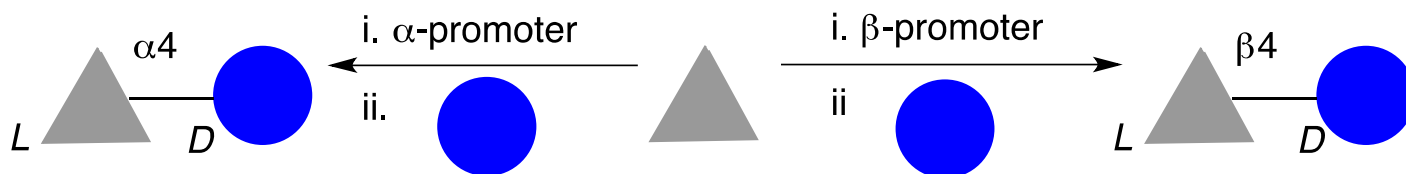
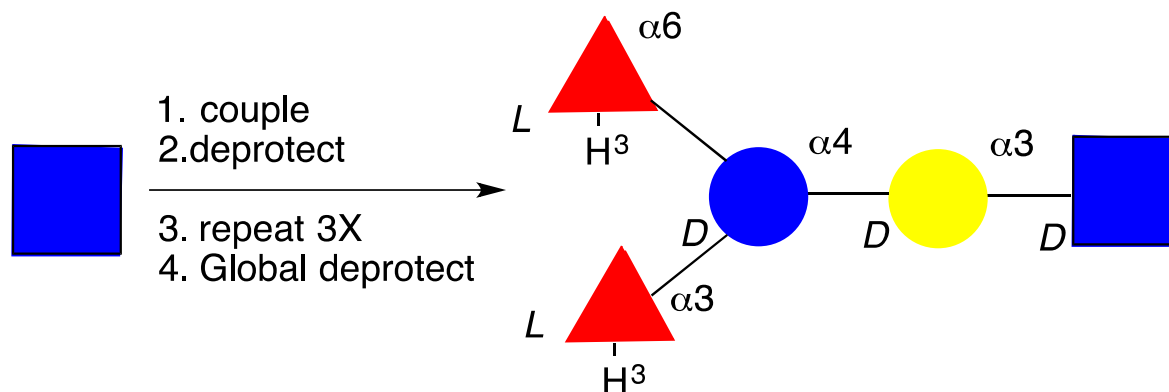


Common Fund Research in the Bennett Group: Reagent Controlled Glycosylation (U01GM120414-01)

Developing Chemical Promoters that Permit Absolute Control Over the Stereochemical Outcome of Glycosylation Reactions:



These Promoters will Make Oligosaccharide Construction Similar to Peptide Synthesis:



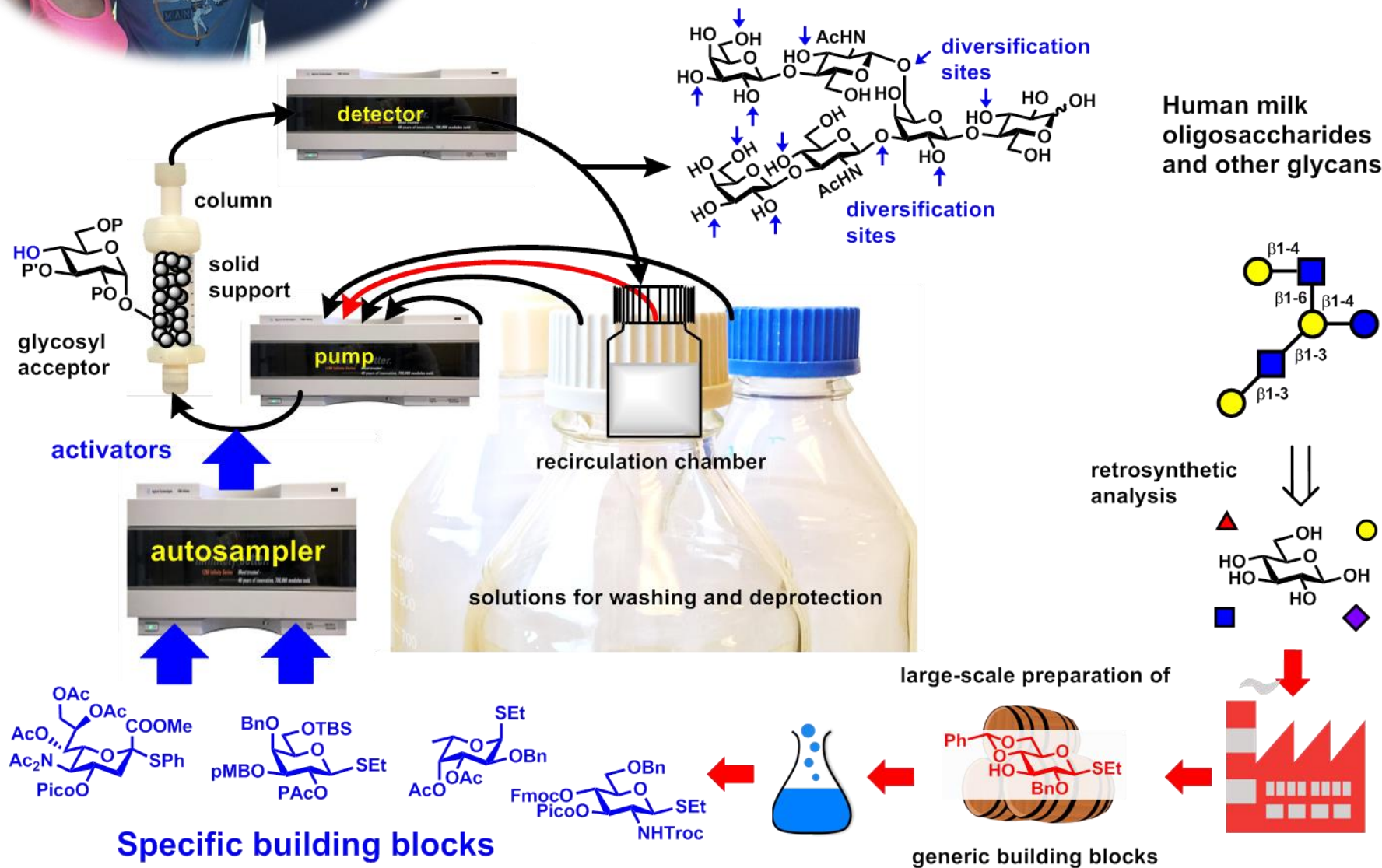
Rapid Construction of Oligosaccharides for Analytical Standards and Therapeutic Development!

