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Abstract: Equilibrium of liquid droplets on soft deformable substrates is investigated.. Disjoining pressure action in the vicinity of the apparent three phase contact line is taken into account. It is shown that the disjoining pressure action determines the substrate deformation. A simplified linear disjoining pressure isotherm and simple Winkler's model to account for the substrate deformation are used which allows to deduce an analytical solutions for both the liquid profile and substrate deformation. The apparent equilibrium contact angle that the liquid makes with the substrate is calculated and it's dependency on the system parameters is investigated.

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Dear Madam/Sir,

We submit of our manuscript **Equilibrium of droplets on a deformable substrate: influence of disjoining pressure** by Gulraiz Ahmed, Victor M. Starov, Vasily V. Kalinin for a publication in a Special Issue "Victor Starov".

Regards, Victor Starov

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Graphical abstract

Highlights

Deformable substrates

Disjoining pressure

Transition zone

Equilibrium of droplets on a deformable substrate: influence of disjoining pressure

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Abstract

Equilibrium of liquid droplets on soft deformable substrates is investigated.. Disjoining pressure action in the vicinity of the apparent three phase contact line is taken into account. It is shown that the disjoining pressure action determines the substrate deformation. A simplified linear disjoining pressure isotherm and simple Winkler's model to account for the substrate deformation are used which allows to deduce an analytical solutions for both the liquid profile and substrate deformation. The apparent equilibrium contact angle that the liquid makes with the substrate is calculated and it's dependency on the system parameters is investigated.

Key words: equilibrium droplets, deformable substrate, disjoining pressure.

1 Introduction

Equilibrium of a liquid droplet on a solid substrate is frequently described based on Young's equation [1]. This simplified equation involves the balance of the horizontal forces leaving the vertical force unbalanced. The latter is possible in the case of a rigid substrate but should be reconsidered in the case of soft deformable substrates. It has been shown in [2] that disjoining pressure action in the vicinity of the apparent three phase contact line results in a deformation of a soft solid substrate.

Note, that in the case of direct application of Young's equation, at the three phase contact line there exists deformation singularity, i.e. the substrate deformation goes to infinity [3-7]. Their investigation revealed that all the equilibrium properties (i.e. contact angle, droplet radius, droplet volume, etc.) of the system under consideration rely upon the selected artificial length parameter which determines a width of zone near the contact line where surface tension is applied. However,

disjoining pressure is a real physical phenomenon, which determines the substrate deformation of soft surfaces.

The shape of the disjoining pressure isotherm determines the shape and the contact angle of the droplet at equilibrium [2].

The problem of equilibrium of the droplet on a deformable substrate has recently gained a lot of interest. There have been a number of experimental studies recently that have considered liquids on deformable substrates [8-14], where deformation of soft solids was investigated near the apparent contact line, but there exists a gap in theoretically understanding of the problem. This void is primarily due to the stress singularity present at the three phase contact line.

Below a mathematical model is presented which incorporates the effect of both capillary and disjoining pressure isotherm on the substrate deformation.

2 Disjoining pressure and deformation of soft solids

In the case of partial wetting, that is, equilibrium contact angle, $\theta_e > 0$, the shape of the transition zone between the bulk of the liquid and the thin film on substrate can be expressed via the disjoining pressure isotherm, $\Pi(h)$ [2]. The equilibrium excess pressure, P_e , inside the droplet can be expressed according to Kelvin's equation [2] and the equilibrium contact angle of a two dimensional droplet can be expressed using the disjoining pressure isotherm as follows [2]:

$$\cos \theta_{e,2D} = 1 + \frac{\Pi(h_e)h_e}{\gamma} + \frac{1}{\gamma} \int_{h_e}^{\infty} \Pi(h)dh \quad (1)$$

Eq. (1) corresponds to an S-shaped disjoining pressure isotherm, $\Pi(h)$. Below the transition zone between the flat meniscus and a film having $\theta_e > 0$ is examined below as an example, following [15]. The liquid droplet forms a wedge changing to flat equilibrium film of thickness, h_e , far from the droplet (Fig. 1).

Fig. 1. 1- Profile of transition zone $h(x)$ between bulk liquid and flat wetting film, 2- s shaped disjoining pressure isotherm, $\Pi(h)$, and 3 - profile of normal pressures acting on substrate; x_Y is the position where the vertical force is exerted.

