Article

Interspecies-Extrapolated Biotic Ligand Model to Predict Arsenate Toxicity to Terrestrial Plants with Consideration of Cell Membrane Surface Electrical Potential

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Abstract: Arsenic is a metalloid that is highly toxic to living organisms in the environment. In this study, toxicity caused by inorganic arsenate (As(V)) to terrestrial plants, such as barley \textit{Hordeum vulgare} and wheat \textit{Triticum aestivum}, was predicted using the existing biotic ligand model (BLM) for bioluminescent \textit{Aliivibrio fischeri} via interspecies extrapolation. Concurrently, the concept of cell plasma membrane electrical potential (Ψ\textsubscript{0}) was incorporated into the extrapolated BLM to improve the model predictability in the presence of major cations such as Ca\textsuperscript{2+}. The 50% effective As(V) toxicity (EC\textsubscript{50}(\text{HAsO}_4^{2-})) to \textit{H. vulgare} decreased from 45.1 ± 4.34 to 15.0 ± 2.60 µM as Ca\textsuperscript{2+} concentration increased from 0.2 to 20 mM owing to the accumulation of H\textsubscript{2}AsO\textsubscript{4}\textsuperscript{-} and HAsO\textsubscript{4}^{2-} on the cell membrane surface. The extrapolated BLM, which only considered inherent sensitivity, explained well the alteration of As(V) toxicity to \textit{H. vulgare} and \textit{T. aestivum} by Ca\textsuperscript{2+} with an order of magnitude, when considering a linear relationship between Ψ\textsubscript{0} and EC\textsubscript{50}(\text{HAsO}_4^{2-}).

Keywords: acute toxicity prediction model; species specificity; inherent sensitivity; environmental modulator; root cell plasma membrane surface electrical potential

1. Introduction

Arsenic (As) accumulates in environmental media via both natural and anthropogenic sources and can cause severe adverse effects on human health as well as ecosystems [1–5]. Because the total As concentration in the environment media cannot accurately indicate bioavailability or ecotoxicity, it is important to rationally assess the ecotoxicity of As using suitable bioassays [6]. The ecotoxicity of As varies greatly depending on environmental factors (e.g., phosphate concentration) based on which the bioassay is performed [7,8] and on the sensitivity of the species tested [9]. The environmental factors can be considered using the biotic ligand model (BLM), which is a semi-mathematical and equilibrium model that predicts site-specific toxicity of cationic metals such as Cd, Cu, Ni, Pb, and Zn [10–13]. An et al. [14] recently developed a BLM for inorganic arsenate (As(V)), which is a predominant species in surface water and surface soil porewater. This BLM can effectively predict As(V) toxicity to the bioluminescence of \textit{Aliivibrio fischeri} considering phosphate competition (toxicity alleviation) and As species alteration by pH. The sensitivity of the species tested can be considered using species sensitivity distribution (SSD), which is the cumulative distribution of toxicity of a single compound to a set of species that constitutes a community [15]. SSD can estimate the 5% hazardous concentration (HC5), which can be defined as the total dissolved concentration of the contaminant that protects 95% of the species in the ecosystem.

To simultaneously overcome site-specificity and species-specificity while assessing ecotoxicity, thereby deriving a safe concentration (e.g., HC5) in the ecosystem by combining BLM and SSD, the full SSD bioavailability normalization approach [16–19] can be used. In this approach, HC5 is estimated using an SSD obtained from the entire normalized (i.e., site-specific physicochemical properties that can affect the toxicity of metals or metalloids...
determined using the BLM) ecotoxicity data set. This approach also assumes that some species likely share the BLM parameters, that is, conditional binding constants between free metal ions or major cations and active binding sites (ABSs; i.e., biotic ligand) at the cell membrane surface of the organism [19]. This assumption has been supported by the fact that BLMs for Cu, Ni, and Zn can be extrapolated across species [16–18,20].

