Melting temperatures deduced from molar volumes: a consequence of the combination of enthalpy/entropy compensation with linear cohesive free-energy densities†

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Enthalpy/entropy compensation is a general issue of intermolecular binding processes when the interaction between the partners can be roughly modelled with a single harmonic potential. Whereas linear H/S correlations are wished for by experimentalists, and often graphically justified, no inflexible law of thermodynamics supports the latter statement. On the contrary, the non-directional Ford’s approach suggests logarithmic H/S relationships, which can be linearized only over narrow enthalpy/entropy ranges. Predictions covering larger domains require mathematical mapping obeying specific boundary conditions which are not compatible with linear plots. The analysis of solvent-free melting processes operating in six different classes of organic and inorganic materials shows that reciprocal Hill plots are acceptable functions for correlating melting enthalpies and entropies. The combination of H/S compensation with the observed linear dependence of the cohesive free energy densities with respect to the melting temperature eventually provides an unprecedented interdependence between melting temperatures and molar volumes. This procedure is exploited for the prediction of melting temperatures in substituted cyanobiphenyls.

Introduction

A rational control and programming of the various contributions to the changes in standard thermodynamic free energy \( \Delta G_{\text{asso}} \), enthalpy \( \Delta H_{\text{asso}} \), and entropy \( \Delta S_{\text{asso}} \) of association accompanying the binding process depicted in eqn (1) is at the heart of self-assemblies occurring in biology,1 in physics2 and in chemistry† (|| refers to the activities of various partners).

\[
A + B \iff [AB] \quad \Delta G_{\text{asso}} = \Delta H_{\text{asso}} - T \Delta S_{\text{asso}} = -RT \ln \left( \frac{[AB]}{[A][B]} \right) 
\] (1)

Beyond the theoretically justified Gibbs free energy relationship (eqn (1), right-hand side),† the simple modelling of the interaction operating between the partners in the [AB] pair by using a harmonic potential (i.e. a spring) suggests that the stronger the binding energy (measured as the potential well depth \( u_{\text{min}} \propto \Delta H_{\text{asso}} \)), the larger its force constant \( k \) (measured as its mean vibrational frequency \( \nu = (1/2\pi) \sqrt{k/m} \propto -\Delta S_{\text{asso}} \) where \( m_r \) is the reduced mass of the harmonic oscillator).3 Consequently, enthalpy and entropy changes are usually correlated for a simple intermolecular binding event with a propensity for compensation, i.e. \( \Delta G_{\text{asso}} \) and \( \Delta H_{\text{asso}} \) concomitantly decrease or increase.† Recurrent, but empirical observations collected from series of association processes investigated in biology, in physics and in chemistry show that minor structural perturbations of the two partners lead to apparent linear correlations (eqn (2), left-hand side). The slope \( \alpha \) of the \( H_i \) versus \( \Delta S_i \) plot has Kelvin units and is often referred to as the compensation temperature \( T_{\text{comp}} \), i.e. the temperature at which all association processes \( i \) within a homogeneous series display the same compensation free energy change \( \Delta G_{\text{comp}} = \gamma \) (eqn (2), center). Its re-writing at the right-hand side of eqn (2) is reminiscent of the Gibbs relationship, which may explain the rather common belief in the statement that linear enthalpy/entropy compensation corresponds to a ‘fourth law’ of thermodynamics.6,7

\[
\Delta H_i = \alpha \Delta S_i + \gamma = \frac{T_{\text{comp}}}{\Delta S_i} + \Delta G_{\text{comp}} \Rightarrow \Delta G_{\text{comp}} = \Delta H_i - T_{\text{comp}} \Delta S_i 
\] (2)

Theoretical approaches based on statistical thermodynamics include both attempts to refute the relevance of enthalpy–entropy correlation,8 as well as to establish its signature as the result of (i) minor perturbations of equilibrium constants,9 (ii) partition functions governed by a Gaussian density of states8b or (iii) underlying hidden thermodynamic processes.9 In 2005,
Ford derived a non-quantum justification (see next section) claiming that enthalpy/entropy compensation ($T^{\text{comp}} > 0$ in eqn (2)) occurs when the minimum host–guest separation $r_0$ in the [AB] pairs remains constant within a series of intermolecular binding events. However, the linearity proposed in eqn (2) does not result from Ford’s model, and alternative physical justifications for parabolic or rectangular-hyperbolic correlation for H/S compensation have been proposed. In this context, Liu and Guo documented that the general emergence of linear H/S compensation is often the consequence of statistical and mathematical artefacts arising from the data analysis. The dimerization process depicted in Fig. 1a illustrates this statement since the convincing linear relationship found for H/S compensation (Fig. 1b) is significantly discrepant when one considers the alternative, but equivalent van’t Hoff plots which should cross at a common ($T^{\text{comp}}$, $\Delta G^{\text{comp}}$) couple (red rectangle in Fig. 1c).

