

FACILE SYNTHESIS OF V₂O₃/CARBON CORE/SHELL HYBRIDS AS AN ANODE FOR LITHIUM-ION BATTERIES

eM⁺
energy Materials

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INTRODUCTION



Fossil fuels are currently humankind's major energy source. However, their use leads to environmental issues, such as air pollution, CO₂ emission, acid rain, and global warming. A more sustainable alternative is the widespread use of **renewable energy**, such as solar and wind. However, the generation of renewable energy is uncontrollable and intermittent, which mandates reliable electrical energy storage systems for stable and consistent power delivery. Among them, **Lithium-ion batteries (LIBs)** are the most promising candidates because of their high energy density and efficiency.^[1]

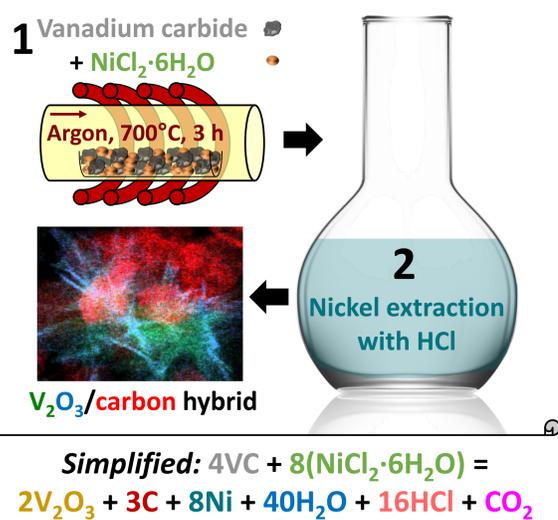


V₂O₃ is a **promising** LIB electrode material but there have been only very few studies exploring V₂O₃ as an **anode** for LIBs so far.^[2-5] The current state of the art assumes that V₂O₃ undergoes structural volume change during cycling; this effect and the **low electronic conductivity explain the poor cycling stability**.^[4] For example, there is a 50-60 % capacity loss of bulk V₂O₃ after 50-100 charge/discharge cycles.^[2, 4]
So how can we overcome these limitations?

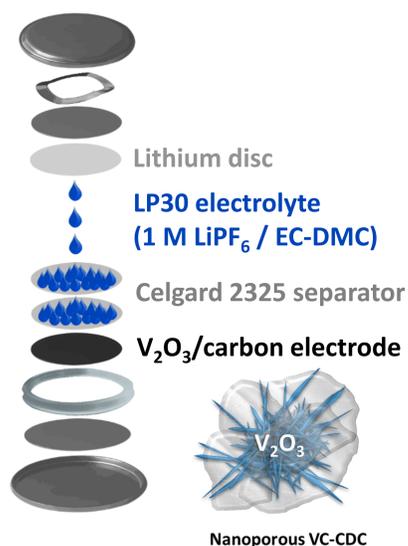
We introduce V₂O₃/C core/shell hybrids where vanadium carbide serves as the precursor for the metal oxide and carbon.^[6]

METHODS

Two-step synthesis

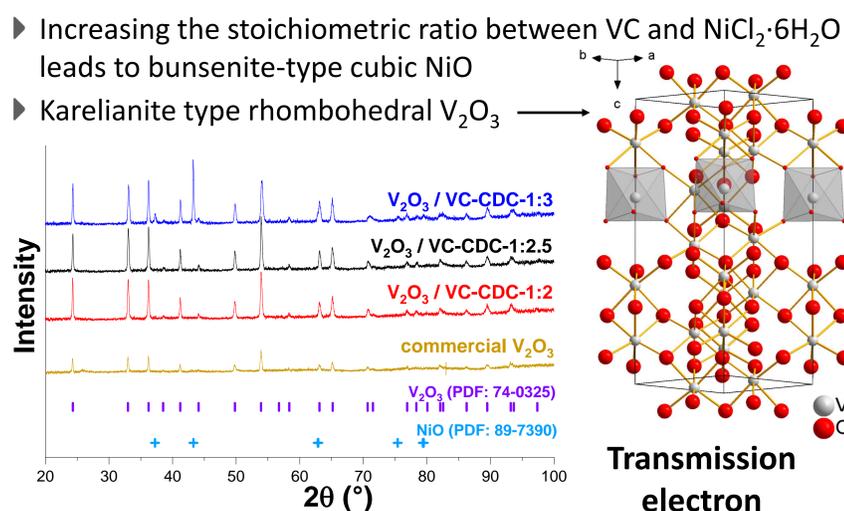


Coin cell assembling



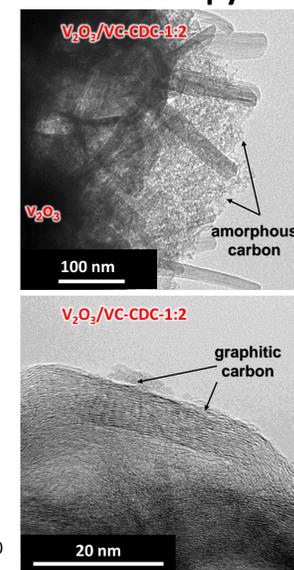
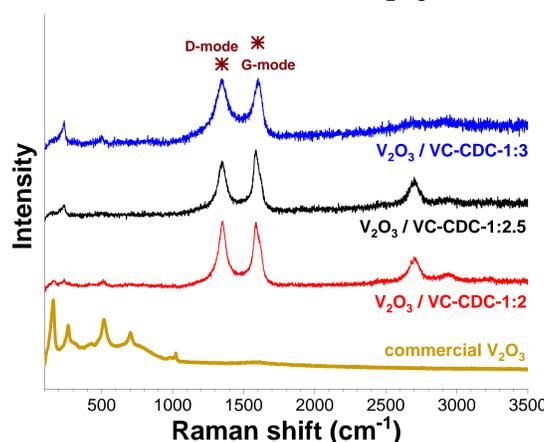
STRUCTURAL CHARACTERIZATION

