

Exfoliated PdSe₂ supported on graphene oxide as an electrocatalyst for efficient water splitting

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INTRODUCTION & AIM

Efficient hydrogen evolution reaction (HER) in alkaline media is crucial for advancing sustainable hydrogen production but remains a significant challenge due to sluggish reaction kinetics. Two-dimensional transition metal chalcogenides (TMCs) have emerged as promising electrocatalysts, offering tunable electronic properties and abundant active sites. Among them, palladium selenide (PdSe₂) stands out for its catalytic potential. However, the intrinsic conductivity and dispersion of PdSe₂ limit its performance. To overcome these drawbacks, graphene oxide (GO) can serve as a conductive scaffold, improving charge transport and catalyst stability.

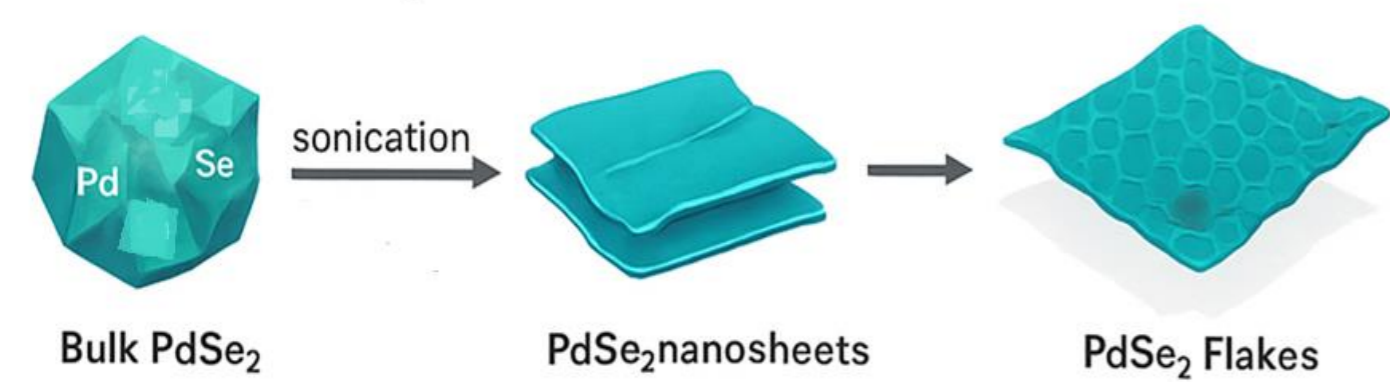
This study aims to develop a composite catalyst based on exfoliated PdSe₂ nanosheets integrated with GO to enhance HER activity in alkaline electrolyte.

The specific objectives are to:

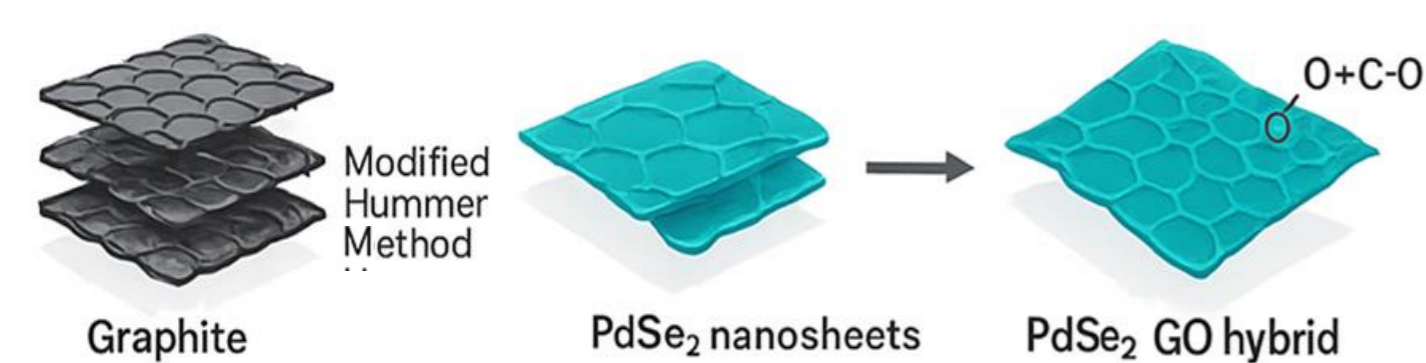
- ✓ Exfoliate bulk PdSe₂ to increase the density of accessible active sites.
- ✓ Utilize GO as a conductive and dispersive support for the nanosheets.
- ✓ Evaluate the electrocatalytic performance of PdSe₂-GO toward HER in alkaline media and compare it with pristine PdSe₂.

