The Dissipative Particle Dynamics Simulation of Macromolecular Suspension in Micro-channels

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Abstract

This paper investigated the transport and conformation of macromolecules in micro-channels using the dissipative particle dynamics (DPD) and finite extensible non-linear elastic (FENE) bead spring chains model. The dynamic behavior of macromolecules with different number of beads and different chain length in three kinds of micro-channels, straight quadrate contraction sloping contraction micro-channel are comparatively analyzed. It is found that the macromolecules tend to drag the simple DPD particles, reducing their velocity, and leading to density fluctuations. The dragging effect is more important as the number of macromolecules or the length of the macromolecular chain increases.

1 Introduction

Understanding the dynamic behavior of macromolecules, such as DNA, is very important for fundamental research and practical applications in bio, chemical and medical engineering, especially in the designing micro-devices. Recently, micro-devices enable processing, analyzing, and delivering biochemical materials in a wide range of biomedical and biological applications[1, 2].

2 Methodology of the dissipative particle dynamics

In Dissipative particle dynamics system, a particle is represented a cluster of molecules or small regions of fluid material. The forces between particles are assumed to be pair-wise additive. The motion of DPD particles is governed by Newton's equations of motion. For simple DPD particle i , we have the governing equations

$$\frac{\mathrm{d}r_i}{\mathrm{d}t} = v_i, \quad \frac{\mathrm{d}v_i}{\mathrm{d}t} = \sum_{j \neq i}^{N} f_{ij_i}^{\mathrm{ext}} \tag{1}$$

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Where r_i and v_i denote the position and velocity of particle i. The mass of the all DPD particle has been taken to be the same and unity; and f_{ij} denote the total force between particles i and j. f_i^{ext} is the external force, such as the gravity. The inter-particle force f_{ij} consists of three parts, namely: conservative force F_{ij}^C, dissipative force F_{ij}^D and random force F_{ij}^R.

$$\mathbf{F}_{ij}^C = \begin{cases} a_{ij}(1 - r_{ij}/r_c)\widehat{\mathbf{r}}_{ij} & r_{ij} < r_C \\ 0 & r_{ij} \geq r_C \end{cases} \tag{2}$$

3 Parameters

To construct a working DPD, we need select values of some necessary parameters. In this section, we will talk abut how to select model parameters' values. Table 1 listed the some model parameters that we need determined before we could sufficient to construct a working DPD system. For a simple single component DPD system, we just set all particles' mass to be unity, and cutoff radius is also set to be unity.

Table 1: Model parameters

Model parameters	Symbol	Value
Mass of DPD particle	m	unity
Cutoff radius of DPD particle	r_C	unity
Simulation time step	Δt	
Friction coefficient	γ	$\sigma = 2\gamma k_b T$

4 Channel flow of FENE chain suspension

We use DPD particles and FENE chains to model the suspension of macromolecules in three kinds of micro-channels. Quadrate contraction micro-channel and sloping contraction micro-channel are comparatively analyzed. The conformation evolution of macromolecules passing through quadrate contraction micro-channel and sloping contraction micro-channel at $t = 4000$ are show in figure 1 and figure 2 respectively.

5 Conclusion

Our numerical results show that macromolecules are mainly concentrated in the middle channel. Macromolecules tend to drag simple fluid particles, reducing their velocity, and leading to density and velocity fluctuations. The dragging effect is more important as the number of macromolecules or the length of the macromolecular chain increases.

The conclusions of this paper is very important for fundamental research and practical applications in bio, chemical and medical engineering. By control the flow of macromolecular suspensions, which carried drugs and DNA molecules, we can efficiently and precisely deliver a small amount of drug or DNA into local tissue, skin regions, and even cells.
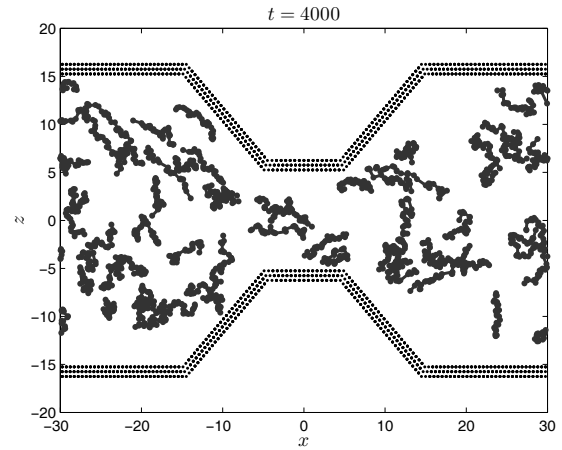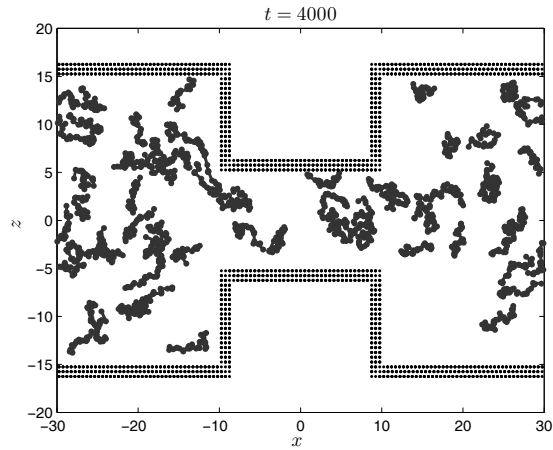
Figure 1: Quadrate contraction micro-channel   Figure 2: Sloping contraction micro-channel

Acknowledgements

I would like to thank LIU Mou-Bin.

