# Next-generation all-solid-state battery


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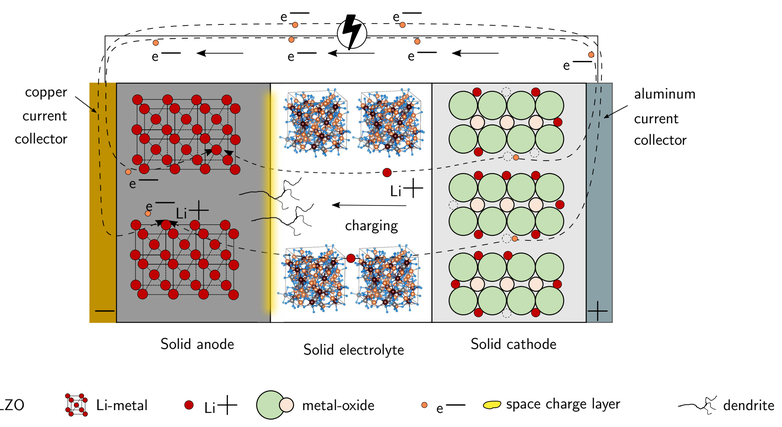
## Mathematical modelling for the next-generation All-solid-state batteries: Nucleation (SE|SSE)$^{(*)}$-interface

**Rechargeable Lithium-ion battery** (LIB) is at the heart of every electric vehicle (EV), portable electronic device, and energy storage system [1]. Nowadays, LIBs enable human life more efficient and help to solve global environment issues thanks to EVs' zero emission. However, conventional LIB (c-LIB) is sensible to temperature and pressure, hence, flammable and explosive. This is mainly due to liquid-based electrolyte found in cLIBs.
**All-solid-state battery** (ASSB) is one of promising candidates to overcome bottlenecks of c-LIBs. Thanks to solid-state electrolyte (SSE), ASSB is highly stable towards temperature and pressure. Nevertheless, metallic Li-dendrite triggered at (SE|SSE)-interface is the main drawback as these dendritic threads extrapolate into grain boundary network of SSE, causing crevice, degradation of ionic conductivity, and the probability of short-circuit.
**Next-generation All-solid-state battery** (ng-ASSB) with a consideration of nucleation criterion defined by

$$a_{\text{Griffith}} := a^* = \arg\min_{a\in\mathbb{R}} \iiint_{\Omega} f(a, \boldsymbol{u}; \lambda, \mu, \boldsymbol{d}\otimes\boldsymbol{d})\, d\Omega - \iint_{\Gamma} f(a;\gamma)\, d\Gamma \Big|_{\boldsymbol{u}^{(s)}}$$

where fdassa,
can help to improve ASSB performance.




