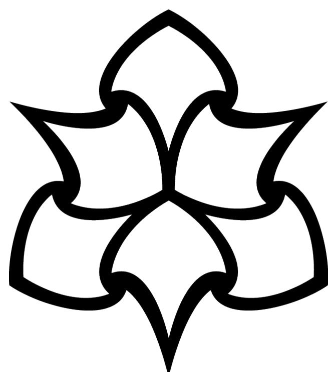

Manchester Metropolitan University

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A Genetic Circuit Compiler:

Generating Combinatorial Genetic Circuits with Web Semantics and Inference

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Abstract

A central strategy of synthetic biology is to understand the basic processes of living creatures through engineering organisms using the same building blocks. Biological machines described in terms of parts can be studied by computer simulation in any of several languages or robotically assembled *in vitro*. In this paper we present a language, the Genetic Circuit Description Language (GCDL) and a compiler, the Genetic Circuit Compiler (GCC). This language describes genetic circuits at a level of granularity appropriate both for automated assembly in the laboratory and deriving simulation code. The GCDL follows Semantic Web practice and the compiler makes novel use of the logical inference facilities that are therefore available. We present the GCDL and compiler structure as a study of a tool for generating κ-language simulations from semantic descriptions of genetic circuits.

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Keywords

Semantic Web, Inference, Program Generation, Synthetic Biology, Genetic Circuits

Synthetic biology extends classical genetic engineering with concepts of modularity, standardisation, and abstraction drawn largely from computer engineering. The goal is ambitious: to design complex biological systems, perhaps entire genomes, from first principles[1]. This enterprise has met with some success such as the microbial production of drug synthesis[2,3], new biofuels production[4] and alternative approaches to disease treatment[5]. However, most applications are still small and mostly designed manually.

The are several obstacles to designing more complex circuits. The design space of potential circuits is very large. Even when a design is chosen, there is large *a priori* uncertainty about what its behaviour will be. In many cases the available information about molecular interactions in a cell is incomplete. A secondary obstacle is that designs can be brittle and very sensitive to the host environment in which they execute. In this context computational techniques become important for identifying biologically feasible solutions to problems of biological system synthesis. Beyond the challenges of the huge design space and associated uncertainties, writing these programs by hand is time-consuming and error prone, and there are very few tools available for verification and debugging them. Descriptions of models in terms of simulation code are tightly coupled to the language of the simulation program, and it may be difficult or impossible to use a different interpreter without completely rewriting the code.

We solve these problems by providing a high-level, modular, implementation-independent language for describing gene circuits called the Genetic Circuit Description Language (GCDL) and a compiler called Genetic Circuit Compiler (GCC). We use a strategy of contextual reasoning to obtain flexible output from this succinct input, and *templates* to support any number of output languages and modelling granularities. An overview of in-

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formation flow through the compiler is shown in Figure 1. We demonstrate the utility of this approach by describing, compiling and simulating a complete genetic circuit, the well-known Elowitz repressilator[6]. The compiler and example code are available at `https://github.com/rulebased/composition`.

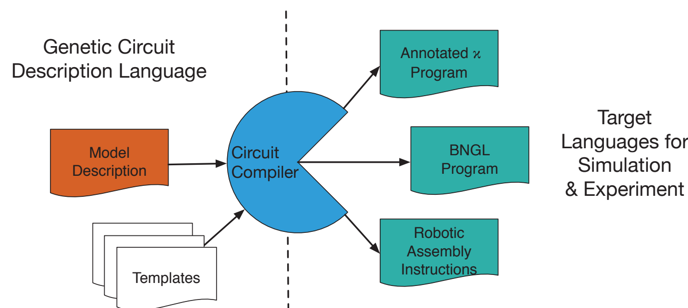

Figure 1: High-level data flow through the compiler. The compiler for synthetic gene circuits takes a model description written in GCDL and, using language-appropriate appropriate templates, creates code for simulation and laboratory assembly. We have implemented templates for annotated-κ for the KaSim software, and envision similar for the BNGL as well as SBOL.

Code generation from this high-level description to a low-level language for simulation greatly reduces the scope for error in coding simulations. Because the language is implementation-independent it is not tightly coupled to any particular interpreter or hardware. In this way GCDL facilitates *evergreen models*, models that are specified sufficiently well to be unambiguous but not so specifically that they can only be executed or constructed in one software package or environment.

Domain specific languages and examples of compilers processing these languages have previously been shown[7–10]. These languages are designed to allow for simulations using a particular methodology such as solving systems of ordinary differential equations or using Monte-Carlo simulation. Unlike previous approaches, we emphasise the use of abstraction to facilitate *retargeting* or production of output suitable for different simulation environments and techniques as well as automated circuit assembly in the laboratory from a single description. Compiler targets are implemented using conditional inference, defining the semantics of the terms used in the description of the circuit in a way that is determined by the desired

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output type. The design of the compiler is general, and not limited to the present context of genetic circuits. The design shown schematically in Figure 2.

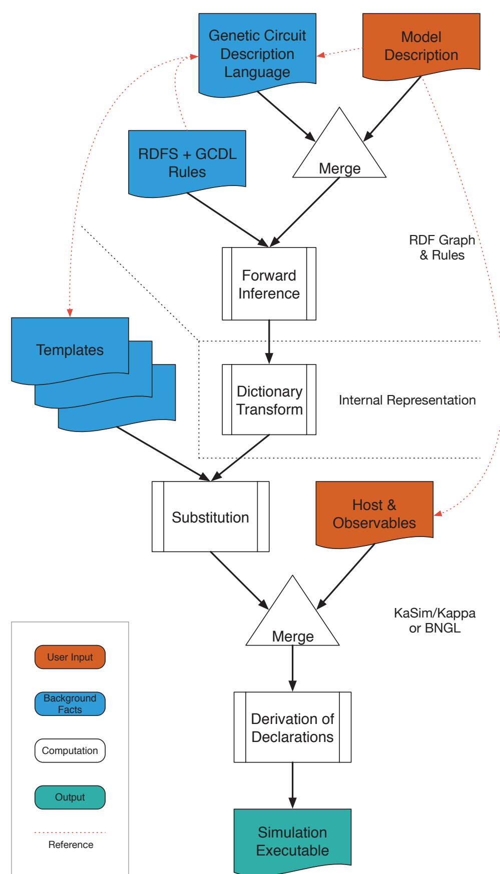

Figure 2: Detailed data flow through the compiler. This illustrates the use of inference to expand the GCDL model to derive consequent information appropriate to producing the next stage of output in the specific target language.

