



Showcasing collaborative research led by Dr Sarah Guerin, from the modelling theme of the Science Foundation Ireland Research Centre for Pharmaceuticals, SSPC.

Density functional theory predictions of the mechanical properties of crystalline materials

Computational chemistry, specifically Density Functional Theory, is an invaluable tool in predicting, understanding, and engineering the mechanical properties of crystalline materials. Extensive benchmarking of functionals and dispersion corrections leads to high predictability versus experiments in pharma and materials science research.

As featured in:



See Sarah Guerin *et al.*,
CrystEngComm, 2021, **23**, 5697.



Cite this: *CrystEngComm*, 2021, 23, 5697

Density functional theory predictions of the mechanical properties of crystalline materials

Evan Kiely,^a Reabetswe Zwane,^{bc} Robert Fox,^{bc}
Anthony M. Reilly ^{bc} and Sarah Guerin ^{*ac}

The mechanical properties of crystalline materials are crucial knowledge for their screening, design, and exploitation. Density functional theory (DFT), remains one of the most effective computational tools for quantitatively predicting and rationalising the mechanical response of these materials. DFT predictions have been shown to quantitatively correlate to a number of experimental techniques, such as nanoindentation, high-pressure X-ray crystallography, impedance spectroscopy, and spectroscopic ellipsometry. Not only can bulk mechanical properties be derived from DFT calculations, this computational methodology allows for a full understanding of the elastic anisotropy in complex crystalline systems. Here we introduce the concepts behind DFT, and highlight a number of case studies and methodologies for predicting the elastic constants of materials that span ice, biomolecular crystals, polymer crystals, and metal-organic frameworks (MOFs). Key parameters that should be considered for theorists are discussed, including exchange-correlation functionals and dispersion corrections. The broad range of software packages and post-analysis tools are also brought to the attention of current and future DFT users. It is envisioned that the accuracy of DFT predictions of elastic constants will continue to improve with advances in high-performance computing power, as well as the incorporation of many-body interactions with quasi-harmonic approximations to overcome the negative effects of calculations carried out at absolute zero.

Received 4th April 2021,
Accepted 22nd May 2021

DOI: 10.1039/d1ce00453k

rsc.li/crystengcomm

Introduction

Knowing the mechanical properties of a material is a necessary requirement for the vast majority of materials scientists and engineers, from composites in the aeronautical and automotive industries,^{1–3} to pharmaceutical crystals^{4–6} and biomaterials.^{7–9} The mechanical properties of a material tell us a lot about its longevity, its ability to withstand damage, and its potential applications.¹⁰ We are now in a period where shape-memory alloys,¹¹ flexible electronics,^{12,13} ferroelastics,^{14–16} and other mechanically-complex and exciting materials are becoming commonplace. As well this, modelling tools such as finite element analysis (FEA) often require prior knowledge of properties such as the Young's modulus to accurately predict the behaviour and performance of materials.^{17,18} More recently, expansions of Pugh mechanical analysis have been used in the derivation of hardness descriptors,¹⁹ in the search for new materials for

hard coating applications. This has also guided researchers in the search for an inorganic compound with a hardness greater than diamond.²⁰ Elastic tensors can also be used to screen for materials with specific thermal properties, as it allows for the estimation of trends in heat capacities and thermal conductivities.^{21–23} By knowing the full anisotropic elastic tensor the elastic response of composite materials can be predicted when combined with mathematical homogenization theories. This has allowed for the design of such materials with engineered stiffness.^{24,25} Additionally, elastic properties are widely used in the field of geophysics, where acoustic velocities can be used to interpret seismic data.^{26,27}

Of all the molecular modelling tools, density functional theory^{28–30} remains one of the most efficient methods of calculating the mechanical properties of a material.^{31–33} While DFT can be used to study almost any material of suitable size, its strength lies in predicting the properties of crystals.^{34–36} Herein we discuss the various DFT methodologies that can be used to calculate the mechanical properties of crystals, as well as exploring other computational chemistry methods that are used today in this endeavour. The limitations of and challenges facing DFT-based predictions mechanical properties are discussed, as well as the exciting future that lies ahead if these challenges are overcome.

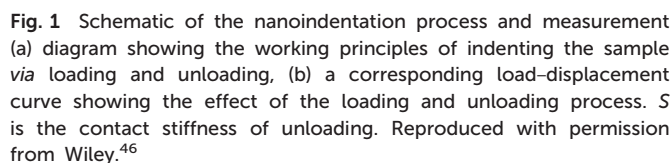
^a Department of Physics, Bernal Institute, University of Limerick, V94 T9PX, Ireland. E-mail: sarah.guerin@ul.ie

^b School of Chemical Sciences, Dublin City University (DCU), Glasnevin, D09 C7F8 Dublin, Ireland

^c SSPC, Science Foundation Ireland Research Centre for Pharmaceuticals, University of Limerick, V94 T9PX, Ireland



In the pharmaceutical industry, it is of utmost importance to understand the elastic and mechanical properties of active pharmaceutical ingredients (APIs). For the API, mechanical properties govern physicochemical properties such as solubility, tabletability, stability and the bioavailability of a drug substance. It is especially important as roughly one in two APIs can exist in multiple solid forms, with each form markedly showing different physicochemical and mechanical properties.^{47,48} Unwanted phase transformations during the development stages (the handling, manufacturing, processing and even the storage) of the API and drug can occur.^{49–51} In the case of polymorphism, the solid–solid transformations can also cause formulation problems. Since polymorphs of the same molecular crystal have differences in interaction energies, other polymorphs tend to transform into the polymorph with the least free energy, and therefore the most stable polymorph. The resulting polymorphs can have undesirable properties, such as in the case of ritonavir and rotigotine.^{52,53} Unwanted phase transformations can affect the drug stability during its lifespan and in the handling of the drug, particularly if a shearing stress is applied. The



piroxicam-succinic acid co-crystal for example is formed *via* the application of mechanical stress to the two components, but undergoes decomposition when shearing occurs. These process-induced transformations are difficult to predict and control due to lack of understanding of the mechanochemical process at an atomistic level. DFT calculations can be a crucial tool to understand and quantify polymorph stability, and can be used to study the interactions between APIs, co-formers, and excipients in both amorphous and crystalline environments.

What is density functional theory?

For single electron systems (2D potential well, hydrogen atom) it is possible to obtain an exact solution to the Schrödinger equation.⁵⁴ However, for any system more complicated than this it is impossible to precisely solve the Schrödinger equation and obtain a mathematical description of the system *i.e.* the wavefunction.⁵⁵ The simplest goal of density functional theory (DFT) calculations is to get an approximate solution to the many-body Schrödinger equation, which gives the ground state properties of the system.

- ion–electron potential energy
- ion–ion potential energy
- electron–electron energy
- kinetic energy
- exchange–correlation energy.

