# Bioorthogonal non-canonical amino acid tagging - BONCAT -

Hatzenpichler lab www.environmental-microbiology.com doi: 10.13140/RG.2.1.3698.7040/1

# **First, some definitions**

## bioorthogonal

non-interacting with cellular functionalities

### non-canonical

synthetic, not part of biological machinery

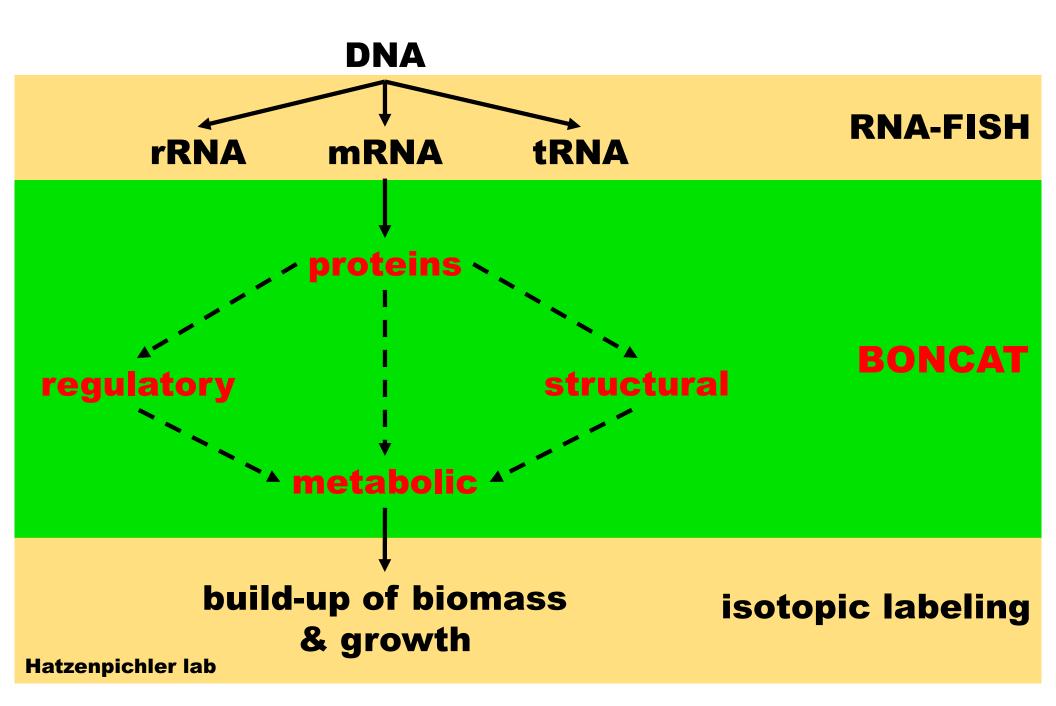
## **Click chemistry**

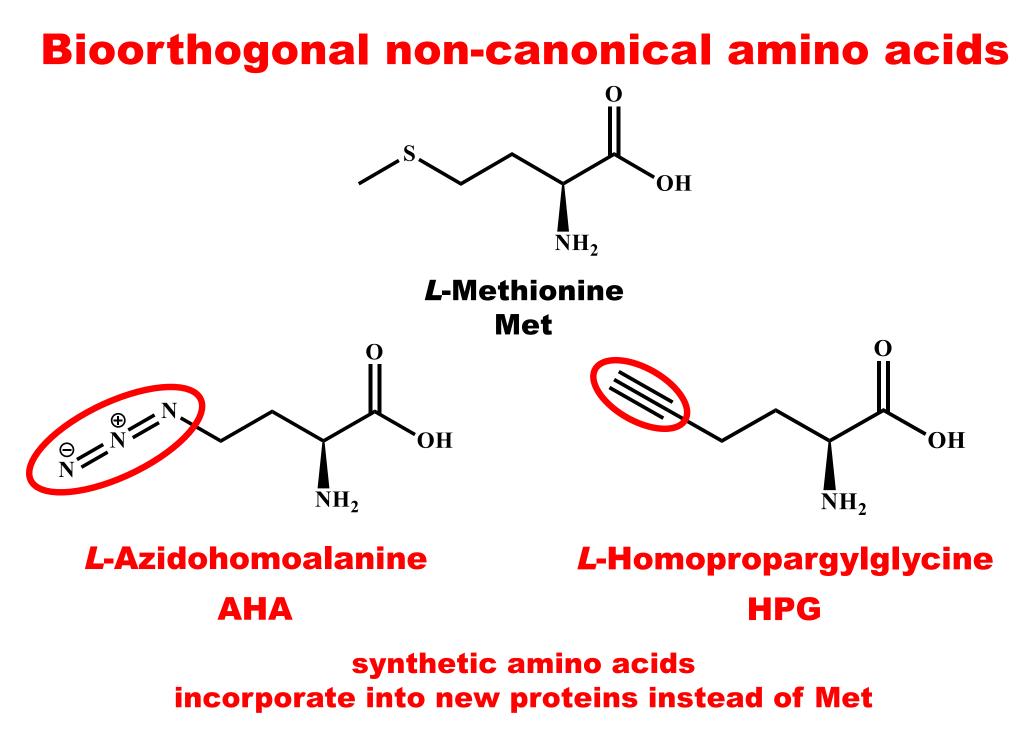
## complete conversion of reagents to single product

