A Platform for the Biomedical Application of Large Language Models

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Abstract

The wealth of knowledge we have amassed in the context of biomedical science has grown exponentially in the last decades. Consequently, understanding and contextualising scientific results has become increasingly difficult for any single individual. In contrast, current Large Language Models (LLMs) can remember an enormous amount of information, but have notable shortcomings, such as a lack of generalised awareness, logical deficits, and a propensity to hallucinate. To improve biomedical analyses, we propose to combine human ingenuity and machine memory by means of an open and modular conversational platform, biochatter, exemplified in the web application ChatGSE. We safeguard against common LLM shortcomings using general and biomedicine-specific measures and allow automated integration of popular bioinformatics methods. Ultimately, we aim to improve the Alreadiness of biomedicine and make LLMs more useful and trustworthy in research applications.

Main

Despite our technological advances, biology and biomedicine continue to pose incredible challenges [1]. We measure more and more data points with ever-increasing resolution to such a degree that their analysis and interpretation have become the bottleneck for their exploitation. One reason for this challenge may be the inherent limitation of human knowledge [2]. Even seasoned domain experts cannot know the implications of every molecule, be it metabolite, DNA, RNA, or protein, even in their own domain. In addition, biological events are context-dependent, for instance with respect to a cell type or specific disease.

Large Language Models (LLMs) of the current generation, on the other hand, can access enormous amounts of knowledge, encoded (incomprehensibly) in their billions of parameters [3,4,5]. Trained correctly, they can recall and combine virtually limitless knowledge from their training set. ChatGPT has taken the world by storm, and many biomedical researchers already use LLMs in their daily work, for general as well as bioinformatics-specific tasks [6,7]. However, the current, predominantly manual, way of interacting with LLMs is virtually non-reproducible, and their behaviour can be erratic. For instance, they are known to hallucinate: they make up facts as they go along, and, to make matters worse, are convinced - and convincing - regarding the truth of their hallucinations [7,8]. While current efforts towards AGI (Artificial General Intelligence) manage to ameliorate some of the shortcomings by ensembling multiple models [9] with long-term memory stores [10], the current generation of AI does not inspire adequate trust to be applied to biomedical problems without supervision [8]. Additionally, biomedicine demands greater care in data privacy, licensing, and transparency than most other real-world issues.

A major aim of computational biology is to distil high-dimensional molecular measurements into a humanly digestible form by projecting the measurements into a lower-dimensional space composed of gene programs, pathways, or other functional groupings of biological entities, for example via gene set enrichment analyses. However, even this distilled knowledge requires advanced expertise and thorough literature research to effectively interpret and exploit, and benchmarking the methods' performance is non-trivial [11].

To improve and accelerate this interpretation and exploration, we have developed BioChatter, a platform for communicating with LLMs specifically tuned to biomedical research, the use of which we demonstrate in a conversational web interface, ChatGSE (Figure 1). The platform guides the human researcher intuitively through the interaction with the model, while counteracting the problematic behaviours of the LLM. Since the interaction is mainly based on plain text, it can be used by virtually any researcher. We engineer prompts around the queries of the user to improve model performance

with regard to biomedicine, and automate the integration of popular bioinformatics methods, such as differential expression and gene set enrichment (Supplementary Note Prompt Engineering).

