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pH-dependent cross-linking of catechols through oxidation via Fe³⁺ and potential implications for mussel adhesion

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Abstract

The mussel byssus is a remarkable attachment structure that is formed by injection molding and rapid in-situ hardening of concentrated solutions of proteins enriched in the catecholic amino acid 3,4-dihydroxy-L-phenylalanine (DOPA). Fe³⁺, found in high concentrations in the byssus, has been speculated to participate in redox reactions with DOPA that lead to protein polymerization, however direct evidence to support this hypothesis has been lacking. Using small molecule catechols, DOPA-containing peptides, and native mussel foot proteins, we report the first direct observation of catechol oxidation and polymerization accompanied by reduction of Fe³⁺ to Fe²⁺. In the case of the small molecule catechol, we identified two dominant dimer species and characterized their connectivities by nuclear magnetic resonance (NMR), with the C6-C6 and C5-C6 linked species as the major and minor products, respectively. For the DOPA-containing peptide, we studied the pH dependence of the reaction and demonstrated that catechol polymerization occurs readily at low pH, but is increasingly diminished in favor of metal-catechol

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Supporting Information. Supplementary methods for the oxidation of 4MC and characterization of products by MS and synthesis of Ac-Ser-DOPA-NH₂ are available. Figure S1, DHPA oxidation of by Fe³⁺ digital images; Figure S2, MS DHPA oxidation; Figure S3, MS 4MC oxidation; Figure S4, HPLC-MS with EICs of DHPA oxidation; Figure S5, labeled DHPA dimer structures; Figure S6, Dimer 1 ¹H NMR; Figure S7, Dimer 1 ¹³C NMR; Figure S8, Dimer 1 ¹H-¹H COSY; Figure S9, Dimer 1 ¹H-¹³C HSQC; Figure S10, Dimer 1 ¹H-¹³C HMBC; Figure S11, Dimer 1 2D-J NMR; Figure S12, Dimer 2 ¹H NMR; Figure S13, Dimer 2 ¹H-¹H COSY; Figure S14, Dimer 2 ¹H-¹³C HSQC; Figure S15, Dimer 2 ¹H-¹³C HMBC; Figure S16, Dimer 2 2D-J NMR; Figure S17, UV-Vis demonstrating Fe³⁺ reduction to Fe²⁺ with phenanthroline; Figure S18, HPLC-MS with EICs of dipeptide oxidation. Table S1, Relative abundance of DHPA, *o*-quinone, and dimers under different reaction conditions.

coordination interactions at higher pH. Finally, we demonstrate that Fe^{3+} can induce cross-links in native byssal mussel proteins *mefp-1* and *mcfp-1* at acidic pH. Based on these findings, we discuss the potential implications to the chemistry of mussel adhesion.

Introduction

Mussels attach to underwater surfaces via a cluster of threads (the mussel byssus) that adhere strongly to solid surfaces via a terminal adhesive plaque (Figure 1a). Each thread of the mussel byssus is individually formed with an adhesive plaque in a fascinating manner akin to injection molding. Liquid protein secretions are extruded into the ventral groove and distal depression of the mussel foot, whereupon they solidify in only a few minutes to yield a newly formed byssal thread that, along with other threads, functions to tether the organism to the surface. Using this process, mussels are able to strongly bind to organic and inorganic surfaces in aqueous environments where many other glues fail.¹⁻⁵

Previous studies revealed that 3,4-dihydroxy-L-phenylalanine (DOPA), an amino acid formed by post-translational modification of tyrosine, is distributed throughout the byssus with especially high concentrations in proteins located near the plaque-substrate interface.⁶⁻⁹ Although a comprehensive understanding of the role of DOPA in mussel adhesion has not been reached, existing experimental evidence points to both adhesive and cohesive functions. Strong interfacial interactions between DOPA or oxidized DOPA residues and surfaces likely contribute to adhesion between the mussel adhesive plaque and substrate,¹⁻⁵ whereas, within the bulk of the adhesive plaque, DOPA-DOPA polymerization is thought to be a source of cross-linking that ultimately leads to solidification into a cohesive elastic solid.¹⁰⁻¹²

The concentration of Fe^{3+} in the mussel byssus has been measured at greater than 1 part per thousand - many orders of magnitude greater than the concentration of Fe^{3+} in seawater (typically 10 parts per billion).¹³⁻¹⁶ Through pulse ^{59}Fe experiments, ^{59}Fe has been found to be taken up by the mussel and redistribute into byssal thread.¹⁷ Fe^{3+} has been shown to form strong coordination bonds with catechols, in which one, two, or three catecholates can bind a single ferric ion.^{13, 18-21} Such noncovalent cross-links may result in a network of coordination bonds serving a mechanical role within adhesive plaques and byssal threads,²² as demonstrated recently using the surface forces apparatus and in biomimetic catechol polymer gels.^{4, 5, 23-27}

Aside from these coordination interactions and literature reports of catechol oxidation to *o*-quinone by Fe^{3+} ,²⁸⁻³⁰ many aspects of catechol- Fe^{3+} interactions in the mussel byssal protein system remain unclear. Native oxide surfaces of steel have been reported to cure DOPA-containing polypeptides, which the authors suggest is due to the presence of soluble Fe^{3+} leading to the oxidation of DOPA.³¹ We recently demonstrated that catechol-modified poly(ethylene glycol) polymers could be covalently cross-linked at acidic pH in the presence of Fe^{3+} .²⁴ Others have speculated that Fe^{3+} is involved in redox-mediated covalent curing of mussel adhesives.^{15, 32-34} Though this type of metal-mediated dimerization has been known for decades to occur with phenols and naphthols,^{35, 36} to our knowledge, no direct evidence

of similar reactions with catechols exists to support the hypothesis that Fe^{3+} could covalently cure mussel adhesives.

In an attempt to elucidate more details of Fe^{3+} -catechol interactions relevant to mussel adhesion, we studied the reaction of small molecule catechols and DOPA-containing peptides (Figure 1b) with Fe^{3+} using spectroscopic and chromatographic methods that allow for conclusive detection of covalent catechol cross-linking, including the pH dependence of such reactions. In our initial studies, monofunctional catechols were selected over native mussel proteins and multivalent catechol polymers in order to more precisely characterize the reaction products.¹¹ The reaction between a catechol-containing small molecule mimic of DOPA, 3-(3,4-dihydroxyphenyl)propionic acid (DHPA, **1**, Figure 1c), and Fe^{3+} was studied as a function of time and metal:catechol stoichiometry by high performance liquid chromatography (HPLC), mass spectrometry (MS), and ultraviolet-visible spectroscopy (UV-Vis). At acidic pH, DHPA and Fe^{3+} mixtures formed catechol-catechol dimers within 1 min and higher oligomers at longer reaction times, along with the reduction of Fe^{3+} to Fe^{2+} . The carbon-carbon connectivities of the main dimer reaction products were determined by nuclear magnetic resonance spectroscopy (NMR). The pH dependence of Fe^{3+} -DOPA interactions was assessed with a DOPA-containing peptide (Ac-Ser-DOPA- NH_2 , **2**, Figure 1c). Covalent cross-linking was demonstrated to occur at acidic pH but not at alkaline marine pH in the presence of Fe^{3+} . Lastly, Fe^{3+} mediated cross-linking of the DOPA-rich mussel foot proteins *mefp-1* and *mcfp-1* was demonstrated at acidic pH.

