

On the high cycle fatigue resistance of austenitic stainless steels with surface gradient microstructures: Effect of load ratio and associated residual stress modification

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Clément Dureau, Mandana Arzaghi, Roxane Massion, Yves Nadot, Thierry Grosdidier. On the high cycle fatigue resistance of austenitic stainless steels with surface gradient microstructures: Effect of load ratio and associated residual stress modification. Materials Science and Engineering: A, 2022, pp.142916. 10.1016/j.msea.2022.142916. hal-03601305

HAL Id: hal-03601305 https://hal.univ-lorraine.fr/hal-03601305

Submitted on 9 Mar 2022

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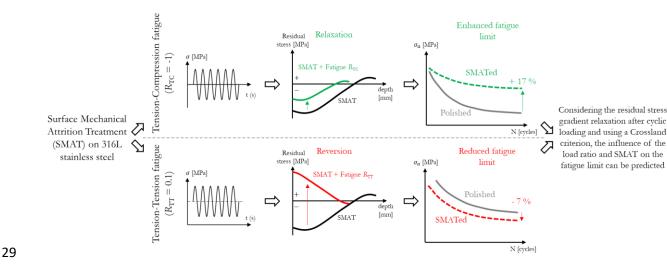
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13 <u>Abstract</u>

The effect of gradient microstructure generated by ultrasonic surface mechanical attrition treatment (SMAT) on the fatigue performance of an austenitic stainless steel has been evaluated in the context of localized and general plasticity by comparing two load ratios: fully reversed tension-compression (TC) R_{TC} =-1 and tension-tension (TT) R_{TT} =+0.1, respectively.

After identical SMAT processing conditions, the fatigue limit was enhanced by +17 % for R_{TC} while it was reduced by -7 % for R_{TT} . Under R_{TC} cyclic loading, self-heating of the specimens and sub-surface martensitic transformation occurred but not for R_{TT} . The residual stress measurements have also revealed that while the stress gradient was smoothed after the R_{TC} fatigue loading, it was completely reverted for the R_{TT} one. In turns, the fatigue crack initiation sites and propagation were modified between the R_{TT} and R_{TC} loading conditions.

Considering the stabilized surface residual stress after fatigue loading, the use of a Crossland criterion allowed to explain both the effects of load ratio and SMAT on the high cycle fatigue behavior of the stainless steel. It turns out that under the studied loading conditions the modifications of residual stress state can be considered as the primary factor governing the varying fatigue performances and the observed triggering at different initiation sites.



Keywords: Surface mechanical attrition treatment (SMAT), ultrasonic shot peening (USP), 316L austenitic
 stainless steel, fatigue, load ratio, residual stresses

32 <u>1. Introduction</u>

In the last two decades, functionally graded materials (FGMs) have been broadly studied, particularly those with varying properties from surface to core. In addition, such materials offer unique mechanical or chemical properties through microstructure or hardening gradients. As a result, automotive, aeronautic, or energy production industries expect them to produce higher efficiency and more reliable components [1, 2], especially those subjected to fatigue issues.

38 In cases such as corrosion, friction wear, or fatigue for which the surface is a critical issue, surface 39 modifications may be sufficient to improve the properties of the mechanical part. In terms of fatigue, as most 40 cracks initiate from the surface, inducing a compressive residual stress state by shot peening improves the 41 fatigue limit of a wide variety of materials [3-7]. Also, surface severe plastic deformation (SPD) processes that 42 lead to FGMs having an ultrafine grain (UFG) surface microstructure represent a way to enhance the functional 43 properties of materials. Among those processes, surface mechanical attrition treatment (SMAT) or ultrasonic 44 shot peening (USP) are derived from conventional shot peening (CSP) [8-12]. They differ from CSP because 45 the motion of the shots is made within a confined chamber so that the shots can impact the surface of the 46 sample, by colliding and bouncing on the chamber walls, for a potentially longer time and with a wider range 47 of incidence angles. This leads to an increased number of activated slip systems allowing to very effectively 48 refine the microstructure leading to a UFG state at the surface and the formation of a gradient 49 microstructure [13, 16]. SMAT or similar treatments were suggested to be efficient ways to enhance the fatigue 50 behavior of many metallic alloys due to the coupled effect of microstructure, hardening gradient, and 51 compressive residual stresses [17-22]. However, some manuscripts suggested that SMAT can be detrimental 52 for some materials or if the chosen set of parameters for the SMAT treatment is not optimized [23-27].

53 The 316L stainless steel is an example of highly used austenitic steel that, despite its good oxidation 54 and corrosion resistances as well as its good formability, suffers from moderate mechanical properties. 55 Different strengthening processes have been studied as a way to improve the mechanical properties of such 56 alloys without giving up too much of their other characteristics [28-30]. By inducing hardening, residual 57 stresses, and UFG microstructure layer, SMAT or similar processes have also been suggested to improve the 58 fatigue behavior of 316L stainless steel. Indeed, Masaki et al. measured a nearly 100 % increase of the fatigue 59 limit of 316L samples after hard shot-peening (HSP) when loaded in rotating bending [31]. Uusitalo et al. [18] 60 and Roland et al. [32] respectively observed an 83 % and 21 % higher fatigue limit after performing SMAT 61 with 3 mm diameter beads on 316L. Comparatively, with similar SMAT treated 316L samples, Sun et al. 62 highlighted that the number of cycles to failure was unchanged compared to the untreated ones in the low cycle 63 fatigue (LCF) regime despite the enhanced stress amplitudes for the SMAT-processed specimens [33]. Here 64 again, on austenitic stainless steels, the literature gives apparently conflicting results. This unchanged fatigue 65 life was proposed to be due to the higher residual stress relaxations induced by the higher strain amplitudes in 66 LCF than in high cycle fatigue (HCF). This was then further investigated by Sun et al., who reported an almost 67 fully relaxed residual stress gradient after a fatigue test conducted at 1.25 % of strain amplitude [34]. 68 Huang et al. even measured a slight tensile residual stresses gradient after 3 % strain [35]. Therefore, this 69 suggests that the macroscopic plastic strains could be affecting the residual stress gradient relaxation after 70 fatigue loading, and then the fatigue life of SMAT processed samples. But, most of the fatigue testing available 71 in the literature [30, 32, 35] have been conducted in fully reversed loading condition (R = -1), and as in this 72 configuration the measured fatigue limits were in the same range of magnitude as the yield strength of the 73 material, no data about the effect of macroscopic strain on the HCF behavior of SMAT processed 316L are 74 available.

75 Thus, the goal of the present work is to shed some light on the exact capability of the SMAT to enhance 76 the high cycle fatigue behavior for different loading conditions by taking the well-known commercial 316L 77 stainless steel as a good test subject. Special attention will be paid to the effect of surface roughness and, as the applied load ratio directly influences the macroscopic strains, to the evolutions of hardness and residual stress
redistributions after fatigue loading.

80 2. Materials and Methods

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2.1. Sample preparation and surface treatment

82 The studied material was an AISI 316L stainless steel formed by swaging above 1040 °C into a 20 mm 83 diameter bar. Table 1 summarizes the chemical composition of this steel together with the resulting martensite 84 start temperature $M_{\rm S}$ and the $M_{\rm d30}$ temperature at which 50 % of martensite is formed when the material is 85 subjected to a 30 % true strain. A heat treatment of 40 min at 900 °C was done on this raw material, resulting 86 in a homogeneous microstructure with an average grain size of approximately 30 µm. Tensile tests revealed a yield stress $\sigma_{vs} = 220$ MPa, an ultimate tensile stress $\sigma_{UTS} = 650$ MPa and an elongation to failure of 105 %. 87 88 Fatigue specimens with a 9 mm diameter and a 12.5 mm gauge length were machined. In order to remove the 89 plastically deformed and heat-affected zone induced by machining, a layer of about 75 µm was removed by 90 grinding the samples before polishing them to a mirror finish surface. While half of the polished samples were 91 kept in this condition (initially polished condition - Table 2 a), the other half was treated by SMAT (SMAT 92 condition - Table 2 a). The treatment was carried at a hitting distance of 20 mm for 60 min using a SONATS 93 ultrasonic shot peening device [36] equipped with a sonotrode made of Ti-6Al-4V that was vibrating at a 60 µm 94 amplitude and a 20 kHz frequency. A total of 4.1 grams of brand new 2 mm diameter 100C6 beads were 95 inserted in the dedicated polyoxymethylene chamber before each treatment and two samples were treated 96 simultaneously. The samples were rotated at a speed of 20 revolutions per minute during this surface 97 mechanical attrition treatment. For specific analysis that will be described later on, some of these SMATed 98 samples were mirror polished after a delicate grinding procedure with 1000 grit SiC abrasive 99 paper (SMAT + polished condition – Table 2 a), resulting in a removed thickness of approximately 35 µm. All 100 the above-mentioned surface conditions are gathered in Table 2 a with their respective surface roughness.

