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Catheryn Jackson, Gregory B. Mckenna. The melting behavior of organic materials confined in porous solids. *Journal of Chemical Physics*, 1990, 93 (12), pp.9002-9011. 10.1063/1.459240 . hal-02557807

HAL Id: hal-02557807

<https://hal.science/hal-02557807>

Submitted on 29 Apr 2020

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The melting behavior of organic materials confined in porous solids

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(Received 6 July 1990; accepted 4 September 1990)

The solid-liquid phase transition temperatures and heats of fusion ΔH_f of nonpolar organic solids confined in the pores of controlled pore glasses were measured by differential scanning calorimetry. The pore diameters d were in the range of 40–730 Å and the organics studied were *cis*-decalin, *trans*-decalin, cyclohexane, benzene, chlorobenzene, naphthalene, and heptane. In accordance with previous reports on studies of primarily inorganic materials, the melting point of the pore solid $T(d)$ decreased with decreasing pore diameter. In addition, a large reduction in the bulk enthalpy of fusion ΔH_f of the pore solid was measured, which apparently has not been studied in detail by other workers. A linear correlation was found between the melting point depression (ΔT_m) and the reciprocal diameter, as predicted by theories of solidification in a capillary. The calculated values of the solid-liquid interfacial energy σ_{sl} were in reasonable agreement with values reported in the literature based on other methods of measurement.

INTRODUCTION

The behavior of liquids and solids in very small pores is relevant from both a fundamental and practical perspective. There is fundamental interest on the effect of finite size constraints on bulk properties. Crystalline melting,^{1–11} solid-solid⁵ and superfluid¹² phase transitions are observed to shift to lower temperatures for materials confined to small pores. Also, the behavior of liquid in small pores has importance in many commercial areas such as catalysis, interfacial adhesion, and separation science. Water has been widely examined because of practical interest in frost prevention in biological systems,¹³ agricultural products, soil,¹⁴ and building materials.¹⁵

We are concerned with the crystalline melting temperature T_m of organic solids formed in small pores of controlled pore glasses¹⁶ of varying pore diameter d . Although many earlier studies of the melting behavior of solids formed in small pores (pore solid) exist, the common feature is generally an estimate of ΔT_m at a single pore size, with no specific information of the effect of pore size on the melting behavior. In fact, very few studies of T_m as a function of pore size exist, since narrow pore size distribution materials of a sufficiently small pore size were not readily available until the mid-1970's. A brief review of the literature will be given to place the present study in proper perspective.

The lowered melting temperature of pore solids has frequently been ascribed to a very small crystal size in the pore^{6,14} and the large surface to volume ratio of material adsorbed in a capillary.¹⁴ The limitation of crystal size is known to lower the equilibrium melting point in other systems such as metal particles,^{17,18} finely dispersed nonporous powders,¹⁹ and polymer lamellae of different thicknesses.²⁰ Very large effects have been reported for gold particles of 50 Å size, which exhibit a ΔT_m of more than 500 K.¹⁷ One expected and important feature of melting in small systems is the "smearing out" of the transition as the particle size decreases. This is due to the diffuse nature of

the solid-liquid interface, which has been predicted by theory and shown by experiment for metal particles.²¹

The formation of small crystals is described theoretically in a classical discussion by Gibbs,²² and results from the effect of surface curvature on the equilibrium state of a pure substance.⁶ A related theory was derived by Thomson²³ for the effect of curvature on the vapor pressure of liquid droplets. This theory may be extended to small crystals to give the same equation as that of Gibbs, as noted by Defay *et al.*⁶ and the equation for the temperature shift of melting in confined geometries is often called the Gibbs-Thomson equation. The Kelvin equation in its original form was found to correctly describe the facile condensation of a vapor in the pores of a solid, i.e., capillary condensation.⁷ A number of similar equations exist in the literature which have been derived for crystals formed in cylindrical pores^{3,8,14} which specifically consider the contact angle between the liquid, solid and the wall of the cylindrical pore, which is very important if freezing behavior is of interest. For crystal melting behavior, however, it is assumed that the contact angle is 180° and the equation becomes identical with the Gibbs-Thomson equation, given below. We will reconsider the effect of the contact angle later in the Analysis section.

The Gibbs-Thomson equation predicts that the melting point depression ΔT_m for a small crystal of size d is given by

$$\Delta T_m = T_m - T_m(d) = 4\sigma_{sl}T_m/(d\Delta H_f\rho_s), \quad (1)$$

where σ_{sl} is the surface energy of the solid-liquid interface, T_m is the normal (bulk) melting point, $T_m(d)$ is the melting point of crystals of size d , ΔH_f is the bulk enthalpy of fusion (per g of material), and ρ_s is the density of the solid. Presented in this form, it is assumed that σ_{sl} is isotropic and that the crystal size is sufficiently large that the material retains its bulk properties for ΔH_f and ρ_s . The possible failure of these assumptions is considered in the Analysis section.

Experimental reports of the depression of the bulk melting temperature of solids confined to a small system size are widespread, but many questions remain unanswered. For example, early calorimetric measurements on water adsorbed in porous silica gel,² showed a broad melting range of 50 K and a reduction in the peak melting point of nearly 12 K. For benzene and naphthalene in the same system, however, an even larger reduction in the peak melting point (~ 40 K) was found and the differences could not be adequately explained by the authors.² In general, low molar mass inorganic liquids such as water,^{2,4,9–11} hydrogen,⁸ neon,⁸ and oxygen,^{3,5} have been studied more frequently than organic liquids.^{2,10}

Another effect which is not well understood is the relationship between small system size and the enthalpy of fusion ΔH_f of the solid formed in the pores. Conflicting results on measurements made for ΔH_f of water in silica gel and other porous media have been noted, with ΔH_f reported to be reduced,⁴ increased,² or unchanged¹¹ for the pore solid. The hysteresis between freezing and melting in the liquid–solid transition of liquids confined in porous media has been a topic of study because of the large degree of supercooling which occurs in liquids such as hydrogen,⁸ where no significant supercooling occurs in the bulk.

Part of the diversity in reports of the melting behavior may be due to the different nature of the many porous “host” materials employed in the studies, including silica gels,² porous sol-gel glasses,^{3,5} porous Vycor²⁴ glass,^{4,8–10} and controlled pore glasses (CPGs).¹¹ The different nature is manifested in the shape and breadth of the pore size distribution, surface roughness, and other features of these materials. One of the earliest materials examined was silica gel, made in a sol-gel process from a suspension of colloidal particles. The sol-gel process usually results in a very broad pore size distribution, although recent developments in sol-gel technology provide better control of the pore geometry, yielding samples with well-defined pore diameters.³ These materials are called “porous sol-gel glasses.” Vycor brand porous glass (Corning Glass²⁴ #7930, pore diameter ~ 50 Å) and controlled pore glasses (CPG Incorporated²⁴) are produced by a phase separation process which is fundamentally different from the sol-gel process.¹⁶ CPGs have the advantage of being available in a number of different pore diameters. The morphology of these materials is a network of cylindrical pores, rather than isolated pores with one end closed. This allows the assumption that the contact angle remains constant within the pore and minimizes changes in ΔT_m as the pore is filled. The pore size distributions of sol-gel glasses,³ porous Vycor glass and controlled pore glasses¹⁶ are quite narrow compared to silica gels, which simplifies the interpretation of the experimental data.