The GCDL is an RDF[11] vocabulary and attendant inference rules which facilitates gathering and collation of information about the constituent parts of a genetic circuit[12]. The output programs can be specialised to various languages, such as the KaSim flavour of κ[13,14], BioNetGen's BNGL[15,16], other representations such as SBOL[17] or indeed whichever form is required by robotic laboratory equipment that assembles circuits *in vitro*. This output flexibility is accomplished using *templates* that use facts derived by inference rules[18] from the input model.

We now proceed as follows. We begin with an overview of those aspects of synthetic

biology and genetic engineering that are necessary to contextualise our work. Next, we explain the representation of this kind of genetic circuit model in GCDL, this is the main input to the compiler. In order to understand the desired output of the compiler, we then illustrate how these constructs are represented as rule-based code for the κ language simulator, KaSim. There follows a discussion of how the compiler infers the executable model from the input description. Finally, we discuss some possible uses and limitations of our technique.

Background

Rule-based Modelling of Genetic Processes

A weakness of reaction-based methods for modelling the processes of transcription, translation and the production of chains of proteins is that they require chemical species for each bound state of the reagents. This in turn requires specification of reactions for each combination of these reagents. To solve this problem of needing combinatorially many reactions to describe substantially the same process, a generalisation of reactions called *rules* are used[19–21].

In the rule-based representation, *agents* correspond to reagents and they can have slots or *sites* that can be bound, or not. They can also have internal state. Unlike reactions which have no preconditions apart from the presence of the reagents, with rules, a configuration of the sites — bound in a particular way, bound in some way, unbound, or unspecified — is a precondition for the application of the rule. A rule may re-arrange the bonds, creating or destroying them, without the need to invent new agents in order to represent different configurations of a given set of molecules.

The reader should note that the word *rule* is used in two distinct senses in this article. The first is as we have just described. The second is in the sense of *inference rule* as used in logic and in particular the way in which we deduce executable rule-based models from their

declarative representations in RDF.

The κ Language

To briefly illustrate the essentials of rule-based modelling we will use the language of the Kappa simulation software, KaSim[14]. An agent declaration and rule expressing the formation of a polymer can be written as,

```
%agent: A(d,u)

'binding' A(u[.]), A(d[.]) -> A(u[1]), A(d[1]) @k
```

We can gloss this as an agent with two sites, `u` and `d` for upstream and downstream, and a rule. The rule concerns two agent patterns one of which has an unbound upstream site, and the other an unbound downstream site, and the action of the rule is to bind them, the notation `[1]` denoting the bond. This process happens at some rate `k`.

The state of the other site of each agent is left unspecified, so implicit in this rule is the possibility that either or both the agents may already be bound to others and so part of arbitrarily long chains. In other words this expression covers not only two monomers joining together but an n-mer and an m-mer for arbitrary n and m. This is the essence of the expressive advantage that rule-based modelling provides. To express a similar concept using a reaction network would in fact require infinitely many reagents for every possible n (and m) and infinitely many reactions for every possible combination.

Biological Parts and Annotation

For efficiency, and economy of representation, we claim that the description of a computational model should include minimum information necessary for simulation. However, in order to use these models in an automated design process, additional metadata, or *annotations*, about the meaning of different modelling entities is needed[12]. Annotation facilitates

the drawing specific parts from a database such as the Virtual Parts Repository[22]. Models in that database are annotated with machine-readable metadata intended for combination into larger models. Myers and his colleagues have used annotations to derive simulatable models from descriptions of genetic circuits[23] and vice versa[24], though these use reaction-based techniques and so inherit the poor scaling properties of that method.

To facilitate the *in silico* evaluation of potential synthetic gene circuits, a library of descriptions of genetic parts, together with their modular models is suggested in[22,25]. These parts are intended to be large enough to have a particular meaning or function (i.e. larger than individual base pairs) but not so large that they lack the flexibility to be recombined (i.e. entire genes). Thus we are concerned with coding sequences for particular proteins, promoters that, when activated, start the transcription process, operators that activate or suppress promoters according to whether they are bound or not by a given protein, and a small number of other objects. A sequence of these objects is a genetic circuit, and our goal is to have a good language for describing such sequences.

Annotation in this setting means machine-readable descriptions of entities of biological interest. This is done with statements, triples of the form (subject, predicate, object) according Semantic Web standards[11,26]. Entities are identified with Universal Resource Identifiers (URIs)[27]. This provides the dual benefit of globally unique identifiers for entities and a built-in mechanism for retrieving more information about them providing that some care is taken to publish data according to best practises[28,29]. Large bodies of such information about biologically relevant information are published on the Web[30,31] and the use of Semantic Web standards for annotating our models allows us to express how an entity in a model description corresponds to a real world protein, or gene sequence or other entity.

The Semantic Web also affords us a technical advantage: inference rules. These can be either explicit as in Notation3[32,33] or implicit as in OWL Description Logics[34,35]. In either case this facility makes it possible, given a set of statements, to derive new statements according to inference rules. We use this to improve the ergonomics of our high-level language:

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while the compiler itself will make use, internally, of a large amount of information, we do not expect the user to supply it in painstaking detail. Rather, we allow the user to specify the minimum possible and provide rules to derive the necessary detail. Inference rules provide for both economy of representation for the high-level model description and flexibility for the different implementations.