To the best of our knowledge, the BLM for As(V) only considers the effect of phosphate competition on A. fischeri, as suggested in a previous study [14], and not on individual species. Moreover, evidence supporting extrapolation with respect to the BLM for As(V) is still scarce. Therefore, in this study, the feasibility of interspecies extrapolation was tested to extend the BLM for As(V) to terrestrial plants.

The phenomenon through which major cations alleviate metal toxicity can be explained in terms of BLM, as well as the concept of cell plasma membrane (PM) electrical potential ($\Psi_0$) [21–23]. The root cell PM surface, which is negatively charged, attracts cations in the bulk-phase medium. The addition of major cations, such as Ca$^{2+}$, to the solution increases $\Psi_0$ and decreases the cationic free metal ions moving to the cell PM surface via electrostatic attraction [21]. In contrast, As(V), which is normally present as H$_2$AsO$_4^-$ and HAsO$_4^{2-}$, can accumulate on the PM surface. Wang et al. [22] revealed a significant increase in As(V) toxicity to wheat (Triticum aestivum) with increasing Ca$^{2+}$ activity in the bulk solution. This was because the increase in Ca$^{2+}$ activity from 0.2 to 2.5 mM in the solution increased $\Psi_0$ from $-40$ to $-20$ mV, resulting in an increased accumulation of H$_2$AsO$_4^-$ and HAsO$_4^{2-}$ on the root cell PM surface of T. aestivum. However, such an increase in As(V) toxicity cannot be explained using the current BLM for As(V) [14].

The aim of this study was to confirm that the BLM for As(V), which was originally developed using A. fischeri, can predict As(V) toxicity to terrestrial plant species (i.e., Hordeum vulgare and T. aestivum) by optimizing the inherent sensitivity (IS). In this regard, the feasibility of interspecies extrapolation of BLM for As(V) was evaluated, and the predicted and measured 50% effective As(V) activities (EC$_{50}$(HAsO$_4^{2-}$)) were compared to assess the predictability of BLM for As(V) after adjusting for the inherent sensitivity (IS) of plants. Furthermore, a methodology was established to incorporate the concept of $\Psi_0$ into the extrapolated BLM for As(V) for considering the increased toxicity of As(V) by Ca$^{2+}$.

2. Materials and Methods

2.1. Toxicity Test

The root growth inhibition test using barley H. vulgare was performed according to ISO guideline 11269-1 [24] with some modifications. Seeds were sterilized in a 5% NaOCl solution (Daejung, Korea) for 10 min and rinsed three times with deionized (DI) water, with a resistance of 18.2 M$\Omega$ cm (Milli-Q, Millipore, Bedford, MA, USA). The seeds were then germinated on filter paper moistened with DI water for 36 h at room temperature (25°C) in complete darkness. Each germinated seed was fixed on a polyethylene sheet that floated on the medium surface. For each medium, a toxicity test was conducted, comprising seven treatments (control + six different concentrations of As(V)). Each treatment was performed using six germinated seeds in a polyethylene beaker filled with 500 mL of the test medium as a hydroponic culture. All test media were incubated for 5 days at 20°C with a light:dark cycle of 16:8 in a growth chamber (E15, Conviron, Winnipeg, Canada). The toxicity of As(V), measured as the relative root elongation (RRE), after exposure to a given concentration of As(V) was calculated [25] as follows:

$$\text{RRE} = 100 \times \frac{L_{f,t} - L_{i,t}}{L_{f,c} - L_{i,c}}$$

where RRE is the relative root elongation (%), $L_{f,t}$ is the average root length (mm) of each treatment after 5 days of exposure, $L_{i,t}$ is the average root length (mm) of each treatment at the initial time (t = 0), $L_{f,c}$ is the average root length (mm) of the control after 5 days of exposure, and $L_{i,c}$ is the average root length (mm) of the control at the initial time.
The changes in the RRE in response to the changes in As(V) concentrations or activities were fitted to a sigmoidal dose–response curve (Equation (2)) to calculate the 50% effective concentration (EC50).

\[ y = y_0 + \frac{a}{1 + e^{-\frac{x - x_0}{b}}} \]  

where y is the observed RRE; x is the natural logarithm of the exposed As(V) concentration or activity; a, b, x₀, and y₀ are the fitting parameters that determine the shape of the dose–response curve.

