Universally valid reduction of multiscale stochastic biochemical systems using simple non-elementary propensities

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Abstract

Biochemical systems consist of numerous elementary reactions governed by the law of mass action. However, experimentally characterizing all the elementary reactions is nearly impossible. Thus, over a century, their deterministic models that typically contain rapid reversible bindings have been simplified with non-elementary reaction functions (e.g., Michaelis-Menten and Morrison equations). Although the non-elementary reaction functions are derived by applying the quasi-steady-state approximation (QSSA) to deterministic systems, they have also been widely used to derive propensities for stochastic simulations due to computational efficiency and simplicity. However, the validity condition for this heuristic approach has not been identified even for the reversible binding between molecules, such as protein-DNA, enzyme-substrate, and receptor-ligand, which is the basis for living cells. Here, we find that the non-elementary propensities based on the deterministic total QSSA can accurately capture the stochastic dynamics of the reversible binding in general. However, serious errors occur when reactant molecules with similar levels tightly bind, unlike deterministic systems. In that case, the non-elementary propensities distort the stochastic dynamics of a bistable switch in the cell cycle and an oscillator in the circadian clock. Accordingly, we derive alternative non-elementary propensities with the stochastic low-state QSSA, developed in this study. This provides a universally valid framework for simplifying multiscale stochastic biochemical systems with rapid reversible bindings, critical for efficient stochastic simulations of cell signaling and gene regulation. To facilitate the framework, we provide a user-friendly open-source computational package, ASSISTER, that automatically performs the present framework.

Introduction

To understand the complex dynamics of numerous molecular interactions in living cells, quantitative analysis using mathematical models is essential [1]. While elementary reactions in living cells can be modeled by the law of mass action, characterizing all their kinetics is challenging. Thus, over a century, the combined effect of a set of elementary reactions such as rapid reversible bindings has been described with non-elementary reaction functions (e.g., Michaelis-Menten and
Morrison equations) to simplify deterministic models \cite{2-7}. Since the early 2000s, these deterministically driven non-elementary reaction functions have also been widely used to derive propensity functions for stochastic simulations, which greatly reduces the computational cost \cite{8-33}. This heuristic approach for efficient stochastic simulations was believed to be valid as long as the non-elementary reaction functions are accurate in the deterministic sense. However, unfortunately, this was not the case \cite{33-40}. The reason for the discrepancy between the deterministic and stochastic simulations has been recently identified for some cases \cite{37-40}, but not for all \cite{41}. Currently, guidelines for this popular but heuristic method for efficient stochastic simulations with non-elementary propensity functions are absent.

The non-elementary reaction functions are mainly the result of the reduction of deterministic models with the following reversible binding reactions:

\begin{equation}
A + B \xrightleftharpoons[k_b]{k_f} C.
\end{equation}

The reversible binding between molecules, such as enzyme-substrate, receptor-ligand, and protein-DNA, is the first step for nearly all biological functions of living cells \cite{42}. However, rather than the reversible binding itself, its outcome is usually our major interest. For instance, rather than the binding between a transcription factor and DNA, we are interested in its outcome, the transcription. Furthermore, the transcription factor binding to DNA takes at most one second while transcription takes about 30 minutes in a mammalian gene \cite{43}, which causes stiffness in numerical simulations \cite{44}. Fortunately, such rapid reversible binding reactions can be eliminated from models using the property: the levels of the species (A, B, and C) regulated by the reversible binding more quickly equilibriate to their quasi-steady-states (QSSs) compared with the total levels of the bound and unbound species, which are not affected by the reversible binding. In deterministic models, their quasi-steady-state approximations (QSSAs), which are non-elementary reaction functions, can be obtained by finding the steady-state solution of the associated differential equation in terms of the total variables. Because the QSSAs are determined by the total variables, they are known as the "total" QSSA (tQSSA). After replacing the variables that represent the levels of A, B, and C with their tQSSAs, rapid reversible bindings have been successfully eliminated from various deterministic models describing enzyme catalysis, gene regulation, and cell cycle regulation \cite{5-7, 23, 45-51}. Note that adopting the total variables leads to timescale separation among variables, while the sole rapidity of the reversible binding reactions does not guarantee timescale separation between the original variables, A, B, and C \cite{7} (see Discussion for details).

In stochastic models, the QSSAs for the numbers of A, B, and C are their stationary average numbers (i.e., the first moment) conditioned on the total numbers of the bound and unbound species for uni- or bi-molecular reactions \cite{27-30} (see SI Appendix for details). These stochastic QSSAs can be obtained by finding the steady-state solution of the chemical master equation (CME). However, unlike the deterministic tQSSA, the stochastic QSSA has a complex form (Eq. (4)), which does not provide any intuition, and importantly, increases computational cost. Thus, its approximation has been derived with the deterministic tQSSA. This approximation, often referred to as the stochastic tQSSA (stQSSA) \cite{7, 31, 38, 39}, leads to non-elementary propensity functions for stochastic simulations using the Gillespie algorithm \cite{52}. In this way, the stochastic dynamics of various systems have been accurately captured with low computational cost \cite{7, 30, 33, 38, 39, 53, 54}. However, a recent study reported that the stQSSA can be inaccurate \cite{41}, which raises the question of validity conditions for the stQSSA.
Here, we identify the complete validity condition for using the stQSSA to simplify stochastic models containing rapid reversible bindings. Specifically, we find that the stQSSA is accurate for a wide range of conditions. However, when two species whose molar ratio is \( \sim 1:1 \) tightly bind, the stQSSA highly overestimates the number of unbound species. In this case, using the stQSSA to simplify stochastic models distorts the stochastic dynamics of the transcriptional repression, the transcriptional negative feedback loop of the circadian clock, and the bistable switch for mitosis. Importantly, by using the fact that the number of the unbound species is low due to the tight binding when the stQSSA is inaccurate, we develop an alternative approach, stochastic “low-state” QSSA (slQSSA). In this way, when reversible bindings are tight and not tight, slQSSA and stQSSA can be used, respectively, which enables one to obtain accurately reduced stochastic models for any case. This proposes a complete and straightforward strategy for efficiently simulating multiscale stochastic biochemical systems containing the fundamental elementary reaction, i.e., rapid reversible binding. To facilitate this framework, we provide a user-friendly open-source computational package, ASSISTER (Adaptive Simplification of Stochastic SystEm with Reversible binding).

