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## Nano structured materials' mechanical characteristics

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### To Cite this Article

Dr.A.Padmanabham,Mr.V.N.V.Radha Kishna Murty, "Nano structured materials' mechanical characteristics",  
Journal of Science and Technology, Vol. 05, Issue 03, May-June 2020, pp280-286.

### Article Info

Received: 01-03-2020

Revised: 25-05-2020

Accepted: 02-06-2020

Published: 08-06-2020

### Abstract:

The mechanical characteristics of nanocrystalline (n-) materials are said to be affected by numerous microstructural factors, including as grain size and shape, distribution of pores, other flaws/defects, surface condition, impurity level, second phases/dopants, stress, duration of application, and temperature. There aren't any studies that go into great detail on the impact of each of these factors. The findings of both unequivocal and contradicting experiments are summarised below. The current theoretical levelThe mechanical behaviour of n-materials is discussed in some depth here. All rights reserved by Elsevier Science B.V.In terms of mechanical characteristics, microhardness, tensile and compressive behaviour, wear resistance, grain boundary sliding, and nanostructured materialsa steady stream of water

## 1. Introduction

Recent research has focused on 'nanostructured' (n-) materials with grain sizes less than 100 nm [1–3]. Even materials with a grain size have received favourable ratings in certain cases. There are 500 nm in this group. This should not be done. superplastic ceramics and highly deformed metallic materials are to blame. Materials with grain sizes ranging from 300 to 500 nm are often used. The high strain rate and/or low temperature superplasticity may be caused by Sub-micron grains are those with a grain size less than 100 nm. even though there are several uses for nanostructured materials Important, but not covered in this lecture.

## 2. Nanoscale structures in one or two dimensions and hybrid structures

Limiting the scope of the experiment yields positive outcomes. Nanoscale effects in a material that is otherwise normal. As This topic has previously been extensively discussed [3]. Some of the strategies will be briefly described. The advantages of adding a few to a few precipitates' materials with dimensions in the nanoscale range (The hardening of precipitation) is well-known. The moment n-Moan increase in hardness and fracture resistance was achieved by dispersing molybdenum in micrometre-sized Al<sub>2</sub>O<sub>3</sub>. strength and toughness were produced, however each characteristic peaked at a certain level. Composition in a different way the fracture strength of Al<sub>2</sub>O<sub>3</sub> with n-SiC dispersions of grain size 200 nm was increased. The high temperature mechanical characteristics were also improved by a factor of three. it's at its most thick, it's hard to tell the hardness of n-composites based on Al<sub>2</sub>O<sub>3</sub>, MgO, and Si<sub>3</sub>N<sub>4</sub> was the hardness change was consistent with the norm of increasing grain size and increasing SiC content mixtures. The hardness of n-composite Al<sub>2</sub>O<sub>3</sub>/SiC decreased much less with rising temperature than in Al<sub>2</sub>O<sub>3</sub>/SiC. Al<sub>2</sub>O<sub>3</sub> is a single piece.

This gap grew as the amount of time passed. Sic content, however, the temperature at which brittleness changes to ductility also went up at the same time. In the case of Mg–, the tensile strength and elongation to fracture Alloys of Zn and La with amorphous or hcp Mg phases particulates in the phase (interparticle distance: 5–10

nm<sup>3</sup>–10 nm) were much higher than in amorphous the compositions of equivalent Mg–Zn–La alloys Homogeneous shear was inhibited by the n-particles Same here. Al-based alloys have also been promoted for this method. A significant increase in resistivity is seen when metallic filaments of diameter 20–90 nm are incorporated into bulk conductors. Introducing a new conception Cu–Nb<sub>3</sub>Sn composites, n-scale multifilament's of Nb<sub>3</sub>Sn Unusual marriages of superconducting and mechanical technologies qualities may be obtained if desired. Composites created in situ using ultrafine number of permanent applications have been suggested for filaments. The use of magnets, especially when combined with other technologies mechanical strength and conductivity are essential. The mechanical and thermal characteristics of nanoscale mullite multilayers with a thickness of 1mm were found to be better. Comparison to their micrometre-sized counterparts. Partially smaller grains of stabilised tetragonal zirconia the performance of 70 nm silicon wafers covered with platinum was excellent. Sensors and non-volatile memory need ferroelectric characteristics. Applications that use memory Supermodulus and super compliance phenomena described in multi-layered thin film are at the centre of a debate.

