

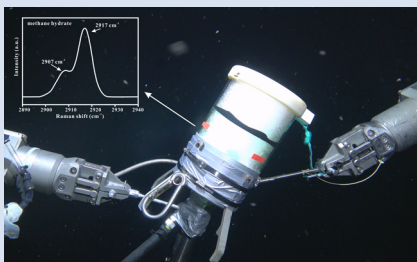
The direct observation and interpretation of gas hydrate decomposition with ocean depth

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Abstract



An exposed gas hydrate (EGH) evolutionary model as a function of water depth was established by *in situ* EGHs ascent experiments in cold seep areas of Haima (1509 m, 2.88 °C and 15.22 MPa), Lingshui (1760 m, 2.56 °C, 17.76 MPa and pH = 7.97) and Site F (1100 m, 3.57 °C, 11.09 MPa and pH = 7.69) in the South China Sea. A remotely operated vehicle was used to reproduce the *in situ* EGH ascent. Changes and temperature variations during EGH ascent were monitored in real time using a Raman insertion probe and dissolved oxygen sensor. The EGH ascent involved three stages of change: i) the metastable stage where no morphology changes, but where gas escapes and there is a decrease in internal temperature; ii) a second stage of coexistence of peripheral hydrate decomposition and internal hydrate growth; iii) a third stage of internal hydrate decomposition at shallower depths. Experimental results indicated that EGHs can carry gas bubbles to shallow depths and even to the sea surface. This could be an important transport mode for cold seep gases affecting shallow waters or the atmosphere.

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Introduction

Exposed gas hydrates (EGHs) have been frequently reported in shallow surface sediments, and are common beneath authigenic carbonate rocks and empty mussel shells in active cold seeps (Fig. 1) (Sassen *et al.*, 2004; Pohlman *et al.*, 2005; Hester *et al.*, 2007). EGHs are formed by the accumulation of rising methane bubbles, which are rapidly covered with hydrate film. These EGHs contain a large amount of free methane gas and have a soft and loose structure, representing the initial state of hydrate formation (Zhang *et al.*, 2017a; Du *et al.*, 2018). Disturbances such as submarine turbidity currents, processes related to climate change, and geological activity can cause the ascent of EGHs due to destabilisation (Rehder *et al.*, 2009). Rising decomposition may occur when EGHs are destabilised, but thermodynamic conditions in the deep ocean environment have the potential to convert the gas back to hydrates (Zheng *et al.*, 2020). The complex marine environment makes it difficult to predict the fate of EGHs in the natural state after destabilisation occurs and they escape into seawater. Previous research has shown that methane fluid from cold seep vents can be dissolved into the water column or consumed by methanotrophic microorganisms (Thornton *et al.*, 2016; Egger *et al.*, 2018). It is unknown whether EGHs destabilised into seawater can improve the survival ability of methane gas, and the depths to which EGHs can carry methane gas after a destabilised ascent remain uncertain.

The investigation of the hydrate evolution after their initial formation has been conducted *via* laboratory simulations in recent decades (Zhong *et al.*, 2016; Lei *et al.*, 2019). The morphology, thickening pattern, and growth resistance of hydrate film have been explored in detail (Zeng *et al.*, 2019; Qureshi *et al.*, 2022a; Dhamu *et al.*, 2023). Research conducted on hydrates in laboratories has typically been limited to steady-state systems at specific temperature and pressure conditions. Laboratory techniques are unavailable for use in real and complex ocean environments (Brewer *et al.*, 1998; Warzinski *et al.*, 2014). Compared to freshwater environments in the laboratory, EGHs in the marine environment are more unstable due to factors such as salinity and currents (Qureshi *et al.*, 2022b, 2022c). With the development of underwater vehicle technology and the increase in our understanding of cold seep regions, *in situ* technologies that can more accurately analyse natural gas hydrates have gradually matured. Initially, Brewer *et al.* (1998) made detailed recordings of the formation and growth patterns of gas hydrates as a function of depth, using a remotely operated vehicle (ROV). Hester *et al.* (2009) concluded that mass-transfer is the rate-controlling mechanism for the dissolution of EGHs and recorded the *in situ* decomposition of EGHs located in the Barkley Canyon area, off Vancouver Island, Canada. Recently, Du *et al.* (2018) characterised the structural features and evolution process of EGHs using Raman spectroscopy in the cold seep area of the South China Sea. A series of investigations of hydrate formation processes and their natural evolution have been performed through *in situ* experiments in recent decades. One similarity

