


Cohesion-controlled granular materialAdrien Gans¹,* Olivier Pouliquen,[†] and Maxime Nicolas¹‡*Aix Marseille Univ, CNRS, IUSTI, Marseille, France* (Received 4 October 2019; accepted 19 February 2020; published 11 March 2020)

We present a simple method to prepare a granular material with a controlled cohesion between particles. The granular material is made of spherical glass beads coated with a polyborosiloxane polymer. This material is proved to be stable in time and nonsensitive to temperature and humidity. The interparticle force is measured and related to the size of the grain and the polymer coating thickness. Classical measurements (packing fraction, repose angle, macroscopic cohesion) are performed with this cohesion-controlled granular material. This model material opens many perspectives to study in a controlled manner the flow of cohesive grains.

DOI: [10.1103/PhysRevE.101.032904](https://doi.org/10.1103/PhysRevE.101.032904)**I. INTRODUCTION**

Handling powders and cohesive materials is an everyday challenge in many industrial processes. Beyond the difficulty of handling a very large number of particles, the cohesion between grains is a serious source of difficulty. Cohesion often comes into play when manipulating small particles, for which humidity [1,2] or attractive forces like Van der Waals [3] or electrostatic forces [4] become important. The adhesion between the grains decreases the ability of powders to flow easily, and the dynamics then differs from dry granular materials made of large particles interacting solely through frictional dry contact interactions. Whereas our knowledge of granular flows has improved over the last 20 years, the behavior of powders and cohesive granular media still represents a real challenge.

From an industrial point of view, different techniques have been developed to characterize and to quantify the “flowability” of powders [5,6]. This concept based on a compilation of different measurements (compaction, repose angles, etc.), although very useful when checking the quality of a product or when comparing different materials, lacks clear physical bases.

From a fundamental point of view, one difficulty in studying cohesive material is the control of the cohesion forces between the particles. Whereas in numerical studies, simplified interaction laws might be introduced in DEM approaches to simulated cohesive grains [7], experimentally, the control of the cohesion is more challenging. Working with actual powders faces many difficulties. Powder particles are often very polydispersed with a great variety of shape and surface roughness. A humidity and temperature control of the ambient air is required to ensure a good reproducibility of the experiments [8,9]. While some comparative tests between benchmark powders (such as lactose) are useful [10], it is still very difficult to link the particle properties to the bulk flow behavior of the powder [3], mainly because of a lack of a controlled particle interaction. Most of the recent experimental works

on cohesive granular materials have focused on capillary cohesion, studying the properties of an assembly of beads mixed with a small amount of a viscous liquid. The influence of the amount of liquid on the packing fraction [11,12], on the repose or avalanche angle [13–17], and on the stability of a column [18] has been studied. The strength of a wet granular assembly has been studied through the measurement of the shear modulus [19], and more recently, rheology experiments have been also conducted [20,21] in a pressure-imposed rheometer for glass beads coated by viscous silicon oil. Despite a good knowledge of the cohesion force arising from capillary bridges [22] the bulk behavior and the rheological measurements suggest a complex dynamics due to the evolution of the capillary bridges during shear, which may migrate and coalesce [23,24]. Recently [25], silanization has been used to modify the surface properties of glass particles and build a cohesive force of 0.1 mN order of magnitude. Experimental research on cohesive materials would strongly benefit from a model system making it possible to tune at will, and through a simple process, the cohesion between the grains.

In this paper we present a new method to prepare a cohesion-controlled granular material (CCGM) made of polyborosiloxane (PBS) coated glass particles, which suits many of the requirements to achieve experiments with a controlled cohesion. The main point is that the cohesion force between particles can be easily tuned through the PBS coating. The coating of the particles is easy and does not require heavy chemical equipment. Furthermore, the CCGM is very stable on a long timescale, is insensitive to humidity of the ambient air, and is also insensitive to room temperature.

The preparation method is first presented in Sec. II. A detailed study of the interparticle cohesive force induced by the presence of the PBS coating is presented in Sec. III. In Sec. IV the CCGM is tested in different classical configurations used for characterizing granular media: measurements of the bulk density, pile angle, and inclined plane experiments. Concluding remarks and perspectives are given in Sec. V.

II. DESIGN OF A COHESION CONTROLLED GRANULAR MATERIAL

The particle-coating material is a polyborosiloxane (PBS) made from a –OH terminated polydimethylsiloxane (PDMS)

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FIG. 1. Example of a cohesion-controlled granular material: a pile of glass beads $d = 3$ mm with a PBS coating layer of thickness $b = 2.2$ μm

cross-linked with boric acid (H_3BO_3) [26,27]. Each batch of CCGM is prepared with a mass m_g of spherical glass beads (diameter d and density $\rho_G = 2600$ kg m^{-3}) with a small polydispersity, a mass m_P of PDMS (density $\rho_P = 970$ kg m^{-3} , viscosity 750 mPa s, and gyration radius $r_g = 17$ \AA), and a mass m_A of boric acid. We kept a constant mass ratio of boric acid over PDMS $m_A/m_P = 0.14$. The boric acid is first dissolved in a small volume (50 ml) of purified water heated at 60 $^\circ\text{C}$. The particles, the PDMS, and the H_3BO_3 solution are then mixed together in a heating mixer (Kenwood Cooking Chef) at 110 $^\circ\text{C}$ over 90 min to ensure the evaporation of the water and a homogeneous PBS coating of the spherical particles. After cooling, the batch is ready to use for experiments. The PBS has been characterized in a rheometer using a plane-plane geometry, and the real and imaginary part of the shear modulus are $G' = 35$ kPa and $G'' = 7.5$ kPa with a relaxation time constant of 3.8 s.

