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Dense dispersed shear bands in gradient-structured Ni

Yanfei Wang^{a,b}, Chongxiang Huang^{a,*}, Yusheng Li^c, Fengjiao Guo^a, Qiong He^a, Mingsai Wang^a, Xiaolei Wu^d, Ronald O. Scattergood^b, Yuntian Zhu^{b,c,**}

^a School of Aeronautics and Astronautics, Sichuan University, Chengdu, 610065, China

^b Department of Materials Science and Engineering, North Carolina State University, Raleigh, NC, 27695, USA

^c Nano and Heterogeneous Structural Materials Center, School of Materials Science and Engineering, Nanjing University of Science and Technology,

Nanjing, 210094, China

^d State Key Laboratory of Nonlinear Mechanics, Institute of Mechanics, Chinese Academy of Sciences, Beijing, 100190, China

ARTICLE INFO

Keywords: Gradient structure Shear bands Ductility Synergistic constraint Strength heterogeneity Nanostructured metal

ABSTRACT

During tensile deformation, nanostructured (NS) metals often fail soon after yielding by forming a localized shear band. Here we report the observation of high density of shear bands that are homogeneously dispersed in the NS layer of a gradient Ni sample. These shear bands were nucleated at early elastic/plastic strain stage, reached number saturation at $\sim 3\%$ strain, and remained arrested by the central coarse-grained (CG) matrix during the entire plastic deformation, resulting in a uniform tensile plasticity comparable to that of CG matrix. The formation of dispersed shear bands was promoted by the elastic/plastic interaction between NS surface layer and CG matrix, and affected by the surface roughness and the hardness variation in the NS surface layer. The width of shear bands remained constant, but the intensity of strain accumulation increased almost linearly with applied tensile strain, suggesting a stable shear banding process. Microstructure examination revealed that the strain in shear bands was accommodated by mechanically driven grain boundary migration and grain coarsening. These results clarify the fundamental questions: why/how does the NS layer supported by CG matrix achieve large uniform elongation? Moreover, the findings demonstrate the possibility of activating dispersed stable shear bands by synthesizing gradient architecture for optimized mechanical performances, i.e., a new strategy for evading the strength-ductility tradeoff in NS metals.

1. Introduction

Gradient-structured materials with increasing grain sizes from nanostructured (NS) surface layers to coarse-grained (CG) central layer have attracted intensive interests due to their superior combination of strength and ductility (Wu et al., 2014a, 2016; Cheng et al., 2018, 2014b; Wu and Zhu, 2017; Lu, 2014; Fang et al., 2011; Wei et al., 2014; Lin et al., 2018; Zhu et al., 2019; Wang et al., 2018a; Lu et al., 2019). They were found to have synergistic strengthening to produce yield strength higher than what is predicted by the rule of mixture (Cheng et al., 2018; Wang et al., 2018a; Wu et al., 2014b) and extra strain hardening (Li et al., 2017; Wu et al., 2014a) to retain good ductility.

The mechanical incompatibility between different layers is believed responsible for the observed mechanical behavior and superior properties (Wu et al., 2014a; Wu and Zhu, 2017). Specifically, during tensile testing the softer CG layer yields first to start

* Corresponding author.

https://doi.org/10.1016/j.ijplas.2019.08.012

Received 28 April 2019; Received in revised form 1 August 2019; Accepted 13 August 2019 Available online 19 August 2019 0749-6419/ © 2019 Elsevier Ltd. All rights reserved.

^{**} Corresponding author. Department of Materials Science and Engineering, North Carolina State University, Raleigh, NC, 27695, USA. *E-mail addresses:* chxhuang@scu.edu.cn (C. Huang), ytzhu@ncsu.edu (Y. Zhu).

plastic deformation, creating plastic/elastic interfaces where geometrically necessary dislocations are piled-up to create back-stress strengthening and extra work-hardening (Cheng et al., 2018; Kassner et al., 2013; Ming et al., 2019; Wang et al., 2018b; Yang et al., 2016; Zhu and Wu, 2019). When the whole sample is plastically deforming, the NS surface layers start unstable necking first, causing larger lateral shrinking in the NS layers than in the CG layer. This converts the applied uniaxial strain into multiaxial strains, which further helps with activating multiple slip systems and promoting dislocation hardening (Asaro, 1983; Wu et al., 2014a). In addition, residual stress produced during the processing of the gradient structure was also found to enhance mechanical properties (Moon et al., 2019; Yang et al., 2019).

