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Highly Dispersed Pseudo-Homogeneous and Heterogeneous Catalysts Synthesized via Inverse Micelle Solutions for the Liquefaction of Coal

Anthony Martino, Jeffrey S. Kawola, Allen G. Sault, Mark Hampden-Smith,
and Stacey A. Yamanaka

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Highly Dispersed Pseudo-Homogeneous and Heterogeneous Catalysts Synthesized via Inverse Micelle Solutions for the Liquefaction of Coal

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Abstract

The mission of this project was to use inverse micelle solutions to synthesize nanometer sized metal particles and test the particles as catalysts in the liquefaction of coal and other related reactions. The initial focus of the project was the synthesis of iron based materials in pseudo-homogeneous form. The first three chapters discuss the synthesis, characterization, and catalyst testing in coal liquefaction and model coal liquefaction reactions of iron based pseudo-homogeneous materials. Later, we became interested in highly dispersed catalysts for coprocessing of coal and plastic waste. Bifunctional catalysts to hydrogenate the coal and depolymerize the plastic waste are ideal. We began studying, based on our previously devised synthesis strategies, the synthesis of heterogeneous catalysts with a bifunctional nature. In chapter 4, we discuss the fundamental principles in heterogeneous catalysis synthesis with inverse micelle solutions. In chapter 5, we extend the synthesis of chapter 4 to practical systems and use the materials in catalyst testing. Finally in chapter 6, we return to iron and coal liquefaction now studied with the heterogeneous catalysts.

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Executive Summary

The mission of this project was to use inverse micelle solutions to synthesize nanometer sized metal particles and test the particles as catalysts in the liquefaction of coal and other related reactions. The initial focus of the project was the synthesis of iron based materials in pseudo-homogeneous form. The first three chapters discuss the synthesis, characterization, and catalyst testing in coal liquefaction and model coal liquefaction reactions of iron based pseudo-homogeneous materials. Later, we became interested in highly dispersed catalysts for coprocessing of coal and plastic waste. Bifunctional catalysts to hydrogenate the coal and depolymerize the plastic waste are ideal. We began studying, based on our previously devised synthesis strategies, the synthesis of heterogeneous catalysts with a bifunctional nature. In chapter 4, we discuss the fundamental principles in heterogeneous catalysis synthesis with inverse micelle solutions. In chapter 5, we extend the synthesis of chapter 4 to practical systems and use the materials in catalyst testing. Finally in chapter 6, we return to iron and coal liquefaction now studied with the heterogeneous catalysts.

In chapter 1, we discuss the synthesis of nanometer sized particles of Fe, Pd, and FeS₂ (pyrite) in inverse micelle solutions and testing of the particles as catalysts in coal conversion processes. TEM shows that the smallest particles are less than 3 nm in diameter. Electron diffraction confirms a presence of α -Fe (not crystalline oxides or borides), FCC-Pd, and the cubic pyrite structure of FeS₂. Several chemical manipulation techniques have been devised to isolate the particles as powders. Methanol washing and forced precipitation causes uncontrolled growth of the particles and results in materials of relatively high surface area. The ultra-small clusters can be isolated in freeze dried surfactant powders, and these powders can be re-dispersed in a variety of organic solvents. Catalytic activity of the ultra-small particles is determined in model compound hydrogenolysis, coal hydrolysis, and coal liquefaction. When surfactant is present, catalytic activity is hindered by two mechanisms. Surfactant chemically poisons the catalyst, and by-products of surfactant disintegration at reaction conditions scavenge hydrogen from hydrogen donating solvents. This latter mechanism is important in model compound hydrogenolysis and liquefaction as these processes depend strongly on hydrogen transfer. Moderate catalytic activity is observed in hydrogenolysis and liquefaction when the catalyst is used in a form free of

surfactant. In coal hydropyrolysis, a process that does not depend on hydrogen donating solvents, significant catalytic activity is observed despite the presence of surfactant. We attribute the high activity to high surface areas of the ultra-small particles and increased contact between the coal and metal due to the wetting properties of the surfactant solutions.

Chapter 2 is a continuation of the work begun in chapter 1, but with more focus on the characterization of the Fe particles. We have synthesized nanometer sized, highly dispersed Fe based particles in inverse micelle systems. TEM shows that the average particle size is less than 3 nm in diameter. Mossbauer spectroscopy confirms that the products of the reduction reaction of iron salts with LiBH_4 are 83% FeB, 15% $\text{Fe}^{2+}\text{BO}_x$, and 2% Fe^0 . Methanol washing and forced precipitation causes uncontrolled growth of the particles and results in relatively high surface area powders. FeB is converted to α -Fe then to an undetermined Fe^{2+} phase and then finally to Fe_3O_4 as the degree of oxidation by exposure to air increases. Exposure to methanol converts approximately 25% of FeB to Fe^0 . A shell and core picture of the clusters develops with oxidation as detected by XPS. Iron based metals are of interest as active, selective catalysts for a number of hydrogenation reactions including methanation and Fischer-Tropsch synthesis. We report again catalytic results of iron clusters in the hydrogenolysis of naphthyl bibenzyl methane (NBM), a model reaction for coal liquefaction.

In chapter 3, we study another methodology to synthesize highly dispersed metal particles on Fe based supports. We have prepared 2-4 nm sized Rh particles dispersed on a variety of supports by the photochemical reduction of $[(\text{C}_2\text{H}_4)_2\text{Rh}(\text{OEt})]_2 \cdot \text{H}_2\text{O}$ in the presence of the supports. Rh particles were dispersed on support materials of different morphology and composition (rod-, spindle-, cube-shaped α - Fe_2O_3 , β - FeOOH , and SiO_2) indicating the procedure is general. The mild photochemical reduction does not effect the microstructure of the support. Quantitative deposition of the Rh on the support resulted as no unreacted $[(\text{C}_2\text{H}_4)_2\text{Rh}(\text{OEt})]_2 \cdot \text{H}_2\text{O}$ or Rh particles exist in solution after reaction. Catalytic hydrogenation testing of the materials indicate that activity is a function of the composition of the support but not the morphology of the support at the length scales studied. Activity is also a function of particle size as effected by an increase in the metal loading of the material. Thermolysis of the β - FeOOH supported material shows that the dispersion of the metal and

the morphology of the support remain constant, the crystalline β -FeOOH becomes amorphous, and a channel structure in the amorphous material develops. As a result, catalytic activity of the heat treated Rh on β -FeOOH decreases.

At this time in the project, we became interested in highly dispersed catalysts for coprocessing of coal and plastic waste. Bifunctional catalysts to hydrogenate the coal and depolymerize the plastic waste are ideal. We began studying, based on our previously devised synthesis strategies, the synthesis of heterogeneous catalysts with a bifunctional nature. In chapter 4, the discussion is of nanometer sized gold particles encapsulated in the micropores of xerogels and aerogels. Gold was studied as a model system. The synthesis, characterization, and testing of practical materials are described in chapter 5. We have encapsulated nanometer sized Au particles in the micropores of xerogels and aerogels. The synthesis is a sequential reduction of a gold salt and sol-gel processing in an inverse micelle solution. The inverse micelle solution is used to solubilize the metal salt and provide a micro-reactor for the nucleation, growth, and stabilization of the nanometer sized clusters. Hydrolysis and condensation of an added siloxane precursor produces a wet gel embedding the particles. The presence of gel precursors de-stabilizes the inverse micelle structure resulting in larger particle sizes compared to typical inverse micelle synthesis techniques. Particle size control is complicated by the gel precursor effect on inverse micelle structure and the production of water and alcohol in the hydrolysis and condensation reactions. Finally, a unique gelation technique is outlined in these micro-heterogeneous solutions. Gelation occurs across the surfactant interface increasing the effective $H_2O:Si$ ratio. Gelation occurs even at low $H_2O:Si$ ratios, and condensation rates are high. Sol-gel parameters like the $H_2O:Si$ ratio and the surfactant concentration have unique and sometimes non-intuitive effects on the hydrolysis and condensation rates and the resulting material properties.

Having now developed the theoretical foundations for the synthesis of heterogeneous catalysis based on an inverse micelle synthesis, in chapter 5 we turn to a practical catalyst system. We introduce two novel synthesis strategies to make nanoclusters on silica and alumina supports using inverse micelle technology and sol-gel processing. In the first methodology, sol-gel chemistry is performed in non-alcohol containing cluster, inverse micelle solutions. Hydrolysis and condensation reactions are inefficient, but nonetheless

monoliths form around the clusters. In the second method, clusters formed in traditional inverse micelle solutions are allowed to diffuse into pre-existing monoliths formed using traditional sol-gel techniques (hydrolysis and condensation in ethanol). The two techniques give materials with similar cluster sizes, but different pore dimensions. At intermediate metal concentrations, particle sintering occurs even though metal size and pore dimensions are equal. At low metal concentrations, no appreciable sintering occurs even though the pores are much larger than the metal particles. We conclude that sintering is more a function of metal loading than the relative sizes of metal particles to pore diameters. Thus, even though sintering by particle to particle diffusion may be prevented by matching particle size with pore size, Ostwald ripening by atomic cleavage and migration from small to large particles dominates even in systems with equal particle and pore sizes. Our samples achieved relatively high catalytic activity and stability over time for propane dehydrogenation indicating that the synthesis methods described here result in catalyst performance more typical of PtSn/Al₂O₃ or PtSn/SiO₂ catalysts prepared by impregnation. This behavior may indicate a high resistance to coke formation in our catalysts.

Finally, in chapter 6 we return to the study of coal liquefaction. We continue our systematic characterization and catalyst activity testing of highly dispersed heterogeneous nanostructured materials by applying the above developed technology to the synthesis of iron based supported xerogel materials. Several iron based samples were synthesized and tested as catalysts in coal liquefaction. One key element of the testing involved the control of the surface acidity of the support. Surface acidity provides the cracking sites for either coal liquefaction or coal-waste coprocessing.

Chapter 1

Synthesis and Characterization of Coal Liquefaction Catalysts in Inverse Micelles

Abstract

We have synthesized nanometer sized particles of Fe, Pd, and FeS₂ (pyrite) in inverse micelle solutions and tested the particles as catalysts in coal conversion processes. The synthesis procedure produces a variety of high surface area, highly dispersed, unsupported metal and metal compound catalysts. While Pd is used to test the viability of the particles formed by this synthesis technique, only disposable iron based particles are of economic interest. The particles are prepared by reduction or chemical reaction of salts solubilized in micro-heterogeneous surfactant and oil solutions. Chemical manipulation produces metal powders. Particle size and composition are determined by transmission electron microscopy, electron diffraction, and UV-visible spectrophotometry. Catalysts in solution and as powders are explored in three reactions: the hydrogenolysis of naphthyl bibenzyl methane, coal hydrolysis, and coal liquefaction. The model hydrogenolysis reaction provides information on the role of surfactant on catalytic activity. Results indicate good catalytic activity for coal hydrolysis and liquefaction.

Introduction

Surfactant molecules possess two distinct moieties: a hydrophilic head group and a hydrophobic tail group. Because of the dual nature of surfactant molecules, they self-assemble in various solvents¹. In many two component systems of surfactant and oil, surfactants aggregate to form inverse micelles. Here, the hydrophilic head groups shield themselves by forming a polar core, and the hydrophobic tail groups are free to move about in the surrounding oleic phase. In three component microemulsion systems of water, oil, and surfactant, water is solubilized within the inverse micelle structure if the water concentration remains low. The solution takes on the structure of water droplets approximately 10-100Å in

diameter dispersed in the oleic solvent with surfactant forming the boundary between the two components.

In structured solutions such as inverse micellar and oil-rich microemulsion solutions, polar chemical reactants can be compartmentalized in the interior of the surfactant aggregates. Mixing two different solutions initiates a reaction sequence sustained by material exchange between inverse micelles. The surfactant interface provides a spatial constraint on the reaction volume, and reactions carried out in these micro-heterogeneous solutions produce colloidal sized particles (10-100Å) stabilized in solution against flocculation by surfactant^{2,3}. Metal alkoxide reactions⁴, the formation of polymer particles^{5,6}, and the formation of metal⁷⁻¹² and semiconductor¹³⁻²¹ clusters are examples of chemical processes that have been carried out in structured surfactant solutions.

The formation of ultra-small metal particles is of particular interest in the area of chemical catalysis. The clusters are high surface area, highly dispersed, unsupported materials. In addition, catalytic enhancement due to unique material properties (i.e. quantum size effects) is possible. Metal clusters prepared by a number of techniques have been studied as potential catalysts²². Reactant adsorption and the reactivity in various processes depends strongly on particle size. The effect is not just a matter of surface area, but of a fundamental change in the material's properties as the number of metal atoms per cluster decreases. Pt, Pd, Rh, and Ir particles prepared via inverse micelle techniques²³ have been studied in the hydrogenation and isomerization of but-1-ene^{24,25} and the hydrogenolysis and isomerization of hexanes²⁶. The catalysts were tested *in-situ* in surfactant, oil solutions and on supports. The activity of these catalysts increases when supported, and the selectivity of the colloidal sized particles is a function of the particle size.

We have studied the formation of iron based clusters and palladium clusters in inverse micelles and their use as catalysts in coal liquefaction reactions. Iron or palladium salts are solubilized within the polar interior of inverse micelles. The addition of an organic based reducing agent initiates a chemical reduction to produce Fe or Pd metal particles. The addition of a water based sulfiding agent initiates a chemical reaction to produce FeS₂ (pyrite) semiconductor particles. The clusters are characterized with respect to size and elemental structure with transmission electron microscopy, electron diffraction, and UV-visible spectrophotometry.

We have tested these iron based clusters for catalytic activity in a model reaction and in coal conversion reactions. The hydrogenolysis of naphthyl bibenzyl methane is used as a model for coal pyrolysis²⁷. We have studied this reaction with our iron based clusters and with a commercial catalyst for comparison. The Fe, Pd, and FeS₂ particles have also been tested in fixed bed coal hydrolypyrolysis and micro-batch coal liquefaction reactions. Once again, comparison is made with commercial catalysts.

Experimental

Octane, cyclohexane, and toluene were purchased from Aldrich at 99.9+% purity. Surfactants used include didodecyldimethylammonium bromide (DDAB) from Kodak, butyl-ethylene glycol n-dodecyl ether (C₁₂E₄) from Nikkol, and POE (6) nonylphenol (Ph₉E₆) from Chem Services Inc. Metal salts used include iron(II) chloride tetrahydrate, iron(III) chloride hexahydrate, iron(II) perchlorate hexahydrate, and palladium chloride from Aldrich, and iron(II) tetrafluoroborate hexahydrate from Alfa Chemicals. Lithium borohydride in tetrahydrofuran (2M) was purchased from Aldrich. Lithium sulfide was purchased from Aldrich and prepared in a 1M solution with de-ionized, distilled water.

The model hydrogenolysis compound, naphthyl bibenzyl methane (NBM), was synthesized in our laboratories using a previously developed recipe²⁸. Coal hydrolypyrolysis was carried out on bituminous Herrin Burning Star No. 2, and coal liquefaction was carried out on DECS-17 Blind Canyon Coal obtained from the Penn State Coal Sample Bank. DECS-17 is a HVA bituminous coal with 0.36% iron, 0.02% pyritic sulfur, and 7.34% mineral matter (on a dry basis). Hydrogen donating solvents 1,2,3,6,7,8 hexahydropyrene (H₆Py, 98% purity) and 9,10 dihydrophenanthrene (DHP, 94% purity) were both purchased from Aldrich. Histological grade, stabilized tetrahydrofuran (THF) and reagent grade heptane from Fisher Scientific were used in product workups.

Synthesis of Metal Colloids (Fe, Pd). Four different iron samples and two different palladium samples were prepared and studied (Table 1). The synthesis technique is general and specific samples differ only in the type and concentration of salts and surfactant added to the apolar solvents. First, the inverse micelle solutions are prepared by adding surfactant to the solvents (i.e. DDAB in toluene, C₁₂E₄ in octane). Then, the metal salts are introduced and the precursor salt solutions are mixed overnight on a stirring plate to assure complete

solubilization. Transparent yellow Fe salt solutions are formed, and transparent clear Pd salt solutions are formed. 2M LiBH_4/THF solution is directly injected into the salt solutions under rapid stirring to initiate the reduction of iron or palladium. The reaction is run at a 5:1 molar ratio of $[\text{BH}_4^-]:[\text{Fe}^{3+}]$ or $[\text{BH}_4^-]:[\text{Pd}^{2+}]$. Fe solutions immediately turn clear and transparent or black. Pd solutions turn dark purple. The reactions are carried out under dry, oxygen-free conditions.

Two iron samples were further worked up to isolate the iron particles from solution. In the first sample (**FeIII**), 15 vol. % methanol was added to the particle solution. After several hours a black precipitate was formed at the bottom of the flask below a clear and transparent solution. The precipitate was separated from the solution by centrifugation and washed with methanol. The process was repeated two or three times, and the powder was finally dried. In the second sample (**FeIV**), the volatile liquid was striped off to form a surfactant paste that contained the iron clusters. Freeze-drying of the paste formed a particle embedded surfactant powder. In this sample, less surfactant was used to increase the metal concentration in the final powder. The powder is easily re-dispersed in organic solvents.

Synthesis of FeS_2 : Four different FeS_2 samples were prepared and studied (Table 1). The synthesis is similar to the synthesis of metal colloids except 1M $\text{Li}_2\text{S}/\text{H}_2\text{O}$ is injected into the salt precursor solutions rather than a reducing agent. The surfactant solutions must be chosen so that some degree of water can be solubilized. In **FeS₂IV**, the precursor salt concentration was high enough that uncontrolled growth of the particles occurred upon reaction. The limited amount of surfactant was unable to stabilize particle growth, and the green product precipitated from solution. The particles were separated by centrifugation, washed, and dried to form a powder.

The samples used for UV-visible spectroscopy (**FeS₂II**) were prepared with different amounts of water. The synthesis is slightly different here. The precursor salt is added to H_2O first, and then the salt water is added to the surfactant, oil solution. The molarity of the salt in water and the amount of the salt water added to the inverse micelle solutions is adjusted to prepare samples of constant salt concentration and varying water amounts. The inverse micelle system ($\text{Ph}_9\text{E}_6/\text{cyclohexane}$) solubilizes a high degree of water. Two different synthesis routes were studied: (1) the reaction was run at a 5:1 molar ratio of $[\text{S}_2^{2-}]$

];[Fe²⁺], (2) the reaction was run at a 1:1 molar ratio of [S₂²⁻]:[Fe²⁺]. The solutions immediately turned dark green at low water concentrations and olive green at high water concentrations.

Characterization of the Colloids: The cluster solutions are studied by electron microscopy, electron diffraction, and UV-visible spectrophotometry. Particle size and structure are characterized with transmission electron microscopy (TEM) and electron diffraction. These tests are performed with a Joel 1200EX electron microscope. The colloidal solutions are applied directly on a holey carbon grid. The solution is wicked away by adsorbent towels under the grid and the particles are deposited on the grid. UV-visible spectrophotometry of the particle solutions is completed *in-situ* with a Hewlett-Packard 8452A diode array spectrophotometer. BET surface area analysis of the methanol washed iron powder (**FeIII**) was carried out with a Quantachrome Autosorb-6 surface analysis apparatus.

