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J.P. Goulmy, L. Toualbi, V. Boyer, P. Kanoute, D. Retraint, et al.. A model for the influence of work hardening and microstructure on the evolution of residual stresses under thermal loading – Application to Inconel 718. European Journal of Mechanics - A/Solids, 2024, 107, pp.105361. 10.1016/j.euromechsol.2024.105361. hal-04634663

HAL Id: hal-04634663 https://hal.science/hal-04634663

Submitted on 4 Jul 2024

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1	A model for the influence of work hardening and microstructure on the
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3	Inconel 718
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11	(+33) 4 42 93 82 23
12	
13	Abstract
14	This study proposes a model for the influence of work hardening and microstructure on the thermal
15	relaxation of residual stresses. To construct such a model, an experimental campaign is first conducted
16	on shot peened samples of Inconel 718 to generate different levels of residual stress and work
17	hardening. The effect of the grain size and the size of the strengthening precipitates is investigated by
18	producing two modified microstructures. Two shot peening conditions are used to introduce several
19	profiles of residual stress and work hardening. These profiles are evaluated using X-ray diffraction. A
20	thermal loading is then applied at 550°C with varying holding times, leading to a rapid but not complete
21	relaxation of the residual stresses and work hardening. The experimental results exhibit the fact that
22	the work hardening levels have a significant influence on this relaxation while the grain size and the
23	size of the strengthening precipitates have a very moderate influence. Based on these experimental
24	results, a model is proposed that considers the influence of work hardening on the thermal relaxation
25	of residual stresses with some predictive applications. It is therefore possible to estimate the relaxation
26	of residual stresses at any point on a shot peened part.
27	
28	Keyword: Inconel 718, Shot Peening, Residual stresses, Work hardening, Thermal relaxation

31 Introduction

32 In order to improve the fatigue life of critical components, surface improvement methods such as shot 33 peening are widely used in the aerospace industry. This process generates superficial compressive 34 residual stresses that tend to delay fatigue crack initiation and prevent small crack propagation [1], [2], 35 [3], [4], [5]. For structures that experience high temperature variations in service, these surface 36 enhancement methods are only efficient if the thermo-mechanical relaxation of the generated residual 37 stress field is moderate at operating temperatures. It is however observed that relaxation due to 38 thermal effects is far from being negligible in high-temperature fatigue tests [6]. It is therefore 39 important to understand the mechanisms that induce the thermal relaxation of residual stresses in 40 order to further input these effects into a model. One of the critical points is to estimate the kinematic 41 and time constants of this relaxation.

42 The nickel-based superalloy Inconel 718 is widely used in the aeronautical industry. In its standard 43 metallurgical state known as Direct Aged (DA), this alloy features a microstructure with a y matrix composed of grains of about 5 μ m, γ'/γ'' strengthening precipitates of a few nanometers, a δ phase 44 45 anchored at the grain boundaries, as well as carbides and nitrides that locally harden the matrix [7], 46 [8], [9], [10]. Subjected to temperatures close to 550°C, Inconel 718 turbine discs undergo thermo-47 mechanical stresses in service. These discs can present microstructural heterogeneities because of their manufacturing process history; they can also undergo more than ten conditions of shot peening 48 49 treatment, depending on their geometry. The in-service evolution of residual stresses in these parts 50 can therefore be particularly diverse and complex. In order to evaluate the benefits of shot peening on 51 the fatigue life of these parts, it is necessary to understand the microstructural changes induced by 52 shot peening and their influence on the thermal relaxation of residual stresses. Inconel 718 thus 53 constitutes a material of interest to study and model residual stress relaxation due to thermal loading. 54 Clearly, microstructural parameters of Inconel 718 have a significant influence on the tensile, fatigue, 55 and creep properties [7], [8], [9]. Furthermore, previous studies [11], [12] have highlighted the 56 influence of microstructural modifications on the residual stress and work hardening profiles induced 57 by shot peening. It was shown that work hardening is significantly affected by a change in grain size. 58 On the contrary, changing the grain size or the size of the strengthening precipitates γ'/γ'' appears to 59 have minimal effect on the residual stress profile. It is then useful to define how these different profiles 60 relax with time.

Many authors [13], [14], [15], [16], [17] have described the thermal relaxation of residual stresses caused by shot peening. By relaxation, it is here meant a lowering of the local absolute value of the residual stress level. The main observations are: (i) the higher the temperature, the higher the relaxation of the residual stresses combined with the fact that relaxation is negligible below a threshold temperature, except at the surface where a change may occur depending on the shot peening condition used; (ii) relaxation is observed at each depth of the profile, hence, the general shape of the profiles (position of the maximum stress, position of the tensile transition) is not changed, except above a certain temperature where a shift from the top of the profile to the core of the part is observed (iii) for a given temperature, the main evolution of the residual stresses takes place during the first instants of temperature holding, then, the residual stress profiles tend to asymptotic profiles with time.

71 The impact of work hardening on the relaxation of residual stresses has also previously been 72 characterized in the literature [6], [18], [19], [20], [21], [22]. Prevéy observed that there is a threshold 73 value (3% of work hardening) below which no relaxation of residual stresses is observed [6]. Above this 74 threshold, many authors have shown that the greater the work hardening induced by shot peening, 75 the greater the relaxation of residual stress [6], [18], [19], [21], [22]. These studies explain that residual 76 stress relaxation is related to the density of dislocations in the material: the higher the number of 77 dislocations, the more the residual stresses relax. It is in particular observed that the laser shock 78 process, known to induce low work hardening levels, results in much lower residual stress relaxation 79 than shot peening [6]. Prevéy also established that residual stress relaxation is influenced by the level 80 of work hardening and not the method of work hardening introduction.

81 The residual stress and work hardening profiles resulting from the different manufacturing processes 82 used for industrial parts vary considerably. Consequently, controlling the relaxation of residual stresses 83 has become challenging and requires the development of appropriate models. The most widely used 84 model for the thermal relaxation of residual stresses is the Zener-Wert-Avrami model [16], [20], [23], 85 [24], [25]. It predicts stress relaxation without taking work hardening into account. Its parameters are 86 therefore only identified for a specific operating point (a microstructure, a position in a sample, a 87 manufacturing process, and a temperature). Hoffmeister et al. proposed a more complex model to 88 account for microstructural effects [16]. Here again, the model depends on a specific operating point 89 and is not able to predict the relaxation of residual stresses in depth unless a large set of measurements is available to identify a set of parameters for each test condition. Thus, although the influence of work 90 91 hardening on the relaxation of residual stresses has been clearly established through several 92 experimental observations, to date, there is no model available in the literature taking these effects 93 into account. It should also be noted that the influence of the microstructure and its relationship to the 94 effect of work hardening is relatively rarely, if ever, discussed in the literature.

