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Unusual seeding effect in the liquid-assisted high-pressure polymorphism of chlorpropamide†

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When crystals of four polymorphs – a stable α -form and three metastable ones (β -, γ -, δ -forms) – are simultaneously present in a pressure cell, immersed in a 1:1 pentane–isopentane mixture, the least stable β -form recrystallises exclusively into the γ -form (if no δ -form seed is present) or concomitantly into the γ - and δ -forms (if a seed of the δ -form is present). However, the β -form never recrystallises to the stable α -form, even if the latter is also present as a potential seed.

Studies of polymorphic molecular crystals are important in many fields of chemistry, including, among others, theory of intermolecular interactions, supramolecular chemistry, crystal engineering, crystallisation theory, chemical engineering, pharmaceutical chemistry, and high-pressure science.^{1–11} For a compound prone to polymorphism, it is not possible to predict which phase will be formed at a selected (T , P) point based solely on a thermodynamic phase diagram. Instead, there exists a complex interplay of nucleation and growth kinetics, alongside thermodynamics, that leads to these unpredictable results. This is reflected in phenomena which were termed “concomitant polymorphism” (several phases coexist under the same conditions, often in the same batch),¹² “disappearing polymorphs” (a fast-growing metastable polymorph obtained once can no longer be crystallised after a seed of the stable form has been obtained),^{13,14} “pre-nucleation” (molecular clusters in solution pre-determine the structure of a crystalline nucleus formed at later stages),¹⁵

“seeding-triggered crystallisation” (a seed of a phase triggers the crystallisation of the same phase),¹⁶ and “template crystallisation” (molecules or molecular clusters in solution influence crystallisation of a certain crystalline form, even if not included in the final crystal structure themselves).^{17,18} High pressure has been recently proposed as a tool to obtain new polymorphs of pharmaceuticals, which, if preserved on flash-decompression, can be used as seeds for mass crystallisation under ambient conditions.^{19–23} However, the control of high-pressure polymorphism is not a trivial task.^{19,24–28} The role of kinetic factors known to influence crystallisation under ambient conditions becomes enormously significant at high pressure, when molecular motions are even more restricted, the sample is located in a confined space of a gasket hole, and the pressure-transmitting fluid is not mixed and is often viscous. Thus diffusion and convection processes are hindered. As a result of the kinetic control of nucleation and nuclei growth,²⁸ different phases can form, depending on the choice of the starting polymorph,^{29–31} the hydrostatic medium,^{32–37} or the compression/decompression protocol.^{26,38–42}

An antidiabetic drug, chlorpropamide (4-chloro-*N*-(propylaminocarbonyl)benzenesulfonamide, $C_{10}H_{13}ClN_2O_3S$), is particularly prone to polymorphism.^{43,44} In addition to the commercially available α -polymorph, which is the thermodynamically stable form under ambient conditions,⁴⁵ four other polymorphs (β -, γ -, δ -, ϵ -) can be preserved for an indefinitely long period time under ambient conditions. These four additional phases can be obtained either from solution crystallisation in different solvents (β -, γ -, δ -) or at high temperature (ϵ -).^{46–48} All known polymorphs have similar hydrogen-bonded chains involving the urea-like core, but differ in the conformation adopted by the phenyl ring and alkyl tails, and the relative orientation of the hydrogen-bonded chains (Fig. 1 and Table S1 in the ESI†). Chlorpropamide is therefore an ideal model system with which the roles of molecular flexibility and intermolecular interactions in forming and rearranging the crystal structures of organic crystals can be studied. The five polymorphs respond differently to

