

Article

Effect of Multi-Step Austempering Treatment on the Microstructure and Mechanical Properties of a High Silicon Carbide-Free Bainitic Steel with Bimodal Bainite Distribution

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Abstract: The effect of multi-step austempering treatments on the microstructure and mechanical properties of a novel medium carbon high silicon carbide-free bainitic steel was studied. Five different isothermal treatment processes were selected, including single-step isothermal treatments above martensite start temperature (at 350 °C and 370 °C, respectively), and three kinds of two-step routes (370 °C + 300 °C, 370 °C + 250 °C, and 350 °C + 250 °C). In comparison with single-step austempering treatment adopting a two-step process, a microstructure with a bimodal-size distribution of bainitic ferrite and without martensite was obtained. Bainitic transformation was studied using dilatometry both for single-step and two-step routes and the specimens were completely characterised by electron microscopy (SEM and TEM), X-ray diffraction (XRD) and standard tensile tests. The mechanical response of the samples subjected to two-step routes was superior to those treated at a single temperature.

Keywords: carbide-free bainite; high silicon steel; bimodal distribution; multi-step austempering; heat treatments



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1. Introduction

Steels with nanostructured bainite are object of extensive research due to their excellent combination of high strength and ductility [1–5]. Their ability to achieve tensile strength largely above 1 GPa with total elongation above 15% is given from their unique multiphase microstructure, which includes bainitic ferrite and carbon enriched austenite [6–8]. This composite microstructure consisting in ultrafine constituents resulting from the diffusionless transformations is the responsible of high strength, since cracks need to traverse interphase but also different crystal structures, as demonstrated several times by Bhadeshia and Edmonds [9].

These steel grades are characterised by a significant amount of silicon, which inhibits cementite precipitation, that deplete mechanical properties [9], favouring austenite carbon enrichment during isothermal treatments [10–12]. During isothermal treatments, when bainitic formation occurs, carbon diffuses from supersaturated ferrite into austenite, increasing austenite stability, therefore, part of it is retained at room temperature.

When austenite is enriched in carbon, its M_s (martensite start) temperature decreases, avoiding martensitic transformation during final cooling and giving the possibility of further bainitic transformation at lower temperatures [1,13–15]. Bainitic microstructures with carbon enriched austenite are generally produced by isothermal treatments, after complete austenitisation, in the temperature range between B_s (bainite start) and M_s [16,17] or below M_s [18,19]. Discussing the morphology of retained austenite it is possible to distinguish between two different forms generated from prior austenite partitioning (i) austenite films

confined by ferritic plates and (ii) austenite blocks, located between bainitic ferrite sheaves and at prior grain boundaries, respectively. [1,16,20–24].

Blocks of austenite, due to the lower carbon content in comparison with films [16,25], have poor stability and a high tendency to undergo martensitic transformation during applied stress, leading to large brittle martensitic islands, thus decreasing material toughness [26]. On the contrary, the presence of austenitic films generally leads to an enhancement in material properties [27–31].

Hence, avoiding the formation of hard brittle martensite and blocky austenite islands is the primary design goal of novel bainitic steels. This could be completed by conducting bainitic transformation at the lowest possible temperature, refining the microstructural constituents [32], or varying the carbon content of the alloys and, as a consequence, Ms temperature. In fact, no thermodynamic limit characterises bainitic transformation and it can be conducted also close to room temperature, but, as drawback, there is a dramatic increase in the time for achieving completion [33]. According to many authors a multi-step austempering bainitic treatment approach, called (MBAT), could represent an innovative solution, as proposed in literature [25,26,34–36]. The MBAT process consists of austenitisation, and two/three isothermal holding periods at different temperatures for bainitic transformation.

High temperature austempering induces bainitic transformation with high kinetics and formation of coarser plates of bainitic ferrite while holding period at lower period allows austenite decomposition in finer bainitic ferrite plates, leading to the formation of microstructure with bimodal size distribution of bainite as shown by Hase and Bhadeshia [1,37], and strong removal of austenite blocks that can deplete the mechanical response, as observed by Avishan [33]. In addition, according to the authors, from mechanical point of view, the application of MBAT process leads to a response enhancement, both in terms of ductility and toughness, that are doubled, respect to the structure produced with single isothermal holding treatments.

The process design of MBAT is also still debated in the literature, from the point of view of the heat-treatment design. Such MBAT treatments were proposed in different variants: (i) by Hase and co-workers and Avishan et al. [33,37], who performed the change in temperature when the transformation at the first step was completed, (ii) while Soliman and co-workers interrupted the transformation at 50% the first step [38].

An inverted MBAT process called IBAT was proposed by Gao and co-workers [25]. In their work, the first step of the process was performed at a lower temperature, and then the material was heated to a higher temperature for the transformation completion IBAT process, which, according to the author, allowed higher microstructural refinement in comparison with standard MBAT.

The aim of the present study was to investigate the details of transformation, microstructures and tensile properties of multi-step ausformed newly developed medium carbon, with 3 wt.% Si carbide-free bainitic steel.

2. Materials and Methods

Alloy Production and Heat Treatments

A 40 kg ingot was produced through vacuum induction melting, heated at 1200 °C and hot rolled down to 20 mm in seven passes, followed by air-cooling to room temperature. The chemical composition was measured with optical emission spectroscopy (OES) and it is reported in Table 1.

Table 1. Chemical composition (wt.%) of the Investigated alloy.

Fe	C	Si	Mn	Al	Cr	Ni	Mo	Cu	Ti	V	P	S
Bal.	0.38	3.20	2.60	0.10	0.05	0.05	0.02	0.05	0.001	0.006	0.008	0.007

ThermoCalc 2021 software (Thermo-Calc Software, Solna, Sweden) (with TCFE10 database) simulations were performed to calculate critical transformation temperatures, both for austenitisation and martensitic transformation (A_1 , A_3 , M_s) to design the heat treatments. Bainite start temperature (B_s) was calculated by means of the equation proposed by Zhao et al. [39]. Results are presented further in the manuscript.

Samples for dilatometry, with lengths 10 mm and 4 mm diameter, were machined employing electro-discharge machining from the steel plate in the direction perpendicular to the rolling direction. Dilatometric studies related to bainitic transformation were performed with a TA instrument quenching dilatometer DIL805A/D (TA Instrument, Hüllhorst, Germany). Based on ThermoCalc simulations a thermal cycle was performed for measuring the real A_{c1} , A_{c3} , and M_s of the investigated alloy. Specimen were heated at $10\text{ }^{\circ}\text{C/s}$ at $900\text{ }^{\circ}\text{C}$ and held for 5 min under vacuum (10^{-4} mbar) and cooled to room temperature in argon at $10\text{ }^{\circ}\text{C/s}$.