A Language for Synthetic Gene Circuits

This section describes the GCDL, the high-level language for describing genetic circuits made from standard biological parts[25 22]. We begin by stating the properties that we want in such a language and showing how we achieve them. There follows a synopsis of the vocabulary terms essential to the language. Finally, we illustrate salient language features applied to example circuits.

Desired Language Features

Our desired language features for high-level representation of a genetic circuit are as follows,

1. sufficiency, there should be enough information to derive executable code for the circuit,
2. identifiability, it should be possible to determine to which biological entities (DNA sequences, proteins) the representation refers,
3. extensibility, it should be straightforward to add information or constructs that are not presently foreseen,
4. generality, there should be no requirement that information about biological parts comes from any particular set or source, and
5. concision, there should be a minimum of extraneous detail or syntax.

The third and fourth requirements are readily met by using RDF as the underlying data model. The *open world* presumption[36] means that adding information as necessary is straightforward. The use of URIs[27] which can be dereferenced to obtain the required information means that information from different web-accessible databases can be obtained, mixed and matched as desired. The use of URIs goes some way towards meeting the second requirement, albeit with some well-known caveats[37].

The first and last of the desired features are the primary areas of innovation of the present work. We suggest (but do not require) the use of Turtle[38] or indeed Notation3[18] as the concrete surface syntax for writing models. This goes some way towards a representation that is intelligible by humans. Even then, we aim to minimise what needs to be written and we do this using inference rules — if a needed fact can be derived from the model under the provided rule-set, it is unnecessary to write it explicitly in the model. Indeed it may even be undesirable to do so since it is a possible source of errors, for example some kinds of assertions may be correct in the context of some output types and incorrect in others. We aim for a minimal, yet complete under the inference rules, description of the model.

Vocabulary Terms

New terms introduced in this paper have the prefix `gcc` which can be read as the "Genetic Circuit Compiler" vocabulary. The list of terms is reproduced in Table 3 and their complete definitions are given together with the accompanying rules in the supplementary materials. The GCDL is the union of terms from the *gcc* namespace with those from the Rule-Based Model Ontology (RBMO) that we previously defined[39] together with terms from the Simple Knowledge Organization System (SKOS)[40] vocabulary, RDF Schema (RDFS)[35] and Resource Description Framework (RDF)[11].

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Table 3: Selected terms from the GCC vocabulary

Classes	
gcc:Part	Generic biological part
gcc:Operator	Operator
gcc:Promoter	Promoter
gcc:RibosomeBindingSite	Ribosome Binding Site
gcc:CodingSequence	Coding Sequence
gcc:Terminator	Terminator
gcc:Token	Token or symbol in a template

Predicates	
gcc:include	Include a low-level model fragment
gcc:prefix	The prefix to use for generated annotations
gcc:init	Specifies initial copy numbers
gcc:part	Links a part to its token or symbol
gcc:overlaps	Indicates that two parts overlap (symmetric)
gcc:linear	Linear circuit type
gcc:circular	Circular circuit type
gcc:transcriptionFactor	Relates an operator to its transcription factor
gcc:transcriptionFactorBindingRate	Various rates
gcc:transcriptionFactorUnbindingRate	
gcc:rnapBindingRate	
gcc:rnapUnbindingRate	
gcc:rnapRNAUnbindingRate	
gcc:ribosomeBindingRate	
gcc:ribosomeRNAUnbindingRate	
gcc:ribosomeProteinUnbindingRate	
gcc:transcriptionInitiationRate	
gcc:transcriptionElongationRate	
gcc:translationElongationRate	
gcc:rnaDegradationRate	
gcc:proteinDegradationRate	

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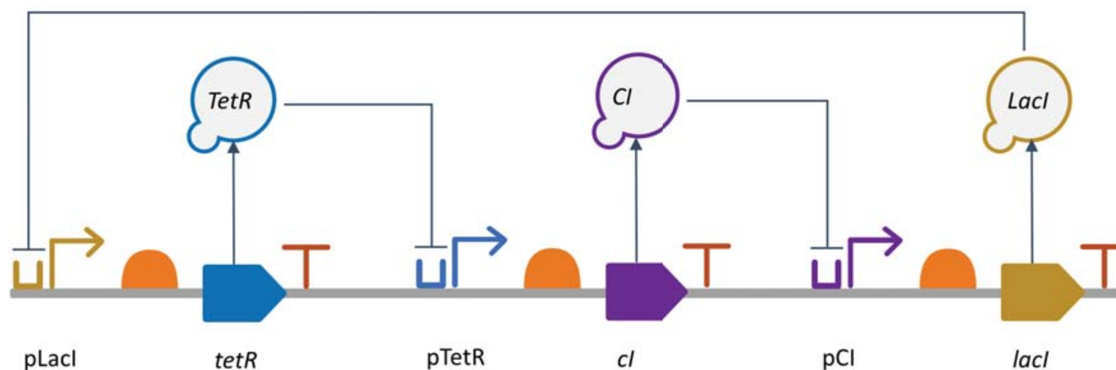
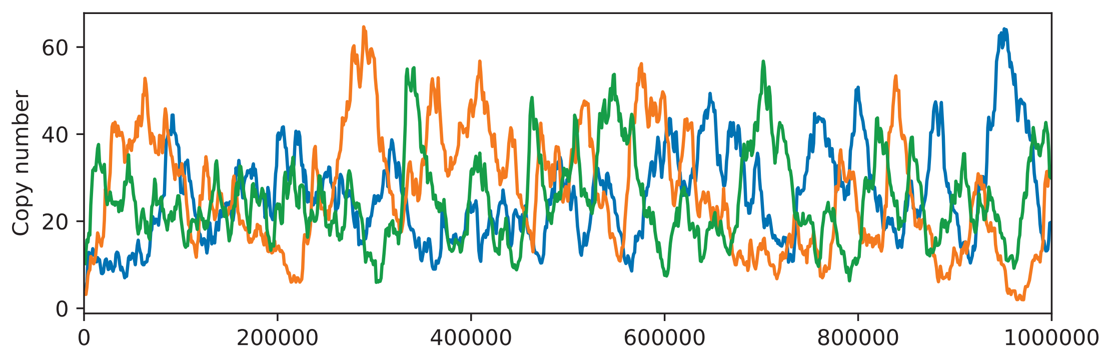


(a) An example genetic circuit: the Elowitz repressilator. It is a negative feedback oscillator. The circuit is arranged linearly. Protein production and inhibitory protein-operator relationships are shown using the SBOL visual standard.



