A study on the thermodynamic consistency of the Park–Paulino–Roesler (PPR) cohesive fracture model

Daniel W. Spring, Oliver Giraldo-Londoño, Glaucio H. Paulino

Department of Civil and Environmental Engineering, University of Illinois at Urbana-Champaign, Urbana, IL, United States
School of Civil and Environmental Engineering, Georgia Institute of Technology, Atlanta, GA, United States

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ABSTRACT

Although the Park–Paulino–Roesler (PPR) potential-based cohesive zone fracture model was not derived based on a thermodynamics consistency principle, we investigate the thermodynamic consistency of the PPR model under conditions of loading, unloading and reloading. First, we present a general anisotropic Helmholtz free energy function. Then, we reformulate the PPR model into the anisotropic Helmholtz form, and investigate its consistency and the various unloading/reloading relations which have been proposed for use with the model. By recasting the PPR model into the Helmholtz form, we illustrate that the PPR cohesive potential, while not designed with thermodynamic consistency in mind, is thermodynamically consistent under the pure loading conditions for which it was designed (as expected). We also demonstrate that the unloading/reloading relations, which are commonly used with the PPR model, are not thermodynamically consistent; however, through our investigation, we develop a new coupled unloading/reloading relation, which maintains the thermodynamic consistency of the PPR cohesive model. The considerations addressed in this paper are aimed at achieving a better understanding of the PPR model and other models of similar nature.

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1. Introduction

Thermodynamically consistent cohesive models are derived from a potential function [1]. There are two primary classes of potential functions. In the first class, the potential is a function of the displacement jump. This form is prevalent in phenomenological cohesive models, and is the class to which the Park–Paulino–Roesler (PPR) cohesive model belongs [2]. Potential functions of this form do not contain a built-in unloading/reloading relation. This is an attractive feature of the model, as any desired feature can be easily incorporated in the basic PPR framework [3,4]. However, without the inclusion of external history parameters and an unloading/reloading relation, these types of models are reversible. The reversibility of the potential function leads to the commonly applied critique that models of this type do not satisfy the second law of thermodynamics (i.e. what is done can be undone) [1,5–8]. Alternatively, some researchers use a thermodynamic potential, specifically a Helmholtz free energy function, to derive a thermodynamically consistent cohesive model [9–11]. In this class of functions, the potential depends on both the displacement jump and a set of internal variables related to the deformation history. Thus, the unloading/reloading relation is intrinsic to the potential, and all states of loading, unloading and reloading are defined by a single function. In the following discussion, we will recast the PPR model into the Helmholtz form, to illustrate that the PPR cohesive potential, while not designed with thermodynamic consistency in mind, is thermodynamically consistent under the pure loading conditions for which it was designed. We will also demonstrate that the unloading/reloading relations, which are commonly used with the PPR model, are not thermodynamically consistent; however, we can extract a new unloading/reloading relation from the Helmholtz form which is thermodynamically consistent.

The remainder of the paper is organized as follows. In the next section, we present a general form for the anisotropic Helmholtz free energy function. In Section 3, we reformulate the PPR potential function as an anisotropic Helmholtz function. In Section 4, we investigate the thermodynamic consistency of the PPR cohesive model under conditions of loading and unloading/reloading. Based on our investigation, we develop a new coupled unloading/reloading relation in Section 3; which maintains the thermodynamic consistency of the PPR cohesive model. In Section 6, we present an example of an interface undergoing
mixed-mode loading and unloading; which highlights the significance of the new formulation. Finally, we provide some concluding remarks in Section 7.

2. The anisotropic Helmholtz free energy function

Here, we outline a framework for generating a thermodynamically consistent constitutive relation for inelastic materials [12]. We consider the inelasticity to be associated with stiffness degradation of the material, and thus work within the framework of damage mechanics, i.e., damage is herein defined as a reduction in the material’s secant stiffness [13]. For most generality, only a couple of assumptions are made. First, following the frequently applied assumption in stress-strain based constitutive models [14,15], we assume a decomposition of the potential energy into shear and normal components. Second, we assume that damage may be captured by a set of scalar-valued damage parameters [1]. With these two assumptions, the general Helmholtz free energy function, $\Psi$, for anisotropic materials, takes the following form:

$$\Psi = \sum_{i=1}^{n} (1 - d_i) \Psi_i = (1 - d_n) \Psi_n + (1 - d_t) \Psi_t,$$  \hspace{1cm} (1)

where $\Psi_n$ is the energy related to normal separation and $\Psi_t$ is the energy related to tangential (or shear) separation. Moreover, $d_n$ and $d_t$ are scalar damage parameters related to the evolution of damage in the normal and shear directions, respectively. The scalar damage parameters are continuous and take values between 0 and 1 (i.e. $d_n \in [0, 1]$ and $d_t \in [0, 1]$) where 0 indicates no damage, and 1 indicates complete damage. In a related investigation, Mosler and Scheider [1] included a third assumption: that the different damage mechanisms in the normal and shear directions are multiplicatively coupled. Thus, they proposed the following form for the anisotropic Helmholtz free energy function:

$$\Psi = \prod_{i=1}^{n} \left(1 - d_i^m\right) \Psi_i.$$  \hspace{1cm} (2)