2.2. Effect of Ca²⁺ Concentration on Arsenate Toxicity to H. vulgare

All reagents used in this study were of analytical grade. DI water was used throughout. A stock solution of As(V) (10 mM) was prepared by dissolving Na₂HAsO₄·7H₂O (98.0–102%, Sigma–Aldrich) in DI water and stored in the dark at 4 °C prior to use. The As(V) concentrations in each test medium (i.e., in direct contact with H. vulgare) were 0, 0.5, 2, 10, 50, 200, and 1000 µM. The pH of each test medium was adjusted to 7 ± 0.05 using 3.6 mM 3-[N-morpholino] propanesulfonic acid (MOPS, >99.5%, Sigma–Aldrich), by adding 1 M NaOH or 1 M HCl. Background concentrations of major cations and anions in each test medium were as follows: Ca²⁺ = 0.2 mM, Mg²⁺ = 0.05 mM, K⁺ = 0.2 mM, Na⁺ = 1 mM, PO₄³⁻ = 0.05 mM, SO₄²⁻ = 0.05 mM, and NO₃⁻ = 0.2 mM. The independent effect of Ca²⁺ on As(V) toxicity to H. vulgare was assessed at pH 7 in 3.6 mM MOPS buffer. Each set comprised a series of solutions with five different Ca²⁺ concentrations (i.e., 0.2, 1, 5, 10, and 20 mM).

2.3. Chemical Analysis

The concentrations of major cations and total As were determined via inductively coupled plasma atomic emission spectrometry (ICP-AES) (ICAP 7400 DUO, Thermo Scientific, Waltham, MA, USA). The concentrations of the major anions were determined via ion chromatography (DX500, Dionex, Sunnyvale, CA, USA). Preliminary tests revealed that the measured concentrations did not differ significantly from their nominal values within 5% variability. Since the salt of pentavalent arsenate (i.e., Na₂HAsO₄·7H₂O) was dissolved in DI water to prepare As(V) stock solution in this study, the total As concentration determined using ICP-AES was considered as the concentration of As(V). In order to validate this assumption, the test medium containing 50 µM As was stored 5 days, and then analyzed using high performance liquid chromatography (1260 In-finity LC system, Agilent Technologies, Santa Clara, CA, USA) linked to inductively coupled plasma mass spectrometry (7700x, Agilent Technologies, Santa Clara, CA, USA). It was confirmed that As(V) species did not alter significantly within 5% variability.

Visual MINTEQ 3.1 [26] was used to calculate As(V) activities (H₂AsO₄⁻ and HAsO₄²⁻) and the major cations/anions in the tested solution. Temperature, pH, ion concentrations, and partial pressure of CO₂ (P_CO₂ = 0.00038 atm) were the input data. The analysis of variance (ANOVA) was performed using Microsoft Excel 2010 Analysis ToolPak to evaluate whether Ca²⁺ significantly (p < 0.05) affected As(V) toxicity.