The availability of CPGs and sol-gel glasses of different pore diameters has led to recent studies which give attention to the melting behavior as a function of pore size. Rennie and Clifford¹¹ conducted a calorimetric study of water in CPG with pore diameters from 20–2500 Å. The results indicated that beyond a few monolayers, water exists essentially as bulk liquid in the pores and that the

melting of pore ice was well described by the Gibbs–Thomson equation. The authors also observed a direct proportionality between the amount of water determined calorimetrically and the total water present, indicating a negligible deviation in the heat of melting due to surface effects. This is in contrast to the earlier report of Litvan⁴ that ΔH_f for water was significantly reduced in porous Vycor glass and contrary to our finding for organic liquids as will be seen subsequently. In addition, Rennie and Clifford¹¹ calculated experimental values for σ_{sl} of water at each pore diameter using Eq. (1), and concluded that it was not a function of diameter. Another study³ utilized a picosecond optical technique to study molecular dynamics of liquid oxygen in porous sol-gel glasses having pore diameters which ranged from 40–400 Å. From these experiments a $1/d$ dependence of ΔT_m was also observed, however, no calorimetric data were available to investigate the enthalpy of fusion of the pore solid.

The present study investigates the melting behavior of a variety of organic solids in CPGs as a function of pore diameter. The melting point depression ΔT_m and the enthalpy of fusion ΔH_f were measured by differential scanning calorimetry (DSC). The CPG surfaces were derivatized to promote wetting by nonpolar, organic liquids. The organic liquids studied in detail were *cis*-decalin, *trans*-decalin, cyclohexane, benzene, chlorobenzene, naphthalene and heptane. Most of the data were obtained with a slight excess loading of the pores. Under this condition, the bulk melting temperature and the lower pore solid melting temperature were both observed, as previously reported for water, benzene, and naphthalene in silica gel,² and water in CPG.¹¹ This condition has the added feature of minimizing the effect of surface forces on the pore material, therefore emphasizing the small size and isolation of the organic crystal. The data were analyzed according to the prediction of a linear relationship between ΔT_m and $1/d$ given by Eq. (1). Values for σ_{sl} were calculated for each material and compared to empirical calculations and literature values based on other techniques, when available.

EXPERIMENTAL

Materials

The controlled pore glasses (CPG) used in this study were supplied by Dr. Wolfgang Haller of NIST.^{16,25} These materials are also available commercially from CPG Inc. in Fairfield, New Jersey. The CPG is a white, free-flowing powder of 120/200 mesh size. The mean pore diameter, the range of diameters which describes 90% of the pores, and the pore volume for each CPG were determined by mercury intrusion and are given in Table I. The surface area of the glass is also given in Table I, as determined by nitrogen adsorption using the Branauer–Emmett–Teller (BET) equation.

The surface of the CPG was derivatized with hexamethyldisilazane to convert the surface hydroxyl groups to trimethylsilyl groups. This treatment makes the glass more hydrophobic and promotes wetting by organic liquids. Each CPG was first cleaned by boiling in nitric acid on a

TABLE I. Controlled pore glass properties.

Mean pore diameter ^a , (Å)	± % ^b	Surface area ^c (m ² /g)	Pore volume ^a (cc/g)
39.5	12.8	144	0.13
85	13.5	175	0.59
156	5.7	166	0.90
255	3.7	95.5	0.96
324	4.2	79.0	1.39
486	3.9	43.3	0.92
729	8.6	33.9	0.98

^aDetermined by mercury intrusion method.^b90% of the pore diameters are within this range.^cDetermined by nitrogen adsorption method (BET equation).

steam bath for 10 h. The nitric acid was then decanted off and the CPG was rinsed copiously with distilled, deionized water, and dried thoroughly in an oven. The hexamethyldisilazane was then added to cover the glass, stirred with a spatula to release trapped air bubbles, and heated at 50–55° C for 20 h. Following this treatment, the reagent was decanted off and the CPG was rinsed well with chloroform and dried thoroughly in an oven. The CPGs were stored in a desiccator when not in use. It should be noted that the surface derivatization does not significantly effect the pore dimensions.²⁶

The organic solvents used in this study were obtained from the following manufacturers²⁴ and used without further purification: *cis*-decalin, *trans*-decalin, and hexadecane from Aldrich Chemical Company, *n*-heptane, cyclohexane, and naphthalene from Mallinckrodt, benzene from J. T. Baker, and chlorobenzene from Fisher Scientific.

Calorimetry

The Perkin–Elmer²⁴ DSC 2C used in this study was specially modified for low temperature work. The modification involved building a new heat sink attachment for the DSC head to allow a continuous flow of liquid nitrogen through the system. This keeps the block at a constant low temperature with minimal operator attention, and improves the stability of the baseline and calibration. The attachment was made from a cylindrical piece of brass (7.5 cm in diameter and 3.0 cm in height) with a 1.27 cm (1/2 in.) hole in the center to bolt it to the DSC head in the

usual manner. The brass piece was machined out to make two cylindrical chambers (around the center hole) separated by a thin sheet of brass. A hole was then drilled into the side of each chamber and fit with a 0.64 cm (1/4 in.) fitting to pipe the liquid nitrogen in and out of the attachment. Flow between the chambers was controlled with a 0.32 cm (1/8 in.) hole. The open end of each chamber was sealed with a piece of brass. The lowest starting temperature achievable with this attachment was ~140 K.

The DSC sample pans used in this study were large volume, stainless-steel pans sealed with an O-ring to prevent evaporation of the solvent (Perkin–Elmer #0319-0218). The method of sample preparation was to weigh the CPG into the preweighed sample pan, add the liquid (or solid in the case of naphthalene), seal the pan, and reweigh. The precision of the balance was ±0.01 mg. The concentrations reported in cm³/g are in cm³ of liquid, calculated using the liquid densities,²⁷ per g of glass. The CPG/liquid mixtures were equilibrated overnight at room temperature while the solid solvents were heated to ~90° C for a few hours to liquify the material so that it could flow into the pores. Before analysis the samples were reweighed to insure that the pans were sealed properly. In most cases, the reweighed samples were within ±0.05 mg. A few pans showed a large weight loss due to improper sealing and were discarded. The prepared sample pans were stable over many months for all of the mixtures except CPG/benzene, which showed a small weight loss with time due to diffusion of the benzene through the rubber O ring. The CPG/benzene mixtures were always weighed both before and after the sample run to assure that the weight was correct. No calorimetric effects due to the O ring were observed.

