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BASIC MECHANICAL PROPERTIES OF WET GRANULAR MATERIALS: A DEM STUDY

Vinh Du Than¹, Saeed Khamseh¹,

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ABSTRACT

We use discrete, grain-level numerical simulations of a model granular assembly, made of spherical balls, to investigate the influence of a small amount of an interstitial wetting liquid, forming capillary bridges between adjacent particles, on two basic aspects of granular material rheology: (i) the plastic response in isotropic compression, and (ii) the critical state under monotonic shear strain, and its generalization to steady, inertial flow. Tensile strength $F_0 = \pi\Gamma a$, in contacts between beads of diameter a joined by a small meniscus of a liquid with surface tension Γ , introduces a new force scale and a new dimensionless control parameter, $P^* = a^2P/F_0$, for grains of diameter a under confining stress P . Under low P^* , as cohesion dominates, capillary cohesion may stabilize very loose structures. Upon increasing pressure P in isotropic compression, such structures gradually collapse. The resulting irreversible compaction is well described by the classical linear relation between $\log P^*$ and void ratio in some range, until a dense structure forms which retains its stability without cohesion as confinement dominates for large P^* . In steady shear flow, with uniform velocity gradient $\dot{\gamma} = \frac{\partial v_1}{\partial x_2}$ under normal stress $P = \sigma_{22}$, the apparent internal friction coefficient, which we define as $\mu^* = \frac{\sigma_{12}}{\sigma_{22}}$, depends on P^* and inertial number (reduced shear rate) $I = \dot{\gamma}\sqrt{\frac{m}{aP}}$, and so does solid fraction Φ . The material exhibits, as P^* decreases, a strongly enhanced resistance to shear (larger μ^*). In the quasistatic limit, for $I \rightarrow 0$, it is roughly predicted by a simple effective pressure assumption, by which the capillary forces are deemed equivalent to

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23 an isotropic pressure increase applied to the dry material, as long as $P^* \geq 1$, while the yield
24 criterion approximately assumes the Mohr-Coulomb form. At lower P^* , such models tend to
25 break down as liquid bonding, causing connected clusters to survive over significant strain
26 intervals, strongly influences the microstructure. Systematic shear banding is observed at
27 very small P^* .

28 **Keywords:** granular materials, cohesion, capillary forces, effective pressure, Mohr-Coulomb,
29 DEM

30 INTRODUCTION

31 Over the last decades our understanding of the microstructural and micromechanical
32 origins of macroscopic granular material rheology have greatly benefitted from the devel-
33 opment of numerical simulation methods of the so-called discrete element type (DEM), in
34 which the motions and interactions of individual grains are modeled (Radjaï and Dubois
35 2011). In particular, two basic concepts, which had previously been identified and exploited
36 in process, chemical and geomechanical engineering, were revisited, and supported by mi-
37 cromechanical interpretations. One is the *random close packing (RCP) state* (Cumberland
38 and Crawford 1987), the configuration of a granular assembly maximizing density under the
39 constraint of mutual non-interpenetrability of the grains, without any specific ordering. The
40 RCP state is a stable equilibrium state of an isotropically compressed assembly of rigid,
41 frictionless grains (Agnolin and Roux 2007a; Donev et al. 2005; O’Hern et al. 2003), and
42 its characteristics, most notably its solid fraction, are remarkably reproducible, irrespective
43 of the dynamical assembling method. Furthermore, because frictionless particle assemblies
44 appear to be devoid of dilatancy – which has been explicitly checked for disks and spherical
45 beads (Peyneau and Roux 2008a), and for polygonal shapes in 2D (Azéma et al. 2015) – the
46 same solid fraction (about 0.64 for identical beads) is obtained on preparing, without any
47 friction mobilization, packings under different, possibly anisotropic, conditions (Silbert et al.
48 2002; Peyneau and Roux 2008b). In the presence of friction, many different states can be ob-
49 served, varying in density and coordination number (Agnolin and Roux 2007a; Magnanimo

50 et al. 2008), even if stresses and microstructure are isotropic. The second traditional notion
51 which has been revisited by DEM, with due attention to its microscopic foundations, is that
52 of the *critical state*, in the sense of soil mechanics (Wood 1990; Mitchell and Soga 2005): the
53 steady state of plastic flow attained, irrespective of the initial state, after large enough strain
54 in monotonically, quasistatically sheared material. The critical state has been shown (Radjaï
55 et al. 2004; Rothenburg and Kruyt 2004; Radjaï et al. 2012; Kruyt and Rothenburg 2014)
56 to be an attractor state for all variables characterizing internal structure, micromorphology
57 and forces, such as coordination numbers, fabric tensors or friction mobilization, as well
58 as for stresses and solid fraction. Upon increasing the shear rate, the material behavior is
59 affected by inertial effects, and the internal state of the homogeneously sheared material
60 depends, under controlled normal stress P , on inertial number I (as defined in the abstract),
61 the quasi-static critical state corresponding to the limit of $I \rightarrow 0$. This generalization of the
62 critical state to I -dependent steady homogeneous shear flows has led to the formulation of
63 efficient constitutive laws for dense granular flows (Forterre and Pouliquen 2008; Andreotti
64 et al. 2013), in terms of the I dependence of internal friction coefficient μ^* and solid frac-
65 tion Φ . The RCP state (or another well-controlled homogeneous isotropic packing state)
66 on the one hand, and the critical state, on the other hand, correspond to the initial and
67 the final states in many relevant mechanical tests – typically one starts from some isotropic
68 packing, of which the RCP is an important limiting case, and one imposes a loading path
69 leading to the critical state (Thornton 2000; Radjaï et al. 2004). Their interest also stems
70 from their lack of dependence on many features and parameters governing contact behavior,
71 especially dynamical ones, but also elastic contact stiffnesses, in the frequent case of negli-
72 gible contact deflections (Roux and Chevoir 2011). On introducing new models for grains,
73 with such features as rolling resistance or angularity (Azéma et al. 2013; Saint-Cyr et al.
74 2012; Estrada et al. 2011), it is natural to first investigate microstructural and mechanical
75 properties of RCP and critical states. Cohesive forces in contacts significantly affect both
76 isotropic packings (Gilabert et al. 2007) and steady shear flows (Rognon et al. 2006).

77 The present paper states some essential results obtained by DEM simulations, for both
78 isotropically compressed static assemblies, and I -dependent steady uniform shear flows, with
79 special emphasis on the critical state in the limit of $I \rightarrow 0$, in the case of a model of wet
80 spherical grains. Compared to similar numerical studies in the literature (Richefeu et al.
81 2006; Scholtès et al. 2009b) the ones presented here investigate looser structures, which
82 could not be observed with dry grains – as evidenced in experiments with sands (Bruchon
83 et al. 2013a; Bruchon et al. 2013b). While both situations should be more extensively
84 studied, in more detailed publications (Khamseh et al. 2015; Than et al. 2015), in which
85 thorougher investigations of microscopic aspects will be presented, some salient features of
86 isotropic compression and steady shear flows are described, stressing the differences with
87 dry, cohesionless materials.

88 The paper is organized in the following way. Once the model material and the inter-
89 actions are suitably described in the forthcoming section (“*Model material and simulation*
90 *ingredients*”), the two main parts of the paper separately address these two important as-
91 pects of wet granular material rheology: “*Isotropic assembly and compression*”, and then
92 “*Dense shear flow and critical states*”. The final *Conclusion* section sums up the results and
93 puts them in perspective.

94 **MODEL MATERIAL AND SIMULATION INGREDIENTS**

95 **Stress and Strain Control**

96 Our model material is an assembly of N equal-sized spherical beads of diameter a . The
97 simulation cell is a rectangular cuboid, with edges, of lengths $(L_\alpha)_{1 \leq \alpha < 3}$, parallel to coordi-
98 nate axes, periodic in all three directions. We control all three diagonal stress components in
99 isotropic compression, and wait for equilibrium conditions, as in Agnolin and Roux (2007a),
100 to be satisfied within a preset tolerance. In shear tests a granular flow is imposed in direction
101 1, with a gradient in direction x_2 , defining shear rate $\dot{\gamma} = \frac{\partial v_1}{\partial x_2}$. In that case, the periodic-
102 ity in direction 2 is implemented through the Lees-Edwards procedure (Allen and Tildesley

103 1987), length L_2 is allowed to vary in response to the enforced condition of constant normal
 104 stress σ_{22} , while lengths L_1 and L_3 are kept fixed – as in Peyneau and Roux (2008a).

105 **Force Model: Elasticity and Friction**

106 The spherical beads are assumed to be made of a material with Young modulus E
 107 and Poisson ratio ν . Coulomb friction applies in the contacts, with friction coefficient μ .
 108 Elastic-frictional contact forces are modeled with a simplified Hertz-Mindlin-Deresiewicz
 109 force law (Agnolin and Roux 2007a). The normal Hertz force F_N depends on contact deflec-
 110 tion h as

$$111 F_N = \frac{\tilde{E}\sqrt{a}}{3}h^{3/2}, \quad (1)$$

112 in which we introduced notation $\tilde{E} = E/(1 - \nu^2)$. The adopted simplified form of tangential
 113 elasticity (Agnolin and Roux 2007a) involves a constant ratio $(2 - 2\nu)/(2 - \nu)$ of tangential
 114 (K_T) to normal (K_N) stiffnesses in contacts. Both depend on F_N , as, from (1), $K_N = \frac{dF_N}{dh} \propto$
 115 $F^{1/3}$. With such laws one should avoid spurious creation of elastic energy, and therefore K_T
 116 is suitably rescaled in cases of decreasing normal force (Elata and Berryman 1996). Details
 117 on the elastic model, on the enforcement of the Coulomb condition,

$$118 \|\mathbf{F}_T\| \leq \mu F_N, \quad (2)$$

119 and on the objective implementation of the force law, with due account of all possible motions
 120 of a pair of contacting grains, are given by Agnolin and Roux (2007a). Our simulations are
 121 carried out with the elastic properties of glass beads ($\tilde{E} = 77$ GPa) and the intergranular
 122 friction coefficient, μ , is kept equal to 0.3 in the present study.

