

A unified framework for Navier–Stokes Cahn–Hilliard models with non-matching densities

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Over the last decades, many diffuse-interface Navier–Stokes Cahn–Hilliard (NSCH) models with non-matching densities have appeared in the literature. These models claim to describe the same physical phenomena, yet they are distinct from one another. The overarching objective of this work is to bring all of these models together by laying down a unified framework of NSCH models with non-zero mass fluxes. Our development is

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based on three unifying principles: (1) there is only one system of balance laws based on continuum mixture theory that describes the physical model, (2) there is only one natural energy-dissipation law that leads to quasi-incompressible NSCH models, (3) variations between the models only appear in the constitutive choices. The framework presented in this work now completes the fundamental exploration of alternate non-matching density NSCH models that utilize a single momentum equation for the mixture velocity, but leaves open room for further sophistication in the energy functional and constitutive dependence.

Keywords: Navier–Stokes Cahn–Hilliard equations; phase-field models; incompressible two-phase flow; mixture theory; thermodynamic consistency.

AMS Subject Classification: Primary: 76T99; Secondary: 35Q30, 35Q35, 35R35, 76D05, 76D45, 80A99

1. Introduction

Phase-field models have emerged as a powerful tool for describing interface problems that appear in various fields in science. These models are also termed diffuse-interface models due to the smooth representation of the interface.⁶ The theory of general diffuse-interface models in solid and fluid mechanics has been presented in Ref. 31. Phase-field models are typically equipped with a thermodynamical framework^{26, 28, 34, 40} and stability properties.¹⁴ A key factor in the success of phase-field models is their ability to be directly applied in computer simulations¹⁶ ranging from phase transitions in fluids²⁴ to fracture mechanics.⁷

In the context of free-surface fluid mechanics, other popular methodologies are the volume-of-fluid methods²¹ and level-set methods.^{32, 36} Level-set methods are popular for incompressible flows^{5, 10, 41} and volume-of-fluid methods are employed for both incompressible^{17, 33} and compressible flows.^{30, 35, 42} The main distinguishing feature of a phase-field model, as compared with volume-of-fluid and level-set models, is that the interface is determined by a physical model with a fixed interface width. In computations with the volume-of-fluid approach often an interpolation technique or a compression algorithm is applied near the interface, whereas level-set methods typically use a redistancing algorithm to maintain a stable interface. Apart from the thermodynamical structure of the underlying phase-field formulations, redundancy of such algorithms is often the most important practical advantage of phase-field simulations.

The phase-field model that describes incompressible isothermal two constituent flows with non-matching densities is the Navier–Stokes Cahn–Hilliard (NSCH) model. Over the years many NSCH models with non-matching densities have been proposed with distinct fundamental variables for the velocity and phase-field. It is the purpose of this paper to present a unified framework for non-matching density NSCH models which is invariant to the choice of fundamental variables. The framework encompasses variations of existing NSCH models.

1.1. *Historical overview*

The first coupling between the Navier–Stokes equations, describing viscous fluid flow, and the Cahn–Hilliard equation, describing spinoidal decomposition, has been

established by Hohenberg and Halperin.²² They proposed the system, referred to as *model H*, that reads

$$\rho \partial_t \mathbf{u} + \rho (\mathbf{u} \cdot \nabla) \mathbf{u} - \operatorname{div}(2\nu(c)\mathbf{D}) + \nabla p = -\sigma \epsilon \operatorname{div}(\nabla c \otimes \nabla c), \quad (1.1a)$$

$$\operatorname{div} \mathbf{u} = 0, \quad (1.1b)$$

$$\partial_t c + \mathbf{u} \cdot \nabla c = \operatorname{div}(m(c)\nabla \mu), \quad (1.1c)$$

$$\mu = \sigma \epsilon^{-1} \varphi'(c) - \sigma \epsilon \Delta c, \quad (1.1d)$$

in domain $\Omega \subset \mathbb{R}^d$, with dimension $d = 2, 3$, both open and bounded that is occupied with two constituents $j = 1, 2$. We adopt the standard notation, where \mathbf{x} represents the (Eulerian) position vector, t is the time, ∂_t the partial time-derivative, ∇ the gradient, div the divergence (defined as $(\operatorname{div} \mathbf{A})_i := \partial A_{ik} / \partial x_k$) and Δ the Laplace operator. Here, \mathbf{u} is the mean velocity, ρ is the density, p is the pressure, c is the so-called order parameter representing a concentration related quantity. Moreover, \mathbf{D} represents the symmetric gradient of the mean velocity, $\mathbf{D} = \frac{1}{2}(\nabla \mathbf{u} + (\nabla \mathbf{u})^T)$, $\nu(c)$ is the concentration-dependent dynamic viscosity of the mixture, σ is the surface tension coefficient and ϵ is an interface thickness parameter. The surface tension coefficient is assumed constant, i.e. Marangoni-type effects are precluded in this model. The quantity $\varphi = \varphi(c)$ is the homogeneous free energy and $m = m(c) \geq 0$ is the mobility. In model (1.1), Eq. (1.1a) describes the linear momentum equation in which the right-hand side, $-\sigma \epsilon \operatorname{div}(\nabla c \otimes \nabla c)$, represents a contribution of the stress tensor that models capillary forces due to surface tension. Equation (1.1b) dictates that the mean velocity is divergence-free. Lastly, the Cahn–Hilliard equation (1.1c) describes the evolution of the order parameter in which the right-hand side represents the divergence of a diffusive flux, and the variable μ given by (1.1d) is often referred to as the chemical potential.

The major limitation of model *H* is its assumption of constant density, i.e. the density of the mixture as well as the density of the individual constituents is constant. As such, this precludes the applicability of the model to problems with large density ratios (e.g. water–air flows). Model *H* has been used to study the fluid behavior at critical points of single and binary fluids. The derivation of the model initially relied on phenomenological arguments and a rigorous derivation in the framework of rational continuum mechanics was absent until 1996.

In that year Gurtin²⁰ provided this missing derivation. The core ideas in this derivation are (i) the separation of balance laws from constitutive relations, (ii) the usage of microforce balance laws, and (iii) the selection of constitutive equations compatible with an energy-dissipation law referred to as the second law of thermodynamics (the so-called Coleman–Noll procedure). Gurtin introduces the balance of mass per constituent while he directly presents a single momentum equation for the mixture. In the same year Gurtin applied this procedure to present the Ginzburg–Landau and Cahn–Hilliard equations with a derivation in a rational continuum mechanics framework.¹⁹ Several extensions of the microforce continuum framework

of Gurtin have been proposed, including a phase-field gradient theory for enriched continua.¹⁵

One of the first efforts of extending model H to the case of non-matching densities is the work of Lowengrub and Truskinovsky.²⁵ Lowengrub and Truskinovsky present the following *quasi-incompressible model*:

$$\left\{ \begin{array}{l} \partial_t(\rho \mathbf{v}) + \operatorname{div}(\rho \mathbf{v} \otimes \mathbf{v}) + \nabla p - \operatorname{div}(\nu(c)(2\mathbf{D} + \lambda(c)(\operatorname{div} \mathbf{v})\mathbf{I})) \\ \quad + \sigma \epsilon \operatorname{div}(\rho \nabla c \otimes \nabla c) = 0, \\ \partial_t \rho + \operatorname{div}(\rho \mathbf{v}) = 0, \\ \rho(\partial_t c + \mathbf{v} \cdot \nabla c) - \operatorname{div}(m \nabla \mu) = 0, \\ \mu - \frac{\sigma}{\epsilon} \frac{\partial \varphi}{\partial c} + \frac{\sigma \epsilon}{\rho} \operatorname{div}(\rho \nabla c) + \frac{p}{\rho^2} \frac{\partial \rho}{\partial c} = 0, \end{array} \right. \quad \begin{array}{l} (1.2a) \\ (1.2b) \\ (1.2c) \\ (1.2d) \end{array}$$

with free energy density per unit volume $\rho \sigma (\epsilon^{-1} \varphi(c) + \epsilon |\nabla c|^2/2)$ and $m \geq 0$ a non-degenerate mobility (the mobility is called *non-degenerate* if it is constant and *degenerate* if it vanishes in the single-fluid regime). The *mixture density* ρ is $\rho = \tilde{\rho}_1 + \tilde{\rho}_2$ and the *mass-averaged velocity* \mathbf{v} is given by $\rho \mathbf{v} = \tilde{\rho}_1 \mathbf{v}_1 + \tilde{\rho}_2 \mathbf{v}_2$ where $\tilde{\rho}_j = \rho c_j$ represents the variable density of constituent j with c_j the concentration of constituent j . In comparison with model H , this model has two key differences. First, the incompressibility constraint (1.1b) is replaced by the conservation of mass of the mixture (1.2b). Expanding the divergence in (1.2b) reveals the quasi-incompressible nature of the model:

$$\operatorname{div} \mathbf{v} + \frac{1}{\rho} \frac{\partial \rho}{\partial c} (\partial_t c + \mathbf{v} \cdot \nabla c) = 0. \quad (1.3)$$

Second, the evolution equation of the order parameter (1.2c) explicitly contains a density which distinguishes it from the evolution equation (1.1c). It seems challenging to design algorithms that solve (1.2) and often a simplified version of the model is employed, see e.g. Ref. 23. The challenge is often linked to the quasi-incompressible nature of the model.³ We note that recently a numerical method for a reformulation of the model (1.2) has been proposed in Ref. 18.

Noteworthy alternative models have been proposed by Boyer⁹ and Ding *et al.*¹² Both models deviate from the model of Lowengrub and Truskinovsky²⁵ in that they use a mean velocity that is fully incompressible:

$$\operatorname{div} \mathbf{u} = 0. \quad (1.4)$$

This mean velocity, defined as $\mathbf{u} = \phi_1 \mathbf{v}_1 + \phi_2 \mathbf{v}_2$ where ϕ_j is the volume fraction of constituent j , is referred to as the *volume-averaged velocity*. Actually the model¹² is, apart from the variable density, identical to the model H (model (1.1)). Analogously to Gurtin²⁰ and Lowengrub and Truskinovsky,²⁵ the starting point of Ding *et al.*¹² consists of the individual mass balance equations of the constituents and the momentum equation of the mixture. In contrast, the point of departure adopted

in the model of Boyer⁹ is composed of individual mass and momentum balance equations, which leads to a different form of the momentum and phase equations.

Unfortunately the models Refs. 9 and 12 are not presented with some (approximate) form of the second law of thermodynamics. This observation has led to the development of the model of Abels *et al.*³ which reads

$$\left\{ \begin{array}{l} \partial_t(\rho \mathbf{u}) + \operatorname{div}(\rho \mathbf{u} \otimes \mathbf{u}) + \operatorname{div} \left(\mathbf{u} \otimes \frac{\rho_1 - \rho_2}{2} m(\phi) \nabla \mu \right) + \nabla p \\ \quad - \operatorname{div}(2\nu(\phi) \mathbf{D}) + \sigma \epsilon \operatorname{div}(\nabla \phi \otimes \nabla \phi) = 0, \\ \operatorname{div} \mathbf{u} = 0, \\ \partial_t \phi + \mathbf{u} \cdot \nabla \phi - \operatorname{div}(m(\phi) \nabla \mu) = 0, \\ \mu - \frac{\sigma}{\epsilon} \frac{\partial \varphi}{\partial \phi} + \sigma \epsilon \Delta \phi = 0, \end{array} \right. \quad \begin{array}{l} (1.5a) \\ (1.5b) \\ (1.5c) \\ (1.5d) \end{array}$$

and is presented with some form of the second law of thermodynamics. This form of the second law is distinct from what appears in Lowengrub and Truskinovsky²⁵ in the sense that a different kinetic energy is used. This model also uses the volume-averaged mean velocity for the velocity of the mixture and makes use of the difference of volume fractions as order parameter. The distinguishing feature of (1.5) is the, somehow surprising, additional convective term in the momentum equation.

Since then several quasi-incompressible models that employ the difference of volume fractions as order parameter have been proposed. Noteworthy contributions are the work of Shen *et al.*,³⁷ Aki *et al.*⁴ and Shokrpour Roudbari *et al.*,³⁸ which are all presented with an approximate form of the second law. The models Refs. 4 and 38 have been derived in a similar way, namely, using balance laws of the individual constituents and the Coleman–Noll procedure. The derivation of the model of Shen *et al.* follows different considerations. An important contribution in the work of Shokrpour Roudbari *et al.*³⁸ is the observation that, up to the definition of the mobility and the definition of mass fluxes, the models^{4, 37, 38} are all equivalent.

1.2. Objective and main results

It is obvious that the literature on NSCH models is divided. The various proposed models (almost) all aim to represent the same physical behavior but are either clearly different or are presented in a (sometimes seemingly) different form. The four key flavors are

- (1) the mixture velocity: mass-averaged velocity (\mathbf{v}) or a volume-averaged velocity (\mathbf{u})
- (2) the order parameter: based on volume fractions (ϕ) or concentration (c)

Table 1. Overview of various NSCH models. The columns indicate the mean velocity (either mass-averaged \mathbf{v} or volume-averaged \mathbf{u}), the order parameter (either volume fraction based ϕ or concentration based c), the type of the free energy (either volume-measure-based Ψ or mass-measure-based ψ), the type of the mobility (either degenerate or non-degenerate) and lastly whether the model is equipped with an energy-dissipation law. The model of Abels *et al.* is presented with both non-degenerate and degenerate mobilities. Additionally, the energy-dissipation law of that model differs from that of the other models, due to the usage of the volume-averaged velocity, and we indicate this by the symbol \star .

Model	Velocity	Order parameter	Free energy	Mobility	Energy law
Abels <i>et al.</i> ³	\mathbf{u}	ϕ	Ψ	non-deg./degen.	\star
Aki <i>et al.</i> ⁴	\mathbf{v}	ϕ	Ψ	non-deg.	\checkmark
Boyer ⁹	\mathbf{u}	ϕ	Ψ	degen.	\times
Ding <i>et al.</i> ¹²	\mathbf{u}	ϕ	Ψ	degen.	\times
Lowengrub and Truskinovsky ²⁵	\mathbf{v}	c	ψ	non-deg.	\checkmark
Shen <i>et al.</i> ³⁷	\mathbf{v}	ϕ	Ψ	non-deg.	\checkmark
Shokrpour Roudbari <i>et al.</i> ³⁸	\mathbf{v}	ϕ	Ψ	non-deg.	\checkmark

- (3) the type of the free energy: volume-measure-based (Ψ) or a mass-measure-based (ψ)
- (4) the mobility: non-degenerate (i.e. constant) or degenerate (i.e. it vanishes in the single-fluid regime)

We provide a summary of several existing models in Table 1.

Even though each of these works has provided new and useful insights into diffuse-interface modeling, and some provide elegant and physically sound derivations, the current status is far from optimal. We have two main objections:

- (1) the systems of balance laws of the various models are distinct before constitutive choices have been applied,
- (2) the energy-dissipation laws of the models are not identical.

Since all the models represent the same physics, we lay down the three unifying principles:

- (1) there is only one system of continuum mechanics balance laws that describes the physical model,
- (2) there is only one natural energy-dissipation law that leads to quasi-incompressible NSCH models,
- (3) variations between the models only appear in the constitutive choices.

The main objective of this work is to lay down a *unified framework of incompressible NSCH models with non-zero mass fluxes* on the basis of these three unifying principles. In particular, we establish one incompressible NSCH system of balance laws and show that many alternate forms are connected via variable transformations. Two (equivalent) formulations that result after the constitutive choices are (i) a formulation in terms of the mass-averaged velocity \mathbf{v} :

$$\left\{ \begin{array}{l} \partial_t(\rho \mathbf{v}) + \operatorname{div}(\rho \mathbf{v} \otimes \mathbf{v}) + \nabla p + \operatorname{div} \left(\nabla \phi \otimes \frac{\partial \Psi}{\partial \nabla \phi} + (\mu \phi - \Psi) \mathbf{I} \right) \\ \quad - \operatorname{div}(\nu(2\mathbf{D} + \lambda(\operatorname{div} \mathbf{v})\mathbf{I})) - \rho \mathbf{g} = 0, \end{array} \right. \quad (1.6a)$$

$$\partial_t \rho + \operatorname{div}(\rho \mathbf{v}) = 0, \quad (1.6b)$$

$$\partial_t \phi + \operatorname{div}(\phi \mathbf{v}) - \operatorname{div}(\mathbf{M}^v \nabla(\mu + \alpha p)) + \zeta m(\mu + \alpha p) = 0, \quad (1.6c)$$

$$\left\{ \begin{array}{l} \mu - \frac{\partial \Psi}{\partial \phi} + \operatorname{div} \left(\frac{\partial \Psi}{\partial \nabla \phi} \right) = 0, \end{array} \right. \quad (1.6d)$$

and (ii) a formulation in terms of the volume-averaged velocity \mathbf{u} :

$$\left\{ \begin{array}{l} \partial_t(\rho \mathbf{u} + \tilde{\mathbf{J}}^u) + \operatorname{div} \left(\rho \mathbf{u} \otimes \mathbf{u} + \tilde{\mathbf{J}}^u \otimes \mathbf{u} + \mathbf{u} \otimes \tilde{\mathbf{J}}^u + \frac{1}{\rho} \tilde{\mathbf{J}}^u \otimes \tilde{\mathbf{J}}^u \right) \\ \quad + \nabla p + \operatorname{div} \left(\nabla \phi \otimes \frac{\partial \Psi}{\partial \nabla \phi} + (\mu \phi - \Psi) \mathbf{I} \right) \end{array} \right. \quad (1.7a)$$

$$- \operatorname{div}(\nu(2\nabla^s(\mathbf{u} + \rho^{-1}\tilde{\mathbf{J}}^u) + \lambda(\operatorname{div}(\mathbf{u} + \rho^{-1}\tilde{\mathbf{J}}^u))\mathbf{I})) - \rho \mathbf{g} = 0, \quad (1.7b)$$

$$\operatorname{div} \mathbf{u} - \beta \gamma = 0, \quad (1.7c)$$

$$\partial_t \phi + \mathbf{u} \cdot \nabla \phi - \operatorname{div}(\mathbf{M}^u \nabla(\mu + \alpha p)) + \frac{\rho}{2\rho_1\rho_2} m(\mu + \alpha p) = 0, \quad (1.7c)$$

$$\left\{ \begin{array}{l} \mu - \frac{\partial \Psi}{\partial \phi} + \operatorname{div} \left(\frac{\partial \Psi}{\partial \nabla \phi} \right) = 0, \end{array} \right. \quad (1.7d)$$

with $\tilde{\mathbf{J}}^u = -(\rho_1 - \rho_2)\mathbf{M}^u \nabla(\mu + \alpha p)/2$. Here, $\mathbf{M}^v = \mathbf{M}^v(\phi, \nabla \phi, \mu, \nabla \mu, p)$, $\mathbf{M}^u(\phi, \nabla \phi, \mu, \nabla \mu, p)$ and $m = m(\phi, \mu, p)$ are degenerate mobilities, $\nu = \nu(\phi)$ is the dynamic viscosity, p is the pressure, \mathbf{g} is the gravitational acceleration, ρ_1 and ρ_2 are the constant specific densities of the constituents and we have introduced the constants $\alpha = (\rho_2 - \rho_1)/(\rho_1 + \rho_2)$, $\beta = (\rho_2 - \rho_1)/(2\rho_1\rho_2)$ and $\zeta = (\rho_1 + \rho_2)/(2\rho_1\rho_2)$. We refer the reader for details on the specific choices and scaling of the mobility parameters, and free energy to Refs. 3, 4 and 20. Moreover, we show that (i) many existing models are identical (up to the definition of the mobility), (ii) existing volume-averaged velocity based models are inconsistent with the mixture theory framework, however have a consistent rectification. As a side observation it turns out that, within our framework, models with a non-degenerate mobility are incompatible in the single-fluid region (see Remark 3.12).

