Efficient Multiscale Platelets Modeling Using Supercomputers

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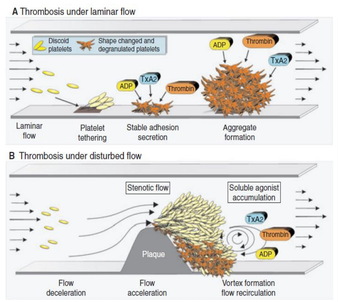
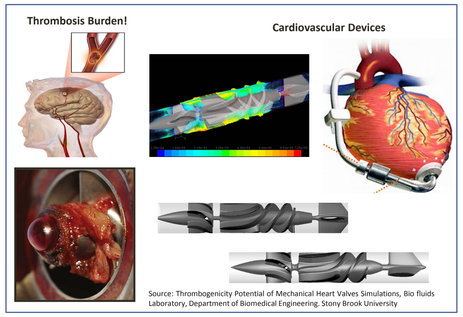
Motivation



Jackson et al., Dynamics of Platelet Thrombus Formation, Journal of Thrombus and Haemostasis, 2009

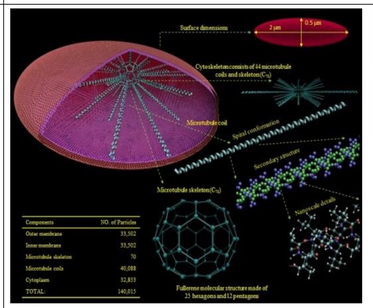
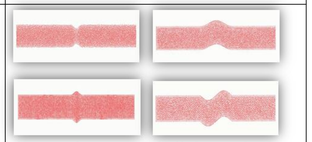
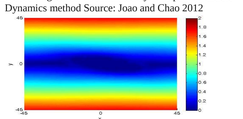
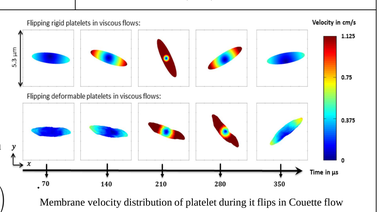


Source: Thrombogenicity Potential of Mechanical Heart Valves Simulations, Bio fluids Laboratory, Department of Biomedical Engineering. Stony Brook University

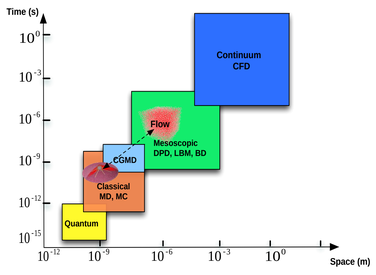
This objective of this project is to efficiently simulate flow-induced platelet activation in order to better understand thrombosis formation mechanisms. The methodologies include:

- Mathematical modeling of viscous blood flows and human platelet cell
- Multiscale coupling methods
- Data analysis of thermodynamic properties
- Algorithmic acceleration
- Heterogeneous computing
- Visualization

