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Critical mineral demands may limit scaling of green hydrogen production

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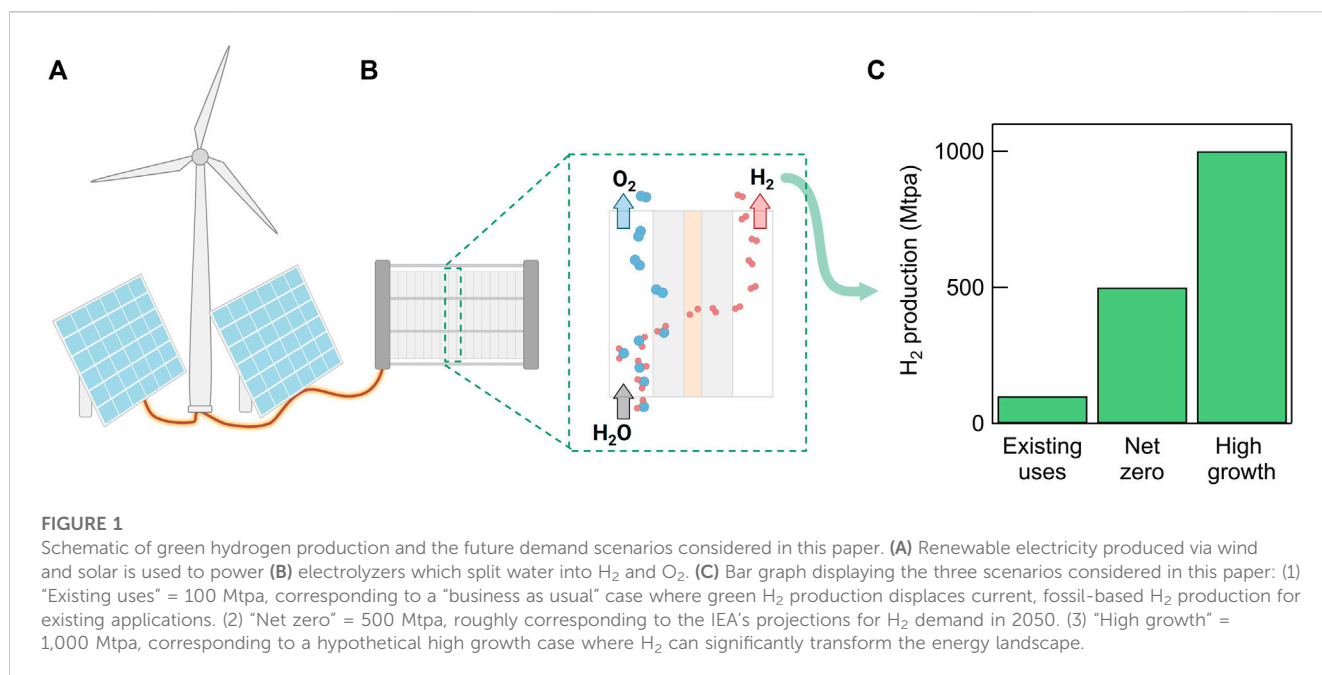
Hydrogen (H₂) is widely viewed as critical to the decarbonization of industry and transportation. Water electrolysis powered by renewable electricity, commonly referred to as green H₂, can be used to generate H₂ with low carbon dioxide emissions. Herein, we analyze the critical mineral and energy demands associated with green H₂ production under three different hypothetical future demand scenarios, ranging from 100–1,000 Mtpa H₂. For each scenario, we calculate the critical mineral demands required to build water electrolyzers (*i.e.*, electrodes and electrolyte) and to build dedicated or additional renewable electricity sources (*i.e.*, wind and solar) to power the electrolyzers. Our analysis shows that scaling electrolyzer and renewable energy technologies that use platinum group metals and rare earth elements will likely face supply constraints. Specifically, larger quantities of lanthanum, yttrium, or iridium will be needed to increase electrolyzer capacity and even more neodymium, silicon, zinc, molybdenum, aluminum, and copper will be needed to build dedicated renewable electricity sources. We find that scaling green H₂ production to meet projected net-zero targets will require ~24,000 TWh of dedicated renewable energy generation, which is roughly the total amount of solar and wind projected to be on the grid in 2050 according to some energy transition models. In summary, critical mineral constraints may hinder the scaling of green H₂ to meet global net-zero emissions targets, motivating the need for the research and development of alternative, low-emission methods of generating H₂.

