Abstract: Huge amounts of natural gas hydrate are trapped in an ice-like structure (hydrate). Most of these hydrates have been formed from biogenic degradation of organic waste in the upper crust and are almost pure methane hydrates. With up to 14 mol% methane, concentrated inside a water phase, this is an attractive energy source. Unlike conventional hydrocarbons, these hydrates are widely distributed around the world, and might in total amount to more than twice the energy in all known sources of conventional fossil fuels. A variety of methods for producing methane from hydrate-filled sediments have been proposed and developed through laboratory scale experiments, pilot scale experiments, and theoretical considerations. Thermal stimulation (steam, hot water) and pressure reduction has by far been the dominating technology platforms during the latest three decades. Thermal stimulation as the primary method is too expensive. There are many challenges related to pressure reduction as a method. Conditions of pressure can be changed to outside the hydrate stability zone, but dissociation energy still needs to be supplied. Pressure release will set up a temperature gradient and heat can be transferred from the surrounding formation, but it has never been proven that the capacity and transport ability will ever be enough to sustain a commercial production rate. On the contrary, some recent pilot tests have been terminated due to freezing down. Other problems include sand production and water production. A more novel approach of injecting CO₂ into natural gas hydrate-filled sediments have also been investigated in various laboratories around the world with varying success. In this work, we focus on some frequent misunderstandings related to this concept. The only feasible mechanism for the use of CO₂ goes though the formation of a new CO₂ hydrate film rapidly forms a mass transport barrier that slows down any further growth of the CO₂ hydrate. Addition of small amounts of surfactants can break these hydrate films. We also demonstrate that the free energy of the CO₂ hydrate is roughly 2 kJ/mol lower than the free energy of the CH₄ hydrate. In addition to heat release from the formation of the new CO₂ hydrate, the increase in ion content of the remaining water will dissociate CH₄ hydrate before the CO₂ hydrate due to the difference in free energy.

Keywords: methane production; hydrate; carbon dioxide storage

1. Introduction

Industrial problems related to hydrate formation in pipelines and process equipment has historically motivated much of the hydrate research. In many of these systems, the heat released during hydrate formation is rapidly distributed through liquid water and solid metal. Related kinetic models have therefore typically focused on the thermodynamic driving force and related mass transport.

Classically, most hydrate risk evaluations have been based on the existence of a free liquid water phase, or the creation of a free water phase through condensation from water dissolved in gas (or liquid). The next step has typically been a consideration of how methanol, or other hydrate...
inhibitors, can shift the hydrate formation region; an example with methanol as inhibitor is illustrated in Figure 1 below. Experimental or theoretical curves like those presented in Figure 1b have been utilized by the oil industry for several decades in order to decide how much methanol they should inject for specific situations of potential hydrate formation risk. Theoretical details and calculation procedures are discussed in Section 2. A typical example is given in Figure 2 below. A typical industrial risk evaluation approach is based on water condensing out as liquid (Figure 2a). During recent years, also the possibility of water dropping out from gas to adsorb on rusty pipeline walls has been investigated and typical calculations are presented in Figure 2b. Typically, the classical approach of calculating maximum water content before condensation will permit in the order of 20 times more water than a corresponding criteria based on water adsorption on rust.

During the latest few decades, the interest in natural gas hydrates as an energy source has increased substantially. Any method utilized to produce methane gas from methane gas hydrates will involve a transfer of heat. Most efforts have been on reducing the pressure to outside hydrate stability, which addresses the thermodynamic driving force, but the heat still needs to be supplied from the surroundings or added in some other way. Some very short pilot tests have been conducted in Alaska [1,2]. In more recent years, two tests were conducted offshore of Japan. The first one lasted for 6 days and was stopped due to problems with production of sand and water, as well as a freezing down due to limited heat supply capacity from the surroundings [3]. The second test was designed and planned for 6 months continuous production but closed down after 24 days due to freezing down problems [4]. Thermal stimulation, through for instance steam or hot water, is also possible but considered far too expensive as the only method. A more novel technology is to inject carbon dioxide. A solid-state mechanism has been proven by Ripmeester et al. [5] and Kuhs et al. [6] for the ice region of water. For temperatures above zero, a substantially faster mechanism [7,8] involves the formation of a new CO$_2$-dominated hydrate inside the pores using free water. Even in permafrost areas it is hard to find any natural gas hydrate deposits with higher hydrate saturations than 85% of the available pore volume. The more typical is 75% or lower like also in the Ignik Sikumi pilot test [9,10]. Injection of CO$_2$ will lead to the formation of a new CO$_2$ hydrate from the free water in the pores and the injected CO$_2$. The use of pure CO$_2$ will rapidly block the pores with CO$_2$ hydrate films and slow down any exchange process with the in situ CH$_4$ hydrate. In the Ignik Sikumi pilot test [9,10], a mixture of 77.5 vol% N$_2$ and the rest CO$_2$ was injected in order to limit and slow down the formation of a new CO$_2$ dominated hydrate. Injecting very diluted CO$_2$ mixtures in N$_2$ will increase permeability for the injected gas but the damping of the new hydrate formation may be far too much for CO$_2$ [10] to even be efficient as a mechanism for CO$_2$/CH$_4$ exchange in the pores. The addition of small amounts of surfactant will keep the interface free of hydrates [11] and stimulate hydrate nucleation and growth below the CO$_2$/water interface region. The effects of surfactant on interface free energy and increased solubility of CO$_2$ below the surfactant stimulated interface are additional effects that will stimulate hydrate growth from solution. Practically it has been observed that small amounts of methanol will act as a promoter for hydrate formation in a pipeline [12].

A brief review of available experimental data for methane hydrate and carbon dioxide hydrate is given in Section 2. Clausius–Clapeyron and Clapeyron have been typical methods for estimating enthalpies of hydrate formation and these methods are briefly discussed in Section 3. In Section 4, we outline our approach based the use of residual thermodynamics for all components in all phases, including hydrate. Example calculations are presented in Section 5, followed by our conclusions.
More precisely, the difference in chemical potential between pure liquid water and chemical potential to hydrate is not straightforward within the more typical ways to calculate hydrate equilibrium today. The Clapeyron method is based on the fact that for the Clapeyron-based methods for the formation of hydrate from a separate gas (or liquid) hydrate the possibility of water dropping out from gas to adsorb on rusty pipeline walls has been investigated.

Figure 1. (a) Hydrate equilibrium curves for CH₄ hydrate with temperature in kelvin and pressure in bar. The solid curve is calculated according to theory and procedures described in Section 2. * are experimental data from Bavoh et al. [13], + are experimental data from Tumba et al. [14] and o are experimental data from Sabil et al. [15]. (b) Hydrate equilibrium curves for hydrate formed from CH₄ and water containing methanol. The solid curve is the calculated equilibrium curve for an amount of 5.44 weight per cent methanol in water. Circles are experimental data from Svartas and Fadnes [16]. The dashed curve is the calculated equilibrium curve for 10 wt% methanol in water and * are experimental data from Ng and Robinson [17]. The dash-dot curve is the calculated equilibrium curve for 20 wt% methanol in water and x are experimental data from Ng & Robinson [17]. + are similar data from Svartás and Fadnes [16] but for 20.01 wt% methanol in water.

