

# Enhanced Hydrogen Storage in Mg thin Flakes with dispersed Ni Nanoparticles prepared by High Energy Ball Milling.

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# Introduction | solid state storage

## Solutions for H<sub>2</sub> storage

Gas

Liquid (LH<sub>2</sub> - LHCs)

Solid

Lower risk in transportation and higher gravimetric capacities.

## Magnesium hydride (MgH<sub>2</sub>)

Lightweight, reversible and abundant on Earth's crust

Theoretical storage capacity (7.67 wt.%)

Sluggish kinetics and thermodynamic stability

$E_a$   
146 kJ/mol

Sorption /  
Desorption  
>350 °C

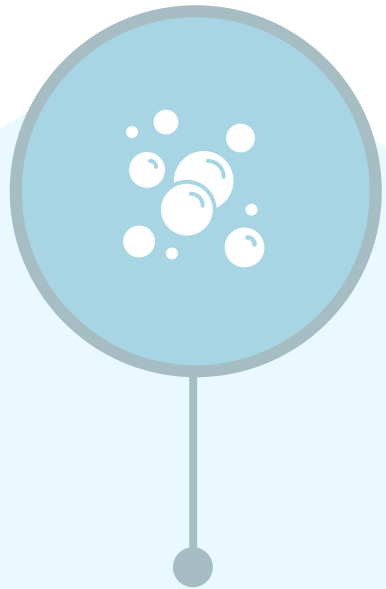


## Introduction | Mg modifications

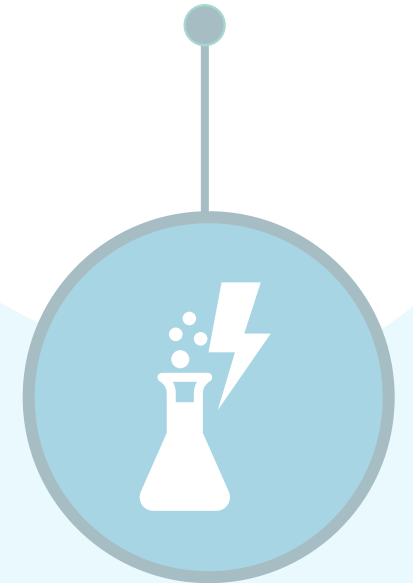
They Improve kinetics or thermodynamics  
But decrease storage capacity

*An approach that involves various modifications is required*

Metal-oxide  
catalysts



Nanoparticles or  
nanostructures



Alloying  
elements and  
additives

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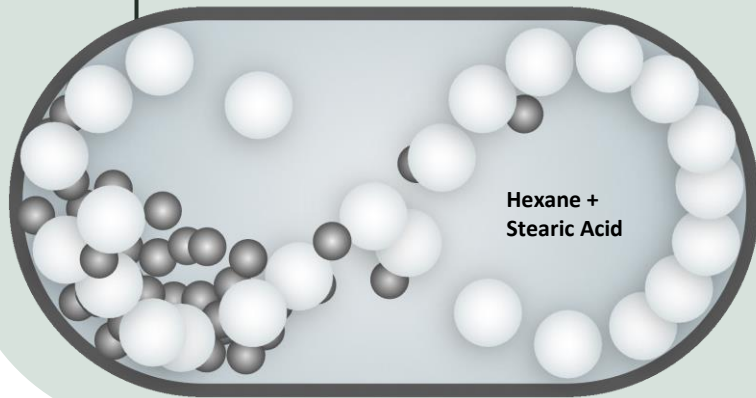
Conclusions



## Two step method of Surfactant Assisted – High Energy Ball Milling

1

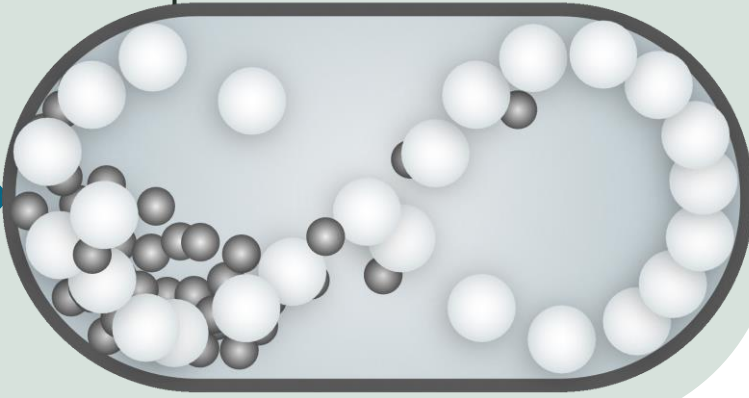
- BPR 40:1
- 1400 rpm
- ZrO<sub>2</sub> balls, 1 mm



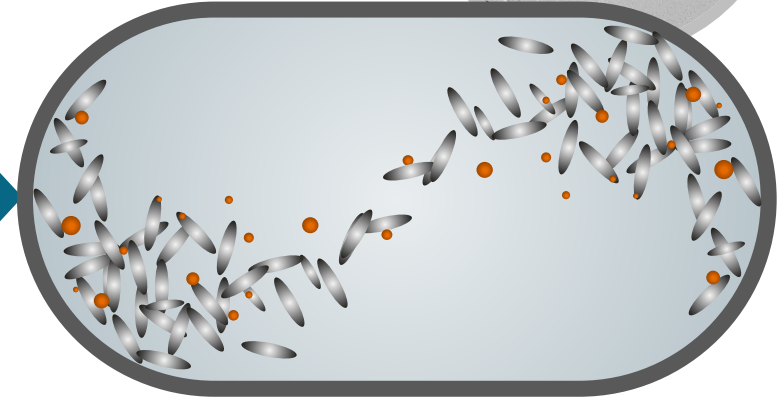
*Comminution*

2

- BPR 60:1
- 1400 rpm
- ZrO<sub>2</sub> balls, 3 mm



*Deformation*



*Blending*

5 wt.%Ni

15 wt.%Ni



## Materials and Methods | Characterization

Absorption /  
Desorption  
tests  
(350°C)

Differential  
Scanning  
Calorimetry

X-Ray  
Diffraction

SEM-EDX



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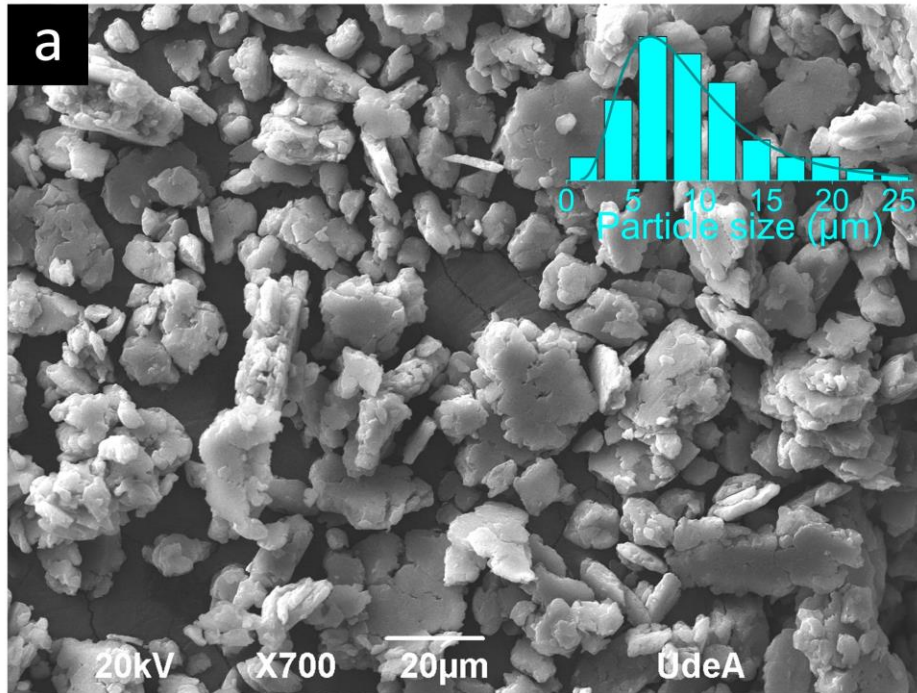


