Determination of the Solubility Limit of Tris(dibenzylideneacetone) dipalladium(0) in Tetrahydrofuran/Water Mixtures

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Supporting Information

ABSTRACT: Determination of the solubility limit of a strongly colored organometallic reagent in a mixed-solvent system provides an example of quantitative solubility measurement appropriate to understand polymer, nanoparticle, and other macromolecular aggregation processes. The specific example chosen involves a solution of tris(dibenzylideneacetone) dipalladium(0), Pd2(dba)3, in THF/H2O mixtures, which has the desired properties of high absorption in the visible range (λ_max = 528 nm) and significant difference in solubility in the two solvents THF and H2O. Pd2(dba)3 is readily available, safe, and commonly used in nanoparticle syntheses. The application of Flory–Huggins theory permits the fit of a quantitative model to data obtained from a straightforward laboratory experiment that can be conducted in less than 3 h. The absorbance of this compound is easily measured by UV–vis absorption spectroscopy. The application of the Beer–Lambert law to the data in this experiment is a useful exercise for students because the absorbances span three orders of magnitude.

KEYWORDS: Upper-Division Undergraduate, Laboratory Instruction, Physical Chemistry, Hands-On Learning/Manipulatives, Computer-Based Learning, Solutions/Solvents, Thermodynamics, Nanotechnology, Colloids, UV-Vis Spectroscopy

The study of molecular solubility and the study of solvent mixtures are both classic problems in physical chemistry taught to undergraduate students. However, the solubility of solutes in mixed organic−aqueous solution is not commonly discussed, although it has increasing application in the synthesis of nanoparticles, polymer synthesis, metal extraction, and crystallization. Control of aggregation by means of the solvent conditions is important in the growing field of nanoparticle synthesis.1−11 The solubility of organic molecules relevant to polymer synthesis has been considered most often in the context of Flory–Huggins theory.12,13 The application of theory to the solubility of organic molecules differs from that of ions, which has been considered in detail owing to the many practical applications, such as extraction of metals from ore.14 The increase in metal ion solubility in solvent mixtures arises largely from specific coordination by solvents (e.g., acetone or coordinating Ag+).15,16 Counter ion effects have also been considered, as well as exceptions to solubility rules.17 Crystallization, on the other hand, involves balancing solubility at the point where controlled growth competes effectively with aggregation. This can be done by introducing a second solvent to reduce the solubility.18,19 The solvent with poor ability to dissolve the solute is known as the nonsolvent. Because of the need to train scientists to understand the technological significance, there are numerous recent examples of laboratory demonstrations for understanding solubility in complex systems.20−22 The experiment described in this article is aimed at understanding this phenomenon and related solubility rules.