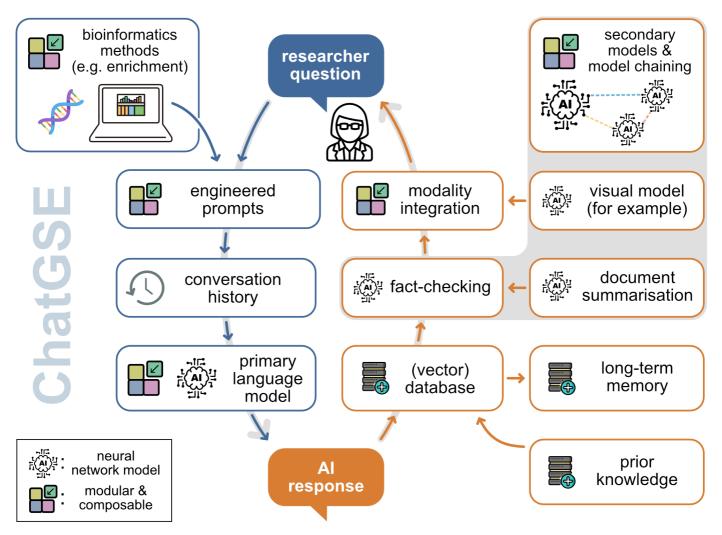


Figure 1: The ChatGSE composable platform architecture (simplified). The user submits a question about a topic of interest (e.g., an experiment) along with the low-dimensional results of a bioinformatics analysis (top left). The platform's main response circuit (blue) composes a number of specifically engineered prompts and passes them (and a conversation history) to the primary LLM, which generates a response for the user based on all inputs. This response is simultaneously used to prompt the secondary circuit (orange), which fulfils auxiliary tasks to complement the primary response. In particular, using search, the secondary circuit queries a database as prior knowledge repository and compares annotations to the primary response. The knowledge graph can also serve as long-term memory extension of the model. Further, an independent LLM receives the primary response for fact-checking, which can be supplemented with context-specific information by a document summarisation model. If this "second opinion" differs from the primary response, a warning is issued. The platform is composable in all aspects, in principle allowing arbitrary extensions to other, specialised models for additional tasks orchestrated by the primary LLM.

On the model side, we implement several measures in addition to the prompt engineering around the user's queries. For instance, we deploy a second model to safeguard the factual correctness of the primary LLM's responses (Supplementary Note Correcting Agent). These interactions are handled by a pre-programmed conversational "Assistant," which dynamically orchestrates LLM agents with distinct tasks using a Python model chaining framework [9]. Using vector database approaches, the user's prompts can be further supplemented with information extracted from pertinent, user-provided literature (Supplementary Note In-context Learning).

To increase data-awareness of the AI agents, we introduce connectivity to databases, which can extend the long-term memory of the models, semantically ground the biological entities with respect to suitable ontologies, and compare the model's responses to prior knowledge ground truth. By integrating a flexible knowledge graph creation framework [12], we allow versatile use cases across the entire research spectrum. For example, connecting to a knowledge graph of cell markers based on

Cell Ontology [doi:10.1186/s13326-016-0088-7], the task of annotating single cell data sets can be automated and made more reproducible (Supplementary Note <u>Cell Type Annotation</u>) by abstracting the pioneering efforts of manually executed studies [6]. We engineer the prompts around knowledge graph queries around the semantic configuration of BioCypher knowledge graphs, which allows efficient communication of the LLM with the knowledge graph and avoids hallucination-based issues. This allows querying any knowledge graph using natural language as well as the development of advanced multi-stage reasoning strategies (Supplementary Note <u>Knowledge Graphs</u>).

Currently, the most powerful conversational AI platform, ChatGPT (OpenAI), is surrounded by data privacy concerns [13]. We address this issue in two ways. Firstly, we provide access to the OpenAI models through the API (Application Programming Interface), which is subject to different, more stringent data protection than the web interface [14]. Secondly, we aim to preferentially support open-source LLMs to facilitate more transparency in their application and increase data privacy by being able to run a model locally [15]. Our orchestration tool supports dozens of LLM providers [9], such as the Hugging Face API, which can be used to query the recently released open-source ChatGPT-alternative HuggingChat or any other of the more than 100 000 open-source models on Hugging Face Hub [16]. Hugging Face also provide an open LLM leaderboard with up-to-date benchmarks of open-source LLMs [17]. Although OpenAI's models currently vastly outperform any alternatives in terms of both LLM performance and API convenience, we expect many open-source developments in this area in the future. Therefore, we support plug-and-play exchange of models to enhance biomedical AI-readiness.

