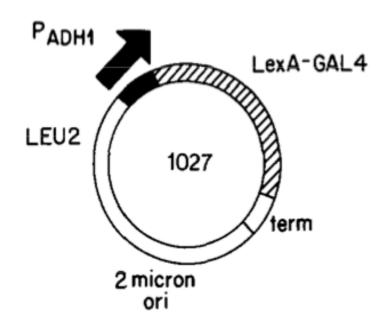
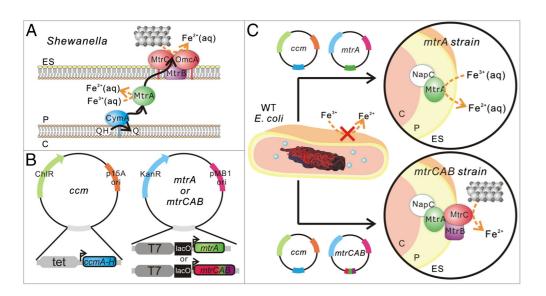
Synthetic Biology Open Language Visual An open-source graphical standard for synthetic biology

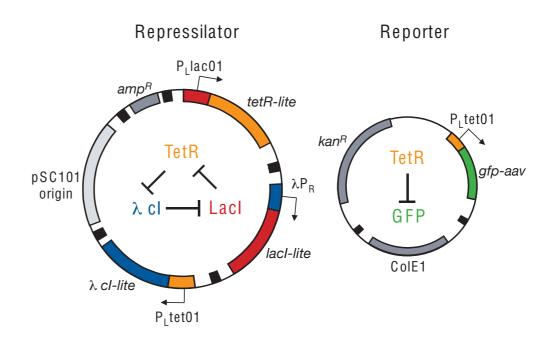
Jacqueline Quinn, Jacob Beal, Swapnil Bhatia, Patrick Cai, Joanna Chen, Kevin Clancy, Robert Sidney Cox III, Michal Galdzicki, Nathan Hillson, Akshay Maheshwari, Chris Myers, Umesh P, Matthew Pocock, Cesar Rodriguez, Herbert M Sauro, Larisa Soldatova, Guy-Bart Stan, Mandy Wilson and Drew Endy

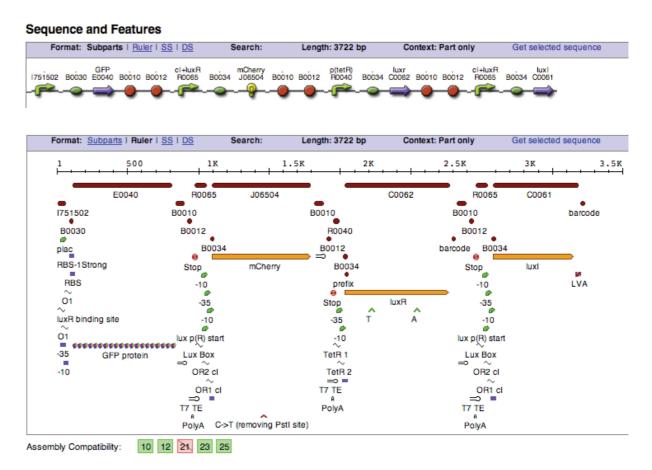
The Fifth International Workshop on Bio-Design Automation July 12, 2013

sbol visual (the basics)





