



Correction: Levelized cost of CO₂ mitigation from hydrogen production routes

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Correction for 'Levelized cost of CO₂ mitigation from hydrogen production routes' by B. Parkinson *et al.*, *Energy Environ. Sci.*, 2019, 12, 19–40, <https://doi.org/10.1039/C8EE02079E>.

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Some of the references had missing or incorrect details; corrected sections of the affected text are provided below. The reference list has also been corrected and is reproduced in full at the end of this correction.

In Section 3.1, “For literature studies including natural gas supply chain contributions to GHG emissions, the reported total range of LCE values are 10.72–15.86 kg CO₂e kg⁻¹ H₂ (average of 12.4 of kg CO₂e kg⁻¹ H₂)^{26–34} without CCS and 3.1–5.9 kg CO₂e kg⁻¹ H₂ (average of 4.3 kg CO₂e kg⁻¹ H₂) with CCS at 90% capture.^{27,28,32,33,35}” should read as “For literature studies including natural gas supply chain contributions to GHG emissions, the reported total range of LCE values are 10.72–15.86 kg CO₂e kg⁻¹ H₂ (average of 12.4 of kg CO₂e kg⁻¹ H₂)^{30–38} without CCS and 3.1–5.9 kg CO₂e kg⁻¹ H₂ (average of 4.3 kg CO₂e kg⁻¹ H₂) with CCS at 90% capture.^{31,32,36,37,39}”

“Direct GHG emissions from the SMR hydrogen production phase are approximately 8–10 t CO₂e t⁻¹ H₂, 60% of which is generated from the process chemistry, while the remaining 40% arises from heat and power sources required.³⁶” should read as “Direct GHG emissions from the SMR hydrogen production phase are approximately 8–10 t CO₂e t⁻¹ H₂, 60% of which is generated from the process chemistry, while the remaining 40% arises from heat and power sources required.²⁶”

“The majority of CO₂ produced exits in two streams, a diluted stream (stack gases with CO₂ concentration 5–10 vol%) and a concentrated stream (approximately 50% by vol after pressure swing adsorption).³⁷” should read as “The majority of CO₂ produced exits in two streams, a diluted stream (stack gases with CO₂ concentration 5–10 vol%) and a concentrated stream (approximately 50% by vol after pressure swing adsorption).²⁷”

“If deep decarbonisation is required and emissions must be further reduced from the entire process, then an amine solvent (MEA) based CCS process might be used to capture up to 90% of the CO₂ contained in the stack gases,³⁸ although demonstrated removal rates are typically 80%.³⁹” should read as “If deep decarbonisation is required and emissions must be further reduced from the entire process, then an amine solvent (MEA) based CCS process might be used to capture up to 90% of the CO₂ contained in the stack gases,²⁸ although demonstrated removal rates are typically 80%.²⁹”

In Section 3.1.2, “In contrast, the post-demonstration of CCS in the EU reports by the European Technology Platform for Zero Emission Fossil Fuel Power Plants (ZEP) detail the costs of CO₂ transport *via* pipeline (higher for shipping) to range from \$2.4–13 t⁻¹ CO₂,⁹⁹ and storage and monitoring from \$8–23 t⁻¹ CO₂,⁷³ depending on the storage site location and degree of characterization.” should read as “In contrast, the post-demonstration of CCS in the EU reports by the European Technology Platform for Zero Emission Fossil Fuel Power Plants (ZEP) detail the costs of CO₂ transport *via* pipeline (higher for shipping) to range from \$2.4–13 t⁻¹ CO₂,⁷³ and storage and monitoring from \$8–23 t⁻¹ CO₂,⁷³ depending on the storage site location and degree of characterization.”

In Section 3.2, “The lower C : H ratio in coal relative to natural gas results in significantly higher direct CO₂e emissions from the process (14.4–25.31 kg CO₂e kg⁻¹ H₂), with an average value of 19.14 kg CO₂e kg⁻¹ H₂.^{8,22,28,43–45}” should read as “The lower C : H ratio in coal relative to natural gas results in significantly higher direct CO₂e emissions from the process (14.4–25.31 kg CO₂e kg⁻¹ H₂), with an average value of 19.14 kg CO₂e kg⁻¹ H₂.^{8,22,32,43–45}”

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In Section 3.2.1, “This represents a relatively small portion of the average $0.45 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$ (range $0.32\text{--}0.77 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$) for coal extraction, processing and transportation reported in the literature for subbituminous coal supply chains.^{27,30,45} The range of LCE from coal gasification with CCS ($\geq 90\%$ capture) presented in the literature range from $0.77\text{--}5.2 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$ ^{27,30,44–46} with an average of $4.56 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$,” should read as “This represents a relatively small portion of the average $0.45 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$ (range $0.32\text{--}0.77 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$) for coal extraction, processing and transportation reported in the literature for subbituminous coal supply chains.^{31,34,45} The range of LCE from coal gasification with CCS ($\geq 90\%$ capture) presented in the literature range from $0.77\text{--}5.2 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$ ^{31,34,44,45,48} with an average of $4.56 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$.”

In Section 3.2.2, “Whilst several studies for coal gasification integrated with CCS for power generation are available,^{47,48}” should read as “Whilst several studies for coal gasification integrated with CCS for power generation are available,^{46,47}”.

“CAC reference plant for IGCC facilities for power production is a supercritical pulverized coal (SCPC) plant without capture (not a similar IGCC plant without capture), which would be the lower cost route for coal-fired power plants without capture.⁴⁵ This results in an increase in the average CAC from $\$43.92 \text{ t}^{-1} \text{ CO}_2$ (for IGCC with CCS to IGCC without) to $\$77.34 \text{ t}^{-1} \text{ CO}_2$ (excluding transport and storage for both) as reported in a recent review by Rubin *et al.*⁴⁵ An average value of $\$43.92 \text{ t}^{-1} \text{ CO}_2$ for IGCC compared to a SMR baseline has been used.⁴⁵” should read as “CAC reference plant for IGCC facilities for power production is a supercritical pulverized coal (SCPC) plant without capture (not a similar IGCC plant without capture), which would be the lower cost route for coal-fired power plants without capture.⁴⁶ This results in an increase in the average CAC from $\$43.92 \text{ t}^{-1} \text{ CO}_2$ (for IGCC with CCS to IGCC without) to $\$77.34 \text{ t}^{-1} \text{ CO}_2$ (excluding transport and storage for both) as reported in a recent review by Rubin *et al.*⁴⁶ An average value of $\$43.92 \text{ t}^{-1} \text{ CO}_2$ for IGCC compared to a SMR baseline has been used.⁴⁶”

In Section 3.4, “For a conservative approach in this study, a LCOH of $\$1.76 \text{ kg}^{-1} \text{ H}_2$ is estimated based on a recent model developed by Parkinson *et al.*,⁷¹” should read as “For a conservative approach in this study, a LCOH of $\$1.76 \text{ kg}^{-1} \text{ H}_2$ is estimated based on a recent model developed by Parkinson *et al.*,⁷⁰”.