Figure 2. (a) Calculated maximum amounts of water in CH₄ before condensing out as liquid water. (b) Maximum amounts of water in CH₄ before adsorption on Hematite. For details on the theory and algorithms for these calculations, as well as other examples, the reader is directed to open literature. Some examples are given in References [10,18–28]. Highest curve is for 50 bars and lowest curve is for 270 bars in both graphs. Other pressures are evenly distributed with 44 bars increasing for each curve from the top 50 bars curves downwards.

2. Residual Thermodynamic Modeling of Hydrate Formation from Water and a Separate Hydrate Former Phase

By definition, a hydrate phase transition is reversible along the equilibrium curve. This is utilized for the Clapeyron-based methods for the formation of hydrate from a separate gas (or liquid) hydrate former phase and a free water phase. More specifically the Clapeyron method is based on the fact that dG = 0 for a phase equilibrium point of pressure and temperature. The application of this simple result to hydrate is not straightforward within the more typical ways to calculate hydrate equilibrium today. In view of Equation (3) below, the empty hydrate is frequently treated as an empirical fitting property. More precisely, the difference in chemical potential between pure liquid water and chemical potential...
in empty clathrate of structure I or II are treated as fitting parameters. Most often, this chemical potential difference at a reference state is fitted as one parameter. An associated enthalpy difference is also fitted, along with fitted differences for specific heat capacities and volume differences so as to be able to adjust chemical potential differences to the pressures and temperatures in consideration. The conditions for hydrate equilibrium are therefore, in this difference method, just a reformulation of Equation (2) below being equal to Equation (3). Empirical fitting of a fundamental property like chemical potential may not be the best way to treat hydrate, but there are also other challenges related to the Clausius approach for hydrate. This is, however, not an important focus in this work. A separate study is dedicated to a more critical review of that method, as well as simplified versions along the lines of Clausius–Clapeyron. These latter simplifications basically imply that condensed phase volumes are neglected. Clausius–Clapeyron models for hydrates are therefore typically only suitable for moderate pressures.

The free energy change for this phase transition can be written as:

$$\Delta G^{(H_1)} = \left[ \sum_i x_i^H \left( \mu_i^H(T,P,x^H) - \mu_{i}^{\text{water}}(T,P,x) \right) + \sum_i x_i^H \left( \mu_i^H(T,P,x^H) - \mu_{i}^{\text{gas}}(T,P,y) \right) \right]$$

(1)

The superscript $H_1$ is used to distinguish the specific heterogeneous phase transition from other hydrate formation phase transitions. $T$ is temperature and $P$ is pressure. $x$ is the mole-fraction in either liquid or hydrate (denoted with a subscript $H$), while $y$ is mole-fraction in the gas (or liquid) hydrate former phase. $i$ is an index for hydrate formers. The superscript water denotes the water phase that is converted into hydrate. Generally, this is ice or liquid, but in this work, we only consider liquid water. $\mu$ is the chemical potential.

The liquid water chemical potential is calculated from the symmetric excess conventions as:

$$\mu_{i}^{\text{water}}(T,P,x) = \mu_{i}^{\text{pure},H_2O}(T,P) + RT \ln \left[ x_{i}^{H_2O} \gamma_{H_2O}(T,P,x^H) \right]$$

(2)

where \( \lim(\gamma_{H_2O}) = 1.0 \) when $x_{H_2O}$ approaches unity.

The chemical potential for water in the hydrate structure is given as [29]:

$$\mu_i^H = \mu_i^{O,H} - \sum_{k=1,2} RT v_k \ln \left( 1 + \sum_i h_{ki} \right)$$

(3)

$$h_{ki} = e^\beta [\mu_k - \Delta s_k]$$

(4)

where $\beta$ is the inverse of the universal gas constant times temperature. At equilibrium, the chemical potential of the guest molecules $i$ in hydrate cavity $k$ is equal to the chemical potential of molecules $i$ in the co-existing phase it comes from. For non-equilibrium, the chemical potential is adjusted for distance from equilibrium through a Taylor expansion as discussed later. The free energies of inclusion (latter term in the exponent) are reported elsewhere [10,18–29]. At thermodynamic equilibrium between a free hydrate former phase, $\mu_{ij}$ is the chemical potential of the guest molecule in the hydrate former phase (gas, liquid, or fluid) at the hydrate equilibrium temperature and pressure.

Hydrate equilibrium is then solved by fixing one of either temperature and pressure and using the fact that the chemical potential of water in hydrate (Equation (3)) and liquid water (Equation (2)) have to be equal. This is very similar to any other approach that utilizes a fugacity time Langmuir constant approach instead of Equation (4). As illustrated by Kvamme & Tanaka [29] a version of Equation (4) for rigid water lattice is the best for small guest molecules. For CH$_4$ the results from Equation (4), and from the more common integration over the Boltzmann factors for a rigid water lattice, the results are almost the same. CO$_2$, on the other hand, is larger. Movements of CO$_2$ in a large cavity of structure I interferes with water liberation movements and the impact is roughly a 1 kJ/mol
destabilization effect. However, the net effect of CO₂ is still a substantially stronger attraction with water than CH₄ in the same cavity type. The advantage of the formulation in Equation (4) is such that it can theoretically correct for large molecules that interfere with water lattice movements, and as such, alters the properties of the water. An example for methane is given in Figure 1a.

For the methanol addition, we have utilized the following model for water activity coefficients:

\[
\gamma_{H₂O} = a₀ + a₁x_{H₂O} + a₂x_{H₂O}² + a₃x_{H₂O}³
\]

where each parameter takes the form (see Table 1 for values):

\[
a_i = a_i,1 + \frac{a_i,2}{T_R} + \frac{a_i,3}{T_R²}
\]

**Table 1. Parameters for Equation (6). Reduced temperature** \(T_R\) is \(T/273.15\).

<table>
<thead>
<tr>
<th>Parameters</th>
<th>(k = 1)</th>
<th>(k = 2)</th>
<th>(k = 3)</th>
</tr>
</thead>
<tbody>
<tr>
<td>(a₀,k)</td>
<td>0.748208</td>
<td>0.520769</td>
<td>-0.599363</td>
</tr>
<tr>
<td>(a₁,k)</td>
<td>0.541737</td>
<td>-0.473879</td>
<td>0.547213</td>
</tr>
<tr>
<td>(a₂,k)</td>
<td>-0.538589</td>
<td>0.567666</td>
<td>-0.525522</td>
</tr>
<tr>
<td>(a₃,k)</td>
<td>0.350678</td>
<td>-0.531175</td>
<td>0.373238</td>
</tr>
</tbody>
</table>

Examples of the changes in chemical potential for water for various mole-fractions of methanol is illustrated in Figure 3a. Note that these values include all changes from pure liquid water, and as such, also contains the ideal missing term in the last term on right hand side of Equation (2). The effect on hydrate equilibrium is visible from the solutions of Equations (2) and (3). The same chemical potential for liquid water and hydrate water results in higher equilibrium pressures for increasing methanol mole-fractions. Some illustrations are given in Figure 1b above. Equilibrium for CH₄ is included through the same hydrate CH₄ chemical potential in Equation (4) and in the separate CH₄ gas phase. The change in hydrate-forming conditions, as well as the change in water enthalpy (see Figure 3b), also shifts the enthalpies of hydrate formation and dissociation.