Naphthyl Bibenzyl Methane (NBM) Hydrogenolysis: Hydrogenolysis testing of NBM has been developed as a model reaction for coal liquefaction²⁷. We add 100 mg of NBM, 400 mg of the hydrogen donating solvent 9,10 dihydrophenanthrene (9, 10 DHP), and up to 15 wt. % catalyst on an NBM basis to a 10 ml flame sealed glass test tube. If used, 15 mg of elemental sulfur is added. The test tube is sealed under ambient air pressure. The tube is placed in a 400°C sand bath for one hour. Previously, NBM hydrogenolysis has been used to study the selectivity of catalysts for five different cleavage reactions²⁷. We do not quantify the selectivity of the catalysts used here. Conversion is calculated as the ratio of the molar sum of all products to the initial amount of NBM. The products are worked up in THF and analyzed by gas chromatography. Methanol washed and freeze dried Fe powders, and freeze dried FeS₂ powders were tested as catalysts. These results were compared to the commercial catalyst Shell 324 (12.4 wt. % Mo, 2.8 wt. % Ni on Al₂O₃).

Coal Hydropyrolysis: Fixed-bed hydropyrolysis tests of bituminous Herrin Burning Star No. 2 were performed at 520°C and 150 bar of hydrogen²⁹. Approximately 5 gm of coal are mixed with 10g sand in a reactor tube and heated at 5°C/sec. Char and ash are removed and weighed after each experiment. Conversion (100% - % char yield) is based on dry, mineral matter free coal. *In-situ* Fe and Pd colloids were tested as catalysts. The coal is wet with *in-situ* particle solutions, then dried, then the process is repeated until the desired

amount of material is loaded on the coal. For comparison, a commercially available sulfided Mo catalyst is dispersed by wetting the coal with a solution of $(\text{NH}_4)_2\text{MoO}_7$. This complex decomposes to MoS_2 at 400°C .

Coal Liquefaction: Batch liquefaction of Blind Canyon DECS-17 coal is performed in a stainless steel micro-reactor at a temperature of 400°C , a cold pressure of 800 psi H_2 , and a reaction time of 30 minutes³⁰. 2.5g of coal, 1g of the hydrogen donating solvent hexahydropyrene, and up to 2.5 wt. % of catalyst on a coal basis are added to the reactor. Reaction analysis consists of extracting products with THF and heptane (C7) in two separate stages, and the products are separated into categories depending on their solubility for the solvents. All soluble organic matter is extracted by THF (THF solubles), and the low molecular weight fraction of this material is extracted by C7 (C7 solubles). Insoluble organic matter (IOM) is neither soluble in THF or C7. The difference between THF solubles and C7 solubles represents the amount of organic solubles not soluble in C7 (represented by THF solubles/C7 insolubles). The percentage of product refers to that percentage of the original amount of dry, mineral matter free coal added before reaction. Three particle samples were tested: freeze dried Fe and FeS_2 samples as well as a surfactant free FeS_2 powder (**FeS₂IV**).

Results

Particle Characterization: Transmission electron micrographs of Fe and FeS_2 particles show ultra-small particles of uniform size and shape highly dispersed on the grid (Figure 1). The number average diameter by TEM of **FeI**, **PdI** (not shown), and **FeS₂I** particles is 1.5 ± 0.2 nm, 1.8 ± 0.2 nm, and 3.1 ± 0.1 nm, respectively. Fe particle size as determined by TEM shows no discernible trend and is roughly constant in the ranges of 0.001M - 0.01M FeCl_3 salt concentration and 1-10 wt. % DDAB in toluene. Average particle size in this system is 2.4 nm in diameter. The size of FeS_2 particles depends on the amount of water solubilized in the inverse micelle solutions and will be discussed later.

The methanol extracted Fe powder (**FeIII**) has been characterized by TEM and BET surface analysis. TEM pictures show agglomeration of the highly dispersed particles. Large agglomeration structures approximately 100 nm in size are observed. These structures

consist of the ultra small Fe particles (approximately 2 nm in diameter) flocculated together. Multi-point BET analysis gives a surface area of 156 m²/gm for this sample.

Selected area electron diffraction patterns of **FeII** particles are consistent with the presence of α -Fe. The electron diffraction results are not consistent with the presence of any crystalline iron oxides or borides. Electron diffraction patterns of **PdI** particles are consistent with the presence of FCC-Pd. Electron diffraction patterns from **FeS₂I** particles are consistent with the presence of cubic pyrite FeS₂. Cubic pyrite FeS₂ and cubic FeS have nearly equal theoretical scattering patterns, but the pattern we observe is distinguished from the pattern one gets from cubic FeS because of the presence of a scattering ring from the [220] plane in pyrite. This plane does not exist in cubic FeS. The electron diffraction pattern is not consistent with any other crystalline Fe and S compounds.

UV-visible spectra of the **FeS₂II** particles in solution show four distinct absorption maxima (Figure 2a). When [S₂²⁻]:[Fe²⁺] is held in excess (= 5:1), the absorption maxima shift to lower energies as the water content of the precursor inverse micelle microemulsion increases. When [S₂²⁻]:[Fe²⁺] = 1, the maxima in the absorption spectra do not red shift with increasing water content, but decrease in magnitude and move toward the spectrum observed in the precursor salt solutions (Figure 2b). The absorption spectra also decay if the samples are held in the presence of oxygen over the course of a day (Figure 3).

Catalyst Testing: Hydrogenolysis of naphthyl bibenzyl methane (NBM) was tested with several inverse micelle synthesized iron based catalysts and compared with the commercial catalyst Shell 324. We tested the methanol extracted Fe powder (**FeIII**) with and without sulfur additives, and we tested freeze dried Fe and FeS₂ particle embedded surfactant powders (**FeIV** and **FeS₂III**). In each experiment we monitored the percentage recovery of the reactant NBM and the percentage conversion to products²⁷ by gas chromatography. In each experiment we also monitored the percentage recovery of the hydrogen donating solvent 9, 10 DHP and the percentage converted to its dehydrogenated form phenanthrene. In an average of six runs, NBM was recovered at nearly 97+/-3% in thermal runs (no catalyst) and no product was obtained (Figures 4a and b). Some NBM was converted to products in thermal runs with added elemental sulfur. Little 9, 10 DHP is dehydrogenated to phenanthrene in the thermal runs. Methanol extracted Fe powder as catalyst decreased the

amount of NBM recovered to 87% and to 75% with added S. Product recovery correspondingly increased. In the Fe sample tests, hydrogens are given up to the hydrogenolysis products, and some 9, 10 DHP is dehydrogenated to phenanthrene. **FeIV** and **FeS₂III** showed less catalytic activity according to the recovery of NBM. Product conversion could not be followed, however, because of the thermal cracking of the surfactant (bars not present in Figure 5a, D and E). Surfactant pyrolysis by-products disrupt GC analysis. Large amounts of the 9, 10 DHP was dehydrogenated to phenanthrene.

Much higher product yields were obtained with Shell 324 (Figure 5). Higher conversions of 9,10 DHP to phenanthrene are consistent with these higher yields. In order to study the effect of surfactant on catalytic activity, we doped Shell 324 with increasing amounts of DDAB. As the Shell 324:DDAB mass ratio decreases, the trends in reactant recovery and product conversion move back to the thermal results indicating loss of catalytic activity. Initially the conversion of 9, 10 DHP to phenanthrene moves toward the thermal results as well. At the lower Shell 324 : DDAB ratios, however, this trend reverses itself and nearly all of the 9,10 DHP is converted to phenanthrene.

Coal hydrolysis testing was done as a function of catalyst concentration for **PdII** catalysts. The coal was incipient wet as described previously to the desired active metal concentration. The activity of the catalyst increases with Pd concentration before flattening out at approximately 0.1% (Figure 6). The conversion at this point is roughly equal to that achieved with the commercial MoS₂ catalyst. The weight percentage of the MoS₂ catalyst is approximately 1% (not shown on Figure 6). Results for **FeI** particles are shown at one concentration and represent a lowering in overall coal conversion by approximately 10% compared to Pd results.

Freeze dried Fe and FeS₂ particle embedded surfactant powders (**FeIV** and **FeS₂III**) and a surfactant free FeS₂ powder (**FeS₂IV**) were tested in micro-batch coal liquefaction reactions. Little to no improvement over the thermal runs is noticed in the freeze dried samples (Figure 7). The surfactant free FeS₂ results show substantial improvement in the conversion to C₇ solubles. Overall conversion to THF solubles decreases.

Discussion

Catalyst Synthesis and Characterization: Introduction of LiBH_4 to the inverse micelle solutions containing iron or palladium salts causes immediate reduction to Fe or Pd, respectively. The particles are formed by the following chemical reactions: $[\text{Fe}^{3+}] + 3[\text{H}^-] \rightarrow \text{Fe (particles)} + 3/2\text{H}_2$ and $[\text{Pd}^{2+}] + 2[\text{H}^-] \rightarrow \text{Pd (particles)} + \text{H}_2$. Introduction of Li_2S to the Fe salt solubilized inverse micelle solutions causes immediate sulfiding of the Fe. The particles are formed by the following reaction: $[\text{Fe}^{2+}] + [\text{S}_2^{2-}] \rightarrow \text{FeS}_2 \text{ (particles)}$. Characterization of the particles by electron diffraction give strong evidence that the chemistry described here produces α -Fe, FCC-Pd, and FeS_2 (pyrite). LiBH_4 or Li_2S enters the surfactant, oil solution and is solubilized within the inverse micelle structure. The reaction is initiated and is sustained through material exchange between inverse micelles. After the particle growth has equilibrated, the surfactant stabilizes the particles and prevents flocculation and precipitation.

The final particle size of nanometer sized clusters formed in inverse micelles depends on a complicated nucleation and growth process³¹. Inverse micelles effect this process primarily in two ways: (1) diffusion of the reacting nucleation sites and ions is governed by exchange rates between micelles and (2) the critical nucleation size depends inherently on the size of the inverse micelles. Size dependence of Au, CdS, and CdSe clusters produced in inverse micelles has been discussed previously^{32,13-21}. The Fe particles synthesized in DDAB/toluene mixtures showed no size dependence by TEM on the salt to surfactant concentration ratio within the range studied.

Size dependence of FeS_2 particles on water concentration in the precursor inverse micelle solutions is inferred from UV-visible spectrophotometry. Particle growth is observed by UV-visible spectrophotometry, because semiconductor band gap excitations in the UV-visible region decrease with increasing particle size when quantum size effects are important³³⁻³⁵. We have observed that the absorption maxima red shift in FeS_2 with increasing water concentration (Figure 2a). We attribute the red shift in the UV-visible spectra with increasing water concentration to the formation of larger particles primarily due to the increase in inverse micelle size. This conclusion assumes the absorption maxima in FeS_2 (pyrite), a semiconductor, is due to a complicated photogenerated band gap excitation in the material. If so, we have illustrated that the FeS_2 particles are small enough to exhibit

quantum confinement effects, and that we can increase FeS₂ particle size by increasing the water concentration in inverse micelle microemulsions.

The chemical sensitivity of FeS₂ particles in solution is illustrated by the dependence of the absorption maxima on reactant concentration and exposure to air. The absorption maxima in FeS₂ do not shift with increasing water concentration when the [S₂²⁻]:[Fe²⁺] molar ratio is not in excess (Figure 2b). The maxima decrease, and at the highest water concentration the spectra resembles the spectra of the iron salt solution. We conclude that the reaction governing the production of FeS₂ colloids is reversible, and that water in the microemulsion mixtures favor the decomposition of the colloids into the ions. It is necessary to use excess reactants in order to push the chemical reaction to the right and favor the formation of the particles. The high surface reactivity of CdSe particles has been demonstrated previously³¹. The stability of the FeS₂ particles further demonstrates surface reactivity. Oxidation of FeS₂ is observed by the disappearance of the absorption maxima when the sample is left in the presence of air. Oxidation of Fe clusters is visible to the naked eye as the solutions turn yellow if exposed to air.

Catalyst Testing: Hydrogenolysis of NBM indicates catalytic activity with **FeIII**. A more than linear increase in activity with added S indicates that there is a synergistic effect between the Fe and S.

Catalyst activity of Shell 324 far exceeds the activity of the iron based clusters. Doping Shell 324 with surfactant illustrates the effect of surfactant on catalyst activity. When Shell 324 is doped with surfactant, catalyst activity decreases (Figure 5). The behavior of the hydrogen donating solvent indicates two mechanisms are at work. At low surfactant concentrations, a decrease in product conversion and a decrease in the amount of 9, 10 DHP dehydrogenated to phenanthrene indicates less activity due to a chemical or steric poisoning of the catalyst. At high surfactant concentrations, we propose that a decrease in NBM conversion and an increase in the amount of 9, 10 DHP dehydrogenated to phenanthrene indicates that the thermal pyrolysis products of the surfactant (observed by GC) scavenge the hydrogens from the donor solvent. Decreased activity with high dehydrogenation conversion of 9,10 DHP to phenanthrene of **FeIV** and **FeS₂III** is consistent with the presence of surfactant and the above stated mechanism.

Coal hydrolysis tests indicate that catalytic activity of ultra-small Pd particles rivals the activity of the MoS₂ catalyst. The adverse effect of surfactant is less important in hydrolysis. Hydrolysis is independent of hydrogen transfer from a hydrogen donating solvent, thus eliminating a mechanism by which surfactant hinders the reaction. Any negative effect of surfactant may also be offset by increased contact of metal and coal due to wetting properties of the surfactant solution. FeI particles show relatively high conversions, but less than Pd and the commercial catalyst. The activity is significant considering the concentration of Fe used is more than an order of magnitude lower than the concentration of the MoS₂ catalyst.

Iron sulfide has been studied extensively as a coal liquefaction catalyst. It is believed that pyrrhotite (Fe_{1-x}S) is the active form of the iron catalyst³⁶, and that pyrite is reduced to pyrrhotite under liquefaction conditions³⁷. We have attempted to use our Fe and FeS₂ (pyrite) particles as coal liquefaction catalysts. FeIV and FeS₂III powders show no improvement over thermal reactions. Liquefaction is highly dependent on a hydrogen donating solvent and surfactant pyrolysis by-products may scavenge hydrogen. Surfactant free FeS₂IV powder shows a decrease in the overall conversion to organic products, but an actual increase in the amount of low molecular weight organic products. High IOM percentages may be due to high selectivity for cracking polar functionalities within DECS-17.

Conclusions

We have synthesized ultra-small, highly dispersed Fe, Pd, and FeS₂ (pyrite) particles in a variety of inverse micelle systems. TEM shows that the smallest particles are less than 3 nm in diameter. Electron diffraction confirms a presence of α -Fe (not crystalline oxides or borides), FCC-Pd, and the cubic pyrite structure of FeS₂. Several chemical manipulation techniques have been devised to isolate the particles as powders. Methanol washing and forced precipitation causes uncontrolled growth of the particles and results in materials of relatively high surface area. The ultra-small clusters can be isolated in freeze dried surfactant powders, and these powders can be re-dispersed in a variety of organic solvents. Catalytic activity of the ultra-small particles is determined in model compound hydrogenolysis, coal

hydropyrolysis, and coal liquefaction. When surfactant is present, catalytic activity is hindered by two mechanisms. Surfactant chemically poisons the catalyst, and by-products of surfactant disintegration at reaction conditions scavenge hydrogen from hydrogen donating solvents. This latter mechanism is important in model compound hydrogenolysis and liquefaction as these processes depend strongly on hydrogen transfer. Moderate catalytic activity is observed in hydrogenolysis and liquefaction when the catalyst is used in a form free of surfactant. In coal hydropyrolysis, a process that does not depend on hydrogen donating solvents, significant catalytic activity is observed in spite of the presence of surfactant. We attribute the high activity to high surface areas of the ultra-small particles and increased contact between the coal and metal due to the wetting properties of the surfactant solutions.

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Table 1. Listing of the samples prepared and used in this study.

Name	Description	Uses
FeI	0.01M FeCl ₃ , 10 wt. % DDAB/toluene	TEM, Hydropyrolysis
FeII	0.01 Fe(BF ₄) ₂ , 10 wt. % C ₁₂ E ₄ /octane	Electron Diffraction
FeIII	FeI , powder from methanol washing	TEM, Hydrogenolysis, BET
FeIV	0.01M FeCl ₃ , 5 wt. % DDAB/toluene, freeze dried powder	Hydrogenolysis, Liquefaction
PdI	0.005M PdCl ₂ 10 wt. % DDAB/toluene	TEM, Electron Diffraction
PdII	0.01M PdCl ₂ 10 wt. % DDAB/toluene	Hydropyrolysis
FeS₂I	0.001M Fe(ClO ₄) ₂ , 10 wt. % C ₁₂ E ₄ /octane	TEM, Electron Diffraction
FeS₂II	0.001M FeCl ₂ , 10 wt. % Ph ₉ E ₆ /cyclohexane, w/ H ₂ O	UV-Visible Spectrophotometry
FeS₂III	0.01M Fe(ClO ₄) ₂ , 10 wt. % C ₁₂ E ₄ /octane, freeze dried powder	Hydrogenolysis, Liquefaction
FeS₂IV	0.05M Fe(ClO ₄) ₂ , 10 wt. % C ₁₂ E ₄ /octane, uncontrolled precipitate powder	Liquefaction

Figure 1a.

Figure 1b.

Figure 1. (a) TEM of FeI particles. The number average diameter of the particles shown is 1.5 ± 0.2 nm. (b) TEM of FeS₂I. The number average diameter of the particles shown is 3.1 ± 0.2 nm.

a)

b)

Figure 2. UV-visible spectra of FeS_2II particles in solution. (a) $[\text{S}_2^{2-}]:[\text{Fe}^{2+}] = 5:1$. The electronic spectra of the semiconductor shifts to lower energies with increasing water concentration in the inverse micelle solutions. (b) $[\text{S}_2^{2-}]:[\text{Fe}^{2+}] = 1:1$. With increasing water concentration, the spectra contains elements of FeS_2 colloids and the precursor salt.

Figure 3. UV-visible spectra of FeS_2II particles in solution. The colloids are stable in the absence of air, but degrade when exposed to O_2 .

a)

b)

Figure 4. NBM hydrogenolysis results for Fe clusters. (a) The percentage of reactant recovered and percentage converted to product. (b) The percentage of the hydrogen donating solvent 9, 10 DHP recovered and the percentage dehydrogenated to phenanthrene. A = thermal average; B = thermal with elemental S average; C = FeIII (~ 15% active metal basis); D = FeIII with elemental S (~ 15% active metal basis); E = FeIV (~ 1.8% active metal basis); F = FeS_2III (~1% active metal basis).

a)

b)

Figure 5. NBM hydrogenolysis results for Shell 324 as a function of DDAB surfactant doping. (a) The percentage of reactant recovered and percentage converted to product. (b) The percentage of the hydrogen donating solvent 9, 10 DHP recovered and the percentage reduced to phenanthrene. A = thermal average; B = Shell 324; C = Shell 324:DDAB = 1:4; D = Shell 324:DDAB = 1:20; E = Shell 324:DDAB = 1:50. Active metal basis ~ 15% for all catalyst runs.

