

# Influence of residual stresses of sputtered thin film electrodes for dielectric elastomer applications

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**Abstract:** This paper focuses on the electromechanical properties of novel sub-micron compliant metallic thin film electrodes for dielectric elastomer membranes. Electrodes with thicknesses within the range 10-20 nm and different residual stress states are explored. Both pure nickel films and sandwiches of nickel (Ni) and carbon (C) are deposited by DC magnetron sputtering onto pre-stretched silicone elastomer membranes. Both 37.5 % biaxial pre-stretch and 57.5% uniaxial pre-stretch under pure shear condition (PSC) are considered in the conducted investigation. After the coating process is completed, the elastomer is allowed to relax. In the contracted configuration, it exhibits a wrinkled surface. After this state is reached, the electromechanical characterization is performed. All types of films reveal a low initial resistance (around 100  $\Omega$ /square). Depending on the kind of pre-stretch and the electrode material, a strain of 100 % without any major degradation is achieved. It is also shown how the residual stress of the layers can be influenced by suitable sputtering parameters. As a result, low residual film stress significantly improves the electromechanical properties of PSC pre-stretched elastomers, but have only minor influence on the biaxially pre-stretched ones, regarding the Ni and the Ni+C thin films. This phenomenon is directly connected to the failure mechanisms observed on the two types of pre-stretched membranes. With reversed layer order, i.e. C+Ni electrodes, the residual stress state of Ni does not influence the electromechanical properties for both, the biaxially pre-stretched and the PSC pre-stretched coated membranes. The results are of fundamental importance for understanding the role of residual stresses for the creation of electromechanically stable and highly conductive electrode films, to be used in DE applications.

**Keywords:** metallic electrode; compliant electrode; electroactive polymer; dielectric elastomer; sputtered thin film; wrinkles; pre-stretch;

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## 1. Introduction

Dielectric elastomer (DE) transducers represent a unique class of sensors and actuators that combine many interesting features, such as large deformation, lightweight, silent operations, scalability, and energy efficiency, amongst others [1,2].

A DE can be interpreted as a flexible capacitor consisting of a flexible polymeric membrane sandwiched by two compliant electrodes [3]. The application of a few kilovolts results in an attraction of the electrodes, accompanied by a simultaneous in plane expansion of the polymeric membrane. The electro-mechanical compressive stress induced on the membrane is described by the Maxwell stress  $\sigma_{Max}$ , given as follows:

$$\sigma_{Max} = \epsilon_r \epsilon_0 \left( \frac{V}{d} \right)^2, \quad (1)$$

with the relative permittivity  $\epsilon_r$ , vacuum dielectric constant  $\epsilon_0$ , voltage  $V$ , and distance between the electrodes  $d$  [4].