KEYWORDS

hydrogen, green hydrogen, critical minerals, electrolysis, solar, wind, renewables

1 Introduction

Hydrogen (H₂) is widely viewed as critical to the decarbonization of industry and transportation. Currently ~100 million tonnes per annum (Mtpa) of H₂ are produced globally (IEA, 2023a), the primary uses of which are petroleum refining, fertilizer production, and oxygen (O₂) scavenging in metal treatments (Ramachandran and Menon, 1998; IRENA, 2018; EIA, 2023). Many proposed decarbonization strategies rely on expanding the uses of H₂ to include use as a chemical reductant in industrial sectors, such as steel refining, and as a chemical fuel for heavy-duty transportation and shipping, either directly via fuel cells or indirectly via ammonia (Hydrogen Council, McKinsey & Company, 2021; IEA, 2023b). However, the majority of H₂ is currently produced from fossil-based feedstocks, such as steam methane reforming (~60 Mtpa) or coal gasification (~20 Mtpa), which yields a carbon footprint of ~10–20 kg carbon dioxide equivalent (CO₂eq)/kg H₂ (IEA, 2023b) and ~900 Mtpa of direct CO₂ emissions. Developing low-emission methods of producing H₂ and scaling the total volume of H₂ produced are required to realize significant CO₂ abatement via new H₂-related applications. Water electrolysis powered by renewable electricity, commonly referred to as “green



hydrogen”, is one method of generating H₂ with low CO₂ emissions. Although water electrolysis has historically been associated with higher costs (van Renssen, 2020; Yu et al., 2021), it is generally considered the most viable technology for producing low-emission H₂ and reaching net-zero emissions targets by 2050 (Castelvecchi, 2022; IRENA, 2022; IEA, 2023a). More than 40 governments around the world have H₂ strategies in place and many offer incentives for green H₂ production to make it more cost competitive with fossil-based production methods (DOE, 2023a; IEA, 2023a).

Herein, we analyze the critical mineral and energy demands associated with green H₂ production. We use the term “critical minerals” to refer to all “critical materials for energy,” as defined by the United States Energy Act of 2020 (Bauer et al., 2023). This definition includes elemental metals, alloys, and natural minerals, though the following analysis mostly focuses on elemental metals. We include both the minerals required to construct dedicated renewable electricity sources (Figure 1A) and the minerals required for water electrolyzers (Figure 1B). In this analysis, we consider the following three scenarios for future green H₂ demand:

- (1) 100 Mtpa, corresponding to an “existing uses” (or “business as usual”) case where green H₂ production displaces current, fossil-based H₂ production for existing uses and applications (Hydrogen Council, McKinsey & Company, 2023):
- (2) 500 Mtpa, corresponding to a “net-zero” case with increased demand and applications for H₂. This is based on the average projected demand for H₂ in 2050 according to the International Energy Agency (IEA)’s Net Zero Roadmaps (IEA, 2021a; 2023b), which forecast that H₂ will primarily be produced via renewable electricity in 2050.
- (3) 1,000 Mtpa, corresponding to a hypothetical “high growth” scenario where increased H₂ production could significantly transform the energy landscape, *i.e.*, H₂ could potentially be widely used as a fuel and replace natural gas. In this scenario, H₂

plays a more significant role in energy storage and transportation as total demand reaches the gigatonne scale (Energy Transitions Commission, 2020; Oliveira et al., 2021).

