Potential of Mechanochemically Activated Sulfidic Mining Waste Rock for Alkali Activation

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Abstract
Sulfidic mining waste rock is a side stream from the mining industry with a potential environmental burden. Alkali activation is a promising method for transforming mining waste into construction materials. However, the low reactivity of minerals can be a sizeable challenge in alkali activation. In the present study, the reactivity of waste rock was enhanced by mechanochemical treatment with a LiCl-containing grinding aid. X-ray diffraction (XRD) and diffuse reflectance infrared Fourier transform (DRIFT) analysis were utilized to display the structural alteration of individual minerals. A schematic implication of the grinding mechanism of mica was provided according to the results of transmission electron microscopy (TEM) and scanning electron microscopy (SEM). The alkaline solubility displayed the enhanced chemical reactivity of the waste rock, in which Si and Al solubility increased by roughly 10 times and 40 times, respectively. The amorphization of aluminosilicate is achieved through chemical assisted mechanochemical activation. Sulfidic waste rock, as the sole precursor in alkali activation, achieved a 28-day compressive strength exceeding 10 MPa under ambient curing conditions. The simulation of the upscaled grinding process was conducted via the HSC Chemistry® software with a life-cycle assessment. The results showed that mining waste rock can be a promising candidate for geopolymer production with a lower carbon footprint, compared to traditional Portland cement.

Graphical Abstract

Keywords Mechanochemical activation · Sulfidic mining waste rock · Geopolymer · Valorization · Life-cycle assessment · HSC Chemistry®

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Introduction

Wastes from mining and mineral processing pose potential environmental and chronic burdens across the world. In North America, the stream of mining waste is as much as 10 times larger than the municipal solid waste per capita. According to a recent European statistics report, almost two thirds (65%) of the total waste generated in the EU-27 in 2018 were composed of major mineral waste [1]. Waste rock is the material generated from excavation of ore, after which sections rich in ore are sent to further ore processing [2]. Mining waste rock is dumped near the mining site or stored in landfills, which can be a potential of pollutants. Leachable hazardous metals can dissolve into soil, and acid leachate can contaminate underground water through acid mine drainage (AMD). The utilization of mining wastes is concerning because it poses potential negative impacts on the landscape and local community. Mining waste rock has been used as a secondary raw material in building materials, such as aggregates and raw meals for Portland cement production [3]. However, construction materials produced solely with waste rock from sulfidic mining have hardly been studied.

Alkali activation is a promising method for transforming mining wastes into construction materials. Alkali-activated materials (AAM) are produced from aluminosilicate rich precursors in alkaline conditions. Glukhovsky [4] proposed the conceptual model of the alkali activation of aluminosilicate materials in three stages: (1) dissolution of aluminates and silicates, (2) reorientation and reorganization, and (3) polymerization and hardening. A subgroup of alkali-activated materials is geopolymers, which consist of SiO$_4$, AlO$_4$, tetrahedra linked with sharing oxygens, while positively charged ions balance the negative charge of Al$^{3+}$ in fourfold coordination [5]. The general precursors for geopolymers are metakaolin, fly ash, and low-Ca slags, which are chemically reactive. Crystalline materials such as clay minerals, mine tailings, and waste rock cannot be directly used as precursors due to their low chemical reactivity.

As mentioned previously, mining waste rocks are crystalline materials that cannot be directly utilized as precursors for alkali activation. Multiples attempts have been made to increase the reactivity of the mineral precursors. Torgal et al. [6] conducted heat treatment on mud waste containing muscovite and found that 11.5% of crystalline muscovite survived after calcination at 950 °C for 2 h. Another study on the activation of raw clay via a mechanochemical process showed that a 25-min ball milling of muscovite-containing clay minerals under optimized grinding parameters can sufficiently disrupt the crystalline structure [7]. However, it should be noted that those clay minerals were originally from Swedish regions, which are known for having “quick clay” that has been skeleton collapsed under high crust strain [8]. This means that the raw materials can be, to some extent, amorphous before mechanochemical activation. Moukannaa et al. [9] prepared a mortar by fusing phosphate mine tailings with 10% NaOH at 550 °C. They found that the compressive strength was 40 MPa after 28 days after the synergy with metakaolin. The aforementioned pre-treatments are relatively time and energy consuming. Although mechanochemical activation by ball milling was successfully implemented on “quick clay,” it is unlikely to disrupt a highly crystalline structure within several minutes. Yao et al. [10] ball milled the highly crystalline muscovite at 500 rpm for 200 min; afterwards, its enhanced pozzolanic reactivity was evaluated by substituting 30% cement in mortar without detriment to compressive strength. However, waste rock is a more complex mixture in which the individual minerals can interact with each other. Consequently, it is important to investigate the amorphization of mica in a complicated system. In the current paper, sulfidic waste rock, resulting from sulfidic ores, was investigated.