Experimental Section

Materials

Acetonitrile, DHPA, ferric chloride hexahydrate, ferrous chloride tetrahydrate, formic acid, 1,10-phenanthroline hydrochloride monohydrate (phenanthroline), glacial acetic acid, sodium acetate trihydrate, bis-tris, bis-tris HCl, bicine, and ethylenediaminetetraacetic acid (EDTA) were purchased from Sigma Aldrich (Milwaukee, WI). All solutions of FeCl_2 were made fresh so that spontaneous oxidation of Fe^{2+} to Fe^{3+} was minimized. All chemicals were used without further purification.

Oxidation of DHPA by Fe^{3+}

Stock solutions of DHPA (75 mM) and FeCl_3 (100 mM, 200 mM, 300 mM) were prepared in Nanopure H_2O . Reactions were initiated by combining 800 μL of the DHPA stock solution and 200 μL of the appropriate FeCl_3 solution. Fe^{3+} :DHPA ratios of 1:3, 2:3, and 3:3 were accomplished by adding 200 μL of 100, 200, and 300 mM FeCl_3 , respectively. For reactions with Fe^{3+} :DHPA ratios of 4:3 and 6:3, 800 μL of the DHPA stock solution was mixed with 400 μL of 200 mM FeCl_3 and 300 mM FeCl_3 , respectively.

HPLC of small molecule reactions

The reaction products were separated by reversed-phase HPLC (Waters; Milford, MA) with a diphenyl column (Grace, Vydac # 219TP54). Separations were performed isocratically with a mobile phase of 90 % water with 0.1 % formic acid and 10 % acetonitrile and a flow rate of 1.0 mL/min. UV detection was performed at 220 nm and 280 nm.

MS of DHPA reaction

The reaction solutions were analyzed at Northwestern University's Integrated Molecular Structure Education and Research Center (IMSERC) on an Agilent 1100 LC/MSD high performance ion trap mass spectrometer by direct injection in negative ion mode with water as the mobile phase.

HPLC-MS of DHPA reactions

The reaction solutions described above were diluted tenfold and the reaction products were separated by reversed-phase HPLC at IMSERC on an Agilent 1100 LC/MSD high performance ion trap mass spectrometer in negative ion mode equipped with a diphenyl column (Grace, Vydac # 219TP54). The isocratic solvent system used for separations was 90 % water with 0.1 % formic acid and 10 % acetonitrile at a flow rate of 1.0 mL/min. UV detection was performed at 220 nm and 280 nm.

NMR characterization of dimer products

HPLC was performed as described above, except that the solvent system was composed of 90 % water with 0.1 % trifluoroacetic acid (TFA) and 10 % acetonitrile with 0.1 % TFA. The products were collected, frozen, lyophilized, and dissolved in D₂O. NMR spectra were acquired at IMSERC at 25 °C. 1H 1D and 2D-J-resolved spectra were acquired on a 500 MHz Varian Inova spectrometer equipped with a 5 mm Nalorac ¹³C{¹H} dual broadband probe bearing z-axis pulsed field gradients. Data were acquired using VNMRJ 2.2D software. The 2D-J experiment employed Varian's pulse sequence "hom2dj.c". ¹H, ¹³C, ¹H-¹³C HSQC and HMBC spectra were acquired in automation on a 500 MHz Bruker Avance-III spectrometer equipped with a 5 mm ¹³C-detect DCH cryoprobe and Z-axis pulsed field gradients. The sample of Dimer 2 (the minor dimer product) was too dilute to obtain a ¹³C spectrum, but ¹³C resonance assignments were made using 2D HSQC and HMBC spectra.

UV-Vis characterization of Fe³⁺ reduction

Spectra of the following solutions were acquired on an Agilent HP8452 spectrometer at Northwestern University's Keck Biophysics Facility: phenanthroline (20 μM); DHPA (30 μM) + phenanthroline (20 μM); FeCl₃ (20 μM) + phenanthroline (20 μM); FeCl₂ (20 μM) + phenanthroline (20 μM); DHPA (30 μM) + FeCl₃ (20 μM) + phenanthroline (20 μM); DHPA (30 μM) + NaIO₄ (15 μM) + phenanthroline (20 μM).

Synthesis of DOPA-containing dipeptide

Ac-Ser-DOPA-NH₂ was prepared by standard Fmoc solid phase peptide synthesis methods (Supplementary Methods).³⁷

pH-dependent oxidation of DOPA-containing dipeptide by Fe³⁺

Ac-Ser-DOPA-NH₂ was dissolved in water or buffer. For 1× FeCl₃ samples, 1× FeCl₃ was added to the sample from a 100 mM FeCl₃ stock within 1 min of dissolving the dipeptide in buffer. The final dipeptide concentration was 50 mM, and the final buffer concentration was 250 mM for all reactions. Reactions were studied at pH 5, 7, and 9, using acetate, bis-tris,

and bicine buffers for each pH, respectively. After 1 h, 10× EDTA was added from a 0.100 M EDTA stock and the final peptide concentration was diluted to 1 mg/mL with water.

HPLC-MS of dipeptide reactions

The reaction solution from dipeptide reactions (10 μ L) were injected onto an Agilent 6210 LC-TOF operated in negative ion mode and equipped with a Waters dC18 Atlantis column (#186001346) at a flow rate of 1.0 mL/min and column temperature of 45 °C at IMSERC. Mobile phases used were A, water with 0.1% formic acid, and B, acetonitrile with 0.1 % formic acid. A linear gradient method was employed: 0–5 min, 3 % B isocratic; 5–20 min, 3–50 % B; 20–25 min, 50–90 % B; 25–30 min, 90 % B. UV absorbance was collected at 280 and 220 nm. Approximately 50 % of the eluent was diverted after the column and UV detector, before reaching the MS. Extracted ion chromatograms were used to identify corresponding HPLC peaks.

