

Synchronization of Biochemical Oscillations

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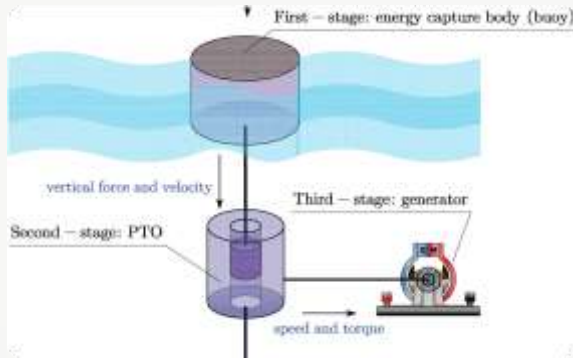
Under the Guidance of

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Introduction

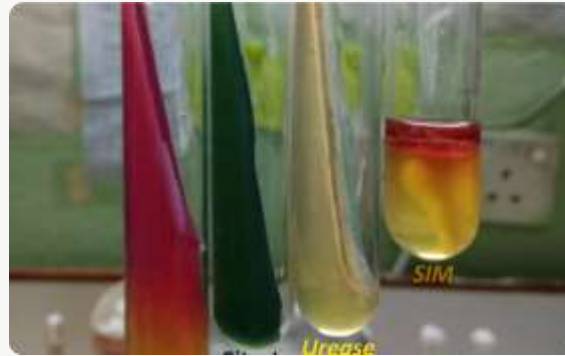
This study explores the synchronization dynamics of biochemical oscillators through numerical simulations. By analyzing the onset of oscillations in a single oscillator and the synchronization behavior of interconnected oscillators, we investigate the influence of critical parameters such as decay rate and time delay on the emergence and coordination of oscillatory behavior. Our results provide insights into the impact of parameter variations on the dynamics of oscillators, contributing to a deeper understanding of regulatory mechanisms in biological systems.

Biochemical Oscillators



Oscillating Waves

Visual representation of biochemical oscillations showcasing their rhythmic and periodic nature.



Regulatory Mechanisms

Diagram illustrating the intricate feedback mechanisms in biochemical systems responsible for oscillatory behavior.



Emergent Dynamics

Image depicting the synchronization of biochemical oscillators and the emergence of collective dynamics.

Understanding the Hill Function

Hill Function

The Hill function is a mathematical model used to describe cooperative interactions in biological systems. It introduces non-linearity to the model, representing phenomena such as cooperative binding or inhibition.

Single Oscillator

The Hill function is expressed as a vital element in understanding the behavior of a single biochemical oscillator.

Non-linear Response

The Hill coefficient dictates the steepness of the curve, capturing the non-linear response in the system.

Dynamics of a Single Biochemical Oscillator

$H(x,h)=1/(1+x^h)$ in Single Oscillator

- $H(x,h)$: Hill function output.
- x : Input variable
- h : Hill coefficient, determining the steepness of the curve.

$$H(x_1,x_2,h_1,h_2)=1/(1+x_1^{h_1}+x_2^{h_2})$$

1

Hill Function Application

The differential equation governing the behavior of a single biochemical oscillator involves the application of the Hill function to a delayed term, capturing the non-linear response.

2

Critical Parameters

The model incorporates critical parameters such as decay rate and time delay, exploring their influence on the emergence and coordination of oscillatory behavior.

3

Insights from Simulations

Numerical simulations based on these equations provide insights into the onset and behavior of oscillations in biochemical systems.

Differential Equation for a Single Oscillator

Rate of Change

The differential equation dx/dt represents the rate of change of the oscillator's state over time.