Calibration of the DSC was done with pure samples of the organic liquids. The heating rate used was 5 K/min and most of the data discussed in this paper was obtained on heating. A cooling rate of 5 K/min was used to prepare each sample for analysis. The normal melting point and enthalpy of fusion for each solvent determined by DSC, as well as the literature values, are given in Table II. At the relatively low heating rate used in this study, good agreement was observed between the experimental peak melting points and the literature melting points. The experimentally measured values of bulk ΔH_f are also listed in Table II

TABLE II. DSC calibration.

Substance	Literature values			Experimental (peak) values		
	T_m (K)	ΔH_f (cal/g)	Source (Ref.)	T_m (K)	ΔH_f (cal/g)	T_m^a (K)
<i>n</i> -heptane	182.4	33.78	27	180.2	30.2	181.0
Chlorobenzene	227.8	20.40	27	229.0	19.7	229.1
<i>c</i> -decalin	230.2	16.4	28	230.1	15.7	229.9
<i>t</i> -decalin	242.6	24.9	28	241.7	23.2	243.5
<i>n</i> -decane	243.3	48.34	27	244.1	47.3	244.7
Benzene	278.5	30.45	27	279.0	29.3	280.8
Cyclohexane	279.5	7.47	27			280.1
<i>o</i> -terphenyl	329.3	17.84	29	331.3	17.5	331.0
Naphthalene	353.4	35.06	27	357.0	34.9	356.1

^aAverage value for bulk melting peak in CPG/liquid mixtures where pore and bulk melting peaks are clearly resolved.

and agree satisfactorily with the literature values. In all cases, the experimentally measured value of the bulk ΔH_f was used for calculations involving the pore melting peak enthalpy. The final column in Table II lists the average peak melting temperature of the excess bulk solvent in CPG/solvent samples where the pore and bulk melting peaks were clearly resolved. Because of the presence of the CPG, this peak tended to be slightly broader than that of the pure solvent, but in general was still in good agreement with the expected melting temperature.

A minimum temperature range of 100 K was studied for each system, from 80 below to 20 K above the normal melting point. The exceptions were the low melting materials *n*-heptane and chlorobenzene, where a slightly smaller range had to be used because the minimum temperature attainable with the instrument was 140 K. In some cases a larger range was studied and no significant differences in the results were observed. Annealing of several of the samples for 3–4 h at low temperatures also did not significantly alter the observed melting temperature, the shape of the melting peak or ΔH_f .

The melting point depression ΔT_m was calculated in two different ways, depending on the resolution of the pore and bulk melting peaks. In the CPGs of smallest pore size, usually 40, 85, and 156 Å, the two peaks were well resolved and a direct measurement of ΔT_m was possible for each case. In the CPGs of larger pore size, the value of the bulk melting peak was taken as the last column in Table II. The broadened peak melting temperature (or a clearly resolved left-hand shoulder if visible) was then subtracted from this bulk melting temperature to obtain ΔT_m .

RESULTS AND ANALYSIS

Effect of pore fullness

The effect of pore fullness on ΔT_m and ΔH_f was examined for both *cis*-decalin and benzene. The 85 Å CPG was used for these studies, which clearly shows the separation between the pore-melting and bulk-melting temperatures, and allows an accurate measurement of ΔH_f . Typical normalized data for *cis*-decalin is shown in Fig. 1 for concentrations of 0.133–0.998 cm³/g. The peak melting temperatures of the pore solid and corresponding ΔH_f are given in Table III. At concentrations of 0.630 cm³/g and below, a single melting endotherm was observed at a peak melting temperature of 212–213 K, about 17–18 K below the normal melting point (230 K). At concentrations of ≥ 0.718 cm³/g and above, the bulk melting peak appeared at the normal melting point for *cis*-decalin, which is due to excess material which cannot fit in the pores. The expected pore volume for the 85 Å CPG (obtained by mercury intrusion data given in Table I) is 0.59 cm³/g, slightly lower than that observed experimentally for *cis*-decalin. This may be a result of different wetting characteristics of the two liquids on the glass surface. A small increase of ~ 1 K in $T_m(d)$ for *cis*-decalin is seen when the pores are completely filled, and the precision in the measurement is improved, as listed in Table III.

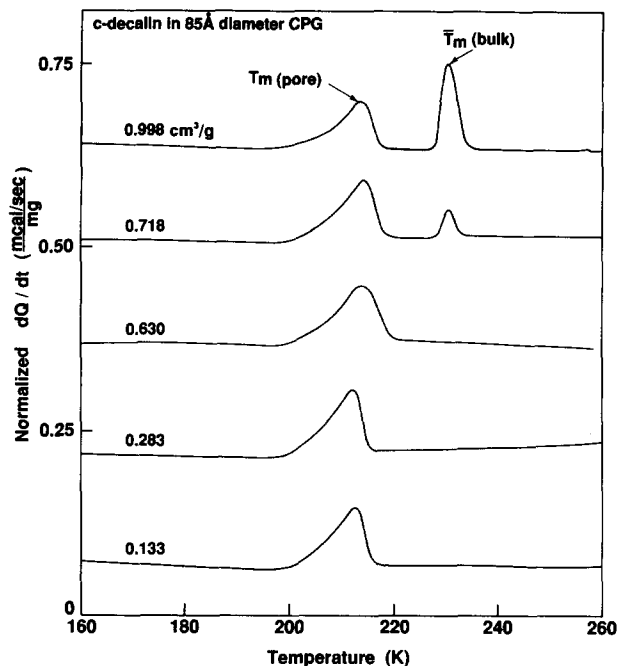


FIG. 1. Effect of pore fullness on the melting behavior: DSC heating curves of *c*-decalin in 85 Å pores of CPG (heating rate = 5 K/min). At 0.133–0.630 cm³/g, the pores are underfilled and only the pore solid melting endotherm appears, while at 0.718–0.998 cm³/g the pores are overfilled and a second melting endotherm appears due to the excess pore bulk solid. See text for further analysis.

Within the error of this measurement, the shape, melting temperature, and enthalpy of the pore melting peak shown in Fig. 1 for *cis*-decalin is independent of pore fullness. This suggests that the liquid prefers to condense as a plug in the center of a few pores rather than to wet the entire inside surface of the CPG uniformly, and that the glass surface is not playing a highly interactive role in these studies. If a uniform wetting of the interior surface were occurring, our results would more closely resemble the earlier study of water in untreated CPG by Rennie and Clifford,¹¹ in which water was undetectable at < 15 wt. % because of the presence of a few monolayers coating the entire surface before any pores were full. *Cis*-decalin was easily detected in the 85 Å CPG as low as 0.067 cm³/g, which corresponds to 5.7 wt. %. Since ΔH_f of *cis*-decalin is about one-fifth that of water, this represents a very great difference in the detection limit. The breadth of the pore melting endotherms shown in Fig. 1 are approximately 15 K, with the peak shape somewhat skewed toward higher temperature. The peak shape and breadth is indicative of the pore size distribution as discussed previously by Homshaw,³⁰ for water saturated ion-exchange resins and clays, and Brun *et al.*^{31,32} with regard to thermoporometry.

