

# Nanostructured Metal Oxide Anodes

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# Overview



## Timeline

- October 1, 2007
- September 30, 2010
- 50% complete

## Budget

- Total project funding  
FY08: \$250K, FY09: \$350K
- Project lead: Anne Dillon

## Barriers

- Cost: developing metal oxide based anodes from abundant, inexpensive metals
- Energy density: improvements in both gravimetric and volumetric energy densities have been demonstrated
- Safety: Anodes operate at higher potential relative to Li metal than graphite, eliminating the risk of Li plating
- Lifetime: Durable and reversible cycling has been achieved

## Partners

- M.M. Thackeray and S-H. Kang, Argonne
- M.S. Whittingham, SUNY-Binghamton
- A. Greenshields, fortu
- S-H. Lee, Univ. of Colorado
- S.M. George, Univ. of Colorado
- A. Pesaran, NREL

# Objectives



**The ultimate goal of this activity is to develop optimized metal oxide nanostructured electrode materials to enable high-performance, durable, and affordable Li-ion batteries for power-assist HEVs and PHEVs that meet the DOE/FreedomCAR targets.**

- Optimize  $\text{MoO}_3$  nanoparticle electrodes in coin cell configuration and compare to previous results for electrophoresis deposited thin film  $\text{MoO}_3$  electrodes.
- Demonstrate a full cell with an  $\text{MoO}_3$  anode and state-of-the-art cathode with a high energy density and stable cycling performance.
- Employ first principles calculations to obtain better understanding of Li-insertion processes and for the prediction of new materials.
- Synthesize  $\text{MoO}_2$  nanoparticles to test theoretical prediction that Li will be extracted at a lower potential ( $\sim 1$  V).
- Explore possibility of other metal oxide nanostructures made from even less expensive starting materials.

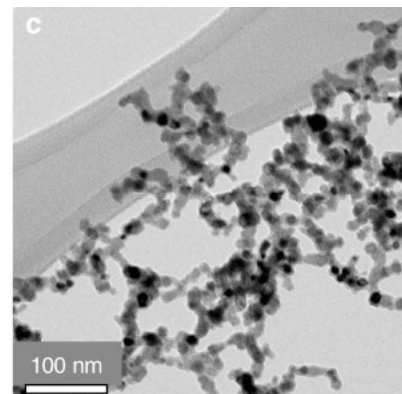
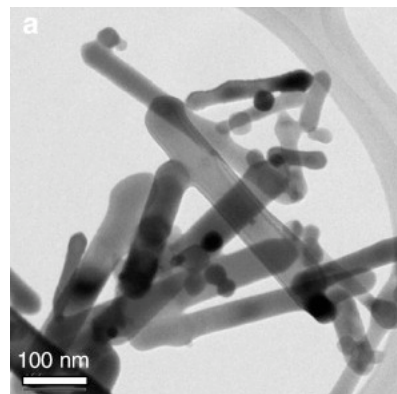
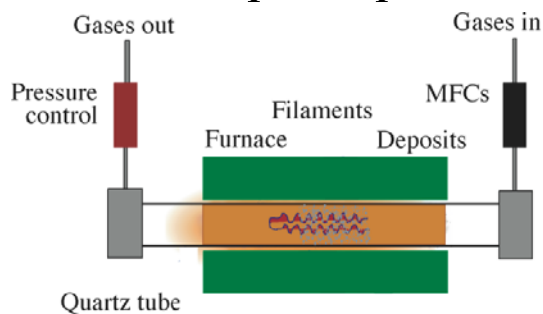
# Milestones



- Sept 2008-report on optimization of  $\text{MoO}_3$  thick electrodes tested in a coin cell configuration, complete. (In this report a reversible capacity of  $\sim 1050$  mAh/g was demonstrated with good cycling and rate capability. This high capacity represents a 60% improvement compared to the thin film  $\text{MoO}_3$  electrodes, 630 mAh/g)
- July 2009-report on optimization of  $\text{MoO}_3$  anodes in a full cell with cathodes supplied by ANL. (Full cell data for the  $\text{MoO}_3$  anodes coupled with both  $\text{Li}_{1.05}\text{M}_{0.95}\text{O}_2$ ,  $\text{M} = \text{Ni}_{1/3}$ ,  $\text{Co}_{1/3}$ ,  $\text{Mn}_{1/3}$  and the state-of-the-art lithium rich cathode  $0.5\text{Li}_2\text{MnO}_3 \cdot 0.5\text{Li}(\text{Mn}_{0.31}\text{Ni}_{0.44}\text{Co}_{0.25})\text{O}_2$  is presented here.)

# Approach

- MoO<sub>3</sub> nanoparticles (nano-rods and nanospheroides) are produced using hot-wire chemical vapor deposition (HWCVD) at different reactor pressures.



Electrophoresis



Material slurry

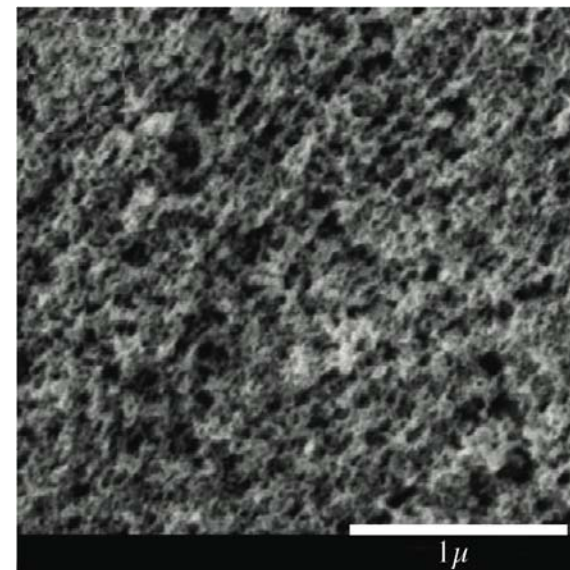
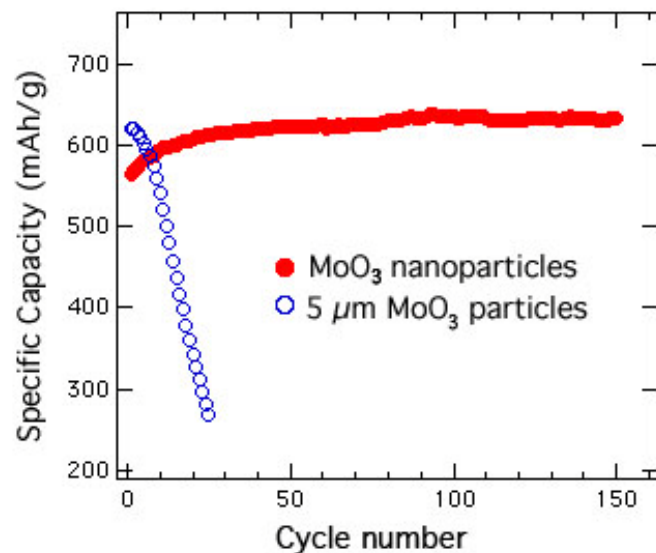


Coin Cell

- Thin film battery electrodes (2-3  $\mu\text{m}$ ) have been fabricated with novel electrophoresis.
- Thick film electrodes ( $\sim 35 \mu\text{m}$ ) for coin cell testing have been optimized versus a Li counter electrode by varying: binder/conductive additive composition and electrode pretreatment.
- Full cell has been also optimized with ANL cathodes.

