

#### Dynamics of the prion proliferation

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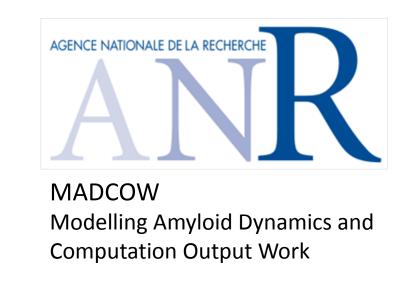
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# Dynamics of the prion proliferation





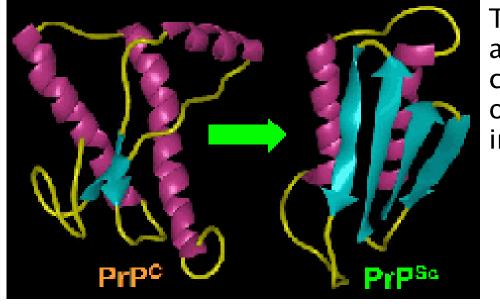
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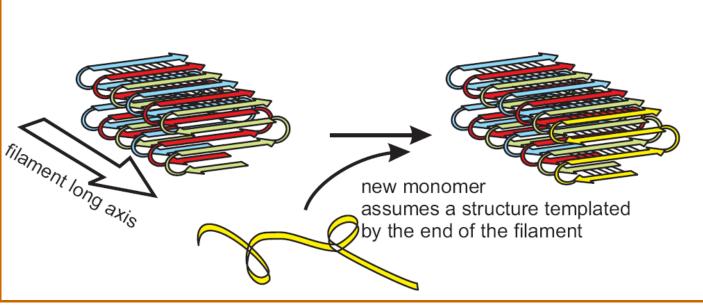
## TSE (Transmissible Spongiform Encephalopathies) and prion hypothesis

Prion protein is responsible for some of the transmissible spongiform encephalopathies diseases. These long-time incubation neurodegenerative dieases are infectious and always fatal. They are due to amyloid formation in the brain. S. Prusiner established infectiousity of the protein and the existence of two different conformational states. He received the Nobel Price in 1997.

Prion  $PrP^c$  is a protein naturally produced in a conventional configuration.



the protein. The latter is the



The appearance of the disease can have two causes :

- A sporadic (spontaneous) apparition of a misfolded protein.
- · A contamination by the infectious agent  $PrP^{Sc}$  , a seed or an amyloïd.

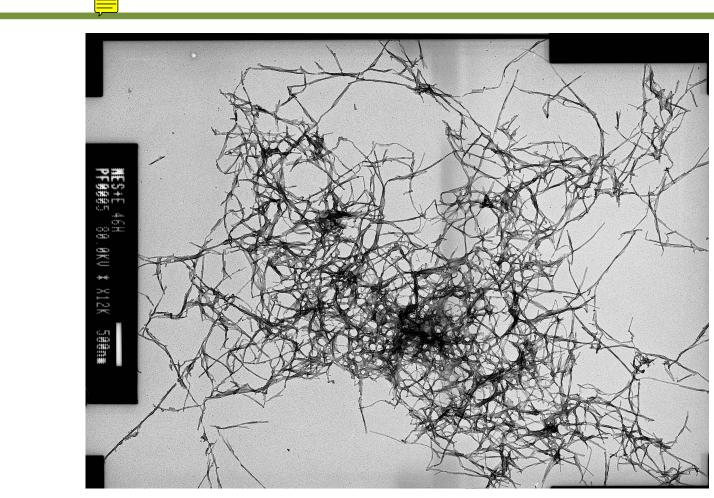
#### Well-known diseases:

- Bovine spongiform encephalopathy
- Scrapie in sheep Creutzfeldt-Jakob disease
- ·Kuru

0.4 mg/ml 0.4 mg/ml + 1% Agr B 0.4 mg/ml + 1% B 1P

0.4 mg/ml + 1% B 2P 0.4 mg/ml + 1% B 3P

**OUR GOAL**: some experiments suggest the existence of different strains. We would like to understand the replication mechanism and the sporadic appearance of amyloid to identify these strains. Our goal is to specify the parameters involved in the prion polymerization and determine prion strain according to these parameters.







