

Neural Schrödinger Bridge for Minimum Effort Stochastic Control of Colloidal Self-assembly

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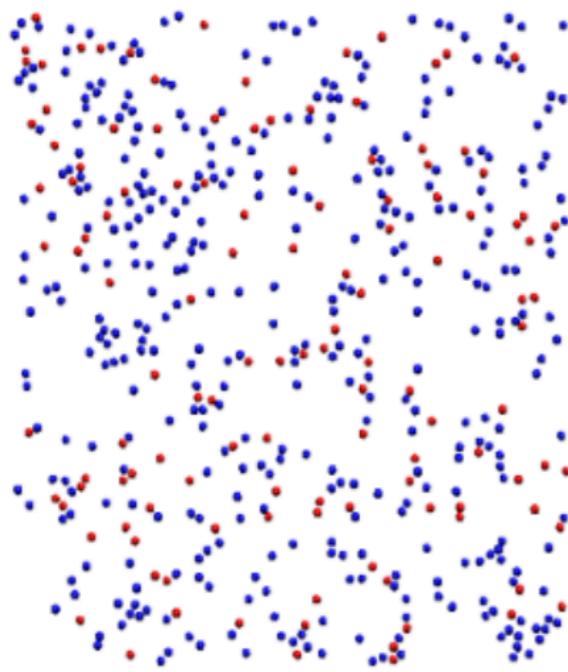
Ali Mesbah
(UC Berkeley)

Mini-symposium on Optimal Transport: Theory and Applications in Systems and Control
SIAM Conference on Control and Its Applications, Philadelphia

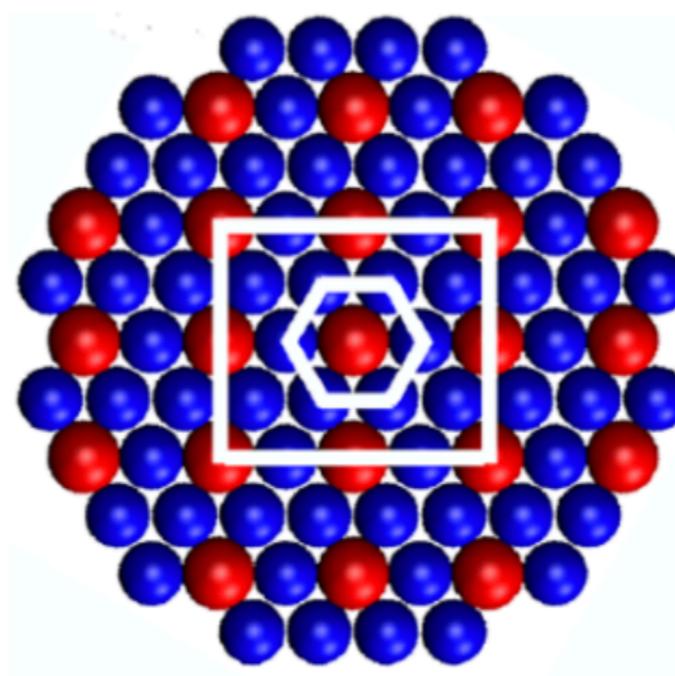
July 25, 2023



Colloidal Self-assembly (SA)



Dispersed particles



Ordered structure

Applications: Precision (sub nm scale) manufacturing of materials with advanced electrical, optical or magnetic properties

This talk: Two controlled colloidal SA case studies:
(1) model-based, (2) data-driven

SIAM news
Newsjournal of the Society for Industrial and Applied Mathematics
Volume 56 Issue 6 July/August 2023
Self-assembly and Particle Aggregation in Stratified Fluids
Experimental Discovery and First-principle Mechanisms of Diffusion-induced Flows in Exterior Domains

By Roberto Camassa and Richard M. McLaughlin

The motion of particulates in the ocean as they fall under gravitation constitutes a complex hydrodynamic cycle. Phytoplankton can photosynthesize dissolved carbon dioxide gas into dense, solid carbon debris called "marine snow," which facilitates the transport of carbon from the surface waters to the ocean depths. Over the course of years, the carbon eventually turns into oil. The prevailing view maintains that particulate aggregation occurs through random turbulent oceanic motions that generate collisions between small-scale sticky particulates, which then form aggregates [4]. However, it is commonly observed at similar depths in stratified water—that is, water in which different layers have different densities—such particles—when suspended at similar depths in stratified water—can slowly form aggregates in the absence of both external random flows and adhesion [7]. Interestingly, these particles' orientation and effective size and attractive force that originates from the no-slip boundary condition. This attraction leads to self-assembly.

that particulate aggregation occurs through random turbulent oceanic motions that subsequently appear to dynamically solve jigsaw-like puzzles while en route to large-scale particulate discs.

Figure 1a depicts a time-series sequence of the motion of a particulate molecule¹ that eventually merges with a larger aggregate, and Figure 1b portrays a schematic of the experimental setup. Figures 1c and 1d present an initial state of the system and a final state 200 minutes later. Our experiments revealed this self-assembly phenomenon. Our demo—which seeks to exemplify

during which particles form molecule-like assemblies that subsequently appear to dynamically solve jigsaw-like puzzles while en route to large-scale particulate discs.

Figure 1a depicts a time-series sequence of the motion of a particulate molecule¹ that eventually merges with a larger aggregate, and Figure 1b portrays a schematic of the experimental setup. Figures 1c and 1d present an initial state of the system and a final state 200 minutes later. Our experiments revealed this self-assembly phenomenon. Our demo—which seeks to exemplify

of physical systems, including the ubiquitous stratified ocean waters, potential microscale organization mechanisms, magnetic dynamics on Mercury and Ganymede, and even organic molecules such as matter's self-organization in deep oceans.

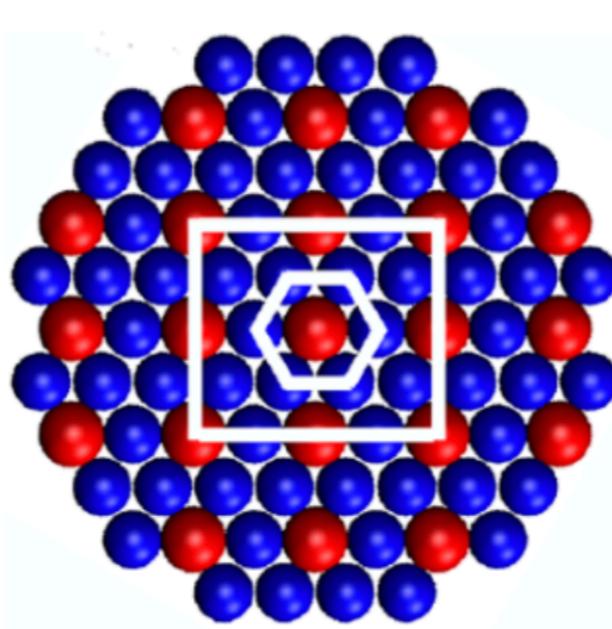
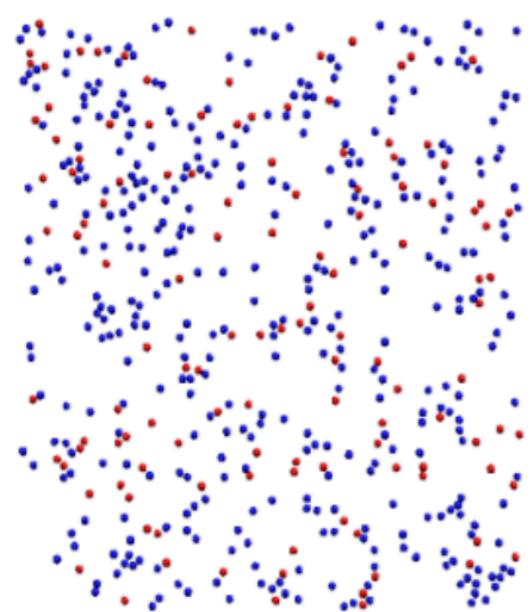
The discovery itself arose through a fortunate series of laboratory mistakes that ultimately led to a "eureka" moment. During a routine outreach demonstration, a setup error revealed this self-assembly phenomenon. Our demo—which seeks to exemplify

See Particle Aggregation on page 5

Figure 1. Experimental self-assembly. **1a.** Time-series montage that illustrates the formation of a molecule. Five different output times—with 10 minutes between each instance—appear from left to right (artistic blurring depicts the motion). The spheres' radii and densities are respectively $0.025\text{--}0.05\text{ cm}$ and 1.05 g cc^{-3} ; the top fluid is fresh water of density 0.997 g cc^{-3} and the bottom is a sodium chloride water solution of density 1.01 g cc^{-3} , with a transition thickness of approximately 2 cm . **1b.** Schematic of the experimental setup. **1c.** Initial cluster.

1d. Final self-assembled cluster. Figure adapted from [7].

