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Investigation of Endothermal Doublets Occurring in the Thermal Decomposition of $\text{Ca}(\text{OH})_2$

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ABSTRACT

The presence of at least two different forms of $\text{Ca}(\text{OH})_2$ is confirmed by the results of a thermoanalytical study on the decomposition and re-hydration characteristics of $\text{Ca}(\text{OH})_2$. The study comprised a series of differential scanning calorimetry (DSC) experiments (at temperatures up to 700°C) performed in an atmosphere of a continuous flow of nitrogen pre-conditioned to various relative humidities ranging from 12 to 100%. The emergence of two separate and distinct decomposition endotherms occurred. The endothermic peak temperatures were dependent on the relative humidity and temperature history of the $\text{Ca}(\text{OH})_2$ samples. Exposure of CaO to higher temperatures and different humidities would appear to promote the formation of both crystalline and microcrystalline products. Under these conditions, higher temperatures give rise to formation of very crystalline products and after some time, vapor phase exposure promotes the formation of microcrystalline products. Factors affecting the formation of the different forms of $\text{Ca}(\text{OH})_2$ are described.

Introduction

The emergence of nanoscience applications in cement chemistry is likely to have a significant impact on the understanding of classical reactions that occur in the cement-water system [1]. $\text{Ca}(\text{OH})_2$, (cement chemistry notation, CH) is not only a reaction product, but it is also an important reactant that can participate in many reactions relevant to the durability of cement-based products [2]. Development work involving the use of nano-particulate CH has the potential to tailor the formation of engineered C-S-H microstructures favorable to high performance.

Thermoanalytical methods for determining the extent of cement hydration have been utilized for decades [3]. The decomposition of CH has been the focus of many studies [4]. Generally a single endotherm (in the temperature range, 450-550°C) is detected by thermogravimetry (TG), differential thermal analysis (DTA), and differential scanning calorimetry (DSC) methods. Two thermal events associated with the decomposition of CH have, however, previously been observed. Greene observed these phenomena in studies of the hydration of Portland cement [5]. The low temperature peak was attributed to chemisorbed water on the surface of free lime particles. The typical 2nd peak was due to the decomposition of a more coarsely crystalline CH. Herrick et al. also observed two endothermal peaks in their DTA curves for hydrating expansive clinker pastes [6]. A metastable CH with cubic morphology was said to decompose at the lower temperature. The higher temperature peak was attributed to decomposition of hexagonal CH. The authors have demonstrated that different forms of CH with varying degrees of crystallinity can be produced through vapor and liquid phase hydration of CaO [7]. The vapor phase hydrated CH generally showed a lower thermal decomposition temperature and lower degree of crystallinity than the liquid hydrated CH.

It is apparent that the application of analytical techniques to characterize the nature of reactions in systems containing cement-based nanomaterials may yield new spectral information and data interpretation. This paper focuses on the factors affecting the thermal decomposition of CH of nanoparticle range. Specifically the influence of the dehydration and re-hydration environment on the nature of the CH formed and the decomposition peak temperatures are investigated. Results of experiments on the decomposition of CH in pre-conditioned humidity environments (12-100% RH) are reported.

Thermal Decomposition and Crystallinity of CH

Previous work by the authors has demonstrated that the degree of crystallinity of CH is related to its decomposition temperature [7]. TG was conducted on CH formed by heating CaCO_3 for 1 hour at 1050°C followed by hydration in liquid water for 7 days (referred to as CH-6) or heating reagent grade CH for 1 hour at 600°C followed by vapor phase hydration at 11% RH (referred to as CH-1). Separate decomposition endotherms were observed in a derivative form of TG (DTG) for each sample with the vapor phase hydration product having the lowest endothermic peak temperature. Mixtures of the two samples exhibited doublets in the DTG spectra. The CH that gave the low temperature

endotherm had a significantly lower degree of crystallinity as determined by XRD analysis. Typical DTG curves for the CH mixtures are illustrated in Figure 1.

Experimental

The materials used in the work reported here are as follows:

Materials

CH: The CH was produced by calcining reagent grade CaCO_3 at 1050°C for 3 hours followed by hydration in de-aerated liquid water for 24 hours. This material was used for all the experiments reported in this paper.