Three different multi-step treatment routes were selected in this research and are shown in Figure 1. The first route, called MULTISTEP1, consists of austenitisation at $900\text{ }^{\circ}\text{C}$ for 5 min, followed by rapid cooling ($10\text{ }^{\circ}\text{C/s}$) to the first bainite formation step at $370\text{ }^{\circ}\text{C}$ until completion of bainitic reaction, and a second bainite formation step at $300\text{ }^{\circ}\text{C}$ for further bainitic transformation, followed by final cooling to room temperature. The second one, named MULTISTEP2, is characterised by a second austempering treatment at $250\text{ }^{\circ}\text{C}$. Finally, the so-called MULTISTEP3 consists of a first austempering treatment at $350\text{ }^{\circ}\text{C}$ till the cessation of bainitic transformation and the second step at $250\text{ }^{\circ}\text{C}$. Single-step treatment with isothermal holding at $350\text{ }^{\circ}\text{C}$ and $370\text{ }^{\circ}\text{C}$ were also performed to compare the results. The heating rate (HR) and cooling rate (CR) of all heat treatments were set to be $10\text{ }^{\circ}\text{C/s}$. All the parameters for the heat treatments are listed in Table 2.

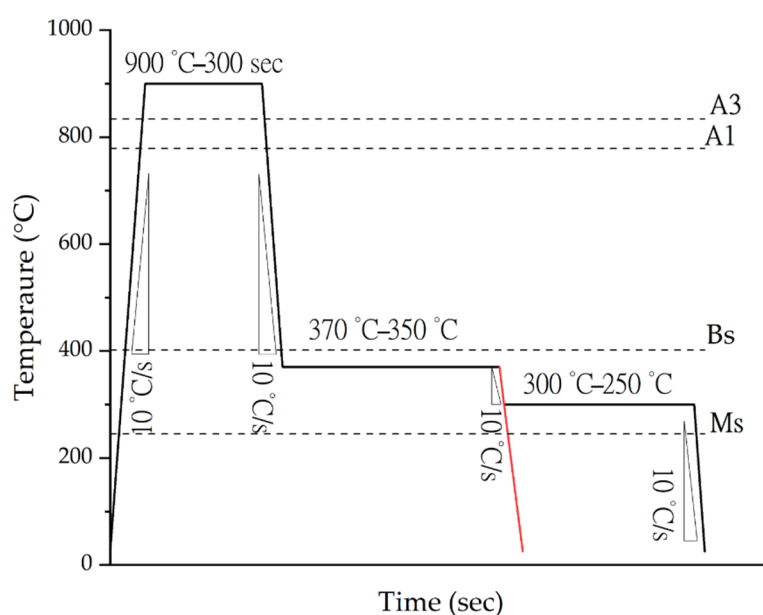


Figure 1. Bainitic multi-step heat treatment cycles after austenitisation at $900\text{ }^{\circ}\text{C}$ for 5 min.

Table 2. Resume of the performed heat treatment routes.

Sample ID	Austenitisation ($^{\circ}\text{C}/\text{time}$)	1st Isothermal Treatment ($^{\circ}\text{C}/\text{time}$)	2nd Isothermal Treatment ($^{\circ}\text{C}$)
MULTISTEP 1	900/300 s	370/2.5 h	300/2.5 h
MULTISTEP 2	900/300 s	370/2.5 h	250/2.5 h
MULTISTEP 3	900/300 s	350/2.5 h	250/2.5 h
350 $^{\circ}\text{C}$ single step	900/300 s	350/2.5 h	-
370 $^{\circ}\text{C}$ single step	900/300 s	370/2.5 h	-

Microstructure characterisation was performed on heat-treated specimens following a standard metallographic practice. Final polishing was performed with diamond polycrystalline suspension of $\frac{1}{4}$ μm , followed by etching using a 5% Nital solution. Observation of the microstructure was performed with a Zeiss ULTRA55 Schottky field emission scanning electron microscope (Carl Zeiss NV, Zaventem, Belgium), operating at 15 kV and with an InLens detector. Thickness of bainitic ferrite was performed on SEM pictures taken at high magnification and analysed with the support of ImageJ software [40]. TEM observations, with a Philips CM30 (Philips NV, Amsterdam, The Netherlands) transmission electron microscope operating at 300 kV, were performed to resolve nanostructured microstructural features. Thin foil preparation involved mechanical thinning to 60 μm and twin-jet electropolished, with a Struers TENUPOL-3 (Struers S.A.S., Ballerup, Denmark), for the perforation of the 3 mm disk. The employed electrolyte consisted of 5% perchloric acid and 95% acetic acid [41–43]. X-ray diffraction measurements were carried out on polished samples to determine the volume fraction of retained austenite. Such characterisations were performed with a Bruker-D8 ADVANCE diffractometer (Bruker, Billerica, MA, USA), with a Cu $K\alpha$ radiation at a voltage of 30 kV and a current of 30 mA. Analysis was performed in the 2θ of 40° – 105° , with step scan 0.02° and acquisition time of 2 s. The volume fraction of retained austenite was calculated by performing Rietveld analysis with the support of Maud software (Luca Lutterotti, University of Trento, Trento, Italy) [44]. Carbon content (c_γ) in retained austenite was calculated according to Equation (1), as a function of austenite lattice parameter (a_γ), estimated from XRD Rietveld refinements [45]:

$$a_\gamma = 3.5467 + 0.0467c_\gamma \quad (1)$$

For tensile tests, flat samples with a gauge length of 25 mm, a width of 6.25 mm, and 3 mm thickness mm were prepared by water-jet cutting and milling, according to ASTM A370-20. At least three samples were tested for each condition to ensure reproducibility and the average value was presented. Heat treatments on dog-bone samples for the tensile tests were performed with a Fours H&C muffle furnace (Fours H&C, Fernelmont, Belgium) for austenitisation, and lead and tin-lead bath furnaces, with eutectic composition, were used for bainitic transformation steps. An MTS (MTS System Corporation, Eden Prairie, MN, USA) with a 50 kN load cell was used for tensile tests, the force was measured using the machine load cell, while displacement was measured using the crosshead movement. Finally, the yield strength was calculated with the 0.2% method.

