Macroscopically measurable force induced by temperature discontinuities at solid-gas interfaces

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We consider a freely movable solid that separates a long tube into two regions, each of which is filled with a dilute gas. The gases in each region are initially prepared at the same pressure but different temperatures. Under the assumption that the pressure and temperatures of gas particles before colliding with the solid are kept constant over time, we show that temperature gaps appearing on the solid surface generate a force. We provide a quantitative estimation of the force, which turns out to be large enough to be observed by a macroscopic measurement.

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I. INTRODUCTION

Macroscopic nonequilibrium phenomena are described by evolution equations for slow modes associated with conservation laws and symmetry breaking [1]. The equations for standard liquids and gases are well established as the hydrodynamic equations of mass, momentum, and energy density fields, and heat conduction and sound propagation in solids are also well established [2]. The validity of the description was carefully investigated in small-scale experiments [3], which suggested that the behavior near a solid wall shows deviations from calculation results based on the standard hydrodynamic equations. Furthermore, a stimulating prediction that a liquid droplet is nucleated in a sheared solid may be another example that is not described by the established continuum equations [4].

In these examples, nontrivial phenomena occur at the interface between a solid and a fluid. Indeed, the description of behavior near the interface has not been established, because its characteristic length scale is too small for macroscopic phenomenological descriptions. Although imposing appropriate boundary conditions at the interface for macroscopic equations often gives a good description, there are cases where the assumptions of the boundary conditions should be seriously considered. In this paper, we study a phenomenon associated with temperature gaps at the interfaces between a solid and gases.

In order to demonstrate our findings clearly, we employ the special-purpose systems shown in Fig. 1. A solid (say, silicon), which consists of many atoms, is placed in a long tube of cross-sectional area $S$. Dilute gases (say, helium) at the same pressure $p$ but different temperatures $T_L$ and $T_R$ are contained in the left and right regions, respectively, at an initial time. The gases are well approximated by ideal gases and cannot mix with each other because the solid acts as a separating wall. It is assumed that the pressure and temperatures of gas particles before colliding with the solid are kept constant over time. In this setup, despite the equal pressure, momentum flows from one gas to the other owing to the energy transfer from the hot side to the cold side. Recently, a phenomenological mechanism for the emergence of a force from such cross-coupling has been proposed in Refs. [5, 6]. In this paper, we provide a quantitative estimation of the force acting on the solid on the basis of a microscopic description of the system under some assumptions. An important finding is that the force is determined by the temperature gaps at the interface of the solid and gases.

More precisely, we denote by $T'_L$ and $T'_R$ the kinetic temperatures of the solid particles at the left and right ends, respectively, which are different from $T_L$ and $T_R$ in general. Such a temperature discontinuity at an interface has been measured in experiments [7–9]. For the mass of gas particles $m_G$ and the mass of solid particles $m$, we define $\epsilon \equiv \sqrt{m_G/m}$, which is assumed to be small.

We then show that the temperature gaps $T'_L - T_L$ and $T'_R - T_R$ generate the force $F_{\text{gap}}$ given by

$$F_{\text{gap}} = \epsilon^2 pS \left( \frac{T'_L - T_L}{T_L} + \frac{T'_R - T_R}{T_R} \right).$$

By assuming Fourier’s law in the solid, we can estimate the temperature gaps in terms of the thermal conductivity of the solid. Surprisingly, the result shows that $F_{\text{gap}}$ takes a macroscopically measurable value. It should be noted that steady-state motion of the solid is observed.
because the force $F_{\text{fap}}$ may be balanced with a friction force induced by collision with gas particles.

In the argument below, we describe the microscopic model that we employ. We then derive the aforementioned result. Finally, we discuss the possibility of experimental realization of the phenomenon in laboratories. Throughout the paper, $\beta$ represents the inverse temperature and $k_B$ the Boltzmann constant. The subscripts or superscripts $L$ and $R$ represent quantities on the left and right sides, respectively.

II. MODEL

We provide a three-dimensional mechanical description of the solid in Fig. 2. We take the $x$ axis along the axial direction of the tube. We assume that the solid consists of $N \times M$ particles of mass $m$, where $N$ and $M$ are the number of particles along the $x$ direction and in a plane perpendicular to the $x$ axis, respectively. A collection of the positions and momenta of $N \times M$ solid particles, which we distinguish by subscripts $i$ and $j$ ($1 \leq i \leq N$, $1 \leq j \leq M$), are denoted by $\Gamma = (r_{i1,1}, \ldots, r_{IN,M}; p_{11,1}, \ldots, p_{NM,M})$, which gives the microscopic state of the solid. The $x$ components of $r_{i,j}$ and $p_{i,j}$ are denoted by $x_{i,j}$ and $p_{i,j}$, respectively, and the corresponding velocity is given by $v_{i,j} = p_{i,j}/m$. The position and velocity of the center of mass of the solid in the $x$ direction are denoted by $X$ and $V$, respectively.