Conclusions

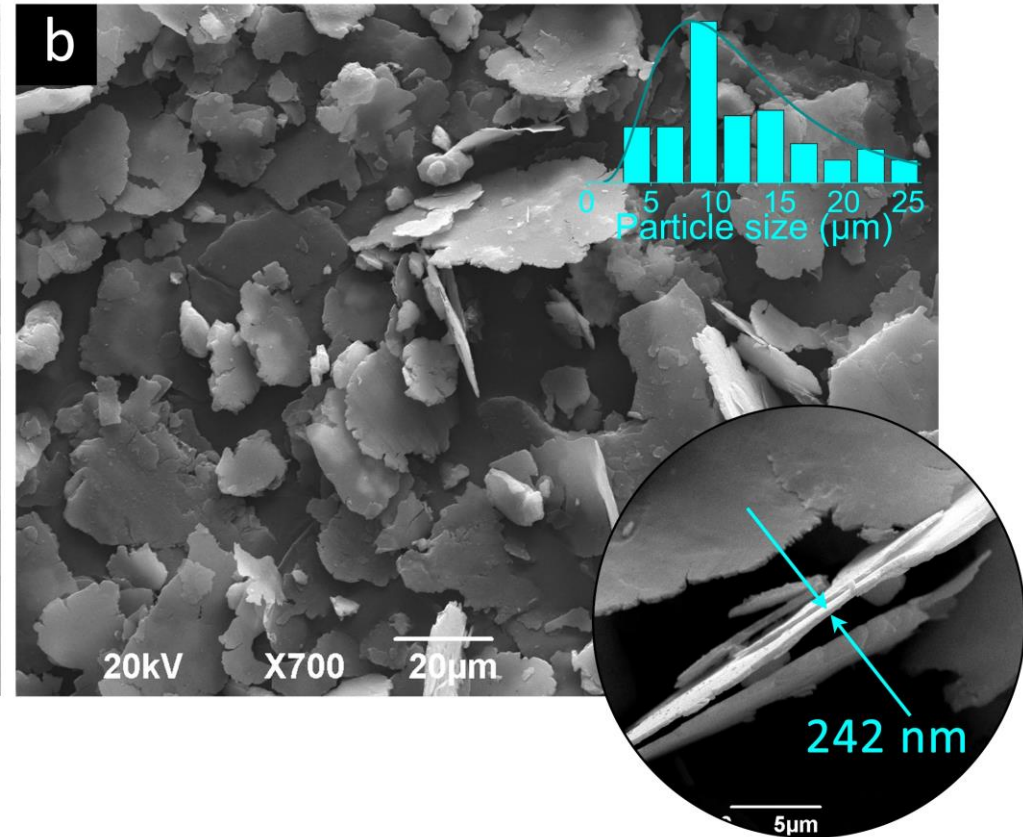


# Results | Mg thin flakes

1st step



2nd step

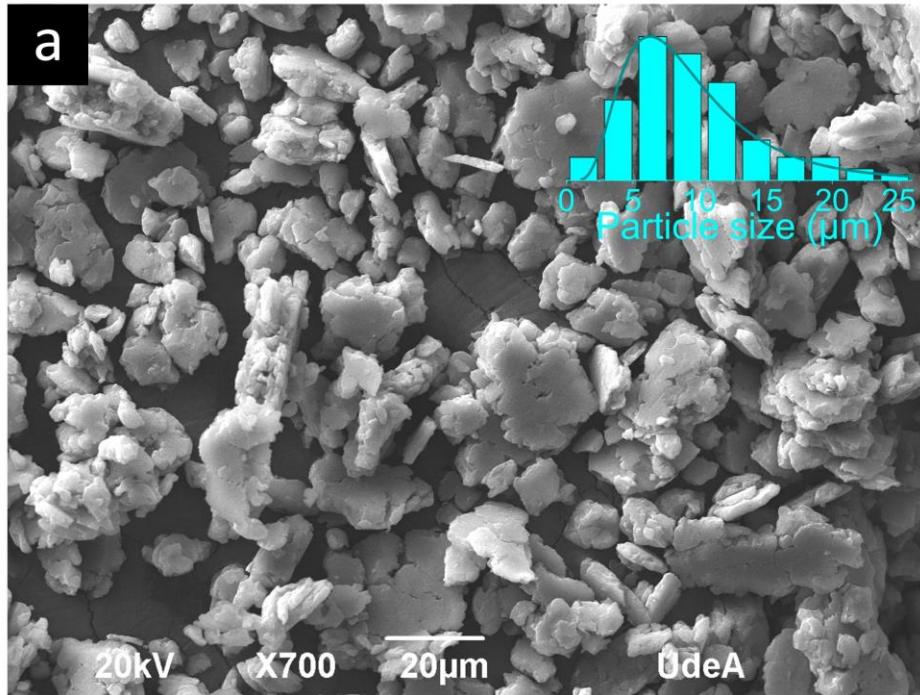


Starting crystallite size  
197.9 nm

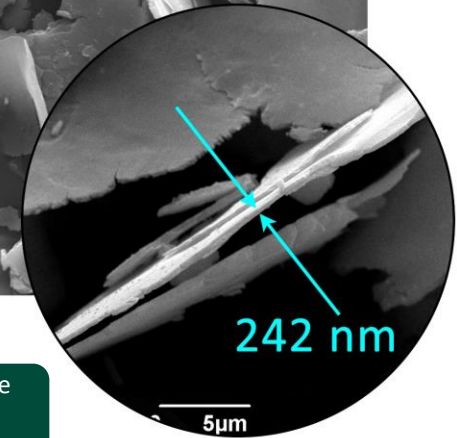
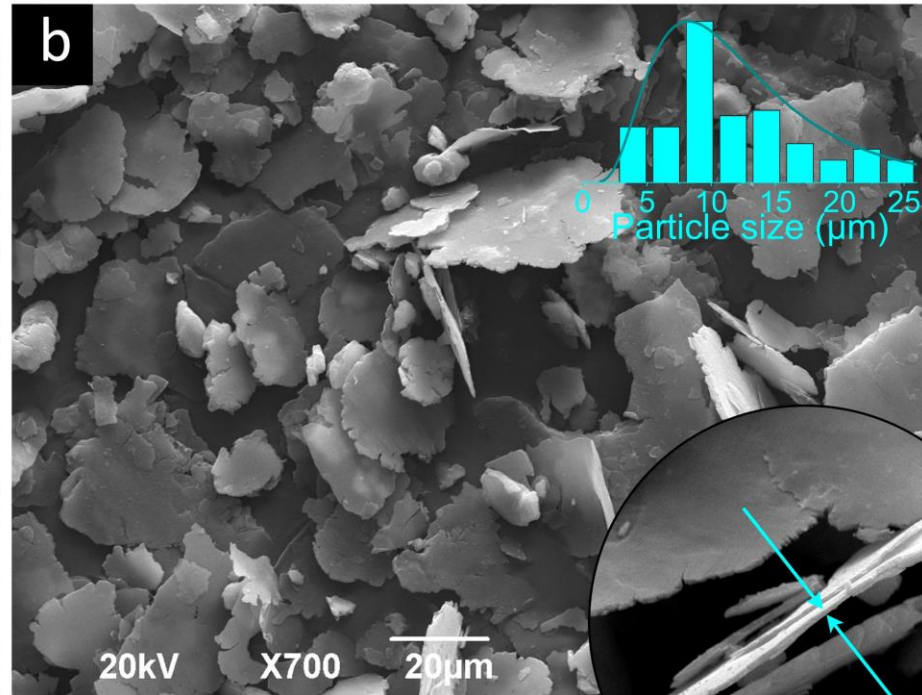