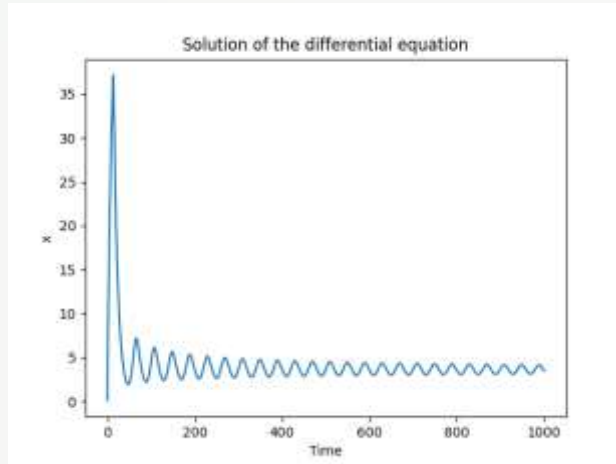
Influencing Factors

The equation incorporates the birth rate (K), decay rate (r), and the Hill function applied to a delayed term.

Time-delayed Influence

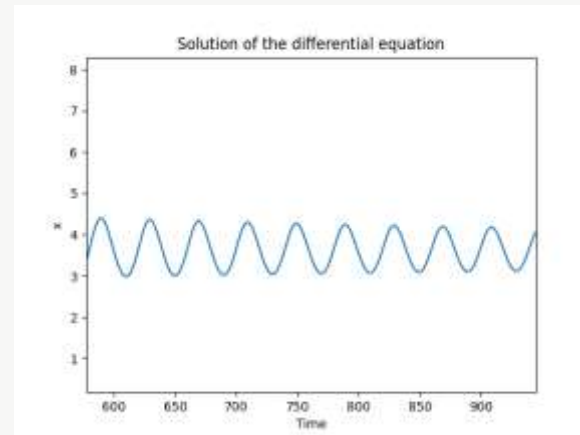
The delayed term represents the influence from the past state of the oscillator, incorporating a time delay (τ).

Time Delay



Dynamics and Period

Time delay (τ) influences the period and dynamics of the oscillator, affecting the timing of oscillatory behavior.



Response to Changes

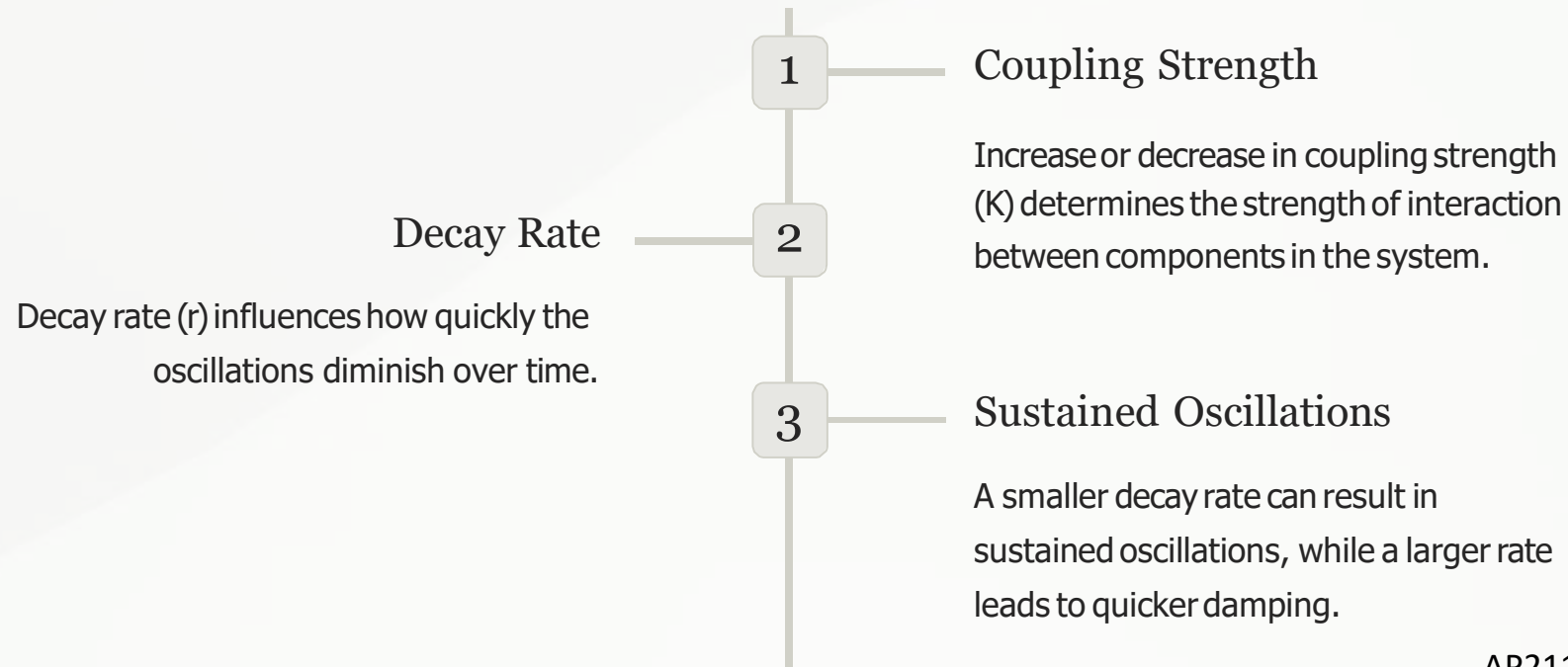
Time delay represents the time it takes for the system to respond to changes, affecting synchrony and coordination.