The Hamiltonian of the solid, $H(\Gamma)$, is given by

$$H(\Gamma) = \sum_{i,j} \left[ \frac{|p_{i,j}|^2}{2m} + U_w(r_{i,j}) \right] + \sum_{(i,j; i',j')} U_{\text{int}}(r_{i,j}, r_{i',j'}),$$

(2)

where $(i,j; i',j')$ represents the nearest- and second-nearest-neighbor pair of solid particles. $U_{\text{int}}$ is the interaction potential between two solid particles, and $U_w$ is the potential of a solid particle and the tube wall. The tube wall is assumed to be frictionless, and $U_w(r_{i,j})$ does not depend on $x_{i,j}$. The motion of solid particles except for left and right ends is described by the Hamiltonian equations.

Next, we provide an effective description of the gases. We focus on the gas on the left side; the gas on the right side can be described similarly. Employing a dilute gas that consists of particles of mass $m_G$, we may assume that the characteristic time of the dissipation process inside each gas is much longer than the time during which we observe the steady-state motion of the solid. Therefore, gas particles that have yet to collide with the solid are in equilibrium at the temperature $T_L$, the pressure $p$, and the number density $n_L = p \beta L$. We also assume that gas particles elastically and instantaneously collide with the solid only once. More precisely, the instantaneous collision means that the characteristic time of the solid-gas interaction is much shorter than the relaxation time of all solid particles. It should be noted that this situation is completely different from a case where the solid is effectively described as a wall with one degree of freedom. For convenience, we make a list of characteristic time scales in Table I. Our assumption means that $\tau_1 \gg \tau_2 > \tau_3 \gg \tau_4$. Furthermore, for simplicity, we assume that the tangent plane at the collision point is perpendicular to the $x$ axis, so that the $x$ component of the velocity of each solid particle at both ends is independent of the other components at the collisions.

For this setup, the interaction between the solid and the gas on the left side can be described by random collisions with the collision rate $\lambda_L(v_G, v_{1,j})$ per unit area for the gas particle velocity $v_G$ and the solid particle velocity $v_{1,j}$. The collision rate is explicitly written as

$$\lambda_L(v_G, v_{1,j}) = n_L(v_G - v_{1,j}) \theta(v_G - v_{1,j}) J_{\text{eq}}^{L}(v_G),$$

(3)

where $\theta$ represents the Heaviside step function and $J_{\text{eq}}(v_G) = \sqrt{3} n_L m_G / 2 \pi \exp(-\beta L m_G v_G^2 / 2)$ is the Maxwell–Boltzmann distribution.

The $t$-th collision time of a gas particle and the $j$-th solid particle at the left end is determined according to the Poisson process. Suppose that a gas particle with a velocity in the $x$ direction $v^{L,l}_{G,j}$ collides with the $j$-th solid particle at $t = t^{L,l}_{G,j}$. The equation of motion for the $j$-th solid particle in the $x$ direction is written as

$$\frac{dp_{1,j}}{dt} = -\frac{\partial H(\Gamma)}{\partial x_{1,j}} + F_{L,j},$$

(4)

with

$$F_{L,j} = \sum_l I_{v_{L,l}}(v_{1,j}) \delta \left(t - t^{L,l}_{G,j}\right),$$

(5)

where $F_{L,j}$ is the force exerted by the elastic collisions of the gas particles, $I(v_G, v) = 2m_G m(v_G - v) / (m_G + m)$ the

| TABLE I. Characteristic time scales. |
|-------------------------------------|
| Symbol | Definition |
|--------|------------|
| $\tau_1$ | relaxation time of the dissipation process inside gas |
| $\tau_2$ | relaxation time of $V$ |
| $\tau_3$ | relaxation time of all solid particles |
| $\tau_4$ | solid-gas interaction time |
impulse of the collision, and \( \tilde{v}_{1,j}(t) \equiv \lim_{t \to t^+} v_{1,j}(t) \) the velocity just before the collision when \( t = T_{L,j} \). Similarly, the collision rate of the gas on the right side is given by \( \lambda_R(v_G, v_{N,j}) = n_R(v_{N,j} - v_G) \theta(v_{N,j} - v_G) \int_R \langle J \rangle_{v_G} \), and the equation of motion for the solid particles at the right end is determined as well.