We have chosen a material that is known to form both nano- and microcrystals in solution, the organometallic reagent trisdibenzylideneacetone dipalladium(0), Pd2(dba)3, which is best known as a means of delivering active Pd in solution for homogeneous catalysis of bond-forming reactions such as the Heck reaction, Sonogashira coupling, Negishi coupling, Carroll rearrangement, and Trost asymmetric allylic alkylation.27−29 However, metallic Pd is formed by allylic and phosphine groups in organic solvents and not by the THF/H2O mixture, which is relatively inert. The precipitate that is discarded in this exercise contains many nanoparticles, which are similar in structure23−25 to those studied elsewhere.5,7,8 Pd2(dba)3 is best known as a means of delivering active Pd in solution for homogeneous catalysis of bond-forming reactions such as the Heck reaction, Sonogashira coupling, Negishi coupling, Carroll rearrangement, and Trost asymmetric allylic alkylation.27−29 However, metallic Pd is formed by allylic and phosphine groups in organic solvents and not by the THF/H2O mixture, which is relatively inert. The precipitate that is discarded in this exercise contains many nanoparticles, which are similar in structure23−25 to those studied elsewhere.5,7,8 Pd2(dba)3 is best known as a means of delivering active Pd in solution for homogeneous catalysis of bond-forming reactions such as the Heck reaction, Sonogashira coupling, Negishi coupling, Carroll rearrangement, and Trost asymmetric allylic alkylation.27−29 However, metallic Pd is formed by allylic and phosphine groups in organic solvents and not by the THF/H2O mixture, which is relatively inert. The precipitate that is discarded in this exercise contains many nanoparticles, which are similar in structure23−25 to those studied elsewhere.5,7,8 Pd2(dba)3 is best known as a means of delivering active Pd in solution for homogeneous catalysis of bond-forming reactions such as the Heck reaction, Sonogashira coupling, Negishi coupling, Carroll rearrangement, and Trost asymmetric allylic alkylation.27−29 However, metallic Pd is formed by allylic and phosphine groups in organic solvents and not by the THF/H2O mixture, which is relatively inert. The precipitate that is discarded in this exercise contains many nanoparticles, which are similar in structure23−25 to those studied elsewhere.5,7,8 Pd2(dba)3 is best known as a means of delivering active Pd in solution for homogeneous catalysis of bond-forming reactions such as the Heck reaction, Sonogashira coupling, Negishi coupling, Carroll rearrangement, and Trost asymmetric allylic alkylation.27−29 However, metallic Pd is formed by allylic and phosphine groups in organic solvents and not by the THF/H2O mixture, which is relatively inert. The precipitate that is discarded in this exercise contains many nanoparticles, which are similar in structure23−25 to those studied elsewhere.5,7,8 Pd2(dba)3 is best known as a means of delivering active Pd in solution for homogeneous catalysis of bond-forming reactions such as the Heck reaction, Sonogashira coupling, Negishi coupling, Carroll rearrangement, and Trost asymmetric allylic alkylation.27−29 However, metallic Pd is formed by allylic and phosphine groups in organic solvents and not by the THF/H2O mixture, which is relatively inert. The precipitate that is discarded in this exercise contains many nanoparticles, which are similar in structure23−25 to those studied elsewhere.5,7,8 Pd2(dba)3 is best known as a means of delivering active Pd in solution for homogeneous catalysis of bond-forming reactions such as the Heck reaction, Sonogashira coupling, Negishi coupling, Carroll rearrangement, and Trost asymmetric allylic alkylation.27−29 However, metallic Pd is formed by allylic and phosphine groups in organic solvents and not by the THF/H2O mixture, which is relatively inert. The precipitate that is discarded in this exercise contains many nanoparticles, which are similar in structure23−25 to those studied elsewhere.5,7,8
solubility. Pd$_2$(dba)$_3$ is a strongly colored organometallic molecule that shows a wide range of solubility in THF/H$_2$O mixtures, ranging from soluble in pure THF to virtually insoluble in pure H$_2$O. Thus, THF and H$_2$O are the solvent and nonsolvent, respectively, in this system. In the experiment described here, the solubility of an organometallic molecule in a mixed-solvent system illustrates solubility in mixed solvents as it applies to aggregation and crystallization. The solubility properties of organometallic Pd$_2$(dba)$_3$ are applicable to a wide range of organic molecules that lack the conveniently observable UV-vis absorption band. The solutions that can be easily prepared in this exercise permit direct observation of rapid aggregation leading to precipitation, on the one hand, and slow aggregation leading to crystal formation, on the other. Finally, because THF/H$_2$O solutions are somewhat unstable and tend to lose Pd(0) to form spherical nanoparticles, there is interesting behavior in the formation of precipitates or mixtures of crystals. These can be observed readily under a light microscope. Consequently, Pd$_2$(dba)$_3$ in THF/H$_2$O mixtures provide an excellent example relevant to particle formation, which can be incorporated into the observations recorded in the laboratory experiment. Such a simple system that exemplifies all of these aspects of the important role played by the nonsolvent has practical benefit for training students. The THF/H$_2$O two-component mixture is widely studied as an example of a “closed-loop” system, which has both an upper and lower critical point such that the two-phase region is completely surrounded by single-phase regions. The critical point is above 70 °C in THF/H$_2$O at 1 atm of pressure so that the complication of the critical behavior can be avoided in the laboratory exercise. The laboratory example is a much-needed study of organic solubility that complements the more commonly studied ionic solubility.