**Results**

**stQSSA can overestimate the number of the unbound species**

In the reversible binding reaction (Eq. (1)), the concentration of \( A \), denoted by \( \tilde{A} \), is governed by the following ODE:

\[
\frac{d\tilde{A}}{dt} = -k_f \tilde{A} \cdot \tilde{B} + k_b \tilde{C} = -k_f \tilde{A} \cdot (\tilde{B} + \tilde{C}) + k_b (\tilde{B} + \tilde{C}) - k_i \tilde{A},
\]

where \( \tilde{A} = \tilde{A} + \tilde{C} \) and \( \tilde{B} = \tilde{B} + \tilde{C} \) are the total concentrations of the bound and unbound species. By solving \( \frac{d\tilde{A}}{dt} = 0 \) in terms of \( \tilde{A} \) and \( \tilde{B} \), the tQSSA for \( \tilde{A} \) can be obtained as follows:

\[
\tilde{A}_{eq} := \frac{1}{2} \left\{ (\tilde{A} + \tilde{B} - \tilde{K}_d) + \sqrt{(\tilde{A} + \tilde{B} - \tilde{K}_d)^2 + 4\tilde{A}\tilde{K}_d} \right\},
\]

where the \( \tilde{K}_d = k_b/k_i \) is the dissociation constant. Note that if the reversible binding (Eq. (1)) is embedded in a larger system, there could be other reactions affecting the dynamics of \( \tilde{A} \) and thus additional terms in Eq. (2). However, as long as the reversible binding is fast (i.e., \( k_i \) and \( k_b \) are much larger than the other reaction rates), \( \tilde{A}_{eq} \) is still an accurate tQSSA for \( \tilde{A} \). Similarly, by solving \( \frac{d\tilde{B}}{dt} = 0 \) and \( \frac{d\tilde{C}}{dt} = 0 \), the tQSSAs for \( \tilde{B} \) and \( \tilde{C} \) can be obtained. These tQSSAs, also known as the Morrison equations [6], are generally valid, unlike the Michaelis-Menten type equations which are valid only when the enzyme concentration is negligible [7]. Thus, the tQSSAs have been used to simplify models containing not only interactions between metabolites but also proteins whose concentrations are typically comparable.

Unlike the deterministic QSSA (Eq. (3)), the stochastic QSSA, which is the stationary average number conditioned on the total numbers of the bound and unbound species, has a complex form [41,45,50]. For instance, the stochastic QSSA for the number of \( A \) (\( \langle A \rangle \)) can be expressed in terms of the dimensionless variables and parameters, \( X = \tilde{X}\Omega \), where \( \Omega \) is the volume of a system (e.g., \( A = \tilde{A}\Omega \)).
$K_d = \tilde{K}_d \Omega$) as follows (see Methods for details):

$$
\langle A \rangle = \left( \sum_{l=0}^{A_T} \frac{lK_d}{l!(A_T - l)!(B_T - A_T + l)!} \right) \cdot \left( \sum_{l=0}^{A_T} \frac{K_d^2}{l!(A_T - l)!(B_T - A_T + l)!} \right)^{-1},
$$

(4)

where $A_0 = \max \{0, A_T - B_T\}$. This complex form of the stochastic QSSA does not provide any intuition and importantly increases computational cost. Thus, as an alternative to the stochastic QSSA, its approximation, the stQSSA was derived with the deterministic tQSSA \[7, 22–25, 31\]. Specifically, the stQSSA for $A$ ($A_{tq}$) can be derived from $\tilde{A}_{tq}$ (Eq. 3) after replacing the concentration-based variables and parameters $(\tilde{X})$ with dimensionless variables and parameters $(X)$ as follows:

$$
\langle A \rangle \approx A_{tq} := \frac{1}{2} \left\{ (A_T - B_T - K_d) + \sqrt{(A_T - B_T - K_d)^2 + 4A_T K_d} \right\}.
$$

(5)

Similarly, the stQSSA for $B$ and $C$ ($B_{tq}$ and $C_{tq}$) can be obtained from their deterministic tQSSAs as follows:

$$
\langle B \rangle \approx B_{tq} := \frac{1}{2} \left\{ (B_T - A_T - K_d) + \sqrt{(B_T - A_T - K_d)^2 + 4B_T K_d} \right\},
$$

$$
\langle C \rangle \approx C_{tq} := \frac{1}{2} \left\{ (A_T + B_T + K_d) - \sqrt{(A_T - B_T - K_d)^2 + 4A_T K_d} \right\}.
$$

(6)

To identify the validity conditions for these stQSSAs, we calculated the relative error $(R_X := \frac{X_{tq}(\tilde{X})}{X_{tq}(\tilde{X})})$ of the stQSSA ($X_{tq}$) to the stochastic QSSA ($\langle X \rangle$) (Fig. 1A, 1B). The errors are nearly zero in most of the parameter regions, which explains why various stochastic models reduced with the stQSSA have been accurate in most previous studies \[7, 30–32, 38–40, 44\]. However, the relative errors of the unbound species ($R_A$ and $R_B$) are high when $A_T \approx B_T$. Specifically, the relative error of the bound species ($R_C$) is at most $\sim 0.2$ but that of the unbound species ($R_A$, $R_B$) can be $\sim 100$.

To investigate why $R_A$ is high when $A_T \approx B_T$, we derived the exact upper and lower bounds for $R_A$ (see Methods for details):

$$
F_A S_A \leq R_A \leq 2F_A S_A,
$$

(7)

where $F_A$ is the Fano factor of $A$ (i.e., $\frac{\text{Var}(A)}{(A)}$), and $S_A$ is the relative sensitivity of $A_{tq}$ with respect to $B_T$ (i.e., $\left. \frac{dA_{tq}}{d B_T} \right|_{A_T}$). Furthermore, we proved that the Fano factor ($F_A$) is less than 1 (i.e., $A$ has a sub-Poissonian stationary distribution; see S1 Appendix for details). Therefore, $R_A$, especially its upper bound, mainly depends on $S_A$ (Figs 1A, 1B, and S1) whose formula can be derived in the following simple form, unlike $R_A$:

$$
S_A = \frac{1}{A_{tq}} \left. \frac{dA_{tq}}{d B_T} \right| = \frac{1}{\sqrt{(A_T - B_T - K_d)^2 + 4A_T K_d}}.
$$

(8)