The range of particle size was $d = [0.8\text{--}10]$ mm in experiments of Sec. III concerning the measurement of the interparticle force, and $d = [0.2\text{--}1.4]$ mm in the macroscopic experiments of Sec. IV. From atomic force microscopy imaging, the average roughness of 0.8 mm particles was 30 ± 2 nm. The effect of the PBS coating is qualitatively illustrated in Fig. 1, where a sample of 3-mm-diameter coated glass beads reveals the cohesive nature of the material. The parameter controlling the cohesion in our study is the averaged thickness b . Assuming an homogeneous coating of a $\Omega_P = m_P/\rho_P$ volume of PBS over perfectly spherical particles of volume $\Omega_G = m_G/\rho_G$ gives an average thickness

$$b = \frac{d}{2} \left[\left(1 + \frac{\Omega_P}{\Omega_G} \right)^{1/3} - 1 \right]. \quad (1)$$

In the limit of a small coating averaged thickness ($b \ll d$), the PBS volume ratio $\Omega_P/\Omega_G \approx 6b/d$. An attempt to visualize the PBS layer is presented in Fig. 2. In the pictures, the PBS appears as a gray and foamy fluid, which is optically different from the clean glass surface. The pictures of the clean surfaces have been obtained after a careful removal of the PBS with a spray of heptane without touching or moving the particle. The iridescence seen in Fig. 2(a) indicates that

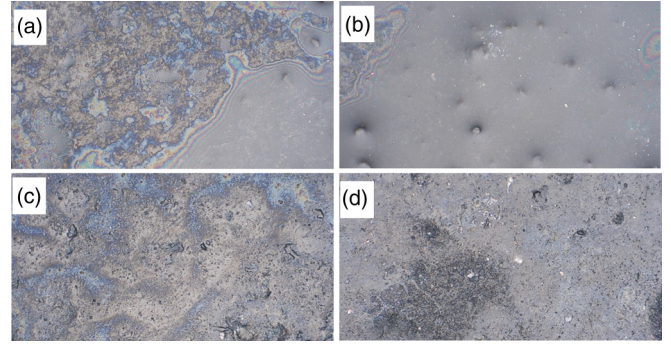


FIG. 2. Close-up visualization of the PBS coating on two different sample glass beads ($d = 10$ mm) with an optical microscope (magnification $\times 700$). (a) An irregularly coated area of particle 1 and (c) the same area after leaning with heptane. (b) A well coated area of particle 2 and (d) the same clean area. Iridescence can be seen where the PBS layer is not homogeneous.

the PBS layer is not perfectly uniform and may suffer from thickness variation, and that some “holes” in the coating layer may also exist [Fig. 2(b)]. However, a statistical analysis of the interparticle force discussed later in the paper (see Sec. III) shows that the presence of defects weakly affects the cohesion between two particles. Since the coating layer is very thin, no capillary structure has been observed in optical microscopy when putting two beads in contact.

A last important remark is that the strong Si-OH link between the polymer and the glass bead surface helps the PBS to stick permanently on the particles. No drainage of the fluid was observed even for a very long time (on the timescale of a year), making the material very stable in time, as will be discussed in Sec. IV.

III. INTERPARTICLE COHESION FORCE MEASUREMENTS

The cohesion force between two particles is a key parameter to control the macroscopic cohesion of a granular assembly. In this section we present the results of different experiments designed to measure the contact force between two particles due to the PBS coating. As presented in the previous section, the control parameters are the size of the beads (diameter d), the PBS averaged layer thickness b , and the duration of the contact. We designed two methods to accurately measure the cohesion force. The first method uses the torque-meter of a commercial rheometer (MCR501 Anton Paar). It provides accurate measurements but performing a statistical analysis is tedious too. The second method is a home-designed force measurement device allowing simultaneous measurements on 10 pairs of particles.

A. Role of the precompression load

The cohesion force between two CCGM particles has been first measured using the rotating head of a Anton-Parr MCR501 rheometer. A sketch of the experimental setup is given in Fig. 3(a). A coated particle is attached to a fixed rigid structure through a linear spring, and a similar coated particle from the same batch is glued at the end of an arm

