

Anisotropic acoustic waves in rarefied nematic liquid crystals

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Why rarefied nematic liquid crystals?



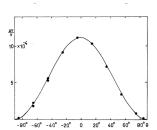


FIG. 2. Angular dependence of sound velocity. T =21°C, ν =10 MHz, and H=5 kOe. θ is the angle between the field direction and propagation direction. Solid line is $12.5 \times 10^{-4} \cos^2 \theta$.

Acoustic waves travel in NLC faster First order theory better fits in the direction parallel to the nematic director [MLS72].

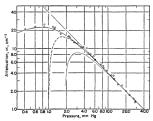


Fig. 1. Attenuation of sound at 1 Mc/sec. in helium. Circles—experi-mental results. Heavy full line—exact hydrodynamic, Light full line first approximation, hydrodynamic and Burnett. Dashed line-second approximation, hydrodynamic. Dotted line-second approximation, Burnett.

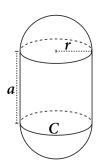
experimental data on acoustic attenuation at low pressure [Gre49].

Curtiss collision operator



In his seminal paper, Curtiss [Cur56] proposed a kinetic theory for spherocylindrical molecules as an idealisation of a polyatomic gas.

- ▶ He considered a larger configuration space made by position, velocity, the Euler angles for describing the orientation of each molecule, and the angular velocity with respect to a fixed coordinate system.
- Molecules would interact by excluded volume, which give rise to short range interactions, hence the nematic ordering.





This led Curtiss to formulate the following **Boltzmann** type equation,

$$\partial_t f + \nabla_r \cdot (\mathbf{v}f) + \nabla_\alpha \cdot (\dot{\alpha}f) = C[f, f] \tag{1}$$

where $f(\mathbf{r}, \mathbf{v}, \alpha, \omega)$ is the usual first reduced distribution function and C[f, f] is the collision operator defined as

$$C[f,f] = -\iiint (f_1^{'}f^{'} - f_1f)(\mathbf{k} \cdot \mathbf{g})S(\mathbf{k})d\mathbf{k}d\mathbf{v}_1d\alpha_1d\omega_1$$

with S(k)dk being the surface element of the excluded volume and $g = v - v_1$. Here without loss of generality the equation is stated in absence of external force and torque.

Collision invariants



It is possible to prove that the following quantities are **collision** invariants for C[f, f], i.e.

$$\iiint \psi^{(i)} d \mathbf{v}_1 d \omega_1 d \alpha_1 = 0.$$

- $\psi^{(1)} = 1$, the **number of particles** in the system;
- $\blacktriangleright \psi^{(2)} = m\mathbf{v}$, the linear momentum;
- $\psi^{(3)} = \mathbb{I}^1 \cdot \omega + \mathbf{r} \times m\mathbf{v}$, the angular momentum;
- $\psi^{(4)} = \frac{1}{2}m\mathbf{v}\cdot\mathbf{v} + \frac{1}{2}\boldsymbol{\omega}\cdot\mathbb{I}\cdot\boldsymbol{\omega}$, the kinetic energy of the system.

¹The inertia tensor for the spherocylinder we are considering.

The hydrodynamic equations - notation



We first introduce the number density, i.e.

$$n(\mathbf{r}) = \iiint f(\mathbf{r}, \mathbf{v}, \alpha, \omega) d\mathbf{v} d\alpha d\omega.$$

Then we can give a meaning to the following chevrons, i.e.

$$\langle\!\langle \cdot \rangle\!\rangle(\mathbf{r}) \coloneqq \frac{1}{n(\mathbf{r})} \iiint \cdot f(\mathbf{r}, \mathbf{v}, \alpha, \omega) d\mathbf{v} d\alpha d\omega.$$

Using this notation we can define macroscopic stream velocity and macroscopic stream angular velocity respectively as:

$$\mathbf{v}_0 \coloneqq \langle \langle \mathbf{v} \rangle \rangle, \qquad \omega_0 \coloneqq \langle \langle \omega \rangle \rangle.$$

The Hydrodynamic Equations – Curtiss Balance Laws



Testing (1) against the first two **collision invariants** and integrating, Curtiss obtained the following **balance laws**:

$$egin{aligned} \partial_t
ho +
abla_{m r} \cdot (
ho m v_0) &= 0, \ \\
ho \Big[\partial_t m v_0 + (
abla_{m r} m v_0) m v_0 \Big] +
abla_{m r} \cdot (
ho \mathbb{P}) &= 0, \end{aligned}$$

where ρ is the **density** defined as $\rho(r) = mn(r)$ and \mathbb{P} is the **pressure tensor** defined as $\mathbb{P} := \langle \langle V \otimes V \rangle \rangle$, with V being the **peculiar velocity** $V := v - v_0$.

