Jammed hard-particle packings: From Kepler to Bernal and beyond

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Understanding the characteristics of jammed particle packings provides basic insights into the structure and bulk properties of crystals, glasses, and granular media and into selected aspects of biological systems. This review describes the diversity of jammed configurations attainable by frictionless convex nonoverlapping (hard) particles in Euclidean spaces and for that purpose it stresses individual-packing geometric analysis. A fundamental feature of that diversity is the necessity to classify individual jammed configurations according to whether they are locally, collectively, or strictly jammed. Each of these categories contains a multitude of jammed configurations spanning a wide and (in the large system limit) continuous range of intensive properties, including packing fraction $\phi$, mean contact number $Z$, and several scalar order metrics. Application of these analytical tools to spheres in three dimensions (an analog to the venerable Ising model) covers a myriad of jammed states, including maximally dense packings (as Kepler conjectured), low-density strictly jammed tunneled crystals, and a substantial family of amorphous packings. With respect to the last of these, the current approach displaces the historically prominent but ambiguous idea of “random close packing” with the precise concept of “maximally random jamming.” Both laboratory procedures and numerical simulation protocols can and, frequently, have been used for creation of ensembles of jammed states. But while the resulting distributions of intensive properties may individually approach narrow distributions in the large system limit, the distinguishing varieties of possible operational details in these procedures and protocols lead to substantial variability among the resulting distributions, some examples of which are presented here. This review also covers recent advances in understanding jammed packings of polydisperse sphere mixtures, as well as convex nonspherical particles, e.g., ellipsoids, “superballs,” and polyhedra. Because of their relevance to error-correcting codes and information theory, sphere packings in high-dimensional Euclidean spaces have been included as well. Some remarks are also made about packings in (curved) non-Euclidean spaces. In closing this review, several basic open questions for future research to consider have been identified.

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The importance of packing hard particles into various kinds of vessels and the questions it raises have an ancient history. Bernal remarked that “heaps (close-packed arrangements of particles) were the first things that were ever measured in the form of basketfuls of grain for the purpose of trading or the collection of taxes” (Bernal, 1965). Although packing problems are easy to pose, they are notoriously difficult to solve rigorously. In 1611, Kepler was asked: What is the densest way to stack equal-sized cannon balls? His solution, known as “Kepler’s conjecture,” was the face-centered-cubic (fcc) arrangement (the way your greengrocer stacks oranges). Gauss (1831) proved that this is the densest Bravais lattice packing (defined below). But almost four centuries passed before Hales proved the general conjecture that there is no other arrangement of spheres in three-dimensional Euclidean space whose density can exceed that of the fcc packing (Hales, 2005); see Aste and Weaire (2008) for a popular account of the proof. Even the proof of the densest packing of congruent (identical) circles in the plane, the two-dimensional analog of Kepler’s problem, appeared only 70 years ago (Rogers, 1964; Conway and Sloane, 1998); see Fig. 1.

Packing problems are ubiquitous and arise in a variety of applications. These exist in the transportation, packaging, agricultural, and communication industries. Furthermore, they have been studied to help understand the symmetry, structure, and macroscopic physical properties of condensed-matter phases, liquids, glasses, and crystals (Mayer and Mayer, 1940; Bernal, 1960, 1965; Stillinger, DiMarzio, and Korney, 1964; Stillinger and Salsburg, 1969; Weeks et al., 1971; Ashcroft and Mermin, 1976; Hansen and McDonald, 1986; Woodcock and Angell, 1981; Speedy, 1994; Chaikin and Lubensky, 1995).

Packing problems are also relevant for the analysis of heterogeneous materials (Torquato, 2002), colloids (Russel et al., 1989; Chaikin and Lubensky, 1995; Torquato, 2009), and granular media (Edwards, 1994). Understanding the symmetries and other mathematical characteristics of the densest sphere packings in various spaces and dimensions is a challenging area of long-standing interest in discrete geometry and number theory (Rogers, 1964; Conway and Sloane, 1998; Cohn and Elkies, 2003) as well as coding theory (Shannon, 1948; Conway and Sloane, 1998; Cohn and Kumar, 2007b).

It is appropriate to mention that packing issues also arise in numerous biological contexts, spanning a wide spectrum of length scales. This includes “crowding” of macromolecules within living cells (Ellis, 2001), the packing of cells to form tissue (Torquato, 2002; Gevertz and Torquato, 2008), the spiral patterns seen in plant shoots and flowers (phyllotaxis) (Prusinkiewicz and Lindenmayer, 1990; Nisoli et al., 2010) and the competitive settlement of territories by animals, the patterns of which can be modeled as random sequential packings (Tanemura and Hasegawa, 1980; Torquato, 2002). Figure 2 shows macromolecular crowding and a familiar phyllotactic pattern.

We call a packing a large collection of nonoverlapping (i.e., hard) particles in either a finite-sized container or in d-dimensional Euclidean space R^d. The packing fraction φ is the fraction of space covered by (interior to) the hard particles. “Jammed” packings are those particle configurations in which each particle is in contact with its nearest neighbors in such a way that mechanical stability of a specific type is conferred to the packing (see Sec. IV). Jammed packings and their properties have received considerable attention in the literature, both
experimental and theoretical. Within the domains of analytical theory and computer simulations, two conceptual approaches for their study have emerged. One is the “ensemble” approach (Bernal, 1960, 1965; Edwards, 1994; Liu and Nagel, 1998; Edwards and Grivev, 2001; Makse and Kurchan, 2002; Silbert et al., 2002, 2005; O’Hern et al., 2003; Wyart et al., 2005; Gao et al., 2006; Song et al., 2008; Parisi and Zamponi, 2010), which for a given packing procedure aims to understand typical configurations and their frequency of occurrence. The other, more recently, is the “geometric-structure” approach (Torquato et al., 2000, 2003; Torquato and Stillinger, 2001, 2007; Kansal et al., 2002b; Donev, Cisse, et al., 2004; Donev et al., 2004a; Donev, Connelly, et al., 2007), which emphasizes quantitative characterization of single-packing configurations, without regard to their occurrence frequency in the algorithmic method used to produce them. Our primary objective is to review the latter approach, while at the same time to show that these two approaches are complementary in that they represent different aspects of the larger context of hard-particle jamming phenomena. In particular, a wide range of jammed packing ensembles can be created by the choice of the generating algorithm, and the geometric-structure approach analyzes and classifies individual members of those ensembles, whether they be crystalline or amorphous at any achievable packing fraction $\phi$.

The process of cooling an initially hot liquid ultimately to absolute zero temperature provides a close and useful analogy for the subject of hard-particle jamming. Figure 3 summarizes this analogy in schematic form, showing typical paths for different isobaric cooling rates in the temperature-volume plane. While these paths are essentially reproducible for a given cooling schedule, i.e., giving a narrow distribution of results, that distribution depends sensitively on the specific cooling schedule or protocol that has been used. A very rapid quench that starts with a hot liquid well above its freezing temperature will avoid crystal nucleation, producing finally a glassy solid at absolute zero temperature. A somewhat slower quench from the same initial condition can also avoid nucleation but will yield at its $T=0$ end point a glassy solid with lower volume and potential energy. An infinitesimal cooling rate in principle will follow a thermodynamically reversible path of equilibrium states, will permit nucleation, and will display a volume discontinuity due to the first-order freezing transition on its way to attaining the structurally perfect crystal ground state. By analogy, for hard-particle systems compression qualitatively plays the same role as decreasing the temperature in an atomic or molecular system. Thus it is well known that compressing a monodisperse hard-sphere fluid very slowly leads to a first-order freezing transition, and the resulting crystal phase corresponds to the closest packing arrangement of those spheres (Mau and Huse, 1999). The resulting hard-sphere stacking variants are configurational images of mechanically stable structures exhibited, for example, by the venerable Lennard-Jones model system. By contrast, rapid compression rates applied to a hard-sphere fluid will create random amorphous jammed packings (Rintoul and Torquato, 1996b), the densities of which can be controlled by the compression rate utilized (see Fig. 11). In both the cases of cooling liquid glass formers and of compressing monodisperse hard spheres, it is valuable to be able to analyze the individual many-particle configurations that emerge from the respective protocols.

We begin this review, after introducing relevant terminology, by specifically considering packings of frictionless identical spheres in the absence of gravity, which represents an idealization of the laboratory situation for investigations of jammed packings; see Sec. III. This simplification follows that tradition in condensed-matter science to exploit idealized models, such as the Ising model, which is regarded as one of the pillars of statistical mechanics (Onsager, 1944; Domb, 1960; Gallavotti, 1999). In that tradition, this idealization offers the opportunity to obtain fundamental as well as practical insights and to uncover unifying concepts that describe a broad range of phenomena. The stripped-down hard-sphere “Ising model” for jammed packings (i.e., jammed, frictionless, and identical spheres in the absence of gravity) embodies the primary attributes of real packings while simultaneously generating mathematical challenges. The geometric-structure approach to analyzing individual jammed states produced by this model...
 covers not only the maximally dense packings (e.g., Kepler’s conjecture) and amorphous “Bernal” packings but an unbounded collection of other jammed configurations. This approach naturally leads to the inevitable conclusion that there is great diversity in the types of attainable jammed packings with varying degrees of order, mechanical stability, and density.

Important insights arise when jammed sphere packings are placed in a broader context that includes jammed states of noncongruent spheres as well as nonspherical objects, as discussed in Secs. IX and X. These extensions include polydisperse spheres, ellipsoids, superballs, and polyhedra in three dimensions. In addition, this broader context also involves sphere packings in Euclidean space with high dimensions (Sec. XI), which is relevant to error correcting codes and information theory (Shannon, 1948; Conway and Sloane, 1998), and packings in non-Euclidean spaces (Sec. XII). Finally, in Sec. XIII, we identify a number of basic open questions for future research.

II. PRELIMINARIES AND DEFINITIONS

Some basic definitions concerning packings are given here. A packing $P$ is a collection of nonoverlapping solid objects or particles in $d$-dimensional Euclidean space $\mathbb{R}^d$. Packings can be defined in other spaces (e.g., hyperbolic spaces and compact spaces, such as the surface of a $d$-dimensional sphere), but our primary focus in this review is $\mathbb{R}^d$. A saturated packing is one in which there is no space available to add another particle of the same kind to the packing.

We will see subsequently that whether a particle possesses central symmetry plays a fundamental role in determining its dense packing characteristics. A $d$-dimensional particle is centrally symmetric if it has a center $C$ that bisects every chord through $C$ connecting any two boundary points of the particle, i.e., the center is a point of inversion symmetry. Examples of centrally symmetric particles in $\mathbb{R}^d$ are spheres, ellipsoids, and superballs (defined in Sec. X). A triangle and tetrahedron are examples of noncentrally symmetric two- and three-dimensional particles, respectively. Figure 4 shows examples of centrally and noncentrally symmetric two-dimensional particles. A $d$-dimensional centrally symmetric particle for $d \geq 2$ is said to possess $d$ equivalent principal (orthogonal) axes (directions) associated with the moment of inertia tensor if those directions are twofold rotational symmetry axes such that the $d$ chords along those directions and connecting the respective pair of particle-boundary points are equal. [For $d=2$, the twofold (out-of-plane) rotation along an orthogonal axis brings the shape to itself, implying the rotation axis is a “mirror image” axis.] Whereas a $d$-dimensional superball has $d$ equivalent directions, a $d$-dimensional ellipsoid generally does not (see Fig. 4).

A lattice $\Lambda$ in $\mathbb{R}^d$ is a subgroup consisting of the integer linear combinations of vectors that constitute a basis for $\mathbb{R}^d$. In the physical sciences and engineering, this is referred to as a Bravais lattice. Unless otherwise stated, the term “lattice” will refer here to a Bravais lattice only. A lattice packing $P_L$ is one in which the centroids of the nonoverlapping identical particles are located at the points of $\Lambda$, and all particles have a common orientation. The set of lattice packings is a subset of all possible packings in $\mathbb{R}^d$. In a lattice packing, the space $\mathbb{R}^d$ can be geometrically divided into identical regions $F$ called fundamental cells, each of which contains the centroid of just one particle. Thus, the density of a lattice packing is given by

$$\phi = v_1 / \text{Vol}(F),$$

where $v_1$ is the volume of a single $d$-dimensional particle and $\text{Vol}(F)$ is the $d$-dimensional volume of the fundamental cell. For example, the volume $v_1(R)$ of a $d$-dimensional spherical particle of radius $R$ is given explicitly by

$$v_1(R) = \pi^{d/2} R^d / \Gamma(1 + d/2),$$

where $\Gamma(x)$ is the Euler gamma function. Figure 5 shows...
lattice packings of congruent spheres and congruent nonspherical particles.

A more general notion than a lattice packing is a periodic packing. A periodic packing of congruent particles is obtained by placing a fixed configuration of \( N \) particles (where \( N \geq 1 \)) with "arbitrary nonoverlapping orientations" in one fundamental cell of a lattice \( \Lambda \), which is then periodically replicated without overlaps. Thus, the packing is still periodic under translations by \( \Lambda \), but the \( N \) particles can occur anywhere in the chosen fundamental cell subject to the overall nonoverlap condition. The packing density of a periodic packing is given by

\[
\phi = Nv_1/V = \rho v_1,
\]

where \( \rho = N/V \) is the number density, i.e., the number of particles per unit volume. Figure 6 shows a periodic nonlattice packing of congruent spheres and congruent nonspherical particles. Note that the particle orientations within a fundamental cell in the latter case are generally not identical to one another.

Consider any discrete (possibly infinite) set of points with position vectors \( X = \{r_1, r_2, \ldots\} \) in \( \mathbb{R}^d \). Associated with each point \( r_i \in X \) is its Voronoi cell \( \text{Vor}(r_i) \), which is defined to be the region of space no farther from the point at \( r_i \) than to any other point \( r_j \) in the set, i.e.,

\[
\text{Vor}(r_i) = \{r: |r - r_i| \leq |r - r_j| \text{ for all } r_j \in X\}. \tag{4}
\]

The Voronoi cells are convex polyhedra whose interiors are disjoint but share common faces, and therefore the union of all of the polyhedra is the whole of \( \mathbb{R}^d \). This partition of space is called the Voronoi tessellation. While the Voronoi polyhedra of a lattice are congruent (identical) to one another, the Voronoi polyhedra of a non-Bravais lattice are not identical to one another. Attached to each vertex of a Voronoi polyhedron is a Delaunay cell, which can be defined as the convex hull of the Voronoi-cell centroids nearest to it, and these Delaunay cells also tile space. Often the Delaunay tessellation is a triangulation of space, i.e., it is a partitioning of \( \mathbb{R}^d \) into \( d \)-dimensional simplices (Torquato, 2002). Geometrically the Voronoi and Delaunay tessellations are dual to each other. The contact network is only defined for a packing in which a subset of the particles form interparticle contacts. For example, when the set of points \( X \) defines the centers of spheres in a sphere packing, the network of interparticle contacts forms the contact network of the packing by associating with every sphere a "node" for each contact point and edges that connect all of the nodes. As we will see in Sec. IV, the contact network is crucial to determining the rigidity properties of the packing and corresponds to a subclass of the class of fascinating objects called tensegrity frameworks, namely, strut frameworks; see Connelly and Whiteley (1996) for details. Figure 7 shows the Voronoi, Delaunay, and contact networks for a portion of a packing of congruent circular disks.

Some of the infinite packings that we consider in this review can only be characterized spatially via statistical correlation functions. For simplicity, consider a nonoverlapping configuration of \( N \) identical \( d \)-dimensional spheres centered at the positions \( r^N = \{r_1, r_2, \ldots, r_N\} \) in a region of volume \( V \) in \( d \)-dimensional Euclidean space \( \mathbb{R}^d \). Ultimately, we pass to the thermodynamic limit, i.e., \( N \to \infty, V \to \infty \) such that the number density \( \rho = N/V \) is a fixed positive constant. For statistically homogeneous sphere packings in \( \mathbb{R}^d \), the quantity \( \rho^n g_n(r^n) \) is proportional to the probability density for simultaneously finding \( n \) sphere centers at locations \( r^n = \{r_1, r_2, \ldots, r_n\} \) in \( \mathbb{R}^d \) (Hansen and McDonald, 1986). With this convention, each \( n \)-particle correlation function \( g_n \) approaches unity when all particle positions become widely separated from one another. Statistical homogeneity implies that \( g_n \) is translationally invariant and therefore only depends on the relative displacements of the positions with respect to some arbitrarily chosen origin of the system, i.e.,

\[
g_n = g_n(r_{12}, r_{13}, \ldots, r_{1n}), \tag{5}
\]

where \( r_0 = r_j - r_i \).

The pair correlation function \( g_2(r) \) is the one of primary interest in this review. If the system is also rotationally invariant (statistically isotropic), then \( g_2 \) depends on the radial distance \( r = |r| \) only, i.e., \( g_2(r) = g_2(r) \).
It is important to introduce the total correlation function \( h(r) = g_2(r) - 1 \), which, for a disordered packing, decays to zero for large \( |r| \) sufficiently rapidly (Torquato and Stillinger, 2006b). We define the structure factor \( S(k) \) for a statistically homogeneous packing via
\[
S(k) = 1 + \rho \tilde{h}(k),
\]
(6)
where \( \tilde{h}(k) \) is the Fourier transform of the total correlation function \( h(r) = g_2(r) - 1 \) and \( k \) is the wave vector. Since the structure factor is the Fourier transform of an autocovariance function (involving the “microscopic” density) (Hansen and McDonald, 1986; Torquato and Stillinger, 2006b), then it follows it is a non-negative quantity for all \( k \), i.e.,
\[
S(k) \geq 0 \quad \text{for all } k.
\]
The non-negativity condition follows physically from the fact that \( S(k) \) is proportional to the intensity of the scattering of incident radiation on a many-particle system (Hansen and McDonald, 1986). The structure factor \( S(k) \) provides a measure of the density fluctuations in the packing at a particular wave vector \( k \).