However, we note that this is a subset of the more general form (1), where:

$$(1 - d_n) = \left(1 - d_n^m\right) \left(1 - d_n^n\right), \quad (1 - d_t) = \left(1 - d_t^m\right) \left(1 - d_t^n\right).$$  \hspace{1cm} (3)

In both cases, the effective damage parameters are continuous and vary between 0 and 1.

In the context of a purely mechanical theory, we can use the classical Coleman and Noll procedure [16] to derive a thermodynamically consistent, anisotropic constitutive relation. In this setting, the Clausius–Duhem dissipation inequality reads:

$$D = w - \dot{\Psi} \geq 0,$$  \hspace{1cm} (4)

where $w$ is the stress power and $\dot{\Psi}$ is the time derivative of the Helmholtz free energy function. In the current context of cohesive zone models, the stress power is written as [1]:

$$\dot{w} = T : [[\dot{u}]].$$  \hspace{1cm} (5)

where $T$ is the traction vector, and $[[\dot{u}]]$ is the time derivative of the displacement jump. Inserting the expressions for the stress power and the time derivative of the general Helmholtz free energy function (1) into the dissipation inequality (4), we obtain:

$$D = \dot{w} - \dot{\Psi} = T : [[\dot{u}]] - (1 - d_n) \frac{\partial \Psi_n}{\partial [\dot{[u]]]} : [[\dot{u}]]$$

$$+ \Psi_n d_n - (1 - d_t) \frac{\partial \Psi_t}{\partial [\dot{[u]]]} : [[\dot{u}]] + \Psi_t \dot{t}$$

$$\geq 0.$$  \hspace{1cm} (6)

From the above expression, we can define the following relations for the cohesive tractions:

$$T_n = (1 - d_n) \frac{\partial \Psi_n}{\partial [\dot{[u]]]} \quad \text{and} \quad T_t = (1 - d_t) \frac{\partial \Psi_t}{\partial [\dot{[u]]]}$$  \hspace{1cm} (7)

which results in the following form for the dissipation inequality:

$$D = \Psi_n d_n + \Psi_t \dot{t} \geq 0.$$  \hspace{1cm} (8)

Since the elastic energies are assumed to be non-negative, the dissipation inequality, and thus the second law of thermodynamics [16], is automatically satisfied if the damage parameters are monotonically increasing:

$$d_n \geq 0 \quad \text{and} \quad \dot{t} \geq 0.$$  \hspace{1cm} (9)

Therefore, in order for an anisotropic constitutive relation to be thermodynamically consistent, the above constraints need to be met.

3. Reformulating the PPR potential as an anisotropic Helmholtz function

To demonstrate the thermodynamic consistency of the PPR potential function [2], under the pure loading conditions it was designed for, we recast the potential function into the form of the Helmholtz free energy function (1). With some algebraic maneuvering, the energies, $\Psi_n$ and $\Psi_t$, consistent with the PPR model, take the form:

$$\Psi_n = \frac{1}{2} E_n \Delta_n^2 \quad \text{and} \quad \Psi_t = \frac{1}{2} E_t \Delta_t^2,$$  \hspace{1cm} (10)

where $\Delta_n$ and $\Delta_t$ are the normal and tangential crack opening widths, respectively. The variables $E_n$ and $E_t$ shown above are initial stiffness parameters in the normal and tangential directions, respectively. These parameters are consistent with the original PPR cohesive model proposed by Park et al. [2] (see Appendix A), and are defined as:

$$E_n = - \frac{\Gamma_n}{\sigma_n^m} \left( \frac{m}{\alpha} \right)^{m-1} \left( m + \alpha \right) \left[ \Gamma_n \left( \frac{n}{\beta} \right) + (\phi - \phi_n) \right],$$  \hspace{1cm} (11)

$$E_t = - \frac{\Gamma_t}{\sigma_t^m} \left( \frac{n}{\beta} \right)^{n-1} \left( n + \beta \right) \left[ \Gamma_n \left( \frac{m}{\alpha} \right) + (\phi - \phi_t) \right],$$  \hspace{1cm} (12)

where $\delta_n$ and $\delta_t$ are the final crack opening widths in the normal and tangential directions:

$$\delta_n = \frac{\phi_n}{\sigma^\text{max}} \alpha \lambda_n (1 - \lambda_n)^{\sigma - 1} \left( \frac{\alpha}{m} \right)^{m-1} \left( \frac{m}{\alpha} \lambda_n + 1 \right)^{m-1},$$  \hspace{1cm} (13)

$$\delta_t = \frac{\phi_t}{\sigma^\text{max}} \beta \lambda_t (1 - \lambda_t)^{\beta - 1} \left( \frac{\beta}{n} \right)^{n-1} \left( \frac{n}{\beta} \lambda_t + 1 \right)^{n-1}.$$  \hspace{1cm} (14)