Similar results on the effect of pore fullness were found for benzene in the 85 Å CPG, shown in Fig. 2 for concentrations of 0.15–0.923 cm³/g. In this case, ΔT_m is ~ 14 –15 K. The shape of the pore melting peak for benzene at low pore filling (0.150–0.553 cm³/g) differs from that observed for *cis*-decalin, with a small shoulder on the right-hand side (high temperature side) of the peak. A bulk melting peak

TABLE III. Melting behavior of pore solid as a function of pore fullness in 85 Å diameter CPG.

Substance	Experimental condition	(cc/g)	$T_m(d)^a$ (K)	ΔH_f (cal/g)	
<i>c</i> -decalin	Underfilled pores	0.067	211.2	5.8	
		0.133	212.2	8.0	
		0.283	211.6	8.1	
		0.373	212.1	8.1	
		0.630	213.3	8.8	
	Overfilled pores		avg. ^b	212.3	8.3
			σ^c	0.7	0.4
		0.718	213.4	8.6	
		0.998	212.8	9.0	
	Benzene	Underfilled pores		avg. ^b	213.1
			σ^c	0.4	0.3
0.066			263.4	14.7	
0.150			266.7	18.6	
Overfilled pores		0.334	264.7	19.9	
		0.553	263.8	19.4	
			avg. ^b	265.1	19.3
			σ^c	1.5	0.7
	0.704	267.0	19.0		
	0.791	266.0	21.1		
	0.923	266.6	19.7		
	1.56	266.6	20.0		
1.76	266.1	19.5			
	avg. ^b	266.5	19.9		
	σ^c	0.4	0.8		

^aPeak melting temperature.^bAvg. = mean of data, for underfilled pores the lowest cc/g data is not included.^c σ = one standard deviation.

also appears at lower pore filling, perhaps due to the higher vapor pressure of benzene compared to *cis*-decalin and the possible transport of some of the pore liquid out of the CPG. The shoulder disappears when the pores are filled to slight excess (0.704–0.923 cm³/g). As noted previously for *cis*-decalin, a small increase of ~1K in $T_m(d)$ is measured with the pores completely filled, as compared to partially filled, and the precision in the measurement of $T_m(d)$ is increased (Table III). The breadth of the pore melting endotherm is similar for both compounds.

Although ΔH_f of the pore solid is fairly independent of pore fullness (>0.13 cm³/g) for both *cis*-decalin and benzene, it is greatly reduced compared to the bulk ΔH_f , as shown in Table III. For *cis*-decalin, the value of ΔH_f for the pore solid is ~8–9 cal/g compared to a bulk ΔH_f of 15.7 cal/g. For benzene, the reduction is slightly less but still significant. At the lowest pore filling, used to find the detection limit of the technique, an additional reduction in ΔH_f was measured. This may be caused in part by the error in measuring such a small peak. The reduction in ΔH_f of pore solid was found to vary with pore diameter (or crystal size) and will be considered in detail later.

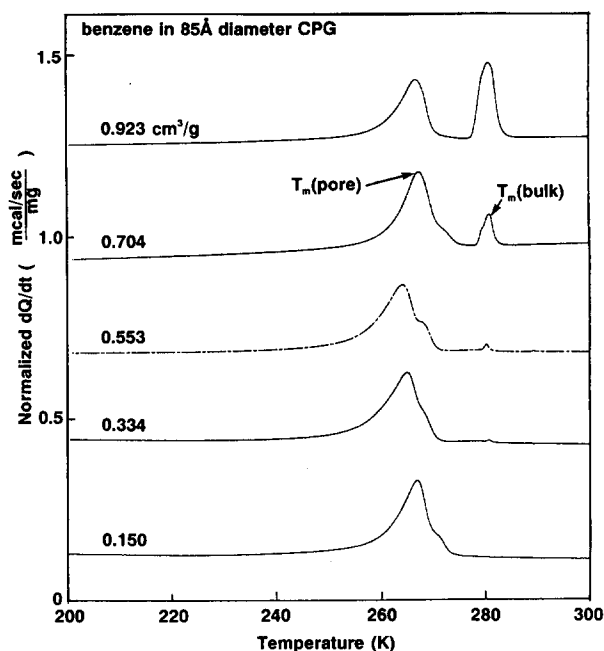


FIG. 2. Effect of pore fullness on the melting behavior: DSC heating curves of benzene in 85 Å pores of CPG (heating rate = 5K/min). At 0.150–0.553 cm³/g the pores are underfilled and at 0.704–0.923 cm³/g the pores are filled to excess, as described in Fig. 1.

Effect of pore diameter on ΔT_m

Studies of the effect of pore fullness allowed us to establish appropriate conditions to study the effect of pore diameter on ΔT_m for a number of different organic materials. Because a slight excess filling of the pores provides an internal standard for the bulk melting temperature, we chose this condition to obtain most of the data described here. Direct measurement of ΔT_m is then possible when a large enough separation of the pore and bulk melting peaks occurs. One set of experimental DSC data will be given for benzene as a function of pore diameter to illustrate ΔT_m and ΔH_f ; similar data obtained for the remaining compounds will then be summarized graphically.

The effect of CPG pore diameter on the melting temperature of the benzene frozen in the pores is shown in Fig. 3. In the 40 Å pores, ΔT_m is ~26–27 K, based on the peak melting temperatures, and the breadth of the peak is ~45 K. The broader nature of this peak may be partially due to the broader CPG pore size distribution at the 40 Å size (Table I) and to the smearing effect of the liquid–solid transition at small particle size.²¹ The pore solid melting endotherm moves to higher temperatures as the pore diameter is increased to 85 and 156 Å, and eventually merges with the bulk (excess pore) melting peak. A typical DSC trace for pure, bulk benzene is shown at the bottom of Fig. 3 for reference.

