Spatial Kappa Simulator User Guide v2.0.1

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July 11, 2012

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Chapter 1

Spatial Kappa simulator User Guide

1.1 Obtaining the simulator

The simulator is available from GitHub as the source Eclipse project, or as a single executable jar file. Both are available at https://github.com/donal-s/SpatialKappa/downloads.

1.2 Starting the simulator

1.2.1 Running the executable jar

The simulator can be started by running the executable jar file: java -jar SpatialKappa-v2.0.1.jar

Double clicking the jar file usually works too.

1.2.2 Running from the Eclipse project

The main class of the simulator is org.demonsoft.spatialkappa.ui.SpatialKappaSimulator

Running as a Java Application will bring up the simulator.

1.3 Using the simulator

The initial screen appears as figure 1.1.

The toolbar options are:

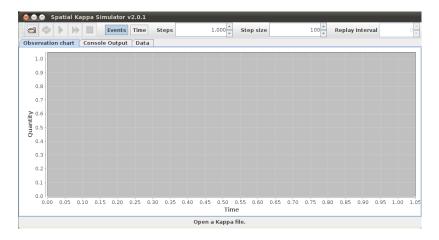
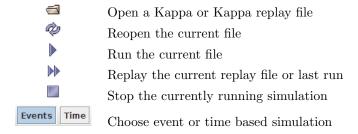


Figure 1.1: Initial view



1.3.1 Opening a Kappa or Kappa replay file

Select the 'Open' button on the toolbar and select the file to open. The current implementation expects Kappa source files to have the suffix .ka and Kappa replay files (discussed later) to have the suffix .kareplay. If the file is parsed successfully, a summary of the Kappa model is displayed in the 'Data' pane (see figure 1.2).

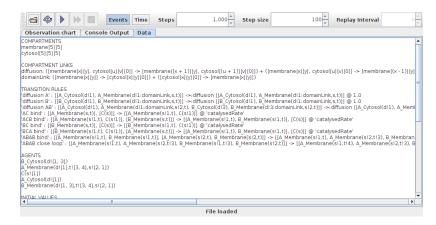


Figure 1.2: Data pane showing loaded Kappa model

Any errors in reading the Kappa file are shown in the 'Console Output' pane. The currently open Kappa file can be refreshed from disk by selecting the 'Reopen' button. Useful when editing the Kappa model.

1.3.2 Running a simulation

With a successfully opened Kappa model, one can run a simulation by selecting the 'Run' button. Simulation parameters can be set on the toolbar before running. There is the option to do an event or a time based simulation. For an event based simulation, the number of steps for the simulation (i.e. data points on the time series chart), and the number of finite rate events per step can be set. Equivalent options for time based simulation can also be set.

The simulation can be halted at any point by selecting the 'Stop' button. Note that complex simulations may take some time to start up while data structures are being generated.

A replay file of the simulation is created in the same directory as the Kappa model. The file format is similar to that produced by KaSim.

1.3.3 Running a simulation replay

As the simulation runs, the state of the simulation observables are logged to disk in a replay file after every step. Once the simulation is complete, this replay file can be rerun by selecting the 'Replay' button. The 'Replay Interval' field allows a delay (in ms) to be added between each step.

The file format is similar to that produced by KaSim. Renaming KaSim data output to have the suffix .kareplay will allow KaSim output to also be visualised using this tool.

1.4 Time series chart

While the raw data produced from simulations is useful, visualisation of the data is important. There is a simple visualisation panel in the simulator. This is dynamically updated as the simulation runs to give the user an idea of how the simulation is progressing. It is however basic in comparison to some of the commercial simulation data visualisation tools available.

The time series chart is similar to the standard Gnuplot output from KaSim. It is a line graph showing observable quantity against time for all observable definitions in the model. The excellent JFreeChart (Gilbert et al., 2010) library was used for generating the charting component. The chart has formatting and save capability, and is zoomable.

