BBF RFC 66: A RESTful API for Supporting Automated BioBrick Model Assembly

Steyn, J. S.; Boyd, R. M; Essa, Y.; Hall, P.; Koh, A.; Sheth, H.; Tsu, D; Woodhouse, S; Hallinan, J.; Wipat, A.; Pocock, M. R.

23 September 2010

Purpose

Constructing simulatable models for BioBricks by hand is a complex and time-consuming task. The time taken could be reduced by using Computer Aided Design (CAD) tools to aid in designing models, but these tools need to be augmented with domain-specific knowledge. Here we propose a standard for a RESTful (Richardson, 2007) API which facilitates the discovery and publication of models of functional biological units. This API is designed to produce parts models which can be automatically combined into complete, simulatable models of entire systems.

Relation to other BBF RFCs

BBF RFC 66 does not update or replace any earlier BBF RFC.

Copyright Notice

Copyright (C) The BioBricks Foundation (2010). All Rights Reserved.

Motivation

Computational simulation is an integral part of the BioBrick design process. However, given that realistic systems often involve multiple BioBricks, and that each BioBrick may contain multiple functional components (Knight, 2003), constructing simulatable models of biological systems can be a daunting task.

Genetic circuits can be designed either manually, using CAD tools (e.g. Chandran, Bergmann & Sauro, 2009), or automatically (Rodrigo, Carrera & Jaramillo, 2007). To support these approaches several researchers have proposed methods for the bottom-up assembly of BioBricks (Marchisio & Stelling, 2008, Cooling et al., 2010). To achieve this task these tools require access to a library of models for individual parts, and enough additional information about these parts to allow the user to quickly identify valid combinations which allow the software to combine model fragments into larger models.

Here we describe a RESTful service API for querying and fetching parts, model fragments for these parts, and combinatorial information about these parts, to support both automated and human-directed construction of complex models.

Terminology and Typography

Text appearing in fixed-width font denote parts of the RESTful API, such as data-types and operations. Text appearing in *italic* font indicate names and identifiers. The words 'part', 'device' and 'system' in standard font have the usual BioBrick meaning.

Data Model

The data model is hierarchical (Fig. 1). At the top level, an assembly bundles parts, connections and other assemblies together, producing re-usable, simulatable systems. Parts represent biological primitives and their associated models, including any publicly visible variables. Connections join parts together via their variables.



Figure 1. Data model class diagram

Data model types

There are four data types in the data model. Their structure is described in this section.

•Assemb | y: A device or system that can be independently simulated

An Assembly has the following attributes:

- •name: The assembly name.
- •description: An optional human-readable description of the assembly.
- •subAssembly: Zero or more assemblies of which this assembly is composed.

•hasParts: The parts assembled by this assembly. Individual Parts may be assembled into any number of assemblies.

•hasConnections: The connections that wire parts together. For example, the *POPs* output of a promoter may be wired to the *POPs* input of a coding sequence.

•Part: An encapsulation of a quantified entity

A Part consists of:

•name: The part name.

•description: An optional human-readable description of the part.

•type: The biological type of the part, such as *Promoter*, *CDS*, *RBS*. This SHOULD come from an appropriate controlled vocabulary.

•model: A MathML block describing the relationships between the variables of this part.

•hasVariables: The variables exposed.

•Variable: An exposed quantity

A Variable has:

•name: No two variables within the same part may share the same name. However, variables from different parts may have the same name.

•units: The physical units of the variable. Defaults to *dimensionless*.

•isin: A flag to indicate if this is an input variable for the part. Defaults to false

• isOut: A flag to indicate if this is an output variable for the part. Defaults to *false*

•Connection: A binding of an out-variable from one component to an invariable of another component

• in: the in-variable to connect

•out: the out-variable to connect

Data model well-formedness constraints

For data to be well-formed, it must both conform to the data model described above, and additionally adhere to the following constraints:

•assembly. name (scope): each name uniquely identifies a single assembly. Assemblies MUST NOT share names. These MAY be re-used between entities of different types, for example between an assembly. name and a part. name.

•part. name (scope): each name uniquely identifies a single part. Parts MUST NOT share names. These MAY be re-used between entities of different types, for example between a part. name and a variable. name.

•part. hasVariables and part. model (scope): Every name of every variable of a part MUST appear in the model of that part.

•Variable. name (scope): Every name of every variable of a part MUST be unique within that part. Variables of different parts MAY (and in some circumstances MUST) use the same name.

•connection. in and connection. out (flags): These MUST refer to variables with the isln and isOut flag set to *true*, respectively.

•connection. in and connection. out (scope): These MUST refer to variables that are within parts that are within the assembly holding the connection.

Data Rendering

Individual implementers are free to chose from one of several renderings of the data model. These include XML and JavaScript Object Notation (JSON).

To keep per-fetch costs low, this specification states what data MUST be present in a particular rendering. In particular, it states what MUST be directly embedded in-line as a value, and what MUST be included by reference to a name.

