

Supplemental Material

Phase Diagram and Structural Diversity of a Family of Truncated Cubes: Degenerate Close-Packed Structures and Vacancy-Rich States

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I. DEFINITION OF THE FAMILY OF TRUNCATED CUBES

We define our family of truncated cubes using a simple mathematical expression for the location of the vertices. The line segments that connect these vertices can only be assigned in one (unique) way to obtain a truncated cube. The vertices of a truncated cube may be written as a function of the shape parameter $s \in [0, 1]$:

$$\{\mathbf{v}(s)\} = \begin{cases} \left(1 - \frac{4}{3}s^3\right)^{(-1/3)} \mathcal{P}_D \left(\pm \left(\frac{1}{2} - s\right), \pm \frac{1}{2}, \pm \frac{1}{2}\right)^T & s \in [0, \frac{1}{2}] \\ \left(\frac{4}{3} - 4\lambda^3\right)^{(-1/3)} \mathcal{P}_D \left(\pm(1 - \lambda), \pm\lambda, 0\right)^T & \lambda \equiv 1 - s \in [0, \frac{1}{2}] \end{cases}, \quad (1)$$

where $\mathcal{P}_D$ is a permutation operation that generates all permutations of each element in the sets of 8 and 4 vertices spanned by the $\pm$ -operations, respectively. All duplicate vertices are removed after letting $\mathcal{P}_D$ act. The ‘ T ’ indicates transposition. The prefactors ensure that the truncated cubes are normalized to unit volume. As mentioned in the main text, several Platonic and Archimedean solids are members of this family: $s = 0$ a cube, $s = (2 - \sqrt{2})/2 \approx 0.292893$ a truncated cube, $s = 1/2$ a cuboctahedron, $s = 2/3$ a truncated octahedron, and $s = 1$ an octahedron; these are depicted in Fig. 1a.

II. METHODS

A. Determining the Close-Packed Structures

The simulations by which the close-packed structures were derived, are based on the floppy-box Monte Carlo (FBMC) method [1, 2] in combination with the separating-axis-based overlap algorithm [3]. We obtained

the densest crystal structure and the corresponding packing fraction ϕ as a function of the level of particle truncation s by considering 1,000 equidistant points in $s \in [0, 1]$. For each point we prepared systems of truncated particles in a dilute phase, typically with packing fraction $\phi \approx 0.001$. We increased the reduced pressure in 100 steps according to a geometric series from $p = 1$ to $p \approx 10^5$ over $4 \cdot 10^6$ Monte Carlo (MC) cycles in order to compress these systems to a high-density crystalline state. This pressure increase was typically applied a total of 1,000 times for $N = 1$ particles in the unit cell and for each shape. We restricted ourselves to $N = 1$ particles in the unit cell, because the truncated cubes are all centrosymmetric. We only considered $N = 2, \dots, 6$ for 14 conveniently chosen values of s , located in the center of the regions indicated in Fig. 1, as will be justified shortly. For these $N > 1$ systems we obtained roughly the same value of ϕ and also the same crystal structures. The densest crystal-structure candidate was selected and allowed to compress further for another 10^6 MC cycles at $p = 10^6$ to achieve 5 decimals of precision in ϕ . In practice, these final cycles of compression did not improve the packing fraction substantially. Figure 1b shows ϕ as a function of s . Note that the packing fraction ‘curve’ is continuous, but has discontinuities in its first derivative. To double check our result, we considered another set of FBMC runs. We used several of the 1,000 densely-packed crystals as our initial configuration and we varied s around these points at high pressure to study the evolution of their structure. Steps of 10^{-5} in s were used and for each step the system is expanded to remove any overlaps, before re-compressing it at $p \approx 10^5$. The packing fractions we obtained showed good correspondence with our original result, but this correspondence failed for a transition between two crystal structures. The consecutive method would often become stuck in the lower density structure that corresponded to the morphology of the crystal phase it came from.

The unit cell for $N = 1$ truncated cubes can be specified by three vectors $\mathbf{v}_i$ ($i = 1, 2, 3$) that are implicitly s dependent. The structure spanned by these three vectors can also be described by the length $v_i = |\mathbf{v}_i|$ of the vectors and the angles θ_{ij} ($i < j = 1, 2, 3$) between them. Note that we ignored the orientation of the particle with respect to the unit cell here. In order to give an unbiased comparison of the different vectors we used lattice