<https://commonfund.nih.gov/Glycoscience/fundedresearch#>

Refinement and implementation of the automated oligosaccharide synthesizer (U01GM120673, 2016-)



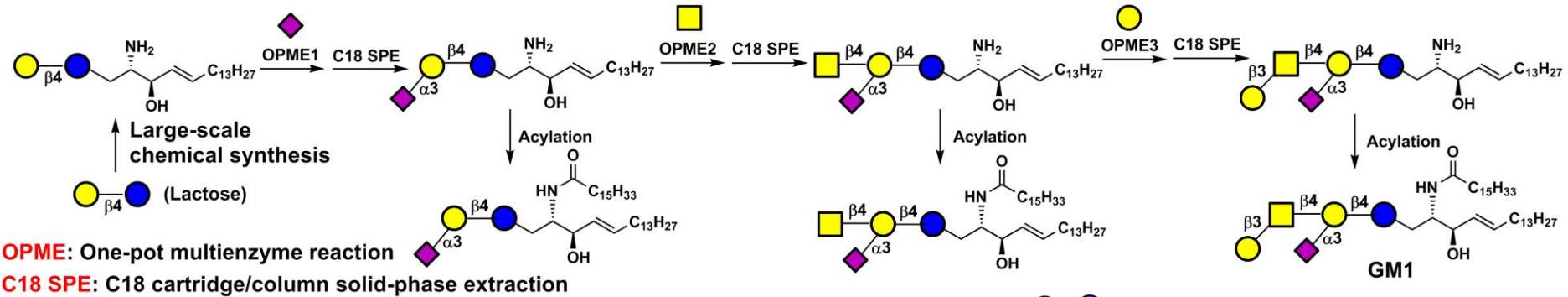
Alexei V. Demchenko, Keith J. Stine, University of Missouri - St. Louis
& Cristina De Meo, Southern Illinois University, Edwardsville



Facile chemoenzymatic synthesis and purification of glycolipids

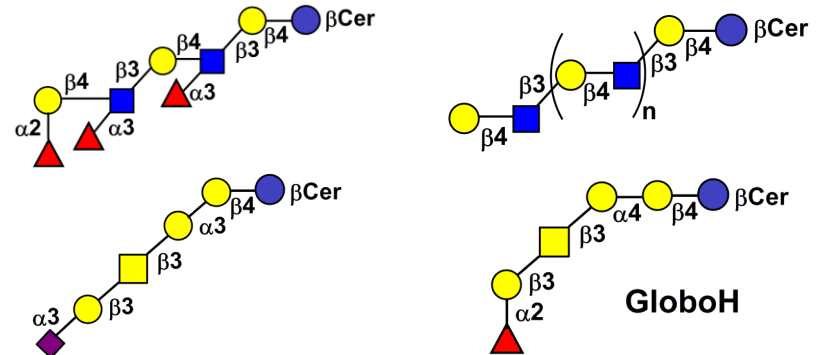
NIH Common Fund Glyco-science Program (U01GM120419)

Xi Chen, U. of California-Davis, xiichen@ucdavis.edu, <http://chengglyco.faculty.ucdavis.edu/>
Peng G. Wang, Georgia State U., pwang11@gsu.edu, <http://lithium.gsu.edu/faculty/PWang/>



