Life Cycle Inventories for Palladium on Niobium Phosphate (Pd/NbOPO₄) and Zirconium Oxide (ZrO₂) Catalysts

Energy Systems Division

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by

Kathryn Kingsbury and P. Thathiana Benavides

Energy Systems Division, Argonne National Laboratory

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CONTENTS

FIGURES	vi
TABLES	vi
ACKNOWLEDGEMENTS	vii
ABBREVIATIONS AND ACRONYMS	ix
EXECUTIVE SUMMARY	xi
1. INTRODUCTION	1
2. DATA AND METHODOLOGY	3
2.1 LIFE CYCLE INVENTORY OF Pd/NbOPO ₄ CATALYST	3
2.1.1 Niobium oxide production from pyrochlore mining and processing	4
2.1.2 Preparation of potassium niobate precursor by hydrothermal synthesis	6
2.1.3 Production of niobium phosphate support by sol–gel synthesis	7
2.1.4 Palladium metal production from PGM mining and processing	9
2.1.5 Production of Pd/NbOPO ₄ catalyst by incipient wetness impregnation	11
2.2 LIFE CYCLE INVENTORY OF ZrO ₂ CATALYST	13
2.2.1 Production of zircon by mining, beneficiation, and separation of mineral sands	14
2.2.2 Production of ZrO ₂ catalyst by caustic fusion and calcination	16
3. RESULTS AND DISCUSSION	18
3.1 SUMMARY OF LCA RESULTS	18
3.2 ANALYSIS OF FACTORS CONTRIBUTING TO CATALYST BURDEN	20
4. CONCLUSION	24
APPENDIX A: Calculation of Energy Consumption in KNbO ₃ Production	26
APPENDIX B: Calculation of Energy Consumption in NbOPO ₄ Production	27
APPENDIX C: Calculation of Energy Consumption in Pd/NbOPO ₄ Production	29
APPENDIX D: Inputs and Results using Market Allocation for Pd Production	30
REFERENCES	32

FIGURES

Figure 1. Supply chain featuring the primary material inputs and outputs for production of the Pd/NbOPO ₄ catalyst.....	3
Figure 2. Supply chain featuring the primary material inputs and outputs for production of the ZrO ₂ catalyst	13
Figure 3. Contribution of material inputs to the production of a) Pd/NbOPO ₄ and b) ZrO ₂	20
Figure 4. Breakdown of cradle-to-gate GHG emissions of a) Pd/NbOPO ₄ and b) ZrO ₂	21
Figure 5. Breakdown of cradle-to-gate fossil fuel consumption of a) Pd/NbOPO ₄ and b) ZrO ₂ .	21
Figure 6. Breakdown of cradle-to-gate water consumption of a) Pd/NbOPO ₄ and b) ZrO ₂	23

TABLES

Table 1. Material and energy inputs for the production of Nb ₂ O ₅	5
Table 2. Material and energy inputs for the production of KNbO ₃	7
Table 3. Material and energy inputs for the production of NbOPO ₄	9
Table 4. Material and energy inputs for the production of palladium	10
Table 5. Material and energy inputs for the production of Pd/NbOPO ₄	12
Table 6. Material and energy inputs for the production of zircon	15
Table 7. Material and energy inputs for the production of ZrO ₂	17
Table 8. Cradle-to-gate LCA results for new catalysts and associated materials	18
Table 9. Comparison of new Pd/NbOPO ₄ and ZrO ₂ catalysts to existing catalysts in GREET ...	19
Table D-1. Market allocation material and energy inputs for the production of palladium	30
Table D-2. Cradle-to-gate LCA results for Pd and Pd/NbOPO ₄ using market allocation inputs .	30

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ABBREVIATIONS AND ACRONYMS

4-BH	4-butoxyheptane
Al ₂ O ₃	aluminum oxide
Al ₂ (SO ₄) ₃	aluminum sulfate
Btu	British thermal units
CBMM	Companhia Brasileira de Metalurgia e Mineração
CO ₂ e	carbon dioxide equivalents
Co-Optima	DOE Co-Optimization of Fuels & Engines initiative
CSTR	Continuous-stir tank reactor
g	gram
gal	gallon
GHG	greenhouse gas
GREET	Greenhouse gases, Regulated Emissions, and Energy use in Technology
GWP	global warming potential
H ₂	hydrogen gas
HMC	heavy-mineral concentrate
HNO ₃	nitric acid
H ₃ PO ₄	phosphoric acid
H ₂ SO ₄	sulfuric acid
IWI	incipient wetness impregnation
K	Kelvin unit of temperature
kg	kilogram
kW	kilowatt
KNbO ₃	potassium niobate
KOH	potassium hydroxide
L	liter
LCA	life cycle analysis
LCI	life cycle inventory
M	molarity
MJ	megajoule
MSP	mineral separation plant
mmBtu	million British thermal units

N ₂	nitrogen gas
Na ₂ SiO ₃	sodium silicate
Na ₂ ZrO ₃	sodium zirconate
Nb	niobium
Nb ₂ O ₅	niobium oxide or niobium pentaoxide
NbOPO ₄	niobium phosphate
NREL	National Renewable Energy Laboratory
Pd	palladium
Pd/NbOPO ₄	palladium on niobium phosphate catalyst
PGM	platinum group metal
Pt	platinum
t	metric tonne
TEA	technoeconomic analysis
ton	U.S. short ton
ZIA	Zircon Industry Association
ZrO ₂	zirconium oxide or zirconia
ZrSiO ₄	zircon, zircon sand, or zirconium silicate

EXECUTIVE SUMMARY

Purpose

Catalysts are used in refining and chemical processes to maximize product yield and reaction selectivity, and they are critically important in optimizing biofuel production to make biofuels a viable alternative to traditional fossil-based fuels. However, catalysts are rarely included in life cycle analysis (LCA) studies of biofuels due to lack of information. The catalyst module in Argonne National Laboratory's Greenhouse gases, Regulated Emissions, and Energy use in Technologies (GREET) model addresses this issue with catalyst energy and material data for use in biofuels LCAs (GREET 2020, Wang et al. 2015). Nevertheless, many catalysts remain to be analyzed, especially as new biofuel production pathways emerge, and catalyst technology advances. We therefore proposed to expand the GREET catalyst module with the addition of new catalyst materials applicable to current biofuel production pathways.

Approach

Before beginning this study, we performed an extensive literature review to assess current catalyst technology for biofuel production, and we selected two catalysts for incorporation into GREET: 1) palladium on a niobium phosphate support (Pd/NbOPO₄), and 2) zirconium oxide (ZrO₂). We then framed our analysis by tracing a cradle-to-gate supply chain for each catalyst. The supply chain for the Pd/NbOPO₄ catalyst required assessment of four precursor materials that were not available in GREET: niobium phosphate (NbOPO₄), potassium niobate (KNbO₃), niobium oxide (Nb₂O₅), and palladium (Pd). The supply chain for the ZrO₂ catalyst required assessment of only one precursor material not already in GREET: zircon (ZrSiO₄). We collected life cycle inventory (LCI) data for the two catalysts and five associated materials using publicly-available information, and we applied engineering calculations to estimate energy inputs for processes that lacked industrial data. Finally, we used the GREET model to calculate cradle-to-gate greenhouse gas (GHG) emissions, fossil fuel consumption, and water consumption for the Pd/NbOPO₄ and ZrO₂ catalysts and their associated materials.

Outcomes

We report the cradle-to-gate GHG emissions, fossil fuel consumption, and water consumption for Nb₂O₅, KNbO₃, NbOPO₄, Pd, Pd/NbOPO₄, ZrSiO₄, and ZrO₂. Notably, the net GHG emissions impact of the Pd/NbOPO₄ catalyst is 8.5 kg CO₂e/kg catalyst, which is comparable to other catalysts in GREET, while the net GHG emissions of the ZrO₂ catalyst is considerably lower at 1.8 kg CO₂e/kg catalyst. We also identify the primary contributors to each catalyst's cradle-to-gate environmental burden. For the Pd/NbOPO₄ catalyst, Pd metal is the main driver of GHG emissions and fossil fuel consumption, while Pd and NbOPO₄ contribute almost equally to water consumption. For the ZrO₂ catalyst, sodium hydroxide (NaOH) is the principal driver of GHG emissions and fossil fuel consumption, while ZrSiO₄ is the main consumer of water. The Pd/NbOPO₄ and ZrO₂ catalysts, as well as Nb₂O₅, KNbO₃, NbOPO₄, and ZrSiO₄, are implemented in the GREET catalyst module, and Pd has been implemented in GREET2 (Kingsbury and Benavides 2021). The material and energy flows for these new materials will be useful to future LCAs, particularly those involving biofuel production.

1. INTRODUCTION

Catalysts are employed across all sectors of the chemical industry to control selectivity and improve product yield in chemical processes. They are especially vital to the production of biofuels, which face many technical and economic obstacles to their industrial viability (Gaspar et al. 2019). A biofuel's eligibility for widespread adoption depends on several factors, including whether its life cycle greenhouse gas (GHG) emissions meet the guidelines of policies such as the U.S. Renewable Fuel Standard (RFS) and the California Low Carbon Fuel Standard (LCFS) (Benavides et al. 2017). Life cycle analysis (LCA) is used to determine these life cycle GHG emissions and other sustainability information about a biofuel. LCA is a valuable method for evaluating a product's environmental impact because it examines all stages in the product's supply chain, from extraction and processing of all raw materials to use and disposal of the final product. However, when LCA of a biofuel is performed, the contribution of the catalysts used in its production is frequently not accounted for due to lack of data and their minimal mass compared with other process inputs. As the impacts of catalysts may be more significant than would be indicated by their small mass input to the process, it is relevant to provide inventories for the catalysts associated with new biofuel production pathways.

