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Energy mechanism for the instability of liquid jets with thermocapillarity ⊘

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ABSTRACT

Xu and Davis [J. Fluid Mech. **161**, 1–25 (1985)] examined the stability of long axisymmetric liquid jet subjected to an axial temperature gradient, finding capillary, surface-wave, and hydrodynamic modes. They showed that capillary breakup can be retarded or even suppressed for a small Prandtl number (Pr < 1) and a large Biot number ($Bi \ge 1$). In the present work, the energy mechanism is carried out for these three kinds of flow instabilities, and the mechanism of suppressing capillary breakup is clarified. When the Reynolds number (R_B) is not large, the work done by the pressure on the free surface (P_S) is the main energy source of the capillary instability. At small Pr and large Bi, the phase difference between the radial velocity and surface deformation increases with R_B , leading to the decrease in P_S , which prevents the occurrence of capillary breakup. Meanwhile, the work done by thermocapillary force becomes the main energy source, making hydrodynamic modes unstable. The perturbation flow fields are displayed, which shows that the temperature fluctuations of three modes differ from each other.

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I. INTRODUCTION

If a temperature difference exists along a gas–liquid interface, the variation of surface tension will induce a convection in the liquid. This phenomenon is called the thermocapillary effect. Since the pioneering work of Pearson,¹ there has been a much greater understanding in this field. If a liquid jet is exposed to a temperature difference from its surroundings, additional instabilities may arise, which can impact the breakup properties of the jet. This situation is found in various processes, such as inkjet printing,^{2,3} fiber preparation,⁴ droplet formation,^{5,6} injection of rocket engine, and gas turbine.^{7,8}

Different types of instabilities may occur based on the direction of temperature gradient. Marangoni instabilities arise from the radial temperature gradient, and extensive research works have been conducted in this area. Imaishi *et al.*⁹ have linearly analyzed the effect of the Prandtl number (*Pr*) on the stability of thermocapillary flow in shallow annular pools. By means of energy budget analysis, they have revealed that instabilities in low-*Pr* range are hydrodynamic in nature. Conversely, instabilities in the middle- and high-*Pr* regions are hydrodynamic instabilities. Qiao *et al.*¹⁰ have investigated temporal linear instability of coaxial jets under a radial temperature gradient, exploring the relative importance of various energy components in the most unstable mode through the application of perturbation kinetic energy balance. Li *et al.*¹¹ have analyzed the stability of thermocapillary flow for a medium Prandtl number (Pr = 6.7) in rotating annular pools with different aspect ratios, showing that all instabilities are hydrothermal. Liu *et al.*¹² have studied thermocapillary flow instabilities in annular pools with different aspect ratios using both linear stability analysis and energy analysis, reporting that the instability mechanism in cases of a small Prandtl number (Pr = 0.011) is hydrodynamic, while for a medium Prandtl number (Pr = 1.4), it is influenced by both hydrodynamic and Marangoni effects.

On the other hand, thermocapillary effect induced by the axial temperature gradient also has a significant impact on the stability of jet flow. Xu and Davis¹³ have investigated the stability of an infinitely long axisymmetric liquid column with interfacial deformation under such a gradient. They have found a new type of instability known as

the hydrothermal instability, in addition to the previously identified capillary instability¹⁴ and surface-wave instability.¹⁵ Furthermore, they have demonstrated that in simplified isothermal jets (the Biot number $Bi \rightarrow \infty$), the surface-wave instabilities may eradicate completely the tendency for capillary breakup. Similarly, the capillary breakup can be retarded or even suppressed for a small Prandtl number (Pr < 1) and a large Biot number ($Bi \ge 1$). Subsequent studies on annular liquid films¹⁶ and encapsulated liquid floating zones¹⁷ have further shown that the thermocapillary-driven flow within the liquid column significantly weakens the capillary instability, thereby enhancing the jet stability.

Wanschura *et al.*¹⁸ have conducted linear stability analysis and energy analysis to investigate thermocapillary flow in a cylindrical liquid bridge with different *Pr*. They have found that the low Prandtl number instability is an instability of the basic axial shear flow, whereas the instability at high *Pr* is attributed to the convective radial heat transport coupled with the Marangoni effect. In a separate study, Levenstam and Amberg¹⁹ have numerically simulated a half-zone model and determined that the instability at low *Pr* is purely hydrodynamic. Later, these findings were further supported by Leypoldt *et al.*²⁰ Recently, thermocapillary flows in liquid bridges between coaxial disks of unequal radii ratios have been widely examined.^{21–23} Based on the methods of linear stability analysis and energy analysis, these studies have revealed that the instability modes at small *Pr* are primarily driven by hydrodynamic effects, while the instability mechanism at larger *Pr* cannot neglect the influence of thermocapillary effect.

Most previous studies have predominantly concentrated on understanding the mechanism of thermocapillary instabilities in liquid columns of finite volume. However, there is still limited investigations on long liquid columns, where the destabilization mechanism of unstable modes and the factors inhibiting capillary breakup remain unclear. Expanding upon the study conducted by Xu and Davis,¹³ the present work explores the energy mechanism for the instability of liquid jets with thermocapillarity, presenting the perturbation flow fields and providing insight into the mechanisms that prevent the capillary breakup of jets.

The paper is organized as follows: Sec. II recalls the model of thermocapillary liquid jets, including dimensionless governing equations and boundary conditions, along with the basic state of flow and temperature profiles. Section III constructs the equation for perturbation kinetic energy. The results of energy components and relevant discussion are presented. Section IV is devoted to the analysis of perturbation fields and an explanation for the mechanism that suppresses capillary breakup. Finally, in Sec. V, we itemize the conclusions.

II. MATHEMATICAL FORMULATION

We consider an infinitely long liquid column of basic radius a with a deformable free surface $r = R(\theta, z, t)$, as illustrated in Fig. 1. The column is described using a cylindrical coordinate system (r, θ, z) ; the corresponding velocity components are (u, v, w). A temperature gradient dT/dz = -b < 0 is applied along the axial direction of the liquid column, while the surface tension σ varies linearly with the temperature,

$$\sigma = \sigma_0 - \gamma (T - T_0), \tag{1}$$

where T_0 is the temperature on the surface at z=0, and $\gamma = -d\sigma/dT > 0$ is the rate of change in surface tension with respect to the temperature.



FIG. 1. The sketch of a basic-state liquid jet with thermocapillarity.

We consider an incompressible Newtonian fluid, whose dynamic viscosity μ , density ρ , specific heat c_p , thermal conductivity rate χ , and unit thermal surface conductance h are constants. Length $L \sim a$, velocity $\mathbf{u} \sim w_* = \gamma b a / \mu$, velocity $\mathbf{u} \sim w_{**} = (\sigma_0 / \rho a)^{\frac{1}{2}}$, time $t \sim (\rho a^3 / \sigma_0)^{\frac{1}{2}}$, pressure $p \sim \sigma_0 / a$, temperature difference $T - T_0 \sim b a$, and surface tension $\sigma \sim \sigma_0$ are selected as characteristic scales. Here, the characteristic velocity w_{**} is related to the capillary instability, whereas w_* is consistent with the thermocapillary effect. This particular configuration considers both the capillary breakup and the thermocapillary effect in a long jet.

The following dimensionless numbers are generated:

$$S = \frac{\rho a \sigma_0}{\mu^2}, \quad R_{\rm B} = \frac{\rho \gamma b a^2}{\mu^2}, \quad Pr = \frac{\mu c_p}{\chi}, \quad Bi = \frac{ha}{\chi}.$$
 (2)

Here, $S^{\frac{1}{2}}$ is the Reynolds number for the capillary instability, which is associated with mean surface tension.¹³ $R_{\rm B}$ is the Reynolds number for the steady thermocapillary flow. *Pr* is the Prandtl number, and *Bi* is the surface Biot number that measures the heat transfer between the liquid and the surrounding gas.

A. Governing equations

In a cylindrical coordinate system, the mathematical model is expressed by the following dimensionless governing equations:

$$\nabla \cdot \mathbf{u} = 0, \tag{3a}$$

$$\partial_t \mathbf{u} + \mathbf{u} \cdot \nabla \mathbf{u} = -\nabla p + S^{-\frac{1}{2}} \nabla \cdot \boldsymbol{\tau}, \tag{3b}$$

$$Pr(\partial_t T + \mathbf{u} \cdot \nabla T) = S^{-\frac{1}{2}} \nabla^2 T.$$
(3c)

Here, **u**, *p*, *T*, and τ stand for the velocity, pressure, temperature, and stress tensor, respectively.