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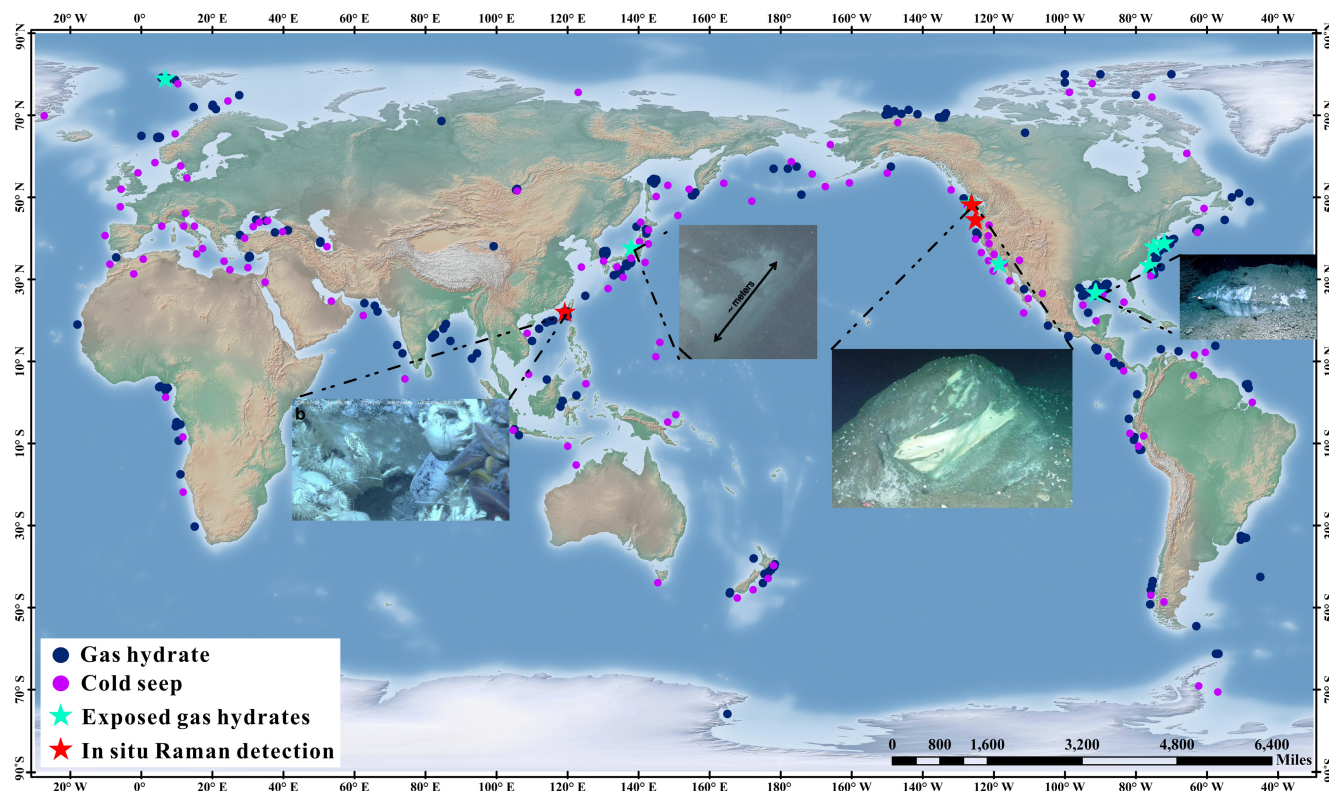


Figure 1 Global map of gas hydrates (Kvenvolden, 1988; Waite *et al.*, 2020) and cold seeps (Mazurenko and Soloviev, 2003; Suess, 2014). Cyan stars represent the locations of exposed gas hydrate occurrence (MacDonald *et al.*, 2003; Sassen *et al.*, 2004; Pohlman *et al.*, 2005; Roberts *et al.*, 2006; Hester *et al.*, 2007; Case *et al.*, 2017). Red stars indicate locations where *in situ* Raman investigations of exposed hydrates have been performed (Hester *et al.*, 2007; Zhang *et al.*, 2017a).

among all these *in situ* experiments was that they were conducted in stationary environments, and there were no records obtained of EGHs as they rose to the surface. Thus, the evolution of EGHs during their rise is not yet clear.