101

2.2. Fatigue tests

102 The fatigue tests were load-controlled to evaluate the overall HCF performance of the studied material 103 for different load ratios. The load ratio is defined by Equation (1), and in order to study its effect on the uniaxial 104 fatigue behavior of the polished and SMAT states of the 316L stainless steel, both tension-compression (TC) 105 and tension-tension (TT) fatigue tests were conducted.

$$R = \frac{\sigma_{\min}}{\sigma_{\max}} \tag{1}$$

107 The following values of the load ratio were selected: $R_{TC} = -1$ and $R_{TT} = +0.1$. The fatigue tests were carried out 108 at room temperature at a loading frequency of 10 Hz with an 810MTS servo-hydraulic machine. Any sample 109 reaching 10⁶ cycles without breaking was considered a run-out and was not reused.

110 <u>2.3. Samples characterization</u>

Fracture surfaces of the broken samples were analysed using optical microscopy (OM) and scanning
 electron microscopy (SEM) to reveal the differences in crack initiation and propagation between the different
 tested conditions.

114 In order to decorrelate the influence of the gradients of microstructure, hardness, and residual stresses, 115 reference SMATed samples were compared to the run-out samples that experienced the highest fatigue tested 116 stress amplitude under R_{TC} and R_{TT} without failure. They were denominated as "SMAT + Fat R_{TC} " and 117 "SMAT + Fat R_{TT}". A final condition was studied, for which a SMATed sample was tested for one single cycle at R_{TT} at the same stress level as the SMAT + Fat R_{TT} one. This sample was analysed to learn about the effect 118 of the number of fatigue cycles; it was named "SMAT + $1Cy R_{TT}$ ". The testing conditions and their labelling 119 120 are gathered in Table 2 b. For these four conditions (SMAT, SMAT + Fat R_{TC} , SMAT + Fat R_{TT} and 121 SMAT + 1Cy R_{TT}), the hardening state characterization was done by standard HV 0.3/20 Vickers micro-122 hardness measurements. They were conducted on specimen cross sections perpendicularly to the SMAT layer 123 starting 30 µm below the surface until reaching the core material. Each indent was spaced by 100 µm to avoid 124 interference, and 10 measurements were conducted for each given depth. In parallel, the in-depth residual stress 125 gradients before and after fatigue loading of those four samples were characterized by X-ray diffraction after 126 successive material layer removal using electrochemical polishing. For electro-polishing, a solution consisting 127 of a mix of 6 % perchloric acid, 35 % butoxyethanol and 59 % methanol was used. The cylindrical fatigue 128 samples were rotated during the XRD measurement to increase the number of diffracting grains and average 129 potential local effects. A Pulstec µ-X360s device with a mounted Cr tube operating at 30 kV and 1.5 mA 130 allowed to obtain full diffraction rings from the two-dimensional detector. The "Analysis Tools for Electron 131 and X-ray diffraction" (ATEX) software [37], developed in LEM3, was used for post-processing the acquired 132 data using the " $\cos \alpha$ " method. Finally, due to the residual stress redistribution occurring during successive 133 material layer removal, the experimentally measured gradient was corrected using the Moore and Evans 134 relations [38].

135 <u>3. Results</u>

136 The results will be presented hereafter into different subsections. The first subsection gathers the 137 fatigue test results. Fatigue tests were carried out at $R_{\rm TC}$ for both the polished and SMAT conditions to set the 138 reference data for easier comparison with the literature. Then similar tests were conducted at R_{TT} in order to 139 highlight the effect of the load ratio. The second subsection details the analysis of the fracture surfaces. The 140 third subsection deals with the analysis of the run-out samples. First, the surface of the run-out samples were 141 investigated using SEM to look for potential fatigue cracks. Also, residual stress as well as hardness evolutions 142 were studied to get further information about the material behavior in the fatigue limit regime of the SMATed 143 samples for both R_{TC} and R_{TT} . Finally, this result section will be concluded by a Crossland initiation criterion 144 analysis taking into account the relaxed residual stress fields to analyse the combined effects of load ratio and 145 SMAT.

146 <u>*3.1. Fatigue properties*</u>

Figure 1 gives the Wöhler curves obtained for the polished and the SMAT conditions for both testing conditions: TC in Figure 1 a and TT in Figure 1 b. Circles represent the polished samples while the squares and stars the SMAT and SMAT + polished conditions, respectively. The full and hollow symbols indicate the broken and run-out specimens, respectively. The experimental data were fitted using the Stromeyer equation [39] defined by Equation (2), a full line was used for fitting the polished state and a dotted line for the SMAT one, while the SMAT + polished data were not fitted due to the small number of tested specimens.

153
$$\sigma_{\rm a} = \sigma_{\rm D} + \left(\frac{c}{N}\right)^{(1/m)} \tag{2}$$

In Figure 1 a, the results obtained by Dureau et al. in a previous study on the 304L stainless steel [40] were superimposed. Indeed, the data reported in Figure 1 a show that the 304L and 316L samples exhibit very similar fatigue behavior for both surface conditions.

For the Wöhler curves of the samples loaded at R_{TC} (Figure 1 a), the fatigue limits are estimated to be 205 MPa and 240 MPa for the polished and SMAT conditions, respectively. Clearly, the SMAT provided an increase of about +17 % of the fatigue limit at R_{TC} . For this R_{TC} loading condition, self-heating of the surface of the specimens as high as 150 and 250 °C was measured for the highest tested stress amplitudes for the polished and SMATed samples respectively. For the samples loaded in the range of the fatigue limit, the self-heating was less than 50 °C and considered negligible, similarly to what was reported in a previous 163 work [40]. For the R_{TT} loading, Figure 1 b indicates that the fatigue life of the SMATed samples was very 164 similar to the polished ones in the 10^5 cycle regime, while it tended to be slightly lower in the 10^6 cycle domain. Under R_{TT} , the fatigue limits were estimated to be 230 MPa and 215 MPa for the polished and SMAT 165 166 conditions, respectively. Comparatively, for the SMAT + polished specimens also loaded at R_{TT} , the fatigue 167 life as well as the fatigue limit of those specimens appeared to be slightly increased compared to the SMATed 168 ones due to the post-polishing procedure. Contrary to what was observed for the R_{TC} condition, a reduction of 169 about -7 % is induced by the surface mechanical attrition treatment when the samples are loaded at $R_{\rm TT}$. The 170 initially polished and SMAT + polished data almost superimpose for the $R_{\rm TT}$ conditions. This indicates that the 171 actual surface state has an influence since the post-polishing procedure restored the overall fatigue properties. Interestingly, contrary to what was reported at R_{TC} , no self-heating of the specimens was measured for this load 172 173 ratio.

174 <u>3.2. Broken samples analysis</u>

Figure 2 presents typical SEM images of the characteristic initiation sites observed on the different tested specimens. In Figure 2, the three horizontal lines correspond to the surface state of the samples (initial polished, SMAT, or SMAT + polished) while the columns rank the observations made for the two load ratios, R_{TC} or R_{TT} , with images of samples that broke under high or low stress amplitudes.

179 For the $R_{\rm TC}$ loaded samples, the comparative analysis of the fracture surfaces - at a given stress 180 amplitude for the polished and SMAT conditions - revealed very similar fatigue crack initiation features. As 181 marked by the red arrows in Figure 2 a1, the high stress amplitude is witnessed by the presence of a secondary 182 crack intercepting the fracture surface in the vicinity of the initiation site of the polished sample. For the SMAT 183 samples subjected to a high stress amplitude, as illustrated in Figure 2 b1, no such crack was observed; instead, 184 a very distinct river line aspect was revealed. For the low stress amplitude conditions, as shown in Figures 2 a2 185 and b2, no difference was noticed on the initiation sites regardless of the surface condition. Thus, the observed 186 modification of the crack initiation was more influenced by the applied stress amplitude than by the exact 187 nature of the surface.

