Vapor flows in the continuum limit in the presence of a small amount of noncondensable gas

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(Received 11 May 2004; accepted 30 July 2004; published online 6 October 2004)

Steady flows of a vapor around its condensed phase of arbitrary shape, on the surface of which evaporation and condensation of the vapor may take place, are considered in the presence of a small amount of a noncondensable gas. By a systematic asymptotic analysis of the Boltzmann system, the present authors have derived the fluid-dynamic system describing such flows in the continuum limit in the case where the amount of the noncondensable gas is infinitesimal compared with that of the vapor [K. Aoki, S. Takata, and S. Taguchi, Eur. J. Mech. B/Fluids **22**, 51 (2003)]. In the present study, the fluid-dynamic system is applied to some physical problems, and it is demonstrated with concrete examples that such a tiny amount of the noncondensable gas causes dramatic changes of the overall vapor flows. © 2004 American Institute of Physics. [DOI: 10.1063/1.1795271]

I. INTRODUCTION

In flows of a vapor with evaporation or condensation taking place on the boundary, the vapor is not in a local equilibrium state near the boundary even in the continuum limit, i.e., the limit in which the Knudsen number, defined by the ratio of the mean free path of the vapor molecules to the characteristic length of the system, goes to zero. Therefore, an approach based on kinetic theory is required to describe such flows. In fact, this type of flow has been one of the important subjects in kinetic theory of gases for more than three decades.^{1–6}

Concerning the continuum limit (and its neighborhood), the fluid-dynamic systems that describe steady flows of the vapor have been established by means of systematic asymptotic analyses of the Boltzmann system.^{5,7–10} Some of these fluid-dynamic systems have been extended to the vapor flows in the presence of another noncondensable gas that neither evaporates nor condenses on the boundary.^{11,12} However, some interesting and important phenomena peculiar to the continuum limit for the mixture of the vapor and the noncondensable gas were clarified only recently.^{13–18}

The phenomenon that is investigated in Ref. 18 is the effect of a tiny amount of the noncondensable gas on the overall vapor flows. For the sake of simplicity we restrict ourselves to the case of a closed domain, and we suppose that steady evaporation and condensation are taking place on the boundary. Let n_r be the reference molecular number density of the vapor, n_{av}^B the average molecular number density of the noncondensable gas over the domain, and Kn the Knudsen number defined by the mean free path of the vapor molecules in a reference equilibrium state divided by the characteristic length of the domain. We assume that the Knudsen number Kn is small and that the average concentration of the noncondensable gas is as small as the Knudsen number. The latter condition is equivalent to $n_{av}^B/n_r = O(Kn)$.

The fluid-dynamic equations, which are the set of compressible Euler equations, and their boundary conditions for the continuum limit under this condition were derived in Ref. 18 with the help of a systematic asymptotic analysis of the Boltzmann system for small Kn and a complementary numerical analysis performed in Refs. 19 and 20. In this limit, the average concentration becomes infinitesimal $(n_{av}^B/n_r \rightarrow 0)$. However, it was shown that the noncondensable gas of an infinitesimal average concentration accumulates in the Knudsen layer with an infinitesimal thickness on the condensing boundary and has a significant effect on the overall vapor flow through the fluid-dynamic boundary condition on the condensing boundary. An example that demonstrates a dramatic change of the streamlines of the vapor flow caused by the presence of the noncondensable gas with an infinitesimal average concentration is also given in Ref. 18.

High-speed flows of a single-component vapor that undergo strong evaporation and condensation play an important role in various applications such as heat pipes and laser ablation. In practical situations, however, the vapor flows may contain tiny impurities. The fluid-dynamic system established in Ref. 18 reveals that such impurities have a crucial effect on the vapor flows. Moreover, it enables quantitative assessment of the effect of the impurities. Therefore, the results given in Ref. 18 are not only of theoretical interest but also of practical importance.

In the present paper, we will give some more applications of the fluid-dynamic system derived in Ref. 18 and demonstrate the effect of the trace of the noncondensable gas on the global vapor flow. We first summarize the fluiddynamic system (Sec. II) and then investigate three concrete problems numerically (Sec. III).

II. BASIC EQUATIONS

In this section, we first describe the physical situation that we consider and then summarize the fluid-dynamic equations and their boundary conditions obtained in Ref. 18.

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A. Physical situation

Let us consider a vapor and its condensed phase of arbitrarily smooth shape, on the surface of which evaporation or condensation of the vapor is taking place. For the sake of simplicity, we restrict ourselves to the case of a closed domain (see Ref. 18 for the case of an infinite domain). A small amount of the noncondensable gas is contained in the system. In the following, we assign the label A to the vapor (it will also be called A component) and B to the noncondensable gas (it will also be called B component).

We first introduce the reference quantities: *L* is the reference length of the system, n_r is the reference number density of the vapor molecules, T_r is the reference temperature, p_r is the reference pressure of the vapor defined by $p_r = kn_rT_r$ (*k* is the Boltzmann constant), m^A is the mass of a molecule of the vapor, l_r is the mean free path of the vapor molecules in the equilibrium state at rest with temperature T_r and number density n_r , and $Kn = l_r/L$ is the Knudsen number with respect to the vapor. Further, we denote by n_{av}^B the average number density of the noncondensable gas contained in the system.

We consider steady flows of the vapor in the following situation: (i) The Knudsen number Kn is small. (ii) The amount of the noncondensable gas contained in the system is small; to be more specific, the amount is such that the average concentration of the noncondensable gas is of the order of the Knudsen number Kn. The latter condition is equivalent to setting

$$\frac{n_{av}^B}{n_r} = O(\mathrm{Kn}). \tag{1}$$

We will focus on the continuum limit where $Kn \rightarrow 0$, or equivalently, the zeroth-order quantities in Kn. It should be noted that in the situation of Eq. (1), the average concentration of the noncondensable gas becomes infinitesimal in the continuum limit. Nevertheless, as we will see, its trace has an important effect on the overall flow of the vapor.

B. Fluid-dynamic equations

In Ref. 18, the asymptotic behavior of the vapor and the noncondensable gas was investigated in the situation described in Sec. II A. More specifically, a systematic asymptotic analysis of the Boltzmann equation for small Knudsen numbers has been carried out under the assumption that the interaction of the vapor molecules with the boundary is described by the conventional condition for evaporation and condensation (the so-called complete condensation condition), and that of the noncondensable-gas molecules with the boundary by the diffuse reflection condition. The result is summarized here and in Sec. II C.

Let X_i denote the space rectangular coordinate system, n^{α} , ρ^{α} , and p^{α} the number density, the mass density, and the pressure of the α component ($\alpha = A, B$), respectively, and v_i and T the flow velocity and the temperature of the vapor,

respectively. We introduce their dimensionless counterparts by the following relations:

$$\begin{aligned} x_i &= X_i/L, \quad \hat{n}^{\alpha} = n^{\alpha}/n_r, \\ \hat{\rho}^{\alpha} &= \rho^{\alpha}/m^A n_r, \quad \hat{p}^{\alpha} = p^{\alpha}/p_r, \\ \hat{v}_i &= v_i/(2kT_r/m^A)^{1/2}, \quad \hat{T} = T/T_r. \end{aligned}$$
(2)

According to Ref. 18, the macroscopic quantities h ($h = \hat{n}^{\alpha}, \hat{\rho}^{\alpha}, \hat{p}^{\alpha}, \hat{v}_{i}$, or \hat{T}) are expressed in the following form:

$$h = h_H + h_K,\tag{3}$$

where h_H is the moderately varying overall solution whose length scale of variation is the reference length of the system [i.e., $\partial h_H / \partial x_i = O(1)$] and h_K is a correction term to h_H in a thin layer (the Knudsen layer) with thickness of the order of the mean free path (or of the order of Kn in the dimensionless x_i space) adjacent to the boundary. The h_H is the socalled Hilbert solution, and we call h_K the Knudsen-layer correction. We consider the situation in which there is no closed streamlines of the vapor flow or in which closed streamlines of the vapor flow, if any, do not carry any noncondensable gas. Then, as discussed in Ref. 18, we can assume, consistently with Eq. (1), that the noncondensable gas is absent in the overall flow field, that is,

$$\hat{n}_{H}^{B} = \hat{\rho}_{H}^{B} = \hat{p}_{H}^{B} = 0, \tag{4}$$

and that the noncondensable gas is also absent in the Knudsen layer on the evaporating boundary (this fact can be shown rigorously for the Maxwell molecules and for most of the model equations¹⁸). Consequently, the noncondensable gas can be present only in the Knudsen layer on the condensing boundary.

The macroscopic quantities h_H in the continuum limit $\text{Kn} \rightarrow 0$ (or, equivalently, the zeroth-order terms of their Kn-expansions) are described by the Euler set of equations for an ideal gas, which is written in the following dimensionless form:¹⁸

$$\frac{\partial \hat{\rho}^A \hat{v}_j}{\partial x_j} = 0, \tag{5a}$$

$$\hat{\rho}^{A}\hat{v}_{j}\frac{\partial\hat{v}_{i}}{\partial x_{j}} + \frac{1}{2}\frac{\partial\hat{p}^{A}}{\partial x_{i}} = 0,$$
(5b)

$$\hat{v}_j \frac{\partial}{\partial x_j} \left(\frac{5}{2} \hat{T} + \hat{v}_k^2 \right) = 0, \qquad (5c)$$

$$\hat{p}^A = \hat{\rho}^A \hat{T},\tag{5d}$$

where $\hat{\rho}^A = \hat{n}^A$. Here and in what follows, we omit the subscript *H* for the Hilbert solution because no confusion is expected.