“Several studies^{35,68–70} report the direct emissions of a pyrolysis facility to range from $0.2\text{--}2.5 \text{ kg CO}_2 \text{ kg}^{-1} \text{ H}_2$,” should read as “Several studies^{39,68–70} report the direct emissions of a pyrolysis facility to range from $0.2\text{--}2.5 \text{ kg CO}_2 \text{ kg}^{-1} \text{ H}_2$.”

“For methane pyrolysis using the updated supply chain emissions discussed in Section 3.1, this study estimates the supply chain emissions contribute $4.28 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$ for the median case, with a range of $3.26\text{--}6.44 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$ using an overall process efficiency of $53\% \text{ HHV}$.⁷¹” should read as “For methane pyrolysis using the updated supply chain emissions discussed in Section 3.1, this study estimates the supply chain emissions contribute $4.28 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$ for the median case, with a range of $3.26\text{--}6.44 \text{ kg CO}_2\text{e kg}^{-1} \text{ H}_2$ using an overall process efficiency of $53\% \text{ HHV}$.⁷⁰”

In Section 3.5, “SOEC are the most electrically efficient of the three technologies, but are currently in the R&D stage facing challenges with corrosion, seals, thermal cycling and chromium migration.⁷³ PEM electrolysis is currently more expensive than alkaline technologies, but is an attractive future technology due to higher current densities ($> 2 \text{ A cm}^{-2}$), higher efficiencies (the largest cost driver), dynamic operation and compact system design.⁷⁴ An excellent summary of the technical specifications of each technology has been compiled by Bhandari *et al.*²⁸” should read as “SOEC are the most electrically efficient of the three technologies, but are currently in the R&D stage facing challenges with corrosion, seals, thermal cycling and chromium migration.⁵² PEM electrolysis is currently more expensive than alkaline technologies, but is an attractive future technology due to higher current densities ($> 2 \text{ A cm}^{-2}$), higher efficiencies (the largest cost driver), dynamic operation and compact system design.⁷⁴ An excellent summary of the technical specifications of each technology has been compiled by Bhandari *et al.*³²”

In Section 3.5.1, “The direct emissions from electrolytic hydrogen production are very low, but the indirect emissions associated with electricity feedstock and electrolyzer systems must be considered.²⁸” should read as “The direct emissions from electrolytic hydrogen production are very low, but the indirect emissions associated with electricity feedstock and electrolyzer systems must be considered.³²”

“Little data is available on the LCE contribution from electrolyser manufacturing and replacements, with estimates suggesting a relatively small contribution in the range of $30\text{--}50 \text{ g CO}_2\text{e kg}^{-1} \text{ H}_2$ ^{28,76,77},” should read as “Little data is available on the LCE contribution from electrolyser manufacturing and replacements, with estimates suggesting a relatively small contribution in the range of $30\text{--}50 \text{ g CO}_2\text{e kg}^{-1} \text{ H}_2$ ^{32,76,77}.”

In Section 3.5.2, “PEM was chosen as the reference electrolyser technology operating with an 85% cell voltage efficiency ($\sim 51 \text{ kW h kg}^{-1} \text{ H}_2$).¹⁰⁶” should read as “PEM was chosen as the reference electrolyser technology operating with an 85% cell voltage efficiency ($\sim 51 \text{ kW h kg}^{-1} \text{ H}_2$).⁷⁵”

In Section 4.3, “These decarbonization fractions are not commensurate with the decarbonization targets required in transport, heat and industry under many scenarios, particularly as global aspirations turn to net zero emissions in the second half of the 21st century.⁸⁴ This highlights that technologies such as carbon capture and sequestration may potentially be an expensive exercise in heroic futility if all aspects of hydrogen supply chains are not addressed. No cost is currently applied to methane emissions and the costs of mitigating supply chain emissions are less well understood than emissions at the point of conversion. It has been suggested that 75% of global methane emissions from oil and gas supply chains are mitigatable, with half of these achieved at a positive net present value.⁸⁵” should read as “These decarbonization fractions are not commensurate with the decarbonization targets required in transport, heat and industry under many scenarios, particularly as global aspirations turn to net zero emissions



in the second half of the 21st century.¹²⁶ This highlights that technologies such as carbon capture and sequestration may potentially be an expensive exercise in heroic futility if all aspects of hydrogen supply chains are not addressed. No cost is currently applied to methane emissions and the costs of mitigating supply chain emissions are less well understood than emissions at the point of conversion. It has been suggested that 75% of global methane emissions from oil and gas supply chains are mitigatable, with half of these achieved at a positive net present value.^{86,}

“With enough incentive (*e.g. via* pollution tax as per ref. 86)” should read as “With enough incentive (*e.g. via* pollution tax as per ref. 127)”.

In Section 4.3.1, “The deployment of CCS at-scale is the only method widely considered to be available to substantially mitigate CO₂ emissions from fossil fuels, which so far has failed to meet ambitious expectations, and continued delays of demonstration projects are causing considerable uncertainty about the role CCS will play in carbon mitigation.⁸⁷ This study has shown that equipping CCS to SMR (90% capture) only effectively reduces the life cycle GHG footprint of hydrogen production by 38–76%, depending on the contribution of supply chain emissions. Additionally, the capture rates may currently be lower than 90%.^{39,}” should read as “The deployment of CCS at-scale is the only method widely considered to be available to substantially mitigate CO₂ emissions from fossil fuels, which so far has failed to meet ambitious expectations, and continued delays of demonstration projects are causing considerable uncertainty about the role CCS will play in carbon mitigation.¹¹² This study has shown that equipping CCS to SMR (90% capture) only effectively reduces the life cycle GHG footprint of hydrogen production by 38–76%, depending on the contribution of supply chain emissions. Additionally, the capture rates may currently be lower than 90%.^{29,}”

“The operational and capital costs would also change with different capture rates, with higher capture rates requiring greater fuel duty and equipment.^{88,}” should read as “The operational and capital costs would also change with different capture rates, with higher capture rates requiring greater fuel duty and equipment.^{113,}”

“A detailed review of carbon capture and utilization options is presented by Hunt *et al.*^{89,}” should read as “A detailed review of carbon capture and utilization options is presented by Hunt *et al.*^{114,}”