The unconventional proteins  $PrP^{Sc}$  create polymers that aggregate in amyloids.

## Spontaneous polymerization experiment

Spontaneous polymerization starts 1 with an initially population of recombinant  $PrP^c$  Prion protein  $\S$ with continuous shaking.

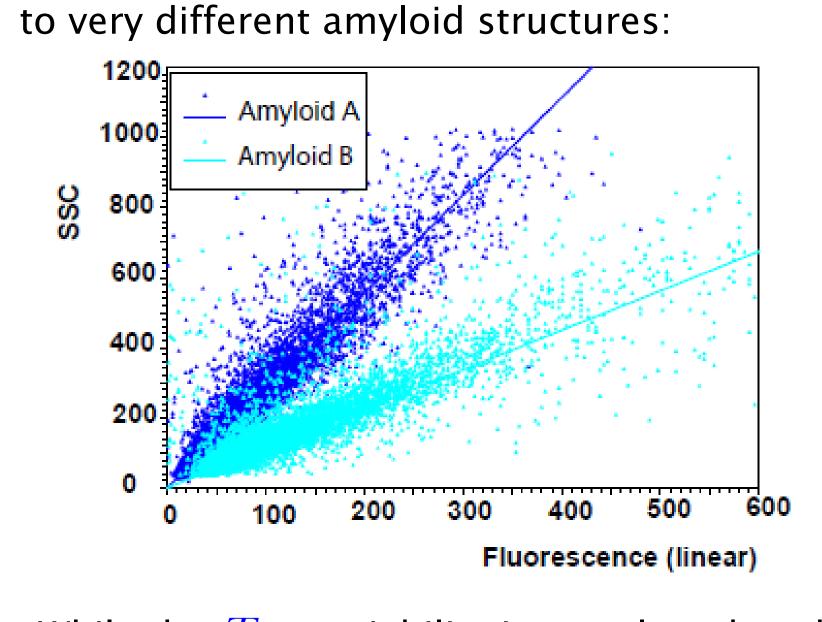
All experiments display a sigmoid shape,  $= y_0 + \frac{\alpha}{1 + e^{\frac{-2(t-T_i)}{\tau * a}}}$  $T_{laq}, \tau, \text{ and } F_{max}$ slope=  $\frac{1}{\tau}$ 

 $T_{lag}$  (10%)

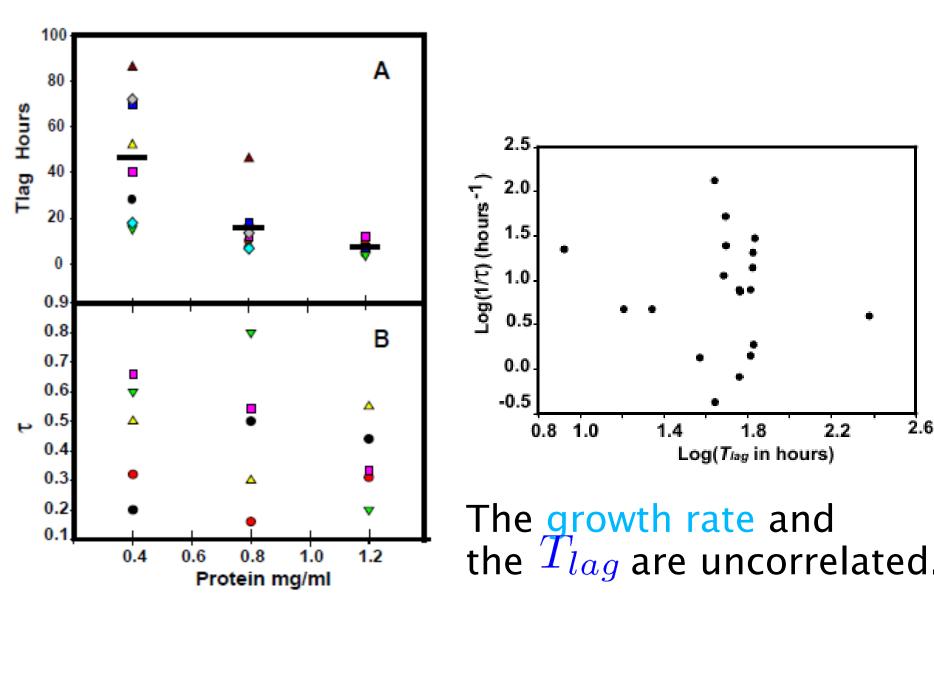
 $T_{lag} = T_i - \tau(y_0 + \frac{a}{2})$ 