123 Estimating the typical contact deflection under confining stress P leads to the definition
 124 of a dimensionless stiffness parameter κ (Radjaï and Dubois 2011), such that $h/a \propto \kappa^{-1}$.
 125 For a Hertzian contact, one may use (Agnolin and Roux 2007a)

$$126 \kappa = \left(\frac{\tilde{E}}{P}\right)^{2/3}. \quad (3)$$

127 In our shear test simulations we keep $\kappa = 8400$, corresponding to glass beads under pressure
 128 100 kPa. It is deemed large enough to approach the limit of rigid grains with good accuracy.

129 Viscous damping terms oppose normal relative motion of contacting grains, and are
 130 chosen to correspond to a restitution coefficient close to zero in normal collisions. Such terms
 131 were shown (da Cruz et al. 2005; Peyneau and Roux 2008a) to have negligible influence in
 132 the slow compression steps and shear flows of the present study.

133 **Force Model : Capillary Attraction**

134 An interstitial wetting liquid, introduced in small amounts, preferentially localizes at
 135 contacts or between close neighboring grains, forming liquid bridges transmitting capillary
 136 forces. Such a liquid bridge, or meniscus, is sketched in Fig. 1. We consider a perfectly wet-
 137 ting liquid, with contact angle θ equal to zero. In accordance with some observations (Her-
 138 minghaus 2005), we assume that the menisci only form as particles come into contact. If
 139 contacting grains move apart from one another, and are separated by distance h , the liquid
 140 bridge remains stable, transmitting an h -dependent force, until the gap, h , reaches a certain
 141 rupture distance D_0 , as observed in (Kohonen et al. 2004). D_0 relates to meniscus volume
 142 V as $D_0 \simeq V^{1/3}$ (Lian et al. 1993; Willett et al. 2000; Pitois et al. 2000).

143 The attractive force between particles separated by distance $h \leq D_0$ is modeled within
 144 the Maugis approximation (Maugis 1987), appropriate for small enough meniscus volume.
 145 The maximum attractive force (tensile strength) is reached for contacting particles, and
 146 equal, according to this model, to $F_0 = \pi a \Gamma$ (Γ is the liquid surface tension) independent of
 147 the meniscus volume. The capillary force varies with gap h between particle surfaces as

$$148 \quad F^{\text{Cap}} = \begin{cases} -F_0 & h \leq 0 \\ -F_0 \left[1 - \frac{1}{\sqrt{1 + \frac{2V}{\pi a h^2}}} \right] & 0 < h \leq D_0 \\ 0 & h > D_0 \end{cases} \quad (4)$$

149 (One should note that $h < 0$ corresponds to an elastic deflection of the particles in contact,

150 and keep in mind that a nonvanishing distant force, $F^{\text{Cap}}(h)$ with $h > 0$, is only possible
151 if the grains have been in contact and did not move apart farther than distance D_0 ever
152 since). This formula is a simpler, analytical form of the toroidal approximation with the
153 “gorge method” (Lian et al. 1993) for the capillary force in a meniscus. Alternative forms
154 of the attractive force law (Willett et al. 2000; Soulié et al. 2006; Radjaï and Richefeu 2009)
155 might actually be more accurate. We found (Khamseh et al. 2015) that the macroscopic
156 results were not affected upon changing the force law, were very moderately influenced
157 by saturation within the pendular range, but did significantly change upon suppressing the
158 capillary hysteresis (i.e., assuming menisci to form as soon as a pair of grains approach within
159 rupture distance D_0). It is important to recall that the Coulomb inequality, as written in (2),
160 applies to the elastic component of the normal force only, to which the negative (attractive)
161 capillary term should be added. Thus, in an isolated grain pair bonded by a meniscus,
162 at equilibrium the repulsive elastic force is equal to F_0 , and the contact may transmit a
163 tangential force at most as large as μF_0 .

164 The morphology of partially saturated granular materials depends on the liquid con-
165 tent (Mitarai and Nori 2006; Kudrolli 2008). The present study, like a number of previous
166 ones (Richefeu et al. 2006; Radjaï and Richefeu 2009; Scholtès et al. 2009a), is restricted to
167 the *pendular state* of low saturations, in which the wetting liquid is confined in bonds or
168 menisci joining contacting grains. Liquid saturation S is defined as the ratio of liquid volume
169 Ω_l to interstitial volume Ω_v , the total system volume being denoted as Ω . Writing Ω_g for the
170 volume of all N solid grains in the system (such that $\Omega = \Omega_g + \Omega_v$, S is related to meniscus
171 volume V , solid fraction $\Phi = \Omega_g/\Omega = 1 - \Omega_v/\Omega = \frac{N\pi a^3}{6\Omega}$ and wet coordination number z (the
172 average number of liquid bonds on one grain). As the liquid volume is $\Omega_l = \frac{zNV}{2}$, one has

$$173 \quad S = \frac{\Omega_l}{\Omega_v} = \frac{zNV}{2(1 - \Phi)\Omega} = \frac{3z}{\pi} \frac{\Phi}{1 - \Phi} \frac{V}{a^3}. \quad (5)$$

174 In our study, we fix the value of meniscus volume V , equal to $10^{-3}a^3$ in all results presented

175 in this paper. Such a choice does not conserve the total liquid volume (proportional to the
 176 varying coordination number z of liquid bonds) – but this is, as we could check (Khamseh
 177 et al. 2015), quite an innocuous drawback.

178 The pendular state to which our model applies is relevant in some low (but not too
 179 small) saturation range. The upper saturation limit for the pendular state corresponds to the
 180 merging of the menisci pertaining to the same grain, which, considering a triangle of spherical
 181 grains in mutual contact, happens as soon as the filling angle φ (see Fig. 1) reaches $\pi/6$. The
 182 analytical formula for V (Lian et al. 1993), within the toroidal approximation, as a function
 183 of φ (setting $h = 0$, and $\theta = 0$), then yields $\frac{V}{a^3} \simeq 8.10^{-3}$. Eq. (5) then predicts a maximum
 184 saturation between 0.05 and 0.1, similar to experimental observations (Herminghaus 2005;
 185 Mitarai and Nori 2006). On the other hand, the minimum saturation S_{\min} for bridges to
 186 form and join neighbouring grains might be roughly estimated upon introducing a roughness
 187 scale δ , assuming a liquid layer of thickness δ coats the surface of the grains, as

$$188 \quad S_{\min} = \frac{6\Phi\delta}{(1 - \Phi)a}. \quad (6)$$

189 For $\Phi = 0.5$ and $\delta \sim 10^{-4}a$, S_{\min} is of the order of 10^{-3} , as observed in experiments (Her-
 190 minghaus 2005). Using (5), and typical values of z and Φ , this sets a lower bound to meniscus
 191 volume, of order $10^{-4}a^3$.

192 Capillary attraction introduces force scale F_0 in the model, whence the definition of a
 193 reduced pressure, comparing applied stress P (isotropic pressure in compression, or controlled
 194 normal stress $\sigma_{22} = P$ for shear flows) to contact tensile strength F_0 , as

$$195 \quad P^* = \frac{a^2 P}{F_0} = \frac{aP}{\pi\Gamma}. \quad (7)$$

196 As observed with different cohesive granular models, mostly in two dimensions (2D) (Gilabert
 197 et al. 2007), we expect strong effects of cohesive forces, possibly very loose equilibrium
 198 microstructures for $P^* \ll 1$, while the properties of cohesionless systems are retrieved in the

199 limit of large P^* . Our simulations are carried out with spheres of diameter $a = 0.115$ mm,
200 perfectly wet by water, with surface tension $\Gamma = 7.3 \times 10^{-2}$ J.m⁻². For such parameter
201 values, one has $P^* = 1$ for a pressure P equal to 2 kPa. While this is admittedly a rather
202 low pressure for most geotechnical applications, it might be relevant in other fields (e.g.,
203 in some chemical engineering process), and the results are also, beyond wet grain models,
204 more generally indicative of the influence of attractive forces of small range in granular
205 assemblies. Another important issue is the possible influence of the initial microstructure
206 assembled under low P^* on the material properties under larger confining stresses. In the
207 following most results are expressed in terms of dimensionless control parameters.

208 **ISOTROPIC ASSEMBLY AND COMPRESSION**

209 We studied the important irreversible configuration changes entailed by a pressure cycle
210 starting with a low value of P^* , of order 10^{-2} or 10^{-3} , with an initial state that cannot
211 be observed without cohesive forces. Maximum pressures are such that $P^* \gg 1$. Our
212 parameter choice is such that $\kappa = 114000$ for $P^* = 1$ and $\kappa = 5300$ for $P^* = 100$, which is
213 still high enough a value for contact deflections to be irrelevant (Roux and Chevoir 2011).
214 Consequently, our results, if expressed as dimensionless quantities functions of P^* (κ being
215 large enough to be irrelevant), may apply to systems of wet spherical grains with arbitrary
216 diameter, liquid surface tension and wetting angle.

