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Residual stress measurement in thin films using a slitting method with geometric phase analysis under a dual beam (FIB/SEM) system

Ronghua Zhu¹, Huimin Xie¹, Xianglu Dai¹, Jianguo Zhu² and Aizi Jin³

 AML, Department of Engineering Mechanics, Tsinghua University, Beijing 100084, People's Republic of China
 Faculty of Civil Engineering and Mechanics, Jiangsu University, Jiangsu 212013, People's Republic of China
 Institute of Physics, Chinese Academic of Sciences, Beijing 100081, People's Republic of China

E-mail: xiehm@mail.tsinghua.edu.cn

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Abstract

Stress generated during thin film deposition is a critical issue for many applications. In general, the possible origins of the residual stress include intrinsic and extrinsic stresses. Since high residual stresses can cause detrimental effects on the film, such as delamination and wrinkle, it is of great importance to quantify the residual stress for the optimal design and the evaluation of its mechanical behavior. In this study, a method combining focused ion beam (FIB) milling and geometric phase analysis (GPA) is developed to assess the residual stress of thin films. The procedures of the residual stress measurement using this method include grating fabrication and slot milling by FIB, high-resolution scanning electron microscope (SEM) imaging of the grating before and after stress relaxation, and deformation analysis by GPA. The residual stress can be inferred from the released deformation using the reference displacements of the finite element model. As an application, this method was utilized to measure the residual stress in a TiAlSiN film, and the measured result is in good agreement with that obtained by the curvature method. In order to analyze the measurement error, the influence factors of Ga⁺ bombardment and the deposited platinum layer on the stress calculation are also discussed in detail.

Keywords: residual stress, thin film, focused ion beam, geometric phase analysis, finite element model

(Some figures may appear in colour only in the online journal)

1. Introduction

In recent years, with the development of micromachining technology, interest has risen in understanding the mechanical properties of thin films. In particular, residual stress evaluation has become a hot issue. Residual stresses in thin films are generated during the fabrication process such as during physical/chemical vapor deposition or the magnetron sputtering process, and can be mainly divided into two types: extrinsic stress and intrinsic stress. Extrinsic stress results from the difference in thermal expansion coefficients between the thin film and the substrate, while intrinsic stress is derived from the strong orientation of the crystal, which strongly depends on the deposition condition and the inherent attributes of the material. Considering that the excessive tensile stress in the film may lead to cracking, and that compressive stress may result in film delamination or buckling, the quantitative evaluation of residual stress in the film appears to be particularly important.

A range of non-destructive techniques are currently available for the estimation of residual stresses, such as x-ray diffraction [1], neutron diffraction [2] and the curvature method [3]. For x-ray/neutron diffraction measurement, since the thickness of the thin film/coating generally ranges from nanoscale to microscale, the corresponding diffraction peaks will be too weak to be conveniently measured. As for the curvature method, the curvatures of the substrate before and after deposition are measured by a white-light interferometer. With the measured data, the average residual stress can be calculated with Stoney's equation. However, due to its poor spatial resolution, the localized residual stress distribution cannot be obtained. In addition, surface micromachining techniques can also be utilized for in situ stress measurement, including the rotating method, buckling method, micro strain gauge method, long-short beam strain sensor method [4] and microbridge method [5, 6]. Xie and Li [7-9] proposed focused ion beam (FIB) moiré technology for the investigation of strain in microstructures. By fabricating high-frequency gratings on the cantilever beam surface of a microelectromechanical system (MEMS) structure, a moiré pattern could be formed at a specific magnification of the scanning ion microscope. In order to improve the accuracy of the in-plane displacement measurement, the FIB moiré method was combined with phase-shifting technology to measure the deformation of the micro-cantilever [10]. In the aforementioned literature, the stress relief depends on the specific machining process for the removal of the sacrificial layer, which are relatively complicated.

In order to simplify the approach of stress relief, FIB, an effective micromachining tool, has been utilized. Massl et al [11] proposed a micro-cantilever technique under the FIB workstation. The stress distribution along the film thickness direction could be determined based on the deflection of the micro-cantilever. This method is only suitable for residual stress measurement at the edge of the specimen. Kang et al [12] devised a novel method combining the FIB slitting and DIC technique under the SEM for in situ measurement of residual stress in thin films. Subsequently, this method has been applied to microscale residual stress assessments of various specimens. Vogel et al [13] successfully measured the residual stress in MEMS and nanoelectromechanical systems (NEMS). Winiarski et al [14, 15] obtained the residual stress distribution in bulk metallic glass. For higher measurement accuracy, Krottenthaler et al [16] provided a new H-bar geometry for measuring the residual stresses in thin films with the advantages of a linear displacement field and large deformation. In addition, the hole-drilling method [17] and the ring-core method [18, 19] have also been applied to microscale residual stress analysis.

