



# Screening LCA for CCU routes connected to CO<sub>2</sub> Smart Grid



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# Screening LCA for CCU routes connected to CO<sub>2</sub> Smart Grid

This report is prepared by:

Harry Croezen  
Sanne Nusselder  
Erik Roos Lindgreen  
Diederik Jaspers

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Further information on this study can be obtained from the contact person Diederik Jaspers (CE Delft)

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# Summary

The Paris agreement requires significant steps in order to achieve reduction of greenhouse gases of 90-95% CO<sub>2</sub> eq. compared to 1990 levels. One of the possible steps is the application of novel technologies like the reuse of CO<sub>2</sub> in a value chain. This is attractive if can cover the cost of the capture of CO<sub>2</sub>, while the ETS-price level is still very low. In the two Dutch provinces of North- and South-Holland, a consortium of more than 20 public and private parties is launching an initiative (CO<sub>2</sub> Smart Grid) aimed at utilizing CO<sub>2</sub> as a raw material for a circular economy (Carbon Capture and Utilization, or shortly CCU). To this end, a network will be developed in which CO<sub>2</sub> from different sources can be made available to different users. The proposed backbone of this network is the existing OCAP CO<sub>2</sub> pipeline, which already provides CO<sub>2</sub> from Shell in Pernis and ethanol producer Alco in Pernis to the horticulture sector in the Westland region for growth promotion of crops.

This study focusses on the Life Cycle Assessment (LCA) of different CCU routes applicable in the CO<sub>2</sub> Smart Grid. The results of this study can serve as input for a to-be conducted Social Cost Benefit Analysis.

## CCU routes

This study compares the environmental impact of nine different CCU routes on the basis of '1 tonne of CO<sub>2</sub> captured in 2030 and subsequent utilization'. The nine routes are a combination of CO<sub>2</sub> capture options from three different sources and utilization of the CO<sub>2</sub> in three different applications.

Table 1 gives an overview of the CCU routes considered in this LCA. Furthermore these nine different CCU routes are compared with Carbon Capture and Storage (CCS) as a reference.

Table 1 - CCU routes

CO <sub>2</sub> source   Utilization	Horticulture	Mineralisation	Methanol production
Waste incineration	Route 1	Route 2	Route 3
Blast furnace process and blast furnace gas	Route 4	Route 5	Route 6
Fossil oil refining	Route 7	Route 8	Route 9

Of course there are many more CCU routes possible in the Netherlands, but this study has been limited to nine different routes which are considered relevant for the region of North-Holland (Tata Steel) and South-Holland (Rotterdam Harbour Industrial Complex).

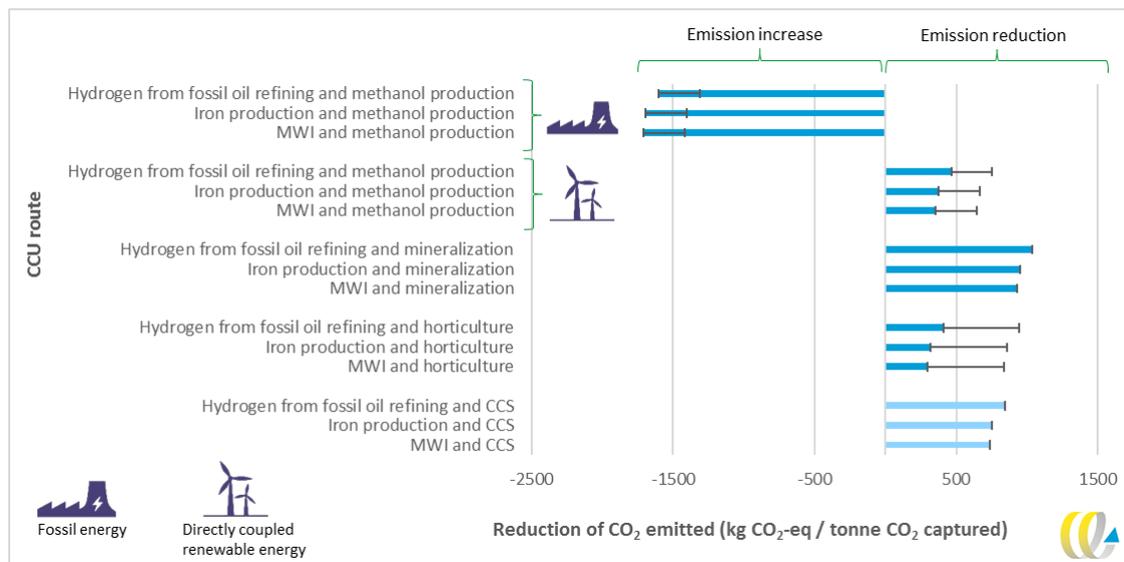


# Climate change impact of CCU routes

## Reduction of climate change impact

All of the routes considered lead to a reduction of climate change impact compared to non-capture, as can be seen in Figure 1.

Figure 1 - Reduction of climate change impact per CCU route in comparison to non-capture



Note: The black bar indicates CO<sub>2</sub> emission reduction due to different situations. For methanol production use as fuel (lowest reduction) and use as chemical where CO<sub>2</sub> is stored for more than 100 years (highest reduction). For utilization in horticulture addition of captured CO<sub>2</sub> in the current OCAP pipeline (lowest reduction) and the addition of CO<sub>2</sub> during the summer (highest reduction).

The extent to which this is the case is dependent on:

- the duration of carbon storage in the produced products (e.g. permanent storage in case of mineralisation of CO<sub>2</sub> in mineral construction materials);
- the quantity of energy used by the capture technology;
- the quantity of energy used by the utilization technology;
- the carbon footprint of the product that is replaced (e.g. avoidance of natural gas burner to supply Dutch horticulture with CO<sub>2</sub> for increased plant growth).

## Utilization in mineralisation

Utilization of CO<sub>2</sub> for mineralisation, the production of one type of mineral material (compensatiesteen), leads to net avoided CO<sub>2</sub> emissions of around 1 tonne of CO<sub>2</sub> per tonne of CO<sub>2</sub> captured. Despite the carbon footprint of the capture technologies, the produced Compensatiesteen avoids production of conventional sand-lime brick. When the capture technologies have a lower carbon footprint (e.g. when in the future renewable electricity mix is used), utilization in mineralisation could even lead to net negative CO<sub>2</sub> emissions. This means that more CO<sub>2</sub> emission is prevented than CO<sub>2</sub> captured.

## Utilization in horticulture

The utilization of CO<sub>2</sub> in horticulture leads to net avoided CO<sub>2</sub> emissions of between 300 and 950 kg CO<sub>2</sub> per tonne of CO<sub>2</sub> captured. The lowest net avoided CO<sub>2</sub> emission is as currently reported for captured CO<sub>2</sub> added to the OCAP-pipeline while the highest reduction occurs when CO<sub>2</sub> is added to a greenhouse during summer. The net avoided CO<sub>2</sub> emission is caused by the avoided use of natural gas for the production of CO<sub>2</sub> in horticulture. This conclusion is valid until the horticulture sector made a transition to a renewable heat source (e.g. geothermal heat).

The possible sources for CO<sub>2</sub> supply in horticulture (the reference) in towards 2040-2050 is unknown because the future benchmark for heat supply in greenhouses has yet to be determined. A possibility is the use of biomass in CHP for both heat and CO<sub>2</sub> production, but also geothermal heat supply without associated CO<sub>2</sub> emissions is an option. The geothermal scenario would fully depend on an external source of CO<sub>2</sub>, which can be delivered by the CO<sub>2</sub> Smart Grid. Whether or not the application of captured CO<sub>2</sub> aids the shift towards renewable energy and what would be the appropriate reference CO<sub>2</sub> source to consider in the future is a topic that needs further discussion. Therefore the exact carbon footprint reduction after a switch to a fully renewable heat source in the horticulture sector is uncertain and depends on the outcome of different scenario's.

## Utilization in methanol production

Utilization in methanol production will lead to net avoided CO<sub>2</sub> emissions when 100% renewable energy is used for methanol and hydrogen production. If fossil-based electricity is used in the process, more CO<sub>2</sub> is emitted than captured. The net avoided CO<sub>2</sub> emissions will increase when the CO<sub>2</sub> is used in durable products. 'Durable' in this context implies that CO<sub>2</sub> is sequestered for more than 100 years. In that case, this utilization method could reach net avoided CO<sub>2</sub> emissions of around 700 kg CO<sub>2</sub> per tonne of CO<sub>2</sub> captured. This is comparable to CCS (see Figure 1). A lot of renewable electricity is required to produce hydrogen for methanol production on a large scale. We assume additional renewable electricity supply (e.g. directly linked windfarms), ample availability of this renewable electricity for producing hydrogen, and that the use of this electricity does not compete with utilization in applications leading to lower net CO<sub>2</sub> emissions.

It must be noted that methanol production is not the only possible application of CO<sub>2</sub> in the chemical industry. The reason that methanol was selected is, apart from the availability of data from the demonstration plant in Iceland, that it is a so called platform chemical with a wide range of products which are currently based on fossil oil and gas. Other possible CO<sub>2</sub> utilization routes in the chemical industry include the production of polyols for the production of polyurethanes. Conclusions drawn on methanol production should thus not be seen as exemplary for CO<sub>2</sub> utilization in the chemical industry.

## Other environmental benefits

For several reasons, no conclusions could be drawn on other environmental impacts:

- additional benefits caused by the additional cleaning of CO<sub>2</sub> containing (flue) gas during the capture process are unknown;
- emissions from degradation of absorbents are unknown.



## Interpretation of the conclusions

The orders of magnitude of CCS and CCU applicability in 2030 are expected to be incomparable. E.g. the potential storage by means of CCS is expected to be much higher than the potential for use of CO<sub>2</sub> in mineralization in the Netherlands. Results must therefore only be seen on a per tonne basis and cannot be extrapolated. The spatial application of the technologies also differ, e.g. CCS can be applied the whole year round while the peak of CO<sub>2</sub> utilization in horticulture is during the growing season and less so in winter.

Because the study carried out is a screening LCA, the drawn conclusions should be seen as indicative figures; they offer an order of magnitude estimation and cannot be seen as representative for individual (industrial) plants present in the Netherlands. Furthermore the results are not appropriate for national carbon accounting. This means that when calculating the emissions of the Netherlands as a whole the presented reduction in CO<sub>2</sub> emissions cannot be taken into consideration. The same holds for using the outcomes for corporate carbon accounting practices.

To make the results applicable to individual CCU routes e.g. CO<sub>2</sub> capture at the AEB MWI in Amsterdam and application of the CO<sub>2</sub> in horticulture in Aalsmeer, a full scale LCA study will need to be conducted based on the actual variables chosen for the specific installations.



# 1 Introduction

In the two Dutch provinces of North- and South-Holland, a consortium of more than twenty public and private parties is launching an initiative - CO<sub>2</sub> Smart Grid - aimed at utilizing CO<sub>2</sub> as a raw material for a circular economy (Carbon Capture and Utilization, or shortly CCU). To this end, a network will be developed in which CO<sub>2</sub> from different sources can be made available to different users. The proposed backbone of this network is the existing OCAP CO<sub>2</sub> pipeline, which already provides CO<sub>2</sub> from Shell in Pernis and ethanol producer Alco in Pernis to the horticulture in the Westland region for growth promotion of crops.

Ecofys has conducted a pre-feasibility study in which they identified in which applications CO<sub>2</sub> could be utilized in North- and South-Holland in the short term (5-10 years) (Ecofys, 2017) Table 2 shows the results of the pre-feasibility study. In this pre-feasibility study, Ecofys did not analyse the source of the used CO<sub>2</sub>.

**Table 2 - Overview of identified prospective utilization application of CO<sub>2</sub> as raw material**

CCU technology	TRL	Current (2017) (kt CO <sub>2</sub> )	Near term (5 years) (kt CO <sub>2</sub> )	Long term (10 years) (ktCO <sub>2</sub> )
Horticulture	9	400-500	850-1,000	1,200
Carbonate mineralization	4-8	0	100-200	100-300
Polymer processing	8	-	12-23	30-45
Concrete curing	7-8	-	-	30
Synthetic methanol (including methane)	8	-	-	220
Methanol yield boosting	9	630	900	1,250
<b>Rounded total</b>		<b>~400</b>	<b>~1,000</b>	<b>~1,700</b>

Source: Table from (Ecofys, 2017). 'Methanol yield boosting' is specifically related to methanol production at BIOMCN in Delfzijl.

According to studies of Ecofys and CE Delft the various capture and application routes are not profitable under current market conditions. The various capture and application routes could have a social advantage, in particular because they could lead to a CO<sub>2</sub> emission reduction, and application might therefore provide a benefit from a societal perspective. This potential benefit can be made explicit by means of a Social Cost Benefit Analysis (SCBA). The basis of such a SCBA is a Life Cycle Analysis (LCA) in which environmental impacts are quantified. The LCA is commissioned by the Ministry of Infrastructure & Water Affairs, the MKBA will be commissioned by BLOC, both on behalf of the CO<sub>2</sub> Smart Grid consortium.

This study focusses on the LCA of different CCU routes applicable in the CO<sub>2</sub> Smart Grid. The results of this study can serve as input for the later SCBA. The study is conducted under supervision of the client, process supervisor BLOC and the core working group of the consortium.