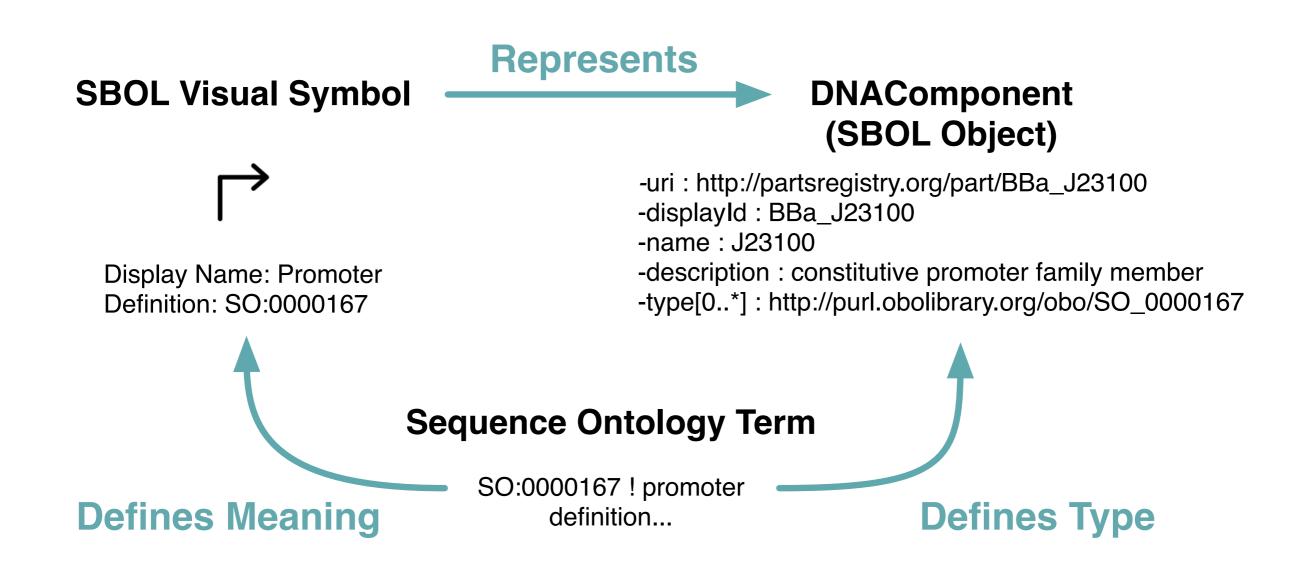




visual representation of genetic design

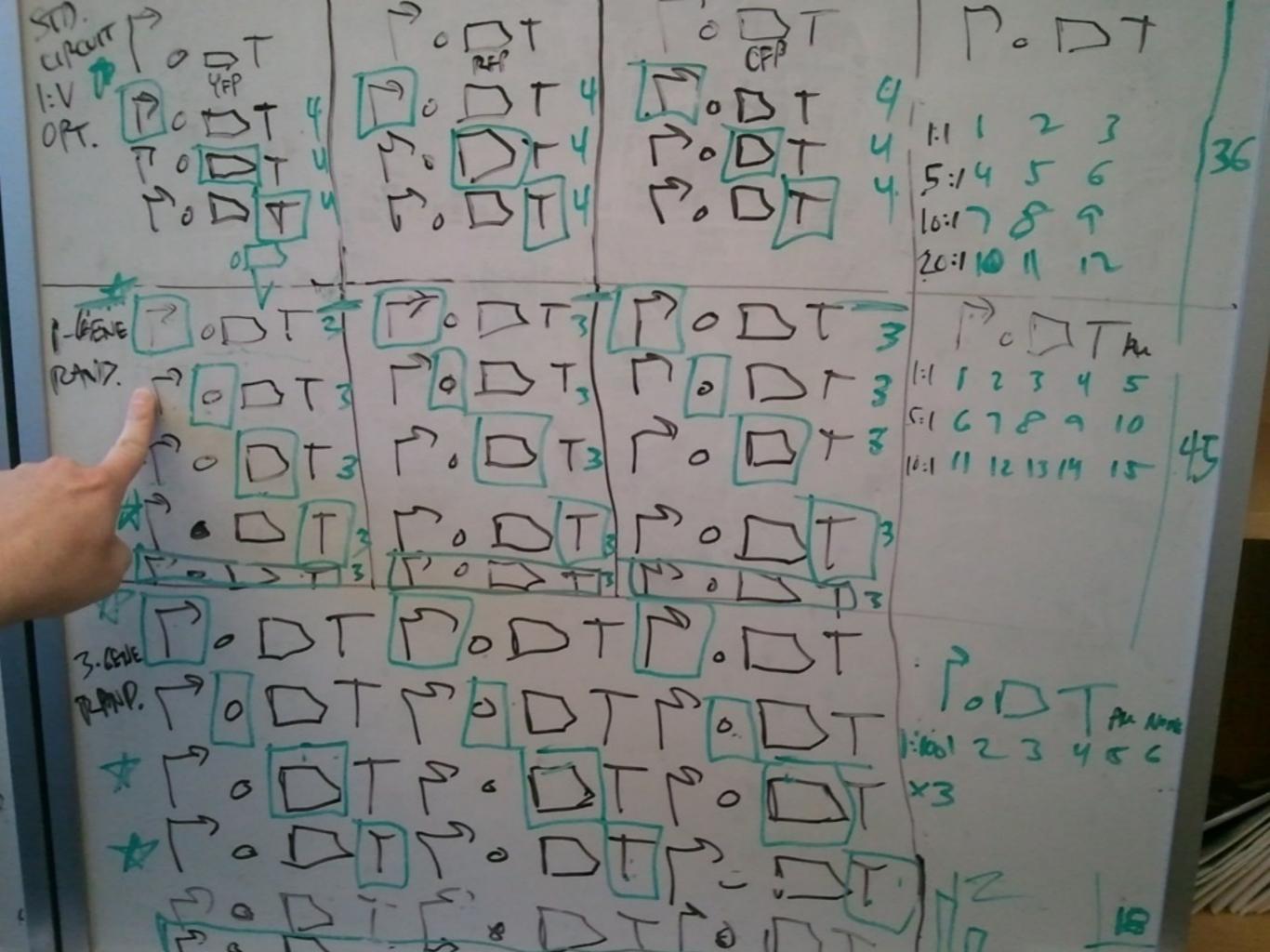
promoter	O origin of replication
cds	primer binding site
ribosome entry site	blunt restriction site
terminator	sticky restriction site
operator	— 5' overhang
insulator	= 3' overhang
ribonuclease site	= assembly scar
rna stability element	× signature
X protease site	user defined
P protein stability element	