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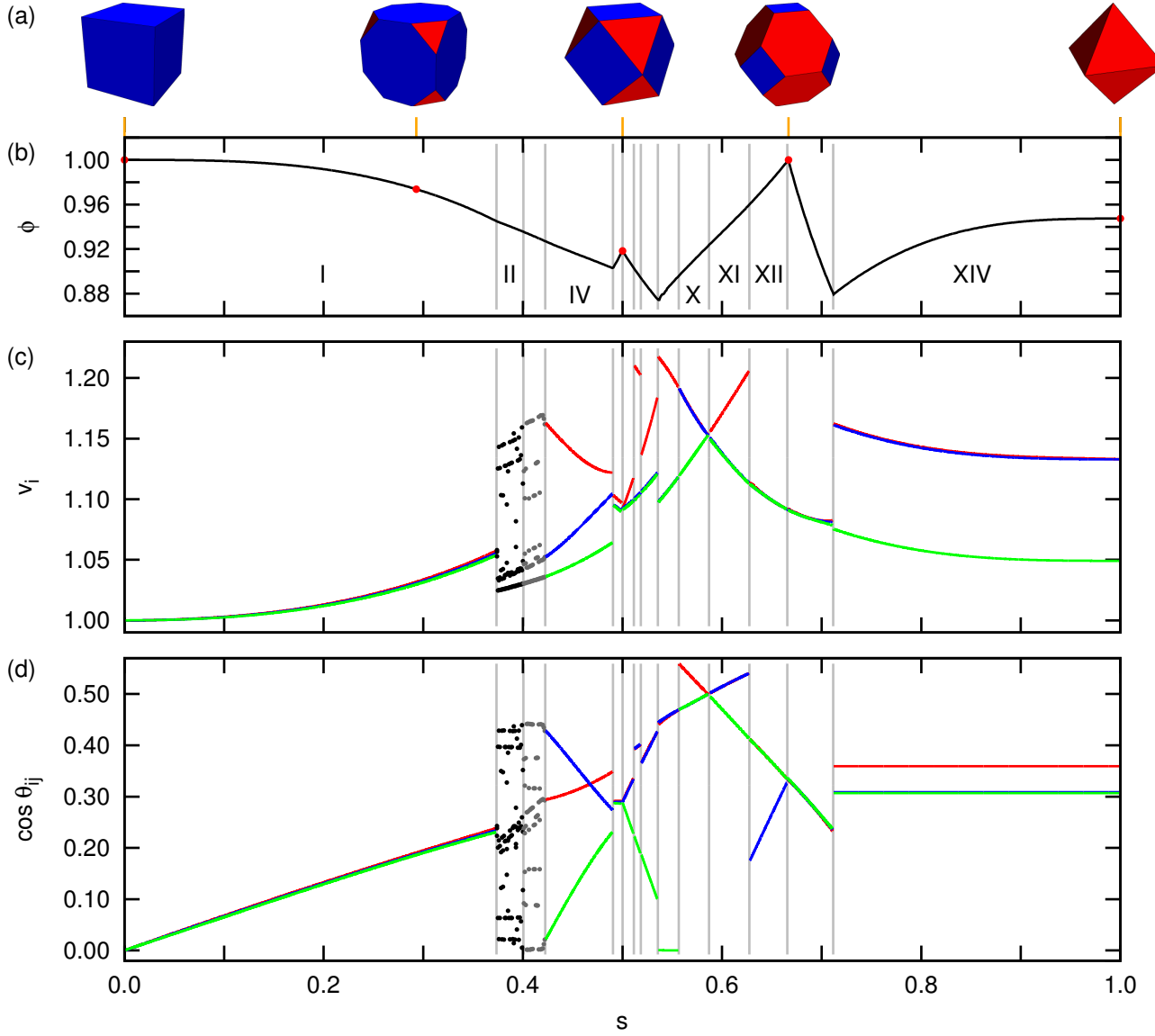


Figure 1: (a) Five examples of truncated cubes (Platonic and Archimedean solids only) for levels of truncation s corresponding to the orange lines: $s = 1$ a cube, $s = (2 - \sqrt{2})/2 \approx 0.293$ a truncated cube, $s = 1/2$ a cuboctahedron, $s = 2/3$ a truncated octahedron, and $s = 1$ an octahedron. (b) The packing fraction ϕ for the close-packed structures as a function of s . The values for the five solids shown in (a) are given by red dots. (c) The length v_i ($i = 1, 2$, and 3) of the three lattice vectors, indicated in red, green, and blue, that span the unit cell of the densest crystal structure as a function of s . Not every line is clearly visible, since there is some overlap. In the region where the black and gray dots are used ($s \in [0.37, 0.40]$ and $s \in [0.40, 0.42]$), there appears to be a degeneracy in the crystal structures, as is explained in the text. (d) The cosine of the angles θ_{ij} ($i < j = 1, 2$, and 3) between the three vectors that span the unit cell as a function of s . Gray vertical lines partition the s -domain into 14 pieces with a ‘different’ crystal structure, based on the discontinuities shown in the v_i and $\cos \theta_{ij}$ results. These regions are numbered with roman numerals in (b); only those regions large enough to accommodate a label are numbered, but the numbering can be continued from left to right, in the unnumbered regions.

reduction [4] to ensure that for each unit cell the surface to volume ratio is minimal. These numerical results are shown in Fig. 1(c,d). By analyzing the v_i and θ_{ij} , as well as the location of the kinks in the ϕ -curve, we were able to partition the $s \in [0, 1]$ domain into 14 distinct regions. This is the reason behind our choice of 14 verification points for $N > 1$ simulations. For a discussion

of the crystal structures in the different regions, we refer the reader to Sec. IV.