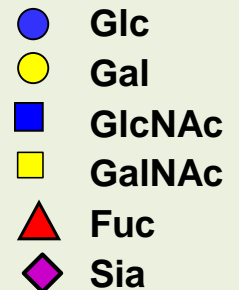
Glycosphingolipids

- ganglio-series
- (neo)lacto-series, fucosylated and sialylated
- (iso)globo-series



Goal: To allow non-specialists to synthesize, functionalize, purify, and study glycosphingolipids

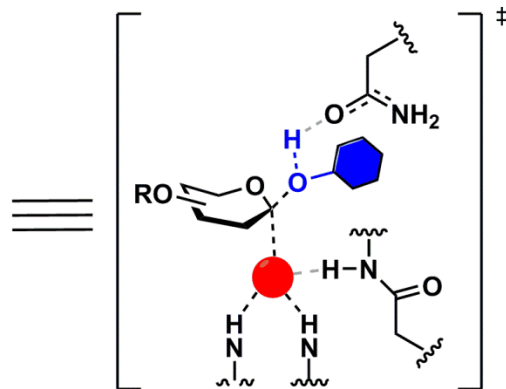
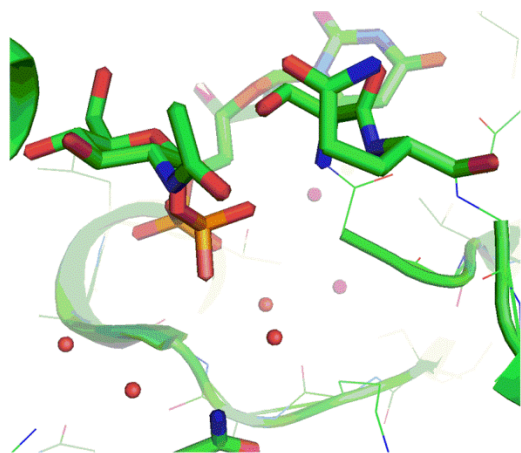
- Identify stable storage conditions for enzymes and reagents
- Assemble OPME enzyme and reagent kits
- Optimize reaction and purification conditions
- Establish protocols for OPME synthesis and C18 cartridge/column purification
- Cross-validation
- For more information, see <http://chengglyco.faculty.ucdavis.edu/glycosphingolipids/>



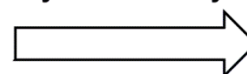
Jacobsen Group

Developing Catalysts for Selective Glycosylation

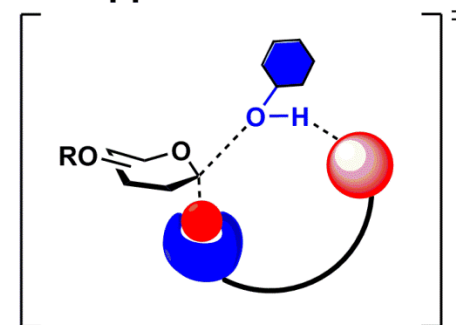
Nature's solution:



Transferase-inspired
synthetic catalysts



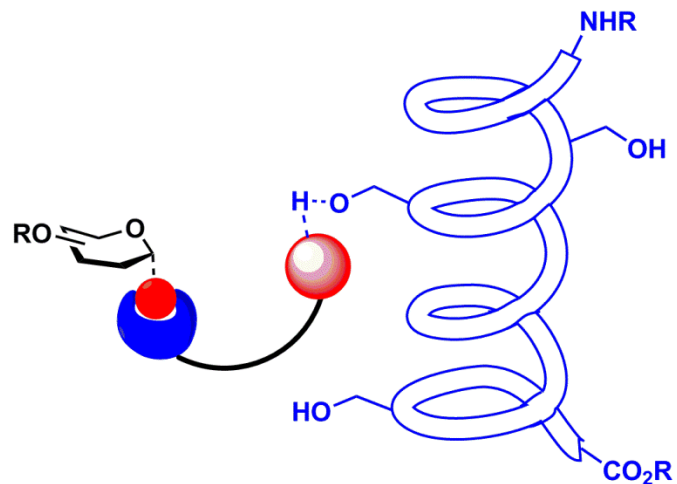
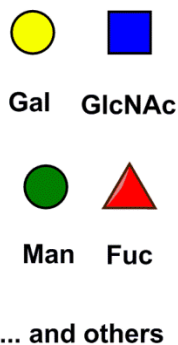
Our approach:



- Mild reaction conditions through dual-activation strategy
- Catalyst-controlled selectivity

New strategy enables:

