Pentacene Organic Thin Film Transistors with Anodized Gate Dielectric

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ABSTRACT

A low temperature high quality gate dielectric process for bottom gate organic thin film transistors (OTFT) is introduced which is compatible to plastic substrates. The Al\textsubscript{2}O\textsubscript{3} dielectric is grown from the aluminum gate electrode by anodic oxidation at room temperature and exhibits an exceptionally good electrical performance even for thin layers of 50 nm. Finding an electrolyte which significantly reduces dielectric charges was instrumental for the desired OTFT application. The electrolyte and substrate dependent behaviour was characterized and compared to different dielectrics to point out the advantages of anodic oxidized aluminum. The characteristics of pentacene bottom contact OTFTs realized with anodized Al\textsubscript{2}O\textsubscript{3} gate dielectric on glass and plastic substrates are presented.

Keywords: Organic Thin Film Transistors, Pentacene, Gate Dielectric, Anodisation

1. INTRODUCTION

Thin film transistors (TFT) with organic semiconducting material instead of silicon as active layer have a big potential for flexible display technologies. These so called organic thin film transistors (OTFT) can be used in a wide range of applications where flexible materials and low cost are required. In contrast to silicon, the thermal, structural and mechanical properties of organic semiconductors are similar to those of flexible plastic substrates. Additionally, organic semiconductors are deposited with low temperature processes. Since the maximal thermal load for many low cost substrates is less than 200°C, such low temperature processes are essential. One disadvantage of OTFTs is the low mobility resulting from weak Van der Waal bonds. These Van der Waal bonds dominate in organic semiconductors in contrast to the covalent bonds dominating in conventional semiconductors.

Developing a process for OTFTs on flexible plastic substrates requires low temperature processes for all layers. For conventional display applications SiO\textsubscript{2} and Si\textsubscript{3}N\textsubscript{4} are used as dielectric layer. Since these insulating materials are deposited at around 280°C they have to be replaced by low temperature alternatives. Finding a low temperature dielectric with good electrical properties is one of the most important steps in developing OTFTs.

One basic requirement for a switching TFT in an active matrix addressed display is a high on-current $I_{\text{ON}}$ and a low off-current $I_{\text{OFF}}$. A high $I_{\text{ON}}$ is required to allow fast switching of the pixel and a low $I_{\text{OFF}}$ to avoid discharging of the pixel in its OFF-state. For a low leakage current through the gate dielectric, a high quality insulating layer with a low defect density must be used.

The influence of the mobility $\mu$, the gate capacity $C_{\text{GS}}$, the width $w$ and length $l$ of the transistor channel, the voltage between gate and source $V_{\text{GS}}$ and threshold voltage $V_{\text{th}}$ on $I_{\text{ON}}$ is given by

$$I_{\text{ON}} = \mu C_{\text{GS}} \frac{w}{2l} (V_{\text{GS}} - V_{\text{th}})^2$$

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The mobility $\mu$ and thus the $I_{ON}$ is limited by the weak van der Waal bonds. It is possible to decrease the operating voltage $V_{GS}$ and the threshold voltage $V_{th}$ by using a dielectric with a high specific gate capacitance $C_{GS}$. From

$$C_{GS} = \frac{\varepsilon_0 \varepsilon_r}{d}$$  \hspace{1cm} (2)

with $\varepsilon_0 = 8.85 \cdot 10^{-12}$ As/Vm can be seen that $C_{GS}$, can be increased by selecting a dielectric with a high dielectric constant $\varepsilon_r$ and a low thickness $d$. To achieve dielectric layers with a high $C_{GS}$ different attempts can be found in literature.\textsuperscript{2-4} The properties of oxides which are grown from Aluminum or Tantalum by anodic oxidation are suitable as dielectric layers in OTFTs since thin layers of about 50 nm can be reliably produced having $\varepsilon_r = 9$ and $\varepsilon_r = 27$ respectively. Anodic oxidation is a low temperature process which is important for the use with plastic substrates. Additionally, our experiments showed that the surface roughness decreased during the process. The surface of the anodized metals is smoothed and the defect density is reduced. A low surface roughness is important for the growth of evaporated organic semiconductors.\textsuperscript{5}