In this study, we specifically examine two catalysts that are relevant to biofuel production. The first is a single-phase catalyst composed of palladium metal on a niobium phosphate support (Pd/NbOPO_4) developed by Hafenstine et al. (2020), and the second is a traditional zirconium oxide (ZrO_2) catalyst. We have selected these catalysts for a variety of reasons. Both catalysts may be applied in the conversion of biomass to 4-butoxyheptane (4-BH), an ether diesel bioblendstock that is currently of interest to the U.S. Department of Energy's (DOE's) Co-Optimization of Fuels & Engines (Co-Optima) initiative due to its outstanding fuel properties and potential for industrial viability (Huq et al. 2019). The production of 4-BH from biomass requires multiple chemical reactions and thus multiple catalysts: ZrO_2 serves as a ketonization catalyst and Pd/NbOPO_4 as a reductive etherification catalyst. The Pd/NbOPO_4 catalyst is especially noteworthy because the Pd active phase assists in hydrogenation chemistry, while the NbOPO_4 support acts as a solid acid catalyst for ketalization and dehydration chemistry (Hafenstine et al. 2020). Both the ZrO_2 and Pd/NbOPO_4 catalysts have displayed high conversion rates for their respective reactions during 4-BH production (Hafenstine et al. 2020).

Both ZrO_2 and Pd/NbOPO_4 also show promise as effective catalysts for producing biofuels other than 4-BH. For the synthesis of biodiesel, ZrO_2 has been successfully used as a catalyst in a sulfated form, in an acid-modified form, and as a support for tungstates (ZIA 2019). There are also ongoing efforts to use ZrO_2 -supported metal catalysts in the reforming of gasoline, diesel, butane, propane, and ethanol (ZIA 2019). Most notably, ZrO_2 has already been employed in a study by the National Renewable Energy Laboratory (NREL) as a ketonization catalyst during production of a diesel and jet fuel range blendstock (Davis et al. 2018). The Pd/NbOPO_4 catalyst is also very likely to be applicable in additional biofuel production pathways due to the valuable catalytic properties of Pd and Nb compounds. Palladium is a

member of the platinum group metals (PGMs), which are widely used in catalysis due to their unique chemical properties and high degree of stability (Kettler 2003). Palladium and other PGM catalysts are especially relevant to the biofuel industry because they can assist reactions such as hydrogenations and oxidations that are common in many different biofuel production pathways. Niobium compounds are also of great interest in the field of catalysis due to their ability to function in different scenarios as catalyst active phases, catalyst supports, solid acid catalysts, or redox materials (Nowak and Ziolk 1999). Niobic acid, which is analogous to hydrated Nb_2O_5 , has been used as a ketone condensation catalyst in the same pathway as ZrO_2 to produce diesel and jet fuel range blendstock (Davis et al. 2018). Niobium phosphate is also especially useful due to its strong surface acidity even at the high temperatures used in pretreatment processes (Nowak and Ziolk 1999). Many opportunities therefore exist for application of our chosen catalysts in future LCA studies.

In this work, we examine the consumption of material and energy inputs throughout the life cycle of the Pd/NbOPO_4 and ZrO_2 catalysts. The scope of our analysis for both catalysts is cradle-to-gate, which refers to all stages in catalyst production from the extraction and processing of raw materials up to the manufacturing facility gate. In Section 2.1, we define a supply chain for Pd/NbOPO_4 and describe the collection of life cycle inventory (LCI) data from various sources detailing the individual stages in the Pd/NbOPO_4 catalyst supply chain. We specifically examine five primary stages: 1) production of Nb_2O_5 from the mining of pyrochlore, 2) chemical synthesis of KNbO_3 from Nb_2O_5 , 3) chemical synthesis of NbOPO_4 from KNbO_3 , 4) production of Pd metal from the mining and processing of PGMs, and 5) synthesis of the Pd/NbOPO_4 catalyst by incipient wetness impregnation (IWI) of the Pd metal onto the NbOPO_4 support. In Section 2.2, we define the supply chain and LCI collection for the ZrO_2 catalyst and examine two primary production stages: 1) production of zircon from the mining and beneficiation of heavy-mineral sands, and 2) ZrO_2 catalyst production by caustic fusion and calcination of zircon. If co-products are produced alongside the catalyst, we apply a mass allocation approach that allocates material and energy consumption to the major product, thus remaining consistent with existing materials in the GREET catalyst module (GREET 2020). In Section 3, we use the GREET model to calculate cradle-to-gate values for GHG emissions, fossil fuel consumption, and water consumption for all new materials. GHG emissions are reported in units of $\text{kg CO}_2\text{e}$, fossil fuel consumption in units of MJ, and water consumption in units of gal. All three metrics are reported per kg of catalyst or associated material produced. While we do not report or discuss LCA results for the generation of criteria air pollutants (CAPs) in this work, these values are available in GREET for the new catalysts and associated materials (GREET 2020). Finally, we provide a discussion of the factors that contribute to the net environmental impact of the Pd/NbOPO_4 and ZrO_2 catalysts that will be beneficial to the design and selection of new catalyst materials. Both catalysts and their associated materials are implemented in the GREET model for use in future LCA studies.

2. DATA AND METHODOLOGY

2.1 LIFE CYCLE INVENTORY OF Pd/NbOPO₄ CATALYST

A product's LCI identifies and quantifies all resources and energy used throughout the product's life cycle. Using the LCI, an LCA can then assess the product's total environmental impact. Before compiling LCI data for the Pd/NbOPO₄ catalyst, it is necessary to identify a supply chain for the catalyst that traces each contributing material back to the beginning of its own life cycle. Many different pathways may be followed to eventually produce the Pd/NbOPO₄ catalyst, but data on industrial catalyst production is very limited. The Pd/NbOPO₄ supply chain that we show in Figure 1 is thus based on a combination of processes for which material and energy inputs are either publicly available or can be calculated using chemical and engineering principles. As depicted in Figure 1, niobium oxide (Nb₂O₅) is produced from the mining and processing of pyrochlore ore. Niobium oxide is then converted to an intermediate chemical, potassium niobate (KNbO₃), which is subsequently converted to the niobium phosphate (NbOPO₄) catalyst support. Palladium (Pd) metal, which is produced from the mining and processing of the platinum group metals (PGMs), is then added to the niobium phosphate support, and several processing steps are performed to produce the final Pd/NbOPO₄ catalyst. In the following subsections, we describe each of these production stages in greater detail.

Figure 1. Supply chain featuring the primary material inputs and outputs for production of the Pd/NbOPO₄ catalyst

To obtain LCI data for the Pd/NbOPO₄ catalyst, we consulted a variety of sources, including scientific literature, technical reports from Argonne and other national laboratories, sustainability reports from mining and chemical companies, information already in the GREET model, and direct correspondence with experts in the field. The final step in the production of the Pd/NbOPO₄ catalyst, IWI, is described by Hafenstine et al. (2020). This process requires Pd metal, NbOPO₄, nitric acid (HNO₃), hydrogen (H₂) gas, and nitrogen (N₂) gas. Data on the last three materials can be found in the GREET model (GREET 2020). The production of Pd as a PGM and its implementation in GREET is described by Kingsbury and Benavides (2021). The NbOPO₄ support can be produced from the reaction of potassium niobate (KNbO₃) and phosphoric acid (H₃PO₄) using a sol–gel method detailed by both Wang et al. (2018) and He et al. (2020). Life-cycle data for H₃PO₄ is already available in the GREET model (GREET 2020), and we traced KNbO₃ through its synthesis from the hydrothermal reaction of Nb₂O₅ with potassium hydroxide (KOH) as described by Lu et al. (1998). Life-cycle data for KOH is found in the GREET model (GREET 2020), and we describe the production of Nb₂O₅ from pyrochlore as reported by Companhia Brasileira de Metalurgia e Mineração (CBMM 2019). Note that for the IWI synthesis of the Pd/NbOPO₄ catalyst, the sol–gel synthesis of NbOPO₄, and the hydrothermal synthesis of KNbO₃, only laboratory-scale material inputs and processing steps were available. We therefore use stoichiometric ratios to estimate material requirements on an industrial scale, and we utilize an estimation technique devised by Majeau-Bettez et al. (2011) and presented in Dunn et al. (2015) to calculate the energy consumption required for these processes.

2.1.1 Niobium oxide production from pyrochlore mining and processing

Niobium is not found in nature as a free element and is most commonly sourced from the mineral pyrochlore, which is found within alkaline igneous rock deposits (Schulz et al. 2017). Brazil contains the world’s largest deposits of pyrochlore and serves as the global leader in niobium production, responsible for 88% of all niobium production worldwide (Padilla 2020). Based in Araxà, Brazil, CBMM owns the largest share of pyrochlore mines and produces more niobium products each year than any other company, so for the purposes of our study we focused on the production methods and data from this company’s official sustainability reports (CBMM 2019). These pyrochlore deposits contain an average of 2.5% Nb₂O₅ content, so several processes are required to concentrate the Nb₂O₅ and separate it from the other materials also obtained from pyrochlore (CBMM 2019).

Pyrochlore ore is mined using open-pit methods and transported by electric-powered conveyor belts to the blending yard and concentration plant where the non-niobium-containing components of the pyrochlore are removed by grinding, magnetic separation, desliming, and flotation. Pyrometallurgy processing of the niobium concentrate removes contaminants such as phosphorous, sulfur, and lead. Depending on the desired end product, the niobium concentrate is further processed in a variety of ways. Although approximately 90% of the niobium that is extracted from pyrochlore is processed and sold as ferroniobium, an iron-niobium alloy used in steel materials, CBMM also produces a significant amount of laboratory-grade niobium

chemicals like Nb₂O₅. In this case, the niobium concentrate is taken to a high-purity niobium oxide plant and a specialized oxides plant where it is purified to 99.5% Nb₂O₅ (CBMM 2019). CBMM conducts all mining, concentrating, and production processes on-site and exports only finished niobium products.

