



# Unraveling the Function of Protein Interactions: Insights and Future Predictions

2024 M.Sc. Biology & Biomedicine Module: Proteinbiochemie und Bioinformatik

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### 1. Why protein interactions are important?



They help us to:

- Study biological processes and molecular functions
- Understand disease mechanisms
- Develop new therapies
- Evolutionary and functional annotations

# 2. Classification of protein interactions

### a. Composition

#### Homo-oligomers:

• If interacting partners are identical

#### Hetero-oligomers:

• If interacting partners are nonidentical (Homo-oligomer) PO4792 04792

Heat Shock Protein 27



Hemoglobin (Hetero-oligomer)

### 2. Classification of protein interactions

### b. <u>Affinity</u>

#### Obligate:

• Constituents of a complex are unstable on their own in vivo

#### Non obligate:

- The components of non-obligate interactions can exist independently
- Ofter are regulated by environmental or cellular conditions





RNAse-Antibody (Non- obligate)

### 2. Classification of protein interactions

### c. Lifetime

#### Transient:

• The components of transient interaction associate and dissociate temporarily in vivo

#### Permanent:

 Permanent interactions are usually very stable and irreversible



# 3. Studying Protein-Protein Interaction Networks (PPINs)



3. How do we study protein interactions on a large scale computationally?

### Graph theory!

"[...] the study of graphs, mathematical structures used to model pairwise relations between objects. A graph in this context is made up of vertices, nodes, or points which are connected by edges, arcs, or lines".





Wikipedia

# 3. Graph Theory: types of graphs



 protein protein interaction networks

- metabolic networks
- regulatory networks

• gene co-expression networks

### 3. Graph Theory: adjacency matrices



## 3. Graph Theory: network topological properties



### **Degree Centrality**

- node property
- number of edges of a vertex (node)

### Average degree

- network property
- mean over all degrees in the network

### **Degree distribution**

- network property
- informs about the topology of the network

# 3. Graph Theory: degree distribution



### 3. Graph Theory: shortest path



#### <u>Path</u>

 between two vertices is formed by the edges that lead from one vertex to another

#### Shortest path

- shortest path between the two vertices
- used to model how information flows

The shortest path between two proteins in a PPI network often represent **the most efficient routes** for signaling

# 3. Graph Theory: betweenness and closeness centrality



# Networks in Biology



### Hyperbolic network model



### 4. How to link ontologies and PPIs

The Molecular Interactions (MI) ontology forms a structured controlled vocabulary for the annotation of experiments concerned with protein-protein interactions.



translating the code of life

Perspective | Published: 30 January 2004

### The HUPO PSI's Molecular Interaction format—a community standard for the representation of protein interaction data

Henning Hermjakob <sup>CD</sup>, Luisa Montecchi-Palazzi, Gary Bader, Jérôme Wojcik, Lukasz Salwinski, Arnaud Ceol, Susan Moore, Sandra Orchard, Ugis Sarkans, Christian von Mering, Bernd Roechert, Sylvain Poux, Eva Jung, Henning Mersch, Paul Kersey, Michael Lappe, Yixue Li, Rong Zeng, Debashis Rana, Macha Nikolski, Holger Husi, Christine Brun, K Shanker, Seth G N Grant, ... Rolf Apweiler + Show authors

Nature Biotechnology 22, 177–183 (2004) Cite this article

2776 Accesses | 449 Citations | 9 Altmetric | Metrics

#### **PSI-MI TAB Format (MITAB):**

- 1. Interactor A & B: Unique identifiers (e.g., UniProt IDs)
- 2. Interaction Type: Nature of the interaction (e.g., physical association, enxymatic activity)
- 3. Detection Method: Experimental approach used (e.g., yeast two-hybrid, co-IP)
- 4. Confidence Score: Quantifies the reliability of the interaction
- 5. Source Database: Where the data originated (e.g., IntAct, DIP)

### 4. How to link ontologies and PPIs



# 5. Databases for protein interactions



### 6. Computational predictors of PPIs



Grassmann et al. (2024), Chemical Reviews

# 7. PPINs and diseases



- Investigate disease pathogenesis
- Identification of critical nodes
- Drug discovery
- Protein networks can model how a mutation affects cellular signaling over time, offering predictions about disease onset and progression.

