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**Effects of Cycling Conditions of Active Material
from Discharged Ni Positive Plates Studied by
Inelastic Neutron Scattering Spectroscopy**

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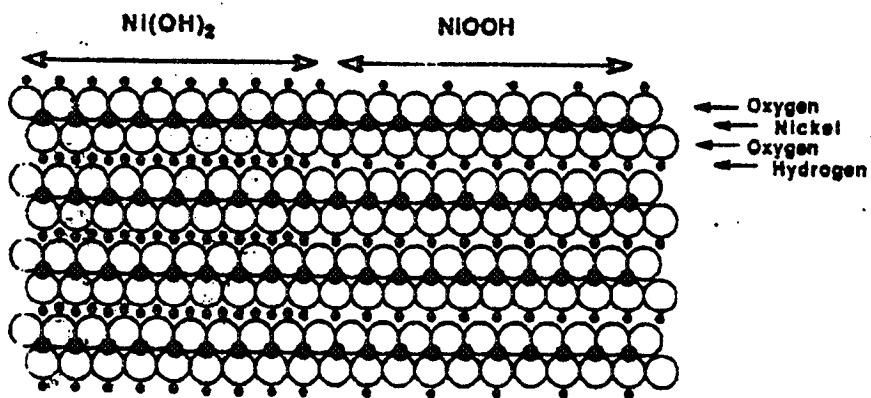
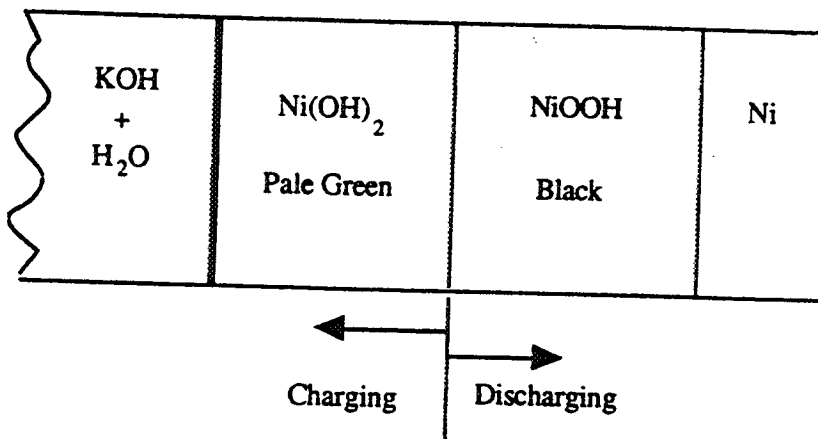
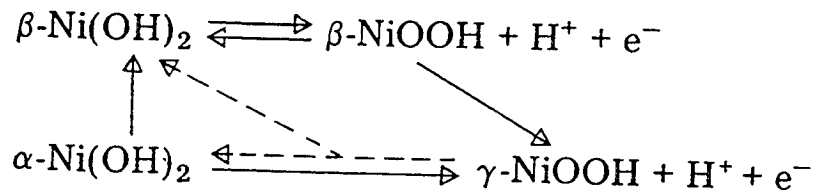
and