standardized symbol set



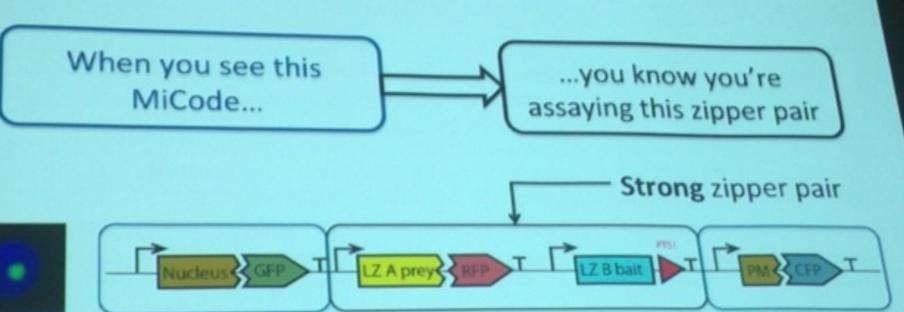
relationship to SBOL

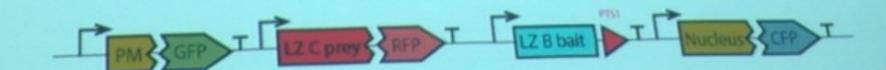
use in various contexts





Using MiCodes in a zipper assay





MiCodes

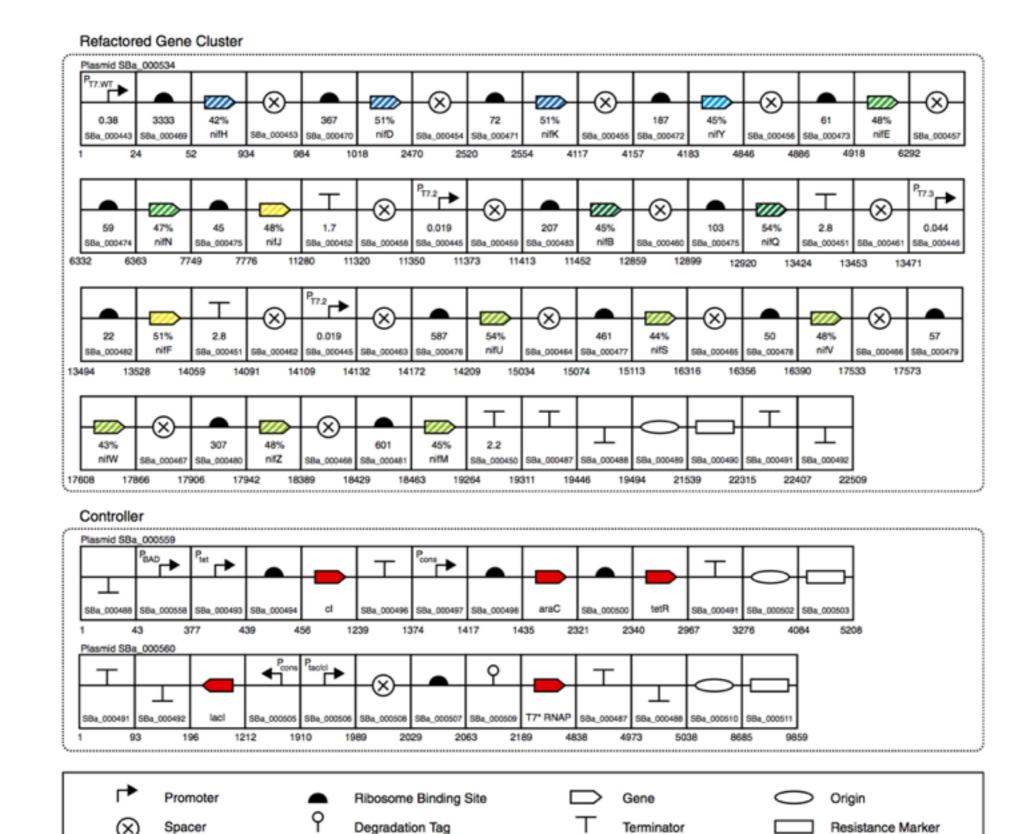
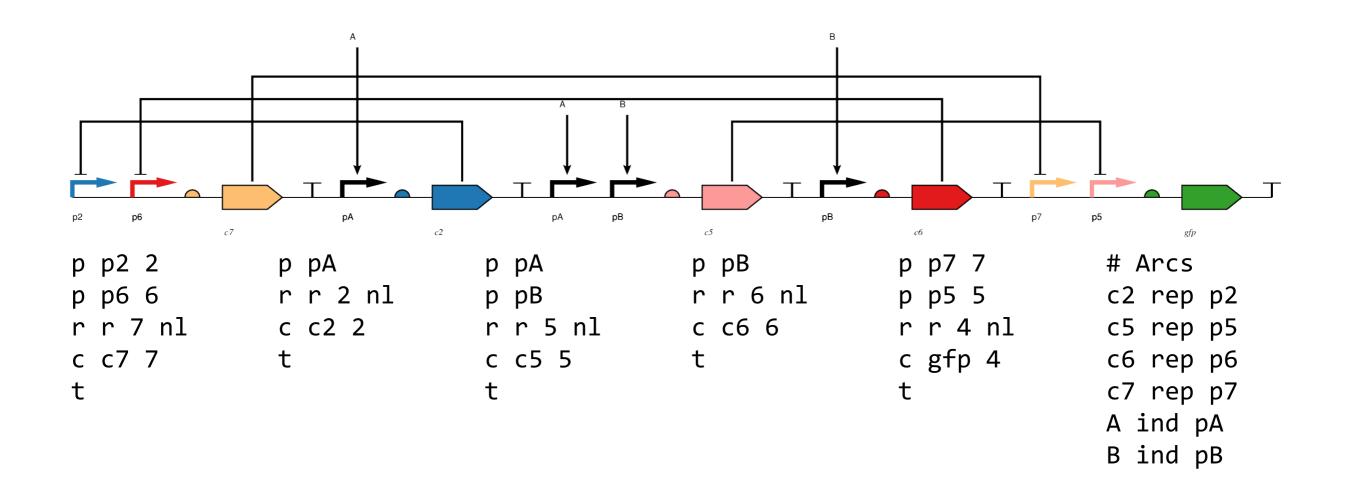
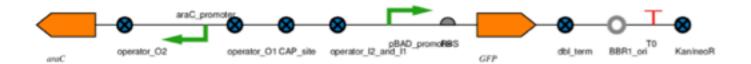


