Data Article

Dataset of the electrochemical potential windows for the Au(hkl)|ionic liquid interfaces defined by the cut-off current densities

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\textbf{A R T I C L E  I N F O}

\textbf{Article history:}
Received 2 November 2021
Revised 11 November 2021
Accepted 12 November 2021
Available online 18 November 2021

\textbf{Keywords:}
Au(hkl)
Electrical double layer
Electrochemical potential window
Interfacial processes
Ionic liquids
Linear sweep voltammetry
Single crystal electrodes
Specific desorption

\textbf{A B S T R A C T}

This data article describes the linear sweep voltammetry (LSV) profiles of five ionic liquids (ILs) at the low-index (hkl) (hkl = 111, 100, and 110) planes of Au. The LSV profiles were recorded at 25 ± 1°C for the Au(hkl)|IL interfaces maintained in a hanging meniscus configuration in an inert Ar atmosphere (with H\textsubscript{2}O and O\textsubscript{2} concentrations being lower than 5 ppm). The width of the electrical double-layer regions (E\textsubscript{dl}) and the electrochemical potential windows (E\textsubscript{pw}) of the ILs were evaluated based on the cut-off current densities (j\textsubscript{cut-off}): ±5, ±10, and ±20 μA cm\textsuperscript{-2} for E\textsubscript{dl} and ±0.1, ±0.5, and ±1.0 mA cm\textsuperscript{-2} for E\textsubscript{pw}. The potential values were calibrated to the redox potential of ferrocene/ferrocenium in each IL. A detailed discussion on the electrochemical behaviors of the ILs on Au(hkl) is provided in the related article “Voltammetric Investigation of Anodic and Cathodic Processes at Au(hkl)|Ionic Liquid Interfaces”, published in the Journal of Electroanalytical Chemistry (Ueda and Yoshimoto, 2021).

DOI of original article: 10.1016/j.jelechem.2021.115691
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https://doi.org/10.1016/j.jelechem.2021.115691
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Specifications Table

| Subject               | Electrochemistry                     |
|-----------------------|--------------------------------------|
| Specific subject area | Surface electrochemistry of ionic liquids (ILs) |
| Type of data          | Table                                |
| How the data were acquired | A CH Instruments potentiostat (Model 610D) was used for linear sweep voltammetry (LSV). The scan rate was 50 mV s⁻¹. LSV was performed for Au(hkl) working electrodes (hkl = 111, 100, and 110) contacted with vacuum-dried ILs at 25 ± 1°C in three-electrode cells with Pt wires as counter and quasi-reference electrodes. Electrochemical measurements for each electrode were conducted in an Ar atmosphere (H₂O and O₂ <5 ppm) in four steps: the electrode potential was (1) swept to the positive direction until the current density reached 20 μA cm⁻², (2) swept to the negative direction until the current density reached ~20 μA cm⁻², (3) swept to the positive direction until the current density reached 1 mA cm⁻², and (4) swept to the negative direction until the current density reached ~1 mA cm⁻². Prior to LSV, the electrode was maintained at ~0.1 V vs. Pt during the holding time of 10 s for (1) and 2 min for (2), (3), and (4). Voltammograms obtained via (1) and (2) were used to evaluate the widths of the electrical double-layer region (εdl), while those obtained via (3) and (4) were used to determine the electrochemical potential windows (Epw). |
| Data format          | Raw                                  |
| Description of data collection | Raw LSV data were exported to Microsoft Excel to plot the voltammograms and analyze the Edl and Epw of ILs on Au(hkl). The cut-off current densities (jcut-off) for Edl were ±5, ±10, and ±20 μA cm⁻², whereas jcut-off for Epw were ±0.1, ±0.5, and ±1.0 mA cm⁻². The anodic and cathodic limits of Edl and Epw were determined based on the jcut-off values. |
| Data source location | • Institution: Kumamoto University  |
|                      | • City/Region: Kumamoto             |
|                      | • Country: Japan                    |
|                      | • Latitude and longitude (and GPS coordinates, if possible) for collected samples/data: 32.81291, 130.72578 |
| Data accessibility   | Repository name: Mendeley Data      |
| Data identification number (DOI): | http://doi.org/10.17632/tv4cm845wv.1 [2] |
| Direct URL to data   | http://doi.org/10.17632/tv4cm845wv.1 |
| Related research article | [1] H. Ueda, S. Yoshimoto, Voltammetric Investigation of Anodic and Cathodic Processes at Au(hkl)|IL Ionic Liquid Interfaces, J. Electroanal. Chem. 900 (2021) 115691. |