[1] We assume here, for simplicity, that the Helmholtz free energy is not a function of any structural tensor. In the remainder of the paper, we consider the crack faces to be parallel, limiting our investigation to the constitutive component of the model and eliminating the need to include energetically consistent conjugate stresses [1].
Moreover, $\alpha$ and $\beta$ are shape parameters which control the softening shape of the traction–separation relation in the normal and shear directions, respectively. Both $\alpha$ and $\beta$ are $\geq 1$. If $\alpha$ (or $\beta$) is set equal to 2, the softening relation is almost linear, whereas if $\alpha$ (or $\beta$) is less than or greater than 2, the relation is concave or convex, respectively. Moreover, the non-dimensional exponents, $m$ and $n$, are evaluated from the shape parameters and the initial slope indicators ($\lambda_n$, $\lambda_t$):

$$m = \frac{\alpha(\alpha - 1)\lambda_n^2}{1 - \alpha \lambda_n^2}, \quad n = \frac{\beta(\beta - 1)\lambda_t^2}{1 - \beta \lambda_t^2}$$

(15)

where the initial slope indicators relate the final crack opening widths to the crack opening widths at peak cohesive stress. The energy constants $\Gamma_n$ and $\Gamma_t$ are related to the normal and tangential fracture energies [2]. When the normal and tangential fracture energies are different ($\phi_n \neq \phi_t$), the energy constants are:

$$\Gamma_n = (-\phi_n)^{(\phi_n - \phi_t)/(|\phi_n - \phi_t|)} \left(\frac{\alpha}{m}\right)^m, \quad \Gamma_t = (-\phi_t)^{(\phi_n - \phi_t)/(|\phi_n - \phi_t|)} \left(\frac{\beta}{n}\right)^n$$

(16)

where the Macaulay bracket is defined such that $\langle x \rangle = (|x| + x)/2$. When the normal and tangential fracture energies are equal ($\phi_n = \phi_t$), the energy constants are:

$$\Gamma_n = -\phi_n \frac{\alpha}{m}, \quad \Gamma_t = \left(\frac{\beta}{n}\right)^n$$

(17)

The damage parameters consistent with the PPR model are functions of two internal history parameters ($\kappa_n, \kappa_t$) [12], and take the form:

$$d_n(\kappa_n, \kappa_t) = 1 - \frac{T_n(\kappa_n, \kappa_t)}{E_n \kappa_n}, \quad d_t(\kappa_n, \kappa_t) = 1 - \frac{T_t(\kappa_n, \kappa_t)}{E_t \kappa_t}$$

(18)

(19)

where

$$T_n(\kappa_n, \kappa_t) = \frac{\Gamma_n}{\delta_n} \left[ m \left( 1 - \frac{\kappa_n}{\delta_n} \right)^{\alpha - 1} \left( \frac{m + \kappa_n}{\alpha \delta_n} \right)^{m-1} - \alpha \left( 1 - \frac{\kappa_n}{\delta_n} \right)^{-1} \left( \frac{m + \kappa_n}{\alpha \delta_n} \right)^{m} \right] \times \left[ \Gamma_t \left( 1 - \frac{\kappa_t}{\delta_t} \right) \left( \frac{n}{\beta} + \frac{\kappa_t}{\delta_t} \right)^n + (\phi_t - \phi_n) \right]$$

(20)

$$T_t(\kappa_n, \kappa_t) = \frac{\Gamma_t}{\delta_t} \left[ n \left( 1 - \frac{\kappa_t}{\delta_t} \right)^{\beta - 1} \left( \frac{n + \kappa_t}{\beta \delta_t} \right)^{n-1} - \beta \left( 1 - \frac{\kappa_t}{\delta_t} \right)^{-1} \left( \frac{n + \kappa_t}{\beta \delta_t} \right)^{n} \right] \times \left[ \Gamma_n \left( 1 - \frac{\kappa_n}{\delta_n} \right) \left( \frac{m}{\alpha} + \frac{\kappa_n}{\delta_n} \right)^{m} + (\phi_n - \phi_t) \right]$$