C. Boundary conditions

The Knudsen-layer correction h_K [see Eq. (3)] is obtained by solving a half-space boundary-value problem of

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the Boltzmann equation. More specifically, the Knudsenlayer correction for the evaporating surface corresponds to the solution for the steady flow of a pure vapor evaporating from a plane condensed phase, $^{21-24}$ and that for the condensing surface corresponds to the solution for the steady flow of a vapor condensing onto a plane condensed phase in the presence of the noncondensable gas. 19,20,25,26 The boundary conditions for the Euler equations (5a)–(5d) are obtained together with the solution for the Knudsen-layer corrections. We will summarize, in the following, the boundary conditions thus obtained (see Ref. 18 for the details of the derivation).

Let $Lx_{wi}(\chi_1, \chi_2)$ denote the boundary, where χ_1 and χ_2 are the (dimensionless) coordinates on the boundary orthogonal to each other, $T_r \hat{T}_w$ the temperature of the boundary, $(2kT_r/m^A)^{1/2} \hat{v}_{wi}$ its velocity, and $p_r \hat{p}_w^A$ the saturation pressure of the vapor at temperature $T_r \hat{T}_w$. Further, we denote by n_i the unit normal vector to the boundary pointing to the gas region and by $t_i^{(1)}$ and $t_i^{(2)}$ the unit tangential vectors to the boundary in the direction of increasing χ_1 and χ_2 , which are taken in such a way that $t_i^{(1)}$, $t_i^{(2)}$, and n_i form a right-hand system. Since we consider steady flows, $\hat{v}_{wi}n_i=0$ holds.

First, we introduce the following local Mach numbers based on the normal and tangential flow velocities on the boundary relative to the velocity of the boundary:

$$M_n = \sqrt{6/5(\hat{T})_b^{-1/2}} |(\hat{v}_j)_b n_j|, \qquad (6a)$$

$$M_{t} = \sqrt{6/5} (\hat{T})_{b}^{-1/2} |(\hat{v}_{i})_{b} - \hat{v}_{wi} - (\hat{v}_{j})_{b} n_{j} n_{i}|,$$
(6b)

where $(\cdot)_b$ indicates that the quantity in the parentheses is evaluated on the boundary. Then, the boundary conditions for the Euler equations in the continuum limit are expressed as follows.

1. Boundary condition on the evaporating surface

The boundary condition for the Euler set of equations on the evaporating surface $[(\hat{v}_j)_b n_j > 0]$ is given in the following form:

$$M_n \le 1, \quad M_t = 0, \tag{7a}$$

$$\frac{(\hat{p}^{A})_{b}}{\hat{p}_{w}^{A}} = h_{1}(M_{n}), \quad \frac{(\hat{T})_{b}}{\hat{T}_{w}} = h_{2}(M_{n}).$$
(7b)

The functions $h_1(M_n)$ and $h_2(M_n)$ in Eq. (7b) have been constructed numerically by the use of the Bhatnagar–Gross– Krook (BGK) model^{27–29} of the Boltzmann equation in Ref. 23. The analytical form of these functions for $M_n \ll 1$ was obtained in Ref. 22 (see also Refs. 5, 24, and 30).

2. Boundary condition on the condensing surface

The boundary condition for the Euler set of equations on the condensing surface $[(\hat{v}_j)_b n_j < 0]$ is summarized as follows:

$$\frac{(\hat{p}^A)_b}{\hat{p}^A_w} = \mathcal{F}_s \left(M_n, M_l, \frac{(\hat{T})_b}{\hat{T}_w}, \frac{l_r}{(\hat{n}^A)_b l_b} \widetilde{\Gamma} \right) \quad (\text{when } M_n < 1), \quad (8a)$$

$$\frac{(\hat{p}^A)_b}{\hat{p}_w^A} \ge \mathcal{F}_b \left(1, M_t, \frac{(\hat{T})_b}{\hat{T}_w}, \frac{l_r}{(\hat{n}^A)_b l_b} \widetilde{\Gamma} \right) \quad (\text{when } M_n = 1), \qquad (8b)$$

$$\frac{(\hat{p}^A)_b}{\hat{p}^A_w} > \mathcal{F}_b\left(M_n, M_t, \frac{(\hat{T})_b}{\hat{T}_w}, \frac{l_r}{(\hat{n}^A)_b l_b}\tilde{\Gamma}\right) \quad (\text{when } M_n > 1), \quad (8c)$$

$$\chi_{1,1} \frac{\partial}{\partial \chi_1} (\hat{\mathcal{N}}_i^{\mathcal{B}} t_i^{(1)}) + \chi_{2,2} \frac{\partial}{\partial \chi_2} (\hat{\mathcal{N}}_i^{\mathcal{B}} t_i^{(2)}) + g_2 \hat{\mathcal{N}}_i^{\mathcal{B}} t_i^{(1)} - g_1 \hat{\mathcal{N}}_i^{\mathcal{B}} t_i^{(2)} = 0,$$
(9)

$$\begin{split} \hat{\mathcal{N}}_{i}^{\mathcal{B}} &= \hat{v}_{wi} \widetilde{\Gamma} + \frac{l_{b}}{l_{r}} (\hat{n}^{A})_{b} (\hat{T})_{b}^{1/2} \\ &\times G_{s} \Biggl(M_{n}, M_{t}, \frac{(\hat{T})_{b}}{\hat{T}_{w}}, \frac{l_{r}}{(\hat{n}^{A})_{b} l_{b}} \widetilde{\Gamma} \Biggr) a_{i} \quad \text{(when } M_{n} < 1\text{)}, \end{split}$$

$$(10a)$$

$$\begin{split} \hat{\mathcal{N}}_{i}^{\mathcal{B}} &= \hat{v}_{wi} \widetilde{\Gamma} + \frac{l_{b}}{l_{r}} (\hat{n}^{A})_{b} (\hat{T})_{b}^{1/2} \\ &\times G_{b} \Biggl(M_{n}, M_{l}, \frac{(\hat{T})_{b}}{\hat{T}_{w}}, \frac{(\hat{p}^{A})_{b}}{\hat{p}_{w}^{A}}, \frac{l_{r}}{(\hat{n}^{A})_{b} l_{b}} \widetilde{\Gamma} \Biggr) a_{i} \\ & (\text{when } M_{n} \ge 1), \end{split}$$
(10b)

where l_b is the mean free path of the vapor molecules in the equilibrium state at rest with number density $n_r(\hat{n}^A)_b$ and temperature $T_r(\hat{T})_b$ and is therefore a function of χ_1 and χ_2 ; $\chi_{1,1}$ and $\chi_{2,2}$ are defined by

$$\chi_{1,1} = \left(\frac{\partial \chi_1}{\partial x_j}\right)_b t_j^{(1)}, \quad \chi_{2,2} = \left(\frac{\partial \chi_2}{\partial x_j}\right)_b t_j^{(2)}, \tag{11}$$

and g_1 and g_2 are, respectively, the geodesic curvatures³¹ (in the dimensionless x_i space) of the χ_1 and χ_2 coordinate lines on the boundary; and a_i , which is defined by

$$a_{i} = \frac{(\hat{v}_{i})_{b} - \hat{v}_{wi} - (\hat{v}_{j})_{b} n_{j} n_{i}}{|(\hat{v}_{i})_{b} - \hat{v}_{wi} - (\hat{v}_{j})_{b} n_{i} n_{i}|},$$
(12)

is the tangential unit vector in the direction of the tangential component of the flow velocity of the vapor relative to the boundary. When the tangential component is zero $[(\hat{v}_i)_b - \hat{v}_{wi} - (\hat{v}_j)_b n_j n_i = 0 \text{ or } M_i = 0]$, we can define a_i as an arbitrary unit tangential vector because G_s and G_b vanish [see Eq. (13) below]. Here, we note the following: In Ref. 18, $|M_n|$ instead of M_n appears in the equations corresponding to Eqs. (8) and (10) because M_n is defined by $M_n = \sqrt{6/5}(\hat{T})_b^{-1/2}(\hat{v}_j)_b n_j$ [cf. Eq. (6a)]; in the same reference, $\mathcal{F}_s(1_-, \cdots)$, which is the same as $\mathcal{F}_b(1, \cdots)$, is used in the equation corresponding to

Eq. (8b); and $\hat{\mathcal{N}}_{i0}$ in Ref. 18 corresponds to $\hat{\mathcal{N}}_{i}^{\mathcal{B}}$ here. The $\tilde{\Gamma}$ and thus $\hat{\mathcal{N}}_{i}^{\mathcal{B}}$ are undetermined functions of χ_{1} and χ_{2} , determined together with the solution of the Euler set of equations. The physical meaning of these functions will be explained below. The functions \mathcal{F}_s and \mathcal{F}_b of four independent variables occurring in Eq. (8) have been constructed numerically in Refs. 19 and 20 (see also Refs. 25 and 26), on the basis of the model Boltzmann equation proposed by Garzó et al.³² [the Garzó–Santos–Brey (GSB) model]. It should be mentioned that these numerical functions are restricted to the case where the molecules of the noncondensable gas are mechanically identical to those of the vapor. The numerical data for these functions are given in Refs. 19 and 20: see Eq. (25) in Ref. 19 and related figures and tables for \mathcal{F}_s ; and see Eq. (15a) in Ref. 20 and related figures and tables for \mathcal{F}_b . The numerical data for the functions G_s in Eq. (10a) and G_b in Eq. (10b), based on the GSB model and restricted to the case where the molecules of the noncondensable gas and those of the vapor are mechanically the same, are also given in Refs. 19 and 20. More specifically, G_s corresponds to $\hat{\mathcal{N}}_f = \Gamma \mathcal{G}$ in Ref. 19 [see Eq. (32) there; Γ corresponds to $l_r \tilde{\Gamma} / (\hat{n}^A)_b l_b$ here], and \mathcal{G} is tabulated there (see Fig. 9 and Table III in Ref. 19 and also Tables IX–XII in Ref. 33); and G_b corresponds to $\hat{\mathcal{N}}_f$ in Ref. 20 (see Fig. 8 and Table III in Ref. 20 and also Tables IX-XII in Ref. 34). We note that the functions G_s and G_b are nonnegative and have the property that

$$G_s = G_b = 0$$
 only when $M_t = 0$ or $\tilde{\Gamma} = 0$. (13)