The Hamiltonian equations in combination with the Poisson processes yield a unique steady state. The expectation value in the steady state is denoted by \( \langle \cdot \rangle \). We then have \( \langle v_{i,j} \rangle = \langle V \rangle \) for any \( i \) and \( j \), and we assume that the statistical properties in the steady state are homogeneous in the vertical direction.

### III. ANALYSIS

We consider the equation of motion for the center of mass. From the law of action and reaction, the equation in the \( x \) direction is written as \( m N M \dot{d}V/dt = \sum_{j=1}^M (F_{L,j} + F_{R,j}) \). Thus, the total force acting on the solid in the \( x \) direction is generated by the elastic collisions of the gas particles, where the collision rate depends on the velocity of the solid particles. Because \( d \langle V \rangle /dt = 0 \), we have the force balance equation in the steady state as \( \sum_{j=1}^M \langle F_{L,j} \rangle + \langle F_{R,j} \rangle = 0 \). Here, we note that

\[
\langle F_{L,j} \rangle = \int dv_G \langle J \rangle_{v_G}(v_G, v_{1,j}) \langle v_{1,j} \rangle S/M. \tag{6}
\]

We then expand the total force \( \sum_{j=1}^M \langle F_{L,j} \rangle + \langle F_{R,j} \rangle \) in \( \epsilon \equiv \sqrt{m_G/m} \). Defining \( \gamma_L \equiv \epsilon n_L \sqrt{8mk_BT_L/\pi} \) and \( \gamma_R \equiv \epsilon n_R \sqrt{8mk_BT_R/\pi} \), we obtain

\[
\sum_{j=1}^M \left[ -\gamma_L \langle v_{1,j} \rangle - \gamma_R \langle v_{N,j} \rangle + \epsilon^2 \beta_L m \langle v_{1,j}^2 \rangle - \epsilon^2 \beta_R m \langle v_{N,j}^2 \rangle \right] S/M + O(\epsilon^3) = 0. \tag{7}
\]

See Appendix A for the derivation. The first and second terms in \( \langle J \rangle \) are interpreted as the friction force that originates from the change in the collision rate due to the motion of the solid particles. The terms proportional to \( \epsilon^2 \) in \( \langle J \rangle \) are expressed in the form \( \langle J \rangle \), where \( T_{L,j} \equiv m(\langle v_{1,j}^2 \rangle - \langle v_{1,j} \rangle^2)/k_B \) and \( T_{R,j} \equiv m(\langle v_{N,j}^2 \rangle - \langle v_{N,j} \rangle^2)/k_B \) are different from \( T_L \) and \( T_R \), respectively. By using \( F_{\text{gap}} \) given in \( \langle 7 \rangle \), we rewrite \( \langle J \rangle \) as

\[
- (\gamma_L + \gamma_R) S \langle V \rangle + F_{\text{gap}} + O(\epsilon^3) = 0. \tag{8}
\]

It should be noted that \( \langle v_{i,j} \rangle = \langle V \rangle = O(\epsilon) \).

We now derive the temperature gaps for the model we consider. First, we shall find a relation connecting the temperature gap with the heat flux. Let \( J_{L,j} \) be the heat flux transferred from the gas to the \( j \)-th solid particle at the left end, and \( \dot{K}_{L,j} \) be the increasing rate of the kinetic energy per unit area of the \( j \)-th solid particle at the left end. The energy conservation law leads to \( J_{L,j} = \dot{K}_{L,j} - p v_{1,j} \). Similarly,

\[
J_{R,j} = \dot{K}_{R,j} + pv_{N,j}. \tag{10}
\]

We denote by \( \Delta K(v_G, v) \) the change in the kinetic energy of a solid particle for the collision of a solid particle of velocity \( v \) with a gas particle of velocity \( v_G \), which is given by \( \Delta K(v_G, v) = 2m_G m/(m_G v_G + m v)/(m_G + m)^2 \). We then calculate \( \langle J_{L,j} \rangle \) as \( \int dv_G \langle J \rangle_{v_G}(v_G, v_{1,j}) \langle v_{1,j} \rangle - p \langle v_{1,j} \rangle \). Expanding this expression in \( \epsilon \), we obtain

\[
\langle J_{L,j} \rangle = \frac{\gamma_R}{m} k_B (T_L - T_{L,j}) + O(\epsilon^2), \tag{9}
\]

\[
\langle J_{R,j} \rangle = \frac{\gamma_L}{m} k_B (T_R - T_{R,j}) + O(\epsilon^2). \tag{10}
\]

See Appendix A for the derivation. These equations mean that the heat flux is related to the temperature gap \( [11, 12] \). The average heat flux through the solid from the left to the right is written as \( \langle J \rangle \equiv \langle J_{L,j} \rangle = -\langle J_{R,j} \rangle \) for any \( j \).