composed of tiny, granular metal-containing layers sandwiched between two other materials. Additional testing is required before it can be determined whether these are inherent qualities of the layers or just artefacts. A list of the books referenced in the preceding paragraph may be found here.) [3] is the place to look. They have been eliminated from this section to save space.) Classification of n-composites by Niihara et al. The systems of ceramic–metal and metal–ceramic inter-, intra/inter- and nano–nanocomposites. within as well as the performance of inter-granular nanocomposites is greatly enhanced. High temperatures without losing any of their mechanical qualities. Novel characteristics of nano–nanocomposites are often observed. Machinability and super plasticity are two examples. The end objective is to creation of ceramics that are very hard and robust can only be achieved in hybrids of nanoscale and micrometre components, according to these scientists. Gas turbine applications requiring high temperature and high specific strength the future of vehicle and aeronautical technology is expected. The remainder of this document is dedicated to supporting information. Three-dimensionally organised nanostructures

### 3. Synthesis, consolidation and sintering

There are several techniques for creating nano powders (for a list of references see, e.g., [1,5]). Metals are readily consolidated and sintered regardless of the powder processing technique. Ceramic powders, on the other hand, are more difficult to densify. After a period of consolidation at ambient temperature, the

The porosity level in a ceramic may be as high as 60%. is around 20% in metals [6]. (In the finest practise to yet, the the porosity level in a compressed metallic specimen is about 5 percent, while ceramics have a range of 15–25 percent [5].) The sintering reaction is affected by the material, powder morphology, temperature, and the specimen's consolidated condition. Exerted pressure, as well as the technique by which it was administered the fortitude/The hardness of a material rises almost linearly with its density. [7–10] of the compact Powder compact densification Densification under an indenter in a microhardness test, for example, may be modelled as a quasi-steady state process.[3]. Sintering kinetics may be accelerated without the use of additives by hot pressing, sinter forging, or hot isostatic pressing. Putting pressure (HIPing). HIPing may also be used to get rid of Following pressure less sintering, there are closed pores. To Hot pressing should be used to maintain the final grain size below 100 nm. Sintering should be done at temperatures less than 0.5 T<sub>m</sub> (T<sub>m</sub> = temperature). On an absolute scale, melting temperature). (These techniques are also anticipated to be effective in the formation of near-net shapes.) consists of nanostructured components.) When the grain size is small, pore curvature dominates the driving force for densification during pressure assisted sintering, is around 2–3 nm. However, when the grain size is about 100 nm, it is more necessary to consider the amount of the applied pressure. In Both impacts are considerable in the range in between. The The findings are believed to show that there is a stress threshold, which is inversely proportional to grain size (and is required to produce increased surface area due to grain boundary movement), must be considered. Overcome before the rate of densification may be raised For Sintering is the process of producing a material with a certain grain size and density. Hot pressing is less effective than forging. These are empirical findings. Semi-quantitative concepts are quite useful in practise.

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Microstructures significant number of atoms in a nanostructured substance situated near the grain/interphase interface. Keeping such constructions exposed to high temperatures/service conditions quite tough.

When this is completed, it is shown that several characteristics are changed compared to those of conventional materials. Microstructural characteristics include grain size, form, holes and their distribution, other faults, surface quality, impurity level and crystal defect all impact mechanical characteristics. Grains are fundamentally equiaxed and their size distribution is narrow and lognormal.

Dislocations are infrequently detected within these grains, especially when the grain size in the lower region of the nanoscale scale. Even those that were observed, frequently had sessile configurations. The porosity level might be substantial and pore size distribution also is commonly lognormal. Enhanced self- and solute-solubilities and a distinct thermodynamic behaviour because the grain boundary phases are present in this class of materials compared with the typical variety [11–13]. Significant (unquantified) porosity is also present when the n-materials are created via deformation or crystallisation from an amorphous condition. In the absence of agglomeration, the pore size is lower than or equal to grain size. Grain boundary triple connections are favourable places for porosity formation. Grain boundary migration is inhibited by an open porous structure. Doping with an insoluble ingredient, grain border contamination, producing a composite at the grain border, adding a grain growth inhibitor or sintering under pressure are some of the techniques of inhibiting grain growth. Impure metals, e.g., n-Pd, considerable grain growth is present even at room temperature. Grain growth is also fast when the relative density is larger than 90 percent. Detailed study on the consequences of porosity, other flaws and their distribution, stress, time of its application, temperature, second phases, dopants, type of bonding in the material and microstructures on a range of length scales on the mechanical properties are absent. When the material was completely dense, its diffusion behaviour was comparable to that of the typical variation [1–3]. Nanostructured high melting point compounds (HMCs) have gotten fairly scant attention. There are just two recorded occasions when the final grain size was less than 100–150 nm: b-SiC of 92 percent relative density and a final grain size of 100 nm; MgO of 99 percent relative density and a final particle size of 100–150 nm. Dynamic recrystallisation induced via plastic deformation at a high temperature and pressure