The most computationally challenging energy contributions are the kinetic energy and the exchange–correlation energy. The kinetic energy is calculated using the Kohn–Sham orbitals.⁵⁹ Generally, these do not correspond to actual electron orbitals – they are orthonormal orbitals. The Kohn–Sham orbitals map the system of interacting electrons on to a system of non-interacting electrons moving in an effective potential. The exchange–correlation energy accounts for the exchange interaction due to repulsion between electrons with parallel spins, and the correlation interaction, which is the correlated motion between electrons of anti-parallel spins due to their mutual coulombic repulsion. In its simplest implementation, exchange–correlation effects are treated *via* the Perdew, Burke, and Ernzerhof (PBE)⁶⁰ implementation of the generalised gradient approximation (GGA).⁶¹ GGA builds on what is known as the local density approximation (LDA), by considering both the local electron density and its gradient, as the electron density can vary rapidly over a small region of space.

Long-range considerations: what is dispersion-corrected DFT?

Considering dispersion corrections in DFT calculations is important as the exchange functionals have difficulty in reproducing long-range behaviour, even those explicitly parameterised for long range behaviour.⁶² This is because approximations such as GGA within these functionals⁶³ result in potential energy surfaces that lack sufficient crystal packing effects (CPEs) without the addition of a dispersion correction.⁶⁴ These are influenced by van der Waals forces that are better estimated through dispersion corrections as their inclusion follows a simple formula where the displacement energy is added to the exchange correlation function so as not to be included in the initial DFT calculations.

A vital aspect to the dispersion correction is dampening, with methods that lack an adequate dampening failing to give consistent results for crystal structure and energies.⁶⁵ The dampening function within the dispersion correction determines the range at which the dispersion correction acts⁶⁶ as well as the steepness of the cut off of the dispersion correction.⁶⁷ The dampening function means that the

dispersion effects approach 1 at long distances, this meaning that it is purely a dispersion interaction, but at as the distance between the dipoles shortens the dispersion correction gets dampened eventually going to zero.⁶⁸ This means that there is no dispersion effect at shorter distances where the XC functionals perform better. The dampening effects are also intended to reduce double counting effects.⁶⁹ Further studies are needed on which dampening function performs best with each dispersion correction and exchange functional.⁶⁸ However with increasing numbers of atoms in the unit cell the difficulty arises in deciding where dampening should take place, as this needs to be symmetric.⁷⁰

Dispersion corrections can be calculated in several ways. This can be with a pairwise approach, a three-body approach or a many body approach. The pairwise approach which is chosen in dispersion corrections schemes such as Grimme-D2 (ref. 71 and 72) and Tkatchenko–Scheffler (TS)⁷³ summates over the C_6R^{-6} potential where R is the atomic distance and the C_6 is the dispersion coefficient. For the Grimme-D2 method these are multiplied by a global scaling factor,⁷¹ where the TS scheme calculates the pairwise dispersion energy using the formula:

$$E_{\text{disp}} = -\frac{1}{2} \sum_{i \neq j} f_{\text{damp}} \frac{\left(R_{ij}, R_{ij}^0\right) C_{6ij}}{R_{ij}^6}$$

where f_{damp} is the dampening function, R_{ij} is the distance between atom i and j and R_{ij}^0 is the sum of the van der Waals equilibrium radii.^{74,75} These methods diverge as the DFT-D2 method developed by Grimme utilises the atom-type model whereas the TS model utilises the atomic volume. This use of the atomic volume allows for the approximation of many-body effects.⁷⁰ An example of the three bodied approach is used in the Grimme-D3 correction scheme. The simplest way to employ a three-body approach is to use a non-additive third order Axilrod-Teller-Muto (ATM) term:

$$E_{\text{disp}} = \frac{C_9^{ABC}(3 \cos \theta_a \cos \theta_b \cos \theta_c + 1)}{(R_{AB}R_{BC}R_{CA})^3}$$

where θ represents the internal angles formed by R_{AB} , R_{BC} and R_{CA} .⁷⁶ This uses a C_9 triple dipole constant that can be approximated in two possible ways, either through integrating the average dipole polarizability at an imaginary frequency for all three atoms, or through calculating the square root of the three-atoms dispersion coefficient.⁷⁶ This three-body treatment has been found to contribute minimally to the overall dispersion energy with one paper saying that the three body treatment contributes <5-10% of the dispersion energy,⁷⁷ and another paper stating that the three body effect contributes 7.2% of the lattice energy.⁷⁸

The many body dispersion method, as used in the many body dispersion scheme (MBD), builds on the pairwise TS approach and addresses the fact that the nature of long-range energy is many-body in nature.⁷⁴ The main drawback

This is where $E^{(2)}$ is a pairwise and $E^{(3)}$ is three body energies.⁶⁵ This is extended up to N atoms up to the Nth order this is coupled with a self-consistent screening in order to reduce the error.⁷⁹

Our work to date studying biomolecular crystals using DFT relies on the Vienna *ab initio* simulation package (VASP)⁸⁰ which uses plane wave basis sets,⁸¹ and the projector augmented-wave (PAW) method.⁸² However, many DFT packages, such as CASTEP,⁸³ ABINIT,⁸⁴ and CP2K⁸⁵ also have efficient schemes for calculating elastic properties. For VASP standard PAW pseudopotentials are used in all calculations, as supplied with the software. As our focus has been on the piezoelectric properties⁸⁶ of crystals, elastic constant calculation is part of a calculation workflow, as the elastic stiffness constants c_{kj} are required to calculate the piezoelectric strain constants d_{ik} . The elastic compliance (s_{kj}) can easily be derived from the stiffness and measured alongside electromechanical properties using impedance spectroscopy,^{87,88} as can the various elastic moduli. Using the piezoelectric charge coefficients, e_{ij} , which are also calculated directly by VASP,⁸⁹ and the elastic stiffness constants, c_{kj} , we can calculate the more useful piezoelectric strain coefficient, d_{ik} , using the relationship

$$d_{ik} = e_{ij}/c_{kj}$$

A finite difference method can be used to calculate the elastic stiffness tensor, with every atom in the unit cell being displaced along each Cartesian axis by a default value of ± 0.01 Å. A Γ -centred k -point grid increases mechanical stability, with an observed negligible dependence of predicted elastic constants on the number of k -points (once $N > 1$).⁹⁰ For organic materials high plane wave energy cut-offs of up to 1000 eV are recommended allow the stress tensor to fully converge due to the presence of oxygen and nitrogen atoms.^{91,92} Young's moduli can be derived from the stiffness matrix components. This can be done using the approximations of Nye,⁹³ or Voigt–Reuss–Hill (VRH).^{94,95} Where the software does not automatically derive elastic moduli the ELATE online tool⁹⁶ is recommended for quick validation of mechanical stability, VRH derivation, and 3D visualisation of elastic properties. The ELAM program⁹⁷ can also be used to derive and visualise elastic properties and their anisotropy, with more thorough options for plot generation and customisation.