The purpose of this paper is to investigate the impact of work hardening and microstructure on the thermal relaxation of residual stresses caused by shot peening on Inconel 718. The main objective is to propose a model that accurately accounts for these effects. This paper is composed of three sections. The first section describes the experimental methods starting with the description of the reference material microstructure of Inconel 718. In addition, this section presents the different thermal

treatments proposed to obtain two complementary microstructures, as well as shot peening conditions and procedures to characterize residual stresses and work hardening. The second section presents and compares the results obtained after shot peening and after thermal relaxation at 550°C of shot peened parts. The influence of work hardening and microstructure on the relaxation of residual stresses is discussed. The final section describes the proposed model for thermal relaxation of residual stresses accounting for the influence of work hardening. The developed analytical model is validated on a large set of data using complementary results from the literature.

107 1 Materials and methods

108 **1.1 Inconel 718**

109 The investigated material is Inconel 718 with a γ matrix; its composition is given in Table 1. The influence of the microstructure is evaluated through the study of three different microstructures. First, 110 111 the *reference* microstructure, obtained after manufacturing and corresponding to the "as-received" material, known as Direct Aged (DA). The other two microstructures, namely coarse grain 112 microstructure (CG) and coarse grain and coarse strengthening precipitates microstructure (CGCP), 113 114 exhibit parameters that have been modified by means of a specific heat treatment history. The 115 microstructure of CG was achieved by annealing at 1040°C for 30 minutes, followed by δ phase precipitation treatment at 955°C for one hour, and finally by conventional γ'/γ'' strengthening 116 117 precipitation treatment (8 hours at 720°C followed by cooling at 50°C/h and aging at 620°C for 8 hours). 118 The CGCP microstructure is characterized by coarse grains and large strengthening precipitates. The 119 heat treatment sequence applied to the CG microstructure was also used, except for the final 120 precipitation treatment of the γ' and γ'' phases. To increase the size of the precipitates, an overaging process was applied at 750°C for 50 hours. Please note that throughout the rest of this article, a color 121 122 code has been used to refer to the three microstructures: the DA microstructure is referenced in blue, the CG microstructure in red and CGCP microstructure in orange. 123

Ni	Fe	Cr	Мо	Al	Ti	Nb	Si	С
54.18	17.31	17.97	2.97	0.56	1	5.39	0.1	0.023

124

Table 1. Composition of the l	nconel 718 DA Alloy (Wt%).
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125 **1.2** Shot peening conditions

The shot peened samples are parallelepipeds of 20x15x10 mm in size. The samples underwent mechanical polishing to achieve a mirror finish before shot peening. The aim was to reduce residual stresses and work hardening caused by machining. XRD measurements were conducted to confirm the absence of residual stress (see Section 2.2). The samples underwent shot peening on one of their 130 largest surfaces using a conventional nozzle and S110 steel shot. Two shot peening conditions (SP1 and 131 SP2) were defined according to AFNOR standard NFL 06-832 [26]. The shot peening parameters (see 132 Table 2) were chosen for two reasons: (i) they are commonly applied to industrial components and (ii) they induce very different residual stress and work hardening gradients, which is necessary to validate 133 the proposed methodology. The shot peening conditions were adjusted to take account for the fact 134 that the change in grain size results in a reduction in the strength of the modified microstructures. As 135 136 a consequence, the conditions shown in Table 2 were chosen so that the coverage was identical for all 137 three microstructures.

Name	Shot	Almen Intensity	Coverage
SP1	S110 steel	12-13A	125%
SP2	S110 steel	22-23A	200%



Table 2. Shot peening conditions applied to the different microstructures.

139 **1.3 Thermal loading conditions**

After shot peening, the samples were placed in a vacuum furnace and subjected to a temperature cycle without any mechanical loading; Figure 1 illustrates the corresponding cycle. The rise in temperature lasts 20 minutes; the temperature is then maintained at 550°C during a time lapse t_h called the "holding time" in the present article. Five different holding times were applied in this experimental campaign, ranging from 0 to 500 hours. The samples were then placed in an argon stream for rapid cooling. Note that $t_h = 0$ corresponds to a sample that has undergone an increase and a decrease in temperature without any holding time.



- 148 Figure 1. Diagram of the thermal loading cycle applied to the samples with the definition of the holding time.
- 149
- 150

151 **1.4 Material characterization**

152 **1.4.1 Scanning Electron Microscope observations**

The microstructural observations were conducted using a MERLIN Scanning Electron Microscope (SEM) with an accelerating voltage of 5 kV and a nominal current of 40 nA. To reveal the microstructure, including the strengthening precipitates γ' and γ'' , an ion polishing system was used to obtain a mirrorpolished surface. The grain size was quantified using the Electron Backscattered Diffraction (EBSD) method. The data was acquired using a NORDIF CD camera operating at 30 fps coupled to a MERLIN SEM operating at 20 kV acceleration voltage and 40 nA nominal current.

159 **1.4.2 X-ray diffraction measurements**

160 X-ray diffraction measurements were conducted on the 311-diffraction peak at a 2ϑ angle of 161 approximately 111° using a Seifert PTS diffractometer. The radiation source was a Co-K α tube with a wavelength of 1.79 Å, an accelerating voltage of 20 kV, and a nominal current of 4 mA. The sample 162 163 preparation was identical to that used for microstructure characterization, as described above. Under 164 these conditions, the X-ray penetration depth was estimated to be between 2 and 4 μ m, and the 165 analyzed surface was a few millimeters wide. The X-ray data were processed using classical methods 166 [27], specifically the $sin^2\psi$ method with analysis of eleven ψ angles to determine the depth profiles. 167 Line broadening was used as a measure for work hardening (see Section 1.4.3 for more details). This 168 was defined by the full width at half maximum (FWHM) of the peaks obtained by a Voigt pseudo-profile fit. For in-depth evaluation of the quantities of interest, electrolytic polishing with an average step of 169 170 40 µm was carried out. A correction method was applied which showed that the material removal had 171 a negligible effect on the residual stress values. Residual stresses and peak widths were measured in 172 two directions, but no significant differences were found. This confirms that shot peening induces an 173 in-plane equi-biaxial stress state. The FWHM depth profile was evaluated before shot peening, and a 174 consistent FWHM value was observed for all untreated samples, as shown by the triangular data points 175 in Figure 5. After shot peening, three residual stress and FWHM profiles were determined at different 176 locations on the surface of the samples for each microstructure to study the dispersion of these 177 parameters. Following heat treatment, one residual stress and FWHM profile was measured for each 178 microstructure.

