

1 **Regionalized life cycle assessment of present and future lithium**
2 **production for Li-ion batteries**

3 **Authors:** Vanessa Schenker^{1,2}, Christopher Oberschelp^{1,2}, Stephan Pfister¹

4 **Affiliations:**

5 ¹ Institute of Environmental Engineering, ETH Zürich, John-von-Neumann-Weg 9, 8093 Zürich,
6 Switzerland

7 ² National Centre of Competence in Research (NCCR) Catalysis, ETH Zürich, Zürich, Switzerland

8 **Corresponding author:** Vanessa Schenker, vanessa.schenker@ifu.baug.ethz.ch, twitter: @schvanes

9 **Contacts of co-author:** Dr. Christopher Oberschelp, christopher.oberschelp@ifu.baug.ethz.ch

10 **Contacts of co-author:** Dr. Stephan Pfister, stephan.pfister@ifu.baug.ethz.ch

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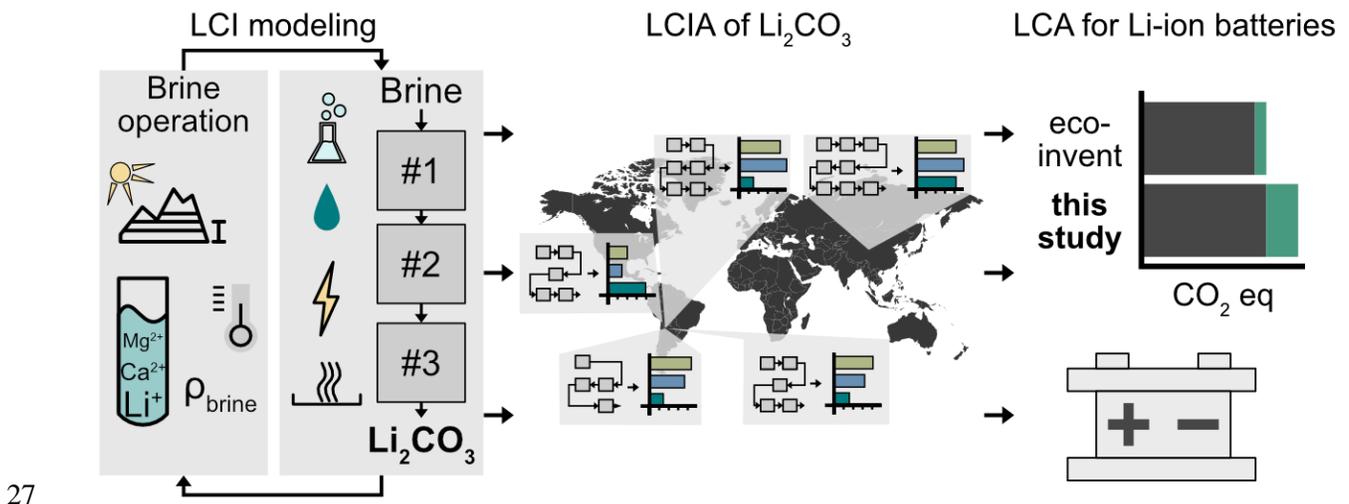
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14 **Abstract**

15 Existing life cycle assessments (LCA) of lithium carbonate production from brines are mainly based on one
16 single brine operation site, while many different lithium carbonate production routes have been developed
17 in the past. Hence, current life cycle inventories do not capture the variability of brine sites and misestimate
18 life cycle impacts. This study presents a systematic approach for LCA of existing and future lithium
19 carbonate production from brines, which can furthermore be applied to geothermal brines or seawater. It
20 has been used to model life cycle inventories of three existing and two upcoming brine operations in
21 Argentina, Chile, and China and combined with regionalized life cycle impact assessment. Impacts on
22 climate change, particulate matter human health impacts, and water scarcity from lithium carbonate
23 production differ substantially among sites. Existing life cycle inventories for lithium-ion battery
24 production underestimate climate change impacts by up to 19% compared to one from our study.

25 **Keywords:** Lithium, Environmental impacts, Life cycle assessment, Life cycle inventory, Brines

26 **Graphical abstract**



28 **1 Introduction**

29 The development of energy storage led to an increased demand for battery metals (Christmann et al., 2015;
30 Wanger, 2011; World Economic Forum, 2019). By 2030, the battery demand is forecasted to grow by 1400
31 % and hence, the demand for Lithium (Li) used in Li-ion batteries is expected to increase by a factor of 6
32 with respect to 2018 (World Economic Forum, 2019).

33 More than two-thirds of the Li resources are located in Argentina, Bolivia, Chile, and China as brine
34 deposits which hold a great supplying potential in the future (Bertau et al., 2017; Kesler et al., 2012; Munk
35 et al., 2016). The primary producer of lithium carbonate (Li_2CO_3) from brines is Chile, followed by
36 Argentina and China (S&P Global, 2021). Brine operations produce Li_2CO_3 with a technical grade (min.
37 99 wt. %) and battery grade (99.5 wt. %). The latter is used to manufacture Li-ion batteries (Dai et al.,
38 2020).

39 Various production routes for Li_2CO_3 from brines have been developed in the past (Tran and Luong, 2015).
40 Generally, the processing can be subdivided into three main processes: Brine's mass reduction in solar
41 evaporation ponds, brine purification, and Li_2CO_3 precipitation (Garrett, 2004; Tran and Luong, 2015). The
42 brine is pumped from an aquifer/salt lake into evaporation ponds to reduce the brine volume by solar
43 evaporation. When a specific Li concentration of the brine is reached, the brine is sent to the processing
44 plant. The purification part consists of a variety of processes in different arrangements to remove impurities
45 (Calcium (Ca), magnesium (Mg), or boron (B)) from the Li-enriched brine, such as by adding quicklime to
46 remove Mg, using organic solvent extraction to remove B, or using ion exchangers to remove Mg, Ca, or
47 B. The selected processes and their order depend on the site-specific brine compositions. Once the pulp has
48 been purified, Li_2CO_3 is precipitated by heating the pulp and adding soda ash. Crystallized Li_2CO_3
49 (technical grade) is dissolved in water at low temperature. The solution is re-heated at 80 °C, and Li_2CO_3
50 (battery grade) precipitates. The final product is dried in a rotary dryer (Garrett, 2004; Tran and Luong,
51 2015). In addition to this approach of extracting Li from brines, other processing techniques include
52 selective Li recovery. The Li-ion selective adsorption technique uses adsorbents (manganese oxide,

53 titanium oxide, or alumina) to selectively uptake Li from the brine sent through the ion exchangers (Garrett,
54 2004; Tran and Luong, 2015). Once the adsorbents are saturated with Li, the columns are washed and take
55 up Li. The Li-containing solution is then sent to evaporation ponds to further concentrate Li (Garrett, 2004).
56 Once a specific concentration is reached in the solution, Li_2CO_3 precipitation (technical grade) is forced by
57 heating the pulp and adding soda ash, as explained for the previous production route. Crystallized Li_2CO_3
58 (technical grade) is dissolved in water at low temperature by adding pressurized gaseous CO_2 . The solution
59 is re-heated at 80 °C, and Li_2CO_3 (battery grade) precipitates. The product is dried in a rotary dryer (Garrett,
60 2004; Tran and Luong, 2015).