For Newtonian fluids,

$$\tau = \dot{\gamma}, \tag{4}$$

where
$$\dot{\boldsymbol{\gamma}} = \nabla \mathbf{u} + (\nabla \mathbf{u})^{\mathrm{T}}$$
.

The boundary conditions are listed as follows. At r = 0, all physical quantities must be finite values:

$$|u|, |v|, |w|, |p|, |T| < \infty.$$
(5)

On the liquid surface at
$$r = R(\theta, z, t)$$
,

$$\partial_t R + \mathbf{u}_\perp \cdot \nabla R = u,$$
 (6)

$$\mathbf{t}_j \cdot \boldsymbol{\tau} \cdot \mathbf{n} = -S^{-\frac{1}{2}} R_{\mathrm{B}} \nabla T \cdot \mathbf{t}_j, \tag{7}$$

$$-p + S^{-\frac{1}{2}}\mathbf{n} \cdot \boldsymbol{\tau} \cdot \mathbf{n} = K(1 - Ca T), \qquad (8)$$

$$-\nabla T \cdot \mathbf{n} = Bi(T - T_a). \tag{9}$$

Here, j = 1, 2, $\nabla f = (\partial_r f, \frac{1}{r} \partial_0 f, \partial_z f)$, and f denotes a scalar. $\mathbf{u}_{\perp} = (v, w)$ represents a projection of \mathbf{u} on the $\theta - z$ orthogonal face. The unit normal vector \mathbf{n} and unit tangent vectors \mathbf{t}_1 and \mathbf{t}_2 at any point on the interface are given by

$$\mathbf{n} = \frac{1}{\sqrt{E}} \left(1, -\frac{1}{R} R_{\theta}, -R_z \right), \tag{10a}$$

$$\mathbf{t}_{1} = \frac{1}{\sqrt{(1+R_{z}^{2})E}} \left(\frac{1}{R}R_{\theta}, 1+R_{z}^{2}, -\frac{1}{R}R_{\theta}R_{z}\right),$$
(10b)

$$\mathbf{t}_2 = \frac{1}{\sqrt{1 + R_z^2}} (R_z, 0, 1), \tag{10c}$$

$$E = 1 + \frac{1}{R^2} R_{\theta}^2 + R_z^2.$$
 (10d)

The curvature K is defined as

$$K = -\nabla \cdot \mathbf{n} = -\left(\frac{1}{r}\frac{\partial(rn_1)}{\partial r} + \frac{1}{r}\frac{\partial(n_2)}{\partial \theta} + \frac{\partial(n_3)}{\partial z}\right)$$
$$= \left[R^2(1+R_z^2) + R_\theta^2\right]^{-\frac{3}{2}} \left\{RR_{zz}(R^2+R_\theta^2) - (1+R_z^2)(R^2-RR_{\theta\theta}) - 2RR_\theta R_z R_{\theta z} - 2R_\theta^2\right\}, \quad (11)$$

where the subscript represents the partial derivative. $Ca = S^{-1}R_B$ denotes the capillary number. T_a is the gas temperature near the interface.

Equation (6) shows the kinematic boundary condition, guaranteeing that the gas–liquid interface is smooth. Equation (7) indicates that the shear stress is balanced with the surface tension gradient generated by the thermocapillary effect. Equation (8) shows the stress balance in the normal direction. Equation (9) stands for the continuity of heat flux cross the liquid surface.

When the capillary number Ca satisfies

$$Ca = S^{-1}R_{\rm B} \ll 1, \tag{12}$$

the jet can be approximated as a parallel flow with perfectly cylindrical interface. We assume that the basic temperature exhibits a linear distribution in the *z*-direction,

$$\bar{\mathbf{u}} = (0, 0, \bar{w}(r)), \quad \bar{T}(r, z) = -z + \overline{T_b}(r).$$
(13)

Here, the superscript line denotes the basic flow, and $\overline{T_b}$ represents the radial temperature distribution. Therefore, the approximate solution of basic flow is determined as follows:¹³

$$(\bar{u},\bar{\nu})pprox 0,$$
 (14a)

$$\bar{w} \approx \frac{1}{2} S^{-\frac{1}{2}} R_{\rm B} \left(r^2 - \frac{1}{2} \right),$$
 (14b)

$$\bar{T} \approx -z - \frac{1}{32} Pr R_{\rm B} (1 - r^2)^2,$$
 (14c)

$$T_a \approx -z,$$
 (14d)

$$\bar{p} \approx 1 + 2Caz,$$
 (14e)

$$R \approx 1,$$
 (14f)

$$\bar{\tau} \approx \partial_r \bar{w}(r) \begin{bmatrix} 0 & 0 & 1 \\ 0 & 0 & 0 \\ 1 & 0 & 0 \end{bmatrix}.$$
(14g)

Next, the stability of the flow is investigated by the modal analysis. A small disturbance in the normal mode form is superimposed on the basic flow,

$$(\mathbf{u}, T, p, \tau, R) = \left(\bar{\mathbf{u}}, \bar{T}, \bar{p}, \bar{\tau}, 1\right) + \operatorname{Re}\left\{\left(\hat{\mathbf{u}}(r), \hat{T}(r), \hat{p}(r), \hat{\tau}(r), \hat{\xi}\right) \\ \times \exp\left[\sigma t + i(\alpha z + m\theta)\right]\right\}.$$
(15)

Here, the variables without superscript line stand for the perturbation. The eigenvalue $\sigma = \sigma_r + i\sigma_i$ consists of the growth rate σ_r and frequency σ_i . α denotes the axial wavenumber, and the integer *m* represents the azimuthal wavenumber.

Substituting Eq. (15) in the fundamental equations and linearizing the equations with respect to the perturbation quantities, we can derive that

$$\hat{u} + r\mathrm{D}\hat{u} + \mathrm{i}m\hat{v} + \mathrm{i}\alpha r\hat{w} = 0, \tag{16}$$

$$im\hat{\tau}_{12} + i\alpha r\hat{\tau}_{13} + rD\hat{\tau}_{11} + \hat{\tau}_{11} - \hat{\tau}_{22} - S^{\frac{1}{2}}i\alpha r\bar{w}\hat{u} = S^{\frac{1}{2}}rD\hat{p} + \sigma S^{\frac{1}{2}}r\hat{u},$$
(17a)

 $im\hat{\tau}_{22} + i\alpha r\hat{\tau}_{23} + rD\hat{\tau}_{12} + 2\hat{\tau}_{12} - S^{\frac{1}{2}}i\alpha r\bar{w}\hat{v} = S^{\frac{1}{2}}im\hat{p} + \sigma S^{\frac{1}{2}}r\hat{v}, \quad (17b)$

$$im\hat{\tau}_{23} + i\alpha r\hat{\tau}_{33} + rD\hat{\tau}_{13} + \hat{\tau}_{13} - S^{\frac{1}{2}}rD\bar{w}\hat{u} - S^{\frac{1}{2}}i\alpha r\bar{w}\hat{w}$$

= $S^{\frac{1}{2}}i\alpha r\hat{p} + \sigma S^{\frac{1}{2}}r\hat{w},$ (17c)

$$\{ r^2 D^2 + rD - \alpha^2 r^2 - m^2 - PrS^{\frac{1}{2}} \alpha r^2 \bar{w} \} \hat{T} - PrS^{\frac{1}{2}} r^2 (D\bar{T}\hat{u} + \bar{T}_z \hat{w})$$

= $\sigma PrS^{\frac{1}{2}} r^2 \hat{T},$ (18)

where $D \equiv \partial_r$ denotes the partial derivative with respect to *r*. Following Chaudhary *et al.*,²⁴ Eqs. (17) are written in a general form represented by the perturbation stress, which can extend to non-Newtonian fluids.