# Results | Mg thin flakes

1st step



2nd step

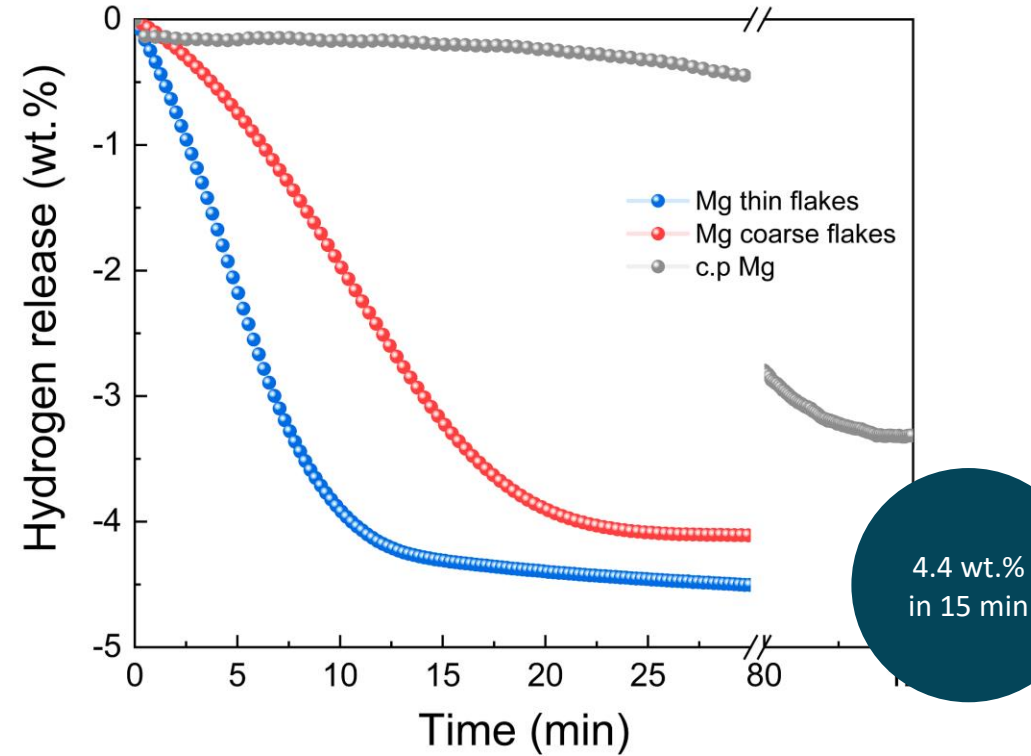
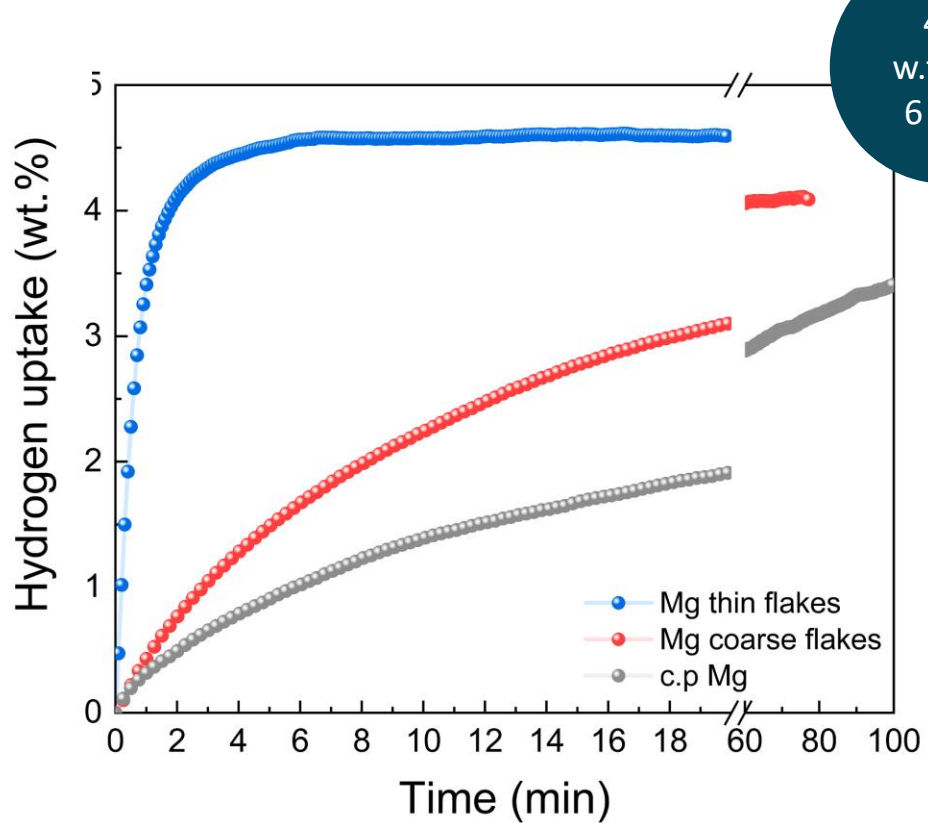