When the problem is spatially two dimensional, we may assume that the physical quantities are independent of χ_2 . For simplicity, let us denote χ_1 by χ and $t_i^{(1)}$ by t_i . Then, because $g_1=g_2=0$ in this case, it follows from Eq. (9) that

$$\hat{\mathcal{N}}_i^B t_i = \text{const.} \tag{14}$$

The functions $\tilde{\Gamma}$ and $\hat{\mathcal{N}}_{i}^{B}$ of χ_{1} and χ_{2} are auxiliary functions in the boundary condition, and their physical meaning is not necessary to obtain the solution of the Euler set of equations with the boundary conditions (7)–(10). However, these functions themselves have clear physical meaning related to the behavior of the noncondensable gas in the Knudsen layer. Let $n_r \hat{n}_K^B$ and $n_r (2kT_r/m^A)^{1/2} \Phi_{iK}^B$ denote, respectively, the number density and the (local) particle flux of the noncondensable gas molecules in the Knudsen layer, and let η be the stretched coordinate normal to the boundary defined by $x_i = x_{wi} + \epsilon \eta n_i$, where $\epsilon = (\sqrt{\pi}/2) \text{Kn}$ ($\eta = 0$ corresponds to the boundary). Then, \hat{n}_K^B and Φ_{iK}^B are functions of χ_1 , χ_2 , and η , and $\tilde{\Gamma}$ and $\hat{\mathcal{N}}_i^B$ are defined as¹⁸

$$\widetilde{\Gamma} = \int_0^\infty \hat{n}_K^B d\eta, \qquad (15a)$$

$$\hat{\mathcal{N}}_{i}^{\mathcal{B}} = \int_{0}^{\infty} \Phi_{ik}^{\mathcal{B}} d\,\eta.$$
(15b)

Thus, $(\sqrt{\pi}/2)n_r l_r \tilde{\Gamma}$ and $(\sqrt{\pi}/2)n_r (2kT_r/m^A)^{1/2} l_r \hat{\mathcal{N}}_i^{\mathcal{B}}$ are, respectively, the total number and the total particle flux of the



FIG. 1. Two-surface problem of evaporation and condensation in the presence of a noncondensable gas.

noncondensable-gas molecules contained in the Knudsen layer per unit area on the boundary.

D. Summary

To summarize, the vapor flows in the continuum limit in the situation explained in Sec. II A are described by the Euler set of equations, Eqs. (5a)–(5d), with the boundary condition, Eqs. (7a) and (7b), when evaporation of the vapor is taking place and with the boundary condition, Eqs. (8a)–(8c), (9), (10a), and (10b), when condensation is taking place. In this physical situation, the noncondensable gas of an infinitesimal average concentration accumulates in the Knudsen layer with an infinitesimal thickness on the condensing boundary. As a result, its local concentration on the boundary rises to the order of unity. In this way, the noncondensable gas has a finite effect on the vapor flow through the boundary condition on the condensing boundary (or more precisely, through $\tilde{\Gamma}$ in the boundary condition).

III. APPLICATIONS

In this section, we show some applications of the Euler system summarized in the preceding section. To be consistent with the numerical data for the functions \mathcal{F}_s , \mathcal{F}_b , G_s , and G_b given in Refs. 19 and 20 [see Sec. II C for the correspondence between (G_s, G_b) and the symbols in Refs. 19 and 20], we assume that the molecules of the noncondensable gas are mechanically identical to those of the vapor and that the behavior of the mixture is described by the GSB model. Then, we make use of these data as well as those for the functions h_1 and h_2 based on the BGK model given in Ref. 23.

A. Two-surface problem of evaporation and condensation

The first problem is the two-surface problem of evaporation and condensation. We consider a vapor (A component) in the gap $0 \le X_1 \le L$ between two parallel plane condensed phases (see Fig. 1). Let us suppose that the surface at X_1 =0 is kept at temperature T_I and is set at rest, whereas that at $X_1=L$ is kept at temperature T_{II} and may be moving in its surface in the X_2 direction with a constant speed U_{II} . We denote by p_I and p_{II} the saturation pressure of the vapor at

TABLE I. The quantities $\hat{\rho}^A$, \hat{v}_1 , \hat{T} , \hat{p}^A , M_n , and $\hat{\mathcal{M}}$ for different values of the parameters p_{II}/p_I , \hat{U}_{II} and Δ ($T_{II}/T_I=1$). Here, the molecules of the noncondensable gas are assumed to be mechanically identical with those of the vapor.

		$p_{\rm II}/p_{\rm I}=1.2$						$p_{\rm II}/p_{\rm I} = 1.5$					
\hat{U}_{II}	Δ	$\hat{ ho}^A$	$-\hat{v}_1$	\hat{T}	\hat{p}^A	M_n	$\hat{\mathcal{M}}$	$\hat{ ho}^A$	$-\hat{v}_1$	\hat{T}	\hat{p}^A	M_n	$\hat{\mathcal{M}}$
0	0	1.118	0.0424	0.981	1.097	0.0469	0.0474	1.284	0.0930	0.958	1.231	0.1041	0.1194
0	0.5	1.128	0.0370	0.983	1.109	0.0409	0.0417	1.312	0.0801	0.964	1.265	0.0894	0.1051
0	1	1.136	0.0329	0.985	1.119	0.0363	0.0374	1.333	0.0704	0.969	1.291	0.0783	0.0938
0	2	1.147	0.0269	0.988	1.134	0.0297	0.0309	1.364	0.0567	0.975	1.329	0.0629	0.0773
0.5	0	1.120	0.0414	0.981	1.099	0.0458	0.0464	1.288	0.0912	0.959	1.235	0.1020	0.1175
0.5	0.5	1.129	0.0363	0.984	1.111	0.0401	0.0410	1.315	0.0787	0.965	1.269	0.0878	0.1035
0.5	1	1.137	0.0324	0.986	1.120	0.0357	0.0368	1.336	0.0693	0.969	1.294	0.0771	0.0926
0.5	2	1.148	0.0265	0.988	1.135	0.0292	0.0304	1.365	0.0560	0.975	1.331	0.0621	0.0764
1	0	1.124	0.0388	0.983	1.105	0.0429	0.0436	1.299	0.0861	0.961	1.249	0.0962	0.1118
1	0.5	1.133	0.0343	0.985	1.116	0.0379	0.0389	1.323	0.0749	0.967	1.279	0.0834	0.0991
1	1	1.140	0.0307	0.986	1.124	0.0339	0.0350	1.342	0.0663	0.970	1.303	0.0737	0.0890
1	2	1.150	0.0254	0.989	1.137	0.0280	0.0292	1.370	0.0540	0.976	1.337	0.0598	0.0740
2	0	1.139	0.0312	0.986	1.123	0.0344	0.0355	1.332	0.0708	0.968	1.290	0.0788	0.0943
2	0.5	1.145	0.0282	0.987	1.131	0.0310	0.0323	1.350	0.0629	0.972	1.312	0.0699	0.0849
2	1	1.150	0.0257	0.989	1.137	0.0283	0.0296	1.364	0.0566	0.975	1.330	0.0628	0.0772
2	2	1.157	0.0218	0.990	1.146	0.0240	0.0252	1.385	0.0473	0.979	1.356	0.0524	0.0655

temperature $T_{\rm I}$ and that at temperature $T_{\rm II}$, respectively. A noncondensable gas (*B* component) is contained in the gap, whose average molecular number density over the gap is denoted by n_{av}^B . Here, we take *L* as the reference length and $T_{\rm I}$, $p_{\rm I}$, and $n_{\rm I}=p_{\rm I}/kT_{\rm I}$ as the reference quantities, i.e., $T_r=T_{\rm I}$, $p_r=p_{\rm I}$, and $n_r=n_{\rm I}$. Then, the problem is characterized by the following parameters:

$$\frac{T_{\rm II}}{T_{\rm I}}, \quad \frac{p_{\rm II}}{p_{\rm I}}, \quad \hat{U}_{\rm II} = \frac{U_{\rm II}}{(2kT_{\rm I}/m^A)^{1/2}}, \quad \mathrm{Kn} = \frac{l_r}{L}, \quad \frac{n_{av}^B}{n_{\rm I}}, \quad (16)$$

where l_r is the mean free path of the vapor molecules in the equilibrium state at rest with temperature $T_{\rm I}$ and pressure $p_{\rm I}$ (or molecular number density $n_{\rm I}$). We assume that $T_{\rm I} \leq T_{\rm II}$ and $p_{\rm I} < p_{\rm II}$, so that evaporation takes place on the surface at

TABLE II. The quantities \hat{p}^A , \hat{v}_1 , \hat{T} , \hat{p}^A , M_n , and $\hat{\mathcal{M}}$ for different values of the parameters p_{II}/p_1 , \hat{U}_{II} and $\Delta (T_{\text{II}}/T_1=1.1)$. See the caption of Table I.