## + mild conditions

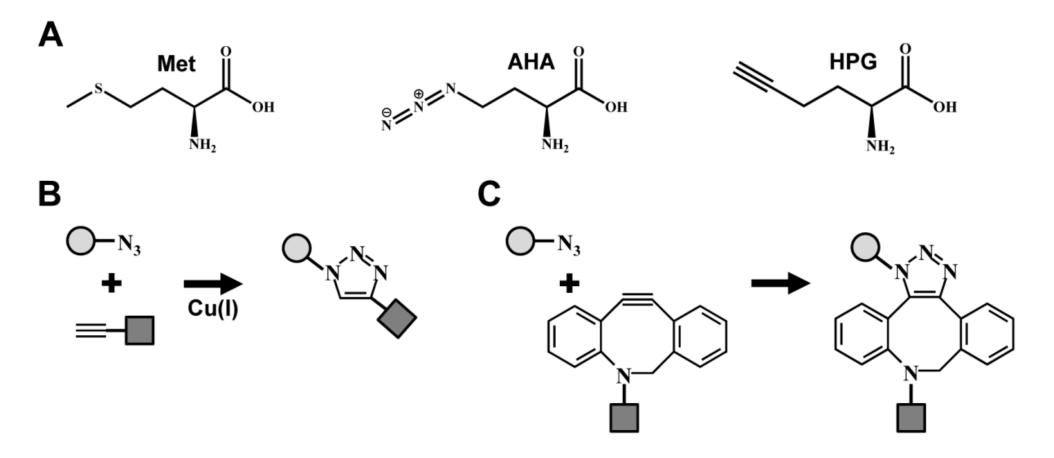
- + very fast
- + in water

# Activity assays on individual cell level





# **Azide-alkyne click reactions**



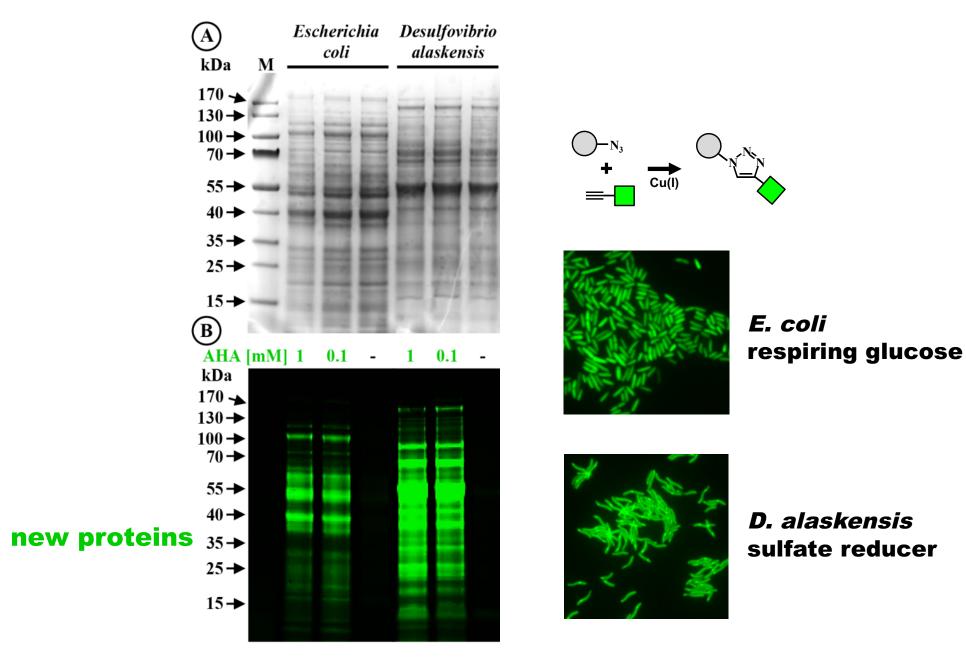
A. Structures of Met and its surrogates AHA and HPG, which compete with Met during translation.

