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Adsorption Mechanism of Myelin Basic Protein on Model Substrates and Its Bridging Interaction between the Two Surfaces

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Supporting Information

ABSTRACT: Myelin basic protein (MBP) is an intrinsically disordered (unstructured) protein known to play an important role in the stability of myelin’s multilamellar membrane structure in the central nervous system. The adsorption of MBP and its capacity to interact with and bridge solid substrates has been studied using a surface forces apparatus (SFA) and a quartz crystal microbalance with dissipation (QCM-D). Adsorption experiments show that MBP molecules adsorb to the surfaces in a swollen state before undergoing a conformational change into a more compact structure with a thickness of ∼3 nm. Moreover, this compact structure is able to interact with nearby mica surfaces to form adhesive bridges. The measured adhesion force (energy) between two bridged surfaces is 1.0 ± 0.1 mN/m (E_ad = 0.21 ± 0.02 mJ/m²), which is slightly smaller than our previously reported adhesion force of 1.7 mN/m (E_ad = 0.36 mJ/m²) for MBP adsorbed on two supported lipid bilayers (Lee et al., Proc. Natl. Acad. Sci. U.S.A. 2014, 111, E768–E775). The saturated surface concentration of compact MBP on a single SiO₂ surface reaches a stable value of 310 ± 10 ng/cm² regardless of the bulk MBP concentration. A kinetic three-step adsorption model was developed that accurately fits the adsorption data. The developed model is a general model, not limited to intrinsically disordered proteins, that can be extended to the adsorption of various chemical compounds that undergo chemical reactions and/or conformational changes upon adsorbing to surfaces. Taken together with our previously published data (Lee et al., Proc. Natl. Acad. Sci. U.S.A. 2014, 111, E768–E775), the present results confirm that conformational changes of MBP upon adsorption are a key for strong adhesion, and that such conformational changes are strongly dependent on the nature of the surfaces.

INTRODUCTION

Myelin basic protein (MBP) is an essential protein for stability of the myelin sheath.1–3 MBP is one of the major proteins (20% of proteins) found in the myelin sheath that surrounds the axons of the central nervous system (CNS). The myelin sheath is a compact stack of lipid bilayers that alternate between cytoplasmic and extracellular leaflets in the radial direction (see Figure 1). MBP acts as a protein glue that bridges and compacts the cytoplasmic leaflets of myelin. Healthy myelin exhibits a very compact structure with a thin water gap distance (∼3–4
nm) between bilayers of the cytoplasmic side, providing a low dielectric constant through the compact bilayers, which in turn allows the axon to transmit electrical impulses in a more efficient and faster manner compared to demyelinated axons. Structural changes in the myelin sheath, such as water gap swelling, vacuolization, vesiculation, lesion formation, and delamination are indications of neurological disorders. Such neurological disorders are usually embodied by a wide spectrum of symptoms including physical and cognitive disabilities, with multiple sclerosis (MS) being the most common disorder.

Figure 1. Schematics of the (a) nervous system, (b) myelin, and (c) myelin basic protein (MBP). For illustrative purposes, the intermembrane separation of the cytoplasmic space appears larger than the extracellular space. However, the extracellular space is actually larger than the cytoplasmic space.

Myelin basic protein belongs to a family of intrinsically unstructured (disordered) proteins with a predominant isoform named C1 with a molecular weight of 18.5 kDa and a net positive charge of 19. Previous studies conducted with model and extracted bilayers have shown that MBP can bind to a negatively charged bilayer via electrostatic and hydrophobic interactions. Recent studies have shown that minute changes in lipid composition lead to changes in lipid domains, where the structure and size of these domains significantly affects the MBP adsorption mechanism, eventually leading to swelling and loss of adhesion between myelin layers.