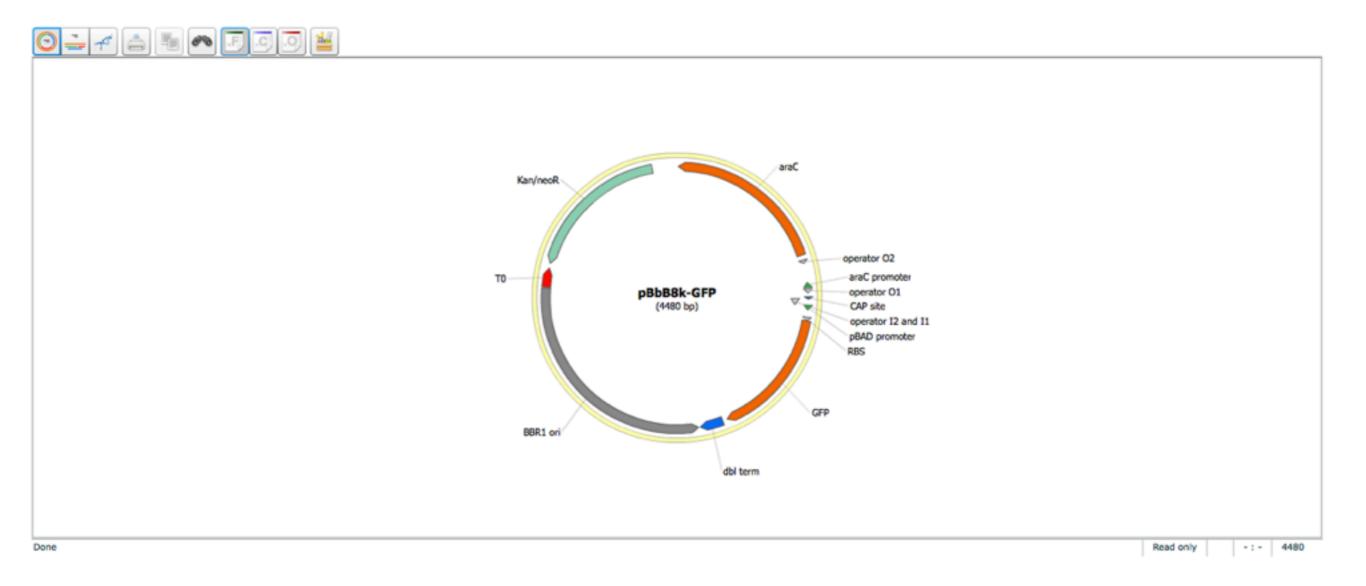
Fig. 4. Comprehensive schematic illustration for the complete refactored gene cluster and controller. Each of the 89 parts is represented according to the Synthetic Biology Open Language visual standard (www.sbolstandard.org), and the SynBERC Registry part number (registry.synberc.org) and part activity are shown. The full sequences of each plasmid have been deposited in GenBank (SBa_000534, JQ903614; SBa_000559, JQ903615; SBa_000560, JQ903616). The T7 promoter strengths are measured with monomeric red fluorescent protein and reported in REUs (Materials and Methods). Terminator strengths are measured in a reporter plasmid and reported as the fold reduction in monomeric red fluorescent protein (RFP) expression compared with a reporter without a terminator. The RBS strength is reported in as arbitrary units of expression from the induced Ptac promoter (1 mM IPTG) and a fusion gene between the first 90 nt of the gene and RFP. The nucleotide numbers for the plasmids containing the refactored cluster and controller are shown. The codon identity of each recoded gene compared with WT is shown as a percentage.

