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Enthalpy of hydration of mgso4

Hydrated enthalpy is an energy change for converting one mole of an anrus substance into one mole of hydrated substance. To find this number, it is necessary to first calculate the enthalpy of elution of each substance separately, and then find a difference between the two. The enthalpy of dissolution is an energy change that dissolves one mole of the substance in water. It is calculated using the temperature change of water, the heat capacity of the substance, and the weight of the mixture. In this experiment, mgso4 and mgso4 · 7h2o were used to calculate the enthalpy of hydration between the two. Experiment A Synroam cup and stirring bar were obtained first and weighed together. This mass was recorded. 100 mL of deionized water was measured in a step-by-step cylinder and put in a cup with a stirring rod. The cup was weighed again and this new mass was recorded. The cup was then placed on a mixing plate set high from the medium and its temperature was recorded every 30 seconds for 4.5 minutes. An unknown amount of MgSO4 salt was added to the cup. The cup was placed in a mixing plate and kept medium to high, and its temperature was recorded for 15 minutes per minute. Finally, we weighed the cup and recorded the final mass. MgSO4 and MgSO4 were repeatedly arranged at 7H2O. Results Measurement MgSO4 · 7 H2O Trial MgSO4 Cup and Stirring Rod Trial Mass (g) 7.85 7.41 Cup Mass, Stirring Rod, Water (g) 107.21 106.70 Mass of Water (g) Mass of Cup, Stirring rod, stirring rod, water, salt (g) 119.50 113.06 Mg salt mass (g) 12.29 6.36 solated molar mass (g) 246.476 120.369 Soled mole addition (mole) 0.3 0.4986 0.0528 Salt and water mass (g) 11.88 15.15 Temperature Mixing time (°C) 20.90 21.60 Extrapolation final temperature of reaction mixture (°C) 19.27 32.15 65 ΔT = Tfinal – Tinitial (°C) 1.63 12.05 Reaction mixture heat capacity (J/(g°C)) 3.84 3.84 Heat transferred during dissolution -4890.4Jdissolution (J/mole) 14000. (14.0 kJ) -92600. (-92.6 kJ) Enthalpy of Hydrogen -106.6 kJ Time (minutes) Temperature of MgSO4 · 7H2O solution (°C) 0.0 n/a n/a 0.5 20.90 21.63 1 0.20 20 21.63 2 0.20 20 21.62 3 0.20 20 21.60 4 0.20 20 21.59 5 0.20 20 21.57 6 0.20 21.54 7 0.20 21.52 8 0.20 21.50 9 0.20 21.48 10 0.20 21.46 11 0.20 21.44 12 0.19 30.10 13 0.19 30.10 14 0.19 30.10 15 0.19 30.10 16 0.19 30.10 17 0.19 30.10 18 0.19 30.10 19 0.19 30.10 20 0.19 30.10 21 0.19 30.10 22 0.19 30.10 23 0.19 30.10 To find the mass of water, I subtracted the weight of the cup with the stirring rod from the weight of the cup with the weight of the salt used. To find the weight of the salt used, I subtracted the weight of the stirr the water from the final weight of the cup. To find the mole of the melt used, I divided the mass of salt by its molar mass. To find the temperature change, the final temperature was deducted from the initial temperature. To find the heat capacity Q of the reaction mixture, equation Q = (mass of the mixture) * (heat capacity of the mixture) * (ΔT) was used. To find ΔHdissolution, a formula of ΔH = Q / (moles of solum) was used. Finally, the hydration enthalpy was calculated and ΔDissolution of MgSO4 was deducted from ΔHdissolution of MgSO4 · 7H2O. Discussion/Conclusion MgSO4 was surprised that mgso4 · 7 H2O salt cooled the water while the salt was heating the water. It was interesting that two substances very close to chemical make up could have such a different reaction in the water. MgSO4's graph on water temperature changes seems to be gradually jumping after adding salt. I think this is because my lab partner forgot to turn on the mixer, so the salt wasn't completely mixed at first. Other than that, the procedure worked - 106.6 kJ hydration enthalpy seems pretty high. Water take 4.184 kJ to raise only 1°C, so 106.6 kJ seems like a lot of energy. C. Delve into the chemistry of scientific experiments for chemistry. Not only are these chemical experiments fun, but they are also very educational for everyone interested in scientific chemical reactions and properties. In this video tutorial, we will show you how to determine the enthalpy change in hydration of MgSO4 (magnesium anhydrous). IT Chemistry: Magnesium sulfate is used as an example of how enthalpy changes are determined. Hydrated MgSO4 and Anemia MgSO4 dissolve separately to record temperature changes. DeltaH of Anemia MgSO4 · Hydrated MgSO4 = Delta