

# Experimental study on the kinetics of the natural gas hydration process with a NiMnGa micro-/nanofluid in a static suspension system

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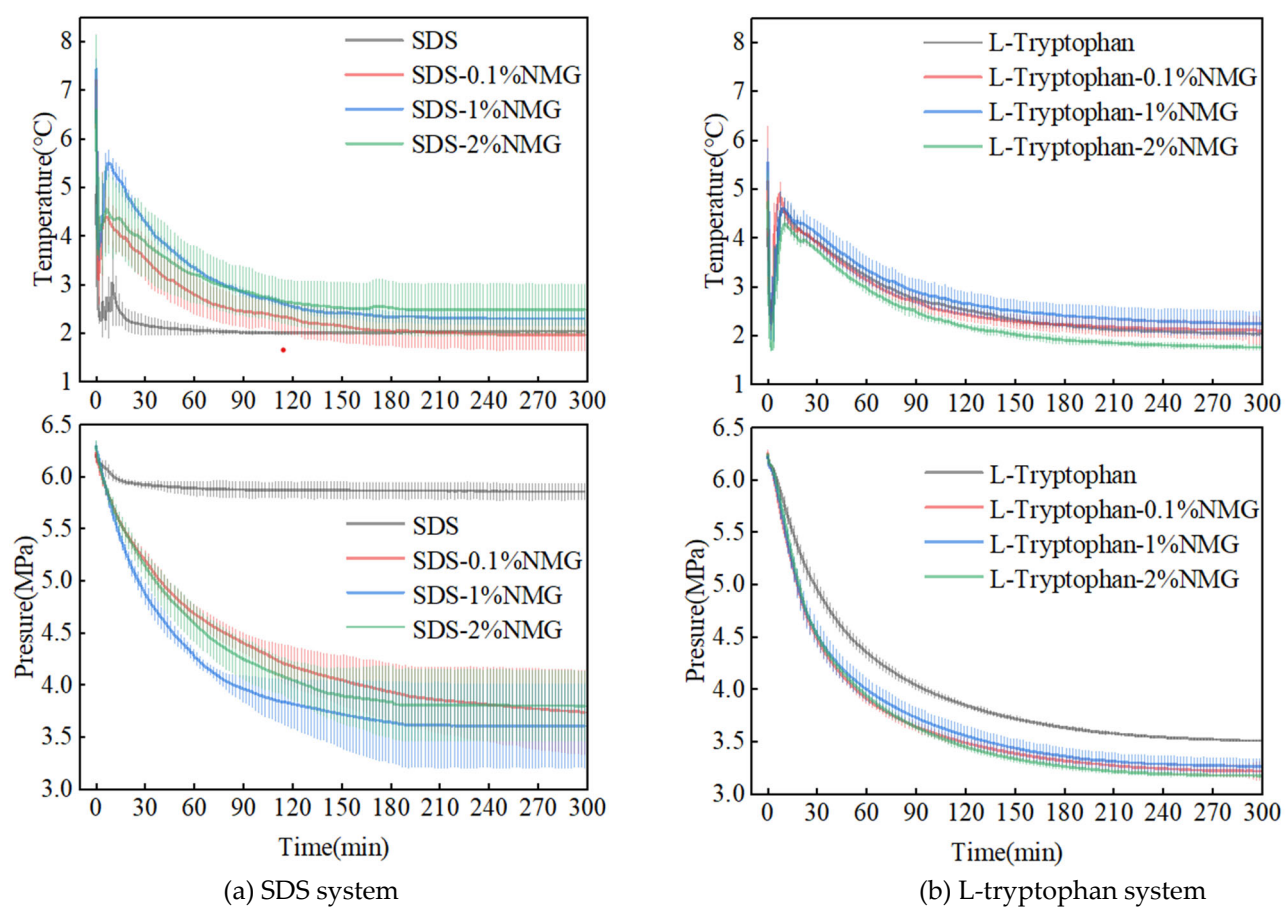
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**Table S1.** Experimental conditions and results of methane hydrate in different systems(P=6.2MPa , T=2°C).

Gas sample	System	Gellan gum /%	SDS /%	L-Tryptophan /%	NMG /%	No.	Induction time /min	Gas consumption	Gas consumption rate/mmol of gas/mol of water/min		
								ion/mmol of gas /mol of water	V <sub>15min</sub>	V <sub>avg</sub>	
99.99% CH <sub>4</sub>	I	0.3	0.05	0	0	1	3	2.763	0.175	0.125	
						2	6	5.105	0.215	0.083	
						3	8.83	2.673	0.151	0.062	
						4	1.83	30.969	0.699	0.167	
						0.1	5	2.67	32.458	0.593	0.116
							6	1.67	39.135	0.612	0.152
							7	1.17	43.797	0.896	0.238
						1	8	1.67	37.864	0.831	0.199
							9	1.87	30.612	0.951	0.177
							10	1.67	32.779	0.637	0.179
						2	11	1.33	40.090	0.707	0.220
							12	1.5	30.848	0.699	0.153
	II		LG	0	1	0.1	1	2.67	38.417	0.889	0.151
							2	3.67	38.718	0.775	0.153
							3	3.33	38.295	0.798	0.146
			LNG				4	2.50	41.948	1.071	0.198
							5	3.17	41.116	1.112	0.189
							6	2.67	42.165	1.241	0.180
			LNG				7	2.17	40.662	1.170	0.190
							8	3.50	41.221	1.024	0.190
							9	3.67	42.022	1.242	0.195
			LNG				10	2.83	43.114	1.190	0.210
							11	3.00	43.013	1.098	0.210
							12	3.00	42.853	1.075	0.198



