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A new constitutive model for polymeric matrices: Application to biomedical materials

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Abstract: Semi-crystalline polymeric composites are increasingly used as bearing material in the biomedical sector, mainly because of their specific mechanical properties and the new advances in 3D printing technologies that allows for customised devices. Among these applications, total or partial prostheses for surgical purposes must consider the influence of temperature and loading rate. This paper proposes a new constitutive model for semi-crystalline polymers, commonly used as matrix material in a wide variety of biomedical composites, that enables reliable predictions under a wide range of loading conditions. The most recent models present limitations to predict the non-linear behaviour of the polymer when it is exposed to large deformations at high strain rates. The proposed model takes into account characteristic behaviours of injected and 3D printed thermoplastic polymers such as material hardening due to strain rate sensitivity, thermal softening, thermal expansion and combines viscoelastic and viscoplastic responses. These viscous-behaviours are relevant for biomedical applications where temperature evolution is expected during the deformation process due to heat generation induced by inelastic dissipation, being essential the thermo-mechanical coupling consideration. The constitutive model is formulated for finite deformations within a thermodynamically consistent framework. Additionally, the model is implemented in a finite element code and its parameters are identified for two biomedical polymers: ultra-high-molecular-weight-polyethylene (UHMWPE) and high density polyethylene (HDPE). Finally, the influence of viscous behaviours on dynamic deformation of semi-crystalline polymeric matrices is analysed. This constitutive model predicts the mechanical behaviour of semi-crystalline polymeric matrices for a wide range of strain rate and temperature conditions, allowing for the optimisation of new composite materials potentially used as effective joint replacement prostheses.

Keywords: Biomedical materials; Polymer–matrix composites (PMCs); Constitutive model; UHMWPE composites.

1. Introduction

Semi-crystalline polymers and their composites are currently used in a wide range of applications that are subjected to dynamic loading [1-4]. In several applications, the dynamic deformation of these polymers is an essential consideration, such as in the biomedical, the aeronautical and the automotive industries [5]. Among the materials employed as matrix in a wide variety of composites, these polymers stand out due to their wear resistance, biocompatibility and good mechanical properties [6-8]. In the biomedical sector, injection moulded and 3D printed polymers are used as bearing material for prostheses [9-12], highlighting their use in hip and knee joints [13]. In such applications, the pure polymeric material still presents mechanical limitations in terms of severe material degradation, Fig.1, and damage arising from its low hardness, its

relatively low stiffness, as well as high creep rate under load [14,15]. In order to enhance its mechanical properties, a wide range of fillers are used: carbon fibres, hard particles, hydroxyapatite, boron carbide and graphene oxide nanoparticles [2,16,17]. The advance in the 3D printed polymeric composites field requires, then, the proper understanding of the mechanical behaviour of matrix materials. In this regard, semi-crystalline polymers behave in a complex manner that presents a strong non-linearity and depends on many factors such as strain rate [18], temperature [19], stress state [20], large deformations and plastic flow [21]. Therefore, the proper understanding of the polymeric matrix is essential for the study of the mechanical response of the composite. To this end, constitutive models for reliable predictions of the mechanical behaviour of such materials must account for strain rate, temperature and stress state dependences combining both viscoelastic and viscoplastic behaviours.

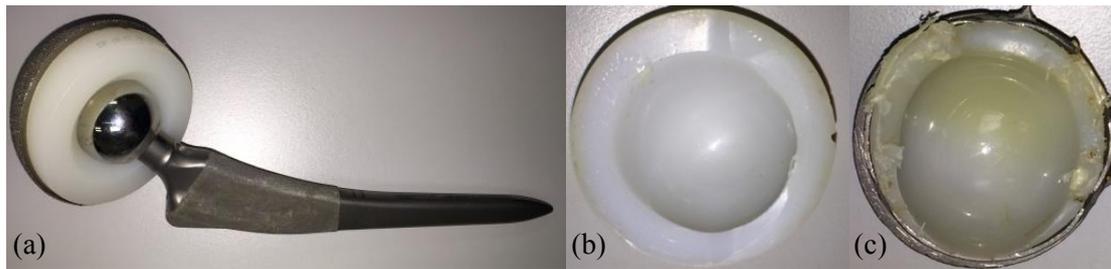


Fig. 1. (a) Real polyethylene-metal hip prosthetic. (b)-(c) Polyethylene acetabular cup removed from patients due to excessive material erosion. Materials provided by Dra. I. Gómez Arrayás (Hospital Ruber Internacional of Madrid).

Moreover, fibre reinforced semi-crystalline composites are also used in structural components for applications that require excellent impact performance [22]. In this regard, when semi-crystalline polymers are used in aeronautical -for example, as bulk space filters- and automotive applications -for example, as vehicle crashworthiness structural components-, other effects must be also accounted for faithfully describing its mechanical behaviour. The exposure of this polymer to high strain rates leads to variations in its response [23]. In this sense, there is a positive dependence of elastic modulus and yield stress on strain rate [18,24]. In addition, there is a strong coupling between the mechanical and thermal behaviours of the polymer which becomes more relevant at high strain rates [12,19,25]. When the polymer exhibits large deformations, there is a heat generation due to inelastic-viscous dissipation that induces an increase in material temperature leading to thermal softening. These thermal effects have a strong influence on the mechanical behaviour of semi-crystalline polymers at high strain rates and should be considered by constitutive models for providing reliable predictions of their behaviour. Under dynamic conditions, the understanding of the composite matrix becomes essential when strain rate and temperature play an important role in its behaviour, resulting in variations of the composite mechanical response [25-29].

Many constitutive approaches have been developed for modelling the mechanical behaviour of semi-crystalline polymers and, particularly, of several types of polyethylene. The first approaches, as the work developed by Hughes [30], described the mechanical response of these polymers by classical isotropic, rate-independent plasticity using Mises yield criterion. Following this line, several authors have used constitutive models to analyse the mechanical response of ultra-high-molecular-weight-polyethylene (UHMWPE) components of hip and knee prosthesis under different loading conditions, where a Mises yield surface is associated with a flow rule followed by isotropic hardening [31-33]. However, these models cannot provide reliable predictions of the non-linear behaviour of the polymer when it is exposed to large deformations.

In addition, these models present considerable limitations when the structural component deforms at high strain rates. In order to develop more robust and reliable constitutive models for semi-crystalline polymers, some authors have proposed viscoelastic-viscoplastic rheological models combining springs and dashpots to capture the inelastic response [34,35]. These models allow for taking into account the non-linear response through the constitutive definition of the springs and the rate-dependent response through the flow rule definition of the dashpots. In this regard, Bergström et al. [36] developed a constitutive model, called hybrid model, for predicting the large strain time-dependent behaviour of UHMWPE. More recently, based on previous models [37,38], Ayoub et al. [39] presented a constitutive model for polyethylene that includes a visco-hyperelastic network resistance acting in parallel with a viscoelastic-viscoplastic intermolecular resistance, where the amorphous and crystalline phases are taken under consideration. Moreover, Garcia-Gonzalez et al. [19] proposed a constitutive model for semi-crystalline polymers formulated in finite deformations where temperature dependence is additionally taken into account.

For reliable modelling of a group of semi-crystalline polymeric matrices is necessary to take into account not only viscoplasticity but also viscoelasticity. The model developed herein considers strain rate and temperature dependences, pressure sensitivity, thermal expansion and thermo-mechanical coupling considering temperature evolution due to inelastic dissipation from viscoelastic and viscoplastic deformation. Regarding temperature dependency, this influence is incorporated for both intermolecular and network stretching resistances. The constitutive equations are formulated in finite deformations within a thermodynamically consistent framework. The constitutive framework is implemented in a VUMAT subroutine for the commercial finite element solver Abaqus/Explicit [40]. The model parameters are identified for UHMWPE and for high density polyethylene (HDPE), two widely used materials in orthopaedics applications, from experimental data reported by Brown et al. [18]. Moreover, the constitutive model is suitable for predicting the mechanical behaviour of a wide variety of polymeric matrices. The application of the model to the study of stretching tests on polymers has demonstrated the importance of taking into account viscous behaviours and thermo-mechanical coupling. This study shows how specific deformation mechanisms govern the mechanical response of polymers and can lead to global or local deformations. The model developed herein is physically motivated on the deformation mechanisms behind the mechanical behaviour of semi-crystalline polymeric matrices and is observed to faithfully describe their mechanical response under a wide range of loading conditions. The work aims at providing a proper understanding of the mechanical behaviour of polymeric matrices, allowing for the development of new composite materials, especially when dealing with dynamic applications.

2. Description of the constitutive model

Semi-crystalline polymeric matrices often exhibit complex viscous effects where both viscoelastic and viscoplastic behaviours are combined. In addition, their stress-strain response can be physically interpreted as the combination of overcoming an intermolecular resistance which is increased by the development of strain-induced crystallization; and a network resistance caused by molecular orientation. The intermolecular resistance may exhibit rate-dependency that can be described by viscoplasticity. The network resistance is commonly defined as purely elastic but it may also exhibit rate-dependency that can be potentially described by viscoelasticity. In addition, viscous contributions to the mechanical behaviour of thermoplastic polymers lead to

temperature increments due to inelastic dissipation and, consequently, induce thermal softening in the material behaviour.

Motivated on the evidence introduced above, this work proposes a constitutive model that incorporates viscous behaviours in the mechanical response of semi-crystalline thermoplastic materials. The model is based on three constitutive branches according to the rheological scheme shown in Fig. 2. Here, the intermolecular resistance (I) introduces a rate- and temperature-dependent part that is defined by a Neo-Hookean hyperelastic spring and a nonlinear viscoplastic dashpot. The network resistance (N) introduces a non-linear temperature-dependent hyperelastic part that is defined by a modified eight-chain spring. The third viscous constitutive branch introduces viscoelasticity through a non-linear hyperelastic response which depends on strain rate.

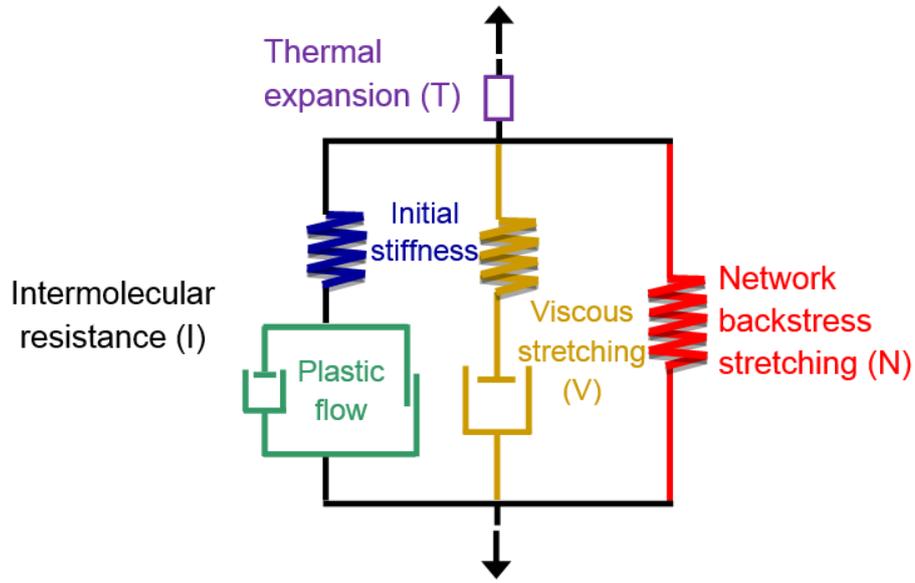


Fig. 2. Rheological scheme of the proposed constitutive model.

According to the arrangement of the rheological model elements depicted in Fig. 2, the total Cauchy stress σ is determined by the contribution of the intermolecular σ_I , the network backstress σ_N and the viscous σ_V resistances:

$$\sigma = \sigma_I + \sigma_N + \sigma_V \quad (1)$$

In order to facilitate the understanding of this section, a summary with the nomenclature used along the formulation is presented in Appendix A.

2.1. Kinematics

The kinematics of the model proposed herein are based on the establishment of five spatial configurations, see Fig. 3. This kinematics goes from an initial reference configuration Ω_0 to a final deformed or current configuration Ω . Three more spatial configurations are defined to allow the determination of the constitutive equations associated to each constitutive element. The first one is referred to as a dilated configuration $\bar{\Omega}$ in which only thermal deformation is accounted for; the second one is referred to as a plastic dilated relaxed configuration $\bar{\bar{\Omega}}$ in which both thermal

and plastic deformations are accounted for; and the third one is referred to as a viscous dilated relaxed configuration $\tilde{\Omega}$ in which both thermal and viscous deformations are accounted for.

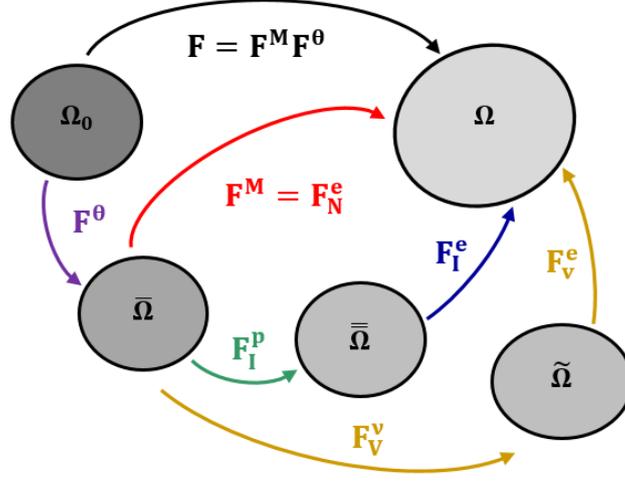


Fig. 3. Kinematics of the model showing the reference or initial configuration Ω_0 , the dilated configuration $\bar{\Omega}$, the plastic dilated relaxed configuration $\bar{\bar{\Omega}}$, the viscous dilated relaxed configuration $\tilde{\Omega}$, and the current or loaded configuration Ω .

