
Preface

The electrochemical oxygen evolution reaction (OER), an anodic half-reaction of water electrolysis, has attained enormous attention in the field of energy conversion. However, OER is a bottleneck for the large-scale production of hydrogen gas due to the involvement of high energy reaction intermediate following a four-proton and four-electron transfer mechanism. Therefore, the OER kinetics is sluggish and it becomes a thermodynamically unfavorable reaction. OER requires additional energy input to cross the high energy barrier. Recently, efforts have been provided to develop transition metal-based electrocatalysts for efficient electrochemical OER. Among different OER active materials, layered double hydroxides (LDHs) have emerged as one of the most outstanding electrocatalysts because of the flexibility in their composition and structure and facile fabrication techniques. In this reference, Co-LDHs have been considered to show promising OER activity but their OER activity is constrained due to the low Gibbs free binding energy of *O on Co-sites and their low electronic conductivity.

Recently, MOF-derived self-supported LDHs have come out as one of the most promising groups of electrocatalysts because of their unique layered structure and potent electrocatalytic activity. In this context, we have utilized self-supported ZIF-67 (zeolitic imidazolate framework, a subclass of metal organic framework, MOF) as the precatalyst to form the active catalysts. ZIF-67-derived LDHs have high surface area and tunable pore structure, which make them suitable OER catalysts.

In addition to this, the incorporation of high-valent metals in Co-LDHs can greatly enhance the OER performance. Simultaneously, the use of a self-supported catalyst on a conductive substrate can enhance the electronic conductivity and strengthen the interaction between catalysts

and support which improves the activity and stability. Therefore, in this thesis, we have explored the self-supported and heterometal-doped Co-LDHs to attain the required OER catalytic activity.

Chapter-1 demonstrates the objectives of the thesis work looking at progress in this field. In this chapter, we have covered the principles of electrochemical water oxidation, its significance in the field of energy conversion and storage, and its evaluation parameters. Further, we have discussed the previously reported MOF-derived LDHs for OER along with various shortcomings in this field. We have discussed the advantages of using ZIF-67 as the precursor for the synthesis of OER catalysts.

Chapter-2 demonstrates the synthesis of noble-metal free V-doped Co-LDHs ($V_x\text{Co-LDH}$) hollow nanocages, synthesized from ZIF-67. ZIF-67 was converted into LDH along with the incorporation of vanadium when treated with VCl_3 . The V doping has been found to be crucial to enhance the OER activity of Co-LDH in the alkaline medium. The effect of changing the amount of V on the activity of $V_x\text{Co-LDH}$ was also studied and it was observed that all $V_x\text{Co-LDH}$ exhibited better OER activity than Co-LDH while the activity of $V_{0.2}\text{Co-LDH}$ was found to be best among all the synthesized $V_x\text{Co-LDH}$ s. $V_{0.2}\text{Co-LDH}$ required only 280 mV of overpotential to produce 100 mA cm^{-2} of current density with stability for 24 h.

Chapter-3 explains the Lewis acid-assisted hydroxylation of ZIF-67 to form $\text{Fe}_x\text{Co-LDH}$ nanosheets via hydrothermal technique. The most appropriate reaction condition for the complete reconstruction and optimal OER activity has been determined. The effect of variation in the quantity of Fe(III) on the OER activity of $\text{Fe}_x\text{Co-LDH}$ has been explored where it was found that

Fe_{0.4}Co-LDH exhibited the best OER activity. The enhanced electrochemical OER activity of ultrathin nanosheets has also been compared with Co-LDH, ZnCo-LDH, and hydrothermally synthesized Fe_{0.4}Co-HC.

In **Chapter-4**, we have shown electrochemical reconstruction of ZIF-67 during anodic water oxidation resulting in the formation of FeCo-LDH-x [Fe-Co(OH)₂/Co(O)OH] (where x = 1, 2, 3, 4, 5). The Fe(III)-ions, present in the electrolyte solution, are introduced in the structure of the active catalyst during the anodic reconstruction of ZIF-67. Electrochemically derived FeCo-LDH-x demonstrates high electrochemical surface area, exposed active sites, atomic-level thickness, and adjusted structural and electronic features to improve the OER activity. The activity of all FeCo-LDH-x was found to be better than that of as synthesized Co-LDH-3. Among the synthesized catalysts, FeCo-LDH-3 exhibited the best OER activity generating 50 mA cm⁻² current density at 180 mV overpotential and with the lowest Tafel slope value and charge transfer resistance.

Chapter-5 demonstrates a facile technique for the synthesis of a self-supported heterostructure catalyst system CeO₂/Ce-Co-LDH (CeCo-x, where, x= 1, 2, 3) using nickel foam supported ZIF-67. The precursor was treated with various amount of Ce(NO₃)₃.9H₂O by hydrothermal technique. During hydrothermal treatment, the insertion of Ce into Co-LDH resulted in 4f-2p-3d orbital overlap along with the formation of heterojunction between CeO₂ and CeCo-LDH. Out of all synthesized CeCo-x catalysts, CeCo-2 demonstrated excellent OER activity as it produces 50 mA cm⁻² current density at a very low overpotential of 220 mV. 4f-2p-3d orbital overlap modulated the electronic structure and the heterostructure regulated the transportation of charges during the OER process. The 4f-2p-3d orbital overlap demonstrated the more proficient delocalization of π -

electron density over metal center via oxide bridging in comparison to 3d-2p-3d overlap. Thus, CeO₂/Ce-Co-LDH heterostructures exhibited excellent OER performance and stability.

In Chapter-6 we have presented a succinct overview of the scientific contributions made in this thesis and summarized our findings, which provide valuable insights for the development of ZIF-67 derived new efficient electrocatalysts for electrochemical energy conversion. We have also explored the future prospects of these research endeavors.