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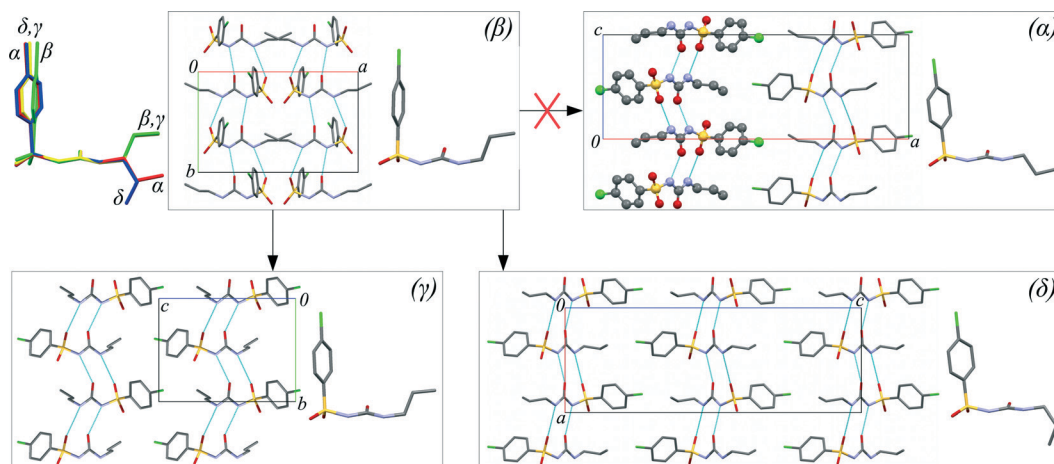


Fig. 1 Molecular packing and molecular conformations of different chlorpropamide polymorphs (α -, γ -, δ -, β -forms). Possible routes of β -polymorph recrystallisation in a pentane-isopentane mixture (1:1) in a diamond anvil cell at 0.3 GPa are shown by arrows. The molecular overlay is shown with the “urea core” taken as a common reference plane. The chain with an inverted orientation (every second chain) in the structure of the α -form is highlighted in a ball-and-stick representation.

cooling^{49–51} and heating,⁴⁸ with each ambient-condition polymorph transforming into a unique non-ambient-temperature form. One could expect similarly high versatility in behavior on hydrostatic compression.

To date, only the effect of pressure on the α -polymorph, which is the stable form at ambient pressure,⁴⁵ has been reported.^{34–37} The α -polymorph undergoes at least one phase transition with increasing pressure in saturated ethanol solution (recrystallisation is possible).³⁷ The same phase transition seems to also take place in dry powder samples without adding any pressure-transmitting fluids,³⁴ though the transition in this case was reported to be kinetically hindered, and thus believed to be solvent-assisted.³⁴ In the present work, we aimed to monitor the pressure-induced structural response of four polymorphs in a fluid that does not dissolve any of the polymorphs at ambient pressure. However, our experiments have shown that one of the metastable polymorphs, the β -form, starts recrystallising as the pressure increases. Interestingly, the other polymorphs do not dissolve in the same fluid, even at high pressures. Even more interestingly, a crystal of the most stable polymorph (the α -form) present in the pressure cell simultaneously with the β -polymorph does not act as a seed, whereas another metastable polymorph (the δ -form) does. We report these observations and rationalise them in the present paper.

Crystals of the α -, β -, and δ -polymorphs were simultaneously loaded into a diamond anvil cell (DAC) to ensure identical compression conditions for a better comparison of their response to increasing pressure. Samples were kept at ambient temperature in all experiments. A pentane-isopentane mixture (1:1) was selected as the pressure-transmitting medium, which does not visibly dissolve any of the three chlorpropamide polymorphs under ambient conditions and has a high hydrostatic limit of 7 GPa.⁵² The pressure was increased directly to 0.5 GPa and held for two hours. Following this period, the β -polymorph had dissolved very slightly (the

crystal edges had become slightly rounded), whereas the α - and δ -polymorphs had not changed (Fig. 2a and S1†). The following day, the pressure in the DAC had spontaneously decreased to 0.3 GPa, and small, thin needles had appeared. Additionally, the crystal of the δ -polymorph had increased in size and the crystal of the β -polymorph had begun dissolving, whereas the crystal of the α -form had remained visually unchanged (Fig. 2a and S1†).