According to the kinematics, the deformation gradient \mathbf{F} is broken down into thermal, \mathbf{F}^θ , and mechanical, \mathbf{F}^M , parts [41,42]. Thus, the total deformation gradient reads as:

$$\mathbf{F} = \mathbf{F}^M \mathbf{F}^\theta \quad (2)$$

The mechanical part of the deformation gradient, \mathbf{F}^M , is equivalent for the three constitutive branches according to the rheological model. The mechanical part associated to the network resistance is defined as purely elastic, \mathbf{F}_N^e ; the part associated to the intermolecular resistance is divided into elastic, \mathbf{F}_I^e , and plastic, \mathbf{F}_I^p , components; and the part associated to the viscoelastic resistance is divided into elastic, \mathbf{F}_V^e , and viscous, \mathbf{F}_V^v , components. Therefore, the total deformation gradient can be decomposed depending on the constitutive branch as:

$$\mathbf{F} = \mathbf{F}_I^e \mathbf{F}_I^p \mathbf{F}^\theta = \mathbf{F}_N^e \mathbf{F}^\theta = \mathbf{F}_V^e \mathbf{F}_V^v \mathbf{F}^\theta \quad (3)$$

The velocity gradient \mathbf{I} , in terms of the kinematics associated with the viscous resistance elements, can be written using Eq. (3) as:

$$\mathbf{I} = \dot{\mathbf{F}} \mathbf{F}^{-1} = \mathbf{I}_V^e + \mathbf{F}_V^e \tilde{\mathbf{L}}_V^v \mathbf{F}_V^{-e} + \mathbf{F}_V^e \mathbf{F}_V^v \bar{\mathbf{L}}_V^p \mathbf{F}_V^{-v} \mathbf{F}_V^{-e} \quad (4)$$

where $\mathbf{I}_V^e = \dot{\mathbf{F}}_V^e \mathbf{F}_V^{-e}$ is the elastic component of the velocity gradient in the current configuration and the viscous component $\tilde{\mathbf{L}}_V^v$ can be defined in the viscous dilated relaxed configuration $\tilde{\Omega}$ as:

$$\tilde{\mathbf{L}}_V^v = \dot{\mathbf{F}}_V^v \mathbf{F}_V^{-v} \quad (5)$$

This velocity gradient, as well as the plastic velocity gradient of the intermolecular resistance $\bar{\mathbf{L}}_I^p = \bar{\mathbf{D}}_I^p + \bar{\mathbf{W}}_I^p$, can be decomposed into its symmetric and skew parts by $\tilde{\mathbf{L}}_V^v = \tilde{\mathbf{D}}_V^v + \tilde{\mathbf{W}}_V^v$. In this work, $\tilde{\Omega}$ and $\bar{\Omega}$ are assumed to be invariant to the rigid body rotations of the current configuration, that is $\tilde{\mathbf{W}}_V^v = \bar{\mathbf{W}}_I^p = \mathbf{0}$, and therefore $\tilde{\mathbf{D}}_V^v = \tilde{\mathbf{L}}_V^v$ and $\bar{\mathbf{D}}_I^p = \bar{\mathbf{L}}_I^p$ [43,44]. The thermal contribution is assumed

isotropic according to Bouvard et al. [45], so that \mathbf{F}^θ is spherical and it is possible to assume $\bar{\mathbf{W}}^\theta = \mathbf{0}$.

The kinematics of the intermolecular and network resistances have been defined in agreement with standard approaches from literature [19].

2.2. Thermodynamics

This section provides the modelling assumptions made for the formulation of the Helmholtz free energy function from which the constitutive equations derive. In addition, the thermodynamic consistency is verified from the development of the Clausius-Duhem inequality expressed in the dilated configuration (this is the common configuration for the three constitutive branches).

The Helmholtz free energy function per unit volume in the dilated configuration $\bar{\Psi}$, is defined as the additive combination of the deformation resistances [46,47] as:

$$\bar{\Psi}(\mathbf{C}_I^e, \mathbf{C}_V^e, \mathbf{C}_N^e, \theta) = \bar{\Psi}^I(\mathbf{C}_I^e, \theta) + \bar{\Psi}^V(\mathbf{C}_V^e) + \bar{\Psi}^N(\mathbf{C}_N^e, \theta) \quad (6)$$

where the Helmholtz free energy depends on the elastic right Cauchy-Green deformation tensors $\mathbf{C}_I^e = \mathbf{F}_I^{eT} \mathbf{F}_I^e$, $\mathbf{C}_V^e = \mathbf{F}_V^{eT} \mathbf{F}_V^e$ and $\mathbf{C}_N^e = \mathbf{F}_N^{eT} \mathbf{F}_N^e$, and temperature θ .

From the definition of the Helmholtz free energy $\bar{\Psi} = \bar{e} - \theta \bar{\eta}$, the time derivative of $\bar{\Psi}$ can be calculated as:

$$\dot{\bar{\Psi}} = \frac{\partial \bar{\Psi}}{\partial \mathbf{C}_I^e} : \dot{\mathbf{C}}_I^e + \frac{\partial \bar{\Psi}}{\partial \mathbf{C}_V^e} : \dot{\mathbf{C}}_V^e + \frac{\partial \bar{\Psi}}{\partial \mathbf{C}_N^e} : \dot{\mathbf{C}}_N^e + \frac{\partial \bar{\Psi}}{\partial \theta} \dot{\theta} \quad (7)$$

Using these modelling assumptions for the definition of $\bar{\Psi}$, the Clausius-Duhem inequality can be obtained from the combination of the first and second thermodynamics principles following the procedure used by Garcia-Gonzalez et al. [19] as:

$$\left(\mathbf{F}_I^p \bar{\mathbf{S}}_I \mathbf{F}_I^{pT} - 2 \frac{\partial \bar{\Psi}}{\partial \mathbf{C}_I^e} \right) : \mathbf{F}_I^{eT} \mathbf{d}_I^e \mathbf{F}_I^e + \left(\mathbf{F}_V^v \bar{\mathbf{S}}_V \mathbf{F}_V^{vT} - 2 \frac{\partial \bar{\Psi}}{\partial \mathbf{C}_V^e} \right) : \mathbf{F}_V^{eT} \mathbf{d}_V^e \mathbf{F}_V^e + \left(\bar{\mathbf{S}}_N - 2 \frac{\partial \bar{\Psi}}{\partial \mathbf{C}_N^e} \right) : \mathbf{F}_N^{eT} \mathbf{d}_N^e \mathbf{F}_N^e + \bar{\mathbf{M}}_I : \bar{\mathbf{D}}^p + \bar{\mathbf{M}}_V : \bar{\mathbf{D}}^v + \left(-\frac{\partial \bar{\Psi}}{\partial \theta} - 3f_\theta \bar{\Psi} - \bar{\eta} + f_\theta (\bar{\mathbf{M}}_I + \bar{\mathbf{M}}_V + \bar{\mathbf{M}}_N) : \mathbf{I} \right) \dot{\theta} - \frac{1}{\theta} \bar{\mathbf{Q}}_x \theta \geq 0 \quad (8)$$

where $\bar{\mathbf{S}}_I$, $\bar{\mathbf{S}}_N$ and $\bar{\mathbf{S}}_V$ are respectively the corresponding second Piola-Kirchhoff stress tensors of the intermolecular, network and viscous resistances expressed in the configuration $\bar{\Omega}$ as $\bar{\mathbf{S}}_I = \mathbf{J}^M \mathbf{F}_I^{-M} \boldsymbol{\sigma}_I \mathbf{F}_I^{-MT}$ and $\bar{\mathbf{M}}_I$, $\bar{\mathbf{M}}_N$ and $\bar{\mathbf{M}}_V$ are respectively the Mandel stress tensors of the intermolecular, network and viscous resistances in $\bar{\Omega}$ as $\bar{\mathbf{M}}_i = \mathbf{F}_i^{MT} \mathbf{F}_i^M \bar{\mathbf{S}}_i$ with $i=\{I, N, V\}$. \mathbf{d}_I^e , \mathbf{d}_N^e and \mathbf{d}_V^e are the symmetric part of the velocity gradient tensors in Ω of the intermolecular, the network and the viscous resistances; f_θ is a temperature-dependent function; $\bar{\eta}$ is the specific entropy per unit volume in $\bar{\Omega}$; and $\bar{\mathbf{Q}}$ is the heat flux per unit volume in $\bar{\Omega}$.

Using standard arguments of the Coleman and Noll method [48,49], the second Piola-Kirchhoff stress tensor associated with each constitutive branch and the specific internal entropy per unit volume that satisfy the second law of thermodynamics along arbitrary thermodynamic processes must read as:

$$\bar{\mathbf{S}}_I = \mathbf{F}_I^{-p} 2 \frac{\partial \bar{\Psi}}{\partial \mathbf{C}_I^e} \mathbf{F}_I^{-pT} \quad (9.1)$$

$$\bar{\mathbf{S}}_N = 2 \frac{\partial \bar{\Psi}}{\partial \mathbf{C}_N^e} \quad (9.2)$$

$$\bar{\mathbf{S}}_V = \mathbf{F}_V^{-\nu} 2 \frac{\partial \bar{\Psi}}{\partial \mathbf{C}_V^e} \mathbf{F}_V^{-\nu T} \quad (9.3)$$

$$\bar{\eta} = -\frac{\partial \bar{\Psi}}{\partial \theta} - 3f_\theta \bar{\Psi} + f_\theta (\bar{\mathbf{M}}_I + \bar{\mathbf{M}}_V + \bar{\mathbf{M}}_N) : \mathbf{I} \quad (9.4)$$

2.3. Thermal expansion

The thermal expansion is assumed to be isotropic, being the contribution of the thermal part to the deformation gradient defined in the form:

$$\dot{\mathbf{F}}^\theta = f_\theta \mathbf{F}^\theta \dot{\theta} \quad (10)$$

where $\dot{\theta}$ is the time derivative of current temperature and $f_\theta = \alpha_\theta$ is the thermal expansion coefficient. The temperature evolution equation can be deduced following the methodology used by Garcia-Gonzalez et al. [19] as:

$$\begin{aligned} & \left(\bar{C} + 3f_\theta \bar{e} - f_\theta (\mathbf{F}_I^{pT} \mathbf{C}_I^e \mathbf{F}_I^p : \bar{\mathbf{S}}_I + \mathbf{C}_N^e : \bar{\mathbf{S}}_N + \mathbf{F}_V^{vT} \mathbf{C}_V^e \mathbf{F}_V^v : \bar{\mathbf{S}}_V) \right) \dot{\theta} = \bar{\mathbf{M}}_I : \bar{\mathbf{D}}^p + \bar{\mathbf{M}}_V : \bar{\mathbf{D}}^v \\ & \theta \left[\frac{3}{2} f_\theta \mathbf{F}_I^p \bar{\mathbf{S}}_I \mathbf{F}_I^{pT} - f_\theta \frac{\partial (\mathbf{F}_I^{pT} \mathbf{C}_I^e \mathbf{F}_I^p : \bar{\mathbf{S}}_I)}{\partial \mathbf{C}_I^e} + \frac{1}{2} \frac{\partial (\mathbf{F}_I^p \bar{\mathbf{S}}_I \mathbf{F}_I^{pT})}{\partial \theta} \right] : \dot{\mathbf{C}}_I^e + \theta \left[\frac{3}{2} f_\theta \bar{\mathbf{S}}_N - f_\theta \frac{\partial (\mathbf{C}_N^e : \bar{\mathbf{S}}_N)}{\partial \mathbf{C}_N^e} + \frac{1}{2} \frac{\partial (\bar{\mathbf{S}}_N)}{\partial \theta} \right] : \dot{\mathbf{C}}_N^e + \\ & \theta \left[\frac{3}{2} f_\theta \mathbf{F}_V^v \bar{\mathbf{S}}_V \mathbf{F}_V^{vT} - f_\theta \frac{\partial (\mathbf{F}_V^{vT} \mathbf{C}_V^e \mathbf{F}_V^v : \bar{\mathbf{S}}_V)}{\partial \mathbf{C}_V^e} + \frac{1}{2} \frac{\partial (\mathbf{F}_V^v \bar{\mathbf{S}}_V \mathbf{F}_V^{vT})}{\partial \theta} \right] : \dot{\mathbf{C}}_V^e - \bar{\nabla}_x \bar{\mathbf{Q}} + \bar{\mathbf{R}} \end{aligned} \quad (11)$$

where \bar{C} is the heat capacity per unit volume.