2.4. Calculation of the Electrical Potential of Root Cell PM Surface

To calculate Ψ₀ and the corresponding ion activities at the PM surface, the Gouy–Chapman–Stern (GCS) model [27] was used. The GCS model comprises two parts: (i) electrostatic theory (Gouy–Chapman theory) and (ii) interaction between ions and the PM surface (Stern portion). The calculation process is described as follows:

First, the actual PM surface charge density (σ, unit: C/m²) can be described using Equation (3):

\[ \sigma^2 = 2\varepsilon\varepsilon_0 RT \sum_i [F_i]_b \exp \left( \frac{-Z_i F \Psi_0}{RT} \right) - 1 \]  

where \( 2\varepsilon\varepsilon_0 RT = 0.00345 \), when the concentrations are expressed in units of mol/L at 25 °C; \( \varepsilon \) is the dielectric constant of water; \( \varepsilon_0 \) is the permittivity of vacuum; \( F, R, \) and \( T \) are the...
Faraday constant, gas constant, and absolute temperature, respectively; \([I^Z]_b\) is the free ion concentration in the bulk solution; \(Z_i\) is the charge of the \(i\)th ion.

Second, if the PM surface binding sites comprising negatively charged sites (\(R^−\)) and neutral sites (\(P^0\)) are occupied by ions (\(I^Z\)), the PM surface species and their equilibrium reactions can be represented as Equations (4) and (5):

\[
K_{RI} = \frac{[RI^{Z-1}]}{[R^-][I^Z]_0} \quad (4)
\]

\[
K_{PI} = \frac{[PI^{Z}]}{[R^0][I^Z]_0} \quad (5)
\]

where \([R^-], [P^0], [RI^{Z-1}], \text{and } [PI^{Z}]\) denote the PM surface densities expressed in mol/m²; and \([I^Z]_0\) denotes the concentration of the unbound \(I^Z\) ion at the PM surface.

Finally, \(\sigma\) in the Graham equation (Equation (3)) can be expressed by Equation (6) as follows:

\[
\sigma = \left\{-[R^-] + \sum_i(Z_i - 1)\left[RI^{Z-1}\right] + \sum_i(Z_i)\left[PI^{Z}\right] \right\}F \quad (6)
\]

To calculate the values of \([I^Z]_0\), the Boltzmann equation (Equation (7)) can be used:

\[
[I^Z]_0 = [I^Z]_b \exp\left(\frac{-ZZF\Psi_0}{RT}\right) \quad (7)
\]

To determine \(\Psi_0\), the trial values were assigned to \(\Psi_0\) in Equations (3)–(7) until the values of \(\sigma\) computed in Equations (3)–(6) converge. The equilibrium constants and total surface densities of the binding sites R and P in Equations (4) and (5) were previously determined and are available from Kopittke et al. [27].

2.5. Interspecies Extrapolation of BLM

The BLM for predicting As(V) toxicity to \(A. fischeri\) [14] is presented in Equation (8):

\[
EC_{50}\left\{HAsO_4^{2-}\right\} = f_{mix}^{50\%} \left\{1 + K_{H2PO4}^{K_{XH2PO4}}\left[H_2PO_4^-\right] + K_{HPO4^{2-}}^{K_{XHPO4}}\right\}
\]

\[
\left(1 - f_{mix}^{50\%}\right) \left[K_{H2AsO_4^{2-}}^{K_{XA}}\left[H^+\right] + K_{HAsO_4}\right]
\]

where \(K_{XA}\) is the conditional binding constant for As(V) or phosphate bound to the ABS (L/mol); \([A^{n-}]\) is the activity of As(V) or phosphate (mol/L) in the bulk solution; \(K_{As}\) is the acid dissociation constant of As(V) (\(pK_{As} = 6.76\)); \(f_{mix}^{50\%}\) is the ABS required to be occupied by As(V) (both \(H_2AsO_4^-\) and \(HAsO_4^{2-}\)) for inducing 50% toxicity [14]. The BLM parameters, including the conditional binding constants and \(f_{mix}^{50\%}\), are presented in Table 1a.

