been constructed and (optionally) pruned. ARAX_resultify can either ignore or enforce edge direction when determining whether a subgraph matches the query graph.

#### 2.1.5 Score and rank result graphs: ARAX_ranker

The ARAX_ranker module assigns a score between 0 and 1 (where a higher score represents a higher-confidence answer) to each result graph and orders the results by descending score. Scores are computed as follows:

1. For each "result graph":
   a. Each edge in the result graph is assigned a score [0,1], as follows:
      i. Any scalar attribute (e.g., Fisher Exact Test $p$-value, or Jaccard Index) on the edge is normalized to a value between 0 and 1 using a sigmoid function parameterized for the specific type of attribute.
      ii. For any edges originating from SemMedDB, the attribute containing PubMed references for publications supporting the given assertion is converted into a scalar value via the function:
      $$\left[1 + \left(\frac{b}{n}\right)^a\right]^{-1}, \tag{2}$$
      where $n$ represents the number of publications listed in the attribute. The values of $a = 3.16993$ and $b = 4$ were chosen so that 4 supporting publications results in a value of 0.5 (i.e. 50% confidence), and 8 publications gives a value of 0.9.
      iii. The given edge's normalized, weighted scalar values are then multiplied to create one single score between 0 and 1 that is assigned to that edge.
   b. Using the normalized, weighted edge scores from Step 1a, three graph weights (calculated using max-flow, Frobenius norm, and weight-of-longest-geodesic-path) are obtained for the result graph.
2. Results are then ranked in descending order by their three metric scores calculated in Step 1b (in three separate lists).
3. Each result is assigned a final score that is the average of its three ranks from Step 2 (normalized between 0 and 1).

The ARAX system automatically runs ARAX_ranker after ARAX_resultify, but ARAX_ranker can also be run individually via the `rank_results` ARAXi command; in principle, a list of result subgraphs produced by another Translator tool could be ranked using this ARAX service.

Table 1: ARAX integrates knowledge from 15 provider services. Abbreviations: Meth., Query Method; Cat., Categories; Pred., Predicates; Meta., Meta-triples. Links to the APIs are available in the SmartAPI registry (Footnote 7).

| Knowledge provider | Meth. | Cat. | Pred. | Meta. |
|---|---|---|---|---|
| Biothings Explorer (BTE) | TRAPI | 15 | 265 | 75,554 |
| Columbia Open Health Data (COHD) | TRAPI | 5 | 2 | 50 |
| Connections Hypothesis Provider | TRAPI | 6 | 12 | 32 |
| Drug-Treats-Disease (Sec. 2.1.2) | SQL | 5 | 1 | 25 |
| Genetics Provider | TRAPI | 4 | 3 | 8 |
| ICEES Asthma Instance | TRAPI | 13 | 2 | 338 |
| ICEES Drug-Induced Liver Injury Instance | TRAPI | 9 | 2 | 162 |
| Molecular Provider (MolePro) | TRAPI | 12 | 125 | 287 |
| Multiomics Clinical HER Risk Provider | TRAPI | 15 | 2 | 24 |
| Multiomics Drug Response Provider | TRAPI | 15 | 4 | 4 |
| Multiomics Tumor Gene Mutation Provider | TRAPI | 15 | 2 | 2 |
| Multiomics Wellness Provider | TRAPI | 15 | 2 | 17 |
| Normalized PubMed Distance (2.1.2) | SQL | 56 | 1 | 3,136 |
| RTX-KG2 | TRAPI | 57 | 86 | 45,300 |
| SPOKE | TRAPI | 18 | 24 | 44 |

## 2.2 Knowledge/Data

### 2.2.1 Richness

ARAX uses 15 different Knowledge Providers to answer queries, 13 of which are accessed via their TRAPI APIs (Table 1). The KPs themselves obtain information from various knowledge sources, giving ARAX access to a combined total of over 100 knowledge sources (e.g., UniProt, DrugBank, DisGeNET, ChEMBL, MONDO, and SemMedDB). Calculating the exact count of underlying knowledge sources is difficult due to the lack of such information in structured form for some KPs, but between RTX-KG2's 73 knowledge sources [19], SPOKE's approximately 40 knowledge sources[5], and BTE's 32 integrated sources[6], ARAX has access to at least 100 distinct underlying knowledge sources (we note that there is some overlap in knowledge sources between KPs). The true count of underlying knowledge sources is likely higher since ARAX uses 12 other KPs that were not included in the above estimate.