1.3. Plan of the paper

The structure of the remainder of the paper is as follows. In Sec. 2, the system of balance laws is established. We present the framework of balance laws for binary mixtures of incompressible viscous fluids. Starting from constituent balance laws, we use mixture theory to derive the balance laws of the mixtures. Additionally, we present some new and important identities and evolution equations. In Sec. 3, we perform constitutive modeling via the Coleman–Noll procedure. Here we highlight the modeling assumptions of the NSCH model from the viewpoint of mixture theory. Additionally, we show that applying the Coleman–Noll procedure to alternative derivations provides the same modeling restriction. In Sec. 4, we discuss the relation of the novel model to existing NSCH models. Finally, in Sec. 5, we summarize our findings and present some possible further research directions.

2. Mixture Theory

In this section, we lay down the mixture theory as well as the necessary definitions. In this work, we focus on incompressible isothermal constituents, which we will specify in Sec. 2.1. We restrict ourselves to the case of binary mixtures for the sake of simplicity and note that the multi-component case is a straightforward extension. This section is fully compatible (i.e. no approximations are introduced) with the mixture theory metaphysical principles proposed by Truesdell⁴³:

- (1) All properties of the mixture must be mathematical consequences of properties of the constituents.
- (2) So as to describe the motion of a constituent, we may in imagination isolate it from the rest of the mixture, provided we allow properly for the actions of the other constituents upon it.
- (3) The motion of the mixture is governed by the same equations as in a single body.

Section 2.1 provides the necessary (kinematic) definitions. Sections 2.2 and 2.3, we introduce the balance laws of the individual constituents and subsequently, relying on metaphysical principles of Ref. 44, balance laws of the mixtures. Finally, in Sec. 2.4, we present alternative, but equivalent, formulations of the balance laws, diffusive fluxes and stress tensors.

2.1. Preliminaries

In domain Ω , we denote with \mathbf{X}_j the position of a particle of constituent j in the Lagrangian (reference) configuration. Denoting the position of the mixture in the Eulerian frame by \mathbf{x} , we identify its relation to the initial configurations of the constituents with an invertible deformation map χ_j :

$$\mathbf{x} := \chi_j(\mathbf{X}_j, t). \quad (2.1)$$

The velocity of constituent j is given by

$$\mathbf{v}_j(\mathbf{x}, t) := \partial_t \chi_j(\mathbf{X}_j, t) = \partial_t \chi_j(\chi_j^{-1}(\mathbf{x}, t), t). \quad (2.2)$$

Consider now an arbitrary control volume $V \subset \Omega$ around spatial position \mathbf{x} in the mixture that contains the masses $M_j = M_j(V)$ of the constituents $j = 1, 2$ at time t . We denote the total mass in V as $M = M(V) = \sum_j M_j(V)$. We define the partial mass density $\tilde{\rho}_j$ as the mass of constituent j per unit volume of the mixture as

$$\tilde{\rho}_j(\mathbf{x}, t) := \lim_{|V| \rightarrow 0} \frac{M_j(V)}{|V|}, \quad (2.3)$$

where $|V|$ denotes the measure of control volume V . The mass density of the total mixture at position \mathbf{x} and time t is now defined as the sum of the partial mass densities of the constituents:

$$\rho(\mathbf{x}, t) := \sum_j \tilde{\rho}_j(\mathbf{x}, t). \quad (2.4)$$

Next, we introduce $V_j \subset V$ as the control volume occupied by constituent j . We define the volume fraction of constituent j as

$$\phi_j(\mathbf{x}, t) := \lim_{|V| \rightarrow 0} \frac{|V_j|}{|V|}, \quad (2.5)$$

where $|V_j|$ denotes the measure of V_j . We assume that

$$\sum_j \phi_j = 1, \quad (2.6)$$

and thus exclude the existence of interstitial voids. Additionally, we introduce the concentration of constituent j as

$$c_j(\mathbf{x}, t) := \lim_{|V| \rightarrow 0} \frac{M_j(V)}{M(V)}. \quad (2.7)$$

The above definitions (2.3), (2.4) and (2.7) imply the relation:

$$\tilde{\rho}_j(\mathbf{x}, t) = \rho(\mathbf{x}, t) c_j(\mathbf{x}, t). \quad (2.8)$$

Next, we define the specific mass density ρ_j as the mass of constituent j per volume occupied by that constituent as

$$\rho_j(\mathbf{x}, t) := \lim_{|V_j| \rightarrow 0} \frac{M_j(V)}{|V_j|}. \quad (2.9)$$

We assume $\rho_j(\mathbf{x}, t) > 0$. From the definitions (2.3), (2.5) and (2.9) we deduce

$$\tilde{\rho}_j(\mathbf{x}, t) = \rho_j(\mathbf{x}, t) \phi_j(\mathbf{x}, t). \quad (2.10)$$

Additionally, we have the relation:

$$\sum_j c_j = 1. \quad (2.11)$$

In general the specific mass densities ρ_j may vary due to compressibility and thermal effects. In this paper, we restrict to constant specific mass densities ρ_j representing incompressible isothermal constituents. The momentum associated with constituent j is:

$$\mathbf{m}_j(\mathbf{x}, t) := \tilde{\rho}_j(\mathbf{x}, t) \mathbf{v}_j(\mathbf{x}, t). \quad (2.12)$$

The momentum of the mixture is the sum of that of the individual constituents:

$$\mathbf{m}(\mathbf{x}, t) := \sum_j \mathbf{m}_j(\mathbf{x}, t). \quad (2.13)$$

The *mixture velocity* \mathbf{v} is a *mass-averaged velocity* or *barycentric velocity* and is identified via the relation:

$$\mathbf{m}(\mathbf{x}, t) = \rho(\mathbf{x}, t) \mathbf{v}(\mathbf{x}, t). \quad (2.14)$$

The *diffusion velocity* or *peculiar velocity* of constituent j is the constituent velocity relative to the gross motion of the mixture:

$$\mathbf{w}_j(\mathbf{x}, t) := \mathbf{v}_j(\mathbf{x}, t) - \mathbf{v}(\mathbf{x}, t). \quad (2.15)$$

An immediate consequence is the observation that the superposition of the momenta relative to the gross motion of the mixture vanishes:

$$\sum_j \tilde{\rho}_j \mathbf{w}_j = \sum_j \tilde{\rho}_j \mathbf{v}_j - \sum_j \tilde{\rho}_j \mathbf{v} = 0. \quad (2.16)$$

From (2.16), we deduce the identity:

$$\sum_j c_j \mathbf{w}_j = 0, \quad (2.17)$$

that will be employed later in this section. We now introduce two different material derivatives, one that follows the individual motion of constituent j and one that follows the mean motion, respectively, given by

$$\dot{\psi} = \partial_t \psi + \mathbf{v}_j \cdot \nabla \psi, \quad (2.18a)$$

$$\dot{\psi} = \partial_t \psi + \mathbf{v} \cdot \nabla \psi. \quad (2.18b)$$

We define the jump and average of a constituent-related (vector-valued) quantity as

$$\llbracket \mathbf{w} \rrbracket := \frac{1}{2}(\mathbf{w}_1 - \mathbf{w}_2), \quad (2.19a)$$

$$\{\mathbf{w}\} := \frac{1}{2}(\mathbf{w}_1 + \mathbf{w}_2), \quad (2.19b)$$

respectively, where the subscripts refer to the constituent numbers. Lastly, we introduce the constants:

$$\alpha := \frac{\rho_2 - \rho_1}{\rho_1 + \rho_2} = -\frac{[\![\rho]\!]}{\{\rho\}}, \quad (2.20a)$$

$$\beta := \frac{\rho_2 - \rho_1}{2\rho_1\rho_2} = -\frac{[\![\rho]\!]}{\rho_1\rho_2}, \quad (2.20b)$$

$$\zeta := \frac{\rho_1 + \rho_2}{2\rho_1\rho_2} = \frac{\{\rho\}}{\rho_1\rho_2}. \quad (2.20c)$$

2.2. Balance laws of single constituents

Denoting by γ_j the mass supply of constituent j due to reaction, we introduce the local evolution equation of the mass of constituent j :

$$\partial_t \tilde{\rho}_j + \operatorname{div}(\tilde{\rho}_j \mathbf{v}_j) = \gamma_j. \quad (2.21)$$

The associated convective form is

$$\dot{\tilde{\rho}}_j + \tilde{\rho}_j \operatorname{div} \mathbf{v}_j = \gamma_j. \quad (2.22)$$

We assume that mass fluxes γ_j vanish in the single fluid region (i.e. when $\phi_j = \pm 1$). As a consequence, the mass balance in the single fluid region reads:

$$\partial_t \rho_j + \operatorname{div}(\rho_j \mathbf{v}_j) = 0, \quad \text{for } \phi_j = 1. \quad (2.23)$$

Since the specific constituent densities ρ_j are constant we deduce the incompressible flow constraint:

$$\operatorname{div} \mathbf{v}_j = 0, \quad \text{for } \phi_j = 1. \quad (2.24)$$

The linear momentum of constituent j satisfies the balance law:

$$\partial_t \mathbf{m}_j + \operatorname{div}(\mathbf{m}_j \otimes \mathbf{v}_j) = \operatorname{div} \mathbf{T}_j + \tilde{\rho}_j \mathbf{b}_j + \boldsymbol{\pi}_j + \mathbf{v}_j \gamma_j. \quad (2.25)$$

Here, \mathbf{T}_j is the Cauchy stress tensor of constituent j and \mathbf{b}_j is the external body force. The term $\boldsymbol{\pi}_j$ represents the momentum supply of constituent j by the other constituents (see e.g. Ref. 44). The balance of angular momentum implies that the Cauchy stress tensors of constituents have the following form:

$$\mathbf{N}_j = \mathbf{T}_j - \mathbf{T}_j^T, \quad (2.26)$$

where \mathbf{N}_j represents the intrinsic moment of momentum vector. Note that the last member on the right-hand side of (2.25) vanishes when switching to a convective form with the aid of the local mass balance (2.21):

$$\tilde{\rho}_j \dot{\mathbf{v}}_j = \operatorname{div} \mathbf{T}_j + \tilde{\rho}_j \mathbf{b}_j + \boldsymbol{\pi}_j. \quad (2.27)$$

2.3. Balance laws of mixtures

The balance of mass of the mixture density follows by summing (2.21) over the constituents:

$$\partial_t \rho + \operatorname{div}(\rho \mathbf{v}) = 0, \quad (2.28)$$

where we have postulated the sum of the mass fluxes to vanish:

$$\sum_j \gamma_j = 0. \quad (2.29)$$

The mixture velocity \mathbf{v} is in general not divergence-free and this property is in literature referred to as a *quasi-incompressible* mixture.²⁵ The postulate (2.29) complies with the third metaphysical principle of mixture theory and precludes the creation of mixture mass. We denote the difference of the mass fluxes by $\gamma = \gamma_1 - \gamma_2$, which provides:

$$\gamma_1 = \frac{1}{2}\gamma, \quad \gamma_2 = -\frac{1}{2}\gamma. \quad (2.30)$$

To proceed, we introduce the order parameters (phase-fields) based on the volume fractions and concentrations of the individual constituents: $\phi = \phi_1 - \phi_2 \in [-1, 1]$ and $c = c_1 - c_2 \in [-1, 1]$. By recalling (2.6) and (2.11), we deduce the relations:

$$\phi_1 = \frac{1 + \phi}{2}, \quad \phi_2 = \frac{1 - \phi}{2}, \quad (2.31a)$$

$$c_1 = \frac{1 + c}{2}, \quad c_2 = \frac{1 - c}{2}. \quad (2.31b)$$

Additionally, the density of the mixture ρ can be expressed in terms ϕ and c via $\rho = \hat{\rho}(\phi) = \check{\rho}(c)$ where $\hat{\rho}(\phi)$ and $\check{\rho}(c)$ are defined as

$$\hat{\rho}(\phi) = \rho_1 \frac{1 + \phi}{2} + \rho_2 \frac{1 - \phi}{2}, \quad (2.32a)$$

$$\frac{1}{\check{\rho}(c)} = \frac{1}{\rho_1} \frac{1 + c}{2} + \frac{1}{\rho_2} \frac{1 - c}{2}. \quad (2.32b)$$

The relation between ϕ and c is given by

$$c = \frac{[\rho] + \{\rho\}\phi}{\{\rho\} + [\rho]\phi}, \quad \phi = \frac{-[\rho] + \{\rho\}c}{\{\rho\} - [\rho]c}, \quad c'(\phi) = \frac{\rho_1 \rho_2}{\rho^2}, \quad \phi'(c) = \frac{\rho^2}{\rho_1 \rho_2}. \quad (2.33)$$

The *diffusive fluxes* are defined as

$$\mathbf{h}^v := \phi_1 \mathbf{w}_1 - \phi_2 \mathbf{w}_2, \quad (2.34a)$$

$$\mathbf{J}^v := \tilde{\rho}_1 \mathbf{w}_1 - \tilde{\rho}_2 \mathbf{w}_2. \quad (2.34b)$$

The evolution equations of the order parameters ϕ and c follow from taking the difference of the mass balance equations of the constituents (2.21):

$$\dot{\phi} + \phi \operatorname{div} \mathbf{v} + \operatorname{div} \mathbf{h}^v = \zeta \gamma, \quad (2.35a)$$

$$\rho \dot{c} + \operatorname{div} \mathbf{J}^v = \gamma, \quad (2.35b)$$

where we recall that the dot $\dot{}$ denotes the material derivative with respect to the mixture velocity \mathbf{v} . Next, to obtain the linear momentum equation of the mixture we take the sum of (2.25):

$$\partial_t \mathbf{m} + \operatorname{div}(\mathbf{m} \otimes \mathbf{v}) = \operatorname{div} \mathbf{T} + \rho \mathbf{b}, \quad (2.36)$$

where the Cauchy stress tensor and the body force of the mixture are, respectively, identified as

$$\mathbf{T} := \sum_j \mathbf{T}_j - \tilde{\rho}_j \mathbf{w}_j \otimes \mathbf{w}_j, \quad (2.37a)$$

$$\rho \mathbf{b} := \sum_j \tilde{\rho}_j \mathbf{b}_j. \quad (2.37b)$$

Here, we have postulated the balance of momentum supplies:

$$\sum_j \pi_j + \gamma_j \mathbf{v}_j = 0, \quad (2.38)$$

which is consistent with the third metaphysical principle of mixture theory and precludes the creation of mixture momentum. We denote the difference of the growth of linear momentum of the constituents by $\mathbf{p} := (\pi_1 + \gamma_1 \mathbf{v}_1) - (\pi_2 + \gamma_2 \mathbf{v}_2)$, which leads to:

$$\pi_1 + \gamma_1 \mathbf{v}_1 = \frac{1}{2} \mathbf{p}, \quad \pi_2 + \gamma_2 \mathbf{v}_2 = -\frac{1}{2} \mathbf{p}. \quad (2.39)$$

Symmetry of the dyadic product implies symmetry of the Cauchy stress tensor of the mixture:

$$\mathbf{T}^T = \mathbf{T}, \quad (2.40)$$

where we have postulated the balance of the intrinsic moment of momentum vectors:

$$\sum_j \mathbf{N}_j = 0. \quad (2.41)$$

The corresponding convective form reads:

$$\rho \dot{\mathbf{v}} = \operatorname{div} \mathbf{T} + \rho \mathbf{b}. \quad (2.42)$$

2.4. Non-typical identities and evolution equations

Apart from the mixture velocity (2.14) one might define other mean velocities. A typical example of another mean velocity is the *volume-averaged velocity* \mathbf{u} given by

$$\mathbf{u}(\mathbf{x}, t) := \sum_j \phi_j(\mathbf{x}, t) \mathbf{v}_j(\mathbf{x}, t). \quad (2.43)$$

We define the peculiar velocity relative to the volume-averaged velocity:

$$\boldsymbol{\omega}_j(\mathbf{x}, t) := \mathbf{v}_j(\mathbf{x}, t) - \mathbf{u}(\mathbf{x}, t), \quad (2.44)$$

which has the consequence:

$$\sum_j \phi_j(\mathbf{x}, t) \boldsymbol{\omega}_j(\mathbf{x}, t) = 0. \quad (2.45)$$

A direct consequence of the constituent mass balance equations (2.21) and (2.45) is the relation:

$$\operatorname{div} \mathbf{u} = \beta \gamma. \quad (2.46)$$

Thus, in absence of mass fluxes ($\gamma = 0$), the volume-averaged velocity inherits the single-constituent solenoidal property of the individual constituents. We additionally introduce the diffusive fluxes with respect to the volume-averaged velocity:

$$\mathbf{h}^u := \phi_1 \boldsymbol{\omega}_1 - \phi_2 \boldsymbol{\omega}_2, \quad (2.47a)$$

$$\mathbf{J}^u := \tilde{\rho}_1 \boldsymbol{\omega}_1 - \tilde{\rho}_2 \boldsymbol{\omega}_2, \quad (2.47b)$$

$$\tilde{\mathbf{J}}^u := \tilde{\rho}_1 \boldsymbol{\omega}_1 + \tilde{\rho}_2 \boldsymbol{\omega}_2. \quad (2.47c)$$

Substitution of the mixture velocity (2.14) and the volume-averaged velocity (2.43) into (2.47c) provides the key identity that reveals the difference between the momentum \mathbf{m} and $\rho \mathbf{u}$:

$$\mathbf{m} = \rho \mathbf{v} = \rho \mathbf{u} + \tilde{\mathbf{J}}^u. \quad (2.48)$$

Remark 2.1. (Mixture momentum) The mixture momentum \mathbf{m} is the sum of the constituent momenta \mathbf{m}_j and thus complies with the first metaphysical principle of mixture theory. Relation (2.48) distinguishes the momentum of the mixture \mathbf{m} from $\rho \mathbf{u}$. We return to this observation later in this section in the context of volume-averaged velocity NSCH models. \triangle

Substituting the key identity (2.48) and the property of the velocity field \mathbf{u} (2.46) into (2.28) provides the alternative form of the mixture mass balance:

$$\overset{\circ}{\rho} + \operatorname{div} \tilde{\mathbf{J}}^u = -\rho \beta \gamma, \quad (2.49)$$

where \circ denotes the material derivative with respect to the volume-averaged velocity \mathbf{u} . Furthermore, substituting the volume-averaged velocity \mathbf{u} and diffusive fluxes (2.47) into (2.35) yields the alternative forms of the phase equations:

$$\overset{\circ}{\phi} + \operatorname{div} \mathbf{h}^u = \frac{\rho}{2\rho_1\rho_2} \gamma, \quad (2.50a)$$

$$\rho \overset{\circ}{c} - c \operatorname{div} \tilde{\mathbf{J}}^u + \operatorname{div} \mathbf{J}^u = \gamma. \quad (2.50b)$$

With the aim of unifying the various formulation of NSCH models in Sec. 3, we relate the various diffusive fluxes in the following lemma.

