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Mechanics of thin-film transistors and solar cells on flexible substrates

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Abstract

When devices are fabricated on thin foil substrates, any mismatch strain in the device structure makes the work piece curve. Any change of the radius of curvature produces a change in the size of the work piece, and thereby misalignment between individual device layers. To achieve tight tolerances, changes of curvature must be minimized throughout the fabrication process.

Amorphous silicon thin-film transistors and solar cells respond differently to externally applied tensile strain. The elastic deformation of the transistor is correlated with small increase in the electron mobility. When the tensile strain reaches $\sim 0.34\%$, crack formation starts and causes an abrupt change in the transistor performance. The performance of solar cells, on the other hand, does not change for tensile strain up to $\sim 0.7\%$. At larger strain the short-circuit current, open-circuit voltage, fill factor, and the efficiency gradually decrease.

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Keywords: Amorphous silicon; Thin-film transistor; Solar cell; Flexible electronics

1. Introduction

Recent research in thin-film electronics has been focused on the replacement of the traditional rigid glass plate substrate with plastic or metallic foils. Among metallic materials, stainless steel and molyb-

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denum foils have been utilized as substrates in the fabrication of thin-film transistors (Theiss and Wagner, 1996; Wu et al., 1997, 2002; Howell et al., 2000; Park et al., 2003) and solar cells (Yang et al., 2003). A number of plastic materials (organic polymers) also have been tested successfully in a variety of thin-film applications (Constant et al., 1994; Young et al., 1997; Burns et al., 1997; Burrows et al., 1997; Gleskova et al., 1998; Parsons et al., 1998; Lueder et al., 1998; Thomasson et al., 1998; Sandoe, 1998;

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Nomenclature			
d FF J_{sc} R T V_{oc} Y $Y^* = \frac{Y}{1-x}$	layer thickness (m); subscript s denotes substrate, f denotes film solar cell fill factor short circuit current (A) radius of curvature (m) temperature (K) open-circuit voltage (V) Young's modulus (Pa); subscript s de- notes substrate, f denotes film $\frac{1}{\sqrt{2}}$ biaxial stain modulus (Pa); subscript s denotes substrate, f denotes film	$Y'=rac{1}{1}$ $lpha$ arepsilon $arepsilon_{ m bi}$ arepsilon arepsilon arepsilon arepsilon arepsilon arepsilon arepsilon arepsilon arepsilon arepsilon arepsilon	$\frac{Y}{(-v^2)}$ plain strain modulus (Pa); subscript s denotes substrate, f denotes film coefficient of thermal expansion (K ⁻¹); subscript s denotes substrate, f denotes film mechanical strain built-in strain in the film Poisson ratio; subscript s denotes sub- strate, f denotes film solar cell efficiency mechanical stress (Pa)

Carey et al., 2000; Sazonov and Nathan, 2000; Boucinha et al., 2000; Kane et al., 2001; Ichikawa et al., 2001; Hsu et al., 2002a; Brida et al., 2002; Takano et al., 2003; Cheng and Wagner, 2004; Gelinck et al., 2004; Shahrjerdi et al., 2004; Nomura et al., 2004; Monacelli et al., 2004; Choi et al., 2004).

There are three main reasons for the attraction of plastic and metallic foils. Unlike glass, the thickness of these materials can be substantially reduced while maintaining their integrity, leading to thin and lightweight products. At the same time, these thin substrates add new functionality to thin-film electronics, namely the flexing and non-planar shaping. Finally, the foil substrates lend themselves to roll-to-roll fabrication.

One faces several new issues when fabricating devices on thin foils. These are usually not encountered during the fabrication of these devices on thick plates of glass. Firstly, the devices experience variable stresses during the manufacturing process that may lead to substantial change in curvature. This leads to a change in the size of the work piece and ultimately to misalignment between different layers of the device. This is important for devices where mask overlay alignment is critical. Therefore, the radius of curvature must be carefully controlled during the fabrication. Secondly, the device application may require intentional bending, stretching, or non-planar shaping after the fabrication. Therefore, one needs to understand the behavior of thin-film devices under strain, and the fracture strain and fracture mechanism of the device layers.