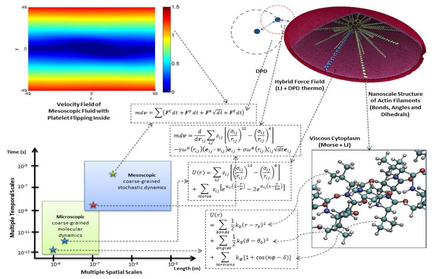
Multiscale Fluid-Platelet Models

Scales	Nanoscale	Mesoscale
Simulation Domain	Platelet Cell	Blood Plasma
Methods	Coarse-Grained Molecular Dynamics (CGMD)	Dissipative Particle Dynamics (DPD)
Time Step	10~100 *fs*	0.01~1 μs
Length	1~20 *A*	0.1 ~ 1 μm
Model Abstraction	 Physical structures and constitutes of Multiscale model of human platelets	 Various vascular geometries simulated by Dissipative Particle Dynamics method Source: Joao and Chao 2012  Couette flow by applying two counter body forces on all boundary particle; in such case, a uniform shear stress will be emulated; and platelet can flip inside such environment. It's the simplest fluid-platelet simulation setup
Force Fields	$V(r) = \sum k_b(r-r_0)^2 + \sum k_\theta(\theta-\theta_0)^2$ Bond and Angle Terms $+ \sum k_\phi[1+\cos(n\phi-\delta)] + \sum \frac{q_i q_j}{4\pi\varepsilon_0 r}$ Dihedral and Electrostatic $+ \sum 4\epsilon\left[\left(\frac{\sigma}{r}\right)^{12} - \left(\frac{\sigma}{r}\right)^{6}\right]$ Van Der Waals (L-J) $+ \sum \epsilon\left[\alpha\left(1-\frac{r}{R(\mu)}\right)-2\exp\left(\frac{\alpha}{2}\left(1-\frac{r}{R(\mu)}\right)\right)\right]$ Modified Morse V is the total energy on each particle composed of platelet. It includes a classical MD potential for describing the actin filament structure, a modified Morse potential for describing the viscous cytoplasm structures, and a CGMD for describing the filamentous core and the membrane structures	$\boldsymbol{F_{ij}} = \boldsymbol{F_{ij}^C} + \boldsymbol{F_{ij}^D} + \boldsymbol{F_{ij}^R}$ (Groot and Warren 1997) $\boldsymbol{F_{ij}^C} = \alpha\omega(r_{ij})\boldsymbol{e}_{ij}$ Conservative Term $\boldsymbol{F^D} = -\gamma\omega^2(r_{ij})(\boldsymbol{e}_{ij}\cdot v_{ij})\boldsymbol{e}_{ij}$ Dissipative Term $\boldsymbol{F^R} = \sigma\omega(r_{ij})\zeta_{ij}\boldsymbol{e}_{ij}$ Random Term Where $\boldsymbol{r_{ij}} = \boldsymbol{r_i} - \boldsymbol{r_j}$, $r_{ij} = \lvert\boldsymbol{r_{ij}}\rvert$, $\boldsymbol{e_{ij}} = \frac{\boldsymbol{r_{ij}}}{\lvert\boldsymbol{r_{ij}}\rvert}$ $\omega(r_{ij}) = 1 - \frac{r_{ij}}{r_c}$ *for* $r_{ij} \le r_c$; *o.w.* $\omega(r_{ij}) = 0$ The ξ_{ij} are symmetric random variables with zero mean and unit variance, uncorrelated for different pairs of particles and different times.
Properties Considerations	Parameterize the undetermined parameters to match physical properties: ▪ Platelet Cell Size ▪ Membrane Young's Modulus ▪ Membrane Shear Modulus ▪ Stretching Response ▪ Cell Plasma Compressibility ▪ Cell Plasma Viscosity ▪ Cell Plasma Pressure ▪ Cell Plasma Density Besides, we also need to consider computational feasibility and the ability of platelet model to become activated.	Parameterize the undetermined parameters and modify boundary conditions to match the physical properties: ▪ Viscosity ▪ Compressibility ▪ Reynolds Number ▪ Density ▪ Viscous Boundary Layers
Spatial Interfacing	Hybrid force filed containing the dissipative and random terms from DPD and Lenard-Jones potential from MD. It's exploited to mimic friction between platelet membranes and surrounding blood flows. $\boldsymbol{F_{ij}} = \boldsymbol{F_{ij}^{LJ}} + \boldsymbol{F_{ij}^D} + \boldsymbol{F_{ij}^R}$ Parameterize the undetermined parameters to match: ▪ platelets flipping trajectory with analytical solution (Jeffery's orbit) in Couette flow ▪ Rotation angle: $\phi = \phi(\dot{\gamma}t) = \text{atan}\left(\frac{1}{r_e}\tan\left[-\dot{\gamma}t\frac{r_e}{r_e^2+1} + \tan^{-1}(r_e\tan\phi_0)\right]\right)$	 Membrane velocity distribution of platelet during it flips in Couette flow

Computational Complexities

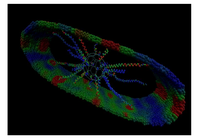


Complexity I: Disparate Temporal and Spatial Scales



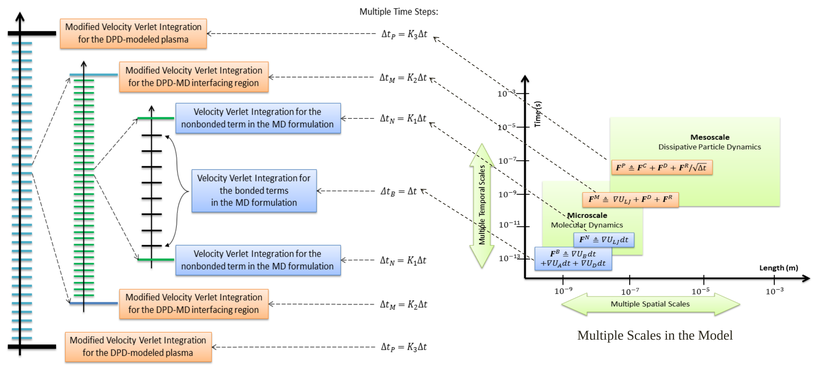
Complexity II: Complicated Model and Force Fields

