



MetalWalls: A classical molecular dynamics software dedicated to the simulation of electrochemical systems

Abel Marin-Lafèche, Matthieu Haefele, Laura Scalfi, Alessandro Coretti, Thomas Dufils, Guillaume Jeanmairet, Stewart K. Reed, Serva Alessandra, Roxanne Berthin, Camille Bacon, et al.

► To cite this version:

Abel Marin-Lafèche, Matthieu Haefele, Laura Scalfi, Alessandro Coretti, Thomas Dufils, et al.. MetalWalls: A classical molecular dynamics software dedicated to the simulation of electrochemical systems. Journal of Open Source Software, 2020, 5 (53), pp.2373. 10.21105/joss.02373 . hal-02975415

HAL Id: hal-02975415

<https://hal.sorbonne-universite.fr/hal-02975415>

Submitted on 22 Oct 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.




MetalWalls: A classical molecular dynamics software dedicated to the simulation of electrochemical systems

Abel Marin-Lafleche¹, Matthieu Haefele¹, Laura Scalfi², Alessandro Coretti^{3, 4}, Thomas Dufils², Guillaume Jeanmairet², Stewart K. Reed⁵, Alessandra Serva², Roxanne Berthin², Camille Bacon², Sara Bonella⁴, Benjamin Rotenberg², Paul A. Madden⁶, and Mathieu Salanne^{1, 2}

1 Maison de la Simulation, CEA, CNRS, Univ. Paris-Sud, UVSQ, Université Paris-Saclay, F-91191 Gif-sur-Yvette, France **2** Sorbonne Université, CNRS, Physico-chimie des Électrolytes et Nanosystèmes Interfaciaux, PHENIX, F-75005 Paris, France **3** Department of Mathematical Sciences, Politecnico di Torino, I-10129 Torino, Italy **4** Centre Européen de Calcul Atomique et Moléculaire (CECAM), École Polytechnique Fédérale de Lausanne, 1015 Lausanne, Switzerland **5** School of Chemistry, University of Leeds, Leeds, UK **6** Department of Materials, University of Oxford, Oxford, UK

DOI: [10.21105/joss.02373](https://doi.org/10.21105/joss.02373)

Software

- [Review](#) 
- [Repository](#) 
- [Archive](#) 

Editor: Richard Gowers 

Reviewers:

- [@mattwthompson](#)
- [@hmacdope](#)

Submitted: 10 June 2020

Published: 25 September 2020

License

Authors of papers retain copyright and release the work under a Creative Commons Attribution 4.0 International License ([CC BY 4.0](https://creativecommons.org/licenses/by/4.0/)).

Summary

Applied electrochemistry plays a key role in many technologies, such as Li-ion batteries, fuel cells, supercapacitors, solar cells, etc. It is therefore at the core of many research programs all over the world. However, fundamental electrochemical investigations remain scarce. In particular, electrochemistry is among the fields for which the gap between theory and experiment is the largest. From the computational point of view, there is no classical molecular dynamics (MD) software devoted to the simulation of electrochemical systems while other fields such as biochemistry or material science have dedicated tools. MetalWalls, a MD code dedicated to electrochemistry, fills this gap. Its main originality is the inclusion of a series of methods which allow a constant electrical potential to be applied to the electrode materials. It also allows the simulation of bulk liquids or solids using the polarizable ion model and the aspherical ion model. MetalWalls is designed to be used on high-performance computers and it has already been employed in a number of scientific publications. It was for example used to study the charging mechanism of supercapacitors (Merlet et al., 2012), nanoelectrowetting (Choudhuri et al., 2016) and water desalination devices (Simoncelli et al., 2018). A typical snapshot of a simulation cell for the latter is shown on [Figure 1](#).

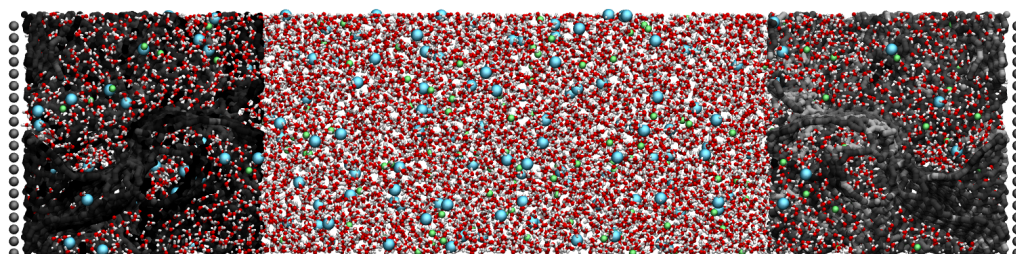


Figure 1: Typical system simulated using MetalWalls. An aqueous NaCl electrolyte is put in contact with two electrified nanoporous carbon electrodes (left: positive electrode, right: negative electrode). Water molecules are in red and white, Cl^- ions in cyan, Na^+ ions in green and the carbon atoms are colored according to their charge (from light grey to black when the charge increases).