Lithium-based grinding aids have been found to display promising results in enhancing the disruption of the micaeous structure (alumino-silicate), thereby improving its chemical reactivity before alkali activation. For example, disruption of the muscovite mica structure [11] has been achieved by immersing mica in a LiCl solution for 24 h before wet-ball milling. Since the late 1990s chemically assisted dry comminution (CADC) has been used as a processing method for improving the surface area of solid materials. The application of CADC on sericite in the presence of lithium nitride accompanied by ion exchange with potassium significantly increases the specific surface area of mica [12]. Solvent-assisted dry grinding is a simpler pathway, in which only a few drops of grinding aid are added without using a large volume of the chemicals involved in both CADC and wet-ball mills.

Therefore, this work aims to investigate the effect of a small amount (0.2 wt%) of grinding aid on mica-containing waste rocks and how it affects the application of mechanochemically activated waste rock as the sole precursor for geopolimerization.

To fully investigate the potential of using sulfidic waste rocks as secondary raw materials for geopolymer production, a simulation-based method was applied by a dynamic process simulator (HSC Chemistry®, developed by Metso Outotec) [13]. A perspective life-cycle assessment (LCA) was conducted for geopolymer production based on simulated upscaling strategies. A process model based on case studies, such as ore processing has been previously investigated [14–18]; however, the simulation of geopolymer binder production has barely been studied before.
Materials and Methods

Waste Rock from Neves Corvo (SOMINCOR by Lundin Mining)

Neves Corvo is an underground high-grade Cu–(Sn)–Zn mine located in the Portuguese sector of the Iberian Pyrite Belt (IPB) in the district of Beja, 220 km southeast of Lisbon (Fig. 1). Copper production started in 1989, followed by tin production between 1990 and 2001, then by zinc/lead production in 2006 [19]. At present, Neves Corvo’s production capacity reaches 2.6 million tons per annum (mtpa) of copper and 1.1 mtpa of zinc and lead combined [20].

Mining activity and the production of base metal concentrates in Neves Corvo generates two different residues: (i) waste rock, the hosting rock extracted and crushed during mining operations, and (ii) tailings, the final residues from the two processing plants. The waste rock consists of variably mineralized felsic volcanic rocks and black schist with disseminated sulfides (mostly composed of muscovite, plagioclase, quartz, chlorite, pyrite, ± chalcopyrite, ± sphalerite, ± gypsum ± carbonates, other oxides, and silicates). The tailings are essentially composed of pyrite (80 to 90 wt%), sphalerite, chalcopyrite, galena, and gangue minerals. Over a 9-year period from 2010 to 2019, the Neves Corvo operation accumulated 7.3 Mt of waste rock stored at the Cerro do Lobo Tailings Management Facility (TMF) and disposed of 17 Mt of thickened tailings, while 3.1 million tons of oxidized waste rock had been stored in a temporary stockpile by the end of 2019. Neves Corvo mine had implemented a series of geochemical and geotechnical strategies where risks associated with potential acid mine drainage (AMD) are kept controlled and stable at the stockpile and the tailings dam, reducing environmental impacts [21].

Materials Preparation

The fresh sulfidic mining waste rock was jaw crushed with a Wedag jaw crusher and double roller crushed using a Mesto Minerals Marcy roller crusher until a particle size lower than 1 mm was achieved for the preparation for the subsequent mechanochemical activation and synthesis of alkali-activated materials. Lithium chloride (BioXtra, ≥ 99.0%) and methanol (99.9%) were purchased from Sigma-Aldrich. Prior to the mechanochemical activation, the waste rocks were ground using a vibratory disk mill (Retsch RS 100) for 1 min at 1500 rpm (dubbed as G1). Mechanochemically activated waste rock was done via grinding with the addition of 0.2 wt% grinding aid (40 wt% LiCl methanol solution) for up to 8 and 16 min (dubbed as G8LiCl and G16LiCl, respectively). The control group was subjected to 8- and 16-min grinding with 0.2 wt% grinding aids (isopropanol; ≥ 99.7%; VWR Chemicals), dubbed as G8 and G16, respectively. The chemical composition of waste rock is given in Table 1, and the sample coding is presented in Table 2.