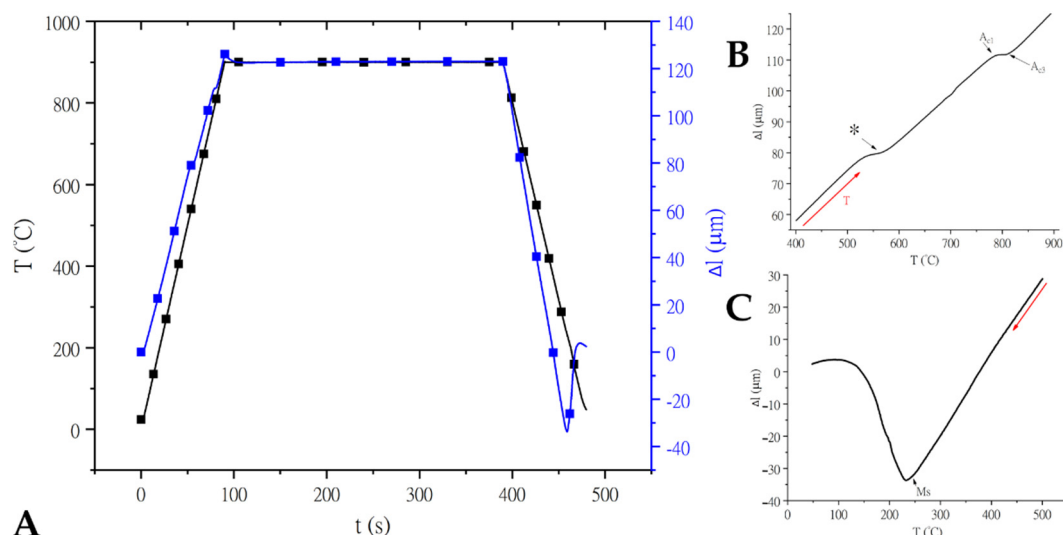
3. Results and Discussion

3.1. Thermocalc Analysis and Dilatometry

Calculated and measured critical transformation temperatures for the investigated alloy are reported in Table 3. Figure 2A shows the heat treatment cycle (black line) used for the estimation of the transformation temperature and the dilatation curve (blue line) obtained during the cycle. A linear contraction confirms that the chosen cooling rate is appropriate to prevent diffusional transformation. Figure 2B shows a detail of the heating branch of the change in length vs. temperature curve. Firstly, the heating is accompanied by a dilatation, with linear behaviour, connected to specimen thermal expansion, while at $\approx 520^\circ\text{C}$, a first contraction, indicated by a star in the figure, can be observed. This refers to the microstructural evolution of the material in the as-received state, which was characterised by martensitic microstructure, at those temperature martensite tempers and cementite precipitates. The hypothesis is supported by El-Fallah and Bhadeshia for analogous heat treatments of medium carbon nickel alloyed steel with starting martensitic microstructure [46]. Once cementite precipitation is completed, at around 580°C , change in length restarted following a linear pattern, due to linear expansion related to temperature increase. A second contraction was observed at 734°C (A1) due to austenite growth and furthermore, the data indicated also that at 839°C (corresponding to A3) the material is fully austenitic.

Table 3. Calculated (Thermocalc) and estimated (from dilatometry measured with spot welded K-thermocouple) transformation temperatures of the investigated alloys.

Calculation Method	A1 (°C)	A3 (°C)	Ms (°C)	Bs (°C)
Thermocalc	718	838	231	-
Dilatometry	779	834	245	-
Zhao et al. [39]	-	-	-	402

**Figure 2.** (A) Thermal cycle for the dilatometric study of critical temperature A_{c1} , A_{c3} , M_s (black line), and dilatation curve (blue line); (B) relative change in length against temperature curve indicating austenitisation transformation and the temperature A_{c1} and A_{c3} ; the star (*) = indicates cementite precipitation during heating of martensite due to carbon rejection; (C) cooling branch of the thermal cycle indicating M_s temperature and martensitic transformation.

Finally, Figure 2C shows the lowest part of the cooling branch of the dilatation experiment, in which martensite starts temperature (M_s) at 245 °C can be observed.

Dilatometric results for single-stage treatments are shown in Figure 3A,B. Figure 3A shows the cooling branch of the thermal cycle, showing length variation during the isothermal holding period due to bainitic reaction for a single step at 350 °C and 370 °C. In both cases, final cooling to room temperature is followed by martensitic transformation, at 122 °C and 226 °C, respectively. Variation in the M_s values between the two specimens and respect to the previously measured value, for the as-received material, is connected with austenite carbon enrichment occurring during bainitic transformation, as reported by Bhadeshia [47]. Furthermore, Chupatanakul and Nash [48] demonstrated, combining dilatometry and relationship between carbon content and M_s , that as austenite carbon concentration increases the M_s values of the untransformed austenite decreases. The decrease in M_s from 226 °C to 122 °C reducing the isothermal holding temperature from 370 °C and 350 °C is explained considering the T_0 curve theory and the thermodynamic of bainitic reaction [10,47]. As the temperature for bainite formation is reduced, higher is the maximum capability of austenite to accept carbon and, as consequence, lower M_s are achieved.

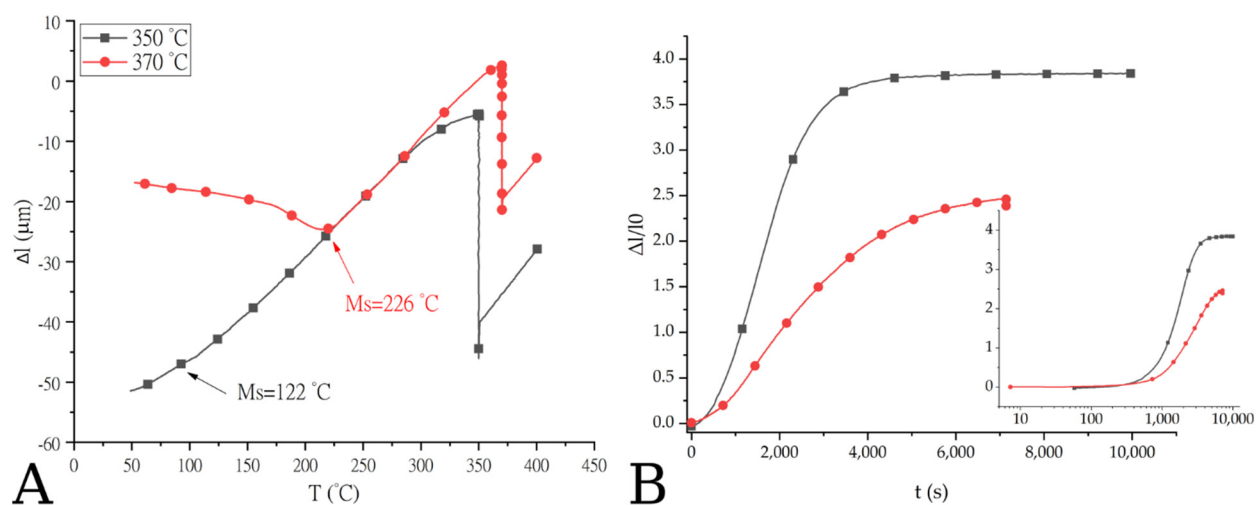


Figure 3. (A) Cooling branch of the dilatometric curve showing the change in length (Δl) against temperature for the single-step austempering treatment (A,B) Cooling branch of the dilatometric curve showing the change in length (Δl) against temperature for the single-step austempering treatment (B) dilatometric curve showing the change in length with time. The insert in (B) better shows the incubation time for both bainite transformations.