# Previously Reported

## Thin Film Electrodes by Novel Electrophoresis



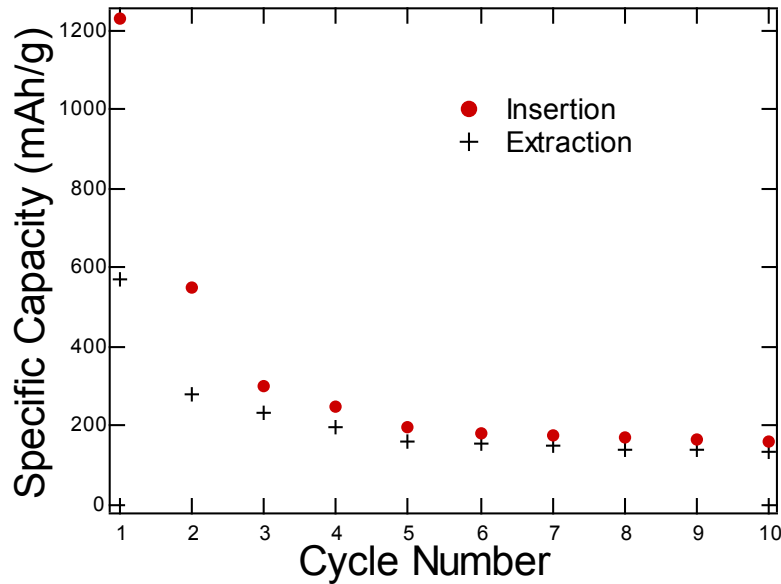
- Porous thin film without binder or conductive additive obtained after electrophoresis.
- Improved durable capacity ( $\sim 600$  mAh/g) found when using the thin film as anode and cycling between 3.0- 0.005 V.

# Technical Accomplishments

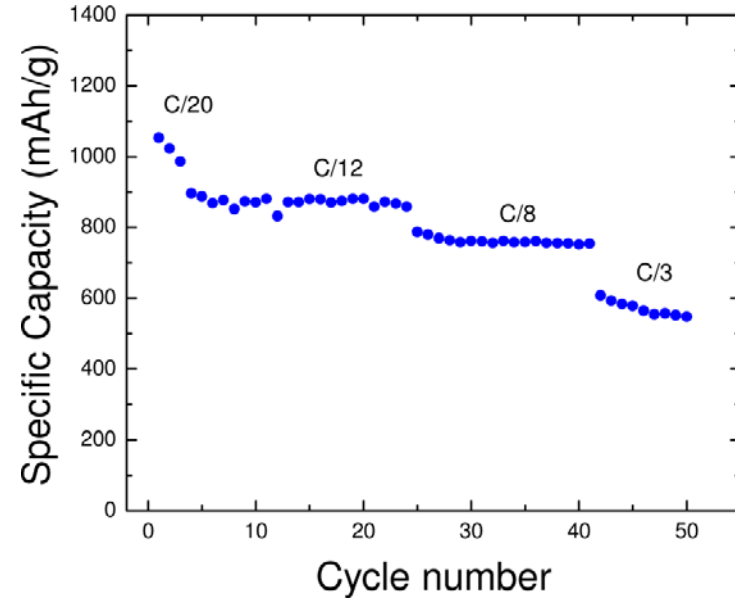
## Optimization by Pre-heat Treatment



70:15:15, 150 °C



70:15:15, 250 °C

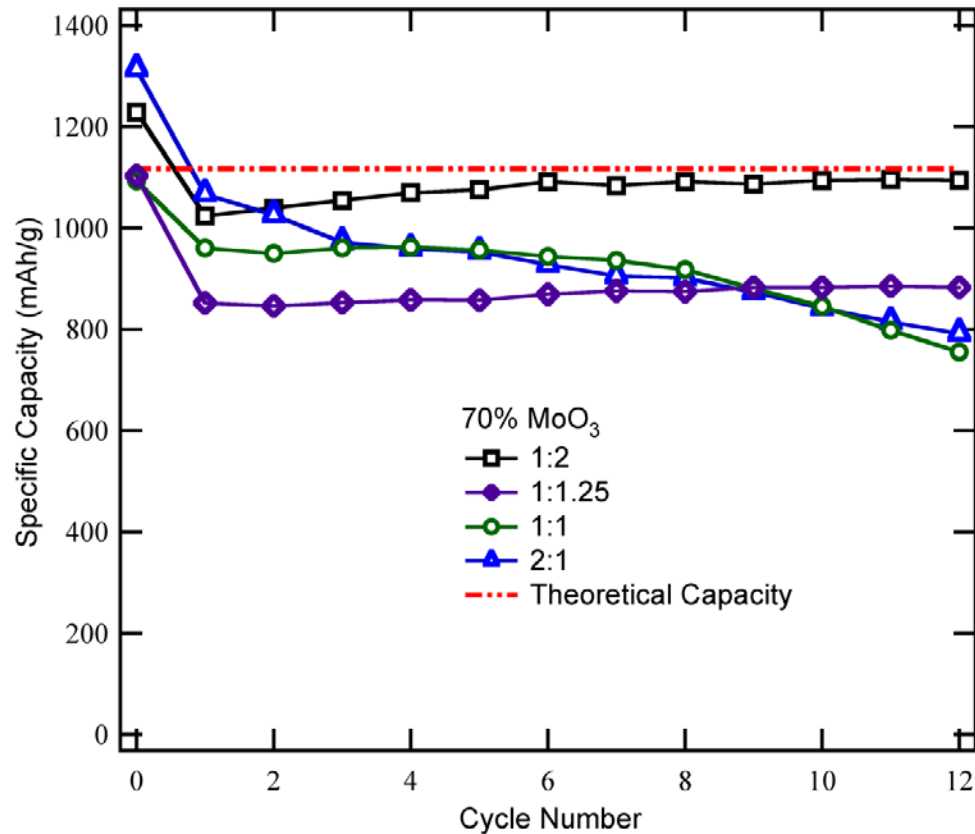


- Slightly less reversible capacity initially reproduced in coin cell configuration with a ratio of 70:15:15 (active material: acetylene black: polyvinylidene fluoride).
- Better reversibility achieved by pre-heating the electrode at 250 °C:
  - ~ 600 mAh/g was observed at C/3;
  - ~ 400 mAh/g delivered at 2C.

# Technical Accomplishments



## Optimization by Varying the Ratio of AB : PVDF



- Polymer rich electrodes provide continuous adhesion through the film.
- Maximum cycling capacity of ~1050 mAh/g (theoretical 1170 mAh/g) achieved at a ratio of 70:10:20 (MoO<sub>3</sub>:AB: PVDF).