## Next-generation All-solid-state battery

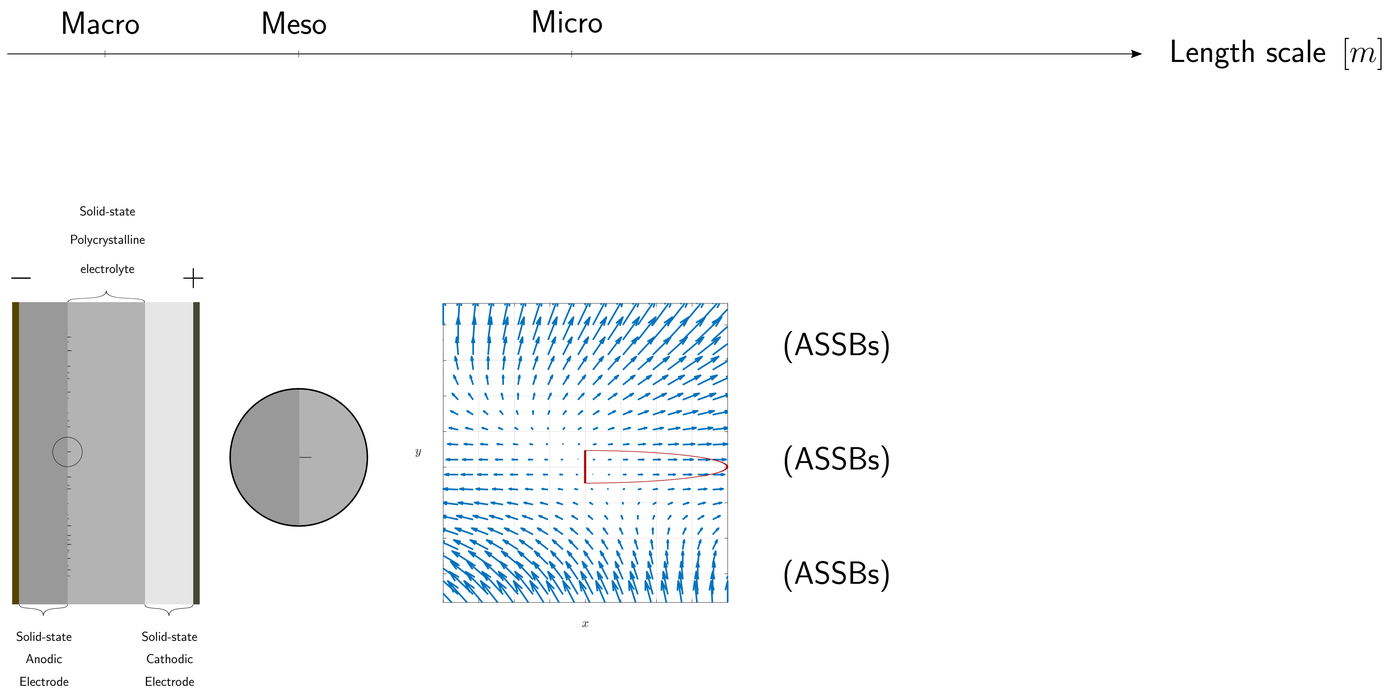
**Interface** between solid electrode and solid-state electrolyte (SE|SSE) taking place at space charge layer (SCL) [2] found in all-solid-state lithium-ion batteries (ASSLiBs) critically exhibits mechanical and electrochemical instability [3]. This evidence points directly to the fact that the soft metallic lithium negative electrode is erroneously prone to triggering dendritic by-products of silvery lithium metal, under cycles of electric charge & discharge [4]. Besides, polycrystalline garnet-typed solid-state electrolyte such as LLZO exhibit grain boundaries and various sizes and shapes of grains under microscopic observation. Therefore, this type of microstructure distinctively leads to nuance destruction of ceramic-like materials. Consequentially, dendritic by-products contribute to degradation of ionic conductivity and trace along grain boundaries in SSE.

$$\rho\,\partial^2_{t^2}\boldsymbol{u}^{(s)} + \nabla\cdot\left(\overset{4}{\mathbb{C}}{}^{\mathbb{D}(\Omega)}_{(\lambda,\mu)} : \nabla\boldsymbol{u}^{(s)}\right) + \rho\nabla V_e = \boldsymbol{0},$$

$$\text{s.t. } a_{\text{Griffith}} := a^* = \arg\min_{a\in\mathbb{R}} \iiint_{\Omega} f(a, \boldsymbol{u}; \lambda, \mu, \boldsymbol{d}\otimes\boldsymbol{d})\, d\Omega - \iint_{\Gamma} f(a;\gamma)\, d\Gamma \Big|_{\boldsymbol{u}^{(s)}}$$

