Vagelos Report Summer 2017

This manuscript was automatically generated from [zietzm/Vagelos2017@570c599](#) on August 17, 2017.

Authors

- **Michael N. Zietz**
  - [zietzm](#)

  Greene Lab, Department of Systems Pharmacology and Translational Therapeutics, University of Pennsylvania

Scientific background

Drug development

Drug development times and R&D expenditures have risen considerably in the last decades. In fact, to bring a new compound to market can take a 14 years on average [1]. This development process, according to a 2016 estimate by DiMasi et al. [2], costs an average of $2.87 billion dollars, including post-approval R&D. This amount is up from two previous studies by the same authors which found the average captialized R&D costs to be $802 million in 2003 [3] and $318 million in 1991 [4]. Despite increases in drug development time and expenditures, the rate of R&D failure has increased since the 1990s [5]. Only about one in 5,000 compounds which begin preclinical testing are eventually approved [6]. Even among those which enter phase I of clinical development, only an estimated one in ten drugs will receive FDA approval [7].

Drug repurposing

Drug repurposing refers to the application of an existing therapeutic to a different disease than the one for which it was originally intended. Because candidates for drug repurposing have already been approved for other diseases, the time and cost associated with repurposing a drug are very small compared to the development of a new drug [8]. In fact, even compounds which have been deemed safe but failed in clinical development for other reasons can be candidates for drug repurposing. Several examples exist of drugs which have been repurposed, for example aspirin to treat coronary artery disease, sildenafil to treat erectile dysfunction, and gabapentin to treat postherpetic neuralgia. Most successfully repurposed drugs were discovered serendipitously and not through any systematic discovery mechanism. Our goal is to make accurate predictions of drugs which are candidates for drug repurposing. We hope to do this using heterogeneous networks of biomedical information to learn the patterns of connections between compounds and diseases.

Summer aims

This summer I worked within the Greene Lab project called 'Hetmech' [9]. Hetmech aims to improve upon and expand the existing method for predicting drug repurposing targets.

I had three primary aims for my work this summer. The details of all are covered in the Methods section.

First, I wanted to add capability within Hetmech to compute the degree-weighted path count (DWPC). Previously, the two options had been an inaccurate DWPC method and a degree-weighted walk count (DWWC), whose limitations will be covered in the Methods section.

Second, I hoped to decrease the time required to make calculations of DWPC. Towards this aim, I hoped to speed up computation by at least an order of magnitude. Sparse matrices and parallel computation were planned solutions to this problem.

Finally, it was my goal to add a multiple search capability. This involves querying a set of nodes to return a set of predictions for nodes which connect the queried nodes. For example, one should be able to search a set of symptoms and return a list of potential diseases that connect the symptoms. As with the previous problems, moving to strictly matrix computation makes this much more simple.

My secondary aim for the summer was to become familiar with and contribute to open-source, reproducible, computational science in biology. I hoped to contribute to a collaboratively written review of deep learning methods in the fields of biology and medicine [10].

Methods

Heterogeneous networks

Heterogeneous networks ('hetnets') are networks with multiple node types and edge types. In the network used this summer, titled Hetionet v1.0 (Figure 1 B), nodes represent instances of 11 biomedical entity types, and edges correspond to one of 24 edge types, or relationships between entities. 'Graph' in this context refers to the entire network of nodes and edges. We define 'metagraph' to mean a graph of the types of nodes and edges in Hetionet (Figure 1 A).
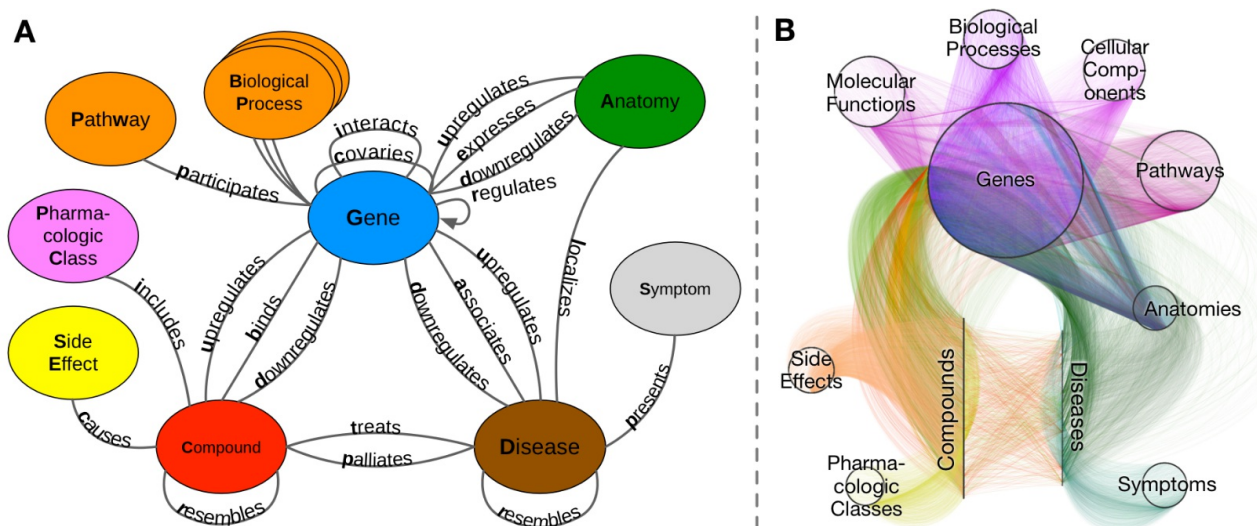
Figure 1: A. Metagraph. The graph of metandoes (node types) and metaedges (edge) type. B. Graph (Hetionet v1.0) The circles and lines represent nodes within the labeled types. For example, within the metanode 'Anatomy' we could have the node 'Leukocyte'.