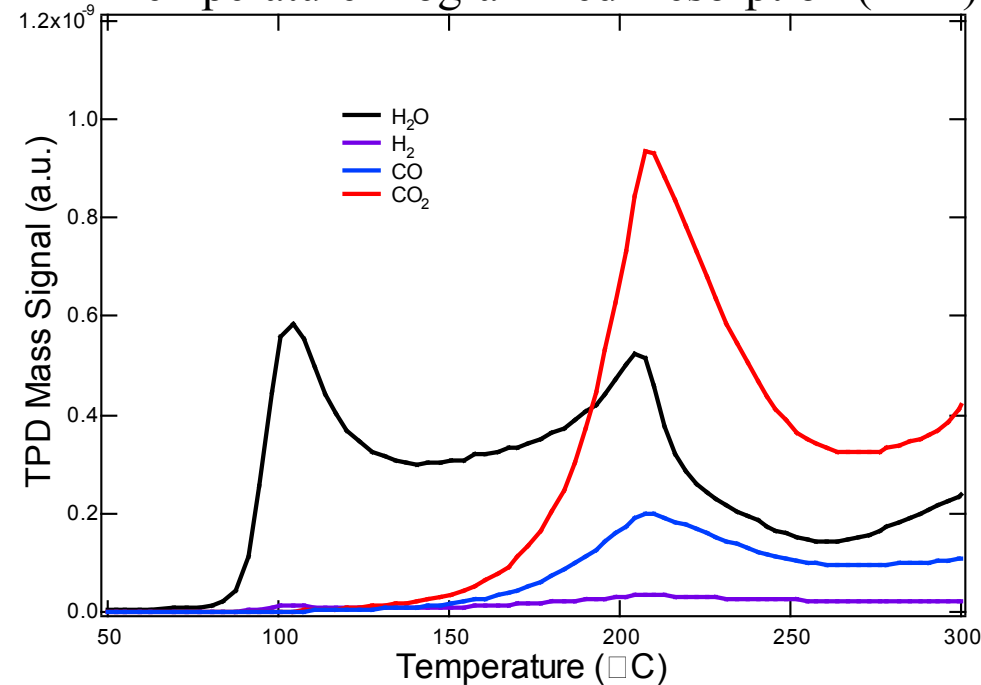


# Technical Accomplishments

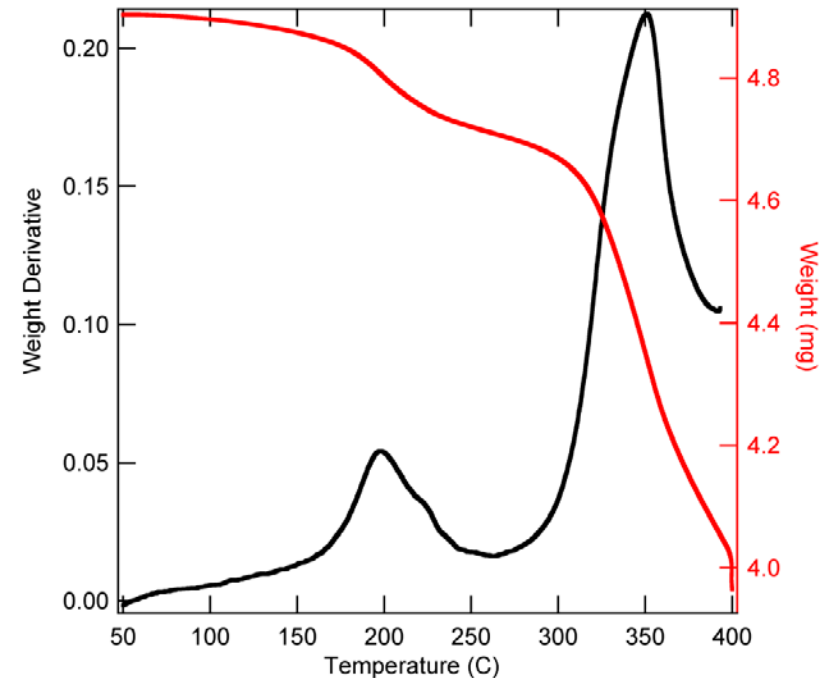
## Explanation of Pre-heat Requirements



### Temperature Programmed Desorption (TPD)



### Thermalgravimetric Analysis (TGA)



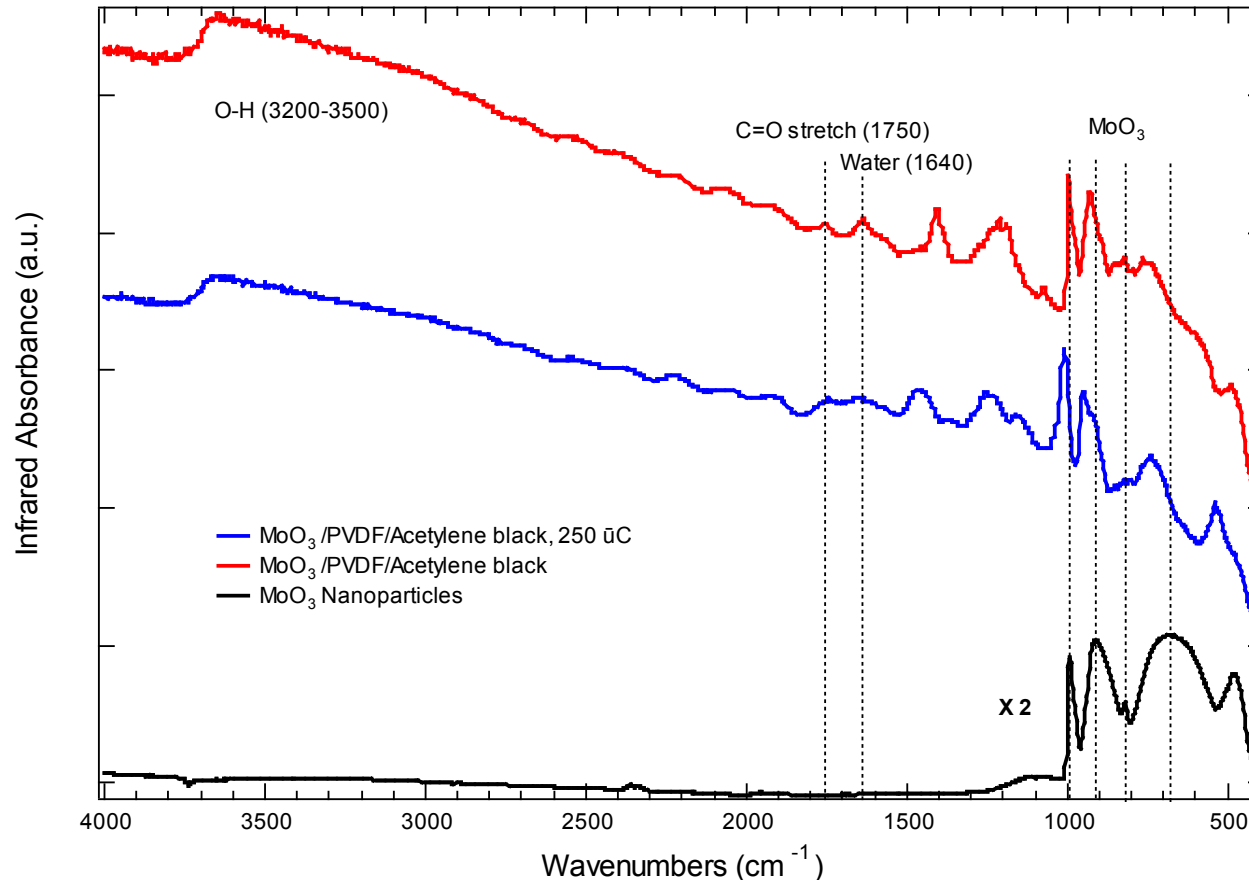
- Water desorption from the electrode observed at a high temperature ( $> 200$  °C).
- CO<sub>2</sub> species are also observed at a higher temperature perhaps due to oxidation of the acetylene black.
- Polymer decomposition observed at a surprising low temperature (300 °C).
- Early decomposition may be catalyzed by nanostructured MoO<sub>3</sub>