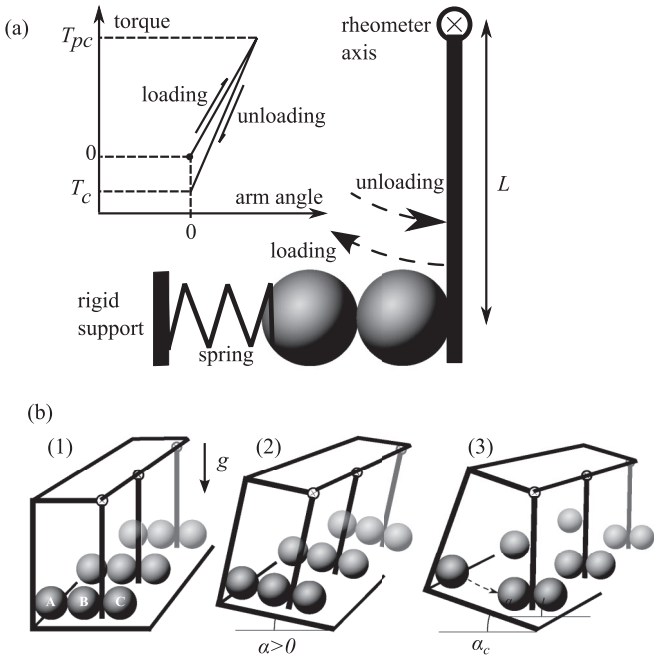


FIG. 3. (a) Sketch of the setup to measure the cohesion force for different precompression force using the rheometer torque meter. The two particles are put in contact with a precompression force $F_{pc} = T_{pc}/L$, where $L = 3.5$ cm is the arm length and the cohesion force $F_c = T_c/L$ is measured when the two particles detach. (b) Sketch of the pendulum experimental setup. Particle A is attached to a rigid structure, particles B and C are attached to the two sides of a pendulum. F_c is measured by inclining the setup. Ten pendulums were mounted in parallel.

attached to the rheometer head. The two particles were put in contact, and a precompression torque T_{pc} corresponding to a precompression force F_{pc} was applied before slowly reversing the applied torque up to the point when the two particles suddenly detach. The critical torque when detachment occurs provides the measurement of the cohesion force F_c . [Fig. 3(a)].

We measured the cohesion force using $d = 10$ mm particles and with a coating $b = 5 \mu\text{m}$, and the precompression force F_{pc} was varied from 0.08 N to 2 N. Figure 4(a) shows that the cohesion force F_c does not depend on the precompression force F_{pc} , and that the order of magnitude of the cohesion force is $F_c \approx 5$ mN (dashed line). This independence of the cohesion with the compression force has been also observed in a different system by Kobayashi *et al.* [28]. With this setup, we also studied if the cohesion force was affected by the number of successive contacts. One can wonder if the polymer layer can be altered after the first sticking contact. Figure 4(b) shows for three different precompression forces that the cohesion force is independent of the number of successive contacts. We therefore conclude that the PBS layer is strongly attached to the glass bead surface and that the stick-pull process occurring for a binary contact is reversible. A last important remark is that a variation in the mean cohesion force is observed in Fig. 4(b): the mean cohesion is $F_c = 4.3$ mN for the triangle symbols and $F_c = 3.7$ mN mean force for the star symbols. This is an indication that the cohesion force may vary from one pair of particles to another and that

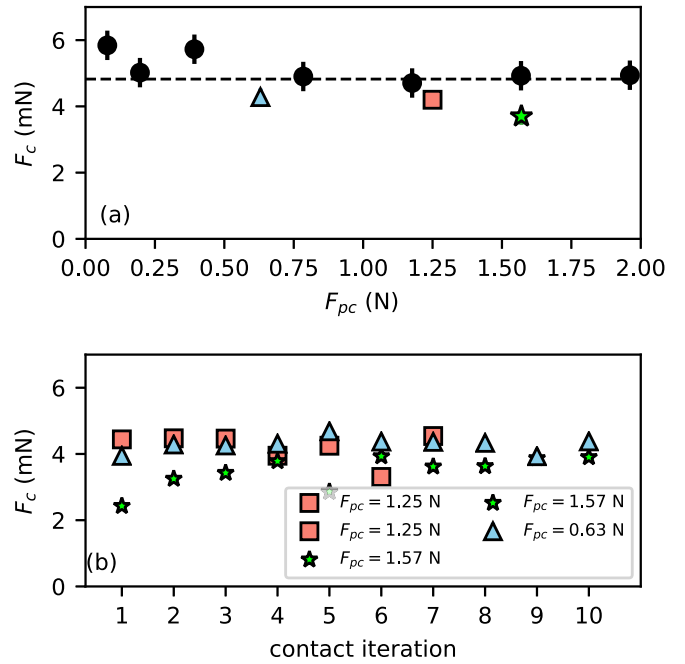


FIG. 4. (a) Cohesion force measured for different precompression forces $d = 10$ mm, $b = 5 \mu\text{m}$. The dashed line indicates the mean cohesion force. Empty colored symbols refer to the legend of (b). (b) Cohesion force for successive contacts, and for different precompression forces. The contact waiting time was kept constant equal to 10 min. If not visible, the error bars are smaller than the symbol size.

a statistical analysis is necessary. This has motivated us to develop a second experimental setup to measure in parallel the cohesion force for 10 pairs of particles.

B. Role of the contact waiting time

The second home-made force measurement device is sketched in Fig. 3(b) and consists of a set of 10 independent parallel pendulums. Each pendulum had one particle (B) attached at the bottom of the arm [Fig. 3(b)], which came into contact with a fixed particle (A). A third particle (C) was also glued on the other side of the pendulum arm and played the role of a counterweight. The setup was mounted on a table that can be inclined. Starting from a nearly horizontal position (step 1 in Fig. 3), the table was slowly inclined with a rate 10°min^{-1} (step 2) until all the pairs of particles detached (step 3). The whole measurement process was recorded with a camera, and each time a pair of particles was detaching, the angle α_c was recorded and the cohesion force F_c was computed from the torque balance.

