

Hydrate morphology: Physical properties of sands with patchy hydrate saturation

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[1] The physical properties of gas hydrate-bearing sediments depend on the volume fraction and spatial distribution of the hydrate phase. The host sediment grain size and the state of effective stress determine the hydrate morphology in sediments; this information can be used to significantly constrain estimates of the physical properties of hydrate-bearing sediments, including the coarse-grained sands subjected to high effective stress that are of interest as potential energy resources. Reported data and physical analyses suggest hydrate-bearing sands contain a heterogeneous, patchy hydrate distribution, whereby zones with 100% pore-space hydrate saturation are embedded in hydrate-free sand. Accounting for patchy rather than homogeneous hydrate distribution yields more tightly constrained estimates of physical properties in hydrate-bearing sands and captures observed physical-property dependencies on hydrate saturation. For example, numerical modeling results of sands with patchy saturation agree with experimental observation, showing a transition in stiffness starting near the series bound at low hydrate saturations but moving toward the parallel bound at high hydrate saturations. The hydrate-patch size itself impacts the physical properties of hydrate-bearing sediments; for example, at constant hydrate saturation, we find that conductivity (electrical, hydraulic and thermal) increases as the number of hydrate-saturated patches increases. This increase reflects the larger number of conductive flow paths that exist in specimens with many small hydrate-saturated patches in comparison to specimens in which a few large hydrate saturated patches can block flow over a significant cross-section of the specimen.

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1. Introduction

[2] Engineering protocols and stability analyses needed for safe gas production from hydrate-bearing sediments require accurate information about the sediment's physical properties and their dependence on hydrate volume fraction and spatial distribution. Prior to the early 1980s, methane gas hydrate was assumed to be uniformly distributed wherever pressure and temperature conditions were appropriate [Milkov, 2004; Trofimuk *et al.*, 1973]. Subsequent studies have revealed a heterogeneous global distribution, due in part to the importance of advective transport in supplying sufficient methane for hydrate formation [McIver, 1981; Gornitz and Fung, 1994; Buffett and Archer, 2004; Wood and Jung, 2008], as well as

spatially varying sediment properties such as grain size and porosity [Clennell *et al.*, 1999; Waite *et al.*, 2009].

[3] Connections between the sediment type, burial conditions, and resulting hydrate distributions are becoming more apparent as the inventory of high-resolution images of recovered hydrate-bearing cores increases. We have taken advantage of X-ray computed tomography [Holland *et al.*, 2008; Kneafsey *et al.*, 2010], micro-CT [Jin *et al.*, 2006; Kerker *et al.*, 2009], and magnetic resonance [Kleinberg *et al.*, 2003; Stevens *et al.*, 2008; Erslund *et al.*, 2010] to explore hydrate occurrence patterns in all types of natural sediments, and present a first-order physical model to analyze these patterns in terms of environmental conditions and sediment characteristics. Focusing on high value, energy resource environments, we then estimate a suite of physical properties for hydrate-bearing sands, containing patches of 100% hydrate-saturated sand embedded in water-filled sand.

2. Hydrate in Porous Media: Patterns

2.1. Nucleation and Growth

[4] Hydrate nucleation is favored at the water-gas interface when free gas is present [Tohidi *et al.*, 2001; Kleinberg

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et al., 2003; *Waite et al.*, 2004], and on mineral surfaces in the absence of free gas due to the reduced thermal activity and spatial distribution of water molecules in the vicinity of mineral surfaces [*Dominguez et al.*, 2000; *Page and Sear*, 2006]. Under stable pressure and temperature conditions, molecules continuously jump from one phase to the other in hydrate-water or hydrate-gas systems, and a minimum crystal size d_{cr} of several nanometers must be reached to ensure a stable nucleus for gas-hydrate growth [*Lee*, 2007]. Eventually, stable nuclei grow into the pore space between sediment grains. Continued molecular transfer across phases can gradually change the spatial hydrate distribution in sediments, concentrating hydrate into fewer, larger crystals [*Myerson*, 2002]. This “Ostwald-ripening” phenomenon occurs because the methane concentration is higher around smaller hydrate crystals according to the Gibbs-Thompson effect [*Henry et al.*, 1999; *Kwon et al.*, 2008], and diffusive transport develops from the higher methane concentrations surrounding smaller crystals toward larger crystals.

[5] Eventually, hydrate growth becomes constrained by the mineral grains. Further growth takes place by either invading neighboring pores or displacing particles to enlarge the occupied pores. Whether growth occurs in a grain-displacing or pore-filling manner depends on the balance between the local effective stress and the capillary pressure developed by the hydrate mass. At equilibrium, the pressure in the hydrate phase u_h must balance the pressure in the water phase u_w and the effective stress acting on the grain skeleton σ' for a large interface:

$$u_h = u_w + \sigma'. \quad (1)$$

In addition, pressures in the hydrate u_h and water u_w phases are related through Laplace’s equation [*Clennell et al.*, 1999; *Coussy*, 2010]

$$(u_h - u_w) = \frac{4\gamma_{hw} \cos \theta}{d_{th}}, \quad (2)$$

where the contact angle is assumed here to be $\theta = 0^\circ$, and the hydrate-water interfacial tension is $\gamma_{hw} = 0.032\text{--}0.039$ N/m [*Uchida et al.*, 1999; *Anderson et al.*, 2003]. The pore throat diameter d_{th} is determined by the fine sediment fraction, taken here to be the finest 10th percentile in the grain size distribution. The pore throat diameter is $d_{th} \cong (\sqrt{2} - 1)d_{10}$ for simple cubic packing. These equations are combined to define a dimensionless parameter:

$$\psi = \frac{\left(\frac{4\gamma_{hw}}{d_{th}}\right)}{\sigma'} = \frac{4\gamma_{hw}}{(\sqrt{2} - 1)d_{10}\sigma'} \cong \frac{10\gamma_{hw}}{d_{10}\sigma'}, \quad (3)$$

which reflects the relative balance between hydrate-water capillarity and effective stress. When $\psi > 1$, capillarity prevails and hydrate can displace particles more readily than invade pore throats; thus, we anticipate that fine-grained soils (small d_{10}) under low effective stress σ' are prone to grain-displacement and segregated hydrate formation. Conversely, $\psi < 1$ corresponds to coarse-grained sediments (large d_{10}) subjected to high effective stress σ' , and hydrate growth will fill and invade existing pores rather than displace grains. Therefore, Ostwald ripening in coarse-grained sediments will promote patchy hydrate distributions, whereby hydrate-filled

sediment patches ($S_h = 100\%$) are embedded in a hydrate-free sediment ($S_h = 0\%$).