217 **Loose initial states**

218 Previous 2D simulations of cohesive systems by Gilabert et al. (2007) made it clear
219 that cohesive forces play an important part in the assembling stage. It is thus necessary to
220 assemble wet grains, rather than introduce liquid bridges into previously assembled dry grain
221 configurations. As in the 2D studies, we found it possible to assemble low density initial
222 configurations by the following procedure. First, disordered assemblies of grains (comprising
223 4000 particles) with solid fraction Φ_0 are prepared, using either random insertion or crystal
224 melting with event-driven, energy-preserving dynamics (Agnolin and Roux 2007a). Then,
225 particles are attributed random velocities, drawn according to a Gaussian distribution with

226 zero mean and variance V_0^2 , and left to interact with the force laws introduced in the previous
 227 section, within a periodic simulation cell of fixed size and shape. Capillary attraction induces
 228 the formation of clusters of grains joined by liquid bridges, and this step of the calculation is
 229 stopped when all particles belong to one single such cluster, and the configuration is regarded
 230 as sufficiently equilibrated. (Tolerances on equilibrium requirements are expressed in terms
 231 of typical contact force F_1 as, $10^{-4}F_1$ for force balance, or $10^{-4}aF_1$ for torque balance – with
 232 $F_1 = F_0$ in the assembling stage). The structure of such initial configurations depends on
 233 imposed solid fraction Φ_0 , which we chose equal to 0.3 in the main simulation series. It
 234 also depends on velocity V_0 . The latter should be compared to the characteristic velocity
 235 $V^* = \sqrt{\frac{D_0 F_0}{m}}$, which is proportional to the relative velocity that is necessary to separate
 236 a pair of grains, overcoming the potential energy of capillary force (4). Here, choosing
 237 $V_0/V^* = 0.2$, we could observe that the results corresponded to a low initial agitation limit.
 238 An important variable characterizing equilibrium configurations, the contact coordination
 239 number, z_c , is then barely larger than 4 – the isostatic (barely rigid) value (Agnolin and
 240 Roux 2007a). The coordination number of distant interactions, i.e., the average number of
 241 non-contacting neighbors connected to one grain through a capillary meniscus, which we
 242 denote as z_d , is equal to zero. This latter observation is explained by the capillary hysteresis:
 243 liquid bonds without contact only exist in pairs that have been in contact in the past. Thus,
 244 for low enough initial agitation velocity V_0 , contacts, once formed in the assembling stage,
 245 do not break in the constant density aggregation stage.

246 **Compression curves**

247 *Loading procedure and measurements*

248 To study the compression of initially loose configurations, stabilized at $\Phi_0 = 0.3$ thanks to
 249 adhesive forces, a loading program is applied in which the isotropic pressure, P , is stepwise
 250 incremented, from a low value (corresponding to $P^* = 10^{-3}$) up to $P^* = 10000$. Steps
 251 are uniform on a logarithmic scale (i.e., pressure is multiplied by $10^{1/4}$ at each step). For
 252 each new value of applied pressure P , one waits for equilibrium to be approached with good

253 accuracy, and records the new configuration. To appreciate the irreversibility of the observed
254 evolution, the loading program is a compression cycle with a decompression branch, on which
255 the previously applied pressure levels are retraced back, down to the initial small value. This
256 paper being only a brief account of salient behaviours in compression and in quasistatic shear
257 flow, we do not present here a complete study of all properties and internal states of P^* -
258 dependent isotropic configurations. We mainly focus on solid fraction Φ , or, according to
259 the soil mechanics presentation, on void ratio $e = -1 + 1/\Phi$, and on contact (z_c) and distant
260 (z_d) coordination numbers. A more complete parametric study (dependence on velocity
261 V_0 , meniscus volume, or various aspects of interaction laws such as capillary hysteresis and
262 possible rolling friction) is also postponed to a forthcoming, more detailed publication (Than
263 et al. 2015).

264 *Irreversible compression*

265 Fig. 2(a) shows the evolution of the void ratio in the isotropic compression of a system
266 of 4000 beads, with the characteristics as described in the previous sections. The loading
267 curve is composed of three parts: first (regime I), in range $P^* \leq 0.01$, the initial structure
268 supports the pressure increase, and void ratio e hardly departs from its initial value (equal to
269 2.33). Then, in a second stage (regime II), extending up to $P^* \simeq 10$, the system undergoes
270 a fast compression, which becomes considerably slower at high pressures (regime III). On
271 reducing P^* , only the density change occurring within regime III is reversed. As apparent
272 on the second graph, Fig. 2(b), in which the reference wet system is compared to a dry
273 (cohesionless) assembly of identical grains, regime III is parallel to the compression curve
274 of dry grains, in which the small compression is due to contact elasticity, and, as shown by
275 Agnolin and Roux (2007b), nearly reversible (in terms of density at least). Thus regime III
276 marks the end of the plastic collapse of the loose structure stabilized by capillary forces.

277 The plastic compression behavior of the wet material is closely similar to the 2D results
278 of Gilabert et al. (2008), and the void ratio curve in regime II might be represented with
279 a linear variation with $\log P^*$, assuming e_{ref} is the void ratio for some reference reduced

280 pressure, P_{ref}^* :

$$281 \quad e - e_{\text{ref}} = -\lambda \log \frac{P^*}{P_{\text{ref}}^*}. \quad (8)$$

282 Coefficient $\lambda \simeq 0.36$ successfully describes the curve in interval $0.04 \leq P^* \leq 2$. Eq. 8 is clas-
283 sically used in soil mechanics for cohesive systems (Mitchell and Soga 2005). Upon unloading
284 and reloading at various pressure levels along the compression curve, it is observed (as shown
285 in Fig. 3) that a plastic response (irreversible structural rearrangement with density change)
286 under isotropic pressure will be observed only if the maximum pressure the system has been
287 subjected to in the past (the “overconsolidation pressure” of soil mechanics) is exceeded.
288 This maximum pressure value appears to fully characterize the history dependence of the
289 system in isotropic compression.

290 *Coordination numbers*

291 In the compression cycle, the coordination numbers of contact (z_c) and distant (z_d) inter-
292 actions are shown in Fig. 4. Compared to density changes, those of the contact coordination
293 number are remarkably small, as it increases from nearly 4 to about 4.8 after the full pres-
294 sure cycle. As to the coordination number of menisci between distant grains, it starts at
295 zero, due to the absence of contact opening in the assembling stage with small agitation
296 velocity. Its increase to 2 in the course of irreversible compression signals the failure of the
297 contact structure: the network gets rearranged as old contacts break, and are replaced by
298 menisci connecting receding grain pairs, and new contacts form. The small change in z_c is
299 the net effect of contact creations and destructions. The final increase of z_c , accompanied by
300 a decrease of z_d , in regime III (high pressures) is caused by the elastic compression, closing
301 the gaps between non-contacting pairs, as in dry systems (Agnolin and Roux 2007b). We
302 could check that the coordination numbers remain very nearly constant along the reversible
303 paths of Fig. 3, which corresponds to unperturbed contact and meniscus networks.

304 Because of the capillary hysteresis of meniscus formation and breakage, only a proportion
305 of neighbour pairs within rupture distance D_0 (defined in the presentation of the force model,

306 in connection with Eq. 4), are connected by a liquid bridge. This proportion is initially zero,
307 it increases with P^* , peaking at 70% in regime II, decreasing to about 55% in the denser
308 systems (similar to the value $\sim 50\%$ reported in the experiments by Kohonen et al. (2004)).

309 *Effect of drying or of saturating*

310 In practice, one may act on a wet system by changing its saturation. The most drastic
311 change should be obtained on entirely suppressing the capillary cohesion, either by drying,
312 or by completely saturating the intergranular voids by the liquid. In numerical simulations,
313 one may simply remove all capillary forces, leaving only the interactions present in a dry
314 system. It is interesting to observe the effects of such an ideal transformation, carried out
315 at various points along the irreversible compression curve. Fig. 5 shows the resulting void
316 ratio evolution, if the system is deprived of capillary forces immediately before unloading at
317 different pressure levels. This ideal drying or saturation step produces a sudden collapse (a
318 brutal compression step), unless all irreversible compression has already taken place (as for
319 points F and G on Fig. 5). (More gradual collapse due to progressive imbibition is reported
320 in some experiments (Bruchon et al. 2013b)). In such a case, one may remove all capillary
321 forces, as their mechanical role, at high P^* , is negligible. Remarkably, the final state after
322 decompression keeps the same density, whether or not the system has been deprived of
323 capillary cohesion.

324 **DENSE SHEAR FLOW AND CRITICAL STATE**

325 **Model Parameters**

326 The results reported here pertain to the same reference systems studied in isotropic
327 compression, with $N = 4000$, friction coefficient $\mu = 0.3$ in the contacts, meniscus volume
328 $V = 10^{-3}a^3$. While stiffness number κ is fixed, reduced pressure takes values $P^* = \infty$ (i.e.,
329 the dry case), 10, 5, 2, 1, 0.436 and 0.1. The investigated range of I values (varying from
330 10^{-4} to 0.562 by factors of $\sqrt{10}$) enables an accurate determination of the quasistatic limit,
331 as well as an assessment of inertial effects.

332 We focus on situations in which a uniform steady state might be identified under constant
333 macroscopic shear rate $\dot{\gamma}$, after a transient stage (of a few unit strains at most). This turns out
334 to exclude small values of P^* : we could record homogeneous state parameters for $P^* = 0.436$,
335 but only partial information was gathered on the material state under $P^* = 0.1$, since such
336 systems flow inhomogeneously, localizing the velocity gradient within a narrow shear band
337 (save for a restricted range of I values of order 0.01).

338 Constitutive laws

339 Restricting their measurement to the higher values of P^* , we measure the (apparent)
340 macroscopic friction coefficient $\mu^* = \frac{\sigma_{12}}{\sigma_{22}}$, and solid fraction Φ , in steady homogeneous shear
341 flows, with the results shown in Fig. 6. As in some published results (Rognon et al. 2006;
342 Rognon et al. 2008), obtained in 2D with a model of cohesive disks, lower P^* values increase
343 μ^* and decrease the density of the sheared material. One may note, though, that the effect
344 on μ^* is considerably larger in the 3D assembly of wet particles: even for $P^* = 1$, when the
345 attractive forces and the confining ones are of similar magnitude, the quasistatic internal
346 friction coefficient, compared to its dry value ($P^* = \infty$), $\mu_0^\infty = 0.335$, nearly doubles, at
347 about 0.61. It reaches 0.867 ± 0.003 at $P^* = 0.436$. In addition to the values shown in the
348 figure, limited data are available for $P^* = 0.1$, in a range of I (of order 0.01) for which shear
349 banding does not occur. Such I values are close enough, judging from the I dependence of
350 μ^* at different P^* levels, to the quasistatic limit of $I \rightarrow 0$. We could then measure $\mu^* \simeq 1.62$
351 and $\Phi \simeq 0.435$.