The hydrodynamic equations – surprise balance laws



For the third collision invariant we took a different route than Curtiss, which led to the following balance law

$$\rho \Big[\partial_t \boldsymbol{\eta} + (\nabla_r \boldsymbol{\eta}) \boldsymbol{v}_0 \Big] + \nabla_r \cdot (\rho \mathbb{N}) = \boldsymbol{\xi}, \tag{2}$$

where η is the macroscopic intrinsic angular momentum defined as $\eta(\mathbf{r}) \coloneqq \langle \langle \mathbb{I} \cdot \omega \rangle \rangle$ and \mathbb{N} is the couple tensor defined as $\mathbb{N} \coloneqq \langle \langle \mathbf{V} \otimes (\mathbb{I}\omega) \rangle \rangle$. Here ξ_I is defined in tensor notation as $\langle \langle mn(\varepsilon_{Iki}v_iv_k)\mathbf{e}_I \rangle \rangle$ and we proved that $\boldsymbol{\xi}$ vanishes (as stated by Curtiss in [Cur56]) in this particular setting.

Maxwell-Boltzmann distribution



In [Cur56] Curtiss gives an expression for the Maxwell–Boltzmann distribution, i.e. the distribution $f^{(0)}$ such that $C[f^{(0)}, f^{(0)}]$ vanishes:

$$f^{(0)}(\mathbf{v}, \boldsymbol{\omega}) = n \frac{\sin(\alpha_2)Q}{\int Q \sin(\alpha_2) d\alpha} \frac{m^{\frac{3}{2}}}{2\pi\theta}^{3} (\Gamma_1 \Gamma_2 \Gamma_3)^{\frac{1}{2}} \exp\left[-m \frac{|\mathbf{V}|}{2\theta} - \frac{\mathbf{\Omega} \cdot \mathbb{I} \cdot \mathbf{\Omega}}{2\theta}\right],$$

where the **peculiar angular velocity** defined as $\Omega \coloneqq \omega - \omega_0$, Γ_i are the moments of inertia of the spherocylinder we are considering and $Q \coloneqq \exp\left[\frac{\omega_0 \cdot \mathbb{I} \cdot \omega_0}{2\theta}\right]$.

Notice in particular that we assumed ω_0 and the **kinetic** temperature $\theta = \frac{m}{2} \mathbf{V} \cdot \mathbf{V} + \frac{1}{2} \mathbf{\Omega} \cdot \mathbb{I} \cdot \mathbf{\Omega}$ are fixed.

Momentum closure around the equilibrium



Now we can use the Maxwell–Boltzmann distribution to compute an approximation of the **pressure tensor** near the equilibrium, i.e.

$$\mathbb{P}^{(0)} = \theta Id$$
.

We can define the **pressure** as $p = \rho \theta$ and rewrite,

$$\left[\partial_t \mathbf{v}_0 + (\nabla_r \mathbf{v}_0) \mathbf{v}_0
ight] = -rac{1}{
ho}
abla p,$$

which is the well known **Euler equation** that if linearised yields the wave equation. Unfortunately the same procedure results in a vanishing $\mathbb{N}^{(0)}$.

Balance laws for kinetic temperature



We need another way to formulate the **constitutive relation** for the **couple tensor**. We begin by observing that from $\psi^{(4)}$ we get the following balance law:

$$\dot{\psi} + \nabla_{r} \mathbf{v}_{0} : \mathbb{P} + \nabla_{r} \omega_{0} : \mathbb{N} - \nabla \cdot \left[\mathbb{P}^{T} \mathbf{v}_{0} + \mathbb{N}^{T} \omega_{0} \right] \geq 0$$

where $\psi = \langle\!\langle \theta \rangle\!\rangle$. We add ξ and observe that if we integrate with appropriate boundary condition the expression is the **rate of work** theorem that was the starting point of Leslie–Ericksen theory:

$$\dot{\psi} + \nabla_{r} \mathbf{v}_{0} : \mathbb{P} + \nabla_{r} \omega_{0} : \mathbb{N} - \nabla \cdot \left[\mathbb{P}^{T} \mathbf{v}_{0} + \mathbb{N}^{T} \omega_{0} \right] + \boldsymbol{\xi} \ge 0.$$
 (3)