Lemma 2.1. (Relations diffusive fluxes) *The various diffusive fluxes are related by the identities:*

$$\tilde{\mathbf{J}}^u = \llbracket \rho \rrbracket \mathbf{h}^u, \quad \mathbf{J}^u = \{\rho\} \mathbf{h}^u, \quad \mathbf{h}^v = \frac{\{\rho\}}{\rho} \mathbf{h}^u, \quad \mathbf{J}^v = \frac{\rho_1 \rho_2}{\rho} \mathbf{h}^u. \quad (2.51)$$

Proof. These identities are all consequences of (2.17) and (2.45). For example, to obtain the first identity, partition $\tilde{\mathbf{J}}^u$ as

$$\begin{aligned} \tilde{\mathbf{J}}^u &= \tilde{\rho}_1 \boldsymbol{\omega}_1 + \tilde{\rho}_2 \boldsymbol{\omega}_2 \\ &= \frac{\rho_1}{2} (\phi_1 \boldsymbol{\omega}_1 - \phi_2 \boldsymbol{\omega}_2) - \frac{\rho_2}{2} (\phi_1 \boldsymbol{\omega}_1 - \phi_2 \boldsymbol{\omega}_2) \\ &\quad + \frac{\rho_1}{2} (\phi_1 \boldsymbol{\omega}_1 + \phi_2 \boldsymbol{\omega}_2) + \frac{\rho_2}{2} (\phi_1 \boldsymbol{\omega}_1 + \phi_2 \boldsymbol{\omega}_2). \end{aligned} \quad (2.52)$$

The first line collapses to $\llbracket \rho \rrbracket \mathbf{h}^u$ and the second line vanishes due to (2.45). \square

Analogously to the mass balance and phase equations, we may also formulate the momentum balance in terms of the volume-averaged velocity. To this purpose, we first introduce the following lemma.

Lemma 2.2. (Relation peculiar velocities to stress tensor) *We have the identity:*

$$\sum_j \tilde{\rho}_j \mathbf{w}_j \otimes \mathbf{w}_j = \sum_j \tilde{\rho}_j \boldsymbol{\omega}_j \otimes \boldsymbol{\omega}_j - \frac{1}{\rho} \tilde{\mathbf{J}}^u \otimes \tilde{\mathbf{J}}^u. \quad (2.53)$$

Proof. Identity (2.53) is a direct consequence of the following two identities:

$$\sum_j \tilde{\rho}_j \mathbf{w}_j \otimes \mathbf{w}_j = \sum_j \tilde{\rho}_j \mathbf{v}_j \otimes \mathbf{v}_j - \rho \mathbf{v} \otimes \mathbf{v}, \quad (2.54a)$$

$$\sum_j \tilde{\rho}_j \boldsymbol{\omega}_j \otimes \boldsymbol{\omega}_j = \sum_j \tilde{\rho}_j \mathbf{v}_j \otimes \mathbf{v}_j - \rho \mathbf{v} \otimes \mathbf{v} + \frac{1}{\rho} \tilde{\mathbf{J}}^u \otimes \tilde{\mathbf{J}}^u. \quad (2.54b)$$

To see the first identity, we substitute the definition of the diffuse-velocity \mathbf{w}_j (2.15) into the left-hand side of (2.54a) and expand:

$$\begin{aligned} \sum_j \tilde{\rho}_j \mathbf{w}_j \otimes \mathbf{w}_j &= \sum_j \tilde{\rho}_j (\mathbf{v}_j - \mathbf{v}) \otimes (\mathbf{v}_j - \mathbf{v}) \\ &= \sum_j (\tilde{\rho}_j \mathbf{v}_j \otimes \mathbf{v}_j - \tilde{\rho}_j \mathbf{v}_j \otimes \mathbf{v} - \tilde{\rho}_j \mathbf{v} \otimes \mathbf{v}_j + \tilde{\rho}_j \mathbf{v} \otimes \mathbf{v}) \\ &= \sum_j \tilde{\rho}_j \mathbf{v}_j \otimes \mathbf{v}_j - \rho \mathbf{v} \otimes \mathbf{v} - \rho \mathbf{v} \otimes \mathbf{v} + \rho \mathbf{v} \otimes \mathbf{v} \\ &= \sum_j \tilde{\rho}_j \mathbf{v}_j \otimes \mathbf{v}_j - \rho \mathbf{v} \otimes \mathbf{v}. \end{aligned} \quad (2.55)$$

The second follows in the same fashion via the substitution of the definition of ω_j (2.44) and the key identity (2.48) into the left-hand side of (2.54b):

$$\begin{aligned}
 \sum_j \tilde{\rho}_j \omega_j \otimes \omega_j &= \sum_j \tilde{\rho}_j (\mathbf{v}_j - \mathbf{u}) \otimes (\mathbf{v}_j - \mathbf{u}) \\
 &= \sum_j (\tilde{\rho}_j \mathbf{v}_j \otimes \mathbf{v}_j - \tilde{\rho}_j \mathbf{v}_j \otimes \mathbf{u} - \tilde{\rho}_j \mathbf{u} \otimes \mathbf{v}_j + \tilde{\rho}_j \mathbf{u} \otimes \mathbf{u}) \\
 &= \sum_j \tilde{\rho}_j \mathbf{v}_j \otimes \mathbf{v}_j - \rho \mathbf{v} \otimes \mathbf{u} - \rho \mathbf{u} \otimes \mathbf{v} + \rho \mathbf{u} \otimes \mathbf{u} \\
 &= \sum_j \tilde{\rho}_j \mathbf{v}_j \otimes \mathbf{v}_j - \rho \mathbf{v} \otimes \mathbf{v} + \frac{1}{\rho} \tilde{\mathbf{J}}^u \otimes \tilde{\mathbf{J}}^u. \quad (2.56)
 \end{aligned}$$

By substituting the key identity (2.48) into the mixture linear momentum equation (2.36) and making use of Lemma 2.2, one can show that the mixture linear momentum equation formulated in terms of the volume-averaged velocity takes the following form:

$$\begin{aligned}
 \partial_t(\rho \mathbf{u} + \tilde{\mathbf{J}}^u) + \operatorname{div} \left(\rho \mathbf{u} \otimes \mathbf{u} + \tilde{\mathbf{J}}^u \otimes \mathbf{u} + \mathbf{u} \otimes \tilde{\mathbf{J}}^u + \frac{1}{\rho} \tilde{\mathbf{J}}^u \otimes \tilde{\mathbf{J}}^u \right) \\
 = \operatorname{div} \mathbf{T} + \rho \mathbf{g}. \quad (2.57)
 \end{aligned}$$

Analogously to taking the difference of the constituent mass conservation equations, we may evaluate the difference of the constituent linear momentum equations. This provides a balance law for the diffusive flux. To do so, we first note that by (2.22) and (2.27) we can rewrite the constituent balance law of linear momentum as:

$$(\tilde{\rho}_j \mathbf{v}_j)^\cdot = \operatorname{div} \mathbf{T}_j + \tilde{\rho}_j \mathbf{b}_j + \boldsymbol{\pi}_j + \gamma_j \mathbf{v}_j - \tilde{\rho}_j \mathbf{v}_j \operatorname{div} \mathbf{v}_j. \quad (2.58)$$

Denoting $\mathbf{J}_j^v = \tilde{\rho}_j \mathbf{w}_j$, we have the identity:

$$(\tilde{\rho}_j \mathbf{v}_j)^\cdot = \dot{\mathbf{J}}_j^v + \dot{\tilde{\rho}}_j \mathbf{v} + (\nabla \mathbf{v}) \mathbf{J}_j^v + \tilde{\rho}_j \dot{\mathbf{v}}. \quad (2.59)$$

Substituting this into (2.58) provides:

$$\dot{\mathbf{J}}_j^v + (\nabla \mathbf{v}) \mathbf{J}_j^v = \operatorname{div} \mathbf{T}_j + \tilde{\rho}_j \mathbf{b}_j + \boldsymbol{\pi}_j + \gamma_j \mathbf{v}_j - \dot{\tilde{\rho}}_j \mathbf{v} - \tilde{\rho}_j \dot{\mathbf{v}} - \tilde{\rho}_j \mathbf{v}_j \operatorname{div} \mathbf{v}_j. \quad (2.60)$$

By expanding the peculiar derivative as

$$\begin{aligned}
 \dot{\mathbf{J}}_j^v &= \dot{\mathbf{J}}_j^v + \mathbf{w}_j \cdot \nabla \mathbf{J}_j^v \\
 &= \dot{\mathbf{J}}_j^v + \operatorname{div}(\mathbf{J}_j^v \otimes \mathbf{w}_j) - (\operatorname{div} \mathbf{v}_j) \mathbf{J}_j^v + (\operatorname{div} \mathbf{v}) \mathbf{J}_j^v, \quad (2.61)
 \end{aligned}$$

we arrive at

$$\begin{aligned}
 \dot{\mathbf{J}}_j^v + (\nabla \mathbf{v} + (\operatorname{div} \mathbf{v}) \mathbf{I}) \mathbf{J}_j^v &= \operatorname{div}(\mathbf{T}_j - \mathbf{J}_j^v \otimes \mathbf{w}_j) + \tilde{\rho}_j \mathbf{b}_j \\
 &\quad + \boldsymbol{\pi}_j + \gamma_j \mathbf{v}_j - \gamma_j \mathbf{v} - \tilde{\rho}_j \dot{\mathbf{v}}, \quad (2.62)
 \end{aligned}$$

where we have used the identity:

$$-\dot{\tilde{\rho}}_j \mathbf{v} + (\operatorname{div} \mathbf{v}_j) \mathbf{J}_j^v - \tilde{\rho}_j \mathbf{v}_j \operatorname{div} \mathbf{v}_j = -\gamma_j \mathbf{v}. \quad (2.63)$$

By substituting for the last member in (2.62) the mixture linear momentum evolution equation (2.42) in the form

$$\tilde{\rho}_j \dot{\mathbf{v}} = c_j \operatorname{div} \mathbf{T} + \tilde{\rho}_j \mathbf{b}, \quad (2.64)$$

we obtain

$$\begin{aligned} \dot{\mathbf{J}}_j^v + (\nabla \mathbf{v} + (\operatorname{div} \mathbf{v}) \mathbf{I}) \mathbf{J}_j^v &= \operatorname{div}(\mathbf{T}_j - \mathbf{J}_j^v \otimes \mathbf{w}_j) - c_j \operatorname{div} \mathbf{T} \\ &\quad + \pi_j + \gamma_j \mathbf{v}_j - \gamma_j \mathbf{v} + \tilde{\rho}_j \mathbf{b}_j - \tilde{\rho}_j \mathbf{b}. \end{aligned} \quad (2.65)$$

Note that in case the body force is the same for all components, i.e. $\mathbf{b}_j = \mathbf{b}$, the body force drops out of the evolution equation (2.65). By subtracting (2.65) for constituent 2 from that of constituent 1 we arrive at *the evolution equation of the diffusive flux*:

$$\begin{aligned} \dot{\mathbf{J}}^v + (\nabla \mathbf{v} + (\operatorname{div} \mathbf{v}) \mathbf{I}) \mathbf{J}^v &= \operatorname{div} \left(\mathbf{T}_1 - \mathbf{T}_2 + \frac{\tilde{\rho}_1 - \tilde{\rho}_2}{4\tilde{\rho}_1\tilde{\rho}_2} \mathbf{J}^v \otimes \mathbf{J}^v \right) - c \operatorname{div} \mathbf{T} \\ &\quad + \tilde{\rho}_1(\mathbf{b}_1 - \mathbf{b}) - \tilde{\rho}_2(\mathbf{b}_2 - \mathbf{b}) + \mathbf{p} - \gamma \mathbf{v}, \end{aligned} \quad (2.66)$$

where we have utilized the identity:

$$-\mathbf{J}_1^v \otimes \mathbf{w}_1 + \mathbf{J}_2^v \otimes \mathbf{w}_2 = \frac{\tilde{\rho}_1 - \tilde{\rho}_2}{4\tilde{\rho}_1\tilde{\rho}_2} \mathbf{J}^v \otimes \mathbf{J}^v, \quad \text{for } \phi_1 \phi_2 \neq 0, \quad (2.67)$$

that results from (2.17). This term vanishes for $\phi_1 \phi_2 = 0$.

With the aid of Lemma 2.1, the key identity (2.48) and the mixture mass balance (2.28), one can formulate the evolution equation of the diffusive flux in terms of \mathbf{J}^u . First, the material derivative of the diffusive flux \mathbf{J}^v may be expressed in terms of \mathbf{J}^u via:

$$\dot{\mathbf{J}}^v = \zeta^{-1} \rho^{-1} (\dot{\mathbf{J}}^u - \alpha \mathbf{J}^u \operatorname{div}(\rho^{-1} \mathbf{J}^u) + \beta \gamma \mathbf{J}^u). \quad (2.68)$$

Next, we may express the material derivative $\dot{\mathbf{J}}^u$ as

$$\dot{\mathbf{J}}^u = \overset{\circ}{\mathbf{J}}^u - \alpha \rho^{-1} \mathbf{J}^u \cdot \nabla \mathbf{J}^u, \quad (2.69)$$

and substitution into (2.68) provides:

$$\dot{\mathbf{J}}^v = \zeta^{-1} \rho^{-1} (\overset{\circ}{\mathbf{J}}^u - \alpha \operatorname{div}(\rho^{-1} \mathbf{J}^u \otimes \mathbf{J}^u) + \beta \gamma \mathbf{J}^u). \quad (2.70)$$

Thus the evolution equation of \mathbf{J}^u may be written as

$$\begin{aligned} \overset{\circ}{\mathbf{J}}^u - \alpha \operatorname{div}(\rho^{-1} \mathbf{J}^u \otimes \mathbf{J}^u) + \beta \gamma \mathbf{J}^u &+ (\nabla(\mathbf{u} - \alpha \rho^{-1} \mathbf{J}^u) + \operatorname{div}(-\alpha \rho^{-1} \mathbf{J}^u)) \mathbf{J}^u \\ &= \zeta \rho \left(\operatorname{div} \left(\mathbf{T}_1 - \mathbf{T}_2 + \zeta^{-2} \rho^{-2} \frac{\tilde{\rho}_1 - \tilde{\rho}_2}{4\tilde{\rho}_1\tilde{\rho}_2} \mathbf{J}^u \otimes \mathbf{J}^u \right) - c \operatorname{div} \mathbf{T} \right. \\ &\quad \left. + \tilde{\rho}_1(\mathbf{b}_1 - \mathbf{b}) - \tilde{\rho}_2(\mathbf{b}_2 - \mathbf{b}) + \mathbf{p} - \gamma \mathbf{v} \right). \end{aligned} \quad (2.71)$$

Remark 2.2. (Mixture momentum equation) An equivalent form of (2.57) has appeared in the Ph.D. thesis of Simsek.³⁹ To the best knowledge of the authors, the linear mixture momentum equation of NSCH models formulated in terms of

the volume-averaged velocity appears in two distinct forms. The first is of the form (2.36) with \mathbf{u} instead of the mixture velocity \mathbf{v} and $\rho\mathbf{u}$ instead of \mathbf{m} . The other proposed by Abels *et al.*,³ follows, as observed by Simsek,³⁹ when taking $\mathring{\mathbf{J}}^u = 0$ in (2.57). From the standpoint of mixture theory, the first class relies on the hidden assumption that all the terms containing $\tilde{\mathbf{J}}^u$ vanish, whereas the model of Abels *et al.* contains the hidden assumption $\mathring{\mathbf{J}}^u = 0$. Both assumptions are incompatible with (2.66) and the linear mixture momentum equation in these models does not match with mixture theory. \triangle

Remark 2.3. (Evolution equation diffusive flux) The evolution equation of the diffusive flux seems to be not well known in the phase-field community. We note however that it was first derived in 1975 by Müller²⁹ and revisited in e.g. Refs. 27 and 28. \triangle

To summarize, in agreement with the first unification principle, we have obtained the following *equivalent formulations of the same model*, one formulated in terms of the mixture velocity \mathbf{v} :

$$\left\{ \begin{array}{l} \partial_t(\rho\mathbf{v}) + \operatorname{div}(\rho\mathbf{v} \otimes \mathbf{v}) - \operatorname{div}\mathbf{T} - \rho\mathbf{g} = 0, \end{array} \right. \quad (2.72a)$$

$$\partial_t\rho + \operatorname{div}(\rho\mathbf{v}) = 0, \quad (2.72b)$$

$$\left\{ \begin{array}{l} \mathring{\mathbf{J}}^v + (\nabla\mathbf{v} + (\operatorname{div}\mathbf{v})\mathbf{I})\mathbf{J}^v - \operatorname{div}\left(\mathbf{T}_1 - \mathbf{T}_2 + \frac{\tilde{\rho}_1 - \tilde{\rho}_2}{4\tilde{\rho}_1\tilde{\rho}_2}\mathbf{J}^v \otimes \mathbf{J}^v\right) \\ + c\operatorname{div}\mathbf{T} - \tilde{\rho}_1(\mathbf{b}_1 - \mathbf{b}) + \tilde{\rho}_2(\mathbf{b}_2 - \mathbf{b}) - (\mathbf{p} - \gamma\mathbf{v}) = 0, \end{array} \right. \quad (2.72c)$$

with the evolution equation of an order parameter

$$\text{either: } \dot{\phi} + \phi\operatorname{div}\mathbf{v} + \operatorname{div}\mathbf{h}^v - \zeta\gamma = 0, \quad (2.72d)$$

$$\text{or: } \rho\dot{c} + \operatorname{div}\mathbf{J}^v - \gamma = 0, \quad (2.72e)$$

and one formulated in terms of the volume-averaged velocity \mathbf{u} :

$$\left\{ \begin{array}{l} \partial_t(\rho\mathbf{u} + \tilde{\mathbf{J}}^u) + \operatorname{div}\left(\rho\mathbf{u} \otimes \mathbf{u} + \tilde{\mathbf{J}}^u \otimes \mathbf{u} + \mathbf{u} \otimes \tilde{\mathbf{J}}^u + \frac{1}{\rho}\tilde{\mathbf{J}}^u \otimes \tilde{\mathbf{J}}^u\right) \\ - \operatorname{div}\mathbf{T} - \rho\mathbf{g} = 0, \end{array} \right. \quad (2.73a)$$

$$\operatorname{div}\mathbf{u} - \beta\gamma = 0, \quad (2.73b)$$

$$\left\{ \begin{array}{l} \mathring{\mathbf{J}}^u - \alpha\operatorname{div}(\rho^{-1}\mathbf{J}^u \otimes \mathbf{J}^u) + \beta\gamma\mathbf{J}^u \\ + (\nabla(\mathbf{u} - \alpha\rho^{-1}\mathbf{J}^u) + \operatorname{div}(-\alpha\rho^{-1}\mathbf{J}^u))\mathbf{J}^u \\ - \frac{\beta\rho}{\alpha}\left(\operatorname{div}\left(\mathbf{T}_1 - \mathbf{T}_2 + \alpha^2\beta^{-2}\rho^{-2}\frac{\tilde{\rho}_1 - \tilde{\rho}_2}{4\tilde{\rho}_1\tilde{\rho}_2}\mathbf{J}^u \otimes \mathbf{J}^u\right) - c\operatorname{div}\mathbf{T} \right. \\ \left. + \tilde{\rho}_1(\mathbf{b}_1 - \mathbf{b}) - \tilde{\rho}_2(\mathbf{b}_2 - \mathbf{b}) + \mathbf{p} - \gamma\mathbf{v}\right) = 0, \end{array} \right. \quad (2.73c)$$

with the evolution equation of an order parameter,

$$\text{either: } \mathring{\phi} + \operatorname{div}\mathbf{h}^u - \frac{\rho}{2\rho_1\rho_2}\gamma = 0, \quad (2.73d)$$

$$\text{or: } \rho\mathring{c} - c\operatorname{div}\tilde{\mathbf{J}}^u + \operatorname{div}\mathbf{J}^u - \gamma = 0. \quad (2.73e)$$

3. Constitutive Modeling

In this section, we discuss the constitutive modeling. We will use the Coleman–Noll procedure¹¹ to obtain constitutive equations that guarantee an energy-dissipation equation. Section 3.1 provides the necessary definitions and assumptions. In Sec. 3.2, we establish the constitutive modeling restriction. Then, in Sec. 3.3, we discuss the equivalence of alternative modeling restrictions. The actual selection of the constitutive models appears in Sec. 3.4.