The origin is taken at $x = 0$, which is a point on the profile lying beyond the influence of surface forces (Fig. 1). The profile of the transition zone, $h(x)$, can be calculated according to Eq. (2), which includes the influence of capillary and disjoining pressure isotherm [2]:

$$\gamma K(x) + \Pi(h) = \frac{\gamma h''}{[1+(h')^2]^{3/2}} + \Pi(h) = P_e, \quad (2)$$

where, $h' = dh/dx$; $h'' = d^2h/dx^2$; and P_e is the excess pressure in the droplet. In the region of the flat equilibrium film, $h'' = 0$ and $\Pi(h_e) = P_e$. In bulk of the liquid, beyond the influence of the surface forces, $\Pi = 0$ and $P_e = \gamma/R$, where R is the radius of curvature of the droplet. In case of planar wedge (Fig. 1, $R = \infty$ and $P_e = 0$). Therefore from Eq. (2) can be concluded:

$$\Pi(h) = -\frac{\gamma h''}{[1+(h')^2]^{3/2}}. \quad (3)$$

The resultant forces on the substrate is given by the following equation:

$$F = \int_0^\infty \Pi(h) dx. \quad (4)$$

Substituting the disjoining pressure isotherm $\Pi(h)$ according to Eq. (3) in equation (4) results in:

$$\begin{aligned} F &= -\gamma \int_0^\infty \left\{ \frac{h''}{[1+(h')^2]^{3/2}} \right\} dx = \gamma \frac{h'(0)}{\sqrt{[1+(h'(0))^2]}} = \frac{\gamma \tan \theta_e}{\sqrt{[1+\tan^2 \theta_e]}} \\ &= \gamma \sin \theta_e \end{aligned} \quad (5)$$

Boundary conditions used in the expression above (equation (5)) are, $h'(\infty) = 0$ and $h'(0) = \tan \theta_e$. This shows that the integration performed over the local values leads to same expression as the vertical component of the surface tension from the Young's equation. In contrast to Young's, the force is not exerted on a specific point, but is distributed over the region where disjoining pressure acts, i.e. transition zone. Based on this conclusion a mathematical model is derived in the next section.

3 Mathematical model

In this section a mathematical model is derived for a liquid droplet on a deformable substrate. According to Winkler's model there is a linear relationship between the local deformation and the applied local stress [16, 17]. Winkler's model is used below.

Fig. 2. Schematic diagram of the liquid droplet on a deformable substrate.

Deformation in the soft substrate is local and is directly proportional to the applied pressure, P . According to Winkler's model,

$$h_s = -KP, \quad (6)$$

where K is the elasticity coefficient, h_s is the local deformation of the substrate due to the presence of the droplet above.

The pressure in the ambient air is P_{air} . Under the action of the pressure from the ambient air the solid deformation is:

$$h_{se} = -KP_{air} \quad (7)$$

The deformed solid substrate is covered by equilibrium liquid thin film, which is calculated according to well-known Kelvin's equation:

$$\Pi(h_e) = P_e = \frac{RT}{v_m} \ln \frac{p_{sat}}{p} \quad (8)$$

where, v_m is the molar volume of the liquid, T is the temperature, R is the gas constant; vapour pressure, p , which is higher than the saturated pressure p_{sat} . Reminder, a droplet can be at the equilibrium with oversaturated vapour pressure.

The excess free energy of the equilibrium thin film of deformed solid per unit area is given by,

$$\frac{F_{e,film}}{S_{film}} = \gamma + \gamma_s + P_e h_e + \frac{h_{se}^2}{2K} + \int_{h_e}^{\infty} \Pi(h) dh, \quad (9)$$

where, $P_e = P_{air} - P_{liquid}$, γ and γ_s are liquid-vapour and solid-vapour interfacial tensions. This free energy should be subtracted from the free energy of the droplet, otherwise the excess free energy of the droplet will be infinite. The excess free energy of the droplet is as follows,

$$F - F_{e,film} = \gamma \Delta S + \gamma_s \Delta S_s + \Delta V + F_{surface\ forces} + F_{deformation} \quad (10)$$

where Δ means “as compared with a flat equilibrium film”. Therefore Eq. (10) can be rewritten as:

$$F - F_{e,film} = 2\pi \int_0^{\infty} r f(h, h', h_s, h'_s) dr \quad (11)$$

where

$$f(h, h', h_s, h'_s) = r \left\{ \gamma \sqrt{1 + h'^2(r)} - \gamma + \gamma_s \sqrt{1 + h_s'^2(r)} - \gamma_s + \frac{P_e(h - h_s) - P_e h_e + \left[\frac{h_s^2}{2K} - \frac{h_{se}^2}{2K} \right] + \int_{h-h_s}^{\infty} \Pi(h) dh - \int_{h_e}^{\infty} \Pi(h) dh}{r} \right\} \quad (12)$$

The expression under the integral in Eq. (11) tends to zero as r tends to infinity.

