

MODELLING THE EVOLUTION OF NEGATIVE PRESSURE MEASURED DURING FLASHING OF WATER

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First results of experimental investigations confirming the possibility of obtaining negative pressures measured during flashing of water were published by Briggs¹ in 1950, then by Borkar et al.² in 1977. The results provide a testimony that the bulk viscosity of a two-phase system in the first stage of flashing can assume considerable values. The present paper is devoted to modeling of evolution of kinetic, or hydraulic pressure $P^h = \frac{1}{3} \text{tr} P_{ij}$, where P_{ij} is a pressure tensor. The kinetic pressure undergoes relaxation to the thermodynamic pressure P which is always positive.

It is shown that the equation to describe the evolution of P^h can be obtained based on Classical Irreversible Thermodynamics (CIT) in the form

$$P^h - P = L_{21}(\mu_l - \mu_g) + L_{22} \frac{\partial w_i}{\partial z_i} \quad (1)$$

where L_{ij} are the phenomenological coefficients, and L_{22} corresponds to the bulk viscosity

$L_{22} = -\zeta$, μ_l and μ_g are chemical potentials of the liquid and vapor respectively and $\frac{\partial w_i}{\partial z_i}$ denotes divergence of velocity. The equation (1) says that the generalized thermodynamic flux $P^v = P^h - P$ is effected by two thermodynamic forces, namely the difference of the chemical potentials $\mu_l - \mu_g$ and the relative rate of change of flow volume $v(t)$ which is

expressed by $\frac{\partial w_i}{\partial z_i}$. At the same time, another thermodynamic flux of the zero tensorial rank

$\frac{dx}{dt}$, where x stands for the dryness fraction, is also a function of the same forces and is expressed in the following form

$$\dot{x} \equiv \frac{dx}{dt} = L_{11}(\mu_l - \mu_g) + L_{12} \frac{\partial w_i}{\partial z_i} \quad (2)$$

A different form of the evolution equation for P^v can be derived from Extended Irreversible Thermodynamics (EIT), namely

$$\theta_p \frac{d(P^h - P)}{dt} + P^h - P = L_{21}(\mu_l - \mu_g) + L_{22} \frac{\partial w_i}{\partial z_i} \quad (3)$$

where θ_p means the relaxation time and $\frac{d}{dt}$ denotes the substantial derivative. Equation (3) can be conjugated with an appropriate equation describing the evolution of \dot{x} namely

$$\theta_x \frac{d\dot{x}}{dt} + \dot{x} = L_{11}(\mu_l - \mu_g) + L_{12} \frac{\partial w_i}{\partial z_i} \quad (4)$$

where θ_x means the relaxation time prescribed to rate of evaporation \dot{x} .

The third equation applied to describe the evolution of P^h is taken from a simple expansion of the equilibrium - thermodynamic pressure P

$$\frac{dP^h}{dt} = - \frac{P^h - P}{\theta_p} \quad (5)$$

However the paper also exhibits differences in evolution of P^h during flashing of water as found from these three approaches.

REFERENCES

1. Briggs, L.J., Limiting Negative Pressure of Water, *J. Appl. Phys.*, Vol. 21, pp. 721-722, 1950.
2. Borkar, G.S., Lienhard, J.H. & Trela, A Rapid Hot-Water Depressurization Experiment, *Report EPRI NP-527 Project RP687-1*, 1977.