

# PCCP

Accepted Manuscript



This is an *Accepted Manuscript*, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

*Accepted Manuscripts* are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. We will replace this *Accepted Manuscript* with the edited and formatted *Advance Article* as soon as it is available.

You can find more information about *Accepted Manuscripts* in the [Information for Authors](#).

Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal's standard [Terms & Conditions](#) and the [Ethical guidelines](#) still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this *Accepted Manuscript* or any consequences arising from the use of any information it contains.

**A Study of the Relationship between Water and Anions of the Hofmeister Series using Pressure Perturbation Calorimetry**

Jordan W. Bye and Robert J. Falconer\*

Department of Chemical & Biological Engineering, ChELSI Institute, University of Sheffield, Sheffield, S1 3JD, England

\* To whom the correspondence should be addressed. Telephone +44 114 2228253, Fax +44 114 2227501 Email r.j.falconer@sheffield.ac.uk

**Abstract**

Pressure perturbation calorimetry (PPC) was used to study the relationship between water and sodium salts with a range of different anions. At temperatures around 25 °C the heat on pressurisation ( $\Delta Q$ ) from 1 to 5 bar was negative for all solutions relative to pure water. The raw data showed that as the temperature rose, the gradient was positive relative to pure water and the transition temperature where  $\Delta Q$  was zero was related to anion surface charge density and was more pronounced for the low-charge density anions. A three component model was developed comprising bulk water, the hydration layer and the solute to calculate the molar expansivity of the hydration layer around the ions in solution. The calculated molar expansivities of water in the hydration layer around the ions were consistently less than pure water.  $\Delta Q$  at different disodium hydrogen phosphate concentrations showed that the change in molar enthalpy relative to pure water was not linear even as it approached infinite dilution suggesting that while hydration layers can be allocated to the water around ions this does not rule out interactions between water and ions extending beyond the immediate hydration layer.

**Keywords:**

Kosmotrope, chaotrope, DSC, thermal expansion coefficient, solvation shell

**Introduction**

Water is a unique solvent that plays a critical role in supporting life on earth. Despite this there is still ongoing discussion between scientists on the nature of water and its association with inorganic and organic molecules. A unique property of water is its propensity to rapidly swap protons between water molecules at a picosecond timescale, referred to as hydrogen bonding. When a salt is added to water it dissociates into anions and cations as described by Arrhenius in 1887.<sup>1</sup> These ions have electrical fields which interact with the dipolar water molecules that arrange themselves around the ions. These arranged water molecules are often referred to as the hydration layer. There is discussion around whether the ion's effect extends just to the water molecules at the interface with the ion, into a second hydration layer or beyond into more distant water.

In 1888 Franz Hofmeister published two papers on the effect salts had on protein solubility in water.<sup>2,3</sup> An English translation of these papers is available in Kunz et al. 2004.<sup>4</sup> The observation was made that ions could be ordered according their ability to precipitate or solubilise protein ensembles. Hofmeister explained his observations in terms of the hydration strength of the ions (their ability to absorb water). Ions with strong hydration compete with the protein for the water and it is this that causes the protein's precipitation. A similar ordering of the ions was observed in surface tension measurements by Adolf Heydweiller in 1910.<sup>5</sup> An alternative explanation for the Hofmeister effect was published in 1930 by Cox and Wolfenden which explained the observed effect of ions on viscosity in terms of the degree of polymerisation of the solvent in the presence of the ions.<sup>6</sup> Over a period of years evolved the explanation for the Hofmeister effect as the ability of an ion to alter the hydrogen bond population of the solvent<sup>7</sup> often referred to as the structure making and structure breaking theory. This theory relied on the ions having the ability to interact with water molecules beyond its first hydration layer. In 2003, the structure making and structure breaking theory was directly challenged using data from femtosecond pump-probe spectroscopy that suggested there was no measurable interaction with water beyond the first hydration layer.<sup>8</sup> Since then evidence using neutron diffraction,<sup>9</sup> molecular dynamic simulation<sup>10</sup> and calorimetry techniques<sup>11</sup> have suggested long-range interactions beyond the first hydration layer are likely.

In this paper we add pressure perturbation calorimetry (PPC) at low pressure changes to the existing palette of techniques for studying electrolyte solutions and discuss the results with reference to the Hofmeister effect. The current PPC technology was preceded by instrumentation that subject the samples to high pressure changes (up to 4,000 bar) which were suitable for studying phase transitions in organic solvents and polymers, and had been around since the 1970s.<sup>12-15</sup> More recently a modification to differential scanning microcalorimetry (DSC) instrumentation enabled this type of analysis to be carried out at lower pressure changes (4 bar) with a high degree of sensitivity which enabled the analysis of the heat change on pressurization for diverse solutes in water.<sup>16,17</sup> These devices measure the heat change in the solution as the pressure above the solution is altered. From this the thermal expansion coefficient ( $\alpha$ ) of the solute has been calculated and the hydrogen-bond population of the water at the solute-water interface studied.<sup>18,19</sup> This approach has been applied to a diverse range of solutes including polymers,<sup>16</sup> amino acids,<sup>17</sup> small inorganic and organic molecules,<sup>18</sup> proteins,<sup>16-28</sup> lipid micelles and bilayers,<sup>29-31</sup> and nucleic acid.<sup>32,33</sup>

The calculation of the thermal expansion coefficient from PPC data assumed a two-state model for water with a relatively low density and a denser liquid species. PPC relies on the Le Chatelier's principle, that on pressurization the water will try to equilibrate by moving towards the higher density form. The derivation of the equation that has been used to calculate the thermal expansion coefficient uses a two-component system for small molecules in solution where  $V_{tot}$  is the total volume,  $V_0$  is the molar volume of solvent and  $\bar{V}$  is the partial specific volume of the solute in the solution which includes any volume changes induced in the solvent;  $x_0$  is molar fraction of the solvent and  $x_s$  is molar fraction of the solute.<sup>17,18</sup>

$$V_{tot} = x_0 V_0 + x_s \bar{V} \quad \text{Equation 1}$$

An alternative approach to the two-component model is to directly take into account the population of water around the solute (referred to as the hydration layer or solvation shell in the literature) that has a molar volume which may be different to bulk water (unperturbed or pure water). Whichever

model is selected for defining the boundary of the hydration layer around a solute this population of water has to be defined in terms of its own average molar volume, molar fraction and average thermal expansion coefficient. In this paper the authors propose an alternative approach to studying solutes in water taking the hydration layer into account and present the results for sodium salts with different anions to illustrate how this approach can be implemented.