![Figure 3](image-url)

**Figure 3.** (a) Corrections to liquid water chemical potential as function of temperature, including the ideal mixing term. Upper curve is for 2 mol% methanol, next is for 4 mol% methanol, then 6 mol% methanol, 8 mol% methanol, 10 mol% methanol, and finally 12 mol% methanol for the lowest curve. (b) Corrections to liquid water enthalpy. Order of the curves from top to bottom is as explained in Figure 3a.
The corresponding filling fractions and mole-fractions of methane in the hydrate is given by:

\[ \theta_{ki} = \frac{h_{ki}}{1 + \sum_j h_{kj}} \]  

(7)

\[ \theta_{ki} \] is the filling fraction of component \( i \) in cavity type \( k \). Also:

\[ x^H_i = \frac{\theta_{\text{large}, i} v_{\text{large}, i} + \theta_{\text{small}, i} v_{\text{small}, i}}{1 + \theta_{\text{large}, i} v_{\text{large}, i} + \theta_{\text{small}, i} v_{\text{small}, i}} \]  

(8)

where \( v \) is the fraction of cavity per water for the actual cavity type, as indicated by subscripts. The corresponding mole-fraction water is then given by:

\[ x^H_{\text{H}_2\text{O}} = 1 - \sum_i x^H_i \]  

(9)

and the associated hydrate free energy is then:

\[ G^{(H)} = x^H_{\text{H}_2\text{O}} \mu_{\text{H}_2\text{O}}^H + \sum_i x^H_i \mu_i^H \]  

(10)

in which \( \mu \) denotes the chemical potential. Subscripts \( \text{H}_2\text{O} \) and \( i \) denote water and hydrate formers, respectively. Superscripts \( \text{H} \), water, and gas denote hydrate, liquid water, and gas phases, respectively. \( x \) is the mole-fraction in liquid or hydrate (superscript \( \text{H} \)) and \( y \) is the mole-fraction in hydrate former phase. \( T \) and \( P \) are temperature and pressure, respectively, and \( G \) is the free energy. The \( \Delta \) symbol, the change in free energy, and superscript \( \text{H}_1 \) indicate this hydrate formation route.

The chemical potential for guest molecule \( j \) (in the case of this work either \( \text{CO}_2 \) or \( \text{CH}_4 \)), which enters Equations (4) and (8) at equilibrium is, according to residual thermodynamics:

\[ \mu_j(T, P, y) = \mu^\text{pure,ideal gas}_j(T, P, y) + RT \ln \left[ y_i \phi_j(T, P, y) \right] \]  

(11)

where \( y_i \) is mole-fraction of component \( i \) in the gas mixture. \( \phi_j \) is the fugacity coefficient for \( i \). Ideal gas chemical potential for pure \( i \) can be trivially calculated for any model molecule via statistical mechanics from mass and intramolecular structure (bond lengths and bond angles). Together with density and temperature, the ideal gas chemical potential is available from the momentum space canonical partition function. We have utilized the SRK [30] equation of state for calculating the fugacity coefficient and the density needed for the ideal gas free energy calculations.

Calculated hydrate equilibrium curves for \( \text{CH}_4 \) and \( \text{CO}_2 \) are plotted in Figure 4a. \( \text{CH}_4 \) is supercritical and the corresponding hydrate equilibrium curve is smooth. \( \text{CO}_2 \), on the other hand, goes through a phase transition that changes the density of \( \text{CO}_2 \). This changes the fugacity coefficient for \( \text{CO}_2 \) substantially (see Equation (11)) and results in a steep change in \( \text{CO}_2 \) hydrate equilibrium pressures over the narrow temperature range for the phase transition. It is beyond the scope of this work to discuss the rapid change in the \( \text{CO}_2 \) hydrate equilibrium curve. However, there are some frequent misunderstandings in the open literature on experimental data for \( \text{CO}_2 \) hydrate; see, for instance, Reference [20] for some examples and associated discussion.
The enthalpy change is trivially related to the corresponding free energy change using the thermodynamic relationship:

$$
\Delta H^{Total}_{\text{Total}} = \Delta G^{Total}_{\text{Total}}
$$

The superscript total is introduced to also include the penalty of pushing aside the old phases. Practically, the total free energy change will be Equation (1) plus the interface free energy times the
contact area between the water and hydrate forming phase during the nucleation stage divided by number of molecules in the specific core size. Since critical nuclei sizes are small [23–25], the whole particle can be considered as covered with water due to capillary forces. Above the critical core size, the penalty diminishes rapidly relative to the free energy benefits from Equation (1).

\[
\frac{\partial \left[ \mu_{H_2O}^H \right]}{\partial T}_{P,N} = \frac{\partial \left[ \mu_{H_2O}^H \right]}{\partial T}_{P,N} - \left[ \frac{\partial }{\partial T} \right]_{P,N} \sum_{k=1,2} v_k \ln \left( 1 + \sum_i h_{ki} \right) \tag{13}
\]

For the liquid water phase in Equation (1), as well as for the empty hydrate chemical potential on the right hand side of Equation (3), results are trivially obtained from Kvamme and Tanaka [29], while the second term on the right hand side is reorganized as:

\[
\left[ \frac{\partial }{\partial T} \right]_{P,N} \sum_{k=1,2} v_k \ln \left( 1 + \sum_i h_{ki} \right) = \left[ \sum_{k=1,2} v_k \left( \frac{\partial h_{ki}}{\partial T} \right) \right]_{P,N} \left( 1 + \sum_i h_{ki} \right) \tag{14}
\]

Furthermore, the derivatives of the cavity partition functions can be written as:

\[
\left[ \frac{\partial h_{ki}}{\partial T} \right]_{P,N} = h_{ki} \left[ \frac{-1}{RT^2} \left( \mu_{ki} - \Delta g_{ki} \right) + \frac{1}{RT} \left( \frac{\partial \mu_{ki}}{\partial T} - \frac{\partial \Delta g_{ki}}{\partial T} \right) \right] \tag{15}
\]

The partial derivatives in the last term on right hand side is numerically differentiated from the polynomial fits of Kvamme and Tanaka [29]:

\[
\frac{\partial \left[ \mu_{H_2O}^H \right]}{\partial T}_{P,N} = \frac{\partial \left[ \mu_{H_2O}^H \right]}{\partial T}_{P,N} + \sum_{k=1,2} v_k \left[ \frac{1}{RT^2} \left( \mu_{ki} - \Delta g_{ki} \right) - \frac{1}{RT} \left( \frac{\partial \mu_{ki}}{\partial T} - \frac{\partial \Delta g_{ki}}{\partial T} \right) \right] \left( 1 + \sum_i h_{ki} \right) \tag{16}
\]

\[
H_{H_2O}^H = -RT^2 \left[ \frac{\partial \left[ \mu_{H_2O}^H \right]}{\partial T} \right]_{P,N} + \sum_{k=1,2} v_k \left[ \frac{1}{RT} \sum_i h_{ki} \left( \mu_{ki} - \Delta g_{ki} \right) - T \left( \frac{\partial \mu_{ki}}{\partial T} - \frac{\partial \Delta g_{ki}}{\partial T} \right) \right] \left( 1 + \sum_i h_{ki} \right) \tag{17}
\]

The final term on the right-hand side is also sampled from molecular simulation (MD) sampling along with the free energies of inclusion. The largest distinction in that term is between methane in large and small cavity. For temperatures between 273.15 and 290 K, these are almost straight lines, but a second-order fit of the MD sampled result to the following second order polynomial:

\[
\frac{\partial \Delta g_{ki}}{\partial T} = a_0 + a_1 T_R + a_2 T_R^2 \tag{18}
\]

Equation (18) gives a good fit to the sampled data. Subscript \( R \) on \( T \) on the right-hand side of Equation (18) indicate the reduced temperature. This is defined as \( T \) divided by critical temperature for the components in consideration. Parameters are given in Table 2 below.
Table 2. Parameters for Equation (16). Small cavity in structure I are not stabilized for the actual temperature range and \( h_{ki} = 0 \) for CO\(_2\) in small cavities. The critical temperature for CH\(_4\) is 190.55 K and critical temperature for CO\(_2\) is 304.25 K.