Ravi Varma, Lisa Diebolt and Margaret Reid

Objectives

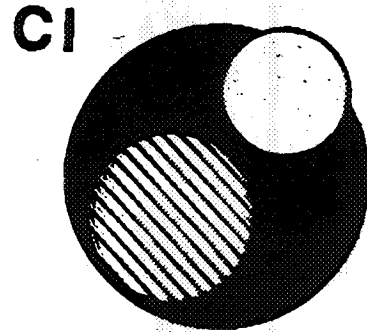
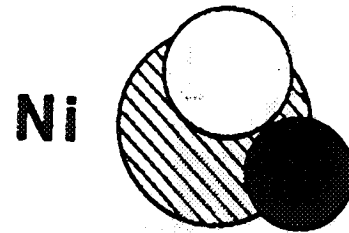
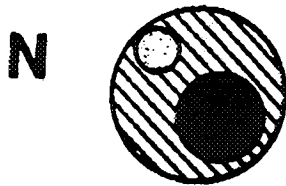
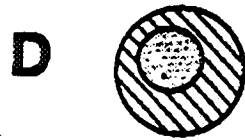
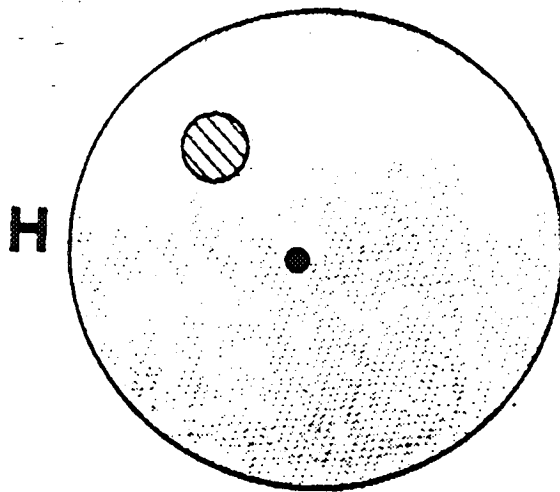
- Identify atomic-level signatures of electrochemical activity of the active material on the Ni positive plate of Ni-H₂ batteries.
- Relate findings to cycling conditions and histories
- Develop INS spectroscopy as a non-destructive testing technique for the evaluation of Ni-positive plates of Ni-H₂ batteries.

Charge/Discharge of (α, β) -Ni(OH)₂ / (γ, β) -NiOOH Couples



Fundamentals of Vibrational Spectroscopy by Inelastic Neutron Scattering

- neutrons are scattered by the atomic nuclei and not the electrons (as are photons)
 - scattering cross-sections a nuclear property
 - H scatters neutrons >10 times more strongly than other atoms
- absorption cross-sections for neutrons are very low:
 - probe the bulk of the sample
 - in-situ methods are easy(no windows required)
- all vibrational modes are observable
 - intensities are weighed by nuclear cross-sections:INS spectra are dominated by modes involving large displacements of H atoms.
 - intensities are readily quantifiable and are proportional to the number of scatterers.