The relative changes in length during single-step treatment are reported in Figure 3B. The two curves indicate that lower temperature for the bainitic reaction leads to an increase in the extent of the bainitic reaction, as shown by the larger dilatation occurring during the isothermal treatment, as confirmed by Xu [49]. Furthermore, both for the isothermal treatment at 350 $^{\circ}\text{C}$ and 370 $^{\circ}\text{C}$, the transformation is characterised by an incubation time (160 s and 300 s, respectively) as it is displayed in Figure 3B. Bainitic reaction at 350 $^{\circ}\text{C}$ is characterised by a shorter incubation period thanks to an increase in the driving force for bainitic transformation, in agreement with Khare et al. [50] and Zhou and co-authors [51]. In addition, the kinetics of the transformation at 350 $^{\circ}\text{C}$ is faster than the transformation at a higher temperature, as indicated by the more rapid increase in change in length, in agreement with Xu and co-authors [49].

Changes in length vs. temperature curves for multi-step treatments are then depicted in Figure 4. As expected, during isothermal holding an increase in specimen length is observed due to bainitic transformation [26,48,52]. For all the multistep heat treatments, no martensite formation is detected while cooling to room temperature. This suggests that after the second stage bainitic transformation, the retained austenite was enriched in carbon, thus lowering the Ms below room temperature [48,52].

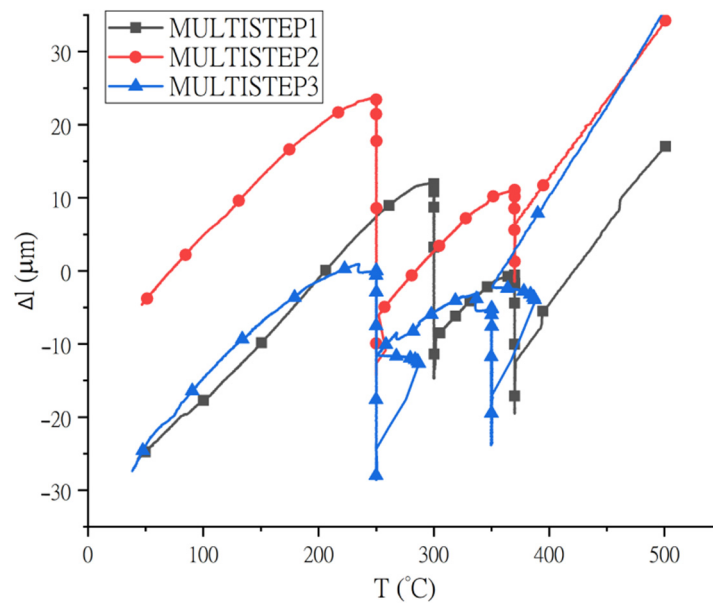


Figure 4. Cooling branch of the dilatometric curve showing the change in length (Δl) against temperature for the multi-step austempering treatments indicating the bainitic transformation during all the austempering steps.

Figure 5 shows the variation of the relative change in length as a function of the holding time for the different multi-stage treatments. All curves are characterised by the typical “S” shape of the isothermal solid-solid phase transformation curves in agreement with the curves proposed by Xu and San-Martin [49,53]. In the case of Figure 5A,C,E it is possible to observe for each step that the transformation starts slowly, and, in all the cases the incubation time that characterises the first stage of phase transformation is compatible with the one of the single-stage treatment. After that, a progressive increase in the transformation rate is observed, as testified by the derivative curves in Figure 5B,D,F. Comparing dilatometric curves of MULTISTEP1 and MULTISTEP2 routes, it is possible to observe that in the second stage of the treatment, performed, respectively, at 300 °C and 250 °C, more bainite is formed decreasing the temperature, thanks to the higher driving force [47]. Similarly, during the first stage of MULTISTEP3, the variation in specimen length is higher than the other two routes treated at higher temperatures in the first stage.

For both austempering stages in all multi-step heat treatments, the transformation rate increases until a “peak time transformation rate” (PTTR), followed by a decrease in the transformation rate [49,53,54].