2.4. Intermolecular and network resistances: thermo-viscoplasticity

Intermolecular resistance

The intermolecular contribution to the Cauchy stress is defined by a temperature-dependent Neo-Hookean model as:

$$\boldsymbol{\sigma}_I = \frac{\lambda_0(\theta) \ln(J_I^p) \mathbf{I}}{J_I} + \frac{\mu_0(\theta)}{J_I} (\mathbf{F}_I^e \mathbf{F}_I^{eT} - \mathbf{I}) \quad (12)$$

where λ_0 and μ_0 are the classical Lamé constants linearly depending on temperature through the Young's modulus $E(\theta) = E_{\text{ref}} + E_1(\theta - \theta_{\text{ref}})$ [45,50].

The plastic contribution is activated when a yield criterion $f_I = \bar{\sigma}_{\text{eqI}} - \sigma_T = 0$ is satisfied. Here,

the Rhagava equivalent stress $\bar{\sigma}_{\text{eqI}} = \frac{(\alpha-1)I_{11} + \sqrt{(\alpha-1)^2 I_{11}^2 + 12\alpha J_{21}}}{2\alpha}$ is used to include the pressure dependency in the yield function [51]. Where α is a material parameter describing the pressure sensitivity and $I_{11} = \text{tr} \bar{\mathbf{M}}_I$ and $J_{21} = \frac{1}{2} \bar{\mathbf{M}}_I^{\text{dev}} : \bar{\mathbf{M}}_I^{\text{dev}}$ are stress invariants of the Mandel stress tensor $\bar{\mathbf{M}}_I = \mathbf{C}_I^e \bar{\mathbf{S}}_I$ expressed in the $\bar{\Omega}$. The scalar term $\sigma_T = \sigma_{T0} \left(1 - \left(\frac{\theta - \theta_{\text{min}}}{\theta_{\text{melt}} - \theta_{\text{min}}} \right)^m \right)$ is defined depending on temperature, where σ_{T0} is the value of σ_T at reference temperature in uniaxial tension, m is a temperature sensitivity parameter, θ_{min} is the lowest temperature considered and θ_{melt} is the melting temperature.

The flow equation that describes the evolution of the plastic deformation is expressed in the plastic dilated relaxed configuration $\bar{\Omega}$ as:

$$\bar{\mathbf{L}}_I^p = \dot{\bar{\gamma}}_I^p \bar{\mathbf{r}}_I; \quad \bar{\mathbf{r}}_I = \frac{\partial g_I}{\partial \bar{\mathbf{M}}_I} \quad (13)$$

where $\dot{\bar{\gamma}}_I^p$ is the viscoplastic multiplier, $\bar{\mathbf{r}}_I$ is the gradient of the plastic potential $g_I = \frac{(\beta-1)I_{11} + \sqrt{(\beta-1)^2 I_{11}^2 + 12\beta J_{21}}}{2\beta}$ on $\bar{\Omega}$, with β being a parameter which controls the volumetric plastic strain.

The viscoplastic multiplier $\dot{\bar{\gamma}}_I^p$ is defined depending on the rate-sensitivity parameters $\dot{\epsilon}_0$ and C as:

$$\dot{\bar{\gamma}}_I^p = \begin{cases} 0 & \text{if } f_I \leq 0 \\ \dot{\epsilon}_0 \left\{ \exp \left[\frac{1}{C} \left(\frac{\bar{\sigma}_{eqI}}{\sigma_T} - 1 \right) \right] - 1 \right\} & \text{if } f_I > 0 \end{cases} \quad (14)$$

Network resistance

This part of the model describes a hyperelastic entropic resistance defined by a modification of the originally eight-chain model proposed by Arruda and Boyce [52]. This modification follows the formulation introduced by Anand [53] but including temperature sensitivity. The Cauchy stress associated to this resistance is defined as:

$$\boldsymbol{\sigma}_N = \frac{(C_R + C_\theta(\theta - \theta_{ref})) \bar{\lambda}_L}{3J_N \bar{\lambda}} \mathfrak{L}^{-1} \left(\frac{\bar{\lambda}}{\bar{\lambda}_L} \right) (\mathbf{B}_N^* - \bar{\lambda}^2 \mathbf{I}) \quad (15)$$

where \mathfrak{L}^{-1} is the inverse of the Langevin function, C_R is the initial elastic modulus of the network backstress resistance, C_θ is a material parameter controlling the elastic modulus dependence on temperature and $\bar{\lambda}_L$ is the locking stretch. $\bar{\lambda} = \sqrt{\frac{1}{3} \text{tr}(\mathbf{F}_N^{e*} (\mathbf{F}_N^{e*})^T)}$ is the average total stretch ratio, with $\mathbf{F}_N^{e*} = J_N^{-1/3} \mathbf{F}_N^e$ being the distortional part of \mathbf{F}_N^e and $J_N = \det(\mathbf{F}_N^e)$.

2.5. Viscous resistance: viscoelasticity

This part of the model describes a viscous material response related to the hardening induced by rate-dependent effects on network stretching. To define this constitutive branch, an eight-chain model is defined for the non-linear spring that depends on the elastic part of the deformation gradient associated with the viscous resistance. In addition, a viscous flow rule is introduced to define the contribution of the linear dashpot. This idea was originally proposed by Bergström and Boyce [54] for elastomers modelling. The Cauchy stress associated with the viscous resistance is defined as:

$$\boldsymbol{\sigma}_V = \frac{c_v \bar{\lambda}_{VL}}{3J_V \bar{\lambda}_V} \mathfrak{L}^{-1} \left(\frac{\bar{\lambda}_V}{\bar{\lambda}_{VL}} \right) (\mathbf{B}_V^{e*} - \bar{\lambda}_V^2 \mathbf{I}) \quad (16)$$

where C_v and $\bar{\lambda}_{vL}$ are material parameters and the average viscous stretch ratio $\bar{\lambda}_v$ is calculated as:

$$\bar{\lambda}_v = \sqrt{\frac{1}{3} \text{tr}(\mathbf{B}_v^{e*})} \quad (17)$$

The distortional left Cauchy-Green viscous deformation tensor, \mathbf{B}_v^{e*} , is determined by the distortional part of \mathbf{F}_v^e defined by Eq. (19).

$$\mathbf{B}_v^{e*} = \mathbf{F}_v^{e*} (\mathbf{F}_v^{e*})^T \quad (18)$$

$$\mathbf{F}_v^{e*} = J_v^{-1/3} \mathbf{F}_v^e \quad (19)$$

In order to define the viscous velocity gradient in the viscous dilated relaxed configuration $\tilde{\Omega}$, a viscoelastic flow rule is assumed following the formulation proposed by Bergström [55] as:

$$\tilde{\mathbf{L}}_v = \dot{\gamma}_v^v \tilde{\mathbf{N}}_v; \quad \tilde{\mathbf{N}}_v = \frac{\boldsymbol{\sigma}_v^{\text{dev}}}{\tau_v} \quad (20)$$

where $\dot{\gamma}_v^v$ is the viscoelastic multiplier, $\tilde{\mathbf{N}}_v$ provides the direction of the viscoelastic flow, $\boldsymbol{\sigma}_v^{\text{dev}}$ is the deviatoric part of $\boldsymbol{\sigma}_v$ and $\tau_v = \sqrt{\text{tr}(\boldsymbol{\sigma}_v^{\text{dev}} \boldsymbol{\sigma}_v^{\text{dev}})}$ is the effective stress driving the viscous flow.

The rate equation for viscous flow is given by:

$$\dot{\gamma}_v^v = \frac{\tau_v}{\sigma_{vT}} \quad (21)$$

where σ_{vT} is a material constant.

3. Identification of model parameters and model predictions

This section summarizes the correspondence of the model parameters with the mechanical response of semi-crystalline matrix materials and, then, the identification of the parameters for UHMWPE is presented. In addition, the parameters of the model are also identified for HDPE in order to compare the predictive capability of the model for polymers that present different deformation behaviour.

3.1. Baseline material

Polyethylene is a highly ductile polymer that can be subjected to large deformations without fracture. This polymer presents different commercial forms whose mechanical properties depend on variables such as crystallinity degree, molecular weight and type of branching: HDPE, low density polyethylene (LDPE), cross-linked polyethylene (PEX) and UHMWPE. Among these polyethylene types, UHMWPE is one of the most employed because of its wear resistance, biocompatibility and good mechanical properties [18]. This thermoplastic polymer exhibits yielding-dependence on strain rate and temperature, increasing the yield stress with strain rate and decreasing with temperature. In addition, this material presents strain-hardening after yield. Although UHMWPE has no clear glass transition, a melting temperature of 405-409 K is observed

[22,56,57] and a thermal expansion coefficient of $124.5 \cdot 10^6 \text{ K}^{-1}$ [13]. In addition, these polymers allow for enhancing their mechanical properties through a wide range of fillers [15].

3.2. Correspondence of the model parameters with mechanical response of the material

The parameters of the proposed model present correspondence with the mechanical response of the material. This makes it easier to identify the proper values for a specific polymer. Such correspondence is introduced in terms of the following blocks:

(i) *Linear response*: the model parameters E_{ref} , E_1 and ν determine the initial elastic response of the material depending on temperature.

(ii) *Yield stress*: the parameters σ_{T0} , C , $\dot{\epsilon}_0$ and m define the yield stress of the material. C and $\dot{\epsilon}_0$ determine the yield stress' strain rate sensitivity and m the temperature sensitivity.

(iii) *Viscous response*: the model parameters C_v and $\bar{\lambda}_{vL}$ determine the stress contribution of the spring of the viscous constitutive branch. C_v is related to the initial elastic modulus of viscous resistance and $\bar{\lambda}_{vL}$ is related to the maximum (fully extended) stretch that a molecule can be exposed to. The material parameter σ_{vT} along with the stress state determines the viscoelastic multiplier that governs the flow rule associated with the linear dashpot.

(iv) *Network response*: the network contribution to the stress state is established by the parameters C_R , C_θ , $\bar{\lambda}_L$ and κ . C_R and C_θ determine the initial elastic modulus of network resistance depending on temperature. The parameter $\bar{\lambda}_L$ is related to the maximum stretch that a molecule can be exposed to and κ is a bulk modulus used in applications where only the network contribution is active (e.g. rubber modelling).

(v) *Volumetric plastic strain sensitivity*: the parameter β introduces the sensitivity of the material with inelastic volume change. This parameter can be defined as $\beta = 1$ assuming volume preserving.

(vi) *Stress state sensitivity*: the parameter α represents the relationship between yield stress in compression and tension and determines the pressure sensitivity.

Taking into account the correspondence of the model parameters with the mechanical response of the material, the parameters of the proposed model are identified for UHMWPE. For this identification process, experimental data covering a wide range of strain rate and temperature loading conditions is used. This experimental data and details about the experiments performance are reported by Brown et al. [18]. The model parameters identified from these experiments are provided in Table 1.

As UHMWPE, HDPE also exhibits yielding-dependence on strain rate and temperature. However, while UHMWPE or others polymers like PEX exhibit significant strain hardening after yield, other polymers such as HDPE present flat flow behaviour [18]. In order to compare the predictive capability of the model for different semi-crystalline polymers and to analyse the effects of particular deformation mechanisms on the mechanical response of these polymers, the model parameters are also identified for HDPE, Table 2.

Table 1
Material parameters for UHMWPE.

| Initial elastic properties | | | | | Intermolecular resistance | | | | | | |
|-----------------------------|----------------------|------------------------------------|---------------------------------------|--------------------|---------------------------|-----|-----------------------|------------------------|-----------------------|----------|---------|
| E_{ref} (GPa) | E_1 (MPa/K) | ν | $\dot{\epsilon}_0$ (s ⁻¹) | C | σ_{T0} (MPa) | m | θ_{ref} (K) | θ_{melt} (K) | θ_{min} (K) | α | β |
| 0.541 | -6.5 | 0.46 | 0.1 | 0.048 | 37.5 | 0.7 | 293 | 406 | 198 | 1.0 | 1.0 |
| Viscous resistance | | | | Network resistance | | | | | | | |
| C_V (Pa) | $\bar{\lambda}_{VL}$ | σ_{VT} (MPa) | C_R (MPa) | C_θ (MPa/K) | $\bar{\lambda}_L$ | | | | | | |
| 12.5 | 5 | 0.05 | 23 | -0.15 | 5 | | | | | | |
| General properties | | | | | | | | | | | |
| ρ (kg/m ³) | C_p (J/Kg K) | α_θ (K ⁻¹) | | | | | | | | | |
| 940 | 1900 | $124.5 \cdot 10^{-6}$ | | | | | | | | | |

Table 2
Material parameters for HDPE.

| Initial elastic properties | | | | | Intermolecular resistance | | | | | | |
|-----------------------------|----------------------|------------------------------------|---------------------------------------|--------------------|---------------------------|------|-----------------------|------------------------|-----------------------|----------|---------|
| E_{ref} (GPa) | E_1 (MPa/K) | ν | $\dot{\epsilon}_0$ (s ⁻¹) | C | σ_{T0} (MPa) | m | θ_{ref} (K) | θ_{melt} (K) | θ_{min} (K) | α | β |
| 0.782 | -6.5 | 0.46 | 0.01 | 0.028 | 80 | 0.76 | 293 | 406 | 198 | 1.0 | 1.0 |
| Viscous resistance | | | | Network resistance | | | | | | | |
| C_V (Pa) | $\bar{\lambda}_{VL}$ | σ_{VT} (Pa) | C_R (MPa) | C_θ (MPa/K) | $\bar{\lambda}_L$ | | | | | | |
| (-) | (-) | (-) | 12 | -0.15 | 5 | | | | | | |
| General properties | | | | | | | | | | | |
| ρ (kg/m ³) | C_p (J/Kg K) | α_θ (K ⁻¹) | | | | | | | | | |
| 970 | 1900 | $93.6 \cdot 10^{-6}$ | | | | | | | | | |

3.3. Model predictions

In this section, the model predictions are compared with experimental data for a wide range of strain rate and temperature conditions. The constitutive model proposed herein is implemented in a VUMAT subroutine and applied to a numerical model with the dimensions of the specimens used in compression tests by Brown et al. [18]. This numerical model was defined with C3D8R elements and developed in Abaqus/Explicit.