(b) Sample simulation data from a program produced by the compiler showing the expected oscillations. Note in particular the relatively small copy numbers of the proteins for which stochastic simulation in the κ language is well suited.

Model Description

To illustrate the syntax of the high-level language, we use the well known Elowitz repressilator shown diagrammatically in Figure 3a. The complete model can be found in the supplementary materials as well as distributed in the `examples/` subdirectory of the compiler distribution. Also included with the compiler is a hand-assembled implementation of this circuit for comparison. A sample trace produced by generated program is shown in Figure 3b. Figure 4 shows a description of this the core of the model, in the GCDL. Some bibliographic metadata is included, using the standard Dublin Core[41] vocabulary, as well as a generic pointer (`rdfs:seeAlso`) to a publication about this model.

The term `gcc:prefix` is necessary in every model, it instructs the compiler that any

```
## Model declaration
:m a rbmo:Model;
  ## bibliographic metadata
  dct:title "The Elowitz repressilator constructed from BioBrick parts";
  dct:description "Representation of the Elowitz repressilator given in the Kappa BioBricks Framework
       book chapter";
  rdfs:seeAlso <http://link.springer.com/protocol/10.1007/978-1-4939-1878-2_6>;
  gcc:prefix <http://id.inf.ed.ac.uk/rbm/examples/repressilator#>;
  ## include the host environment
  gcc:include <.../host.ka>;
  ## initialisations
  gcc:init
    [ rbmo:agent :RNAp; gcc:value 700 ],
    [ rbmo:agent :Ribosome; gcc:value 1000 ];
  ## The circuit itself, a list of parts
  gcc:linear (
    :R0040o :R0040p :B0034a :C0051 :B0011a
    :R0051o :R0051p :B0034b :C0012 :B0011b
    :R0010o :R0010p :B0034c :C0040 :B0011c
  ).
```

Figure 4: Example model for a synthetic gene circuit, Elowitz' repressilator.

entities that it creates should be created under the given prefix. Ultimately annotated rules will be generated for the low-level representation and the annotated entities require names. To give them names, a namespace is required and this is how it is provided.

Next there is a `gcc:include` statement. This is a facility for including extra information in the low-level language. Extra information typically means rules for protein-protein interactions which are beyond the scope of the current work and as such it is simply supplied as a program fragment in the output language. This corresponds roughly to calling an assembly or machine language routine to perform a specialised task when programming a computer in a high-level language like C.

There follows initialisation for specific variables. In this case these are the copy numbers for RNA polymerase molecules and ribosomes. These are denoted using `rbmo:agent` because of our choice to support rule-based modelling for greater generality than reaction-based methods. Finally, the circuit itself is specified. The argument, or object is an `rdf:List` which simply contains identifiers for the parts, in order.

The circuit itself is now defined. However at this juncture, we simply have a list of parts without having specified what they are or what their intended behaviour is. To obtain a working model, we need more.

```
:C0012 a gcc:CodingSequence;
    gcc:label "Coding sequence for LacI";
    gcc:part "C0012";
    gcc:protein :P0010;
    gcc:proteinDegradationRate 0.0001.

:P0010 a gcc:Protein;
    bqbiol:is uniprot:P03023;
    skos:prefLabel "P0010";
    rdfs:label "LacI".
```

Figure 5: A coding sequence part description from the repressilator model. Notice how the coding sequence is linked to the protein that it codes for.

A Part Description

A simple example of a part description is shown in Figure 5. This is a coding sequence, as is clear from the type annotation on the part. It codes for a particular protein, specified with `gcc:protein`. This term is specific to proteins because under normal circumstances other kinds of part do not code for proteins. It is given a part symbol using `gcc:part` because the output language will not typically permit the use of URIs as identifiers, so this symbol via the implied `skos:prefLabel`[40] is what will appear instead. The protein produced by this coding sequence is also specified and linked using `gcc:protein`. It too is given a label using `skos:prefLabel` for the same reason, and its degradation rate is also specified with `gcc:proteinDegradationRate`. It is equally possible to specify the rates for transcription and translation in a similar manner though not shown here. In practice, rates are known primarily from experiment and this is an important reason to have accessible databases or repositories of part specifications.

Importantly, following the practice in our previous paper on rule annotation[39], a weak identity assertion is made with identifiers in external databases for the parts. This uses `bqbiol:is` instead of `owl:sameAs` because the strong replacement semantics (Leibniz' Law[42]) of the latter can yield unwanted inferences when terms are not used perfectly rigorously[37]. This weaker identify assertion permits the identification of the `:P0010` in the example with the identifier for the protein in the well-known UniProt[31] database.

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A More Complex Part Description

A more involved example demonstrating how an operator-promoter combination is encoded is shown in Figure 6. Here we have an operator with the rates for binding and unbinding of the transcription factor specified explicitly. If the operator is bound by the transcription factor, the neighbouring promoter is repressed — an RNA polymerase will not be able to bind. By contrast if the operator is unbound, the promoter will accept binding of RNA polymerase easily and frequently. The language supports an arbitrary amount of operator context for operators and promoters enabling the specification of complex regulatory structures such as combinatorial logic gates[43–45] and some forms of cooperative binding.