2.2. Hydrate Occurrence Patterns: Field Evidence

[6] Reported hydrate morphologies in natural hydrate-bearing sediments are summarized in Table 1, together with reservoir location, core depth, and sediment characteristics. Figure 1 shows the same data plotted in terms of vertical effective stress σ'_v versus effective grain size d_{10} . The vertical effective stress is calculated using the sediment depth and an assumed sediment density of 2000 kg/m³. The effective grain size d_{10} is estimated from reported grain size distributions and sediment descriptions.

[7] Observations show that naturally occurring hydrate exhibits three general morphologies: pore-fill, lenses/veins, and nodules/chunks [*Collett et al.*, 2009; *Holland et al.*, 2008]. As shown in Figure 1, pore-filling hydrate morphology corresponds to deep, coarse-grained sediments for which $\psi < 10^{-3}$. Hydrate forms in nodules and chunks when hydrate crystals can grow by displacing particles in every direction in an entirely hydrate-water capillarity dominated regime where $\psi > 1$. At intermediate conditions, $10^{-3} < \psi < 1$, hydrate growth is capillarity controlled but grain displacement is preferentially aligned normal to the local minimum principal stress σ'_3 and leads to lens formation [*Shin and Santamarina*, 2011; *Cook et al.*, 2008; *Daigle and Dugan*, 2010].

3. Physical Property Estimation for Patchy Saturation: Methods

[8] The physical properties of hydrate-bearing sediments depend on the spatial distribution of gas hydrate, which is generally not known a priori. Observations summarized in Figure 1 allow us to constrain the range of possible morphologies, and thereby tighten realistic upper and lower bounds for physical property values.

[9] In particular, pore-filling conditions ($\psi < 10^{-3}$) and Ostwald ripening combine to generate “patchy hydrate saturation” in mature hydrate-bearing coarse-grained sediments. From a modeling perspective, patchy saturation is a two-component combination of hydrate-saturated sand patches ($S_h = 100\%$) embedded in water-saturated, hydrate-free sediment ($S_h = 0\%$), rather than a three-component system (either mineral-hydrate-water or mineral-hydrate-gas). In the following sections, we use analytical models and numerical simulations to estimate core-scale physical properties of patchy hydrate-bearing sediments, and demonstrate the benefits of utilizing the two-component nature of the patchy hydrate distribution to provide a tighter range of physical property estimates than can be obtained for a three-component system. Note that logging and seismic investigations operate at lower resolution [*Shankar and Riedel*, 2011] and are less sensitive to hydrate morphology [*Knight and Nolenhoeksema*, 1990; *Li et al.*, 2001]; instead, they reflect larger lithologic-scale heterogeneities (reviewed in *Waite et al.* [2009]).

3.1. Upper and Lower Bounds

[10] Upper and lower bounds can be computed for a given physical property by assuming end-member arrangements of the components (Tables 2a and 2b). The parallel and series configurations, also known as the Voigt and Reuss models, are easily calculated but are broader than necessary. Tighter

Table 1. Hydrate Morphologies in Natural Hydrate-Bearing Sediments^a

Location	No.	Depth ^b (m)	Water Depth (m)	Hydrate Morphology	Sediment Description	Index Properties ^c	d_{10} (μm)		Typical Values	Reference ^d
							Mean	s.dev.		
Mallik	1	913.7	913.7	pore filling	granular/pebbles (max 1 cm)	$d_{10} = 275.9\text{--}292.9 \mu\text{m}$ (at 912.3 m)	285	8.5		a,b
	2	903	903	pore filling	medium-grained sand	$d_{10} = 108.4 \mu\text{m}$ (at 908.3 m)	108.4	76.02		a,b
	3	892.8–1088.1	990.45	pore filling	medium/fine sands	$d_{50} = 149.9\text{--}502.5 \mu\text{m}$	150	50	fine/medium sands: 0.06–0.6 mm	a,b
Blake Ridge	4	259	3058.1	nodules	nanofossil-rich clay	$d_{50} = 1.6 \mu\text{m}$	0.16	0.11		c, d
	5	330	3100.1	chunks	55%illite + 45%Smectite		0.001	0.0007	Smectite: ~ 1 nm	c, d
	6	3.09	2781.59	nodules/chunks	soft clay (39.2% clay)	LL = 68; PL = 24	0.2	0.14	Kaolinite: $\sim 0.1\text{--}1 \mu\text{m}$	e, f
Nankai Trough	7	207.8–260	1178.9	pore filling	sandy sediments		100	70.13	fine/medium sands: 0.06–0.6 mm	g
	8	90–140	4799.3	veins	60% sand/granules + 40% silt	$S_s = 87\text{--}94 \text{ m}^2/\text{g}$;	10	7.01	silt: 2–60 μm	h
NGHP	9	60	1300	veins (nodules?)	fine grained clay minerals	LL = 73–75; PL = 34–36	0.2	0.14	Kaolinite: $\sim 0.1\text{--}1 \mu\text{m}$	i, j
	10	58.44	1107.44	veins		$\phi_{10} = 9.28$; $\phi_{50} = 6.08$	1.6	1.12		k
Cascadia Margin	11	3.7	678.2	nodules	soft clay sediments	$\phi_{50} = 5.7$; S = 1.2;	4	0.3		l, m
	12	13.05	687.55	nodules		SK = -0.81 (at 3.59 m)	2.5	0.2		m, n
Offshore Peru	13	141	3961	nodules	diatomaceous muds	SK = -0.58	0.01	0.01	Illite: $\sim 10\text{--}100$ nm	o, p
	14	165.6	5235.6	nodules	(44%Illite)		23	8		p, q
Okushiri Ridge	15	88.3–97.9	2715.7	pore filling	Sandy sediments	$\phi_{10} = 5\text{--}6$	100	70.13	fine/medium sands: 0.06–0.6 mm	r
	16	88.3–97.9	2715.7	nodule	diatom-bearing clays, silty clay	LL = 113–143 (at 83.75–189.9 m, 799A)	0.01	0.01	Illite: $\sim 10\text{--}100$ nm	r, s
Ulleung Basin	17	91.07	2661.67	nodule	abundant diatoms	75% sand + 15% silt + 10%clay	0.2	0.1	Kaolinite: $\sim 0.1\text{--}1 \mu\text{m}$	r
	18	136	2100	veins/lenses	microfossile, clay	$d_{50} = 3.041 \mu\text{m}$; $d_{10} = 1.5 \mu\text{m}$	1.5	1.05		t
Orca Basin, GOM	19	140	2218	vein and nodule		18%clay + 82% silt $S_s = 71$ (at 141 m)	0.2	0.14	Kaolinite: $\sim 0.1\text{--}1 \mu\text{m}$	u, v
	20	20–40	2425	nodules	smectite dominant clay	LL = 83; PL = 31	0.001	0.0007	Smectite ~ 1 nm	w, x, y
Alaminos Canyon, GOM	21	~ 400	~ 3130	pore filling	fine sands	$\phi_{50} = 6.13$; S = 1.73;	80	56.10	fine sands: 60–200 μm	z
	22	2.3–2.7	~ 840	nodules	20.42% clay	SK = 1.4; Kt = 2.16	0.45	0.25		aa, ab
Mt. Elbert	23	1.65–2.65	~ 670	nodules/veins	23.4% clay	$\phi_{50} = 6.35$; S = 1.70;	0.4	0.2		aa, ab
	24	619.9	619.9	pore filling	fine quartz sand	SK = 1.31; Kt = 2.09	200	140.26		ac, ad
Atwater Valley	25	661	661	pore filling	fine-grained quarts	$d_{10} = \sim 0.2$ mm (at 618.1 m)	80	56.10		ac, ad
	26	620	620	vein	silty sand	$d_{10} = 0.017$ mm	17	11.92		ae
Hydrate Ridge	27	0–158	1370	nodules	clay	$S_s = 94.2\text{--}143.1 \text{ m}^2/\text{g}$	0.01	0.01	Illite: $\sim 10\text{--}100$ nm	af
	28	5–106	945.5	nodules	clay	$d_{50} = 6 \mu\text{m}$; $d_{10} = \sim 0.2$ mm	0.2	0.14		ag