**B.** In Cu(I)-catalyzed click chemistry an azide group ( $N_3$ ) is linked to a terminal alkyne residue, yielding a triazole conjugate.

C. Strain-promoted click chemistry allows the copper-less conjugation of an azide group ( $N_3$ ) with a cyclo-octyne-carrying molecule, yielding a triazole conjugate.

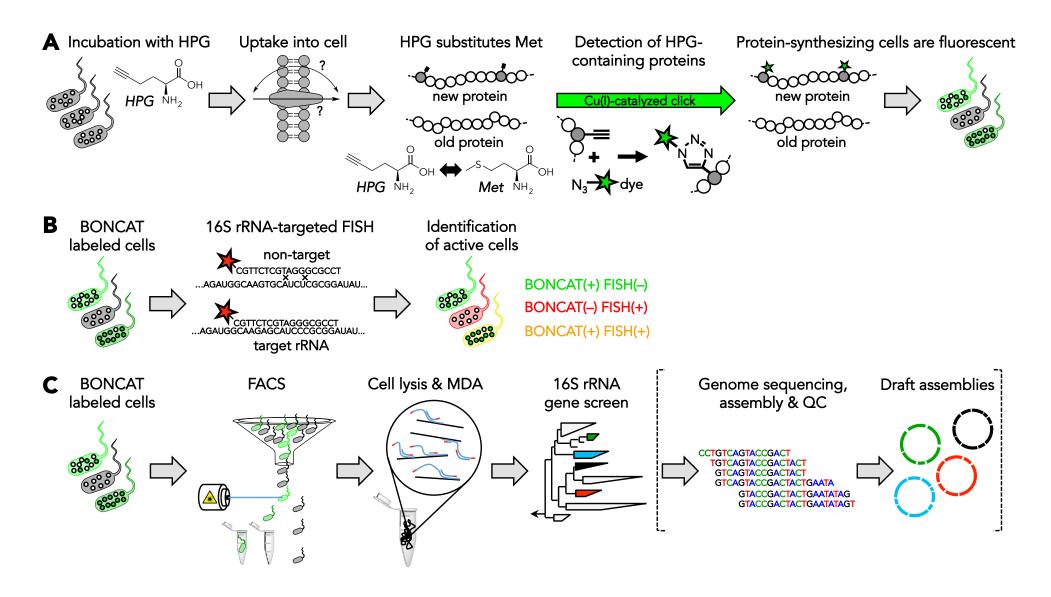
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# **Incorporation into newly made proteins**