Using this methodology, we have calculated to a high accuracy *versus* experiment the elastic constants of amino acid^{90,98–100} and peptide crystals,^{101,102} co-crystals,^{103,104} and biominerals¹⁰⁵ (Fig. 2). We have also recently calculated the elastic properties of large protein crystals using classical force fields. Elastic constants of the transmembrane protein ba3 cytochrome c oxidase, as well as lysozyme, and aldehyde dehydrogenase were predicted using the classical CHARMM forcefield model for the protein, ions and water with structures calculated using the CP2K modelling software augmented with homemade subroutines to impose crystal symmetry. In any calculation of elastic properties if crystal symmetry is not preserved then stiffness and compliance tensors will be incorrect. To evaluate the mechanical stability of the crystal we can apply the Born–Huang elastic stability criteria for the appropriate crystal system. As an example for the cubic crystal system, it is required that $c_{11} - c_{12} > 0$, $c_{11} + 2c_{12} > 0$, $c_{44} > 0$. If one or more of these criteria is violated, one or more of the elastic tensor eigenvalues is negative and the crystal is mechanically unstable.

Fig. 2 shows the high quantitative accuracy that can be obtained using the above PBE-only methodology for individual stiffness tensor components and derived Young's modulus values for a small sample of the different classes of crystal that we have studied. As the piezoelectric response is inversely proportional to the elastic stiffness if the material is predicted to be more flexible than it is this will lead to an overestimation of the electromechanical coupling. Full mechanical testing of piezoelectric materials is always recommended as properties such as fracture limit and hardness ultimately determine the specific applications and environments the material can be used in.

Pei & Zeng¹⁰⁶ computed structural and elastic properties of nine phases of piezoelectric polyvinylidene fluoride (PVDF) crystals using DFT with and without a variety of dispersion corrections. In addition to the four known crystalline forms the mechanical properties of five theoretically predicted crystalline forms of PVDF were also investigated. The DFT/PBE calculations show that the cell parameters of four known crystalline forms are in good agreement with experiment. However, they identified that including empirical van der Waals corrections, specifically the Grimme-D2 method, led to a large error in the calculated unit cell lattice parameters. By comparing conventional PBE (without dispersion corrections) and DFT-D2 calculations the authors could highlight that the PBE method provides a better description of the structural and mechanic properties of PVDF crystals.

Complex hybrid systems: limitations of DFT

Kosa *et al.* recently demonstrated a combined computational and experimental approach in studying the elastic anisotropy of a zinc phosphate phosphonoacetate (ZnPA)¹⁰⁷ metal-organic framework (MOF). This 3D framework is composed of Zn-O-Zn layers that are connected by phosphate groups

a. Inorganic Crystals



Quartz

$c_{11} = 89 \text{ GPa}$ (87)
 $c_{33} = 105 \text{ GPa}$ (107)
 $c_{66} = 39 \text{ GPa}$ (40)



Aluminium Nitrate

$c_{11} = 377 \text{ GPa}$ (378)
 $c_{33} = 358 \text{ GPa}$ (392)
 $c_{66} = 123 \text{ GPa}$ (121)



Zinc Oxide

$c_{11} = 189 \text{ GPa}$ (199)
 $c_{33} = 187 \text{ GPa}$ (233)
 $c_{66} = 43 \text{ GPa}$ (44)

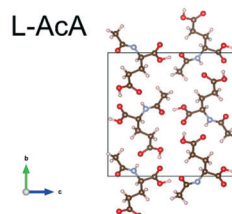
b. Biomineral



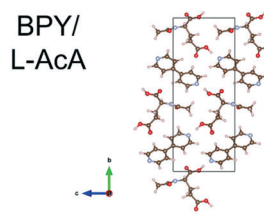
Calcite

Young's Modulus
 $= 73.62 \text{ GPa}$ (72.35)

c. Organic Crystals



Young's Modulus
 $= 8.5 \text{ GPa}$ (10.7)



Young's Modulus
 $= 26.4 \text{ GPa}$ (23.8)



Young's Modulus
 $= 4.2 \text{ GPa}$ (4.3)

Fig. 2 Crystal structures and mechanical properties of different classes of crystal calculated using a PBE-only DFT methodology as published in previous literature^{90,103,105} and summarised in the previous section a. inorganic piezoelectric crystals quartz (SiO_2), aluminium nitrate (AlN), and zinc oxide (ZnO) b. the biomineral calcite (CaCO_3) c. molecular crystals 4,4'-bipyridine (BPY), *N*-acetyl-L-alanine (AcA), and their combined BPY/AcA cocystal. Experimental values are shown in brackets.

bridging the ZnO_4 tetrahedra. This in turn sections off small channels that run parallel to the crystallographic *a*-axis. DFT calculations rationalise that the pore morphology contributes towards the elastic anisotropy. An efficient computational scheme Again a simple and efficient GGA calculation scheme was used, in this case to predict the Young's modulus and the Poisson's ratio (*n*) along the primary crystallographic axes. Notably, they avoided a finite difference methodology in order to circumvent computing the full elastic stiffness tensor (c_{ij}), which can be a computationally heavy approach for low-symmetrical crystal systems.^{38,108} The relative stiffness of the different crystal facets was found to be in reasonably good agreement with experiments. In terms of their absolute values, however, the calculated moduli are consistently higher, by as much as 25% to 40%, than those determined experimentally. Given that the hybrid compound considered here was not moisture sensitive the Young's moduli measured by nanoindentation

are expected to be reliable,¹⁰⁹ as in the authors' previous work. This study pointed out that the higher stiffness calculated by DFT could be due both from the derivation scheme and deficiencies associated with the electronic structure methods used. They discuss how theoretical calculations at 0 K are expected to overestimate the elastic stiffness when compared to experiments performed at ambient 300 K. In relation to the electronic structure method, the authors note that an incorrect description of the position of transition metal d-states can cause over-hybridization of the Zn–O bonds within GGA because of a self-interaction error (SIE). This can lead to artificial stiffening in the zinc–oxygen cores of the ZnPA framework and thus an overestimation of mechanical strength. They emphasise that due to the current limitations of DFT, elastic property predictions for complex hybrid systems are potentially sensitive to the choice of, among others, the exchange–correlation functional and the pseudopotential.



Choosing your functional for high accuracy elastic constants

Dengetal *et al.* highlight the benefits of testing multiple functionals to obtain accurate bulk moduli predictions. The elastic moduli from all functionals (PBE, PBEsol, optB88-vdW) are significantly higher than the values reported experimentally, with PBEsol deviating the least from experimental measurements. The authors attribute this difference to many factors. For example, the samples for characterization in the reported experiments were polycrystalline with both finite grain size and porosity. They are modelled however, as infinite single crystals in plane-wave DFT. The elastic moduli of samples with different grain sizes or porosities can vary, which can also lead to discrepancies in experimental values. The authors also derive Pugh's ratio, G/B , which is commonly used to evaluate the brittleness of materials, with a larger G/B indicating that the material is more brittle.