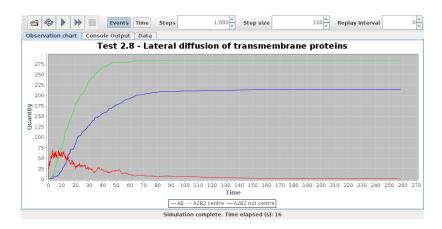


Figure 1.3: Sample time series chart output

Chapter 2

Kappa Language Extensions

Note on terminology. In the following 'voxel' means a subunit of a defined compartment. Species is mostly used to refer to a particular agent or complex in the model.

2.1 Existing Kappa language

The current Kappa language consists of the following constructs, as described in the KaSim documentation (Feret and Krivine, 2012). A more formal description of the language grammar (including the spatial extensions) is given in Appendix A

Comments:

```
# This is a comment
```

Agents:

```
AgentName ()
```

AgentName(stateWithValue~stateValue, stateWithNoValue, unboundSite, boundSite!1)

Non-spatial transition rules:

```
'state change' A(state old) -> A(state new) @ 0.1
'binding' A(bindsite),B(bindsite) -> A(bindsite!1),B(bindsite!1) @ 0.1
'unbinding' A(bindsite!1),B(bindsite!1) -> A(bindsite),B(bindsite) @ 0.1
'creation' -> A() @ 0.1
'degradation' A() -> @ 0.1
A(state old) -> A(state new) @ 0.1 # Unnamed transform rule
```

Initial species values:

```
%init 1000 A(state~old),C() # 1000 of each of A(..) and C() %init 2000 A(bindsite!1),B(bindsite!1) # 2000 of the bound complex A(..),B(..)
```

Observables and named variables: Variables can be referenced in perturbation calculations, but do not show in outputs as observations.

```
%obs 'Label' A() # All agents A()
%obs A() # Unnamed observation, will default to 'A()' in outputs
%obs A(state~old) # All agents matching A(state~old)
%obs 'binding' # Activity of the transform rule named 'binding'
%var 'Named variable' B()
```

2.2 Concepts to encapsulate

The spatial Kappa language requires some encoding of the following concepts to be useful.

- A description of **compartments** and their **dimensions**. A model may have multiple compartments each containing reacting species. The dimensions are necessary to define the shape of a compartment, be it a single cell, a 1 dimensional linear array of voxels, a 2 dimensional grid of some form, or a 3 dimensional lattice structure. Relative differences in size of different compartments can be specified, e.g. the reacting volume of a nucleus relative to the surrounding cytosol. Relative differences in shape can also be specified, for example the thin layer of cytosol next to the inner surface of the plasma membrane versus the rest of the cytosol.
- A description of **channels**, both intra-compartment and inter-compartment. The intra-compartment channel specification should be rich enough to allow description of multiple structures, e.g. in 1D linear arrays or circles, in 2D square or hexagonal meshes, cylinders or tori, in 3D cubes, filled cylinders, spheres, etc.
- A means of **locating species** within compartments, e.g. all DNA would reside within the nucleus, or cell receptors would be limited to the plasma membrane. Note that a multi-agent species need not be confined to a single voxel, but may span neighbouring, connected voxels.
- A means of **locating transition rules** within compartments, e.g. DNA transcription is isolated to the nucleus. The language should also allow the same transform rule to be specified with different rates depending on the location of the reacting species.
- A description of the **transport** (active or diffusive) of species within a compartment or between compartments along previously described channel structures. The rates of transport should be general to all species, or species specific. Note that transport of multi-voxel species should also be possible.

Additional concepts

- Granularity within compartments. It would be useful to be able to specify locations at the level of compartments or single voxels within a compartment. This would allow the model to represent, for example, a signal cascade being initiated as one point in the cytosol, and the resulting signal molecules being diffused through the cytosol.
- Backwards compatibility with basic Kappa. Given the quantity of existing models, the extended language should allow the existing models to work as before without modification. The user should have the choice of not using the spatial aspects of the extended language with no rework penalty.

2.3 New language constructs

A full BNF description of the extended Kappa grammar is given in Appendix A. The new constructs are identifiable as new rule types (%channel and %compartment), and location or channel identifiers in existing rule types prefixed with ':'.