Second, we consider the heat flux. In general, the heat flux depends on the interaction potential between solid particles, and it is difficult to calculate it from a microscopic description. Nevertheless, by selecting a proper short-range interaction between solid particles in our model, we may phenomenologically assume Fourier's law in the form

\[
\langle J \rangle = \kappa(T_L - T_R)/L, \tag{11}
\]

where \( \kappa \) and \( L \) represent the thermal conductivity and the axial length of the solid, respectively, and the temperature dependence of \( \kappa \) is ignored. From \( \langle 9 \rangle \), \( \langle 10 \rangle \), and \( \langle 11 \rangle \), we obtain

\[
\langle J \rangle = \frac{\kappa(T_L - T_R)/L}{1 + \kappa m (1/\gamma_L + 1/\gamma_R)/(k_B L)} + O(\epsilon^2), \tag{12}
\]

and

\[
T_{L,j} - T_L = \frac{\gamma_R/(\gamma_R + \gamma_R)}{1 + k_B L \gamma_R/(k_B m (\gamma_L + \gamma_R))} (T_R - T_{L,j}) + O(\epsilon). \tag{13}
\]

By substituting this result into \( \langle 11 \rangle \), we obtain an expression for \( F_{\text{gap}} \) in terms of the experimental parameters. Furthermore, \( \langle 11 \rangle \), \( \langle 9 \rangle \), \( \langle 10 \rangle \), and \( \langle 12 \rangle \) lead to the simple relation

\[
\langle V \rangle = -\frac{\pi \langle J \rangle}{8p} + O(\epsilon^2), \tag{14}
\]

which connects the moving velocity with the heat flux passing inside the solid. See Refs. [3, 4] for an intuitive explanation of the result.

In order to directly demonstrate the validity of our theory, we performed numerical experiments by solving the Hamiltonian equations in combination with the Poisson process. Here, for simplicity, we consider the case that the system is defined in two dimensions. Concretely, we use the potentials \( U_{\text{int}}(r, r') = k(|r - r'| - \sqrt{d})^2/2 \) for the \( d \)-th nearest neighbor pair of solid particles \( (d = 1, 2) \) and \( U_{\text{gap}}(r) = 1/|r - r'_{\text{gap}}(r)|^6 \), where \( k \) is the spring constant, \( \sqrt{d} \) the natural length, and \( r'_{\text{gap}}(r) \) the nearest position of
the tube wall from $r$. All the quantities are converted into dimensionless forms by setting $k = a = m = 1$. We then set the parameter values as $k_B T_L = 0.07$, $k_B T_R = 0.1$, and $\epsilon = \sqrt{1/10}$. Because the equations are nonlinear, we observed the temperature gradient inside the solid and the temperature gaps at the surface of the solid. We then measured $\langle V \rangle$ and $\langle J \rangle$ for several values of $p$ and $N = M$, and plotted $\langle -\langle J \rangle/p , \langle V \rangle \rangle$ in Fig. 3. We find that the obtained data is consistent with the nontrivial relation (14).

IV. EXPERIMENTS

Let us discuss the experimental feasibility of the phenomenon under consideration. As one example of laboratory experiments, we consider silicon and helium of atomic weight 28 and 4, respectively, where $\epsilon = \sqrt{1/7}$. The thermal conductivity of silicon at room temperature is $\kappa \simeq 149 \text{ J/(m} \cdot \text{s} \cdot \text{K)}$, and the density of silicon is $\rho \simeq 2.33 \text{ g/cm}^3$. We set $T_L = 293 \text{ K}$, $T_R = 303 \text{ K}$, $p = 1 \text{ atm}$, $S = 7 \text{ cm}^2$, and $L = 1 \text{ cm}$. By using (1), (12), (13), and (14), we obtain the velocity of the solid $\langle V \rangle \simeq 3.9 \times 10^{-8} \text{ cm/s}$, the temperature gap $T_L' - T_L \simeq 1.6 \text{ K}$, and the temperature-gap-induced force $F_{\text{gap}} \simeq 1.1 \times 10^{-4} \text{ N}$. These estimated values are large enough to be measured in careful experiments. We next consider several possible difficulties that may arise in experiments.

