

Exploring ligand-transport pathways in proteins Structure, dynamics, function & dysfunction

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Journey to the protein core and back

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Outline

- Transport pathways in soluble/globular proteins
- Bioinformatics methods for pathway analysis
- Effect of mutations in transport pathways
- Roles of transport pathways in pathology and drug discovery

Cellular environment

- Proteins
- Nucleic acids
- Membranes
- Metabolites
 - lipids, peptides & sugars
- Water & ions

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Ligand size













Proteins with functional sites located in occluded cavities



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Some basic questions

- How widespread are the tunnels?
- What types of tunnels do exist?
- What are functional roles of the tunnels?

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Some basic questions

- How widespread are the tunnels?
- What types of tunnels do exist?
- What are functional roles of the tunnels?

□ Need tools/methods to detect, evaluate and design the tunnels

G Software tool that accounts for protein dynamics by analyzing tunnels

- to identify transient tunnels
- to estimate importance of tunnels



Development of software for analysis of tunnel dynamics



analysis of pathways in Voronoi diagrams

Development of software for analysis of tunnel dynamics



analysis of pathways in Voronoi diagrams



Development of software for analysis of tunnel dynamics



pathway from a cavity to the bulk solvent



Development of software for analysis of tunnel dynamics



starting from a point in the cavity



Development of software for analysis of tunnel dynamics



the shortest and widest pathway identified



Development of software for analysis of tunnel dynamics



the shortest and widest pathway identified





















identification of tunnels in each structure







merging all identified tunnels





clustering of tunnels





analysis of tunnels



analysis of tunnel dynamics
Ligand-transport pathways – spread



Ligand-transport pathways – types

What types of tunnels exist?

single tunnel connecting the active site cavity with the bulk solvent



. Candida rugosa lipase **E.C. 3.1.1.3** (PDB-ID 1CRL)

Marques et al. 2016, Understanding Enzymes - Function, Design, Engineering and Analysis, PanStanford, pp. 421-464.

Ligand-transport pathways – types

What types of tunnels exist?

- single tunnel connecting the active site cavity with the bulk solvent
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Marques et al. 2016, Understanding Enzymes - Function, Design, Engineering and Analysis, PanStanford, pp. 421-464.

Ligand-transport pathways – types

What types of tunnels exist?

- single tunnel connecting the active site cavity with the bulk solvent
- multiple tunnels connecting the active site cavity with the bulk solvent
- multiple tunnels connecting several active sites



Marques et al. 2016, Understanding Enzymes - Function, Design, Engineering and Analysis, PanStanford, pp. 421-464.

Ligand-transport pathways – dynamics



Ligand-transport pathways – gates













2. Methane monooxygenase hydroxylase E.C. 1.14.13.25; PDB-ID 1MHY, 1XVG







3. Acetylcholinesterase E.C. 3.1.1.7; PDB-ID 2XI4





Gora et al. 2013, Chem. Rev. 113: 5871–5923.

Ligand-transport pathways – gates



1. α -amylase E.C. 2.4.1.18; PDB-ID 3N98



- 2. Methane monooxygenase hydroxylase E.C. 1.14.13.25; PDB-ID 1MHY, 1XVG







4. Triosephosphate isomerase E.C. 5.3.1.1; PDB-ID 1TIM, 1TPH









5. HIV Protease E.C. 3.4.23.16; PDB-ID 1HVR, 2PC0







3. Acetylcholinesterase E.C. 3.1.1.7; PDB-ID 2XI4







6. Acylaminoacyl peptidase E.C. 3.4.19.1; PDB-ID 304G





Gora et al. 2013, Chem. Rev. 113: 5871–5923.

Ligand-transport pathways – gates



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- control access of various solvents to the active sites
- prevents dissipation of electrons by solvent

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Mutations in existing pathways

alter properties permanent pathways



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- introduce gate (permanent -> transient)



Koudelakova et al. 2013, Angew. Chem. Int. Ed Engl. 52: 1959–1963. Marques et al. 2017, J. Chem. Inf. Model. 50: 1970–1989.

Mutations in existing pathways

- alter properties permanent pathways
- introduce gate (permanent -> transient)
- remove gate (transient -> permanent)
- modulate gating frequency or amplitude



Liskova et al. 2015, ChemCatChem 7: 648–659.

Mutations in existing pathways

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Gain-of-function mutations in potential pathways

activate new functional pathways

D Potential pathways – globular proteins

leading through voids of proteins with only sub-Å dimension



D Potential pathways – globular proteins

upon gain-of-function mutations, leading to well-defined tunnel



D Potential pathways – globular proteins

• the open tunnel enables efficient transport of waters to the active site



D Potential pathways – globular proteins

the opening of the tunnel have profound functional consequences



Soluble/globular proteins



Soluble/globular proteins

Transmembrane proteins





Soluble/globular proteins

Transmembrane proteins





Soluble/globular proteins

Transmembrane proteins





D Potential pathways – transmembrane channels

leading through voids of proteins with only sub-Å dimension



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D Potential pathways – transmembrane channels

viable upon gain-of-function mutations



Potential pathways – transmembrane channels

notable functional consequences



Ligand-transport pathways – pathology

Only rarely considered when interpreting molecular bases of diseases

- tunnels in soluble proteins have been accepted as functionally important just recently
- hindered by the lack of information on the presence of transient tunnels and potential ones with high propensity for opening

Pathologies linked to ligand-transport pathways

Protein	Disease/pathology		
Dihydroorotate dehydrogenase	autoimmune and parasitic diseases, immunosuppression, cancer, inflammation		
Nitric oxide synthase	neurological diseases, inflammation, rheumatoid arthritis, immune-type diabetes, stroke, cancer, thrombosis, infection susceptibilities		
Glycogen phosphorylase	diabetes		
Leukotriene-A4 hydrolase	inflammatory diseases		
Neurolysin	nervous and endocrine systems disorders		
Plasma cholesteryl ester transfer protein CETP	atherosclerosis		
β-hydroxyacyl-acyl carrier protein dehydratase FabZ	gastric diseases		
voltage-gated Na, K, Ca channels	periodic paralyses, mixed arrhythmias, dilated cardiomyopathy, neuronal hyperexcitability,		

Ligand-transport pathways – drug discovery

U Tunnels promising targets in drug discovery

- new functional locations to target
- selective drugs due to relatively lower evolutionary conservation



Marques et al. 2016, Med. Res. Rev. 37: 1095-1139 Santos et al. 2017, Nat. Rev. Drug Discov. 16: 19-34
Ligand-transport pathways – drug discovery

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Leukotriene A4 hydrolase/aminopeptidase

Ligand-transport pathways – drug discovery

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□ Ligand-transport pathways are of functional importance not only in transmembrane proteins but also in a wide range of soluble proteins



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 - pronounced functional impact
 - in regions otherwise functionally irrelevant

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 - **D** Pathways represents interesting targets for drug discovery