Noll-Coleman procedure



Since we are happy with our **pressure tensor** so far we make the following **ansatz**

$$\psi = \psi(\nu, \nabla \nu)$$

where ν is the **nematic director**. Expanding the total derivative and using the Ericksen identity we get the following expression in tensor notation

$$\dot{\psi} = \varepsilon_{iqp} \Big[(\nu_q \frac{\partial \psi}{\partial (\nu_p)} + \partial_k (\nu_q) \frac{\partial \psi}{\partial (\partial_k \nu_p)}) \omega_i^0 + \nu_q \frac{\partial \psi}{\partial (\partial_k \nu_p)} \partial_k \omega_i^0 \Big] - \frac{\partial \psi}{\partial (\partial_k \nu_p)} \partial_q (\nu_p) \partial(\nu_q^0)$$

Noll-Coleman procedure



Substituting this expression into (3) and considering thermodynamic processes for which the exact divergences disappear, we get:

$$\begin{split} \left[\mathbb{P}_{ij} + \frac{\partial \psi}{\partial (\partial_{j} \nu_{p})} \partial_{i}(\nu_{p})\right] \partial_{j}(\nu_{i}) + \left[N_{ij} - \varepsilon_{iqp} \nu_{q} \frac{\partial \psi}{\partial (\partial_{j} \nu_{p})}\right] \partial_{j}(\omega_{i}^{0}) \\ \left[P_{pq} - \frac{\partial \psi}{\partial (\partial_{p} \nu_{k}) \partial_{q}(\nu_{k})}\right] \varepsilon_{iqp} \omega_{i}^{0} \geq 0. \end{split}$$

Since the above expression must hold for all thermodynamic processes for which the exact divergences disappear, we get the following **constitutive relations**:

$$\mathbb{P} = \nabla \boldsymbol{\nu}^{\mathsf{T}} \frac{\partial \psi}{\partial (\nabla \boldsymbol{\nu})} + \mathbb{P}^{(0)}, \qquad \mathsf{N}_{ij} = \varepsilon_{iqp} \nu_q \frac{\partial \psi}{\partial (\partial_i \nu_p)} = \boldsymbol{\nu} \times \frac{\partial \psi}{\partial (\nabla \boldsymbol{\nu})}.$$

Anisotropic waves



It can be shown that steady spherical solutions of (2) are of the form $\nu = \frac{\mathbf{r}}{|\mathbf{r}|}$ and that $\nabla \nu^T \nabla \nu = \mathbf{Id} - \nu \otimes \nu$. Therefore for this particular case we have the following choice of **pressure tensor**:

$$\mathbb{P} = \mathbb{P}^{(0)} + \mathbf{Id} + \mathbf{\nu} \otimes \mathbf{\nu}.$$

If we linearise the **Euler equation** with this choice of **pressure tensor** we get the wave equation:

$$\partial_t^2 \rho - \nabla \cdot \left[(\mathbf{Id} + \boldsymbol{\nu} \otimes \boldsymbol{\nu}) \nabla \rho \right] = 0.$$

Anisotropic waves



It is well known that a planar wave

$$p(\mathbf{r},t) = A\cos(\mathbf{k}\cdot\mathbf{r} - \omega t)$$

travelling in a transversely isotropic medium has speed of sound

$$c_s = (1 + \cos(\theta))^2$$

where θ is the angle between k and ν . A similar reasoning was presented in [BDT14], where a theory for anisotropic waves in **dense** liquid crystals is developed.

Conclusions



- ► First order theory arise naturally from Boltzman type equation even for rod-like molecules.
- Using a Noll-Coleman argument for the closure of the the momentum hierarchy allows us to capture the anisotropy of acoustic waves in rarefied liquid crystals.
- ▶ We hope to use the relation that arise from the closure procedure presented today to compute Frank constants from I, the inertia tensor of the spherocylinder we are considering.

References





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