With this device, the influence of the contact waiting time between two coated particles has been investigated from 5 s to 2 hr. We also investigated a 24 hr waiting time, but the results were not significantly different from the 2 hr result. Figure 5 shows that the cohesion force varies with contact time t_c for $t_c \leq 600$ s but eventually saturates for long contact times $t_c > 1000$ s. This confirms the qualitative observations made when handling the CCGM out of storage. A CCGM stored during a long time looks more cohesive, although a vigorous shaking of

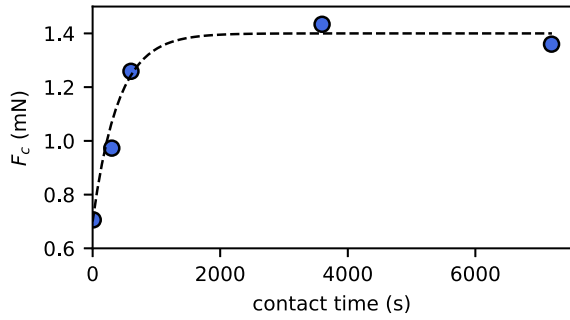


FIG. 5. Cohesion force as a function of the duration of the contact. The dashed line is a qualitative trend illustrating an exponential relaxation with time.

the packing which renews all the contacts seems to diminish the cohesive nature of the sample. In the following, we now refer to “short” waiting time experiments when $t_c = 10$ s, and “long” waiting time experiments when $t_c = 10$ min.

The cohesion force distribution has been measured for 100 pairs of particles out of the same batch ($d = 5$ mm, $b = 2$ μ m). The probability distribution function $p(F_c)$ is shown in Fig. 6 for short and long contact time. The averaged cohesion force is 0.56 ± 0.1 mN and 1.14 ± 0.3 mN for short and long contact time, respectively. For $t_c = 10$ s the cohesion force distribution is narrower than for $t_c = 10$ min. We have not investigated in more detail the influence of the contact time and the origin of the force distribution, which are certainly related to the coating property of the particle and to the entanglement dynamics of the polymer chains.

C. Scaling of the cohesion force

To understand the physical origin of the cohesion force, we have systematically studied how it varies with the particle diameter d and the averaged PBS coating thickness b . In Fig. 7 the cohesion force is plotted as a function of the particle

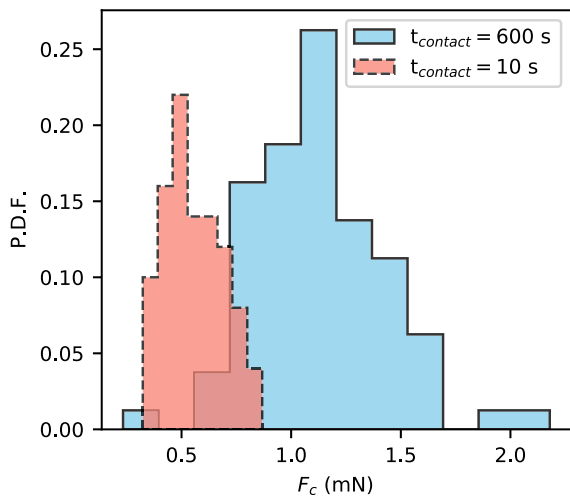


FIG. 6. Probability distribution function of the cohesion force measured for approximately 100 pairs of particles, for two different contact times ($t_c = 10$ s and $t_c = 10$ min) and for $d = 5$ mm, $b = 2$ μ m coated particles.

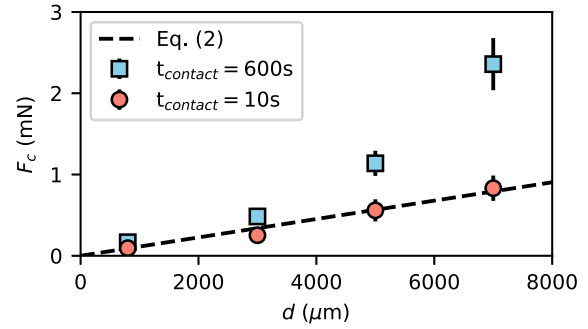


FIG. 7. The cohesion force F_c as a function of the particle diameter d for short (10 s) (circles) and long (10 min) (squares) contact times. The dashed line is the linear expression (2).

diameter d for a constant layer thickness $b = 2$ μ m, and for the short and long waiting contact times. In the range $0.8 < d < 7$ mm the cohesion force increases with the particle diameter. The cohesion force varies linearly with the diameter for short contact time but exhibits a more rapid increase for long contact time. The linear variation can be well described by a capillary model at contact [29]

$$F_c = \frac{3}{2}\pi\gamma d, \tag{2}$$

where the surface tension $\gamma \approx 24$ mN m^{-1} , a relevant order of magnitude for PDMS. For long contact times, other molecular phenomena may occur, such as a slow polymer entanglement between PDMS polymers, and we did not investigate further the long-time correlation between the cohesion force and the particle radius.