Under equilibrium conditions the excess free energy should reach to a minimum value. To satisfy this condition the first variation of free excess energy should be zero, which results in two Euler equations for the droplet and soft substrate profiles,

$$\frac{d}{dr} \left(\frac{\partial f}{\partial h'} \right) - \frac{\partial f}{\partial h} = 0 \quad (13)$$

$$\frac{d}{dr} \left(\frac{\partial f}{\partial h'_s} \right) - \frac{\partial f}{\partial h_s} = 0 \quad (14)$$

Substitution the expression for f from (12) into equations (13) and (14) results in the following equations:

$$\frac{\gamma}{r} \frac{d}{dr} \frac{r h'}{(1 + h'^2)^{1/2}} + \Pi(h - h_s) = P_e \quad (15)$$

$$\frac{\gamma_s}{r} \frac{d}{dr} \frac{r h'_s}{(1 + h'^2_s)^{1/2}} - \Pi(h - h_s) - \frac{h_s}{K} = -P_e \quad (16)$$

These two equations (15) and (16) forms a system of two differential equations for two unknown profiles: the liquid droplet, $h(r)$, and deformed solid substrates, $h_s(r)$.

Below the simplest case is considered, $\gamma_s = 0$. Eq.(16) in this case results in:

$$h_s = -K[\Pi(h - h_s) - P_e] \quad (17)$$

Region where surface forces are negligible, i.e. bulk of the droplet, equation (17) gives:

$$h_s = K P_e < 0 \quad (18)$$

because P_e is negative. Hence, Eq. (17) is in agreement with definition (6). In the case of a low slope approximation, $h'^2 \ll 1$ the profile of the droplet $h(r)$ and profile of the substrate $h_s(r)$ satisfy the following set of a second order ordinary differential equation and an algebraic equation:

$$\gamma \left(h'' + \frac{h'}{r} \right) + \Pi(h - h_s) = P_e \quad (19)$$

$$h_s = -K[\Pi(h - h_s) - P_e] \quad (20)$$

Eq. (19) is different from the usual capillary equation for the droplet on a rigid substrate, because now the disjoining pressure term depends on the profile of the deformable substrate, $h_s(r)$, which is determined according to Eq. (20). Both Eq. (19) and (20) are coupled and can be solved numerically only, however, below the problem is simplified even further to obtain an analytical solution. For this purpose we adopt a very simple disjoining pressure isotherm (linear function of h (Fig. 3)):

$$\Pi(h) = \begin{cases} P_1 - ah & \text{at } h \leq t_1 \\ 0 & \text{at } h > t_1 \end{cases} \quad (21)$$

Fig.3. Simplified disjoining pressure isotherm adopted below for calculations.

where P_1 and t_0 are defined in Fig. 3, t_1 is the range of surface forces action. The slop a of the $\Pi(h)$ dependency is given by:

$$a = \frac{P_1 - P_e}{h_e} \quad (22)$$

The selected linear dependency of the disjoining pressure isotherm $\Pi(h)$ on h according to Eq. (21) in spite of considerable simplification still captures the essential properties of the disjoining pressure isotherm (i) it satisfies the stability condition, $\Pi'(h) < 0$ when $h < t_1$, and (ii) the influence of surface forces is short range and radius of it's action is defined by t_1 ; (iii) at the proper choice of the disjoining pressure parameters (see below) it corresponds to the particle wetting case.

3.1 Droplets on a non-deformable substrate

Fig. 4 shows a schematic diagram of the droplet on the non-deformable solid substrate, where L_1 is the length from the origin to the point where the influence of the surface forces come into play, L is the effective radius of the droplet base, $\theta_{e,nd}$ is the equilibrium contact angle the droplet makes with the solid substrate, R is the radius of the droplet.

Fig. 4. Schematic diagram of a droplet on a non-deformable solid substrate. Here, $R = -\frac{\gamma}{P_e}$.