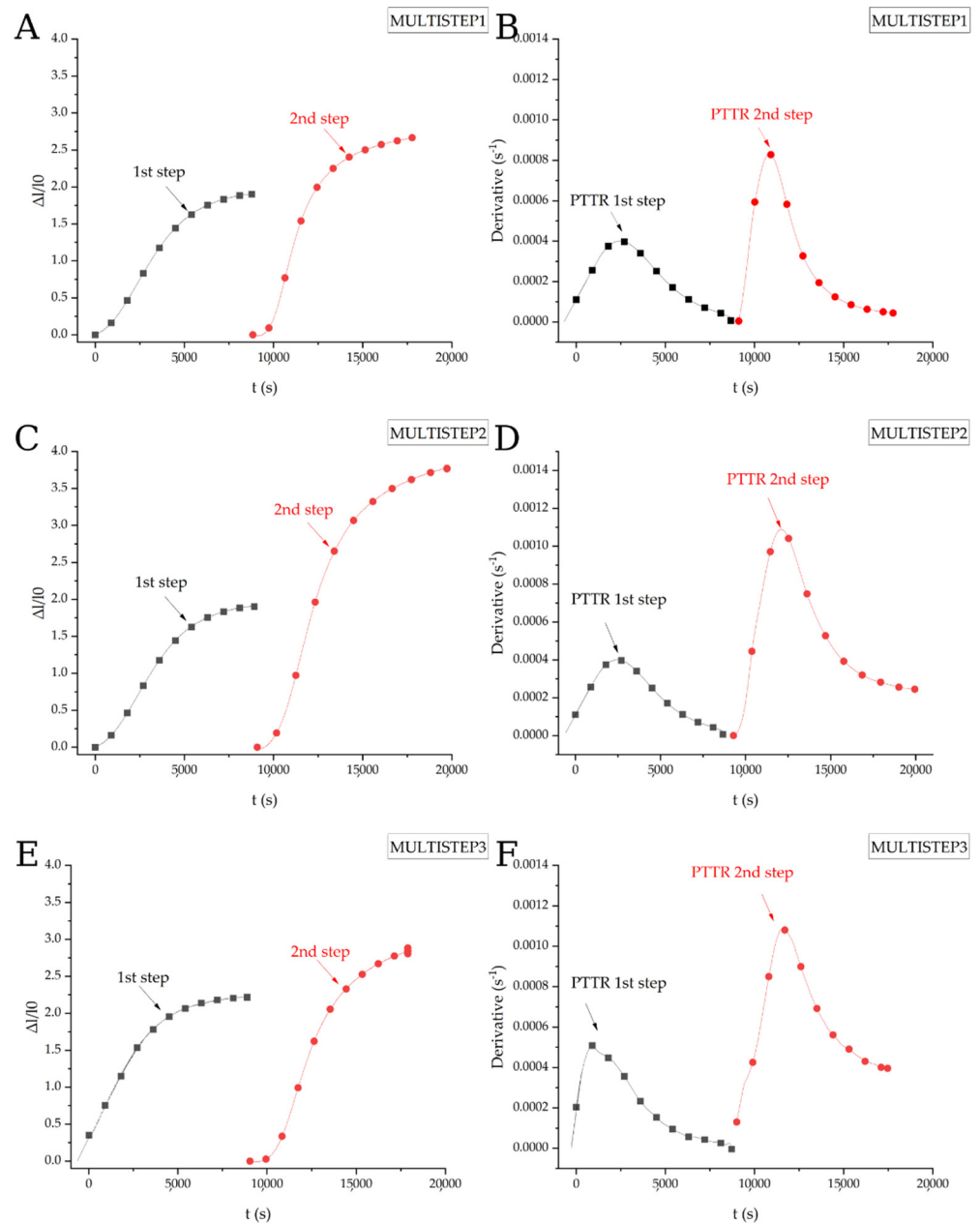


Figure 5. (A,C,E) relative change in length vs. austempering time curve for the heat treatment MULTISTEP 1, MULTISTEP 2, MULTISTEP 3, respectively. (B,D,F) Derivative of the relative change in length against time treatment describing the kinetic of the bainitic transformation during MULTISTEP 1, MULTISTEP 2, MULTISTEP 3 respectively.

3.2. Microstructural Characterization

Figure 6A,B depicts SEM micrographs of the specimens after the single-step austempering treatments. Figure 6A shows the microstructure after isothermal holding at 370 °C for 2.5 h, time at which bainite transformation is completed according to Figure 3B. Although in Figure 6B the product of the bainitic transformation at 350 °C is shown. Both images illustrate that the microstructure consists of bainitic ferrite plates (dark grey) separated by high carbon enriched austenite in form of films (R_{AF}) (light grey), as reported in the literature for similar steels [6,55–57]. Moreover, martensitic islands (M) and retained austenite blocks (R_{AB}) are present in correspondence with the prior austenite grain boundaries, according to Zhao [52]. The formers are the product of partial martensite transformation of retained austenite, as supported by the dilatometric curves shown in Figure 3. Concerning

the conventional single-step austempering process at 350 °C, the produced microstructures are also finer respect to the one formed at 370 °C, as demonstrated by the measurement of bainitic ferrite thickness, showed in Table 4. These results agree with those reported by other authors in the literature [37].

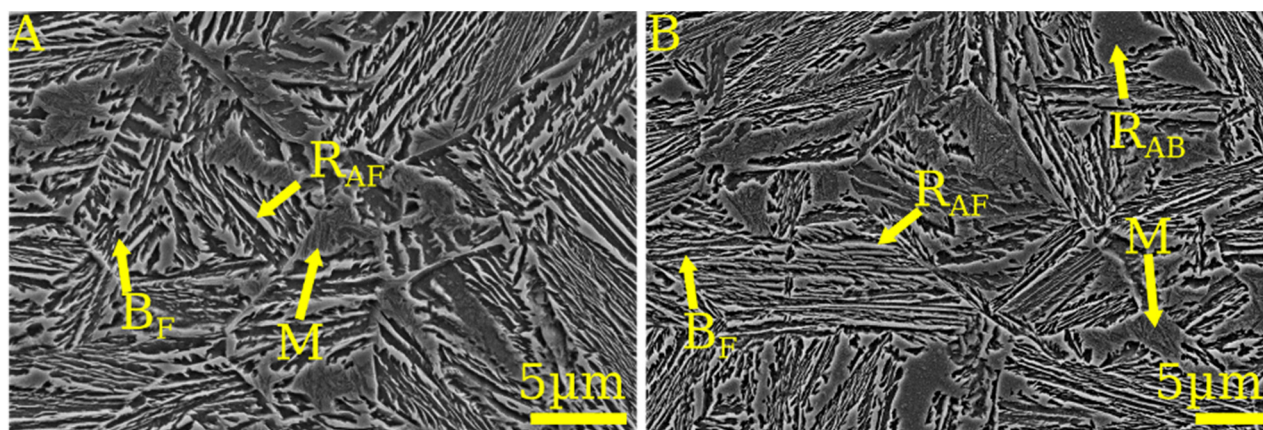


Figure 6. SEM micrograph of the sample treated with a single-step austempering until completion of the bainitic transformation at (A) 370 °C and (B) 350 °C. RAF: retained austenite film; BF: bainitic ferrite; RAB: retained austenite block; M: martensite.

Table 4. Resume on the measurements on the bainitic ferrite formed at the various isothermal holding temperature.

Temperature for Bainite Transformation (°C)	Bainitic Ferrite Thickness (nm)
370	345 ± 120
350	306 ± 65
300	146 ± 37
250	106 ± 33

Figure 7A reports the microstructure of the sample treated with the MULTISTEP 1 heat treatment route. A carbide-free bainitic microstructure is observed with two different generations of bainitic shaves which are formed during the two steps of the isothermal bainitic transformation, which are indicated with yellow dashed circles and B1 and B2 in the images. There is a first coarser generation (B1) formed during the holding period at 370 °C, while the second is finer (B2) and formed during isothermal holding at 300 °C, as shown in Table 4 from the measurement of bainitic ferrite thickness at various isothermal holding temperature, and in accordance with Bhadeshia [37,47]. Moreover, the microstructure displays the product of martensitic transformation, suggesting a beneficial effect of the second isothermal treatment on retained austenite stability. The results lead to confirm the previous hypothesis in the discussion of the dilatometric curves, where the deviations from linearity were related to martensitic formation after single-step heat treatments, during the final cooling. During the second isothermal stage, untransformed austenite transforms in bainitic ferrite leading to further carbon enrichment, an increase in the amount of thin films able to contain more carbon, and a reduction in the volume fraction of austenite blocks.