CaCO_3 : The reagent grade CaCO_3 was supplied by Fisher Scientific. The BET surface area was $2.6 \text{ m}^2/\text{g}$.

Apparatus

DSC: A Dupont 1090 Thermal Analyzer was used in conjunction with a DSC cell.

In each experiment 4 mg of sample was placed in a gold pan (sample holder).

A heating rate of $10^\circ\text{C}/\text{min}$ with a nitrogen flow of $100 \text{ ml}/\text{min}$ was used. The nitrogen was preconditioned at different values of RH (12-100%). The apparatus used to condition the nitrogen gas for delivery to the DSC cell is shown in Figure 2. The nitrogen is passed over a salt solution or water at a predetermined RH. This is followed by flow into a vessel equipped with an RH sensor to verify the RH of the carrier gas and through a flow meter into the DSC cell. In this work the RH value refers to room temperature equilibrium conditions. As the temperature in the DSC cell increases, the saturation vapor pressure increases as well. The effective RH would therefore be less.

Test Procedures

Series I: CH samples were heated in the DSC cell from room temperature to 600°C at $10^\circ\text{C}/\text{min}$ with a nitrogen flow of $100 \text{ ml}/\text{min}$ preconditioned at different values of RH (12, 38, 60, 80 and 100%). Five continuous heating and cooling (to 100°C) cycles were performed.

Series II: CH samples were subjected to heating/cooling cycles as for Series I with the exception that the samples were held isothermally for different times (1 min to 22 h) at 600°C before cooling. All experiments were performed with the preconditioned environment at 100% RH.

Series III: CH samples were subjected to heating/cooling cycles as for Series II except that the isothermal temperature was 700°C . Isothermal time ranged from 1 to 60 min. All experiments were performed at 100% RH.

Results and Discussion

The results of the DSC experiments generally confirm the presence of two decomposition endotherms for the CH used in this study. The characteristics of the DSC curves are described for the various test conditions as follows:

Series I:

The effect of various RH environments on the decomposition of CH is represented by a single endotherm on the 1st heating as shown in Figure 3. As the RH increases, the CH decomposes at higher temperatures. It is apparent that the CH can decompose more readily in the lower RH environments as the onset of endotherm and peak temperatures are reduced..

The liquid hydrated CH decomposes to CaO, when the temperature increases from room temperature to 600°C. The CaO re-hydrates to form CH, when the temperature decreases from 600°C to 100°C on cooling. The DSC curves for the 4th heating (after 3 cycles of heating and cooling) in various RH environments are typical and are illustrated in Figure 4. The shift of the curves with the RH observed in Figure 3 is also seen in Figure 4. More energy is required to decompose the CH in the higher RH environments. More well-defined doublets also appear in these higher RH environments. This can be explained by examining the 3rd cooling and 4th heating curves for various RH environments shown in Figure 5. The DSC curves for the 38% RH and 100% RH environments are shown. The peaks on cooling and heating indicate the re-hydration and decomposition of the CH, respectively. The CaO starts re-hydrating at 285°C in the 100% RH environments whereas it only starts re-hydrating at 235°C in the 38% RH environment. As mentioned earlier, the RH values in this work refer to room temperature equilibrium conditions. The effective RH in the DSC cell constantly changes while the temperature decreases on cooling, that is, the CaO re-hydrates while the effective RH changes. The range of the effective RH variation with temperature in the 100% RH cell is much wider than the one in the 38% RH cell. The CaO therefore re-hydrates in the wider range of effective RH in the 100% RH cell than the 38% RH cell. This might be the reason for the more distinctive doublets in the 100% RH environment during decomposition in the 4th heating as opposed to those produced in the 38% RH environment. Hydration of a sample produced in different RH environments would result in a difference in degree of crystallinity and therefore difference in a decomposition temperature. It has been shown by the authors that the surface area of CH has no significant influence on the decomposition temperature[7]. In this work it was demonstrated that ΔG , the change in Gibbs Free Energy, for the formation of CH by hydrating CaO is more negative as relative humidity increases (i.e. the thermodynamic driving force for the reaction increases). This is consistent with the variation in decomposition temperature of CH observed in the DTG analysis. Further X-ray evidence (in the form of Williamson-Hall plots) also indicated that the CH formed by hydration of CaO at low relative humidities was less ordered. It is also possible (in practice) that some cements which show dual peaks may have been partly hydrated during exposure to moisture, producing CH of different crystallinity.