Incoherent scattering cross section



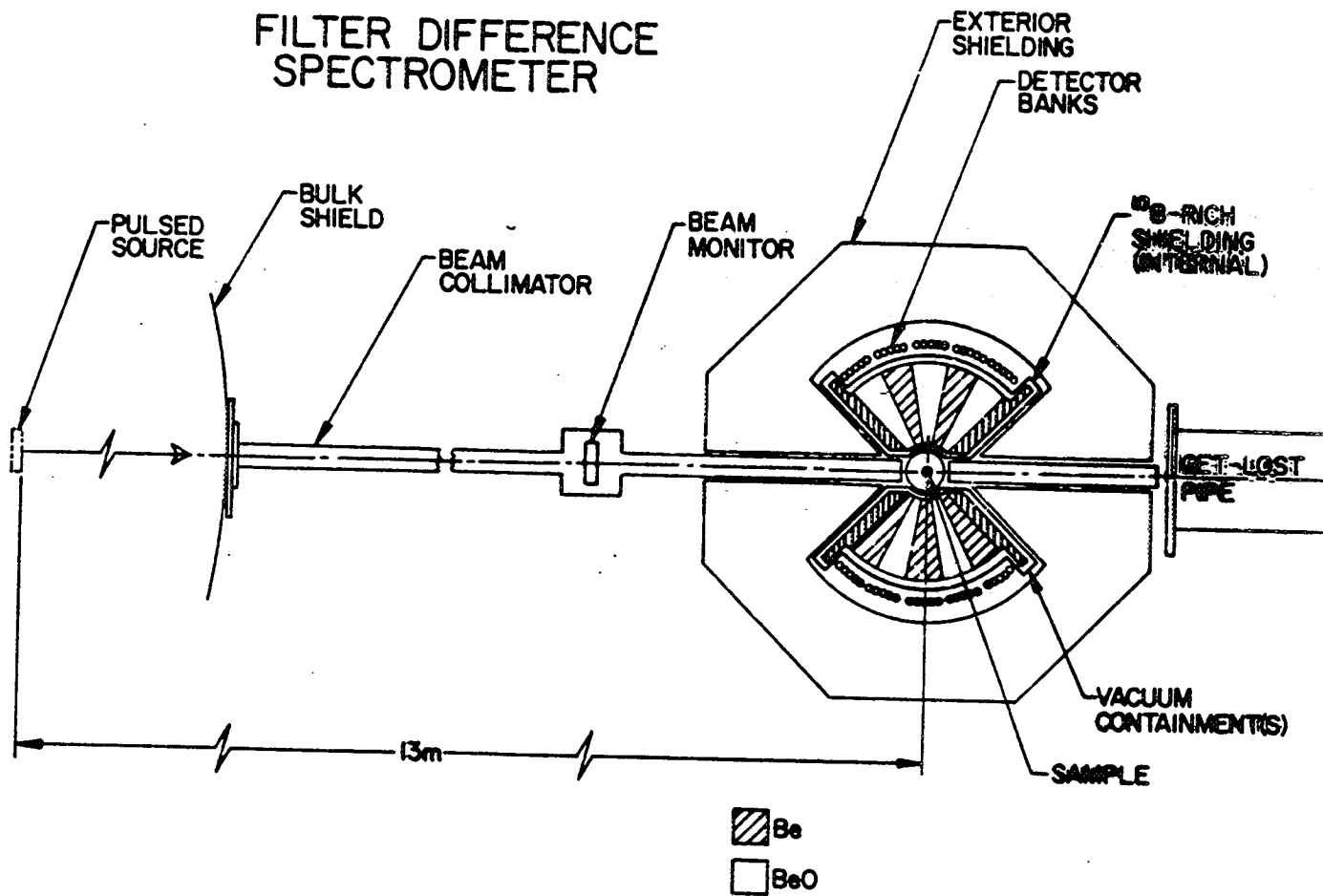
Coherent scattering cross section



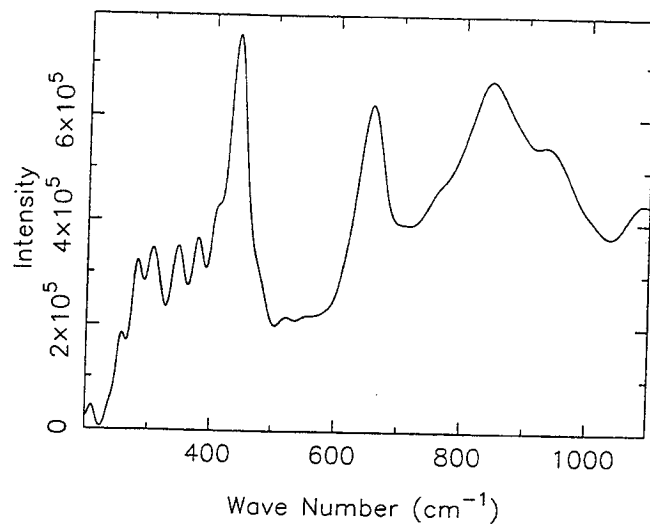
Absorption cross section

INS Vibrational Spectroscopy

- technique is well suited for application to battery material
 - bulk probe
 - sensitivity to protons (H)
- experiments are carried out at the Lujan Center of LANL
 - 5 - 10g samples from battery plates
 - FDS instrument; $\Delta E = 50 - 4000 \text{ cm}^{-1}$
 - 12-24 hrs. data collection time
 - T=15K