Value of the Data

- The electrochemical data reported herein are valuable because they can provide fundamental information on Au(hkl)|IL interfaces for electrochemical studies.
- Electrochemists can benefit from these data because it will aid them in selecting appropriate potential ranges for studies using Au(hkl)|IL interfaces. For instance, the decomposition of ILs can be significantly reduced by limiting the potential of the Au(hkl) working electrode to within the Edl.
- These data can be used to gain further insights into the origin of each anodic or cathodic process occurring at Au(hkl)|IL interfaces by means of microscopic or spectroscopic techniques such as scanning tunneling microscopy [3–5] and differential electrochemical mass spectroscopy [6].
These data were analyzed at different $j_{cut-off}$ values, thereby providing a basis for a fair comparison of $E_{dl}$ and $E_{pw}$ between different electrode|IL interfaces.

In addition, these data are useful for identifying potential regions in which ILs exhibit nearly ideal capacitive behavior. Such potential regions are essential for ensuring the accuracy of microcalorimetric measurements [7] and amperometric sensors using ILs [8].

1. Data Description

This data article summarizes the LSV profiles, $E_{dl}$, and $E_{pw}$ of the Au(hkl)|IL interfaces. The chemical structures of the five ILs are shown in Fig. 1. The raw data of all LSV profiles and Tables can be found in the repository (see “Data accessibility” in the Specifications Table) [2].

Fig. 2 shows the LSV profiles of 1-butyl-3-methylimidazolium hexafluorophosphate ([C$_4$mim][PF$_6$]) on Au(hkl). In the enlarged voltammograms (the dotted lines), two reductive peaks at −1.11 and −1.63 V vs. Fc/Fc$^+$ appeared for the Au(111) surface, whereas the reductive peaks were unclear for Au(100) and Au(110). As shown using the solid lines, the oxidation onset potential of [C$_4$mim][PF$_6$] on Au(110) was more negative than that on Au(111) and Au(100).

![Fig. 1. The chemical structures of the ILs.](image)

![Fig. 2. LSV profiles of [C$_4$mim][PF$_6$] on Au(hkl) recorded at the scan rate of 50 mV s$^{-1}$.](image)
Similarly, the reduction onset potential of [C₄mim][PF₆] on Au(110) was more positive than that on the other crystal faces. In addition, an oxidation process was observed prior to a massive increase in the oxidation current density of [C₄mim][PF₆] on Au(110).

Fig. 3 depicts the LSV profiles of N-butyl-N-methylpyrrolidinium bis(trifluoromethylsulfonyl)amide ([C₄mppyrr][Tf₂N]) on Au(hkl). The enlarged voltammogram at the anodic scan of [C₄mppyrr][Tf₂N] on Au(111) was nearly identical to that of Au(110), except for the presence of a shoulder peak at approximately 1.40 V vs. Fc/Fc⁺. Furthermore, both crystal faces generated nearly identical voltammetric shapes during the cathodic scan prior to reaching −20 μA cm⁻². Conversely, in the Eₜ₁ region, the anodic and cathodic processes on Au(100) were milder than those on the other crystal faces. In the voltammograms recorded to determine the Eₚ𝑤 (the solid lines), the voltammetric shape between 1.20 V vs. Fc/Fc⁺ and Eₚ𝑤 than −1.50 V vs. Fc/Fc⁺ were dependent on the crystallographic orientation of gold.

Fig. 4 shows the LSV profiles of tributylmethylammonium bis(trifluoromethylsulfonyl)amide ([N₁,4,4,4][Tf₂N]) on Au(hkl). In the Eₜ₁ region (indicated using the dotted lines), a small anodic process was observed at approximately 0.1 V vs. Fc/Fc⁺ solely for the Au(111) surface. All crystal faces exhibited a cathodic peak at approximately −1.00 V vs. Fc/Fc⁺. The absolute value of the peak current density for this cathodic process was evaluated to be in the following order: Au(100) < Au(111) < Au(110). In the Eₚ𝑤 region (indicated using the solid lines), the voltammetric shapes of [N₁,4,4,4][Tf₂N] on Au(111) and Au(110) were nearly identical, except for the difference in the peak current density of the cathodic process at approximately −1.50 V vs. Fc/Fc⁺. In contrast, the absolute value of the current density (|j|) for Au(100) tended to be the lowest over the entire potential range.