Structural studies on MBP with circular dichroism (CD) spectroscopy have shown that no structural order exists in the protein when in solution. Electron microscopy and solid-state NMR (SSNMR) studies have found that MBP bound to lipid monolayers exhibits a C-shape or hairpin-like structure. However, no study has been performed on the kinetics of MBP adsorption while simultaneously monitoring the changes in conformation of MBP after adsorption to a surface. This study aims to establish a kinetic model that will explain the adsorption and conformational changes of MBP onto a model surface, and how the conformation affects the bridging interaction between the surfaces. In order to measure the bridging force between model surfaces, a surface forces apparatus (SFA) is used. A quartz crystal microbalance with dissipation (QCM-D) was used to measure the adsorption of MBP to SiO2 surfaces as a function of adsorption time and bulk MBP concentration. A 3-step adsorption model was derived to explain the detailed kinetic information obtained from the QCM-D.

MATERIALS AND METHODS

Materials. The C1 isoform of myelin basic protein was isolated from bovine brain white matter and kept in a deep freezer until use. MBP was dissolved in buffer (pH 7.4) composed of 150 mM sodium nitrate (Sigma-Aldrich, Inc. ReagentPlus ≥ 99%), 10 mM Mops.

Figure 2. (a) Force–distance (F–D) profiles of MBP between two mica surfaces at two difference concentrations of 25 and 100 μg/mL after 3 h equilibration (3 h after injecting MBP solution between the surfaces), and (b) the schematics showing the possible structuring of MBP during the force runs.

3160
sodium salt (Sigma-Aldrich, Inc. ≥ 99.5%), and 2 mM calcium nitrate tetrahydrate (Sigma-Aldrich, Inc. ≥ 99%) to make a desired concentration of MBP solution. Aliquots of MBP solutions were kept frozen (−50 °C) in acid-cleaned vials until use.

**Methods.** **SFA Experiments.** An SFA 2000 (SurForce, LLC) was used for the force measurements.25 Freshly cleaved and back-silvered mica sheets were glued onto two cylindrical disks with curvature radii (R) of ~2 cm. The surfaces were mounted in the SFA in a cross-cylindrical geometry, which is mathematically equivalent to a sphere-on-flat geometry (see Figure 2b). The separation distance between two surfaces was measured using optical interferometry,26 and the forces between the two surfaces was measured using a double cantilever spring holding the lower disk. MBP solution was injected between the surfaces, followed by a 15 min equilibration time. For the force-distance measurements, the lower surface was advanced toward the upper surface using a fine-control motorized micrometer with an approach velocity of 0.5–1.0 nm/s, followed by separation of the two surfaces at the same velocity, in order to measure the adhesion force. After the first force run, MBP was allowed to further and completely adsorb and equilibrate on the surfaces for 3 h. This was followed by a second and subsequent force runs, as MBP was found to undergo time-dependent conformational changes.

**QCM-D Experiments.** A Quartz Crystal Microbalance with Dissipation (QCM-D E4, Biolin Scientific)27 was used to measure the mass density of adsorbed MBP. A silicon dioxide (SiO2) sensor (Bolin Scientific) was used as the substrate since the SiO2 surface is negatively charged like the myelin sheath. SiO2 sensors were rinsed with ethanol, followed by exposure to UV-ozone plasma for 30 min before use. The baseline was calibrated by injecting the buffer (mentioned in the Materials section) using a peristaltic pump with a flow rate of Q = 400 μL/min. Based on the information from QSense, Q = 400 μL/min gives a flow velocity of 0.96 m/s, corresponding to a Reynolds number of Re = 9.1, which indicates that the system is in the laminar flow regime (Re < 2300). After the frequency and dissipation values became stable, MBP solution at three different concentrations (25, 10, and 5 μg/mL) was injected simultaneously to three different QCM-D chambers for 1 h. The frequency changes (of the third and fifth harmonics) were converted to surface concentrations using the Sauerbrey equation,28 which is an excellent approximation for a low-dissipation system (∆F/∆D < 10−5/10 Hz),29 where ∆D is the dissipation change.