352 As suggested by Rognon et al. (2006), for each P^* , a power law fit can describe the I
353 dependence of μ^* and Φ , as in a number of studies of dry granular flows (Hatano 2007;
354 Peyneau and Roux 2008a):

$$355 \mu^* = \mu_0^* + CI^\alpha \quad (9)$$

356 The data are compatible with a P^* -independent value of exponent α , $\alpha \simeq 0.8$, while coeffi-
357 cient C decreases for smaller P^* . In comparison with μ^* , the solid fraction changes relatively

358 little as a function of both P^* and I in the investigated range. Both quantities tend to depart
 359 slower from their quasistatic limit for $I \rightarrow 0$ as cohesive effects get stronger (for smaller P^*).

360 Void ratios $e_0 = -1 + 1/\Phi_0$ in the P^* -dependent critical states are compared to the
 361 values $e_{\text{iso}}(P^*)$ obtained in direct isotropic compression (normally consolidated states) in
 362 Fig. 7. The difference $e_{\text{iso}} - e_0$ is a decreasing function of P^* , but remains positive, and
 363 critical states are denser than isotropically compressed ones (this also applies to dry grains,
 364 $P^* = \infty$ – see the inserted subplot of Fig. 7). All normally consolidated isotropic states
 365 should therefore be regarded as loose: they have to contract under shear before approaching
 366 the critical state.

367 **Internal States, Microscopic Aspects**

368 *Coordination numbers*

369 Coordination numbers are shown in Fig. 8, as functions of P^* for different I , showing
 370 a quasistatic limit to be closely approached at small I . A comparison to isotropic states
 371 obtained in compression (Fig. 4) reveals, unlike for the density, quite similar values of both
 372 z_c and z_d at given P^* . The number of contacts does not change much with P^* , while the
 373 number of distant interactions tends to increase with P^* . Faster flows (larger I values) tend
 374 to break contacts, which results in smaller z_c values, an effect partly compensated by the
 375 increase of z_d : menisci survive contact openings with separation distances below D_0 .

376 *Contributions to stresses*

377 Throughout the studied parameter range, stresses in the flow are dominated by force
 378 contributions:

$$379 \quad \sigma_{\alpha\beta} = \frac{1}{\Omega} \sum_{i < j} F_{ij}^{\alpha} r_{ij}^{\beta}, \quad (10)$$

380 the sum running over all pairs i, j of grains interacting by force F_{ij} , r_{ij} pointing from the
 381 center of i to the center of j . This suggests a decomposition into contributions of different
 382 interactions. One may, e.g., isolate the contribution of distant interactions ($\sigma_{\alpha\beta}^d$) and contact
 383 interactions ($\sigma_{\alpha\beta}^c$), the latter being split into the contributions of tangential forces ($\sigma_{\alpha\beta}^T$) and

384 normal ones ($\sigma_{\alpha\beta}^{\text{N,c}}$):

$$385 \quad \sigma_{\alpha\beta} = \sigma_{\alpha\beta}^{\text{c}} + \sigma_{\alpha\beta}^{\text{d}} = \sigma_{\alpha\beta}^{\text{N,c}} + \sigma_{\alpha\beta}^{\text{T}} + \sigma_{\alpha\beta}^{\text{d}} \quad (11)$$

386 Alternatively, one may split force F_{ij} into its tangential and normal components, and
 387 isolate, in the latter, the capillary force from the elastic one. This results in a decomposition
 388 of stresses into the contribution $\sigma_{\alpha\beta}^{\text{Ne}}$ of normal elastic forces in contacts, the one of tangential
 389 contact forces, $\sigma_{\alpha\beta}^{\text{T}}$, and that of capillary forces, $\sigma_{\alpha\beta}^{\text{cap}}$, the latter incorporating both contacts
 390 and distant interactions through liquid bridges:

$$391 \quad \sigma_{\alpha\beta} = \sigma_{\alpha\beta}^{\text{Ne}} + \sigma_{\alpha\beta}^{\text{T}} + \sigma_{\alpha\beta}^{\text{cap}}. \quad (12)$$

392 To understand the large values of σ_{12} observed at small P^* , one may use Eq. 11, in which
 393 all terms of the sum have the same sign. Distant interactions contribute at most (for small
 394 P^*) 8% of the sum. Tangential forces account for about 18% of the total at $P^* = 0.436$,
 395 decreasing to 10% for $P^* = 10$. Thus the essential contribution to shear stress is that of
 396 normal contact forces, $\sigma_{12}^{\text{N,c}}$.

397 For normal stresses, it is instructive to use decomposition (12). Capillary forces are
 398 attractive, and thus contribute negatively to $\sigma_{22} = P$, as shown in Fig. 9. As the contribution
 399 of tangential forces is vanishingly small, normal elastic forces have to compensate the effect
 400 of σ_{22}^{cap} , whence $\frac{\sigma_{22}^{\text{Ne}}}{\sigma_{22}} > 3$ for $P^* = 0.436$. Remarkably, the contribution of capillary forces to
 401 shear stress, which is also opposite to that of normal repulsive forces, remains modest: $\frac{\sigma_{12}^{\text{cap}}}{\sigma_{12}}$
 402 evolves from about -0.12 at $P^* = 0.436$ to -0.03 at $P^* = 10$.

403 Following a number of recent micromechanical studies of granular materials (Peyneau
 404 and Roux 2008b; Azéma and Radjaï 2014), one may relate stresses $\sigma_{\alpha\beta}^{\text{N}}$ due to normal forces
 405 to fabric and force anisotropy parameters, an approach that we do not pursue any further
 406 here (more indications are provided by Khamseh et al. (2015)).

407 *Age of contacts and menisci.*

408 One remarkable feature by which systems with capillary cohesion differ in shear flow from
409 dry granular assemblies is the distribution of contact and interaction ages, expressed in terms
410 of strain intervals. Thus Fig. 10 shows that the same pairs of grains may stay in interaction,
411 joined by a meniscus, over several units of strain, the more often the lower P^* . Those
412 distribution functions decay exponentially for large values, with a characteristic time growing
413 from $1.1\dot{\gamma}^{-1}$ for $P^* = 10$ to $1.7\dot{\gamma}^{-1}$ for $P^* = 0.436$ – contrasting with the corresponding decay
414 time for the *contact* age distribution in a dry cohesionless system ($P^* = \infty$), which is about
415 $0.12\dot{\gamma}^{-1}$ for $I = 0.1$. Interestingly, contact lifetimes might also exceed a few strain units
416 but are considerably smaller, and, unlike meniscus lifetimes, decrease for increasing I in
417 the investigated range. Whereas pairs of dry grains tend to come into contact if oriented
418 within the compression quadrants of the shear flow, and then separate once in the extension
419 quadrant, grains connected by liquid bonds tend to form clusters that survive tumbling
420 motions in the average shear flow over notable strain intervals. Upon increasing I , although
421 contacting pairs separate more easily, they tend to remain joined by menisci. Qualitatively,
422 such a feature might explain the slow I dependence of μ^* and Φ in strongly cohesive systems,
423 and should be related to the reduced fabric anisotropy at small P^* , as well as to the influence
424 of meniscus volume or force range D_0 (Khamseh et al. 2015).

425 **Effective Pressure Approach**

426 Although the discussion of the different contributions to shear stress σ_{12} does not lead to
427 an explanation of the observed large values of μ^* , the large tensile contribution of capillary
428 forces to normal stress (Fig. 9) provides a clue. One may write

$$429 \quad \sigma_{22}^{\text{cap}} = -\beta\sigma_{22}, \quad (13)$$

430 with a coefficient β ranging, in the quasistatic limit, from about 0.15 ($P^* = 10$) to 2.1
431 ($P^* = 0.436$). (If the result for $P^* = 0.1$ and $I \sim 0.01$ is added β then reaches about 7.2).

432 Incidentally, the independence of coefficient β on inertia parameter I for $I \leq 0.1$ confirms
 433 that the rheological effect of liquid bonds is not easily disrupted by collisions in the presence
 434 of moderate inertial effects, as noticed from the distribution of their ages in the previous
 435 section.

436 One may invoke the concept of *effective pressure* to describe the effect of capillary forces
 437 on the shear resistance of the material: the attractive forces create larger repulsive elastic
 438 reactions in the contact, corresponding to an effective pressure equal to $(1 + \beta)\mathcal{P}$. (Note
 439 that one ignores here the small contribution of capillary forces to shear stress). One assumes
 440 then that the shear behavior of the material is identical to that of a dry material under
 441 such effective normal stress σ_{22}^{eff} . This approach leads to the following prediction for the
 442 P^* -dependent quasistatic friction coefficient μ_0^* :

$$443 \quad \mu_0^* = (1 + \beta)\mu_0^\infty, \quad (14)$$

444 in which μ_0^∞ denotes the quasistatic internal friction coefficient for dry grains, $P^* = \infty$.

445 The performance of the simple effective pressure prediction for the P^* dependence of μ_0^*
 446 is visualized in Fig. 11. Although the global increase of μ_0^* is captured, it is overestimated for
 447 the smallest P^* values ($P^* = 1$ and below). The relative error in the prediction of μ_0^* , using
 448 the exact, measured value of β , is about 5% at $P^* = 10$, increasing to 20% at $P^* = 0.436$ (and
 449 the value of μ_0^* for $P^* = 0.1$, about 1.62, from the data for $I \sim 0.01$ is largely overestimated,
 450 at 2.7). Thus, the effective stress approach provides a rough estimate for internal friction
 451 increase at small P^* , but becomes inaccurate in the strong cohesion regime. It cannot be
 452 exact for various reasons: while the mechanical properties are supposed to be the same once
 453 effective stresses are applied to the dry material, the density is different in the dry and
 454 the wet case (with Φ varying between 0.525 and 0.596 as P^* grows from 0.436 to infinity);
 455 capillary forces also contribute to shear stress, the force network is bound to be different,
 456 etc.