61 Regarding environmental impacts, energy provision for Li_2CO_3 production is mainly based on fossil fuels
62 contributing to climate change (Stamp et al. 2012, Kelly et al. 2021). Stamp et al. (2012) published life
63 cycle inventory (LCI) data for brine-related Li_2CO_3 production from the Salar de Atacama in Chile. This
64 data was integrated into the ecoinvent LCI database in 2012 and has not been updated or expanded
65 (ecoinvent, 2021). Kelly et al. (2021) used more updated recent technical data from the Salar de Atacama
66 to quantify impacts on climate change and water scarcity but also did not improve on the coverage in terms
67 of different sites. Ambrose and Kendall (2019) slightly extended the coverage by including lab-scale data
68 from the Salar de Uyuni in Bolivia when assessing climate change impacts.

69 When assessing water scarcity impacts related to Li-ion battery storage, water scarcity impacts of Li from
70 brines were classified as critical, according to Schomberg et al. (2021). They included brine consumption
71 in the water scarcity footprint (WSF) when applying the LCA midpoint indicator AWARE. However, brines
72 are not directly used by ecosystems or humans as a water source and should, thus, not be considered when
73 applying this LCA method (Boulay et al., 2018). Brine pumping affects the hydrogeological systems with
74 wetland and lake ecosystems at the Salar de Atacama but these direct and indirect effects of brine pumping
75 are only measurable by assessing the hydrogeology of these salt flats (Liu et al., 2019; Liu and Agusdinata,
76 2021; Marazuela et al., 2019).

77 The scientific focus has been LCIs of Li_2CO_3 production from Salar de Atacama in Chile so far (57 % of
78 the LCE production from brines in 2018 (S&P Global, 2021)). However, processing techniques from other
79 brine operations differ from the one used at Salar de Atacama since they vary in their chemical composition
80 (Flexer et al., 2018; Houston et al., 2011; Munk et al., 2016; Tran and Luong, 2015). Thus, production
81 routes adapted to the brine chemistry and other environmental parameters were developed (Garrett, 2004;
82 Swain, 2017; Tran and Luong, 2015). Hence, what is missing so far in literature is a detailed assessment of
83 other Li_2CO_3 production pathways and their related environmental impacts. Furthermore, the existing LCA
84 studies are difficult to compare since the goal and scope of these studies vary. Differences in system
85 boundaries and degree of transparency hamper the direct comparison of these studies. Hence, the main
86 objective of this paper is to develop a systematic approach to model site-specific LCIs of Li_2CO_3 production
87 from brines when operational data from the companies are not publicly available. We apply our approach
88 by assessing environmental impacts of Li_2CO_3 (battery grade) production from five brine operations in
89 Chile, Argentina, and China. We cover climate change impacts, regionalized human health impacts from
90 fine particulate matter (PM) formation and partly regionalized WSFs. Finally, by integrating Li_2CO_3 from
91 different brine operations, the consequences on climate change impacts related to Li-ion battery production
92 were assessed.

93 **2 Methods**

94 **2.1 Framework to assess environmental impacts of Li_2CO_3 from brines**

95 We present an approach to quantify environmental impacts of Li_2CO_3 production from brines. Specifically,
96 we developed a modular approach to model site-specific LCIs, which allows for flexible adjustments to
97 future process updates at each extraction site and can also be applied to other brines in future research. The
98 approach follows the ISO 14040:2006 and ISO 14044:2006 standards to allow a standardized LCA (ISO,
99 2006a, 2006b). Hence, four steps need to be examined: Goal and scope definition, LCI analysis, life cycle
100 impacts assessment, and the final interpretation.

101 **Step 1: Goal and scope**

102 The goal and scope should be defined in the first step. We suggest setting the functional unit to 1 kg Li_2CO_3
103 (battery grade) which facilitates the integration of LCIs in Li-ion batteries. System boundaries should be
104 set accordingly to the project's scope (e.g., the system boundaries (cradle-to-gate approach) could be set
105 from pumping the brines to the surface until the final product (Li_2CO_3 , battery grade) leaves the processing
106 plant).

107 **Step 2: LCI analysis**

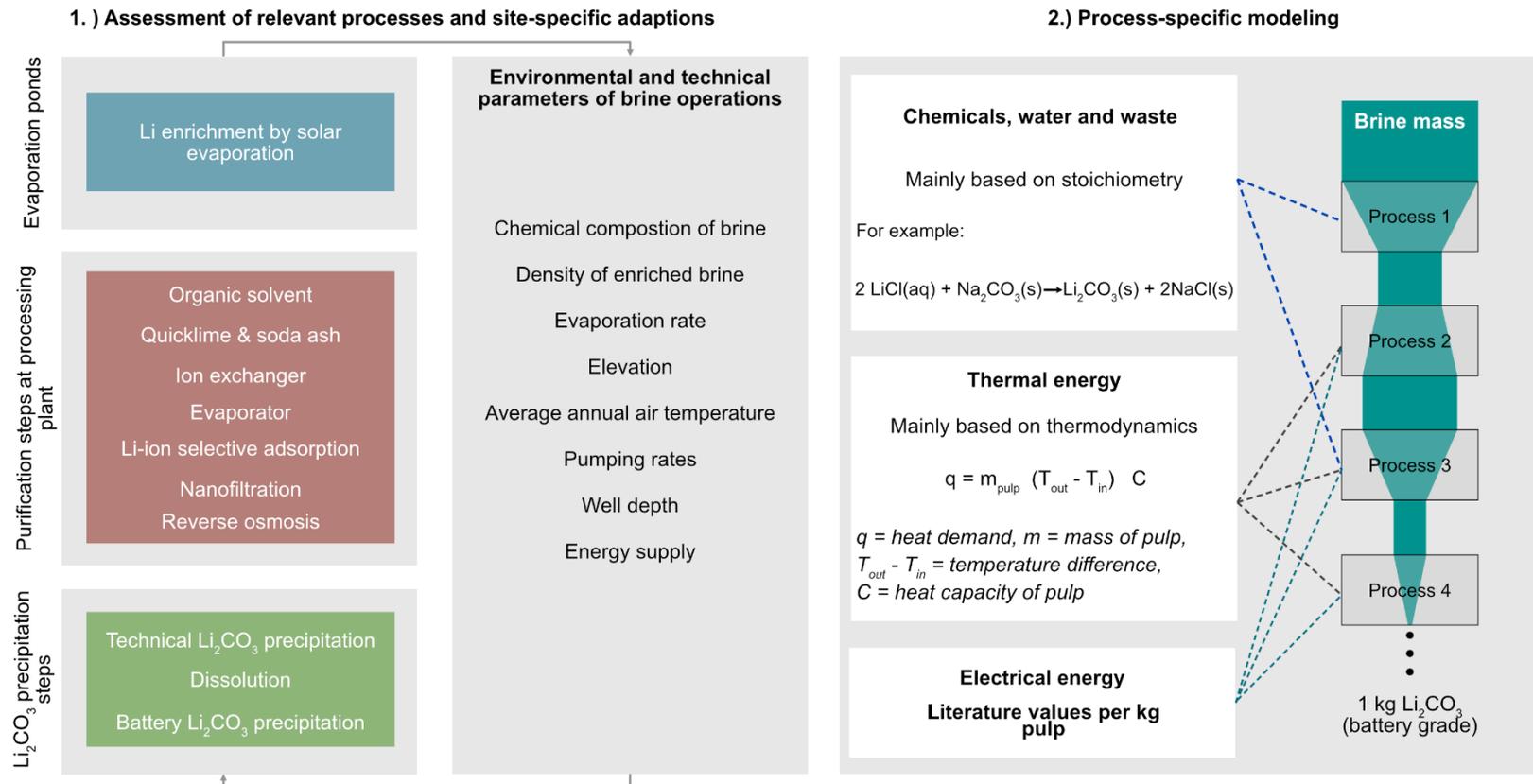
108 Step 2 consists of modeling the LCI for Li_2CO_3 production from one or multiple brine sites if site data are
109 not publicly available. We propose using the approach as developed in this study (Figure 1): (1)
110 identification of relevant processes and site-specific environmental and technical parameters and (2)
111 process-specific modeling of energy and material demand.