High-pressure vibrational spectra were measured *in situ* for each individual polymorph crystal by using a Raman confocal microscope. The Raman spectra of the newly formed needle-shaped crystals did not match those of any of the polymorphs originally loaded into the DAC (α -, β -, or δ -forms). However, the Raman data were of insufficient quality to reliably identify the new phase (Fig. 3). To identify the new needle-shaped phase unambiguously, a single crystal of β -chlorpropamide was placed alone in the DAC, again in a pentane-isopentane mixture (1:1). Partial dissolution of the crystal and the growth of multiple tiny needles (with similar appearance to those described above) were observed (Fig. 2b). An attempt to characterize this phase *in situ* by single-crystal X-ray diffraction, even using synchrotron radiation, was unsuccessful; the sample contained too many crystallites for a successful single-crystal diffraction approach, and too few with a similar orientation for a powder diffraction experiment. In order to study the structure of a selected crystal of the new phase, the DAC was thus decompressed and opened. The hydrostatic fluid evaporated immediately upon opening the cell, and the needles of the high-pressure phase did not dissolve or recrystallize. An individual needle (of approximate size $0.15 \times 0.01 \times 0.01$ mm³) was selected and single-crystal diffraction showed that the needles corresponded to γ -chlorpropamide.⁴⁷ This was particularly surprising as it is known to be less dense under ambient pressure (1.416 g cm^{−3}) than the α - and δ -polymorphs (1.450 and 1.455 g cm^{−3}, respectively^{48,49}). An additional experiment





Fig. 2 Crystals of the α -, β -, and δ -polymorphs of chlorpropamide (a) and the crystals of the β -polymorph of chlorpropamide (b) loaded in a pentane-isopentane mixture (1:1) in a diamond anvil cell at different time moments. The growth of new needle-shaped crystals (γ -polymorph) and growth of the δ -polymorph (increase in size of the originally available crystals) are clearly seen in the presence of the α - and δ -phases (a), whereas only the recrystallisation of the β - into the γ -polymorph (needles) is clearly seen if the β -phase is present alone (b).

was carried out to confirm that the γ -polymorph in pentane-isopentane does not undergo any phase transitions in the pressure range of these experiments (below 1 GPa), either on compression or on decompression. Thus, in addition to optical microscopy observations, X-ray diffraction completely excludes the possibility that the γ -polymorph might be formed during decompression of another high-pressure phase or upon subsequent opening of the DAC.

These experiments proved to be highly reproducible. Invariably, in the absence of δ -form seeds, needles of the γ -form recrystallised from the β -polymorph. Alternatively, in the presence of a δ -form seed, simultaneous slow recrystallisation of the β -crystal into both the γ -form (new crystals) and the δ -form (an increase in the size of the seed crystals) was observed. It therefore appears that the high-pressure trans-



Fig. 3 Raman spectra of the α -, β -, and δ -polymorphs of chlorpropamide and the new needle-shaped (γ) polymorph at the first pressure point (0.3 GPa). No solid-state transformations occur in the starting α - and δ -forms at this pressure. Characteristic features of the spectra for the different polymorphs are shown by arrows.

formations of β -chlorpropamide below 0.5 GPa are not solid-state phase transitions, but result from recrystallisation. Interestingly, the α -form – the most stable polymorph under ambient conditions, though not the densest one⁴⁵ – neither acted as a seed in the recrystallisation of the β - or γ -polymorphs nor recrystallised into the denser δ -form.

It would not be expected that the relative densities of the polymorphs could change in the absence of any phase transitions before 0.5 GPa. To verify this assumption, the unit cell volumes of the α -, γ -, and δ -polymorphs were measured at high pressure using single-crystal X-ray diffraction. All three crystals were loaded into the same DAC simultaneously, ensuring identical conditions for each polymorph. Paraffin was used as the pressure-transmitting medium in lieu of pentane-isopentane to exclude any recrystallisation events and allow hydrostatic conditions up to 5 GPa.⁵³ The relative densities were calculated from unit cell parameters measured at 0.35 and 0.50 GPa (see the ESI† for cell parameters). The densities at 0.35 and 0.50 GPa were as follows: α -polymorph – 1.528 and 1.546 g cm⁻³; γ -polymorph – 1.522 and 1.549 g cm⁻³; δ -polymorph – 1.534 and 1.554 g cm⁻³, respectively.

