



Does Maxwell's hypothesis of air saturation near the surface of evaporating liquid hold at all spatial scales?

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The classical model of evaporation of liquids hinges on Maxwell's assumption that the air near the liquid's surface is saturated. It allows one to find the evaporative flux without considering the interface separating liquid and air. Maxwell's hypothesis is based on an implicit assumption that the vapour-emission capacity of the interface exceeds the throughput of air (i.e. its ability to pass the vapour on to infinity). If this is indeed so, then the air adjacent to the liquid would get quickly saturated, justifying Maxwell's hypothesis. In the present paper, the so-called diffuse-interface model is used to account for the interfacial physics and thus derive a generalised version of Maxwell's boundary condition for the near-interface vapour density. It is then applied to a spherical drop floating in air. It turns out that the vapour-emission capacity of the interface exceeds the throughput of air only if the drop's radius is $r_d \gtrsim 10 \mu\text{m}$, but for $r_d \approx 2 \mu\text{m}$, the two are comparable. For $r_d \lesssim 1 \mu\text{m}$, evaporation is interface-driven, and the resulting evaporation rate is noticeably smaller than that predicted by the classical model.

Key words: condensation/evaporation, drops

1. Introduction

Consider a liquid drop floating in the air and evaporating. In its simplest formulation, this problem was examined more than 140 years ago by Maxwell (1877, 1890), and its more elaborate versions are still explored now. Several of these are relevant to the present study: Sobac *et al.* (2015), Cossali & Tonini (2019) and Finneran (2021) argued that the dependence of the fluid's properties on the temperature can affect the evaporation rate strongly; Talbot *et al.* (2016) examined transient deviations of drop evaporation from the

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quasi-steady regime; Finneran, Garner & Nadal (2021) showed that the transient effects can cause a noticeable departure from the so-called ‘ d^2 law’ (stating that the time of a drop’s evaporation is proportional to its diameter squared); Long, Micci & Wong (1996), Landry *et al.* (2007), Jakubczyk *et al.* (2012), Hołyst *et al.* (2013), Hołyst, Litniewski & Jakubczyk (2017), Rana, Lockerby & Sprittles (2018, 2019) and Zhao & Nadal (2023) showed that the ‘ d^2 law’ can be violated by out-of-equilibrium gas kinetics. A more detailed review of the recent literature on evaporating drops can be found in Sazhin (2017).

Most work on evaporation has been based on Maxwell’s hypothesis that the air near the drop’s surface – or, more generally, near any liquid/air interface – is close to saturation. This allows one to solve the problem without knowing anything about the interfacial dynamics. The only exceptions are models accounting for the so-called Knudsen layer – a layer where the out-of-equilibrium kinetics plays a part (e.g. Rana *et al.* 2018, 2019; Zhao & Nadal 2023).

The physics behind Maxwell’s hypothesis becomes clear if evaporation is viewed as a two-tier process: first, the interface emits molecules of the liquid into the air; second, the air passes them on to infinity. If the former emits more vapour than the latter can pass on, then the vapour accumulates near the drop – bringing the adjacent air layer close to saturation and the interface close to thermodynamic equilibrium. As a result, the interface reduces vapour emission in line with the air throughput. In terms of this mechanism, Maxwell’s hypothesis amounts to an assumption that the vapour-emission capacity of the interface exceeds the throughput of air.

Maxwell could not have tested this assumption, as there were no quantitative models of interfaces at the time. Two such models exist now: the Enskog–Vlasov kinetic equation (de Sobrino 1967; Grmela 1971; Grmela & Garcia-Colin 1980*a,b*; Frezzotti, Gibelli & Lorenzani 2005; Barbante, Frezzotti & Gibelli 2015; Frezzotti & Barbante 2017; Frezzotti *et al.* 2018; Benilov & Benilov 2018, 2019*a,b*; Struchtrup & Frezzotti 2022) and the diffuse-interface model (see a review by Anderson, McFadden & Wheeler 1998). The latter is much simpler, thus it makes a better starting point.

The diffuse-interface model (DIM) is based on two assumptions regarding the van der Waals force.

- (i) Van der Waals interactions are pairwise – hence the net force affecting a molecule is an algebraic sum of the forces exerted by the other molecules.
- (ii) The thickness of the interface is much greater than the spatial scale of van der Waals interactions.

Admittedly, assumption (ii) does not hold well at room temperature, in which case the interfacial thickness rarely exceeds the molecular size by an order of magnitude (e.g. Liu, Harder & Berne 2005; Verde, Bolhuis & Campen 2012; Pezzotti, Galimberti & Gaigeot 2017; Pezzotti, Serva & Gaigeot 2018; Dodia *et al.* 2019; and references therein). Thus the DIM should be viewed as a means to estimate qualitatively the importance of interfacial dynamics for evaporation, whereas quantitative predictions should be left to the Enskog–Vlasov kinetic equation (which can handle thin interfaces). These two models are to each other what the first term of a Taylor series is to the exact value of the function.

The DIM has been used many times before – for phase transitions in ferroelectrics (Ginzburg 1960), spinodal decomposition (e.g. Cahn 1961; Lowengrub & Truskinovsky 1998), crystal growth (e.g. Collins & Levine 1985; Tang, Carter & Cannon 2006; Heinonen *et al.* 2016; Philippe, Henry & Plapp 2020), solidification (e.g. Stinner, Nestler & Garcke 2004; Nestler, Garcke & Stinner 2005), polymers (e.g. Thiele, Madruga & Frastia 2007;

Madruga & Thiele 2009), electrowetting (e.g. Lu *et al.* 2007), contact lines in liquids (e.g. Pismen & Pomeau 2000; Ding & Spelt 2007; Yue, Zhou & Feng 2010; Yue & Feng 2011; Sibley *et al.* 2014; Ding *et al.* 2017; Borcia *et al.* 2019; Zhu *et al.* 2019, 2020), Faraday instability (e.g. Borcia & Bestehorn 2014; Bestehorn *et al.* 2021), Rayleigh–Taylor instability (e.g. Zanella *et al.* 2020, 2021), cavitation (e.g. Petitpas *et al.* 2009), nucleation (e.g. Magaletti, Marino & Casciola 2015; Magaletti *et al.* 2016; Gallo, Magaletti & Casciola 2018; Gallo *et al.* 2020), liquid films (e.g. Benilov 2020, 2022a), tumour growth (e.g. Frigeri, Grasselli & Rocca 2015; Rocca & Scala 2016; Dai *et al.* 2017), classification of data (e.g. Bertozzi & Flenner 2012, 2016), and so on. The DIM has also been applied to evaporation and condensation (e.g. Pomeau 1986; Barbante, Frezzotti & Gibelli 2014; Benilov 2022a,b, 2023b), but only in application to liquids evaporating into, or condensating from, their own vapour, not air.

In the present paper, the DIM is used to examine evaporation of liquids into air under ‘normal conditions’, i.e. when the temperature is between 15 °C and 35 °C, and the air pressure is at 1 atm. A generalised Maxwell boundary condition is derived for this case; to facilitate its use without going into the detail of the derivation, this condition is summarised here.

The following notation will be used: ρ_1 is the density of the liquid or its vapour, ρ_2 is that of air. Outside the drop, the air is almost homogeneous, i.e. $\rho_2 \approx \rho_2^{(a)}$. Let $\rho_1^{(v.sat)}(T)$ be the saturated vapour density at a temperature T , and let $\rho_1^{(l.sat)}(T)$ be the matching liquid density. Introduce also the vapour-in-the-air diffusivity \mathcal{D} , the shear and bulk viscosities of air, μ_s and μ_b , respectively, and the specific gas constant R_i of the liquid or its vapour ($i = 1$), or the air ($i = 2$). The interface as a whole is characterised by its curvature C and surface tension σ .

All these parameters are macroscopic, i.e. they characterise the fluid as a continuum, whereas the Korteweg matrix K_{ij} is a microscopic characteristic. It describes the intermolecular (van der Waals) force due to the liquid–liquid (K_{11}), air–air (K_{22}) and liquid–air ($K_{12} = K_{21}$) interactions. Macroscopically, K_{ij} determine the capillary properties of the fluid and thus can be deduced from measurements of its surface tension (see Appendix B). The numerical values of K_{ij} determined this way for water and air are presented later, in (3.12).

Let $\rho_1^{(+)}$ be the macroscopic vapour density immediately outside the interface, and let $(\mathbf{n} \cdot \nabla \rho_1)^{(\pm)}$ be its normal derivative (where \mathbf{n} is directed towards the air). Then the generalised Maxwell boundary condition has the form

$$\rho_1^{(+)} = \rho_1^{(v.sat)} \exp \left[-\frac{R_2}{R_1} \xi A(\xi) + \frac{\sigma C}{\rho_1^{(l.sat)} R_1 T} \right], \quad (1.1)$$

where ξ is the relative vapour flux,

$$\xi = \frac{-\mathcal{D}(\mathbf{n} \cdot \nabla \rho_1)^{(\pm)}}{E_0}, \quad (1.2)$$

and the parameter

$$E_0 = \frac{K_{22}^{1/2} \rho_2^{(a)5/2} (R_2 T)^{1/2}}{\frac{4}{3} \mu_s + \mu_b} \quad (1.3)$$

will be referred to as the ‘vapour-emission capacity’ of the interface. Physically, E_0 is determined by a narrow zone at the outskirts of the interface, driven by the van der Waals

force, pressure gradient and viscosity. The location of this zone suggests that it is a DIM analogue of the Knudsen layer, i.e. the Knudsen layer plus the van der Waals force – which is an order-one effect in the DIM, but is neglected by the standard Hertz–Knudsen kinetic model.

The universal non-dimensional function $A(\xi)$ that appears in (1.1) is also a characteristic of the above-mentioned zone at the outskirts of the interface. It can be approximated by the expression

$$A(\xi) = \frac{\frac{2}{5} \ln(\xi + 3.2867) - 0.81571}{\xi + 3.2867} + \frac{1.3074}{(\xi + 3.2867)^{7/5}}, \quad (1.4)$$

obtained through a combination of asymptotics, numerics and curve fitting (see Appendix C).

The term involving the curvature C in condition (1.1) describes the Kelvin effect, i.e. the shift of the saturated vapour pressure near a curved surface (e.g. Eggers & Pismen 2010; Colinet & Rednikov 2011; Rednikov & Colinet 2013, 2017, 2019; Morris 2014; Janeček *et al.* 2015; Saxton *et al.* 2017). If the interface is flat ($C = 0$) and its vapour-emission capacity exceeds the diffusive flux,

$$E_0 \gg |-D(\mathbf{n} \cdot \nabla \rho_1)^{(+)}|, \quad (1.5)$$

then (1.1) reduces to the classical Maxwell boundary condition.

This paper has the following structure. In § 2, Maxwell’s boundary condition is reviewed briefly in application to evaporation of floating drops. Its generalised version is derived in § 3, and applied to drops in § 4. In § 5, the proposed model is compared to the Hertz–Knudsen kinetic model, and further applications of the former are outlined.

2. The classical model

2.1. Formulation

Consider a mixture of two fluids with partial densities ρ_i , where $i = 1, 2$. The first fluid will be referred to as either ‘liquid’ or ‘vapour’ (depending on the phase it is in), and the second fluid will be referred to as ‘air’.

Under the assumption of spherical symmetry, ρ_i generally vary with the radial coordinate r and time t . In the classical model, however, one assumes that inside the drop

$$\rho_1 = \rho_1^{(l)}, \quad \rho_2 = 0 \quad \text{if } r \in (0, r_d), \quad (2.1)$$

where $\rho_1^{(l)}$ is a known constant, and the drop’s radius r_d depends on t .

Outside the drop, both densities do vary. The air density is much larger than that of the vapour, but their variations are of the same order (as, physically, the pressure should be spatially uniform). As a result, the variations of ρ_2 are relatively weak, so one assumes

$$\rho_2 = \rho_2^{(a)} \quad \text{if } r \in (r_d, \infty), \quad (2.2)$$

where the dry-air density $\rho_2^{(a)}$ is a known parameter. Note that, physically, $\rho_1^{(l)}$ and $\rho_2^{(a)}$ are inter-related by the requirement that the pressure difference between the inside and outside of the drop equals the capillary correction due to the curvature of the interface.

The classical model of evaporation also involves the liquid’s thermal conductivity $\kappa^{(l)}$, air’s thermal conductivity $\kappa^{(a)}$, saturated vapour density $\rho_1^{(v.sat)}$, vaporisation heat Δ , and

vapour-in-the-air diffusivity \mathcal{D} . The dependence of these parameters on the temperature T is supposed to be known.