METHOD

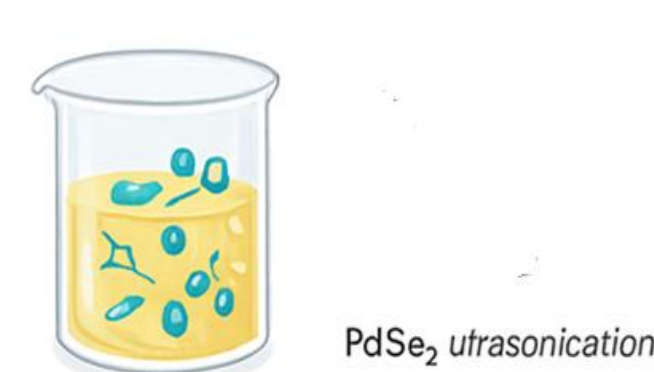
Exfoliation of PdSe₂



Synthesis of Graphene Oxide (GO)



Composite Formation



Electrode Preparation

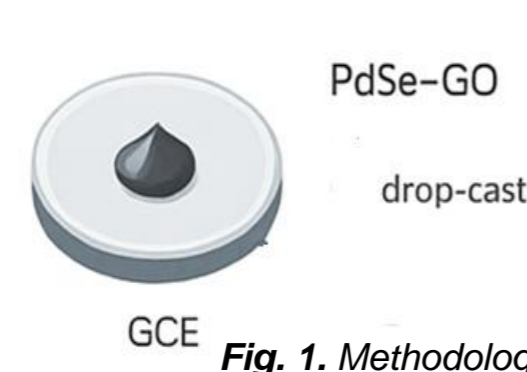


Fig. 1. Methodology workflow

RESULTS & DISCUSSION

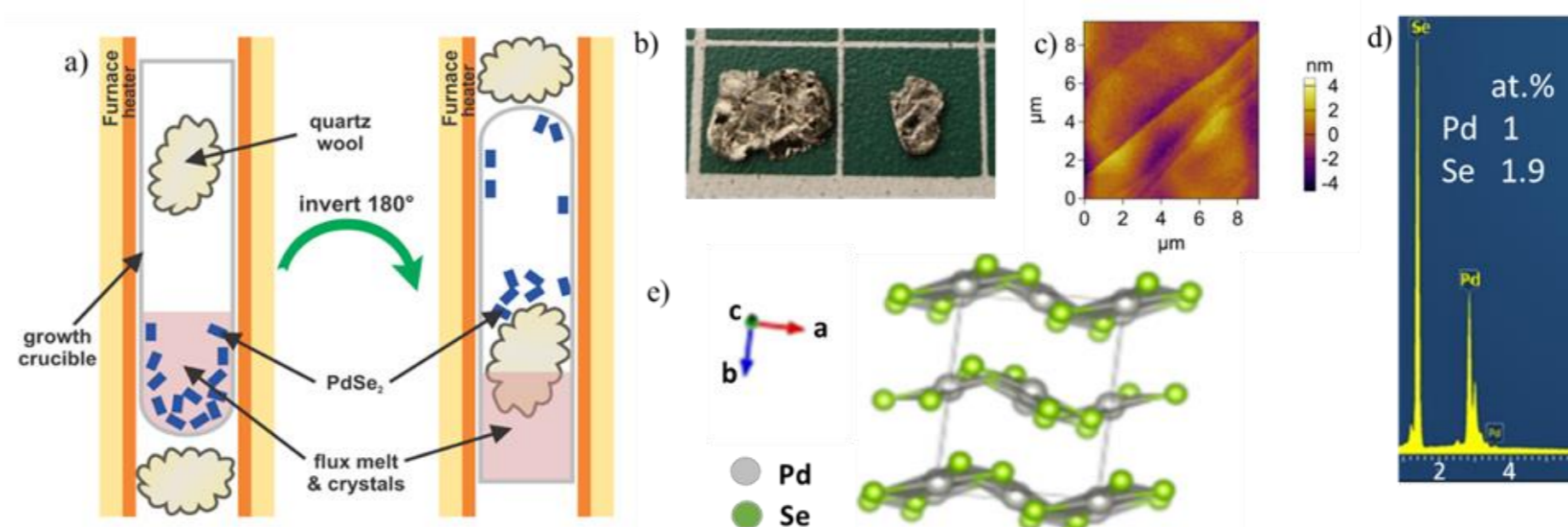


Fig. 2. (a) Diagram of self-flux (high-temperature solution) growth method, (b) photo of shiny crystals of PdSe₂; (c) AFM image of cleaved crystal layered structure; (d) EDS spectrum analysis (atomic %), Pd/Se ratio: 1/1.9 and (e) representation of the crystal structure of PdSe₂ along the b-axis (gray spheres – Pd, green spheres – Se ions).