Refined crystallite size

17.8 nm



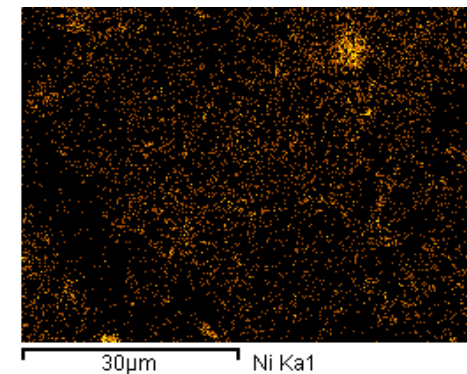
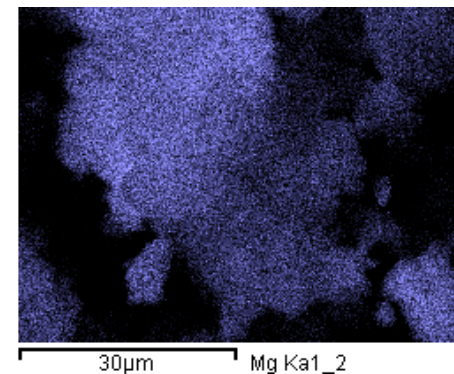
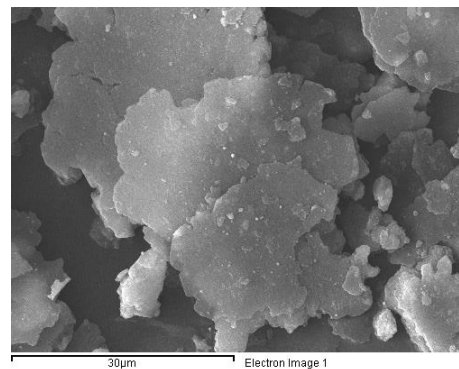
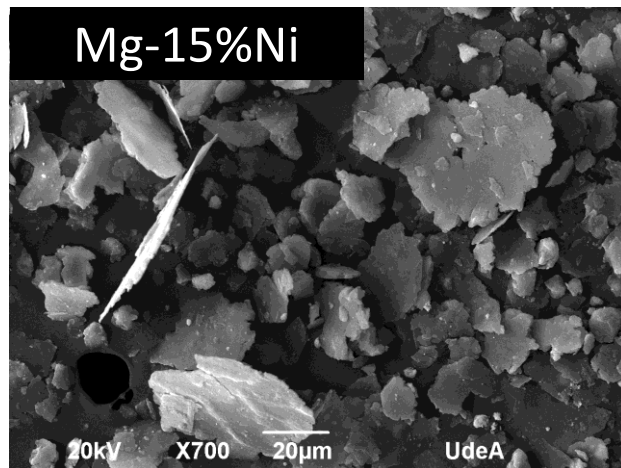
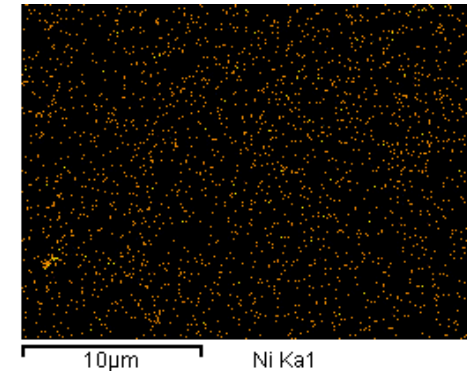
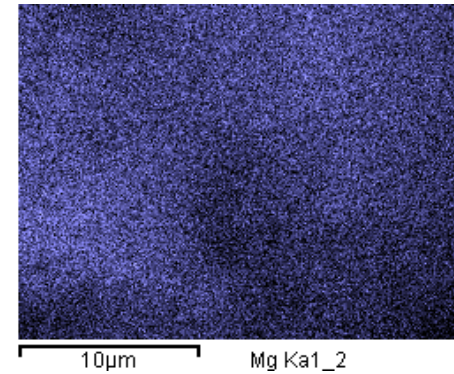
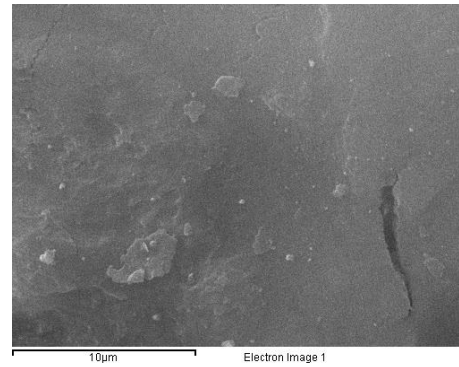
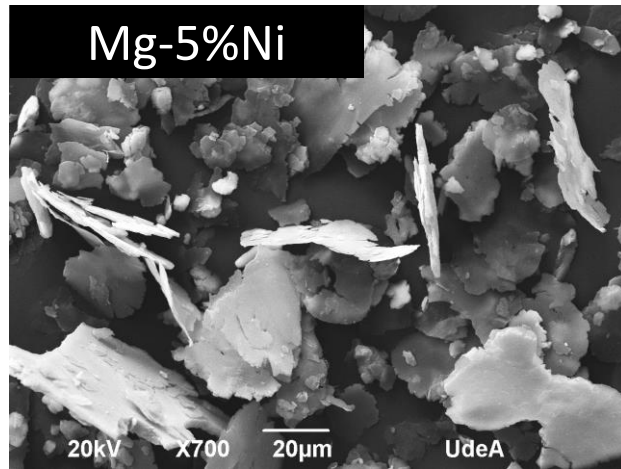
## Results | Mg thin flakes



Improved kinetics (94% less time) and capacity (increase in 25%)



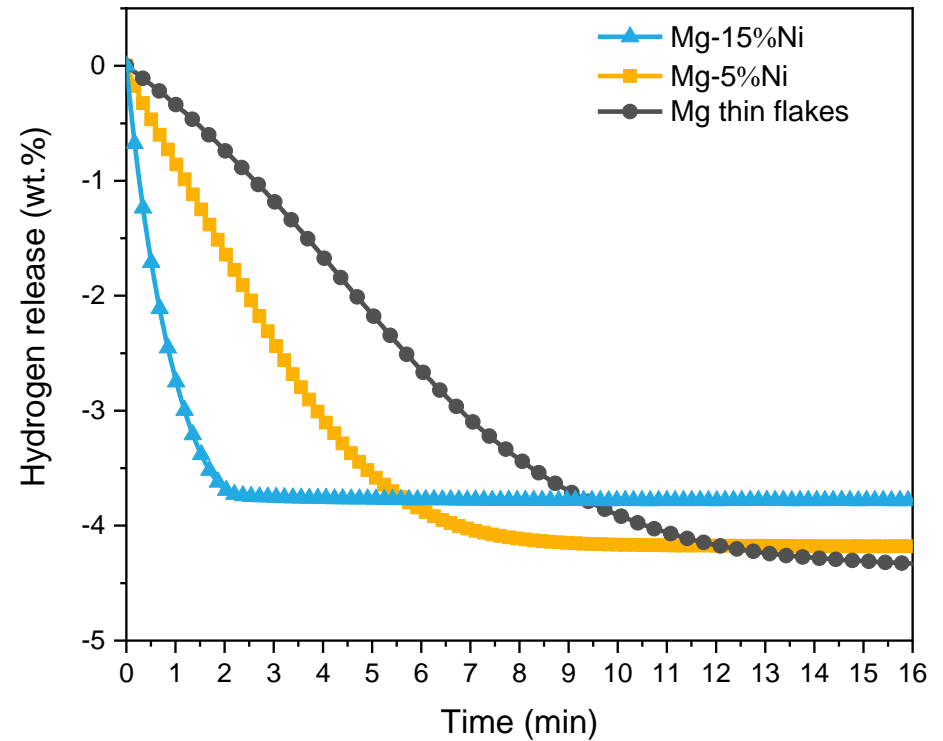
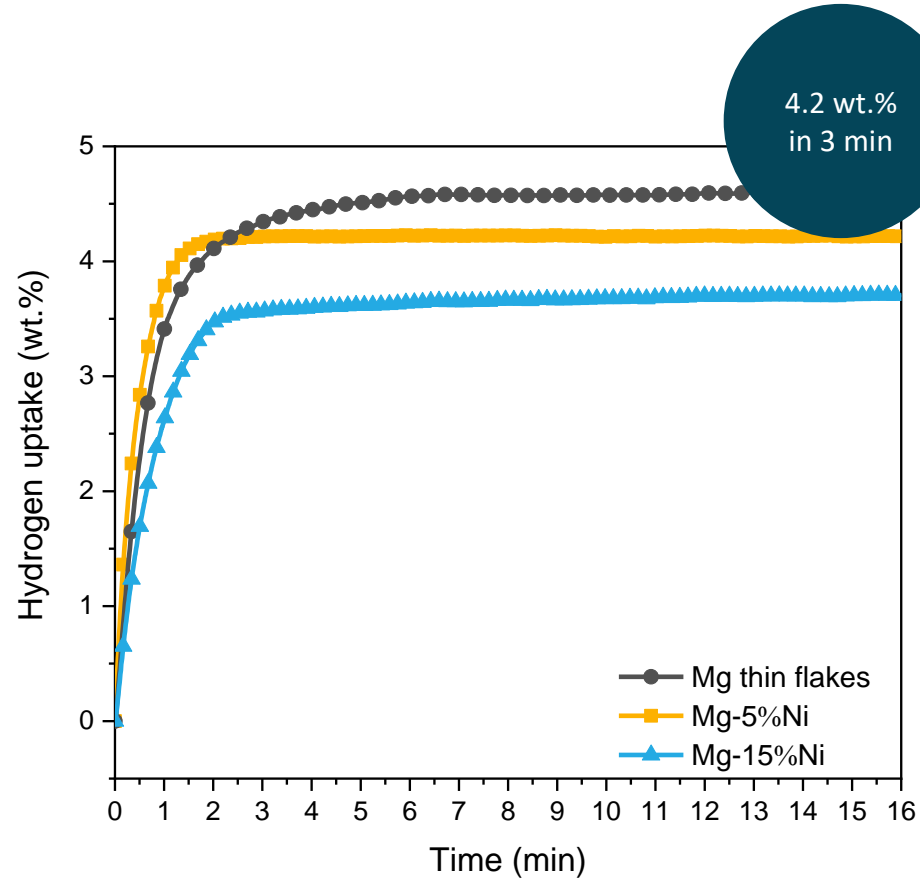
# Results | Mg with Ni decoration