\[
EC_{50_{site}} = IS \times EM_{site} \quad (9)
\]

\[
IS = \frac{EC_{50_{test}}}{EM_{test}} = \frac{f_{mix}^{50\%}}{1 - f_{mix}^{50\%}} \quad (10)
\]

\[
EM = \frac{1 + K_{H2PO4}^{K_{XH2PO4}}\left[H_2PO_4^-\right] + K_{HPO4^{2-}}^{K_{XHPO4}}\right\}
\]

\[
\left(\frac{K_{H2AsO_4^{2-}}^{K_{XA}}\left[H^+\right] + K_{HAsO_4}}{K_{H2PO4}^{K_{XH2PO4}}}\right) \quad (11)
\]
Table 1. (a) Conditional binding constants of the biotic ligand model (BLM) for inorganic arsenate (As(V)) obtained from \textit{A. fischeri} [14] and (b) calculated inherent sensitivity ($f_{50\%}$) of \textit{A. fischeri} and terrestrial plants.

| Conditional binding constant | $\log K_{XH2AsO4}$ | $\log K_{XHAsO4}$ | $\log K_{XH2PO4}$ | $\log K_{XHPO4}$ |
|-----------------------------|---------------------|--------------------|--------------------|--------------------|
| Value (L/mol)               | 3.067               | 4.802              | 3.424              | 4.588              |

(b) Inherent sensitivity of \textit{A. fischeri} and terrestrial plants

| Species                  | Toxic endpoint                  | $f_{50\%}$ (dimensionless) |
|--------------------------|--------------------------------|-----------------------------|
| \textit{Aliivibrio fischeri} | 5 min bioluminescence inhibition | 0.616 $^b$                 |
| \textit{Hordeum vulgare}  | 5 days relative root elongation  | 0.626 $^{c,d}$             |
| \textit{Triticum aestivum} | 2 days relative root elongation | 0.015 $^{c,e}$             |

$^a$ Conditional binding constants for As(V) ($\text{HAsO}_4^{2-}$ and $\text{H}_2\text{AsO}_4^{−}$) or phosphate ($\text{HPO}_4^{2-}$ and $\text{H}_2\text{PO}_4^{−}$) bound to active binding sites on the cell membrane surface of \textit{A. fischeri}; $^b$ obtained from An et al. [14]; $^c$ Calculated in this study; $^d$ Toxicity data obtained from An et al. [14]; $^e$ Toxicity data obtained from Wang et al. [22].

The site-specific EC50 of \textit{H. vulgare} was calculated by multiplying the IS of the tested species and the environmental modulator (EM) [19]. IS was derived from the measured EC50 and chemical analysis data of the media (i.e., phosphate activities and pH) in toxicity tests. Hence, to determine the IS (corresponding $f_{50\% \text{mix}}$ value), a single EC50 of \textit{H. vulgare} for As(V) obtained from the toxicity test was required.