software

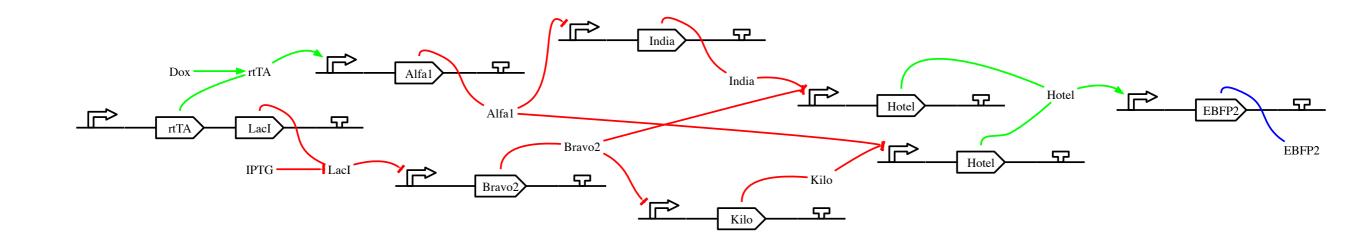


PigeonCAD pigeoncad.org

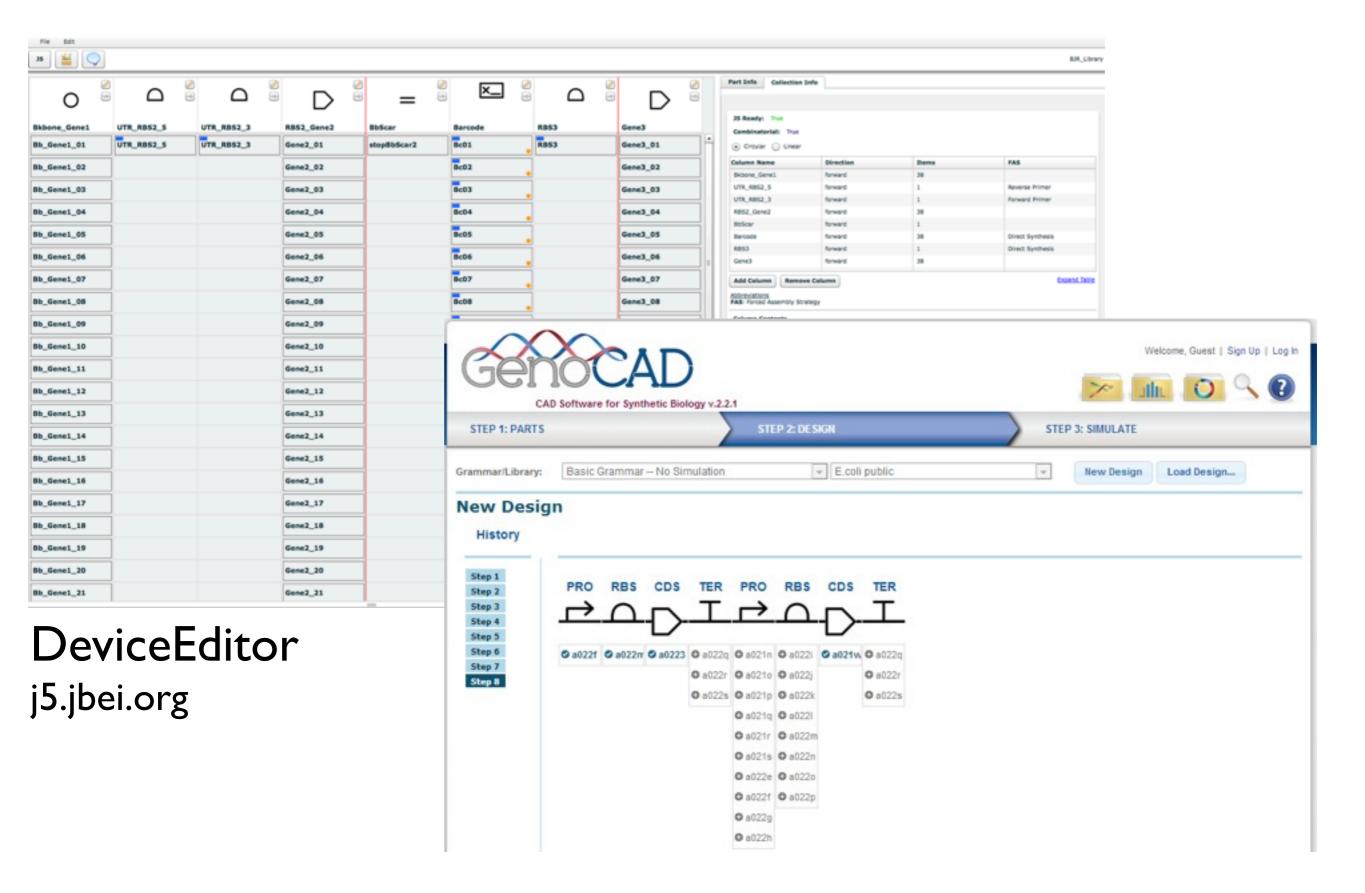




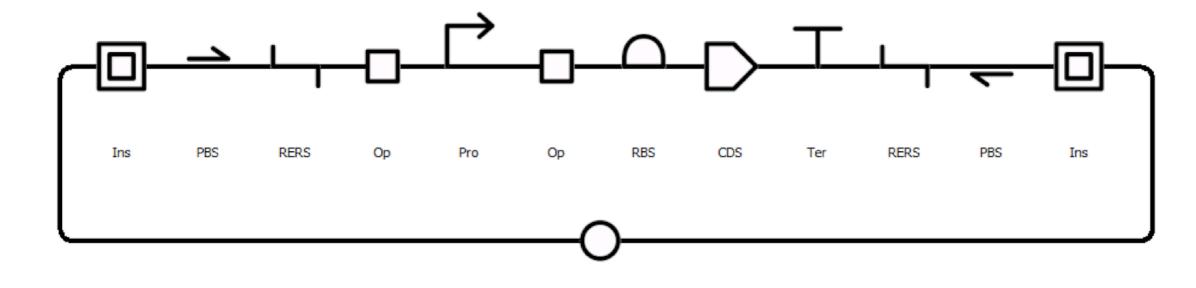
JBEI-ICE public-registry.jbei.org

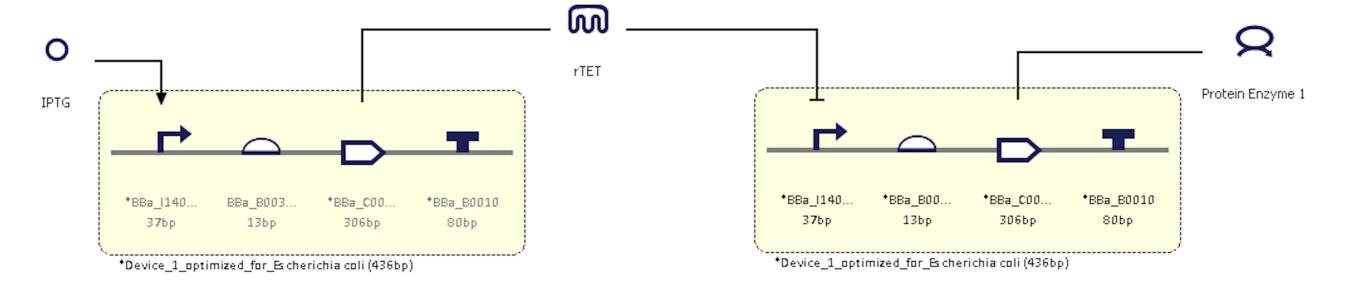


BioCompiler synbiotools.bbn.com



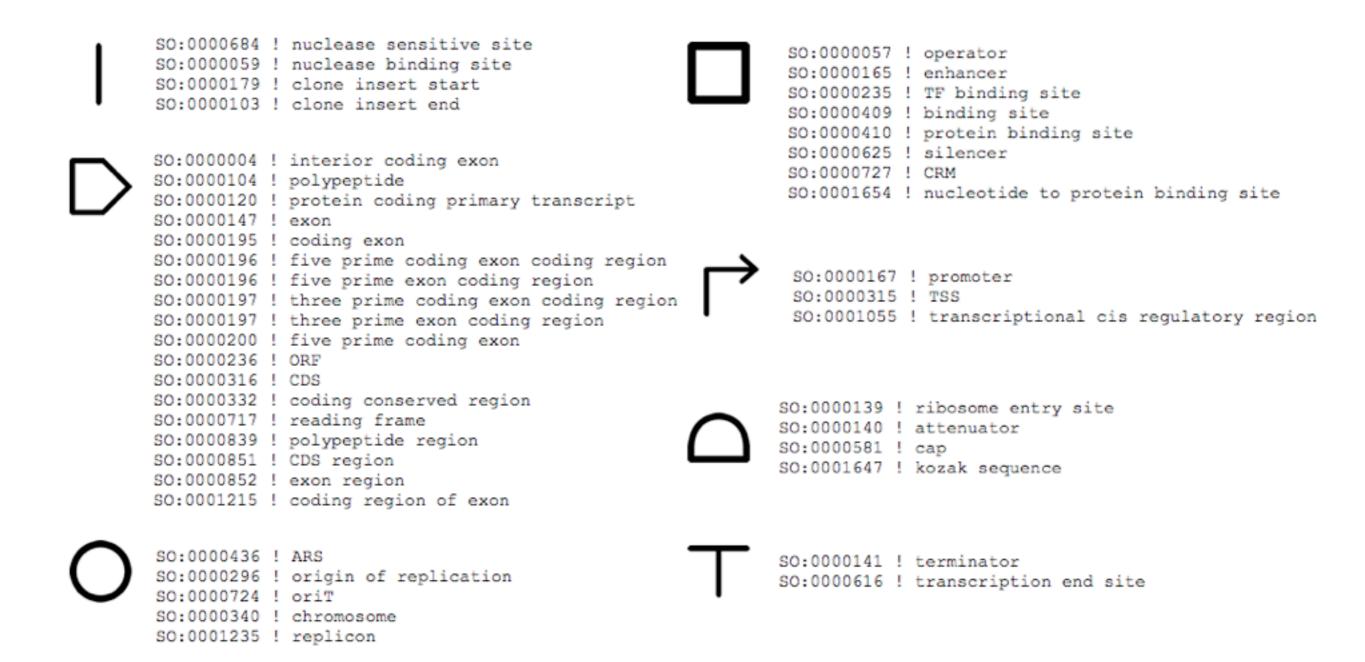
GenoCAD genocad.org



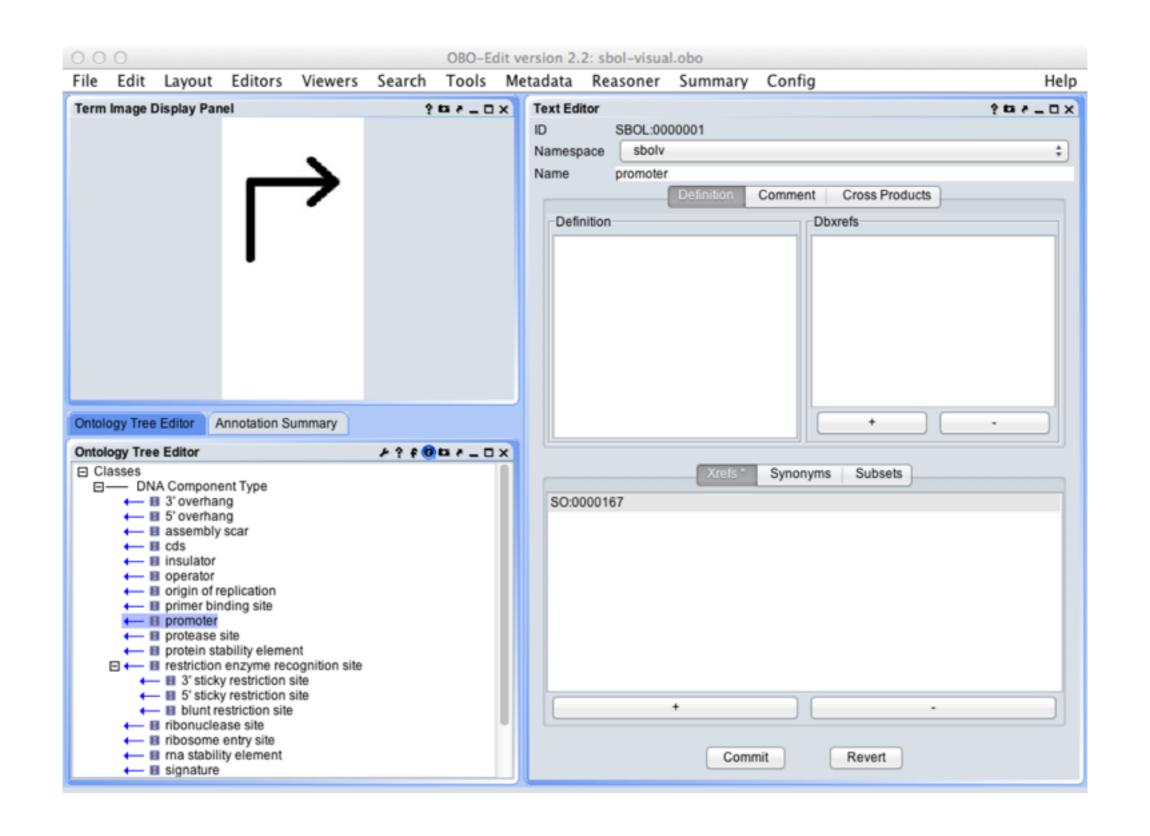


VectorNTI Express Designer

next steps



further ontological mappings



serializations and versioning



