Covalent Functionalization of Nickel Phosphide Nanocrystals with Aryl-Diazonium Salts

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Abstract
Covalent functionalization of Ni2P nanocrystals was demonstrated using aryl-diazonium salts. Spontaneous adsorption of aryl functional groups was observed, with surface coverages ranging from 20-96% depending on the native reactivity of the salt as determined by the aryl substitution pattern. Increased coverage was possible for low reactivity species using a sacrificial reductant. Functionalization was confirmed using thermogravimetric analysis, FTIR and X-ray photoelectron spectroscopy. The structure and energetics of this nanocrystal electrocatalyst system, as a function of ligand coverage, was explored with density functional theory calculations. The Hammett parameter of the surface functional group was found to linearly correlate with the change in Ni and P core-electron binding energies and the nanocrystal’s work-function. The electrocatalytic activity and stability of the functionalized nanocrystals for hydrogen evolution were also improved when compared to the unfunctionalized material, but a simple trend based on electrostatics was not evident.

TOC Graphic

Introduction
Great progress has been made over the past few decades in the development and deployment of renewable energy technologies. This global effort has positioned sustainably generated electricity to, in some cases, be cheaper than carbon-based competitors.1 While renewably-sourced electricity will be a key factor in decarbonizing the power and light passenger transportation sectors, there is still significant work to be done in decarbonizing other carbon intensive sectors such as industrial transportation and chemical commodities.2-4 To achieve the same emissions reductions in these sectors, we must fundamentally alter the way we produce and recycle small molecule chemical feedstocks such as H2, O2, N2, and CO2. The key component to enabling this
kind of sustainable circular economy is the development of efficient, selective, and economically viable (photo)electrocatalysts.5

Today, the most active catalytic materials are scarce noble metals such as platinum, palladium, and iridium. Specifically in the case of the hydrogen evolution reaction (HER), the high activity of these materials has been ascribed to their ideal metal-hydride (M-H) bond dissociation free energies (BDFEs), their diffuse frontier orbitals that span the Fermi level, and their stability in highly acidic and basic media.6–8 The relationship between these types of “descriptors” and the intrinsic activity of a catalyst is still being investigated and qualified by many computational and electroanalytical groups.5,9–15 Descriptor-based models for understanding catalytic activity have led to the discovery of earth-abundant catalytic materials with relatively high intrinsic activity.5 Initial efforts in this space were focused, in some instances, on the investigation of molybdenum disulfide due to its solution processability, low toxicity, and long history of use as a hydrodesulfurization catalyst.16–19 This same line of theory driven inquiry led to the investigation of transition metal phosphides as a new class of materials with excellent catalytic activity for HER and high stability in acidic media. Research efforts over the past decade have explored many aspects of these materials including synthetic methodology and mechanisms20,21,30,22–29, the effects of varying atomic composition31–34, nano-structuring35–37, and computational modeling of their catalytically active surface sites.31,38–44

Despite great progress in developing this class of materials, earth abundant alternatives still present orders of magnitude lower activity than their precious metal counterparts. This begs the question many researchers are now investigating, “what are we missing in the current descriptor-based models?” If we can already achieve thermoneutral surface binding sites, maybe the answer lies just beyond the surface in the secondary and outer substrate coordination spheres?

More than a century of metal-based molecular (homogeneous) catalyst development highlighted the pivotal role of ligands in determining the chemical properties of a metal center, such as redox potential and metal-substrate BDFE.9,45–48 It is also clear that the way ligands interact with substrate and solvent, through secondary and outer-coordination sphere effects, have a dramatic influence on the rate of catalytic turnover and product selectivity.49–52 We propose hybrid inorganic nanomaterials with tailor-made organic surface ligands could offer a strategy to translate the lessons learned in molecular systems to nanoparticles and represent a vast and underexplored parameter space in heterogeneous electrocatalysis.53–55

Previous studies from our laboratory have demonstrated the importance of the interface between heterogeneous nanoparticle surfaces and the bulk solvent.56,57 Through this work, for instance, we found that the predominant reason long chain aliphatic ligands hinder catalytic activity in aqueous media is because of the poor interface formed between the non-polar surface and the polar electrolyte. Just as aliphatic ligands make a poor interface with polar solvents, it has been shown that polar ligands that are able to hydrogen bond with a polar solvent can induce some degree of local order near the surface of a nanoparticle.58 We also know that surface ligands are able to shift the Fermi level of a nanomaterial based on their electron withdrawing/donating ability by inducing or enhancing the surface dipole.59–61 These isolated studies provide, in concert, a road map to tune a variety of catalytically relevant parameters in heterogenous nanoparticle catalysts. In colloidal nanoparticle systems, ligand chemistry is most commonly explored using dative Lewis acid/base interactions between surface metal cation sites and electron rich binding groups such as amines, carboxylates, and phosphonates.62,63 Well understood mechanisms for ligand exchange have been developed based on concentration gradients, salt metathesis, and displacement by higher binding affinity head groups.64–70 Unfortunately, the same chemical principles that make
these ligand exchange reactions viable also make them unstable in catalytically relevant conditions. Especially the strong acid conditions, typically 0.5 M sulfuric acid, required for HER lead to the protonation and desorption of such ligands.

Thus, to explore the influence that ligand surface chemistry can have in altering the catalytic activity and stability of colloidal nanoparticles, we must first establish synthetic methods to create more robust nanoparticle-ligand bonds. One such method that has been explored in the literature is the covalent functionalization of surfaces by substituted aryldiazonium salts. Diazonium salts are a diverse class of organic reagents that have been shown to form covalent bonds with electron-rich surfaces.\textsuperscript{71–76} Specifically, Saveant showed decades ago that these reagents could functionalize the surface of glassy carbon electrodes and that these functional groups were stable to physical abrasion and electrochemical cycling.\textsuperscript{77} More recently, diazonium reactivity has been explored with two-dimensional materials, namely MoS\textsubscript{2} nanosheets.\textsuperscript{78,79} Miller and co-workers explored the effects of covalent surface functionalization on the catalytic activity of MoS\textsubscript{2} for HER and found strong correlations between the functional group Hammett parameter, MoS\textsubscript{2} electronic structure, and overpotential for HER.\textsuperscript{61} Though this chemistry has never been explored on colloidal heteroatomic nanoparticles, specifically transition metal phosphides, there is precedent for the approach being viable for P-functionalization based on studies on phosphorene.\textsuperscript{78,80}

Herein, we explore the synthetic conditions required to make a well-defined covalent ligand shell on the surface of colloidally synthesized Ni\textsubscript{2}P nanocrystals. The effects of substituted aryl ligands on electronic structure, catalytic activity, and stability are presented using a combination of spectroscopic, electrochemical measurements, and density functional theory (DFT) calculations. Finally, synthetic challenges and design principles for future development of colloidal nanocrystal electrocatalysts are presented.