179 **1.4.3 Procedure of work hardening characterization**

180 If the method for evaluating the residual stress values from X-Ray diffraction measurements is well
181 established, the evaluation of the work hardening requires the implementation of a calibration method
182 [17], [22], [28], [29], [30]. This method has been successfully applied to the microstructures studied in

this paper [11], [12] and has demonstrated its interest in validating the modeling of the shot peening process [31]. Work hardening is identified with the equivalent plastic strain noted ε_p as discussed in [11]. The relationship between the FWHM obtained by X-ray diffraction and the work hardening is presented in Figure 2 for the three Inconel 718 microstructures and is modeled by [17]:

$$FWHM(\varepsilon_p) = A_{FWHM}(1 - e^{-B_{FWHM}\varepsilon_p}) + C_{FWHM}\varepsilon_p + D_{FWHM}$$
 Eq. 1

187 The parameters that have been evaluated are A_{FWHM} , B_{FWHM} , C_{FWHM} and D_{FWHM} [11]. They are given in 188 Table 3. The work hardening induced by shot peening can be determined by inverting Eq. 1. A typical 189 work hardening profile obtained by this method is shown in Figure 3.

190 The parameters for the work hardening characterization were identified for each microstructure; this 191 method can be used only if no microstructural modification is observed during any of the treatments 192 (thermal or mechanical) that the samples undergo [11], [12]. Scanning electron microscope analyses 193 were therefore performed after shot peening and after the thermal treatment to assess the evolution 194 of the microstructure.



195

Figure 2. The full width at half maximum (FWHM) as a function of equivalent plastic strain for different load
 histories, temperatures, and microstructures with corresponding calibration curves. [11].

	Direct Aged (DA)	Coarse grain (CG)	Coarse grain and coarse precipitates (CGCP)
A_{FWHM}	0.66	0.56	0.35
B _{FWHM}	31.12	23.92	20.00
C _{FWHM}	0.99	0.	.90
D _{FWHM}	1.04	1.00	1.18

199Table 3. The constants' values obtained for calibrating the full width at half maximum (FWHM) for the three200microstructures, as defined by Eq.1. [11]. Note that all constants are estimated to the nearest 0.01.

201 1.4.4 Experimental plan

To conclude this Materials and Methods Section, the complete X-ray diffraction measurement plan is presented in Table 4. For each condition and each microstructure, the number of profiles acquired in depth is indicated. Figure 3 also presents the syntax used to describe the shape of the shot peen induced residual stress and work hardening profiles.

	Microstructure								
	Di	rect Aged (D/	A)	Coarse grain (CG)			Coarse grain and coarse precipi- tates (CGCP)		
Holding time t_h (h)	Unshot	SP1	SP2	Unshot	SP1	SP2	Unshot	SP1	SP2
Untreated	3 p	3 p	3 p	3 p	3 p	3 p	3 p	3 p	3 p
0	-	1 p	1 p	-	1 p	1 p	-	1 p	1 p
0.5	-	1 p	1 p	-	1 p	1 p	-	1 p	1 p
1	-	1 p	1 p	-	1 p	1 p	-	1 p	1 p
5	-	1 p	1 p	-	1 p	1 p	-	1 p	1 p
50	-	1 p	1 p	-	1 p	1 p	-	1 p	1 p
500	-	1 pp	1 p	-	1 pp	1 pp	-	1 pp	1 pp

206 207

208

Table 4. X-ray diffraction measurement matrix. p: profile with measurements up to at least 300 μm depth for residual stresses and work hardening; pp: partial profile, up to 150 μm.



Figure 3. Syntax used to describe the profiles of residual stress and work hardening induced by shot peening induced.

211 **2 Experimental results**

212 **2.1 Microstructural observations**

213 To detect any microstructural evolution, SEM observations were carried out on the three investigated 214 microstructures before shot peening, after shot peening alone, and after shot peening followed by a 215 holding time of 500 hours at 550°C. The results of these observations are presented in Table 5. Only 216 the results with the higher shot peening condition (SP2) are detailed here, as similar results were 217 obtained for the SP1 shot peening condition. Table 6 summarizes the microstructural characterizations 218 obtained from SEM observations and EBSD analyses before shot peening. An average grain size of about 219 5 μ m is observed for the DA microstructure, while the other two microstructures have average grain 220 sizes around 35 μ m. The size of the strengthening precipitates is similar for the DA and CG 221 microstructures (10 to 20 nm), but much larger for the CGCS microstructure with a size of 100 to 200 222 nm. Binarization of the SEM images was performed to assess the morphology and distribution of the δ phase. A form factor of 0.9 was arbitrarily chosen to differentiate needle-shaped from globular 223 224 precipitates. The proportion of the δ phase in the CG and CGCP microstructures is between 1.4% and 225 1.7%, values that remain in the same order of magnitude as for the DA microstructure (2.3%). The 226 proportion of needles and globules (40% and 60%, respectively) is also similar for the three 227 microstructures. Comparing the DA microstructure with the CG microstructure thus gives access to the 228 effect of grain size. Similarly, the comparison of the CG and the CGCP microstructures allows studying 229 the influence of the size of the strengthening precipitates.

230 The SEM characterization of a sample shot peened under SP2 conditions, the most severe, shows a 231 great stability of the microstructure (Figure 4). It confirms that the shot peening conditions applied in 232 this work do not induce any change in grain size or orientation, or in the size of the strengthening 233 precipitates. Similarly, the SEM characterization performed after the thermal loadings at 550°C 234 revealed no microstructural change in terms of grain size or strengthening precipitate size, showing 235 morphologically stable microstructures over time (Table 5). The temperature of 550°C used in this study 236 for the thermal loading does not induce any microstructural change in the strengthening precipitates, 237 even in the presence of work hardening at the material surface. It should be noted that an increase in 238 the rate of y" precipitates was observed in a previous study for shot peened samples subjected to aging 239 treatments at 700°C [32]; this can be explained by the higher temperature of the treatment. These 240 analyses validate that, since no microstructural change is observed, the calibration curves shown in 241 Figure 2 can be used to evaluate the work hardening gradient introduced by shot peening, before and 242 after the thermal loading.



peening, after shot peening (SP2) and after shot peening (SP2) and 500 hours at 550°C. The results obtained

for the SP1 shot peening condition are similar.