![Fig. 1 General view of waste rock stockpiles at the Neves Corvo mine (Portugal)](image)

Table 1 Chemical composition of waste rock

| Main component | (Wt%) |
|----------------|-------|
| SiO₂           | 52.31 |
| Al₂O₃          | 12.15 |
| CaO            | 0.56  |
| MgO            | 1.98  |
| K₂O            | 1.49  |
| Fe₂O₃          | 16.69 |
| SO₃            | 12.34 |
| TiO₂           | 0.56  |
| MnO            | 0.13  |
| L.O.I          | 9.7   |

Table 2 Sample coding

| Sample coding | Description                              |
|---------------|------------------------------------------|
| G1            | 1-min grinding waste rock                |
| G8            | 8-min grinding waste rock                |
| G16           | 16-min grinding waste rock               |
| G8LiCl        | 8-min grinding waste rock with LiCl     |
| G16LiCl       | 16-min grinding waste rock with LiCl    |
| IPG16         | Geopolymer produced from G16            |
| IPG16LiCl     | Geopolymer produced from G16LiCl        |
Synthesis of Waste Rock-Based Geopolymer

The mix design is given in Table 3. The alkali activator (molar ratio of SiO$_2$/Na$_2$O = 1.33) was prepared by adding dry sodium hydroxide pellets (VWR Chemicals, > 97%) to sodium silicate solution (VWR Chemicals, SiO$_2$: 26.5%, Na$_2$O: 8%, H$_2$O: 65.5%). No extra water was added. The alkali-activated waste rock was produced by mixing mechanically activated waste rock and an alkali activator using a shear mixer (IKA® WERKE) for 3 min at 700 rpm. Thereafter, the paste was molded in the Teflon molds with dimensions of 2 cm × 2 cm × 8 cm. A vibrating machine (Vortex-Genie 2, Prolab Oy) was used to remove all air bubbles. The sample was demolded after curing under ambient conditions for 24 h and then continuously cured in a plastic bag until the test. This was nominated as IPG16LiCl, where IP and t represent the geopolymer and the curing time, respectively. In this work, the precursor from mining waste rock was relatively rich in aluminosilicates.

Characterization

The chemical composition of mining waste rock was determined using a PANalytical AXiosmAX XRF spectrometer with a rhodium tube, which has a maximum power rate of 4 kW. The particle-size distribution (PSD) and Brunauer–Emmett–Teller (BET) surface area were determined using a laser diffraction particle-size analyzer (LS 13320, Beckman Coulter, Inc., Brea, CA, USA) and ASAP 2000 Micrometrics, respectively. XRD analysis was done using a Rigaku SmartLab 4.5 kW, with the equipment parameters of Co source (40 kV and 135 mA) $K_{α1}(K_{α1} = 1.78892$ Å; $K_{α2} = 1.79278$ Å; $K_{α1}/K_{α2} = 0.5$), scan rate of 3°/min, and 0.02°/step. Quantitative analysis was performed by using the PDXL2 Software Suite with integrated access to the PDF-4 (2019) database and 10 wt% rutile (powder; < 5 μm; ≥ 99.9% trace; metal basis) as the internal standard. The alkaline solubility of the ground powder was evaluated by mixing with a 6 M NaOH solution in polypropylene bottles at a liquid/solid mass ratio of 200/1. A horizontal shaking table (IKA KS 260 orbital shaker) was utilized at Mot = 150/min for 24 h in a moderate environment, which was set at 23 ± 0.5 °C with 50% humidity. Thereafter, the resulting specimens were filtered by using a 0.2 μm filter paper, and the filtrate was acidified with 6 M HNO$_3$ to a pH lower than 1. The concentration of Si and Al elements was measured using the inductively coupled plasma-optical emission spectroscopy (ICP-OES) technique. A vibrational analysis of waste rocks was performed by using diffuse reflectance infrared Fourier transform (DRIFT). The spectra were collected via a Bruker Vertex 80v spectrometer (USA) with a range of 400–4000 cm$^{-1}$, and 40 scans were taken at a resolution of 1 cm$^{-1}$ for each sample. The morphology of the specimen was characterized using Zeiss ULTRA plus (scanning electron microscopy) SEM with an acceleration voltage of 5 kV. An EDS (Energy-Dispersive X-Ray Spectroscopy) analysis was conducted with an acceleration voltage of 15 kV and a beam current of $120 \times 10^{-8}$ A. The polished cross section of the geopolymer sample was subjected to a 50-point analysis under the magnification of × 3000 and a working distance of 6 to 8 mm. The morphological features of ground waste rock samples were measured by a transmission electron microscope (TEM) (JEOL JEM-2200FS, Japan). The preparation of the samples was performed by dispersing a small amount of powder (about 0.05 g) in high-purity ethanol (5 ml); the sample was then vibrated in a sonicated bath for 5 min. A drop of diluted powder suspension was placed onto the standard Cu TEM grid covered with a perforated lacey carbon film. The standard conditions of 200 kV were used during the TEM analysis. Unconfined compressive strength was assessed using a Zwick Roell 100kN machine with a loading force of 2.4 kN/s until failure.