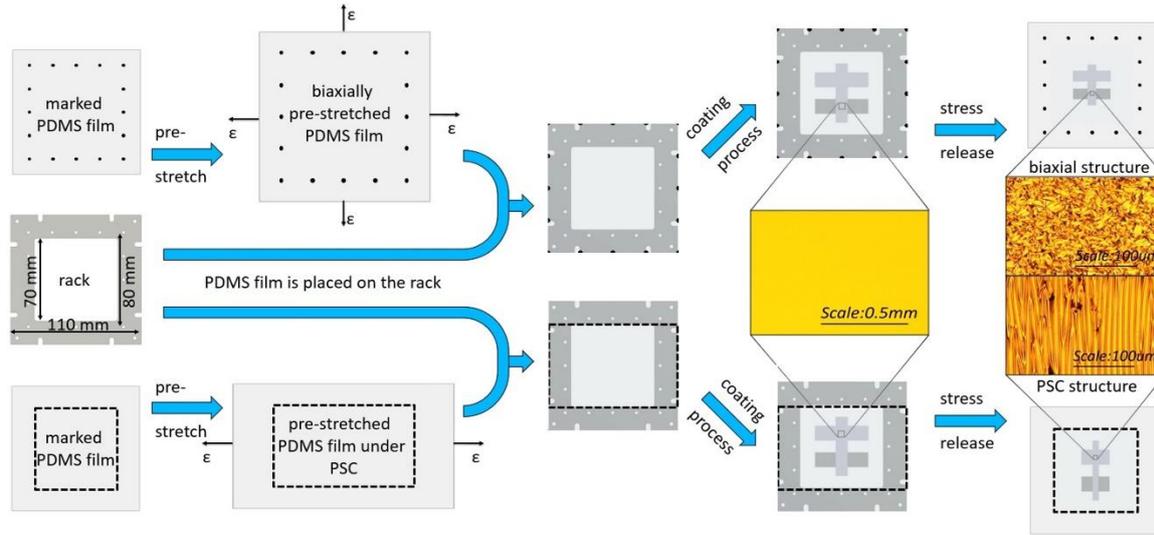
The electrodes are of crucial importance regarding the functionality of the DE. They have to be soft and highly conductive even at high strain levels up to 100 %. Furthermore, patternability and stretchability are required [5]. Usually, carbon black electrodes, i.e., a mixture of carbon and silicone, with a thickness of several  $\mu\text{m}$ , are used to obtain soft and stretchable electrodes. Such types of electrodes, however, exhibit a relatively high initial resistance of around 50 k $\Omega$  per square [6]. Metallic thin film electrodes with a thickness in the nm- range provide better conductivity in the order of 100  $\Omega$  per square, however they stiffen the system and have only limited elasticity. Scientists tried to overcome these drawbacks by creating corrugated [7] or wrinkled surfaces [8], by generating stretchable zig-zag [9], meander [10] or kirigami structures [11], or by applying sophisticated manufacturing methods such as ion implantation [12].

This study focuses on sputtered wrinkled metallic electrodes, a well-known principle in the field of flexible electronics [13,14]. In here, an either biaxially or under pure-shear condition uniaxially (PSC) pre-stretched silicone membrane is sputter coated with either a pure nickel thin film or a sandwich consisting of nickel and carbon with a thickness of 10 nm or 20 nm respectively. After the deposition process, the membrane is allowed to relax whereby a wrinkled metallic electrode is obtained. These wrinkles act as mechanical buffer when a strain is applied during actuation. Electromechanical tensile tests are performed to characterize the thin film electrodes. The influence of the residual stress state of the respective thin film electrode on the electromechanical properties of the DE is also clarified. The material selection of nickel and carbon is motivated by our own previous research on compliant electrodes [15] and sensor thin films [16,17].

## 2. Materials and Methods

In this work, Wacker Elastosil 2030 with a thickness of  $47.5 \mu\text{m} \pm 5\%$  is used as silicone membrane (PDMS membrane). The membrane is sputter coated in a pre-stretched state. Either a 37.5 % biaxial pre-stretch or a 57.5 % uniaxial pre-stretch under pure-shear conditions (PSC) are considered in this paper. As soon as the pre-stretch is applied, the silicone film is placed on a metal rack, dedusted with deionized air, and then coated in the vacuum chamber. A mirror-like surface is obtained after the sputtering. By carefully removing the silicone from the rack after the coating, the membrane is allowed to relax. It shrinks to its initial dimension, thus forcing the metallic thin film to form wrinkles. Depending on the type of pre-stretch, an aligned wrinkle structure (PSC pre-stretch) or a random wrinkle orientation (biaxial pre-stretch) is obtained (cf. **Figure 1**).