Figure 6. Herrin Burning Star No. 2 hydropyrolysis results for **PdII**, **FeI**, and a commercial MoS_2 catalyst.

Figure 7. DECS-17 Blind Canyon coal liquefaction results for Fe based clusters. A = thermal average; B = FeIV (~ 2% active metal basis); C = FeS_2III (~ 1.2% active metal basis); D = FeS_2IV (~ 2.4% active metal basis).

Chapter 2

The Synthesis and Characterization of Iron Colloid Catalysts in Inverse Micelle Solutions

Abstract

We have studied the synthesis and characterization of Fe based nanometer sized clusters formed in inverse micelle solutions. Inverse micelles provide a colloidal sized reaction template in an oleic solvent. Metal salts are solubilized within the interior of inverse micelles, and the addition of a reducing agent initiates nucleation and growth to produce the clusters. Surfactant acts to stabilize the particles. TEM shows that the average particle size is less than 3 nm in diameter. The reduction reaction of iron salts with LiBH_4 in inverse micelle solutions results in the formation of FeB, $\text{Fe}^{2+}\text{BO}_x$ (i.e. pyroborate), and $\alpha\text{-Fe}$ as determined by Mossbauer spectroscopy, electron diffraction, and x-ray photoelectron spectroscopy (XPS). Oxidation after exposure to air leads to the conversion of the FeB to $\alpha\text{-Fe}$ then to an undetermined Fe^{2+} phase and then finally to Fe_3O_4 . Iron based metals are of interest as active, selective catalysts for a number of hydrogenation reactions including methanation and Fischer-Tropsch synthesis. We report catalytic results of iron clusters in the hydrogenolysis of naphthyl bibenzyl methane (NBM), a model reaction for coal liquefaction. The role of surfactant in the reaction mechanism is determined.

Introduction

Colloidal sized metal and semiconductor particles (1 - 20 nm) are of current interest, because they mark a material transition range between molecular and bulk properties. With decreasing colloid size, bulk properties are lost as the continuum of electronic states breaks down (i.e. quantum size effects) and as the fraction of surface atoms becomes large. There are a number of ways to synthesize metal particles with diameters between 1-20 nm (nanoclusters) [1-3]. All synthesis routines include nucleation, growth, and stabilization of the particles and attempt to control particle size, size distribution, chemical composition, and structure. Chemical or photolytic reduction [4-6], thermal decomposition [7-9], and vapor

phase condensation [10-12] of metal salts and organometallic reagents initiates nucleation and growth. Controlling double layer forces with buffers and electrolytes [13,14] and the use of steric stabilizing agents such as polymers [15-24], surfactants [25-33], and bulky ligand appendages [34-41] stabilize the particles in solution.

A common nanocluster synthesis technique is the reduction of metal salts in inverse micelle solutions [25-33]. Inverse micelles are solution structures formed by the self-assembly of surfactants in apolar solvents (i.e. toluene, alkanes). Surfactants possess two distinct moieties, a hydrophilic head group and a hydrophobic tail group, and they self-assemble in apolar solvents so that the hydrophilic head groups shield themselves from the oleic surroundings [42]. The relatively polar head group regions solubilize and confine added metal salts and act as reaction cages when a reducing agent is introduced. The reaction sequence is sustained by material exchange between inverse micelles, and final particle size is dependent on stabilization by the surfactant. Ultra-small, monodispersed particles sterically stabilized in solution by surfactant are formed.

There are numerous advantages to synthesizing colloidal sized metal particles in inverse micelle solutions. The inverse micelle structure provides a reaction template in the appropriate size range, and size control is achieved by regulating the nucleation and growth process [2]. The size of particles formed by the reduction of metal salts in inverse micelle solutions depends on the inherent size of the inverse micelles and the material exchange rate between inverse micelles. Both micelle size and the material exchange rate depend on experimental parameters like the surfactant/solvent system, the addition of water which swells the inverse micelles, and the salt to surfactant ratio. The inverse micelle technique provides a single synthesis route to producing a number of different metals, metal oxides, and metal sulfides. Metal type is dependent only on the initial metal salt chosen, and Au, Ag, Pt, Pd, Rh, and Ni and Fe based metals are examples of different colloids synthesized in inverse micelles [43-47]. Bimetallic colloids are formed by starting with mixtures of metal salts. Metal sulfides are synthesized by using sulfiding agents rather than reducing agents [48-61], and metal oxides are produced by using reducing agents in a basic environment [62-64].

We have explored the synthesis of iron based nanoclusters in inverse micelles. Characterization of the size of the clusters is completed with transmission electron microscopy (TEM), and chemical composition of the clusters synthesized by an Fe salt, borohydrid reduction reaction is completed with Mossbauer spectroscopy, electron

diffraction, and x-ray photoelectron spectroscopy (XPS). The effect of oxidation on the clusters is determined. Our interest in iron colloids stem from their potential use as coal liquefaction catalysts [79-81]. Iron based metals are also of interest as active, selective catalysts for a number of hydrogenation reactions including methanation and Fischer-Tropsch synthesis [82,83]. Here, we test the iron clusters in the hydrogenolysis of naphthyl bibenzyl methane (NBM), a model reaction for coal liquefaction.

Metal clusters prepared by a number of techniques have been studied as potential catalysts [68-71]. Reactant adsorption and the reactivity in various processes depend strongly on particle size. Pt, Pd, Rh, and Ir particles prepared via inverse micelle techniques [72] have been studied in the hydrogenation and isomerization of but-1-ene [73,74], and the hydrogenolysis and isomerization of hexanes [75]. The activity of these catalysts increases when supported, and the selectivity of the colloidal sized particles is a function of the particle size. These studies found that activity is hindered by the presence of surfactant most likely due to steric effects on the metal surface. In other studies, the presence of surfactant on Ni₂B enhances activity by increasing the dispersion of the particles despite steric effects, but adsorbed species on Co₂B particles decrease activity again most likely due to steric effects [76]. In catalyst tests performed on colloids in normal micelles, the presence of surfactant enhances selectivity [77,78] as hydrogenation is preferred when the reactant mimics the surfactant and the double bond is accessible to the metal surface.

Experimental

Octane and toluene were purchased from Aldrich at 99.9+% purity. Surfactants used include didodecyldimethylammonium bromide (DDAB) from Kodak and butyl-ethylene glycol n-dodecyl ether (C₁₂E₄) from Nikkol. Metal salts used include iron(III) chloride hexahydrate from Aldrich and iron(II) tetrafluoroborate hexahydrate from Alfa Chemicals. Lithium borohydride in tetrahydrofuran (2M) was purchased from Aldrich.

A. Synthesis of Fe Based Colloids: Four different iron samples (FeI, FeII, FeIII, and FeIV) were prepared and studied (Table 1). The reactions and all chemical manipulations are carried out in a glove box under dry, oxygen-free conditions. The synthesis technique is general and specific samples differ only in the type and concentration of salts and surfactant added to the apolar solvents. First, the inverse micelle solutions are prepared by adding

surfactant to the solvents (i.e. DDAB in toluene, C₁₂E₄ in octane). Then, the metal salts are introduced and the precursor salt solutions are mixed overnight on a stirring plate to assure complete solubilization. Transparent yellow Fe salt solutions are formed. 2M LiBH₄/THF solution is directly injected into the salt solutions under rapid stirring to initiate the reduction of iron. The reaction is run at a 3:1 molar ratio of [BH₄⁻]:[Fe²⁺]. The reacted Fe solutions turn black upon reduction, and the colloids formed in solution are stabilized by surfactant indefinitely. As an example of the synthesis, FeI is made by first mixing 10 wt. % DDAB in toluene. 0.01M FeCl₃ is added to the solution and mixed overnight. The 2M LiBH₄ in THF is then directly injected to total 0.03M LiBH₄ in the Fe salt, inverse micelle solution.

Further manipulation of the DDAB/toluene solutions produces iron powders. When approximately 15 vol. % methanol is added to the colloidal solution, the surfactant is washed from the colloid surface, and the particles aggregate. A black precipitate forms below a clear and transparent solution after several hours. The precipitate is separated from the solution by centrifugation and washed with methanol. The process is repeated two or three times, and the powder is finally dried (FeIII). We have also coated the particles with poly(vinylpyrrolidone) (MW = 40,000) by adding the polymer and iron powder to methanol. The particles remain in solution for days without the presence of surfactant. Removing the methanol produces Fe particles embedded in PVP powder (FeIV). Replacing surfactant with polymer allows us to control the ratio of stabilizing agent to metal, remove the relatively volatile surfactant for x-ray photoelectron spectroscopy studies, and protect the particles from oxidation during air exposure.

B. Characterization of the Colloids: Particle size and composition in the colloidal solutions are characterized with transmission electron microscopy (TEM) and electron diffraction. These tests are performed with a Jeol 1200EX electron microscope. The colloidal solutions are applied directly on a holey carbon grid, and the solution is wicked away by adsorbent towels under the grid leaving the particles on the grid. Transfer of the particle coated grid into the microscope briefly exposes the sample to air.

Mossbauer spectroscopy is used to determine the chemical state of the iron *in-situ* (FeII) and in the PVP coated iron powders (FeIV). The Mossbauer Spectrometer (Austin Science) and its operation are described in detail elsewhere [84]. The spectra are obtained using a 60 mCi⁵⁷Co in palladium foil source and computer fitted to Lorentzian lines with a

least-squares optimization procedure. The spectra are corrected during the collection procedure to remove the curved background of instrumental origin, and resonant absorption areas are found from integration of the background curvature-corrected spectra. With the use of an absolute laser velocity calibrator it is possible to measure isomer shifts to within an absolute accuracy of ± 0.005 mm/s. The *in-situ* sample was tested in the frozen state ($T = -176^\circ\text{C}$), and the particle embedded polymer powder was tested under four conditions: at room temperature without air exposure, at room temperature after air exposure, at -183°C , and at -183°C with an applied magnetic field of 11 kGauss.

X-ray photoelectron spectroscopy (XPS) is used to determine the chemical state of the iron particles. XPS is performed with a VG Microtech Clam2 analyzer operating at a resolution of 1 eV and a 600 W Al K_{α} x-ray source. Assignment of the iron oxidation state relies on comparison with the Fe 2p peak positions and lineshapes of iron metal and the various iron oxides [85]. These standard lineshapes have been verified on the same instrument used to analyze the colloidal iron particles. Charge referencing was performed for all samples by adjusting the C 1s binding energy to 284.6 eV. To prepare samples of the surfactant stabilized iron particles suitable for XPS analysis, the *in-situ* solution (FeII) was evaporated to concentrate the particles and a drop of the resulting solution was placed on a Ta foil and allowed to evaporate completely overnight. For analysis of the washed iron powders (FeIII), the particles were re-dispersed in methanol and a drop of the dispersed solution was placed on a Ta foil and again allowed to dry. Finally, for the particles embedded in PVP (FeIV), the PVP matrix was dissolved in methanol and a drop of the resulting solution was placed on a Ta foil and the methanol evaporated to form a PVP film. All samples were prepared and mounted on a sample holder inside a dry, oxygen free glove box. As with microscopy, transfer of the sample to the XPS chamber requires brief exposure to air.

Complexometric experiments were used to determine the extent of reaction of the iron salts upon reduction. 1, 10 phenanthroline forms a complex with ionic Fe^{2+} species that has a distinctive maximum at 530 nm. The peak is followed before and after reaction with UV-visible spectrophotometry (Hewlett-Packard 8452A diode array spectrophotometer) by adding 1, 10 phenanthroline to the Fe salt precursor solutions and reacted Fe colloid solutions. The quantity and nature of the Fe^{2+} species present is determined. Two salt solutions were studied: FeCl_3 and $\text{Fe}(\text{BF}_4)_2$ in 10 wt. % C_{12}E_4 / octane.

Finally, BET surface area analysis of the methanol washed iron powder was carried out with a Quantachrome Autosorb-6 surface analysis apparatus.

C. Naphthyl Bibenzyl Methane (NBM) Hydrogenolysis: Hydrogenolysis testing of NBM has been developed as a model reaction for coal liquefaction [86]. We add 100 mg of NBM, 400 mg of the hydrogen donating solvent 9,10 dihydrophenanthrene (9, 10 DHP), and up to 15 wt. % catalyst on an NBM basis to a 10 ml flame sealed glass test tube. If used, 15 mg of elemental sulfur is added. The test tube is sealed under ambient air pressure. The tube is placed in a 400°C sand bath for one hour. Previously, NBM hydrogenolysis has been used to study the selectivity of catalysts for five different cleavage reactions. We do not quantify the selectivity of the catalysts used here. Conversion is calculated as the ratio of the molar sum of all products to the initial amount of NBM. The products are worked up in THF and analyzed by gas chromatography. Methanol washed Fe powders were tested as catalysts (FeIII). These results were compared to the commercial catalyst Shell 324 (12.4 wt. % Mo, 2.8 wt. % Ni on Al₂O₃).

Results

Transmission electron micrographs show that the product of the reaction is highly dispersed nanometer sized particles of uniform size and shape (Figure 1). The number average diameter by TEM is 1.5 +/- 0.2 nm (FeI). Fe particle size as determined by TEM shows no discernible trend and is roughly constant in the ranges of 0.001M - 0.01M FeCl₃ salt concentration and 1-10 wt. % DDAB in toluene. In this range, the average particle diameter is 2.4 nm. The methanol extracted Fe powder (FeIII) characterized by TEM and BET surface analysis consists of agglomerated particles. Large agglomeration structures larger than 100 nm in size are observed (Figure 2). These structures consist of the ultra small particles (approximately 2 nm in diameter) flocculated together. Multi-point BET analysis gives a surface area of 156 m²/gm for this sample. The degree of re-dispersion upon addition of PVP has not been determined. As the particles are stable in solution for only a couple of days with PVP, it is likely that the degree of dispersion is not as great as after the original reaction in the presence of surfactant.

Mossbauer spectroscopy completed on the *in-situ* sample (FeII) frozen at -176°C (Figure 3) indicates the presence of three different forms of iron. The spectrum is fit with

two doublets and a singlet (corresponding Mossbauer parameters are found in Table 2). We assign the first doublet to the presence of fine superparamagnetic particles of FeB, the second doublet to the presence of $\text{Fe}^{2+}\text{BO}_x$, and the singlet to the presence of superparamagnetic Fe metal. On the basis of spectral areas, the iron phases are present at 83, 15, and 2 mole %, respectively with the species order listed above. The mole percentages are based on equal recoil-free fractions of each phase, and this assumption is only approximately correct in the case of small clusters.

Mossbauer spectroscopy completed on the particle embedded polymer powder (FeIV) indicates the presence of the same three forms of iron. The spectrum for the powder at 21°C before exposure to air (Figure 4) is fit with two doublets and a singlet (corresponding Mossbauer parameters are found in Table 3). The mole % of FeB, $\text{Fe}^{2+}\text{BO}_x$, and Fe metal are 60, 10, and 30 mole %, respectively. There is no discernible change in the spectrum or fit after the sample is exposed to air. The spectrum of the sample at -183°C is quite different from the room temperature spectrum. It appears that the two large peaks in the room temperature spectrum have broadened. The broadened spectrum is fit with a sextet and two doublets. The sextet has a magnetic hyperfine splitting of 128 kOe and an isomer shift of 0.35 mm/sec consistent with the presence of superparamagnetic FeB. When a magnetic field is applied to the sample, the spectrum is further broadened.

Selected area electron diffraction patterns of *in-situ* iron particles (FeII) deposited on the microscope grid are consistent with the presence of α -Fe. Diffraction parameters, d , from scattering patterns depend on instrument settings and the radius of each scattering maxima ($d = L\lambda/R$, where L is the instrument sample to detector distance, λ is the electron beam wavelength, and R is the diffraction ring radius). The ratios of theoretical diffraction parameters to the first diffraction parameter (d/d_1) are known for many crystalline materials. The diffraction pattern of FeII shows several maxima, and experimentally we observe d/d_1 equal to 1, 0.701, 0.573, 0.482, 0.436, and 0.381. Theoretical values of d/d_1 for α -Fe equal 1, 0.707, 0.577, 0.500, 0.447, and 0.408. Theoretical values of d/d_1 for FeB equal 1, 0.842, 0.728, 0.696, 0.670, and 0.615. The electron diffraction results, therefore, indicate the presence of α -Fe and do not indicate the presence of any crystalline iron oxides, borides, or

borates. A separate diffraction pattern consistent with the presence of crystalline B_2O_3 as a reaction by-product is also detected.

XPS detects Fe^{2+} for both the iron particles embedded in PVP (FeIV) and the *in-situ* iron particles (FeII). For FeIV, an Fe 2p_{3/2} binding energy of ~709.0 eV and a prominent shakeup feature at ~715 eV is observed (Figure 5a). Zero valent Fe either as a metal or boride is not detected. The signal to noise ratios for these spectra are poor due to the presence of either surfactant or PVP near the sample surface. The low signal to noise ratios prevent the detection of boron in these samples and prevent the possible detection of small amounts of zero valent Fe, which would appear at 707.0 eV. Signal to noise ratios are improved slightly by mild heating in vacuum, but not enough to allow detection of boron or positive identification of a metallic iron component. For the uncoated iron particles (FeIII), the signal to noise ratio is an order of magnitude higher, and the initial state of the iron is clearly a mixture of Fe^{2+} and Fe^{3+} (Figure 5b). The resulting spectrum is quite similar to that of magnetite (Fe_3O_4), and no more than 2-3 % of the detected iron is present as metallic iron or iron boride. An oxidized form of boron is also detected with a B 1s binding energy of 192.0 eV consistent with the presence of either B_2O_3 or an iron borate. Heating the washed iron particles to 300°C under vacuum results in elimination of the Fe^{3+} contribution and the emergence of a pure Fe^{2+} spectrum.

Complexometric experiments indicate that the reaction is complete upon the addition of $LiBH_4$. In the $Fe(BF_4)_2$ precursor solutions, the Fe^{2+} species complexes with 1,10 phenanthroline, and the distinctive peak is observed at 530 nm. The peak is non existent in $Fe(BF_4)_2$ samples after the addition of $LiBH_4$ indicating the reaction goes to completion. In $FeCl_3$ precursor solutions, no complex is formed with the Fe^{3+} species. After reaction, a slight peak at 530 nm is detected indicating some conversion of Fe^{3+} to Fe^{2+} but too little to quantify.