The linear perturbation constitutive equation for a Newtonian fluid is

$$\begin{cases} \hat{\tau}_{11} - 2D\hat{u} = 0, \quad r\hat{\tau}_{12} - im\hat{u} - rD\hat{v} + \hat{v} = 0, \\ \hat{\tau}_{13} - i\alpha\hat{u} - D\hat{w} = 0, \quad r\hat{\tau}_{22} - 2(im\hat{v} + \hat{u}) = 0, \\ r\hat{\tau}_{23} - i\alpha r\hat{v} - im\hat{w} = 0, \quad \hat{\tau}_{33} - 2i\alpha\hat{w} = 0. \end{cases}$$
(19)

The linear perturbation boundary conditions are written as follows. At the center axis r = 0,²⁵

$$\begin{cases} m = 0: \hat{u} = 0, \quad \hat{v} = 0, \quad D\hat{w} = 0, \quad D\hat{T} = 0; \\ m = 1: D\hat{u} = 0, \quad \hat{u} + i\hat{v} = 0, \quad \hat{w} = 0, \quad \hat{T} = 0; \\ m > 1: \hat{u} = 0, \quad \hat{v} = 0, \quad \hat{w} = 0, \quad \hat{T} = 0. \end{cases}$$
(20)

On the liquid surface at r = 1,

$$\hat{u} - i\alpha \bar{w}\hat{\xi} = \sigma \hat{\xi},\tag{21}$$

$$\hat{\tau}_{12} + S^{-\frac{1}{2}} R_{\rm B} im \left(2 D \bar{T} \hat{\xi} + \hat{T} \right) = 0,$$
 (22)

$$D\bar{\tau}_{13}\hat{\xi} + \hat{\tau}_{13} + S^{-\frac{1}{2}}R_{\rm B}i\alpha(2D\bar{T}\hat{\xi} + \hat{T}) = 0, \qquad (23)$$

$$-\hat{p} + S^{-\frac{1}{2}}(\hat{\tau}_{11} - 2i\alpha\bar{\tau}_{13}\hat{\xi}) - (1 - \alpha^2 - m^2)\hat{\xi} - Ca(D\bar{T}\hat{\xi} + \hat{T}) = 0,$$
(24)

$$Bi(\hat{T} + D\bar{T}\hat{\xi}) + D\hat{T} + (D^2\bar{T} - i\alpha\bar{T}_z)\hat{\xi} = 0.$$
(25)

The normal stress boundary condition (24) requires that the surface tension does not vary much over the characteristic axial wavelength of the disturbance, i.e.,

$$Ca \cdot 2\pi/\alpha \ll 1,$$
 (26)

thus, $(1 - Ca \,\overline{T}(z, 1))(1 - \alpha^2 - m^2)\hat{\xi} \approx (1 - \alpha^2 - m^2)\hat{\xi}$, where $\overline{T}(z, 1) = -z$.

We can use the Chebyshev collocation method to solve the general eigenvalue problem expressed as $Wg = \sigma Zg$, where W and Z are two matrices, and g is the eigenvector. Before solving the equations, \hat{p} in Eqs. (17) and (24) can be eliminated. The details on this process can be found in Appendix A. N_c Gauss–Lobatto points are set for the governing equations in the flow region $r = (1 - \cos(\frac{j\pi}{N_c+1}))/2$, $j = 1 \sim N_c$, and two extra points are set for the boundary conditions at r = 0,1.

Table I presents the eigenvalues of the neutral capillary mode calculated by using different numbers of collocation points. We choose 70–80 collocation points for our numerical solution, balancing the accuracy and the efficiency. Xu and Davis¹³ have given an approximate expression (6–2) that describes the dispersion relation for a non-isothermal jet when α is small. Table II presents a comparison of the eigenvalues obtained by Xu and Davis (XD) with those calculated in our present study (PW). The relative errors are less than 2.5×10^{-2} , confirming the dependability of our computational findings.

III. ENERGY ANALYSIS

The rate of change of perturbation kinetic energy is given by¹⁷

TABLE I. The eigenvalues of neutral capillary mode computed by different numbers of collocation points at Bi = 1, Pr = 0.1, $R_{\rm B} = 200$, $S = 10^4$, $\alpha = 0.864$, and m = 0.

N _c	$\sigma_r (\times 10^{-4})$	$\sigma_i (imes 10^{-2})$
30	4.179 40	4.843 199 3
50	4.179 40	4.843 199 3
80	4.179 40	4.843 199 3
100	4.179 40	4.843 199 4

TABLE II. A comparison of eigenvalues for the non-isothermal case at Bi = 1, Pr = 0.1, $R_{\rm B} = 10$, and $S = 10^2$. Here, "XD" and "PW" stand for the results of Xu and Davis and the present work, respectively.

α	XR	PW
0.01	0.02236 + 0.02211i	0.022 86 + 0.021 61i
	-0.02236 - 0.02261i	-0.02289 - 0.02211i
0.001	0.00707 + 0.00705i	0.00706 + 0.00703i
	$-0.00707 - 0.00710\mathrm{i}$	-0.00706 - 0.00708i

$$\partial_t E_k = -\frac{1}{2} S^{-\frac{1}{2}} \int (\boldsymbol{\tau} : \dot{\boldsymbol{\gamma}}) d^3 r + S^{-\frac{1}{2}} \int (\boldsymbol{u} \cdot \boldsymbol{\tau} \cdot \boldsymbol{n}) d^2 r$$
$$-\int \boldsymbol{u} \cdot ((\boldsymbol{u} \cdot \nabla) \bar{\boldsymbol{u}}) d^3 r - \int (p \boldsymbol{u} \cdot \boldsymbol{n}) d^2 r$$
$$= -N + M + I + P_S. \tag{27}$$

Here, $E_k = \frac{1}{2} \int |\mathbf{u}|^2 d^3 r$, whose integral area is $0 \le r \le 1, 0 \le \theta \le 2\pi$, $0 \le z \le 2\pi/\alpha$.

 $\int f d^3 r$ and $\int f d^2 r$ represent volume integral and surface integral, respectively. *N* is the work done by stress in the bulk of the jet, *M* is the work done by the stress on the surface, *I* is the energy from the basic flow, and *P*_S is the work done by the pressure on the surface. For Newtonian fluids, Eq. (18) indicates that N > 0, meaning that viscous dissipation always occurs in the liquid column, the same as the liquid layer.^{26,27} The perturbation energy is normalized as $\int |\mathbf{u}|^2 d^3 r = 1$ in the following.

On the deformable surface, there are horizontal and vertical components for the stress. As a result, M can be decomposed into two following terms:²⁸

$$M = S^{-\frac{1}{2}} \int u\tau_{11} d^2r + S^{-\frac{1}{2}} \int (v\tau_{12} + w\tau_{13}) d^2r = M_v + M_h.$$
 (28)

Here, M_v and M_h represent the work done by vertical and horizontal components of the stress, respectively. According to boundary conditions (21) and (22), the perturbed shear stress on the surface can be caused by both the thermocapillary effect and interfacial deformation. Therefore, M_h can be further decomposed into two parts as follows:

$$M_h = M_{hT} + M_{hR}, (29a)$$

$$M_{hT} = -S^{-1}R_{\rm B} \int \left(v \frac{\partial T}{\partial \theta} + w \frac{\partial T}{\partial z} \right) {\rm d}^2 r, \qquad (29b)$$

$$M_{hR} = -2S^{-1}R_{\rm B} \int v \mathrm{D}\bar{T} \frac{\partial\xi}{\partial\theta} \,\mathrm{d}^{2}r -S^{-\frac{1}{2}} \int w \left(\mathrm{D}\bar{\tau}_{13}\xi + 2S^{-\frac{1}{2}}R_{\rm B}\mathrm{D}\bar{T} \frac{\partial\xi}{\partial z} \right) \mathrm{d}^{2}r.$$
(29c)

Here, M_{hT} and M_{hR} stand for the work done by thermocapillary force and interfacial deformation-induced shear stress, respectively.

This paper uses the terminology suggested by Xu and Davis¹³ to label the capillary mode, surface-wave mode, and hydrodynamic mode with azimuthal wavenumber m = 0 as $W_0^{(c)}$, $W_0^{(-)}$, and $W_0^{(+)}$, respectively. When the dispersion relations of the capillary mode and surface wave mode are connected in a curve, α_B is the wave number corresponding to the transition point of the two modes. The capillary mode $W_0^{(c)}$ is for $\alpha < \alpha_{\rm B}$, and surface-wave mode $W_0^{(-)}$ is for $\alpha > \alpha_{\rm B}$. Xu and Davis¹³ have also found that for isothermal jets (axial wavenumber α approximately less than 2) or non-isothermal jets, some unstable modes m = 0 become the most dangerous mode. In some cases, the increase in $R_{\rm B}$ can suppress the unstable capillary modes. In order to comprehend the underlying energy mechanism, we conduct the energy analysis on the capillary mode $W_0^{(c)}$, surface-wave mode $W_0^{(-)}$, and hydrodynamic mode $W_0^{(+)}$.

