

1 Predicting Evaporation Rates of Droplet Arrays 2 [20-10-21]

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7 The evaporation of multiple sessile droplets is both scientifically interesting and practically
8 important, occurring in many natural and industrial applications. Although there are simple
9 analytic expressions to predict evaporation rates of single droplets, there are no such
10 frameworks for general configurations of droplets of arbitrary size, contact angle or spacing.
11 However, a recent theoretical contribution by Masoud *et al.* (2021) shows how considerable
12 insight can be obtained into the evaporation of arbitrary configurations of droplets without
13 having either to obtain the solution for the concentration of vapour in the atmosphere or to
14 perform direct numerical simulations of the full problem. The theoretical predictions show
15 excellent agreement with simulations for all configurations, only deviating by 25% for the
16 most confined droplets.

17 **Key words:** drops, condensation/evaporation, contact lines

18 1. Introduction

19 Sweat evaporating from an athlete's skin, agrochemicals sprayed onto crops, inkjet printers,
20 industrial spray coolers and virus transmission from infected surfaces all depend on the
21 collective evaporation of many droplets on a surface. However, despite these and many other
22 applications, nearly all of the considerable analytical, experimental and numerical work on
23 droplet evaporation has focused on a single droplet. Typically, the rate of evaporation is
24 controlled by the diffusion of vapour in the quiescent atmosphere, and is therefore described
25 by the "diffusion-limited model". In its simplest form this model involves solving Laplace's
26 equation for the concentration of vapour in the atmosphere subject to mixed boundary
27 conditions representing complete saturation at the free surface of the droplet, no flux of
28 vapour through the unwetted part of the substrate, and a far-field condition representing
29 the ambient vapour concentration. Lebedev (1965) and Popov (2005) provide a well known
30 analytic solution to this problem, giving a simple form for the diffusive vapour field and
31 evaporation rate for a single droplet.

32 However, in practice, droplets rarely occur in isolation, and so understanding the inter-
33 actions between multiple droplets is of considerable scientific and practical importance.

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34 Although research on 3-dimensional arrays of droplets (e.g. aerosols) is an extensive field,
 35 2-dimensional arrays of interacting *sessile* droplets on a surface are much less studied, and
 36 we summarise the key findings below. Kokalj *et al.* (2010) applied computational methods
 37 to droplet arrays and demonstrated that cooling was greatest for small dense droplet arrays,
 38 which would lead to a reduction in evaporation rate. Sokuler *et al.* (2010) show that in contrast
 39 to isolated droplets with constant contact angle, for which the evaporation rate reduces over
 40 time as $J = \frac{dV}{dt} \propto t^{1/2}$, for large droplet arrays, approximated as a continuous film, the
 41 evaporation rate is constant over time. For droplets with a pinned contact line, Carrier *et al.*
 42 (2016) also find, both experimentally and theoretically, that the total evaporation rate depends
 43 on droplet size and configurations. For small droplets, evaporation is diffusive-limited and
 44 proportional to the droplet’s size, whereas for droplets larger than around 20 μm , evaporation
 45 becomes convective with the rate proportional to the droplet area. They introduce the idea
 46 of a “superdrop” to predict the evaporation rate of droplet arrays and give a simple analytic
 47 expression to describe how the evaporation is hindered due to the presence of other droplets.
 48 For droplets dissolving in a surrounding fluid (an analogous situation also described by the
 49 Laplace equation), Chong *et al.* (2020) found evidence for a similar transition from diffusion
 50 to a convective plume. However, in their case, convection in the more dense arrays led to a
 51 surprising *increase* in dissolution rate.

52 A key concept throughout these studies is the so-called “shielding” effect, in which the
 53 presence of vapour from the other droplets reduces the evaporation rate (and hence increases
 54 the lifetime) of a droplet relative to that of the same droplet in isolation. However, none of
 55 the works mentioned above explicitly consider the variation in evaporation rate from one
 56 droplet to another, which will depend strongly on each droplet’s position within the array.
 57 This more involved problem was in fact first solved by Fabrikant (1985) for potential flow
 58 through a perforated membrane. Although a seemingly unrelated problem, Wray *et al.* (2020)
 59 recognised it as being analogous to the evaporation of zero-thickness circular droplets, and
 60 were able to integrate the expression for evaporation rate to obtain droplet drying times.
 61 These results are formally valid in the asymptotic limit where the droplets are well separated,
 62 with the problem reducing to a system of N linear equations describing the evaporation rate
 63 from each droplet. Both Fabrikant (1985) and Wray *et al.* (2020) found good agreement
 64 between the theoretical predictions for a pair of identical droplets with those of numerical
 65 calculations right up to the limit of touching droplets. In this case the effect of shielding
 66 increases the lifetime of the droplets by one third. In addition Wray *et al.* (2020) obtained
 67 expressions describing the variation of flux across the surface of each droplet.

68 Very recently, Edwards *et al.* (2021) found very good agreement between the theoretical
 69 predictions of Fabrikant (1985) and experimental results obtained using an interferometric
 70 technique to directly measure the individual evaporation rate of up to 25 droplets in ten
 71 different configurations.