**Experimental Section**

2-propanol (anhydrous, 99.5%), triethyloxonium tetrafluoroborate (≥97.0%, Meerwein’s reagent), acetonitrile (anhydrous, 99.8%), chloroform (anhydrous, ≥99), 4-nitrobenzenediazonium tetra fluoroborate (97%), 4-methoxybenezenediazonium tetrafluoroborate (98%), 3,5-dichlorobenzenediazonium tetrafluoroborate, and bi(cyclopentylidienyl)cobalt(II) were all purchase from Millipore-Sigma, stored in a nitrogen atmosphere glovebox, and used without further purification. Diazonium salts and Meerwein’s reagent were stored at -20 °C in a glovebox freezer. Tris-diethylaminophosphine (97%, Millipore-Sigma) was stored in ambient conditions and used as received. Nickel (II) chloride (98%, Millipore-Sigma) was dried at 100 °C under vacuum overnight before being stored in a nitrogen glovebox until use. Oleylamine (technical grade, 90%), was dried over CaH\textsubscript{2}, distilled, and stored over 4 Å sieves in a nitrogen glovebox.

**Synthetic Details.**

Ni\textsubscript{2}P nanocrystals were synthesized using air-free Schlenk techniques under an N\textsubscript{2} atmosphere. Glassware, including a 100 mL three neck round bottom flask (RBF), two condensing columns (used without flowing water), a hose adapter, glass thermal well, Teflon magnetic stir bar, 100 mL Schlenk flash, canula needle, and 14/20 ground glass stopper were all dried in the oven at 160 °C overnight prior to use. The next day, the reaction vessel was assembled on top of a heating mantle and connected to the Schlenk line, evacuated, and refilled with dry N\textsubscript{2}. In an N\textsubscript{2} glovebox, 944 mg (7.2 mmol) of NiCl\textsubscript{2} was measured. NiCl\textsubscript{2} was quickly transferred out of the glovebox and added to the reaction vessel. The vessel was purged with high vacuum/N\textsubscript{2} flushes three times. 48 mL (146 mmol or 20 mol eq. relative to Ni) of oleylamine was added to the reaction vessel under an
N₂ atmosphere through a rubber stopper. The mixture of NiCl₂ and oleylamine was heated to 120 °C and vigorously stirred under vacuum for 60 min to remove trace O₂/H₂O and volatile organic contaminants. The vessel was wrapped in glass wool to promote uniform heating. The reaction temperature was measured by an internal probe, submerged in the reaction vessel through contact with a glass thermal well. After 60 min, the reaction mixture was cooled to ~50 °C. 8 mL (28.8 mmol or 4 mol eq. relative to Ni) tris-diethylaminophosphine was injected through the rubber stopper. The reaction vessel was then heated to 250 °C at a ramp rate of roughly 10 °C min⁻¹ and held for 60 min. After about 12 min, at 178 °C, a red hue was observed. The solution quickly became black as the temperature climbed above 200 °C. At this point vapors were observed in the vessel. After 1 hour at 250 °C the vessel was lifted from the heating mantle and the glass wool was removed, allowing the solution to cool.

As the solution cooled, an oven dried Schlenk flask was attached to the Schlenk line and purged with three flushes of vacuum/N₂ and finally held under N₂ flow. Once the reaction mixture reached ~80 °C it was transferred to the Schlenk flash via canula. The Schlenk flask was then sealed with a greased ground glass stopper which was fastened with electrical tape and brought into the glovebox. The reaction mixture was separated into 6 individual 45 mL Falcon tubes, diluted with 2x volume of 2-propanol, and centrifuged at 7.83 kRPM for 15 min. This was repeated until the supernatant for all tubes was clear. The nanocrystals were redispersed in minimal pentane and diluted with 10x volume of 2-propanol before centrifugation at 7.83 kRPM for 15 min. This process was repeated three times, and then another three times using a toluene solvent/acetonitrile antisolvent (1:10) mixture. Finally, the cleaned product was dissolved in minimal chloroform and transferred to a tared 20 mL scintillation vial. Once the solvent was removed by vacuum the product appeared as a free-flowing black powder. 842 mg of Ni₄P was recovered.

The treatment of Ni₂P with Meerwein’s salt was carried out in a 250 mL round bottom flask that was dried a 160 °C overnight before use. 100 mg of Ni₂P was dissolved in 40 mL of toluene and 1.425 g of Meerwein’s salt was dissolved in 50 mL of acetonitrile (0.1 M and 50 mol eq. relative to oleylamine, assuming it accounts for 20% of the Ni₂P sample mass). These solutions were mixed and stirred vigorously for a few minutes to make a homogeneous dark black solution, then left still overnight. The next day a black precipitate was observed at the bottom of the flask. The clear solution was poured off, and the black powder was collected by addition of minimal fresh acetonitrile. Stripped particles, Ni₂Ps, were then moved to a 45 mL Falcon tube where another 10 mL of acetonitrile was added. The sample was centrifuged at 7.83 kRPM for 15 min, resulting in a clear supernatant that was decanted. This process was repeated twice more. Washed Ni₂Ps was then transferred to a tared 20 mL scintillation vial as a solution in toluene, and then dried by vacuum.

Surface functionalization reactions were carried out in an N₂ glovebox in 20 mL scintillation vials that were oven dried overnight at 160 °C before use. 40 mg of Ni₂Ps (when corrected for 15 mass% oleylamine, this translates to 34 mg Ni₂P or 0.229 mmol P) was dissolved in 10 mL of chloroform and added to a 20 mL scintillation vial with a Teflon stir bar and set to stir at 700 RPM. While stirring, in the cases that a chemical reductant was added, 0.115 mmol (2 mol eq. to surface P, SI. S3) of the reducing agent (bi(cyclopentylidenyl)cobalt(II) or cobaltocene) was measured and added to the chloroform solution. Then, 0.115 mmol (2 mol eq. to surface P, SI. S3) of the respective diazonium salt was weighed out and dissolved in 10 mL of acetonitrile. The diazonium solution was added to the vial of stirring Ni₂Ps in a slow, dropwise manner. The effervescence of N₂ in the form of small bubbles was observed after addition of the diazonium, signaling the initiation of radical formation. Solutions were allowed to stir while lightly capped for 48 hours.
For purifying the functionalized nanocrystals, the solvent was first removed under vacuum. Once dried, 5 mL of clean acetonitrile was added to the residue and shaken to disperse. This solution was transferred to a 45 mL Falcon tube and centrifuged at 5 kRPM for 10 min. The supernatant, which had a faint yellow color, was either discarded or collected for analysis. Washing and centrifugation using fresh acetonitrile was repeated 3 more times. Samples were then dissolved in minimal chloroform, transferred to tared 20 mL scintillation vials, and pumped down to a fine, free flowing, black powder.