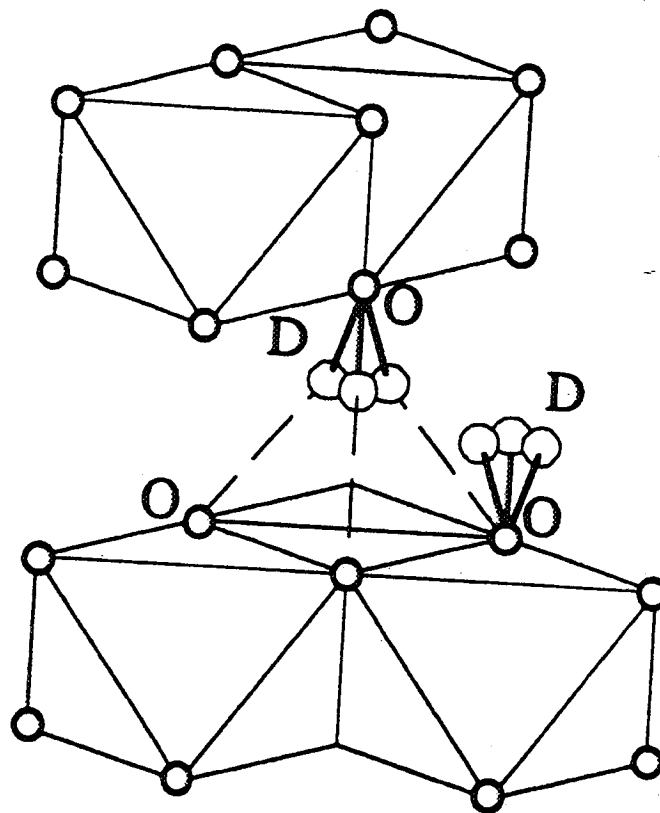


Assignment of β -Ni(OH)₂ vibrational bands



	INS (cm ⁻¹)	IR, Raman (cm ⁻¹)	Assignment
	315,290	318	$\nu(\text{NiO}); A_{1g}$
	358	350	$\delta(\text{ONiO}); E_u$
	390		$\nu(\text{NiO}); A_{2u}$
	451,412	449	$\delta(\text{ONiO}); E_g$
		452	accoust. + E_u
		530	
	673		$\gamma(\text{OH}) E_u$
	867		$\gamma(\text{OH}) E_g$
	929		