Series II:

The effect of an isothermal hold after heating to 600°C followed by cooling to 100°C and reheating to 600°C in a 100% RH environment was investigated. The DSC results are plotted in Figure 6. The DSC curves for the 1st heating are shown to the right of the figure and illustrated by a single endotherm with a peak temperature occurring at about 435°C. To the left of the figure, the DSC curves for the 2nd heating are shown with much lower peak temperatures occurring than the 1st heating. The CH that decomposed on the 1st heating was initially hydrated in the liquid phase whereas the CH that decomposed on the 2nd heating was hydrated in the DSC cell on cooling, that is, the vapor phase. The resulting difference in the decomposition temperature between the CH hydrated in the liquid and vapor phase was observed in the previous work by the authors [7]. The DSC curves for the 2nd heating also indicate the well-defined doublets appearing for the longer isothermal hold. All display two endothermic peaks- low temperature peaks at about 402°C and high temperature peaks at 420°C. The low temperature peaks increase in magnitude with a hold time up to about 5 hours with a slight decrease occurring at 22 hours. The CaO is calcined while holding isothermally at 600°C. Depending on the isothermal holding time, the nature of the CaO as well as the characteristics of the CH on re-hydration and subsequent decomposition are changed.

Series III:

The effect of a higher isothermal temperature (700°C) for a heating-cooling-reheating cycle on the thermal decomposition of CH was determined. The DSC curves for samples subjected to an isothermal hold (after initially heating CH to 700°C) followed by cooling to 100°C and reheating to 700°C in a 100% RH environment are presented in Figure 7. The DSC curves all exhibit two separate endotherms with peak temperatures at 402°C and 420°C. The magnitude of the smaller peak at 402°C decreased as the isothermal hold time at 700°C increased and nearly disappeared at a hold time of 60 min. The magnitude of the larger peak at 420°C decreased only slightly with hold time. It appears that the increased hold time at 700°C favors the formation of a more crystalline form of CH that decomposes at a higher temperature. The microcrystalline form of CH is inhibited from forming possibly due to the decreased reactivity of the CaO at the longer hold times. In this work, the term “microcrystalline” is used to denote the less ordered arrangement of atoms in the crystal structure. This effect is much more pronounced than that observed for Series II specimens where the CaO promoting formation of microcrystalline CH remained reactive for up to 24 hours.

Conclusions

1. Thermograms for the decomposition of CH indicate the presence of two distinct forms (reflected by two endotherms) when the decomposition and re-hydration process occurs in a controlled humidity and CO₂ free environment.
2. The magnitude of both the low and high temperature endothermic peak heights is dependent on the relative humidity and temperature in the DSC test environment.
3. High humidity (in a DSC cell) promotes the hydration and decomposition of larger amounts of CH as indicated by the growth of both DSC endotherms.

4. Higher temperature is conducive to formation of a more crystalline CH product, while at the same time, vapor phase hydration promotes formation of a microcrystalline product.
5. The relative magnitude of the low temperature endotherm (reflecting formation of microcrystalline CH) is sensitive to heating temperatures in excess of 600°C.
6. Previously observed doublets (in the temperature range generally associated with the decomposition of CH) for DTA traces of hydrating cement systems can be explained in terms of the degree of crystallinity of the CH present (rather than the particle size or surface area effects).
7. When a cement or C₃S exhibits dual peaks it is possible the exposure conditions especially humidity cycles will determine the development of dual CH peaks.

