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Original Article

Microstructural evolution and microhardness in a low carbon steel processed by high-pressure torsion☆

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ABSTRACT

A low-carbon triple-alloyed steel was processed by high-pressure torsion at room temperature for up to 5 turns under a pressure of 6.0 GPa. Microhardness, scanning electron microscopy and X-ray diffraction were used to investigate the hardness and microstructural evolution of the steel. Values of the Vickers microhardness were recorded across the sample diameters. The results show that there is a gradual evolution in both the hardness and the microstructure with increasing numbers of turns. However, the microhardness does not become fully homogeneous across the sample diameter after five turns and there remain significantly lower values in the center of the disk. These results indicate that complete homogeneity across the disks for this steel requires applied pressures higher than 6.0 GPa and/or torsional straining through more than 5 turns.

1. Introduction

The bulk properties of a material are of great importance for engineering applications. Comprehensive investigations on techniques for improving these properties are crucial, especially for materials for technological use such as with the iron-carbon system and its alloys [1]. Among these, the AISI 8620 low carbon – triple-alloyed steel is of great interest because of its use in manufacturing as a gear material and for hard-wearing machine parts when it is hardened and formed through carburizing or boronizing [2].

Improving the mechanical properties of this steel requires the use of techniques that modify its internal structure. Among these techniques, processing by high-pressure torsion (HPT) has attracted attention because of the capacity for achieving exceptional grain refinement, often to the nanometer level, and exceptionally high strength [3]. It was shown recently that HPT processing causes grain refinement and the decomposition of a supersaturated solid solution [1].

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The aim of this study was to investigate the microstructural evolution and the corresponding changes in microhardness of the AISI 8620 steel subjected to processing by HPT. This steel has fractions of ferrite and pearlite which are expected to deform during the processing operation.

2. Experimental materials and procedures

The material used in this work was a low carbon – triple-alloyed steel containing 0.2% C, 0.5% Cr, 0.6% Ni, 0.2% Mo, 0.8 Mo with the balance as Fe where the composition is expressed in wt.%. The as-received material in the normalized state was machined to cylindrical rods of 10 mm diameter and 50 mm length. HPT disks with thicknesses of ~1 mm were sliced from the rod and these disks were polished to final thicknesses of ~0.86 mm. The samples were heat treated at a temperature of 650 °C for 15 min in order to relieve the stresses introduced by machining. It was anticipated that this treatment would not produce any significant changes in the material structure or mechanical properties.

The HPT processing was conducted under quasi-constrained conditions [4,5] using a facility consisting of upper and lower anvils having central depressions with diameters of 10 mm and depths of 0.25 mm [6]. The processing was conducted at room temperature by rotating the lower anvil at a speed of 1 rpm under an applied load of ~470 kN, which is equivalent to an imposed pressure of $P = 6.0$ GPa. Separate disks were processed through totals of $N = 1/4$, 1, 2 and 5 turns. Two disks were prepared for each condition, for microstructure and microhardness measurements. The upper surface of each disk was marked immediately after HPT and prior to any microstructural analysis and hardness measurements.

Following HPT, X-ray diffraction patterns of the processed disks were recorded using an X'PertPro Panalytical diffractometer working with the following settings: θ–2θ varying from 10° to 90° with 0.02° step size, monochromatic CuKα radiation ($\lambda = 1.5409$ Å), 45 kV and 40 mA. For microhardness measurements, each processed disk was mounted and polished to have a mirror-like surface and measurements were taken at positions separated by 0.3 mm across the diameters of each disk with four individual points recorded around each selected position at distances of 0.15 mm. Full details of this procedure were given in an earlier report [7]. The microhardness measurements were taken using an Esseway model 600 hardness tester at a load of 50 kg-F and 20 s dwell time.

For microstructural observation using scanning electron microscopy (SEM), the disks were polished to have a mirror-like surface and then they were etched with nital solution. Images were taken using a JEOL model JSM 6490-LV SEM working at 10 kV near the mid-radius positions at distances of ~2 and ~4 mm from the center of each disk.

3. Results and discussion

Fig. 1 shows the X-ray diffraction patterns of the samples processed to different numbers of turns. Only the ferrite phase in orientations (1 1 0), (2 0 0) and (2 1 1) is observed according to the JCPDS 00-006-0696 database. Diffraction peaks of pearlite were not present due to its low volume fraction in this steel. It is readily apparent that no phase transformation occurs after the HPT process. However, there is a change in the peak intensities after five turns that indicates the occurrence of microstructural changes due to the formation of nanostructures and the dissolution of the pearlite lamella in the steel [8,9]. On the other hand, the full width at half maximum of the peaks does not show a significant variation and this is probably because the patterns were taken at the centers of the disks in the low deformation zone.

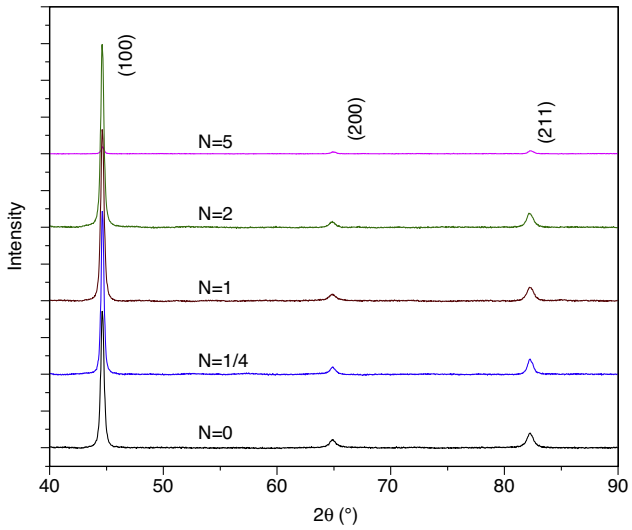

Fig. 1 – X-ray diffraction patterns for the samples processed through different numbers of turns.

Fig. 2 shows images of the surfaces of the samples processed through different numbers of rotations in HPT. These images were recorded near the mid-radius position at a distance of ~2 mm from the center of the disk. It can be seen that the structure at this position after 1/4 and 1 turn is not highly deformed and the microstructural features of ferrite and pearlite, shown in dark and bright contrast respectively, are clearly visible. At this stage, the soft ferrite grains were probably deformed by dislocation subdivision [10]. This type of SPD-induced grain refinement was reported earlier in ferrite [9]. After 2 revolutions in Fig. 2(c) ferrite and pearlite are present in the form of elongated grains and the spacing of the cementite lamella has decreased. The original grain boundaries appear to have been essentially compressed and they are elongated in the direction of grain flow with the thickness of the grains decreasing with increasing strain [11]. This characteristic of elongated grains is expected when deformed by a shear strain, even in the early stages of deformation [8]. After five turns in Fig. 2(d), no clear patterns or structural features are observed. This is probably due to the development of a very small grain size and the deformation of pearlite that occurs by further decreasing the interlamellar distance to form a structure of alternate ferrite and carbon enriched areas [12].

Fig. 3 shows images recorded at a distance of ~4 mm from the centers of the disks. Microstructural changes are observed at this position even after processing for only 1/4 revolution. After 1 revolution in Fig. 3(b), the pearlite is highly deformed