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1 Precious Metal-Free Nickel Nitride Catalyst for the Oxygen 2 Reduction Reaction

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24 Abstract:

25 With promising activity and stability for the oxygen reduction reaction (ORR), transition
26 metal (TM) nitrides are an interesting class of non-platinum group catalysts for polymer
27 electrolyte membrane fuel cells (PEMFCs). Here we report an active thin film nickel nitride
28 catalyst synthesized through a reactive sputtering method. In RDE testing in 0.1M HClO₄
29 electrolyte, the crystalline nickel nitride film achieved high ORR activity and selectivity to
30 4 electron ORR. It also exhibited good stability during 10 h and 40 h chronoamperometry
31 (CA) measurements in acid and alkaline, respectively. A combined experiment-theory
32 approach, with detailed *ex-situ* characterization with TEM and NEXAFS to reveal a mixed
33 Ni₄N/Ni₃N structure with an amorphous surface oxide and DFT calculations to provide
34 insight into the surface structure during catalysis, is highlighted. Design strategies for
35 activity and stability improvement through alloying and nanostructuring are discussed.

36

37 Keywords:

38 Electrocatalysis, oxygen reduction reaction, non-precious metal catalysts, transition metal
39 nitrides, reactive sputter deposition, density functional theory

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Introduction

Proton-exchange membrane fuel cells (PEMFCs) are a promising technology towards enabling efficient and clean electricity production for transportation and industrial applications. In a PEMFC, hydrogen fuel is oxidized at the anode and oxygen is reduced at the cathode, generating electricity and releasing water as a benign by-product. Current commercial PEMFCs utilize expensive platinum-based catalysts for both the anode and cathode.¹ In particular, prohibitively large Pt loadings are required at the cathode to overcome the sluggish kinetics of the oxygen reduction reaction (ORR).^{2,3} While much work has focused on improving the mass activity of Pt ORR catalysts,⁴⁻⁷ limitations, including scarcity, instability, and susceptibility to carbon monoxide poisoning, remain.⁸⁻¹⁰ Therefore, the discovery of earth abundant, non-precious metal ORR catalysts is of critical interest for the large scale implementation of fuel cell technology.

Many non-platinum group (non-PGM) materials have been explored for ORR, including transition metal oxide^{11,12} and nitrogen-doped carbon¹³ catalysts that have shown promising activity in alkaline electrolyte, and metal-nitrogen-carbon¹⁴⁻¹⁶ catalysts that have achieved Pt-like activity in acid. Another promising class of earth-abundant materials are the transition metal nitrides, which are electronically conductive and have shown enhanced catalytic activity relative to their parent metal due to beneficial changes in the d-band structure.¹⁷ Moreover, nitrides have achieved activity approaching that of the noble metals for a variety of reactions.¹⁸ ORR activity has been demonstrated for several nitrides, including Co, Fe, and Ti.¹⁹⁻²⁴ Nitrides of Mo and W have also shown promising stability in acidic electrolyte due to their high corrosion resistance.²⁵⁻²⁷ Particular interest has focused on the bimetallic nitride $\text{Co}_{0.6}\text{Mo}_{1.4}\text{N}_2$, which combines the activity of Co_xN with the acid stability of Mo_xN .²⁸⁻³⁰

Nickel-based nitride catalysts have also shown some ORR activity. Specifically, a high surface area Ni_3N on Ni-foam catalyst demonstrated low overpotential and good stability for HER and OER, as well as preliminary evidence of activity for ORR in alkaline media.³¹ Ni_3FeN nanostructures have shown promising bifunctional activity, both as supports and catalysts, for metal-air batteries in alkaline media.³²⁻³⁴ Additionally, Ni_3N quantum dots supported on NiO nanosheets have shown good activity and stability for ORR in alkaline.³⁵ Herein, we demonstrate an ORR active nickel nitride catalyst in acid for the first time. Through extensive electrochemical, materials, and theoretical characterization of a well-defined, carbon-free thin film catalyst, we investigate the fundamental ORR activity, selectivity, and stability of nickel nitride in both acidic and alkaline electrolyte. DFT is utilized to construct Pourbaix diagrams and identify the active surface of the catalyst.

1

2 **Materials and Methods**

3 **Materials**

4 Glassy carbon electrodes (Pine Research Instrument, 0.196 cm² geometrical area),
5 graphite counter electrode, platinum wire, silicon wafers (WRS, 100 mm, P/Bor <100>,
6 10-20 Ohm-cm), Ni sputtering target (Kurt J Lesker, 99.99% purity, 2" diameter), Ti
7 sputtering target (Kurt J Lesker, 99.99% purity, 2" diameter), perchloric acid (Honeywell
8 Fluka, 70%), potassium hydroxide (Fisher Chemical, 86.4% assay), were all used as
9 received and without further purification unless otherwise stated.

10

11 **Synthesis**

12 Nickel nitride thin films were prepared by DC reactive sputtering using a Lesker Sputter.
13 Prior to the nitride synthesis, a 20 nm thick Ti sticking layer was synthesized by sputter
14 deposition (3 minutes, 200W, 100% Ar, 3 mTorr) onto the polished glassy carbon disk
15 electrode. Subsequently, without breaking vacuum, the Ni target was sputtered in a
16 mixture of 75% Ar, 25% N₂ plasma to deposit ~130 nm of Ni_xN. The magnetron power
17 supply was maintained at 200 W and the chamber pressure was 8 mTorr. The substrate
18 was held at 180 ± 10 °C with a substrate bias of 100 V throughout the deposition. The
19 Ni₃N and Ni₄N standards used for near edge X-ray absorption fine structure (NEXAFS)
20 spectroscopy were synthesized on silicon substrates; Ni₃N was made by increasing
21 chamber pressure to 18 mTorr, while Ni₄N was made by decreasing the substrate
22 temperature to 90 °C. Ni and NiO thin films were synthesized using the same conditions,
23 but with 100% Ar and 20% O₂ & 80% Ar plasma, respectively. Films were also
24 synthesized on Si wafers, with native SiO₂ layer, for structural characterization.