Hydrogenolysis of naphthyl bibenzyl methane (NBM) was tested with several inverse micelle synthesized iron based catalysts and compared with the commercial catalyst Shell 324. We tested the methanol extracted Fe powder (FeIII) with and without sulfur additives and the dispersed catalysts with surfactant (FeI). In each experiment we monitored the percentage recovery of the reactant NBM and the percentage conversion to products by gas

chromatography. In each experiment we also monitored the percentage recovery of the hydrogen donating solvent 9, 10 DHP and the percentage converted to its dehydrogenated form phenanthrene. In an average of six runs, NBM was recovered at nearly 97 $\pm$ 3% in thermal runs (no catalyst) and no product was obtained (Figure 6a). Some NBM was converted to products in thermal runs with added elemental sulfur. Little 9, 10 DHP was dehydrogenated to phenanthrene in the thermal runs. FeIII as catalyst decreases the amount of NBM recovered to 87% and with added S to 75%, and product recovery correspondingly increases. Hydrogens are given up to the hydrogenolysis products, and some 9, 10 DHP is dehydrogenated to phenanthrene (Figure 6b). The dispersed catalysts with surfactant present (FeI) showed little activity compared to thermal runs.

Much higher product yields were obtained with Shell 324 (Figure 7). Higher conversions of 9,10 DHP to phenanthrene are consistent with these higher yields. In order to study the effect of surfactant on catalytic activity, we doped Shell 324 with increasing amounts of DDAB. As the Shell 324:DDAB mass ratio decreases, the trends in reactant recovery and product conversion move back to the thermal results indicating loss of catalytic activity. Initially the conversion of 9, 10 DHP to phenanthrene moves toward the thermal results as well. At the lower Shell 324:DDAB ratios, however, this trend reverses itself and nearly all of the 9,10 DHP is converted to phenanthrene.

Discussion

Introduction of LiBH_4 to iron salt containing inverse micelle solutions causes immediate reduction of the salt to form nanometer sized, highly dispersed particles of equal size and shape. The reaction goes to completion. No initial Fe salt remains as determined by the complexometric experiments. LiBH_4 enters the surfactant-oil solution and is solubilized within the inverse micelle structure. The reaction is initiated and is sustained through material exchange between inverse micelles. After the particle growth has equilibrated, the surfactant stabilizes the particles and prevents flocculation and precipitation. The final particle size of nanometer sized clusters formed in inverse micelles depends on a complicated nucleation and growth process [2]. Inverse micelles effect this process primarily in two ways: (1) diffusion of the reacting nucleation sites and ions is governed by exchange rates between micelles and (2) the critical nucleation size depends inherently on the size of the inverse micelles. Size dependence of metal borides, CdS, and CdSe clusters produced in

inverse micelles has been discussed previously [48-61]. The Fe particles synthesized in DDAB/toluene mixtures showed no size dependence by TEM on the salt to surfactant concentration ratio within the range studied.

Mossbauer spectroscopy of *in-situ* Fe colloids (FeII) indicates three products are formed by the reduction of iron salts with LiBH₄ in inverse micelle solutions. FeB, Fe²⁺BO_x, and metallic Fe are synthesized to 83 mole %, 15 mole %, and 2 mole %, respectively. Similar type spectra were observed for the NaBH₄ reduction of Fe salts in diglyme [82], and the doublet was assigned to the presence of FeB. The isomer shifts we report are higher, but they are consistent with those reported by Shinjo et al. [88]. Metallic Fe produces a singlet with an isomer shift close to zero [89,90]. Clearly, α -Fe is not the majority phase as determined by Mossbauer. The following reaction accounts for the presence of the three products: $3\text{Fe}^{2+} + 4\text{BH}_4^- + x\text{H}_2\text{O} \rightarrow \text{FeB} + \text{Fe(s)} + \text{Fe}^{2+}\text{BO}_x + (5 + x)\text{H}_2 + \text{B}_2\text{H}_6$. The waters of hydration of the metal salt provide the source of water, and pyroborate, Fe₂B₂O₃, is one possible Fe²⁺ borate phase represented by Fe²⁺BO_x. Similar reaction products and a similar reaction mechanism are found for the borohydride reduction of iron salts in a nonaqueous solvent [91, 92]. Mossbauer spectroscopy on the particle embedded polymer powder (FeIV) detects the three phases, FeB, Fe²⁺BO_x, and metallic Fe, in approximate mole percentages of 60%, 10%, and 30%, respectively. The conversion of approximately 25 % of FeB to Fe(s) results from the methanol workup necessary to extract the particles from solution and prepare the powder. There is no discernible change in the spectra or fit after the powder is exposed to air. The iron is apparently protected from oxidation by the polymer.

Mossbauer results of the powder at low temperatures and in the presence of a magnetic field further support the presence of FeB. The spectral broadening is consistent with longer magnetic relaxation times in a superparamagnetic species, and the sextet has a hyperfine field consistent with FeB. In the limit of a sufficiently large magnetic field or at sufficiently low temperatures, the spectrum would resolve into six lines.

Electron diffraction of the *in-situ* Fe colloids (FeII) gives strong evidence for the presence of α -Fe. FeB is not detected. It has been reported that the presence of air during the reduction process favors the formation of metallic metal over metal borides by the

following mechanism: $MB + O_2 \rightarrow M(s) + B_2O_3$ [91,92]. In our procedure, the reaction is done in an argon environment, but the TEM grid is exposed to air prior to microscopy. The above stated mechanism is most likely responsible for the detection of α -Fe as the majority phase by electron diffraction while FeB is the majority phase detected by Mossbauer. The detection of crystalline B_2O_3 by diffraction supports this theory.

XPS of the *in-situ* solution (FeII) and the polymer coated particles (FeIV) detects an undetermined Fe^{2+} species. We postulate that the Fe^0 particles (FeB and α -Fe) formed by the reduction reaction and the oxidation of FeB undergo further surface oxidation to form a thin, poorly crystalline oxide layer over an Fe^0 core. The presence of metallic iron detected by Mossbauer is most likely obscured by the oxide layer and the poor signal to noise ratio. The surface oxidation does not continue on to the formation of Fe_3O_4 as in bulk iron samples [87]. The small size of the iron particles (in the case of FeII) or the small grain sizes (in the case of FeIV) and the presence of the surfactant or polymer may result in oxidation behavior that differs from bulk iron. XPS of the iron powder (FeIII) indicates the presence of Fe_3O_4 or a mixture of Fe^{2+} and Fe^{3+} . Since the washed sample is severely agglomerated and not protected by either surfactant or polymer, it is not surprising that the oxidation behavior upon exposure to air is similar to the oxidation behavior of bulk iron metal. Also, XPS detection of an oxidized boron species (not a boride, however) in the washed sample is consistent with the conversion of FeB into iron metal and B_2O_3 upon air exposure. Conversion of Fe^{3+} to Fe^{2+} upon heating FeIII to $300^\circ C$ under vacuum suggests that the Fe^{3+} is a superficial surface layer formed during brief air exposure during sample loading.

Hydrogenolysis of NBM indicates catalytic activity with FeIII. A more than linear increase in activity with added S indicates that there is a synergistic effect between the Fe and S. Catalyst activity of Shell 324 far exceeds the activity of the iron based clusters. Doping Shell 324 with surfactant illustrates the effect of surfactant on catalyst activity. When Shell 324 is doped with surfactant, catalyst activity decreases (Figure 7). The behavior of the hydrogen donating solvent indicates two mechanisms are at work. At low surfactant concentrations, a decrease in product conversion and a decrease in the amount of 9, 10 DHP dehydrogenated to phenanthrene indicates less activity due to a chemical or steric poisoning of the catalyst. At high surfactant concentrations, we propose that a decrease in NBM

conversion and an increase in the amount of 9, 10 DHP dehydrogenated to phenanthrene indicates that the thermal pyrolysis products of the surfactant (observed by GC) scavenge the hydrogens from the donor solvent. The dispersed catalyst (FeI) showed little activity most likely due to the presence of surfactant. The apparent advantage of the nanometer size of the particles is eliminated by the surfactant.

Conclusions

We have synthesized nanometer sized, highly dispersed Fe based particles in inverse micelle systems. TEM shows that the average particle size is less than 3 nm in diameter. Mossbauer spectroscopy confirms that the products of the reduction reaction of iron salts with LiBH_4 are 83% FeB, 15% $\text{Fe}^{2+}\text{BO}_x$, and 2% Fe^0 . Methanol washing and forced precipitation causes uncontrolled growth of the particles and results in relatively high surface area powders. FeB is converted to α -Fe then to an undetermined Fe^{2+} phase and then finally to Fe_3O_4 as the degree of oxidation by exposure to air increases. Exposure to methanol converts approximately 25% of FeB to Fe^0 . A shell and core picture of the clusters develops with oxidation as detected by XPS. Catalytic activity of the ultra-small particles is determined in a model compound hydrogenolysis reaction. When surfactant is present, catalytic activity is hindered by two mechanisms. Surfactant chemically poisons the catalyst, and by-products of surfactant disintegration at reaction conditions scavenge hydrogen from hydrogen donating solvents. This latter mechanism is important in coal liquefaction as this process depends strongly on hydrogen transfer.

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Table 1. Listing of the samples used in this study.

NAME	DESCRIPTION	CHARACTERIZATION
FeI	0.01M FeCl ₃ , 10 wt. % DDAB/toluene	TEM
FeII	0.01M Fe(BF ₄) ₂ , 10 wt. % C ₁₂ E ₄ /octane	Mossbauer, electron diffraction, XPS
FeIII	FeI powder from methanol washing	TEM, BET, XPS, NBM hydrogenolysis
FeIV	FeIII re-dispersed with PVP and re-dried	Mossbauer, XPS

Table 2. Mossbauer Parameters for FeII (Figure 3). T = -176°C.

Assignment	Isomer Shift ^a (mm/sec)	Quadrupole Splitting (mm/sec)	Magnetic Hyperfine Splitting (kOe)	% Area
Doublet 1 (FeB)	0.470	0.69	----	83.3
Doublet 2 (Fe ²⁺ BO _x)	1.300	3.08	----	14.7
Singlet (Fe ⁰)	-0.110	----	----	2.0

^arelative to iron

Table 3. Mossbauer Parameters for FeIV (Figure 4). T = 21°C.

Assignment	Isomer Shift ^a (mm/sec)	Quadrupole Splitting (mm/sec)	Magnetic Hyperfine Splitting (kOe)	% Area
Doublet 1 (FeB)	0.450	0.27	----	60.4
Doublet 2 (Fe ²⁺ BO _x)	1.108	2.11	----	10.4
Singlet (Fe ⁰)	-0.136	----	----	29.2

^arelative to iron

Figure 1. TEM of FeI particles. The number average diameter of the particles is 1.5 ± 0.2 nm.

Figure 2. TEM of FeIII particles. After methanol washing the clusters aggregate and form particulate powders.

Figure 3. Mossbauer spectrum of FeII . $T = -176^\circ\text{C}$. Fit analysis listed in Table 2.

Figure 4. Mossbauer spectrum of FeIV. $T = 21^{\circ}\text{C}$. Fit analysis listed in Table 3.

Figure 5a: XPS of the Fe 2p region of Fe particles embedded in PVP (FeIV). a) As prepared, b) heated to 250°C in vacuum. Note the single Fe 2p peak at 709.9 eV with a shakeup feature at ~715 eV characteristic of Fe²⁺. No appreciable intensity is observed at the position expected for zero valent iron.

Figure 5b: XPS of the Fe 2p region of uncoated Fe particles (FeII). a) As prepared, b) heated to 300°C in vacuum. The absence of shakeup features in the as prepared spectrum is characteristic of a mixture of Fe^{2+} and Fe^{3+} , while the shift of the 2p $3/2$ peak and the appearance of a shakeup feature after heating to 300°C indicate transformation of Fe^{3+} into Fe^{2+} . No zero valent iron is observed at 707.0 eV.

Figure 6a. NBM hydrogenolysis results for Fe clusters. The percentage of reactant recovered and percentage converted to product. A = thermal average; B = thermal with elemental S average; C = FeIII (~ 5% active metal basis); D = FeIII with elemental S (~ 5% active metal basis).

Figure 6b. NBM hydrogenolysis results for Fe clusters. The percentage of the hydrogen donating solvent 9, 10 DHP recovered and the percentage dehydrogenated to phenanthrene. A = thermal average; B = thermal with elemental S average; C = FeIII (~ 5% active metal basis); D = FeIII with elemental S (~ 5% active metal basis).

Figure 7a. NBM hydrogenolysis results for Shell 324 as a function of DDAB surfactant doping. The percentage of reactant recovered and percentage converted to product. A = thermal average; B = Shell 324; C = Shell 324:DDAB = 1:4; D = Shell 324:DDAB = 1:20; E = Shell 324:DDAB = 1:50. Active metal basis ~ 5% for all catalyst runs.

Figure 7b. NBM hydrogenolysis results for Shell 324 as a function of DDAB surfactant doping. The percentage of the hydrogen donating solvent 9, 10 DHP recovered and the percentage dehydrogenated to phenanthrene. A = thermal average; B = Shell 324; C = Shell 324:DDAB = 1:4; D = Shell 324:DDAB = 1:20; E = Shell 324:DDAB = 1:50. Active metal basis ~ 5% for all catalyst runs.

Chapter 3

Preparation, Characterization, and Catalyst Testing of Highly Dispersed Rhodium Particles On Different Morphology Iron Oxide Supports.

Abstract

The deposition of uniform 2 - 4 nm sized Rh particles on different shaped α -Fe₂O₃ particles (rod shaped, spindle shaped, and cube shaped) and on β -FeOOH is achieved by the photochemical reduction of $[(C_2H_4)_2Rh(OEt)]_2 \cdot H_2O$ in the presence of the iron oxide. Deposition of the metal and synthesis of the support materials are independent, because the mild photochemical reduction does not effect the microstructure of the support. The Rh-coated iron oxide particles are characterized by electron microscopy, x-ray diffraction, and thermal analysis. We have also studied the catalytic activity of the highly dispersed, high surface area Rh particles in a model hydrogenation reaction. Catalyst activity is dependent on the support composition (β -FeOOH versus α -Fe₂O₃) and independent on support morphology at the length scales studied (~200 nm). Finally, we have studied the effects of thermolysis on the dispersion of the rhodium particles, the microstructure of the support materials, and the consequent effects on the activity of the catalyst.

Introduction

Studies of nanometer sized metal particles are generating a great deal of interest due to their size-dependent properties.^{1,2} These size-dependent properties exhibit themselves in both physical and chemical phenomena. Changes in physical phenomena such as electrical conductivity, optical absorption, and magnetism with size result from quantum-confinement effects.³⁻⁵ Changes in chemical phenomena such as catalytic reactivity and selectivity result from the high proportion of surface

atoms having unsatisfied valences and low coordination numbers in ultra small particles.⁶

We are interested in the preparation of nanometer sized Rh particles for heterogeneous (supported) catalysis applications.⁷⁻¹² While there are a number of synthetic methods available to prepare heterogeneous catalysts^{7-9,13-21}, photochemical reduction methods²²⁻²⁴ are attractive because they avoid high temperatures that change the support microstructure.²⁵ Photochemical preparation of metal on metal oxide materials from inorganic or metal-organic precursors is broadly divided into two strategies. The first strategy requires a photoactive support material, e.g. TiO₂, to provide the electrons necessary to reduce the precursor complex. In this case, the metal is only deposited on the surface of the photoactive material and not on any insulating substrates. In the alternative strategy that we employ, the molecular precursor acts as the chromophore and metal may be deposited on the entire support surface regardless of composition.

In this work, we describe a method to deposit Rh on various supports by the photochemical reduction of a Rh(I) complex in the presence of the support. We study the synthesis, characterization, and activity of heterogeneous catalysts formed by the deposition of Rh on rod shaped, spindle shaped, and cube shaped α -Fe₂O₃. This work is an extension of our previous work¹² in which we reported on the deposition of 2 - 4 nm sized Rh particles on rod-shaped β -FeOOH and spherical SiO₂ particles. We further demonstrate the generality of the process and provide a comprehensive report on the catalytic activity of the materials. Because the deposition of the metal and the synthesis of the support are independent, we are able to study the deposition procedure and catalyst activity as a function of the support material. The catalysts are tested in the hydrogenation of pyrene, a model compound reaction representative of coal liquefaction processes. Finally, we investigate the thermal stability of the materials and the subsequent effect on catalytic activity.

Experimental

General Procedures and Instrumentation

Hydrocarbon and ethereal solvents (toluene and tetrahydrofuran) were purchased from Aldrich Chemical Co. and pre-dried over sodium, distilled from sodium benzophenone ketyl, and stored over 4 Å molecular sieves. Pyrene and anhydrous hexadecane was purchased from Aldrich Chemical Co. and used as delivered. Reagents $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ and NaH_2PO_4 were purchased from J.T. Baker Inc. The compound $[(\text{C}_2\text{H}_4)_2\text{RhOEt}]_2 \cdot \text{H}_2\text{O}$ was prepared as described previously.⁹

Scanning electron microscopy (SEM) was completed with a Hitachi S-800, and transmission electron microscopy (TEM) was performed on a JOEL 2000-FX operating at 200 KeV. The reaction products were dispersed in acetone. A silicon wafer was dip-coated with the particles for SEM, and a carbon-coated copper grid was dip-coated for TEM. Electron diffraction revealed the presence or absence of crystallinity. X-ray powder diffraction was completed on a PVD-V Scintag X-ray diffractometer using a smear mount method to form a thin layer of sample onto a 3 x 3 cm² piece of glass. Surface areas were measured the BET nitrogen adsorption method. Thermogravimetric analysis (TGA) was performed on a Perkin-Elmer TGA 7 instrument.

Rod-shaped, spindle-shaped, and cube-shaped particles of $\alpha\text{-Fe}_2\text{O}_3$ were prepared by modifications of literature procedures as described below.²⁶⁻²⁸ The deposition of Rh on these materials is also described below. Rod shaped $\beta\text{-FeOOH}$ particles and rod-shaped $\beta\text{-FeOOH}$ with 2.4 wt.% and 11.4 wt.% Rh were prepared as described previously.¹² All manipulations were carried out under a dinitrogen atmosphere using standard Schlenk techniques.²⁹

Syntheses

1. Rod, Spindle, and Cube-Shaped $\alpha\text{-Fe}_2\text{O}_3$

Rod-Shaped $\alpha\text{-Fe}_2\text{O}_3$: Rod-shaped $\alpha\text{-Fe}_2\text{O}_3$ is produced from the thermal decomposition of rod-shaped $\beta\text{-FeOOH}$. Rod-shaped $\beta\text{-FeOOH}$, 0.304 g (3.42 mmole), was heated at 400 °C *in vacuo* for 2 hour, and 0.244 g (1.53 mmole) of the red-brown product was obtained. The corresponding yield of the process is 89% based on Fe_2O_3 .

Spindle-Shaped α -Fe₂O₃: 2.703 g (10 mmole) of FeCl₃•6H₂O and 0.138 g (1.0 mmole) of NaH₂PO₄ were dissolved in about 100 cm³ of distilled water. The solution was filtered and diluted to 500 cm³, transferred into a 1000 cm³-Erlenmeyer flask, and sealed with a screw cap. The Erlenmeyer flask was then placed into a preheated oven and kept at 100°C for 48.5 hours. The color of the solution changed from yellow to dark-brown. After cooling, the solid was separated by centrifugation, washed six times with distilled water, three times with acetone, and dried at ambient temperature to give 0.674 g of final product. The corresponding yield of the process is 84% based on Fe₂O₃.