Colloidal SA Case Study 1: Model Based



Ordered structure

Typical state: $\langle C_6 \rangle \in [0, 6]$

Averaged order parameter

Average number of hexagonally close-packed neighboring particles in 2D assembly

~ measure of crystallinity order

Dispersed particles

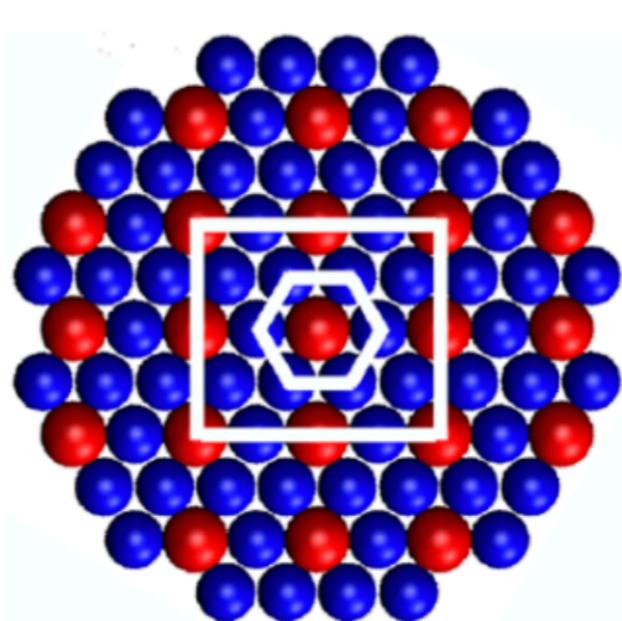
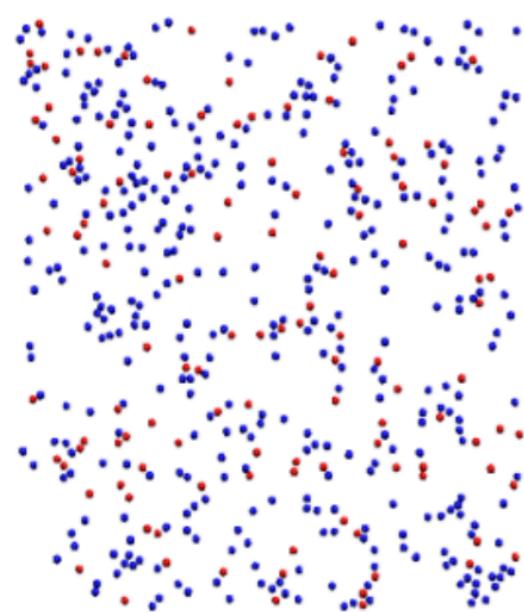
Typical control: u

Electric field voltage

Technical challenge:

Nonlinear + noisy molecular dynamics $\rightsquigarrow \langle C_6 \rangle$ is a controlled stochastic process

Colloidal SA Case Study 2: Data Driven



Ordered structure

Dispersed particles

Typical state: $(\langle C_{10} \rangle, \langle C_{12} \rangle) \in [0, 1]^2$

Steinhart bond order parameters

useful for distinguishing between
BCC and FCC structures

Typical control:

$(u_1, u_2) = (\text{temperature}, \text{pressure})$

Technical challenge:

Difficult to deduce first principle physics-based controlled dynamics over $(\langle C_{10} \rangle, \langle C_{12} \rangle)$

Controlled SA as Generalized Schrödinger Bridge

Intuition for Case Study 1: $\langle C_6 \rangle \approx 0 \Leftrightarrow$ Crystalline disorder

$\langle C_6 \rangle \approx 6 \Leftrightarrow$ Crystalline order

↷ Steer the stochastic state $\langle C_6 \rangle$ from disordered at $t = 0$ to ordered at $t = T$

In typical applications, prescribed time horizon 200 s

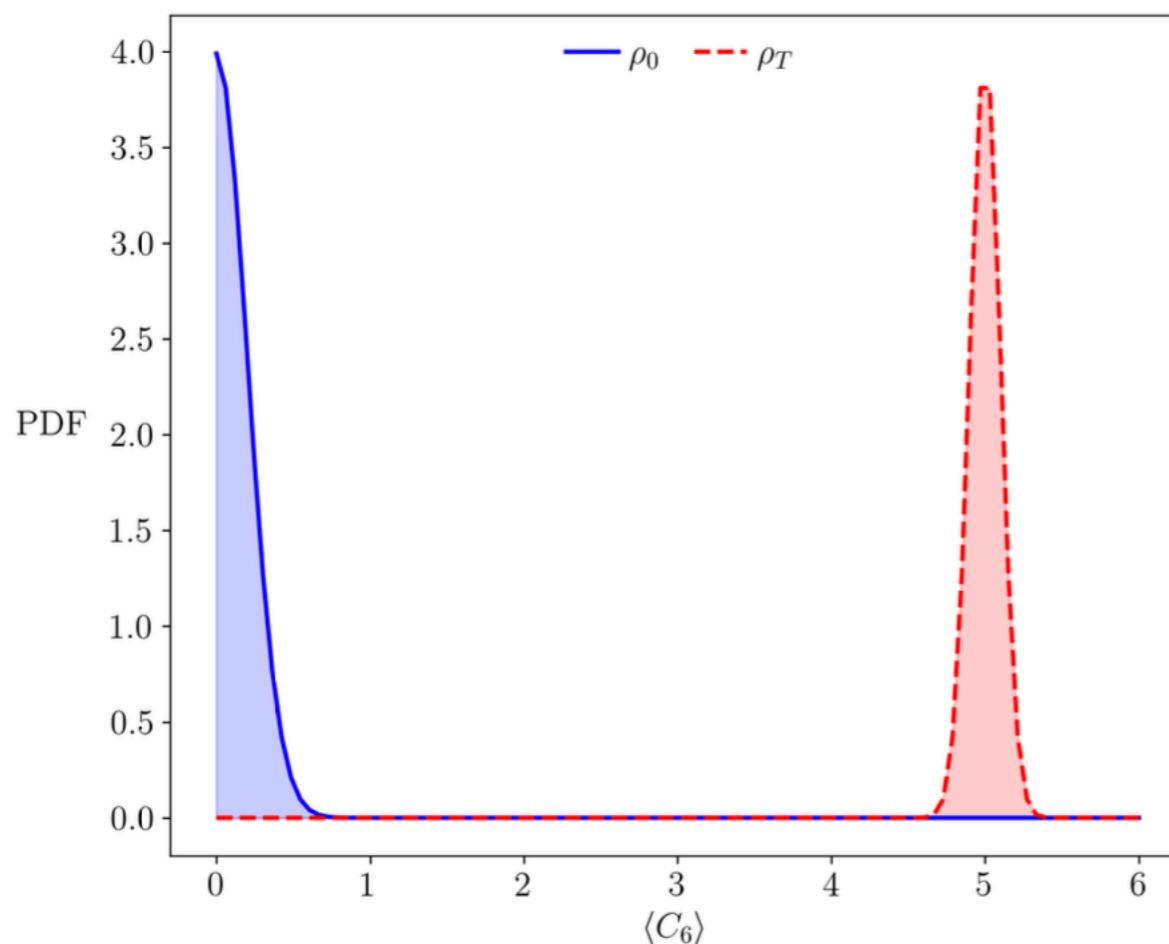
Controlled SA as Generalized Schrödinger Bridge

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In typical SA applications, prescribed time horizon 200 s



Endpoint PDF constraints:

$\langle C_6 \rangle(t = 0) \sim \rho_0$ given

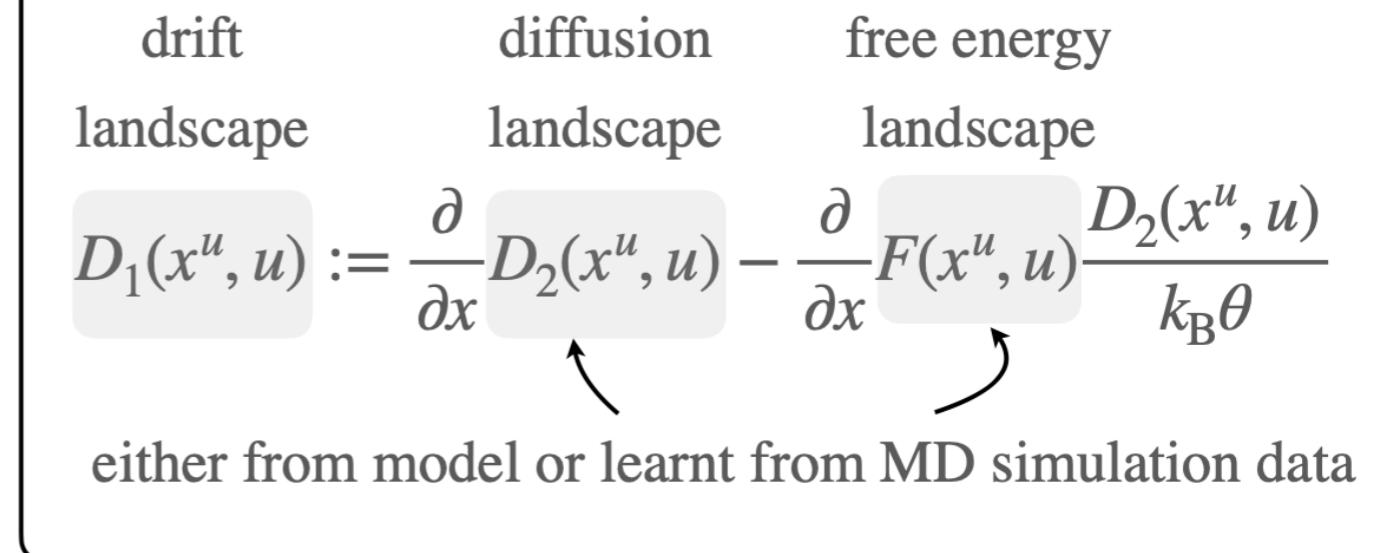
$\langle C_6 \rangle(t = T) \sim \rho_T$ given