### III. Lessons from Disordered Jammed Packings of Spheres

The classical statistical mechanics of hard-sphere systems has generated a large collection of scientific publications, dating back at least to Boltzmann, 1898. That collection includes examinations of equilibrium, transport, and jammed packing phenomena. With respect to the last of these, the concept of a unique random close packing (RCP) state, pioneered by Bernal (1960, 1965) to model the structure of liquids, has been one of the more persistent themes with a venerable history (Scott and Kilgour, 1969; Anonymous, 1972; Visscher and Bolsterli, 1972; Gotoh and Finney, 1974; Berryman, 1983; Jodrey and Tory, 1985; Tobochnik and Chapin, 1988; Zinchenko, 1994; Jullien et al., 1997; Pouliquen et al., 1997; Kamien and Liu, 2007). Until about a decade ago, the prevailing notion of the RCP state was that it is the maximum density that a large random collection of congruent (identical) spheres can attain and that this density is a well-defined quantity. This traditional view has been summarized as follows: “Ball bearings and similar objects have been shaken, settled in oil, stuck with paint, kneaded inside rubber balloons and all with no better result than (a packing fraction of) ... 0.636” (Anonymous, 1972). Torquato et al. (2000) argued that this RCP-state concept is actually ill-defined and thus should be abandoned in favor of a more precise alternative.

It is instructive to review these developments because they will point to the need for a geometric-structure approach generally to understand jammed packings, whether disordered or not. It has been observed (Torquato et al., 2000) that there has existed ample evidence, in the form of actual and computer-simulation experiments, to suggest strongly that the RCP state is indeed ill-defined and, in particular, dependent on the protocol used to produce the packings and on other system characteristics. In a classic experiment, Scott and Kilgour (1969) obtained the “RCP” packing fraction value \( \phi = 0.637 \) by pouring ball bearings into a large container, vertically vibrating the system for sufficiently long times to achieve a putative maximum densification, and extrapolating the measured volume fractions to eliminate finite-size effects. Important dynamical parameters for this kind of experiment include the pouring rate as well as the amplitude, frequency, and direction of the vibrations. The shape, smoothness, and rigidity of the container boundary are other crucial characteristics. For example, containers with curved or flat boundaries could frustrate or induce crystallization, respectively, in the packings, and hence the choice of container shape can limit the portion of configuration space that can be sampled. The key interactions are interparticle forces, including (ideally) repulsive hard-sphere interactions, friction between the particles (which inhibits densification), and gravity. The final packing fraction will inevitably be sensitive to these system characteristics. Indeed, one can achieve denser (partially and imperfectly crystalline) packings when the particles are poured at low rates into horizontally shaken containers with flat boundaries (Pouliquen et al., 1997).

It is tempting to compare experimentally observed statistics of so-called RCP configurations (packing fraction, correlation functions, and Voronoi statistics) to those generated on a computer. One must be careful in making such comparisons since it is difficult to simulate the features of real systems, such as the method of preparation and system characteristics (shaking, friction, gravity, etc.). Nonetheless, computer algorithms are valuable because they can be used to generate and study idealized random packings, but the final states are clearly protocol dependent. For example, a popular rate-dependent densification algorithm (Jodrey and Tory, 1985; Jullien et al., 1997) achieves \( \phi \) between 0.642 and 0.649, a Monte Carlo scheme (Tobochnik and Chapin, 1988) gives \( \phi \approx 0.68 \), a differential-equation densification scheme produces \( \phi = 0.64 \) (Zinchenko, 1994), and a “drop and roll” procedure (Visscher and Bolsterli, 1972) yields \( \phi \approx 0.60 \), and each of these protocols yields different sphere contact statistics.

As noted it has been argued that these variabilities of RCP arise because it is an ambiguous concept, explaining why there is no rigorous prediction of the RCP density, in spite of attempts to estimate it (Gotoh and Finney, 1974; Berryman, 1983; Song et al., 2008). The phrase “close packed” implies that the spheres are in contact with one another with the highest possible average contact number \( Z \). This would be consistent with the aforementioned traditional view that RCP presents the highest density that a random packing of close-packed spheres can possess. However, the terms “random” and “close packed” are at odds with one another. Increasing the degree of coordination (nearest-neighbor contacts) and thus the bulk system density comes at the expense of disorder. The precise proportion of each of these competing effects is arbitrary and therein lies a funda-
FIG. 8. (Color online) Typical protocols used to generate disordered sphere packings in three dimensions produce highly crystalline packings in two dimensions. Left panel: A three-dimensional MRJ-like configuration of 500 spheres with \( \phi \approx 0.64 \) produced using the Lubachevsky-Stillinger (LS) algorithm with a fast expansion rate (Torquato et al., 2000). Right panel: A crystalline collectively jammed configuration (Sec. IV.A) of 1000 disks with \( \phi = 0.88 \) produced using the LS algorithm with a fast expansion rate (Donev et al., 2004a).

mental problem. Moreover, since “randomness” of selected jammed packings has never been quantified, the proportion of these competing effects could not be specified. To remedy these serious flaws, Torquato et al. (2000) replaced the notion of close packing with jamming categories (defined in Sec. IV), which requires that each particle of a particular packing has a minimal number of properly arranged contacting particles. Furthermore, they introduced the notion of an “order metric” to quantify the degree of order (or disorder) of a single packing configuration.

Using the Lubachevsky-Stillinger (LS) (Lubachevsky and Stillinger, 1990) molecular-dynamics growth algorithm to generate jammed packings, it was shown (Torquato et al., 2000) that fastest particle growth rates generated the most disordered sphere (MRJ) packings (with \( \phi = 0.64 \); see the left panel of Fig. 8), but that by slowing the growth rates larger packing fractions could be continuously achieved up to the densest value \( \pi / \sqrt{18} \approx 0.74048 \ldots \) such that the degree of order increased monotonically with \( \phi \). Those results demonstrated that the notion of RCP as the highest possible density that a random sphere packing can attain is ill-defined since one can achieve packings with arbitrarily small increases in density at the expense of correspondingly small increases in order. This led Torquato et al. (2000) to supplant the concept of RCP with the maximally random jammed (MRJ) state, which is defined to be that jammed state with a minimal value of an order metric (see Sec. V). This work pointed the way toward a quantitative means of characterizing all packings, namely, the geometric-structure approach.

We note that the same LS packing protocol that leads to a uniformly disordered jammed state in three dimensions typically yields a highly crystalline “collectively” jammed packing in two dimensions. Figure 8 shows the vivid visual difference between the textures produced in three and two dimensions (see Sec. VII for further remarks). The low-concentration occurrence of crystal defects in the latter is evidence for the notion that there are far fewer collectively jammed states for \( N \) hard disks in two dimensions compared to \( N \) hard spheres in three dimensions. This distinction can be placed in a wider context by recalling that there is only one type of jammed state for hard rods in one dimension, and it is a defect-free perfect one-dimensional crystal. These cases for \( d = 1, 2 \), and 3, numerical results for MRJ packing for \( d = 4, 5 \), and 6, and theoretical results (Torquato and Stillinger, 2006b), indicating that packings in large dimensions are highly degenerate, suggest that the number of distinct collectively jammed packings (defined in Sec. IV.A) for a fixed large number \( N \) of identical hard spheres rises monotonically with Euclidean dimension \( d \). The questions and issues raised by these differences in the degree of disorder across dimensions emphasizes the need for a geometric-structure approach, to be elaborated in the following.

IV. JAMMING CATEGORIES, ISOSTATICITY, AND POLYTOPES

A. Jamming categories

In much of the ensuing discussion, we treat packings of frictionless congruent spheres of diameter \( D \) in \( \mathbb{R}^d \) in the absence of gravity, i.e., the “Ising model” of jammed sphere packings. Packing spheres is inherently a geometrical problem due to exclusion-volume effects. Indeed, the singular nature of the hard-sphere pair potential (plus infinity or zero for \( r < D \) or \( r \geq D \), respectively, where \( r \) is the pair separation) is crucial because it enables one to be precise about the concept of jamming. Analyzing this model directly is clearly preferable to methods that begin with particle systems having “soft” interactions, which are then intended to mimic packings upon passing to the hard-sphere limit (Donev et al., 2004c).

Three broad and mathematically precise jamming categories of sphere packings can be distinguished depending on the nature of their mechanical stability (Torquato and Stillinger, 2001, 2003). In order of increasing stringency (stability), for a finite sphere packing, these are the following: (1) Local jamming: each particle in the packing is locally trapped by its neighbors (at least \( d + 1 \) contacting particles, not all in the same hemisphere), i.e., it cannot be translated while fixing the positions of all other particles; (2) collective jamming: any locally jammed configuration is collectively jammed if no subset of particles can simultaneously be displaced so that its members move out of contact with one another and with the remainder set; and (3) strict jamming: any collectively jammed configuration that disallows all uniform volume-nonincreasing strains of the system boundary is strictly jammed.

We stress that these hierarchical jamming categories do not exhaust the universe of possible distinctions (Bezdek et al., 1998; Torquato and Stillinger, 2001; Donev et al., 2004a, 2004b), but they span a reasonable spectrum of possibilities. Importantly, the jamming cat-
category of a given sphere configuration depends on the boundary conditions employed. For example, hard-wall boundary conditions (Torquato and Stillinger, 2001) generally yield different jamming classifications from periodic boundary conditions (Donev et al., 2004a). These jamming categories, which are closely related to the concepts of “rigid” and “stable” packings found in the mathematics literature (Bézdek et al., 1998), mean that there can be no “rattlers” (i.e., movable but caged particles) in the packing. Nevertheless, it is the significant majority of spheres that compose the underlying jammed network that confers rigidity to the packing, and in any case, the rattlers could be removed (in computer simulations) without disrupting the jammed remainder. Figure 9 shows examples of ordered locally and collectively jammed packings of disks in two dimensions within hard-wall containers. Observe the square-lattice packing with square hard-wall boundary conditions can only be collectively jammed (not strictly jammed) even in the infinite-volume limit. This is to be contrasted with MRJ packings, where the distinction between collective and strict jamming vanishes in the infinite-volume limit, as discussed in Sec. IVB. Note that the configurations shown in Fig. 8, generated under periodic boundary conditions, are at least collectively jammed.

To emphasize the fact that the jamming category depends on the boundary conditions of a packing, we tabulate whether common periodic structures are locally, collectively, or strictly jammed. Table I gives the jamming classification for such packings with hard-wall boundary conditions. These results are compared to corresponding jamming categories for periodic boundary conditions in Table II. The latter results will depend on the choice of the number of particles $N$ within the fundamental cell.

Rigorous and efficient linear-programming algorithms have been devised to assess whether a particular sphere packing is locally, collectively, or strictly jammed (Donev et al., 2004a, 2004b). It is noteworthy that the jamming categories can now be ascertained in real-system experiments using imaging techniques that enable one to determine configurational coordinates of a packing [e.g., tomography (Aste et al., 2006), confocal microscopy (Brujic et al., 2003), and magnetic resonance imaging (Man et al., 2005)]. Given these coordinates (with high precision), one can rigorously test the jamming category of the experimentally generated packing using the aforementioned linear programming techniques.

### Table I. Classification of some of the common jammed periodic (crystal) packings of identical spheres in two and three dimensions, where $Z$ denotes the contact number per particle and $\phi$ is the packing fraction for the infinite packing (Torquato and Stillinger, 2001). Here hard boundaries are applicable: in two dimensions we use commensurate rectangular boundaries and in three dimensions we use a cubical boundary, with the exception of the hexagonal close-packed crystal in which the natural choice is a hexagonal prism.

| Periodic (crystal) structures | Locally jammed | Collectively jammed | Strictly jammed |
|------------------------------|----------------|---------------------|----------------|
| Honeycomb ($Z=3, \phi=0.605\ldots$) | Yes | No | No |
| Kagomé ($Z=4, \phi=0.680\ldots$) | No\textsuperscript{a} | No\textsuperscript{a} | No\textsuperscript{a} |
| Square ($Z=4, \phi=0.785\ldots$) | Yes | Yes | No |
| Triangular ($Z=6, \phi=0.907\ldots$) | Yes | Yes | Yes |
| Diamond ($Z=4, \phi=0.340\ldots$) | Yes | No | No |
| Simple cubic ($Z=6, \phi=0.524\ldots$) | Yes | Yes | No |
| Body-centered cubic ($Z=8, \phi=0.680\ldots$) | Yes | Yes | No |
| Face-centered cubic ($Z=12, \phi=0.740\ldots$) | Yes | Yes | Yes |
| Hexagonal close packed ($Z=12, \phi=0.740\ldots$) | Yes | Yes | Yes |

\textsuperscript{a}With appropriately placed regular triangular- or hexagonal-shaped boundaries, the kagomé structure is locally, collectively, and strictly jammed.
B. Isostaticity

A packing of \( N \) hard spheres of diameter \( D \) in a \textit{jammed} framework in \( d \)-dimensional Euclidean space is characterized by the \((Nd)\)-dimensional configuration vector of centroid positions \( \mathbf{R} = r^N = \{r_1, \ldots, r_N\} \). Assume that a configuration \( \mathbf{R}_i \) represents a collectively jammed \textit{framework} of a packing (i.e., excluding rattlers) with packing fraction \( \phi_p \), where there are \( M \) interparticle contacts.

Isostatic packings are jammed packings that possess the minimal number of contacts for a jamming category; namely, under periodic boundary conditions, for collective jamming, \( M = 2N - 1 \) and \( 3N - 2 \) for \( d = 2 \) and \( d = 3 \), respectively, and for strict jamming, \( M = 2N + 1 \) and \( 3N + 3 \) for \( d = 2 \) and \( d = 3 \), respectively (Donev et al., 2005c). Thus, we see that the relative differences between isostatic collective and strict jammed packings diminish as \( N \) becomes large, and since the number of degrees of freedom is essentially equal to \( Nd \) [depending on the jamming category and boundary conditions (Donev et al., 2005c)], an isostatic packing has a mean contact number per particle \( Z \) equal to \( 2d \). Collectively or strictly jammed MRJ packings in the infinite-volume limit are isostatic. Note that packings in which \( Z = 2d \) are not necessarily collectively or strictly jammed; for example, we see from Table II that the square and kagomé lattices (with \( Z = 4 \)) and the simple-cubic lattice (with \( Z = 6 \)) are neither collectively nor strictly jammed. Isostaticity has attained a special status in the field and has been closely linked to “generic” or “random” packings (Moukarzel, 1998). In fact, as we will show, isostatic packings can be perfectly ordered; see also Mari et al. (2009). Packings having more contacts than isostatic ones are \textit{hyperstatic} and those having fewer contacts than isostatic packings are \textit{hyperstatic}; for sphere packings, these latter packings cannot be collectively or strictly jammed in the above sense (Donev, Connelly, et al., 2007). The terms \textit{overconstrained} and \textit{underconstrained} (or hypoconstrained), respectively, were used by Donev, Connelly, et al. (2007) to describe such packings.

C. Polytope picture of configuration space

Full understanding of the many-body properties of \( N \) particles in a \( d \)-dimensional container of content (volume) \( V \) is facilitated by viewing the system in its \( dN \)-dimensional configuration space. This is an especially useful approach for hard-particle models and helps us to understand the full range of issues concerning the approach to a jammed state. For the moment, we restrict attention specifically the cases of \( d \)-dimensional hard spheres. When container content \( V \) is very large for fixed \( N \), i.e., when packing fraction \( \phi \approx 0 \), the hard spheres are free to move virtually independently. Consequently, the measure (content) \( C \) of the available multidimensional configuration space is simply \( C \approx V^N \). But decreasing \( V \) enhances the chance for sphere collisions and correspondingly reduces \( C \), the remaining fraction of configuration space that is free of sphere overlaps. The amount of reduction is related exponentially to the excess entropy \( S^{(e)}(N, V) \) for the \( N \)-sphere system,

\[
C(N, V) = V^N \exp[S^{(e)}(N, V)/k_B],
\]

where \( k_B \) is Boltzmann’s constant.

In the low-density regime, the excess entropy admits of a power series expansion in covering fraction \( \phi \),

\[
\frac{S^{(e)}(N, V)}{Nk_B} = N \sum_{n=1} \left( \frac{\beta_n}{n + 1} \right) \left( \frac{\phi}{v_1} \right)^n.
\]

Here \( v_1 \) is the volume of a particle, as indicated in Eq. (2). The \( \beta_n \) are the \textit{irreducible} Mayer cluster integral sums for \( n + 1 \) particles that determine the virial coefficient of order \( n + 1 \) (Mayer and Mayer, 1940). For hard spheres in dimensions \( 1 \leq d \leq 8 \), these coefficients for low orders \( 1 \leq n \leq 3 \) are known exactly, and accurate numerical estimates are available for \( n + 1 \leq 10 \) (Cisby and McCoy, 2006). This power series represents a function of \( \phi \) obtainable by analytic continuation along the positive real axis to represent the thermodynamic behavior for the fluid phase from \( \phi = 0 \) up to the freezing transition, which occurs at \( \phi \approx 0.4911 \) for hard spheres in three di-
dimensions (Noya et al., 2008). This value is slightly below the minimum density $\phi \approx 0.4937$ at which collective jamming of $d=3$ hard spheres is suspected first to occur (Torquato and Stillinger, 2007). Consequently, the available configuration space measured by $C(N,V)$ remains connected in this density range, i.e., any nonoverlap configuration of the $N$ spheres can be connected to any other one by a continuous displacement of the spheres that does not violate the nonoverlap condition.