Hetionet v1.0 incorporated 47,031 nodes and 2,250,197 edges [11]. A further breakdown of the nodes and edges can be found below in Tables 1 and 2, respectively.

Table 1: Breakdown of nodes by type

Metanode	Nodes
Anatomy	402
Biological Process	11,381
Cellular Component	1,391
Compound	1,552
Disease	137
Gene	20,945
Molecular Function	2,884
Pathway	1,822
Pharmacologic Class	345
Side Effect	5,734
Symptom	438

Table 2: Breakdown of edges by type and data source

Metaedge	Edges	Source

Metaedge	Edges	Source
Anatomy-downregulates-Gene	102,240	Bgee
Anatomy-expresses-Gene	526,407	Bgee and TISSUES
Anatomy-upregulates-Gene	97,848	Bgee
Compound-binds-Gene	11,571	BindingDB, DrugBank, DrugCentral
Compound-causes-Side Effect	138,944	SIDER
Compound-downregulates-Gene	21,102	LINCS L1000
Compound-palliates-Disease	390	PharmacotherapyDB
Compound-resembles-Compound	6,486	Dice coefficient = 0.5
Compound-treats-Disease	755	PharmacotherapyDB
Compound-upregulates-Gene	18,756	LINCS L1000
Disease-associates-Gene	12,623	GWAS Catalog, DISEASES, DisGeNET, DOAF
Disease-downregulates-Gene	7,623	STARGEO
Disease-localizes-Anatomy	3,602	MEDLINE
Disease-presents-Symptom	3,357	MEDLINE
Disease-resembles-Disease	543	MEDLINE
Disease-upregulates-Gene	7,731	STARGEO
Gene-covaries-Gene	61,690	Evolutionary rate covariation = 0.75
Gene-interacts-Gene	147,164	Evolutionary rate covariation = 0.75
Gene-participates-Biological Process	559,504	Gene Ontology
Gene-participates-Cellular Component	73,566	Gene Ontology
Gene-participates-Molecular Function	97,222	Gene Ontology
Gene-participates-Pathway	84,372	Gene Ontology
Gene-regulates-Gene	265,672	Gene Ontology
Pharmacologic Class-includes-Compound	1,029	DrugCentral