Characterization.
Powder X-ray diffraction (XRD) data was collected on a Bruker Microfocus instrument. Powder XRD samples were prepared by drop-casting a solution of Ni$_2$P onto silicon single crystal wafers. Diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) was collected on a Bruker Alpha IR instrument. Samples were prepared by grinding the Ni$_2$P together with FT-IR grade potassium bromide (KBr, >99%; Alfa Aesar) using an agate mortar and pestle. Background measurements were taken with a golden puck. $^1$H and $^{31}$P spectra were recorded on Bruker AV300 and Bruker AV500 spectrometers, respectively. Deuterated chloroform (CDCl$_3$) and acetonitrile (CD$_3$CN) were purchased from Cambridge isotope labs and dried over calcium hydride, vacuum-transferred, and stored over 4 Å sieves in a N$_2$ glovebox. CDCl$_3$ contained a 1% internal standard of tetramethyl silane. $^{31}$P spectra were corrected to an 85% phosphoric acid standard in water. X-ray photoelectron spectroscopy (XPS) was conducted on a Kratos AXIS Ultra DLD. Pass energy for survey spectra (to calculate composition) was 150 eV. Data point spacing was 1.0 eV per step for survey spectra, and 0.4 eV per step for detailed spectra. Pass energy for high-resolution spectra was 50 eV, with a data point spacing of 0.065 eV. All spectra were calibrated to align carbon peak intensities at 284.8 eV. Ultraviolet photoelectron spectroscopy (UPS) was also conducted on a Kratos AXIS Ultra DLD. The UV source was a He(I) lamp with photon energy of 21.2 eV. Pass energy was 5 eV with a step size of 0.01 eV. Aperture (spot) size was 100 μm. Take off angle was 0 degrees, and the stage was biased by a 9V battery while collecting data. All spectra were corrected to a sputter cleaned gold sample’s fermi edge at 0 eV. Transmission electron microscopy (TEM) images were collected on an FEI Tecnai G2 F20 microscope. Samples were drop cast from toluene onto lacey carbon grids and vacuum dried overnight. Scanning electron microscopy (SEM) images were collected on a ThermoFisher Scientific Apreo Variable Pressure SEM with integrated energy dispersive x-ray spectrometer (EDS) from Oxford. Elemental quantification was carried out using a Perkin Elmer Optima 8300 Inductively Coupled Plasma-Optical Emission Spectrophotometer (ICP-OES). Thermogravimetric Analysis (TGA) measurements experiments were carried out using a TA Instruments TGA Q5000. ~10 mg of material was loaded onto an aluminum pan at ambient conditions, then raised to 500 °C at a ramp rate of 10 °C/min under N$_2$ with a flow rate of 25 mL/min. All electrochemical measurements were performed with a Gamry Interface 1000. Measurements were conducted under Ar flow in 0.5M H$_2$SO$_4$ with 18 MΩ Millipore water in a custom four-neck cell fitted with a graphite rod counter electrode separated in a fritted compartment, a Ag/Ag$_2$SO$_4$ reference electrode separated by a Vycor frit, and a glassy carbon working electrode. A desired volume of a known mass% solution was drop cast onto the glassy carbon electrode to disperse the catalyst. Glassy carbon electrodes were consecutively polished with 5, 1, and 0.05 μm alumina paste and sonicated/washed with 18 MΩ Millipore water prior to use.

DFT Calculations.
Spin-polarized DFT calculations were carried out using the Quantum ESPRESSO\textsuperscript{81} package (v6.5). Exchange-correlation effects are described using the Perdew–Burke–Ernzerhof (PBE) functional\textsuperscript{82} along with the semi-empirical Grimme’s D2 van der Waals (vdW) corrections.\textsuperscript{83} The bonding environment of each system is described under the projector augmented wave method (PAW)\textsuperscript{84}. The expansion of valence wave functions has been accounted for with a plane wave cut-off energy of 680 eV. All simulations used a 3x3x1 Γ-centered Monkhorst-Pack\textsuperscript{85} k-point integration of the Brillouin zone. Surface reactions are modelled using an 8-layer periodically repeated hexagonal supercell (11.8 Å x 11.8 Å x 41.6 Å) belonging to the P6\textsubscript{2}m space group to model the coverage effect of surface adsorbates. A 25 Å vacuum space was employed to prevent spurious interactions between the periodically repeated images. Atomic coordinates were relaxed using the Broyden–Fletcher–Goldfarb–Shanno\textsuperscript{86} (BFGS) algorithm until the Hellmann-Feynman forces on all relaxed atoms fell below 0.05 eV/Å. For geometry optimization the electron occupancies were determined by the Gaussian smearing method with a smearing value of 0.07 eV, while for accurate total energy and partial density of states analysis we adopt the optimized tetrahedron method.\textsuperscript{87} Due to the relatively large system sizes in this study (144-204 atoms), only the top two layers are relaxed, with the bottom 6 layers remaining fixed to that of the bulk values. Preliminary analysis indicated that C\textsubscript{6}H\textsubscript{4}NO\textsubscript{2} preferentially binds to the P top site with high coordination Ni and P sites being disfavored in terms of total energy (See SI for more details). The Ni\textsubscript{2}P(0001) slab with the Ni\textsubscript{3}P\textsubscript{2} surface termination was chosen for three main reasons: (i) it closely resembles the experimental synthesis which has an observed stoichiometry of (Ni\textsubscript{1.78}P), (ii) it is known to be stable under the electrochemical conditions relevant in this study\textsuperscript{81} and (iii) it provides a point of reference with the current published literature.

Results and Discussion

Nickel phosphide synthesis and ligand stripping.

Nickel phosphide nanocrystals were prepared using a modified literature procedure.\textsuperscript{20} Briefly, nickel(II) chloride was mixed with tris(diethylamino)phosphine in neat oleylamine. After heating to 250 °C for 60 min, phase pure Ni\textsubscript{2}P nanocrystals with an average diameter of 5 nm (Fig. 1A) were recovered from a viscous black reaction mixture after successive rounds of precipitation, centrifugation at 7830 RPM, and redissolution with isopropanol/pentane followed by toluene/acetonitrile. TEM images reveal an approximately spherical morphology with an average diameter of 5 nm ± 0.9 nm (Fig. 1A). The powder XRD pattern is in agreement with the reported reference powder pattern (PDF 01-074-1385). (Fig. 1B). Scherrer analysis reflects an average crystalline domain size of 5.18 nm, consistent with the TEM analysis. ICP-OES reveals a composition that is phosphorus rich, with an average stoichiometry of Ni\textsubscript{1.78}P. This is a notable departure from the cation rich nature of more ionic nanomaterials synthesized with aminophosphine precursors, namely InP.\textsuperscript{88,89} We believe this composition is reflective of the inorganic nanocrystal core composition since there was no trace of excess molecular precursor or phosphonium salt byproduct found by NMR or FTIR spectroscopy of the purified end material (SI Fig. S1 and S4a).

