A new reactive sputtering technique for the low temperature deposition of transparent light emitting ZnS:Mn thin films

Steve Wakeham, Mike Thwaites, Costas Tsakonas, Wayne Cranton, Robert Ranson, Gabriel Boutaud and Demosthenes Koutsegeorgis

1 Plasma Quest Limited, Unit 1B Rose Estate, Osborn Way, Hook, UK
2 Nottingham Trent University, School of Science and Technology, Clifton lane, Nottingham, NG11 8NS, UK

Received ZZZ, revised ZZZ, accepted ZZZ
Published online ZZZ  (Dates will be provided by the publisher.)

PACS 81.15.Cd, 81.15.Ef, 42.79.Kr, 52.50.Dg, 78.20.-e

Abstract The temperature sensitive nature of the substrates used in the flexible displays market necessitates a low temperature deposition technique for processing them. ZnS:Mn exhibiting high intensity photoluminescence and good crystallinity has been deposited onto Si wafers, glass microscope slides and polymeric substrates using a new reactive sputtering technology referred to as HiTUS. This technique enables very high deposition rates and requires no substrate heating. When incorporated as part of a complete EL device, as-deposited ZnS:Mn films are seen to exhibit stable electroluminescence on Si, glass and planarised PET substrate materials. Post annealing of the devices on Si and glass at temperatures of up to 600 °C show that the HiTUS films perform better than equivalent ZnS:Mn films deposited using RF magnetron sputtering.

1 Introduction This work compares two different sputtering techniques for the deposition of alternating current thin film electroluminescent (ACTFEL) devices. High target utilisation sputtering (HiTUS) is a new method for sputtering thin films that allows for the deposition of working electroluminescent (EL) structures with no substrate heating or post deposition annealing [1]. However, here we look at the effect of post deposition annealing HiTUS devices that have been deposited onto silicon and glass substrates with no substrate heating. Radio frequency (RF) magnetron sputtering has also been used for the fabrication of comparable devices with both substrate heating and post deposition annealing. The results of quantitative EL measurements for structures deposited using these two techniques are presented here.

2 Experimental HiTUS and RF magnetron sputtering have both been used to deposit devices with the double insulating layer structure incorporating ZnS:Mn as the active, light emitting layer, Y2O3 as the dielectric layers and ITO as the front and back contact electrodes.

2.1 High Target Utilisation Sputtering Unlike RF magnetron sputtering, HiTUS relies on the generation of a plasma remotely from the targets. An RF coil antennae surrounding a quartz glass tube is located in a side arm (referred to as the plasma launch system, or PLS) adjacent to the vacuum chamber. The plasma is initiated here and amplified by a launch electromagnet at the exit of the PLS. A steering electromagnet is then used to focus and control the direction of the plasma. This system uses similar principles to an RF inductively coupled plasma but the magnetic enhancement provided by HiTUS results in higher plasma densities, of the order of 1015 ions cm−3. Further details are described elsewhere [2]. With careful control of the current flowing through each of the electromagnets, the plasma beam can be directed such as to cover the full surface area of the target. With the application of a sufficient negative bias, argon ions are then accelerated into the target resulting in a high current density over the full surface area of the target. Hence the target is uniformly eroded resulting in a significant reduction in target poisoning over conventional magnetron sputtering and hence also a much im-

Copyright line will be provided by the publisher
proved deposition rate when depositing non-metallic thin films.

The HiTUS system used for this work accommodates four 4" diameter, 6 mm thick water cooled targets. This allows for the deposition of complete devices without needing to break the vacuum and potentially introduce contaminants.

2.1.2 Deposition of ITO, Y₂O₃ and ZnS:Mn using HITUS All the materials used in this work were deposited from metallic targets in a reactive gas mixture of argon and either oxygen or hydrogen sulphide. Prior to depositing each layer, the substrates were immersed in a diffuse, high density plasma. This is easily achieved using HiTUS by switching off the steering electromagnet. The impinging low energy (less than 50 eV) argon ions remove volatile species from the substrate surface helping to promote adhesion of subsequent layers. It is the intention to also investigate the effect this diffuse plasma could have on the interface states between the ZnS:Mn and dielectric layers.

After this substrate pre-conditioning, the target is sputter cleaned for up to 5 minutes prior to admission of the reactive gas. Deposition conditions are then allowed to stabilise for a further 5 minutes before the shutter, which is located immediately adjacent to the substrates, is opened and deposition commences.

Other than the devices deposited onto silicon wafers, an ITO back contact was used. This process has been previously developed to be compatible with polymeric substrate materials and hence is a low temperature process. Details can be found elsewhere [2].