## Data Analysis

The authors initially calculated values for the coefficient of thermal expansion using the equation derived in Lin et. al. 2002<sup>17</sup> using published apparent partial volume ( $\bar{V}$ ) values<sup>34</sup> and the experimental PPC results for 100 mM salt solutions (see Figure S1 in the supplementary information). While the coefficient of thermal expansion for sodium chloride, sodium bromide, sodium iodide, sodium thiocyanide and sodium perchlorate are plausible those for sodium fluoride and disodium sulphate were unusual. Sodium fluoride and disodium sulphate both have high charge density and their conventional apparent partial volume ( $\bar{V}$ ) values<sup>34</sup> are markedly lower than their intrinsic volumes ( $V_{int}$ )<sup>35</sup> (see Figure S2 in the supplementary information). It was concluded that the two-component model was problematic for high charge density ions. It was this observation that led the authors to suggest a three-component model for salt solutions where  $V_b$  is the molar volume of bulk solvent,  $V_h$  is the average molar volume of the solvent within the hydration layer and  $V_s$  is the molar volume of solute;  $x_b$  is the molar fraction of the bulk solvent,  $x_h$  is the molar fraction of the solvent within the hydration layer and  $x_s$  is the molar fraction of the solute.

$$V_{tot} = x_b V_b + x_h V_h + x_s V_s \quad \text{Equation 2}$$

The heat ( $Q$ ) is derived from first principles for a single component system where  $T$  is temperature,  $\Delta P$  is change in pressure,  $\alpha$  is the thermal expansion coefficient and  $V$  is the volume.<sup>17</sup>

$$Q = -T \Delta P \alpha V \quad \text{Equation 3}$$

For a three component system this becomes

$$Q = -T \Delta P [x_b V_b \alpha_b + x_h V_h \alpha_h + x_s V_s \alpha_s] \quad \text{Equation 4}$$

Where  $\alpha_b$ ,  $\alpha_h$  and  $\alpha_s$  is the thermal expansion coefficient of the bulk water, hydration layer and solute, respectively.

The difference in heat between the sample and reference cells is

$$\Delta Q = T \Delta P [x_0 V_0 \alpha_0 - x_b V_b \alpha_b - x_h V_h \alpha_h - x_s V_s \alpha_s] \quad \text{Equation 5}$$

Where  $x_0$ ,  $V_0$  and  $\alpha_0$  is the molar fraction, molar volume and thermal expansion coefficient of the pure water in the reference cell.

As  $V_0 = V_b$ ,  $\alpha_0 = \alpha_b$  and  $x_0$  is 1 the equation can be simplified to

$$\Delta Q = T \Delta P [(1 - x_b)(V_b \alpha_b) - x_h V_h \alpha_h - x_s V_s \alpha_s] \quad \text{Equation 6}$$

As  $x_b = 1 - x_h - x_s$

$$\Delta Q = T\Delta P[(x_s + x_h)(V_b\alpha_b) - x_hV_h\alpha_h - x_sV_s\alpha_s] \quad \text{Equation 7}$$

As the  $x_h$  is defined by the multiple number ( $n$ ) of water molecules around the ions  $x_h = nx_s$ , the equation can be further simplified to

$$\Delta Q = T\Delta P[(n + 1)(x_sV_b\alpha_b) - nx_sV_h\alpha_h - x_sV_s\alpha_s] \quad \text{Equation 8}$$

$$\Delta Q = T\Delta Px_s[(n + 1)(V_b\alpha_b) - nV_h\alpha_h - V_s\alpha_s] \quad \text{Equation 9}$$

The experimental results show that as  $\Delta P$  and  $x_s$  tend to zero  $\Delta Q$  tend to zero but as  $T$  tends to zero  $\Delta Q$  is negative therefore a constant ( $A$ ) is added to the equation. Note  $A$  is undefined and may be due to the presence of the sodium cation and its interaction with the anion. Where  $T$  is the variable

$$\Delta Q = T\Delta Px_s[(n + 1)(V_b\alpha_b) - nV_h\alpha_h - V_s\alpha_s] + A \quad \text{Equation 10}$$

Accurate values for density, specific volume ( $V_b$ ), thermal expansivity ( $\alpha_b$ ), and compressibility of ordinary water are available in the literature.<sup>36</sup> The number of water molecules for the first and second hydration layers around anions and cations has been estimated by diffraction methods<sup>37</sup> and molecular dynamic simulation<sup>38</sup> so a value can be attributed to  $n$ . The  $V_s\alpha_s$  is known for crystalline sodium halides.<sup>39</sup> Molar expansivity of the hydration layer ( $E_h$ ) can be derived using equation 10 as  $E_h = V_h\alpha_h$ .

Determining the thermal expansion coefficient ( $\alpha_h$ ) is more difficult as an estimate as a value for  $V_h$  is needed. The difference between experimental determined apparent partial molal volumes<sup>34</sup> and the intrinsic molal volume<sup>35</sup> could be used to estimate the change in molar volume of the water within the hydration layer. An alternative approach would be to calculate the effect of electrostriction on the molar volume of the water within the hydration layer from molecular dynamic simulation data.<sup>38</sup> Both these methods would be estimates at best so the calculations used by the authors were confined to calculating the molar expansivity of the hydration layer around the ions.

137

## 138 Methods and Materials

**139 Sample Preparation.** Ultra-pure water and all salts were sourced from Sigma Aldrich, Gillingham, UK  
 140 with purities >99.9%. Sodium was used as the counter-ion for all anions. 2000 mM stock solutions at  
 141 pH 7 were made for all salts (sodium fluoride stock solutions were made to 100 mM at pH 8.4 due to  
 142 solubility limits); stock solution dilutions were performed by adding the required amount of pure  
 143 water.

**144 Pressure perturbation calorimetry (PPC) measurements.** PPC measurements were obtained using a  
 145 capillary Nano-DSC (TA Instruments, New Castle, DE, USA). Samples were degassed for 1 hour at 30  
 146 °C to remove dissolved gas from samples and eliminate bubble formation during the scan. Heat  
 147 effects ( $\Delta Q$ ) were measured during alternating pressure pulses of  $\pm 4$  bar from 1 bar to 5 bar at 1 °C  
 148 intervals from 7 – 92 °C, giving a usable data range of 10 – 90 °C. A heating rate of 0.1 °C/min was  
 149 used to satisfy isothermal conditions required during pressure pulses; this scanning rate is slower  
 150 than the instrument feedback.<sup>33</sup> The instrument was held at a constant temperature for 30 minutes  
 151 before each scan to ensure that any asymmetry between the reference and sample cells was

minimal. Heat changes during pressurization steps were used for data analysis and were calculated using the NanoAnalyze software (TA Instruments, New Castle, DE, USA) provided by the manufacturer. Water baseline scans were performed with pure water in reference and sample cell, scans with salt present were performed with pure water in the reference cell and salt solution in the sample cell. The area under each thermal power spike was calculated by integration using NanoAnalyze software (TA Instruments, New Castle, DE, USA) and was used as the heat change during pressurization for that temperature, shown in Figures 1a, b, c and d.

**Calculation of molar expansivity.** Application of Equation 10 is simple in practice for solutes where the specific volume ( $V_s$ ), thermal expansivity ( $\alpha_s$ ) are known and  $n$  can be estimated. The specific volume ( $V_b$ ), thermal expansivity ( $\alpha_b$ ) of pure water are known.<sup>36</sup>  $A$  was determined by adjusting it till the calculated  $\Delta Q$  fitted experimental  $\Delta Q$  data.

Equation 10 was rearranged as follows so molar expansivity ( $V_h\alpha_h$ ) can be calculated.

$$V_h\alpha_h = \frac{[(n+1)(V_b\alpha_b)] - V_s\alpha_s - \frac{\Delta Q - A}{T\Delta Px_s}}{n} \quad \text{Equation 11}$$

## Results

2 M stock solutions were made for disodium hydrogen phosphate (at pH 7.0 this is a mixture of sodium dihydrogen phosphate and disodium hydrogen phosphate), disodium sulphate, sodium fluoride, sodium chloride, sodium bromide, sodium iodide, sodium thiocyanate and sodium perchlorate at pH 7.0 (sodium fluoride at pH 8.4 was only made to 100 mM due to solubility limits). These salts were chosen so that a range of anions from the Hofmeister series could be studied, including the simple monoatomic halogen anions and more complex oxoanions.