Wanted control policy: $u = \pi(t, \langle C_6 \rangle)$

Minimum Effort Colloidal SA

Proposed formulation:

$$\inf_{u \in \mathcal{U}} \mathbb{E}_{\mu^u} \left[\int_0^T \frac{1}{2} u^2 dt \right],$$



subject to $dx^u = D_1(x^u, u) dt + \sqrt{2D_2(x^u, u)} dw,$

$\curvearrowleft \langle C_6 \rangle$ $\curvearrowleft \text{standard Wiener process}$

$$x^u(t=0) \sim d\mu_0 = \rho_0 dx^u, \quad x^u(t=T) \sim d\mu_T = \rho_T dx^u$$

Minimum Effort Colloidal SA

Equivalent formulation:

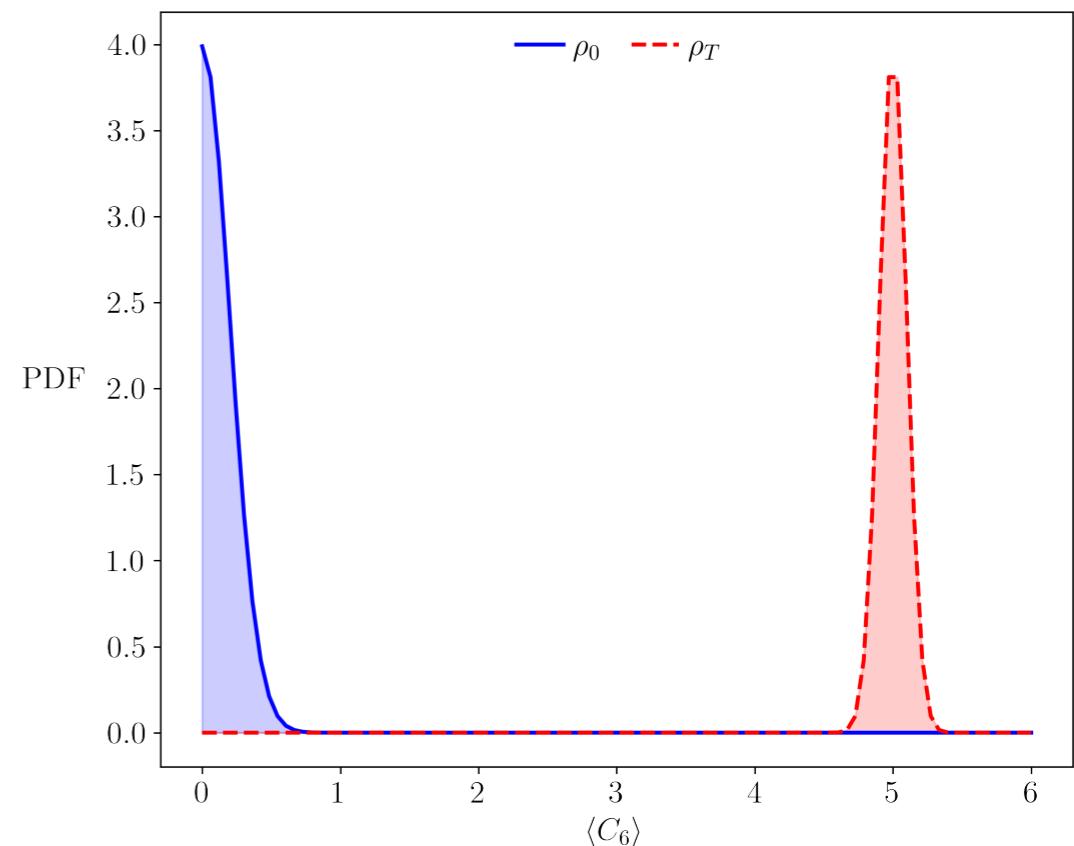
$$\inf_{(\rho^u, u)} \int_0^T \int_{\mathbb{R}} \frac{1}{2} u^2(x^u, t) \rho^u(x^u, t) dx^u dt$$

subject to $\frac{\partial \rho^u}{\partial t} = - \frac{\partial}{\partial x^u} (D_1 \rho^u) + \frac{\partial^2}{\partial x^{u2}} (D_2 \rho^u)$

$$\rho^u(x^u, t = 0) = \rho_0, \quad \rho^u(x^u, t = T) = \rho_T$$

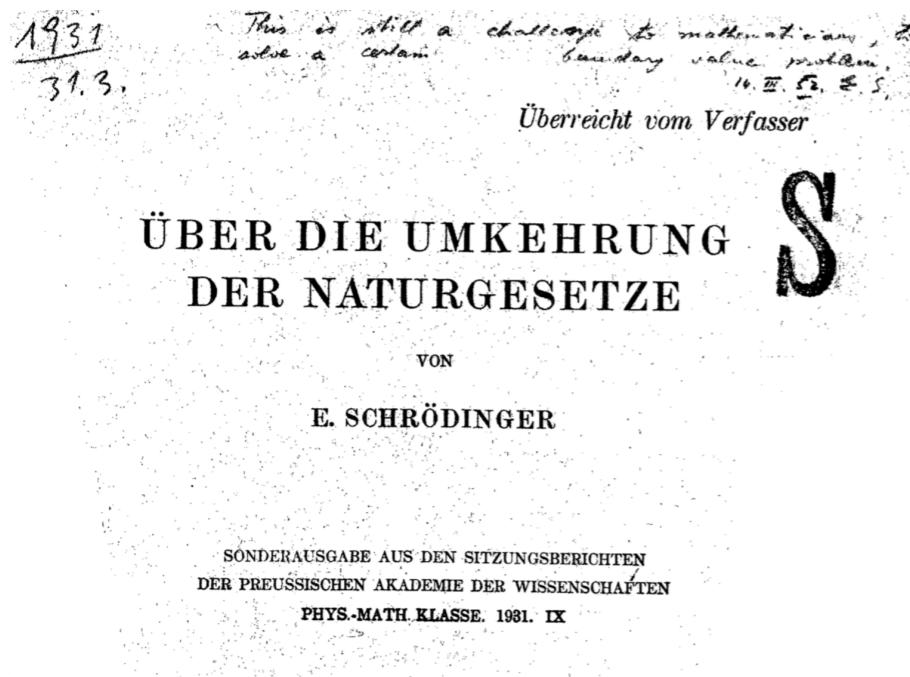
Controlled Fokker-Planck-Kolmogorov PDE

Guaranteed existence-uniqueness
for compactly supported ρ_0, ρ_T



Generalized Schrödinger Bridge Problem (GSBP)

Classical SBP: $D_1 \equiv u$, $D_2 \equiv \text{Identity}$



ÜBER DIE UMKEHRUNG DER NATURGESETZE

VON

E. SCHRÖDINGER

SONDERAUSGABE AUS DEN SITZUNGSBERICHTEN
DER PREUSSISCHEN AKADEMIE DER WISSENSCHAFTEN
PHYS.-MATH. KLASSE. 1931. IX

S

Sur la théorie relativiste de l'électron
et l'interprétation de la mécanique quantique

PAR

E. SCHRÖDINGER

I. — Introduction

J'ai l'intention d'exposer dans ces conférences diverses idées concernant la mécanique quantique et l'interprétation qu'on en donne généralement à l'heure actuelle ; je parlerai principalement de la théorie quantique relativiste du mouvement de l'électron. Autant que nous pouvons nous en rendre compte aujourd'hui, il semble à peu près sûr que la mécanique quantique de l'électron, sous sa forme idéale, que nous ne possédons pas encore, doit former un jour la base de toute la physique. A cet intérêt tout à fait général, s'ajoute, ici à Paris, un intérêt particulier : vous savez tous que les bases de la théorie moderne de l'électron ont été posées à Paris par votre célèbre compatriote Louis de BROGLIE.