Based on the relative densities, one might expect three high-pressure recrystallisation pathways for β -chlorpropamide: $\beta \rightarrow \alpha$, $\beta \rightarrow \gamma$, or $\beta \rightarrow \delta$. However, only the two latter processes are observed. It is rather common that the presence of a stable polymorph suppresses the crystallisation of a metastable form.⁵⁴ In contrast, the high-pressure behavior of chlorpropamide, when a seed of the most stable (α -) polymorph does not grow, but instead, one metastable polymorph (γ -) grows as new crystals even without seeding, and another metastable form present as a seed (δ -) increases in size, is highly unusual. It therefore becomes important to consider the relative rates of the required molecular re-

The complex mutual effect of chlorpropamide polymorphs at relatively low (0.5 GPa) pressures is not the only manifestation of the kinetic control of phase transitions in this system. As crystals of the β -polymorph recrystallised

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Conclusions

The example of chlorpropamide adds a new dimension to studying the complexities of the polymorphism of organic crystals in general, and of high-pressure polymorphism, in particular. Kinetic barriers for the structural rearrangement in the solid state and crystallisation at high pressure from solution can be significant and result in the formation of metastable materials. The resulting products may simply be those with the lowest kinetic barrier for nucleation.⁶⁰ This barrier can obviously be changed in the presence of a seed, but not every polymorph can act as a seed (even being denser and even being more stable; these two criteria do not always coincide). Further, even in the presence of the most stable phase as a seed, other crystallisation routes to relatively less stable forms remain possible. The solid-state transformations can also give not the thermodynamically most stable phase, but the polymorph which can be formed more easily from the starting crystal structure.⁶¹ Transformations in a diamond anvil pressure cell can combine the restrictions of confined crystallisation in a small volume with the limitations of convection, the diffusion of molecules through a viscous medium and the difficulties of structural rearrangement of molecules and molecular packing in the solid state.

Experimental

Single crystals of α - and δ -chlorpropamide were grown using techniques described in ref. 45. β -Chlorpropamide was grown from saturated ethanol solution.

Pressure was generated using a Boehler-Almax⁶² diamond anvil cell (DAC) with a stainless steel gasket (hole diameter 300 μm). The ruby fluorescence line was used for pressure calibration with an accuracy of ± 0.05 GPa.⁶³ A pentane–isopentane mixture was used as the pressure-transmitting medium.⁵² Paraffin was used as the pressure-transmitting medium⁵³ in the high-pressure X-ray diffraction experiment aimed at determining the relative densities of the α -, γ -, and δ -polymorphs of chlorpropamide. An Oxford Diffraction Gemini R Ultra diffractometer (Mo $K\alpha$) and CrysAlisPro⁶⁴ software were used to determine the cell parameters.

Optical microscopy experiments were carried out using a Nikon AZ100 microscope.

A Horiba Jobin Yvon LabRam HR800 spectrometer was used for Raman spectroscopy. The spectra were obtained with the 532 nm line of a 40 mW neodymium solid-state laser (Nd:YAG neodymium-doped yttrium aluminium garnet $\text{Nd:Y}_3\text{Al}_5\text{O}_{12}$ with a frequency-doubled line). The spectra were recorded on a 1024-channel LN/CCD detector from -10 to 3800 cm^{-1} with 2 cm^{-1} spectral resolution. The power of the laser beam on the sample inside the DAC was below 10 mW. Raman spectra were collected in back-scattering geometry using an Olympus BX41 microscope. This microscope used an Olympus $50\times$ objective lens with a long working distance of $\text{LWD} = 11\text{ mm}$. Using a 0.5 numerical aperture produced a focal spot diameter of $\sim 2\text{ }\mu\text{m}$. An edge filter almost completely suppressed the low-

frequency Raman spectrum below 60 cm^{-1} and attenuated that over the range $60\text{--}100\text{ cm}^{-1}$. The laser light that remained after scattering and collection in the spectrometer, subsequently suppressed by the edge filter to the residual low-intensity line at 0 cm^{-1} , was used for additional calibration correction of each recorded spectrum.