(21)

Substituting Eqs. (10), (18) and (19) into Eq. (7), the cohesive tractions can be compactly written as:

$$T_n = T_n(\kappa_n, \kappa_t) \frac{\Delta_n}{\kappa_n}, \quad T_t = T_t(\kappa_n, \kappa_t) \frac{\Delta_t}{\kappa_t}$$

(22)

The internal history parameters $\kappa_n$ and $\kappa_t$ represent the maximum normal opening and absolute tangential opening in the history of loading, respectively. Because the damage parameters are constrained to be irreversible in Eq. (9), to satisfy the dissipation inequality in Eq. (4), the history parameter must also be irreversible, i.e.:

$$\kappa_n = \max \left\{ \Delta_n \right\} \quad \text{and} \quad \kappa_t = \max \left\{ \left| \Delta_t \right| \right\}$$

(23)

In a numerical setting, these parameters satisfy the following relations from one increment (i) to the next (i+1):

$$\kappa_n^{i+1} = \max \left\{ \kappa_n^i, \Delta_n^{i+1} \right\} \quad \text{and} \quad \kappa_t^{i+1} = \max \left\{ \kappa_t^i, \left| \Delta_t^{i+1} \right| \right\}$$

(24)

Therefore, when unloading occurs, i.e., $\kappa_n^{i+1} = \kappa_n^i$ and $\kappa_t^{i+1} = \kappa_t^i$, the corresponding damage parameter does not decrease. In the pure loading scenario, $\kappa_n^{i+1} = \Delta_n^{i+1}$ and $\kappa_t^{i+1} = \left| \Delta_t^{i+1} \right|$; and the tractions resulting from the Helmholtz free energy function are equivalent to those from the PPR potential function (see Appendix A).

There are three distinct scenarios where unloading occurs, as illustrated in Fig. 1. In order for an unloading/reloading relation to be thermodynamically consistent, the damage parameters during unloading need to satisfy the constraints in Eq. (9) in all three scenarios. In the following section, we will plot the evolution of the damage parameters for the PPR model under conditions of loading and unloading/reloading. In the case of unloading/reloading, we will consider both the coupled and uncoupled relations proposed for use with the PPR model [17].

4. Assessing the thermodynamic consistency of the PPR cohesive model

As discussed previously, the primary critique of potential-based models which only depend on the displacement jump is the reversibility of the model without the inclusion of external history parameters. Thus, it is not surprising that, in conditions of pure loading, cohesive models of this type are indeed thermodynamically consistent (i.e. damage monotonically increases in the model). To show this, we plot a typical traction–separation relation derived from the Helmholtz form of the PPR model, and the corresponding evolution of damage parameters in Fig. 2. As illustrated, each damage parameter begins at 0 and monotonically increase to 1 as the separation ($\Delta_n, \Delta_t$) increases to the final crack opening width ($\delta_n, \delta_t$). If this were not the case (i.e. the damage parameter, and thus the damage in the material, decreased), it would indicate that the material has self-healed; a behavior not accounted for in the original formulation of the PPR cohesive model.

Fig. 1. At each state (reference point) of the loading progress, there are four possible scenarios for loading to progress. The pure loading condition constitutes the white region ($\Delta_n^\text{max} \geq \Delta_t^\text{max}$ and $\Delta_t^\text{max} \geq \Delta_t^\text{max}$); partial unloading constitutes the two light gray regions ($\Delta_n^\text{max} < \Delta_t^\text{max} \geq \Delta_t^\text{max}$; or $\Delta_n^\text{max} \geq \Delta_t^\text{max} < \Delta_t^\text{max}$); and pure unloading constitutes the dark gray region ($\Delta_n^\text{max} < \Delta_t^\text{max} < \Delta_t^\text{max}$). The reference point corresponds to the point of intersection between the historical maximum normal separation and the historical maximum tangential separation.
Since the unloading/reloading relations which have been proposed for use with the PPR model are both phenomenological and independent of the original potential that it is based upon, their thermodynamic consistency is not guaranteed (as expected). Here, we examine both the uncoupled and coupled unloading/reloading relations [17] and compute their effective damage parameters, to demonstrate the issues inherent in their formulation. A summary of the coupled and uncoupled unloading/reloading relations

### Table 1
Summary of the coupled, uncoupled and thermodynamically consistent unloading/reloading relations for the PPR cohesive model.