Even though a detailed understanding and comprehensive mechanical theory do not yet exist, a number of experimental results are available and simple mechanical theories have been worked out (Gleskova and Wagner, 1999a,b, 2001; Suo et al., 1999; Gleskova et al., 2000, 2002, 2004; Wagner et al., 2000, 2002; Hsu et al., 2002b, 2004; Jones et al., 2002; Servati and Nathan, 2005). The purpose of this paper is to summarize the current knowledge of the mechanics of thin-film electronics with a focus on amorphous silicon thin-film technology. In the calculations we emphasize two-layer structures of substrate and film. Such structures are simple enough to be treated analytically, yet they provide a basic understanding of the mechanics of thin-film devices on flexible substrates.

2. Curvature induced during manufacturing

Thin-film devices are built on substrates layer-bylayer, often at elevated temperature. Strain develops in the structure by built-in stresses in the deposited layers (Hooke's law: $\sigma = Y \cdot \varepsilon$, where σ is stress, Y Young's modulus, and ε strain), or, upon cooling down, by the differences in the thermal expansion and humidity coefficients between the deposited film and the substrate, or between different films. The mechanics of the film-on-substrate structure depends strongly on the elastic (Young's) moduli and thicknesses of the substrate Y_s , d_s and the thin film Y_f , d_f .

When $Y_f \cdot d_f \ll Y_s \cdot d_s$, the substrate dominates and the film complies with it, as a thin-film transistor (TFT) or solar cell do on a plate glass substrate. The stress in the substrate is small, and the film/substrate couple curves only slightly, even when the film is highly stressed. The strain is biaxial, and the structure forms a spherical cap with the radius of curvature given by:

$$R = \frac{d_{s}}{6 \cdot \frac{Y_{f}^{*}}{Y_{s}^{*}} \cdot \frac{d_{f}}{d_{s}} \cdot \varepsilon} \cdot \varepsilon$$
$$\cdot \frac{\left[1 - \frac{Y_{f}^{*}}{Y_{s}^{*}} \cdot \left(\frac{d_{f}}{d_{s}}\right)^{2}\right]^{2} + 4 \cdot \frac{Y_{f}^{*}}{Y_{s}^{*}} \cdot \frac{d_{f}}{d_{s}} \cdot \left(1 + \frac{d_{f}}{d_{s}}\right)^{2}}{1 + \frac{d_{f}}{d_{s}}} \tag{1}$$

Here $Y_{\rm f}^* = \frac{Y_{\rm f}}{1-\nu_{\rm f}}$ and $Y_{\rm s}^* = \frac{Y_{\rm s}}{1-\nu_{\rm s}}$ are the biaxial strain moduli of the film and substrate, respectively. $\nu_{\rm f}$ and $\nu_{\rm s}$ are the corresponding Poisson ratios. ε is the mismatch strain. The mismatch strain ε has two dominant components. One is the thermal mismatch strain caused by the difference between the coefficient of thermal expansion of the substrate, $\alpha_{\rm s}$, and that of the film, $\alpha_{\rm f}$. The other is the builtin strain $\varepsilon_{\rm bi}$ in the deposited film. Therefore,

$$\varepsilon = (\alpha_{\rm f} - \alpha_{\rm s}) \cdot \Delta T + \varepsilon_{\rm bi} \tag{2}$$

where ΔT is the difference between the deposition and the room temperatures. *R* is negative when the film is under compression and the structure curves with the film being on the convex side. *R* is positive when the film is under tension and the structure curves with the film being on the concave side.

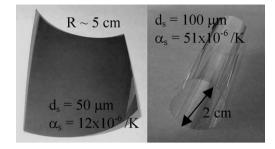


Fig. 1. PECVD silicon nitride film deposited on two different plastic substrates. The substrate thickness and the coefficient of the thermal expansion are shown for each substrate. The film is deposited on the top and the structures roll into cylinder.