For a solid non-deformable substrate Eq. (19) reduces to:

$$\gamma \left(h'' + \frac{h'}{r} \right) + \Pi(h) = P_e. \quad (23)$$

For the bulk of the liquid droplet, i.e. the spherical region, at $h > t_1$ and $\Pi(h) = 0$. Hence, in this region Eq.0 (23) transforms into:

$$\gamma \left(h'' + \frac{h'}{r} \right) = P_e. \quad (24)$$

Solution of this equation is:

$$h(r) = Ar^2 + B, \quad (25)$$

where integration constants, A and B , can be determined from boundary conditions at $r=0$ and $r=L$ and this solutions is marked by a subscript *out*:

$$h_{out}(r) = -\frac{r^2}{2R} + \frac{R \sin^2 \theta_{e,nd}}{2} \text{ or } h_{out}(r) = \frac{-r^2 + L^2}{2R} \quad (26)$$

For the transition region in the case of non-deformable solid substrate Eq. (23) along with the disjoining pressure isotherm given by Eq. (21) results in:

$$h'' + \frac{h'}{r} - \frac{ah}{\gamma} = \frac{P_e - P_1}{\gamma} \quad (27)$$

Solution of Eq. **Error! Reference source not found.** is as follows:

$$h_{in} = h_e + C_{1,nd} K_0(r\lambda) \quad (28)$$

where, $\lambda = \sqrt{\frac{a}{\gamma}}$, K_0 is the modified Bessel function of the zero order, $C_{1,nd}$ is an integration constant and this solution is marked by the subscript *in*. Solutions (25) and (28) should satisfy the following boundary conditions:

$h_{out}(L_1) = h_{in}(L_1) = t_1$; $h'_{out}(L_1) = h'_{in}(L_1)$, which gives three equations for determination of the three unknown coefficients: $C_{1,nd}$, $\theta_{e,nd}$ and L_1 .

Solution results in

$$C_{1,nd} K_0(L_1 \lambda) = \frac{L_1}{R \lambda}, \quad (29)$$

$$\theta_{e,nd} = \sqrt{\frac{2}{R} \left(h_e + \frac{L_1}{R \lambda} + \frac{L_1^2}{2R} \right)}, \quad (280)$$

$$L_1 = R \lambda (t_1 - h_e). \quad (291)$$

The latter allows determining the effective radius of the droplet, L :

$$L = \sqrt{L_1^2 + 2R t_1}. \quad (32)$$

Introducing: $P_2 = a(t_0 - t_1)$, Eq. (30) can be rewritten as

$$\theta_{e,nd} = \sqrt{\frac{1}{\gamma a} [P_e^2 - P_1 P_e - P_2 P_e + P_2^2]} \quad (33)$$

The equilibrium contact angle was previously determined according to Eq. (1). According to the simplifies isotherm (Fig. 3) this contact angle can be calculated as:

$$\theta_{e,2D} = \sqrt{2 \left\{ -\frac{P_e h_e}{\gamma} - \frac{a}{\gamma} \left[t_0(t_1 - h_e) - \frac{1}{2}(t_1^2 - h_e^2) \right] \right\}} \quad (34)$$

3.2 Deformable substrate

Fig. 5 shows a schematic diagram of a droplet on the deformable solid substrate based on the assumption of a linear disjoining pressure, according to Eq. (21) (Fig.3). In Fig. 5 $\theta_{e,d}$ is the equilibrium macroscopic contact angle of the droplet with the soft substrate.

Fig. 5. Schematic diagram of droplet on a deformable/soft substrate

For the bulk of the liquid droplet, i.e. the spherical region, $h - h_s > t_1$, that is at $0 < r < L_1$ (Fig. 5) from Eq. (21):

$$h_s = K P_e \quad (35)$$

For the transition region, that is at $r > L_1$ Eq. (20) gives:

$$h_s = \frac{K(P_e - P_1 + ah)}{aK + 1} \quad (36)$$

Substitution of the latter expression into Eq. (20) results in:

$$h'' + \frac{h'}{r} - \frac{a}{\gamma(aK + 1)} h = \frac{P_e - P_1}{\gamma(aK + 1)}$$

Solution of the latter equation is:

$$h = h_e + C_{1,d} K_0(r\beta),$$

where $\beta = \sqrt{\frac{a}{\gamma(aK+1)}}$ and $C_{1,d}$ is an integration constant.

The droplet profile and its derivative should be continuous at $r=L_1$, but the solid surface should be discontinuous and $h(L_1)-h_s(L_1)=t_1$. Using these boundary conditions all unknown constants $C_{1,d}$, $\theta_{e,d}$ and L , can be determined. Below only expression for the contact angle on the deformable substrate is presented:

$$\theta_{e,d}^2 = \frac{KP_2}{\gamma} (P_2 - P_e) + \theta_{e,nd}^2 \quad (37)$$

where $P_2 = a(t_0 - t_1)$. That is if $K = 0$ (non-deformable substrate), then $\theta_{e,d} = \theta_{e,nd}$. Expression $\frac{KP_2}{\gamma} (P_2 - P_e)$ in Eq. (37) is always positive, that is, $\theta_{e,d} > \theta_{e,nd}$ the contact angle on a deformable substrate is always bigger than the contact angle on the corresponding non-deformable substrate.