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<td>Criterion</td>
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<td>$\kappa = \max \left{ \sqrt{\Delta s^2 + \Delta t^2} \right}$</td>
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<td>$\sqrt{\Delta s^2 + \Delta t^2} &lt; \kappa$</td>
<td>$\Delta_s &lt; \Delta_k$, $\Delta_t &lt; \Delta_k$</td>
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<td>$\Delta s = \frac{\Delta s \kappa}{\sqrt{\Delta s^2 + \Delta t^2}}$, and $\Delta t = \frac{\Delta s \kappa}{\sqrt{\Delta s^2 + \Delta t^2}}$</td>
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### Normal region

![Normal region diagram](image)

### Tangential region

![Tangential region diagram](image)
is listed in Table 1. First, in the uncoupled case, unloading in the normal direction is viewed independently from that in the tangential direction. Typical uncoupled unloading tractions are illustrated in Fig. 3(a) and (b). Because this unloading relation is not extracted from the cohesive model itself, there is no explicit form assumed for the damage parameters; however, we can compute the effective damage parameters numerically, as illustrated in Fig. 3(c) and (d).

In Fig. 3(c), the normal damage parameter monotonically increases with normal separation. However, when simultaneous unloading in the normal and tangential directions occurs (the pure unloading scenario in Fig. 1), the damage parameter decreases with decreasing tangential separation. This is not a physically realistic response. In the absence of self-healing, the damage parameters should not decrease as the material unloads. Similarly, the tangential damage parameter monotonically increases with tangential separation, as illustrated in Fig. 3(d), but decreases with normal separation when simultaneous unloading occurs in both directions.

Finally, for the coupled unloading/reloading case, unloading in the normal direction is viewed as dependent on that in the tangential direction. Typical coupled unloading tractions are illustrated in Fig. 4(a) and (b). This unloading relation is also phenomenological, and not extracted from the cohesive model itself (i.e., there is no explicit form assumed for the damage parameters), thus we compute the effective damage parameters numerically, and illustrate them in Fig. 4(c) and (d). From the figures, it is clear that neither the normal damage parameter nor the tangential damage parameter demonstrate monotonicity in the region of unloading.

In a thermodynamically consistent model, we would expect each of the damage parameters to display monotonically increasing behavior, regardless of the unloading condition. It is well-known that this behavior is not possible with an uncoupled unloading/reloading relation, thus the unloading/reloading relation must be coupled. In the following section, we derive a new, thermodynamically consistent, coupled unloading/reloading relation for use with the PPR cohesive model.

5. A thermodynamically consistent unloading/reloading relation

In the previous section, we verified that neither the coupled nor the uncoupled unloading/reloading relations, which have been proposed for use with the PPR cohesive model, are thermodynamically consistent (as expected). However, since the anisotropic Helmholtz form of the PPR model describes all conditions of loading, it contains an intrinsic unloading/reloading relation. The corresponding thermodynamically consistent relations, extracted from the Helmholtz form of the PPR model, are depicted in Fig. 5. The damage parameters, equivalent to those computed using Eqs. (18) and (19), are illustrated in Fig. 5(c) and (d). As shown, both the normal and tangential damage parameters monotonically increase under both normal and tangential separation. When unloading/reloading occurs simultaneously in both directions, both damage parameters remain constant; which is the only thermodynamically consistent form of these parameters in this scenario (when self-healing is neglected), as per the dissipation inequality (8).

In addition to the cohesive traction vectors in Eq. (22), we outline the form of the material tangent stiffness matrix for the thermodynamically consistent unloading/reloading relation:

\[
D^\nu(\Delta_n, \Delta_t) = \begin{bmatrix}
D^\nu_{nn} & D^\nu_{nt} \\
D^\nu_{tn} & D^\nu_{tt}
\end{bmatrix} = \begin{bmatrix}
\frac{\partial T_n}{\partial \Delta_n} & \frac{\partial T_t}{\partial \Delta_t} \\
\frac{\partial T_t}{\partial \Delta_n} & \frac{\partial T_t}{\partial \Delta_t}
\end{bmatrix}.
\] (25)

For each of the three distinct unloading scenarios, illustrated in Fig. 1, the components \(D^\nu_{nn}, D^\nu_{nt}, D^\nu_{tn}, \) and \(D^\nu_{tt}\) of the tangent matrix become:

1) If \(\Delta_n < \kappa_n\) and \(\Delta_t < \kappa_t\)

\[
D^\nu_{nn} = T_{nt}(\kappa_n, \kappa_t) \frac{1}{\kappa_n}, \quad D^\nu_{tt} = T_{tt}(\kappa_n, \kappa_t) \frac{1}{\kappa_t}
\] (26)
Fig. 4. Depiction of the coupled unloading/reloading relation for the PPR cohesive model: (a) traction in the normal direction; (b) traction in the tangential direction; (c) effective damage parameter in the normal direction; (d) effective damage parameter in the tangential direction. The cohesive parameters are: \( \phi_n = 100 \text{ N/m}, \phi_t = 200 \text{ N/m}, \sigma_{\text{max}} = 40 \text{ MPa}, \tau_{\text{max}} = 30 \text{ MPa}, \alpha_n = 5, \beta_n = 1.3, \alpha_t = 1, \beta_t = 1, \lambda_n = 0.2, \lambda_t = 0.1, \Delta_{\text{max}} = 0.35 \delta_n, \) and \( \Delta_{\text{max}} = 0.25 \delta_t. \)