A question arises on how the NS layer deforms in a gradient-structured sample. This question is critical for understanding the deformation mechanism and mechanical behavior of gradient materials. When deformed alone, NS metals often quickly start strain localization after yielding without much uniform elongation due to their low stain hardening capability (Ovid'ko et al., 2018; Valiev et al., 2016; Zhu and Wu, 2018). In gradient materials produced by surface mechanical attrition treatment, the NS layers have been reported to sustain very high plastic strains without apparent failure (Wu et al., 2014a, 2016; Cheng et al., 2018; Lu, 2014; Fang et al., 2011; Wei et al., 2014; Lin et al., 2018; Zhu et al., 2019). For a long time it was believed that the NS layers were deformed uniformly due to the constraint and support by the CG central layer (Fang et al., 2011; Lu, 2014; Wei et al., 2014; Wu et al., 2014a). However, it was recently reported that large shear bands were formed in the NS layer of gradient IF steel, which was delocalized along the gauge length to develop into a large strain accumulation zone with increasing tensile strain (Yuan et al., 2019). The shear bands accommodated the majority of applied tensile strain and provided some strain hardening to help with improving the overall strain hardening and ductility. This finding is surprising, because in both homogeneous NS and work-hardened polycrystalline materials the formation of such macroscopic SBs generally indicates the development of catastrophic strain localization, which will induce early fracture (Cheng et al., 2005; Ovid'ko et al., 2018; Yang et al., 2010). But it is not clear whether such stable shear banding is a universal behavior of the NS layers in gradient materials.

Shear banding is shear strain localization in a narrow zone, which often runs across multiple grain boundaries and twin boundaries during plastic deformation of polycrystalline metals. It is caused by local strain instability and accompanied with dramatic local orientation and texture change (Hong et al., 2010; Jia et al., 2013, 2012). Strain hardening was believed necessary to prevent shear banding (Borg, 2007; Mahesh, 2006). Nanostructured materials are especially prone to shear banding due to their low strain hardening capability. In the gradient structured metals, shear bands may be initiated in the NS layers. However, it may be difficult for the shear bands to propagate through the thickness/cross-section of the sample because the central CG layer typically has much higher strain hardening capability, which may act to stabilize the shear bands (Yuan et al., 2019). This needs to be further studied.

In this study we systematically studied shear band formation in the NS layer of gradient structured Ni plate with a large thickness of 3.6 mm using *in-situ* digital image correlation (DIC) technique. Nanostructured layers and microstructure gradients were produced with different processing techniques and parameters to study the mechanism of shear band nucleation and growth. It is found that high density of uniformly dispersed shear bands was formed instead of individual catastrophic shear bands that are typically reported in conventional homogeneous polycrystalline samples. Shear banding appears to be a primary mechanism for the NS layers to accommodate large applied plastic strain instead of being a pre-failure phenomenon.

2. Experimental procedures

2.1. Materials and processing

Commercial-pure Ni (99.60 wt%) plates with a dimension of 100 mm \times 90 mm \times 3.6 mm were used for this study. The Ni plates were firstly annealed in vacuum at 750 °C for 4 h, forming a fully recrystallized CG structure. Gradient plates with different surface roughness were symmetrically processed by means of rotationally accelerated shot peening (RASP) and piezoelectric surface nanocrystallization (PSNC) on both sides of as-annealed plates. RASP is a new surface mechanical attrition technique, which can vary both the speed and diameter of impacting balls (Wang et al., 2017). PSNC has relatively low processing efficiency but can produce extremely smooth NS surfaces (Li et al., 2016). Table 1 lists the main processing parameters for three different types of gradient samples Ni_{RASP- ϕ_2}, Ni_{RASP- ϕ_2}, In the RASP process, the samples peened by big balls are further treated using small balls for a longer time in order to reduce surface roughness.

Table 1

The processing parameters and referential label of gradient samples, where ϕ is the ball diameter, v is velocity, t is the processing time in RASP, and d is the penetration depth of the indenter in PSNC.

Sample	RASP step I			RASP step II	PSNC		
	ϕ , mm	<i>ν</i> , m/s	t, min	φ, mm	<i>v</i> , m/s	t, min	d, mm
Ni _{RASP-\phi2}	2	40	5	0.5	40	10	-
Ni _{RASP-01}	1	40	5	0.5	40	10	-
Ni _{PSNC}	-	-	-	-	-	-	0.3



Fig. 1. (*A*) Illustration of the geometry of gradient tensile specimen. In the coordinate, *Y* is the tensile loading direction, *X* is the sample width direction, and *Z* is the sample thickness direction with microstructure gradient. The *XOY* plane parallels to the NS surface. (*B*) Typical speckle pattern (the left subgraph) and detailed distribution of gray scale (the right subgraph) on the NS surface on gauge section. The red frame marks the 1452×450 pixel² effective calculation area.