Life-Cycle Assessment (LCA)

System Definition and Functional Unit

LCA is a standardized method for assessing the potential environmental impacts of a product or system. The goal of this study is to evaluate the environmental performance of alkali-activated binder from waste rock in comparison to that of ordinary Portland cement (OPC). Thus, the LCA was calculated by defining the functional unit as the production of “1 kg of binder material”. To make the analysis comparable to that of standard Portland cement, a multiplier factor was introduced as a substitution ratio [22, 23], which is formulated as follows:

Substitution ratio = $\frac{28\text{-day compressive strength (MPa) of geopolymer}}{28\text{-day compressive strength (MPa) of base case material}}.$

Table 3  Mix design of geopolymer

| Sample name | Waste rock (wt%) | NaOH (wt%) | Sodium silicate solution (wt%) | Water/binder ratio |
|-------------|-----------------|------------|-----------------------------|-------------------|
| IPG16LiCl   | 73.15           | 1.74       | 25.11                       | 0.2               |

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When comparing the LCA results of the developed geopolymer binder and Portland cements, the impacts of the geopolymer were divided by the substitution ratio above, (i.e., 0.26). All the environmental impacts in this study were evaluated using the following six midpoint methods: (1) global warming potential (IPCC 2013), (2) cumulative energy demand, non-renewable (CED), (3) ReCipe water depletion, (4) ReCipe particulate matter formation, (5) CML 2002 resource depletion, (6) CML 2002 acidification potential [24], and (7) ReCipe 2016 Endpoint (H) to account for overall impacts across categories [22].

Data Collection and Life-Cycle Inventory (LCI) Calculations

The first step comprises a conversion of lab protocols into a lab-scale process diagram (Fig. 2). Central to this step are minerals and particle-size characterizations, i.e., mineralogy and grain sizes (refer to electronic supplementary materials, Supplementary Fig. S1, Tables S1, S2), and the results are obtained from experimental steps in “Characterization” section. The information was used as necessary input data in the subsequent step, mass, and energy balance computer simulation, within HSC Sim 10 [25].

In the second step, various possible large-scale process flow diagrams were accounted by building three configurations for the mechanochemical activation of waste rocks by translating lab-scale processes to their equivalent industrial equipment (see Supplementary Table S3, Figs. S2–S4). These three configurations represent expert and technology experts’ knowledge in the upscaling aspects of this study. When looking for equivalent industrial-scale processes, the equipment was chosen based on the functionalities and performances that align with actual requirements. For example, the lab-scale jaw crusher, double-roller crusher, sieving equipment, and vibratory disk mill are used as the main comminution equipment on a large scale. Specifically, for geopolymer binder making, the mix design ratio, as shown in Table 3, is kept the same for both lab-scale and all large-scale configurations.

In the third step, model configurations were constructed in HSC Sim 10 by providing input data from experiments and external resources, i.e., industrial data and expert opinions (see Supplementary Section B for design considerations). The established flowsheet models embrace a range of controls, such as working parameters, residence time, and feed properties [26]. It is notable that the modularity of unit processes in HSC Sim 10 allows for parameterizations in the models. In combination, the previous three steps are completed to generate the mass and energy balance that will be used for the life-cycle inventory of the study. The generated inventories are then transferred to the LCA modeling software for calculating the environmental impacts using the life-cycle impact assessment methods chosen in the previous section. In this study, Activity Browser software was used to run the LCA modeling [27]. The background data, such as European electricity mix and chemicals usage in the technosphere, were taken directly from the well-established LCA database Ecoinvent 3.7 [28], while all foreground data were obtained from the results of flowsheet simulations. It was assumed that the production facilities were near the waste rock storage facility; hence, the transporting efforts were excluded. The overall LCA procedure and the process for conducting the analysis for comparative purposes are shown in Fig. 3.