Acknowledgment

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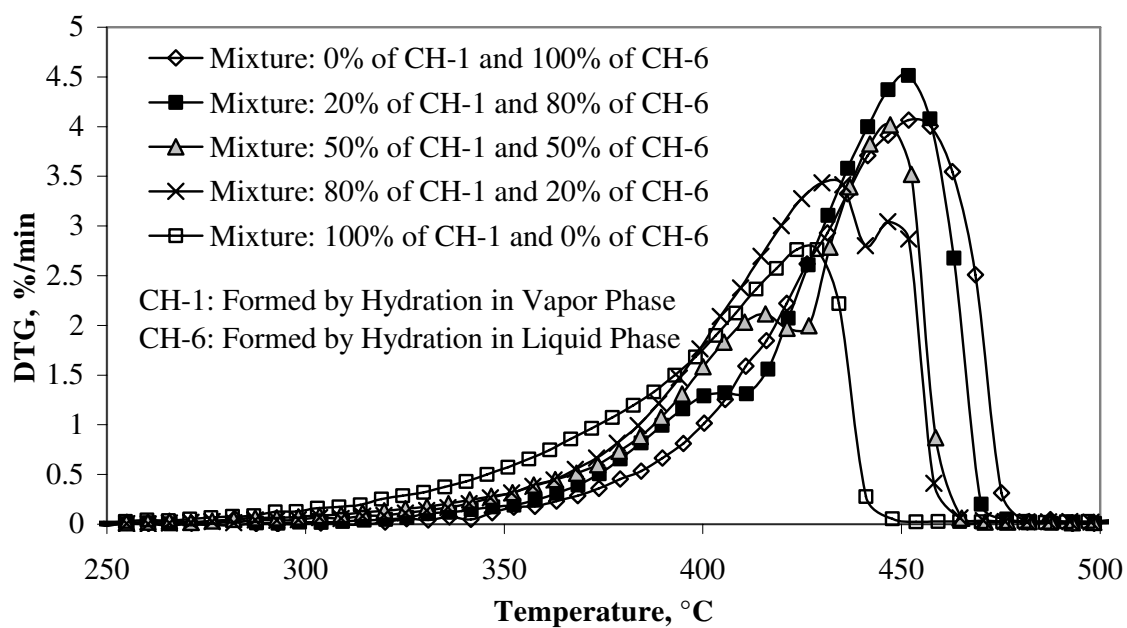


Figure 1 DTG curves for samples CH-1 and CH-6 and their mixtures with different proportions

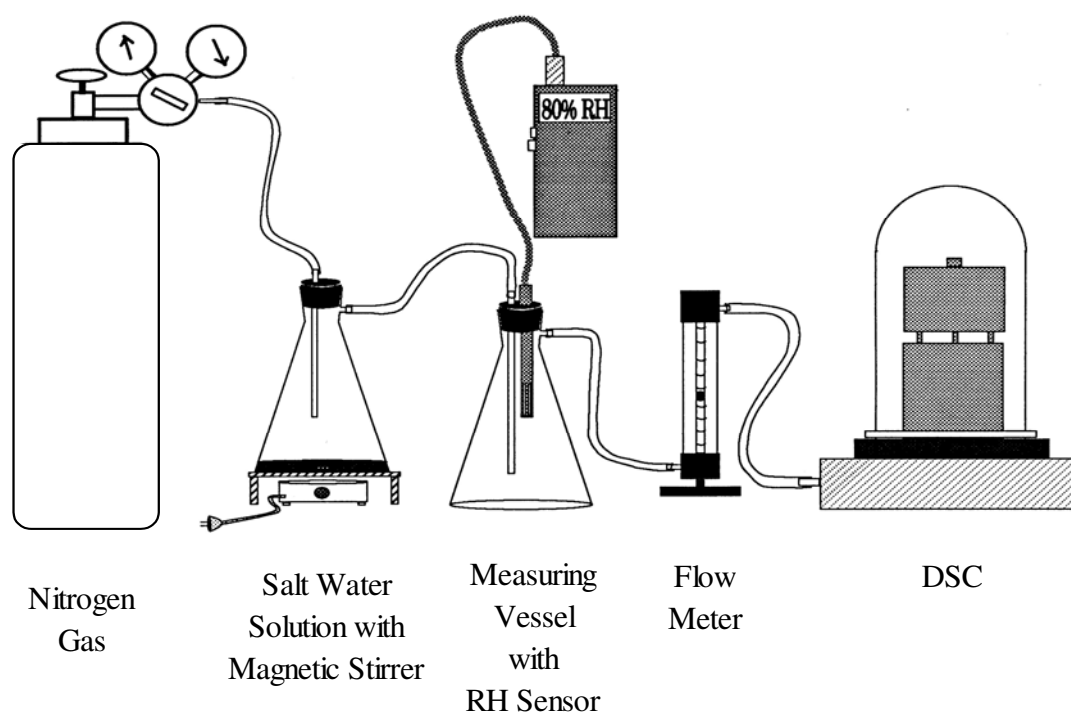


Figure 2 The apparatus to condition the nitrogen gas for delivery to the DSC cell