2.2. Microstructural characterization and mechanical tests

The microstructures of gradient samples were characterized using scanning electron microscopy (SEM) and transmission electron microscopy (TEM). Focused ion beam was used to extract TEM foils precisely from selected positions. A Ni coating was deposited on the surface to protect the microstructure of topmost layer before ion beam cutting. TEM observation was performed in an FEI Tecnai G2 T20 microscope at 200 KV.

The variation of Vickers hardness along the depth was measured using a standard pyramid indenter at a load of 25 g for 15 s. Tests for each sample were repeated at four independent locations. Indentations were arranged along a zigzag line, and the space between neighboring indentations was three times longer than the diagonal of impression. Dog-bone shaped tensile specimens with a gauge dimension of $18 \times 4.9 \times 3.6 \text{ mm}^3$ were machined from as-processed gradient plates. Fig. 1A illustrates the geometry of gradient tensile specimen. Tensile specimens with only the topmost 400-µm-thick layer or the central 2.0-mm-thick core layer of Ni_{RASP- ϕ 1} material were prepared by polishing away the other layers of gradient tensile specimen. All tensile tests were performed at a strain rate of $5 \times 10^{-4} \text{ s}^{-1}$.

2.3. DIC strain characterization

The strain distribution and evolution on the nanostructured surface and the lateral surface (the surface parallel to the *YOZ* plane) of gradient tensile specimens during tension were *in-situly* recorded by DIC, using a short-focus optical lens. A random speckle pattern was prepared by spraying black paints on white background before performing DIC imaging. Fig. 1B presents a typical speckle image taken from the nanostructured surface. As indicated by the dotted red frame, an effective area of 1452×450 pixel² with a resolution of 9.7 µm/pixel was used for DIC calculation. The right subgraph shows the distribution of gray scale in a representative correlation window (36 × 36 pixel²). The mean intensity gradient of the gray profile of present speckle pattern is calculated as 48.6, meaning a high correlation coefficient for strain calculation (Pan et al., 2010).

3. Results

3.1. Surface roughness of the gradient samples

Fig. 2 shows the linear distribution of the surface roughness of the gradient samples, which were measured using a white light interferometry. The height resolution is \sim 20 nm. The maximum height difference between the convex peak and concave valley of



Fig. 2. The linear distribution of surface roughness measured in the three types of gradient samples.



Fig. 3. (*A*) A gradient microstructure from nanostructured surface layer to coarse-grained central layer; (*B*) Bright-field TEM image showing the NS at the topmost surface. (*C*) Micro-hardness profiles measured in the mechanical gradient layer of as-received gradient samples. The double-arrowed yellow line in (*B*) indicates the direction parallel to surface. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

roughness contour for $Ni_{RASP-\phi_2}$, $Ni_{RASP-\phi_1}$ and Ni_{PSNC} samples are measured as 80.8 µm, 37.1 µm and 2.1 µm, respectively. As shown, the sample processed with larger balls has higher roughness. The PSNC produced a very smooth surface due to its high striking frequency.

3.2. Gradient microstructure and microhardness

The Ni_{RASP- ϕ_1} sample was used to show typical gradient microstructure. Fig. 3*A* is a representative SEM image from the top NS layer to the CG central layer, which reveals a severely deformed sub-surface layer with obliterated initial grain boundary. Fig. 3*B* is a typical TEM image taken at the topmost surface layer, showing a mixture of elongated and equiaxed nanostructures with high density of dislocations. The solid symbols in Fig. 3*C* mark the microhardness profiles of the three types of gradient samples before tensile testing. As shown, the topmost layers of all samples have a hardness of ~270 Hv, which is about twice that of the CG matrix. The thicknesses of the gradient layers of the Ni_{RASP- ϕ_1} and Ni_{PSNC} samples measured from these hardness profiles are ~900 µm, ~780 µm and ~450 µm, respectively. The detailed gradient microstructure and formation mechanism of such gradient samples were reported in previous works (Liu et al., 2015; Wang et al., 2006).

3.3. Strength-ductility synergy

Tensile specimens with only the topmost 400- μ m-thick layer and the central 2.0-mm-thick core layer of Ni_{RASP- ϕ 1} material are referred to as freestanding nanostructured gradient surface layer (NGSL) and homogeneous CG core, respectively. Fig. 4 compares the



Fig. 4. Tensile engineering stress-strain curves for gradient samples and the freestanding NGSL and homogeneous CG matrix layer.