Mohr-Coulomb Model for Critical States

Coefficient β might actually be predicted as follows. From Eq. 10, one may relate (Agnolin and Roux 2007a) the average pressure, $\mathcal{P} = \text{tr}\underline{\underline{\sigma}}/3$, to the average normal force $\langle F^N \rangle$ for all interactions, and to the average, $\langle F^N h \rangle_d$, over pairs in distant interaction, of the product of force by distance $h \leq D_0$:

$$\mathcal{P} = \frac{\Phi z}{\pi a^2} \langle F^N \rangle + \frac{\Phi z_d}{\pi a^3} \langle F^N h \rangle_d \quad (15)$$

As normal stress differences are small, ratio $\frac{\mathcal{P}}{\sigma_{22}}$ only slightly differs from 1 (about 0.95) at small I . In formula 15, the second term of the r.h.s. might be neglected, as it contributes less than 2% of the pressure. Contacts (z_c , on average, per grain) carry capillary force $-F_0$, while distant forces (z_d per grain) average to a fraction of $-F_0$. From (15) the capillary contribution to pressure \mathcal{P} is bracketed as $-\frac{\Phi z F_0}{\pi a^2} \leq \mathcal{P}^{\text{cap}} \leq -\frac{\Phi z_c F_0}{\pi a^2}$, in which z denotes the total coordination number, $z = z_c + z_d$. Dividing by σ_{22} , one obtains:

$$-\frac{\Phi z}{\pi P^*} \leq \frac{\mathcal{P}^{\text{cap}}}{\sigma_{22}} \leq -\frac{\Phi z_c}{\pi P^*}. \quad (16)$$

Ignoring the small difference between \mathcal{P}^{cap} and σ_{22}^{cap} , (16) provides an estimate of coefficient β defined in (13). Thus the value of β for reduced pressure $P^* = 0.436$ is predicted between 1.9 and 2.3 (and for $P^* = 0.1$, it should reach about 8). Relation 16 also suggests that β is roughly proportional to $1/P^*$:

$$\beta \simeq b/P^*, \quad \text{with} \quad \frac{z_c \Phi}{\pi} \leq b \leq \frac{z \Phi}{\pi}. \quad (17)$$

Given the moderate variations of coordination numbers and density with P^* in the investigated range, one might choose a constant coefficient b in (17). Eq. 14, on multiplying by σ_{22} , then takes the form of a Mohr-Coulomb relation:

$$\sigma_{12} = c + \mu_1^* \sigma_{22}. \quad (18)$$

479 This relation, a classical criterion for plastic failure (Wood 1990; Biarez and Hicher 1993;
 480 Richefeu et al. 2006; Andreotti et al. 2013), defines a macroscopic cohesion c , and an internal
 481 friction coefficient μ_1^* , valid in simple shear for whatever normal stress σ_{22} . Here, assuming a
 482 constant coefficient b in (17), the Mohr-Coulomb relation is predicted to hold with the same
 483 value of internal friction as in the dry case, $\mu_1^* = \mu_0^\infty$, while macroscopic cohesion c is given
 484 by

$$485 \quad c = \frac{b\mu_0^\infty F_0}{a^2} = \frac{b\pi\mu_0^\infty\Gamma}{a} \quad (19)$$

486 This estimate of the macroscopic cohesion in the Mohr-Coulomb sense is very similar to
 487 the one obtained by Richefeu et al. (2006), by a different route. In Fig. 11, the prediction
 488 of static friction coefficient μ_0^* as a function of $1/P^*$ using β as deduced from (17), with
 489 coefficient b equal to the middle point of the specified interval, *viz.* $b = \frac{(z_c+z)\Phi}{2\pi}$, is shown to
 490 perform quite well for $P^* \geq 1$, failing at small P^* , when the effective stress approach with
 491 the exact value of β fails too.

492 In general, assuming a Mohr-Coulomb criterion for critical states to apply with a P^* -
 493 independent value of cohesion c implies, upon dividing (18) by $\sigma_{22} = P^*F_0/a^2$, that the
 494 quasistatic friction coefficient μ_0^* should vary linearly with $1/P^*$:

$$495 \quad \mu_0^* = \frac{\sigma_{12}}{\sigma_{22}} = \mu_1^* + \frac{a^2c}{F_0P^*} \quad (20)$$

496 The Mohr-Coulomb representation of yield stresses might thus be used as an approximation
 497 for $P^* \geq 1$, with $a^2c/F_0 \simeq 0.27$, but the observed sublinear increase of μ_0^* with $1/P^*$
 498 in Fig. 11 (see the result for $P^* = 0.436$, and the subplot including value $\mu_0^* \simeq 1.6$ for
 499 $P^* = 0.1$) clearly precludes the definition of unique values of macroscopic cohesion and
 500 friction coefficient according to (18) for smaller pressures.

501 CONCLUSIONS

502 We now provide a quick summary of the main results on both compression and steady
 503 shear flow, and end with a discussion, in which perspectives for future work are evoked.

504 **Isotropic Compression**

505 Model wet granular assemblies exhibit the same striking differences with cohesionless sys-
506 tems under compression that the simpler, mostly 2D models of the recent literature: stability
507 of loose structures under low P^* , plastic behaviour in isotropic compression with hardening
508 expressed by the overconsolidation pressure, linear variation of void ratio with $\log P$ in some
509 range – a qualitative behaviour common to many cohesive particulate materials. Compar-
510 isons with experimental observations are possible. Although the sensitivity of the results to
511 quite a few model features still needs to be assessed, one may tentatively conclude that the
512 final state, obtained after sufficient overconsolidation, should be independent of the initial
513 configurations and of some aspects of the compression procedure. This state is not affected
514 by the removal of capillary forces, and may be regarded as the result of an ideal, homoge-
515 neous and isotropic version of the moist tamping assembling process (Frost and Park 2003;
516 Benahmed et al. 2004). It is looser (see Figs. 2(b) and 5) than directly compressed packs of
517 dry grains, and could qualify as a reference loose state.

518 **Shear Flow and Critical States**

519 The main rheological influence of capillary adhesion on critical state and shear flow is a
520 strong increase of shear resistance (or apparent friction μ^*) as P^* decreases, even though
521 as P^* reaches values of order 0.1, the strong localization tendency hinders the identification
522 of constitutive laws for homogeneous flow. Meanwhile, density and coordination numbers
523 variations with P^* are slower. In the presence of capillary forces, clusters of particles joined
524 by liquid bridges may survive strain intervals of several units, and the compressive role of
525 attractive forces is not as immediately disturbed as the one of the externally applied normal
526 stress upon increasing shear rate and inertial effects. A simple effective stress approach
527 may quantitatively account for the shear resistance trend in good approximation as long as
528 $P^* \geq 1$. The Mohr-Coulomb criterion approximately describes critical states in the same
529 reduced pressure range, but is no longer applicable at lower P^* . Many results (regarding,
530 in particular, normal stress differences, fabric anisotropy, sensitivity to meniscus volume...)

531 are deferred to a more detailed paper submitted by some of the present authors (Khamseh
532 et al. 2015). A major concern, the shear banding instability affecting low P^* shear flows,
533 should certainly be addressed in a systematic study as well.

534 Discussion

535 Our simple model of wet grains reveals many new behaviors, compared to dry materials,
536 and provides means for a critical review of macroscopic phenomenological laws (compression
537 curve, Mohr-Coulomb criterion). While some phenomena were already explored in 2D
538 cohesive models, modeling more realistic 3D systems reveals quantitative differences (e.g.,
539 a much stronger enhancement of shear resistance), and should permit experimental con-
540 frontations (Pierrat et al. 1998; Richefeu et al. 2006). The present paper did not discuss
541 the influence of liquid saturation within the pendular range, and our model with constant
542 meniscus volume does not strictly maintain a fixed water content in the material. Such
543 issues are discussed separately for compression (Than et al. 2015) and shear flow (Khamseh
544 et al. 2015) in forthcoming publications. We could check that a correction of the model in
545 which capillary forces are slightly more accurately described does not significantly change
546 the results. Similarly, a correction of meniscus volume to ensure a constant total liquid
547 volume brings only hardly noticeable changes to compression or shear behavior. Rheological
548 properties vary moderately through the pendular range (Khamseh et al. 2015) (with a very
549 small density change and a variation of about 20% of μ_0^* at $P^* = 0.436$).

550 A more serious limitation of our model is its inability to deal with saturations exceeding
551 the pendular regime. Numerical models for higher saturation levels, resorting, e.g., to a
552 lattice Boltzmann discretization of the interstitial liquid, are currently being developed (De-
553 lenne et al. 2015). Even in the small saturation range, though, the results for compression
554 and shear show that the material behavior is considerably enriched compared to dry granular
555 systems. One obvious, broad perspective is the exploration of the large unknown territory
556 that separates isotropically assembled states from critical states, the simulation of deviatoric
557 loads and the assessment of the applicability of macroscopic models of cohesive soils. It

558 would be interesting to explore the effects of resistance to rolling, due to surface asperities,
559 on the material behavior in the presence of capillary cohesion. Even a small rolling resis-
560 tance was observed to have important effects on the behavior of cohesive systems in two
561 dimensions (Gilabert et al. 2008). It might be viewed as a first step towards the modeling of
562 non-spherical objects (Estrada et al. 2011), but the geometry of liquid bridges joining objects
563 with flat or angular surfaces might entail different force laws, and the effects of capillarity
564 should be investigated in such cases.