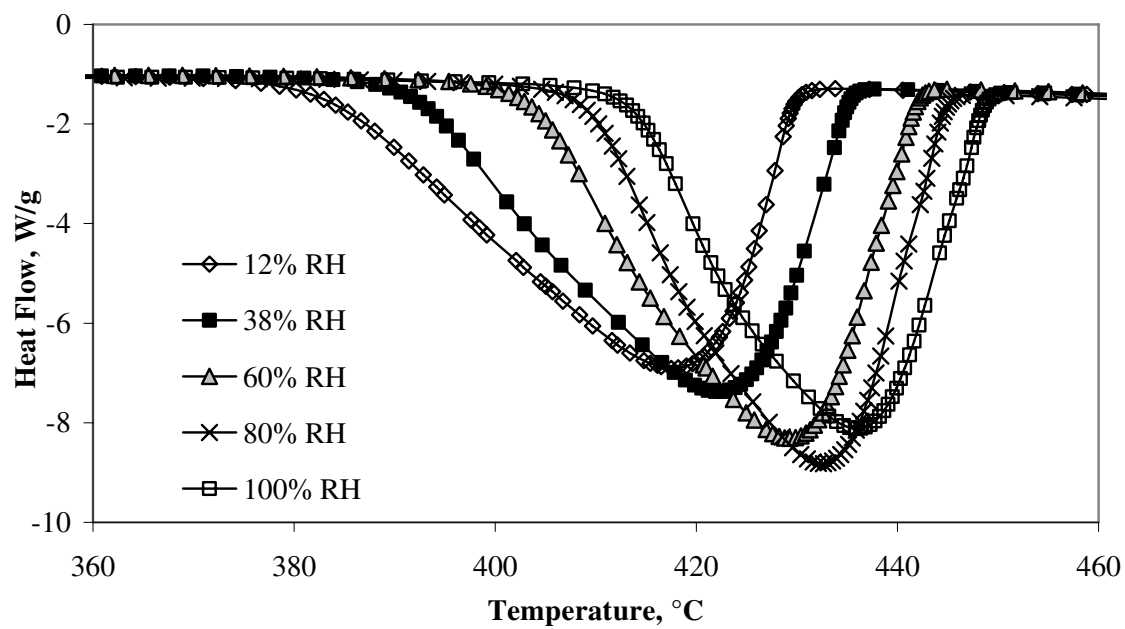


Figure 3 DSC curves for the 1st heating of CH in various RH environments

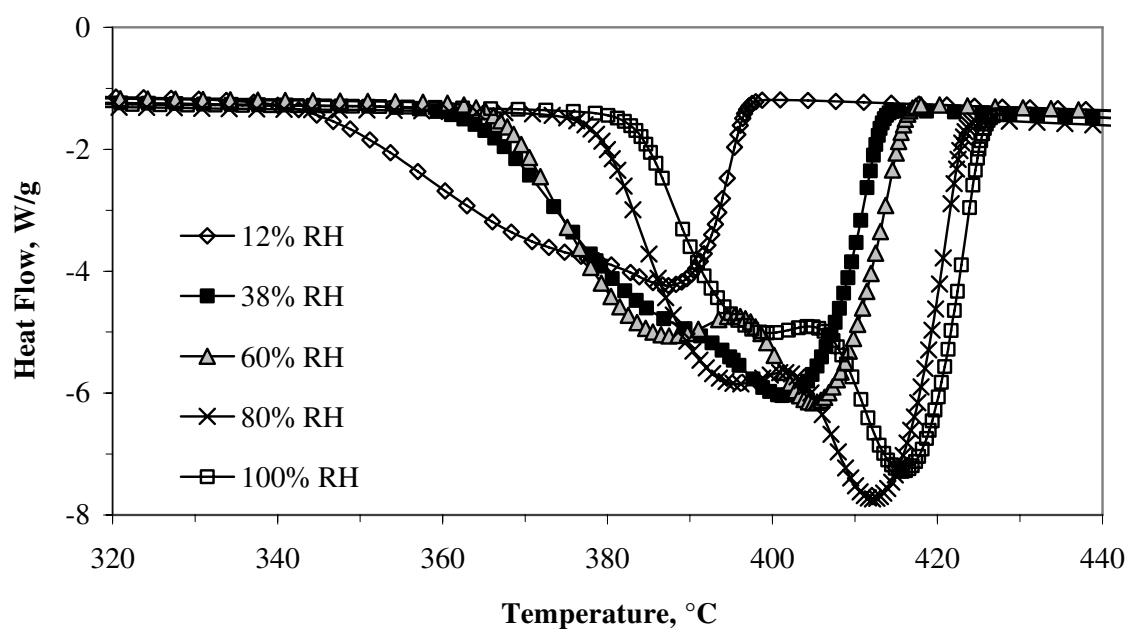


Figure 4 DSC curves for the 