• Broad glycosyl donor scope



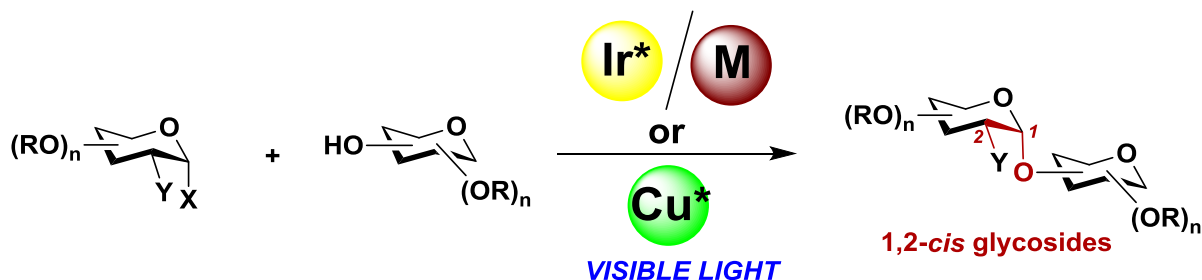
• Catalyst control of multiple reactive sites

• Broad functional group compatibility
-Amino acids
-Glycosyl acceptors

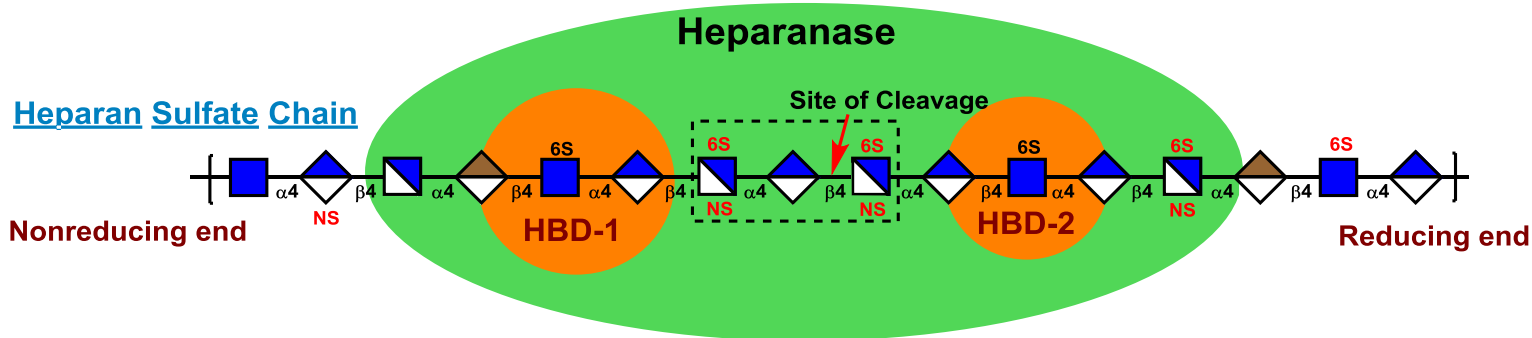
• Conjugates selectively to:
-Amines
-Thiols
-Alcohols

NIH Common Fund Research in the Nguyen Group: Stereoselective 1,2-Cis Glycosylation

Developing predictable and stereoselective 1,2-cis glycosylation reactions via either dual catalytic photoredox catalysis or photoinduced copper catalysis



Rapid and stereoselective synthesis of bioactive oligosaccharides for analytical standards and therapeutic applications



NIH-U01 GM120293

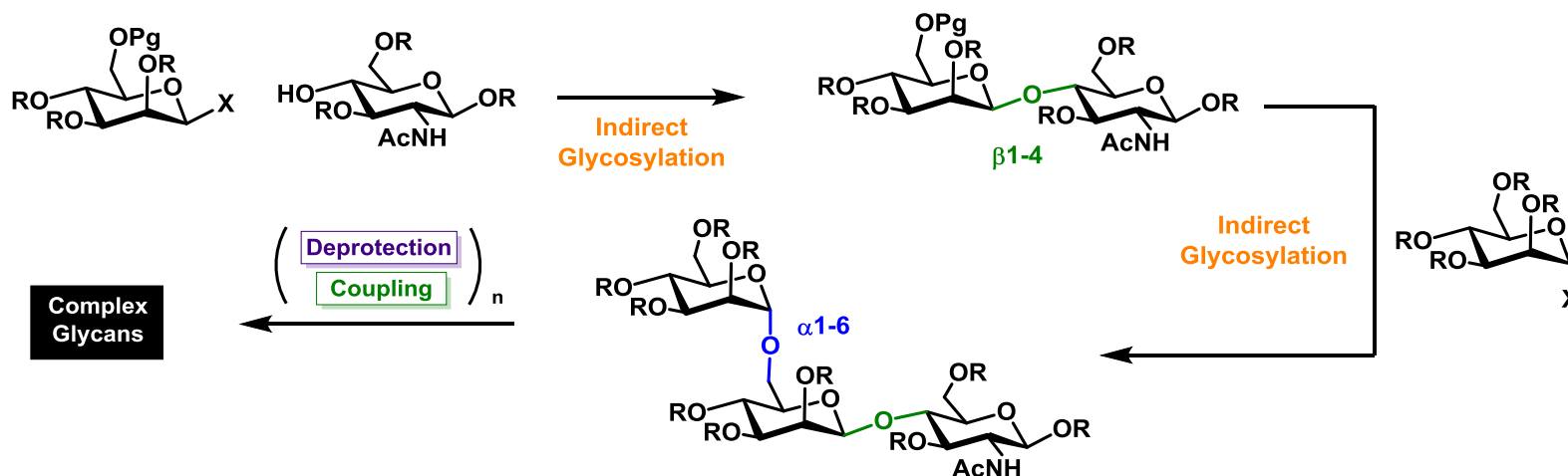
<https://commonfund.nih.gov/Glycoscience>