2. EXPERIMENTS AND RESULTS

Different dielectric materials are investigated for their compatibility in organic thin film transistors. Two different organic and four inorganic materials are investigated. The organic dielectrics, a polycrystalline and a polyimide, are deposited by spin coating. The inorganic materials are plasma enhanced chemical vapour deposited (PECVD) Si$_3$N$_4$ and SiO$_2$ layers which have been developed for polysilicon TFTs\textsuperscript{6} and anodically oxidized Ta$_2$O$_5$ and Al$_2$O$_3$.

In this work the transfer of OTFT processes to plastic substrates is important. Therefore a low temperature dielectric with good insulating properties is needed. Spin coating and anodic oxidation are low temperature processes. The deposition temperature for the PECVD SiO$_2$ and Si$_3$N$_4$ layers is 280 °C and is not suitable for the application on plastic substrates. But they are useful to compare the proved dielectric properties with those of the low temperature ones. Since the anodized oxides fullfill the above mentioned requirements the anodization process is discussed in more detail.

2.1. Anodisation process

Figure 1 shows schematically the anodization process.
The substrate with a structured metal layer of aluminum or tantalum is used as the anode and a metal plate as the cathode in an electrolyte bath. The electrolyte must be electrically conductive and for the aluminum layer the pH-value must be between 6 and 8 otherwise the aluminum layer is dissolved. During anodisation the current density is kept constant while the voltage increases linearly with the oxide thickness. Metal ions migrate onto the surface of the metal and react with $OH^-$ anions of the electrolyte. One part of the oxide grows into the metal and the other grows on the metal. The voltage defines the thickness of the oxide. When the desired thickness is reached, the voltage is kept constant so that the current density drops exponentially. Ethyleneglycol and $H_2O_2$ buffered with ammoniumhydroxid to obtain a pH-value of 7 are used as electrolyte to form $Al_2O_3$. Citric acid and $H_2O_2$ are used to grow $Ta_2O_5$. A varity of different possible electrolytes can be found in literature.\(^7\)\(^8\)

2.2. Characterisation of the dielectric

Capacitors are first built on glass substrates to characterize and compare the different materials. The current density is measured while increasing the electrical field strength linearly by \(\frac{dE}{dt}\) = 25 kV/cm·s. The resulting total current density \(j_{\text{tot}}\) consists of the addition of the displacement current \(j_{\text{dis}}\) and the leakage current \(j_{\text{leak}}\) according to

\[
j_{\text{tot}} = j_{\text{dis}} + j_{\text{leak}} = \varepsilon_r\varepsilon_0 \frac{dE}{dt} + j_{\text{leak}}.
\]

In this equation \(\varepsilon_r\) and \(\varepsilon_0\) are the dielectric constants for the material and the vacuum respectively. In the resulting graph the constant part of the curve is due to \(j_{\text{dis}}\) which depends on the dielectric material constant and the electrical field strength ramp. The slope results from \(j_{\text{leak}}\). The breakthrough field strength is defined as the value when the current density reaches \(j_{\text{tot}} = 10^{-6} \text{A/cm}^2\).

The measured current densities over the electrical field strength of the different materials are presented in figure 2. Since the maximum voltage \(U_{\text{max}}\) of the used picoamperemeter is limited to \(U_{\text{max}} = \pm 100\) V, the maximum applied electrical field strength \(E_{\text{max}} = U_{\text{max}} / d\) for each capacitor depends on the dielectric thickness \(d\). From figure 2 it can be seen that the current density for the spin coated polyacryl and polyimide increases strongly with the applied electrical field. For $SiO_2$ and $Si_3N_4 j_{\text{tot}}$ is influenced only by \(j_{\text{dis}}\) for a wide area before the breakthrough occurs. Both anodically oxidized dielectrics $Al_2O_3$ and $Ta_2O_5$ show an unsymmetry between positive and negative field strength. While a definite breakthrough can be observed in the negative region it is not well defined for positive field strength. The definite breakthrough in the negative region can be related to the direction of the applied electrical field strength, which is the same as during the oxidation process. We assume that negative anions in the oxide are causing the undefined breakthrough for positive field strengths.