3.1. Definitions, assumptions and modeling choices

We make the following *simplification assumptions* (S):

- the kinetic energies of the constituents are negligible when computed relative to the gross motion of the constituent.
- the body force acting on the constituents is a constant gravitational force, i.e. $\mathbf{b}_j = \mathbf{b} = \mathbf{g} = -g\mathbf{j}$ with g constant and \mathbf{j} the vertical unit vector.

We define the total energy \mathcal{E} associated with the system of balance laws as the sum of the Helmholtz free energy, kinetic energy and gravitational energy:

$$\mathcal{E}(\phi, \mathbf{v}) = \int_{\mathcal{R}(t)} (\Psi + \mathcal{K}(\mathcal{R}(t)) + \mathcal{G}(\mathcal{R}(t))) \, dv. \quad (3.1)$$

Here, $\mathcal{R} = \mathcal{R}(t)$ denotes an arbitrary time-dependent control volume in Ω with volume element dv and unit outward normal $\boldsymbol{\nu}$ that is transported by \mathbf{v} (and thus the normal velocity of $\partial\mathcal{R}(t)$ is $\mathbf{v} \cdot \boldsymbol{\nu}$). The kinetic and gravitational energies are given by

$$\mathcal{K}(\mathcal{R}(t)) = \frac{1}{2} \rho \|\mathbf{v}\|^2, \quad (3.2a)$$

$$\mathcal{G}(\mathcal{R}(t)) = \rho g y. \quad (3.2b)$$

Remark 3.1. (Free energy) The free energy Ψ is defined with respect to the volume element dv . We call this free energy a *volume-measure-based* free energy. The common alternative is to introduce a free energy ψ with respect to the mass element ρdv . This free energy is referred to as a *mass-measure-based* free energy. Throughout this paper, we use Ψ and ψ for volume-measure-based, and mass-measure-based free energies, respectively.

Remark 3.2. (Deviation from mixture theory) At this point, we deviate from mixture theory of rational mechanics laid down by Truesdell.⁴³ Specifically, we diverge from mixture theory in that the first metaphysical principle is violated. Namely, mixture theory dictates that the kinetic energy of the mixture \mathcal{K}_M is the superposition of the individual kinetic energies of the constituents:

$$\mathcal{K}_M = \sum_j \mathcal{K}_j. \quad (3.3)$$

A straightforward substitution of the mixture quantities ρ and $\rho \mathbf{v}$ reveals:

$$\mathcal{K}_M = \mathcal{K} + \sum_j \frac{1}{2} \tilde{\rho}_j \|\mathbf{w}_j\|^2, \quad (3.4)$$

in which the second member represents the kinetic energy of the constituents relative to the gross motion of the constituent. Remark that the second member vanishes in the single constituent and thus the discrepancy appears solely in the interface region. The simplification assumption (S) requires the vanishing of the second member. \triangle

Remark 3.3. (Simplification assumption) The first part of the simplification assumption (S) is closely related to the one adopted by Gurtin²⁰ which states: ‘the momenta and kinetic energies of the constituents are negligible when computed relative to the gross motion of the constituent’. We recall that the first part is not an assumption since the momenta relative to the gross motion of the constituent is absent as stated in (2.16). \triangle

Remark 3.4. (Kinetic energy volume-averaged velocity models) The kinetic energy of diffuse-interface models formulated in terms of the volume-averaged velocity \mathbf{u} is taken as $\rho \|\mathbf{u}\|^2/2$, see e.g. Ref. 3. Just as using \mathcal{K} instead of the kinetic energy of the mixture \mathcal{K}_M , see Remark 3.2, the mismatch exclusively occurs in the interface region. An important difference is that a relation like (3.4), in which the kinetic energy of the mixture is the sum of the approximate kinetic energy and a kinetic energy of with respect to the gross motion of the constituent, does not hold. Instead we have the relation:

$$\mathcal{K}_M = \frac{1}{2} \rho \|\mathbf{u}\|^2 + \sum_j \frac{1}{2} \tilde{\rho}_j \|\mathbf{w}_j\|^2 + \frac{1}{2\rho} \tilde{\mathbf{J}}^u \cdot \tilde{\mathbf{J}}^u + \tilde{\mathbf{J}}^u \cdot \mathbf{u}, \quad (3.5)$$

which follows from the identity relation (2.48) and reveals a non-obvious approximation of the kinetic energy of the mixture by $\rho \|\mathbf{u}\|^2/2$. The last term in (3.5) is not guaranteed positive and thus does not represent a kinetic energy relative to some different velocity. Relation (2.48) reveals that the kinetic energy \mathcal{K} formulated in terms of the volume-averaged velocity reads:

$$\mathcal{K}(\mathcal{R}(t)) = \frac{(\rho \mathbf{u} + \tilde{\mathbf{J}}^u) \cdot (\rho \mathbf{u} + \tilde{\mathbf{J}}^u)}{2\rho}. \quad (3.6)$$

\triangle

We proceed with selecting the volume fraction difference ϕ as order parameter and postulating the free energy to pertain to the constitutive class:

$$\Psi = \hat{\Psi}(\phi, \nabla \phi, \mathbf{D}), \quad (3.7)$$

where we recall that \mathbf{D} is the symmetric gradient of the mixture velocity:

$$\mathbf{D} = \frac{1}{2}(\nabla \mathbf{v} + (\nabla \mathbf{v})^T). \quad (3.8)$$

Next, we define a chemical potential-like quantity in the usual way as

$$\hat{\mu} := \frac{\partial \hat{\Psi}}{\partial \phi} - \operatorname{div} \frac{\partial \hat{\Psi}}{\partial \nabla \phi}. \quad (3.9)$$

In the phase-field community this is sometimes referred to as the Fréchet (or variational) derivative of the total Helmholtz free energy. We postpone the selection of the constitutive classes of the stress tensor, diffusive flux and the mass flux, denoted $\hat{\mathbf{T}}$, $\hat{\mathbf{h}}^v$ and $\hat{\gamma}$, to the end of Sec. 3.2. With the aim of deriving NSCH models, we postulate the energy-dissipation law:

$$\frac{d}{dt} \mathcal{E}(\phi, \mathbf{v}) = \mathcal{W}(\mathcal{R}(t)) - \mathcal{D}(\mathcal{R}(t)), \quad (3.10)$$

where $\mathcal{W}(\mathcal{R}(t))$ is the rate of work performed by macro- and micro stresses coming through the boundary $\partial \mathcal{R}(t)$ and $\mathcal{D}(\mathcal{R}(t))$ is the dissipation for which we demand $\mathcal{D}(\mathcal{R}(t)) \geq 0$.

Remark 3.5. (Arbitrariness of modeling choices) The choice of working with the volume fraction difference ϕ as the order parameter seems arbitrary. One could instead work with the concentration difference c . Additionally, one may work with a mass-measure-based free energy instead of a volume-measure-based free energy. In Sec. 3.3, we discuss the relation of the modeling choices in detail. \triangle

Remark 3.6. (Constitutive class diffusive flux) From the viewpoint of mixture theory, the usage of a constitutive class for the diffusive flux should come as a surprise. Namely, in Sec. 2, we have established an evolution equation for the mass diffusion \mathbf{J}^v , i.e. (2.66). Here, we discard the PDE (2.66). This is an approximation of the mixture theory framework, or at least, it is not obvious how it fits in the framework. This approximation however is essential in order to work with a reduced model of NSCH type. In other words, without the approximation one does not retrieve an NSCH-type model. \triangle

Remark 3.7. (Energy-dissipation law) The energy-dissipation statement is a necessary and core element in the derivation of NSCH models. Equation (3.10) is closely linked to the first law of thermodynamics whereas the requirement $\mathcal{D}(\mathcal{R}(t)) \geq 0$ is associated with the second law of thermodynamics. In the literature on NSCH models the energy-dissipation law is often called the second law of thermodynamics and models that satisfy it are referred to as thermodynamically consistent. However, we note that the energy-dissipation statement is not obviously compatible with the second law of thermodynamics presented in mixture theory, see e.g. Ref. 43. As such, referring to NSCH models as thermodynamically consistent is not justified. We refer to Refs. 4 and 13 for some remarks on approximations in the energy-dissipation statement. \triangle

Remark 3.8. (Rate of work term) By using concepts like the microforce balance, the rate of work term may be already defined at this point, see Ref. 20. \triangle

Remark 3.9. (Alternative modeling approach) Instead of working with constitutive classes and the Coleman–Noll procedure, an alternative is to postulate the dissipation, see Ref. 26. \triangle

3.2. Derivation of the constitutive modeling restriction

We proceed with the evaluation of the evolution of the energy (3.1). By applying Reynolds transport theorem to the free energy $\hat{\Psi}$, we have

$$\frac{d}{dt} \int_{\mathcal{R}(t)} \hat{\Psi} \, dv = \int_{\mathcal{R}(t)} \partial_t \hat{\Psi} \, dv + \int_{\partial \mathcal{R}(t)} \hat{\Psi} \mathbf{v} \cdot \boldsymbol{\nu} \, da. \quad (3.11)$$

In the next step, we apply the divergence theorem and expand the derivatives:

$$\frac{d}{dt} \int_{\mathcal{R}(t)} \hat{\Psi} \, dv = \int_{\mathcal{R}(t)} \frac{\partial \hat{\Psi}}{\partial \phi} \dot{\phi} + \frac{\partial \hat{\Psi}}{\partial \nabla \phi} \cdot (\nabla \phi)^\cdot + \frac{\partial \hat{\Psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} + \hat{\Psi} \operatorname{div} \mathbf{v} \, dv, \quad (3.12)$$

where we recall that the dot $\dot{}$ is the material derivative with respect to the mixture velocity \mathbf{v} . By substituting the identity

$$(\nabla \phi)^\cdot = \nabla(\dot{\phi}) - (\nabla \phi)^T \nabla \mathbf{v}, \quad (3.13)$$

into (3.12) and subsequently integrating by parts, we arrive at

$$\begin{aligned} \frac{d}{dt} \int_{\mathcal{R}(t)} \hat{\Psi} \, dv &= \int_{\mathcal{R}(t)} \hat{\mu} \dot{\phi} - \left(\nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} \right) : \nabla \mathbf{v} + \frac{\partial \hat{\Psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} + \hat{\Psi} \operatorname{div} \mathbf{v} \, dv \\ &\quad + \int_{\partial \mathcal{R}(t)} \dot{\phi} \frac{\partial \hat{\Psi}}{\partial \nabla \phi} \cdot \boldsymbol{\nu} \, da. \end{aligned} \quad (3.14)$$

We substitute the phase evolution equation (2.35a) for the material derivative $\dot{\phi}$ and apply integration by parts to obtain

$$\begin{aligned} \frac{d}{dt} \int_{\mathcal{R}(t)} \hat{\Psi} \, dv &= \int_{\mathcal{R}(t)} \nabla \hat{\mu} \cdot \hat{\mathbf{h}}^v - \left(\nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} \right) : \nabla \mathbf{v} + \frac{\partial \hat{\Psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} \\ &\quad + (\hat{\Psi} - \hat{\mu} \phi) \operatorname{div} \mathbf{v} + \hat{\mu} \zeta \dot{\gamma} \, dv \\ &\quad + \int_{\partial \mathcal{R}(t)} \left(-\hat{\mu} \hat{\mathbf{h}}^v + \dot{\phi} \frac{\partial \hat{\Psi}}{\partial \nabla \phi} \right) \cdot \boldsymbol{\nu} \, da. \end{aligned} \quad (3.15)$$

Next, we turn our focus on the kinetic energy. In a similar fashion we apply Reynolds transport theorem and find:

$$\begin{aligned} \frac{d}{dt} \int_{\mathcal{R}(t)} \mathcal{K}(\mathcal{R}(t)) \, dv &= \int_{\mathcal{R}(t)} \mathbf{v} \cdot \partial_t(\rho \mathbf{v}) - \frac{1}{2} \|\mathbf{v}\|^2 \partial_t \rho \, dv \\ &\quad + \int_{\partial \mathcal{R}(t)} \mathcal{K}(\mathcal{R}(t)) \mathbf{v} \cdot \boldsymbol{\nu} \, da. \end{aligned} \quad (3.16)$$

Substitution of the mass and momentum equations provides:

$$\begin{aligned} \frac{d}{dt} \int_{\mathcal{R}(t)} \mathcal{K}(\mathcal{R}(t)) \, dv &= \int_{\mathcal{R}(t)} -\mathbf{v} \cdot \operatorname{div}(\rho \mathbf{v} \otimes \mathbf{v}) + \frac{1}{2} \|\mathbf{v}\|^2 \operatorname{div}(\rho \mathbf{v}) + \mathbf{v} \cdot \operatorname{div} \hat{\mathbf{T}} \\ &\quad + \rho \mathbf{v} \cdot \mathbf{g} \, dv + \int_{\partial \mathcal{R}(t)} \mathcal{K}(\mathcal{R}(t)) \mathbf{v} \cdot \boldsymbol{\nu} \, da. \end{aligned} \quad (3.17)$$

With the aid of the identity

$$-\mathbf{v} \cdot \operatorname{div}(\rho \mathbf{v} \otimes \mathbf{v}) + \frac{1}{2} \|\mathbf{v}\|_2^2 \operatorname{div}(\rho \mathbf{v}) = -\operatorname{div} \left(\frac{1}{2} \rho \|\mathbf{v}\|^2 \mathbf{u} \right), \quad (3.18)$$

and using integration by parts on the stress tensor, the evolution of the kinetic energy simplifies to:

$$\frac{d}{dt} \int_{\mathcal{R}(t)} \mathcal{K}(\mathcal{R}(t)) \, dv = \int_{\mathcal{R}(t)} -\nabla \mathbf{v} : \hat{\mathbf{T}} + \rho \mathbf{v} \cdot \mathbf{g} \, dv + \int_{\partial \mathcal{R}(t)} \mathbf{v} \cdot \hat{\mathbf{T}} \boldsymbol{\nu} \, da. \quad (3.19)$$

Finally, for the gravitational energy evolution we also apply Reynolds transport theorem to find:

$$\frac{d}{dt} \int_{\mathcal{R}(t)} \mathcal{G}(\mathcal{R}(t)) \, dv = \int_{\mathcal{R}(t)} g y \partial_t \rho \, dv + \int_{\partial \mathcal{R}(t)} \mathcal{G}(\mathcal{R}(t)) \mathbf{v} \cdot \boldsymbol{\nu} \, da. \quad (3.20)$$

Substituting the mass evolution equation (2.28) and subsequently providing integration by part leads to

$$\frac{d}{dt} \int_{\mathcal{R}(t)} \mathcal{G}(\mathcal{R}(t)) \, dv = - \int_{\mathcal{R}(t)} \rho \mathbf{v} \cdot \mathbf{g} \, dv. \quad (3.21)$$

Taking the sum of (3.15), (3.19) and (3.21) we arrive at

$$\begin{aligned} \frac{d}{dt} \mathcal{E}(\phi, \mathbf{v}) &= \int_{\partial \mathcal{R}(t)} \left(\mathbf{v}^T \hat{\mathbf{T}} - \hat{\mu} \hat{\mathbf{h}}^v + \dot{\phi} \frac{\partial \hat{\Psi}}{\partial \nabla \phi} \right) \cdot \boldsymbol{\nu} \, da \\ &\quad - \int_{\mathcal{R}(t)} \left(\hat{\mathbf{T}} + \nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} + (\hat{\mu} \phi - \hat{\Psi}) \mathbf{I} \right) : \nabla \mathbf{v} - \nabla \hat{\mu} \cdot \hat{\mathbf{h}}^v \\ &\quad - \frac{\partial \hat{\Psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} - \hat{\mu} \zeta \hat{\gamma} \, dv. \end{aligned} \quad (3.22)$$

Next, we observe that the components $\nabla \mathbf{v}$, $\hat{\mathbf{h}}^v$ and $\hat{\gamma}$ are not independent. Namely, we have the sequence of identities:

$$-\operatorname{div} \mathbf{v} = -\alpha(\dot{\phi} + \phi \operatorname{div} \mathbf{v}) = \alpha \operatorname{div} \hat{\mathbf{h}}^v - \beta \hat{\gamma}. \quad (3.23)$$

