Lawrence Berkeley National Laboratory

Recent Work

Title

ELECTRIC FIELD QUENCHING OF THE PHOTOLUMINESCENCE OF ZnO PELLETS

Permalink

https://escholarship.org/uc/item/9wz6z5dj

Author

Petermann, Gunter.

Publication Date

1972-12-01

RECEIVED
Submitted to Journal of Applied Physics FADMAINNE LBL-1546
RADIATION FABORATORY Preprint c.

CIBRARY AND DOCUMENTS SECTION

FIRE WA

ELECTRIC FIELD QUENCHING OF THE PHOTOLUMINESCENCE OF ZnO PELLETS

Günter Petermann

December 1972

Prepared for the U. S. Atomic Energy Commission under Contract W-7405-ENG-48

For Reference

Not to be taken from this room



DISCLAIMER

This document was prepared as an account of work sponsored by the United States Government. While this document is believed to contain correct information, neither the United States Government nor any agency thereof, nor the Regents of the University of California, nor any of their employees, makes any warranty, express or implied, or assumes any legal responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by its trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof, or the Regents of the University of California. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof or the Regents of the University of California.

0003993479

ELECTRIC FIELD QUENCHING OF THE PHOTOLUMINESCENCE OF ZnO PELLETS Günter Petermann

ABSTRACT

The influence of an electric field on the photoluminescence of pressed ZnO pellets was investigated. A quenching of the luminescence intensity was found. Suggestions are made to improve the original luminescence intensity and use this effect in practical applications.

ELECTRIC FIELD QUENCHING OF THE PHOTOLUMINESCENCE OF ZnO PELLETS Günter Petermann*

Laboratory of Chemical Biodynamics, Lawrence Berkeley Laboratory,
University of California, Berkeley, California 94720

The quenching of photoluminescence by an electric field (Dechène effect) has been observed for several inorganic ¹⁻⁴ and, to the author's knowledge, for one organic (poly-N-vinylcarbazole) material. The inorganic compounds are usually Zn and Cd sulfides, and the investigations have been performed on single crystals as well as on polycrystalline thin layers. The effect is explained in terms of a decrease in the electron density due to the extraction of charge carriers from the illuminated surface by the field. 4

The electric field induced changes of zinc oxide luminescence have been observed by Filinski and Skettrup, who have used this effect to distinguish between various types of exciton complexes at low temperatures. Those investigations were confined to the luminescence in the narrow (360 to 380 nm) spectral range near the absorption edge of ZnO. In addition to this blue emission, zinc oxide possesses a broad green emission band; the relative intensity of both bands is strongly dependent on the doping and on the preparation of the material. The quenching of the green luminescence by an electric field and its relationship to the photocurrent has been reported in crystals of ZnO. Since the energy efficiency of the green luminescence in ZnO can be rather large (approximately 20%) and the observed effect is very pronounced (a few volts were sufficient to yield extinction of the luminescence of these crystals), the development of practical applications such as display devices seems to be feasible.

Single crystals of ZnO are rather difficult to obtain, 10 but the technology for preparing large areas of ZnO seems to be well established (Electrofax paper). 11 Therefore it was of interest to compare the findings made on single crystals of ZnO to thin layers of polycrystalline material. The most obvious method would have been to use ZnO paper in the experiments, especially since its ZnO coating shows a slight green luminescence under ultraviolet illumination. However, it was not practical to use the Electrofax paper because of the necessity of making electric contacts, and a different arrangement was subsequently chosen for the preliminary investigations.

Zinc carbonate was calcined in air at 1000°C to obtain the green fluorescing ZnO. ¹² This material was pressed into pellets and contacted with a wire on one side into which Indium had been diffused by heating to 600°C to obtain an ohmic contact. The pellets were then mounted as electrodes in an electrochemical cell, using a 1 M KCl solution as the second contact. The luminescence was excited with a Xenon lap (wavelengths range: 310 to 390 nm) and monitored with a photomultiplier while a positive voltage of 12 volts was switched on and off to the pellet. Further experimental details have been described in Ref. 13, where the same arrangement was used.