## THEORY

The theory of solubility of organic compounds in two-component mixtures is related to the theory of aggregation, which branches into the theory of micelle formation, polymer formation, and crystallization. One can write the equilibrium as

\[ N \text{Pd}_2(\text{dba})_3(\text{THF}/\text{H}_2\text{O soln}) \rightleftharpoons [\text{Pd}_2(\text{dba})_3]_N (N\text{-mer aggregate}) \] (1)

where \( N \) is the stoichiometric coefficient and the associated aggregation Gibbs energy is \( \Delta_{ag}G_N \). The aggregate may remain in a suspension or may precipitate. The suspending is given by

\[ \text{[Pd}_2(\text{dba})_3]_N (\text{suspension}) \rightarrow \text{[Pd}_2(\text{dba})_3]_N (\text{precipitate}) \] (2)

The suspending of the \([\text{Pd}_2(\text{dba})_3]_N \) aggregate is not governed by an equilibrium constant as the aggregate is intrinsically unstable with respect to solidification. Thus, there is an important distinction that should be made between a solution and a suspension. This is extremely relevant in the chemical literature today given the number of nanoparticle syntheses that cross the boundary from a solution to a suspension as part of the protocol for synthesis.

The rules of colloid chemistry determine the time scale on which aggregates will remain suspended. Derjaguin—Landau—Verwey—Overbeek (DLVO) theory, the theory of colloidal stability, is based on the role of charge stabilization as a balance to ubiquitous van der Waals forces that lead to flocculation and aggregation. Since \([\text{Pd}_2(\text{dba})_3]_N \) is a neutral aggregate, there is only a van der Waals force without a balancing electrostatic repulsion term and rapid precipitation is expected. Visual observation confirms that precipitation is rapid in this system. The aggregation equilibrium provides an intuitive picture of formation of a “nanoparticle” that is no longer soluble and hence precipitates, but the theory of mixed solvents is complicated in this case. On the other hand, the theory of solubility in mixed solvents provides a quantitative explanation that relates to the solubility in the pure solvents, without explicitly mentioning how the solute is destabilized. Hence, Flory—Huggins and aggregation equilibrium—DLVO theory provide two complementary views to the Gibbs energy that governs solubility.

## THE AGGREGATION EQUILIBRIUM

The fact that Pd$_2$(dba)$_3$ forms organic crystals, as observed by transmission electron microscopy (TEM) and light microscopy, makes this system a test system for exploring the factors that govern the aggregation equilibria and solution stability in mixed solvents. Thus, Pd$_2$(dba)$_3$ in THF/H$_2$O solutions provides a range of solution behaviors with which to test solubility, suspendability, and crystal formation in the laboratory. The thermodynamics of aggregation due to the hydrophobic effect can be expressed in terms of an equilibrium constant for N-mer aggregate formation. It is understood that this aggregate, once formed, is metastable according to DLVO theory. Thus, we expect that on some time scale precipitation will occur. First, we will derive the aggregation equilibrium and then we will show a complementary view based on Flory—Huggins theory.

The chemical potential of the solute, \( \mu_1 \), and the N-mer aggregate, \( \mu_N \), with activities \( a_1 \) and \( a_N \), respectively, in a two-component mixture can be written as

\[
\mu_1 = \mu_1^o + k_B T \ln a_1
\] (3)

\[
\mu_N = \mu_N^o + k_B T \ln a_N
\] (4)

where \( k_B \) is the Boltzmann constant and \( T \) is the absolute temperature. The Gibbs energy change for the reaction, eq 1, is

\[
\Delta_{ag}G_N = \mu_N^o - N\mu_1^o
\] (5)

At equilibrium \( \Delta_{ag}G_N = 0 \), and we can write

\[
\mu_N^o + k_B T \ln a_N = N(\mu_1^o + k_B T \ln a_1)
\] (6)

If we define \( \Delta_{ag}G_N^o = \mu_N^o - N\mu_1^o \), then the expression becomes

\[
\ln a_1 = \left( \frac{[\ln a_N]}{N} + \frac{[\Delta_{ag}G_N^o]}{(Nk_B T)} \right)
\] (7)

which determines the extents of aggregation of N-mers in equilibrium with monomers. Technically speaking, the aggregate is in a suspension and not a solution. If the aggregate is sufficiently large or colloidally unstable, it will precipitate. Because N-mer is removed from suspension, the equilibrium in eq 1 is shifted toward the right. At some point the solubility limit is reached and no further precipitation occurs. At this point, we can consider the system a solution, rather than a suspension. We define the solubility limit as the maximum concentration of the monomer solute for which a solution occurs.