The expression developed for the determination of temperature evolution, Eq. (11), is reduced taking into account uniquely specific energy due to plastic dissipation of the intermolecular resistance; viscous dissipation of the viscoelastic resistance; and heat conduction. Thermoelastic coupling is neglected in line with published studies for thermoplastic polymers [19,45]. The temperature evolution is thus reduced to:

$$\dot{\theta} = \frac{\bar{M}_I \cdot \bar{D}^P}{\bar{c}} + \frac{\bar{M}_V \cdot \bar{D}^V}{\bar{c}} - \frac{\bar{\nabla}_x \bar{Q}}{\bar{c}} \quad (22)$$

The model predictions show a good agreement with experiments in terms of strain rate and temperature sensitivities [18]. In this regard, Figs. 4 and 5 show the yield stress model predictions depending on strain rate and temperature; and their comparison with the experimental values for UHMWPE and HDPE respectively. This predictive capacity is determined by the intermolecular constitutive branch that accounts for strain rate sensitivity through the viscoplastic flow rule, Eq. (13), and for temperature sensitivity through the term σ_T that introduced thermal softening in the viscoplastic multiplier through Eq. (14).

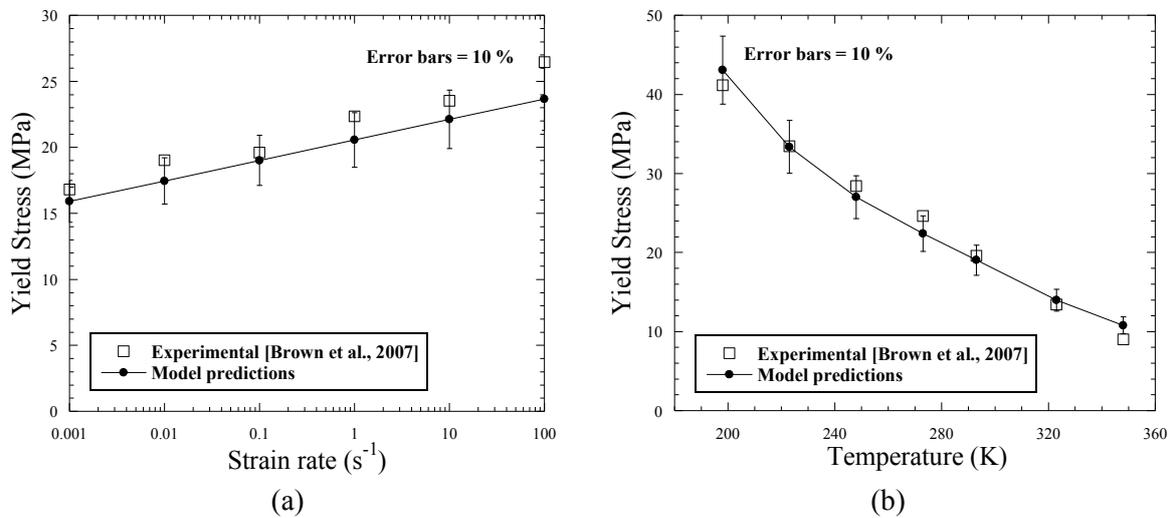
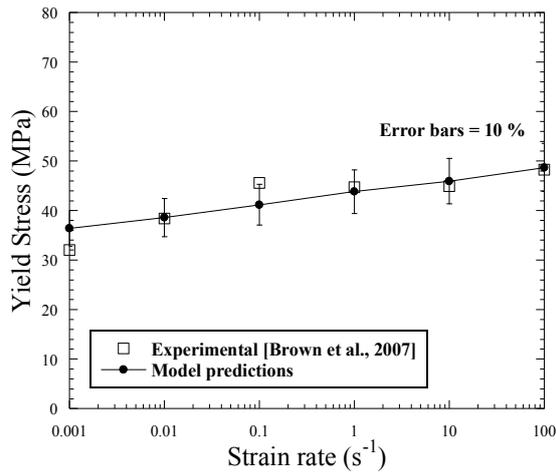
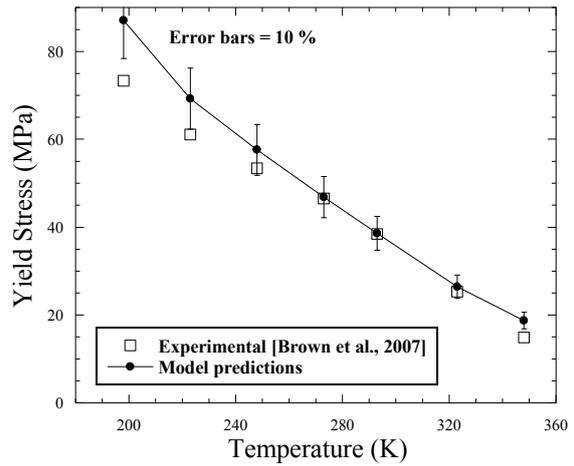


Fig. 4. Experimental data [18] versus model predictions in terms of: (a) strain rate sensitivity; and (b) temperature sensitivity of UHMWPE for uniaxial compression tests.

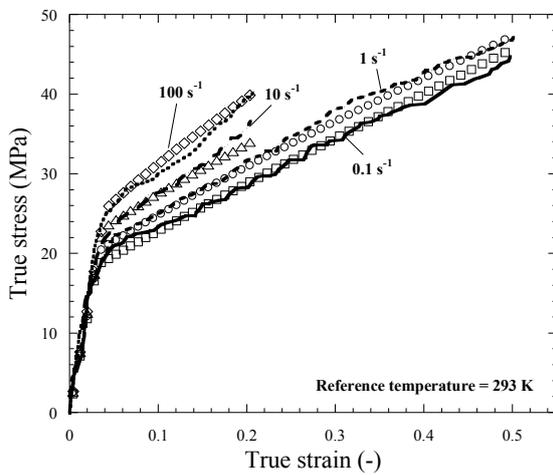


(a)

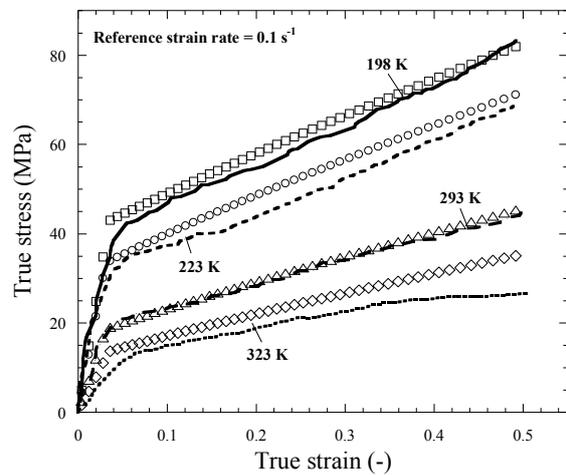


(b)

Fig. 5. Experimental data [18] versus model predictions in terms of: (a) strain rate sensitivity; and (b) temperature sensitivity of HDPE for uniaxial compression tests.

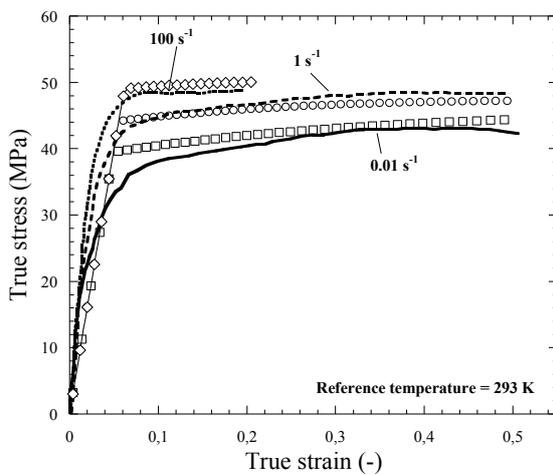


(a)

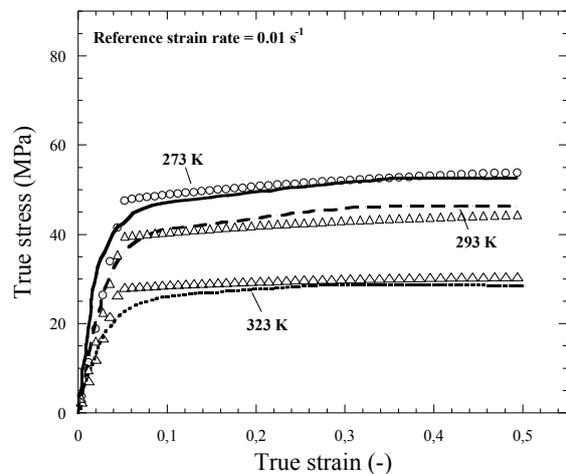


(b)

Fig. 6. Stress-strain experimental curves of UHMWPE [18] versus model predictions for: (a) different strain rates; and (b) different temperatures.



(a)



(b)

Fig. 7. Stress-strain experimental curves of HDPE [18] versus model predictions for: (a) different strain rates; and (b) different temperatures.

Furthermore, the model not only predicts the strain rate and temperature dependences on yield stress, but also on the stress response along the whole deformation process. With the aim of highlighting the predictive capacity of the proposed model, a comparison between model predictions and experimental data [18] is provided in Figs. 6 and 7. These figures show stress-strain curves for UHMWPE and HDPE covering a wide range of strain rate and temperature conditions. A good agreement between model predictions and experiments is observed in these terms. In this regard, the model faithfully predicts the initial slope of the stress-strain curve depending on temperature. This is determined by the definition of $E(\theta)$ in the intermolecular resistance. In addition, in the case of UHMWPE, once the yield point is reached, the stress-strain is mainly governed by the viscous and network resistances, especially at large deformations where their contribution is more relevant. The rate-dependent non-linear behaviour at large deformations is determined by the viscous resistance through the viscoelastic dashpot, Fig. 6a. Moreover, the network resistance introduces temperature dependence in the non-linear behaviour at large deformations, Fig. 6b. Unlike UHMWPE, the viscous flow of HDPE is defined uniquely by the intermolecular resistance since presents a linear behaviour without significant hardening after yielding, Fig. 7. Consequently, the viscoelastic branch was neglected here for HDPE.

Therefore, it can be concluded that the model allows for the introduction of temperature-dependent behaviour associated to both intermolecular and network resistances and couple viscoelastic and viscoplastic deformation mechanisms. This model, calibrated for different biomaterials, is potentially applied to problems in biomedical engineering allowing for the analysis of prosthesis devices that can be subjected to dynamic loading. In addition, the model offers other possibilities in terms of analysis of deformation mechanisms and their implications in semi-crystalline polymeric matrices, also allowing the extrapolation of results to a wider range of temperature and strain rate. In the next section, it is applied to the analysis of viscous behaviours and thermo-mechanical coupling on the deformation process of both polyethylene materials.

4. Analysis of viscous behaviours and thermo-mechanical coupling on necking

This section presents an analysis of the influence of viscous behaviours and deformation mechanisms on necking localization and temperature evolution due to inelastic dissipation. To this end, numerical simulations of tensile tests at different strain rate conditions were conducted.

4.1. Definition of the numerical model

A FE model of tensile specimen with a normalized ASTM D638 Type V geometry and 3 mm of thickness was developed, Fig. 8(a). The specimen presents a gauge section of 9.53 mm in length and 3.18 mm in width. This geometry was chosen in accordance with the specimens used by Mohagheghian et al. [58] in the experimental tensile tests carried out on HDPE and UHMWPE samples. A total number of 176,464 linear elements with reduced integration (C3D8R in Abaqus notation) were employed to mesh the whole region. A higher mesh density was defined in the central zone of the specimen, progressively decreasing the element size along the Z axis to the extremes of the specimen.

Regarding the boundary conditions, a constant stretching velocity is applied on one end face, while the nodes on the opposite face are embedded, Fig. 8 (b). These imposed loading conditions can be formulated as $V_Z(L_t, t) = \dot{\epsilon}_0 \cdot L_0$ and $V_Z(0, t) = 0$ where t is the time, $\dot{\epsilon}_0$ the initial strain

rate, L_t the specimen length and L_0 the gauge length. Note that the stretching velocity was defined assuming that the deformation localizes along the gauge length. Thus, the simulations were performed at two different strain rates, $\dot{\epsilon}_0 = 0.1 \text{ s}^{-1}$ and $\dot{\epsilon}_0 = 100 \text{ s}^{-1}$, allowing for evaluating the influence of this variable on material hardening and on thermo-mechanical coupling. Although an initial temperature of 293 K was defined for all simulations, it evolves along the deformation process due to the consideration of inelastic dissipation following Eq. (22). According to observations of other authors about temperature evolution for semi-crystalline polymers [19,59], Eq. (22) can be reduced neglecting the term associated to thermal conduction for high strain rate conditions assuming adiabatic heating.