Fig. 4. SEM image of exfoliated PdSe₂ crystal

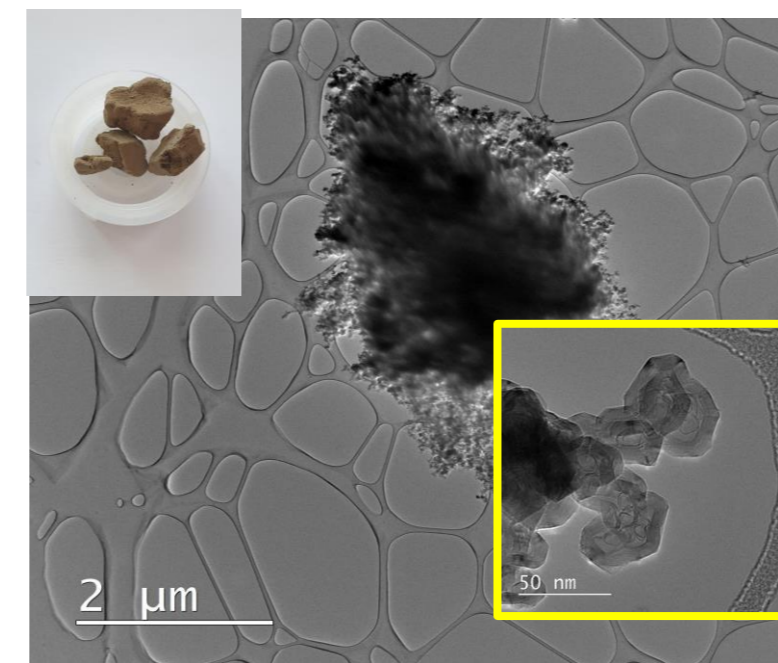


Fig. 5. Bright Field TEM image of GO – inset zoomed image and real image of the as prepared GO

- ✓ The PdSe₂ flakes with sizes from 500 nm were successfully exfoliated from PdSe₂ single crystal grown by self-flux method.
- ✓ The as prepared GO has uniform and homogeneous structure with size of around 50 nm

Electrochemical test set-up

1. Electrochemical Test Cell – Standart three electrode
2. Erf- RHE
3. Ec- Pt-wire
4. Ew- GC or Freudenberg
5. Electrolyte – 25% KOH
6. S_{EW} = 0.28 /1 cm²

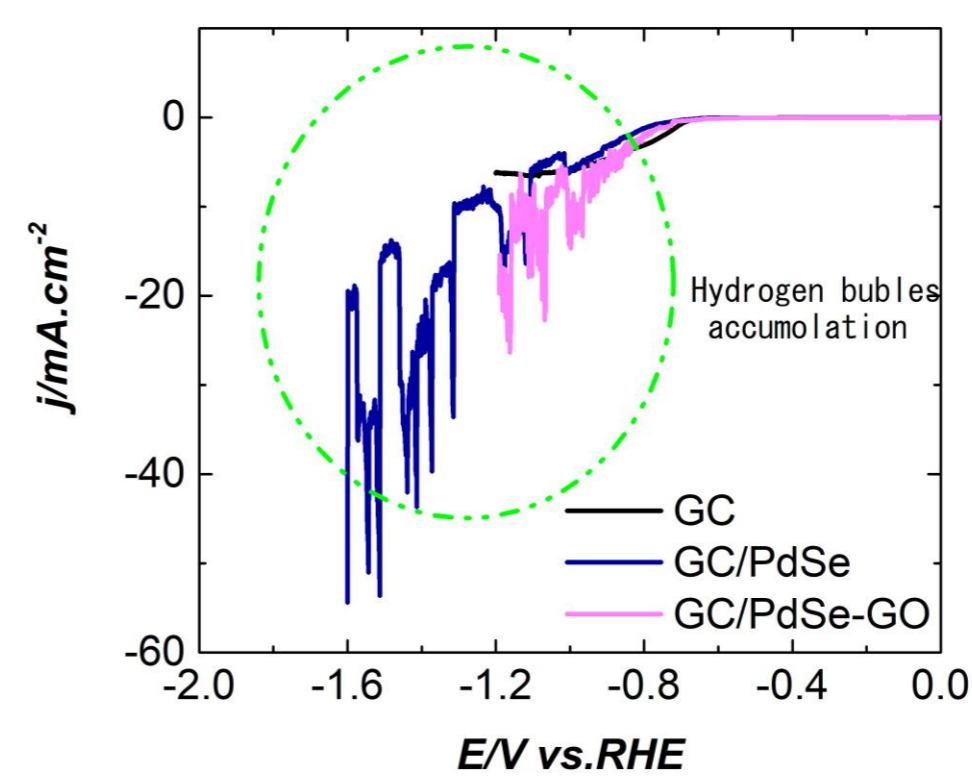


Fig. 6 Cathodic polarisation curves of the PdSe and PdSe GO/GC electrodes in 25% KOH, scan rate 1 mV.s⁻¹