Analysis by diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) and \textsuperscript{1}H NMR spectroscopy supports a ligand shell composed of oleylamine (SI Fig. S1). The IR spectra showed signatures of long-chain aliphatic hydrocarbons at 1,450, 2,850, and 2,920 cm\textsuperscript{-1}, as well as the expected alkene resonances at 1,600 and 3,000 cm\textsuperscript{-1}.\textsuperscript{90,91} \textsuperscript{1}H NMR spectra show resonances associated with aliphatic protons in the range of 1-2 ppm and the alkene protons at 5.3 ppm.\textsuperscript{92} Further, the broadness of these peaks is indicative of the long relaxation times expected for
molecules bound to a nanoparticle surface.\textsuperscript{65,93} Though previous reports of nanocrystals prepared using combinations of metal halide salts and aminophosphine precursors show evidence of a mixed ligand shell comprised of both amine and chloride ions, no evidence of Cl atoms are observed in the XPS survey spectrum (SI Fig. S2a).\textsuperscript{90} Further, thermogravimetric analysis suggests that the surface ligands compose 19\% of the sample by mass (SI Fig. S3). The first derivative of mass loss for this sample shows a peak at \textasciitilde350 °C, matching closely with the boiling point of neat oleylamine (365 °C).

\textbf{Figure 1.} A) General reaction scheme for the synthesis of Ni\textsubscript{2}P nanocrystals with aminophosphines. B) XRD of Ni\textsubscript{2}P nanocrystals references against powder pattern (PDF 01-074-1385). C) TEM of Ni\textsubscript{2}P product, showing 5 ± 0.9 nm spherical nanocrystals. Inset shows a single nanocrystal with a 5 nm scale bar for reference. Full image reflects a larger sample set of nanocrystals all reflecting similar sizes and morphologies with a 50 nm scale bar for reference.

High resolution XPS analysis of the Ni and P 2p electron binding energies are consistent with a covalent metal-phosphide material.\textsuperscript{38,28} The P 2p spectrum (SI Fig S2c) shows a spin-orbit doublet with the center of the parent peak located at 129 eV consistent with P in the 0 oxidation state and representing 72\% of the total P in the sample. A higher energy shoulder centered at 131.5 eV was fit to a broad doublet peak, representing what is likely multiple ill-defined higher oxidation state P components within the material. This species comes at a slightly lower binding energy than is expected of phosphate salts (132.5 eV), which leads us to believe it is likely a partially oxidized surface site.\textsuperscript{94} Similarly, the Ni 2p spectrum shows an unusual Ni binding energy of 853 eV, which lies in-between that of Ni\textsuperscript{0} and Ni\textsuperscript{2+} (852 eV and 854 eV, respectively).\textsuperscript{95} This result is indicative of the relatively high amount of electron density that remains on the Ni sites within the phosphide lattice. The 2p\textsubscript{3/2} environment can be deconvoluted into two contributions. The first being the metal-phosphide species centered at 853 eV which represents 72\% of the sample. The second, centered at 853.5 eV, accounts for 28\% of the sample and is likely the slightly reduced surface oxide, similar to that observed in the P 2p spectrum.

Prior to covalent functionalization of Ni\textsubscript{2}P, the concentration of native oleylamine ligands was reduced via alkylation using Meerwein’s salt. Alkylation is a widely used method for the removal
of common ligands including oleylamine, oleate, and phosphate species.\textsuperscript{96,97} Typically, ligand stripping procedures for colloidally dispersed nanoparticles consist of preparing a biphasic solution where the non-polar phase contains the native nanocrystals dissolved in hexanes and the polar phase is a solution of Meerwein’s salt in acetonitrile or DMF. Vortex mixing leads to rapid precipitation of the nanocrystals from solution. The recovered nanocrystals can be washed to remove residual aliphatic ligands before the particles are able to be resuspended by solvents with weak binding affinities such as DMF. In the most successful examples, complete loss of aliphatic C-H vibrations in FTIR spectra coupled with elemental analysis showing a lower concentration of C atoms are observed.\textsuperscript{96,98} Interestingly, this method of ligand stripping was not as efficient in the case of the oleylamine-ligated Ni$_2$P nanocrystals studied here. Following synthesis and purification, the Ni$_2$P nanocrystals showed only sparing solubility in hexanes, limiting our ability to directly use the procedure as previously reported. Additionally, vortex mixing the biphasic solution resulted in formation of a highly viscous gel. Ultimately, the ligand exchange was accomplished using a homogenous solution of the nanocrystals (5 mg/mL) in toluene with a large excess of Meerwein’s salt (50 mol equiv. vs oleylamine) in acetonitrile. This solution was mixed vigorously for several minutes in a round bottom flask and left to sit overnight, resulting in the formation of a black precipitate. The supernatant was decanted, and the resulting nanocrystal powder was dissolved in toluene and subjected to repeated rounds of precipitation and redissolution from acetonitrile.

The Meerwein’s salt-treated nanocrystals were characterized by TGA, IR, and NMR spectroscopy (SI Fig. S3-4). These data suggest that roughly 30\% of the initial oleylamine surface ligands were removed. However, oleylamine still accounted for 15\% of the sample by mass, and signature spectroscopic features were still observed in the DRIFTS and $^1$H NMR spectra. We do note that changes in the $^{31}$P NMR spectra suggest that alkylation is an effective method to remove the surface phosphate species that formed from ambient oxidation. Successive alkylation treatments had limited added effect and we have not managed to realize complete ligand removal in this system (SI Fig. S3), suggesting that the starting nanocrystals have a range of amine binding energies and that only the weakest bound subset can be removed using this method. This result stands in contrast to earlier examples of Meerwein’s salt treatment of nanocrystals for ligand stripping, a fact that may be attributable to differences in the covalent bonding character of the Ni$_2$P. Early examples from Helms and co-workers showcased the method’s efficacy on highly ionic materials such as CdSe, TiO$_2$, and ITO.\textsuperscript{96} Later studies, however, have shown less consistent results. In a study from Suntivich \textit{et al.} TGA suggested that alkylation did not remove all oleate ligands from the surface of Mn$_3$O$_4$ particles, despite the ionic nature of that lattice.\textsuperscript{97} In contrast, a recent paper from Buonsanti \textit{et al.} showed complete removal of native phosphate and tri-alkylamine ligands from the surface of Cu$^0$ nanospheres.\textsuperscript{98}