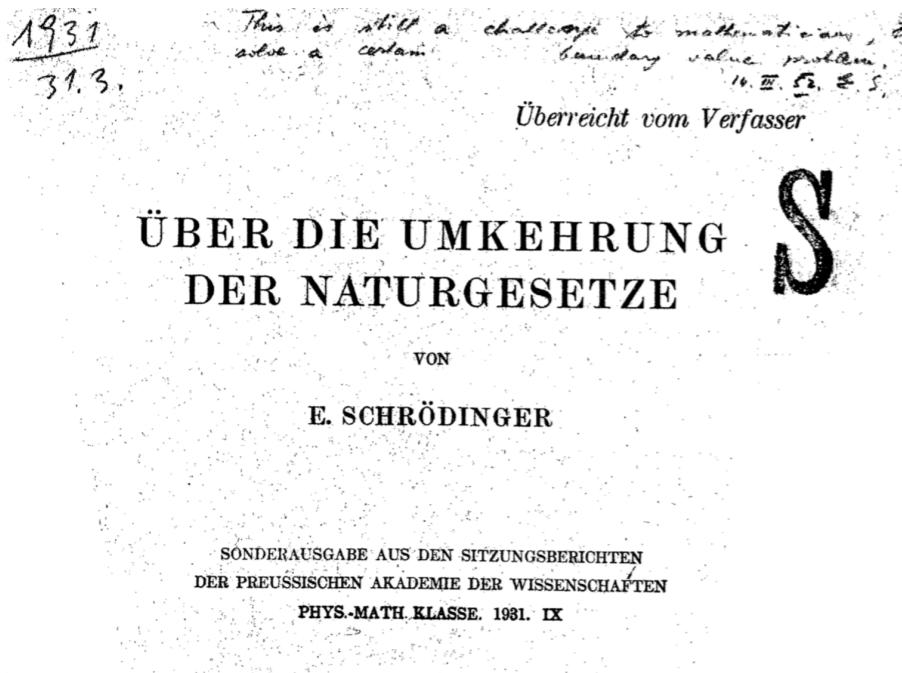


Classical SBP \equiv Stochastic version of dynamic OMT

Minimum effort \equiv Most likely evolution between observed distributions

Generalized Schrödinger Bridge Problem (GSBP)

Classical SBP: $D_1 \equiv u$, $D_2 \equiv \text{Identity}$



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Classical SBP \equiv Stochastic version of dynamic OMT

Minimum effort \equiv Most likely evolution between observed distributions

In our colloidal SA:

Both D_1, D_2 are nonlinear in state + non-affine in control

Conditions for Optimality: Case 1

Three coupled PDEs with endpoint boundary conditions:

$\frac{\partial \psi}{\partial t} = \frac{1}{2} (\pi^{\text{opt}})^2 - \frac{\partial \psi}{\partial x} D_1 - \frac{\partial^2 \psi}{\partial x^{u2}} D_2$	HJB PDE
$\frac{\partial \rho^u}{\partial t} = - \frac{\partial}{\partial x^u} (D_1 \rho^u) + \frac{\partial^2}{\partial x^{u2}} (D_2 \rho^u)$	Controlled FPK PDE
$\pi^{\text{opt}}(x^u, t) = \frac{\partial \psi}{\partial x^u} \frac{\partial D_1}{\partial u} + \frac{\partial^2 \psi}{\partial x^{u2}} \frac{\partial D_2}{\partial u}$	Optimal policy
$\rho^u(x^u, t = 0) = \rho_0, \quad \rho^u(x^u, t = T) = \rho_T$	Boundary conditions

value function	optimally controlled PDF	optimal policy
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to be solved for the triple: $\psi(x^u, t), \rho^u(x^u, t), \pi^{\text{opt}}(x^u, t)$

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HJB PDE

$$\frac{\partial \rho^u}{\partial t} = - \frac{\partial}{\partial x^u} (D_1 \rho^u) + \frac{\partial^2}{\partial x^{u2}} (D_2 \rho^u)$$

Controlled FPK PDE

$$\pi^{\text{opt}}(x^u, t) = \frac{\partial \psi}{\partial x^u} \frac{\partial D_1}{\partial u} + \frac{\partial^2 \psi}{\partial x^{u2}} \frac{\partial D_2}{\partial u}$$

Optimal policy

$$\rho^u(x^u, t = 0) = \rho_0, \quad \rho^u(x^u, t = T) = \rho_T$$

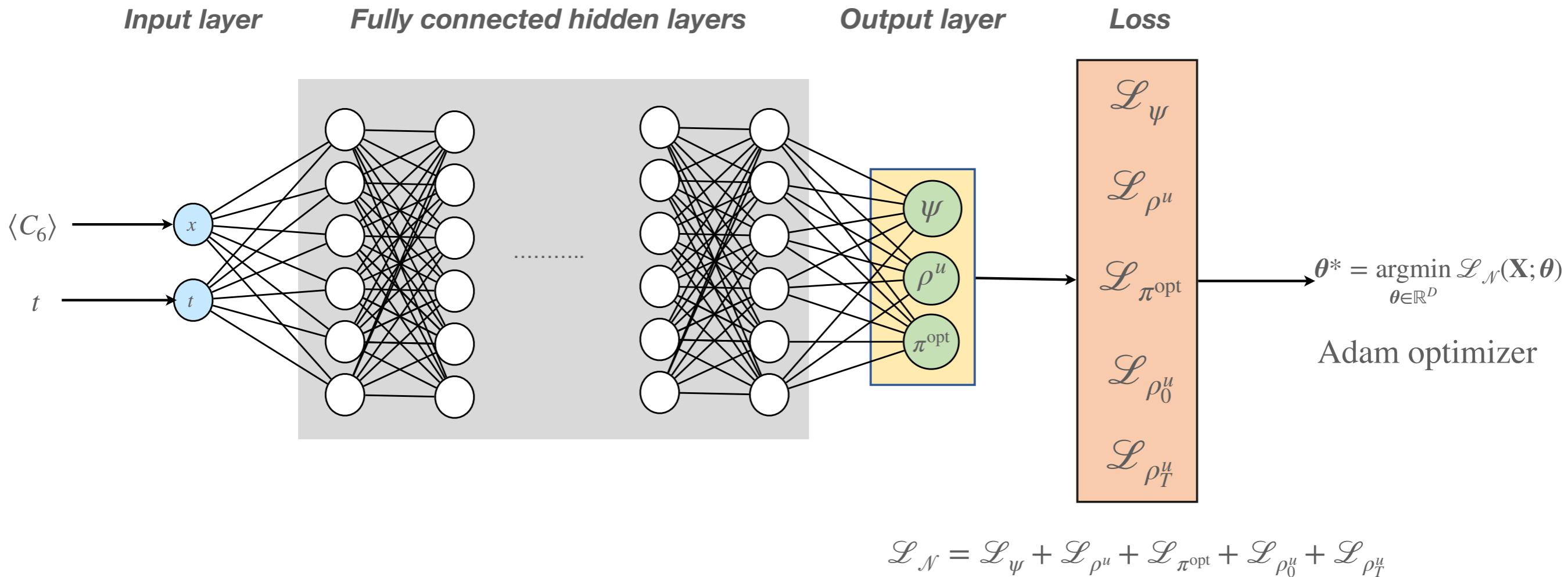
Boundary conditions

value function	optimally controlled PDF	optimal policy
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Cf. classical SBP: two coupled PDEs + optimal policy explicit in value fn ψ

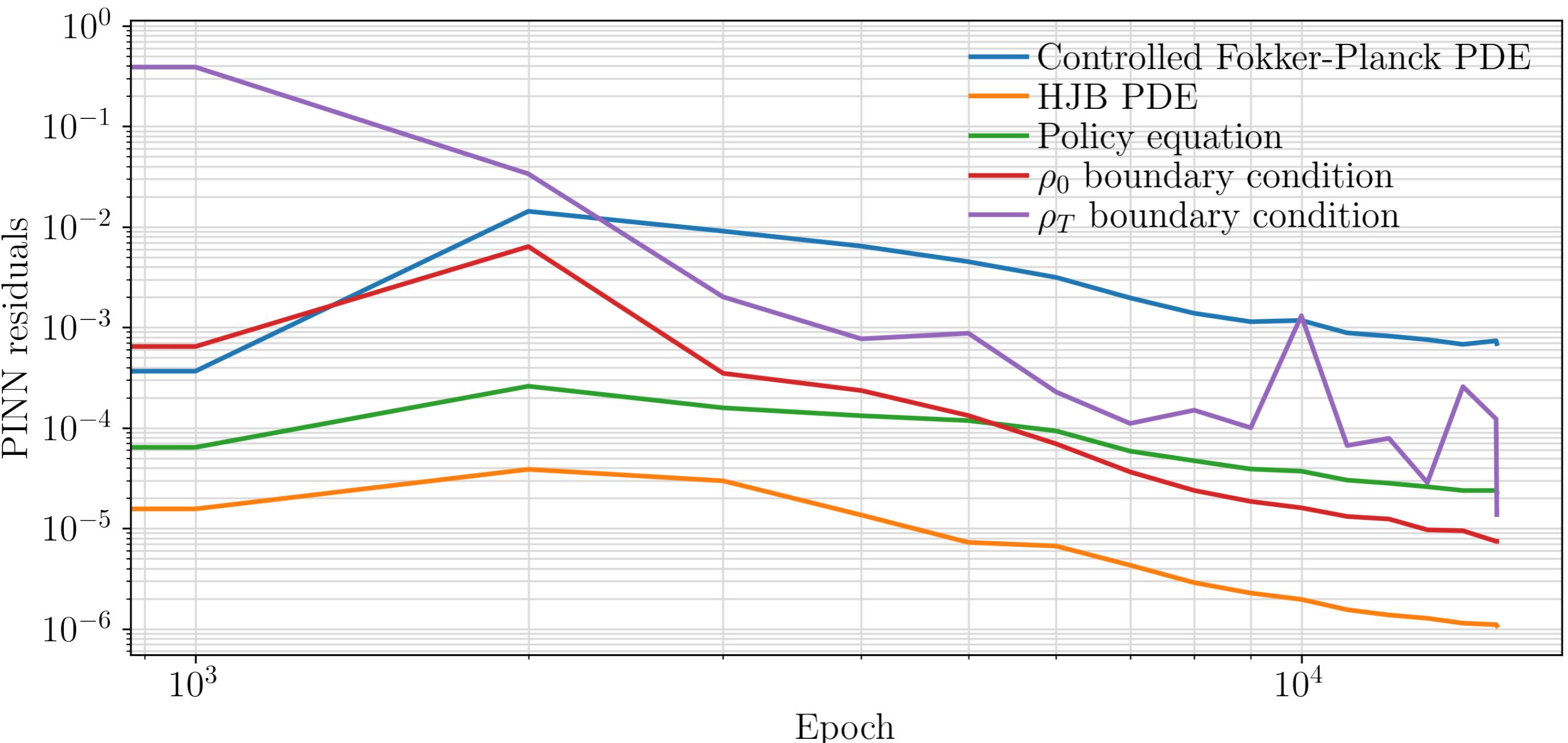
Idea: Train Physics Informed Neural Network (PINN) to Learn the Solution of the GSBP