Cube-Shaped α -Fe₂O₃: 2.4336 g (9 mmole) of FeCl₃•6H₂O was dissolved in 100 cm³ of distilled water with 5 cm³ of 0.1 M HCl. The solution was filtered and diluted to 500 cm³, transferred to a 1000 cm³-Erlenmeyer flask, and sealed with a screw cap. The Erlenmeyer flask was then placed into a preheated oven and kept at 100°C for 24 hours. The color of the solution changed from yellow to brown. After cooling, the solid was separated by centrifugation, washed six times with distilled water, three times with acetone, and dried at ambient temperature to give 0.287 g of final product. The corresponding yield of the process is 40% based on Fe₂O₃.

2. Deposition of Rh Particles by Photochemical Reduction

Rh particles were dispersed on each of the three morphologies of α -Fe₂O₃ through photoreduction of [(C₂H₄)₂RhOEt]₂•H₂O in the presence of the α -Fe₂O₃ samples. Three Schlenk flasks were charged separately under N₂ with 7.5 mg (0.018 mmole) of [(C₂H₄)₂RhOEt]₂•H₂O and 150 mg (0.939 mmole) of rod-shaped α -Fe₂O₃, 10 mg (0.023 mmole) of [(C₂H₄)₂RhOEt]₂•H₂O and 200 mg (1.25 mmole) of spindle-shaped α -Fe₂O₃, and 8 mg (0.019 mmole) of [(C₂H₄)₂RhOEt]₂•H₂O and 154 mg (0.964 mmole) of cube-shaped α -Fe₂O₃. A stir bar and 15 cm³ of THF was added to each flask. The suspensions were dispersed by indirect sonication for one minute and stirred at room temperature in the dark for about 30 minutes. The mixtures were then irradiated with a mercury lamp for 2 hours. The products were separated by filtration, washed with acetone, and dried in air. The filtrates of Rh on

rod-shaped and spindle-shaped $\alpha\text{-Fe}_2\text{O}_3$ are colorless while the filtrate of Rh on cube-shaped $\alpha\text{-Fe}_2\text{O}_3$ shows a light yellow color. Assuming quantitative reaction of the Rh(I) complex, the metal loading on the supports is 2.4 wt. %.

Catalytic Pyrene Hydrogenation

Hydrogenation of pyrene was carried out in a Berghof Teflon lined microbatch reactor. The reactor was charged with 100 mg pyrene and 1000 mg of hexadecane solvent, and the reactions were run at $100 \pm 3^\circ\text{C}$, 100 psig cold H_2 , for 30 min.. Two catalyst loadings were tested: 0.5 wt. % and 2.3 wt.% catalyst on a Rh to pyrene basis. Hydrogenation products for this reaction include iso-decahydropyrene, decahydropyrene, tetrahydropyrene, iso-hexahydropyrene, hexahydropyrene, and dihydropyrene. Products were worked up in toluene and product analysis was completed by gas chromatography. Activity is reported in turnover numbers (TON) defined by: [(number of moles of pyrene reacted)/(number of moles of Rh $\cdot$ time)].

Results and Discussion

Characterization of 2-4 nm Rh Particles on Rod-Shaped, Spindle Shaped, and Cube-Shaped $\alpha\text{-Fe}_2\text{O}_3$

The synthesis described in the previous section produced rod-shaped, spindle-shaped, and cube-shaped $\alpha\text{-Fe}_2\text{O}_3$. X-ray powder diffraction confirmed the presence of crystalline $\alpha\text{-Fe}_2\text{O}_3$ in the rod-shaped material, and X-ray powder diffraction and electron diffraction confirmed the presence of crystalline $\alpha\text{-Fe}_2\text{O}_3$ in the spindle and cube-shaped material. The lattice fringe images observed by high resolution TEM verify that individual particles of spindle and cube-shaped $\alpha\text{-Fe}_2\text{O}_3$ are single crystals. The spindle-shaped particles are approximately 325 nm long and 125 nm in diameter, and the average cube-shaped particle size is 150 nm. BET surface areas of the rod, spindle, and cube shaped $\alpha\text{-Fe}_2\text{O}_3$ powders are $8.6 \text{ m}^2/\text{g}$, $11.3 \text{ m}^2/\text{g}$, and $7.8 \text{ m}^2/\text{g}$, respectively.

After deposition of Rh by photochemical reduction, the red-brown rod-shaped powders turned brown, and the brown spindle-shaped powders and red-brown cube-shaped powders darkened. In all cases, it is seen that 2 - 4 nm particles of Rh are uniformly dispersed on the support materials. As in our previous work, we have shown that the photochemical deposition procedure is independent of support material (Fe_2O_3 , FeOOH , and SiO_2). Quantitative deposition of the Rh on the support resulted as no unreacted $[(\text{C}_2\text{H}_4)_2\text{Rh}(\text{OEt})]_2 \cdot \text{H}_2\text{O}$ exists in solution after reaction as determined by ^1H NMR spectroscopy of the material contained in the supernatant. Energy dispersive spectroscopy confirms the presence of Rh in all samples. No Rh particles were detected in the dispersed solution (even after sonication) indicating the particles adhere strongly to the support surface.

Thermolysis of Rod-, Spindle-, and Cube-Shaped $\alpha\text{-Fe}_2\text{O}_3$, and of $\beta\text{-FeOOH}$ and Rh coated $\beta\text{-FeOOH}$

In order to determine the effect of heat treatment in these materials, model experiments were carried out wherein the Fe_2O_3 and the $\beta\text{-FeOOH}$ supports were heated and characterized by crystallography and microscopy. The spindle and cube-shaped $\alpha\text{-Fe}_2\text{O}_3$ particles exhibited a total weight loss of 3.5% and 1.0%, respectively, on heating in static air up to 800 °C. The weight losses are probably due to the dehydration of surface hydroxyl groups. Heating these samples up to 800°C in a 7% H_2 in N_2 atmosphere resulted in weight losses of 32.3% and 31.0% for spindle- and cube-shaped powders, respectively. The weight losses are consistent with the formation of Fe.

We have shown that rod-shaped $\alpha\text{-Fe}_2\text{O}_3$ is produced by the thermal decomposition of rod-shaped $\beta\text{-FeOOH}$ at 400°C *in vacuo*. It is interesting to note that the synthesis is strongly dependent on temperature and atmosphere. At 300°C, an amorphous iron oxide is produced that gives no diffraction pattern. The amorphous material has a BET surface area of 54.2 m^2/gm , more than twice that of the original rod-shaped $\beta\text{-FeOOH}$ powder. At 400°C, in air rather than *in vacuo*, an amorphous iron oxide is once again the product. The rod-shaped morphology remains, but no

diffraction pattern is detected. TEM indicates the creation of an approximately 10 nm channel structure that does not exist in the β -FeOOH. An increase in microporosity of the amorphous material could account for the increase in the BET surface area. At present we speculate that the amorphous structure is derived from the removal of water and interstitial ions (Cl^-) from the β -FeOOH structure³⁰, and that thermolysis under air rather than vacuum allows the gross morphology to be retained while the porous internal microstructure can develop. Further investigation is clearly warranted.

On heating Rh loaded β -FeOOH particles (synthesis described previously¹²) to 400°C in air, the transition to the same amorphous iron oxide material occurs. X-ray diffraction shows only the pattern consistent with fcc-Rh. No agglomeration of the Rh particles is detected.

Catalytic Hydrogenation of Pyrene

Polynuclear aromatic hydrocarbons (PAHs) are vital in the liquefaction of coal to viable fuels, because they assist hydrogen transfer between a hydrogen source and free radical coal fragments formed in the liquefaction process.³¹ Hydrogenation of the PAH pyrene models the kinetics and thermodynamics of the hydrogenation of PAHs in liquefaction reactions³² and so the reaction is studied. Hydrogenation of pyrene to form 4,5-dihdropyrene is modeled by pseudo first order reversible kinetics, and the reaction proceeds at moderate conditions ($T = 100^\circ\text{C}$, $P = 100$ psi H_2) in the presence of a catalyst. Catalysts studied in the past include Ni³³, NiMo³⁴, Pd³⁵, Fe³⁶, MoS₂³⁷, and Rh⁸. We have studied the heterogeneous Rh catalysts synthesized here as a function of the support material, thermal treatment, and metal loading.

Background runs on the three types of α -Fe₂O₃, β -FeOOH, and SiO₂ show no catalytic activity; no products are detected giving a TON of zero. The materials become catalytic in the presence of Rh, and activity is a function of the active metal concentration (Figure 1). Support composition dependent catalytic activity is clearly noted (Figure 2). The activity of the Rh on β -FeOOH catalyst is twice that of

previously studied Rh catalysts⁸ and four times greater than Pd catalysts³⁵ studied at the same conditions. Activity of the heterogeneous Rh catalysts decreases by factors of two as a function of support material for α -Fe₂O₃ and then SiO₂. The activity of Rh on rod-, spindle-, and cube-shaped α -Fe₂O₃ catalysts is relatively constant (Figure 2) indicating that activity is independent on the morphology of the support material.

The effect of the thermolysis of Rh on β -FeOOH is minimal (Figure 3). Catalytic activity decreases slightly in the temperature treated sample. In the temperature treated sample, the dispersion of the metal and the morphology of the support remain constant, the crystalline β -FeOOH becomes amorphous, and a channel structure in the amorphous material develops. Decrease in the catalytic activity must result from the chemical change in the support that produces the amorphous structure.

Finally, catalytic activity is determined as a function of the metal loading on the support (Figure 4). At constant active metal concentration with respect to the reactant pyrene, catalytic activity drops substantially in the material highly loaded with Rh. Lower activity is consistent with lower dispersion and larger particle sizes observed in the highly loaded Rh on β -FeOOH material.

Conclusions:

We have prepared 2-4 nm sized Rh particles dispersed on a variety of supports by the photochemical reduction of $[(C_2H_4)_2Rh(OEt)]_2 \cdot H_2O$ in the presence of the supports. Rh particles were dispersed on support materials of different morphology and composition (rod-, spindle-, cube-shaped α -Fe₂O₃, β -FeOOH, and SiO₂) indicating the procedure is general. The mild photochemical reduction does not effect the microstructure of the support. Quantitative deposition of the Rh on the support resulted as no unreacted $[(C_2H_4)_2Rh(OEt)]_2 \cdot H_2O$ or Rh particles exist in solution after reaction. Catalytic hydrogenation testing of the materials indicate that activity is a function of the composition of the support but not the morphology of the support at the length scales studied. Activity is also a function of particle size as effected by an increase in the metal loading of the material. Thermolysis of the β -FeOOH supported

material shows that the dispersion of the metal and the morphology of the support remain constant, the crystalline β -FeOOH becomes amorphous, and a channel structure in the amorphous material develops. As a result, catalytic activity of the heat treated Rh on β -FeOOH decreases.

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Figure 1: Effect of Rh concentration on the catalytic hydrogenation of pyrene for 2.4 wt. % Rh on cube-shaped $\alpha\text{-Fe}_2\text{O}_3$ materials. TON for cube-shaped $\alpha\text{-Fe}_2\text{O}_3$, all other iron oxide materials, and SiO_2 without Rh is zero (no products formed).

Figure 2: Effect of support composition and morphology on the catalytic hydrogenation of pyrene (2.4 wt. % Rh on each support, 0.5 wt. % Rh to pyrene). β -FeOOH supported materials are roughly two times more active than α -Fe₂O₃ supported materials and six times for active than SiO₂ supported materials.

Figure 3: Effect of heat treatment on synthesis of β -FeOOH supported catalysts (2.4 wt. % Rh on each support, 0.5 wt. % Rh to pyrene). Catalytic activity decreases slightly with the crystalline to amorphous change in the support material. The crystalline material is synthesized in vacuo at 400°C while the amorphous material is synthesized at 300°C.

Figure 4: Effect of metal loading on the β -FeOOH support (at constant metal concentration in the reaction) on catalytic hydrogenation of pyrene. The material with high metal loadings shows a marked decrease in activity due to loss of dispersion of the metal.

Chapter 4

Encapsulation of Gold Nanoclusters in Silica Materials via an Inverse Micelle / Sol-Gel Synthesis

Abstract

Nanometer sized gold particles were encapsulated in the micropores of xerogels and aerogels. The synthesis involves the sequential reduction of a gold salt followed by sol-gel processing in an inverse micelle solution. The inverse micelle solution solubilizes the metal salt and provides a micro-reactor for the nucleation, growth, and stabilization of the nanometer sized clusters. Hydrolysis and condensation of an added siloxane precursor produces a wet gel embedding the particles. Characterization of the particle size and composition and the particle growth process was completed with transmission electron microscopy (TEM), electron diffraction, and UV-visible absorption spectrometry. Characterization of the gel surface areas was completed with N₂ porosimetry. Material properties determined as a function of the gel precursor (TEOS vs. a pre-hydrolyzed form of TEOS), the water to gel precursor reaction stoichiometry, and surfactant concentration are discussed in terms of the unique solution chemistry occurring in the micro-heterogeneous inverse micelle solutions.

Introduction

Colloidal sized metal and semiconductor particles with diameters of 1 - 20 nm (nanoclusters) are of current interest because they mark a material transition range between molecular and bulk properties. With decreasing colloid size, bulk properties are lost as the continuum of electronic states breaks down (i.e. quantum size effects) and as the fraction of surface atoms becomes large. There are a number of ways to synthesize metal particles with diameters between 1-20 nm¹⁻³. All synthesis routes include nucleation, growth, and stabilization of the particles and attempt to control particle size, size distribution, chemical composition, and structure. Chemical or photolytic reduction⁴⁻⁶, thermal decomposition⁷⁻⁹, and vapor phase condensation¹⁰⁻¹² of metal salts and organometallic reagents initiates

nucleation and growth. Controlling double layer forces with buffers and electrolytes^{13,14} and the use of steric stabilizing agents such as polymers¹⁵⁻²⁴, surfactants²⁵⁻³⁴, and bulky ligand appendages³⁵⁻⁴² stabilize the particles in solution.

A common nanocluster synthesis technique is the reduction of metal salts in inverse micelle solutions²⁵⁻³⁴. Inverse micelles are solution structures formed by the self-assembly of surfactants in apolar solvents (i.e. toluene, alkanes). Surfactants possess two distinct moieties, a hydrophilic head group and a hydrophobic tail group, and they self-assemble in apolar solvents so that the hydrophilic head groups shield themselves from the oleic surroundings⁴³. The relatively polar head group regions solubilize and confine added metal salts and act as reaction cages when a reducing agent is introduced. Ultra-small, monodispersed particles sterically stabilized in solution by surfactant are formed. Advantages of the inverse micelle synthesis technique include mean particle size control and compositional variety. Particle size control results from regulation of the nucleation site size (inverse micelle size) and the particle growth rate (material exchange rate between inverse micelles). A variety of metals and bimetallics are produced by choosing different metals salts and mixtures of metal salts. Unfortunately, the presence of surfactant hinders potential applications (i.e. catalysis), and removal of the surfactant causes immediate flocculation of the particles with loss of the unique and interesting material properties.

Our goal is to isolate nanoclusters and preserve their unique properties without the presence of any stabilizing agents that attach themselves to the particle surface. We have developed sol-gel processing in inverse micelle, nanocluster solutions to encapsulate or sterically entrap nanoclusters in the micropores of xerogels and aerogels. The synthesis is a sequential reduction of a metal salt and sol-gel processing of an added siloxane precursor in an inverse micelle solution. Sol-gel processing of porous silica gels in polar solvents has been used to encapsulate a variety of large molecules including laser dyes, photochromics, and proteins⁴⁴⁻⁵⁰. Sol-gel encapsulation involves polymerization of a gel precursor, usually tetramethoxysilane or tetraethoxysilane, to build the silica gel structure (host) around the dopant (guest). Steric entrapment occurs when the gel pore size is comparable to the dopant size. Also, nanometer sized gold particles have been stabilized with a silica coating using silane coupling agents as surface primers⁵¹. The product of this work includes nanoclusters in a silica monolith that remain small, of one size, and highly dispersed even after the removal of the surfactant.

Other works involving sol-gel processing in surfactant, apolar solvent solutions use high surfactant or alcohol concentrations and are not suitable for monodisperse cluster formation⁵²⁻⁵⁸. We have established an entirely new system with low surfactant concentrations and no alcohol. The inverse micelle solutions and precursor salt solutions are characterized by small angle x-ray scattering (SAXS). After chemical reduction of the metal salt and hydrolysis and condensation of an added siloxane precursor, cluster size, chemical composition, and the cluster growth process are characterized by transmission electron microscopy (TEM), electron diffraction, and UV-visible absorption spectrometry. Gel surface areas are determined by N₂ porosimetry. Material properties are determined as a function of the gel precursor (TEOS vs. the pre-hydrolyzed form of TEOS), the water to gel precursor reaction stoichiometry (molar ratio = 4:1, 3:1, 2:1, and 1:1), surfactant concentration, drying mechanism, and process washing with hexanol.

We are currently interested in nanocluster-gel materials as catalysts. Metal-gel matrixed materials have been extensively studied as catalysts, because high surface area, high porosity gels act to disperse the active metal, minimize mass transfer and pore diffusion limitations, and promote activity through tunable surface acidities. Traditionally, atomic metals are added to gels when salts are ion-exchanged with terminal hydroxyl groups on the gel^{59,60}, through impregnation into the gel backbone during gel processing⁶¹⁻⁶⁴, or through the use of chelating agents^{65,66}. Clusters are formed later through chemical, thermal, and photolytic reduction. No current synthesis method prevents metal particle sintering under robust catalytic conditions, however. Our work is the first where formation of clusters proceeds formation of the gel, thus resulting in encapsulation and potentially new advantages. Clusters are trapped in the gel pores of comparable size limiting the modes of particle sintering. The advantages of the inverse micelle technique are retained including formation of ultra-small, monodisperse, highly dispersed particles, and particle size and composition control (i.e. various metals, mixtures of different metals, alloys, layered particles, metal oxides, and metal sulfides).

Experimental

Materials: The surfactant didodecyldimethylammonium bromide (DDAB), apolar solvent toluene (99.9+% purity), reducing agent lithium borohydride in tetrahydrofuran (2M), gold trichloride, tetraethylorthosilicate (TEOS), and a 40 wt. % tetrabutylammonium

hydroxide (TBAOH) in water solution were purchased from Aldrich and used as delivered. Polydiethoxy siloxane (M.W. = 610 gm/mol), an oligomer of TEOS formed by a proprietary pre-hydrolysis reaction, was purchased from United Chemical Technologies of Bristol, PA and used as delivered. Hexanol is used as a washing solvent and is purchased from Aldrich at 99.9+% purity.