Another excellent screening of DFT functionals for elastic constant prediction was carried out by Rego & de Koning in their recent study on hexagonal proton-disordered ice using the Quantum ESPRESSO software package.¹¹⁰ They evaluated nine different exchange–correlation functionals, four of which include long-range dispersion interactions through the non-local van der Waals (vdW) approach. While we have observed that dispersion corrections can over-bind crystals and induce polymorphic transitions in small crystals,^{90,106} they are known to play an important role in the condensed phases of water. The authors utilise the well-established energy-strain approach to calculate the elastic constants, in which one exploits the relation between the energy of a crystal and its state of deformation, using the equation

$$E = E_0 + V_0/2 \sum_{m,n=1}^6 C_{mn} \epsilon_m \epsilon_n$$

With the exception of the inferior metaGGA SCAN functional all functionals predict an excessively stiff response to tensile/compressive distortions, as well as shear deformations along the basal plane. These overestimates correlate with lattice constants and other intermolecular distances that are underestimated, and intramolecular separations that are too large. The inclusion of non-local van der Waals corrections generally improves these structural parameters and softens the elastic response functions (Fig. 3).

DFT screening of mechanical properties and database generation

Efforts aimed at developing databases of elastic moduli from first-principles computational methods have been undertaken in numerous studies.^{112,113} These computational approaches are advantageous as all of the data can be derived in a consistent manner, allowing for unambiguous comparisons across many classes of material. De Jong and



Fig. 3 Relative difference of DFT values with respect to experimental data for the structural parameters of ice.^{110,111} Reproduced with permission from AIP Publishing.

colleagues have expanded on this approach, producing the largest database of calculated elastic properties of crystalline inorganic compounds to date, ranging from metals and metallic compounds to semiconductors and insulators. The calculations formed part of the larger high-throughput effort,¹¹⁴ undertaken by the Materials Project.¹¹⁵ Using DFT the calculated elastic constants were consistently shown to be within 15% of experimental values, which represents a smaller scatter than that observed in experimental measurements in some cases (Fig. 4). Pearson (r) and Spearman (ρ) coefficients indicate that the calculations performed in this work yield elastic properties that show an excellent correlation with experimental values, making the database useful for screening materials with properties based on elastic tensors.

For this study the elastic constants were calculated using a stress-strain methodology. Starting from a relaxed structure

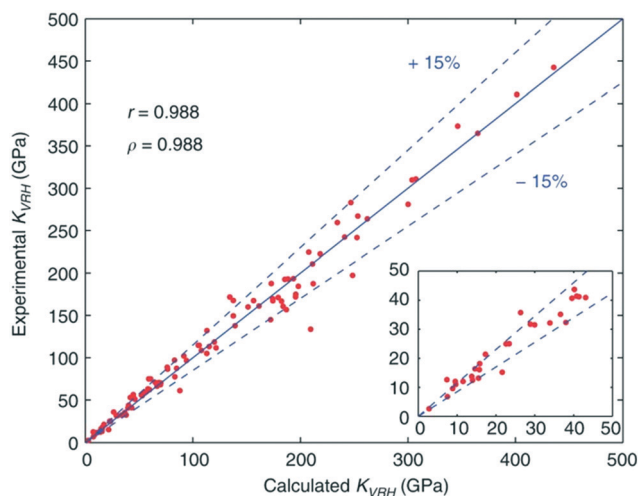


Fig. 4 Comparison of experimental and calculated bulk moduli for a selected set of systems, with calculated Pearson correlation coefficient r and Spearman correlation coefficient ρ reported.¹⁹



Mechanical properties of crystal polymorphs

dielectric,¹¹⁶ or physiochemical¹²³ can also be used to distinguish and rationalize polymorphic behaviour. Since both polymorphism and distinct mechanical properties arise from differences in crystal packing of molecules (in the case of packing polymorphism), polymorphs can be identified by their unique mechanical properties. In addition to colour or morphology, mechanical properties can also be used for the rapid screening and sorting of concomitant polymorphic forms.⁴⁷ This is especially useful for structurally similar polymorphic forms, such as the aspirin polytypes. The structural differences in aspirin I and aspirin II are not obvious. They both constitute hydrogen-bonded dimers ordered in layers, with the layers arranged differently relative to each other.¹²⁴ Reilly and Tkatchenko have shown that their Young's moduli exhibit distinct anisotropies.¹²⁵ In their work they use dispersion-inclusive DFT, by incorporating many-body van der Waals (vdW) interactions to calculate the elastic tensor of the two polymorphic forms. Their elastic property analysis shows that both forms are expected to be mechanically stable under compression and shearing at room temperature – shedding light on the widely-discussed mechanical stability of form II.^{38,126}

The work of Reilly and Tkatchenko shows that extensive characterization of polymorphic forms can be achieved from the determination of elastic tensor using DFT and other *in silico* methods. A strong interconnect between molecular modelling and experiment is preferable, as even with structure–property correlations, a polycrystalline lens is needed to fully understand properties like plasticity and tableability. The tableability of a polycrystalline material is influenced by factors such as the relative movements of grains and grain size, which are difficult to extrapolate from single crystal DFT. Karki *et al.* have successfully used the eigenvalues of the compliance tensor to rationalize the differences in mechanical behavior of paracetamol polymorphic forms – which show significant differences in structural features – as well as paracetamol form II cocrystals.⁴⁸ They used the value of the highest compliance eigenvalue to draw conclusions about the compliance of a crystal and the relative strengths of shear planes. From that, they establish that form II shows a higher compliance eigenvalue compared to form I, because it is compliant to shearing, which leads to plastic deformation during tableting. This result is consistent with the layered structure of form II which grants it preferable compaction properties.¹²⁷ Despite these milestones of linking mechanical behavior to structure using DFT, crystal engineering of pharmaceuticals with desired mechanical behavior is still to be achieved. With the maturity of DFT, crystal structure prediction, and ever-increasing computational power, there is continued opportunity for new insights.

The right direction: the importance of elastic anisotropy

The mechanical properties of crystals are naturally anisotropic, with different stress-strain responses depending on the crystallographic axis of interest. Azuri *et al.*, in their