Fig. 5. The distributions of strain ε_y (the left column) and corresponding ε_x (the right column) on the surface of Ni_{RASP- ψ^2} (A1-A2), Ni_{RASP- ψ^1} (B1–B2) and Ni_{PSNC} (C1–C2) gradient samples and homogeneous CG (D1-D2) sample. They reveal the distribution and evolution of dispersive shear bands (SBs) in gradient sample. The shear bands are warm-colored in ε_y contour and cold-colored in ε_x contour. In the coordinate, Y is the tensile loading direction and X is the sample width direction. The number above each subgraph represents the average true tensile strain applied to the sample.}

tensile behaviors of gradient samples, the freestanding NGSL and homogeneous CG layer. The gradient samples exhibit improved yield strength and excellent ductility (uniform elongation), which indicates a superior strength-ductility combination. Interestingly, the yield strength of the freestanding NGSL reaches 504 \pm 6 MPa, while its ductility is lowered to ~6.8%. These results indicate that the NS surface layers were well supported by the CG layer and did not fail prematurely.

3.4. Dense dispersed shear bands in nanostructured layer

Fig. 5A–C present the distribution and evolution of strains in the tensile direction ε_y (the left five columns) and width direction ε_x (the right five columns) on the nanostructured surface of gradient samples during uniform elongation. Surprisingly, dense macroscopic shear bands (SBs), orientated at 49° ~55° with respect to the tensile axis, are uniformly distributed over the whole surface of all



Fig. 6. Strain distribution and evolution in SBs. (*A*) and (*B*) are the statistically averaged distribution of e_y and e_x across a representative SBs in Ni_{RASP- ϕ_2}. Curves in (*A*) and (*B*) with the same color are strain distributions at the same strain state. (*C*) Gauss fitting of the strain peak of a SB, from which the strain intensity (*I*) and the full width at half maximum (*W*) are measured. (*D*) The evolution of $|\overline{I}|$ and \overline{W} in the e_x peak of SBs in three types of gradient samples, with increasing applied strain. Each data point in (*D*) was averaged from the values of 8 randomly selected SBs. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

gradient samples. In contrast, such SBs are not found in the homogeneous CG sample (Fig. 5*D*), which implies that the low strain hardening capability of the NS layer is a precondition for the formation of SBs. It is also clear that the three types of gradient samples exhibit obvious difference in the morphology of SBs and the extent of strain inhomogeneity, which indicates significant effect of surface roughness and gradient layer thickness on the SB dimension, density and strain intensity.

For every type of gradient sample, the distribution and evolution of strains in 8 randomly selected SBs were analyzed. Taking a representative SB in the Ni_{RASP- ϕ_2} sample as an example, the distributions of ε_y and ε_x across the band at different tensile strains were statistically averaged along the shear banding direction and plotted in Fig. 6A and B, respectively. As shown, the peak strain in SB is nearly twice that in the background (i.e. non-shear banding zone). It should be noted that the strains in both SBs and background increase with applied strain.

Fig. 6C demonstrates that the strain peak profile across a SB can be fitted with a Gauss function to extract two important parameters: the strain intensity (*I*) and width at half maximum (*W*). Fig. 6D compares the evolutions of the average strain intensity (\overline{I}) and width at half maximum (W). Fig. 6D compares the evolutions of the average strain intensity (\overline{I}) and width at half maximum (W) extracted from the ε_x profile of the selected SBs, with increasing applied strain. As shown, the $|\overline{I}|$ increased linearly with increasing applied strain, suggesting a stable plastic deformation in SBs. In addition, the $|\overline{I}|$

Increased faster in the Ni_{RASP- ϕ 2} sample than in the Ni_{PSNC} sample. The \bar{W} remains largely constant during the whole plastic straining process, indicating constant SB widths, which contrasts the width growth in gradient IF steel (Yuan et al., 2019). The \bar{W} of ε_x strain peaks in the Ni_{RASP- ϕ 2}, Ni_{RASP- ϕ 1} and Ni_{PSNC} samples are ~0.33 mm, ~0.27 mm and ~0.24 mm, respectively. These values are much larger than those in homogeneous NS/UFG bulks (Jia et al., 2003, 2013; Hong et al., 2010; Carsley et al., 1995, 1998).

Fig. 7 shows SB nucleation at early strain stages in a $Ni_{RASP-\phi2}$ sample. SBs started to nucleate at the elastic-plastic transition stage (indicated by the white arrows). The density evolution of SBs, total length of SBs per unit area, as a function of applied strain is shown in Fig. 8. It is revealed that for all samples the SBs density increased quickly at the elastic-plastic transition and low plastic-strain stages, and then reached saturation at ~3% strain. It is also clear that the saturated density of SBs in the Ni_{PSNC} sample with smoother surface is nearly double that in the Ni_{RASP- $\phi2}$} sample.