112 **(1) Identification of relevant processes and site-specific parameters**

113 Literature research (e.g., company reports, patents, scientific papers) is required to identify the relevant
114 processes to produce Li_2CO_3 . The process configuration determines the mass flows of the Li-containing
115 pulp and thus, requires a detailed assessment for the latter LCI modeling. The types of input and waste
116 production for all identified processes need to be defined. Environmental and technical parameters (e.g.,
117 chemical composition of the brine, evaporation rate, or annual average air temperature) need to be
118 considered because they influence resource demand of Li_2CO_3 production (e.g., heating demand of Li_2CO_3
119 precipitation or the chemical demand to remove B from the pulp). Sources of thermal energy can be
120 retrieved from company reports. Identifying processes and relevant parameters is an iterative process since
121 the type of Li_2CO_3 production also guides the number and type of parameters required for LCI modeling.

122 **(2) Process-specific modeling of material and energy demand**

123 The reported Li concentration of the enriched brine serves as an approximation for the required mass of
124 brine entering the processing plant. The input demand for processes, such as purification (Mg removal by
125 adding quicklime) and Li_2CO_3 precipitation (adding soda ash), is dependent on the mass flows (pulp
126 entering the process). The chemical composition of the brine affects required industrial chemical demand
127 (e.g., the mass of Mg in the brine is proportional to the amount of quicklime if it is added to brine stored in
128 the evaporation ponds (Flexer et al. 2018)). Mass of process-related chemicals (e.g., quicklime to remove
129 Mg in the pulp) and produced waste (e.g., NaCl precipitation due to soda ash) are suggested to be
130 stoichiometrically calculated. The required mass of chemicals should be adapted by adding a percentage to
131 account for the incompleteness of chemical reactions according to available data. Those inputs and outputs
132 need to be investigated to determine the mass of pulp going into the next process. Further affecting mass
133 flows within the processing plant is residual pulp re-circulation to previous purification steps, which needs
134 to be considered if reported. Energy demand can be quantified once process-specific mass flows are
135 determined. Thermal energy demands are influenced by the mass, temperature difference, and heat capacity
136 of the pulp (Figure 1). Literature values per kg pulp for process-specific electricity demands are available
137 and can be used to determine the operational electricity requirement on-site.



138

139 *Figure 1: Proposed approach to model LCI for Li₂CO₃ production from brines.*

140

141 **Step 3a: Life cycle impact assessment**

142 To assess environmental impacts of Li_2CO_3 (battery grade) production from each brine operation, we
143 suggest to consider impacts on climate change (GWP 100a) (IPCC 2013), fully regionalized LCA impact
144 assessment of fine PM formation (Oberschelp et al. 2020), and partly regionalized WSF based on AWARE
145 (Boulay et al., 2018). The selection of impact categories (impacts on climate change and water scarcity) is
146 based on existing literature (e.g., Stamp et al. 2012; Schomberg et al. 2021; Kelly et al. 2021). Since the
147 energy requirement is mainly based on fossil fuels (Kelly et al., 2021), PM-related human health impacts
148 should also be assessed, as it was shown in Oberschelp et al. (2019).

149 **Step 3b: Sensitivity analyses**

150 Monte Carlo simulations for brine operations allow to analyze the robustness of the results. Uncertainty
151 distribution types (e.g., triangular or log-normal distributions) need to be carefully chosen. If many data are
152 available, a random sampling of several datasets should be performed. Changing parameters based on
153 physical relations could be a less preferred option to test the results. For the case of limited data, the data
154 quality should be expressed by the Pedigree matrix described in Wernet et al. (2016).

155 **3 Results and discussion**

156 **3.1 Application of the approach to present and future brine sites**

157 To test our presented framework, the lithium extraction sites (Salar de Atacama, Salar de Olaroz, Salar de
158 Cauchari-Olaroz, Salar del Hombre Muerto (North), Chaerhan salt lake) were environmentally assessed.
159 Of the five selected sites, Salar de Atacama, Salar de Olaroz, and Chaerhan salt lake have been producing,
160 whereas Salar de Cauchari-Olaroz is currently in the construction phase and plans to start Li_2CO_3
161 production in 2022 (S&P Global, 2021) while Salar del Hombre Muerto (North) is at an early exploration
162 stage. For the latter, the start of mining activity and extraction technology is not yet clearly set (S&P Global,

163 2021), but a Li_2CO_3 production pathway has been suggested by Knight Piésold Ltd. and JDS Energy &
164 Mining Inc. (2019).

165 This paper covers 70 % of the brine-related lithium carbonate production worldwide (sum of current
166 production from the Salar de Atacama, Salar de Olaroz, and Chaerhan salt lake) (S&P Global, 2021). We
167 also cover future production sites at Salar de Cauchari-Olaroz and Salar del Hombre Muerto North (reported
168 and estimated production given in Table 1).

169 **Step 1: Goal and scope**

170 The goal is to quantify environmental impacts of Li_2CO_3 (battery grade) production from brine operations
171 and to allow a comparison among them using an attributional LCA with the cut-off allocation approach
172 from ecoinvent (Wernet et al., 2016). The functional unit was 1 kg Li_2CO_3 (battery grade). We used a
173 cradle-to-gate approach. The system boundaries from pumping the brine to the surface until the final
174 product (Li_2CO_3 , battery grade) leaves the processing plant for the South American salt lakes. At Chaerhan
175 salt lake, the brine is first sent to a K-fertilizer plant, and Li_2CO_3 production uses the effluent, which is
176 considered as a waste stream and, therefore, without burden in the cut-off allocation approach. Since only
177 Li_2CO_3 (technical grade) is produced (Gansu United testing services Co Ltd (2018), Lanke Lithium (2018)),
178 we added the processes (dilution and re-heating the Li-bearing solution) required to manufacture Li_2CO_3
179 (battery grade). We further assessed the environmental impacts of Li-ion battery production. We
180 incorporated our modeled LCIs of Chaerhan salt lake and Salar de Atacama in the Li-ion battery production
181 based on ecoinvent v3.8 cut-off (ecoinvent, 2021). The functional unit was 1 kg of rechargeable Li-ion
182 battery. Brightway 2 by Mutel (2017) and Activity Browser by Steubing et al. (2020) were used to conduct
183 the assessment.