4 Results and Discussion

4.1 Non-deformable substrate: Effect of variation of P_e on $\theta_{e,nd}$

Dependency of equilibrium contact angles $\theta_{e,nd}$ and $\theta_{e,2D}$ from according to Eqs. (280) and (34) on P_e are presented in this section. Below $|P_e|$ is varied from 0 to $|P_2|$. Fig. 6 shows the behaviour of equilibrium contact angle when the excess pressure is varied. Fig. 6a shows the variation at $t_0 = 7 \times 10^{-7}$ cm and Fig. 6b shows variation at $t_0 = 2 \times 10^{-6}$ cm. In both cases, two-dimensional equilibrium contact angle $\theta_{e,2D}$ calculated according to Eq (34) turns up to be higher as compared with three-dimensional equilibrium contact angle calculated according to Eq. (33). It was shown earlier [2] $\theta_{e,2D}$ can only increase with the decrease in P_e , which is in agreement with both Figs. 6a and 6b. However, it turns out that the dependency of the contact angle in the three dimensional case, $\theta_{e,nd}$, differs substantially from the two dimensional case (always higher) and can go via minimum as a function of P_e (Fig. 6a). If the value of t_0 is increased and brought closer to t_1 then $\theta_{e,nd}$ is an increasing function of P_e with no minimum.

(b) $t_0 = 2 \times 10^{-6}$ cm

Fig. 6. No-deformable substrate. Effect of variation of excess pressure, P_e on two-dimensional, $\theta_{e,2D}$, and three-dimensional, $\theta_{e,nd}$, equilibrium contact angles. Selected parameters are as shown

Physical Property	Value
Surface tension, γ	72 dyn/cm
t_1	3×10^{-6} cm
a	1×10^{11} dyn/cm ³
$ P_e $	$0 - P_2 $ dyn/cm ²

4.2 Deformable substrate

The profile of the droplet and the subsequent changes in the deformable substrate are plotted in Fig. using equations deduced in section 3.2. The soft substrate behaves in the same manner as predicted in Fig. 5.

Fig. 6. Deformable substrate. Calculated profile of the droplet and deformable substrate.

Physical Property	Value
Surface tension, γ	72 dyn/cm
t_1	3×10^{-6} cm
t_0	7×10^{-7} cm
a	1×10^{11} dyn/cm ³
K	1×10^{-11} cm ³ /dyn

4.3. Effect of variation of P_e on equilibrium contact angle $\theta_{e,d}$

Excess pressure is varied as before $0 \leq |P_e| \leq |P_2|$. In order to compare the results with the $\theta_{e,nd}$, the value of K is increased gradually from 0 to 1×10^{-11} . All other physical properties of the disjoining pressure isotherm are kept the same as shown in Fig. 6. for t_0 which is equal to 7×10^{-7} cm in Fig. 7a and is equal to 2×10^{-6} cm in Fig. 7b. Fig. 7a has a minima for $\theta_{e,d}$ and it becomes an increasing function as t_0 is increased and brought closer to t_1 (Fig. 7b). It is apparent from Fig. 7 that $\theta_{e,d}$ is bigger than $\theta_{e,nd}$ for all values of $K > 0$.

(a) $t_0 = 7 \times 10^{-7} \text{ cm}$

(a) $t_0 = 2 \times 10^{-6} \text{ cm}$

Fig. 7. Effect of variation of excess pressure, P_e on $\theta_{e,d}$ for different values of elasticity coefficient K .

5 Conclusions

Equilibrium of liquid droplets on soft deformable substrates is investigated. Disjoining pressure action in the vicinity of the apparent three phase contact line is taken into account. It is shown that the disjoining pressure action determines the substrate deformation. A simplified linear disjoining pressure isotherm and simple Winkler's model to account for the substrate deformation are used which allows deducing an analytical solutions for both the liquid profile and substrate deformation. The apparent equilibrium contact angle that the liquid makes with the substrate is calculated and its dependency on the system parameters is investigated. It is shown also that the calculated equilibrium contact angles in the case of non-deformable substrates in the case of two-dimensional droplets are bigger as compared with three dimensional case. It is shown that equilibrium contact angles on deformable substrates are always higher than on the corresponding non-deformable substrates.

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Abstract

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