Thermal data was collected as microwatts ( $\mu W$ ) of power required to keep the reference and sample cell temperatures identical upon pressurization and depressurization steps, the raw data for 1000 mM disodium hydrogen phosphate, sodium chloride and sodium thiocyanate, is shown in Figures 1a, b and c, respectively. Figure 1d shows a small temperature range in which the area under the thermal spikes (which was integrated) can be more clearly seen, along with the pressure change which caused the thermal spike. The power function was converted into heat absorbed or released ( $\Delta Q$ ) in micro joules ( $\mu J$ ) by the sample during pressure changes between 1-5 atm.

The  $\Delta Q$  associated with pressurization of the salt solutions at 100 mM and 1000 mM, respectively, is shown in Figures 2a and 2b. The plotted heat energies show the difference in energy added to the sample cell compared to the reference cell which contained pure water. Sodium fluoride was not tested at 1000 mM due to its low solubility. For all salts tested a negative  $\Delta Q$  is observed at lower temperatures upon sample pressurization, this indicates that an exothermic process is taking place. The temperature where  $\Delta Q$  upon sample pressurization is zero ( $T_i$ ) was found to be salt dependent and independent of salt concentration. The  $T_i$  were found to be 60.5 °C for disodium hydrogen phosphate, 59.5 °C for disodium sulphate, 54 °C for sodium fluoride, 59 °C for sodium chloride, 65.5 °C for sodium bromide, 76.5 °C for sodium iodide, 90.5 °C for sodium thiocyanate and 104 °C for sodium perchlorate at all salt concentrations. The temperature where  $\Delta Q$  upon pressurization



became endothermic was not reached for sodium perchlorate concentrations. The temperature where its  $\Delta Q$  became endothermic was calculated by extrapolation reached for sodium perchlorate. The surface charge density of the ions tested plotted against the temperature where the heat changes upon pressurization of the salt solution becomes endothermic at the same temperature is shown in Figure 3. Charge, ionic and apparent ionic radii for sulphate, fluoride, chloride, bromide, iodide, thiocyanate and perchlorate were used to calculate the surface charge density of each ion.<sup>29,30</sup> It should be noted that phosphate was omitted from this graph due to phosphate being composed of a mixture of  $\text{H}_2\text{PO}_4^-$  and  $\text{HPO}_4^{2-}$  at pH 7.0. There is a trend of increasing  $T_i$  values with a decrease in surface charge density. Thiocyanate does not fit the trend as well as the other ions which may be due to it being treated as a sphere to obtain its ionic radius, when it is not spherical. It should be noted that the ordering of these ions by the  $T_i$  value closely agrees with the order seen in the Hofmeister series.<sup>2-4</sup>

The average gradient on the  $\Delta Q$  versus temperature plot for the eight salts tested at 100 mM from 9.5 – 91.5 °C. The average gradient for each salt (100 mM) was calculated by dividing the difference in  $\Delta Q$  upon pressurization at 9.5 °C and 91.5 °C. From Figure 2 it can clearly be seen that ions with a -2 charge release more energy at lower temperatures and require more energy at higher temperatures upon pressurization than ions with a -1 charge. It is worth remembering that the divalent anion, sulphate has two sodium cations so the gradient cannot be ascribed to the anion alone. The gradient of ions with a -1 charge was similar to each other, so is not dependent on ion charge density, size or shape.

The effect of different phosphate concentrations on  $\Delta Q$  versus temperature is shown in Figure 4. The change in molar enthalpy relative to pure water of the  $\Delta Q$  versus temperature was dependent on phosphate concentration. The  $\Delta Q$  versus phosphate concentration at different temperatures is shown in Figure 5a and b. It is worth noting that change in molar enthalpy relative to pure water is not linear even at concentrations around 100 mM where there are over 500 water molecules per anion and cation.

The values for the molal volume ( $V_b$ ) and thermal expansion coefficient of pure water ( $\alpha_b$ ) are known from the literature,<sup>36</sup> and  $T$ ,  $\Delta P$  and  $x_s$  values are controlled during each experiment. The variables are  $n$  and molar expansivity ( $V_h\alpha_h$ ).  $V_h$  and  $\alpha_h$  cannot be separated by this approach.  $V_s\alpha_s$  is known for crystalline salts at a set temperature.<sup>38</sup> In this paper we assume the  $V_s\alpha_s$  value for salt and the  $n$  does not vary with temperature calculations. The  $\Delta Q$  values calculated using equation 10 for 100 mM NaCl (Figure 6) is consistent with an  $n$  value of 13.3 and a  $V_h\alpha_h$  value 0.55 times that of the known  $V_b\alpha_b$  for bulk water at temperatures between 5 and 95 °C. The calculated values for all the sodium halide salts using both the single hydration layer and double hydration layer models are shown in Table 1. The  $V_h\alpha_h$  values approach that of pure water as the charge density reduces but are all lower than pure water.

## Discussion

Interpretation of the PPC results for the different salt solutions has to take into account that the measured  $\Delta Q$  is a result of both the sodium and the anion and cannot be separated. The monovalent anions all have the same concentration of sodium so differences between the  $\Delta Q$  values will be due to the anion alone. The second factor that has to be considered is that the measured  $\Delta Q$  is relative

to the pure water in the reference cell. By applying equation 10 the  $V_h\alpha_h$  values for the hydration layer were calculated.

Heat release has been interpreted as bond formation between water molecules<sup>40</sup> so the initial exothermic  $\Delta Q$  can be interpreted as the hydration layer strengthening or forming additional hydrogen bonds relative to pure water. At temperatures over the  $T_i$  value this relationship reverses and  $\Delta Q$  became endothermic can be interpreted as the hydration layer weakening or breaking more hydrogen bonds relative to pure water. The temperature where this transition between bond formation and bond breaking is observed is dependent on charge density of the anion. This is clearly seen for the halides which are simple spheres but the general trend is also evident with the more complex structure of the oxoanions and thiocyanate (which is more linear than spherical). It is possible that PPC is providing evidence about the hydrogen bond population of water around the anions. Electrostatic interaction impact on the hydration layer is also evident as valency of the anion has an obvious effect on the average gradient on the  $\Delta Q$  versus temperature plot though this work does not differentiate between this being driven by the valency of the anion or the number of sodium ions present.

The experimental data where the concentration of phosphate was varied (Figure 4, 5a and 5b) demonstrated that change in molar enthalpy relative to pure water at a temperature below the  $T_i$  value was reduced as the  $x_s$  increased. The non-linearity occurs below 100 mM phosphate suggests that there are longer-range interactions between water and ions that extends beyond the first two layers of water and are probably electrostatic interactions.<sup>10</sup> This does call into question where the  $n$  value should be set so the use of equation 10 should have the assumptions used to select the  $n$  value clearly explained.

The equation for the analysis of PPC data presented in this paper (equation 10) is based on a three component model comprising bulk water (unperturbed water); hydration water (water that is measurably perturbed by the ions) and the ions. Equation 10 accepts that the hydration water has its own molar volume and thermal expansion coefficient that may differ from the bulk water and this should not be ignored. The  $n$  value is dependent on where boundary between the hydration layer and bulk water is set. Molecular dynamic simulation has been used to estimate the  $n$  value around the halides, see Table 1.<sup>38</sup> For example, if the boundary is set at a single layer of water molecules per ion, a sodium and chloride would have an  $n$  value of 13.3; but if a double layer is chosen the  $n$  value will be 34.3. The researcher has a choice of a single thickness or double thickness model when using equation 10 and should remember that the calculated  $V_h\alpha_h$  will be an average for the water molecules within the hydration layer.