H of hydrated enthalpy. Check out C's details in WonderHowTo's chemical science video. Music: Blues Corp by Kevin McLeod @IncompetechLicensed under Creative Commons Attribution 3.0 has mastered Microsoft Excel and take work prospects from your job to the next level Want to raise your career with premium A-to-Z Microsoft Excel training bundles from the new gadget hack shop and get lifetime access to over 40 hours of functions, formulas, tools, and more basic advanced instructions. Buy Now (97% Off) &P: Personalize Academics.edu content, customize ads, and use cookies to improve the user experience. By using our site, you use cookies to consent to our collection of information. For more information see the Privacy Policy.X This is a video on how to determine hydration enthalpy, I take magnesium sulfate as an example explaining how it was determined. Hydrated MgSO4 and Anhydrous MgSO4 dissolved respectively and changes are recorded and Delta H of DeltaH-hydrated MgSO4 hydrated enthalpy of Anion MgSO4. Please rate, comment and subscribe! Bruce Corporation by Kevin McLeod @ incompetech.com Licensed under Creative Commons Attribution 3.0 1 Geotechnical Institute of Geotechnology (Mineralogy). You will find articles from Friedrich Schiller University, Jena, Germany 2 Institute of Geology, Mineralogy and Geophysics, University of R le, Bochum, Germany Klaus Dieter Grevel 1 Institute of Geogeology (Mineralogy), Friedrich Schiller University, Jena, Germany. Find articles from Austria Artur Benisek 3 Fachbereich Materialforschung ∓ Fisk, Abteilm Mineralogy, University of Salzburg Salzburg and Austria. Find articles by Edgar Dux 4 Department of the Faculty of Chemistry, University of Hamburg, Germany. A. Dominic Fortes 2: Institute of Geology, Mineralogy and Geophysics, University of R le, Bochum, Germany. Bernd Mahler 1 Institute of Geogeology (Mineralogy), Friedrich Schiller University, Jena, Germany 2 Institute of Geology, Mineralogy and Geophysics, University of R le, Bochum, Germany. Abteilm Mineralogy, University of Salzburg, University of Salzburg, University of Hamburg, University of Hamburg, University of London. Uk corresponding author. Address Correspondence: Dr. Klaus Dieter Grevel, Jena, University of Friedrich Schiller, Institute of Geo-Scientific Research, Faculty of Minerals, Karl Zeiss Promenade 10, D-07745 Yenah, Germany. Email: ed.bur@leiverG-reilo-suaikReceived January 28, 2012; July 7, 2012. Copyright 2012. Mary Ann Liebert, inc., enthalpy in the formation of synthetic MgSO4.4H2O (Star Kate) and MgSO4.3H2O were obtained by solution heat measurement of 7±286.15 K. The formation enthalpy obtained from the element is (Star Kate)=−2498.7±1.1 kJ·mol−1 and (MgSO4.3H2O)=−2210.3±1.3 kJ·mol−1. Star Kate's standard entropy is the temperature range 5 KJt±4t300 K; (Star Kate)=254.48±2.0 J. In addition, differential scanning heat measurement (DSC) measurement using Parkin Elmer diamond DSC with a temperature range of 270KJt±4t300K was performed to confirm the reproducibility of PPMs measurement around the ambient temperature. Star Kate's experimental Cp data between 229 and 303K, fitted with the Meyer-Kelly polynomial, produced Cp(T)=107.925+0.5532. T−1048894. T^2. Hydration status of all Mg sulfate hydrate/s in response to local temperature and humidity conditions. Based on the recently reported equilibrium relative humidity and the above new standard characteristics, the internal consistent thermodynamic database of mgso4·nH2O systems has been refined by mathematical programming (MAP) analysis. As can be seen from the resulting phase diagram, the star kate is unstable throughout the T–P RH range. Due to the motion limits of kiselite formation, star kate's unstable development may be possible under Martian conditions. Keywords: Mg sulfate · star kate · thermodynamic data · entrophy – enthalpy–thermodynamics.astrobiology 12.1042.1054-A Three different multi-shapes are described from laboratory experiments: MgSO4 and nH2O (1≤n11) in many different hydration forms (Table 1). Most of them have been found as minerals on Earth (Peterson, 2011), and recent space missions have supported the presence of significant amounts of Mg sulfuric acid in Martian soils and sediments (e.g., Chiphera and Bannman, 2007; Roach et al. referred to in 2009 and in 0). Recently, enthalpy of formation from elements of kyselite (n=1), sandierite (n=2), hexahydrate (n=6), and epsomite (n=7) (2009) was measured (Grevel and Maglian, 2009). These authors then combined enthalpy data as a function of temperature and relative humidity from other experimental studies on the stability of some Mg-sulfate hydrates, including Epsilite, Hexahydrate, Star Kate (n=4), and kiselite (vart Hoffman et al., 1912; Chou and Seal, 2003, 2007). Obtain an internal and consistent thermodynamic database of systems MgSO4–H2O with rigorous mathematical programming (MAP) analysis. As a result, a phase diagram shown in FIG. As concluded by Grevel and Maglian (2009), star kate is unstable when it comes to kyselite formation, while kyselite formation is known to be very sluggish under typical dehydration (Steiger et al., 2011). Thus, FIG. 1 shows an extended field of star kate's metastatic presence, making this phase a major reaction product of the dehydration reaction. However, star kate's data in the database of Grevel and Maglian (2009) was derived from experiments. Therefore, direct measurement of this stage was strongly desired to define the main purpose of this study. We obtained synthetic starkey by solutoning the amount of heat to water at room temperature. Star Kate's standard entrophy was derived from low temperature thermal capacity measurements obtained with the Physical Property Measurement System (PPMS). Furthermore, enthalpy in the formation of n hydrate (MgSO4.3H2O) was measured. Different hydration forms of MgSO4 and nH2O (0≤n11) as described in the literature are literature step-by-step references: n-MgSO4Differnterperis and Soldnerse, 1958; Fortes et al., 1982; Yamaguchi and Kato, 1972; Fortes and others, 2007; n-MgSO4DiffrtRowe, 1967; Dimon and Kato, 1984; Key Sesserie 1 Hawthorne, 1, 1987;Reagent one hydrate 1 Spera and Bannman, 2007; Gindrod and others, 2010; Steiger and others, 2011;54; hydrogen 1,25van't Hoff and Dawson, 1899; Hodenberg and Kyun, 1967 Sandierite 2Ma and others., 2009a;2.4 Hydrate 2.4 Chiterra and Bannman, 2007;2.5 Hydrate 2.5Ma and others., 2009b;7Hydrate3Hodenberg and K hn, 1967; Fortes and others, 2010;Starkey4tBaur, 1962, 1964a;Cranwickite4tPeterson, 2011; Pentahidrite 5Baur and Lonn, 1972; Hexahidrite 6 Salkin and others, 1964a;Epsomite7Baur, 1964a; Ferrat and others, 1973; Fortes and others, 2006 Meridian At 11 Peterson and the King, 2006; Peterson and others, 2007; Fortes et al., 2008;Steiger et al. (2011) reported new measurements of equilibrium relative humidity to some stable and metastatic hydrated dehydration equilibrium in the MgSO4-H2O system (Fig. 1). These measurements serve as an additional constraint for MAP procedures to obtain a revised internally consistent thermodynamic dataset of magnesium sulfate hydrates. Star Kate MgSO4.6H2O, in a drying cabinet (4 days) at about 320K (Steiger et al., 2011) (Puka 00627 p.a.) produced in the Department of Chemistry (University of Hamburg, Germany) by dehydration. After verification by X-ray powder diffracting (XRD), the powdered substance was stored in a sealed vial in a drying cabinet and then sent to Bochum by regular mail. The XRD pattern was obtained by a transmission sted p powder diffrometer (STOE, Germany) using Cu-K  radiation. The XRD pattern was compared with the JCPDS standard 24-720. Furthermore, powder XRD measurements obtained with Siemens D500 diffracts were scanned every 2s between 3° and 65°2  and collected before and after Cp measurements in Salzburg. MgSO 4 and 3H2O crystals were prepared at 105 °C, without stirring 25% mgso4 solution without stirring (Fortes and Lemme Calliot, 2010). Crystals grown in this way are clusters of euhedral colorless plates 1 to 2 mm long and 0.1 to 0.5 mm thick (Figure 2). After extraction from the solution, the crystals of MgSO4 and 3H2O became somewhat dull. However, batches of crystals that were not used, as reported by Hodenberg and Kyun (1967), were stored in small glass jars for 10 days and endured repeated exposure to warm air (~30 °C) and examination under a microscope. Thus, some crystal fragments can be transported to Jena by email for use in heat measurement experiments. Once powdered under a fine phase hydrates so rapidly that measurements requiring powder samples were not made with this material. The crystal structure of mgso4 and 3D2O in critical conditions were previously measured by neutron single crystal diffraction measurements.Lau Langevin, Grenoble, France (Fortes and Lemme Calliau, 2010). Thermal expansion behavior were measured by high-resolution