A stiff film and a compliant substrate, for example amorphous silicon device on an organic polymer foil, may have similar products of elastic modulus and thickness, $Y_f \cdot d_f \approx Y_s \cdot d_s$. Such equal strength of film and substrate gives rise to complicated mechanical situations. The structure rolls into a cylinder instead of forming a spherical cap. An example is shown in Fig. 1, for a 500-nm thick layer of silicon nitride deposited on two different plastic substrates by plasma enhanced chemical vapor deposition (PECVD) at 150 °C. The Stoney formula is no longer valid. Using the procedure outlined in (Suo et al., 1999), the radius of curvature *R* is given by:

$$R = \frac{d_{s}}{6 \cdot \frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} \cdot \varepsilon} \cdot \left\{ \frac{\left\{ \left[1 - \frac{Y'_{f}}{Y'_{s}} \cdot \left(\frac{d_{f}}{d_{s}} \right)^{2} \right]^{2} + 4 \cdot \frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} \cdot \left(1 + \frac{d_{f}}{d_{s}} \right)^{2} \right\} \left[(1 - v_{s}^{2}) + \left(\frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} \right)^{2} (1 - v_{f}^{2}) \right]}{\left(1 + \frac{d_{f}}{d_{s}} \right) \cdot \left(1 + \frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} \right) \cdot \left[(1 - v_{s}^{2})(1 + v_{f}) + \frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} (1 - v_{f}^{2})(1 + v_{s}) \right]} + \frac{3 \cdot \left(\frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} \right)^{2} \left(1 + \frac{d_{f}}{d_{s}} \right)^{2} \left[(1 - v_{s}^{2}) + (1 - v_{f}^{2}) \right] + 2 \cdot \frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} \cdot (1 - v_{s}v_{f}) \cdot \left(1 + \frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} \right) \cdot \left(1 + \frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} \right)^{2} \left(1 + \frac{d_{f}}{d_{s}} \right)^{2} \left[(1 - v_{s}^{2}) + (1 - v_{f}^{2}) \right] + 2 \cdot \frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} \cdot (1 - v_{s}v_{f}) \cdot \left(1 + \frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} \right) \cdot \left(1 + \frac{Y'_{f}}{Y'_{s}} \cdot \frac{d_{f}}{d_{s}} \right) \right\}$$

$$(4)$$

For amorphous silicon devices on glass substrates $Y_f \cdot d_f \ll Y_s \cdot d_s$, which allows to neglect the second fraction in Eq. (1) and to simplify to the Stoney formula (Freund, 1996; Finot et al., 1997; Cu, 1998):

$$R = \frac{d_s}{6 \cdot \frac{Y_t^s}{Y_s^s} \cdot \frac{d_f}{d_s} \cdot \varepsilon}$$
(3)

In such cases R is very large.

Here $Y'_{\rm f} = \frac{Y_{\rm f}}{1-\nu_{\rm f}^2}$ and $Y'_{\rm s} = \frac{Y_{\rm s}}{1-\nu_{\rm s}^2}$ are the plane strain moduli of the film and the substrate, respectively. If the Poisson ratios ν of the film and the substrate are identical, Eq. (4) simplifies into the form published previously (Suo et al., 1999):

$$R = \frac{d_{s}}{6 \cdot \frac{Y_{f}}{Y_{s}} \cdot \frac{d_{f}}{d_{s}} \cdot \varepsilon} \cdot \frac{\left[1 - \frac{Y_{f}}{Y_{s}} \cdot \left(\frac{d_{f}}{d_{s}}\right)^{2}\right]^{2} + 4\frac{Y_{f}}{Y_{s}} \cdot \frac{d_{f}}{d_{s}} \cdot \left(1 + \frac{d_{f}}{d_{s}}\right)^{2}}{(1 + v) \cdot \left(1 + \frac{d_{f}}{d_{s}}\right)}$$
(5)

The mismatch strain ε is again given by Eq. (2). When the mechanical properties of the film and the substrate and the radius of curvature *R* are known, one can easily extract the built-in strain ε_{bi} in the film (Cheng et al., 2005). Our current focus, however, is on the understanding of how to keep the radius of curvature large, and preferably infinite, since this is an important factor in the device fabrication.