Fig. 9 shows the SB evolution in a stand-alone NGSL. As indicate by the white arrow, a dominant SB with extremely high strain concentration quickly developed, penetrated through the cross-section and then caused fracture, leading to low ductility. Meanwhile, the SBs in uniform gauge section exhibit a lower density than in the integrated Ni_{RASP- ϕ_1} sample (Fig. 5*B*). These results suggest that dense dispersive SBs can be activated only when NS surface layers and CG matrix are deformed together.



Fig. 7. (*A*) SBs on the surface of $Ni_{RASP-\phi2}$ sample at different strain stages, revealing the fast increase of SBs density in elastic-plastic transition stage (P2–P3) and low plastic-strain stage (P3–P4). The white arrows indicate the initial embryos of inclined strain bands. (*B*) True stress-strain curves, showing the strain stage of P1–P4.



Fig. 8. The density evolution of SBs in gradient samples.



Fig. 9. SBs on the surface of freestanding NGSL that peeled from Ni_{RASP- ϕ_1} material: (A1) ε_{y} ; (A2) ε_x . The double-headed arrow indicates the runaway SB.

4. Discussion

The experimental results show that the enhanced plasticity of NS surface layer in gradient structure can be primarily attributed to the formation of dense macroscopic SBs and their stable evolution. Such dispersed and stable shear banding deformation was never observed in homogeneous metals under tension, which suggests that this mechanism is unique to the NS-CG gradient structure.

4.1. Unique characteristics of dispersed shear bands

As a non-crystallographic deformation mode, SBs are also often observed in materials without enough strain hardening to maintain uniform plastic flow, such as NS/UFG bulk metals and metallic glasses (Yang et al., 2010; Cheng et al., 2005; Hong et al., 2010; Jia et al., 2003; Carsley et al., 1995, 1998; Lu et al., 2013; Hays et al., 2000). It is interesting that both the configuration and evolution of the SBs observed here are very different from those in conventional homogeneous materials.

First, the SBs in gradient Ni are observed to distribute uniformly over the whole gauge section, and the plastic deformation in the SBs evolved stably with increasing applied strain (Fig. 6D). In contrast, in NS/UFG metals, SBs propagate quickly to fail the sample



Fig. 10. Schematic illustration of the load and constraint applied in the NS surface layer of gradient sample at the elastic-plastic transition stage. σ_y is external load, and σ_x^- is the compressive constraint from plastic CG matrix. θ is the preferential shear banding plane. The tilted short bands indicate the formation of early SBs.

due to the low strain hardening capacity and lack of constraint, and the gauge region far away from neck or fracture zone generally exhibits no SBs (Cheng et al., 2005; Yang et al., 2010). Second, the SBs in the gradient sample have macroscopic sizes in the order of millimeters (Figs. 6 and 5(A-C)). In contrast, except for the predominant unstable SB, the height of most SBs in the NS/UFG bulks is in the range of several tens or hundreds of nanometers, and their width generally ranges from submicron to several microns, i.e., micro SBs (Jia et al., 2003; Lu et al., 2013; Yang et al., 2010). Third, the density of SBs in gradient Ni reached saturation at \sim 3% strain (Fig. 8), while new SBs and branches in NS materials are nucleated through the whole plastic straining process (Jia et al., 2003). Fourth, the constant width of SBs in the gradient Ni differs greatly from the continuous thickening behavior of micro SBs in NS metals (Carsley et al., 1995; Hong et al., 2010; Jia et al., 2003). This is also different from the earlier report of SB widening in gradient IF steel (Yuan et al., 2019).

4.2. Nucleation of dispersed shear bands

SBs may be nucleated when the local shear stress (τ_{θ}) on the preferential shear banding plane (θ) reaches the critical value for shear instability ($\tau_{c\theta}$), which is preferentially activated at stress concentration sites under load (Ardeljan et al., 2015; He et al., 2003; Jia et al., 2013). In heterostructured materials, the boundaries between soft and hard domains are likely sites for stress concentration due to the mutual constraint-induced complex stress state and dislocation pile-up (C. X. Huang et al., 2018; M. Huang et al., 2018; Wang et al., 2019c; Wu et al., 2015).