Recently, the geometric phase analysis (GPA) method [20, 21] has been developed for full-field measurement of displacement and strain fields. In this method, regular gratings or grids are used as deformation carriers. The displacement can be measured by analyzing the gratings or grids before and after deformation. This technique has been utilized to measure the deformation at macro- and microscales [22–26]. In this study, a method based on FIB slot milling and GPA is developed for residual stress measurement. In the experiment, FIB is used to fabricate high-frequency gratings on the sample surface. SEM images of gratings before and after stress release are analyzed by GPA to obtain the deformation, from which the residual stress can be inferred by reference displacements of the finite element model. Furthermore, we analyze the influence factors of the experimental process on the measurement accuracy, where the Ga^+ bombardment on the surface and the error caused by the deposited platinum (Pt) layer are mainly discussed.

2. Geometric phase analysis

2.1. Principles of geometric phase analysis

The GPA method was firstly introduced by Takeda [20] and Hÿtch [21], and has already been applied in the displacement/ strain field analysis of crystal structures with high-resolution electron microscopy. Details concerning the theoretical discussions of this method can be found in the literature [27].

For an image with perfect periodic structures, the gray level at the position \vec{r} in the image is defined as:

$$I(\vec{r}) = \sum_{g} H_{g}(\vec{r}) e^{2\pi i \vec{f} \cdot \vec{r}}$$
(1)

where \vec{f} is the spatial frequency and $H_g(\vec{r}) = A_g(\vec{r}) e^{iP_g(\vec{r})}$ are the local Fourier components concerning the amplitude $A_g(\vec{r})$ and the phase $P_g(\vec{r})$.

With the grating frequency varying from \vec{f} to $\vec{f} + \Delta f$ after deformation, the displacement distribution can be calculated from the phase image with a crucial relative formula:

$$\Delta P_f(\vec{r}) = -2\pi \vec{f} \cdot u(\vec{r}).$$
⁽²⁾

For a one-way grating situation, that is to say that the image only contains the periodical structure in the x direction, equation (2) is simplified as:

$$\Delta P_f(x) = -2\pi f \cdot u_x. \tag{3}$$

Thus the displacement u_x and the strain ε_x along the *x* direction can be obtained from the phase image:

$$u_x = -\frac{1}{2\pi f} \Delta P_f(x) \tag{4}$$

$$\varepsilon_x = -\frac{1}{2\pi f} \frac{\partial \Delta P_f(x)}{\partial x}.$$
 (5)

For the cross-grating with the periodical structure along the mutually perpendicular x and y axes, the displacement field and the strain field can be expressed as the matrix form:

$$\begin{pmatrix} u_{x} \\ u_{y} \end{pmatrix} = -\frac{1}{2\pi} \begin{pmatrix} f_{1x} & f_{1y} \\ f_{2x} & f_{2y} \end{pmatrix}^{-1} \begin{pmatrix} P_{f_{1}} \\ P_{f_{2}} \end{pmatrix}$$
(6)
$$\vec{e} = \begin{pmatrix} \varepsilon_{xx} & \varepsilon_{xy} \\ \varepsilon_{yx} & \varepsilon_{yy} \end{pmatrix} = \begin{pmatrix} \frac{\partial u_{x}}{\partial x} & \frac{\partial u_{x}}{\partial y} \\ \frac{\partial u_{y}}{\partial x} & \frac{\partial u_{y}}{\partial y} \end{pmatrix}$$
(7)
$$= -\frac{1}{2\pi} \begin{pmatrix} f_{1x} & f_{1y} \\ f_{2x} & f_{2y} \end{pmatrix}^{-1} \begin{pmatrix} \frac{\partial P_{f_{1}}}{\partial x} & \frac{\partial P_{f_{1}}}{\partial y} \\ \frac{\partial P_{f_{2}}}{\partial x} & \frac{\partial P_{f_{2}}}{\partial y} \end{pmatrix}$$
(7)

where $u_x(u_y)$ is the displacement component in the x(y) direction and \vec{e} is the strain tensor of the deformed grating.