<table>
<thead>
<tr>
<th>Comp. ( i )</th>
<th>Cavity ( k )</th>
<th>( a_0 )</th>
<th>( a_1 )</th>
<th>( a_2 )</th>
</tr>
</thead>
<tbody>
<tr>
<td>CH(_4)</td>
<td>Large</td>
<td>−0.458007</td>
<td>0.266046</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>Small</td>
<td>0.689560</td>
<td>−0.796062</td>
<td>0.234249</td>
</tr>
<tr>
<td>CO(_2)</td>
<td>Large</td>
<td>−0.652590</td>
<td>0.649258</td>
<td>0</td>
</tr>
</tbody>
</table>

For liquid water, the enthalpy is even more trivially obtained using numerical differentiation of the polynomial fit of chemical potential as a function of \( T \) given by Kvamme and Tanaka [29].

In an equilibrium situation, the chemical potential of the same guest in the two cavity types must be the same. Furthermore, these guest chemical potentials must be equal to the chemical potential of the same molecule in the phase that it came from (gas, dissolved in water, adsorbed on mineral surface). For the heterogeneous case, this means the chemical potential of the molecule in gas (or liquid) hydrate former phase. However, outside of equilibrium, the gradients in chemical potentials as a function of \( T \), \( P \), and mole-fractions have to reflect how the molecule behaves in the cavity.

Enthalpies for various guest molecules in the two types of cavities can be evaluated using Monte Carlo simulations along the lines described by Kvamme and Lund [31] and Kvamme and Førreisdahl [32] by sampling guest water interaction energies and efficient volumes from the movements of the guest molecules. The final result needed here:

\[
H_{ki}^R = U_{ki}^R + (z_{ki} - 1)RT
\]  

(19)

where \( U \) is energy and superscript \( R \) denotes the residual (interaction) contribution. \( z_{ki} \) is the compressibility factor for the guest molecule \( i \) in cavity \( k \). Calculations of consistent ideal gas values for the same interaction models that were used in calculation of the residual values is trivial:

\[
z_{ki} = \frac{PV_{ki}}{k_BT}
\]  

(20)

in which \( k_B \) is Boltzmann’s constant and \( V_{ki} \) is the excluded volume of a molecule of type \( i \) in cavity of type \( k \). This latter volume is calculated from the sampled volume of the center of mass movements plus the excluded volume due to water/guest occupation. The sampled values for residual energies and occupation volumes for the various cavities are given in Table 3 below. Slightly more complex sampling and calculation for molecules that are not monoatomic (or approximated as monoatomic like methane) but still fairly standard [31,32] and explicit discussion on this is not needed here.

For a relevant temperature span in the order of 10 K (273–283 K), the differences in residual energies from Monte Carlo sampled data does not vary substantially, and that is why they are listed as constant values for the indicated temperature span. The same values are also used up to 290 K. This is as expected since the hydrate water lattice is fairly rigid and the average movements are almost the same for the limited temperature range. The sampled cavity partition functions will of course vary significantly over the same temperature range due to the direct exponential (Boltzmann factor) dependency. The interaction models for CH\(_4\) and CO\(_2\) utilized are the same as those utilized by Kvamme and Tanaka [29]. An average attraction is also indicated for CO\(_2\) in small cavities. However, the sampled Langmuir constant is very small and not significant. This is also confirmed by the molecular dynamics studies along the lines of Kvamme & Tanaka [29] for which the movements of CO\(_2\) in the small cavity interferes with several water libration frequencies. The resulting free energy of inclusion is not beneficial for CO\(_2\) in the small cavity. Small cavity occupation of CO\(_2\) has been found at extreme conditions in the ice range of temperatures in some studies [29]. However, it remains unclear and unverified whether there would be any significant filling of CO\(_2\) in small cavities for temperatures above zero degrees Celsius.
Table 3. Sample residual energies, $U^R$, and cavity occupation volumes for CH$_4$ and CO$_2$.

<table>
<thead>
<tr>
<th>Property</th>
<th>CH$_4$</th>
<th>CO$_2$</th>
</tr>
</thead>
<tbody>
<tr>
<td>$U^R_k$ (kJ/mole)</td>
<td>Large Cavity</td>
<td>Small Cavity</td>
</tr>
<tr>
<td>$V_{ki}$ (Å$^3$)</td>
<td>164.2</td>
<td>89.2</td>
</tr>
</tbody>
</table>

The energies for CO$_2$ in small cavities was sampled after a structure I containing only CO$_2$ in large cavities was stabilized. Then, small cavities were gradually filled and simulations were run until the average fluctuations in the sampled interaction energies were symmetrical and on average less than 0.5% of the average energy for the hydrate crystal. Practically, this energy value does not have any implication on the enthalpy since the canonical partition function for CO$_2$ in small cavity is practically zero, and as such also the filling fraction of CO$_2$ in small cavity is almost zero. At pressures below 95 bars along the equilibrium curve, it is zero to the third digit in mole-fraction, while the maximum filling fraction in small cavity contributes with 0.006 to the mole-fraction at (290.00 K, 403.0 bars).

In contrast, the calculated mole-fractions of methane in structure I varied between 0.134 at (276.16 K, 25.2 bars) along the equilibrium curve to 0.138 at (290.00 K, 164.7 bars).

The derivative of the chemical potential of a guest molecule i in cavity type k with respect to temperature, as needed in Equation (15), is the negative of partial molar entropy for the same guest molecule and can be calculated according to:

$$\left[\frac{\partial \mu_{ki}}{\partial T}\right]_P = \frac{\mu_{ki} - H_{ki}}{T}$$

Equation (15) can then be rearranged into:

$$H_H^{H_2O} = -RT^2 \frac{\partial}{\partial T} \left[ \frac{\mu_{0,H_2O}^H}{RT} \right]_{P,N} + \left\{ \sum_{k=1,2} v_k \frac{\sum_{i} h_{ki} \left[ (H_{ki} - \Delta g_{ki} + T \frac{\partial \Delta g_{ki}}{\partial T} ) \right]}{1 + \sum_{i} h_{ki}} \right\}$$

The sampled temperature derivatives of guest inclusion free energies for methane in small cavity exhibits very different behavior due to the slightly higher density (roughly 85% higher density as compared to CH$_4$ in large cavities). CO$_2$ in small cavities is not relevant since the canonical partition functions are practically zero within significance.

Residual enthalpies for a hydrate former in a separate hydrate former phase are trivially given using:

$$H^R_i = -RT^2 \sum_{i} y_i \left[ \frac{\partial \ln \phi_{i,gas}^R}{\partial T} \right]_{P,y_i}$$

in which the same equation of state (SRK) is utilized as the one used for calculating fugacity coefficients for the chemical potentials. As examples, we plot the calculated equilibrium curves for CH$_4$ and CO$_2$ in Figure 2a and the corresponding calculated heats of hydrate formation along the equilibrium curves in Figure 2b. The calculated results for CH$_4$ in Figure 1a is compared to experimental data in Kvamme et al. [11], and comparison between calculated results and experimental data for the CO$_2$ equilibrium curve can be found in Kvamme [10].