Homogeneous distribution through the dry blending method



# Results | Kinetics test at 350°C, 20 bar

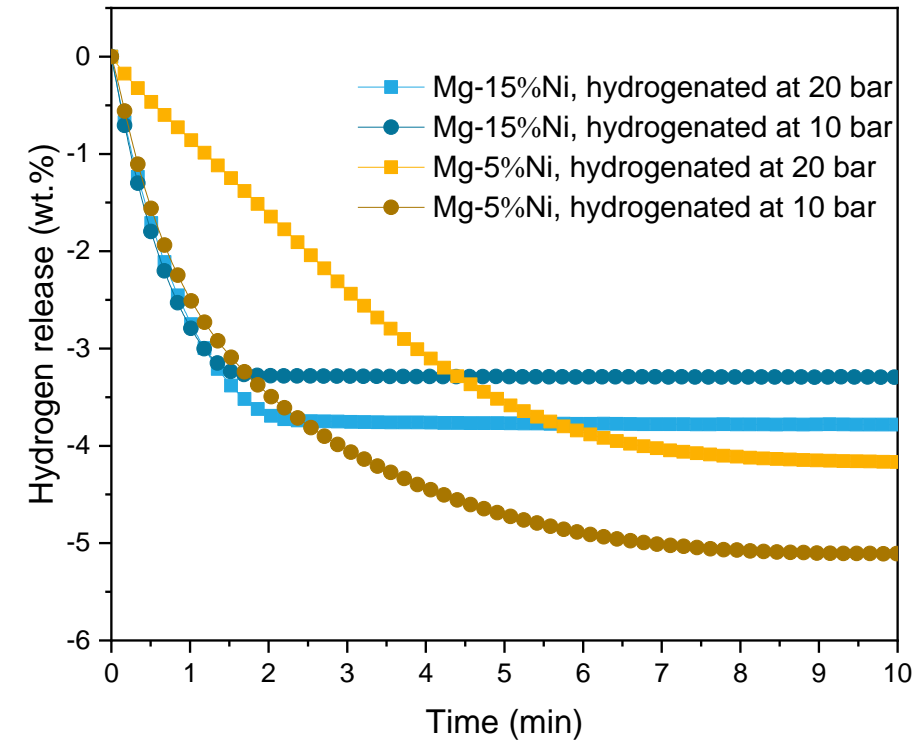
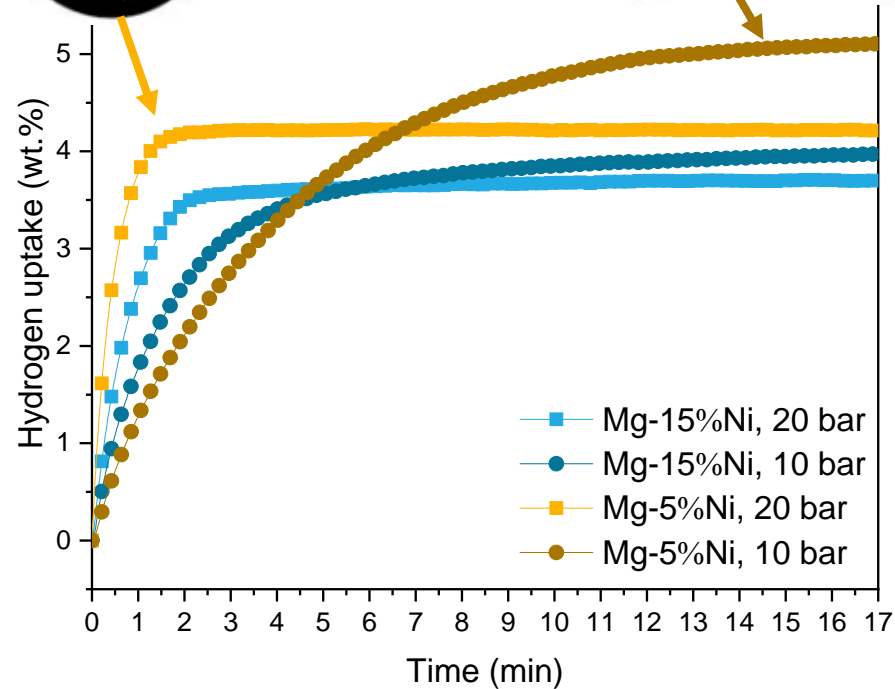


Improved kinetics (50% of the time) but decreased capacity (9%)