In this study, we performed *in situ* experiments on the ascent decomposition of EGHs in the Haima, Lingshui, and Site F cold seep areas in the South China Sea (SCS) (Fig. 2a). The EGH samples were formed directly using gas-rich fluids that erupted from cold seep vents (Video S-1). We monitored the morphological changes that occurred during the EGH ascent using a camera mounted on the remotely operated vehicle (ROV) “Faxian” (Fig. 2b–g). A Raman insertion probe for gas hydrates (RiP-Gh), previously developed by Zhang *et al.* (2017b), and a dissolved oxygen sensor (DOS, JFE RINKO I ARO-USB) were used to capture the kinetic and thermodynamic behaviour of EGHs during their ascent (analytical methods are provided in detail in the Supplementary Information). Here, we report data from the *in situ* ascent of EGHs in a natural environment and analyse the corresponding impact on the upper water layers.

In Situ Raman Spectra of Exposed Gas Hydrate Samples

The Raman spectral data indicated that the gaseous component of the cold seep fluids consisted primarily of methane gas (Fig. S-1). When conducting the *in situ* experiments using cold seep vent fluids in the seafloor, we found that the formation of hydrates occurred nearly instantaneously (Video S-1). Methane gas bubbles were rapidly wrapped with a hydrate film during their ascent, which provided separation between the internal gas and external water (Fig. 2h). Multiple bubbles carrying hydrate films

accumulated in cells to form a honeycomb structure in the EGH sample (Video S-1, Fig. 2h₁). After a stationary period, an RiP-Gh probe was inserted into the EGH sample at approximately 20 cm for *in situ* Raman spectral observation (Videos S-2 to S-4). Notably, the time that hydrates remained on the seafloor after formation varied in each *in situ* experiment. Hydrate samples remained on the seafloor for half an hour at the Haima cold seep, 2.6 hours at the Lingshui cold seep and 48 hours at the Site F cold seep (Table S-2). The different stationary periods of hydrates make our experiments more generalisable.

The Raman spectra of EGHs indicated two apparent Raman peaks occurring at 2907 cm^{-1} and 2917 cm^{-1} . Both Raman spectra and the guest molecule species indicated that the EGHs in our *in situ* experiments were all type I structures. Notably, the peak intensity at 2907 cm^{-1} was less than that at 2917 cm^{-1} (Fig. S-2). The Raman peak located at 2917 cm^{-1} consisted of a composite peak that was influenced by both a small cage vibration and the gaseous CH_4 vibration.

Stage Changes in the Rising Process of Exposed Gas Hydrates

In the Haima *in situ* experiments, the entire monitoring time of the rising EGHs was 0.79 h. There were three significant stage changes during the rising of the EGHs towards the surface (Video S-2). The first stage occurred at pressures of 15.22–7.05 MPa (1509–700 m). There were no significant changes in the morphology of the EGHs during this stage (Fig. S-3). However, the 2917 cm^{-1} composite peak intensity exhibited a decreasing trend, which was associated with the free CH_4 gas escaping as the hydrate migrated towards the surface