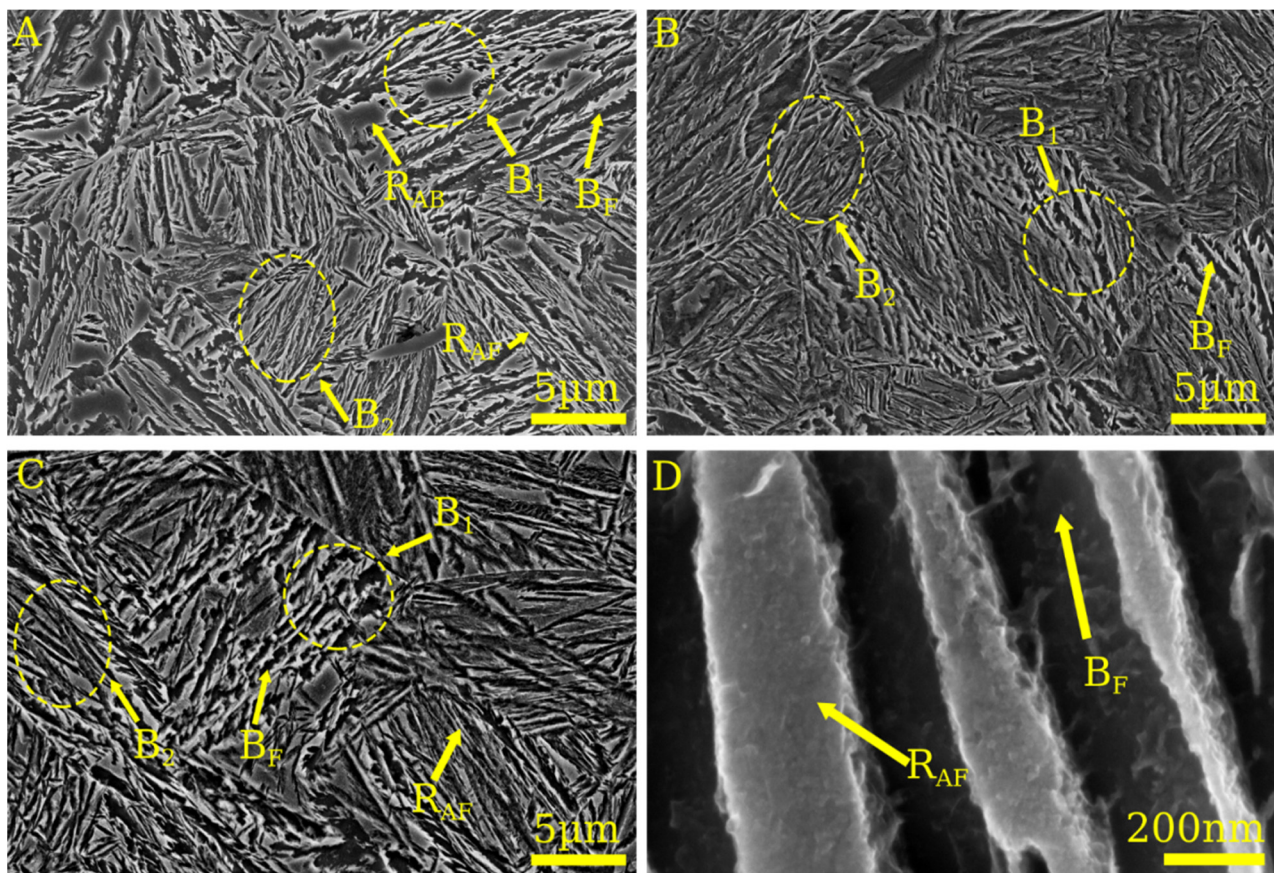


Figure 7. (A–C) SEM micrograph of the sample treated according to route MULTISTEP 1, MULTISTEP2, MULTISTEP3, respectively. (D) High magnification SEM micrograph showing film of retained austenite and bainitic ferrite. B1: carbide-free bainite formed during the first isothermal holding step; B2: carbide-free bainite formed during the second isothermal holding step. RAF: retained austenite film; BF: bainitic ferrite; RAB: retained austenite block; M: martensite.

Figure 7B shows the microstructural characterisation of MULTISTEP 2 which, as in the previous case, displays a microstructure consisting of two bainite generations. Finally, as for MULTISTEP 1, no martensite islands are present. A bimodal distribution on bainite is also observed after the MULTISTEP 3 route, as shown in Figure 7C. Thick plates of bainitic ferrite transform during the isothermal stage at 350 °C and the finer during the second stage at 250 °C, as reported in Table 4 as consequence of the thickness measurement of bainitic ferrite plates. One highlight of SEM observations is that the amount of blocky austenite is larger in MULTISTEP 1 than MULTISTEP 2 and MULTISTEP 3.

Finally, in Figure 7D is a high magnification SEM micrograph of bainitic sheaves of the sample treated according to MULTISTEP1, showing bainitic subunits with a thin films of retained austenite with thickness in nanoscale. Moreover, no carbides were observed thanks to the effect of silicon on suppression of cementite [58].

TEM analysis was performed on samples treated according to MULTISTEP3 route. Selected micrographs are shown in Figure 8. Bright-field image reported in Figure 8A depicts a shave of carbide-free bainite, with austenite films entrapped by bainitic ferrite sub-units, while in Figure 8B a block of retained austenite localized between different variants of bainite, with the typical polygonal shape, in 2D section [23].

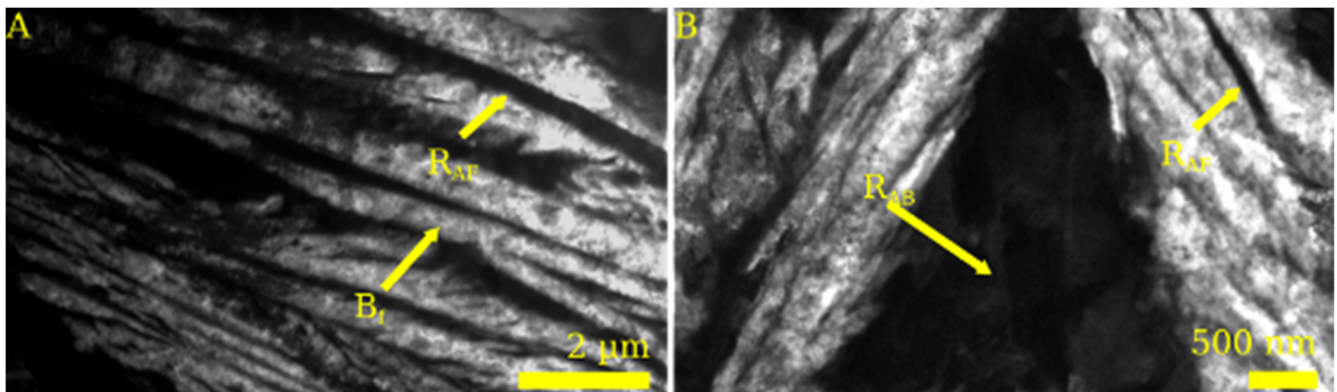


Figure 8. TEM micrographs on the samples treated according to MULTISTEP3 route (A) Bainitic sheave, (B) Retained austenite block between bainitic sheaves.