CBMM sustainability reports for 2016, 2017, 2018, and 2019 include the total niobium production, material inputs, and energy consumption per year. The average yearly production of total niobium products during this time period was 862,750 tons. Approximately 89% of this total consisted of ferroniobium products, with the remaining 11% of production being composed of other niobium products such as Nb₂O₅. CBMM details the amount of petroleum coke, charcoal, water, electricity, diesel fuel, and liquefied petroleum gas that is consumed during the overall production process each year. This data represents all of the materials and energy that went into producing all niobium products, beginning with the mining of pyrochlore ore and ending at the point of commercial distribution of the final niobium products. CBMM also reports minimal consumption of aviation fuel and biodiesel, but we omitted these from our inventory because their combined contribution to the total energy input was less than 0.1% and because detailed information regarding their exact application in Nb₂O₅ production was not provided. Using a mass allocation approach, we determined the material and energy inputs per ton of Nb₂O₅ product (Table 1). Upstream information regarding petroleum coke, diesel fuel, liquefied petroleum gas, and electricity generation in Brazil is available in GREET (GREET 2020). To determine upstream information for charcoal, we use the same approach presented in Benavides et al. (2015) and model charcoal production from the pyrolysis of willow, the energy and emissions burdens of which are detailed in GREET (GREET 2020).

Table 1. Material and energy inputs for the production of Nb₂O₅

Material	Amount per ton Nb₂O₅	
Petroleum coke	0.0037 tons	
Charcoal	0.0135 tons	
Water	542 gal	
Energy	mmBtu/ton Nb₂O₅	Share %
Electricity	1.212	72%
Diesel	0.149	9%
Liquefied petroleum gas	0.320	19%
Total energy input	1.681	

2.1.2 Preparation of potassium niobate precursor by hydrothermal synthesis

Potassium niobate (KNbO_3) can be synthesized from Nb_2O_5 and KOH using a variety of methods such as hydrothermal, solid-state, and precipitation reactions. We chose to explore the hydrothermal method because it requires significantly lower reaction temperatures compared to the other two methods (Lu et al. 1998), and it is associated with high product yield and quick reaction time (Rashid et al. 2018). Hydrothermal synthesis occurs within a sealed vessel, usually an autoclave, where extremely high pressures enable a solvent to be heated past its normal boiling point without evaporating. The high temperature and pressure promote the rapid interaction and assembly of the particles dissolved in the solvent (Rashid et al. 2018).

We examined a procedure for the laboratory-scale hydrothermal synthesis of KNbO_3 described by Lu et al. (1998). In this procedure, solid KOH is dissolved in water to prepare an 8 M KOH solution, which is then mixed with 3.32 g of Nb_2O_5 . The total volume of solution is set to 125 mL, so the Nb_2O_5 has a concentration of 0.1 M. The solution is placed into a sealed autoclave reactor and heated at 200°C for 2 hours while being continuously stirred. After hydrothermal reaction (see Equation 1) the KNbO_3 product obtained is washed and dried.

To estimate the quantities of materials used, we assumed a 100% yield of the KNbO_3 product and used the reaction's stoichiometric ratios to relate the mass inputs of Nb_2O_5 , the limiting reagent, to the mass of KNbO_3 produced. Since excess KOH is used, we assumed that all KOH that is not consumed during the reaction is recycled and may be re-used as a reactant. Finally, we assumed that the H_2O formed during reaction can be recycled for use as a solvent, and we used a conservative estimate that 50% of the water used as a solvent may be recycled by purification.

To estimate energy consumption for the production of KNbO_3 , we adopted the methodology of Majeau-Bettez et al. (2011) and Dunn et al. (2015). In this method, the total energy of production is equivalent to the sum of the energy needed to heat any materials plus the energy required to power any equipment used. On an industrial scale, the hydrothermal reaction may be performed in a continuous-stir tank reactor (CSTR). The total energy (q_{t,KNbO_3}) required to produce 1 ton of KNbO_3 is calculated as the sum of the heat required to heat the reaction solvent (q_{solvent}), the heat loss of the reactor (q_{CSTR}), and the energy used in stirring the materials (q_{stir}). The energy required to dry the product is assumed to be negligible in this scenario as the product does not require intensive heat-treating. A key assumption in the total energy calculation is that half of the heat used to heat the solvent may be recovered and re-incorporated by heat integration, causing the value of q_{solvent} to be halved (Equation 2).

$$q_{t,\text{KNbO}_3} = \frac{q_{\text{solvent}}}{2} + q_{\text{CSTR}} + q_{\text{stir}} \quad (2)$$

A detailed account of the calculations for each term in Equation 2 can be found in Appendix A. The volume of the reactor is assumed to be 10,000 L, which we used to scale the energy consumption according to the amount of material required to produce 1 ton of KNbO_3 . Heating of the reactor and solvent consumes natural gas that is combusted in a boiler with a 90% efficiency, while the energy used for stirring is electricity. Table 2 summarizes the total material and energy inputs for the production of KNbO_3 from the hydrothermal reaction of Nb_2O_5 and KOH .

Table 2. Material and energy inputs for the production of KNbO_3

Material	Amount per ton KNbO_3	
Nb_2O_5	0.738 tons	
KOH	0.312 tons	
Water	3,325 gal	
Energy	mmBtu/ton KNbO_3	Share %
Electricity	0.100	1%
Natural gas	7.886	98%
<i>Total energy input</i>	7.986	

Because the hydrothermal method requires continuous maintenance of high pressures and involves the heating of solvents with high specific heat capacities, it is quite energy intensive in terms of natural gas and water consumption.

2.1.3 Production of niobium phosphate support by sol–gel synthesis

Niobium phosphate (NbOPO_4) is commonly produced by the reaction of potassium niobate (KNbO_3) with phosphoric acid (H_3PO_4). This approach is used by Wang et al. (2018) and He et al. (2020), who both describe a sol–gel method for this reaction. This method involves the stimulation of small molecules to form a colloidal solution, which then evolves to form a gel with a liquid and a solid phase. The liquid phase is then removed by drying the solvent (Rashid et al. 2018).

We analyzed a laboratory-scale sol–gel synthesis of NbOPO_4 that draws from the procedures of both Wang et al. (2018) and He et al. (2020). In this process, 1.15 g of 85% H_3PO_4 is dissolved in 25 mL of water, and a 0.1 M KNbO_3 solution is added to the H_3PO_4 solution. Using the stoichiometric ratios of the reaction (see Equation 3), we determined that 1.80 g of pure KNbO_3 is reacted with the H_3PO_4 and that the total volume of solution is 124.75 mL, giving a concentration of 0.4 M for H_3PO_4 . The solution is stirred for 1 hour at 40°C and for 2 additional hours at 80°C. It is then heated in a sealed autoclave reactor at 160°C for 24 hours. After cooling, the product is filtered, washed, and dried at 110°C overnight or for approximately 12 hours. Finally, the solid product is calcined at 400°C for 3 hours to obtain a catalyst-quality NbOPO_4 product.

In order to estimate the quantities of material in industrial production, we assumed a 100% yield of the NbOPO₄ product and used the reaction's stoichiometric ratios to relate the mass inputs of KNbO₃ and H₃PO₄ to the mass of NbOPO₄ produced. We considered KOH to be a coproduct of NbOPO₄, and we used a mass allocation approach to calculate the environmental burden of this synthesis based on 79% of the products consisting of NbOPO₄ and 21% consisting of KOH. We again assumed that the H₂O formed during reaction can be recycled for use as a solvent, and we used a conservative estimate that 50% of the total water that is used as a solvent may be recycled after purification.

To estimate energy consumption for the production of NbOPO₄, we adopted the methodology of Majeau-Bettez et al. (2011) and Dunn et al. (2015) as we did with the production of KNbO₃, but we expanded the total energy consumption to include drying and calcining, as these are important steps in achieving a catalyst-grade product. On an industrial scale, reaction and solvent heating within the autoclave may be modeled using a CSTR, while drying occurs in a chamber oven and calcining in a chamber furnace. The total energy (q_{t,NbOPO_4}) required to produce 1 ton of NbOPO₄ is calculated as the sum of the heat required to heat the reaction solvent (q_{solvent}), the heat loss by the reactor (q_{CSTR}), the heat required to dry the NbOPO₄ product (q_{dry}), the energy used in heating the oven (q_{oven}), the heat required to calcine the NbOPO₄ product (q_{calcine}), the energy used in heating the furnace (q_{furnace}), and any energy used in stirring the materials (q_{stir}). We assume that half of the heat consumed during the heating of any materials, either in solution or as dry NbOPO₄, may be recovered and re-incorporated into the process by heat integration, causing the values of q_{solvent} , q_{dry} , and q_{calcine} to be halved (Equation 4).

$$q_{t,\text{NbOPO}_4} = \frac{q_{\text{solvent}}}{2} + q_{\text{CSTR}} + \frac{q_{\text{dry}}}{2} + q_{\text{oven}} + \frac{q_{\text{calcine}}}{2} + q_{\text{furnace}} + q_{\text{stir}} \quad (4)$$

More details regarding the calculations for each term in Equation 4 can be found in Appendix B. As with the production of KNbO₃, we scaled the energy consumption to 1 ton of NbOPO₄ by relating the interior volume of any equipment used to the amount of material that must be loaded into the equipment in order to produce 1 ton of NbOPO₄. The same steam-powered CSTR with a volume of 10,000 L that is described by Majeau-Bettez et al. (2011) may be used to heat the solvent in the first step. We modeled the drying step using a 1,000 L chamber oven and the calcining step using an 8,300 L chamber furnace. Both the chamber oven and chamber furnace are from Nabertherm and are powered by electricity (Nabertherm 2016).