25 **Physical characterization**

26 Grazing-incidence X-ray diffraction (XRD) data were obtained using a D8 Venture single
27 crystal diffractometer (Bruker, λ=1.5418 Å) at an incidence angle of 5°. Top-down and
28 cross-sectional scanning electron microscopy (SEM) was conducted with a FEI Magellan
29 400 XHR SEM. X-ray photoelectron spectroscopy (XPS) was performed with a Phi
30 Versaprobe 1 using monochromatized Al Kα (1486 eV) radiation. All XPS spectra were
31 calibrated to the C 1s peak at a binding energy of 284.8 eV. CasaXPS software was used
32 to perform peak fitting with Shirley backgrounds.

33 The cross-section TEM specimen was prepared conventionally.³⁶ The samples were
34 glued and mechanically grounded to approximately 15 μm in thickness. The cross-section
35 specimen was further ion milled until electron transparent in a Gatan PIPS II ion milling
36 machine. Argon ion beams with 5 keV energy were induced at an incident angle of 5
37 degrees to create a hole at the center of the specimen. After the formation of the hole,

1 the beam energy was gradually reduced to 0.5 keV for final cleaning. TEM experiments
2 were conducted in a FEI Titan 80-300 environmental transmission electron microscope
3 equipped with a spherical aberration corrector in the image-forming (objective) lens.
4 Selected Area Diffraction Patterns (SADP) were calibrated using a standard Si [110]
5 diffraction pattern taken under the same conditions.

6 NEXAFS data for N and O K-edge and Ni L-edge were collected at the Stanford
7 Synchrotron Radiation Light Source (SSRL) on BL 8-2 and 10-1 in total fluorescence yield
8 (TFY) and total electron yield (TEY) modes, operating the monochromator with 0.2 eV
9 (for N and O K-edges) and 0.3 eV (for Ni L-edge) resolution, respectively. NiO, Ni₃N, and
10 Ni₄N were used as standards. The collected data were normalized to the edge jump after
11 subtraction of the pre-edge region. TFY enables bulk sensitivity with a penetration depth
12 on the order of 100 nm, while TEY has a penetration depth of 5-10 nm.

13 NEXAFS simulations for the N K-edge were conducted with OCEAN code^{37,38} and FEFF
14 9 code³⁹ using a broadening of 1.0 eV and 0.52 eV, respectively, and the crystallographic
15 structures of Ni₄N and Ni₃N. O K-edge was simulated with the FEFF 9 code using a SCF
16 cluster of 6 Å and increasing the radius of the FMS cluster from 3 Å to 9 Å starting from
17 the crystallographic structure of NiO. A broadening of 0.8 eV was applied to simulate the
18 experimental broadening (primarily instrumental, core-hole, and vibrational broadening).
19 The screening of the core-hole was computed using random phase approximation (RPA)
20 and the H-L self-energy was used for the calculation of the exchange correlation potential.

21 **Electrochemical Testing**

22 Electrochemistry was performed using a rotating disk electrode (Pine Research
23 Instrument) in a three-electrode glass cell, with 0.1 M HClO₄ or 0.1 M KOH electrolyte
24 purged with oxygen or nitrogen. An in-house built reversible hydrogen electrode (RHE)
25 was used as the reference electrode and a graphite rod as the counter electrode. The
26 series resistance of the cell was measured at 100 kHz and the iR losses were
27 compensated at 85%. The remaining 15% was corrected after testing. Electrochemical
28 activity was assessed using cyclic voltammetry (CV), sweeping reversibly from 0.8 V to
29 0.05 V at a scan rate of 20 mV/s using a Biologic VSP-300 Potentiostat. Unless otherwise
30 stated, the oxygen purged voltammograms were corrected for background current by
31 subtracting the baseline (nitrogen) sweep. For simplicity, only cathodic CV sweeps are
32 shown in the main text; full CVs are shown in the SI. Stability was evaluated using
33 chronoamperometry, holding at a constant potential of 0.15 V. Selectivity measurements
34 were conducted using a rotating ring disk electrode (Pine Research Instrument) with a Pt
35 ring held at 1.2 V vs RHE to measure the H₂O₂ produced in the reaction. See the SI for
36 details of ring calibration and selectivity calculations.

37 **Computational Details**

1 We performed density functional theory (DFT) calculations using the Vienna Ab Initio
2 Simulation Package (VASP)^{40,41} with the PBE exchange-correlation functional⁴² and
3 projector augmented-wave (PAW) pseudopotentials.⁴³ We added a Hubbard-U correction
4 (PBE+U)⁴⁴ on insulating Ni oxide materials to include the on-site Coulomb interaction of
5 localized electrons. The value of the Hubbard interaction, U, for Ni was chosen to be 6.2
6 eV in accordance with the literature.^{45,46} The energy cutoff, convergence criteria for self-
7 consistent iterations, and for geometry relaxation were set to 500 eV, 10^{-4} eV, 0.05 eV/Å,
8 respectively. Further computational details can be found in the SI.