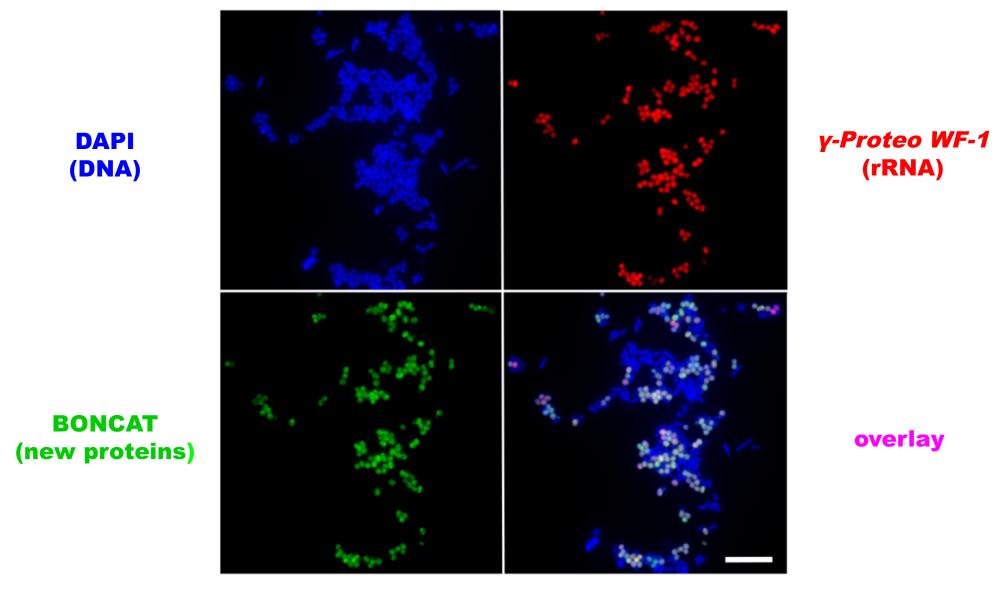


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# Visualizing, identifying, and sorting translationally active microbes



## **Identification of translationally active cells**



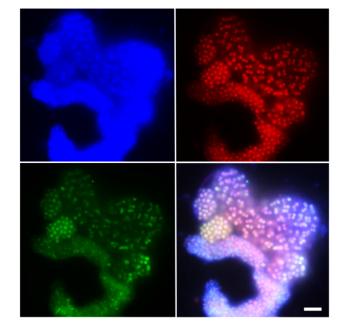
Bar = 10 µm

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# **BONCAT-FISH of uncultured microbes**

**EUB338 I-III** 

**Gam42a + competitor** 



Methane seep ANME-SRB consortium

Tongue biofilm and saliva

Freshwater from Lily pond on Caltech campus



Hatzenpichler et al., 2014; Hatzenpichler et al., 2015; Hatzenpichler et al., 2016

# Visualizing new proteins in situ

generally applicable (works for all taxonomies and physiologies tested so far)

detectable after 2% of generation time

**FISH-BONCAT** links function and identity of a cell

**BONCAT** correlates with <sup>15</sup>NH<sub>3</sub> incorporation (nanoSIMS)

no change in protein expression (Bagert *et al*., 2014)

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## **Limitations and advantages of BONCAT-FISH**

uptake and incorporation

**Methionine-rich samples are tough** 

hard to quantitate amount of new proteins in uncultured cells

potential for cell inactivation or community shifts

links cellular identity and function

fluorescence-based in situ activity studies

metabolic screening

activity-based cell-sorting

fast + highly selective + cheap + easily available1 hazide-alkyne~\$500epi-scope

## **BONCAT in environmental microbiology** (as of August 2017)

Hatzenpichler R, Connon SA, Goudeau D, Malmstrom RR, Woyke T, Orphan VJ Visualizing *in situ* translational activity for identifying and sorting slow-growing archaeal-bacterial consortia Proc Natl Acad Sci USA, 113: E4069-E4078 (2016)

application of BONCAT-FISH and BONCAT-FACS to ANME-SRB consortia from three methane seep sediments; development of activity-based cell-sorting via bioorthogonal labeling

Hatzenpichler R and Orphan VJ

Detection of protein-synthesizing microorganisms in the environment via bioorthogonal non-canonical amino acid tagging (BONCAT)

Book chapter for Hydrocarbon and Lipid Microbiology Protocols, Vol. 7: Single-cell and single-molecule methods

Springer Protocols Handbooks, doi 10.1007/8623\_2015\_61 (2015)

> description of how to design and perform BONCAT-experiments using AHA and HPG

Samo TJ, Smriga S, Malfatti F, Sherwood B, and Azam F

Broad distribution and high proportion of protein synthesis active marine bacteria revealed by click chemistry at the single cell level

Front Mar Sci, doi: 10.3389/fmars.2014.00048 (2014)

> application of BONCAT to seawater; correlation of BONCAT with MAR

Hatzenpichler R, Scheller S, Tavormina PL, Babin B, Tirrell D, and Orphan VJ

*In situ* visualization of newly synthesized proteins in environmental microbes using amino acid tagging and click chemistry

Environ Microbiol, 16: 2568-2590 (2014)

first application of BONCAT to uncultured microbes in the environment; development of BONCAT-FISH; correlation of BONCAT with nanoSIMS