Spontaneous polymerization can lead

with three main characteristics:

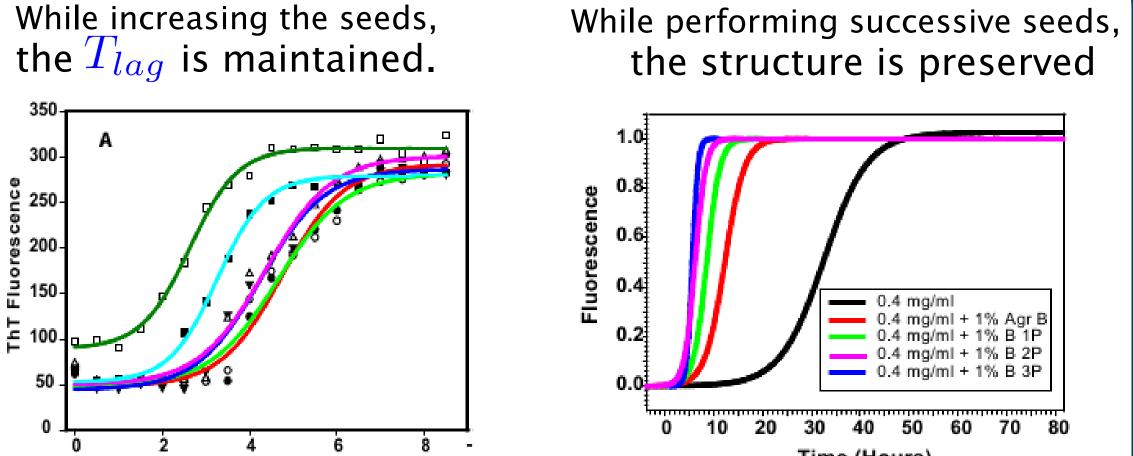


While the  $T_{lag}$  variability is greatly reduced with increased initial concentration, growth rate variability is maintained.