The apparent partial molar volumes of water around salts at infinite dilution<sup>32</sup> and the ionic volume<sup>43</sup> has been observed to be quite different for over 50 years,<sup>6</sup> see Figure S1 in the supplementary material. This has been explained as electrostriction, a reduction in the volume of the water interacting with the ion's electrical field and is particularly noticeable for high charge density ions like sodium, and fluoride.<sup>34</sup> The  $V_h\alpha_h$  values calculated are consistently lower than bulk water suggesting that any decrease in the  $V_h$  value is due to electrostriction, the  $\alpha_h$  value or a combination of the two.

Partial molal adiabatic compressibility sodium salts in water at 25°C determined from the velocity of sound through the sample<sup>41</sup> like the molar expansivity of the hydration layer ( $V_h\alpha_h$ ) calculated here



also follows a series that mirrors the Hofmeister series. It is attractive to think the three phenomenon are related and have to do with the interaction of the salt with water.

Franz Hofmeister's original observation was that certain salts were consistent in their ability to precipitate and dissolve proteins.<sup>2-4</sup> Hofmeister put this observation down to the salt absorbing the water. In the 1930s the idea that ions have a long-range effect on the hydrogen bond structure of water to explain the viscosity of electrolyte solutions.<sup>5</sup> This led to the theory of some ions being "structure makers" and others being "structure breakers". Since the 1930s there has been extensive research undertaken to better study the interaction between water and ions using an incredibly diverse collection of analytical techniques including nuclear magnetic resonance, optical Kerr effect spectroscopy, dielectric relaxation spectroscopy, transient vibrational adsorption spectroscopy, terahertz spectroscopy, x-ray scattering and molecular dynamic simulation. The femtosecond pump-probe spectroscopy results first used to challenge the long-range effect of ions on water and came to the conclusion that ions had little effect on water beyond its immediate single molecule thick hydration layer.<sup>8</sup> A hydration layer composed of a single water layer has been used to explain experimental data for a range of techniques including nuclear magnetic resonance,<sup>42</sup> optical Kerr effect spectroscopy,<sup>43</sup> transient vibrational adsorption spectroscopy,<sup>42</sup> terahertz spectroscopy.<sup>44</sup> The problem with the model of water being limited to a single layer of water around the ion is that it failed to explain the Hofmeister effect while destroying the established structure maker structure breaker theory. The new orthodoxy of a single water molecule thick hydration layer has been criticized using data from neutron scattering<sup>9</sup> and femtosecond infrared spectroscopy<sup>45</sup> which detected a second layer of water around ions. When the potential long-range effects of the ion's electrical field is taken into consideration along with the dynamic nature of water the selection of a hydration layer's boundary is probably a measure of each analytical method's sensitivity. PPC can be used to justify either a single or double water layer around ions. It can also be used to challenge the idea of the interaction between water and ions being solely short range. The slope on the  $\Delta Q$  versus phosphate ion concentration plot shown in Figures 5 and 6 did not show linearity as it approached zero concentration. This phenomenon was also observed when determining apparent partial molar volumes of water around salts where an infinite dilution method was needed to estimate the partial molar volume.<sup>34</sup> Both observations could be used to argue for long-range interactions between ions and water beyond a one or two molecule thick hydration layer. While the authors would not argue that structure making structure breaking theory is an adequate explanation for the Hofmeister effect the femtosecond pump-probe spectroscopy evidence that was used to criticise the theory<sup>8</sup> was not as conclusive as it looked at the time.

The majority of the papers that used PPC to study amino acids,<sup>17</sup> small inorganic and organic molecules,<sup>18</sup> proteins<sup>17-28</sup> and DNA<sup>32,33</sup> used the equation derived in Lin et al 2002<sup>17</sup> to calculate the thermal coefficient of expansion of the solvent volume. The authors of this paper suggest that equation 10 is worth applying to PPC studies of diverse solutes in aqueous solutions. Small organic and inorganic molecules as well as macromolecules have hydration layers. There is an increasing body of work using techniques such as terahertz spectroscopy that have been used to detect hydration layers around diverse organic molecules including sugars,<sup>46</sup> peptides<sup>47</sup> and proteins<sup>48-50</sup> suggesting hydration in many cases is not limited to a single layer at the solutes surface but can be extensive. Whether the boundary of the hydration layer around a solute is a single layer, a double layer or more complex this population of water has to be defined in terms of its own average molar volume and thermal expansion coefficient like the water around salts in this paper. In studies using

macromolecules such as globular proteins as the solute, it is also worth noting that the molar expansivity ( $V_s\alpha_s$ ) will have to be estimated from the known partial volumes of the protein's constituent groups<sup>51</sup> and that estimating a value for  $n$  in equation 10 will be problematic with the data available at the time of writing.

## Acknowledgements

The authors would like to thank the Department of Chemical and Biological Engineering at the University of Sheffield for financing Jordan Bye's studentship, and the Engineering and Physical Sciences Research Council (EPSRC) KnowledgeTransfer Account for part funding the purchase of the DSC. The authors would also like to thank Dr James McGregor of the University of Sheffield for his advice into the interpretation of the PPC results.

## References

- 1 S. Arrhenius, *Z. Phys. Chem.* 1887, **1**, 631-648.
- 2 F. Hofmeister, *Arch. Exp. Pathol. Pharmacol.* 1888, **24**, 247-260.
- 3 F. Hofmeister, *Arch. Exp. Pathol. Pharmacol.* 1888, **25**, 1-30.
- 4 W. Kunz, J. Henle, B. W. Ninham, *Curr. Opin. Colloid Interface Sci.* 2004, **9**, 19-37.
- 5 A. Heydweiller, *Annalen der Physik* 1910, **33**, 145-185.
- 6 W. M. Cox, J. H. Wolfenden, *R. Soc. Ser. A* 1934, **92**, 475-488.
- 7 R. W. Gurney, *Ionic Processes in Solution*; McGraw-Hill: New York, 1953.
- 8 A. W. Omta, M. F. Kropman, S. Woutersen, H. J. Bakker, *Science*, 2003, **301**, 347-349.
- 9 R. Mancinelli, A. Botti, F. Bruni, M. A. Ricci, A. K. Soper, *Phys. Chem. Chem. Phys.* 2007, **9**, 2959-2967.
- 10 S. J. Irudayam and R. H. Henchman, *J. Chem. Phys.* 2013, **137**, 034508.
- 11 J. W. Bye, and R. J. Falconer, *J. Phys. Chem. B* 2014, **118**, 4282-4286.
- 12 L. Ter-Minassian, J. C. Petit, N. Vankiet, C. Brunaud, *J. Chim. Phys.* 1970, **67**, 265-269.
- 13 L. Ter-Minassian, F. Milliou, *J. Phys. E*, 1983, **16**, 450-455.
- 14 S. L. Ranzio, *J. Phys. E*, 1984, **11**, 1058-1061.
- 15 S. L. Ranzio, *Thermochim. Acta*, 1997, **300**, 29-41.
- 16 P. Kujawa and F. M. Winnik, *Macromolecules*, 2001, **34**, 4130-4135.
- 17 L. N. Lin, J. F. Brandts, J. M. Brandts, V. Plotnikov, *Anal. Biochem.* 2002, **302**, 144-60.
- 18 J. D. Batchelor, A. Olteanu, A. Tripathy, G. J. Pielak, *J. Am. Chem. Soc.* 2004, **126**, 1958-1961.
- 19 R. Ravindra, R. Winter, *Z. Phys. Chem.* 2003, **217**, 1221-1243.
- 20 W. Dzwolak, R. Ravindra, J. Lendermann, R. Winter, *Biochemistry*, 2003, **42**, 11347-11355.
- 21 C. Nicolini, R. Ravindra, B. Ludolph, R. Winter, *Biophys. J.* 2004, **86**, 1385-1392.
- 22 R. Ravindra, R. Winter, *Chemphyschem*, 2004, **5**, 566-571.
- 23 R. Ravindra, C. Royer, R. Winter, *Phys. Chem. Chem. Phys.* 2004, **8**, 1938-1943.
- 24 L. Mitra, N. Smolin, R. Ravindra, R. Winter, *Phys. Chem. Chem. Phys.* 2006, **8**, 1249-1265.
- 25 R. Winter, D. Lopes, S. Grundzielanek, K. Vogtt, *J. Non-Equil. Thermody.* 2007, **32**, 41-97.
- 26 A. Cooper, D. L. Cameron, J. Jakus, G. W. Pettigrew, *Biochem. Soc. Trans.* 2007, **35**, 1547-1550.