Table 3 summarizes the total material and energy inputs for the production of NbOPO₄ from the sol–gel reaction of KNbO₃ and H₃PO₄.

Table 3. Material and energy inputs for the production of NbOPO₄

Material	Amount per ton NbOPO₄	
KNbO ₃	0.692 tons	
H ₃ PO ₄	0.377 tons	
Water	5,756 gal	
Energy	mmBtu/ton NbOPO₄	Share %
Electricity	0.446	3%
Natural gas	13.341	97%
<i>Total energy input</i>	13.788	

The production of NbOPO₄ is quite intensive in terms of natural gas and water consumption. A large amount of water is required to obtain the correct concentration of reactants during the sol–gel synthesis procedure. The need to heat all of this water in the CSTR leads to heavy consumption of the natural gas used to power the CSTR.

2.1.4 Palladium metal production from PGM mining and processing

Palladium metal is produced in conjunction with the other platinum group metals (PGMs): platinum, palladium, rhodium, ruthenium, iridium, and osmium. PGMs are found as mixed metal and mineral deposits along with gold, nickel and copper within basic igneous rocks (Seymour and O’Farrelly 2012). The Bushveld Igneous Complex in South Africa contains the largest PGM deposits in the world and produces over 70% of the world’s supply of PGMs. These deposits contain 5-8 g of PGM content per metric tonne (t) of ore, and 20% to 25% of this PGM content is palladium (Seymour and O’Farrelly 2012). This gives a maximum palladium content of only 0.0002% in the ore, so intensive processing is required to procure palladium from these mixed ore deposits. The typical process stages for PGM production are mining, concentrating, smelting, and refining (Benavides et al. 2015).

Conventional underground or open cut techniques are used for mining the PGM ore, which is located at depths between 500 m and 2 km in South African deposits, requiring large energy inputs for drilling, ore hauling, and refrigeration (Benavides et al. 2015). The mined ore is then crushed and milled into small particles and mixed with water to form a slurry for concentration, where froth flotation is employed to separate waste from the valuable minerals, producing a concentrate with a 100-150 g/t PGM content (Seymour and O’Farrelly 2012). Gravity separation may be used at this point to further separate larger PGM particles, while the majority of the concentrate is smelted in electric furnaces and slow-cooled to produce a matte with a 1500 g/t PGM content as well as nickel, copper, and iron sulfides. (Seymour and O’Farrelly 2012). The iron and sulfur content are decreased with oxygen-enriched air, and

magnetic separation removes nickel and copper from the rest of the PGMs, producing a final concentrate with a 50%-60% PGM content. Finally, a variety of refining techniques such as solvent extraction, distillation, and ion-exchange further increase PGM content in the concentrate and separate the PGMs from each other. In the case of palladium, an aqua regia process is generally used to dissolve the PGM concentrate before treatment with ammonium hydroxide and hydrochloric acid to precipitate a solid palladium complex (Seymour and O’Farrelly 2012).

Kingsbury and Benavides (2021) provide water and energy inputs for the production of palladium by mining, concentrating, smelting, and refining the PGMs. This data, along with inputs for three other PGM production pathways, has already been incorporated into the GREET2 model (GREET 2020). The data reported by Kingsbury and Benavides (2021) is sourced from Anglo American Platinum, a South African mining company that operates seven mines in the Bushveld Complex. From 2015 to 2019, Anglo American Platinum reported its average annual energy and water consumption for the production of all their products: palladium, platinum, rhodium, gold, iridium, ruthenium, nickel, and copper. These reported values correspond to the total inputs across the mining, concentrating, smelting, and refining stages. Information about potential process chemicals or materials used in PGM refining and separation was not provided by Anglo American Platinum and thus is not included in this inventory, but we assume that their contribution to the total energy consumption is minimal. Kingsbury and Benavides (2021) determined the division of the total energy input by fuel type (electricity, diesel, coal, and natural gas). For consistency with other catalyst materials in the GREET model (GREET 2020), we used energy inputs for palladium that were calculated from the total production energy by mass allocation, which is set as the default method in GREET. Kingsbury and Benavides (2021) also discuss market allocation as an option for calculating the energy inputs per ton of Pd, so we provide these inputs and the corresponding LCA results in Appendix D. Palladium is a high-value metal, especially in comparison to nickel and copper which are also produced by Anglo American Platinum, so the energy consumption of Pd production is considerably higher when using market allocation as opposed to mass allocation. The inputs for the production of 1 ton of palladium using the results of the mass allocation approach (Kingsbury and Benavides 2021) are presented in Table 4.

Table 4. Material and energy inputs for the production of palladium

Material	Amount per ton Pd	
Water	171,176 gal	
Energy	mmBtu/ton Pd	Share %
Electricity	340.1	71%
Diesel	54.8	11%
Coal	80.1	17%
Natural gas	4.9	1%
Total energy input	479.1	

Palladium production is extremely resource-intensive due to the very low PGM content ($\leq 0.0002\%$) in mined ore, which requires considerable energy and water inputs for extraction and beneficiation. For comparison, the Nb_2O_5 content in the pyrochlore ore from which it is extracted is approximately 2.5%, leading to significantly smaller energy and material requirements. We note that palladium and other PGMs may also be recovered by recycling, or secondary production, which would significantly reduce their environmental burden. Secondary production of palladium involves smelting or dissolving a palladium-containing product, and then refining this molten matte or solution to separate out the palladium in a purified form (Benavides et al. 2015). Palladium and other PGMs display an extremely high recovery rate at 98%-99% (Seymour and O'Farrelly 2012) after undergoing the processes of secondary production. Information about the material and energy inputs for the secondary production of palladium was not available, so we did not account for this pathway in our inventory, but it would be worthwhile to examine in future LCA studies.

2.1.5 Production of Pd/NbOPO₄ catalyst by incipient wetness impregnation

To produce the Pd/NbOPO₄ catalyst, Pd is deposited on the NbOPO₄ support using IWI. In this method, the active phase metal is first dissolved in an acid and then drawn into the pores of the catalyst support through capillary action. Calcination then eliminates any volatile compounds from the metal or support (Rashid et al. 2018).

We examined the procedure of Hafenstine et al. (2020) for the IWI synthesis of Pd/NbOPO₄. Pd metal is dissolved in a nitric acid (HNO_3) solution and mixed with NbOPO₄ for 12 hours. The catalyst slurry is dried under air at 107°C for 6 hours and at 265°C for 2 hours. The dried product is ground and sieved into mesh particles. The product is then purged under nitrogen (N_2) gas for 1 hour. Finally, Pd ions are reduced to Pd metal by heating the product to 265°C and reducing with hydrogen (H_2) gas for 5 hours, forming the Pd/NbOPO₄ catalyst.

Quantities of materials for industrial production of Pd/NbOPO₄ were provided by direct correspondence with manufacturers of the catalyst (G. Hafenstine, pers. comm.). We also consulted the parameters for Pd/NbOPO₄ production listed in the CatCost software, a tool that allows researchers to estimate the large-scale production costs of pre-commercial catalysts (CatCost 2019) and which Hafenstine et al. (2020) used to incorporate the Pd/NbOPO₄ catalyst into a technoeconomic analysis (TEA). We verified that the material inputs and process steps that we used to estimate energy inputs for catalyst production were reflective of the procedure in CatCost. Hafenstine et al. (2020) designed the catalyst synthesis so that the Pd active phase in the Pd/NbOPO₄ catalyst is 5% by weight. The amount of N_2 and H_2 gas required for purging is based on a flow rate of 200 mL/min for both gases. We again assumed that 50% of the water that is used as a solvent may be recycled after purification.

We again estimated the energy consumption for the production of Pd/NbOPO₄ by taking the sum of the energy consumed in heating any materials plus the energy required to power any equipment (Majeau-Bettez et al. 2011, Dunn et al. 2015). In this scenario, we modeled the

stirring of materials using an electrically powered industrial chemical mixer that is capable of agitating catalyst slurries. The catalyst is dried in two stages at different temperatures, both of which may be modeled using a chamber oven, while heating of the catalyst under H₂ gas requires a tube furnace. The total energy ($q_{t,Pd/NbOPO_4}$) required to produce 1 ton of the Pd/NbOPO₄ catalyst is calculated as the sum of the heat required to dry the catalyst in two stages at 107°C (q_{dry_1}) and 265°C (q_{dry_2}), the heat used in heating the oven to 107°C (q_{oven_1}) and to 265°C (q_{oven_2}), the heat required to reduce the Pd/NbOPO₄ catalyst under H₂ gas (q_{reduce}), the energy used in heating the tube furnace for this reduction (q_{tube}), and the energy used in stirring the materials (q_{stir}). We assumed that half of the heat consumed by the Pd/NbOPO₄ material in any stage may be recovered and re-incorporated, causing the values of q_{dry_1} , q_{dry_2} , and q_{reduce} to be halved (Equation 5).

$$q_{t,Pd/NbOPO_4} = \frac{q_{dry_1}}{2} + q_{oven_1} + \frac{q_{dry_2}}{2} + q_{oven_2} + \frac{q_{reduce}}{2} + q_{tube} + q_{stir} \quad (5)$$

An explanation of each individual term in Equation 5 is provided in Appendix C. In order to scale the energy consumption to the production of 1 ton of Pd/NbOPO₄, we modeled both drying steps using a chamber oven with a volume of 1,050 L (Nabertherm 2016). While stirring was performed in the CSTR for the production of KNbO₃ and NbOPO₄, in this case the stirring is performed using a chemical mixer that can agitate catalyst slurries up to 1,892 L (Madden Pump 2020). The interior volume of the tube furnace used for reduction is calculated based on a provided tube radius of 50 mm and a heated length of 100 mm (Nabertherm 2017). Table 5 summarizes the total material and energy inputs for the production of Pd/NbOPO₄. Compared to the rest of the processing steps in the supply chain of the Pd/NbOPO₄ catalyst, the final synthesis of the catalyst by IWI is minimally energy intensive.