## 2 Methodology

The Life Cycle Assessment (LCA) methodology is used to determine the environmental impact of a product or service throughout the entire life cycle. It can be used to compare the environmental impact of different products or services. Because the different CCU routes do not provide the same product, although CO<sub>2</sub> is captured in all CCU routes, a substitution approach is used. See Section 2.3.

The reporting methodology for LCA is set by the ISO14040 and ISO14044 guidelines for Life Cycle Assessment. The main lines of these methodological guidelines are followed with the important note that this study is a screening LCA, and not a full scale Life Cycle Assessment. A screening LCA aims to give an indication of the comparative environmental impact and recognizes the uncertainties because of the short span in which this study is carried out.

A number of important methodological choices are described in this chapter.

### 2.1 Goal and scope definition

#### 2.1.1 Goal of the study

The main goal of the study is to identify the environmental hotspots in the different CCU routes, and make a comparison of the different routes.

The main goal is reached by:

- examining the net avoided CO<sub>2</sub> emission for the CCU routes;
- examining the implication of the different storage times during which the CO<sub>2</sub> is utilized in the intended applications;
- examining possible other environmental impacts of the CCU routes.

#### 2.1.2 Scope of the study

In order to make a comparison, a unit of comparison needs to be defined. This unit of comparison is called the **functional unit**. The functional unit is defined as:

1 tonne of CO<sub>2</sub> captured in 2030 and subsequent utilization.

Different utilization-routes produce different products/services. CCU is a multifunctional system generating both the service of capturing of CO<sub>2</sub> as well as utilizing the CO<sub>2</sub> in a product/service. Since the aim of this study is to provide insight into the environmental benefit of the entire CCU process, and not into a single product, the functional unit has been set in such a way that it follows one tonne of captured CO<sub>2</sub> through the entire process.

There are many CCU routes possible in the Netherlands of which nine different CCU routes are compared in this study. These nine routes are based on CO<sub>2</sub> capture options from three different sources and utilization of the CO<sub>2</sub> in three different applications. A selection of routes has been made based on expected availability of CO<sub>2</sub> in 2030, technology readiness level and the compatibility with current industry. Table 3 gives an overview of the nine different CCU routes that are considered in this LCA. Furthermore these nine different CCU routes are compared with Carbon Capture and Storage (CCS) as a reference.



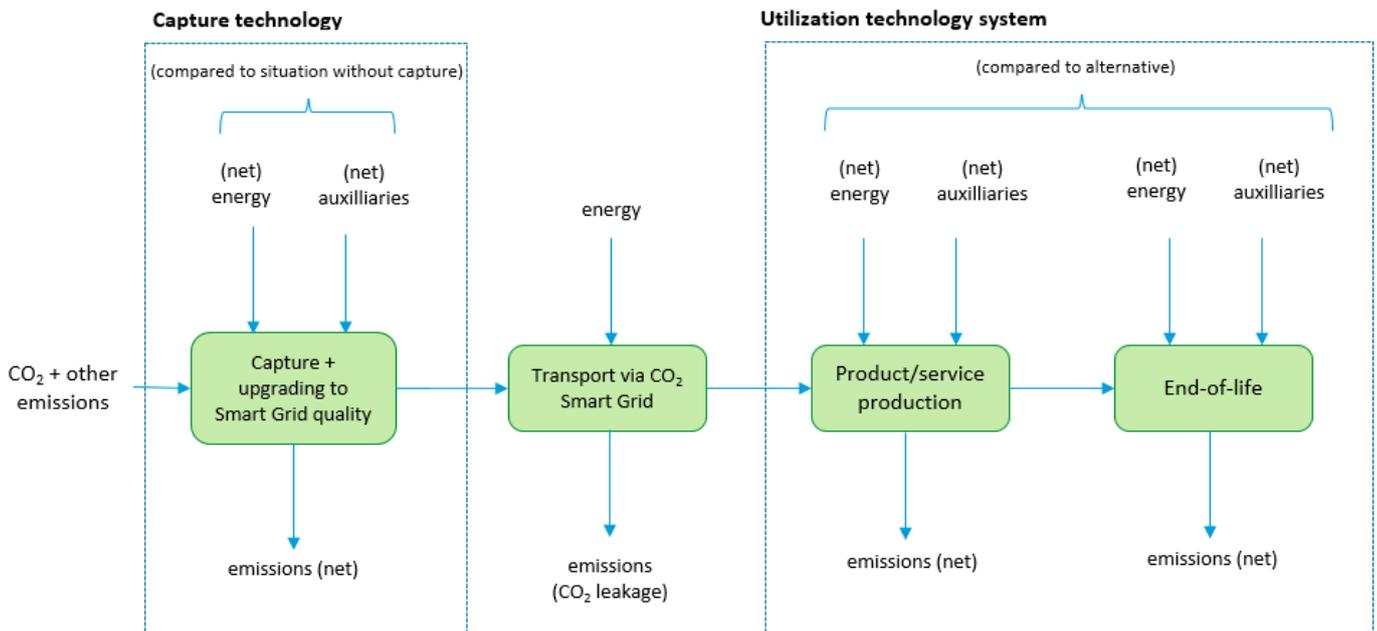
**Table 3 - CCU routes**

CO <sub>2</sub> source   Utilization	Horticulture	Mineralisation	Methanol production
Waste incineration	Route 1	Route 2	Route 3
Blast furnace process and blast furnace gas	Route 4	Route 5	Route 6
Fossil oil refining	Route 7	Route 8	Route 9

This study compares the fulfilment of the functional unit of these nine CCU routes within the **system boundaries** as shown in Figure 2. All green filled boxes represent life cycle phases that are taken into consideration in this study.

For every life cycle phase, material and energy use are taken into consideration as well as all environmental emissions relevant to the environmental impact categories considered in this study (see Section 2.2). Figure 2 shows the general system boundaries. The exact capture technology and utilization system differ per CCU route. The product or service produced because of the utilization of CO<sub>2</sub> also differs per utilization method. For the system description per CCU route see Chapter 3. The CO<sub>2</sub> source is outside of the system boundaries this means that e.g. the production of iron is considered to occur whether or not CCU is applied. The systems of iron production and CCU are therefore seen as two different production systems.

**Figure 2 - System boundaries of CCU**



Note: All life cycle phases with a green filling are taken into consideration in this study, including energy and auxiliary use as well as emissions.



## 2.2 Environmental impact categories

This study uses the ReCiPe2016 methodology to examine the environmental impact of the different CCU routes<sup>1</sup>. The ReCiPe2016 Midpoint Hierarchist Approach (v.1) has been chosen as it is included in the SimaPro Software (v.8.4). A wide range of different environmental impacts are included in the ReCiPe-methodology and can be studied with LCA. Within the limited time frame of the study only global warming potential (CO<sub>2</sub> eq. emissions) are quantified. Qualitative statements will be made on other relevant environmental impacts such as fine particulate matter formation and acidification.

## 2.3 Dealing with a multifunctional system

As described earlier the different utilization-routes produce different products/services. CCU is a multifunctional system generating both the service of capturing of CO<sub>2</sub> as well as utilizing the CO<sub>2</sub> in one or multiple products.

The choice of functional unit leaves us with the issue of how to show the benefit of the produced product/service per utilization method. According to ISO14044, there are different approaches if a system under study has multiple functions. The preferred approach according to ISO14044 is to prevent needing to allocate environmental burdens between the different products/services delivered by a CCU route. Allocation of environmental burdens based on economic or physical relationships introduces uncertainties into an LCA study. In the case of the produced product/service we therefore opt for preventing allocation.

Different approaches can be taken to prevent allocation, the most common ones being system expansion and substitution. The different CO<sub>2</sub> utilization routes produce different products/services. Using system expansion would require that all possible products are accounted for in all different options, creating very large systems that make the comparison of the actual CO<sub>2</sub> utilization options complex. We therefore apply substitution by assuming prevention of the currently applied production method for the product or service. The products/services prevented are described per utilization method in Chapter 3.

## 2.4 Fossil and biogenic CO<sub>2</sub>

LCA convention such as e.g. the EN 16760 norm states that to assess climate change impact, all biogenic and fossil CO<sub>2</sub> emissions and removals should be considered. In this study not all life cycle stages are included for the CO<sub>2</sub> sources. For example the biogenic CO<sub>2</sub> uptake (removal) in biogenic products that are eventually treated as waste in an MWI are not taken into consideration in this study.

This means that no comparison can be made of the difference in impact over the entire life cycle of the biobased material and e.g. the fossil-based material in case of the coal-fired power plant. That is also not the purpose of this study. Therefore no environmental distinction is made in this study on the environmental impact of the emission of biogenic and fossil-based CO<sub>2</sub>. In the case future studies are carried out in which the production phase of the CO<sub>2</sub> source is taken into consideration, the emission of the two types of CO<sub>2</sub> is distinguished in the figures and tables in this study.

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<sup>1</sup> For the full methodological report see (Huijbregts, et al., n.d.).



## 2.5 CO<sub>2</sub> storage period

CO<sub>2</sub> is stored for different time periods in the different products considered in this study.

The ILCD-guidelines (JRC European Commission, 2010) state that:

*“temporary carbon storage and the equivalent delayed emissions and delayed reuse/recycling/recovery within the first 100 years from the time of the study shall not be considered quantitatively”.*

We therefore only consider only two different CO<sub>2</sub> storage periods:

- 100 years or less, not leading to CO<sub>2</sub> emission reduction;
- more than 100 years, leading to CO<sub>2</sub> emission reduction.

In reality also a temporary storage of CO<sub>2</sub> (e.g. for 40 years) can have an environmental impact.

Considering those differences is outside of the scope of this study, and not (yet) common in carbon accounting.

## 2.6 Electricity use: changes in energy demand and energy production

Changes in the energy demand and energy production are compensated for by extra production of fossil energy (Agentschap NL, CBS, ECN, PBL, 2012). This method ('de referentiepark-methode') is used in the monitoring and evaluation of energy- and climate policies in the Netherlands. In this study we use this marginal approach to the energy system in line with Dutch convention.

Per year ECN determines a CO<sub>2</sub> emission factor for the exact energy production facilities being used to compensate for the increased energy demand or decreased energy production. ECN has also determined a projection for this CO<sub>2</sub> emission factor for the years 2020, 2023 and 2030.

The CO<sub>2</sub> emission factor is 0.67 kg CO<sub>2</sub>/kWh in each of these years (ECN, 2017). Since this study looks at CCU options in the year 2030, we use this emission factor.

## 3 CCU routes & system boundaries

The CO<sub>2</sub> sources and the technologies used for carbon capture from the three different CO<sub>2</sub> sources are further described in Section 3.1, purification and compression is described in Section 3.2 and the utilization technologies are described in Section 3.3. Combining the three sources with the three utilization technologies leads to nine CCU routes that are examined in this study.

### 3.1 CO<sub>2</sub> sources and carbon capture

Three different industrial processes are considered as source for CO<sub>2</sub> capture:

- waste incineration;
- blast furnace gas from the blast furnace process (iron production);
- fossil oil refining.

These different CO<sub>2</sub> sources were selected based on the expectation that these sources will still be available in 2030 and beyond and because these sources emit significant amounts of CO<sub>2</sub> annually and can hence supply a relevant amount of CO<sub>2</sub> to a CCU grid. Furthermore, these sources are through individual plants already connected to the OCAP infrastructure, which forms the basis of the CO<sub>2</sub> Smart Grid, or can in the near future be connected without large (technological) obstacles. Each of these sources and the technology used to capture the CO<sub>2</sub> are discussed per source below. For each source the carbon capture is assumed to be an addition to the current practice (tailpipe capture of CO<sub>2</sub>) and no more amendments are assumed to be made to the current business of an industrial plant except the accommodation CO<sub>2</sub> capture.

#### 3.1.1 Municipal waste incineration plants (MWI)

The Dutch circular economy policies aim to reduce the quantity of waste being used for energy recovery and instead to increase recycling of waste streams. We expect, however, that considering the speed at which the circular economy is taking shape in the Netherlands, waste incineration still has a role in 2030. Waste incineration plants are therefore considered to be a relevant source for CO<sub>2</sub> capture. Other reasons for their relevance as CO<sub>2</sub> source include:

- Flue gas of MWI is a point CO<sub>2</sub> source.
- There is an incentive for reducing waste production from the circular economy policies and the public opinion of MWI-plants is that they are not as favourable as recycling of material. The application of carbon capture at an MWI will therefore not lead to continued waste incineration when this would not be the case without CO<sub>2</sub> capture (no lock-in is created).
- A part of the CO<sub>2</sub> emissions from MWI are biogenic, since the MWI incinerates biogenic material such as garden and kitchen waste.