184 **Step 2: LCI analysis**

185 **(1) Identification of relevant processes and parameters**

186 We identified the relevant processes and their input demand for the chosen brine sites (Table 1, graphical
187 illustration Figure 2). Required environmental and technical parameters are presented in the SI for each
188 brine operation. Salar de Atacama, Salar de Cauchari-Olaroz, and Salar del Hombre Muerto (North) have
189 similar general Li_2CO_3 production routes but with varying purification steps. While Salar de Olaroz uses
190 ion exchangers to remove impurities from the pulp, Chaerhan salt lake uses Li-ion selective ion exchangers
191 to adsorb Li.

192 *Table 1 Differences in main processing techniques for the investigated brine operations. Production in metric tons of*
193 *Salar de Atacama, Salar de Olaroz, and Chaerhan salt lake for the year 2018 are based on S&P Global (2021).*
194 *Estimated production at Salar de Cauchari-Olaroz is given by Andeburg Consulting Services Inc and Montgomery &*
195 *Associates (2019) and at Salar del Hombre Muerto (North) by Knight Piésold Ltd. and JDS Energy & Mining Inc.,*
196 *(2019).*

Brine operation	Processing techniques
Salar de Atacama, Chile (88 100 t Li_2CO_3)	The brine is pumped into evaporation ponds to enrich Li from 0.15 wt. % to 6 wt. % and then transported to the processing plant (Garrett 2004). Subsequently, purification steps consist of organic solvent extraction to remove B and adding quicklime, respectively soda ash to remove Mg and Ca (Wilkomirsky, 1999). Then soda ash is added to the heated brine to let Li_2CO_3 (technical grade) precipitate (Kelly et al., 2021; Wilkomirsky, 1999). Li_2CO_3 (technical grade) is dissolved at low temperatures and re-heated to produce Li_2CO_3 (battery grade).
Salar de Olaroz, Argentina (12 000 t Li_2CO_3)	The pumped brine first reacts with quicklime and is then enriched from 0.06 wt. % Li to 1.2 wt. % Li in the evaporation ponds. In the processing

	<p>plant, the brine reacts with soda ash to let impure Li_2CO_3 precipitate (Ehren and De Castro Alem, 2018; Orocobre, 2019). Then, Li_2CO_3 is dissolved in deionized water at low temperatures and the solution is sent through ion exchangers to remove residual Mg, Ca, and B. To precipitate Li_2CO_3 (battery grade), the pulp is re-heated as a last step (Ehren and De Castro Alem, 2018).</p>
<p>Salar de Cauchari-Olaroz, Argentina (40 000 t Li_2CO_3)</p>	<p>The pumped brine is enriched from 0.05 wt. % until 4 wt. % Li in evaporation ponds. Quicklime is added to remove Mg (Tran and Luong, 2015). B is removed via organic solvent extraction followed by removing Mg and Ca salts by adding quicklime and soda ash (Perez et al., 2014). The pulp is then heated to remove residual sulfates (Andeburg Consulting Services Inc and Montgomery & Associates, 2019). An evaporator is then used to decrease the volume of the Li-containing pulp, which is followed by an ion exchanger to remove any residual impurities (Andeburg Consulting Services Inc and Montgomery & Associates, 2019). In the next step, Li_2CO_3 (technical grade) is forced to precipitate by heating the pulp and adding soda ash. Subsequently, Li_2CO_3 is dissolved in water at low temperatures, and the solution is re-heated to produce Li_2CO_3 (battery grade) (Perez et al., 2014).</p>
<p>Salar del Hombre Muerto (North), Argentina (5 000 t Li_2CO_3)</p>	<p>A processing sequence similar to Salar de Atacama has been suggested (Knight Piésold Ltd. and JDS Energy & Mining Inc., 2019). As a first step, the brine would be enriched from 0.07 wt. % Li until 4 wt. % Li in evaporation ponds. Quicklime would be added to the evaporation ponds (Knight Piésold Ltd. and JDS Energy & Mining Inc., 2019). In the processing plant, purification steps would consist of B removal by organic</p>