Since complexation processes occurring in dilute solutions are complicated by unavoidable solvation changes, which may induce additional H/S compensation phenomena, we resorted to the melting of solids into isotropic liquid conducted in absence of solvents or of additives for getting a rough estimation of the strength of the intermolecular interactions occurring in a solid. These processes can be idealized as reversible $n$th order chemical reactions, in which $n$ identical monomeric units A associate into fully assembled entities $A_n$ (eqn (3)).

\[
A_n \frac{\Delta G_{\text{asso}}^{0} - \Delta H_{\text{m}}^{0} - T \Delta S_{\text{m}}^{0}}{\Delta C_{\text{d}}^{O} - \Delta C_{\text{d}}^{\text{asso}} - \Delta C_{\text{d}}^{\text{m}}} A_n
\]

At the melting temperature $T = T_m$, the solid–liquid phase equilibrium implies that $\Delta G_{\text{asso}} = \Delta G_{\text{m}} = 0$ and $T_m = \Delta H_{\text{m}}^{0} / \Delta S_{\text{m}}^{0} = \Delta H_{\text{asso}} / \Delta S_{\text{asso}} = \Delta H_{\text{m}} / \Delta S_{\text{m}}$ when one reasonably assumes that the melting enthalpies $\Delta H_{\text{m}}$ and entropies $\Delta S_{\text{m}}$ operating at the melting temperatures are satisfyingly estimated by those defined in the standard conditions ($i.e. \Delta H_{\text{m}} \equiv \Delta H_{\text{m}}^{0}$ and $\Delta S_{\text{m}} \equiv \Delta S_{\text{m}}^{0}$). Expressed at a common reference temperature $T^0$, each melting process is further characterized by its specific standard free energy change $\Delta G_{\text{m}}^{0} = \Delta H_{\text{m}}^{0} - T^0 \Delta S_{\text{m}}^{0} = \Delta H_{\text{m}} - T^0 \Delta S_{\text{m}}$ which is further used to compute the standard cohesive free energy density CFED $= \Delta G_{\text{m}}^{0} / V_{\text{mol}}$ ($V_{\text{mol}}$ is the molar volume), a parameter estimating the average cohesive forces operating in the solid.

Fig. 1 (a) Schematic dimerization of complexes [Eu(L$^{1+3}$)(NO$_3$)$_3$] occurring in dichloromethane, (b) associated ‘linear’ $\Delta H_{\text{dim}}^{0}$ vs. $\Delta S_{\text{dim}}^{0}$ plots and (c) individual van’t Hoff $\Delta G_{\text{dim}} / T$ vs. $T^{-1}$ plots for nine different $R$ substituents. Adapted from ref. 17.

Fig. 2. Plots of (a) the cohesive free energy densities (CFED) versus the melting temperature ($T_m$) and (b) the melting temperature ($T_m$) versus the molar volumes ($V_{\text{mol}}$) for saturated linear hydrocarbons $C_nH_{2n+2}$ ($n = 2–20$). The dotted red traces show the theoretical curves. Adapted from ref. 17.

### Theoretical background

Following the formalism of molecular association proposed by Luo and Sharp, the equilibrium constant $K_{\text{asso}}^{AB}$ associated with the simple binding event shown in eqn (1) can be written as eqn (4), where $c$ is the standard concentration of the reference state (fixed to 1 M in this contribution), $H(r, \Omega)$ is a bonding function depending on the separation ($r$) and orientation ($\Omega$) of the two partners in the [AB] pair ($H(r, \Omega) = 1$ when the complex [AB] exists and $H(r, \Omega) = 0$ otherwise), $\beta = (k_B T)^{-1}$ stands for the thermal factor and $\omega(r, \Omega)$ is the potential mean force between A and B.

\[
K_{\text{asso}}^{AB} = \frac{|AB|}{|A||B|} = \frac{c}{8\pi} \int H(r, \Omega) e^{-\beta \omega(r, \Omega)} dr d\Omega
\]