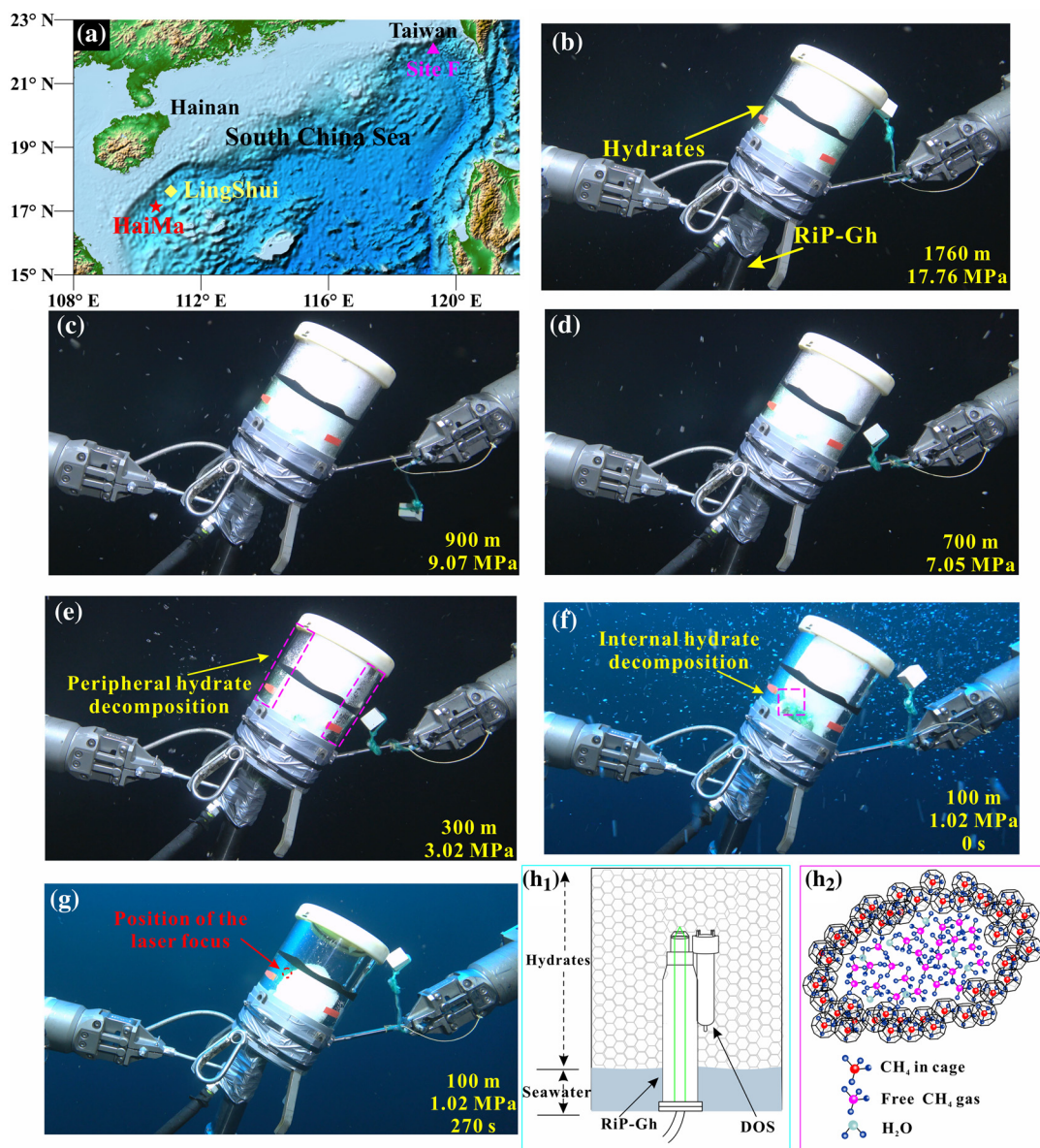


Figure 2 (a) Map showing the locations of the Haima, Lingshui and Site F cold seep vents. (b–g) Images showing hydrate collection and monitoring controlled by the ROV manipulator in Lingshui *in situ* experiment. (h₁) Schematic diagram of the RiP-Gh and DOS probes touching and being inserted into the EGHs for dynamic monitoring. (h₂) Schematic diagram of the hydrate film formed in the gas bubbles.

(Fig. 3b). In normal circumstances, the further growth of the hydrate can only occur *via* the diffusion of water molecules through the hydrate film, which occurs slowly (Dhamu *et al.*, 2023). However, during the ROV ascent, the hydrate film can rupture due to the decrease in pressure exerted by the seawater. Consequently, the gas inside the EGH samples will escape, leading to an accelerated exchange with water molecules and other surrounding media. In contrast, the peak intensity that occurred at 2907 cm⁻¹ showed a weaker enhancement trend (Fig. 3b). Simultaneously, the large cage vibration after Gaussian fitting also shows an enhanced trend (Fig. 4a). This indicates the growth of the EGH samples at this stage. By comparing the hydrate stability curves, we found that the internal environment of the EGHs was consistent with the stable existence of hydrates (Fig. 4b).