B. Equations of State and Mesophase Structures

We used the close-packed crystal structures obtained from the FBMC calculations as initial configurations for variable-box-shape isothermal-isobaric (NPT) Monte Carlo simulations, to study the phase behavior at intermediate pressures. Initial configurations of 300 to 600 particles were prepared and melted to determine the equations of state (EOSs) for the various phases. Typical equilibration and production times were around $1.2 \cdot 10^6$ and $2 \cdot 10^6$ Monte Carlo Sweeps (MCS), respectively. One MCS is defined as N Monte Carlo trial moves (translation, rotation, volume change, or deformation of the box, respectively), where N is the number of particles in the system. We sampled the lattice vectors, as well as the average positions and orientations of the particles as a function of packing fraction and truncation parameter s . The sampling was done on an interval of 100 MCS to avoid correlated configurations. Using these results we set up regular NPT simulations (possibly with a triclinic box shape) to more accurately sample the EOSs for all phases with larger system sizes of 1,000 to 2,000 particles, including the liquid phase. In addition, we employed these results to set up the Einstein crystal [5] for the free-energy calculations.

Several typical configurations of crystalline phases slightly above fluid coexistence are displayed in Fig. 2 to give an impression of the mesophases that occur in the systems we studied. We have also added a movie file to the supplemental information ‘pbct.mpg’, which shows the plastic-body-centered tetragonal (pbct) phase for truncation $s = 0.546$ and pressure $p^* = 11.0$, to illustrate that a bct structure can be present for particles that rotate. On the top-left corner the entire simulation box is shown for an NPT simulation containing 512 particles. An enlarged pbct unit cell is shown from the front (top-right corner), from the bottom (bottom-left corner), and from the side (bottom-right corner); with all the neighboring particles hidden. The change in the color of the particles is related to the orientation of the particles, the more the particle is aligned with the lattice vectors of the Einstein crystal, the more bright green the color is; the more the particle is misaligned, the more it tends to a blue color.

C. Free-Energy Calculations and Confining Potentials

We obtained the dimensionless free energy per particle $f = \beta F/N$ as a function of packing fraction ϕ (or equivalently the density ρ) for the entire density range by thermodynamic integration [6] over the EOS, from reference density ρ_0 to the density of interest ρ :

$$f(\rho) = f(\rho_0) + \int_{\rho_0}^{\rho} \frac{\beta P(\rho')}{\rho'^2} d\rho'. \quad (2)$$

Here $f(\rho_0) \equiv \beta\mu(\rho_0) - \beta P(\rho_0)/\rho_0$ is the reduced Helmholtz free energy per particle at density ρ_0 , with $\beta = 1/k_B T$, T the temperature and k_B Boltzmann’s constant, $\mu(\rho_0)$ the chemical potential, and $P(\rho_0)$ the pressure. The Helmholtz free energy at reference density ρ_0 was obtained as follows.

- In the fluid phase we used Widom’s particle insertion method [7] to obtain the free energy. This method was employed at relatively low densities to obtain small error bars. We performed the calculations at $\phi \approx 0.2$. We note that there were no finite size effects within the computational accuracy for the particle insertion method.
- In the crystal phase we used the Einstein integration method [6, 8, 9]. The reduced Helmholtz free energy per particle $f = \beta F/N$ of a crystal is given by:

$$f(\rho) = f_{\text{Einst}}(\lambda_{\text{max}}) - \frac{1}{N} \int_0^{\lambda_{\text{max}}} d\lambda \left\langle \frac{\partial \beta U_{\text{Einst}}(\lambda)}{\partial \lambda} \right\rangle, \quad (3)$$

where f_{Einst} denotes the reduced free energy per particle of the ideal Einstein crystal, which is given by:

$$\begin{aligned} f_{\text{Einst}}(\lambda_{\text{max}}) = & -\frac{3(N-1)}{2N} \log \left(\frac{\pi}{\lambda_{\text{max}}} \right) + \\ & \log \left(\frac{\Lambda_i^3 \Lambda_r}{v_p} \right) + \frac{1}{N} \log \left(\frac{v_p}{V N^{1/2}} \right) - \\ & \frac{1}{N} \log \left\{ \frac{1}{8\pi^2} \int d\theta \sin(\theta) d\phi d\chi \times \right. \\ & \left. \exp \left[-\frac{\lambda_{\text{max}}}{k_B T} (\sin^2 \psi_{ia} + \sin^2 \psi_{ib}) \right] \right\}, \end{aligned}$$

with v_p the volume of a particle. $U_{\text{Einst}}(\lambda)$ denotes the harmonic potential that fixes the particles to the respective Einstein lattice positions:

$$\beta U_{\text{Einst}}(\lambda) = \lambda \sum_{i=1}^N [(\mathbf{r}_i - \mathbf{r}_{i,0})^2 / v_p^{2/3} + (\sin^2 \psi_{ia} + \sin^2 \psi_{ib})], \quad (4)$$

with $(\mathbf{r}_i - \mathbf{r}_{i,0})$ the displacement of particle i from its position in the ideal Einstein crystal. The angles ψ_{ia} and ψ_{ib} are the minimum angles between vectors, $\mathbf{a}$ and $\mathbf{b}$, describing the orientations of the particles in the ideal Einstein crystal and the equivalent vectors that describe the orientation of the particle in the actual crystal, respectively. When λ is large the particles translational and orientational displacements are frozen, while at lower λ ’s the particles freely displace and rotate. They thereby not only explore the possible states, but also the underlying degeneracy coming from the symmetry of