2.2. Sensitivity analysis

According to equation (2), the sensitivity of the GPA method depends on the phase information measurement, similar to the moiré fringe analysis [28, 29]. Considering that the fringe phase information is not directly reflected in the spectrum image, it is necessary to build the relation between the fringe phase and the spatial frequency to obtain the sensitivity of the phase in the frequency domain. The local spatial frequency f_n of the *n*th spectrum can be expressed as:

$$f_n = nf_0 + \frac{n}{2\pi} \frac{\partial p}{\partial x}, \ n = 1, 2, ...$$
 (8)

where f_0 is the initial frequency and p is the phase. The larger the value of n, the more noise is included in the frequency domain; thus in the GPA method, the displacement/strain distributions are calculated based on the first harmonic, as shown in figure 2(a). Substituting n = 1 into equation (8), the phase change between two adjacent points is derived [28]:

$$\Delta p = 2\pi (f_1 - f_0) \Delta x, \qquad (9)$$

where Δx is the image size in the spatial domain and f_0 is the fundamental frequency. The local spatial frequency f_1 varies between $f_{1\min}$ and $f_{1\max}$ (see figure 2(*d*)). Theoretically, the value of $\Delta \varphi$ tends to zero if the sidelobe width has a minimum. However, in practice, the first harmonic of the grating always scatters around the harmonic frequency in the frequency domain, so the detectable deviation between f_1 and f_0 is limited by the pixel size. Thus, the value of $\Delta f = f_1 - f_0$ is limited to one pixel in the frequency domain (without considering the noise effect). In addition, from the magic relationship between Δx and Δf :

$$\Delta x \cdot \Delta f = 1 / N.$$

The ultimate sensitivity can be obtained: $\Delta p_{\min} = 2\pi/N$, where N is the number of pixels of the image in the spatial domain in the GPA calculation. In other words, if a 500-pixel spatial domain with a grating pitch of 10 pixels is selected for analysis, the ultimate sensitivity can reach 0.02 pixels in theory.

2.3. Verification of the GPA performance under the FIB system

A digital grating with a cosinusoidal intensity distribution was generated. The intensity of the (i, j) pixel was defined by $I(i, j) = \frac{A}{4} \left[\cos\left(\frac{2\pi i}{T}\right) + \cos\left(\frac{2\pi j}{T}\right) + 2 \right]$, where the pitch *T* was 10 pixels and the amplitude *A* was 255. Considering that the pixel resolution used in SEM imaging is normally set as

 1024×884 pixels² or 2048×1536 pixels², an image with a similar resolution of 1024×1024 pixels² was generated. In order to present a typical case, a homogeneous strain field was imposed with the functions of the vertical displacement and strain expressed as $u_y = \varepsilon_0 y$, $\varepsilon_y = \varepsilon_0$. Figure 1 shows the generated digital image, in which the central deformed domain has a vertical strain of 0.001, which is marked by the yellow



Figure 1. Computer generated image with the periodic structure (the side parts are undeformed and the central part is deformed).

rectangle, and the other parts are the undeformed state. The main process of GPA is displayed in figure 2. By choosing the deformed domain, the frequency spectrum can be obtained by Fourier transform (see figure 2(a)). The unwrapped phase field and the displacement field can be obtained via the inverse Fourier transform and phase unwrapping process (see figures 2(b), (c)).

In this study, all the experiments are conducted in the dual beam system, which incorporates an ion beam and an electron beam. The ion beam can be utilized for specific material removal or repair, while the electron beam is used for nondestructive high-resolution imaging. The dual beam system has two critical issues: (i) secondary electron images are relatively noisy in comparison with CCD images; (ii) the ion beam may often cause undesired damage to the sample, and a significant deterioration of the local image pattern may occur, resulting in the decorrelation effect if DIC analysis is used.

For the first issue, in the DIC analysis, a systematic study of the simulated image shows that the error in the displacement measurement may vary from 0.05 pixels to 0.16 pixels due to the standard deviation of Gaussian noise ranging from 10 to 40 grey levels (classical value for SEM images acquired under the usual conditions) on the 256 grey level scale [30]. To assess the effect of noise, simulated images with Gaussian noise of different levels are analyzed by GPA. For comparison, the commercial DIC package Vic 2D from Correlation Solutions Inc. (Columbia, SC) was applied with a subset size of 29 \times 29 pixels² and a step size of 5 pixels to calculate the strain field. Still taking the example in figure 1, noise from 5 to 40 grey levels was introduced in the undeformed and deformed images. The results of the strain analysis by DIC and GPA are shown in figure 3. With the noise level increasing, the obtained average strain by GPA has an error of less than 3%, while the results by DIC have larger errors, indicating that for SEM images under the usual conditions, GPA has a good antinoise performance in deformation measurement.