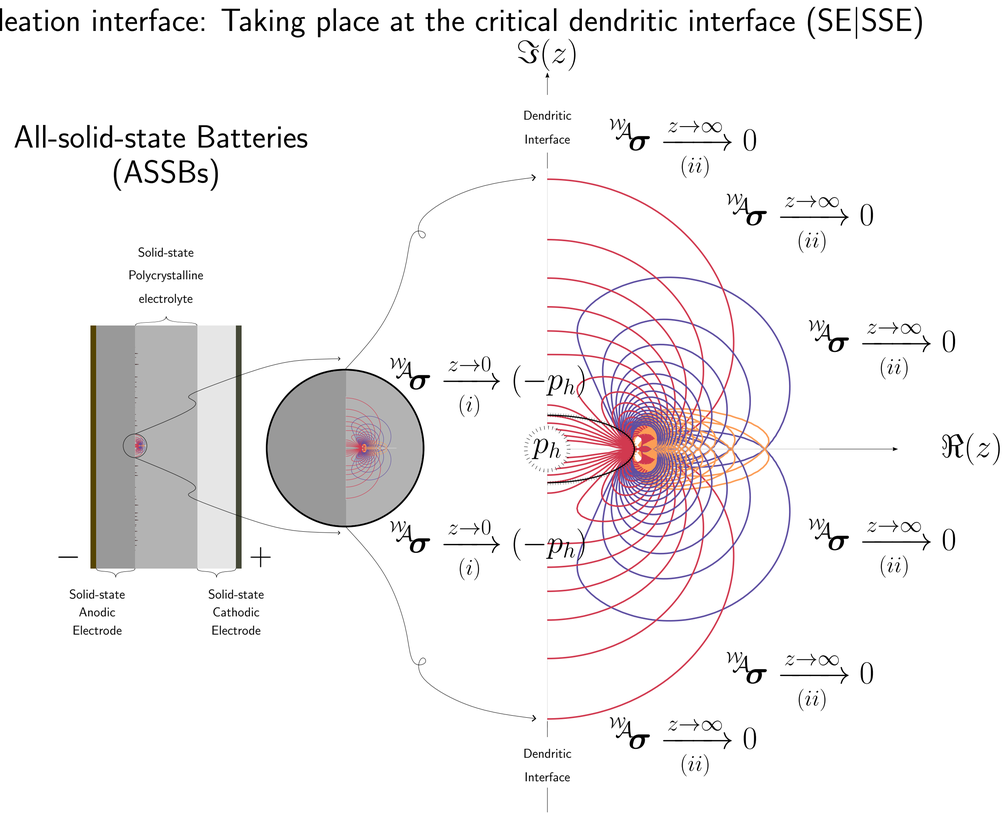
where

## Showcase

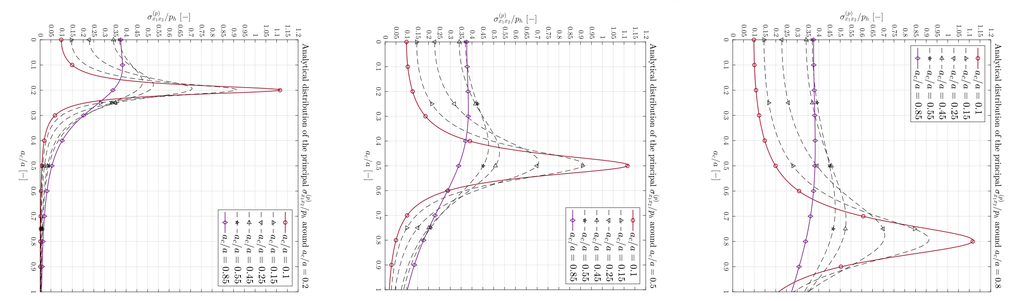



## Nucleation interface: Taking place at the critical dendritic interface

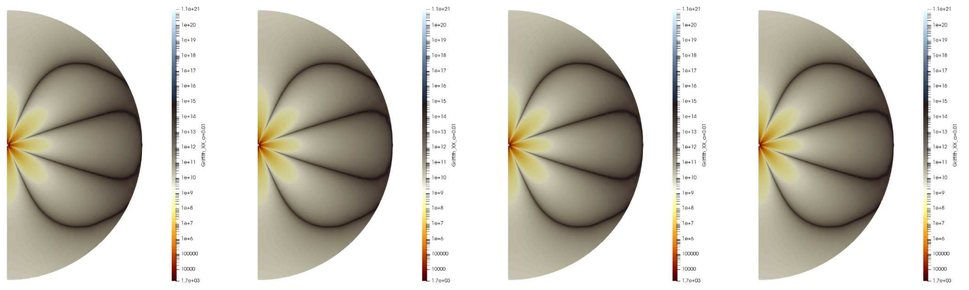
Nucleation interface: Taking place at the critical dendritic interface (SE|SSE)



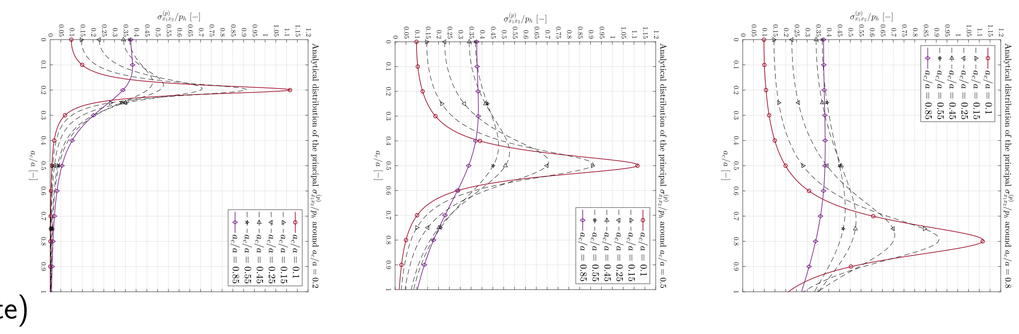

Distribution of the analytical maximal shear stress component ${}^{w_A}\sigma^{\Pi}_{x_1x_2}$, around the crack tip $a_c$.




The set of boundary conditions is likewise the path of the pressure-centric dendritic crack.



Comparison




Nucleation interface: Taking place at the critical dendritic interface (Solid electrode | Solid-state electrolyte)

## Contact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## References


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