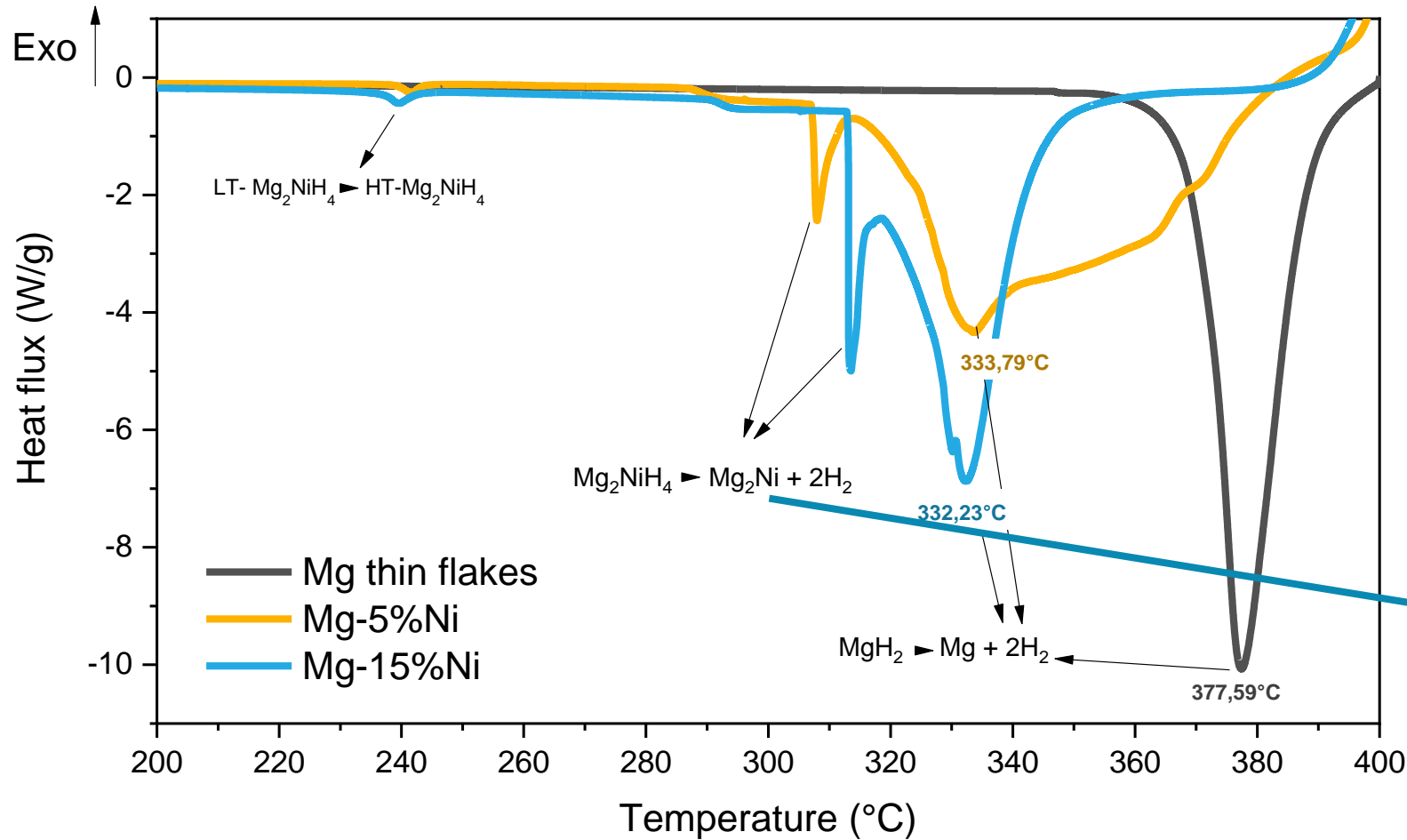
## Results | Kinetics at 350°C, 10 bar

Lower pressures resulted in higher capacities but slower kinetics due to a change in sorption mechanism.





# Results | Mg with Ni decoration



Decrease in operation temperature

$\text{Mg}_2\text{Ni}$  promotes desorption reaction

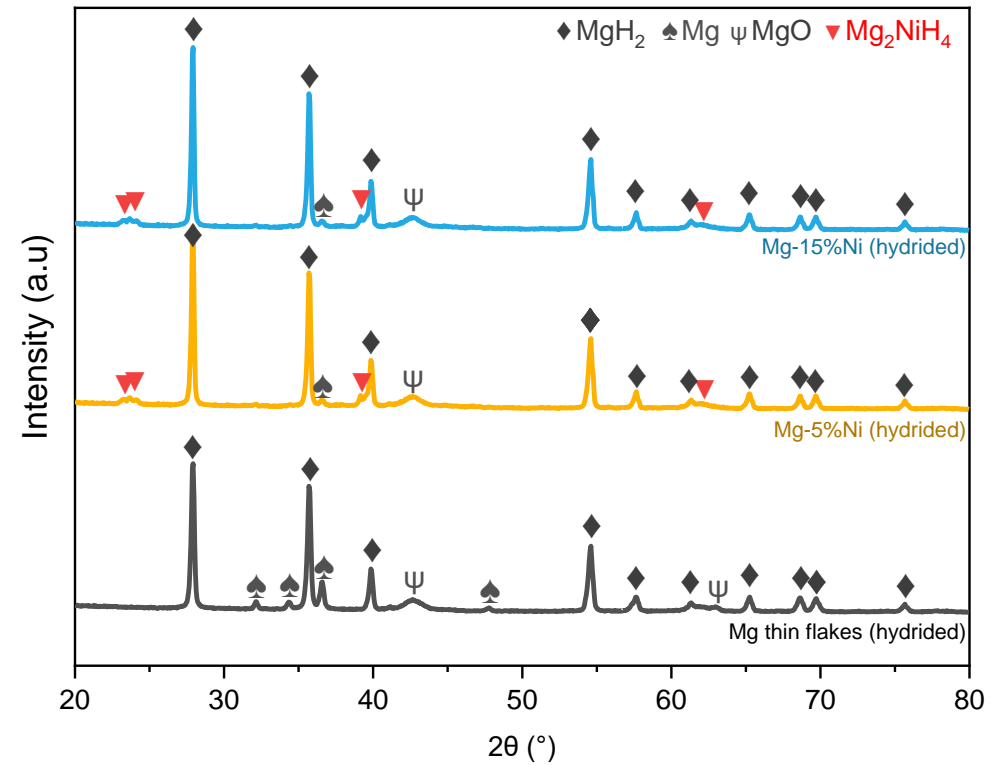
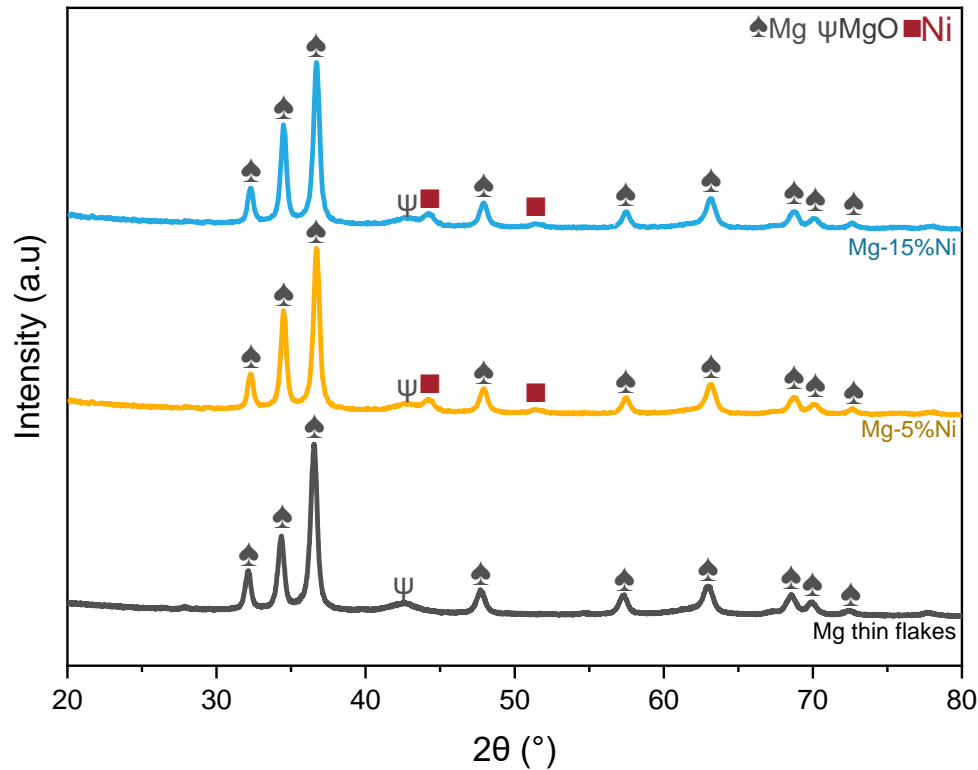
This was later confirmed by XRD analysis





