Thermal Stability of Titanium Alloy VT8M-1 with Ultrafine-Grained Structure

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Abstract

The paper considers the effect of a long-duration heating at a service temperature of 450°C on VT8M-1 with a coarse-grained (CG) and ultrafine-grained (UGF) microstructure. A duplex ultrafine-grained microstructure, composed of equiaxed grains of primary α-phase and an ultrafine constituent of α- and β-phases, was processed by thermal treatment and further rotary swaging. This type of a microstructure demonstrates a best combination of strength and ductility at room temperature in comparison with the CG structure. A thermal stability of an UGF state was studied at 450°C for 50, 100, 200, 300, 400, and 500 hours. The evolution of the alloy microstructure against the duration of heating was considered by transmission electronic microscopy (TEM), scanning electronic microscopy (SEM). No increase in the alloy structural elements and strength decrease resulting from a long-term annealing (up to 500 hours) at T=450°C have been observed. This proves a high thermal stability of the UGF structure and mechanical properties of VT8M-1 processed via rotary swaging.

Keywords: Ti alloys, rotary swaging, ultrafine-grained structure, mechanical properties, long-term annealing, thermal stability.

1. Introduction

Two-phase (α+β) Ti alloys that perform well within a wide temperature range are used to produce a number of critical parts both in aircraft engineering and engine manufacturing. In particular, VT8M-1 (Ti-5.7Al-3.8Mo-1.22V-1.35Sn), which is used to produce blades for a gas-turbine compressor (GTC), can be operated within a temperature range of 450–500 °C. The service conditions of modern constructions are currently increasingly demanding, while applied thermal or thermomechanical treatment (TMT) cannot provide the necessary strength of part [1,2]. One of the most promising technique to enhance the mechanical properties of metallic materials is the formation of a bulk ultrafine-grained (UGF) structure via severe plastic deformation (SPD) methods [3]. It has been shown earlier that the formation of UGF structure in two-phase Ti alloys (with Ti-6Al-4V as an example) leads to the increase in specific strength, fatigue resistance, and the enhancement in service properties of parts made of such materials [4]. However, UFG metals processed via SPD often demonstrate a reduced thermal stability arising from a high accumulated internal energy, which leads to a fast relaxation and decreases the recrystallization temperature [5]. As a result, the practical application of UFG Ti alloys at a service temperature of GTC parts is seriously limited.

It is well known that during rotary swaging billets are slightly reduced in a gradual manner, which enables to achieve a higher strain degree in the material along with a more homogeneous deformation of a billet [6]. As a result, an UFG structure is formed in a bulk billet from VTBM-1, which is reported in [7]. The possibility to produce long-sized semi-finished parts supporting further shape-generating operations is another advantage of this technique. The goal of this work is to research a thermal stability of UGF structures at a service temperature and mechanical properties of VTBM-1 processed by a rotary swaging with a view to assess the innovative capacity of the material in terms of its usage for GTC production.

2. Materials and methods

VTBM-1 Ti alloy was taken as a study material. The chemical composition of the alloy is shown in Table 1. Hot-rolled rods 70 mm in diameter were considered as an as-delivered state. The material was produced by VSMPO-AVISO Company (Verkhnyaya Salda, Russian Federation). Rods were obtained by a vacuum-arc melting. The initial material was annealed at 750°C for 1 hour (TT) in order to obtaining an equilibrium duplex structure.

Table 1. The chemical composition of VTBM-1 according to the Manufacturer’s Certificate (weight, %)

| Ti   | Al | Mo | Zr | Sn | Si | Fe | C  | N  | O  | H  | Impurities |
|------|----|----|----|----|----|----|----|----|----|----|------------|
| 87.86| 5.27| 4.00| 1.20| 1.26| 0.20| 0.034| 0.007| <0.003| 0.085| 0.0021| 0.1094     |

Rotary swaging took place at 750°C with a gradual reduction along the diameter of billets from 70 to 32 mm. As a result, true strain of 1.56 was reached. Strain degree was calculated from the following ratio: \( \varepsilon = \ln(S_2/S_1) \), where \( S_2 \) and \( S_1 \) are cross-section areas prior and after deformation, correspondingly.