2 Water electrolyzer analysis

Water electrolyzer technologies use metallic electrode catalysts to convert water into H₂ and O₂ under an applied electrical current. Alkaline water electrolysis (AE) and solid oxide electrolysis cells (SOEC) also use metal oxides in electrochemical separators. Significant research efforts have been devoted to increasing electrolyzer efficiency and the discovery of new materials for electrolysis technology (Roger et al., 2017; Hauch et al., 2020; Tong et al., 2020; Chatenet et al., 2022). The first commercial water electrolysis technology to be developed was AE. While AE is relatively inexpensive and typically uses nickel (Ni) catalysts and modern separators comprising zirconium dioxide (ZrO₂) in a polysulfone matrix, the limited current density, gas crossover, and corrosion of the Ni-based electrodes led to the development of membrane-based electrolysis. In proton exchange membrane (PEM) electrolysis, a solid polymer electrolyte composed of perfluorosulfonic acids (PFSA) is directly integrated with the electrodes. While PEM allows for higher current densities and increased H₂ purity, PEM uses platinum group metals (PGMs, *e.g.*, platinum, Pt, palladium, Pd, and iridium, Ir) as electrode catalysts, making the raw materials more expensive than those required for AE. Both AE and PEM operate at relatively low temperatures (<100°C) compared to SOECs, which operate at temperatures >700°C. SOECs may enable higher efficiency H₂ production (Ni et al., 2008), but require transition metals and rare earth elements (REEs). Typically, SOEC electrodes are made from composites of Ni with yttrium (Y)-stabilized-zirconia and use strontium

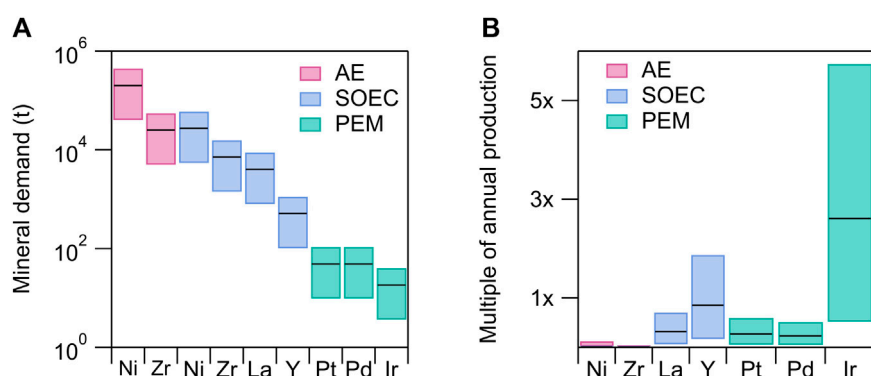


FIGURE 2

For each floating bar, the lower and upper limits correspond to the quantity of H₂ required for the “existing uses” (100 Mtpa) and “high growth” (1,000 Mtpa) scenarios, respectively. Black lines correspond to the “net zero” (500 Mtpa) scenario (defined in Figure 1C). (A) Total mineral demand required to produce 100–1,000 Mt H₂ using different electrolyzer technologies: Alkaline water electrolyzers, AE (pink), solid oxide electrolysis cells, SOEC (blue) and proton exchange membranes, PEM (green). Values are calculated according to Eq. 1 assuming the leveled mineral demand for each technology (IEA, 2021b) and target electrolyzer efficiencies of 45 kWh/kg H₂ for AE (DOE, 2023b) and PEM (DOE, 2023c), and 40 kWh/kg H₂ for SOEC (DOE, 2023d). (B) Mineral demand normalized to annual, global production values (USGS, 2023). Note that in (B) the AE and SOEC values for Ni and Zr are stacked on top of each other but are too small to be observed on the linear scale shown.

(Sr)-doped lanthanum (La)-based manganese (Mn) or iron (Fe) oxides as separator materials.

We assess the demand for select minerals used in the aforementioned electrolyzers (AE, PEM and SOEC) in Figure 2, according to the three H₂ production scenarios described above. The selected metals are essential to each technology’s performance and currently cannot be easily substituted. We use reported leveled mineral demands (IEA, 2021b) and assume each electrolyzer supplies 100% of global green H₂ demand. Each electrolyzer type is assumed to operate at the system efficiency targets set by the U.S. Department of Energy: 45 kWh/kgH₂ for AE (DOE, 2023b) and PEM (DOE, 2023c), and 40 kWh/kgH₂ for SOEC (DOE, 2023d). We approximate the total mineral quantities required for each H₂ demand scenario following Eq. 1:

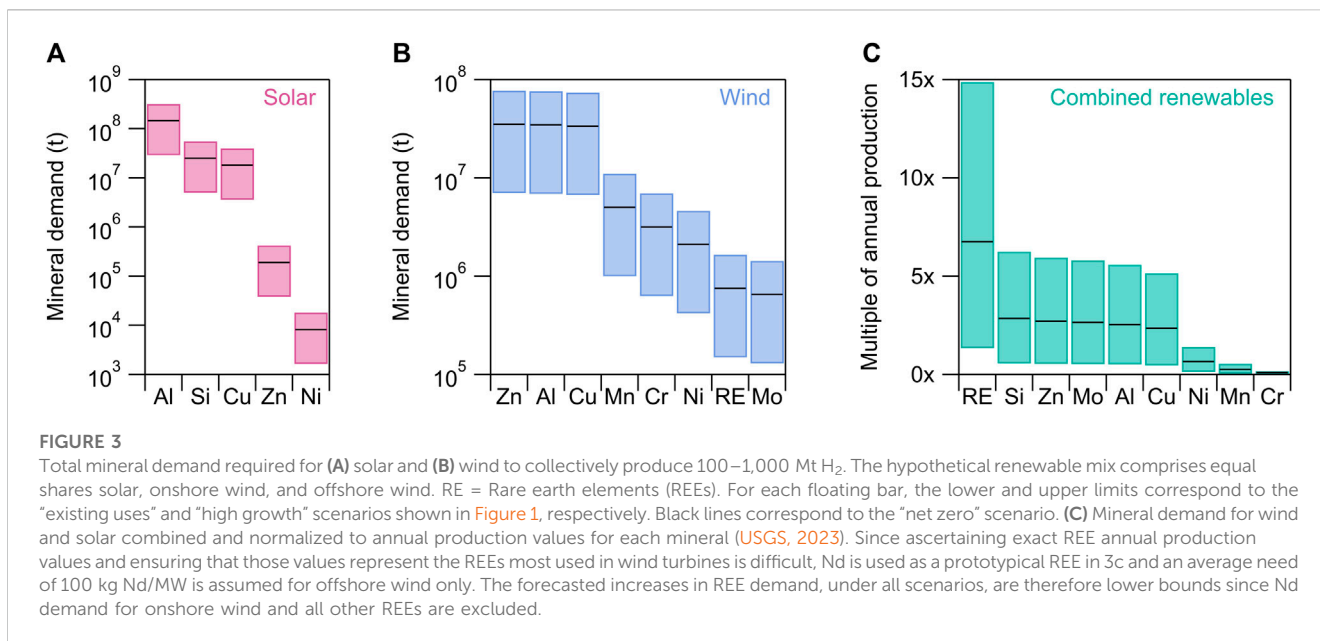
$$\text{Mineral demand} \left(\frac{\text{Tonnes mineral}}{\text{GWh output}} \right) \times \text{Electrolyzer efficiency} \left(\frac{\text{GWh}}{\text{Mt H}_2} \right) \times \text{H}_2 \text{ demand (Mtpa H}_2) = \text{Tonnes mineral} \quad (1)$$

PEM electrolyzers require the least amount of raw material, requiring less than 500 tonnes of PGMs to generate total anticipated electrolyzer capacity (Figure 2A), even in the “high growth” scenario (1,000 Mtpa H₂). However as all current PEM technologies rely on PGMs, PEM becomes the most resource constrained (Figure 2B) when demand is normalized to current global annual production values (USGS, 2023). Even under the “existing uses” scenario (100 Mtpa H₂), half of all annual global Ir production is needed to build sufficient PEM electrolyzer capacity. We acknowledge that the electrolyzers would only need to be built once, yet we are normalizing to mineral production on a per annum basis. However since green H₂ currently accounts for less than 1% of global H₂ production (IEA, 2023a), the projected growth of PEM electrolyzers would represent new demand for Ir and other PGMs. Scaling PEM electrolyzers to the “net zero” or “high growth” scenarios would result in vastly higher (~2.5–5×) Ir demand. The high cost of PGMs will likely lead to PEM being more expensive than

other electrolyzer technologies. In addition, potential regulations and restrictions surrounding the production of fluorinated compounds, such as PFSA (Lim, 2023), increase the uncertainty of PEM as a long-term technology solution for water electrolysis.