First, there is the friction between the solid and the tube. Because the coefficient of static friction of a lubricant is at most 0.5 [13], the static friction force is about $0.5 \times 9.8 \text{ m/s}^2 \times \rho S L \simeq 8.0 \times 10^{-2} \text{ N}$, which is smaller than $F_{\text{gap}} \simeq 1.1 \times 10^{-4} \text{ N}$. Thus, the effect of the friction can be mitigated by the use of a lubricant.

Second, one may worry that the relaxation time of the motion of the solid is longer than the observation time. However, since the relaxation time is estimated as $m N M / (\gamma_L + \gamma_R) S = \rho L / (\gamma_L + \gamma_R) \simeq 5.7 \times 10^{-2} \text{ s}$, the velocity of the solid is rapidly relaxed to the steady-state value.

Lastly, the most difficult experimental setup may be the control of the temperatures of the gases. One method is to connect the tube with baths of dilute gas, where the length of the tube is chosen to be shorter than the mean-free path of the dilute gas as shown in Fig. 4. The mean-free path of the helium atoms at 1 atm and 300 K is about 200 nm and is inversely proportional to the pressure. Therefore, when the length of the tube is 20 cm, we have to set the pressure at $1 \times 10^{-6} \text{ atm}$, and $F_{\text{gap}}$ becomes very small. On the contrary, when the mean-free path is much shorter than the length of the tube, a temperature gradient appears in the gases. This makes $T_R - T_L$ small, and as a result $F_{\text{gap}}$ becomes small. The simplest realization is to control the temperatures from the side wall of the tube. In this case, we immobilize the solid by attaching a spring to it (see Fig. 5). Because the value of the stall force is equal to $F_{\text{gap}}$, we can measure its value, whereas we cannot observe the steady-state motion.

Setting aside the quantitative aspects, we may observe phenomena to which $F_{\text{gap}}$ makes a dominating contribution. One example is a Brownian particle under a temperature gradient [14,17]. If the thermal conductivity of the particle is much larger than that of a solution, heat flux passes inside the particle. Then, because the temperature gap appears on the solid surface, the force $F_{\text{gap}}$ is generated. It should be noted, however, that other types of forces appear on the particle under a temperature gradient [17]. It is a stimulating challenge to separate $F_{\text{gap}}$.
from the total force.

V. CONCLUDING REMARKS

In this paper, we have predicted that the temperature gap at the interface between a solid and a gas yields the force $F_{\text{gap}}$. Because this force is not described by standard continuum theory such as hydrodynamics and elasticity theory, further experimental and theoretical studies are necessary so as to obtain a systematic understanding of the nature of the force. Before ending the paper, we provide a few remarks.

With regard to our setup, similar models were studied in the context of the so-called adiabatic piston problem [18-21]. Indeed, for the case $\kappa \gg L_k B \gamma_L/[m(\gamma_L + \gamma_R)]$ or $N = 1$, we obtain $\langle J \rangle = k T_{\text{L}} - T_{\text{R}}/[m(\gamma_L + \gamma_R)] + O(\epsilon^2)$ and $\langle V \rangle = \sqrt{\pi/S}(\sqrt{k T_{\text{L}}} / m - \sqrt{k T_{\text{R}}}/m) + O(\epsilon^2)$, where $T_{\text{L}} = T_{\text{R}}$. These expressions for $\langle J \rangle$ and $\langle V \rangle$ are identical to those derived in Refs. [22, 23]. Here, it should be noted that $m$ in the preceding studies was assumed to be the total mass of the solid, which gives much smaller values of $\langle J \rangle$ and $\langle V \rangle$ than ours.

In contrast to the interface case, there exists no force due to temperature differences at the atomic scale in the bulk. In order to clearly understand the difference between the two cases, we should derive $F_{\text{gap}}$ on the basis of a mechanical description of solids and gases. This fundamental question might be solved by considering hydrodynamics of a binary mixture fluid in the phase separation state. Because hydrodynamics may involve the discontinuity of the temperature profile at the interface between the two materials, the standard assumption of slowly varying thermodynamic quantities may not be valid. The derivation may be obtained as an extension of a recent work [24] in which the hydrodynamic equations for a simple fluid are derived from a Hamiltonian description of identical particles. Obviously, the experimental measurement of $F_{\text{gap}}$ is of great importance even for the theory. By clarifying the mechanism of nonstandard forces, we hope to develop the understanding of nonequilibrium systems.