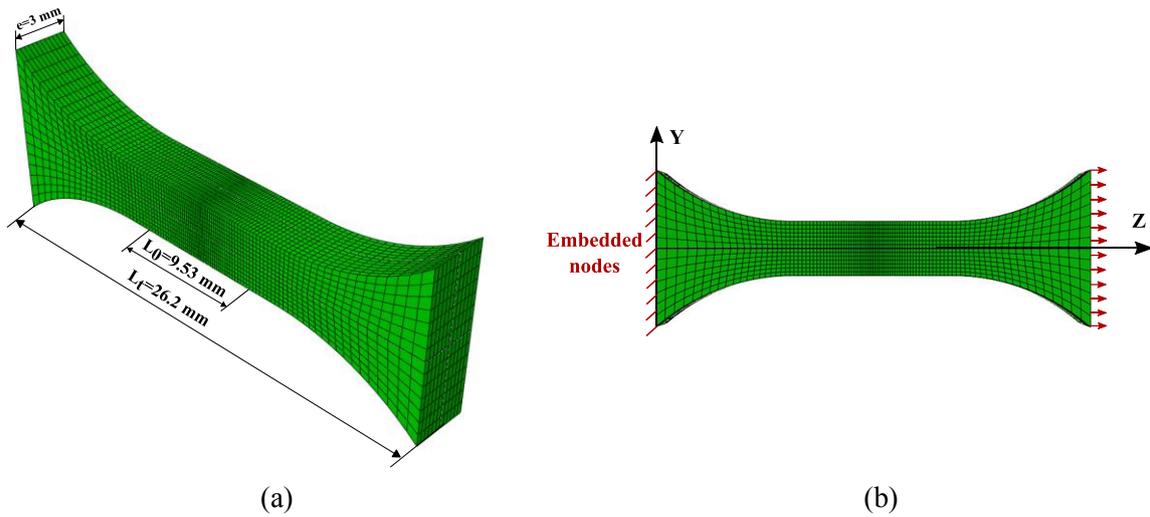


Fig. 8. Numerical model: (a) geometrical information; (b) boundary conditions applied.

4.2. Results and analysis

A series of numerical simulations were performed to study and compare the influence of viscoelastic behaviours and thermo-mechanical coupling during the deformation process of two types of polyethylene. In these simulations, specimens of both materials were subjected to stretching conditions varying the initial strain rate. Experimentally, the mechanical behaviour of both materials presents a strong dependence on strain rate and temperature, increasing the yield stress with strain rate and decreasing with temperature. However, while the UHMWPE exhibits significant strain hardening after yield, Fig. 6, the HDPE presents flat flow behaviour Fig. 7. In the former, the viscoelastic resistance plays an important role in the deformation process, whereas in the HDPE this contribution is not relevant. In this regard, the mechanical response of these materials can be understood as the competition between hardening due to strain and strain rate effects and softening due to thermal effects. This competition becomes more complex at high strain rates where this variable induces an important hardening but thermal softening is more relevant because of the strong thermo-mechanical coupling under these conditions, as can be deduced from Eq. (22).

Figs. 9 and 10 show two stages of the tensile deformation process for UHMWPE and HDPE at strain rate conditions of 0.1 s^{-1} and 100 s^{-1} . These stages are identified by the engineering strain defined as $\epsilon = \Delta Z/L_0$, where ΔZ is the current displacement applied in the longitudinal direction. A uniform deformation along the whole gauge length is observed for UHMWPE whereas HDPE presents a local deformation which results in a pronounced necking process. This difference in

the behaviour of both materials can be explained by the higher viscosity and strain-hardening after yield of UHMWPE with respect to HDPE. In this sense, viscous contributions to the mechanical behaviour have been demonstrated to stabilize necking formation leading to higher ductility [19]. Moreover, strain hardening also plays a stabilizing role in necking formation [58-60]. In addition, thermal softening also influences the deformation process of thermoplastic polymers [61-65], as observed in other ductile materials [66,67]. In this regard, understanding this as a negative hardening, if temperature evolution occurs heterogeneously by inelastic dissipation, local necking is favoured leading to instability in the zone that reaches higher temperatures. Fig. 11 shows the temperature distribution along the gauge length for both polyethylene materials at strains of 0.2 and 0.5 and strain rates of 0.1 s^{-1} and 100 s^{-1} . Temperature remains practically uniform along the gauge length of UHMWPE specimen whereas it shows a relevant concentration in the necking zone of HDPE specimen. This temperature increment due to inelastic dissipation is more relevant at high strain rates and leads to a more pronounced necking formation, see Fig. 10(d) where the temperature reaches a maximum up to $\Delta\theta = 60 \text{ K}$ in the necking zone of HDPE. Comparing the results in terms of temperature, a higher increment of this variable is reached by HDPE. Although this material does not present contribution of the viscous branch, the higher values of stress flow result in higher contribution of inelastic dissipation according to Eq. (22). This higher temperature increment in HDPE favours necking formation in comparison with UHMWPE.

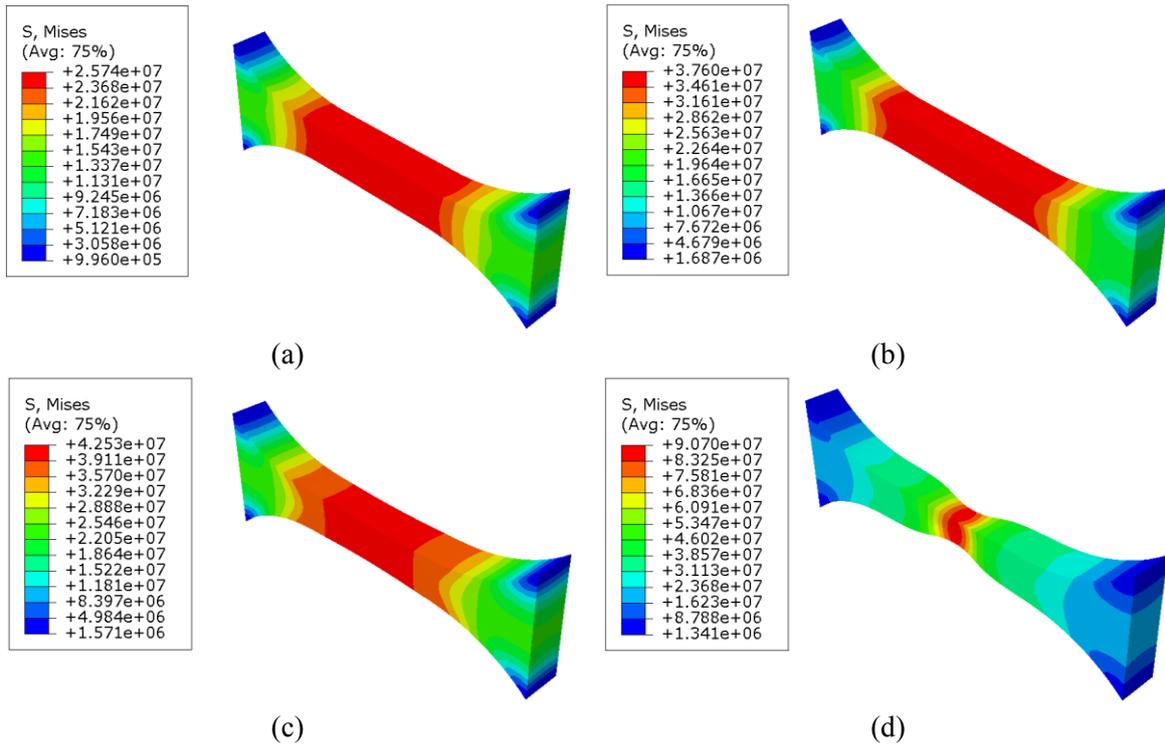


Fig. 9. Stress distribution at $\dot{\epsilon}_0 = 0.1 \text{ s}^{-1}$ for: (a) UHMWPE at $\epsilon = 0.2$ (b) UHMWPE at $\epsilon = 0.5$ (c) HDPE at $\epsilon = 0.2$ (d) HDPE at $\epsilon = 0.5$.

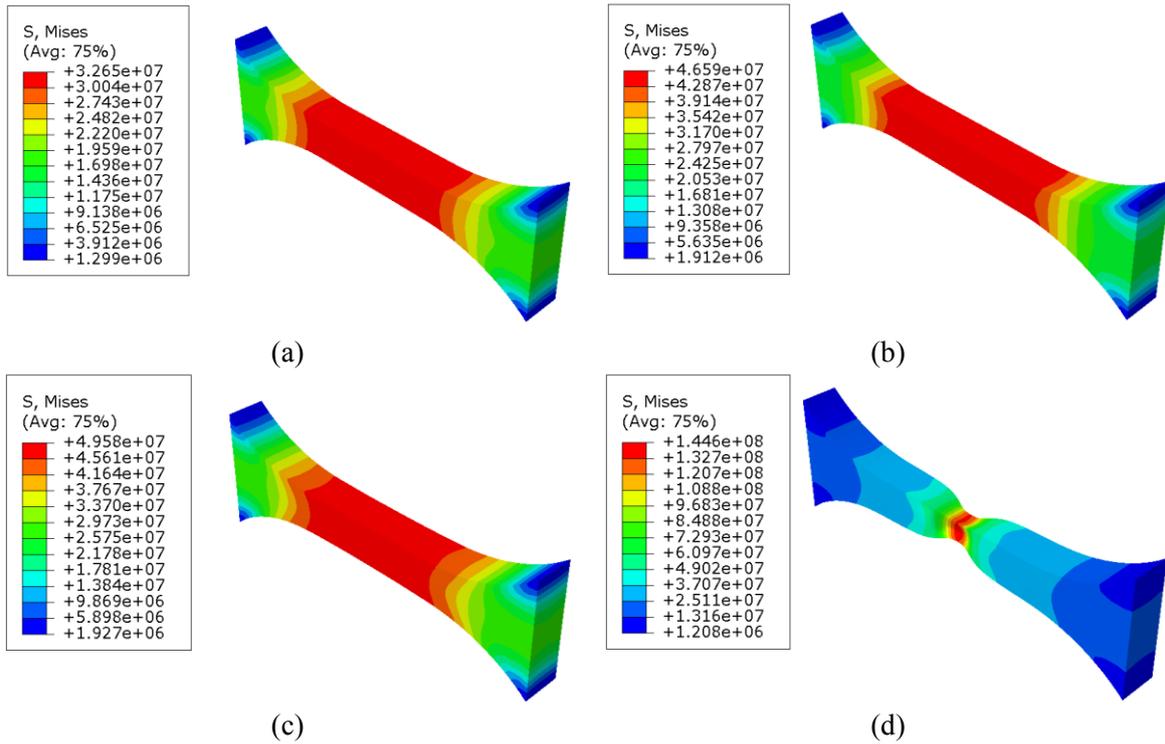


Fig. 10. Stress distribution at $\dot{\epsilon}_0 = 100 \text{ s}^{-1}$ for: (a) UHMWPE at $\epsilon = 0.2$ (b) UHMWPE at $\epsilon = 0.5$ (c) HDPE at $\epsilon = 0.2$ (d) HDPE at $\epsilon = 0.5$.

The results numerically obtained here in terms of necking formation correlate with experiments conducted by others authors [58,59]. Mohagheghian et al. [58] studied the response of different types of polyethylene at nominal strain rates of 0.01 s^{-1} , 0.1 s^{-1} and 1 s^{-1} over specimens with the same geometry used in our numerical simulations. At low strain rates, HDPE presents neck formation while at high strain rates HDPE fails during neck propagation phases. In contrast, no necking process was showed in the UHMWPE specimen for all strain rates studied. This response was explained by the stabilising effect associated to strain hardening. Moreover, the same way to deform was observed by Torres et al. [68] for HDPE. These authors also observed a necking process with an increase in temperature along necking length in agreement with the results obtained here.

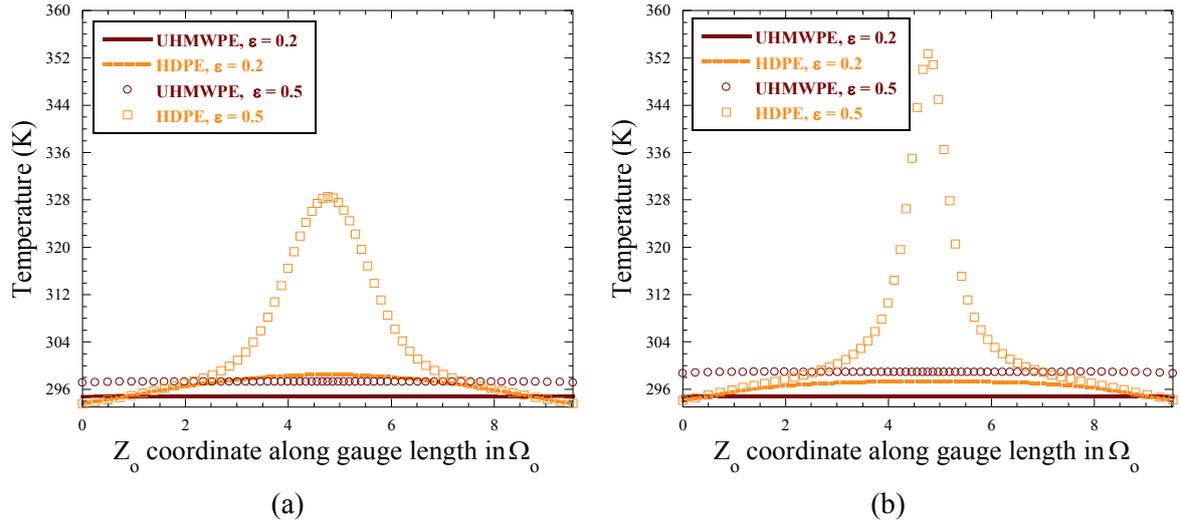


Fig. 11. Temperature distribution along gauge length at strains of 0.2 and 0.5 in UHMWPE and HDPE specimens at: (a) strain rate of 0.1 s^{-1} ; and (b) strain rate of 100 s^{-1} .