Fig. 5 illustrates the LSV profiles of 1-butyl-3-methylimidazolium iodide ([C₄mim][I]) on Au(hkl). As indicated using the dotted lines, the order of the onset oxidation potential was evaluated as Au(111) < Au(100) < Au(110). The cathodic peak potentials were −0.92 V vs. Fc/Fc⁺ for Au(111), −1.35 V vs. Fc/Fc⁺ for Au(100), and −1.29 V vs. Fc/Fc⁺ for Au(110). As indicated using the solid lines, no significant differences in the voltammetric shape at the anodic scan were identified between Au(hkl). During the cathodic scan, a voltammetric peak generated by the from the reductive desorption of the iodine adlayer on Au(hkl) appeared at approximately −2.20 V vs.
Fig. 4. LSV profiles of [N_{1,4,4,4}]Tf_2N on Au(hkl) recorded at the scan rate of 50 mV s\(^{-1}\).

Fig. 5. LSV profiles of [C_4mim][I] on Au(hkl) recorded at the scan rate of 50 mV s\(^{-1}\).

Fc/Fc^+ [1,9]. Furthermore, the \(|j|\) value during the \(E_{pw-CL}\) determining reduction was lowest for Au(110).

Fig. 6 shows the LSV profiles of 1-hexyl-3-methylimidazolium iodide ([C_6mim][I]) on Au(hkl). The enlarged voltammograms of Au(hkl) in the anodic scan were nearly identical. The cathodic peak appeared at \(-1.40\) V vs. Fc/Fc^+ for Au(111) and Au(100). As for the Au(110) surface, the two cathodic peaks were observed at \(-0.97\) and \(-1.64\) V vs. Fc/Fc^+. In the \(E_{pw}\) region (indicated using the solid lines), all the voltammograms exhibited the maximum \(|j|\) values during the anodic and cathodic scans, which were between 0.5 and 1.0 mA cm\(^{-2}\) and in the following order: Au(111) \(<\) Au(100) \(<\) Au(110). The voltammetric shape for the reductive desorption of the iodine adlayer at approximately \(-2.20\) V vs. Fc/Fc^+ was dependent on the crystallographic orientation of gold.
The $E_{dl}$ of [C₄mim][PF₆] on Au(hkl) was estimated to be Au(110) < Au(100) < Au(111) and $E_{dl}$ for [C₄mim][PF₆] followed the order: Au(111) < Au(100) < Au(110). Therefore, the $E_{dl}$ value of [C₄mim][PF₆] was evaluated as Au(110) < Au(100) < Au(111). Similarly, the $E_{pw}$ of [C₄mim][PF₆] followed the order: Au(110) < Au(100) < Au(111), and the $E_{pw}$ of [C₄mim][PF₆] was regarded as Au(111) < Au(100) < Au(110), suggesting that the electrochemical stability of [C₄mim][PF₆] on the electrode surface followed the order: Au(110) < Au(100) < Au(111).

Tables 1 and 2 summarize the $E_{dl}$ and $E_{pw}$ of [C₄mim][PF₆] on Au(hkl), respectively. The $E_{dl}$ of [C₄mim][PF₆] was estimated to be Au(110) < Au(100) < Au(111) and $E_{dl}$ for [C₄mim][PF₆] followed the order: Au(111) < Au(100) < Au(110). Therefore, the $E_{dl}$ value of [C₄mim][PF₆] was evaluated as Au(110) < Au(100) < Au(111). Similarly, the $E_{pw}$ of [C₄mim][PF₆] followed the order: Au(110) < Au(100) < Au(111), and the $E_{pw}$ of [C₄mim][PF₆] was regarded as Au(111) < Au(100) < Au(110), suggesting that the electrochemical stability of [C₄mim][PF₆] on the electrode surface followed the order: Au(110) < Au(100) < Au(111).