The transcription factor is specified by using `gcc:transcriptionFactor` to refer to the protein that will turn the operator on or off. Like `gcc:protein` for coding sequences, the term is unique to operators.

```
:R0040o a gcc:Operator;
  rdfs:label "TetR activated operator";
  gcc:part "R0040o";
  gcc:transcriptionFactor :P0040;
  gcc:transcriptionFactorBindingRate 0.01;
  gcc:transcriptionFactorUnbindingRate 0.01.

:R0040p a gcc:Promoter;
  rdfs:label "TetR repressible promoter";
  gcc:part "R0040p";
  gcc:rnapBindingRate
    [
      gcc:upstream ([ a rbmo:BoundState;
                      rbmo:stateOf :R0040o ]);
      gcc:value 7e-7
    ], [
      gcc:upstream ([ a rbmo:UnboundState;
                      rbmo:stateOf :R0040o ]);
      gcc:value 0.0007
    ].
```

Figure 6: An operator and promoter from the repressilator model. The binding rates for the promoter depend on the state of the adjacent operator.

The promoter comes next and it is the most complex part to specify. Because the rate for binding of RNA polymerase depends on the state of the operator, two rates must be specified. States of the nearby parts are specified using the `rbmo` vocabulary which makes available the full range of expressiveness for rule-based output languages. For generality, a

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list of parts, upstream or downstream on the DNA strand may be specified along with their states. This enables a promoter to be controlled by two or more operators. The rate itself in this case is given with `gcc:value` for each case.

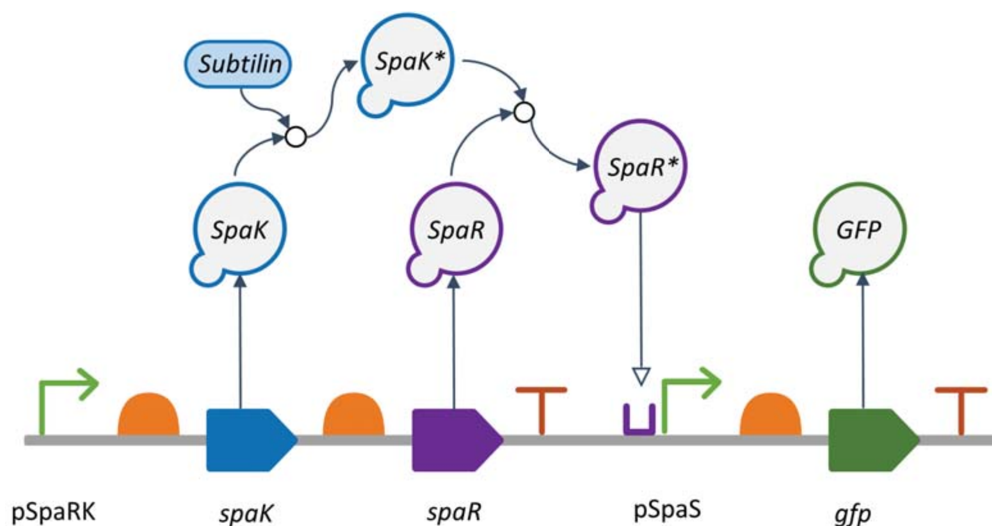
Host and Protein-Protein Interactions

The language can also support protein–protein interactions in a basic way. To see why these are useful, consider an example from the engineering of a bacterial communication system where the subtilin molecule is used to control population level dynamics. Cells have the receiver device [22,46] to sense the existence of subtilin, and the reporter device to initiate downstream cellular processes (Figures 7a and 7b). In the subtilin receiver, the interactions among the proteins produced by translation and the operator-promoters are mediated by a cascade reaction initiated by the subtilin molecule. Subtilin combines to phosphorylate the *SpaK* protein, which in turn phosphorylates the *SpaR* protein that finally binds to the promoter that controls the emission of a fluorescent green protein.

While the genetic circuit can straightforwardly be described similarly to the previous repressilator example, the protein–protein interactions cannot. We do not attempt here to model these interactions in the GCDL though a future extension could do so. Instead we simply allow for inclusion of the relevant program, as a file in the output language (in this case κ-language). It is possible to supply arbitrary code in the low-level language using the `gcc:include` term. This facility makes it feasible to represent such genetic circuits which depend strongly on the host environment in order to operate.

Protein Fusion

It is also worth noting that this example illustrates that in the high-level language it is immediately possible to represent devices that produce chains of proteins. This is known as protein fusion and is interesting for some applications[47]. A chain of proteins is produced by adding adjacent (and appropriate) coding sequences. It is enough to simply list the coding



(a) Diagram of the subtilin genetic circuit. The figure shows the multirelay phosphorylation, and hence the activation, of SpaR TFs to induce the downstream gene expression. As a result, GFP reporter proteins are produced in the presence of Subtilin molecules.

```
:m a rbmo:Model;
    dct:title "Subtilin Receiver Two-Component System";
    gcc:include <.../subtilin-host.ka>;
    gcc:linear (
        :pSpaRK :RBSa :spaK :RBSb :spaR :Ta
        :pSpaS  :RBSc :gfp :Tb
    ).
```

(b) Corresponding semantic model.

Figure 7: Representations of the Subtilin Receiver model.

sequences in the circuit; nothing else need be done.

Other Parts

The descriptions for the other kinds of biological parts, terminators, coding sequences, follow a similar pattern. There are terms for specifying the rates for the rules in which they participate, and a few specialised terms according to the function of the specific part. It is possible to find the available terms out by inspecting the `gcc` vocabulary included in the supplementary materials.

Output Representation

We now briefly consider the form of the output representation. By using different templates, the compiler can produce output in different languages. We focus on rule-based representations here and use the language of the KaSim simulator[14] for concrete illustration as it is widely adopted for stochastic simulation of rule-based models[48]. The rule-based modelling approach is merely outlined here and follows that used in Kappa BioBricks Framework (KBBF)[48] closely. We stress that though output as executable program in the KaSim language is demonstrated here, alternative rule-based representations like BioNetGen are equally possible as are descriptions in a language like SBOL as input to an experimental process in the laboratory. A more detailed account of the modelling methodology and corresponding output can be found in the supplementary materials.