Fig. 5. Depiction of the thermodynamically consistent unloading/reloading relation for the PPR cohesive model, derived from the Helmholtz form of the model: (a) traction in the normal direction; (b) traction in the tangential direction; (c) damage parameter in the normal direction; and (d) damage parameter in the tangential direction. The cohesive parameters are: \( \phi_n = 100 \text{ N/m}, \phi_t = 200 \text{ N/m}, \sigma_{\text{max}} = 40 \text{ MPa}, \tau_{\text{max}} = 30 \text{ MPa}, \alpha_n = 5, \beta_n = 1.3, \alpha_t = 1, \beta_t = 1, \lambda_n = 0.2, \lambda_t = 0.1, \Delta_{\text{max}} = 0.35 \delta_n, \) and \( \Delta_{\text{max}} = 0.25 \delta_t. \)

\[
\begin{align*}
D_{rn}^n &= 0 & D_{rt}^n &= 0 \\
D_{rn}^t &= T_n(\kappa_n, \Delta_t) \frac{1}{\kappa_n} & D_{rt}^n &= D_{nt}(\Delta_n, \kappa_t) \\
D_{rt}^n &= T_t(\kappa_n, \Delta_t) \left( \frac{\Delta_n}{\kappa_n} \right) & D_{rt}^t &= T_t(\Delta_n, \kappa_t) \frac{1}{\kappa_t} \\
D_{rn}^t &= D_{nt}(\kappa_n, \Delta_t) & D_{rt}^t &= D_{nt}(\Delta_n, \kappa_t) \left( \frac{\Delta_t}{\kappa_t} \right)
\end{align*}
\]

where \( D_{rn}(\Delta_n, \Delta_t) = \partial T_n/\partial \Delta_t \) and \( D_{rt}(\Delta_n, \Delta_t) = \partial T_t/\partial \Delta_n \) [18,19].
6. Example: mixed-mode sinusoidal loading and unloading

In this section, we evaluate the response of a cohesive interface during nontrivial mixed-mode loading and unloading. The crack faces are assumed to remain parallel to one another, however, the crack opening in the normal direction is assumed to follow a relation independent of that in the tangential direction. The respective opening separations are assumed to follow the relations:

\[
\Delta_n = 2\sin(0.15\tau) \quad \text{and} \quad \Delta_t = 3.5\sin(0.12\tau),
\]

where \( \tau \) is a time-like parameter \( (0 \leq \tau \leq 100) \), as illustrated in Figs. 6 and 7. The response of the interface is investigated for each of the three unloading/reloading relations summarized in Table 1. The cohesive parameters are selected as: \( \phi_n = \phi_t = 100 \, \text{N/m} \), \( \sigma_{\text{max}} = 40 \, \text{MPa} \), \( \tau_{\text{max}} = 15 \, \text{MPa} \), \( \lambda_n = 0.1 \), \( \lambda_t = 0.2 \), \( \alpha = 5 \), \( \beta = 1.3 \), \( \alpha_n = 1 \), and \( \beta_t = 1 \). As noted previously, during unloading/reloading of the interface, damage is expected to remain constant, and thus, the slope of the traction–separation relation is also expected to remain constant.

For each unloading/reloading relation, the computed normal and tangential traction–separation relations are illustrated in Figs. 8 and 9, respectively. From the results, it is clear that the only unloading/reloading relation able to capture the expected response is the one derived from the thermodynamically consistent formulation. The coupled and uncoupled unloading/reloading relations produce significant deviations from the expected results, due to non-monotonic damage accumulation in these relations. Note that, even though the violation of monotonicity is small (see Figs. 3 and 4), the impact of the violation on the global response is large.

The unexpected unloading/reloading behavior, observed using both the uncoupled and coupled relations, could lead to incorrect global behavior if one were to use these relations for problems which experience a significant amount of mixed-mode crack face unloading. One prominent example of this is in the case of fatigue damage, wherein the cohesive zone experiences repeated cycles of loading, unloading and reloading. Fatigue damage has been modeled with the cohesive zone concept, and is generally introduced through unloading/reloading relations [20–22].
aforementioned thermodynamically consistent unloading/reloading relation might be a viable alternative to address fatigue modeling.