For the samples loaded at R_{TT} at high stress amplitudes, all the samples experienced fatigue crack initiation on surface inclusions whatever the exact nature of the surface condition as seen in Figures 2 c1, d1 and e1. Moreover, all the observed inclusions exhibited a rather circular shape except for the SMAT conditions. Indeed, compared to the round solid inclusion shown in Figure 2 c1 for the polished sample, the inclusion seen 192 in Figure 2 d1 for the SMAT sample exhibits numerous sharp angles and facets. Indeed, in the case of rather 193 large particles located at or just beneath the surface, the repeated impacts can result in their fragmentation 194 directly under the action of SMAT. This can explain why fragmented inclusions were observed in some SMAT 195 samples (Figure 2 d1) but not in the polished ones (Figure 2 c1). Figures 2 e1 highlights, once again, an 196 initiation on a surface solid inclusion for the SMAT + polished condition; even if the inclusion was even ejected 197 from the fracture surface for this specific sample, leaving a cavity. For the lowest stress amplitudes, initiation 198 also occurred at the surface for the three conditions. However, inclusions were never observed at the initiation 199 site of the analysed initial polished and SMAT conditions. Figure 2 c2 demonstrates that, as no inclusion is 200 found, fatigue cracks are probably initiated on surface roughness induced by plasticity such as slip bands. In 201 contrast, Figure 2 d2 shows that for the SMAT, initiation occurred on a dimple (i.e., a surface asperity induced 202 by SMAT), as will be confirmed in Figure 3. For the SMAT + polished samples loaded at low 203 stress (Figure 2 e2), initiation occurred at a surface inclusion as previously described for the R_{TT} high stress 204 conditions. However, due to the lower number of tested samples, it seems speculative to conclude that no 205 change in the initiation mechanism is observed when reducing the stress amplitude in the case of 206 SMAT + polished samples.

207 Figure 3 gathers the EDX maps corresponding to the four areas surrounded by red rectangles in 208 Figure 2. Only the case of R_{TT} samples is presented because the analysis carried out on the initiation sites of 209 the $R_{\rm TC}$ samples did not reveal any inclusion or chemical heterogeneity. As shown in Figure 3 a, the inclusion 210 found at the initiation site of the polished sample loaded at high stress amplitude is a rather complex calcium-211 aluminate oxide also containing magnesium. The inclusion visible in Figure 2 e2 for the case of a low stress 212 SMAT + polished sample loaded at R_{TT} was another calcium aluminate oxide containing magnesium and 213 silicon. The inclusions could also be simple aluminium oxides, as shown in Figure 3 b for a SMAT sample 214 loaded at high amplitude. The most probable explanation for the presence of these different oxides is that they 215 are initially formed between the molten metal and the slag by the addition of deoxidation products during the 216 argon oxygen decarburation process. As explained by Zhang et al. [41], this usually leads to the formation of 217 characteristic oxides based on SiO₂, CaO, and Al₂O₃. These oxides being solid in the molten metal due to their 218 higher melting temperature, they are present in the final product as inclusions of different oxide chemistry. 219 These observed inclusions have a fairly nodular shape with a diameter of about 10 µm regardless of their 220 composition. On the other hand, Figure 3 c reveals that the dimple defect observed in Figure 2 d2 is filled by 221 aluminium, titanium, and vanadium. This chemical composition corresponds to the characteristic

222 contamination layer that can form during sufficiently long surface mechanical attrition treatment. Such type of 223 contamination layer is known to form by erosion ablated scraps coming from the 100C6 shots, the Ti-6Al-4V 224 sole of the sonotrode or even particles from the chambers. Such contamination layer was already observed by 225 Samih et al. and was even shown to reach depths of approximately 5 µm due to the pile-up, ablation, and 226 deposition mechanisms for similar SMAT conditions [42]. These observations were later confirmed by 227 Chemkhi et al. on SMATed 316L stainless steel [43]. This contamination layer is known to affect the reactivity 228 and corrosion resistance of the SMAT treated surfaces [10]. Its chemistry was also shown to have determining 229 consequences regarding the wear behavior of the shot peened surfaces due to its effect on the exact nature of 230 the abrasive oxides that form during dry contact friction [44].

231 Figure 4 compares the macroscopic optical observations of the fracture surfaces of the different tested 232 specimen conditions (line a) with sketches of the fracture sequence with topology of these fracture surfaces 233 (line b). For the R_{TT} loading conditions, the observed features for the SMAT and the SMAT + polished samples 234 were similar; thus, only the SMAT condition is shown. For the R_{TC} loading condition of the polished sample 235 (Figure 4 a1), the observation shows a fairly straight propagation front perpendicular to the propagation 236 direction (i.e., sample radius) before the final fracture. Comparatively, the propagation region forms an internal 237 circular-like area for the SMAT samples (Figure 4 a2). The same behavior was observed in a previous study 238 on a 304L stainless steel treated by SMAT and loaded at R_{TC} [40]. For the R_{TT} loading conditions (Figure 4 a3 239 and a4), semi-circular crack fronts were observed before the occurrence of a ductile fracture for all conditions. 240 At this scale, no obvious difference in the shape of the fracture surfaces was observed when comparing the 241 different polished and SMAT samples loaded at R_{TT}. Figures 4 b1, b2, b3, and b4, give the topology of the 242 fracture surfaces and the initiation site was taken as the reference point for the measured altitude. The 243 propagation plateau in the case of the $R_{\rm TC}$ polished sample is perfectly perpendicular to the loading axis, as 244 indicated in Figure 4 b1 by the large domain of blue colour pointed by the red arrow. Comparatively, in the 245 central region of the fracture surface of the $R_{\rm TC}$ SMAT sample, a slight angle of about 7 ° was observed 246 perpendicular to the crack propagation direction due to the higher stress amplitude, as shown in Figure 4 b2. 247 In addition as indicated by the dotted black arrows, the fatigue fracture surface delimited by the circle-like 248 dotted line in Figure 4 b2 continued toward the hardened surface by a propagation zone inclined at 45 °. Minor 249 differences can be noticed on the fracture surfaces when comparing the $R_{\rm TT}$ conditions (Figures 4 b3 and b4). 250 However, inclined propagation zones are visible in the vicinity of the surface for the shot peened samples, as 251 outlined in Figure 4 b4 by the regions delimited by the curved dotted lines and pointed by the red arrows. These

252 specific crack propagation features for the SMAT samples loaded at R_{TC} and R_{TT}, because they were not visible 253 for the initial polished samples, are then linked to the presence of the SMAT affected layer. They can either 254 result from the microstructure and especially texture effects, as was already discussed for several UFG metallic 255 materials such as interstitial-free steel [45], aluminium alloy [46], or commercially pure copper [47]. As the 256 texture modifications under SMAT are rather weak [48], the modifications in crack propagation are more likely 257 due to the tri-axial field of residual stress generated in this region. It leads to a curved crack propagation path, 258 as discussed by Cotterell and Rice [49]. As shown by comparing Figures 4 b1, b2, b3, and b4, the failure 259 sequence always consists of three stages. For all conditions, it starts with a surface crack initiation followed by 260 a propagation phase and finishes with a ductile fracture. However, as confirmed by the larger domains of grey 261 contrast in Figures 4 b3 and b4, the ductile fracture occurred after much shorter fatigue crack propagation for 262 the $R_{\rm TT}$ than for the $R_{\rm TC}$ ones (Figures 4 b1 and b2). This is expected since, for an identical stress amplitude, 263 the maximum macroscopic stress is more than twice as high for the $R_{\rm TT}$ loading condition than for the $R_{\rm TC}$ one. 264 Also, due to this much higher stress while breaking, samples loaded at R_{TT} experienced necking and bending, 265 leading to a much more deformed final section (Figures 4 b3 and b4) compared to the R_{TC} samples for which 266 the original circular section can still be clearly distinguished (Figures 4 b1 and b2).