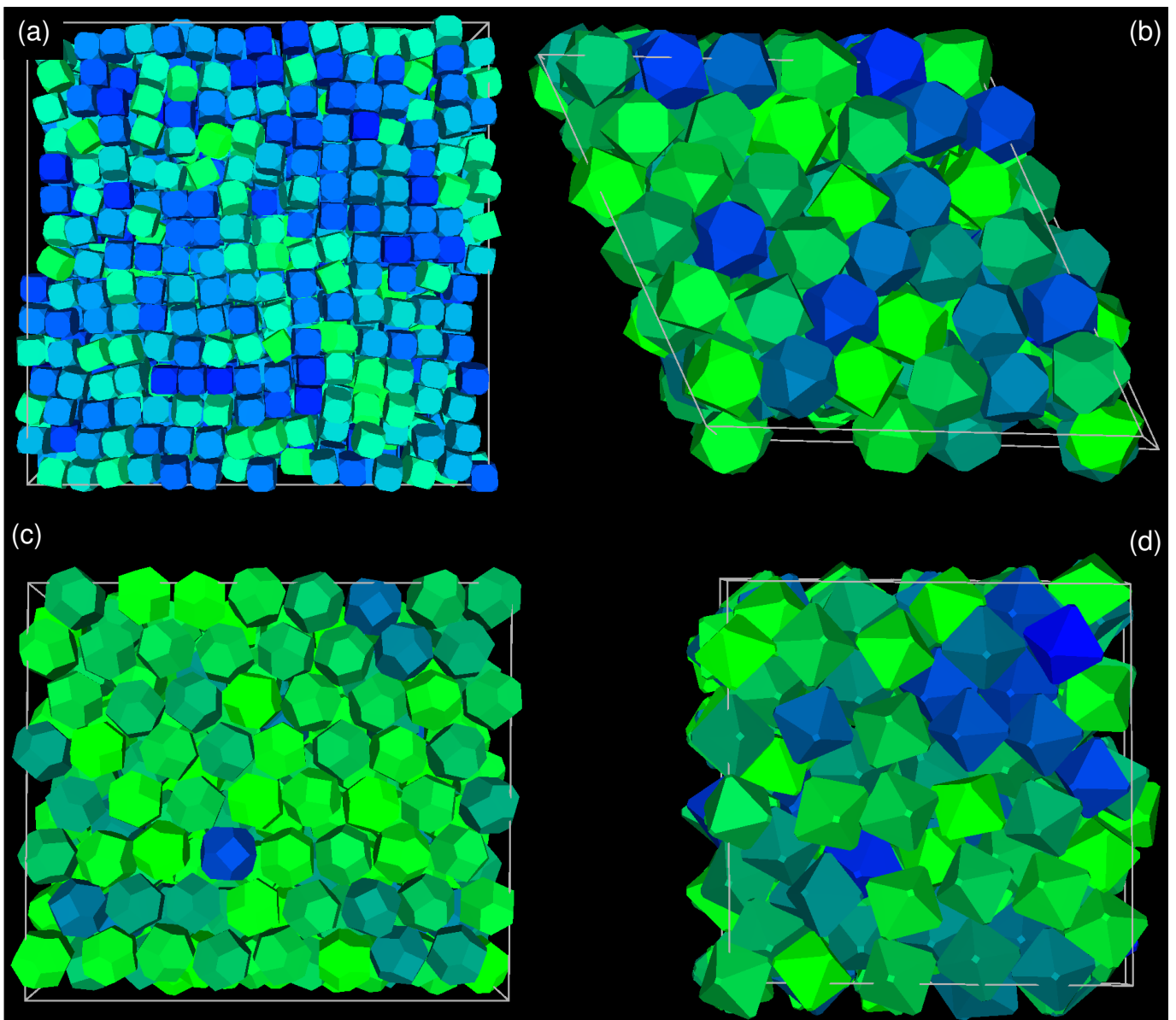


Figure 2: Several snapshots of our isothermal-isobaric (NPT) simulations showing the various crystal structures that form in our family of truncated cubes. (a) Equilibrium vacancy-rich simple cubic crystal for a truncation of $s = 0.25$ at packing fraction $\phi = 0.56$. The simulation was performed for $N = 3,235$ particles. In this system the vacancy concentration was found to be $\alpha = 0.032$. (b) Plastic hexagonal-close-packed (PHCP) phase for $s = 0.411$ and $\phi = 0.6$ in a box containing $N = 216$ particles. (c) Plastic body-centered-tetragonal (pbct) phase for $s = 0.607$ and $\phi = 0.58$ in a box containing $N = 512$ particles. (d) Plastic body-centered-cubic (PBCC) phase for $s = 0.900$ and $\phi = 0.52$ in a box containing $N = 250$ particles. The coloring used here indicates the level of alignment of these particles with the lattice vectors of the Einstein crystal, as explained in the text.