Purification of *mfp-1*

Mfp-1 from *Mytilus californianus* (*mcfp-1*) was purified from mussels harvested off Goleta Pier, (Santa Barbara, CA), and held in circulation tanks before shucking the mussels and extracting the foot. *Mfp-1* from *Mytilus edulis* (*mefp-1*) was purified from mussels feet supplied by Northeast Transport (Waldoboro, ME). Both proteins were purified as described previously,³⁸ with some modifications. Briefly, the feet were frozen to –70 °C before flaying away the pigmented epithelium. Approximately 50 prepared feet were homogenized in four equivalents (w/v) of 5% acetic acid (v/v) with 10 μ M leupeptin, 10 μ M pepstatin, and 1 mM EDTA in a glass Kontes tissue grinder (Vineland, NJ) on ice and centrifuged at 20,000 \times g, 4 °C for 40 min. The supernatant was acidified with 70 % perchloric acid to a final concentration of 1.5 % (v/v). After centrifugation at 20,000 \times g, 4 °C for 40 min, the supernatant was dialyzed 4 \times 4 L of 5 % acetic acid (v/v) for 4 h in 1,000 kDa molecular weight cutoff dialysis tubing (Spectrum Industries, Los Angeles, CA) before freeze drying. The lyophilized protein was resuspended in 200 μ L of 5 % acetic acid (v/v) and 50 μ L aliquots were run over a size exclusion column (Shodex KW-803 5 μ m, 8 \times 300 mm, New York, NY). The eluent was monitored at 280 nm and those fractions positive for protein were subjected to acid-urea polyacrylamide gel electrophoresis (7.5% acrylamide and 0.2% *N,N*-methylenebisacrylamide) containing 5% acetic acid and 8 M urea.³⁹ After electrophoresis, gels were stained with Sigma-Aldrich Coomassie Blue R-250 (Brooklyn, NY). Pure *mfp-1* fractions were pooled and aliquoted before freeze-drying and stored at –70 °C for future use.

Fe³⁺-mediated protein cross-linking

Protein (*mefp-1* or *mcfp-1*) was resuspended to 50 μ M in 0.1 M formic acid buffer (pH 3.0) (Fisher), mixed with ferric chloride adjusted to three different concentrations (0, 0.005, 0.05, 0.5 μ M), and incubated for 5 min at room temperature. Solutions were diluted (1/5) with 0.1 M formic acid pH 3.0 with a small amount of tracking dye (0.3 μ M methyl green, 5% acetic acid, 8M urea) and loaded onto an acid-urea polyacrylamide gel and ran for 1–2 h at 110 V and stained with Coomassie blue.

Results

Fe³⁺-induced cross-linking of a DOPA mimic

DHPA (Figure 1c, 1) was chosen as the catechol-containing small molecule because the presence of the carboxylic acid allowed the products to remain soluble throughout the reaction (other catechols studied formed precipitates). Initial reactions were carried out in unbuffered water, which was acidic (pH ~2) due to the presence of FeCl₃. Upon mixing Fe³⁺ (orange) and DHPA (pale yellow) solutions, the mixture immediately turned deep green (Figure S1), a color associated previously with a noncovalent coordination interaction between one Fe³⁺ ion and one catechol.^{18, 28} As time progressed, the solution transitioned to yellow (~1 h) and brown (~2 d), reminiscent of catechol-quinone tanning.⁴⁰ Ion trap mass spectrometric analysis of the reactant solutions revealed a mixture of unreacted DHPA, dimers and higher oligomers (Figure 2 and Figure S2). Several Fe³⁺:DHPA ratios ranging from iron-rich to iron-deficient were examined (Table 1) in order to determine the dependence of the reaction on reactant stoichiometry. At excess DHPA (Reaction 1, Fe³⁺:DHPA of 1:3), a large portion of DHPA remained, with limited dimer evident in the mass spectrum (Figure S2). With increasing Fe³⁺, the amount of unreacted DHPA decreased and the amount of higher mass products increased. At excess Fe³⁺ (Reaction 5, Fe³⁺:DHPA of 6:3), only a small portion of DHPA remained (Figure 2), in favor of dimers and trace quantities of trimers and tetramers. Similar results were obtained when the reaction was performed with 4-methylcatechol (4MC) (Supplementary Methods and Figure S3), indicating that the carboxylic acid of DHPA did not significantly influence the reaction.

In order to gain a more complete understanding of the reaction between Fe³⁺ and DHPA, HPLC was used to separate the products in the reaction mixture. Due to the aromatic character of DHPA and its dimers, a diphenyl column provided maximal separation of the reaction components. Initially, HPLC-MS was performed in order to identify the molecule associated with each HPLC peak (Figure S4). Two aspects of the HPLC-MS data were extremely interesting: the *o*-quinone form of DHPA (*m/z* = 179) could be specifically identified, and two different peaks associated with the dimer (*m/z* = 361) were detected, each separated by 3.5 min.

Using the assignments from HPLC-MS, the effect of Fe³⁺:DHPA on the reaction products was further studied (Figure 3a). By determining the area under the DHPA peak relative to the total area under the peaks of DHPA and dimers, the amount of DHPA consumed in the reaction could be estimated (Table 1). At low Fe³⁺:DHPA (1:3), over 77 % of the DHPA remained unreacted. However, the amount of unreacted DHPA decreased to 57 %, 43 %, 29 %, and 12 % at Fe³⁺:DHPA of 2:3, 3:3, 4:3, and 6:3, respectively. An interesting trend was noted in the relative abundance of the two dimer peaks appearing at 7.5 and 10.5 min (Table S1). The ratio of the peak areas of major (7.5 min; Dimer 1) to minor (10.5 min; Dimer 2) dimer products increased from 2.5 at Fe³⁺:DHPA of 1:3 to 5.8 at Fe³⁺:DHPA of 6:3. Thus, increasing the amount of Fe³⁺ relative to DHPA favors production of Dimer 1.

The time dependence of the reaction between Fe³⁺ and DHPA was also studied (Figure 3b). Using Fe³⁺:DHPA equal to 3:3 as a representative reaction condition, the reaction mixture was analyzed by HPLC after 1 min, 37 min, 189 min, 618 min, and 24 h. A general

observation about this chemical transformation is that a significant amount of DHPA is consumed within the first minute, yielding dimer products (Figure 3b and Table S1). Between the two forms of dimers, Dimer 1 was the dominant species at all time points and was roughly 3.2–3.6 times more abundant than Dimer 2. An interesting feature of the time-dependent HPLC is the transient presence of the *o*-quinone form of DHPA. This species was observed after 1 min of reaction time, peaked at 37 min, decreased significantly thereafter, and was not observed at 7 days.