Fig. 6. LSV profiles of [C₄mim][I] on Au(hkl) recorded at the scan rate of 50 mV s⁻¹.
Table 2
The $E_{\text{pw-AL}}$, $E_{\text{pw-CL}}$, and $E_{\text{pw}}$ of [C₄mim][PF₆] on Au(hkl) at different $j_{\text{cut-off}}$ values.

| $j_{\text{cut-off}}/\pm$mA cm⁻² | Crystal face | $E_{\text{pw-CL}}/V$ vs.Fc/Fc⁺ | $E_{\text{pw-AL}}/V$ vs. Fc/Fc⁺ | $E_{\text{pw}}/V$ |
|-------------------------------|-------------|--------------------------------|-------------------------------|----------------|
| 0.1                           | Au(111)     | 2.80                           | 1.80                          | 4.60           |
|                               | Au(100)     | 2.79                           | 1.78                          | 4.57           |
|                               | Au(110)     | 2.69                           | 1.75                          | 4.44           |
| 0.5                           | Au(111)     | 2.92                           | 1.99                          | 4.91           |
|                               | Au(100)     | 2.89                           | 1.96                          | 4.85           |
|                               | Au(110)     | 2.86                           | 1.95                          | 4.81           |
| 1.0                           | Au(111)     | 2.99                           | 2.05                          | 5.05           |
|                               | Au(100)     | 2.94                           | 2.01                          | 4.94           |
|                               | Au(110)     | 2.92                           | 2.00                          | 4.91           |

Table 3
The $E_{\text{dl-AL}}$, $E_{\text{dl-CL}}$, and $E_{\text{dl}}$ of [C₄mpyr][Tf₂N] on Au(hkl) at different $j_{\text{cut-off}}$ values.

| $j_{\text{cut-off}}/\pm$μA cm⁻² | Crystal face | $E_{\text{dl-CL}}/V$ vs. Fc/Fc⁺ | $E_{\text{dl-AL}}/V$ vs. Fc/Fc⁺ | $E_{\text{dl}}/V$ |
|-------------------------------|-------------|--------------------------------|-------------------------------|----------------|
| 5                             | Au(111)     | -1.07                          | 1.09                          | 2.16           |
|                               | Au(100)     | -1.14                          | 1.19                          | 2.33           |
|                               | Au(110)     | -1.07                          | 1.07                          | 2.15           |
| 10                            | Au(111)     | -1.17                          | 1.38                          | 2.55           |
|                               | Au(100)     | -1.23                          | 1.53                          | 2.76           |
|                               | Au(110)     | -1.18                          | 1.33                          | 2.51           |
| 20                            | Au(111)     | -1.23                          | 1.84                          | 3.07           |
|                               | Au(100)     | -1.31                          | 1.85                          | 3.17           |
|                               | Au(110)     | -1.26                          | 1.78                          | 3.04           |

Table 4
The $E_{\text{pw-AL}}$, $E_{\text{pw-CL}}$, and $E_{\text{pw}}$ of [C₄mpyr][Tf₂N] on Au(hkl) at different $j_{\text{cut-off}}$ values.

| $j_{\text{cut-off}}/\pm$mA cm⁻² | Crystal face | $E_{\text{pw-CL}}/V$ vs. Fc/Fc⁺ | $E_{\text{pw-AL}}/V$ vs. Fc/Fc⁺ | $E_{\text{pw}}/V$ |
|-------------------------------|-------------|--------------------------------|-------------------------------|----------------|
| 0.1                           | Au(111)     | -3.13                          | 2.26                          | 5.40           |
|                               | Au(100)     | -3.14                          | 2.30                          | 5.44           |
|                               | Au(110)     | -3.13                          | 2.30                          | 5.43           |
| 0.5                           | Au(111)     | -3.34                          | 2.40                          | 5.74           |
|                               | Au(100)     | -3.33                          | 2.40                          | 5.73           |
|                               | Au(110)     | -3.36                          | 2.41                          | 5.77           |
| 1.0                           | Au(111)     | -3.42                          | 2.46                          | 5.87           |
|                               | Au(100)     | -3.40                          | 2.45                          | 5.85           |
|                               | Au(110)     | -3.43                          | 2.45                          | 5.88           |

the $j_{\text{cut-off}}$ values, whereas the Au(100) surface afforded the widest $E_{\text{pw}}$. Specifically, when $j_{\text{cut-off}}$ was $\pm 1.0$ mA cm⁻², the $E_{\text{pw}}$ was 6.99 V for the reaction on the Au(100) surface, whereas the $E_{\text{pw}}$ was 6.54 V for that on Au(111), and 6.60 V in the case of the Au(110) surface.