# Technical Accomplishments



## Explanation of Pre-heat Requirements

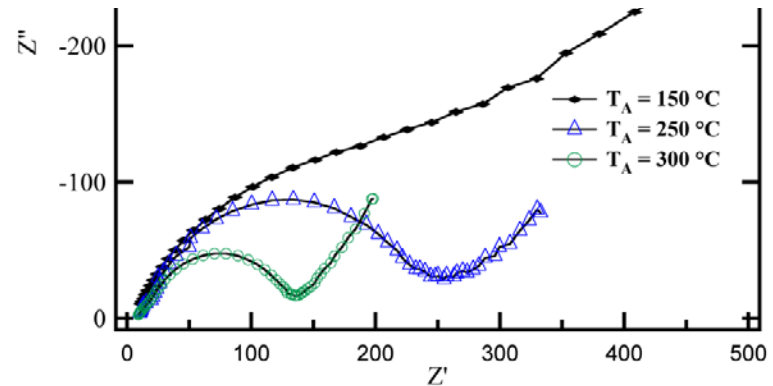
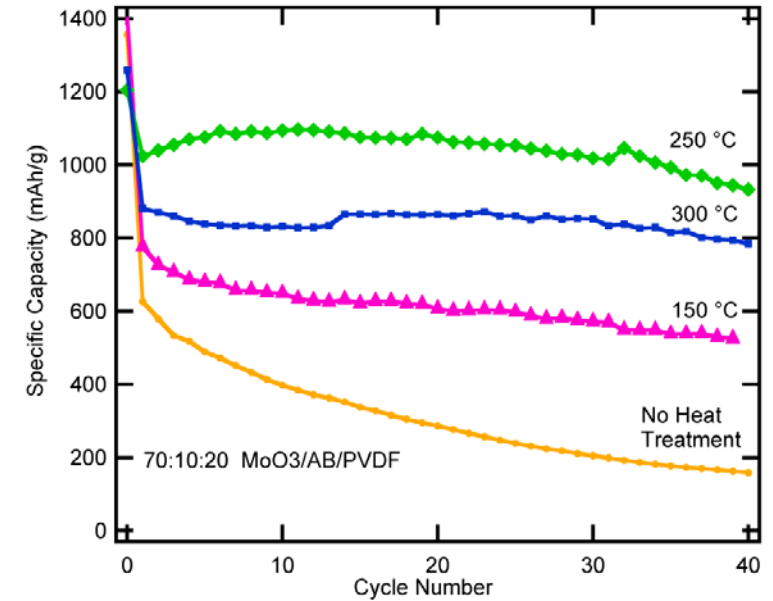
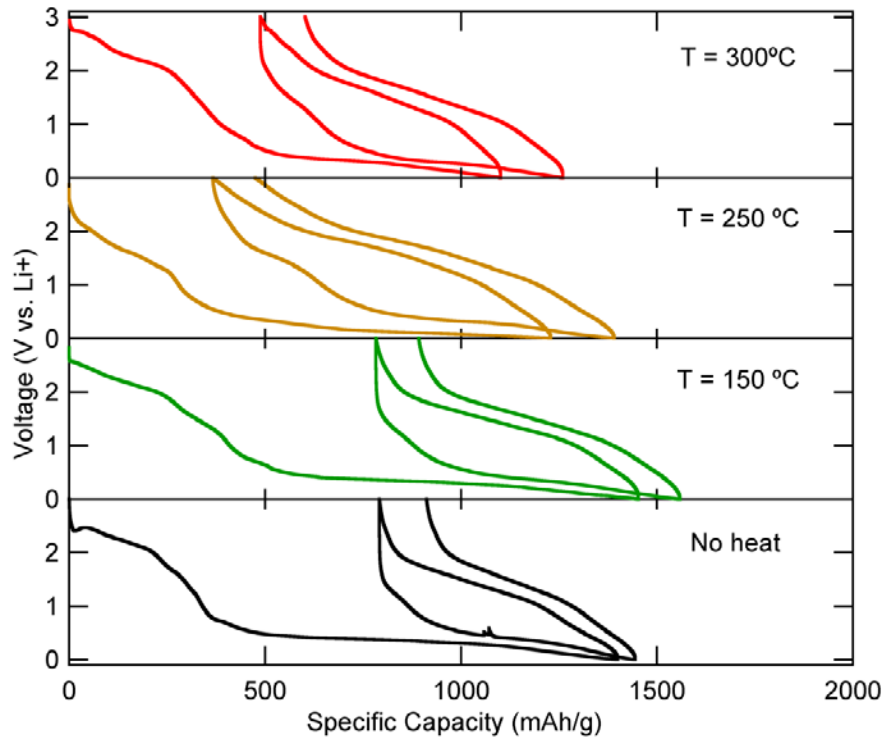
Results confirmed by Infrared Spectroscopy (IR)



- H<sub>2</sub>O/OH originated from acetylene black, PVDF and NMP solvent.
- Weakly bound water removed by pre-heat treatment.
- Presence of bound water is one reason for irreversibility in cycling without pre-heating treatment.

# Technical Accomplishments

## Optimization by Pre-heat Treatment



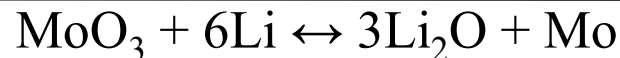
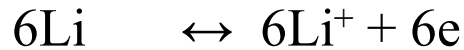
- Highly improved capacity of 1050 mAh/g is achieved by using a ratio of 70:10:20 and pre-heating at 250 °C.
- Electrical resistance steadily decreases with increase of temperature.
- Decreased capacity at 300 °C likely due to the binder breakdown and isolation of certain particle clusters.

# Technical Accomplishment



## Nano-sized $\text{Li}_x\text{MoO}_3$ : Displacement redox reaction?

Displacement redox reaction\* for  $\text{MoO}_3$  nanoparticles :



What is size distribution of Mo clusters?

First-principles molecular dynamics (FPMD)

- $(\text{Li}_4\text{MoO}_3)_{36}$  &  $(\text{Li}_6\text{MoO}_3)_{36}$
- Start from uniformly lithiated alpha phase of  $\text{MoO}_3$
- $T = 600$  K (to speed up the MD simulations)
- VASP code

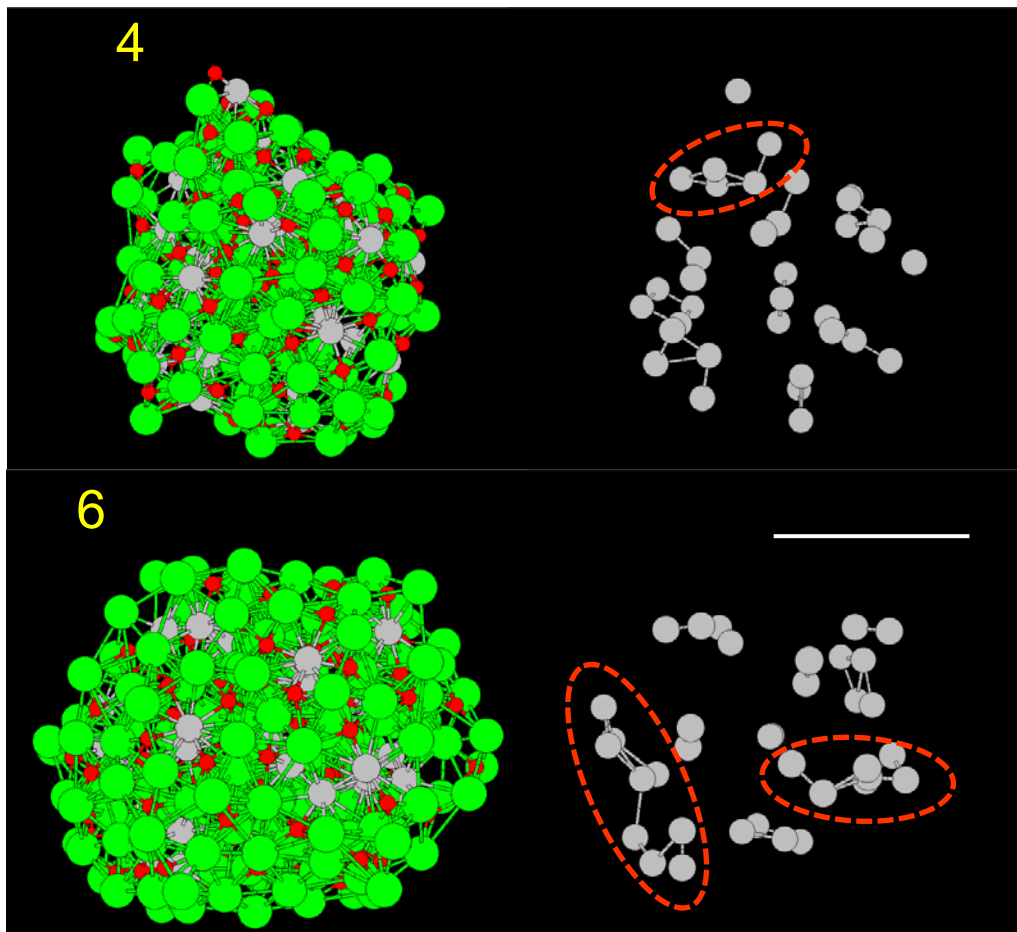
\*P. Poizot et al., Nature 407, 496 (2000)

