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PREPARATION AND LUMINESCENT PROPERTIES OF ZnTe-Cd_{1-x}Mn_xTe_{1-y}Se_y HETEROJUNCTION*

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ZnTe-Cd_{1-x}Mn_xTe_{1-y}Se_y heterojunctions were prepared by vapor-transport epitaxy of ZnTe on In-doped Cd_{1-x}Mn_xTe_{1-y}Se_y ($x = 0.05$, $y = 0.03$) single crystalline substrate in vacuum. At temperatures lower than 120 K the infrared and red electroluminescence were observed from the ZnTe-Cd_{1-x}Mn_xTe_{1-y}Se_y diode with forward current density in the range 0.003-4.0 A/cm².

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1. Introduction

Diluted magnetic semiconductors (DMS) have recently generated a great deal of interest not only for basic investigations but also for device applications [1]. Heterojunction between the DMS of A^{II}_{1-x}Mn_xB^{VI} type and A^{II}B^{VI} compounds may give new possibility to fabricate electroluminescence devices. In this paper we report the fabrication method of ZnTe-Cd_{1-x}Mn_xTe_{1-y}Se_y heterojunction and its luminescent properties.

2. Preparation of ZnTe-Cd_{1-x}Mn_xTe_{1-y}Se_y heterojunction

ZnTe-Cd_{1-x}Mn_xTe_{1-y}Se_y heterojunctions were formed by vapor-transport epitaxy of ZnTe on In-doped Cd_{1-x}Mn_xTe_{1-y}Se_y ($x = 0.05$, $y = 0.03$, In: 5×10^{18} cm⁻³) single crystalline substrates, cleaved along (110) surfaces from the as-grown crystal. The electron concentration in substrate material was from 1 to 3×10^{17} cm⁻³ at room temperature. Single crystals of ZnTe prepared by Bridgman method were used as the evaporating source. The growth of the ZnTe

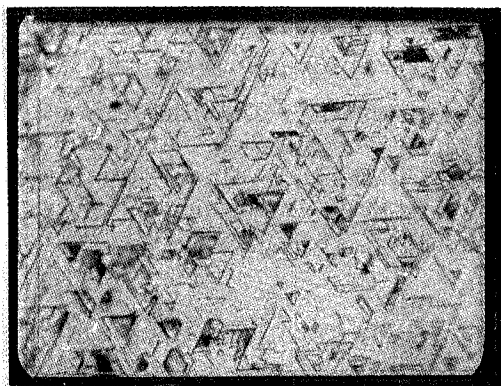
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epitaxial layer was performed in a quartz reactor pumped down to 1×10^{-6} mm Hg and placed in a horizontal two-zone temperature furnace.

The results of electron diffraction and metallographic observation showed that the quality of the layer structure depends on the source and substrate temperatures. Single crystalline ZnTe films can be obtained at the source and substrate temperatures $1150 \pm 5^\circ\text{C}$ and $580 \pm 5^\circ\text{C}$, respectively. Figure 1 shows the surfaces of non-etched ZnTe layers grown on $\text{Cd}_{1-x}\text{Mn}_x\text{Te}_{1-y}\text{Se}_y$ single crystalline substrate and on mica. It can be seen from Fig. 1 that the ZnTe epitaxial layer grows on $\text{Cd}_{1-x}\text{Mn}_x\text{Te}_{1-y}\text{Se}_y$ substrate in (110) direction, while on mica in (111) direction.



(a)



(b)

Fig. 1. Surface of ZnTe layers grown on $\text{Cd}_{1-x}\text{Mn}_x\text{Te}_{1-y}\text{Se}_y$ (a) and mica (b).

ZnTe-Cd_{1-x}Mn_xTe_{1-y}Se_y diodes were formed by soldering In contact to Cd_{1-x}Mn_xTe_{1-y}Se_y substrate and by evaporating Au film on the ZnTe layer, followed by thermal annealing at 410°C for 10 s. The size of the diodes was about 1.0 × 1.0 × 0.5 mm³. The current-voltage characteristics of the obtained diodes were measured at temperature range 2–300 K. They showed that in the dark, the forward current was about 10⁴ times higher than the reverse one at 1 V. Almost all the diodes exhibit interesting switching effect when the applied voltage changes. The switching threshold depends on temperature and on external illumination. The reverse current varies intensively with external illumination. These effects depend on the "history" of the diode*.