The second stage occurs at pressures of 7.05–2.59 MPa (700–257 m) during the ascent. At this stage, the composite peak at 2917 cm⁻¹ also shows a decreasing trend (Fig. 3b), indicating that the gas escapes. Raman spectral data suggest that the gas

escapes throughout the experiment (Fig. 3a). The same process was also observed in the *in situ* experiments conducted at Lingshui and Site F (Figs. S-5 and S-6). The hydrate samples started decomposing from the periphery boundary at a pressure of approximately 7.05 MPa (700 m), when the ambient temperature of the seawater was 6.9 °C (Fig. 4a). The hydrate stability curve (Dickens and Quinby-Hunt, 1994) shows that the theoretical depth at which the decomposition of the EGHs begins is at the pressure of 6.19 MPa (614 m) (Fig. 4b). Here, the depth where decomposition of the EGH sample occurred was advanced during the ascent. Noticeably, although the EGH sample started to decompose from the periphery boundary, there was also a continued increase in the Raman intensity of the large cage inside the EGHs (Fig. 4a). This suggests that there exists a continued growth inside the EGHs. Moreover, the temperature and pressure conditions inside the EGHs at this stage were consistent with the stable existence of hydrates (Fig. 4b). We monitored the continued hydrate growth process up to a pressure of 2.59 MPa (257 m) (Fig. 4a).

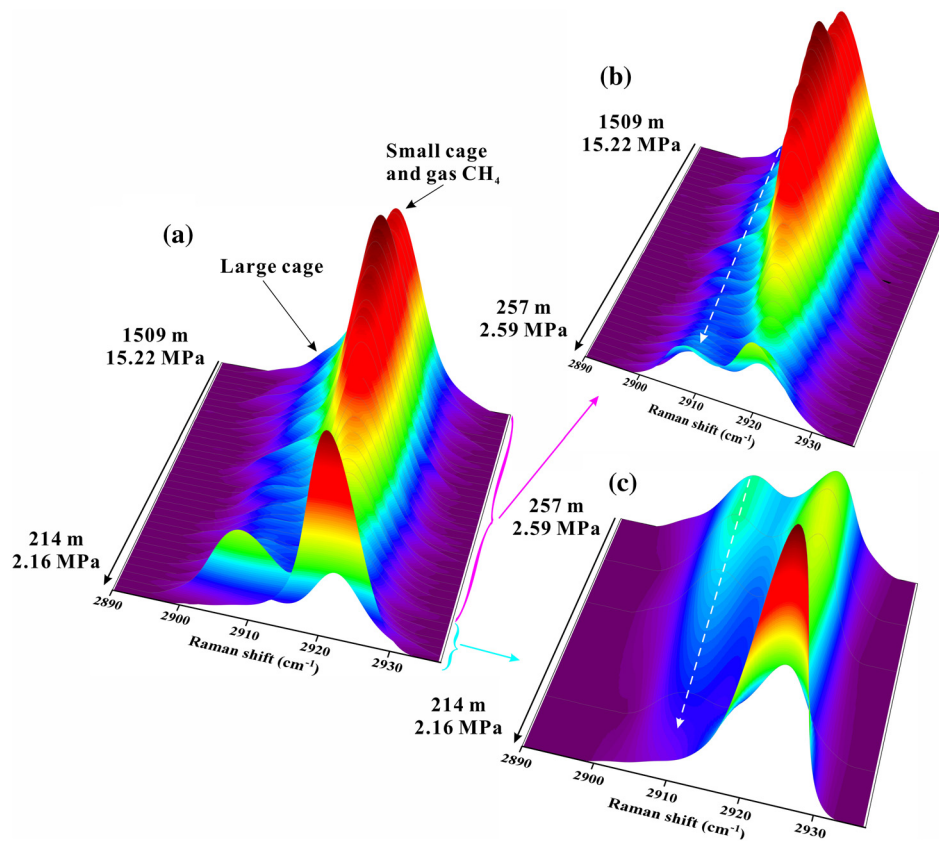


Figure 3 *In situ* Raman spectra for EGHS during their ascent towards the surface in the Haima cold seep. (a) Overall ascent over the pressure range of 15.22–2.16 MPa (1509–214 m). (b) The continuous growth stage at a Raman intensity of 2907 cm^{-1} , and the continuous decrease at 2917 cm^{-1} . (c) Hydrate decomposition stage.

The third stage occurs at pressures of 2.59–2.16 MPa (257–214 m) during the ascent. At this stage, the peak intensity at 2907 cm^{-1} and the large cage vibration (2905 cm^{-1}) abruptly decreased, indicating that the inside of the EGHS sample started to decompose (Figs. 3c and 4a). At a pressure of 2.16 MPa (214 m), the Raman signal of the hydrate disappeared, and only gaseous CH_4 remained (Fig. 3c). One interesting point we observed is that internal hydrate decomposition started at 2.59 MPa (257 m) during this stage. However, the hydrate equilibrium curve shows that the temperature and pressure inside the EGHS did not satisfy the conditions for the stable existence of EGHS as early as 3.34 MPa (332 m) (Fig. 4b). This indicates that the peripheral hydrate decomposition causes a hysteresis effect on the internal hydrate decomposition. Notably, at a pressure of 2.16 MPa (214 m), the complete decomposition of EGHS was recorded by the RiP-Gh probe. The residue of the EGHS, however, was still present above the sample cell.