References

[1] K. Chun, G. Hashiguchi, H. Toshiyoshi, and H. Fujita, "Fabrication of array of hollow microcapillaries used for injection of genetic materials into animal/plant cells," Jpn. J. Appl. Phys., Part 2 38, L279(1999).

[2] Fan X, Phan-Thien N, Yong N T, Wu X, Xu D. "Microchannel flow of a macromolecular suspension". Phys Fluids, 2003, 15(1): 11-21

Appendices

A   Latex Code

```latex
% copyright by Zhou Lvwen. zhou.lv.wen@gmail.com
\documentclass[12pt,a4paper]{article}   % 文档说明[字号, 纸张类型]{文档类型}
%%%%%%%%%%%%%%%%%%%%%%%%%%%%% 导 言 区 %%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%
\usepackage{amsmath,fancyhdr,graphicx,appendix,lastpage,extramarks,array}
\usepackage{listings}
\usepackage[usenames,dvipsnames]{color}

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\usepackage[left=0.8in,right=0.8in,top=1.2in,bottom=1in]{geometry}
%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%
\usepackage{xeCJK}
%\usepackage{fontspec}
\setCJKmainfont[BoldFont=simhei.ttf]{simsun.ttf}
%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%
\setlength{\parskip}{2pt}                % 定义段落间距

% 定义页眉: 左页眉{中文姓名, 学号}, 便为老师打分, 所以这里用中文姓名
%           中页眉{英语学术论文写作}为课程名称
%           右页眉{第X页,共X页}
\pagestyle{fancy} \lhead{周吕文, 201128000718065}
\chead{英语学术论文写作 (排版作业)}
\rhead{第\ \thepage\ 页,{~} 共\ \protect\pageref{LastPage} 页}

% 注: 各位同学改改页边距, 段落间距及页眉. 别搞得大家版式一样
%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%
% Latex code 环境
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\definecolor{hellgelb}{rgb}{0.96,0.96,0.96}
\usepackage{showexpl}
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52        captionpos=t,%
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54        xrightmargin=\fboxsep%
55 }
56
57 %%%%%%%%%%%%%%%%%%%%%%%%%% 正    文 %%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%
58 \begin{document}
59 \title{The Dissipative Particle Dynamics Simulation of Macromolecular
     Suspension in Micro-channels}    % 论文标题
60
61 % 作者 \\单位
62 \author{ZHOU Lv-Wen$^1$\footnote{zhou.lv.wen@gmail.com}, LIU Mou-Bin$^2$\\
63        \textit{$^{1,2}$Institute of Mechanics, CAS, Beijing 100190, China}}
64 \date{July 1, 2012}                      % 写作日期，省略则为当前计算机日期
65 \maketitle
66 \thispagestyle{fancy}                    % 首页添加页眉
67
68 % .................................................................
69 \begin{abstract}                         % 摘要
70 This paper investigated the transport and conformation of macromolecules in
     micro-channels using the dissipative particle dynamics (DPD) and finite
     extensible non-linear elastic (FENE) bead spring chains model. The dynamic
     behavior of macromolecules with different number of beads and different
     chain length in three kinds of micro-channels, straight quadrate contraction
      sloping contraction micro-channel are comparatively analyzed. It is found
     that the macromolecules tend to drag the simple DPD particles, reducing
     their velocity, and leading to density fluctuations. The dragging effect is
     more important as the number of macromolecules or the length of the
     macromolecular chain increases.
71 \end{abstract}
72 % .................................................................
73
74 \section{Introduction}                   % 引言
75  Understanding the dynamic behavior of macromolecules, such as DNA, is very
      important for fundamental research and practical applications in bio,
     chemical and medical engineering, especially in the designing micro-devices
     . Recently, micro-devices enable processing, analyzing, and delivering
     biochemical materials in a wide range of biomedical and biological
     applications\cite{KChun,FanX}.
76
77 %.................................................................
78 \section{Methodology of the dissipative particle dynamics}    % 正文
79 In Dissipative particle dynamics system, a particle is represented a cluster
     of molecules or small regions of fluid material. The forces between
     particles are assumed to be pair-wise additive. The motion of DPD particles
     is governed by Newton's equations of motion. For simple DPD particle $i$ ,
     we have the governing equations
80 \begin{equation}
81 \frac{\mathrm{d}r_i}{\mathrm{d}t} = v_i,\,\,\,\,\,
82 \frac{\mathrm{d}v_i}{\mathrm{d}t} = \sum_{j\neq i}^N f_{ij_{i}}^{\mathrm{ext}}
83 \end{equation}
84 Where $r_i$ and $v_i$ denote the position and velocity of particle $i$. The