Experimental data on enthalpies of hydrate formation (or dissociation) are frequently missing various pieces of information that is important for quantitative comparison. The hydrate number is often missing or set to a fixed (guessed) number. Pressure is sometimes missing and temperature is not always given. Therefore, the degree of superheating relative to the hydrate stability limit is not always available. It is beyond the scope of this work to conduct a detailed review of the various data we have.
used for comparison. Readers are directed to the original sources listed in the caption to Figure 6. Work is in progress on a more detailed review of experimental data for the heats of dissociation of hydrates.

Figure 6. (a) Estimated heats of formation based residual thermodynamics for CH$_4$ (solid) as a function of methanol mole-fraction in water. Lowest curve is for pure water and highest curve is for 12 mol% methanol. Second curve from bottom is for 2 mol% methanol, then 4 mol% methanol, 6 mol% methanol, and 8 mol% methanol. Some results from open sources for pure water are also plotted: data from Nakamura et al. [31] (x), Deaton and Frost [32] (*), Kang [33] (o), and Lievois [34] (+). (b) Estimated heats of formation based residual thermodynamics for CO$_2$ (solid). Same notations as in Figure 6a as a function of mol% methanol in water. One available value from Kang [33] for pure water is also plotted using o.

Outside of equilibrium, the only phase that needs special attention is the hydrate phase since description of all fluids phase are continuous while the theory for hydrates is based on a Langmuir type of equilibrium adsorption theory:

$$ G_{\text{Non-equilibrium}}^{H}(T,P,x) = \left[ G^{H,\text{Eq}}(T, P_{\text{Eq}}, x_{\text{Eq}}) + \sum_{r} \frac{\partial G^{H}}{\partial x_{r}} \bigg|_{P,T,x_{\text{Eq}}} (x_{r} - x_{r}^{\text{Eq}}) + \frac{\partial G^{H}}{\partial P} \bigg|_{T,x_{\text{Eq}}} (P - P_{\text{Eq}}) + \frac{\partial G^{H}}{\partial T} \bigg|_{P,x_{\text{Eq}}} (T - T_{\text{Eq}}) \right] \tag{24} $$

in which the equilibrium properties, as a reference state, is available from the calculations leading to Figure 1a. The third term on the right-hand side is a trivial volume term and the final temperature term on the right-hand side can be avoided by expanding in a constant temperature from the equilibrium curve. The second term involves an iteration in which a new set of filling fractions are calculated from the properties at the new pressure. The correction is very limited and no further iteration is needed. The corresponding chemical potentials needed for Equation (8) in the non-equilibrium situation is then trivially given using:

$$ \mu_{m}^{H}(T,P,x) = \left[ \frac{\partial \left( N G_{\text{Non-equilibrium}}^{H}(T, P, x) \right)}{\partial N_{m}} \right]_{T,P,N_{i\neq m}} \tag{25} $$

This then, in turn, can be used for the calculation of the phase transition enthalpies outside of hydrate equilibrium.

As discussed by Kvamme et al. [11], small amounts of methanol will concentrate on the interface between liquid water and a non-polar (or slightly polar) hydrate former phase. This will keep the interface free of hydrate. Heterogeneous hydrate nucleation is therefore only examined for methanol
concentrations up to 2 mol%. For this purpose, we utilize the classical nucleation theory (CNT) because it is simple enough to also be implemented in industrial tools, as well as in reservoir modelling.

The primary reason for conducting these calculations is that we want to illustrate the common misunderstanding that CO₂ injection is not a feasible way to produce hydrate because it is slow. As discussed by Kvamme et al. [11], CH₄ hydrate nucleation is very fast and a nanoscale process in time and volume. In terms of physics, this is not surprising, and it falls well into nucleation times for ice and other phases of complexity similar to hydrates. Since CNT is described in more detail in many other places, including our paper on CH₄ hydrate nucleation [11], only a very brief presentation is given here so as to explain symbols.

Kinetic models for hydrates are implicit in terms of mass transport, heat transport, and thermodynamic control of the phase transition. This is true even for the simplest theory of all of the classical theory (CNT).

The mass transport fluxes need various kinetic theories, e.g., Multicomponent Diffuse Interface Theory (MDIT) theory [33–35], which reduces to the classical theory for a multi-component system when the interface thickness in MDIT theory goes to zero. For an illustration of the coupled transport and thermodynamic control of the phase transition kinetics, CNT serves as an easy method here because of the separation of contributions:

\[ J = J_0 e^{-\beta \Delta G_{\text{Total}}} \]  \hspace{1cm} (26)

where \( J_0 \) is the mass transport flux supplying the hydrate growth. For the phase transition in Equation (1), it will be the supply of CO₂ across an interface of gradually more structured water towards the hydrate core, as discussed in Kvamme et al. [11]. In Equation (10), it will be the diffusion rate for dissolved CO₂ to crystal growth from aqueous solution. The units of \( J_0 \) will be mol/m²s for heterogeneous hydrate formation on the growing surface area of the hydrate crystal. \( \beta \) is the inverse of the gas constant times temperature and \( \Delta G_{\text{Total}} \) is the molar free energy change of the phase transition. This molar free energy consists of two contributions: (1) the phase transition free energy as described by Equation (1), and (2) the penalty work of pushing aside old phases. Since the molar densities of liquid water and hydrate are reasonably close, it is a fair approximation to multiply the molar free energy of the phase transition with molar density of hydrate times the volume of the hydrate core. The push-work penalty term is simply the interface free energy times the surface area of the hydrate crystal. Using lines below symbols to indicate extensive properties (unit: J):

\[ \Delta G_{\text{Total}} = \Delta G_{\text{Phasetransition}} + \Delta G_{\text{Pushwork}} \]  \hspace{1cm} (27)

For the simplest possible geometry of a crystal, which is a sphere with radius \( R \), we then get:

\[ \Delta G_{\text{Total}} = \frac{4}{3} \pi R^3 \rho_{N}^{H} \Delta G_{\text{Phasetransition}} + 4 \pi R^2 \gamma \]  \hspace{1cm} (28)

where \( \rho_{N}^{H} \) is the molar density of the hydrate and \( \gamma \) is the interface free energy between the hydrate and surrounding phase. Even if a hydrate core that grows on the surface of water is floating, it is expected that small crystals are likely to also be covered by water towards the gas side due to capillary forces that will facilitate transport and adsorption of water molecules from the liquid water side to also cover the hydrate particle on the gas side.

Differentiation of Equation (25) with respect to \( R \) and solving for the maximum free energy radius (the critical core size) gives the usual result:

\[ R^* = \frac{2 \gamma}{\rho_{N}^{H} \Delta G_{\text{Phasetransition}}} \]  \hspace{1cm} (29)
in which the superscript * denotes the critical nuclei radius. Critical radius for three different temperatures and four concentrations of methanol in water is given in Figure 7 below. For the two temperatures below the transition over to high density CO$_2$, the trend is very similar to what was observed from phase field theory (PFT) modelling [27,36]. The exception is for hydrate forming from liquid CO$_2$ as seen from the dash-dot curves in Figure 7 by the large critical radii, even for pure liquid water.