Hien M. Nguyen (hien-nguyen@uiowa.edu)
University of Iowa

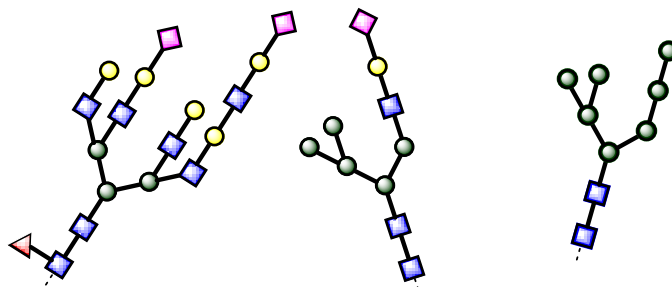
<https://nguyenresearchgroup.lab.uiowa.edu/>

NIH Common Fund Research in the Brichacek Group: Novel Glycosylation Mechanisms

Indirect Glycosylation Methods to Facilitate More Efficient and Selective Couplings

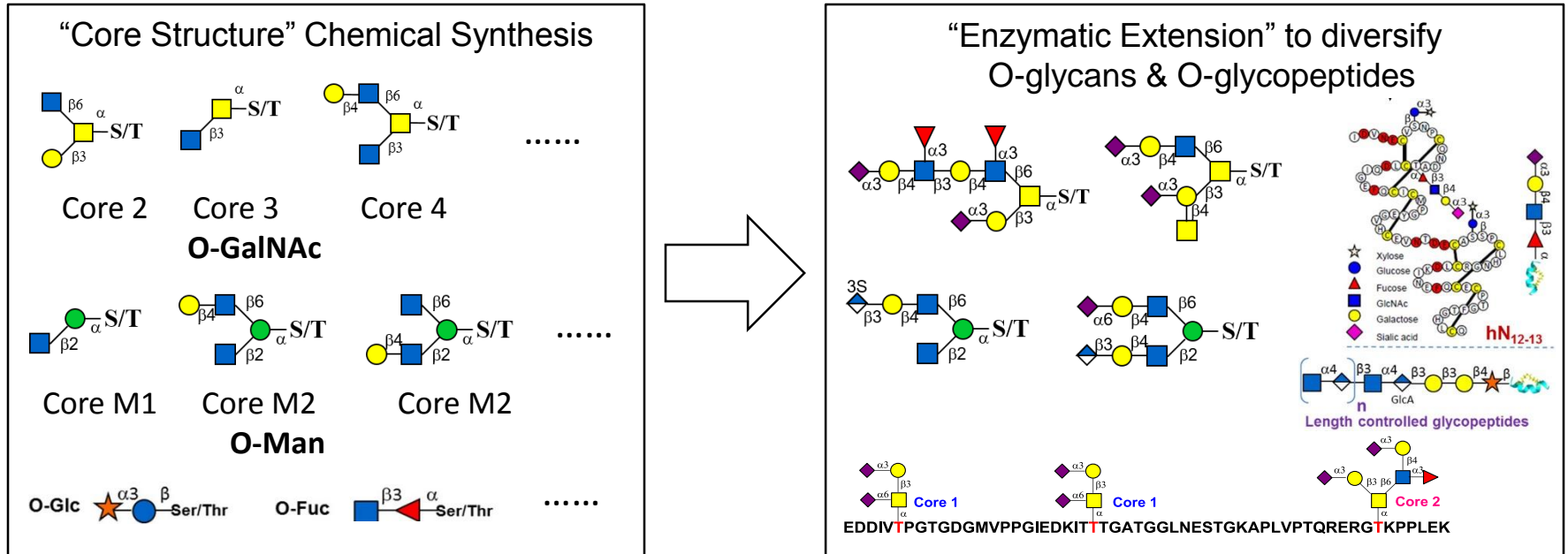


Enable access of oligosaccharides of defined sequence, branching, and stereochemistry on demand to a diverse range of biomedical researchers.