2.3.1 Compartments and voxels

'%compartment:' id ('[' INT ']')*

Compartments are defined as single voxels or regular multidimensional arrays of voxels as follows

```
For example
%compartment: SingleCell
%compartment: 1dArray' [10]
                                 # 10 voxels in size
%compartment: 2dArray' [10][5]
                                 # 10x5 voxels in size
%compartment: 3dArray' [10][5][4] # 10x5x4 voxels in size
   Compartments or individual voxels within a compartment can be referenced
using the following location syntax
':' id ( '[' cellIndexExpr ']' )*
where
cellIndexExpr :
 cellIndexAtom operator_cell_index cellIndexAtom
 | cellIndexAtom
cellIndexAtom :
  '(' cellIndexExpr ')'
 | INT
 lid
operator_cell_index :
  ·+· | ·-· | ·*· | ·/· | ·%·
For example
:myCompartment
                                   # the compartment as a whole
:myCompartment [0][0][0]
                                   # the first voxel in a 3d array compartment
:myCompartment [4]
                                   # the fifth voxel in a 1d linear arrays
:myCompartment [x][y][z]
                                   # variable name usage described in
:myCompartment [x*2][y -1][z+(x*3)] #
                                       channel section below
```

In all situations where locations are used, it is only legal to refer to the compartment as a whole by omitting the cell indices, or refer to a single voxel, by fully defining the correct number of cell indices to match the dimensions of the compartment. Also, variable names are only permitted in locations within channel definitions, described below.

2.3.2 Channels

The structure of a compartment is further defined by how voxels within the compartment are linked to each other and to connected compartments. The channels are then used in defining both static links between agents, and movement of agents through the geometry of the model. These channels are defined as follows

```
'%channel:' id channel
| '%channel:' id '(' channel ')' ('+' '(' channel ')')*
where
channel:
  source=locations '->' target=locations
locations :
  location (',' location)*
Where location is as described above. For example
%compartment: 2dArray [10][200]
                                   # 10x200 voxels in size
# Link all voxels to their horizontally adjacent neighbours
# Link all voxels to their vertically adjacent neighbours
# Wrap around the voxels on the left and right edges to create a cylinder
# Wrap around the voxels on the top and bottom edges to create a torus
%channel: meshlinks \
    (:2dArray[x][y] -> :2dArray[x+1][y]) + (:2dArray[x][y] -> :2dArray[x -1][y]) +\
    (:2dArray[x][y] -> :2dArray[x][y+1]) + (:2dArray[x][y] -> :2dArray[x][y -1]) +\
    (:2dArray[0][y] -> :2dArray[9][y]) + (:2dArray[9][y] -> :2dArray[0][y]) +\
    (:2dArray[x][0] -> :2dArray[x][199]) + (:2dArray[x][199] -> :2dArray[x][0])
```

The above code defines a thin torus composed of a 2d mesh.

Locations on the left hand side of the channel definitions above may contain either constant values or single variable names, not complex expressions. The variable names are used to define the dimensions which will be iterated through to produce links. Locations on the right hand side allow constant values or complex expressions involving the variables defined on the left hand side of the expression. It is invalid for the right hand expression to use variables not defined on the left. If setting the values of variables references valid voxels on both the left and right, then those voxels are deemed to be linked. References which refer to voxels outside the dimensions of the compartment are ignored, and no link is created. The references in a channel expression can refer to the same compartment, or two different compartments. The modulus operator % is useful in defining regular, repeating linkage patterns within the compartment, for example the 2D hexagonal mesh described in Appendix B.1.

Channels can make use of multiple source voxels simultaneously. For example if a model was to represent the movement of transmembrane proteins laterally along the surface of a membrane, then the channel used to describe the lateral motion would need to include simultaneous movement in two compartments (cytosol and membrane). This is represented as follows:

```
%compartment: membrane [5][5]
%compartment: cytosol [5][5][5]
%channel: diffusion \
    (:membrane [x][y], :cytosol [u][v][0] -> :membrane [x+1][y], :cytosol [u+1][v][0]) + \
    (:membrane [x][y], :cytosol [u][v][0] -> :membrane [x -1][y], :cytosol [u -1][v][0]) + \
    (:membrane [x][y], :cytosol [u][v][0] -> :membrane [x][y+1], :cytosol [u][v+1][0]) + \
    (:membrane [x][y], :cytosol [u][v][0] -> :membrane [x][y -1], :cytosol [u][v -1][0])
```

Here, variables x,y represent locations in the membrane, and u,v represent locations in the cytosol. The definition updates the locations in these two compartments in unison.