The classical model is usually combined with the quasi-steady approximation, based on an assumption that the diffusivity and thermal conductivity are sufficiently large (which they are indeed for water under normal conditions). Then the term $\partial\rho_1/\partial t$ in the diffusion equation and the term $\partial T/\partial t$ in the heat-conduction equation can both be neglected. The simplest version of the classical model also neglects the Stefan flow (which carries the vapour away from the surface of the liquid), the Soret effects (the mass flux due to the temperature gradient), and the Dufour effect (the heat flux due to the density gradient).

Then, for the spherically symmetric case, one obtains

$$\frac{\partial}{\partial r} \left(r^2 \frac{\partial T}{\partial r} \right) = 0 \quad \text{if } r \in (0, r_d), \quad (2.3)$$

$$\frac{\partial}{\partial r} \left(r^2 \frac{\partial \rho_1}{\partial r} \right) = 0, \quad \frac{\partial}{\partial r} \left(r^2 \frac{\partial T}{\partial r} \right) = 0 \quad \text{if } r \in (r_d, \infty). \quad (2.4)$$

At the drop's centre, the usual boundary condition applies:

$$\frac{\partial T}{\partial r} = 0 \quad \text{at } r = 0. \quad (2.5)$$

To formulate the boundary conditions at the drop's surface, introduce

$$T^{(\pm)} = \lim_{r \rightarrow r_d \pm 0} T, \quad \left(\frac{\partial T}{\partial r} \right)^{(\pm)} = \lim_{r \rightarrow r_d \pm 0} \frac{\partial T}{\partial r}, \quad (2.6a,b)$$

where $(+)$ and $(-)$ mean 'just outside' and 'just inside' the drop, respectively. Then the continuity of the temperature and heat flux imply

$$T^{(-)} = T^{(+)} \quad \text{at } r = r_d, \quad (2.7)$$

$$\kappa^{(l)} \left(\frac{\partial T}{\partial r} \right)^{(-)} - \kappa^{(a)} \left(\frac{\partial T}{\partial r} \right)^{(+)} = \dot{r}_d \rho_1^{(l)} \Delta \quad \text{at } r = r_d, \quad (2.8)$$

where

$$\dot{r}_d = \frac{dr_d}{dt}. \quad (2.9)$$

As discussed in the Introduction, the classical model assumes the Maxwell boundary condition for the vapour density at the drop's surface, i.e.

$$\rho_1 = \rho_1^{(v.sat)} \quad \text{at } r = r_d, \quad (2.10)$$

and the following conditions should be imposed on the vapour-air mixture far away from the drop:

$$\rho_1 \rightarrow \rho_1^{(a)}, \quad T \rightarrow T^{(a)} \quad \text{as } r \rightarrow \infty. \quad (2.11)$$

To ensure that the liquid is evaporating – not condensating or being in equilibrium – let the vapour at infinity be undersaturated:

$$\rho_1^{(a)} < \rho_1^{(a.sat)(a)}, \quad (2.12)$$

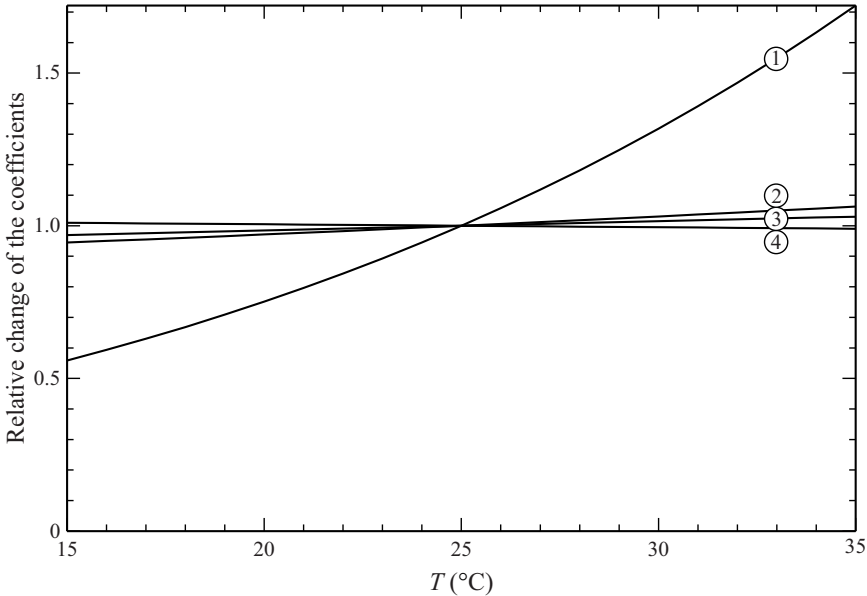


Figure 1. The temperature dependencies of the saturated vapour density $\rho_1^{(v.sat)}$, vapour-in-the-air diffusivity \mathcal{D} , thermal conductivity $\kappa^{(a)}$ of the air, and vaporisation heat Δ (all non-dimensionalised on their reference values): (1) $\rho_1^{(v.sat)}(T)/\rho_1^{(v.sat)}(25^\circ\text{C})$, (2) $\mathcal{D}(T)/\mathcal{D}(25^\circ\text{C})$, (3) $\kappa^{(a)}(T)/\kappa^{(a)}(25^\circ\text{C})$, and (4) $\Delta(T)/\Delta(25^\circ\text{C})$. Curves (1)–(4) correspond to the empirical formulae of Wagner & Pruss (1993), Engineering Toolbox (2018), White (2005) and Lindstrom & Mallard (1997), respectively.

where $\rho_1^{(a.sat)(a)} = \rho_1^{(a.sat)}(T^{(a)})$. Finally, the velocity of the drop’s (receding) surface is related to the evaporative flux by

$$\rho_1^{(l)} \dot{r}_d = \mathcal{D} \frac{\partial \rho}{\partial r} \quad \text{at } r = r_d. \tag{2.13}$$

Boundary-value problem (2.3)–(2.13) governs $\rho_1(r, t)$, $T(r, t)$ and $r_d(t)$.

2.2. The dependence of $\rho_1^{(v.sat)}$ on T

One’s experience with raindrops sitting on one’s skin suggests that evaporative cooling does not exceed several degrees. This seems to enable one to neglect the dependence of the external parameters on the temperature – say, ‘freeze’ them at $T = T^{(a)}$. Such an approximation is indeed applicable to all parameters but one – namely, the saturated vapour density $\rho_1^{(v.sat)}$.

This claim is illustrated for water in figure 1. One can see that the relative change of $\rho_1^{(v.sat)}$ across the room temperature range exceeds those of the remaining parameters.

In this paper, the dependence $\rho_1^{(v.sat)}(T)$ is approximated by Antoine’s equation (Antoine 1888). If written in terms of the density – as opposed to the pressure, as in the original formulation – it takes the form

$$\frac{\rho_1^{(v.sat)}}{\rho_A} = \frac{T_A}{T} \exp\left(-\frac{T_A}{T}\right), \tag{2.14}$$

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where, for water,

$$\rho_A = 6.6326 \times 10^4 \text{ kg m}^{-3}, \quad T_A = 5292 \text{ K.} \quad (2.15a,b)$$

These values have been obtained by fitting (2.14) to the high-accuracy IAPWS empirical formula (Wagner & Pruss 1993) for the room temperature range. The relative deviation of the two formulae is less than 0.23 %, which is much better than the accuracy of Antoine's equation with ρ_A and T_A taken from a thermodynamics handbook.

Not only does Antoine's equation provide an accurate approximation, but it also has physical meaning. As shown by Benilov (2020), (2.14) describes the generic van der Waals fluid in the low-temperature limit – i.e. when T is much closer to the freezing point than to the critical point. Observe also that T_A is large, which is what makes the dependence of $\rho_1^{(v.sat)}$ on T strong.

Equation (2.14) can be rearranged conveniently using a reference temperature (as done by Sobac *et al.* 2015). Choosing that to be $T^{(a)}$, one obtains

$$\frac{\rho_1^{(v.sat)}}{\rho_1^{(v.sat)(a)}} = \frac{T^{(a)}}{T} \exp\left(-\frac{T_A}{T} + \frac{T_A}{T^{(a)}}\right), \quad (2.16)$$

where $\rho_1^{(v.sat)(a)}$ is the saturated vapour density at infinity. One can further assume $(T^{(a)} - T)/T^{(a)} \ll 1$ and reduce (2.16) to

$$\rho_1^{(v.sat)} = \rho_1^{(v.sat)(a)} \exp\left[\frac{T_A}{T^{(a)2}} (T - T^{(a)})\right]. \quad (2.17)$$

Observe that even if T is close to $T^{(a)}$, the expression in the square brackets can still be order-one (because $T_A \gg T^{(a)}$).

2.3. Solution of boundary-value problem (2.3)–(2.13)

Equations (2.3)–(2.4) can be solved readily as

$$T = \text{const}_1 + \frac{\text{const}_2}{r} \quad \text{if } r \in (0, r_d), \quad (2.18)$$

$$\rho_1 = \text{const}_3 + \frac{\text{const}_4}{r}, \quad T = \text{const}_5 + \frac{\text{const}_6}{r} \quad \text{if } r \in (r_d, \infty), \quad (2.19)$$

where the constants of integration can be fixed via boundary conditions (2.5)–(2.11):

$$\left. \begin{aligned} \text{const}_1 &= T^{(a)} + r_d \dot{r}_d \frac{\rho_1^{(l)} \Delta}{\kappa^{(a)}}, & \text{const}_2 &= 0, & \text{const}_3 &= \rho_1^{(a)}, \\ \text{const}_4 &= r_d \left[\rho_1^{(v.sat)} \left(T^{(a)} + r_d \dot{r}_d \frac{\rho_1^{(l)} \Delta}{\kappa^{(a)}} \right) - \rho_1^{(a)} \right], & \text{const}_5 &= T^{(a)}, \\ \text{const}_6 &= r_d^2 \dot{r}_d \frac{\rho_1^{(l)} \Delta}{\kappa^{(a)}}. \end{aligned} \right\} \quad (2.20a-f)$$

Recalling (2.16) for $\rho_1^{(v.sat)}$ and (2.13) for \dot{r}_d , one obtains

$$\underbrace{\frac{\rho_1^{(l)}}{\mathcal{D}\rho_1^{(v.sat)(a)}} \dot{r}_d r_d}_{\text{diffusion}} + \exp \underbrace{\frac{T_A \rho_1^{(l)} \Delta}{T^{(a)2 \kappa^{(a)}} \dot{r}_d r_d}}_{\text{non-isothermality}} = H, \quad (2.21)$$

where

$$H = \frac{\rho_1^{(a)}}{\rho_1^{(v.sat)(a)}} \tag{2.22}$$

is the relative humidity of air far away from the drop.

Ordinary differential equation (2.21) determines $r_d(t)$; the terms on its left-hand side are labelled according to their physical meanings.

Equation (2.21) yields the following expression for a drop’s time of evaporation:

$$t_e = r_d^2(0) \frac{x\rho_1^{(l)}}{2\mathcal{D}\rho_1^{(v.sat)(a)}}, \tag{2.23}$$

where $r_d(0)$ is the drop’s initial radius, and x is the solution of the non-dimensional algebraic equation

$$-x + \exp(-\alpha x) = H, \tag{2.24}$$

with

$$\alpha = \frac{T_A \mathcal{D} \rho_1^{(v.sat)(a)} \Delta}{T^{(a)2} \kappa^{(a)}}. \tag{2.25}$$

Formula (2.23) is often referred to as the ‘ d^2 law’, where d is the drop’s diameter.