Facile Synthesis of O-glycans & O-glycopeptides

NIH Common Fund Glyco-science Program (U01GM116263)
Peng George Wang & Lei Li, Georgia State University

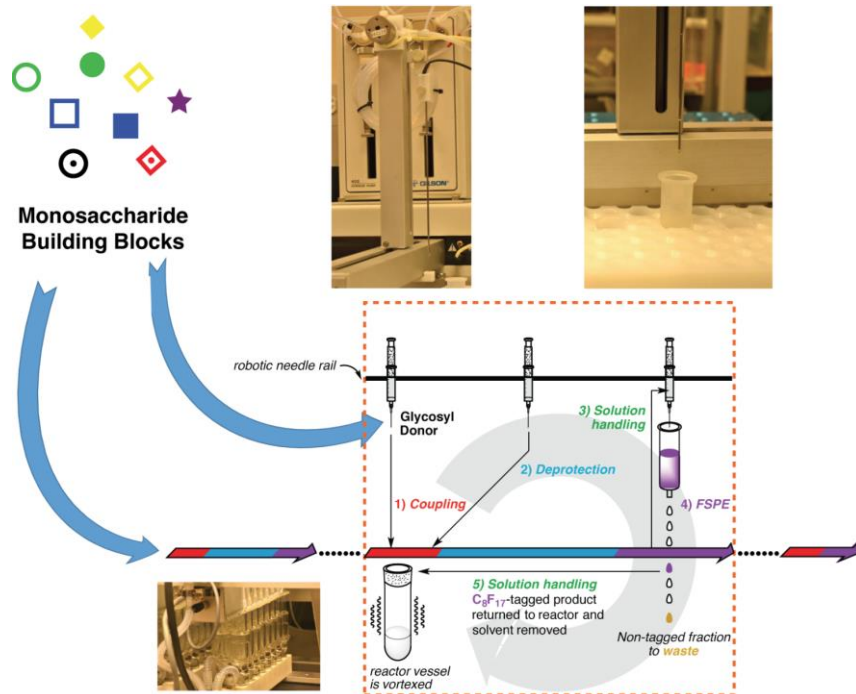


Goal: Develop “Core Synthesis/Enzymatic Extension” strategy for the access of O-glycan and O-glycopeptide libraries, automated glycopeptide synthesis

- Convergent chemical synthesis of O-glycan core structures in gram scale;
- Enzymatic extension strategy allows diversity of core structures;
- Automatic glycopeptide synthesis on solid phase/water soluble supports;
- Synthesis of hundreds of O-glycans and O-glycopeptides;
- Cross-validation

Common Fund Research in the Pohl/Dong Groups: Sugar Building Blocks and Automated Synthesis of Biomedically-Relevant Glycans

*Developing Chemical Methods to Access Building Blocks and Create Oligosaccharides
Using Solution-Phase Automation Platforms*

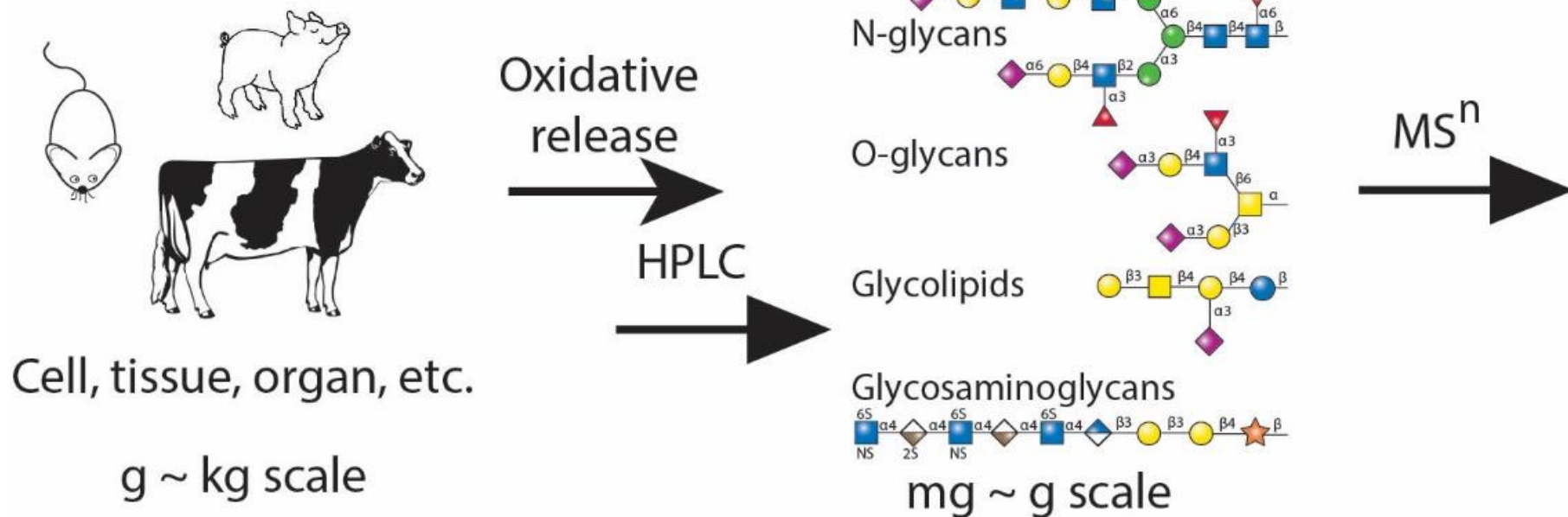


*Human Milk Oligosaccharides,
Bacterial Rhamnans, and
Mammalian O- and N-Glycans*