A. The "flying jet"

When $Bi \to \infty$, we can obtain from Eq. (25) that T(1) = 0, and there are no temperature perturbations on the free surface. Therefore, all unstable modes are actually isothermal and independent of *Pr*. The basic temperature field is important only in that the thermocapillary force drives the basic flow, which can be considered as driven by the "wind stress" having the same magnitude as the thermocapillary gradient γb . This situation is referred to as the isothermal "flying jet" by Xu and Davis.¹³ Thus, *R*_B serves as the Reynolds number of the flying jet. Such isothermal cases do not require the "slowly varying" approximation, i.e., Eq. (26).²⁹ Therefore, all physical modes need only satisfy the constraint (12). In these cases, the dispersion curves of the capillary mode $W_0^{(c)}$ and surface-wave mode $W_0^{(-)}$ are connected smoothly.

Tables III and IV present the terms of the perturbation energy growth for the isothermal jet for the capillary mode $W_0^{(c)}$, surfacewave mode $W_0^{(-)}$, and hydrodynamic mode $W_0^{(+)}$ at $\alpha = 0.4 < \alpha_B$, $S = 10^4$. Tables V and VI present the cases for the isothermal jet for surface-wave mode $W_0^{(-)}$ and hydrodynamic mode $W_0^{(+)}$ at $\alpha = 1.4 > \alpha_B$ and $S = 10^4$. It is evident that thermocapillary forces do no work ($M_{hT} = 0$), which is obvious in the isothermal case.

For $0 < \alpha < \alpha_{\rm B}$, N is the viscous dissipating in the capillary mode $W_0^{(c)}$ and hydrodynamic mode $W_0^{(+)}$. P_S is the main energy dissipation for the stable hydrodynamic mode $W_0^{(+)}$, but the main energy source in the unstable capillary mode $W_0^{(c)}$. When P_S significantly decreases with R_B , the capillary mode $W_0^{(c)}$ is stabilized. For $\alpha > \alpha_{\rm B}$, P_S , M and I serve as energy sources, while N is the only pathway of energy dissipation for the surface-wave mode $W_0^{(-)}$. For the

TABLE III. The terms of the perturbation energy growth for the isothermal jet for the capillary mode $W_0^{(c)}$ and the surface-wave mode $W_0^{(-)}$ at $\alpha = 0.4$ and $S = 10^4$.

Mode		$W_0^{(c)}$		$W_0^{(-)}$
R _B	100	200	300	500
$\alpha_{\rm B}$	0.9439	0.8175	0.5158	0.2140
$\partial_t E_k$	0.237 63	0.18071	0.046 31	-0.01107
P_S	0.229 42	0.14878	0.024 52	0.00071
-N	-0.00599	-0.00940	-0.01340	-0.00700
Ι	0.004 66	0.01208	-0.00782	-0.03030
M	0.009 54	0.029 25	0.043 01	0.025 51
M_{hR}	0.007 83	0.027 14	0.040 52	0.023 65
M_{hT}	0.000 00	0.000 00	0.000 00	0.000 00
M_{ν}	0.001 71	0.002 11	0.002 49	0.001 86

TABLE IV.	The terms of	the perturbation	on energy	growth f	or the	isothermal	jet for	the
hydrodynar	nic mode $W_0^{(+)}$) at $\alpha = 0.4$ a	and $S = 1$	0 ⁴ .				

Mode	$W_0^{(+)}$					
R _B	100	200	300	500		
$\partial_t E_k$	-0.23703	-0.19520	-0.11406	-0.05977		
P_S	-0.24232	-0.20449	-0.15339	-0.20345		
-N	-0.01250	-0.03022	-0.04336	-0.15402		
Ι	0.010 11	0.00947	0.006 58	0.032 45		
Μ	0.007 68	0.030 04	0.07610	0.265 25		
M_{hR}	0.005 68	0.027 36	0.07294	0.261 00		
M_{hT}	0.00000	0.00000	0.00000	0.000 00		
M_{ν}	0.001 99	0.002 68	0.003 17	0.004 24		

TABLE V. The terms of the perturbation energy growth for the isothermal jet for the surface-wave mode $W_0^{(-)}$ at $\alpha = 1.4 > \alpha_B$ and $S = 10^4$.

Mode	$W_{0}^{(-)}$					
R _B	100	200	300	500		
$\alpha_{\rm B}$	0.9439	0.8175	0.5158	0.2140		
$\partial_t E_k$	-0.01209	-0.00045	0.006 08	0.010 91		
P_S	0.004 56	0.006 55	0.00944	0.013 03		
-N	-0.05871	-0.05966	-0.06026	-0.06084		
Ι	0.011 02	0.01495	0.01572	0.014 53		
M	0.031 03	0.037 70	0.041 18	0.044 19		
M_{hR}	0.01073	0.016 45	0.01944	0.022 00		
M_{hT}	0.000 00	0.000 00	0.000 00	0.000 00		
M_{ν}	0.020 31	0.021 25	0.02174	0.022 19		

TABLE VI. The terms of the perturbation energy growth for the isothermal jet for the hydrodynamic mode $W_0^{(+)}$ at $\alpha = 1.4$ and $S = 10^4$.

Mode	$W_0^{(+)}$						
R _B	100	200	300	500			
$\partial_t E_k$	-0.04723	-0.063 10	-0.07520	-0.09114			
P_S	0.039 66	0.113 27	0.245 46	0.663 77			
-N	-0.05868	-0.06794	-0.09690	-0.26321			
Ι	-0.02623	-0.08102	-0.17868	-0.51614			
М	-0.00197	-0.02740	-0.04508	0.02444			
M_{hR}	-0.01742	-0.03823	-0.05016	0.030 20			
M_{hT}	0.000 00	0.000 00	0.000 00	0.000 00			
M_{ν}	0.015 44	0.01084	0.005 09	-0.00576			

hydrodynamic mode $W_0^{(+)}$, P_S is still the energy source, while *N* and *I* dissipate the energy. The sign of *M* depends on the parameters.

The physical reason for these phenomena can be explained as follows. In the inviscid jet $[R_B = O(1) \text{ as } S \rightarrow \infty]$, ¹³ N = M = I = 0 in Eq. (27), suggesting that P_S is the main energy source for the unstable

Mode	$R_{\rm B}$	α_M	$\partial_t E_k$	P_S	-N	Ι	М	M_{hR}	M_{hT}	M_{ν}
$W_0^{(c)}$	0	0.690	0.3363	0.3457	-0.0138	0.0000	0.0045	0.0000	0.0000	0.0045
	100	0.654	0.3015	0.2928	-0.0143	0.0075	0.0156	0.0110	0.0000	0.0046
	200	0.532	0.1989	0.1643	-0.0138	0.0138	0.0346	0.0308	0.0000	0.0038
	300	0.273	0.0646	0.0330	-0.0100	0.0018	0.0399	0.0387	0.0000	0.0012
	500	0.051	0.0241	0.0024	-0.0051	-0.0002	0.0270	0.0270	0.0000	0.0000
$W_0^{(-)}$	200	1.051	0.0084	0.0082	-0.0344	0.0008	0.0337	0.0214	0.0000	0.0123
	300	1.141	0.0096	0.0091	-0.0403	0.0050	0.0358	0.0213	0.0000	0.0145
	500	1.302	0.0112	0.0122	-0.0526	0.0104	0.0412	0.0221	0.0000	0.0191

TABLE VII. The terms of perturbation energy growth for the most unstable modes at $S = 10^4$.

capillary mode $W_0^{(c)}$. This mechanism still keeps in the flying jet. For the hydrodynamic mode $W_0^{(+)}$ at $S = 10^4$, it can be seen from Eq. (24) that $-\hat{p}(1) \approx (1 - \alpha^2)\hat{\xi}$. The phase difference between the $\hat{p}(1)$ and $\hat{\xi}$ at $\alpha < 1$ is opposite to that at $\alpha > 1$, while the phase difference between \hat{u} and $\hat{\xi}$ is not very sensitive to α . As a result, the sign of P_S is different in these two cases of $W_0^{(+)}$. The surface-wave mode $W_0^{(-)}$ is caused by the "wind stress,"²⁹ so the work done by the stress on the surface (*M*) is crucial for $W_0^{(-)}$.