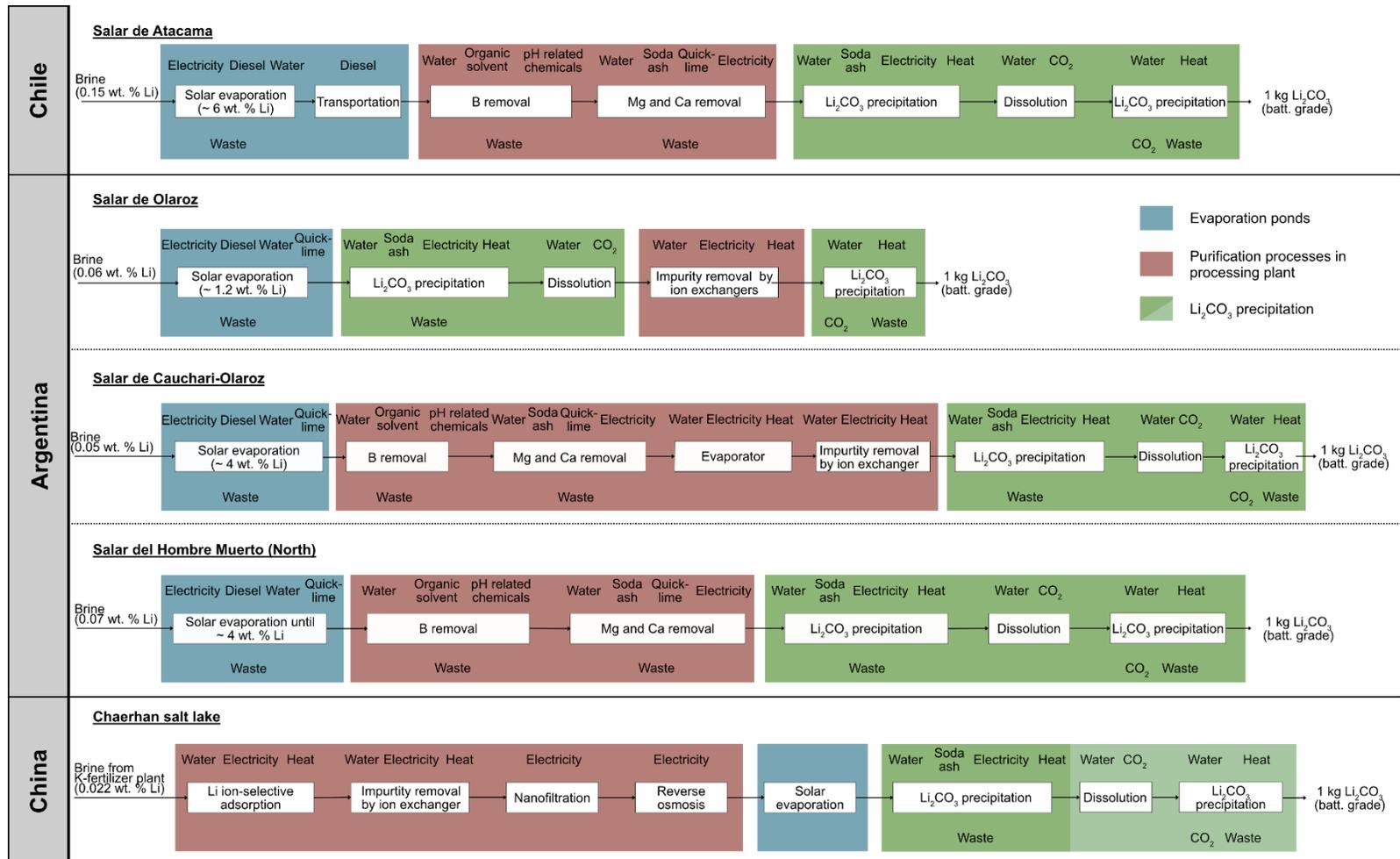
	<p>solvent extraction, Mg and Ca removal by adding soda ash and quicklime. Then, Li_2CO_3 (technical grade) precipitates by heating the pulp and adding soda ash. Li_2CO_3 (technical grade) is then dissolved in water at low temperatures and re-heated to precipitate Li_2CO_3 (battery grade).</p>
<p>Chaerhan salt lake, China (8 000 t Li_2CO_3)</p>	<p>This Li_2CO_3 production plant uses the residual K-depleted pulp of a K fertilizers production plant. This pulp contains 0.022 wt. % Li (Lanke Lithium, 2018). The first step is a Li-adsorption technique, where alumina hydroxide is used as a resin to adsorb Li selectively. Deionized water is used to remove Li from the adsorbent again. The Li-containing solution is sent through ion exchangers to remove still existing impurities. Nanofiltration and reverse osmosis are then required to reduce the volume of the Li-containing solution (Wen et al., 2006). Solar evaporation continuously reduces the brine volume from which subsequently Li_2CO_3 (technical grade) is produced (Gansu United testing services Co Ltd, 2018; Lanke Lithium, 2018; Li et al., 2020). Due to the aforementioned required system expansion, we added the process sequence for Li_2CO_3 (battery grade).</p>

197

198 **(2) Process-specific modeling of material and energy demands**

199 The resource demand of each process was calculated based on mass and energy balances as proposed by
200 our approach. We calculated the mass required in the evaporation ponds to produce 1 kg Li_2CO_3 (battery
201 grade) based on the reported Li concentration of the brine. For the evaporation ponds, waste production is
202 not considered because the wastes from the different sites consist mainly of precipitated salts discarded in
203 the near-by salt flats and are thus expected to have limited environmental impacts for the covered types of
204 impact categories. The Li concentration of the enriched brine then served as an approximation of the mass

205 going into the processing plant. Based on the calculated mass flow, chemicals were estimated based on
206 stoichiometries. To account for the incompleteness of the chemical reaction, 20 % mass is added to the
207 modeled quicklime consumption based on Flexer et al. (2018) and 10 % to the modeled soda ash
208 consumption (Li et al., 2020). 98.5 % of the organic solvent required in the B removal step is assumed to
209 be recycled (CELIMIN, personal communication). Since deionized water is required in various processes,
210 we estimated the required mass of water for each process. The brine operators reported water purification
211 steps. However, they are not explicitly stated. Hence, for all brine operations we assumed that brackish
212 water is treated by a reverse osmosis and an ion exchanger at all salt lakes (e.g., Lithium Americas Corp.
213 2019). Due to the lack of site-specific information regarding waste treatment at the processing plants, we
214 did not include waste treatment in the LCI, in contrast to the existing dataset in ecoinvent v3.8 (ecoinvent
215 2021). Sources of thermal and electrical energy (i.e., heat from natural gas, heat and power co-generation
216 from natural gas, and power from the location-specific grid) were based on company reports. Efficiencies
217 of thermal processes are assumed to be 85 % due to the lack of information in used literature (U.S.
218 Department of Energy Energy Efficiency and Renewable Energy, 2003).



219

220 Figure 2 Process sequence of the five brine sites. The blue squares indicate that the processes are related to evaporation ponds, while red squares represent
 221 purification steps in the processing plant. Green squares represent Li_2CO_3 precipitation steps. Bright green squares within the processing sequence of Chaerhan
 222 salt lake indicate the required system expansion.

223 **Modeled resource consumption**

224 Table 2 presents the modeled inputs per kg Li₂CO₃ (battery grade) from the five sites. We find that Chaerhan
 225 salt lake has the highest consumption in electricity (27.8 kWh/kg Li₂CO₃), heat (298 MJ/kg Li₂CO₃), and
 226 water (474 kg/kg Li₂CO₃) due to the specific Li-ion adsorption, ion exchangers and the following
 227 nanofiltration, respectively reverse osmosis used for purification. Water (219 kg/kg Li₂CO₃) and electricity
 228 (1.5 kWh/kg Li₂CO₃) demand at Salar de Olaroz are higher than the other South American salt lakes. The
 229 water and electricity consumption originates from the intensive use of ion exchangers for removing
 230 impurities and re-generating ion exchangers. Since Salar de Olaroz relies on removing impurities by ion
 231 exchangers, Salar de Olaroz does not require any B removal-related chemicals in contrast to Salar de
 232 Atacama, Salar de Cauchari-Olaroz, and Salar del Hombre Muerto (North). Quicklime demand in the
 233 evaporation ponds to remove Mg is highest at Salar de Olaroz, while Salar de Atacama requires the lowest
 234 quicklime demand per kg Li₂CO₃. At Salar de Atacama, quicklime is only required to remove residual Mg
 235 from the pulp in the processing plant. Soda ash is used at all sites, mainly due to the Li₂CO₃ precipitation
 236 step.