- *Analytical standards and compounds for bioassays with the potential to incorporate fluorescent and other labels*
- *New methods to purify synthetic glycans to 99.5%+ purity for immunological studies (Chem Commun. 2016, 52, 13253)*

<https://commonfund.nih.gov/Glycoscience/fundedresearch#>

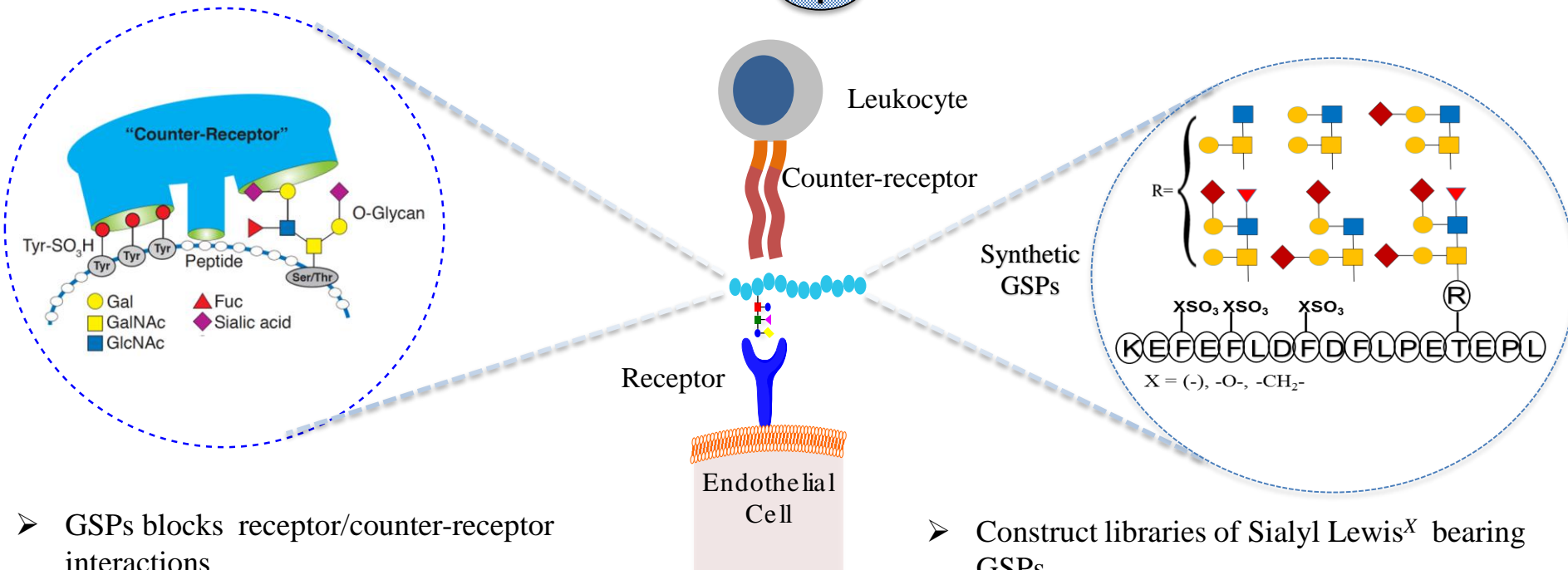
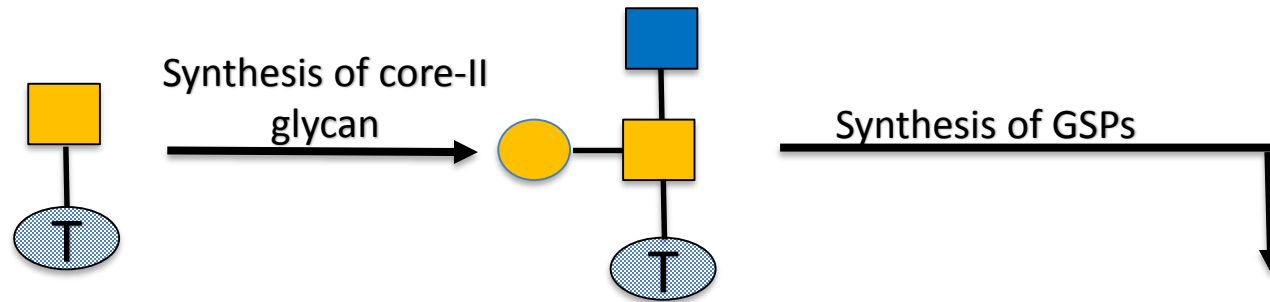
Large scale chemical preparation of glycans from natural sources



- Novel methods for large scale chemical release and multi-dimensional HPLC separation of natural glycans *Xuezheng Song, Emory University*
- Detailed structure characterization/confirmation of natural glycans *Vernon Reinhold, University of New Hampshire Glycomics Center*