It is instructive to estimate α for, say, the midpoint of the room temperature range, $T^{(a)} = 25^\circ\text{C}$, and pressure 1 atm. Letting the liquid be water, one can use the NIST online calculator (Lindstrom & Mallard 1997) to obtain

$$\rho_1^{(l)} = 997.05 \text{ kg m}^{-3}, \quad \rho_1^{(v.sat)(a)} = 0.023075 \text{ kg m}^{-3}, \quad \Delta = 2441.7 \text{ kJ kg}^{-1}, \tag{2.26a-c}$$

whereas for air, Engineering ToolBox (2003, 2009, 2018) yield

$$\rho_2^{(a)} = 1.184 \text{ kg m}^{-3}, \quad \kappa^{(a)} = 0.02624 \text{ W m}^{-1} \text{ K}^{-1}, \quad \mathcal{D} = 2.49 \times 10^{-5} \text{ m}^2 \text{ s}^{-1}. \tag{2.27a-c}$$

Substituting values (2.26)–(2.27) into (2.25), and recalling the value in (2.15) for T_A , one obtains

$$\alpha \approx 3.1828. \tag{2.28}$$

2.4. Discussion

It is interesting to assess the importance of non-isothermal effects by comparing the above model with its isothermal reduction. It can be verified readily that the latter amounts to taking in (2.24) the limit $\alpha \rightarrow 0$, so that $x = 1 - H$. Denoting the isothermal time of evaporation by t'_e , one obtains

$$\frac{t_e}{t'_e} = \frac{\mathcal{H}}{1 - H}. \tag{2.29}$$

It turns out that the isothermal approach noticeably underpredicts the evaporation time. The largest discrepancy is observed at high humidity, in which case (2.24) can be

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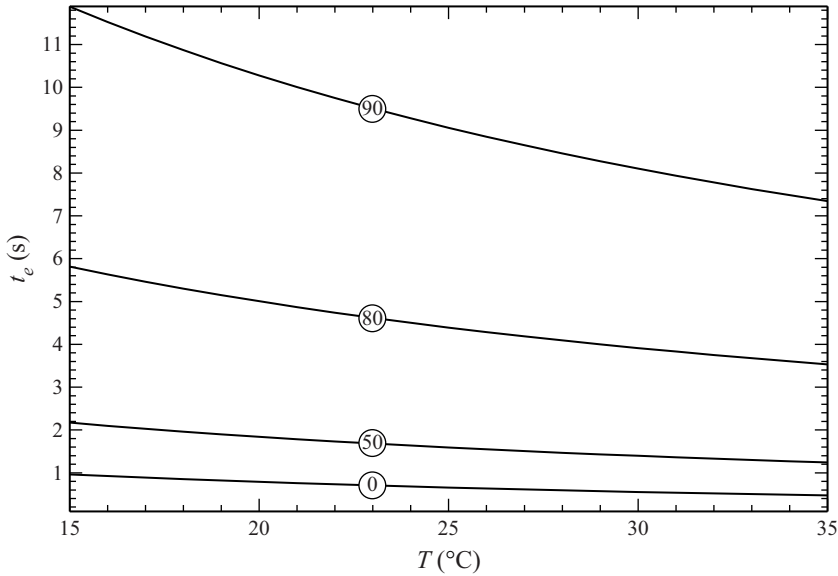


Figure 2. The time of full evaporation of a water drop of radius 16 μm versus the temperature. The curves are marked with the corresponding relative humidity (in percentage).

solved asymptotically. Assuming estimate (2.28) for α , one obtains

$$\frac{t_e}{t'_e} \rightarrow \alpha + 1 \approx 4.18 \quad \text{as } H \rightarrow 1. \quad (2.30)$$

The smallest discrepancy, in turn, is observed at low humidity. Solving (2.24) with $H = 0$ numerically, one obtains

$$\frac{t_e}{t'_e} \approx 2.95 \quad \text{for } H = 0. \quad (2.31)$$

Evidently, the isothermal model may yield a significant error in the whole range of H , which agrees broadly with conclusions of Sobac *et al.* (2015), Cossali & Tonini (2019) and Finneran (2021).

It is also interesting to calculate the time of evaporation of a typical ‘cough drop’ (through which COVID-19, flu, etc. are spread). The radii of such drops are typically between 0.5 and 16.0 μm (Yang *et al.* 2007). The largest drops from this range survive the longest – hence are the most dangerous when spreading the disease. The dependence t_e on T for this case is illustrated in figure 2.

3. The generalised Maxwell boundary condition as described by the diffuse-interface model

Unlike the classical model where the drop’s boundary has zero thickness, the DIM assumes the liquid and air to be separated by a finitely thin interface where the partial densities ρ_i vary smoothly. In this section, the DIM will be used to derive a model of evaporation that takes into account interfacial physics.

Mathematically, the interface should be viewed as an inner asymptotic region separating the outer regions of liquid and air. The DIM describes them all, but it can be simplified depending on the region to which it is applied.

First, one can take advantage of the fact that the temperature field associated with evaporative cooling occurs at a macroscopic scale (e.g. the drop’s radius). The interface, on the other hand, is microscopic – hence the temperature change in it is negligible, and it can be examined via the isothermal version of the DIM.

One should still account for the temperature dependence of the saturated vapour density (which was found to be important in the classical model) – but there is no need to use the full non-isothermal equations just for this. Instead, one can use the classical (macroscopic) model to calculate the temperature field and then use it to prescribe the correct value of $\rho_1^{(v.sat)}(T)$ near the drop. This workaround allows one to avoid cumbersome calculations associated with non-isothermality, yet obtain the same result.

One can also consider first a flat interface. Once the generalised Maxwell boundary condition has been derived for this case, the pressure difference due to the capillary effect can be incorporated readily into it.

3.1. The governing equations

Let z be the vertical Cartesian coordinate, and let w be the vertical velocity of the mixture as a whole. Introduce also the pressure p , and keep in mind that it is not an independent unknown: since the DIM assumes the fluid to be compressible, p is a known function of ρ_1 , ρ_2 and T (the equation of state). The shear and bulk viscosities – μ_s and μ_b , respectively – are also known functions of the same arguments.

Introduce the partial chemical potentials G_1 and G_2 . Their thermodynamic definition is given in [Appendix A.1](#) – but technically, one can simply treat them as given functions of ρ_1 , ρ_2 and T , such that

$$\frac{\partial p}{\partial \rho_j} = \sum_i \rho_i \frac{\partial G_i}{\partial \rho_j}. \tag{3.1}$$

In the low-density limit, p and G_i tend to their ideal gas approximations:

$$p \sim \sum_i R_i \rho_i, \quad G_i \sim R_i T \ln \rho_i \quad \text{as } \rho_i \rightarrow 0, \tag{3.2}$$

where R_i are the specific gas constants. For water and air, they are

$$R_1 = 461.53 \text{ m}^2 \text{ s}^{-2} \text{ K}^{-1}, \quad R_2 = 289.41 \text{ m}^2 \text{ s}^{-2} \text{ K}^{-1}, \tag{3.3a,b}$$

with R_2 calculated as the 70/30 weighted average of the gas constants of nitrogen and oxygen.

The DIM comprises the standard hydrodynamic equations involving an undetermined external force F_i , which will be identified later with the van der Waals force. Under the slow-flow (Stokes) approximation, a compressible isothermal binary fluid is governed by

$$\frac{\partial(\rho_1 + \rho_2)}{\partial t} + \frac{\partial[(\rho_1 + \rho_2)w]}{\partial z} = 0, \quad \frac{\partial \rho_1}{\partial t} + \frac{\partial(\rho_1 w + J)}{\partial z} = 0, \tag{3.4a,b}$$

$$\frac{\partial p}{\partial z} = \frac{\partial}{\partial z} \left(\eta \frac{\partial w}{\partial z} \right) + \rho_1 F_1 + \rho_2 F_2, \tag{3.5}$$

where the effective viscosity is

$$\eta = \frac{4\mu_s}{3} + \mu_b, \tag{3.6}$$

the diffusive flux is (e.g. Giovangigli 2021)

$$J = D \left(F_1 - \frac{\partial G_1}{\partial z} - F_2 + \frac{\partial G_2}{\partial z} \right), \quad (3.7)$$

and D is the diffusion coefficient.

To understand the relationship of D with the diffusivity \mathcal{D} from the classical model, assume that the fluid is diluted, so that the van der Waals force is negligible ($F_i = 0$) and the chemical potentials are given by (3.2). As a result, (3.7) reduces to

$$J \sim D \left(-\frac{R_1 T}{\rho_1} \frac{\partial \rho_1}{\partial z} + \frac{R_2 T}{\rho_2} \frac{\partial \rho_2}{\partial z} \right) \quad \text{as } \rho_i \rightarrow 0. \quad (3.8)$$

Let the vapour density be much smaller than that of the air ($\rho_1 \ll \rho_2$), but let their variations be comparable ($\partial \rho_1 / \partial z \sim \partial \rho_2 / \partial z$) – so that the above expression reduces to

$$J \sim -D \frac{R_1 T}{\rho_1} \frac{\partial \rho_1}{\partial z} \quad \text{as } \rho_i \rightarrow 0, \frac{\rho_1}{\rho_2} \rightarrow 0. \quad (3.9a,b)$$

A comparison of this expression with that for the classical diffusive flux yields

$$D \sim \frac{\rho_1}{R_1 T} \mathcal{D} \quad \text{as } \rho_i \rightarrow 0, \frac{\rho_1}{\rho_2} \rightarrow 0. \quad (3.10a,b)$$

For a dense mixture of fluids with comparable densities, D cannot be deduced from \mathcal{D} , but its dependence on (ρ_1, ρ_2, T) is supposed to be known from measurements.

For a multicomponent flow, the (one-dimensional) DIM approximation of the van der Waals force is (Benilov 2023a)

$$F_i = \sum_j K_{ij} \frac{\partial^3 \rho_j}{\partial z^3}, \quad (3.11)$$

where the Korteweg matrix K_{ij} characterises the interaction of molecules of components i and j (Newton's third law implies $K_{ij} = K_{ji}$). For the water–air combination, the Korteweg matrix is (see Appendix B)

$$\begin{bmatrix} K_{11} & K_{12} \\ K_{21} & K_{22} \end{bmatrix} = \begin{bmatrix} 1.87810 & 0.84978 \\ 0.84978 & 1.36880 \end{bmatrix} \times 10^{-17} \text{ m}^7 \text{ s}^{-2} \text{ kg}^{-1}. \quad (3.12)$$

Next, the pressure term in the momentum equation (3.5) can be rewritten conveniently in terms of the chemical potentials. Using identity (3.1), one obtains

$$\rho_1 \frac{\partial G_1}{\partial z} + \rho_2 \frac{\partial G_2}{\partial z} = \frac{\partial}{\partial z} \left(\eta \frac{\partial w}{\partial z} \right) + \rho_1 F_1 + \rho_2 F_2. \quad (3.13)$$

This equation and (3.7) can be viewed as a set of algebraic equations for $F_1 - \partial G_1 / \partial z$ and $F_2 - \partial G_2 / \partial z$. Solving these equations and recalling (3.11) for F_i , one obtains

$$\frac{\partial}{\partial z} \left(\sum_j K_{1j} \frac{\partial^2 \rho_j}{\partial z^2} - G_1 \right) = -\frac{1}{\rho_1 + \rho_2} \left[\frac{\partial}{\partial z} \left(\eta \frac{\partial w}{\partial z} \right) - \frac{\rho_2 J}{D} \right], \quad (3.14)$$

$$\frac{\partial}{\partial z} \left(\sum_j K_{2j} \frac{\partial^2 \rho_j}{\partial z^2} - G_2 \right) = -\frac{1}{\rho_1 + \rho_2} \left[\frac{\partial}{\partial z} \left(\eta \frac{\partial w}{\partial z} \right) + \frac{\rho_1 J}{D} \right]. \quad (3.15)$$

Equations (3.4) and (3.14)–(3.15) determine the unknowns ρ_1, ρ_2, w and J .

3.2. Non-dimensionalisation

In what follows, the vertical coordinate z will be non-dimensionalised on the interfacial thickness \bar{z} , which implies that distances should be measured from the current height $Z(t)$ of the interface. Mathematically, this corresponds to the change of variables $(z, t) \rightarrow (z_{new}, t_{new})$, where

$$z_{new} = z - Z(t), \quad t_{new} = t. \tag{3.16a,b}$$

Rewriting (3.4) in terms of the new variables, introducing $\dot{Z} = dZ/dt$, and omitting the subscript $_{new}$, one obtains

$$\frac{\partial(\rho_1 + \rho_2)}{\partial t} - \dot{Z} \frac{\partial \rho_2}{\partial z} + \frac{\partial[(\rho_1 + \rho_2)w]}{\partial z} = 0, \tag{3.17a}$$

$$\frac{\partial \rho_1}{\partial t} - \dot{Z} \frac{\partial \rho_1}{\partial z} + \frac{\partial(\rho_1 w + J)}{\partial z} = 0, \tag{3.17b}$$

whereas (3.14)–(3.15) remain the same as before.

There are three density scales in the problem – the liquid density, the air density and the saturated vapour density. Since they differ from each other by orders of magnitude, no single scaling of ρ_i would fit all the asymptotic regions. Still, a ‘generic’ non-dimensionalisation is needed, so let the density scale be that of air, $\bar{\rho} = \rho_2^{(a)}$.

Introducing the air viscosity scale $\bar{\eta}$, define the characteristic pressure and velocity by

$$\bar{p} = \frac{\bar{K}\bar{\rho}^2}{\bar{z}^2}, \quad \bar{w} = \frac{\bar{p}\bar{z}}{\bar{\eta}}, \tag{3.18a,b}$$

respectively. These choices for \bar{p} and \bar{w} correspond to an asymptotic regime where the pressure gradient, viscous stress and van der Waals force in the momentum equation (3.5) are of the same order.

