Sandia National Laboratories



IMPROVING ALKALINE ZINC-COPPER OXIDE BATTERIES THROUGH CHEMICAL MODIFICATIONS

Bryan R.Wygant, ¹Ciara N.Wright, ²Timothy N. Lambert^{2,3*}

¹Nanoscale Sciences, ²Department of Photovoltaics & Materials Technologies, ³Center for Integrated Nanotechnologies, Sandia National Laboratories, Albuquerque, New Mexico 87185, USA *tnlambe@sandia.gov



Other Routes to Improvement

In addition to changes to the quantity of the Bi_2O_3 additive, there are several other ways to further affect Zn-CuO battery performance and lifetime. These include alternative additives, novel coatings, and modifications to the electrolyte.







Schorr et al., ACS Appl. Energy Mater., 2021, 4, 7073-7082

- Zn-CuO has long been known as a high capacity, low voltage primary battery, but recent work has shown the chemistry is reversible in the presence of Bi_2O_3
- Optimization of Bi_2O_3 quantity, as well as other chemical modifications, expected to yield further improvements to battery performance



Tuning the total concentration of Bi_2O_3 (solid and soluble) has a small, but notable, impact on both the capacity and energy density of the Zn-CuO battery



Total quantity of solid Bi_2O_3 (in cathode) appears to impact charge/discharge behavior ~I wt% Bi_2O_3 appears to be reasonable balance all of performance metrics



f

<u>20 nm</u>

Use of gelling agents in electrolyte the and chemical additives like Bi can also influence the capacity/energy density and cycling performance of Zn-CuO batteries

- I. Reducing Bi quantity improves energy density while retaining capacity benefits
- 2. Both soluble and solid Bi play a role in mediating Zn-CuO battery cycling
- 3. Other additives, the use of coatings, and electrolyte modifications can also impact/improve battery performance
- 4. Understanding capacity decay and improving battery lifetimes are important next steps

Where Does the Bi Go?

Presence of Bi (Bi^0 or Bi_2O_3) reduces charge transfer resistance, improves cathode conductivity

- STEM/EDX imaging of discharged cathodes shows that the Bi is evenly co-located with the Cu, and but is likely concentrated at the particle surface
- Bi electrochemistry likely mediated through the soluble $Bi(OH)_4^{2-}$ species

ENERGY

Sandia National Laboratories is a multimission laboratory managed and operated by National Technology & Engineering Solutions of The views expressed in this article do not necessarily represent the views of the U.S. Department of Energy or the United States Government. This Sandia, LLC, a wholly owned subsidiary of Honeywell International Inc., for the U.S. Department of Energy's National Nuclear article has been authored by an employee of National Technology & Engineering Solutions of Sandia, LLC under Contract No. DE-NA0003525 with Security Administration under contract DE-NA0003525. This work was supported by the U.S. Department of Energy (DOE). The National Technology & Engineering Solutions of Sandia, LLC employee owns all right, title and interest to Electricity. Dr. Imre Gyuk, Director of Energy Storage Research at the U.S. Department of Energy Office of Electricity, is thanked for their contribution to the article and is responsible for its contents. The United States Government retains and the publisher, by accepting the article his financial support of this project. This work was performed, in part, at the Center for Integrated Nanotechnologies, an Office of for publication, acknowledges that the United States Government retains a non-exclusive, paid-up, irrevocable, world-wide license to publish or Science User Facility operated for the U.S. Department of Energy (DOE) Office of Science. reproduce the published form of this article or allow others to do so, for United States Government purposes. The DOE will provide public access to these results of federally sponsored research in accordance with the DOE Public Access Plan https://www.energy.gov/downloads/doe-public-

access-plan."

SAND2023-10686C