From analysis of Eq. (5) one can easily conclude that making $Y_f \cdot d_f \ll Y_s \cdot d_s$ makes R large. This can be achieved by: (1) choosing a thick substrate (not desirable for flexing or "shaping" applications), (2) keeping the device structure very thin (may not be possible), (3) choosing a substrate with large Young's modulus (not possible for plastic substrates), or a combination of these measures. R can also be made large by keeping the mismatch strain ε close to zero. One can: (1) choose a substrate with a coefficient of thermal expansion (CTE) close to that of the device layers (This is not possible for plastic substrates whose CTE usually is much larger than that of silicon device films. Stainless steel substrates have relatively low CTE.), (2) minimize ΔT by lowering the deposition temperature. (In thin-film silicon technology, lower deposition temperature leads to worse electronic properties. Here, the organic electronics, for example polymer light emitting diodes and thin-film transistors have an advantage.), (3) compensate the CTE mismatch with built-in strain in the film, such that $(\alpha_{\rm f} - \alpha_{\rm s}) \cdot \Delta T = \varepsilon_{\rm bi}$ (The built-in strain depends on the deposition conditions and sometimes can be easily adjusted.).

There are two more solutions for dealing with undesirable curving. One is to attach the substrate foil to a rigid carrier for the duration of the fabrication. The rigid carrier suppresses the bending of the structure, and sets the strain in the plane of the film by the thermal strain of the carrier, which is small. This approach, used by a number of research groups, allows a larger selection of plastic materials. However, one needs an adhesive whose glass or decomposition temperature is higher than the highest process temperature. In addition, the adhesive should not outgas in the vacuum and should resist wet and dry processing. Once the fabrication is completed, the adhesive should permit easy separation of the carrier and the work piece. Even though this approach has been demonstrated in the laboratory, it is not an ideal manufacturing solution. The second solution is based on the fact that the patterning of continuous layers into islands relieves the global mismatch strain. However, strain is concentrated around the edges of the islands.

3. Misalignment caused by curvature

Fig. 2 shows several thin films typically used in the amorphous silicon thin-film transistor fabrication (a-Si:H TFT) deposited on 50 µm thick Kapton E. As described in the previous section, different thin film materials exhibit different built-in strains resulting in different radii of curvature R. It has been shown experimentally that a change in curvature affects the alignment between different layers of the device (Cheng et al., 2005). If one starts with a flat substrate, the misalignment increases with decreasing radius of curvature, because a larger strain is developed when the work piece is flattened for mask alignment. The larger the radius of curvature, the less flattening is required and the smaller is the misalignment. In a-Si:H TFTs on Kapton E, the misalignment between the gate and the source/drain

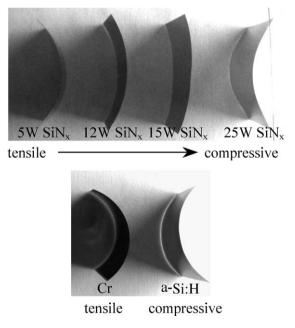


Fig. 2. Curvature induced by mismatch strain in films of Cr, a-Si:H, and SiN_x deposited over a range of RF power, all on 50- μ m thick Kapton[®] 200E polyimide substrates. All films are facing to the left. The 300-nm to 500-nm thick SiN_x and the 250-nm thick a-Si:H films were deposited by PECVD at 150 °C, and the 80-nm thick Cr was deposited by thermal evaporation without control of substrate temperature. The built-in stress of Cr is tensile and that of a-Si:H is compressive. The built-in stress in SiN_x changes with the deposition power (Wagner et al., 2005).

electrodes as a function of the deposition power of the gate dielectric easily can reach values of 500 ppm. In an optimized structure the misalignment was reduced to \sim 100 ppm (Cheng et al., 2005). Therefore, the radius of curvature needs to be carefully controlled throughout the fabrication. We are currently developing a mathematical model of this phenomenon with the aim to control it.