Structural determination of DHPA dimers by NMR

Evidence from HPLC-MS pointing to the presence of two distinct dimer reaction products prompted us to undertake structural analysis. HPLC fractions containing Dimer 1 and Dimer 2 were collected, frozen, lyophilized, and characterized by NMR with D₂O as the solvent (Figs. S5 – S16). Assuming C-C bond formation, DHPA dimers potentially contain an element of symmetry that would make the four ¹H atoms on the two aromatic rings appear at two frequencies instead of four. The relative position (ortho, meta, or para) of the two ¹H's on each ring can therefore be determined by the ¹H-¹H scalar coupling constant.⁴¹

Dimer 1, the compound isolated from the major dimer peak at ~7.5 min, presented two resonances in the aromatic portion of the ¹H spectrum (Figure 4a), indicating both rings had the same substitution pattern. The ¹H-¹H scalar coupling constant between these nuclei was determined to be 0.3 Hz by analysis of the 2D-J-resolved NMR spectrum (Figure S11), indicating that the ¹H atoms have a para relationship to one another.⁴¹ Thus, Dimer 1 was concluded to contain a C6-C6 linkage between the aromatic rings (Figure 4b and Figure S5).

In contrast, Dimer 2, the compound isolated from the minor dimer peak at ~10.5 min, presented four resonances in the aromatic portion of the ¹H spectrum (Figure 4a), indicating that the two rings had different substitution patterns. Analysis of the 2D-J-resolved NMR spectrum (Figure S16) revealed that one pair of resonances possessed a similar coupling constant to that of Dimer 1 (0.3 Hz), indicating that one ring has two hydrogens para to one another. The other pair of resonances exhibited a scalar coupling constant of 2.1 Hz, indicating a meta relationship between them. Thus, Dimer 2, was assigned a C5-C6 bond between the aromatic rings (Figure 4b and Figure S5).

Reduction of Fe³⁺ to Fe²⁺

A UV-Vis assay based on phenanthroline²⁸ was employed in order to demonstrate that oxidation of the catechol to the *o*-quinone was accompanied by reduction of Fe³⁺ to Fe²⁺. In this simple colorimetric assay, addition of phenanthroline to a Fe²⁺ solution produces a deep red color, whereas addition of phenanthroline to a Fe³⁺ solution (orange) produces no visible color change.²⁸ Therefore, the UV-Vis spectra of dilute solutions containing phenanthroline combined with DHPA, Fe³⁺, Fe²⁺, Fe³⁺+DHPA, and IO₄-DHPA were recorded (Figure S17). Notably, the reaction mixture of DHPA and Fe³⁺ yielded a strong absorbance between ~350 nm and ~550 nm in the presence of phenanthroline, identical to the spectrum obtained from a control mixture of phenanthroline and Fe²⁺ but different from that arising from the reaction of DHPA with NaIO₄. These data show that Fe³⁺ is reduced to Fe²⁺ during the dimerization of DHPA, likely through a quinone-mediated process.

Fe³⁺-induced cross-linking of a DOPA-containing peptide

The reactions between DHPA and Fe³⁺ described above were performed under unbuffered acidic conditions (pH ~2) as a consequence of the acidity of FeCl₃. Recognizing that mussel byssus formation occurs at higher pH, we investigated the effect of pH on the oxidation and cross-linking of catechols by Fe³⁺. Analysis of DHPA was complicated by the formation of insoluble precipitates at pH >4, presumably due to metal carboxylate salt formation. Instead, pH-dependent reactions were investigated using the DOPA-containing dipeptide Ac-Ser-DOPA-NH₂ (**2**, Figure 1c), in unbuffered water (pH ~2 due to FeCl₃ acidity) and at pH 5, 7, and 9 buffered with acetate, bis-tris, and bicine, respectively. Ac-Ser-DOPA-NH₂ is based on the repeating decapeptide motif of the mussel adhesive protein *mfp-1*.⁷ The buffers for these experiments were carefully chosen in order to eliminate unwanted side reactions, precipitation, and chelation of Fe³⁺. Figure 5 shows HPLC-MS chromatograms of the reactions between Ac-Ser-DOPA-NH₂ and Fe³⁺. Extracted ion chromatograms (EIC) were used to identify peaks (Supplemental Figure S18). In the presence of Fe³⁺, the dimer of the dipeptide is seen to form at pH 5, but significantly less so at higher pHs. Four dimers of the dipeptide peaks were identified (+), three of which appear to be present in significant quantities at pH 5 with 1× Fe³⁺. The dimer peak seen at ~6 min in the EIC (Figure S18) at pH 5 with 1× Fe³⁺, only appears to be significant in acidic unbuffered water with 1× Fe³⁺, along with many small unidentified peaks (seen in the chromatogram baseline). It appears that reactions of DOPA with 1× Fe³⁺ in unbuffered water are more promiscuous than in the buffered systems. At pH 9 in the absence of Fe³⁺, a peak at ~9 min was seen, however it could not be identified by HPLC-MS. It is likely attributable to an autoxidation product, as the solution was noted to develop a light pink color at this basic pH that remained even after acidification with EDTA. In the presence of Fe³⁺ at pH 9, this peak is not seen, supporting the idea that coordination of Fe³⁺ protects DOPA from autoxidation at high pH. The color of the reaction solutions in the pH 5–9 range was consistent with the formation of Fe-catechol coordination interactions.^{13, 18, 20}

Fe³⁺-induced cross-linking of *mefp-3* and *mcfp-3*

Purified DOPA-rich *mefp-1* and *mcfp-1*, MW ~110 kDa^{19, 42} and ~92 kDa,⁴³ respectively, were studied to determine whether Fe³⁺ could induce covalent cross-links in native proteins. *Mefp-1* has a DOPA content of approximately 11 % (~62 DOPA residues/protein),^{7, 43} while *mcfp-1* has an approximate DOPA content of 13 % (~95 DOPA residues/protein).⁴³ In a 0.1 M formic acid buffer solution (pH 3), 50 μM *mefp-1* or *mcfp-1* was incubated with increasing concentrations of Fe³⁺, up to 0.5 μM. As shown in Figure 6, the *mefp-1* and *mcfp-1* monomer bands are seen to decrease, while a new, higher MW band is seen to appear with increasing concentrations of Fe³⁺. The higher MW band is absent from the Fe³⁺ free experiment, indicating that Fe³⁺ is able to cross-link *mefp-1* and *mcfp-1*, likely through DOPA-DOPA covalent cross-links. Given the acidity of the solution, Fe³⁺-DOPA coordination cross-links would not be expected to occur.¹⁸

Discussion

Using catechol-containing small molecules and peptides, our results provide the first direct characterization of the covalent cross-linking of catechols induced by redox coupling with

Fe³⁺. For the reaction between Fe³⁺ and DHPA under acidic conditions, analysis of the reaction solutions revealed that formation of the *o*-quinone form of DHPA was accompanied by the reduction of Fe³⁺ to Fe²⁺ and the formation of dimers, trimers, and tetramers. One- and two-dimensional NMR analyses of HPLC fractions containing DHPA dimers unequivocally identified the major reaction product formed under these conditions to be coupled via a C6-C6 linkage between aromatic rings, with the remainder coupled via a C5-C6 linkage. Covalent multimer formation was also observed for reactions between Fe³⁺ and a DOPA-containing peptide at low pH (pH 2–3). However, the formation of covalent multimers in the presence of Fe³⁺ decreased as the pH was increased, and became nearly undetectable at pH 9.