A general argument has been advanced that thermodynamic functions must experience a subtle but distinctive essential singularity at first-order phase transition points (Andreev, 1964; Fisher and Felderhof, 1970). In particular, this applies to the hard-sphere freezing transition and implies that attempts to analytically continue fluid behavior into a metastable overcompressed state are dubious. Aside from any other arguments that might be brought to bear, this indicates that such extrapolations are fundamentally incapable of identifying unique random jammed states of the hard-sphere system. Nevertheless, it is clear that increasing $\phi$ beyond its value at the thermodynamic freezing point soon initiates partial fragmentation of the previously connected nonoverlap configuration space in finite systems. That is, locally disconnected portions are shed, each to become an individual jammed state displaying its own geometric characteristics. (The jammed tunneled crystals mentioned in Sec. VI are examples of such localized regions near the freezing point.) We elaborate on this point within this section after discussing the polytope picture of configuration space near jamming points.

Consider decreasing the packing fraction slightly in a sphere packing that is at least collectively jammed by reducing the particle diameter by $\Delta D$, $\delta = \Delta D/D \ll 1$, so that the packing fraction is lowered to $\phi = \phi_0 (1 - \delta d)$. We call $\delta$ the jamming gap or distance to jamming. It can be shown that there is a sufficiently small $\delta$ that does not destroy the jamming confinement property, in the sense that the configuration point $R = R_J + \Delta R$ remains trapped in a small neighborhood $J_{AR}$ around $R_J$ (Connelly, 1982). Indeed, there exists a range of positive values of $\delta$ that depends on $N$ and the particle arrangements that maintains the jamming confinement property. We call $\delta$, the threshold value at which jamming is lost. How does $\delta$ scale with $N$ for a particular $d$? An elementary analysis based on the idea that in order for a neighbor pair (or some larger local group) of particles to change places, the surrounding $N-2$ (or $N-3$, . . .) particles must be radially displaced and compressed outward so as to concentrate the requisite free volume around that local interchangeable group concludes that $\delta \sim C N^{-1/d}$, where the constant $C$ depends on the dimension $d$ and the original jammed particle configuration.

It is noteworthy that for fixed $N$ and sufficiently small $\delta$, it can be shown that asymptotically (through first order in $\delta$) the set of displacements that are accessible to the packing approaches a convex limiting polytope (a closed polyhedron in high dimension) $P_{AR} \subseteq J_{AR}$ (Salsburg and Wood, 1962; Stillinger and Salsburg, 1969).

This polytope $P_{AR}$ is determined from the linearized impenetrability equations (Donev et al., 2004a, 2004b) and, for a fixed system center of mass, is necessarily bounded for a jammed configuration. This implies that the number of interparticle contacts $M$ is at least one larger than the dimensionality $d_{CS}$ of the relevant configuration space. Examples of such low-dimensional polytopes for a single locally jammed disk are shown in Fig. 10.

Importantly, for an isostatic contact network, $P_{AR}$ is a simplex (Donev et al., 2005c). A $d$-dimensional simplex in $R^d$ is a closed convex polytope whose $d+1$ vertices (zero-dimensional points) do not all lie in a $(d-1)$-dimensional flat subspace, or alternatively, it is a finite region of $R^d$ enclosed by $d+1$ hyperplanes $[(d-1)$-dimensional “faces”] (e.g., a triangle for $d=2$, a tetrahedron for $d=3$ or a pentatope for $d=4$). For overconstrained jammed packings (e.g., ordered maximally dense states), the limiting high-dimensional polytopes have more faces than simplices do and can be geometrically very complex (Salsburg and Wood, 1962; Stillinger and Salsburg, 1969). The fact that $P_{AR}$ is a simplex for an isostatic packing enables one to derive rigorous results, as we now describe.

Consider adding thermal kinetic energy to a nearly jammed sphere packing in the absence of rattlers. While the system will not be globally ergodic over the full system configuration space and thus not in thermodynamic equilibrium, one can still define a macroscopic pressure $p$ for the trapped but locally ergodic system by considering time averages as the system executes a tightly confined motion around the particular configuration $R_f$. The probability distribution $P(f)$ of the time-averaged interparticle forces $f$ has been rigorously linked to the contact value $r = D$ of the pair-correlation function $g_2$ defined in Sec. II, and this in turn can be related to the distribution of simplex face hyperareas for the limiting polytope (Donev et al., 2005c). Moreover, since the available (free) configuration volume scales in a predictable way with the jamming gap $\delta$, one can show that the reduced pressure is asymptotically given by the free-
measuring the pressure and extrapolating to jamming point has not quite yet been reached, just by since it enables one to determine accurately the true suggests that it is correct. Equation proof yet for this claim, all numerical evidence strongly allowed to go to infinity. Although there is no rigorous leading behavior of the pressure, even as system size is concerned, the extremely narrow connecting filaments that density end points are random jammed packings (Rintoul and Torquato, 1996b; Torquato, 2002), which can be regarded to be glasses. A rapid compression leads to a lower random jammed density than that for a slow compression. The most rapid compression presumably leads to the MRJ state with \( \phi = 0.64 \) (Torquato, 2002). Torquato (1995a, 1995b) reasoned that the functional form of the pressure of the stable liquid branch (which appears to be dominated by an unphysical pole at \( \phi = 1 \)) must be fundamentally different from the free-volume form (10) that applies near jammed states, implying that the equation of state is nonanalytic at the freezing point and proposed the following expression along any constrained metastable branch:

\[
\frac{p}{\rho k_B T} = 1 + 4 \phi \phi_f \frac{1 - \phi_f \phi_f}{1 - \phi \phi_f} \quad \text{for} \quad \phi \leq \phi \leq \phi_f, \tag{11}
\]

where \( \phi_f = 0.491 \) is the packing fraction at the freezing point, \( g_f = 5.72 \) is the corresponding value of the pair-correlation function at contact, and \( \phi_f \) is the jamming density, whose value will depend on which metastable path is chosen. [Torquato (1995a, 1995b) actually considered the more general problem of nearest-neighbor statistics of hard-sphere systems, which required an expression for the equation of state.] Unfortunately, there is no unique metastable branch (see Fig. 11) because it depends on the particular constraints used to generate the metastable states or, in other words, the protocol employed, which again emphasizes one of the themes of this review. Moreover, in practice, metastable states of identical spheres in \( \mathbb{R}^3 \) have an inevitable tendency to crystallize (Rintoul and Torquato, 1996b), but even in binary mixtures of hard spheres chosen to avoid crystallization the dispersion of results and ultimate non-uniqueness of the jammed states still apply. We note that Kamien and Liu (2007) assumed the same free-volume form to fit the pressure of “metastable” states for monodisperse hard spheres as obtained from both numerical and experimental data to determine \( \phi_f \). Their best fit yielded \( \phi_f = 0.6465 \).

We note that density of states (vibrational modes) in packings of soft spheres has been the subject of recent interest (Silbert et al., 2005; Wyart et al., 2005). Collective jamming in hard-sphere packings corresponds to having no “soft modes” in soft-sphere systems, i.e., no unconstrained local or global particle translations are allowed, except those corresponding to rattlers. Observe that it immediately follows that if a hard-sphere packing is collectively jammed to first order in \( \delta \), a corresponding configuration of purely soft repelling particles will pos-

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**FIG. 11.** (Color) The isothermal phase behavior of three-dimensional hard-sphere model in the pressure-packing fraction plane. Increasing the density plays the same role as decreasing temperature of a molecular liquid; see Fig. 3. Three different isothermal densification paths by which a hard-sphere liquid may jam are shown. An infinitesimal compression rate of the liquid traces out the thermodynamic equilibrium path (shown in green), including a discontinuity resulting from the first-order freezing transition to a crystal branch. Rapid compressions of the liquid while suppressing some degree of local order (blue curves) can avoid crystal nucleation (on short time scales) and produce a range of amorphous metastable extensions of the liquid branch that jam only at their density maxima. Adapted from Torquato, 2002.
sess quadratic modes in the vibrational energy spectrum for such a system of soft spheres.

V. ORDER METRICS

The enumeration and classification of both ordered and disordered jammed sphere packings for the various jamming categories is an unresolved problem. Since the difficulty of the complete enumeration of jammed packing configurations rises exponentially with the number of particles, it is desirable to devise a small set of intensive parameters that can characterize packings well. One obvious property of a sphere packing is the packing fraction $\phi$. Another important characteristic of a packing is some measure of its “randomness” or degree of disorder. We have stressed that one ambiguity of the old RCP concept was that randomness was never quantified. To do so is a nontrivial challenge, but even the tentative concept was that randomness was never quantified. To do so is a nontrivial challenge, but even the tentative

classification has been profitable not only to characterize such a system of soft spheres.

A more local measure of bond-orientational order $Q_b$ can be obtained by evaluating the bond order at a fixed orientation. It is seen that $Q = Q_b/Q_b^{\infty}$ lies in the closed interval $[0,1]$ and therefore qualifies as an order metric (Torquato et al., 2000).

A. Specific order metrics

Many relevant order metrics have been devised, but here we describe only some of them. The bond-orientational order $Q_6$ (Steinhardt et al., 1983) in three dimensions is defined in terms of the spherical harmonics $Y_{6m}(\theta, \varphi)$, where $\theta$ and $\varphi$ are the polar and azimuthal angles (relative to a fixed coordinate system) of the near-neighbor bond for particle pair $i$. A near neighbor could be defined as any sphere within a specified local radius (e.g., set by the first minimum of the pair-correlation function beyond contact) or by a sphere within a face-sharing Voronoi polyhedron (Torquato, 2002). The average of $Q_6$ over all of the near-neighbor bonds $N_b$ provides a global measure of symmetries in many-particle systems. Of particular interest is the average when $\ell = 6$, i.e.,

$$Q_6 = \left( \frac{4\pi}{13} \sum_{m=-6}^{6} \frac{1}{N_b} \sum_{i=1}^{N_b} Y_{6m}(\theta_i, \varphi_i) \right)^{1/2},$$

since it reaches its maximum value for the perfect fcc lattice and is zero for a Poisson (uncorrelated) point distribution in the infinite-volume limit (Rintoul and Torquato, 1996a); see Fig. 12 for the two-dimensional analog. It is seen that $Q = Q_b/Q_b^{\infty}$ lies in the closed interval $[0,1]$ and therefore qualifies as an order metric (Torquato et al., 2000).

FIG. 12. (Color online) Two types of order. (a) Bond-orientational order contains information about the orientation of the vectors connecting neighboring particles in the packing of interest (left). If these orientations persist throughout the packing, as they do in a triangular lattice (right), the packing is considered to be perfectly bond-orientationally ordered. (b) Translational order contains information about the relative spacing of particles in the packing of interest at some fixed number density (left) relative to that of the densest-particle configuration at the same number density (right).
each sphere individually, and then averaging over all spheres (Kansal et al., 2002b), i.e.,

$$Q_{6,\text{local}} = \frac{1}{N} \sum_{j=1}^{N} \left( \frac{4\pi}{13} \sum_{m=-6}^{6} \frac{1}{n_{j}} \sum_{i=1}^{12} Y_{6m}(\theta_{i}, \varphi_{i}) \right)^{2} \frac{1}{12},$$

(13)

where $n_{j}$ is the number of nearest neighbors of sphere $j$. This is analogous to the two-dimensional definition of local bond-orientational order studied by Kansal et al. (2000). As noted in that work, such a local measure of order is more sensitive to small crystalline regions within a packing than is its global counterpart $Q_{6}$, and thus avoids the possibility of “destructive” interference between differently oriented crystalline regions.

Using the fcc lattice radial coordination structure as a reference system, one can define a translational order metric $T$ based upon the mean occupation of sphere centers within thin concentric shells around each sphere in the packing as compared to the mean occupation of the same shells in the fcc lattice and the ideal gas at the same packing fraction (Torquato et al., 2000; Truskett et al., 2000), i.e.,

$$T = \frac{\sum_{i=1}^{N_{\text{shells}}} (n_{i} - n_{i}^{\text{ideal}})}{\sum_{i=1}^{N_{\text{shells}}} (n_{i}^{\text{fcc}} - n_{i}^{\text{ideal}})}.$$

Here $n_{i}$ is the average occupancy of the $i^{th}$ shell and $N_{\text{shells}}$ is the total number of shells employed. Figure 12 shows the two-dimensional analog of this order metric.

All of the order metrics defined above were constructed to yield their maximum values of unity (when appropriately normalized) for the densest and most symmetrical (close-packed crystal) packing. Order metrics that are not based on any specific crystal structure have also been devised. For example, two different translational order metrics have been constructed that are based on functionals of the pair-correlation function (Truskett et al., 2000). It has been suggested that local density fluctuations within a “window” of a given size also can be a useful order metric (Torquato and Stillinger, 2003). In particular, calculation of the local number variance for a variety of crystal, quasicrystal, and “hyperuniform” disordered point patterns reveals that it provides a useful rank order of these hyperuniform spatial patterns at large length scales (Torquato and Stillinger, 2003; Zachary and Torquato, 2009). A hyperuniform point pattern is one which the infinite-wavelength density fluctuations vanish or, equivalently, possesses a structure factor $S(k)$ (defined in Sec. II) that tends to zero in the limit $k \rightarrow 0$ (Torquato and Stillinger, 2003).

VI. ORDER MAPS AND OPTIMAL PACKINGS

The geometric-structure classification naturally emphasizes that there is a great diversity in the types of attainable jammed packings with varying magnitudes of overall order, density, and other intensive parameters. The notions of “order maps” in combination with the mathematically precise “jamming categories” enable one to view and characterize well-known packing states, such as the densest sphere packing (Kepler’s conjecture) and MRJ packings as extremal states in the order map for a given jamming category. Indeed, this picture encompasses not only these special jammed states, but an uncountably infinite number of other packings, some of which have only recently been identified as physically significant, e.g., the jamming-threshold states (least dense jammed packings) as well as states between these and MRJ.

The so-called order map (Torquato et al., 2000) provides a useful means to classify packings, jammed or not. It represents any attainable hard-sphere configuration as a point in the $\phi$-$\psi$ plane. This two-parameter description is but a very small subset of the relevant parameters that are necessary to fully characterize a configuration, but it nonetheless enables one to draw important conclusions. For collective jamming, a highly schematic order map has previously been proposed (Torquato et al., 2000).

Here we present a set of refined order maps for each of three jamming categories in $\mathbb{R}^{3}$ (see Fig. 13) based both upon early work (Torquato et al., 2000; Kansal et al., 2002b) and recent investigations (Donev et al., 2004a; Torquato and Stillinger, 2007). Crucially, the order maps shown in Fig. 13 are generally different across jamming categories and independent of the protocols used to generate hard-sphere configurations and for present purposes include rattlers. In practice, one needs to use a variety of protocols to produce jammed configurations in order to populate the interior and to delineate the

B. Characteristics of a good order metric

The specific order metrics have both strengths and weaknesses. This raises the question of what are the characteristics of a good order metric? There is clearly an enormous family of scalar functions that possess the aforementioned three generic properties of an order metric $\phi$, but they may not necessarily be useful ones. It has been suggested that a good order metric should have the following additional properties (Kansal et al., 2002b):

1. sensitivity to any type of ordering without bias toward any reference system;
2. ability to reflect the hierarchy of ordering between prototypical systems given by common physical intuition (e.g., perfect crystals with high symmetry should be highly ordered, followed by quasicrystals, correlated disordered packings without long-range order, and finally spatially uncorrelated or Poisson distributed particles);
3. capacity to detect order at any length scale; and
4. incorporation of both the variety of local coordination patterns and the spatial distribution of such patterns should be included. Moreover, any useful set of order metrics should consistently produce results that are positively correlated with one another (Torquato et al., 2000; Torquato, 2002). The development of improved order metrics deserves continued research attention.
boundary of the jammed regions shown in Fig. 13 (Kantsal et al., 2002b). Moreover, the frequency of occurrence of a particular configuration is irrelevant insofar as the order map is concerned. In other words, the order map emphasizes a geometric-structure approach to packing by characterizing single configurations, regardless of how they were generated or their occurrence probability. In Fig. 13, the white and blue regions represent geometrically possible configurations, while the dark shaded regions are devoid of packings (e.g., maximally dense packings with very low order-metric values do not exist). Clearly, an appreciably reduced region of attainable packings will be occupied by jammed packings, and for any finite packing, its size must decrease as the stringency of the jamming category increases. In the infinite-size limit (not shown in Fig. 13), the regions occupied by collectively and strictly jammed sets become identical. The following extremal points or loci in each jammed category increases. In the infinite-space, and dark shaded regions contain no packings. The locus of points A-A’ correspond to the lowest-density jammed packings. The locus of points B-B’ correspond to the densest jammed packings. Points MRJ represent the maximally random jammed states, i.e., the most disordered states subject to the jamming constraint. It should be noted that the packings represented are not subject to rattler exclusion.

responding displaced position in the order map involves a small reduction in packing fraction from $\phi = 0.64$ and a slight increase in order measure.

(4) More generally, any point along the boundary of the region is an extremal point, residing at the limit of attainability for the jamming category under consideration.