Because $S_A$ attains the maximum value $\frac{1}{\sqrt{4A_T K_d}}$ at $B_T = A_T - K_d$, $S_A$ has a large maximum value when $K_d < 1$ at $A_T = B_T + K_d \approx B_T$. This explains why $R_A$, whose upper bound is mainly determined by $2S_A$, is large when the binding is tight ($K_d < 1$) and the total numbers of the bound and unbound species are similar ($A_T \approx B_T$) (Fig. 1B). In this case, the majority of $A$ is bound with $B$, and thus $A = 0$ most of the time (Fig. 1B left). That is, $A$ rarely becomes 1 by the weak unbinding reaction and then immediately $A$ becomes 0 by the strong binding reaction. As a result, the probability that $A = 1$ is approximately 1% (i.e., $\langle A \rangle \approx 0.01$), but the stQSSA for $A$ ($A_{tq}$) overestimates it as 10%, which is 10 times larger (Fig. 1B right). Since $A$ and $B$ are symmetric, the above analysis can be applied to $B$, analogously.
Fig 1. stQSSA overestimates the number of the unbound species when their molar ratio is $\sim$1:1 and binding is tight. (a-c) Heat maps of the relative errors ($R_X = \frac{|X_{stQSSA} - \langle X \rangle|}{\langle X \rangle}$) of the stQSSA ($X_{stQSSA}$) to the stochastic QSSA ($\langle X \rangle$) for $X = A, B, C$ in the reversible binding reaction (Eq. (1)). Color in the heat maps represents the maximum value of $R_X$ calculated by varying $K_d$ from $10^{-4}$ to $10^2$ for each total number of the bound and unbound species ($A_T = A + C$ and $B_T = B + C$). $R_A$ and $R_B$ can be extremely large when $A_T \approx B_T$ while $R_C$ is always small. (d) $R_A$ calculated over $B_T/A_T$ between 0 and 2 (gray arrow in a) for three fixed $K_d$ values ($10^{-4}$, $10^{-3}$ and $10^{-2}$). $R_A$ becomes larger as $B_T/A_T$ is similar to 1 and the $K_d$ becomes smaller (i.e., the binding becomes tighter). (e) $R_A$ mainly depends on the relative sensitivity of $A_{stQSSA}$ (i.e., $2S_A$), which can be derived in a simple form, unlike $R_A$ (Eq. (3)). The maximum value of $2S_A$ is given by $\sqrt{A_T K_d}$, which is achieved when $B_T/A_T$ is similar to 1 as in the case of $R_A$. (f) A trajectory (left) and the stationary probability distribution (right) of $A$ for a parameter set where $R_A$ is large (green triangle in d, $A_T = B_T = 100, k_f/\Omega = 10^4 s^{-1}, k_b = 1 s^{-1}$), simulated using the Gillespie algorithm. Since $A_T = B_T$ and $A$ binds with $B$ tightly, $A = 0$ (i.e., every $A$ is bound) most of the time, and it rarely becomes 1 by the weak unbinding reaction (solid arrow) and immediately comes back to 0 by the strong binding reaction (dotted arrow). As a result, when $K_d = 10^{-4}$, the probability that $A = 1$ is $\sim$0.01, but the stQSSA for $A$ overestimates it as $\sim$0.1, which is 10 times larger (i.e., a 10-fold error).
stQSSA can overestimate the transcriptional activity

We found that the stQSSA for the number of the unbound species is inaccurate if their molar ratio is ~1:1 and their binding is tight (Fig 1d:1f). Thus, we expected that in such cases, using the stQSSA to eliminate a rapid reversible binding in a stochastic model can distort its dynamics. To illustrate this, we constructed a simple gene regulatory network where gene expressions are determined by a reversible binding between transcription factors and genes (Fig 2a, left, Table S1); DNA (D) and a transcription factor (P) reversibly bind to form a complex (D:P). As P acts as a repressor of MR transcription, the transcription rate of MR is proportional to the number of the unbound DNA (D). On the other hand, as P acts as an activator of MA transcription, the transcription rate of MA is proportional to the number of the bound DNA (D:P). Note that the number of unbound and bound DNA can be interpreted as the number of unbound and bound DNA binding sites. In this model, because the reversible binding reaction between D and P is much faster than the other reactions (i.e., the production and the decay of MR and MA), the variables (D and D:P) rapidly reach their QSS. Thus, by replacing them with their stQSSAs (D_{tq} and D:P_{tq}), we can obtain a reduced model (Fig 2a, right, Table S2). The reduced model consists of only the slow variables, MR and MA, because D_{tq} and D:P_{tq} are fully determined by the conserved total number of the DNA (D_{T} = D + D:P) and the conserved total number of the transcription factor (P_{T} = P + D:P), as illustrated in Table S2. This elimination of the fast variables, which are the major source of computational cost, greatly reduces the computation time of stochastic simulations.

To test whether the reduced model accurately captures the dynamics of the full model, we compared their stochastic simulations with the Gillespie algorithm (see Tables S1 and S2 for propensity functions). When D_{T} and P_{T} are the same and the binding between D and P is tight, MR simulated with the reduced model largely exceeds MR simulated with the full model (Fig 2f top) because the stQSSA (D_{tq}) overestimates the stochastic QSSA for the number of the unbound DNA (⟨D⟩) which determines the transcription rate of MR (Fig 2a), as seen in Fig 1. On the other hand, when D_{T} is not similar to P_{T} (Fig 2d top) or the binding is weak (Fig 2f top), D_{tq} accurately approximates ⟨D⟩ as seen in Fig 1, and thus the reduced model accurately captures the dynamics of MR in the full model.

Unlike MR (Fig 2a top), the stochastic dynamics of MA of the reduced model and the full model are identical (Fig 2f bottom) because the stQSSA for D:P (D:P_{tq}) always accurately approximates the stochastic QSSA for the number of the bound DNA (⟨D:P⟩) which determines the transcription of MA (Fig 1). Taken together, the stQSSA can be used to describe transcriptional activation depending on bound DNA under any conditions (Fig 2f bottom). On the other hand, it needs to be restrictively used to describe transcriptional repression depending on unbound DNA (Fig 2f bottom).