Table 5. Material and energy inputs for the production of Pd/NbOPO₄

Material	Amount per ton Pd/NbOPO ₄	
Pd	0.05 tons	
HNO ₃	0.05 tons	
NbOPO ₄	0.95 tons	
N ₂ gas	0.0058 tons	
H ₂ gas	0.0021 tons	
Water	72 gal	
Energy	mmBtu/ton Pd/NbOPO ₄	Share %
Electricity	0.496	100%
Total energy input	0.496	

2.2 LIFE CYCLE INVENTORY OF ZrO_2 CATALYST

As with our analysis of the Pd/NbOPO_4 catalyst, it is first necessary to identify a cradle-to-gate supply chain for production of the ZrO_2 catalyst based on publicly available data. ZrO_2 is a commonly used material in many industrial processes, and it may adopt several different crystal structures and morphologies with differing properties and applications (ZIA 2019). For the purposes of this study, we specifically outlined a pathway for the production of a pure monoclinic ZrO_2 catalyst, as this is the form used in the production pathway of the 4-BH biofuel (G. Hafenstine, pers. comm.). Zirconium compounds such as ZrO_2 may be sourced from many different zirconium-containing minerals, such as baddeleyite and eudialyte, but we limited our focus to zircon (ZrSiO_4) since this is the most commonly used commercial zirconium mineral (Nielsen et al. 2012).

As depicted in Figure 2, zircon is produced by the mining, beneficiation, and separation of mineral sands. Zircon is then decomposed by caustic fusion with NaOH to yield sodium zirconate (Na_2ZrO_3) as well as a sodium silicate (Na_2SiO_3) byproduct. Na_2ZrO_3 then hydrolyzes to form hydrous zirconia which is calcined at the production plant to form commercial ZrO_2 . Finally, the commercial ZrO_2 is further calcined at high temperatures to yield catalyst-grade ZrO_2 with a monoclinic crystal structure. We describe these production processes in further detail in the following subsections.

Figure 2. Supply chain featuring the primary material inputs and outputs for production of the ZrO_2 catalyst

We again consulted a variety of sources, from scientific literature to the GREET database, in order to determine the material and energy inputs for the LCI of the ZrO_2 catalyst. For the final calcination of commercial ZrO_2 to a monoclinic catalyst product, we examined a laboratory procedure from Shylesh et al. (2017), which requires no material inputs other than commercial ZrO_2 . For the decomposition of ZrSiO_4 to Na_2ZrO_3 , hydrolysis of Na_2ZrO_3 to hydrous zirconia, and calcination of hydrous zirconia to commercial ZrO_2 , we used an inventory from Primas (2007). Other than the ZrSiO_4 material input, the process steps in this inventory required only one additional material input, NaOH , the life cycle data of which is available in

GREET (GREET 2020). All inputs for the production of zircon from the mining, beneficiation, and separation of mineral sands are described by Gediga et al. (2019). Additional material inputs of NaOH, sulfuric acid (H_2SO_4), aluminum sulfate ($\text{Al}_2(\text{SO}_4)_3$), and lubricating oil were required for the production of ZrSiO_4 . Both NaOH and H_2SO_4 are detailed in GREET (GREET 2020). We used estimations to input $\text{Al}_2(\text{SO}_4)_3$ and lubricating oil, since these materials are not in GREET and contribute only very slightly to ZrSiO_4 production. $\text{Al}_2(\text{SO}_4)_3$ may be produced from the reaction of H_2SO_4 with aluminum oxide (Al_2O_3) (Darragh and Ertell 2003), both of which are available in the GREET model (GREET 2020), so we used stoichiometry to calculate the amount of Al_2O_3 and H_2SO_4 needed to produce the necessary amount of $\text{Al}_2(\text{SO}_4)_3$ for ZrSiO_4 production. We assumed no additional energy is required for this chemical reaction and included Al_2O_3 and H_2SO_4 as material inputs in place of $\text{Al}_2(\text{SO}_4)_3$. We used gasoline blendstock as a surrogate for lubricating oil, which is produced from the refining and distillation of crude oil much like gasoline (Sullivan 2005). When implementing vehicle fluids such as engine oil, which is analogous to lubricating oil, into GREET2, Burnham et al. (2006) estimated energy and emissions using gasoline manufacturing, so we adopted the same approach in our inventory.

2.2.1 Production of zircon by mining, beneficiation, and separation of mineral sands

Zircon forms naturally by crystallization as a mineral within silica-containing rocks such as granites, syenites, and diorites (ZIA 2019). The zircon contained in these primary ore deposits is largely inaccessible or minimally concentrated, so the majority (97%) of zircon is obtained from secondary placer deposits of heavy-mineral sands (ZIA 2019). These sand deposits have developed over millions of years as weathering and erosion have worn away at zircon-containing rocks, while wind and water have concentrated and transported the heavy zircon grains to coastline areas. Other minerals such as ilmenite (FeTiO_3) and rutile (TiO_2) undergo this same process and are found within the same heavy-mineral sands as zircon (ZIA 2019).

The production of zircon from heavy-mineral sands consists of three primary process stages: wet or dry mining, beneficiation (also referred to as concentration), and separation. These processes are described by ZIA (2019) and Gediga et al. (2019). Dry mining involves using equipment such as scrapers, bulldozers, and excavators to obtain heavy-mineral sands from solid inland deposits, while wet mining involves employing floating dredges to remove heavy-mineral sands from water ponds. The mined product, which contains about 10%-20% heavy-mineral content, is then concentrated. Concentration is typically performed by wet gravity separation methods that use equipment such as hydrocyclones or spirals to divide the heavy-mineral sands from other lighter and non-valuable materials, producing a heavy-mineral concentrate (HMC) with an 85%-95% heavy-mineral content. For wet mining, concentration occurs within the water source using a floating concentrator in conjunction with the floating dredge, while for dry mining, a slurry is produced from the mined heavy-mineral sands to input into the concentrator on site. After concentration, the HMC is transported to a mineral separation plant (MSP) where a combination of techniques, including screening as well as magnetic, electrostatic, and gravity separation circuits, is employed to separate zircon from other minerals in the HMC such as ilmenite and rutile.

Gediga et al. (2019) provide a comprehensive life cycle inventory for zircon production from mineral sands that covers more than 77% of global zircon production. Industry data for zircon production was collected from ten different production sites from eight companies across all major heavy-mineral sand-producing countries, including Australia, South Africa, Senegal, Kenya, and the United States. Both wet and dry mining methods were included in this survey. Unit process inputs and outputs for mining and beneficiation at each site's mining facilities and for separation at each site's MSP were obtained, and Gediga et al. (2019) report the material and energy inputs and outputs for these processes as a weighted average based on each site's contribution to the total production mass.

The final sand product consists of 37% zircon, 48% ilmenite, and 15% rutile. From this data, we calculated the total material and energy inputs for the mining, beneficiation, and separation of heavy-mineral sands, and we used mass allocation to determine the material and energy inputs per ton of zircon product. These inputs are shown in Table 6. As stated in the previous section, upstream information for all material and energy inputs are found in GREET (GREET 2020), with gasoline blendstock serving as a surrogate for lubricating oil, and Al_2O_3 and H_2SO_4 acting as stoichiometrically equivalent inputs in place of $\text{Al}_2(\text{SO}_4)_3$.

Table 6. Material and energy inputs for the production of zircon

Material	Amount per ton zircon	
H_2SO_4	0.00276 tons	
Al_2O_3	0.00005 tons	
NaOH	0.00010 tons	
Lubricating oil	0.00110 tons	
Water	5,981 gal	
Energy	mmBtu/ton zircon	Share %
Electricity	0.728	47%
Natural gas	0.289	18%
Diesel	0.536	35%
Total energy input	1.552	

The production of zircon by mining, beneficiation, and separation of heavy-mineral sands is much less energy-intensive than palladium production by PGM mining and is comparable to niobium production by pyrochlore mining. The large water consumption can be attributed to the wet concentration process in which the heavy-mineral sands are separated from lighter material in water using hydrocyclones and spirals.

2.2.2 Production of ZrO₂ catalyst by caustic fusion and calcination

Commercial ZrO₂ can be obtained from zircon by a variety of chemical, thermal, or mechanical decomposition methods (ZIA 2019). We focused on the decomposition of zircon by caustic fusion with sodium hydroxide, as this is the most commonly used zircon decomposition method (Nielsen and Wilfing 2012) and the process for which the most detailed data could be obtained. Caustic fusion entails reacting zircon (ZrSiO₄) with a slight excess of NaOH at 650°C to form sodium zirconate, sodium silicate, and water, as displayed in Equation 6.

After cooling, the products of this reaction are crushed to form a slurry. The majority of the sodium silicate is dissolved in water, while the sodium zirconate is hydrolyzed to form soluble NaOH and insoluble hydrous zirconia (Nielsen and Wilfing 2012). The hydrous zirconia can then be separated from the other products by filtration and calcined at temperatures between 800°C and 1000°C to form commercial ZrO₂ (Primas 2007).