The above data and discussions are based on the experiment conducted at the Haima cold seep vent. The *in situ* experiments conducted at Site F (Video S-4) and the Lingshui cold seep (Video S-3) vent locations also showed similar results to those at the Haima cold seep vent. The monitoring time for the rising process of EGHS in the Site F cold seep area was 0.61 h, while in the Lingshui cold seep area the monitoring time was 1.34 h. There were also three distinct stage changes during the EGH ascent (Figs. S-3 to S-7). During the *in situ* ascent experiment performed at Site F, the first stage occurred at pressures of 11.09–7.05 MPa (1100–700 m), the second stage occurred at pressures of 7.05–0.65 MPa (700–65 m), and the third stage occurred at pressures of 0.65–0.28 MPa (65–27 m) (Fig. S-7b). The first stage of the *in situ* experiment conducted at the

Lingshui cold seep vent occurred at pressures of 17.76–7.05 MPa (1760–700 m) during the ascent process, and the second stage occurred at pressures of 7.05–1.01 MPa (700–100 m) (Fig. S-7a). The ROV did not continue to ascend in the third stage during the *in situ* experiment performed at the Lingshui cold seep vent. The ROV instead remained stationary at a water depth of 100 m (1.01 MPa), and we monitored the hydrate decomposition process of the third stage over time (Fig. S-7a). The third stage of the *in situ* experiment for the Lingshui cold seep vent lasted approximately 270 s (Fig. S-7a).

Specific Temperature Variations Inside Hydrates

The temperature of the EGHS showed a distinct trend from that of seawater. This appeared inconsistent with both the commonly believed exothermic and absorbed nature of hydrate formation and decomposition. In the first stage, the internal temperature of the hydrate samples dropped by 2.7 °C when the ambient temperature–pressure conditions fit the zone of hydrate stability (Fig. 4b). The ambient temperature of the seawater increased by 4 °C. At this stage, the hydrate periphery had not yet begun to dissipate, thus protecting the internal temperature of the gas hydrate from the influence of the ambient temperature of the seawater. The decrease in internal temperature was attributed to the dissolution of EGH samples. Various factors, such as the high methane chemical potential in the sample cell, the presence of oxygen, and the unsaturated methane concentration in the seawater, are likely reasons contributing to the dissolution of the EGHS.