Single-crystal synchrotron X-ray diffraction data were collected at the Swiss-Norwegian Beamline BM01A at the European Synchrotron Radiation Facility (ESRF, Grenoble, France, experiment CH-4526 (PILATUS 2M hybrid pixel detector, wavelength $0.81984\text{ }\text{\AA}$). The data were converted and integrated using the SNBL toolbox⁶⁵ and CrysAlisPro⁶⁴ software packages. The crystal structure was refined with SHELXL.⁶⁶ Complete structural data were deposited in the CSD⁶⁷ with refcode CCDC 1437552.

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References

- 1 A. J. Cruz-Cabeza, S. M. Reutzel-Edens and J. Bernstein, *Chem. Soc. Rev.*, 2015, **44**, 8619.
- 2 M. A. Neumann, F. J. J. Leusen and J. Kendrick, *Angew. Chem., Int. Ed.*, 2008, **47**, 2427.
- 3 X.-Z. Li, M. Li, Z. Li, J.-Z. Hou, X.-C. Huang and D. Li, *Angew. Chem., Int. Ed.*, 2008, **47**, 6371.
- 4 S. A. Moggach, T. D. Bennett and A. K. Cheetham, *Angew. Chem., Int. Ed.*, 2009, **48**, 7087.
- 5 E. Patyk, J. Skumiel, M. Podsiadło and A. Katrusiak, *Angew. Chem., Int. Ed.*, 2012, **51**, 2146.
- 6 N. Marom, R. A. DiStasio, V. Atalla, S. Levchenko, A. M. Reilly, J. R. Chelikowsky, L. Leiserowitz and A. Tkatchenko, *Angew. Chem., Int. Ed.*, 2013, **52**, 6629.
- 7 A. Lemmerer, J. Bernstein, U. J. Griesser, V. Kahlenberg, D. M. Többsen, S. H. Lapidus, P. W. Stephens and C. Esterhuysen, *Chem. – Eur. J.*, 2011, **17**, 13445.
- 8 J. Bernstein, *Cryst. Growth Des.*, 2011, **11**, 632.
- 9 D. Singhal and W. Curatolo, *Adv. Drug Delivery Rev.*, 2004, **56**, 335.
- 10 R. J. Davey, S. L. M. Schroeder and J. H. ter Horst, *Angew. Chem., Int. Ed.*, 2013, **52**, 2167.
- 11 A. Gavezzotti, *J. Pharm. Sci.*, 2007, **96**, 2232.
- 12 J. Bernstein, R. J. Davey and J.-O. Henck, *Angew. Chem., Int. Ed.*, 1999, **38**, 3440.