the particle itself. The translational and rotational thermal wavelengths Λ_t and Λ_r are set to 1 in our calculations. We mostly used the same system sizes of 700 to 1,500 particles to compute the free energies for the (plastic) crystal phases. We found that finite-size scaling (FSS) was only necessary in the octahedron regime, *i.e.*, $s \approx 1$, to establish the phase diagram. For such high levels of truncation the free-energy differences between the various

phases at coexistence proved to be very small, see Ref. [5]. For the other phase transitions the free energies obtained without FSS proved to be sufficient to accurately determine the phase boundaries.

- For the free-energy calculations of a plastic-crystal (rotator) phase, we followed the approach of the authors of Ref. [10], who introduced a method, which allows for a continuous transition from a non-

interacting plastic-crystal to an interacting plastic-crystal phase of hard truncated cubes. We used a tunable soft-to-hard interaction potential between the particles

$$\varphi(i, j) = \begin{cases} \gamma[1 - A(1 + \zeta(i, j))] & \text{if } \zeta(i, j) < 0 \\ 0 & \text{otherwise} \end{cases}, \quad (5)$$

where $\zeta(i, j)$ is the overlap potential defined in Ref. [11], which is negative when two particles i and j overlap and positive otherwise, and γ is the integration parameter, which runs from $\gamma = 0$ (non-interacting) to $\gamma = \gamma_{\max}$, for which the system interacts fully. In our calculations we used $A = 0.9$ following Ref. [12]. The dimensionless Helmholtz free energy per particle $f = \beta F/N$ of the plastic crystal is given by:

$$f(\rho) = f_{\text{Einst}}(\lambda_{\max}) - \frac{1}{N} \int_0^{\lambda_{\max}} d\lambda \left\langle \frac{\partial \beta U_{\text{Einst}}(\lambda)}{\partial \lambda} \right\rangle_{\gamma_{\max}} + \frac{1}{N} \int_0^{\gamma_{\max}} d\gamma \left\langle \frac{\partial \sum_{i \neq j}^N \beta \varphi(i, j)}{\partial \gamma} \right\rangle_{\lambda_{\max}}.$$

- We calculated the free energy of a solid with a fixed vacancy concentration $\alpha = (N_L - N)/N_L$, with N the number of particles and N_L the number of lattice sites, according to the procedures outlined in Ref. [13]. The dimensionless free energy per particle $f_{\text{Einst}}^{\text{vac}}$ of a non-interacting Einstein crystal with vacancies is given by:

$$f_{\text{Einst}}^{\text{vac}}(\lambda_{\max}) = f_{\text{Einst}}(\lambda_{\max}) + f_{\text{comb}}, \quad (6)$$

where f_{comb} is the combinatorial entropy associated with placing N particles on N_L lattice sites:

$$f_{\text{comb}} = -\frac{1}{N} \log \left(\frac{N_L!}{N!(N_L - N)!} \right). \quad (7)$$

For more details about the implementation of the free-energy calculations, we refer to the methods section of Ref. [13]. We used $N_L = 15^3 = 3,375$ for the free-energy calculation at $\phi = 0.56$.

III. FREE ENERGY OF VACANCIES

We calculated the free energies for a simple cubic (sc) crystal phase of truncated hard cubes with a shape parameter $s = 0.0, 0.05, 0.15$, and 0.25 and packing fraction $\phi = 0.56$ as a function of the vacancy concentration $\alpha = (N_L - N)/N_L$ as described in Sec. II C. In Fig. 3, we present the dimensionless free energy per particle f_{def} as a function of α without taking into account the combinatorial entropy, *i.e.*, $f_{\text{def}}(\alpha) = f(\alpha) - f_{\text{comb}}(\alpha)$. In accordance with the observations of Ref. [13], f_{def} vs.

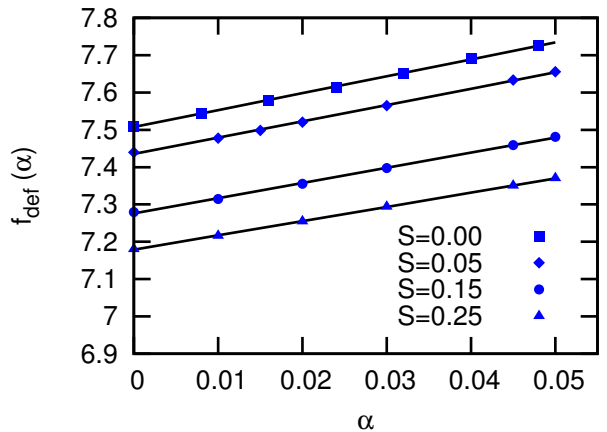


Figure 3: The reduced free energy per particle $f_{\text{def}}(\alpha)$ as a function of vacancy concentration α for a simple cubic crystal phase of truncated cubes with shape parameter $s = 0, 0.05, 0.15$, and 0.25 and packing fraction $\phi = 0.56$. $f_{\text{def}}(\alpha)$ does not account for the combinatorial entropy, *i.e.*, $f_{\text{def}}(\alpha) = f(\alpha) - f_{\text{comb}}(\alpha)$.