^aThese data reflect properties of each recovered core only, and may not necessarily represent the properties of entire reservoirs.

^bDepth (m); sediment depth, which is counted as the depth from ground surface for permafrost sediments or from seafloor for marine sediments.

^cProperties: LL: liquid limit; PL: plastic limit; S_s : specific surface; ϕ : the ϕ scale of particle size; S: sorting; SK: skewness; Kt: kurtosis. The mean values of d_{10} are chosen either based on reported data or from typical particle sizes listed in the Notes column. The s. dev. (standard deviation) is computed as $s.d.[X] = \text{Mean} \times \sqrt{\phi^{0.4} - 1}$, since most natural sediments present log-normal distributions with standard deviation $N(\sigma^2) = 0.4 \pm 0.2$ [Phadnis and Santamarina, 2011].

^dReferences are as follows: a, Uchida et al. [2000]; b, Kulenkampff and Spangenberg [2005]; c, Matsumoto et al. [2000]; d, Collett and Wendlandt [2000]; e, Winters [2000a]; f, Winters [2000b]; g, Uchida et al. [2004]; h, Taira et al. [1991]; i, Yun et al. [2010]; j, Collett et al. [2008]; k, Winters et al. [2008]; l, Hovland et al. [1995]; m, Camerlenghi et al. [1995]; n, Westbrook et al. [1994]; o, von Huene et al. [1987]; p, Kvernolden and Kasner [1990]; q, Hill and Marsters [1990]; r, Tamaki et al. [1990]; s, Holler [1992]; t, Yun et al. [2011]; u, Kim et al. [2011]; v, Kwon et al. [2011]; w, Pflaum et al. [1985]; x, Thayer et al. [1986]; y, Bryant et al. [1986]; z, Boswell et al. [2009]; aa, Luan et al. [2008]; ab, Dang et al. [2010]; ac, Stern et al. [2011]; ad, Winters et al. [2011]; ae, Dai et al. [2011]; af, Yun et al. [2006]; ag, Tan et al. [2006].

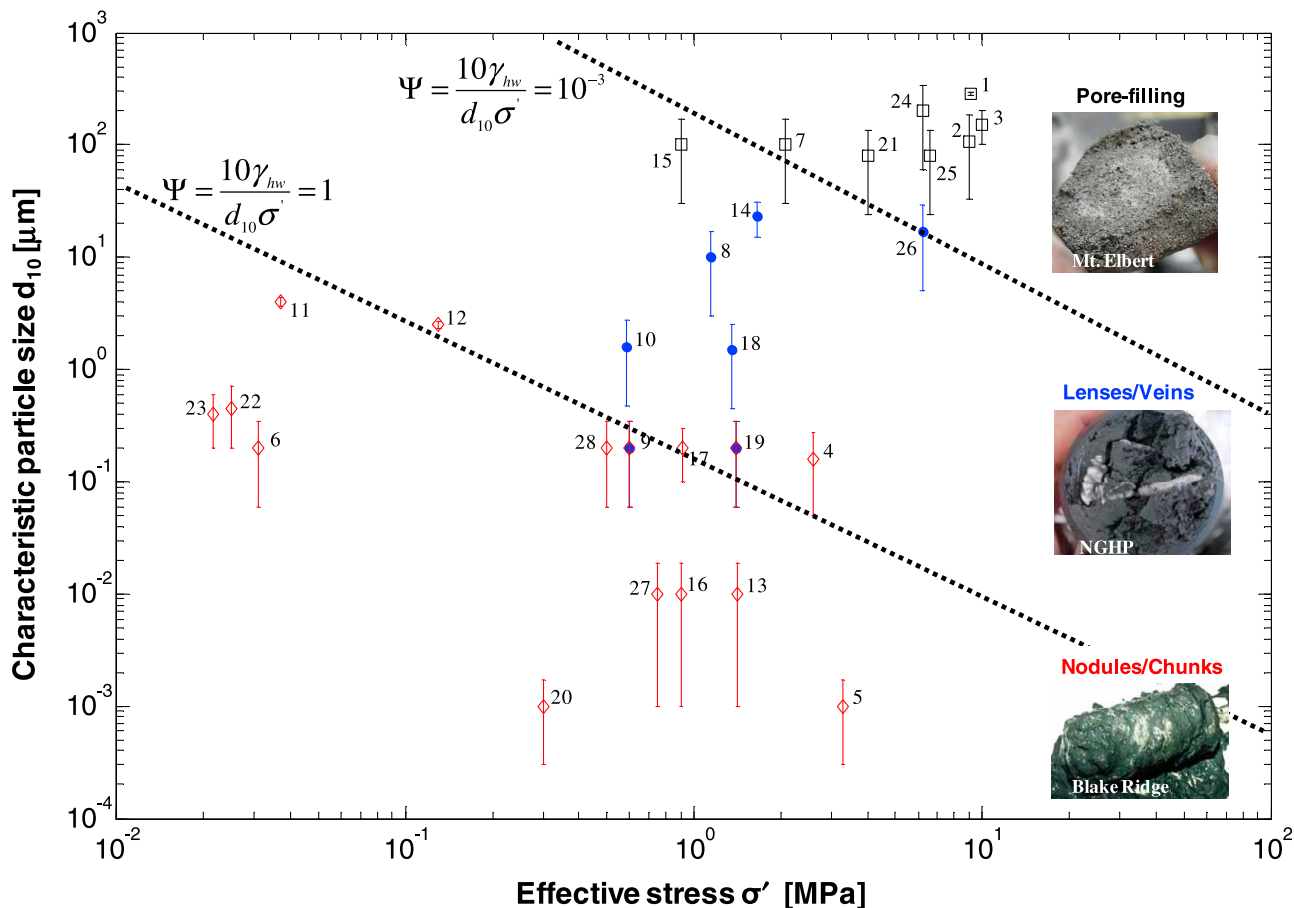


Figure 1. Hydrate morphologies in natural sediments – Field evidence and ψ -boundaries. In response to a combination of sediment particle size and effective stress, hydrate formation occurs in one of three general morphologies: Environments with relatively large sediment grains and high effective stress can host pore-filling hydrates (top inset); in contrast, hydrates forming in fine-grained sediment with low effective stress can displace sediment grains in all directions and create hydrate nodules or chunks (bottom inset). Between these two extremes, hydrate formation responds to variations in the local stress, displacing sediment grains in the directions of the smallest stress to form hydrate lenses and veins (middle inset). All data are based on core analysis and do not necessarily reflect reservoir properties (refer to Table 1 for details).