The influence of the coating PBS thickness b on F_c is studied in Fig. 8 for different particle diameters d . The cohesion force normalized by the expression (2) is plotted as a function of b . We first observe that all the data obtained for different particle diameters collapse on a single curve. The normalized cohesion force starts from zero when there is no coating, increases and reaches a plateau equal to 1 when the averaged

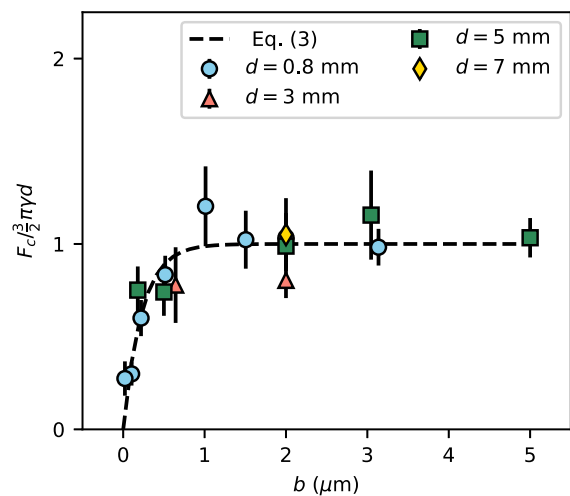


FIG. 8. Cohesion force normalized by $\frac{3}{2}\pi\gamma d$ as a function of the mean PBS layer b for short contact times and for different particle sizes.

thickness of PBS is larger than $1 \mu\text{m}$. An *ad hoc* expression for the cohesion force can be proposed:

$$F_c = \frac{3}{2}\pi\gamma d(1 - e^{-b/B}), \quad (3)$$

where $B \approx 230 \text{ nm}$ is a characteristic thickness which is a few times the particle roughness value ($30 \pm 2 \text{ nm}$). This behavior is reminiscent of what is observed with liquid capillary bridges. In this latter case the cohesion force increases when increasing the amount of liquid, up to the point where the liquid screens the surface roughness and that a single bridge exists, giving rise to the saturated force given by Eq. (2) [15].

From this expression, we can write the expression for a Bond number, i.e., the ratio of the weight of the particle over the cohesion force, a dimensionless number that will be useful in the following:

$$\text{Bo} = \frac{1}{9} \frac{\rho_G g d^2}{\gamma(1 - e^{-b/B})}. \quad (4)$$

The threshold value $\text{Bo} = 1$ gives a critical particle size for which the weight is balanced by the cohesion force. A typical example of $\text{Bo} \approx 1$ is given by a coating thickness $b = 2 \mu\text{m}$ with particles $d = 3 \text{ mm}$, as illustrated in Fig. 1.

IV. BULK BEHAVIOR OF THE CCGM

The previous sections were dedicated to the measure of the cohesion force between pairs of particles. In this section we discuss the collective behavior of the coated particles in several classical configurations used in the literature for characterizing granular media. The goal is to show that the ability to precisely control the cohesion force between the grains open new perspectives to understand the influence of the interparticle force in the dynamics of cohesive granular media. This section presents experimental results for the bulk density, for the angle of repose of static piles, and for the onset of the flow of a layer of CCGM resting on an inclined plane.

A. Packing fraction

In many industrial processes, the bulk density of a granular assembly is a qualitative indicator of the cohesive property of the medium [30,31]. The Haussner ratio ($H = \rho_T/\rho_B$) or the Carr index [defined as $100(1 - 1/H)$] is often used, both implying the measurement of the bulk density in two different compaction states: the aerated density ρ_B (similar to the random loose packing state) and the tapped density ρ_T (similar to the random close packing). In the following we investigated how the loose packing fraction changes for CCGM batches when varying the PBS coating. The random loose packing fraction ϕ_{rlp} was evaluated through mass and volume measurements in a 250 cm^3 graduated cylindrical test tube. The results are presented in Fig. 9 for $d = 680 \text{ mm}$ particles and for a coating layer thickness varying from 72 to 624 nm. The best collapse of experimental data is obtained with a plot of the packing fraction versus the volume ratio Ω_P/Ω_G , which is proportional to b/d .

For very low values of the coating ($\Omega_P/\Omega_G \approx 10^{-4}$), the packing fraction is equal to the packing fraction of clean

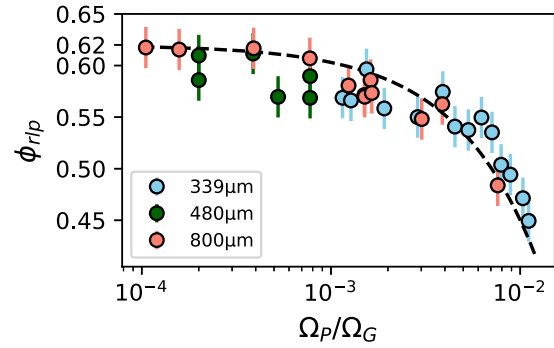


FIG. 9. Random loose packing fraction of various CCGM with different particle sizes and different coating values. The packing fraction decreases with the volume ratio Ω_P/Ω_G . The legend indicates the beads' diameter.

and dry glass beads. The packing fraction decreases for an increasing PBS content, and a low packing fraction $\phi_{rlp} \approx 0.45$ may be reached for a typical $\Omega_P/\Omega_G \approx 10^{-2}$ value. This can be explained by the existence of large-scale voids and arches in the bulk sustained by strong cohesive links between particles.