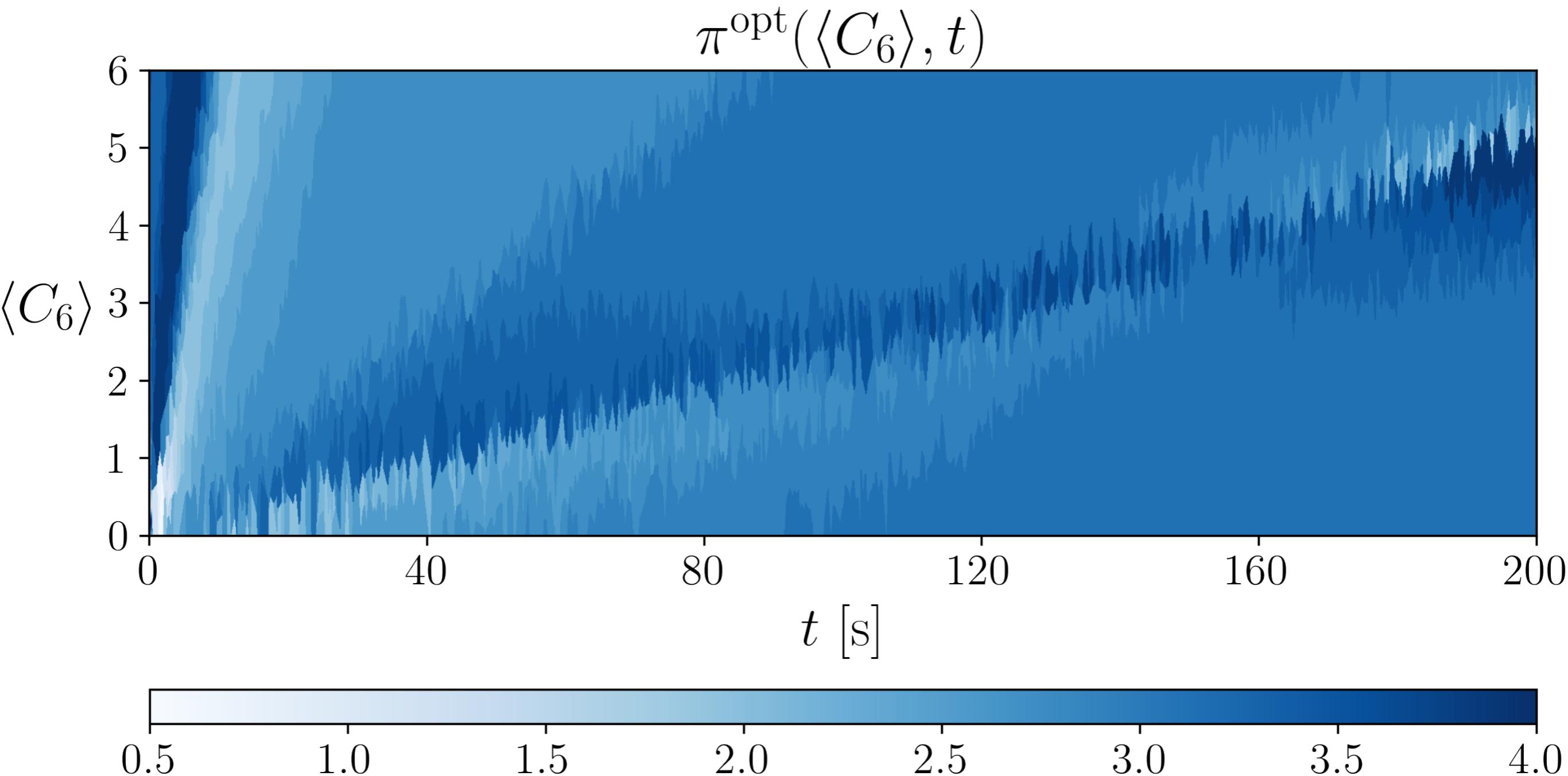
[Lu Lu, et al, 2021] [Niaki, et al, 2021]

Case Study 1: Residuals for PINN Training

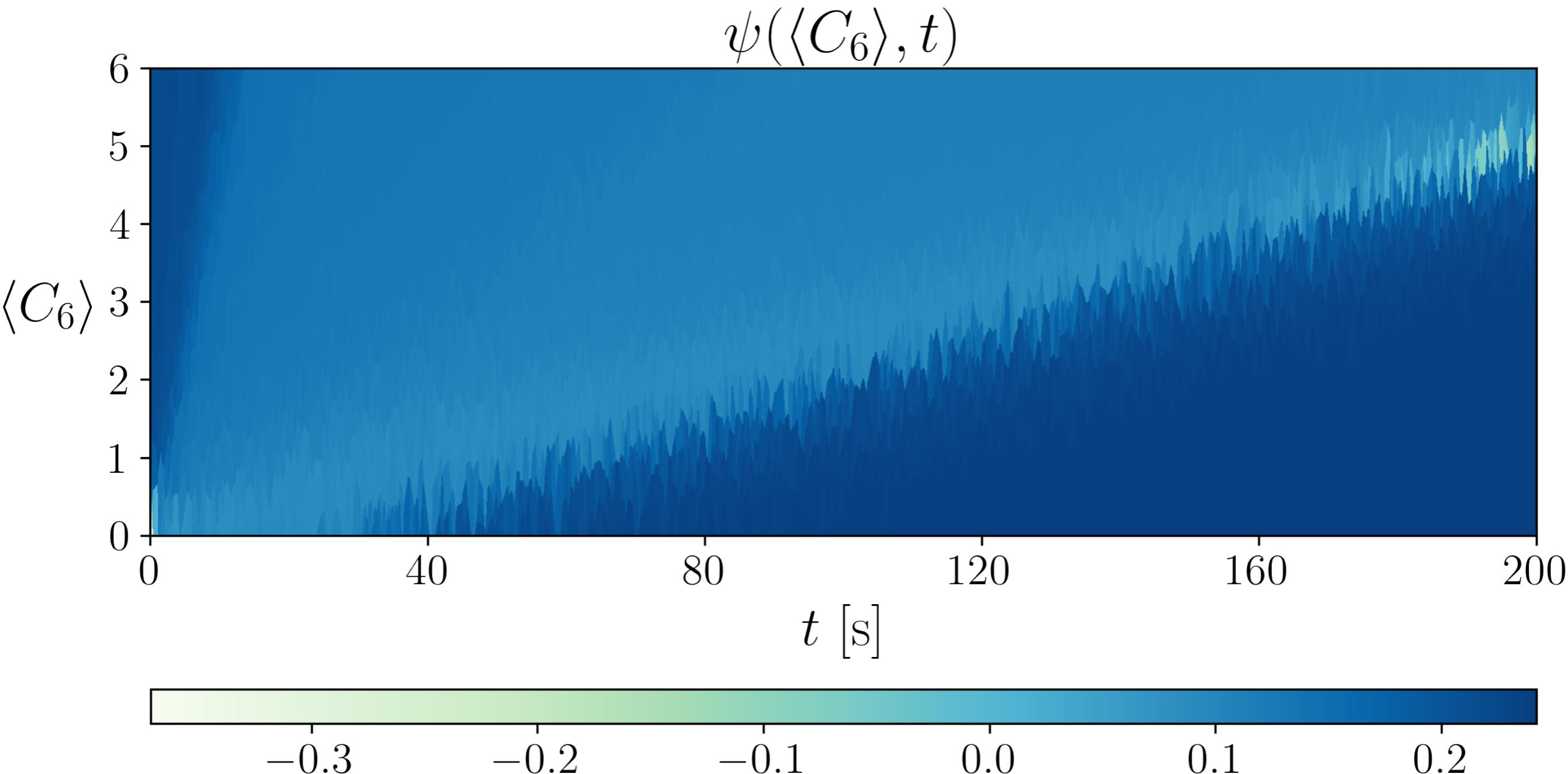
Benchmark controlled self-assembly system: [Y Xue, et al, *IEEE Trans. Control Sys. Technology*, 2014]