Material and energy inputs for ZrO₂ production from zircon are presented by Primas (2007) as overall values for zircon decomposition by caustic fusion and calcination of hydrous zirconia to form commercial ZrO₂. Primas notes that the energy inputs reported for these processes are approximations based on very similar caustic fusion and calcination processes commonly used to produce sodium silicate and aluminum oxide. Primas calculated the material inputs using stoichiometry, with an assumed 10% excess of NaOH and a 95% product yield for zircon decomposition. The intermediate hydrolysis of sodium zirconate to hydrous zirconia is assumed to require no additional material or energy inputs. Primas reports a final product composition of 50% ZrO₂ and 50% sodium silicate, so we applied mass allocation to determine the material and energy inputs per ton of commercial ZrO₂.

Because we are specifically examining the production pathway for catalyst-grade, monoclinic ZrO₂, we incorporated an additional calcination step modeled after a laboratory procedure from Shylesh et al. (2017) that produces a monoclinic ZrO₂ catalyst by calcining purchased commercial ZrO₂ at a temperature of 1073 K for 3 hours. Using the same approach explained in the previous sections and in the appendices for the laboratory production of KNbO₃, NbOPO₄, and the Pd/NbOPO₄ catalyst, we calculated the energy requirement for calcination by modeling thermodynamic heating of ZrO₂ in an 8,300 L Nabertherm chamber furnace (Nabertherm 2016) according to the Majeau-Bettez et al. (2011) and Dunn et al. (2015) methodology. The energy required for this calcining step is equivalent to the sum of the energy needed to heat solid ZrO₂ to the required temperature (1073 K) and the energy used to heat an 8,300 L chamber furnace to that temperature for 3 hours. Only a small additional electricity input is required for this final calcination step, and we added this value to the energy consumption for the previous stages in ZrO₂ production. In Table 7, we present the material and energy inputs for the production of 1 ton of ZrO₂ catalyst as combined totals for the caustic fusion of zircon,

calcination of hydrous zirconia, and calcination of commercial ZrO_2 . All inputs apart from ZrSiO_4 , whose production we describe, are detailed in GREET (GREET 2020).

Table 7. Material and energy inputs for the production of ZrO_2

Material	Amount per ton ZrO_2	
ZrSiO ₄	0.785 tons	
NaOH	0.720 tons	
Water	194 gal	
Energy	mmBtu/ton ZrO_2	Share %
Electricity	0.440	20%
Natural gas	0.914	41%
Coal	0.023	1%
Residual oil	0.860	38%
<i>Total energy input</i>	2.237	

3. RESULTS AND DISCUSSION

3.1 SUMMARY OF LCA RESULTS

We used Argonne’s GREET model (GREET 2020) to perform cradle-to-gate LCAs for Nb_2O_5 , KNbO_3 , NbOPO_4 , Pd, the Pd/ NbOPO_4 catalyst, ZrSiO_4 , and the ZrO_2 catalyst. Fossil fuel consumption is reported in terms of energy unit (MJ) per mass unit (kg) of material produced. GHG emissions are reported in terms of mass unit (kg) of carbon dioxide equivalents (CO_2e) per mass unit (kg) of material produced. In order to report GHG emissions in terms of CO_2e , other GHGs such as CH_4 and N_2O are converted to their equivalent amounts of CO_2 using each gas’s global warming potential (GWP). Water consumption is reported in terms of volume unit (gal) per mass unit (kg) of material produced. The LCA results for all new catalyst materials examined in this study are shown in Table 8.

Table 8. Cradle-to-gate LCA results for new catalysts and associated materials

Cradle-to-gate result	Nb_2O_5	KNbO_3	NbOPO_4	Pd	Pd/ NbOPO_4 catalyst	ZrSiO_4	ZrO_2 catalyst
GHG emissions (kg CO_2e /kg material)	0.1	1.2	2.4	121	8.5	0.2	1.8
Fossil fuel consumption (MJ/kg material)	1.4	18.3	36.1	1,245	98.6	2.6	25.8
Water consumption (gal/kg material)	2.2	6.8	14.8	251	26.9	6.8	8.3

The cradle-to-gate GHG emissions, fossil fuel consumption, and water consumption values for Pd are significantly higher than the other catalyst materials, especially in comparison to the results for Nb_2O_5 and ZrSiO_4 , which are also produced by mining and beneficiation methods. This can largely be attributed to the fact that the Pd content within the mixed metal PGM ore from which it is mined is less than 0.0002%, compared to an Nb_2O_5 content of 2.5% in pyrochlore ore and a ZrSiO_4 content of 10%-20% in heavy-mineral sands. Much more intensive processes are thus required to extract and isolate Pd from its mined ore. However, it is important to note that metals, particularly those with a high economic value like the PGMs, often display a large environmental burden during their production, but their use and end-of-life recycling typically counterbalance this impact (Santero and Hendry 2016). Catalysts that incorporate PGMs generally display higher activity than base-metal catalysts, and PGM recovery rates after recycling are very high at 98%-99% (Seymour and O’Farrelly 2012). While the system boundary of this study is cradle-to-gate, a cradle-to-grave analysis that accounts for usage and recycling would be especially beneficial in the case of Pd and the Pd/ NbOPO_4 catalyst.

Below we provide a comparison of the LCA results for the new Pd/ NbOPO_4 and ZrO_2 catalysts to the results for existing catalysts in the GREET model. Table 9 shows the Pd/ NbOPO_4

and ZrO₂ catalyst results alongside results for three of the five catalysts previously implemented in GREET: Pt/ γ -Al₂O₃, MoCo/ γ -Al₂O₃, and ZSM-5 (GREET 2020, Wang et al. 2015).

Table 9. Comparison of new Pd/NbOPO₄ and ZrO₂ catalysts to existing catalysts in GREET

Cradle-to-gate result	Pd/NbOPO₄	ZrO₂	Pt/γ-Al₂O₃^a	MoCo/γ-Al₂O₃^a	ZSM-5^a
GHG emissions (kg CO ₂ e/kg material)	8.5	1.8	6.8	9.3	6.8
Fossil fuel consumption (MJ/kg material)	98.6	25.8	88.4	128.1	111.7
Water consumption (gal/kg material)	26.9	8.3	11.0	9.1	3.5

^a Results for Pt/ γ -Al₂O₃, MoCo/ γ -Al₂O₃, and ZSM-5 catalysts are from the GREET model (GREET 2020).

The net GHG emissions and fossil fuel consumption for Pd/NbOPO₄ are very comparable to the existing catalysts in GREET, while these values for the ZrO₂ catalyst are considerably lower. On the other hand, water consumption for ZrO₂ is very similar to the existing catalysts in GREET, while the water consumption for Pd/NbOPO₄ is markedly higher. We explore the contributing factors to the Pd/NbOPO₄ and ZrO₂ LCA results in the following section.

3.2 ANALYSIS OF FACTORS CONTRIBUTING TO CATALYST BURDEN

To assist our interpretation of the LCA results for the Pd/NbOPO₄ and ZrO₂ catalysts, we first broke down the different materials contributing to the production of these catalysts, as shown in Figure 3. The major material input for the Pd/NbOPO₄ catalyst is the NbOPO₄ support, which accounts for 90% of material inputs for the catalyst. We also show the contribution of Nb₂O₅, H₃PO₄, and KOH as material precursors for the NbOPO₄ support; the KNbO₃ intermediate is accounted for by the Nb₂O₅ and KOH inputs. As the primary source of Nb, Nb₂O₅ accounts for nearly half of the material inputs for the NbOPO₄ support. Palladium and HNO₃ each contribute approximately 4.5% of material inputs for the catalyst, while N₂ and H₂ gas inputs are minimal. For the ZrO₂ catalyst, only two major material inputs are used: zircon and NaOH. The percentage contributions of zircon and NaOH to the total material inputs are nearly the same, with zircon contributing just over 50% and NaOH just under 50% of the material composition.

Figure 3. Contribution of material inputs to the production of a) Pd/NbOPO₄ and b) ZrO₂

The figures below show the contributions of the various material and energy inputs for both catalysts to the cradle-to-gate GHG emissions (Figure 4) and fossil fuel consumption (Figure 5). The proportions of each input's contribution to GHG emissions and fossil fuel consumption are very similar, which is to be expected because processes that consume large amounts of fossil fuel are also likely to exhibit large amounts of GHG emissions. Although a small amount of the energy and emissions burden is associated with additional energy inputs (electricity, natural gas, residual oil, and coal) and estimated transportation requirements for catalyst production, the majority of the environmental impact for both catalysts comes from the upstream production of their material precursors. However, for both the Pd/NbOPO₄ and ZrO₂

catalysts, the percent mass contribution of each material input as shown in Figure 3 is not indicative of that material's contribution to GHG emissions and fossil fuel consumption.

Figure 4. Breakdown of cradle-to-gate GHG emissions of a) Pd/NbOPO₄ and b) ZrO₂

Figure 5. Breakdown of cradle-to-gate fossil fuel consumption of a) Pd/NbOPO₄ and b) ZrO₂

Although the primary material component of the Pd/NbOPO₄ catalyst by mass is the NbOPO₄ support, the Pd metal is by far the major contributor to cradle-to-gate GHG emissions and fossil fuel consumption—despite accounting for less than 5% of the catalyst’s material inputs. This is because Pd is far more carbon-intensive than any of the other materials that are used, and its own cradle-to-gate GHG emissions and fossil fuel consumption are much higher than any other material that contributes to the Pd/NbOPO₄ catalyst, as we observed in Table 8. It is notable that the primary source of purchased energy for the production of Pd by PGM mining is electricity, contributing to 71% of the total energy consumption of Pd production, as shown in Table 4. In South Africa, 88% of electricity is sourced from coal (Kingsbury and Benavides 2021) which is also a major source of GHG emissions, leading to a high fossil fuel and emissions impact for Pd.