microstructure in this type of materials, e.g., in boron nitride. The objective is to build super hard and ductile HMCs [3]. Two conflicting viewpoints exist with respect to the nature of grain limits in n-materials. One asserts that the structure of grain boundaries in n-materials is no different from those found in coarse grained materials [14]. But in another research based on computer simulation [15] the following conclusions have been reached: (i) long range periodicity is missing; (ii) grain boundary energy distribution is narrower than in bicrystals; (iii) grain boundaries are broader by approximately 50 percent but the width distribution is narrower than in bicrystals; (iv) in the zeroth order, the grain boundaries are anisotropic, cement-like phase, an image distinct from the structures found in glasses (which have short range order) and those comes from bicrystal investigations. It is crucial to agree upon the major aspects of grain boundary structure in n-materials at an early stage since any model for the deformation of n-materials will have to rely crucially on this description.

## 5. Hardness

When the consolidation temperature was larger than the temperature of hardness testing, microhardness was fairly uniform throughout the whole specimen. But defect size in a specimen might decrease with rising consolidation temperature can alter the outcome if the grain size is adjusted via shift in the temperature of consolidation [16]. In high melting compounds in the bulk, hardness was irrespective of grain size. In thin films toughness increased with decreasing grain size down to the nanometre range (lowest limit is not mentioned) [17]. In many n-ceramics and n-metal/ceramic composites hardness is basically unaffected up to such temperatures at which diffusion and grain boundary deformation processes become relevant. These temperatures are lower for n-materials than for their coarse-grained versions. At higher the hardness diminishes with rising temperature and this finding is due primarily to creep flow [1–3]. The grain size dependence of hardness of nanostructured metallic and intermetallic materials at ambient conditions at subambient temperatures has been the topic of several investigations (for a list of references, see [1–3]). The hardness of n-materials is much higher (typically around

five times) than that of their coarse-grained equivalents. But in one research on n-Cu and n-Pd [16] logarithmic creep might be noticed at room temperature.

There has been a controversy as to whether the “inverse Hall–Petch effect” (increasing hardness/strength with increasing grain size) observed at room temperature is real or not. It has been warned [9,18] that when annealing or a greater consolidation temperature is utilised to enhance grain size, the size of defects might reduce the density and/or the internal tensions in the compacts might alter. Then, these impacts will also have to be considered together with that of grain size when addressing the inverse Hall–Petch effect. Even Morris [5], who has taken a fairly dim view of most of the experimental evidence for the inverse Hall–Petch impact, has underlined that the findings of Wang et al. [19] in this regard on fully dense n-Ni (6–40 nm) seem to be real. In n-Cu, TiAl, Ni3Al, Fe3Al and TiAl–TiB2 specimen sat near full density furthermore grain boundary processes were present at room temperature. In TiAl specimens of 96 percent theoretical density an inverse Hall–Petch relation was observed for particle sizes lower roughly 30 nm at both ambient temperature and  $-30^{\circ}\text{C}$ .