APPENDIX I. REFERENCES

- 565
566 Agnolin, I. and Roux, J.-N. (2007a). “Internal states of model isotropic granular packings.
567 I. Assembling process, geometry, and contact networks.” *Phys. Rev. E*, 76(6), 061302.
- 568 Agnolin, I. and Roux, J.-N. (2007b). “Internal states of model isotropic granular packings.
569 II. Compression and pressure cycles.” *Phys. Rev. E*, 76(6), 061303.
- 570 Allen, M. and Tildesley, D. (1987). *Computer simulations of liquids*. Oxford University Press,
571 Oxford.
- 572 Andreotti, B., Forterre, Y., and Pouliquen, O. (2013). *Granular Media: Between Fluid and*
573 *Solid*. Cambridge University Press, Cambridge, UK.
- 574 Azéma, E. and Radjaï, F. (2014). “Internal structure of inertial granular flows.” *Phys. Rev.*
575 *Lett.*, 112, 078001.
- 576 Azéma, E., Radjaï, F., and Dubois, F. (2013). “Packings of irregular polyhedral particles:
577 Strength, structure, and effects of angularity.” *Phys. Rev. E*, 87, 062223.
- 578 Azéma, E., Radjaï, F., and Roux, J.-N. (2015). “Internal friction and absence of dilatancy
579 of packings of frictionless polygons.” *Phys. Rev. E*, 91, 010202(R).
- 580 Benahmed, N., Canou, J., and Dupla, J.-C. (2004). “Structure initiale et propriétés de
581 liquéfaction statique d’un sable.” *Comptes-Rendus Académie des Sciences, Mécanique*,
582 332, 887–894.
- 583 Biarez, J. and Hicher, P.-Y. (1993). *Elementary Mechanics of Soil Behaviour*. A. A. Balkema,
584 Rotterdam.
- 585 Bruchon, J.-F., Pereira, J.-M., Vandamme, M., Lenoir, N., Delage, P., and Bornert, M.
586 (2013a). “Full 3d investigation and characterisation of capillary collapse of a loose unsat-
587 urated sand using x-ray ct.” *Granular Matter*, 15(6), 783–800.
- 588 Bruchon, J.-F., Pereira, J.-M., Vandamme, M., Lenoir, N., Delage, P., and Bornert, M.
589 (2013b). “X-ray microtomography characterisation of the changes in statistical homogene-
590 ity of an unsaturated sand during imbibition.” *Géotechnique Letters*, 3, 84–88.
- 591 Cumberland, D. and Crawford, R. (1987). *The Packing of Particles*. Elsevier, Amsterdam.

592 da Cruz, F., Emam, S., Prochnow, M., Roux, J.-N., and Chevoir, F. (2005). “Rheophysics
593 of dense granular materials: discrete simulation of plane shear flows.” *Phys. Rev. E*, 72,
594 021309.

595 Delenne, J.-Y., Richefeu, V., and Radjaï, F. (2015). “Liquid clustering and capillary pressure
596 in granular media.” *J. Fluid Mech.*, 762, R5.

597 Donev, A., Torquato, S., and Stillinger, F. H. (2005). “Pair correlation function character-
598 istics of nearly jammed disordered and ordered hard-sphere solids.” *Phys. Rev. E*, 71(1),
599 011105.

600 Elata, D. and Berryman, J. G. (1996). “Contact force-displacement laws and the mechanical
601 behavior of random packs of identical spheres.” *Mechanics of Materials*, 24, 229–240.

602 Estrada, N., Azéma, E., Radjaï, F., and Taboada, A. (2011). “Identification of rolling resis-
603 tance as a shape parameter in sheared granular media..” *Phys. Rev. E*, 84, 011306.

604 Forterre, Y. and Pouliquen, O. (2008). “Flows of dense granular media.” *Annu. Rev. Fluid
605 Mech.*, 40, 1–24.

606 Frost, J. D. and Park, J.-Y. (2003). “A critical assessment of the moist tamping technique.”
607 *ASTM Geotechnical Testing Journal*, GTJ20039850–261.

608 Gilabert, F. A., Roux, J.-N., and Castellanos, A. (2007). “Computer simulation of model
609 cohesive powders: Influence of assembling procedure and contact laws on low consolidation
610 states.” *Phys. Rev. E*, 75(1), 011303.

611 Gilabert, F. A., Roux, J.-N., and Castellanos, A. (2008). “Computer simulation of model
612 cohesive powders: Plastic consolidation, structural changes, and elasticity under isotropic
613 loads.” *Phys. Rev. E*, 78(1), 031305.

614 Hatano, T. (2007). “Power-law friction in closely packed granular materials.” *Phys. Rev. E*,
615 75, 060301(R).

616 Herminghaus, S. (2005). “Dynamics of wet granular matter.” *Advances in Physics*, 54, 3,
617 221–261.

618 Khamseh, S., Roux, J.-N., and Chevoir, F. (2015). “Flow of wet granular materials: a

619 numerical study.”, submitted for publication in *Phys. Rev. E*.

620 Kohonen, M. M., Geromichalos, D., Scheel, M., Schier, C., and Herminghaus, S. (2004). “On
621 capillary bridges in wet granular materials.” *Physica A*, 339, 7–15.

622 Kruyt, N. P. and Rothenburg, L. (2014). “On micromechanical characteristics of the critical
623 state of two-dimensional granular materials.” *Acta Mechanica*, 225(8), 2301 – 2318.

624 Kudrolli, A. (2008). “Granular matter: Sticky sand.” *Nature Materials*, 7, 174–175.

625 Lian, G., Thornton, C., and Adams, M. (1993). “A theoretical-study of the liquid bridge
626 forces between 2 rigid spherical bodies.” *J. Coll. Inter. Sci.*, 161, 138–147.

627 Magnanimo, V., La Ragione, L., Jenkins, J. T., Wang, P., and Makse, H. A. (2008). “Char-
628 acterizing the shear and bulk moduli of an idealized granular material.” *Europhys. Lett.*,
629 81, 34006.

630 Maugis, D. (1987). “Adherence of elastomers: Fracture mechanics aspects.” *Journal of Ad-
631 hesion Science and Technology*, 1, 105–134.

632 Mitarai, N. and Nori, F. (2006). “Wet granular materials.” *Advances in Physics*, 55, 1, 1–45.

633 Mitchell, J. and Soga, K. (2005). *Fundamentals of Soil Behavior*. Wiley.

634 O’Hern, C., Silbert, L. E., Liu, A. J., and Nagel, S. R. (2003). “Jamming at zero temperature
635 and zero applied stress: The epitome of disorder.” *Phys. Rev. E*, 68(1), 011306.

636 Peyneau, P.-E. and Roux, J.-N. (2008a). “Frictionless bead packs have macroscopic friction,
637 but no dilatancy.” *Phys. Rev. E*, 78, 011307.

638 Peyneau, P.-E. and Roux, J.-N. (2008b). “Solidlike behavior and anisotropy in rigid friction-
639 less bead assemblies.” *Phys. Rev. E*, 78, 041307.

640 Pierrat, P., Agrawal, D. K., and Caram, H. S. (1998). “Effect of moisture on the yield locus
641 of granular materials: theory of shift.” *Powder Tech.*, 99, 220–227.

642 Pitois, O., Moucheron, P., and Chateau, X. (2000). “Liquid bridge between two moving
643 spheres: An experimental study of viscosity effects.” *J. Coll. Interf. Sci.*, 231, 26–31.

644 Radjaï, F., Delenne, J.-Y., Azéma, E., and Roux, S. (2012). “Fabric evolution and accessible
645 geometrical states in granular materials.” *Granular Matter*, 14, 259–264.

646 Radjaï, F. and Dubois, F. (2011). *Discrete-element modeling of granular materials*. Wiley.

647 Radjaï, F. and Richefeu, V. (2009). “Bond anisotropy and cohesion of wet granular materi-
648 als.” *Phil. Trans. R. Soc. A*, 367, 5123–5138.

649 Radjaï, F., Troadec, H., and Roux, S. (2004). “Basic features of granular plasticity.” *Granular*
650 *Materials: Fundamentals and Applications*, S. J. Antony, W. Hoyle, and Y. Ding, eds.,
651 Cambridge, Royal Society of Chemistry, 157–183.

652 Richefeu, V., El Youssoufi, M., and Radjaï, F. (2006). “Shear strength properties of wet
653 granular materials.” *Phys. Rev. E*, 73, 051304.

654 Rognon, P., Roux, J.-N., Naaïm, M., and Chevoir, F. (2008). “Dense flows of cohesive
655 granular materials.” *J. Fluid Mech.*, 596, 21–47.

656 Rognon, P., Roux, J.-N., Wolf, D., Naaïm, M., and Chevoir, F. (2006). “Rheophysics of
657 cohesive granular materials.” *Europhysics Letters*, 74, 644–650.

658 Rothenburg, L. and Kruyt, N. P. (2004). “Critical state and evolution of coordination number
659 in simulated granular materials.” *International Journal of Solids and Structures*, 41(2),
660 5763–5774.

661 Roux, J.-N. and Chevoir, F. (2011). “Dimensional Analysis and Control Parameters.” In
662 (Radjaï and Dubois 2011), Chapter 8, 199–232.

663 Saint-Cyr, B., Szarf, K., Voivret, C., Azéma, E., Richefeu, V., Delenne, J.-Y., Combe, G.,
664 Nouguièr-Lehon, C., Villard, P., Sornay, P., Chaze, M., and Radjaï, F. (2012). “Particle
665 shape dependence in 2d granular media.” *EPL*, 98(4), 44008.

666 Scholtès, L., Chareyre, B., Nicot, F., and Darve, F. (2009a). “Micromechanics of granular
667 materials with capillary effects.” *International Journal of Engineering Science*, 47, 64–75.

668 Scholtès, L., Hicher, P.-Y., Nicot, F., Chareyre, B., and Darve, F. (2009b). “On the capillary
669 stress tensor in wet granular materials.” *Int. Journal for Numerical and Analytical Methods*
670 *in Geomechanics*, 33, 1289–1313.

671 Silbert, L. E., Ertas, D., Grest, G. S., Halsey, T. C., and Levine, D. (2002). “Geometry of
672 frictionless and frictional sphere packings.” *Phys. Rev. E*, 65(3), 031304.