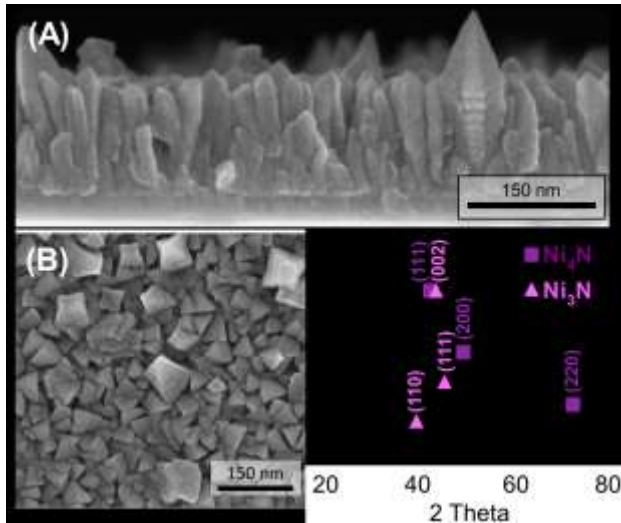
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10 **Results and Discussion**

11 **Characterization**

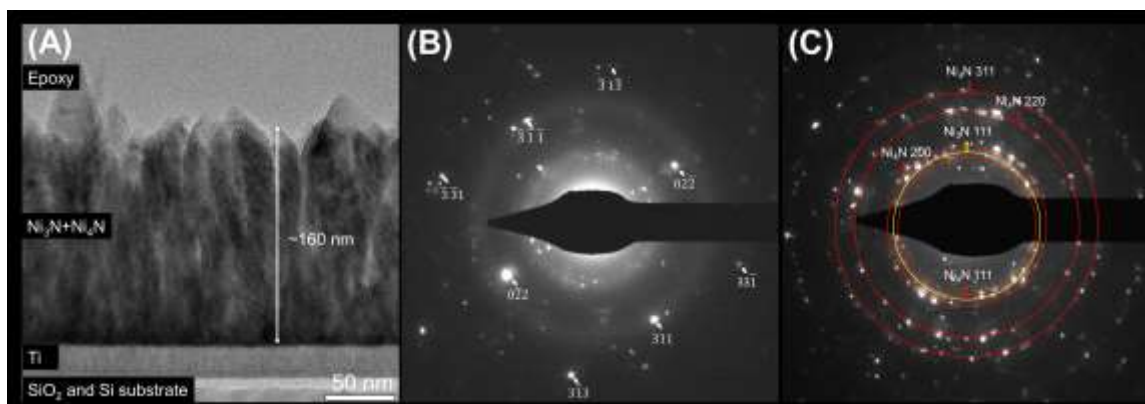
12 The reactively sputtered nickel nitride films are ~160 nm in thickness as shown by cross-
13 sectional scanning electron microscopy (SEM) in Figure 1A. The film consists of well-
14 defined nanopillars, which are variable in height but closely packed to form a rough, dense
15 film. The 20-30 nm Ti sticking layer is visible beneath the pillars. The top down SEM in
16 Figure 1B shows that the nanopillars are pyramids with either square or triangular base
17 shapes. The larger square prisms have an average diameter (distance from opposite
18 corners, viewed from above) of 70 nm, while the triangular prisms are 30-40 nm.

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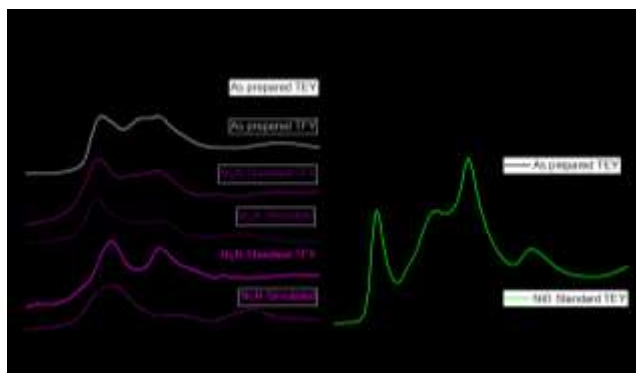
20 **Figure 1.** Characterization of as-prepared, sputtered Ni_xN thin films. (A) Cross-sectional
21 and (B) top down SEM micrographs, (C) XRD pattern with ICDD references and
22 corresponding crystal planes.

23 X-ray diffraction (XRD) of the thin films shows peaks corresponding to both hexagonal
24 Ni_3N (ICDD 00-010-0280 at 2Θ of 39.9° , 42.7° , and 45.2°) and cubic Ni_4N (ICDD 00-036-
25 1300 at 42.0° , 49.3° , and 72.0°), confirming that no other crystalline species are present
26 (Figure 1C).



1
2 **Figure 2.** TEM-SADP (selected area diffraction pattern) characterization of as-prepared
3 Ni_xN thin films. (A) Cross-sectional view, (B) SADP taken from top surface of the film
4 matching diffraction pattern of Ni₄N from $[\bar{2}33]$ zone axis, (C) SADP taken from the bulk
5 region of the film showing diffraction rings of Ni₄N and Ni₃N.

6
7 The films were also analyzed using cross-sectional transmission electron microscopy
8 (TEM). The TEM image of the film in Figure 2A shows a pillar morphology, as seen in
9 SEM, as well as discrete titanium (20 nm thick), silicon dioxide (7 nm thick), and silicon
10 layers. Distinct crystalline regions are also visible (Fig. S1). Selected area diffraction
11 patterns (SADP) were used to determine the crystal structures at the surface and in the
12 bulk, as shown in Figure 2B and C, respectively. The selected area aperture positions are
13 shown in Figure S2. In Figure 2B, the diffraction pattern measured at the surface matches
14 the Ni₄N cubic structure. The measured d-spacings of diffraction spots along the two
15 orthogonal directions are 0.131 nm and 0.111 nm. These match well with the (0 $\bar{2}2$) and
16 (311) planes of the reference Ni₄N. These also suggest that the diffraction pattern was
17 taken near a $[\bar{2}33]$ zone axis of Ni₄N. The diffraction pattern from the bulk region shows
18 diffraction rings (labeled in red) of Ni₄N which correspond to the (111), (200), (220), and
19 (311) planes. This indicates that the bulk of the thin film is polycrystalline. An additional
20 ring (labeled in yellow) was observed between the (111) and (200) rings of Ni₄N. The d-
21 spacing of this ring matches the (111) plane spacing of Ni₃N. The SADP indicates that
22 the film is primarily composed of Ni₄N, with a minor phase of Ni₃N in the bulk. However,
23 it is important to note that the analyzed volume of our polycrystalline film from this method
24 is limited. To probe a broader region of the sample, we employed NEXAFS spectroscopy,
25 as discussed in the next section.