In the future, we aim to integrate biological prior knowledge representation with LLM reasoning. Using the emergent strategies of in-context learning, instruction learning, and chain-of-thought prompting [18], this can enable causal inference on relationships between biological entities, for instance via protein-protein interactions (Supplementary Note Causal Inference), and automated validation of literature references provided by the LLM (Supplementary Note Literature Reference Database). While the current models do not yet appear suited for unsupervised reasoning in the biomedical space, they can already save much time otherwise spent on web and literature searches. Additionally, the ChatGSE platform provides a reproducible environment for benchmarking of models and engineered prompts to gauge their biomedical reliability. The ability to chain arbitrary types of models enables advanced applications, for instance connecting to visual modalities such as spatial omics. We provide further details and application scenarios in our Supplementary Notes.

While we focus on the biomedical field, the concept of the tool can easily be extended to other scientific domains by adjusting domain-specific prompts and data inputs, which in our framework are accessible in a composable and user-friendly manner. The Python library to interact with LLMs, vector databases, and all other features is developed openly on GitHub (https://github.com/biocypher/biochatter), and can be integrated into any number of user interface solutions apart from our own, for instance, INDRA [19] and drugst.one [20]. We develop under the permissive MIT licence and encourage contributions and suggestions from the community with regard to the addition of bioinformatics tool inputs, prompt engineering, safeguarding mechanisms, and any other feature.

Supplementary Materials

In our Supplementary Notes, we explain the functions of our platform in more detail. Please note that several of the mentioned features, particularly more advanced ones, are in early developmental stages. For practical reasons, we divide our software into two distinct packages, the user interface (UI) component ChatGSE and the Python backend library biochatter. This also reflects how we expect the platform to be used: as a generic backend library for the development of custom UIs. For an up-to-date overview and preview of current functionality of the platform, please visit the online preview. Both libraries are developed in Python (version 3.10), according to modern standards of software development [21]. We use Streamlit (version 1.21.0, https://streamlit.io) for the web UI. We include a code of conduct and contributor guidelines to offer accessibility and inclusivity to all that are interested in contributing to the framework.

Prompt Engineering

Recent experience with Large Language Models (LLMs) shows that the clever engineering of model prompts can yield drastic performance increases. For example, when performing logical inference, simply adding "Let's think step by step" to the end of a question prompt increased LLM performance from ~20% to ~80% [22]. As such, we highly prioritise the identification and adjustment of prompts tuned exactly to the requirements of the specific task to be performed, respecting "general" rules of interacting with LLMs as well as biomedicine-specific issues.

We designed the backend of ChatGSE to be completely flexible with regard to the application and rearrangement of prompts and prompt templates (which allow the insertion of variables such as the user input question or data). For templating, we use the corresponding generic functionality provided by LangChain [9], while adding our biomedicine-specific layer on top. We provide general prompts to the primary model, setting it up to be helpful and concise in its responses, and individual prompts for each tool we want the primary model to "understand." This includes explanation of the method itself as well as structural information about the data file containing the results.

To make this functionality available to all users, we provide a "Prompt Engineering" tab in the ChatGSE application, which allows the modification (or removal) of existing prompts, as well as the addition of new ones. We also provide functionality to import and export these prompts in JSON format to facilitate reproducible and shareable biomedical prompt engineering. To discuss and share examples of useful or ineffective prompts, we encourage all users to join the #chatgse stream in our freely accessible Zulip channel at https://biocypher.zulipchat.com, or the GitHub discussion thread at https://github.com/biocypher/ChatGSE/discussions/11.

To further improve reproducible prompt engineering, we will establish online datasets for benchmarking LLM performance on biomedical tasks. These datasets will be available through the ChatGSE frontend and the performance of specific prompt sets on these benchmarks will be captured using an online leaderboard system. This way, the most useful prompt sets for specific tasks can be maintained concurrently with the development of the LLMs.

Correcting Agent

The propensity of LLMs to hallucinate untrue facts necessitates control mechanisms and guardrails for their application in research, particularly in high-stakes fields such as biomedicine [7]. Some corrective incentive can be included in the instructions to the primary model, for instance by including a "Criticism" directive such as "Constructively self-criticise your big-picture behaviour constantly."