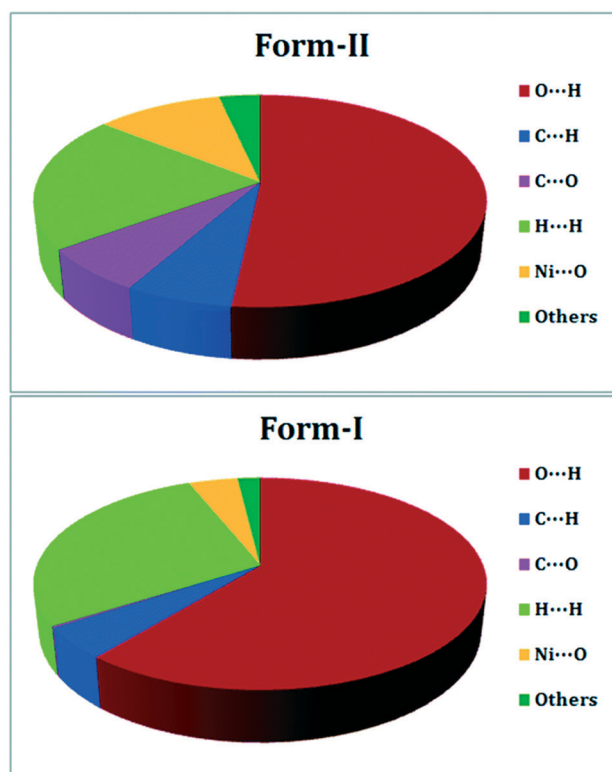
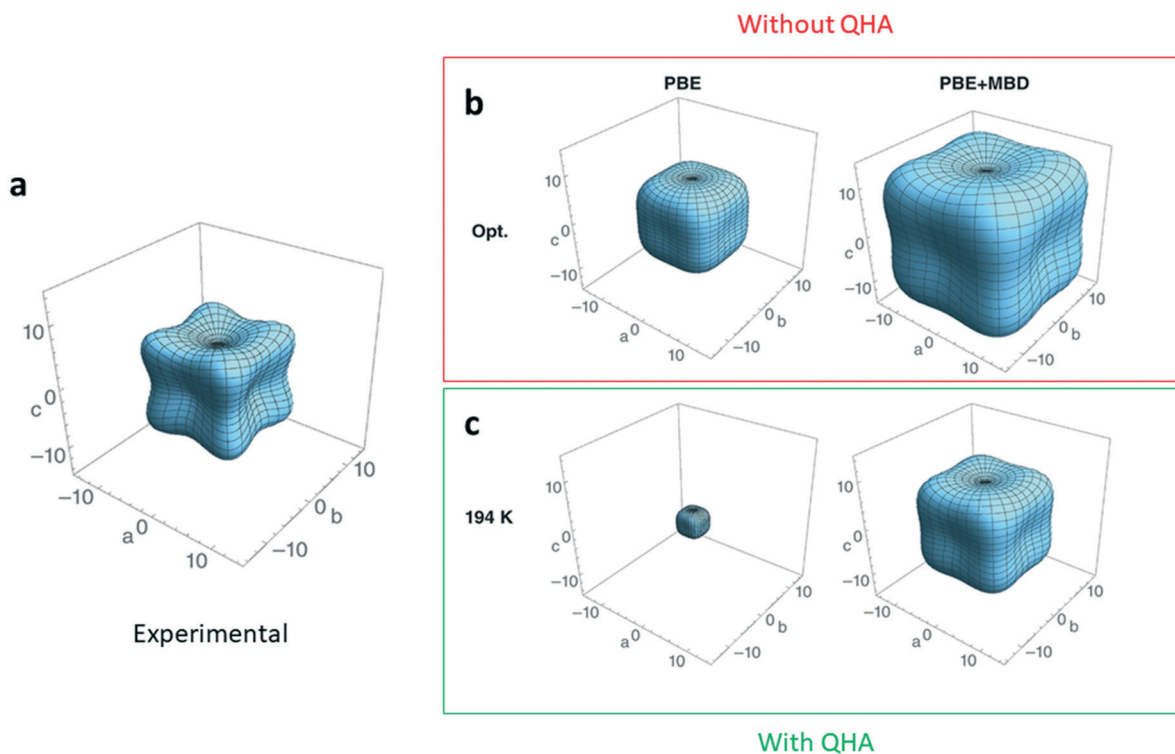


Fig. 5 DFT-calculated intermolecular interactions that contribute to the Hirshfeld surfaces of polymorphic form-II (top) and form-I (bottom) of a 1D Ni(II) polymer with iminodiacetic acid (IDA), namely diaquaiminodiacetonickel(II). Reproduced from ref. 117 with permission from the Royal Society of Chemistry.

The elastic stiffness tensor predicted by some DFT methods is calculated as a 6×6 matrix that naturally describes the elastic anisotropy of a crystalline system. However, it is only recently that elastic anisotropy has begun to be considered in experimental measurement of mechanical properties. Mishra and co-workers systematically examined the mechanical properties of dimorphic forms, forms I and II, of a 1:1 caffeine–glutaric acid cocrystal on multiple faces. Here nanoindentation was used to fully understand the co-crystal mechanical anisotropy and mechanical stability under an applied load.¹³³ The higher hardness and elastic modulus of stable form II was rationalized on the basis of its corrugated layers, higher interlayer energy, lower interlayer separation, and the presence of more intermolecular interactions in the crystal structure compared to metastable form I. The results show that mechanical anisotropy in both polymorphs arises due to the difference in orientation of the identical 2D structural features, namely, the number of possible slip systems and the strength of the intermolecular interactions with respect to the indentation direction. It is hoped that studies like these will influence future experimental investigations, where directional elastic stiffness can be correlated with DFT-