- 27 D. L. Cameron, J. Jakus, S. R. Pauleta, G. W. Pettigrew, A. Cooper, *J. Phys. Chem. B*, 2010, **114**, 16228-16235.
- 28 A. D. Tsamaloukas, N. K. Pyzocha, G. I. Makhatadze, *J. Phys. Chem. B*, 2010, **114**, 16166-16170.
- 29 M. Nazari, H. Y. Fan, H. Heerklotz, *Langmuir*, 2012, **28**, 14129-14136.
- 30 N. Tamai, Y. Nambu, S. Tanaka, M. Goto, H. Matsuki, S. Kaneshina, *Colloid Surface B*, 2012, **92**, 232-239.
- 31 H. Y. Fan, M. Nazari, S. Chowdury, H. Heerklotz, *Langmuir*, 2011, **27**, 1693-1699.
- 32 G. Rayan, A. D. Tsamaloukas, R. B. Macgregor, H. Heerklotz, *J. Phys. Chem. B*, 2009, **113**, 1738-1742.
- 33 A. I. Dragan, D. J. Russel, P. L. Privalov, *Biopolymers*, 2009, **91**, 95-101.
- 34 F. J. Millero, *Chem. Rev.* 1971, **71**, 147-176.
- 35 Y. Marcus, *Ions Properties*, CRC Press, Boca Raton, 1997.
- 36 G. S. Kell, *J. Chem. Eng. Data*, 1967, **12**, 66-69.
- 37 H. Ohtaki, T. Radai, *Chem. Rev.* **93**, 1157-1204.
- 38 S. H. Lee, J. C. Rasaiah, *J. Phys. Chem.* 1996, **100**, 1420-1425.
- 39 D. R. Lide *CRC Handbook of Chemistry and Physics*, Ed. 88, CRC Press, Boca Raton, 2008.
- 40 B. Hribar, N. T. Southall, V. Vlachy, K. A. Dill, *J. Am. Chem. Soc.* 2002, **124**, 12302-12311.
- 41 F. J. Millero, J. D. Sharp, *J. Chem. Eng. Data*, 2013, **58**, 3458-3463.
- 42 H. J. Bakker, *Chem. Rev.* 2008, **108**, 1456-1473.
- 43 I. A. Heisler, K. Mazur, S. R. Meech *J. Phys. Chem. B* 2011, **115**, 1863-1973.
- 44 M. Heyden, E. Bründermann, U. Heugen, G. Niehues, D. M. Leitner, M. Havenith, *J. Am. Chem. Soc.* 2008, **130**, 5773-5779.
- 45 K. J. Tielrooij, N. Garcia-Araez, M. Bonn, H. J. Bakker, *Science* 2010, **328**, 1006-1009.
- 46 M. Heyden, E. Bründermann, U. Heugen, G. Niehues, D. M. Leitner, M. Havenith, *J. Am. Chem. Soc.* 2008, **130**, 5773-5779.
- 47 T. Ding, R. Li, J. A. Zeitler, T. L. Huber, L. F. Gladden, A. P. J. Middelberg, R. J. Falconer, *Opt. Express*, 2010 **18**, 27431-27444.
- 48 S. Ebbinghaus, S. J. Kim, M. Heyden, X. Yu, U. Heugen, M. Gruebele, D. M. Leitner, M. Havenith, *M. Proc. Natl. Acad. Sci.* 2007, **104**, 20749-20752
- 49 B. Born, H. Weingärtner, E. Bründermann, M. Havenith, *J. Am. Chem. Soc.* 2009, **131**, 3752-3755.
- 50 J. W. Bye, S. C. Meliga, D. Ferachou, J. A. Zeitler, R. J. Falconer, *J. Phys. Chem. A* 2014, **118**, 83-88.
- 51 G. I. Makhatadze, V. N. Medvedkin, P. L. Privalov, *Biopolymers* 1990, **30**, 1001-1010.

Tables

Table 1 The number of water molecules in the 1<sup>st</sup> and 2<sup>nd</sup> water layers around sodium ( $n$ ) and halide ions, molar expansivity of a salt ( $V_s\alpha_s$ ), the relationship between the average molar expansivity of water in the hydration layer and the bulk phase ( $V_h\alpha_h / V_b\alpha_b$ ) and the constant  $A$  for NaF, NaCl, NaBr and NaI. The later two values derived from 100 mM experimental data using equation 10.

1 layer model	$n^a$	$V_s\alpha_s(\text{m}^3\text{mol}^{-1}\text{K}^{-1})^b$	$V_h\alpha_h / V_b\alpha_b^c$	$A(\text{J})$
NaF	12.3	$1.48 \times 10^{-9}$	0.54	$7.35 \times 10^{-7}$
NaCl	13.3	$3.16 \times 10^{-9}$	0.55	$6.60 \times 10^{-7}$
NaBr	13.1	$4.46 \times 10^{-9}$	0.62	$7.05 \times 10^{-7}$
NaI	14.0	$5.51 \times 10^{-9}$	0.65	$7.80 \times 10^{-7}$
2 layer model				
NaF	34.3	$1.48 \times 10^{-9}$	0.83	$7.50 \times 10^{-7}$
NaCl	40.8	$3.16 \times 10^{-9}$	0.85	$6.75 \times 10^{-7}$
NaBr	40.5	$4.46 \times 10^{-9}$	0.87	$7.50 \times 10^{-7}$
NaI	46.1	$5.51 \times 10^{-9}$	0.89	$8.10 \times 10^{-7}$

<sup>a</sup> The number of water molecules per sodium and halide ions was determined by MD simulation.<sup>40</sup>

<sup>b</sup> The  $V_s\alpha_s$  is for crystalline sodium halides.<sup>41</sup>

<sup>c</sup> The  $V_b\alpha_b$  is derived from pure water density measurements published in Kell, 1967.<sup>38</sup> The  $V_h\alpha_h / V_b\alpha_b$  and  $A$  values were calculated using equation 10.