The coating is performed in a  $45 \times 45 \times 45 \text{ cm}^3$  vacuum chamber from the company CCR. The chamber is equipped with a turbo molecular pump, a downstream throttle valve (VAT), and two heads for magnetron targets with a diameter of 127 mm. One head carries a nickel target, while the



**Figure 1:** Process steps of manufacturing the dielectric elastomer membranes: a pre-stretched silicone membrane is fixed on the metal frame and then coated. Afterwards, the membrane is allowed to relax, and a wrinkled surface is achieved.

other one a carbon target. The samples are placed with a sample to target distance of 45 mm on a movable carrier. The chamber has no load lock, hence it is vented with nitrogen after every process. Once the vacuum chamber is evacuated to a background pressure less than  $1 \times 10^{-5}$  mbar, three purge-pump cycles with argon to a pressure of  $1 \times 10^{-1}$  mbar are executed, to ensure clean sputtering conditions. Then, argon (purity 99.999 %) is let into the vacuum chamber with a constant flow of 15 sccm. The process pressure is controlled in the range between 1.5  $\mu$ bar and 36  $\mu$ bar, thanks to appropriate setting of the throttle valve. When the process pressure is adjusted, the first target is pre-sputtered for 1.5 minutes with 300 W DC power. Then, the sample is transferred under the target for the coating process. After a certain process time, the sample is removed again. If a sandwich layer is desired, the procedure is subsequently executed in the same manner at the second target.

The thin films are deposited on both surfaces of the PDMS, by applying cross-shaped shadow masks rotated against each other by 180 degrees (see **Figure 2**). The long cross bar (10 mm  $\times$  50 mm) represents the active area to be examined, while the perpendicular bars serve as electrical contacts. Three types of electrodes are researched in this study. Pure nickel thin films with a thickness of 10 nm or a bi-layer sandwiches, with an overall thickness of 20 nm, consisting of carbon (10 nm) and Ni (10 nm), are manufactured. Both electrode layer sequences, Ni+C as well as C+Ni, are investigated in this study. Due to different deposition rates of nickel and carbon, the 10 nm thick Ni-layer corresponds to a sputter time of 5 sec, whereas 60 sec is needed to achieve a 10 nm thick carbon layer. Further details on the manufacturing process are given in [15].

The residual stress state of the deposited thin films is examined on glass slides. Glass slides are sputter coated, and the curvature of the slides is measured before and after the coating process. The residual stress of the deposited thin film can then be calculated with the aid of the Stoney equation [18,19]:

$$\sigma_R = \frac{E_s}{6(1-\nu_s)} \frac{h_s^2}{h_f} \left( \frac{1}{R} - \frac{1}{R_0} \right), \quad (2)$$

with the Young's Modulus of the substrate  $E_s$ , the Poisson ratio of the substrate  $\nu_s$ , the thickness of the substrate  $h_s$ , the thickness of the film  $h_f$ , the initial radius of the glass slide  $R_0$ , and the radius of the glass slide after the sputter coating  $R$ .

The radius of the glass slides is determined by means of a white light profilometer *Microprof* from the company *FRT*. The surface profile of the slides is measured with a line scan. Then, the

software fits a section of a circle to the 2D profile, and calculates the corresponding radius of curvature. Overall, nickel layers produced at 13 different process pressures in the range between 1.5  $\mu\text{bar}$  and 36  $\mu\text{bar}$  are investigated. The sputter time of 45 sec, the power of 300 W DC, and the sample to target distance of 45 mm are kept constant for this series of experiments. For each pressure level, the residual stress is calculated and plotted as a function of the process pressure. A fit of the measured data is then performed with *OriginLab*.