In Table III, the growth rate of perturbation energy for the capillary mode $W_0^{(c)}$ gradually decreases with $R_{\rm B}$. The explanation is given in the last part of Sec. IV A. However, the case is opposite for the surface-wave mode $W_0^{(-)}$ in Table V. The reason is that $W_0^{(-)}$ is caused by the wind stress, whose magnitude can be measured by $R_{\rm B}$. This indicates that the velocity distribution in Eq. (14b) caused by the wind stress²⁹ can suppress capillary instabilities, allowing for longer continuous jets, but trigger surface-wave instabilities.

Table VII displays the perturbation energy growth for the most unstable modes at $S = 10^4$, where α_M is the corresponding wavenumber. As $R_{\rm B}$ increases, the perturbation energy growth decreases for the capillary mode $W_0^{(c)}$, while the opposite occurs for the surface-wave mode $W_0^{(-)}$, in agreement with the preceding analysis. It should be noted that for $R_{\rm B} \ge 200$, both $W_0^{(c)}$ and $W_0^{(-)}$ are unstable. However, the growth rates of $W_0^{(c)}$ and $W_0^{(-)}$ are smaller than those of the capillary mode $W_0^{(c)}$ at $R_{\rm B} = 0$ and $R_{\rm B} = 100$. Therefore, the increase in $R_{\rm B}$ can still slow down the process of capillary breakup, extending the jet zone.

TABLE VIII. The terms of the perturbation energy growth for the capillary mode $W_0^{(c)}$ at $R_B = 100$, $\alpha = 0.4$, and $S = 10^4$.

$W_0^{(c)}$	$Bi \rightarrow \infty$	Bi = 100 $Pr = 0.1$	Bi = 1 $Pr = 0.1$	Bi = 1 $Pr = 0.5$
$\partial_t E_k$	0.237 63	0.237 58	0.234 97	0.232 20
P_S	0.229 42	0.229 57	0.236 87	0.243 12
-N	-0.00599	-0.00595	-0.00479	-0.00494
Ι	0.004 66	0.004 59	0.001 09	0.000 14
M_{hR}	0.007 83	0.007 80	0.006 49	0.005 92
M_{hT}	0.00000	-0.00013	-0.00625	-0.01342
M_{ν}	0.001 71	0.001 71	0.001 57	0.001 37

B. Thermocapillary instabilities

When *Bi* is finite, the jet becomes non-isothermal, both restrictions (12) and (26) should be satisfied for the physical modes.¹³ If $R_{\rm B} \neq 0$, thermocapillary forces become the driving force for the surface flow.

Xu and Davis¹³ have suggested that the modes m = 0 at large Bi $(Bi \ge 1)$ and small Pr (Pr < 1) are similar to those in the isothermal cases. Table VIII shows the terms of the perturbation energy growth for the capillary mode $W_0^{(c)}$ at $R_{\rm B} = 100$, $\alpha = 0.4$, and $S = 10^4$. It can be seen that for unstable capillary modes $W_0^{(c)}$, P_S is dominant and changes little. Thus, their energy mechanisms are also similar.

1. The effect of R_B

For large Bi ($Bi \ge 1$) and small Pr (Pr < 1), as $R_{\rm B}$ increases, the capillary mode $W_0^{\rm (c)}$ and surface-wave mode $W_0^{\rm (-)}$ split. In addition, when $R_{\rm B}$ is not large, the surface-wave mode $W_0^{\rm (-)}$ generates a new branch $W_0^{\rm (-)'}$, and the hydrodynamic mode $W_0^{\rm (+)}$ also has a new branch $W_0^{\rm (+)'}$. As $R_{\rm B}$ increases, the branch $W_0^{\rm (-)'}$ disappears, while branch $W_0^{\rm (+)'}$ becomes more stable (see Figs. 6 and 7 in Ref. 13).

Tables IX and X show the terms of the perturbation energy growth of the non-isothermal jet for the modes $W_0^{(c)}$ and $W_0^{(+)}$, respectively. In Table IX, when R_B is small, the capillary mode $W_0^{(c)}$ mainly absorbs energy from P_S . However, as R_B increases, the energy

TABLE IX. The terms of the perturbation energy growth for the non-isothermal jet for the capillary mode $W_0^{(c)}$ at $\alpha = 0.6$, Pr = 0.1, Bi = 1, and $S = 10^4$.

Mode	$W_0^{(c)}$						
R _B	40	100	200	300			
$\partial_t E_k$	0.32177	0.294 12	0.181 26	0.013 89			
P_S	0.328 80	0.298 84	0.157 17	-0.00280			
-N	-0.01052	-0.01063	-0.01370	-0.04451			
Ι	0.000 22	0.002 30	0.01403	-0.04033			
M	0.003 27	0.003 61	0.02376	0.101 53			
M_{hR}	0.001 35	0.008 51	0.03212	0.051 67			
M_{hT}	-0.00150	-0.00833	-0.01270	0.042 80			
M_{ν}	0.003 42	0.003 43	0.004 34	0.007 05			

TABLE X.	The terms of t	ne perturbation	energy grow	th for the nor	i-isothermal jet for
the hydrod	ynamic mode V	$V_0^{(+)}$ at $\alpha = 0.0$	6, <i>Pr</i> = 0.1, <i>B</i>	i = 1, and S	= 10 ⁴ .

TABLE XII. The terms of the perturbation energy growth for the non-isothermal jet for the surface-wave mode $W_0^{(-)}$ at $\alpha = 1.4$, Pr = 0.1, Bi = 1, and $S = 10^4$.

Mode	$W_0^{(+)}$						
R _B	40	100	200	300			
$\partial_t E_k$	-0.192 36	-0.20102	-0.15456	-0.09443			
P_S	-0.03237	-0.18837	-0.31204	-0.36133			
-N	-0.16742	-0.17061	-0.18979	-0.36929			
Ι	0.006 30	0.051 55	0.113 12	0.15170			
M	0.001 13	0.106 40	0.23415	0.484 49			
M_{hR}	-0.00191	-0.00661	0.009 82	-0.01264			
M_{hT}	0.002 96	0.109 65	0.217 53	0.490 55			
M_{ν}	0.000 08	0.003 36	0.006 81	0.006 57			

contribution from P_S decreases rapidly. As a result, the rate of change of perturbation kinetic energy also decreases. When R_B further increases, M becomes the main energy source, while the other terms are dissipative. However, the growth rate of capillary mode $W_0^{(c)}$ largely decreases, and the flow becomes more stable. Table X indicates that the energy for hydrodynamic mode $W_0^{(+)}$ mainly comes from M_{hT} , but is dissipated by N and P_S . When R_B is large enough, M_{hT} dominates.

If Pr becomes larger, new unstable modes appear. Table XI shows the terms of the perturbation energy growth for the non-isothermal jet for the hydrodynamic mode $W_0^{(+)}$ at different $R_{\rm B}$. It can be seen that when $R_{\rm B}$ increases, M_{hT} becomes dominant and $W_0^{(+)}$ becomes more unstable. It should be noted that $W_0^{(+)}$ gives rise to a new unstable mode known as the hydrothermal instability,¹³ which exhibits a completely different energy mechanism compared to the unstable modes in isothermal conditions.

Table XII presents the energy growth for the non-isothermal jet for the surface-wave mode $W_0^{(-)}$ at $\alpha = 1.4$, Pr = 0.1, Bi = 1, and $S = 10^4$. It is found that the mode is dissipated by N and M_{hT} . As R_B increases, M_{hT} becomes more negative, N changes little, while other items increase. The growth rate of perturbation energy becomes positive, indicating the appearance of surface-wave instability.

Table XIII shows the perturbation energy growth for the nonisothermal jet for the new branch of surface-wave mode $W_0^{(-)'}$, the

TABLE XI. The terms of the perturbation energy growth for the non-isothermal jet for the hydrodynamic mode $W_0^{(+)}$ at $\alpha = 1$, Pr = 5, Bi = 1, and $S = 10^4$.

