A High-Capacity and Long-Cycle-Life Lithium-Ion **Battery Anode Materials; Graphene Encapsulated** Co₃O₄ Nanoparticles

Dong Sung Choi

Chanhoon Kim, Il-Doo Kim, Sang Ouk Kim

KAIST, Department of Materials Science and Engineering, 291 Daehak-ro, Yuseong-gu Daejeon 34141, Republic of Korea

ongyarchoi@kaist.ac.kr

For practical large-scale production for catalyst synthesis, the preparation process must be simple, scalable, and cost effective. To this end, many methods have been proposed. Nonetheless, these methods suffer from intrinsic drawbacks, such as (i) the requirement of complicated multistep processes that include mixing, precipitation, cleaning, drying, calcining, and so on; (ii) catalyst poisoning by chemical residues; (iii) severe conditions required for alloying; (iv) poor control over the mean size and size distribution of catalyst particles; and (v) poor adhesion with supporting materials. [1] specifically, long-term More catalytic efficiency can be greatly improved by addressina typical degradation the mechanisms of catalysts, such as agglomeration, dissolution into the electrolyte, poisoning of catalytic sites and degradation of the catalytic support. [2] Here, we introduce a facile synthetic process for inorganic-organic hybrid catalyst systems, specifically those of graphene encapsulated cobalt oxide for Li-ion battery anode materials.

References

- Dong Sung Choi et al., Advanced [1] Materials, 33 (2016) 7115-7122
- [2] Heeyeon Kim et al., ACS Nano, 6 (2015) 5947-5957

Figures

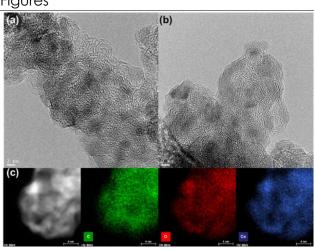


Figure 1: High resolution TEM images of graphene encapsulated Co₃O₄ nanoparticles and HAADF image and EDS maps of C, O and Co.

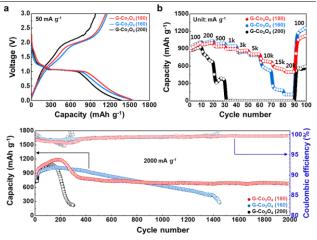


Figure 2: (a) Galvanostatic first cycle discharge /charge voltage profiles and (b) rate capability encapsulated the araphene CO3O4 of nanoparticles. (c) Long-term cycling stability of graphene encapsulated Co₃O₄ nanoparticles.