Solving eqn (4) within the frame of the van’t Hoff isotherm for a non-directional harmonic potential $\omega(r, \Omega) = u_{\text{min}}^{AB} + (\kappa_A B / 2) r^2$ operating between A and B, whereby $u_{\text{min}}^{AB}$ is the minimum potential energy and $\kappa_A B$ is the force constant, leads to eqn (5) and (6), where $N_A$ is Avogadro number.

\[
\Delta H_{\text{asso}}^{AB} / N_A = u_{\text{min}}^{AB} + \frac{3}{2} k_B T
\]
Interestingly, the enthalpy change $\Delta H_{asso}^{AB}$ mainly depends on the magnitude of the interaction energy $u_{min}$, while the entropy change $\Delta S_{asso}$ is controlled by the force constant $k_{A,B}$, the latter term being an estimation of the capacity of the bound system to gain residual degrees of freedom. The development of the harmonic potential at the minimum of a standard Lennard-Jones potential $V_{lj}$ shows that the absolute minimum energy of the attractive well depth corresponds to $u_{min}$ when the equilibrium A⋯B separation amounts to $2^{1/6}r^0$ ($r^0$ is the critical intermolecular A⋯B distance at which the interaction potential is zero: $V_{lj}(r = r^0) = 0$, see Fig. S1 in the ESIT). Consequently, the total energy of the harmonic oscillator for the special motion amplitude $(2^{1/6}r^0 - r^0)$ exactly corresponds to the well depth $u_{min}$ and eqn (7) results:

$$-u_{min}^{A,B} = k_{A,B}^{A,B} \left( \frac{1}{2} - 2^{1/6} \right)^2 f^{A,B} = -\frac{2}{(1 - 2^{1/6})^2} f^{A,B} u_{min}$$

For a minor structural perturbation affecting a series of A and B partners, the minimum contact distance $r_0$ is constant within the resulting [AB] pairs and Ford’s model (eqn (7), righthand side) predicts that the force constants $k_{A,B}$ (which affects the entropy changes, eqn (6)) are linearly correlated with the potential well depths $u_{min}^{A,B}$ (which measures the enthalpy change, eqn (5)). Since the coefficient $f = 2/(1 - 2^{1/6})^2 r_0^2$ is positive, a larger cohesive energy between A and B in the [AB] pair (i.e. $u_{min}^{A,B}$ and $\Delta H_{asso}^{AB}$ become more negative) produces an increase in the force constant $k_{A,B}$, hence in the mean vibrational frequency in the deeper potential. Since $\Delta S_{asso}^{A,B}/N_{Av} \propto -3/2 \ln(k_{A,B})$ in eqn (6), larger $k_{A,B}$ induces more negative association entropies and H/S compensation occurs. This model is nothing but the Einstein model for crystals, from which Lindemann postulated that the melting of a solid occurs when the amplitude of the atomic thermal vibrations reaches some critical fraction of the equilibrium lattice spacing. Introducing eqn (7) into eqn (5) provides a simple correlation between the enthalpy change of the association process and the force constant in the bound state (eqn (8); $R = k_B N_{Av}$ is the ideal gas constant). Further introduction into eqn (6) gives the searched (logarithmic) dependence between $\Delta H_{asso}^{AB}$ and $\Delta S_{asso}$ (eqn (9)).

$$\frac{\Delta H_{asso}^{AB}}{N_{Av}} = -k_{A,B}^{A,B} + \frac{3}{2} k_B T \Rightarrow k_{A,B}^{A,B} = \frac{f}{N_{Av}} \left( \frac{3}{2} RT - \Delta H_{asso}^{A,B} \right)$$

$$\Delta S_{asso}^{A,B} = R \left[ \ln(c^f) + \frac{3}{2} \ln \left( \frac{2\pi e RT}{f} \right) - \frac{3}{2} \ln \left( \frac{3}{2} RT - \Delta H_{asso}^{A,B} \right) \right]$$

Its application to the melting of the $A_x$ assembly shown in eqn (3) requires $\Delta H_m = -\Delta H_{asso}$ and $\Delta S_m = -\Delta S_{asso}$, and eqn (10) eventually results

$$\Delta S_m = k_B \ln \left[ \frac{2\pi e}{f k_{A,B}^{A,B}} \right]^{3/2}$$

Applied to the melting of a series of similar compounds, the force constant $k_{A,B}$ can be approximated by a first-order Taylor series around its average magnitude $k_{0,B}^{A,B}$, and $\ln \left( \frac{3}{2} RT + \Delta H_m \right)$ in eqn (10) can be replaced with $

$$\ln \left( \frac{3}{2} RT + \Delta H_{m,0} \right) + \left( \Delta H_m - \Delta H_{m,0} \right) \left( \frac{3}{2} RT + \Delta H_{m,0} \right)$$

where $\Delta H_{m,0}$ is the average melting enthalpy in the series. However, the rapid divergence of the logarithmic function from a linear approximation restricts the use of linear H/S compensation within this model to a narrow thermodynamic range of melting processes. In this contribution, we propose to improve the predictive capacity of the latter thermodynamic model via a thorough investigation of the melting processes occurring in metals, in various organic compounds and in inorganic oxides.