bounds are computed assuming the shell-like arrangement of individual components described by Hashin and Shtrikman [1962, 1963], and illustrated in Tables 2a and 2b. Hashin-Shtrikman bounds apply to hydraulic, thermal, and electrical conductivities, electrical permittivity, magnetic permeability, and both bulk and shear moduli.

3.2. Analytical Models

[11] Effective medium models can be used to estimate physical properties of mixtures when their spatial configuration is known (Tables 3a and 3b). The Kuster-Toksöz effective medium model estimates effective properties of a combined medium with known host and patch properties, and it can account for patch shape, e.g., spherical, needle-like, and disk-shaped cracks [Kuster and Toksöz, 1974; Berryman, 1995]. The model converges to the Hashin-Shtrikman upper or lower bounds for spherical patches. Due to the asymmetric nature of this model, results depend on the material selected to be either the patch or the host. In contrast, the self-consistent model is symmetric, meaning all components in the system are considered to be patchy inclusions, and the numerical results

are the same regardless of which material is defined as the patch, and which is defined as the host [Berryman, 1980]. The bulk and shear moduli must be iteratively solved in this model, but other properties are computed directly, including conductivity, permeability, and permittivity (see Tables 3a and 3b). These analytical models address first order quasi-static effects,

Table 2a. Physical Properties and Upper and Lower Bound Models: General Expressions^a

Model	Bound	Equation	Equation Number
Voigt (parallel) ^b	Upper	$X = \sum f_i x_i$	T1
Reuss (series) ^b	Lower	$X = (\sum f_i / x_i)^{-1}$	T2
Hashin-Shtrikman ^c	Upper	$X = x_{\max} + \frac{\xi A_{\max}}{1 - \xi A_{\max} / x_{\max}}$	T3
	Lower	$X = x_{\min} + \frac{\xi A_{\min}}{1 - \xi A_{\min} / x_{\min}}$	T4

^aX – bulk property of the mixture; x_i – physical property of the i th constituent; f_i - volume fraction of the i th constituent; $A_{\max, \min} =$

$\sum \frac{f_i}{\alpha_{\max, \min} + \xi / (x_i - x_{\max, \min})}$, see Table 2b for α and ξ .

^bMavko et al. [1998].

^cHashin and Shtrikman [1962, 1963].

Table 2b. Property-Dependent α and ξ Parameters, to Be Substituted Into the Expressions for X and $A_{\max,\min}$ From Table 2a^a

Properties	α		ξ
	max	min	
Hydraulic conductivity $k_H^{b,c}$	$\alpha_{\max} = \frac{1}{3x_{\max}}$	$\alpha_{\min} = \frac{1}{3x_{\min}}$	$\xi = 1$
Thermal conductivity λ^d			
Electrical conductivity σ_e^d			
Magnetic permeability μ^d			
Dielectric permittivity ϵ_r^d			
Bulk modulus K^c	$\alpha_{\max} = \frac{3}{3K_{\max}+4G_{\max}}$	$\alpha_{\min} = \frac{3}{3K_{\min}+4G_{\min}}$	$\xi = 1$
Shear modulus G^c	$\alpha_{\max} = \frac{3(K_{\max}+2G_{\max})}{5G_{\max}(3K_{\max}+4G_{\max})}$	$\alpha_{\min} = \frac{3(K_{\min}+2G_{\min})}{5G_{\min}(3K_{\min}+4G_{\min})}$	$\xi = 0.5$

^aAs in Table 2a, x_{\max} and x_{\min} are the maximum and minimum physical property values, respectively, for the components in the mixture.

^bZimmerman and Bodvarsson [1995].

^cZimmerman et al. [1991].

^dHarter and Knudby [2004].

^eHashin and Shtrikman [1962, 1963].

and are well-suited for providing a unified approach for estimating a suite of physical properties even in cases where only basic field data are available.

[12] More specialized physical property treatments of patchy-saturated rocks [e.g., Dvorkin and Nur, 1998; Vogelaar et al., 2010; Quintal et al., 2011; see also Mavko and Mukerji, 1998] generally deal with gas patches embedded within fluid-saturated media, where both gas-filled and fluid-fill patches have identical shear stiffness (or shear wave velocity). This uniform shear stiffness simplification cannot be applied to hydrate-filled patches where solid hydrate significantly enhances the stiffness of hydrate-filled sediment. Our low-frequency models do not consider dispersion due to either viscous losses or polarization effects [Asami, 2005].

3.3. Numerical Simulations

[13] Numerical simulations allow us to capture conditions beyond the assumptions made in analytical models. In this study, simulations were implemented in COMSOL Multiphysics. We use cylindrical specimens (radius $R = 0.05$ m and height $H = 0.2$ m) containing spherical hydrate-saturated sand patches. The number of patches varies from 1 to 200 while the specimen's hydrate saturation is held constant ($S_h = 0.2$). Several realizations of randomly distributed patches are generated for a given number of hydrate patches. The properties of hydrate-saturated sand patches and hydrate-free sand matrix listed in Table 4 are used as inputs to the numerical model.

[14] The 3D numerical model is meshed using tetrahedral element, and the number of elements increases with the number of patches, varying from $\sim 10,000$ to $\sim 300,000$. To compute moduli, the cylindrical specimen is loaded with an isotropic stress σ_i and the bulk modulus K is calculated as

$K = \Delta\sigma_i/\Delta\varepsilon_v$, where $\Delta\varepsilon_v$ is the computed volumetric strain increment for a change in isotropic stress $\Delta\sigma_i$. When computing conductivities, the cylindrical specimen is surrounded by non-conductive lateral boundaries. A longitudinal gradient is then imposed to generate the flux appropriate to the type of conductivity being considered. Conductivities are calculated as the ratio between the computed flux and the applied gradient.

[15] A complementary set of simulations is conducted for 20% hydrate-free sand patches embedded in a hydrate-saturated sand host (i.e., $S_h = 80\%$).