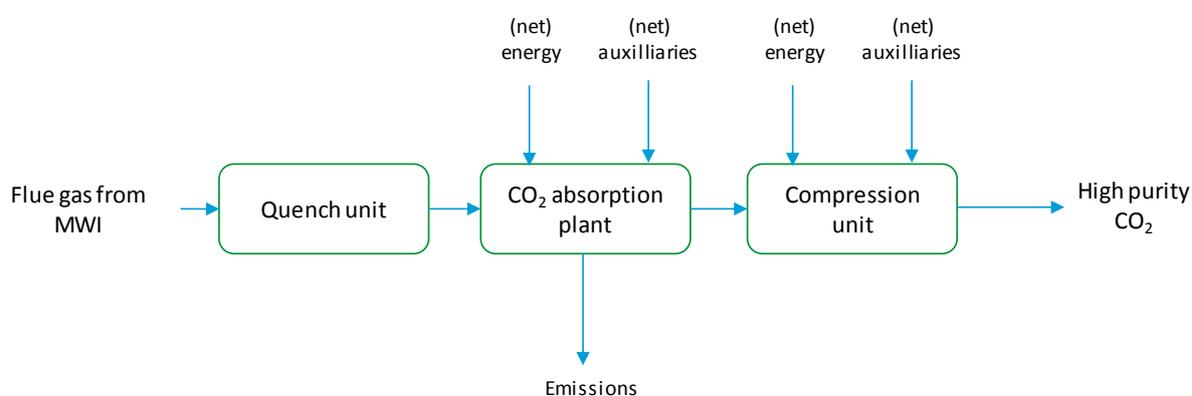
CO<sub>2</sub> emissions at MWIs are assumed to be captured by an innovative absorption technology in a CO<sub>2</sub> absorption plant. This technology has been developed by Procede Gas Treating, and is selected for its high Technology Readiness Level. The technology is currently applied in Delta (British Columbia) and at Twence in the Netherlands.



This technology uses Bilisol as an absorbent. This is a biodegradable solvent developed by Procede with a low degradation rate and very low volatility. A schematic representation of the processes is given in Figure 3. The hot flue gases are cooled to approximately 50°C and cleaned in a quench. CO<sub>2</sub> is next captured by scrubbing the flue gas with a Bilisol solution, after which Bilisol is regenerated in a separate reactor vessel heated with low-pressure steam from the MWI. Recovered high purity CO<sub>2</sub> (≥ 99.95 vol%) is next dehydrated and compressed to the necessary pressure for the CO<sub>2</sub> Smart Grid.

The use of low-pressure steam from the MWI leads to a reduction of the production of electricity. The reduction in electricity production is approximately 0.25 MWe per MW heat extracted<sup>2</sup>. As described in Section 2.6 the reduction in electricity production is compensated by extra production of fossil electricity.

**Figure 3 - Schematic representation of carbon capture at MWIs**



The capture of CO<sub>2</sub> emissions at MWIs is a special case, as the input, and therefore CO<sub>2</sub> emissions, are partly of biogenic origin. The biogenic content of the waste incinerated at MWIs is approximately 64%.<sup>3</sup> As described in Section 2.4, we present the biogenic CO<sub>2</sub> emissions but do not make a distinction between the environmental impact of biogenic and fossil-based CO<sub>2</sub> emissions.

For the LCI of this capture technology see Annex A.

### 3.1.2 Blast furnace gas from the blast furnace process

Even in a circular economy there will be demand for primary, ore based high-grade flat steel as used in e.g. car manufacturing, due to losses and downgrading in quality of materials. Such high-grade steel can only be produced by way of the blast furnace production route for iron, as utilized at e.g. Tata IJmuiden. Tata IJmuiden is globally one of the most technologically advanced producers of such high-grade steel and is also one of the few producers operating competitively (Tata Steel, 2016) in a market plagued by overcapacity. It would hence be likely that Tata IJmuiden is still operational in 2030 and beyond. Based on this perspective, CO<sub>2</sub> capture from blast furnace gas at Tata IJmuiden is proposed as one of the options as feedstock for the CO<sub>2</sub> Smart Grid.

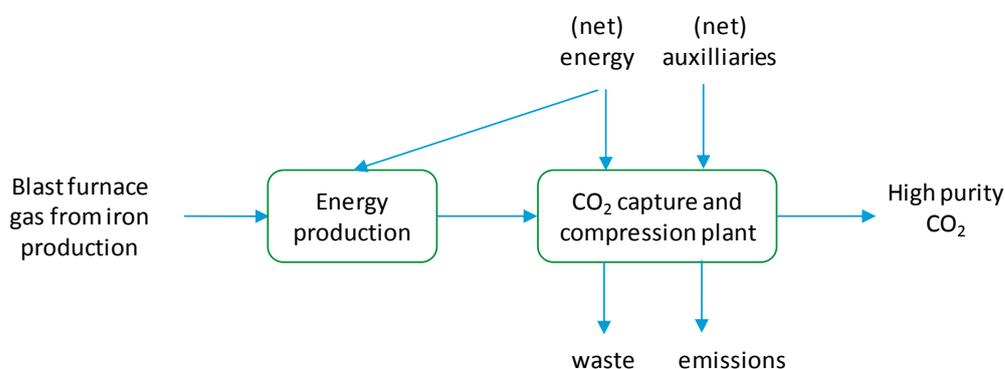
<sup>2</sup> Personal communication AVR, December 2017. Als reported by (ECN, DNV-GL, 2014).

<sup>3</sup> Based on data from (RIVM, 2017). Number applicable to 2015.

The blast furnace gas from Tata IJmuiden is currently fed into two different power plants (Velsen 25 and IJmond 1) where it is being incinerated to produce electricity. In case these two plants are not operational, a third plant (Velsen 24) will be used. Velsen 25 has the largest capacity of the three plants (375 MW). Therefore, this study looks at an amine-based capture method for the blast furnace process at the Velsen 25-plant. This technology is listed by the IEA as one of the primary technologies for CO<sub>2</sub> capture in iron production (IEA, 2013). For capture of CO<sub>2</sub> from blast furnace gas the amine considered is methyldiethanolamine (MDEA). After capture the CO<sub>2</sub> is compressed to the necessary pressure for the CO<sub>2</sub> Smart Grid.

Figure 4 shows a schematic representation of carbon capture from blast furnace gas from iron production for the iron production at Tata IJmuiden.

**Figure 4 - Schematic representation of carbon capture from blast furnace gas from iron production**



The CO<sub>2</sub> capture at the Velsen 25-plant leads to a reduction in electricity production of the plant. As described in Section 2.6, the reduction in electricity production is compensated by extra production of fossil electricity. An additional benefit of CO<sub>2</sub> capture in this way is that the heating value of the blast furnace gas increases (Zhang, et al., 2013). It has not been possible to quantify the impact of the increased heating value on the Velsen 25-plant, and the possible environmental benefit due to this is therefore not included. For the LCI of this capture technology see Annex A.

### 3.1.3 Fossil oil refining

The timeframe of realization of the large scale implementation of alternatives for conventional fuels (NH<sub>3</sub>, biobased), especially for shipping, is still unclear. There are several risks that pose serious barriers to the development and implementation of e.g. biofuels. The most important risks are related to strong fluctuations in oil price, which has recently negatively impacted bioenergy manufacturers (World Energy Council, 2016). This is also acknowledged by the European Commission, stating that, in 2030, 'fossil fuels continue to be by far the dominant energy source' (GAIN, 2017). Therefore, we assume in this study that fossil oil refining is likely to remain in place until 2030. However, it should be noted that fossil oil refining is likely to lose some market share to other fuel types. The International Maritime Organization (IMO), for instance, will introduce regulations on CO<sub>2</sub> emissions from shipping by 2023 (IMO, 2016).

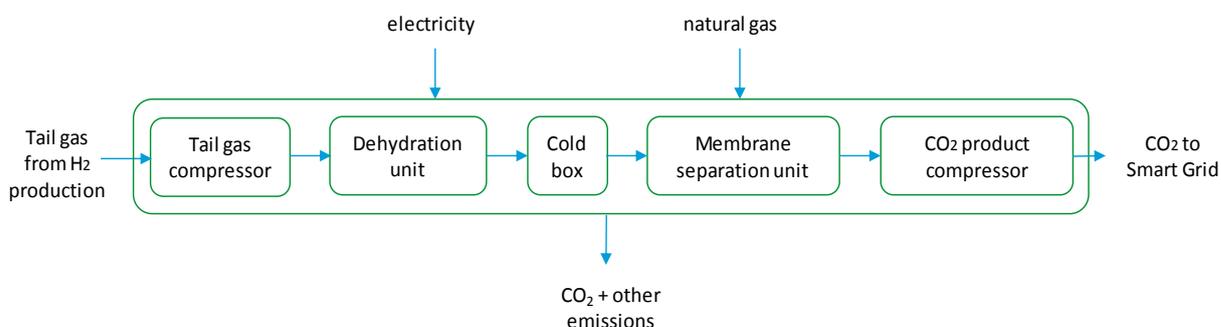
For fossil oil refineries, there are several different CO<sub>2</sub> emissions sources, such cracking reactors and hydrogen plants. For this sector we will consider CO<sub>2</sub> capture at the hydrogen plant. For capture at a hydrogen plant several different technologies are being applied commercially or demonstrated at

commercial scale<sup>4</sup>. In this case, cryogenic capture of the tail gas released during H<sub>2</sub> production will be considered. Benefits of this technology compared with alternative capture technologies include a very high purity CO<sub>2</sub> product.

Cryogenic capture (or 'low temperature separation') is based on separation principles involving the partial condensation of CO<sub>2</sub> and separating it from the gas phase in a distillation- or flash column (IEA, 2013). The selected specific technology is based on case 2B from IEA (2017), and includes the use of membranes in the setup of the CO<sub>2</sub> purification and compression unit. While not going into detail on its technological specifics, we briefly describe its components (see Figure 5):

- tail gas compressor: compresses tail gas to the required pressure of the cold box (see below);
- dehydration unit (dryers): dries compressed tail gas and lowers its temperature to below -55°C;
- cold box: contains coupled flash columns to separate the partially condensed CO<sub>2</sub> from the gas phase;
- membrane separation unit: recovers additional CO<sub>2</sub> from the output of the cold box;
- CO<sub>2</sub> product compressor: compresses the CO<sub>2</sub> to 110 bar(a).

Figure 5 - Carbon capture at H<sub>2</sub> production (cryogenic technology and membranes)



For the LCI of this capture technology see Annex A.

### 3.2 CO<sub>2</sub> upgrading: purification and compression

For utilization and for transport by means of the OCAP-pipeline system the captured CO<sub>2</sub> will have to meet specification requirements (see Table 4). In case specifications of the captured CO<sub>2</sub> do not meet requirements for utilization and/or transport, the CO<sub>2</sub> will have to be upgraded.

Table 4 - Specification requirements for applications and transportation

	Horticulture*	Mineralisation (Compensatiesteent)	MeOH production	CCS
CO <sub>2</sub> (vol%)	≥ 99.3%	60%	≥ 99.9%	≥ 99.9%
Pressure (bar(a))	≥ 21	unknown	50 - 100	130

\* Specifications as currently met in the OCAP pipeline.

<sup>4</sup> These include VPSA, amine based capture (BASF MDEA, Shell ADIP X), cryogenic capture and a combination of cryogenic separation and cold methanol (see e.g. (Zero Emissions Platform (ZEP), 2017).



The current OCAP-pipeline pressure is standardized at 21 bar(a). For a doubling of capacity, when realising the CO<sub>2</sub> Smart Grid, the pressure will need to be higher. We assume a necessity of approx. 40 bar(a) in pressure. This assumption was agreed on in the project meeting of December 5<sup>th</sup> 2017. For the CO<sub>2</sub> Smart Grid the purity of the CO<sub>2</sub> will need to be 99.9 vol% to meet the requirements of all the three studied applications.

For CCS an extra compression step until 130 bar(a) is required before injection in supercritical state.

Efficient compression to the required high pressure level of the CO<sub>2</sub> gas takes place in several stages. Based on polytropic efficiency in an electrically driven compressor the work per ton of CO<sub>2</sub> is calculated per stage using the following input variables:

- mass flow in kg/s;
- input pressure;
- output pressure;
- input temperature;
- gas compressibility;
- molar weight of the gas;
- polytropic efficiency of the compressor stage;
- electric motor efficiency.

The output pressure of the previous stage is used as the input pressure of the next stage. The next stage input temperature is after intercooling when applied. When the required pressure is reached no more stages are added. This results in the work of compression per stage which are added to deliver the total work of compression in kJ/kg CO<sub>2</sub> compressed.

### **3.3 CO<sub>2</sub> utilization**

#### **3.3.1 Horticulture**

Enhanced CO<sub>2</sub> levels in horticulture in greenhouses are essential for creating optimal growing conditions for commercial crops. The CO<sub>2</sub> used in Dutch greenhouses is currently supplied either by CO<sub>2</sub> produced from the combustion of natural gas in a gas burner, a CHP-unit or delivered from the OCAP-pipeline network. This latter network is a network in South-Holland currently supplying CO<sub>2</sub> from Alco and Shell to horticulture in South-Holland.

The horticulture sector is strongly committed to sustainability, and has the ambition to become carbon neutral by 2040. A boundary condition for realizing this goal is an abundance of externally available CO<sub>2</sub>. The availability of external CO<sub>2</sub> is seen by the sector as a key enabling factor in realizing this transition. Under these developments, application of captured CO<sub>2</sub> in horticulture provides one of most interesting and well-developed opportunities for CCU application (Ecofys, 2017). In the provinces North-Holland and South-Holland (i.e. roughly the area around the OCAP pipeline), horticulture is said to provide a CCU potential of 500 ktonnes at the moment, with the potential to increase to 1.2 Mtonne in 10 years. For the Netherlands, this potential is estimated at 2.1 Mtonne in 2030 due to the creation of new CCU projects (Berenschot ; EEI ; MEC, 2013).



## Description of utilization technology

For this utilization system we present two figures:

1. figure that shows the utilization system of the application of CO<sub>2</sub> from the CO<sub>2</sub> Smart Grid as plant growth enhancer in horticulture;
2. figure of the reference case (the alternative): using a gas burner for the generation of (useless) heat and CO<sub>2</sub>.