These findings show a clear pH dependence of the interactions between Fe³⁺ and catechols, suggesting a refinement of existing views regarding the role of Fe³⁺ in mussel adhesion. To place our results into context, it is useful to consider the mechanism and conditions under which mussel byssal proteins are secreted.⁴⁴ Byssal threads and adhesive plaques are formed by injection molding of liquid precursors within the distal depression and ventral groove of the mussel foot, followed by release of the newly formed thread and adhesive plaque after several minutes of ‘curing’. Recent unpublished work has determined that the initial pH of secreted byssal proteins is ~4,⁴⁵ while published work from the same group had earlier indicated an initial pH of ~5.8 ± 0.3.⁴⁶ After release of the newly formed thread, rapid neutralization to marine pH (8–8.5) presumably occurs upon exposure to seawater. Therefore, the interactions between Fe³⁺ and catechols that are most relevant to mussel adhesion are likely to occur in the mildly acidic to mildly alkaline pH range (pH 4–8.5).

The formation of coordination complexes between Fe³⁺ and catechols within this pH range is well documented in the literature.^{13, 18–21} At catechol:Fe³⁺ ratios greater than 3, the bis Fe(DOPA)₂ complex is favored in the moderately acidic and neutral pH range, whereas tris coordination is favored at alkaline pH.²⁰ Tris Fe³⁺(DOPA)₃ coordination interactions may be responsible for the unique physical properties of the byssal thread cuticle,^{22, 46} which is rich in Fe³⁺ and the DOPA-containing protein *mfp-1*. The mechanical behavior of biomimetic polymer hydrogels formed through tris Fe³⁺-catechol coordination interactions leaves little doubt that such interactions are mechanically active and can contribute to solidification of the mussel byssus.^{23–26} Although covalent DOPA-DOPA cross-links have been detected in the mussel byssus,¹² the reaction pathway has not been well characterized. Given our results, one would expect that either Fe³⁺ is stored separately from the DOPA-containing mfps or an additional factor inhibits the oxidation of DOPA by Fe³⁺ observed in our studies. If Fe³⁺ was stored together with mfps premature cross-linking in byssal secretions would likely result, leading to a more viscous liquid or even a solid-like gel, making byssal formation difficult or even intractable.

A primary role for covalent cross-linking of mussel adhesive proteins by Fe³⁺ has been suggested,^{15, 32} however, the subtleties of byssal precursor storage and processing by the mussel are not well understood. The polyphenolic gland has been identified and demonstrated to contain *o*-diphenols.^{47, 48} If dilute FeCl₃ is incubated with the foot, this structure turns grey-green in color, and if Na₂CO₃ is subsequently added, the structure turns red.⁴⁷ This, together with research suggesting iron is secreted from the foot into the

thread,¹⁷ provides limited support for the idea that Fe³⁺ and the DOPA-rich *mfps* are stored separately in the glandular mussel foot. If this case were true, our results suggest that the mussel may use Fe³⁺ as an oxidant for DOPA during the initial (acidic) stages of byssal secretion to induce covalent DOPA-DOPA crosslinks. DOPA may then coordinate Fe³⁺ after the byssus is released into the basic marine environment to enhance the thread's mechanical properties (Figure 1b).

In a synthetic catechol-modified poly(ethylene glycol) system, we recently demonstrated that the mechanical properties of hydrogels could be dramatically enhanced by applying a similar pH switch in the presence of Fe³⁺.²⁴ In that system, Fe³⁺ was first used to covalently cross-link the hydrogel at acidic pH and then the pH was adjusted to a basic pH to allow for Fe³⁺-catechol coordination bonds to form. At higher pH, gels had an increased modulus and an improved energy dissipating capacity, mechanical features that are important for the byssal thread. Similarly, a recent DOPA-modified peptide amphiphile network demonstrated increased modulus and self-healing properties in the presence of Fe³⁺ at basic pH.²⁷

While further work is needed to definitively elucidate the interactions between Fe³⁺ and the various *mfps* before, during, and after byssus formation, we hypothesize that a complex role for Fe³⁺ may exist – one that includes both covalent and coordination interactions that are strongly modulated by pH. The mechanical implications of covalent versus coordination cross-links for the performance of the mussel byssus cannot be ignored. Evidence at the single molecule³ and macroscopic gel^{23–26} levels indicate that fundamentally different mechanical behavior emerges when catechol polymers are cross-linked via coordination and covalent interactions. It may be that mussels achieve the optimal balance between these two types of cross-links through modulation of pH during byssus formation. While an oxidative role for Fe³⁺ in the mussel adhesive remains hypothetical at this point, these results are highly relevant to the many bioinspired systems where catechols and iron are present.^{4, 5, 23–27}