237 *Table 2 Modeled life cycle inputs for 1 kg Li₂CO₃ (battery grade) production at selected salt lakes.*

Input demand/kg Li₂CO₃	Salar de Atacama	Salar de Olaroz	Salar de Cauchari-Olaroz	Salar del Hombre Muerto	Chaerhan salt lake
<i>Electricity [kWh]</i>	0.4	1.5	0.7	0.8	28
<i>Heat [MJ]</i>	19	19	28	14	298
<i>Water [kg]</i>	38	219	46	43	474
<i>Quicklime [kg]</i>	0.04	4.1	2.7	3.1	-
<i>Sodium hydroxide [kg]</i>	0.06	-	0.35	0.08	-
<i>Organic solvent [kg]</i>	0.04	-	0.1	0.7	-
<i>Hydrochloric acid [kg]</i>	0.10	-	0.5	0.9	-
<i>Soda ash [kg]</i>	1.9	1.6	2.1	1.6	1.6

238 **Step 3a: Life cycle impact assessment**

239 To assess environmental impacts of Li₂CO₃ (battery grade) production from each salt lake, GWP 100a
 240 (IPCC 2013), globally regionalized LCA impact assessment of PM formation (Oberschelp et al. 2020), and
 241 partly regionalized WSF based on AWARE (Boulay et al., 2018) were chosen.

242 **Climate change impacts**

243 Figure 4-A presents the climate change impacts to produce 1 kg Li_2CO_3 (battery grade) and their causes. In
244 addition, we compare our results with two datasets (for Li_2CO_3 from brines and spodumene-bearing
245 pegmatites) provided in ecoinvent v3.8 (ecoinvent, 2021).

246 We find that Li_2CO_3 production from the Salar de Atacama has the lowest climate change impacts (3.4 kg
247 $\text{CO}_2\text{eq/kg Li}_2\text{CO}_3$). Heat (41 % of the climate change impacts) and on-site chemicals use (38 %) are the
248 predominant contributors to the overall impacts. Soda ash for Li_2CO_3 precipitation is responsible for 31 %
249 of the total climate change impacts, while other chemicals, like organic solvents, quicklime, and
250 hydrochloric acid, only contribute minor shares. Hence, we find that Li_2CO_3 (technical grade) precipitation
251 followed by Li_2CO_3 (battery grade) precipitation are the major processes contributing to climate change per
252 kg Li_2CO_3 (battery grade) at Salar de Atacama (Figure 3). This is also in accordance with the findings by
253 Stamp et al. (2012) and Kelly et al. (2021) when assessing environmental impacts related to the Li_2CO_3
254 production at the Salar de Atacama.

255 Climate change impacts related to the Argentinian brines are up to 235 % higher than for Li_2CO_3 extracted
256 from the Salar de Atacama (Salar de Olaroz: 7.4 kg $\text{CO}_2\text{eq/kg Li}_2\text{CO}_3$; Salar de Cauchari-Olaroz: 7.7 kg
257 $\text{CO}_2\text{eq/kg Li}_2\text{CO}_3$; Salar del Hombre Muerto (North): 8 kg $\text{CO}_2\text{eq/kg Li}_2\text{CO}_3$). Quicklime required in the
258 evaporation ponds to remove impurities contributes to the total climate change impacts (Salar de Olaroz:
259 67 % of the total impact; Salar de Cauchari-Olaroz: 43 %; Salar del Hombre Muerto (North): 49 %), while
260 heat and electricity on-site contribute to a minor extent. Quicklime production, in general, is associated
261 with significant GHG emissions (Eula 2014). First, the production is heavily energy-intensive and mainly
262 based on fossil fuels. Second, the chemical reaction to produce quicklime from limestone emits CO_2 as a
263 co-product (European Lime Association, 2014). Those two factors are reflected in the overall impacts if
264 quicklime is used in these evaporation ponds (Figures 3 and 4-A). Hence, evaporation ponds and Li_2CO_3
265 (technical grade) precipitation are the most critical contributors due to the usage of quicklime, respectively

266 soda ash for the Argentinian salt lakes. At Salar del Hombre Muerto (North) and Salar de Cauchari-Olaroz
267 specifically, the B removal step additionally makes up a significant share of climate change impacts.

268 We find that Li_2CO_3 from Chaerhan salt lake has by far the highest climate change impacts (31.6 kg
269 $\text{CO}_2\text{eq/kg Li}_2\text{CO}_3$) resulting from the heat and electricity demand for the Li-ion selective adsorption
270 technique. This technique mainly includes two phases (adsorption and desorption phase) which require the
271 solution being heated up to a specific temperature using natural gas. Furthermore, electricity from the
272 provincial electricity grid is required for the ion exchangers and membrane separation (nanofiltration and
273 reverse osmosis) to remove impurities in the Li_2CO_3 bearing solution.

274 Climate change impacts of Li_2CO_3 from brines in ecoinvent v3.8 (2.1 kg $\text{CO}_2\text{eq/kg Li}_2\text{CO}_3$) and a recent
275 study by Kelly et al. (2021) (2.7 – 3.1 kg $\text{CO}_2\text{eq/kg Li}_2\text{CO}_3$) are in the same range as the ones of our modeled
276 LCI for Salar de Atacama. However, these numbers are lower than the other brines in Argentina and China,
277 underestimating the climate change impacts of average Li_2CO_3 . If Li_2CO_3 is extracted from spodumene-
278 bearing pegmatites, as described with the dataset provided by ecoinvent v3.8 (ecoinvent, 2021), the climate
279 change impacts add up to 10.7 kg $\text{CO}_2\text{eq/kg Li}_2\text{CO}_3$. Kelly et al. (2021) estimated 20.4 kg $\text{CO}_2\text{eq/kg Li}_2\text{CO}_3$
280 from the Australian pegmatitic mine. Both estimations are higher than our results for the Argentinian salt
281 lakes but still lower than the one for Chaerhan salt lake.

282



283

284 *Figure 3 Process-related contributive analyses regarding climate change impacts assessed with GWP (100 years)*
 285 *shown in percentage. The blue squares indicate that the processes are related to evaporation ponds, while red squares*
 286 *represent purification steps in the processing plant. Green squares indicate Li_2CO_3 precipitation steps. Bright green*
 287 *squares within the processing sequence of Chaerhan salt lake indicate that processes were added to hypothetically*
 288 *produce Li_2CO_3 (battery grade) to provide the same functional unit as the other systems.*

289 **PM-related human health impacts**

290 The results indicate a large variability in PM-related health impacts (Figure 4-B), which underlines the
291 necessity to perform regionalized LCAs for brine operations in this impact category. The highest PM health
292 impacts occur due to the Li_2CO_3 production at Chaerhan salt lake (1.2 micro-disability adjusted life years
293 (μDALY)/kg Li_2CO_3) followed by Salar del Hombre Muerto (0.67 μDALY /kg), Salar de Cauchari-Olaroz
294 (0.67 μDALY /kg), Salar de Atacama (0.43 μDALY /kg), and Salar de Olaroz (0.31 μDALY /kg). For all
295 sites, the background processes predominantly contribute to PM health impacts in contrast to foreground
296 processes (i.e., heat or diesel consumption) from the remote location of all salt lakes and low local
297 population densities. Electricity use in China, India, and Indonesia for various products and services in the
298 background system makes up a significant share of the overall impacts of all brine sites (Salar de Atacama:
299 31 %; Salar de Olaroz: 44 %; Salar de Cauchari-Olaroz and Salar del Hombre Muerto (North): 56 %;
300 Chaerhan salt lake: 48 %). The contribution of electricity required or generated on-site varies largely for
301 PM health impacts. While the Argentinian electricity mix contributes to the overall PM health impacts with
302 less than 1 %, the Chilean and Chinese electricity mix significantly contribute (27 % and 17 %) due to
303 particulates, $< 2.5 \mu\text{m}$, sulfur dioxide, and nitrogen oxides from coal power generation. 43 % of the Chilean
304 electricity mix comes from coal power. The Qinghai province-specific electricity mix, which was chosen
305 for Chaerhan salt lake, consists of 22 % coal power, while the main source in this region is hydropower. In
306 contrast, the Argentinian electricity mix includes little coal power (1 %) and does not significantly
307 contribute to the PM health impacts.