- 13 D.-K. Bučar, R. W. Lancaster and J. Bernstein, *Angew. Chem., Int. Ed.*, 2015, **54**, 6972.
- 14 J. D. Dunitz and J. Bernstein, *Acc. Chem. Res.*, 1995, **28**, 193.
- 15 C. Shahar, S. Dutta, H. Weissman, L. J. W. Shimon, H. Ott and B. Rybtchinski, *Angew. Chem., Int. Ed.*, 2016, **55**, 179.
- 16 T. Seki, K. Sakurada and H. Ito, *Angew. Chem., Int. Ed.*, 2013, **52**, 12828.
- 17 M. J. Prakash, P. Raghavaiah, Y. S. R. Krishna and T. P. Radhakrishnan, *Angew. Chem., Int. Ed.*, 2008, **120**(21), 3969.
- 18 I. Weissbuch, V. Y. Torbeev, L. Leiserowitz and M. Lahav, *Angew. Chem., Int. Ed.*, 2005, **44**, 3226.
- 19 M. A. Neumann, J. van de Streek, F. P. A. Fabbiani, P. Hidber and O. Grassmann, *Nat. Commun.*, 2015, **6**, 7793.
- 20 F. P. A. Fabbiani, G. Buth, D. C. Levendis and A. J. Cruz-Cabeza, *Chem. Commun.*, 2014, **50**, 1817.
- 21 F. P. A. Fabbiani, D. C. Levendis, G. Buth, W. F. Kuhs, N. Shankland and H. Sowa, *CrystEngComm*, 2010, **12**, 2354.
- 22 F. P. A. Fabbiani, *NATO Sci. Peace Secur. Ser. B Phys. Biophys.*, 2010, 545.
- 23 I. D. H. Oswald, I. Chataigner, S. Elphick, F. P. A. Fabbiani, A. R. Lennie, J. Maddaluno, W. G. Marshall, T. J. Prior, C. R. Pulham and R. I. Smith, *CrystEngComm*, 2009, **11**, 359.
- 24 E. Patyk, M. Podsiadło and A. Katrusiak, *Cryst. Growth Des.*, 2015, **15**, 5670.
- 25 M. Aniola and A. Katrusiak, *Cryst. Growth Des.*, 2015, **15**, 764.
- 26 R. Gajda, A. Katrusiak and J. Crassous, *CrystEngComm*, 2009, **11**, 2668.
- 27 R. Lee, J. A. K. Howard, M. R. Probert and J. W. Steed, *Chem. Soc. Rev.*, 2014, **43**, 4300.
- 28 E. Boldyreva, *Cryst. Growth Des.*, 2007, **7**, 1662.
- 29 E. V. Boldyreva, S. N. Ivashevskaya and H. Sowa, *Z. Kristallogr.*, 2005, **220**, 50.
- 30 S. V. Goryainov, E. N. Kolesnik and E. V. Boldyreva, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2005, **357**, 340.
- 31 E. V. Boldyreva, S. N. Ivashevskaya, H. Sowa, H. Ahsbahs and H. P. Weber, *Dokl. Akad. Nauk*, 2004, **396**, 358.
- 32 E. Eikeland, M. K. Thomsen, S. R. Madsen, J. Overgaard, M. A. Spackman and B. B. Iversen, *Chem. – Eur. J.*, 2016, **22**(12), 4061–4069.
- 33 F. P. A. Fabbiani, D. R. Allan, W. I. F. David, A. J. Davidson, A. R. Lennie, S. Parsons, C. R. Pulham and J. E. Warren, *Cryst. Growth Des.*, 2007, **7**, 1115.
- 34 E. V. Boldyreva, V. Dmitriev and B. C. Hancock, *Int. J. Pharm.*, 2006, **327**, 51.
- 35 J. Wąsicki, D. P. Kozlenko, S. E. Pankov, P. Bilski, A. Pajzdarska, B. C. Hancock, A. Medek, W. Nawrocik and B. N. Savenko, *J. Pharm. Sci.*, 2009, **98**, 1426.
- 36 S. E. Kichanov, D. P. Kozlenko, J. Wąsicki, W. Nawrocik, L. S. Dubrovinsky, H.-P. Liermann, W. Morgenroth and B. N. Savenko, *J. Pharm. Sci.*, 2015, **104**, 81.
- 37 Y. V. Seryotkin, T. N. Drebuschak and E. V. Boldyreva, *Acta Crystallogr., Sect. B: Struct. Sci., Cryst. Eng. Mater.*, 2013, **69**, 77.
- 38 J. Ridout, L. S. Price, J. A. K. Howard and M. R. Probert, *Cryst. Growth Des.*, 2014, **14**, 3384.
- 39 B. A. Zakharov, N. A. Tumanov and E. V. Boldyreva, *CrystEngComm*, 2015, **17**, 2074.