In the complete set of tests, no evidence for dislocation activity or a density shift might be identified. And hence the impact was recognised as real [20]. (A full list of sources on this subject is provided in [1–3].) Thus, there is a case to argue that the slope of plots of ambient and sub-ambient temperature hardness ( $H_v$ ) of metallic materials vs the inverse of the square root of grain size ( $d^{-0.5}$ ) begins dropping as the grain size ( $d$ ) goes below roughly a micrometre, across a restricted lower grain size range  $H_v$  stays almost independent of  $d$  and finally the slope obtains a negative value as one approaches the lower limitations of the nm particle size range. In our perspective, then proof is required that the inverse Hall–Petch effect is really high-temperature creep and high-temperature creep have been hotly discussed. This impact is manifested in the form of superplasticity. The relationship between grain size and strength is obvious. Additionally, it's known as: Creep/super plasticity begins earlier in some areas. When the grain size is smaller, the temperature rises. Hence, it follows suggesting that n-metal materials have a very small grain size. The lesser the porosity, the smaller the size. Where the inverse Hall–Petch effect first manifests itself witnessed, sometimes down to temperatures as low as the outside air. In very porous materials, the hardness fluctuates. Time will be broken up and may be discarded at any point. In any case, incompatibilities in density, and the time-dependent nature of Like a power law, hardness changes in a similar way. It might be seen in very thick substances (see later). Other studies have shown an opposite Hall–Petch effect at room temperature. SnO2 and two metal–ceramic composites at various temperatures Near-full density (Fe–Tin, Ni–Tin) If these are the case, studies show that they can be replicated, ceramics processing. And a low-temperature composites manufacturing process is now conceivable.

## 6. Tensile and compressive response

The tensile strength and ductility of n-materials may be improved by annealing them after consolidation and/or polishing the specimen surface. There were no significant differences in stiffness between completely dense and coarse-grained samples when it came to the Young's modulus. When the skin's pores become visible in the presence of imperfections, the Young's modulus might be reduced by a factor of two to six. n-Pd showed signs of tensional creep. With a stress level well below the yield stress, at room temperature. When it comes to n-materials, there is no indication of super plasticity in tension in powder metallurgical processes. A304–306 Materials Science and Engineering A200–205 (2001) 203 203 route. More than 70 percent of participants in a recent trial [23] were found to be in this category. At  $1120^{\circ}\text{C}$ , n-ZrO2 doped with ZrO2 exhibited elongation in conjunction with Y2O3. Because the fracture began near the surface in this instance, after polishing, a fault was accidentally left in the specimen, there is room for improvement in this material's ductility. In comparison, the first grain of a strongly deformed It alloy At  $500^{\circ}\text{C}$ , the length of a 20-nm molecule may be stretched by 600%. Micrometre-sized grains of the same alloy show different properties. Only in the  $900\text{--}950^{\circ}\text{C}$  temperature range is there superplasticity. Nanocrystalline metallic metals may be produced in large quantities with this method since there is no residual porosity. Materials.

However, with materials of this sort, the grain boundaries even in non-equilibrium states with greater energy material with a given grain size may have a wide range of characteristics. It was found [24]. This necessitates grain-based property control. Boundary engineering is a field that will shape the future. Except for n-SnO2 and n-TiO2 of near-full purity, most ceramics tested under compression at room temperature had a density that

showed strain rate sensitivity. Brittle behaviour was shown by the temperature. The yield strength of an n-intermetallic Fe<sub>28</sub>Al<sub>2</sub>Cr was A compressive strain of around 1.4 could be achieved at a ten-fold increase in the fine grain (75 nm) version of the material. The temperature of a room. There was a higher creep rate when the porosity in the material was less than 20%. Ceramic powder compacts were just a little compressed. It is based on the porosity of the material. The microstructure of n-ceramics Lower temperatures may reveal alterations and/or creep mechanisms. Temperatures than those seen in fine-grained ceramics. strain rate dropped with increasing load in experiments involving constant load at a tension equivalent to densification and/or grain growth to limit of tolerable strain (see Section 3). It's possible to get the flow back by increasing the burden or stress. N-TiO<sub>2</sub> and n-TiO<sub>2</sub>/Y<sub>2</sub>O<sub>3</sub>At 810°C (0.5 Tm), compacts with a density close to their maximum might be compressed. The Power law creep equations were used to analyse the data.