The dotted line indicates the elements of the system that are taken into account in assessing the environmental benefits of using captured CO<sub>2</sub> in this application.

In the reference case system, natural gas is burned to generate CO<sub>2</sub>. The heat that is unwanted in the greenhouse is released to the air. When using CO<sub>2</sub> from the CO<sub>2</sub> Smart Grid or OCAP-pipeline, the burning of natural gas in the summer is no longer needed. The quantity of natural gas incinerated that can be replaced by CO<sub>2</sub> from the CO<sub>2</sub> Smart Grid is determined based on the current incineration of natural gas in the summer, when the heat is not necessary for plant growth. Another method to determine the gas replaced by CO<sub>2</sub> is taken the average as has occurred in the current OCAP-pipeline. The choice for this approach is in line with previous research by CE Delft (CE Delft, 2017).

Figure 6 - System boundaries of utilization in horticulture – case A (current situation)

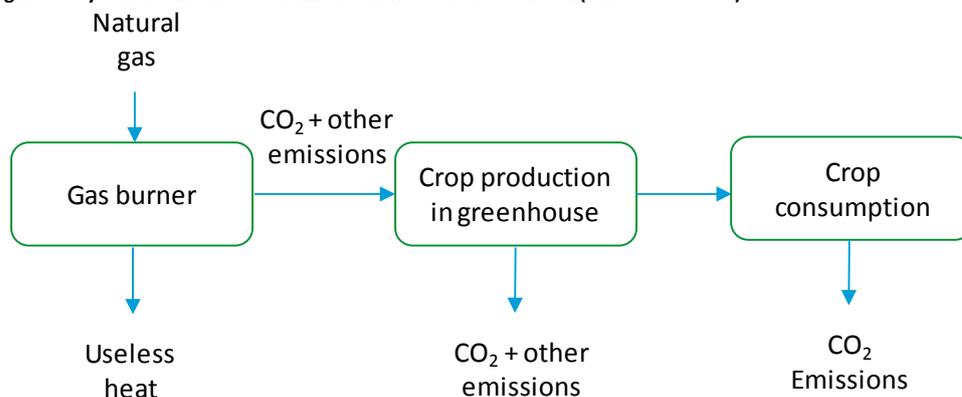
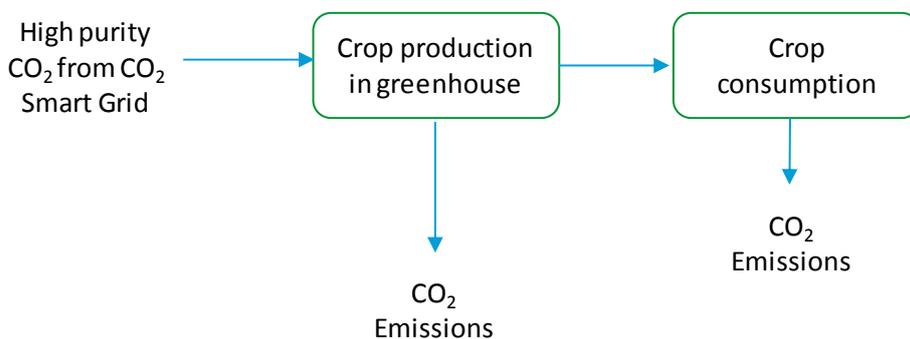


Figure 7 - System boundaries of utilization in horticulture – case B



## Produced products/services

All CCU routes that include utilization in horticulture produce the following products/services:

- capture of one tonne of CO<sub>2</sub>;
- increased plant growth.

## CO<sub>2</sub> storage time

The CO<sub>2</sub> that is sequestered in plants is released back into the atmosphere relatively quickly. Therefore, the storage of CO<sub>2</sub> in agricultural and horticultural crops is short-cyclical. In line with conventional CO<sub>2</sub> accounting practices, short-cyclical CO<sub>2</sub> is in this study not accounted for as a reduction in CO<sub>2</sub> emissions.

## Excluded: increased plant growth

A side effect of using a gas burner or combined heat and power (CHP) generator in summer to generate CO<sub>2</sub> for use in horticulture is that the production of CO<sub>2</sub> is limited by the production of heat. After all, crops are only able to grow properly at a certain maximum temperature. Therefore, when no heat is produced in the process of generating CO<sub>2</sub>, i.e. by using external CO<sub>2</sub>, the used amount of CO<sub>2</sub> per m<sup>2</sup> can be larger. This is likely to have a positive effect on the production efficiency of greenhouses (energy used per weight of crop produced) (Dieleman, et al., 2009). However, since no quantitative data is available on this issue, it has not been taken into account in this study.

## Excluded: alternative CO<sub>2</sub> source makes energy transition possible

Currently most greenhouses in North- and South-Holland are heated by means of a combined heat and power (CHP) unit. These CHPs use natural gas to produce three products: heat, electricity (also supplied to the grid) and CO<sub>2</sub> used as plant growth enhancer. This means that the supply of an alternative affordable CO<sub>2</sub> for use as plant growth enhancer can have the effect of making a transition towards a different heating technology for greenhouses possible. This is a situation in which the abundance of external CO<sub>2</sub> and its application in horticulture has enabled a transition to carbon-neutral heat. Carbon-neutral heat could for example be geothermal heat, residual heat, or a combination of these and other options. Since the exact impact that using an external CO<sub>2</sub> source has on the energy supply is unknown, this is not included in the LCA.

### 3.3.2 Mineralisation

In this application route, a mineral feedstock reacts with captured CO<sub>2</sub> to form an inert carbonate rock. Hereby, the carbon is chemically trapped and permanently sequestered. According to Ecofys, the market potential for carbonate mineralisation is somewhere between 100 and 300 ktonnes per year within ten years (Ecofys, 2017).

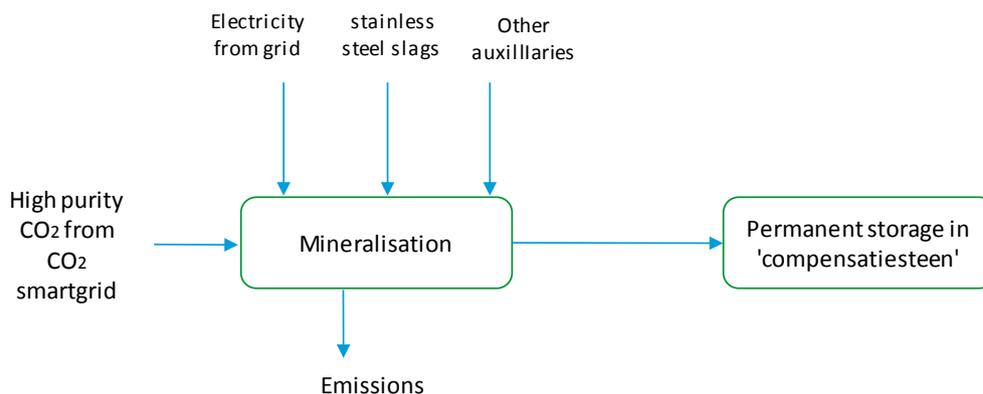
There are several technologies possible for carbonate mineralisation these include reaction of several waste products with CO<sub>2</sub> as well as the reaction of olivine (a mineral) with CO<sub>2</sub>. In this study, we consider the Carbstone-process, as developed by the Belgium research institute VITO, as an example for mineralisation. This process has been amended and is commercially applied (TRL 9) by the RuwBouw Groep, who sells a 'compensation stone' (Compensatiesteent) made through this technology. The RuwBouw Groep uses slags from stainless steel production, sand and CO<sub>2</sub> and converts this into a stone that can be used as a substitution for sand-lime bricks.



## Description of utilization technology

The current pilot plant of the RuwBouw Groep produces 3,000 m<sup>3</sup> of Compensatiesteen per year. The organisation is investigating the possibilities for setting up a full-scale production plant with a capacity of sequestering 80 ktonnes of CO<sub>2</sub>, equivalent to the production of 164,000 m<sup>3</sup> compensatiesteen. Compensatiesteen is produced by means of a hydraulic press, which uses little electricity. The stone is then cured in a CO<sub>2</sub> rich environment until it is fully saturated. Figure 8 shows the production and end-of-life of Compensatiesteen. For a full life cycle inventory see Annex A.

Figure 8 - System boundaries of utilization in carbonate mineralisation



## Produced products/services

All CCU routes that include utilization of CO<sub>2</sub> for mineralisation in Compensatiesteen produce the following products/services:

- capture of one tonne of CO<sub>2</sub>;
- Compensatiesteen.

## CO<sub>2</sub> storage time

The CO<sub>2</sub> used in mineralisation is permanently stored, and will only come free again with continuous weathering of rock or when treated in an industrial process.

## Prevention of sand-lime brick production

Compensatiesteen is a hard, stone-like material that is currently used in construction applications where originally sand-lime bricks would be used. RuwBouw Groep expects that the stone can also be used in conventional non-constructive concrete applications if the permit procedure for this application has been completed. Non-constructive applications include concrete parts which, with the exception of any transport and auxiliary reinforcement, do not contain any structural reinforcement. In this LCA we consider the prevention of sand-lime brick production.

## Conventional use stainless steel slags

Stainless steel slags are currently treated and used as aggregates or sand in road construction<sup>5</sup>. If the stainless steel slags are used to produce Compensatiesteent, the aggregate will need to come from elsewhere. The environmental impact of aggregate production elsewhere is taken into consideration in this study.

## Excluded: cleaning stainless steel slags

The stainless steel slags used by the RuwBouw Group are cleaned before being used in the Compensatiesteent. However, it is currently unclear where the cleaning process takes place, and whether this process requires a large amount of associated energy use and/or other inputs. We expect that in comparison to the conventional application of stainless steel slags as granulate or sand in road construction, no extra treatment is needed.

### 3.3.3 Methanol

According to Ecofys, the Dutch market potential for CO<sub>2</sub> based methanol amounts to 220 ktonnes/year within ten years (Ecofys, 2017). In methanol production the captured CO<sub>2</sub> is hydrogenated with separately produced hydrogen. This hydrogen in the studied CCU route is produced through electrolysis: the process of using electricity to split water into hydrogen and oxygen. We study the production of methanol and the electrolysis based on a fossil fuel mix (as described in Section 2.6) as well as based on directly coupled renewable energy.

## Description of utilization technology

We consider the process as it is currently applied by Carbon Recycling International (CRI). CRI runs a demonstration installation with a 4,000 tonnes/year production capacity of 'Vulcanol' which has been operational since 2012<sup>6</sup>. CRI aims at a commercial scale of 35-40 ktonnes/year. The TRL level of this technology is estimated to be TRL 7-8. Vulcanol is fuel grade methanol which can be blended with gasoline for automobiles and used in the production of biodiesel or fuel ether. In addition, Vulcanol can be used in the production of several synthetic materials.

Figure 9 shows the utilization system of CO<sub>2</sub> from the CO<sub>2</sub> Smart Grid as feedstock for the production of methanol production based on this technology. The process yields methanol and water and some combustible by-products, which may be marketed/supplied to external customers. The heat of the exothermic CO<sub>2</sub> hydrogenation reaction is partially used to heat feed streams and for distillation of the raw product.

We study this CCU in the following four cases:

- complete renewable electricity use, short term sequestration of CO<sub>2</sub> (e.g. fuel);
- complete renewable electricity use, long term sequestration of CO<sub>2</sub> (e.g. chemical);
- complete fossil electricity use, short term sequestration of CO<sub>2</sub> (e.g. fuel);
- complete fossil electricity use, long term sequestration of CO<sub>2</sub> (e.g. chemical).

In the case of production with completely renewable energy use, the hydrogen is considered to be produced with renewable energy with a direct connection to the hydrogen plant, e.g. hydrogen produced by water electrolysis with electricity from *directly coupled* wind power or photovoltaic power. Hydrogen production by way of electrolysis and methanol production need not take place at

<sup>5</sup> See for example the products sold by Orbix: [www.orbix.be/nl/materialen](http://www.orbix.be/nl/materialen)

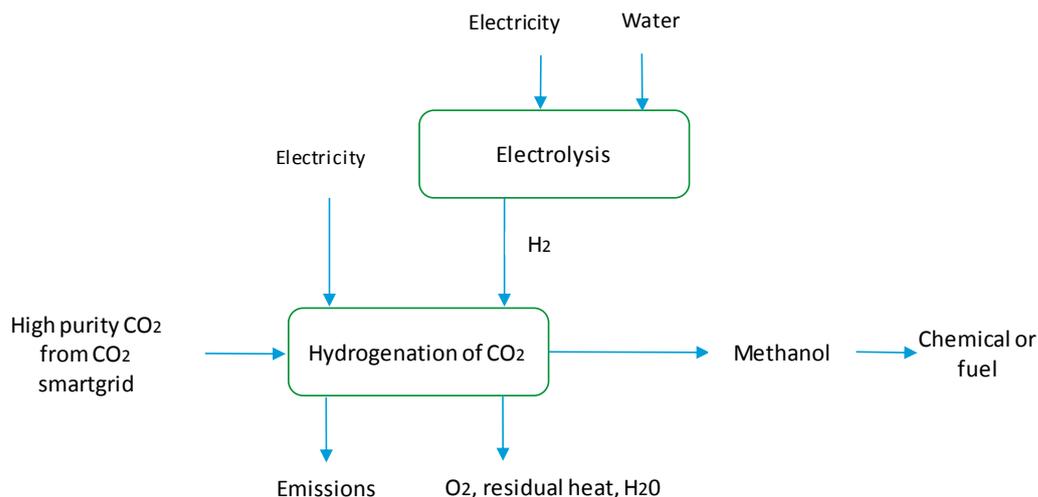
<sup>6</sup> A second technology-provider is Japanese company Mitsui Chemicals Inc., but their technology seems less evolved.



the same location if hydrogen production can be connected with methanol production by way of a pipeline, e.g. the existing Air Liquide North-western high pressure H<sub>2</sub> network. Such a high pressure system may act or be utilized as a H<sub>2</sub> buffer by way of the ‘line pack’<sup>7</sup> of the system.