Hydrogen disorder in brucite structures



Raman Scattering Spectra of Ni electrode materials

B. C. Cornilsen and collaborators

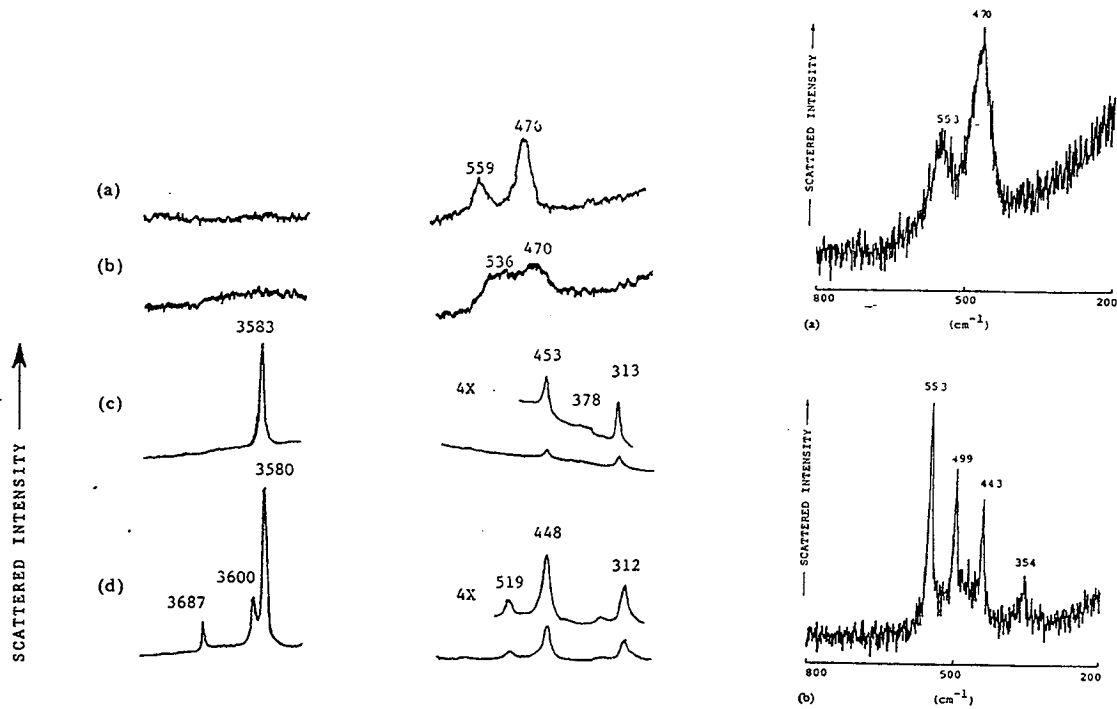
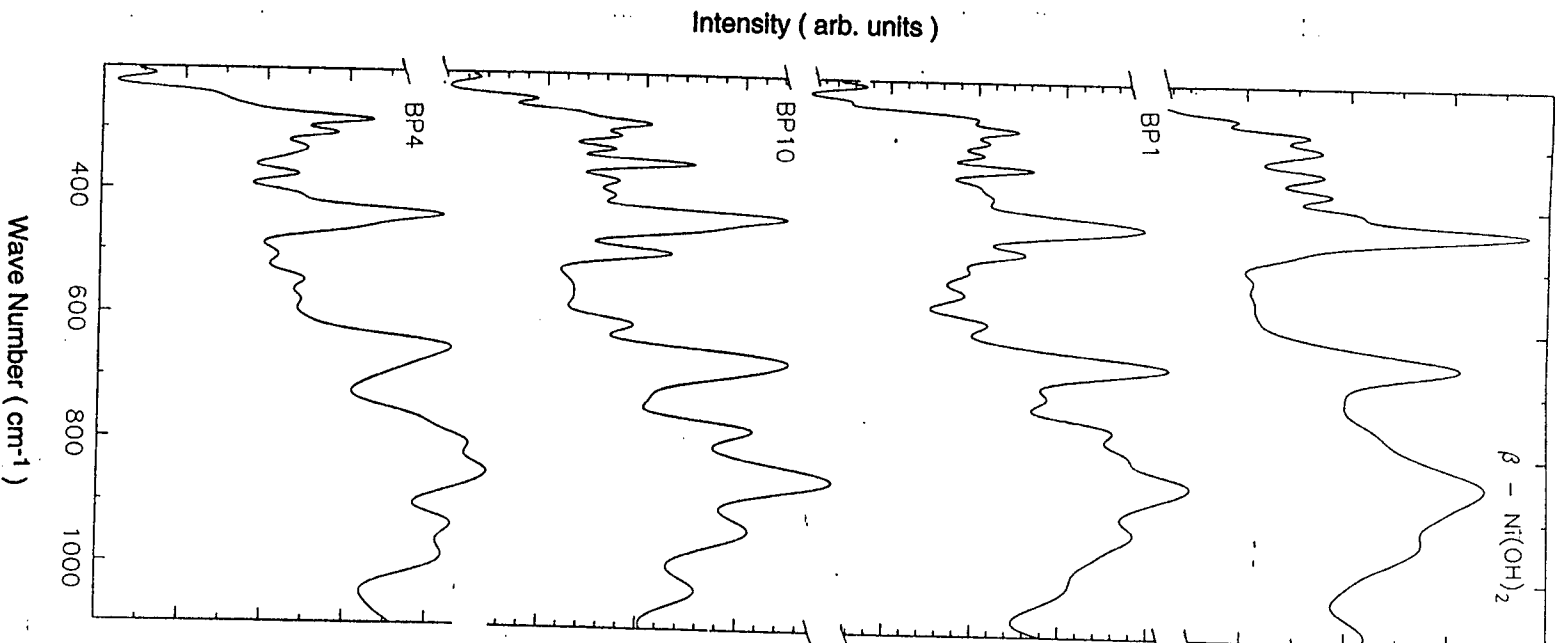