“Amine-based CO₂ absorption systems are also commercially mature technologies in the chemicals sector.^{90,}” should read as “Amine-based CO₂ absorption systems are also commercially mature technologies in the chemicals sector.^{115,}”

“Several proposals for equipping CCS to industrially active clusters with shared transport and storage infrastructure have been proposed and it is widely agreed the largest cost reductions can be achieved here.^{91–95,}” should read as “Several proposals for equipping CCS to industrially active clusters with shared transport and storage infrastructure have been proposed and it is widely agreed the largest cost reductions can be achieved here.^{116–120,}”

“Long term monitoring of storage sites is also required to ensure leakage rates to the atmosphere of less 0.1% year^{−1} needed to ensure effective climate change abatement are maintained.^{96,97} Commercial scale storage sites will also have to manage the risks of financial penalties for unforeseen leakages.^{98,}” should read as “Long term monitoring of storage sites is also required to ensure leakage rates to the atmosphere of less 0.1% year^{−1} needed to ensure effective climate change abatement are maintained.^{121,122} Commercial scale storage sites will also have to manage the risks of financial penalties for unforeseen leakages.^{123,}”

In Section 4.3.2, “It should be noted however, there is a large body of literature that suggests up to 60% of coal mine methane mitigation is achievable at relatively low cost.^{99,100,}” should read as “It should be noted however, there is a large body of literature that suggests up to 60% of coal mine methane mitigation is achievable at relatively low cost.^{128,129,}”

“Coal-to-hydrogen processes are however less prevalent as dedicated hydrogen producers, with advanced coal gasification concepts aiming to integrate CO₂ separation with water–gas shift reactions to achieve higher process efficiencies with ease of CO₂ capture.^{101,}” should read as “Coal-to-hydrogen processes are however less prevalent as dedicated hydrogen producers, with advanced coal gasification concepts aiming to integrate CO₂ separation with water–gas shift reactions to achieve higher process efficiencies with ease of CO₂ capture.^{130,}”

“The order of magnitude estimates for coal-to-hydrogen with CCS are likely to remain around the \$60–100 t^{−1} CO₂.^{47,}” should read as “The order of magnitude estimates for coal-to-hydrogen with CCS are likely to remain around the \$60–100 t^{−1} CO₂.^{46,}”

In Section 4.3.3, “This finding is of particular importance as the technology is regularly cited in academic literature as a means of producing CO₂-free hydrogen,^{2,4,6,9,14,19,21,66,68–70,102–106} which is only achievable if supply chain emissions are low. It should be noted, however, in contrast to technologies such as CCS the storage of a solid by-product carbon is relatively simple either temporarily or permanently, an idea first proposed as the ‘carbon moratorium’ by Kreysa.^{104,}” should read as “This finding is of particular importance as the technology is regularly cited in academic literature as a means of producing CO₂-free hydrogen,^{2,4,6,9,14,19,21,66,68–70,103,104,124,125,131,132} which is only achievable if supply chain emissions are low. It should be noted, however, in contrast to technologies such as CCS the storage of a solid by-product carbon is relatively simple either temporarily or permanently, an idea first proposed as the ‘carbon moratorium’ by Kreysa.^{125,}”

“There are several technical challenges associated with management of the solid carbon produced which fouls (coke) solid catalysts.^{102,107–109} However, proponents of the technology claim these limitations can be overcome with appropriate process and reactor design to allow easy separation of the carbon product.^{10,52,106,110,}” should read as “There are several technical challenges associated with management of the solid carbon produced which fouls (coke) solid catalysts.^{131,133–135} However, proponents of the technology claim these limitations can be overcome with appropriate process and reactor design to allow easy separation of the carbon product.^{10,52,132,136,}”



In Section 4.4, “It is interesting to note that electrolysis avoidance costs in their current status are comparable to negative emissions technologies such as direct capture, estimated to be \$600–700 t⁻¹ CO₂ avoided.^{111,112}” should read as “It is interesting to note that electrolysis avoidance costs in their current status are comparable to negative emissions technologies such as direct capture, estimated to be \$600–700 t⁻¹ CO₂ avoided.^{137,138}”

“For example, wind capacity factors have increased markedly in the past decade and in some sites (*e.g.* Denmark) may exceed 50%.¹¹³” should read as “For example, wind capacity factors have increased markedly in the past decade and in some sites (*e.g.* Denmark) may exceed 50%.¹³⁹”

“This is illustrated in Fig. 10(b), which shows the effect of increasing the utilization of an optimistic 50% capacity factor future wind electrolysis scenario using the average EU (~0.275 kg CO₂ kW h⁻¹, \$0.081 kW h⁻¹^{114,115}) and French (~0.050 kg CO₂ kW h⁻¹, \$0.0596 kW h⁻¹^{114,115}) electricity grids.” should read as “This is illustrated in Fig. 10(b), which shows the effect of increasing the utilization of an optimistic 50% capacity factor future wind electrolysis scenario using the average EU (~0.275 kg CO₂ kW h⁻¹, \$0.081 kW h⁻¹^{140,141}) and French (~0.050 kg CO₂ kW h⁻¹, \$0.0596 kW h⁻¹^{140,141}) electricity grids.”

“Wind and solar power have witnessed substantial cost reductions over the past two decades and potential areas of future cost reduction have been identified by others.¹¹⁶ The ability of solar PV to meet the required cost reductions to be a more cost competitive mitigation option requires a balance between the investments required to produce and install a module, the total energy provided by that module and its lifetime and conversion efficiency.¹¹⁷ New materials realising sufficient efficiency, stability and cost will be required.¹¹⁸” should read as “Wind and solar power have witnessed substantial cost reductions over the past two decades and potential areas of future cost reduction have been identified by others.¹⁴² The ability of solar PV to meet the required cost reductions to be a more cost competitive mitigation option requires a balance between the investments required to produce and install a module, the total energy provided by that module and its lifetime and conversion efficiency.¹⁴³ New materials realising sufficient efficiency, stability and cost will be required.¹¹¹”

“Continuous increases in the average capacity of turbines, hub-heights and swept areas have allowed higher utilization factors and reduced costs.¹¹⁹” should read as “Continuous increases in the average capacity of turbines, hub-heights and swept areas have allowed higher utilization factors and reduced costs.¹⁴⁴”

“Thermal efficiencies of nuclear and combustion power plants to electricity operating through Rankine cycles range from 31% for a Magnox type to around 40% for an advanced gas cooled reactor.¹²⁰” should read as “Thermal efficiencies of nuclear and combustion power plants to electricity operating through Rankine cycles range from 31% for a Magnox type to around 40% for an advanced gas cooled reactor.¹⁴⁵”