3.3. X-ray Diffraction

Figure 9 shows the diffraction patterns of all the specimens after the different treatment processes, both single-step and multistep. Then, the results of retained austenite quantification performed utilizing Rietveld analysis and evaluation of the carbon content are shown in Table 5. Peaks relative to BCC and FCC phases characterize all the patterns displayed in the figure.

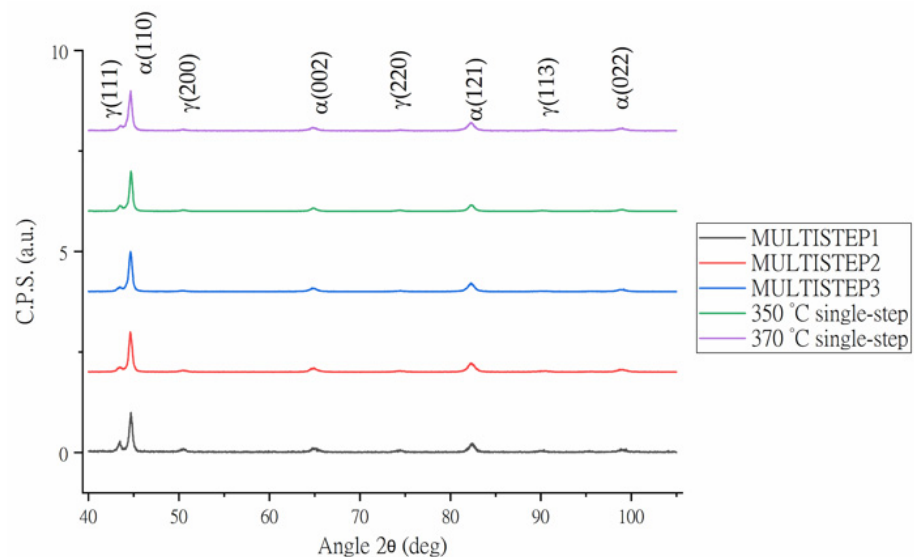


Figure 9. XRD patterns for the different heat treatment routes.

For the single-step treatment, it is possible to observe that a reduction in holding temperature, during the isothermal treatment, from 370 °C to 350 °C leads to an increase in the austenite final amount, once the material is cooled down to ambient temperature. This result is in contrast with some results present in the literature, for example by Son et al. and Zhao and co-authors [7,16], that observed a reduction in the final total amount, thanks to the higher driving force associated with the bainitic reaction that is reduced when the temperature for the bainitic reaction is closer to B_s [50]. However, the opposite behaviour is explained considering the lower austenite carbon enrichment (0.81% vs. 0.87%) that occurs at a higher temperature, shown in the table, and therefore a lower austenite stabilization. This explanation is confirmed by both dilatometry and SEM observations, where martensitic transformation at 226 °C for the specimen treated at 370 °C occurs and whose product is visible in the micrographs. In addition, results are supported also by the work published by Putatunda et al. [59], in the same range of temperature.

Two-step treatments are characterised by significant variations in the final content of retained austenite. In the case of MULTISTEP 1 and 2, the final austenite volume fraction is significantly higher than the one obtained with the single step at 370 °C. Such an increase from 12.9% to 23.4% and 17.97%, respectively, are associated with the double bainitic reaction mechanism. On the one hand, during a single-stage treatment at 370 °C the carbon enrichment and the extent of bainitic reaction is limited, as shown by dilatometry, i.e., retained austenite could not be stabilized at room temperature and undergo martensitic transformation during final cooling, as shown by SEM micrographs. On the other hand, performing a second step leads to a secondary bainitic reaction, which promotes a further carbon enrichment of surrounding austenite, leading to an increase in austenite stability, which is, therefore, retained at room temperature [16].

In the case of MULTISTEP3, a reduction in the final volume of austenite is observed thanks to the larger extent of the bainitic reaction occurring performing a two-stage treatment, in agreement with Avishan [33].

Comparing the different multi-step routes it is clear that depending on the temperature of the second isothermal stage the amount within the microstructure varies significantly. MULTISTEP2 leads to a decrease in austenite volume fraction with respect to MULTISTEP1 because of the second stage performed at a lower temperature that leads to larger decomposition of blocky austenite, in the second generation of bainite, at 250 °C respect to the one that occurred at 300 °C, as shown by Avishan and Mousalou [1,33].

In the case of multi-step treatment, a reduction in the value of the austenite carbon content (c_γ) presented in Table 5 is observed passing from MULTISTEP2 to MULTISTEP3 due to the higher extent of the bainitic reaction occurring at lower temperature and consequent larger carbon enrichment from supersaturated ferrite. However, the presented value are the results of the applied model, cited in the previous section, considering an average lattice parameter of austenite without the possibility to distinguish different morphology of retained austenite (blocks and films) and the two different bainite distributions. Different techniques, as neutron diffraction, can provide more attainable results [60].

Table 5. Variations in volume fraction of retained austenite and carbon concentration (c_γ) as a function of the austempering cycle of the investigated steel.

Treatment	V_γ (vol, %)	a_γ (Å)	c_γ (wt. %)
MULTISTEP 1	23.42 ± 0.95	3.611605	0.95
MULTISTEP 2	17.97 ± 0.47	3.607816	0.88
MULTISTEP 3	11.63 ± 0.42	3.610771	0.94
350 °C single step	15.64 ± 0.36	3.607232	0.87
370 °C single step	12.87 ± 0.50	3.604915	0.81

3.4. Mechanical Properties

Figure 10 shows the stress–strain engineering curves of the heat-treated samples, while results are resumed in Table 6. In the case of single isothermal treatment processes both yield strength and ultimate tensile strength decrease with the bainitic transformation temperature. This mechanical behaviour is consistent with the microstructural observation and results exposed in literature by many researchers, as Zhao and co-authors [61]. For lower isothermal holding conditions (350 °C) an improvement in terms of elongation to fracturing (from 0.028 to 0.066) was also observed, in agreement with Zhao [16]. Higher nucleation rate and yield strength of undercooled austenite at lower isothermal holding temperature produce finer bainitic ferrite plates, that combined with smaller and more stable blocks affects positively the mechanical response [16]. Moreover, in single-step processes continuous yielding is observable and the fracture is reached without necking and just showed uniform elongation, similarly with Avishan and co-authors [23]. As explained by the authors, the presence of austenite with low stability, transformation induced phenomena occur in the early stage of the tensile test. TRIP effect occurring at low strain level is, as matter of fact, detrimental.