Table I: The dimensionless free energy per particle for a simple cubic crystal of truncated cubes with shape parameter $s = 0, 0.05, 0.15$, and 0.25 and packing fraction $\phi = 0.56$. From left to right we specify the truncation parameter s , the free energy for a system with zero vacancies $f(0)$ and the free energy cost to create a single vacancy f_1 (without the combinatorial contribution) as obtained by Monte Carlo simulations.

s	$f(0)$	f_1
0.00	7.5077	4.52640
0.05	7.4354	4.36892
0.15	7.2762	4.06024
0.25	7.1788	3.81633

α in Fig. 3 shows a linear behavior suggesting that the vacancies are only weakly interacting. Assuming the vacancies to be non-interacting, as explained in the main text, one can write the free energy of a crystal phase as

$$f_{\text{def}}(\alpha) = f(0) + \alpha f_1 \quad (8)$$

where $f(0)$ is the dimensionless free energy per particle of a crystal with no vacancies and $f_1 > 0$ is the reduced free energy cost to create a single vacancy (without the combinatorial component). Fitting the results as displayed in Fig. 3 with Eq. 8, we determined f_1 for truncated cubes as a function of shape parameter s . The results are tabulated in Table I. Surprisingly, we found that the free energy cost to create a vacancy f_1 decreases with increasing s at packing fraction $\phi = 0.56$. Hence, the equilibrium vacancy concentration increases with particle truncation s , also see the main text.

In the phase diagram of truncated cubes, shown in Fig. 2 of the manuscript, we did not take into account the shift in the coexistence densities effected by the high vacancy concentrations in the sc phase. Smallegange *et al.* have

shown that the coexistence densities for the phase diagram change by only a few percent if the vacancies are taken into account for the high-vacancy phases [13]. We therefore feel justified in this omission, since the influence in the vacancies leads to a minute change in the location of the phase boundaries for the sc-fluid coexistence densities. There is no indication that the other phases suffered from such high vacancy concentrations and any phase boundaries determined for these coexistences can therefore be taken as precise within the numerical uncertainty.

IV. DENSE PACKINGS AND CRYSTAL STRUCTURES

In this section we discuss the properties of the close-packed structures that are obtained using the methods described in Sec. II A. Figure 5 shows the crystal structure in the center of each of the 14 regions that we found in Fig. 1. There is a strong difference between the domains $s < 1/2$ and $s > 1/2$. Geometrically the cuboctahedron ($s = 1/2$) is the transition point between shapes which have a more cube-like nature and shapes which have a more octahedron-like nature. It is therefore not surprising that the crystal structures in the two regions ($s < 1/2$ and $s > 1/2$) appear to have a deformed simple cubic (dsc) symmetry and a deformed body-centered tetragonal (dbct) symmetry, respectively. We illustrate this in Fig. 5 where we show the most orthorhombic unit cell: $N = 1$ for $s < 1/2$ and $N = 2$ for $s > 1/2$. A remarkable result is the stability of the Minkowski crystal [14], which is the densest-packed Bravais-lattice structure for octahedra [15], under variations in s . For all $s \in [0.71, 1]$, we find a Minkowski crystal in the dense-packed limit, which can be inferred from the horizontal $\cos \theta_{ij}$ lines in Fig. 1d. The scaled length of the vectors $v_i \phi^{-1/3}$ is also constant on this domain.

Let us now examine the crystal structures in the 14 regions identified by the discontinuities in the vectors of the unit cell. In literature it has become commonplace to assign atomic equivalents to structures observed in simulations or experiments. For example, this is done for binary mixtures of spheres [1, 16], a family of truncated tetrahedra [17], several faceted particles [18], and systems of nanoparticles [19, 20]. We attempted to follow suit by determining the symmetry group of the structures in Fig. 5 using *FindSym* [21] and by subsequently assigning an atomic equivalent [22]. However, we found that a description in terms of atomic equivalents inadequately captures the richness in crystal structure, since particle orientation is not taken into account. Moreover, for many of our structures we are unable to determine a nontrivial space group using *FindSym*. We therefore resorted to visual analysis and we used this to group the 14 regions in Fig. 1 based on similarities between the respective structures.