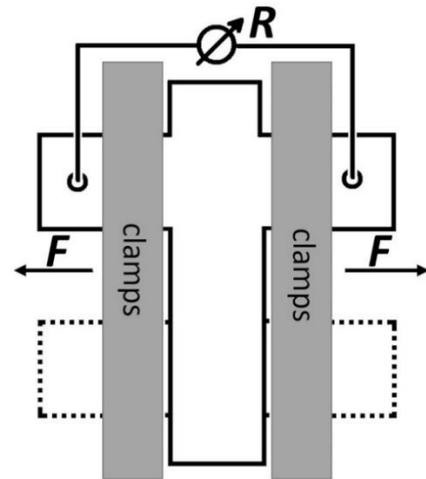
A direct thickness measurement of the thin film electrode on the soft and transparent PDMS is not possible. Hence, the thickness of the layers is determined by means of the deposition rate of the respective target materials. A sputter deposited thin film of either carbon or nickel is created on glass slides, with a small cover glass in the center to produce a step with the thickness of the film. The deposited thin film is produced with the same sputter conditions as those used for manufacturing the DE membranes. Then, this step is measured ascending and descending with a *Dektak 150* tactile profilometer. The deposition rate is calculated from the step height and the corresponding sputter time. The associated linear fit is done by *OriginLab*. For carbon, a series of coatings with sputter times between 0 and 5 min are performed at a process pressure of 1.5  $\mu\text{bar}$ . For nickel, sputter times between 0 and 2 min were chosen with pressures of 1.5  $\mu\text{bar}$  and 18  $\mu\text{bar}$ , respectively.

Both PSC and biaxially pre-stretched membranes, coated on only one side with either Ni, Ni+C or C+Ni thin film electrodes, are tested on a uniaxial tensile tester. For each type of electrode, both electrodes with nickel thin films deposited at high and low process pressure are manufactured and examined. The schematic test arrangement is shown in **Figure 2**. (for a detailed description of the tensile tester, the author may refer to [20]). With the tensile direction being perpendicular to the long side, and a length to width ratio of the active area equal to 5:1, the uniaxial tensile test can be interpreted as a pure shear tensile test [21]. The transverse contraction is negligible. During the tests, a LCR meter (*Hameg MH8118*) is used to measure the resistance versus the applied strain. Starting at 0 % strain, the strain level is increased stepwise with a complete strain release between the steps. Every strain level is kept constant for 25 seconds, and the electrical resistance is recorded and averaged over time. The averaged resistance is then plotted as a function of the corresponding strain. An increment of 10 % is chosen up to an absolute strain of 30 %, followed by an increase of 5 % up to the electrical failure (defined to be 100 k $\Omega$  in this study). Two smaller steps of 2.5 % strain increments are also included around the level of pre-stretch.

### 3. Results and discussion

#### 3.1. Deposition rate

The result of the deposition rate analysis is shown in Figure 3. The deposition rate of carbon is 0.16 nm per second, so 10 nm of a thin carbon film is produced in about one minute. Nickel layers sputtered at two very different process pressures (1.5  $\mu\text{bar}$  and 18.1  $\mu\text{bar}$ ) have comparable deposition rates of 2.55 nm/sec and of 2.59 nm/sec, respectively. In both cases, although they differ greatly in their residual stresses (see below), the actual thickness of nickel thin films produced in five seconds is slightly lower than the theoretical thickness calculated by the deposition rate. This is related to the process control during the manufacturing (for further details see [15]). In general, the



**Figure 2:** Tensile test arrangement.

deposition rate is dependent from several parameters such as the process pressure, the target to substrate distance, the vacuum chamber, and the deposition power, amongst others [22–24]. It increases first with a higher process pressure, due to a rise of the number of ions [24]. With a further increase of the process pressure, the probability of an expelled atom to reach the substrate decreases due to interatomic collisions and back diffusion [22,24]. These two competing processes could be responsible for the obtained results, i.e., the deposition rate for both Ni-processes are hardly distinguishable.