**RESULTS AND DISCUSSION**

Effects of Bulk MBP Concentration on the Interaction Forces between Mica Surfaces.** Figure 2a shows the normalized force (F/R) versus distance (D) curves of MBP at two bulk concentrations (C = 25 and 100 μg/mL) after equilibrating for 3 h. Zero separation distance (D = 0) is defined as mica–mica contact in air. At C = 25 μg/mL, the steric “hard-wall” thickness, D_{stenso} was 3 nm, similar to the thickness of MBP (σ_{SMBP}) in its compact C-shape conformation (see schematic in Figure 2b).16 On the approach, the steric repulsion due to trapped MBP between the mica surfaces starts around D = 6 nm, which is comparable to 2σ_{MBP}. The measured adhesion force, F_{dR} = 1.1 mN/m, which corresponds to adhesion energy of E_{ad} = F_{dR}/1.5πR = 0.23 mJ/m² using Johnson–Kendall–Roberts (JKR) theory,30 is slightly smaller compared to a previously measured adhesion force, F_{dR} = 1.7 mN/m (E_{ad} = 0.36 mJ/m²), of MBP between normal (nonpathological) myelin lipid bilayers.22 During separation from adhesive contact, the MBP molecules were found to stretch 5 nm before adhesive detachment occurred. This length is less than the contour length (15 nm)16,34 and much less than the fully extended length (50 nm) of MBP.19

The force–distance profiles at MBP concentration of C = 100 μg/mL showed the following differences compared to the...
profiles at \( C = 25 \ \mu g/mL \): (i) the steric repulsion on approach is more pronounced with the repulsion starting at \( D = 30 \ \text{nm} \); (ii) \( D_{\text{steric}} \) was equal to 6 nm or \( 2D_{\text{MBP}} \) at \( F/R = 6 \ \text{mN/m} \), indicating a double layer of MBP (see schematic in Figure 2b); (iii) the adhesion force is weaker \( (F_{ad}/R = 0.7 \ \text{mN/m}, E_{ad} = 0.23 \ \text{mJ/m}^2) \), and (iv) the surfaces detached at a larger distance \( (D_1 = 18 \ \text{nm}) \) before they jumped apart, which indicates that the MBP molecules stretch farther than at \( C = 25 \ \mu g/mL \) (Figure 2b).

These results demonstrate that at low solution concentration of MBP \( (C = 25 \ \mu g/mL) \), MBP molecules form a more compact monolayer film with a higher adhesion force, while at a high MBP concentration \( (C = 100 \ \mu g/mL) \), the protein forms a multilayer (gel-like) film with stronger steric repulsion, thicker steric wall thickness, lower adhesion force/energy (due to cohesion between the MBP molecules), and more pronounced stretching of the molecules before the surfaces detach (jump apart in SFA experiments).

**Conformational Changes of Adsorbed MBP upon Adsorption.** The structure of MBP changes in the presence of lipids; however, it is a slow process that can take several minutes. Figure 3 shows how the force–distance profile changes as MBP is allowed to structurally rearrange on the mica surfaces. The first force runs (black circles) show the force distance profiles 15 min after MBP injection, while subsequent force runs (red circles) were performed after 3 h. After 3 h, no further changes are observed in the force distance profiles, which led to the conclusion that MBP reached its equilibrium conformation. To test for reproducibility, we also performed force runs immediately after the first, which gave the same force profiles as in the first run, and had no effects on the second run after 3 h, which was also obtained if the two surfaces were brought together for the first time after 3 h. Thus, the same force profiles are obtained 3 h after the adsorption irrespective of whether or not the surfaces are previously brought together. These tests show that any long-term changes, e.g., after 3 h, after the adsorption time, are due to “natural” relaxations of the MBP and not due to the “push-pull” effects of previous force runs.

At \( C = 25 \ \mu g/mL \), no adhesion (i.e., purely repulsive forces) was measured between the two mica surfaces with a layer of MBP trapped between them after 15 min. However, after 3 h, the adhesion \( F_{ad}/R \) increases up to \( \sim 1.1 \ \text{mN/m} \) \( (E_{ad} = 0.23 \ \text{mJ/m}^2) \), even though only minor changes in the steric wall distance, \( D_{\text{steric}} \), were found. The appearance of an adhesive contact indicates that MBP molecules underwent a structural change that favors adhesion between MBP molecules and the mica surface. The positive charges on MBP will conform to the negative mica surface, while the hydrophobic groups of MBP will oppose contact with the hydrophilic mica surface. Also, hydrophobic groups of MBP molecules will attract similar groups on other MBP molecules in close proximity. The two effects of MBP–mica (adhesion) and MBP–MBP (cohesion) interactions that lead to structural changes are modeled later (see below).