Current SOEC technologies are potentially constrained by two REEs, La and Y (Figure 2). The limited distribution of REE reserves, challenges associated with separating and purifying specific minerals, and increasing competing demand for these elements from other clean energy technologies make scaling SOECs to meet higher H₂ demands potentially difficult. Compared to PEM and SOECs, AE is the least resource constrained, as it requires only a small fraction of total Ni and Zr annual production. However, AE is the most resource intensive in terms of the total tonnage of materials required, representing potentially a higher CO₂ footprint associated with producing the materials for these electrolyzers (Azadi et al., 2020). Our findings suggest securing sufficient PGMs and REEs will be a key challenge to scaling green H₂ production and are consistent with recent analysis of U.S. electrolyzer supply chain vulnerabilities (Badgett et al., 2022).

Technological improvements, such as anion exchange membrane (AEM)-based electrolysis, may alleviate some of these potential supply constraints. AEM electrolysis combines the inexpensive and more abundant mineral benefits of AE with the efficiencies of PEM, making it an attractive electrolysis technology under development (Du et al., 2022). While AEM device lifetimes are currently less than those of AE and PEM (Mustain et al., 2020), active research into improving membrane stability and recovering metals from degraded devices may make AEM a viable path for future water electrolyzers with low mineral demands and high performance. On the other hand, regardless of electrolyzer technology, system efficiency is limited by the thermodynamic limit of ~39 kWh/kg H₂, assuming standard temperature and pressure (Levene et al., 2007).



3 Renewable electricity analysis

We next consider the mineral demands that arise from building dedicated renewable electricity sources to power water electrolyzers. Currently, water electrolysis using the existing, fossil-based global grid can produce greater CO₂ emissions than directly producing H₂ from fossil fuels (IEA, 2023c). Therefore, either “behind-the-meter” renewable systems, *i.e.*, electrolyzers directly connected to renewable electricity sources, or an electrical grid based solely on a renewable mix are required to enable low-emission water electrolysis. For simplicity, we assume a hypothetical future energy supply comprising equal parts solar, onshore wind, and offshore wind. We calculate the capacity factor (CF) for each electricity source by taking the ratio of 2021 global generation (GWh) to 2021 global capacity (GW), as shown in Eq. 2 (IRENA, 2023a). This yields global CFs of 14%, 25%, and 29% for solar, onshore wind, and offshore wind, respectively. These CFs agree with global and regional analyses (Tong et al., 2021; Bolson et al., 2022), though higher CFs can be achieved in select regions. For example, in Chile’s Atacama Desert >30% CF can be achieved with standard photovoltaics and >80% with concentrated solar power (Gallardo et al., 2021), and >50% CF can be achieved for offshore wind in Scotland (Carreno-Madinabeitia et al., 2021).

$$\left(\frac{\text{Generation (GWh)}}{\text{Capacity (GW)}} \right) \times \left(\frac{1}{8760 \text{ hours per year}} \right) = \text{Capacity factor (CF)} \quad (2)$$

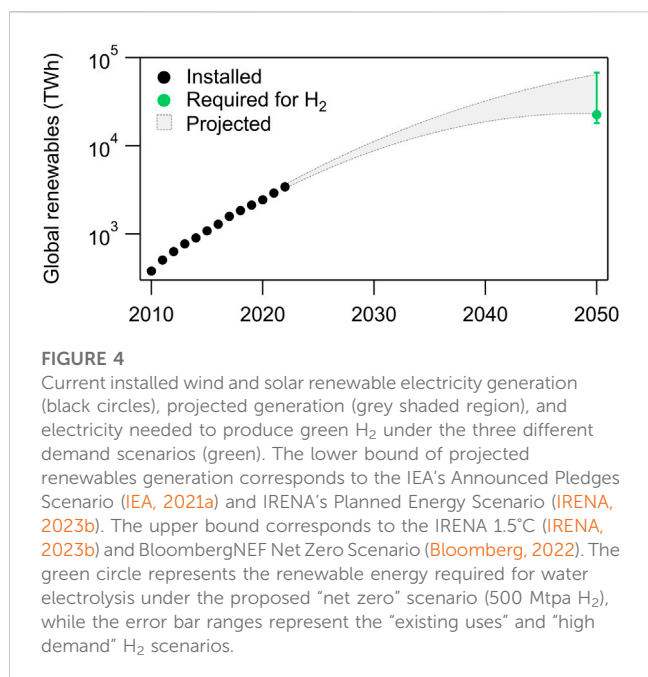
Assuming a target system efficiency of 45 kWh/kg H₂, total mineral demand (Figures 3A,B) is calculated by multiplying the levelized mineral demands for solar and wind (IEA, 2021c) by the energy demand required for each H₂ scenario, as given in Eq. 3. Offshore and onshore wind are binned together as they share similar mineral requirements, though the quantities required for specific minerals may vary.