\textit{Covalent functionalization.}

To a chloroform solution of Meerwein’s salt-treated Ni$_2$P nanocrystals (Ni$_2$Ps) a solution of diazonium salt in acetonitrile was added dropwise (Fig. 2A). A spontaneous reaction was indicated by effervescence as N$_2$ gas was released. In some instances, a toluene solution of cobaltocene was added dropwise prior to addition of diazonium. In either case, the solution was capped loosely and left to stir for 48 hours. The functionalized nanocrystals were purified by first removing the solvent from the crude mixture under vacuum. Once the sample was dry, the powder was dispersed in acetonitrile by vigorous shaking or sonication, followed by multiple rounds of precipitation via centrifugation, removal of supernatant, and redissolution in acetonitrile to remove excess
diazonium salt. Subsequent washes with isopropanol and/or pentane were performed to remove any aliphatic reaction co-products. After multiple washes, the particles remained highly soluble in chloroform and/or toluene. Solvent removal under vacuum resulted in a free-flowing black powder.

We note that during initial washing steps the supernatant was observed to have a slightly yellow hue. Upon further investigation, we determined this side product was likely the result of reactivity between the diazonium ions and oleylamine. In fact, diazonium salts have been previously documented to act as ligand stripping reagents for colloidal quantum dots. Though the authors did not speak much to the mechanism, they showed the loss of prominent aliphatic vibrations after treatment and the growth of a BF$_4$ signal – suggesting some sort of ion exchange reaction. We propose that the mechanism is likely much more complicated due to the complex reactivity of highly electrophilic diazonium ions and nucleophilic primary amines.

$^{99}$ 1H NMR spectra of oleylamine, 4-nitrobenezene diazonium tetrafluoroborate, and an equimolar mixture of the two can be found in the supporting information (Fig. S6). Here we can observe the degradation of the clean aryl doublets (8.8 and 8.6 ppm) found in the parent diazonium into a convolution of multiple aryl species that are more electron rich, as suggested by their downfield shift (spanning from 7.6 to 8.3 ppm). We also observe that the reaction continues to evolve over the course of 2 days, by which point the original aryl resonances are no longer observed. We compare this control sample to the supernatant from our Ni$_2$P-C$_6$H$_4$NO$_2$ sample and observe many similar features, including the convoluted aryl signals and overwhelming aliphatic protons from the oleylamine.

A survey of early work studying the reactivity of diazonium ions lays out multiple paths of reactivity between these species, including nucleophilic attack at the beta-N, substitution at the C-2 and C-4 positions, heterolytic dediazoniations, and homomolecular coupling. This wide array of possible products is even further muddled by the influence of catalytic nanoparticles that provide competing bonding affinities, reducing equivalents, and catalytically active surfaces that can alter the reaction landscape. Further, the comparison of reaction supernatant and control samples suggest the reaction products are different, or at least are formed in very different ratios. However, none of these spectral features are observed in the purified products (Fig. S7) suggesting that these species can be completely removed from the system by thorough washing and do not result in persistent contamination.
Evidence of spontaneous functionalization of the Ni$_2$P nanocrystals was provided by TGA and DRIFTS of the purified and dried nanocrystal powders. The DRIFTS spectra reflected a mixed ligand shell, composed of remnant oleylamine and substituted aryl groups. Importantly, when comparing the spectra of functionalized nanocrystals to the parent diazonium salt (Fig. 2C and S8-10) we observed the absence of the N-N triple bond at ~2200 cm$^{-1}$. This is evidence that the nanocrystals were able to reduce the parent diazonium, cleaving the N$_2$-Ar bond. Further, this implies that the new spectral features observed in the fingerprint region are from surface bound functional groups rather than remnant/physiosorbed diazonium salt. Closer analysis of the features found in the fingerprint region show key vibrational modes associated with the specific functional groups expected in each sample, despite overlapping signal from oleylamine at ~1450 cm$^{-1}$. In the Ni$_2$P-C$_6$H$_4$OMe sample, the Ar-O stretch at 1291 cm$^{-1}$ as well as the O-CH$_3$ rocking mode at 1180 cm$^{-1}$ were observed. An additional peak at 1493 cm$^{-1}$ was attributed to aryl ring deformations induced by electron donating groups, thus only being observed in this sample. The peak observed from 1586-1590 cm$^{-1}$ in each sample is another characteristic aryl ring deformation attributed to C=C–C stretches. Characteristic symmetric and asymmetric N-O stretches were observed in the Ni$_2$P-C$_6$H$_4$NO$_2$ sample at 1513 and 1346 cm$^{-1}$, respectively. The in and out of plane NO scissor modes were difficult to resolve but could be observed at 856 and 724 cm$^{-1}$, respectively.

Interestingly these values were shifted to slightly lower frequencies than those of the parent diazonium salt, implying that the nanoparticle surface is acting as an electron donor, weakening the N-O bond. We note that all of the observed vibrational frequencies in our sample are in agreement with those previously reported in the functionalization of MoS$_2$ by aryl diazonium salts, implying the distortion of bond energies via electronic effects from the underlying material is a common theme.

TGA data (Fig. 2B) confirm a higher degree of functionalization with increasing Hammett parameter of the aryl substituent (NO$_2$>Cl>OMe), and hence increasing reactivity of the diazonium salt. This is consistent with previous observations in the literature, with the note that many papers use different methods of reporting the degree of functionalization depending on the system under investigation. For example, in studies of electrografting diazoniums onto glassy carbon surfaces authors report the magnitude of reductive current as a measure of functionalization yield. Another common method is reporting atomic ratios determined by XPS or S-Mo:S-C ratios in the case of MoS$_2$ functionalization. Though many of these studies supplement their findings with TGA and find the same result. We note, however, that the TGA data collected here is convoluted by the persistence of oleylamine in our sample, precluding direct correlation of the absolute mass loss values to covalently bound functional groups. In any case, we estimate the surface coverage of each sample with the assumption that 15% mass loss from oleylamine is constant across all samples, and that any further mass loss is from covalently bound surface functional groups. With these assumptions, we observed an upper bound for functionalization of 14% by mass for the nitrophenyl group. Assuming a 100 mg sample, this translates to 14 mg, or
114 μmol of nitrobenzene equivalents. In the same sample, 71 mg of Ni₂P would translate to 119 μmol of surface P sites, translating to 96% surface P sites being functionalized. Following this logic, methoxybenzene and 3,5-dichlorobenzene samples translate to 27 and 68 μmol, or 20% and 54% surface P-Ar sites, respectively (example calculation in SI). All functionalization densities were below 100% saturation and were less than the molar equivalents of parent diazonium added to the reaction mixture.