72 **2. Overview of Masoud *et al.* (2021)**

73 The work of Masoud *et al.* (2021) is important and novel as it extends the findings of
 74 Fabrikant (1985), removing the restrictions of thin droplets and circular contact lines. They
 75 used Green’s second identity to simply and elegantly obtain an exact relationship between the
 76 local flux and total evaporation rate from the droplets. Using the method of reflections and
 77 assuming that the droplets are well separated, they obtained a system of N linear equations
 78 for the evaporation rates from each droplet J_n involving only the rate for the isolated droplet
 79 \hat{J}_n and $\phi(r_{nm})$, the normalised vapour concentration at the location of the m^{th} droplet:

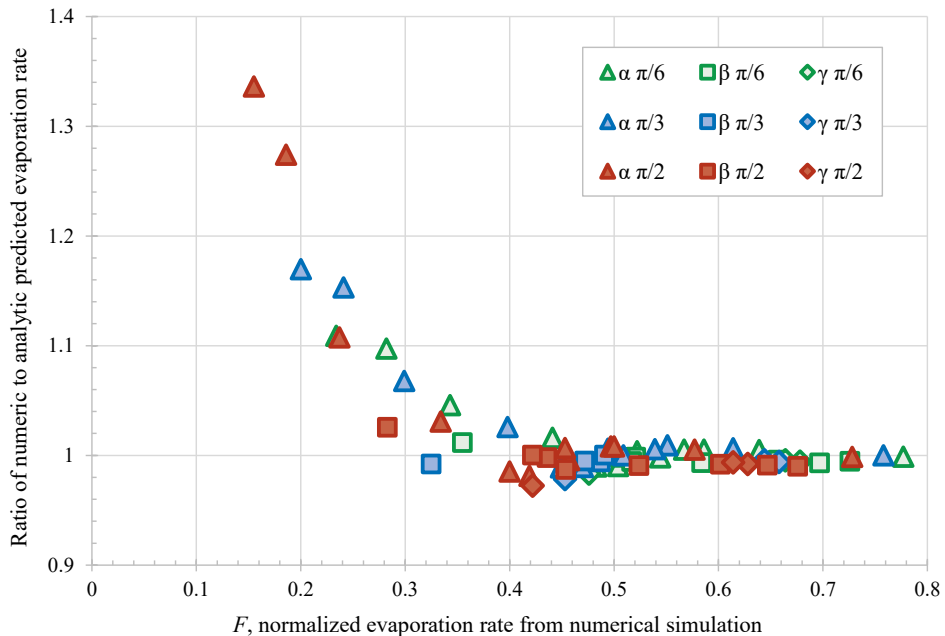


Figure 1: Comparison of the predictions using the new analytical approach to direct numerical simulations, for various droplet arrangements and confinements (α - most confined, β and γ - least confined) and initial contact angles ($\pi/6$, $\pi/3$ and $\pi/2$) for droplet separation of 2.5 radii.

$$F = \frac{J_n}{\hat{J}_n} = 1 - \sum_{m=1}^N \phi(r_{nm}) \frac{J_m}{\hat{J}_n}. \quad (2.1)$$

Using for example the the expression of Popov (2005) for $\phi(r_{nm})$, this set of N equations can be solved giving the evaporation rates for each droplet. For low contact angles the vapour concentration term reduces to $\phi(r_{nm}) = \frac{2}{\pi} \arcsin\left(\frac{a_n}{r_{nm}}\right)$, recovering the simpler form derived by Fabrikant (1985).

The authors evaluated the accuracy of their theoretical predictions by comparing with the results of direct numerical simulations for twelve different configurations of identical spherical-cap droplets using three different contact angles ($\pi/6$, $\pi/3$ and $\pi/2$) and droplet separations of 2.5 and 3 radii. (i.e., 72 different calculations). Fig. 1 replots the data-set provided in Table 1 for the closest separations, confirming the excellent agreement for the faster evaporation droplets, with $F > 0.4$. For slower evaporating droplets ($F < 0.4$) which are more confined and have larger contact angles, the theoretical results are systematically slower than the numerical results.

3. Future

The great merit of this work is that it quantifies the significance of the shielding effect in arbitrary configurations of droplets of different sizes and contact angles without having either to obtain the solution for the concentration of vapour in the atmosphere or to perform time-consuming and technically challenging direct numerical simulations. Moreover, it opens the

98 door to a greater understanding of the many applications of this effect and could lead to
 99 improvements in inkjet printers or cooling systems, for example.

100 Here we briefly mention three specific directions for potential future work. Firstly, the
 101 systematic discrepancy seen for the most confined droplets could be investigated further.
 102 As the droplets have a centre to centre separation of 2.5 radii the theory is expected to
 103 be accurate, so the disagreement is most likely due to a more subtle effect of confinement
 104 rather than the prediction being applied beyond its valid range. Any improvement to the
 105 theory should be verified against additional numerical and experimental work. Secondly, it
 106 would be interesting to explore the collective behaviour of an increasing number of droplets
 107 and thereby determine to what extent a collection of many droplets can be considered as
 108 one large “super droplet” of an appropriate shape and size, as discussed by Carrier *et al.*
 109 (2016). Thirdly, and most generally, the same theoretical approach can be applied to other
 110 physical situations governed by Laplace’s equation such as for example the dissolution of
 111 microbubbles, as reviewed by Lohse & Zhang (2015); Qian *et al.* (2019).

112 **Declaration of interests.** The author reports no conflict of interest.

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