3. Results and Discussion

3.1. Effect of Increasing Ca$^{2+}$ Concentrations on As(V) Toxicity

The changes in the EC$_{50}$(HAsO$_4^{2−}$) values with varying Ca$^{2+}$ concentrations are shown in Figure 1. The EC$_{50}$(HAsO$_4^{2−}$) of \textit{H. vulgare} significantly decreased from 45.1 ± 4.34 to 15.0 ± 2.60 µM as the Ca$^{2+}$ concentrations increased from 0.2 to 20 mM (Figure 1a). However, the EC$_{50}$(HAsO$_4^{2−}$) of \textit{A. fischeri}, which was the species used for developing the original BLM for As(V), was not significantly altered, although the Ca$^{2+}$ concentrations increased from 0 to 25 mM (Figure 1b) [14]. This may be due to the sharp increase in $\Psi_0$ of \textit{H. vulgare} from $−53.8$ to $−3.3$ mV. However, only a slight increase in $\Psi_0$ from $−13.7$ to $−0.4$ mV was obtained for \textit{A. fischeri} because the experimental solution containing 0.342 M NaCl was completely used for the osmotic pressure control in the toxicity test using \textit{A. fischeri} and, thus, the solution already had a considerable number of positive ions. As reported in previous studies [21–23], an increase in $\Psi_0$ accumulates more $\text{H}_2\text{AsO}_4^{−}$ and $\text{HAsO}_4^{2−}$, thereby increasing As(V) toxicity. The EC$_{50}$(HAsO$_4^{2−}$) of \textit{H. vulgare} decreased 3.02 times, indicating its significance for more accurate prediction and evaluation of As(V) toxicity; however, it cannot be obtained using the current BLM for As(V).
A negative linear relationship (slope = −0.615, R² = 0.771) between the calculated Ψ₀ and measured EC₅₀[HAsO₄²⁻] of H. vulgare is shown in Figure 2a. Wang et al. [22] revealed a similar trend in wheat T. aestivum (slope = −0.003, R² = 0.729; Figure 2b). The slopes of the linear relationships between the calculated Ψ₀ and measured EC₅₀[HAsO₄²⁻] were different owing to the differences in species sensitivity to As(V). Therefore, the normalized EC50, which can be defined as the measured EC₅₀[HAsO₄²⁻] divided by the y-axis intercept of the negative linear relationship (i.e., a situation where Ψ₀ = 0), was used to quantify the effect of Ψ₀ when compensating for species sensitivity (Figure 2c). Herein, the same negative linear relationship (slope = −0.055, R² = 0.742) was obtained for both species (H. vulgare and T. aestivum). This indicates that the influence of Ψ₀ is not a plant-specific property but a physicochemical property based on the attraction or repulsion of As(V) with the PM surface. Consequently, the effect of Ψ₀ could be considered in a similar way for various plants that have a negatively charged PM surface.

Figure 1. Measured EC₅₀ values expressed as HAsO₄²⁻ activity (left y-axis) obtained from (a) root elongation tests using Hordeum vulgare and (b) bioluminescence inhibition tests using Aliivibrio fischeri (data obtained from An et al. [14]). Calculated cell membrane surface electrical potential (Ψ₀) (right y-axis) for (a) H. vulgare and (b) A. fischeri with varying Ca²⁺ concentrations. Solid bars and error bars indicate the EC50 values and their standard deviations (n = 3), respectively. Open circles represent the calculated Ψ₀ values. The p-value of the ANOVA test was >0.05, indicating that Ca²⁺ addition did not significantly affect As(V) toxicity.
calculated PM surface electrical potential (Ψ0). Normalized EC50 calculated by dividing the measured EC50[HAsO4−2]− by the y-axis intercept (i.e., a situation where Ψ0 = 0) of the negative linear relationship of each species presented in Figure 2a,b is also plotted against Ψ0 (c) to quantify the effect of Ψ0 by eliminating species sensitivity. The solid line represents the linear regression curve.

3.2. Interspecies Extrapolation of BLM to Predict As(V) Toxicity to H. vulgare and T. aestivum

To extrapolate the BLM originally developed from A. fischeri to barley H. vulgare, the IS (f50mix%) had to be determined first. The EC50 value (i.e., EC50[HAsO4−2]) of H. vulgare was 45.1 ± 4.34 μM, when the concentrations of major cations and anions were set to background levels by adjusting the pH to 7. Using the calculated EM value (2.70 × 10−5 M) in Equation (11), the IS and f50mix0% were determined to be 1.675 and 0.626, respectively (Table 1b). The calculated IS of H. vulgare was similar to that of A. fischeri (f50mix0% = 0.616) (Table 1b). The IS of wheat T. aestivum was also calculated to assess its applicability to other terrestrial plants. Wang et al. [22] reported the EC50 values of T. aestivum (toxic endpoint = root elongation change over 48 h) in hydroponic cultures. The calculated IS (f50mix%) of T. aestivum was 0.015 (Table 1b), implying that the sensitivity to As(V) is considerably higher in T. aestivum than in H. vulgare.

