

Waste biorefinery technologies for accelerating sustainable energy processes

# Enhancing Oxygen Reduction via N-Doped Graphene Derived from PET Bottle Waste as an Efficient Carbon Support for PdNi Nanoparticles

#### A. Balčiūnaitė<sup>1</sup>, B. Šljukić<sup>2</sup>, N.A. Elessawy<sup>3</sup>, M.H. Gouda<sup>3</sup>, D.M.F. Santos<sup>2</sup>

<sup>1</sup>Department of Catalysis, Center for Physical Sciences and Technology, 10257 Vilnius, Lithuania <sup>2</sup>Center of Physics and Engineering of Advanced Materials, Instituto Superior Técnico, Universidade de Lisboa, 1049-001 Lisbon, Portugal <sup>3</sup>City of Scientific Research & Technological Applications (SRTA-City), 21934 Alexandria, Egypt



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# Introduction

Why materials from PET bottles waste?

What is the problem with PET plastic waste?

- plastics like PET can break down into tiny pieces called microplastics
- are pervasive in our oceans, bays, lakes, and even drinking water

Using waste PET as an electrocatalyst support for electrochemical energy conversion devices can lower its environmental impact, including greenhouse gas emissions, energy demand, water consumption, acid rain, and eutrophication potential.







# Experimental



## Materials from PET

The composition of the prepared catalysts as determined by ICP-OES.

Catalant	Loading		
Catalyst	Pd, µg cm <sup>-2</sup>	Ni, mg cm <sup>-2</sup>	
Ni@NG	-	0.70	
PdNi_5@NG	0.70	0.43	
PdNi_10@NG	0.94	0.41	
PdNi_15@NG	2.05	0.44	









TEM image of PdNi\_15@NG.

Catalysts	a <sub>S,BET</sub> (m <sup>2</sup> g <sup>-1</sup> )	Total pore volume (cm <sup>3</sup> g <sup>-1</sup> )	Average pore diameter (nm)
NG	57.3	0.053	3.72
Ni@NG	97.3	0.062	2.56
PdNi_5@NG	70.6	0.076	4.31
PdNi_10@NG	39.5	0.089	8.99
PdNi_15@NG	6.05	0.033	21.5



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#### Characterization



The XPS elemental survey spectra of PdNi\_15@NG surface.

# **Experimental**





1. Catalyst



#### 2. Catalyst ink

5 mg catalyst
100 μL of 2 wt.% solution of

PVDF in NMP

Sonicated for 30 min



3. Drying



PVDF - polyvinylidene fluoride; NMP - N-methyl-2- pyrrolydone



#### Oxygen reduction



Current densities are ~2 times higher at Ni@NG than at the NG support

CVs of NG (a) and Ni@NG (b) recorded in 0.1 M KOH at 50 mV s<sup>-1</sup>.







CVs of PdNi\_5@NG (a), PdNi\_10@NG (b), and PdNi\_15@NG (c) in 0.1 M KOH at 50 mV s<sup>-1</sup>.

# Current densities are ~2 and over 3 times higher at PdNi\_15@NG than at PdNi\_10@NG and PdNi\_5@NG catalysts, respectively.





Oxygen reduction

Koutecky-Levich (K-L) equation:  $\frac{1}{I} = \frac{1}{I_{k}} + \frac{1}{I_{d}} = -\frac{1}{nFAkc_{O_{2}}^{b}} - \frac{1}{0.62 nFAD_{O_{2}}^{2/3} v^{-1/6} c_{O_{2}}^{b} \omega^{1/2}}$ 

Current densities increase ~ 3.9 and 6.9 times with the increase of rotation rate from 0 to 2400 rpm on NG and Ni@NG catalysts, respectively.

LSVs of NG (a) and Ni@NG (c) recorded in 0.1 M KOH at 10 mV s<sup>-1</sup> at different rotation rates. (b,d) K-L plots of the NG (b) and Ni@NG (d) collected at different potentials.



Oxygen reduction

Current densities increase ~2.7 and 4.0 times with the increase of rotation rate from 0 to 2400 rpm on PdNi\_5@NG and PdNi\_10@NG catalysts, respectively.

LSVs of PdNi\_5@NG (a) and PdNi\_10@NG (c) recorded in 0.1 M KOH at 10 mV s<sup>-1</sup> at different rpms. (b,d) K-L plots of PdNi\_5@NG (b) and PdNi\_10@NG (d) collected at different potentials.

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#### Oxygen reduction

(a) LSVs of PdNi\_15@NG recorded in 0.1 M KOH at 10 mV s<sup>-1</sup> at different rpms. (b) K-L plots of PdNi\_15@NG collected at different potentials. (c) Comparison of cathodic currents of different catalyst compositions at 1600 rpm. (d) K-L plots of the different catalysts collected at -0.5 V.

#### The current densities are ~ 1.1-2.5 times higher on PdNi\_15@NG.

 Table 3. ORR performance of different catalysts.

	E <sub>onset</sub> (V)	E <sub>1/2</sub> (V)	$\mathbf{j}_{\mathrm{d}}$ (mA cm <sup>-2</sup> )
GN	-0.274	-0.371	-0.63
Ni@GN	-0.293	-0.410	-1.16
PdNi_5@NG	-0.258	-0.376	-0.71
PdNi_10@NG	-0.249	-0.396	-1.41
PdNi_15@NG	-0.161	-0.366	-1.53

- The studies showed that the prepared PdNi@NG catalysts have good electrochemical stability in alkaline solution.
- The PdNi\_15@NG catalyst exhibited the best electrocatalytic activity for the oxygen reduction reaction.
- The prepared PdNi@NG catalysts are promising electrode materials for alkaline fuel cells.
- N-doped graphene from PET bottle wastes was demonstrated to be an effective electrocatalyst support.