5. Conclusions

In this work, a hyperelastic constitutive model which combines viscoelasticity and viscoplasticity with thermo-mechanical coupling is developed to predict the mechanical behaviour of semi-crystalline polymeric matrices. This model is physically motivated on the interpretation of the stress-strain response of semi-crystalline polymers as the combination of overcoming an intermolecular resistance, a viscous resistance and a network resistance. To this end, the intermolecular resistance introduces the thermo-viscoplastic behaviour of the polymer and the viscous and network resistances introduce the thermo-viscoelastic behaviour. In addition, temperature evolution due to inelastic dissipation is accounted for, as well as thermal softening associated to temperature increment.

The constitutive equations were implemented in a VUMAT subroutine for Abaqus/Explicit and then, the model parameters were identified for UHMWPE and HDPE from experimental data covering a wide range of strain rate and temperature conditions. Finally, the model was employed to study the influence of viscous contributions and thermo-mechanical coupling on necking process when semi-crystalline specimens are subjected to large stretching conditions. Numerical predictions show that UHMWPE can undergo higher strain values without neck formation leading to higher ductility with respect to HDPE.

The consideration of viscous behaviours and their thermo-mechanical coupling in the constitutive modelling of semi-crystalline polymers were identified as key points in the faithful prediction of the characteristic mechanical behaviour of both injection moulded and 3D printed thermoplastic polymers. In this sense, the constitutive model developed in this work is able not only to be considered as a predictive tool, but also to be useful in the understanding of the material behaviour of semi-crystalline polymers. The proposed model provides insights into the mechanical behaviour of polymeric matrices, allowing for the development of new composite materials with dynamic applications.

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Appendix A. Nomenclature used in the formulation

| Nomenclature | | | |
|--|---|-----------------------------|--|
| $\{\Omega_o, \bar{\Omega}, \bar{\bar{\Omega}}, \tilde{\Omega}, \Omega\}$ | initial, dilated, plastic dilated relaxed, viscous dilated relaxed and current configurations | $\bar{\bar{W}}_I^P$ | skew part of the plastic velocity gradient in Ω |
| \bar{e} | specific internal energy per unit volume in $\bar{\Omega}$ | \bar{W}_V^V | skew part of the viscous velocity gradient in $\bar{\Omega}$ |
| \bar{Q} | heat flux per unit volume in $\bar{\Omega}$ | σ | Cauchy stress tensor |
| \bar{R} | heat resource per unit volume in $\bar{\Omega}$ | σ_N | network Cauchy stress tensor |
| $\bar{\eta}$ | specific entropy per unit volume in $\bar{\Omega}$ | σ_I | intermolecular Cauchy stress tensor |
| \bar{C} | heat capacity per unit volume in $\bar{\Omega}$ | σ_V | viscous Cauchy stress tensor |
| $\{\theta, \theta_{ref}, \theta_{melt}, \theta_{min}\}$ | current, reference, melting and lowest temperature considered | \bar{S}_N | network second Piola-Kirchhoff stress tensor in $\bar{\Omega}$ |
| f_θ | temperature-dependent function | $\{\bar{S}_I, \bar{S}_I\}$ | intermolecular second Piola-Kirchhoff stress tensor in $\bar{\Omega}$ and $\bar{\bar{\Omega}}$ |
| $\bar{\Psi}$ | Helmholtz free energy per unit volume in $\bar{\Omega}$ | \bar{S}_V | viscous second Piola-Kirchhoff stress tensor in $\bar{\Omega}$ |
| $\bar{\nabla}_x$ | gradient with respect to the point x in $\bar{\Omega}$ | \bar{M}_N | network Mandel stress tensor in $\bar{\Omega}$ |
| $\bar{\mathfrak{L}}$ | Langevin function | $\{\bar{M}_I, \bar{M}_I\}$ | intermolecular Mandel stress tensor in $\bar{\Omega}$ and $\bar{\bar{\Omega}}$ |
| I | Identity matrix | \bar{M}_V | viscous Mandel stress tensor in $\bar{\Omega}$ |
| J_I | determinant of the intermolecular deformation gradient | $\bar{\lambda}$ | average total stretch ratio |
| J_N | determinant of the network deformation gradient | $\bar{\lambda}_L$ | locking stretch |
| J_V | determinant of the viscous deformation gradient | $\{I_{1I}, J_{2I}\}$ | stress invariants of the intermolecular Mandel stress tensor |
| F | deformation gradient | $\bar{\sigma}_{eqI}$ | Rhagava equivalent stress |
| F^θ | thermal deformation gradient | g_I | plastic potential |
| F^M | mechanical deformation gradient | \bar{r}_I | gradient of the plastic potential |
| F_N^e | network elastic deformation gradient | $\dot{\bar{Y}}_I^P$ | viscoplastic multiplier |
| F_I^e | intermolecular elastic deformation gradient | α_θ | thermal expansion coefficient |
| F_I^P | intermolecular plastic deformation gradient | $\{\lambda_0, \mu_0\}$ | classical Lamé constants |
| F_V^e | viscoelastic elastic deformation gradient | E | Young's modulus |
| F_V^V | viscoelastic viscous deformation gradient | $\{E_{ref}, E_1\}$ | Young's modulus at the reference temperature and a specified material parameter |
| $\{C_N^e, B_N^e\}$ | network elastic right and left Cauchy-Green tensor | $\{\sigma_T, \sigma_{T0}\}$ | yield stress in uniaxial tension and its value at reference temperature |
| C_I^e | intermolecular elastic right Cauchy-Green tensor | α | pressure sensitivity parameter |
| $\{C_V^e, B_V^e\}$ | viscous elastic right Cauchy-Green tensor | β | volumetric plastic strain parameter |
| l | velocity gradient | $\dot{\epsilon}_0$ | reference strain rate |
| l_V^e | viscous elastic velocity spatial gradient | C | rate sensitivity parameter |

| | | | |
|---------------------------|---|--------------------------|---|
| $\bar{\mathbf{L}}^\theta$ | thermal velocity gradient in $\bar{\Omega}$ | m | temperature sensitivity parameter |
| $\bar{\mathbf{L}}_V^v$ | viscous velocity gradient in $\bar{\Omega}$ | C_R | initial elastic modulus of the network backstress resistance |
| $\bar{\mathbf{L}}_I^p$ | plastic velocity gradient in $\bar{\Omega}$ | C_θ | temperature-dependent material parameter controlling the elastic modulus of network |
| \mathbf{d}_N^e | symmetric part of the network elastic velocity gradient in Ω | C_v | initial elastic modulus of the viscous backstress resistance |
| \mathbf{d}_I^e | symmetric part of the intermolecular elastic velocity gradient in Ω | $\bar{\lambda}_V$ | average viscous stretch ratio |
| \mathbf{d}_V^e | symmetric part of the viscous elastic velocity gradient in Ω | $\bar{\lambda}_{VL}$ | viscous locking stretch |
| $\bar{\mathbf{D}}^p$ | symmetric part of the plastic velocity gradient in $\bar{\Omega}$ | $\bar{\mathbf{N}}_V$ | gradient of the viscoelastic flow |
| $\bar{\mathbf{D}}^v$ | symmetric part of the viscous velocity gradient in $\bar{\Omega}$ | $\dot{\bar{\gamma}}_V^v$ | viscoelastic multiplier |
| $\bar{\mathbf{D}}_I^p$ | symmetric part of the intermolecular plastic velocity gradient in $\bar{\Omega}$ | τ_v | effective stress driving the viscous flow |
| $\bar{\mathbf{D}}_V^v$ | symmetric part of the intermolecular viscous velocity gradient in $\bar{\Omega}$ | σ_{VT} | Material parameter of the viscous resistance |
| $\bar{\mathbf{W}}^\theta$ | skew part of the thermal velocity gradient in $\bar{\Omega}$ | | |

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Figures

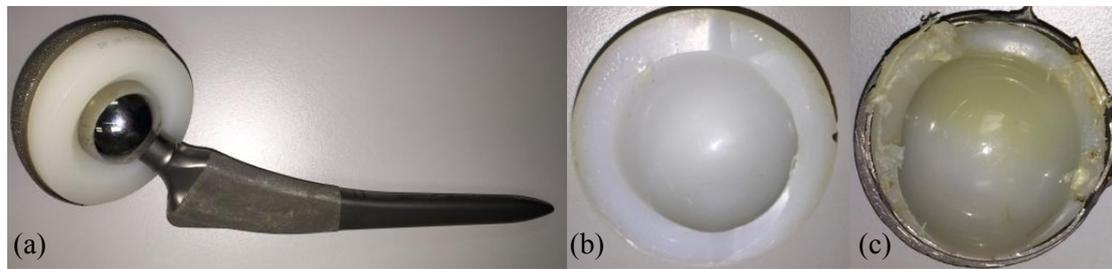


Fig. 1. (a) Real polyethylene-metal hip prosthetic. (b)-(c) Polyethylene acetabular cup removed from patients due to excessive material erosion. Materials provided by Dra. I. Gómez Arrayás (Hospital Ruber Internacional of Madrid).

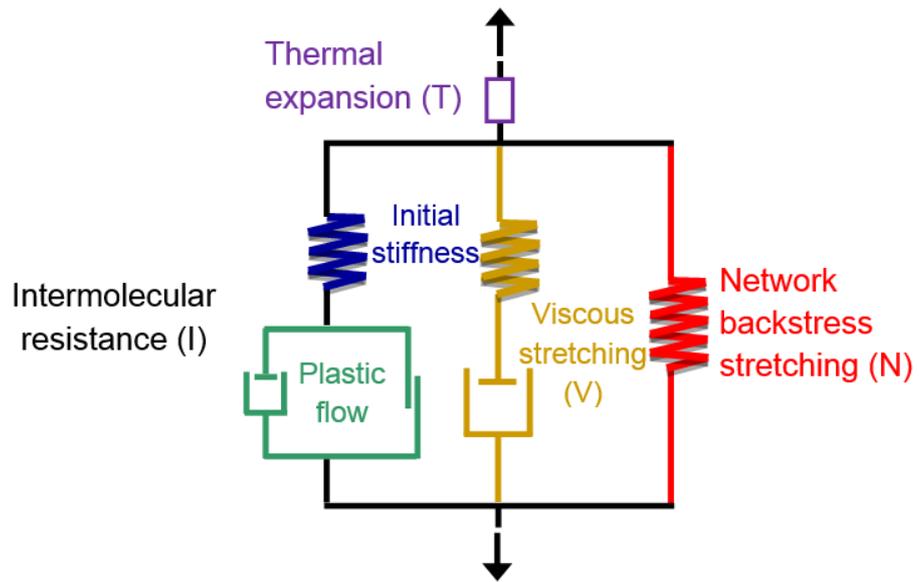


Fig. 2. Rheological scheme of the proposed constitutive model.