In the case of production based in fossil electricity mix we use the carbon footprint of electricity as given in Section 2.6.

**Figure 9 - System boundaries of production of methanol from CO<sub>2</sub> through hydrogenation**



### Produced products/services

All CCU routes that include utilization of CO<sub>2</sub> for methanol production produce the following products/services:

- capture of one tonne of CO<sub>2</sub>;
- methanol.

The combustible by-products, residual heat and O<sub>2</sub> could be marketed as products but are not considered to be so in the base case modelling because not enough data has been obtained to do so.

### CO<sub>2</sub> storage time

Given the wide range of applications for methanol it is undoable in this project to consider each of them. We will therefore indicate the time period during which the CO<sub>2</sub> utilized in methanol production is ‘sequestered’ in these applications. This will be done for two extremes in terms of duration:

- use in fuels (e.g. as oxygenate or as a component in biodiesel methyl esters or MTBE/TAME);
- use as chemical for use as a component in technical plastics.

<sup>7</sup> The intrinsic volume of the pipeline system.



In the case of use in fuels the carbon storage is short-cyclical, as the fuel is relatively quickly burned. In line with conventional CO<sub>2</sub> accounting practices, short-cyclical CO<sub>2</sub> is in this study not accounted for as a reduction in CO<sub>2</sub> emissions (see Section 2.4). In the case of use as chemical we assume CO<sub>2</sub> storage time of more than 100 years when used for technical plastic production that can be recycled several times.

### **Prevention of diesel production and use (application as fuel)**

The reference technology for CO<sub>2</sub> based methanol production used as fuel is the production and use of conventional diesel for transportation.

### **Prevention of conventional methanol production (application as chemical)**

The reference technology for CO<sub>2</sub> based methanol production is conventional methanol production in world scale units, utilizing stranded gas.



## 4 Reference technology: CCS

### 4.1 Introduction

One of the main questions to be considered and evaluated in this report is whether it is worthwhile in terms of CO<sub>2</sub> sequestration and/or other environmental aspects to utilize captured CO<sub>2</sub> for each of the considered applications instead of immediate geological storage in offshore abandoned gas fields or in offshore deep aquifers. Therefore an introduction is given into the carbon capture and storage (CCS) technology.

### 4.2 Background

CCS deposits captured carbon from large point sources to storage sites such that it will not enter the atmosphere, normally deposition occurs in underground geological formation such as abandoned gas fields or offshore deep aquifers. The CO<sub>2</sub> is captured, compressed, transport and subsequently injected.

For CCS as a reference case to this study, the injection step is that only step that differs from the CCU routes. The capture and transportation of CO<sub>2</sub> is also included for all CCU cases. For the injection, a compressor is used, which compresses the captured CO<sub>2</sub> into a supercritical fluid. The CO<sub>2</sub> is then injected under pressure into the geological formation, where it is trapped under an impermeable layer of rock. In this study, the electricity that is needed to inject the captured CO<sub>2</sub> into the geological formation is taken into account, as well as (possible) carbon leakage from the compressor.

### 4.3 Literature review

Several studies have assessed the carbon footprint of CCS technologies. In addition, a number of meta-studies that critically compare a variety of LCAs involved with this topic have been published.

(Cuéllar-France & Azapagic, 2015) published a well-cited comprehensive article in which numerous LCAs of CCS and CCU technologies are compared. The authors conclude that, on average, the Global Warming Potential (GWP) of CCS is significantly lower than that of CCU options. However, other environmental impacts, such as acidification potential and human toxicity potential might be higher compared to CCU. A number of CCS studies specifically address lowering the GWP of power plants. In this case, the GWP is reduced by 63-82%.

Another well-cited article describes the LCA of a pulverized coal power plant with post-combustion capture, transport and storage of CO<sub>2</sub> (Koornneef, et al., 2008). While the study is slightly older, it is situated in the Netherlands, and therefore relevant to this study. The authors show that GHG-emissions per kWh produced are reduced by 71-78%, depending on the technological advancement of the power plant. The International Energy Agency published a synthesis report of LCAs of CCS technologies in 2010 (Marx, et al., 2011). The results of the LCAs of the coal power generation systems with CCS clearly indicate a substantial reduction in GWP of around 80%. Similar results are shown for application of CCS at lignite power plants.

A German study from 2007 presents an LCA and cost assessment of CCS technologies at hard coal-fired power plants and compares this to renewable energy solutions (Viebahn, et al., 2007).



The conclusion of the study is that CO<sub>2</sub> emissions per kWh for CCS technologies are 72-90% lower than for coal-fired power plants without CCS.

A more recent Norwegian study assesses the environmental impact of carbon capture in the context of a natural gas combined cycle electricity generation plant (Singh & Hertwich, 2011). The authors show that, when sequestering 90% CO<sub>2</sub> from the flue gas, 70% of CO<sub>2</sub> emissions per kWh are avoided. The Global Warming Potential is reduced by 64%. However, a number of environmental impact on midpoint level are influenced conversely: for example, both acidification (43%) and eutrophication (35%) increase. This is a similar result as (Cuéllar-France & Azapagic, 2015).

#### **4.4 Conclusion**

The consulted peer-reviewed academic references present that the reduction in carbon dioxide emissions from power plants range between 63-90%, strongly depending on the carbon capture technology and carbon source. This means that between 630 and 900 kg of CO<sub>2</sub> is sequestered per tonne of captured CO<sub>2</sub> for more than 100 years in a CO<sub>2</sub> storage location. The carbon sources studied in this study are different than those looked at in the literature, but the literature gives a good insight in the order of magnitude of sequestered CO<sub>2</sub>. Some studies indicate that trade-offs might occur on other environmental effects. This points towards the importance of, in further studies, also taking into account e.g. acidification and eutrophication effects.



## 5 Results: Global warming

The results for this screening LCA of CCU routes are presented in two ways: per carbon capture technology/carbon source and per utilization technology. Subsequently, different forms of utilization can be more easily compared, whereas it also enables us to draw more attention to the environmental performance of the different capture methods.

### Global warming, CO<sub>2</sub> and CO<sub>2</sub> eq.

Global warming is caused by greenhouse gasses. The most commonly known greenhouse gas is carbon dioxide (CO<sub>2</sub>). This is, however, not the only greenhouse gas, other such gasses include methane and dinitrogen monoxide. All other greenhouse gasses can be expressed in CO<sub>2</sub> eq.; the global warming potential of a greenhouse gas compared to carbon dioxide. In this chapter we look at the impact of the CCU routes on global warming. We have not only looked at CO<sub>2</sub> emissions, but also other greenhouse gas emissions. *When referring to CO<sub>2</sub> emissions or reduction of CO<sub>2</sub> emissions we are therefore technically speaking about CO<sub>2</sub> eq. and not only CO<sub>2</sub>.*

### 5.1 Results per carbon capture technology/carbon source

In this section, the results are shown separately for each the three carbon capture technologies/carbon sources.

#### 5.1.1 Carbon capture at a MWI

Table 5 shows the emitted CO<sub>2</sub> and the net avoided CO<sub>2</sub> emission of the different utilization-routes for CO<sub>2</sub> captured at an MWI in comparison with not capturing CO<sub>2</sub> at a municipal waste incinerator, including a breakdown. Figure 10 shows the emitted CO<sub>2</sub> of the different utilization-routes for CO<sub>2</sub> captured at an MWI.

Table 5 - Net avoided CO<sub>2</sub> emission per CCU/CCS route compared to non-capture

	Capture from MWI and utilization in horticulture#	Capture from MWI and utilization for mineralisation	Capture from MWI and utilization for methanol production* 100% renewable energy	Capture from MWI and utilization for methanol production* 100% fossil-based energy	Capture from MWI and storage (CCS)
CO <sub>2</sub> emission capture technology (kg/tonne captured)	239 kg	239 kg	239 kg	239 kg	239 kg
CO <sub>2</sub> emission product/service production (kg/tonne captured)	0 kg	116 kg	568 kg	2634 kg	24 kg
CO <sub>2</sub> emission end-of-life (within 100 years) (kg/tonne captured)	1,000 kg, of which: 361 kg fossil based 639 kg biogenic	0 kg	1,000 kg, of which: 361 kg fossil based 639 kg biogenic	1,000 kg, of which: 361 kg fossil based 639 kg biogenic	0 kg



	Capture from MWI and utilization in horticulture#	Capture from MWI and utilization for mineralisation	Capture from MWI and utilization for methanol production* 100% renewable energy	Capture from MWI and utilization for methanol production* 100% fossil-based energy	Capture from MWI and storage (CCS)
CO <sub>2</sub> emission reduction replacement (kg/tonne captured)	-1,076 kg	-286 kg	-1,163 kg	-1,163 kg	0 kg
Total CO <sub>2</sub> emitted (kg/tonne captured)	162 kg	69 kg	644 kg	2710 kg	262 kg
CO <sub>2</sub> emitted without CO <sub>2</sub> capture at MWI	- 1,000 kg	- 1,000 kg	- 1,000 kg	- 1,000 kg	- 1,000 kg
Reduction of CO <sub>2</sub> emission in comparison to current situation (kg/tonne captured)	- 838 kg	- 931 kg	-356 kg	1710 kg (emission increase)	- 738 kg

# Results for utilization in horticulture are based on utilization in summer. For more horticulture-results see Section 5.2.1.

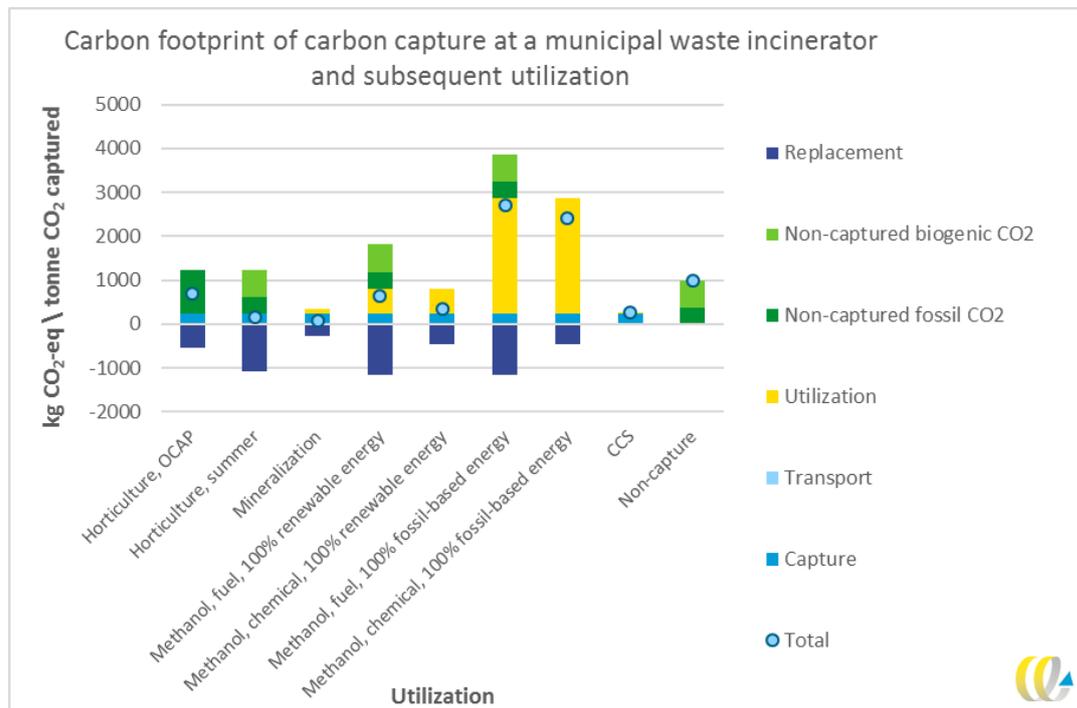
\* Results for methanol production are based utilization of methanol used as fuel. For more methanol-results see Section 5.2.3.

Table 5 and Figure 10 show that the utilization-route of methanol is the least preferable option, especially when fossil fuel is used. There are several reasons for this. Firstly, all captured CO<sub>2</sub> is emitted very rapidly again if the methanol is applied in an application that sequesters the CO<sub>2</sub> shorter than 100 years, for example when the methanol is used as a fuel. In addition, the current available production technique for methanol from CO<sub>2</sub> is not so efficient, which is reflected in the relatively high value for the emissions associated with utilization. When methanol is used for the production of a chemical for an application where the CO<sub>2</sub> is sequestered for more than 100 years, the methanol utilization-route comes closer to CCS.