# Technical Accomplishments



FPMD simulations for  $\text{Li}_x\text{MoO}_3$  nanoparticles ( $x = 4, 6$ )

$\text{Mo}_n$	1	2	3	4	5	6	7	8	9
$(\text{Li}_4\text{MoO}_3)_{36}$	4	1	3	4	1	-	-	-	-
$(\text{Li}_6\text{MoO}_3)_{36}$	-	6	-	2	-	-	1	-	1



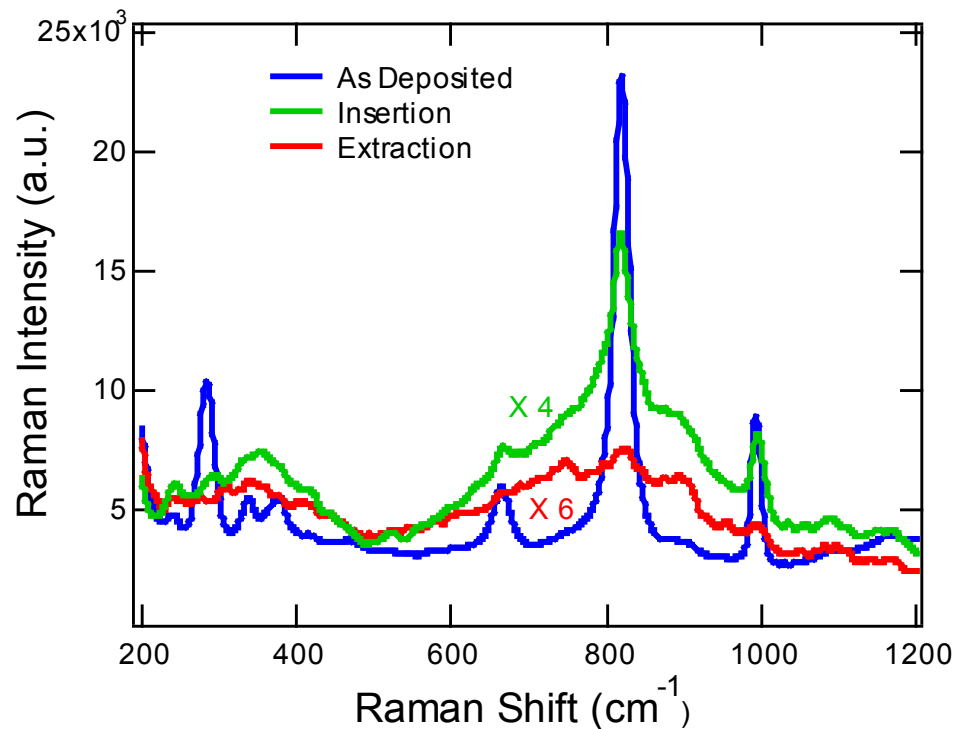
Small clusters of  $\text{Mo}_n$  are easily formed within the nanoparticle.

The size  $n$  of the Mo cluster ranges from 2 to 9. The Mo nanoclusters are small enough to enable reversible Li insertion/desertion.

Our theoretical results support a displacement redox reaction which involves the formation and decomposition of metal nanoclusters.

# Technical Accomplishments

## *In Situ* Raman Showing Disordered Structure after Cycles

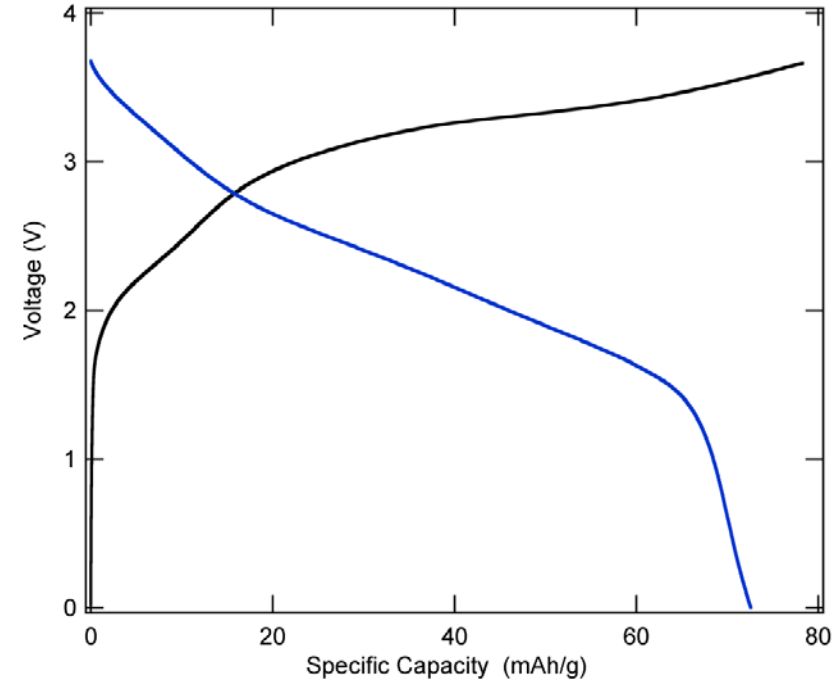
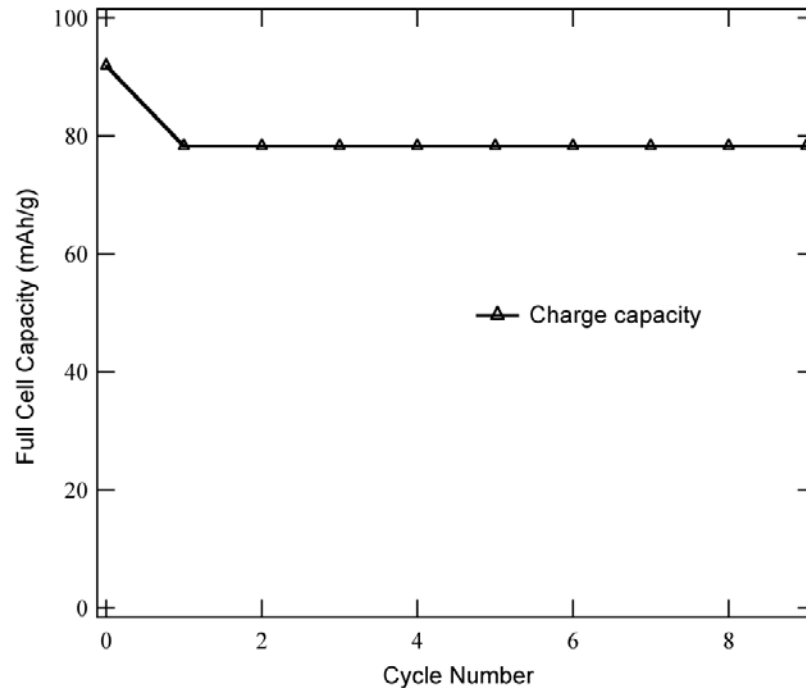


*In situ* Raman confirms significant loss in structural order in first insertion cycle consistent with molecular dynamics simulations.