- **I** In this region ($s \in [0.00, 0.37]$) we obtained a con-

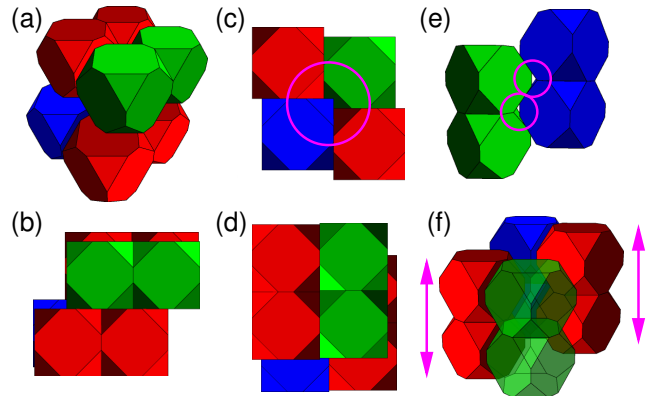


Figure 4: A visual representation of the degenerate crystal structure; we consider the value $s = 0.387$ in this figure. Pairs of truncated cubes, for which the octahedral faces are aligned (columns), are color-coded. Different viewpoints are displayed for a piece of crystal consisting of 8 particles: (a) Bird's eye view, (b) side view, (c), front view, and (d) top view of this structure. In (c) we use a magenta circle to indicate that the blue column is interlocking with the green column in a diagonal way. (e) A diagonal view of the crystal structure, where the red columns have been removed. Magenta circles show the interlocking. (f) The two red columns are *not* interlocking with the blue and green column, allowing for freedom of motion in the direction of the magenta arrows. The green column is made translucent to better illustrate the properties of this crystal structure.

tinuous and uniform distortion of the simple cubic structure for cubes. For $s = 0$ the particles form a sc crystal, which has the same morphology as α Po (α -Polonium) [22]. The uniformly deformed simple cubic (udsc) structure we found for $s > 0$ is similar to that of β Po [22]. We verified this deformed quality for values as low as $s = 10^{-5}$.

- **II & III** For these two regions ($s \in [0.37, 0.40]$ and $s \in [0.40, 0.42]$) we found that there is a degeneracy in the crystal-structure candidates that achieve the densest-known packing. Although certain structures appear favored over others, there is no clear relation between the structure and s . However, the packing fraction ϕ of the close-packed crystals is continuous in these regions. The observed degeneracy can be explained by the formation of sheets consisting of diagonally-interlocked columns, which can slide up or down (in the direction of the columns) with respect to each other, as shown in Fig. 4. For $s = 0.387$, the truncated cubes are arranged in a dsc crystal lattice, where the particles form columns that are interlocked in a diagonal way. These structures are referred to as mono-interlocking distorted simple cubic (MI dsc) crystals. This diagonal interlocking together with the close-packing condition prevents lateral motion in the plane normal to the column's direc-