stQSSA can distort oscillatory dynamics

To illustrate how the stQSSA distorts the dynamics when the molar ratio between tightly binding species is ~1:1, we investigated the simple model where the molar ratio is conserved (Fig 2). However, the total copy numbers of binding species and thus their molar ratio can be varied (e.g., oscillate) in a living cell due to other reactions in a larger system. This raises the question of whether the model reduction based on the stQSSA is accurate or not if the molar ratio is temporarily ~1:1. To investigate this, we used a modified Kim-Forger model, which describes the transcriptional negative feedback loop of the mammalian circadian clock. In this model (Fig 3a, top, Table S3), free activator (A) promotes the transcription of mRNA (M), and the
protein translated from M produces repressor (R) passing through several steps (P_i, i = 1, 2, 3). Then R reversibly binds with A to form a complex (A:R) which no longer promotes the transcription, and thus represses its own transcription. In this model, the reversible binding between R and A is much faster than the other reactions (i.e., production and decay). Thus, by replacing the fast variable A, which determines the transcription rate of M, with its stQSSA (A_tq), we can obtain a reduced model (Fig 3b, bottom, Table S4). The reduced model consists of only the slow variables, R_T, M and P_i, because A_tq is fully determined by the conserved total number of the activator (A_T = A + A:R) and the slowly varying total number of the repressor (R_T = R + A:R), as illustrated in Table S4.
**Fig 3. stQSSA can distort the dynamics of a biological oscillator.** (a) Full model diagram of an oscillatory transcriptional negative feedback loop (top, Table S3). Unbound activator (A) promotes the transcription of mRNA (M), and the protein translated from M produces repressor (R) passing through several steps (P_1, i = 1, 2, 3). Then R binds with A to form a complex (A:R) which is transcriptionally inactive, and thus represses its own transcription. As the reversible binding between R and A is rapid, by replacing A with its stQSSA (A_tq), we can obtain a reduced model which consists of only slowly varying R_T, M, and P_i (bottom, Table S4). (b-c) Oscillatory trajectories of M (green) and R_T/A_T (orange) simulated with the full model (b top) and the reduced model (b bottom), using the Gillespie algorithm (see Tables S3 and S4 for propensity functions). When R binds with A tightly (K_d = 10^{-4}) both the full model and the reduced model show the oscillatory behaviors. However, when the trajectory of R_T/A_T stays near 1 (dashed lines in b), A_tq overestimates the stochastic QSSA for A (⟨A⟩), and thus the transcription more frequently occurs in the reduced model (b bottom) compared to the full model (b top). As a result, the reduced model predicts a shorter period than the full model (c). (d-e) On the other hand, when the degradation rate of R increases and thus the trajectory of R_T/A_T stays near 1 for a short time (d; dashed lines), the reduced model accurately captures the dynamics of the full model (e).
In the model, the inactive form of cyclin B/Cdc2 (P) is converted to an active form (M) by Cdc25 (D). Furthermore, as M activates D which converts P to M, M promotes its own activation (i.e., form a positive feedback loop; see [46, 57] for details). The positive feedback loop is suppressed by Suc1 protein (B) as it binds with M to form a complex (M:B) which no longer activates D. The total activated cyclin B/Cdc2 (M and M:B) become P with the same constant rate. In this model, the reversible binding between M and B is much faster than the other reactions. Thus, by replacing the fast variable \( M \) with its stQSSA \( M_{\text{tq}} \), a reduced model can be derived (Fig 4b bottom, Table S6). The reduced model consists of only the slow variables, \( M_{\text{T}} \) and P, because \( M_{\text{tq}} \) is fully determined by the conserved total number of Suc1 \( (B_T = B + M:B) \) and the slowly varying total number of the activated cyclin B/Cdc2 \( (M_T = M + M:B) \), as illustrated in Table S6.

**Fig 4. stQSSA can distort the dynamics of a bistable switch** (a) Full model diagram of a bistable switch for mitosis (top, Table S5). The inactive form of cyclin B/Cdc2 (P) becomes an active form (M) by Cdc25 (D). In this process, M enhances its own activation by activating D, and thus forms a positive feedback loop (see [46, 57] for details). The positive feedback loop is suppressed as Suc1 protein (B) binds with M to form a complex (M:B) which does not activates D. The total activated cyclin B/Cdc2, M and M:B, becomes P with the same constant rate. As the reversible binding between M and B is rapid, by replacing M with its stQSSA \( (M_{\text{tq}}) \) a reduced model can be derived (Fig 4b bottom, Table S6). The reduced model consists of only the slow variables, \( M_{\text{T}} \) and P, because \( M_{\text{tq}} \) is fully determined by the conserved total number of Suc1 \( (B_T = B + M:B) \) and the slowly varying total number of the activated cyclin B/Cdc2 \( (M_T = M + M:B) \), as illustrated in Table S6.

(b-c) Simulated trajectories (b) and the stationary distributions (c) of \( M_{\text{T}} \) from the full model and the reduced model using the Gillespie algorithm (see Tables S5 and S6 for propensity functions). When M binds with B tightly \( (K_d = 10^{-3}) \), both the full model and the reduced model show the bistable behaviors (i.e., bimodal stationary distributions) of \( M_{\text{T}} \) (Fig 4b). However, the trajectory of the reduced model is more attracted to the upper mode of \( M_{\text{T}} \) compared to the full model (Fig 4b). As a result, the bimodal distribution of \( M_{\text{T}} \) from the reduced model is biased to the upper mode (c). (d-e) On the other hand, when the binding between M and B becomes weak \( (K_d = 10) \), \( M_{\text{tq}} \) accurately estimates \( \langle M \rangle \), and thus the reduced model accurately captures the dynamics of the full model, which no longer shows the bistable behavior.

When M and B tightly bind, both the full model and the reduced model show the bistable behaviors (i.e., bimodal stationary distributions) of \( M_{\text{T}} \) (Fig 4b). However, the trajectory of the reduced model is more attracted to the upper mode of \( M_{\text{T}} \) compared to the full model (Fig 4b and 4c). This dynamics biased to the upper mode occurs because \( M_{\text{tq}} \) overestimates the stochastic QSSA for M \( (\langle M \rangle) \) near the...
\( M_T / B_T = 1 \) region (Fig 4b dashed line) which separates the upper and lower modes. On the other hand, when the binding between M and B becomes weak, \( M_{th} \) accurately approximates the stochastic QSSA for \( M \) even when \( M_T \) is similar to \( B_T \). Thus, the reduced model accurately captures the dynamics of the full model, which no longer shows bistable behavior (Fig 4a and 4b). As expected, when \( A_T \) decreases because \( T \)

An alternative approach when the stQSSA is not applicable

In the presence of a rapid and tight reversible binding between species whose molar ratio is \( \sim 1:1 \), the reduction of stochastic models with the stQSSA for the number of the unbound species can cause errors (Figs 3a, 3b, and 3c). In such cases, due to the tight binding, the two species tend to bind until no molecules of one species are left (Fig 3d). Specifically, if \( A_T \leq B_T \) (\( A_T \geq B_T \)), the majority of the \( A \) (\( B \)) will be bound. Thus, in the presence of tight binding, we can assume that the stationary distributions could be used to derive an alternative approximation.