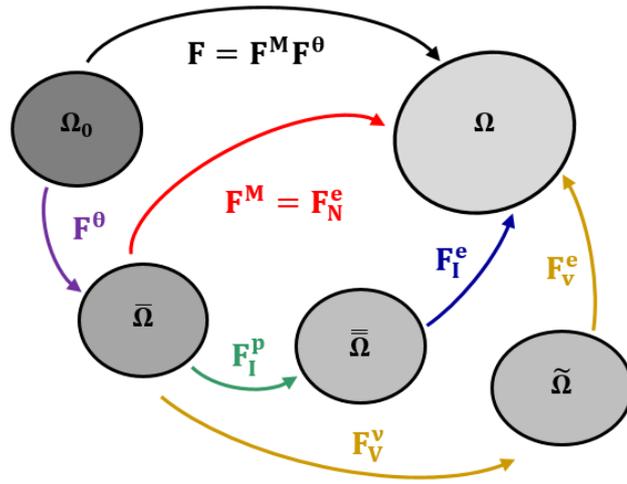
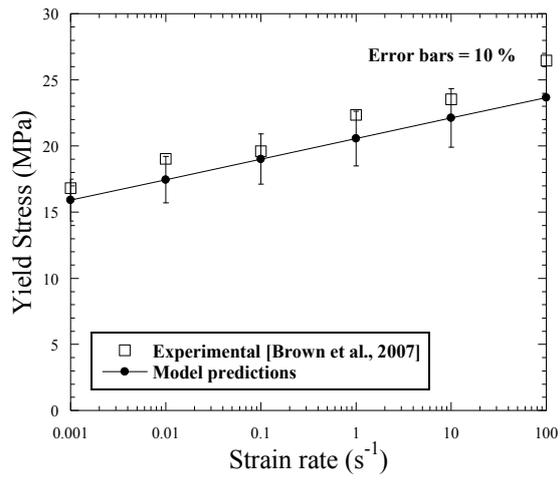
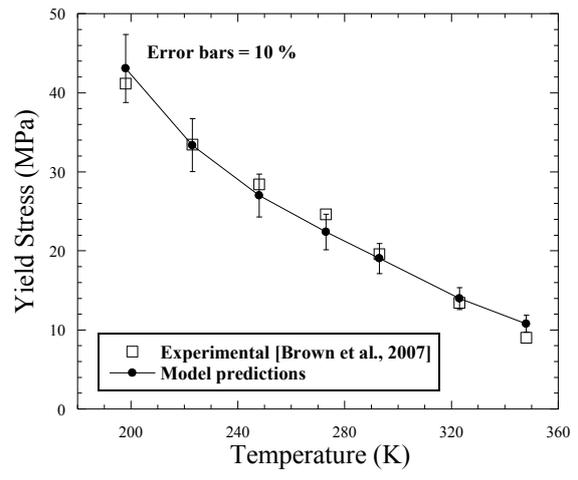


Fig. 3. Kinematics of the model showing the reference or initial configuration Ω_0 , the dilated configuration $\bar{\Omega}$, the plastic dilated relaxed configuration $\bar{\bar{\Omega}}$, the viscous dilated relaxed configuration $\tilde{\Omega}$, and the current or loaded configuration Ω .

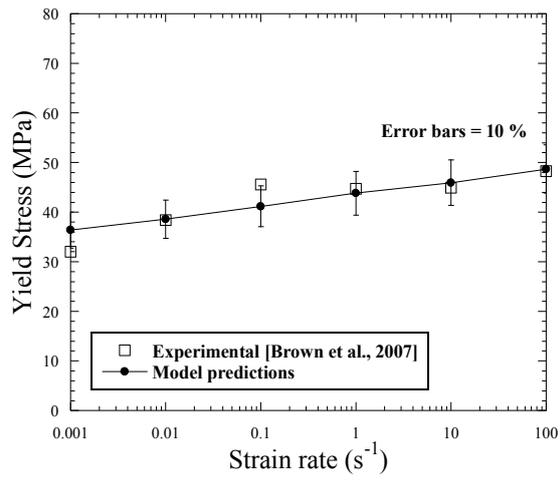


(a)

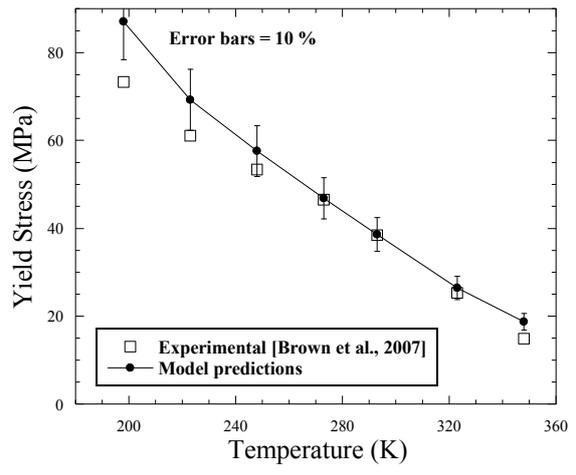


(b)

Fig. 4. Experimental data [18] versus model predictions in terms of: (a) strain rate sensitivity; and (b) temperature sensitivity of UHMWPE for uniaxial compression tests.

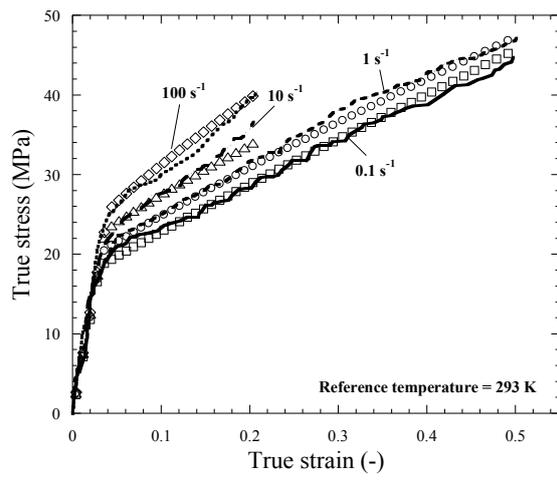


(a)

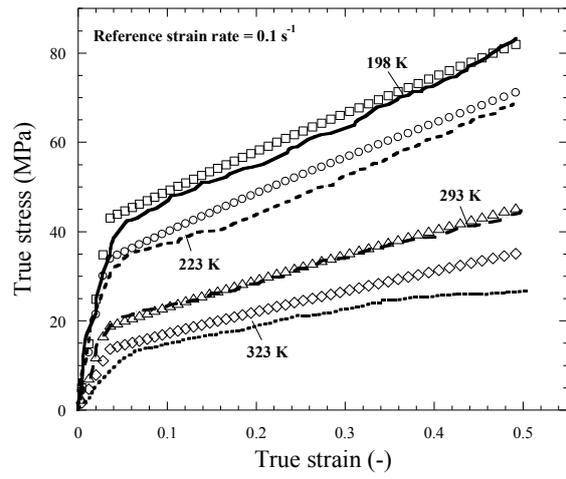


(b)

Fig. 5. Experimental data [18] versus model predictions in terms of: (a) strain rate sensitivity; and (b) temperature sensitivity of HDPE for uniaxial compression tests.

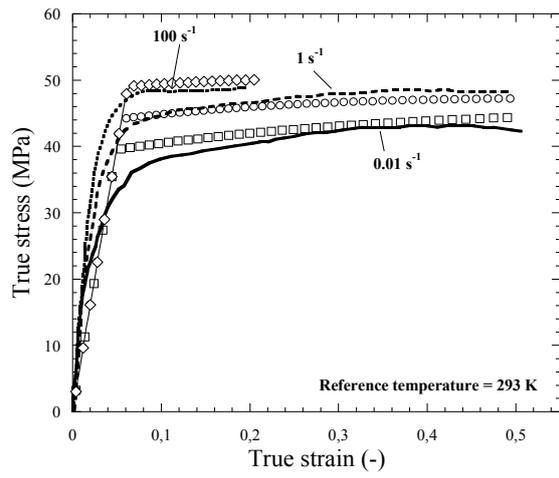


(a)

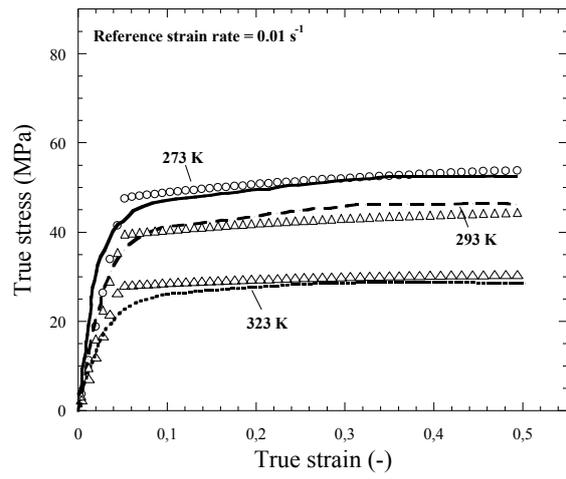


(b)

Fig. 6. Stress-strain experimental curves of UHMWPE [18] versus model predictions for: (a) different strain rates; and (b) different temperatures.

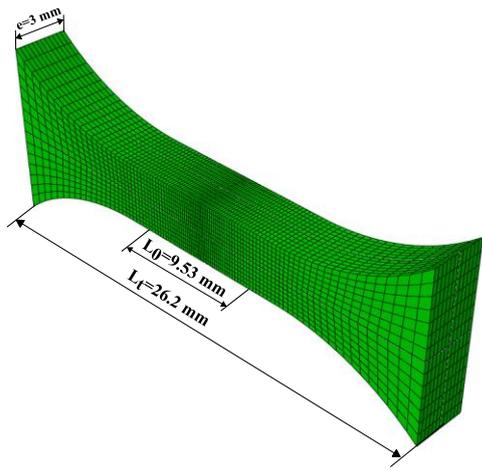


(a)

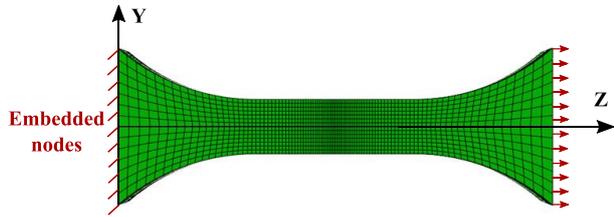


(b)

Fig. 7. Stress-strain experimental curves of HDPE [18] versus model predictions for: (a) different strain rates; and (b) different temperatures.



(a)



(b)

Fig. 8. Numerical model: (a) geometrical information; (b) boundary conditions applied.

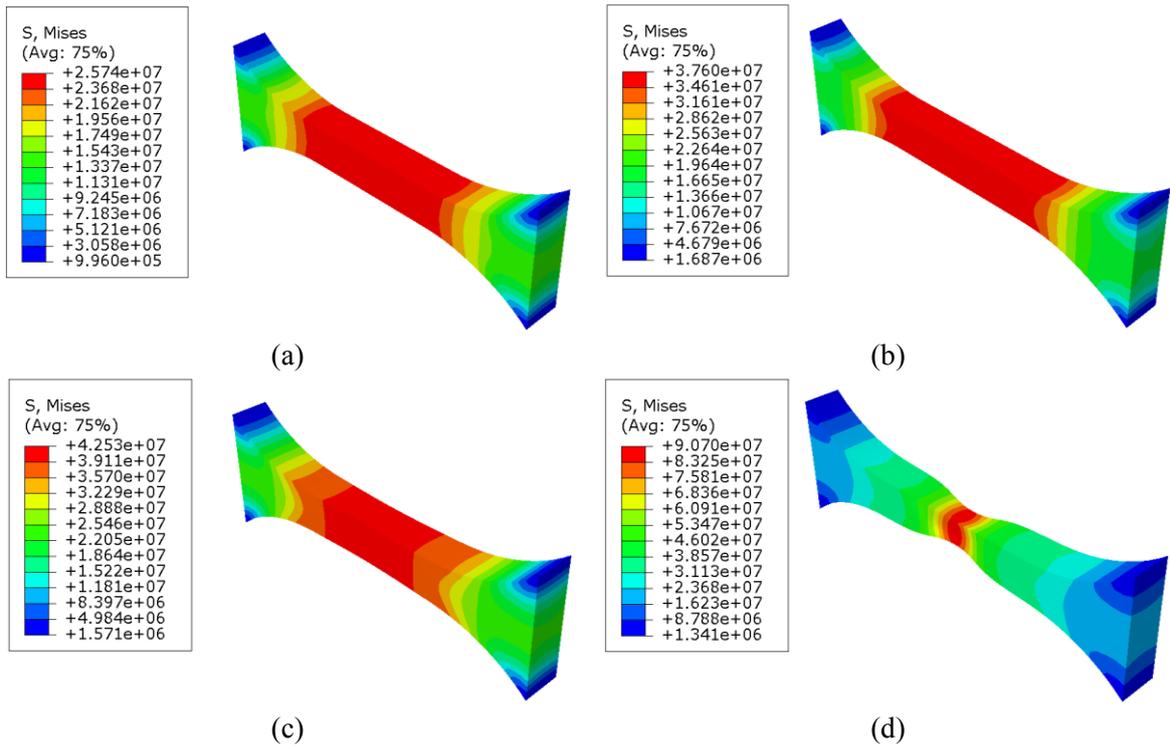


Fig. 9. Stress distribution at $\dot{\varepsilon}_0 = 0.1 \text{ s}^{-1}$ for: (a) UHMWPE at $\varepsilon = 0.2$ (b) UHMWPE at $\varepsilon = 0.5$ (c) HDPE at $\varepsilon = 0.2$ (d) HDPE at $\varepsilon = 0.5$.