At \( C = 100 \ \mu g/mL \), MBP molecules formed a thicker layer \( (D_{\text{steric}} = 10 \ \text{nm} \) at \( F/R = 6 \ \text{mN/m} \)) compared to at \( 25 \ \mu g/mL \) \( (D_{\text{steric}} = 3 \ \text{nm} \) at \( F/R = 6 \ \text{mN/m} \)), which became more compact after 3 h \( (D_{\text{steric}} = 6 \ \text{nm} \) at \( F/R = 6 \ \text{mN/m} \)). On the other hand, significant adhesion \( (F_{ad}/R = 0.8 \ \text{mN/m}, E_{ad} = 0.17 \ \text{mJ/m}^2) \) was observed shortly after the injection of MBP, which only slightly increases \( (F_{ad}/R = 0.9 \ \text{mN/m}, E_{ad} = 0.19 \ \text{mJ/m}^2) \) after 3 h. The significant inward shift of the steric hard wall distance indicates a structural change of the MBP film. Furthermore, the cohesive interaction force seems to be independent of conformation or protein layer thickness.

QCM-D measurements provided further detailed information on the MBP adsorption mechanism and its associated structural changes. Figure 4 shows plots of surface concentration \( (C_{\theta} \) obtained by the Sauerbrey equation) of MBP on an SiO2 surface as a function of the adsorption time \( (t) \). At \( C = 25 \ \mu g/mL \), the surface concentration of MBP, \( C_{\theta} \), increased rapidly and peaked at \( 450 \ \text{ng/cm}^2 \) at \( t = 90 \ \text{s} \) after injection of the MBP solution, and then settled to \( C_{\theta} = 320 \ \text{ng/cm}^2 \) after \( t = 15 \ \text{min} \) \( (900 \ \text{s}) \). The rate of surface coverage after injection of MBP solution decreases with decreasing concentrations, and the initial overshoot of surface coverage also decreases with decreasing concentration and was nonexistent at concentrations below \( C = 5 \ \mu g/mL \). At \( C = 10 \ \mu g/mL \), the \( C_{\theta} \) reached a peak value of \( 355 \ \text{ng/cm}^2 \) after \( t = 4 \ \text{min} \), which equilibrated to \( C_{\theta} = 305 \ \text{ng/cm}^2 \) after \( t = 30 \ \text{min} \). At \( C = 5 \ \mu g/mL \), no peak was observed, and \( C_{\theta} \) slowly increased to an equilibrium value of \( C_{\theta} = 305 \ \text{ng/cm}^2 \).

It is interesting to note that, regardless of the bulk MBP concentration, \( C_{\theta} \), the surface concentration equilibrated to similar values \( (C_{\theta} = 310 \pm 10 \ \text{ng/cm}^2) \) after 30 min. Together with the dissipation data measured with QCM-D (Figure 5), these results suggest that MBP molecules form a weakly bound and soft preadsorbed layer on an SiO2 surface, followed by conformational changes to a more compact (and thinner) and more stable structure (see the inset schematic in Figure 5) that occupies a larger area per molecule. The results also agree with the SFA results (see Figure 3b), which showed an initial thick layer that equilibrates into a more compact layer after 3 h.

**Modeling the MBP Adsorption Mechanism.** Based on the SFA and QCM-D results, we propose a “three-step model” (see Figure 6a) for the MBP adsorption mechanism: (i) Bulk MBP moves from the bulk, across the concentration boundary layer, to the “subsurface” (bulk region in close proximity to the silica surface); (ii) MBP molecules from the subsurface adsorb...
to the surface, and (iii) the adsorbed MBP molecules undergo a conformational change. This model can be expressed using the following equation:

\[
A \xleftrightarrow{k_m} B \xrightarrow{k_1} C \xrightarrow{k_2} D
\]  