4. Physical Properties of a Patchy System: Results

[16] We use the analytical models and numerical simulations described above to estimate the physical properties of hydrate-bearing sands with patchy-hydrate saturation. Table 4 lists the physical property values selected for these computations. These values correspond to hydrate-bearing sands with porosity $n = 0.4$ and effective stress $\sigma' = 1$ MPa.

4.1. Bulk Modulus

[17] Upper and lower bounds provide quick estimates to assess the plausibility of measured values. Parallel, series, and Hashin-Shtrikman bounds are shown in Figure 2 for the bulk modulus K of hydrate-bearing sands, either assuming a three-component system (i.e., mineral, hydrate, and water) or a two-component patchy mixture (i.e., patches with $S_h = 100\%$ and sand with $S_h = 0\%$). The patchy saturation assumption significantly reduces the gap between upper and lower bounds because it removes the effect of extreme values for single components, such as the null shear stiffness of water in a series

Table 3a. Physical Properties: Analytical Estimates^a

Model	Equation	Equation Number
Kuster-Toksoz ^b	$(K_{KT} - K_m) \frac{K_m + 4G_m/3}{K_{KT} + 4G_m/3} = \sum f_i (K_i - K_m) P_{m,i}$	T5
	$(G_{KT} - G_m) \frac{G_m + 4G_m/3}{G_{KT} + 4G_m/3} = \sum f_i (G_i - G_m) Q_{m,i}$	T6
Self-consistent ^c	$0 = \sum f_i (X_i - X) Z_{SC,i}$	T7

^aAnalytical models for estimating, rather than bounding, the physical properties of gas hydrate-bearing sediment. X is the estimated bulk property of the mixture. The terms $P_{m,i}$, $Q_{m,i}$, and $Z_{SC,i}$ depend on the patch shapes and physical properties, as expressed in Table 3b.

^bKuster and Toksoz [1974].

^cBerryman [1980].

Table 3b. Shape Factors for the Kuster-Toksoz ($P_{m,i}$, $Q_{m,i}$) and Self-Consistent Models ($Z_{sc,i}$)^a

	Spherical	Needle	Disk ^b
Conductivity ^c $Z_{sc,i} =$	$\frac{1}{X_i + 2X_m}$	$\frac{1}{9} \left(\frac{1}{X_m} + \frac{4}{X_i + X_m} \right)$	$\frac{1}{9} \left(\frac{1}{X_i} + \frac{2}{X_m} \right)$
Bulk modulus $Z_{sc,i}$ or $P_{m,i} =$	$\frac{K_m + 4G_m/3}{K_i + 4G_m/3}$	$\frac{K_m + G_m + G_i/3}{K_i + G_m + G_i/3}$	$\frac{K_m + 4G_i/3}{K_i + 4G_i/3}$
Shear modulus ^d $Z_{sc,i}$ or $Q_{m,i} =$	$\frac{G_m + \zeta_m}{G_i + \zeta_m}$	$\frac{1}{5} \left(\frac{4G_m}{G_i + G_m} + 2 \frac{G_m + \vartheta_m}{G_i + \vartheta_m} + \frac{K_i + 4G_m/3}{K_i + G_m + G_i/3} \right)$	$\frac{G_m + \zeta_i}{G_i + \zeta_i}$

^aFor $P_{m,i}$ and $Q_{m,i}$, the subscript m refers to the property value for the maximum or minimum constituent, depending on whether the upper or lower bound is being calculated. For $Z_{sc,i}$, m refers to the value for the final self-consistent material. In all cases, the subscript i refers to the value for i th component in the system [Berryman, 1995].

^bBecause of the $1/X_i$ term, the disk approximation is calculated in Figure 4 (bottom) with a hydrate patch conductivity of $X_i = (1 \times 10^{-5}) \cdot X_{hydrate-free\ patches}$, rather than setting $X_i = 0$.

^cAppropriate for the conductivity, permeability, and permittivity properties listed in Table 3b. Though Berryman [1995] does not recommend this approach for hydraulic conductivity, equivalent relations have been derived specifically for hydraulic conductivity by Pozdniakov and Tsang [2004].

$$\zeta_m = \frac{G_m}{6} \frac{9K_m + 8G_m}{K_m + 2G_m}, \zeta_i = \frac{G_i}{6} \frac{9K_i + 8G_i}{K_i + 2G_i}, \vartheta_m = G_m \frac{3K_m + G_m}{3K_m + 7G_m}, \vartheta_i = G_i \frac{3K_i + G_i}{3K_i + 7G_i}$$

configuration, or the high stiffness of quartz for a parallel arrangement (Table 4).

[18] Numerical simulations (solid dots in Figure 2) are obtained by assuming the host medium and inclusions are linearly elastic (values listed in Table 4). The numerically computed bulk modulus for hydrate-bearing sediments approaches the lower bound when the global hydrate saturation is low ($S_h < \sim 20\%$) and the upper bound when the global hydrate saturation is high ($S_h > \sim 80\%$). Moduli calculated from laboratory velocity measurements for various types of sediments containing THF hydrate reveal similar trends over a range of hydrate saturations [Lee et al., 2010].

[19] Numerical results show the increase in bulk stiffness of hydrate-bearing sediments with the number of hydrate-saturated patches for a given hydrate saturation (Figure 2).

This result reflects interaction among patches as the number of patches increases and their relative distance becomes less than two times the patch size. Grain-based discrete-element simulations confirm that strong force chains form between large, hydrate-saturated sand patches [Jung et al., 2012]. Because the geometric distribution of patches in cylindrical specimens deviates from Hashin-Shtrikman assumptions, some numerical results fall outside the Hashin-Shtrikman bounds but remain within the series and parallel bounds.

[20] The Kuster-Toksöz effective media model compares well with finite element simulation results only when the patch concentration is low because the Kuster-Toksöz approach inherently disregards the collaborative coupling among inclusions, or patches, within the host material [see also Mavko et al., 1998]. The self-consistent model, in which

Table 4. Selected Physical Properties for Two- and Three-Component Mixtures in Hydrate-Bearing Sediment Systems (Assumed Porosity $n = 0.4$ and Mean Effective Stress $\sigma'_v = 1$ MPa)