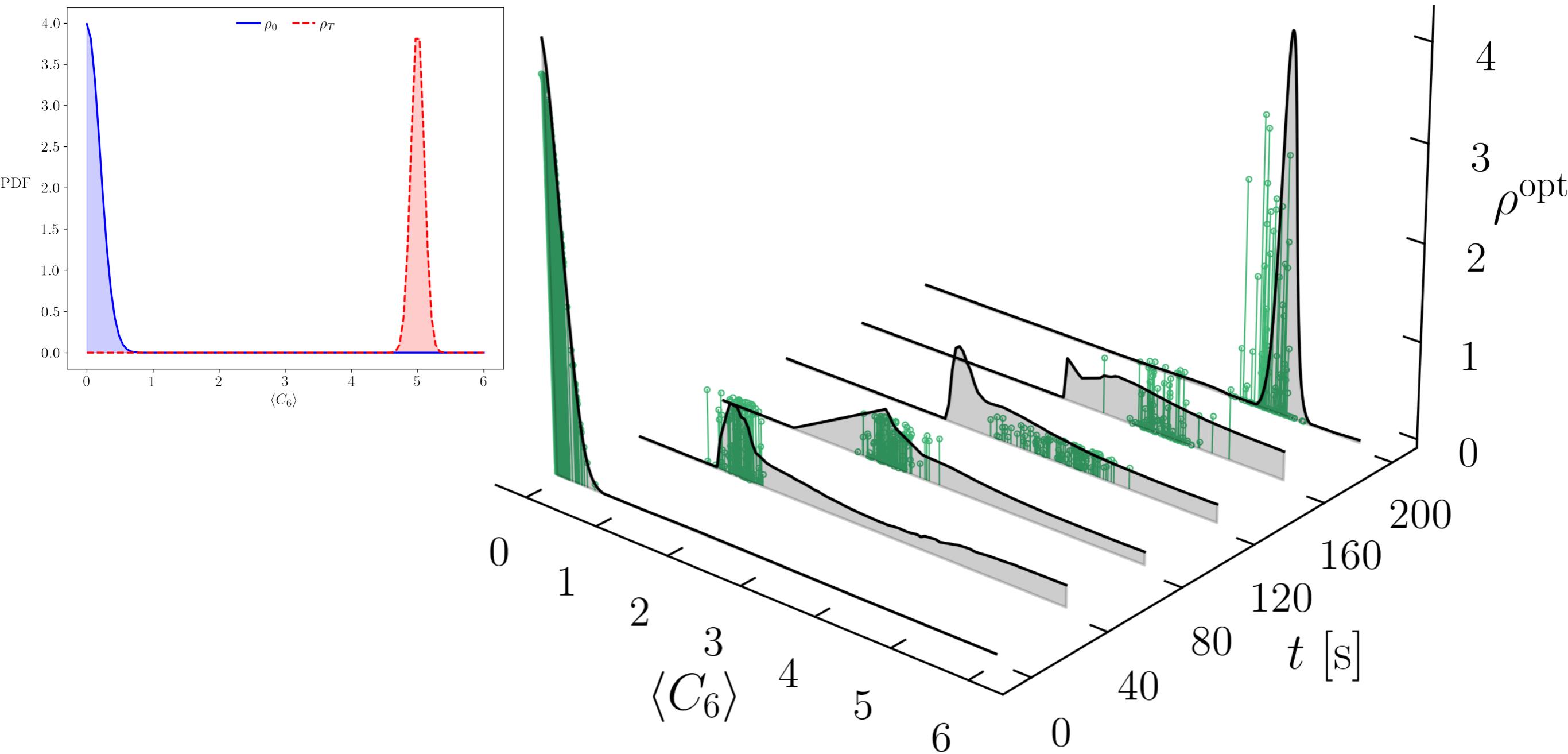
Case Study 1: Optimal Policy



Case Study 1: Value Function

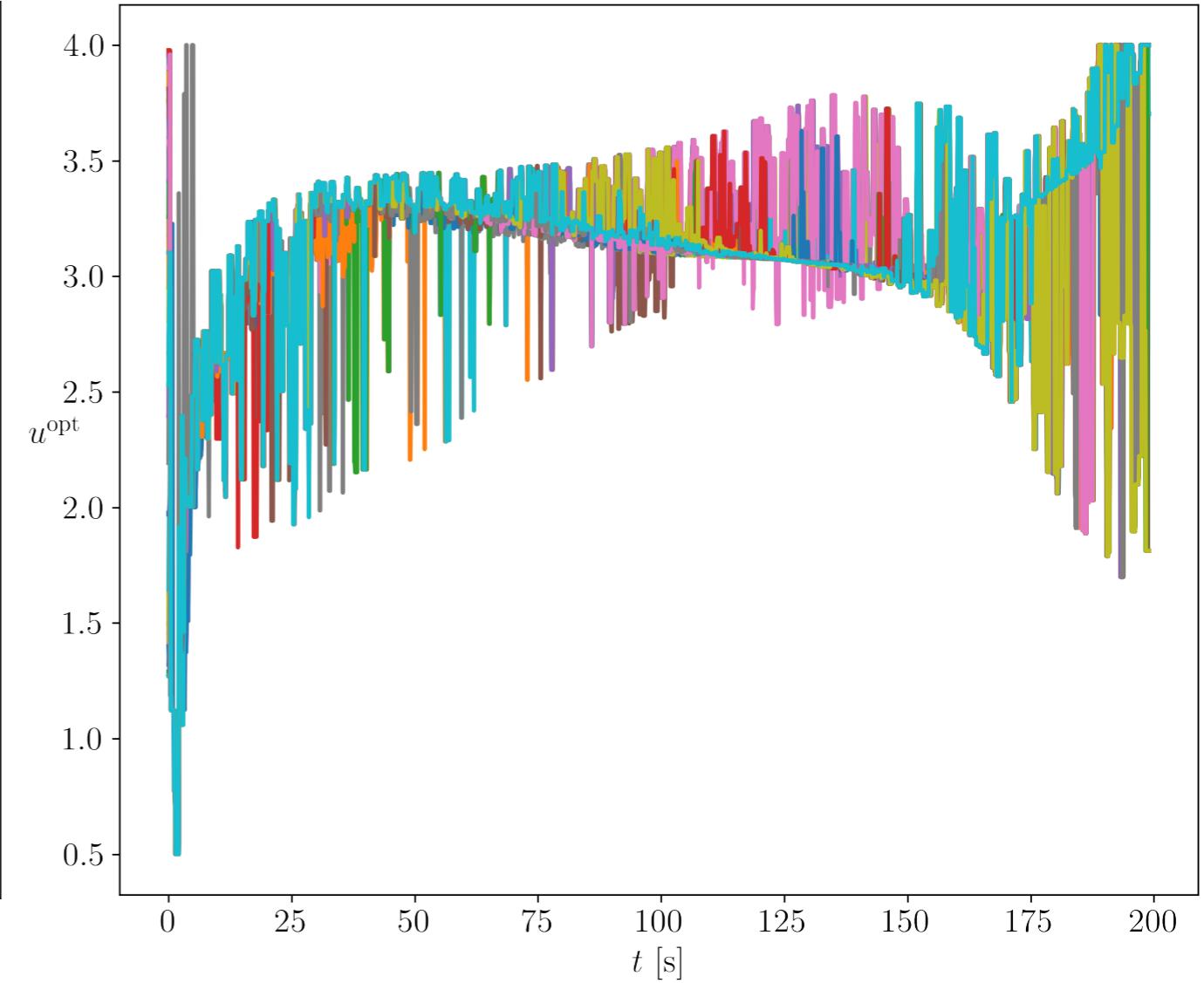
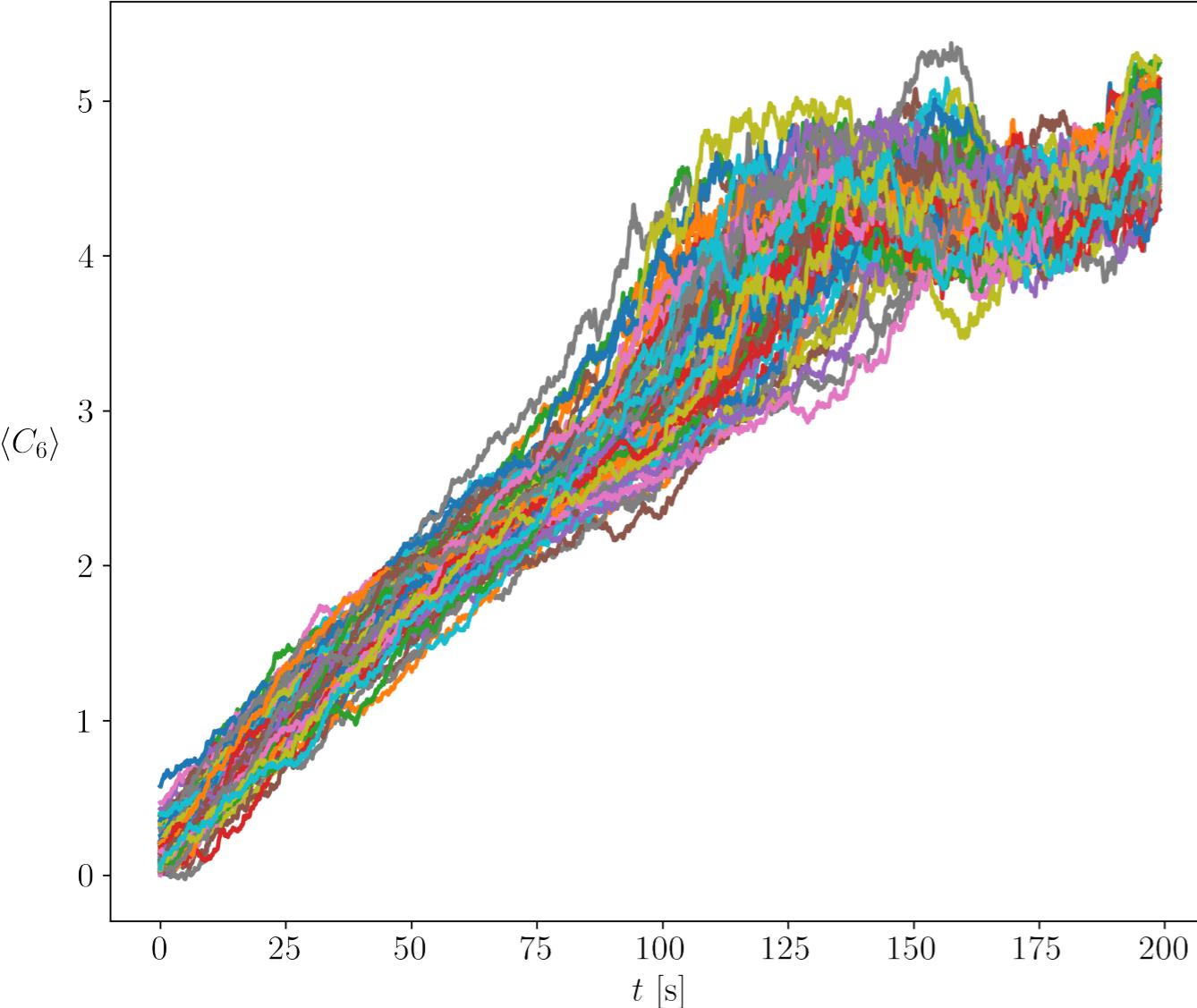