In total, ARAX's KPs can answer queries about 66 different categories of nodes (e.g., `biolink:Protein`, `biolink:Disease`) and 267 different edge predicates (e.g., `biolink:interacts_with`, `biolink:treats`). This results in a total of 75,587 distinct meta-triples (combinations of subject node category, edge predicate, and object node category) that ARAX has access to. The breakdown of these counts by KP is shown in Table 1. Figure 3 shows a small selection of ARAX's overall meta-graph, including some of the most commonly queried node categories. The counts for each KP are based on their TRAPI v1.2.0 API `/meta_knowledge_graph` endpoints, accessed on May 9, 2022. All such APIs are listed in the SmartAPI Registry online.[7])

### 2.2.2 Knowledge Federation

Two aspects of ARAX's architecture are crucial for knowledge federation: its adherence to standards for query APIs and for its semantic layer (i.e., TRAPI and Biolink, respectively) and its comprehensive service for mapping between equivalent node identifiers (i.e., node synonymization).

**TRAPI and the Biolink Model**
ARAX bypasses many of the data integration challenges that arise when combining information from multiple sources by adhering to a standard web API format (TRAPI [2]) that itself adheres to a standard semantic format (Biolink[3]). Not only does ARAX's API conform to TRAPI, but in the course of answering queries, ARAX's design favors using KPs that themselves speak TRAPI. This means that ARAX has to do very minimal processing to relay queries to and combine answers from different KPs, since they all represent their answers in the same format, using the same node categories and edge predicates. Because of this, any KP that speaks TRAPI can be plugged into the ARAX_expander module with ease. In the future, we expect ARAX_expander to dynamically select TRAPI KPs that are registered in the SmartAPI [38] registry, meaning ARAX will be able to add new KPs without any human intervention.

[5] `spoke.ucsf.edu/data-tools`
[6] `github.com/biothings/biothings_explorer`
[7] `smart-api.info/registry`

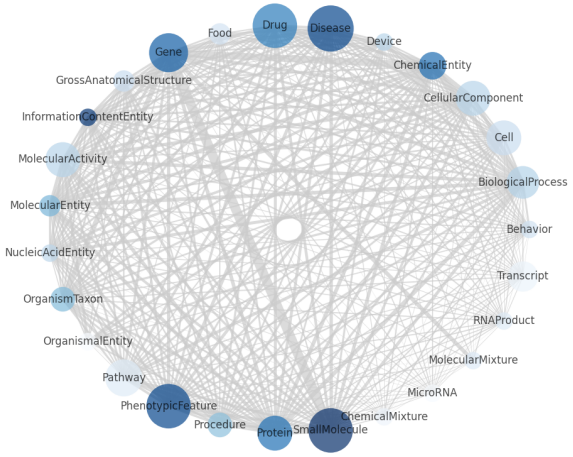


Figure 3: A small portion of ARAX's meta-graph including a selection of the most commonly queried node categories. The size of each node represents the number of ARAX's KPs that can answer queries involving that category, the color of each node represents the number of node identifier types ARAX has access to (through its KPs) for that category, and the thickness of each edge represents the number of distinct predicates ARAX has access to between two nodes with the given categories.