However, this does not guarantee to prevent hallucinations completely, as the same model that hallucinates also is responsible for correction.

Thus, we implement a modular combination of primary and corrective model, where the corrective agent – a "second opinion" – is set up with instructions tuned exactly to the task of fact-checking the responses of the primary model ("You are a fact-checker. Please judge the following statement for its factual correctness. ..."). The performance of the corrective agent can be further increased by using a more powerful model (e.g., moving from gpt-3.5-turbo to gpt-4), by pre-processing the primary model's response (e.g., splitting the response into single sentences and fact-checking each sentence individually), and comparing the primary model's statements to prior knowledge stored in a knowledge graph connected to the chat platform. As model development advances and other parties create models as powerful as OpenAl's, we foresee it will be useful to combine models from different suppliers to increase diversity between primary and corrective models. Using ChatGSE's modular framework, arbitrary numbers of corrective models can be added to the LLM chain.

To allow all users to interact with correcting functionality, we include a dedicated "Corrective Agent" tab for adjusting settings of the corrective model independent of the settings for the primary model. We also facilitate the testing of corrective agents and their prompts by providing a free-text field for sending false information to the corrective agent. This is necessary since some models, particularly those from OpenAI, are heavily regulated to not purposely provide false information, even for testing purposes.

The comparative power of the LLM agents can be further increased by connecting to a vector database containing embeddings of the contents of specific user-supplied documents. For more information, see the following Supplementary Note 3: In-context Learning.

In-context Learning

While the general knowledge of current LLMs is extensive, they may not know how to prioritise very specific scientific results, or they may not have had access to some research articles in their training data (e.g., due to their recency or licensing issues). To bridge this gap, we can provide additional information from relevant publications to the model via the prompt. However, we cannot add entire publications to the prompt, since the input length of current models still is restricted; we need to isolate the information that is specifically relevant to the question given by the user. To find this information, we perform a similarity search between the user's question and the contents of user-provided scientific articles (or other texts). The most efficient way to do this mapping is by using a vector database.

The contextual background information provided by the user (e.g., by uploading a scientific article of prior work related to the experiment to be interpreted) is split into pieces suitable to be digested by the LLM, which are individually embedded by the model. These embeddings (represented by vectors) are used to store the text fragments in a vector database; the storage as vectors allows fast and efficient retrieval of similar entities via the comparison of individual vectors. For example, the two sentences "Amyloid beta levels are associated with Alzheimer's Disease stage." and "One of the most important clinical markers of AD progression is the amount of deposited A-beta 42." would be closely associated in a vector database (given the embedding model is of sufficient quality, i.e., similar to GPT-3 or better), while traditional text-based similarity metrics probably would not identify them as highly similar.

By comparing the user's question to prior knowledge in the vector database, we can extract the relevant pieces of information from the entire background. These pieces (for instance, single sentences directly related to the topic of the question) are then sufficiently small to be directly added

to the prompt. In this way, the model can learn from additional context without the need for retraining or fine-tuning. This method is sometimes described as in-context learning [18].

To provide access to this functionality in ChatGSE, we add a "Document Summarisation" tab to the platform that allows the upload of text documents to be added to a vector database, which then can be queried to add contextual information to the prompt sent to the primary model. This contextual information is transparently displayed in the main chat window. Since this functionality requires a connection to a vector database system, we provide modular connectivity to several standard vector database providers, such as Pinecone, Weaviate, or Milvus.

Cell Type Annotation

A common repetitive task in bioinformatics is to annotate single-cell datasets with cell type labels. This task is usually performed by a human expert, who will look at the expression of marker genes and assign a cell type label based on their knowledge of the cell types present in the tissue of interest. LLMs have been shown to be able to perform this task with high accuracy, and can be used to automate cell type annotation with minimal human input [6].