For the ZrO₂ catalyst, the NaOH material input is the primary contributor to cradle-to-gate GHG emissions and fossil fuel consumption despite contributing less mass than zircon to the total material inputs in catalyst production. As discussed in Section 3.1, the zircon content in the heavy-mineral sands from which it is produced is very high, so its production is minimally resource-intensive in comparison to other mining and beneficiation processes. Material and energy flows for NaOH are already detailed in GREET, so we do not describe those inputs in this study, but we note that the cradle-to-gate fossil fuel consumption of NaOH as reported in GREET (28.5 MJ/kg NaOH) is much higher than that of zircon (2.6 MJ/kg zircon), and that 73% of the fossil fuel consumption for NaOH production is from natural gas (GREET 2020).

We also examine the contributions of the various material and energy inputs of both Pd/NbOPO₄ and ZrO₂ to the cradle-to-gate water consumption in Figure 6. Process water consumption for the synthesis of both catalysts is very minimal, as is water consumed by transportation and upstream water consumption from energy inputs. The majority of water consumption is from the upstream production of the major material inputs: Pd and NbOPO₄ for Pd/NbOPO₄, and zircon and NaOH for ZrO₂. The contributions of Pd and NbOPO₄ to the total water consumption of the Pd/NbOPO₄ catalyst are nearly equivalent, while zircon has a larger impact than NaOH on water consumption for ZrO₂.

Figure 6. Breakdown of cradle-to-gate water consumption of a) Pd/NbOPO₄ and b) ZrO₂

The sizeable impact of zircon on the cradle-to-gate water consumption of ZrO₂ can be attributed to the wet concentration process used during heavy-mineral sands mining to increase the zircon content of the sands after mining. This explains why the ZrO₂ catalyst has a cradle-to-gate water consumption similar to other catalysts in GREET despite having lower values for cradle-to-gate GHG emissions and fossil fuel consumption. As with zircon, the concentration process for palladium production uses large amounts of water when creating a slurry to separate less valuable materials from the PGMs. A sizeable water consumption attributed to the NbOPO₄ support is also expected, both because the support makes up 90% of the catalyst's material inputs and because the sol-gel synthesis of NbOPO₄ from KNbO₃ and H₃PO₄ requires a large amount of water for use as a reaction solvent. The combined impact of Pd and NbOPO₄ causes the cradle-to-gate water consumption for this catalyst to be higher than that of the other catalysts in the GREET model. However, we note that the water consumed during production of the NbOPO₄ catalyst support and its KNbO₃ precursor is estimated based on stoichiometric assumptions from laboratory processes, which unlike industrial processes are not designed to minimize water consumption or to incorporate water recirculation. The cradle-to-gate water consumption of the Pd/NbOPO₄ catalyst is thus an overestimation, and there is potential to reduce water consumption if these processes were industrially optimized or if water recirculation was considered.

4. CONCLUSION

In this study, the supply chains of a Pd/NbOPO₄ catalyst and a ZrO₂ catalyst were identified, and LCI data was collected for each stage in the catalysts' supply chains. Material and energy flows for many materials used in the production of the Pd/NbOPO₄ and ZrO₂ catalysts are already detailed in GREET, while other materials such as Nb₂O₅, KNbO₃, NbOPO₄, Pd, and ZrSiO₄ required the compilation of new LCI data and analysis. To procure material and energy inputs for these previously undefined materials, we consulted a combination of scientific literature, technical reports, sustainability reports from mining and chemical industries, and direct correspondence with professionals in the field. Material and energy flows for the new catalysts and associated materials were implemented in GREET, and cradle-to-gate LCAs were performed. We reported cradle-to-gate values for GHG emissions, fossil fuel consumption, and water consumption for the new materials, and we examined the various factors that contribute to the environmental burden of the Pd/NbOPO₄ and ZrO₂ catalysts.

The cradle-to-gate GHG emissions and fossil fuel consumption of the Pd/NbOPO₄ catalyst are very similar to results for existing catalysts in GREET, such as Pt/γ-Al₂O₃, MoCo/γ-Al₂O₃, and ZSM-5. The primary contributing factor to emissions and fossil fuel consumption for Pd/NbOPO₄ is the mining and processing of the PGMs to produce Pd. The initial Pd content in PGM ore is extremely low, at less than 0.0002%, so very intensive processes of open cut mining, wet concentration, high-temperature smelting, and both physical and chemical refining are required. In addition, the main energy input for PGM production is electricity, which is almost entirely sourced from coal in South Africa, where PGMs are produced. However, the large environmental burden of Pd does not necessarily indicate that alternative metals need to be considered. PGMs have valuable catalytic properties, and their high recyclability rate (98%-99%) should be considered when putting their environmental burden in context. Although data for the secondary production of Pd was not available at this time, it would be very beneficial to incorporate recycling as a pathway for Pd production in the catalyst's LCA. There are likely to be more opportunities for reducing GHG emissions and fossil fuel consumption of Pd/NbOPO₄ by targeting the various processes used to synthesize the NbOPO₄ support, particularly those derived from laboratory procedures that are not optimized for industrial performance.

Cradle-to-gate GHG emissions and fossil fuel consumption for the ZrO₂ catalyst are smaller than those of the Pd/NbOPO₄ catalyst and other catalysts in GREET. This can be attributed to the facts that the ZrO₂ catalyst supply chain requires fewer production steps and that the mining and beneficiation processes for zircon are minimally intensive due to the very high initial zircon content (10%-20%) in heavy-mineral sands. It is the upstream production of NaOH, not zircon, that is the primary driver of the energy and emissions burden of ZrO₂. The environmental burden of NaOH production is mainly due to the heavy use of natural gas. While the cradle-to-gate GHG emissions and fossil fuel consumption results for the ZrO₂ catalyst are already low when compared to the other catalysts, a reduction in NaOH consumption during ZrO₂ production could be an effective option for minimizing the environmental burden of this catalyst. However, this may be difficult to achieve as excess NaOH is required for the caustic

fusion of zircon. It could be beneficial to examine potential recycling of NaOH in order to address this issue.

Both catalysts require water-intensive processes during their production. The cradle-to-gate water consumption of the Pd/NbOPO₄ catalyst is higher than that of the other catalysts in GREET that have comparable emissions and energy consumption results. Since one of the catalysts that we used for comparison is Pt/ γ -Al₂O₃, which sources platinum (Pt) from the same PGM mining process as Pd, we can attribute the higher water consumption of the Pd/NbOPO₄ catalyst to the contribution of the NbOPO₄ support. Data on industrial-scale production of NbOPO₄ was unavailable, so we consulted several laboratory-scale synthetic procedures when building this inventory. Many of the chemical reactions from these laboratory procedures are conducted in water or water-based solvents, and we utilized the volume of water that the authors of these procedures report as necessary for the synthesis of a given mass of product. However, these processes are not optimized to minimize water consumption as industrial processes would be. The water consumption of NbOPO₄ is thus an overestimate, and it would be beneficial to examine whether the reactions involved in its production can be performed using less water. For the ZrO₂ catalyst, the main contributor to water consumption is zircon, the production of which utilizes wet gravity separation methods to separate heavy-mineral sands from less valuable mining products. Although less impactful than zircon, NaOH also contributes to the water consumption of ZrO₂, so reduction of NaOH could also help reduce water consumption.

The results of this study may be used to inform the selection of materials for the design of other catalysts or to make targeted adjustments to various stages of Pd/NbOPO₄ and ZrO₂ production in order to minimize environmental impacts. The system boundary of this study is cradle-to-gate, so there are more conclusions to be drawn about the ultimate environmental burden of the catalysts by accounting for end-of-life recycling and performing a cradle-to-grave analysis. In addition, the environmental burden of Pd/NbOPO₄ and ZrO₂ may be properly contextualized by incorporating these catalysts into LCAs of the related biofuels. Both Pd/NbOPO₄ and ZrO₂ have been successfully used in the production of the ether diesel bioblendstock 4-BH, so an LCA of this biofuel that includes these catalysts would be beneficial. ZrO₂ and niobic acid, a hydrated form of Nb₂O₅, have also been used in the production of diesel and jet fuel range blendstocks that merit LCAs. Due to the heavy environmental burden of producing metals, catalyst production is typically intensive, but oftentimes only a very small amount of catalyst is required for the production of a large amount of biofuel. By including catalysts in a biofuel's LCA, we can evaluate the catalyst's environmental impact in the proper context as well as obtain more accurate estimations of the life-cycle energy consumption and GHG emissions of the biofuel.

APPENDIX A: Calculation of Energy Consumption in KNbO₃ Production

The total energy input for the production of KNbO₃ is described by Equation 2. It includes three terms that require prior determination: $q_{solvent}$, q_{CSTR} , and q_{stir} . The value of $q_{solvent}$ corresponds to the heat required to raise the temperature of the solvent to the desired reaction conditions. In this particular reaction, an 8 M KOH aqueous solution is heated from room temperature (25°C) to 200°C. We assume ideal thermodynamic heating of the KOH solution and use Equation A-1 to calculate $q_{solvent}$.

$$q_{solvent} = mC_p\Delta T \quad (A-1)$$

where:

m is the mass of solution used to produce 1 ton of KNbO₃ (25,174 kg);

C_p is the average specific heat capacity of an 8 M KOH solution (3387 J/kg K); and

ΔT is the change in temperature (175 K).