267

3.3. Unbroken samples analysis

268 Figure 5 presents the characteristic surface state of the run-out samples observed by SEM. For R_{TC} the 269 images are not presented because, for the polished samples, the observation only revealed fine witness marks 270 of plastic activity while, for the SMAT samples, observation was nearly impossible due to the roughness of the 271 surface and the presence of the contamination layer (previously discussed in Figure 3 c) partially covering the 272 sample surface. For R_{TC} , Figure 5 a reveals fine witness marks of plastic activity visible on the polished 273 surface. Some favourably oriented grains exhibit slip bands which usually are parallel. Such traces were not 274 visible for the SMAT samples (Figure 5 b) due to the roughness of the surface and the presence of the 275 contamination layer (previously discussed in Figure 3 c) partially covering the sample surface. For R_{TT} , as 276 shown in Figure 5 c, the surface of the polished samples is very affected by the fatigue loading. Indeed, it is 277 covered with highly pronounced slip bands and the deformation of the grains creates a micro-roughness leading 278 to the classic "orange-peel" aspect of the deformed samples. For the $R_{\rm TT}$ loaded SMAT samples, Figure 5 d 279 illustrates that the contamination layer covering the SMAT sample surface is not homogeneous. Indeed, in 280 Figure 5 d, the yellow dashed curve delimits the left end side where the rough surface of the fatigue sample is 281 visible from the right end side where the contamination layer appears in a darker colour. Once again, the rough 282 SMAT surface does not allow to identify surface plasticity. Comparatively, as indicated by the red arrows, it 283 is interesting to notice that the contamination layer of the samples has fractured during the fatigue loading due 284 to the elongation of the stainless steel substrate. For the SMAT + polished condition, slip bands are visible at 285 the surface of the samples (Figure 5 e). However, due to the refined microstructure generated during SMAT, 286 the slip bands are not as straight as for the initial polished state. Figure 5 has shown that different surface states 287 and plastic activity were identified for the different studied conditions. Nevertheless, fatigue crack initiation 288 was never detected for run-out samples regardless of their surface condition or the imposed stress amplitude 289 and load ratio. This indicates that initiation seems to be the key aspect to consider when studying the effect of 290 SMAT on the HCF behavior of stainless steel.

291 Figure 6 gives the evolutions of the longitudinal component of the residual stress tensor with respect 292 to the depth below the surface for different conditions. Figure 6 a compares the stress gradient after fatigue on 293 a run-out sample for the $R_{\rm TC}$ loading (SMAT + Fat $R_{\rm TC}$ condition for $\sigma_{\rm a} = 240$ MPa) with the reference SMAT 294 one. Figure 6 b shows the evolutions of the stress state for R_{TT} loading after one cycle (SMAT + 1Cy R_{TT} for 295 $\sigma_a = 215$ MPa) and after another fatigue run-out (SMAT + Fat R_{TT} also for $\sigma_a = 215$ MPa). The reference 296 SMAT sample characterization is presented by solid black curves, while dotted and dashed lines are used after 297 loading. As the 316L stainless steel samples underwent stress- or strain-induced martensitic phase 298 transformation during the SMAT process, two phases did coexist. Thus, in the graphs, circles are the measured 299 values of residual stress in austenite (γ), while squares are used for martensite (α '). The error bars represent 300 the standard deviation of each linear fit used in the " $\cos \alpha$ " method for each measured point. For clarity, they 301 are only displayed when the standard error is more than 50 MPa. As shown by the solid black curves in 302 Figures 6 a and b, SMAT allowed to create a compressive residual stress gradient along about 600 µm below 303 the treated surface. Due to the stress and strain fields, a partial martensitic transformation occurred in the first 304 100 µm below the surface. This is rather consistent with the observations of Novelli et al. on the 304L stainless 305 steel [50], even if it has to be recalled that the austenite in the 304L is less stable and then more susceptible to 306 undergo martensitic transformation than for the 316L. For the reference SMAT condition, surface residual 307 stresses of approximately -600 MPa and -900 MPa were evaluated in austenite and martensite, respectively. 308 The maximum values of residual stresses were recorded slightly below the surface. They were -750 MPa for 309 austenite at 100 µm below the surface and -1 GPa for martensite at a depth of 25 µm. These results are 310 consistent with observations made by different authors [34, 40, 51]. Interestingly, after fatigue, stress 311 relaxation took place in very different ways depending on the loading conditions. As shown in Figure 6 a for 312 the $R_{\rm TC}$ loading, the surface stresses were relaxed by approximately 50 % in both austenite and sub-surface 313 martensite for the SMAT + Fat $R_{\rm TC}$ sample. This resulted in approximately constant internal stress levels along 314 the first 100 µm below the surface with values of -250 MPa and -500 MPa in the austenite and martensite 315 phases, respectively. In addition to this compressive residual stress relaxation, the depth of the gradient was 316 also reduced down to 275 µm. Moreover, it is very interesting to notice that additional martensite was created 317 during the cyclic loading at depths below 200 µm (down to about 350 µm because deeper investigations were 318 not conducted). Thus, due to the load condition of SMAT followed by fatigue at R_{TC} , martensite was present 319 in a fait amount at depth in the range 0-100 µm and bellow 200 µm but not detected in the 100-200 µm depth 320 domain. For the $R_{\rm TT}$ loading conditions, a reversion of the residual stress state was revealed. Indeed, for the 321 SMAT + Fat R_{TT} sample, surface stresses of +850 MPa and +1250 MPa were measured in austenite and 322 martensite, respectively. Similarly to what was observed on the SMAT + Fat R_{TC} , the residual stresses stored 323 in martensite were relatively constant but, in this case, no additional stress- or strain-induced martensite was 324 formed in the subsurface. In austenite, the internal stress level decreased almost linearly as a function of depth 325 down to 350 µm below the surface. Interestingly, the stress gradient was almost completely reverted after only 326 one cycle of $R_{\rm TT}$ fatigue. Indeed, surface stress of +750 MPa in austenite and +1150 MPa in martensite were 327 measured for the SMAT + $1Cy R_{TT}$ sample.

328 Figure 7 compares the SMAT reference hardness gradient with the ones measured after the different 329 loading conditions. Figures 7 a, b, and c present the results for the SMAT + Fat R_{TC} , SMAT + Fat R_{TT} , and 330 SMAT + 1Cy $R_{\rm TT}$ samples, respectively. Each point represents the average value and the error bars are the 331 minimum and maximum hardness values measured at the corresponding depth. The hardness gradient for the 332 reference SMAT sample before fatigue is displayed in every graph. It is characterized by a maximum value of 333 500 HV measured 50 µm below the surface and an estimated total hardened layer of about 600 µm. These 334 values are consistent with data from the literature on the 316L and 304L alloys after comparable surface 335 mechanical attrition treatments [34, 40, 52, 53]. After cyclic loading for one cycle or more, the curves of the 336 SMAT + Fat R_{TC}, SMAT + Fat R_{TT}, and SMAT + 1Cy R_{TT} samples depict a general gradient evolution that is 337 quite similar to the reference SMAT one. However, when sufficient fatigue cycles were applied (i.e., for the 338 two SMAT + Fat conditions in Figures 7 a and b), a slight softening was detected at the subsurface 339 between 100 and 200 µm (as indicated by the single red arrows), while, as pointed by the double red arrows 340 below 600 µm, a hardening was measured in the so-called "core material". Differences in the behavior of 341 dislocations can explain these two opposite phenomena. Indeed, during cyclic loading, the annihilation of dislocations created under SMAT occurs, thus leading to softening in the subsurface. Similar explanation was given in the work of Sun et al. in which a similar behavior was observed [33]. In the core material, new dislocations were created during fatigue, and the material was hardened. The stabilized value in the core is 220 HV, which represents an increase of about 10 % compared to the initial hardness of the material. The absence of significant differences for the SMAT + 1Cy R_{TT} sample indicates that several cycles of fatigue are needed to reach the stabilized hardened state.

348

3.4. Crossland criterion

349 The fact that the residual stress gradient after fatigue was shown to be drastically different for both 350 loading conditions (RTC and RTT) indicates that not only the macroscopic loading has to be considered to 351 understand the influence of SMAT on the fatigue limits for those two load ratios. Moreover, as for each tested 352 conditions the fatigue limit was shown to be characterised by the incapacity of fatigue cracks to initiate, an 353 initiation endurance criterion taking into account the multiaxial stress field at the surface of the fatigue samples, 354 such as the Crossland criterion [54], seems to be a way to understand the difference of effectiveness of such 355 treatment in the HCF regime. Three hypotheses were considered to establish the Crossland criterion. The first 356 one is that the initial polished samples were exempt from residual stress due to the surface polishing after 357 machining. The second one is that the free surface of the SMAT samples is considered to be under a plane 358 stress state and that the shear components σ_{12} and σ_{21} are negligible compared to the longitudinal and tangential 359 ones σ_{11} and σ_{22} . The third one is that only the residual stress state measured in austenite was considered 360 because the proportion of martensite was limited and not quantified in detail, so that a "rule of mixtures" could not be used. Considering that the estimated fatigue limits are $\sigma_{R_{TC}}^{\text{polished}} = 205 \text{ MPa}, \sigma_{R_{TC}}^{\text{SMAT}} = 240 \text{ MPa},$ 361 $\sigma_{R_{\text{TT}}}^{\text{polished}} = 230 \text{ MPa}$, and $\sigma_{R_{\text{TT}}}^{\text{SMAT}} = 215 \text{ MPa}$, and that the stabilized surface residual stress tensors after fatigue 362 363 loading of the SMAT samples are:

364
$$\underline{RS}_{R_{TC}}^{SMAT} \begin{bmatrix} -229 & \sim 0 & 0 \\ \sim 0 & -512 & 0 \\ 0 & 0 & 0 \end{bmatrix} \text{ and } \underline{RS}_{R_{TT}}^{SMAT} \begin{bmatrix} +872 & \sim 0 & 0 \\ \sim 0 & -225 & 0 \\ 0 & 0 & 0 \end{bmatrix}$$

The following two stress tensors invariants were calculated. $J_{1,\max} = \frac{1}{3} tr\left(\underline{\sigma}_{max}\right)$ and $\sqrt{J_{2,a}} = \sqrt{\frac{1}{2}} \underline{S}_a : \underline{S}_a$ where \underline{S}_a is the stress amplitude tensor deviator defined by $\underline{S}_a = \underline{\sigma}_a - \frac{1}{3} tr\left(\underline{\sigma}_a\right) \cdot \underline{I}$. The results are plotted in Figure 8 for the different load ratios and surface state conditions. The two points corresponding to the fatigue limits of the polished conditions for R_{TC} (hollow circle) and R_{TT} (filled circle) are located in a similar $\sqrt{J_{2,a}}$ area 369 of the graph due to the comparable measured fatigue limits. However, the point representing the polished $R_{\rm TT}$ 370 (filled circle) condition is shifted to the right on the $J_{1,\text{max}}$ axis compared to the R_{TC} (hollow circle) one due to 371 the higher maximum stress during cycle for this load ratio. Comparatively, for both SMAT conditions, the 372 residual stress state acts as a mean stress during cyclic loading. Thus, due to the compressive stress field, the 373 point corresponding to the R_{TC} SMAT condition (hollow square) is pushed in the region of the graph where 374 $J_{1,\text{max}}$ is negative. Because of the surface tensile residual stresses for R_{TT} SMAT (filled square), the 375 corresponding point is shifted to the right end side of the graph. The four points are almost aligned in a slightly 376 decreasing trend indicating the negative effect of the maximum hydrostatic pressure, which is usually observed 377 in the Crossland criterion representations [54, 55]. Moreover, the obtained fitted linear curve separates the 378 graph into two regions. The region below the curve corresponds to the "safety domain" where failure will not 379 occur. In contrast, the region above the line represents the "failure domain" in which cracking will occur, and 380 fracture of the sample should happen.

381 <u>4. Discussions</u>

382 The previous section has shown that the load ratio applied to a specimen has a non-negligible impact 383 on the potential effect of SMAT on the HCF behavior of austenitic stainless steels. Indeed, our results have 384 shown that the SMAT allowed to increase the fatigue limit of samples loaded at R_{TC} = -1 by about +17 % for 385 both the 316L and 304L steels, while a reduction of -7 % was observed when samples were loaded at 386 $R_{\rm TT} = +0.1$. For $R_{\rm TC}$, such improvement has already been observed in fully reversed loading (R = -1), whether 387 the steel was loaded in tension-compression [32, 40] or alternating flexural or rotating bending [18, 40]. In 388 comparison, for R_{TT}, no direct comparison with data from the literature is available for stainless steels treated 389 by SMAT. While mechanical surface treatments are carried out to enhance the fatigue limit of materials, they 390 appear to potentially have the opposite effect and actually reduce it. In such case, it is most of the time attributed 391 to the modified surface roughness and generated surface defects. Thus, the first section of this discussion will 392 deal with the direct effect of the surface integrity on the fatigue behavior. This will be shown to be insufficient 393 to fully explain the differences in behavior. Then, the influence of the plastic deformation mechanisms on the 394 microstructure and hardness as well as the role of the residual stress evolutions and their effect on the fatigue 395 crack initiation mechanisms will be discussed.

396

4.1. Effect of the surface integrity on fatigue crack initiation

397 It is often proposed that the reduction in fatigue properties is due to surface defects that lead to faster 398 fatigue crack initiation and compensate the benefits of the surface compressive residual stresses and hardened 399 layer. This hypothesis can be found in several works focusing on a wide variety of materials and loading 400 conditions. Kumar et al. studied uniaxial tension-tension fatigue at R = +0.1 after 30 min and 60 min of SMAT 401 on the 718 Inconel superalloy [56] and on a Ti-6Al-4V titanium alloy [24]. While minimal effect was obtained 402 on the superalloy [56], the reduction of the fatigue limit observed after 60 min of SMAT in the Ti-alloy [24] 403 was proposed to be due to the increase of micro-damage linked to the increased treatment time. Tian et al. drew 404 a similar conclusion for a C-2000 nickel-based superalloy loaded at R = +0.1 under four-point bending for 405 which, after an improvement of 50 % in the fatigue limit after 30 min of SMAT, the fatigue life was 406 systematically reduced when the treatment duration was increased (60 min, 90 min and 180 min). This was 407 also discussed in the LCF and very high cycle fatigue (VHCF) regimes by Pandey et al. [57] and by 408 Gao et al. [58], who respectively studied a 7075 aluminium alloy and a TC11 titanium alloy loaded at R = -1409 after different SMAT-processing times. For the Al-alloy, the number of cycles to failure was almost unchanged 410 after 30 sec and 60 sec of SMAT, increased for 180 sec but was reduced back to the initial performances after 411 300 sec of surface modification treatment [57]. The number of cycles to failure of the Ti-alloy was reduced by 412 approximately one to two decades for a "mild" surface modification condition and even further for a more 413 drastic SMAT one [58]. These negative behaviours were again attributed to the easier initiation of the fatigue 414 cracks which was cancelling the improved crack propagation resistance due to the compressive residual stress 415 layer [57, 58]. Maurel et al. studied the effect of SMAT duration when combined with ageing heat treatment 416 on the fatigue behavior of two aluminium alloys, namely a 2024 and 7075 series [26]. Some samples were aged 417 before SMAT and some after SMAT. By carrying SMAT on the soft solution treated condition (SMAT before 418 ageing), the fatigue limits were unchanged for the 2024 alloy and reduced by about 15 % for the 7075 one. 419 Comparatively, for both materials, conducting SMAT on the hardened material (SMAT after ageing) led to a 420 reduced samples surface roughness resulting in an increase of about 20 % of the fatigue limits.

421 Considering the non-negligible number of publications highlighting the fatigue life reduction after 422 severe SMAT-processing as the consequence of the increased roughness, it would be tempting to attribute the 423 reduction of the fatigue limit observed in this work at R_{TT} to the formation of dimples due to the high number 424 of impacts. However, as shown in Figure 2, such initiation mechanism on dimples was only detected for the 425 SMATed samples loaded at R_{TT} under low stress amplitudes. Instead, for the other R_{TT} conditions, crack 426 initiation mainly occurred around inclusions that were already present in the raw material while no obvious 427 difference was found in the initiation sites comparing polished and SMATed samples for the R_{TC} conditions. 428 However, it is clear that the "quality" of the surface has an effect on the fatigue properties. Indeed, for the 429 SMAT samples that were mirror polished and tested at $R_{\rm TT}$, the properties were restored to those of the initially 430 polished material (Figure 1 b). Thus, the action of smoothing the surface after SMAT had a positive effect on 431 the fatigue properties. From this point of view, Bagherifard et al. also observed an additional (+10 %) increase 432 in the fatigue limit of severely shot peened low-alloy steel samples after conducting a grinding 433 post-processing [59]. An additional confirmation of the importance of the surface state in initiating the fracture 434 is the absence of any surface crack on the run-out samples (Figure 5). However, our overall results clearly 435 show that, even if the roughness of the surface and the presence of stress raisers must have an effect on the 436 fatigue properties, it is not the primary reason for the difference in behavior observed between the $R_{\rm TC}$ and $R_{\rm TT}$ 437 loading conditions.