B. Angle of repose

The measure of the slope angle of a granular heap is also a way to emphasize the role of the cohesion in a powder or in a granular material [14,32–36]. With our CCGM, static piles were made from a chute flow from a hopper on a 5-cm-diameter rough disk. A side-view camera captured the image of a pile and the angle of repose α_r is obtained from image analysis. Examples of images of piles are given in Fig. 10. The value of the repose angle is averaged over 20 measurements. The repose angle is observed to increase when increasing the PBS thickness. Without coating [Fig. 10(a)], the heap presents a smooth surface with a constant angle. Adding some cohesion gives rise to steeper slopes and also to abrupt local variations of the local angle as seen illustrated in Figs. 10(b)–10(d).

Figure 11 shows that the angle θ_r increases with the coating thickness b , with a sharp increase for $b \approx 40 \text{ nm}$, and seems to saturate for coatings larger than 50 nm. We compare our

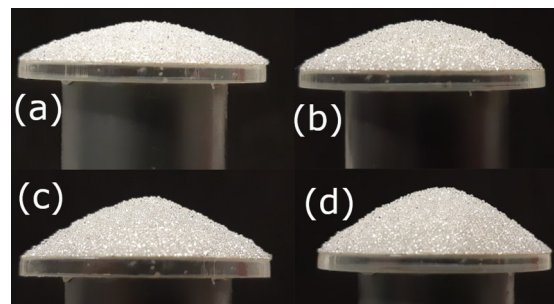


FIG. 10. Images of piles for a CCGM with $d = 480 \mu\text{m}$ with increasing PBS coating: (a) no coating, $\alpha_r = 27.7 \pm 0.8$, (b) $b = 31 \text{ nm}$, $\alpha_r = 30.1 \pm 0.9$, (c) $b = 52 \text{ nm}$, $\alpha_r = 40.3 \pm 1.9$, (d) $b = 62 \text{ nm}$, $\alpha_r = 42.5 \pm 2.2$

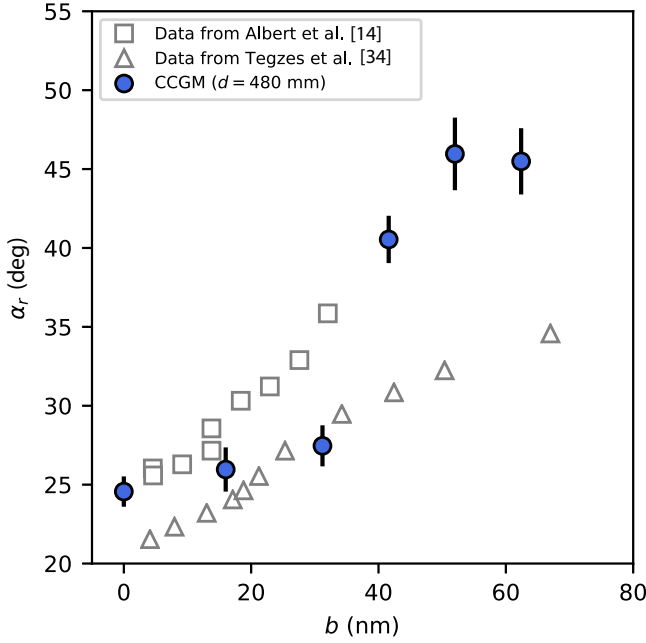


FIG. 11. Heap repose angle for $d = 480 \mu\text{m}$ particles and various coatings. Empty symbols are data from the literature with capillary cohesion: $d = 800 \mu\text{m}$ [14] (squares) and $d = 900 \mu\text{m}$ [34] (triangles).

results with the repose angle obtained with the crater method using vacuum pump oil-coated particles [14,34] in Fig. 11. While the heap formation methods are different, the CCGM presents a similar trend. Our experimental setup is limited to low-cohesion values because the materials prepared with a large cohesion ($b > 100 \text{ nm}$) experience difficulties to flow through the hopper under gravity only.

The heap angle experiment is also a benchmark test to assess the stability and the durability of the CCGM. The first test concerns the stability with temperature since the cohesion is based on a cross-linked polymer. Piles have been prepared with a CCGM stored in controlled-temperature devices. As shown in Fig. 12(a), the repose angle is nearly independent of temperature from 0°C to 60°C . No noticeable difference was found between experiments at 20°C and 60°C . This means that no specific care is needed for experiments at a standard room temperature.

We have also investigated the stability of the CCGM with time by measuring the heap repose angle for the same batch of particles at different ages. Figure 12(b) shows the evolution of the repose angle α_r for three different materials at different ages from preparation. This plot shows that for thin coatings ($b = 16$ or $b = 31 \text{ nm}$), the heap angle remains identical even for a sample prepared 1 yr ago. For a thicker coating ($b = 62 \text{ nm}$), a slow decrease of the repose angle has been observed. Nevertheless, the PBS-coated CCGM seems to be stable for months, and large batches can then be prepared before performing large-scale experiments. This CCGM also seems to be unaffected by the moisture content of the ambient air since our experiments were performed without humidity control across different temperature and humidity conditions over a year.

