



A hydrate shell growth model in bubble flow of water-dominated system considering intrinsic kinetics, mass and heat transfer mechanisms



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ABSTRACT

The gas hydrate formed in the wellbore or pipeline may pose severe challenges to pressure control and flow assurance. One key issue to address the hydrate problem is the intercoupling of hydrate shell growth characteristics and multiphase flow behaviors, which has not been studied thoroughly in the bubble flow of a water-dominated system. In this study, we develop a fully coupled hydro-thermo-hydrate model considering the interactions of hydrate intrinsic kinetics, mass and heat transfer, and hydrodynamics mechanisms. In the model, the varying concentration of gas on the outside of a hydrate shell, is introduced to describe the dynamic equilibrium between the gas outward diffusion within the hydrate shell, hydrate formation kinetics at the hydrate/liquid interface, and gas dissolution into liquid. The simulation results agree well with the experimental data. Using the proposed model, we study a special kick development mechanism caused by the phase transition of gas traps during deepwater horizontal drilling. The simulated results show that there exists a hydrate phase stability field in the wellbore during deepwater drilling. As the migration of gas traps, the hydrate growth on the bubble surface, which is closely related to the formation and decomposition of hydrate and the dissolution and desolvation of gas, may result in an abrupt and rapid gas kick. The proposed model adds further insights into quantitatively characterizing the hydrate growth and interphase mass transfer rules in the bubble flow of water-dominated systems.

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1. Introduction

Formed under low temperature and high pressure, the gas hydrates are crystalline inclusion compounds, in which the gas molecules are trapped in the hydrogen-bonded cages of water molecules [1]. Hydrate utilization technology has been widely used in the field of energy [2–4], such as the development of hydrate resources, solute separation and energy storage. However, as oil and gas exploration enters into the deepwater reservoirs, the thermodynamic and material conditions of hydrate formation are satisfied perfectly in the wellbore and subsea pipelines [5–7]. The formation of solid hydrate will alter the flow behaviors of gas and liquid, which can lead to severe flow assurance problems including the drilled gas kick [6], pipe blockage [7,8], etc. Therefore, it is necessary to study the hydrate phase transition rules on the gas bubbles of a water-dominated system.

The hydrate growth mechanisms are the key issues to address the flow assurance challenge. A wide variety of evidence in the lit-

erature indicates that the hydrate formation phenomenon is limited by the mechanisms of intrinsic kinetics, and mass and heat transfer [1,9]. At present, several hydrate kinetics models under different experimental conditions have been developed [10–13], which have been widely utilized to analyze the hydrate formation rules in the water-dominated systems. Using the hydrate kinetics models, Zheng et al. [14], Yapa et al. [5], and Li and Huang [15] simulated the hydrate formation characteristics for the gas bubbles during free rising, and gas bubbles in an intense plume. The hydrate shell growth was found to have an important effect on the migration process and lifetime of a gas bubble. As for the multiphase flow scenarios, Wang et al. [6], Boxall et al. [7], and Wang et al. [8] demonstrated the interactions of gas, liquid, and hydrate phases in wellbore and pipeline considering the hydrate kinetic limitation; moreover, the multiphase flow behaviors of kick migration during drilling, and the flow assurance during pipeline transportation were analyzed. The gas hydrate was observed to occur and cover the bubble surface rapidly in bubble flow (the gas void fraction is less than 25% [16]) of a water-dominated system [17–19]. The hydrate film will not prevent, but only retard, the transports of gas and liquid through it [3,17]. Therefore, the mass transfer process becomes the control step and dominating factor for the hydrate shell thickening.