To study thermal stability, coarse-grained VTBM-1 samples processed by TT (hereinafter referred to as CG) and by TT + rotary swaging (hereinafter referred to as RS) were held in a furnace at 450 °C. Continuous long-term annealing took place in furnaces produced by Noberterm Company. Holding time was 50, 100, 200, 300, 400, and 500 hours.

Tensile mechanical tests were performed using Instron universal testing machine at room temperature with a strain rate of 1×10^{-3} s^{-1}, according to ISO 6892-1-2009. Cylindrical specimens cut out in the longitudinal direction were tested.

The microstructure in various states was studied both in longitudinal and cross-sectional directions using JEOL JSM 6390 scanning electron microscope and JEOL JEM 2100 transmission electron microscope. Samples for TEM-foils were cut out using electrical discharge machining, mechanically thinned to a thickness of 100 μm and then electro-polished using a Tenupol-5 facility with a solution of 5% perchloric acid, 35% butanol and 60% methanol, at a polishing temperature within the range from ~20 to ~35 °C.

The X-ray diffraction (XRD) analysis was conducted on a Rigaku Ultima IV diffractometer. The samples were examined with CuKα-radiation (40 kV, 30mA) and the phase composition of the alloy was determined using the Rietveld method.

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3. Results and discussion

VTBM-1 processed by TT has a duplex microstructure (Fig. 1a). A mean grain size in a CG state of a primary α-phase is about 3 μm with a volume fraction α_glob = 65%.

Upon RS, the material microstructure is explicitly oriented along the axial direction (Fig. 1b), which is associated with the flow of the material during a RS. α-globules are elongated in the longitudinal section (Fig. 1b), and are twisted quite a lot in the cross section (Fig. 1c). The lamellar constituent upon deformation is divided into the fragments of oval and globular shapes. However, this process is rather heterogeneous with both almost unaffected laminas and strongly fragmented areas (which used to be integral laminas) observed.

The tensile mechanical testing of VTBM-1 in a CG state and after RS at room temperature shows an increase by 20% in the ultimate tensile strength in the RS-processed state as compared to a CG one (i.e. by 200 MPa) and constituted 1290 MPa, and a decrease by 5% in the elongation as compared to a CG state constituting 6-9%. Also, it should be noted that a uniform elongation in the material after RS changed slightly making up about 3.4% (Fig. 2). In a recent work, the ultrafine-grained structure in VTBM-1 was obtained by ECAP [7]. These specimens demonstrated a slight (about 1.2%) uniform elongation and rather rapid strain localization typical for many SPD-processed metals. It is reasonable to expect that the material subjected to RS has a higher fracture toughness compared to the ECAP-processed state.

By comparing the microstructure in CG and RS states subjected to a long-term annealing for 500 hours, it can be seen that annealing has almost no effect on the structure morphology and element dimensions (Fig. 1 and 3). Volume fraction α_glob remained at the level of 65% (Fig. 4) with a similar mean size of α_glob, (3 μm). The same picture was observed in an UFG state of VTBM-1 with grain size of primary α-phase and its volume fraction being at the same level after 500 hour heating (Fig. 3 and 4).

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TEM method was applied to study the microstructure after annealing at 450°C with a holding time of 50, 300, and 500 hours more thoroughly (Fig. 5). RS processing resulted in an increased density of both grain-boundary and intragranular dislocations leading to less distinct interphase boundaries. The resulting microstructure has the combination of a lamellar constituent retained after deformation (with a thickness of lamellas of about 150 nm) and grains as small as 300 nm of an oval shape. Such grains appeared during the formation of a cross interphase boundary, which resembles the fragmentation of lamellas. Local contractions with the formation of cross boundaries suggest that the fragmentation of lamellas take place by the mechanisms of groove formation and propagation.
The annealing for 50 hours at 450°C led to the decrease in dislocation density with clearly distinct boundaries of lamellas and grains as a result of dislocation redistribution and annihilation. The longer holding duration (up to 300 and 500 hours) gave rise to recovery processes as well as to the generation of some more perfect grain boundaries (marked with the arrow in Fig. 5). The appearance of moire contouring of such grains points to a low level of internal stress.