Synthesis: Surfactant is added to toluene (1-5 wt. %) and stirred by hand shaking to form the inverse micelle solution. AuCl_3 (0.001M) and the gel precursor (0.4M) are added, and the solution is stirred until the salt is fully solubilized. A gold colored transparent solution is formed. The LiBH_4/THF solution is injected into the salt precursor solution under rapid stirring so that the $[\text{BH}_4^-] : [\text{Au}^{3+}] = 3:1$. The gold solution immediately turns dark purple. The 40 wt. % TBAOH in water solution is added 1 to 5 minutes afterwards, and gelation time is marked when the solution no longer flows under gravity. A deep purple, viscous gel is formed. The water to gel precursor molar ratio is studied as an experimental variable and is set at 1:1, 2:1, 3:1, and 4:1. Two different gel precursors are tested: TEOS and a pre-hydrolyzed form of TEOS. The gels are aged for fourteen days prior to drying under ambient conditions to form xerogels or under supercritical CO_2 extraction to form aerogels. Some samples were washed to remove surfactant and reaction by-products by passing warm hexanol over the dried, crushed gels through an aspirator funnel. Samples discussed in 'Results' were not washed unless indicated.

Characterization: SAXS was completed at the Small-Angle Scattering Center at the University of New Mexico. A 12 kW rotating anode source was used with a Kratky camera and an M-Braun linear position sensitive detector. A Ni filter was used as a monochromator, and the samples were sealed in quartz capillary tubes. Particle size and composition in the colloidal solutions were characterized with transmission electron microscopy (TEM) and electron diffraction. These tests were performed with a 300 keV Phillips CM30 electron microscope. Dried, crushed gels were dispersed over a holey carbon substrate. UV-visible absorption spectroscopy was completed with a Hewlett Packard 8452A Diode Array spectrophotometer. Quartz cuvettes were used in liquid samples, and gels were sliced and mounted on quartz microscope slides. BET surface area analysis of the crushed xerogels and aerogels was carried out with a Quantachrome Autosorb-6 surface analysis apparatus.

Results

Small angle x-ray scattering was used to characterize the inverse micelle solutions and the effect of the addition of the gold salt and gel precursors to the solution structure. SAXS of DDAB/toluene inverse micelle solutions, gold salt in DDAB/toluene inverse micelle solutions, and gold salt and TEOS in DDAB/toluene inverse micelle solutions is weak but existent. Scattering curves are flat and bend only at relatively large scattering angles. Guinier analysis indicates the aggregate size, $R_g = 0.49$ nm, remains constant in all three cases (Figure 1). No small angle x-ray scattering is observed when only TEOS is added to DDAB/toluene mixtures. Finally, scattering and Guinier analysis of gold salt and TEOS solutions after reduction (i.e. after cluster formation, but before hydrolysis and condensation) indicates a cluster size of $R_g = 2.34$ nm.

Formation of the clusters occurs within seconds after addition of the reducing agent. The gold solutions turn purple, the color indicative of the presence of colloidal sized gold. Gelation occurs in the cluster, surfactant solutions at 1.5, 2.5, 3.3, and 24 hours after the addition of TBAOH/H₂O for samples with the reaction stoichiometry of 1:1, 2:1, 3:1, and 4:1 molar ratio of water to TEOS. The pre-hydrolyzed TEOS precursor samples gel much more quickly compared to the TEOS precursor. With the molar ratio of water to TEOS oligomer = 4:1, gelation occurs in 15 minutes. For reaction stoichiometries of 3:1 and less, gelation occurs within 5 minutes. The basic water solution is fully solubilized in the inverse micelle solution before gelation. The clusters are embedded in the gels upon gelation. The solutions are transparent and clear (yet colored) at all water concentrations and no cloudiness or indication of phase separation is visibly apparent.

All wet gels are transparent, homogeneously deep purple, viscous, and undergo little to no syneresis (Figure 2). Drying to form xerogels causes the gels to shrink 1/3 to 1/2 of their original volume independent of reaction stoichiometry and precursor type. Deep purple monoliths are formed. Drying to form aerogels causes little shrinking in pre-hydrolyzed TEOS gels and shrinks TEOS precursor gels by 1/2 of their original volume. Aerogels appear chalky and light purple, but regain their color when wetted.

The synthesis produces highly dispersed, monodispersed, nanometer sized gold particles embedded in silica matrices (Figure 3). Particle size is tested as a function of reaction stoichiometry, the gel precursor type, the drying procedure, and washed vs. unwashed samples. Particle size is independent of reaction stoichiometry, gel precursor type,

and the effect of washing (Table 1). The size distribution of a typical sample is relatively narrow (Figure 4). For washed samples, particle size is independent of the drying procedure (xerogels vs. aerogels). It appears that the particles are washed out of samples made with the pre-hydrolyzed form of the TEOS precursor. Few are observed within the gel by TEM. Particle size is roughly twice as large in gels as in pure inverse micelle solutions of DDAB and toluene⁶⁷. Electron diffraction from the clusters show five rings that index to the presence of FCC-Au. The relative diffraction peak radii for the five rings are 1.0, 0.86, 0.61, 0.52, and 0.40. It should be noted that our work with Pd systems indicate 0.01M metal salt precursor solutions result in approximately 1 wt. % Pd on SiO₂ materials as measured by atomic adsorption spectroscopy.

Monitoring cluster size in solution and in wet gels as a function of time provides information on the growth mechanism and the effect of gelation on particle growth. Size is monitored by measuring the gold plasmon resonance observed by UV-visible spectroscopy. Colloidal sized gold in solution exhibits a plasmon resonance between 500 and 550 nm depending on particle size. Increasing particle size results in a red shift in the plasmon resonance. Immediately after salt reduction, the gold plasmon resonance occurs at 502 nm in gel precursor, DDAB, and toluene solutions. The plasmon resonance red shifts to 509 nm after three hours and stabilizes at 526 nm after seven days (Figure 5). If gelation is initiated to a new sample by adding TBAOH/water to the solution, the gold plasmon resonance is 518 nm and 527 nm after three hours and after seven days, respectively (Figure 5). The particles slowly grow after the reduction before stabilizing in both un-gelled and gelled samples. The particle growth mechanism is faster, but the final particle size is unaffected by the gelation process (assuming constant medium refractive indices between ungelled and gelled samples).

Surface areas as determined by N₂ porosimetry are strongly dependent on the reaction stoichiometry, surfactant concentration, and gel precursor type. With some exceptions, aerogel surface areas increase as the molar ratio of water to gel precursor increases and as surfactant concentration decreases (Figures 6 and 7). Aerogels formed with pre-hydrolyzed TEOS precursor exhibit higher surface areas than when formed with TEOS (compare Figures 6a and 7a). When gold is embedded in the silica matrices, the surface areas increase and follow the same trend with reaction stoichiometry (Figures 6b and 7b). To determine if surfactant changes the reaction chemistry and thus the material properties or obstructs the silica surface, the materials were washed with warm hexanol. Surface areas increase and

follow similar trends with reaction stoichiometry (Figures 6b and 7b). TEOS xerogels exhibit negligible surface areas (not shown). Pre-hydrolyzed TEOS xerogels (not shown) exhibit similar surface areas and trends as do pre-hydrolyzed TEOS aerogels (Figure 7a). The silica matrices are mesoporous with pore diameters ranging from 5 - 30 nm.

Discussion

SAXS indicates that inverse micelles formed in DDAB/toluene solutions are small ($R_g = 0.49$ nm) with only a few surfactant molecules forming each aggregate structure. SAXS also indicates that the addition of gold salt or the addition of both gold salt and TEOS have no effect on aggregate size. However, SAXS is not observed when only TEOS is added to inverse micelle solutions. TEOS alone causes dissolution of the surfactant aggregate structures. TEOS is miscible in toluene, but relatively polar. The driving force for surfactant aggregation decreases as the gel precursor acts as a co-solvent for toluene and the polar moieties of the surfactant. It is likely that there is some effect of TEOS on gold salt inverse micelle structures, and the effect is offset by the salt or not detected due to weak scattering.

The size of particles formed by the reduction of metal salts in typical inverse micelle solutions depends on the inherent size of the inverse micelles and the material exchange rate between inverse micelles. Both micelle size and the material exchange rate depend on experimental parameters like the surfactant/solvent system, the addition of water which swells the inverse micelles, and the salt to surfactant ratio. Particle size control is often complicated but feasible. Our work with the TEOS, DDAB, and toluene system establishes three results: (1) the reduction reaction in TEOS inverse micelle solutions and in gelled samples produces particles roughly two times larger than particles formed in only DDAB/toluene mixtures⁶⁷, (2) the gelation has little effect on the final particle size as determined by the plasmon resonance, but does influence the initial growth rate, and (3) the particle size is not dependent on the gel precursor type, the reaction stoichiometry, or the drying method.

The effect of TEOS on the inverse micelle structure and the cluster growth and stabilization processes is the likely cause for the increased particle size in the mixtures studied here compared to particles synthesized in traditional DDAB/toluene mixtures. Cluster nucleation and growth is sustained within the inverse micelles. A change in micelle structure and the growth and stabilization process is inferred from the effects of TEOS on the

surfactant aggregation phenomena as detected by SAXS. With the addition of the base and water solution, hydrolysis and condensation of the TEOS starts. The presence of water and the further production of water and alcohol in the reactions further destabilizes the inverse micelle structure and causes the clusters to grow at a faster rate as evidenced by the UV-visible absorption spectra. However, the final particle size is unchanged in gelled systems compared to ungelled systems. Apparently, an upper bound in the stabilization process is reached.

Some of the variables adjusted in this study to control material properties include the gel precursor type, the water to gel precursor reaction stoichiometry, and the drying process (xerogels vs. aerogels). The precursor type and drying process would have no direct effect on particle size control and none was observed. More typical methods to control particle size include, the surfactant/solvent system, the metal salt to surfactant ratio, and water content. Different surfactant/solvent systems were explored in this study (Polyoxyethylene (6) nonyl phenyl in cyclohexane and tetraethyleneglycol mono n-dodecyl ether in octane), but results were less fruitful. Silica particles resulted, or phase separation occurred with only the water rich phase gelling⁶⁸. Salt to surfactant ratios were not varied in this study. We attempted to control particle size through the use of water content. No particle size control is apparent in this effort most likely due to the destabilization of the inverse micelle structure in the presence of TEOS and the hydrolysis and condensation by-products. However, the principles of particle size control remain in this methodology and further effort is required.

Despite the lack of particle size control, the role of the surfactant is clear. The inverse micelles act to solubilize the metal salt, stabilize the clusters against growth, and solubilize the TBAOH/H₂O mixture during gelation. If surfactant is not used, little AuCl₃ is solubilized in TEOS, toluene mixtures (<< 0.001M). Reduction with LiBH₄ results in large particles that are only stable for a few hours. Finally, introduction of the TBAOH/H₂O mixture results in immediate precipitation of the clusters, phase separation of a water rich and an oil rich phase, and no gelation.

In traditional sol-gel chemistry, the relative rates of hydrolysis and condensation as effected by sol-gel processing parameters like acid vs. base catalyzed reactions and the H₂O:Si ratio control the final gel properties^{69,70}. In general, hydrolysis dominates under acid-catalyzed, low H₂O:Si ratio conditions and weakly branched, polymeric sols and gels are produced. Condensation dominates under base-catalyzed, high H₂O:Si ratio conditions to

produce highly condensed particulate sols and gels. As condensation rates increase, gelation time and gel surface areas generally decrease.

For gels prepared with TEOS in inverse micelle solutions as the $H_2O:Si$ ratio increases, gel times and surface areas increase in direct contradiction to the above stated conventional wisdom. We propose that the mechanism of gelation in the cluster solutions depends on the relative solubilities of the reactants in various 'areas' of the micro-heterogeneous solutions. TEOS is soluble in toluene, and DDAB is soluble in TEOS. Thus, TEOS most likely exists within the inverse micelle and throughout the solution. Decreased aggregation of DDAB in toluene with the introduction of TEOS as detected by SAXS supports this hypothesis. Added water is only soluble within the inverse micelle. Thus, it is likely that gelation occurs mostly at the surfactant interface. Two important results stem from this hypothesis. First, the effective $H_2O:Si$ ratio is much higher than expected from the set $H_2O:Si$ recipe resulting in high condensation rates. As a result, gels are formed in this study even at low $H_2O:Si$ recipes, and surface areas of TEOS gels are low. Second, with increasing $H_2O:Si$ ratio, the $H_2O:DDAB$ ratio increases. Increasing the $H_2O:DDAB$ leads to larger, more polydisperse inverse micelles and a subsequent decrease in the surface area of reaction. Increasing the $H_2O:Si$ recipe actually leads to a decrease in the effective $H_2O:Si$ ratio, and thus, lower condensation rates and the observed higher surface areas and faster gel times. While the proposed gelation mechanism explains the existing data, it is clearly not universally applicable. Monoliths are not formed in other inverse micelles systems, and the use of other structured surfactant phases produce periodic silica phases⁷¹⁻⁷³.

Gel surface areas increase with decreasing surfactant concentration. This result is consistent with our proposed mechanism. As surfactant concentration decreases, the $H_2O:DDAB$ ratio increases resulting in a decrease in the surface area of the reaction. Decreasing surfactant concentration leads to lower effective $H_2O:Si$ ratios, lower condensation rates, and the observed higher surface areas. It is apparent from the increase in the gel surface area after the surfactant is washed away that surfactant also blocks the available gel surface.

Gelation times are faster and surface areas are higher when the pre-hydrolyzed form of TEOS is used. These results indicate that the condensation rates for the pre-hydrolyzed TEOS precursor reactions are slower than for the TEOS precursor reactions. Steric effects are most likely responsible for the slower condensation rates with the bulkier precursor.

Conclusions

We have encapsulated nanometer sized Au particles in the micropores of xerogels and aerogels. The synthesis is a sequential reduction of a gold salt and sol-gel processing in an inverse micelle solution. The inverse micelle solution is used to solubilize the metal salt and provide a micro-reactor for the nucleation, growth, and stabilization of the nanometer sized clusters. Hydrolysis and condensation of an added siloxane precursor produces a wet gel embedding the particles. The presence of gel precursors de-stabilizes the inverse micelle structure resulting in larger particle sizes compared to typical inverse micelle synthesis techniques. Particle size control is complicated by the gel precursor effect on inverse micelle structure and the production of water and alcohol in the hydrolysis and condensation reactions. Finally, a unique gelation technique is outlined in these micro-heterogeneous solutions. Gelation occurs across the surfactant interface increasing the effective $H_2O:Si$ ratio. Gelation occurs even at low $H_2O:Si$ ratios, and condensation rates are high. Sol-gel parameters like the $H_2O:Si$ ratio and the surfactant concentration have unique and sometimes not intuitive effects on the hydrolysis and condensation rates and the resulting material properties.

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Table 1. Particle diameters in nanometers as a function of reaction stoichiometry, drying process, precursor type, and hexanol washing.

H ₂ O:Si	AEROGELS				XEROGELS			
	TEOS		pre-hydrolyzed TEOS		TEOS		pre-hydrolyzed TEOS	
	washed	not washed	washed	not washed	washed	not washed	washed	not washed
2:1	5.8 +/- 1.3	5.6 +/- 1.5	5.7 +/- 1.3	N/A	N/A	6.0 +/- 1.5	N/A	N/A
4:1	5.6 +/- 1.1	6.7 +/- 1.1	N/A	N/A	N/A	6.7 +/- 2.2	N/A	7.1 +/- 1.2

Figure 1. Small angle x-ray scattering of DDAB/toluene inverse micelles with a) added Au salt, b) added Au salt and TEOS, c) Au clusters in the presence of TEOS. No SAXS is observed when only TEOS is added to DDAB/toluene. The precursor solutions show small aggregate structures ($R_g = 0.49$ nm). The cluster size as detected by SAXS ($R_g = 2.34$ nm) is slightly smaller than when measured by TEM. $q = (4\pi/\lambda)\sin\theta$ where λ is the wavelength of the radiation and 2θ is the scattering angle.

Figure 2. Picture of typical wet gels, xerogels, and aerogels with stericly entrapped Au clusters. Aged gels and xerogels retain the purple color of colloidal Au. Aerogels lose most color, but become purple when re-wet with toluene.

Figure 3. TEM of Au clusters embedded in silica matrix. The particles are 5.7 nm in diameter and highly dispersed. Particle size is roughly twice that of Au clusters synthesized in traditional DDAB/toluene solutions without TEOS. Particle size is independent of gel precursor type, drying procedure, or the $\text{H}_2\text{O}:\text{Si}$ ratio.

Figure 4. Size distribution as determined by TEM indicating average particle size and monodispersity.

Figure 5. UV-Visible spectrum of Au cluster as a function of time and gelled and ungelled samples. The plasmon resonance red shifts with time as the particles grow. The growth rate is faster in gelled samples, but final particle size is independent of the gelation process.

Figure 6. (a) BET surface area's of TEOS aerogels as a function of the H₂O:Si ratio and surfactant concentration. Au salt or clusters are not present. The materials are mesoporous. Surface areas generally increase with the reaction stoichiometry and decreasing surfactant concentration. BET surface area's of TEOS xerogels are negligible. (b) BET surface area's of TEOS aerogels as a function of hexanol washing and cluster presence. Surface area's increase when surfactant is washed from the pores. The presence of Au salt or clusters has no effect of gel surface areas.

Figure 7. (a) BET surface area's of pre-hydrolyzed TEOS aerogels as a function of the $H_2O:Si$ ratio and surfactant concentration. Au salt or clusters are not present. The trends established for TEOS materials are repeated. Surface areas generally increase with the reaction stoichiometry and decreasing surfactant concentration. Pre-hydrolyzed TEOS xerogel surface areas are smaller, but follow similar trends. (b) BET surface area's of pre-hydrolyzed TEOS aerogels as a function of hexanol washing and cluster presence. The trends established for TEOS materials are repeated. Surface area's increase when surfactant is washed from the pores. The presence of Au salt or clusters has no effect of gel surface areas.

Chapter 5

Novel Sol-Gel Based Nanocluster Materials as Catalysts in the Dehydrogenation of Propane

Abstract

We introduce two novel synthesis strategies to make nanoclusters on silica and alumina supports using inverse micelle technology and sol-gel processing. In the first method, sol-gel chemistry (hydrolysis and condensation of metal alkoxides) is performed in alcohol-free inverse micelle, cluster solutions. In the second method, clusters formed in traditional inverse micelle solutions are allowed to diffuse into pre-existing wet monoliths formed using traditional sol-gel techniques. The different materials produced are characterized and compared with respect to particle size and the substrate properties using N_2 porosimetry methods, chemisorption, atomic absorption, and transmission electron microscopy. The effect of calcination on particle and support stability is determined and discussed in terms of the metal loading and the relationship between particle size and pore dimensions. We conclude that the relative sizes of particles and pores has no clear effect on sintering behavior. Sintering appears to be predominately dependent on metal loading suggesting Ostwald ripening as the sintering mechanism, but even this analysis is oversimplified. Finally, we study the materials as catalysts in the selective dehydrogenation of propane with particular interest in stability over time.

Introduction

Nanometer sized metal clusters embedded in sol-gel prepared metal oxide substrates form a unique class of catalytic materials. Sol-gel processing loosely defines the inorganic polymerization reactions of hydrolysis and condensation of metal alkoxides to form solid metal oxides (1). Because many synthesis parameters are available, sol-gel processing offers versatility in final material properties not available by other catalyst synthesis methods (2-5). For instance, by adjusting the oxide precursor type and concentration, the water to precursor reaction ratio, and whether the reactions are acid or base catalyzed, materials with controlled surface areas and pore dimensions are produced. Formation of metal oxide colloidal dispersions (sols) precedes final condensation into monolith, particulate, or thin film gels.