1
 2 **Figure 3.** NEXAFS spectra of as-prepared, sputtered Ni_xN thin films. (A) N K-edge of the
 3 sample, Ni₃N and Ni₄N standards, and simulated spectra; (B) O K-edge of the sample
 4 and NiO standard. Dashed lines are intended as guides for the reader. TEY: surface;
 5 TFY: bulk.

6 To further probe the structure and chemical state of the catalyst, the films were
 7 characterized using NEXAFS spectroscopy. In Figure 3A, the N K-edge NEXAFS spectra
 8 for the as-prepared catalyst are compared with experimental and simulated spectra for
 9 Ni₃N and Ni₄N standards. The spectra were collected in TEY and TFY modes, allowing
 10 for a comparison between the surface (TEY: few nm) and the bulk of the film (TFY). Two
 11 main peaks are present both for the surface and the bulk of the as-prepared sample,
 12 though differences are evident between the two spectra. For TEY, the feature at 397.5
 13 eV matches well with both the experimental and simulated spectrum of Ni₄N. The TFY
 14 signal shows a shift of the peak at 397.5 eV towards higher energy and the appearance
 15 of a shoulder corresponding to Ni₃N. We can thus confirm that the catalyst is a mixture of
 16 Ni₄N and Ni₃N, as indicated by XRD and TEM analyses, and locate Ni₃N in the bulk. The
 17 feature at 400.5 eV is associated with the oxidation of the surface of the nitride film upon
 18 exposure to air after synthesis, as has been previously reported for other transition metal
 19 nitrides.^{47,48} Notably, the fact that the intensity of this feature considerably decreases in
 20 the bulk of the material provides evidence for the presence of a surface N state different
 21 than in the pure bulk nitride film. This is also observed in the experimental spectra for the
 22 Ni₄N and Ni₃N standards (Figure S3).

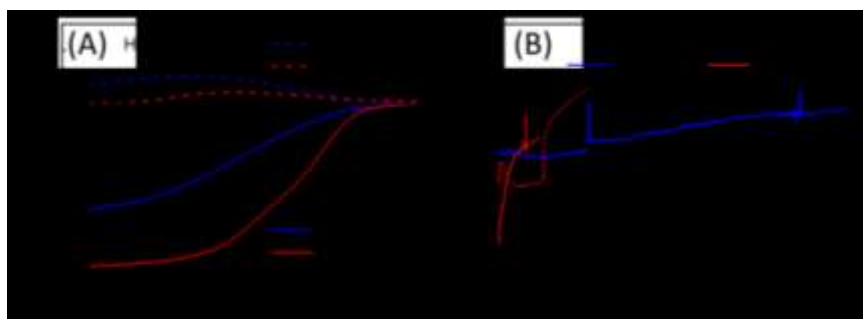
23 Figure 3B shows experimental O K-edge spectra for the catalyst and the NiO standard,
 24 which allow for further characterization of surface oxidation. The TFY signal for O was
 25 negligible, confirming that the oxidation process is limited to the surface and there is no
 26 oxygen incorporation in the bulk. The signal for the oxidized layer on top of the nitride film
 27 consists of a peak at 532.8 eV, shifted to higher energy compared to the bulk NiO, and a
 28 broad feature at 540.5 eV, which has been reported for small native Ni oxide clusters on
 29 top of Ni films.⁴⁹ It was concluded that none of the defined features typical of crystalline
 30 NiO were detected due to the amorphous nature of the small NiO domains. O K-edge
 31 spectra for NiO clusters of various dimensions were simulated (Fig. S4). The simulated

1 spectrum for a NiO cluster with ca. 5 Å radius (57 atom cluster) is the best match to the
2 experimental, suggesting the presence of these clusters on the nitride surface. The Ni L3-
3 and L2-edge spectra also support the presence of oxidized Ni₄N on top of Ni₃N (Fig. S5).
4 Our spectroscopic measurements thus indicate that prior to testing, the catalyst structure
5 consists of small clusters of NiO on top of Ni₄N, with Ni₃N crystallites also present in the
6 bulk.

7

8 **Electrochemistry**

9 The ORR activity and selectivity of the nickel nitride catalyst was assessed using a
10 rotating ring disk electrode (RRDE) in both acidic (0.1 M perchloric acid) and alkaline
11 (0.1 M potassium hydroxide) electrolyte. Figure 4A shows the activity and selectivity of
12 the synthesized catalyst, with the ORR polarization curve measured on the disk
13 electrode (solid lines) and the H₂O₂ oxidation current measured on the platinum ring
14 electrode (dashed lines). Comparison with the activity of a commercial Pt/C (46.6% from
15 TKK) catalyst is shown in Figure S6.⁵⁰ In acid, the ORR onsets (defined as the potential
16 required to reach 100 μA/cm² geometric current density) at 0.68 V vs RHE and reaches
17 the mass transport limited current density of 6.1 mA/cm² at 0.2 V vs RHE. Furthermore,
18 the halfwave potential is 0.49 V vs RHE, indicating a relatively sharp onset. This is
19 promising activity for a non-Pt group metal catalyst in acid and is comparable to the
20 metal nitrides currently reported in the literature (Table S3).^{51–53} Nickel nitrogen-doped
21 carbon (Ni-NC) type catalysts have also demonstrated good ORR performance, but
22 they cannot be directly compared with this work due to their high surface area and the
23 contributions of the NC support.^{54,55} In alkaline electrolyte, nickel nitride shows similar
24 activity with an onset at 0.68 V vs RHE but has a lower mass transport limited current
25 density of 4 mA/cm², as well as a slower onset with a halfwave potential of 0.42 V vs
26 RHE. This difference in activity is reflected in the Tafel slopes: 89.7 mV/decade in acid
27 compared to 175 mV/decade in alkaline (Fig. S7). Similar pH dependent Tafel slope
28 trends have been reported in the literature.⁵⁶ Aside from the Tafel slopes, PGM-free
29 catalysts are generally more ORR active in alkaline, with higher onset and halfwave
30 potentials (Table S3). The reversal in that trend for this nickel nitride system can likely
31 be attributed to different H₂O₂ selectivity and different surface structures under reaction
32 conditions, which will be discussed in the following sections.