The value of q_{CSTR} corresponds to the heat loss of the CSTR, and it is calculated using Equation A-2 according to the methodology of Majeau-Bettez et al. (2011). The parameters λ , A , T_0 , and x are specific to the reactor and are provided by Majeau-Bettez et al. (2011), while t_r and T_r are specific to the reaction and are sourced from the synthesis procedure of Lu et al. (1998).

$$q_{CSTR} = \frac{\lambda A (T_r - T_0)t_r}{x} \quad (A-2)$$

where:

λ is thermal conductivity specific to the reactor (0.04 W/mK);

A is the surface area of the reactor (25 m²);

t_r is the reaction time (2 hours);

T_r is the temperature of the reactor (200°C);

T_0 is the temperature of the reactor wall (50°C); and

x is the thickness of the reactor insulation (0.1 m).

In this scenario, the reactants are stirred in the CSTR, so the value of q_{stir} depends on how much electricity the specific CSTR uses to agitate the reactants. Majeau-Bettez et al. (2011) report an electricity consumption of 20.92 MJ/hr for stirring the contents of the 10,000 L reactor, which must be stirred for 2 hours.

APPENDIX B: Calculation of Energy Consumption in NbOPO₄ Production

The total energy input for the production of NbOPO₄ is described by Equation 4 above. It includes seven terms: $q_{solvent}$, q_{CSTR} , q_{dry} , $q_{calcine}$, q_{oven} , $q_{furnace}$, and q_{stir} . Using the same approach as for the production of KNbO₃ in Appendix A, the value of $q_{solvent}$ is again calculated using Equation A-1. In the case of NbOPO₄ production, a 0.4 M H₃OPO₄ solution is heated incrementally from room temperature (approximately 25°C) to 40°C, to 80°C, and ultimately to 160°C. The input values for m , C_p , and ΔT in Equation A-1 are adjusted to correspond to the mass of H₃PO₄ solution used to produce 1 ton of NbOPO₄ (55,568 kg), the average specific heat capacity of a 0.4 M H₃PO₄ solution (4148 J/kg K), and the total change in temperature (135 K), respectively. The value of q_{CSTR} is calculated using Equation A-2, with the reaction-specific values for t_r and T_r being adjusted to the total heating time for NbOPO₄ production (27 hours) and the temperature of the reactor (160°C), respectively.

The values of q_{dry} and $q_{calcine}$ are determined in the same manner as $q_{solvent}$, using Equation A-1, assuming ideal thermodynamic heating of solid NbOPO₄ from room temperature to the desired final temperature for either drying (110°C) or calcining (400°C). The specific heat capacity of NbOPO₄ is estimated as 418 J/kg K according to the Neumann-Kopp rule, which states that the specific heat capacity of a compound may be approximated as a weighted sum of the heat capacities of its elemental constituents (Atkins and Escudier, 2019). Both q_{dry} and $q_{calcine}$ are calculated for the heating of 1 ton, or 907.185 kg, of solid NbOPO₄ product.

The energy used to heat the oven to 110°C for the drying step (q_{oven}) is calculated using Equation B-1 according to the methodology of Majeau-Bettez et al. (2011) and Dunn et al. (2015). The value for the heating duration t is provided by the synthesis procedure of He et al. (2020), while the value for power consumption P is estimated using data for a Nabertherm chamber oven (Nabertherm 2016). Majeau-Bettez et al. (2011) assume in their calculations that an oven does not require the electricity needed to reach its maximum heating power if it is not being heated to its maximum temperature.

We therefore apply an estimation method described by Tschernitz (2001) in which the operating power of the oven is approximated by the ratio of the operating temperature to the maximum temperature taken to the third power and multiplied by the maximum power consumption of the oven. The volume of the oven is 1,000 L, which we use to scale q_{oven} to the electricity that the oven requires for the drying of 1 ton of NbOPO₄.

$$q_{oven} = P \times t \quad (B-1)$$

where:

P is the operating power of the oven at 110°C (0.68 kW); and
 t is the duration of heating in the oven (12 hours).

The value for $q_{furnace}$ is calculated using the same method as q_{oven} in Equation B-1, with P being adjusted to the operating power of a Nabertherm chamber furnace at 400°C (12.5 kW) and t being adjusted to the duration of heating in the furnace during the calcining step (3 hours). The volume of the furnace, 8,300 L, is used to scale the electricity consumption to 1 ton of NbOPO₄. Finally, q_{stir} is again determined based on an electricity consumption of 20.92 MJ/hr for stirring the contents of the 10,000 L continuous-stir tank reactor (Majeau-Bettez et al. 2011) for 3 hours.

APPENDIX C: Calculation of Energy Consumption in Pd/NbOPO₄ Production

The total energy input for the production of Pd/NbOPO₄ is described by Equation 5. It includes seven terms: q_{dry_1} , q_{dry_2} , q_{reduce} , q_{oven_1} , q_{oven_2} , q_{tube} , and q_{stir} . The values for q_{dry_1} , q_{dry_2} , and q_{reduce} are all calculated using Equation A-1 assuming ideal thermodynamic heating of solid Pd/NbOPO₄. For all three calculations, the input for mass m is 1 ton, or 907.185 kg, and the input for the average specific heat C_p of Pd/NbOPO₄ is 409 J/kg K. The change in temperature ΔT is 82 K, 240 K, and 240 K for the first drying step, the second drying step, and the reducing step, respectively.

The values of q_{oven_1} , q_{oven_2} , and q_{tube} are all calculated with Equation B-1 using the same method of estimating power consumption based on the operating temperature. We use a Nabertherm oven with a volume of 1,050 L to model drying steps at 110°C and 265°C (Nabertherm 2017). The first drying step has an operating power consumption P of 0.21 kW and a duration t of 6 hours. The second drying step has an operating power consumption P of 3.2 kW and a duration t of 2 hours. We use a Nabertherm tube furnace with a loading volume of 0.8 L to model the reduction of Pd/NbOPO₄ at 265°C (Nabertherm 2017). Reduction in the tube furnace requires an operating power consumption P of 0.065 kW and a duration t of 5 hours. The volumes of the oven and tube furnace are used to scale the results to the production of 1 ton of Pd/NbOPO₄.

The reactants used to form Pd/NbOPO₄ are not stirred in an autoclave or CSTR, as they are in the production of KNbO₃ and NbOPO₄, so we model the agitation of the reactants using a chemical mixer that can agitate catalyst slurries up to 1,892L (Madden Pump 2020). The mixer operates at 0.33 horsepower when stirring its maximum volume of contents, and the duration of stirring is 12 hours. Again, the energy consumption is scaled to the volume of reactants that must be stirred to produce 1 ton of Pd/NbOPO₄.

APPENDIX D: Inputs and Results using Market Allocation for Pd Production

In Table D-1, we provide inputs for palladium production that were calculated by Kingsbury and Benavides (2021) using a market allocation approach. These inputs serve as an alternative to the values in Table 4 that were calculated using mass allocation, which is the default allocation method in GREET (GREET 2020). The total energy input of Pd production by market allocation is more than two orders of magnitude greater than the energy input by mass allocation. This is primarily due to the high market price of Pd, which is second only to Pt when compared to the market prices of the other outputs of PGM production. Specific values for the variables used in the market allocation calculation, including the market prices and production masses of the PGMs, can be found in the report by Kingsbury and Benavides (2021).

Table D-1. Market allocation material and energy inputs for the production of palladium

Material	Amount per ton Pd	
Water	171,176 gal	
Energy	mmBtu/ton Pd	Share %
Electricity	90,370	71%
Diesel	14,571	11%
Coal	21,292	17%
Natural gas	1,292	1%
Total energy input	127,525	

In Table D-2, we provide the results of the cradle-to-gate LCA for both Pd and the Pd/NbOPO₄ catalyst using the market allocation inputs for Pd production from Table D-1.

Table D-2. Cradle-to-gate LCA results for Pd and Pd/NbOPO₄ using market allocation inputs

Cradle-to-gate result	Pd	Pd/NbOPO₄ catalyst
GHG emissions (kg CO ₂ e/kg material)	32,223	1,614
Fossil fuel consumption (MJ/kg material)	330,918	16,582
Water consumption (gal/kg material)	16,786	854

When compared to the cradle-to-gate LCA results for Pd and Pd/NbOPO₄ shown in Table 8 that used the mass allocation inputs for Pd production, the results shown in Table D-2 are significantly higher for both Pd and Pd/NbOPO₄. Notably, the cradle-to-gate GHG emissions and fossil fuel consumption for the Pd/NbOPO₄ catalyst are both more than two orders of magnitude

larger when using the market allocation inputs compared to the mass allocation inputs, and the cradle-to-gate water consumption is more than one order of magnitude larger. This causes Pd to have a much larger percent contribution to the cradle-to-gate GHG emissions, fossil fuel consumption, and water consumption of the Pd/NbOPO₄ catalyst since the environmental burden of Pd increases while the other components of the catalyst remain the same. When using the mass allocation inputs for Pd production, Pd accounts for 71% of the GHG emissions, 63% of the fossil fuel consumption, and 47% of the water consumption of the Pd/NbOPO₄ catalyst. When using the market allocation inputs, Pd accounts for over 99% of the GHG emissions and fossil fuel consumption of the Pd/NbOPO₄ catalyst and for over 98% of the catalyst's water consumption. Although Pd makes up only a small amount of the Pd/NbOPO₄ catalyst by mass, the choice of allocation method for Pd production has a significant effect on the catalyst's LCA results.

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Energy Systems Division

Argonne National Laboratory

9700 South Cass Avenue

Argonne, IL 60439

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