The accuracy of the sQSSA for \( A \) (Eq. (5)) is expected to increase when \( A_T K_d \) decreases because \( A_T K_d \) is an approximated number of the unbound \( A \). On the other hand, the accuracy of the stQSSA for \( A \) decreases as \( A_T K_d \) decreases (Fig 1c). To investigate this, we calculated the maximum relative error of \( A_{th} \) (Eq. (4)) (see Methods for details):

\[
⟨A⟩ \approx A_{th} = \left\{ \begin{array}{ll}
\frac{(A_T - B_T + 1)(A_T - B_T + B_T K_d)}{A_T - B_T + B_T K_d - 1} & \text{if } A_T \geq B_T, \\
\frac{A_T K_d}{B_T - A_T + A_T K_d + 1} & \text{if } A_T < B_T.
\end{array} \right.
\]

We will refer to this approximation as the stochastic “low-state” QSSA (sQSSA). Since this approximation relies on the property that the state space is restricted at the low level, the stochastic QSSA with singular perturbation analysis introduced in [58] could be used to derive an alternative approximation.

The parameters used in Figs 2a (triangle), 3b (circle), and 4b (square) are located in the region where the stQSSA is inaccurate (Fig 5a) but the sQSSA is accurate. In particular, when \( A_T K_d < 10^{-1} \) and \( A_T K_d > 10^1 \), \( R_{th} \) and \( R_A \) are less than 0.1 (i.e., the relative errors are less than 10%), respectively. The parameters used in Figs 2a (triangle), 3b (circle), and 4b (square) are located in the region where the sQSSA is inaccurate (Fig 5a) but the sQSSA is accurate (Fig 5b). Therefore, with these parameters, the reduced models obtained by using the sQSSAs accurately capture the dynamics of the full models for the simple gene regulatory network (Fig 5). Table S2, the transcriptional negative feedback loop (Fig 4d, Table S4), and the bistable switch for mitosis (Fig 5a, Table S6), unlike the stQSSA (Figs 2a, 3b, and 3c). Furthermore, by allowing \( A \) or \( B \) to reach more than two states (e.g., 0, 1, and 2), more accurate sQSSAs can be derived (see Methods for details). In particular, the relative errors of the sQSSAs derived by allowing the 3/4/5 states are less than 0.1 when \( A_T K_d \) is less than 2/5/10, respectively (Fig 5c).

Consequently, for the error tolerance of 0.1, if \( A_T K_d < 10^1 \) and thus the stQSSA is inaccurate, the sQSSA can be used to approximate the stochastic QSSA for \( A \) (Fig 5a). Taken together, by using either stQSSA or sQSSA depending on \( A_T K_d \), we can always accurately reduce multiscale stochastic biochemical systems with rapid reversible bindings. Of course, for a different error tolerance, we need a different threshold of \( A_T K_d \) and the number of states for the sQSSA. To facilitate the calculation of such change depending on the error tolerance, we have developed a
Fig 5. slQSSA can be used to reduce multiscale stochastic biochemical systems containing rapid reversible bindings when the stQSSA is not applicable. (a-b) Heat maps of the relative errors ($R_A = \frac{|A_{\text{slQSSA}} - \langle A \rangle|}{\langle A \rangle}$) and $R_{\text{slQSSA}} = \frac{|A_{\text{slQSSA}} - \langle A \rangle|}{\langle A \rangle}$) when the stQSSA ($A_{\text{stQSSA}}$) and the two-state slQSSA ($A_{\text{slQSSA}}$) approximate the stochastic QSSA for $A$ ($\langle A \rangle$) in the reversible binding reaction (Eq. 1). Color represents the maximum value of $R_A$ and $R_{\text{slQSSA}}$ for each $A_TK_d$ and $K_d$ when $B_T$ varies, and the dashed lines represent when those values are 0.1. When $A_TK_d$ are high and low, the stQSSA and the slQSSA are accurate, respectively. The parameters used in Figs 2b (triangle), 3b (circle), and 4b (square) are located in the region where the slQSSA (b), but not the stQSSA (a), is accurate (the circle is actually located outside of the heat maps; $A_TK_d = 5 \times 10^{-4}$ and $K_d = 10^{-4}$). (c-e) As a result, the full models are successfully reduced with the slQSSA (c-e) but not the stQSSA (Figs 2b, 3c, and 4c). See Tables S2, S4, and S6 for the propensity functions used for the simulations and Fig S3 for a benchmark comparison with GillesPy2, one of the major, standard software suites for stochastic simulation [59]. (f) The adaptive use of the stQSSA and the slQSSA to approximate the stochastic QSSA for $A$ when $A_TK_d > 10^1$ and otherwise, respectively, guarantees the successful reduction of stochastic models containing rapid reversible bindings. Note that when $10^{-1} < A_TK_d < 10^1$, the slQSSAs with more than two states need to be used (see Fig S2 for details).
user-friendly open-source computational package, ASSISTER (Fig 6). In particular, the function Gillespie_Reduction in the package automatically constructs a reduced model adaptively using the more accurate one between the two approximation methods and performs accurate and efficient stochastic simulations (see SI Appendix for the manual).

![Schematic diagram for the computational package, ASSISTER.](https://example.com/schematic-diagram)

**Fig 6. Schematic diagram for the computational package, ASSISTER.** When a stochastic model containing a rapid reversible binding (D + P ⇌ D:P) and the error tolerance (ε) are given as inputs, the auxiliary function QSSA_Threshold determines the threshold of the total number of binding molecules, \( D_{\text{thres}} \), and the number of states for the slQSSA, \( L \). When the \( D_T = D + D:P \) is less (larger) than \( D_{\text{thres}} \), the \( L \)-state slQSSA (stQSSA) approximates the exact stochastic QSSA with a relative error less than \( \epsilon \). Based on the relationship between \( D_{\text{thres}} \) and \( D_T \), the more accurate one between the two models is adaptively chosen. In this way, Gillespie_Reduction performs efficient and accurate stochastic simulations, yielding the simulated trajectories as the final output. See the manual in SI Appendix for a more detailed description of the input and output.