	Direct Aged Coarse grain microstructure microstructure		Coarse strengthening precipitates microstructure
Average grain size (µm)	4.2	34.7	36.8
Strengthening precipitates size (nm)	Strengthening precipitates size (nm) ~ 10 - 20		~ 100 - 200
δ phase area fraction	2.3%	1.4%	1.7%
δ phase morphology	43 ± 2% needles 57 ± 2% globules	36 ± 2% needles 64 ± 2% globules	41 ± 2% needles 59 ± 2% globules

247

Table 6. For the three investigated microstructures: mean size of the grain and of the strengthening



precipitates, and δ phase characteristics.





Figure 4. SEM images at different depth for DA microstructure after SP2 shot peening.

251

252 2.2 Residual stresses and work hardening after shot peening

The residual stress and work hardening profiles determined by X-ray diffraction for the three microstructures before shot peening and after *SP1* and *SP2* conditions are now compared (Figure 5). The dashed lines correspond to a fit obtained for each microstructure and each shot peening condition. Despite the fluctuations from one sample to another, clear trends are obtained between the two investigated shot peening conditions.

The work hardening is clearly affected by the shot peening conditions, both in terms of value and depth. For example, for the *DA* microstructure, the average surface value is changed by almost 50% between the *SP1* and *SP2* conditions ($WH_{surf}^{\Box} = 20\%$ and $WH_{surf}^{\Box} = 40\%$ respectively). An increase in shot peening coverage and intensity results in a higher maximum compressive residual stress σ_{max}^{\Box} . The associated depth is also affected ($z_{\sigma_{max}^{\Box}SP1} = 50 \ \mu m$; $z_{\sigma_{max}^{\Box}SP2} = 80 \ \mu m$), as the size of the compression zone ($z_{\sigma_{0}^{\Box}SP1} = 175 \ \mu m$; $z_{\sigma_{0}^{\Box}SP2} = 225 \ \mu m$). At the surface, the residual stress values are relatively similar 264 $(\sigma_{surf}^{\Box}SP1 \approx \sigma_{surf}^{\Box}SP2 \approx -700 \text{ MPa})$. Similar trends are observed for both *CG* and *CGCP* microstructures. 265 The residual stress differences remain below 100 MPa along the profiles when comparing the three 266 microstructures. Thus, changing the grain size or the strengthening precipitates size has little impact 267 on the residual stresses. However, the work hardening is strongly affected by the microstructural 268 modifications. An increase in grain size implies an increase in surface work hardening of more than 25% 269 for both investigated shot peening conditions. However, the thickness of the work hardened area is 270 only slightly altered.



274

These results highlight the influence of both the shot peening conditions and the microstructure onthe residual stress and work hardening profiles.

Table 7 presents the minimum and maximum values encountered for each condition for the *DA* microstructure (see notations in Figure 3) with the purpose of characterizing experimental dispersion. This experimental dispersion may come from: measurement errors, microstructural variation from one sample to another, or the non-repeatability of the shot peening process. These three possible sources of dispersion have been investigated and characterized. They are described in detail below. The measurement conditions are the same for all the samples, so the error is evaluated for all
 measurements around 20 MPa. Therefore, the observed variations cannot be specifically due
 to this aspect.

Concerning the influence of the microstructure, the analyses in terms of grain size, distribution,
 and size of strengthening precipitates or crystalline orientation do not show any notable
 difference between the different samples. Looking at the *DA* microstructure compared to the
 two modified microstructures, the small differences in terms of the average value of the
 measured residual stresses or work hardening (Figure 5) show that the microstructural
 variations, even of significant, can be neglected in front of the experimental variability related
 to the shot peening itself.

292 Indeed, it is classical observed that the shot peening process shows a wide scatter of 293 experimental residual stress results [23]. This dispersion is generally related to an inherent 294 variability of many parameters related to the shot stream, such as size, shape, hardness, impact 295 angle and shot peening time. One way to account for this dispersion is to use the Monte Carlo 296 simulation technique by introducing variability in the process parameters to predict the 297 residual stress field [33], [34]. The influence of the shot peening process on the repeatability 298 of the measurements appears to be the most detrimental aspect of the experimental 299 dispersion. Since it is difficult to control, a repetition of the measurements on different samples 300 is often recommended to obtain a better accuracy. Experimental standard deviations of the 301 order of 100 MPa for the residual stresses and of the order of 10% for the work hardening were 302 obtained at the surface for all the microstructures. The standard deviation seems to decrease 303 with depth. These results are consistent with those obtained by Daoud et al. in the simulation 304 of the shot peening process [35].

	SF	21	SI	2
	Min	Max	Min	Max
σ^{III}_{surf} (MPa)	-600	-800	-600	-850
$\sigma_{max}^{[]]}$ (MPa)	-800	-1050	-900	-1150
z _{σmax} (μm)	40	60	70	90
$z_{\sigma_0^{\square}}$ (μm)	120	200	200	250
$WH_{max}^{[]}$ (%)	15	30	20	45
$WH_{surf}^{[]]}$ (%)	15	30	20	45
z _{WH} max (μm)	0	0	0	0
z _{WH0} Ξ (μm)	75	170	100	200

305

Table 7. Summary of variations encountered for the two SP1 and SP2 shot peening conditions for the DA

307 2.3 Residual stresses and work hardening after a thermal loading

The residual stresses and work hardening determined after thermal loading are now being studied. The first part is devoted to the study of the profiles. The second part focuses on the evolution of the key parameters identified in Figure 3, which will serve as the basis for a model to estimate the relaxation of residual stresses.

312 2.3.1 Residual stress and work hardening profiles after thermal loading

The residual stress and work hardening profiles were analyzed to track their evolution after the shot 313 314 peened samples were subjected to different holding times at 550°C. As the relaxation under thermal 315 loading is isotropic, only the component in the \vec{x} direction of the stress was studied (see caption of 316 Figure 1). Note that a distinction is made between samples that have not undergone thermal loading 317 and those that have undergone a temperature rise and fall ($t_h = 0$). Except for a few measured points 318 related to the SP2 shot peening condition, the difference is small between the reference and the 319 samples that have undergone a holding time $t_h = 0$: the rise and fall in temperature do not trigger any 320 relaxation.