Then-TiO<sub>2</sub> has a stress exponent N of 2.2–3.0, while n-TiO<sub>2</sub>/Y<sub>2</sub>O<sub>3</sub>. The number of atoms in the compact varied with its density. The inverse Hall–Petch effect is likewise compatible with this (as noted above). Assembled grains clear results are thwarted by growth/microstructural alterations Determination of the grain size effect on flow rate (3). Compression (and bending) ductility was examined in one research. Between 40 and 60 nanometres in diameter, the difference was more 60 nanometres. Reduced strain hardening and a weaker consolidation Indexes were considered to be the cause of this phenomenon. Fracture behaviour is poorly understood. Toughness and a lower transition from brittle to ductile the nanoscale temperature range has not yet been established with decreasing particle size as has been done in instance comprises micrometre-sized grains of materials. In-depth investigation of intermetallic TiAl (grain size range: 100 nanometres to the hardness reduced as the grain size dropped (70 nm). as intergranular fracture replaced intragranular cracking. There has been no research in the range of 5–100 nm particle size.) may be found in various sizes.) It was during this procedure that the transmission electron microscope was used. Sputtered gold specimens (TEM) will be examined using this technique. Intergranular crack formation in grains with a diameter of 10 nm was seen, as was dissemination via the creation of pores and linkages. Grain boundary activity, diffusion, and movement are all depending on strain rate. Sliding grain boundaries were critical in deciding the final product. Final toughness and ductility. There was no certainty as to whether or not there was a presence. If there was any mass deformation or activity at all, it was concentrated in the grained border areas [5].

## 7. Exhaustion and wear and tear

Studies on fatigue are few and far between. The cyclical nature corrosion resistance of n-Ni bulk specimens' comparable range of grain sizes (50–100 nm) was found in electro-deposition. Compared to the standard version. There isn't any hard data on the fatigue life and the development rate of fatigue cracks may be found. Any kind of n-materials Researchers have discovered that shear bands that are particularly powerful Cracks develop as a result of the formation of protrusions and intrusions. Infancy [5] All the results of Morris [5] have been condensed into a single summary. Researchers are interested in how n-materials wear. Surface coatings and bulk materials for cutting tools have taken into account However, despite the increased hardness. Toughness, on the other hand, was seen as a separate attribute from strength. Hardness. Extremely strong resistance to abrasive wear. Moreover, the wear resistance of the sliding surface was excellent. When the grain size was reduced, tool life increased dramatically. Up to 300 nanometres in length. For example, it has previously been used in industry for this purpose the production of tiny and medium-sized tools complicated forms on the cutting edge.) Is it possible to get the same benefits with even lower grain sizes? Also. So far, the reasons for the above-mentioned impacts have been given. The details of how clothes wear out [5] are a little hazy.

## 8. Mechanics-related

**modelling** More in-depth discussions may be found in [5,26–28]. The drawbacks of the improved dislocation methods models and other impromptu ideas work nicely documented. Because grain boundary slippage and diffusion are interdependent, one or the other may be used as a rate of return depends on the procedure being referred to as opposed to the more leisurely one. Dispersion has been largely ignored by academics. Regulating the speed at which something happens. Padmanabhan and his colleagues, on the other hand, Grain/interphase

boundary sliding has been considered by [29]. optimum superplastic flow and deformation control superplastic flow in metallic alloys [26–28] The glasses [30]. It has various benefits, the most noteworthy of which is that boundary sliding may be used as a rate-controlling mechanism. Superplastic flow [29] and molecular dynamics modelling studies have shown that it is possible to accurately anticipate grain rotations. With respect to n-materials [31,32]. (There will be no grain rotations in) regulated flow in which the boundary shear forces are kept to a minimum assumes that they will all quickly return to their original values This

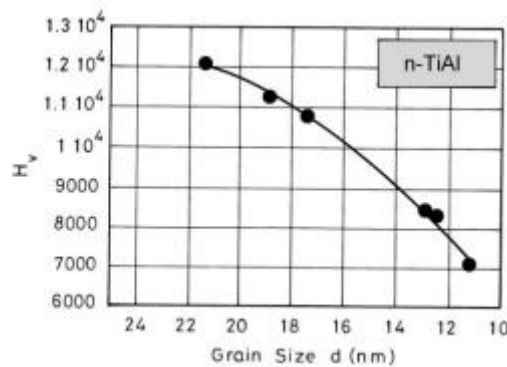
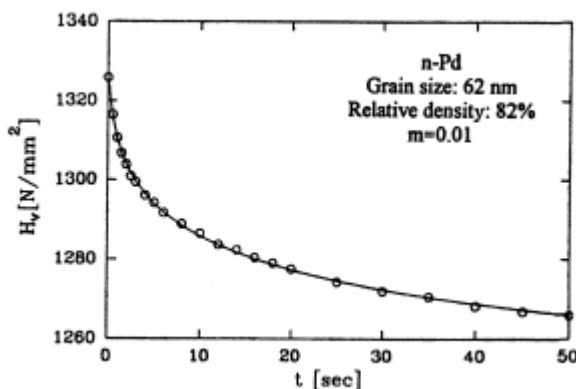