The Pd$_2$(dba)$_3$ unit cell is approximately $1.3 \times 1.3 \times 1.5 \text{ nm}^3$ or 2.5 nm$^3$. In pure THF, a typical size for a hexagonal aggregate of Pd$_2$(dba)$_3$ is 500 nm in diameter, in 50% THF/50% H$_2$O.
H₂O,⁵⁻,⁷⁻,⁸⁻,²³⁻,²⁵ giving a volume of 2 × 10⁻⁷ nm³ composed of ~10⁴ monomers. Because Pd₂(dba)₃ is virtually insoluble in pure H₂O,⁶⁻ it is difficult to assess the magnitude of N in that solvent. However, we have observed a decrease in aggregate size as the solvent hydrophobicity increases, which may be a result of the greater solubility of the monomer. The decrease in aggregate size is analogous to the disappearance of a well-defined micelle for surfactants in nonaqueous solvents.⁵⁻ The aggregation and colloidal instability of Pd₂(dba)₃ in THF/H₂O solutions appears to be comparable to the stability of polymers in solvent mixtures. These considerations justify the use of Flory—Huggins theory to model the solubility limit of Pd₂(dba)₃ in equilibrium with its aggregates.

**SOLUBILITY OF SOLUTES IN MIXED-SOLVENT SYSTEMS: FLORY—HUGGINS THEORY**

Pd₂(dba)₃ monomer is observed to be soluble only in the THF fraction, and hence, Pd₂(dba)₃ aggregates will tend to precipitate as the volume fraction of H₂O increases in a mixed-solvent system. Starting with the standard Gibbs energies of solvation of Pd₂(dba)₃ in pure H₂O and THF, which are Δ_solG_W° and Δ_solG_THF°, respectively, the Gibbs energy for solution in the mixture, Δ_solG_mix°, can be expressed using Flory—Huggins theory

\[
\Delta_{sol}G_{mix}°/(RT) = \phi_W\Delta_{sol}G_W°/(RT) + \phi_{THF}\Delta_{sol}G_{THF}°/(RT)
\]

\[
+ \phi_W\ln \phi_W + \phi_{THF}\ln \phi_{THF} + \chi_{W,THF}\phi_W\phi_{THF}
\]

where ϕ_W = V_W/V and ϕ_{THF} = V_{THF}/V are the volume fractions of water and THF, respectively, and \(\chi_{W,THF}\) is the Flory—Huggins interaction parameter for THF and H₂O.¹²⁻,³⁸⁻,⁴⁰ By knowing the solubility in THF, we can calculate

\[
\ln a_1 = -\Delta_{sol}G_{THF}°/(RT)
\]

in the mixed-solvent system. Given estimates of Δ_{sol}G_{THF}° = 12.2 kJ/mol and Δ_{sol}G_W° = 42.5 kJ/mol, based on the experiment shown below, combined with an estimate of 1.5 for the Flory—Huggins parameter of a THF/H₂O mixture, we can calculate the theoretical solubility plot in Figure 1. The plot of solubility is linear on a semi-log plot in accord with eq 9, which makes a simple presentation for the student.