7. Concluding remarks

This paper addresses the thermodynamic consistency of the PPR potential-based cohesive fracture model. We demonstrate that the PPR model, while not designed with thermodynamic consistency in mind, is thermodynamically consistent under the pure loading conditions for which it was designed. We do this by proposing a general form for the anisotropic Helmholtz free energy function, and then by recasting the PPR cohesive model to this form. Through the reformulation of the PPR model, we are able to demonstrate that damage accumulates in the model in an irreversible manner when the crack opening width increases monotonically. Alternatively, when the crack undergoes unloading/reloading, we verify that the commonly applied coupled and uncoupled unloading/reloading relations produce self-healing behavior within the formulation. To address this issue, we present a new coupled unloading/reloading relation which maintains the thermodynamic consistency of the PPR model.

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Appendix A. Comparison between the present thermodynamically consistent formulation and the original PPR model

Here, the proposed thermodynamically consistent form of the PPR model is compared to the original form of the model, presented in [2], for the case of monotonic loading. First, we present the derivation of the initial stiffness parameters $E_n$ and $E_i$ in Eqs. (11) and (12). Next, the damage parameters $d_n$ and $d_i$ in Eqs. (18) and (19) are simplified for the case of monotonic loading. Lastly, the proposed thermodynamically consistent model is compared to the original PPR model. The result of this investigation illustrates that the proposed thermodynamically consistent form of the PPR model is equivalent to the original model under monotonic loading conditions.

The initial stiffness parameters $E_n$ and $E_i$, in Eqs. (11) and (12), are obtained by evaluating $D_m(0, 0)$ and $D_n(0, 0)$ (from Eq. 35) in [18]:

\[
E_n = D_m(0, 0) = \frac{\Gamma_n}{\sigma_n} \left[ (m^2 - m) \left(\frac{m}{\alpha} \right)^{m-2} + (\alpha^2 - \alpha) \left(\frac{m}{\alpha} \right)^m \right] \\
- 2m\alpha \left(\frac{m}{\alpha} \right)^{m-1} \left[ \Gamma_i \left(\frac{n}{\beta} \right)^n + (\phi_t - \phi_n) \right] \\
= - \frac{\Gamma_n}{\sigma_n} \left(\frac{m}{\alpha} \right)^{m-1} \left( m + \alpha \right) \left[ \Gamma_i \left(\frac{n}{\beta} \right)^n + (\phi_t - \phi_n) \right], \tag{A.1}
\]

\[
E_i = D_n(0, 0) = \frac{\Gamma_i}{n} \left[ (n^2 - n) \left(\frac{n}{\beta} \right)^{n-2} + (\beta^2 - \beta) \left(\frac{n}{\beta} \right)^n \right] \\
- 2n\beta \left(\frac{n}{\beta} \right)^{n-1} \left[ \Gamma_n \left(\frac{m}{\alpha} \right)^m + (\phi_n - \phi_i) \right] \\
= - \frac{\Gamma_i}{n} \left(\frac{n}{\beta} \right)^{n-1} \left( n + \beta \right) \left[ \Gamma_n \left(\frac{m}{\alpha} \right)^m + (\phi_n - \phi_i) \right]. \tag{A.2}
\]

The damage parameters $d_n$ and $d_i$ in Eqs. (18) and (19) account for the degradation of the cohesive tractions as the displacement jump increases. Conceptually, the proposed damage parameters $d_n$ and $d_i$ account for difference between the elastic stresses in the normal and tangential directions and the corresponding inelastic stresses predicted by the original formulation of the model [2]. If monotonic loading is assumed, the internal history parameters become $\kappa_n = \Delta_n$ and $\kappa_i = |\Delta_i|$. Replacing these expressions for $\kappa_n$ and $\kappa_i$ in Eqs. (18) and (21), we obtain:

\[
1 - d_n \left( \Delta_n, |\Delta_i| \right) = \frac{\Gamma_n}{E_n |\Delta_i| \sigma_n} \left[ n \left( 1 - \frac{|\Delta_i|}{\beta_i} \right)^\beta \left( \frac{n}{\beta} + \frac{|\Delta_i|}{\beta_i} \right)^{n-1} \right] \\
- \alpha \left( 1 - \frac{\Delta_n}{\delta_n} \right)^{-1} \left( \frac{m}{\alpha} + \frac{\Delta_n}{\delta_n} \right)^m \left[ \Gamma_i \left( 1 - \frac{|\Delta_i|}{\beta_i} \right)^\beta \left( \frac{n}{\beta} + \frac{|\Delta_i|}{\beta_i} \right)^n + (\phi_n - \phi_i) \right]. \tag{A.3}
\]