1
 2 **Figure 4.** Electrochemical performance of Ni_xN in acid (red) and base (blue). (A) RRDE
 3 measurements showing ORR polarization curves and H₂O₂ current density for nickel
 4 nitride. (B) Chronoamperometry (CA) profile at 0.15 V vs RHE in acid, for 2 samples
 5 tested under different cycling parameters, and in base.

6 For PEMFCs, high selectivity to the 4-electron reduction, corresponding to a low H₂O₂
 7 current, is preferred. In acid, the selectivity of the catalyst to H₂O₂ (2-electron reduction)
 8 was found to decrease with increasing overpotential and was negligible at potentials
 9 below 0.3 V vs RHE. (Fig. S10). The H₂O₂ current was used to calculate the electron
 10 transfer number, *n*, which was found to range from 3.2 (at 0.6 V vs RHE) to 4.0 (at 0.1 V
 11 vs RHE) (see SI for calculations). In alkaline electrolyte, nickel nitride has higher
 12 selectivity towards H₂O₂, with *n* values ranging from 2.4 at 0.6 V vs RHE up to 3.4 at 0.1
 13 V vs RHE. This high selectivity to H₂O₂, as well as the onset of ORR activity close to the
 14 thermodynamic limiting potential of 0.70 V vs RHE for 2-electron ORR, indicates that in
 15 alkaline electrolyte the catalyst is unable to turnover 4-electron ORR until the onset of 2-
 16 electron ORR. This limitation to the thermodynamic potential of 2-electron ORR likely
 17 contributes to the lower activity in alkaline electrolyte.

18 In addition to activity and selectivity, stability is a crucial performance metric for any
 19 catalyst. Here, electrochemical stability was assessed using chronoamperometry, and the
 20 use of post-test characterization and theoretical calculations to evaluate chemical stability
 21 will be discussed in later sections. Figure 4B shows stability tests for nickel nitride in acid
 22 and alkaline electrolyte. After 3 CVs, catalyst stability was evaluated by
 23 chronoamperometry (CA) at 0.15 V vs RHE. CVs were run every 5 h to assess changes
 24 in activity (Fig. S11).

25 In alkaline, the catalyst improved over the first 5 h of the chronoamperometry hold,
 26 increasing from 3.3 to 3.5 mA/cm² of current density in the CA, and the CV after 5 h
 27 demonstrated improvement of 25 mV and 75 mV in onset and half-wave potential,
 28 respectively (Fig. S11B). Subsequent CVs show a slow degradation, with a loss of
 29 approximately 0.2 mA/cm² every 5 h from 10 h to 30 h. The noise in the data, observed
 30 at 3 h, is due to bubble formation on the surface of the catalyst. The bubble was removed
 31 mechanically and had no lasting effect on catalyst performance. All other spikes
 32 correspond to switching between the CA and CVs. The catalyst continued to degrade

1 slowly, with current density losses of 25% at 24 h and 40% at 40 h, at which time the
2 stability test was ended.

3 Interestingly, in acid, the activity and stability of the nickel nitride catalyst was found to be
4 dependent on the cycling parameters of the cyclic voltammogram (CV). To illustrate this
5 dependence, Figure S12 shows cathodic sweeps, with no background subtraction, for
6 two sibling catalysts in acid. The catalysts were tested identically, except that the sweeps
7 started either at more positive (0.8 V vs RHE), as was shown in Figure 4A, or less positive
8 (0.15 V vs RHE) potential. The “more positive start” catalyst shows superior activity, with
9 both an earlier onset and larger limiting current density. However, it loses activity more
10 rapidly, producing only 3 mA/cm² after 5 h, corresponding to a 50% loss in activity. CVs
11 during the stability test for the “less positive start” catalyst are shown in Figure S11A, with
12 relatively similar ORR activity at 0 and 5 h, but a significant loss of activity during the cycle
13 at 10 h. These differences in activity and stability between these testing protocols, as well
14 as the losses caused by the CVs during the CA, provide evidence that the catalyst is very
15 sensitive to applied potential.