In Section 4.5, “Although the thermochemical cycle efficiencies (~50%) have yet to be demonstrated at scale, it has been the subject of significant research by the Japan Atomic Energy Agency,¹²¹ General Atomics¹²² and Westinghouse.¹²³” should read as “Although the thermochemical cycle efficiencies (~50%) have yet to be demonstrated at scale, it has been the subject of significant research by the Japan Atomic Energy Agency,¹⁴⁶ General Atomics¹⁴⁷ and Westinghouse.¹⁴⁸”

“Although unit costs for technologies usually decrease with increasing volume of production, nuclear power has consistently seen the opposite within the United States; which reflects the idiosyncrasies of the regulatory environment as public opposition grew and regulations were tightened.¹²⁴ Lovering *et al.*¹²⁵” should read as “Although unit costs for technologies usually decrease with increasing volume of production, nuclear power has consistently seen the opposite within the United States; which reflects the idiosyncrasies of the regulatory environment as public opposition grew and regulations were tightened.¹⁴⁹ Lovering *et al.*¹⁵⁰”

“They concluded that there is no inherent cost escalation trend associated with nuclear technology, and the large variance witnessed in cost trends over time and across different countries, even with similar nuclear reactor technologies, suggests that cost drivers other than learning-by-doing have dominated the cost experience of nuclear power construction.¹²⁵ Additionally, as the majority of literature on nuclear power costs have focussed almost exclusively on the United States and France, there is an incomplete picture of the economic evolution of the technology.¹²⁵” should read as “They concluded that there is no inherent cost escalation trend associated with nuclear technology, and the large variance witnessed in cost trends over time and across different countries, even with similar nuclear reactor technologies, suggests that cost drivers other than learning-by-doing have dominated the cost experience of nuclear power construction.¹⁵⁰ Additionally, as the majority of literature on nuclear power costs have focussed almost exclusively on the United States and France, there is an incomplete picture of the economic evolution of the technology.¹⁵⁰”

In Section 4.6, “Only a very small percentage of solar energy is converted to hydrogen (~6–12 wt% H₂ kg⁻¹ biomass¹²⁶)” should read as “Only a very small percentage of solar energy is converted to hydrogen (~6–12 wt% H₂ kg⁻¹ biomass¹⁵¹)”.

“A recent report¹²⁷ suggests with anticipated improvements in agricultural practices and plant breeding, feedstocks may exceed 244 million dry tons at a farm-gate price of \$60 dry ton⁻¹. It has also been proposed that waste biomass feedstocks could be co-fired in coal gasification facilities to further reduce the net CO₂ closer to zero with minimal impact on the downstream flue gas treatment unit.¹²⁸ In a plant with post-combustion capture this increases the cost of electricity by 6% and has no impact on the cost of CO₂ avoidance, but the cost depends strongly on the cost of biomass.¹²⁸” should read as “A recent report⁸¹ suggests with anticipated improvements in agricultural practices and plant breeding, feedstocks may exceed 244 million dry tons at a farm-gate price of \$60 dry ton⁻¹. It has also been proposed that waste biomass feedstocks could be co-fired in coal gasification facilities to



further reduce the net CO₂ closer to zero with minimal impact on the downstream flue gas treatment unit.¹⁰⁰ In a plant with post-combustion capture this increases the cost of electricity by 6% and has no impact on the cost of CO₂ avoidance, but the cost depends strongly on the cost of biomass.¹⁰⁰

“The production of hydrogen from biomass with CCS is one of only a few technologies that may deliver negative emissions at relatively modest costs, which may become important for global decarbonization issues in the second half of this century.¹²⁹” should read as “The production of hydrogen from biomass with CCS is one of only a few technologies that may deliver negative emissions at relatively modest costs, which may become important for global decarbonization issues in the second half of this century.⁹⁹”

“This is reflected by several commercial examples of biomass for heat and power¹³⁰ but no completed industrial-scale demonstrations of any biomass technology for hydrogen production.¹²⁶” should read as “This is reflected by several commercial examples of biomass for heat and power¹⁰⁶ but no completed industrial-scale demonstrations of any biomass technology for hydrogen production.¹⁵¹”

In Section 5, “These decarbonization fractions are not commensurate with the decarbonization targets required in transport, heat and industry under many scenarios, particularly as global aspirations turn to net zero emissions in the second half of the 21st century in line with the Paris Agreement.⁸⁴” should read as “These decarbonization fractions are not commensurate with the decarbonization targets required in transport, heat and industry under many scenarios, particularly as global aspirations turn to net zero emissions in the second half of the 21st century in line with the Paris Agreement.¹²⁶”

“This finding is of particular importance as the technology is regularly cited in academic literature as a means of producing CO₂-free hydrogen,^{2,4,6,9,14,19,21,66,68–70,102–106}” should read as “This finding is of particular importance as the technology is regularly cited in academic literature as a means of producing CO₂-free hydrogen,^{2,4,6,9,14,19,21,66,68–70,103,104,125,131,132}”.

The Royal Society of Chemistry apologises for these errors and any consequent inconvenience to authors and readers.