		$p_{\rm II}/p_{\rm I}$ =2						$p_{\rm II}/p_{\rm I}$ =5					
\hat{U}_{II}	Δ	$\hat{ ho}^A$	$-\hat{v}_1$	\hat{T}	\hat{p}^A	M_n	$\hat{\mathcal{M}}$	$\hat{ ho}^A$	$-\hat{v}_1$	\hat{T}	\hat{p}^A	M_n	$\hat{\mathcal{M}}$
0	0	1.402	0.1643	1.023	1.434	0.1779	0.2303	2.600	0.3602	0.928	2.412	0.4096	0.9365
0	0.5	1.457	0.1396	1.034	1.507	0.1504	0.2033	2.923	0.2824	0.966	2.823	0.3147	0.8255
0	1	1.499	0.1216	1.043	1.563	0.1305	0.1823	3.143	0.2347	0.989	3.109	0.2585	0.7377
0	2	1.558	0.0969	1.054	1.643	0.1034	0.1510	3.434	0.1775	1.016	3.490	0.1929	0.6095
0.5	0	1.408	0.1615	1.024	1.442	0.1749	0.2274	2.616	0.3559	0.930	2.433	0.4043	0.9310
0.5	0.5	1.462	0.1376	1.035	1.513	0.1482	0.2011	2.936	0.2794	0.967	2.840	0.3112	0.8203
0.5	1	1.502	0.1201	1.044	1.568	0.1287	0.1803	3.154	0.2325	0.990	3.122	0.2560	0.7333
0.5	2	1.561	0.0959	1.055	1.646	0.1023	0.1497	3.441	0.1761	1.017	3.499	0.1913	0.6060
1	0	1.425	0.1540	1.027	1.464	0.1664	0.2195	2.663	0.3440	0.936	2.492	0.3895	0.9161
1	0.5	1.475	0.1319	1.038	1.530	0.1419	0.1946	2.973	0.2711	0.971	2.888	0.3013	0.8060
1	1	1.513	0.1156	1.046	1.582	0.1239	0.1749	3.184	0.2263	0.993	3.162	0.2488	0.7205
1	2	1.568	0.0930	1.056	1.656	0.0991	0.1458	3.462	0.1723	1.019	3.527	0.1870	0.5965
2	0	1.478	0.1304	1.039	1.535	0.1401	0.1927	2.822	0.3054	0.955	2.694	0.3424	0.8618
2	0.5	1.517	0.1139	1.046	1.588	0.1219	0.1728	3.098	0.2441	0.984	3.050	0.2695	0.7562
2	1	1.547	0.1013	1.052	1.628	0.1082	0.1567	3.285	0.2060	1.003	3.294	0.2253	0.6767
2	2	1.592	0.0833	1.061	1.689	0.0886	0.1327	3.532	0.1593	1.025	3.620	0.1724	0.5626

TABLE III. The quantities \hat{p}^A , \hat{v}_1 , \hat{T} , \hat{p}^A , M_n , and $\hat{\mathcal{M}}$ for different values of the parameters p_{II}/p_I , \hat{U}_{II} and $\Delta (T_{II}/T_I=1.2)$. See the caption of Table I.

		$p_{\rm II}/p_{\rm I}=5$						$p_{\rm II}/p_{\rm I} = 10$						
\hat{U}_{II}	Δ	$\hat{ ho}^A$	$-\hat{v}_1$	\hat{T}	\hat{p}^A	M_n	$\hat{\mathcal{M}}$	$\hat{ ho}^A$	$-\hat{v}_1$	\hat{T}	\hat{p}^A	M_n	$\hat{\mathcal{M}}$	
0	0	2.381	0.3768	1.012	2.409	0.4103	0.8972	3.929	0.5136	0.940	3.694	0.5803	2.018	
0	0.5	2.665	0.2986	1.052	2.803	0.3190	0.7958	4.754	0.3779	1.011	4.808	0.4117	1.797	
0	1	2.862	0.2497	1.077	3.081	0.2636	0.7146	5.271	0.3063	1.048	5.523	0.3278	1.615	
0	2	3.125	0.1902	1.106	3.458	0.1980	0.5943	5.924	0.2264	1.088	6.447	0.2377	1.341	
0.5	0	2.395	0.3726	1.014	2.429	0.4053	0.8924	3.954	0.5090	0.942	3.727	0.5744	2.013	
0.5	0.5	2.677	0.2956	1.053	2.819	0.3155	0.7913	4.775	0.3748	1.013	4.837	0.4079	1.789	
0.5	1	2.872	0.2475	1.078	3.095	0.2611	0.7108	5.289	0.3040	1.049	5.548	0.3251	1.608	
0.5	2	3.132	0.1887	1.107	3.467	0.1965	0.5910	5.937	0.2249	1.089	6.465	0.2361	1.335	
1	0	2.437	0.3606	1.020	2.486	0.3910	0.8788	4.025	0.4962	0.949	3.822	0.5578	1.997	
1	0.5	2.710	0.2871	1.058	2.866	0.3058	0.7780	4.836	0.3659	1.018	4.921	0.3974	1.769	
1	1	2.899	0.2411	1.081	3.133	0.2540	0.6989	5.340	0.2973	1.052	5.620	0.3175	1.588	
1	2	3.151	0.1847	1.109	3.494	0.1921	0.5820	5.973	0.2208	1.091	6.517	0.2315	1.319	
2	0	2.578	0.3215	1.040	2.681	0.3453	0.8288	4.273	0.4534	0.972	4.153	0.5037	1.937	
2	0.5	2.822	0.2594	1.072	3.024	0.2745	0.7320	5.042	0.3368	1.032	5.206	0.3631	1.698	
2	1	2.990	0.2200	1.091	3.263	0.2307	0.6578	5.513	0.2754	1.064	5.863	0.2926	1.518	
2	2	3.215	0.1712	1.116	3.587	0.1775	0.5504	6.097	0.2069	1.098	6.694	0.2163	1.261	

 $X_1 = L$ (or $x_1 = 1$) and condensation at $X_1 = 0$ (or $x_1 = 0$); therefore $\hat{v}_1 < 0$. In view of Eq. (1), we let

$$\frac{n_{av}^B}{n_1} = \Delta \text{ Kn},\tag{17}$$

where Δ is a constant, and we specify Δ rather than $n_{av}^B/n_{\rm I}$. We investigate the steady flow of the vapor caused by evaporation and condensation on the condensed phases and the effect of the noncondensable gas on the vapor flow in the continuum limit with respect to the vapor (Kn \rightarrow 0). The present problem has been investigated in Ref. 13 in the case of $\hat{U}_{\rm II}$ =0. The following analysis and discussion are straightforward extensions of those given there.

Now, let us apply the Euler equations and their boundary conditions to the present problem. Since the problem is spatially one-dimensional, $\tilde{\Gamma}$ in Eqs. (8a)–(8c) is a constant, which is related to Δ by¹³

$$\Delta = \frac{\sqrt{\pi}}{2}\tilde{\Gamma}.$$
(18)

This relation is derived from the following fact: since there is no noncondensable gas except in the Knudsen layer at the condensing surface $(x_1=0)$, the quantity $(\sqrt{\pi}/2)n_I l_r \tilde{\Gamma}$, which is the total number of the noncondensable-gas molecules contained in the Knudsen layer per unit area of the condensing surface [see the paragraph containing Eq. (15)], is equal to $n_{av}^B L$, i.e., the total number of the noncondensable-gas molecules contained in the column of length *L* between two condensed phases, perpendicular to them, with a unit base area. In the present one-dimensional problem, $\hat{\mathcal{N}}_i^B$ = $(0, \hat{\mathcal{N}}_2^B, 0)$ is also constant, so that Eq. (9) or Eq. (14) is satisfied. It follows from Eqs. (5a)–(5d) that $\hat{\rho}^A(=\hat{n}^A)$, \hat{v}_i ($\hat{v}_3=0$), \hat{T} , and \hat{p}^A are all constant. The boundary condition on the evaporating surface (7) leads to the following condition at $x_1=1$:

$$M_n \le 1, \quad \hat{v}_2 = \hat{U}_{\mathrm{II}}, \tag{19a}$$

$$\hat{p}^{A} = \frac{p_{\mathrm{II}}}{p_{\mathrm{I}}} h_{1}(M_{n}), \quad \hat{T} = \frac{T_{\mathrm{II}}}{T_{\mathrm{I}}} h_{2}(M_{n}),$$
(19b)

where

$$M_n = \sqrt{6/5} \hat{T}^{-1/2} |\hat{v}_1|. \tag{20}$$

The notation $(\cdot)_b$ is omitted in Eqs. (19) and (20) as well as in Eqs. (21) and (22) below because \hat{p}^A , \hat{T} , etc., are constants. The boundary condition on the condensing surface (8)–(10) reduces to the following condition at $x_1=0$:

$$\hat{p}^{A} = \mathcal{F}_{s}\left(M_{n}, M_{t}, \hat{T}, \frac{l_{r}}{\hat{n}^{A}l_{b}}\tilde{\Gamma}\right) \quad (M_{n} < 1),$$
(21a)

$$\hat{p}^A \ge \mathcal{F}_b \left(1, M_l, \hat{T}, \frac{l_r}{\hat{n}^A l_b} \tilde{\Gamma} \right) \quad (M_n = 1),$$
(21b)

$$\hat{p}^{A} > \mathcal{F}_{b}\left(M_{n}, M_{t}, \hat{T}, \frac{l_{r}}{\hat{n}^{A}l_{b}}\tilde{\Gamma}\right) \quad (M_{n} > 1),$$
(21c)

where M_n is the same as Eq. (20), M_t is given by

$$M_t = \sqrt{6/5} \hat{T}^{-1/2} |\hat{v}_2|, \qquad (22)$$

and l_b is the mean free path of the vapor molecules in the equilibrium state at rest with temperature $T_1 \hat{T}$ and number density $n_1 \hat{n}^A$. The ratio l_r/l_b , appearing in the functions \mathcal{F}_s

and \mathcal{F}_b , depends on the molecular model: $l_r/l_b = \hat{n}^A / \hat{T}^{1/2}$ for the GSB model (for example, $l_r/l_b = \hat{n}^A$ for hard-sphere molecules).