Figure 7. (a) Calculated critical radius for pure water and three different temperatures: 273.16 K (solid), 278.16 K (dash), and 283.16 K (dash-dot). (b) Same as (a) but now for 1 mol% methanol in water. (c) Same as (a) but now for 2 mol% methanol in water. (d) Same as (a) but for 3 mol% methanol in water.

The change in interface concentrations of CO$_2$ as function of distance from the liquid side ($z = 0$) of the interface to hydrate side ($z = 12$) has been sampled using molecular dynamics simulations [36,37] and fitted to a mathematical model for the profile as given by Equation (30) below:

$$C(R) = \sum_{i=1}^{7} a_i \left[ \arctan \left( 0.6 \left( \frac{z}{12} \right) \left( \frac{\pi}{2} \right) \right) \right]^{(i-1)}$$

$z \in (0,12)$ with coefficient is Table 4 below.

Table 4. Coefficients for the fitted model of interface concentrations of CO$_2$ from Equation (30).

<table>
<thead>
<tr>
<th>Coefficients</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>$a_1$</td>
<td>$3.649712 \times 10^{-4}$</td>
</tr>
<tr>
<td>$a_2$</td>
<td>$9.895427 \times 10^{-4}$</td>
</tr>
<tr>
<td>$a_3$</td>
<td>$1.091582 \times 10^{-2}$</td>
</tr>
<tr>
<td>$a_4$</td>
<td>$-1.112258 \times 10^{-1}$</td>
</tr>
<tr>
<td>$a_5$</td>
<td>$4.381832 \times 10^{-1}$</td>
</tr>
<tr>
<td>$a_6$</td>
<td>$-6.201765 \times 10^{-1}$</td>
</tr>
<tr>
<td>$a_7$</td>
<td>$2.896391 \times 10^{-1}$</td>
</tr>
</tbody>
</table>
Fick’s law for the mass transport part related to $J_0$ in Equation (23) can be expressed as:

$$\frac{\partial C(z)}{\partial t} = -D_{CO_2} \frac{\partial^2 C(z)}{\partial z^2}$$  (31)

where $C$ is concentration of $CO_2$, $t$ is time, and $D_{CO_2}$ is the diffusivity of $CO_2$ through the interface.

An empirical and simple interpolation of diffusivities from liquid side to hydrate side is utilized just to illustrate the order of magnitude of nucleation times:

$$-\log_{10} \left[ \frac{D(R)}{D_{\text{liquid},j}} \right] = \sum_{i=1}^{9} a_i \left( a \tan \left[ \left( 0.5 R \pi / 2 \right) \right] \right)^{i-1}$$  (32)

where $j = CO_2$ or $CH_4$.

This equation is an interpolation of some sampled molecular dynamics (MD) data from a hydrate block during dissociation. The samplings are not rigorous enough to serve as any real theoretical result. Diffusivities across an interface between liquid water and hydrate is complex and hard to calculate using MD. Equation (28) is therefore to be considered as empirical but with some qualified guessing based on molecular dynamics observations. Work is in progress on the development of other methods to estimate these diffusivities. We consider it as good enough for illustrating the order of magnitude of nucleation times. $R$ is the distance from the liquid side of the interface. $D_{\text{liquid},j}$ is the diffusivity of the guest molecule ($CO_2$ or $CH_4$) in the liquid outside the liquid side interface boundary. Even though $CO_2$ is the focus of this work, $CH_4$ is also listed since we did not find substantial differences in the water structure in the interface between hydrate of these two types of guest molecules. The parameters in Equation (28) are given in Table 5 below.

\[
\begin{align*}
& t(R) - t(0) = \int_{C(R=0)}^{C(R=12)} \frac{\partial C(z)}{\partial t} \left[ -D_{CO_2} \frac{\partial^2 C(z)}{\partial z^2} \right] \\
\end{align*}
\]  (33)

**Table 5. Parameters for Equation (29).**

<table>
<thead>
<tr>
<th>$i$</th>
<th>Parameter</th>
<th>$i$</th>
<th>Parameter</th>
<th>$i$</th>
<th>Parameter</th>
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<td>6</td>
<td>1939.55</td>
<td>9</td>
<td>-7496.15</td>
</tr>
</tbody>
</table>

Diffusivities of $CO_2$ through hydrate vary between different sources in the open literature. All values are based on modelling, mainly using Monte Carlo approaches. It is beyond the scope of this work to screen all available studies and corresponding reported values. Reported values vary between $10^{-15}$ m$^2$/s to $10^{-17}$ m$^2$/s. We simply choose $10^{-15}$ m$^2$/s for $D_{H_4}$ and five conservative values for the liquid side of the interface; $10^{-9}$ m$^2$/s, $10^{-10}$ m$^2$/s, $10^{-11}$ m$^2$/s, $10^{-12}$ m$^2$/s, and $10^{-13}$ m$^2$/s just to illustrate a fairly conservative choice of mass transport characteristics across the interface. The lowest of these values for liquid side diffusivity coefficient gives a transport time for $CO_2$ across the interface of 51 ns, while the second smallest diffusivity coefficient results in a transport time of 5.1 ns. Based on the structure and density of the $CO_2$ on the liquid side of the profile in Figure 8b, these small diffusivities are unrealistically small and a liquid side diffusivity of $10^{-9}$ m$^2$/s is used to calculate nucleation associated with critical nuclei sizes in Figure 7. Results for pure water and three different concentrations of methanol in water are given in Figure 9.
Figure 8. (a) Relative diffusivities for CO₂ across the interface between water and the hydrate. D is the interface diffusivity and \( D_{\text{liquid}} \) is the diffusivity at the liquid side of the interface. (b) The concentration profile for CO₂ as a function of distance from the liquid side of the interface.

Figure 9. (a) Calculated nucleation times in nanoseconds using the concentration profiles as mathematically fitted using Equation (26) in Equation (27), together with Equation (28), for changes in diffusivity across the liquid water/hydrate interface. The applied diffusivity coefficient on the liquid side is \( 10^{-9} \) m². The three different temperatures are 273.16 K (solid), 278.16 K (dash), and 283.16 K (dash-dot). (b) Same as (a) but now for 1 mol% methanol in water. (c) Same as (a) but now for 2 mol% methanol in water. (d) Same as (a) but for 3 mol% methanol in water.

As mentioned, the interpolation in Equation (28) is empirical, but any type of model that contains some very slow diffusivity rates close to a hydrate (in accordance with the structure development of water) would still result in nanoscale nucleation times. The calculations behind the plots in Figure 9 are very conservative because of the very small value for the liquid-side diffusivity coefficient applied in Equation (28). Nucleation times may therefore very well be substantially faster than the curves indicate.
The difference between 273.16 K and 278.16 K is not very significant, with asymptotic values for 166 ns and 172 ns, respectively, for 400 bars pressure, while the corresponding value is 485 ns for 283.16 K.

When nucleation times of several hours are reported, it is normally the first visible (human eye or microscope) hydrate. The development of this hydrate will be substantially delayed by slow transport through the hydrate film. We have also seen this when we utilized magnetic resonance imaging (MRI) [38] to monitor the development of CH\textsubscript{4} hydrate formation in a compartment of plastic walls. The onset of massive growth that results in visible hydrate is given in terms of induction times. Several mechanisms are involved in the development towards induction time. The rearrangement of hydrates in lack of new hydrate building blocks, capillary transport towards solid walls, and heat release during formation of new hydrate are some of the factors.