### 7. PPINs and diseases



### 7. PPIs and diseases

#### A survey of SCA1 RBM17 binds to pS776 and U2AF65 binds to pS776 and enhances pathogenesis regulates alternative splicing PPA2 modulates **RBM17** interaction ataxin-1 interactions **U2AF65** interaction with splicing factors PPA2-mediated dephosphorylation 14-3-3 binding retards SCA1 degeneration LANP deletion ameliorates S776 phosphorylation 14-3-3 binding increases toxicity SCA1 pathology LANP interaction AKT-mediated phosphorylation **CIC** binding increases SCA1 toxicity **CIC** interaction PolyQ SCA1 36-88 **AXH** Ataxin-1 Gif-1 interaction luclear localization K746 Gif-1 is necessary for Absence of NLS diminishes survival of Purkinje cells SCA1 phenotypes RORa interaction Partial loss of RORa enhances pathogenicity **RNF4** interaction SUMOylation **RNF4** ubiquitinates SUMOylated SUMOylation enhances ataxin-1 degradation and reduces toxicity ataxin-1 and enhances its degradation

Graphic representation of the SCA1 protein, its domains, interactions, and modifications.

> JCI Insight. 2021 Feb 8;6(3):e144955. doi: 10.1172/jci.insight.144955.

#### Modulation of ATXN1 S776 phosphorylation reveals the importance of allele-specific targeting in SCA1

Larissa Nitschke <sup>1</sup> <sup>2</sup> <sup>3</sup>, Stephanie L Coffin <sup>2</sup> <sup>3</sup> <sup>4</sup>, Eder Xhako <sup>2</sup> <sup>3</sup> <sup>4</sup>, Dany B El-Najjar <sup>2</sup> <sup>3</sup>, James P Orengo <sup>3</sup> <sup>5</sup>, Elizabeth Alcala <sup>2</sup> <sup>3</sup>, Yanwan Dai <sup>3</sup> <sup>6</sup>, Ying-Wooi Wan <sup>2</sup> <sup>3</sup>, Zhandong Liu <sup>3</sup> <sup>6</sup>, Harry T Orr <sup>7</sup>, Huda Y Zoghbi <sup>1</sup> <sup>2</sup> <sup>3</sup> <sup>4</sup> <sup>5</sup> <sup>6</sup> <sup>8</sup>

Affiliations + expand PMID: 33554954 PMCID: PMC7934855 DOI: 10.1172/jci.insight.144955

Disruption of S776 phosphorylation on the polyQ-expanded ATXN1 results in an **improvement** of SCA1 pathogenesis!

### 7. PPIs and diseases



### 7. PPIs and diseases



# Takeaways for therapeutic usage of PPINs

**PPI networks reveal disease mechanisms**: From hijacked host pathways in infections to disrupted molecular interactions in neurodegeneration.

**They guide drug discovery**: Many successful therapies were developed by targeting interactions within these networks.

**They facilitate multi-omics integration**: Combining genetics, transcriptomics, and proteomics with PPI networks provides a comprehensive disease understanding.

# Lovely to meet you all !!!!!

Thank you! Any questions?



NETWORK OUERY

 ${
m IE}\,$  » Human Integrated Protein-Protein Interaction rEference

SCREEN ANNOTATION

DOWNI OAD

INFORMATION



- Google "hippie database" and go to <u>https://cbdm-01.zdv.uni-mainz.de/~mschaefer/hippie/</u>
- 2) Click on "NETWORK QUERY" and type on the box: ATXN1
- 3) Scroll down on the website and set the output type as: show in browser-text, set the HIPPIE confidence score = 0,7 and select on the tissue filter the "brain-cerebellum"
- 4) Click on search

### QUESTIONS

- 1) How many interactor ATAXIN 1 has?
- 2) Is ATAXIN 1 interacting with CIC protein?
- 3) If yes, how many publications validate this interaction?