A. Strict jamming

We first discuss the strict-jamming order map. The densest sphere packings in three dimensions which lie along the locus B-B’ are strictly jammed (Torquato and Stillinger, 2001; Donev et al., 2004a) implying that their shear moduli are infinitely large (Torquato et al., 2003). We take point B to correspond to the fcc packing, i.e., it is the most ordered and symmetric densest packing. The other points along the line B-B’ represent the stacking variants of the fcc packing. All can be conveniently viewed as stacks of planar triangular arrays of spheres, within which each sphere contacts six neighbors. These triangular layers can be stacked on one another, fitting spheres of one layer into “pockets” formed by nearest-neighbor triangles in the layer below. At each such layer addition there are two choices of which set of pockets in the layer below are to be filled. Thus, all of the stacking variants can individually be encoded by an infinite binary sequence and therefore constitute an uncountably infinite number of maximally dense packings called the Barlow packings (Barlow, 1883). The most disordered

![FIG. 14. (Color online) Three different optimal strictly jammed packings identified in the rightmost graph of Fig. 13. Left panel: (A) $Z = 7$. Middle panel: MRJ: $Z = 6$ (isostatic). Right panel: (B) $Z = 12$.](image-url)
subset of these is denoted by point $B'$. A rigorous proof that $\phi_{\text{max}} = \pi/\sqrt{18} = 0.74048\ldots$ has only recently appeared (Hales, 2005). In two dimensions, the strictly jammed triangular lattice is the unique densest packing (Fejes Tóth, 1964) and so for $d=2$ the line $B-B'$ collapses to a single point $B$.

The MRJ state is a well-defined minimum in an order map in that for a particular choice of jamming category and order metric it can be identified unambiguously. The MRJ concept is automatically compromised by passing to either the maximal packing density (fcc and its stacking variants) or the minimal possible density for strict jamming (tunneled crystals), thereby causing any reasonable order metric to rise on either side. This eliminates the possibility of a flat horizontal portion of the lower boundary of the jammed accessible region in the $\phi-\psi$ plane in Fig. 13 (multiple MRJ states with different densities) and therefore indicates the uniqueness of the MRJ state in density for a particular order metric. Indeed, at least for collective and strict jamming in three dimensions, a variety of sensible order metrics produce an MRJ state with a packing fraction approximately equal to 0.64 (Kansal et al., 2002b) (see Fig. 8), close to the traditionally advocated density of the RCP state, and with an isostatic mean contact number $Z=6$. This consistency among the different order metrics speaks to the utility of the order-metric concept, even if a perfect order metric has not yet been identified. However, the packing fraction of the MRJ state should not be confused with the MRJ state itself. It is possible to have a rather ordered strictly jammed packing at this same density (Kansal et al., 2002b), as shown in Fig. 13; for example, a jammed but vacancy-diluted fcc lattice packing. This is one reason why the two-parameter order map description of packings is not only useful but necessary. In other words, density alone is far from sufficient in characterizing a jammed packing.

The packings corresponding to the locus of points $A-A'$ have received little attention until recently. Although it has not yet been rigorously established as such, a candidate for the lower limiting packing fraction $\phi_{\text{min}}$ for strictly jammed packings is the subset of “tunneled crystals” that contain linear arrays of vacancies (Torquato and Stillinger, 2007). These relatively sparse structures are generated by stacking planar “honeycomb” layers one upon another, and they all amount to removal of one-third of the spheres from the maximally dense structures with packing fraction $\phi_{\text{max}}$. Consequently, $\phi_{\text{min}} = 2\phi_{\text{max}}/3 = 0.49365\ldots$. Every sphere in a tunneled crystal contacts seven immediate neighbors in one of two possible coordination geometries, and all of the stacking variants exhibit some form of long-range order. It is appropriate to view the two families of maximum-density and minimum-density strictly jammed packings as structural siblings of one another. Note that jammed packings can trivially be created whose densities span the entire range between these extremal cases simply by filling an arbitrary fraction of the vacant sites in any one of the tunneled structures. The dashed lines joining the points $A$ to $B$ and points $A'$ to $B'$ shown in Fig. 13 are the result of sequentially filling the most ordered and disordered tunneled crystals with spheres until the filling process ends with the most ordered and disordered densest Barlow packings, respectively. Interestingly, the tunneled crystals exist at the edge of mechanical stability since removal of any one sphere from the interior would cause the entire packing to collapse. It is noteworthy that Burnell and Sondhi (2008) showed that an infinite subclass of the tunneled crystals has an underlying topology that greatly simplifies the determination of their magnetic phase structure for nearest-neighbor antiferromagnetic interactions and $O(N)$ spins.

It should come as no surprise that ensemble methods that produce “most probable” configurations typically miss interesting extremal points in the order map, such as the locus of points $A-A'$ and the rest of the jamming-region boundary, including remarkably enough the line $B-B'$. However, numerical protocols can be devised to yield unusual extremal jammed states, as discussed in Sec. VII, for example.

Observe that irregular jammed packings can be created in the entire nontrivial range of packing fraction $0.64 < \phi < 0.74048\ldots$ (Torquato et al., 2000; Kansal et al., 2002b) using the LS algorithm. Thus, in the rightmost plot in Fig. 13, the MRJ-$B'$ portion of the boundary of the jammed set, possessing the lowest order metric, is demonstrably achievable. Until recently, no algorithms have produced disordered strictly jammed packings to the left of the MRJ point. A new algorithm described elsewhere (Torquato and Jiao, 2010c) has indeed yielded such packings with $\phi = 0.60$, which are overconstrained with $Z=6.4$, implying that they are more ordered than the MRJ state (see Sec. VII for details). The existence of disordered strictly jammed packings with such anomalously low densities expands conventional thinking about the nature and diversity of disordered packings and places in a broader context those protocols that produce “typical” configurations.

Indeed, there is no fundamental reason why the entire lower boundary of the jammed set between the low-density jamming threshold and MRJ point cannot also be realized. Note that such low-density disordered packings are not so-called “random loose” packings, which are even less well defined than RCP states. For example, it is not clear that the former are even collectively jammed. A necessary first step would be to classify the jamming category of a random loose packing (RLP), which has yet to be done. Therefore, the current tendency in the literature to put so-called RCP and RLP on the same footing as far as jamming is concerned (Song et al., 2008) is premature at best.

In $\mathbb{R}^2$, the so-called “reinforced” kagomé packing with precisely four contacts per particle (in the infinite-packing limit) is evidently the lowest density strictly jammed subpacking of the triangular lattice packing (Donev et al., 2004a) with $\phi_{\text{min}} = \sqrt{3}/8 = 0.68017\ldots$. Note that this packing has the isostatic contact number $Z=4$ and yet is an ordered packing, which runs counter to the
prevalent notion that isostaticity is a consequence of “genericity” or randomness (Moukarzel, 1998).

B. Collective and local jamming

Observe that the locus of points $B-B'$ is invariant under change of the jamming category, as shown in Fig. 13. This is not true of the MRJ state, which will generally have a different location in the local-jamming and collective-jamming order maps. Another important distinction is that it is possible to pack spheres subject only to the weak locally jammed criterion so that the resulting packing fraction is arbitrarily close to zero (Böröczky, 1964; Stillinger et al., 2003). But demanding either collective jamming or strict jamming evidently forces $\phi$ to equal or exceed a lower limit $\phi_{\text{min}}$ that is well above zero.

C. Broader applications to other condensed states of matter

Although methods for characterizing structural order in regular crystalline solids are well established (Ashcroft and Mermin, 1976; Chaikin and Lubensky, 1995), similar techniques for noncrystalline condensed states of matter are not nearly as advanced. The notions of order metrics and order maps have been fruitfully extended to characterize the degree of structural order in condensed phases of matter in which the constituent molecules (jammed or not) possess both attractive and repulsive interactions. This includes the determination of the order maps of models of simple liquids, glasses, and crystals with isotropic interactions (Truskett et al., 2000; Errington et al., 2003), models of water (Errington and Debenedetti, 2001; Errington et al., 2002), and models of amorphous polymers (Stachurski, 2003).

FIG. 15. (Color online) A schematic of the disjoint set of nearly jammed packings (dark shaded regions) that develop in multidimensional configuration space as covering fraction $\phi$ increases. The two axes represent the collection of configurational coordinates. Note that each individual region approaches a polytope in the jamming limit, as discussed in Sec. IV.C.

Figure 15 offers a simple schematic to illustrate this $dN$-dimensional disconnection feature. Several allowed regions with different sizes and shapes are shown. Their boundaries consist of sets of slightly curved hypersurfaces, each of which corresponds to a particle pair contact or contact with a hard wall if present. Particle growth or system compression causes hypersurfaces (numbering at least $dN+1$ for hard walls) to move inward, reducing region content toward zero. The larger the region shown, the larger should be understood its jamming $\phi$ value. The basic issue involved in either laboratory or computer experiments is how and why the various jamming protocols used populate the disconnected regions. Presumably any given algorithm has associated with it a characteristic set of occupation weights, leading in turn to well-defined averages for any property of interest, including packing fraction $\phi$ and any chosen order metric $\psi$. The fact that these averages indeed vary with algorithm is a major point of the present review.

Ensemble methods have been invoked to attach special significance to so-called “typical” or “unique” packings because of their frequency of occurrence in the specific method employed. In particular, significance has been attached to the so-called unique $J$ ( jammed) point, which is suggested to correspond to the onset of collective jamming in soft sphere systems (O’Hern et al., 2003). The order maps described in Sec. V as well as the ensuing discussion demonstrate that claims of such uniqueness overlook the wide variability of packing algorithms and the distribution of configurations that they generate. Individual packing protocols (numerical or experimental) produce jammed packings that are strongly concentrated in isolated pockets of configuration space that are individually selected by those protocols. Therefore, conclusions drawn from any particular protocol are highly specific rather than general in our view.

Indeed, one can create protocols that can lead to
jammed packings at any preselected density with a high probability of occurrence anywhere over a wide density range. Unless it were chosen to be highly restrictive, a typical disk or sphere jamming algorithm applied to a large number \( N \) of particles would be capable of producing a large number of geometrically distinguishable results. In particular, these distinguishable jammed configurations from a given algorithm would show some dispersion in their \( \phi \) and \( \psi \) values. However, upon comparing the distributions of obtained results for a substantial range of particle numbers \( N \) (with fixed boundary conditions), one must expect a narrowing of those distributions with increasing \( N \) owing to operation of a central limit theorem. Indeed, this narrowing would converge individually onto values that are algorithm specific, i.e., different from one another. Figure 16 provides a clear illustration of such narrowing with respect to \( \phi \) distributions, with evident variation over algorithms, as obtained by Jiao et al. (2010b). The examples shown contrast results for two distinctly different sphere system sizes (\( \sim 250 \) and \( \sim 2500 \) particles) and for two different algorithms that have results for disordered jammed packings converging, respectively, onto packing fractions of about 0.60, 0.64, 0.68, and 0.72. The histogram for the lowest density was produced using the new algorithm (Torquato and Jiao, 2010c) noted in Sec. VI, while the other two histograms were generated using the LS algorithm. We stress again that any temptation to select a specific \( \phi \) value as uniquely significant (e.g., 0.64) is primarily based on inadequate sampling of the full range of algorithmic richness and diversity that is available at least in the underlying mathematical theory of sphere jamming.

VIII. ATTRIBUTES OF THE MAXIMALLY RANDOM JAMMED STATE

The MRJ state under the strict-jamming constraint is a prototypical glass (Torquato and Stillinger, 2007) in that it is maximally disordered without any long-range order and perfectly rigid [the elastic moduli are indeed unbounded (Torquato et al., 2003)]. This endows such packings, which are isostatic (\( Z=2d \)), with special attributes (Donev et al., 2005, 2005c). For example, the pair-correlation function \( g_z(r) \) (which provides the distribution of pair distances) of three-dimensional MRJ packings possesses a split second peak (Zallen, 1983), with a prominent discontinuity at twice the sphere diameter, as shown in the left panel of Fig. 17, which is a well-known characteristic of disordered jammed packings. The values \( r=\sqrt{3}D \) and \( r=2D \) are highlighted in Fig. 17, and match the two observed singularities. Interestingly, an integrable power-law divergence \( 1/(r/D-1)^{\alpha} \) with \( \alpha \approx 0.4 \) exists for near contacts (Donev et al., 2005c). No peaks are observed at \( r=\sqrt{2}D \) or \( r=\sqrt{5}D \), which are typical of crystal packings, indicating that there is no detectable undistorted crystal ordering in the packing. We note that in a computational study of stiff “soft” spheres (Silbert et al., 2002), a nearly square-root divergence for near contacts was found.

The MRJ state possesses unusual spatial density fluctuations. It was conjectured (Torquato and Stillinger, 2003) that all strictly jammed saturated packings of congruent spheres (disordered or not) are hyperuniform, i.e., infinite-wavelength density fluctuations vanish or, equivalently, the structure factor \( S(k) \) vanishes in the limit \( k \rightarrow 0 \). (Recall that a saturated packing is one in which no space exists to insert additional particles.) Disordered hyperuniform point distributions are uncommon. Not only was this conjecture verified numerically for an MRJ-like state using a million-particle packing of monodisperse spheres, but it was shown that the structure factor has an unusual nonanalytic linear dependence near the origin (Donev et al., 2005), namely, \( S(k) \sim |k| \) for \( k \rightarrow 0 \), or equivalently, a quasi-long-ranged tail of the total pair-correlation function \( h(r) \sim -r^{-4} \). This same linear nonanalytic behavior of \( S(k) \) near the origin

FIG. 16. (Color online) Packing protocols can be devised that lead to strictly jammed states at any specific density with a high probability of occurrence anywhere over a wide density range. Shown are histograms of jammed packings that are centered around four different packing fractions: \( \phi=0.60, 0.64, 0.68, \) and 0.72, as obtained by Jiao et al. (2010b). The distributions become narrower as the system size increases.
PACKINGS TO MODEL LIQUID STRUCTURE WERE IMPRECISE. THE RIGHT PANEL OF FIG. 8 AND IS ONLY COLLECTIVELY JAMMED AT THIS HIGH DENSITY. BECAUSE SUCH HIGHLY ORDERED PACKINGS OF 10 000 SPHERES IS ALSO FOUND IN SUCH DIVERSE THREE-DIMENSIONAL SYSTEMS AS THE EARLY UNIVERSE (PEEBLES, 1993), GROUND STATE OF LIQUID HELIUM (FEYNMAN AND COHEN, 1956; REATTO AND CHESTER, 1967), AND NONINTERACTING SPIN-POLARIZED FERMI-ONIC GROUND STATES (TORQUATO ET AL., 2008; SCARDICCHIO ET AL., 2009). THE GENERALIZATION OF THE AFOREMENTIONED CONJECTURE THAT DESCRIBES WHEN STRICTLY JAMMED SATURATED PACKINGS OF NONCONGRUENT SPHERE PACKINGS AS WELL AS OTHER PARTICLE SHAPES HAS BEEN GIVEN BY ZACHARY ET AL. (2010). SPECIFICALLY, THE VOID SPACES OF GENERAL MRJ PACKINGS ARE HIGHLY CONSTRAINED BY THE UNDERLYING CONTACT NETWORK, WHICH INDUCE HYPERUNIFORMITY AND QUASI-LONG-RANGE BEHAVIOR OF THE TWO-POINT PROBABILITY FUNCTION FOR THE VOID PHASE.

THE QUASI-LONG-RANGE BEHAVIOR OF \( g_2(r) \) AS WELL AS THE AFOREMENTIONED PAIR-CORRELATION FEATURES DISTINGUISH THE MRJ STATE STRONGLY FROM THAT OF THE EQUILIBRIUM HARD-SPHERE FLUID (HANSEN AND McCLAIN, 1986), WHICH IS CHARACTERIZED BY A STRUCTURE FACTOR THAT IS ANALYTIC AT \( k = 0 \) AND THUS HAS A PAIR-CORRELATION FUNCTION THAT DECAYS EXPONENTIALLY FAST TO UNITY FOR LARGE R. CONSEQUENTLY, EARLY ATTEMPTS (BERNAL, 1960) TO USE DISORDERED JAMMED PACKINGS TO MODEL LIQUID STRUCTURE WERE IMPRECISE.

IT SHOULD BE RECOGNIZED THAT MRJ-LIKE SPHERE PACKINGS CREATED IN PRACTICE VIA COMPUTER ALGORITHMS (TORQUATO ET AL., 2000; O’HERN ET AL., 2002; DONEV ET AL., 2005) OR ACTUAL EXPERIMENTS MAY CONTAIN A SMALL CONCENTRATION OF RATTLERS, THE AVERAGE CONCENTRATION OF WHICH IS PROTOCOL DEPENDENT. THE PACKINGS LEADING TO THE DATA SHOWN IN FIG. 17 CONTAIN BETWEEN 2% AND 3% RATTLERS. THUS, THE HYPERUNIFORMITY PROPERTY OF THE MRJ STATE REQUIRES THAT THE RATTLERS BE RETAINED IN THE PACKING.