Table 1 summarizes the thicknesses, the dielectric constants \(\varepsilon_r\), the breakthrough field strengths \(E_d\) and the average roughness \(R_a\) determined by atomic force microscope (AFM) measurements for the investigated materials. The average roughness of anodically oxidized dielectrics is higher than for the spin coated and for $SiO_2$ layers but it is comparable to the roughness values measured on $Si_3N_4$. The anodically oxidized materials have been further investigated due to their electrical performance, the low deposition temperature, the high dielectric constants and the possibility to form thin layers which reduces the required OTFT operating voltages.

<table>
<thead>
<tr>
<th>material</th>
<th>thickness [nm]</th>
<th>(\varepsilon_r)</th>
<th>$E_d^+ [\text{MV/cm}]$</th>
<th>$E_d^- [\text{MV/cm}]$</th>
<th>roughness (R_a) [pm]</th>
</tr>
</thead>
<tbody>
<tr>
<td>$Al_2O_3$</td>
<td>60</td>
<td>9</td>
<td>5&gt;</td>
<td>6</td>
<td>935</td>
</tr>
<tr>
<td>$Ta_2O_5$</td>
<td>50</td>
<td>27</td>
<td>2&gt;</td>
<td>4.5</td>
<td>1150</td>
</tr>
<tr>
<td>$SiO_2$</td>
<td>100</td>
<td>3.9</td>
<td>8&gt;</td>
<td>10&gt;</td>
<td>667</td>
</tr>
<tr>
<td>$Si_3N_4$</td>
<td>200</td>
<td>7.5</td>
<td>7&gt;</td>
<td>7&gt;</td>
<td>978</td>
</tr>
<tr>
<td>polyimide</td>
<td>400</td>
<td>3.9</td>
<td>&lt;3</td>
<td>3</td>
<td>561</td>
</tr>
<tr>
<td>polyacryl</td>
<td>400</td>
<td>3.9</td>
<td>&lt;4</td>
<td>3</td>
<td>521</td>
</tr>
</tbody>
</table>

Table 1. Thickness, dielectric constant, breakthrough field strength and surface roughness of investigated dielectrics.
To improve the electrical and structural properties of anodically oxidized dielectrics, different electrolytes are investigated. The results for the electrical performance are summarized in table 2 and in figure 4(a). Characteristic anions which are included in the ethyleneglycol and aqueous citric acid electrolytes are supposed to be incorporated in the dielectric during oxidation and negatively influence the breakthrough performance. By using H$_2$O$_2$ as electrolyte the electrical performance of both Al$_2$O$_3$ and Ta$_2$O$_5$ is improved especially for positive field strengths. Beside that the average roughness $R_a$ is decreased with the use of H$_2$O$_2$.

![Figure 2](image-url)

**Figure 2.** Measured total current density $j_{tot}$ depending on the electrical field strength of different dielectric materials.