**ARAX Node Synonymizer**

Because there are many overlapping controlled vocabularies within biomedicine, concepts (or the nodes that represent them) can often be described using multiple identifiers. For instance, MONDO:0019391, DOID:13636, and ORPHANET:84 are all valid identifiers for the disease Fanconi anemia, coming from the MONDO Disease Ontology [39], the Disease Ontology [40], and the Orphanet Rare Disease Ontology [41, 42], respectively. Different knowledge providers may refer to the same concept using different identifiers, making it challenging to integrate their results. To address this problem, we created the ARAX Node Synonymizer, a node synonym mapping service built into ARAX. The Node Synonymizer determines node equivalencies by combining four kinds of evidence:

1. Concept equivalence information provided by a Translator web service called the Standards and Reference Implementation Node Normalizer[8],
2. Nodes connected by `biolink:same_as` relations in RTX-KG2,
3. Nodes with identical names, and
4. Node semantic type compatibility.

The Node Synonymizer uses this information to partition a given set of node identifiers into sets of semantically equivalent identifiers. Each of these clusters is assigned a single representative concept identifier from among the cluster's members. ARAX uses the Node Synonymizer to map all node identifiers returned from knowledge providers to these canonical identifiers, facilitating merging of the knowledge providers' responses.

## 2.3 Interfaces for accessing ARAX

### 2.3.1 ARAX Web Browser Interface

The ARAX Web Browser User Interface ("ARAX UI"; arax.rtx.ai) provides an intuitive, human-friendly mechanism for querying ARAX and exploring the answers it returns. It allows users to formulate biomedical questions in four different formats: an interactive visual query graph builder, TRAPI Operations, TRAPI JSON, and ARAXi. The interactive query graph builder allows users to construct a query graph in a visual fashion via drop-down lists and clickable buttons, with no JSON or TRAPI knowledge required. The TRAPI Operations input aids users in creating a TRAPI-compliant Workflow by providing a drop-down list of allowable Operations and guided entry of Operation parameters. The TRAPI JSON input method facilitates sharing and re-use of graph analysis workflows. The ARAXi input method (described in Section 2.1) provides users with easy-to-understand syntax to formulate more complex query workflows, along with drop-down menus of ARAXi commands to facilitate discovery. For each query posted to the ARAX UI, the output includes five sections:

[8] `github.com/TranslatorSRI/NodeNormalization`

1. **Summary**: Summarizes the answers to the query in a simple table sorted by ARAX-defined scores that can be considered a measure of confidence (Sec. 2.1.5).
2. **Knowledge Graph**: Provides a visual representation of the answer KG for easy viewing of its topology.
3. **Results**: Provides a display of all the results that satisfy the input query graph, ranked as in Section 2.1.5. Clicking on individual results displays an interactive graphical representation of the result. Within each result, clicking on nodes and edges displays detailed information including evidence/provenance information and concept descriptions.
4. **Messages**: Shows all messages logged during processing of the query, including any errors or warnings.
5. **Provenance**: When available, shows the number of edges in the final KG based on their predicate types and the knowledge sources that they came from.

The ARAX UI provides persistent URLs that enable fast retrieval of results from previously run queries. Apart from these main functions, the ARAX UI also provides three additional tools/services named as follows:

- **Synonym Lookup**: Exposes the ARAX Node Synonymizer (described in "ARAX Node Synonymizer" in Section 2.2.2); a user can input a concept identifier or name and see its set of equivalent identifiers and other synonym information.
- **Dev Info**: Provides a JSON-formatted view of UI-server communications for development purposes.
- **System Activity**: Displays the status and activity of previous queries submitted to ARAX over a user-selectable time interval and retrieves corresponding queries and results (if they completed successfully).

#### 2.3.2 ARAX API

ARAX can also be used via its publicly-accessible API[9], which is listed in the SmartAPI registry [38]. The ARAX API includes `/query` and `/asyncquery` endpoints that accept TRAPI input, as well as an `/entity` endpoint that exposes the ARAX Node Synonymizer service, allowing users to programmatically retrieve equivalent identifiers and other synonym information for a given concept.

## 3 Results

To illustrate how ARAX can be used to explore biomedical knowledge in a focused manner, we evaluated ARAX using two different queries, one to search for molecular mechanisms in bipolar disorder and one to search for the mechanism of action for an antiviral for COVID-19 disease.