r (Decay rate)	τ (Time delay)
0.1	13.5
0.12	11.6
0.2	7.32
0.21	7.15
0.25	6.15
0.3	5.3
0.35	4.7
0.5	3.6

Impact on Synchronization

The interplay between time delay and the delayed term affects the synchronization behavior of interconnected oscillators.

Coupling Strength and Decay Rate



Single Oscillator Observations

Critical Parameters	Effect of Decay Rate (r)	Periodic Peaks
High r	Decreased critical time delay (τ)	Amplitude decreases with successive peaks
Low r	Increased critical time delay (τ)	Persistent periodic oscillations

Role of Dual Time Delays in Synchronization

1

Effect of Decay Rate

As the decay rate increases, the required time delays for synchronization decrease, highlighting the critical role of decay rate in synchronization.

2

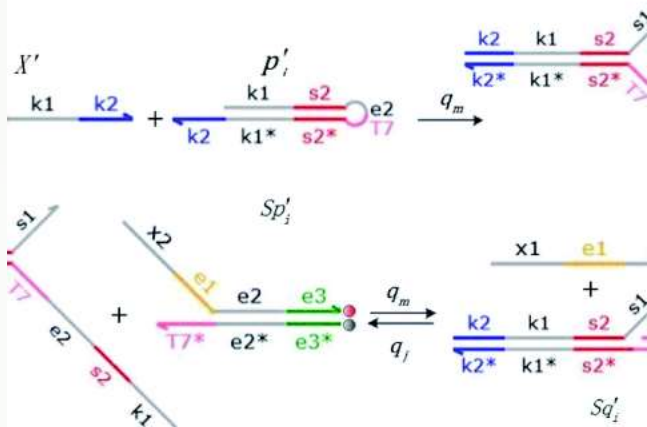
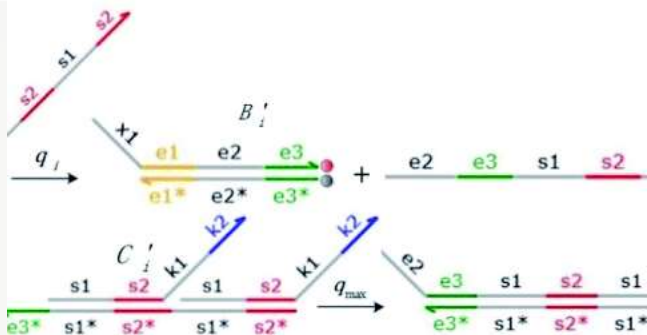
Symmetry in Peak

When τ_2 is equal to $2 \tau_1$ or vice versa, the two interconnected oscillators exhibit symmetry in their peaks, contributing to synchronization.

3

Transition to Square-like Format

As the decay rate increases, both graphs tend to take on a square-like format, indicating a more synchronized and periodic behavior.