IT IS WELL KNOWN THAT LACK OF “FRUSTRATION” (JULIEN ET AL., 1997; TORQUATO, 2002) IN TWO-DIMENSIONAL ANALOGS OF THREE-DIMENSIONAL COMPUTATIONAL AND EXPERIMENTAL PROTOCOLS THAT LEAD TO PUTATIVE RCP STATES RESULT IN PACKINGS OF IDENTICAL DISKS THAT ARE HIGHLY CRYSTALLINE, FORMING RATHER LARGE TRIANGULAR COORDINATION DOMAINS (GRAINS). SUCH A 1000-PARTICLE PACKING WITH \( \phi = 0.88 \) IS SHOWN IN THE RIGHT PANEL OF FIG. 8 AND IS ONLY COLLECTIVELY JAMMED AT THIS HIGH DENSITY. BECAUSE SUCH HIGHLY ORDERED PACKINGS ARE THE MOST PROBABLE OUTCOMES FOR THESE TYPICAL PROTOCOLS, “ENTROPIC MEASURES” OF DISORDER WOULD IDENTIFY THESE AS THE MOST DISORDERED, A MISLEADING CONCLUSION. AN APPROPRIATE ORDER METRIC, ON THE OTHER HAND, IS CAPABLE OF IDENTIFYING A PARTICULAR CONFIGURATION (NOT AN ENSEMBLE OF CONFIGURATIONS) OF CONSIDERABLY LOWER DENSITY (E.G., A JAMMED VACANCY-DILUTED TRIANGULAR LATTICE OR ITS MULTIDOMAIN VARIANT) THAT IS CONSISTENT WITH OUR INTUITIVE NOTIONS OF MAXIMAL DISORDER. HOWEVER, TYPICAL PACKING PROTOCOLS WOULD ALMOST NEVER GENERATE SUCH DISORDERED DISK CONFIGURATIONS BECAUSE OF THEIR INHERENT IMPLICIT BIAS TOWARD UNDILUTED CRYSTALLIZATION. NOTE THAT THE SAME PROBLEMS PERSIST EVEN FOR POLYDISPERSE DISK PACKINGS PROVIDED THAT THE SIZE DISTRIBUTION IS SUFFICIENTLY NARROW.

IMPORTANTLY, PREVIOUSLY REPORTED LOW PACKING FRACTIONS OF 0.82–0.84 FOR SO-CALLED RCP DISK ARRANGEMENTS (BERRYMAN, 1983) WERE FOUND NOT EVEN TO BE COLLECTIVELY JAMMED (DONEV ET AL., 2004A). THIS CONCLUSION DEMONSTRATES THAT THE DISTINCTIONS BETWEEN THE DIFFERENT JAMMING CATEGORIES ARE CRUCIAL. MOREOVER, THE GEOMETRIC-STRUCTURE APPROACH TO JAMMING REVEALS THE BASIC IMPORTANCE OF COLLECTIVE MOTIONS POTENTIALLY INVOLVING AN ARBITRARILY LARGE NUMBER OF PARTICLES. THEREFORE, METHODS THAT ASSUME COLLECTIVE JAMMING BASED ON PACKING FRACTION AND LOCAL CRITERIA, SUCH AS NEAREST-NEIGHBOR COORDINATION AND VORONOI STATISTICS (SONG ET AL., 2008), ARE INCOMPLETE.

IX. PACKINGS OF SPHERES WITH A SIZE DISTRIBUTION

POLYDISPERSY IN THE SIZE OF THE PARTICLES CONSTITUTES A FUNDAMENTAL FEATURE OF THE MICROSTRUCTURE OF A WIDE CLASS OF DISPERSIONS OF TECHNOLOGICAL IMPORTANCE, INCLUDING THOSE INVOLVED IN COMPOSITE SOLID PROPELLANT COMBUSTION (KERSTEIN, 1987), SINTERING OF POWDERS (RAHAMAN, 1995), COLLOIDS (RUSSEL ET AL., 1989), TRANSPORT AND MECHANICAL PROPERTIES OF PARTICULATE COMPOSITE MATERIALS (CHRISTENSEN, 1979), AND FLOW IN PACKED BEDS (SCHEIDEGGER, 1974).
The spheres generally possess a distribution in radius $R$ characterized by a probability density $f(R)$ that normalizes to unity, i.e.,

$$\int_0^{\infty} f(R) dR = 1. \quad (15)$$

The average of any function $w(R)$ is defined by

$$\langle w(R) \rangle = \int_0^{\infty} w(R) f(R) dR. \quad (16)$$

The overall packing fraction $\phi$ of the system is defined as

$$\phi = \rho \langle v_1(R) \rangle, \quad (17)$$

where $\rho$ is the total number density, $v_1(R)$ is given by Eq. (2), and $\langle v_1(R) \rangle$ is the average sphere volume defined by

$$\langle v_1(R) \rangle = \frac{\pi^{d/2}}{\Gamma(1 + d/2)} \langle R^d \rangle. \quad (18)$$

There is a variety of choices for the size distribution $f(R)$ that deserve consideration. Two continuous probability densities that have been widely used to characterize physical phenomena are the Schulz (1939) and logarithmic-normal (Cramer, 1954) distributions. The Schulz distribution is defined as

$$f(R) = \frac{1}{\Gamma(m+1)} \left( \frac{m+1}{\langle R \rangle} \right)^{m+1} R^m \exp \left[ - \frac{(m+1)R}{\langle R \rangle} \right], \quad (19)$$

where $\Gamma(x)$ is the Gamma function. When the parameter $m$ is restricted to non-negative integer values, $\Gamma(m+1) = m!$, and the $n$th moment of this distribution is given by

$$\langle R^n \rangle = \frac{(m+n)!}{m!} \frac{1}{(m+1)^n} \langle R \rangle^n. \quad (20)$$

By increasing $m$, the variance decreases, i.e., the distribution becomes sharper. In the monodisperse limit $m \to \infty$, $f(R) \to \delta(R - \langle R \rangle)$. The case $m = 0$ gives an exponential distribution in which many particles have extremely small radii. By contrast, the logarithmic-normal distribution is defined as

$$f(R) = \frac{1}{\sqrt{2\pi \beta^2}} \exp \left[ - \frac{[\ln(R/\langle R \rangle)]^2}{2 \beta^2} \right], \quad (21)$$

where $\beta^2 = (\langle \ln R \rangle^2) - (\langle \ln R \rangle)^2$. The quantity $\ln R$ has a normal or Gaussian distribution. The $n$th moment is given by

$$\langle R^n \rangle = \exp(n^2 \beta^2/2)(\langle R \rangle)^n. \quad (22)$$

As $\beta^2 \to 0$, $f(R) \to \delta(R - \langle R \rangle)$. Figure 18 shows examples of the Schulz and logarithmic-normal size distributions.

One can obtain corresponding results for spheres with $M$ discrete different sizes from the continuous case by letting

$$f(R) = \sum_{i=1}^{M} \rho_i \delta(R - R_i), \quad (23)$$

where $\rho_i$ and $R_i$ are number density and radius of type-$i$ particles, respectively, and $\rho$ is the total number density. Therefore, the overall volume fraction using Eq. (17) is given by

$$\phi = \sum_{i=1}^{M} \phi^{(i)}, \quad (24)$$

where

$$\phi^{(i)} = \rho_i v_1(R_i) \quad (25)$$

is the packing fraction of the $i$th component.

Sphere packings with a size distribution exhibit intriguing structural features, some of which are only beginning to be understood. It is known, for example, that a relatively small degree of polydispersity can suppress the disorder-order phase transition seen in monodisperse hard-sphere systems (Henderson et al., 1996). Interestingly, equilibrium mixtures of small and large hard spheres can “phase separate” (i.e., the small and large spheres demix) at sufficiently high densities but the precise nature of such phase transitions has not yet been established and is a subject of intense interest; see Dijkstra et al. (1999), and references therein.
Our main interest here is in dense polydisperse packings of spheres, especially jammed ones. Little is rigorously known about the characteristics of such systems. For example, the maximal overall packing fraction of even a monodisperse packing fraction in \( \mathbb{R}^d \), which we denote by \( \phi_{\text{max}}^{(1)} \), for arbitrary values of the mole fractions and radii \( R_1 \) and \( R_2 \) is unknown, not to mention the determination of the corresponding structures. However, one can bound \( \phi_{\text{max}}^{(2)} \) from above and below in terms of the maximal packing fraction \( \phi_{\text{max}}^{(1)} \) for a monodisperse sphere packing in the infinite-volume limit using the following analysis of Torquato (2002). It is clear that \( \phi_{\text{max}}^{(2)} \) is bounded from below by the upper bound on \( \phi_{\text{max}}^{(1)} \). The lower bound \( \phi_{\text{max}}^{(2)} \) is independent of the radii and corresponds to the case when the two components are completely phase separated (demixed), each at the packing fraction \( \phi_{\text{max}}^{(1)} \). Moreover, one can bound \( \phi_{\text{max}}^{(2)} \) from above in terms of the monodisperse value \( \phi_{\text{max}}^{(1)} \) for arbitrary values of \( R_1 \) and \( R_2 \). Specifically, consider a wide separation of sizes \( (R_1 \ll R_2) \) and imagine a sequential process in which the larger spheres are first packed at the maximum density \( \phi_{\text{max}}^{(1)} \) for a monodisperse packing. The remaining interstitial space between the larger spheres can now be packed with the smaller spheres at the packing fraction \( \phi_{\text{max}}^{(1)} \) provided that \( R_1/R_2 \to 0 \). The overall packing fraction in this limit is given by \( 1-(1-\phi_{\text{max}}^{(1)})^2 \), which is an upper bound for any binary packing. Thus, \( \phi_{\text{max}}^{(2)} \leq 1-(1-\pi/(12))^{2}=0.991 \) for \( d=2 \) and \( \phi_{\text{max}}^{(2)} \leq 1-(1-\pi/(18))^{2}=0.933 \) for \( d=3 \), where \( \phi_{\text{max}}^{(1)} \) corresponds to the maximal packing fraction in two and three dimensions, respectively.

The same arguments extend to systems of \( M \) different hard spheres with radii \( R_1, R_2, \ldots, R_M \) in \( \mathbb{R}^d \) (Torquato, 2002). Specifically, the overall maximal packing fraction \( \phi_{\text{max}}^{(M)} \) of such a general mixture in \( \mathbb{R}^d \) (where \( \phi \) is defined by Eq. (17) with Eq. (23)) is bounded from above and below by

\[
\phi_{\text{max}}^{(1)} \leq \phi_{\text{max}}^{(M)} \leq 1 - (1 - \phi_{\text{max}}^{(1)})^M. \tag{26}
\]

The lower bound corresponds to the case when the \( M \) components completely demix, each at the density \( \phi_{\text{max}}^{(1)} \). The upper bound corresponds to the generalization of the aforementioned ideal sequential packing process for arbitrary \( M \) in which we take the limits \( R_1/R_2 \to 0 \), \( R_2/R_3 \to 0 \), \ldots, \( R_{M-1}/R_M \to 0 \). Specific nonsequential protocols (algorithmic or otherwise) that can generate structures that approach the upper bound [Eq. (26)] for arbitrary values of \( M \) are currently unknown and thus the development of such protocols is an open area of research. We see that in the limit \( M \to \infty \) the upper bound approaches unity, corresponding to space-filling polydisperse spheres with an infinitely wide separation in sizes (Herrmann et al., 1990). Furthermore, one can also imagine constructing space-filling polydisperse spheres with a continuous size distribution with sizes ranging to the infinitesimally small (Torquato, 2002).

Jammed binary packings have received some attention but their characterization is far from complete. Here we note work concerned with maximally dense binary packings as well as disordered jammed binary packings in two and three dimensions. Among these cases, we know most about the determination of the maximally dense binary packings in \( \mathbb{R}^2 \). Let \( R_S \) and \( R_L \) denote the radii of the small and large disks (\( R_S \ll R_L \)), the radii ratio \( \alpha = R_S/R_L \) and \( x_S \) be the number fraction of small disks in the entire packing. Ideally, it is desired to obtain \( \phi_{\text{max}} \) as a function of \( \alpha \) and \( x_S \). In practice, we have a sketchy understanding of the surface \( \phi_{\text{max}}(\alpha, x_S) \). Fejes Tóth (1964) reported a number of candidate maximally dense packing arrangements for certain values of the radii ratio in the range \( \alpha \approx 0.154 \ldots \). Maximally dense binary disk packings have been also investigated to determine the stable crystalline phase diagram of such alloys (Likos and Henley, 1993). The determination of \( \phi_{\text{max}} \) for sufficiently small \( \alpha \) amounts to finding the optimal arrangement of the small disks within a tricusp: the nonconvex cavity between three close-packed large disks. A particle-growth Monte Carlo algorithm was used to generate the densest arrangements of small identical disks (ranging in number from 1 to 19) within such a tricusp (Uche et al., 2004). All of these results can be compared to a relatively sharp upper bound on \( \phi_{\text{max}} \) given by

\[
\phi_{\text{max}} \leq \phi_U = \frac{\pi \alpha^2 + 2(1 - \alpha^2) \arcsin(\alpha/(1+\alpha))]}{2\alpha(1+2\alpha)^{1/2}}. \tag{27}
\]

The fraction \( \phi_U \) corresponds to the densest local packing arrangement for three binary disks shown in Fig. 19 and

![FIG. 19. One large disk and two small disks in mutual contact provide the densest local arrangement of binary disks (Florian, 1960). The intersection of the shaded triangle with the three disks yields the local packing fraction \( \phi_U = \frac{\pi \alpha^2 + 2(1 - \alpha^2) \arcsin(\alpha/(1+\alpha))]}{2\alpha(1+2\alpha)^{1/2}} \), where \( \alpha = R_S/R_L \).](image-url)
hence bounds $\phi_{\text{max}}$ from above (Florian, 1960). Inequality (27) also applies to general multicomponent packings, where $\alpha$ is taken to be the ratio of the smallest disk radius to the largest disk radius.

The most comprehensive study of the densest possible packings of binary spheres in $\mathbb{R}^3$ as well as more general size-discrete mixtures has recently been reported by Hudson and Harowell (2008). They generated candidate maximally dense polydisperse packings based on filling the interstices in uniform three-dimensional tilings of space with spheres of different sizes. They were able to find for certain size ratios and compositions a number of space with spheres of different sizes. They were able to fill the interstices in uniform three-dimensional tilings date maximally dense polydisperse packings based on dered jammed packings of binary disks in

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and Torquato, 2007 has been argued that there is no "ideal glass transition" (Parisi and Zamponi, 2005). The existence of an ideal glass transition remains a much debated topic of research.

In three dimensions, it was shown by Schaeft and Sillec (1994) that increasing polydispersity increases the packing fraction over the monodisperse value that an amorphous hard-sphere system can possess. The LS algorithm has been extended to generate jammed sphere packings with a polydispersity in size (Kansal et al., 2002a). It was applied to show that disordered packings with a wide range of packing fractions that exceed 0.64 and varying degrees of disorder can be achieved; see also Chaudhuri et al. (2010). Not surprisingly, the determination of the maximally random jammed (MRJ) state for an arbitrary polydisperse sphere packing is a wide open question. Clusel et al. (2009) carried out a series of experiments to understand polydisperse random packings of spheres. Specifically, they produced three-dimensional random packings of frictionless emulsion droplets with a high degree of size polydispersity and visualize and characterize them using confocal microscopy.

The aforementioned investigations and the many co-
nundrums that remain serve to illustrate the richness of polydisperse jammed packings but further discussion is beyond the scope of this review.

X. PACKINGS OF NONSPHERICAL PARTICLES

Jammed packing characteristics become considerably more complex by allowing for nonspherical particle shapes (Betke and Henk, 2000; Roux, 2000; Williams and Philipse, 2003; Donev, Cisse et al., 2004; Donev, Stillinger et al., 2004; Donev et al., 2005a, 2005b; Man et al., 2005; Conway and Torquato, 2006; Chaikin et al., 2007; Donev, Connelly et al., 2007; Chen, 2008; Hajji-Akbari et al., 2009; Torquato and Jiao, 2009a, 2009b, 2010b; Chen et al., 2010; Jiao et al., 2010a; Kallus et al., 2010). We focus here on the latest developments in this category, specifically for particle shapes that are continuous deformations of a sphere (ellipsoids and superballs) as well as polyhedra. Nonsphericity introduces rotational degrees of freedom not present in sphere packings and can dramatically alter the jamming characteristics from those of sphere packings. [We note in passing that there has been deep and productive examination of the equilibrium phase behavior and transport properties of hard nonspherical particles (Frenkel and Maguire, 1983; Frenkel et al., 1984; Allen, 1993; Yatsenko and Schweizer, 2008).]

Recent developments have provided organizing principles to characterize and classify jammed packings of nonspherical particles in terms of shape symmetry of the particles (Torquato and Jiao, 2009a). We elaborate on these principles in Sec. X.C where we discuss packings of polyhedra. We begin the discussion by considering developments within the last several years on ellipsoid packings, which has spurred much of the resurgent interest in dense packings of nonspherical particles.

A. Ellipsoid packings

One simple generalization of the sphere is an ellipsoid, the family of which is a continuous deformation of a sphere. A three-dimensional ellipsoid is a centrally symmetric body occupying the region

\[(x_1/a)^2 + (x_2/b)^2 + (x_3/c)^2 \leq 1,\]

where $x_i$ (i=1,2,3) are Cartesian coordinates and $a$, $b$, and $c$ are the semiaxes of the ellipsoid. Thus, we see that an ellipsoid is an affine (linear) transformation of the sphere. A spheroid is an ellipsoid in which two of the semiaxes are equal, say, $a=c$, and is a prolate (elongated) spheroid if $b \geq a$ and an oblate (flattened) spheroid if $b \leq a$.