### 3.1 Case 1: Bipolar disorder

Bipolar disorder is a mental health condition that affects over 46M people worldwide [43]. It is characterized by mood swings that can include excessive happiness (mania) and/or excessive sadness (depression). The causes of bipolar disorder are not well understood, but are thought to include both genetic and environmental factors, with stress and substance abuse contributing to disease severity. It has been recently proposed [44] that decreased expression of D-amino acid oxidase (DAO) in cerebellar neurons might increase the risk for bipolar disorder by affecting the regulation of N-methyl-D-aspartate receptors (NMDARs). Using the ARAX web browser interface, we explored the connection between DAO and bipolar disorder via protein intermediaries in the context of a two-hop query graph (e.g., "DAO-—$\star$-—bipolar disorder" where $\star$ represents a peptide/protein mediator). Within the browser interface, we posted the ARAXi commands of a two-hop query graph to ARAX, which returned 17 result subgraphs (Fig. 4). The query and its results are available as online supplementary material.[10] The "N-methyl-D-aspartate receptors" result (which we expected to see based on prior literature [44]) is in the top 10 highest-scoring results, based on ARAX's result-graph ranking algorithm (see Sec. 2.1.5). For some of the other peptide/protein results returned, ARAX provided hyperlinks to publications that support the relevant relations in the specific DAO-to-bipolar result graph such as the amino acid proline (Fig. 5).

[9] `arax.rtx.ai/api/arax/v1.2/openapi.json`

[10] `github:RTXteam/RTX/notes/vignettes.md#bipolar-disorder`

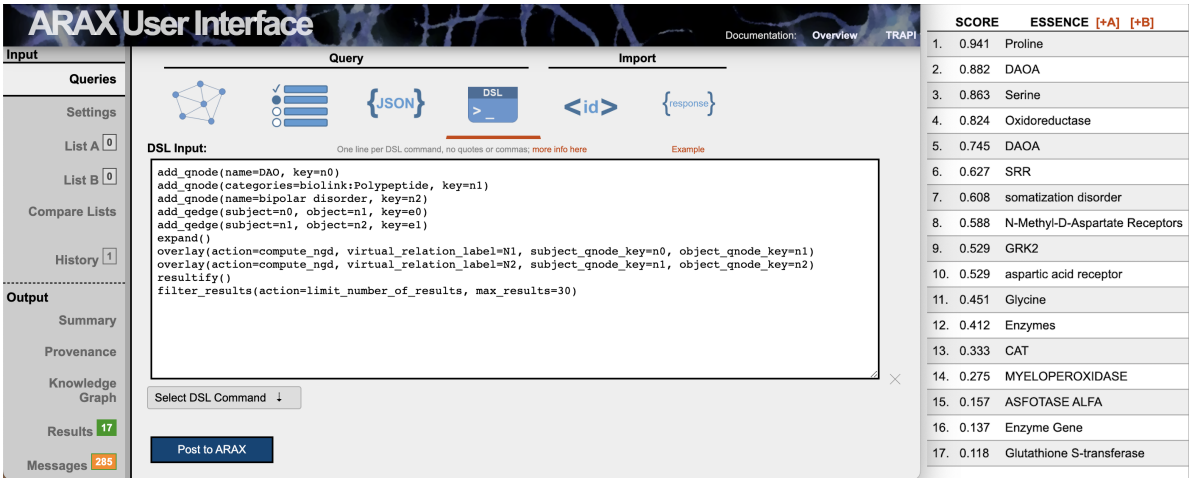


Figure 4: ARAX UI results with its corresponding ARAXi commands for Case 1 (amino acid connections between DAO and bipolar disorder).