This journal is © The Royal Society of Chemistry 2021

- 12 A. Nathan, A. Ahnood, M. T. Cole, S. Lee, Y. Suzuki, P. Hiralal, F. Bonaccorso, T. Hasan, L. Garcia-Gancedo and A. Dyadyusha, Flexible electronics: the next ubiquitous platform, *Proc. IEEE*, 2012, **100**(Special Centennial Issue), 1486–1517.
- 13 L. Wang, D. Chen, K. Jiang and G. Shen, New insights and perspectives into biological materials for flexible electronics, *Chem. Soc. Rev.*, 2017, **46**(22), 6764–6815.
- 14 S. Baek, H. Jang, C. Folkman, Y. Li, B. Winchester, J. Zhang, Q. He, Y. Chu, C. Nelson and M. Rzechowski, Ferroelastic switching for nanoscale non-volatile magnetoelectric devices, *Nat. Mater.*, 2010, **9**(4), 309–314.
- 15 Y. Hu, L. You, B. Xu, T. Li, S. A. Morris, Y. Li, Y. Zhang, X. Wang, P. S. Lee and H. J. Fan, Ferroelastic-switching-driven large shear strain and piezoelectricity in a hybrid ferroelectric, *Nat. Mater.*, 2021, 1–6.
- 16 T. Seki, C. Feng, K. Kashiwayama, S. Sakamoto, Y. Takasaki, T. Sasaki, S. Takamizawa and H. Ito, Photoluminescent ferroelastic molecular crystals, *Angew. Chem.*, 2020, **132**(23), 8924–8928.
- 17 K. Balasubramanian, N. Rajeswari and K. Vaidheeswaran, Analysis of mechanical properties of natural fibre composites by experimental with FEA, *Mater. Today: Proc.*, 2020, **28**, 1149–1153.
- 18 S. Eshraghi and S. Das, Micromechanical finite-element modeling and experimental characterization of the compressive mechanical properties of polycaprolactone-hydroxyapatite composite scaffolds prepared by selective laser sintering for bone tissue engineering, *Acta Biomater.*, 2012, **8**(8), 3138–3143.
- 19 M. De Jong, W. Chen, T. Angsten, A. Jain, R. Notestine, A. Gamst, M. Sluiter, C. K. Ande, S. Van Der Zwaag and J. J. Plata, Charting the complete elastic properties of inorganic crystalline compounds, *Sci. Data*, 2015, **2**(1), 1–13.
- 20 H. Niu, X.-Q. Chen, P. Liu, W. Xing, X. Cheng, D. Li and Y. Li, Extra-electron induced covalent strengthening and generalization of intrinsic ductile-to-brittle criterion, *Sci. Rep.*, 2012, **2**(1), 1–6.
- 21 D. G. Cahill, S. K. Watson and R. O. Pohl, Lower limit to the thermal conductivity of disordered crystals, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1992, **46**(10), 6131.
- 22 D. R. Clarke, Materials selection guidelines for low thermal conductivity thermal barrier coatings, *Surf. Coat. Technol.*, 2003, **163**, 67–74.
- 23 G. J. Snyder and E. S. Toberer, Complex thermoelectric materials, *Materials for sustainable energy: a collection of peer-reviewed research and review articles from Nature Publishing Group*, 2011, pp. 101–110.
- 24 Z. Hashin and S. Shtrikman, A variational approach to the theory of the elastic behaviour of multiphase materials, *J. Mech. Phys. Solids*, 1963, **11**(2), 127–140.
- 25 T. Zohdi and P. Wriggers, Aspects of the computational testing of the mechanical properties of microheterogeneous material samples, *Int. J. Numer. Methods Eng.*, 2001, **50**(11), 2573–2599.
- 26 O. L. Anderson, E. Schreiber, R. C. Liebermann and N. Soga, Some elastic constant data on minerals relevant to geophysics, *Rev. Geophys.*, 1968, **6**(4), 491–524.
- 27 B. B. Karki, L. Stixrude and R. M. Wentzcovitch, High-pressure elastic properties of major materials of Earth's mantle from first principles, *Rev. Geophys.*, 2001, **39**(4), 507–534.
- 28 K. Burke, Perspective on density functional theory, *J. Chem. Phys.*, 2012, **136**(15), 150901.
- 29 C. Fiolhais, F. Nogueira and M. A. Marques, *A primer in density functional theory*, Springer Science & Business Media, 2003, vol. 620.
- 30 R. G. Parr, Density functional theory of atoms and molecules, in *Horizons of quantum chemistry*, Springer, 1980, pp. 5–15.
- 31 R. Astala, S. M. Auerbach and P. Monson, Density functional theory study of silica zeolite structures: Stabilities and mechanical properties of SOD, LTA, CHA, MOR, and MFI, *J. Phys. Chem. B*, 2004, **108**(26), 9208–9215.
- 32 F. Colmenero, J. Cobos and V. Timón, Periodic density functional theory study of the structure, Raman spectrum, and mechanical properties of Schoepite mineral, *Inorg. Chem.*, 2018, **57**(8), 4470–4481.
- 33 A. Fereidoon, M. G. Ahangari, M. D. Ganji and M. Jahanshahi, Density functional theory investigation of the mechanical properties of single-walled carbon nanotubes, *Comput. Mater. Sci.*, 2012, **53**(1), 377–381.
- 34 M. A. Neumann and M.-A. Perrin, Energy ranking of molecular crystals using density functional theory calculations and an empirical van der Waals correction, *J. Phys. Chem. B*, 2005, **109**(32), 15531–15541.
- 35 P. Ravindran, L. Fast, P. A. Korzhavyi, B. Johansson, J. Wills and O. Eriksson, Density functional theory for calculation of elastic properties of orthorhombic crystals: Application to TiSi₂, *J. Appl. Phys.*, 1998, **84**(9), 4891–4904.
- 36 S. Refaely-Abramson, S. Sharifzadeh, M. Jain, R. Baer, J. B. Neaton and L. Kronik, Gap renormalization of molecular crystals from density-functional theory, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2013, **88**(8), 081204.
- 37 R. Golesorkhtabar, P. Pavone, J. Spitaler, P. Puschnig and C. Draxl, ElaStic: A tool for calculating second-order elastic constants from first principles, *Comput. Phys. Commun.*, 2013, **184**(8), 1861–1873.
- 38 J. D. Bauer, E. Haussühl, B. R. Winkler, D. Arbeck, V. Milman and S. Robertson, Elastic properties, thermal expansion, and polymorphism of acetylsalicylic acid, *Cryst. Growth Des.*, 2010, **10**(7), 3132–3140.
- 39 J. C. Tan and A. K. Cheetham, Mechanical properties of hybrid inorganic-organic framework materials: establishing fundamental structure-property relationships, *Chem. Soc. Rev.*, 2011, **40**(2), 1059–1080.
- 40 T. D. Bennett, J. C. Tan, S. A. Moggach, R. Galvelis, C. Mellot-Draznieks, B. A. Reisner, A. Thirumurugan, D. R. Allan and A. K. Cheetham, Mechanical Properties of Dense Zeolitic Imidazolate Frameworks(ZIFs): A High-Pressure X-ray Diffraction, Nanoindentation and Computational



