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Surface studies of p-GaN:Cs photocathodes with in-situ X-ray photoelectron spectroscopy (XPS)

Introduction

- Higher beam currents and brightness are desired, therefore
- new photocathodes with higher QE are required
- p-type GaN is able to produce a negative electron affinity (NEA) surface when cesium is deposited on it
- A thermal cleaning under vacuum was carried out to achieve an atomically clean surface prior to the Cs deposition



Methods and Material

- The p-GaN surface is studied with *in-situ* XPS without leaving the UHV environment
- A PHI 5600 spectrometer (average pressure of 4 × 10⁻⁹ Torr and Al Ka line (*hv* =1486.6 eV) from non-monochromatized x-ray source was used
- p-GaN (5 μm), grown on sapphire with

Achieved QE values for p-GaN:Cs

- QE depends on the surface conditions
- The QE values correlate with the temperature used in the thermal cleaning
- Samples were re-activated several times



Fig. 1: The interior of the UHV preparation chamber (showing a sample holder, a halogen lamp, an anode, and cesium dispensers) and the XPS analysis chamber connected to the preparation chamber.

Surface Studies of thermal cleaning

- p-GaN was rinsed in 99 % pure EtOH
- EtOH solvent residues (C-OH) attached to the p-GaN surface
- thermal cleaning of (450 °C, 1h) was applied: peak intensity of EtOH residuals was reduced



metalorganic chemical vapor deposition (MOCVD) was used with a Mg conc.: 5 × $10^{16} - 1 \times 10^{17}$ cm⁻³

p-GaN:Cs photocathode

- Cs was deposited on the p-GaN to achieve a NEA surface
- Cs current: 3.0 4.0 A (3 × 10⁻⁹ mbar)
- p-GaN was illuminated with 310 nm UVlight during the Cs deposition
- In-situ photocurrent was observed until a maximum was achieved



- *Figure 2:* Achieved QE values of the p-GaN:Cs photocathodes depending on their surface conditions (applied T in thermal cleaning).
- Higher QE (max. 11.5 %) values were achieved @ moderate temperature (400—500 °C)
- Less QE values @ T > 600 °C
- During the first experiments from A1-A6
 no *in-situ* XPS was available
 → no surface studies

Surface after Cs deposition (XPS)

- Cs caused a shift toward higher binding energies, but not in O 1s
 → O located in bulk, not at surface
- most influence observed in C 1s peak: new component appeared at 286 eV

 \rightarrow cesium carbide species



- *Figure 3:* O 1s and C 1s photoelectron spectra for the p-GaN surface cleaned with 99 % EtOH (line 0) and after thermal cleaning (line 1).
- Thermal cleaning was not able to remove O and C entirely
- C and O contaminants remained
- \rightarrow Derive from MOCVD ?

Photocathode degradation

- QE decays usually 1/e, but X-rays accelerated the degradation
- Photoemission peaks shifted toward lower binding energies



Figure 4: In-situ photocurrent and vacuum value during the cesium activation of the p-GaN surface.

- 7.7 % QE was achieved although C and O remained on the surface
- Simple and manageable preparation compared to other semiconductor photocathodes

Conclusion

- Our study showed that p-GaN has an high potential to become a future electron source
- Thermal cleaning cannot remove O and C contaminations entirely
- $(\rightarrow$ C and O were residuals from MOCVD?)
- Cs caused a shift toward higher binding energy
- In C 1s: new species was created
 Formation of cesium carbide species caused an external degradation

Figure 6: The QE decay of the p-GaN:Cs photocathode and the Ga 3d_{3/2} and C 1s photoemission spectra at different times during its decay. The C 1s spectrum showing the evolution of the cesium carbide species.

- peak intensity of the cesium carbide species (286 eV) increased with ongoing degradation
- X-rays accelerated the photocathode degradation additional

Outlook

- p-GaN with higher quality (MBE or HVPE)
- On different substrates ? (SiC, Si)
- Remove C and O with low energy ions ?
- Influence of Mg concentration on QE ?

References related to p-GaN:Cs photocathodes

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