References

- 1 M. Hulme, *Nat. Clim. Change*, 2016, **6**, 222–224.
- 2 N. Z. Muradov and T. N. Veziroğlu, *Int. J. Hydrogen Energy*, 2008, **33**, 6804–6839.
- 3 G. Simbollotti, *IEA Energy Technology Essentials – Hydrogen Production and Distribution*, 2017.
- 4 A. Konieczny, K. Mondal, T. Wiltowski and P. Dydo, *Int. J. Hydrogen Energy*, 2008, **33**, 264–272.
- 5 A. H. Fakeeha, A. A. Ibrahim, W. U. Khan, K. Seshan, R. L. Al Otaibi and A. S. Al-Fatesh, *Arabian J. Chem.*, 2018, **11**, 405–414.
- 6 U. P. M. Ashik, W. M. A. Wan Daud and H. F. Abbas, *Renewable Sustainable Energy Rev.*, 2015, **44**, 221–256.
- 7 M. Granovskii, I. Dincer and M. A. Rosen, *J. Power Sources*, 2006, **157**, 411–421.
- 8 N. Muradov, *Int. J. Hydrogen Energy*, 2017, **42**, 14058–14088.
- 9 N. Muradov and T. Veziroğlu, *Int. J. Hydrogen Energy*, 2005, **30**, 225–237.
- 10 L. Weger, A. Abánades and T. Butler, *Int. J. Hydrogen Energy*, 2017, **42**, 720–731.
- 11 B. C. R. Ewan and R. W. K. Allen, *Int. J. Hydrogen Energy*, 2005, **30**, 809–819.
- 12 C. Acar and I. Dincer, *Int. J. Hydrogen Energy*, 2014, **39**, 1–12.
- 13 I. Dincer and C. Acar, *Int. J. Hydrogen Energy*, 2015, **40**, 11094–11111.
- 14 O. Machhammer, A. Bode and W. Hormuth, *Chem. Eng. Technol.*, 2016, **39**, 1185–1193.
- 15 J. Speirs, P. Balcombe, E. Johnson, J. Martin, N. Brandon and A. Hawkes, *Energy Policy*, 2018, **118**, 291–297.
- 16 A. Haryanto, S. Fernando, N. Murali and S. Adhikari, *Energy Fuels*, 2005, **19**, 2098–2106.
- 17 J. W. Ager, M. R. Shaner, K. A. Walczak, I. D. Sharp and S. Ardo, *Energy Environ. Sci.*, 2015, **8**, 2811–2824.
- 18 J. R. McKone, N. S. Lewis and H. B. Gray, *Chem. Mater.*, 2014, **26**, 407–414.
- 19 N. Z. Muradov and T. N. Veziroğlu, *Carbon-neutral fuels and energy carriers*, CRC Press, 2011.
- 20 The Royal Society, *Options for producing low-carbon hydrogen at scale*, Report DES4801_2, The Royal Society, 2018.
- 21 B. Parkinson, M. Tabatabaei, D. C. Upham, B. Ballinger, C. Greig, S. Smart and E. McFarland, *Int. J. Hydrogen Energy*, 2018, **43**, 2540–2555.
- 22 National Research Council, *The hydrogen economy: opportunities, costs, barriers, and R&D needs*, Report 0309091632, National Academies Press, 2004.
- 23 D. Gray and G. Tomlinson, *MTR 2002-31 Mitretek Technical Paper Hydrogen From Coal*, 2001.
- 24 F. Mueller-Langer, E. Tzimas, M. Kaltschmitt and S. Peteves, *Int. J. Hydrogen Energy*, 2007, **32**, 3797–3810.
- 25 S. Penner, *Energy*, 2006, **31**, 33–43.
- 26 M. Melania, *Current central hydrogen production from natural gas without CO₂ sequestration*, version 3.101.
- 27 G. Collodi and F. Wheeler, *Chem. Eng. Trans.*, 2010, **19**, 37–42.
- 28 G. Collodi, G. Azzaro, N. Ferrari and S. Santos, *Energy Procedia*, 2017, **114**, 2690–2712.



- 29 P. Balcombe, J. Speirs, E. Johnson, J. Martin, N. Brandon and A. Hawkes, *Renewable Sustainable Energy Rev.*, 2018, **91**, 1077–1088.
- 30 A. H. Strømman and E. Hertwich, *Hybrid life cycle assessment of large-scale hydrogen production facilities*, 2004.
- 31 J. Ruether, M. Ramezan and E. Grol, *Life-cycle analysis of greenhouse gas emissions for hydrogen fuel production in the United States from LNG and coal*, DOE/NETL-2006/1227, November, 2005.
- 32 R. Bhandari, C. A. Trudewind and P. Zapp, *J. Cleaner Prod.*, 2014, **85**, 151–163.
- 33 J. Lane and P. Spath, *Technoeconomic analysis of the thermocatalytic decomposition of natural gas*, National Renewable Energy Lab., Golden, CO (US), 2001.
- 34 E. Cetinkaya, I. Dincer and G. F. Naterer, *Int. J. Hydrogen Energy*, 2012, **37**, 2071–2080.
- 35 C. Koroneos, A. Dompros, G. Roumbas and N. Moussiopoulos, *Int. J. Hydrogen Energy*, 2004, **29**, 1443–1450.
- 36 D. Sadler, *H21 Leeds City Gate Report*, Northern Gas Networks, 2016.
- 37 J. Dufour, D. P. Serrano, J. L. Gálvez, A. González, E. Soria and J. L. Fierro, *Int. J. Hydrogen Energy*, 2012, **37**, 1173–1183.
- 38 L. Tock and F. Maréchal, *Int. J. Hydrogen Energy*, 2012, **37**, 11785–11795.
- 39 J. Dufour, D. P. Serrano, J. L. Gálvez, J. Moreno and C. García, *Int. J. Hydrogen Energy*, 2009, **34**, 1370–1376.
- 40 R. Kothari, D. Buddhi and R. L. Sawhney, *Renewable Sustainable Energy Rev.*, 2008, **12**, 553–563.
- 41 N. V. S. N. M. Konda, N. Shah and N. P. Brandon, *Int. J. Hydrogen Energy*, 2011, **36**, 4619–4635.
- 42 C. E. G. Padro and V. Putsche, *Survey of the economics of hydrogen technologies*, National Renewable Energy Lab., Golden, CO (US), 1999.
- 43 Royal Belgian Academy Council of Applied Science, *Hydrogen as an energy carrier*, 2006.
- 44 P. Chiesa, S. Consonni, T. Kreutz and R. Williams, *Int. J. Hydrogen Energy*, 2005, **30**, 747–767.
- 45 P. Burmistrz, T. Chmielniak, L. Czepirski and M. Gazda-Grzywacz, *J. Cleaner Prod.*, 2016, **139**, 858–865.
- 46 E. S. Rubin, J. E. Davison and H. J. Herzog, *Int. J. Greenhouse Gas Control*, 2015, **40**, 378–400.
- 47 IEAGHG, *Co-production of hydrogen and electricity by coal gasification with CO₂ capture – updated economic analysis*, 2008.
- 48 A. Verma and A. Kumar, *Appl. Energy*, 2015, **147**, 556–568.
- 49 P. Nikolaidis and A. Poullikkas, *Renewable Sustainable Energy Rev.*, 2017, **67**, 597–611.
- 50 GoGreenGas, *BioSNG Demonstration Plant Project Close-Down Report*, 2017.
- 51 T. L. Group, Green Hydrogen from Biomass, https://www.linde-engineering.com/en/innovations/green_hydrogen_from_biomass/index.html, accessed April 2018.
- 52 J. D. Holladay, J. Hu, D. L. King and Y. Wang, *Catal. Today*, 2009, **139**, 244–260.
- 53 D. Wang, S. Czernik, D. Montane, M. Mann and E. Chornet, *Ind. Eng. Chem. Res.*, 1997, **36**, 1507–1518.
- 54 J. R. Bartels, M. B. Pate and N. K. Olson, *Int. J. Hydrogen Energy*, 2010, **35**, 8371–8384.
- 55 J. Yao, M. Kraussler, F. Benedikt and H. Hofbauer, *Energy Convers. Manage.*, 2017, **145**, 278–292.
- 56 A. Mehmeti, A. Angelis-Dimakis, G. Arampatzis, S. J. McPhail and S. Ulgiati, *Environments*, 2018, **5**, 24.
- 57 A. Valente, D. Iribarren and J. Dufour, *Int. J. Life Cycle Assess.*, 2017, **22**, 346–363.
- 58 M. Martín-Gamboa, D. Iribarren, A. Susmozas and J. Dufour, *Bioresour. Technol.*, 2016, **214**, 376–385.
- 59 A. Susmozas, D. Iribarren and J. Dufour, *Int. J. Hydrogen Energy*, 2013, **38**, 9961–9972.
- 60 Y. Kalinci, A. Hepbasli and I. Dincer, *Int. J. Hydrogen Energy*, 2012, **37**, 14026–14039.
- 61 J. Moreno and J. Dufour, *Int. J. Hydrogen Energy*, 2013, **38**, 7616–7622.
- 62 D. Iribarren, A. Susmozas, F. Petrakopoulou and J. Dufour, *J. Cleaner Prod.*, 2014, **69**, 165–175.
- 63 A. Susmozas, D. Iribarren, P. Zapp, J. Linßen and J. Dufour, *Int. J. Hydrogen Energy*, 2016, **41**, 19484–19491.
- 64 D. Mahajan, C. E. Taylor and G. A. Mansoori, *J. Pet. Sci. Eng.*, 2007, **56**, 1–8.
- 65 A. M. Amin, E. Croiset and W. Epling, *Int. J. Hydrogen Energy*, 2011, **36**, 2904–2935.
- 66 R. A. Dagle, V. Dagle, M. D. Bearden, J. D. Holladay, T. R. Krause and S. Ahmed, *An Overview of Natural Gas Conversion Technologies for Co-Production of Hydrogen and Value-Added Solid Carbon Products*, Pacific Northwest National Lab. (PNNL), Richland, WA (United States); Argonne National Lab. (ANL), Argonne, IL (United States), 2017.
- 67 N. Shah, D. Panjala and G. P. Huffman, *Energy Fuels*, 2001, **15**, 1528–1534.
- 68 M. Steinberg, *Int. J. Hydrogen Energy*, 1999, **24**, 771–777.
- 69 N. Z. Muradov, *Energy Fuels*, 1998, **12**, 41–48.
- 70 B. Parkinson, J. W. Matthews, T. B. McConaughy, D. C. Upham and E. W. McFarland, *Chem. Eng. Technol.*, 2017, **40**, 1022–1030.
- 71 S. Postels, A. Abánades, N. von der Assen, R. K. Rathnam, S. Stückrad and A. Bardow, *Int. J. Hydrogen Energy*, 2016, **41**, 23204–23212.
- 72 S. M. Saba, M. Müller, M. Robinius and D. Stolten, *Int. J. Hydrogen Energy*, 2017, **43**, 1209–1223.
- 73 Zero Emissions Platform, *The Costs of CO₂-Storage—Post-Demonstration CCS in the EU*, 2011.
- 74 K. E. Ayers, C. Capuano and E. B. Anderson, *ECS Trans.*, 2012, **41**, 15–22.
- 75 G. R. Saur, B. James and W. Colella, *Hydrogen and fuel cells program: production case studies*.