**EXPERIMENTAL PROCEDURE**

For measurements of the solubility limit, 73.2 mg of tris(dibenzyldieneacetone dipalladium(0), Pd₂(dba)₃ (STREM catalog no. 51364-51-3, Mᵣ = 915.7), was dissolved in 20 mL of anhydrous tetrahydrofuran (THF) to make a 4 mM stock solution. When employed as a catalyst, Pd₂(dba)₃ is often mixed immediately with H₂O and exposed to air, so that a glove box is not necessary. The resulting solution had a deep purple color. Solutions of varying percentage of THF and H₂O were prepared by quickly pipetting an appropriate volume of the 4 mM stock solution of Pd₂(dba)₃ in THF into vials containing H₂O to make THF/H₂O mixtures ranging from 90% THF to 10% THF with total volume of 2 mL. The nominal Pd₂(dba)₃ concentration obtained was equal to (THF%/100) × 4 mM. For example, a 10% THF/90% H₂O solution would correspond to 3.60 mM Pd₂(dba)₃. If Pd₂(dba)₃ were completely soluble. However, a precipitate was immediately observed in all solutions studied. Therefore, the mixtures were immediately centrifuged at 10,000 rpm for 5 min, the precipitate was removed, and the supernatant containing a saturated solution of Pd₂(dba)₃ was used for the solubility measurements. Filtration of the precipitates through a Whatman 0.02 μm filter gives results that are similar to those obtained by centrifugation.

Absorption spectra of the supernatant from the centrifuged solutions were measured using a HP8542 photodiode array spectrophotometer. Measurements under a variety of conditions showed that exposure to oxygen for periods of many hours or even days did not affect the spectra.

To convert the absorbance measurements to concentration using the Beer—Lambert law, the extinction coefficient, ε, of the 528 nm (18,940 cm⁻¹) metal-to-ligand charge transfer (MLCT) absorption band of Pd₂(dba)₃ shown in Figure 2 was determined to be 8,970 L mol⁻¹ cm⁻¹ by repeated measurements of the absorbance using quantities of Pd₂(dba)₃ ranging from 4.6 to 70.3 mg dissolved in 20 mL of THF solution and assuming complete dissolution of Pd₂(dba)₃. This value is significantly lower than the 30,000 L mol⁻¹ cm⁻¹ given by Kitaigorodsky et al. for tris(dibenzyldieneacetone) palladium(0) in THF. The dba ligand transition is an aromatic π → π* transition at 375 nm.

Figure 1. Calculated solubility limit based on Flory–Huggins theory. The solubility limit was calculated using eq 8 with values of Δ_solG_{THF}° = 12.2 kJ/mol, Δ_solG_W° = 42.5 kJ/mol, and χ_{W,THF} = 1.5, as described in the text.

Figure 2. Absorption spectrum of a 500 μM solution of Pd₂(dba)₃ in THF. The metal-to-ligand charge transfer band at 528 nm is labeled MLCT in the figure. The dba ligand transition is an aromatic π → π* transition at 375 nm.
lower than the value of 19,500 L mol\(^{-1}\) cm\(^{-1}\) measured previously by Gray and co-workers in pure 2-methyl THF,\(^{41}\) which may be due to the tendency of Pd\(_2\)(dba)\(_3\) to spontaneously form aggregates even in pure THF.\(^{24}\) All of the solutions had a measurable absorption at 26,670 cm\(^{-1}\) (375 nm) due to the \(\pi \rightarrow \pi^*\) transition of free dba. As observed in Figure 2, this band did not interfere with the solubility measurements based on the MLCT band.

The optimum method for measurement of solubility would involve the use of a variable path length cell. Because such cells are usually not available, the recommended procedure uses a combination of standard 1 mm and 1 cm path length cells to span an absorbance range of the saturated solutions from \(A \sim 0.001\) in 30% THF solutions (1 cm path length) to \(A > 20\) in a 1 cm path length equivalent. Owing its high absorbance, the 90% THF concentration was diluted by a factor 10, and then the absorbance was measured in a 1 mm cuvette (Figure 3A). Below 30% THF, the absorbance was too low to obtain a measurement in a 1 cm path length cell (\(A < 0.0001\)), which is the longest convenient path length in the HP8542 spectrophotometer (Figure 3A).