Figure 2. (*a*) Fourier picture of deformed domain. (*b*) Filtered image of the spot f_{1y} . (*c*) Phase/displacement distribution in the analysis area. (*d*) Intensity distribution along the *x* direction in the spectrum.



Figure 3. The average strain value of the deformed domain obtained from GPA and DIC. The Gaussian noises vary from 0 to 40 grey levels.

For the second issue, material re-deposition around the milling area and material removal owing to Ga⁺ ion imaging are two main reasons for this phenomenon. The influencing degree of the former effect varies with different materials. For example, the low-carbon mild steel and ceramic coating have different experimental phenomena. The latter effect is difficult to avoid, because the positioning process of the SEM image is necessary during the ion milling. A verification experiment on the latter effect is conducted on an aluminum film under the dual beam system to assess the surface deterioration effect by ion bombardment. Generally, the current of the ion beam ranges from hundreds of picoamperes to several nanoamperes during deposition or etching. Thus, the ion beam current was respectively set as 200 pA, 1 nA and 5 nA in the experiment, with a dwell time of 10 µs/pixel. After different scanning times of the ion beam, a clear deterioration of the image quality occurred, as shown in figure 4. With the increase of the ion beam current and scanning times, the deterioration of the sample surface becomes more pronounced. As efficient parameters to assess the image quality in DIC, the correlation coefficient (zero-normalized cross-correlation) and mean intensity gradient [31] are utilized to analyze the deterioration of the image pattern after Ga⁺ bombardment. Figure 5 shows that the mean intensity gradient of the image drops sharply after the first few times of scanning. The ion beam will erode the surface of the sample, and the correlation calculation can still proceed well in this case. After ion scanning more times, the upper surface of the specimen is etched and then more materials appear below the surface. The analysis results verify that the mean intensity gradient will slightly increase with more scanning time. Due to the change in the original surface morphology, decorrelated points in these areas will be computed incorrectly with low correlation coefficients, and mismatch phenomena may occur. As shown in figure 5, upper limits of scanning times that have no unsolved areas in the images are 300 (500 pA), 35 (1 nA) and 6 (5 nA), respectively.

Compared with DIC, the gratings instead of speckles are utilized as deformation carriers in the GPA calculation, and in this case the ion beam will have no influence on the grating frequency. In addition, the ion scanning times can be limited



Figure 4. The bombarded surface with an ion beam current of 1 nA. From (a)-(e), the scanning times are 0, 10, 20, 50 and 100.

to at least one time with the optimal operation in the following experiment.

3. Experimental procedures

3.1. Sample preparation

The materials utilized in this study were a $1.5 \,\mu m$ TiAlSiN film on a quartz substrate and a $2\,\mu m$ TiAlSiN film on a silicon wafer by ion beam-assisted deposition, as shown in figure 6. The mechanical parameters of the films in terms of the Young's modulus were measured by a nanoindentation test. The nanoindenter XP with a Berkovich tip was set to continuous stiffness mode, and the average elastic modulus with an indention depth of 300 nm was evaluated by the Oliver-Pharr method. For this film, an elastic modulus of 378 GPa and a Poisson's ratio of 0.3 were utilized (see table 1). Prior to the FIB milling process, the film was measured with the curvature method to obtain an independent value of residual stress. Figure 7 shows the wafer curvatures after the film deposition with a radius of 245.4 ± 2.4 mm (quartz substrate) and $4743 \pm 53.7 \,\text{mm}$ (silicon substrate). The substrate before deposition can be regarded as a flat surface with a curvature close to zero. Based on the classical Stoney formula [32], the tests yielded the residual stress values of -1552 ± 15 MPa for the film with the quartz substrate and -1397 ± 16 MPa for the film with the silicon wafer substrate.

3.2. Grating fabrication and slot milling by FIB technique

In this study, the dual beam system (FEI DB 235) was utilized in the experiment for residual stress assessment. The direct writing capability of the ion beam is utilized to mill periodic gratings. There are two modes normally utilized in the milling process: the parallel mode and the serial mode. In the parallel mode, all the periodic lines or points in the whole region are etched in a scanning period, and repetitive scanning is necessary until the expected depth is reached, while in in the serial mode, the FIB is set to etch every periodic line or point to the expected depth, and then etch the next lines or points in a specified order. The latter mode may cause serious re-deposition during the milling process. Therefore, the parallel mode was chosen for the grating production in the experiment [23].