<table>
<thead>
<tr>
<th>material</th>
<th>electrolyte</th>
<th>thickness [nm]</th>
<th>$E_d^+ [MV/cm]$</th>
<th>$E_d^- [MV/cm]$</th>
<th>$R_a [pm]$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Al$_2$O$_3$</td>
<td>ethyleneglycol</td>
<td>60</td>
<td>$\approx 3$</td>
<td>5.5</td>
<td>1570</td>
</tr>
<tr>
<td>Al$_2$O$_3$</td>
<td>H$_2$O$_2$</td>
<td>60</td>
<td>$\approx 5$</td>
<td>6</td>
<td>935</td>
</tr>
<tr>
<td>Ta$_2$O$_5$</td>
<td>citric acid</td>
<td>100</td>
<td>0.1</td>
<td>0.5</td>
<td>1150</td>
</tr>
<tr>
<td>Ta$_2$O$_5$</td>
<td>H$_2$O$_2$</td>
<td>50</td>
<td>2</td>
<td>4.5</td>
<td>997</td>
</tr>
</tbody>
</table>

**Table 2.** Thickness and breakthrough field strength depending on the electrolyte.

### 2.3. Kiessig Interference measurements

The structure of Al$_2$O$_3$ is examined by Kiessig Interference measurements. For this measurement the substrate is radiated with a x-ray under small angles of incidence. The x-ray penetrates into the materials and is partly reflected at the layer interfaces. The thickness of the different layers is obtained from the distance between two of the occurring interferences by using equation 4 wherein $d$ is the thickness and $q_{\perp i}$ and $q_{\perp i-1}$ are the orthogonal wave vectors of two proximated interference maxima:

$$d = \frac{2 \cdot \pi}{q_{\perp i} - q_{\perp i-1}}.$$ (4)

The damping of the curve depends on the surface roughness of the investigated layers.
As shown in figure 3 the structural performance of Al₂O₃ is improved with the use of H₂O₂. From the reduced damping of the oscillation for the H₂O₂ electrolyte it can be concluded that the smoothness of the surface and the transition region between the metal and the oxide are enhanced. The thickness d of the oxide layer can be extracted with equation 4. It is d = 65 nm for the Al₂O₃ grown in the H₂O₂ electrolyte and d = 57 nm when using Ethylenglycol. Both Al₂O₃ layers are not stoichiometric as it can be concluded from the maxima and minima positions.

![Kiessig oscillation of Al₂O₃ samples using different electrolytes.](image)

**Figure 3.** Kiessig oscillation of Al₂O₃ samples using different electrolytes.

### 2.4. Transfer to plastic substrates

After optimizing the process for Al₂O₃ on glass substrates, it was transferred to a polyethersulphone (PES) plastic substrate. The anodisation of aluminum is chosen as the first process to be transfered on plastic substrate since the bandgap, which influences the intrinsic breakdown, of Al₂O₃ is 8.8 eV in comparison to Ta₂O₅ where it is 4.4 eV.⁹ Additionally, aluminum can be structured by wet-etching while plasma etching processes required for tantalum are affecting the plastic. Thus none of the processes required for fabricating capacitors with Al₂O₃ dielectric on plastic (sputtering and structuring the metals, anodisation with electrolytes) has a visible detrimental effect on the plastic substrates. The electrical performance of capacitors with Al₂O₃ as insulator are measured on glass and plastic substrates. The results are shown in figure 4(b). The breakdown behaviour is not influenced by the substrate material. The leakage current determined by the slope of the current density is even less than on the glass substrate. The structure of the anodized films on both substrates are analysed again with Kiessig interference. The resulting characteristics are shown in figure 5.

The resulting thickness extracted from the oscillation caused by the Al/Al₂O₃ interface is d = 63 nm on the plastic substrate. The thickness is not influenced by the substrate material. From the higher damping of the curve measured on plastic substrate it can be concluded that the surface of the different layers on the plastic substrate are rougher than on glass. The less defined absorption edge measured on the plastic substrate is related to fixture problems resulting in a not perfectly flat substrate during the measurement.

### 2.5. AFM measurements

Since the surface roughness influences the growth of the following layers, it is examined with an atomic force microscope (AFM). Figure 6(a) and 6(b) show the resulting picture of Al₂O₃ on glass and on plastic substrates. While small grains are visible on the Al₂O₃ surface on plastic the surface on glass is smooth. This indicates
different growth mechanisms for the sputtered aluminum layer on PES and glass. The resulting values for the average surface roughness $R_a$ of $\text{Al}_2\text{O}_3$ are $R_a = 935 \text{ pm}$ on glass and $R_a = 1610 \text{ pm}$ on plastic (table 3). The difference in the surface roughness corresponds to the results of the Kiessig interference measurements (figure 5).