- 673 Soulié, F., Cherblanc, F., El Youssoufi, M., and Saix, C. (2006). “Influence of liquid bridges
674 on the mechanical behaviour of polydisperse granular materials.” *Int. J. Numer. Anal.*
675 *Meth. Geomech.*, 30, 213–228.
- 676 Than, V.-D., Roux, J.-N., Tang, A. M., and Pereira, J.-M. (2015). “Numerical simulation of
677 plastic compression of wet granular soils.”, to be submitted.
- 678 Thornton, C. (2000). “Numerical simulations of deviatoric shear deformation of granular
679 media.” *Géotechnique*, 50, 43–53.
- 680 Willett, C., Adams, M., Johnson, S., and Seville, J. (2000). “Capillary bridges between two
681 spherical bodies.” *Langmuir*, 16 (24), 9396–9405.
- 682 Wood, D. M. (1990). *Soil Behaviour and Critical State Soil Mechanics*. Cambridge University
683 Press.

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APPENDIX II. NOTATION

685

The following symbols are used in this paper:

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 a = spherical grain diameter; α = exponent of power law expressing internal friction increase with I ; b = proportionality coefficient relating β to $1/P^*$; β = ratio of capillary contribution to total normal stress in shear flow; C = coefficient of I^α for power law increase of μ^* ; c = macroscopic cohesion according to Mohr-Coulomb criterion; D_0 = rupture distance of liquid bridge; \tilde{E} = Modulus appearing in Hertz law; e = void ratio; e_0 = void ratio in critical state; e_{iso} = void ratio in isotropic compression;

687

 F_0 = maximum capillary tensile force through liquid bridge; Φ = solid fraction; Φ_0 = solid fraction in critical state; Φ_{iso} = solid fraction in isotropic compression; φ = filling angle in liquid meniscus; Γ = surface tension of nonsaturating interstitial liquid; $\dot{\gamma}$ = shear rate; h = distance between particles or contact deflection; I = inertial number; κ = dimensionless contact stiffness number; m = grain mass;

- μ = intergranular friction coefficient;
 μ^* = apparent internal, macroscopic friction coefficient;
 μ_1^* = internal friction coefficient according to Mohr-Coulomb criterion;
 μ_0^∞ = internal friction coefficient in critical state for dry grains;
 Ω = total sample volume;
 Ω_l = liquid volume;
 P = controlled stress: isotropic pressure in compression, normal stress σ_{22} in shear flow;
 P^* = dimensionless, reduced pressure;
 \mathcal{P} = mean pressure;
 S = saturation;
 $\sigma_{\alpha\beta}$ = stress tensor;
 $\sigma_{\alpha\beta}^{\text{eff}}$ = effective stress tensor;
 $\sigma_{\alpha\beta}^{\text{cap}}$ = contribution of all capillary forces to stress tensor;
 $\sigma_{\alpha\beta}^{\text{T}}$ = contribution of tangential forces to stress tensor;
 $\sigma_{\alpha\beta}^{\text{N}}$ = contribution of normal forces to stress tensor;
 $\sigma_{\alpha\beta}^{\text{c}}$ = contribution of contact forces to stress tensor;
 $\sigma_{\alpha\beta}^{\text{Ne}}$ = contribution of normal, elastic contact forces to stress tensor;
 $\sigma_{\alpha\beta}^{\text{N,c}}$ = contribution of capillary forces in contacts to stress tensor;
 $\sigma_{\alpha\beta}^{\text{d}}$ = contribution of distant capillary forces to stress tensor;
 V = meniscus volume;
 V_0 = initial mean quadratic agitation velocity, in assembling stage;
 V^* = characteristic velocity, associated with attractive force;
 z_c = coordination number of contacts;
 z_d = coordination number of distant interactions;
 z = total coordination number.

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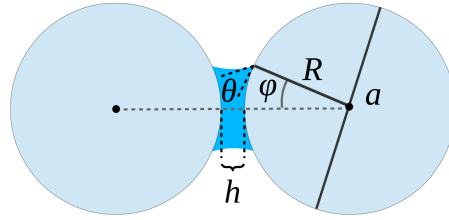


FIG. 1. A meniscus between two spherical grains of diameter $a = 2R$, with distance h between solid surfaces, filling angle φ , contact angle θ .

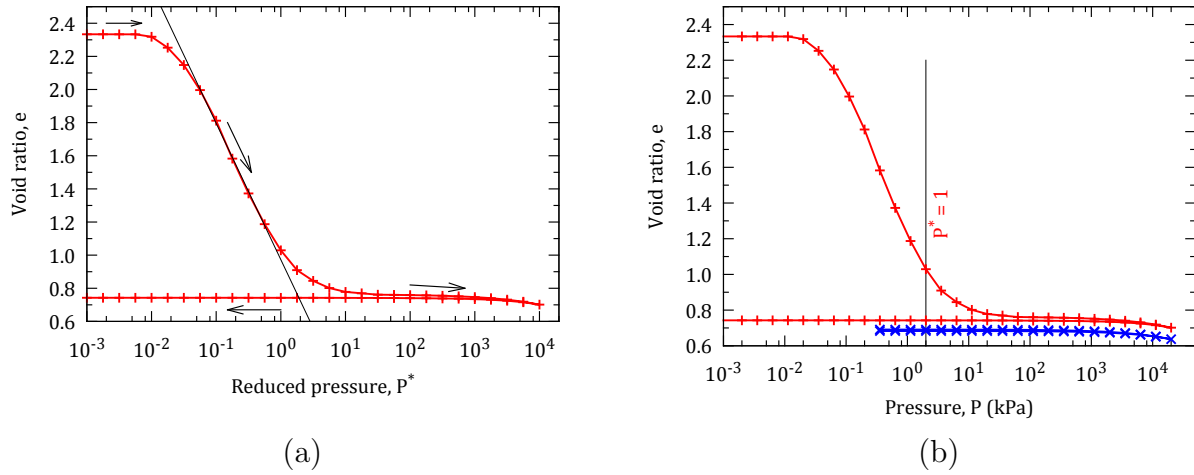


FIG. 2. (a) Effect of cycle of pressure (in dimensionless form) on void ratio. The straight line fits the curve for intermediate P^* values. (b) Same curve, compared to result obtained with dry, cohesionless grains (bottom curve, crosses joined by dark line), with pressure in kPa, for glass beads with $a = 0.115$ mm, wet by water.

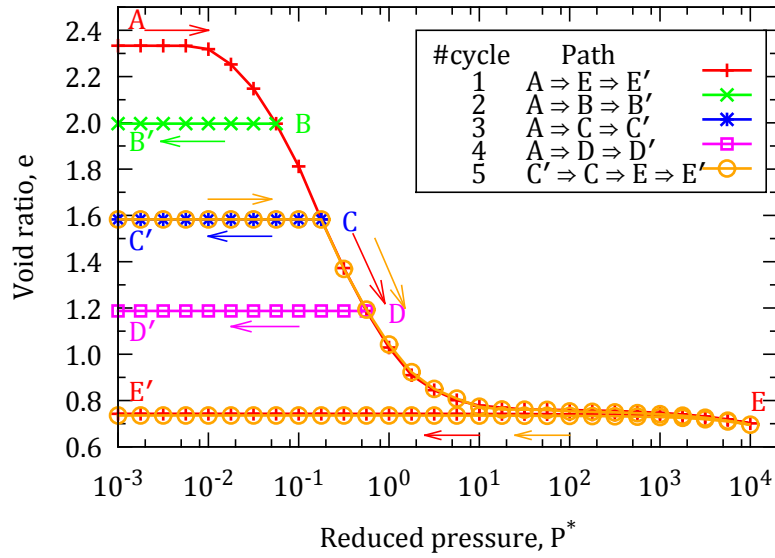


FIG. 3. Effect of different (isotropic) unloading and reloading histories on void ratio. The system does not rearrange along unloading paths BB' , CC' , DD' , EE' , which are reversible. Path 5 causes plastic response in section CE, along which pressure increases beyond its past maximum. The primary curve (path 4) is then retraced.

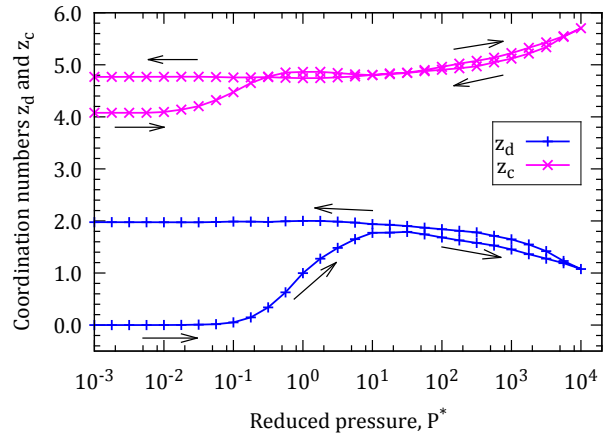


FIG. 4. Evolution of coordination numbers of contacts z_c and of distant interactions z_d , in the pressure cycle of Fig. 2(a).

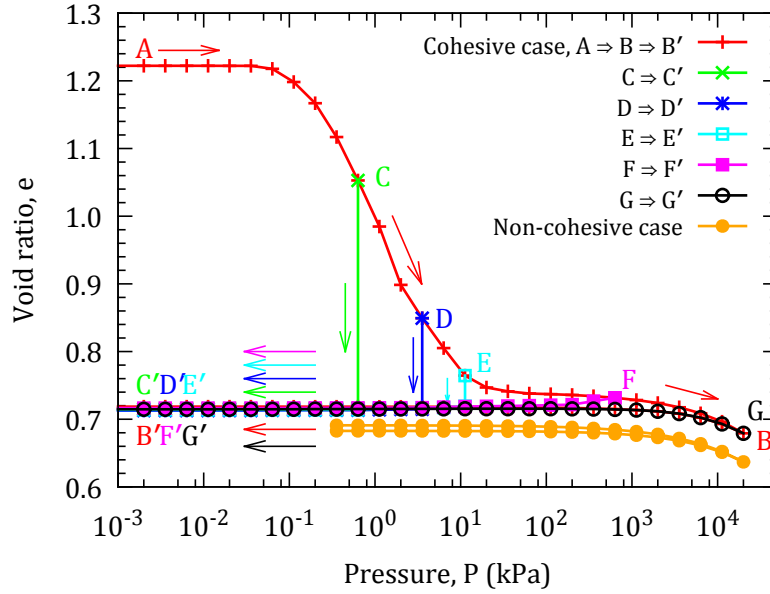


FIG. 5. Void ratio versus reduced pressure P^* , as cohesive forces are suppressed at the beginning of unloading, starting at different points on the primary compression curve. (In this case the initial state had solid fraction $\Phi_0 = 0.45$).