Further examples of compartment and channel specifications for common structures are given in Appendix B.1.

2.3.3 Locating agents

The definitions above can now be used to locate species within the model. Any rule that accepts a definition of agents (i.e. transition, var, obs, init rules), now allows these agents to be located. For each group of agents, a prefixed location constrains the agents to that location. For example

When locations are specified both for agent groups and individual agents, the individual agent location takes precedence. This allows for concise definition of agent groups where all but one of the agents in the group share a location.

2.3.4 Agent links

Agents in neighbouring voxels linked by a defined channel can be linked together. This is an extension of the basic Kappa link syntax to name the channel used to link the agents. For example

The above describes a model where the species A-B exists in two compartments, B in the cytosol and A embedded in the membrane. When specifying agent links using channels, only one end of the link needs to specify the channel. If a link does not specify the channel, it is assumed that both agents party to the link exist in the same voxel.

Links including channels can be created or broken in the same way as basic Kappa links in transition rules.

2.3.5 Species movement

Species can move along defined channels. Species movement is described using the ->: operator.

```
(source=location)? '->:' channelName=id (target=location)?
| (a=agentGroup)? '->:' channelName=id (b=agentGroup)?
```

Movement transition rules can either constrain the movement by species chosen, or by source location. For example

```
%compartment: membrane [5][5]
%channel: diffusion \ (:membrane [x][y] -> :membrane [x+1][y]) + (:membrane [x][y] -> :membrane [x - 1][y]) + \
    (:membrane [x][y] -> :membrane [x][y+1]) + (:membrane [x][y] -> :membrane [x][y - 1])
'diffusion A' A(s,t) ->:diffusion A(s,t) @ 1.0
'diffusion B' B(s,t) ->:diffusion B(s,t) @ 1.0
'diffusion AB' A(s!1,t), B(s!1,t) ->:diffusion A(s!1,t), B(s!1,t) @ 0.5
'diffusion all' ->:diffusion @ 1.0 # All species located in a single voxel will match this rule
   To describe movement of species which span more than one voxel, use the
multi agent channel definition above. For example
%compartment: membrane [5][5]
%compartment: cytosol [5][5][5]
%channel: diffusion \setminus
    (:membrane [x][y], :cytosol [u][v][0] \rightarrow :membrane [x+1][y], :cytosol [u+1][v][0]) + \
     (:membrane \ [x] \ [y] \ , \ :cytosol \ [u] \ [v] \ [0] \ \rightarrow \ :membrane \ [x \ -1] \ [y] \ , \ :cytosol \ [u \ -1] \ [v] \ [0]) \ + \ \\ 
     \hbox{(:membrane $[x][y]$, :cytosol $[u][v][0]$ $\rightarrow$ :membrane $[x][y+1]$, :cytosol $[u][v+1][0]$) $+$ $$ $$ $$
    (:membrane [x][y], :cytosol [u][v][0] -> :membrane [x][y -1], :cytosol [u][v -1][0])
%channel: domainLink \setminus
    (:membrane [x][y] -> :cytosol [x][y][0]) + (:cytosol [x][y][0] -> :membrane [x][y])
'diffusion A' A_m:membrane(s,t,d!1:domainLink),A_c(d!1) ->:diffusion \
               A_m:membrane(s,t,d!1:domainLink),A_c(d!1) @ 1.0
B_m:membrane(s,t,d!1:domainLink),B_c(d!1) @ 1.0
'diffusion AB' A_m:membrane(s!2,t,d!1:domainLink),A_c(d!1), \
               B_m:membrane(s!2,t,d!3:domainLink), B_c(d!3) \rightarrow:diffusion \setminus
                A_m:membrane(s!2,t,d!1:domainLink),A_c(d!1), \
               B_m:membrane(s!2,t,d!3:domainLink),B_c(d!3) @ 0.5
```

For example models demonstrating the use of the language extensions, refer to Appendix B.