DFT calculations support the idea that adsorption of C₆H₄-NO₂, C₆H₃-Cl₂ and C₆H₄-OMe is spontaneous (Fig. 2D). Here, adsorbate-adsorbate and adsorbate-substrate interactions dictate the binding energy trends, with the electron donating/withdrawing character playing a key role. For instance, the adsorption energy of the electron withdrawing radicals (C₆H₃-Cl₂ and C₆H₄-NO₂) become less favorable as we increase the surface coverage. While the electron donating propensity of C₆H₄-OMe counteracts the effects of adsorbate-adsorbate interaction, highlighted by the near constant binding energy as a function of surface coverage. Similar trends in binding energy have been observed in the literature for black phosphorous and molybdenum disulfide. It should be noted that these calculations do not include thermal effects, which could significantly destabilize the adsorption of diazonium salts under reaction conditions. Other factors, such as, the presence of excess oleylamine capping ligands is shown to further destabilize the adsorption energetics (Fig. S23). That said, the low coverage trend of these species is consistent with the observed experimental coverages: 96% (C₆H₄-NO₂), 54% (C₆H₃-Cl₂) and 20% (C₆H₄-OMe).

To address the issue of unsaturated surface coverage, presumably due to the insufficient reducing potential or equivalents in the reaction mixture, the effect of adding a molecular reductant was investigated. This concept was recently validated by the Lewis group who showed that the functionalization density of MoS₂ by alkyl iodide species could be increased by adding sacrificial reductants to the reaction mixture. This study demonstrated a strong correlation between the strength of the reductant and the coverage of functional groups, supporting our hypothesis that each surface functional group equivalent becomes increasingly difficult to generate and is strongly dependent on the thermodynamic driving force for both generating the radical species and facilitating surface-C bond formation.

We experimentally validated this hypothesis in our own system by adding equimolar concentrations of cobaltocene (relative to diazonium) to our reaction mixtures. The resulting differences between batches of Ni₂P-C₆H₄NO₂ is shown in Fig. 3. The TGA data show a clear increase in mass loss and suggest that the organic content has increased to roughly 40% of the total sample mass. Following our earlier estimate of surface coverage, this would reflect a large excess (225% of surface P sites) of nitrobenzene groups. This super saturation can be rationalized by the formation of hyperbranched nitrophenyl oligomers on the nanocrystal surface, as shown in Fig. 3D. Uncontrolled reactivity, or hyperbranching, is a common phenomenon observed in diazonium chemistry. It is well known that aryl radicals react readily with sp² carbons, hence their usefulness in functionalizing graphitic carbons. As such, they are also able to react with the sp³ carbons present in surface arenes. This can lead to the formation of large polymeric films, sometimes on the order of tens of nanometers thick.
**Figure 3.** A) Reaction scheme showing functionalization of Ni$_2$P using [N$_2$-C$_6$H$_4$NO$_2$][BF$_4$] with the addition of cobaltocene as a sacrificial reductant. B) TGA of both samples, showing an increase in mass loss percent for samples made with added cobaltocene. Dotted lines show the first derivative of the mass loss percent line. C) TEM of Ni$_2$P-C$_6$H$_4$NO$_2$ nanocrystals synthesized with no added reductant. Particles are still uniform and maintain their morphological integrity. Scale bar is 10 nm. D) TEM of Ni$_2$P-C$_6$H$_4$NO$_2$ synthesized with added cobaltocene. Particles show aggregation induced by the extensive aryl-multilayer film formation caused by aryl radical hyperbranching. Scale bar is 10 nm.

**Effect of Functionalization on the Electronic Structure.**

The higher concentration of functional groups in the samples, derived from reactions in which a reductant was added, makes observation of functional groups with XPS easier. Spectra showing the presence of Cl and NO$_2$ functional groups can be found in the supporting information (Fig. S13). What’s more interesting, is the comparison of the Ni and P 2p high resolution scans between Ni$_2$P-C$_6$H$_4$NO$_2$ samples made with and without added reductant (Fig. 4A and 4B). Rather than distinct peaks indicating the formation of a P-Ar or Ni-Ar bonds, we observe systematic shifts of the entire ‘metal phosphide’ signal. Although this result is surprising, it can be interpreted as the functionalization perturbing core-level electron density throughout the entire nanoparticle, which is reflected in the electronic density of states (DOS) inferred from the DFT calculations (Fig. S25). For the bare surface, the partial DOS plot shows that Ni$_2$P exhibits metallic type conductivity with the states around the Fermi level consisting largely of Ni 3d-states with only a minor contribution from the overlapping P 3p-states. On adsorption of nitrobenzene groups, we observe a clear perturbation of these states and redistribution of electrons due to the electron withdrawing properties of the adsorbate. Despite our model only considering diazonium species adsorbed on the P-top sites, the change in total DOS can be attributed to a shift of both the P 3p-states and Ni 3d-states as well as a local increase in the P 3p-states. While we would typically expect this shift
in electron density to be localized and correlated with the functionalized surface sites, we propose the high level of covalency, and the metallic character of these metal phosphide nanocrystals allow localized perturbations to be compensated for by the ‘bulk’ material. Further, the same trend is reflected in the work function of these materials. The work function is made more positive with higher degrees of functionalization, which can be interpreted as the highly polar nitrophenyl functional groups inducing a stronger surface dipole moment away from the crystal’s core, in-turn requiring more energy to remove an electron from valence orbitals to vacuum.

Figure 4. A) High resolution XPS spectra showing the gradual progression of the Ni 2p3/2 signal shifting to higher binding energies as the coverage of nitrobenzene groups increases (blue > red). B) P 2p spectra of the same sample set, showing a similar systematic shift. C) UPS spectra of the same samples, showing the increased binding energy of the secondary electron cut off point, or work function. This illustrates increasing difficulty to eject an electron from the surface of the material as the coverage of nitrobenzene groups increases.

Although this phenomenon is unexpected when compared to studies of exfoliated MoS2 and black phosphorous nanosheets, it is reminiscent of the effects observed when doping metal phosphides with metals of varying electronegativity.111 In this study, Mar and coworkers observed a systematic shift in the P and Ni 2p3/2 binding energy over the range of ~500 meV. The authors find that as more electropositive ions (Co<Fe<Mn<Cr) are doped into the lattice the P 2p3/2 binding energy systematically decreases. They rationalize this finding by considering a charge potential model to balance the inter and intraatomic effects of dopants on the P atoms in the lattice.112 Based on this model, intraatomic effects would lead to a decrease in binding energy due to increased nuclear screening and greater negative charge of P core electrons induced by their more electropositive environment. In contrast, interatomic effects would lead to a positive shift in binding energy due to the general increase in Madelung potential felt by each atom in the more electropositive lattice. We believe this rationale supports our interpretation of highly coupled covalent surface functional groups being able to perturb the ‘bulk’ electronic structure of the nanocrystal by inducing strong interatomic screening effects. The result of which is a systematic perturbation of the core-level binding energies in both Ni and P based on the degree of electron density that is donated or withdrawn by the surface functional group.