The first identity is a direct consequence of the mass balance of the mixture (2.28) and the second follows from the phase equation (2.35a). We now introduce the following partition:

$$\hat{\mathbf{T}} = \hat{\mathbf{T}}_0 - \hat{p} \mathbf{I} \quad \text{with} \quad \hat{\mathbf{T}}_0 := \hat{\mathbf{T}} + \hat{p} \mathbf{I}, \quad (3.24)$$

in which \hat{p} is a scalar field that represents the mechanical pressure and $-\hat{p}\mathbf{I}$ corresponds to the hydro-static part of the stress tensor $\hat{\mathbf{T}}$. To exploit the degeneracy, we let the scalar field \hat{p} act as a Lagrange multiplier on (3.23) by multiplying this equation by \hat{p} :

$$\begin{aligned} -\hat{p} \operatorname{div} \mathbf{v} &= \hat{p} \alpha \operatorname{div} \hat{\mathbf{h}}^v - \hat{p} \beta \hat{\gamma} \\ &= -\nabla(\hat{p} \alpha) \cdot \hat{\mathbf{h}}^v + \operatorname{div}(\hat{p} \alpha \hat{\mathbf{h}}^v) - \hat{p} \beta \hat{\gamma}. \end{aligned} \quad (3.25)$$

If we substitute the partition (3.24) and the identity (3.25) into (3.22) we can identify the rate of work performed by macro- and micro stresses and the dissipation as

$$\mathcal{W}(\mathcal{R}(t)) = \int_{\partial \mathcal{R}(t)} \left(\mathbf{v}^T (\hat{\mathbf{T}}_0 - \hat{p} \mathbf{I}) - (\hat{\mu} + \alpha \hat{p}) \hat{\mathbf{h}}^v + \dot{\phi} \frac{\partial \hat{\Psi}}{\partial \nabla \phi} \right) \cdot \boldsymbol{\nu} \, da, \quad (3.26a)$$

$$\begin{aligned} \mathcal{D}(\mathcal{R}(t)) &= \int_{\mathcal{R}(t)} \left(\hat{\mathbf{T}}_0 + \nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} + (\hat{\mu} \phi - \hat{\Psi}) \mathbf{I} \right) : \nabla \mathbf{v} \\ &\quad - \nabla(\hat{\mu} + \hat{p} \alpha) \cdot \hat{\mathbf{h}}^v - \frac{\partial \hat{\Psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} - (\hat{\mu} + \alpha \hat{p}) \zeta \hat{\gamma} \, dv. \end{aligned} \quad (3.26b)$$

Positivity (\geq) of the diffusion $\mathcal{D}(\mathcal{R}(t))$ implies, since the control volume $\mathcal{R}(t)$ is arbitrary, positivity of the integrand. To ensure this, we impose positivity of the members in the integrand independently. The constitutive class of the free energy Ψ , Eq. (3.7), restricts dependence of Ψ on \mathbf{D} . For a more elaborate discussion on the possible dependence of Ψ on \mathbf{D} we refer to Ref. 20. This implies that positivity of the last member of the integrand of $\mathcal{D}(\mathcal{R}(t))$ can only be achieved by requiring it to vanish via $\partial \Psi / \partial \mathbf{D} = 0$. This means the constitutive class of Ψ reduces to

$$\Psi = \hat{\Psi}(\phi, \nabla \phi), \quad (3.27)$$

and as a consequence the second to last member in the integrand in (3.26b) vanishes. The energy-dissipation law is thus satisfied when the local inequality is fulfilled:

$$\begin{aligned} &\left(\hat{\mathbf{T}}_0 + \nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} + (\hat{\mu} \phi - \hat{\Psi}) \mathbf{I} \right) : \nabla \mathbf{v} \\ &\quad - \nabla(\hat{\mu} + \alpha \hat{p}) \cdot \hat{\mathbf{h}}^v - (\hat{\mu} + \alpha \hat{p}) \zeta \hat{\gamma} \geq 0. \end{aligned} \quad (3.28)$$

Based on the inequality (3.28), we now restrict ourselves to stress tensors \mathbf{T} , diffusive fluxes \mathbf{h}^v and mass fluxes γ belonging to the constitutive classes:

$$\mathbf{T} = \hat{\mathbf{T}}(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \mathbf{D}, \hat{p}), \quad (3.29a)$$

$$\mathbf{h}^v = \hat{\mathbf{h}}^v(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \nabla \hat{p}), \quad (3.29b)$$

$$\gamma = \hat{\gamma}(\phi, \hat{\mu}, \hat{p}). \quad (3.29c)$$

3.3. Equivalence of alternative modeling restrictions

As mentioned in Remark 3.5, some of the modeling choices in Sec. 3.1 seem arbitrary. As a consequence, the modeling restriction obtained in Sec. 3.2 may seem discretionary. In Appendix A, we provide the derivations of the most obvious alternative modeling restrictions. An overview of the variations is as follows:

- (1) Sections 3.1 and 3.2: a volume-measure-based free energy Ψ and the difference of volume fractions ϕ as order parameter,
- (2) A.1: a mass-measure-based free energy ψ and the difference of volume fractions ϕ as order parameter,
- (3) A.2: a volume-measure-based free energy Ψ and the difference of concentrations c as order parameter,
- (4) A.3: a mass-measure-based free energy ψ and the difference of concentrations c as order parameter.

The associated constitutive classes of these modeling choices are

$$\begin{aligned} (1) : \quad \Psi &= \hat{\Psi}(\phi, \nabla \phi), & \mathbf{T} &= \hat{\mathbf{T}}(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \mathbf{D}, \hat{p}), \\ \mathbf{h}^v &= \hat{\mathbf{h}}^v(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \nabla \hat{p}), & \gamma &= \hat{\gamma}(\phi, \hat{\mu}, \hat{p}), \end{aligned} \quad (3.30a)$$

$$\begin{aligned} (2) : \quad \psi &= \hat{\psi}(\phi, \nabla \phi), & \mathbf{T} &= \hat{\mathbf{T}}(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \mathbf{D}, \hat{p}), \\ \mathbf{h}^v &= \hat{\mathbf{h}}^v(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \nabla \hat{p}), & \gamma &= \hat{\gamma}(\phi, \hat{\mu}, \hat{p}), \end{aligned} \quad (3.30b)$$

$$\begin{aligned} (3) : \quad \Psi &= \check{\Psi}(c, \nabla c), & \mathbf{T} &= \check{\mathbf{T}}(c, \nabla c, \check{\mu}, \nabla \check{\mu}, \mathbf{D}, \check{p}), \\ \mathbf{J}^v &= \check{\mathbf{J}}^v(c, \nabla c, \check{\mu}, \nabla \check{\mu}, \nabla \check{p}), & \gamma &= \check{\gamma}(c, \check{\mu}, \check{p}), \end{aligned} \quad (3.30c)$$

$$\begin{aligned} (4) : \quad \psi &= \check{\psi}(c, \nabla c), & \mathbf{T} &= \check{\mathbf{T}}(c, \nabla c, \check{\mu}, \nabla \check{\mu}, \mathbf{D}, \check{p}), \\ \mathbf{J}^v &= \check{\mathbf{J}}^v(c, \nabla c, \check{\mu}, \nabla \check{\mu}, \nabla \check{p}), & \gamma &= \check{\gamma}(c, \check{\mu}, \check{p}), \end{aligned} \quad (3.30d)$$

where \hat{p} , $\hat{\hat{p}}$, \check{p} and $\check{\check{p}}$ are the pressures associated with the modeling choices and the chemical potential-like variables are, respectively, given by

$$(1) : \quad \hat{\mu} = \frac{\partial \hat{\Psi}}{\partial \phi} - \operatorname{div} \frac{\partial \hat{\Psi}}{\partial \nabla \phi}, \quad (3.31a)$$

$$(2) : \quad \hat{\hat{\mu}} = \frac{\partial \hat{\psi}}{\partial \phi} - \frac{1}{\rho} \operatorname{div} \left(\rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} \right), \quad (3.31b)$$

$$(3) : \quad \check{\mu} = \frac{\partial \check{\Psi}}{\partial c} - \operatorname{div} \frac{\partial \check{\Psi}}{\partial \nabla c}, \quad (3.31c)$$

$$(4) : \quad \check{\check{\mu}} = \frac{\partial \check{\psi}}{\partial c} - \frac{1}{\rho} \operatorname{div} \left(\rho \frac{\partial \check{\psi}}{\partial \nabla c} \right). \quad (3.31d)$$

Collecting the final restrictions provides:

$$(1) : \left(\hat{\mathbf{T}}_0 + \nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} + (\hat{\mu} \phi - \hat{\Psi}) \mathbf{I} \right) : \nabla \mathbf{v} \\ - \nabla(\hat{\mu} + \alpha \hat{p}) \cdot \hat{\mathbf{h}}^v - \zeta(\hat{\mu} + \alpha \hat{p}) \hat{\gamma} \geq 0, \quad (3.32a)$$

$$(2) : \left(\hat{\hat{\mathbf{T}}}_0 + \nabla \phi \otimes \rho \frac{\partial \hat{\hat{\Psi}}}{\partial \nabla \phi} + \rho \hat{\hat{\mu}} \phi \mathbf{I} \right) : \nabla \mathbf{v} \\ - \nabla(\rho \hat{\hat{\mu}} + \alpha \hat{\hat{p}}) \cdot \hat{\hat{\mathbf{h}}}^v - \zeta(\rho \hat{\hat{\mu}} + \alpha \hat{\hat{p}}) \hat{\hat{\gamma}} \geq 0, \quad (3.32b)$$

$$(3) : \left(\check{\mathbf{T}}_0 + \nabla c \otimes \frac{\partial \check{\Psi}}{\partial \nabla c} - \check{\Psi} \mathbf{I} \right) : \nabla \mathbf{v} \\ - \nabla \left(\frac{\check{\mu}}{\rho} + \beta \check{p} \right) \cdot \check{\mathbf{J}}^v - \left(\frac{\check{\mu}}{\rho} + \beta \check{p} \right) \check{\gamma} \geq 0, \quad (3.32c)$$

$$(4) : \left(\check{\check{\mathbf{T}}}_0 + \nabla c \otimes \rho \frac{\partial \check{\check{\Psi}}}{\partial \nabla c} \right) : \nabla \mathbf{v} \\ - \nabla(\check{\check{\mu}} + \check{\beta} \check{\check{p}}) \cdot \check{\check{\mathbf{J}}}^v - (\check{\check{\mu}} + \check{\beta} \check{\check{p}}) \check{\check{\gamma}} \geq 0. \quad (3.32d)$$

In order to allow comparison of these modeling restrictions, the relation between these modeling classes needs to be specified. We select the following relations:

free energy classes:

$$\hat{\Psi}(\phi, \nabla \phi) \equiv \hat{\rho}(\phi) \hat{\psi}(\phi, \nabla \phi) \equiv \check{\rho}(c) \check{\psi}(c, \nabla c) \equiv \check{\Psi}(c, \nabla c), \quad (3.33)$$

stress tensor classes:

$$\hat{\mathbf{T}}(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \hat{p}) \equiv \hat{\hat{\mathbf{T}}}(\phi, \nabla \phi, \hat{\hat{\mu}}, \nabla \hat{\hat{\mu}}, \hat{\hat{p}}) \\ \equiv \check{\check{\mathbf{T}}}(c, \nabla c, \check{\check{\mu}}, \nabla \check{\check{\mu}}, \check{\check{p}}) \\ \equiv \check{\mathbf{T}}(c, \nabla c, \check{\mu}, \nabla \check{\mu}, \check{p}), \quad (3.34)$$

diffusive flux classes:

$$\hat{\mathbf{h}}^v(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \nabla \hat{p}) \equiv \hat{\hat{\mathbf{h}}}^v(\phi, \nabla \phi, \hat{\hat{\mu}}, \nabla \hat{\hat{\mu}}, \nabla \hat{\hat{p}}) \\ \equiv \zeta \check{\check{\mathbf{J}}}^v(c, \nabla c, \check{\check{\mu}}, \nabla \check{\check{\mu}}, \nabla \check{\check{p}}) \\ \equiv \zeta \check{\mathbf{J}}^v(c, \nabla c, \check{\mu}, \nabla \check{\mu}, \nabla \check{p}), \quad (3.35)$$

mass flux classes:

$$\hat{\gamma}(\phi, \hat{\mu}, \hat{p}) \equiv \hat{\hat{\gamma}}(\phi, \hat{\hat{\mu}}, \hat{\hat{p}}) \equiv \check{\gamma}(c, \check{\mu}, \check{p}) \equiv \check{\check{\gamma}}(c, \check{\check{\mu}}, \check{\check{p}}). \quad (3.36)$$

The particular identification choice of the diffusive fluxes originates from the relation between the diffusive fluxes before constitutive modeling (Lemma 2.1).

Note that a direct consequence of the free energy classes (3.33) is the equivalence of the associated Korteweg tensors:

$$\nabla\phi \otimes \frac{\partial\hat{\Psi}}{\partial\nabla\phi} = \nabla\phi \otimes \rho \frac{\partial\hat{\psi}}{\partial\nabla\phi} = \nabla c \otimes \rho \frac{\partial\check{\psi}}{\partial\nabla c} = \nabla c \otimes \frac{\partial\check{\Psi}}{\partial\nabla c}, \quad (3.37)$$

where we recall the chain rule and the observations $\phi = \phi(c)$ and $c = c(\phi)$.

Lemma 3.1. (Relations between chemical potential-like quantities) *The chemical potential-like quantities are related via the following identities:*

$$\hat{\mu} = \rho\hat{\mu} + \hat{\psi}[\rho], \quad (3.38a)$$

$$\check{\mu} = \rho\check{\mu} - \beta\rho^2\check{\psi}, \quad (3.38b)$$

$$\check{\mu} = \frac{\rho^2}{\rho_1\rho_2}\hat{\mu}. \quad (3.38c)$$

Proof. See Appendix B. □

We are now ready to state one of the main results of this paper: *the choice of the order parameter and the type of the free energy do not influence the modeling restrictions*. The selection of a particular order parameter should not be regarded as a modeling step but simply as part of the variable selection as one can easily alter via a variable transformation.

Theorem 3.1. (Equivalence modeling restrictions) *Selecting the following relations between the pressures of the various modeling choices:*

$$\hat{p} = \hat{p} + \hat{\psi}\{\rho\}, \quad (3.39a)$$

$$\check{p} = \check{p} + \rho\check{\psi}, \quad (3.39b)$$

$$\check{p} = \hat{p} + \hat{\mu}\phi, \quad (3.39c)$$

yields equivalence of the restrictions in (3.32).

Proof. The proof relies on the identifications (3.33)–(3.36), the relation (3.37) and Lemma 3.1. Details are provided in Appendix B. □

3.4. Selection of constitutive models

Now that we have established equivalence of the modeling restrictions, we proceed with the one derived in this section:

$$\begin{aligned} & \left(\hat{\mathbf{T}}_0 + \nabla\phi \otimes \frac{\partial\hat{\Psi}}{\partial\nabla\phi} + (\hat{\mu}\phi - \hat{\Psi})\mathbf{I} \right) : \nabla\mathbf{v} \\ & - \nabla(\hat{\mu} + \alpha\hat{p}) \cdot \hat{\mathbf{h}}^v - \zeta(\hat{\mu} + \alpha\hat{p})\hat{\gamma} \geq 0. \end{aligned} \quad (3.40)$$

To avoid that variations of $\nabla \mathbf{v}$ lead to a violation of the modeling restriction, we make the constitutive choice:

$$\hat{\mathbf{T}}_0 = -\nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} - (\hat{\mu} \phi - \hat{\Psi}) \mathbf{I} + \nu(\phi)(2\mathbf{D} + \lambda(\operatorname{div} \mathbf{v}) \mathbf{I}), \quad (3.41)$$

which is in agreement with (2.40) and precludes violation of the energy-dissipation law by the stress contribution. Here, $\nu(\phi) \geq 0$ is the dynamic phase-dependent viscosity and $\lambda \geq -2/d$ is a scalar, where we recall that d is the number of dimensions. This form of the viscous stress tensor assumes an isotropic Newtonian mixture. A direct consequence is the expression of the stress tensor \mathbf{T} :

$$\hat{\mathbf{T}} = -\nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} - (\mu \phi - \hat{\Psi}) \mathbf{I} + \nu(\phi)(2\mathbf{D} + \lambda(\operatorname{div} \mathbf{v}) \mathbf{I}) - \hat{p} \mathbf{I}. \quad (3.42)$$

Next, we focus on the diffusive flux $\hat{\mathbf{h}}^v$. Insisting positivity of the second member in (3.40) implies the following form:

$$\hat{\mathbf{h}}^v = -\hat{\mathbf{M}}^v \nabla (\hat{\mu} + \alpha \hat{p}), \quad (3.43)$$

for some constitutive quantity $\hat{\mathbf{M}}^v = \hat{\mathbf{M}}^v(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \nabla \hat{p})$, referred to as *mobility tensor*, that is consistent with the inequality

$$-\nabla (\hat{\mu} + \alpha \hat{p}) \cdot (\hat{\mathbf{M}}^v \nabla (\hat{\mu} + \alpha \hat{p})) \leq 0 \quad (3.44)$$

for all $(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \nabla \hat{p})$. This may be understood as the *generalized Fick laws of diffusion*.

Lastly, we insist positivity of the third term in (3.40). Noting that $\zeta > 0$, this implies the following form:

$$\hat{\gamma} = -\hat{m}(\phi, \hat{\mu}, \hat{p})(\hat{\mu} + \alpha \hat{p}), \quad (3.45)$$

where the mobility $m = \hat{m}(\phi, \hat{\mu}, \hat{p})$ only attains positive values ($m \geq 0$). Ensuring vanishing fluxes in the single fluid region requires $\hat{m}(\phi = \pm 1, \hat{\mu}, \hat{p}) = 0$.