Let \bar{t} be the evaporation time scale, and introduce

$$\varepsilon = \sqrt{\frac{\bar{z}}{\bar{t}\bar{w}}}. \tag{3.19}$$

This parameter is responsible for the quasi-steady approximation – hence it is expected to be small. Another important parameter is the advection/diffusion ratio, defined by

$$\delta = \frac{\bar{w}\bar{\rho}^2\bar{z}}{\bar{D}\bar{p}}, \tag{3.20}$$

where \bar{D} is the characteristic value of the diffusion coefficient D .

The following non-dimensional variables will be used:

$$\left. \begin{aligned} t_{nd} &= \frac{t}{\bar{t}}, & z_{nd} &= \frac{z}{\bar{z}}, \\ (\rho_i)_{nd} &= \frac{\rho_i}{\bar{\rho}}, & w_{nd} &= \frac{w}{\varepsilon\bar{w}}, & J_{nd} &= \frac{J}{\varepsilon\bar{\rho}\bar{w}}, & Z_{nd} &= \varepsilon Z, \\ \eta_{nd} &= \frac{\eta}{\bar{\eta}}, & D_{nd} &= \frac{D}{\bar{D}}, & (K_{ij})_{nd} &= \frac{K_{ij}}{\bar{K}}, & (G_i)_{nd} &= \frac{\bar{\rho}}{\bar{p}} G_i. \end{aligned} \right\} \tag{3.21a-j}$$

Rewriting (3.14)–(3.17) in terms of the non-dimensional variables, and omitting the subscript nd , one obtains

$$\frac{\partial}{\partial z} \left(\sum_j K_{1j} \frac{\partial^2 \rho_j}{\partial z^2} - G_1 \right) = -\frac{\varepsilon}{\rho_1 + \rho_2} \left[\frac{\partial}{\partial z} \left(\eta \frac{\partial w}{\partial z} \right) - \delta \frac{\rho_2 J}{D} \right], \quad (3.22)$$

$$\frac{\partial}{\partial z} \left(\sum_j K_{2j} \frac{\partial^2 \rho_j}{\partial z^2} - G_2 \right) = -\frac{\varepsilon}{\rho_1 + \rho_2} \left[\frac{\partial}{\partial z} \left(\eta \frac{\partial w}{\partial z} \right) + \delta \frac{\rho_1 J}{D} \right], \quad (3.23)$$

$$\varepsilon \frac{\partial(\rho_1 + \rho_2)}{\partial t} + \frac{\partial[(\rho_1 + \rho_2)(w - \dot{Z})]}{\partial z} = 0, \quad \varepsilon \frac{\partial \rho_1}{\partial t} + \frac{\partial[\rho_1(w - \dot{Z}) + J]}{\partial z} = 0. \quad (3.24a,b)$$

Evidently, the non-dimensional equations (3.22)–(3.24) can be transformed back to their dimensional versions, (3.14)–(3.17), by setting $\varepsilon = \delta = 1$. This shortcut will be used for re-dimensionalising the results obtained.

In what follows, one will need the non-dimensional versions of the low-density expressions (3.2) for p and G_i , and (3.10) for D . To obtain these, introduce

$$T_{nd} = \frac{\bar{R}\bar{\rho}}{\bar{p}} T, \quad \mathcal{D}_{nd} = \frac{\bar{\rho}^2}{\bar{D}\bar{p}} \mathcal{D}, \quad (3.25a,b)$$

where \bar{R} is the characteristic gas constant. Rewriting (3.2) and (3.10) in terms of T_{nd} and \mathcal{D}_{nd} , and omitting the subscript nd , one can verify that the dimensional and non-dimensional versions of these expressions look exactly the same.

3.3. Non-dimensional parameters

Let the characteristic value of the Korteweg parameter be that of dry air, $\bar{K} = K_{22}$, given by (3.12). The characteristic values of the rest of the parameters are given by (2.26)–(2.27) and (3.3).

For common fluids, the room temperature range is much closer to the freezing point than the critical point (e.g. for water, the former is 0 °C and the latter is 374 °C). In such cases, the interfacial thickness \bar{z} is comparable to the liquid's molecular size – i.e. several ångströms – as confirmed by both experiments (e.g. Verde *et al.* 2012; Pezzotti *et al.* 2017) and molecular simulations (e.g. Liu *et al.* 2005; Pezzotti *et al.* 2017; Dodia *et al.* 2019; and references therein).

To obtain an estimate of \bar{z} from within the DIM, one can use the particular case of an equilibrium interface. It is examined briefly in Appendix A.2 for the parameters specific to the water–air combination listed in Appendix B. The resulting profiles of the water and air densities are shown in figure 3, suggesting that the interfacial thickness is

$$\bar{z} \sim 5\text{Å}. \quad (3.26)$$

This value exceeds the water's molecular size by a factor of 2–3 (depending on how it is defined).

According to (3.19), ε depends on the time scale \bar{t} – which, in turn, depends on the evaporation rate – hence ε can be estimated only in application to a specific setting.

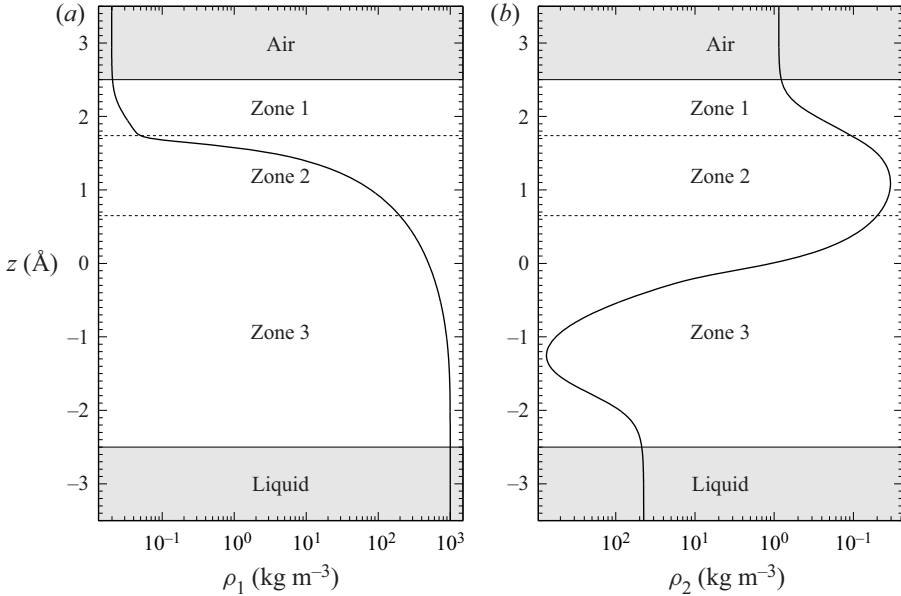


Figure 3. A schematic illustrating the asymptotic structure of the solution. The curves $\rho_1(z)$ and $\rho_2(z)$ describe a flat equilibrium water/air interface at $T = 25^\circ\text{C}$, computed using the DIM (see Appendix A.2). The scale of the horizontal axes in both panels is logarithmic. The height of the non-shaded region is used as an estimate for the interfacial thickness \bar{z} .

Assuming that to be a drop of radius r_d , one can use the continuity of the advective mass flux to relate \dot{r}_d to \bar{w} , we have

$$\dot{r}_d \rho_1^{(l)} \sim \bar{w} \rho_2^{(a)}. \tag{3.27}$$

Expressing \bar{w} from this equality, substituting it into definition (3.19) of ε , and approximating \dot{r}_d by r_d/\bar{t} , one obtains

$$\varepsilon = \sqrt{\frac{\rho_2^{(a)} \bar{z}}{\rho_1^{(l)} r_d}}. \tag{3.28}$$

Luckily, this expression no longer involves \bar{t} (which is unknown without further calculations). Then for r_d between $1\ \mu\text{m}$ and $1\ \text{nm}$, the above expression shows that ε is between 0.771×10^{-3} and 2.44×10^{-2} – i.e. it is small.

To estimate δ , estimate the diffusion coefficient by $\bar{D} = \mathcal{D} \bar{\rho}^2 / \bar{\rho}$, where \mathcal{D} is the standard vapour-in-the-air diffusivity. Then (3.18) and (3.20) yield

$$\delta = \frac{\bar{K} \bar{\rho}^2}{\mathcal{D} \bar{\eta}}. \tag{3.29}$$

Estimating the viscosity of air at 25°C using the results of Shang *et al.* (2019), one obtains

$$\bar{\eta} = 4.20 \times 10^{-5} \text{ kg m}^{-1} \text{ s}^{-1}, \tag{3.30}$$

for which (3.29) yields $\delta \approx 1.83 \times 10^{-8}$. Thus one can assume that

$$\delta \ll \varepsilon \ll 1. \tag{3.31}$$

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In addition to ϵ and δ , the problem involves two 'hidden' small parameters:

$$\epsilon^{(a/l)} = \frac{\rho_2^{(a)}}{\rho_1^{(l)}} \approx 1.19 \times 10^{-3}, \quad \epsilon^{(v/a)} = \frac{\rho_1^{(v.sat)}}{\rho_2^{(a)}} \approx 1.95 \times 10^{-2}. \quad (3.32a,b)$$

The presence of four independent small parameters makes the analysis awkward: they arise in various combinations, and each time, one has to check whether these are large, small or order-one. In particular, the following estimates will be needed:

$$\frac{\epsilon^2 \epsilon^{(a/l)} \epsilon^{(v/a)}}{\delta} \ll \frac{\epsilon^2 \epsilon^{(a/l)}}{\delta} \approx 3.84 \times 10^{-2} \ll 1, \quad (3.33)$$

$$\frac{\delta}{\epsilon^{(v/a)}} \approx 0.938 \times 10^{-6} \ll 1. \quad (3.34)$$

Thus all of the above parameter combinations will be assumed small.

3.4. *The boundary/matching conditions*

When applied to the interface (inner region), (3.22)–(3.24) need boundary conditions at large z . These are to be obtained via matching the inner solution to those in the liquid and air (outer regions). The former is trivial: since the liquid is homogeneous and at rest, the interfacial solution should satisfy

$$\rho_i \rightarrow \rho_i^{(l)} \quad \text{as } z \rightarrow -\infty, \quad (3.35)$$

$$w \rightarrow 0, \quad J \rightarrow 0 \quad \text{as } z \rightarrow -\infty. \quad (3.36)$$

Unlike the classical model where $\rho_i^{(l)}$ are external parameters, the DIM determines them as functions of the pressure and humidity of air. The DIM also describes dissolution of air in liquid – hence $\rho_2^{(l)} \neq 0$. This quantity is very small and thus unimportant both dynamically and thermodynamically.

The matching between the interface and air is less trivial: it can be carried out only after examining the large-distance behaviours of the inner solution and choosing the one that matches the outer solution. The properties of the latter are easy to guess: it should be such that $\rho_1 \ll \rho_2$ (small vapour concentration in the air), and the variations of ρ_1 and ρ_2 should be such that the total pressure is spatially uniform (e.g. Ricci & Rocca 1984; Mills & Coimbra 2016).

To verify that (3.22)–(3.24) admit such an asymptotic solution, let

$$\rho_i = \rho_i^{(+)} + \epsilon \delta \rho_i'^{(+)} z + O(\epsilon^2 \delta, \epsilon \delta^2), \quad w = w^{(+)} + O(\epsilon), \quad J = J^{(+)} + O(\epsilon), \quad (3.37a-c)$$

where $\rho_i^{(+)}$, $\rho_i'^{(+)}$, etc. depend on t (but not on z) and may involve the hidden small parameters $\epsilon^{(a/l)}$ and $\epsilon^{(v/a)}$. It turns out that only $\rho_i^{(+)}$ is independent in these expansions, with the other coefficients being expressible through $\rho_i^{(+)}$.