Case Study 1: Optimally Controlled State PDFs



... the MSE losses are not appropriate for enforcing the endpoint PDF constraints

Case Study 1: Closed Loop Sample Paths



GSBP Conditions for Optimality with m Inputs

$m + 2$ coupled PDEs with endpoint boundary conditions:

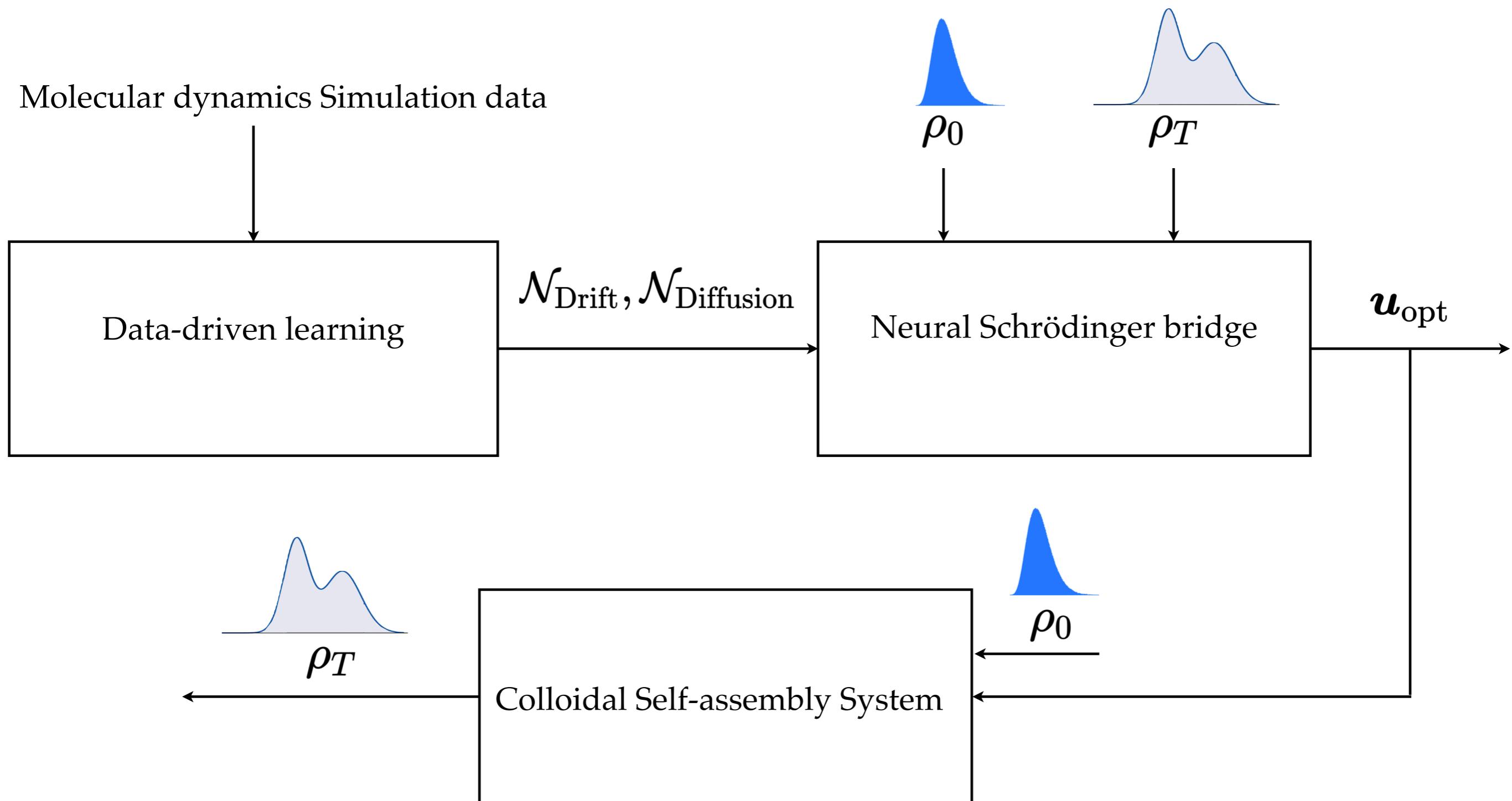
$$\begin{aligned} \frac{\partial \psi}{\partial t} &= \frac{1}{2} \|u_{\text{opt}}\|_2^2 - \langle \nabla_x \psi, f \rangle - \langle G, \text{Hess}(\psi) \rangle, \\ \frac{\partial \rho_{\text{opt}}^u}{\partial t} &= -\nabla \cdot (\rho_{\text{opt}}^u f) + \langle G, \text{Hess}(\rho_{\text{opt}}^u) \rangle, \\ u_{\text{opt}} &= \nabla_{u_{\text{opt}}} (\langle \nabla_x \psi, f \rangle + \langle G, \text{Hess}(\psi) \rangle), \\ \rho_{\text{opt}}^u(0, x) &= \rho_0, \quad \rho_{\text{opt}}^u(T, x) = \rho_T, \end{aligned}$$

Drift coefficient Diffusion tensor

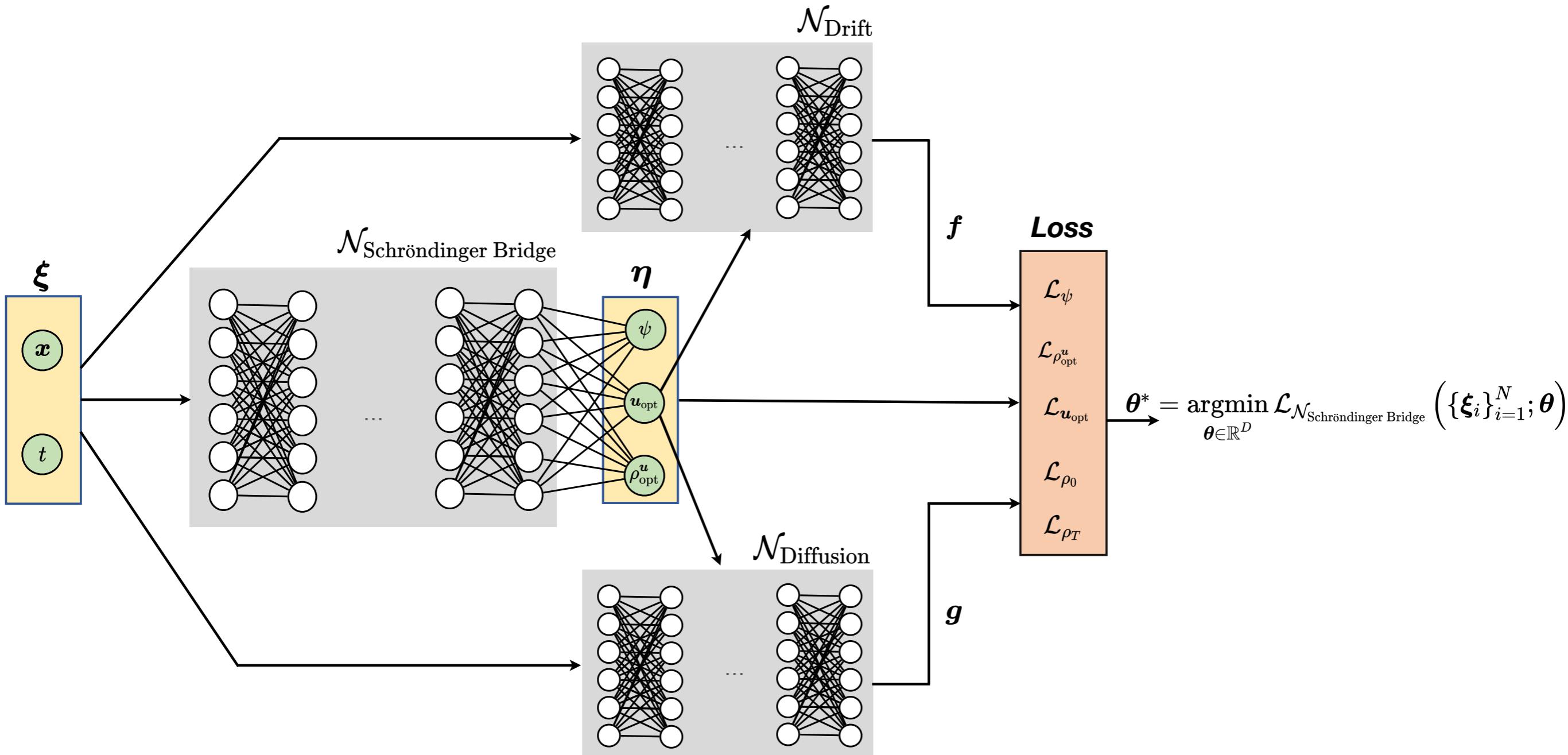
The diagram consists of two arrows. One arrow points downwards from the text 'Drift coefficient' to the term f in the first equation. Another arrow points diagonally upwards and to the right from the text 'Diffusion tensor' to the term G in the second equation.