Mode	$W^{(+)}_0$					
R _B	40	100	200	300		
$\partial_t E_k$	-0.003 55	0.02667	0.07283	0.115 87		
P_S	-0.07425	-0.08102	-0.09008	-0.09719		
-N	-0.31896	-0.54984	-0.86176	-1.13326		
Ι	-0.01303	-0.03578	-0.06688	-0.09362		
M_{hR}	-0.04571	-0.05471	-0.06291	-0.06847		
M_{hT}	0.429 03	0.728 25	1.13412	1.487 67		
M_{ν}	0.019 36	0.01976	0.02034	0.02073		

Mode	$W_0^{(-)}$				
R _B	40	100	200	300	
$\partial_t E_k$	-0.02244	-0.01271	0.001 67	0.011 58	
Р	0.009 98	0.010 82	0.01422	0.018 00	
-N	-0.05713	-0.05713	-0.05766	-0.05806	
Ι	0.006 27	0.01440	0.02464	0.030 69	
M	0.018 44	0.01921	0.02047	0.020 95	
M_{hR}	0.004 91	0.010 19	0.01544	0.018 19	
M_{hT}	-0.00481	-0.00947	-0.01399	-0.01660	
M_{ν}	0.018 34	0.018 50	0.019 02	0.019 37	

surface-wave mode $W_0^{(-)}$, and the capillary mode $W_0^{(c)}$. Figure 7 in Ref. 13 suggests that $W_0^{(-)'}$ gradually disappears as $R_{\rm B}$ increases. In Table XIII, the branch $W_0^{(-)'}$ at $R_{\rm B} = 40$ and the surface-wave mode $W_0^{(-)}$ at $R_{\rm B} = 100$ have similar values for each term. Hence, it is reasonable to infer that when $R_{\rm B}$ increases, the branch $W_0^{(-)'}$ at $0 < \alpha < \alpha_{\rm B}$ becomes the surface-wave mode $W_0^{(-)}$, while the branch $W_0^{(-)'}$ at $\alpha > \alpha_{\rm B}$ becomes the capillary mode $W_0^{(c)}$. As the branch of hydrodynamic mode $W_0^{(+)}$ becomes more stable as $R_{\rm B}$ increases, its energy analysis is omitted.

Table XIV shows the energy mechanism for the most unstable modes at Pr=0.5, Bi=1 and $S=10^4$. As $R_{\rm B}$ increases, $\partial_t E_k$ decreases for $W_0^{\rm (c)}$, while the variation of $\partial_t E_k$ for $W_0^{\rm (-)}$ is not monotonous. On the other hand, $\partial_t E_k$ of $W_0^{\rm (+)}$ increases. This suggests that the appearance of surface-wave and hydrothermal instabilities suppress the capillary breakup and allow for a longer jet region. Although unstable capillary modes $W_0^{\rm (c)}$ at $R_{\rm B}=350$ and 500 do not fully satisfy (26), the suppression of capillary instability still occurs.

TABLE XIII. The terms of the perturbation energy growth for the non-isothermal jet for the new branch of surface-wave mode $W_0^{(-)'}$ at Pr = 0.1, Bi = 1, $S = 10^4$, and $R_B = 40$ and the surface-wave mode $W_0^{(-)}$ and the capillary mode $W_0^{(c)}$ at Pr = 0.1, Bi = 1, $S = 10^4$, and $R_B = 100$.

Mode	$W_{0}^{(-)'}$	$W_{0}^{(-)}$	$W_{0}^{(-)'}$	$W_0^{(c)}$
$R_{\rm B}$	40	100	40	100
$\alpha_{\rm B}$	0.9789	_	0.9789	_
α	$\alpha = 0.4$	$4 < \alpha_{\rm B}$	$\alpha = 1.2$	$2 > \alpha_{\rm B}$
P_S	-0.00498	-0.02141	0.008 06	0.010 67
-N	-0.14649	-0.13183	-0.04106	-0.04106
Ι	0.004 58	0.011 58	0.005 55	0.012 47
M_{hR}	-0.00108	-0.00371	0.00661	0.012 50
M_{hT}	0.000 51	0.010 58	-0.00544	-0.01040
M_{ν}	-0.00026	-0.00096	0.012 82	0.012 89

Mode	$R_{\rm B}$	α_M	$\partial_t E_k$	P_S	-N	Ι	М	M_{hR}	M_{hT}	M_{ν}
$W_0^{(c)}$	40	0.683	0.3292	0.3416	-0.0135	-0.0004	0.0014	0.0014	-0.0042	0.0042
-	100	0.649	0.2934	0.3088	-0.0126	0.0035	-0.0064	0.0091	-0.0189	0.0034
	200	0.574	0.1957	0.1519	-0.0175	0.0313	0.0299	0.0320	-0.0062	0.0041
	350	0.562	0.0814	-0.0045	-0.0818	-0.0193	0.1871	0.0576	0.1220	0.0075
	500	0.680	0.0546	-0.0453	-0.1915	-0.0663	0.3577	0.0693	0.2760	0.0124
$W_0^{(-)}$	100	1.015	0.0129	0.0207	-0.0287	0.0166	0.0044	0.0169	-0.0205	0.0079
	200	0.974	0.0288	0.0261	-0.0272	0.0303	-0.0004	0.0186	-0.0262	0.0073
	350	0.853	0.0414	0.0291	-0.0223	0.0434	-0.0089	0.0194	-0.0342	0.0058
	500	0.820	0.0373	0.0281	-0.0212	0.0422	-0.0118	0.0202	-0.0380	0.0060
$W_0^{(+)}$	350	1.547	0.0086	-0.3108	-0.8601	-0.0008	1.1803	-0.1390	1.2427	0.0767
	500	1.821	0.0418	-0.4490	-1.4131	0.0046	1.8993	-0.1630	1.9501	0.1123

TABLE XIV. The rate of change of perturbation energy for the most unstable modes at Pr = 0.5, Bi = 1, and $S = 10^4$.

TABLE XV. The terms of the perturbation energy growth for the non-isothermal jet for the capillary mode $W_0^{(c)}$ at $\alpha = 0.6$, $R_{\rm B} = 300$, Bi = 1, and $S = 10^4$.

Mode	$W_0^{(c)}$				
Pr	1	5	10		
$\partial_t E_k$	0.139 129	0.197 955	0.210 977		
P_S	0.022 032	0.025 869	0.027774		
-N	-0.078454	-0.124960	-0.141570		
Ι	0.006 044	0.002 586	-0.005104		
M	0.189 507	0.294 459	0.329 876		
M_{hR}	0.055 316	0.058 873	0.059 445		
M_{hT}	0.125715	0.226 050	0.260 630		
M_{ν}	0.008 476	0.009 536	0.009 801		

2. The effect of Pr

For large *Bi*, as *Pr* increases, the capillary mode $W_0^{(c)}$ and surfacewave mode $W_0^{(-)}$ are also distinctly separated (see Figs. 11 and 12 in Ref. 13). Tables XV and XVI present the perturbation energy growth of the non-isothermal jet for the capillary mode $W_0^{(c)}$ and hydrodynamic mode $W_0^{(+)}$, respectively. In Table XV, as *Pr* increases, the

TABLE XVI. The terms of the perturbation energy growth for the non-isothermal jet for the hydrodynamic mode $W_0^{(+)}$ at $\alpha = 1$, $R_B = 300$, Bi = 1, and $S = 10^4$.

Mode	$W_0^{(+)}$				
Pr	1	5	10		
$\partial_t E_k$	0.020 581	0.115 874	0.151 824		
P_S	-0.103330	-0.097185	-0.097212		
-N	-0.950970	-1.133257	-1.200462		
Ι	-0.095425	-0.093619	-0.091977		
M_{hR}	-0.102894	-0.068470	-0.060731		
M_{hT}	1.252 300	1.487 674	1.581 481		
M_{ν}	0.020 900	0.020730	0.020725		

energy dissipated by *N* increases and *I* changes from positive to negative. However, both P_S and *M* increase. The increase in $\partial_t E_k$ indicates that $W_0^{(c)}$ becomes more unstable.