A mean cross lamella dimension remains at the level of 150 nm with a grain size about 300 nm. The analysis of TEM microstructure images of the annealed alloy conforms well to the X-ray results (Tab. 2). In particular, dislocation density fell from 10.8 to \(7.5 \times 10^{15} \text{m}^{-2}\) and crystallite size increased from 25 to 41 nm resulting from the longer duration of annealing up to 100 hours (Tab. 2).

Closer examination of the fine microstructure of samples subjected to RS + annealing for 300h revealed some interesting results. A diffraction pattern of the refined microstructure was obtained with an area of about 3µm². Both Ti α- and β-phases were observed as well as extra reflections forming a ring near the reflection \((200)\) (Fig. 6).

![Fig. 5. TEM images of VTBM-1 samples in the longitudinal section: following RS - (a); RS + annealing at T= 450°C for: (b) - 50h; (c) - 300h; (d) - 500h](image)

![Fig. 4. Volume fraction of globular α-phase versus holding time curve for both CG and RS-processed states of VTBM-1 treated at 450°C](image)

| State                | Lattice strain, % | Crystallite size, nm | Disl. density, \(10^{15} \text{m}^{-2}\) | Vol. fraction of beta phase, % |
|----------------------|-------------------|----------------------|------------------------------------------|-------------------------------|
| VTBM-1 CG+TT         | 0.07              | 124                  | 0.7                                      | 14.9                          |
| VTBM-1 HT+RS         | 0.48              | 25                   | 10.8                                     | 12.0                          |
| VTBM-1 HT+RS+ 50h annealing | 0.32            | 28                   | 9.6                                      | 9.9                           |
| VTBM-1 HT+RS+ 100h annealing | 0.21            | 41                   | 7.5                                      | 11.6                          |
It may be suggested that ageing processes caused by long-term annealing result in the precipitation of Ti-, Zr-, Si-based particles [8]. According to the literature on the subject (Hirsch), the formation of such ring-like reflections can actually point to the precipitation of a finely dispersed phase. However, we have not yet managed to accurately identify the precipitated phases. This will require some more time and a greater number of diffraction patterns.

Upon SPD, a volume fraction of β-phase in VT8M-1 decreases (Tab. 2), which relates to β→α transformation under the effect of large compressive stress [9].

Following long-term annealing, samples in a CG state and RS-processed were subjected to mechanical tensile tests. A long-term annealing of a CG state with a holding time of up to 300 hours resulted in a slight strengthening (Fig. 7a). At the same time, a notable reduction in a relative elongation starts only after annealing for 100 hours (Fig. 7b). The opposite situation in terms of both strength and ductility was observed after heating for 400 and 500 hours (Fig. 7a,b). This behavior can be attributed to β→α+β₂ decay resulting from a long-term heating at 450°С: initial strength increase and ductility drop is conditioned by the precipitation of secondary α-phase disperse particles with further particle coagulation resulting in the opposite situation [10].

The study of the thermal stability of a RS-processed state reveals slight fluctuations of ultimate stress, yield strength and relative elongation, which are within the limits of error. These minor variations, like an increase by 30-40 MPa, can be observed at the first point, following the annealing for 50 hours. This behavior is associated with the action of two competing mechanisms: namely, the strengthening in a RS-processed state (due to ageing during a long-term holding, like in a CG state) and the recovery in the strongly deformed material.

Thus, the rotary swaging results in both enhanced strength and ductility in the VT8M-1 alloy as compared to ECAP processing. Up to date, this technique is the most efficient way of producing rods on a commercial scale. High strength and thermal stability of the UFG VT8M-1 at 450°С open the way for successful application of the material to manufacture GTC parts.

4. Conclusions

The research of the effect of a duplex UFG microstructure in VT8M-1 processed via rotary swaging on the modification of its mechanical properties with an increase in a heating time up to 500 hours at 450°C leads to the following conclusions:

1. VT8M-1 processed by a rotary swaging preserves its thermal stability at 450°C after 500 hours of holding. At the same time, there has not been observed any notable change in a volume fraction of a primary α-phase of VT8M-1 samples both for a GG state and that subjected to a rotary swaging.

2. The mechanical properties of the UFG materials change slightly due to the action of two competing mechanisms: ageing during heating at 450°C and the structure recovery. The yield strength of the UFG alloy at room temperature exceeds the characteristics of a CG state by 200 MPa and remains at the same level after long-term annealing up to 500 hours.

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