We further validate this hypothesis by showing systematic variation of core-level binding energies and NC work function across our Hammett series (Fig. 5). There is a clear linear trend observed in both Ni and P 2p high-resolution spectra where a larger Hammett parameter increases the core-level binding energy. This effect is most pronounced in the comparison of work functions.
across the Hammett series shown in Fig. 5C. To ensure this wasn’t purely an electrostatic screening effect, we validated our hypothesis against a computational model where the same trend is observed (Fig. S24E).

Figure 5. A) High resolution XPS spectra showing the shift of Ni 2p3/2 signal in response to the Hammett parameter of the surface functional group. B) P 2p spectra of the same sample set, showing a similar systematic shift. C) UPS spectra of the same samples, showing the response of material work function to the Hammett parameter of surface functional group. Inlet shows the linear correlation between these descriptors.

**Electrocatalytic HER Activity.**

To test the effect that surface functionalization has on the electrocatalytic activity for HER of the Ni2P nanocrystals, 20 μL of a 10 mg/mL solution of Ni2P nanocrystals in toluene were drop casted onto a 1.5 mm diameter glassy carbon electrode. Samples were prepared in a glovebox and transferred to a vessel of 0.5 M H2SO4 that was sparged with argon and held under a constant argon flow. Here we report the current density normalized by the geometric surface area of the glassy carbon electrode due to inconsistencies in estimating the electrochemically active surface area of the nanocrystal films (Figs. S14-S18). Metal phosphides are known to have exceptionally high specific capacitance values due to pseudocapacitive effects, which typically leads to over estimation of electrochemically active area. These pseudocapacitive properties would be affected by alteration of surface charge induced by the surface functional groups. This added complication makes it difficult to uniformly attribute “capacitive” current to strictly ion adsorption on a metallic surface, which is the base assumption in the estimation of electrochemically active surface area. This is reflected in our inability to find a potential region that reflects a purely capacitive response characterized by a square-wave with equal anodic and cathodic current. Instead, we observe variable cathodic current around the open-circuit potential with added resistive losses, potentially due to mass-transport limitation through the nanocrystal film (Figs. S15-S18). Though this may preclude us from reporting the true intrinsic activity of our catalysts, we can still compare relative activity within the series.

The catalytic activity results are summarized in Fig. 6 and reflect multiple key results. First, we observe over the course of 24 hours and 75 polarization cycles that the catalytic activity does not track the Hammett parameter, instead Ni2P-C6H4OMe is the most active catalyst followed by Ni2P-C6H4NO2 and then Ni2P-C6H3Cl2 being the least active. This contrasts with previous studies by Miller and co-workers on covalently functionalized MoS2, where they found that catalytic activity was correlated with the functional group’s Hammett parameter, although the nitrobenzene
functionalized sample was not measured in that instance.\textsuperscript{61} Comparisons between these systems brings us to our second key result, which is improved catalytic activity in all functionalized samples when compared to Ni$_2$P in the first 20 polarization cycles. We propose these discrepancies reflect that the influence of covalently tethered ligands is more nuanced than simple electronic effects and that their impact may vary depending on the nature of the catalyst’s surface chemistry and the density of grafted aryl groups. For example, the documented active site for MoS$_2$ is the edge atoms of the 2D sheet while the basal plane surface is relatively inert for HER. Thus, it is possible that functional groups that bond at the basal plane surface do not have added steric effects because of spatial separation from the catalytically active site. As we show below, that is not the case in Ni$_2$P nanocrystals, for which the aryl functional groups bond to atoms directly adjacent to active sites. This could allow for the same functional group to result in different effects based on the intrinsic reactivity and structure of the underlying material.

**Figure 6.** (a) LSV of the 10\textsuperscript{th} polarization cycle for Ni$_2$P and Hammett series electrodes collected at 5 mVs\textsuperscript{-1}. (b) Overpotential at 10 mA/cm\textsuperscript{2} averaged across three separate electrodes for each sample over the course of 24 hours and 75 polarization cycles. (c) Calculated free energy profile at 0.00 V, pH = 0 and 300 K for the adsorption of H onto the Ni$_2$P(0001) Ni$_3$ hollow site, with and without surface functional groups.

For efficient hydrogen evolution, the hydrogen adsorption free energy should be slightly negative, so that there is a driving force for hydrogen to adsorb onto the catalyst surface, but the adsorption energy is low enough to allow for the subsequent proton coupling steps. As can be seen from Fig. 6C, DFT calculations predict that the free energy for hydrogen adsorption at the Ni$_2$P(0001) Ni 3-fold hollow site (possible surface binding sites are shown in Fig. S21) can be modified substantially with surface functionalization of diazonium salts. The trend follows: Ni$_2$P(-0.47 eV) > Ni$_2$P-C$_6$H$_3$Cl$_3$(-0.26 eV) > Ni$_2$P-C$_6$H$_4$NO$_2$ (-0.22 eV) > Ni$_2$P-C$_6$H$_4$OMe(-0.14 eV), which is in agreement with our experimental findings. In turn, this suggests that the adsorption energies of H on the nanocrystal surface do not directly follow the electrostatic trend.
Figure 7. Charge density difference plots highlighting the interactions of key species at the Ni$_2$P(0001) interface. (a) H adsorbed at the Ni$_3$ hollow site, (b) C$_6$H$_4$-NO$_2$, (c) C$_6$H$_3$-Cl$_2$ and (d) C$_6$H$_4$-OMe adsorbed on the P top sites. Red and blue shading represent electron depletion and electron accumulation respectively. The iso-surface level is set to 0.013 e/Å$^3$.