Chaikof Group

Facile Synthesis of Glycosulfopeptides (GSPs) and Related Bioconjugates

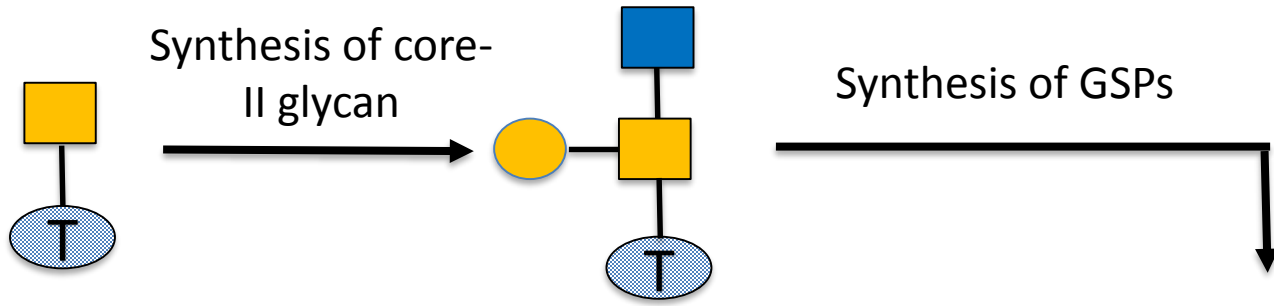


➤ GSPs blocks receptor/counter-receptor interactions

- Construct libraries of Sialyl Lewis^X bearing GSPs
- Total synthesis Of GSnP-6 and other glycosulfopeptides

Chaikof Group

Facile Synthesis of Glycosulfopeptides (GSPs) and Related Bioconjugates



Human PSGL-1 -K-E-F-Y-E-Y-L-D-Y⁴⁶-D-F-L-P-P-E-T-E-P-P-E-
 Murine PSGL-1 -G-E-D-P-D-Y⁴⁶-T-Y-N-T-D-P-P-E-
 Human GPIba -D-T-D-L-Y²³⁶-D-Y-Y-P-E-E-D-T-E-G-D-K-V-R-A-T-R-T-V-
 Murine GPIba -D-T-D-Y²⁸⁹-D-D-Y-D-D-I-P-D-V-P-A-T-R-T-E-V-K-F-S-T-N-T-
 Human Endoglycan -Q-P-P-Q⁹⁷-F-W-E-E-E-E-L¹⁷⁴-L-D-L-G-P-T-A-D-Y¹¹⁸-V-F-P-D-L-T-E-
 Murine Endoglycan -L-Q-P-P-Q⁹⁹-F-W-E-E-E-E-L¹⁷⁴-L-D-L-G-P-T-A-D-Y¹¹⁸-V-F-P-D-L-T-E-
 Endoglycan -V-P-S¹⁸⁴-V-T-P-S-T-V-A-P-G-V-Q¹⁸⁴-N-Y-S¹⁸⁴-Q-E-S-G-G-T-E-W-P-T-G-G-
 Human C3aR -N-N-R-C-G¹⁷⁴-K-F-G-L-S-S-L-D-Y-P-D-F-Y-G-D-P-L-E-
 -Q-G-F-Q-D-Y³⁷⁷-N-L-G-Q-F-
 Human CCR5 -M-D-Y³⁷⁷-Q-V-S-S-P-I-Y-D-I-N-Y-Y-T-S-E-P-C-
 Human Factor VIII -K-N-T-G-D⁷³⁷-Y-E-D-S-Y⁷⁴²-E-D-I-S-A-Y-L-L-S-K-N-N-A-I-E-P-R-S-F-S-Q-
 -N-S-R-H-P-S⁷⁶⁵-T-R-Q-K-Q-F⁷⁶⁵-N-A-T-I-P-

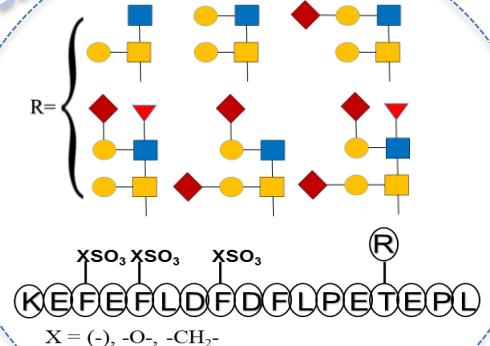
Y -Potential Tyr-Sulfation Site
 S,T -Potential O-glycosylation Site
 □ -Potential N-glycosylation Site

Natural GSPs

Counter-receptor

Synthetic GSPs

Receptor



- GSPs blocks receptor/counter-receptor interactions

- Construct libraries of sialyl Lewis^X bearing GSPs
- Total synthesis of GS_nP-6 and other glycosulfopeptides

Develop Catalytic Methods to Streamline the Assembly of Oligosaccharides

Weiping Tang, University of Wisconsin – Madison; Peng Liu, University of Pittsburgh

This program focuses on two essential issues in carbohydrate chemical synthesis.

- 1) Streamline the synthesis of carbohydrate building blocks (BBs) by site-selective functionalization of OH groups.
- 2) Streamline the assembly of carbohydrate building blocks with minimal protecting groups via stereoselective glycosylation.