### 3.2 Residual stress analysis

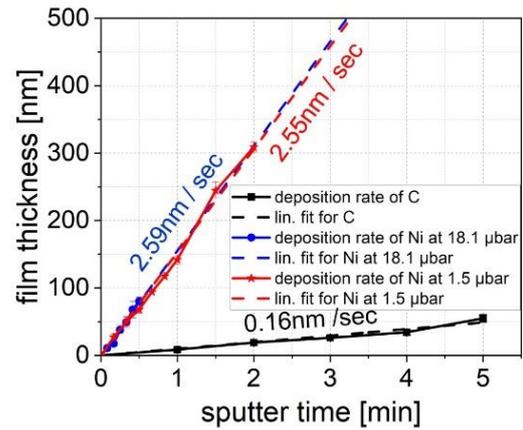
In **Figure 4**, the measured data of the residual stress analysis and the corresponding fit is plotted as a function of the process pressure. All measured stresses are tensile residual stresses. These curves are shaped similarly to a concave parabola, which starts with a local minimum of 75 MPa at around 1.5  $\mu\text{bar}$ , followed by an increase of the residual stress up to 917 MPa at 18  $\mu\text{bar}$ . Then, the residual stress drops again to 208 MPa at 36  $\mu\text{bar}$ . These results are based on equation 2 with a Young's modulus (of glass) of 64 GPa [25], a Poisson ratio of 0.2 [25], a thickness of the glass slides of 0.5 mm, and a film thickness of 115 nm. The difference of film thickness between a Ni (low stress) and a Ni (high stress) layer due to the different deposition rate is 2 nm. For the interpretation of the residual stress state of the examined thin films, this thickness deviation is negligible.

Ni (high stress) layers have a good adhesion on glass slides, as shown in previous tests. Hence, for the examination of the series of nickel electrodes, 18.1  $\mu\text{bar}$  was first chosen as process pressure. It could be observed, however, that the Ni (high stress) layers tend to form cracks when deposited onto PSC pre-stretched membranes. By depositing Ni (low stress) electrodes, sputtered at 1.5  $\mu\text{bar}$  on silicon membranes, the crack formation can be avoided completely. However, the nickel thin film manufactured at 36.3  $\mu\text{bar}$  is still prone to crack formation on silicone. Based on this observation, the residual stress study focuses on the Ni (low stress) films deposited at low process pressures.

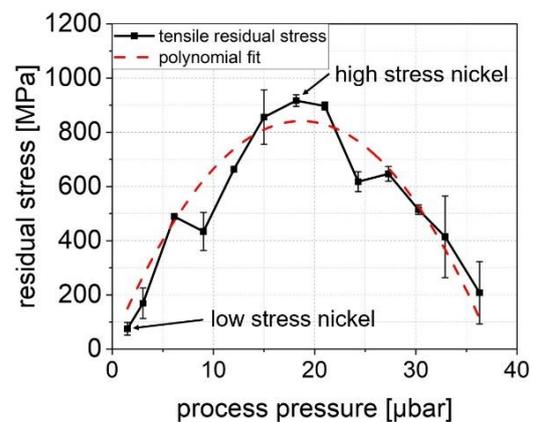
Regarding the wrinkle formation after the relaxation of the electrodes with low stress nickel and high stress nickel, no significant optical differences can be observed.