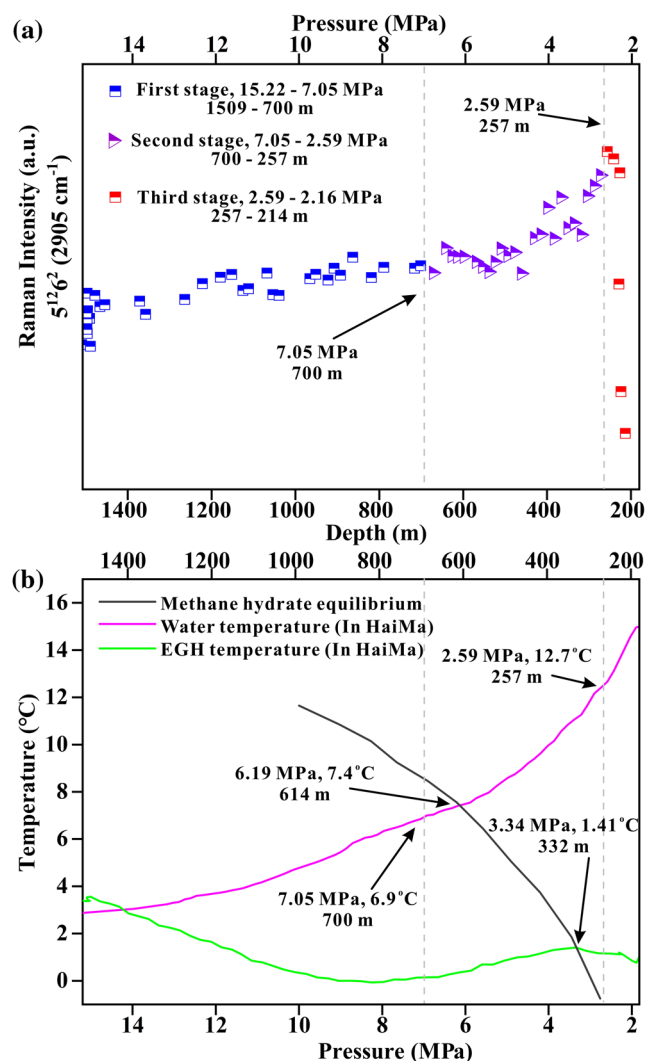


Figure 4 (a) Variation in the large cage Raman intensity with pressure during EGH ascent. (b) Temperature-pressure conditions present in the ambient seawater and the interior of the hydrate during EGH ascent. The black curve represents the phase equilibrium conditions for methane hydrate in seawater ($S = 35\%$) (Dickens and Quinby-Hunt, 1994).

However, when the EGH samples decomposed at the periphery boundary during the second stage, the internal temperature increased by $1.02\text{ }^{\circ}\text{C}$ (Fig. 4b), and the temperature of the ambient seawater increased by $5.7\text{ }^{\circ}\text{C}$. During this stage, the inner environment of the hydrate was gradually exposed due to the periphery of the EGHs decomposing. The influence of seawater ambient temperature on the internal area of the hydrates intensified. The increase in the EGH inner temperature was likely influenced by the wide variation in the ambient temperature of the seawater. In the third stage, an approximately $1.5\text{ }^{\circ}\text{C}$ increase in ambient temperature occurred at water depths of 257–214 m. Since the third stage only spanned 43 m, the influence of the ambient temperature of the seawater on the internal hydrates was small. The EGH internal temperature decreased by $0.31\text{ }^{\circ}\text{C}$ at this stage due to the hydrate decomposition and endothermic processes.

New Insights into the Vertical Spreading of Bubbles in Cold Seep Areas

The depth of the location at which the Raman signal disappeared during the third stage represents the complete EGH

decomposition. We estimated the variation in the hydrate volume with depth in the sample cells (Fig. S-8). All experiments showed that the gas hydrate was still retained above the sample cell at the end of the third stage. The rate of gas hydrate decomposition during the third stage was the fastest. The gas hydrate decomposition rates of the third stage in our three *in situ* experiments were estimated to be approximately $7.1 \pm 0.32\text{ cm}^3/\text{m}$ at the Haima cold seep vent site, $21.8 \pm 0.31\text{ cm}^3/\text{m}$ at the Site F, and $3.7 \pm 0.22\text{ cm}^3/\text{s}$ at the Lingshui site (Fig. S-8). Even considering the rapid decomposition rate that occurred at the third stage, the ROV retained some of the hydrate when it rose to the water surface. The EGHs can significantly enhance the survival of methane bubbles. The EGHs easily form in the cold seep areas on the seafloor, in shallow surface sediments, beneath authigenic carbonate rocks, or even in an empty mussel shell (Fig. S-9). Given the abundance of active cold seep vents worldwide, the potential environmental impact caused by the ascent and subsequent decomposition of EGHs carrying methane bubbles cannot be ignored.

Conclusions

Here, we provide the first record of the destabilising ascent of EGHs. The evolutionary patterns of EGHs throughout the water depth scale were established, including the gas escape processes, and the specific temperature variations. Notably, the gas escape process occurred throughout all depths in the experiment. At water pressures $> 7.05\text{ MPa}$ (depths $> 700\text{ m}$), the first stage of the evolution of EGHs was observed, during which the morphology of EGHs remained stable. Pressures of 7.05–2.59 MPa (depths of 700–257 m) corresponded to the second stage of EGH ascent, during which peripheral hydrate decomposition and internal hydrate growth coexisted. The decomposition of the peripheral hydrates will cause a hysteresis effect on the decomposition of the hydrates. The third stage (2.59–2.16 MPa, 257–214 m) was the period of internal hydrate decomposition. From our *in situ* experiments, it appears that the presence of EGHs results in methane gas influencing the overlying water column or even the atmospheric environment. It can also provide *in situ* data for gas hydrate exploitation to avoid construction-induced block hydrate uplift, which may increase greenhouse gas emissions.

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Additional Information

Supplementary Information accompanies this letter at <https://www.geochemicalperspectivesletters.org/article2327>.



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