Field	Туре	Encoding	
Assembly			
name	string	value	
description	string	value	
subAssembly	List of assembly	reference	
		\${subAssembly.name}	
hasPart	List of part	reference	
		\${hasParts.name}	
hasConnections	List of connection	value	
Part			
name	string	value	
description	string	value	
type	string(cv)	value	
model	MathML	value	
hasVariables	List of variable	value	
Connection			
in	variable	reference \${in.name}	
out	variable	reference \$ {out. name}	
Variable			
name	string	value	
units	string(cv)	value	
isIn	boolean	value	

isOut boo	lean	value
-----------	------	-------

Component Assembly

A CAD client is responsible for allowing the user to combine parts and assemblies to build a model of the system of interest. This is achieved by binding all unbound variables. A variable is unbound if it is not referred to by any connection. To ensure that the model can be simulated, the CAD program will need to bind each unbound in-variable to either an unbound out-variable, or to a constant value. For example, if an assembly modeling the *Lac* operon exposes in-variables named *lac_repressor* and *inactivated_lac_repressor*, these variables must be bound either to out-variables giving the levels of these molecules, or to suitable constant values.

Resources

REST is based around resources, identified by URLs. Given, for example, a base URL called BASE, the following resources MUST be provided by any implementation of this API: BASE/parts/ representing all parts; BASE/parts/ $\$ [name] for each part; BASE/assemblies/ representing all assemblies; and BASE/assemblies/ $\$ [name] for each assembly. For example, a server located at 'http://myuni.ac.uk/modelrepository' which published an assembly $BA_K302012$, would be exposed as: 'http://myuni.ac.uk/modelrepository/assemblies/BA_K302012'.

URL	Description			
parts/	All parts			
	Operation	Result	Example	
	GET	The names	Response: ['generic_promoter', 'Pspac',	
		of all parts	'Pveg']	
	POST	Create a	Query: {	
		new part	name: 'Plac',	
			description: 'Lac Operon promoter',	
			type: 'promoter',	
			model: '', hasVariables: [{	
			name: 'RNA_polymerase',	
			isln: true }, {	
			name: 'inact_lac_repressor',	
			isln: true }, {	
			name: 'POPs'.	
			isOut: true }]}	
			Result: <u>parts/Plac</u> created	
parts/	A part			
\${name}	Operation	Result	Example	
	GET	Fetch the	Response: { name: '\${name}',	
		part	··· }	
assemblies/	All assemblies			
	Operation	Result	Example	

	GET	The names of all assemblies	Response: ['Lac Operon', 'Bba_K302012']
	POST	Create a newassem bly	Query: { name: 'BBA_K302012', description: 'Filamentous cell phenotype', subAssembly: ['BBA_K302003', 'BBA_K302005'], hasConnections = [{ in: 'BBA_K302003.POPs', out: 'BBA_K302005.POPs'}] Result: assemblies/BBA_K302012created
assemblies/	An assembly		
\${name}	Operation	Result	Example
φ(name)	GET	Fetch an assembly	Response: { name: '\${name}', }

Authors' Contact Information

Jannetta S. Steyn: jannetta.steyn@ncl.ac.uk Rachel May Boyd: rachel.boyd@ncl.ac.uk Younus Essa: younus.essa@ncl.ac.uk Phil Hall: phillip.hall@ncl.ac.uk Alan Koh: s.c.a.koh@ncl.ac.uk Harsh Sheth: harsh.sheth@ncl.ac.uk Deena Tsu: deena.tsu@ncl.ac.uk Steven Woodhouse: steven.woodhouse@ncl.ac.uk Anil Wipat: anil.wipat@ncl.ac.uk Jennifer Hallinan: j.s.hallinan@ncl.ac.uk Matthew Pocock: <u>matthew.pocock@ncl.ac.uk</u>

References

Chandran, D., Bergmann, F. T. & Sauro, H. M. (2009). TinkerCell: modular CAD tool for synthetic biology. Journal of Biological Engineering 3(19): doi:10.1186/1754-1611-3-19.

Cooling, M. T., Rouilly, V., Misirli, G., Lawson, J., Yu, T., Hallinan, J. & Wipat, A. (2010). Standard virtual biological parts: a repository of modular modeling components for synthetic biology. Bioinformatics 26: 925 - 931.

Knight, T. (2003). Idempotent vector design for standard assembly of Biobricks. In Knight, T. MIT Synthetic Biology Working Group Technical Report 0 http://hdl.handle.net/1721.1/21168.

Marchisio, M. A. & Stelling, A. (2008). Computational design of synthetic gene circuits with composable parts. Bioinformatics 24(17): 1903 - 1910.

Richardson, L. (2007). RESTful Web Services. Pragma.

Rodrigo, G., Carrera, J. & Jaramillo, A. (2007). Genetdes: Automatic design of transcriptional networks. Bioinformatics 23(14): 1857 - 1858.