438

4.2. Effects of the cyclic stress on the plastic deformation mechanisms

Exploring alternative hypotheses for the variable effectiveness of SMAT to enhance the fatigue limit 439 440 at $R_{\rm TC}$ and $R_{\rm TT}$, the effect of the different type of loading has to be discussed. Indeed, the fact that half of the 441 fatigue life of the samples tested at R_{TC} occurs under a compressive state and that the maximum stress during 442 cycle is higher for R_{TT} imply very different macro- and micro- strains that may lead to the occurrence of 443 different damage mechanisms. The difference in material response to the different loading conditions can be 444 witnessed by two major observations. The first one is the self-heating of the samples observed at R_{TC} for a 445 10 Hz frequency but not at $R_{\rm TT}$. The self-heating at $R_{\rm TC}$ is not detailed in the present paper but is completely 446 identical to the one observed on the 304L stainless steel studied in a previous work [40] in which the same 447 fatigue behavior at $R_{\rm TC}$ was obtained (Figure 1 a). In contrast, no self-heating was recorded at $R_{\rm TT}$. To observe 448 such self-heating, a much higher loading frequency should probably be used. Indeed, Tian et al. also measured 449 a self-heating on a 316LN loaded at R = -1 at 10 Hz but had to increase the testing frequency to 70 Hz to start 450 observing relevant self-heating at R = +0.1 [60]. The second evidence is the difference in microstructure 451 evolution in the core of the sample between the $R_{\rm TC}$ and $R_{\rm TT}$ loading conditions. Indeed, during the tension-452 compression cyclic loading of the SMATed samples, the martensitic transformation occurred at depth below 453 200 µm (Figure 6 a). Interestingly, this transformation did not occur in the regions closer to the surface at 454 100-200 µm, probably due to the too high density of dislocations already present in the SMATed highly 455 hardened domain that prevented the austenite to martensite phase transformation. This is in good accordance 456 with the literature about the conditions required for athermal martensitic transformation in steels. Indeed, it 457 was shown that enhancing the yield stress of the parent phase (austenite) by a moderate increase of the 458 dislocation density [61, 62] or decreasing the grain size [63] both lead to a reduction in the $M_{\rm S}$ temperature. 459 Such lowered value of $M_{\rm S}$ implies that a greater mechanical work is required to allow martensitic 460 transformation; in other words, a greater stability of the austenite in this region. In contrast, neither stress- nor 461 strain-assisted martensitic transformation was triggered under the tension-tension fatigue condition used in this 462 work. Due to the negative values of $M_{\rm S}$ and $M_{\rm d30}$ for this alloy (Table 1), a significant strain is required to 463 induce $\gamma \to \alpha'$ phase transformation, as it was the case in the first 100 µm due to the severe plastic deformation 464 induced by SMAT. The fact that phase transformation occurred during cyclic loading under R_{TC} and not under 465 $R_{\rm TT}$ conditions seems rather counter intuitive. Indeed, it is widely admitted that the martensite volume increases 466 with strain [61, 64] and such phenomena is considered in recent modelling [65]. Also, Lebedev et al. showed 467 that for a 18Cr-10Ni stainless steel, the martensitic transformation is more advanced when the sample is 468 subjected to tension than compression [66]. Then, considering that the samples are subjected to higher strains 469 for R_{TT} than R_{TC} and are exclusively undergoing tensile loading at R_{TT} , it would be fair to expect a higher 470 volume fraction of martensite for $R_{\rm TT}$ than $R_{\rm TC}$. The exact opposite is observed here, indicating that other 471 parameters are affecting the stability of austenite when subjected to cyclic loading. Lebedev et al. also reported 472 that it is unusual to find martensite in fatigue loaded samples except on the crack path where the stress triaxiality 473 is much higher than in the rest of the samples [66] which might be an essential aspect to consider in order to 474 understand the observed phase transformation.

475 Nevertheless, even if these differences in the plastic deformation mechanisms and resulting 476 microstructures were observed due to the different loading conditions, they are not at the origin of the 477 differences in the fatigue properties. Indeed, the observed phase transformation is supposed to have negligible 478 influence on the surface fatigue crack initiation because it occurred 200 µm below the surface. In addition, in 479 Figure 1 the 304L and the 316L stainless steels were shown to have identical fatigue behavior at R_{TC} even if 480 Dureau et al. highlighted that the surface of the 304L was fully martensitic after SMAT [40]. Moreover, the 481 analysis of hardness gradient after different fatigue loading conditions revealed limited changes in the 482 hardening state in the SMAT affected region for R_{TC} and R_{TT} conditions. As the evolutions of hardness followed 483 the same trends, it should have an equally beneficial contribution to the fatigue life enhancement for both load 484 ratios after SMAT and, consequently, it is also not responsible for the fatigue life reduction at $R_{\rm TT}$.

485 One of the major difference revealed by the analysis of the run-out samples is that the residual stress 486 gradient evolutions after fatigue were drastically different under R_{TC} and R_{TT} . The compressive stress state 487 relaxation observed for R_{TC} was commonly observed after fatigue loading for many materials and testing 488 conditions [4, 5, 19, 31, 40, 51] and sometimes even led to a fully relaxed stress field [34]. Comparatively, the 489 gradient reversion leading to the formation of tensile residual stress gradient generally occurs for quasi-static 490 loading or when overloads occur during fatigue [35, 67-71]. In our case, the inverted residual stress state was 491 formed after the first cycle of fatigue and remained during the whole fatigue life of the samples. As the internal 492 stresses redistributions are essentially a matter of mechanical equilibrium between the sub-surface layer 493 affected by the SMAT and the core, the maximum applied stress during cyclic loading governs the residual 494 stress evolutions. For the studied samples, the maximum applied loads during cycles were respectively $\sigma_{R_{TC} \max}^{SMAT} = 240$ MPa and $\sigma_{R_{TT} \max}^{SMAT} = 477$ MPa. $\sigma_{R_{TC} \max}^{SMAT}$ is then equal to 35% of the maximum tensile stress 495 σ_{UTS} (=650 MPa), and 110 % of the yield stress σ_{ys} (=220 MPa). On the other hand, $\sigma_{R_{\text{TT}}\,\text{max}}^{\text{SMAT}}$ is equal to 72% 496 of σ_{UTS} thus 215 % of σ_{ys} . As $\sigma_{R_{\text{TC}} \text{ max}}^{\text{SMAT}}$ was close to the yield stress, the core of the samples experienced small 497 strains at R_{TC} , leading to moderate residual stress relaxations. In contrast, $\sigma_{R_{TT}}^{SMAT}$ being twice the yield stress, 498 499 the applied tensile strain has reverted the distribution of residual stresses. In both cases, those redistributed and 500 stabilized residual stresses act as a local mean stress superimposed to the macroscopic loading, thus changing 501 the stress field during cycling. This can help to understand the differences in the observed fatigue crack 502 initiation mechanisms.

503 For the R_{TC} conditions, the maximum applied loads are just a few tens of MPa above the material 504 elastic limit, causing only moderate and localized plastic strain. As confirmed by the surface analysis in 505 Figure 2, these levels of plasticity are not large enough under the $R_{\rm TC}$ conditions to initiate cracking on 506 inclusions regardless of the stress amplitude, especially for the SMAT samples due to the surface compressive 507 residual stress. If Kakiuchi et al. observed one initiation on such calcium-aluminate inclusion in a 304L 508 stainless steel treated by SMAT and then loaded at R = -1 under rotating bending [72], it should be noticed that 509 their fatigue tests were conducted at a temperature of 300 °C. It is possible that this elevated temperature can 510 facilitate the use of these particles as stress raisers because it enhances the differences in mechanical properties 511 between the austenitic matrix and the hard inclusion.

512 For the R_{TT} conditions, the much higher applied stress than for R_{TC} led to macroscopic straining of the 513 samples highlighted by the high density of slip bands in Figure 2, thus damaging the inclusions interface with 514 the steel matrix. This allowed the inclusions to be activated as initiation sites. This is particularly true for the 515 SMATed and SMAT + polished samples due to the superimposed macroscopic tensile loading and surface high
516 tensile residual stress states.

517 Finally, as the surface inspection of the run-out samples never revealed non-propagating fatigue 518 cracks, the resistance of the material to fatigue crack initiation was shown to rule the fatigue limit in the high 519 cycle fatigue regime. Then, the redistributed residual stress fields were taken into account into a Crossland 520 fatigue initiation criterion and provided a good description of both the effect of load ratio and SMAT. The 521 residual stress state was then considered the first-order parameter governing the fatigue performance of 522 SMATed austenitic stainless steels samples. Therefore, the development of numerical models to predict the 523 cyclic response of SMATed samples and the residual stress field evolutions after such cyclic loading seems to 524 be a very interesting way of predicting the fatigue life of gradient microstructure stainless steels components 525 under various loading conditions.