## Seeding experiment

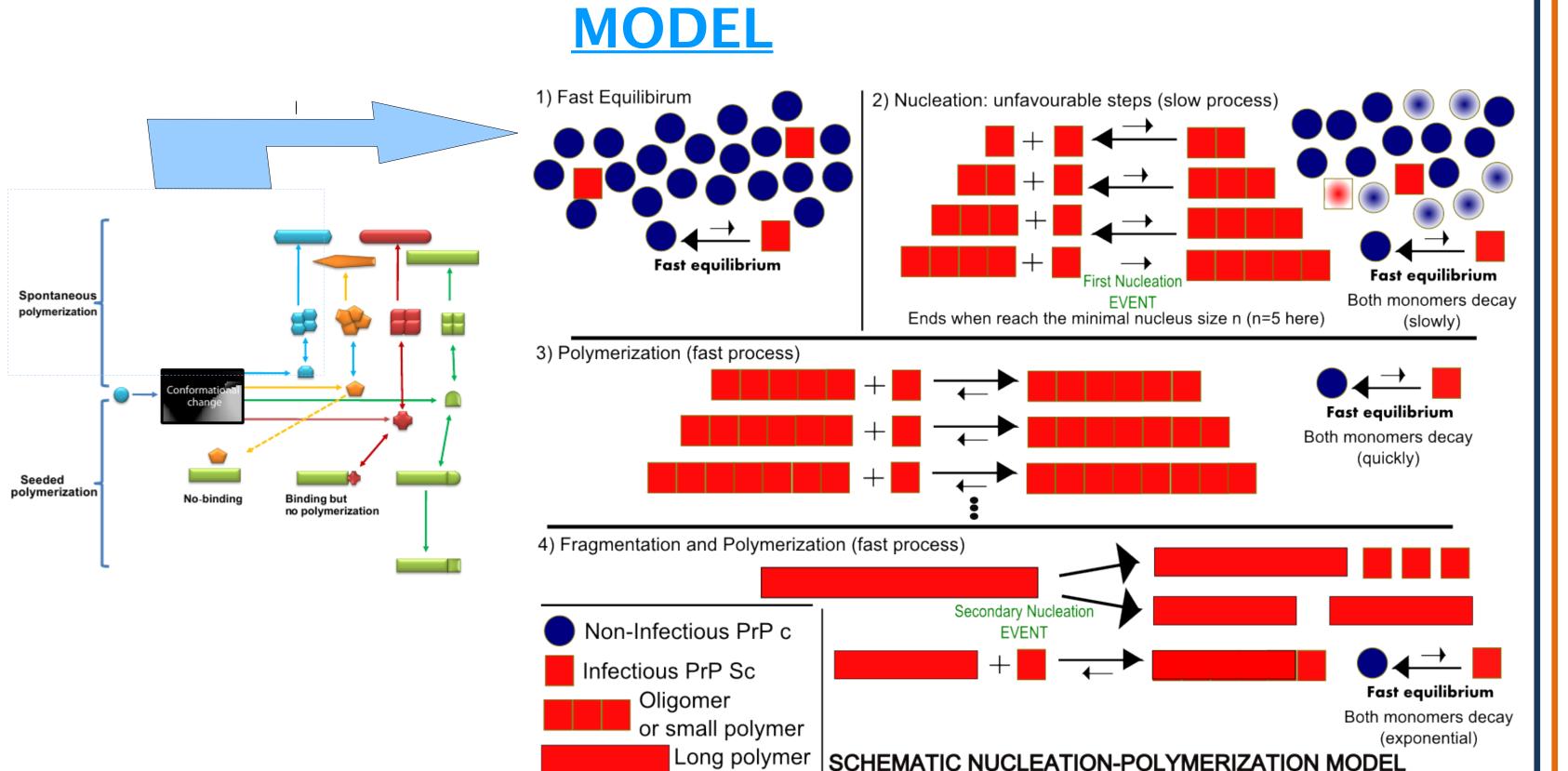
These experiments starts with pre-formed seeds.



In vitro Prion amyloids formation present a primarily conformational change step ans a nucleation-dependent polymerization.

