

11.8: Colligative Properties of Ionic Solutes

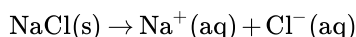
Learning Objective

- Determine the colligative properties of solutions of ionic solutes.

Previously, we considered the colligative properties of solutions with molecular solutes. What about solutions with ionic solutes? Do they exhibit colligative properties?

There is a complicating factor: ionic solutes separate into ions when they dissolve. This increases the total number of particles dissolved in solution and *increases the impact on the resulting colligative property*. Historically, this greater-than-expected impact on colligative properties was one main piece of evidence for ionic compounds separating into ions (increased electrical conductivity was another piece of evidence).

For example, when NaCl dissolves, it separates into two ions:



This means that a 1 M solution of NaCl actually has a net particle concentration of 2 M. The observed colligative property will then be twice as large as expected for a 1 M solution.

It is easy to incorporate this concept into our equations to calculate the respective colligative property. We define the **van 't Hoff factor** (i) as the number of particles each solute formula unit breaks apart into when it dissolves. Previously, we have always tacitly assumed that the van 't Hoff factor is simply 1. But for some ionic compounds, i is not 1, as shown in Table 11.8.1.

Table 11.8.1 Ideal van 't Hoff Factors for Ionic Compounds

Compound	i
NaCl	2
KBr	2
LiNO ₃	2
CaCl ₂	3
Mg(C ₂ H ₃ O ₂) ₂	3
FeCl ₃	4
Al ₂ (SO ₄) ₃	5

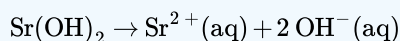
The ideal van 't Hoff factor is equal to the number of ions that form when an ionic compound dissolves.

✓ Example 11.8.1

Predict the van 't Hoff factor for Sr(OH)₂.

Solution

When Sr(OH)₂ dissolves, it separates into one Sr²⁺ ion and two OH⁻ ions:



Because it breaks up into three ions, its van 't Hoff factor is 3.

? Exercise 11.8.1

What is the van 't Hoff factor for Fe(NO₃)₃?

Answer

4

It is the "ideal" van 't Hoff factor because this is what we expect from the ionic formula. However, this factor is usually correct only for dilute solutions (solutions less than 0.001 M). At concentrations greater than 0.001 M, there are enough interactions between ions of opposite charge that the net concentration of the ions is less than expected—sometimes significantly. The actual van 't Hoff factor is thus less than the ideal one. Here, we will use ideal van 't Hoff factors.

Revised equations to calculate the effect of ionization are then easily produced:

$$\Delta T_b = imK_b$$

$$\Delta T_f = imK_f$$

$$\Pi = iMRT$$

where all variables have been previously defined. To calculate vapor pressure depression according to Raoult's law, the mole fraction of solvent particles must be recalculated to take into account the increased number of particles formed on ionization.

✓ Example 11.8.2

Determine the freezing point of a 1.77 *m* solution of NaCl in H₂O.

Solution

For NaCl, we need to remember to include the van 't Hoff factor, which is 2. Otherwise, the calculation of the freezing point is straightforward:

$$\Delta T_f = (2)(1.77\text{ }m)(1.86\text{ }^{\circ}\text{C}/m) = 6.58\text{ }^{\circ}\text{C}$$

This represents the change in the freezing point, which is decreasing. So we have to subtract this change from the normal freezing point of water, 0.00°C:

$$0.00 - 6.58 = -6.58^{\circ}\text{C}$$

? Exercise 11.8.2

Determine the boiling point of a 0.887 *m* solution of CaCl₂ in H₂O.

Answer

$$101.36\text{ }^{\circ}\text{C}$$

✓ Food and Drink App: Salting Pasta Cooking Water

When cooking dried pasta, many recipes call for salting the water before cooking the pasta. Some argue—with colligative properties on their side—that adding salt to the water raises the boiling point, thus cooking the pasta faster. Is there any truth to this?



Figure 11.8.1: Cooking dried pasta. Why do so many recipes call for adding salt to water when boiling pasta? Is it to raise the boiling temperature of the water? (Public Domain; Jan Vašek via pixabay)

To judge the veracity of this claim, we can calculate how much salt should be added to the water to raise the boiling temperature by 1.0°C, with the presumption that dried pasta cooks noticeably faster at 101°C than at 100°C (although a 1° difference may make only a negligible change in cooking times). We can calculate the molality that the water should have:

$$1.0^{\circ}\text{C} = m(0.512^{\circ}\text{C}/m)m = 1.95$$

We have ignored the van 't Hoff factor in our estimation because this obviously is not a dilute solution. Let us further assume that we are using 4 L of water (which is very close to 4 qt, which in turn equals 1 gal). Because 4 L of water is about 4 kg (it is actually slightly less at 100°C), we can determine how much salt (NaCl) to add:

$$4 \text{ kg } \cancel{H_2O} \times \frac{1.95 \text{ mol } \cancel{NaCl}}{\text{kg } \cancel{H_2O}} \times \frac{58.5 \text{ g } NaCl}{1 \text{ mol } \cancel{NaCl}} = 456.3 \text{ g } NaCl$$

This is just over 1 lb of salt and is equivalent to nearly 1 cup in the kitchen. In your experience, do you add almost a cup of salt to a pot of water to make pasta? Certainly not! A few pinches, perhaps one-fourth of a teaspoon, but not almost a cup! It is obvious that the little amount of salt that most people add to their pasta water is not going to significantly raise the boiling point of the water.

So why do people add some salt to boiling water? There are several possible reasons, the most obvious of which is taste: adding salt adds a little bit of salt flavor to the pasta. It cannot be much because most of the salt remains in the water, not in the cooked pasta. However, it may be enough to detect with our taste buds. The other obvious reason is habit; recipes tell us to add salt, so we do, even if there is little scientific or culinary reason to do so.

Summary

For ionic solutes, the calculation of colligative properties must include the fact that the solutes separate into multiple particles when they dissolve. The equations for calculating colligative properties of solutions of ionic solvents include the van 't Hoff factor, i .

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