Constant potential electrodes

MetalWalls employs most of the standard methods and algorithms available in typical MD packages. Here we discuss only its main originality, which lies in the possibility of simulating materials at constant electric potential instead of constant charge. This is interesting for simulating conductors where the charges are not fixed on the atoms but can spread in the material and adapt to the external medium. In MetalWalls, this is done by allowing the charge on the material atoms to fluctuate under the condition of constant potential. The method was first introduced by Siepmann and Sprik (Siepmann & Sprik, 1995). It was later adapted to the case of 2D periodic boundary conditions and extended to a Born-Oppenheimer-like approach in the first version of MetalWalls (Reed, Lanning, & Madden, 2007). The method was also discussed in the framework of Mass-Zero (MaZe) constrained molecular dynamics in (Coretti et al., 2020). An updated description of the statistical mechanics of the constant potential ensemble can be found in the reference (Scalfi et al., 2020).

Two different models are employed for the charge distributions of the electrode and electrolyte atoms constituting the electrochemical cell. On the one hand, electrode atoms carry a Gaussian charge distribution of amplitude Q_i and width η_i^{-1}

$$\rho_i(\mathbf{r}) = Q_i \eta_i^3 \pi^{-3/2} \exp(-\eta_i^2 |\mathbf{r} - \mathbf{R}_i|^2) \quad (1)$$

where \mathbf{R}_i are the electrode atoms positions.

On the other hand, electrolyte atoms have a point charge distribution

$$\rho_i(\mathbf{r}) = q_i \delta(\mathbf{r} - \mathbf{r}_i) \quad (2)$$

where \mathbf{r}_i are the electrolyte atoms positions. The Coulomb potential energy for the full charge distribution $\rho = \sum_i \rho_i$ is given (in atomic units) by

$$V_c[\rho] = \frac{1}{2} \int d^3\mathbf{r} d^3\mathbf{r}' \frac{\rho(\mathbf{r})\rho(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|} \quad (3)$$

As it is long-ranged, it cannot be truncated at a cut-off distance. In MetalWalls, it is calculated using the Ewald summation method, whose specific expression depends on the periodicity (2D or 3D) of the system (Reed et al., 2007). Given the above form of the charge density, for fixed atomic positions it can also be expressed as a $V_c(Q)$, where Q is a vector containing all the N electrode atom charges (Q_1, \dots, Q_N).

The electric potential generated by the charge distribution associated with an electrode atom is $\frac{\partial V_c(Q)}{\partial Q_i}$. It follows that the constant electric potential condition within the electrode materials is given by

$$\frac{\partial V_c(Q)}{\partial Q_i} = \Psi_i \quad (4)$$

where Ψ_i is the applied potential difference on the electrode sites. This is equivalent to

$$\frac{\partial V(Q)}{\partial Q_i} = 0 \quad (5)$$

where the total potential energy $V(Q)$ is defined as

$$V(Q) = V_c(Q) - \sum_{i=1}^N \Psi_i Q_i \quad (6)$$

and includes the work necessary to charge the electrodes. The charges Q that satisfy the constant potential condition are found by solving the system of N equations given by Eq. (5).

It is also frequent, in constant potential simulations, to enforce the so-called electroneutrality constraint, defined by the equation

$$\sum_{i=1}^N Q_i = 0 \quad (7)$$

As the electrolyte ions undergo thermal motion, thereby changing the external potential, these equations must be solved to determine the electrode charges consistent with the constant potential condition. Two methods, described below, have been applied to achieve this.

Born-Oppenheimer approach

A first method to simulate the dynamics of the electrolyte atoms under the constant potential condition is to solve directly the equation

$$\frac{\partial V_c(Q)}{\partial Q_i} = \Psi_i \quad (8)$$

for each electrolyte configuration. This method relies on the quadratic form of the Coulomb energy with respect to the electrode charges to solve efficiently the equation for the Q . In particular, V_c can be written in the form

$$V_c(Q) = \frac{1}{2} Q^T A Q - Q^T b + c \quad (9)$$

which yields a set of equations for the charges given by $AQ - b = \Psi$, where the matrix A depends only on the geometry of the electrodes while the vector b depends also on the positions of the electrolyte atoms (Ψ is a vector containing the N electrode atom potentials (Ψ_1, \dots, Ψ_N)). The set of equations can then be solved by direct numerical matrix inversion yielding the charges

$$Q = A^{-1}(b + \Psi) \quad (10)$$

This is very efficient when the electrodes are fixed as the matrix A is constant and the inversion has to be performed only once at the beginning of the simulation. MetalWalls also allows constant pressure P_z simulations to be run by treating the electrodes as rigid pistons. In this case, a more efficient way to solve the linear system is by using the conjugate gradient iterative method.

The fulfillment of the electroneutrality constraint in this framework and its repercussions on the statistical mechanics of the constant-potential ensemble are discussed at length in (Scalfi et al., 2020).