The result of this experiment is shown in Fig. 1. The application of the positive voltage leads to a reduction (less than 1%) of the green luminescence intensity of the ZnO pellet. The time response is approximately equivalent to the time response of the recorder; as yet, no attempts have been made to measure the real response time.

The voltage necessary to achieve an effect is much higher for the polycrystalline thin layers than it is for the single crystals of ZnO, 8,13 and the effect itself is rather small. Even the original luminescence of the pellets is much smaller, which is probably due to pressure quenching 14 as a result of the preparation method. The change in the luminescence intensity when the voltage is applied seems to be restricted; this is in contrast to the large reduction of the luminescence intensity of the single crystals. However, the experiments have shown that the effect also exists in thin layers of polycrystalline ZnO. Different preparation methods for the ZnO and the electrodes (e.g., reductive treatment after calcining which increases the luminescence intensity 11 and sintering of the ZnO powder onto a suitable base) might result in properties equivalent to those of the single crystals and lead to practical applications.

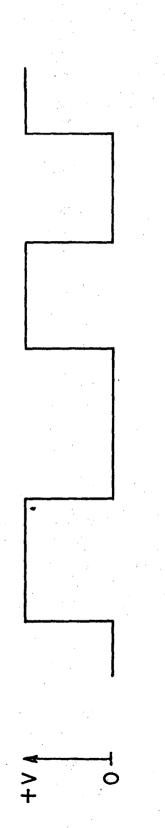
The author gratefully acknowledges the grant of a fellowship by the Deutsche Forschungsgemeinschaft, and the helpful suggestions and criticisms of M. Tomkiewicz. He also wishes to thank M. Klein for reading the manuscript. This work was supported, in part, by the U. S. Atomic Energy Commission.

References

- 1. T. H. Yee and G. A. Condas, J. Appl. Phys. 40, 2317 (1969).
- 2. F. Matossi and S. Nudelman, Phys. Rev. <u>89</u>, 660 (1953).
- 3. G. A. Zholkevich and V. P. Dudnik, Optika i Spektroskopiya 17, 745 (1964).
- 4. P. J. Daniel, R. G. Schwarz, M. E. Lasser and L. W. Hershinger, Phys. Rev. 111, 1240 (1958).
- 5. R. B. Comizzoli, Photochem. Photobiol. 15, 399 (1972).
- 6. I. Filinski and T. Skettrup, "Electric-Field and Temperature Dependence of Luminescence in ZnO," Proc. Intern. Conf. Semicond. Phys., 9th, Moscow, 1968, p. 216 (1968).
- 7. B. Andress, Z. Physik. <u>170</u>, 1 (1962).
- 8. G. Petermann, H. Tributsch and R. Bogomolni, J. Chem. Phys. <u>57</u>, 1026 (1972).
- 9. G. Heiland, E. Mollwo and F. Stöckmann, "Electronic Processes in ZnO," Solid State Physics, Vol. 8, Academic Press (1959), p. 230.
- 10. Ibid., p. 200.
- 11. Ibid., p. 295.
- 12. H. Gobrecht, D. Hahn and K. Scheffler, Z. Physik. 139, 365 (1954).
- 13. G. Petermann, M. Tomkiewicz and M. P. Klein, to be published.
- 14. See Ref. 9, p. 228.
 - *Present address: Inst. f. Physik. Chemie, Universitaet Duesseldorf, 4 Duesseldorf, Germany.

FIGURE CAPTION

Fig. 1. Influence of an electric field on the photoluminescence of \mbox{ZnO} pellets.



XBL7211-4810



LUMINESCENCE INTENSITY (rel. units)

LEGAL NOTICE-

This report was prepared as an account of work sponsored by the United States Government. Neither the United States nor the United States Atomic Energy Commission, nor any of their employees, nor any of their contractors, subcontractors, or their employees, makes any warranty, express or implied, or assumes any legal liability or responsibility for the accuracy, completeness or usefulness of any information, apparatus, product or process disclosed, or represents that its use would not infringe privately owned rights.

TECHNICAL INFORMATION DIVISION
LAWRENCE BERKELEY LABORATORY
UNIVERSITY OF CALIFORNIA
BERKELEY, CALIFORNIA 94720