- 40 M. Fisch, A. Lanza, E. Boldyreva, P. Macchi and N. Casati, *J. Phys. Chem. C*, 2015, **119**, 18611.
- 41 E. V. Boldyreva, T. P. Shakhshneider and H. Ahsbahs, *J. Therm. Anal. Calorim.*, 2002, **68**, 437.
- 42 N. A. Tumanov, E. V. Boldyreva, B. A. Kolesov, A. V. Kurnosov and R. Quesada Cabrera, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2010, **66**, 458.
- 43 A. P. Ayala, M. W. C. Caetano, S. B. Honorato, J. Mendes Filho, H. W. Siesler, S. N. Faudone, S. L. Cuffini, F. T. Martins, C. C. P. da Silva and J. Ellena, *J. Raman Spectrosc.*, 2012, **43**, 263.
- 44 A. J. Cruz-Cabeza and J. Bernstein, *Chem. Rev.*, 2014, **114**, 2170.
- 45 V. A. Drebuschak, T. N. Drebuschak, N. V. Chukanov and E. V. Boldyreva, *J. Therm. Anal. Calorim.*, 2008, **93**, 343.
- 46 T. N. Drebuschak, N. V. Chukanov and E. V. Boldyreva, *Acta Crystallogr., Sect. E: Struct. Rep. Online*, 2006, **62**, o4393.
- 47 T. N. Drebuschak, N. V. Chukanov and E. V. Boldyreva, *Acta Crystallogr., Sect. C: Cryst. Struct. Commun.*, 2007, **63**, o355.
- 48 T. N. Drebuschak, N. V. Chukanov and E. V. Boldyreva, *Acta Crystallogr., Sect. C: Cryst. Struct. Commun.*, 2008, **64**, o623.
- 49 T. N. Drebuschak, Y. A. Chesalov and E. V. Boldyreva, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2009, **65**, 770.
- 50 T. N. Drebuschak, V. A. Drebuschak and E. V. Boldyreva, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2011, **67**, 163.
- 51 T. N. Drebuschak, A. A. Ogienko and E. V. Boldyreva, *CrystEngComm*, 2011, **13**, 4405.
- 52 G. J. Piermarini, *J. Appl. Phys.*, 1973, **44**, 5377.
- 53 J. W. Otto, J. K. Vassiliou and G. Frommeyer, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1998, **57**, 3253.
- 54 J. R. Cox, L. A. Ferris and V. R. Thalladi, *Angew. Chem., Int. Ed.*, 2007, **46**, 4333.
- 55 A. Gavezzotti, *Molecular Aggregation, International Union of Crystallography Monographs on Crystallography*, Oxford University Press, 2006, p. 442.
- 56 A. Gavezzotti, *Cryst. Res. Technol.*, 2013, **48**, 793.
- 57 N. B. Leonidov, *Russ. Chem. J.*, 1997, **41**, 5.
- 58 N. B. Leonidov, P. M. Zorky and A. E. Masunov, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 1993, **49**, c185.
- 59 D. Gidalevitz, R. Feidenhans'l, S. Matlis, D.-M. Smilgies, M. J. Christensen and L. Leiserowitz, *Angew. Chem., Int. Ed. Engl.*, 1997, **36**, 955.
- 60 W. Ostwald, *Z. Phys. Chem.*, 1897, **22**, 289.
- 61 *Reactivity of Molecular Solids, Molecular Solid State Series*, ed. E. V. Boldyreva and V. V. Boldyrev, Wiley, Chichester, 1999.
- 62 R. Boehler, *Rev. Sci. Instrum.*, 2006, **77**, 115103.
- 63 G. J. Piermarini, S. Block, J. D. Barnett and R. A. Forman, *J. Appl. Phys.*, 1975, **46**, 2774.
- 64 *CrysAlisPro*, Agilent Technologies UK, Yarnton, Oxfordshire, UK, 2013.
- 65 V. Dyadkin, *SNBL Tool box*, Swiss Norwegian Beamline at ESRF, Grenoble, France, 2013.
- 66 G. M. Sheldrick, *Acta Crystallogr., Sect. C: Struct. Chem.*, 2015, **71**, 3.
- 67 F. H. Allen, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2002, **58**, 380.

