purnal of Luminescence 40 & 41 (1988) 521-522 worth-Holland, Amsterdam

THE LUMINESCENCE OF HIGHLY EXCITED CdS

Xianping LIU, Qingcheng BAO, Renshong DAI and Xurong XU Changchun Institute of Physics, Academia Sinica, Changchun, China

We measure the emission spectra of single crystal CdS from  $10^6 \text{W/cm}^2$  to  $10^7 \text{W/cm}^2$  excitation intensity at 4.2K, and observe a new luminescence peaking at 4910Å after the excitation intensity increases to 1.0 X  $10^7 \text{W/cm}^2$ .

# 1. INTRODUCTION

cdS single crystal is a very typical direct band semiconductor material, which is paid wide attention to in basic and applied research. There are a large number of papers dicussing the emission spectra of CdS, the luminescence characteristic of CdS under very low and very high excitation intensities having been known basically  $^{1-4}$ , that is, the luminescence of bound excitons in CdS is important on low excitation intensity, and the luminescence of electron-hole plasma is dominant on very high excitation intensity. The luminescence of CdS is quite complex on middle excitation intensity, so the causes of M band P band are not well known.

# 2. THE RESULTS OF EXPERIMENTS

The samples used in our experiments are very pure CdS crystals which are grown by the vapour phase epitaxy method, whose thickness is about hundreds  $\mu$  m. The samples are held in cryostat ESR-900 and cooled down at 4.2K±0.5, excited by a light beam of wavelength 308nm from the excimer laser EMG-102 XeCl, the single pulse energy is about 150mJ, the pulse duration is about 10ns and the pulse repetitions rate is 10 Mz.

Fig. 1 shows the four emission spectra of CMS in the range of excitation intensities  $10^6-10^7 \text{W/cm}^3$  at 4.2K. unter lower excitation intensity (a), the luminescence has mainly both  $^{4869}\text{Å}$  peak and 4888Å peak, which are the luminescence has mainly both

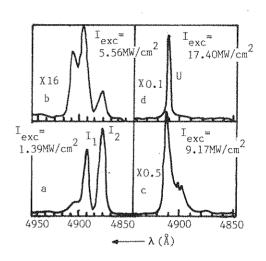


FIGURE 1 Luminescence spectra of CdS for different excitation intensities at 4.2K,  $hv_{\rm exc}$ =4.02eV.

nescence of excitons bounded to neutral  $donors(I_2)$  and excitons bounded to neutral accepters  $(I_1)$ , respectively. A new luminescence band arises on the long wavelength of  $I_1$  peak with an increase of excitation intensity while the intensity of  ${
m I_2}$  peak is weakened. After the excitation intensity is increased to 1.74 X $10^7 \text{W/cm}^2$  the spectra is led by a new peak -4910Å, which we call U peak. U peak has only 3Å halfwidth. Fig. 2 shows variation with excitation intensities of U peak luminescence intensity. The front of the curve is approxiamately quadratic and the rear straight. We measure variation with temperature of U peak on constant exciation intensity, see Fig. 3.

<sup>®</sup>22~2313/88/\$03.50 © Elsevier Science Publishers B.V.

North-Holland Physics Publishing Division)

12000 C in polarisations o) to the chain c

24000

S.McClure, J.Chem.

J.R.G.Thorne,

m. Soc. Faraday

ersity of Oxford,

, Mol. Phys. 30

ala, Mol. Phys.

H.J.Guggenheim,

il. Mag. 3 (1957)

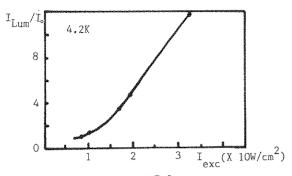


FIGURE 2

Variation with excitation intensities of U peak luminescence intensity I at 4.2%. (The luminescence intensity of I = 0.17MW/cm² is regarded as an unit of luminescence.)

## 3. DISCUSSION

According to a reduction of  $I_2$  peak when U peak arise, and ultra-linear variation of its luminescence intensity with excitation intensity, we think U peak as the luminescence of inelastic scattering of excitons with excitons bounded to neutral donors. When the density of excitons is very high in luminescence bodies, the scattering possibility of excitons with bound excitons becomes greater. After the scattering takes place, excitons are excited to higher energy states (n=2,3,...,) and bound excitons turn to luminescence by recombination. According to a prinple of energy conversation in scattering course, we give energy formula of the peak,

$$h\omega_{\text{max}} = Eg(T) - E_{x}^{b} - E_{bx}^{b} (1-1/n^{2})$$
 (1)  
 $n = 2, 3, ...$ 

where Eg(T) is a width of forbidden band;  $E_{x}^{b}$  is a bound energy of excitons;  $E_{bx}^{b}$  is a bound energy of exciton bounded to neutral donor. When n = 2, h $\omega_{max}$  = 4909Å (T = OK). We derivate the relation between luminescence intensity of U peak and excitation intensity by means of the luminescence model of inelastic scattering of excitons with bound excitons

$$I_{Lum} = \frac{a}{1 + b I_{exc}} I_{exc}^{2} \qquad (2)$$

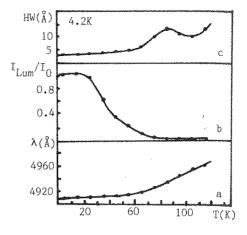


FIGURE 3
Variation with teperature of the emission maxima (a), luminescence intensity (b) and half width (c) of U peak. (Luminescence intensity I at 4.2K is regarded as an unit of luminescence in (c).)

where a and b are all constant not relative to excitation intensity  $I_{\rm exc}$ . We see that  $I_{\rm Lum}$   $I_{\rm exc}$  is ultra-linear on low excitation intensity and linear on high excitation intensity. This is basically agreement with behaviour of U peak.

### 4. CONCLUSION

We observed a new luminescence peak in CdS at 4.2K after excitation intensity is increased to 1.0  $\times$  10  $^{7}$  W/cm<sup>2</sup>. The peak is at 4910Å of the spectra and has only 3A halfwidth and quite high luminescence intensity. We propose the luminescence model of inelastic scattering of excitons with excitons bounded to neutral donors for the peak.

### REFERENCES

- H. Schrey and C. Klingshirn, Phys. Stat. Sol. (b) 90, 67(1978).
- 2. C. Klingshirn and H. Haug, Phys. Rep. 70, 315-398 (1981).
- Baldassare Di Bartolo, Collective Excitations in Solid (1983).
- H. Yoshida and S. Shionoya, Phys. Stat. Sol. (b) 115, 203 (1983).

THE INTERA

Li MA, Zil Changchun

We found a under high

1. INTRODUCTI ZnSe is or for applicati al. 1 in ZnSe tributed the excitons scat combination. the interact cant roles to this paper a band, associ attributed to plained by to ZnSe epilaye

2. EXPERIMEN
ZhSe epil
strate by va
laboratory.
QJD-9 was us
peak power ]
spectra were
of Model 444
multiplier.

3. RESULTS .
In Fig.
bands, E<sub>S</sub> at respectivel
intensity,
tion shifts
has no chan

0022-2313/88, North-Holland