3. Luminescent properties of ZnTe-Cd_{1-x}Mn_xTe_{1-y}Se_y heterojunction

We measured the reflectivity and photoluminescence spectra of the CdMnTeSe substrate material and the ZnTe epitaxial layers grown on CdMnTeSe (110) substrate.

The photoluminescence (PL) spectrum of Cd_{0.95}Mn_{0.05}Te_{0.97}Se_{0.03} crystal excited by He-Ne or Ar laser at 4.2 K is characterized by four bands: the most intensive band with peak at 1.45 eV and three broad bands of weaker intensities with peaks at 1.6, 1.63 and 1.65 eV. From the PL and reflectivity spectra of the crystal, we determined the band gap energy of this material equal to 1.65 eV.

Electroluminescence (EL) measurement was performed in the temperature region 2–120 K by passing through the diode a stabilized forward current ranging from 0.03 to 40 mA (about 0.003–4.0 A/cm²). The radiation from the diode was led out for measurement in the plane parallel to the junction. The spectral distribution was analysed by a GMD 1000 double grating monochromator with a dispersion of about 3.0 cm⁻¹ and detected by a HAMAMATSU R 943-02 photomultiplier in photon-counting regime.

Figure 2 shows the spectra of the electroluminescence (EL) of the ZnTe-Cd_{1-x}Mn_xTe_{1-y}Se_y diode and photoluminescence (PL) of the substrate materials at 4.2 K. The EL spectrum consists of two regions: a relatively intensive and narrow peak at about 1.45 eV (so far called the main peak) and a long "tail" in the range from 1.6 up to 1.9 eV. It is worth noting that the intensity of the main peak depends on the density of the forward current (*J*) and reaches its maximum at *J* ~ 16 A/cm² (see insert in Fig. 3). Besides, the relative intensity of the main peak to the tail increased with the forward current (see Fig. 3), but the position of the observed bands is independent of the current.

Measurements of EL in magnetic fields up to 5 T showed no change in peaks position for both circular polarizations.

From the analysis of the EL spectra of the ZnTe-Cd_{1-x}Mn_xTe_{1-y}Se_y diode, for the moment, we interpret the emission band with peak at 1.45 eV as caused by radiative recombination of the acceptor-donor pairs in the region of the junction situated at the Cd_{1-x}Mn_xTe_{1-y}Se_y substrate side. The wide band in the region from 1.6 eV to 1.9 eV would be due to recombination of the injected electrons on

*The details of these studies will be published elsewhere.

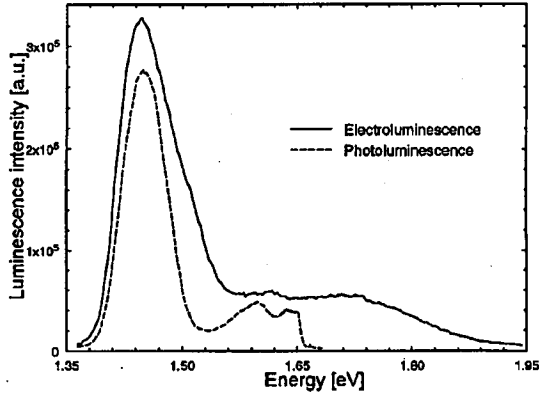


Fig. 2. Electroluminescence spectrum of ZnTe-Cd_{0.95}Mn_{0.05}Te_{0.97}Se_{0.03} diode (solid line) and photoluminescence spectrum of Cd_{0.95}Mn_{0.05}Te_{0.97}Se_{0.03} substrate (dashed line).

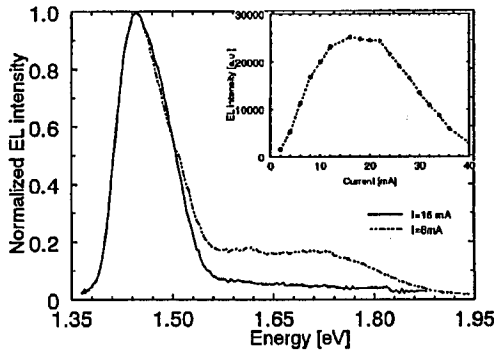


Fig. 3. Electroluminescence spectra normalized to the main peak, for two values of the current: 16 mA (solid line) and 8 mA (dashed line). In the insert: EL intensity at the main line as a function of the current intensity.

deep levels in ZnTe layer side of the junction and probably, partially by excitonic recombination in the substrate side.

Acknowledgments

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References

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