Similar calorimetric data were obtained for *cis*-decalin, *trans*-decalin, cyclohexane, chlorobenzene, naphthalene, and heptane in the investigation of the effect of molecular size on ΔT_m and ΔH_f . The results for ΔT_m are summarized in Fig. 4, where ΔT_m is plotted versus $1/d$. In general, a linear relationship is observed, in qualitative agreement

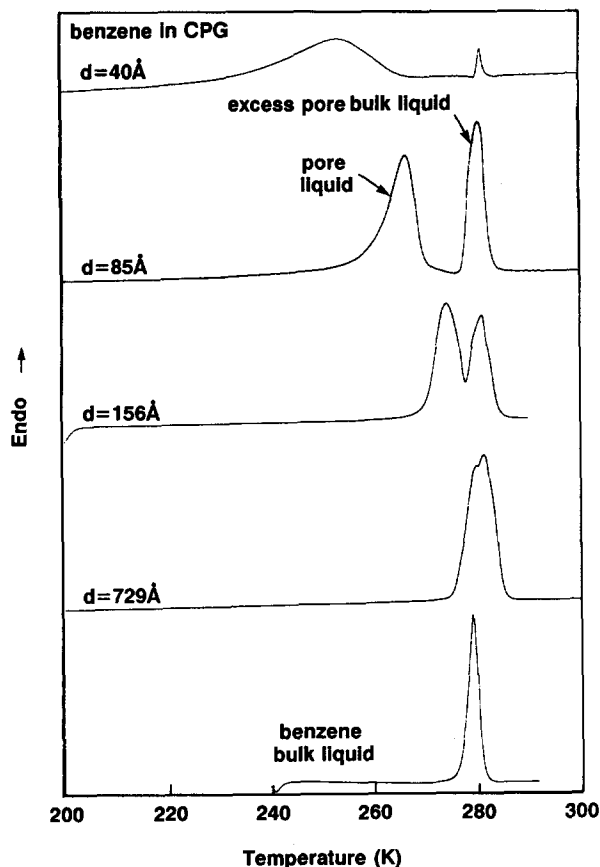


FIG. 3. Effect of pore diameter on the melting behavior: DSC heating curves of benzene in pore diameters of 40, 85, 156 and 729 Å CPG and compared to pure benzene standard (heating rate 5 K/min). The pores are filled to slight excess to show ΔT_m directly.

with Eq. (1). These results and the evaluation of σ_{sl} will be discussed in the Analysis section.

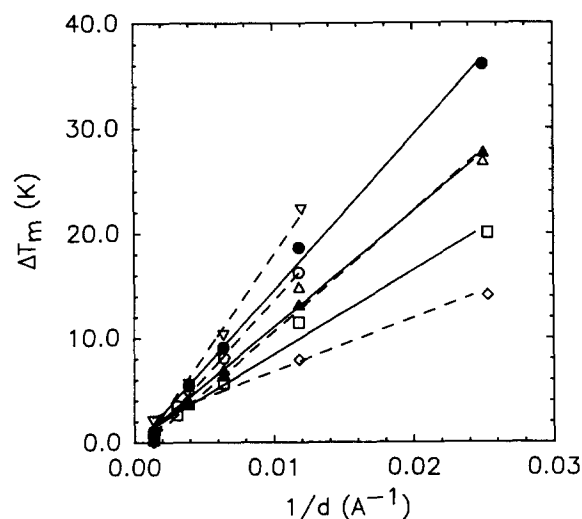


FIG. 4. Experimental values of ΔT_m plotted as a function of the reciprocal pore diameter for (○) *cis*-decalin, (●) *trans*-decalin, (▽) cyclohexane, (△) benzene, (▲) chlorobenzene, (◆) naphthalene and (□) heptane. The lines through the data are linear regressions fits used to calculate values of σ_{sl} . Melting endotherms were not observed in the smallest pores (40 Å) for cyclohexane and *cis*-decalin, see text for details.

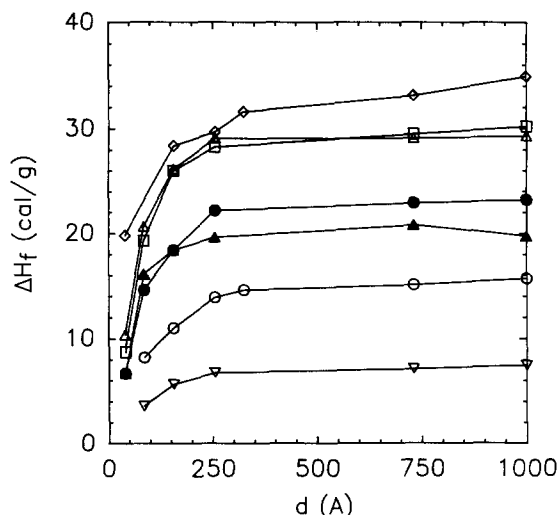


FIG. 5. Experimental values of ΔH_f of the pore solid vs d , showing a large reduction in the bulk ΔH_f (plotted at 1000 Å for reference) in the smallest pores. Symbols are the same as given in Fig. 4 caption.

In the case of the *cis*-decalin and cyclohexane in the smallest diameter pores (40 Å), no melting endotherm was detected down to 145 K.³³ Assuming the $1/d$ dependence of ΔT_m (Fig. 4) to be valid, T_m at 40 Å is expected at about 205 K for *c*-decalin and 233 K for cyclohexane, well within the range of study. The lack of a melting endotherm may be due to the rather low bulk ΔH_f of these compounds compared to the other organics studied (Table II) in combination with the reduction of the bulk ΔH_f observed in the smallest pores, to be discussed in the next section. The possibility that these two liquids might not flow into the smallest pores was discarded based on the fact that a reduced enthalpy was measured for the bulk (excess liquid) melting peak. In fact, the calculation of the pore volume for *cis*-decalin and cyclohexane in the 40 Å pores (from the measurement of the ΔH_f of the excess liquid) gave similar results to the other compounds studied. Annealing at low temperature for 3–4 h also did not change the result. This suggests that these liquids may be stable in a highly undercooled state in the 40 Å pores, although a greater than $1/d$ dependence at this small size may also explain the behavior. We also note that no evidence was obtained which suggests vitrification of these two systems.

Effect of pore diameter on ΔH_f

The effect of pore diameter on the measured value of ΔH_f at small pore sizes is dramatic, as shown in Fig. 5. The bulk value is plotted at a diameter of 1000 Å, for reference. The observed ratio of $\Delta H_f(\text{pore})/\Delta H_f(\text{bulk})$ at the 40 Å pore diameter ranges from 0.28 for heptane to 0.57 for naphthalene. This may be the reason a pore melting endotherm is not detected in *cis*-decalin and cyclohexane in the 40 Å pores. In the 85 Å pores, these materials show low ratios of $\Delta H_f(\text{pore})/\Delta H_f(\text{bulk})$ of ~ 0.5 , compared to 0.65–0.88 for the other materials. Although there is some error in the calculated pore melting enthalpy due to the breadth of the peak at small pore size, the error is not large

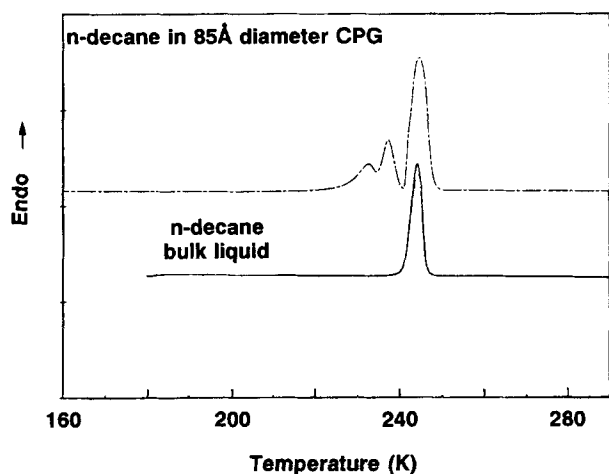


FIG. 6. Anomalous melting behavior of *n*-decane: DSC heating curves of *n*-decane in 85 Å CPG (heating rate 5 K/min). A scan of pure decane is shown for reference, see the text for details.