Figure 6 shows the residual stress and work hardening profiles obtained after thermal loading for the 321 322 DA microstructure, shot peened under SP1 (Figure 6.a and b.) and SP2 (Figure 6.c and d.) conditions. 323 Both shot peening conditions present the same evolution of the profiles. The residual stresses and 324 work hardening induced by shot peening are only partially relaxed during thermal loading, even for the 325 longest holding times. The depth associated with the maximum value of compressive residual stresses $z_{\sigma \square n}$ is not affected by the duration of the thermal holding at 550°C. On the contrary, for the SP1 326 condition, the depth at which the residual stresses become positive $z_{\sigma_{\alpha}^{\square}}$ undergoes a slight decrease 327 328 with a holding time at 550°C, which is not observed for the SP2 conditions. It is interesting to note that 329 the surface residual stress that corresponds to the most intense level of work hardening, experiences 330 the greatest relaxation. This relaxation is even more pronounced for the SP2 condition: at the surface 331 where the work hardening value is about 40%, the compressive residual stress value evolves from -600 MPa to -200 MPa. 332

Figure 7 and Figure 8 show the residual stress and work hardening profiles for the *CG* and *CGCP* microstructures, respectively. It is noteworthy for these modified microstructures that even at very high levels of surface work hardening (over 60% for the *CG* and *CGCP* microstructures under *SP2* conditions), the relaxation of residual stresses is only partial after several hundred hours at 550°C. Again, the depth associated with the maximum value of compressive residual stresses $z_{\sigma_{max}}$ and that at which the residual stresses are positive $z_{\sigma_0^{\Box}}$ appear to be relatively unaffected by the increase holding time. Similar to what was observed in the *DA* microstructure, the higher the level of work

hardening, the higher the relaxation of residual stresses. The highest levels of relaxation are then foundat the surface.

The main information to be drawn from these results is that the effects of shot peening, although partially reduced, are still present even after being exposed to a purely thermal loading at 550°C for several hundred hours. It is therefore of particular interest to be able to model the residual stress relaxation at any point on the part, in order to know the residual stress profile at any point in time.

Figure 6. Residual stress and work hardening profiles, for different holding times at 550°C, for Inconel 718 DA
 microstructure, SP1 and SP2 conditions.

Figure 7. Residual stress and work hardening profiles for different holding times at 550°C, for Inconel 718
 coarse grain microstructure, SP1 and SP2 conditions.

352

353 354

Figure 8. Residual stress and work hardening profiles for different holding times at 550°C, for Inconel 718 coarse grain and coarse strengthening microstructure, *SP1* and *SP2* conditions.

2.3.2 Definition of ratios to quantify relaxation

Ratios are here defined to quantify the relaxation of the residual stresses and the work hardening due to thermal loading. We first define reference profiles corresponding to the profiles determined after shot peening, for which, by definition, there is no relaxation. These reference profiles are respectively noted, $\sigma_{ref}(z)$ for the residual stresses and $WH_{ref}(z)$ for work hardening. The experimental reference profiles are all presented in Figure 5 as a function of the depth *z* for different microstructures and shot peening conditions.

Several profiles of residual stress and work hardening were evaluated after thermal loading for different holding times t_h (see Figure 1 for the definition of t_h). These profiles are noted $\sigma(t_h, z)$ for the residual stresses and $WH(t_h, z)$ for work hardening, respectively, and are presented in Figure 6 to Figure 8 as a

- 365 function of the depth *z* for different microstructures and shot peening conditions.
- To quantify the relaxation, we define the ratios $R^{\sigma}(t_h, z)$ for residual stresses and $R^{WH}(t_h, z)$ for work hardening such that:

$$R^{\sigma}(t_h, z) = \frac{\sigma(t_h, z)}{\sigma_{ref}(z)} \quad and \qquad R^{WH}(t_h, z) = \frac{WH(t_h, z)}{WH_{ref}(z)}$$
 Eq. 2

Thus, when the ratios are equal to 1, no relaxation is observed; if the ratios were equal to 0, all the residual stresses or work hardening would have vanished.

2.3.3 Evolution of the surface and maximum values of residual stresses and work hardening with thermal loading

To simplify the analysis, representative points have been identified on the profiles of the residual stress and work hardening shown in Figure 6 to Figure 8. The evolution of these specific values is here studied as a function of the holding time t_h . For residual stress, the surface residual stress σ_{surf}^{\Box} and the maximum value σ_{max}^{\Box} at depth $z_{\sigma_{max}}^{\Box}$ were considered. For work hardening, the value at the surface WH_{surf}^{\Box} alone has been considered since work hardening is maximum at the surface (see Figure 3 for the definition of σ_{surf}^{\Box} , $z_{\sigma_{max}}^{\Box}$ and WH_{surf}^{\Box}). The following ratios as detailed in Section 2.3.2 can then be defined:

$$R_{surf}^{\sigma}(t_h) = \frac{\sigma(t_h, 0)}{\sigma_{ref}(0)} \quad R_{max}^{\sigma}(t_h) = \frac{\sigma(t_h, z_{\sigma_{max}})}{\sigma_{ref}\left(z_{\sigma_{max}}\right)} \quad R_{max}^{WH}(t_h) = \frac{WH(t_h, 0)}{WH_{ref}(0)} \qquad \text{Eq. 3}$$

The evolution of R_{surf}^{σ} as a function of holding time is shown in Figure 9 for the *DA* microstructure and the two shot peening conditions. A significant evolution of the surface residual stresses occurs in the early stages (before 5 hours); between 5 and 500 hours, the surface residual stresses relax at a significantly lower rate. Very similar results were obtained for the other two microstructures and are thus not presented here.

Figure 9. Evolution of R_{surf}^{σ} as a function of time at 550°C for the *DA* microstructure and the two *SP1* and *SP2* shot peening conditions; note the change in scale for the smaller holding time axis.

The evolutions of R_{max}^{σ} and R_{max}^{WH} are shown in Figure 10 as a function of holding time, for the three 387 388 microstructures and the two shot peening conditions; note that a logarithmic scale has been adopted 389 for the holding time. As a whole, the results enable to identify some trends concerning the evolution 390 of the different variables as a function of holding time (note the scale is logarithmic): (i) a significant 391 evolution of the maximum residual stresses and work hardening occurs for holding times below 5 392 hours; this was also observed for the surface residual stress; (ii) for holding times above 5 hours, the 393 relaxation of the residual stresses is reduced and the work hardening is slowed down, particularly for 394 SP2. Comparing Figure 9 and Figure 10, it is also clear that R_{max}^{σ} is higher than R_{surf}^{σ} . For example, for 395 the DA microstructure, SP2 shot peening condition, and after a holding time t_h = 5h, R_{max}^{σ} = 0.8 while 396 R_{surf}^{σ} = 0.55. This means that the relaxation of residual stresses is maximum at the surface. These 397 results are related to the initial work hardening value introduced during shot peening. The higher the 398 work hardening value, the higher the relaxation of the residual stresses. To visualize this, the values 399 R_{surf}^{σ} and R_{max}^{σ} can be plotted as a function of the initial work hardening. Figure 11 presents the results 400 obtained for two holding times (1 h and 500 h). For a given holding time, a linear relationship is 401 obtained between the normalized unrelaxed residual stress R^{σ} and the initial work hardening. The 402 longer the holding time, the greater the influence of work hardening. It is interesting to note that the 403 three microstructures follow a very similar trend. These results show work hardening is the dominant 404 parameter in the relaxation of residual stresses. The influence of the microstructure on the relaxation 405 of the residual stresses is only of second order.