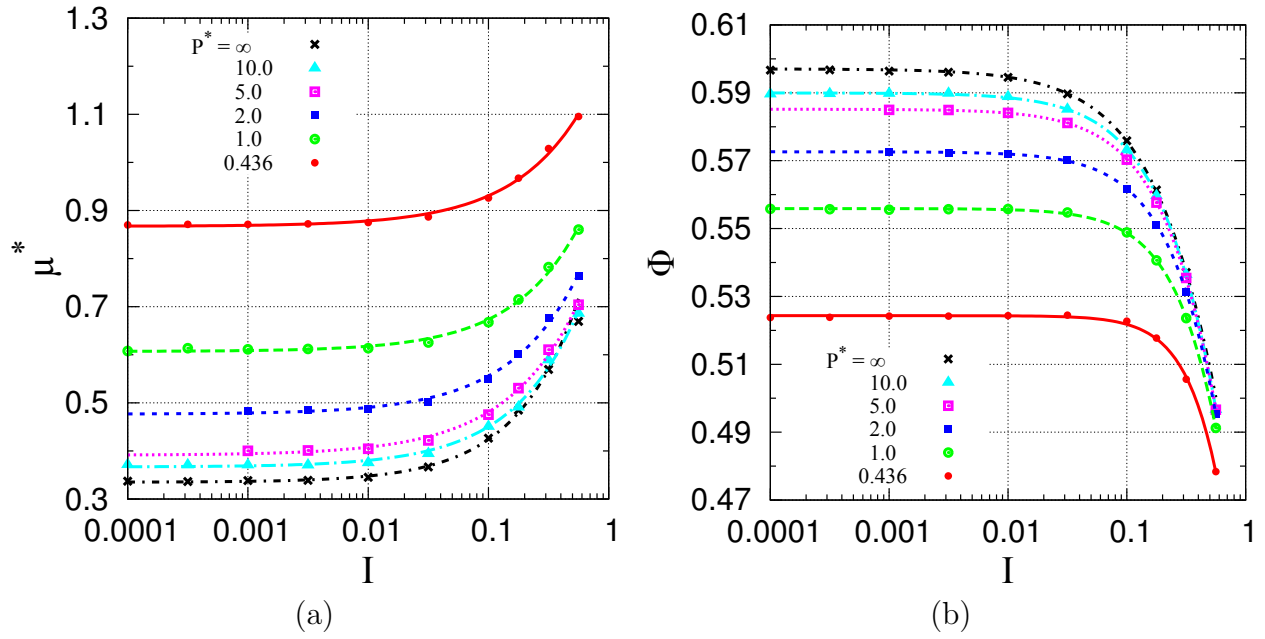


FIG. 6. (a) Internal friction coefficient μ^* and (b) solid fraction Φ versus I for different values of P^* .

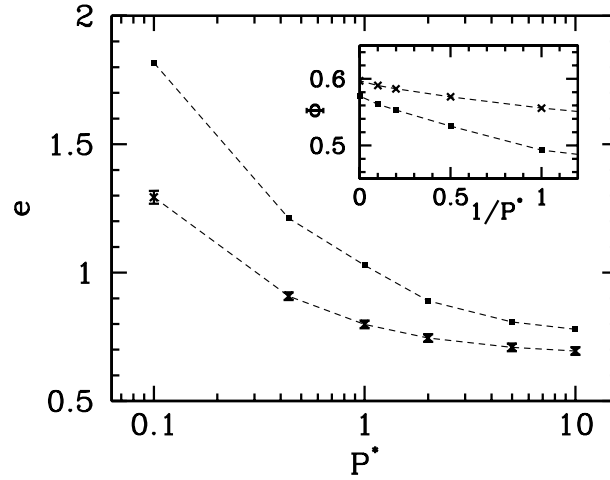


FIG. 7. Main plot: void ratios in primary isotropic compression, e_{iso} (square dots), and in critical state, e_0 (crosses with error bars) versus $\log P^*$. Inset: detail of variations of solid fraction Φ_{iso} in isotropic compression (square dots), and Φ_0 , in critical state (crosses), for large P^* (including the dry case of infinite P^*), versus $1/P^*$. Dashed lines are drawn to guide the eye.

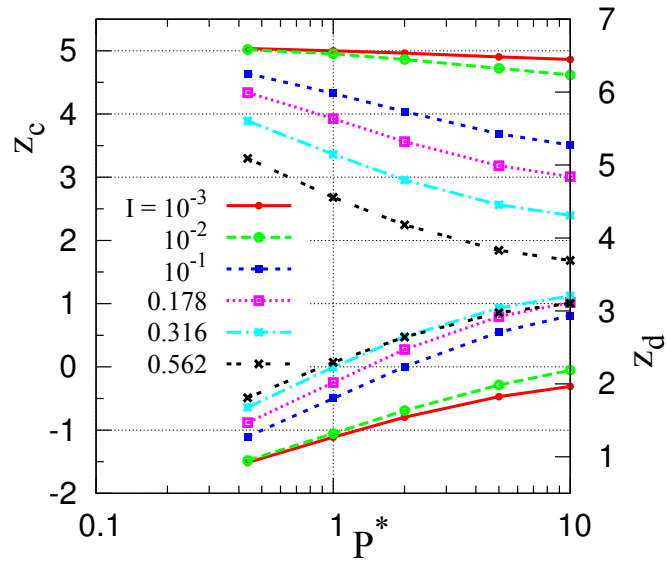


FIG. 8. Coordination numbers of contacts z_c (left axis, upper curves) and of distant interactions z_d (right axis, bottom curves), versus reduced pressure for different values of I .

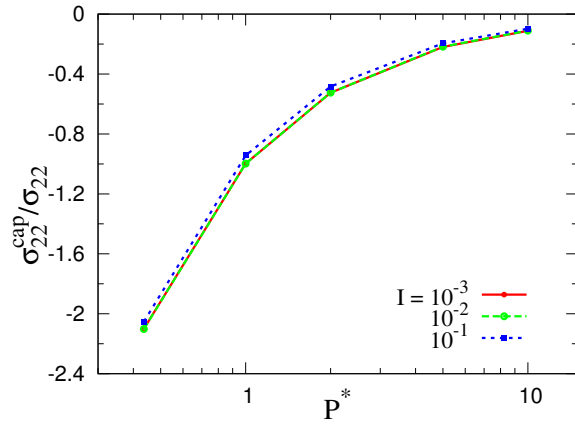


FIG. 9. Contribution of capillary forces to normal stress σ_{22} .

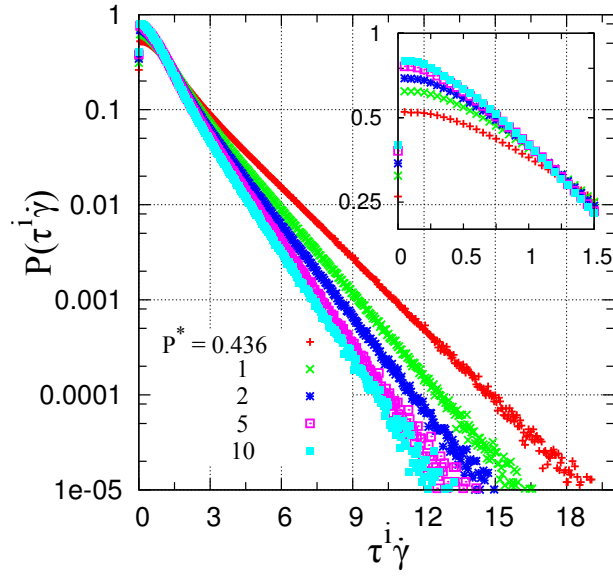


FIG. 10. Distribution of the age of menisci for different P^* values (results for different I indistinguishable on this plot). Inset shows detail of initial decay.

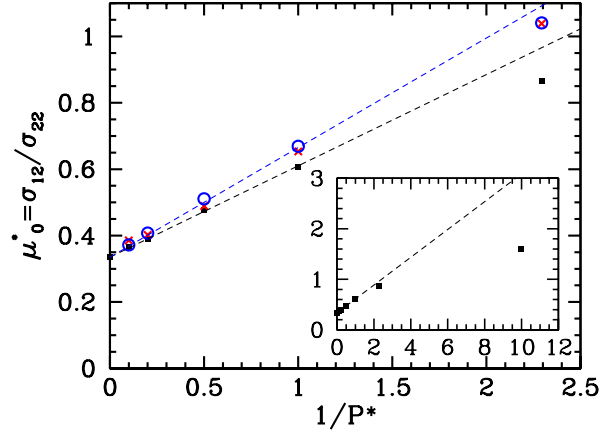


FIG. 11. Apparent quasistatic friction coefficient μ_0^* versus $1/P^*$ – showing the value of μ_0^∞ for $1/P^* = 0$. Square dots: numerical results (error bars are smaller); (red) crosses: predictions of (14), with exact coefficient β ; (blue) circles: same with estimated β . Dotted lines: Mohr-Coulomb models, predicted from (19) (upper line), or fitted to the data in range $P^* \geq 1$ (lower straight line). Inset: measured μ_0^* versus $1/P^*$, including data point for $P^* = 0.1$, with Mohr-Coulomb fit to $P^* \geq 1$ data.