### Results and discussion

#### Modelling H/S compensation

Thanks to Yaws’ efforts for gathering melting enthalpies and entropies (Fig. 3), we were able to select six families of compounds, linear alkanes (Table S1†), linear alkanolic acids (Table S2†), organosilanes (Table S3†), lanthanide metals (Table S4†), transition metals (Table S5†) and transition metal oxides (Table S6†), for which (i) the internal cohesion forces result from various types of intermolecular interactions (permanent and/or induced electric multipolar, H-bonds, covalency) and (ii) the melting temperatures span a broad domain. Assuming the justified approximation that $\Delta H_m$ and $\Delta S_m$ are essentially constant over a reasonable temperature range around a reference temperature $T^0$ taken as the average melting temperature within each series (Tables S1–S6 in the ESIT), we were able to select six families of compounds for which (i) the internal cohesion forces result from various types of intermolecular interactions (permanent and/or induced electric multipolar, H-bonds, covalency) and (ii) the melting temperatures span a broad domain. Assuming the justified approximation that $\Delta H_m$ and $\Delta S_m$ are essentially constant over a reasonable temperature range around a reference temperature $T^0$ taken as the average melting temperature within each series (Tables S1–S6 in the ESIT), the experimental $\Delta S_m$ versus $\Delta H_m$ plots of the six individual series were fitted with Ford’s model by tuning $f$ in eqn (10), but only a poor match
of the use of first-order, respectively second-order Taylor polynomials for approaching eqn (10).\(^2\) Higher-order Taylor series are precluded by the well-known Runge’s phenomenon, which strongly limits the approximation of logarithmic mapping by using power polynomials of increasing orders.\(^2\) On the contrary, the use of a Hill function has no mathematical justification for fitting logarithmic plots, but its stability over a large enthalpy range appeared to be a considerable advantage for fitting \(H/S\) compensation (vide infra). A simple look at Fig. 3 indicates that \(\Delta S_m \to 0\) when \(\Delta H_m \to 0\) for melting processes recorded at normal pressure (in other words, the melting temperature \(T_m = \Delta H_m/\Delta S_m\) does not diverge for \(\Delta S_m \to 0\)),\(^2\) a boundary condition which is not satisfied by the fitting traces collected in Fig. 4 and S2–S6 (ESI†). Including the latter constraint into \(H/S\) compensation results in the loss of one degree of freedom by fixing one parameter during the fitting process (Tables 1 and S7–S11,\(^\dagger\) column 1). Consequently, Ford’s approach lacks of acceptable correlation (magenta traces in Fig. 5 and S7–S11,\(^\dagger\) no tuneable parameter), while linear approximations (red traces in Fig. 5 and S7–S11,\(^\dagger\) one tuneable parameter, \(\Delta S_m = b\Delta H_m\)) are dismissed because only a single melting temperature \(T_m = b^{\text{−1}}\) is tolerated along the complete series of compounds.

Therefore, among the four mathematical functions selected for extending the logarithmic \(H/S\) dependence suggested by the simple Ford’s model, only parabolic (blue traces, two tuneable parameters) and Hill (green traces, three tuneable parameters) plots provide satisfying fitted traces compatible with the inclusion of the extra boundary condition \(\Delta S_m \to 0\) when \(\Delta H_m \to 0\) (Fig. 5 and S7–S11, Tables 1 and S7–S11, ESI†).\(^\dagger\)

### Cohesive free energy densities and three-dimensional \(\Delta H_m, \Delta S_m, V_{\text{mol}}\) plots

Taking Trouton’s rule into account for the vaporization of liquids,\(^2\) Hildebrand introduced the concept of cohesion energy density \(\text{CED} = (\Delta H_v - RT)/V_{\text{mol}}\) for estimating the average cohesive forces operating within liquids and for predicting their vaporization temperatures (\(\Delta H_v\), is the vaporization enthalpy).\(^1\)\(^9\) In absence of valuable alternative to Trouton’s rule fixing the melting entropy in solids,\(^1\)\(^8\) we have resorted to standard melting Gibbs free energy \(\Delta G_m = \Delta H_m - T \Delta S_m\) computed at a