## Figure Titles

*Figures 1a-d: Figures 1a-c show the raw data from a pressure perturbation scan of 1000 mM disodium hydrogen phosphate, sodium chloride and sodium thiocyanate in pure water at pH 7 respectively. Alternating pressure pulses from 1-5 bar and then 5-1 bar were applied to the sample at 1 °C intervals from 7 °C – 92 °C, with a heating rate of 0.1 °C/min. Figure 1d shows a close up of figure 1a from 70 °C to 75 °C, with data for pressure increases and decreases shown (---). Here the area under the power spike for the pressure pulses was integrated by the software provided to calculate the energy absorbed and released by the sample relative to the reference cell containing pure water. For the pressurization steps shown in figure 1d the change in heat is positive (endothermic) with reference to the pure water baseline. During the depressurization steps the heat change is negative (exothermic) with reference to the pure water baseline.*

*Figure 2a: Heat changes for pressure increase (1 to 5 bar) from 7 °C – 92 °C for (a) 100 mM and (b) 1000 mM disodium hydrogen phosphate (●), disodium sulphate (■), sodium fluoride (✕), sodium chloride (▲), sodium bromide (⊕), sodium iodide (⊙), sodium thiocyanate (⊖) and sodium perchlorate (▼). The sample size was 300 µL. Lines between data points do not represent experimental data and are only shown for guidance.*

*Figure 3: Ion surface charge density against the temperature ( $T_i$ ) where the  $\Delta Q$  equals zero for 100 mM disodium sulphate (■), sodium fluoride (✕), sodium chloride (▲), sodium bromide (⊕), sodium iodide (⊙), and sodium perchlorate (▼). Note thiocyanide was omitted as it is not spherical.*

*Figure 4: Heat changes for pressure increase (1-5 bar) from 7 °C – 92 °C at different concentrations of disodium hydrogen phosphate; 50 (◆), 100 (▲), 250 (●), 500 (▼), 750 (■) and 1000 mM (●). The sample size was 300 µL.*

*Figure 5: Heat changes for pressure increase (1-5 bar) at different concentrations of disodium hydrogen phosphate and different temperatures; (a) 11.5 (●), 21.5 (■), 31.5 (▲), (b) 41.5 (●), 61.5 (◆), 71.5 (▲), 81.5 (✕) and 91.5 °C (▼). The sample size was 300 µL.*

*Figure 6: Experimental values for the  $\Delta Q$  for 100 mM NaCl fitted against values for  $\Delta Q$  calculated using equation 10 using the assumptions that  $n$  is 13.3 and the molar expansivity of the hydration layer is 0.55 that of pure water.*

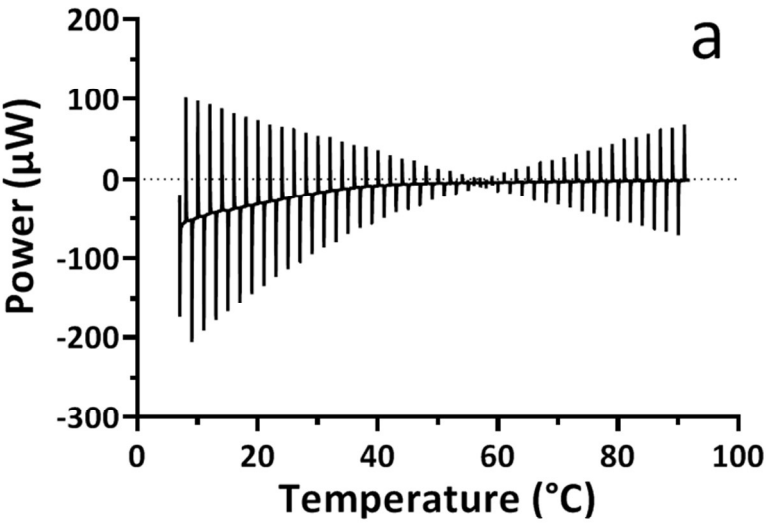


Figure 1a  
254x190mm (96 x 96 DPI)



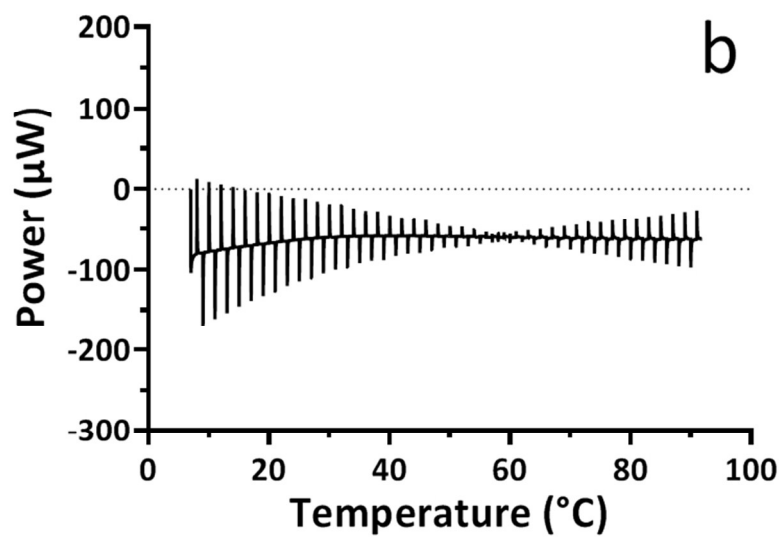


Figure 1b  
254x190mm (96 x 96 DPI)

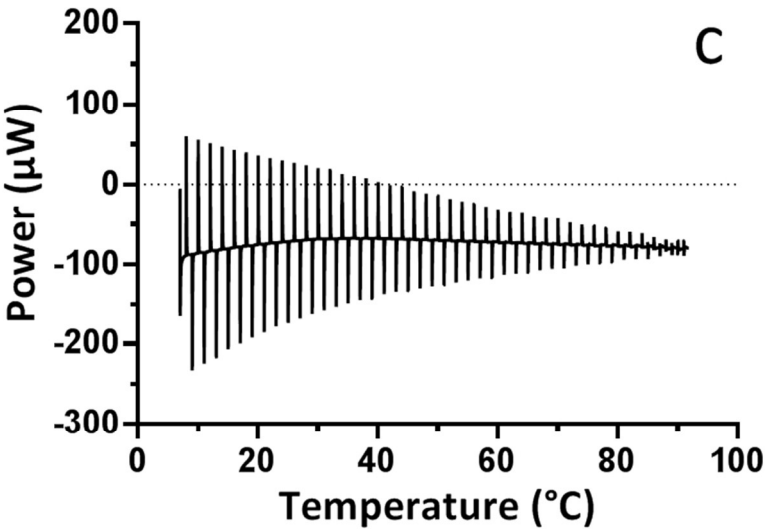


Figure 1c  
254x190mm (96 x 96 DPI)