# Technical Accomplishments

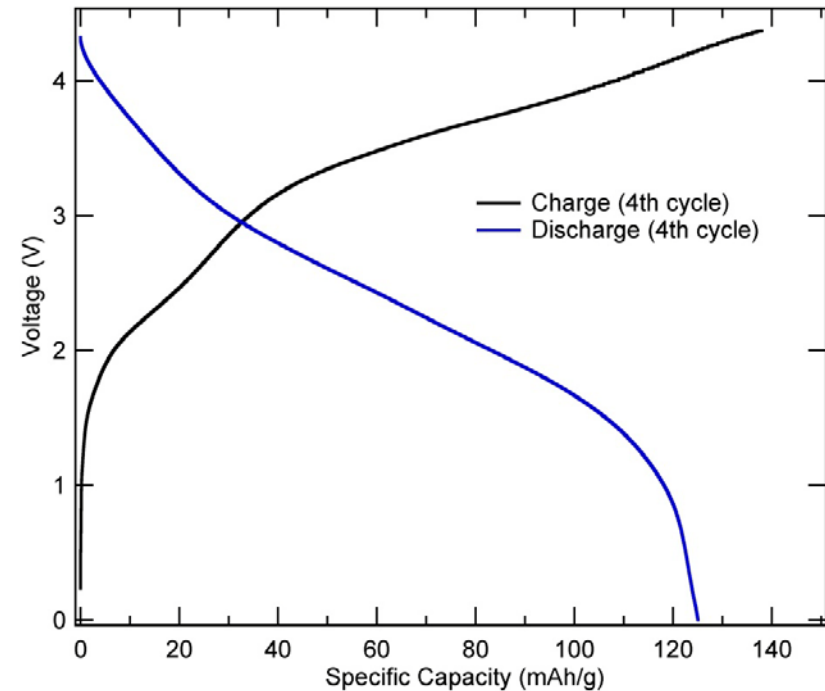
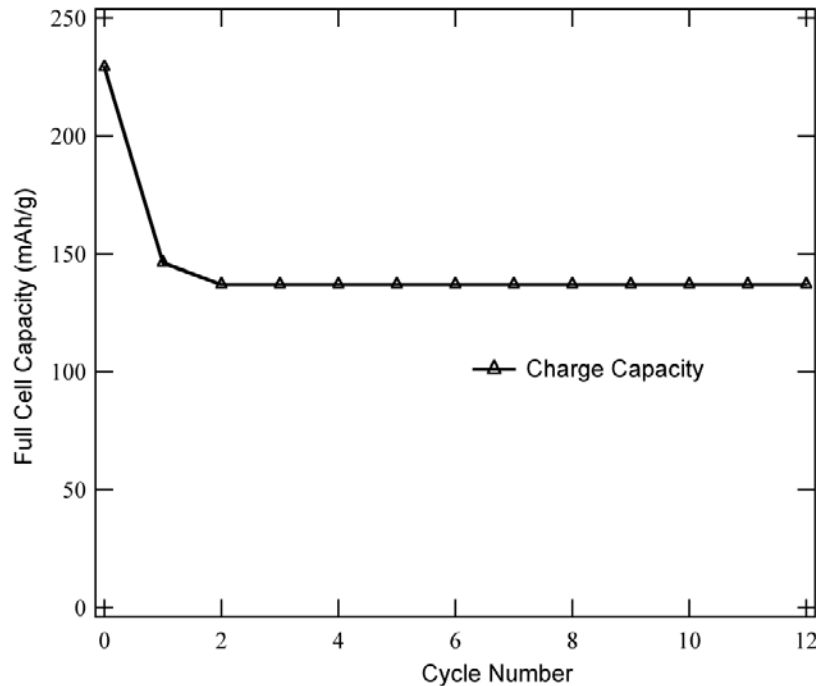


## Full Cell Testing Using ANL Cathode



- Full cell capacity of  $\sim 80$  mAh/g achieved when cycling between 4.0-1.0 V by coupling with Gen 2 cathode obtained from M. Thackeray and S-H. Kang (ANL).
- Cell contains 12 mg cathode material and 2.5 mg anode material.
- In the full cell  $\text{MoO}_3$  has a reversible capacity of  $\sim 677$  mAh/g

## Full Cell Testing Using ANL Cathode



- Stable capacity of 140 mAh/g (commercial capacity: ~80 mAh/g) when cycling between 4.0-0.01 V at a constant capacity and coupled with lithium rich cathode (250mAh/g) obtained from M. Thackeray and S-H. Kang at Argonne
- Cell contains 7.4 mg cathode material and 1.6 mg anode material.
- In the full cell  $\text{MoO}_3$  has a reversible capacity of ~ 776 mAh/g.

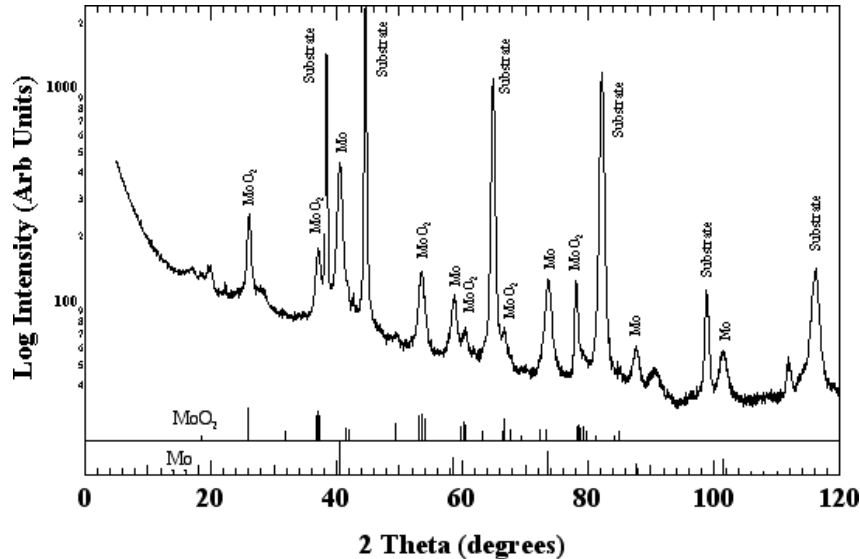


# Technical Accomplishments

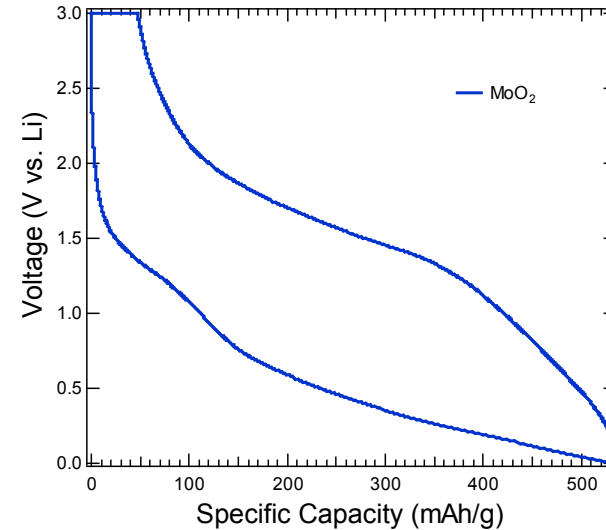
## HWCVD Production of Nano-MoO<sub>2</sub>