PdSe₂ and PdSe₂-GO in HER

- Pure PdSe₂ shows excellent long-term stability
- PdSe₂-GO composite enhances catalytic activity
- Hydrogen adsorption confirms HER interaction
- PdSe/rGO catalysts on Freudenberg**
- PdSe shifts ORR to positive potentials
- HER activity order: PdSe > rGO > PdSe-rGO > FR

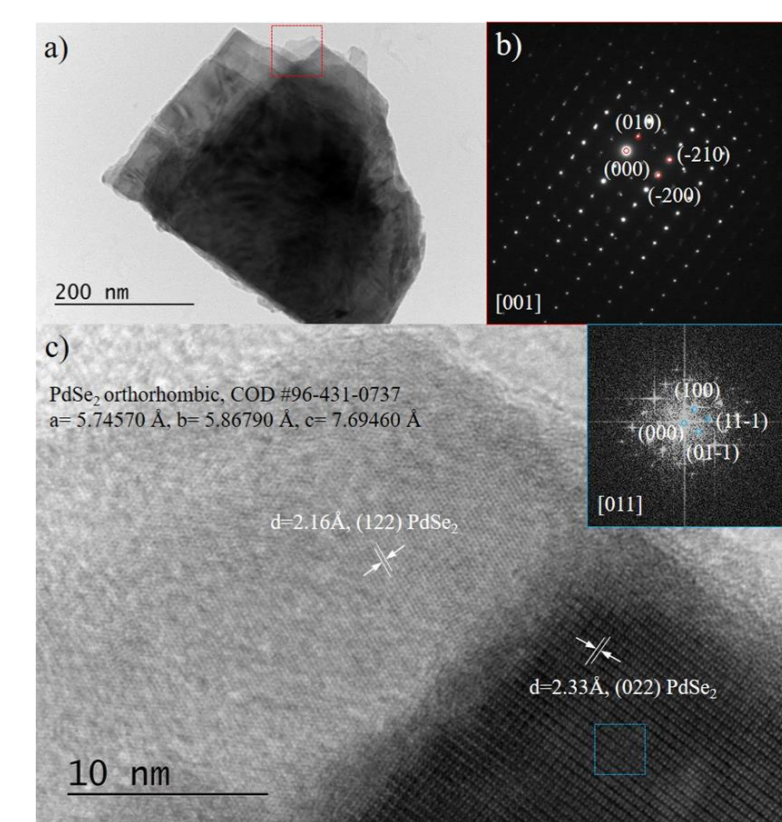


Fig. 3. (a) Bright Field TEM image, (b) the corresponding SAED pattern and (c) HRTEM image and the Fast Fourier Transform pattern from the blue square area (inset) of PdSe₂ crystal