Conclusions

Herein, we have characterized the products of Fe³⁺-induced oxidation of the catechol DHPA. Through HPLC-MS and NMR, we have shown that the dominant products are the C6-C6 and C5-C6 connected dimers. HPLC-MS experiments involving a DOPA-containing peptide showed that at acidic pH Fe³⁺ induces covalent bond formation, whereas at basic pH, DOPA is protected through coordination bond formation and covalent bonds are not formed. Using the DOPA-rich mussel foot proteins, *mefp-1* and *mcfp-1*, we have shown that Fe³⁺ can induce covalent bond formation between the proteins at acidic pH. We have discussed the implications of this chemistry to byssal thread and mussel adhesive formation, hypothesizing that Fe³⁺ may play both an oxidant role to induce covalent DOPA-DOPA bond formation and act as a coordination center for DOPA, and that these roles may be switched with the temporal acidic to basic pH change that occurs during byssal secretion and release. We further discuss the mechanical implications of such chemistry and the possible advantages.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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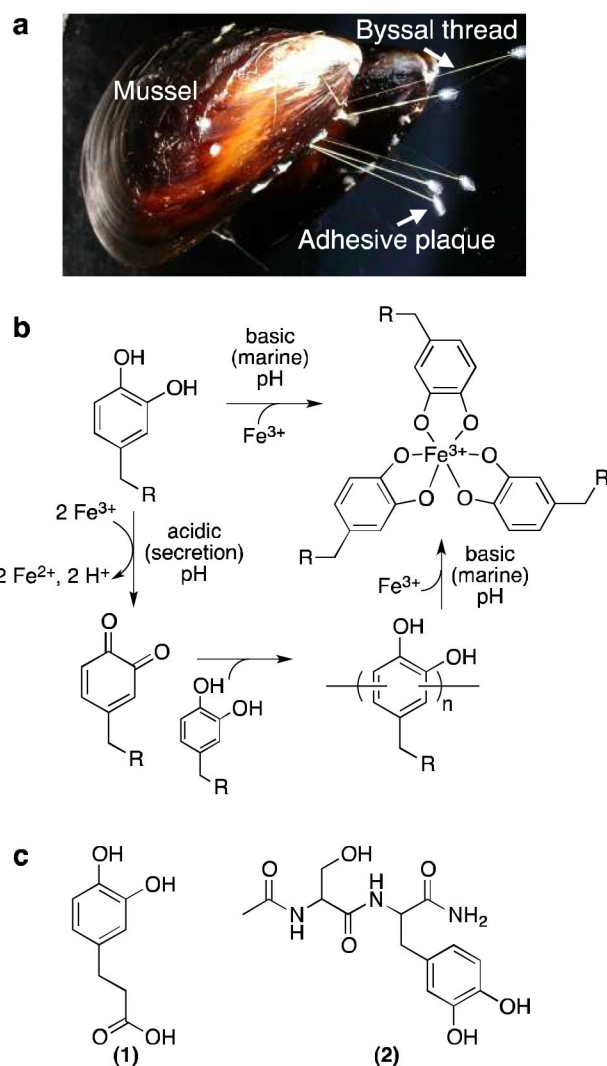


Figure 1. Mussel byssus, model for pH-dependent Fe^{3+} -catechol chemistry, and molecules studied. (a) Picture of mussel with labeled byssal structures. (b) Model for pH-dependent interactions between Fe^{3+} and catechol. At acidic pH, catechols and Fe^{3+} react to produce *o*-quinones, leading to catechol-catechol oligomerization. At basic pH, catechols form coordination bonds with Fe^{3+} that prevent oxidation of the catechol. In the context of the mussel byssus, Fe^{3+} could hypothetically induce covalent cross-linking during acidic processing of byssal protein secretions and enhance the mechanical properties of the byssal thread and adhesive by formation of mechanically active Fe^{3+} -catechol coordination interactions upon equilibration to basic marine pH. Note that Fe^{3+} -catechol coordination bonds would be expected to be mechanically reversible, whereas covalent catechol-catechol bonds would not.²³ (c) Catechol-containing molecules studied: (1) DHPA and (2) Ac-Ser-DOPA-NH₂ (dipeptide).

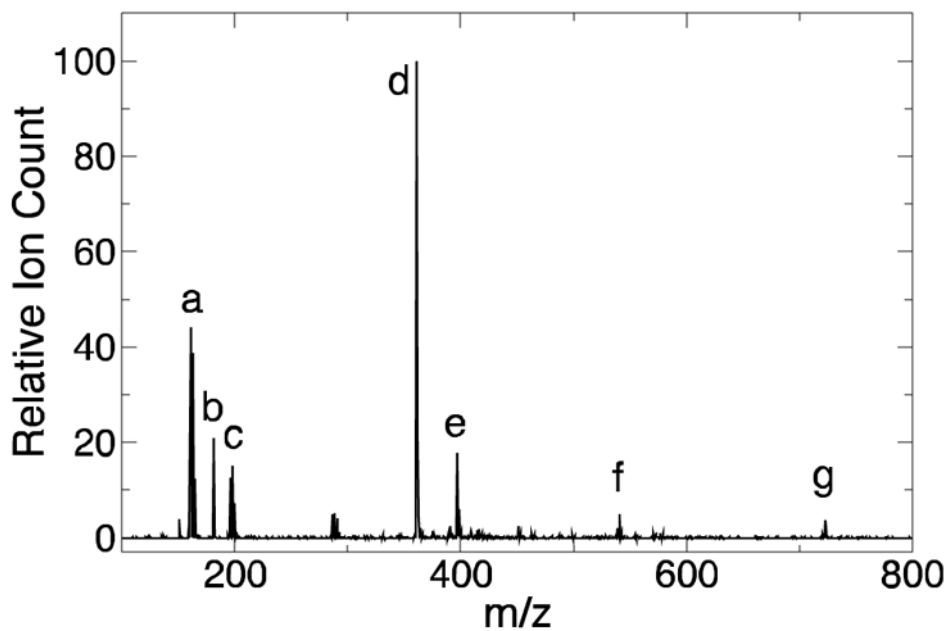


Figure 2. Fe^{3+} induces DHPA multimer formation. Representative negative ion MS spectrum of direct injection of reaction mixture (Reaction 5, Fe^{3+} :DHPA of 6:3) after 7 d. The solution contained iron species (peaks a and c at m/z of 161 and 198, respectively), DHPA (peak b at m/z of 181), dimers (peak d at m/z of 361), trimers (peak f at m/z of 541), and tetramers (peak g at m/z of 721). Peak e represents dimers associating with a chloride ion (m/z of 397).

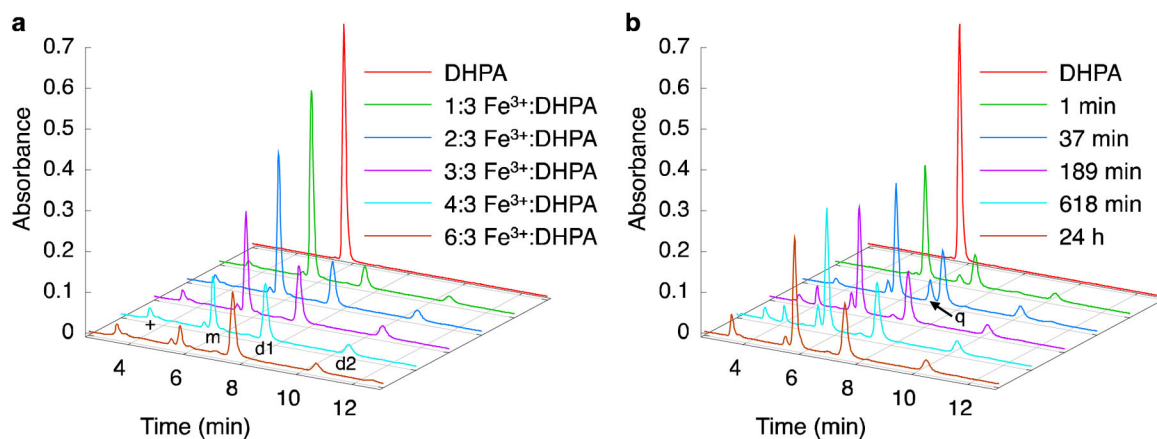


Figure 3.