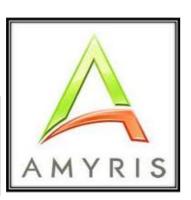




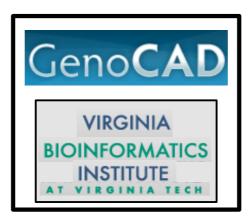


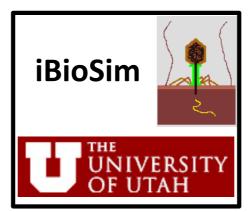










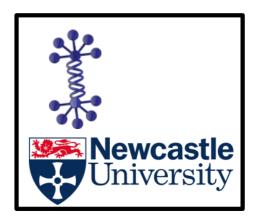












Use SBOL Visual!

- Use SBOL Visual!
- Tell us how it goes

- Use SBOL Visual!
- Tell us how it goes
- Submit new symbols

- Use SBOL Visual!
- Tell us how it goes
- Submit new symbols
- Join the Working Group

SBOL Visual Working Group



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Chris Myers

Umesh P
Matthew Pocock
Cesar Rodriguez
Herbert Sauro
Larisa Soldatova
Guy-Bart Stan
Grimaldo Urena
Alan Villalobos
Mandy Wilson

Thank You!

www.sbolstandard.org/visual visual@sbolstandard.org

promoter	O origin of replication
cds	primer binding site
ribosome entry site	blunt restriction site
terminator	sticky restriction site
operator	— 5' overhang
insulator	= 3' overhang
ribonuclease site	assembly scar
rna stability element	× signature
X protease site	user defined
P protein stability element	