To further probe the origins of the observed trend, we examined the effects of electron transfer between the nanocrystal atoms and surface adsorbates by calculating their charge density difference (CDD) (Fig. 7). Fig. 7A shows, that once adsorbed on the surface, hydrogen accumulates electron density from nearby Ni atoms, forming a surface hydride. Interestingly, from Figs. 7B to 7D, adsorption of diazonium salts on P top sites exhibit non-local charge redistribution at the Ni$_3$ hollow site. The net effect of functionalization is the tuning of the hydrogen adsorption free energy at the Ni$_3$ hollow site via electron redistribution resulting in a weaker adsorption energy for H atoms, thus making the catalytic landscape more thermoneutral relative to the bare surface. These local fluctuations in Ni charge density can facilitate H atom diffusion onto the Ni-Ni bridge site (Figs. S27 to S28). The Ni-Ni bridge site is expected to be a catalytically active site in computational studies but is generally only accessed at higher H coverages.$^{40,117}$ We further note that the transition from a surface hollow site to lower coordination sites would be hindered by the Pauli repulsion exhibited by the Cl atoms of the C$_6$H$_3$-Cl$_2$ functional groups and the hydride nature of adsorbed H (H$^\delta$-Cl$^\delta$). Additionally, this charge redistribution depends strongly on the alignment of diazonium species above the P top site. Most noticeable is the repulsive interactions exhibited by C$_6$H$_3$Cl$_2$ (Fig. 7C), which is reflected in the binding energy discussed previously. The Cl$^\delta$-Cl$^\delta$ repulsion results in a staggered array of C$_6$H$_3$Cl$_2$ species and a disfavored alignment of the C p$_z$-orbitals with P 3p-states. To compensate for this, some P atoms are pulled away from the surface plane, resulting in significant distortion and a weaker interaction with the surface atoms. The trend in P-atom distortions C$_6$H$_3$-Cl$_2$(0.57 Å) > C$_6$H$_4$-NO$_2$(0.31 Å) > C$_6$H$_4$-OMe(0.29 Å) suggests that para substitution minimizes adsorbate-adsorbate interactions leading to an optimal surface packing of diazonium salts relative to the meta substituted functional groups. Overall, we find that the local electronic structure and bonding interactions of the diazonium salts influence the charge density across the surface, which we hypothesize being the cause of our observed trend.
Electrocatalytic Stability.

The next key finding is the variation of activity over multiple polarization cycles. The first and most important point is that the trend in activity induced by the functional groups is held over 24 hours of exposure to catalytic conditions. We propose this is strong evidence in favor of covalent functional groups being a viable pathway to stable hybrid nanomaterials. Another interesting point is the variability, or lack thereof, in activity over time. We observed a gradual increase in activity after successive polarizations for the ligand stripped Ni$_2$P sample, as shown in Fig. 6B. This is presumably due to the removal of remnant oleylamine from the surface, allowing for a better wetting of the electrolyte as previously demonstrated in our lab.\textsuperscript{56} However, we cannot count out the possibility of surface reconstruction, which has been theoretically predicted and experimentally measured to take place during catalysis.\textsuperscript{40,42,118–120} If, after prolonged cycling, the dominant surface facet is no longer the Ni$_3$P$_2$ (0001) termination it could account for the gradual disagreement with the predicted thermodynamic landscape that we observe in earlier cycles (Fig. 6C). Further, this swing in activity is quite large, showing an improvement of 150 mV over the course of our measurements. This contrasts with the functionalized samples which show very little variability under the same conditions, reflecting a maximum average deviation of only 25 mV. We note that although the activity of the Ni$_2$P electrodes did improve over time, they never achieved higher current densities or lower overpotential than the Ni$_2$P-C$_6$H$_4$OMe sample, even in the few samples we measured out to 100 polarization cycles (Fig. S19, S20). Another practically relevant observation during our analysis was the I-V response during cycling. Consistently, we observed wild variations in the current response from Ni$_2$P electrodes, presumably due to the formation of hydrogen bubbles. This is reflected in the noisy LSV reported in Fig. 6A but becomes even more apparent at high current densities (Fig. S20). Finally, we were able to observe the retention of key atomic handles by SEM-EDS after repeated cycling, presented in Fig. 8. Critically, this illustrates that the chlorine atoms from the 3,5-dichlorobenzene functional group remain observable and are specifically observed in regions of the image where Ni and P are also detected, rather than randomly scattered across the sample.

Figure 8. SEM-EDS map of Ni$_2$P-C$_6$H$_3$Cl$_2$ electrode after 100 polarization cycles showing the presence and spatial overlap of P (a), Ni (b), and Cl (c) on the electrode after cycling.
Conclusions and Outlook

In conclusion, we have formulated synthetic conditions for generating (sub)monolayer coverages of substituted aryl functional groups covalently bound to the surface of Ni2P nanocrystals and measured the resultant effects on the nanomaterial’s electronic and electrocatalytic properties.

Through our study we have highlighted the need for further investigation into soft ligand removal strategies for colloidal metal phosphide nanocrystals. We have shown that native ligands can participate in side reactions with functionalization precursors and convolute characterization. Still, we show that Ni2P nanocrystals can be spontaneously functionalized by substituted aryl diazonium salts. Though we did not observe explicit experimental evidence, computation modeling suggests this functionalization occurs via formation of covalent P-C bonds. The degree of functionalization is dictated by the electron density in the aryl ring, where substituents that withdraw more electron density result in higher functionalization densities. We find that the electron donating/withdrawing nature of the surface functional group systematically alters core-level electron density and the bulk Fermi-level of the materials. Interestingly, we find that the influence on electrocatalytic activity is improved by the addition of substituted aryl groups, but the trend does not clearly track the electrostatic influence of the substituent functional groups. Though unexpected, these findings are validated by DFT modeling of our system which identifies the same broken trend. Through this model we propose that the functional groups have a nuanced effect on activity, which go beyond simple surface dipole effects, and can alter the binding energetics at active sites through surface deformation and steric repulsion. Further, we find that this trend in altered electrocatalytic activity for the HER is maintained over the course of 24 hours and more than 75 polarization cycles, signaling the promise for covalently bound ligands to be robust to catalytic conditions.

Though sterically rigid aryl functional groups are likely not the key ingredient for making an earth abundant catalyst rival the activity of platinum, the utilization of this well-studied class of surface functionalization agents is a promising starting point. We propose that building models and further refining this relatively simple system will lay the groundwork for the development of synthetic procedures to produce elegant systems of covalently tethered proton relay ligands that fully utilize secondary and outer coordination sphere effects. However, there is still much work to be done as a community before such a system can be conceived. For example, exploring the range of electro/nucleophilicities required to form bonds with surface atoms so we can move away from harsh and difficult-to-control radical chemistry. These ideas lead to a further question: would we install an already synthesized complex ligand on the surface, or build it from the bottom-up using post-synthetic transformations once a suitable reactive fragment is bound to the particle surface? Surendranath and co-workers have covalently tethered organometallic catalysts to the edges and defect sites of carbon materials.121,122 In those systems, the authors were able to leverage native o-quinone moieties that irreversibly condense with diamines to form their covalent linkages. Though such native defects are not as well documented in nanocrystalline phosphide or other more complex materials, a better understanding of the chemical nature of these materials’ surface atoms should allow for the development of more elegant functionalization strategies. In fact, collaborative studies between our group and the Mayer group have taken aim at building a more molecular understanding of colloidal metal phosphide nanocrystals.123 This concept and line of study will directly bolster our on-going efforts to refine surface functionalization procedures and create better defined hybrid nanomaterials.
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