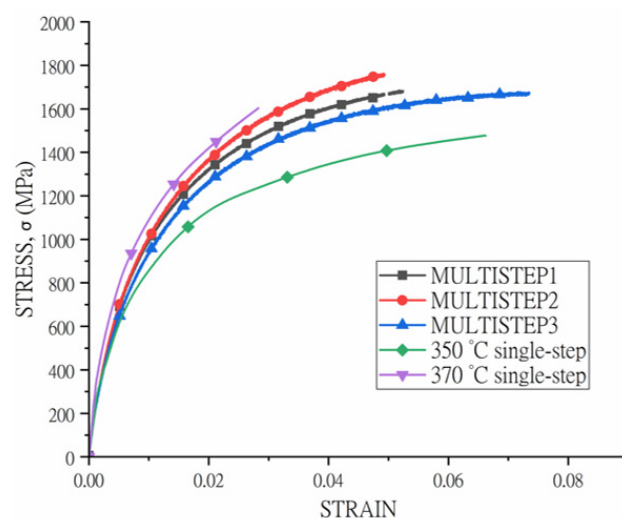


Figure 10. Engineering stress–engineering strain curves of the heat-treated steel samples (yield strength was calculated with the 0.2 criteria).

Table 6. Resume of mechanical properties of the heat-treated steel samples.

Treatment	σ_Y (MPa)	σ_{UTS} (MPa)	ϵ_f
MULTISTEP 1	950	1685	0.052
MULTISTEP 2	970	1761	0.049
MULTISTEP 3	857	1677	0.073
350 °C single-step	658	1477	0.066
370 °C single-step	1046	1603	0.028

Considering the multi-step routes, the results indicated that the treatments were successfully improving both ultimate tensile strength levels and elongations to fracture. In particular, MULTISTEP1 and MULTISTEP2 routes allow to improve fracture elongation from 0.028 to 0.052 and 0.049, respectively, and ultimate tensile strength from 1603 MPa to 1685 to 1761 MPa, respectively. Greater fracture elongation observed in the two-stage treatment, in comparison with the single-step treatment at 370 °C, is associated with lower content of untempered brittle martensite and lower amount and size of blocks of retained austenite [37]. The finer microstructure, deriving from the second isothermal stage, can compensate more efficiently, the damage occurring during stress application, giving as results better comprehensive properties, as proved also by [26]. Results agree with microstructural observation and X-ray diffraction and literature results proposed by Bhadeshia [37].

Moreover, both routes exhibit just uniform elongation without formation of neck and localized deformation before reaching fracture and without reduction in the sample area. Comparing MULTISTEP1 and 2, the improvement in terms of UTS is connected to the finer bainite during the second stage of the isothermal treatment in the case of MULTISTEP2. Concerning the MULTISTEP3 route greater values yielding point and UTS point are observable with respect to the single-stage treatment performed at 350 °C, while just a slight increase in total elongation characterises the two-step treated material. Higher yield strength is explained considering the increase in dislocation density and the carbon trapping in dislocation occurring during the two-step treatment [57].

3.5. Fracture Surface Analysis

Figure 11 shows SEM micrographs of the fracture surface for all the multistep treatments. It is evident that the fractures are mainly composed of facets, evidenced in the figure by the yellow dashed circles, and are present in all the specimens. The aspect of the fracture surface is clearly in agreement with the tensile test curves, characterized by

uniform elongation and the absence of necking and localised deformation, similarly to the results proposed by Avishan [23].

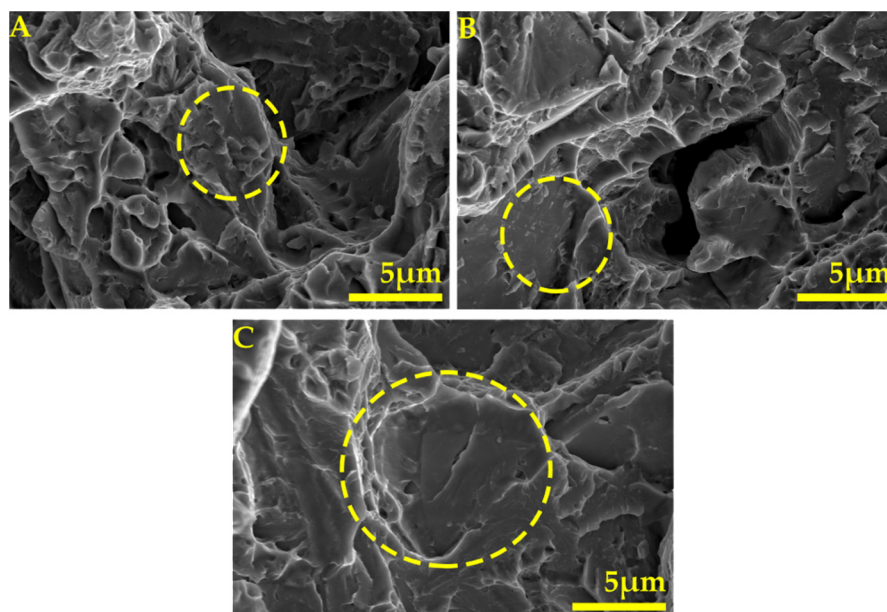


Figure 11. Fracture surfaces of the heat-treated steel samples after (A) MULTISTEP1, (B) MULTISTEP2, (C) MULTISTEP3.

However, the mechanical behaviour and the morphology of the fracture surfaces are in contrast with several works in literature where specimens showed necking. This can find an explanation considering the mechanical austenite stability, as discussed by Garcia-Mateo [6,62]. Very fine filmy retained austenite can rich very high stability and it is not possible to obtain full advantage of TRIP effect, due to the small number of potential nucleation sites for martensitic transformation and to the large carbon content and other elements such Mn and Si that enhance austenite mechanical stability. As a consequence of high stable austenite, therefore, cannot provide effective TRIP effect and necking formation.

4. Conclusions

Five different heat treatment routes have been performed in this research. In particular, two single-step isothermal treatments were performed above M_s temperature, at 350 °C and 370 °C, and three two-step austempering routes. The following main conclusions can be drawn from the results:

- (1) Single-step isothermal treatments leads to microstructure containing bainitic ferrite, films of stable austenite, unstable block and hard brittle martensite. Such combination of microstructural constituents is responsible of poor elongation due to unfavourable TRIP effect;
- (2) Two steps routes lead to the formation of a bimodal bainite distribution, coarser bainite formed at a higher temperature, and finer bainite formed during the second stage. Moreover, the second step of the bainite reaction avoids the presence of large martensitic islands in the final microstructure;
- (3) In addition, the second step of bainitic reaction leads to a further increase in austenite carbon content, improving the thermal stability of retained austenite;
- (4) Two-step treatments lead to improve mechanical properties in terms both of ductility (fracture elongation) and ultimate tensile strength.

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writing—review and editing, L.P., A.M.B.; visualization, M.F., A.M.B.; supervision, M.D., P.J.J.; project administration, M.D., P.J.J., funding acquisition, M.D. All authors have read and agreed to the published version of the manuscript.

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