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Figure 10. Evolution of a) R_{max}^{σ} and b) R_{max}^{WH} as a function of holding time at 550°C for the three microstructures and the two shot peening conditions.

410 Figure 11. Evolution of R^{σ} as a function of the initial work hardening value. R_{surf}^{σ} and R_{max}^{σ} values are 411 indicated by circles and squares respectively. The definition of these parameters is given in Eq. 3. Only results 412 for two holding times (1h and 500h) are presented.

413 **2.3.4 Influence of the microstructure**

If we want to identify the second order influence of the microstructure on the relaxation of residual stresses, it is necessary to remove the influence of work hardening. To do this, it is essential to analyze points with similar work hardening levels, or at least with sufficiently low work hardening levels, so 417 that this parameter has the minimum impact on the analysis. It is interesting to notice that, for the depth $z_{\sigma_{max}}$, work hardening is five times lower than the surface work hardening (see for example 418 419 Figure 6). Moreover, it is relatively similar for the two shot peening conditions and the three 420 microstructures (comparing Figure 6 to Figure 8). The influence of the microstructure can then be determined by observing the evolution of R_{max}^{σ} (Figure 10.a). A holding time of 500 h is chosen to 421 422 analyze the results because the effects are more pronounced in this case. The results are shown in 423 Table 8. An increase in grain size leads to a decrease in the thermal relaxation of residual stresses. The 424 influence of a change in the size of the strengthening precipitates is more questionable. An increase in 425 the size of the precipitates leads to a slight increase in the relaxation of residual stresses.

426 To explain this dependence on the microstructure, we consider the mobility of dislocations. 427 Dislocations are the main vector of the plastic deformation introduced into the microstructure by shot 428 peening. Because of the strong plastic strain introduced during the process, a rearrangement of the 429 dislocations towards lower distortion energies occurs during the first moments at high temperature 430 [23]. The more the dislocation structure is disturbed, the more it has to rearrange itself to reach an 431 equilibrium level. This reorganization within the material leads to a relaxation of the residual stresses. 432 The phenomena observed here are of the recovery process type. However, similarities can be drawn 433 from studies dealing with the influence of microstructure on the creep behavior of Inconel 718. For a 434 creep load, it has been shown in the literature that the influence of microstructural parameters in order 435 of importance in Inconel 718 are: grain size, morphology and distribution of the δ phase and the size 436 of the strengthening precipitates γ'/γ'' [7], [8], [9]. As the grain size increases, the linear density of grain 437 boundaries decreases, leading to a limitation of grain boundary slip and diffusion mechanisms at high 438 temperatures. The δ phase appears to be the most important microstructural parameter on creep at 439 an equivalent grain size. The lower the proportion of δ phase at the grain boundaries, the better the 440 creep resistance. Moreover, a globular morphology is to be preferred to an acicular one to limit creep [7]. Finally, the size of the strengthening precipitates γ'/γ'' can play a major role at fixed grain size and 441 442 δ phase fraction, depending on the structural hardening mechanism involved. In the modified 443 microstructures of this study, the transition from DA to CG microstructure mainly leads to an increase 444 in grain size by a factor of 5 (see Table 5). Thus, the differences observed between the DA and CG 445 microstructures are essentially related to the increase in grain size. Grain size, δ phase density, and 446 distribution are similar between the two modified microstructures (CG and CGCP). The size of the 447 strengthening precipitates is the only factor that causes a factor of ten difference between the two 448 microstructures. According to Sundararaman, the size of the strengthening precipitates that causes a 449 change in the precipitate shearing mechanisms is around 10 nm in the case of Inconel 718 [36]. In the 450 present work, the size of the precipitates is greater than this value (20 and 200 nm see Table 6) for all

451 the studied microstructures, so that a bypass type of Orowan mechanism occurs. When a dislocation 452 bypasses an obstacle, it may create more dislocations. These newly formed dislocations are usually 453 mobile and can move through the crystal lattice. This mechanism can thus increase the number of 454 mobile dislocations in the material [36]. As a result, during holdings at 550°C, the dislocations rearrange more easily as the size of the precipitates increases. This leads to an easier return to an equilibrium 455 456 state and, thus, to a more significant relaxation of the residual stresses. However, since the impact of 457 the size of the strengthening precipitates is of third order, the influence of a modification of their size 458 on the relaxation of the residual stresses appears to be moderate in this work, and this despite a factor 459 of 10 between the two modified microstructures.

	Direct Aged microstructure	Coarse grain microstructure	Coarse strengthening precipitates microstructure
SP1	0.59	0.75	0.71
SP2	0.59	0.73	0.65

Table 8. Value of R_{max}^{σ} after 500 hours of holding time at 550°C for the three microstructures and the two

shot peening conditions.

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3 A model for the thermal relaxation of residual stresses

The experimental results section confirms the previously obtained results in the literature [14], [15], [16], [37]. It also clearly demonstrates the influence of work hardening on residual stress relaxation. It is now proposed to build on these results to provide a model that allows the assessment of residual stress relaxation. The first proposal is to evaluate the existing Zener-Wert-Avrami model using these data. Then, a new model considering work hardening is proposed. The model's validation is conducted by comparing it with our experimental data and complementary results from the literature. Finally, an example is provided to demonstrate the model's application.

471 3.1 The Zener-Wert-Avrami model

The Zener-Wert-Avrami model is an empirical equation widely used in the literature to describe the evolution of a thermally activated process [38]. It was first used by Vöhringer in 1987 to describe the relaxation of residual stresses [23]. Since its introduction, this model has been used by many authors on different materials (steels, nickel-based alloys, titanium and aluminum alloys) [16], [20], [23], [24], [25]. This model allows for the determination of the stress relaxation $\sigma(T, t_h)$ at a given depth relative to the initial stress at the same depth $\sigma_{ref}^{[:]}$, as a function of the holding time t_h and the temperature Tusing the following equation:

$$\frac{\sigma(T, t_h)}{\sigma_{ref}^{\text{III}}} = exp\left(-\left[C \ exp\left(-\frac{\Delta H_A}{k \ T}\right) t_h\right]^m\right)$$
 Eq. 4

- 479 where ΔH_A is the activation enthalpy, *k* is the Boltzmann constant, *C* and *m* are material parameters to 480 be identified for each position in the component.
- 481 In this work, since the tests are all performed at the same temperature, Eq. 4 is reduced to:

$$\frac{\sigma(t_h)}{\sigma_{ref}^{[m]}} = exp(-[A, t_h]^m)$$
 Eq. 5

482 A and *m* have then to be identified for each manufacturing process, each microstructure, and each483 point in the structure.