Results and Discussion

PSD and BET

The cumulative particle-size distribution curve displays the difference between G1, G16, and G16LiCl (Fig. 4). Grinding without the aid of chemicals hardly increases the BET surface area and results in only a small decrease in particle size. Lithium chloride can efficiently enhance mechanochemical activation, whereby almost 98% of particles are smaller than 20 µm and 50% smaller than 3 µm. The corresponding BET surface area increases from 2.10 to 18.01 m²/g, whereas the BET surface area of the samples milled without grinding aid only increases to 3.81 m²/g. The effect of mechanochemical activation also results in the amorphization of minerals in the waste rock. The grinding aid has a significant influence on aluminosilicate minerals, such as muscovite and clinochlore. On the contrary, stable minerals, such as quartz and pyrite, are kept intact during mechanochemical activation. The morphology depicts samples with markedly different features. G8 and G16 display a pristine muscovite lamellar structure, while G8LiCl and G16LiCl have irregularly shaped particles (Fig. 5).
The XRD patterns were identified using PDXL2 software (Fig. 6a). These phases include quartz (ICDD; PDF 04-014-7569), pyrite (ICDD; PDF 04-004-6511), muscovite (ICDD; PDF 04-016-3068), clinochlore (ICDD; PDF 04-022-2452), and gypsum (ICDD; PDF 04-010-9409) with internal standard rutile (ICDD; PDF 04-008-7847). The structure of layered aluminosilicate (muscovite and clinochlore) is considerably disrupted after a 16-min grinding with LiCl, while ground aluminosilicates without LiCl remain intact, comparable to the untreated waste rock. The XRD pattern of G16LiCl shows line broadening, and the peaks of layered aluminosilicate almost disappeared in comparison with G16 and untreated waste rock. The results of quantitative XRD analysis further confirm the diffraction line broadening and decrease in intensity in XRD patterns (Fig. 7). The quantitative analysis errors may result from the heterogeneity of waste rock during crushing and grinding. The DRIFT patterns reflect the structural alteration of aluminosilicate minerals in Fig. 6b. The Si–O stretching band at 1062 cm\(^{-1}\) and Si–O bending at 520 cm\(^{-1}\) disappear, indicating the collapse of silicate structure. In addition, a splitting band appears at 1465 cm\(^{-1}\) and 1277 cm\(^{-1}\). The heavy distortion during grinding results in a misfit between octahedral and tetrahedral sheets, thereby resulting in tetrahedral rotation [29]. The bands noticeable at 1670 cm\(^{-1}\) and 1610 cm\(^{-1}\) are assigned to molecular water bending modes, while the bands between 2200 and 1700 cm\(^{-1}\) correspond to the harmonic combination of bands between 400 and 1000 cm\(^{-1}\) [30]. For clinochlore, the OH liberation band around 600 to 800 cm\(^{-1}\) remains for all ground samples, as is the case for heat-treated/mechanically treated clinochlore [31–33]. Nevertheless, the decreased intensity also occurs in clinochlore: the 3550 cm\(^{-1}\) band practically vanishes after grinding with LiCl grinding aid. The stretching vibration of absorbed water molecules is ascribed to a broad band at around 3430 cm\(^{-1}\). There was no significant difference seen when treated with 16-min grinding. However, the broad band at 933 cm\(^{-1}\) for Al–OH disappears, a result that differs from the heat-treated case [34]. The band at 3627 cm\(^{-1}\) is assigned to the OH stretching absorption of muscovite, during which the intensity decreases [35]. In conclusion, dehydroxylation occurred during the mechanochemical process, resulting in a change...
in aluminum coordination for further geopolymerization with silicate species.