For the gradient structure, during the elastic-plastic transition stage (see Fig. 7*B*), the elastic NS surface layer and the plastic inner layers form an elastic-plastic domain boundary. Due to the incompatibility in lateral shrinking rate between the inner plastic CG (apparent Poisson ratio $\nu \approx 0.5$) and the elastic NS surface layer ($\nu \approx 0.33$), a lateral compression stress (σ_x^-) is added to the NS surface layer (see Fig. 10) (Wang et al., 2018b; Wu et al., 2014b). This increases the shear stress τ_0 from $\sigma_y \sin \theta \cos \theta$ to ($\sigma_y + |\sigma_x^-|)\sin \theta \cos \theta$. In addition, the dislocation pile-up near this elastic/plastic interface also contributes to stress concentration (Wu et al., 2015; Zeng et al., 2016). As verified by the Vickers hardness map in Fig. 11, the surface layers themselves exhibit significant hardness variation, which indicates the existence of in-layer soft/hard domain boundary. This is expected to introduce stress/strain inhomogeneity and concentration sites as well (Sun et al., 2009; Wu and Zhu, 2017). Hence, the NS surface layers in gradient materials are believed to have abundant dispersed potential sites for nucleating SBs.

During deformation, once the stress concentration is built high enough (reaches τ_{cc}) to induce plastic deformation across a domain boundary, the local plastic deformation may become unstable in the NS surface layer because the harder domain likely has finer grain size and low strain hardening capability (Ardeljan et al., 2015; Wang et al., 2019b). This will nucleate a local SB, whose propagation in the length direction can only be stopped by another hard domain with even higher strength in the NS layer. This effect is similar to the activation of SBs in NS Ti composite (He et al., 2003), bimodal structured Cu (Wang et al., 2019b), and heterophase laminate (Jia et al., 2013), where the SB nucleation was promoted by the interaction-induced high internal stress at domain boundaries.

As discussed above, SBs can be effectively arrested by the neighboring harder domains in the NS layer due to the strength inhomogeneity. In the depth direction, a propagating SB will meet layers with larger grains whose higher plasticity will lower the stress at the band front, while their stronger work hardening helps with arresting the propagation. The arrest of propagating SBs provides opportunity to nucleate more SBs in less optimal regions, which eventually leads to the nucleation of high density of dispersed SBs that are homogeneously distributed over the whole NS layer. This is the primary mechanism for the fast increase of SB density at the yielding and low plastic-strain stages (Figs. 7 and 8). This process is similar to the arrestment of propagating SBs and the activation of multiple SBs in inhomogeneous metallic glasses consisting of ductile phase or layers (Hays et al., 2000; Kosiba et al., 2019; Lu et al., 2013; Sha et al., 2017; Zhou et al., 2013).



Fig. 11. Vickers hardness contour measured on the sub-surface layer (at the depth of $130\,\mu$ m) of as-received Ni_{RASP- ϕ 2} sample. The data map contains the value of 2400 homogeneously distributed indentations. The indent depth is ~2.75–3.15 μ m.

The orientation of SBs observed here is not along the plane with maximum shear stress, i.e., 45° with respect to the loading axis. This is a glasslike local shear instability response (Carsley et al., 1995; Donovan, 1989; Zhang et al., 2003), which has also been observed in high-strength NS and UFG metals. For example, micro SBs in UFG Pd–Ag alloy oriented ~50° to the tensile direction (Yang et al., 2010). Asymmetric shear banding orientations under tension and compression were observed in NS Fe–Cu alloy (Carsley et al., 1998, 1995). It has been proposed that such behaviors can be attributed to the yielding-related shear banding process and the normal stress-dependent yielding mechanism (Carsley et al., 1998). As described in the Mohr-Coulomb criterion ($\tau_{y0} = \tau_0 - \mu\sigma_0$), the shear yielding stress (τ_{y0}) on inclined θ plane of NS metals is sensitive to the in-plane normal stress (σ_0) (Donovan, 1989; Zhang et al., 2003). For the NS surface layer of gradient Ni sample under tension, the τ_{y0} is met more easily on planes 49° ~ 55° to the tensile axis, which induces plastic shear banding in this angle range.

4.3. Stable evolution of dispersed shear bands

As demonstrated by the linear evolution of strain intensity in SBs (Fig. 6D), SBs in NS layer remain stable to very large applied tensile strains without serious strain localization to fracture the sample.

Without the CG central layer the freestanding NGSL formed a dominant SB that caused the early fracture (Fig. 9), which verifies the critical role of the CG central layer in stabilizing SBs propagation. Fig. 12 presents the DIC strain mappings characterized on the lateral surface of gradient sample. As shown, SBs were originated from NS surface layers and weakened along depth with a gradual decrease in strain intensity. They partially intersected with each other, and were terminated by CG interior at the depth of $\sim 650 \,\mu\text{m}$. On the other hand, the in-layer mutual intersection of SBs constrained their in-layer propagation (Fig. 5*A*-*C*). These processes blunted bands sharpness, and hindered their catastrophic development, leading to a stable propagation process. Stabilization of strain localization by the neighboring softer domain or incompatible interface has also been reported in other heterogeneous structures (Azizi et al., 2018; Chen et al., 2008; Wu et al., 2017; Zhu and Lu, 2012).