     mass of the all DPD particle has been taken to be the same and unity; and $f
     _{ij}$ denote the total force between particles $i$ and $j$. $f_i^{\mathrm{
     ext}}$ is the external force, such as the gravity. The inter-particle force
     $f_{ij}$ consists of three parts, namely: conservative force $F_{ij}^C$,
```

```latex
    dissipative force $F_{ij}^D$ and random force $F_{ij}^R$.

\begin{equation}
\mathbf{F}_{ij}^{C}=\begin{cases}
a_{ij}(1-r_{ij}/r_{c})\mathbf{\hat{r}}_{ij} & r_{ij}<r_{C}\\
0 & r_{ij}\geq r_{C}
\end{cases}
\end{equation}

\section{Parameters}

To construct a working DPD, we need select values of some necessary parameters
   . In this section, we will talk abut how to select model parameters' values.
    Table \ref{parameters} listed the some model parameters that we need
   determined before we could sufficient to construct a working DPD system. For
    a simple single component DPD system, we just set all particles' mass to be
    unity, and cutoff radius is also set to be unity.
\begin{table}[!htb]
\centering
\caption{\label{parameters}Model parameters}
\begin{tabular}{|l|l|l|}
\textbf{Model parameters} & \multicolumn{1}{c|}{\textbf{Symbol}} & \
  multicolumn{1}{c|}{\textbf{Value}} \\
\hline
Mass of DPD particle & \multicolumn{1}{c|}{$m$} & \multicolumn{1}{c|}{unity}
   \\
\hline
Cutoff radius of DPD particle & \multicolumn{1}{c|}{$r_C$} & \multicolumn{1}{c
   |}{unity} \\
\hline
Simulation time step & \multicolumn{1}{c|}{$\Delta t$} & \multicolumn{1}{c|}{}
    \\
\hline
Friction coefficient & \multicolumn{1}{c|}{$\gamma$} & \multicolumn{1}{c|}{$\
   sigma=2\gamma k_bT$} \\
\hline
\end{tabular}
\end{table}

\section{Channel flow of FENE chain suspension}

We use DPD particles and FENE chains to model the suspension of macromolecules
    in three kinds of micro-channels. Quadrate contraction micro-channel and
   sloping contraction micro-channel are comparatively analyzed. The
   conformation evolution of macromolecules passing through quadrate
   contraction micro-channel and sloping contraction micro-channel at $t =
   4000$ are show in figure \ref{chainT} and figure \ref{chainY} respectively.

\begin{figure}[!htb]
\centering
\begin{minipage}[c]{0.5\textwidth}
\centering
\includegraphics[width=0.85\textwidth]{./figures/chainT4000s.pdf}
\caption{\label{chainT} Quadrate contraction micro-channel}
\end{minipage}%
\begin{minipage}[c]{0.5\textwidth}
\centering
```

```latex
126 \includegraphics[width=0.85\textwidth]{./figures/chainY4000s.pdf}
127 \caption{\label{chainY} Sloping contraction micro-channel}
128 \end{minipage}
129 \end{figure}
130
131 %............................................................
132
133
134 \section{Conclusion}                        % 结论
135 Our numerical results show that macromolecules are mainly concentrated in the
     middle channel. Macromolecules tend to drag simple fluid particles, reducing
      their velocity, and leading to density and velocity fluctuations. The
     dragging effect is more important as the number of macromolecules or the
     length of the macromolecular chain increases.
136
137 The conclusions of this paper is very important for fundamental research and
     practical applications in bio, chemical and medical engineering. By control
     the flow of macromolecular suspensions, which carried drugs and DNA
     molecules, we can efficiently and precisely deliver a small amount of drug
     or DNA into local tissue, skin regions, and even cells.
138
139 %............................................................
140 \section*{Acknowledgements}                 % 致谢
141 I would like to thank LIU Mou-Bin.
142
143 %............................................................
144 \begin{thebibliography}{99}                  % 参考文献
145 \bibitem{KChun} K. Chun, G. Hashiguchi, H. Toshiyoshi, and H. Fujita, ``
     Fabrication of array of hollow microcapillaries used for injection of
     genetic materials into animal/plant cells,'' Jpn. J. Appl. Phys., Part 2 38,
     L279(1999).
146 \bibitem{FanX} Fan X, Phan-Thien N, Yong N T, Wu X, Xu D. ``Microchannel flow
     of a macromolecular suspension''. Phys Fluids, 2003, 15(1): 11-21
147 \end{thebibliography}
148
149
150 \newpage
151 %............................................................
152 \appendix
153 \appendixpage                               % 附录
154 \section{Latex Code}                        % 附录 A
155 \lstinputlisting{main.tex}
156 \end{document}
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