2.6. OTFT performance

The $\text{Al}_2\text{O}_3$ is used as the dielectric layer in organic thin film transistors (OTFT). OTFT structures are fabricated on glass and plastic substrates. Bottom gate, bottom contact OTFTs are processed using aluminum as gate metallization. The surface of the structured aluminum is anodized forming the oxide layer. As drain- and source contact material gold is sputtered and patterned. Finally, pentacene is evaporated as semiconducting layer. The cross section of the fabricated OTFTs is presented in figure 7. For the OTFTs a channel length of $l = 5 \mu \text{ m}$ and $l = 10 \mu \text{ m}$ is chosen and the channel width is varied between $w = 50 \mu \text{ m}$ and $w = 850 \mu \text{ m}$. The mobility $\mu$
and the threshold voltage $V_{th}$ are extracted from saturation region of the transfer characteristic using equation 1. The ON/OFF ratio is based on the maximum and the minimum value of $I_{DS}$. The resulting values of the OTFT on plastic and glass substrates are compared with each other.

The distortion which occurs during the process on plastic substrates is reduced by two tempering steps, one before the first layer has been deposited and the other one after the anodisation. The first tempering step is for the preshrinking of the foil to reduce film stress between the aluminum layer and the substrate. The second one is to drive out the moisture which gets into the substrate during the anodic oxidation.

The resulting transfer characteristic of OTFT on glass and plastic substrates are presented in figures 8(a) and 8(b). The extracted parameters for the mobility $\mu$, the threshold voltage $V_{th}$ and the ON/OFF ratio of the presented characteristics are shown in table 3.

The electrical performance of these OTFTs on glass and plastic substrates are comparable. Remarkable is the excellent blocking behaviour resulting in low off-currents $I_{OFF} < 1 \cdot 10^{-12}$ A for both substrates. $I_{OFF}$ is not influenced by leakage currents through the optimized gate dielectric Al$_2$O$_3$. The mobility and the ON/OFF ratio are not influenced by the substrate material as it can be seen from the values in table 3. The values are uniformly distributed over the 2 inch test substrates and they are reproducible. Another interesting effect is the threshold voltage shift when measuring from positive to negative gate voltages followed by a reverse sweep. This effect occurs for both substrate materials but is more distinct on plastic. This shift depends on the value of the most negative gate voltage which is applied to the OTFT. Reasons for this shift could be the interface defects.
between the dielectric and the pentacene. Since the surface of Al₂O₃ on plastic substrate is rougher than on glass this would explain the more pronounced shift for OTFTs on plastic substrates. Similar effects in OTFT and different explanations for this can be found in literature.¹⁰,¹¹ These effects have to be investigated in more detail for a better understanding and further improvements of OTFTs.

3. CONCLUSIONS

Different low temperature gate dielectrics are investigated and compared to conventionally used TFT gate dielectrics SiO₂ and Si₃N₄ to find a convenient gate dielectric for OTFT application on plastic substrates. Anodically oxidized Al₂O₃ is found to fulfill all requirements. Before the process is transferred to plastic substrates the influence of different electrolytes on the electrical and structural properties of the dielectric is investigated. Using breakthrough and x-ray measurements it has been shown that buffered H₂O₂ improves both properties. After the process is successfully transferred to plastic substrates, OTFTs on PES are built which show similar electrical behaviour as those on glass substrates.

ACKNOWLEDGMENTS

This project has been partially funded by the EU funded programme FlexiDis (Contract IST-2004-4354) and the DFG focus program (project No. PF385/2-3).

The authors thank Stefan Hirschmann from the 3rd Institute of Physics, University of Stuttgart for the purification of pentacene.

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