![Table 1](image)

Table 1: Entropy–enthalpy correlations fitted for linear alkanes \(\text{C}_n\text{H}_{2n+2} (n = 1–30)^a\)

| Function | \(f/m^2\) | \(a\) | \(b\) | \(c\) | \(d\) |
|----------|---------|------|------|------|------|
| **Unconstrained** | | | | | |
| Eqn (10) | | 6(7) \(\times 10^5\) | | | | |
| \(\Delta S_m = a + b \Delta H_m\) | | | 33(2) | 2.69(4) \(\times 10^{-3}\) | | |
| \(\Delta S_m = a + b \Delta H_m + c(\Delta H_m)^2\) | | | 26(2) | 3.2(1) \(\times 10^{-3}\) | \(-5(1) \times 10^{-9}\) | |
| \(\Delta S_m = a(\Delta H_m - b)/(c - \Delta H_m)^2\) | | | 4.9(8) \(\times 10^2\) | \(-7(8) \times 10^2\) | 3.3(5) \(\times 10^3\) | 6.2(3) \(\times 10^{-1}\) |
| **Constrained: \(\Delta S_m \to 0\) when \(\Delta H_m \to 0\)** | | | | | | |
| Eqn (10) | 11.39 | | | | | |
| \(\Delta S_m = a + b \Delta H_m\) | | | 0 | 3.17(6) \(\times 10^{-3}\) | | |
| \(\Delta S_m = a + b \Delta H_m + c(\Delta H_m)^2\) | | | 0 | 4.2(2) \(\times 10^{-3}\) | \(-1.3(2) \times 10^{-8}\) | |
| \(\Delta S_m = a(\Delta H_m - b)/(c - \Delta H_m)^2\) | | | 4.5(4) \(\times 10^2\) | 0 | 3.0(3) \(\times 10^3\) | 5.9(2) \(\times 10^{-1}\) |

\(^a\) Units used for polynomial fits: \(a/J\) mol\(^{-1}\) K\(^{-1}\), \(b/K^{-1}\) and \(c/J\) mol K\(^{-1}\). Units used for reciprocal Hill fits: \(a/J\) mol\(^{-1}\) K\(^{-1}\), \(b\) and \(c\) in J mol\(^{-1}\).
reference temperature $T^0$ for estimating the standard cohesive free energy densities $\text{CFED} = \frac{\Delta G^0_m}{V_{\text{mol}}}$ along each homogeneous series of compounds (Tables S1–S6, ESI†). We are aware that the melting enthalpies ($\Delta H_m$) and entropies ($\Delta S_m$) are indeed collected at different temperatures for each compound (i.e. at $T_m$) but their dependence on temperature is limited, and the choice of a reference temperature $T^0$ around the center of each series obeying $H/S$ compensation is thus acceptable for computing $\Delta G^0_m$. The experimental linear correlations observed between CFED and the associated melting temperatures $T_m$ in Fig. 6 provide eqn (11), which interconnects $\Delta H_m$, $\Delta S_m$ and $V_{\text{mol}}$ within each series (note that $T_m = \Delta H_m/\Delta S_m$)

$$\text{CFED} = \frac{\Delta G^0_m}{V_{\text{mol}}} = \frac{-\Delta H_m - T^0 \Delta S_m}{V_{\text{mol}}} = \lambda T_m + \mu \quad (11)$$

Taken separately, eqn (11) is of limited interest for tuning melting temperatures by molecular design since only the molar volume $V_{\text{mol}}$ can be easily computed for unknown compounds, while satisfying estimations and/or modelling of melting...
entropies or enthalpies are difficult. However, the consideration of H/S compensations operating in each series (Fig. 5 and S7–S11, ESI†) provides a correlation $\Delta S_m = g(D_H_m)$ which can be introduced into eqn (11) to give eqn (12) when one reminds that $T_m = \Delta H_m/\Delta S_m = \Delta H_m/g(D_H_m)$ ($g_i$ represents either a parabolic or a reciprocal Hill function).

$$g(D_H_m)\Delta H_m - T^0g(D_H_m) - V_{mol}H - \lambda V_{mol}\Delta H_m = 0 \quad (12)$$