Remark 3.10. (Constitutive models) The particular constitutive models of the diffusive flux (3.43) and mass flux (3.45) are the only constitutive models in the specified classes (3.29b) and (3.29c), that guarantee positivity of the second and third term in (3.40), respectively. This claim results from a theorem on the solution of thermodynamical inequalities proved by Gurtin.¹⁹ One could also employ this theorem to construct the most general form of the stress tensor. In this paper, we only present a simple constitutive choice of the stress tensor in agreement with unification principle three. \triangle

Remark 3.11. (Mobility tensor) The first appearance of the mobility tensor is in Ref. 19. In phase-field literature it is common to work with an isotropic mobility tensor. \triangle

Remark 3.12. (Incompatible mobility in single fluid region) We show that a degenerate mobility is required in general. Consider in the general non-matching

density case ($\alpha \neq 0$) a region $\mathcal{D}_a \in \Omega$ in which $\phi \equiv a$ with $-1 \leq a \leq 1$ constant. Note that in \mathcal{D}_a we have $\hat{\Psi} = \hat{\Psi}(\phi, \nabla\phi) = \hat{\Psi}(\phi) = \hat{\Psi}(a) = \text{constant}$, and thus $\hat{\mu} = 0$ in \mathcal{D}_a . As a consequence we deduce $\hat{\mathbf{M}}^v = \hat{\mathbf{M}}^v(a, \mathbf{0}, 0, \mathbf{0}, \nabla\hat{p})$ in \mathcal{D}_a . The mass conservation equation (2.28) takes the following form:

$$\operatorname{div} \mathbf{v} = 0, \quad \text{in } \mathcal{D}_a, \quad (3.46)$$

whereas the phase equation (2.35a) reduces to

$$a \operatorname{div} \mathbf{v} - \alpha \operatorname{div}(\hat{\mathbf{M}}^v \nabla \hat{p}) = -\beta \hat{m} \hat{p}, \quad \text{in } \mathcal{D}_a. \quad (3.47)$$

Therefore

$$\operatorname{div}(\hat{\mathbf{M}}^v \nabla \hat{p}) = \zeta \hat{m} \hat{p}, \quad \text{in } \mathcal{D}_a, \quad (3.48)$$

representing a balance equation for the pressure. Next, consider the cases $a = \pm 1$. In this situation the mass flux contribution vanishes: $m = 0$ and thus (3.48) reduces to

$$\operatorname{div}(\hat{\mathbf{M}}^v \nabla \hat{p}) = 0, \quad \text{in } \mathcal{D}_{\pm 1}. \quad (3.49)$$

In general (3.49) represents a balance equation, purely formulated in terms of the pressure, that holds in the single fluid. There exists no such *nontrivial equation* that matches with the standard incompressible Navier–Stokes equations in the single fluid regime. In particular, when $\hat{\mathbf{M}}^v$ is a non-zero constant tensor it follows that

$$\Delta \hat{p} = 0, \quad \text{in } \mathcal{D}_{\pm 1}, \quad (3.50)$$

which in general *does not hold* in the single fluid regime (remark that (3.50) holds for Stokes flow with a divergence free-force). Thus in general we are left with the trivial instance of (3.49):

$$\hat{\mathbf{M}}^v = 0, \quad \text{in } \mathcal{D}_{\pm 1}, \quad (3.51)$$

i.e. the mobility tensor is of degenerate type. Note that for many models proposed in literature a non-degenerate mobility was chosen, see Table 2. \triangle

Remark 3.13. (Degenerate mobility) The relevance of a degenerate mobility tensor is not new and is in fact well known in community, see e.g. Refs. 1, 2 and 8. We note that the model presented in Ref. 3 has been studied for a variety of mobility choices. Its sharp interface limit has been rigorously shown to exist for both degenerate and non-degenerate mobilities, and the associated sharp interface free boundary problem depends the choice of the mobility.

Remark 3.14. (Alternative reasoning degenerate mobility) Instead of following the above arguments leading to a degenerate mobility, one could reason as follows. Note that a substitution of the volume fractions (2.31) into the volume-averaged velocity (2.43) and subsequently into the equation for diffusive flux (2.66) provides the alternative form of the diffusive flux \mathbf{h}^u :

$$\mathbf{h}^u = -\xi(\phi) \llbracket \mathbf{v} \rrbracket, \quad (3.52)$$

with

$$\xi(\phi) := \phi^2 - 1 \leq 0. \quad (3.53)$$

Obviously \mathbf{h}^u vanishes whenever $\phi = \pm 1$. Recalling Lemma 2.1, we have

$$\mathbf{h}^v = \frac{\{\rho\}}{\rho} \mathbf{h}^u = -\frac{\{\rho\}}{\rho} \xi(\phi) \llbracket \mathbf{v} \rrbracket, \quad (3.54)$$

and thus the diffusive flux \mathbf{h}^v also vanishes in the single-fluid case $\phi = \pm 1$. Demanding the constitutive model of the diffusive flux $\mathbf{h}^v = \hat{\mathbf{h}}^v$ to be compatible with this restriction we get:

$$\hat{\mathbf{M}}^v \nabla \hat{p} = 0, \quad \text{for } \phi = \pm 1, \quad (3.55)$$

where $\hat{\mathbf{M}}^v$ only depends on $\nabla \hat{p}$. As before, the only solution that matches with the standard incompressible Navier–Stokes equations in the single fluid regime is the trivial one:

$$\hat{\mathbf{M}}^v = 0, \quad \text{for } \phi = \pm 1. \quad (3.56)$$

Note that (3.54) suggests the specific form $\hat{\mathbf{M}}^v = -\xi(\phi) \bar{\mathbf{M}}$ for some $\bar{\mathbf{M}}$.

Of course, this argument can also be made when working with the concentration c as order parameter. On the account of Lemma 2.1 and the identity

$$\xi(\phi) = \frac{\rho^2}{\rho_1 \rho_2} \xi(c), \quad (3.57)$$

we have

$$\mathbf{J}^u = -\frac{\rho^2 \{\rho\}}{\rho_1 \rho_2} \xi(c) \llbracket \mathbf{v} \rrbracket. \quad (3.58)$$

Again employing Lemma 2.1 reveals that the diffusive flux \mathbf{J}^v takes the following form:

$$\mathbf{J}^v = \frac{\rho_1 \rho_2}{\{\rho\} \rho} \mathbf{J}^u = -\rho \xi(c) \llbracket \mathbf{v} \rrbracket. \quad (3.59)$$

This quantity vanishes in the single-fluid regime $c = \pm 1$, and using the constitutive class $\mathbf{J}^v = \check{\mathbf{J}}^v$ we can deduce that the associated mobility tensor vanishes when $c = \pm 1$. \triangle

Remark 3.15. (Alternative diffusive fluxes) The equivalence of the diffusive fluxes, (3.35), provides the form of the diffusive flux $\check{\mathbf{J}}^v$:

$$\check{\mathbf{J}}^v = -\check{\mathbf{M}}^v \nabla (\hat{\mu} + \alpha \hat{p}), \quad (3.60)$$

with $\check{\mathbf{M}}^v = \zeta^{-1} \hat{\mathbf{M}}^v$. Additionally, by identifying the diffusive flux $\hat{\mathbf{h}}^u = (\rho / \{\rho\}) \hat{\mathbf{h}}^v$ we find:

$$\hat{\mathbf{h}}^u = -\hat{\mathbf{M}}^u \nabla (\hat{\mu} + \alpha \hat{p}), \quad (3.61)$$

with $\hat{\mathbf{M}}^u = (\rho / \{\rho\}) \hat{\mathbf{M}}^v$. \triangle

This completes the constitutive modeling. Via substituting (3.42), (3.43) and (3.45) into (2.72) we have obtained *one model* that can be formulated in various

ways using:

- the mixture velocity \mathbf{v} or the volume-averaged velocity \mathbf{u} ,
- the volume fraction difference ϕ or the concentration difference c ,
- the mass-measure-based free energy $\hat{\psi}$ or the volume-measure-based free energy $\hat{\Psi}$.

We withhold from presenting many formulations and only make variations in the mean velocity to obtain one formulation in terms of the mixture velocity:

$$\left\{ \begin{array}{l} \partial_t(\rho\mathbf{v}) + \operatorname{div}(\rho\mathbf{v} \otimes \mathbf{v}) + \nabla\hat{p} + \operatorname{div}\left(\nabla\phi \otimes \frac{\partial\hat{\Psi}}{\partial\nabla\phi} + (\mu\phi - \hat{\Psi})\mathbf{I}\right) \\ \quad - \operatorname{div}(\nu(2\mathbf{D} + \lambda(\operatorname{div}\mathbf{v})\mathbf{I})) - \rho\mathbf{g} = 0, \\ \partial_t\rho + \operatorname{div}(\rho\mathbf{v}) = 0, \\ \partial_t\phi + \operatorname{div}(\phi\mathbf{v}) - \operatorname{div}(\hat{\mathbf{M}}^v\nabla(\hat{\mu} + \alpha\hat{p})) + \zeta\hat{m}(\hat{\mu} + \alpha\hat{p}) = 0, \\ \hat{\mu} - \frac{\partial\hat{\Psi}}{\partial\phi} + \operatorname{div}\left(\frac{\partial\hat{\Psi}}{\partial\nabla\phi}\right) = 0, \end{array} \right. \quad \begin{array}{l} (3.62a) \\ (3.62b) \\ (3.62c) \\ (3.62d) \end{array}$$

and one formulation in terms of the volume-averaged velocity:

$$\left\{ \begin{array}{l} \partial_t(\rho\mathbf{u} + \tilde{\mathbf{J}}^u) + \operatorname{div}\left(\rho\mathbf{u} \otimes \mathbf{u} + \tilde{\mathbf{J}}^u \otimes \mathbf{u} + \mathbf{u} \otimes \tilde{\mathbf{J}}^u + \frac{1}{\rho}\tilde{\mathbf{J}}^u \otimes \tilde{\mathbf{J}}^u\right) \\ \quad + \nabla\hat{p} + \operatorname{div}\left(\nabla\phi \otimes \frac{\partial\hat{\Psi}}{\partial\nabla\phi} + (\hat{\mu}\phi - \hat{\Psi})\mathbf{I}\right) \\ \quad - \operatorname{div}(\nu(2\nabla^s(\mathbf{u} + \rho^{-1}\tilde{\mathbf{J}}^u) + \lambda(\operatorname{div}(\mathbf{u} + \rho^{-1}\tilde{\mathbf{J}}^u))\mathbf{I})) - \rho\mathbf{g} = 0, \\ \operatorname{div}\mathbf{u} + \beta\hat{m}(\hat{\mu} + \alpha\hat{p}) = 0, \\ \partial_t\phi + \mathbf{u} \cdot \nabla\phi - \operatorname{div}(\hat{\mathbf{M}}^u\nabla(\hat{\mu} + \alpha\hat{p})) + \frac{\rho}{2\rho_1\rho_2}\hat{m}(\hat{\mu} + \alpha\hat{p}) = 0, \\ \hat{\mu} - \frac{\partial\hat{\Psi}}{\partial\phi} + \operatorname{div}\left(\frac{\partial\hat{\Psi}}{\partial\nabla\phi}\right) = 0, \end{array} \right. \quad \begin{array}{l} (3.63a) \\ (3.63b) \\ (3.63c) \\ (3.63d) \end{array}$$

with $\tilde{\mathbf{J}}^u = -[\rho]\hat{\mathbf{M}}^u\nabla(\hat{\mu} + \alpha\hat{p})$. Both models obviously satisfy the exact same form of the energy-dissipation statement (3.10), which complies with the second unification principle. We remark that the variable transformation that links the models (3.62) and (3.63) via (2.48) now involves a constitutive model for the diffusive flux $\tilde{\mathbf{J}}^u$.

4. Unification of Existing Navier–Stokes Cahn–Hilliard Models

As discussed in the introductory section, there exists a wide spectrum with many flavors of NSCH models. It is the purpose of this section to unify them.

Lowengrub and Truskinovsky²⁵

To indicate the relation between our model and that of Ref. 25 we set $\gamma = 0$ and formulate our model in terms of $(\mathbf{v}, c, \check{\psi})$:

$$\left\{ \begin{array}{l} \partial_t(\rho\mathbf{v}) + \operatorname{div}(\rho\mathbf{v} \otimes \mathbf{v}) + \nabla \check{p} + \operatorname{div} \left(\nabla c \otimes \rho \frac{\partial \check{\psi}}{\partial \nabla c} \right) \\ \quad - \operatorname{div}(\nu(c)(2\mathbf{D} + \lambda(\operatorname{div}\mathbf{v})\mathbf{I})) - \rho\mathbf{g} = 0, \\ \partial_t \rho + \operatorname{div}(\rho\mathbf{v}) = 0, \\ \rho \dot{c} - \operatorname{div}(\hat{\mathbf{M}}^v(\phi(c))\nabla(\check{\mu} + \beta\check{p})) = 0, \\ \check{\mu} - \frac{\partial \check{\psi}}{\partial c} + \frac{1}{\rho} \operatorname{div} \left(\rho \frac{\partial \check{\psi}}{\partial \nabla c} \right) = 0. \end{array} \right. \quad \begin{array}{l} (4.1a) \\ (4.1b) \\ (4.1c) \\ (4.1d) \end{array}$$

To establish (4.1c) we have employed the relation between the diffusive classes (3.35):

$$\hat{\mathbf{h}}^v(\phi, \nabla\phi, \hat{\mu}, \nabla\hat{\mu}, \nabla\hat{p}) = \zeta \check{\mathbf{J}}^v(c, \nabla c, \check{\mu}, \nabla\check{\mu}, \nabla\check{p}) \quad (4.2)$$

and the identity

$$\check{\mu} + \beta\check{p} = \zeta(\hat{\mu} + \alpha\hat{p}). \quad (4.3)$$

First, we note that the mass balance of the mixture may be written as

$$\operatorname{div}\mathbf{v} + \frac{1}{\rho}\check{\rho}'(c)\dot{c} = 0. \quad (4.4)$$

Next, one recognizes the derivative $\partial\rho/\partial c$ in the evolution of the order parameter (4.1c) via:

$$\rho \dot{c} - \operatorname{div} \left(\hat{\mathbf{M}}^v(\phi(c))\nabla \left(\check{\mu} - \frac{\check{p}}{\rho^2}\check{\rho}'(c) \right) \right) = 0. \quad (4.5)$$

Substitution of (4.1d) into (4.5) provides the alternative form:

$$\rho \dot{c} - \operatorname{div} \left(\hat{\mathbf{M}}^v(\phi(c))\nabla \left(\frac{\partial \check{\psi}}{\partial c} - \frac{1}{\rho} \operatorname{div} \left(\rho \frac{\partial \check{\psi}}{\partial \nabla c} \right) - \frac{\check{p}}{\rho^2}\check{\rho}'(c) \right) \right) = 0. \quad (4.6)$$

To obtain the model of Lowengrub and Truskinovsky²⁵ one has to set $\check{\psi} = \sigma\varphi(c)/\epsilon + \sigma\epsilon|\nabla c|^2/2$ for some $\varphi = \varphi(c)$, and replace the quantity $\hat{\mathbf{M}}^v(\phi(c))$ in (4.6) by a non-zero constant isotropic mobility $\tilde{m}\mathbf{I}$:

$$\left\{ \begin{array}{l} \partial_t(\rho\mathbf{v}) + \operatorname{div}(\rho\mathbf{v} \otimes \mathbf{v}) + \nabla p + \sigma\epsilon \operatorname{div}(\nabla c \otimes \rho \nabla c) \\ \quad - \operatorname{div}(\nu(c)(2\mathbf{D} + \lambda(\operatorname{div}\mathbf{v})\mathbf{I})) = 0, \\ \operatorname{div}\mathbf{v} + \frac{1}{\rho}\check{\rho}'(c)\dot{c} = 0, \\ \rho \dot{c} - \operatorname{div} \left(\tilde{m}\nabla \left(\frac{\sigma}{\epsilon} \frac{\partial \varphi}{\partial c} - \frac{\sigma\epsilon}{\rho} \operatorname{div}(\rho \nabla c) - \frac{p}{\rho^2}\check{\rho}'(c) \right) \right) = 0. \end{array} \right. \quad \begin{array}{l} (4.7a) \\ (4.7b) \\ (4.7c) \end{array}$$

In other words, this comparison reveals that the quasi-incompressible model of Ref. 25 is identical to our model (up to the mobility type, see Remark 3.12).

Shokrpour Roudbari *et al.*,³⁸ Aki *et al.*⁴ and Shen *et al.*³⁷

We now explore the relation of our model with the models Refs. 4, 37 and 38. For the sake of clarity we restrict to $\gamma = 0$ and note that only model Ref. 4 contains mass fluxes. An alternative formulation of model (3.62) follows when instead of identity (3.25) one employs

$$\begin{aligned} -p^* \operatorname{div} \mathbf{v} &= \frac{p^* \llbracket \rho \rrbracket}{\rho} \dot{\phi} \\ &= \frac{p^* \llbracket \rho \rrbracket}{\rho} (-\operatorname{div} \hat{\mathbf{h}}^v - \phi \operatorname{div} \mathbf{v}) \\ &= \nabla \left(\frac{p^* \llbracket \rho \rrbracket}{\rho} \right) \cdot \hat{\mathbf{h}}^v - \frac{p^* \llbracket \rho \rrbracket}{\rho} \phi \operatorname{div} \mathbf{v} - \operatorname{div} \left(\frac{p^* \llbracket \rho \rrbracket}{\rho} \hat{\mathbf{h}}^v \right), \end{aligned} \quad (4.8)$$

in which p^* is the pressure quantity. The associated stress tensor and diffusive flux take the form

$$\begin{cases} \hat{\mathbf{T}} = -\nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} - \left(\hat{\mu} \phi - \hat{\Psi} - \frac{p^* \llbracket \rho \rrbracket}{\rho} \phi \right) \mathbf{I} \\ \quad + \nu(\phi)(2\mathbf{D} + \lambda(\operatorname{div} \mathbf{v})\mathbf{I}) - p\mathbf{I}, \\ \hat{\mathbf{h}}^v = -\hat{\mathbf{M}}^v(\phi) \nabla \left(\hat{\mu} - \frac{p^* \llbracket \rho \rrbracket}{\rho} \phi \right). \end{cases} \quad (4.9a)$$

$$(4.9b)$$

Noting that the mass equation may be written as

$$\operatorname{div} \mathbf{v} - \alpha \operatorname{div} \left(\hat{\mathbf{M}}^v(\phi) \nabla \left(\hat{\mu} - \frac{p^* \llbracket \rho \rrbracket}{\rho} \phi \right) \right) = 0, \quad (4.10)$$

we obtain the equivalent model:

$$\begin{cases} \partial_t(\rho \mathbf{v}) + \operatorname{div}(\rho \mathbf{v} \otimes \mathbf{v}) + \nabla p^* \\ \quad + \operatorname{div} \left(\nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} + \left(\hat{\mu} \phi - \hat{\Psi} - \frac{p^* \llbracket \rho \rrbracket}{\rho} \phi \right) \mathbf{I} \right) \\ \quad - \operatorname{div}(\nu(\phi)(2\mathbf{D} + \lambda(\operatorname{div} \mathbf{v})\mathbf{I})) - \rho \mathbf{g} = 0, \\ \operatorname{div} \mathbf{v} - \alpha \operatorname{div} \left(\hat{\mathbf{M}}^v(\phi) \nabla \left(\hat{\mu} - \frac{p^* \llbracket \rho \rrbracket}{\rho} \phi \right) \right) = 0, \\ \dot{\phi} + \phi \operatorname{div} \mathbf{v} - \operatorname{div} \left(\hat{\mathbf{M}}^v(\phi) \nabla \left(\hat{\mu} - \frac{p^* \llbracket \rho \rrbracket}{\rho} \phi \right) \right) = 0, \\ \hat{\mu} - \frac{\partial \hat{\Psi}}{\partial \phi} + \operatorname{div} \left(\frac{\partial \hat{\Psi}}{\partial \nabla \phi} \right) = 0. \end{cases} \quad (4.11a)$$

$$(4.11b)$$

$$(4.11c)$$

$$(4.11d)$$

The third and fourth terms in the momentum equation (4.11a) may be written as

$$\begin{aligned} \nabla p^* + \operatorname{div} \left(\nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} + \left(\hat{\mu} \phi - \hat{\Psi} - \frac{p^* \llbracket \rho \rrbracket}{\rho} \phi \right) \mathbf{I} \right) \\ = \nabla \left(p^* \frac{\{\rho\}}{\rho} \right) + \operatorname{div} \left(\nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} + \left(\hat{\mu} \phi - \hat{\Psi} \right) \mathbf{I} \right). \end{aligned} \quad (4.12)$$

The last member of the phase equation (4.11c) may be written as

$$-\operatorname{div} \left(\hat{\mathbf{M}}^v(\phi) \nabla \left(\hat{\mu} - \frac{p^* \llbracket \rho \rrbracket}{\rho} \right) \right) = -\operatorname{div} \left(\hat{\mathbf{M}}^v(\phi) \nabla \left(\hat{\mu} + \alpha p^* \frac{\{\rho\}}{\rho} \right) \right). \quad (4.13)$$

Equivalence of model (4.11) with model (3.62) follows from (4.12) and (4.13) via the variable transformation:

$$\hat{p} = p^* \frac{\{\rho\}}{\rho} = \frac{p^*}{1 - \alpha \phi}. \quad (4.14)$$

Up to the definition of the mobility, this model (and thus also model (3.62)) is equivalent to the model proposed by Ref. 38. A variable transformation presented in Ref. 38 reveals that the models Refs. 4 and 37 are equivalent, up to the definition of the mobility, to the model Ref. 38. This reveals that all these models are equivalent to our model (up to the mobility type, see Remark 3.12).