- 76 J. C. Koj, A. Schreiber, P. Zapp and P. Marcuello, *Energy Procedia*, 2015, **75**, 2871–2877.
- 77 M. Fischer, M. Faltenbacher and O. Schuller, *Life Cycle Impact Assessment*, ECTOS, 2005.
- 78 L. C. Brown, G. E. Besenbruch, R. Lentsch, K. R. Schultz, J. Funk, P. Pickard, A. Marshall and S. Showalter, *High efficiency generation of hydrogen fuels using nuclear power*, General Atomics, San Diego, CA (United States), 2003.
- 79 L. Brown, J. Funk and S. Showalter, *Initial screening of thermochemical water-splitting cycles for high efficiency generation of hydrogen fuels using nuclear power*, General Atomics, San Diego, CA (US); University of Kentucky, Lexington, KY (US); Sandia National Labs, Albuquerque, NM (US), 1999.
- 80 M. T. Balta, I. Dincer and A. Hepbasli, *Int. J. Hydrogen Energy*, 2009, **34**, 2925–2939.
- 81 U.S. DOE, *US Billion-Ton Update: Biomass Supply for a Bioenergy and Bioproducts Industry*, 2011.
- 82 Z. L. Wang, G. F. Naterer, K. S. Gabriel, R. Gravelins and V. N. Daggupati, *Int. J. Hydrogen Energy*, 2010, **35**, 4820–4830.
- 83 A. Ozbilen, I. Dincer and M. A. Rosen, *Int. J. Hydrogen Energy*, 2011, **36**, 11321–11327.
- 84 P. Balcombe, K. Anderson, J. Speirs, N. Brandon and A. Hawkes, *ACS Sustainable Chem. Eng.*, 2016, **5**, 3–20.
- 85 M. Rutkowski, Current (2010) hydrogen from natural gas with CO₂ capture and sequestration, http://www.hydrogen.energy.gov/h2a_prod_studies.html, accessed March 2018.
- 86 IEA, *World Energy Outlook 2017*, 2017.
- 87 P. Balcombe, N. P. Brandon and A. D. Hawkes, *J. Cleaner Prod.*, 2018, **172**, 2019–2032.
- 88 G. Myhre, D. Shindell, F.-M. Bréon, W. Collins, J. Fuglestedt, J. Huang, D. Koch, J.-F. Lamarque, D. Lee, B. Mendoza, T. Nakajima, A. Robock, G. Stephens, T. Takemura and H. Zhang, in *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*, ed. T. F. Stocker, D. Qin, G.-K. Plattner, M. Tignor, S.K. Allen, J. Boschung, A. Nauels, Y. Xia, V. Bex and P.M. Midgley, Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 2013.
- 89 M. Saunio, P. Bousquet, B. Poulter, A. Peregon, P. Ciais, J. G. Canadell, E. J. Dlugokencky, G. Etiope, D. Bastviken, S. Houweling, G. Janssens-Maenhout, F. N. Tubiello, S. Castaldi, R. B. Jackson, M. Alexe, V. K. Arora, D. J. Beerling, P. Bergamaschi, D. R. Blake, G. Brailsford, V. Brovkin, L. Bruhwiler, C. Crevoisier, P. Crill, K. Covey, C. Curry, C. Frankenberg, N. Gedney, L. Höglund-Isaksson, M. Ishizawa, A. Ito, F. Joos, H.-S. Kim, T. Kleinen, P. Krummel, J.-F. Lamarque, R. Langenfelds, R. Locatelli, T. Machida, S. Maksyutov, K. C. McDonald, J. Marshall, J. R. Melton, I. Morino, V. Naik, S. O'Doherty, F.-J. W. Parmentier, P. K. Patra, C. Peng, S. Peng, G. P. Peters, I. Pison, C. Prigent, R. Prinn, M. Ramonet, W. J. Riley, M. Saito, M. Santini, R. Schroeder, I. J. Simpson, R. Spahni, P. Steele, A. Takizawa, B. F. Thornton, H. Tian, Y. Tohjima, N. Viovy, A. Voulgarakis, M. van Weele, G. R. van der Werf, R. Weiss, C. Wiedinmyer, D. J. Wilton, A. Wiltshire, D. Worthy, D. Wunch, X. Xu, Y. Yoshida, B. Zhang, Z. Zhang and Q. Zhu, *Earth Syst. Sci. Data*, 2016, **8**, 697–751.
- 90 D. A. Kirchgessner, S. D. Piccot and S. S. Masemore, *J. Air Waste Manag. Assoc.*, 2000, **50**, 1904–1919.
- 91 D. A. Kirchgessner, S. D. Piccot and J. D. Winkler, *Chemosphere*, 1993, **26**, 453–472.
- 92 W. Irving, O. Tailakov and D. Kruger, *Good Practice Guidance and Uncertainty Management in National Greenhouse Gas Inventories*, 1999.
- 93 M. Mann and P. Spath, *Life Cycle Assessment of Renewable Hydrogen Production via Wind/Electrolysis: Milestone Completion Report*, National Renewable Energy Lab., Golden, CO (US), 2004.
- 94 J. E. Mason, *Energy Policy*, 2007, **35**, 1315–1329.
- 95 A. Patyk, T. M. Bachmann and A. Brisse, *Int. J. Hydrogen Energy*, 2013, **38**, 3865–3880.
- 96 D. Nugent and B. K. Sovacool, *Energy Policy*, 2014, **65**, 229–244.
- 97 E. S. Warner and G. A. Heath, *J. Ind. Ecol.*, 2012, **16**, S73–S92.
- 98 S. Santos, Understanding the Potential of CCS in Hydrogen Production (Review of the Current State-of-the-Art), http://www.ieaghg.org/docs/General_Docs/IEAGHG_Presentations/18_-_S._Santos_IEAGHGSECURED.pdf.
- 99 S. Fuss, J. G. Canadell, G. P. Peters, M. Tavoni, R. M. Andrew, P. Ciais, R. B. Jackson, C. D. Jones, F. Kraxner, N. Nakicenovic, C. Le Quere, M. R. Raupach, A. Sharifi, P. Smith and Y. Yamagata, *Nat. Clim. Change*, 2014, **4**, 850–853.
- 100 IEAGHG, *CO₂ capture at coal based power and hydrogen plants, Report 2014/3*, Cheltenham, UK, 2014.
- 101 EPRI, *Program on technology innovation: integrated generation technology options 2012. Technical Update*, Electric Power Research Institute, Palo Alto, CA, 2013.
- 102 J. Black, *Cost and Performance Baseline for Fossil Energy Plants Volume 1: Bituminous Coal and Natural Gas to Electricity – Revision 2a, September 2013*, DOE/2010/1397, National Energy Technology Laboratory, USA, 2013.
- 103 T. Keipi, H. Tolvanen and J. Konttinen, *Energy Convers. Manag.*, 2018, **159**, 264–273.
- 104 N. Muradov, Thermocatalytic CO₂-free production of hydrogen from hydrocarbon fuels, in *Proceedings of the 2000 Hydrogen Program Review*, NREL/CP-570-28890, 2000.
- 105 M. Steinberg and H. C. Cheng, *Int. J. Hydrogen Energy*, 1989, **14**, 797–820.
- 106 V. S. Sikarwar, M. Zhao, P. Clough, J. Yao, X. Zhong, M. Z. Memon, N. Shah, E. J. Anthony and P. S. Fennell, *Energy Environ. Sci.*, 2016, **9**, 2939–2977.
- 107 R. Wisler, K. Jenni, J. Seel, E. Baker, M. Hand, E. Lantz and A. Smith, *Nat. Energy*, 2016, **1**, 16135.