In Table XVI, *N* is the main energy dissipation, while M_{hT} becomes the main energy source. Thus, $W_0^{(+)}$ belongs to the hydrothermal instability. In Tables XI and XVI, it can be found that the increase in $R_{\rm B}$ or Pr leads to the increase in M_{hT} and the hydrothermal instability. Therefore, if we define the Marangoni number as $Ma_{\rm B} \equiv PrR_{\rm B}$, then the increase in $M_{\rm aB}$ can also lead to unstable hydrodynamic modes.

Table XVII presents the perturbation energy growth for the nonisothermal jet for the most unstable surface-wave modes $W_0^{(-)}$ at $R_{\rm B} = 300$, Bi = 1, and $S = 10^4$. It can be seen that $P_{\rm S}$, M_{hR} , and I are energy sources for $W_0^{(-)}$, while N and M_{hT} are dissipative. However, the growth rates of unstable modes in Table XVII are relatively small compared to those in Tables XV and XVI. Since the new branch of the hydrodynamic mode $W_0^{(+)'}$ remains stable, its energy analysis is omitted.

IV. PERTURBATION FLOW FIELD

Why the thermocapillary effect can suppress the capillary breakup? In order to provide a clearer explanation for this question, the perturbed flow and temperature distributions for various unstable modes are presented in this section. In Figs. 2–10, the y axis represents

TABLE XVII. The terms of the perturbation energy growth for the non-isothermal jet for the most unstable surface-wave modes $W_0^{(-)}$ at $R_B = 300$, Bi = 1, and $S = 10^4$.

Mode	$W_0^{(-)}$				
Pr	1	5	10		
α_M	0.839	0.630	0.507		
$\partial_t E_k$	0.049 409	0.062 852	0.053 487		
P_S	0.035 065	0.041 050	0.038 012		
-N	-0.022659	-0.015382	-0.010087		
Ι	0.052736	0.054 590	0.038 231		
M_{hR}	0.018 601	0.020795	0.022 140		
M_{hT}	-0.039304	-0.041185	-0.036842		
M_{ν}	0.004 971	0.002 985	0.002 033		

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FIG. 3. The perturbation flow field of the most unstable modes at $R_{\rm B} = 200$ and $S = 10^4$: (a) the capillary mode $W_0^{(c)}$ and (b) the surface-wave $W_0^{(-)}$.



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FIG. 5. The perturbation flow field of the most unstable modes at $R_{\rm B} = 500$ and $S = 10^4$: (a) the capillary mode $W_0^{(c)}$ and (b) the surface-wave mode $W_0^{(-)}$.

the perturbation phase $\Phi = \alpha z + m\theta$. The x axis represents the jet radius, and $\xi_R = \operatorname{Re}\{\hat{\xi} \exp[i(\alpha z + m\theta)]\}$ represents the surface deformation. For convenience, all complex amplitudes of perturbations $\hat{\phi}$ below are relabeled as ϕ in the following.

A. The "flying jet"

Figures 2–5 show the perturbation flow fields of the most unstable modes at $R_{\rm B} = 0-500$ and $S = 10^4$. As these modes are isothermal, there are no temperature perturbations. The capillary mode $W_0^{(c)}$ in Fig. 2(a) is stationary, while other capillary modes $W_0^{(c)}$ and the surface-wave modes $W_0^{(-)}$ propagate along the negative y axis. As $R_{\rm B}$



FIG. 6. The perturbation flow field of the most unstable capillary mode $W_0^{(c)}$ at $R_B = 40$, Pr = 0.5, Bi = 1, and $S = 10^4$.

increases, the amplitude of ζ_R decays for these modes, which effectively prevents the jet from splitting.

The reason why the capillary breakup in isothermal jets can be suppressed is explained as follows. Tables III and VII suggest that the main energy source of the capillary instability is the work done by the pressure on the surface (P_S). For the capillary modes $W_0^{(c)}$ (m = 0) at $S = 10^4$, the surface pressure perturbation $-p(1) \approx (1 - \alpha_M^2)\xi$, where $\alpha_M < 1$ (Table VII). In Fig. 2(a), $R_B = 0$, $\bar{w}(1) = \frac{1}{4}S^{-\frac{1}{2}}R_B = 0$, $-p(1) \approx 0.5239\xi$. On the liquid surface r = 1, $\Delta \varphi = \arg(u|_{r=1})$ $-\arg(\xi_R) = 0$, $\int (u|_{r=1}\xi_R)dr^2 \propto \cos(\Delta \varphi)$ (as shown in Appendix B). So it can be seen from the expression of P_S in Eq. (27) that the capillary mode $W_0^{(c)}$ can extract energy from the surface pressure, which may cause the capillary breakup. However, $\bar{w}(1)$ increases with R_B . According to the continuous boundary condition (21), the phase difference between $u|_{r=1}$ and ξ_R also increases rapidly, leading to the decrease in P_S . Then, the capillary breakup is retarded.

B. Thermocapillary instabilities

Figures 6–10 show the perturbation flow fields of the most unstable modes at $R_{\rm B} = 40-500$, Pr=0.5, Bi=1, and $S=10^4$. For the capillary mode $W_0^{(c)}$, there are obvious temperature fluctuations throughout the flow region [Figs. 6, 7(a), 8(a), 9(a), and 10(a)]. For the surface-wave mode $W_0^{(-)}$, the temperature fluctuations are very weak near the interface [Figs. 7(b), 8(b), 9(b), and 10(b)]. On the contrary, the perturbation is concentrated in the thin layer near the liquid surface for the hydrodynamic mode $W_0^{(+)}$ [Figs. 9(c) and 10(c)]. The capillary modes $W_0^{(c)}$ and surface-wave modes $W_0^{(-)}$ propagate along the negative y axis, whereas the hydrodynamic modes $W_0^{(+)}$ propagate in the opposite direction. The surface perturbation amplitude always decreases with the increase in $R_{\rm B}$ for all modes, which can prevent the capillary breakup.

We provide an explanation for the suppression of capillary breakup in non-isothermal jets as follows. Table IX shows that the work done by surface pressure (P_S) is still the main energy source of the capillary breakup. For the capillary modes $W_0^{(c)}$ (m=0) at $S=10^4$, Pr=0.5, and Bi=1, the normal stress boundary condition (24) and thermal equilibrium condition (25) on the interface can be simplified as



FIG. 7. The perturbation flow field of the most unstable modes at $R_{\rm B} = 100$, Pr $= 0.5, Bi = 1, and S = 10^4$: (a) the capillary mode $W_0^{(c)}$ and (b) the surface-wave $W_0^{(-)}$.



$$p(1) = 2S^{-1}R_{\rm B}\frac{\partial\xi}{\partial z} - S^{-\frac{1}{2}}\tau_{11} + (1 - \alpha^2)\xi + S^{-1}R_{\rm B}T(1), \quad (30)$$

$$\frac{\partial \xi}{\partial z} = -T(1) - DT(1) + \frac{1}{8}R_{\rm B}\xi,$$
 (31)

respectively. Substituting Eq. (31) in Eq. (30) results in

$$-p(1) = \left(\frac{1}{4}S^{-1}R_{\rm B}^{2} + 1 - \alpha^{2}\right)\xi - S^{-1}R_{\rm B}[T(1) + 2DT(1)] - 2S^{-\frac{1}{2}}Du(1).$$
(32)

When $R_{\rm B}$ is not large, $-p(1) \approx (1 - \alpha_M^2)\xi$ can still be obtained. In Fig. 7, $R_{\rm B} = 40$, $\alpha_M = 0.683$, then $\bar{w}(1) = \frac{1}{4}S^{-\frac{1}{2}}R_{\rm B} = 0.1$, -p(1) $\approx 0.5335\xi$, $\arg(u|_{r=1}) - \arg(\xi_R) \approx 11.6^\circ$. Owing to the proper phase relationship, the key to the capillary breakup is still the work done by surface pressure caused by surface deformation. For $R_{\rm B} > 200, -p(1)$ $\approx (\frac{1}{4}S^{-1}R_{\rm B}^2 + 1 - \alpha_M^2)\xi$. However, since $\bar{w}(1)$ is increased by $R_{\rm B}$, the phase difference between $u|_{r=1}$ and ξ_{R} also increases. For example, when $R_{\rm B} = 500$, $\arg(u|_{r=1}) - \arg(\xi_R) \approx 86.1^{\circ}$. Therefore, P_S decreases significantly. In addition, in Fig. 10(a), due to the axisymmetric perturbation flow field (v=0), $\arg(w|_{r=1}) - \arg(\xi_R) \approx 171.1^{\circ}$ and $\arg(w|_{r=1})$ $-\arg\left(\frac{\partial T}{\partial z}\Big|_{r=1}\right) \approx 167.0^{\circ}$, we can drive that from Eqs. (28) and (29),

$$M \approx -S^{-1}R_{\rm B} \int \left(w\xi + w\frac{\partial T}{\partial z}\right) {\rm d}^2 r.$$
 (33)

Then, M becomes the main energy source for the capillary mode. However, the thermocapillary flow also increases the viscous





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FIG. 9. The perturbation flow field of the most unstable modes at $R_B = 350$, Pr = 0.5, Bi = 1, and $S = 10^4$: (a) the capillary mode $W_0^{(c)}$; (b) the surface-wave mode $W_0^{(-)}$; and (c) the hydrodynamic mode $W_0^{(+)}$.