Appendix A

Spatial Kappa Grammar

The following is a cut down version of the Antlr grammar used in the Kappa simulator. The syntax has been trimmed for readability, as the original Antlr grammar has artificial constructs for dealing with left recursion, etc. It is read basically as BNF notation with assignments (variable=bnfConstruct). The existing basic Kappa grammar is shown in black, with the spatial constructs shown as blue.

```
prog :
  (line)*
line:
  agentDecl NEWLINE!
  | compartmentDecl NEWLINE!
  channelDecl NEWLINE!
  | ruleDecl NEWLINE!
  | initDecl NEWLINE!
  | plotDecl NEWLINE!
  | obsDecl NEWLINE!
  | varDecl NEWLINE!
  | modDecl NEWLINE!
  | COMMENT!
  | NEWLINE!
ruleDecl :
  label? transition rate
transition :
  (source=location)? CHANNEL_TRANSITION channelName=id (target=location)?
  (a=agentGroup)? CHANNEL_TRANSITION channelName=id (b=agentGroup)?
  | (a=agentGroup)? FORWARD_TRANSITION (b=agentGroup)?
agentGroup :
  location? agent (',' agent)*
  id (location)? ('(' (agentInterface (',' agentInterface)*)? ')')?
agentInterface :
  id state? link?
state :
```

```
,~, id
link :
    '!' INT (':' channelName=id)?
    '''    '''

  | '!' '_' (':' channelName=id)?
  , , ? ,
rate:
  '@' varAlgebraExpr
{\tt initDecl} :
  '%init:' (INT | label) agentGroup
agentDecl :
  '%agent:' agentName=id ('(' (agentDeclInterface (',' agentDeclInterface)*)?')')?
agentDeclInterface :
  id state*
compartmentDecl :
   '%compartment:' id ('[' INT ']')*
channelDecl :
   '%channel:' linkName=id channel
   | '%channel:' linkName=id '(' channel ')' ('+' '(' channel ')')*
channel:
  source=locations FORWARD_TRANSITION target=locations
locations :
  location (',' location)*
location :
  ':' sourceCompartment=id compartmentIndexExpr*
compartmentIndexExpr :
  '[' cellIndexExpr ']'
plotDecl :
  '%plot:' label
obsDecl :
  '%obs:' label? agentGroup
varDecl :
  '%var:' label varAlgebraExpr
   | '%var:' label agentGroup
varAlgebraExpr :
  a=varAlgebraMultExpr (op=operator_add b=varAlgebraMultExpr )*
varAlgebraMultExpr :
  a=varAlgebraExpExpr (op=operator_mult b=varAlgebraExpExpr )*
{\tt varAlgebraExpExpr} \ :
  a=varAlgebraAtom operator_exp b=varAlgebraExpExpr
  | a=varAlgebraAtom
varAlgebraAtom :
```

```
'(' varAlgebraExpr ')'
  | number
  | label
  | '[' 'inf' ']'
  ' '[' 'pi' ']'
  , , T, ,],
  | '[' 'E' ']'
  | operator_unary varAlgebraAtom
   \verb|| operator_binary_prefix a=varAlgebraAtom b=varAlgebraAtom \\
modDecl :
  '%mod:' booleanExpression 'do' effect until?
booleanExpression :
  a=booleanAtom (op=booleanOperator b=booleanAtom )*
{\tt boolean Operator} \ :
  '&&' | '||'
relationalOperator :
  '<' | '>' | '='
{\tt booleanAtom} \; : \;
  '(' booleanExpression ')'
  | '[' 'true' ']'
  | '[' 'false' ']'
  '[', 'not', ']' booleanAtom
  effect :
  '$SNAPSHOT'
  / '$STOP'
  ' '$ADD' varAlgebraExpr agentGroup
  | '$DEL' varAlgebraExpr agentGroup
  | label ':=' varAlgebraExpr
until:
  'until' booleanExpression
cellIndexExpr :
  a=cellIndexAtom operator_cell_index b=cellIndexAtom
  | a=cellIndexAtom
cellIndexAtom :
  '(' cellIndexExpr ')'
  | INT
  | id
  ALPHANUMERIC ( ALPHANUMERIC | '_' | '-' )*
label :
 LABEL
number :
  ( INT | FLOAT )
operator_cell_index :
  ( '+' | '*' | '-' | '/' | '%' )
```

```
operator_exp :
  | ,^,
operator_unary :
  '[' 'log' ']'
| '[' 'sin' ']'
| '[' 'cos' ']'
| '[' 'tan' ']'
  | '[' 'sqrt' ']'
operator_binary_prefix :
    '[' 'mod' ']'
operator_mult :
   '*' | '/'
{\tt operator\_add} \; : \;
   ·+ · | ·- ·
CHANNEL_TRANSITION :
  ·->: ·
{\tt FORWARD\_TRANSITION} \ :
  ,->,
INT :
  NUMERIC
FLOAT :
  NUMERIC '.' NUMERIC EXPONENT?
  | '.' NUMERIC EXPONENT?
  | NUMERIC EXPONENT
ALPHANUMERIC : ( NUMERIC | 'a'...'z' | 'A'...'Z' )+
NUMERIC :
  ('0'...'9')+
EXPONENT :
  ('e'|'E') ('+'|'-')? NUMERIC
LABEL :
  '\'' .* '\''
COMMENT :
  '#', ~( '\n', | '\r', )*
NEWLINE :
  '\r'? '\n' | '\r'
  ( ' ' | '\t' | '\\' NEWLINE )+
```