From the first equation of Eq. (19a), the flow is either subsonic $(M_n < 1)$ or sonic $(M_n=1)$. First, we consider the subsonic case. If we eliminate \hat{p}^A , \hat{T} , \hat{v}_2 , and $\tilde{\Gamma}$ from Eqs. (18), (19a), (19b), (21a), and (22), we obtain the following equation for M_n :

$$\frac{p_{\rm II}}{p_{\rm I}} h_1(M_n) = \mathcal{F}_s \left(M_n, \left(\frac{6}{5}\right)^{1/2} \hat{U}_{\rm II} \left[\frac{T_{\rm II}}{T_{\rm I}} h_2(M_n) \right]^{-1/2}, \\ \frac{T_{\rm II}}{T_{\rm I}} h_2(M_n), \frac{2}{\sqrt{\pi}} \left[\frac{T_{\rm II}}{T_{\rm I}} h_2(M_n) \right]^{-1/2} \Delta \right).$$
(23)

The M_n is determined by solving Eq. (23) for a given set of the parameter $(T_{\rm II}/T_{\rm I}, p_{\rm II}/p_{\rm I}, \hat{U}_{\rm II}, \Delta)$. Then, \hat{p}^A and \hat{T} are obtained from Eq. (19b), \hat{v}_1 from Eq. (20), and $\hat{\rho}^A$ (= \hat{n}^A) from Eq. (5d). We show the constants $\hat{\rho}^A$, \hat{v}_1 , \hat{T} , \hat{p}^A , and M_n thus obtained for various values of the set $(T_{\rm II}/T_{\rm I}, p_{\rm II}/p_{\rm I}, \hat{U}_{\rm II}, \Delta)$ in Tables I–III. The results for the case $U_{II}=0$ are given in Table I of Ref. 13, where the parameter $n_{\rm II}/n_{\rm I}$ ($n_{\rm II}=p_{\rm II}/kT_{\rm II}$) is used rather than $p_{\rm II}/p_{\rm I}$. In the present study, we recomputed the case of $U_{II}=0$ using supplemented numerical data of \mathcal{F}_s and included the results in Tables I–III for the sake of completeness. Since the dependence of \mathcal{F}_s on M_t is weak, the solution does not depend much on \hat{U}_{II} . Its dependence on Δ is stronger. The dimensionless mass-flow rate $\hat{\mathcal{M}}(=\hat{\rho}^A | \hat{v}_1 |)$ from the evaporating to the condensing surface, which is related to the (dimensional) mass-flow rate $\mathcal{M}(=\rho^A |v_1|)$ per unit time and unit area (through a plane x_1 =const) as \mathcal{M} $=2p_{\rm I}(2kT_{\rm I}/m^A)^{-1/2}\hat{\mathcal{M}}$, is also shown in Tables I–III. With the increase of Δ (or the amount of the noncondensable gas in the Knudsen layer at $x_1=0$), the condensation becomes weaker, and thus $|\hat{v}_1|$ and $\hat{\mathcal{M}}$ decrease. On the other hand, $\hat{v}_2 = \hat{U}_{\text{II}}$ holds irrespective of Δ . Therefore, when \hat{U}_{II} is finite (e.g., \hat{U}_{II} =0.5, 1, 2 in Tables I–III), the flow becomes more tangential to the plane condensed phases for larger Δ . We will see the same effect in Figs. 5 and 6 in Sec. III B.

Next, we consider the case where the flow speed is sonic $(M_n=1)$. If we eliminate \hat{p}^A , \hat{T} , \hat{v}_2 , and $\tilde{\Gamma}$ from Eqs. (18), (19a), (19b), (21b), and (22), we obtain the following relation:

$$\frac{p_{\mathrm{II}}}{p_{\mathrm{I}}} \ge \mathcal{S}\left(\hat{U}_{\mathrm{II}}, \frac{T_{\mathrm{II}}}{T_{\mathrm{I}}}, \Delta\right),\tag{24a}$$

where

$$S\left(\hat{U}_{\mathrm{II}}, \frac{T_{\mathrm{II}}}{T_{\mathrm{I}}}, \Delta\right) = \frac{1}{h_{1}(1)} \mathcal{F}_{b}\left(1, \left(\frac{6}{5}\right)^{1/2} \hat{U}_{\mathrm{II}}\left[\frac{T_{\mathrm{II}}}{T_{\mathrm{I}}}h_{2}(1)\right]^{-1/2}, \\ \frac{T_{\mathrm{II}}}{T_{\mathrm{I}}}h_{2}(1), \frac{2}{\sqrt{\pi}}\left[\frac{T_{\mathrm{II}}}{T_{\mathrm{I}}}h_{2}(1)\right]^{-1/2}\Delta\right).$$
(24b)

This relation gives the condition for $T_{\rm II}/T_{\rm I}$, $p_{\rm II}/p_{\rm I}$, $\hat{U}_{\rm II}$, and Δ for which a sonic flow $(M_n=1)$ occurs. Some examples of the boundary $S(\hat{U}_{\rm II}, T_{\rm II}/T_{\rm I}, \Delta)$ of the existence range (24a)



FIG. 2. The boundary $S(\hat{U}_{\text{II}}, T_{\text{II}}/T_{1}, \Delta)$ of the existence range vs Δ for various values of \hat{U}_{II} . (a) $T_{\text{II}}/T_{\text{I}}=1$, (b) $T_{\text{II}}/T_{\text{I}}=2$. The dotted lines in the figures indicate the asymptotes ($\Delta = \delta_c$) of the curves. Here, the molecules of the noncondensable gas are assumed to be mechanically identical with those of the vapor.

are shown in Fig. 2, where $S(\hat{U}_{\text{II}}, T_{\text{II}}/T_{\text{I}}, \Delta)$ is shown as a function of Δ for $T_{\text{II}}/T_{\text{I}}=1$ and 2 and for various values of \hat{U}_{II} . The $S(\hat{U}_{\text{II}}, T_{\text{II}}/T_{\text{I}}, \Delta)$ is an increasing function of Δ and becomes infinitely large as Δ approaches a critical value δ_c that depends on \hat{U}_{II} and $T_{\text{II}}/T_{\text{I}}$. That is, $\Delta = \delta_c$ is the asymptote of the curve, and there is no sonic solution in the region $\Delta \ge \delta_c$. The formula of the critical value δ_c is readily obtained from the form of \mathcal{F}_b given in Ref. 20, i.e.,

$$\delta_{c} = \frac{\sqrt{\pi}}{2} \left[\frac{T_{\rm II}}{T_{\rm I}} h_{2}(1) \right]^{1/2} \\ \times \Gamma_{b} \left(1, \left(\frac{6}{5} \right)^{1/2} \hat{U}_{\rm II} \left[\frac{T_{\rm II}}{T_{\rm I}} h_{2}(1) \right]^{-1/2}, \frac{T_{\rm II}}{T_{\rm I}} h_{2}(1) \right), \quad (25)$$

where Γ_b is a function the numerical data of which are given



FIG. 3. The critical value δ_c vs \hat{U}_{II} for various T_{II}/T_I . The δ_c is taken as the abscissa for easy comparison with Fig. 2. See the caption of Fig. 2.



FIG. 4. Vapor flow between a plane and a sinusoidal condensed phase in the presence of a noncondensable gas.

in Ref. 20. In Fig. 3, δ_c versus \hat{U}_{II} is shown for various values of $T_{\text{II}}/T_{\text{I}}$; δ_c is taken as the abscissa for easy comparison with Fig. 2.

Here, we should recall that, because of Eq. (17), n_{av}^B/n_I vanishes in the continuum limit Kn \rightarrow 0. As we have seen, however, the solution in the continuum limit depends on Δ , which is a quantity specifying the vanishing rate of n_{av}^B/n_I and thus related to the amount of the noncondensable gas. This means that, in the continuum limit, the vapor flow is still affected by the noncondensable gas in spite of the fact that its average concentration is infinitesimal.

B. Vapor flows between a plane and a sinusoidal condensed phase

Next we consider a vapor (A component) and a noncondensable gas (B component) in the gap between a sinusoidal condensed phase located at $X_1 = A \cos(\pi X_2/L)$ and a plane condensed phase located at $X_1 = L$ (see Fig. 4). Let us suppose that the sinusoidal condensed phase is kept at temperature $T_{\rm I}$ and is set at rest, whereas the plane condensed phase is kept at temperature $T_{\rm II}$ and may be moving in its surface in the X_2 direction with a constant speed $U_{\rm II}$. We denote by $p_{\rm I}$ the saturation pressure of the vapor at temperature T_{I} and by $p_{\rm II}$ that at temperature $T_{\rm II}$. We take L as the reference length and $T_{\rm I}$, $p_{\rm I}$, and $n_{\rm I} = p_{\rm I}/kT_{\rm I}$ as the reference quantities, i.e., $T_r = T_I$, $p_r = p_I$, and $n_r = n_I$. Then, the Knudsen number Kn of the system is given by $Kn = l_r/L$, where l_r is the mean free path of the vapor molecules in the equilibrium state at rest with temperature $T_{\rm I}$ and pressure $p_{\rm I}$ (or molecular number density $n_{\rm I}$). We assume that the flow field is periodic in the X_2 direction with period 2L and consider the problem in the closed domain

$$A\cos\left(\frac{\pi X_2}{L}\right) \leq X_1 \leq L, \quad 0 \leq X_2 \leq 2L.$$
(26)

We denote by n_{av}^B the average molecular number density of the noncondensable gas over the domain (26) and let it be in the form of Eq. (17), i.e., $n_{av}^B/n_I = \Delta Kn$, in order to be consistent with Eq. (1). The problem is then characterized by the following parameters:

$$\frac{T_{\rm II}}{T_{\rm I}}, \quad \frac{p_{\rm II}}{p_{\rm I}}, \quad \hat{U}_{\rm II} = \frac{U_{\rm II}}{(2kT_{\rm I}/m^A)^{1/2}}, \quad \frac{A}{L}, \quad \text{Kn}, \quad \Delta.$$
 (27)

We investigate the steady behavior of the vapor and the noncondensable gas in the continuum limit (Kn \rightarrow 0), where the average concentration of the noncondensable gas becomes infinitesimal $(n_{av}^B/n_1\rightarrow 0)$.