**Figure S1.** Curve of temperature and pressure change with time in different systems.

## Experimental section

### Materials

(1) High-purity nickel lump, electrolytic manganese, and pure gallium metal were melted in the vacuum medium frequency arc melting furnace (SR830, Beijing Zhongke Keji Co., Ltd.) according to the ratio of Ni<sub>52.5</sub>Mn<sub>22.5</sub>Ga<sub>25</sub>. In order to make the alloy melt more uniform, the ingots were turned over and melted 6 times with electromagnetic stirring during the melting process, and finally, button-shaped ingots were obtained.

(2) The melted ingots were polished for oxidation and cut by a wire cutting machine (BDR7725, Beijing Dimeng Hengda Company). The cut ingots are homogenized in a heat treatment chamber resistance heating furnace (SX-4-10, Harbin E. Excel Heat Treatment Equipment Manufacturing Co., Ltd.), and the surface oxidized skin is polished after water cooling.

(3) The homogenized solid solution treated ingots were prepared into small particles of uniform size and ball milled in a high-speed vibrating ball mill (GB-80, Nanjing Boynton Instruments Technology Co., Ltd.) for 30 min–90 min. Various particle sizes could be prepared by controlling the ball milling time and finally annealed to produce the desired NiMnGa micro-nano particles.

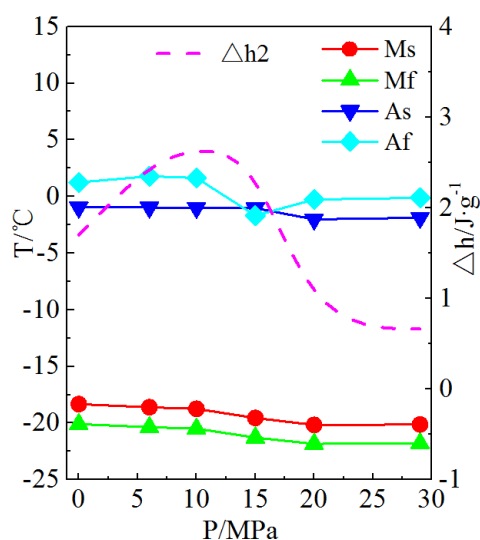
### Test Results

In order to grasp the degree of influence of different NMG and different pressure conditions on the phase change conditions, the parameters of phase change temperature, inverse phase change temperature, and their latent heat were measured. The results of  $\mu$ DSCVII high-pressure differential scanning calorimeter (HP DSC) tests are shown in Table S2. NMG phase change temperature and inverse phase change latent heat diagram are shown in Figure S2. From the test results, it can be

seen that the NMG inverse phase change temperature is 1.794°C in the pressure range of 6~8 MPa, and the latent heat of inverse phase change is 2.493J/g.

**Table S2.** The results of  $\mu$ DSCVII high-pressure differential scanning calorimeter (HP DSC) tests

NO.	Pressure /MPa	Ni <sub>52.5</sub> Mn <sub>22.5</sub> Ga <sub>25</sub>				
		M/g	M <sub>s</sub> ~M <sub>f</sub>	A <sub>s</sub> ~A <sub>f</sub>	$\Delta h_1$	$\Delta h_2$
1.	0	0.0440	-18.343~-20.1	-0.967~1.245	-0.94	1.7
2.	6	0.0485	-18.612~-20.378	-0.976~1.794	-0.814	2.493
3.	10	0.0475	-18.747~-20.506	-0.998~1.636	-0.859	2.666
4.	15	0.0554	-19.562~-21.3	-1.036~-1.685	-0.748	2.569
5.	20	0.0601	-20.171~-21.849	-2.014~-0.264	-0.614	0.684
6.	26	0.0615	-20.142~-21.805	-1.878~-0.111	-0.559	0.661



**Figure S2.** NMG phase change temperature and inverse phase change latent heat diagram

Where, M-phase change start temperature/°C

M<sub>f</sub>-end of phase change temperature/°C

A<sub>s</sub>-reverse phase change start temperature/°C

A<sub>f</sub>-the end of the reverse phase change temperature/°C

$\Delta h_1$ -the latent heat of phase change J/g

$\Delta h_2$ -inverse phase change latent heat J/g

M-sample mass/g