enough to account for such a large reduction in ΔH_f . A small correction for the temperature dependence of ΔH_f is also not enough to account for the large reductions in ΔH_f at the smallest pore sizes.⁷

Many explanations for the reduction in the bulk ΔH_f in the small pores are possible, from both an experimental and theoretical viewpoint. Experimentally, if a liquid layer remains unfrozen on the surface of the glass, all of the material cannot be expected to contribute to the crystalline melting endotherm. If this were the case, however, it is surprising that the proportionality between ΔT_m and $1/d$ continues at small pore diameter. Related to this is the possible smearing out of the liquid–solid transition as the pore size decreases, where the interface between liquid and solid is large. From a theoretical standpoint, it may be that at very small size, the surface to volume ratio of a crystal becomes important and the bulk properties are affected.

Anomalous melting behavior of some compounds

All of the data discussed thus far for the pore melting of various organic compounds were well behaved and appeared essentially as shown in Fig. 3 for the example of benzene. For some other compounds this was not the case, and may be an important clue to the behavior of certain materials in confined geometries. In the selection of compounds for this work, we attempted to include a series of *n*-alkanes to complement the data on ring systems. Heptane ($n = 7$) was the first member of this series, since the melting points of the lower alkanes were below the temperature limit of 140 K for our DSC. Heptane behaved similarly to benzene, described above. The next *n*-alkane studied, *n*-decane ($n = 10$), gave a bimodal pore melting peak shown in Fig. 6, for the 85 Å pores. This was not due to any impurity in the decane, since the pure material gave a single sharp melting endotherm.

Although we do not know the origin of this behavior, diffraction studies on the crystal structure of the pore solid may prove interesting. Neutron diffraction has recently

been shown to be a viable technique to study the structure of deuterium oxide³⁴ and oxygen³⁵ solidified in porous glasses, a rather surprising result considering the small scale of the pores. This suggests that the crystallite size parallel to the cylindrical pore is sufficiently large to yield distinct diffraction peaks and useful information about the effect of confining geometry on crystal structure in the pores. For some *n*-alkanes, multiple solid crystal forms have been reported³⁶ and may be relevant to the anomalous melting behavior observed.

ANALYSIS

In our study of the melting temperature of small organic crystals formed in the pores of CPGs, we analyze the data based on the prediction of a $1/d$ dependence of ΔT_m according to Eq. (1), and calculate a value of σ_{sl} from the slope of the plot of ΔT_m vs $1/d$ (Fig. 4). The necessary assumptions are the same as those adopted previously by Gibbs, namely that the values of σ_{sl} , ΔH_f and ρ_s are independent of pore size (or crystallite size) and are represented by the bulk values. In addition, it is assumed that the pore diameter is equal to the crystal diameter, although this is actually an upper limit on the crystal diameter. The implicit assumption is that the crystal structure or unit cell of the pore solid is the same as the bulk material. We are aware that a number of other factors might be important, such as surface roughness, transport between the pores and the bulk, and the possibility of a liquid monolayer on the pore surface which might reduce the measured value of ΔH_f and alter the effective pore diameter. In addition, if growing a crystal in a small pore results in a different crystal structure, all the thermodynamic parameters would change.

The values of σ_{sl} calculated from the slopes are given in the first column of Table IV. The first entries were calculated without the smallest pore data, which showed the largest reduction in ΔH_f and are most likely to deviate from the equation due to the small scale. The values calculated including the small pore data are given in parentheses. (Because we have been unable to find values for the solid densities of *cis*-decalin and *trans*-decalin, they were calculated by assuming the solid densities are 1.18 times the liquid density;²⁷ the solid densities of the other materials were found in the literature.³⁷) Also given in Table IV are values of σ_{sl} calculated from empirical relations,^{38–40} or literature values reported from different methods of measurement, when available. The empirical relation used was given by Dunning³⁸ based on experimental data of homogeneous nucleation experiments from the melt; where the molar ratio of $\sigma_{sl}/\Delta H_f$ per unit area was approximately 0.33 for a number of organics. This was near the lower limit expected for this ratio. Turnbull³⁹ gives an empirical estimate of this range as 0.32–0.45 and theoretical calculations⁴⁰ have predicted a value of 0.45.

We first compare the empirically calculated values of σ_{sl} to our values calculated from the experimental data. In general, the experimentally calculated values from the depression of the melting point (DMP) method used in this

TABLE IV. Comparison of experimental, empirically calculated, and literature values of σ_{sl} (erg/cm²).

Substance	Experimental calculation		Empirical calculation ^b	Literature values		
	from slope of Eq. (1) ^a			σ_{sl}	Method ^c	Reference
Benzene	15.7	(12.9)	21.5	44 ± 10	GBG	41
				22 ± 2	C	41
				19.8	HN	41
				21.4	HN	38
Chlorobenzene	14.1	(13.3)	18.4			
Cyclohexane	4.6	...	4.8			
Heptane	17.1	(13.5)	23.3			
Naphthalene	8.2	(6.1)	31.7	61 ± 11	GBG	41
				27.2	HN	41
				30.1	HN	38
<i>cis</i> -decalin	11.6	...	14.4			
<i>trans</i> -decalin	18.4	(16.2)	21.4			

^aNumbers in parentheses are obtained including the data from 40 Å pores.

^bCalculated from the empirical relation: $\sigma_{sl,M}/\Delta H_{f,M}$ (Refs. 38 and 39).

^cGBG = grain boundary groove; C = conical capillary; and HN = homogeneous nucleation.

work are ~25% lower than the empirically calculated values (Table IV). Naphthalene is the notable exception, where a ~75% lower value of σ_{sl} was obtained. The low values cannot be easily explained based on the direct application of Eq. (1) to the data obtained in this study. For example, if the lowered ΔH_f observed in the small pores is taken into account, the calculated value of σ_{sl} would be further reduced. Similarly, reduced values of the bulk ρ_s , due to poor packing, would also lower σ_{sl} . The failure of the assumption that the pore diameter is equal to the crystal diameter, also does not account for the small value of σ_{sl} . If a few monolayers of liquid on the surface of the pore do not freeze, the effective pore diameter would be smaller and inclusion of such a correction in Eq. (1) would further reduce the calculated value of σ_{sl} . One factor which is not explicitly given in Eq. (1) is the equilibrium contact angle between the solid-liquid and solid-substrate interfaces, which is important for freezing studies and determines whether the solid nucleus forms on the substrate (wall) or in the bulk.¹⁴ We have assumed that the contact angle is 180° to obtain Eq. (1), as discussed in the Introduction, which implies that the solid is separated from the substrate by a layer of liquid. If the contact angle is actually somewhat less than 180°, this would increase the calculated value of σ_{sl} . We note that the especially low value of σ_{sl} obtained for naphthalene may be due to the large anisotropy reported for σ_{sl} of this compound (20%).⁴¹ It is also possible that changes in the crystallographic form occur when naphthalene is crystallized in a constrained geometry, and may be an interesting subject of future study.