[FIGURE 3] (A) Absorbance data for mixtures made by addition of 4 mM Pd\(_2\)(dba)\(_3\) in dry THF solutions to H\(_2\)O. The 90% THF sample was multiplied by a factor of 10 because a diluted sample was used for the actual measurement to avoid measuring an absorbance \(A_{258\text{ nm}} > 1\). (B) Solubility of Pd\(_2\)(dba)\(_3\) is based on the extinction coefficient of 8,970 L mol\(^{-1}\) cm\(^{-1}\) for the 18,940 cm\(^{-1}\) band of Pd\(_2\)(dba)\(_3\) in THF. Concentrated solutions are purple in color and a 1 mm cuvette was required above 60% THF. The sample at a volume fraction of 90% THF was too concentrated to measure accurately in a 1 mm cuvette so it was diluted by a factor of 10 prior to measurement as indicated. Below 30% THF, suspensions of Pd\(_2\)(dba)\(_3\) are transparent and lack any color.

\[ \text{Concentration} \times \text{Path length} = \text{Absorbance} \times \text{Extinction coefficient} \]

\[ A = \frac{\text{Absorbance}}{\text{Concentration}} \times \text{Path length} \]

\[ \text{Extinction coefficient} = \frac{\text{Absorbance}}{\text{Concentration} \times \text{Path length}} \]

\[ \text{Solubility limit} = \frac{1 \text{ cm path length cell}}{0.001 \text{ absorbance}} \]

\[ \text{MLCT band} = \frac{1 \text{ mm path length cell}}{0.0001 \text{ absorbance}} \]

\[ \text{Path length} = \frac{\text{Absorbance}}{\text{Concentration} \times \text{Solubility limit}} \]

\[ \text{Concentration} = \frac{\text{Absorbance} \times \text{Path length}}{\text{Extinction coefficient}} \]

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\[ \text{Path length} = \frac{\text{Absorbance}}{\text{Concentration} \times \text{Solubility limit}} \]
Students should be cautioned that accurate measurement requires \( A_{528\ nm} < 1 \) for greatest accuracy. Students will need to examine their measurement to see if \( A_{528\ nm} \) exceeds 1 for a given solution. In that case, the students may need to make an appropriate dilution to obtain an accurate measurement. They may need to be reminded of the dilution factor equation \( C_i V_i = C_j V_j \), where \( C_i \) and \( V_i \) are the concentration and volume of sample \( i \), respectively. Since the transmitted intensity depends on \( 10^{-\frac{A}{2}} \), the quantity of light reaching the detector drops below 10%, when \( A > 1 \). Although it is somewhat arbitrary, there is clearly a point at which the signal-to-noise ratio suffers from low light intensity. Because the concentration depends exponentially on the solvent volume fraction according to Flory–Huggins theory, the concentrations, and therefore absorbances of the samples at 528 nm, span more than 3 orders of magnitude in this experiment. The most concentrated solutions have \( A_{528\ nm} > 2 \) even in a 1 mm cuvette. Thus, students will need to reason how to measure absorbance in such concentration solutions. The data in Figure 3 show that the solution was diluted by a factor of 10 at the highest THF volume fraction in order to make a measurement in the linear range.

Students can also compare their results to studies that have been carried out in the chemical literature. \( \text{Pd}_2(\text{dba})_3 \) has been used as a precursor in nanoparticle synthesis in a number of studies. Clearly, the mixed-solvent solubility is of great interest in understanding the mechanism for such syntheses. To increase the student’s awareness of the importance to read the chemical literature critically, students may wish to consult studies that use mixed solvents. One recent reference shows a picture of a solution of 400 \( \mu \text{M} \text{Pd}_2(\text{dba})_3 \) in 10%/THF/90%/H\(_2\)O.\(^{42}\) The data in Figure 3B show that these reported values\(^ {42} \) are >3000 times higher than the solubility limit. It would be instructive for students to compare this claim with their measurement and comment on this in the laboratory report.

**CONCLUSION**

The student laboratory experiment described here dispels a common misconception that solubility scales linearly with the solvent composition. In fact, the exponential dependence of chemical equilibrium, and therefore solubility in this case, on Gibbs energy is revealed by a colorimetric measurement of the concentration of a dye that is soluble in one solvent (THF), but insoluble in another (H\(_2\)O) in a two-component mixture. This fact can be demonstrated within about 2 h using inexpensive reagents and standard laboratory equipment; a weighing balance, UV–vis spectrophotometer (even single wavelength), a pipet, vials, and an optical cuvette.

**ASSOCIATED CONTENT**

**Supporting Information**

Student laboratory procedures; instructor notes. This material is available via the Internet at http://pubs.acs.org.

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