FIG. 10. The perturbation flow field of the most unstable modes at $R_{\rm B} = 500$, Pr = 0.5, Bi = 1, and $S = 10^4$: (a) the capillary mode $W_0^{(c)}$; (b) the surface-wave mode $W_0^{(-)}$; and (c) the hydrodynamic mode $W_0^{(+)}$.

dissipation (*N*) and the energy transferred to the basic flow (*I*). As a result, the capillary instability is still greatly suppressed.

V. CONCLUSION

In this paper, the energy analysis is performed for instabilities in the cylindrical jet with thermocapillarity. The perturbation flow field is displayed for the capillary mode, surface-wave mode, and hydrodynamic mode. The mechanism of suppressing capillary breakup of the jet is elucidated.

When the Biot number tends to infinity $(Bi \rightarrow \infty)$, the flow is an isothermal jet driven by the wind stress of the same magnitude as the thermocapillary gradient. The work done by the pressure on the

surface (P_S) is the main energy source for the capillary mode. As the Reynolds number of the flying jet (R_B) increases, P_S decreased rapidly. Therefore, the capillary breakup is suppressed.

For a small Prandtl number (Pr < 1) and a large Bi ($Bi \ge 1$), when the Reynolds number (R_B) is not too large, P_S remains the main energy source for capillary instability. However, the phase difference between radial velocity and surface deformation increases with R_B , leading to a rapid decrease in P_S . When R_B is large enough, the work done by the stress on the surface (M) is dominant, but the dissipation caused by the viscosity (N) and the energy coming from the basic flow (I) also increase. Therefore, the capillary instability is also suppressed.

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The increase in $R_{\rm B}$ or Pr enhances the energy obtained from the thermocapillary force (M_{hT}), making it the dominant factor and leading to the hydrothermal instability.

For the capillary mode, there are obvious temperature fluctuations throughout the flow region, and for the surface-wave mode, the temperature fluctuations are very weak near the surface, while for the hydrodynamic mode, the hot spot is always found in a thin layer near the liquid surface.

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AUTHOR DECLARATIONS

Conflict of Interest

The authors have no conflicts to disclose.

Author Contributions

YuWen Sun: Conceptualization (equal); Data curation (lead); Formal analysis (equal); Investigation (equal); Methodology (equal); Software (equal); Validation (equal); Visualization (lead); Writing – original draft (lead); Writing – review & editing (equal). Kai-xin Hu: Conceptualization (equal); Funding acquisition (equal); Investigation (equal); Methodology (equal); Project administration (equal); Resources (equal); Supervision (equal); Validation (equal); Writing – original draft (equal); Writing – review & editing (equal). Qi-Sheng Chen: Supervision (equal); Validation (equal).

DATA AVAILABILITY

The data that support the findings of this study are available within the article.

APPENDIX A: THE EXPRESSION AFTER ELIMINATING THE PRESSURE TERM

By utilizing Eqs. (17a)-17(c) to eliminate \hat{p} , we obtain the following equation:

$$\begin{split} m\{\hat{\tau}_{13} + rD\hat{\tau}_{13} + im\hat{\tau}_{23} + i\alpha r\hat{\tau}_{33}\} &= S^{\frac{3}{2}}i\alpha mr\bar{w}\hat{w} \\ &- S^{\frac{1}{2}}mrD\bar{w}\hat{u} - \alpha r\{2\hat{\tau}_{12} + rD\hat{\tau}_{12} + im\hat{\tau}_{22} + i\alpha r\hat{\tau}_{23}\} \\ &+ S^{\frac{1}{2}}i\alpha^{2}r^{2}\bar{w}\hat{v} = \sigma S^{\frac{1}{2}}r(m\hat{w} - \alpha r\hat{v}), \end{split}$$
(A1)

$$\begin{aligned} \alpha \Big\{ r D \hat{\tau}_{13} + r^2 D^2 \hat{\tau}_{13} - \hat{\tau}_{13} - im \hat{\tau}_{23} + imr D \hat{\tau}_{23} + i\alpha r^2 D \hat{\tau}_{33} \Big\} \\ &- S^{\frac{1}{2}} i\alpha^2 r^2 (D\bar{w}\hat{w} + \bar{w}D\hat{w}) - S^{\frac{1}{2}} \alpha r^2 (D^2 \bar{w}\hat{u} + D\bar{w}D\hat{u}) \\ &+ mr^2 \Big\{ 3D \hat{\tau}_{12} + r D^2 \hat{\tau}_{12} + im D \hat{\tau}_{22} + i\alpha \hat{\tau}_{23} + i\alpha r D \hat{\tau}_{23} \Big\} \\ &- S^{\frac{1}{2}} i\alpha mr^2 (\bar{w}\hat{v} + r D\bar{w}\hat{v} + r \bar{w}D\hat{v}) - S^{\frac{1}{2}} k^2 \alpha r^2 \bar{w}\hat{u} \\ &- ik^2 r \{ \hat{\tau}_{11} + r D \hat{\tau}_{11} + im \hat{\tau}_{12} + i\alpha r \hat{\tau}_{13} - \hat{\tau}_{22} \} \\ &= \sigma S^{\frac{1}{2}} r^2 (m \hat{v} + mr D \hat{v} + \alpha D \hat{w} - ik^2 \hat{u}). \end{aligned}$$
(A2)

Using Eqs. (17b) and (17c) to obtain \hat{p} and subsequently substituting it in Eq. (24), we can determine the boundary condition for the normal stress as follows:

$$mr\{2\hat{\tau}_{12} + rD\hat{\tau}_{12} + im\hat{\tau}_{22} + i\alpha r\hat{\tau}_{23}\} - S^{\frac{1}{2}}i\alpha mr^{2}\bar{w}\hat{v} + \alpha\{\hat{\tau}_{13} + rD\hat{\tau}_{13} + im\hat{\tau}_{23} + i\alpha r\hat{\tau}_{33}\} - S^{\frac{1}{2}}\alpha rD\bar{w}\hat{u} - S^{\frac{1}{2}}i\alpha^{2}r\bar{w}\hat{w} - ik^{2}r(\hat{\tau}_{11} - 2i\alpha\bar{\tau}_{13}\hat{R}) + S^{\frac{1}{2}}ik^{2}r(1 - \alpha^{2} - m^{2})\hat{R} + S^{-\frac{1}{2}}R_{B}ik^{2}r(D\bar{T}\hat{R} + \hat{T}) = \sigma S^{\frac{1}{2}}r(mr\hat{v} + \alpha\hat{w}).$$
(A3)

Here, $k^2 = \alpha^2 + m^2$.

APPENDIX B: THE SURFACE INTEGRAL EXPRESSION FOR THE PRODUCT OF PERTURBATIONS

For the perturbations A and B on the liquid surface,

$$\mathbf{A} = |\mathbf{A}| \sin\left(\boldsymbol{\Phi} + \boldsymbol{\varphi}_1\right),\tag{B1}$$

$$\mathbf{B} = |\mathbf{B}|\sin\left(\boldsymbol{\Phi} + \boldsymbol{\varphi}_2\right),\tag{B2}$$

the surface integral of their product is

$$\int_{0}^{2\pi} (AB) dr^{2} = |AB| \int_{0}^{2\pi} \sin \left(\Phi + \varphi_{1} \right) \sin \left(\Phi + \varphi_{2} \right) d\Phi$$
$$= \pi |AB| \cos(\varphi_{1} - \varphi_{2}). \tag{B3}$$

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