# **Subsystem: Ammonia Assimilation**

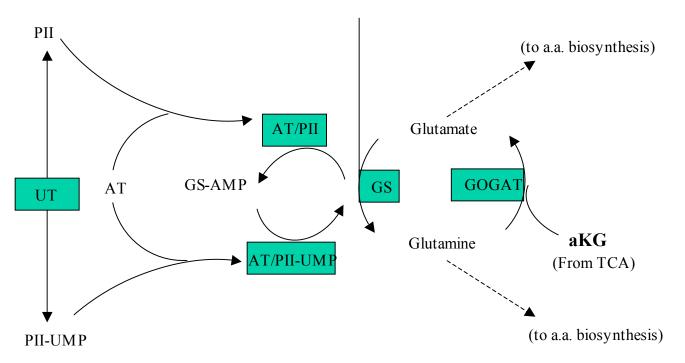
Ed Frank, Argonne National Laboratory, Argonne, IL

## Introduction

Abbrev	Functional Role
GS1	Glutamine synthetase (EC 6.3.1.2)
AT	Glutamate-ammonia-ligase adenylyltransferase (EC 2.7.7.42)
PII	Nitrogen regulatory protein P-II
PIIK	Nitrogen regulatory protein P-II, glnK
UT	[Protein-PII] uridylyltransferase (EC 2.7.7.59)
GOGDP1	Glutamate synthase [NADPH] large chain (EC 1.4.1.13)
GOGDP2	Glutamate synthase [NADPH] small chain (EC 1.4.1.13)
GOGATF	Ferredoxin-dependent glutamate synthase (EC 1.4.7.1)
GOGD	Glutamate synthase [NADH] (EC 1.4.1.14)
NRI	Nitrogen regulation protein NR(I)
NRII	Nitrogen regulation protein NR(II) (EC 2.7.3)
NtcA	Global nitrogen regulatory protein, CRP family of transcriptional regulators
IF7	Glutamine synthetase inactivating factor IF7
IF17	Glutamine synthetase inactivating factor IF17

- Ammonia, nitrite and nitrate assimilation supply nitrogen for biosynthesis.
  - Ammonia assimilation can occur via glutamate dehydrogenase (GDH) or GS-GOGAT pathways. The latter is shown here. When both capabilities exist in an organism, GS-GOGAT route prevails in low ammonia conditions.
  - The subsystem is particularly interesting because of the regulatory aspects involved: posttranslational modification of several enzymes and a signaling system.
- The system holds little functional clustering compared to other subsystems. This and the difficulty of identifying signaling genes made this an interesting test case: Can a clear classification of the variations in the GS, GOGAT, and AT subsets be used to help identify or constrain the possible signaling and regulatory variations?

### Ammonia Assimilation: Subsystem diagram



NH3

• In some organisms, Glutamine synthetase is partly regulated by adenylylation. The effector, Adenylyltransferase is itself regulated by the regulatory protein, PII: binding PII vs. PII-UMP determines whether de-adenylylation or adenylylation occurs. Interconversion of PII and PII-UMP is accomplished by Uridylyltransferase, UT. This is the story in some bacteria (e.g. E. coli) but not all. In cyanobacteria the control is not by adenylylation but perhaps by phosphorylation of a related compound and/or the action of IF7 and IF17.

## Ammonia Assimilation Spreadsheet : Examples of variations

Organism	GS1	AT	PII	PIIK	UT	GOGDP1	GOGDP2	GOGATF	GOGD	NRI	NRII	IF7	IF17
Anabaena variabilis ATCC			<u>3138</u>					<u>4866</u>				<u>3127</u>	
29413													
Synechococcus elongatus	<u>1461</u>							<u>237</u>				<u>247</u>	
PCC 7942													
Synechococcus sp. WH	<u>1068</u>							<u>2125</u>					
8102													
Trichodesmium erythraeum	<u>5780</u>		<u>4778</u>					<u>4553</u>					
IMS101													
Crocosphaera watsonii WH	<u>3868</u>		<u>3936</u>					<u>4121</u>					
8501													
Gloeobacter violaceus PCC	<u>1052</u>		<u>256,</u>					<u>1508</u>					
7421			<u>689</u>										
Prochlorococcus marinus	<u>1377</u>		<u>709</u>					<u>659</u>					
MED4													
Prochlorococcus marinus	<u>601</u>		<u>1475</u>					<u>1771</u>					
str. MIT 9313													
Nostoc sp. PCC7120	<u>2635</u>		<u>2626</u>					<u>4651</u>				<u>2636</u>	
Synechocystis sp. PCC	<u>1933,4</u>		<u>1956</u>			<u>439</u>	<u>198</u>	<u>2982</u>				<u>571</u>	<u>415</u>
6803	<u>76</u>												
Escherichia coli 0157:H7	<u>4803</u>	<u>3936</u>	<u>3417</u>		<u>169</u>	<u>4094</u>	<u>4095</u>			<u>4801</u>	<u>4802</u>		
EDL933													
Escherichia coli CFT073	<u>4717</u>	<u>3715</u>	<u>3000</u>	<u>546</u>	<u>196</u>	<u>3882</u>	<u>3382, 3883</u>			<u>4715</u>	<u>4716</u>		
Escherichia coli K12	<u>3790</u>	<u>3001</u>	<u>2523</u>	<u>447</u>	<u>167</u>	<u>3155</u>	<u>3156</u>			<u>3788</u>	<u>3789</u>		

## **Open problems, comments, conjectures**

- Protein PII (glnB) has a variant (glnK). Clustal alignment yields a tree that can be painted with data from organisms discussed in (Arcondeguy, et al., 2001). The result (a calibrated tree) appears useful for making assignments in other organisms.
- Ferredoxin-dependent, NADH-dependent, and NADPH-dependent GOGAT can be distinguished by length and functional clustering: the NADPH-dependent occurs as a pair while Fd-dependent appears singly and equal in size to the large subunit of the NADPH-dependent. NADH-dependent appears singly but equal in length to the sum of the two chains of NADPH-dependent.
- The NADPH-dependent large and small subunits are functionally clustered to the NRI and NRII. Is this true in all cases when NRI and NRII are present or only in the organisms studied so far? Projection of this cluster across organisms reveals a number of possible incorrectly called genes.
- Are the NADPH-dependent assignments in Synechocystis correct?
- Why are so few IF7 and IF17 found in the cyanobacteria?

#### **References:**

- 1. Merrick MJ, Edwards RA. Nitrogen Control in Bacteria. Microbiological Reviews 59, 604-22 (1995)
- 2. Arcondeguy T, Jack R, Merrick M. PII Signal Transduction Proteins, Pivotal Players in Microbial Nitrogen Control. Microbiology and Molecular Biology Reviews **65**, 80-105 (2001)
- 3. Herroro A, Muro-Pastor AM, Flores E. Nitrogen Control in Cyanobacteria. J. of Bacteriaology. **183**, 411-25.
- 4. Rhodes D. http://www.hort.purdue.edu/rhodcv/hort640c/INDEX.HTM