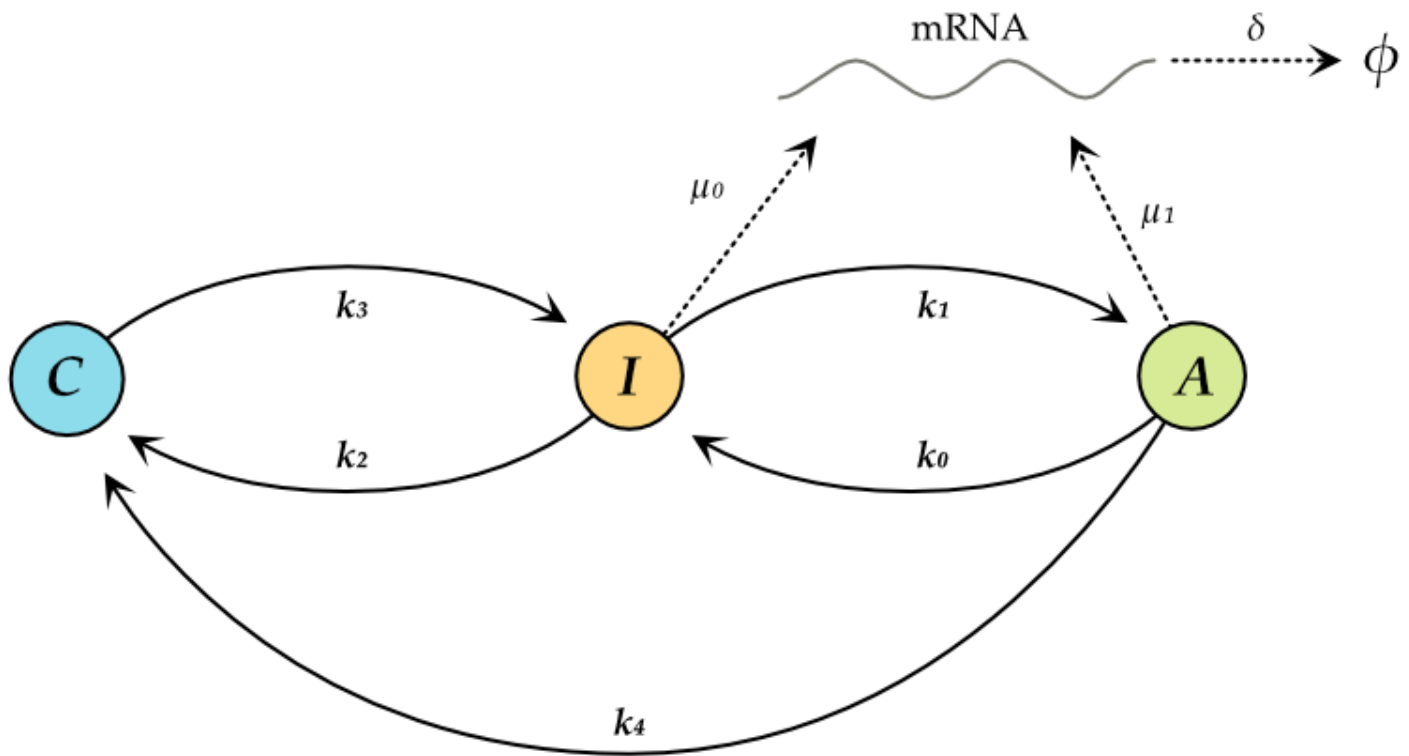


Three State Model

The functions in this file implement a three-state model for gene transcription that is summarised by the figure below



*The three-state model. Each locus can be in one of three states: Closed, Inactive or Active. Transcripts are produced from inactive loci at rate μ_0 per locus, from active loci at rate μ_1 and degrade at a rate δ per transcript.**

Computing probabilities

The main function of interest here is

```
matExpDistrib3( maxRNA, nLoci, mu, k, delta, times, eps )
```

whose arguments are:

maxRNA

the maximal number of transcripts that we expect to see at the longest times. If this is too small the estimates of the distribution will be poor.

nLoci

the number of potential sites of transcription. Typically it's two, but other values are possible. The number states for a cell is $(nLoc i + 2) * (nLoc i + 1) / 2$.

mu

a vector containing the rates of transcription in the form $[\mu_0, \mu_1]$. Both are measured in transcripts per minute per locus.

k

a vector containing the rates of locus-state transitions in the form $[k_0, k_1, k_2, k_3, k_4]$. All are measured in transitions per minute per locus.

times

a vector of times, which should be
* positive,
* measured in minutes,
* arranged in ascending order,

at which we want the distribution.

eps

an optional threshold for small probabilities whose default value, `eps = 0.000001` is set in the file `matExpDistributions.m`, Probabilities less than `eps` are set to zero.

Format of the output

The function `matExpDistributions3()` computes a probability for each of the

$$nModelStates = (maxRNA + 1) * (nLoc i + 2) * (nLoc i + 1) / 2$$

possible states of the model. A state, which describes what is happening in a single cell, is specified by by four non-negative whole numbers (`nc`, `ni`, `na`, `m`) where

nc is the number of loci in the closed state;

ni is the number of loci in the inactive state;

na is the number of loci in the active state;

m is the number of transcripts.

Thus a cell with `nLoc i` possible sites of transcription should always have `nc + ni + na = nLoc i`.

The output of `matExpDistrib3()` is a matrix with one row for each state and one column for each time in the input argument `times`. The way the states get mapped to rows of this matrix is determined by the function `stateToIdx3(ni, na, nLoc, m)` and is illustrated in the table below

The mapping between states (`nc, ni, na, m`) and row numbers in the output of `matExpDistrib3()`.

nc	ni	na	m	rowNum
2	0	0	0	1
1	1	0	0	2
0	2	0	0	3
1	0	1	0	4
0	1	1	0	5
0	0	2	0	6
2	0	0	1	7
1	1	0	1	8
0	2	0	1	9
1	0	1	1	10
0	1	1	1	11
0	0	2	1	12

Probability distribution over number of transcripts

All the states that involve the same number of transcripts are stored in consecutive rows of the output of `matExpDistrib3()`. This makes it easy to implement a convenience function, `marginalise()`, that computes a matrix of probabilities with one row for each of the $(\text{maxRNA} + 1)$ possible values of the number of transcripts and one column for each of the times. The m, j entry in this matrix is the probability of seeing $m-1$ transcripts at time t_j . A typical application is

```
stateProbMat = matExpDistrib3( maxRNA, nLoc, mu, k, delta, times )
transcriptCountProbMat = marginalise( stateProbMat, nLoc )
```