**Discussion**

Reversible binding between molecules—for example, between DNA and a transcription factor, a ligand and a receptor, and an enzyme and a substrate—is a fundamental reaction for numerous biological functions [42]. As the reversible binding reactions occur typically on a timescale of 1~1000ms, which is much faster than the other reactions (e.g., 30min for a mammalian mRNA transcription or a protein translation and 10h for their typical lifetimes) [43], a system containing the rapid reversible binding becomes a multi-timescale system. In such multi-timescale systems, the rapid reversible binding prohibitively increases the computational cost of stochastic...
simulations. Accordingly, to accelerate stochastic simulations, various methods have been developed \[44, 60\]. In particular, the model reduction using the stQSSA has successfully simplified various stochastic models in numerous studies \[7, 30–33, 38, 39\]. Thus, it has been commonly believed that the stQSSA is generally accurate for any conditions, until a recent counterexample was identified \[41\]. In this work, we rigorously derived the validity conditions for using the stQSSA to reduce stochastic models with a rapid reversible binding. Specifically, we showed that the relative error of the stQSSA for the number of unbound species \((R_A)\) mainly depends on the relative sensitivity of the stQSSA \((S_A, \text{Eq. (8)})\), which attains maximum value \(\frac{1}{\sqrt{A_T K_d}}\) at \(A_T = B_T + K_d\). This allowed us to find that the stQSSA for the number of the unbound species is inaccurate if their molar ratio is \(\sim 1:1\) and their binding is tight (Fig 1f). In that case, the stQSSA highly overestimates the number of the unbound species. Therefore, the reduced models obtained by using the stQSSA distort the dynamics of the gene regulatory model (Fig 2b), the transcriptional negative feedback loop model for circadian rhythms (Fig 3c), and the bistable switch model for mitosis (Fig 4c).

When the reversible binding reactions are sufficiently faster than the other reactions, the deterministic tQSSA is known to be accurate \[7, 48, 50\]. Indeed, for all examples considered in our work (Figs 2b, 3b, and 4b), the deterministic simulations with the tQSSA are accurate, unlike the stochastic simulations. This indicates that it is risky to investigate the validity conditions of the stQSSA solely based on the validity conditions of the deterministic tQSSA. Instead, the direct derivation of the relative error of the stQSSA is needed, as demonstrated in this study (Eq. (7)). It would be interesting in future work to perform such error analysis for more complex examples, such as coupled enzymatic networks with multiple rapid reversible bindings \[26, 61, 62\].

The rapid reversible reactions are typical conditions for model reductions using the “partial-equilibrium approximation,” which confines species concentrations to the equilibrium states. In general, this condition does not imply a timescale separation between the variables (\(A, B, \text{and } C\)), limiting the application of the QSSA. However, the rapid reversible binding guarantees that the total variables \(A_T\) and \(B_T\) always evolve more slowly than the variables \(A, B, \text{and } C\). Therefore, the QSSAs in terms of the total variables can lead to accurate model reductions in both deterministic \[7, 47, 48, 50\] and stochastic \[27, 30\] regimes. In this work, we investigated under which conditions the complex stochastic QSSA (Eq. (4)) can be approximated by the corresponding simple deterministic QSSA (Eq. (5)), referred to as the stQSSA. We found that such an approximation becomes inaccurate when two species with similar levels tightly bind. Thus, we derived an alternative approximation for the stochastic QSSA: slQSSA. Using the two approximations for the stochastic QSSA, we developed the universally valid reduction framework for stochastic models containing rapid reversible bindings. We also provided the user-friendly open-source computational package, ASSISTER, for this framework. On the other hand, in the absence of rapid reversible binding, one can reduce a model by assuming that ‘highly reactive species’ are in their QSSs \[58\]. Interestingly, the reduced stochastic models were often different from the heuristically reduced models obtained with the deterministic QSSA. It would be interesting in future work to investigate when such discrepancies occur.

While the deterministic tQSSA (Eq. (3)) was used to approximate the stochastic QSSA for the number of reversibly binding species in this work, a simpler deterministic QSSA referred to as the ‘standard’ QSSA (sQSSA) is more widely used to approximate the stochastic QSSA due to its simplicity \[8, 20, 27, 32\]. For instance,
the stochastic sQSSA for $C$ in Eq. (1), which has the Michaelis-Menten type form

$$C_{sq} = \frac{A_T B}{B + K_d},$$

has been widely used as a propensity function for the Gillespie algorithm. Though this sQSSA has been widely used, it is less accurate than the stQSSA (Eq. (5)) [38, 39]. This is why many examples showing the inaccuracy of the stochastic sQSSA have been reported [33–40], whereas only one example showing the inaccuracy of the stQSSA has been reported [41]. Note that while Eq. (10) is different from the typical “Michaelis-Menten” equation, which uses the Michaelis-Menten constant instead of the dissociation constant ($K_d$), they become nearly the same when the timescale of reversible binding is faster than the catalytic reaction. Importantly, our work also provides the validity condition for using the stochastic sQSSA (Eq. (10)). That is, when $B_T + K_d \gg A_T$, which is known as the low-enzyme concentration condition, $C_{sq} \approx C_{eq}$ [7], indicating that the Michaelis-Menten type sQSSA for the bounded species (Eq. (10)) can be used to reduce models containing the rapid reversible binding. Similarly, when $B_T + K_d \gg A_T$, $A_{sq} = \frac{A_T K_d}{B_T + K_d}$ could also be used. This is consistent with the validity conditions for the stochastic sQSSA derived under the assumption of either a low fluctuation level [32] or a low copy number [40]. Furthermore, the “pre-factor” QSSA (pQSSA), which is more accurate than the sQSSA, has also been used for stochastic simulations [63, 64]. However, recent studies have shown that the stQSSA is more accurate than the stochastic pQSSA (see [33, 39] for details).

The accuracy of the stQSSA for the number of the unbound species depends on both the molar ratio between reversibly binding species and the tightness of their binding (Fig 1A). However, as the molar ratio typically varies in larger models containing reversible binding, practically, the accuracy is mainly determined by the tightness of binding. Specifically, for the relative error of the stQSSA to be less than 0.1, $A_T K_d$ ($\approx$ the number of the unbound $A$) should be larger than 10 (Fig 1A, dashed line). This $A_T K_d$ value-based criteria explains the controversy about the accuracy of the stQSSA in previous studies. That is, $A_T K_d$ was less than 10 in a previous study where the reduced model obtained by using the stQSSA was inaccurate [11]. On the other hand, $A_T K_d$ were much greater than 10 in all of the examples investigated in previous studies reporting the accuracy of the stQSSA [7, 33, 38, 39, 53, 54]. Furthermore, the stQSSA always accurately approximates the stochastic QSSA for the number of the bound species (Fig 1A). This explains why the stQSSA was accurate in previous studies where the stQSSA was used to approximate the number of enzyme-substrate complex [30, 33].