Figure 20 shows how prolate and oblate spheroids are obtained from a sphere by a linear stretch and shrinkage of the space along the axis of symmetry, respectively. This figure also illustrates two other basic points by inscribing the particles within the smallest circular cylinders. The fraction of space occupied by each of the particles within the cylinders is an invariant equal (due to
the affine transformations) to 2/3. This might lead one to believe that the densest packing of ellipsoids is given by an affine transformation of one of the densest sphere packings, but such transformations necessarily lead to ellipsoids that all have exactly the same orientations. Exploiting the rotational degrees of freedom so that the ellipsoids are not all required to have the same orientations turns out to lead to larger packing fractions than that for maximally dense sphere packings. Furthermore, because the fraction of space remains the same in each example shown in Fig. 20, the sometimes popular notion that going to the extreme “needlelike” limit \( b/a \to \infty \) or extreme “disklike” limit \( b/a \to 0 \) can lead to packing fractions \( \phi \) approaching unity is misguided.

Experiments on M&M\textsuperscript{®} candies (spheroidal particles) (Donev, Cisse, et al., 2004; Man et al., 2005) as well as numerical results produced by a modified LS algorithm (Donev et al., 2005a, 2005b) found MRJ-like packings with packing fractions and mean contact numbers that were higher than for spheres. This led to a numerical study of the packing fraction \( \phi \) and mean contact number \( Z \) as a function of the semiaxes (aspect) ratios.

The results were quite dramatic in several respects. It was first shown that \( \phi \) and \( Z \), as a function of aspect ratio, each have a cusp (i.e., nondifferentiable) minimum at the sphere point, and \( \phi \) versus aspect ratio possesses a density maximum; see Fig. 21, which shows the more refined calculations presented by Donev, Connelly, et al. (2007). The existence of a cusp at the sphere point runs counter to the prevailing expectation in the literature that for “generic” (disordered) jammed frictionless particles the total number of (independent) constraints equals the total number of degrees of freedom \( d_f \), implying a mean contact number \( Z=2d_f \) (\( d_f=2 \) for disks, \( d_f=3 \) for ellipses, \( d_f=5 \) for spheres, and \( d_f=6 \) for general ellipsoids). This has been referred to as the isostatic conjecture (Alexander, 1998) or isocounting conjecture (Donev, Connelly, et al., 2007). Since \( d_f \) increases discontinuously with the introduction of rotational degrees of freedom as one makes the particles nonspherical, the isostatic conjecture predicts that \( Z \)

should have a jump increase at aspect ratio \( \alpha=1 \) to a value of \( Z=2 \) for a general ellipsoid. Such a discontinuity was not observed by Donev, Cisse, et al. (2004), rather it was observed that jammed ellipsoid packings are hypostatic \( Z<2d_f \) near the sphere point, and only become nearly isostatic for large aspect ratios. In fact, the isostatic conjecture is only rigorously true for amorphous sphere packings after removal of rattlers; generic nonspherical-particle packings should generally be hypostatic (or subisostatic) (Roux, 2000; Donev, Connelly, et al., 2007).

Until recently, it was accepted that a subisostatic or hypostatic packing of nonspherical particles cannot be rigid (jammed) due to the existence of “floppy” modes (Alexander, 1998), which are unjamming motions (mechanisms) derived within a linear theory of rigidity, i.e., a first-order analysis in the jamming gap \( \delta \) (see Sec. IV.C). The observation that terms of order higher than first generally need to be considered was emphasized by Roux (2000), but this analysis was only developed for spheres. It has recently been rigorously shown that if the curvature of nonspherical particles at their contact points are included in a second-order and higher-order analysis, then hypostatic packings of such particles can indeed be jammed (Donev, Connelly, et al., 2007). For example, ellipsoid packings are generally not jammed to first order in \( \delta \) but are jammed to second order in \( \delta \) (Donev, Connelly, et al., 2007) due to curvature deviations from the sphere.

To illustrate how nonspherical jammed packings can be hypostatic, Fig. 22 shows two simple two-dimensional examples consisting of a few fixed ellipses and a central particle that is translationally and rotationally trapped by the fixed particles. Generally, four contacting particles are required to trap the central one. However, there are special correlated configurations that only require three contacting particles to trap the central one.

FIG. 20. (Color online) A sphere inscribed within the smallest circular cylinder (i) undergoes a linear stretch and shrinkage of the space along the vertical direction leading to a prolate spheroid (ii) and an oblate spheroid (iii). This linear transformation leaves the fraction of space occupied by the spheroids within the cylinders unchanged from that of the fraction of the cylinder volume occupied by the sphere, equal to 2/3.

FIG. 21. (Color online) Density \( \phi \) vs aspect ratio \( \alpha \) for MRJ packings of 10,000 ellipsoids as obtained by Donev, Connelly, et al. (2007). The semiaxes here are 1, \( \alpha \), \( \alpha^3 \). The inset shows the mean contact number \( Z \) as a function of \( \alpha \). Neither the spheroid (oblate or prolate) nor general ellipsoids cases attain their isostatic values of \( Z=10 \) or 12, respectively.
In such instances, the normal vectors at the points of contact intersect at a common point, as is necessary to achieve torque balance. At first glance, such configurations might be dismissed as probability-zero events. However, it was shown that such nongeneric configurations are degenerate (frequently encountered). This “focusing capacity” toward hypostatic values of $Z$ applies to large jammed packings of nonspherical particles and in the case of ellipsoids must be present for sufficiently small aspect ratios for a variety of realistic packing protocols (Donev, Connelly, et al., 2007). It has been suggested that the degree of nongenericity of the packings be quantified by determining the fraction of local coordination configurations in which the central particles have fewer contacting neighbors than the average value $Z$ (Jiao et al., 2010a).

Having established that curvature deviations from the spherical reference shape exert a fundamental influence on constraint counting (Donev, Connelly, et al., 2007), it is clear that similar effects will emerge when the hard-particle interactions are replaced by nonspherical particles interacting with soft short-range repulsive potentials. It immediately follows that jamming to first and second order in $\delta$ for hard nonspherical particles, for example, leads to quadratic and quartic modes in the vibrational energy spectrum for packings of such particles that interact with purely soft repulsive interactions. See Mailman et al. (2009) and Zeravcic et al. (2009) for studies of the latter type for ellipses and ellipsoids, respectively.

It is noteworthy that in striking contrast with MRJ-like sphere packings the rattler concentrations of the MRJ-like ellipsoid packings appear practically to vanish outside of some small neighborhood of the sphere point (Donev, Connelly, et al., 2007). It was shown that MRJ-like packings of nearly spherical ellipsoids can be obtained with $\phi \approx 0.74$, i.e., packing fractions approaching those of the densest three-dimensional sphere packings (Donev, Cisse, et al., 2004). This suggested that there exist ordered ellipsoid packings with appreciably higher densities. Indeed, the densest known ellipsoid packings were subsequently discovered (Donev, Stillinger, et al., 2004; see Fig. 23). These represent a new family of non-Bravais lattice packings of ellipsoids with a packing fraction that always exceeds the density of the densest Bravais lattice packing ($\phi = 0.74048$) as a function of the aspect ratio $\alpha$. The point $\alpha = 1$ corresponding to the face-centered-cubic lattice sphere packing is shown, along with the two sharp maxima in the packing fraction for prolate ellipsoids with $\alpha = 3/2$ and oblate ellipsoids with $\alpha = 1/\sqrt{3}$, as illustrated in the insets. For both $\alpha < 1/\sqrt{3}$ and $\alpha > \sqrt{3}$, the packing fractions of the laminated packings drop off precipitously holding the particle orientations fixed (lines). The presently maximal achievable packing fraction $\phi = 0.7707\ldots$ is highlighted with a thicker line, and is constant for $\alpha \leq 1/\sqrt{3}$ and $\alpha \geq \sqrt{3}$ because there is an affine stretch by an arbitrary factor along a direction in a mirror plane of the particle directions; see Donev, Stillinger, et al. (2004).

For identical ellipse packings in $R^2$, the maximally dense arrangement is obtained by an affine stretching of the optimal triangular-lattice packing of circular disks with $\phi_{\text{max}} = \pi/\sqrt{12}$, which leaves $\phi_{\text{max}}$ unchanged (Fejes Tóth, 1964; Donev, Stillinger, et al., 2004); see Fig. 24. This maximally dense ellipse packing is not rotationally jammed for any noncircular shape, since it can be sheared continuously without introducing overlap or changing the density (Donev, Connelly, et al., 2007). The packing is, however, strictly translationally jammed.

### B. Superball packings

Virtually all systematic investigations of the densest particle packings have been carried out for convex objects. A $d$-dimensional superball is a centrally symmetric body in $d$-dimensional Euclidean space occupying the region...
dimensional counterparts of ellipses ($d \geq 3$) in order to improve on $\phi_{\text{max}}$ for spheres. Even for three-dimensional ellipsoids, $\phi_{\text{max}}$ increases smoothly as the aspect ratios of the semiaxes vary from unity (Donev, Stillinger, et al., 2004) and hence has no cusp at the sphere point. In fact, congruent three-dimensional ellipsoid packings have a cusplike behavior at the sphere point only when they are randomly jammed (Donev, Cisse, et al., 2004).

Increasing the dimensionality of the particle imbues the optimal “superball” packings with structural characteristics that are richer than their two-dimensional counterparts (Jiao et al., 2009). For example, in three dimensions, a superball is a perfect sphere at $p=1$ but can possess two types of shape anisotropy: cubelike shapes (three-dimensional analog of the square symmetry of the superdisk) and octahedronlike shapes, depending on the value of the deformation parameter $p$ (see Fig. 26). As $p$ continuously increases from 1 to $\infty$, we have a family of convex superballs with cubelike shapes; at the limit $p = \infty$, the superball is a perfect cube. As $p$ decreases from 1 to 0.5, a family of convex superballs with octahedronlike shapes are obtained; at $p=0.5$, the superball becomes a regular octahedron. When $p<0.5$, the superball still possesses an octahedronlike shape but is now concave, becoming a three-dimensional “cross” in the limit $p \rightarrow 0$. Note that the cube and regular octahedron (two of the five Platonic polyhedra) have the same group symmetry (i.e., they have the same 48 space-group elements) because they are dual to each other. Two polyhedra are dual to each other if the vertices of one correspond to the faces of the other.

Jiao et al. (2009) obtained analytical constructions for the densest known superball packings for all convex and concave cases. The candidate maximally dense packings are certain families of Bravais lattice packings (in which each particle has 12 contacting neighbors) possessing the global symmetries that are consistent with the symmetries of a superball. Evidence is provided that these packings are indeed optimal, and Torquato and Jiao (2009b) conjectured that the densest packings of all convex superballs are their densest lattice packings; see Fig. 27. For superballs in the cubic regime ($p>1$), the candidate optimal packings are achieved by two families of Bravais lattice packings ($C_0$ and $C_1$ lattices) possessing twofold and threefold rotational symmetries, respectively, which can both be considered to be continuous deformations of the fcc lattice. For superballs in the octahedral regime ($0.5<p<1$), there are also two families of Bravais lattices ($O_0$ and $O_1$ lattices) obtainable from continuous deformations of the fcc lattice keeping its fourfold rotational symmetry and from the densest lat-

\begin{equation}
|x_1|^{2p} + |x_2|^{2p} + \cdots + |x_d|^{2p} \leq 1,
\end{equation}

where $x_i$ ($i=1, \ldots, d$) are Cartesian coordinates and $p \geq 0$ is the deformation parameter (not pressure as denoted in Sec. IV.C), which controls the extent to which the particle shape has deformed from that of a $d$-dimensional sphere ($p=1$). Thus, superballs constitute a large family of both convex ($p>0.5$) and concave ($0 \leq p < 0.5$) particles (see Fig. 26).

In general, a “superdisk,” the designation in the two-dimensional case, possesses square symmetry. As $p$ moves away from unity, two families of superdisks with square symmetry can be obtained depending on whether $p<1$ or $p>1$ such that there is a 45° rotation with respect to the “protuberances.” When $p<0.5$, the superdisk is concave (see Fig. 25). The candidate maximally dense packings were recently proposed for all convex and concave shapes (Jiao et al., 2008). These are achieved by two different families of Bravais lattice packings such that $\phi_{\text{max}}$ is nonanalytic at the “circular-disk” point ($p=1$) and increases significantly as $p$ moves away from unity. The broken rotational symmetry of superdisks influences the packing characteristics in a nontrivial way that is distinctly different from ellipse packings. Recall that for ellipse packings, no improvement over the maximal circle packing density is possible. For superdisks, one can take advantage of the fourfold rotationally symmetric shape of the particle to obtain a substantial improvement on the maximal circle packing density. By contrast, one needs to use higher-

![FIG. 25.](Color online) Superdisks with different values of the deformation parameter $p$.
tice packing for regular octahedra (Minkowski, 1905; Betke and Henk, 2000), keeping the translational symmetry of the projected lattice on the coordinate planes, which are apparently optimal in the vicinity of the sphere point and the octahedron point, respectively (see Fig. 27).

The proposed maximal packing density \( \phi_{\text{max}} \) as a function of deformation parameter \( p \) is plotted in Fig. 28. As \( p \) increases from unity, the initial increase in \( \phi_{\text{max}} \) is linear in \( p - 1 \) and subsequently \( \phi_{\text{max}} \) increases monotonically with \( p \) until it reaches unity as the particle shape becomes more like a cube, which is more efficient at filling space than a sphere. These characteristics stand in contrast to those of the densest known ellipsoid packings, achieved by certain crystal arrangements of congruent spheroids with a two-particle basis, whose packing density as a function of aspect ratios has zero initial slope and is bounded from above by a value of 0.7707... (Donev, Stillinger, et al., 2004). As \( p \) decreases from unity, the initial increase of \( \phi_{\text{max}} \) is linear in \( 1 - p \). Thus, \( \phi_{\text{max}} \) is a nonanalytic function of \( p \) at \( p = 1 \), which is consistent with conclusions made about superdisk packings (Jiao et al., 2008). However, the behavior of \( \phi_{\text{max}} \) as the superball shape moves off the sphere point is distinctly different from that of optimal spheroid packings, for which \( \phi_{\text{max}} \) increases smoothly as the aspect ratios of the semiaxes vary from unity and hence has no cusp at the sphere point (Donev, Stillinger, et al., 2004).

The density of congruent ellipsoid packings (not \( \phi_{\text{max}} \)) has a cusplike behavior at the sphere point only when the packings are randomly jammed (Donev, Cisse, et al., 2004). The distinction between the two systems results from different broken rotational symmetries. For spheroids, the continuous rotational symmetry is only partially broken, i.e., spheroids still possess one rotationally symmetric axis, and the three coordinate directions are not equivalent, which facilitates dense non-Bravais packings. For superballs, the continuous rotational symmetry of a sphere is completely broken and the three coordinate directions are equivalently fourfold rotationally symmetric directions of the particle. Thus, a superball is less symmetric but more isotropic than an ellipsoid, a shape characteristic which apparently favors dense Bravais lattice packings. The broken symmetry of superballs makes their shapes more efficient in tiling space and thus results in a larger and faster increase in the packing density as the shape moves away from the sphere point.

As \( p \) decreases from 0.5, the superballs become concave particles, but they still possess octahedronlike shapes [see Fig. 26(a)]. The lack of simulation techniques to generate concave superball packings makes it difficult to find the optimal packings for the entire range of concave shapes \((0 < p < 0.5)\). However, based on their conclusions for convex superball packings, Jiao et al. (2009) conjectured that near the octahedron point, the optimal packings possess similar translational symmetry to that of the \( O_1 \)-lattice packing, and based on theoretical considerations proposed candidate optimal packings for all concave cases with a density versus \( p \) as shown in Fig. 29.

Jiao et al. (2010a) determined the packing fractions of MRJ packings of binary superdisks in \( R^2 \) and monodispersed superballs in \( R^3 \). They found that the MRJ densities of such packings increase dramatically and nonanalytically as one moves away from the circulardisk and sphere point \((p = 1)\). Moreover, these disordered packings were demonstrated to be hypostatic, i.e., the average number of contacting neighbors is less than twice the total number of degrees of freedom per particle, and the packings are mechanically stable. As a result, the local arrangements of the particles are necessarily nontrivially correlated to achieve jamming and hence “nongeneric.” The degree of nongenericity of the packings was quantitatively characterized by determining the fraction of local coordination structures in which the central particles have fewer contacting neighbors than...
the average value $Z$. Figure 30 shows local packing structures with more contacts than average and those with less contacts than average in MRJ binary superdisk packings for different $p$ values. It was also explicitly shown that such seemingly “special” packing configurations are counterintuitively not rare. As the anisotropy of the particles increases, it was shown that the fraction of rattlers decreases while the minimal orientational order (as measured by the cubic order metric) increases. These novel characteristics result from the unique rotational symmetry breaking manner of superdisk and superball particles.

C. Polyhedron packings

Until some recent developments, little was known about the densest packings of polyhedral particles. The difficulty in obtaining dense packings of polyhedra is related to their complex rotational degrees of freedom and to the nonsmooth nature of their shapes.

The Platonic solids (mentioned in Plato’s Timaeus) are convex polyhedra with faces composed of congruent convex regular polygons. There are exactly five such solids: the tetrahedron (P1), icosahedron (P2), dodecahedron (P3), octahedron (P4), and cube (P5); see Fig. 31. [We note that viral capsids often have icosahedral symmetry; see, for example, Zandi et al. (2004).] Here we focus on the problem of the determination of the densest packings of each of the Platonic solids in three-dimensional Euclidean space $\mathbb{R}^3$, except for the cube, which is the only Platonic solid that tiles space.

It is useful to highlight some basic geometrical properties of the Platonic solids that we employ in subsequent sections of this review. The dihedral angle $\theta$ is the interior angle between any two face planes and is given by

\[
\theta = \frac{\pi}{n}
\]

where $n$ is the number of face angles meeting at a vertex.