4. Externally applied strain

All integrated circuits are fabricated flat. The use of the fabricated circuit may require that it is bent (once or repeatedly), stretched, or shaped after the fabrication. Therefore, it is important to understand the electrical and mechanical behavior of the devices under externally applied strain.

a-Si:H TFTs respond to increasing mechanical strain by elastic deformation followed by fracture. The response of the TFT is determined by the magnitude of the applied strain regardless of its origin, for example, bending, stretching or shaping (Suo et al., 1999; Gleskova and Wagner, 2001, 2002; Hsu et al., 2004; Gleskova et al., 2004). Experimental findings of the mechanical behavior of the a-Si:H TFTs under externally applied strain are summarized in Fig. 3. Three different regimes have been identified. Under elastic deformation (which is the safe regime) the current-voltage characteristics of the TFT change reversibly (Gleskova and Wagner, 2001; Gleskova et al., 2002, 2004; Servati and Nathan, 2005). The electron mobility decreases linearly with compressive strain and increases with tensile strain. The mobility changes correlate with the broadening or steepening of the conduction band tail in a-Si:H channel material. These changes in the electron mobility are relatively small, for example, at the compressive strain of 1%, the mobility is reduced by 25%.

a-Si:H TFTs can be strained more in compression than in tension. No mechanical failure was observed in compression for strains up to $\sim 2\%$. In

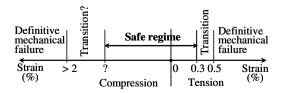


Fig. 3. Summary of the response of a-Si:H TFT on Kapton E to mechanical strain (Wagner et al., 2005).

tension, the crack formation starts at the strain of $\sim 0.3\%$ that marks the onset of the transition regime. In the transition regime the TFT fails but the electrical function is restored when the strain is eliminated (Gleskova et al., 2002). When the tensile strain reaches $\sim 0.5\%$ the cracks become permanent and definitive mechanical failure occurs (Gleskova and Wagner, 1999b). The cracks form perpendicularly to the strain direction. If the source-drain current path and the strain direction are parallel, the cracks interrupt the current path. This is schematically illustrated in Fig. 4.

One study suggested that the failure mechanism of the amorphous silicon-germanium solar cells is somewhat different from that of TFTs. The results of the bending of triple-junction solar cells on stainless steel substrate (Jones et al., 2002) are summarized in Fig. 5. No changes in the electrical performance, namely the short-circuit current J_{sc} , open-circuit voltage $V_{\rm oc}$, fill factor FF, and efficiency η , were observed for tensile strain up to $\sim 0.7\%$ and compressive strain up to $\sim 1.7\%$. For tensile strains larger than 0.7%, a gradual decrease in $J_{\rm sc}$, $V_{\rm oc}$, FF, and η occurred. However, even at the tensile strain of 2% (the largest strain applied) the solar cell efficiency was still equal to 50% of the original value. This behavior can be explained by a vertical current path that is parallel to the cracks as illustrated in Fig. 6.

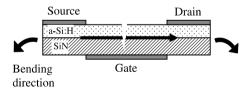


Fig. 4. Crack formation in a TFT when the strain and the sourcedrain current path are parallel. The arrow depicts the current path.

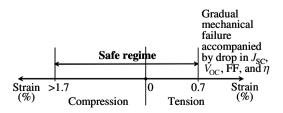


Fig. 5. Summary of the response of a triple-junction amorphous silicon-germanium solar cell on stainless steel to mechanical strain.

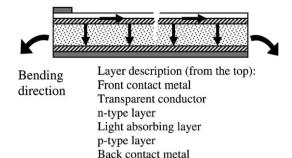


Fig. 6. Crack formation in a solar cell with respect to the current path.

5. Summary

The fabrication of thin-film devices on flexible substrates introduces several new fabrication issues that are not encountered during the fabrication of thin-film devices on rigid substrates. Strain that develops in the structure as a result of the built-in stresses in the deposited layers, or differences in the thermal expansion coefficients between the deposited films and the substrate, lead to a change in curvature of the work piece. Any change in the radius of curvature directly affects the misalignment between device levels and produces mask misalignment. To minimize the misalignment, one needs to carefully control the curvature of the work piece throughout the fabrication process.

Amorphous silicon solar cells and thin-film transistors respond differently to an externally applied strain. This difference is more apparent under tension. TFTs start to fail at a strain of ~0.3% and the failure is abrupt. Solar cells do not exhibit any change in the electrical performance for strains up to ~0.7%. If the tensile strain is further increased, the short-circuit current J_{sc} , open-circuit voltage V_{oc} , fill factor FF, and the efficiency η gradually decrease. The horizontal current flow in a TFT in contrast to the predominantly vertical current flow in a solar cell accounts for the difference.

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