In muscovite, $\text{KAl}_2[\text{AlSi}_3\text{O}_{10}(\text{OH})_2]$, a quarter of $\text{Si}^{4+}$ is substituted by $\text{Al}^{3+}$, leading to a negative unit charge, which is filled in by a $\text{K}^+$. The mineral is also known as dioctahedral mica since $\text{Al}^{3+}$ occupies the M2 octahedral sites, while the M1 octahedral sites are vacant [36]. There are two possible ways for lithium ions to influence the amorphization of the waste rocks: (1) $\text{Li}^+$ shows a tendency to enter the vacant position in the dioctahedral layer of mica, thereby changing its charge balance and expelling $\text{K}^+$ from the interlayer position [37]; (2) $\text{Li}^+$ has the smallest cation size and highest electronegativity out of all alkali metals, causing it to enter into the interlayer and push $\text{K}^+$ out. Figure 8 shows a schematic diagram of the function of the lithium cation during grinding. High-energy grinding impacts the structure of mica not only through delamination but also by the dislocation of the layered structure of mica upon incorporation of the lithium cation. Furthermore, only a small amount (0.2 wt%) of the grinding aid can affect the layered structure.

**TEM**

TEM is a vital tool for investigating detailed features of crystalline and amorphous phases. As reported previously, mica is sensitive to electron beams which means it can decompose under electron radiation [38]. Therefore, the radiation damage rate (RDR) should be considered during TEM.
measurement. The diffraction pattern is obtained within 90 s because the RDR of muscovite is classified between medium and fast. A high-resolution transmission electron microscope (HRTEM) can reflect the mineral structure change on a nanoscale and disordered mica can be found in a sample ground for 16 min (Fig. 9). The figure also displays associated selected area diffraction (SAD) patterns from both G8LiCl and G16LiCl, and G16LiCl explicitly shows SAD patterns with diffuse rings from the completely amorphous region at b’s center. A combined diffused ring with a matrix spot pattern appears in G8LiCl, and this indicates the intermediate transmission from the crystalline phase to the amorphous phase. The lattice parameters of both samples were determined by XRD diffraction. Results indicate that a considerable change occurs in the c axis, that is, probably due to the compression along the (001) cleavage plane, leading to structural distortion [39].

Arrays of the (001) plane have rotated in the boundary across for the diffraction pattern of G8LiCl, the misorientation of (001) becomes slightly rotated, meaning that the complex interleaving of the boundary has occurred, as reported in the literature [38]. This result is consistent with the XRD analysis results indicating that the (001) reflection becomes a hump after 8-min grinding. The amorphous halo of G16LiCl indicates that there is only a partial orientation in the muscovite structure, i.e., it partially loses crystallinity after 16 min of intense grinding with LiCl media. In addition, a delaminated and irregular particle can be seen in the HRTEM (Fig. 9b).

**Alkali Activation Potential**

The alkaline dissolution results show enhanced aluminum and silicon dissolution (Fig. 10a). Since Si is located in a stable Si–O bonding environment within quartz and layered aluminosilicates, a lower enhancement of Si solubility can be seen (Fig. 10a). Al solubility experiences a small increase after grinding without LiCl; however, after grinding with LiCl for 16 min, Al solubility shows up to a 40-fold increase. The XRD reflection patterns and Rietveld refinement of ground samples after solubility tests are shown in Supplementary Figs. S5–S8. There are few changes seen for both G1 and G16, while G16LiCl exhibits a considerable amount of amorphous phase (43.1%). Si solubility also increases tenfold for G16LiCl, and dissolved silicate units come from layered aluminosilicates, as quartz is chemically inert. The morphology of samples after the dissolution test is shown in Supplementary Figs. S9 and S10. G16 retains its regular particle shape, while G16LiCl exhibits an amorphous gel-like structure after the alkaline dissolution test. G16 particles retain their regular shape with clear grain boundaries, and the major elements (Al, Si, and K) are homogenously distributed without conspicuous dissolution. In contrast, higher Al and Si dissolution occurs for G16LiCl particles during the dissolution test, thereby indicating an amorphous
structure without a uniform elemental distribution and clear grain boundary.

Waste rock-based alkali-activated materials were prepared, and the compressive strength was measured on days 7 and 28. IPG16LiCl28 displays a moderate 28-day compressive strength of 10.3 MPa (Fig. 10b). Furthermore, it meets the minimum requirement of compressive strength for bricks under negligible weathering conditions according to ASTM
The inert minerals, such as quartz and pyrite, fill in the matrix as fine aggregates. Although 40% Al solubility is reached, the initial amount of Al is rather low (13 wt%), which results in a moderate compressive strength in this case. It should be mentioned that the sample IPG16 was prepared with untreated waste rock. However, the sample did not harden when cured at room temperature and maintained its slurry form at day 28. Moreover, oxidized pyrite transforms to mobilized sulfate, increasing the possibility of acid leaching, which requires a further leaching assessment of the resultant geopolymer.