(1)

where, A, B, C, and D represent MBP in the bulk, subsurface MBP, adsorbed MBP, and adsorbed MBP after a conformational change, respectively. Here, \(k_m\) is the mass transfer coefficient for the diffusion of MBP across the boundary layer, \(k_1\) is the rate constant for the adsorption of MBP from the subsurface to the surface, \(k_{-1}\) is the rate constant for desorption of the adsorbed MBP, and \(k_2\) is the rate constant for the conformational change of adsorbed MBP. For simplifying the model, we assume that (i) the concentration boundary layer is at steady state (this is justified in Supporting Information section SI#1); (ii) the adsorbed MBP molecules can either detach or change their conformation; (iii) the conformational change of MBP on the surface is irreversible, and (iv) after the conformational change (state C to D), MBP is strongly bound to the surface and does not detach to the subsurface.

The MBP adsorption onto a silica surface was measured by a QCM-D at three different bulk concentrations of MBP, \(C_{A0} = 25, 10, \text{ and } 5 \mu g/mL\) (Figure 4 and 6b–d). These three solutions were guided into the QCM-D chamber (volume = 40 \(\mu\)L and pre-filled with buffer solution) at a constant volumetric flow rate of \(Q = 400 \mu\text{L/min}\). The chamber concentration of MBP, \(C_A\), increases from 0 (pure buffer) to \(C_{A0}\) after a residence time \(\tau\), which is 0.1 min; therefore, data below time scales of this order was ignored when fitting data to the model.
The rate of transfer of MBP across the boundary layer (from state A to B in eq 1) can be modeled using the standard mass transport relationship:

\[ J_A = k_m (C_{A0} - C_B) \]  
(2)

where \( C_B \) is the bulk concentration of MBP at the subsurface. The MBP can go from the subsurface to an adsorbed state (state C in eq 1) at the surface. The net adsorption rate of adsorbed MBP, \( \frac{dC_B}{dt} \), is directly proportional to the subsurface MBP concentration \( (C_B) \) and \( \theta_B \), the fraction of vacant sites at the silica surface, given by the following equation:

\[ \theta_B = 1 - \frac{C_C}{C_{Sat}} - \frac{C_D}{C_{Sat}} \]  
(3)

Here, \( C_D \) is the MBP concentration after the conformational change, and \( C_{Sat} \) and \( C_{Sat} \) are the saturated surface concentrations of MBP when it completely covers the surface with initially adsorbed MBP only or with conformationally changed MBP only, respectively. To calculate \( C_{Sat} \) we approximate the adsorbed form of MBP (MBP in state C) using prolate ellipsoidal geometry. Assuming a hydrodynamic area of 400 Å²/molecule of the adsorbed MBP in state C, we calculate a complete coverage of \( C_{Sat} \) as 768.3 ng/cm². The complete coverage of conformationally-changed MBP at the silica surface, \( C_{Sat} \), is equal to the final surface concentration of MBP molecules found from the QCM-D measurements (see Figure 4). After a sufficiently long time has elapsed (\( t > 60 \) min), the QCM-D data of all concentrations of MBP approach the same value of \( C_{Sat} \) = 304 ng/cm², where the silica surface is completely covered with MBP molecules in their final stable structure.

A balance of the MBP molecules at the subsurface yields (see eq 1):

\[ k_m(C_{A0} - C_B) = k_{B1}C_B - k_{C1}C_C \]  
(4)

This provides the following expression for the subsurface MBP concentration, \( C_B \):

\[ C_B = \frac{k_mC_{A0} + k_{C1}C_C}{k_m + k_{B1}} \]  
(5)

The rate of adsorbed MBP in state C can be expressed as the following Langmuir-type differential equation (see eq 1):

\[ \frac{dC_C}{dt} = k_{C1}C_B\theta_B - k_{C1}C_C - k_{C2}C_C \]  
(6)

Using eq 5, the above equation can be simplified to

\[ \frac{dC_C}{dt} = k_m\left(\frac{k_{B1}C_{A0} - k_{C1}C_C}{k_m + k_{B1}}\right) - k_{C2}C_C \]  
(7)

The rate of the MBP conformation change at the silica surface can be expressed by the following, simple kinetic equation (see eq 1):

\[ \frac{dC_B}{dt} = k_{C2}C_C \]  
(8)

The total surface concentration, \( C_{mb} \) measured from the QCM-D, is the sum of \( C_C \) and \( C_D \):

\[ C_B = C_C + C_D \]  
(9)

Table 1 summarizes the overall equations and units for the proposed three-step adsorption model given in eqs 3 and 7–9.