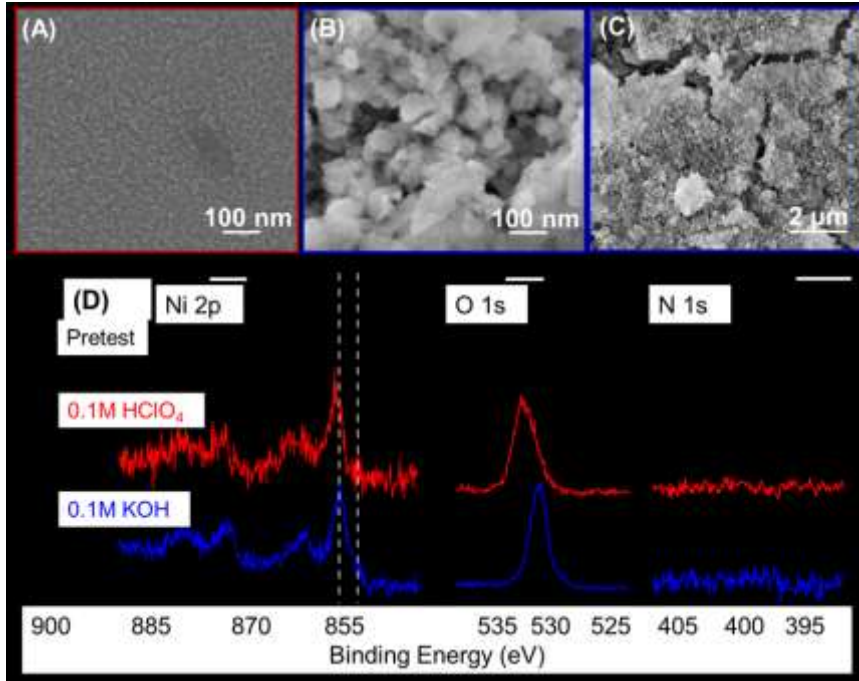
16 Sensitivity to potential is likely a consequence of surface oxidation reactions which occur
17 in the ORR potential window.⁵⁷ In both N₂ and O₂ scans, there was an irreversible
18 oxidative peak on the anodic sweep in the 0.65 – 0.8 V vs RHE region during the first 4
19 sweeps (Figure S13). Between sweeps 1 and 3, this results in a positive shift in onset
20 potential by 100 mV and a 2 mA/cm² gain in mass transport limited-current density. This
21 oxidation peak shrinks after 4 sweeps, indicating that further oxidation of the catalyst was
22 limited. This coincided with a loss of activity, with current density decreasing to zero over
23 the subsequent 4 sweeps. We therefore hypothesize that the active surface is a partially
24 oxidized nitride and that the activity is suppressed, as the oxidation progresses, through
25 decreased conductivity and dissolution. This reveals the sensitivity of the catalyst to open
26 circuit potential and oxidizing potentials; durability could be improved by maintaining the
27 catalyst at reducing currents.

28

29 **Characterization After Electrochemical Testing**

30 To better understand chemical stability and investigate the mechanisms of failure in acid
31 and base, the films were characterized after activity (CV) and stability (CA) testing using
32 SEM and XPS. After the 10 h CA in acid (Figure 5A), SEM shows that the surface
33 structure had changed completely, with most of the material gone. This structural change
34 likely contributes to the loss in performance over time. After the 40 h CA in base (Figure
35 5B,C), the structure is distinct from the as-prepared catalyst. The surface was covered
36 with rod-shaped structures, loosely packed into clusters approximately 0.1 μm² in area,
37 and large cracks (approximately 300 nm across) appeared in the film. We hypothesize

1 that these cracks may have undercut the film, disturbing its electrical connectivity and
2 performance.



3
4 **Figure 5.** Top down SEM images of the nickel nitride films after (A) stability test in acid,
5 and (B, C) stability test in base. (D) High-resolution XPS spectra for as-prepared (black),
6 post-stability acid (red), and post-stability base (blue) for Ni 2p, O 1s, and N 1s regions.
7 Dashed lines are intended as guides for the reader.

8
9 The XPS spectra (Figure 5D) also show a change in elemental composition. Before
10 continuing with this discussion, it is necessary to note that accurate quantitative analysis
11 of this XPS data is difficult due to the complexity of the Ni 2p spectrum, so all statements
12 of composition are meant as best approximation. Elemental quantification of the nickel
13 nitride revealed a ratio of Ni to N of approximately 3.5, consistent with a mixture of Ni₃N
14 and Ni₄N observed in XRD. Prior to testing, peaks were identified in the Ni 2p_{3/2} region at
15 binding energies of 853.0 and 855.2 eV (dashed lines), corresponding to the interstitial
16 nitride (Ni^{δ+}, δ = 0-2) and oxide (Ni²⁺), respectively. There are satellite peaks in the region
17 860 – 865 eV. This matches well with literature spectra for nickel nitride.⁵⁸ In the O 1s
18 region, the low energy peak at 529.3 eV corresponds to the nickel oxide, while we
19 associate the higher energy peak with hydroxide (literature value of 531.1 eV) and
20 carbonyl species (most likely carbonate, literature value of 532.8 eV).⁵⁹ Finally, the N 1s
21 peak at 397.6 eV corresponds to a metal nitride.

22 After stability testing in acid, the surface composition changed drastically. The Ni 2p_{3/2}
23 peaks shifted to higher binding energies, with the largest peak at 855.8 eV, indicating the

1 Ni²⁺ oxidation state. The N 1s peak disappeared, corresponding to a loss of nitrogen from
2 the surface. The O 1s peaks increased in intensity relative to the Ni and indicate the
3 presence of hydroxide and carbonate species (from air exposure prior to
4 characterization).⁶⁰ In conjunction with the structural changes observed by SEM, this
5 indicates an extensive reorganization of the surface during testing, resulting in a surface
6 that more closely resembles Ni(OH)₂ than the original Ni₄N.⁶¹ Similar composition
7 changes were observed after CV testing in acid (Figure S14). From XPS it is not possible
8 to determine when these changes occurred, though it seems likely that they were caused
9 by exposure to oxidizing potentials during testing or open circuit potential when the
10 catalyst is removed from electrolyte. These structural and composition changes indicate
11 that overall chemical stability is poor in acid. However, an analysis of potential corrosion
12 current, as well as the poor ORR activity of the underlying Ti and glassy carbon in acid,
13 indicates that the Ni_xN catalyst is responsible for the current observed during the stability
14 test (see SI for details).