**SEM and EDS of Waste Rock-Based Geopolymer**

The 28-day sample was subjected to an SEM analysis. The dense matrix may be correlated with an N-A-S-H-type gel (area 2 in Fig. 11). The term geopolymer will be used to refer to the N-A-S-H gel in this study, while the waste rock-based geopolymer represents the complex combination of geopolymer and inert fine aggregates such as pyrite, quartz, and unreacted mica. Area 1 of the ternary diagram shows quartz and aluminosilicates, since not all aluminosilicates can be transformed into an amorphous phase after mechanochemical activation. The dissolved Al species can react with silicate species from both dissolved Si and sodium silicate solution to form N-A-S-H gel, as displayed in the ternary diagram. It is important to note that the Na/Al ratio was held constant at 1:1 to achieve the overall charging balance of the resulting geopolymer. The Si mapping indicates the highly crystalline silica, and the black dots are attributed to inert pyrite crystals. Al mapping shows the distribution of geopolymeric gel since Al is dissolved from the amorphous phase. This result is in line with the alkaline solubility test where a large proportion of aluminum dissolves in the presence of an alkali solution. The dissolved aluminate species can react with silicate species in the subsequent condensation, where most of the silicate units come from the sodium silicate solution. Therefore, the condensation was dominated by the reaction between aluninate and silicate which are mostly oligomeric [41]. The ternary diagram indicates the formation of the N-A-S-H gel combined with unreacted aluminosilicates, in which the formation of N-A-S-H gel involves the dissolution of aluminosilicates and geopolymerization, i.e., precipitating and polymerizing to a three-dimensional framework.

**Simulation of Grinding Process and Life-Cycle Assessment**

Figure 12 shows the compilation of results specifically for global warming potential and water depletion. The results for other categories are depicted in Table 4. An industrial-scale
Fig. 12 The global warming potential of geopolymer binder production a and the share of impacts b. Part c compares the water depletion to produce the same output with different configurations.

Table 4 The environmental impacts of supplying 1 kg binder material across different impact assessment methods

| Process option to produce 1 kg binder | IPCC 2013, in CO₂-eq | CED non-renewable, fossil in MJ | ReCipe water depletion, in m³ | ReCipe particulate matter formation, in kg PM10-eq | CML acidification, in kg SO₂-eq | CML resource depletion, in kg Sb-eq | ReCipe endpoint, in Pt |
|--------------------------------------|-----------------------|----------------------------------|-------------------------------|---------------------------------------------|---------------------------------|----------------------------------|-------------------------|
| Route 0: geopolymer lab scale         | 4.43                  | 52.62                            | 2.8E−02                       | 7.5E−03                                     | 3.1E−02                         | 2.1E−02                         | 0.469                   |
| Route 1: upscaled geopolymer with thickener | 0.57                  | 5.88                             | 9.7E−03                       | 1.1E−03                                     | 3.4E−03                         | 3.1E−03                         | 0.063                   |
| Route 2: upscaled geopolymer with pressure filter | 0.51                  | 5.26                             | 4.4E−03                       | 1.0E−03                                     | 3.1E−03                         | 2.8E−03                         | 0.057                   |
| Route 3: upscaled geopolymer with double sieving | 0.49                  | 5.01                             | 4.3E−03                       | 9.7E−04                                     | 2.9E−03                         | 2.7E−03                         | 0.054                   |
| Conventional route: Portland cement   | 0.86                  | 3.03                             | 1.0E−03                       | 1.0E−03                                     | 1.7E−03                         | 1.7E−03                         | 0.053                   |
| The optimum (Route 3) compared to Portland cement | 43%                   | 65%                              | 319%                          | − 6%                                         | 75%                             | 59%                             | 2%                      |
flowsheet grinding process simulation was conducted, and this was modeled in line with the laboratory-scale experimental data [42]. Drastic reductions of more than 90% are expected from the upscaling of geopolymer (route 0 to the respective three large-scale configurations). The conversion of equipment from pulverizers to a set of crushing and milling equipment manages to minimize the amount of electricity needed, decreasing energy consumption in the first step of the process, i.e., mechanochemical activation. Similar to the modeling of other lab-scale technologies, greenhouse gas emissions also decrease due to scaling effects of electricity consumption [43, 44]. Finally, when compared to the production of ordinary Portland cement, all geopolymer production routes can compete with the average 40% global warming potential (GWP) reduction potentials, except for the lab-scale scenario (Fig. 12a). It is important to note that with geopolymer routes, one can eliminate the calcination process that contributes to most of the emissions in the making of Portland cement. Once the system reaches common industrial throughput (i.e., 100 ton/h), the contribution towards GWP is almost equally shared between precursor production and geopolymer binder manufacturing (Fig. 12b).