Also, it is possible to tune molecular composition; SiO_2 , Al_2O_3 , TiO_2 , and other pure species, as well as mixed metal oxides are formed. Atomic metals are added later to gels when salts are ion-exchanged with terminal hydroxyl groups (6,7), through impregnation of metal salts into the gel backbone (8-11), or through the use of chelating agents (12,13). Reduced metal clusters are formed by chemical, thermal, and photolytic reduction. Again, the methods are versatile with respect to metal type, and Pt, Pd, and Ru are common.

In addition to versatility, metal cluster, sol-gel materials have specific catalytic advantages. For clusters synthesized by impregnation techniques and independent of metal type or support composition, some degree of occlusion of the particles within the metal oxide framework occurs (14-17). As a result, higher resistance to metal sintering is noticed. Also, sol-gel processing parameters have been used to control pore dimensions and particle sizes so that the particles are encapsulated (i.e., the pore size is approximately equal to the particle diameter) (18). Resistance to sintering increases without occlusion of the particles. In one case, however, lower activities were observed due to encapsulation (19). In another study, Pt/ Al_2O_3 sol-gel prepared materials showed resistance to coke formation for a number of different support and particle morphologies (20).

In this work, we test the sintering resistance, activity, and selectivity of Pt particles embedded in sol-gel formulated SiO_2 and Al_2O_3 prepared by several novel synthesis methods. In an earlier study, we synthesized Au particles in SiO_2 by combining inverse micelle techniques with sol-gel processing in nonaqueous, alcohol-free surfactant solutions (21). We have now expanded the synthesis to include the formation of Pt particles and the formation of Al_2O_3 supports. We also develop a second synthesis approach in this paper; namely, inverse micelle synthesized Pt clusters are encapsulated in *traditionally* synthesized sol-gel SiO_2 supports. The clusters are allowed to diffuse into the pores of a wet alcogel, and drying of the gel encapsulates the particles in the micropores. We describe the synthesis, characterization, and sintering behavior of these materials, and we show activity and selectivity results of these materials tested as catalysts in the dehydrogenation of propane. The synthesis methodologies we present offer the advantages of the inverse micelle technique to make clusters; namely, ultra-small, monodispersed particles of controlled, tunable sizes are formed (22,23). Also, the methodology offers a novel procedure to encapsulate particles in the support by building the metal oxide framework around the clusters.

The advantages of cluster, sol-gel derived SiO_2 and Al_2O_3 catalysts address especially important problems in selective dehydrogenation processes. In earlier studies, deactivation of SiO_2 and Al_2O_3 supported metals by surface carbon deposition and/or particle sintering is observed (25-32). Tin as a stabilizer, for instance, is often added to counter deactivation. In particle, sol-gel derived materials, we hope a natural resistance to coke formation and prevention of sintering by encapsulation will relieve some of these problems.

Experimental

Materials. The following materials were purchased from Aldrich Chemical Co. and were used as delivered: the surfactant didodecyldimethylammonium bromide (DDAB), tetraethylorthosilicate (TEOS), aluminum tri-*sec*-butoxide (97%), ethyl acetoacetate (99+%), platinum (II) chloride (98%), apolar solvent toluene (99.9+% purity), reducing agent lithium borohydride in tetrahydrofuran (2M), and a 40 wt. % tetrabutylammonium hydroxide (TBAOH) in water solution. Acid and base solutions were made from HCL certified 1N solution and concentrated HNO_3 and NH_4OH certified ACS plus from Fisher Scientific. Absolute ethyl alcohol was supplied for AAPER alcohol.

Synthesis. The synthesis of nanoclusters via inverse micelle solutions is an established technique (22,23). Metal salts are solubilized within the polar regions of inverse micelles. When a reducing agent is added, nucleation and growth of the metal particles commences. Stabilization of the growth is controlled by the surfactant, and final particle size depends on a number of synthesis variables. The synthesis of Au particles encapsulated in SiO_2 via an inverse micelle, sol-gel combined procedure was described earlier (21, Figure 1). Here, we extend the synthesis to include Pt particles and Al_2O_3 supports (Table 1).

Sequential inverse micelle, sol-gel derived Pt/SiO₂: Surfactant is added to toluene (5 wt. %) and stirred by hand shaking to form the inverse micelle solution. PtCl_2 (0.005M) and the gel precursor TEOS (0.4M) are added to the DDAB, toluene mixture, and the solution is stirred until the salt is fully solubilized. Sufficient LiBH_4 /THF solution to result in a 3:1 ratio of BH_4^- : Pt^{2+} is then injected into the salt precursor solution under rapid stirring to form the Pt clusters. The 40 wt. % TBAOH in water solution is added to the mixture 5 minutes after the addition of the LiBH_4 , and gelation is marked when the solution no longer flows under gravity. The water to gel precursor molar ratio is set at 3:1. The gels are aged

for two days at 50°C and are washed with pure toluene (soaked for several hours at 50°C, 3 cycles) to remove any unattached surfactant, metal salt, or excess reducing agent. The samples are then split in two with half being used to form xerogels and the other half being used to form aerogels. Xerogels are formed by three staged drying cycles of 40°C, 50°C, and 120°C each overnight. Aerogels are formed under supercritical CO₂ extraction. The overall molar equivalent ratio of the synthesis is Si : 3H₂O : 23 toluene : 0.2 TBAOH where the toluene actually represents 0.005M PtCl₂ in 5 wt. % DDAB in toluene.

Sequential inverse micelle, sol-gel derived Pt/Al₂O₃: The Pt/Al₂O₃ synthesis is loosely based on a previous alcohol based Al₂O₃ recipe without metal (33). DDAB is added to toluene (5 wt. %) and stirred by hand shaking to form the inverse micelle solution. The gel precursor is a 1:1 molar ratio of aluminum tri-*sec*-butoxide and ethyl acetoacetate. PtCl₂ (0.005M) and the gel precursor (0.8M) are added to the DDAB, toluene mixture, and the solution is stirred until the salt is fully solubilized. The LiBH₄/THF solution is then injected into the salt precursor solution to form the clusters under rapid stirring so that the [BH₄⁻] : [Pt²⁺] = 3:1. After 30 minutes, an aqueous 4.3M HNO₃ solution is added, and gelation is marked when the solution no longer flows under gravity. The water to gel precursor molar ratio is set at 3:1. The gels are aged for two days at 50°C and are washed with pure toluene (soaked for several hours at 50°C, 3 cycles) to remove any unattached surfactant, metal salt, or excess reducing agent. The samples are then split in two with half being used to form xerogels and the other half being used to form aerogels. Xerogels are formed by three staged drying cycles of 40°C, 50°C, and 120°C each overnight. Aerogels are formed under supercritical CO₂ extraction. The overall molar equivalent ratio of the synthesis is Al : 3H₂O : 14 toluene : 0.2 HNO₃ where the toluene represents 0.005M PtCl₂ in 5 wt. % DDAB in toluene.

Concurrent inverse micelle, sol-gel derived Pt/SiO₂: An asterisk denotes samples made by the ‘concurrent’ methodology. Here, Pt clusters are synthesized with a ‘standardized’ inverse micelle recipe. Concurrently, SiO₂ monolith is synthesized with a ‘standardized’ ethanol sol-gel recipe. The clusters are then allowed to diffuse into the gel, and the gel is dried leading to encapsulation of the particles in the micropores. To form the clusters, 0.01M PtCl₂ is solubilized in a 5 wt. % DDAB, toluene mixture. The LiBH₄/THF solution is injected into the salt precursor solution under rapid stirring so that the [BH₄⁻] :

[Pt²⁺] ratio is 3:1. To form the SiO₂ monolith, a stock solution of 61ml TEOS, 61 ml ethanol, 4.87 ml water, and 0.2ml 1M HCl is refluxed at 60°C for 1.5 hrs with stirring. The solution is stable indefinitely at 20°C, but is stored in a freezer to ensure stability. The approximate molar ratio of the solution is TEOS : 4 ethanol : H₂O : 0.007 HCl. Mixing a 10:1 volume ratio of stock solution to 0.5M NH₄OH causes gelation within minutes. The gels are aged and washed three times with excess ethanol at 50°C for several hours. To produce the cluster / SiO₂ matrix, the gels are washed three times with excess toluene at 50°C for several hours. Then, the SiO₂ is washed with excess cluster solution three times at 50°C for several hours. Xerogels are formed by three staged drying cycles of 40°C, 50°C, and 120°C each overnight. The methodology was not extended to aerogels or Al₂O₃ supports, although this should be facile.

All samples were calcined in a tube furnace under air. A two stage temperature ramp was used. The samples were held at 250°C for 1 hour and then the final temperature for 1 hour. Final calcine temperatures were determined by TGA results as discussed below.

Characterization. Metal weight fractions on the supports were determined with a Perkin-Elmer 5100PC atomic absorption spectrophotometer (AA). Thermal gravimetric analysis (TGA) was completed with a TA Instruments Thermal Analyst 2000. Platinum particle size was characterized with transmission electron microscopy (TEM) with a 300 keV Phillips CM30 electron microscope. To prepare TEM samples, dried, crushed gels were dispersed over a holey carbon substrate. N₂ porosimetry, surface area analysis (BET) of the crushed xerogels and aerogels was carried out with a Quantachrome Autosorb-6 surface analysis apparatus. BET samples were outgassed for 20 hours at 100°C. Hydrogen chemisorption was measured using a Coulter Omnisorp 100CX instrument. All chemisorption samples were calcined at 450°C in air for 1 hour except the 'concurrently' produced Pt/SiO₂* sample. Based on TGA results, the concurrently produced Pt/SiO₂* sample was calcined at 600°C for 1 hour in air. In the chemisorption unit, the samples were outgassed in flowing He at 350°C for 1 hour followed by reduction in flowing hydrogen at 400°C for 2 hours. Samples were then evacuated at 405°C for 3 hours prior to cooling to room temperature before measurement of hydrogen uptakes. Dispersions were calculated from total hydrogen uptake and converted to particle sizes by assuming spherical particles and a Pt atom surface density of 1.25×10^{15} atoms/cm². The samples used for TEM, chemisorption, and propane

dehydrogenation are from the set of samples described by the first column of the AA results (i.e., those listed first and on the left in Table 3).

Catalyst Testing. All samples were initially in the form of a single monolithic piece. These monoliths were carefully ground and sieved to obtain a -60/+80 mesh fraction for use in reactor testing. The remainder of the sample disintegrated into a fine powder which was used for the majority of the physical characterization. The 'sequentially' produced Pt/SiO₂ sample was tested for propane dehydrogenation in the powdered state since careful grinding of the monolith did not result in a sufficient quantity of -60/+80 mesh material.

Activity measurements were made in a fixed bed flow reactor. The reaction sequence begins with heating of the catalyst bed to reaction temperature (typically 525-575°C) in flowing nitrogen. Once temperature is reached, the nitrogen flow is stopped and reactant flow begins. Propane is fed through a calibrated mass flow controller and is preheated in a 1/8" coiled stainless steel tube at the reactor inlet. The preheated reactant then enters a 3/8" stainless steel tube containing catalyst. The catalyst is supported on a fine steel mesh located at the bottom of the reactor. Depending on catalyst density, 0.3-0.6 gm of catalyst are typically used to give a catalyst bed volume of 3.6 cc. After exiting the reactor, product gases are exhausted through the sample loop of a gas chromatograph, where C1-C4 hydrocarbons are analyzed. Gas sampling and analysis was performed at 20 minute intervals. Oxygen regeneration and hydrogen reduction treatments were often used alternately to achieve maximum catalyst activity. Reactor temperature is measured by three thermocouples externally attached to the top, middle, and bottom of the reactor, and one thermocouple located inside the reactor at the bottom of the catalyst bed. No significant differences were ever observed among the four thermocouples indicating that the reactor was operating in an isothermal mode.

In general, no consistent activation behavior was observed among the various catalysts. In most cases, hydrogen reduction was found to be beneficial, but for one catalyst (Pt/Al₂O₃ aerogel) hydrogen reduction actually decreased activity. Also, the rate of deactivation of the catalysts varied, as did the efficacy of air oxidation to burn off carbon and regenerate the catalysts. Because of these variations among the catalysts, quantitative comparisons are difficult. The reported activities are therefore the best activities that were achieved on each catalyst, and the prior history of the catalysts varies substantially. In addition to variations in pretreatment, variations in both weight hourly space velocity

(WHSV) and gas hourly space velocity (GHSV) also occurred among the catalysts. The GHSV variations arise as a result of a loss of catalyst bed volume during the reactions, which prevents an *a priori* prediction of the true GHSV. The WHSV velocities result from both the bed volume changes, as well as differences in density among the catalysts. In general, the initial bed volume was kept constant at ~3.6 cc, provided a sufficient quantity of the -60/+80 mesh fraction was available. As a result, variations in WHSV are much greater than variations in GHSV. In order to compensate for the space velocity variations, both propane conversion/selectivity and propane turnover frequencies (TOF, calculated from measured dispersions, WHSV's and Pt loadings) are reported. The conversion/selectivity data allow comparison with a commercial 7.5 wt % Cr/Al₂O₃ catalyst (Girdler G-41), while the TOFs allow comparisons of intrinsic activity among the catalysts. Because conversions are generally less than 10% and therefore far from equilibrium (equilibrium conversion is ~30% at 550°C), the use of TOFs should compensate for space velocity differences and provide a valid measure of intrinsic activity.

Results

TGA weight loss results indicate a large difference between the samples produced by the sequential reactions of metal reduction, hydrolysis and condensation and the SiO₂* sample produced by the concurrent synthesis (Figure 2). The weight loss is large on an absolute scale for the samples produced in the sequential methodology with greater than 50% weight loss. Most weight loss occurs below 400°C so 450°C was chosen as the calcination temperature. The Pt/SiO₂* sample shows two weight loss regions, at approximately 350°C and 550°C. Only 15 wt. % of the total sample is lost. 600°C was therefore chosen as the calcination temperature to ensure complete weight loss prior to analysis and testing. TGA results did not differ in air compared to an inert He atmosphere.

BET surface area results indicate a large difference between the samples produced by the sequential reactions and the SiO₂* sample produced by the concurrent synthesis (Figure 3). 'Sequentially' produced Pt/SiO₂ and Pt/Al₂O₃ xerogels and aerogels are nonporous and show little surface area. Upon calcination surface area increases greatly but then decreases during propane dehydrogenation. The same trend is observed with the Pt/SiO₂* xerogel, but these samples are mesoporous with high surface areas even prior to calcining. Differential

pore size analysis shows the Pt/SiO_2^* xerogel pores are larger with a broader pore size distribution than the other samples (Figure 4).

Particle size was studied with TEM (Figure 5) and H_2 chemisorption. TEM analysis indicates the initial Pt/SiO_2 xerogel, Pt/SiO_2 aerogel, and the Pt/SiO_2^* xerogel particle diameters are 1.5 nm, 1.5 nm, and 2.0 nm, respectively. After calcination, TEM particle sizes increase to 4.5 nm, 4.0 nm, and 3.5 nm, respectively. The Al_2O_3 xerogel and aerogel show aggregated metal particles throughout the substrate. No individual particles are observed. After calcination, individual particles are observed. The $\text{Pt/Al}_2\text{O}_3$ xerogel and aerogel particle diameters are 3.5 nm and >50 nm, respectively. Particle sizes determined by chemisorption are compared to TEM results (Table 2). For the calcined Pt/SiO_2 and Pt/SiO_2^* aerogels and the calcined $\text{Pt/Al}_2\text{O}_3$ xerogel, the measured dispersions are 12.7%, 22.3%, and 11.1%, respectively, corresponding to particle diameters of 4.5, 2.5, and 4.9 nm, in close agreement with TEM results. H_2 chemisorption measurements were not possible on the uncalcined samples due to severe outgassing problems.

Atomic absorption spectroscopy was used to determine metal loadings. Three separate sample sets were tested (Table 3). The first set of numbers (on the left) represent the samples used for BET, TEM, chemisorption, and activity testing. For the Pt/SiO_2 xerogel, Pt/SiO_2 aerogel, Pt/SiO_2^* xerogel, $\text{Pt/Al}_2\text{O}_3$ xerogel and $\text{Pt/Al}_2\text{O}_3$ aerogel, the metal loadings are 0.24 $\pm$ 0.08 wt. %, 0.23 $\pm$ 0.07 wt. %, 0.25 $\pm$ 0.03 wt. %, 0.22 $\pm$ 0.14 wt. %, and 0.24 $\pm$ 0.14 wt. %, respectively. Reproducibility is poor, but it is particularly poor with the Al_2O_3 samples indicating an extreme sensitivity of metal loading to synthesis conditions. It is possible to increase metal loadings by increasing precursor salt concentrations, but particle size and polydispersity generally increases.

The platinum samples were tested for propane dehydrogenation at 550°C (Table 4, Figure 6). Note the WHSV's and pretreatment conditions differ significantly among the six samples studied, as discussed earlier. Because of these variations, quantitative activity comparisons cannot be made, and the activity measurements serve mainly to demonstrate the potential of this class of dehydrogenation catalysts, as will be discussed later.

Discussion

Synthesis Effects on Support Properties. TGA results indicate that for samples synthesized via the sequential methodology (i.e., sol-gel processing is completed in an

alcohol free, cluster, inverse micelle system) over 50 wt. % of the sample is thermally lost by about 400°C. The only single component present to such extent in the initial reaction mixture is the gel precursor TEOS leading us to believe that the large weight loss in the final product is due to unreacted TEOS monomer. It is likely that the unreacted TEOS results from inefficient hydrolysis and condensation reactions in apolar, alcohol free inverse micelle mixtures. The same argument holds for the alumina system.

In samples synthesized via the concurrent methodology (i.e., using a well defined sol-gel process taking place in ethanol followed by the addition of an already prepared cluster, inverse micelle solution), TGA results show two weight loss regions with a total loss of approximately 12 wt. %. No weight loss occurs before 200°C indicating all solvents were removed in the drying procedure. At approximately 200°C, there is a loss of about 6 wt. % mostly due to the thermal breakdown of surfactant. There is little unreacted TEOS due to the highly studied, efficient synthesis procedure used. Finally, starting at approximately 400°C, about 5 wt. % is lost probably due to the thermal breakdown of terminal hydroxy groups.

The N₂ porosimetry results support our TGA analysis. In sequentially synthesized samples, no measurable surface area exists before calcination, because of the presence of surfactant and unreacted TEOS in the pores. The surface area increases greatly upon thermal removal of these components. In the concurrently synthesized sample, surface area increases slightly due to removal of surfactant.

In our earlier work, we studied the synthesis parameters that govern final material properties in sol-gel processed materials in inverse micelle systems (21). We observed, as we observe here, that monoliths form at much lower water:TEOS ratios than in alcohol based systems. We speculate that monoliths form due to localized high water concentration regions in the inverse micelles with the hydrolysis and condensation reaction mechanisms taking place across surfactant monolayers and in transient micelle material exchange states. It appears in these mechanisms that only a fraction of TEOS is exposed to the water rich regions and participates in the reaction, thus providing a high effective water concentration as we speculated earlier. Also in our previous work, we noticed a reverse trend in surface area with the water:TEOS ratio than is typically found in alcohol systems. As the samples in our previous study were not calcined, it is unfortunate that we were measuring surface areas of materials obstructed with unreacted TEOS. True monolithic surfaces were not measured calling into question the analysis of the effect of the water:TEOS ratio in that work.