The real work of modelling the transcription and translation machinery is done with sliding rules. Figure 8 shows how this works for the creation of a protein from a coding sequence. This is our first example of a rule where though the adjacent part figures explicitly in the rule, its *type* does not. It is sufficient to know that it is a piece of RNA. In this case,

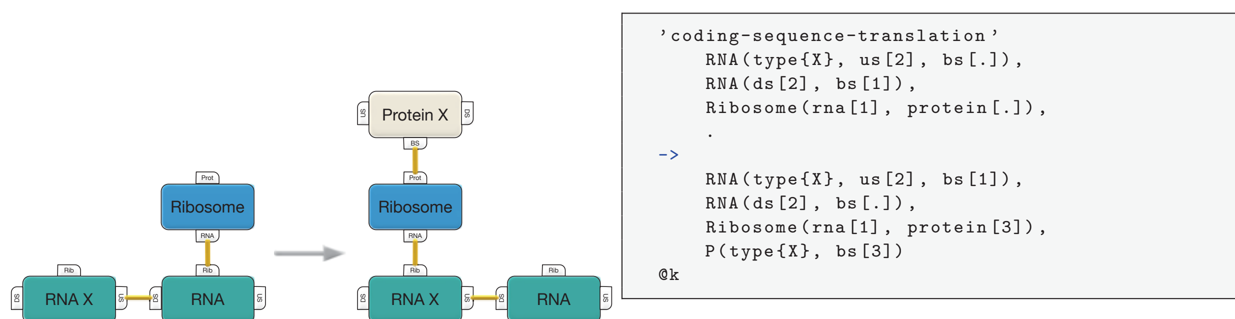


Figure 8: Translation of the RNA segment corresponding to a coding sequence to produce a protein.

two pieces of RNA are involved, the part that is central to this rule corresponds to the coding sequence for X. It is adjacent to another piece of RNA, and the ribosome slides from one to the other (to the left, where sliding on DNA happens, as we will see next, to the right) and in the process, emits a protein of type X.

Genetic Circuit Compiler

Having described the GCDL in some detail, we now briefly sketch our implementation of the compiler. Many compiler implementations are possible; ours innovatively combines the logical inference that is native to the semantic web with the use of templates to generate the target program. The templates define standard models for each type of part in a given output language. Different output languages or model granularities are achieved by choosing a different set of templates. The overall information flow through the compiler is illustrated in Figure 1.

Our strategy is to first gather all the input statements and background facts that are asserted by the various vocabularies in use. In the first inference step, standard RDF rules are used to make available consequent facts that will be needed to produce the ultimate result. The result is a program in a language such as κ and not RDF, and which uses local variable names and not URIs, so the materialised facts are transformed into a suitable internal representation. Substitution into templates is done next, and finally the result is post-processed to derive any remaining program directives that are only knowable once the complete circuit is assembled.

It is interesting to consider that the entire compiler can be thought of as implementing a kind of inference quite different from what is commonly used with the Semantic Web. The consequent, the executable model, is in a different language from the antecedent, the declarative description. Through the use of embedding annotations, however, the original model is nevertheless carried through to the output, and is unambiguously recoverable. There is thus an arrow from the space of declarative models in RDF to the space of annotated executable models. There is an arrow in the other direction that forgets the executable part and retains the declarative part. In an important sense, the two representations contain the same information, only that the executable model has more materialised detail in order that it may be run.

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Semantic Inference

The input from the user is the model description in the high-level language as described above. This description uses terms from, and makes reference to the `gcc` and `rbmo` vocabularies. The *meaning* of these terms, in the context of deriving an equivalent version of the program in the low-level language, is given by the companion inference rules. This is a somewhat subtle concept so let us illustrate what it means. Consider the statement,

```
:R0040a a gcc:Operator.
```

This statement gives the type of `:R0040a` as `gcc:Operator`.

The implications of this statement allows to identify the correct template to use for this part, found from information provided by the `gcc` vocabulary. Indeed, as a background fact, we have,

```
gcc:Operator gcc:kappaTemplate rbmt:operator.ka.
```

or in other words that an `gcc:Operator` corresponds to the template `rbmt:operator.ka`. We also have an inference rule, provided with the `gcc` vocabulary that says,

```
{ ?part a [ gcc:kappaTemplate ?template ] } => { ?part gcc:kappaTemplate ?template }.
```

In the Notation 3[32,33] language this means that, "for all `?part`s that has a type that corresponds to a kappa `?template`, that `?part` itself corresponds to that `?template`". Alternatively,

$$\texttt{type}(p, x) \wedge \texttt{kappa}(x, t) \rightarrow \texttt{kappa}(p, t)$$

It would have been perfectly possible to explicitly write what template should be used for each part in the high-level model description. That is not desirable because it would leak implementation details of the compiler into what ought to be an implementation-independent

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declarative description.

The above rule, and others like it serve to elaborate the high-level description into a more detailed version suitable for the next stage of the compiler and relieve the user of the need to supply the extra details. All implications that can be drawn under the `rdfs` inference rules and the `gcc` specific rules are drawn and become part of the in the in-memory RDF storage as the transitive closure of the rules (given the background facts and the provided model facts).

Internal Representation

The output of the first stage of the compiler contains all the information necessary to completely describe the output, but it is not in a convenient form for providing to the template rendering engine. Our implementation choice for the compiler is the Jinja2[49] rendering engine. This means that the appropriate data-structure is a dictionary or associative list that can be processed natively by these tools without need of external library. The required internal representation is built up by querying the in-memory RDF storage for the specific information required by the templates.

Our implementation does not require modification when new terms are added to the vocabulary and templates. To add support for a new kind of part it is necessary to write a new template for it and possibly add some terms to the vocabulary but does not require changing the compiler software itself. What makes this possible are the inference rules described in the previous section. The queries on the RDF storage that produce the internal representation are posed in terms of the *consequents* of the inference rules rather than the specific form of input.

Template Substitution

The templates that produce the bulk of the low-level output are written in the well-known Jinja2 language. This language is commonly used for the server-side generation of web pages.