The CG interior of present gradient Ni sample may also play a role in constraining the widening of dispersed SBs. In this study, the SB width remained constant with increasing strain (Fig. 6*D*), which is very different from the sustained widening of SB observed in the NS layer of gradient IF steel (Yuan et al., 2019). A closer examination of the gradient IF steel sample found that the gradient layer extended all the way to the center due to thin whole sample thickness (1 mm). In contrast, the Ni sample used here is much thicker (3.6 mm), which resulted in a much thicker CG central layer. Therefore, the likely reason for this difference is that the CG core in the gradient IF steel is too thin to effectively constrain the widening of SBs.

Moreover, the strain intensity of SBs increased gradually with applied strain (Fig. 6), which indicates that large strain gradient was formed in the shear banding zone during straining. As reported in gradient structured IF steel and CG/NS laminates, such strain gradient increase is known to lead to the accumulation of geometrically necessary dislocations and the development of back-stress work hardening (C. X. Huang et al., 2018; Li et al., 2017; Yang et al., 2016). The enhanced strain hardening will help to stabilize the plastic flow in SBs.

4.4. Microstructure evolution in shear bands

Dispersed SBs stabilize the uniform elongation of NS surface layer, which verifies a long-term conjecture that NS metals may serve as ductile materials as long as catastrophic strain localization is effectively suppressed (Fang et al., 2011; Yuan et al., 2019). An issue arising here is how the microstructure in SBs evolves to accommodate such large tensile strain.

We examined the microstructure of the topmost NS layers of $Ni_{RASP-\phi1}$ sample before and after tension (Fig. 13). The microstructure of as-processed surface layer is characterized by largely elongated grains with an average transversal grain size of 68 nm and an aspect ratio of 4.17 (Fig. 13A1 and A2). After deformation, surprisingly, equiaxed grains with even larger size and lower dislocation density are developed in shear banding zone (Fig. 13B1 and B2). These observations suggest that the plastic deformation of shear banding zone is dominated by grain coarsening, which has been interpreted as a mechanically-driven grain boundary migration



Fig. 12. SBs on the lateral surface of Ni_{RASP- ϕ 1} sample, measured at the applied strain of 7.22%. Z represents the thickness direction with microstructure gradient.

mechanism in NS metals (Chen et al., 2017; Fang et al., 2011; Wang et al., 2019a). The increase of grain size and change of aspect ratio are also detected in the zones outside SBs (Fig. 13*C1* and *C2*). However, comparing to that in shear banding zone, the extent of grain coarsening is less significant and the initial NS remains in local regions (marked by dotted cycles), suggesting that the rate and extent of grain boundary migration are positively correlated with stress/strain concentrations (Rupert et al., 2009).

As marked by red arrows in Fig. 13*C1*, under stress grain boundary migration occurred at the expense of neighboring highlydefective microstructures. This procedure results in dismantlement and annihilation of initial defects, thereby accommodating applied strain. The migration of grain boundary can be interpreted as a releasing process of distortion energy (Legros et al., 2008; Wang et al., 2012). Accompanying grain boundary migration in SBs, the formation of equiaxed grains with relatively low dislocation density provides substantial available room for defects storage (Fig. 13*B1*), which implies the regaining of strain hardening capability for shear banding zone (Huang et al., 2015; Yuan et al., 2019). With further increase in applied strain, the dominant plastic mechanism in shear banding zone will shift from grain boundary migration to conventional dislocation slip as the size of grown grains tends to saturate (Fang et al., 2011). This is the reason why some equiaxed coarse grains in shear banding zone exhibit high dislocation density (Fig. 13*B1*). More importantly, the regained strain hardening capability may also play a crucial role in stabilizing shear banding deformation.

4.5. Effects of surface roughness and strength heterogeneity on shear banding

Surface roughness could significantly affect the morphology of SBs, but is not the primary reason for SB nucleation. As shown in Fig. 2, the PSNC sample had a very smooth surface, but dispersed SBs appeared in its NS layer (Fig. 5C) with higher density than in the RASP samples which had much higher surface roughness (Fig. 8).

The statistical structural parameters of gradient samples, and the geometrical parameters and strain intensity of SBs are summarized in Table 2. It can be seen that SBs in samples with bigger surface roughness exhibit greater width, higher strain intensity but lower spatial density. This can be attributed to the effect of concave-convex roughness pattern on the development of stress concentration. Bigger peening balls produced the wider and deeper valleys on the surface layer (Table 2 and Fig. 2), which consequently led to larger stress concentration sites with longer inter-site distance and higher stress concentration under load.