484 To evaluate the ability of the Zener-Wert-Avrami model to reproduce the experimental results of the 485 present study, the parameters A and m have been identified for the experimental data presented Section 2; a least square method has been applied for each microstructure, for each shot peening 486 condition and for each position, $z_{\sigma_{surf}^{RS}}$ and $z_{\sigma_{max}^{RS}}$. The corresponding sets of values for A and m are 487 488 presented in Table 9. Note that the parameters obtained, A and m, are simply the values that best fit 489 the proposed empirical (Eq. 5) for predicting residual stress relaxation. No physical meaning can be 490 attached to these values. As expected, these values depend on the shot peening conditions and the 491 microstructure. Note that the *m* parameter is the most affected by the position and thus by work 492 hardening. The Zener-Wert-Avrami model is mainly devoted to the prediction of relaxation as a function 493 of the temperature. Due to the substantial impact of work hardening on the thermal relaxation of 494 residual stresses, and the large range of hardening in the structure induced by shot peening, this model 495 is not suitable for the description of the thermal relaxation of a complete residual stress profile or 496 others shot peening conditions.

Parameter	Depth	DA-SP1	DA-SP2	CG-SP1	CG-SP2	CGCP-SP1	CGCP-SP2
۵	$Z_{\sigma_{surf}}$	6.56E-02	1.91E-01	8.21E-02	2.05E-01	3.17E-01	1.43E-01
~	$Z_{\sigma_{max}}$	1.96E-01	1.88E-01	1.24E-01	6.82E-02	2.57E-01	2.11E-01
3	$Z_{\sigma_{surf}^{\square}}$	3.04E-07	5.59E-03	2.31E-05	6.84E-03	1.96E-03	3.03E-03
10	Z	6.28E-05	8.11E-05	2.52E-08	9.89E-12	4.95E-05	7.40E-05

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Table 9. Values of the coefficients identified for the different shot peening conditions and the differentmicrostructures at two depths ($z_{\sigma_{surf}}$, $z_{\sigma_{max}}$).

499 3.2 A model for thermal residual stress relaxation accounting for work 500 hardening

A model, which considers the work hardening state and considers holding time and position, is proposed to predict the relaxation of residual stresses. This model is based on the experimental observations presented in Section 2. The aim of the model is to be independent of the manufacturing 504 process that generated the residual stresses. In addition, its relative simplicity (analytical) is crucial to 505 function as a practical tool for validating the conditions of a prestressing manufacturing process and its 506 parameters in an industrial context. It has been shown that the microstructural impact on the thermal 507 relaxation of residual stresses is small compared to that of work hardening (see Section 2.3). It is thus 508 decided to disregard the influence of the microstructure to define the model. We then study the residual stress ratio R^{σ} (defined Eq.3) at two specific locations $z_{\sigma_{surf}^{\square}}$ and $z_{\sigma_{max}^{\square}}$ as a function of the 509 510 initial work hardening level. These results are presented in Figure 12 for all the investigated holding 511 times; note that in this figure, the microstructure has not been identified to simplify the reading since 512 we disregard its influence. Shot peening conditions are considered on the basis of the level of work 513 hardening they cause. Therefore, in the same way, they are not explicitly shown in Figure 12. A linear 514 regression has been conducted for the data points corresponding to each specified holding time. 515 Comparing the experimental results and the straight line obtained with the linear regression, we make 516 the hypothesis that, for a given holding time, the ratio of unrelaxed residual stresses depends linearly 517 on the initial value of the work hardening and can be expressed as follows:

$$R^{\sigma}(t_h, z) = B(t_h) \cdot WH(z) + C(t_h)$$
 Eq. 6

518 where WH(z) is the initial work hardening value at the depth z (with a range between 0 and 1) and, 519 $B(t_h)$ and $C(t_h)$ are functions that depend on the holding time.

The parameters *B* and *C* are evaluated for each straight line in Figure 12 and for each holding time; the corresponding results are shown in Figure 13. These two parameters exhibit a rapid evolution for the shorter holding times, followed by a stabilization for the longest holding time. It is therefore proposed to describe the evolution of $C(t_h)$ and $B(t_h)$ with a logarithmic function such that:

$$B(t_h) = b_1 \log_{10}(t_h) + b_2$$
 Eq. 7

$$C(t_h) = c_1 \log_{10}(t_h) + c_2$$
 Eq. 8

with b_1 , b_2 , c_1 and c_2 being constant parameters that need to be identified. Considering this formulation, for $t_h = 0$, the model gives $B(0) \neq 0$ and $C(0) \neq 1$. Therefore, the model may appear erroneous for holding times less than 0.029 hours (i.e., 2 minutes). However, given the holding times used in the proposed applications (greater than 30 minutes), and in the interest of model simplicity, this model is considered in the following, as is Eq. 6. Other formulation using exponential function could be used to respect that the residual stress profile being then the initial one as expected (B(0) = 0 and C(0) = 1).

Also note that in the model a residual stress relaxation can be obtained even if there is no work hardening (WH(0) = 0) in the material. Finally, the proposed model that estimates the ratio of unrelaxed residual stresses takes the form:

$$R^{\sigma}(t_h, z) = [b_1 \log_{10}(t_h) + b_2] \cdot WH(z) + c_1 \log_{10}(t_h) + c_2$$
 Eq. 9

microstructure and for different holding times and for the two depth locations $z_{\sigma_{surf}}$ and $z_{\sigma_{max}}$.

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534 Figure 12. Evolution of *R^o* as a function of the initial work hardening level without distinction of

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Figure 13. Evolu

Figure 13. Evolution of $B(t_h)$ and $C(t_h)$ as a function of holding time.

The four material parameters b_1 , b_2 , c_1 and c_2 have been identified for Inconel 718 with a least square method on the data presented in Figure 13. The obtained values are given in Table 10. Note that given the model form in Eq.9, only three holding time values are sufficient to identify the model, such as the initial residual stress and work hardening levels and the residual stress and work hardening levels after 5 hours and 500 hours of holding time.

<i>b</i> 1	<i>b</i> ₂	<i>C</i> ₁	C ₂
-0.130	-0.364	-0.061	0.907

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Table 10. Values of the parameters of the model of Eq. 9 for Inconel 718.