We propose to combine LLM inference on cell types with the storage solutions provided by ChatGSE: using a vector database to store embeddings of cells for a more streamlined workflow, and using a traditional database on the basis of BioCypher [12] to inject prior knowledge into the process, as well as store the intermediate decisions of the model/user.

Using the graphical user interface of ChatGSE, we can provide recommendations of inferred cell types to the human user, such that the ultimate decision about a cell type annotation remains with the domain expert, while the tedious aspects of annotation are reduced significantly. If the model reaches a threshold of perfect annotation for any cell type (e.g., a >99% success rate in more than 50 cells), the user can decide to trust the model in instances of this cell type and fully automate the annotation in these instances. Similarly, confidence metrics can be used to trigger user input only at cell type inferences that are not straightforward.

Knowledge Graphs

Causal Inference

More recent models have shown considerable capacity for common sense reasoning, in particular, GPT-4 [5]. There is an unmet need to compare the reasoning outcomes of LLMs to other, more traditional modes of inference, such as the do-calculus [23], logic models [24], and semantic reasoning approaches [25]. With ChatGSE, we provide a Python platform for the side-by-side application of reasoning algorithms, in the "Causal Inference" tab.

The ability to connect to flexible databases created by BioCypher enables the representation of the same basic knowledge, but tailored to each individual reasoning mode (for instance, a labelled property graph as input for the logic model, an RDF graph for the semantic reasoner, and a vector database for the LLM). This increases the ease-of-use as well as the reproducibility of benchmarking the reasoning abilities of different algorithms.

Literature Reference Database

LLMs are very good feature extractors. In addition to the correcting agent described in Supplementary Note <u>Correcting Agent</u>, a "literature agent" can additionally be used to ameliorate the occasionally untruthful statements of the primary model. Given a response from said model, the literature agent is

tasked with extracting references to academic papers from the response (if it contains such), and validating the existence of the claimed reference as well as its attributes (such as title and digital object identifier). The validation of extracted article authors, titles, and identifiers can be performed by a simple search in a connected database of scientific publications.

Experimental Design

Experimental design is a crucial step in any biological experiment. However, it can be a subtle and complex task, requiring a deep understanding of the biological system under study as well as statistical and computational expertise. LLMs can potentially fill the gaps that exist in most research groups, which traditionally focus on either the biological or the statistical aspects of experimental design.

Similar to the general chat functionality, we provide facilities to upload experimental design plans and ask about their feasibility in the biological or statistical context in an "Experimental Design" tab. Coupled with a suitable knowledge graph (for instance containing methods literature) and/or a document summarisation of relevant articles, ChatGSE can be a simple and effective tool to consider the design of an experiment from multiple perspectives.

Podcast my Paper

The recent years have brought significant advances in text-to-speech applications, yielding large accessibility benefits to simple text-based communication and personal assistants. However, the text of scientific papers is riddled with special characters, references, legends, and entities otherwise hard to interpret by text-to-speech algorithms, such as PCR primer sequences.

To process arbitrary scientific manuscripts into "listenable" text, we create an agent that ingests the paper paragraph-wise and distributes individual sections to an LLM for text cleaning, removing all detracting information, such as reference numbers, web links, figure legends, and statistical information. The resulting cleaned sections are also lightly summarised to account for consumption by listening, and then fed into a text-to-speech model to generate individual sections (chapters) of the article. The resulting audio can be played back on any media device.

Journal Club

In many instances, the evaluation of scientific literature is pursued comparatively. Whether it is the discussion of which hypothesis is the correct one of two, or which method should be applied to address a specific question, multiple manuscripts need to be integrated to draw conclusions. We propose to facilitate scientific "discussion" by orchestrating an Al journal club, in which a primary model guides individual models, each responsible for representing one scientific position (e.g., each representing one manuscript).