Tables 7 and 8 list the $E_{\text{dl}}$ and $E_{\text{pw}}$ of [C₄mim][I] on Au(hkl). The order of the $E_{\text{dl}}$ values was dependent on $j_{\text{cut-off}}$, which was due to the difference in the current density measured during the cathodic process. Similarly, the order of the $E_{\text{pw}}$ values at $j_{\text{cut-off}} = \pm 0.1$ mA cm⁻² was affected by the peak current density for reductive desorption of the iodine adlayer. At $j_{\text{cut-off}} = 0.5$ or 1.0 mA cm⁻², $E_{\text{pw}}$ was influenced solely by the cathodic decomposition of [C₄mim][I] and the anodic reaction involving the complexation of gold with iodide, resulting in the following order of $E_{\text{pw}}$: Au(100) < Au(111) < Au(110).
### Table 5
The $E_{dl\text{-AL}}$, $E_{dl\text{-CL}}$, and $E_{dl}$ of $[N_{1,4,4,4}]^2[Tf_2N]$ on Au(hkl) at different $j_{cut\text{-off}}$ values.

| $j_{cut\text{-off}}/\pm \mu A \text{ cm}^{-2}$ | Crystal face | $E_{dl\text{-CL}}/V$ vs. Fc/Fc$^+$ | $E_{dl\text{-AL}}/V$ vs. Fc/Fc$^+$ | $E_{dl}/V$ |
|------------------------------------------|-------------|-------------------------------|-------------------------------|-----------|
| 5                                       | Au(111)     | –0.82                         | 0.49                          | 1.31      |
|                                          | Au(100)     | –1.49                         | 0.52                          | 2.01      |
|                                          | Au(110)     | –0.80                         | 0.52                          | 1.32      |
| 10                                      | Au(111)     | –1.55                         | 0.60                          | 2.14      |
|                                          | Au(100)     | –1.70                         | 0.63                          | 2.33      |
|                                          | Au(110)     | –1.68                         | 0.63                          | 2.31      |
| 20                                      | Au(111)     | –1.77                         | 0.76                          | 2.53      |
|                                          | Au(100)     | –1.98                         | 0.88                          | 2.87      |
|                                          | Au(110)     | –1.95                         | 0.80                          | 2.75      |

### Table 6
The $E_{pw\text{-AL}}$, $E_{pw\text{-CL}}$, and $E_{pw}$ of $[N_{1,4,4,4}]^2[Tf_2N]$ on Au(hkl) at different $j_{cut\text{-off}}$ values.

| $j_{cut\text{-off}}/\pm mA \text{ cm}^{-2}$ | Crystal face | $E_{pw\text{-CL}}/V$ vs. Fc/Fc$^+$ | $E_{pw\text{-AL}}/V$ vs. Fc/Fc$^+$ | $E_{pw}/V$ |
|------------------------------------------|-------------|-------------------------------|-------------------------------|-----------|
| 0.1                                      | Au(111)     | –3.28                         | 2.02                          | 5.29      |
|                                          | Au(100)     | –3.58                         | 2.10                          | 5.68      |
|                                          | Au(110)     | –3.31                         | 2.03                          | 5.34      |
| 0.5                                      | Au(111)     | –3.85                         | 2.28                          | 6.13      |
|                                          | Au(100)     | –3.97                         | 2.41                          | 6.39      |
|                                          | Au(110)     | –3.87                         | 2.30                          | 6.17      |
| 1.0                                      | Au(111)     | –4.05                         | 2.49                          | 6.54      |
|                                          | Au(100)     | –4.27                         | 2.72                          | 6.99      |
|                                          | Au(110)     | –4.08                         | 2.52                          | 6.60      |

### Table 7
The $E_{dl\text{-AL}}$, $E_{dl\text{-CL}}$, and $E_{dl}$ of $[C_4\text{mim}]^+$ on Au(hkl) at different $j_{cut\text{-off}}$ values.