Table 2. Comparison of the various NSCH models. The column ‘MT-consistent balance laws’ indicates whether the balance laws of the model are compatible with mixture theory. In the third column ‘Compatible in single fluid’ we state whether the model has a degenerate or non-degenerate mobility, see Remark 3.12, and in the last column whether the model is energy dissipative. The symbol ★ indicates that there is an energy-dissipation law but that the associated kinetic energy is not an obvious approximation of the kinetic energy of the mixture, see also Remark 3.4.

Model	MT-consistent balance laws	Compatible in single fluid	Energy-dissipation law
Abels <i>et al.</i> ³	✗	✓	★
Aki <i>et al.</i> ⁴	✓	✗	✓
Boyer ⁹	✗	✓	✗
Ding <i>et al.</i> ¹²	✗	✓	✗
Lowengrub and Truskinovsky ²⁵	✓	✗	✓
Shen <i>et al.</i> ³⁷	✓	✗	✓
Shokrpour Roudbari <i>et al.</i> ³⁸	✓	✗	✓
Current	✓	✓	✓

Boyer,⁹ Ding *et al.*¹² and Abels *et al.*³

The models Refs. 9, 12 and 3 are formulated in terms of the volume-averaged mean velocity. The linear momentum equation deviates from that in our model (3.63), and as such these models are incompatible with our model. In contrast to the models Refs. 9 and 12, model Ref. 3 is presented with an energy-dissipation law. The kinetic energy in this law is however not an obvious approximation of the kinetic energy of the mixture, see Remark 3.4. Finally, we note that these models are consistent with the incompressible Navier–Stokes equations in the single-fluid regime.

We close this section with an overview of the various models presented in Table 2.

5. Summary and Outlook

In this paper, we established a unified framework of all existing NSCH models. To this purpose, we used the general continuum mixture theory and laid down three unifying principles:

- (1) there is only one system of continuum mechanics balance laws that describes the physical model,
- (2) there is only one natural energy-dissipation law that leads to quasi-incompressible NSCH models,
- (3) variations between the models can only appear in the constitutive choices.

In Sec. 2, we provided a precise statement of the principles of mixture theory and their consequences. Furthermore, we showed that the mixture framework leads to one system of balance laws that can be formulated using different variable sets, e.g. in terms of a mass-averaged or volume-averaged velocity. Formulating the balance laws using the volume-averaged velocity, we found a system distinct from existing volume-averaged velocity based models. We illustrated the incompatibility with mixture theory of volume-averaged velocity based models that appear in the literature. This can however easily be repaired. Next, in Sec. 3, we demonstrated how an energy-dissipation law naturally leads to quasi-incompressible NSCH models. We proved that the energy-dissipative modeling restriction for the constitutive classes is independent of the variable set. Additionally, we showed that in our framework the mobility tensor of the diffuse flux is of degenerate type. In Sec. 4, we demonstrated that, using the appropriate degenerate mobility, existing NSCH models are often equivalent reformulations of the same physical model. Finally, we presented an overview of the various NSCH models in Table 2.

While we think that the presented framework and the associated analysis are useful to gain insight into NSCH models, we do not claim that they are sufficient in this aspect. We outline some of our thoughts on potential further directions for future research. First, it is essential to establish the sharp interface asymptotics and associated free energy inequalities. We conjecture that the approaches and techniques presented in Refs. 3 and 4 can directly be applied to the model proposed herein as well. A second important direction for future research lies in

the rigorous mathematical analysis of the proposed models. Furthermore, we note that, even though the continuous formulations are equivalent, associated discretization methods are, at least *ab initio*, not identical. As such, it would be valuable to design numerical schemes that inherit useful properties of the model via convenient formulations. Lastly, it would be worthwhile to explore alternative two-phase flow models that utilize the evolution equation of the diffusive flux.

Appendix A. Alternative Constitutive Modeling

In this appendix, we provide the derivation of the most common alternative constitutive modeling approaches:

- A.1: volume fraction ϕ and mass-measure-based free energy ψ ,
- A.2: concentration c and volume-measure-based free energy Ψ ,
- A.3: concentration c and mass-measure-based free energy ψ .

A.1. Constitutive modeling: Volume fraction and mass-measure-based free energy

Using the same simplification assumption (S), the total energy \mathcal{E} is the superposition of the Helmholtz free energy, kinetic energy and gravitational energy:

$$\mathcal{E}(\phi, \mathbf{v}) = \int_{\mathcal{R}(t)} (\rho\psi + \mathcal{K}(\mathcal{R}(t)) + \mathcal{G}(\mathcal{R}(t))) \, dv, \quad (\text{A.1})$$

where $\mathcal{R} = \mathcal{R}(t)$ denotes an arbitrary time-dependent control volume in Ω that is transported by the mixture velocity \mathbf{v} . The mass-measure-based free energy is postulated to pertain to the constitutive class:

$$\psi = \hat{\psi}(\phi, \nabla\phi, \mathbf{D}), \quad (\text{A.2})$$

and we define the chemical potential-like quantity as

$$\hat{\mu} = \frac{\partial \hat{\psi}}{\partial \phi} - \frac{1}{\rho} \operatorname{div} \left(\rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} \right). \quad (\text{A.3})$$

Analogously to Sec. 3.2, we work with constitutive classes for stress tensor $\mathbf{T} = \hat{\mathbf{T}}$, diffusive flux $\mathbf{h}^v = \hat{\mathbf{h}}^v$ and mass flux $\gamma = \hat{\gamma}$ and postpone their specification. Next, we postulate the energy-dissipation law:

$$\frac{d}{dt} \mathcal{E}(\phi, \mathbf{v}) = \mathcal{W}(\mathcal{R}(t)) - \mathcal{D}(\mathcal{R}(t)), \quad (\text{A.4})$$

in which $\mathcal{W}(\mathcal{R}(t))$ and $\mathcal{D}(\mathcal{R}(t)) \geq 0$ have the same interpretation as in Sec. 3.2. With the aid of Reynolds transport theorem and the mixture mass balance law

(2.28), the evolution of the free energy in (A.1) takes the form:

$$\begin{aligned} \frac{d}{dt} \int_{\mathcal{R}(t)} \rho \hat{\psi} \, dv &= \int_{\mathcal{R}(t)} \rho \dot{\hat{\psi}} \, dv \\ &= \int_{\mathcal{R}(t)} \rho \left(\frac{\partial \hat{\psi}}{\partial \phi} \dot{\phi} + \frac{\partial \hat{\psi}}{\partial \nabla \phi} \cdot (\nabla \phi) \cdot + \frac{\partial \hat{\psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} \right) \, dv. \end{aligned} \quad (\text{A.5})$$

Substitution of (3.13) into (A.5) and subsequently the integration by parts provides:

$$\begin{aligned} \frac{d}{dt} \int_{\mathcal{R}(t)} \rho \hat{\psi} \, dv &= \int_{\mathcal{R}(t)} \rho \hat{\mu} \dot{\phi} - \left(\nabla \phi \otimes \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} \right) : \nabla \mathbf{v} + \rho \frac{\partial \hat{\psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} \, dv \\ &\quad + \int_{\partial \mathcal{R}(t)} \dot{\phi} \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} \cdot \boldsymbol{\nu} \, da. \end{aligned} \quad (\text{A.6})$$

We now eliminate the material derivative $\dot{\phi}$ via the substitution of the phase evolution equation (2.35a):

$$\begin{aligned} \frac{d}{dt} \int_{\mathcal{R}(t)} \rho \hat{\psi} \, dv &= \int_{\mathcal{R}(t)} \nabla(\rho \hat{\mu}) \cdot \hat{\mathbf{h}}^v - \left(\nabla \phi \otimes \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} \right) : \nabla \mathbf{v} + \rho \frac{\partial \hat{\psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} \\ &\quad - \rho \hat{\mu} \phi \operatorname{div} \mathbf{v} + \hat{\gamma} \zeta \rho \hat{\mu} \, dv \\ &\quad + \int_{\partial \mathcal{R}(t)} \left(-\rho \hat{\mu} \hat{\mathbf{h}}^v + \dot{\phi} \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} \right) \cdot \boldsymbol{\nu} \, da. \end{aligned} \quad (\text{A.7})$$

Taking now the sum of (A.7), kinetic energy evolution (3.19) and gravitational energy evolution (3.21) we arrive at

$$\begin{aligned} \frac{d}{dt} \mathcal{E}(\phi, \mathbf{v}) &= \int_{\partial \mathcal{R}(t)} \left(\mathbf{v}^T \hat{\mathbf{T}} - \rho \hat{\mu} \hat{\mathbf{h}}^v + \dot{\phi} \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} \right) \cdot \boldsymbol{\nu} \, da \\ &\quad - \int_{\mathcal{R}(t)} \left(\hat{\mathbf{T}} + \nabla \phi \otimes \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} + \mu \phi \mathbf{I} \right) : \nabla \mathbf{v} \\ &\quad - \nabla(\rho \hat{\mu}) \cdot \hat{\mathbf{h}}^v - \frac{\partial \hat{\psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} - \zeta(\rho \hat{\mu} + \alpha \hat{p}) \hat{\gamma} \, dv. \end{aligned} \quad (\text{A.8})$$

Employing the partition:

$$\hat{\mathbf{T}} = \hat{\mathbf{T}}_0 - \hat{p} \mathbf{I} \quad \text{with} \quad \hat{\mathbf{T}}_0 := \hat{\mathbf{T}} + \hat{p} \mathbf{I}, \quad (\text{A.9})$$

in which \hat{p} is a scalar pressure field and the identity (3.25) with pressure \hat{p} in (3.22) allows to identify the rate of work and dissipation as

$$\mathcal{W}(\mathcal{R}(t)) = \int_{\partial \mathcal{R}(t)} \left(\mathbf{v}^T (\hat{\mathbf{T}}_0 - \hat{p} \mathbf{I}) - (\rho \hat{\mu} + \alpha \hat{p}) \hat{\mathbf{h}}^v + \dot{\phi} \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} \right) \cdot \boldsymbol{\nu} \, da, \quad (\text{A.10a})$$

$$\begin{aligned} \mathcal{D}(\mathcal{R}(t)) = \int_{\mathcal{R}(t)} & \left(\hat{\mathbf{T}}_0 + \nabla \phi \otimes \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} + \rho \hat{\mu} \phi \mathbf{I} \right) : \nabla \mathbf{v} \\ & - \nabla(\rho \hat{\mu} + \hat{p} \alpha) \cdot \hat{\mathbf{h}}^v - \frac{\partial \hat{\psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} - \zeta(\rho \hat{\mu} + \alpha \hat{p}) \hat{\gamma} \, dv. \end{aligned} \quad (\text{A.10b})$$

By following the same argument as in Sec. 3.2 the positivity of the diffusion leads to the reduced class:

$$\psi = \hat{\psi}(\phi, \nabla \phi), \quad (\text{A.11})$$

and the modeling restriction readily follows:

$$\begin{aligned} & \left(\hat{\mathbf{T}}_0 + \nabla \phi \otimes \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} + \rho \hat{\mu} \phi \mathbf{I} \right) : \nabla \mathbf{v} \\ & - \nabla(\rho \hat{\mu} + \alpha \hat{p}) \cdot \hat{\mathbf{h}}^v - \zeta(\rho \hat{\mu} + \alpha \hat{p}) \hat{\gamma} \geq 0. \end{aligned} \quad (\text{A.12})$$

Based on the form of the modeling restriction (A.12) we select the following constitutive classes:

$$\mathbf{T} = \hat{\mathbf{T}}(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \mathbf{D}, \hat{p}), \quad (\text{A.13a})$$

$$\mathbf{h}^v = \hat{\mathbf{h}}^v(\phi, \nabla \phi, \hat{\mu}, \nabla \hat{\mu}, \nabla \hat{p}), \quad (\text{A.13b})$$

$$\gamma = \hat{\gamma}(\phi, \hat{\mu}, \hat{p}). \quad (\text{A.13c})$$

A.2. Constitutive modeling: Concentration and volume-measure-based free energy

With the aid of the simplification assumption (S), the total energy \mathcal{E} reads in terms of the concentration c :

$$\mathcal{E}(c, \mathbf{v}) = \int_{\mathcal{R}(t)} (\Psi + \mathcal{K}(\mathcal{R}(t)) + \mathcal{G}(\mathcal{R}(t))) \, dv, \quad (\text{A.14})$$

where again $\mathcal{R} = \mathcal{R}(t)$ denotes an arbitrary time-dependent control volume in Ω that is transported by the mixture velocity \mathbf{v} . The volume-measure-based free energy is postulated to belong to the class:

$$\Psi = \check{\Psi}(c, \nabla c, \mathbf{D}), \quad (\text{A.15})$$

and the concentration-based chemical potential-like quantity is the Fréchet derivative of the Helmholtz free energy Ψ with respect to c :

$$\check{\mu} = \frac{\partial \check{\Psi}}{\partial c} - \operatorname{div} \frac{\partial \check{\Psi}}{\partial \nabla c}. \quad (\text{A.16})$$

We work with constitutive classes for stress tensor $\mathbf{T} = \check{\mathbf{T}}$, diffusive flux $\mathbf{h}^v = \check{\mathbf{h}}^v$ and mass flux $\gamma = \check{\gamma}$ and postpone their specification. We postulate the energy-dissipation law:

$$\frac{d}{dt} \mathcal{E}(c, \mathbf{v}) = \mathcal{W}(\mathcal{R}(t)) - \mathcal{D}(\mathcal{R}(t)), \quad (\text{A.17})$$

in which we use the same interpretation of $\mathscr{W}(\mathcal{R}(t))$ and $\mathscr{D}(\mathcal{R}(t)) \geq 0$ as in Sec. 3. To proceed we follow the same procedure as in Sec. 3 with the concentration evolution equation (2.35b) and find

$$\begin{aligned} \frac{d}{dt} \mathcal{E}(c, \mathbf{v}) &= \int_{\partial \mathcal{R}(t)} \left(\mathbf{v}^T \check{\mathbf{T}} - \check{\mu} \frac{\check{\mathbf{J}}^v}{\rho} + \dot{c} \frac{\partial \check{\Psi}}{\partial \nabla c} \right) \cdot \boldsymbol{\nu} \, da \\ &\quad - \int_{\mathcal{R}(t)} \left(\check{\mathbf{T}} + \nabla c \otimes \frac{\partial \check{\Psi}}{\partial \nabla c} - \check{\Psi} \mathbf{I} \right) : \nabla \mathbf{v} - \nabla \left(\frac{\check{\mu}}{\rho} \right) \cdot \check{\mathbf{J}}^v \\ &\quad - \frac{\partial \check{\Psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} - \frac{\check{\mu}}{\rho} \check{\gamma} \, dv. \end{aligned} \quad (\text{A.18})$$

Next, we introduce the partition:

$$\check{\mathbf{T}} = \check{\mathbf{T}}_0 - \check{p} \mathbf{I} \quad \text{with} \quad \check{\mathbf{T}}_0 := \check{\mathbf{T}} + \check{p} \mathbf{I}, \quad (\text{A.19})$$

in which \check{p} is a scalar pressure field. Via the concentration equation (2.35b) we deduce the identity

$$\begin{aligned} -\check{p} \operatorname{div} \mathbf{v} &= \check{p} \left(\frac{1}{\rho^2} \frac{\partial \rho}{\partial c} \rho \dot{c} \right) \\ &= -\check{p} \frac{1}{\rho^2} \frac{\partial \rho}{\partial c} (\operatorname{div} \check{\mathbf{J}}^v - \check{\gamma}) \\ &= \frac{\check{p}}{2} \left(\frac{1}{\rho_1} - \frac{1}{\rho_2} \right) (\operatorname{div} \check{\mathbf{J}}^v - \check{\gamma}) \\ &= \beta \check{p} \operatorname{div} \check{\mathbf{J}}^v - \beta \check{p} \check{\gamma}, \end{aligned} \quad (\text{A.20})$$

where we recall the definition:

$$\beta = \frac{\rho_2 - \rho_1}{2\rho_1\rho_2} = -\frac{\llbracket \rho \rrbracket}{\rho_1\rho_2}. \quad (\text{A.21})$$

Via substitution of the identity (A.20) we can identify the rate of work and dissipation term:

$$\mathscr{W}(\mathcal{R}(t)) = \int_{\partial \mathcal{R}(t)} \left(\mathbf{v}^T (\check{\mathbf{T}}_0 - \check{p} \mathbf{I}) - \left(\frac{\check{\mu}}{\rho} + \beta \check{p} \right) \check{\mathbf{J}}^v + \dot{c} \frac{\partial \check{\Psi}}{\partial \nabla c} \right) \cdot \boldsymbol{\nu} \, da, \quad (\text{A.22a})$$

$$\begin{aligned} \mathscr{D}(\mathcal{R}(t)) &= \int_{\mathcal{R}(t)} \left(\check{\mathbf{T}}_0 + \nabla c \otimes \frac{\partial \check{\Psi}}{\partial \nabla c} - \check{\Psi} \mathbf{I} \right) : \nabla \mathbf{v} \\ &\quad - \nabla \left(\frac{\check{\mu}}{\rho} + \beta \check{p} \right) \cdot \check{\mathbf{J}}^v - \frac{\partial \check{\Psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} - \left(\frac{\check{\mu}}{\rho} + \beta \check{p} \right) \check{\gamma} \, dv. \end{aligned} \quad (\text{A.22b})$$