Property	Three-Component Mixture			Two-Component Mixture	
	Quartz	Seawater	Hydrate	Sediment + Seawater ($S_h = 0\%$)	Sediment + Hydrate ($S_h = 100\%$)
Mass density ρ (g/cm ³)	2.65	1.02–1.05	0.93	1.99	1.962
Poisson's ratio ν (–)	0.17	0.5	0.33 ^a	0.15 ± 0.05^b	$0.3 \pm 0.05^{b,c}$
Shear wave velocity V_s (m/s)	4120 ⁿ	0	1960 ^d	392 ^e	1881 ^e
Compressional wave velocity V_p (m/s)	6040 ⁿ	1500 ⁿ	3770 ^d	1829	3065
Bulk modulus, K (GPa)	36.6 ⁿ	2.25 ⁿ	8.41 ⁿ	5.44 ^p	15.9 ^p
Shear modulus, G (GPa)	31.2 ^o	0 ⁿ	3.54 ⁿ	0.27 ^p	7.3 ^p
Drained Young's Mod. $E_{drained}$ (GPa)	72	2.2	8.4 ^a	0.7	18.05
Thermal conductivity λ (W/m/K)	7.7–8.4 ^f	0.56 ^g –0.58 ^h	0.5–0.58 ^{c,i,j,k}	3.7 ^j	4.5 ^j
Heat capacity C (J/kg/K)	730	4200 ^h	1980–2080 ^{a,i,l,m}	2118	1270
Hydraulic conductivity k_H (mm/s)	0	n/a	0	0.1	0
Electrical conductivity σ_e (S/m)	0	5	0	1.15 ^c	0 ^e
Permittivity ϵ_r (–)	4.2	80	7	33 ^c	5 ^e

^aSloan and Koh [2008] (Methane hydrate).

^bYun et al. [2005] (drained, skeleton, small strain).

^cLee et al. [2010] (Methane hydrate, 263 K).

^dHelgerud et al. [1999] (Methane hydrate, 5 MPa, 273 K).

^eSantamarina and Ruppel [2010].

^fRevil [2000].

^gKaye and Laby [2008] (273 K).

^hWeast [1987] (273 K).

ⁱWaite et al. [2005] (THF hydrate, 261 K).

^jCortes et al. [2009] (THF hydrate, 261 K).

^kHuang and Fan [2004].

^lHanda et al. [1984].

^mHanda [1986] (Methane hydrate).

ⁿWaite et al. [2009].

^oPeselnick et al. [1967].

^pComputed values.

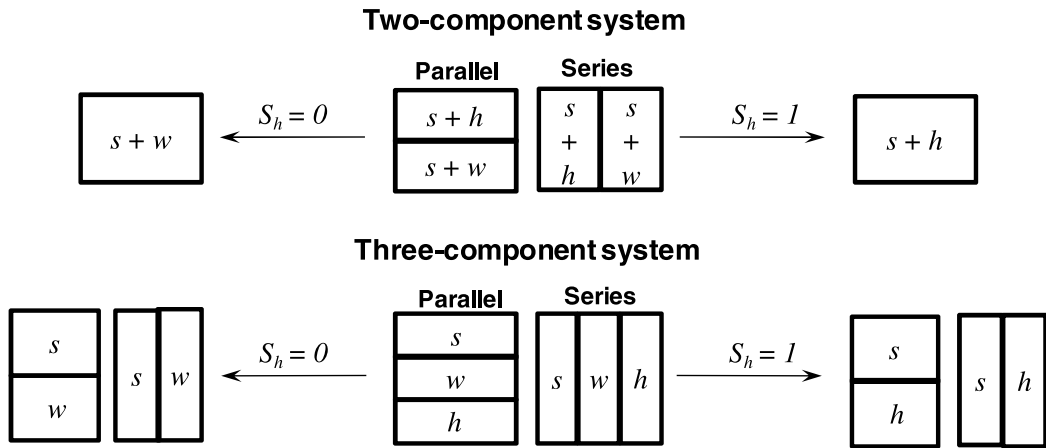
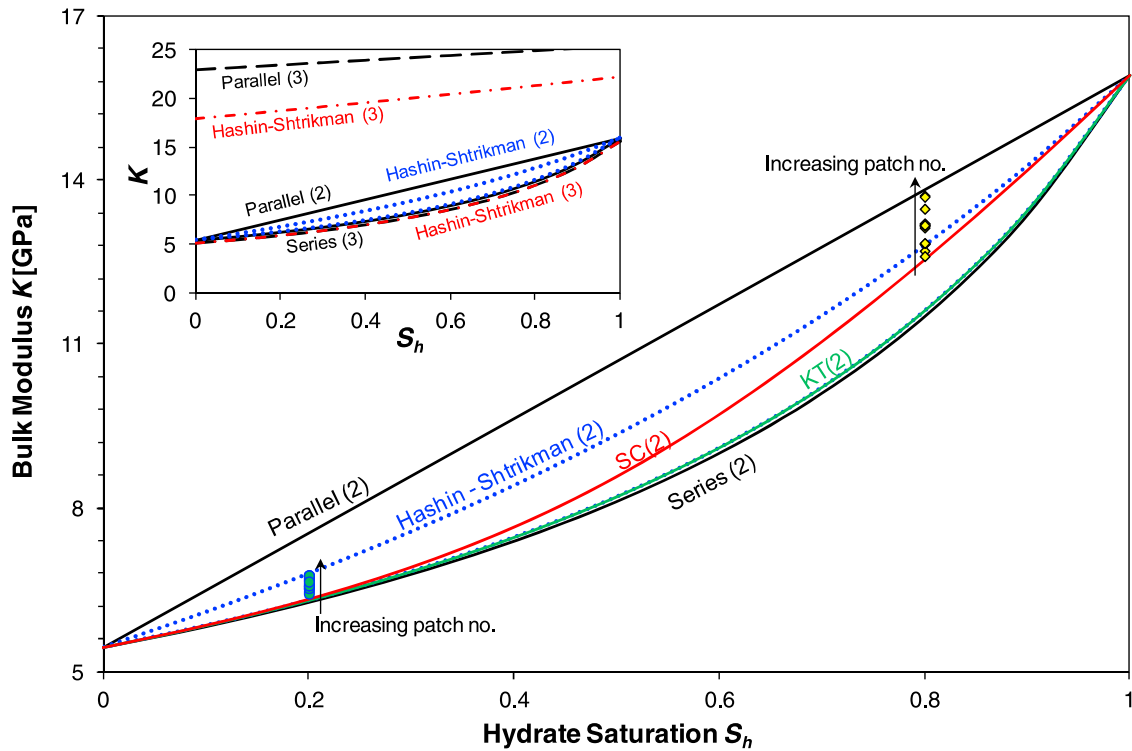


Figure 2. Bulk modulus K : bounds, analytical models, and numerical simulations. Filled circles show numerical simulation results for a fixed hydrate saturation (either $S_h = 0.2$ or 0.8) but different number of patches. KT: Kuster-Toksoz model; SC: self-consistent model. Numbers 2 and 3 after each model designate 2- and 3-component systems. The 2-component model bounds are significantly tighter than three-component bounds, a result that can be exploited when geologic conditions suggest 2-component patchy hydrate saturations can be expected rather than 3-component homogeneous hydrate distributions. Refer to Tables 2a and 2b for material parameters and Tables 3a, 3b, and 4 for mathematical expressions.

all constituents are considered inclusions and there is no set host material, outperforms the Kuster-Toksoz model even at higher hydrate-patch concentrations [Berryman, 1980], in agreement with Figure 2.