Graph analysis

An adjacency matrix refers to a labeled matrix with 1 or 0 at every position, corresponding to the presence or absence of a connection between two nodes [12]. The matrix is labeled, meaning that each row and column correspond to a source and target node, respectively. A function was created, titled `metaedge_to_adjacency_matrix` which performed the conversion from a string metaedge, such as 'DaG', to an adjacency matrix.
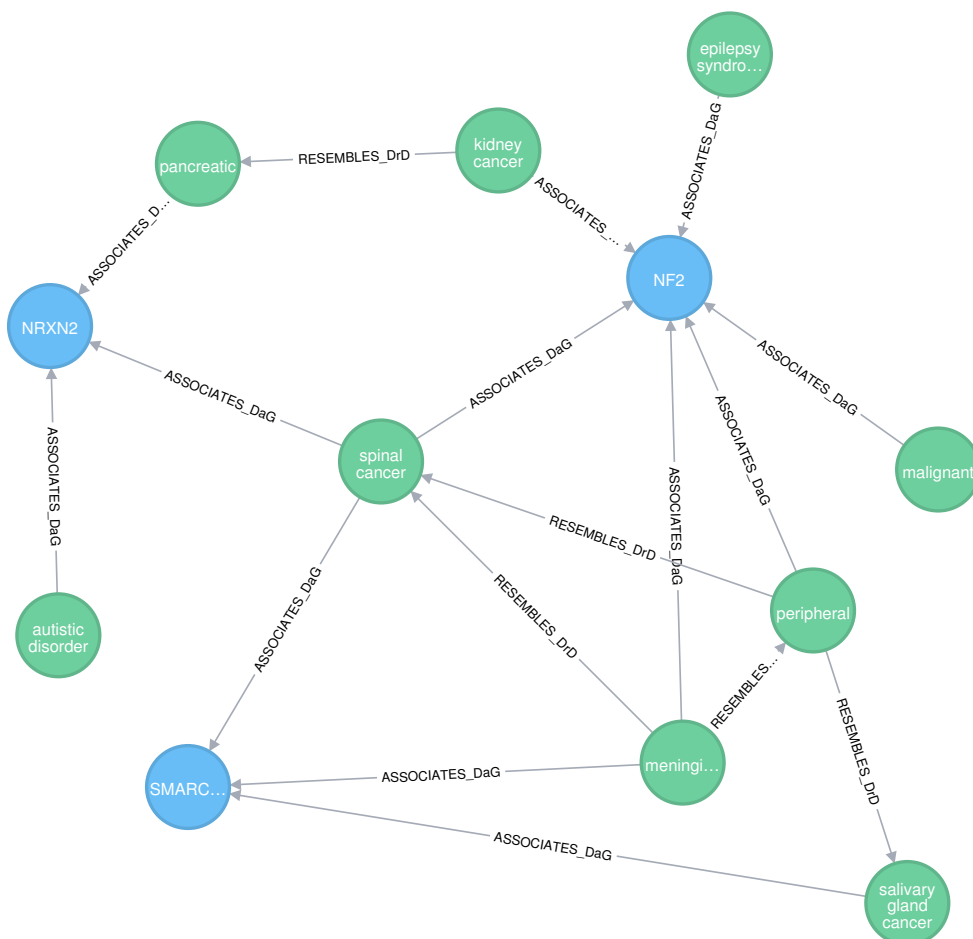

Figure 2: An example graph. Blue nodes are genes, while green nodes are diseases. The edge type between all the nodes in this graph is 'DaG' or 'Disease-associates-Gene'.

For example, the graph in Figure 2 has the following adjacency matrix corresponding to 'GaD':

$$ \begin{equation} \begin{bmatrix} 1 & 1 & 0 & 1 & 1 & 1 & 0 & 1 & 0\\ 0 & 0 & 1 & 1 & 0 & 0 & 1 & 0 & 0\\ 0 & 0 & 0 & 1 & 0 & 0 & 0 & 1 & 1 \end{bmatrix} \end{equation} $$

with labels for rows and columns, respectively:

$$\begin{bmatrix} \textrm{NF2}\\ \textrm{NRXN2}\\ \textrm{SMARCE1} \end{bmatrix}$$ $$\begin{bmatrix} \textrm{epilepsy syndrome}\\ \textrm{kidney cancer}\\ \textrm{pancreatic cancer}\\ \textrm{spinal cancer}\\ \textrm{malignant mesothelioma}\\ \textrm{peripheral nervous system neoplasm}\\ \textrm{autistic disorder}\\ \textrm{meningioma}\\ \textrm{salivary gland cancer} \end{bmatrix}$$

An adjacency matrix is identical to the information about connections between nodes along a given metaedge. Another way to consider an adjacency matrix is as a list of the nodes at which one can arrive in one step from a given start node. In this sense, performing a matrix multiplication with two adjacency matrices gives the nodes at which one can arrive in exactly *two* steps. Further, an arbitrary number of multiplications can be performed between adjacency matrices corresponding to various metaedges, so long as the dimensionality is appropriate to the matrix multiplication in question. Using this method, we can extract what is known as a walk count, or the number of ways to traverse the graph between two nodes. In this way of thinking, an adjacency matrix corresponds to the path count for paths of length one. Using the graph in Figure 2, we could perform a traversal along the meta-*path* 'GaDaG', and would obtain the following matrix:

$$\begin{bmatrix} 6 & 1 & 2\\ 1 & 3 & 1\\ 2 & 1 & 3 \end{bmatrix}$$

Notice that the elements along the main diagonal of the above matrix are not zero. This indiciates that we are accounting for walks in which we traverse from nodes as follows: A -> B -> A. Useful information cannot be gained from looping walks such as these, and they introduce considerable noise in measures of connection between nodes. We therefore wanted to eliminate any usage of walk count, and replace it with path count, where a path is a type of walk which *cannot* loop backwards on itself. In this example and for paths of length two, this is trivial; we simply subtract the main diagonal, and we have converted from a walk count to a path count. However, this conversion becomes non-trivial when dealing with longer paths and overlapping metanode repeats. Using the graph in Figure 2,
`NF2-associates-spinal cancer-resembles-peripheral nervous system neoplasm-associates-NF2`
is considered a walk, but it is not a path because its start and end nodes are the same. Note that it is perfectly acceptable to repeat *metanodes*, meaning that we can have metapaths of the form
`CrCrCrC` . Paths simply exclude the repeat of *specific nodes* within the node traverse order.

Path-counts provide useful information for predicting potential new relationships within a graph. Higher path-counts between two nodes shows good performance as a feature for predicting novel connections [13]. However, as high-degree nodes make many connections, superior performance was acheived by downweighting nodes according to their degree. To do this, row sums and column sums are taken for a matrix at each step. These one-dimensional arrays are exponentiated This represents the 'degree-weight' portion of the 'degree-weighted path count'. My work toward this will be further discussed in the Results section.

Computational tools

All computational work was done in Python version 3.6. Specifically we used an open-source scientific distribution of Python called Anaconda.

Matrices

Python's mathematical and array library, NumPy [14], has an n-dimensional array class called an
`ndarray` . `numpy.ndarray` s are very useful for representing matrix information, and have superior functionality for our purposes than the native `numpy.matrix` class. However, as can be seen in

Table 1, the number of nodes in a given adjacency matrix can be upwards of 20,000. In performing matrix multiplications, this can become a computation-intensive process that requires both significant CPU power and memory. Additionally, since the adjacency matrices are full of more zeros than ones, a majority of the multiplications performed are trivial zero multiplications.

One of my early goals for the summer was the conversion of all walk-count (and subsequently, path count) functions to sparse matrices. Sparse matrices, as employed by the Python library SciPy [15], represent data in matrices which are primarily composed of zeros. The selection of sparse representation and threshold are discussed in the Results section, but sparse matrices warrant a brief description.

In the sparse representation we used, Compressed-sparse-column format (CSC), a matrix is stored as three one-dimensional arrays.

Consider the following matrix:

$$\begin{bmatrix} 4 & 5 & 0 & 0\\ 0 & 0 & 1 & 0\\ 3 & 0 & 0 & 9\\ 0 & 6 & 0 & 5 \end{bmatrix}$$

We represent the nonzero elements with one array, with the elements being taken from top to bottom, from left to right.

$$\begin{bmatrix} 4 & 3 & 5 & 6 & 1 & 9 & 5 \end{bmatrix}$$

Next, we give the row indices of these elements.

$$\begin{bmatrix} 0 & 2 & 0 & 3 & 1 & 2 & 3 \end{bmatrix}$$

The final array represents what is called a column pointer. This array gives the indices where each column starts.

$$\begin{bmatrix} 0 & 2 & 4 & 5 & 7 \end{bmatrix}$$

These three arrays represent the entirety of a matrix.

Other tools

A significant amount of work done later in the summer involved attempting to compare the new functions to the quite slow old functions. As has become relatively standard for much of computational work, we made use of Jupyter notebooks [16] to display the code used to run analysis and its output in a re-usable way. Within this, we were able to effectively incorporate the data manipulation and analysis library pandas [17] and the multiprocessing library concurrent.futures [18]. For visualization, we utilized the online service Neo4j to represent Hetionet v1.0 [19], and the Python graphing library Matplotlib [20].

The most important tool we used this summer to track progress was the version control software, Git. Our repositories were hosted on the online git hosting service, GitHub [21]. Specifically, every contribution I made this summer can be viewed in detail at my GitHub profile [22]. This will be

discussed further in the Results section below.