$$\text{Mineral demand} \left(\frac{\text{Tonnes mineral}}{\text{GW}} \right) \times \left(\frac{1}{\text{CF} \times 8760 \text{ hours}} \right) \times \text{H}_2 \text{ demand (GWh)} = \text{Tonnes mineral} \quad (3)$$

The total tonnage of minerals required for renewable energy (Figure 3) is much greater than the quantity required for electrolyzers (Figure 2). For example, the Ni needed for AE under the “high demand” scenario (~0.40 Mt) is roughly equivalent to the total amount of Ni required for solar and wind under the “existing uses” scenario (~0.002 Mt for solar and ~0.4 Mt for wind). The contrast is even starker for PEM and SOECs, where the mass of all minerals required for electrolysis is several orders of magnitude less than the mass of minerals needed for renewables.

As with the electrolyzer analysis, we normalize renewable energy mineral demand to current annual production values (Figure 3C) (USGS, 2023). We bin together the shared minerals (Al, Cu, Ni, and Zn), assuming equal energy contribution of solar, offshore wind, and onshore wind. Normalizing total demand to current annual production highlights the minerals of particular concern. Higher quantities of Si, Cu, Al, Zn and Mo will be required, ranging from ~0.5–5× annual production, pending the H₂ demand scenario. Though elemental Si and Al are abundant, both depend on energy- and emission-intensive refining processes (Maldonado, 2020; Lennon et al., 2022; Bauer et al., 2023). Greater electrification in other sectors will also increase demand for Si and Al. The increase in demand for Cu is also noteworthy, as it is the most common conductor for wiring (He and Small, 2022), and critical for global renewable generation and electrification efforts, yet global ore grades are in decline (Northey et al., 2014).

While solar has seen the highest growth in capacity over the last decade (IRENA, 2023a), its relatively low global CF (~14%) makes it challenging to use for continuous H₂ production, absent dedicated energy storage systems or dynamic electrolyzers. Offshore wind has the highest CF (~29%), making it an attractive renewable electricity source with lower intermittency. However, offshore wind requires



more Cu than onshore wind due to longer cable lengths. Similarly, more REEs are required for offshore wind due to the greater dependence of offshore turbines on strong permanent magnet generators. Most permanent magnets are based on neodymium-iron-boron alloys (NdFeB) doped with praseodymium (Pr), dysprosium (Dy), and terbium (Tb), though the type and quantity of REEs vary depending on turbine design (Gielen, 2021). We use Nd as a representative REE and assume 100 kg Nd/MW are required for offshore turbines only. With this calculation, we find REEs to be the most supply constrained mineral needed for renewables. More than an order of magnitude increase over current annual REE production values is required to meet the high demand scenario. Accounting for REE demand for onshore turbines and for REEs besides Nd would result in an even greater increase over current production levels and further supply constraints. Additionally, since REEs are essential for electric motors (Smith et al., 2022), future demand for these minerals is also likely to increase with greater electric vehicle adoption.

We next compare the total renewable electricity needed to produce green H₂ relative to current and projected global renewable electricity generation. At 45 kWh/kg H₂, a substantial fraction of global wind and solar energy will be required for water electrolysis to produce green H₂ (Figure 4). Under the "existing uses" scenario (100 Mtpa H₂), water electrolysis will require more electricity (~4,500 TWh) than total wind and solar generation in 2022 (3,427 TWh) (IEA, 2021a). The IRENA Planned Energy Scenario (IRENA, 2023b) and IEA Announced Pledges Scenario (IEA, 2021a), two more conservative energy transition models, estimate that wind and solar will generate ~25,000–30,000 TWh globally in 2050. In contrast, more aggressive targets, such as the IRENA 1.5°C Scenario (IRENA, 2023b) and BloombergNEF Net Zero Scenario (Bloomberg, 2022) project wind and solar will generate ~60,000–65,000 TWh in 2050. Considering these generation ranges, the "net zero" H₂ scenario (500 Mtpa, ~22,500 TWh) will require between ~34 and 90% of all installed

renewables in 2050. Even with technological improvements to reach 100% efficiency (39 kWh/kg H₂), electrolyzers may still require at least ~30% of projected global renewables in 2050 under the "net zero" scenario. Given these projections, it is more realistic that green H₂ uses a dedicated "behind-the-meter" system than connecting to the grid, which will also enable lower costs.