Two DHPA-dimer species are produced after Fe^{3+} -coupled oxidation. HPLC chromatograms (280 nm) of reactions between DHPA and Fe^{3+} . (a) Effect of stoichiometry on the reaction between DHPA and Fe^{3+} after 7 d. The peaks at ~3.5 min (+), ~5.5 min (m), ~7.5 min (d1), and ~10.5 min (d2) correspond to soluble Fe^{3+} and Cl^- ions, DHPA, Dimer 1, and Dimer 2; these results were determined by HPLC-MS (Figure S4). (b) Time dependence of the reaction between DHPA and Fe^{3+} (1:1 molar equiv.). The peaks are identified as in (a), with the addition of a small peak at ~7 min (q) that corresponds to the *o*-quinone form of DHPA.

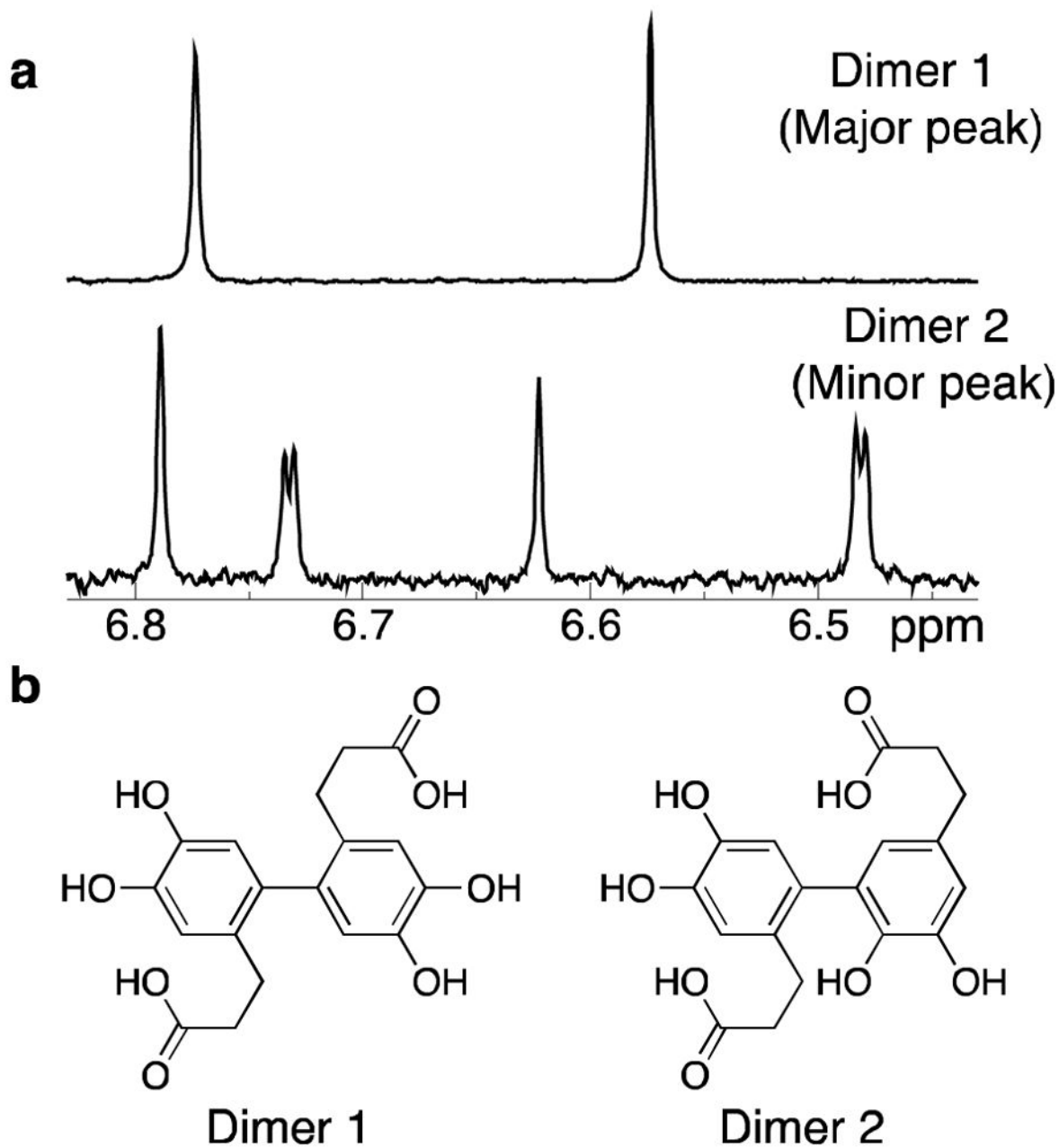


Figure 4.

Connectivity of DHPA dimers determined by NMR. Structural analysis of dimers formed by reaction of DHPA and Fe^{3+} . (a) Aromatic regions of the 500 MHz ^1H 1D NMR spectra of dimers eluting at ~ 7.5 min (Dimer 1) and ~ 10.5 min (Dimer 2) from Figure 3. Dimer 1 exhibits two signals, indicating a symmetric species; the ^1H - ^1H scalar coupling is smaller than the signals' linewidth (~ 1.0 Hz), indicating the hydrogens are para to one another. Dimer 2 exhibits four signals, indicating an asymmetric species; the two scalar couplings are < 1.0 Hz and 2.2 Hz, indicating para and meta ^1H - ^1H relationships, respectively. (b) Dimer 1 is connected in a C6-C6 fashion, resulting in a symmetric molecule with hydrogens that are

para to one another. Dimer 2 has a C5–C6 linkage, producing an asymmetric molecule with two hydrogens para to one another and two hydrogens meta to one another.