The carbon footprint of utilization of CO<sub>2</sub> in horticulture can be the same order of magnitude as that of CCS when the CO<sub>2</sub> is applied during summer. This is mostly linked to the large benefit associated with the avoided incineration of natural gas in the summer months. It is, in this case, the question whether this situation will still be relevant in the (near) future, and especially towards 2050, when heat production in the horticulture sector in the Netherlands will become carbon neutral. The in that case reference is no longer necessarily natural gas incineration but CO<sub>2</sub> could also be supplied by e.g. a wood burner. When adding CO<sub>2</sub> to horticulture year round, as currently applied in the OCAP-pipeline the reduction is approximately 500 kg CO<sub>2</sub> eq. lower, and no longer comparable to CCS. It can therefore be concluded that the CO<sub>2</sub> emission reduction is dependent on the specific situation.

For the mineralisation-route, the results indicate that long-term sequestration of captured carbon could be a good option. In addition, the replacement of sand-lime brick is relatively certain, and still quite a conservative (i.e. simple) avoided product. The energy use of the utilization of this route is also modest in terms of carbon footprint. There are however some uncertainties surrounding the energy use for utilization, since the modelling has been based completely on data supplied by the producer of Compensatiestein. In the sensitivity analysis we will delve further into this uncertainty (see Chapter 7).

Figure 10 - Carbon footprint of carbon capture at a MWI and subsequent utilization per tonne captured CO<sub>2</sub>



### 5.1.2 Carbon capture from blast furnace gas from iron production

Table 6 shows the emitted CO<sub>2</sub> and the net avoided CO<sub>2</sub> emissions of the different utilization-routes for CO<sub>2</sub> captured from blast furnace gas from iron production in comparison with not capturing the CO<sub>2</sub>, including a breakdown. Figure 11 shows a breakdown of the emitted CO<sub>2</sub> of the different utilization-routes for CO<sub>2</sub> captured from blast furnace gas.



**Table 6 - Net avoided CO<sub>2</sub> emission per CCU/CCS route compared to non-capture**

	Capture from blast furnace gas and utilization in horticulture#	Capture from blast furnace gas and utilization for mineralisation	Capture from blast furnace gas and utilization for methanol production* 100% renewable energy	Capture from blast furnace gas and utilization for methanol production* 100% fossil-based energy	Capture from blast furnace gas and storage (CCS)
CO <sub>2</sub> emission capture technology (kg/tonne captured)	220 kg	220 kg	220 kg	220 kg	220 kg
CO <sub>2</sub> emission product/service production (kg/tonne captured)	0 kg	116 kg	568 kg	2,634 kg	24 kg
CO <sub>2</sub> emission end-of-life (within 100 years) (kg/tonne captured)	1,000 kg	0 kg	1,000 kg	1,000 kg	0 kg
CO <sub>2</sub> emission reduction replacement (kg/tonne captured)	- 1,076 kg	- 286 kg	- 1,163 kg	- 1,163 kg	0 kg
Total CO <sub>2</sub> emitted (kg/tonne captured)	144 kg	50 kg	625 kg	2,691 kg	244 kg
<i>CO<sub>2</sub> emitted without CO<sub>2</sub> capture from blast furnace gas</i>	<i>- 1,000 kg</i>	<i>- 1,000 kg</i>	<i>- 1,000 kg</i>	<i>- 1,000 kg</i>	<i>- 1,000 kg</i>
Reduction of CO <sub>2</sub> emission in comparison to current situation (kg/tonne captured)	- 856 kg	- 950 kg	- 375 kg	1,691 kg (emission increase)	- 756 kg

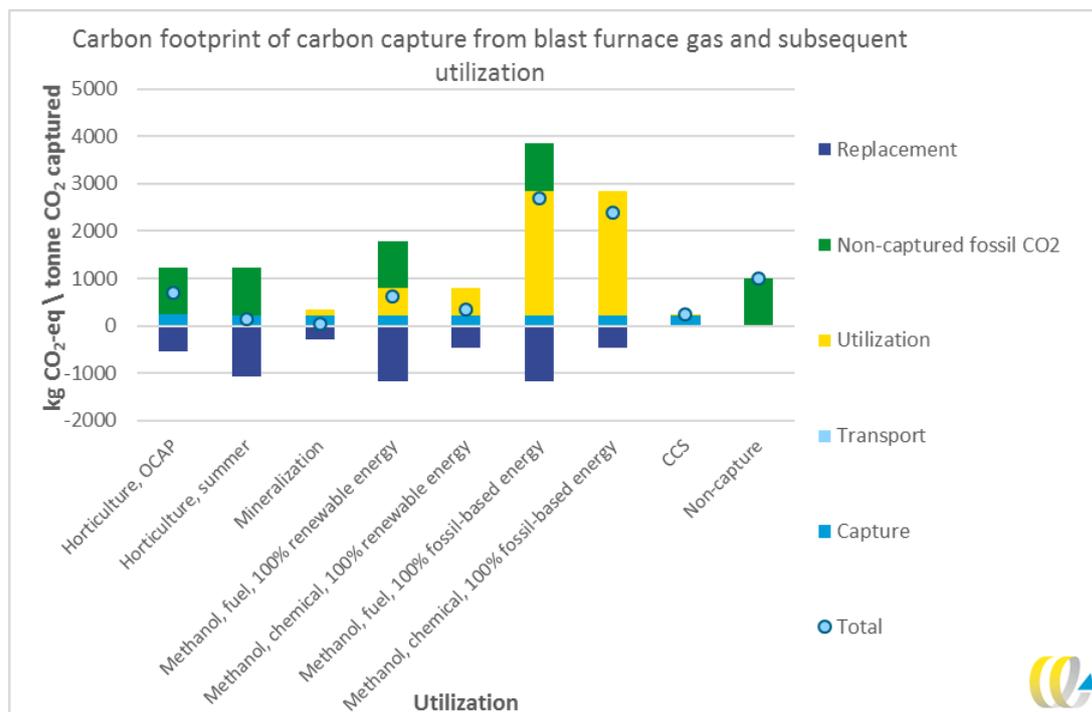
# Results for utilization in horticulture are based on utilization in summer. For more horticulture-results see Section 5.2.1.

\* Results for methanol production are based utilization of methanol used as fuel. For more methanol-results see Section 5.2.3.

For the case of carbon capture from blast furnace gas, Figure 11 shows that the carbon footprint of the capture technique is comparable to that of capture at an MWI. The methanol-route, when produced from fossil-based energy, is again the least favourable option.



Figure 11 - Carbon footprint of carbon capture from blast furnace gas and subsequent utilization per tonne captured CO<sub>2</sub>



### 5.1.3 Carbon capture at hydrogen plant (from fossil oil refining)

Table 7 shows the emitted CO<sub>2</sub> and the net avoided CO<sub>2</sub> emissions of the different utilization-routes for CO<sub>2</sub> captured at a hydrogen plant (from fossil oil refining) in comparison with not capturing the CO<sub>2</sub>, including a breakdown. Figure 12 shows a breakdown of the emitted CO<sub>2</sub> of the different utilization-routes for CO<sub>2</sub> captured at a hydrogen plant.

Table 7 - Net avoided CO<sub>2</sub> emission per CCU/CCS route compared to non-capture

	Capture at a hydrogen plant and utilization in horticulture#	Capture at a hydrogen plant and utilization for mineralisation	Capture at a hydrogen plant and utilization for methanol production* 100% renewable energy	Capture at a hydrogen plant and utilization for methanol production* 100% fossil-based energy	Capture at a hydrogen plant and storage (CCS)
CO <sub>2</sub> emission capture technology (kg/tonne captured)	129 kg	129 kg	129 kg	129 kg	129 kg
CO <sub>2</sub> emission product/service production (kg/tonne captured)	0 kg	116 kg	568 kg	2,634 kg	24 kg
CO <sub>2</sub> emission end-of-life (within 100 years) (kg/tonne captured)	1,000 kg	0 kg	1,000 kg	1,000 kg	0 kg
CO <sub>2</sub> emission reduction replacement	- 1,076 kg	- 286 kg	- 1,163 kg	- 1,163 kg	0 kg



	Capture at a hydrogen plant and utilization in horticulture#	Capture at a hydrogen plant and utilization for mineralisation	Capture at a hydrogen plant and utilization for methanol production* 100% renewable energy	Capture at a hydrogen plant and utilization for methanol production* 100% fossil-based energy	Capture at a hydrogen plant and storage (CCS)
(kg/tonne captured)					
Total CO <sub>2</sub> emitted (kg/tonne captured)	53 kg	-41 kg	535 kg	2,600 kg	153 kg
CO <sub>2</sub> emitted without CO <sub>2</sub> capture at a hydrogen plant	- 1,000 kg	- 1,000 kg	- 1,000 kg	- 1,000 kg	- 1,000 kg
Reduction of CO <sub>2</sub> emission in comparison to current situation (kg/tonne captured)	947 kg	1,041 kg	465 kg	1,600 kg (emission increase)	847 kg

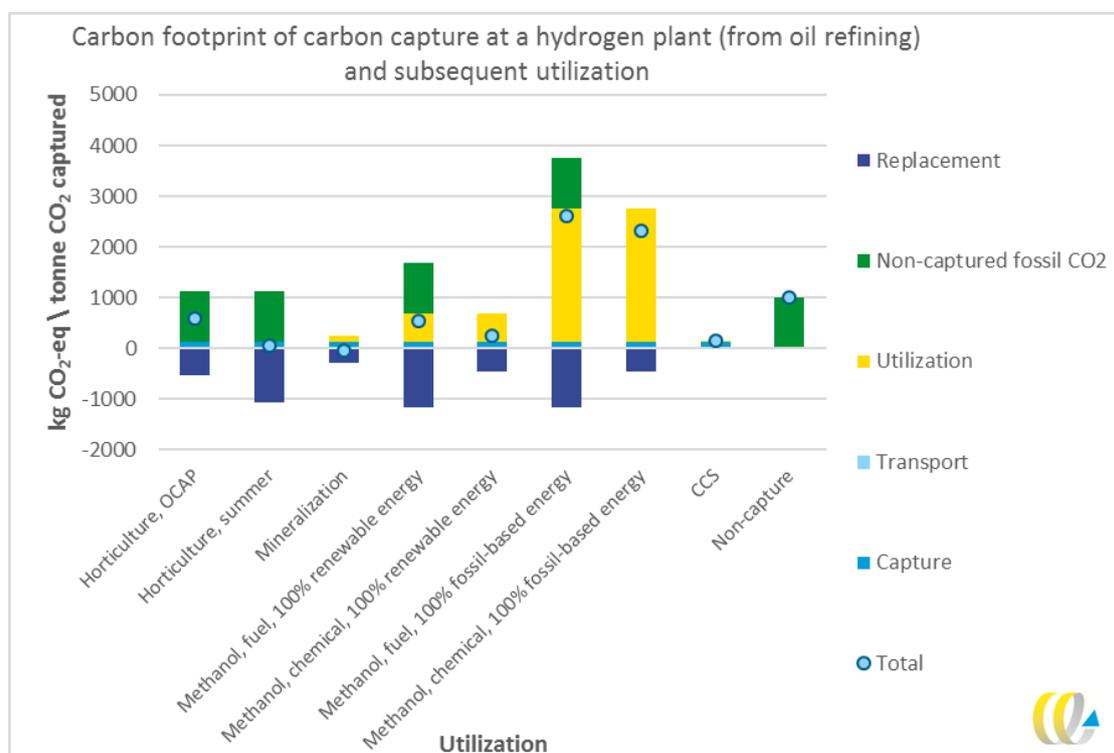
# Results for utilization in horticulture are based on utilization in summer. For more horticulture-results see Section 5.2.1.

\* Results for methanol production are based utilization of methanol used as fuel. For more methanol-results see Section 5.2.3.

Since the carbon footprint of the capture of CO<sub>2</sub> at fossil oil refining is comparable to that of capture from blast furnace gas as described in Section 5.1.2, the results of the different utilization technologies combined with capture do not differ much. Again mineralisation leads to a negative carbon dioxide emission (more carbon dioxide being captured than emitted), application in horticulture (in summer) is comparable to CCS while application in methanol production based on fossil-based energy is the least preferable option.



Figure 12 - Carbon footprint of carbon capture at a hydrogen plant (from fossil oil refining) and subsequent utilization per tonne captured CO<sub>2</sub>



## 5.2 Results per utilization technology

In this section, we present the estimated carbon footprint per utilization technology.

### 5.2.1 Utilization in horticulture

Table 8 shows the emitted CO<sub>2</sub> and the net avoided CO<sub>2</sub> emission of the different CO<sub>2</sub> sources/capture technologies and utilization of CO<sub>2</sub> in horticulture in comparison with not capturing the CO<sub>2</sub>, including a breakdown. Figure 13 shows a breakdown of the emitted CO<sub>2</sub>.