\[
1 - d_i \left( \Delta_n, |\Delta_i| \right) = \frac{\Gamma_i}{E_i |\Delta_i| \sigma_n} \left[ n \left( 1 - \frac{|\Delta_i|}{\beta_i} \right)^\beta \left( \frac{n}{\beta} + \frac{|\Delta_i|}{\beta_i} \right)^{n-1} \right] \\
- \beta \left( 1 - \frac{\Delta_n}{\delta_n} \right)^{-1} \left( \frac{m}{\alpha} + \frac{\Delta_n}{\delta_n} \right)^m \left[ \Gamma_n \left( 1 - \frac{\Delta_n}{\delta_n} \right)^\alpha \left( \frac{m}{\alpha} + \frac{\Delta_n}{\delta_n} \right)^m + (\phi_n - \phi_i) \right]. \tag{A.4}
\]

Substituting Eqs. (A.3), (A.4) and (10) into Eq. (7) yields:

\[
T_n(\Delta_n, \Delta_i) = \frac{\Gamma_n}{\delta_n} \left[ n \left( 1 - \frac{\Delta_n}{\delta_n} \right)^\alpha \left( \frac{m}{\alpha} + \frac{\Delta_n}{\delta_n} \right)^{m-1} \right] \\
- \alpha \left( 1 - \frac{\Delta_n}{\delta_n} \right)^{-1} \left( \frac{m}{\alpha} + \frac{\Delta_n}{\delta_n} \right)^m \left[ \Gamma_i \left( 1 - \frac{|\Delta_i|}{\beta_i} \right)^\beta \left( \frac{n}{\beta} + \frac{|\Delta_i|}{\beta_i} \right)^n + (\phi_n - \phi_i) \right], \tag{A.5}
\]
\[ T_i(\Delta n, \Delta t) = \frac{\Gamma_i}{\delta t} \left[ n \left( 1 - \frac{\Delta t}{\delta t} \right)^\beta \left( \frac{n}{\beta} + \frac{\Delta t}{\delta t} \right)^{n-1} \right] \]

\[ - \beta \left( 1 - \frac{\Delta t}{\delta t} \right)^{\beta-1} \left( \frac{n}{\beta} + \frac{\Delta t}{\delta t} \right)^n \]

\[ \times \left[ \Gamma_n \left( 1 - \frac{\Delta n}{\delta n} \right)^a \left( \frac{m}{\alpha} + \frac{\Delta n}{\delta n} \right)^m + (\phi_n - \phi_i) \right] \frac{\Delta t}{\delta t} \]

\[ (A.6) \]

As expected, these expressions are identical to the cohesive tractions predicted by the original PPR model [2], demonstrating the equivalence between the proposed thermodynamically consistent formulation and the original PPR model formulation under conditions of monotonic loading.

**Appendix B. Nomenclature**

- \( \alpha \): parameter controlling the shape of the cohesive softening curve in the normal direction
- \( \alpha_v \): parameter controlling the shape of the unloading/reloading curve in the normal direction
- \( \frac{\delta_n}{\delta t} \): conjugate normal final crack opening width
- \( \beta \): parameter controlling the shape of the cohesive softening curve in the tangential direction
- \( \beta_v \): parameter controlling the shape of the unloading/reloading curve in the tangential direction
- \( \Delta_n \): normal separation along the fracture surface
- \( \Delta_t \): tangential separation along the fracture surface
- \( \Delta_{n+i} \): normal separation along the fracture surface at numerical increment \( i+1 \)
- \( \Delta_{t+i} \): tangential separation along the fracture surface at numerical increment \( i+1 \)
- \( \Delta_{n+i,1} \): tangential final crack opening width
- \( \Delta_n \): tangential separation along the fracture surface at numerical increment \( i \)
- \( \Delta_t \): tangential separation along the fracture surface at numerical increment \( i \)
- \( \Delta_{n+i,1} \): tangential separation along the fracture surface at numerical increment \( i+1 \)
- \( \Delta_{t+i,1} \): tangential final crack opening width
- \( \Delta_{n+i} \): tangential separation along the fracture surface at numerical increment \( i \)
- \( \Delta_{t+i} \): tangential separation along the fracture surface at numerical increment \( i \)
- \( \Psi \): time derivative of the Helmholtz free energy function
- \( \Gamma_n \): energy constant in the PPR model
- \( \Gamma_t \): energy constant in the PPR model
- \( \kappa_n \): internal history parameter in the normal direction
- \( \kappa_t \): internal history parameter in the tangential direction
- \( \lambda_n \): parameter controlling the hardening slope of the PPR model in the normal direction
- \( \lambda_t \): parameter controlling the hardening slope of the PPR model in the normal direction
- \( \dot{\cdot} \): time derivative
- \( \{\cdot\} \): Macauley bracket

**References**