Using the same argument as before we work with the reduced class of the free energy:

$$\Psi = \check{\Psi}(c, \nabla c), \quad (\text{A.23})$$

and obtain the modeling restriction:

$$\left(\check{\mathbf{T}}_0 + \nabla c \otimes \frac{\partial \check{\Psi}}{\partial \nabla c} - \check{\Psi} \mathbf{I} \right) : \nabla \mathbf{v} - \nabla \left(\frac{\check{\mu}}{\rho} + \beta \check{p} \right) \cdot \check{\mathbf{J}}^v - \left(\frac{\check{\mu}}{\rho} + \beta \check{p} \right) \check{\gamma} \geq 0. \quad (\text{A.24})$$

Based on (A.24) we specify the classes for stress tensor, diffusive flux, and mass flux:

$$\mathbf{T} = \check{\mathbf{T}}(c, \nabla c, \mu, \nabla \mu, \mathbf{D}, \check{p}), \quad (\text{A.25a})$$

$$\mathbf{J}^v = \check{\mathbf{J}}^v(c, \nabla c, \mu, \nabla \mu, \nabla \check{p}), \quad (\text{A.25b})$$

$$\gamma = \check{\gamma}(c, \check{\mu}, \check{p}). \quad (\text{A.25c})$$

A.3. Constitutive modeling: Concentration and mass-measure-based free energy

The simplification assumption (S) provides the total energy \mathcal{E} in terms of the concentration c :

$$\mathcal{E}(c, \mathbf{v}) = \int_{\mathcal{R}(t)} (\rho \psi + \mathcal{K}(\mathcal{R}(t)) + \mathcal{G}(\mathcal{R}(t))) \, dv, \quad (\text{A.26})$$

where again $\mathcal{R} = \mathcal{R}(t)$ denotes an arbitrary time-dependent control volume in Ω that is transported by the mixture velocity \mathbf{v} . The mass-measure-based free energy is postulated to belong to the class:

$$\psi = \check{\psi}(c, \nabla c, \mathbf{D}), \quad (\text{A.27})$$

and the concentration-based chemical potential-like quantity is defined as

$$\check{\mu} = \frac{\partial \check{\psi}}{\partial c} - \frac{1}{\rho} \operatorname{div} \left(\rho \frac{\partial \check{\psi}}{\partial \nabla c} \right). \quad (\text{A.28})$$

Analogously to the other cases, we work with constitutive classes for stress tensor $\mathbf{T} = \check{\mathbf{T}}$, diffusive flux $\mathbf{h}^v = \check{\mathbf{h}}^v$ and mass flux $\gamma = \check{\gamma}$ and postpone their specification. We postulate:

$$\frac{d}{dt} \mathcal{E}(c, \mathbf{v}) = \mathcal{W}(\mathcal{R}(t)) - \mathcal{D}(\mathcal{R}(t)), \quad (\text{A.29})$$

with the same interpretation of $\mathcal{W}(\mathcal{R}(t))$ and $\mathcal{D}(\mathcal{R}(t)) \geq 0$. We proceed with the evaluation of the evolution of the energy (A.26) and find via Reynolds transport theorem and the mass balance equation (2.28):

$$\begin{aligned} \frac{d}{dt} \int_{\mathcal{R}(t)} \rho \check{\psi} \, dv &= \int_{\mathcal{R}(t)} \rho \dot{\check{\psi}} \, dv \\ &= \int_{\mathcal{R}(t)} \rho \left(\frac{\partial \check{\psi}}{\partial c} \dot{c} + \frac{\partial \check{\psi}}{\partial \nabla c} \cdot (\nabla c)^\cdot + \frac{\partial \check{\psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} \right) \, dv. \end{aligned} \quad (\text{A.30})$$

On the account of the relation

$$(\nabla c)^\cdot = \nabla(\dot{c}) - (\nabla c)^T \nabla \mathbf{v}, \quad (\text{A.31})$$

we arrive at

$$\begin{aligned} \frac{d}{dt} \int_{\mathcal{R}(t)} \rho \check{\psi} \, dv &= \int_{\mathcal{R}(t)} \rho \check{\mu} \dot{c} - \left(\nabla c \otimes \frac{\partial \rho \check{\psi}}{\partial \nabla c} \right) : \nabla \mathbf{v} + \rho \frac{\partial \check{\psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} \, dv \\ &\quad + \int_{\partial \mathcal{R}(t)} \dot{c} \rho \frac{\partial \check{\psi}}{\partial \nabla c} \cdot \boldsymbol{\nu} \, da. \end{aligned} \quad (\text{A.32})$$

By substituting the concentration equation (2.35b), we find

$$\begin{aligned} \frac{d}{dt} \int_{\mathcal{R}(t)} \rho \check{\psi} \, dv &= \int_{\mathcal{R}(t)} \nabla \check{\mu} \cdot \check{\mathbf{J}}^v - \left(\nabla c \otimes \frac{\partial \rho \check{\psi}}{\partial \nabla c} \right) : \nabla \mathbf{v} + \rho \frac{\partial \check{\psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} + \check{\mu} \check{\gamma} \, dv \\ &\quad + \int_{\partial \mathcal{R}(t)} \left(-\check{\mu} \check{\mathbf{J}}^v + \dot{c} \rho \frac{\partial \check{\psi}}{\partial \nabla c} \right) \cdot \boldsymbol{\nu} \, da. \end{aligned} \quad (\text{A.33})$$

Following the same procedure as in Sec. 3, we find

$$\begin{aligned} \frac{d}{dt} \mathcal{E}(c, \mathbf{v}) &= \int_{\partial \mathcal{R}(t)} \left(\mathbf{v}^T \check{\mathbf{T}} - \check{\mu} \check{\mathbf{J}}^v + \dot{c} \frac{\partial \rho \check{\psi}}{\partial \nabla c} \right) \cdot \boldsymbol{\nu} \, da \\ &\quad - \int_{\mathcal{R}(t)} \left(\check{\mathbf{T}} + \nabla c \otimes \frac{\partial \rho \check{\psi}}{\partial \nabla c} \right) : \nabla \mathbf{v} - \nabla \check{\mu} \cdot \check{\mathbf{J}}^v \\ &\quad - \frac{\partial \rho \check{\psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} - \check{\mu} \check{\gamma} \, dv. \end{aligned} \quad (\text{A.34})$$

Via substitution of the partition

$$\check{\mathbf{T}} = \check{\mathbf{T}}_0 - \check{p} \mathbf{I} \quad \text{with} \quad \check{\mathbf{T}}_0 := \check{\mathbf{T}} + \check{p} \mathbf{I}, \quad (\text{A.35})$$

and the identity (A.20) with pressure \check{p} we identify the rate of work and dissipation term:

$$\mathcal{W}(\mathcal{R}(t)) = \int_{\partial \mathcal{R}(t)} \left(\mathbf{v}^T (\check{\mathbf{T}}_0 - \check{p} \mathbf{I}) - (\check{\mu} + \check{p} \beta) \check{\mathbf{J}}^v + \dot{c} \frac{\partial \rho \check{\psi}}{\partial \nabla c} \right) \cdot \boldsymbol{\nu} \, da, \quad (\text{A.36a})$$

$$\begin{aligned} \mathcal{D}(\mathcal{R}(t)) &= \int_{\mathcal{R}(t)} \left(\check{\mathbf{T}}_0 + \nabla c \otimes \frac{\partial \rho \check{\psi}}{\partial \nabla c} \right) : \nabla \mathbf{v} \\ &\quad - \nabla (\check{\mu} + \check{p} \beta) \cdot \check{\mathbf{J}}^v - \frac{\partial \rho \check{\psi}}{\partial \mathbf{D}} : \dot{\mathbf{D}} - (\check{\mu} + \check{p} \beta) \check{\gamma} \, dv, \end{aligned} \quad (\text{A.36b})$$

where we recall $\beta = -[\![\rho]\!]/(\rho_1 \rho_2)$. Using the same argument as before, we work with the reduced class of the free energy:

$$\psi = \check{\psi}(c, \nabla c), \quad (\text{A.37})$$

and obtain the modeling restriction

$$\left(\check{\mathbf{T}}_0 + \nabla c \otimes \rho \frac{\partial \check{\psi}}{\partial \nabla c} \right) : \nabla \mathbf{v} - \nabla (\check{\mu} + \check{p} \beta) \cdot \check{\mathbf{J}}^v - (\check{\mu} + \check{p} \beta) \check{\gamma} \geq 0. \quad (\text{A.38})$$

Based on the modeling restriction we restrict to the constitutive classes:

$$\mathbf{T} = \check{\mathbf{T}}(c, \nabla c, \check{\mu}, \nabla \check{\mu}, \mathbf{D}, \check{p}), \quad (\text{A.39a})$$

$$\mathbf{J}^v = \check{\mathbf{J}}^v(c, \nabla c, \check{\mu}, \nabla \check{\mu}, \nabla \check{p}), \quad (\text{A.39b})$$

$$\gamma = \check{\gamma}(c, \check{\mu}, \check{p}). \quad (\text{A.39c})$$

Appendix B. Proofs of Lemmas 3.1 and Theorem 3.1

Lemma B.1. (Relations between chemical potential-like quantities) *The chemical potential-like quantities are related via the following identities:*

$$\hat{\mu} = \rho \hat{\check{\mu}} + \hat{\psi}[\rho], \quad (\text{B.1a})$$

$$\check{\mu} = \rho \check{\check{\mu}} - \beta \rho^2 \check{\psi}, \quad (\text{B.1b})$$

$$\check{\mu} = \frac{\rho^2}{\rho_1 \rho_2} \hat{\mu}. \quad (\text{B.1c})$$

Proof. The first identity (B.1a) follows by (i) substituting the first identification from (3.33) into $\hat{\mu}$, (ii) applying the chain rule and (iii) using the identity $\hat{\rho}'(\phi) = [\rho]$.

The second identity (B.1b) follows in a similar fashion from the third identification in (3.33) by noting that $\check{\rho}'(c) = -\beta \rho^2$.

The last identity (B.1c) is a direct consequence of the chain rule for variational derivatives. Alternatively, one can apply a direct computation which we present here. Substituting the identification (3.33) into $\check{\mu}$ and expanding the derivatives by the chain rule gives:

$$\begin{aligned} \check{\mu} &= \frac{\partial \hat{\Psi}(\phi(c), \phi'(c) \nabla c)}{\partial c} - \operatorname{div} \frac{\partial \hat{\Psi}(\phi(c), \phi'(c) \nabla c)}{\partial \nabla c} \\ &= \frac{\partial \hat{\Psi}(\phi(c), \phi'(c) \nabla c)}{\partial \phi} \phi'(c) + \frac{\partial \hat{\Psi}(\phi(c), \phi'(c) \nabla c)}{\partial \nabla \phi} \cdot \nabla c \phi''(c) \\ &\quad - \operatorname{div} \left(\frac{\partial \hat{\Psi}(\phi(c), \phi'(c) \nabla c)}{\partial \nabla \phi} \right) \phi'(c) - \frac{\partial \hat{\Psi}(\phi(c), \phi'(c) \nabla c)}{\partial \nabla \phi} \cdot \nabla (\phi'(c)) \\ &= \frac{\partial \hat{\Psi}(\phi(c), \phi'(c) \nabla c)}{\partial \phi} \phi'(c) - \operatorname{div} \left(\frac{\partial \hat{\Psi}(\phi(c), \phi'(c) \nabla c)}{\partial \nabla \phi} \right) \phi'(c) \\ &= \hat{\mu} \phi'(c). \end{aligned} \quad (\text{B.2})$$

Recalling the relation between ϕ and c , (2.33), completes the proof. \square

Theorem B.1. (Modeling restrictions independent of order parameter and free energy type) *The restrictions in (3.32) are equivalent. In particular, we have the*

relations between the pressures of the various modeling choices:

$$\hat{p} = \hat{\hat{p}} + \hat{\psi} \{ \rho \}, \quad (\text{B.3a})$$

$$\check{p} = \check{\check{p}} + \rho \check{\check{\psi}}, \quad (\text{B.3b})$$

$$\check{p} = \hat{p} + \hat{\mu} \phi. \quad (\text{B.3c})$$

Proof. We start off by showing equivalence of the restrictions (3.32a) and (3.32b). Consider the term in brackets in the first member of (3.32a) in isolation. Substituting the identification of the equivalence classes and the identity (3.37), and applying Lemma 3.1 provides:

$$\begin{aligned} & \hat{\mathbf{T}}_0 + \nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} + (\hat{\mu} \phi - \hat{\Psi}) \mathbf{I} \\ &= \hat{\mathbf{T}} + \nabla \phi \otimes \frac{\partial \hat{\Psi}}{\partial \nabla \phi} + (\hat{p} + \hat{\mu} \phi - \hat{\Psi}) \mathbf{I} \\ &= \hat{\mathbf{T}} + \nabla \phi \otimes \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} + (\hat{p} + \rho \hat{\mu} \phi + \hat{\psi} \llbracket \rho \rrbracket \phi - \rho \hat{\psi}) \mathbf{I} \\ &= \hat{\mathbf{T}} + \nabla \phi \otimes \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} + \rho \hat{\mu} \phi \mathbf{I} + (\hat{p} - \hat{\psi} \{ \rho \}) \mathbf{I} \\ &= \hat{\mathbf{T}}_0 + \nabla \phi \otimes \rho \frac{\partial \hat{\psi}}{\partial \nabla \phi} + \rho \hat{\mu} \phi \mathbf{I} + (\hat{p} - \hat{\hat{p}} - \hat{\psi} \{ \rho \}) \mathbf{I}. \end{aligned} \quad (\text{B.4})$$

Next, we consider the term in brackets in the second and third members in isolation and apply Lemma 3.1:

$$\begin{aligned} \hat{\mu} + \alpha \hat{p} &= \rho \hat{\mu} + \hat{\psi} \llbracket \rho \rrbracket + \alpha \hat{p} \\ &= \rho \hat{\mu} + \alpha (\hat{p} - \hat{\psi} \{ \rho \}) \\ &= \rho \hat{\mu} + \alpha \hat{\hat{p}} + \alpha (\hat{p} - \hat{\hat{p}} - \hat{\psi} \{ \rho \}). \end{aligned} \quad (\text{B.5})$$

Equivalence of (3.32a) and (3.32b) results with the observation that the last members in (B.4) and (B.5) vanish with the variable transformation:

$$\hat{p} = \hat{\hat{p}} + \hat{\psi} \{ \rho \}. \quad (\text{B.6})$$

We proceed with showing the equivalence of (3.32c) and (3.32d). Applying Lemma 3.1 and the identity (3.37) to the term in brackets in the first member of (3.32c) provides:

$$\begin{aligned} \check{\mathbf{T}}_0 + \nabla c \otimes \frac{\partial \check{\Psi}}{\partial \nabla c} - \check{\Psi} \mathbf{I} &= \check{\mathbf{T}} + \nabla c \otimes \frac{\partial \check{\Psi}}{\partial \nabla c} + (\check{p} - \check{\Psi}) \mathbf{I} \\ &= \check{\mathbf{T}} + \nabla c \otimes \rho \frac{\partial \check{\check{\psi}}}{\partial \nabla c} + (\check{p} - \rho \check{\check{\psi}}) \mathbf{I} \\ &= \check{\mathbf{T}}_0 + \nabla \phi \otimes \rho \frac{\partial \check{\check{\psi}}}{\partial \nabla \phi} + (\check{p} - \check{\check{p}} - \rho \check{\check{\psi}}) \mathbf{I}. \end{aligned} \quad (\text{B.7})$$

Using Lemma 3.1 the term in brackets in the second and third members of (3.32c) takes the form:

$$\frac{\check{\mu}}{\rho} + \beta\check{p} = \check{\mu} - \beta(\rho\check{\psi} - \check{p}) = \check{\mu} + \beta\check{p} + \beta(\check{p} - \check{p} - \rho\check{\psi}). \quad (\text{B.8})$$

Equivalence of (3.32c) and (3.32d) is a consequence of the observation that the last members in (B.7) and (B.8) disappear with the variable transformation:

$$\check{p} = \check{\check{p}} + \rho\check{\psi}. \quad (\text{B.9})$$

We finalize the proof by showing equivalence of (3.32a) and (3.32c). Again on the account of Lemma 3.1 and (3.37), the term in brackets in the first member of (3.32a) can be written as

$$\begin{aligned} \hat{\mathbf{T}}_0 + \nabla\phi \otimes \frac{\partial\hat{\Psi}}{\partial\nabla\phi} + (\hat{\mu}\phi - \hat{\Psi})\mathbf{I} \\ &= \hat{\mathbf{T}} + \nabla\phi \otimes \frac{\partial\hat{\Psi}}{\partial\nabla\phi} + (\hat{p} + \hat{\mu}\phi - \hat{\Psi})\mathbf{I} \\ &= \check{\mathbf{T}} + \nabla c \otimes \frac{\partial\check{\Psi}}{\partial\nabla c} + (\hat{p} + \hat{\mu}\phi - \check{\Psi})\mathbf{I} \\ &= \check{\mathbf{T}}_0 + \nabla c \otimes \frac{\partial\check{\Psi}}{\partial\nabla c} - \check{\Psi}\mathbf{I} + (\hat{p} - \check{p} + \hat{\mu}\phi)\mathbf{I}. \end{aligned} \quad (\text{B.10})$$

Finally, we focus on the second and third member in (3.32a). On the account of Lemma 3.1, we write the sequence of identities:

$$\begin{aligned} \zeta(\hat{\mu} + \alpha\hat{p}) &= \frac{\{\rho\}}{\rho_1\rho_2}\hat{\mu} + \beta\hat{p} = \frac{\{\rho\}}{\rho^2}\check{\mu} - \beta\hat{\mu}\phi + \beta\check{p} + \beta(\hat{p} - \check{p} + \hat{\mu}\phi) \\ &= \frac{\{\rho\}}{\rho^2}\check{\mu} - \frac{\{\rho\}}{\rho^2}\check{\mu}\phi + \beta\check{p} + \beta(\hat{p} - \check{p} + \hat{\mu}\phi) \\ &= \frac{\check{\mu}}{\rho} + \beta\check{p} + \beta(\hat{p} - \check{p} + \hat{\mu}\phi). \end{aligned} \quad (\text{B.11})$$

Elimination of the last terms in (B.10) and (B.11) via the variable transformation

$$\check{p} = \hat{p} + \hat{\mu}\phi, \quad (\text{B.12})$$

and recalling (3.35) and (3.36) concludes the proof. \square

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