# Results | Mg with Ni decoration

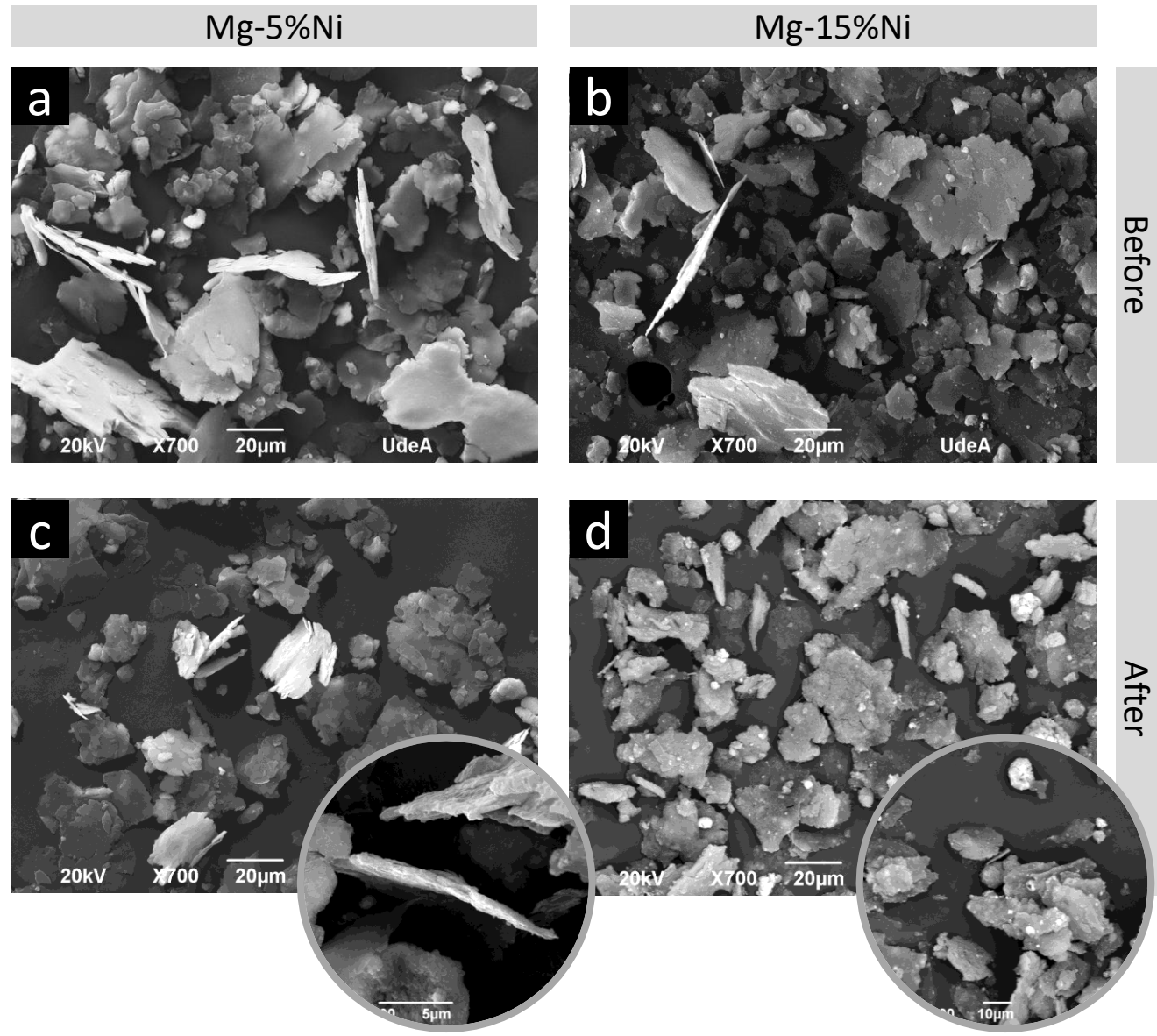
Oxygen content (EDX) below 5 wt.%



Hydrogen sorption at 350°C promotes Mg – Ni reaction to form the complex hydride



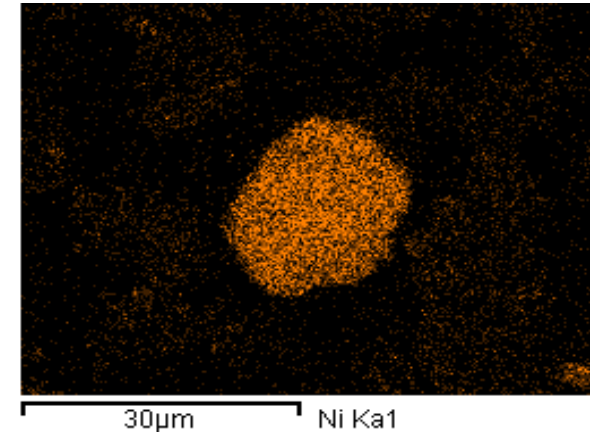
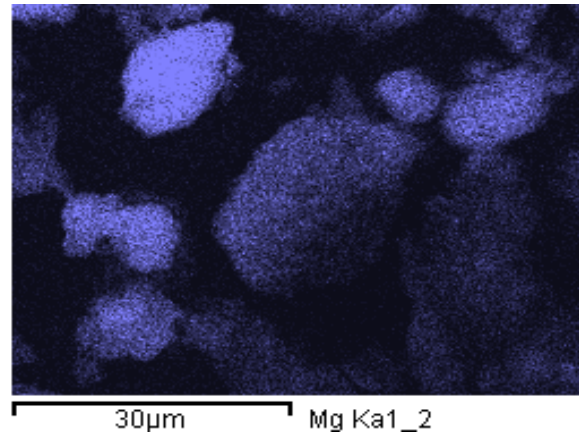
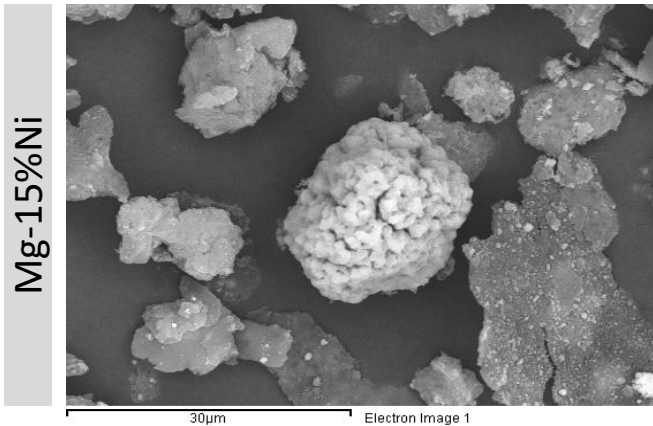
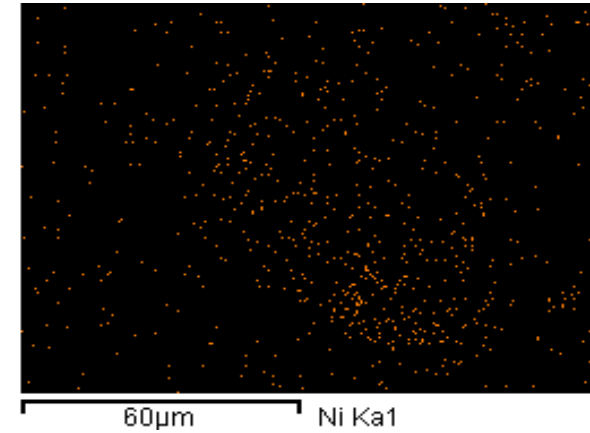
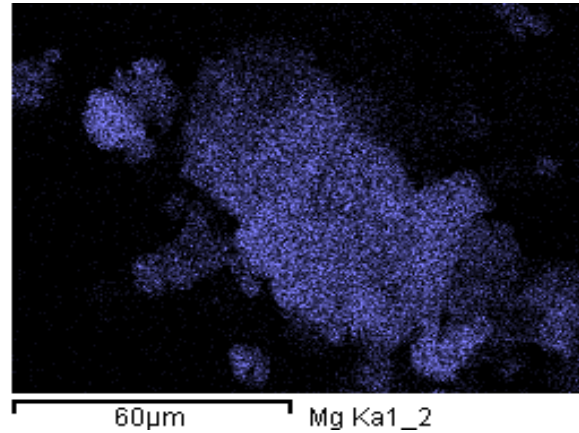
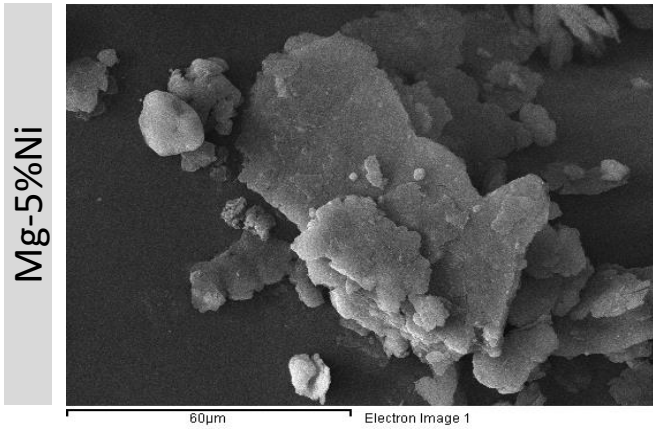
# Results | Mg with Ni decoration