4.2. Thermal, Hydraulic and Electrical Conductivities

[21] Conduction phenomena are mathematically equivalent. Therefore, the thermal, hydraulic, and electrical conductivities are estimated using the same analytical models

(Tables 3a and 3b), inserting the appropriate physical parameters for the property of interest (Table 4).

[22] Figure 3 shows that thermal, hydraulic, and electrical conductivities increase with the number of hydrate patches for a given hydrate saturation. Though measurable, this effect is small relative to the conductivity difference between hydrate-saturated patches and hydrate-free sediment, which is shown on the right side of each plot in Figure 3. At a fixed hydrate saturation, a single large patch inside the cylindrical specimen mimics the series (or Reuss) configuration. The

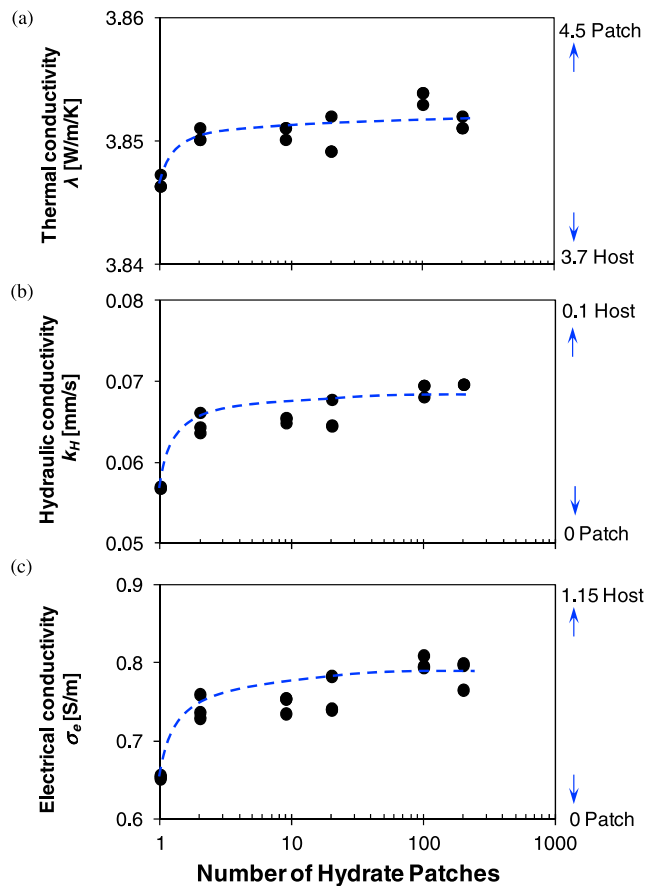


Figure 3. Conductivities versus the number of hydrate patches for a sandy specimen with porosity $n = 0.4$ and hydrate saturation $S_h = 0.2$. Dots are numerical results and dashed lines are trends. The patch size decreases as the number of patches increases to retain a constant hydrate saturation. The upper and lower end-member property values are given at the top and bottom right corner of each plot. Refer to Table 4 for material parameters.

size of each patch decreases when the number of patches increases at a fixed hydrate saturation and conduction improves as more flow paths become available.

4.3. Discussion: Impact of Hydrate Pore Habit on Physical Properties

4.3.1. Compressional P Wave Velocity

[23] Figure 4 (top) shows trends with hydrate saturation for various growth patterns. A small amount of hydrate can dramatically increase the sediment stiffness, and hence the P wave velocity, in “cementing systems” that form when water and gas phases are present (cases A and B – see experimental data in *Priest et al.* [2005, 2009] and a hybrid model in *Chand et al.* [2006]). This dramatic increase occurs because the gas-water interface clings onto water-wet minerals; then, when water converts to hydrate, the new solid phase dramatically stiffens contacts, increasing the P wave velocity even with only a small hydrate saturation. In contrast, there is no significant velocity increase at low hydrate saturations when hydrate forms from dissolved phase methane and nucleates either in the pore fluid (C) or preferentially on mineral surfaces

and grows away from grain contacts (D). Between $S_h \approx 25\text{--}40\%$, enough hydrate forms to transition the pore nucleation case (C) to the structure-building case (D) as hydrate begins contacting more than one sediment grain within a pore and the velocity begins increasing rapidly with increasing hydrate saturation. The rapid stiffness increase in patchy distributions when hydrate saturation exceeds 30–40% results from increased patch size and enhanced interactions between patches (case E). This rapid increase has also been observed experimentally [*Berge et al.*, 1999; *Yun et al.*, 2005, 2007]. Cases A, B, C, and D are calculated using the formulation in *Dvorkin et al.* [2000] (equations also provided in *Mavko et al.* [1998]). In agreement with results in Figure 2, anticipated trends for patchy saturation (E) are estimated with the self-consistent model using the properties of hydrate-free host sediment and hydrate-saturated patches.

4.3.2. Hydraulic Conductivity

[24] Pore-filling hydrate leaves thin flow pathways which significantly reduce hydraulic conductivity (Figure 4, bottom, curve H_1). More conductive, larger diameter flow paths exist in grain-coating hydrate (curve H_2) [*Kleinberg et al.*, 2003; *Spangenberg*, 2001]. Flow is effectively cut off in both of these homogeneous distribution cases by $S_h = 80\%$. Concentrating hydrate into spherical or needle-shaped hydrate saturated patches leaves hydrate-free sections; hence, spherical and needle-shaped patches are less disruptive to fluid flow than homogeneous hydrate distributions (curves P_2 and P_3). Disk-shaped patches aligned normal to flow can cause extreme flow-blocking conditions even at low hydrate saturations (curve P_1). Conversely, high patch aspect ratio in the flow direction lowers the impact of non-conductive patches on global conductivity [*Xia and Thrope*, 1988]. Spherical patches exhibit a conductivity cutoff at a percolation threshold $S_h = 2/3$; for higher hydrate saturation, the needle-shaped patch in the self-consistent model (i.e., curve P_3) may better represent field conditions [*Berryman*, 1995].

[25] The analysis based on patchy saturation can also be applied to natural hydrate-bearing fines where lenses or chunks are made of 100% segregated hydrate, with the caveat that anisotropy can control such systems. Conductivities can become patch shape- and orientation- dependent, as has been observed in recent gas hydrate logging studies in fine-grained sediments [*Cook et al.*, 2008; *Lee and Collett*, 2009].

5. Conclusions

[26] The lithologically determined and spatially varying hydraulic conductivity in sediments control advective methane transport and the reservoir-scale heterogeneous hydrate distribution [*Nimblett and Ruppel*, 2003; *Weinberger and Brown*, 2006]. Yet, hydrate morphology at the meso-scale is inherently controlled by the state of effective stress σ' and the grain size of the finer fraction of the sediment d_{10} . Therefore, the meso-scale hydrate morphology can be anticipated from burial depth and sediment type.