Table 8 - Net avoided CO<sub>2</sub> emission per CCU route compared to non-capture

	Capture at MWI plant and utilization in horticulture	Capture at iron production and utilization in horticulture	Capture at hydrogen plant and utilization in horticulture
CO <sub>2</sub> emission capture technology (kg/tonne captured)	239 kg	220 kg	129 kg
CO <sub>2</sub> emission product/service production (kg/tonne captured)	0 kg	0 kg	0 kg
CO <sub>2</sub> emission end-of-life (within 100 years) (kg/tonne captured)	1,000 kg, of which: 361 kg fossil based 639 kg biogenic	1,000 kg	1,000 kg
CO <sub>2</sub> emission reduction replacement *	-1,076 kg (summer)	-1,076 kg (summer)	-1,076 kg (summer)

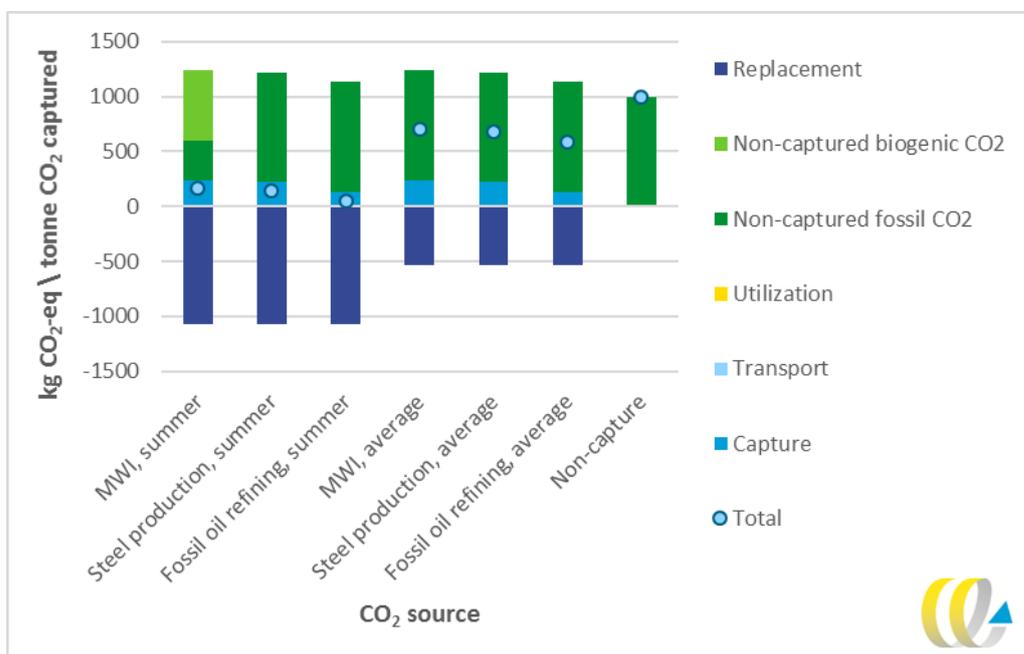
	Capture at MWI plant and utilization in horticulture	Capture at iron production and utilization in horticulture	Capture at hydrogen plant and utilization in horticulture
(kg/tonne captured)	- 538 kg (OCAP)	- 538 kg (OCAP)	- 538 kg (OCAP)
Total CO <sub>2</sub> emitted (kg/tonne captured) *	162 kg (summer) 700 kg (OCAP)	144 kg (summer) 682 kg (OCAP)	53 kg (summer) 591 kg (OCAP)
CO <sub>2</sub> emitted without CO <sub>2</sub> capture	- 1,000 kg	- 1,000 kg	- 1,000 kg
Reduction of CO <sub>2</sub> emission in comparison to current situation (kg/tonne captured) *	838 kg (summer) 300 kg (OCAP)	856 kg (summer) 318 (OCAP)	947 kg (summer) 409 kg (OCAP)

Note: These are indicative figures, and serve to give an order-of-magnitude-estimation.

\* This table gives a range of values for both utilization of CO<sub>2</sub> in summer and the reduction of CO<sub>2</sub> emission as is achieved in the current OCAP-pipeline year round.

Table 8 and Figure 13 shows that for all three carbon capture technologies the utilization of the captured carbon in horticulture leads to net CO<sub>2</sub> emissions and that net more than 800 kg of CO<sub>2</sub> emission avoided per tonne of CO<sub>2</sub> captured when the CO<sub>2</sub> is added to horticulture in summer. This is because currently the CO<sub>2</sub> used in greenhouses in the Netherlands largely originate from natural gas combustion, the prevention of natural gas use (the replacement) compensates for a large part of the CO<sub>2</sub> emissions. In the current OCAP pipeline approximately half of the CO<sub>2</sub> added to the greenhouses prevents the use of natural gas use, on average 300 kg of CO<sub>2</sub> emission is avoided per tonne of CO<sub>2</sub>-captured and added to a greenhouse.

Figure 13 - Carbon footprint of carbon capture and utilization in horticulture per tonne of captured CO<sub>2</sub>



## Future energy supply horticulture

When the incineration of natural gas is no longer the most logical supply for CO<sub>2</sub>, i.e. when the heat supply will become carbon-neutral, it can be argued that the application of captured CO<sub>2</sub> in horticulture no longer needs to lead to the prevention of natural gas use. In that case the CO<sub>2</sub> could also be supplied by e.g. a wood burner. If that is the case the CO<sub>2</sub> emissions from utilizing captured CO<sub>2</sub> in horticulture will be higher than the quantity of CO<sub>2</sub> captured because of the energy demand for the capturing technology.

### 5.2.2 Utilization in mineralisation (Compensatiestein)

Figure 9 shows the emitted CO<sub>2</sub> and the net avoided CO<sub>2</sub> emissions of the different CO<sub>2</sub> sources/capture technologies and utilization of CO<sub>2</sub> for mineralisation in comparison with not capturing the CO<sub>2</sub>, including a breakdown. Figure 14 shows a breakdown of the emitted CO<sub>2</sub>.

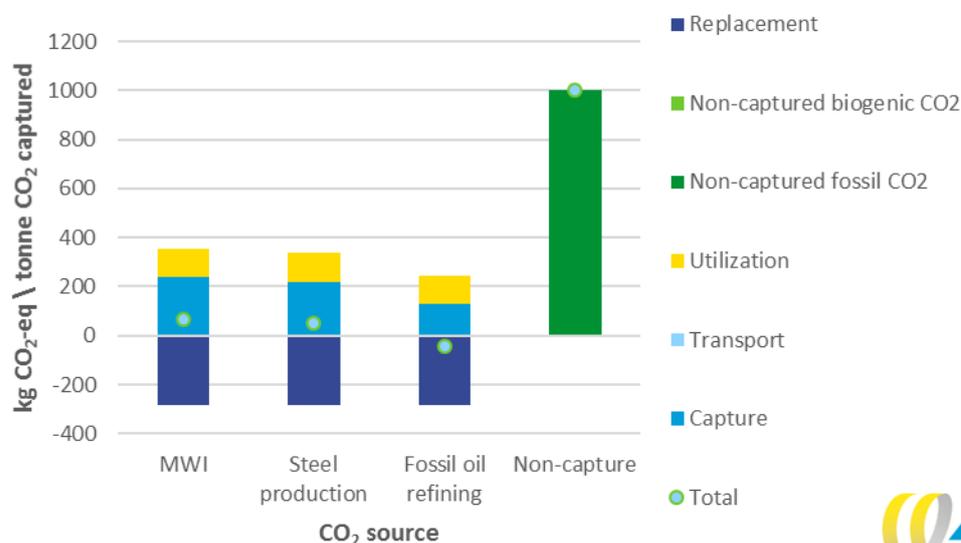
Table 9 - Net avoided CO<sub>2</sub> emission per CCU route compared to non-capture

	Capture at MWI plant and utilization for mineralisation	Capture at iron production and utilization for mineralisation	Capture at hydrogen plant and utilization for mineralisation
CO <sub>2</sub> emission capture technology (kg/tonne captured)	239 kg	220 kg	129 kg
CO <sub>2</sub> emission product/service production (kg/tonne captured)	116 kg	116 kg	116 kg
CO <sub>2</sub> emission end-of-life (within 100 years) (kg/tonne captured)	0 kg	0 kg	0 kg
CO <sub>2</sub> emission reduction replacement (kg/tonne captured)	- 286 kg	- 286 kg	- 286 kg
Total CO <sub>2</sub> emitted (kg/tonne captured)	69 kg	50 kg	-41 kg
CO <sub>2</sub> emitted without CO <sub>2</sub> capture	- 1,000 kg	- 1,000 kg	- 1,000 kg
Reduction of CO <sub>2</sub> emission in comparison to current situation (kg/tonne captured)	931 kg	950 kg	1,041 kg

Note: These are indicative figures, and serve to give an order of magnitude estimation.

Table 9 and Figure 14 show that the lower the carbon footprint of the capture technology is, the more likely that mineralisation of CO<sub>2</sub> in Compensatiestein will lead to a net negative CO<sub>2</sub> emission. The figure also shows that, even in the case of a relatively high carbon footprint of the capture technology, such as capture at the MWI, there is a reduction of more than 90% of the CO<sub>2</sub> emissions compared to non-capture.

**Figure 14 - Carbon footprint of carbon capture and utilization for mineralisation (Compensatiesteen) per tonne of captured CO<sub>2</sub>**



### 5.2.3 Utilization in methanol production

We study this CCU route in the following four cases:

1. Complete renewable electricity use, short term sequestration of CO<sub>2</sub> (e.g. fuel).
2. Complete renewable electricity use, long term sequestration of CO<sub>2</sub> (e.g. chemical).
3. Complete fossil electricity use, short term sequestration of CO<sub>2</sub> (e.g. fuel).
4. Complete fossil electricity use, long term sequestration of CO<sub>2</sub> (e.g. chemical).

To make the comparison as easy as possible the range of values for the four cases with the three studied capture methods/CO<sub>2</sub> sources is shown.

Table 10 shows the emitted CO<sub>2</sub> and the net avoided CO<sub>2</sub> emission of the different cases in comparison with not capturing the CO<sub>2</sub>, including a breakdown. Figure 15 shows a breakdown of the emitted CO<sub>2</sub>.

**Table 10 - Net avoided CO<sub>2</sub> emission for capture and utilization for methanol production compared to non-capture**

	Renewable electricity CO <sub>2</sub> storage <100 years	Renewable electricity CO <sub>2</sub> storage >100 years	Fossil electricity CO <sub>2</sub> storage <100 years	Fossil electricity CO <sub>2</sub> storage >100 years
CO <sub>2</sub> emission capture technology (kg/tonne captured)*	129 – 239 kg	129 – 239 kg	129 – 239 kg	129 – 239 kg
CO <sub>2</sub> emission product/service production (kg/tonne captured)	568 kg	568 kg	2,634 kg	2,634 kg
CO <sub>2</sub> emission end-of-life (within 100 years) (kg/tonne captured)	1,000 kg	0 kg	0 kg	0 kg
CO <sub>2</sub> emission reduction replacement (kg/tonne captured)	- 1,163 kg	- 451kg	- 1,163 kg	- 451 kg



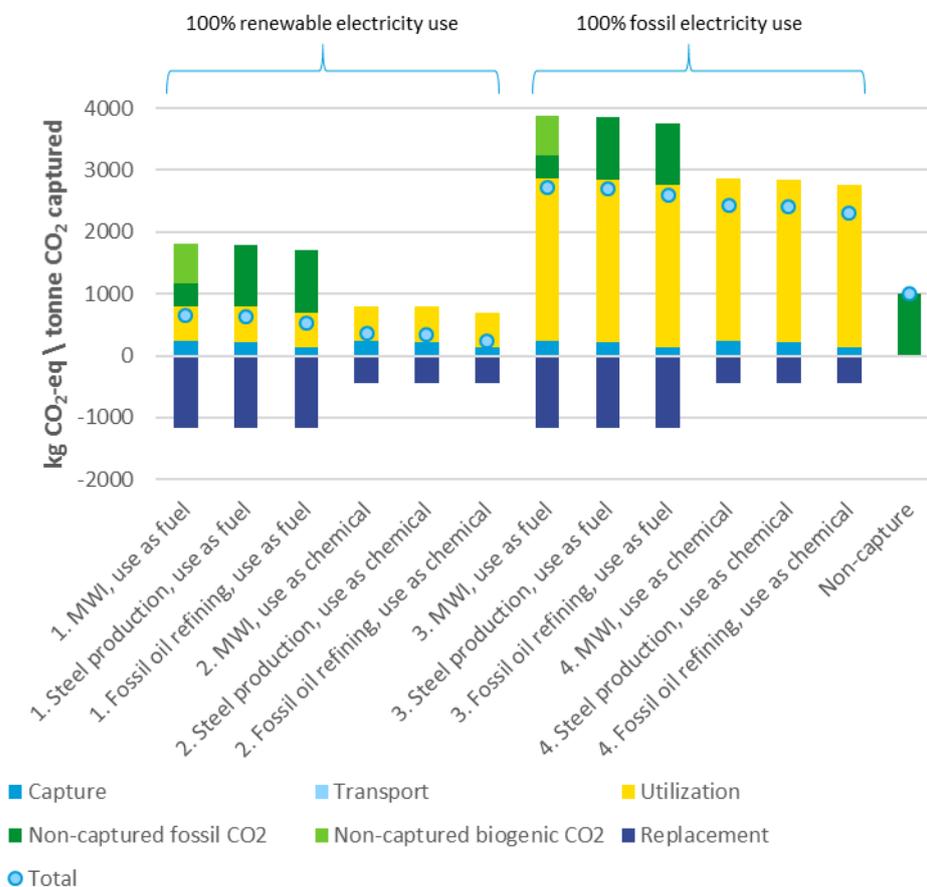
	Renewable electricity CO <sub>2</sub> storage <100 years	Renewable electricity CO <sub>2</sub> storage >100 years	Fossil electricity CO <sub>2</sub> storage <100 years	Fossil electricity CO <sub>2</sub> storage >100 years
Total CO <sub>2</sub> emitted (kg/tonne captured)	535 – 644 kg	246 – 355 kg	2,600 – 2,710 kg	2,312 – 2,421 kg
CO <sub>2</sub> emitted without CO <sub>2</sub> capture	- 1,000 kg	- 1,000 kg	- 1,000 kg	- 1,000 kg
Reduction of CO <sub>2</sub> emission in comparison to current situation (kg/tonne captured)	- 356 – - 465 kg	- 645 – - 754 kg	1,600 – 1,710 kg	1,312 – 1,421 kg

\* This table gives a range of values for all three capture technologies studied.