Note that the parameters $B(t_h)$ and $C(t_h)$ could be modified to include a dependence on temperature. By considering different temperatures in an experimental campaign, an exponential form with temperature dependence could be considered for these parameters. This constitutes a perspective on the present work.

549 3.3 Model validation

550 The model proposed in the previous section is now employed to predict the relaxation of residual 551 stresses under varying conditions. These conditions differ from those of the previous analysis used to 552 identify the parameters. Three sets of experimental data are considered:

i) The part of the results presented in Section 2.3 that have not been used to identify the parameters, that is for positions other than $z_{\sigma_{surf}}$ and $z_{\sigma_{max}}$.

ii) Additional measurements that have been performed in the framework of the present study
for the *DA* microstructure and for 50 hours of holding time at 550°C following the protocol detailed
Section 1.3.

558 iii) Data from Prevéy's work [6] which allows the model to be validated under experimental 559 conditions not investigated here.

560 For all these data, a statistical analysis has been carried out to consider the experimental scatter and 561 the possible errors in the identification of the model. The different steps of this statistical procedure 562 are illustrated in Figure 14. With this objective, a pair of initial residual stress and work hardening 563 profiles is first generated using a normal distribution with the same randomly chosen probability. The 564 initial residual stress and work hardening profiles (solid red lines) have been used as the average values 565 of the normal distribution (Figure 14.a). The standard deviation considered in the normal distribution 566 is characteristic of the observed experimental scatter. Fifty pairs of initial residual stress and work 567 hardening profiles have been generated following this procedure (examples of such profiles are shown 568 in green in the Figure 14.b). Then, the model presented in Section 3.2 is applied to all these pairs of 569 profiles to obtain 50 profiles of relaxed residual stresses (solid grey lines in the Figure 14.c). The average 570 of these profiles is then computed (dashed black line).

573 Figure 14. Steps of the statistical procedure used to validate the model of residual stress relaxation. The red 574 curve represents the reference profiles. The green curves represent an example of a pair of profiles generated 575 with the normal distribution.

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576 This procedure has been first applied to the experimental data obtained in the campaign carried out in 577 this work. The results are presented in Figure 15. For all shot peening conditions and microstructures considered, the average values predicted by the model are in accordance with the experimental data. 578 579 It is worth noting that the model reproduces particularly well the surface relaxation of the residual 580 stress level, which is strongly influenced by the high level of work hardening. The methodology 581 implemented permits estimation of the relaxation of the residual stresses for the whole depth affected 582 by the process. Additionally, by incorporating experimental dispersion into the procedure, significant 583 variations in residual stress profiles within similar surface treatment and heat treatment conditions can 584 be accurately reflected. These results are consistent with the experimental observations.

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Figure 15. Comparison between experimental data and modeling. The labels in the graphs indicate: the microstructure – the shot peening condition – the holding time.

The statistical procedure has also been applied to experimental data obtained by Prevéy for Inconel 718 [6]. In Prevéy's work, the residual stress and work hardening profiles have been evaluated for different surface treatments before and after a heat treatment at 525°C for 100 hours; two shot peening conditions (here denoted *SP3* and *SP4*) as well as gravity peening (here noted *GP*) and low plastic burnishing (here noted *LPB*) treatments have been studied. Figure 16 presents the initial residual stress and work hardening profiles extracted from [6] for the different processes. Note that these four surface treatments provide a variety of residual stress and work hardening profiles. The test conditions 595 differ from those of our study (material, measurement conditions, heat treatment temperature); the 596 microstructure is not given. These results thus constitute an interesting test for the model proposed in 597 Section 3.2. Figure 16 depicts the residual stress relaxation profiles obtained through experimentation 598 [6] and the profiles predicted by the model using the statistical analysis outlined above. Once again, 599 the model successfully replicates the trends of the experimental results obtained by Prevéy. Notably, 600 the model accurately estimates the maximum level of residual stress after 100h of holding time at 601 525°C. The relaxation of surface residual stresses could be more accurate, as in the previous example 602 in Figure 15. However, this analysis confirms the relevance and the pertinence of the proposed 603 modeling.

Figure 16. Residual stresses (a) and work hardening (b) profiles produced by shot peening (SP3 and SP4),
 gravity peening, and low plastic Burnishing in IN718 [6].

608 Figure 17. Comparison between experiment [6] and modeling of residual stress profiles after 100h holding

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time at 525°C.

610 **4** Conclusion

611 The aim of this work was to establish a model for the influence of work hardening and microstructure 612 on the evolution of residual stresses under thermal loading. The first step was to analyze the numerous 613 experimental conditions used in this study to assess the effect of work hardening, grain size and size of 614 strengthening precipitates on the relaxation of residual stresses. Residual stress and work hardening 615 profiles were determined on shot peened components for different holding times at 550°C. The 616 calibration methodology previously proposed [11] allows the influence of different microstructures in 617 terms of grain size and precipitate size to be evaluated. It has been possible to establish the order of 618 influence of the parameters on the relaxation of residual stresses under thermal loading. The first order 619 parameter is work hardening. The higher it is, the greater the relaxation of residual stresses. The second 620 order parameter is grain size. An increase in this factor leads to a decrease in residual stress relaxation. 621 Finally, the parameter with the least influence in the study carried out was the size of the strengthening 622 precipitates, where no clear trend could be deduced from the experimental data analyzed.

Based on the experimental results, an analytical model was then proposed which considers the 623 624 influence of the work hardening induced by the surface treatment. Several advantages of this model 625 can be highlighted. The parameters of the model can be identified on the basis of characteristic points 626 present on the residual stress profiles; at the surface and where the compressive residual stresses are 627 at their maximum. As a result, only a few points are required to calibrate the model. Once identified, it 628 can be used to predict the relaxation of residual stresses in depth if the work hardening profile induced 629 by the surface treatment is known. The proposed model is not limited to the case of shot peening. It 630 can now be applied to estimate the relaxation of residual stresses on components that may have a 631 work hardening gradient in the part. As explained in the introduction, shot-peened Inconel 718 parts 632 are subject to temperature fatigue loading. The study carried out here will therefore make it possible to decouple the effects of mechanical loading and temperature. 633

634 **5** Acknowledgement

This work was conducted with the help of the French Technological Research Institute for Materials, Metallurgy and Processes (IRT M2P) under the CONDOR project. The authors would like to acknowledge IRT M2P and all the partners of the project led by IRT M2P. Safran is warmly thanked for its precious collaboration in this work.

6 Data availability

The raw/processed data required to reproduce these findings cannot be shared at this time due to legal or ethical reasons.

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