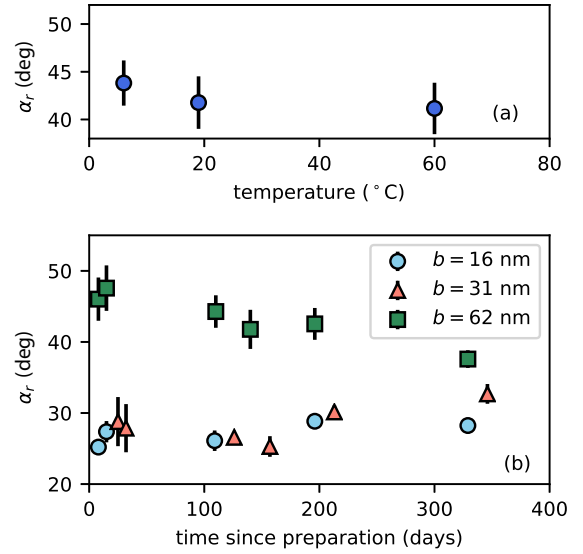


FIG. 12. (a) Effect of the temperature on the repose angle of a CCGM ($d = 480 \mu\text{m}$, $b = 62 \text{ nm}$). (b) Heap repose angle α_r as a function of the time since the preparation of the CCGM. Experiments were made with $d = 480 \mu\text{m}$ particles.

C. Onset of flow on an inclined plane

Measuring the onset of flow of a layer of particles lying on a rough inclined bed is another way to investigate the friction and the cohesion of a material. The simplest description of the plasticity of a granular material assumes that the yield stress follows a cohesive Mohr-Coulomb criterion, $\tau_{\text{yield}} = \mu P + \tau_c$, where μ is the friction coefficient, P the confining pressure and τ_c the cohesive stress. Starting from an horizontal plane and increasing progressively the inclination θ , a layer of thickness h will start to flow at a critical angle θ_{start} when the shear stress at the base reaches the yield stress value:

$$\rho_G \phi g h \sin \theta_{\text{start}} = \mu \rho_G \phi g h \cos \theta_{\text{start}} + \tau_c \quad (5)$$

where ϕ is the volume fraction of the layer. This equation can be simplified as

$$h \sin \theta_{\text{start}} = \mu h \cos \theta_{\text{start}} + h_c \quad (6)$$

where $h_c = \tau_c / (\phi \rho_G g)$ is a characteristic cohesive length, which represents the maximum thickness of a self-standing vertical layer of granular medium stuck to a rough surface under gravity.

Equation (6) shows that the cohesion length h_c and the friction coefficient μ can be identified by systematically measuring the critical starting angle θ_{start} for different thicknesses h . We have conducted such a series of experiments with our model cohesive material. For seek of efficiency, we have not used a uniform layer as initial state, but rather a prismatic deposit, as sketched in Fig. 13(a). With this geometry, it is possible in a single experiment to perform several measurements of $(h, \theta_{\text{start}})$.

A typical experiment is conducted as follows. The CCGM is poured on a $20 \times 6 \text{ cm}^2$ rectangular rough plate (the roughness is made with CCGM particles glued on a double sided adhesive tape) with two prismatic side walls. The free surface of the deposit is then leveled following the two side walls.

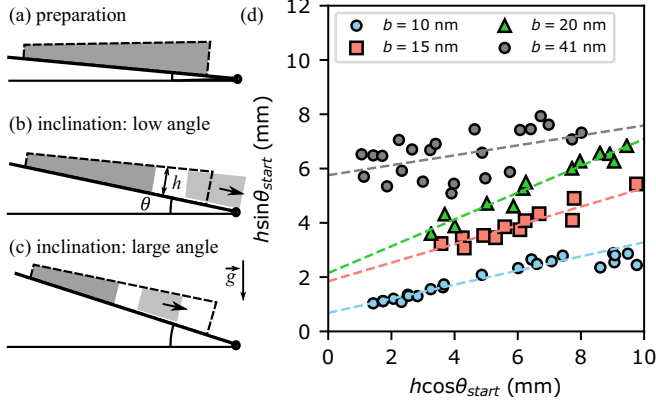


FIG. 13. (a)–(c) Sketch of the inclined plane setup with the variable granular thickness and a progressive inclination. (d) Inclined plane results for $d = 202 \mu\text{m}$ CCGM particles with increasing coating thickness. Dashed lines are best fits using Eq. (6).

The final prismatic volume has a thickness varying linearly from 15 to 25 mm [see Fig. 13(a)]. The thickness of the granular layer is measured with a laser sheet technique and the angle with a clinometer. Starting from a very low angle of inclination (typically 10°), the setup is slowly inclined at a constant rate. A first avalanche occurs at the bottom thick side [Fig. 13(b)], leaving a thinner and shorter prism. When the angle is further increased, a second avalanche occurs [Fig. 13(c)], which corresponds to a different h , and so on. In one experiment, one can then extract the critical angle θ_{start} for four to five different thicknesses. For a single CCGM batch, this experiment is repeated several times. The collected data are then plotted in a $(h \cos \theta_{start}, h \sin \theta_{start})$ plane. According to Eq. (6), a linear fit of experimental data gives the slope μ and the intercept value h_c for a given CCGM. Figure 13(d) shows the $(h \cos \theta_{start}, h \sin \theta_{start})$ plot for small glass beads of diameter $d = 202 \pm 4 \mu\text{m}$. Results are given for four coatings of increasing thickness b . Despite some experimental noise, the linear expression (5) fits well the experimental data. Increasing the coating thickness b increases the cohesive length h_c (the intercept of the linear fit with the vertical axis) but does not significantly affect the friction coefficient (the slope of the lines). From the measure of h_c , one can then estimate the cohesive stress τ_c .