- 108 P. Frankl, S. Nowak, M. Gutschner, S. Gnos and T. Rinke, International Energy Association, 2010, <https://www.iea.org/reports/technology-roadmap-solar-photovoltaic-energy-2010>.
- 109 World Nuclear Association, Economics of Nuclear Power, <http://www.world-nuclear.org/information-library/economic-aspects/economics-of-nuclear-power.aspx>.
- 110 IRENA, Concentrating Sol. Power, IRENA working paper, 2012, https://www.irena.org/-/media/Files/IRENA/Agency/Publication/2012/RE_Technologies_Cost_Analysis-CSP.pdf.
- 111 M. R. Shaner, H. A. Atwater, N. S. Lewis and E. W. McFarland, *Energy Environ. Sci.*, 2016, **9**, 2354–2371.
- 112 J. P. Marshall, *Energy Policy*, 2016, **99**, 288–298.
- 113 A. B. Rao and E. S. Rubin, *Ind. Eng. Chem. Res.*, 2006, **45**, 2421–2429.
- 114 A. J. Hunt, E. H. Sin, R. Marriott and J. H. Clark, *ChemSusChem*, 2010, **3**, 306–322.
- 115 T. S. Chung, D. Patiño-Echeverri and T. L. Johnson, *Energy Policy*, 2011, **39**, 5609–5620.
- 116 R. Bell, *The Production of Low Carbon Gas – Consultation Response: SCCS response to the Carbon Connect consultation on the production of low carbon gas*, 2018.
- 117 M. Bui, C. S. Adjiman, A. Bardow, E. J. Anthony, A. Boston, S. Brown, P. S. Fennell, S. Fuss, A. Galindo and L. A. Hackett, *Energy Environ. Sci.*, 2018, **11**, 1062–1176.
- 118 IEAGHG, *Enabling the deployment of industrial CCS clusters*, 2018.
- 119 Global CCS Institute, *The Global Status of CCS. Special Report: Understanding the Industrial CCS Hubs and Clusters*, Melbourne, Australia, 2016.
- 120 i24c, *Deployment of an industrial carbon capture and storage cluster in Europe: A funding pathway*, Cambridge, UK, 2017.
- 121 I. Enting, D. Etheridge and M. J. Fielding, *Int. J. Greenhouse Gas Control*, 2008, **2**, 289–296.
- 122 P. M. Haugan and F. Joos, Metrics to assess the mitigation of global warming by carbon capture and storage in the ocean and in geological reservoirs, *Geophys. Res. Lett.*, 2004, **31**, L18202, DOI: 10.1029/2004GL020295.
- 123 C. R. Jenkins, P. J. Cook, J. Ennis-King, J. Undershultz, C. Boreham, T. Dance, P. de Caritat, D. M. Etheridge, B. M. Freifeld and A. Hortle, *Proc. Natl. Acad. Sci. U. S. A.*, 2012, **109**, E35–E41.
- 124 K. C. Mondal and S. Ramesh Chandran, *Int. J. Hydrogen Energy*, 2014, **39**, 9670–9674.
- 125 G. Kreysa, *ChemSusChem*, 2009, **2**, 49–55.
- 126 R. K. Pachauri, M. R. Allen, V. R. Barros, J. Broome, W. Cramer, R. Christ, J. A. Church, L. Clarke, Q. Dahe, P. Dasgupta, N. K. Dubash, et al., *Climate change 2014: synthesis report. Contribution of Working Groups I, II and III to the fifth assessment report of the Intergovernmental Panel on Climate Change*, IPCC, 2014, pp. 151.
- 127 C. Munnings and A. Krupnick, *Comparing policies to reduce methane emissions in the natural gas sector*, Washington, DC: Resources for the Future, 2017.
- 128 J. Banks, Barriers and Opportunities for Reducing Methane Emissions from Coal Mines, Clear Air Task Force, https://www.catf.us/wp-content/uploads/2019/10/201209-Barriers_and_Opportunities_in_Coal_Mine_Methane_Abatement.pdf.
- 129 U.S. Environmental Protection Agency, *Coal Mine Methane (CMM) Finance Guide*, 2009.
- 130 H. Yang, Z. Xu, M. Fan, R. Gupta, R. B. Slimane, A. E. Bland and I. Wright, *J. Environ. Sci.*, 2008, **20**, 14–27.
- 131 N. Muradov, *Catal. Commun.*, 2001, **2**, 89–94.
- 132 T. Abbasi and S. Abbasi, *Renewable Sustainable Energy Rev.*, 2011, **15**, 1828–1834.
- 133 M.-S. Liao and Q.-E. Zhang, *J. Mol. Catal. A: Chem.*, 1998, **136**, 185–194.
- 134 R. Aiello, J. E. Fiscus, H.-C. Zur Loye and M. D. Amiridis, *Appl. Catal. A*, 2000, **192**, 227–234.
- 135 P. Tang, Q. Zhu, Z. Wu and D. Ma, *Energy Environ. Sci.*, 2014, **7**, 2580–2591.
- 136 T. Geißler, A. Abánades, A. Heinzl, K. Mehravaran, G. Müller, R. Rathnam, C. Rubbia, D. Salmieri, L. Stoppel and S. Stückrad, *Chem. Eng. J.*, 2016, **299**, 192–200.
- 137 M. D. Eisaman, J. L. B. Rivest, S. D. Karnitz, C.-F. de Lannoy, A. Jose, R. W. DeVaul and K. Hannun, *Int. J. Greenhouse Gas Control*, 2018, **70**, 254–261.
- 138 R. Socolow, M. Desmond, R. Aines, J. Blackstock, O. Bolland, T. Kaarsberg, N. Lewis, M. Mazzotti, A. Pfeffer and K. Sawyer, *Direct air capture of CO₂ with chemicals: a technology assessment for the APS Panel on Public Affairs*, American Physical Society, 2011.
- 139 World Energy Council, *World Energy Resources*, World Energy Council, London, UK, 2016.
- 140 UK Government, International Industrial Energy Prices, <https://www.gov.uk/government/statistical-data-sets/international-industrial-energy-prices>.
- 141 European Environment Agency, Electricity Generation – CO₂ emissions intensity, 2018, [https://www.eea.europa.eu/data-and-maps/daviz/co2-emission-intensity-3#tab-googlechartid_chart_11_filters=%7B%22rowFilters%22%3A%7B%7D%3B%22columnFilters%22%3A%7B%22pre_config_ugeo%22%3A%5B%22European%20Union%20\(28%20countries\)%22%5D%7D%7D](https://www.eea.europa.eu/data-and-maps/daviz/co2-emission-intensity-3#tab-googlechartid_chart_11_filters=%7B%22rowFilters%22%3A%7B%7D%3B%22columnFilters%22%3A%7B%22pre_config_ugeo%22%3A%5B%22European%20Union%20(28%20countries)%22%5D%7D%7D).
- 142 E. Lantz, M. Hand and R. Wisner, Past and Future Cost of Wind Energy: Preprint, National Renewable Energy Laboratory (NREL), Golden, CO, 2012, <https://www.nrel.gov/docs/fy12osti/54526.pdf>.
- 143 S. Almosni, A. Delamarre, Z. Jehl, D. Suchet, L. Cojocar, M. Giteau, B. Behaghel, A. Julian, C. Ibrahim and L. Tatry, *Sci. Technol. Adv. Mater.*, 2018, **19**, 336–369.