[27] Heterogeneous hydrate distributions can be grouped into (1) hydrate-water capillarity-controlled grain-displacing nodules and chunks in fine-grained sediments at low effective stress, (2) lenses and veins in fine-grained sediments at intermediate effective stress where hydrate-water capillarity and effective stress affect growth, and (3) patchy saturation

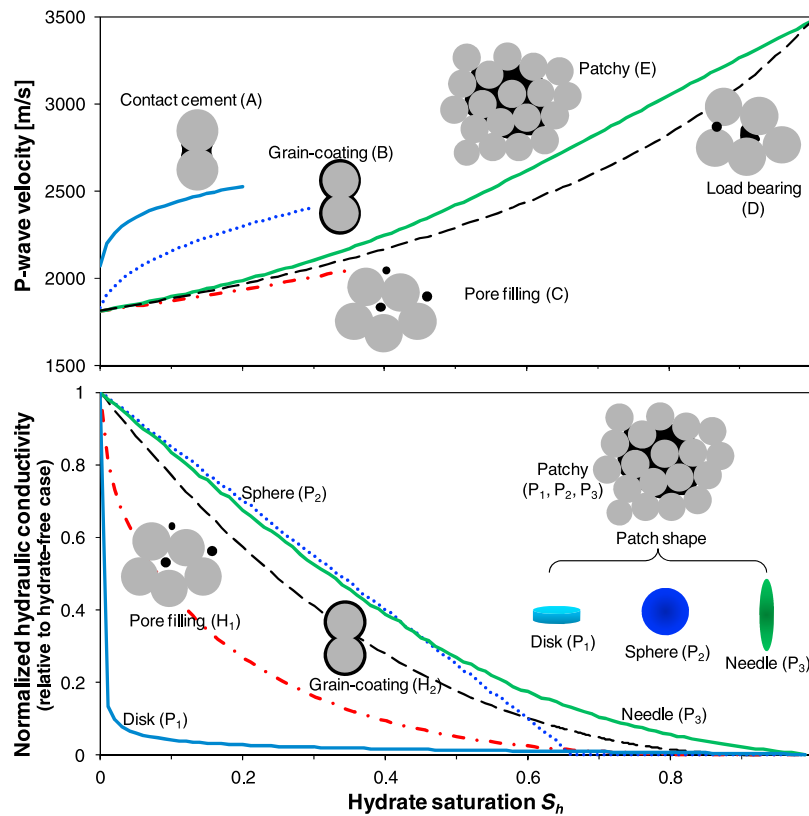


Figure 4. Self-consistent versus pore-habit specific models: predicted dependence of physical properties on gas hydrate saturation. (top) Compressional, P wave velocity: P wave velocity increases rapidly even at very low hydrate saturations when hydrate forms at sediment grain contacts. When hydrate forms away from grain contacts (C, D), the velocity increases more rapidly at high hydrate saturation when hydrate begins bridging between multiple sediment grains across a pore or fully hydrate-saturated patches become large and numerous enough to begin interacting (E). Hydrate contacts sediment grains and can no longer be considered purely part of the pore fluid above $S_h \approx 25\text{--}40\%$. (bottom) Normalized hydraulic conductivity relative to the hydrate-free $S_h = 0$ state: homogeneous hydrate distributions including pore-filling (H_1) or grain coating (H_2) hydrates versus patchy hydrate distributions with disk- (P_1), sphere- (P_2), or needle- shaped (P_3) hydrates. Homogeneous curves H_1 and H_2 are calculated from Kleinberg *et al.* [2003]. Patchy distribution curves P_1 , P_2 , and P_3 are calculated from Table 4 [Berryman, 1995; Pozdniakov and Tsang, 2004].

in coarse sands at high effective stress where the granular skeleton remains unaltered during hydrate growth.

[28] Hydrate-bearing sands are of particular interest as an energy resource. Accounting for patchy saturation, hydrate-bearing sands can be analyzed as a two-component system made of hydrate-saturated sand patches ($S_h = 100\%$) embedded in hydrate-free sand ($S_h = 0\%$). This morphology can be exploited to tighten the bounds of potential physical property values relative to bounds obtained by assuming homogeneous multicomponent mixtures of mineral-hydrate-gas or mineral-hydrate-gas-water.

[29] The bulk stiffness of patchy hydrate-bearing sands approaches the lower Voigt-Reuss bound at low hydrate saturations and the upper bound at high saturations due to the mechanical interaction between hydrate-saturated patches. The self-consistent model tracks this shift from the lower-bound to the upper-bound with increasing hydrate saturation, unlike many morphology-specific models that tend to follow either the lower or upper-bound over the entire saturation range.

[30] Patchy hydrate saturation is generally less disruptive to bulk conductivities than homogeneously distributed hydrates. Smaller patches increase flow tortuosity but have lower possibility of extreme flow-blocking conditions. An increased patch slenderness in the fluid flow direction decreases the impact of patches on the global sediment conductivity.

Notation

u_h	pressure in the hydrate phase, Pa.
u_w	pressure in the water phase, Pa.
σ'	effective stress, Pa.
γ_{wh}	the hydrate-water interfacial tension, N m^{-1} .
θ	contact angle, degrees.
d_{cr}	minimum crystal size, m.
d_{th}	pore throat diameter, m.
d_{10}	the finest 10th percentile in the grain size distribution, m.
Ψ	defined dimensionless stress parameter.
S_h	hydrate saturation.
R	radius, m.

H height, m.
 n porosity.
 E Young's modulus, Pa.
 σ_i isotropic stress, Pa.
 ε_v volumetric strain.
 λ thermal conductivity, $\text{W m}^{-1} \text{K}^{-1}$.
 k_H hydraulic conductivity, m s^{-1} .
 σ_e electric conductivity, S m^{-1} .
 ρ mass density, kg m^{-3} .
 ν Poisson's ratio.
 V_s shear wave velocity, m s^{-1} .
 V_p compressional wave velocity, m s^{-1} .
 K bulk modulus, Pa.
 G shear modulus, Pa.
 E_{drained} drained Young's modulus, Pa.
 C heat capacity, $\text{J kg}^{-1} \text{K}^{-1}$.
 ε_r dielectric permittivity.
 μ magnetic permeability.
 f_i volume fraction of composite i .
 LL liquid limit.
 PL plastic limit.
 S_s specific surface, m^2/g .
 ϕ the ϕ scale of particle size.
 S sorting.
 Sk skewness.
 Kt kurtosis.
 X bulk properties.
 x component properties.
 α, ξ coefficients in Hashin-Shtrikman model, refer to Tables 2a and 2b.
 P, Q, ζ, ν coefficients in Kuster-Toksoz model, refer to Tables 3a and 3b.
 Z, ζ, ν coefficients in self-consistent model, refer to Tables 3a and 3b.

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