We restrict ourselves to the case where evaporation takes place on the plane condensed phase and condensation on the sinusoidal one. In addition, we assume, just for simplicity, that the speed of condensation on the sinusoidal condensed phase is subsonic. The present problem has been investigated on the basis of the same Euler system in Ref. 18 in the case where the plane condensed phase is at rest ($\hat{U}_{\rm II}$ =0). This case will also be included in the following discussions.

Now let us summarize the boundary conditions for the Euler set of equations (5a)–(5d) in the present problem. The boundary condition on the plane condensed phase, where evaporation is assumed to be taking place, is given by Eq. (7), i.e., at $x_1=1$,

$$M_n \le 1, \quad \hat{v}_2 = \hat{U}_{\mathrm{II}}, \tag{28a}$$

$$\hat{p}^{A} = \frac{p_{\mathrm{II}}}{p_{\mathrm{I}}} h_{1}(M_{n}), \quad \hat{T} = \frac{T_{\mathrm{II}}}{T_{\mathrm{I}}} h_{2}(M_{n}),$$
(28b)

with

$$M_n = \sqrt{6/5}\hat{T}^{-1/2}|\hat{v}_1|. \tag{29}$$

In Eqs. (28) and (29) and in Eqs. (30)–(33) below, the notation $(\cdot)_b$ is omitted because it is obvious that the quantities occurring in the boundary conditions are their boundary values. The boundary condition on the sinusoidal condensed phase, where subsonic condensation is assumed to be taking place, is given by Eqs. (8a) and (9). That is, at $x_1 = (A/L)\cos(\pi x_2)$,

$$\hat{p}^{A} = \mathcal{F}_{s}\left(M_{n}, M_{t}, \hat{T}, \frac{l_{r}}{\hat{n}^{A}l_{b}}\tilde{\Gamma}\right),$$
(30)

where $M_n(<1)$ and M_t are given by

$$M_n = \sqrt{6/5} \hat{T}^{-1/2} |\hat{v}_j n_j|, \qquad (31a)$$

$$M_t = \sqrt{6/5} \hat{T}^{-1/2} |\hat{v}_j t_j|, \qquad (31b)$$

with n_i and t_i being unit normal and tangential vectors to the sinusoidal condensed phase, and l_b the mean free path of the molecules of the vapor in the equilibrium state at rest with temperature $T_1 \hat{T}$ and number density $n_1 \hat{n}^A$. In addition, in the present two-dimensional problem, Eq. (14) holds along the sinusoidal boundary with $\hat{\mathcal{N}}_i^B$ given by [see Eq. (10a)]





FIG. 5. Streamlines of the vapor flow for $\Delta = 2$ in the case of $T_{\rm II}/T_{\rm I}=1$, $p_{\rm II}/p_{\rm I}=2$, and A/L=0.2. (a) $\hat{U}_{\rm II}=0$, (b) $\hat{U}_{\rm II}=0.1$, (c) $\hat{U}_{\rm II}=0.2$, (d) $\hat{U}_{\rm II}=0.3$. The corresponding streamlines in the pure-vapor case ($\Delta=0$) are shown by dashed lines. Here, the molecules of the noncondensable gas are assumed to be mechanically identical with those of the vapor.

FIG. 6. Streamlines of the vapor flow for $\Delta = 4$ in the case of $T_{\rm II}/T_{\rm I}=1$, $p_{\rm II}/p_{\rm I}=2$, and A/L=0.2. (a) $\hat{U}_{\rm II}=0$, (b) $\hat{U}_{\rm II}=0.1$, (c) $\hat{U}_{\rm II}=0.2$, (d) $\hat{U}_{\rm II}=0.3$. The corresponding streamlines in the pure-vapor case ($\Delta=0$) are shown by dashed lines. See the caption of Fig. 5.



FIG. 7. Isolines of the flow speed $|v_i|$ for $\hat{U}_{II}=0.1$, $T_{II}/T_I=1$, $p_{II}/p_I=2$, and A/L=0.2. (a) Pure vapor ($\Delta = 0$), $|v_i|/(2kT_1/m^A)^{1/2}=0.164+0.01m (m=0,...,8)$; (b) $\Delta=2$, $|v_i|/(2kT_1/m^A)^{1/2}=0.075+0.015m (m=0,...,10)$. See the caption of Fig. 5.



FIG. 8. Isolines of the number density n^A for $\hat{U}_{II}=0.1$, $T_{II}/T_I=1$, $p_{II}/p_I=2$, and A/L=0.2. (a) Pure vapor ($\Delta =0$), $n^A/n_I=1.504+0.004m$ (m=0,...,8); (b) $\Delta=2$, $n^A/n_I=1.646+0.004m$ (m=0,...,10). See the caption of Fig. 5.

$$\hat{\mathcal{N}}_{i}^{\mathcal{B}} = \frac{l_{b}}{l_{r}} \hat{n}^{A} \hat{T}^{1/2} G_{s} \left(M_{n}, M_{i}, \hat{T}, \frac{l_{r}}{\hat{n}^{A} l_{b}} \tilde{\Gamma} \right) a_{i}, \qquad (32)$$

$$\Delta = \frac{\sqrt{\pi}}{4} \int \tilde{\Gamma} ds, \qquad (34)$$

where

$$a_i = \begin{cases} t_i & (\text{for } \hat{v}_j t_j \ge 0), \\ -t_i & (\text{for } \hat{v}_j t_j < 0). \end{cases}$$
(33)

[Here, a_i for $\hat{v}_j t_j = 0$ is defined as t_i ; see the sentence following that containing Eq. (12).] Note that the ratio l_r/l_b appearing in the functions \mathcal{F}_s and G_s is given by $l_r/l_b = \hat{n}^A / \hat{T}^{1/2}$ for the GSB model. In order to complete the boundary condition, we need the relation between Δ and $\tilde{\Gamma}$. This relation takes the following form:

where ds is the line element along the sinusoidal boundary in the dimensionless x_1x_2 plane, and the range of integration is from $x_2=0$ to 2. This relation is obtained by noting that $(\sqrt{\pi}/2)n_1l_r\tilde{\Gamma}$ is the total number of the noncondensable gas in the Knudsen layer per unit area of the boundary [see the paragraph containing Eq. (15) in Sec. II C] and that its total number in the entire domain per unit width in X_3 , i.e., $n_{av}^B(2L^2)$, is therefore given by $(\sqrt{\pi}/2)n_1l_rL\int \tilde{\Gamma} ds$. Finally, these boundary conditions are supplemented by the condition $h(x_2=2)=h(x_2=0)$ $(h=\hat{\rho}^A, \hat{p}^A, \hat{v}_i, \text{ or } \hat{T})$ and $\tilde{\Gamma}(x_2=2)$





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 $=\widetilde{\Gamma}(x_2=0)$, which corresponds to the periodic boundary condition.

We solve this boundary-value problem numerically by means of a standard finite-difference method. Here, we omit the description about the numerical method and show only the results. The computation is carried out for A/L=0.2, $T_{\rm II}/T_{\rm I}=1$, and $p_{\rm II}/p_{\rm I}=2$.

Figure 5 shows the streamlines of the vapor flow for $\hat{U}_{\text{II}}=0, 0.1, 0.2, \text{ and } 0.3 \text{ in the case of } \Delta=2, \text{ and Fig. 6 shows}$ those for the same \hat{U}_{II} in the case of $\Delta = 4$. In these figures, the corresponding streamlines for a pure-vapor flow ($\Delta = 0$) are also shown by dashed lines. As is seen from the figures, the flow pattern of the vapor strongly depends on Δ . (In the case of $\hat{U}_{\text{II}}=0$, the pattern of the streamlines for $\Delta=2$ and that for $\Delta = 4$ are quite similar [see Figs. 5(a) and 6(a)]. But, as is discussed in Ref. 18, the values of the quantities such as the flow speed $|v_i|$ and the number density n^A are different (see Sec. 4 of Ref. 18).) Let us consider the case where \hat{U}_{II} is finite [cf. Figs. 5(b)-5(d) and 6(b)-6(d)]. The streamlines of the vapor flow near the plane condensed phase show a similar tendency to those in the two-surface problem (Sec. III A). That is, the presence of the noncondensable gas in the Knudsen layer on the sinusoidal condensed phase weakens condensation there and thus evaporation on the plane condensed phase, so that the streamlines become more tangential to the plane condensed phase for larger values of Δ . In the purevapor case ($\Delta = 0$), the shape of the condensing surface has a weak effect on the overall vapor flow, and its streamlines are nearly parallel and do not differ much from those of the two-surface problem. When the noncondensable gas is contained in the Knudsen layer on the condensing surface, the magnitude of the component of the flow velocity of the vapor normal to the condensed phase is reduced there. On the other hand, the magnitude of its tangential component is not reduced much because the noncondensable gas is mobile along the condensed phase and therefore does not give strong resistance. Therefore, the incidence of the streamlines becomes more tangential for larger values of Δ . In this way, the vapor flow is more affected by the geometry of the condensing surface when Δ is large. When $\hat{U}_{\text{II}}=0$ [Figs. 5(a) and 6(a)], the problem is symmetric with respect to the X_1 axis

TABLE IV. Mass-flow rate $\hat{\mathcal{M}}$ for various values of \hat{U}_{II} and Δ in the case $T_{\text{II}}/T_{\text{I}}=1$, $p_{\text{II}}/p_{\text{I}}=2$, and A/L=0.2. Here, the molecules of the noncondensable gas are assumed to be mechanically identical with those of the vapor.