KaSim or BNGL programs are not web pages but they are text documents and Jinja2 is well suited to generating them. It has a notion of inclusion and inheritance that is useful for handling the variations among the different kinds of parts, which typically differ in the rules for one or two of the interactions in which they participate with the others being identical. We provide a total of 15 templates for KaSim, of which there are top-level templates for each of the five distinct types of biological part defined in the `gcc` vocabulary as well as a generic part template, five templates implementing functionality shared among parts, and five consisting of supporting boilerplate required by KaSim.

A full description of the facilities provided by Jinja2 is beyond the scope of this paper, but a flavour is given in Figure 9 which shows an example of a template for a generic part (not having specific functionality like a promoter or operator might) demonstrating substitution of the `name` variable derived from annotation, and include statements referencing several other templates, one of which is reproduced and shows the KaSim code that is produced.

We use specific terms for defining the rates for the rules in which biological parts are involved, and a few other terms according to the function of the biological part of interest. It is possible to find the available terms out by inspecting the `gcc` vocabulary provided in the supplementary materials..

A fragment of the `gcc` vocabulary is reproduced in Figure 10. Though this exposes some implementation detail, it is useful to understand the relationships between the various terms used to describe models. This is also important when supplying customised templates.

There are `gcc:Tokens`, so named because they correspond to tokens in the low-level language that are replaced. Each must have a preferred label that gives the literal token. In cases where there exists a sensible default value, this is given with `gcc:default`. The purpose of these statements is to act as a bridge between the fully materialised RDF representation of the model and the templates that require substitution of locally meaningful names.

For each kind of part (such as the `gcc:Operator` in the example in Figure 10), there are two main annotations that are necessary. For each machine-readable low-level language,

```
## Auto-generated generic part {{ name }}
{% include "header.ka" %}
{% import "context.ka" as context with context %}
{% import "meta.ka" as meta with context %}

{% include "transcription_elongation.ka" %}
{% include "transcription_termination.ka" %}
{% include "translation_chain.ka" %}
{% include "translation_elongation.ka" %}
{% include "translation_termination.ka" %}
{% include "host_maintenance.ka" %}
```

```
{% set rule = "%s-translation-chain" % name %}
//
//^ :{{ rule }} a rbmo:Rule;
//^   bqbiol:isVersionOf go:GO:0006415;
{{ meta.rule() }}{# #}
//^   rdfs:label "{{ name }} formation of \
//^               translational chains, due to \
//^               gene fusion or leakiness of \
//^               stop codons".
// {{ name }} formation of translational chains,
// due to gene fusion or leakiness of stop codons
//
'{{ rule }}' \
    RNA(ds[2], bs[1]), \
    Ribosome(rna[1], protein[3]), \
    RNA(type{{ curly(name) }}, us[2], bs[.]), \
    Protein(ds[.], bs[3]), . \
-> \
    RNA(ds[2], bs[.]), \
    Ribosome(rna[1], protein[3]), \
    RNA(type~{{ name }}, us[2], bs[1]), \
    P(ds[4], bs[.]), \
    P(type{{ curly(name) }}, us[4], bs[3]) \
    @{{ translationElongationRate }}
```

Figure 9: Template examples. On top is the template for a generic part, and it references several other templates, one of which, `translation_chain.ka`, is reproduced on bottom.

a template is specified. The `gcc:tokens` annotations give the tokens that are pertinent to this kind of part. These must be specified in the high-level model or allowed to take on their default values. In addition to documenting the requirements of the templates for each kind of part, these statements are, "operationalised" and used by the compiler. They can equally well be used to check that a supplied high-level model is sufficiently complete and well-formed to produce an output program.

```
gcc:transcriptionFactor a gcc:Token;
    skos:prefLabel "transcriptionFactor".
gcc:transcriptionFactorBindingRate a gcc:Token;
    skos:prefLabel "transcriptionFactorBindingRate";
    gcc:default 1.0.
gcc:transcriptionFactorUnbindingRate a gcc:Token;
    skos:prefLabel "transcriptionFactorUnbindingRate";
    gcc:default 1.0.

gcc:Operator rdfs:subClassOf gcc:Part;
    gcc:kappaTemplate rbmt:operator.ka;
    gcc:bnglTemplate rbmt:operator.bngl;
    gcc:tokens
        gcc:transcriptionFactor,
        gcc:transcriptionFactorBindingRate,
        gcc:transcriptionFactorUnbindingRate.
```

Figure 10: The specification in the `gcc` vocabulary of an `gcc:Operator` and associated terms.

Derivation of Declarations

The KaSim language requires forward declaration of the type signatures of agents. This is by design[50] so that the simulator can check that agents are correctly used where they appear in patterns in the rules. While this design choice can help a modeller that is writing a simulation program in the low-level language by hand, to assist in finding mistakes and typographical errors, it is not possible to know *a priori* what these declarations should be in the present context. The correct declarations for `DNA`, `RNA` and `Protein` depend on the complete set of parts that make up the model so their correct declarations cannot be in any template for an individual part.

To solve this issue, the compiler implements a post-processing step. The rules that are produced by instantiating the templates for each part are concatenated together with any explicitly supplied rules and then the whole is parsed. The use of each agent in each rule in this rule-set is assumed to be correct by construction. From there a declaration that covers each use of each agent is built up.

Initialisation

At this final stage of the compiler, all rules are present, both supplied by the user for the host environment and implied by the parts that form the genetic circuits and all declarations

are also present. What is missing is the statement that creates an initial copy of the DNA sequence itself, which each upstream–downstream bond present. This information is, of course, available in the definition of the circuit, and so an appropriate `%init` statement, creating a single instance of the DNA sequence with correct linkages between the agent-parts is produced and added to the output. The low-level program is finally complete and ready to be executed.