Preserved morphology after 5 cycles



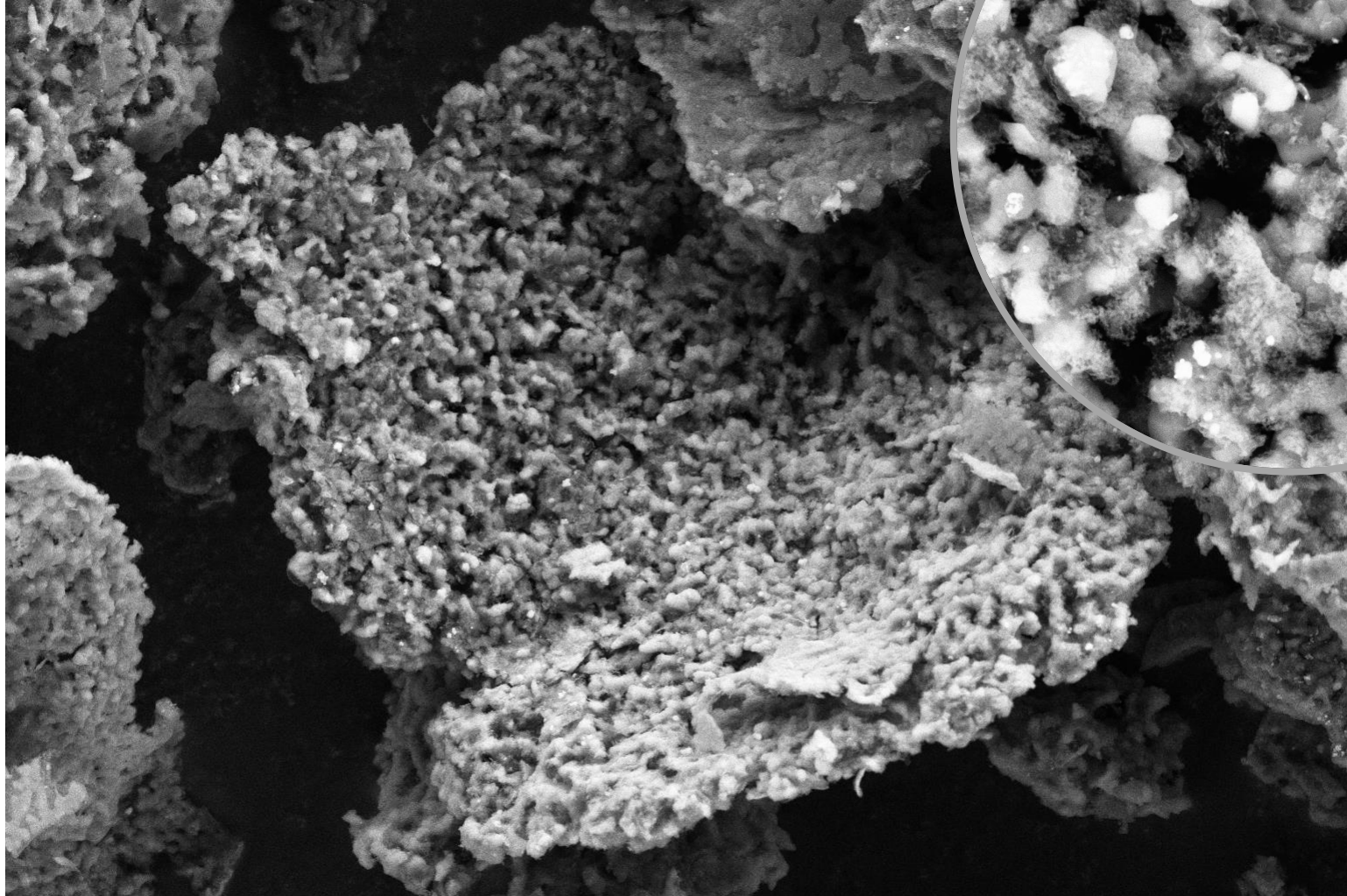
# Results | Mg with Ni decoration



Higher Ni content agglomerated after sorption/desorption tests



# Results | Mg with Ni decoration



**416 ± 127 nm**  
Cycling could lead to finer particles

	use case	det	mode	mag	HV	curr	WD	HFW	vac mode	5 μm
OptiPlan	ETD	SE	5 000 x	5.00 kV	50 pA	2.9 mm	41.4 μm	High vacuum	Scios 2	

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# Conclusions

1

A two step ball milling method led to Mg thin flakes (thickness <300 nm) and improved hydrogen storage capacity

2

5 wt.%Nickel decoration improved sorption/desorption process in 50% of the time with a decrease of 9% in capacity

3

The formation of  $Mg_2NiH_4$  after activation process led to improved dehydrogenation kinetics

# Thanks *for* your attention