neutron powder diffing at ISIS neutron spray sources (Fortes et al., 2010). Dimensions of room temperature cells, as=8.1952(2)  , b=10.9210(2)  , c=12.966(4)   and the space group P63 agree with the previously published values of Hodenberg and K hn (1967) for the protonized form (MgSO4.3H2O) used in this work. For the solution caloric value measurement, a commercially available IWC-400 is temperature microcalorimeter (Caloric Measurement Science Co., Ltd.) explained by Ackerman et al. (2009) at the Institute of Earth Science (University of Jena Friedrich-Schiller). A caloric solvent (25 mL of deionized water) contained in a sealed polyethylene ketone cup of 60 mL, was inserted into a liquid bath of a calorimeter and kept at a constant temperature of 25 °C. (±0.0005 °C). During stabilization (8 hours) and experiments (80 to 100 minutes), the solvent was stirred with a SiO2 glass stirrer driven by a motor located about 40 cm from the active zone of the instrument. A fine-grained star keylet sample was pushed into a pellet of about 20 mg and weighed with a microbalance with an accuracy of 0.002 mg (as stated by the manufacturer). On the other, the crystal fragments of the coarse grain size in hydrate sample were weighed and used for the heat quantity measurement experiment without pulverization. Immediately after weighing, the sample was dropped into the solvent through a SiO2 glass tube to measure the heat generated or consumed during dissolution. The solution was always newly prepared, and the final sugar content of Mg2+ was the same in each run. This type of heat measurement does not require consideration of dilution (or concentration) effects. The caloric effect was calculated by integrating the heat flow rate between the reaction cup and the constant temperature reservoir. This thermometer was calibrated by dissolving 20 mg pellets of KCl in 25 g of deionized water. Before each calibration measurement, potassium chloride was heated overnight in a furnace at 500 °C to remove adsorbed water. The expected thermal effect of the calibration run was calculated from Parker (1965). The weight loss of the two star Kate samples (50 mg each) was measured with the STA 503 thermal analyzer (BAHR thermal analysis GmbH) at Rule University Bochum. The sample was examined with N2 dry air (1 atm) in the temperature range from room temperature to 600 °C, at a heating rate of 5 °C / min. Differential thermal analysis (DTA) and thermogravimetry analysis (TG) signals were recorded simultaneously. Star Kate's low temperature (5–300K) thermal capacity behavior was investigated using a quantum-constructed physical property measurement system (PPMS). Established at the University of Salzburg. About 20 mg of the powder sample was sealed in an Al pan and the powder was compressed into a 1 mm thick layer (Dax and Berthold, 2005). The heat capacity was measured at 60 different temperatures and 3 times at each temperature when cooled from 300 K at a number interval. The complete PPMs experiment for measuring Cp consisted of additional runs and sample runs. During the initial measurement, the heat capacity of the empty sample platform was determined. In the second measurement, the sample was placed in a fying pan and the heat capacity of the entire ensemble was measured. The net thermal capacity of the sample is given by the difference between the two measurements. To convert PPMs data from µJ units, a uncertainty of 0.02 mg a sample weight was adopted. K-1 to 3. Differential scanning heat measurement &t measurement using a Parkin Elmer diamond DSC with a temperature range of 270K&t4t300K was performed to confirm the reproducibility of PPMs measurements around ambient temperature (Dax and Benisek, 2011). The powder sample was placed in parkin elmer al pan covered with a lid. Measurements were made under the arg flow and the manometer block was kept at a constant temperature of 243.3 K in the Parkin Elmer Intracooler. The growth of ice crystals on the thermometer block was prevented by dry air flow (200 mL·min−1). In addition, the cover heater was turned off. The full survey consisted of three separate measurements: blanks, references, and sample measurements. Blanks, references, and sample measurements. Before the sample measurement, the DSC was calibrated with a reference exchange using a synthetic single crystal of Mg2+ was the same in each run. 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