4th heating of CH in various RH environments

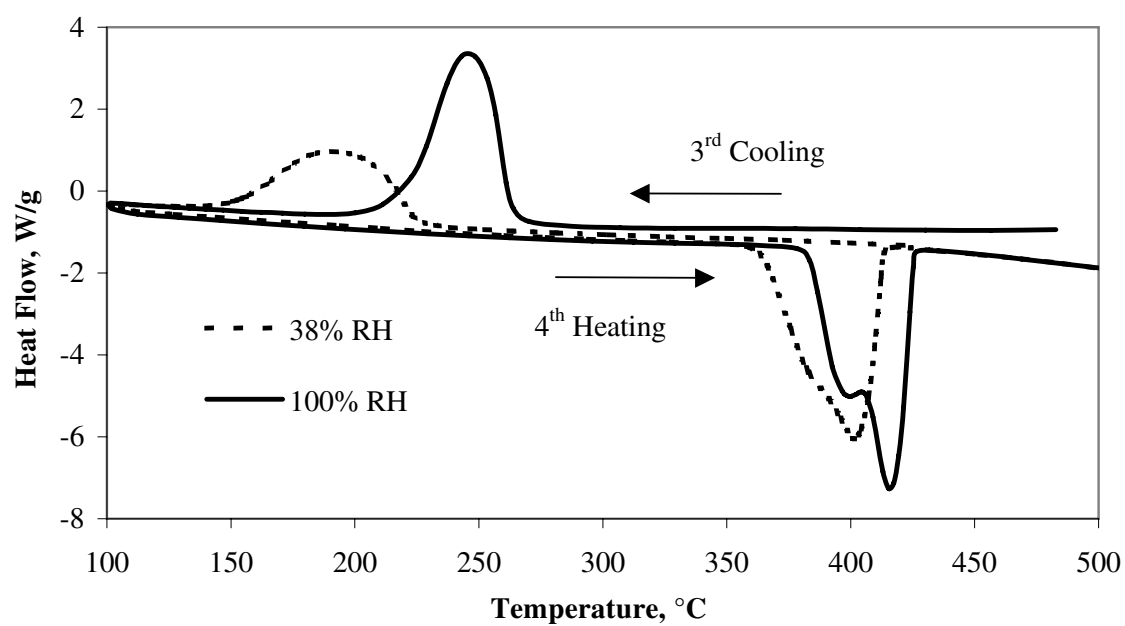


Figure 5 DSC curves for the 3rd cooling of CaO and 4th heating of CH in a 38% and 100% RH environment