In real biological systems, the validity condition of the stQSSA ($A_T K_d > 10$) is not always guaranteed. Specifically, the range of $A_T K_d$ can span approximately from $10^{-3}$ to $10^{10}$ in human cells ($\Omega = 10^{-15} \sim 10^{-14} m^3$) since the protein-protein dissociation constant ($K_d$) is $10 fM \sim 1 \mu M$ (i.e., $10^{12} \sim 10^{20} m^{-3}$), and the numbers of molecules ($A_T$) is $10^9 \sim 10^{14}$ [43, 65]. Moreover, in smaller cells like budding yeast ($\Omega = 10^{-17} \sim 10^{-16} m^3$) or E. Coli ($\Omega = 10^{-19} \sim 10^{-18} m^3$) cells, the range of $A_T K_d$ can span from $10^{-7}$ to $10^8$, which contains the region in which the stQSSA can be extremely inaccurate. Accordingly, the sQSSA, which accurately approximates the stochastic QSSA when $A_T K_d$ is less than 10, is necessary. Specifically, the relative error of the sQSSA, unlike that of the stQSSA (Fig 5A and 5D), decreases as $A_T K_d$ decreases because the sQSSA relies on the assumption that the stationary distributions of the number of the unbound species ($\approx A_T K_d$) are concentrated on the few lowest states. Taken together, by using the stQSSA and the sQSSA when the $A_T K_d$ value is greater and less than 10, respectively, one can always accurately simplify stochastic models containing rapid reversible binding reactions to accelerate
simulation and also facilitate stochastic analysis (Fig 5). This can be facilitated by the computational package, ASSISTER (Fig 6).

**Methods**

**Exact bounds for the relative error of the stQSSA to the stochastic QSSA**

In this section, we derive the exact upper and lower bounds for \( R_A = \left| \frac{A_{tq} - \langle A \rangle}{\langle A \rangle} \right| \) (Eq. 7), where \( A_{tq} \) and \( \langle A \rangle \) are the stQSSA and the stochastic QSSA for \( A \), respectively.

From the CME describing the reversible binding reaction (Eq. (1)), the following steady-state moment equation can be derived:

\[
\langle k_T A \cdot B / \Omega \rangle = \langle k_b C \rangle ,
\]

where \( \langle \cdot \rangle \) is the stationary expectation. Eq. (11) becomes

\[
\langle A \rangle = \frac{K_d}{K_d + [B]} \approx \langle k_T A \rangle = \langle k_b C \rangle,
\]

By dividing Eq. (16) by \( \langle k_T A \rangle \), we get the following quadratic equation:

\[
\langle A \rangle^2 - \langle A \rangle - \langle A \rangle - A_T K_d + \langle A \rangle = 0.
\]

The non-negative root of this quadratic equation becomes \( \langle A \rangle \):

\[
\langle A \rangle = \frac{1}{2} \left\{ (A_T - B_T - K_d) + \sqrt{(A_T - B_T - K_d)^2 + 4A_T K_d - 4 \langle A \rangle} \right\}.
\]

By subtracting Eq. (13) from Eq. (15), we get

\[
A_{tq} - \langle A \rangle = \frac{1}{2} \left\{ \sqrt{(A_T - B_T - K_d)^2 + 4A_T K_d} - \sqrt{(A_T - B_T - K_d)^2 + 4A_T K_d - 4 \langle A \rangle} \right\}
\]

Since \( 0 \leq (A_T - B_T - K_d)^2 + 4A_T K_d - 4 \langle A \rangle \leq (A_T - B_T - K_d)^2 + 4A_T K_d \), we get the bounds for \( A_{tq} - \langle A \rangle \) from Eq. (15):

\[
\frac{2 \langle A \rangle}{\sqrt{(A_T - B_T - K_d)^2 + 4A_T K_d}} \leq A_{tq} - \langle A \rangle \leq \frac{2 \langle A \rangle}{\sqrt{(A_T - B_T - K_d)^2 + 4A_T K_d}}.
\]

By dividing Eq. (10) by \( \langle A \rangle \), we can get the bounds for the relative error, \( R_A = \left| \frac{A_{tq} - \langle A \rangle}{\langle A \rangle} \right| \) as follows:

\[
\frac{\langle A \rangle}{\sqrt{(A_T - B_T - K_d)^2 + 4A_T K_d}} \leq R_A \leq \frac{1}{\sqrt{(A_T - B_T - K_d)^2 + 4A_T K_d}}.
\]

This can be re-expressed as \( F_A S_A \leq R_A \leq 2F_A S_A \) (Eq. (11)) because \( \frac{\langle A \rangle}{\sqrt{(A_T - B_T - K_d)^2 + 4A_T K_d}} \) is the relative sensitivity of \( A_{tq} \), i.e.,

\[
S_A = \frac{1}{\langle A_{tq} \rangle} \left| \frac{dA_{tq}}{dB_T} \right| .
\]
The relative sensitivity, $S_A$, attains the maximum value \( \frac{1}{\sqrt{A_0 K_d}} \) when the term in the square root of the denominator has the minimum value, i.e., $B_T = A_T - K_d$ (Eq. 4). In particular, $S_A$ has a large maximum value when $K_d \ll 1$ at $A_T = B_T + K_d \approx B_T$. On the other hand, if $A_T \ll B_T$, $S_A \approx 0$ because the majority of $A$ presents in the bound state regardless of $B_T$ (i.e., \( \frac{dA_{\text{sequestered}}}{dA_T} \approx 0 \)). When $A_T \geq B_T$, \( \frac{dA_{\text{sequestered}}}{dA_T} \approx 1 \) because as $B_T$ decreases by one, approximately one $A$ is released from the complex. In this case, if $A_T \gg B_T$, the majority of $A$ are free and thus $\frac{1}{A_0} \approx \frac{1}{A_T-B_T} \approx 0$, leading to $S_A \approx 0$. However, if $A_T \approx B_T$, the majority of $A$ is sequestered by $B$, $A_{\text{sequestered}} \approx 0$, leading to $S_A \gg 1$. When binding is weak ($K_d \gg 1$), $S_A \approx 0$ because the number of $A$, which is approximated by $A_{\text{sequestered}}$, changes little as $B_T$ changes (i.e., \( \frac{dA_{\text{sequestered}}}{dB_T} \approx 0 \)). Taken together, $S_A$ is large only when the binding reaction is tight ($K_d \ll 1$) and the binding species are present with 1:1 molar ratio ($A_T \approx B_T$).