526 <u>6. Conclusions</u>

527 Uniaxial fatigue tests have been conducted on an austenitic stainless steel with load ratios $R_{TC} = -1$ and 528 $R_{TT} = +0.1$, to evaluate the high cycle fatigue performance of the gradient microstructure generated by SMAT 529 in a context of local or general plasticity. The fatigue limit change resulting from SMAT was beneficial 530 at $R_{TC} = -1$ (+17 %) and detrimental at $R_{TT} = +0.1$ (-7 %).

After fatigue loading, no cracks were detected at the surface of any of the run-out samples, indicating that the resistance to fatigue crack initiation was governing the fatigue limit in the HCF regime. Processing inclusions already present in the initial material were the predominant initiation sites for crack nucleation for R_{TT} rather that surface defects. The evolution of residual stresses was proved to be the primary parameter governing the fatigue response: the compressive residual stress generated by SMAT was partially relaxed at R_{TC} while it was fully reverted after only one cycle of R_{TT} loading.

The beneficial effect on the fatigue limit (improved by +17 %) for R_{TC} = -1 loading is due to the sub-surface compressive residual stresses present in the hardened gradient microstructure that overcome the negative effect of the surface defects induced by SMAT. During fatigue loading at R_{TC} , self-heating of all the specimens occurred together with sub-surface martensitic transformation for the SMATed samples. Nevertheless, these two modifications had a negligible impact on the fatigue properties. The detrimental effect on the fatigue limit (reduced by -7 %) for R_{TT} = +0.1 is associated to the high imposed macroscopic stresses that caused a complete reversion of the compressive residual stress gradient towards tension. This very high tensile stress leads to faster fatigue crack initiation compared to the untreated material. In this case, no self-heating nor sub-surface martensitic transformation were revealed, but the much higher stress at R_{TT} was proposed to be responsible for the initiations on inclusions. In this case, the combined negative effects of tensile residual stresses and surface roughness leads to a slightly reduced fatigue limit for SMAT specimens.

For the $R_{\rm TT}$ loading condition, a mirror polishing procedure after SMAT allowed to restore the fatigue limit (same as the untreated condition) but did not improve it. This is a confirmation that (i) the redistribution of residual stress was of primary importance for controlling the high cycle fatigue behavior and (ii) the surface defects created by SMAT were not responsible for the incapacity of the SMAT to improve the fatigue limit at $R_{\rm TT}$.

554

555 <u>Acknowledgement</u>

- 556 This work was supported by the French State through the program "Investment in the future" operated by the
- 557 National Research Agency (ANR), referenced by ANR-11-LABX-0008-01 (Labex DAMAS).
- 558 This work was also supported by the CPER FEDER project of Région Nouvelle Aquitaine.

559 *Declaration of competing interest*

The authors declare that they have no known competing financial interests or personal relationships that couldhave appeared to influence the work reported in this paper.

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	С	Mn	Si	S	Р	Cr	Ni	Мо	Cu	Ν	Fe	$M_{\rm S} [^{\circ} \rm C]$	<i>M</i> _{d30} [° C]
Wt [%]	0.029	1.499	0.341	0.03	0.22	16.605	10.370	2.036	0.127	0.047	Balance	-210	-1

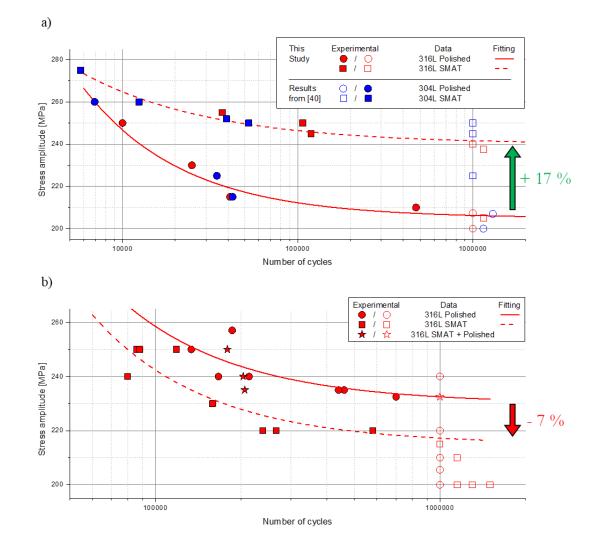
Table 1 Chemical composition of the studied 316L stainless steel

a)

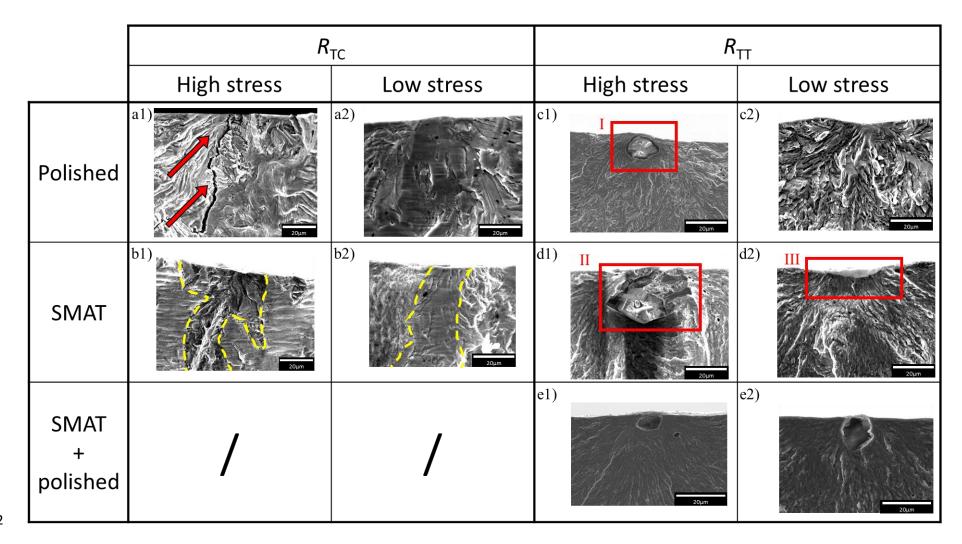
Testing condition name	Step 1	Step 2	Step 3	Step 4	Step 5	Ra [µm]
Initially polished		Machined	Mirror polished	/	/	$0.017 \pm 2.5.10^{-3}$
SMAT	Heat treated 900 °C 40 min			SMATed	/	1.621 ± 0.23
SMAT + polished					Mirror polished	$0.018 \pm 1.4.10^{-3}$

b)

Analysed unbroken sample	Load ratio	Stress amplitude [MPa]	Number of cycles	
SMAT	/	/	/	
SMAT + Fat $R_{\rm TC}$	-1	240	2.106	
SMAT + Fat $R_{\rm TT}$	+0.1	215	4.106	
SMAT + 1Cy $R_{\rm TT}$	+0.1	215	1	



780 Figure 1 Wöhler curves of a) polished and SMAT 316L and 304L samples loaded at R_{TC} =-1 and b) polished, SMAT and SMAT + polished 316L samples loaded at



782

783 Figure 2 SEM images of the characteristic initiation sites for a) R_{TC} polished, b) R_{TC} SMAT, c) R_{TT} polished, d) R_{TT} SMAT and e) R_{TT} SMAT + polished samples for