This way of reasoning has proven effective in the way science is conducted by humans. By installing dedicated representatives of any position, the argument can be led more effectively, since each participant of the discussion is only responsible for knowing the facts that relate to their own position. In AI reasoning, problems could arise when one model is responsible for multiple sides of an argument, although for different reasons. In the case of argument being grounded in a vector database injection process (see Supplementary Note 3: In-context Learning), only one model in charge of representing two or more positions (manuscripts) on the matter may become biassed towards a specific position due to multiple issues. The amount of prompts that can be injected into one model query is limited by the token input length, and sentences from manuscripts that are by chance more similar to the phrase used for the similarity search could become overrepresented in the model's

reasoning. Using one model per position solves this problem by distributing token limits across multiple models, and by accessing only single manuscripts (or several manuscripts, but from the same "camp").

The models interact with one another in the way that human discussions are led: the moderator (primary model) gives tasks to the secondary models according to the human researcher's question, and elicits a response that the adversary needs to address in the next iteration of the discussion. The discussion, while being autonomous apart from the initial question, can be guided step-by-step by the human researcher.

Author Contributions

SL conceptualised and developed the platform and wrote the manuscript. JSR supervised the project, revised the manuscript, and acquired funding.

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Conflict of Interest

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References

1. Study reveals cancer's 'infinite' ability to evolve

BBC News

(2023-04-12) https://www.bbc.com/news/health-65252510

2. Capacity limits of information processing in the brain

René Marois, Jason Ivanoff

Trends in Cognitive Sciences (2005-06) https://doi.org/d5gmqt

DOI: <u>10.1016/j.tics.2005.04.010</u> · PMID: <u>15925809</u>

3. PaLM: Scaling Language Modeling with Pathways

Aakanksha Chowdhery, Sharan Narang, Jacob Devlin, Maarten Bosma, Gaurav Mishra, Adam Roberts, Paul Barham, Hyung Won Chung, Charles Sutton, Sebastian Gehrmann, ... Noah Fiedel *arXiv* (2022) https://doi.org/kfxf

DOI: 10.48550/arxiv.2204.02311

4. LaMDA: Language Models for Dialog Applications

Romal Thoppilan, Daniel De Freitas, Jamie Hall, Noam Shazeer, Apoorv Kulshreshtha, Heng-Tze Cheng, Alicia Jin, Taylor Bos, Leslie Baker, Yu Du, ... Quoc Le

arXiv (2022) https://doi.org/kmfc DOI: 10.48550/arxiv.2201.08239

5. **GPT-4 Technical Report**

OpenAl

arXiv (2023) https://doi.org/grx4cb DOI: 10.48550/arxiv.2303.08774

6. Reference-free and cost-effective automated cell type annotation with GPT-4 in single-cell RNA-seq analysis

Wenpin Hou, Zhicheng Ji

Cold Spring Harbor Laboratory (2023-04-21) https://doi.org/gsznzg

DOI: 10.1101/2023.04.16.537094 · PMID: 37131626 · PMCID: PMC10153208

7. How will generative AI disrupt data science in drug discovery?

Jean-Philippe Vert

Nature Biotechnology (2023-05-08) https://doi.org/gsznzd

DOI: <u>10.1038/s41587-023-01789-6</u> · PMID: <u>37156917</u>

8. Foundation models for generalist medical artificial intelligence

Michael Moor, Oishi Banerjee, Zahra Shakeri Hossein Abad, Harlan M Krumholz, Jure Leskovec, Eric J Topol, Pranav Rajpurkar

Nature (2023-04-12) https://doi.org/gr4td4

DOI: 10.1038/s41586-023-05881-4 · PMID: 37045921

9. Dangchain https://python.langchain.com/

10. **AutoGPT Official**

AutoGPT Official

(2023-09-18) https://autogpt.net/

11. Toward a gold standard for benchmarking gene set enrichment analysis

Ludwig Geistlinger, Gergely Csaba, Mara Santarelli, Marcel Ramos, Lucas Schiffer, Nitesh Turaga, Charity Law, Sean Davis, Vincent Carey, Martin Morgan, ... Levi Waldron

Briefings in Bioinformatics (2020-03-09) https://doi.org/ggs7tp
DOI: 10.1093/bib/bbz158 · PMID: 32026945 · PMCID: PMC7820859