Some estimates of hydrate film growth kinetics using a constant diffusivity coefficient though the hydrate film is given as Equation (33), and two different choices for liquid diffusivity coefficients in Equation (33). Liquid diffusivity coefficient in equation equal to (34). Liquid diffusivity coefficient in equation equal to (a) $10^{-7}$ m\textsuperscript{2}/s, and (b) $5 \times 10^{-8}$ m\textsuperscript{2}/s.

3. Residual Thermodynamic Modeling of Hydrate Formation from Water and Dissolved Hydrate Former in Water

$$\Delta G^{(H_2)} = \left[ x^H_{H_2O} \left( \mu^H_{H_2O}(T, P, x^H_{H_2O}) - \mu^\text{water}(T, P, x) \right) + \sum_j x^H_j \left( \mu^H_j(T, P, x^H_j) - \mu^\text{water}(T, P, x) \right) \right]$$

(34)

where the superscript H\textsubscript{2} now refers to homogeneous hydrate formation from liquid water and dissolved hydrate former in water, in accordance with a similar notation in Kvamme et al. [11]. The chemical potential for dissolved methane in water can then [11] be formulated as:

$$\mu^\text{water}(T, P, x) = \mu^\text{\textsuperscript{\infty,Residual}}_{CH_4}(T, P, x) + \mu_{CH_4}^{\text{\textsuperscript{\infty,Residual}}}(T, P, x) + RT \ln \left( x_{CH_4} \gamma_{CH_4}(T, P, x) \right)$$

(35)

with

$$\mu^\text{\textsuperscript{\infty,Residual}}_{CH_4} = 3.665 + \frac{40.667}{T_R} - \frac{48.860}{T_R^2}$$

(36)

where $T_R$ is temperature divided by the critical temperature for methane. The maximum temperature used in the fitting is 325 K. Ideal gas as a function of temperature and density is trivial to consistently calculate using the TIP4P model moments in inertia for the rotational contribution [29].
The activity coefficient for methane in water, based on the asymmetric excess convention (activity coefficient for CH\textsubscript{4} in water unity as mole-fraction CH\textsubscript{4} goes to zero) has been fitted to the following function:

\[
\ln \gamma_{\text{CH}_4}^\infty(T, P, x_{\text{CH}_4}) = \sum_{i=1,2}^{39} \left[ a_0(i) + \frac{a_1(i+1)}{T_R} \right] (x_{\text{CH}_4})^{0.05 + \frac{i - 1}{40}} \tag{37}
\]

where \(T_R\) is the reduced temperature and defined as actual \(T\) in Kelvin divided by the critical temperature for methane (190.6 K). The lower summation 1,2 indicates starting from 1 and counting in steps of 2. Parameters are given in Kvamme et al. [11] (Table 3 in that paper).

For CO\textsubscript{2}, a slightly different approach is utilized. The density of CO\textsubscript{2} as dissolved in water will correspond to the partial molar volume of CO\textsubscript{2} at infinite dilution. The infinite dilution ideal gas chemical potential is not very sensitive to pressure, so the following approximation to only temperature dependency is considered as adequate:

\[
\mu_{\text{CO}_2}^\infty, \text{ideal gas} = -130.006 + \frac{163.818}{T_{0,R}} - \frac{64.898}{T_{0,R}^2} \tag{38}
\]

where \(T_{0,R}\) is 273.15 K divided by the actual temperature. Equation (34) does not apply to temperatures above 303 K due to the limited range of temperatures for which infinite partial molar volumes are used, and for temperatures above 273.15 K.

The fugacity coefficient for CO\textsubscript{2} in water is fitted using the following function:

\[
\ln \phi_{\text{CO}_2}^\text{water}(T, P, x_{\text{CO}_2}) = \sum_{i=1,2}^{39} \left[ a_0(i) + \frac{a_1(i+1)}{T_R} \right] (x_{\text{CO}_2})^{0.05 + \frac{i - 1}{40}} \tag{39}
\]

where \(T_R\) is reduced temperature and defined as actual \(T\) in Kelvin divided by critical temperature for CO\textsubscript{2} (304.35 K). The lower summation 1, 2 indicates starting from 1 and counting in steps of 2. Parameters are given in Table 6 below. The vector sign on mole-fraction \(x\) denote the vector of mole-fractions \(i\).

### Table 6. Parameters for Equation (28).

<table>
<thead>
<tr>
<th>(i)</th>
<th>(a_0)</th>
<th>(a_1)</th>
<th>(i)</th>
<th>(a_0)</th>
<th>(a_1)</th>
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<td>89.048877</td>
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<td>35.597488</td>
<td>38.125674</td>
</tr>
</tbody>
</table>

The chemical potential for CO\textsubscript{2} that applies to Equations (4) and (8) for an equilibrium case is then given as:

\[
\mu_{\text{CO}_2}(T, P, y_{\text{CO}_2}) = \mu_{\text{CO}_2}^\infty, \text{ideal gas}(T, P, y_{\text{CO}_2}) + RT \ln \left[ x_{\text{CO}_2} \phi_{\text{CO}_2}(T, P, y_{\text{CO}_2}) \right] \tag{40}
\]

Since the chemical potential of CO\textsubscript{2} is not necessarily the same for dissolved CO\textsubscript{2} in water and CO\textsubscript{2} in gas (or liquid) in a non-equilibrium situation, then hydrate formed according to Equation (2) will be different from the first hydrate and accordingly denoted H\textsubscript{2}. The composition of this hydrate will be different as seen from the corresponding compositions, which follows from Equations (4)–(8).
For homogeneous hydrate formation from liquid water and a dissolved hydrate former, the number of degrees of freedom (Gibbs phase rule) is 2. Each specific set of temperature and pressure will have a maximum dissolved content of hydrate former according a Henry’s law type of calculation [11]. These values are trivially found through iterative solutions of chemical potential for CO₂ in water solution being equal to CO₂ in a separate phase. In this work, we have used the SRK [30] equation of state for calculating gas (or liquid) CO₂ fugacity coefficients.

For the same set of temperature and pressure, there is a unique minimum content of hydrate former in the liquid water that is needed to keep the hydrate stable. These limits are found from equal chemical potentials for water in liquid and hydrate when chemical potential for CO₂ from Equation (36) is used in Equation (4), as given by Equation (37) below. Examples for CH₄ are given in Figure 11 below for CH₄ for a larger range of temperatures and pressures than those reported by Kvamme et al. [11]. Similar results for CO₂ are given in Figure 12.

![Figures 11 and 12](https://example.com/figures)

**Figure 11.** (a) Calculated solubility of CH₄ in water. (b) Calculated minimum limits of mole-fraction CH₄ in water to keep the hydrate stable.

**Figure 12.** (a) Calculated solubility of CO₂ in water. (b) Calculated minimum limits of mole-fraction CO₂ in water to keep the hydrate stable.