### 3.3 Resistance vs. strain measurements

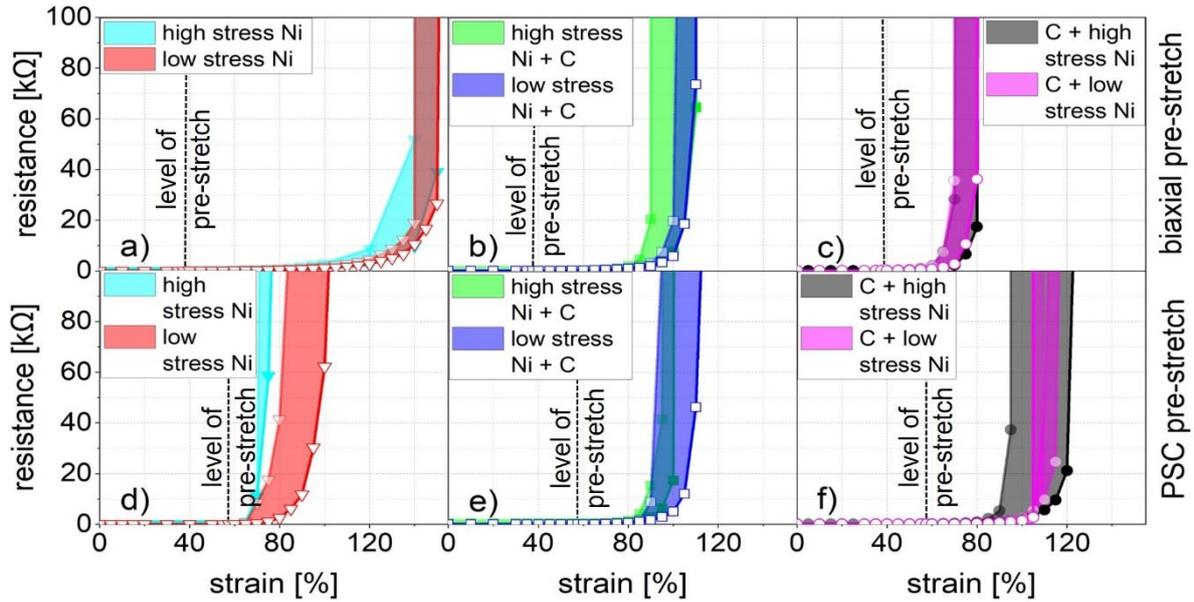
The resistance versus strain for Ni, Ni+C, and C+Ni thin film electrodes, deposited on either biaxially pre-stretched or PSC-pre-stretched membranes, is illustrated in **Figure 5**. In the first row (**Figure 5 a, b and c**), the results of the biaxially pre-stretched silicone membrane are shown,



**Figure 3:** Deposition rate analysis of C and Ni is showed here. The deposition rate of C is 0.16 nm/s (black curve). The rate of Ni is 2.55 (red) and 2.59 nm/s (blue curve), deposited at the different pressures of 1.5  $\mu\text{bar}$  and 18.1  $\mu\text{bar}$  respectively.



**Figure 4:** Tensile residual stress versus process pressure (black curve) and the corresponding polynomial fit.



**Figure 5:** Resistance measurements vs. strain results. Biaxially pre-stretched (a,b,c) and PSC pre-stretched membranes (d,e,f) are coated with different types of thin film electrodes. Each type of thin film is deposited with high and with low residual stress and the results are compared graphically. Every colored cluster represents at least three specimens.

whereas PSC pre-stretched silicone is represented in the second line (**Figure 5 d, e and f**). Each column depicts the results of one type of thin film, starting from pure nickel (**Figure 5 a and d**), followed by Ni+C (**b and e**), and then C+Ni (**c and f**), when moving from left to right. Each graph contains two colored areas, representing the data family of at least 3 specimens of the corresponding thin film type. The maximum and the minimum values of the series of the examined type of electrode frames the colored area.

Overall, there are some characteristics that are independent of the residual stress state. For all layer combinations, the threshold of the resistance increase is located at strain levels above the respective level of pre-stretch (LoP). For the biaxially pre-stretched membranes, this threshold is shifted to higher strain levels with respect to the corresponding LoP. Furthermore, the resistance increase on biaxially pre-stretched membranes is more gradual, while electrodes deposited on PSC membranes show a fast and steep increase of the resistance above the LoP. This phenomenon is related to the difference of the failure mechanisms of the two types of pre-stretch. If a low resistance is needed at high strain levels, a pure Ni thin film can be used on biaxially pre-stretched membranes, whereas on PSC membranes the C+Ni electrodes gives the best results. These results obtained here for Ni layers correspond to the trends of the previous NiCr study [15].

From the results, it can be concluded that both the different thin film configuration and the type of pre-stretch are influenced in different ways by the residual stress state of the electrodes. Ni and Ni+C reveal the same characteristics. On foils with a biaxial pre-stretch, the residual stress state of the electrode does not influence the resistance curve at high strain levels. The width of the colored region is narrowed for the low residual stress specimen. The maximum curves of both the low and the high residual stress layer are congruent, though. When a low stress layer is manufactured onto membranes pre-stretched under PSC conditions, the threshold of the resistance increase can be shifted towards higher strain levels.