Synthesis Effects on Particle Properties. TEM results indicate that the metal reduction reaction to form clusters is highly efficient in the SiO_2 samples. Small, monodisperse particles with diameters of less than 2 nm is typical for Pt particles synthesized in pure DDAB/toluene inverse micelle mixtures. Visually, an abrupt color change and rapid H_2 gas production indicate an instantaneous reaction. In the sequential reduction methodology, the presence of TEOS, the hydrolysis and condensation reactions, and drying have no effect on final particle size. In the concurrent reduction methodology, diffusion of the particles into the gel network and drying have no effect on final particle size.

The reduction reaction efficiency decreases substantially in alumina systems. Visually, color change and H_2 production occurs over at least one-half hour. In reactions in inverse micelles, it is known that slow nucleation and growth results in large, polydisperse size distributions. We observe similar results here. Further, the particles are aggregated indicating surfactant stabilization was not effective. A concurrent reaction mechanism where the particles are synthesized separately maybe required to obtain highly dispersed Pt particles in aluminum gels. We have not yet studied a concurrent reaction mechanism for alumina.

Sintering and the Relationship between Particle and Pore Sizes. In the sequentially synthesized SiO_2 samples, particle size and pore dimensions are relatively equal before calcination. After calcination, the particle size increases to approximately three times larger than the pore dimensions even at the relatively low calcination temperatures of 450°C . Agreement between TEM and chemisorption results indicate the particles are moving out of the pores and unto the surface of the substrates rather than being occluded in the support structure.

In the concurrently synthesized Pt/SiO_2 xerogel, particle size is approximately three times smaller than the pore dimension before calcination. After calcination, the particle size and the pore dimensions grow only slightly (i.e. no sintering), the particle size remains approximately three times smaller than the pore dimension. Agreement between TEM and chemisorption results indicate no particle occlusion.

It is clear from the above results that no clear relationship between particle size and pore dimension effects sintering behavior. Previous work on Pt/SiO_2 catalysts (18) states that there is a strong relationship between particle size and pore size, and that for low metal loadings sintering is prevented when particle sizes match average pore diameters. The conclusions of this previous work were based upon three distinct cases: 1) 'high' Pt loadings

with particle sizes smaller than the pore size; 2) 'high' Pt loadings with particle sizes matched to the pore size; and 3) 'low' Pt loadings with particle sizes matched to the pore size. The fourth case, 'low' Pt loadings with particle sizes smaller than the pore size, was not studied. Thus, the possibility that sintering resistance is merely a reflection of Pt loading only could not be eliminated. The concurrently synthesized Pt/SiO_2^* xerogel reported here falls into the fourth case, and demonstrates resistance to sintering even though the particle size and pore size are not well matched.

The results from the previous work (18) combined with our results for the Pt/SiO_2^* sample is further evidence that there is no clear relationship between particle size and pore dimension effecting sintering behavior. The combined results with Pt/SiO_2^* provide strong evidence that sintering is a function of metal loading. Metal particle sintering on support materials occurs by two mechanisms: (1) particle aggregation due to particle diffusion along the support, and (2) Ostwald ripening where large particles grow at the expense of smaller particles due to atomization and diffusion between particles. As Ostwald ripening is concentration dependent, it appears Ostwald ripening is the predominant mechanism for sintering. It may be possible to effect vapor pressures and influence surface tensions by encapsulation such that the mechanism for Ostwald ripening is hindered, but there is yet no experimental evidence to quantify the effects. The relationship between the initial particle size and pore dimension may effect particle to particle diffusion, thus somewhat preventing the less important form of sintering. Paradoxically, however, sintering is observed in the Pt/SiO_2 xerogel where metal loading is low (while the average metal loading for the Pt/SiO_2 xerogel is relatively high recall the sample tested in BET, TEM, chemisorption, and catalyst testing is the one reported in the left most column of Table III). It appears that sintering is more complicated than concentration dependent Ostwald ripening also.

The effect of calcination on particle size in alumina samples is counter intuitive. Encapsulation is not an issue, because the particles and aggregates are much larger than the pore dimensions. With calcination, aggregated particles separate into smaller domains. In alumina samples, random particle diffusion upon calcination apparently decreases the particle concentration gradient resulting in more dispersed, smaller particles.

Catalyst Testing. The propane conversion data in Table 4 shows that the uncalcined Pt/SiO_2^* xerogel results in the highest conversions of any of the Pt catalysts. Note, however, that the space velocities for this catalyst are substantially lower than for the other catalysts.

Of the other catalysts, the calcined Pt/SiO_2^* xerogel gives the highest conversions, although the performance of the $\text{Pt/Al}_2\text{O}_3$ aerogel is comparable if a linear correction for the difference in GHSV is made. Even with corrections for GHSV differences, none of the catalysts performed comparably to the commercial Cr catalyst, which achieved near equilibrium conversion with a very high selectivity. In general, the selectivities of the Pt catalysts are inferior to the commercial catalyst, particularly at the higher conversions. The selectivities are, however, substantially better than those reported by de Miguel, *et al.* (27) at 853 K, and comparable to those reported by Bariãs, *et al.* (25) at 519°

On a TOF basis, the calcined Pt/SiO_2^* xerogel and the $\text{Pt/Al}_2\text{O}_3$ aerogel have by far the highest activities. No calculations of TOFs are possible for the commercial catalyst or the uncalcined Pt/SiO_2^* xerogel due to the absence of an appropriate active site counting method for the former, and severe outgassing problems for the latter, which prevent hydrogen uptake measurements. The magnitudes of the TOFs on the two best catalysts are substantially higher than those reported by Bariãs, *et al.* (25) for traditionally prepared Pt/SiO_2 or $\text{Pt/Al}_2\text{O}_3$ catalysts at 519 K. Furthermore, while Bariãs, *et al.* noted rapid deactivation over a period of ~30 minutes, resulting in loss of more than 90% of the initial activity, we find less than a 50% loss in activity for the $\text{Pt/Al}_2\text{O}_3$ aerogel, and only a 10% loss for the Pt/SiO_2^* xerogel catalyst over a similar time period. In fact with the exception of the Pt/SiO_2 aerogel, all of our catalysts show a remarkable resistance to deactivation (Table 2). This resistance is comparable to that reported by Bariãs, *et al.* (25) for traditionally prepared $\text{PtSn/Al}_2\text{O}_3$ and PtSn/SiO_2 catalysts. Furthermore, the initial activities of our two best catalysts are comparable to those of the $\text{PtSn/Al}_2\text{O}_3$ and PtSn/SiO_2 catalysts of Bariãs, *et al.* Thus, through the use of sol-gel techniques to prepare Pt clusters inside SiO_2 and Al_2O_3 xerogels and aerogels, we have obtained performance that is comparable to $\text{PtSn/Al}_2\text{O}_3$ and PtSn/SiO_2 catalysts prepared by incipient wetness. Since deactivation in these catalysts is traditionally attributed to coke formation and the addition of Sn is known to inhibit coke formation (25,27), it is probable that the sol-gel derived catalysts also somehow inhibit coke formation, although as yet we have no physical evidence to support this hypothesis. This conclusion is consistent with previous work (20).

Conclusions

We introduce two novel synthesis strategies to make nanoclusters on silica and alumina supports using inverse micelle technology and sol-gel processing. In the first methodology, sol-gel chemistry is performed in alcohol-free inverse micelle, cluster solutions. Hydrolysis and condensation reactions are inefficient, but nonetheless monoliths form around the clusters. In the second method, clusters formed in traditional inverse micelle solutions are allowed to diffuse into pre-existing monoliths formed using traditional sol-gel techniques (hydrolysis and condensation in ethanol). The two techniques give materials with similar cluster sizes, but different pore dimensions. We conclude that the relative sizes of particles and pores has no clear effect on sintering behavior. Sintering appears to be predominately dependent on metal loading suggesting Ostwald ripening as the sintering mechanism, but even this analysis is oversimplified. Our samples achieved relatively high catalytic activity and stability over time for propane dehydrogenation indicating that the synthesis methods described here result in catalyst performance more typical of PtSn/Al₂O₃ or PtSn/SiO₂ catalysts prepared by impregnation. This behavior may indicate a high resistance to coke formation in our catalysts.

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Table 1. Summary of samples, recipes, and post-synthesis treatments used in this study. The asterisk always refers to the sample synthesized by the ‘concurrent’ procedure explained in the text.

Catalyst	Recipe	Post-Synthesis Treatment
Pt/SiO ₂ xerogel	Si • 3H ₂ O • 23DDAB/Tol • 0.2 TBAOH • 0.005M PtCl ₂	washed; 40, 50, 120°C air drying stages
Pt/SiO ₂ aerogel	Si • 3H ₂ O • 23DDAB/Tol • 0.2 TBAOH • 0.005M PtCl ₂	Supercritical extraction: CO ₂ , 40°C
Pt/Al ₂ O ₃ xerogel	Al • 3H ₂ O • 14DDAB/Tol • 0.2HNO ₃ • 0.005M PtCl ₂	washed; 40, 50, 120°C air drying stages
Pt/Al ₂ O ₃ aerogel	Al • 3H ₂ O • 14DDAB/Tol • 0.2HNO ₃ • 0.005M PtCl ₂	Supercritical extraction: CO ₂ , 40°C
Pt/SiO ₂ * xerogel	Si • 4EtOH • H ₂ O • 0.007HCl; 0.1H ₂ O • 0.05M NH ₄ OH; 0.01M PtCl ₂	washed; 40, 50, 120°C air drying stages

Table 2. Summary of particle diameters as determined by TEM. TEM results are compared to chemisorption results for the post-calcined samples.

Catalyst	Particle Diameter (nm)		
	Post-Synthesis	Post-Calcination	
	TEM	TEM	Chemisorption
Pt/SiO ₂ xerogel	1.5	4.5	4.5
Pt/SiO ₂ aerogel	1.5	4.0	
Pt/Al ₂ O ₃ xerogel	aggregated	3.5	4.9
Pt/Al ₂ O ₃ aerogel	aggregated	> 50	
Pt/SiO ₂ * xerogel	2.0	3.5	2.5

Table 3. Summary of weight percent metal loading as determined by atomic absorption (AA). Multiple measurements show the reproducibility of the synthesis.

Catalyst	Metal Loading (wt. %)	
	uncalcined	calcined
Pt/SiO ₂ xerogel	0.16, 0.25, 0.31	0.32, 0.50, 0.62
Pt/SiO ₂ aerogel	0.15, 0.27, 0.26	0.43, 0.77, 0.74
Pt/Al ₂ O ₃ xerogel	0.20, 0.10, 0.37	0.50, 0.25, 0.92
Pt/Al ₂ O ₃ aerogel	0.37, 0.10, 0.26	1.06, 0.29, 0.74
Pt/SiO ₂ * xerogel	0.25	0.30

1. Formation of nanoclusters by the inverse micelle technique.

2. Formation of the gel around the nanoclusters.

Figure 1. Schematic representation of the 'sequential' reaction method to form clusters embedded in gels. All reactants are initially present and the two reactions (metal salt reduction and sol-gel processing) are carried out in sequence. In the 'concurrent' reaction methods, the two reactions are carried out separately and the two products (clusters in inverse micelles and wet monoliths) are mixed.

a) SiO₂ samples

b) Al₂O₃ samples

Figure 2. TGA results to determine weight loss with temperature. In sequentially produced samples, over 50 wt. % is lost due to unreacted gel precursor left by the inefficiency of sol-gel processing in inverse micelle solutions. The concurrently produced samples, using a well studied synthesis procedure to produce silica monoliths results in an efficient reaction and little loss. a) SiO₂ samples. b) Al₂O₃ samples.

a) SiO₂ samples

b) Al₂O₃ samples

Figure 3. N₂ porosimetry results to determine surface area in dried, calcined, and reacted samples. In sequentially produced samples, removal of the unreacted TEOS upon calcination opens the pores and a large increase in surface area is observed. a) SiO₂ samples. b) Al₂O₃ samples.

Figure 4. Pore size distributions calculated from the porosimetry results. In sequentially produced samples, the pores are small and monodisperse falling on the limit of what is defined as mesoporous (pore sizes approximately 1nm in diameter). There is little effect of calcination. In the concurrently synthesized sample, pore sizes are larger and more disperse. The average pore size is approximately 6nm in diameter before calcination (not shown) and grows upon calcination to approximately 11nm in diameter. a) sequentially produced SiO_2 samples. b) concurrently produced SiO_2 sample. c) sequentially produced Al_2O_3 samples.

Pt / silica aerogel

10nm

Figure 5. Representative TEM picture of inverse micelle cluster, sol-gel derived materials. Sample shown is of Pt clusters on a SiO_2 aerogel support before calcination.

Figure 6. Activity of samples with respect to propane conversion to propene. Air regeneration and H_2 reduction was used to achieve maximum activity for each sample. Two samples (Pt/SiO₂* xerogel and Pt/Al₂O₃ aerogel) show relatively large activities compared to previously published results. In addition, stability remains good up to 30 minutes (not tested longer). Previous results show complete loss of activity in less than 10 minutes unless a stabilizer is used. Increased stability could be due to prevention of char on the sol-gel derived supports.

Chapter 6

Novel Sol-Gel Based Nanocluster Materials as Catalysts in the Liquefaction of Coal

Abstract

We continue our systematic characterization and catalyst activity testing of highly dispersed heterogeneous nanostructured materials by applying the above developed technology to the synthesis of iron based supported xerogel materials. Several iron based samples were synthesized and tested as catalysts in coal liquefaction.

Results

Synthesis of Fe Based Nanostructured Heterogeneous Catalysts

We have studied the synthesis of two different iron phases in the heterogeneous catalyst recipes. We have synthesized Fe^0 particles and applied a new synthesis in an attempt to make FeOOH . The synthesis of heterogeneous Fe^0 /xerogel catalysts is fairly straightforward, because we originally synthesized Fe^0 in inverse micelles by the reduction of FeCl_3 with LiBH_4 . Some characterization is necessary, however, because of the addition of basic water in sol-gel processing. A series of four Fe^0 samples of both silica and alumina have been synthesized in the past quarter.

The synthesis of FeOOH is desired, because FeOOH has been shown to be the best Fe based catalyst precursor for coal liquefaction. FeOOH in aqueous solutions is typically synthesized by regulating the pH to 8 and reacting an Fe salt with a solution of H_2O_2 . The synthesis is tricky in inverse micelles with sol-gel processing, because pH in an apolar solvent is undefined and acids and bases catalysis sol-gel processing making pH regulation difficult. Nonetheless, we modified a literature preparation to our synthesis. We have not been able to identify FeOOH by xray diffraction. Modification to the synthesis is needed.

Coal Liquefaction Testing

We have completed initial coal liquefaction catalyst testing with the highly dispersed heterogeneous materials. The ultimate goal will be to develop plastic as a hydrogen source to increase the economic viability of coal liquefaction in a simple, single stage, single catalyst

process. The highly dispersed heterogeneous catalysts are ideal for coal / plastic coprocessing, because they are bifunctional. The dispersed metal acts to hydrogenate the coal, and the silica or alumina support depolymerizes the plastic waste.

Results of the initial liquefaction tests are shown below. There are two trends that are important. First, overall conversion increases markedly with platinum. Strong effects on metal type indicate that despite being embedded in the support the metal is still highly dispersed with respect to the coal. Iron based materials are far less active than platinum. The second important result is that Al_2O_3 as a support consistently shows lower conversion to oil fractions than thermal runs. Al_2O_3 is a far better cracking catalyst than SiO_2 due to surface acidity. We believe that cracking is occurring at a much faster rate than hydrogenation with Al_2O_3 causing regressive reactions with the coal. A support mixture of SiO_2 and Al_2O_3 could be used to balance the hydrogenation and cracking rates to optimize conversion.

Metallic iron shows only an increase of roughly 7-10% in overall conversion compared to thermal. We are currently preparing FeOOH and FeS_2 for testing in the heterogeneous recipes. It is stated in the results below that FeOOH was tested. A nominal recipe for FeOOH was attempted, but characterization does not indicate the presence of FeOOH . The recipe must be adjusted further to the inverse micelle technique. FeS_2 formation has been completed, but testing is not available yet.

Finally, two different $\text{Fe}/\text{Al}_2\text{O}_3$ samples are shown in the results below. In the first as described in the experimental section, a two-step sequential reduction of an Fe salt followed by hydrolysis and condensation of aluminum alkoxides in a microemulsion forms nanoparticles in an alumina gel. In the second synthesis technique, nanoparticles and gels are formed in separate flasks, and the clusters are allowed to diffuse into the gel micropores upon mixing. Steric entrapment and thermal stability result from controlled drying and structural collapse of the gel. This latter methodology is currently being developed further.

The above results indicate two important trends (Figure 1). First, overall conversion increases markedly with platinum. Strong effects on metal type indicate that despite being embedded in the support the metal is still highly dispersed with respect to the coal. Iron based materials are far less active than platinum. The second important trend is that Al_2O_3 as a support consistently shows lower conversion to oil fractions (C7 sols) than thermal runs despite moderate increases in overall conversion. Al_2O_3 is a far better cracking catalyst than

SiO₂ due to surface acidity. We believe that cracking is occurring at a much faster rate than hydrogenation with Al₂O₃ causing regressive reactions with the coal.

It should be possible to optimize surface acidity to increase overall conversion and conversion to oil fractions by making supports of SiO₂ and Al₂O₃ mixtures. In earlier work (Quarterly Report, 4/97), we described the synthesis of SiO₂/Al₂O₃ support mixtures. This quarter we prepared metallic Fe clusters on a 10 wt. % Al₂O₃/SiO₂ support. The nanoparticles and gels were formed in separate flasks, and the clusters were allowed to diffuse into the gel micropores upon mixing. Steric entrapment and thermal stability resulted from controlled drying and structural collapse of the gel. This synthesis methodology provides the best catalyst as noticed from the differences between Fe⁰/Al₂O₃ 1 and Fe⁰/Al₂O₃ 2 (Figure 1. see Quarterly Report, 1/98 for details).

Coal liquefaction results indicate the effect of mixing Al₂O₃ and SiO₂ in the support did not have the intended effect (Figure 2). Percentage of oil fractions is roughly the same as with pure SiO₂ supports and slightly ahead of the thermal result. Overall conversion is down markedly, however. As shown in last quarters' results, overall conversion is highly dependent on metal type. Our leading explanation for this quarter's result is that the support acidity is effecting the nature of the metallic Fe composition, but clearly more investigation is needed.

Figure 1. Conversion to THF sols and C7 sols in the liquefaction of DECS-17 coal. Highly dispersed Fe materials are compared to platinum, and the effects of SiO₂ vs. Al₂O₃ support materials are determined.

Appendix

Publications

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