Note the strange frame that is drawn between the corners of Figures 11b and 12b, which is an artificial bug in the graphical software. This is of course not a physical part of the calculated data, and must accordingly be neglected in the reading of the curves. In Figure 12b, there is as such a fairly steep change in liquid mole-fraction for the lower limit of hydrate stability. For methane, which does not have a sharp density increase in the same range, the contours in Figure 11b are slightly concave, and again, the outer strange “side-walls” are not physical.
The only change in Equation (3) for this case is in the cavity partition functions such that Equation (4), now with more specific notations, can be written at equilibrium as:

$$ h_{ki} = \frac{\beta_j^{\mu_{\text{water}}}(T,P) - \Delta G_0(T)}{\beta} $$

where the chemical potentials are either from Equation (35) for CO$_2$. For the case of equilibrium, it is assumed that chemical potential for the guest molecule has same chemical potential in both cavities. Outside of equilibrium, Equation (39) applies and the Taylor expansion will also imply that the chemical potentials of guest molecules in the two types of cavities are not necessarily the same.

A second set of changes, relative to the case of separate phases for water and hydrate former, lies in the enthalpy for the hydrate former that enters in the changes associated with the phase transition:

$$ H_{CO_2}^R = -RT^2 \left[ \frac{\partial \ln \phi_{\text{water}}}{\partial T} \right] P_{\beta \neq CO_2} $$

and:

$$ H_{CO_2}^\text{water} = -RT^2 \left[ \frac{\partial \ln \phi_{\text{CO}_2}}{\partial T} \right] P_{\beta \neq CO_2} + \left[ \frac{\partial \ln \phi_{\text{water}}}{\partial T} \right] P_{\beta \neq CO_2} $$

An example for nucleation of hydrate from solution is given in Figure 13 below. As expected, the critical nuclei radius decreases substantially when the concentration of CO$_2$ in the water gets closer to the solubility curve. Furthermore, very close to the hydrate stability limit concentrations, the critical radius approaches impossible sizes. The same is reflected in the corresponding nucleation times.

![Figure 13](image-url)  
Figure 13. (a) Critical nuclei size for CO$_2$ hydrate forming from dissolved CO$_2$ in water as a function of CO$_2$ concentration in water at 273.16 K and 100 bar. (b) Calculated nucleation times as a function of CO$_2$ concentration in water.

The heat of formation from liquid solution is smaller than the values for formation from gas and liquid since changes in enthalpy for the guest molecules is smaller. Some examples are plotted in Figure 14b below. Practically, these are interesting since the most likely hydrate formation from solution is towards the existing hydrate film formed from water and a separate CO$_2$ phase. The released heat will then distribute between dissipation into the water below or heating and potentially dissociating the hydrate locally. The dash-dot curves for 283.16 K in Figure 14a,b may be hard to see from the way the graphical software presented them. In Figure 14a, this is the short curve highest up, and in Figure 14b, it is the short curve on the bottom.
4. Discussion

Two very common misunderstandings are related to Figure 4. The first of these is related to the shape of the equilibrium curve. It is not our intention to criticize experimental groups here but as discussed by Kvamme and Aromada [20], the transition over to higher density for CO₂ also results in a sharp transition of equilibrium pressures. This transition is frequently smoothed out when experimental data are published. Another misunderstanding is that the pressure and temperature projection of the stability region is discussed as CH₄ being more stable than CH₄ hydrate after this region of change in density for CO₂ because the equilibrium pressures for CH₄ then become lower, as seen from Figure 4a. Stability is a multidimensional function of all independent thermodynamic variables like pressure, temperature, and concentrations of all components in all phases. As such, the temperature and pressure projection only tells part of the story. Figure 7 tells other parts of the story. Furthermore, the free energy of CO₂ hydrate is lower than the free energy of CH₄ hydrate for all pressures and temperatures. These are comparable free energies since all components and all phases are based on residual thermodynamic descriptions, and as such, are consistent.

The concept of using CO₂ for the combined production of CH₄ and long-term safe storage of CO₂ as hydrate has been misunderstood in many experimental reports. We have used very conservative transport properties in kinetic modeling of CO₂ hydrate formation from separate CO₂ phase and from solution. Furthermore, even with low diffusion rates for the supply of mass to nucleation and growth, the nucleation times hardly exceed some few hundred nanoseconds. Transport through these hydrate films, in order to sustain further growth, is extremely slow, and even to grow 1 millimeter may take several hours. Even though these estimates are oversimplified, there is no doubt that CO₂ diffusivity coefficients through hydrate in the order of $10^{-15}$ m²/s to $10^{-17}$ m²/s creates substantial delays in the growth up to visible hydrate film range. Addition of small amounts of surface-active components can solve these problems as discussed by Kvamme et al. [11] in a recent paper.

Figure 14. (a) Residual enthalpy for CO₂ in liquid water at 100 bar and three temperatures: 273.16 K (solid black), 278.16 K (dash), and 283.16 K (dash-dot). The solid blue line is the residual enthalpy for CO₂ in a large cavity of structure I and not significantly sensitive to temperature for this limited temperature range. (b) Enthalpies of formation of CO₂ hydrate from liquid solution in water at 100 bar and three different temperatures: 273.16 K (solid), 278.16 K (dash), and 283.16 K (dash-dot).
5. Conclusions

Many experimental groups around the world have been conducting experiments on the exchange on CH\textsubscript{4} hydrate with CO\textsubscript{2}. Frequent conclusions end up with the process being far too slow. Some end up adding nitrogen or other gases, which often leads to even slower conversion because the thermodynamic driving forces for the critical parts of the mechanism is reduced. There are two verified mechanisms for the conversion but one of them, the solid-state conversion mechanism, has only been proven in the ice region far below 0 °C. The other mechanism, which has been proven by state-of-the-art theoretical approaches involves the creation of a new hydrate from the injected CO\textsubscript{2}. In this work, we have demonstrated a new approach for calculating enthalpies of hydrate formation and have shown that the released heat of formation of CO\textsubscript{2} hydrate may be 10 kJ/mole of hydrate former higher than similar number for CH\textsubscript{4} hydrate. We have also demonstrated that the free energy of CO\textsubscript{2} hydrate is in the order of 2 kJ/mole hydrate lower for CO\textsubscript{2} hydrate as compared to CH\textsubscript{4} hydrate. This latter piece of information is important since the formation of a new CO\textsubscript{2} hydrate will lead to an increase in ion activity for the remaining liquid water. The lower free energy for the CO\textsubscript{2} hydrate implies that CH\textsubscript{4} hydrate is more affected by the increased ion activity in terms of destabilization. The problem with the CO\textsubscript{2} injection method is that hydrate films form very fast. In this work, we have utilized classical nucleation theory (CNT) with very slow transport characteristics so as to be conservative. Despite this, it is hard to get nucleation times that exceed some few hundred nanoseconds. Furthermore, transport of mass through these hydrates is extremely slow. There are many sources of data that publish diffusivities for CH\textsubscript{4} or CO\textsubscript{2} but reported values vary a lot between $10^{-15}$ m\textsuperscript{2}/s and $10^{-17}$ m\textsuperscript{2}/s. With values like this and oversimplified estimates, it is easy to verify that even creating a 1 mm hydrate film of CO\textsubscript{2} can take hundreds of hours. This is also in accordance with previous results from our group that reported visible (on a 300-micron resolution scale experiment) hydrate from CH\textsubscript{4} and water after 100 h. Based on our earlier results on adding small amounts of alcohols or surfactants, we proposed the addition of these types of chemicals rather than addition of nitrogen or other gases that makes it less likely to generate new hydrate. The key lesson here is to keep the CO\textsubscript{2}/water interface free of hydrate films.

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References