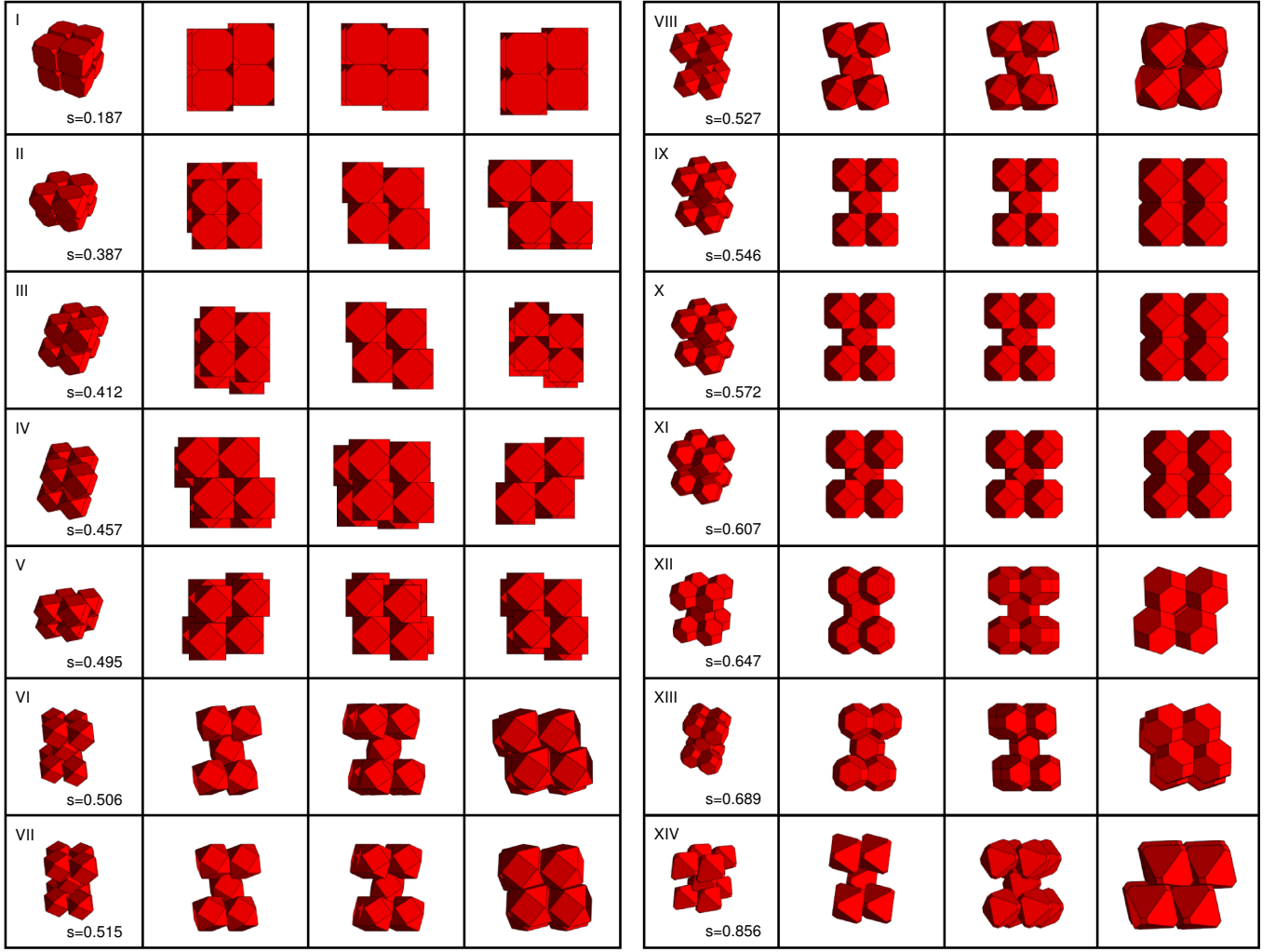


Figure 5: Visual representations of the crystal structures obtained for the first 7 regions (left) and the last 7 regions (right) of Fig. 1. From left to right each entry (row) contains a bird's eye view, the front view, the side view, and the top view of this structure. The Roman numeral in the top-left corner gives the relevant domain in Fig. 1. The truncation parameter s for these structures is given in the bottom-right corner.

tion. However, since the system is not fully interlocked, motion in the direction of the columns is possible for the diagonally interlocked sheets. We refer to these structures as mono-interlocking distorted simple cubic (MI dsc) crystals. This degeneracy is different from the degeneracy that occurs in structures consisting of cubes or hexagonal prisms for instance, since such systems allow lateral freedom of movement of columns or (perpendicular to the columns) of sheets of aligned particles. That is, there is possible freedom of motion in three directions, albeit not necessarily at the same time. The interlocking nature of the MI dsc phase only allows for movement in one direction only, namely parallel to the columns, which may lead to strong rheological differences between this structure and, *e.g.*, the sc structure for cubes.

- **IV** For this region ($s \in [0.42, 0.49]$) we find a dsc phase that is interlocking in two directions: a bi-interlocking dsc (BI dsc) phase. For each instance of interlocking two degrees of translational motion are frozen out. This implies that the BI dsc structure is completely fixed, which is confirmed by the unicity of the v_i and θ_{ij} results in Fig. 1(c,d).
- **V** In this region ($s \in [0.49, 0.50]$) we observed a tri-interlocking dsc (TI dsc) phase.
- **VI - VIII** Here ($s \in [0.50, 0.51]$, $[0.51, 0.52]$, and $[0.52, 0.54]$) we found structures that are best described by a distorted body-centered tetragonal (dbct) structure. The truncated cubes in these crystals are not aligned with the axes of the unit cell. It is unclear to what extent structures in regions VI, VII, and VIII are the same. The smooth

flow of the ϕ -curve (Fig. 1b), as well as their appearance, s implies continuity, but the jumps in the values of v_i and θ_{ij} [Fig. 1(c,d)] suggest otherwise.

- **IX - XI** These structures ($s \in [0.54, 0.56]$, $[0.56, 0.59]$, and $[0.59, 0.63]$) have a body-centered tetragonal (bct) morphology, for which the particles are aligned with the lattice vectors of the unit cell. It is surprising that the structures in region XII exhibit a dbct morphology, since regions IX - XII share the same smooth piece of ϕ -curve, see Fig. 1b. This leads us to conclude that a smooth dependence of ϕ on s is *not* indicative of uniformity in crystal structure. The strong similarity between the crystal structures in regions IX - XI and the apparent smooth transition between structures from region IX to X and from X to XI, also leads us to conclude that discontinuities in the properties of unit cell are not indicative of discontinuities in the properties of the crystal structure.
- **XII & XIII** These two dbct structures ($s \in$

$[0.63, 0.67]$ and $s \in [0.67, 0.71]$) are different from the dbct structures in regions VI - VIII, since the particles appear to be aligned with the lattice vectors of the unit cell. Moreover, crystals in region XIII are unusual, since there are large ‘voids’ in the structure. That is, for all other structures we found that the largest facets of a particle are always in contact with a facet of another particle. This is not the case in region XIII, because there is a substantial gap between some of the hexagonal facets.

- **XIV** The Minkowski crystal of region XIV ($s \in [0.71, 1.00]$) is also peculiar. It is the only structure which does not undergo some form of reorganization upon varying the level of truncation. It is worthwhile to study the origin of this apparent stability, which sharply contrasts with the immediate distortion found around $s = 0$. However, this goes beyond the scope of the current investigation.

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