 $u_i + f_1 \xrightarrow{k_{pf}} u_{i+1}, \ (i \ge n)$ 

 $u_i \xrightarrow{d_2} u_{i-1} + f_1, \ (i \ge n)$ 



# **HYPOTHESIS**

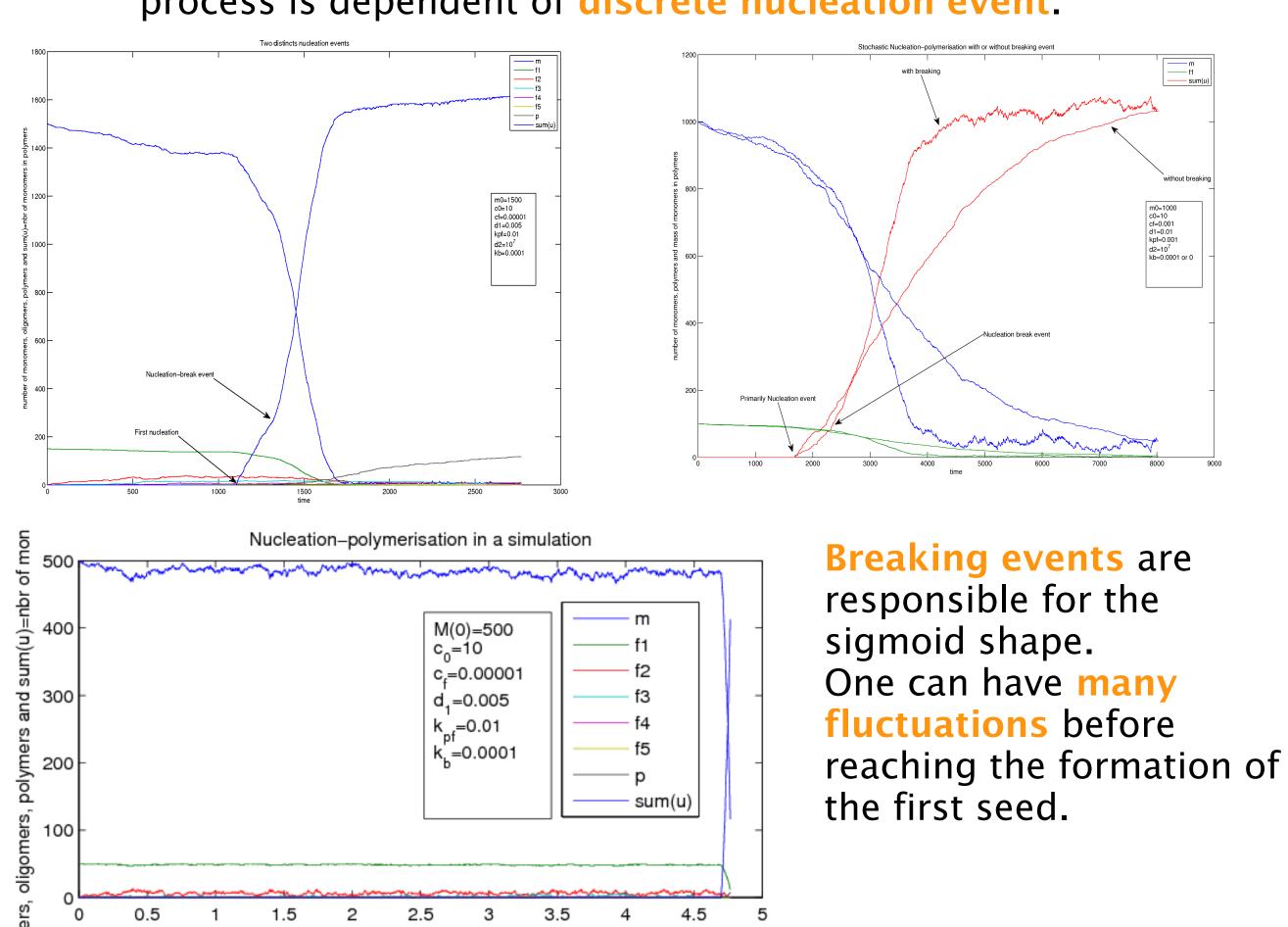
→ Conformational change (fast equilibrium) → Heterogeneous nucleus formation \_\_\_\_\_

→ Specific strain or amyloid

lag are uncorrelated.  $\rightarrow$  Specific polymerization, by binding to amyloid seed

→Fragmentation

Simulation was performed using a Gillespie algorithm. The modelexhibits sigmoid shape, and the polymerisation process is dependent of discrete nucleation event.