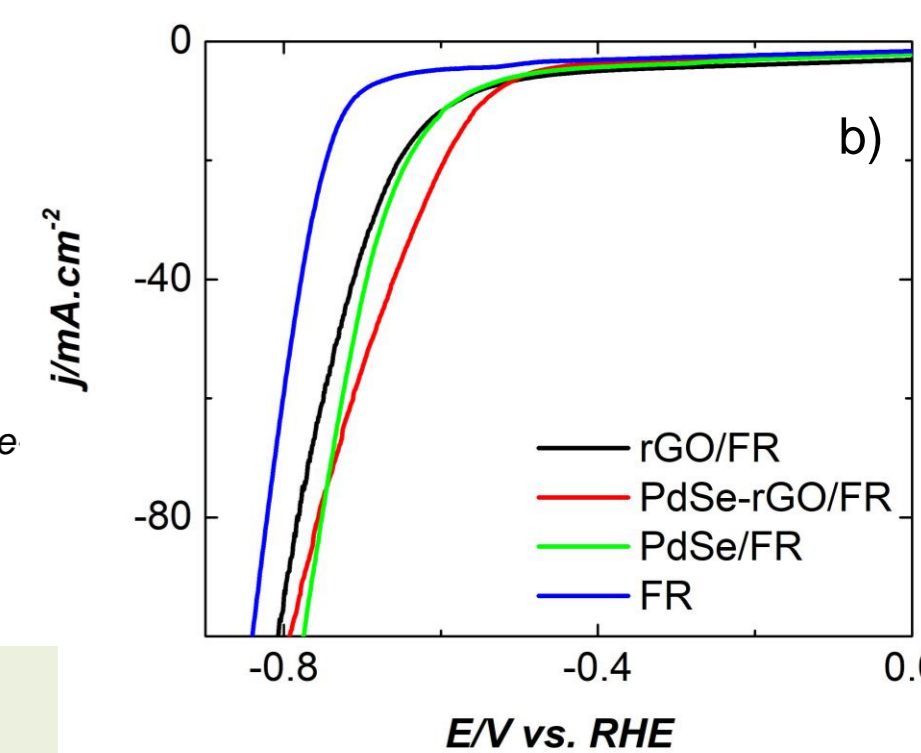
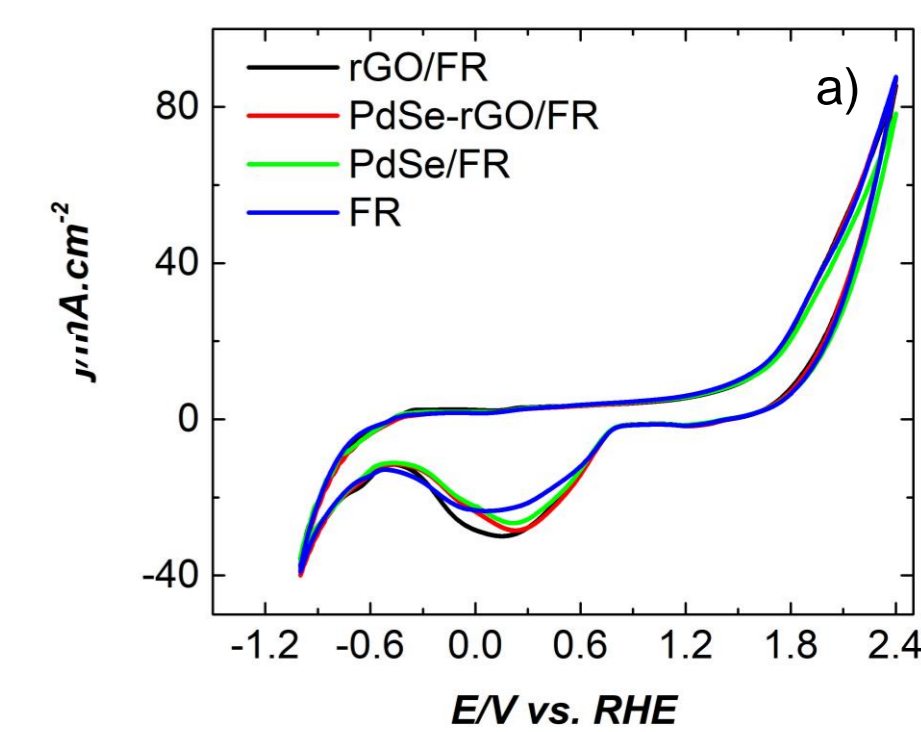


Fig. 7 (a) CV and (b) Cathodic polarisation curves of the PdSe, rGO, PdSe-rGO/ Freudenberg electrodes in 25% KOH, scan rate 1 mV.s⁻¹

CONCLUSION

✦ PdSe₂ and its composites with GO demonstrate promising electrocatalytic properties for HER and ORR in alkaline media.

✦ Pure PdSe₂ exhibits superior HER activity and long-term stability compared to composites and support materials.

✦ The electrochemical reduction of GO in the PdSe₂-GO system shows enhanced conductivity and dispersion, though further optimization is needed to maximize synergy. (also called PdSe₂-rGO)

✦ Compared to reported PdSe-based electrocatalysts (HER overpotential ~540 mV at 10 mA cm⁻² for rGO-PdSe), the investigated PdSe₂ materials display lower overpotentials and improved stability, confirming their advantage as efficient catalysts.

✦ Future work will focus on tuning PdSe₂/rGO ratios and doping with transition metals (Ni, Co, Fe) to further improve performance and reduce Pd content.

Funding: The authors acknowledge the financial support of the Project No. BG16RFPR002-1.014-0009 “Development and Sustainability of the Center of competence HITMOBIL—Technologies and systems for generation, storage and consumption of clean energy”, funded by the Operational Program “Science and Education for Smart Growth” 202114–2020 2027 and co-funded by the EU through the European Regional Development Fund.