Reduction of Contact Resistance in Bottom Contact Pentacene-TFT by Employing Carrier Injection layer

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1. Introduction

Organic thin-film transistors (OTFTs) are receiving considerable attention for use in the backplane of the active-matrix organic light emitting diode (AMOLED) displays fabricated on the flexible substrate[1]-[2]. High performances such as high on current, low operating voltage and rapid rise in subthreshold current are required for OTFT to drive the OLED display. In addition, these improvements of performances are especially needed in the bottom contact OTFT, because the photolithography patterning was used to realize higher resolution in the fabrication process of AMOLED display.

The reduction of contact resistance between source-drain (S-D) electrodes and organic semiconductor is important for the bottom contact OTFT to achieve the high performance[3]-[4]. In top contact OTFT, Chih-Wei Chu and co-worker improved the performance by employing bilayer S-D electrodes with metal oxide such as molybdenum oxide (MoO₃), vanadium oxide (VOₓ) and tungsten oxide (WOₓ)[5]. These metal oxides were well known materials as the carrier injection layer in OLED[6]. Additionally, it can be expected that the effect of the improvement in the bottom contact OTFT is larger than that of top contact OTFT.

We report on the reduction of contact resistance by employing S-D electrodes with MoO₃ as the carrier injection layer in bottom contact pentacene-TFT. The contact resistance was estimated from the gated transfer length method (gated-TLM)[4] and compared with the conventional S-D electrodes (Cr/Au).

2. Fabrication Process

A schematic cross section of the bottom contact pentacene-TFT was shown in Figure 1. A highly doped silicon wafer with thermally grown SiO₂ (200 nm) was served as the common gate electrode and gate insulator. Two types S-D electrodes consisting of MoO₃/Au or Cr/Au were fabricated on the substrate. MoO₃ layer was deposited by thermal evaporation from MoO₃ powder and then Au was formed on the MoO₃ layer in the same vacuum chamber. The S-D electrodes were patterned by photolithography and lift-off process. Cr/Au electrode was deposited by e-beam evaporation and patterned by the same procedure. The channel length L and width W were 50 µm and 500 µm, respectively. After the lift-off process, the substrate was immersed in HMDS solution for 8 h to improve the crystal growth of pentacene layer on the SiO₂ gate insulator. A 30-nm-thick pentacene, which was purified by the train sublimation twice, was formed on the substrate by thermal vapor deposition at room temperature (deposition rate 0.03 nm/s). The electrical characteristics were measured at room temperature by using an HP4140 source-meter system. Electrical characterizations were carried out in vacuum without exposure to air after the deposition of pentacene.

3. Improvement of TFT Performance

Figure 2 shows the plots of the drain current (I₉) (left) and square root of I₀ (right) vs. gate voltage (V₉) at a drain voltage (V_D) of 50 V for the pentacene-TFT with the S-D electrodes of MoO₃ (2 nm)/Au (50 nm) or Cr (2 nm)/Au (50 nm). MoO₃ provided a high adhesiveness between the gate insulator and Au electrode and fine pattern. MoO₃ carrier injection layer significantly improved the performance of the pentacene-TFT. Threshold voltage was reduced from 13 V to 0.2 V compared with the conventional S-D electrodes (Cr/Au). Drain current exceeded 10⁻⁵ A less than Vg of 20V, whereas the Vg of more than 40 V was needed in Cr/Au electrode for achieving the same current. The effect of improvement was especially large at the low operating voltage. The mobility in saturation regime was improved from 0.13 cm²/Vs to 0.42 cm²/Vs (MoO₃/Au) with the on/off ratio of 10⁵. MoO₃ carrier injection layer showed the highest performance at the thickness of few nano meters (~2 nm), which was comparable to the effective channel thickness on the gate insulator in pentacene-TFT[7]. The improvement of TFT performance was clearly attributed to the reduction of contact resistance in the S-D electrode by employing the MoO₃ injection layer.
Rikenkeiki). The Work functions of MoO_2 materials. These values were estimated from the level of the pentacene and work function of electrode of MoO_2 was estimated from the gated-TLM[4]. Contact resistance estimated to be 5.6 eV and 4.6 eV, respectively. The injection barrier height from MoO_2 is smaller than that of Cr/Au. This gate voltage dependence was consistent with the previous report[4]. In addition, the vacuum level of the organic semiconductor is expected to shift downwards at the interface between organic semiconductor and metal[8]. Therefore the injection barrier height at the interface might be higher than the estimated value in Figure 3.

Figure 4 shows the gate voltage dependence of contact resistance in the bottom contact pentacene-TFT with MoO_2 carrier injection layer. By employing MoO_2 carrier injection layer, TFT performance especially improved at the low operating voltage. threshold voltage was reduced from 13 V to 0.2 V and the mobility was improved from 0.13 cm^2/Vs to 0.42 cm^2/Vs. MoO_2 carrier injection layer significantly reduced contact resistance in bottom contact pentacene-TFT. Therefore MoO_2 was effective in the reduction of contact resistance in bottom contact OTFT.

5. Conclusions

We have fabricated the bottom contact pentacene-TFT with MoO_2 carrier injection layer. By employing MoO_2 carrier injection layer, TFT performance especially improved at the low operating voltage. threshold voltage was reduced from 13 V to 0.2 V and the mobility was improved from 0.13 cm^2/Vs to 0.42 cm^2/Vs. MoO_2 carrier injection layer significantly reduced contact resistance in bottom contact pentacene-TFT. Therefore MoO_2 was effective in the reduction of contact resistance in bottom contact OTFT.

References


Fig. 2 I_D-V_G characteristics of pentacene TFTs with Cr/Au or MoO_2/Au S-D electrodes.

Fig. 3 Energy level diagram of pentacene and electrode materials.

Fig. 4 Gate voltage dependence of contact resistance in pentacene TFTs with Cr/Au or MoO_2/Au S-D electrodes.