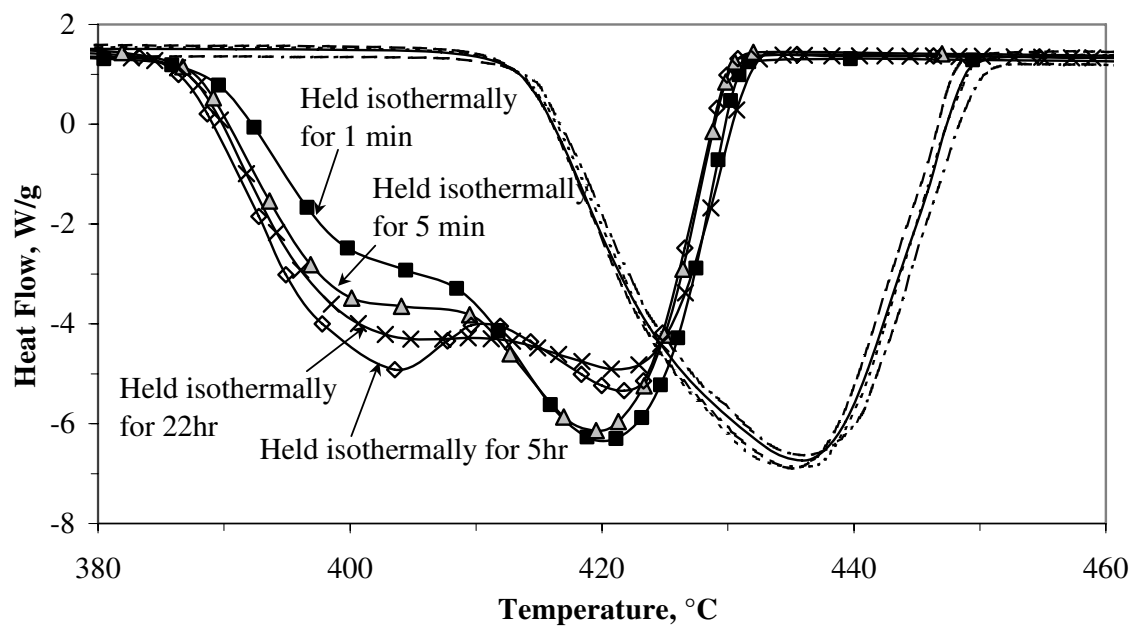


Figure 6 DSC curves for the 1st and 2nd heating of CH in 100% RH held isothermally at 600°C for various times

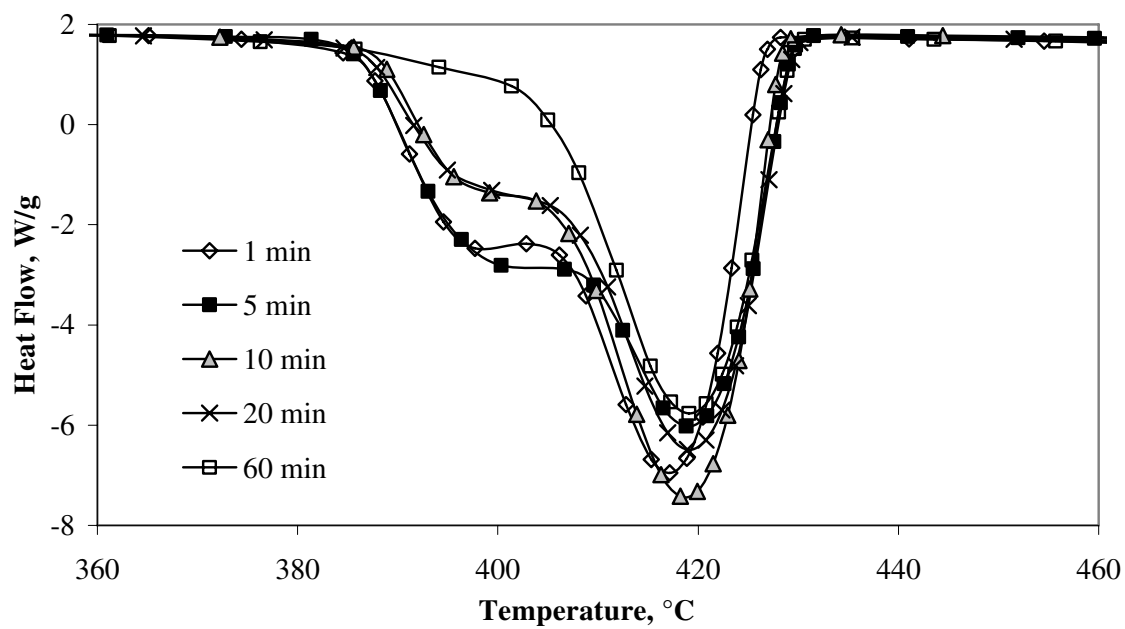


Figure 7 DSC curves for the 2nd heating of CH in 100% RH after holding isothermally at 700°C for various times