15 The composition change after stability testing in base is similar. The primary remaining
16 Ni 2p_{3/2} peak is at 855.9 eV and there is no N 1s signal at the surface, indicating a
17 complete conversion at the surface to a hydroxide structure. The O 1s spectrum shows a
18 small oxide peak, but hydroxide is the dominant species. Comparatively, the composition
19 changes after CV testing are minor, with nitride composition retained (Fig. S14). This was
20 confirmed by the bulk NEXAFS spectra, which showed no significant change between
21 this and the as-deposited sample (Fig. S15). This electrochemical, microscopic, and
22 spectroscopic investigation of stability and failure mechanisms has revealed that activity
23 loss is correlated with the conversion of the nitride to hydroxide and associated material
24 property changes. Due to the sensitivity of the activity and stability of these catalysts to
25 applied potential, future work should focus on probing the stability of these catalysts as a
26 function of applied potential. We hypothesize that stability would be more challenging at
27 more oxidizing conditions, since extensive catalyst oxidation was found to detrimentally
28 impact activity and stability.

29 To gain insight into oxide phases observed experimentally, we evaluated the stability of
30 oxide phases in the presence of Ni₃N and Ni₄N with DFT. The optimized bulk structures
31 of Ni₃N and Ni₄N are in good agreement with the XRD and SADP results, as well as
32 previously reported bulk parameters ($a=b=4.619$ and $c=4.307$ Å for Ni₃N, $a=b=c=3.730$ Å
33 for Ni₄N).^{62,63} We modeled $\sqrt{3} \times \sqrt{3}$ structures for Ni₃N (0001) and Ni₄N (111) to minimize
34 the lattice mismatch between nickel nitrides (Ni₃N and Ni₄N) and nickel oxides (NiOOH
35 and NiO). In the next step we simulated monolayer films of NiOOH and NiO supported on
36 Ni₃N and Ni₄N (Figure 6A). We considered all possible adsorption sites and surface
37 termination to minimize the oxide nitride interface formation energy. The formation
38 energies of nickel oxides films (NiOOH and NiO) on nickel nitrides (Ni₃N and Ni₄N) are
39 calculated as following:

1
$$E_{\text{Formation}} = [E_{\text{slab,oxide+nitride}} - E_{\text{slab,nitride}} - nE_{\text{bulk,oxide}}]/n,$$

2 Where $E_{\text{slab,oxide+nitride}}$, $E_{\text{slab,nitride}}$ and $E_{\text{bulk,nitride}}$ are electronic energies of the total
 3 system, slab structure of nitrides and bulk nickel oxides. The n is the number of oxide
 4 units in the total system. The most stable geometries for NiOOH and NiO on Ni₃N and
 5 Ni₄N are shown in Figure 6A and Table 1 summarizes the results of formation energy
 6 calculations. These results show that NiOOH layer is effectively stabilized by -0.67 and -
 7 0.53 eV/f.u on Ni₃N and Ni₄N, respectively, while NiO is significantly destabilized.

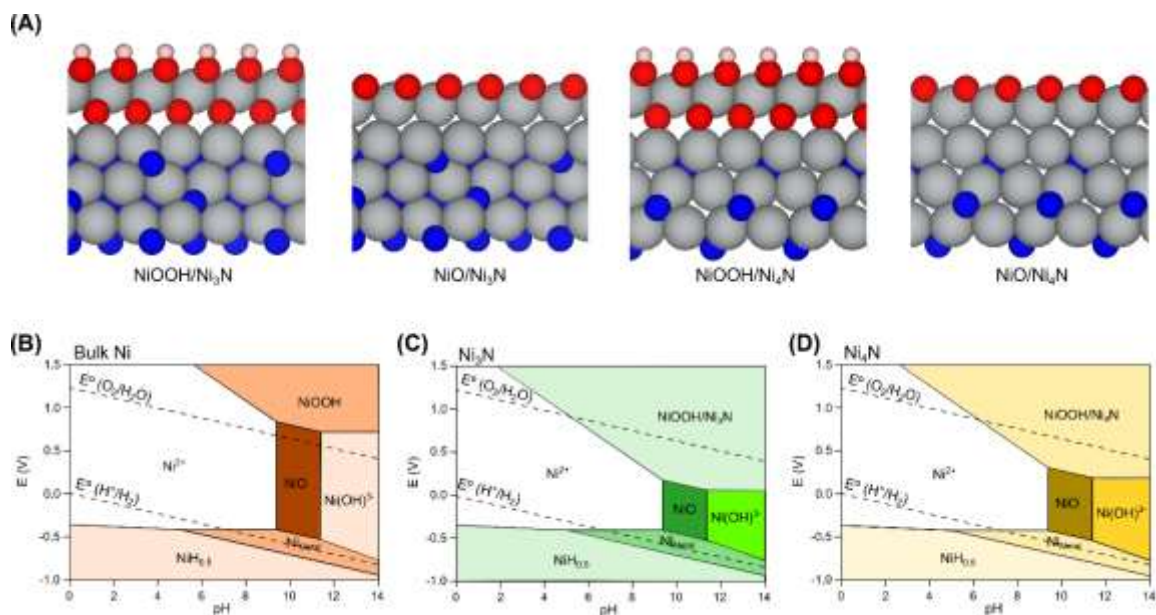
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9 **Table 1.** The calculated formation energy of Ni oxide species on Ni nitrides.