Fig. 1. A least-squares plot for n-Trial [20] in terms of Eq. (1). A grain boundary width of 0.57 nm is predicted.

$$H_v = H_{va} - \frac{m_2}{d} (d - d_0)^{0.5} \quad (1)$$

$$H_v = H_{va} - (H_{va} m) \ln(Bt + 1) \quad (2)$$

Here  $H_{va}$  is the hardness on load application,  $H_v$  the hardness at a given time  $t$ ,  $d$  the grain size,  $d_0 = 2 \sqrt{6\delta}$  (with  $\delta$  the grain boundary width) and  $m_2$  is a well-defined physical constant which at this level is treated as a fitting constant. (For details, see [26,27].) Fig. 1 is an example that uses the experimental data concerning n-TiAl [20]. Another prediction is that the microhardness variation with time will obey the power law (superplastic flow equalton). A simple analysis [30] leads to



## References

- [1] H. Hahn, K.A. Padmanabhan, *Nano structs. Mater.* 6 (1995) 191.
- [2] H. Hahn, K.A. Padmanabhan, *Advanced Materials and Processing, Vol. 3, Korean Institute of Metals and Materials, Kyongju, Korea, 1995, p. 2119.*

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- [3] K.A. Padmanabhan, H. Hahn, *Synthesis and Processing of Nanocrystalline Powder*, Minerals, Metals and Materials Society, Warrendale, PA, 1996, p. 21.
- [4] K. Niihara, A. Nakahira, T. Sekino, *Nanophase and Nanocomposite Materials*, MRS Symposium, Pittsburgh, PA, 1993, p. 405.
- [5] D.G. Morris, *Mechanical Behaviour of Nanostructured Materials*, Mater. Sci. Foundations 2, Trans. Tech. Publ., Aedermannsdorf, Switzerland, 1998, p. 1.
- [6] J. Eastman, R.W. Siegel, *Res. Dev. (January 1989)* 56.
- [7] G. Skandan, H. Hahn, B.H. Kear, M. Roddy, W.R. Cannon, *Mater. Lett.* 20 (1994) 305.
- [8] H.J. Hofler, R.S. Averback, *MRS Symposium*, Vol. 286, Pittsburgh, PA, 1993, p. 9.
- [9] G.E. Fougere, J.R. Weertman, R.W. Siegel, *Nanostruct. Mater.* 5 (1995) 127.
- [10] J.P. Ahn, M.Y. Huh, *Advanced Materials and Processing*, Vol. 3, Korean Institute of Metals and Materials, Kyongju, Korea, 1995, p. 2171.
- [11] R.W. Siegel, *Superplasticity of Metals, Ceramics and Intermetallics*, MRS Symposium, Vol. 196, Pittsburgh, PA, 1990, p. 59.
- [12] H. Hahn, J. Logas, R.S. Averback, *J. Mater. Res.* 5 (1990) 609.
- [13] H. Hahn, H.J. Hofler, R.S. Averback, *Defect and Diffusion Forum* 66–69 (1989) 549.
- [14] R.W. Siegel, *Mater. Sci. Eng. A* 168 (1993) 189.
- [15] S.R. Phillpot, D. Wolf, H. Gleiter, *J. Appl. Phys.* 78 (1995) 847. [16] G.E. Fougere, J.R. Weertman, R.W. Siegel, *Nanostruct. Mater.* 5 (1995) 127.
- [17] R.A. Andrievski, *J. Mater. Sci.* 29 (1994) 614.
- [18] G.E. Fougere, J.R. Weertman, R.W. Siegel, S. Kim, *Scripta Metall. Mater.* 26 (1992) 1879.
- [19] N. Wang, Z. Wang, K.T. Aust, U. Erb, *Mater. Sci. Eng. A* 237 (1997) 150.
- [20] H. Chang, C.J. Alstetter, R.S. Averback, *Advanced Materials and Processing*, Vol. 3, Korean Institute of Metals and Materials, Kyongju, Korea, 1995, p. 2107.
- [21] Y. Ogino, T. Yamasaki, B.L. Shen, *Advanced Materials and Processing*, Vol. 3, Korean Institute of Metals and Materials, Kyongju, Korea, 1995, p. 2149