respectively 1) high and 2) low stress amplitudes

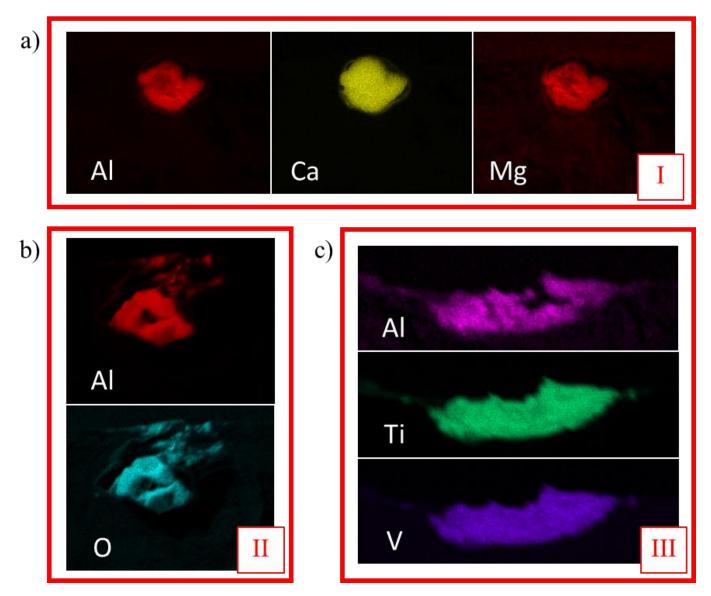
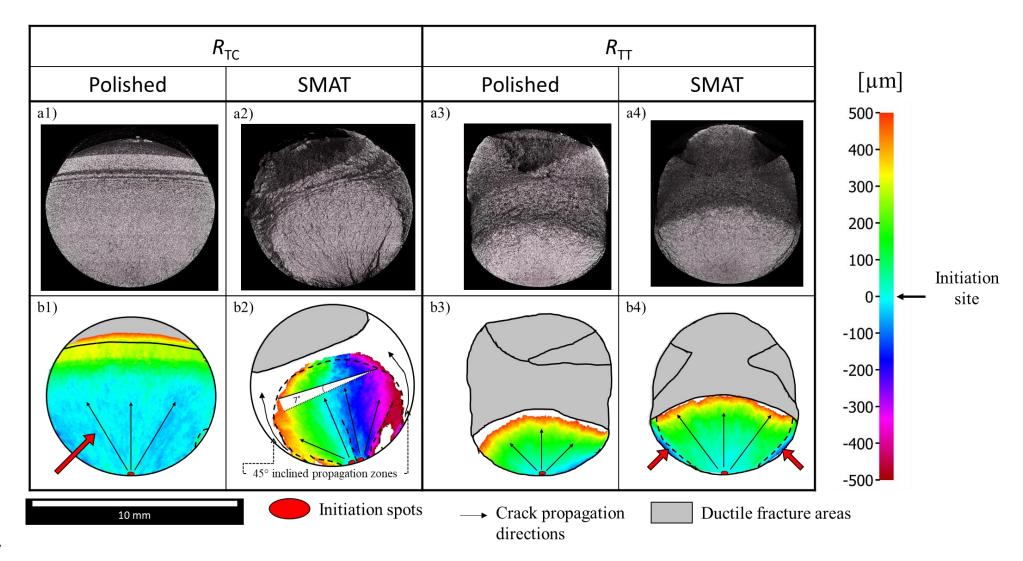


Figure 3 EDX maps corresponding to the regions highlighted in red in Figure 2



788 Figure 4 Observations of fracture surfaces a) optical microscope pictures and b) sketches with topology mapping of 1) R_{TC} polished 210 MPa, 2) R_{TC} SMAT 245 MPa,

3) R_{TT} polished 232.5 MPa and 4) R_{TT} SMAT 220 MPa samples

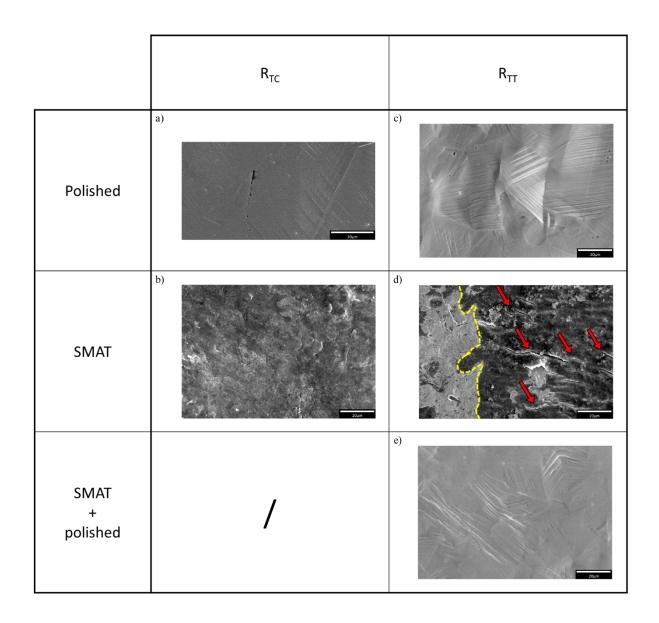
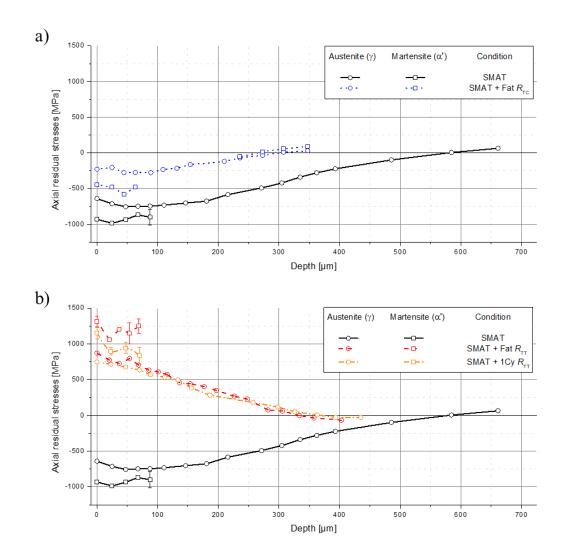
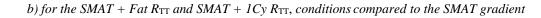
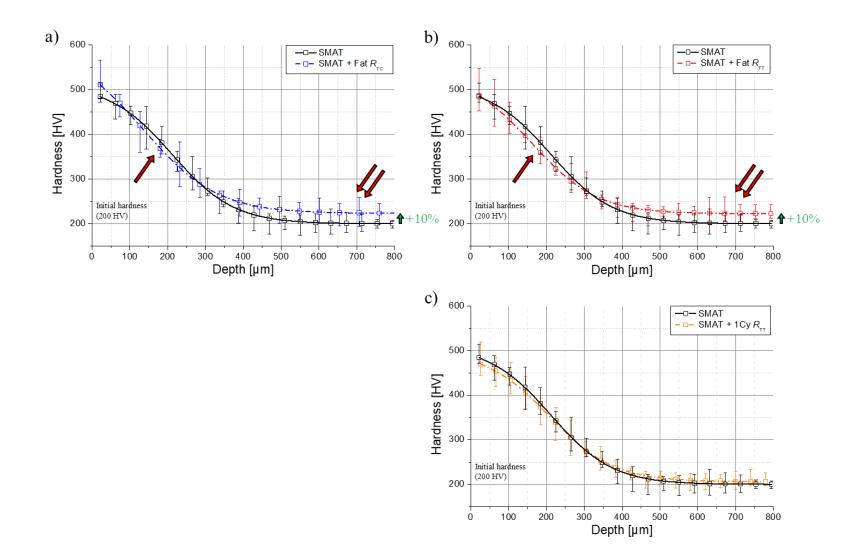


Figure 5 SEM images of the surface of a) R_{TC} polished, b) R_{TC} SMAT, c) R_{TT} polished, d) R_{TT} SMAT and e) R_{TT} SMAT + polished run-out samples



793 Figure 6 Evolution of the residual stresses in both austenite and martensite as a function of the depth below the treated surface a) for the SMAT + Fat R_{TC} condition and





795

796 Figure 7 Evolution of hardness as a function of the depth below the treated surface after a) SMAT + Fat R_{TC}, b) SMAT + Fat R_{TT} and c) SMAT + 1Cy R_{TT}, compared to

the SMAT gradient

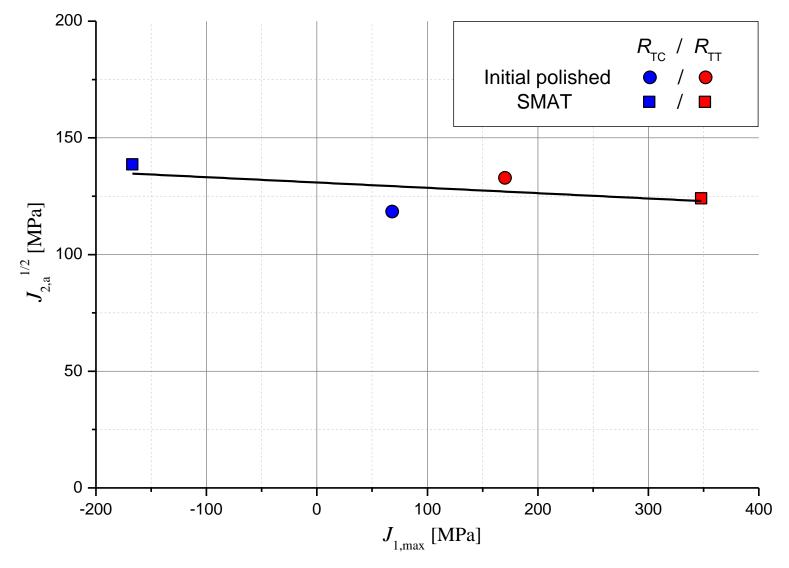


Figure 8 Crossland criterion established for all the tested load ratios and surface states