### XRD MoO<sub>2</sub>



### Voltage Profile

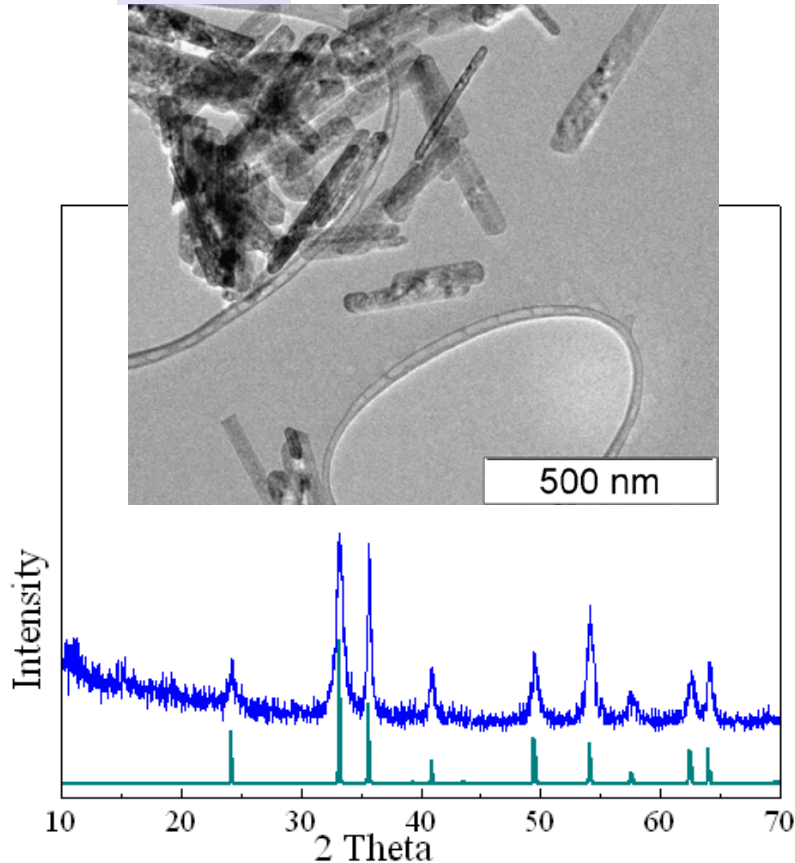


- Previously theoretical predictions indicated crystalline MoO<sub>2</sub> would have a lower lithium extraction potential.
- Nano-MoO<sub>2</sub> was produced by the modified HWCVD process.
- However, upon cycling a thin film of the MoO<sub>2</sub> material, the voltage profile was not significantly different from that of MoO<sub>3</sub>.
- The discrepancy with the theory may be attributed to the fact that the nanoparticles become highly disordered upon cycling, with the calculations performed for crystals.

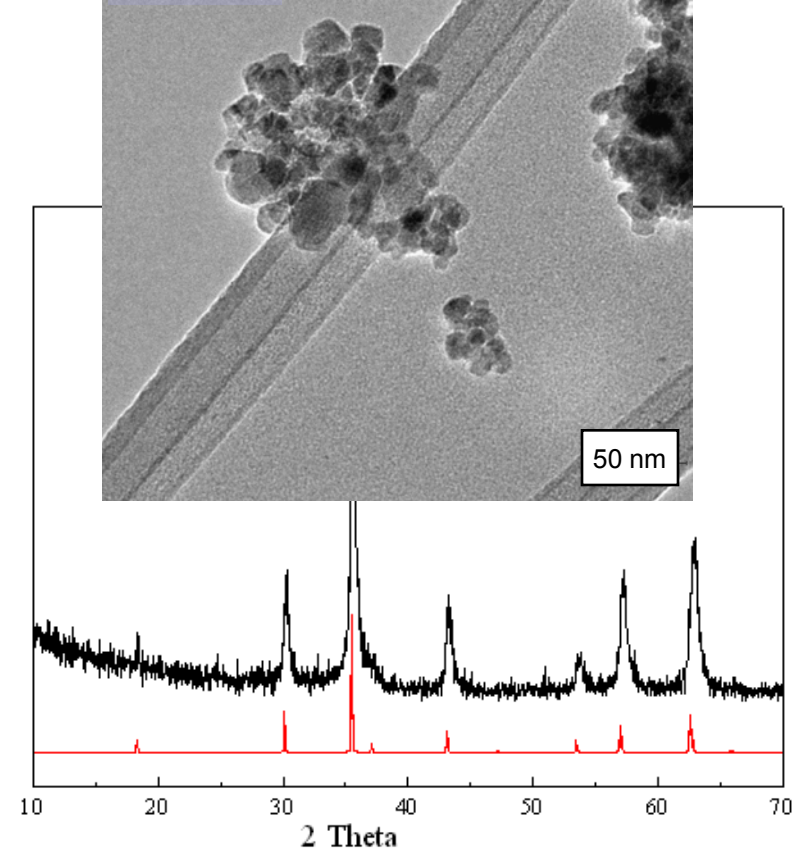
# Recent Development

## Synthesis of Iron Oxide by Hydrothermal Technique

$\text{Fe}_2\text{O}_3$



$\text{Fe}_3\text{O}_4$

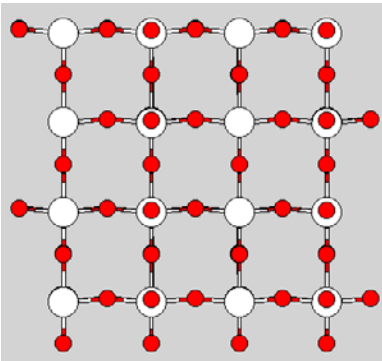


- $\text{Fe}_2\text{O}_3$  nanofibers (40-50 nm width) has been produced using hydrothermal process followed by post-heat at 300 °C.
- $\text{Fe}_3\text{O}_4$  nanoparticles (10-20 nm) obtained by using reducing agent in hydrothermal process.
- Iron oxides allow for a more economical system.