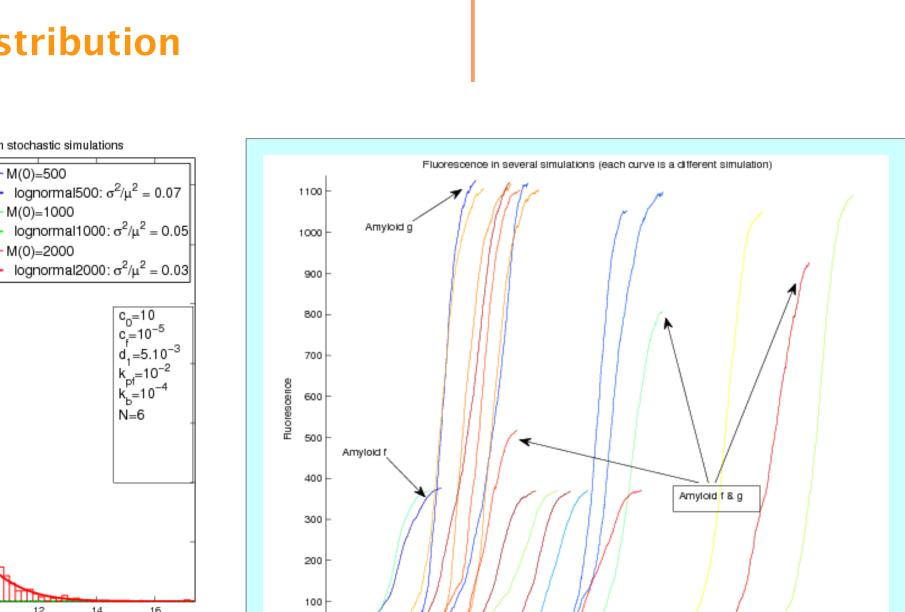


 $T_{lag}$  is much more dependent of the initial amount of protein than the growth rate. Both quantities are following a log-normal distribution

— lognormal2000:  $\sigma^2/\mu^2 = 0.01$ 

logmormal1000:  $\sigma^2/\mu^2 = 0.29$ 

• lognormal500:  $\sigma^2/\mu^2 = 2.12$ 

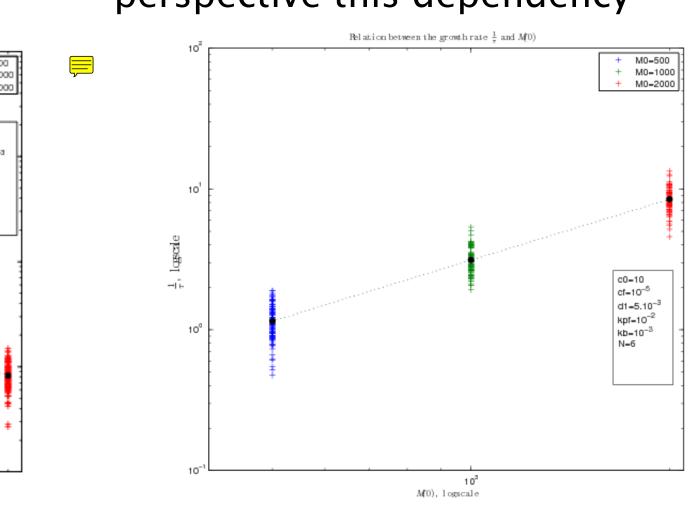


**Stochastic Nucleation-Polymerisation Model** 

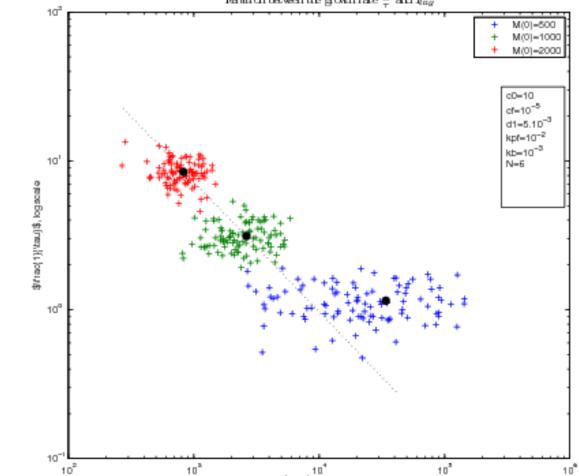
The evolution of  $T_{lag}$  while increased initial population is in good agreement with data.

Ristion between the  $T_{lag}$  and  $\Lambda(0)$ 

But the growth rate is also dependent of the initial population. This contradicts data, although the variability put into perspective this dependency



Both quantities are somewhat correlated, although here again variability could explain the experimental observations.



### **CONCLUSION:**

This preliminary work shows that a stochastic nucleation-polymerization model can explain the observed variability. In a stochastic description, discrete events are responsible for the overall dynamic. Variability can also be an artifact to some observable experimental laws.

#### **GOING FURTHER:**

•Analytical study will be performed to gain more insight into the effects of stochasticity on the dynamics. •General scaling laws such as the value of Lag Time (exit time problem) and the growth rate are expected. •Given this information and improving the algorithm could lead to precise date fitting.

•Interactions between different structures (strain) can then be of importance.

- lognormal500:  $\sigma^2/\mu^2 = 0.07$ 

lognormal2000:  $\sigma^2/\mu^2 = 0.03$ 

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