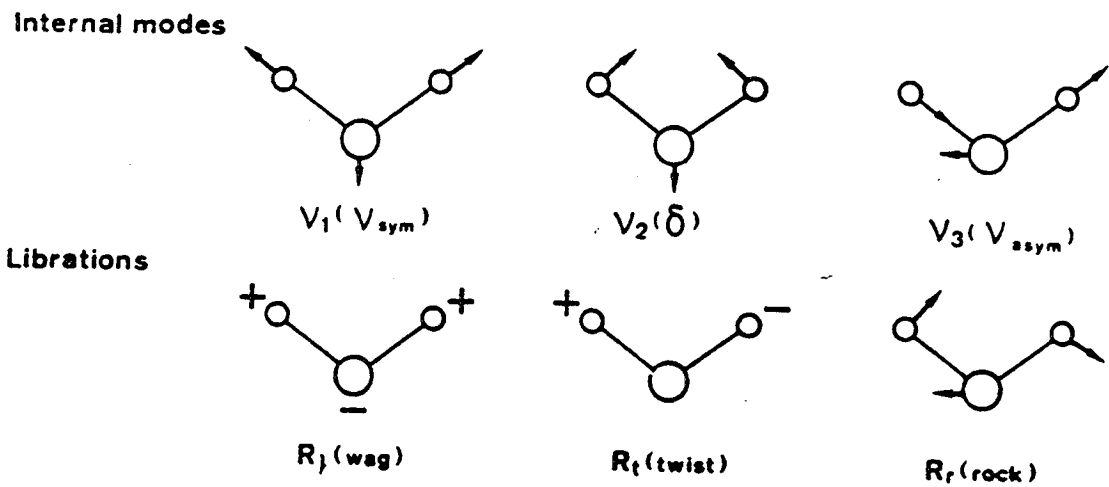
Fig. 1. Raman spectra of nickel electrode active mass and model compounds. (a) Charged γ active mass; (b) discharged α active mass; (c) recrystallized β -Ni(OH)₂; (d) first precipitate β -phase.

Raman spectra of: (a) discharged active mass (ID no. 16-09); (b) 'phase-X'

KOH Concentration	Number of Cycles
BP1 21 %	38,191
BP4 31 %	3,286
BP10 23.5%	28,495



Vibrational Modes of Hydration Water

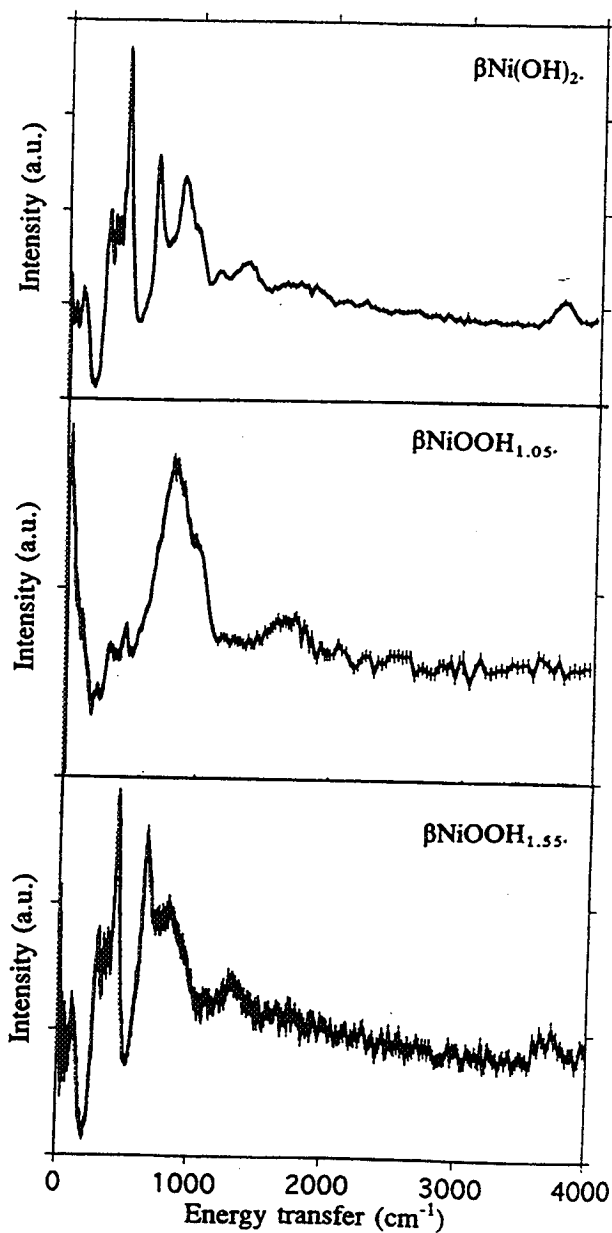


frequency ranges (cm^{-1})