(a)

iate CRNs	ideal formal CRNs
$\left. \begin{array}{l} 1) \\ 2) \end{array} \right\} \Rightarrow X_M \xrightarrow{\theta_1} 2X'_M \quad (5)$	$\left. \begin{array}{l} (3) \\ (4) \end{array} \right\} \Rightarrow X'_M \xrightleftharpoons[\theta_2]{\theta_m} X \quad (6)$
(7)	
$= \frac{\theta_2}{C_m}, q_m = \frac{\theta_m}{C_m}, q_i = \frac{k_1}{C_m}$	



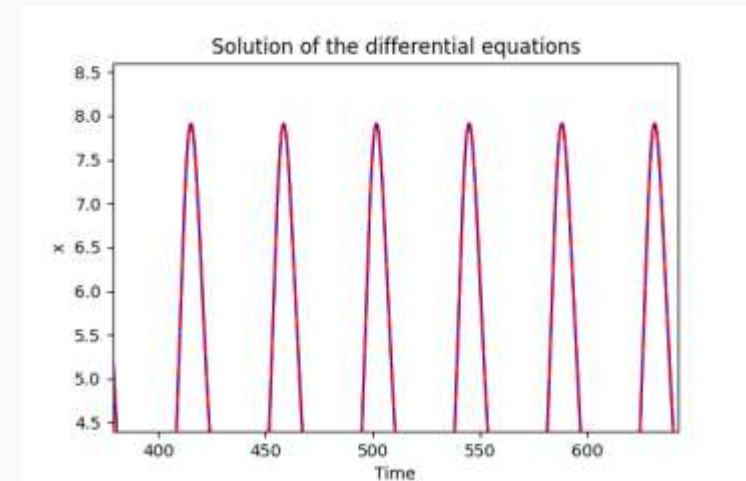
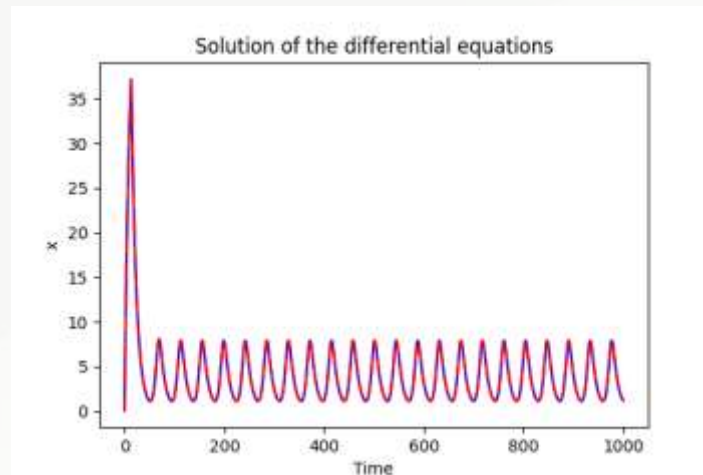
Two Oscillators Observations

Critical Parameters	Impact of Dual Time Delays (τ_1 and τ_2)	Symmetry in Peaks	Transition to Square-like Format
High r	Inverse relationship with dual time delays (τ_1 and τ_2)	Peaks exhibit synchronization influenced by τ_1 and τ_2 ratio	Square-like format indicates synchronized periodic behavior

Synchronization at Equal τ_1 and τ_2

1 Critical Condition

Synchronization occurs when the dual time delays (τ_1 and τ_2) are balanced and become equal.



Overall Trends

Decay Rate and Synchronization

The increase in decay rate (r) tends to stabilize and synchronize the oscillations in both single and dual oscillator scenarios.

Critical Time Delays

Shorter time delays favor sustained oscillations, and synchronization is achieved when the dual time delays (τ_1 and τ_2) are balanced.

Symmetry and Synchronization

Symmetry in the peaks between interconnected oscillators is influenced by the ratio of the dual time delays (τ_1 and τ_2).

Conclusion

The synchronization dynamics of biochemical oscillators exhibit a complex interplay between critical parameters such as decay rate, coupling strength, and time delays. Through numerical simulations, we have gained insights into the emergence and coordination of oscillatory behavior in both single and interconnected oscillators. Our findings contribute to a deeper understanding of regulatory mechanisms in biological systems, aiding future research and potentially informing therapeutic interventions.