Both the Y₂O₃ and the ZnS:Mn were also deposited with no substrate heating, making this procedure directly transferable to plastics. A summary of the deposition conditions used for each of the materials within the EL structure is given in table 1.

<table>
<thead>
<tr>
<th>Material</th>
<th>Process pressure (mbar)</th>
<th>RF power (W)</th>
<th>DC power (kW)</th>
<th>Sub. Temp. (°C)</th>
<th>Dep. Rate (nm/min)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ITO</td>
<td>3.6 x 10⁻³</td>
<td>1.0</td>
<td>0.81</td>
<td>&lt; 70 °C</td>
<td>87</td>
</tr>
<tr>
<td>Y₂O₃</td>
<td>4.0 x 10⁻³</td>
<td>1.99</td>
<td>1.50</td>
<td>&lt; 70 °C</td>
<td>28</td>
</tr>
<tr>
<td>ZnS:Mn</td>
<td>7.7 x 10⁻³</td>
<td>0.79</td>
<td>0.20</td>
<td>&lt; 70 °C</td>
<td>52</td>
</tr>
</tbody>
</table>

2.2 RF magnetron sputtering For a comparative study, two separate RF magnetron sputtering systems (13.56 MHz) were also used for the deposition of Y₂O₃, ZnS:Mn and ITO. A base pressure of 1.2 x 10⁻⁶ mbar was used for the Y₂O₃ and ZnS:Mn layers and the second system used to deposit the ITO was evacuated to a base pressure of 1.6 x 10⁻⁷ mbar. Diffusion pumps backed by rotary pumps were used to attain this vacuum in both plants. The substrates were loaded into the main chamber through a side load lock. Two RF magnetron guns were positioned at 30° to the vertical and 15 cm from the substrate surface for the dielectric/phosphor depositions. The same configuration was used for the ITO coatings but with a target-substrate separation of 10 cm. The two main chambers were equipped with mass flow controlled gas lines providing 100 % Argon for the EL device layers (dielectric and phosphor films) and 0.2 % Oxygen/Argon for the ITO contacts. Alkali-free borosilicate glass and n-type silicon substrates were used in this study. Prior to deposition the glass and Si wafers were baked at 350 °C and 500 °C respectively. All targets were pre-sputtered for between 2 and 5 minutes prior to deposition. The ZnS:Mn layers were deposited by co-sputtering a 99.99 % pure ZnS target and a 99.95 % pure ZnS target doped with 1.0 wt.% Mn. Two RF power supplies providing 80 W to the ZnS target and 40 W to the target doped with Mn gave ZnS thin films with 0.5 wt.% Mn. The Y₂O₃ and ITO layers were deposited from 99.99 % pure compound targets. All targets were 3" in diameter and provided by Testbourne UK. Table 2 shows the growth parameters for all layers (the total power on both targets is shown for the ZnS:Mn layer).

<table>
<thead>
<tr>
<th>Material</th>
<th>Process pressure (mbar)</th>
<th>RF power (W)</th>
<th>Sub. Temp. (°C)</th>
<th>Dep. Rate (nm/min)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ITO</td>
<td>2.66 x 10⁻³</td>
<td>50</td>
<td>180 °C</td>
<td>4</td>
</tr>
<tr>
<td>Y₂O₃</td>
<td>6.66 x 10⁻³</td>
<td>120</td>
<td>200 °C</td>
<td>1.5</td>
</tr>
<tr>
<td>ZnS:Mn</td>
<td>6.66 x 10⁻³</td>
<td>120</td>
<td>200 °C</td>
<td>3.7</td>
</tr>
</tbody>
</table>

2.3 Comparison of sputtering methods The primary difference between the two sputtering techniques described above is that HiTUS decouples the ion density, which is controlled by the RF antennae power supply, from the ion energy, which is controlled by the DC bias applied to the target. This offers greater control over the growth process due to the independent control of multiple sputtering parameters.

From tables 1 and 2 it is clear that the HiTUS system also enables growth rates in excess of an order of magnitude faster than those offered by RF magnetron sputtering. Whilst higher deposition rates can be achieved using magnetron sputtering, this is to the detriment of the film properties since the RF power must be increased, with concomitant increase of the target bias, resulting in damage to film growth due to high energy ion bombardment [3,4]. Also of note is that these high growth rates are achieved using no substrate heating hence making HiTUS ideal for applications that require plastic substrates.