$\hat{U}_{\mathrm{II}} ackslash \Delta$	0	2	4
0	0.4987	0.3734	0.2863
0.1	0.4984	0.3580	0.2773
0.2	0.4979	0.3442	0.2727
0.3	0.4989	0.3435	0.2593

and thus $\hat{\mathcal{N}}_{i}^{\mathcal{B}}t_{i}=0$ holds identically. Therefore, it follows from Eqs. (13) and (32) that $M_{t}=0$ on the sinusoidal condensed phase. In other words, the vapor condenses perpendicularly there. In contrast, the vapor condenses obliquely in the purevapor case. This difference has been pointed out in Ref. 18.

The isolines of the flow speed $|v_i|$, those of the number density n^A , and those of the temperature T are shown in Figs. 7–9, respectively, for $\hat{U}_{\rm II}$ =0.1. In Figs. 7–9, the isolines of the quantities for the pure vapor $(\Delta=0)$ are shown in (a), while those for $\Delta = 2$ in (b). Since the motion of the plane condensed phase is relatively slow ($\hat{U}_{II}=0.1$), the density and temperature variations are small. Therefore, a small error can change isolines significantly. It should be understood that the isolines in Figs. 8 and 9 give information about the almost uniform values of the number density and temperature of the vapor but do not show the detailed structure of their small variations accurately. Let \mathcal{M} be the rate of the mass of the vapor evaporating from the part $0 \le X_1 \le 2L$ of the plane condensed phase per unit time and per unit width in the X_3 direction. Its dimensionless counterpart $\hat{\mathcal{M}}$, defined by $\hat{\mathcal{M}}$ $=\mathcal{M}/m^A n_{\rm I} (2kT_{\rm I}/m^A)^{1/2}L$, is shown in Table IV.

In summary of all the results presented here, the noncondensable gas with an infinitesimal average concentration completely changes the global two-dimensional flow field of the vapor.

The quantity $\tilde{\Gamma}$, which is a function of the position on the boundary, is a measure of accumulation of the noncondensable gas inside the thickless Knudsen layer on the condensing boundary. The distribution of $\tilde{\Gamma}$ along the sinusoidal con-

510 0.30.20.14 8 0.2 0.1 0.3 $\hat{U}_{II}=0$ 3 6 1 ۱Ľ 2° 0<u>`</u> 0^{ι}_0 0.51.52 0.51.51 1 $\mathbf{2}$ X_2/L X_2/L (a) $\Delta = 2$ (b) $\Delta = 4$

FIG. 10. Distribution of $\tilde{\Gamma}$ along the sinusoidal condensed phase for various values of $\hat{U}_{\rm II}$ in the case of $T_{\rm II}/T_{\rm I}$ = 1, $p_{\rm II}/p_{\rm I}$ =2, and A/L=0.2. (a) Δ =2, (b) Δ =4. See the caption of Fig. 5.



densed phase is shown in Fig. 10 for various values of \hat{U}_{II} in the case of $\Delta = 2$ [Fig. 10(a)] and 4 [Fig. 10(b)]. Let us define a kind of average flow speed $\mathcal{V}^{\mathcal{B}}$ of the noncondensable gas inside the Knudsen layer along the sinusoidal condensed phase by

$$\widetilde{\Gamma}\mathcal{V}^{B} = \widehat{\mathcal{N}}_{i}^{B}t_{i}.$$
(35)

Recall that $\hat{\mathcal{N}}_{i}^{B}t_{i}$, which is the total particle flux of the noncondensable gas in the Knudsen layer, is constant along the boundary [Eq. (14)]. The distribution of \mathcal{V}^{B} along the sinusoidal condensed phase is shown in Fig. 11 for $\hat{U}_{II}=0, 0.1,$ 0.2, and 0.3 in the case of $\Delta=2$ [Fig. 11(a)] and 4 [Fig. 11(b)]. When $\hat{U}_{II}=0, \mathcal{V}^{B}$ vanishes identically (see Sec. 4 of Ref. 18). The average speed \mathcal{V}^{B} naturally increases with the increase of \hat{U}_{II} .

Finally, we show some results for relatively small Δ . If the amount of the noncondensable gas is large enough, it distributes over the entire boundary (see Fig. 10). On the other hand, when the amount of the noncondensable gas becomes small, it may vanish (or equivalently, $\tilde{\Gamma}$ may vanish) in a certain part of the boundary. Since we are considering a steady flow, we need to assume that the part which contains the noncondensable gas does not move. This means that its particle flux $\hat{\mathcal{N}}_i^{\mathcal{B}} t_i$ vanishes and thus $M_t=0$ holds because of Eq. (13) with nonzero $\tilde{\Gamma}$. To summarize, Eq. (30) with M_t =0 holds in the part which contains the noncondensable gas, and Eq. (30) with $\tilde{\Gamma}=0$ holds in the part which does not

FIG. 11. Distribution of \mathcal{V}^{B} along the sinusoidal condensed phase for various values of \hat{U}_{II} in the case of $T_{\text{II}}/T_{\text{I}}=1$, $p_{\text{II}}/p_{\text{I}}=2$, and A/L=0.2. (a) $\Delta=2$, (b) $\Delta=4$. Here, $\mathcal{V}^{B}=0$ holds identically for $\hat{U}_{\text{II}}=0$. See the caption of Fig. 5.

contain it. Figure 12 shows the distributions of $\tilde{\Gamma}$ and \mathcal{V}^{B} along the sinusoidal boundary, in the case of \hat{U}_{II} =0.1, for several values of Δ , i.e., $\Delta = 2, 1, 0.5, 0.1$, and 0.05. When Δ is not small, the noncondensable gas distributes over the whole boundary. But for small Δ (Δ =0.1, 0.05), the noncondensable gas tends to disappear from the part $0 \le X_2/L \le 1$ and from the part near $X_2/L=2$. (The \mathcal{V}^B does not have a meaning in the part without the noncondensable gas.) At the left-side edge of the part with the noncondensable gas in Fig. 12(a), the slope of the distribution of $\tilde{\Gamma}$ tends to become very steep. If the distribution of a physical quantity, e.g., $\hat{\rho}^A$, \hat{v}_i , or \hat{T} , along the sinusoidal condensed phase is plotted as a function of X_2/L , the derivative of the curve seems to be discontinuous at the corresponding point, and its limit from the right seems to be (plus or minus) infinity. In contrast, at the right-hand edge of the part with the noncondensable gas, $\overline{\Gamma}$ disappears smoothly and no singularity appears, that is, the curve of any physical quantity is smooth at the corresponding point. Here, we should recall that the Euler set of equations is derived from the Hilbert solution of the Boltzmann equation with a moderately varying property.¹⁸ Therefore, if such a singularity as described above appears in the physical quantities, it violates the assumption for the Hilbert solution. In other words, the present fluid-dynamic system is not theoretically applicable to such situations. In order to treat this type of flow correctly, we need a two-dimensional analysis based on kinetic theory in the neighborhood of the singular point.



FIG. 12. Distribution of $\tilde{\Gamma}$ (a) and that of \mathcal{V}^{B} (b) along the sinusoidal condensed phase for various values of Δ for $\hat{U}_{\text{II}}=0.1$, $T_{\text{II}}/T_{\text{I}}=1$, $p_{\text{II}}/p_{\text{I}}=2$, and A/L=0.2. See the caption of Fig. 5.

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FIG. 13. Vapor flow between cylindrical condensed phases in the presence of a noncondensable gas.

C. Vapor flows between two noncoaxial cylindrical condensed phases

In this section, we consider a vapor (A component) and a noncondensable gas (B component) between two noncoaxial circular cylinders made of the condensed phase as shown in Fig. 13, where the X_1 , X_2 , and X_3 axes are assumed to form the right-hand system. Let $R_{\rm I}$ (or $R_{\rm II}$) be the radius of the inner (or outer) cylinder, T_{I} (or T_{II}) the temperature of the inner (or outer) cylinder, D the distance between the two axes, and $p_{\rm I}$ (or $p_{\rm II}$) the saturation pressure of the vapor at temperature $T_{\rm I}$ (or $T_{\rm II}$). The inner cylinder may be rotating (counterclockwise in Fig. 13) around its axis with a constant angular velocity, whereas the outer cylinder is kept at rest. We denote by $U_{\rm I}$ the speed of the surface of the inner cylinder and take $R_{\rm I}$ as the reference length and $T_{\rm I}$, $p_{\rm I}$, and $n_{\rm I}$ $=p_{\rm I}/kT_{\rm I}$ as the reference quantities, i.e., $L=R_{\rm I}$, $T_r=T_{\rm I}$, p_r $=p_{\rm I}$, and $n_r = n_{\rm I}$. Then the Knudsen number of the system is given by $Kn = l_r/R_I$ with l_r being the mean free path of the vapor molecules in the equilibrium state at rest with temperature $T_{\rm I}$ and pressure $p_{\rm I}$ (or molecular number density $n_{\rm I}$). As before, the average molecular number density of the noncondensable gas over the whole domain is denoted by n_{av}^{B} and is put in the form $n_{av}^B/n_I = \Delta Kn$, where Δ is a constant to be specified. The problem is characterized by the following parameters:

$$\frac{T_{\rm II}}{T_{\rm I}}, \quad \frac{p_{\rm II}}{p_{\rm I}}, \quad \hat{U}_{\rm I} = \frac{U_{\rm I}}{(2kT_{\rm I}/m^{\rm A})^{1/2}}, \quad \frac{R_{\rm II}}{R_{\rm I}}, \quad \frac{D}{R_{\rm I}}, \quad \text{Kn}, \quad \Delta.$$
(36)

We investigate the steady behavior of the vapor and the effect of the trace of the noncondensable gas in the continuum limit (Kn \rightarrow 0).