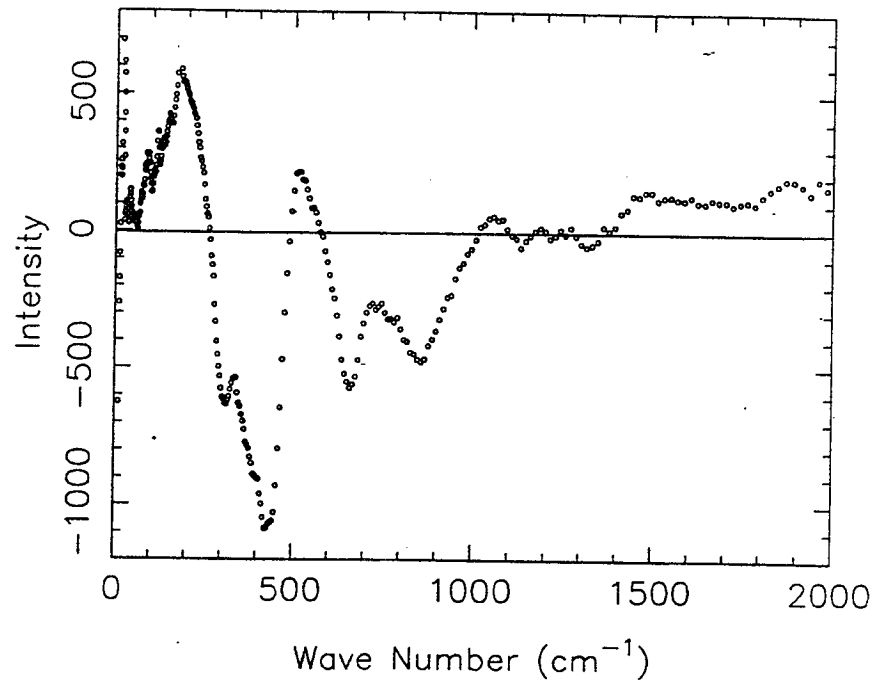
stretching modes (ν)	3600 - 3000
bending modes (δ)	1660 - 1590
librations (R)	1050 - 350
translatory modes (T)	350 - 100