### Table 1. Equations for the Proposed Three-Step Model

| Equation | Description |
|----------|-------------|
| \[ \frac{dC_C}{dt} = k_{C1}C_B\theta_B - k_{C1}C_C - k_{C2}C_C \] | Rate of adsorbed MBP in state C |
| \[ \frac{dC_B}{dt} = k_{C2}C_C \] | Rate of transfer of MBP across the boundary layer |

### Table 2. Parameters for the Fits in Figure 6

| Concentration (μg/mL) | \( k_m \) | \( k_{B1}/k_{C1} \) | \( k_{C2} \) |
|-----------------------|-----------|-----------------|-------------|
| 25                    | 0.0259    | 0.0457          | 0.135       |
| 10                    | 0.0244    | 0.0719          | 0.0461      |
| 5                     | 0.0192    | 0.0222          | 0.570       |

The above equation suggests that the dynamics of \( C_C \) does not depend separately on \( k_1 \) and \( k_{-1,1} \), but only on the ratio \( k_{1} \).
This was experienced during the fitting process; the residual and the agreement between theory and experiment were relatively insensitive to the individual values of $k_1$ and $k_{-1}$ as long as their ratio was close to the optimum value. The mean mass transfer coefficient in the three experiments is 0.023 cm/min. Since $k_1$ has to exceed $k_{-1}$ to be in the mass-transfer-dominated limit, 0.023 cm/min represents an approximate lower bound for the rate constant $k_1$ in the experiments, and the corresponding, approximate lower bounds for $k_{-1}$ for the three bulk MBP concentrations, 25 μg/mL, 10 μg/mL and 5 μg/mL, are 0.503 min$^{-1}$, 0.320 min$^{-1}$ and 1.04 min$^{-1}$, respectively.

**CONCLUSIONS**

Direct measurements using an SFA and QCM-D provide quantitative and qualitative information about the buildup of MBP layers and the bridging adhesions of MBP, and the kinetics of MBP adsorption to surfaces and the conformational changes of MBP with time (energies). The bridging forces (energies) and the thickness of the MBP monolayer between two mica surfaces are $F_{ad}/R = 1.0 \pm 0.1$ mN/m ($E_{ad} = 0.21 \pm 0.02$ mJ/m$^2$) and $D_{Steric} = 3$ nm, respectively (at a bulk concentration of $C = 10$ mg/mL and 3 h after injection of MBP).

On silica surfaces, the final equilibrium structure of MBP occupies 310 ± 10 ng/cm$^2$ on a silica surface. Covering a bare silica surface with MBP molecules at equilibrium takes 30 min. The adsorption of MBP on a SiO$_2$ surface can best be described by a three-step adsorption model, where bulk MBP diffuses across the concentration boundary layer, adsorbs to the SiO$_2$ surface, and undergoes a slow, conformational change to a stable and adhesive structure. The derived adsorption model can be used to fit the adsorption kinetics of a wide range of intrinsically disordered proteins and perhaps extended further to the adsorption of various chemical compounds, which undergo chemical reactions or conformational changes after adsorbing to surfaces.

This study was performed on model solid substrates, rather than on and between myelin lipid bilayers. Therefore, we are likely ignoring some important effects that would be present in naturally-occurring lipid bilayers, such as lipid–protein coupling, hydrophobic interactions between lipids and proteins, undulation of lipid bilayers, and effects from lipid domains. Nevertheless, this study still provides qualitative information on MBP adsorption and bridging mechanisms to negatively charged SiO$_2$, and mica surfaces, resembling charged myelin lipid bilayers.

**ASSOCIATED CONTENT**

Supporting Information

Estimation of boundary layer thickness and time for boundary layer development, non-dimensionalized equations with fitting parameters, and raw QCM data. This material is available free of charge via the Internet at http://pubs.acs.org.

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Notes

The authors declare no competing financial interest.

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