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Appendix A: Derivation of (7), (9), and (10)

By considering $f_{\text{eq}}^L(v) = \epsilon \sqrt{\beta_L m/2\pi} \exp(-\epsilon^2 \beta_L m v^2/2)$ and $f_{\text{eq}}^R(v) = \epsilon \sqrt{\beta_R m/2\pi} \exp(-\epsilon^2 \beta_R m v^2/2)$, we obtain

\[
\int_V v^k f_{\text{eq}}^L(v) dv = \int_0^\infty v^k f_{\text{eq}}^L(v) dv + O(\epsilon) = \frac{\epsilon^{-k}}{2\sqrt{\pi}} \langle \frac{2}{\beta_L m} \rangle^\frac{k}{2} \Gamma \left( \frac{k+1}{2} \right) + O(\epsilon),
\]

(A1)

\[
\int_{-\infty}^V v^k f_{\text{eq}}^R(v) dv = \int_0^\infty v^k f_{\text{eq}}^R(v) dv + O(\epsilon) = \frac{(-\epsilon)^{-k}}{2\sqrt{\pi}} \langle \frac{2}{\beta_R m} \rangle^\frac{k}{2} \Gamma \left( \frac{k+1}{2} \right) + O(\epsilon),
\]

(A2)

where $k$ is a non-negative integer. By using (A1) and (A2), we obtain

\[
\int \lambda_L(v, V) I(v, V) dv = \frac{2 \epsilon^2 m n_L}{1 + \epsilon^2} \int_V (v - V)^2 f_{\text{eq}}^L(v) dv
= (1 - \epsilon^2)^2 \pi / \beta_L V + \epsilon^2 m \beta_L V^2 + O(\epsilon^2),
\]

(A3)

\[
\int dv \lambda_R(v, V) I(v, V)
= \frac{2 \epsilon^2 m n_R}{1 + \epsilon^2} \int_{-\infty}^V dv (V - v)^2 f_{\text{eq}}^R(v)
= -(1 - \epsilon^2)^2 \pi / \beta_R V - \epsilon^2 m \beta_R V^2 + O(\epsilon^2),
\]

(A4)

\[
\int \lambda_L(v, V) \Delta K(v, V) dv
= \frac{2 \epsilon^2 m n_L}{1 + \epsilon^2} \int_V (v - V)^2 (v^2 + V) f_{\text{eq}}^L(v) dv
= p V - \gamma_L V^2 + \frac{\gamma_L k_B T_L}{m} + O(\epsilon^2),
\]

(A5)

\[
\int \lambda_R(v, V) \Delta K(v, V) dv
= \frac{2 \epsilon^2 m n_R}{1 + \epsilon^2} \int_{-\infty}^V (V - v)^2 (v^2 + V) f_{\text{eq}}^R(v) dv
= -p V - \gamma_R V^2 + \frac{\gamma_R k_B T_R}{m} + O(\epsilon^2),
\]

(A6)

where we have used $p = n_L / \beta_L = n_R / \beta_R$, $\gamma_L = c n_L / 8 \pi m k_B T_L / \pi$, and $\gamma_R = c n_R / 8 \pi m k_B T_R / \pi$. Here, we assume that $\langle O(\epsilon^3) \rangle = O(\epsilon^3)$. Then, (A3) and (A4) lead
By using \( \langle v_{1,j} \rangle = \langle V \rangle = O(\epsilon) \), \( \text{(A5)} \) and \( \text{(A6)} \), we obtain

\[
\langle J_{L,j} \rangle = \left\langle \int dv_G \lambda_L(v_G, v_{1,j}) \Delta K(v_G, v_{1,j}) \right\rangle - p \langle v_{1,j} \rangle \\
= \frac{\gamma_L}{m} k_B \left( T_L - m \langle v_{1,j}^2 \rangle / k_B \right) + O(\epsilon^2)
\]

\[
\langle J_{R,j} \rangle = \left\langle \int dv_G \lambda_R(v_G, v_{N,j}) \Delta K(v_G, v_{N,j}) \right\rangle + p \langle v_{N,j} \rangle \\
= \frac{\gamma_R}{m} k_B \left( T_R - m \langle v_{N,j}^2 \rangle / k_B \right) + O(\epsilon^2).
\]

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