12. Democratizing knowledge representation with BioCypher

Sebastian Lobentanzer, Patrick Aloy, Jan Baumbach, Balazs Bohar, Vincent J Carey, Pornpimol Charoentong, Katharina Danhauser, Tunca Doğan, Johann Dreo, Ian Dunham, ... Julio Saez-Rodriguez

Nature Biotechnology (2023-06-19) https://doi.org/gszqjr
DOI: 10.1038/s41587-023-01848-y · PMID: 37337100

13. European privacy watchdog creates ChatGPT task force

Toby Sterling

Reuters (2023-04-14) https://www.reuters.com/technology/european-data-protection-board-discussing-ai-policy-thursday-meeting-2023-04-13/

14. **Terms of use** <u>https://openai.com/policies/terms-of-use</u>

15. Why open-source generative AI models are an ethical way forward for science

Arthur Spirling

Nature (2023-04-18) https://doi.org/gsqx6v

DOI: 10.1038/d41586-023-01295-4 · PMID: 37072520

- 16. Hugging Face Hub documentation https://huggingface.co/docs/hub/index
- 17. **Open LLM Leaderboard a Hugging Face Space by HuggingFaceH4**https://huggingface.co/spaces/HuggingFaceH4/open llm leaderboard

18. HuggingGPT: Solving AI Tasks with ChatGPT and its Friends in Hugging Face

Yongliang Shen, Kaitao Song, Xu Tan, Dongsheng Li, Weiming Lu, Yueting Zhuang *arXiv* (2023) https://doi.org/gskd97

DOI: 10.48550/arxiv.2303.17580

19. From word models to executable models of signaling networks using automated assembly

Benjamin M Gyori, John A Bachman, Kartik Subramanian, Jeremy L Muhlich, Lucian Galescu, Peter K Sorger

Molecular Systems Biology (2017-11) https://doi.org/gcm498

DOI: 10.15252/msb.20177651 · PMID: 29175850 · PMCID: PMC5731347

20. Drugst.One -- A plug-and-play solution for online systems medicine and network-based drug repurposing

Andreas Maier, Michael Hartung, Mark Abovsky, Klaudia Adamowicz, Gary D Bader, Sylvie Baier, David B Blumenthal, Jing Chen, Maria L Elkjaer, Carlos Garcia-Hernandez, ... Jan Baumbach arXiv (2023) https://doi.org/gszqjw

DOI: 10.48550/arxiv.2305.15453

21. The TRUST Principles for digital repositories

Dawei Lin, Jonathan Crabtree, Ingrid Dillo, Robert R Downs, Rorie Edmunds, David Giaretta, Marisa De Giusti, Hervé L'Hours, Wim Hugo, Reyna Jenkyns, ... John Westbrook *Scientific Data* (2020-05-14) https://doi.org/ggwrtj

DOI: 10.1038/s41597-020-0486-7 · PMID: 32409645 · PMCID: PMC7224370

22. Large Language Models are Zero-Shot Reasoners

Takeshi Kojima, Shixiang Shane Gu, Machel Reid, Yutaka Matsuo, Yusuke Iwasawa *arXiv* (2022) https://doi.org/gr263v

DOI: 10.48550/arxiv.2205.11916

23. **Causality**

Judea Pearl

Cambridge University Press (2009-09-14) https://doi.org/ggd72q

DOI: <u>10.1017/cbo9780511803161</u>

24. Quantitative and logic modelling of molecular and gene networks

Nicolas Le Novère

Nature Reviews Genetics (2015-02-03) https://doi.org/f6299z

DOI: <u>10.1038/nrg3885</u> · PMID: <u>25645874</u> · PMCID: <u>PMC4604653</u>

25. Datalog Reasoning over Compressed RDF Knowledge Bases

Pan Hu, Jacopo Urbani, Boris Motik, Ian Horrocks

Proceedings of the 28th ACM International Conference on Information and Knowledge

Management (2019-11-03) https://doi.org/grfjn7

DOI: <u>10.1145/3357384.3358147</u>