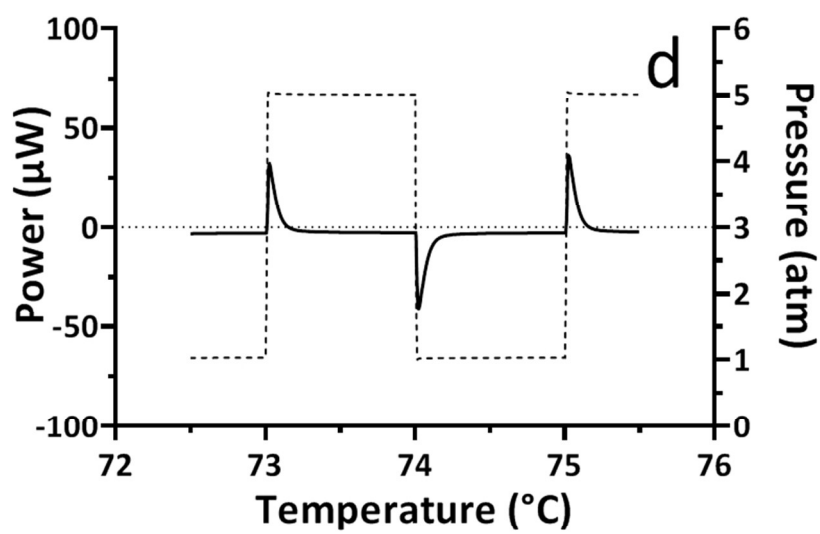


Figure1d  
254x190mm (96 x 96 DPI)

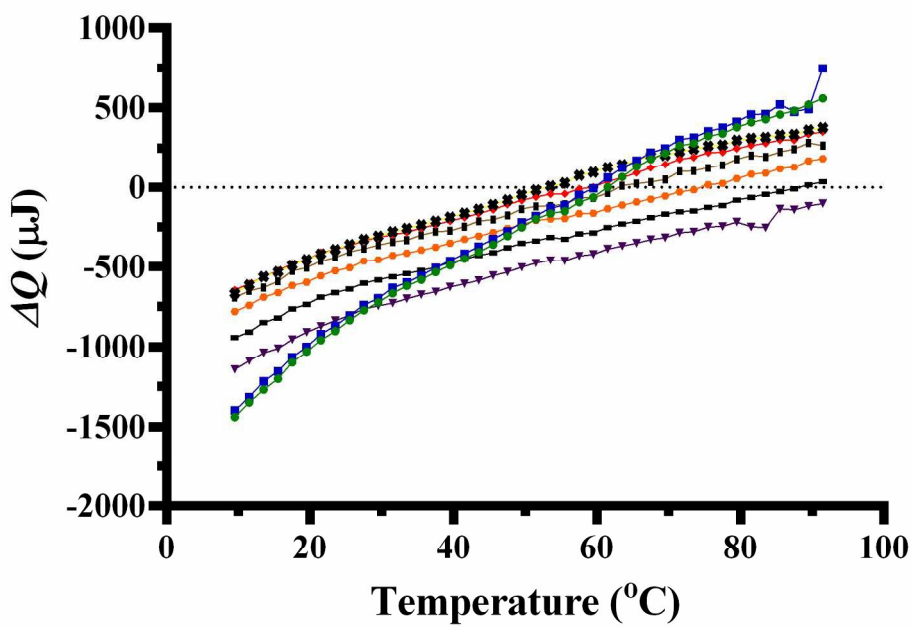


Figure 2a  
187x125mm (300 x 300 DPI)

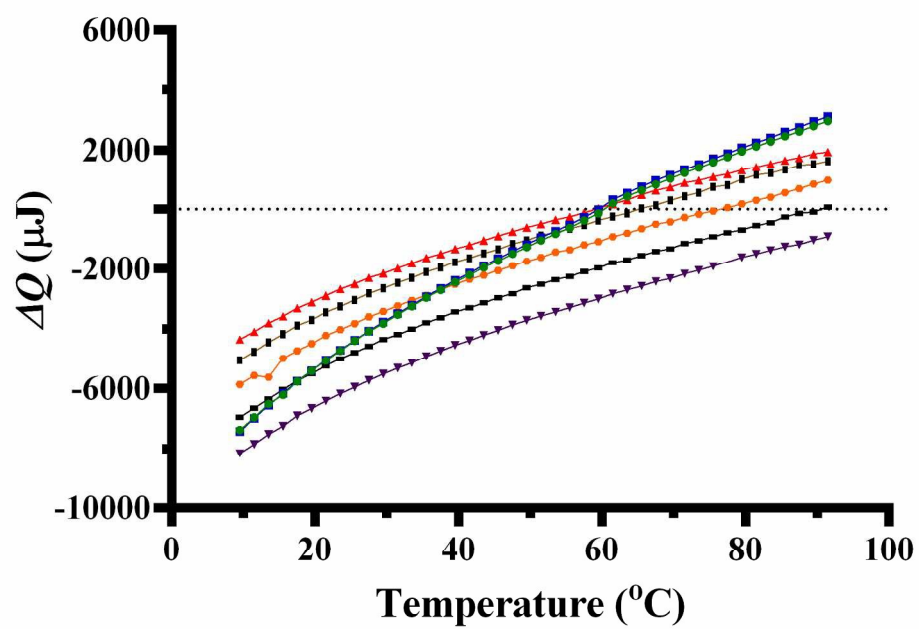


Figure 2b  
187x125mm (300 x 300 DPI)

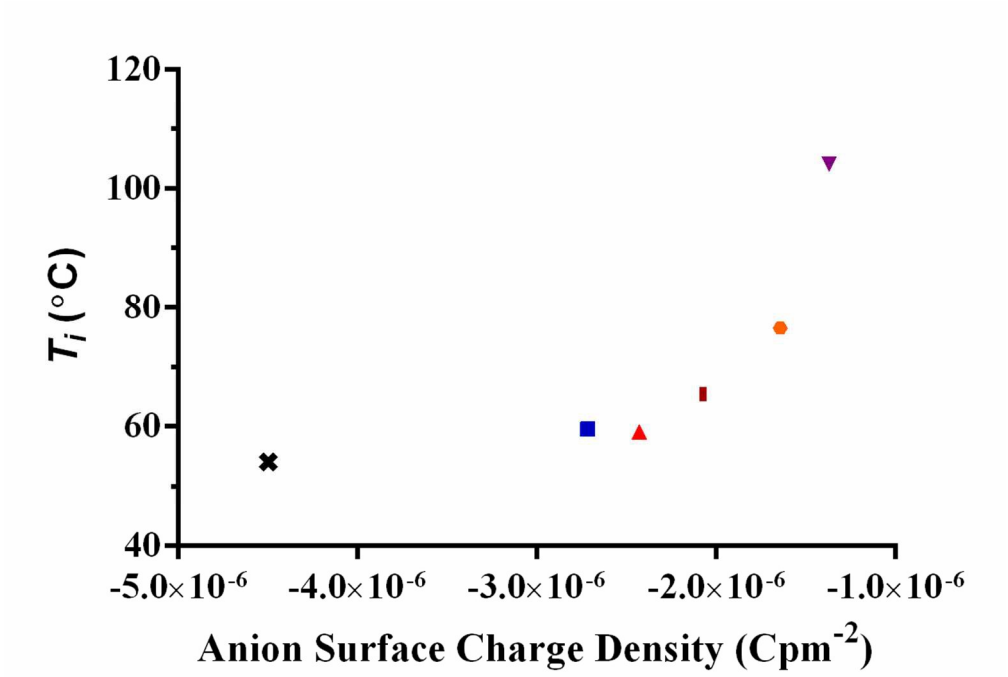


Figure 3  
106x74mm (300 x 300 DPI)