As for the fabrication of the deformation carrier under the FIB system, prior to the milling process, a Pt layer with a size of $20\mu m \times 40\mu m \times 50 nm$ was deposited on the TiAlSiN film using an ion beam-induced deposition technique. Then two 5000 lines/mm gratings at a size of $20 \mu m \times 20 \mu m$ were milled onto the Pt layer in turn. The ion beam current intensity was 50 pA in the milling process. The depth of the gratings was set to be the same as the Pt layer thickness. Figure 8 displays a simple schematic diagram of the gratings and the slot. The slot had a length of $40\mu m$, slightly larger than the width of the gratings, to guarantee deformation uniformity along the slot in the analysis domain. The whole experimental process can be summarized in the following steps: (i) Images of two gratings A and B were captured at a magnification of $12000 \times$, in which one pixel equals 11.2 nm. The parameters of SEM in this step were a working distance of 9.100 mm and an accelerating voltage of the scanning electron beam of 15 kV, which were kept constant in the SEM environment. (ii) The sample stage was tilted and the predesigned slot milled. The current intensity of ion beam was chosen to be 2nA for the higher etching rate. (iii) The sample stage was tilted to the original state and SEM images were captured in the same way as in the first step. The time gap between the two SEM images could be kept within 10-15 min under normal circumstances, which was acceptable for the system drift. Figure 9 shows the SEM images after the above steps.



Figure 5. (left) The mean intensity gradient and average correlation coefficient variation with the increase in scanning times of Ga^+ ion bombardment on the surface (from top to bottom, the beam currents are 200 pA, 1 nA and 5 nA); (right) The image grayscale distribution of the SEM image after different times of ion scanning.

Table 1. Mechanical properties of the TiAlSiN film.

	Elastic modulus/GPa	Hardness/GPa
TiAlSiN film	378.2 ± 8.5	19.5 ± 0.8

Due to the repeating tilt of the sample stage, the rigid body rotation of the specimen always occurs when capturing the image. However, with the unique trench-shaped geometry of the slot, zero displacement points are either along the middle of the slot or at an infinite distance, in theory. It is difficult to find a reference position with zero displacement. The other way to eliminate the influence of the rigid body rotation is the scanning moiré method. As figure 10 shows, a clear moiré pattern was generated at a magnification of $300 \times$ under SEM. If the rotation occurs among the image recording steps, the generated moiré patterns would be slanted (see figures 10(a), (*b*)). By adjusting the rotation angle of the scanning lines in



Figure 6. TiAlSiN films on the quartz substrate (sample A) and the silicon substrate (samples B and C) by ion beam-assisted deposition. (Sample sizes are $30.0 \times 5.0 \times 0.2$ mm and $15.0 \times 15.0 \times 0.5$ mm, respectively.)



Figure 7. Curvature of the quartz wafer measured after the deposition of TiAlSiN film. (*a*) The TiAlSiN film with a quartz substrate. (*b*) The TiAlSiN film with a silicon substrate.



Figure 8. Schematic diagram of the milling process. Gratings A and B are both $20\mu m \times 20\mu m$ and the slot is set to be milled between these two gratings.

SEM, the moiré patterns could be restored to the original state and the fringe should be parallel to the grating lines.

4. Results and discussion

4.1. Released displacement measurement

When applying GPA to the measurement of released deformation, an appropriate region first needs to be chosen. According to the finite element model (see figure 11), the maximum displacement occurs on the edge of the trench, and declines rapidly along the direction perpendicular to the slot. Therefore, the region with relatively large displacement was selected for the calculation, and is marked by the yellow dotted box in figure 9(*b*). Figure 12 displays the 3D/2D displacement field normal to the slot, and only the region between the dotted lines was used in the stress analysis. The average strain of this region, in which one pixel equals 11.2 nm, was $5123 \pm 458 \mu \varepsilon$ (quartz substrate) and $2465 \pm 205 \mu \varepsilon$ (silicon substrate).



Figure 9. (*a*) SEM image of the gratings after the milling step. (*b*) The magnification was increased to $12\ 000 \times$ to capture the domain in the red dotted dashed line, and the region in the yellow dotted box is cropped for calculation. (*c*) The cross-section of the trench after the ion milling.

A detailed analysis of the residual stress assessment can be found in the next section.