To find how $w^{(+)}$ and $J^{(+)}$ depend on $\rho_i^{(+)}$, observe that to leading order, (3.24) can be integrated. Using boundary conditions (3.35)–(3.36) to fix the constants of integration,

one obtains

$$w = -\dot{Z} \left(\frac{\rho_1^{(l)} + \rho_2^{(l)}}{\rho_1 + \rho_2} - 1 \right) + O(\varepsilon), \quad J = -\dot{Z} \frac{\rho_1^{(l)} \rho_2 - \rho_1 \rho_2^{(l)}}{\rho_1 + \rho_2} + O(\varepsilon), \quad (3.38a,b)$$

which entails

$$w^{(+)} = -\dot{Z} \left(\frac{\rho_1^{(l)} + \rho_2^{(l)}}{\rho_1^{(+)} + \rho_2^{(+)}} - 1 \right), \quad J^{(+)} = -\dot{Z} \frac{\rho_1^{(l)} \rho_2^{(+)} - \rho_1^{(+)} \rho_2^{(l)}}{\rho_1^{(+)} + \rho_2^{(+)}}. \quad (3.39a,b)$$

To calculate $\rho_i'^{(\pm)}$, one should substitute the expansions of w and J into (3.22)–(3.23). Since air is a diluted fluid, the effective viscosity does not depend on the density (e.g. Ferziger & Kaper 1972), i.e.

$$\eta \sim \eta_0 \quad \text{as } \rho_i \rightarrow 0, \quad (3.40)$$

where η_0 depends only on T . This property implies that when expressions (3.38) are substituted into (3.22)–(3.23), the linear term in the expansion for w cancels out. The quadratic term does not, but its contribution is much smaller than that of $J^{(0)}$ (due to estimates (3.33)), and to leading order, one obtains the following algebraic equations for $\rho_i'^{(\pm)}$:

$$\sum_i \left(\frac{\partial G_1}{\partial \rho_i} \right)^{(+)} \rho_i'^{(+)} = -\frac{\rho_2^{(+)} J^{(+)}}{D^{(+)}(\rho_1^{(+)} + \rho_2^{(+)})}, \quad (3.41)$$

$$\sum_i \left(\frac{\partial G_2}{\partial \rho_i} \right)^{(+)} \rho_i'^{(+)} = \frac{\rho_1^{(+)} J^{(+)}}{D^{(+)}(\rho_1^{(+)} + \rho_2^{(+)})}, \quad (3.42)$$

where $D^{(+)} = D(\rho_1^{(+)}, \rho_2^{(+)}, T)$ and

$$\left(\frac{\partial G_j}{\partial \rho_i} \right)^{(+)} = \frac{\partial G_j(\rho_1^{(+)}, \rho_2^{(+)}, T)}{\partial \rho_i^{(+)}}. \quad (3.43)$$

Given the smallness of the air-to-liquid and vapour-to-air density ratios (3.32), (3.41)–(3.42) can be simplified. Using the diluted-fluid expressions (3.2) and (3.10) for G_i and D , respectively, and keeping the leading order only, one obtains

$$\rho_1'^{(+)} = \frac{\rho_1^{(l)} \dot{Z}}{D}, \quad (3.44)$$

$$\rho_2'^{(+)} = -\rho_1'^{(+)} \frac{R_1}{R_2}. \quad (3.45)$$

Equality (3.44) is the desired matching condition; it can also be viewed as the relationship of the speed \dot{Z} of the interface to the vapour flux $D\rho'^{(\pm)}$, whereas (3.45) is the standard condition of constant pressure for diffusion in diluted fluids. As mentioned before, both conditions are obvious physically, but it is still important to verify that they are intrinsic to the DIM formalism.

3.5. The interfacial asymptotic region

Three asymptotic zones exist within the inner (interfacial) region,

$$\left. \begin{aligned} \text{Zone 1 (mostly air):} & \quad \rho_1 \ll \rho_2 \ll \rho^{(l)}, \\ \text{Zone 2 (air-vapour mixture):} & \quad \rho_1 \sim \rho_2 \ll \rho^{(l)}, \\ \text{Zone 3 (mostly liquid):} & \quad \rho_2 \ll \rho_1 \sim \rho^{(l)}. \end{aligned} \right\} \quad (3.46)$$

Note that Zone 1 is not part of the air region: in the latter, the air density is nearly uniform, $\rho_2 \approx \rho_2^{(a)}$, whereas in the former, variations of ρ_2 are comparable to $\rho_2^{(a)}$.

A schematic illustrating the asymptotic structure of the problem can be found in [figure 3](#) (which is computed for the equilibrium interface, i.e. that where the relative humidity of air is $H = 1$). Its most counter-intuitive features are the local maximum and minimum of the air density $\rho_2(z)$ in Zone 2. Note that they both disappear in the limit $K_{12} \rightarrow 0$ (see [figure 11](#) of [Benilov 2023a](#)). One can thus assume that the maximum of ρ_2 is created by the van der Waals force exerted by the bulk of the liquid by attracting a certain amount of air towards the interface. The existence of the minimum of ρ_2 is harder to understand, but it is dynamically unimportant anyway. The air density is extremely small there – even smaller than that of the air dissolved in the liquid – and they both have a minuscule effect on the dynamics of the rest of the fluid.

It turns out that only Zone 1 contributes significantly to the evaporation rate, whereas the contributions of Zones 2–3 are negligible. To understand why, recall that the fluid density in the latter two zones exceeds that of the vapour component of air by an order of magnitude. As a result, Zones 2–3 are close to equilibrium (the fact that the ‘outside’ air is undersaturated cannot perturb them too much). Thus Zone 1 is the only region out of equilibrium – hence it is the only place where irreversible processes, such as evaporation, can occur.

The calculations associated with Zones 2–3 can be avoided via a certain workaround developed by [Benilov \(2022b, 2023b\)](#) for evaporation of pure fluids. The workaround involves the following two steps.

- (i) Equations [\(3.22\)–\(3.24\)](#) can be rearranged in such a way that the velocity \dot{Z} of the interface is expressed in terms of the fluid’s known parameters (relative humidity, etc.) and a certain integral of the solution.
- (ii) The contribution of Zone 1 to this integral exceeds the contributions of Zones 2–3 by an order of magnitude. Thus to evaluate this integral asymptotically, it is sufficient to find the solution only in Zone 1.

3.5.1. Calculation of \dot{Z}

Recall that matching conditions [\(3.44\)–\(3.45\)](#) emerged from the governing equations at the order of $O(\varepsilon\delta)$. Finding \dot{Z} does not require going that far, as the terms in [\(3.22\)–\(3.23\)](#) that involve \dot{Z} are $O(\varepsilon)$. The terms $O(\delta)$ in these equations can also be omitted (as suggested by the estimates in [§ 3.3](#)).

Substitute expressions [\(3.38\)](#) for w and J into [\(3.22\)–\(3.23\)](#), and, keeping the terms up to $O(\varepsilon)$, obtain

$$\frac{\partial}{\partial z} \left(\sum_j K_{1j} \frac{\partial^2 \rho_j}{\partial z^2} - G_1 \right) = \frac{\varepsilon \dot{Z}}{\rho_1 + \rho_2} \frac{\partial}{\partial z} \left[\eta \frac{\partial}{\partial z} \left(\frac{\rho_1^{(l)} + \rho_2^{(l)}}{\rho_1 + \rho_2} \right) \right], \quad (3.47)$$

$$\frac{\partial}{\partial z} \left(\sum_j K_{2j} \frac{\partial^2 \rho_j}{\partial z^2} - G_2 \right) = \frac{\varepsilon \dot{Z}}{\rho_1 + \rho_2} \frac{\partial}{\partial z} \left[\eta \frac{\partial}{\partial z} \left(\frac{\rho_1^{(l)} + \rho_2^{(l)}}{\rho_1 + \rho_2} \right) \right]. \quad (3.48)$$

Keeping the same accuracy in boundary condition (3.37), one obtains

$$\rho_i \rightarrow \rho_i^{(+)} \quad \text{as } z \rightarrow +\infty. \quad (3.49)$$

Now, integrate (3.47) and (3.48) with respect to z from $-\infty$ to $+\infty$. Recalling boundary conditions (3.35) and (3.49), one obtains

$$G_i^{(l)} - G_i^{(+)} = -\varepsilon \dot{Z} A, \quad (3.50)$$

where

$$A = \int_{-\infty}^{+\infty} \frac{1}{\rho_1 + \rho_2} \frac{\partial}{\partial z} \left[\frac{(\rho_1^{(l)} + \rho_2^{(l)}) \eta}{(\rho_1 + \rho_2)^2} \frac{\partial(\rho_1 + \rho_2)}{\partial z} \right] dz. \quad (3.51)$$

Next, consider the combination

$$\int_{-\infty}^{+\infty} [\rho_1 \times (3.47) + \rho_2 \times (3.48)] dz. \quad (3.52)$$

Integrating by parts the terms involving the third derivatives, take into account boundary conditions (3.35) and (3.49), and recall identity (3.1) to obtain

$$p^{(l)} - p^{(+)} = 0. \quad (3.53)$$

Thus the pressure in the liquid coincides with that in the air (as expected physically).

Equalities (3.50) and (3.53) relate the (unknown) liquid densities $\rho_i^{(l)}$ and the velocity \dot{Z} of the interface to the air parameters (marked with $(+)$). In addition, \dot{Z} is also related to the air parameters by equality (3.44). If two of these three equalities are used to eliminate $\rho_i^{(l)}$ and \dot{Z} , then the remaining one will inter-relate the air parameters and thus turn into the generalised Maxwell boundary condition (GMBC).

To eliminate $\rho_i^{(l)}$ from equalities (3.50) and (3.53), recall that equations of state of liquids at normal conditions are typically such that even a minuscule change of the density gives rise to a huge change of the pressure. This means effectively that $\rho_i^{(l)}$ is close to its saturated value, $\rho_i^{(l)} \approx \rho_i^{(l.sat)}$ – so that (3.50) becomes

$$G_i^{(l.sat)} - G_i^{(+)} = -\varepsilon \dot{Z} A. \quad (3.54)$$

In the state of saturation, the chemical potentials of the liquid and vapour are equal (see Appendix A.2) – hence one can change in the above equality $G_1^{(l.sat)} \rightarrow G_1^{(v.sat)}$. Then, using for both $G_1^{(v.sat)}$ and $G_1^{(+)}$ the diluted-fluid expression (3.2), one obtains

$$\rho_1^{(l.sat)} R_1 T \ln \frac{\rho_1^{(+)}}{\rho_1^{(v.sat)}} = \varepsilon \mathcal{D} \rho_1'^{(+)} A. \quad (3.55)$$

This is, essentially, the GMBC.

With A calculated (see the next subsection), equality (3.55) inter-relates the near-interface vapour density $\rho_1^{(+)}$ and its gradient $\rho_1'^{(+)}$. If $\varepsilon = 0$, then (3.55) yields

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$\rho_1^{(+)} = \rho_1^{(v.sat)}$, which is the classical Maxwell boundary condition. If, however, ε is small but not zero, then $\rho_1^{(+)}$ can differ significantly from $\rho_1^{(v.sat)}$ – because A can actually be large, so that $\varepsilon A \sim 1$. This does not violate the employed asymptotic approach: it requires only $\varepsilon \ll 1$, which makes the time derivatives in (3.24) small and thus justifies the quasi-steady approximation.

3.5.2. Zone 1 of the interfacial asymptotic region

The coefficient A arises in all problems where the DIM is applied to evaporation or condensation – but, so far, it has been calculated only for pure fluids and the case where the relative humidity is close to unity (Benilov 2020, 2022a,b, 2023b). The calculation of A for a binary mixture and arbitrary humidity is more complex technically, but the underlying idea is the same.

It is based on the observation that the main contribution to (3.51) comes from the zone where $\rho_1 + \rho_2$ is small, but its derivatives are relatively large – i.e. near the air region, but not in it. This means effectively that (3.51) can be calculated asymptotically by considering Zone 1 only (see its definition at the beginning of § 3.5).

Before presenting the scaling of Zone 1, it is instructive to apply the ‘obvious’ approximations – i.e. take advantage of the diluted-fluid expressions (3.2), (3.10) and (3.40), plus neglect ρ_1 and $\rho_2^{(l)}$ if they appear next to ρ_2 and $\rho_1^{(l)}$, respectively. As a result, (3.47)–(3.48) and (3.51) become

$$\frac{\partial}{\partial z} \left(K_{12} \frac{\partial^2 \rho_2}{\partial z^2} - R_1 T \ln \rho_1 \right) = \varepsilon \rho_1^{(l)} \dot{Z} \frac{\eta_0}{\rho_2} \frac{\partial^2}{\partial z^2} \left(\frac{1}{\rho_2} \right), \tag{3.56}$$

$$\frac{\partial}{\partial z} \left(K_{22} \frac{\partial^2 \rho_2}{\partial z^2} - R_2 T \ln \rho_2 \right) = \varepsilon \rho_1^{(l)} \dot{Z} \frac{\eta_0}{\rho_2} \frac{\partial^2}{\partial z^2} \left(\frac{1}{\rho_2} \right), \tag{3.57}$$

$$A = \eta_0 \rho_1^{(l)} \int_{-\infty}^{+\infty} \frac{1}{\rho_2} \frac{\partial}{\partial z} \left(\frac{1}{\rho_2^2} \frac{\partial \rho_2}{\partial z} \right) dz. \tag{3.58}$$

The most general asymptotic limit is when the three terms in (3.57) are of the same order, which implies the rescaling

$$z = z_0 + \left(\frac{K_{22} \rho_2^{(+)}}{R_2 T} \right)^{1/2} z_{new}, \quad \rho_i = \rho_i^{(+)} (\rho_i)_{new}, \quad \dot{Z} = -\frac{E_0}{\varepsilon \rho_1^{(l)}} \xi, \tag{3.59a-c}$$

$$A = \frac{\rho_1^{(l)} R_2 T}{E_0} A_{new}, \tag{3.60}$$

where z_0 is the ‘location’ of Zone 1, E_0 is given by (1.3) of the Introduction, and the minus in the expression for \dot{Z} is introduced to make ξ positive (recall that evaporation corresponds to $\dot{Z} < 0$). Since the problem under consideration is translationally invariant, the actual value of z_0 does not feature in, and is not needed for, any leading-order results. One does not need (3.56) either, because now A depends only on ρ_2 .