Appendix B

Spatial Kappa Examples

B.1 Spatial Kappa patterns

The following are generic shapes, with their equivalent Spatial Kappa representations. These are intended to be copied during model development.

B.1.1 1 dimensional patterns

Linear array

```
%compartment: array [n] # Replace n with length of array
%channel: intra-array (:array [x] -> :array [x+1]) + (:array [x] -> :array [x -1])

Circle
%compartment: circle [n] # Replace n with number of cells in circle
%channel: intra-circle \
    (:circle [x] -> :circle [x+1]) + (:circle [x] -> :circle [x -1]) + \
    (:circle [n-1] -> :circle [0]) + (:circle [0] -> :circle [n-1]) # Replace n-1 as above
```

B.1.2 2 dimensional surfaces

Rectangular mesh

There are 2 variants here, 4-way linked and 8-way linked.

%compartment: mesh [n][m] # Replace n and m with dimensions of mesh

```
# 4-way diffusion
%channel: intra-mesh \
    (:mesh [x][y] -> :mesh [x+1][y]) + (:mesh [x][y] -> :mesh [x -1][y]) + \
    (:mesh [x][y] -> :mesh [x][y+1]) + (:mesh [x][y] -> :mesh [x][y -1])

# or 8-way diffusion
%channel: intra-mesh \
    (:mesh [x][y] -> :mesh [x+1][y]) + (:mesh [x][y] -> :mesh [x -1][y]) + \
    (:mesh [x][y] -> :mesh [x][y+1]) + (:mesh [x][y] -> :mesh [x][y -1]) + \
    (:mesh [x][y] -> :mesh [x+1][y+1]) + (:mesh [x][y] -> :mesh [x -1][y-1]) + \
    (:mesh [x][y] -> :mesh [x+1][y-1]) + (:mesh [x][y] -> :mesh [x -1][y+1])
```

Hexagonal mesh

Again, 2 variants depending on what overall shape is required. The first form has a simpler representation of intra-compartment links, but the overall structure is rhomboid, whereas the second produces an overall rectangular shape at the expense of more complicated link statements.