A comparison of the values of σ_{sl} calculated here with those obtained by other experimental methods is next considered. Our values are generally comparable to values obtained by the methods of homogeneous nucleation (HN) and conical capillary (C), but low compared to the grain boundary groove (CBG) method, within the large variability of such measurements reported in the literature. The inherent difficulties of determining reliable values of σ_{sl} have been reviewed by Bilgram,²¹ Jones,⁴¹ and Woodruff.⁴² The most common methods are HN and GBG

experiments. In the case of HN, widely used for both metals and organics, the most serious questions arise from the uncertainty of applying macroscopic thermodynamics to systems as small as the solid nucleus (typically 10 Å in size), the question of trace impurities which would result in heterogeneous rather than homogeneous nucleation, and the fact that in a nucleation experiment, σ_{sl} is determined as much as 200 K below T_m without consideration of a temperature coefficient for σ_{sl} .⁴¹ The GBG method is also called the method of interface intersections, and the measured angles of intersection can be used to derive a value of σ_{sl} . The GBG method is very difficult to apply to pure materials that are opaque, such as metals, and has more widespread use in the study of metal alloys.⁴² The conical capillary (C) method was one of the earliest to measure σ_{sl} from the direct application of the Gibbs-Thomson equation, but has not been used subsequently. This is due to practical disadvantages, such as the extremely good temperature control required and lack of suitability for impure systems.

The method of obtaining σ_{sl} from measurements of DMP of small crystals used in this work is similar to HN in size scale and may suffer criticisms in this regard, although trace impurities and temperature dependence of σ_{sl} should be much less important. As mentioned earlier, narrow pore size distribution materials of sufficiently small pore diameters have become readily available in recent years, and the method of DMP should be reconsidered as a method of determining σ_{sl} . It may be at least as reliable and easier to use than the HN method. This has also been noted by the workers employing thermoporimetry.³⁰

Some mention should be made to the possible relation of this work to cluster theory,⁴³ for although the geometry is different, the size scale is similar. The melting point of microcrystals, or clusters, should clearly demonstrate the manner in which the properties approach those of the bulk phase as the size of the particles increase, although more work in this area is needed.⁴⁴ Computer simulations indicate that the melting transition of Lennard-Jones microcrystals confined to spherical cavities is related to specific

geometrical factors. The Lennard-Jones clusters are predicted to have a melting temperature that is directly related to the number of atoms in the cluster. The equation for the melting point depression ΔT_m^* (where * denotes quantities reduced by the Lennard-Jones parameters) is

$$\Delta T_m^* = AN^{-1/3} - BN^{-2/3}, \quad (2)$$

where A and B are unknown constants and N is the number of atoms in the cluster. This equation is derived for spherical particles so the assumption, $R_s \propto N^{-1/3}$ is made, and the leading term in Eq. (2) has the same inverse radius dependence as predicted by Eq. (1). However, at very small radius (small N) the $N^{-2/3}$ term becomes significant and Eq. (2) predicts a dependence of ΔT_m which differs from $1/d$. Rather it predicts that $\Delta T_m \propto 1/d^2$. This would not be supported from our data, which include sizes to 40 Å. Alternatively, that continuum equations apply to both organics and metals¹⁸ at the size scales studied implies that bulk properties are still valid. This is in accord with the work⁴⁵ of Lessen, Asher and Brucat on vanadium argon clusters with n atoms, VAr_n^+ , where bulk behavior is approached with just $n = 7$. One may object that the large reduction in ΔH_f reported in this work is counter to the argument of bulk behavior, but at this time we do not know whether the reduction of ΔH_f is due to the diffuse nature of the solid-liquid interface, the presence of a liquid layer, or an actual reduction in the enthalpy of fusion of a very small crystal. Further experimental work using techniques such as scattering and nuclear magnetic resonance might help resolve these questions.

SUMMARY

Studies on the effect of finite size on the bulk properties of materials confined to very small pores is a fundamental bridge of understanding between the molecular scale and its relationship to the bulk scale. In this work, we have studied the crystalline melting transition and melting enthalpy of a series of organic solids confined to small pores, and determined the effect of pore diameter on the magnitude of the melting point depression and the apparent enthalpy of fusion ΔH_f . The porous media used in this work were controlled pore glasses with well characterized and narrow pore size distributions, and pore diameters in the range of 40–730 Å. These unique materials provide a carefully controlled environment in which to grow small organic crystals of specific diameters.

The melting point depression ΔT_m was found to be an inverse linear function of the crystal diameter d in accordance with the continuum theory of Eq. (1). A few earlier studies on ΔT_m as a function of pore diameter for small inorganic molecules such as water¹¹ and oxygen⁵ have been previously reported to have similar ΔT_m vs $1/d$ dependences, but in general, controlled studies on liquids confined in pores of different diameter are rare. Quantitative application of Eq. (1) gave values of the solid-liquid interfacial energy σ_{sl} , which are similar to, but somewhat lower than, those obtained by homogeneous nucleation methods. This may be due to the unknown factor of the contact angle at the solid-liquid interface. Naphthalene

gave an anomalously low value by this method and possible explanations, such as anisotropy of σ_{sl} , were considered.

A unique feature of this work is the quantitative measure of the enthalpy of fusion ΔH_f of the solid in the pores as a function of pore diameter. The values of ΔH_f were found to be a function of pore diameter and greatly reduced (40–70%) in the smallest pores. For some compounds (*cis*-decalin and cyclohexane) no melting endotherm was detected in the 40 Å pores even well below the expected melting temperature. Possible explanations for this behavior were offered, such as the low enthalpy of fusion of these two materials (relative to the other materials studied) and the possible stability of the undercooled liquid in the confined geometry of the pores. Relevant to this may be studies on the liquid-glass transition of organic liquids confined in small pores, which we report in a set of related experiments.⁴⁶

This work is part of an ongoing study of the freezing and melting behavior of organic solvents in thermoreversible⁴⁷ and cross-linked^{48,49} polymer gels. Under certain conditions, the mesh-like, network structure of polymer chains forces the solvent to stay “compartmentalized,”¹³ so that upon freezing, only very small solvent crystals can form. The melting behavior then may mimic that observed in CPG in some regards, and we hope to relate the data of these two studies.

ACKNOWLEDGMENTS

The authors are indebted to Dr. Wolfgang Haller of NIST for providing the samples of controlled pore glass and instruction on how to best derivatize the surfaces. We would also like to thank Dr. Jack Douglas of NIST for his many helpful comments on the manuscript. The low-temperature modification of the DSC was made by Willard Roberts of the Center for Building Technology at NIST, and use of that equipment is gratefully acknowledged.