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Nomenclature

a	profile parameter for low gas fraction in liquid	q	influx rate of fluid from the surrounding, kg/s
A	cross-sectional area of fluid flow, m^2	r	radial distance, m
A_p	interfacial area of gas-liquid contact, m^2	r_{ti}	inner radius of the inner pipe, m
A_0	heat transfer resistances between the annulus and surroundings, $(K\ m)/W$	r_{to}	outer radius of the inner pipe, m
B_0	heat transfer resistances between the inner pipe and annulus, $(K\ m)/W$	r_{ci}	inner radius of the outer pipe, m
c	reservoir compressibility, $1/Pa$	r_{co}	outer radius of the outer pipe, m
C_g	gas concentration, mol/m^3	r_w	wellbore radius, m
C_s	gas concentration on the inner surface of hydrate shell, mol/m^3	R	bubble radius
C_x	gas concentration on the outer surface of hydrate shell, mol/m^3	Re	Reynolds number
C_{w0}	concentration of water, mol/m^3	s	length, m
C_∞	gas concentration in the liquid bulk, mol/m^3	Δs	length of space segment, m
C_0	distribution coefficient	S	skin factor
d_c	equivalent diameter, m	t	time, s
D_g	gas diffusion coefficient in liquid, m^2/s	T	fluid temperature in the annulus, K
D_h	gas diffusion coefficient in hydrate shell, m^2/s	T_t	temperature in the inner pipe, K
e_i	internal energy of the i th phase, J/kg	T_{ei}	surrounding environment temperature, K
f	fugacity of dissolved gas, Pa	T_D	dimensionless temperature
f_{eq}	gas fugacity in three-phase equilibrium, Pa	U	terminal velocity of gas bubble, m/s
g	gravitational acceleration, m/s^2	U_t	overall heat transfer coefficient in the inner pipe, $W/(m^2\ K)$
h_i	enthalpy of the i th phase, J/kg	U_a	overall heat transfer coefficient in the annulus, $W/(m^2\ K)$
h_{oi}	enthalpy of the i th phase under the influx or leak condition, J/kg	v	velocity, m/s
h_c	convective heat transfer coefficients of the fluid, $W/(m^2\ K)$	v_d	gas drift velocity, m/s
H	Henry's constant, Pa	x_{sol}	fraction of dissolved gas in liquid, m^3/m^3
ΔH_{sol}	gas dissolution heat, J/kg		
ΔH_{hyd}	phase transition heat of hydrate, J/kg		
I_{tsfer}	heat transfer rate of the flowing fluid with the surroundings, $J/(m\ s)$	Greek letters	
$J_{diffusion}$	gas diffusion rate in the hydrate shell, mol/s	θ	angle between perpendicular and flow direction, rad
$J_{reaction}$	gas consumption or generation rate, mol/s	φ	reservoir porosity
$J_{dissolve}$	gas dissolution rate, mol/s	σ_{hl}	hydrate/liquid interfacial tension, N/m
k_g	mass transfer coefficient of gas dissolution, m/s	γ	profile parameter reduction term
k_e	thermal conductivity of the surroundings, $W/(m\ K)$	κ	Euler constant
K	reaction constant of hydrate formation or decomposition, $mol/(m^2\ Pa\ s)$	δ	hydrate shell thickness, m
K_e	reservoir permeability, m^2	ρ	density, kg/m^3
M	molar mass, kg/mol	ρ_c	average density of hydrated bubble, kg/m^3
N	number density of gas bubble, $1/m$	μ_f	fluid viscosity at characteristic temperature, Pa s
Nu	Nusselt number	μ_w	fluid viscosities at surface temperature, Pa s
p	fluid pressure, Pa	μ_g	gas viscosity, Pa s
p_{res}	reservoir pressure, Pa	α	volume fraction
Pr	Prandtl number		
		Subscripts	
		g	gas
		h	hydrate
		l	liquid
		m	mixture

Due to the mass transfer limitation, the vertical growth of a hydrate shell is much slower than its lateral growth. The experiments of the hydrate tensile strength and morphology indicate that the hydrate shell can be a porous solid material [18]. Considering the gas diffusion within the hydrate, Holder and Warzinski [20] built the steady-state equation of hydrate shell thickness, and analyzed the rules of hydrate thickening on the bubble surface. Regarding the air bubble in polar ice, Salamatin et al. [21] proposed a coupled model to describe the hydrated bubble behaviors, in which the mass transfer of different phases within the hydrate shell is incorporated. Furthermore, Shi et al. [9], Gong et al. [22], and Turner et al. [23] developed the hydrate inward growth model of water drops in oil emulsions. Gas diffusion across the hydrate shell is revealed as the major factor for hydrate shell growth. However, hydrate formation is a strongly system dependent process [9],

affected by the bubble hydrodynamics, gas-water distributions, etc. Therefore, the hydrate shell growth characteristics of the steady bubble or oil-dominated system, which are mainly studied at present [24], can be significantly different with the scenario here. This is because the gas dissolution of the moving hydrated bubble and the dynamic variation of the dissolved gas distribution can affect the driving forces for mass transfer and intrinsic kinetics.

The hydrate shell growth should be intercoupled with the multiphase flow behaviors of the system. The hydrate formation/decomposition can lead to the heat release/absorption and mass transport between different phases, and then influence the variations of volume fractions, pressure, and temperature in water-dominated systems. Conversely, the multiphase flow behaviors control the hydrate phase stability field and the hydrate formation/decomposition rate.

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