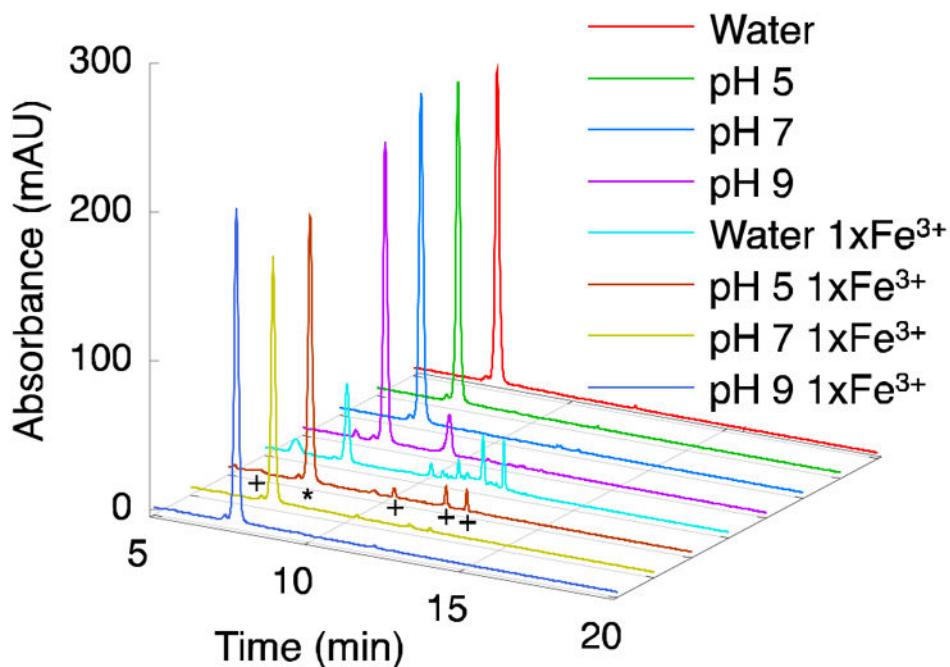


Figure 5.

DOPA-containing dipeptide forms dimers at acidic pH in the presence of Fe^{3+} . DOPA-containing dipeptide (Ac-Ser-DOPA-NH_2) reaction with and without Fe^{3+} as studied by HPLC-MS (280 nm shown) at different reaction pHs. Peaks were identified as: ~7 min, dipeptide (*); dimers of dipeptide (+) were detected at ~6, ~10, ~11, and ~12 min (seen prominently in pH 5 and water $1 \times \text{Fe}^{3+}$ samples); peak in pH 9 sample at ~9 min, unidentified. In unbuffered water (pH ~2) many unidentified products are produced, while at pH 5 three dimers are the dominant species formed after the reactions with Fe^{3+} . Dimer at ~6 min is seen prominently only in water $1 \times \text{Fe}^{3+}$ sample. At higher pH in the presence of Fe^{3+} , the dipeptide appears to be protected from oxidation by the presence of Fe^{3+} .

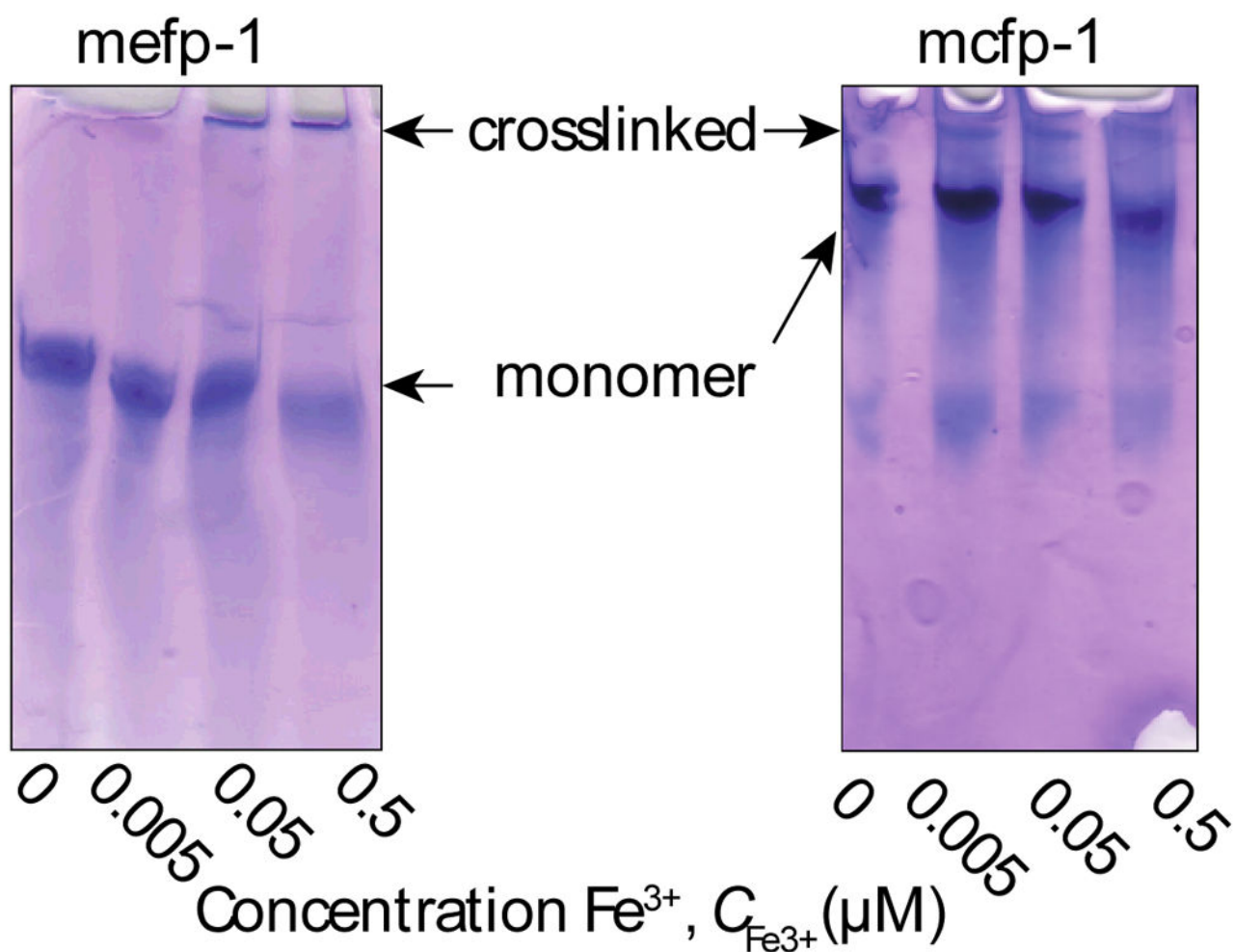


Figure 6.

Effect of Fe^{3+} on protein crosslinking. Purified protein (*mefp-1* and *mcfp-1*) in 0.1 M formic acid pH 3.0 was incubated at room temperature with varying iron concentrations (0–0.5 μM) and run on an acid-urea polyacrylamide gel. Higher molecular weight bands represent dimer or higher oligomer formation. Note that the *mefp-1* gel was run for 2 h while the *mcfp-1* gel was run for 1 h.

Table 1Fe³⁺-DHPA reactions and residual DHPA after 7 d

Reaction	[Fe ³⁺] (mM)	[DHPA] (mM)	[Fe ³⁺]:[DHPA]	% DHPA unreacted ^a
1	20	60	1:3	77
2	40	60	2:3	57
3	60	60	3:3	43
4	67	50	4:3	29
5	100	50	6:3	12

^a As determined by HPLC (Figure 3a)