Since the NS surface layers in all of the gradient samples were produced by the impact of high-speed balls or indenter, it was inevitable for the deformation to be heterogeneous (Panin et al., 2015). This led to inhomogeneous distribution of hardness (strength) in surface layers (Fig. 11), and thus provided extra soft/hard incompatible boundaries for stress concentration and strength heterogeneity to arrest the propagation of early SBs during deformation.

5. Conclusions

In summary, a unique plastic strain accommodation mechanism, formation of dense dispersed SBs, was experimentally revealed in gradient-structured Ni using DIC strain measurements. The main conclusions can be drawn as follow.

- (1) Dense dispersed SBs were uniformly distributed over the NS surface layer, and evolved stably during the entire plastic deformation, resulting in excellent uniform elongations (larger than 30.6%) which were much larger than that of a freestanding NS surface layer (~6.8%).
- (2) The nucleation of dispersed SBs started at an early elastic-plastic transition stage, and reached saturation at $\sim 3\%$ strain. The gradient microstructure, surface roughness and hardness variation introduced abundant soft/hard domain boundaries in the NS



Fig. 13. Bright-filed TEM micrographs and the statistical distribution of grain size in the topmost layer of Ni_{RASP. ϕ_1}: (*A*1, *A*2) As-received; (*B*1, *B*2) shear banding zone with a local ε_y of ~37%; (*C*1, *C*2) non-shear banding matrix with a local ε_y of ~16%. TEM foils of (*B*) and (*C*) were extracted from the SB and the neighboring matrix indicated by red arrows in Fig. 12, at the applied strain (whole-field average) of 21.7%. Dotted cycles and red arrows in (*C*1) mark the residue initial NS and the migrating grain boundary, respectively. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

Table 2

The structural parameters of gradient samples and the geometrical parameters of SBs, including the maximum height difference between the convex peak and concave valley (*H*) and the average space between neighboring concave valleys $(\overline{d_v})$ of surface roughness contour, the thickness of gradient layer (t_G), the thickness ratio of gradient layer to sample (t_G/t_S), the saturated SB density (ρ_b), the average space between neighboring SBs ($\overline{d_b}$), the average strain intensity ($|\overline{I}|$) and width at half intensity (\overline{W}) of ε_x peaks at the tensile strain of ~15.5%.

Material	Roughness	Roughness		Gradient layer		SBs			
	<i>Η</i> , μm	$\overline{d_{v}}$, mm	<i>t</i> _G , μm	t_G/t_I	ρ_b , m/m ²	$\overline{d_b}$, mm	\bar{W} , mm	<u> </u>], %	
Ni _{RASP-ø2} Ni _{RASP-ø1} Ni _{PSNC}	80.8 37.1 2.1	0.82 ± 0.46 0.38 ± 0.23 -	900 780 450	0.5 0.43 0.25	$\begin{array}{rrrr} 1330 \ \pm \ 100 \\ 2140 \ \pm \ 150 \\ 2430 \ \pm \ 120 \end{array}$	$\begin{array}{r} 0.91 \ \pm \ 0.67 \\ 0.56 \ \pm \ 0.44 \\ 0.49 \ \pm \ 0.31 \end{array}$	$\begin{array}{r} 0.33 \ \pm \ 0.12 \\ 0.27 \ \pm \ 0.09 \\ 0.24 \ \pm \ 0.11 \end{array}$	6.20 ± 3.8 5.62 ± 4.0 3.13 ± 2.1	

surface layer. The stress concentration induced by elastic/plastic interaction at domain boundaries promoted the early nucleation of SBs. The CG matrix and neighboring hard domains arrested the propagation of early SBs, which provided an opportunity to nucleate more SBs at dispersed stress concentration sites.

- (3) Stable shear banding at the large strain stage was maintained by the stabilization and mechanical constraints from the central CG matrix and the mutual intersection of SBs.
- (4) The large applied strain in shear bands was accommodated by mechanically-driven grain boundary migration. Grain coarsening in shear bands led to the regaining of strain hardening capability, which helps to stabilize shear banding deformation.
- (5) Surface roughness affected the morphology of SBs by affecting the stress concentration in the surface layer.

Acknowledgements

This work was supported by the National Key R&D Program of China (2017YFA0204403), the National Natural Science Foundation of China (Nos.11672195, 51931003 and 51741106) and Sichuan Youth Science and Technology Foundation (2016JQ0047). Yanfei Wang would like to acknowledge the support from Chinese Scholar Council.

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