- 144 IRENA, *Renewable Power Generation Costs in 2017*, International Renewable Energy Agency, Abu Dhabi, 2018.
- 145 G. F. Hewitt and J. G. Collier, *Introduction to nuclear power*, CRC Press, 2000.
- 146 N. Sakaba, H. Sato, H. Ohashi, T. Nishihara and K. Kunitomi, *J. Nucl. Sci. Technol.*, 2008, **45**, 962–969.
- 147 J. Norman, G. Besenbruch, L. Brown, D. O'keefe and C. Allen, *Thermochemical water-splitting cycle, bench-scale investigations, and process engineering. Final report, February 1977–December 31, 1981*, GA Technologies, Inc., San Diego, CA (USA), 1982.
- 148 L. Brecher, S. Spewock and C. Warde, *Int. J. Hydrogen Energy*, 1977, **2**, 7–15.
- 149 N. E. Hultman, J. G. Koomey and D. M. Kammen, *Environ. Sci. Technol.*, 2007, 2087–2094.
- 150 J. R. Lovering, A. Yip and T. Nordhaus, *Energy Policy*, 2016, **91**, 371–382.
- 151 P. E. Dodds and W. McDowall, *A review of hydrogen production technologies for energy system models*, UKSHEC Working Paper No. 6, UCL Energy Institute, University College London, 2012.