¹S. Brunauer, *The Adsorption of Gases and Vapors, Vol. I, Physical Adsorption* (Princeton University, Princeton, 1943), p. 444.

²W. A. Patrick and W. A. Kemper, *J. Phys. Chem.* **42**, 369 (1938).

³D. D. Awschalom and J. Warnock, in *Molecular Dynamics in Restricted Geometries*, edited by J. Klafter and J. M. Drake (Wiley, New York, 1989), Chap. 12.

⁴G. G. Litvan, *Can. J. Chem.* **44**, 2617 (1966).

⁵D. D. Awschalom and J. Warnock, *Phys. Rev. B* **35**, 6774 (1987).

⁶R. Defay, I. Prigogine, A. Bellemans, and D. H. Everett, *Surface Tension and Adsorption* (Wiley, New York, 1966).

⁷B. R. Puri, D. D. Singh, and Y. P. Myer, *Trans. Faraday Soc.* **53**, 530 (1957).

⁸J. L. Tell and H. J. Maris, *Phys. Rev. B* **28**, 5122 (1983).

⁹B. V. Enustun, H. S. Senturk, and O. Yurdakul, *J. Coll. Int. Sci.* **65**, 509 (1978).

¹⁰C. Hodgson and R. McIntosh, *Can. J. Chem.* **38**, 958 (1960).

¹¹G. K. Rennie and J. Clifford, *J. Chem. Soc. F1* **73**, 680 (1977).

¹²K. Shirahama, M. Kubota, S. Ogawa, N. Wada, and T. Watanabe, *Phys. Rev. Lett.* **64**, 1541 (1990).

¹³N. Murase, K. Gonda, and T. Watanabe, *J. Phys. Chem.* **90**, 5420 (1986), and reference cited therein.

¹⁴K. A. Jackson and B. Chalmers, *J. Appl. Phys.* **29**, 1178 (1958).

¹⁵R. F. Feldman and H. Cheng-yi, *Cement Concrete Res.* **15**, 765 (1985).

¹⁶W. Haller, *Nature* **206**, 693 (1965).

¹⁷Ph. Buffat and J.-P. Borel, *Phys. Rev. A* **13**, 2287 (1976).

¹⁸P. R. Couchman and W. A. Jesser, *Nature* **269**, 481 (1977).

- ¹⁹ F. Ehrburger and J. Lahage, *Colloids Surf.* **23**, 105 (1987).
- ²⁰ J. D. Hoffman, G. T. Davis, and J. I. Lauritzen, in *Treatise on Solid-State Chemistry*, edited by N. B. Hannay (Plenum, New York, 1976).
- ²¹ J. H. Bilgram, *Phys. Rep.* **153**, 1 (1987).
- ²² J. W. Gibbs, *Collected Works* (New York, 1928).
- ²³ W. Thomson (Lord Kelvin), *Philos. Mag.* **42**, 448 (1871).
- ²⁴ Certain commercial companies are named in order to specify adequately the experimental procedure. This in no way implies endorsement or recommendation by NIST.
- ²⁵ W. Haller, in *Solid Phase Biochemistry*, edited by W. H. Scouten (Wiley, New York, 1983), Ch. 11.
- ²⁶ B. W. Sands, Y. S. Kim, and J. L. Bass, *J. Chrom.* **360**, 353 (1986).
- ²⁷ *Handbook of Chemistry and Physics*, 56th ed. edited by R. C. Weast (Chemical Rubber Co., Cleveland, 1975).
- ²⁸ J. P. McCullough, H. L. Finke, J. F. Messerly, S. S. Todd, T. C. Kinchelae, and G. Waddington, *J. Chem. Phys.* **61**, 1105 (1957).
- ²⁹ S. S. Chang and A. B. Bestul, *J. Chem. Phys.* **56**, 503 (1972).
- ³⁰ L. G. Homshaw, *J. Coll. Int. Sci.* **84**, 141 (1981).
- ³¹ M. Brun, A. Lallemand, J.-F. Quinson, and C. Eyraud, *Thermochim. Acta* **21**, 59 (1977).
- ³² J.-F. Quinson, M. Brun, R. Spitz, and M. Bartholin, *Makromol. Chem.* **185**, 1105 (1984).
- ³³ The cyclohexane lower limit is ~ 190 K because the pore melting peak was not observed below the solid-solid phase transition observed for this compound at 186 K [R. Kahn, R. Fourme, D. Andre, and M. Renaud, *Acta Cryst. B* **29**, 131 (1973)].
- ³⁴ M. Dunn, J. C. Dore, and P. Chieux, *J. Cryst. Growth* **92**, 233 (1988).
- ³⁵ P. E. Sokol, K. W. Herwig, and M. Pang, *Bull. APS* **35**, 442 (1990).
- ³⁶ E. F. Westrum and J. P. McCullough, in *Physics and Chemistry of the Organic Solid State*, edited by D. Fox *et al.* (Interscience, New York, 1963), Chap. 1.
- ³⁷ *Crystal Data Determinative Tables*, 3rd. ed., edited by J. D. H. Donnay and H. M. Ondik (U. S. Department of Commerce (NBS) and JCPDS, Washington, D. C., 1972), Vol. 1.
- ³⁸ W. J. Dunning, in *Physics and Chemistry of the Organic Solid State*, edited by D. Fox *et al.* (Interscience, New York, 1963), Chap. 7.
- ³⁹ D. Turnbull, *J. Appl. Phys.* **21**, 1022 (1950).
- ⁴⁰ J. Q. Broughton and G. H. Gilmer, *J. Chem. Phys.* **84**, 5759 (1986).
- ⁴¹ D. R. H. Jones, *J. Mater. Sci.* **9**, 1 (1974).
- ⁴² D. P. Woodruff, *The Solid-Liquid Interface* (Cambridge University, Oxford, 1973), Chap. 2.
- ⁴³ N. Quirke, *Mol. Simul.* **1**, 249 (1988).
- ⁴⁴ D. J. Wales and R. S. Berry, *J. Chem. Phys.* **92**, 4473 (1990).
- ⁴⁵ D. Lessen, R. L. Asher, and P. J. Brucat, *Int. J. Mass Spectrosc. Ion Process.* (in press).
- ⁴⁶ C. L. Jackson and G. B. McKenna, *J. Non-Cryst. Solids* (in press).
- ⁴⁷ C. L. Jackson and G. B. McKenna, *Am. Chem. Soc.: Polym. Preprints* **31**(2), 607 (1990).
- ⁴⁸ C. L. Jackson and G. B. McKenna, *Rubber Chem. Tech.* (in press).
- ⁴⁹ B. B. Boonstra, F. A. Heckman, and G. L. Taylor, *J. Appl. Polym. Sci.* **12**, 223 (1968).