Mass-Zero constrained MD (MaZe)

Another approach, based on the method introduced in (Coretti, Bonella, & Ciccotti, 2018), is to consider the motion of the electrolyte particles subject to the set of holonomic constraints given by

$$\frac{\partial V(Q)}{\partial Q_i} = 0 \quad (11)$$

This approach is based on an extended Lagrangian, where the electrode charges Q are considered additional dynamical variables for the system. Unlike previous extended-Lagrangian

approaches *à la* Car and Parrinello, the mass-zero limit is enforced rigorously in this method and the additional degrees of freedom are properly separated from the physical ones in an adiabatic sense. This yields dynamical equations of motion for the additional variables which can be solved numerically by symplectic and time-reversible algorithms, while keeping the adiabatic separation. In (Bonella, Coretti, Vuilleumier, & Ciccotti, 2020) it has been proven that the dynamics also samples the Born-Oppenheimer distribution exactly. The application of the method to constant potential simulations is described in detail in (Coretti et al., 2020) together with the possibility of enforcing the electroneutrality constraint.

Finite electric field variant

An extension of the constant potential approach was recently proposed, in which the electrochemical cell is simulated by including one electrode only, using 3D periodic boundary conditions. The electrode is set at constant potential and a finite field is applied to the whole cell. Two electrochemical interfaces form on the opposite sides of the metal, and we have shown that this approach yields similar results to the two electrodes setup for bulk electrode materials (Dufils, Jeanmairet, Rotenberg, Sprik, & Salanne, 2019).

Acknowledgements

MS would like to acknowledge the support from Maison de la Simulation through the « Chaire d'excellence » program. This project has received funding from the European Research Council (ERC) under the European Union's Horizon 2020 research and innovation programme (Grant Agreement No. 771294). This work was supported by the French National Research Agency (Labex STORE-EX, Grant No. ANR-10-LABX-0076). We acknowledge support from EoCoE, a project funded by the European Union Contracts No. H2020-EINFRA-2015-1-676629 and H2020-INFRAEDI-2018-824158.

References

- Bonella, S., Coretti, A., Vuilleumier, R., & Ciccotti, G. (2020). Adiabatic motion and statistical mechanics via mass zero constrained dynamics. *Phys. Chem. Chem. Phys.*, 22, 10775–10785. doi:[10.1039/d0cp00163e](https://doi.org/10.1039/d0cp00163e)
- Choudhuri, J. R., Vanzo, D., Madden, P. A., Salanne, M., Bratko, D., & Luzar, A. (2016). Dynamic response in nanoelectrowetting on a dielectric. *ACS Nano*, 10, 8536–8544. doi:[10.1021/acs.nano.6b03753](https://doi.org/10.1021/acs.nano.6b03753)
- Coretti, A., Bonella, S., & Ciccotti, G. (2018). Communication: Constrained molecular dynamics for polarizable models. *J. Chem. Phys.*, 149, 191102. doi:[10.1063/1.5055704](https://doi.org/10.1063/1.5055704)
- Coretti, A., Scalfi, L., Bacon, C., Rotenberg, B., Vuilleumier, R., Ciccotti, G., Salanne, M., et al. (2020). Mass-zero constrained molecular dynamics for electrode charges in simulations of electrochemical systems. *J. Chem. Phys.*, 152, 194701. doi:[10.1063/5.0007192](https://doi.org/10.1063/5.0007192)
- Dufils, T., Jeanmairet, G., Rotenberg, B., Sprik, M., & Salanne, M. (2019). Simulating electrochemical systems by combining the finite field method with a constant potential electrode. *Phys. Rev. Lett.*, 123, 195501. doi:[10.1103/physrevlett.123.195501](https://doi.org/10.1103/physrevlett.123.195501)
- Merlet, C., Rotenberg, B., Madden, P. A., Taberna, P.-L., Simon, P., Gogotsi, Y., & Salanne, M. (2012). On the Molecular Origin of Supercapacitance in Nanoporous Carbon Electrodes. *Nat. Mater.*, 11, 306–310. doi:[10.1038/nmat3260](https://doi.org/10.1038/nmat3260)

- Reed, S. K., Lanning, O. J., & Madden, P. A. (2007). Electrochemical Interface Between an Ionic Liquid and a Model Metallic Electrode. *J. Chem. Phys.*, 126, 084704. doi:[10.1063/1.2464084](https://doi.org/10.1063/1.2464084)
- Scalfi, L., Limmer, D. T., Coretti, A., Bonella, S., Madden, P. A., Salanne, M., & Rotenberg, B. (2020). Charge fluctuations from molecular simulations in the constant-potential ensemble. *Phys. Chem. Chem. Phys.*, 22, 10480–10489. doi:[10.1039/C9CP06285H](https://doi.org/10.1039/C9CP06285H)
- Siepmann, J. I., & Sprik, M. (1995). Influence of Surface-Topology and Electrostatic Potential on Water Electrode Systems. *J. Chem. Phys.*, 102, 511–524. doi:[10.1063/1.469429](https://doi.org/10.1063/1.469429)
- Simoncelli, M., Ganfoud, N., Sene, A., Haefele, M., Daffos, B., Taberna, P.-L., Salanne, M., et al. (2018). Blue energy and desalination with nanoporous carbon electrodes: Capacitance from molecular simulations to continuous models. *Phys. Rev. X*, 8, 021024. doi:[10.1103/physrevx.8.021024](https://doi.org/10.1103/physrevx.8.021024)