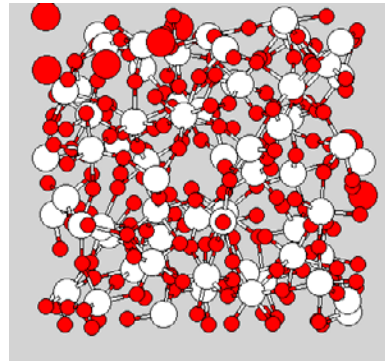
# Recent Development

## Oxygen Vacancy: $\text{WO}_3$ , $\text{MoO}_3$ , and $\text{Fe}_2\text{O}_3$

### Atomic structure

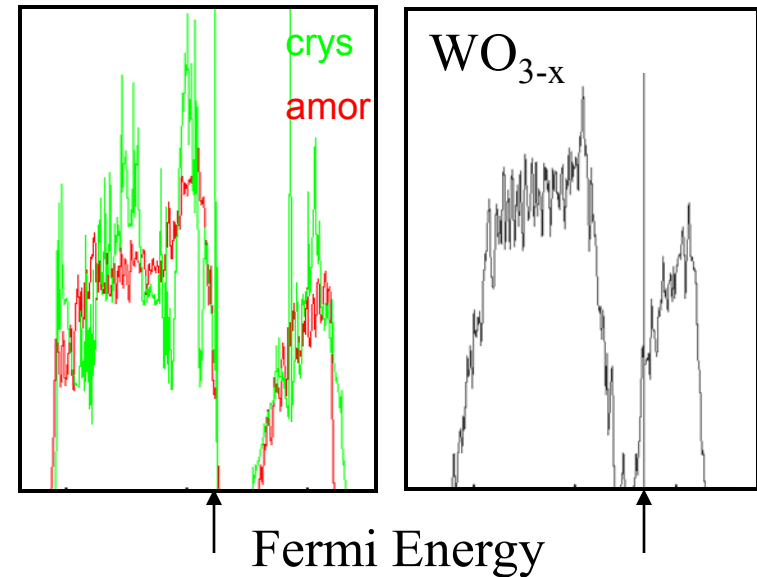


Crystalline  $\text{WO}_3$



Amorphous  $\text{WO}_3$

### Electronic structure



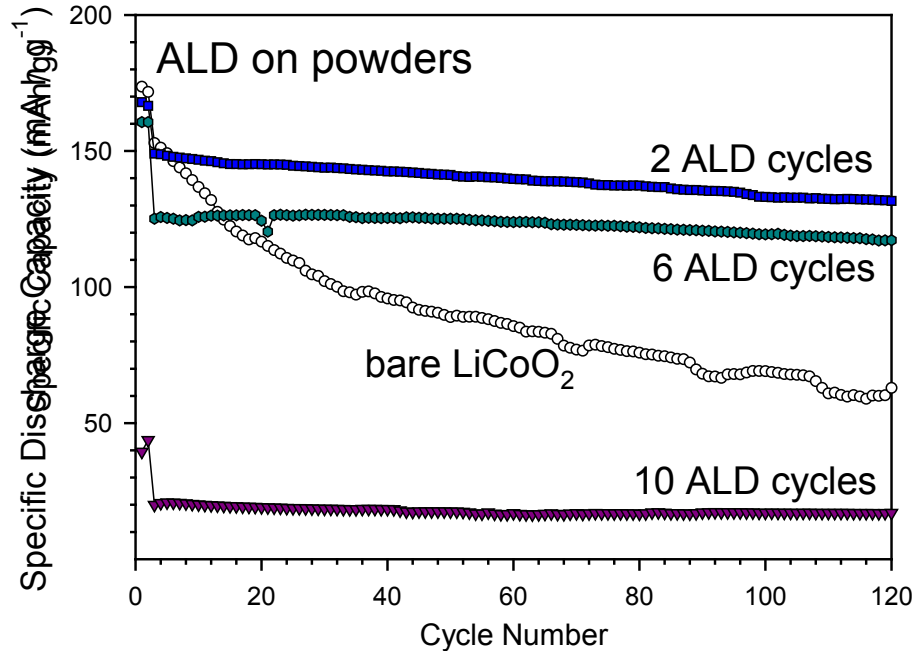
- Oxygen vacancy creates defect states near the conduction band of transition metal oxides such as  $\text{WO}_3$ ,  $\text{MoO}_3$ , and  $\text{Fe}_2\text{O}_3$ .
- For sub-stoichiometric amorphous  $\text{WO}_{3-x}$  the conduction band is populated to a larger extent.
- The conduction band filling will lower the potential inserted Li.
- By creating oxygen vacancies and substoichiometric amorphous samples, we can reduce Li potentials of  $\text{MoO}_3$  and  $\text{Fe}_2\text{O}_3$  to make them more suitable anodes.

# Recent Development

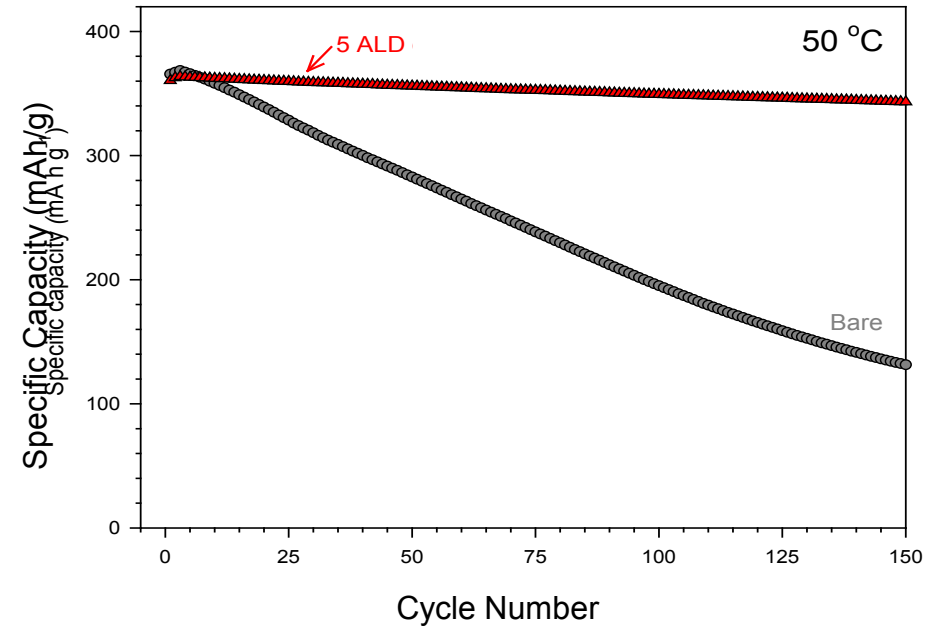


## Atomic Layer Deposition (ALD) Improves Durability

### ALD on $\text{LiCoO}_2$



### ALD on Natural Graphite



Improved cyclability achieved for both cathode and anode by applying a thin ALD coating.

ALD coatings eliminate SEI and surface reactions that cause degradation.

# Proposed FY 09 Future Work



- Optimize full cells with ANL cathodes to improve durable capacity and rate capability (July 2009 Milestone).
- Work with fortu (Switzerland) to develop high-voltage cell.
- Perform theoretical calculations to understand the hysteresis of the charge/discharge for the  $\text{MoO}_3$  nanoparticles. Use theoretical calculations to predict composition and orientation of economical oxides nanoparticles with more desirable voltage profiles.
- Synthesis of alternative nanostructures made from abundant elements, such as  $\text{Fe}_2\text{O}_3$ ,  $\text{Fe}_3\text{O}_4$ , and  $\text{MnO}_2$  will be explored. Inexpensive synthesis routes—including HWCVD, hydrothermal techniques, and electrodeposition—will be employed.
- Apply a protective ALD coating on graphite nanoparticles to eliminate surface degradation mechanisms and improve rate capability.

# Conclusions



- Capacity of  $\text{MoO}_3$  anode has been increased to  $\sim 1050$  mAh/g by optimizing the coin cell configuration. TPD, TGA, and IR employed to facilitate these optimizations.
- Theoretical calculations were performed to explain the mechanism for the increased Li-insertion observed in the coin cell testing.
- The  $\text{MoO}_3$  anode has been successfully paired with two Argonne cathodes:  $\text{Li}_{1.05}\text{M}_{0.95}\text{O}_2$ ,  $\text{M} = \text{Ni}_{1/3}, \text{Co}_{1/3}, \text{Mn}_{1/3}$  and the state-of-the-art lithium rich cathode  $0.5\text{Li}_2\text{MnO}_3 \cdot 0.5\text{Li}(\text{Mn}_{0.31}\text{Ni}_{0.44}\text{Co}_{0.25})\text{O}_2$
- In-situ Raman capabilities, established this year, show that  $\text{MoO}_3$  nanoparticles become highly disordered in the initial cycle.

	Gravimetric Capacity (mAh/g)	Volumetric Capacity (mAh/cm <sup>3</sup> )	Full Cell Capacity (mAh/g)
FY08	630	2200	--
FY09	1050	800	140
Commercial	350 (graphite)	770 (graphite)	80 (graphite/LiCoO <sub>2</sub> ) (J.Power Sources 88, p.237, 2000)

# Acknowledgments

- DOE OVTP Support
  - David Howell



- NREL Program/Project Guidance
  - Ahmad Pesaran
  - Terry Penney

# Publications

- S-H. Lee, Y-H. Kim, R. Deshpande, P.A. Parilla, E. Whitney, D.T. Gillaspie, K.M. Jones, A.H. Mahan, S. Zhang, and A.C. Dillon, “Reversible Lithium Ion insertin in Molybdenum Oxide Nanoparticles,” *Advanced Materials*, 2008 (20), 3627.
- “Electrochemical Reactivity of Ball-milled  $\text{MoO}_{3-y}$  Powders as Anode for Lithium Secondary Batteries,” Yoon S. Jung, Sangkyoo Lee, Dongjoon Ahn, Anne C. Dillon, and Se-Hee Lee, *J. Power Sources*, 188, 286 (2009).
- A.C. Dillon, A.H. Mahan, R. Deshpande, P.A. Parilla, K.M. Jones, S-H. Lee, “Metal Oxide Nanoparticles for Improved Electrochromic and Lithium-Ion Battery Technologies,” *Thin Solid Films* 516 (5), 794-797 (2008).
- N.A. Chernova, M. Roppolo, A.C. Dillon, and M.S. Whittingham, “Layered Vanadium and Molybdenum Oxides: Batteries and Electrochromics,” *Journal of Materials Chemistry* (Invited Review, in press).
- S-H. Lee, R. Deshpande, D Benhammou, P.A. Parilla, A.H. Mahan, and A.C. Dillon, *Thin Solid Films* In Press.
- L.A. Riley, C. Ban, L. Gedvillas, S-H. Lee, and A.C. Dillon, “Optimization of  $\text{MoO}_3$  Lithium Ion Battery Anodes,” *in preparation*.
- Yoon S. Jung, Andrew S. Cavanagh, Anne C. Dillon, Markus D. Groner, Steven M. George, and Se-Hee Lee, “Enhanced Stability of  $\text{LiCoO}_2$  Cathodes in Lithium-ion Batteries Using Surface Modification by Atomic Layer Deposition,” *Advanced Materials*, (submitted).