Results

The main problem towards which I worked this summer was an implementation of the degree-weighted path count. While the degree-weighted walk count is relatively trivial to implement with matrix multiplication, the path count is non-trivial. If computational efficiency is to be considered, each category of metapath will require a different path-count method. As of mid-August 2017, I have merged 7 pull requests into the main Hetmech repository on GitHub, and I have one open development branch. These contributions amounted to 1043 lines added and 264 lines deleted in the repository.

DWPC

I have almost completed an implementation of the degree-weighted path count (DWPC) in the form of several independent functions. When a user calls the `dwpc` function over a metapath, a series of steps occur before any actual path-counting occurs. First, the metapath is categorized according to its repeated metanodes. For example, the metapath `GaDrDaG`, ('Gene-associates-Disease-resembles-Disease-associates-Gene') would be classified `BAAB`. Further examples of this classification method are in Table 3 below.

Next, the metapath was split into segments according to its classification. This step allowed for the abstraction of metapath patterns to metapaths which followed the pattern of a classification but included randomly inserted, metanodes anywhere in the bath. Splitting the metapath essentially allowed us to work with paths like `A-B-C-D-B-A` in the same way that we work with `A-B-B-A`, by abstracting any non-repeating metanodes to within segments.

Table 3: Example metapaths with classifications and segments

Metapath	Classification	Segments
CbGiGbC	BAAB	CbG GiG GbC
DaGaDaG	BABA	DaG GaD DaG
DlAeGaDaG	BABA	DlAeG GaD DaG
CrCrC	short_repeat	CrCrC
DlAeG	no_repeats	DlAeG
CrCrCrC	long_repeat	CrCrCrC
CbGiGiGaDrDpCpD	interior_complete_group	CbG GiGiG GaD DrD DpC CpD

Once the metapath is split into segments, the original metapath classification is used to select the appropriate dwpc function. For example, if the metapath classification is `BAAB`, then the

segmented metapath will be run in the function `dwpc_baab`.

Each DWPC function has a unique method for ensuring that it outputs a path-count rather than a walk count. For many metapaths this was a non-trivial method to unravel, and often involved several steps of additions, subtractions, multiplications, and normalizations. Our work was greatly aided by the help of a mathematician with whom we collaborated on some of the more challenging matrix operations.

In addition to the specific DWPC functions, I created a general method which will work over all metapaths, no matter the length. While slower than the other methods, this function allows us to ensure that every path is covered by the DWPC. The method uses a dictionary of history vectors for every index in the matrix and splits computations whenever a path has the opportunity to diverge into multiple potential paths.

Calculation time

Multiple search capability

As with many graph traversal tasks, using adjacency matrices to calculated DWPC meant that querying a set of nodes has become simpler. Our initial work has been on metapaths starting with compounds and ending with diseases. Having cached our DWPC matrices along every C-[…]-D metapath with five or fewer edges, we can extract path counts for queried nodes by performing matrix-vector multiplication.

Other summer results

I also gained a deeper appreciation of and respect for open-access and collaborative science. In working towards this, I reported a bug in the open source Python repository SciPy which dealt with issues involving matrix multiplication [23].

In the deep learning review, I contributed a section on deep learning applications in the field of protein-protein interaction predictions, with a subsection on deep learning methods for the prediction of MHC-peptide binding [24].

Next steps

The primary next step is to use the functionality created this summer to make test predictions of potential targets for drug repurposing.

By increasing the amount of data within the hetnet, we will expand our capability to make high-quality predictions of potential connections between nodes.

Ideas to add above

- Actually make some C–>D predictions

- Make the hetnet bigger
- Use many more metapaths than just Rephetio ones, as these aren't (to my knowledge) all the metapaths linking compounds and diseases.
- Potentially test the method's performance on other types of metapath, ie. not just C-[…]-D.
- Potentially some sort of analysis of new drugs entering into the later stages of clinical trial, and compare these (whose data shouldn't be in the hetnet) to our predictions for their targets. This would be a further validation of the methodology.
- Create an open-access webserver
- Potentially try new methods to find predictions between nodes, see if some other sort of feature can be extracted for non-existent connections or nodes. Something like [25] where a two-stage deep neural network first learned the characteristics of proteins that make them likely to interact, then secondly combined these already-higher level features into an even higher-level feature in the second stage. There could be any number of other graph-theory concepts which could be applied here, including both features and potential corrections, like graph centrality, etc.

Citations

1. Scannell JW, Blanckley A, Boldon H, Warrington B. 2012 Diagnosing the decline in pharmaceutical R&D efficiency. *Nat Rev Drug Discov* **11**, 191–200. See https://doi.org/10.1038/nrd3681.