Formation energy (eV/f.u.)	Ni ₃ N	Ni ₄ N
NiOOH	-0.67	-0.53
NiO	1.60	1.64

10

11 To further elaborate the formation of different oxide phases during ORR testing, as well
 12 as differences in stability between acid and alkaline electrolyte, we constructed Pourbaix
 13 diagrams. Figure 6 shows (A) the most stable geometries of the NiO and NiOOH
 14 overlayers as well as the theoretically calculated Pourbaix diagrams for (B) bulk Ni, (C)
 15 Ni₃N, and (D) Ni₄N. The region of interest for ORR is 0.2 V_{RHE} to 1.23 V_{RHE} , below the
 16 upper dashed line in the Pourbaix diagram. The strong formation energy of NiOOH on
 17 both Ni₃N and Ni₄N results in the expansion of NiOOH area in the Pourbaix diagrams (Fig
 18 6C and D). Hence, the observed oxidized phase in the experiment can be attributed to
 19 the stabilized NiOOH on the nickel nitride surfaces, particularly in alkaline electrolyte. This
 20 analysis also shows that nickel nitride surfaces are not stable under acidic environment
 21 due to dissolution of Ni²⁺ ions. Differences in surface composition and coverage between
 22 these electrolytes may contribute to the observed differences in ORR activity and
 23 selectivity.



1
 2 **Figure 6.** (A) Optimized geometries of NiO and NiOOH supported on Ni₃N and Ni₄N,
 3 respectively. Pourbaix diagrams of (B) bulk Ni, (C) Ni₃N, and (D) Ni₄N. (B) is based on
 4 the experimental values of formation free energies of bulk phases^{62,63}, and the
 5 stabilization energies of nickel nitrides on NiOOH were added to construct (C) and (D).
 6 NiO is destabilized on the nickel nitrides, thus the bulk formation free energy was used.
 7 Color codes: silver (Ni), blue (N), red (O), pink (H). The equilibrium potentials of O₂/H₂O
 8 (1.23 V_{RHE}) and H⁺/H₂ (0.00 V_{RHE}) are marked with dashed lines.

9 The above theoretical investigation provides a basic understanding of the formation of
 10 possible nickel oxide phases on the examined nickel nitrides, which is of particular interest
 11 due to recent DFT calculations showing that the formation of an oxide over-layer on cobalt
 12 nitride promotes ORR.⁶⁴ This is in agreement with our experimental observation of
 13 extensive surface oxidation after ORR testing. While the XPS and NEXAFS results
 14 indicate a surface hydroxide species, these are *ex situ* measurements, taken after the
 15 catalysts had been exposed to air, and thus cannot be used to determine the surface
 16 composition under reaction conditions. This will be the subject of future investigations
 17 utilizing operando spectroscopies.

18 It is important to note that the Pourbaix diagram is based on thermodynamics and does
 19 not provide any information about the kinetics of the phase transition. Thus, while the
 20 nickel nitride is not predicted to be stable in acid, several hours of stability were achieved,
 21 perhaps because the oxidation or dissolution processes are slow kinetically. This can
 22 serve as a design principle for further catalyst development. Superior stability is observed
 23 in base, where the catalyst is more thermodynamically stable.

24 The use of a well-defined and carbon-free catalyst morphology facilitated a combined
 25 experiment-theory approach to determining the active surface of the catalyst. With the

1 insights into the origins of activity and stability obtained through the study of this model
2 system, efforts to improve this catalyst fall into two main categories.⁶⁵ First, intrinsic
3 activity and stability can be improved through alloying with a more stable transition metal
4 nitride, which has been demonstrated for several nitride systems.^{22,23,29} Based on the
5 Pourbaix calculations and the potential of activation via oxide over-layer formation,⁶⁴ a
6 promising candidate would be a metal nitride which forms a stable, passivating oxide or
7 hydroxide surface in acid and alkaline electrolyte. Second, after discovery of a promising
8 material system, the number of active sites can be increased through nanostructuring and
9 supporting the catalyst on a high surface area material such as N-doped carbon.^{54,55}

10

11 **Conclusions**

12 In summary, nickel nitride thin films were found to be active ORR catalysts, achieving
13 activity comparable to the metal nitrides reported in the literature. Reactive sputtering is
14 highlighted as a promising route to synthesize transition metal nitride electrocatalysts with
15 a variety of compositions and crystal structures, allowing for fundamental investigation of
16 the active surface. By synthesizing the catalyst in a carbon-free morphology, we were
17 able to use a combined experiment-theory approach to isolate the activity of the nickel
18 nitride and found that it is an active and reasonably stable non-precious metal catalyst for
19 ORR in acid, as well as in base. In future work, catalyst activity and stability could be
20 improved through alloying and nanostructuring. *Ex situ* XAS and XPS characterization
21 indicated that an oxide overlayer readily forms on the catalyst surface. DFT calculations
22 predicted similar surface oxidation under reaction conditions, which was shown to
23 correlate with changes in electrochemical activity and stability. This surface oxidation can
24 serve as a design principle for a variety of metal nitride catalysts and demonstrates the
25 importance of complementary experimental and theoretical studies.

26

27 **Associated Content**

28 Supporting Information available: Supplementary experimental characterization (TEM,
29 XRD, NEXAFS, electrochemistry, and XPS), and details for H₂O₂ selectivity, Tafel, and
30 TOF calculations. Raw data from all main text figures are also provided, in line with the
31 U.S. Department of Energy requirements.

32

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9 Notes

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24

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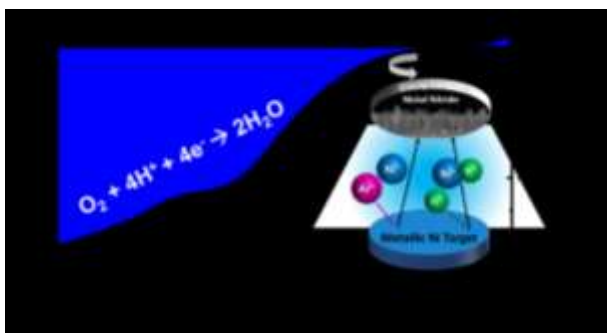
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1 TOC image:



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