The second variant demonstrates handling of alternate odd-even linkage depending on the column of the structure.

```
%compartment: mesh [n][m] # Replace n and m with dimensions of mesh
# Variant 1 - rhomboid mesh
%channel: intra-mesh \
    (:mesh [x][y] -> :mesh [x][y+1]) + (:mesh [x][y] -> :mesh [x][y -1]) + \
    (:mesh [x][y] -> :mesh [x+1][y]) + (:mesh [x][y] -> :mesh [x -1][y]) + \
    (:mesh [x][y] -> :mesh [x+1][y+1]) + (:mesh [x][y] -> :mesh [x -1][y-1])
# Variant 2 - rectangular mesh
%channel: intra-mesh \
    (:mesh [x][y] -> :mesh [x][y+1]) + (:mesh [x][y] -> :mesh [x][y -1]) + \
    (:mesh [x][y] -> :mesh [x+1][y]) + (:mesh [x][y] -> :mesh [x -1][y]) + \
    (:mesh [x][y] -> :mesh [x+1][(y+1)-(2*(x*2))])
# The above statement alternates [x+1][y+1] and [x+1][y-1] as x increases
```

Cylinder and torus

By connecting together the top and bottom edges of a mesh as described above, we get a cylinder. By also connecting together the left and right edges we get a torus.

```
%compartment: mesh [n][m] # Replace n and m with dimensions of mesh
# 4-way diffusion mesh
%channel: intra-mesh \
    (:mesh [x][y] \rightarrow :mesh [x+1][y]) + (:mesh [x][y] \rightarrow :mesh [x-1][y]) + 
    (:mesh [x][y] \rightarrow :mesh [x][y+1]) + (:mesh [x][y] \rightarrow :mesh [x][y-1])
%channel: intra-mesh mesh [x][y] \iff mesh [x+1][y]
%channel: intra-mesh mesh [x][y] <-> mesh [x][y+1]
# cylinder
%channel: intra-mesh \
    (:mesh [x][y] \rightarrow :mesh [x+1][y]) + (:mesh [x][y] \rightarrow :mesh [x-1][y]) + 
    (:mesh [x][y] \rightarrow :mesh [x][m-1]) + (:mesh [x][y] \rightarrow :mesh [x+1][0])
# Replace m-1 as above
# torus
%channel: intra-mesh \
    (:mesh [x][y] \rightarrow :mesh [x+1][y]) + (:mesh [x][y] \rightarrow :mesh [x -1][y]) + 
    (:mesh [x][y] \rightarrow :mesh [x][y+1]) + (:mesh [x][y] \rightarrow :mesh [x][y-1]) + 
    (:mesh [x][m -1] -> :mesh [x][0]) + (:mesh [x][0] -> :mesh [x][m -1]) +
    (:mesh [n -1][y] \rightarrow :mesh [0][y]) + (:mesh [0][y] \rightarrow :mesh [n -1][y])
# Replace n-1 as above
```

B.2 Sample Spatial Kappa models

The following are a couple of simple spatial kappa models to demonstrate the use of the language features.

B.2.1 2d diffusion model

This model shows simple diffusion from a point for three distinct species.

```
%agent: A()
%agent: B()
%agent: C()
%compartment: cytosol [30][30]
# 6-way diffusion
%channel: 6way \
    (:cytosol [x][y] -> :cytosol [x][y+1]) + (:cytosol [x][y] -> :cytosol [x][y -1]) + \
    (:cytosol [x][y] -> :cytosol [x+1][y]) + (:cytosol [x][y] -> :cytosol [x -1][y]) + \
    (:cytosol [x][y] \rightarrow :cytosol [x+1][(y+1)-(2*(x%2))]) + 
    (:cytosol [x][y] \rightarrow :cytosol [x -1][(y -1)+(2*((x -1)%2))])
'diffusion' ->:6way @ 0.4
%init: 10000 :cytosol[10][10] A
%init: 10000 :cytosol[10][14] B
%init: 10000 :cytosol[14][14] C
%obs: 'Red' A
%obs: 'Green' B
%obs: 'Blue' C
```