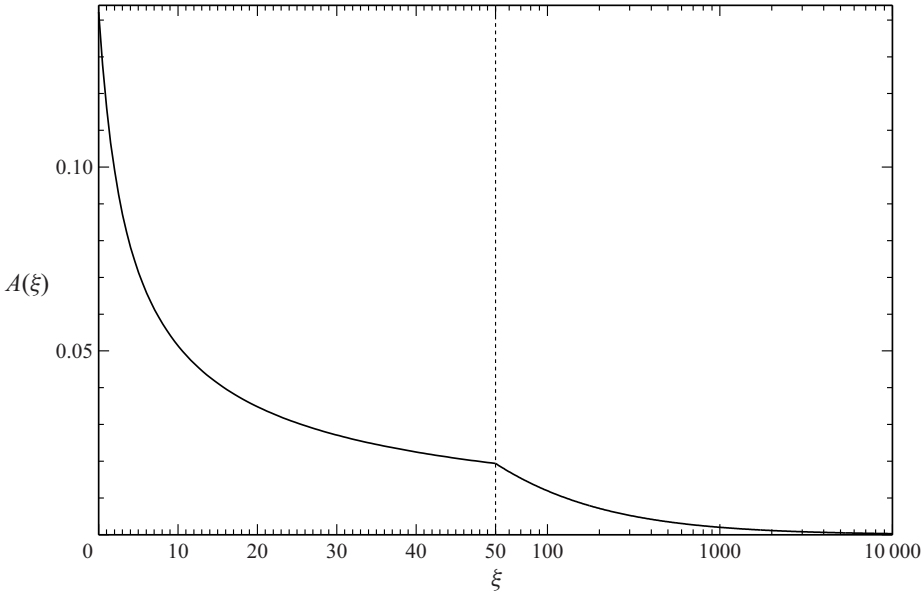


Figure 4. The function $A(\xi)$. Note that in the right-hand half of the figure, the scale of ξ is logarithmic.

Substitution of (3.59)–(3.60) into (3.57), boundary condition (3.49), and (3.58) yield (with the subscript *new* omitted)

$$\frac{\partial}{\partial z} \left(\frac{\partial^2 \rho_2}{\partial z^2} - \ln \rho_2 \right) = -\frac{\xi}{\rho_2} \frac{\partial^2}{\partial z^2} \left(\frac{1}{\rho_2} \right), \tag{3.61}$$

$$\rho_2 \rightarrow 1 \quad \text{as } z \rightarrow +\infty, \tag{3.62}$$

$$A = \int_{-\infty}^{+\infty} \frac{1}{\rho_2} \frac{\partial}{\partial z} \left(\frac{1}{\rho_2^2} \frac{\partial \rho_2}{\partial z} \right) dz. \tag{3.63}$$

As shown in Appendix C, boundary condition (3.62) is sufficient to uniquely fix the solution of (3.61), making it unnecessary to examine the other asymptotic zones of the interfacial region.

It can be derived readily from (3.61) that

$$\rho_2 \sim z^2 \ln(-z) \quad \text{as } z \rightarrow -\infty. \tag{3.64}$$

The growth of $\rho_2(z)$ as $z \rightarrow -\infty$ in Zone 1 agrees with the fact that this function has a local maximum in Zone 2 (see figure 3). Asymptotics (3.64) also guarantees that (3.63) converges at the lower limit.

The function $A(\xi)$ was computed numerically using the algorithm described in § C.2; the result is plotted in figure 4. For practical use, however, it is more convenient to use the approximation formula (1.4) (for more details, see § C.3).

3.5.3. Summarising the GMBC for flat interfaces

Recall that the dimensional variables can be recovered by setting $\varepsilon = \delta = 1$. Then equalities (3.44) and (3.59) yield a dimensional relationship of the near-interface air

density $\rho_1^{(+)}$ and its gradient $\rho_1'^{(\+)}$,

$$\xi = \frac{-\mathcal{D}\rho_1'^{(\+)}}{E_0}, \tag{3.65}$$

and (3.55) yields

$$\rho_1^{(+)} = \rho_1^{(v.sat)} \exp \left[-\frac{R_2}{R_1} \xi A(\xi) \right]. \tag{3.66}$$

Equalities (3.65) and (3.66) inter-relate the near-surface vapour density $\rho_1^{(+)}$ and its normal derivative $\rho_1'^{(\+)}$, hence they constitute the generalised Maxwell boundary condition (for flat interfaces), as required.

3.5.4. Curved interfaces

Interfacial curvature influences evaporation via the so-called Kelvin effect, i.e. by changing the effective saturated vapour pressure near the liquid's (curved) surface. This effect can be incorporated readily into the analysis under the assumption that the curvature radius is much larger than the interfacial thickness. This has been done, and the result has turned out to be physically obvious: the Kelvin effect adds a new term to the pressure condition (3.53),

$$p^{(l)} - p^{(+)} = \varepsilon C \sigma, \tag{3.67}$$

where C is the interfacial curvature, and σ is the surface tension. (The DIM expression for σ is given by (B8) of Appendix B.)

The extra term in the pressure equation entails an extra term in the GMBC: instead of (3.66), it becomes (1.1) of the Introduction, whereas definition (3.65) of the relative evaporation rate ξ coincides with (1.2) if one sets $\rho_1'^{(\+)} = (\mathbf{n} \cdot \nabla \rho_1)^{(\+)}$.

4. Evaporation of drops as described by the generalised model

To compare the generalised model to the classical one, the former will be applied to evaporation of floating spherical drops. To do so, one needs to replace the classical boundary condition (2.10) with its generalised version (1.1), and leave the rest of the governing set (2.3)–(2.13) the same as before. The ensuing algebra is also the same as before, and one eventually obtains the following differential equation for $r_d(t)$:

$$\underbrace{\frac{\rho_1^{(l.sat)}}{\mathcal{D}\rho_1^{(v.sat)(a)}} \dot{r}_d r_d}_{\text{diffusion}} + \exp \left[\underbrace{\frac{T_A \rho_1^{(l.sat)} \Delta}{T^{(a)} 2\kappa^{(a)}} \dot{r}_d r_d}_{\text{non-isothermality}} - \underbrace{\frac{R_2}{R_1} \xi A(\xi)}_{\text{interfacial effects}} + \underbrace{\frac{2\sigma}{\rho_1^{(l.sat)} R_1 T^{(a)} r_d}}_{\text{Kelvin effect}} \right] = H, \tag{4.1}$$

where

$$\xi = \frac{-\rho_1^{(l.sat)} \dot{r}_d}{E_0} \tag{4.2}$$

is the ratio of the evaporative flux to the interfacial emission capacity (the latter given by (1.3)).

Equations (4.1)–(4.2) generalise the corresponding classical result (2.21). The term involving $A(\xi)$ in (4.1) describes emission of vapour by the interface, and the term involving σ gives the shift of the effective saturation pressure near a curved surface. The remaining two terms on the left-hand side of (4.1) are the same as those in its classical counterpart (2.21).

The classical and modified models – described by (2.21) and (4.1), respectively – are compared in figures 5(a,b). For figure 5(a), both were treated as algebraic equations for the non-dimensional evaporation flux ξ with the drop’s radius r_d being a parameter – so that the result is the dependence ξ versus r_d . Figure 5(b), in turn, shows the dependence of the time of full evaporation on the drop’s initial radius.

The following features of these figures should be observed.

- (i) The (classical) d^2 law states that $t_e \sim r_d^2$, which is why curve (M) in figure 5(b) is a straight line with slope 2. In figure 5(a), curve (M) is also a straight line, but its slope is -1 (which agrees with the corresponding result of the classical model).
- (ii) The shaded areas in figures 5(a,b) mark the region $\xi < 1$, where the two models are supposed to yield similar results – and indeed, they do. The value of r_d corresponding to $\xi = 1$ is $r_d \approx 2 \mu\text{m}$.
- (iii) For smaller r_d , curves (M) and (B) begin to diverge. The difference between the classical and generalised results peaks when the drop’s radius is close to, and slightly less than, $r_d = 10 \text{ nm}$.

A similar effect is observed in models resolving or parametrising the Knudsen layer (Long *et al.* 1996; Haut & Colinet 2005; Landry *et al.* 2007; Jakubczyk *et al.* 2012; Hołyst *et al.* 2013, 2017; Rana *et al.* 2018, 2019; Zhao & Nadal 2023).

- (iv) For drops with $r_d \lesssim 10 \text{ nm}$, the Kelvin effect ‘kicks in’: it increases the effective saturated pressure and thus accelerates the vapour emission by the interface – as a result, curves (M) and (B) re-converge.

To verify this explanation, the four terms of (4.1) have been computed individually and plotted in figure 5(c). Evidently, evaporation of small drops is governed by the balance of interfacial effects and the Kelvin effect. Non-isothermality is less important, whereas the diffusion of vapour in the air is unimportant.

- (v) A general criterion of importance of the van der Waals force relative to diffusion can be obtained by estimating the characteristic ratio B of the second term in the square brackets in (4.1) to the first term. Recalling (4.2), one obtains

$$B = \frac{R_2 \bar{A} T^{(a)2} \kappa^{(a)}}{R_1 r_d E_0 T_A \Delta}, \tag{4.3}$$

where \bar{A} is a characteristic value of the function $A(\xi)$. The van der Waals force is important when $B \gtrsim 1$, which amounts to

$$r_d \lesssim \frac{R_2 \bar{A} T^{(a)2} \kappa^{(a)}}{R_1 E_0 T_A \Delta}. \tag{4.4}$$

For the parameters of water at 25°C and with $\bar{A} = A(0) \approx 0.14219$, one obtains $r_d \lesssim 0.4 \mu\text{m}$, which agrees qualitatively with the fact that curves (d) and (i) in figure 5(c) intersect at $r_d \approx 1 \mu\text{m}$.

- (vi) Figure 5(c) also suggests that for large drops (say, $r_d \gtrsim 0.1 \mu\text{m}$), the Kelvin effect is negligible, which agrees broadly with the estimate of Sobac *et al.* (2015).

It is worth emphasising, however, that this conclusion applies only to large floating drops, whereas large sessile drops can be still influenced by the Kelvin effect via the