308 Soda ash is a relevant contributor to the PM health impacts at all sites. The specific contributions range
309 from 4 % at Chaerhan salt lake to 16 % at Salar de Atacama. During soda ash production, ammonia is
310 released into the atmosphere contributing significantly to PM health impacts in highly populated areas, such
311 as Europe. However, the location of soda ash production is highly uncertain since our results rather reflect
312 the LCI in ecoinvent v3.8 than the actual resource supplier due to missing operational data.

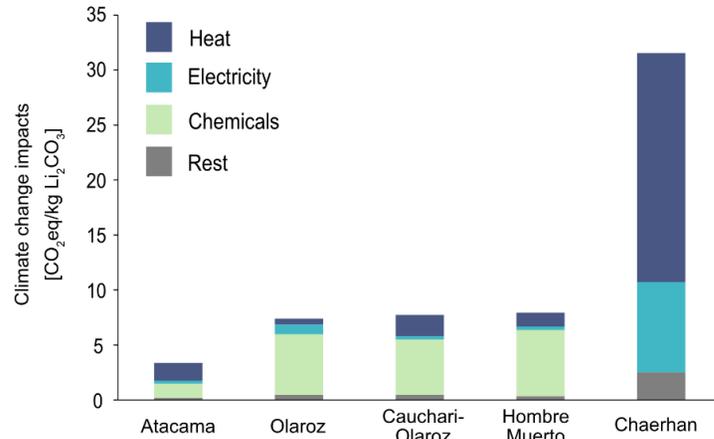
313 **Water scarcity footprint**

314 The WSF for each salt lake using the AWARE method (Boulay et al., 2018) is presented in Figure 4-C.
315 Chaerhan salt lake has the highest impact on water scarcity ($35.25 \text{ m}^3_{\text{world eq}}/\text{kg Li}_2\text{CO}_3$), followed by Salar
316 de Atacama ($4.77 \text{ m}^3_{\text{world eq}}/\text{kg Li}_2\text{CO}_3$). The Argentinian salt lakes are in the same range ($1.36 \text{ m}^3_{\text{world eq}}/\text{kg}$
317 Li_2CO_3 at Salar del Hombre Muerto, $1.62 \text{ m}^3_{\text{world eq}}/\text{kg Li}_2\text{CO}_3$ at Salar de Cauchari-Olaroz, and $1.73 \text{ m}^3_{\text{world}}$
318 $\text{eq}/\text{kg Li}_2\text{CO}_3$ at Salar de Olaroz). The water scarcity impacts of Salar de Atacama predominately originate
319 from the direct use of freshwater at the processing plant (81 %). However, Salar de Atacama has the lowest
320 water demand on-site compared to all other salt lakes. Nevertheless, due to its high aridity (e.g., Munk et
321 al. (2016)) the location-specific characterization factor is the highest with $94.7 \text{ m}^3_{\text{world eq}}/\text{m}^3$ amongst these
322 salt lakes, which is reflected in the overall water scarcity impacts.

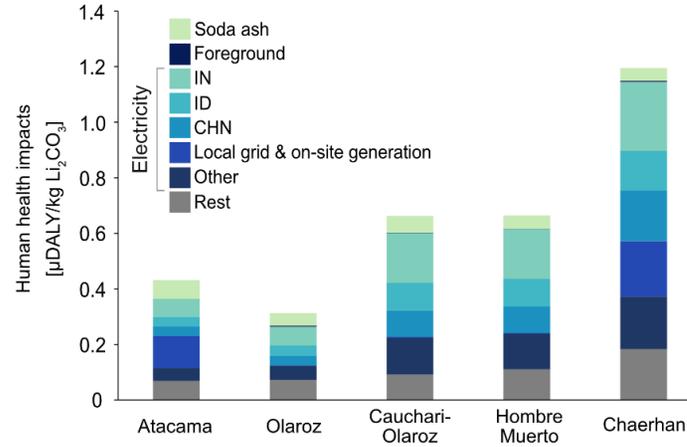
323 The Argentinian brines have the lowest WSF due to their relatively low characterization factor ($2.7 - 5$
324 $\text{m}^3_{\text{world eq}}/\text{m}^3$). This is particularly important for Li_2CO_3 production at Salar de Olaroz, which has a high
325 water requirement due to the intensive use of ion exchangers in the foreground system. The water demand
326 originates from the regeneration of the resin used to remove residual impurities and then allow a Li_2CO_3
327 (battery grade) precipitation. Nevertheless, the overall impact is lower than the one from Salar de Atacama.
328 In contrast to Salar de Olaroz, water scarcity impacts in the foreground are minor compared to the ones in
329 the background at Salar de Cauchari-Olaroz (85 %) and Salar del Hombre Muerto (North) (91 %).

330 The WSF from Li_2CO_3 production at Chaerhan salt lake originates from the extensive water use (see chapter
331 resource consumption) in the processing plant due to the Li-ion specific adsorption technique. Furthermore,
332 the location-specific characterization factor ($70.6 \text{ m}^3_{\text{world eq}}/\text{m}^3$) contributes to the relatively high WSF. The
333 water demand in the background only accounts for 5 % of the total WSF. In general, it has to be noted that
334 the background water consumption was not allocated to specific regions and was assessed with the global
335 average AWARE characterization factor, which is rather high (Boulay et al., 2018). Therefore, background
336 water stress might be overestimated in some cases.

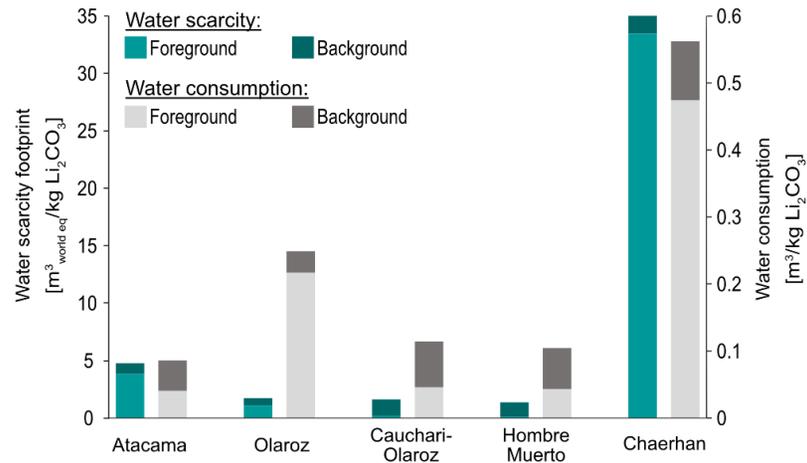
A. Climate change impacts



B. PM-related human health impacts



C. Water scarcity footprint



337

338 *Figure 4 Environmental impacts of Li₂CO₃ production from brines. (A) Impacts on climate change (GWP 100 yrs), (B) PM-related human health impacts and*