B.2.2 Bi-trivalent binding model

A variant of bi-trivalent binding test model (Yang et al., 2008). This spatial model allows unbound species to diffuse through the compartment, but bound species remain within a cell of the compartment.

```
%agent: A(a,b,bindings~0~1~2)
%agent: B(a,b,c)
%compartment: cytosol [20][20]
# 6-way diffusion
%channel: 6way
    (:cytosol [x][y] \rightarrow :cytosol [x][y+1]) + (:cytosol [x][y] \rightarrow :cytosol [x][y-1]) + 
     \hbox{(:cytosol $[x][y]$ $\rightarrow$ :cytosol $[x+1][y]$) $+$ (:cytosol $[x][y]$ $\rightarrow$ :cytosol $[x-1][y]$) $+$ $$ $$ $$
     (:cytosol [x][y] \rightarrow :cytosol [x+1][(y+1)-(2*(x%2))]) +
    (:cytosol [x][y] \rightarrow :cytosol [x -1][(y -1)+(2*((x -1)%2))])
'diffusion-A' A(bindings~0) ->:6way A(bindings~0) @ 0.1
'diffusion-B' B(a,b,c) ->:6way B(a,b,c) @ 1
A(a,b,bindings~0), B(a) -> A(a!1,b,bindings~1),B(a!1) @ 1
A(a,b,bindings~0), B(b) -> A(a!1,b,bindings~1),B(b!1) @ 1
A(a,b,bindings^0), B(c) \rightarrow A(a!1,b,bindings^1), B(c!1) @ 1
A(a,b,bindings~0), B(a) -> A(a,b!1,bindings~1),B(a!1) @ 1
A(a,b,bindings^0), B(b) \rightarrow A(a,b!1,bindings^1), B(b!1) @ 1
A(a,b,bindings^0), B(c) \rightarrow A(a,b!1,bindings^1), B(c!1) @ 1
A(a,b!_,bindings~1), B(a) -> A(a!1,b!_,bindings~2),B(a!1) @ 1
```

```
A(a,b!_,bindings~1), B(b) -> A(a!1,b!_,bindings~2),B(b!1) @ 1
A(a,b!_,bindings~1), B(c) -> A(a!1,b!_,bindings~2),B(c!1) @ 1
A(a!\_,b,bindings^1), B(a) \rightarrow A(a!\_,b!1,bindings^2), B(a!1) @ 1
A(a!_,b,bindings~1), B(b) -> A(a!_,b!1,bindings~2),B(b!1) @ 1
A(a!_,b,bindings~1), B(c) -> A(a!_,b!1,bindings~2),B(c!1) @ 1
A(a!1,b,bindings^1),B(a!1) \rightarrow A(a,b,bindings^0), B(a) @ 0.01
A(a!1,b,bindings~1),B(b!1) -> A(a,b,bindings~0), B(b) @ 0.01
A(a!1,b,bindings~1),B(c!1) -> A(a,b,bindings~0), B(c) @ 0.01
A(a,b!1,bindings~1),B(a!1) -> A(a,b,bindings~0), B(a) @ 0.01
A(a,b!1,bindings~1),B(b!1) -> A(a,b,bindings~0), B(b) @ 0.01
A(a,b!1,bindings~1),B(c!1) -> A(a,b,bindings~0), B(c) @ 0.01
A(a!1,b!\_,bindings^2),B(a!1) \rightarrow A(a,b!\_,bindings^1), B(a) @ 0.01
A(a!1,b!\_,bindings^2),B(b!1) \rightarrow A(a,b!\_,bindings^1),B(b) @ 0.01
A(a!1,b!\_,bindings^2),B(c!1) \rightarrow A(a,b!\_,bindings^1), B(c) @ 0.01
A(a!_,b!1,bindings~2),B(a!1) -> A(a!_,b,bindings~1), B(a) @ 0.01
A(a!_,b!1,bindings~2),B(b!1) -> A(a!_,b,bindings~1), B(b) @ 0.01
A(a!\_,b!1,bindings^2),B(c!1) \rightarrow A(a!\_,b,bindings^1), B(c) @ 0.01
```

%init: 600 A(a,b,bindings~0)

%init: 400 B(a,b,c)

%obs: 'Red' A(bindings~2) %obs: 'Green' A(bindings~1) %obs: 'Blue' A(bindings~0)

Bibliography

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