Since $A_T = A_T \Omega$, $B_T = B_T \Omega$, and $K_d = K_d \Omega$, $S_A = o(\Omega^{-1})$ if the concentrations remain constant. This implies that when the volume $\Omega$ goes to infinity (i.e., thermodynamic limit), $S_A$ and thus $R_A$ become zero (i.e., the stochastic QSSA becomes nearly identical to the deterministic QSSA (tQSSA)). On the other hand, as $\Omega$ goes to zero (i.e., the volume of the system gets smaller), $S_A$ goes to infinity.

### Derivation of the stochastic QSSA and the sQSSA

Here we derive the stochastic QSSA for $A$ (\( \langle A \rangle \), Eq. 4). Let $p(l)$ be the probability that $A = l$ at its stationary distribution (i.e., the probability that $A(\infty) = l$). Then the following recurrence relation of $p(l)$ can be obtained from the steady-state CME:

\[
(l+1)(B_T-A_T+l+1)p(l+1) - (l+1)(B_T-A_T+l)p(l) + K_d(A_T-l+1)p(l-1) - K_d(A_T-l)p(l) = 0.
\]

(17)

Let $A_0 = \max\{A_T - B_T, 0\}$. Since $A_0$ is the lowest state that $A$ can reach, $p(l) = 0$ for $l < A_0$. Then we can inductively prove that the following relation satisfies Eq. (17):

\[
p(l + A_0) = \begin{cases} 
\pi(l + A_0)p(A_0) & \text{for } 0 \leq l \leq A_T - A_0, \\
0 & \text{otherwise},
\end{cases}
\]

(18)

where $\pi(l) = \frac{K_d^{l-A_0} \min(A_T, B_T)!(A_T-l)!}{l!(A_T-l)!(B_T-A_T+l)!}$. Then, because $\sum p(l) = 1$, $p(l) = \pi(l) \cdot \left( \sum_{l=A_0}^{A_T} \pi(l) \right)^{-1}$ if $A_0 \leq l \leq A_T$, and $p(l) = 0$ otherwise by Eq. (18). Therefore, we can obtain the stationary average number of $A$ (Eq. 4) as

\[
\langle A \rangle = \sum_{l=A_0}^{A_T} l \pi(l) \cdot \left( \sum_{l=A_0}^{A_T} \pi(l) \right)^{-1} = \left( \sum_{l=A_0}^{A_T} \frac{lK_d^l}{l!(A_T-l)!(B_T-A_T+l)!} \right) \cdot \left( \sum_{l=A_0}^{A_T} \frac{K_d^l}{l!(A_T-l)!(B_T-A_T+l)!} \right)^{-1}.
\]

Next we derive the sQSSA, which is the approximation for Eq. (4). In the presence of tight binding, we can assume that the stationary distributions of $A$ and $B$ are concentrated on the states \{0, 1\} when $A_T < B_T$ and $A_T \geq B_T$, respectively. Since when the distribution of $B$ is concentrated on 0 and 1, the distribution of $A$ is concentrated on $A_T - B_T$ and $A_T - B_T + 1$, we can simply say that the distribution of $A$ is concentrated on $A_0$ and $A_0 + 1$. Thus, by assuming that $p(l) = \pi(l) \cdot \left( \sum_{l=A_0}^{A_T} \pi(l) \right)^{-1}$ is approximately zero for $l > A_0 + 1$ and...
\[
\sum_{l=A_0}^{A_T} \pi(l) \approx \sum_{l=A_0}^{A_0+1} \pi(l),
\]
we can derive the two-state slQSSA for \( A \) (Eq. (9)) as follows:

\[
\langle A \rangle \approx \left( \sum_{l=A_0}^{A_0+1} \frac{lK_d^l}{l!(A_T - l)!(B_T - A_T + l)!} \right) \cdot \left( \sum_{l=A_0}^{A_0+1} \frac{K_d^l}{l!(A_T - l)!(B_T - A_T + l)!} \right)^{-1}
\]

\[
= \left\{ \begin{array}{ll}
(A_T - B_T + B_T K_d) \cdot \left( 1 + \frac{B_T K_d}{A_T - B_T + B_T K_d} \right)^{-1} & \text{if } A_T \geq B_T \\
\frac{A_T K_d}{B_T - A_T - A_T K_d + 1} \cdot \left( 1 + \frac{A_T K_d}{B_T - A_T - A_T K_d + 1} \right)^{-1} & \text{if } A_T < B_T
\end{array} \right.
\]

In general, for any integer \( k \geq 2 \), we can derive the \( k \)-state slQSSA as

\[
A^k_q := \left( \sum_{l=A_0}^{A_0+k-1} \frac{lK_d^l}{l!(A_T - l)!(B_T - A_T + l)!} \right) \cdot \left( \sum_{l=A_0}^{A_0+k-1} \frac{K_d^l}{l!(A_T - l)!(B_T - A_T + l)!} \right)^{-1}.
\]

(19)

Computational package for universally valid reduction of stochastic models containing rapid reversible binding reactions

We have developed a user-friendly computational package ASSISTER that contains three main codes implemented in MATLAB (Fig 6): \texttt{LQSSA}, \texttt{QSSA\_Threshold}, and \texttt{Gillespie\_Reduction}. \texttt{LQSSA} calculates the \( L \)-state slQSSA (Eq. (19)) for given \( A_T, B_T, K_d \), and \( L \). \texttt{QSSA\_Threshold} determines which of the slQSSA and the \( L \)-state slQSSA ensures a smaller error than a tolerance \( \epsilon \) for a given \( K_d \) value. This allows the function \texttt{Gillespie\_Reduction} to perform accurate stochastic simulations for any values of the parameters with the adaptive choice of the valid approximation method determined by using \texttt{QSSA\_Threshold} (Fig 6). See S1 Appendix for details and the manual. ASSISTER can be found at https://github.com/Mathbiomed/ASSISTER.

Supporting information

S1 Appendix. Supplementary Methods, Tables S1-S6, and Figs S1-S3. (PDF)

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Author Contributions

All authors designed the study and performed mathematical analysis. YS performed the computation and all authors analyzed the computation results. YS and JKK wrote the draft and all authors revised the manuscript.
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