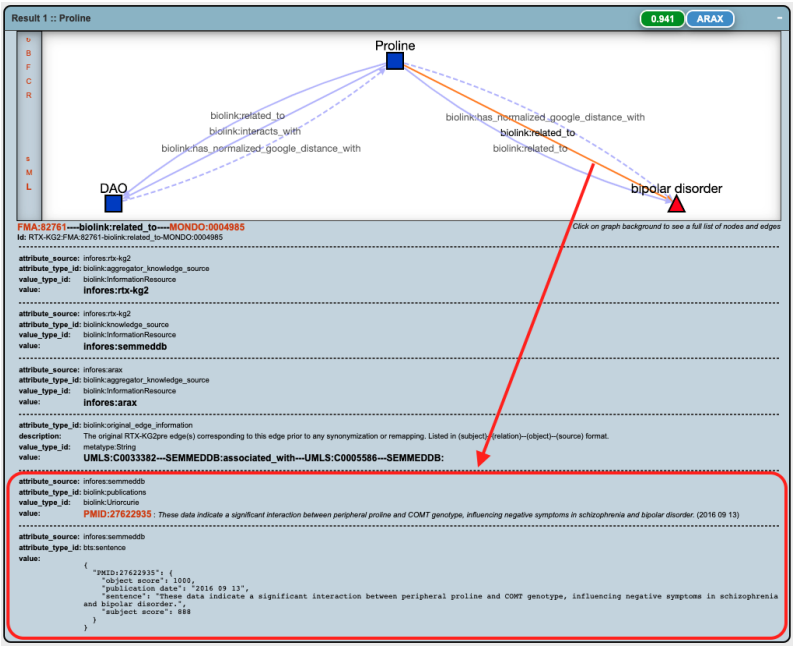


Figure 5: A "result graph" for a single result ("proline") for the two-hop query for mediators between DAO and bipolar disorder, with supporting publication (`PMID:27622935`) and its corresponding relevant sentence shown (highlighted by a red square) for a specific edge “Proline—`biolink:related_to`—bipolar disorder”.

### 3.2 Case 2: Coronavirus disease 2019

Coronavirus disease 2019 (COVID-19), caused by the SARS-CoV-2 pathogen, has resulted in a worldwide pandemic. Researchers have made extensive efforts [45, 46, 47, 48, 49] to repurpose existing drugs to treat COVID-19. Remdesivir was the first repurposed antiviral drug that was approved by the United States Food and Drug Administration for the treatment of COVID-19. Remdesivir was originally developed to treat hepatitis C and then found to be effective against many other viral diseases including Ebola and other coronavirus diseases. Its mechanism of action is to interfere with the viral RNA-dependent RNA polymerase (RdRp) [50]. By using ARAX, we can provide a series of potential three-hop biological-relation paths between remdesivir and COVID-19 via RNA-directed RNA polymerase. Figure 6 shows the top 25 ARAX results with the corresponding ARAXi commands. The query and its results are available as online supplementary material.[11] A top-ranking result, "Virus Replication", is precisely the mechanism of action that has been reported by various researchers [51, 52, 50]. Furthermore, ARAX results can reveal potential downstream mediators of therapeutic efficacy such as release of type I interferon, an antiviral cytokine which SARS-CoV-2 infection is known to inhibit [53] (see result "IFNB1" for this query).