Maxwell's hypothesis of saturation near liquid/air interface

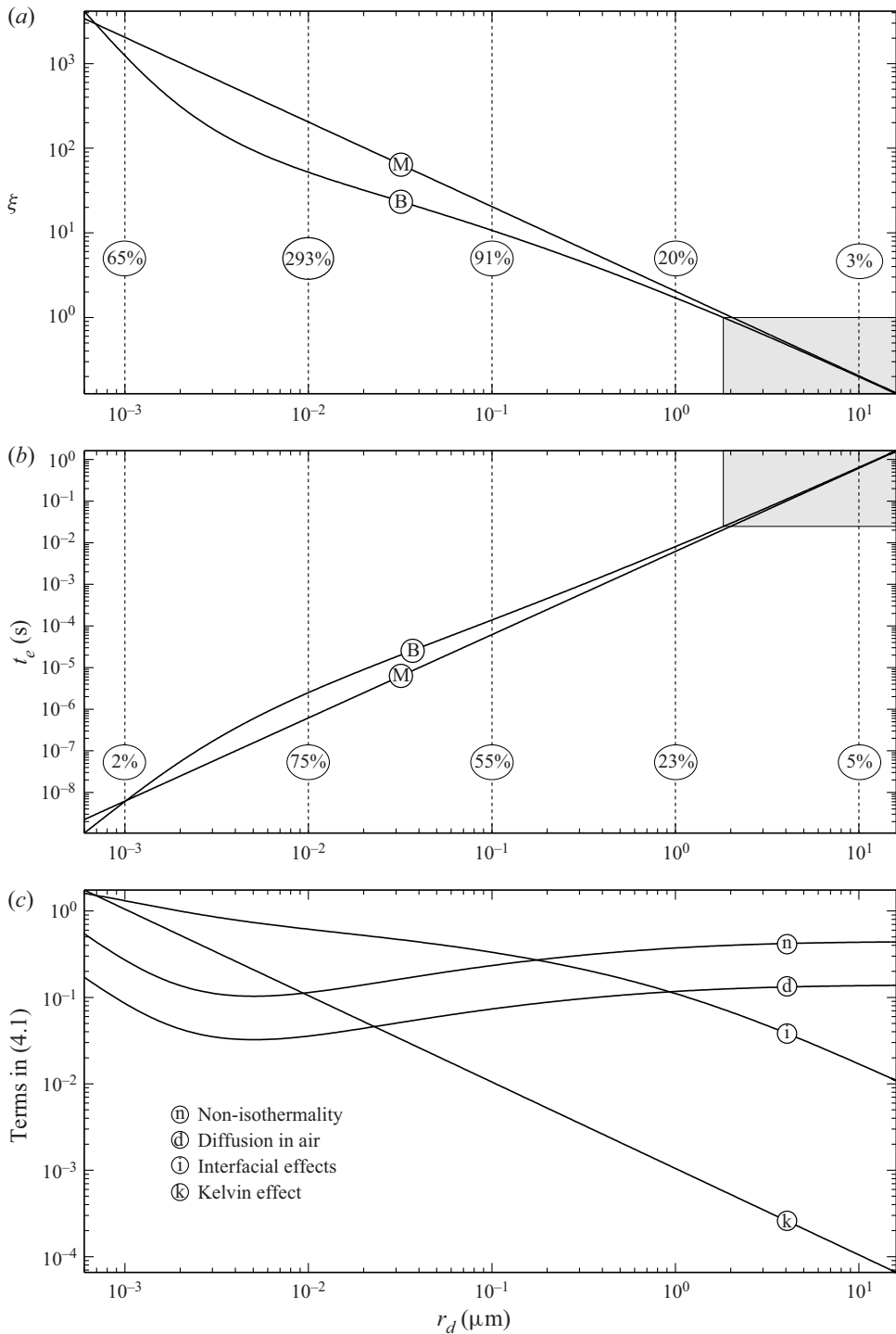


Figure 5. Characteristics of an evaporating spherical drop versus its radius, for $T = 25^\circ\text{C}$ and relative humidity $H = 50\%$. Curves (M) and (B) are computed using Maxwell's classical model and its generalised version, respectively. The numbers in the ellipses show the relative deviations of curve (M) from curve (B). (a) The evaporative flux normalised by the emission capacity of the interface (where ξ is defined by (4.2)). (b) The time of evaporation. The shaded areas of (a) and (b) correspond to $\xi < 1$, which is where the classical model is supposed to work. (c) The four terms (effects) in (4.1).

highly curved part of their surface near the contact line (e.g. Deegan *et al.* 2000; Dunn *et al.* 2009; Eggers & Pismen 2010; Colinet & Rednikov 2011; Rednikov & Colinet 2013, 2019; Morris 2014; Stauber *et al.* 2014, 2015; Janeček *et al.* 2015; Sáenz *et al.* 2015; Saxton *et al.* 2016, 2017; Wray, Duffy & Wilson 2019; Williams *et al.* 2020).

- (vii) Overall, figures 5(a,b) should be viewed as an argument in favour of the GMBC at scales $\lesssim 1 \mu\text{m}$, whereas figure 5(c) provides a physical explanation as to why.

5. Concluding remarks

Since the main result of this paper has been summarised in the Introduction, it only remains to comment on further applications of the proposed model and its connections with other models of evaporation.

- (i) Apart from drops, the generalised Maxwell boundary condition can be applied to menisci – and if their radii are small, then the GMBC should produce results more accurate than those of the classical model. This should be important for diffusion of fluids in porous media with small pores.

Note also that interfacial effects are particularly strong for concave menisci, in which case the Kelvin effect works with the interfacial effects, not against them. This should make the difference between the predictions of the classical and generalised models larger than that for convex menisci and drops.

- (ii) For flat and cylindrical interfaces, the results obtained via the two models converge with time. In both cases, the diffusion equation does not have a one-dimensional solution describing a steady flux of vapour towards infinity, which effectively means that the throughput of air is zero. As a result, after a sufficiently long time, the air near the interface gets almost saturated.

At finite times, however, the classical and generalised models yield different results even for cylindrical and flat interfaces.

- (iii) A sessile or pendant drop – especially that with moving contact lines – evaporates differently from floating ones. Its surface is highly curved near the contact lines, changing the local evaporation rate. To account for this effect is not a straightforward task, however, as the flow inside the drop is not one-dimensional in this case.

One might hope that the generalised model might explain the problem of abnormally small Navier slip arising for several liquids, including water (Podgorski, Flesselles & Limat 2001; Winkels *et al.* 2011; Puthenveetil, Senthilkumar & Hopfinger 2013; Benilov & Benilov 2015).

- (iv) All of the problems listed above are easier to solve using the quasi-static approximation. One should keep in mind, however, that Finneran *et al.* (2021) compared that with the transient solution of the governing equations (for the classical model) and observed that at high pressure, the quasi-steady approximation overpredicts the droplet lifetime by up to 80%.

One might conjecture that the difference between the transient and quasi-steady approaches is due to the fact that the latter implies that the mass of evaporated liquid is infinite. Indeed, the steady solution of the spherically symmetric diffusion equation for vapour is $\sim 1/r$ – hence the integral of this solution over the region between the drop's surface and infinity diverges.

Thus for the quasi-steady solution to establish itself around the drop, a significant part of the drop should have evaporated already. This applies to both classical and generalised models.

- (v) If the drop's radius is small, but the Kelvin effect is nevertheless neglected, the generalised model – i.e. (4.1) with $\sigma = 0$ – predicts that the time of a drop's evaporation is proportional to its radius. A similar behaviour occurs due to the temperature jump associated with the Knudsen layer (e.g. Rana *et al.* 2019; Zhao & Nadal 2023), but this seems to be just a coincidence. Indeed, the DIM does not describe temperature jumps (which are a purely kinetic effect (Bond & Struchtrup 2004; Persad & Ward 2016; Rana *et al.* 2018)), whereas the existing kinetic models do not include the van der Waals force (which is vital for Zone 1 of the present analysis).

Yet the Knudsen layer and Zone 1 are both located at the outskirts of the interface, out of equilibrium and affected by irreversible processes: the former by collisions, and the latter by viscosity (which is the hydrodynamic analogue of collisions). All this suggests that Zone 1 is the DIM equivalent of the Knudsen layer. If this is indeed the case, then the main conclusion of this paper is that the van der Waals force is one of the driving effects in the Knudsen layer and thus should be accounted for.

This conjecture can be verified using the Enskog–Vlasov kinetic equation; it includes the van der Waals force (just like the DIM) and can handle temperature jumps (like the Boltzmann equation). Since both effects work the same way – to slow evaporation down – their joint effect should be stronger than predicted by the DIM and existing models of the Knudsen layer separately.

- (vi) Despite the fact that the DIM cannot handle temperature jumps, it is still closely related to the Enskog–Vlasov equation, with the former being the hydrodynamic approximation of the latter (Giovangigli 2020, 2021). The asymptotic structure of the solution of the two models should be similar, and the results obtained in the present paper should make the analysis of the Enskog–Vlasov equation much easier.
- (vii) Note that, strictly speaking, the surface tension depends on the drop's curvature (Tolman's effect), which can have a profound influence on, say, cavitation (Menzl *et al.* 2016; Magaletti, Gallo & Casciola 2021; Aasen *et al.* 2023; Lamas *et al.* 2023). Since cavitating bubbles are created with zero radii, Tolman's length – no matter how small – may affect the whole 'trajectory' of their growth. Evaporation appears to be different, however; since Tolman's length for, say, water under normal conditions is less than 1 Å (e.g. Wilhelmson, Bedeaux & Reguera 2015; Burian *et al.* 2017), it is much smaller than typical radii of evaporating drops, and the whole effect is likely to be negligible.

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Appendix A. The thermodynamics incorporated into the DIM

A.1. The definition of the chemical potential G_i

Consider an N -component compressible non-ideal fluid, characterised by the temperature T and partial densities ρ_i , where $i = 1, \dots, N$. The fluid's thermodynamic properties are fully described by the internal energy $e(\rho_1, \dots, \rho_N, T)$ and entropy $s(\rho_1, \dots, \rho_N, T)$ – both specific, i.e. per unit mass. Note that e and s are not fully arbitrary, but should satisfy the fundamental thermodynamic relation, which can be written in the form (see Appendix

A of Benilov 2023a)

$$\frac{\partial e}{\partial T} = T \frac{\partial s}{\partial T}. \tag{A1}$$

The dependence of the fluid pressure p on $(\rho_1, \dots, \rho_N, T)$, or the equation of state, is given by (e.g. Giovangigli & Matuszewski 2013)

$$p = \rho \sum_i \rho_i \left(\frac{\partial e}{\partial \rho_i} - T \frac{\partial s}{\partial \rho_i} \right), \tag{A2}$$

where

$$\rho = \sum_i \rho_i \tag{A3}$$

is the total density. The partial chemical potentials, in turn, are given by

$$G_i = \frac{\partial(\rho e)}{\partial \rho_i} - T \frac{\partial(\rho s)}{\partial \rho_i}. \tag{A4}$$

Using (A1)–(A4), one can verify readily identity (3.1).

A.2. The Maxwell construction (conditions of saturation)

Consider a flat interface in a multicomponent fluid in the state of equilibrium. This implies that there should be no flow and fluxes ($w = J_i = 0$), and no dependence on t , so that (3.14)–(3.15) yield

$$\frac{d}{dz} \left(\sum_j K_{ij} \frac{d^2 \rho_j}{dz^2} - G_i \right) = 0. \tag{A5}$$

One should also impose the following boundary conditions:

$$\rho_i \rightarrow \rho_i^{(l.sat)} \quad \text{as } z \rightarrow -\infty, \tag{A6}$$

$$\rho_i \rightarrow \rho_i^{(v.sat)} \quad \text{as } z \rightarrow +\infty, \tag{A7}$$

where $\rho_i^{(l.sat)}$ and $\rho_i^{(v.sat)}$ are the densities of saturated liquid and vapour, respectively.

Integrating (A5) with respect to z , and taking into account conditions (A6)–(A7), one obtains

$$G_i^{(l.sat)} = G_i^{(v.sat)}. \tag{A8}$$

Then consider

$$\int_{-\infty}^{+\infty} \sum_i \rho_i \times (\text{A5}) \, dz. \tag{A9}$$

Integrating the resulting equality by parts, and keeping in mind that

$$\sum_{i,j} K_{ij} \frac{d\rho_i}{dz} \frac{d^2 \rho_j}{dz^2} \, dz = \frac{1}{2} \frac{d}{dz} \sum_{i,j} K_{ij} \frac{d\rho_i}{dz} \frac{d\rho_j}{dz} \, dz, \tag{A10}$$

one can use boundary conditions (A6)–(A7) to obtain

$$-\sum_i \rho_i^{(v.sat)} G_i^{(v.sat)} + \sum_i \rho_i^{(l.sat)} G_i^{(l.sat)} + \int_{-\infty}^{+\infty} \sum_i \frac{d\rho_i}{dz} G_i dz = 0. \quad (A11)$$

Observe that identity (3.1) implies that

$$G_i = \frac{\partial}{\partial \rho_i} \left(\sum_j \rho_j G_j - p \right). \quad (A12)$$

This equality helps one to integrate the last term of (A11) and obtain

$$p^{(l.sat)} = p^{(v.sat)}. \quad (A13)$$

Physically, equalities (A8) and (A13) reflect the famous Maxwell construction (Maxwell 1875), which inter-relates the partial densities of the components in vapour and liquid. One should also require

$$p^{(v.sat)} = p_A, \quad (A14)$$

where p_A is the atmospheric pressure.

An example of a numerical solution of boundary-value problem (A5)–(A7) in application to a water/air interface (with the chemical potentials G_i described in the next appendix) is shown in figure 3.