- Study of the Zinc Framework Zn(Im)₂, and its Lithium Boron Analogue, LiB(Im)₄, *Chem. – Eur. J.*, 2010, **16**(35), 10684–10690.
- 41 C. Boissiere, D. Grosso, S. Lepoutre, L. Nicole, A. B. Bruneau and C. Sanchez, Porosity and mechanical properties of mesoporous thin films assessed by environmental ellipsometric porosimetry, *Langmuir*, 2005, **21**(26), 12362–12371.
 - 42 A. C. Fischer-Cripps and D. Nicholson, Nanoindentation. Mechanical engineering series, *Appl. Mech. Rev.*, 2004, **57**(2), B12.
 - 43 D. M. Ebenstein and L. A. Pruitt, Nanoindentation of biological materials, *Nano Today*, 2006, **1**(3), 26–33.
 - 44 H. Attar, S. Ehtemam-Haghighi, D. Kent, I. Okulov, H. Wendrock, M. Bönisch, A. Volegov, M. Calin, J. Eckert and M. Dargusch, Nanoindentation and wear properties of Ti and Ti-TiB composite materials produced by selective laser melting, *Mater. Sci. Eng., A*, 2017, **688**, 20–26.
 - 45 G. Konstantopoulos, E. P. Koumoulos and C. A. Charitidis, Classification of mechanism of reinforcement in the fiber-matrix interface: Application of Machine Learning on nanoindentation data, *Mater. Des.*, 2020, **192**, 108705.
 - 46 R. He, S. Gahlawat, C. Guo, S. Chen, T. Dahal, H. Zhang, W. Liu, Q. Zhang, E. Chere and K. White, Studies on mechanical properties of thermoelectric materials by nanoindentation, *Phys. Status Solidi A*, 2015, **212**(10), 2191–2195.
 - 47 C. M. Reddy, S. Basavoju and G. R. Desiraju, Sorting of polymorphs based on mechanical properties. Trimorphs of 6-chloro-2, 4-dinitroaniline, *Chem. Commun.*, 2005(19), 2439–2441.
 - 48 S. Karki, T. Friščić, L. Fabian, P. R. Laity, G. M. Day and W. Jones, Improving mechanical properties of crystalline solids by cocrystal formation: new compressible forms of paracetamol, *Adv. Mater.*, 2009, **21**(38–39), 3905–3909.
 - 49 K. R. Morris, U. J. Griesser, C. J. Eckhardt and J. G. Stowell, Theoretical approaches to physical transformations of active pharmaceutical ingredients during manufacturing processes, *Adv. Drug Delivery Rev.*, 2001, **48**(1), 91–114.
 - 50 G. G. Zhang, D. Law, E. A. Schmitt and Y. Qiu, Phase transformation considerations during process development and manufacture of solid oral dosage forms, *Adv. Drug Delivery Rev.*, 2004, **56**(3), 371–390.
 - 51 I. Tumanov, A. Achkasov, S. Myz, E. Boldyreva and V. Boldyrev, in Different effect of impact and shear mechanical treatment on mechanochemical cocrystallization of piroxicam and succinic acid, *Doklady Chemistry*, Pleiades Publishing, 2014, pp. 154–159.
 - 52 J. Bauer, S. Spanton, R. Henry, J. Quick, W. Dziki, W. Porter and J. Morris, Ritonavir: an extraordinary example of conformational polymorphism, *Pharm. Res.*, 2001, **18**(6), 859–866.
 - 53 I. B. Rietveld and R. Céolin, Rotigotine: unexpected polymorphism with predictable overall monotropic behavior, *J. Pharm. Sci.*, 2015, **104**(12), 4117–4122.
 - 54 M. F. Manning, Exact solutions of the Schrödinger equation, *Phys. Rev.*, 1935, **48**(2), 161.
 - 55 P. A. M. Dirac, *The principles of quantum mechanics*, Oxford university press, 1981.
 - 56 W. Kohn, A. D. Becke and R. G. Parr, Density functional theory of electronic structure, *J. Phys. Chem.*, 1996, **100**(31), 12974–12980.
 - 57 P. Hohenberg and W. Kohn, Inhomogeneous electron gas, *Phys. Rev.*, 1964, **136**(3), B864.
 - 58 W. Koch and M. C. Holthausen, *A chemist's guide to density functional theory*, John Wiley & Sons, 2015.
 - 59 J. P. Perdew and M. Levy, Physical content of the exact Kohn-Sham orbital energies: band gaps and derivative discontinuities, *Phys. Rev. Lett.*, 1983, **51**(20), 1884.
 - 60 J. P. Perdew and W. Yue, Accurate and simple density functional for the electronic exchange energy: Generalized gradient approximation, *Phys. Rev. B*, 1986, **33**(12), 8800.
 - 61 J. P. Perdew, K. Burke and M. Ernzerhof, Generalized gradient approximation made simple, *Phys. Rev. Lett.*, 1996, **77**(18), 3865.
 - 62 S. Kristyán and P. Pulay, Can(semi) local density functional theory account for the London dispersion forces?, *Chem. Phys. Lett.*, 1994, **229**(3), 175–180.
 - 63 J. G. Brandenburg and S. Grimme, Dispersion corrected Hartree-Fock and density functional theory for organic crystal structure prediction, *Prediction and Calculation of Crystal Structures*, 2013, pp. 1–23.
 - 64 J. Moellmann and S. Grimme, Importance of London dispersion effects for the packing of molecular crystals: a case study for intramolecular stacking in a bis-thiophene derivative, *Phys. Chem. Chem. Phys.*, 2010, **12**(30), 8500–8504.
 - 65 P. Xu, M. Alkan and M. S. Gordon, Many-Body Dispersion, *Chem. Rev.*, 2020, **120**(22), 12343–12356.
 - 66 S. Grimme, Density functional theory with London dispersion corrections, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2011, **1**(2), 211–228.
 - 67 M. D. King and T. M. Korter, Modified corrections for London forces in solid-state density functional theory calculations of structure and lattice dynamics of molecular crystals, *J. Phys. Chem. A*, 2012, **116**(25), 6927–6934.
 - 68 Y. Liu and W. A. I. Goddard, A universal damping function for empirical dispersion correction on density functional theory, *Mater. Trans.*, 2009, **50**(7), 1664–1670.
 - 69 S. Grimme, S. Ehrlich and L. Goerigk, Effect of the damping function in dispersion corrected density functional theory, *J. Comput. Chem.*, 2011, **32**(7), 1456–1465.
 - 70 A. Otero-de-la-Roza, L. M. LeBlanc and E. R. Johnson, What is “many-body” dispersion and should I worry about it?, *Phys. Chem. Chem. Phys.*, 2020, **22**(16), 8266–8276.
 - 71 S. Grimme, Accurate description of van der Waals complexes by density functional theory including empirical corrections, *J. Comput. Chem.*, 2004, **25**(12), 1463–1473.
 - 72 S. Grimme, Semiempirical GGA-type density functional constructed with a long-range dispersion correction, *J. Comput. Chem.*, 2006, **27**(15), 1787–1799.
 - 73 A. Tkatchenko and M. Scheffler, Accurate molecular van der Waals interactions from ground-state electron density