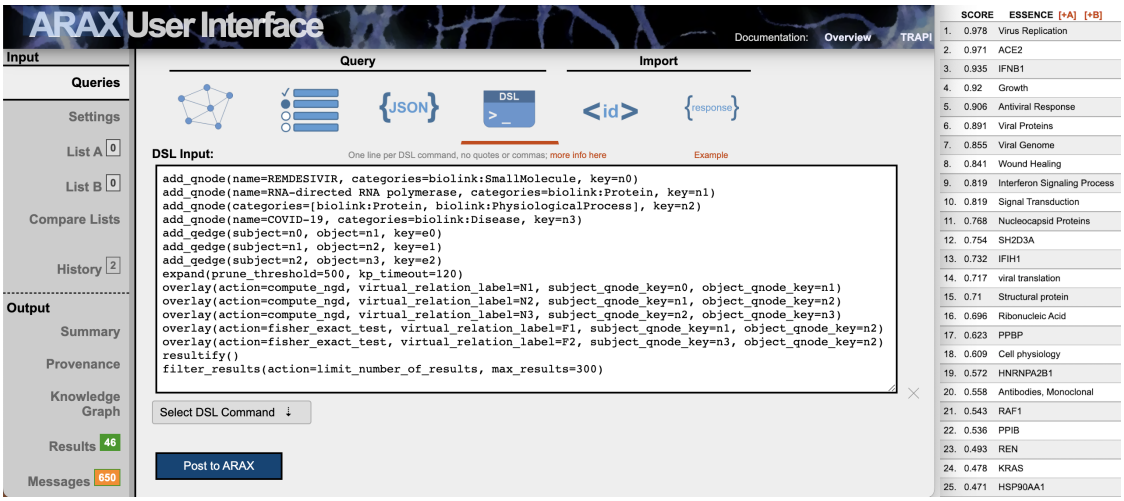


Figure 6: ARAXi workflow for finding the remdesivir mechanism-of-action downstream of RdRp for treating COVID-19 (Case 2), along with the top 25 result pathways.

## 4 Discussion

The purpose of ARAX and Translator is to enable integrated analysis of structured biomedical knowledge in order to provide ranked, coherent answers to translational biomedical questions. ARAX's three key innovations—ARAXi, its 15 knowledge providers, and its result subgraph ranking algorithm—together provide significant leverage to enable a researcher to tackle translational questions that are more complex and rely more on basic science (and not just on clinically validated) knowledge. Further, ARAX's web browser interface enables composing queries and browsing/exploring results, without having to programmatically post-process results or query the API.

While it shares the goal of leveraging knowledge to advance translation, ARAX is not an AI-based Biomedical Question-Answering system such as IBM Watson [54], MedQA [55], or BioSQUASH [56]. Such question-answering systems can provide a direct semantic answer computationally extracted from relevant biomedical documents via natural language processing (NLP) techniques. Thus, the knowledge used in these systems are only limited to document-based knowledge. Further, ARAX differs from the NLP-based document search tool BioMed Explorer[12]; BioMed Explorer is designed to answer a question by finding relevant resources (such as articles) using semantic search and showing excerpts from those resources. ARAX, in contrast, is based on multiple types of data sources (e.g., publications, databases, electronic health records, etc.). Instead of directly providing an intuitive answer, ARAX is more like a "searching & computing engine" where it provides a self-developed domain language to translate a specific biomedical question into a query graph, "search" for answers from different knowledge providers, and then perform computations

[11] `github:RTXteam/RTX/notes/vignettes.md#covid-19`
[12] `sites.research.google/biomedexplorer/`

with the resulting graphs. Thus, it can facilitate querying and exploring a significant fraction of published biomedical knowledge and public biomedical knowledge-bases. However, unlike search engines or semantic search engines, ARAX allows users to create custom knowledge graph analysis workflows, as well as access reasoning modules for specific translational applications like drug repositioning.

ARAX was built to help biomedical researchers explore structured knowledge and help generate new hypotheses. ARAX is for research purposes and is not meant to be used by clinicians in the course of treating patients. As the system is being actively developed, there is no expectation that results from queries run on ARAX will be retained indefinitely.

## 5 Conclusion

ARAX is a computational reasoning tool that allows users to easily extract, explore, and analyze knowledge from diverse biomedical resources. With self-developed ARAXi, ARAX allows users to encode their complex biomedical questions into specific KG analysis workflows in an intuitive manner. These workflows can intelligently integrate data from multiple biomedical knowledge providers, analyze and filter that data, and rank the final results. The ARAX UI further simplifies this process by providing multiple query options and facilitating interactive exploration of query results. The ARAX API enables incorporating ARAX's knowledge retrieval and reasoning capabilities within a workflow involving non-ARAX tools. We believe that ARAX can help life sciences researchers to more effectively interpret new research findings and develop new hypotheses. ARAX is still under active development. Subsequent updates will add more knowledge providers and new capabilities to the system.

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## Abbreviations

KG, knowledge graph
KP, knowledge provider
TRAPI, Translator Reasoner application programming interface

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