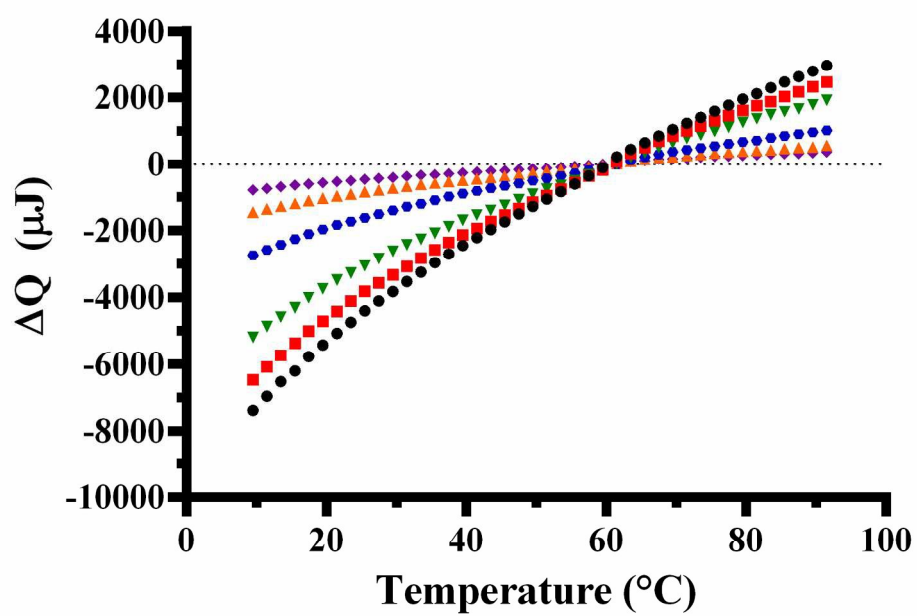


Figure 4  
182x119mm (300 x 300 DPI)

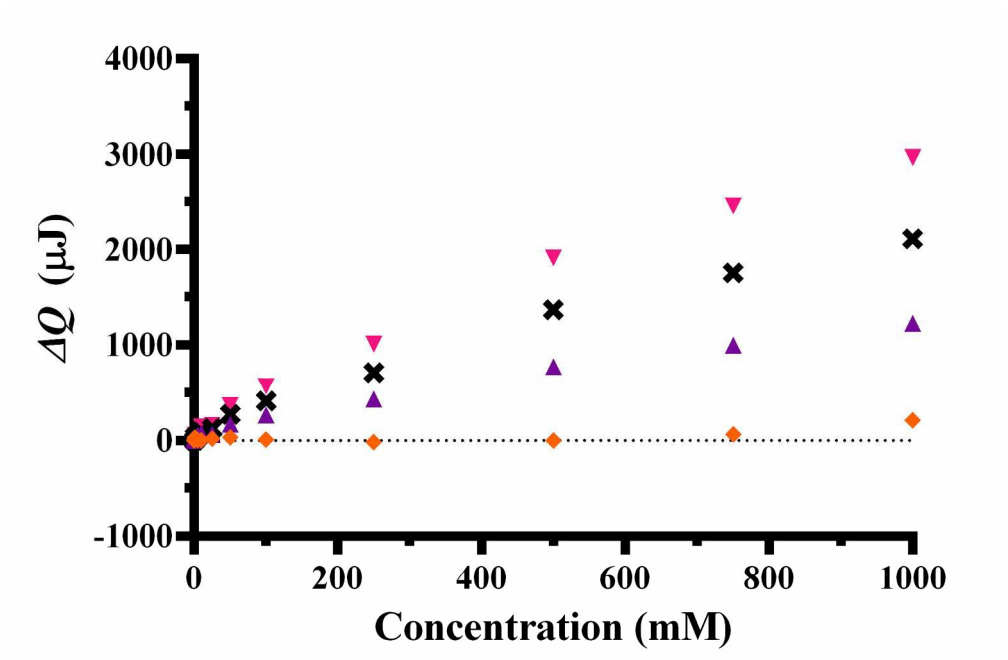


Figure 5a  
187x124mm (300 x 300 DPI)

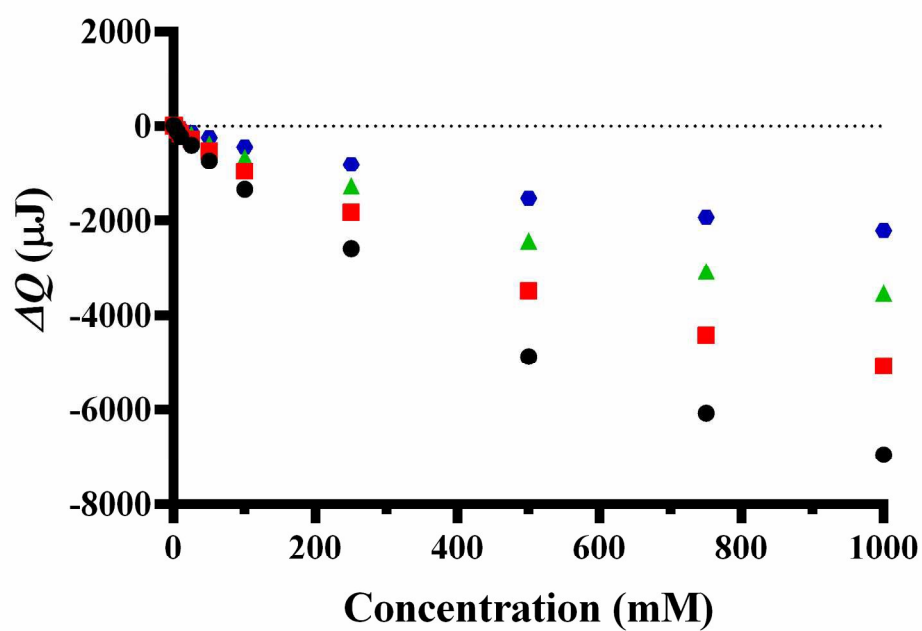


Figure 5b  
181x122mm (300 x 300 DPI)

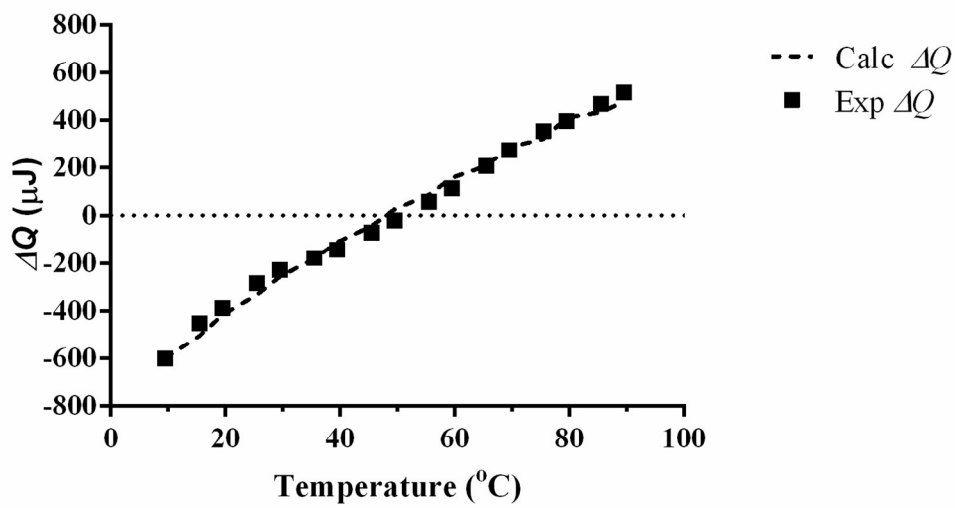


Figure 6  
133x74mm (300 x 300 DPI)