2. DiMasi JA, Grabowski HG, Hansen RW. 2016 Innovation in the pharmaceutical industry: New estimates of R&D costs. *Journal of Health Economics* **47**, 20–33. See https://doi.org/10.1016/j.jhealeco.2016.01.012.

3. DiMasi JA, Hansen RW, Grabowski HG. 2003 The price of innovation: new estimates of drug development costs. *Journal of Health Economics* **22**, 151–185. See https://doi.org/10.1016/s0167-6296(02)00126-1.

4. DiMasi JA, Hansen RW, Grabowski HG, Lasagna L. 1991 Cost of innovation in the pharmaceutical industry. *Journal of Health Economics* **10**, 107–142. See https://doi.org/10.1016/0167-6296(91)90001-4.

5. Pammolli F, Magazzini L, Riccaboni M. 2011 The productivity crisis in pharmaceutical R&D. *Nat Rev Drug Discov* **10**, 428–438. See https://doi.org/10.1038/nrd3405.

6. Schuhmacher A, Gassmann O, Hinder M. 2016 Changing R&D models in research-based pharmaceutical companies. *J Transl Med* **14**, 105. See https://www.ncbi.nlm.nih.gov/pubmed/27118048.

7. Hay M, Thomas DW, Craighead JL, Economides C, Rosenthal J. 2014 Clinical development success rates for investigational drugs. *Nat Biotechnol* **32**, 40–51. See https://doi.org/10.1038/nbt.2786.

8. Nosengo N. 2016 Can you teach old drugs new tricks? *Nature* **534**, 314–316. See https://doi.org/10.1038/534314a.

9. greenelab. In press. greenelab/hetmech. *GitHub*. See https://github.com/greenelab/hetmech.

10. greenelab. In press. greenelab/deep-review. *GitHub*. See https://github.com/greenelab/deep-review.

11. Himmelstein DS, Baranzini SE. 2015 Heterogeneous Network Edge Prediction: A Data Integration Approach to Prioritize Disease-Associated Genes. *PLoS Comput Biol* **11**, e1004259. See https://www.ncbi.nlm.nih.gov/pubmed/26158728.

12. Weisstein EW. In press. Adjacency Matrix. *MathWorld–A Wolfram Web Resource*. See http://mathworld.wolfram.com/AdjacencyMatrix.html.

13. Himmelstein DS, Baranzini SE. 2015 Heterogeneous Network Edge Prediction: A Data Integration Approach to Prioritize Disease-Associated Genes. *PLoS Comput Biol* **11**, e1004259. See https://doi.org/10.1371/journal.pcbi.1004259.

14. numpy. In press. numpy/numpy. *GitHub*. See https://github.com/numpy/numpy.

15. scipy. In press. scipy/scipy. *GitHub*. See https://github.com/scipy/scipy.

16. jupyter. In press. jupyter/notebook. *GitHub*. See https://github.com/jupyter/notebook.

17. pandas-dev. In press. pandas-dev/pandas. *GitHub*. See https://github.com/pandas-dev/pandas.

18. 2017 17.4. concurrent.futures — Launching parallel tasks — Python 3.6.2 documentation. See https://docs.python.org/3/library/concurrent.futures.html.

19. Technology N. 2017 Hetionet · Neo4j Browser. See https://neo4j.het.io/browser/.

20. matplotlib. In press. matplotlib/matplotlib. *GitHub*. See https://github.com/matplotlib/matplotlib.

21. In press. Greene Laboratory. *GitHub*. See https://github.com/greenelab.

22. zietzm. In press. zietzm (Michael Zietz). *GitHub*. See https://github.com/zietzm.

23. scipy. In press. Operations between numpy.array and scipy.sparse matrix return inconsistent array type · Issue #7510 · scipy/scipy. *GitHub*. See https://github.com/scipy/scipy/issues/7510.

24. greenelab. In press. Added PPI section with MHC subsection by zietzm · Pull Request #638 · greenelab/deep-review. *GitHub*. See https://github.com/greenelab/deep-review/pull/638.

25. Du X, Sun S, Hu C, Yao Y, Yan Y, Zhang Y. 2017 DeepPPI: Boosting Prediction of Protein–Protein Interactions with Deep Neural Networks. *J. Chem. Inf. Model.* **57**, 1499–1510. See https://doi.org/10.1021/acs.jcim.7b00028.