X-ray diffraction (XRD)



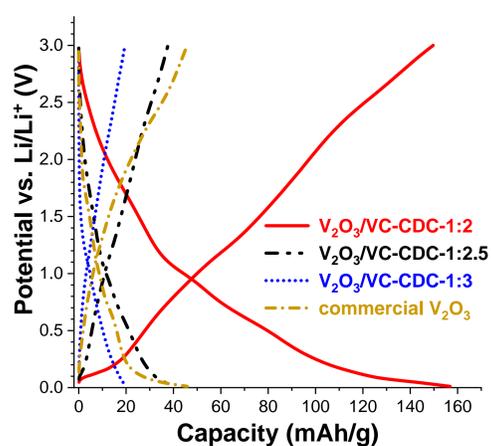
Raman spectroscopy

- Broad peaks of disordered nanocarbon (CDC: carbide-derived carbon)
- Small peaks of nanoscale V₂O₃

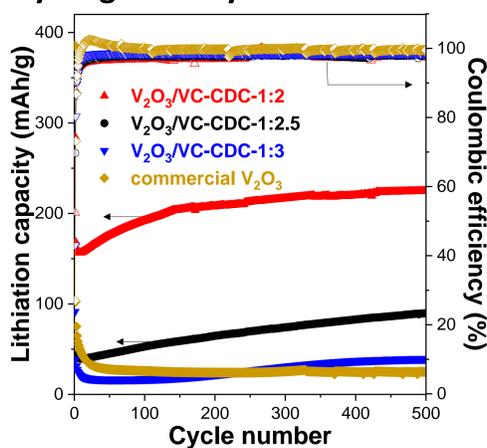


ELECTROCHEMICAL CHARACTERIZATION

Voltage profile



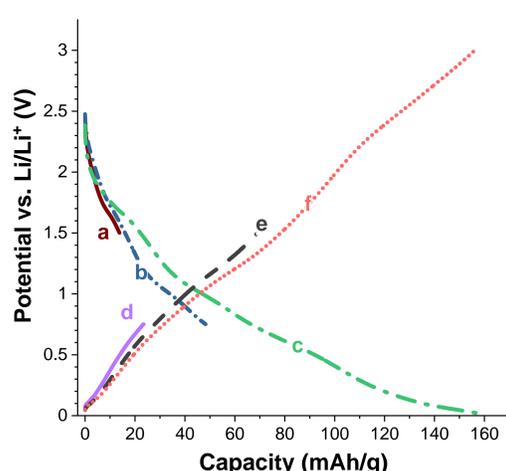
Cycling stability



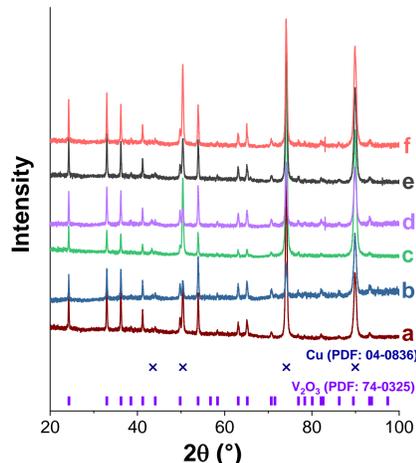
- Highest capacity and best performance stability for V₂O₃/VC-CDC-1:2

POST MORTEM ANALYSIS

Voltage profiles after 40 cycles



XRD patterns at different states



Sample	Phase	Volume (Å ³)
Lithiation at 1.5 V (a)	V ₂ O ₃	298
Lithiation at 0.75 V (b)	V ₂ O ₃	298
Lithiation at 0.01 V (c)	V ₂ O ₃	297
Delithiation at 0.75 V (d)	V ₂ O ₃	297
Delithiation at 1.5 V (e)	V ₂ O ₃	297
Delithiation at 3.0 V (f)	V ₂ O ₃	297

- No significant crystallographic V₂O₃ volume change via electrochemical cycling
- Surprising result per the state of the art in the literature

Raman data analysis after 500 cycles

Sample	State	I _D /I _G ratio
V ₂ O ₃ /VC-CDC-1:2	Initial	1.1
	500 cycles	2.7
V ₂ O ₃ /VC-CDC-1:2.5	Initial	1.1
	500 cycles	2.3
V ₂ O ₃ /VC-CDC-1:3	Initial	2.1
	500 cycles	2.2
commercial V ₂ O ₃	Initial	2.0
	500 cycles	0.9

- VC-CDC becomes structurally more disordered after cycling
- Ergo: carbon plays an active role in the energy storage process

FUNDING

DFG

REFERENCES

- [1] Nitta et al. (2015), *Mater Today* 18, 252-264.
- [2] Jiang et al. (2015), *Appl. Mater. Interfaces* 7, 1595-1601.
- [3] Shi et al. (2015), *J. Power Sources* 275, 392-398.
- [4] Sun et al. (2011), *J. Power Sources* 196, 8644-8650.
- [5] Yan et al. (2016), *J. Power Sources* 329, 148-169.
- [6] Budak et al. (2019), *Batteries&Supercaps*, 2, 74-82.

CONCLUSIONS

- Using an optimized precursor ratio, we synthesized a V₂O₃/VC-CDC hybrid material with a core/shell particle architecture.
- The material afforded a capacity of 160 mAh/g with high performance stability up to 500 cycles.