Discussion

We have presented a language, the GCDL for describing genetic circuits and our compiler for generating simulation executables from it. We have made the case that the succinctness of the GCDL affords the user the benefit of describing the salient aspects of these circuits free of extraneous detail, that this reduces the potential for user error inherent in detailed coding of molecular interactions, and that this approach also affords flexibility in choosing the simulation or experimental methodology for the model. We have further developed the argument that modularity in modelling of genetic circuits has similar benefits of modularity in high-level programming languages, namely encapsulation and clarity. We now consider some of the limitations and benefits of our design choices and explore some areas ripe for future research.

It is important to understand the correctness and verification properties of the compiler and the GCDL. The GCDL is an RDF-based language and models are typically written in Turtle. The syntax[11,38] on a concrete level is well defined and models that are badly formed will be rejected. The standard templates are documented in machine-readable form in the GCDL vocabulary. Annotations that are required for a given part type also cause the model to be rejected by the compiler if they are not present. But the compiler does not perform verification in terms of how the parts are composed. Users are free to choose any DNA parts and in any order. For example, a model that includes a coding sequence part without

preceding promoter and ribosome binding site parts is allowed, though and the resulting model would emit no protein agents and perhaps not be very interesting. Verifying whether a given circuit expressed in the GCDL is accepted by a parts grammar[7,51] verification of part sequences is out of scope for the compiler but could be the subject of future work.

The expressive power afforded by the design choice of modularity — fixing the level of abstraction for a model — comes at a cost. Biological parts are considered as atomic units. While it is straightforward to model complex mechanisms like combinatorial logic operators and cooperative binding it is not straightforward to mix models in terms of the part abstraction with models of the underlying substrate. Phenomena that inherently involve the physical or chemical structure of the DNA molecule or the shape of a protein cannot be modelled directly and we are restricted to simply asserting that they occur or not at some rate. Similarly, while parts which share nucleotide sequences and may overlap can be marked as such with the `gcc:overlaps` term, this has no effect on the modelling. If the fact of parts overlapping is significant in the behaviour of the circuit, those parts are not modular and that would break the abstraction barrier. Such an annotation can, however, be used when selecting parts for assembly *in vitro*. Parts for which overlap is functionally significant can also be treated as an atomic unit with a suitable template. The modelling abstraction, once chosen, is fixed. This is by design, in order that models so expressed remain tractable.

Similar reasoning applies to optimisation of DNA sequences. This is not our focus in the present work. Here, our main goal is to capture the dynamics of genetic circuit designs and to automate the process of model generation. Hence, deriving final DNA sequences encoding the behaviours captured in models is not our focus, and related research can indeed be incorporated in the future[52]. Because the language is based on RDF, custom user based data can be stored as annotations[39] to facilitate later optimisation.

We do, however, envision optimisation of circuits at the level of abstraction that we have chosen, and derivation of circuits to a given specification. A method for doing this, which we only sketch here, is to define a suitable fitness or distance measure on the output of

simulations with respect to the desired specification. A starting candidate circuit is chosen, constructed from a given library of parts, and measured. Parts of the circuit are swapped, added or removed at random, subject to the constraint that the circuit remains well formed according to an operon grammar[7,51] and the new circuit is measured with respect to the specification. If the result is better than the previous circuit, the change is accepted, and the process is repeated until a locally optimal solution is found. This evolutionary algorithm approach is in contrast to the approach of assembling all possible circuits *in vitro* seen elsewhere[53–56] and is likely to be less efficient in cases where the desired behaviour of the circuit can be measured simply, such as by detecting the production of a fluorescent protein. However for cases that may be more difficult to measure *in vitro* such as oscillations or more complex outputs it can be more straightforward to measure the output and compare to the specification when done *in silico*.

Currently, the templates that we have supplied only handle single stranded genetic constructs. Parts are composed using upstream and downstream bonds to create chains of DNA sequences, and our framework currently does not consider whether the other strand is free or not regarding the elongation RNAP or the binding of molecules and so on. One reason why we have chosen to support the single-stranded case first is simplicity. Another is that databases of models for double-stranded parts are not available. Adding support for this in templates, and developing a library of suitable parts is another topic for future research.

Here, we presented the application of rule-based models and Semantic Web technologies to automate the design of genetic circuits. Representations of cellular activities were captured using modular rules to support scalability of designs. The automation process is facilitated by the GCDL high level language, which is built upon the Semantic Web and is used to describe genetic circuits. Furthermore, we presented a compiler that generates rule-based executable models from the high-level description. The implementation of the compiler is notable in its use of semantic inference and the language is sophisticated enough to support several classes of complex regulatory mechanisms. Despite the expressive power afforded by

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this approach, the language maintains a succinctness and simplicity that we hope will be a boon to those modelling genetic circuits in silico. The implicit modularity in our rule-based approach and the high-level language presented will be beneficial to synthetic biologists to model complex regulatory relationships through the use of widely adopted standards.

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Conflicts of Interest The authors have no competing interests.

Resources

Rule-Based Modelling Ontology	`http://purl.org/rbm/rbmo`
Genetic Circuit Description Language	`http://purl.org/rbm/comp`
Rule-Based Modelling Examples	`https://github.com/rulebased/rbmo`
Genetic Circuit Compiler Software	`https://github.com/rulebased/composition`
Kappa BioBricks Framework Software	`https://github.com/sstucki/lms-kappa`

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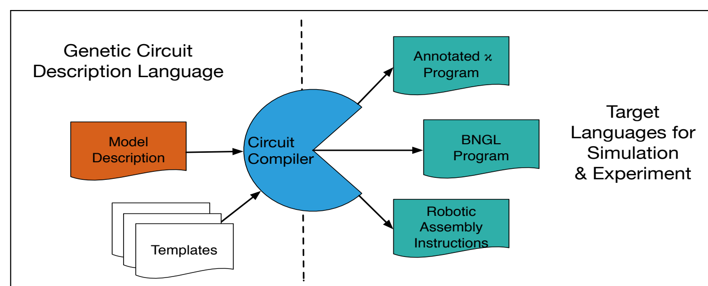

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