The IEA has recently lowered their forecast demand for H₂ in 2050 from 530 (IEA, 2021a) to 470 (IEA, 2023b) Mtpa, perhaps due to some of the aforementioned challenges. Given the quantity of minerals required for renewable generation, we feel the primary focus should be on deployment of renewables for grid electricity generation. At present, the energy and mineral requirements for wind and solar present a major challenge to scaling green H₂ production.

4 Discussion

The above analysis is intended to highlight the challenges associated with scaling green H₂ production using existing technology and mineral production levels, especially with regards to REEs and PGMs. Both REEs and PGMs currently rely on extremely concentrated supply chains with limited geographic diversity. REEs are abundant in the earth's crust (~130 Mt) but are typically found in very low concentrations. Only ~250,000 tonnes of rare earth oxides were mined in 2023 (USGS, 2023), with ~70% mined in China. While there are currently ~70,000 tonnes of known PGM reserves, ~90% are located in South Africa (USGS, 2023). Innovations in surveying technologies to identify new deposits and new technologies to enable mining of REEs and PGMs from low concentration sources would help alleviate potential supply constraints. Within this analysis, we have excluded calculating mineral demands for dedicated battery storage systems, which is expected to increase the demand of other critical minerals, such as Li, Co, and graphite. We also leave the mineral demands of fuel cells out of scope, even though PEM fuel cells may represent additional demand for PGMs (Wang et al., 2022). In addition, we note that locations with high solar CFs are often in water stressed regions, potentially limiting the ease of implementing large-scale electrolysis (Tonelli et al., 2023). Therefore, ongoing research and development efforts dedicated to identifying alternative materials and increasing efficiency are essential. REE-free permanent magnets, significant reductions in catalyst loading, PGM-free AEMs, and coupling water electrolysis with other heat generation sources, are all being investigated and can potentially improve the energy and material efficiency of green H₂ production. Nuclear energy, especially small modular reactors, provides another alternate route to scaling low-emission, electrolysis-based H₂ production, given the high CF and low-emissions profile of nuclear systems (IEA, 2023c). Finally, we note that we have excluded material and energy requirements for "blue hydrogen", which integrates carbon capture and storage with fossil-based methods, due to the nascency of carbon capture and storage technology.

As this special issue highlights, "natural hydrogen" is one potential alternative method for producing low-emission H₂. Natural hydrogen is H₂ created in the subsurface, through serpentinization of Fe-containing rocks or radiolysis of water.

There are potentially large untapped reservoirs of natural hydrogen, ranging from ~0.5 (Lollar et al., 2014) to >1,000 (Zgonnik, 2020) Mtpa of H₂. Natural hydrogen may alleviate some of the mineral supply constraints discussed above and provide a potential source of low-emissions H₂ in the future. However, there are significant challenges to utilizing this unexplored resource and the needed technology is notably less mature than renewables and electrolysis. The exploration and discovery of natural hydrogen accumulations, the ability to stimulate subsurface reservoirs, and the capacity to effectively remove H₂ from the subsurface are currently all not well understood enough for on-demand extraction of H₂. Ensuring that natural H₂ extraction and production does not have unintended environmental impacts, including seismicity, microbial community changes, water usage, and fugitive emissions, is another area in need of research. We hope this opening perspective piece helps frame the exciting and urgent challenges natural hydrogen researchers aim to solve.

Data availability statement

The original contributions presented in the study are included in the article/Supplementary material, further inquiries can be directed to the corresponding author.

Author contributions

JG: Conceptualization, Writing—original draft, Writing—review and editing. MZ: Writing—original draft, Writing—review and editing. DW: Conceptualization, Supervision, Writing—review and editing.

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Conflict of interest

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