| $j_{cut\text{-off}}/\pm \mu A \text{ cm}^{-2}$ | Crystal face | $E_{dl\text{-CL}}/V$ vs. Fc/Fc$^+$ | $E_{dl\text{-AL}}/V$ vs. Fc/Fc$^+$ | $E_{dl}/V$ |
|------------------------------------------|-------------|-------------------------------|-------------------------------|-----------|
| 5                                       | Au(111)     | –1.64                         | –0.55                         | 1.09      |
|                                          | Au(100)     | –1.29                         | –0.53                         | 0.76      |
|                                          | Au(110)     | –1.13                         | –0.48                         | 0.65      |
| 10                                      | Au(111)     | –1.94                         | –0.51                         | 1.42      |
|                                          | Au(100)     | –2.03                         | –0.49                         | 1.54      |
|                                          | Au(110)     | –1.97                         | –0.44                         | 1.52      |
| 20                                      | Au(111)     | –2.03                         | –0.47                         | 1.56      |
|                                          | Au(100)     | –2.09                         | –0.46                         | 1.63      |
|                                          | Au(110)     | –2.03                         | –0.42                         | 1.61      |

Tables 9 and 10 show the $E_{dl}$ and $E_{pw}$ of $[C_6\text{mim}]^+$ on Au(hkl). The difference in the current density measured during the cathodic process affected the order of the $E_{dl}$ values at each $j_{cut\text{-off}}$ value. The lowest value of $E_{pw}$ was observed for Au(111), whereas Au(100) and Au(110) exhibited nearly equal values. $E_{pw}$ at $\pm 1.0 \text{ mA cm}^{-2}$ was not measured because the current density did not reach $\pm 1.0 \text{ mA cm}^{-2}$. 
Table 8
The $E_{\text{pw-CL}}, E_{\text{pw-AL}},$ and $E_{\text{pw}}$ of [C₄mim][I] on Au(hkl) at different $j_{\text{cut-off}}$ values.

| $j_{\text{cut-off}}$/±mA cm⁻² | Crystal face | $E_{\text{pw-CL}}$/V vs. Fc/Fc⁺ | $E_{\text{pw-AL}}$/V vs. Fc/Fc⁺ | $E_{\text{pw}}$/V |
|---------|-----------------|-----------------|-----------------|-----------------|
| 0.1     | Au(111)         | -2.15           | -0.41           | 1.74            |
|         | Au(100)         | -2.49           | -0.41           | 2.08            |
|         | Au(110)         | -2.20           | -0.40           | 1.79            |
| 0.5     | Au(111)         | -2.65           | -0.26           | 2.39            |
|         | Au(100)         | -2.64           | -0.26           | 2.38            |
|         | Au(110)         | -2.67           | -0.25           | 2.42            |
| 1.0     | Au(111)         | -2.80           | -0.13           | 2.68            |
|         | Au(100)         | -2.79           | -0.14           | 2.65            |
|         | Au(110)         | -2.86           | -0.11           | 2.74            |

Table 9
The $E_{\text{dl-AL}}, E_{\text{dl-CL}},$ and $E_{\text{dl}}$ of [C₆mim][I] on Au(hkl) at different $j_{\text{cut-off}}$ values.

| $j_{\text{cut-off}}$/±μA cm⁻² | Crystal face | $E_{\text{dl-CL}}$/V vs. Fc/Fc⁺ | $E_{\text{dl-AL}}$/V vs. Fc/Fc⁺ | $E_{\text{dl}}$/V |
|---------|-----------------|-----------------|-----------------|-----------------|
| 5       | Au(111)         | -1.24           | -0.55           | 0.68            |
|         | Au(100)         | -1.30           | -0.54           | 0.76            |
|         | Au(110)         | -0.78           | -0.54           | 0.23            |
| 10      | Au(111)         | -1.98           | -0.52           | 1.46            |
|         | Au(100)         | -2.11           | -0.51           | 1.60            |
|         | Au(110)         | -0.91           | -0.51           | 0.40            |
| 20      | Au(111)         | -2.09           | -0.50           | 1.59            |
|         | Au(100)         | -2.19           | -0.48           | 1.70            |
|         | Au(110)         | -2.09           | -0.48           | 1.60            |

Table 10
The $E_{\text{pw-AL}}, E_{\text{pw-CL}},$ and $E_{\text{pw}}$ of [C₆mim][I] on Au(hkl) at different $j_{\text{cut-off}}$ values.

| $j_{\text{cut-off}}$/±mA cm⁻² | Crystal face | $E_{\text{pw-CL}}$/V vs. Fc/Fc⁺ | $E_{\text{pw-AL}}$/V vs. Fc/Fc⁺ | $E_{\text{pw}}$/V |
|---------|-----------------|-----------------|-----------------|-----------------|
| 0.1     | Au(111)         | -2.61           | -0.47           | 2.14            |
|         | Au(100)         | -2.61           | -0.43           | 2.17            |
|         | Au(110)         | -2.63           | -0.44           | 2.18            |
| 0.5     | Au(111)         | -2.82           | -0.27           | 2.55            |
|         | Au(100)         | -2.94           | -0.14           | 2.80            |
|         | Au(110)         | -2.95           | -0.14           | 2.81            |
| 1.0     | Au(111)         | ND a            | ND a            | ND a            |
|         | Au(100)         | ND a            | ND a            | ND a            |
|         | Au(110)         | ND a            | ND a            | ND a            |

a Not determined because the decomposition current density did not reach ±1.0 mA cm⁻².