Cf. classical SBP: two coupled PDEs + optimal policy explicit in value fn ψ

Data-driven GSBP for Colloidal SA



Architecture for Data-driven GSBP



Sinkhorn Losses for Boundary Conditions

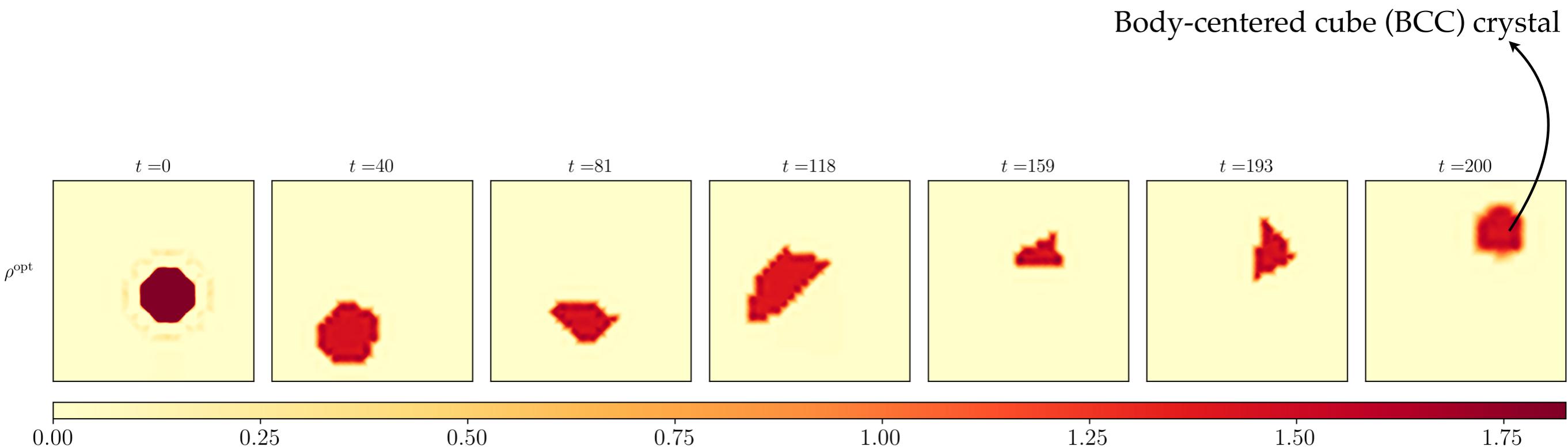
$$W_\varepsilon^2(\mu_0, \mu_1) := \inf_{\pi \in \Pi_2(\mu_0, \mu_1)} \int_{\mathbb{R}^n \times \mathbb{R}^n} \{\|\mathbf{x} - \mathbf{y}\|_2^2 + \varepsilon \log \pi(\mathbf{x}, \mathbf{y})\} d\pi(\mathbf{x}, \mathbf{y})$$

For boundary conditions, use Sinkhorn losses: $\mathcal{L}_{\rho_i} := W_\varepsilon^2\left(\rho_i, \rho_i^{\text{epoch index}}(\boldsymbol{\theta})\right)$

Implementation friendly for PINN training:

$$\text{Autodiff}_{\boldsymbol{\theta}} W_\varepsilon^2\left(\rho_i, \rho_i^{\text{epoch index}}(\boldsymbol{\theta})\right) \quad \forall i \in \{0, T\}$$

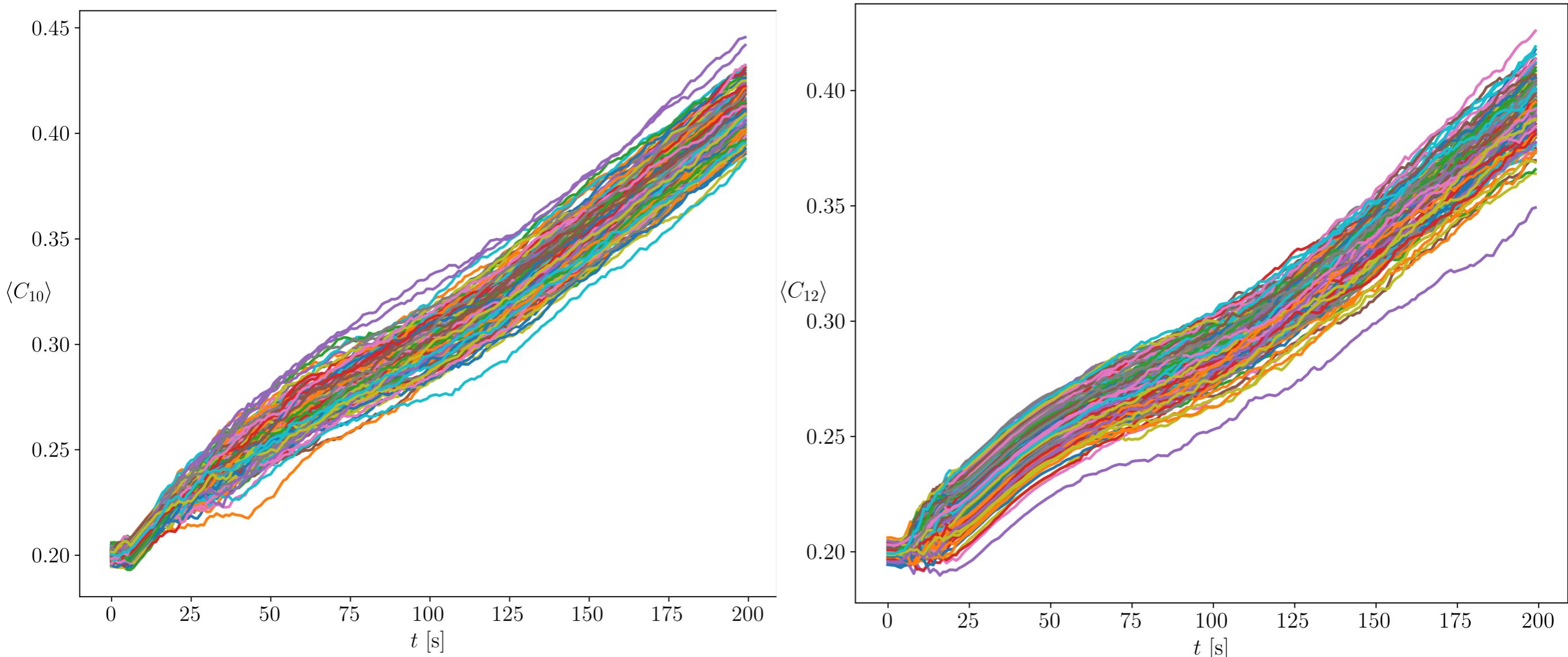
Case Study 2: Synthesize BCC Crystalline Structure by PDF Steering in $(\langle C_{10} \rangle, \langle C_{12} \rangle)$ Space



Data-driven:

Uses PINN with Sinkhorn losses + the drift-diffusion are themselves NNs

Case Study 2: Closed Loop State Sample Paths



Desired transport from mean $(0.2, 0.2)$ to $(0.40, 0.37)$ for BCC structure

Take Home Message

GSBPs arise quite naturally in engineering applications such as colloidal SA

Computational methods for GSBPs need more development

Refs for this work:

I. Nodozi, J. O'Leary, A. Mesbah, A.H., A physics-informed deep learning approach for minimum effort stochastic control of colloidal self-assembly, *ACC 2023*

I. Nodozi, C. Yan, M. Khare, A.H., A. Mesbah, Neural Schrödinger Bridge with Sinkhorn Losses: Application to Data-driven Minimum Effort Control of Colloidal Self-assembly, *arXiv????*

Thank You

Acknowledgement:



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