Categories	Single Platelet	Multiple Platelets
In Vacuum	~0.14 million particles	Complex interactions among platelets
In Blood Plasma	~0.6 million particles	~2.7 million particles for 4 platelets flipping in blood plasm ~10.9 million particles for 16 platelets flipping in blood plasma > 50 million particles for 100 platelets in blood plasma
In Blood Vessels	Many types of blood cells and complex interactions among those cells	
With Shear Stresses & Thermo Conditions	Much more complex inputs and outputs control	



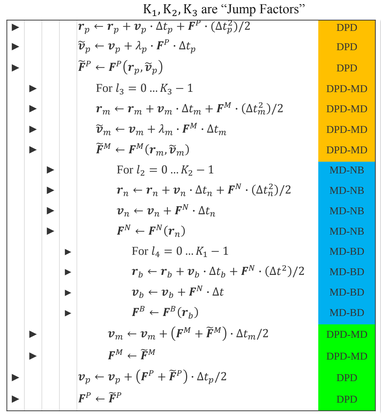
Complexity III: Large Demand of On-the-fly Analysis

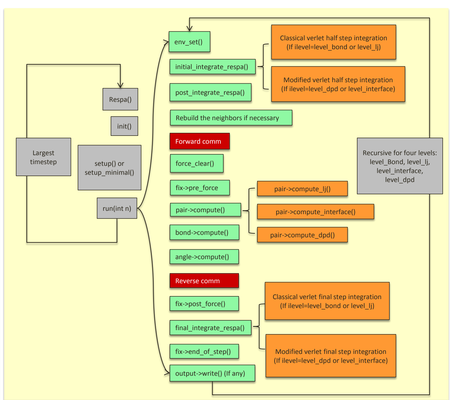
Speedup Strategy I-Multiscale Multiple Time Stepping Algorithm



Multiple Scales in the Model

K_1, K_2, K_3 are "Jump Factors"





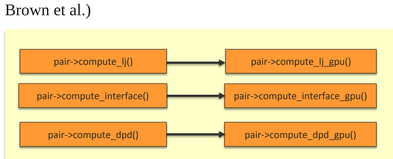
Speedup Strategy II-GPGPU Acceleration

LAMMPS (S. Plimpton et al.) and LAMMPS GPU Package (M. Brown et al.)

- Particle data is exchanged between host and device every step.
- Force evaluations and neighbor list build can be accelerated.
- Three precision options: single, mix, and double

Tailored Modifications:

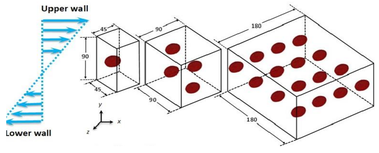
- No-slip boundary condition for DPD flows
- Above 4-level multiscale MTS algorithm
- DPD prediction and correction time integration



Performance Results on Supercomputers

➢ Varying Problem Sizes:

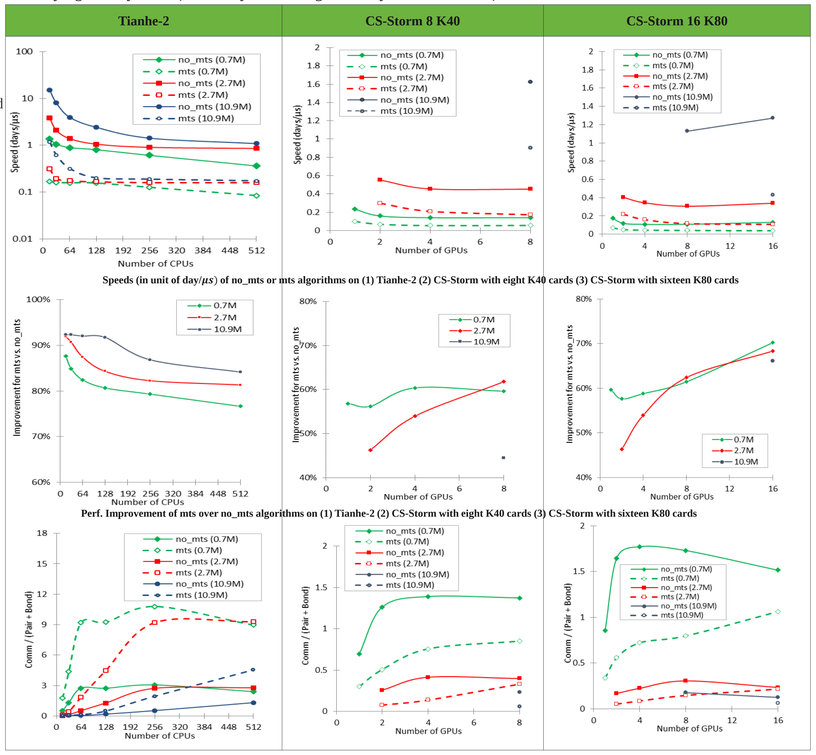
Experiments	# of Platelets	# of Particles	Dimensions
Exp-S	1	680,718	$45\times90\times45$
Exp-M	4	2,722,872	$90\times90\times90$
Exp-L	16	10,891,488	$180\times90\times180$



➢ Varying MTS Jump Factors:

Case	Time steps for each scale				Configurations			
	CGMD-BD ($\Delta t_1\times10^{-6}$)	CGMD-NB ($\Delta t_2\times10^{-6}$)	DPD-CGMD ($\Delta t_3\times10^{-6}$)	DPD ($\Delta t_4\times10^{-6}$)	$\Delta t\times10^{-6}$	K_1	K_2	K_3
CaseA	2.5	2.5	25.0	500.0	500.0	1	10	20
CaseB	5.0	5.0	50.0	1000.0	1000.0	1	10	20
CaseC	5.0	5.0	50.0	500.0	500.0	1	10	10
CaseD	10.0	10.0	100.0	500.0	500.0	1	10	5
CaseE	10.0	10.0	100.0	1000.0	1000.0	1	10	10
STS	1.0	1.0	1.0	1.0	1.0	1	1	1

➢ Varying Test Systems (CPU-only versus High-Density GPGPU Server):



Speeds (in unit of day/μs) of no_mts or mts algorithms on (1) Tianhe-2 (2) CS-Storm with eight K40 cards (3) CS-Storm with sixteen K80 cards

Perf. Improvement of mts over no_mts algorithms on (1) Tianhe-2 (2) CS-Storm with eight K40 cards (3) CS-Storm with sixteen K80 cards

Ratio of communication over computation(1) Tianhe-2 (2) CS-Storm with eight K40 cards (3) CS-Storm with sixteen K80 cards

Summary and Future Work

- With combined algorithmic and hardware accelerations, we can efficiently simulate 1-*ms* the millisecond-scale hematology at resolutions of nanoscale platelets and mesoscale bio-flows using millions of particles.
- The rule of thumb is to consider the balance of speed and accuracy for an optimal MTS scheme and the balance of computation and communication for an optimal load-balancing scheme between accelerators and CPUs.
- Future work involves with the efforts to reduce communication overheads and simulate more complicated multiscale phenomena.

References and Related Published Work

- **N. Zhang**, P. Zhang, L. Zhang, X. Zhu, L. Huang, and Y. Deng, "Performance Examinations of Multiple Time-Stepping Algorithms on Stampede Supercomputer", XSEDE15 Technical Paper Program, St. Louis, MO, 07/2015.
- P. Zhang, **N. Zhang**, Y. Deng, and D. Bluestein, "A Multiple Time Stepping Algorithm for Efficient Multiscale Modeling of Platelets Flowing in Blood Plasma", Journal of Computational Physics, vol. 284, pp. 668-686, 01/2015.
- P. Zhang, C. Gao, **N. Zhang**, M. J. Slepian, Y. Deng, and D. Bluestein, "Multiscale Particle-Based Modeling of Flowing Platelets in Blood Plasma Using Dissipative Particle Dynamics and Coarse Grained Molecular Dynamics", Cellular and Molecular Bioengineering, vol. 7 pp. 552-574, 12/2014.
- **N. Zhang**, P. Zhang, W. Kang, D. Bluestein, and Y. Deng, "Parameterizing the Morse potential for coarse-grained modeling of blood plasma", Journal of Computational Physics, vol. 257, pp. 726-736, 01/2014.

Acknowledgements

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