103 W. Ji, *et al.*, Tunable Mechanical and Optoelectronic Properties of Organic Cocrystals by Unexpected Stacking

- Transformation from H-to J-and X-Aggregation, *ACS Nano*, 2020, **14**(8), 10704–10715.
- 104 W. Ji, *et al.*, Modulation of physical properties of organic cocrystals by amino acid chirality, *Mater. Today*, 2021, **42**, 29–40.
 - 105 S. Guerin, S. A. Tofail and D. Thompson, Longitudinal piezoelectricity in natural calcite materials: Preliminary studies, *IEEE Trans. Dielectr. Electr. Insul.*, 2018, **25**(3), 803–807.
 - 106 Y. Pei and X. C. Zeng, Elastic properties of poly(vinylidene fluoride)(PVDF) crystals: A density functional theory study, *J. Appl. Phys.*, 2011, **109**(9), 093514.
 - 107 M. Kosa, J. C. Tan, C. A. Merrill, M. Krack, A. K. Cheetham and M. Parrinello, Probing the Mechanical Properties of Hybrid Inorganic–Organic Frameworks: A Computational and Experimental Study, *ChemPhysChem*, 2010, **11**(11), 2332–2336.
 - 108 F. M. Capaldi, M. C. Boyce and G. C. Rutledge, The mechanical properties of crystalline cyclopentyl polyhedral oligomeric silsesquioxane, *J. Chem. Phys.*, 2006, **124**(21), 214709.
 - 109 J. Tan, C. Merrill, J. Orton and A. Cheetham, Anisotropic mechanical properties of polymorphic hybrid inorganic–organic framework materials with different dimensionalities, *Acta Mater.*, 2009, **57**(12), 3481–3496.
 - 110 J. Santos Rego and M. de Koning, Density-functional theory prediction of the elastic constants of ice I_h, *J. Chem. Phys.*, 2020, **152**(8), 084502.
 - 111 K. Röttger, A. Endriss, J. Ihringer, S. Doyle and W. Kuhs, Lattice constants and thermal expansion of H₂O and D₂O ice I_h between 10 and 265 K, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1994, **50**(6), 644–648.
 - 112 C. R. Da Silva, P. R. da Silveira, B. Karki, R. M. Wentzcovitch, P. A. Jensen, E. F. Bollig, M. Pierce, G. Erlebacher and D. A. Yuen, Virtual laboratory for planetary materials: System service architecture overview, *Phys. Earth Planet. Inter.*, 2007, **163**(1–4), 321–332.
 - 113 P. R. Da Silveira, C. R. da Silva and R. M. Wentzcovitch, Metadata management for distributed first principles calculations in VLab—A collaborative cyberinfrastructure for materials computation, *Comput. Phys. Commun.*, 2008, **178**(3), 186–198.
 - 114 S. Curtarolo, G. L. Hart, M. B. Nardelli, N. Mingo, S. Sanvito and O. Levy, The high-throughput highway to computational materials design, *Nat. Mater.*, 2013, **12**(3), 191–201.
 - 115 A. Jain, S. P. Ong, G. Hautier, W. Chen, W. D. Richards, S. Dacek, S. Cholia, D. Gunter, D. Skinner and G. Ceder, Commentary: The Materials Project: A materials genome approach to accelerating materials innovation, *APL Mater.*, 2013, **1**(1), 011002.
 - 116 S. Roy, A. Hazra, A. Bandyopadhyay, D. Raut, P. L. Madhuri, D. S. Rao, U. Ramamurty, S. K. Pati, S. Krishna Prasad and T. K. Maji, Reversible Polymorphism, Liquid Crystallinity, and Stimuli-Responsive Luminescence in a Bola-amphiphilic π -System: Structure–Property Correlations Through Nanoindentation and DFT Calculations, *J. Phys. Chem. Lett.*, 2016, **7**(20), 4086–4092.
 - 117 S. K. Seth, A. Bauzá and A. Frontera, Screening polymorphism in a Ni(II) metal–organic framework: experimental observations, Hirshfeld surface analyses and DFT studies, *CrystEngComm*, 2018, **20**(6), 746–754.
 - 118 S. Wen and G. J. Beran, Crystal polymorphism in oxalyl dihydrazide: Is empirical DFT-D accurate enough?, *J. Chem. Theory Comput.*, 2012, **8**(8), 2698–2705.
 - 119 A. Pedone, D. Presti and M. C. Menziani, On the ability of periodic dispersion-corrected DFT calculations to predict molecular crystal polymorphism in para-diiodobenzene, *Chem. Phys. Lett.*, 2012, **541**, 12–15.
 - 120 K. Hongo, M. A. Watson, R. S. Sánchez-Carrera, T. Iitaka and A. Aspuru-Guzik, Failure of conventional density functionals for the prediction of molecular crystal polymorphism: a Quantum Monte Carlo Study, *J. Phys. Chem. Lett.*, 2010, **1**(12), 1789–1794.
 - 121 J. Van De Streek and M. A. Neumann, Validation of experimental molecular crystal structures with dispersion-corrected density functional theory calculations, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2010, **66**(5), 544–558.
 - 122 O. U. O. Mota, R. A. Araujo, H. Wang and T. Çağın, Mechanical properties of metal nitrides for radiation resistant coating applications: a DFT study, *Phys. Procedia*, 2015, **66**, 576–585.
 - 123 L. F. L. Silva, W. Paschoal, G. S. Pinheiro, J. G. da Silva Filho, P. T. Freire, F. F. de Sousa and S. G. Moreira, Understanding the effect of solvent polarity on the polymorphism of octadecanoic acid through spectroscopic techniques and DFT calculations, *CrystEngComm*, 2019, **21**(2), 297–309.
 - 124 A. D. Bond, R. Boese and G. R. Desiraju, On the polymorphism of aspirin: crystalline aspirin as intergrowths of two “polymorphic” domains, *Angew. Chem., Int. Ed.*, 2007, **46**(4), 618–622.
 - 125 A. M. Reilly and A. Tkatchenko, Understanding the role of vibrations, exact exchange, and many-body van der Waals interactions in the cohesive properties of molecular crystals, *J. Chem. Phys.*, 2013, **139**(2), 024705.
 - 126 S. Varughese, M. Kiran, K. A. Solanko, A. D. Bond, U. Ramamurty and G. R. Desiraju, Interaction anisotropy and shear instability of aspirin polymorphs established by nanoindentation, *Chem. Sci.*, 2011, **2**(11), 2236–2242.
 - 127 T. Beyer, G. M. Day and S. L. Price, The prediction, morphology, and mechanical properties of the polymorphs of paracetamol, *J. Am. Chem. Soc.*, 2001, **123**(21), 5086–5094.
 - 128 I. Azuri, E. Meirzadeh, D. Ehre, S. R. Cohen, A. M. Rappe, M. Lahav, I. Lubomirsky and L. Kronik, Unusually large Young's moduli of amino acid molecular crystals, *Angew. Chem., Int. Ed.*, 2015, **54**(46), 13566–13570.
 - 129 I. Diddens, B. Murphy, M. Krisch and M. Müller, Anisotropic elastic properties of cellulose measured using inelastic X-ray scattering, *Macromolecules*, 2008, **41**(24), 9755–9759.



- 130 A. Ishikawa, T. Okano and J. Sugiyama, Fine structure and tensile properties of ramie fibres in the crystalline form of cellulose I, II, III and IV, *Polymer*, 1997, **38**(2), 463–468.
- 131 R. Wagner, R. Moon, J. Pratt, G. Shaw and A. Raman, Uncertainty quantification in nanomechanical measurements using the atomic force microscope, *Nanotechnology*, 2011, **22**(45), 455703.
- 132 F. L. Dri, L. G. Hector, R. J. Moon and P. D. Zavattieri, Anisotropy of the elastic properties of crystalline cellulose I β from first principles density functional theory with Van der Waals interactions, *Cellulose*, 2013, **20**(6), 2703–2718.
- 133 M. K. Mishra, *et al.*, Structural Basis for Mechanical Anisotropy in Polymorphs of a Caffeine–Glutaric Acid Cocrystal, *Cryst. Growth Des.*, 2020, **20**(10), 6306–6315.
- 134 J. Hoja, A. M. Reilly and A. Tkatchenko, First-principles modeling of molecular crystals: structures and stabilities, temperature and pressure, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2017, **7**(1), e1294.
- 135 A. Togo and I. Tanaka, First principles phonon calculations in materials science, *Scr. Mater.*, 2015, **108**, 1–5.
- 136 A. Erba, J. Maul and B. Civalleri, Thermal properties of molecular crystals through dispersion-corrected quasi-harmonic ab initio calculations: the case of urea, *Chem. Commun.*, 2016, **52**(9), 1820–1823.
- 137 H. Kiefte, S. Breckon, R. Penney and M. Clouter, Elastic constants of ammonia by Brillouin spectroscopy, *J. Chem. Phys.*, 1985, **83**(9), 4738–4743.
- 138 T. Kamencek, S. Wieser, H. Kojima, N. Bedoya-Martínez, J. P. Dürholt, R. Schmid and E. Zofer, Evaluating Computational Shortcuts in Supercell-Based Phonon Calculations of Molecular Crystals: The Instructive Case of Naphthalene, *J. Chem. Theory Comput.*, 2020, **16**(4), 2716–2735.
- 139 J. G. Brandenburg and S. Grimme, Accurate modeling of organic molecular crystals by dispersion-corrected density functional tight binding(DFTB), *J. Phys. Chem. Lett.*, 2014, **5**(11), 1785–1789.