The stress τ_c is a macroscopic measurement of the cohesion, which can be compared to the interparticle cohesion force measured in Sec. III. From a dimensional analysis, the scaling relating the cohesive shear stress and the cohesion force is $\tau_c d^2 \propto F_c$. Following Ref. [17], a theoretical expression based on a mean field approach relates the macroscopic cohesion τ_c to the cohesion force:

$$\tau_c = \frac{3\mu\phi Z F_c}{2\pi d^2}, \quad (7)$$

with μ the friction coefficient, ϕ the volume fraction and Z the averaged coordinance number (number of contacts per particle). Figure 14 gathers our data for different particle sizes and different PBS coatings and shows a linear trend between $\tau_c d^2$ and F_c although it is not perfect especially for small particles. The presence of other cohesive forces like

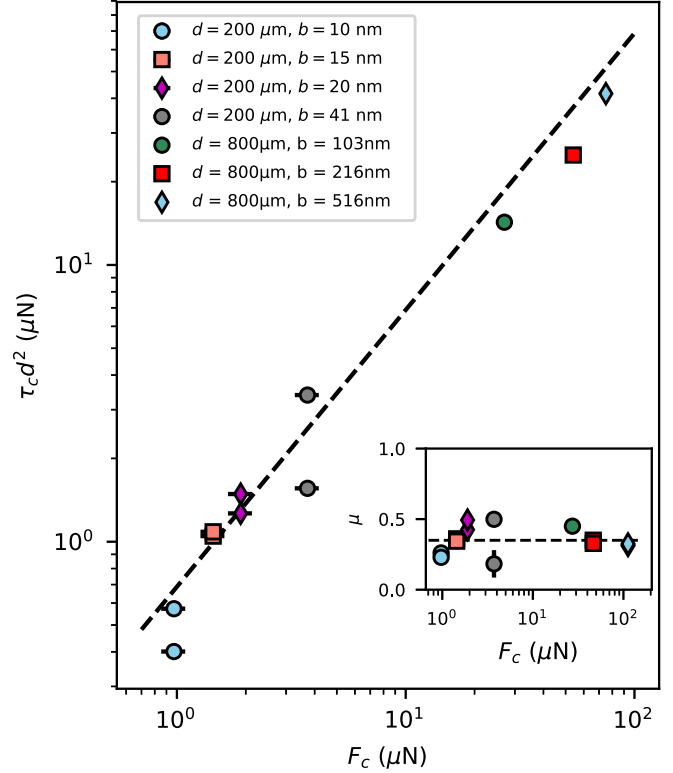


FIG. 14. The macroscopic cohesion force $\tau_c d^2$ measured from inclined plane experiments as a function of the interparticle cohesive force F_c . The dashed line is the prediction from Eq. (7).

humidity may explain this discrepancy observed for small particles. Taking the experimental averaged value $\mu = 0.4$ along with $\phi = 0.6$ and $Z = 6$, the theoretical prediction is plotted in Fig. 14 and gives indeed a good estimate of the measured cohesion. The insert of Fig. 14 also shows that the macroscopic friction coefficient μ seems independent of the coating property of the particles.

V. CONCLUDING REMARKS AND PERSPECTIVES

A cohesion-controlled granular material (CCGM) has been designed with glass beads coated by a thin layer of polyborosiloxane polymer. The preparation technique is simple, the resulting CCGM is very robust, and it can be used several times in various situations to model industrial or natural cohesive granular materials. We have investigated how the interparticle cohesive contact force varies with the thickness of the coating and with the particle size and have showed that the cohesion force does not depend on the precompression force. We also showed that the cohesion force slightly evolves with time and eventually saturates for a long time.

Different experimental setups were used to put in evidence the influence of the cohesion on various macroscopic quantities. We have illustrated the effect of the cohesion on the packing fraction of loose samples, on the repose angle of heaps, and on the onset of flow of a granular deposit on a rough bed. In all cases we have shown that controlling the interparticle force is a way to control the macroscopic properties. This CCGM seems to be very stable in time and weakly affected

by the humidity or the temperature of the laboratory. This is a major advantage compared to previous capillary cohesion techniques based on Newtonian fluids where drainage, evaporation, and capillary bridges recombination may occur during an experiment.

In this paper we have restricted our observation to static configurations. However, the possibility to control the cohesion open many perspectives to study also the flow properties of cohesive materials. Configurations such as flows down inclined planes, flows in hoppers and silos, flows in rotating drums, and mixing or segregation may also be investigated,

with the hope that it will help to better understand the physics beyond the concept of “flowability” of cohesive granular materials.

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