2.4 PL and EL measurement facility The photoluminescence (PL) measurements were carried out in a dark room at ambient temperature using a Laser Science VSL-337 ND nitrogen laser operating at 337 nm and 4 ns
pulses. A power of 400 μJ/pulse with a 20 Hz repetition rate was used. The intensity of the PL emission was measured using an S2000 Ocean Optics fibre optic spectrometer.

The EL measurements were also carried out in a dark room. A TTI TG1010 programmable function generator, a voltage amplifier (maximum supply of 900 V peak to peak) and a Minolta LS-110 luminance meter were used to drive the devices and record their luminance. The input devices were all connected in series to a current limiting 100 kΩ resistor. All EL measurements were carried out using a sinusoidal waveform voltage at a frequency of 1 kHz with voltage increments of 8 V peak to peak and 5 s duration.

3 Photoluminescence results

Prior to making a complete electroluminescent device using ZnS:Mn as the active, light emitting layer, individual films were first optimized for PL.

Three different targets were used to investigate the effect of different Mn concentrations on the PL properties of ZnS:Mn thin films deposited using HiTUS. Single layers of ZnS:Mn with thicknesses of between 0.5 and 1.0μm were initially deposited at room temperature onto glass microscope slides from Zn targets doped with 0.4, 0.5 and 0.6 wt.% Mn. For each of the three Zn:Mn targets the H2S flow rate was optimized to produce films that exhibited intense PL when excited by a nitrogen gas laser. As shown in Figure 1, the Zn target doped with 0.6 wt.% Mn consistently gave films with the highest PL intensity. The peak intensity of the emitted light is centred at approximately 600 nm, consistent with the 580 to 590 nm characteristic emission of Mn phosphor.

By increasing the H2S flow rate during deposition the coatings are seen to become fully transparent. For optimized, fully transparent films, the deposition rate can be increased to 90 nm/min for a target power of 0.5 kW. However, whilst this high deposition rate did produce coatings that displayed intense PL, they exhibited very poor breakdown properties when incorporated as part of an EL structure. ZnS:Mn films produced at lower powers were seen to exhibit excellent PL and EL properties. Figure 2 shows that for optimum PL a target power of approximately 200 W should be used, giving a deposition rate of 52 nm/min. The results of optimised ZnS:Mn films deposited at 200 °C using RF magnetron sputtering have also been included for comparison. Annealing these films at 500 °C sees an increase in the PL intensity but they still do not emit as brightly as those deposited using HiTUS. The HiTUS films were deposited with no substrate heating or post deposition annealing. Whilst the film deposited at 200 W was slightly thicker than the others, this only accounts for a small proportion of the increase in PL intensity.

Figure 2 PL intensity as a function of target power for HiTUS ZnS:Mn films deposited with no substrate heating or post deposition annealing. The results of films deposited at 200 °C and post deposition annealed at 500 °C using RF magnetron sputtering have been included for comparison.

Following the successful deposition of ZnS:Mn onto glass and Si substrates, single layers were also deposited onto Kapton™ (50 μm), planarised polyethylene naphthalate (PEN - 125 μm) and polyethylene terephthalate (PET - 188 μm) using HiTUS. Coatings with a thickness of up to 1.0 μm resulted in no deterioration of the flexible substrates.

The positive PL results indicate that these films should be excellent candidates for EL devices. Earlier work on ITO deposited using HiTUS identified this as a good material for the top and bottom contact electrodes due to its low resistivity (less than 4.0 × 10^{-4} Ωcm), high transmission and flexibility [2,5]. ITO films 200 nm thick deposited onto planarised PEN and PET have been shown to be flexible to a 1.0 cm bend radius with no subsequent increase in sheet resistance.

Dielectric layers of Y2O3 and Ta2O5 were also deposited onto glass and polymeric substrates with excellent transparency and breakdown properties, making possible the fabrication of completely transparent and flexible ACTFEL test devices.

3.1 ZnS:Mn XRD results

XRD analysis of these films showed that they are highly crystalline exhibiting a major diffraction peak at a 2θ angle of 28.5°. Subsidiary
peaks also occur at 59° and 95°. Peak matching revealed that the probable structure is either cubic or cubic with small admixtures of hexagonal phase in line with previous reports [6]. The crystallite sizes, as calculated from the Scherer equation and the FWHM of the main peaks, are found to be 79 nm, 73 nm and 69 nm for target powers of 200 W, 500 W and 100 W respectively. The crystallite size of the films deposited at 200 °C and post deposition annealed at 500 °C using RF magnetron sputtering is significantly smaller at 16 nm. This data is to form part of a further study looking at the correlation between crystalgraphic structure and PL response.