The different failure mechanism of the biaxially and PSC pre-stretched membranes is responsible for the different results. When the biaxially pre-stretched membrane is allowed to relax after the coating, a wrinkled surface with a random wrinkle orientation is obtained. A PSC membrane exhibits aligned wrinkles after relaxation. During the strain tests, this aligned structure is unfolded

and flattened. When the LoP is reached, a perfectly flat electrode surface is obtained in theory. By exceeding the LoP cracks in the metallic electrode can be created, which can easily propagate through the flat surface. Hence, few but large cracks are responsible for the drastic resistance increase. In this case, the intrinsic residual stress of the electrode directly influences the driving force for the crack formation. A high residual tensile stress promotes crack formation and propagation. With low residual stress, on the other hand, crack formation is delayed and the threshold is shifted to higher levels of strain. A biaxially pre-stretched membrane is only partially flattened during a uniaxial tensile test. Even at strain levels above the LoP, wrinkles parallel to the strain direction are still present. The remaining wrinkled structure leads, during a tensile test, to an inhomogeneous stress distribution over the surface. Once a crack is created, it is not able to propagate through the whole electrode in one step, since it stops when it crosses wrinkles oriented perpendicularly to the crack growth direction. Hence, a huge number of small cracks is present. In general, this is advantageous when a low resistance at high strain is desired [26]. This phenomenon of intrinsic impediments of crack propagation was shown before on coated surfaces with different roughnesses [27]. The results of the mentioned study could easily be assigned to a randomly wrinkled surface. For biaxially pre-stretched membranes, the failure mechanism dominates over the residual stress state.

A Ni+C electrode consisting of a nickel layer with low stresses on PSC pre-stretched silicone exhibits an degradation at strain levels 10% higher, compared to the Ni+C electrode with high residual stresses (see **Figure 5 d**). Regarding the pure nickel electrodes on membranes with a PSC pre-stretch, replacing the high stress nickel by a low stress nickel thin film provides an enhancement of almost 30 % (compare **Figure 5 e**). In our study, only the residual stress of the Ni layer is investigated. The properties of the carbon layers are the same for all measured sandwich electrodes. The performance of the Ni+C electrode is determined by the interaction of both thin film layers, and not only by the nickel thin film. Therefore, the change in the properties of sandwich Ni+C layer is not as obvious as the drastic change of a pure nickel electrode.

If Ni is deposited as a top layer on a carbon layer, i.e., a C+Ni sandwich (**Figure 5 c and f**), then the residual stress of the nickel layer does not influence the resistance at high strain levels. The carbon bottom-layer somehow absorbs the high residual stress of the nickel layer. In that case, a reduction of the residual stress of the nickel top layer does not lead to any further improvement, but also to no degradation of the electromechanical properties. The exact mechanism of how carbon works has not been clarified yet, and will be part of the investigation for future research.

#### 4. Conclusion

Based on the conducted study, it can be concluded that the reduction of residual stresses of a thin film electrode can improve, under certain circumstances, the electromechanical properties of a DE membrane. For biaxially pre-stretched membranes tested under pure shear conditions, the resistance increase is solely dominated by the failure mechanism of the randomly oriented wrinkles, and not by the residual stress state of the compliant electrode. Regarding the electromechanical properties, neither benefits nor drawbacks are observed in case of low -stress nickel layers. The same is true for sandwich layers if Ni is the top-layer. In that case, the carbon bottom layer absorbs the residual stresses of the top-layer. For membranes, both pre-stretched and tested under pure shear conditions, the residual stress reduction of the nickel layers results in a shift of the resistance increase towards higher strain levels. The strain threshold is thus enhanced by almost 30 % for pure nickel electrodes. Thus, electrodes with low residual stress states offer the chance of improving the electromechanical properties. Furthermore, the process parameters which are applied to obtain films with low stresses, avoid crack formation during sputtering. Hence, it is highly recommended to use thin films with low residual stresses for all kind of DE applications.

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