FIG. 28. (Color online) Density vs deformation parameter $p$ for the packings of convex superballs (Jiao et al., 2009). Inset: Around $p_c = 1.1509\ldots$, the two curves are almost locally parallel to each other.

FIG. 29. (Color online) Density vs deformation parameter $p$ for the lattice packings of concave superballs (Jiao et al., 2009). Inset: A concave superball with $p = 0.1$, which will become a three-dimensional cross at the limit $p \to 0$.

FIG. 30. (Color online) Local packing structures with more contacts than average (shown in blue) and those with less contacts than average (shown in pink) in two-dimensional MRJ binary superdisk packings for different values of the deformation parameter $p$, as obtained from Jiao et al. (2010a). Note that the degree of nongenericity decreases (fraction of space occupied by the pink colored particles decreases) as $p$ moves away from its sphere-point value. Here the size ratio (diameter of large superdisks divided by the diameter of the small superdisks) is 1.4 and the molar ratio (number of large superdisks divided by the number of small superdisks) is $1/3$.

FIG. 31. (Color online) The five Platonic solids: tetrahedron (P1), icosahedron (P2), dodecahedron (P3), octahedron (P4), and cube (P5).
where \( p \) is the number of sides of each face and \( q \) is the number of faces meeting at each vertex (Coxeter, 1973). Thus, \( \theta = \frac{\cos(\pi/q)}{\sin(\pi/p)} \), (30)

![Diagram of Archimedean solids]

FIG. 32. (Color online) The 13 Archimedean solids: truncated tetrahedron (A1), truncated icosahedron (A2), snub cube (A3), snub dodecahedron (A4), rhombicosidodecahedron (A5), truncated icosidodecahedron (A6), truncated icosahedron (A7), icosidodecahedron (A8), rhombicuboctahedron (A9), truncated dodecahedron (A10), cuboctahedron (A11), truncated cube (A12), and truncated octahedron (A13). This typical enumeration of the Archimedean solids does not count the chiral forms (not shown) of the snub cube (A3) and snub dodecahedron (A4), which implies that the left-handed and right-handed forms of each of these pairs lack central symmetry. The remaining 11 Archimedean solids are nonchiral (i.e., each solid is superposable on its mirror image) and the only noncentrally symmetric one among these is the truncated tetrahedron.

([P2]–[P5] and [A2]–[A13]) distinguish their dense packing arrangements from those of the noncentrally symmetric ones ([P1] and [A1]) in a fundamental way.

Tetrahedral tilings of space underlie many different molecular systems (Conway and Torquato, 2006). Since regular tetrahedra cannot tile space, it is of interest to determine the highest density that such packings of particles can achieve (one of Hilbert’s 18th problem set). It is of interest to note that the densest Bravais lattice packing of tetrahedra (which requires all of the tetrahedra to have the same orientations) has \( \phi = 18/49 = 0.367 \ldots \) and each tetrahedron touches 14 others. Recently, Conway and Torquato (2006) showed that the maximally dense tetrahedron packing cannot be a Bravais lattice (because dense tetrahedron packings favor face-to-face contacts) and found non-Bravais lattice (periodic) packings of regular tetrahedra with \( \phi = 0.72 \). One such packing is based upon the filling of “imaginary” icosahedra with the densest arrangement of 20 tetrahedra and then arranging the imaginary icosahedra in their densest lattice packing configuration. Using “tetrahedral” dice, Chaikin et al. (2007) experimentally generated jammed disordered packings of such dice with \( \phi = 0.75 \); see also Jaoshvili et al. (2010) for a refined version of this work. However, because these dice are not perfect tetrahedra (vertices and edges are slightly rounded), a definitive conclusion could not be reached. Using physical models and computer algebra system, Chen (2008) discovered a dense periodic arrangement of tetrahedra with \( \phi = 0.7786 \ldots \) which exceeds the density of the densest sphere packing by an appreciable amount.
In an attempt to find even denser packings of tetrahedra, Torquato and Jiao (2009a, 2009b) formulated the problem of generating dense packings of polyhedra within an adaptive fundamental cell subject to periodic boundary conditions as an optimization problem, which they call the adaptive shrinking cell (ASC) scheme. Starting from a variety of initial unjammed configurations, this optimization procedure uses both a sequential search of the configurational space of the particles and the space of lattices via an adaptive fundamental cell that shrinks on average to obtain dense packings. This was used to obtain a tetrahedron packing consisting of 72 particles per fundamental cell with packing fraction $\phi = 0.782\ldots$ (Torquato and Jiao, 2009a). Using 314 particles per fundamental cell and starting from an “equilibrated” low-density liquid configuration, they were able to improve the packing fraction to $\phi = 0.823\ldots$ (Torquato and Jiao, 2009b). This packing arrangement interestingly lacks any long-range order. Haji-Akbari et al. (2009) numerically constructed a periodic packing of tetrahedra made of parallel stacks of “rings” around “pentagonal” dipyramids consisting of 82 particles per fundamental cell and a density $\phi = 0.8503\ldots$.

Kallus et al. (2010) found a remarkably simple “uniform” packing of tetrahedra with high symmetry consisting of only four particles per fundamental cell (two “dimers”) with packing fraction $\phi = \frac{100}{117} = 0.854 700\ldots$. A uniform packing has a symmetry (in this case a point inversion symmetry) that takes one tetrahedron to another. A dimer is composed of a pair of regular tetrahedra that exactly share a common face. Torquato and Jiao (2010a) subsequently presented an analytical formulation to construct a three-parameter family of dense uniform dimer packings of tetrahedra again with four particles per fundamental cell. (A uniform dimer packing of tetrahedra has a point of inversion symmetry operation that takes any dimer into another.) Making an assumption about one of these parameters resulted in a two-parameter family, including those with a packing fraction as high as $\phi = \frac{12250}{14319} = 0.855 506\ldots$ (see left panel of Fig. 33). Chen et al. (2010) recognized that such an assumption was made in the formulation of Torquato and Jiao (2010a) and employed a similar formalism to obtain a three-parameter family of tetrahedron packings, including the densest known dimer packings of tetrahedra with the slightly higher packing fraction $\phi = \frac{4671}{5319} = 0.856 347\ldots$ (see right panel of Fig. 33). The most general analytical formulation to date to construct dense periodic packings of tetrahedra with four particles per fundamental cell was carried out by Torquato and Jiao (2010b). This study involved a six-parameter family of dense tetrahedron packings that includes as special cases all of the aforementioned “dimer” packings of tetrahedra, including the densest known packings with packing fraction $\phi = \frac{4671}{5319} = 0.856 347\ldots$. This recent investigation strongly suggests that the latter set of packings are the densest among all packings with a four-particle basis. Whether these packings are the densest packings of tetrahedra among all packings is an open question for reasons given by Torquato and Jiao (2010b).

Using the ASC scheme and a variety of initial conditions with multiple particles in the fundamental cell Torquato and Jiao (2009a, 2009b) were also able to find the densest known packings of other nontiling Platonic solids like the densest tetrahedron packing, which must be a non-Bravais lattice packing, the densest packings of the octahedron, dodecahedron, and icosahedron (three nontiling Platonic solids) with densities 0.947\ldots, 0.904\ldots, and 0.836\ldots, respectively. Unlike the densest tetrahedron packing, which must be a non-Bravais lattice packing, the densest packings of the other nontiling Platonic solids found by the algorithm are their previously known optimal (Bravais) lattice packings (Minkowski, 1905; Betke and Henk, 2000); see Fig. 34. These simulation results as well as other theoretical considerations, described below, led them to general organizing principles concerning the densest packings of a class of nonspherical particles.

Rigorous upper bounds on the maximal packing fraction $\phi_{\text{max}}$ of nonspherical particle packings of general shape can be used to assess the packing efficiency of a particular dense packing of such particles. However, it has been highly challenging to formulate upper bounds for nontiling particle packings that are nontrivially less than unity. It has recently been shown that $\phi_{\text{max}}$ of a packing of congruent nonspherical particles of volume $v_p$ in $\mathbb{R}^3$ is bounded from above according to

$$\phi_{\text{max}} \leq \phi_{\text{max}}^U = \min \left[ \frac{v_p}{\frac{\pi}{\sqrt{18}}}, 1 \right],$$

where $v_s$ is the volume of the largest sphere that can be inscribed in the nonspherical particle and $\pi/\sqrt{18}$ is the maximal sphere-packing density (Torquato and Jiao,
Dodecahedron packing with packing fraction of the densest lattice packing is not a congruent polyhedron without central symmetry generally their corresponding optimal lattice packings. Since bound circumradius to the inradius when particles are central symmetry enables maximal face-to-face contacts for a higher packing fraction. They also showed that substantial face-to-face contacts between any of the centrally symmetric Platonic and Archimedean solids asphericities and explain the corresponding small differences between \( \phi_{t}^{\text{max}} \) and the packing fraction of the densest lattice packing \( \phi_{l}^{\text{max}} \).

Additional remarks

It is noteworthy that the densest known packings of all of the Platonic and Archimedean solids as well as the densest known packings of superballs (Jiao et al., 2009) and ellipsoids (Donev, Stillinger, et al., 2004) in \( \mathbb{R}^3 \) have packing fractions that exceed the optimal sphere packing value \( \phi_{S}^{\text{max}} = \pi / \sqrt{18} = 0.7408... \). These results are consistent with a conjecture of Ulam who proposed without any justification [in a private communication to Gardner (2001)] that the optimal packing fraction for congruent sphere packings is smaller than that for any other convex body. The sphere is perfectly isotropic with an asphericity \( \gamma \) of unity, and therefore its rotational degrees of freedom are irrelevant in affecting its packing characteristics. On the other hand, each of the aforementioned convex nonspherical particles break the continuous rotational symmetry of the sphere and thus its broken symmetry can be exploited to yield the densest possible packings, which might be expected to exceed \( \phi_{S}^{\text{max}} = \pi / \sqrt{18} = 0.7408... \) (Torquato and Jiao, 2009a). However, broken rotational symmetry in and of itself may not be sufficient to satisfy Ulam's conjecture if the convex particle has a little or no symmetry (Torquato and Jiao, 2009a).

Apparently, the two-dimensional analog of Ulam's conjecture [optimal density of congruent circle packings \( \phi_{C}^{\text{max}} = \pi / \sqrt{12} = 0.906899... \) is smaller than that for any other convex two-dimensional body] is false. The "smoothed" octagon constructed by Reinhardt (1934) is conjectured to have smallest optimal packing fraction \( \phi_{C}^{\text{max}} = (8 - 4 \sqrt{2} - \ln 2) / (2 \sqrt{2} - 1) = 0.902414... \) among all congruent centrally symmetric planar particles.

It will also be interesting to determine whether conjecture 1 can be extended to other polyhedral packings. The infinite families of prisms and antiprisms provide such a class of packings. A prism is a polyhedron having bases that are parallel, congruent polygons and sides that are parallelograms. An antiprism is a polyhedron having bases that are parallel congruent polygons and
sides that are alternating bands of triangles. They are generally much less symmetric than either the Platonic or Archimedean solids. Moreover, even the centrally symmetric prisms and antiprisms generally do not possess three equivalent directions. Thus, it is less obvious whether Bravais lattices would still provide the optimal packings for these solids, except for prisms that tile space, e.g., hexagonal prism or rhombical prisms (Torquato and Jiao, 2009b). Torquato and Jiao (2009b) also commented on the validity of conjecture 1 to polytopes in four and higher dimensions.

XI. PACKING SPHERES IN HIGH-DIMENSIONAL EUCLIDEAN SPACES

There has been resurgent interest in sphere packings for \( d > 3 \) in both the physical and mathematical sciences (Conway and Sloane, 1995; Frisch and Percus, 1999; Elkie, 2000; Parisi and Slanina, 2000; Cohn, 2002; Cohn and Elkies, 2003; Parisi and Zamponi, 2006, 2010; Skoge et al., 2006; Torquato, 2006; Torquato and Stillinger, 2006a, 2006b; Rohrmann and Santos, 2007; Adda-Bedia et al., 2008; Scardicchio et al., 2008; van Meel, Charbonneau et al., 2009; van Meel, Frenkel, and Charbonneau, 2009, Cohn and Kumar, 2009; Lue et al., 2010). Remarkably, the optimal way of sending digital signals over noisy channels corresponds to the densest sphere packing in a high-dimensional space (Shannon, 1948; Conway and Sloane, 1998). These “error-correcting” codes underlie a variety of systems in digital communications and storage, including compact disks, cell phones, and the Internet; see Fig. 35. Physicists have studied sphere packings in high dimensions to gain insight into liquid and glassy states of matter as well as phase behavior in lower dimensions (Frisch and Percus, 1999; Parisi and Slanina, 2000; Parisi and Zamponi, 2006, 2010; Skoge et al., 2006; Rohrmann and Santos, 2007; Adda-Bedia et al., 2008; van Meel, Charbonneau et al., 2009). Finding the densest packings in arbitrary dimension is a problem of long-standing interest in discrete geometry (Conway and Sloane, 1998). A comprehensive review of this subject is beyond the scope of this article. We instead summarize the relevant literature leading to a recent development beyond the scope of this article. We instead summarize the relevant literature leading to a recent development.

The remarkably symmetric E8 and Leech lattices in \( \mathbb{R}^8 \) and \( \mathbb{R}^{24} \), respectively, are most likely the densest packings in these dimensions (Cohn and Kumar, 2009). Table III lists the densest known sphere packings in \( \mathbb{R}^d \) for selected \( d \). Interestingly, the nonlattice (periodic) packing \( P_{10c} \) (with 40 spheres per fundamental cell) is the densest known packing in \( \mathbb{R}^{10} \), which is the lowest dimension in which the best known packing is not a (Bravais) lattice. It is noteworthy that for sufficiently large \( d \), lattice packings are most likely not the densest (see Fig. 36), but it becomes increasingly difficult to find explicit dense packing constructions as \( d \) increases. Indeed, the problem of finding the shortest lattice vector in a particular lattice packing (densest lattice packing) grows superexponentially with \( d \) and is in the class of NP-hard (nondeterministic polynomial-time hard) problems (Ajtai, 1998).

For large \( d \), the best that one can do theoretically is to devise upper and lower bounds on \( \phi_{\text{max}} \) (Conway and Sloane, 1998). The nonconstructive lower bound of Minkowski (1905) established the existence of reasonably dense lattice packings. He found that the maximal packing fraction \( \phi_{\text{max}}^{L} \) among all lattice packings for \( d \geq 2 \) satisfies...
TABLE III. The densest known sphere packings in $\mathbb{R}^d$ for selected $d$. Except for the nonlattice packing $P_{10}$, in $\mathbb{R}^{10}$, all of the other densest known packings listed are lattice packings: $Z$ is the integer lattice, $A_2$ is the triangular lattice, $D_3$ is the checkerboard lattice (a generalization of the fcc lattice), $E_8$ is one of the root lattices, and $\Lambda_4$ is the laminated lattice. See Conway and Sloane (1998) for further details.

| Dimension, $d$ | Packing structure | Packing fraction, $\phi$ |
|---------------|------------------|------------------------|
| 1             | $Z$              | 1                      |
| 2             | $A_2$            | $\pi/\sqrt{12} \approx 0.9068...$ |
| 3             | $D_3$            | $\pi/\sqrt{18} \approx 0.7404...$ |
| 4             | $D_4$            | $\pi^2/16 \approx 0.6168...$ |
| 5             | $D_5$            | $2\pi^2/(30\sqrt{2}) \approx 0.4652...$ |
| 6             | $E_6$            | $3\pi^2/(144\sqrt{3}) \approx 0.3729...$ |
| 7             | $E_7$            | $4\pi^3/105 \approx 0.2952...$ |
| 8             | $E_8$            | $4\pi^3/384 \approx 0.2536...$ |
| 9             | $\Lambda_8$      | $2\pi^4/(945\sqrt{2}) \approx 0.1457...$ |
| 10            | $P_{10}$         | $5\pi^3/3072 \approx 0.09961...$ |
| 16            | $\Lambda_{16}$   | $9\pi^8/645120 \approx 0.01470...$ |
| 24            | $\Lambda_{24}$   | $\pi^{12}/479001600 \approx 0.001929...$ |

$$\phi_{\text{max}}^L \geq \zeta(d)/\gamma^{d-1},$$

where $\zeta(d) = \sum_{k=1}^{\infty} k^{-d}$ is the Riemann zeta function. Note that for large values of $d$, the asymptotic behavior of the Minkowski lower bound is controlled by $2^{-d}$.

Since 1905, many extensions and generalizations of Eq. (32) have been obtained (Davenport and Rogers, 1947; Ball, 1992; Conway and Sloane, 1998; Vance, 2009), but none of these investigations have been able to improve upon the dominant exponential term $2^{-d}$. It is useful to note that the packing fraction of a saturated packing of congruent spheres in $\mathbb{R}^d$ for all $d$ satisfies

$$\phi \geq 1/2^d. \quad (33)$$

The proof is trivial. A saturated packing of congruent spheres of unit diameter and packing fraction $\phi$ in $\mathbb{R}^d$ has the property that each point in space lies within a unit distance from the center of some sphere. Thus, a covering of the space is achieved if each sphere center is encompassed by a sphere of unit radius and the packing fraction of this covering is $2^d\phi \geq 1$. Thus, the bound [Eq. (33)], which is sometimes called the “greedy” lower bound, has the same dominant exponential term as Eq. (32).