Nevertheless, when it comes to the six other impact categories, the results are different (Table 4), barring particular matter formation. Although the second and third routes can minimize the water consumption in the grinding circuit (Table 5), the consumption is still higher in all scenarios compared to the production of Portland cement. The other two impact categories also reflect similar trends. The use of lithium chloride as a grinding aid as well as alkali activators (e.g., sodium hydroxide and sodium silicate solution) results in higher resource consumption from the production of such chemicals. One can mitigate these issues by sourcing alkalis from underutilized waste to avoid the need for primary alkali productions in the system [45]. The other cause is due to the definition of the functional unit for this study. With the relatively low mechanical performance of geopolymer developed in our study, the environmental impacts are amplified to almost four times that of OPC. Therefore, even though this specific mine waste and recipe cannot be sustainably scaled up, this amount of mechanochemical treatment and chemical use may be justified, given that an adequate mechanical strength is reached. As this system has not been optimized, this result is expected. The mechanical strength can be increased via further optimization and by extending the assessment boundary until the production of concrete or mortar [46]. With such a methodological choice, the mechanical strength in the geopolymer pathways will increase, and the sustainability benefits can also be better justified. Thus, the study also implies the following promising avenues for further investigations: (i) using the developed geopolymer binders to produce concrete/mortars, (ii) formulating new recipes for alkali activating geopolymers, and (iii) a combination of both strategies. With the right alkali formulations and mix designs [47], the developed geopolymer concrete can compete with standard cement concrete from an environmental perspective. Lithium-containing grinding aid is suitable for lab-scale production; however, it can be critical when upscaling the valorization of millions of tons of waste rock. From the perspective of a circular economy, the recycling of lithium from the battery industry or the seeking of an alternative cheap grinding aid for an industrial application can be the subject of future endeavors.

### Conclusion

In summary, this study indicates that sulfidic mining waste rock can be a potential candidate for the fabrication of geopolymers after mechanochemical activation. Lithium chloride grinding aids can significantly improve the amorphization of the waste rock during grinding, thereby enhancing Si and Al dissolution. In contrast, ground waste rock with typical grinding aid (e.g., isopropanol) retains the waste rock’s crystalline structure. This coincides with the generation of an amorphous phase in which chemically assisted mechanochemically activated samples possess more amorphous phases than mechanochemically treated samples regardless of grinding time. The amorphous aluminosilicates (e.g., muscovite, clinochlore) resulting from structural distortion/dislocation were obtained after mechanochemical treatment in terms shown by the XRD and TEM analyses. The optimal unconfined compressive strength for geopolymer produced from 16-min grinding waste rock with LiCl (G16LiCl) is 10.27 MPa after 28 days of curing under ambient conditions. Despite the complexity of mineralogy, a proper pre-treatment method should be considered before designing the activation route. It should be noted that similar mineralogy (e.g., aluminosilicate-bearing clay resources) can express different stoichiometry compositions and degrees of crystallinity due to mineralization, weathering, and crust conditions (geological origin).

Simulation can be a versatile tool when upscaling lab work to an industrial scale, as it offers a vital method for

| Model type | LCA indicators (For production of precursor 1 kg) |
|------------|--------------------------------------------------|
|            | The summarized data in this table are taken from simulation results in Supplementary Figs. S2–S4 |
|            | Precursor products (kg) | Water recovery (kg) | Fresh water consumption (kg) | Electricity (kWh 10⁻³) |
| Route 1    | 1 | 0.00 | 1.17 | 25.06 |
| Route 2    | 1 | 0.72 | 0.16 | 21.36 |
| Route 3    | 1 | 0.68 | 0.15 | 20.76 |
minimizing costs and predicting the results. When life-cycle assessment is implemented, the environmental impact of the simulated process can pinpoint the advantages and disadvantages of several design choices. Indeed, the environmental performance of geopolymer production increases drastically due to optical energy minimization, a result that is parallel to the continuous proposal of sustainable alkali activators and mix strategies in geopolymer production.

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Declarations

Conflict of interest The authors declare that they have no conflict of interest.

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