The extrapolated BLM to predict As(V) toxicity to H. vulgare and T. aestivum was validated by comparing the measured EC50 values with the predicted EC50 values (Figure 3). To analyze the effect of Ψ0 described in Section 3.1, the slopes of the negative linear relationships (p) in Figure 2a,b were used following Equation (12), when EC50 was predicted. 

\[
\text{EC50} \left\{\text{HAsO}_4^{2-}\right\}_\text{site} = \text{EC50}\left\{\text{HAsO}_4^{2-}\right\}_\text{ini} + p(\Psi_0,\text{site} - \Psi_0,\text{ini})
\]  

(12)

where EC50[HAsO4−2]site is the value indicating the effect of Ψ0 at a site; EC50[HAsO4−2]ini is the predicted EC50 value from the extrapolated BLM; Ψ0,site is the PM surface electrical potential calculated from the chemical analysis of water at the site; Ψ0,ini is the PM surface electrical potential calculated from the chemical analysis of water in the toxicity test solution when determining the EC50 value.
Figure 3. Comparison of measured EC\textsubscript{50}{\{HAsO\textsubscript{4}^{2-}\}} and predicted EC\textsubscript{50}{\{HAsO\textsubscript{4}^{2-}\}} by the extrapolated BLM considering Ψ\textsubscript{0}. Open and closed circles indicate the EC\textsubscript{50}{\{HAsO\textsubscript{4}^{2-}\}} values for \textit{T. aestivum} and \textit{H. vulgare}, respectively. The solid line represents a perfect match between the measured and predicted EC50 values. The dashed lines indicate the difference between the measured and predicted values within an order of magnitude.

The model-predicted EC\textsubscript{50}{\{HAsO\textsubscript{4}^{2-}\}} and experimentally determined EC\textsubscript{50}{\{HAsO\textsubscript{4}^{2-}\}} exhibited a strong linear relationship within an order of magnitude (Figure 3). This demonstrated that although only IS is considered, site-specific As(V) toxicity to terrestrial plants can be predicted via the chemical analysis of water or porewater with significant accuracy. Thus, the BLM parameters for each plant species are not required, that is, the conditional binding constants between As(V) or phosphate and ABs of the cell membrane are the same for both species; however, only one EC50 should be used for sensitivity correction. To achieve improved prediction accuracy, the relationship between EC50 values and Ψ\textsubscript{0} must be derived.

As(V) is assimilated into cells by the existing orthophosphate transporters in both prokaryotes and eukaryotes [28,29], because As(V) is a chemical analog of orthophosphate. This suggests that there are no severe problems in directly using the BLM parameters (e.g., conditional binding constants: \(K_{XH2AsO4}, K_{XHAsO4}, K_{XH2PO4},\) and \(K_{XHPO4}\)) derived from \textit{A. fischeri} to predict As(V) toxicity to terrestrial plants. The BLM for As(V), originally developed for \textit{A. fischeri}, can be extrapolated to terrestrial plants, including \textit{H. vulgare} and \textit{T. aestivum} across, plant kingdoms.

4. Conclusions

In this study, the toxicity caused by As(V) to the terrestrial plants, namely, \textit{H. vulgare} and \textit{T. aestivum}, was predicted using the existing BLM for bioluminescence of \textit{A. fischeri} via interspecies extrapolation. The EC\textsubscript{50}{\{HAsO\textsubscript{4}^{2-}\}} values decreased from 45.1 ± 4.34 to 15.0 ± 2.60 μM as Ca\textsuperscript{2+} concentration increased from 0.2 to 20 mM, owing to the accumulation of H\textsubscript{2}AsO\textsubscript{4}\textsuperscript{−} and HAsO\textsubscript{4}^{2−} on the PM surface. This was successfully predicted in the extrapolated BLM using a linear relationship between Ψ\textsubscript{0} and EC\textsubscript{50}{\{HAsO\textsubscript{4}^{2-}\}}. Consequently, the BLM for As(V) developed for a single bacterium can predict As(V) toxicity to terrestrial plant species among various physicochemical properties by optimizing IS.
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