It is also clear from figure 3 that the optimum, as deposited, HiTUS films exhibit XRD peaks with higher intensity than ZnS:Mn films deposited at 200 °C and post deposition annealed at 500 °C using RF magnetron sputtering are also included.

4 Electroluminescence results The fabrication of ACTFEL test devices using both HiTUS and RF magnetron sputtering has been realised. The double insulating layer structure has been used for all devices with Y2O3 being used as the dielectric. Devices were initially deposited onto silicon substrates with ITO being used as the top contact. The possibility of making completely transparent devices on both glass and flexible substrate materials was then considered. The results in this section focus on a comparison of the two sputtering techniques to deposit working, EL devices on both glass and silicon substrates. Working EL structures were also deposited onto planarised PET using HiTUS. These results are presented elsewhere [1]. Due to the higher processing temperatures employed by RF magnetron sputtering, it was not possible to deposit devices onto plastic substrates using this technique.

4.1 EL devices deposited onto Si wafers Figure 4 shows the luminance as a function of applied peak to peak voltage (L-V) for an EL device deposited using RF magnetron sputtering. It is clear that heating the sample not only reduces the threshold voltage but also increases the maximum attainable luminance. This is to be expected as heating of the ZnS:Mn layer results in a greater number of Mn ions finding their energetically favoured position as substitutes for the Zn atoms within the host lattice. In this position they readily contribute to the emission of light as active, luminescent centres. When located as interstitial defects they simply act as scattering centres for electrons. The effect of annealing the complete EL structure is that the hot electrons injected from the interface states are released at lower fields and hence the threshold voltage for EL emission occurs for lower voltages. Above the threshold voltage the gradient of the L-V curve also becomes flatter for an anneal temperature of 600 °C indicating an increased energy distribution of interface states [7]. After post annealing in a vacuum at 600 °C the luminance is seen to increase to approximately 500 Cd/m² at 750 V.

Figure 4 Luminance as a function of applied peak to peak voltage for an EL device on silicon fabricated using RF magnetron sputtering.

For comparison, the results of an equivalent device incorporating ITO and dielectric layers deposited using RF magnetron sputtering but ZnS:Mn deposited using HiTUS with no substrate heating are shown in figure 5. A maximum luminance of 740 Cd/m² is now obtained for an applied peak to peak voltage of 690 V. Of significance is the sharpness of the turn-on when the structure is heated to 600 °C. This suggests that devices made with HiTUS
ZnS:Mn result in an interface region with a more concentrated distribution of trapped states. The increase in threshold voltage when increasing the anneal temperature from 400 to 600 °C is an interesting result that may imply a modification of the density of interface states via the removal of higher energy states (traps), or possibly a modification of the high field resistivity of the ZnS:Mn leading to increased threshold for impact excitation. This is to be investigated further. An increase in luminance of approximately 315 Cd/m² for a rise in voltage of just 50 V is seen. The same rise in voltage for the devices fabricated using ZnS:Mn deposited by RF magnetron sputtering sees only a 17 Cd/m² increase from 300 to 350 V, the steepest section of the L-V curve. This makes the drive electronics potentially much easier and cheaper to implement for devices deposited using HiTUS ZnS:Mn.

Figure 6 shows an L-V curve for a complete structure deposited using HiTUS, with no substrate heating or post deposition annealing. The effect of annealing at 200 °C is also shown although this results in little improvement of the device. For comparison, the results of post deposition annealing a device deposited at the substrate temperatures quoted in table 2 using RF magnetron sputtering is also shown. The maximum luminance is significantly higher with substrate heating. However, this obviously precludes the use of these devices for plastic applications. Devices deposited at ambient substrate temperature using RF magnetron sputtering exhibit no PL or EL and also breakdown very easily. It should be noted that the Y₂O₃ layer used in the HiTUS devices has not yet been optimised.

5 Conclusions ACTFEL devices have been deposited onto Si and glass substrates using two different sputtering techniques. Equivalent structures have been compared in terms of device performance for these two different technologies. Devices deposited onto silicon wafers using HiTUS ZnS:Mn are seen to offer a brighter maximum luminance as well as sharper turn-on characteristics despite being deposited with no substrate heating. Additionally, HiTUS allows for the deposition of working EL devices with no substrate heating or post deposition annealing; an achievement not possible with RF magnetron sputtering. The results presented thus far suggest that if the Y₂O₃ layers deposited using HiTUS can be optimised, ambient EL devices with comparable luminance to equivalent devices deposited using RF magnetron sputtering at elevated substrate temperatures could be possible.

References