4.2. Finite element calibration

Finite element modeling of the slitting method was performed with the commercial software Simulia Abaqus. Since the slot length is much larger than the slot width or depth, a 2D elasticity analysis was performed with the plane strain model. A 2D mesh with CPE4R finite elements was suitable for the computation and the element density was large enough for sufficient computing accuracy. The modeling process can be mainly divided into two steps. The first step is to bring compressive/tensile residual stress to the elements. In the analysis, an initial stress of 100MPa was first introduced into the film (see figure 11(a)). The second step is to remove the element corresponding to the geometry of the microslot. It is also



Figure 10. The scanning moiré fringes generated at a magnification of $300 \times .$ (*a*) The moiré fringe shows that the grating has a clockwise rotation. (*b*) The moiré fringe shows that the grating has a counterclockwise rotation. (*c*) The grating parallel to the scanning lines of SEM.



Figure 11. (*a*) Step 1: The introduction of the initial stress (100 MPa tensile residual stress was introduced in this step) (*b*) Step 2: The stress field of the film after removing the elements of the slot. (*c*) Step 2: The displacement field of the sample after releasing the constraint.

worth noting that the geometry of the slot in the finite element analysis is a wedge-shaped slot and not the ideal square shape. This influential factor has been taken into account. After the stress release, the displacement/strain perpendicular to the slot could be obtained (see figure 11(b), (c)). A virtual displacement gauge could be imaged to be attached to the surface, and the location of this gauge is based on the region analyzed in the GPA analysis. Through changing the initial stress in the first step, different displacement/strain fields could be acquired in the analysis. The average strain of the same region as that used in the GPA analysis was $-324.6\mu\varepsilon$ (quartz substrate) and $-188.5\,\mu\epsilon$ (silicon substrate). Based on the linear relationship between the initial stress and the released displacement, a true local residual stress of -1.58 ± 0.14 GPa (quartz substrate) and -1.31 ± 0.11 GPa (silicon substrate) could be obtained, which is in reasonable agreement with the values obtained from the curvature method.

4.3. Effect of the Pt layer on the released displacement measurement

Ion beam-induced deposition was utilized for the Pt protection layer. During the Pt layer deposition, the accelerating voltage of the ion beam was 30 kV, the current intensity was 500 pA with the dwell time was $0.2 \mu s$. The thickness of the Pt layer was about 50 nm, which is much thinner than that of the TiAlSiN film. The Pt layer has two main functions: (i) to protect the film underneath it from Ga⁺ bombardment. The influenced depth ranges from ~15 nm to 50 nm in principle, so the effect of ion bombardment on the residual stress of the film can be considered negligible. (ii) to act as the marker layer. Owing to its layer production characteristics, the Pt layer can reduce the surface roughness effectively, which is beneficial to the grating production process.

The composition of the deposited Pt layer is PtC_8 from the results of [33], rather than pure platinum. It is porous and soft with an elastic modulus of ~8 GPa [34], much softer than the bulk platinum and the carbon film. In order to quantitatively evaluate the effect of the Pt layer on the released deformation, a finite element model similar to that in figure 11 is built, supposing an additional 50 nm-thick



Figure 12. (*a*) The 3D displacement of the gratings along the *x* axis. (*b*) The averaged experimental u_x displacements normal to the slot obtained from GPA. (Test 1 and test 2 are conducted on the film with quartz substrate. The calculated residual stresses are -1.48 GPa and -1.68 GPa, respectively. Test 3 and test 4 are conducted on the film with silicon substrate. The residual stresses calculated in the tests are -1.23 GPa and -1.38 GPa, respectively.)

Pt layer is entirely tied to the film, and only elastic deformation occurs after the residual stress release. Comparing the released displacement field of the model with and without the Pt layer, the former estimates the displacement with a discrepancy of less than 1%, which can be considered negligible in the present case.

5. Conclusion

In this study, a method combining FIB slot milling and GPA calculation was developed for the residual stress measurement of the micro-area under the FIB/SEM dual beam system. Taking advantage of sensor fabrication and high-resolution observation, the whole procedure of this method could be realized in one system. The residual stresses of TiAlSiN films were measured using this method, and the experimental results show good agreements with those obtained by the curvature method.

From our analysis, the possible influence factors of the operation process on stress assessment are considered. Due to the low elastic modulus and small thickness of the deposited Pt layer compared with the substrate and film, the influence is negligible in the present case according to analysis results from the finite element method simulation. Furthermore, the Ga⁺ bombardment may cause significant deterioration on the local image pattern, so it is necessary to decrease the exposure time in the FIB system with a lower beam current value.

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