Appendix B. The parameters of the DIM

Benilov (2023a) proposed to use the DIM with the so-called Enskog–Vlasov equation of state. In application to a binary fluid with one diluted component, it amounts to the following expressions for the chemical potentials:

$$G_1 = T \left[R_1 \ln \rho_1 + \Theta(\rho_1) + (\rho_1 + \rho_2) \frac{d\Theta(\rho_1)}{d\rho_1} \right] - 2(a_{11}\rho_1 + a_{12}\rho_2) + T[R_1 + 3R_1(1 - \ln T)], \quad (B1)$$

$$G_2 = T[R_2 \ln \rho_2 + \Theta(\rho_1)] - 2a_{12}\rho_1 + T[R_2 + \frac{5}{2}R_2(1 - \ln T)], \quad (B2)$$

where a_{11} and a_{12} are adjustable parameters, and $\Theta(\rho_1)$ is an adjustable function. For water and air, Benilov (2023a) proposed

$$a_{11} = 2112.1 \text{ m}^5 \text{ s}^{-2} \text{ kg}^{-1}, \quad a_{12} = 208 \text{ m}^5 \text{ s}^{-2} \text{ kg}^{-1}, \quad (B3a,b)$$

$$\Theta(\rho_1) = R \left[-q^{(0)} \ln \left(1 - 0.99 \frac{\rho_1}{\rho_{tp}} \right) + \sum_{n=1}^4 q^{(n)} \left(\frac{\rho_1}{\rho_{tp}} \right)^n \right], \quad (B4)$$

where

$$\rho_{tp} = 999.79 \text{ kg m}^{-3}, \quad (B5)$$

$$q^{(0)} = 0.071894, \quad q^{(1)} = 1.4139, \quad q^{(2)} = 8.1126, \quad (B6a-c)$$

$$q^{(3)} = -8.3669, \quad q^{(4)} = 4.0238. \quad (B7a,b)$$

When using the DIM, one also needs the Korteweg matrix K_{ij} . The value of K_{11} can be deduced from the surface tension of the liquid–water/vapour–water interface, and K_{22}

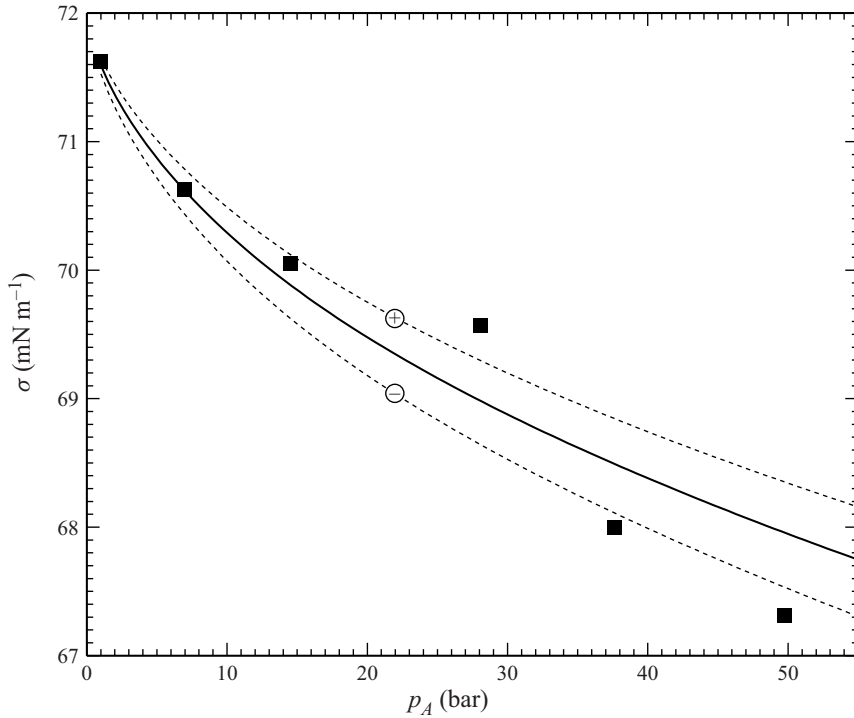


Figure 6. The surface tension of the water/air interface versus the pressure, at $T = 22.5^\circ\text{C}$. The black squares show the experimental data of Hinton & Alvarez (2021), and the solid line shows the theoretical result (B8) computed using the parameter values described in Appendix B and $K_{12} = 0.84978 \times 10^{-17} \text{ m}^7 \text{ s}^{-2} \text{ kg}^{-1}$ (chosen as the best fit of the experimental results). The dashed curves marked with ‘ \pm ’ are computed for $K_{12} = (0.84978 \pm 0.04810) \times 10^{-17} \text{ m}^7 \text{ s}^{-2} \text{ kg}^{-1}$, respectively.

from that of the liquid–air/vapour–air interface. The resulting values, computed by Benilov (2023a), are listed in equality (3.12).

The coefficient K_{12} is harder to calculate. One might think that it can be determined by comparing the liquid–water/vapour–water interface with the liquid–water/air interface – but at normal conditions, their characteristics are almost indistinguishable. Instead, K_{12} was deduced from the surface tension of the liquid–water/air interface at high pressure, measured by Hinton & Alvarez (2021). The following approach was used.

Within the framework of the DIM, the surface tension is given by

$$\sigma = \sum_{i,j} K_{ij} \int_{-\infty}^{\infty} \frac{d\rho_i}{dz} \frac{d\rho_j}{dz} dz, \tag{B8}$$

where $\rho_i(z)$ is the solution of the boundary-value problem (A5)–(A7). It was solved numerically with different values of K_{12} and the rest of the parameters given by (B1)–(B7) and (3.12). For each value of K_{12} , problem (A5)–(A7) was solved for various values of the atmospheric pressure p_A , and this computation was repeated until the best fit with the experimental dependence σ versus p_A is achieved. The results are shown in figure 6.

Unfortunately, Hinton & Alvarez (2021) provide data for only six values of p_A , and the resulting pairs (p_A, σ) do not sit on a smooth curve. Thus these measurements do not seem to merit the use of formal curve-fitting methods; instead, a value of K_{12} was chosen

'manually'. If sought in the form

$$K_{12} = b(K_{11}K_{22})^{1/2}, \tag{B9}$$

the best fit appears to be for $b = 0.53$ – for which (B9) yields the value of K_{12} given by (3.12). For comparison, figure 6 shows curves with K_{12} corresponding to $b = 0.50$ and $b = 0.56$.

Note that in the pure-fluid reduction of (B8), one can replace the coordinate z with the density ρ and then use the pure-fluid reduction of boundary-value problem (A5)–(A7) to obtain a closed-form integral involving the fluid's thermodynamic functions (see equation (7.12) of Benilov 2023a). As a result, σ can be found without solving the equations for the interfacial profile. In the multicomponent case, however, a similar change $z \rightarrow \rho_1$ does allow one to bypass solving the equations, as (B8) would involve the dependence of ρ_2 on ρ_1 .

Appendix C. Equation (3.61)

The ordinary differential equation (ODE) (3.61) is third order, but it can be reduced to a first-order equation of the Chini kind. To do so, multiply (3.61) by ρ_2 and integrate with respect to z . Fixing the constant of integration via boundary condition (3.62), one obtains

$$\rho_2 \frac{d^2 \rho_2}{dz^2} - \frac{1}{2} \left(\frac{d\rho_2}{dz} \right)^2 - \rho_2 + 1 = \frac{\xi}{\rho_2^2} \frac{d\rho_2}{dz}. \tag{C1}$$

Then changing $(\rho_2, z) \rightarrow (q, \rho)$, where

$$q = \frac{d\rho_2}{dz}, \quad \rho = \rho_2, \tag{C2a,b}$$

one obtains

$$\rho q \frac{dq}{d\rho} - \frac{1}{2} q^2 - \rho + 1 = \frac{\xi}{\rho^2} q. \tag{C3}$$

In terms of (q, ρ) , boundary condition (3.62) becomes

$$q = 0 \quad \text{at} \quad \rho = 1. \tag{C4}$$

C.1. Numerical results

Observe that boundary-value problem (C3)–(C4) is singular at the boundary point, hence to solve it numerically, one needs to 'step away' from $\rho = 1$. This can be done using the asymptotic expansion

$$q \sim \sum_{n=1}^{\infty} a_n (\rho - 1)^n \quad \text{as} \quad \rho \rightarrow 1. \tag{C5}$$

Substituting into (C3), one can calculate recursively as many coefficients a_n as one needs, e.g.

$$a_1 = \frac{\xi - \sqrt{\xi^2 + 4}}{2}, \quad a_2 = \frac{4\xi a_1 + a_1^2}{2\xi - 6a_1}, \quad \dots \tag{C6}$$

To rewrite (3.63) for $A(\xi)$ in terms of (q, ρ) , one should first integrate it by parts and use boundary conditions (3.62) and (3.64) to obtain

$$A = \int_{-\infty}^{+\infty} \frac{1}{\rho_2^4} \left(\frac{d\rho_2}{dz} \right)^2 dz. \tag{C7}$$

Rewriting this integral in terms of variables (C2), one obtains

$$A = \int_1^{+\infty} \frac{q(\rho)}{\rho^4} d\rho, \tag{C8}$$

where $q(\rho)$ is determined by (C3).

Boundary-value problem (C3)–(C4) was solved numerically by calculating 7 terms of series (C5) to set up a boundary condition at $\rho = 1.1$ (a smaller number of terms was found to be insufficient for large ξ). Then the solution was shot towards $\rho \rightarrow +\infty$ using the MATLAB function ‘ode23t’ (all other MATLAB functions for ODEs were found to be slow and inaccurate for large ξ). Finally, the computed solution was substituted into (C8), which was evaluated via Simpson’s rule.

C.2. Asymptotic results

As discussed in the main body of the paper, the classical and generalised models do not differ much for $\xi \sim 1$. Thus the asymptotic limit $\xi \rightarrow \infty$ is of particular importance.

If $\xi \gg 1$ and $\rho \sim 1$, then the first two terms in (C3) can be omitted, and one obtains

$$q \approx \xi^{-1} \rho^2 (-\rho + 1). \tag{C9}$$

The omitted terms become important (and the above solution inapplicable) if

$$\rho \gtrsim \xi^{2/5}. \tag{C10}$$

To find the solution for large ρ , introduce (q', ρ') such that

$$q = \xi^{1/5} \rho', \quad \rho = \xi^{2/5} \rho'. \tag{C11a,b}$$

In terms of the new variables, (C3) becomes, to leading order,

$$\rho' q' \frac{dq'}{d\rho'} - \frac{1}{2} q'^2 - \rho' = \frac{q'}{\rho'^2}, \tag{C12}$$

and its solution matches (C9) only if

$$q' \sim -\rho'^3 + O(\rho'^8) \quad \text{as } \rho' \rightarrow 0. \tag{C13}$$

The composite solution, valid for all ρ , is

$$q = \frac{\rho^2(\rho - 1)}{\xi} + \xi^{1/5} q'(\xi^{-2/5} \rho) - \frac{\rho^3}{\xi}, \tag{C14}$$

where the first term is the finite- ρ solution (C9), the second term reflects scaling (C11) for large ρ , and the last term is the common part of the two solutions. Substituting the above

expression into (C8), one obtains

$$A = -\frac{1}{\xi} - \frac{1}{\xi} \int_{\xi^{-1/5}}^{\infty} \frac{q'(\rho')}{\rho'^4} d\rho'. \quad (\text{C15})$$

One might be tempted now to take the limit $\xi \rightarrow \infty$, but the integral in (C15) is actually divergent (due to asymptotics (C13)). One can still use (C13) to express this integral through a convergent one:

$$\int_{\xi^{-1/5}}^{\infty} \frac{q'(\rho')}{\rho'^4} d\rho' = -\frac{2}{5} \ln \xi - \int_{\xi^{-1/5}}^{\infty} \ln \rho' \frac{d}{d\rho'} \left(\frac{q'}{\rho^3} \right) d\rho'. \quad (\text{C16})$$

Now take the limit $\xi \rightarrow \infty$, so that (C15) becomes

$$A \approx \frac{1}{\xi} \left[-1 + \frac{2}{5} \ln \xi + \int_0^{\infty} \ln \rho' \frac{d}{d\rho'} \left(\frac{q'}{\rho^3} \right) d\rho' \right]. \quad (\text{C17})$$

The integral in this expression does not involve ξ , hence it can be evaluated 'once and for all'. This was done by solving boundary-value problem (C12)–(C13) numerically and substituting the resulting solution into (C17), which yields

$$A \approx \frac{\frac{2}{5} \ln \xi - 0.81571}{\xi}. \quad (\text{C18})$$

C.3. Curve-fitting results

The large- ξ asymptotics (C18) can be used to derive an approximation of $A(\xi)$ for all ξ . To do so, assume that

$$A \approx \frac{\frac{2}{5} \ln(\xi + \xi_0) - 0.81571}{\xi + \xi_0} + \frac{A_0}{(\xi + \xi_0)^{7/5}}, \quad (\text{C19})$$

where ξ_0 and A_0 are undetermined constants; the former regularises asymptotics (C18) at $\xi = 0$, and the latter improves the accuracy for large ξ . (Note that a term $\sim \xi^{-7/5}$ would appear as a correction to the leading-order solution if (C18) were extended to higher orders.)

The coefficients ξ_0 and A_0 were fixed by fitting (C19) to the corresponding numerical result for the range $\xi \in [0, 10^5]$, using the MATLAB function 'fit'. The resulting values are those in (1.4) of the Introduction. The relative error of (1.4) within the range $\xi \in [0, 10^5]$ is approximately 0.8 %.

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