339 *(C) WSF (primary y-axis), and water consumption (secondary y-axis). (A) and (C) are shown on a midpoint level and (B) on an endpoint level.*

340 **Step 3b: Robustness and limitations of the approach**

341 We ran Monte Carlo simulations with $n = 5000$ runs for the foreground system of each salt lake using GWP
342 (IPCC 2013) and PM health impacts (Oberschelp et al. 2020) to assess the robustness of our results (more
343 information in the SI). The relative standard deviation ranges from $\pm 33\%$ (Salar de Atacama) to $\pm 98\%$
344 (Chaerhan salt lake) for impacts on climate change, while the Argentinian salt lakes are in-between. For
345 PM-related health impacts, the relative standard deviation lies between $\pm 46\%$ (Salar de Cauchari-Olaroz
346 and Salar del Hombre Muerto (North)) and $\pm 73\%$ (Chaerhan salt lake). The relatively higher standard
347 deviation of Chaerhan salt lake is explained by the lowest data quality of all assessed sites. In general, the
348 origin of inputs is mostly unknown, thus contributing to the uncertainties. We, therefore, relied onecoinvent
349 data. In order to decrease these uncertainties, site-specific information regarding the input supply is crucial.
350 If these data are not available, country-specific trading data could be obtained to decrease the uncertainties
351 in the future.

352 There are also data gaps in our modeling approach which need to be discussed. Energy, water, and chemical
353 demand modeled for this study were compared with annually reported company data or technical reports
354 from exploration activity (more information in SI). We used company data as an indicator rather than as a
355 fixed reference because specific boundaries and further documentation were generally not provided.
356 Quantitative chemical demand at Salar de Cauchari-Olaroz has not been reported to the authors' knowledge
357 and thus, could not be compared with our results. Furthermore, the LCI modeling of Li_2CO_3 (battery grade)
358 from Chaerhan salt lake predominately relies on parameters reported in construction plans by Gansu United
359 testing services Co Ltd (2018), impeding to test robustness of the model for that site. Annual changes of
360 the brine chemistry are challenging to include in the LCI, but may affect resource consumption (especially
361 chemical and heating demand) on-site. We could not estimate the salt crystallization sequence and hence,
362 calculate waste production in evaporation ponds. However, these wastes are mainly deposited on-site,
363 requiring little transport.

364 **3.2 Implications for Li-ion battery production**

365 To set this study into a broader context, modelled LCIs of two brine sites (Chaerhan salt lake and Salar de
366 Atacama) were implemented in an ecoinvent v3.8 dataset, which represents the production of 1 kg
367 rechargeable Li-ion batteries. This type of battery is used for a variety of electrical vehicles (Crenna et al.,
368 2021). Furthermore, we also adjusted the ecoinvent dataset to only use Li from the brine dataset in ecoinvent
369 v3.8. This leads to three battery datasets:

- 370 1. 100 % of the Li_2CO_3 production originates from the dataset for Li_2CO_3 from brines in ecoinvent
371 v3.8 (based on Salar de Atacama).
- 372 2. 100 % of the entire Li_2CO_3 production is assumed to be from Salar de Atacama.
- 373 3. 100 % of the entire Li_2CO_3 production is replaced by Li_2CO_3 from Chaerhan salt lake.

374 As already Stamp et al. (2012) and Kelly et al. (2021) highlighted, the source of lithium affects the amount
375 of GHG emissions related to Li-ion battery production. The maximum increase of climate change impacts
376 is 19 % when implementing 100 % Li_2CO_3 production by Chaerhan salt lake (dataset 3) compared to the
377 baseline (dataset 1). The leading cause is the high thermal and electrical energy demand based on fossil
378 fuels of Chaerhan salt lake. Li_2CO_3 from Salar de Atacama (dataset 2) only increases climate change impact
379 of a Li-ion battery by < 1 %. In the future, more lithium production might be sourced from high-impact
380 mines since an increase in demand and price might make energy-intensive production routes profitable.
381 Multiple studies emphasize the range of reported or modeled GHG emissions related to Li-ion battery
382 production (e.g., Raugei and Winfield 2019; Crenna et al. 2021). For our study, these findings indicate that
383 the overall share of Li_2CO_3 from brines regarding impacts may change in the future due to more detailed
384 and transparent Li-ion battery supply chains. Thus, LCA and carbon footprinting of future battery
385 production should consider the potential for high-impact Li_2CO_3 supply and consider the existing LCIs as
386 highly uncertain.

387 **4 Conclusion and outlook**

388 The Li supply is key for the transition towards a global decarbonized society. Li expects higher growth
389 rates than other metals in the future. Hence, it is inevitable to assess environmental impacts of current and
390 future Li extraction to avoid severe environmental burden shifting. Currently used LCIs of Li_2CO_3 from
391 brines do not represent the global market nowadays and even less for the future. Thus, we developed a
392 framework to update LCIs of Li_2CO_3 production from brines site-specifically. This methodology was
393 applied to existing and future production sites. Our framework helps to treat data gaps and to derive process-
394 specific parameters from patents. Furthermore, an approach to assess sites using waste streams as a Li
395 source, like Chaerhan salt lake, was developed. Regionalization of foreground and background data as well
396 as using regionalized impact assessment methods were examined and discussed in detail. Our results
397 demonstrate the necessity of defining a framework to assess various sites and resulting variabilities in global
398 production. The case studies show that available literature data underestimate environmental impacts of
399 Li_2CO_3 production from brines. This mainly is a consequence of only assessing Li_2CO_3 production at Salar
400 de Atacama and assuming that this data is representative of Li_2CO_3 production in general, which is not the
401 case. Furthermore, the variability of our results is a consequence of the brine composition, the applied
402 processing technique, and the brine location. For instance, water scarcity and PM impacts need to be site-
403 specifically assessed since the location of impact (background or foreground) varies among these sites.
404 Future improvements regarding the assessment of Li should focus on other Li sources, such as pegmatites,
405 geothermal brines, and seawater.

406 The integration of obtained LCIs in Li-ion batteries demonstrates that the overall impacts on climate change
407 increase to up to 19%. Hence, supply chains of Li-ion batteries need to be assessed in detail, especially for
408 future scenarios. Improvements regarding the resolution of supply chains are crucial to transition towards
409 low-carbon technologies sustainably. This includes site-specific assessment of other minerals for batteries
410 like Aluminum and Cobalt, such as recently done for copper mine tailings (Adrianto et al., 2022). Our
411 framework serves as a starting point for enhancing LCI and regionalized LCIA of other battery minerals.

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422 **Declaration of interest**

423 The authors declare no competing interests.

424

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