INS Spectra of Reference Compounds

F. Fillaux et al., Physica B 213&214, 637 (1995)



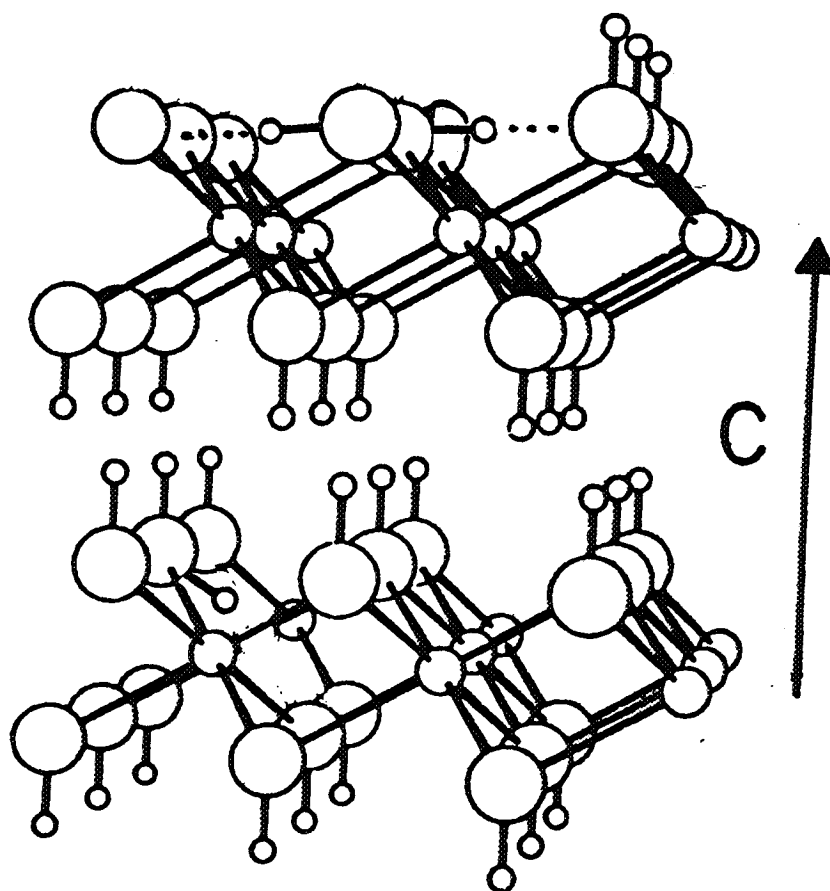
Ni(OH)₂ {BP1 - β}



INS results

- discharged materials are mainly β -Ni(OH)₂
- changes in the Ni-O stretching and bending regions:
 - a decreases from 3.13 Å (β -Ni(OH)₂) to 2.89Å (β -NiOOH)
 - distortion of NiO₆ octahedron
 - frequency shifts and band splittings result
- water librations above $\sim 500 \text{ cm}^{-1}$
 - vacancies may allow formation of Ni(H₂O)
- protons in O-H...O hydrogen bonds: β -NiOOH

Structural Models for Hydrogen in NiOOH and bound H₂O



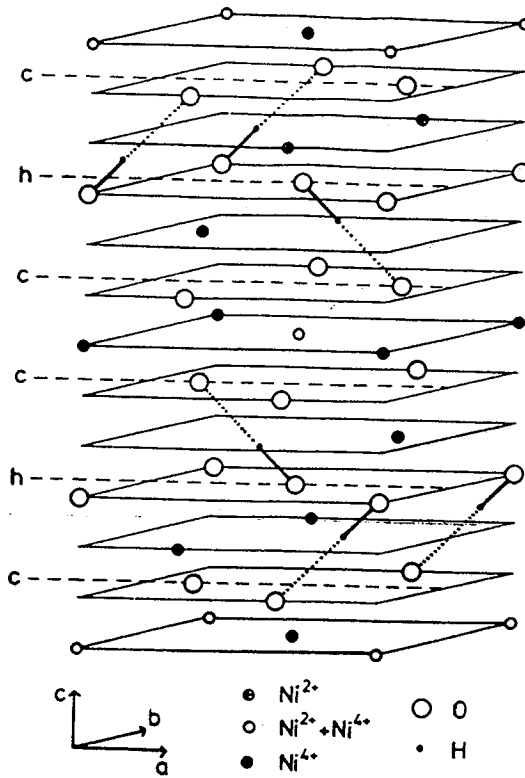


FIGURE 2
Schematic representation of the structure of Ni_2O_3H

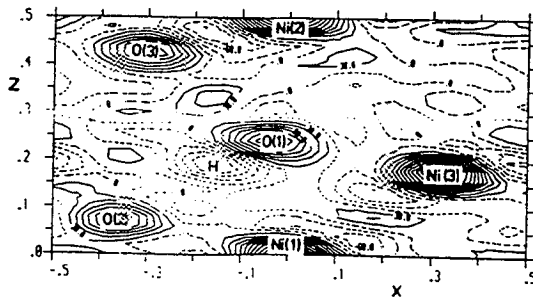


FIGURE 3
Fourier section, $y = 0$, based on observed intensities and calculated phases

Conclusions

- (1) Irreversible formation of NiOOH; scales with number of cycles
- (2) additional protons are bound in the lattice to form Ni-(H₂O) complexes; increases with KOH concentration in the cell.
- (3) These processes occur only in the outermost layers of the plate material but lead to the failure of the battery cells.

Acknowledgments

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