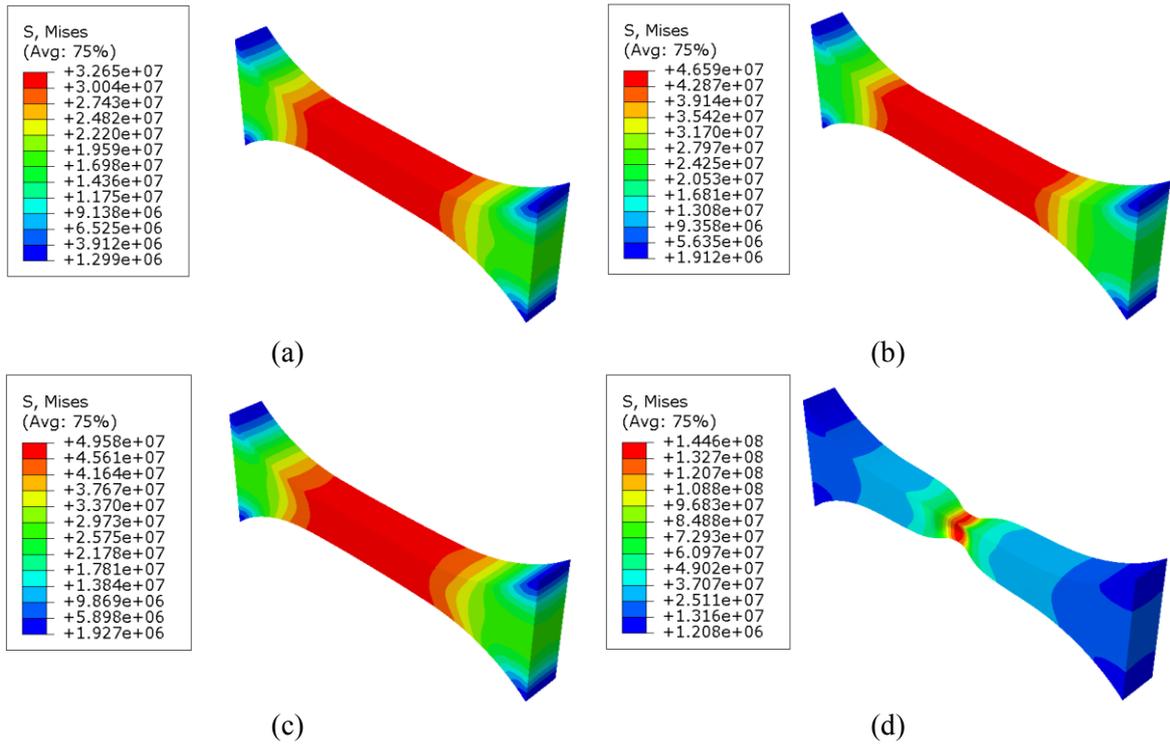


Fig. 10. Stress distribution at $\dot{\epsilon}_0 = 100 \text{ s}^{-1}$ for: (a) UHMWPE at $\epsilon = 0.2$ (b) UHMWPE at $\epsilon = 0.5$ (c) HDPE at $\epsilon = 0.2$ (d) HDPE at $\epsilon = 0.5$.

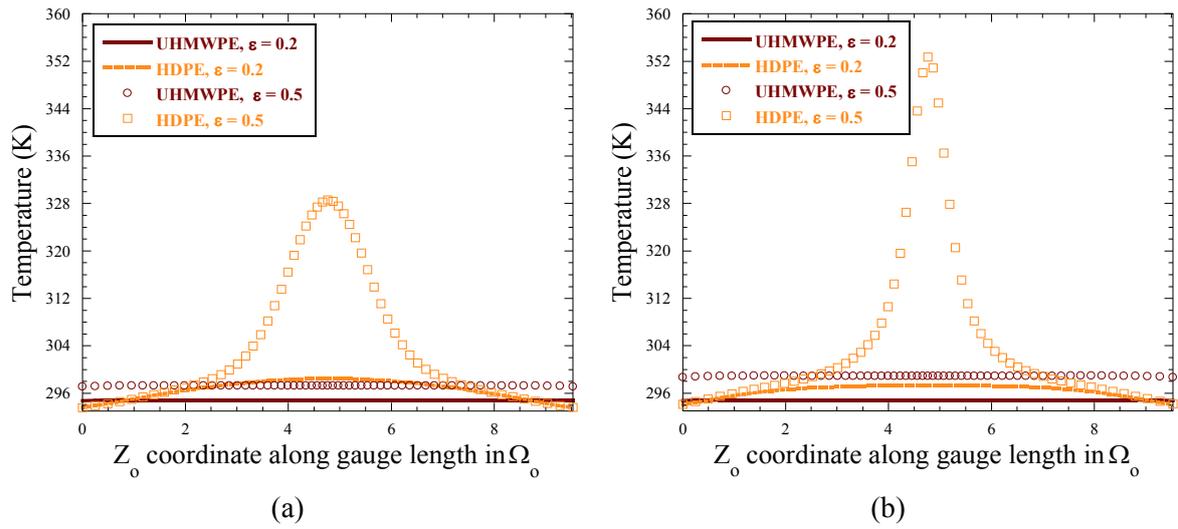


Fig. 11. Temperature distribution along gauge length at strains of 0.2 and 0.5 in UHMWPE and HDPE specimens at: (a) strain rate of 0.1 s^{-1} ; and (b) strain rate of 100 s^{-1} .

Tables

Table 1

Material parameters for UHMWPE.

| Initial elastic properties | | | | | Intermolecular resistance | | | | | | |
|-----------------------------|----------------------|------------------------------------|---------------------------------------|--------------------|---------------------------|-----|------------------------------|-------------------------------|------------------------------|----------|---------|
| E_{ref} (GPa) | E_1 (MPa/K) | ν | $\dot{\epsilon}_0$ (s ⁻¹) | C | σ_{T0} (MPa) | m | θ_{ref} (K) | θ_{melt} (K) | θ_{min} (K) | α | β |
| 0.541 | -6.5 | 0.46 | 0.1 | 0.048 | 37.5 | 0.7 | 293 | 406 | 198 | 1.0 | 1.0 |
| Viscous resistance | | | | Network resistance | | | | | | | |
| C_V (Pa) | $\bar{\lambda}_{VL}$ | σ_{VT} (MPa) | C_R (MPa) | C_θ (MPa/K) | $\bar{\lambda}_L$ | | | | | | |
| 12.5 | 5 | 0.05 | 23 | -0.15 | 5 | | | | | | |
| General properties | | | | | | | | | | | |
| ρ (kg/m ³) | C_p (J/Kg K) | α_θ (K ⁻¹) | | | | | | | | | |
| 940 | 1900 | $124.5 \cdot 10^{-6}$ | | | | | | | | | |

Table 2
Material parameters for HDPE.

| Initial elastic properties | | | | | Intermolecular resistance | | | | | | |
|-----------------------------|----------------------|-------------------------------|---------------------------------------|--------------------|---------------------------|------|------------------------------|-------------------------------|------------------------------|----------|---------|
| E_{ref} (GPa) | E_1 (MPa/K) | ν | $\dot{\epsilon}_0$ (s ⁻¹) | C | σ_{T0} (MPa) | m | θ_{ref} (K) | θ_{melt} (K) | θ_{min} (K) | α | β |
| 0.782 | -6.5 | 0.46 | 0.01 | 0.028 | 80 | 0.76 | 293 | 406 | 198 | 1.0 | 1.0 |
| Viscous resistance | | | | Network resistance | | | | | | | |
| C_V (Pa) | $\bar{\lambda}_{VL}$ | σ_{VT} (Pa) | C_R (MPa) | C_0 (MPa/K) | $\bar{\lambda}_L$ | | | | | | |
| (-) | (-) | (-) | 12 | -0.15 | 5 | | | | | | |
| General properties | | | | | | | | | | | |
| ρ (kg/m ³) | C_p (J/Kg K) | α_0 (K ⁻¹) | | | | | | | | | |
| 970 | 1900 | $93.6 \cdot 10^{-6}$ | | | | | | | | | |

Appendix A. Nomenclature used in the formulation

| Nomenclature | | | |
|--|---|----------------------------------|--|
| $\{\Omega_0, \bar{\Omega}, \bar{\bar{\Omega}}, \tilde{\Omega}, \Omega\}$ | initial, dilated, plastic dilated relaxed, viscous dilated relaxed and current configurations | $\bar{\bar{W}}_I^P$ | skew part of the plastic velocity gradient in Ω |
| \bar{e} | specific internal energy per unit volume in $\bar{\Omega}$ | $\bar{\bar{W}}_V^V$ | skew part of the viscous velocity gradient in $\bar{\Omega}$ |
| \bar{Q} | heat flux per unit volume in $\bar{\Omega}$ | σ | Cauchy stress tensor |
| \bar{R} | heat resource per unit volume in $\bar{\Omega}$ | σ_N | network Cauchy stress tensor |
| $\bar{\eta}$ | specific entropy per unit volume in $\bar{\Omega}$ | σ_I | intermolecular Cauchy stress tensor |
| \bar{C} | heat capacity per unit volume in $\bar{\Omega}$ | σ_V | viscous Cauchy stress tensor |
| $\{\theta, \theta_{ref}, \theta_{melt}, \theta_{min}\}$ | current, reference, melting and lowest temperature considered | \bar{S}_N | network second Piola-Kirchhoff stress tensor in $\bar{\Omega}$ |
| f_θ | temperature-dependent function | $\{\bar{S}_I, \bar{\bar{S}}_I\}$ | intermolecular second Piola-Kirchhoff stress tensor in $\bar{\Omega}$ and $\bar{\bar{\Omega}}$ |
| $\bar{\Psi}$ | Helmholtz free energy per unit volume in $\bar{\Omega}$ | \bar{S}_V | viscous second Piola-Kirchhoff stress tensor in $\bar{\Omega}$ |
| $\bar{\nabla}_x$ | gradient with respect to the point x in $\bar{\Omega}$ | \bar{M}_N | network Mandel stress tensor in $\bar{\Omega}$ |
| $\bar{\mathcal{L}}$ | Langevin function | $\{\bar{M}_I, \bar{\bar{M}}_I\}$ | intermolecular Mandel stress tensor in $\bar{\Omega}$ and $\bar{\bar{\Omega}}$ |
| \mathbf{I} | Identity matrix | \bar{M}_V | viscous Mandel stress tensor in $\bar{\Omega}$ |
| J_I | determinant of the intermolecular deformation gradient | $\bar{\lambda}$ | average total stretch ratio |
| J_N | determinant of the network deformation gradient | $\bar{\lambda}_L$ | locking stretch |
| J_V | determinant of the viscous deformation gradient | $\{I_{11}, J_{21}\}$ | stress invariants of the intermolecular Mandel stress tensor |
| \mathbf{F} | deformation gradient | $\bar{\sigma}_{eqI}$ | Rhagava equivalent stress |
| \mathbf{F}^θ | thermal deformation gradient | g_I | plastic potential |
| \mathbf{F}^M | mechanical deformation gradient | \bar{r}_I | gradient of the plastic potential |
| \mathbf{F}_N^e | network elastic deformation gradient | $\bar{\bar{\gamma}}_I^P$ | viscoplastic multiplier |
| \mathbf{F}_I^e | intermolecular elastic deformation gradient | α_θ | thermal expansion coefficient |
| \mathbf{F}_I^P | intermolecular plastic deformation gradient | $\{\lambda_0, \mu_0\}$ | classical Lamé constants |
| \mathbf{F}_V^e | viscoelastic elastic deformation gradient | E | Young's modulus |
| \mathbf{F}_V^V | viscoelastic viscous deformation gradient | $\{E_{ref}, E_1\}$ | Young's modulus at the reference temperature and a specified material parameter |
| $\{\mathbf{C}_N^e, \mathbf{B}_N^e\}$ | network elastic right and left Cauchy-Green tensor | $\{\sigma_T, \sigma_{T0}\}$ | yield stress in uniaxial tension and its value at reference temperature |
| \mathbf{C}_I^e | intermolecular elastic right Cauchy-Green tensor | α | pressure sensitivity parameter |
| $\{\mathbf{C}_V^e, \mathbf{B}_V^e\}$ | viscous elastic right Cauchy-Green tensor | β | volumetric plastic strain parameter |
| \mathbf{l} | velocity gradient | $\dot{\epsilon}_0$ | reference strain rate |
| \mathbf{l}_V^e | viscous elastic velocity spatial gradient | C | rate sensitivity parameter |
| $\bar{\mathbf{L}}^\theta$ | thermal velocity gradient in $\bar{\Omega}$ | m | temperature sensitivity parameter |
| $\bar{\mathbf{L}}_V^V$ | viscous velocity gradient in $\bar{\Omega}$ | C_R | initial elastic modulus of the network backstress resistance |
| $\bar{\mathbf{L}}_I^P$ | plastic velocity gradient in $\bar{\bar{\Omega}}$ | C_θ | temperature-dependent material parameter controlling the elastic modulus of network |
| \mathbf{d}_N^e | symmetric part of the network elastic velocity gradient in Ω | C_v | initial elastic modulus of the viscous backstress resistance |
| \mathbf{d}_I^e | symmetric part of the intermolecular elastic velocity gradient in Ω | $\bar{\lambda}_V$ | average viscous stretch ratio |

| | | | |
|---------------------------|--|------------------------|--|
| \mathbf{d}_v^e | symmetric part of the viscous elastic velocity gradient in Ω | $\bar{\lambda}_{vL}$ | viscous locking stretch |
| $\bar{\mathbf{D}}^p$ | symmetric part of the plastic velocity gradient in $\bar{\Omega}$ | $\tilde{\mathbf{N}}_v$ | gradient of the viscoelastic flow |
| $\bar{\mathbf{D}}^v$ | symmetric part of the viscous velocity gradient in $\bar{\Omega}$ | $\hat{\gamma}_v^v$ | viscoelastic multiplier |
| $\bar{\mathbf{D}}_I^p$ | symmetric part of the intermolecular plastic velocity gradient in $\bar{\bar{\Omega}}$ | τ_v | effective stress driving the viscous flow |
| $\tilde{\mathbf{D}}_v^v$ | symmetric part of the intermolecular viscous velocity gradient in $\tilde{\Omega}$ | σ_{vT} | Material parameter of the viscous resistance |
| $\bar{\mathbf{W}}^\theta$ | skew part of the thermal velocity gradient in $\bar{\Omega}$ | | |