2. Experimental Design, Materials and Methods

[C₄mim][PF₆] (Merck, >99.0%), [C₄mpyrr][Tf₂N] (Solvionic, 99.9%), [N₁,4,4,4][Tf₂N] (IoLiTec, >99%), [C₄mim][I] (Kanto Chemical Co. Ltd., >99%), and [C₆mim][I] (Kanto Chemical Co. Ltd., >99%) were used in this study. Detailed information about the water, halide, and alkali metal contents, the presence of other impurities, and the color of each IL is provided in Section I of the Supporting Information in ref. [1]. Following the drying of the ILs in vacuum at approximately 80°C for >6 h, they did not exhibit the cathodic stripping peak of gold oxide originating from a trace amount of water in the ILs [10].
Au(111), Au(100), and Au(110) working electrodes were prepared using Clavilier’s method [11]. The area of the working electrode was 0.065 ± 0.005 cm². The working electrodes and Pt wires were annealed in a hydrogen flame and cooled in air for 1 min. Thereafter, they were placed inside the antechamber of a vacuum-type glove box (UN650F, UNICO Corp.), followed by evacuation for >15 min. The antechamber was refilled with Ar gas until the vacuum gauge reached ~0.1 bar with respect to the atmospheric pressure, whereupon it was re-evacuated. This refill/evacuation cycle was repeated two times. Subsequently, the pressure of the antechamber was increased to atmospheric pressure using Ar gas. The electrodes were transferred from the antechamber to the main room of the glove box, which was maintained at sufficiently low H₂O and O₂ concentrations using a gas recycling purification system (MF-71, UNICO). The working electrodes were contacted with the ILs in three-electrode cells using Pt wires as counter and quasi-reference electrodes.

LSV was conducted while maintaining the contact between the working electrode and IL in a hanging-meniscus configuration. The detailed steps of LSV and data analysis have been explained in “How the data were acquired” and “Description of data collection” in the Specifications Table. E_dl and E_pw were calculated using the following equations:

\[
E_{\text{dl}} = E_{\text{dl-AL}} - E_{\text{dl-CL}} \quad \left( J_{\text{cut-off}} = \pm 5, \pm 10, \text{ or } \pm 20 \, \text{μA cm}^{-2} \right) \tag{1}
\]

\[
E_{\text{pw}} = E_{\text{pw-AL}} - E_{\text{pw-CL}} \quad \left( J_{\text{cut-off}} = \pm 0.1, \pm 0.5, \text{ or } \pm 1.0 \, \text{mA cm}^{-2} \right) \tag{2}
\]

where E_{dl-AL} and E_{pw-AL} are the electrode potentials at which the positive J_{cut-off} values are measured, and E_{dl-CL} and E_{pw-CL} are the electrode potentials at which the negative J_{cut-off} values are measured. The J_{cut-off} values for E_{dl} and E_{pw} were chosen based on previous studies [7,12–15]. The potential values of LSV were referenced to the redox potential of 2 mM ferrocene (Fc) in the corresponding IL, as recommended by IUPAC [16]. The Fc/Fc⁺ redox couple has been used widely to characterize ILs [17–20].

Ethics Statement

Not applicable.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

CRediT Author Statement

Hiroyuki Ueda: Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Validation, Visualization, Writing – original draft, Writing – review & editing; Soichiro Yoshimoto: Data curation, Funding acquisition, Project administration, Supervision, Writing – review & editing.

Acknowledgments

This work was supported by JSPS KAKENHI [grant numbers 19H02560 and 19K22115] and by the IIiNa Interdisciplinary Research Project of the Institute of Industrial Nanomaterials, Kumamoto University. We thank Dr. Katsuhiko Nishiyama at Kumamoto University for helpful discussions. We would like to thank Editage (www.editage.com) for English language editing.
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