We know that there exists a disordered but unsaturated packing construction, known as the “ghost” random sequential addition (RSA) packing (Torquato and Stillinger, 2006a), that achieves the packing $2^d$ for any $d$. This packing, shown in Fig. 37 and described in its caption, is a generalization of the standard RSA packing, also described in the caption of Fig. 37. It was shown that all of the $n$-particle correlation functions of this nonequilibrium model, in a certain limit, can be obtained analytically for all allowable densities and in any dimension. This represents the first exactly solvable disordered sphere-packing model in arbitrary dimension.

[Note that Matérn (1986) gave an expression for the pair-correlation function for this model.] The existence of this unjammed disordered packing strongly suggests that Bravais lattice packings (which are almost surely unsaturated for sufficiently large $d$) are far from optimal for large $d$. Further support for this conclusion is the fact that the maximal “saturation” packing fraction of the standard disordered RSA packing apparently scales as $d^{-2}$ or possibly $d \ln(d)2^{-d}$ for large $d$ (Torquato et al., 2006). Spheres in both the ghost and standard RSA packings cannot form interparticle contacts, which appears to be a crucial attribute to obtain exponential improvement on Minkowski’s bound (Torquato and Stillinger, 2006b), as discussed below.

The best currently known lower bound on $\phi_{\text{max}}^L$ for dimensions not divisible by four was obtained by Ball (1992). He found that

$$\phi_{\text{max}}^L \geq 2(d-1)\zeta(d)/2^d. \quad (34)$$

For dimensions divisible by four, Vance (2009) recently found the tightest lower bound on the maximal packing fraction among lattice packings,

$$\phi_{\text{max}}^L \geq 6d/2^d(1-e^{-d}). \quad (35)$$

Table IV gives the dominant asymptotic behavior of several lower bounds on $\phi_{\text{max}}^L$ for large $d$. Note that the best lower bounds on $\phi_{\text{max}}^L$ improve on Minkowski’s bound...
by a linear factor in $d$ rather than providing exponential improvement. This suggests that the packing fraction of the densest lattice packing in high $d$ is controlled by the exponential factor $2^{-d}$.

Nontrivial upper bounds on the maximal packing fraction $\phi_{\text{max}}$ for any sphere packing in $\mathbb{R}^d$ have been derived. Blichfeldt (1929) showed that the maximal packing fraction for all $d$ satisfies $\phi_{\text{max}} \leq (d/2 + 1)2^{-d/2}$. This upper bound was improved by Rogers (1958, 1964) by an analysis of the Voronoi cells. For large $d$, Rogers’ upper bound asymptotically becomes $d^{-2}e^{-d}$. Kabatiansky and Levenshtein (1978) found an even stronger bound, which in the limit $d \to \infty$ yields $\phi_{\text{max}} \leq 2^{-0.5990d}$. All of these upper bounds prove that the maximal packing fraction tends to zero in the limit $d \to \infty$. This rather counterintuitive high-dimensional property of sphere packings can be understood by recognizing that almost all of the volume of a $d$-dimensional sphere for large $d$ is concentrated near the sphere surface. For example, the volume contained with such a sphere up to 99% of it radius is $(99/100)^d$, which tends to zero exponentially fast. Thus, in high-dimensional sphere packings (densest or not), almost all of the volume is occupied by the void space (space exterior to the spheres), which is to be contrasted with the densest sphere packings in low dimensions in which the volume contained within the spheres dominates over the void volume.

Cohn and Elkies (2003) obtained and computed linear programming upper bounds, which provided improvement over Rogers’ upper bounds for dimensions 4–36, but it is not yet known whether they improve upon the Kabatiansky-Levenshtein upper bounds for large $d$. Cohn and Kumar (2009) used these techniques to prove that the Leech lattice is the unique densest lattice in $\mathbb{R}^{24}$. They also proved that no sphere packing in $\mathbb{R}^{24}$ can exceed the density of the Leech lattice by a factor of more than $1 + 1.65 \times 10^{-30}$ and gave a new proof that $E_8$ is the unique densest lattice in $\mathbb{R}^8$. Table V provides the dominant asymptotic behavior of several upper bounds on $\phi_{\text{max}}$ for large $d$. Note that the best upper and lower bounds on $\phi_{\text{max}}$ differ by an exponential factor as $d \to \infty$.

Since 1905, many extensions and generalizations of Minkowski’s bound have been derived (Conway and Sloane, 1998), but none of them have improved upon the dominant exponential term $2^{-d}$. Torquato and Stillinger (2006b) used a conjecture concerning the existence of disordered sphere packings and a $g_2$-invariant optimization procedure that maximizes $\phi$ associated with a radial “test” pair-correlation function $g_2(r)$ to provide the putative exponential improvement on Minkowski’s 100-year-old bound on $\phi_{\text{max}}$. Specifically, a $g_2$-invariant process (Torquato and Stillinger, 2002) is one in which the functional form of a test pair-correlation $g_2(r)$ function remains invariant as density varies, for all $r$, over the range of packing fractions.
The terminal packing fraction $\phi_*$ is the maximum achievable density for the $g_2$-invariant process subject to satisfaction of certain non-negativity conditions on pair correlations. For any test $g_2(r)$ that is a function of radial distance $r = |\mathbf{r}|$ associated with a packing, i.e., $g_2(r) = 0$ for $r < D$, they maximized the corresponding packing fraction,

$$\phi_* = \lim_{\max} \phi,$$

subject to satisfying the following two necessary conditions:

$$g_2(r) \geq 0 \quad \text{for all } r$$

and

$$S(k) \geq 0 \quad \text{for all } k.$$

Condition (39) is a necessary condition for the existence of any point process [cf. Eq. (7)]. When there exist sphere packings with a $g_2$ satisfying these conditions in the interval $[0, \phi_*]$, then one has the lower bound on the maximal packing fraction given by

$$\phi_{\max} \geq \phi_*.$$

Torquato and Stillinger (2006b) conjectured that a test function $g_2(r)$ is a pair-correlation function of a translationally invariant disordered sphere packing in $\mathbb{R}^d$ for $0 \leq \phi \leq \phi_*$ for sufficiently large $d$ if and only if conditions (38) and (39) are satisfied. There is mounting evidence to support this conjecture. First, they identified a decorrelation principle, which states that unconstrained correlations in disordered sphere packings vanish asymptotically in high dimensions and that the $g_n$ for any $n \geq 3$ can be inferred entirely (up to small errors) from a knowledge of $\rho$ and $g_2$. This decorrelation principle, among other results, provides justification for the conjecture of Torquato and Stillinger (2006b), and is exhibited by the exactly solvable ghost RSA packing process (Torquato and Stillinger, 2006a) as well as by computer simulations in high dimensions of the maximally random jammed state (Skoge et al., 2006) and the standard RSA packing (Torquato, 2006). Second, other necessary conditions on $g_2$ (Costin and Lebowitz, 2004; Torquato and Stillinger, 2006b; Hopkins et al., 2009) appear to only have relevance in low dimensions. Third, one can recover the form of known rigorous bounds [cf. Eqs. (32) and (34)] for specific test $g_2$’s when the conjecture is invoked. Finally, in these two instances, configurations of disordered sphere packings on the torus have been numerically constructed with such $g_2$ in low dimensions for densities up to the terminal packing fraction (Crawford et al., 2003; Uche et al., 2006).

Using a particular test pair correlation corresponding to a disordered sphere packing. Torquato and Stillinger (2006b) found a conjectural lower bound on $\phi_{\text{max}}$ that is controlled by $2^{-(0.778 \ldots) d}$ and the associated lower bound on the average contact (kissing) number $Z$ is controlled by $2^{0.22134 \ldots d}$ (a highly overconstrained situation). These results counterintuitively suggest that the densest packings as $d$ increases without bound may exhibit increasingly complex fundamental cells or even become disordered at some sufficiently large $d$ rather than periodic. The latter possibility would imply the existence of disordered classical ground states for some continuous potentials. Scardicchio et al. (2008) demonstrated that there is a wide class of test functions (corresponding to disordered packings) that lead to precisely the same putative exponential improvement on Minkowski’s lower bound and therefore the asymptotic form $2^{-(0.778 \ldots) d}$ is much more general and robust than previously surmised.

Interestingly, the optimization problem defined above is the dual of the infinite-dimensional linear program (LP) devised by Cohn (2002) to obtain upper bounds on the maximal packing fraction; see Cohn and Elkies (2003) for a proof. In particular, let $f(r)$ be a radial function in $\mathbb{R}^d$ such that

$$f(r) \leq 0 \quad \text{for } r \geq D,$$

$$\tilde{f}(k) \geq 0 \quad \text{for all } k,$$

where $\tilde{f}(k)$ is the Fourier transform of $f(r)$. Then the number density $\rho$ is bounded from above by

$$\min_{\rho} \frac{\tilde{f}(0)}{2^d\tilde{f}(0)}.$$

The radial function $f(r)$ can be physically interpreted to be a pair potential. The fact that its Fourier transform must be non-negative for all $k$ is a well-known stability condition for many-particle systems with pairwise interactions (Ruelle, 1999). We see that whereas the LP problem specified by Eqs. (38) and (39) utilizes information about pair correlations, its dual program [Eqs. (41) and (42)] uses information about pair interactions. As noted by Torquato and Stillinger (2006b) even if there does not exist a sphere packing with $g_2$ satisfying conditions (38) and (39) and the hard-core constraint on $g_2$, the terminal packing fraction $\phi_*$ can never exceed the Cohn-Elkies upper bound. Every LP has a dual program and when an optimal solution exists, there is no duality gap between the upper bound and lower bound formulations. Recently, Cohn and Kumar (2007a) proved that there is no duality gap.

XII. REMARKS ON PACKING PROBLEMS IN NON-EUCLIDEAN SPACES

Particle packing problems in non-Euclidean (curved) spaces have been the focus of research in a variety of fields, including physics (Bowick et al., 2006; Modes and Kamien, 2007), biology (Tammes, 1930; Goldberg, 1967; Prusinkiewicz and Lindenmayer, 1990; Torquato et al., 2002; Zandi et al., 2004), communications theory (Conway and Sloane, 1998), and geometry (Conway and Sloane, 1998; Hardin and Saff, 2004; Cohn and Kumar,
TABLE VI. The largest known kissing numbers for identical spheres in $R^d$ for selected $d$. Except for $R^3$ and $R^{10}$, the largest known kissing numbers listed are those found in the densest lattice packings listed in Table III. $P_{9a}$ is a nonlattice packing with an average kissing number of 235 $3/5$ but with a maximum kissing number of 306. $P_{10b}$ is a nonlattice packing with an average kissing number of 340 $1/3$ but with a maximum kissing number of 500. See Conway and Sloane (1998) for further details.

| Dimension, $d$ | Packing structure | Kissing number, $\tau$ |
|---------------|-------------------|-----------------------|
| 1             | $Z$               | 2                     |
| 2             | $A_2$             | 6                     |
| 3             | $D_3$             | 12                    |
| 4             | $D_4$             | 24                    |
| 5             | $D_5$             | 40                    |
| 6             | $E_6$             | 72                    |
| 7             | $E_7$             | 126                   |
| 8             | $E_8$             | 240                   |
| 9             | $P_{9a}$          | 306                   |
| 10            | $P_{10b}$         | 500                   |
| 16            | $\Lambda_{16}$   | 4320                  |
| 24            | $\Lambda_{24}$   | 196560                |

2007b). Although a comprehensive overview of this topic is beyond the scope of this review, we highlight here some of the developments in spaces with constant positive and negative curvatures. We limit the discussion to packing spheres on the positively curved unit sphere $S^{d-1} \subset R^d$ and in negatively curved hyperbolic space $H^d$.

The kissing (or contact) number $\tau$ is the number of spheres of unit radius that can simultaneously touch a unit sphere $S^{d-1}$ (Conway and Sloane, 1998). The kissing number problem asks for the maximal kissing number $\tau_{\text{max}}$ in $R^d$. The determination of the maximal kissing number in $R^3$ spurred a famous debate between Isaac Newton and David Gregory in 1694. The former correctly thought the answer was 12, but the latter wrongly believed that 13 unit spheres could simultaneously contact another unit sphere. The optimal kissing number $\tau_{\text{max}}$ in dimensions greater than three is only known for $R^4$ (Musin, 2008), $R^8$ and $R^{24}$ (Levenshtein, 1979; Odlyzko and Sloane, 1979). Table VI lists the largest known kissing numbers in selected dimensions.

In geometry and coding theory, a spherical code with parameters $(d, N, t)$ is a set of $N$ points on the unit sphere $S^{d-1}$ such that no two distinct points in that set have inner product greater than or equal to $t$, i.e., the angles between them are all at least $\cos^{-1} t$. The fundamental problem is to maximize $N$ for a given value of $t$, or equivalently to minimize $t$ given $N$ [sometimes called the Tammes problem, which was motivated by an application in botany (Tammes, 1930)]. One of the first rigorous studies of spherical codes was by Schütte and van der Waerden (1951). Delsarte et al. (1977) introduced much of the most important mathematical machinery to understand spherical codes and designs. One natural generalization of the best way to distribute points on $S^{d-1}$ (or $R^d$) is the energy minimization problem: Given some potential function depending on the pairwise distances between points, how should the points be arranged so as to minimize the total energy (or what are the ground-state configurations)? The original Thomson problem of “spherical crystallography” seeks the ground states of electron shells interacting via the Coulomb potential; but it is also profitable to study ground states of particles interacting with other potentials on $S^{d-1}$ (Bowers et al., 2006). Cohn and Kumar (2007b) introduced the beautiful idea of a universally optimal configuration, a unique configuration that minimizes a class of potentials. In particular, they proved that for any fixed number of points $N$ on $S^{d-1}$ there is a universally optimal configuration that minimizes all completely monotonic potential functions (e.g., all inverse power laws).

The optimal spherical code problem is related to the densest local packing (DLP) problem in $R^d$ (Hopkins et al., 2010a), which involves the placement of $N$ nonoverlapping spheres of unit diameter near an additional fixed unit-diameter sphere such that the greatest radius $R$ from the center of the fixed sphere to the centers of any of the $N$ surrounding spheres is minimized. We recast the optimal spherical code problem as the placement of the centers of $N$ nonoverlapping spheres of unit diameter onto the surface of a sphere of radius $R$ such that $R$ is minimized. It is has been proved that for any $d$, all solutions for $R$ between unity and the golden ratio $\tau = (1 + \sqrt{5})/2$ to the optimal spherical code problem for $N$ spheres are also solutions to the corresponding DLP problem (Hopkins et al., 2010b). It follows that for any packing of nonoverlapping spheres of unit diameter, a spherical region of radius $R$ less than or equal to $\tau$ centered on an arbitrary sphere center cannot enclose a number of sphere centers greater than one more than the number that than can be placed on the spherical region's surface.

We saw in Sec. VIII that monodisperse circle (circular disk) packings in $R^2$ have a great tendency to crystallize at high densities due to a lack of geometrical frustration. The hyperbolic plane $H^2$ (for a particular constant negative curvature, which measures the deviation from the flat Euclidean plane) provides a two-dimensional space in which global crystalline order in dense circle packings is frustrated, and thus affords a means to use circle packings to understand fundamental features of simple liquids, disordered jammed states, and glasses. Modes and Kamien (2007) formulated an expression for the equation of state for disordered hard disks in $H^2$ and compared it to corresponding results obtained from molecular-dynamics simulations. Modes and Kamien (2007) derived a generalization of the virial equation in $H^2$ relating the pressure to the pair-correlation function and developed the appropriate setting for extending integral-equation approaches of liquid-state theory. For a discussion of the mathematical subtleties associated with finding the densest packings of identical


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d-dimensional spheres in \( \mathbb{H}^d \), see Bowen and Radin (2003).

XIII. CHALLENGES AND OPEN QUESTIONS

The geometric-structure approach advanced and explored in this review provides a comprehensive methodology to analyze and compare jammed disk and sphere packings across their infinitely rich variety. This approach also highlights aspects of present ignorance, thus generating many challenges and open questions for future investigation. Even for identical spheres, detailed characterization of jammed structures across the simple two-dimensional (2D) order maps outlined in Sec. V is still incomplete. A partial list of open and challenging questions in the case of sphere packings includes the following:

1. Are the strictly jammed “tunneled” crystals (Torquato and Stillinger, 2007) the family of lowest density collectively jammed packings under periodic boundary conditions?

2. How can the extremal jammed packings that inhabit the upper and lower boundaries of occupied regions of each of those order maps be unambiguously identified?

3. What would be the shapes of analogous occupied regions if the two-parameter versions shown in Fig. 13 were to be generalized to three or more parameters?

4. To what extent can the rattler concentration in collectively or strictly sphere packings be treated as an independent variable? What is the upper limit to attainable rattler concentrations under periodic boundary conditions?

5. What relations can be established between order metrics and geometry of the corresponding configurational-space polytopes?

6. Can upper and lower bounds be established for the number of collectively and/or strictly jammed states for \( N \) spheres?

7. Upon extending the geometric-structure approach to Euclidean dimensions greater than three, do crystalline arrangements with arbitrarily large unit cells or even disordered jammed packings ever provide the highest attainable densities?

Jamming characteristics of nonspherical and even nonconvex hard particles is an area of research that is still largely undeveloped and therefore deserves research attention. This is true even for simple particle shapes, such as spheres. For example, we know little about the jamming categories of spheres in curved spaces. Does curvature facilitate jamming or not? Does it induce ordering or disordering? These are just a few of the many challenging issues that merit further investigation.

In view of the wide interest in packing problems that the research community displays there is reason to be optimistic that substantial conceptual advances are forthcoming. It will be fascinating to see how future articles covering this subject document those advances.

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