The behavior of the vapor is described by the Euler equations (5a)–(5d). Let us consider the case where the vapor is evaporating from the inner cylinder and condensing onto the outer cylinder and assume that the speed of condensation on the outer cylinder is subsonic. Then the boundary condition on the inner cylinder $(x_1^2+x_2^2=1)$, where evaporation is taking place, is given by

$$M_n \le 1, \quad \hat{v}_j t_j = \hat{U}_{\mathrm{I}}, \tag{37a}$$

$$\hat{p}^{A} = h_{1}(M_{n}), \quad \hat{T} = h_{2}(M_{n}),$$
(37b)

where

$$M_n = \sqrt{6/5} \hat{T}^{-1/2} |\hat{v}_j n_j|, \qquad (38)$$

and n_i and t_i are the unit normal and tangential vectors to the inner cylinder, n_i pointing into the region of the vapor, and n_i , t_i , and the X_3 axis forming the right-hand system. In Eqs. (37) and (38) and Eqs. (39)–(42) below, the notation $(\cdot)_b$ is omitted as in Sec. III B. On the other hand, the boundary condition on the outer cylinder $[(x_1-D/R_I)^2+x_2^2=(R_{II}/R_I)^2]$, where condensation is taking place, is given by

$$\hat{p}^{A} = \frac{p_{\mathrm{II}}}{p_{\mathrm{I}}} \mathcal{F}_{s} \left(M_{n}, M_{l}, \frac{T_{\mathrm{I}}}{T_{\mathrm{II}}} \hat{T}, \frac{l_{r}}{\hat{n}^{A} l_{b}} \widetilde{\Gamma} \right),$$
(39)

where

$$M_n = \sqrt{6/5} \hat{T}^{-1/2} |\hat{v}_j n_j|, \qquad (40a)$$

$$M_t = \sqrt{6/5} \hat{T}^{-1/2} |\hat{v}_j t_j|, \qquad (40b)$$

and n_i and t_i are the unit normal and tangential vectors to the outer cylinder, n_i pointing into the region of the vapor, and n_i , t_i , and the X_3 axis forming the right-hand system. In addition, Eq. (14) holds along the outer cylinder, where $\hat{\mathcal{N}}_i^{\mathcal{B}}$ is given by



FIG. 14. Streamlines of the vapor flow for $\Delta = 2$ in the case of $T_{\rm II}/T_{\rm I}=1$, $p_{\rm II}/p_{\rm I}=0.5$, $R_{\rm II}/R_{\rm I}=2$, and $D/R_{\rm I}=0.5$. (a) $\hat{U}_{\rm I}=0$, (b) $\hat{U}_{\rm I}=0.1$, (c) $\hat{U}_{\rm I}=0.2$. The corresponding streamlines in the pure-vapor case ($\Delta=0$) are shown by dashed lines. Here, the molecules of the noncondensable gas are assumed to be mechanically identical with those of the vapor.

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$$\hat{\mathcal{N}}_{i}^{\mathcal{B}} = \frac{l_{b}}{l_{r}} \hat{n}^{A} \hat{T}^{1/2} G_{s} \left(M_{n}, M_{t}, \frac{T_{\mathrm{II}}}{T_{\mathrm{II}}} \hat{T}, \frac{l_{r}}{\hat{n}^{A} l_{b}} \widetilde{\Gamma} \right) a_{i}, \tag{41}$$

with

$$a_i = \begin{cases} t_i & \text{for } \hat{v}_j t_j \ge 0, \\ -t_i & \text{for } \hat{v}_j t_j < 0. \end{cases}$$
(42)

[Here, a_i for $\hat{v}_j t_j = 0$ is defined as t_i ; see the sentence following that containing Eq. (12).] Finally, the relation between Δ and $\tilde{\Gamma}$ is given by

$$\Delta = \frac{1}{2\sqrt{\pi}} \left[\left(\frac{R_{\rm II}}{R_{\rm I}} \right)^2 - 1 \right]^{-1} \int \tilde{\Gamma} ds, \qquad (43)$$

where ds is the (dimensionless) line element along the outer cylinder, and the integration is carried out over the whole circle. This relation is obtained in a similar way as Eq. (34).

We analyze the above boundary-value problem of the Euler set of equations by a finite-difference method. The result for $T_{\rm II}/T_{\rm I}=1$, $p_{\rm II}/p_{\rm I}=0.5$, $R_{\rm II}/R_{\rm I}=2$, and $D/R_{\rm I}=0.5$ is shown in Figs. 14–17. Figure 14 shows the streamlines of the vapor flow for $\hat{U}_{\rm I}=0$, 0.1, and 0.2 in the case of $\Delta=2$. In the figure, the streamlines for the corresponding pure-vapor case ($\Delta=0$) are also shown by dashed lines. Figure 15 shows the isolines of the molecular number density of the vapor n^A in the case of $\hat{U}_{\rm I}=0.1$, where the isolines for the pure-vapor case ($\Delta=0$) are shown in (a) and those for $\Delta=2$ in (b). Figure 16 shows the distribution of $\tilde{\Gamma}$ and that of \mathcal{V}^B along the

FIG. 15. Isolines of the number density n^{A} for \hat{U}_{I} =0.1 in the case T_{II}/T_{I} =1, p_{II}/p_{I} =0.5, R_{II}/R_{I} =2, and D/R_{I} =0.5. (a) Pure vapor (Δ =0), n^{A}/n_{I} =0.67+0.01m (m =0,...,5); (b) Δ =2, n^{A}/n_{I} =0.744+0.006m (m =0,...,5). See the caption of Fig. 14.

outer cylinder for $\Delta=2$. Here, \mathcal{V}^B is the same as that used in Sec. III B [Eq. (35)], and θ ($0 \le \theta < 2\pi$) is the angle between the X_1 axis and the line connecting a point on the outer cylinder and its axis [the point (D,0) in Fig. 13] measured counterclockwise. Let $\mathcal{M}=m^A n_I (2kT_I/m^A)^{1/2}R_I\hat{\mathcal{M}}$ be the total mass-flow rate of the vapor evaporating from the inner cylinder (per unit width in the X_3 direction and per unit time). The dimensionless mass-flow rate $\hat{\mathcal{M}}$ versus Δ for $\hat{U}_I=0, 0.1, 0.2,$ and 0.3 is shown in Fig. 17.

The features of the streamlines can basically be understood from the discussion in Sec. III B. The dimensionless mass-flow rate $\hat{\mathcal{M}}$ naturally decreases with the increase of Δ . Faster rotation of the inner cylinder leads to a pressure rise on the outer cylinder because of stronger centrifugal effect and thus causes stronger condensation. Therefore, $\hat{\mathcal{M}}$ increases as $\hat{U}_{\rm I}$ increases. The results shown here again demonstrate that the noncondensable gas with an infinitesimal average concentration has a significant effect on the overall vapor flow field.

IV. CONCLUDING REMARKS

In a previous paper,¹⁸ we derived the fluid-dynamic system that describes steady flows of a vapor around its condensed phase, on the surface of which evaporation or condensation is taking place, in the continuum limit $(Kn \rightarrow 0)$ in the presence of a noncondensable gas whose amount is in-



FIG. 16. Distribution of $\tilde{\Gamma}$ and that of \mathcal{V}^{B} along the outer cylinder for various \hat{U}_{I} in the case $\Delta = 2$. (a) $\tilde{\Gamma}$, (b) \mathcal{V}^{B} . Here, $\theta \ (0 \le \theta < 2\pi)$ is the angle between the X_{1} axis and the line connecting a point on the outer cylinder and its axis [the point (D,0) in Fig. 13] measured counterclockwise. In (b), $\mathcal{V}^{B}=0$ holds identically in the case $\hat{U}_{I}=0$. See the caption of Fig. 14.

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FIG. 17. $\hat{\mathcal{M}}$ vs Δ for various \hat{U}_{I} . The symbols $\bigcirc, \Box, \triangle$, and \diamond indicate the numerical data, which are connected by spline curves. See the caption of Fig. 14.

finitesimal in comparison with that of the vapor. In the case of a closed domain, the amount of the noncondensable gas is expressed more precisely in such a way that its average concentration is infinitesimal [O(Kn)]. In the same paper, we pointed out that such a small amount of the noncondensable gas can affect the overall vapor flow significantly, accumulating in the Knudsen layer with an infinitesimal thickness on the condensing surface. The present paper is devoted to the application of the fluid-dynamic system mentioned above to some concrete problems that demonstrate the effect of the trace of the noncondensable gas clearly.

The fluid-dynamic system consists of the set of the compressible Euler equations and its boundary conditions constructed numerically under the following assumptions: (i) the mechanical property of the molecules of the vapor and that of the noncondensable gas are the same and (ii) the model Boltzmann equations (the BGK and GSB models) describe the behavior in the Knudsen layers (Sec. II). On the basis of this system, the two-surface problem of evaporation and condensation (Sec. III A), vapor flows between a plane and a sinusoidal condensed phase (Sec. III B), and vapor flows between two noncoaxial cylindrical condensed phases (Sec. III C) were investigated numerically. In these problems, it is clearly seen that the noncondensable gas with an infinitesimal average concentration has significant effects on the overall vapor flows in the continuum limit.

ACKNOWLEDGMENTS

The present work was supported by the Grant-in-Aid for Scientific Research (Grant No. 14350047) from the Japan Society for the Promotion of Science and by the Center of Excellence for Research and Education on Complex Functional Mechanical Systems (the COE program of the Ministry of Education, Culture, Sports, Science and Technology, Japan).

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