For quadratic fits, $\Delta S_m = g_{quad}(\Delta H_m) = b\Delta H_m + c(\Delta H_m)^2$ and eqn (12) corresponds to a polynomial, which can be solved to give an estimation of the melting enthalpy $\Delta H_m$ as soon as the molar volumes $V_{mol}$ of a target compound belonging to the series is at hand. The associated melting entropy $\Delta S_m$ immediately results from $H/S$ correlation. The same procedure holds when using reciprocal Hill fits characterized by $\Delta S_m = g_{Hill}(\Delta H_m) = a[\Delta H_m/(\Delta H_m + c)]d$. The quality of the correlations can be estimated in the three-dimensional $V_{mol}$, $\Delta H_m$, $\Delta S_m$ plots for each series (Fig. 7 and S12–S16, ESI†).

As an ultimate step, the combination of melting enthalpies and entropies into the melting temperatures $T_m = \Delta H_m/\Delta S_m$ reduces the dimension of the correlation shown in Fig. 7 to give an easy-manageable $T_m$ versus $V_{mol}$ plot (Fig. 8 and S17–S21, ESI†). The careful inspection of the latter two-dimensional plots indicate that, for any series under investigation, reciprocal Hill functions provide the best fits.

Combination of $H/S$ compensation with linear cohesive free energy densities (CFED) for predicting the melting temperatures of substituted lipophilic cyanobiphenyls

Lipophilic cyanobiphenyls with variable lengths ($n$-CB) and substitutions (12Me$^m$-CB) have been considered as precursors for inducing liquid crystalline mesophases (Fig. 9a, Table S12, Fig. 9).

**Fig. 9** (a) Chemical structures of substituted cyanobiphenyls with associated plots corresponding to (b) melting entropy versus melting enthalpy changes, (c) cohesion free energy densities (CFED) versus the melting temperature ($T_m$) and (d) three-dimensional $V_{mol}$ versus $\Delta H_m$ and $\Delta S_m$ fitted with a parabolic $H/S$ function (blue trace) or a reciprocal Hill function (green trace). Reference temperature $T^0 = 325.92 \text{ K}$. 

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support in favor of the Hill plot as the best predictive tool within this novel series, in line with the previous analysis of linear alkane and alkanoic acids, organosilanes, lanthanide metals, transition metals and transition metal oxides.

Conclusion

The non-directional Ford’s approach uses a single harmonic potential with a unique minimum contact distance for catching any intermolecular interaction. It predicts that, for a series of binding reactions involving structurally similar partners, the thermodynamic of association/dissociation is dominated by $H/S$ compensation and logarithmic $\Delta S$ versus $\Delta H$ dependences are expected. Applied to the melting processes occurring in six (arbitrarily selected) sets of chemical compounds, this rough model justifies the emergence of $H/S$ compensation, but it fails in reproducing accurate trends because the modelling of the intermolecular interactions operating in solid with the help of a single harmonic potential is a too big oversimplification. Increasing the number of degrees of freedom for approaching $H/S$ correlations restores satisfying fits when using linear or quadratic polynomials, or reciprocal Hill functions. The inclusion of the boundary condition $\Delta S_m \to 0$ when $\Delta H_m \to 0$ recues linear $H/S$ correlations and only quadratic or Hill plots are able to give satisfying $\Delta S_m$ versus $\Delta H_m$ fits. Standard empirical analysis of enthalpy/entropy compensation within the frame of this simple binding thermodynamics stops at this point, and no predictive tool is at hand for estimating the strength of intermolecular interactions and for programming melting temperatures. The additional concept of standard cohesive free energy density ($\text{CFED} = \Delta G_m/V_{mol}$) and its observed linear correlation with the melting temperatures (eqn (11)) introduces an unprecedented relationship between the molar volume, $V_{mol}$, a parameter accessible to supramolecular chemists and the correlated melting enthalpies and entropies. For the six different series of compounds selected in this contribution, i.e. linear alkanes, linear alkanoic acids, organosilanes, lanthanide metals, transition metals and transition metal oxides, the ultimate reduced $T_m$ versus $V_{mol}$ plots satisfactorily reproduce the experimental data. We notice that the use of reciprocal Hill plots provide the best fits for $H/S$ correlations in melting processes, and its application for rationalizing melting temperatures recently determined for substituted cyanobiphenyls provides reasonable predictions for currently unknown compounds in the series.

Experimental

The mathematical analyses were performed by using Igor Pro® (WaveMetrics Inc.) and Excel® [Microsoft] softwares.

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