In case of production of methanol from CO<sub>2</sub> with non-renewable electricity there will be no reduction in CO<sub>2</sub> emissions in comparison to non-capture. In the case of 100% renewable electricity use for the hydrogen and methanol production a net reduction of CO<sub>2</sub> emission can be achieved ranging between 350 kg and 750 kg per tonne of CO<sub>2</sub> captured. The higher end of this spectrum can be reached with a capture technology with low CO<sub>2</sub> footprint, and utilization of the methanol in an application where the CO<sub>2</sub> is stored for more than 100 years.



Figure 15 - Carbon footprint of carbon capture and utilization for methanol production



Note: The methanol routes consider 100% renewable energy use, and applications where the CO<sub>2</sub> is stored for less than 100 years (e.g. fuels).

**Discussion: renewable energy in methanol production**

The utilization of 100% renewable electricity in the production of methanol using captured CO<sub>2</sub> might naturally lead to a discussion regarding the administration of the environmental benefits of this electricity. In the Netherlands, renewable electricity is largely made possible through the SDE+ subsidy scheme, introduced to accelerate the large-scale implementation of renewable energy technologies. The subsidy itself is made possible by the Dutch government, and mainly Dutch consumers who pay an extra fee for their electricity.

When strictly interpreting LCA rules, the environmental benefits of the renewable energy produced through the SDE+ system should therefore be rewarded to the government and consumers. Parties that make the realization of additional renewable energy possible through e.g. additional funding could make the decision to use the renewable energy for the production of 'green' methanol. However, in this case, it is important to stress that in the coming years, the net CO<sub>2</sub> reduction of this application of renewable electricity will be lower than when it will be used directly to replace fossil electricity.



## 6 Results: Other environmental impacts

### Environmental benefit additional cleaning of CO<sub>2</sub> containing gas

For coal-fired power plants, the deployment of carbon capture results in an additional environmental advantage: additional cleaning of the produced flue gases. This advantage results in lower emissions of e.g. SO<sub>2</sub> and particulate matter of coal-fired power plants: see (Royal Haskoning, 2011). For the three CO<sub>2</sub> sources considered in this study, any possible additional cleaning of CO<sub>2</sub>-containing gases has not been described in detail in literature. Therefore the additional advantages of this additional cleaning are not expressed in the results of this study.

For example, the emissions associated with blast furnace gas include hydrogen sulphide, fine particulate matter and carbonyl sulphide (COS). It is likely, in line with what occurs at coal-fired plants when applying capture, that some hydrogen sulphide and fine particulate matter will be captured along with CO<sub>2</sub>. COS is unlikely to be captured.

### Capture from blast furnace gas and MWI: environmental costs of additional emissions

Furthermore additional emissions from capture associated with the application of absorbent have not been taken into consideration because of a lack of data. It is however known that the use of MEA as an absorbent has in the past led to the production of aerosols. The MDEA absorbent is less prone to degeneration than MEA is but the exact emissions because of the use of this absorbent are unknown. Additionally, engineering measures to prevent the emissions of these aerosols can mitigate this. Also, whether or not Bilisol (the absorbent used for capture from the MWI) degenerates is not known.

### Environmental impacts of utilization technology system

Because of the lack of data described on the possible environmental benefits and environmental costs of the difference between capture and non-capture at the CO<sub>2</sub> capture location we exclusively describe the environmental impacts relevant for air quality that are related to the utilization technology. This study has considered the following environmental impact of the three utilization technology systems:

1. Fine particulate matter formation (PM<sub>2.5</sub> emissions).
2. Terrestrial acidification (SO<sub>2</sub> emissions).
3. Tropospheric ozone formation (NO<sub>x</sub> emissions).

All utilization technology systems lead to a reduction of environmental impacts in these impact categories, even when considering fossil energy use for the capturing technologies. This means that in all cases the conventional production of product that is being replaced (natural gas combustion, sand-lime brick and methanol) has higher emissions than the emissions from the electricity used for the CCU (capture plus utilization).

### **Future research needed**

As indicated in Chapter 4, possible trade-offs between reduction in CO<sub>2</sub> emissions and acidification and eutrophication exist. The acidification and eutrophication impact of the different CCU routes have not been studied. In further research it is important to take these into consideration, as well as other emissions occurring at the CO<sub>2</sub> capture location.



# 7 Sensitivity analysis

Because of the short time frame in which this screening LCA has been carried out there are uncertainties surrounding the results discussed in the previous chapter. This is partly due to the fact that the available data originates from one source, and time was too limited to verify the data. In this chapter we describe the uncertainties that have been identified that make it possible to reach (firmer) conclusions about the studied CCU routes.

Uncertainties can arise because of several reasons. In this study, they mainly originate from a lack of available data (or time to obtain the data) and the difficulty of studying environmental impacts in the future. The most important uncertainties are briefly described below.

## 7.1 Uncertainties because of data availability

Uncertainties related to data availability include missing data on:

- compression energy for capture from MWI;
- carbon footprint of absorbents; carbon footprint of stainless steel slags.

Furthermore uncertainties exist in the used data, because it has not been possible to verify all data obtained from a single data source.

### Capture from MWI: Compression energy

The energy needed for compression of the CO<sub>2</sub> captured from the MWI has been set at its most conservative because the pressure of the produced CO<sub>2</sub> was not mentioned in the used literature. The energy use for the compression accounts for approximately 20% of the CO<sub>2</sub> emissions from the capture at the MWI in the current calculations.

*This uncertainty could lead to a reduction of the carbon footprint of capture at the MWI compared to the results that are presented in Chapter 5, and could make the carbon footprint of this capture technology more comparable to the carbon footprint of the other two capture technologies studied.*

### Capture from blast furnace gas: Carbon footprint absorbent

The carbon footprint of the MDEA-absorbent needed for the capture of CO<sub>2</sub> from blast furnace gas is not publically available. An approximation of the footprint has been made based on the production of methylamine. This is likely to be an underestimation of the actual carbon footprint.

*This uncertainty could lead to a slight increase of the carbon footprint of the capture of CO<sub>2</sub> from iron production compared to the results in Chapter 5.*

### Capture from fossil oil refining

Because the capture technology considered for fossil oil refining does not produce CO<sub>2</sub> with the purity vol% required for use in the CO<sub>2</sub> Smart Grid an extra purification step is needed. The CATOX-technology could be used to do so. The only necessary input for this process, besides infrastructure, is pure O<sub>2</sub>. Since very little O<sub>2</sub> is needed, the production of O<sub>2</sub> has not been taken into consideration



because the exact quantity needed is unknown. An environmental burden is however associated with this production.

*This uncertainty could lead to a slight increase of the carbon footprint of the capture of CO<sub>2</sub> from fossil oil refining compared to the results in Chapter 5. The increase can be expected to be low because of the small quantity of O<sub>2</sub> needed.*

### **Utilization for mineralisation: stainless steel slags**

Stainless steel slags have been modelled as having no environmental impact because of their status as a waste product. However, stainless steel slags are actually used as aggregate in road construction, and therefore, a part of the environmental emissions for the stainless steel production should be attributed to this product. Because the economic value of stainless steel slags is unknown, economic allocation has however not been applied.

*This uncertainty could lead to a slight increase of the carbon footprint of the utilization of CO<sub>2</sub> for mineralisation. The impact can however be expected to be limited since a tonne of steel has a much higher value than a tonne of aggregate.*

### **All CCU routes: verification of data**

Because of the short time span of the study, data for several processes within the CCU route have been obtained from a single data source. This data has not been verified extensively. Some uncertainties exist because of this, including:

- Electricity use for the production of Compensatiestein, which is much lower than that of the Carbstone technology, of which it has been derived from<sup>8</sup>.
- Data on capture at the MWI has been obtained from (Monteiro, et al., 2015), a study conducted by Procede, the owner of the technology. This data has not been verified except for order of magnitude verifications.
- Data on production of methanol has been based on data on a single pilot plant from (Stefansson, 2015). This data has not been verified except for order of magnitude verifications.

## **7.2 Uncertainties due to future development of CO<sub>2</sub> Smart Grid**

This study looks at the implementation of a CO<sub>2</sub> Smart Grid in 2030. Since it is difficult to predict the future there are a number of uncertainties concerning future development of the studied CCU routes. These include:

- other applications of Compensatiestein;
- sustainable heat supply in Dutch horticulture;
- optimization possibilities for methanol production;
- additional cleaning of CO<sub>2</sub> containing gas;
- exact requirements of the CO<sub>2</sub> for the CO<sub>2</sub> Smart Grid;
- uncertainty of electricity use for CCU routes.

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<sup>8</sup> For the Carbstone technology energy use has been determined at 200 kWe per m<sup>3</sup> Carbstone concrete (Vito, 2014). While the Ruwbouwgroupp reports electricity use that is approximately 80% lower.



### **Utilization for mineralisation: other applications of Compensatiesteen**

Compensatiesteen has currently been tested and approved for use in applications where it prevents the use of sand-lime brick. The stone could also be applied in applications where it replaces concrete, but has not been approved for this application. The application of Compensatiesteen instead of concrete would lead to a bigger 'replacement' of CO<sub>2</sub> emissions than when using it instead of sand-lime brick.

*If Compensatiesteen can replace concrete in 2030, the CCU routes including mineralisation would lead to a higher net avoided CO<sub>2</sub> emission than shown in the results in Chapter 5.*

### **Utilization for horticulture: sustainable heat supply**

The future heat supply for Dutch horticulture is unknown, and therefore also the possible sources for CO<sub>2</sub> used in greenhouses. A possibility is the use of biomass for both heat and CO<sub>2</sub> production, but also geothermal heat supply not yielding any CO<sub>2</sub> emissions is an option. Whether or not the application of captured CO<sub>2</sub> aids the shift towards renewable energy and what would be the appropriate reference CO<sub>2</sub> source to consider in the future is a topic that needs further discussion.

*If a sustainable heat supply is in place in 2030 in Dutch horticulture, the CCU routes including utilization for horticulture could lead to a lower net avoided CO<sub>2</sub> emission than shown in the results in Chapter 5.*

### **Utilization for methanol production: marketing of by-products**

In the current study we have only looked at methanol production from captured CO<sub>2</sub> as it is currently applied in a pilot plant in Iceland. A possible optimization of the current practice is the marketing of by-products such as residual heat and O<sub>2</sub> from electrolysis. The O<sub>2</sub> and residual heat will need to meet the specifications required by the market.

*If the by-products of methanol production can be marketed, the CCU-routes including methanol would lead to a lower carbon footprint than shown in Chapter 5.*

### **Requirements CO<sub>2</sub> Smart Grid: compression and purity**

The requirements of the CO<sub>2</sub> Smart Grid are not yet known. The exact compression of CO<sub>2</sub> needed for transport over distance as well as the required purity of CO<sub>2</sub> for the utilization technologies attached to the grid remain to be determined when the exact utilization technologies are known.

*When less compression and a lower vol% of CO<sub>2</sub> is required, the environmental impact of upgrading the CO<sub>2</sub> stream to the desired level will decrease. This means that the carbon footprint of all CCU routes would decrease in comparison those shown in Chapter 5.*



### **All CCU-routes: future renewable electricity use**

In CCU routes electricity is used. We have in line with LCA and Dutch policy practices used the fossil electricity type on the margin (see Section 2.6). It is possible that the different CCU routes ensure that they use renewable electricity. E.g. using directly coupled renewable electricity for the production of Compensatiesteun.

*If the electricity used would be from directly coupled renewable electricity, the carbon footprint of a CCU routes would decrease in comparison those shown in Chapter 5.*



## 8 Conclusion

Here, we summarize the results presented in Chapter 5. In addition, we formulate conclusions based on the sensitivity assessments shown in Chapter 7.

### **Capture technologies and carbon sources considered have comparable carbon footprints**

The three different capture technologies do not differ significantly in carbon footprint. The capture of CO<sub>2</sub> from iron production and capture of CO<sub>2</sub> the MWI are particularly comparable in terms of carbon footprint. The footprint of capture from fossil oil refining (at the hydrogen plant) is slightly lower, but the difference between the technologies in the results could be due to uncertainties surrounding the data gathered.

### **Utilization in mineralisation**

Utilization of CO<sub>2</sub> for mineralisation, the production of Compensatiesteen, leads to net avoided CO<sub>2</sub> emissions of around 1 tonne of CO<sub>2</sub> per tonne of CO<sub>2</sub> captured. Despite the carbon footprint of the capture technologies, the produced Compensatiesteen leads to the avoided production of conventional sand-lime brick. It is possible that the footprint of the utilization technology is slightly higher than portrayed in this report because of an uncertainty surrounding the energy use in the process. This requires further study but will not lead to the technology having a net CO<sub>2</sub> emission compared to non-capture. mineralisation

### **Utilization in horticulture**

In the current situation, the utilization of CO<sub>2</sub> in horticulture leads to net avoided CO<sub>2</sub> emissions between 300 and around 950 kg CO<sub>2</sub> per tonne of CO<sub>2</sub> captured. The highest end of this range is for use of CO<sub>2</sub> in summer and is a comparable or even better performance than well-functioning CCS. The net avoided CO<sub>2</sub> emission is caused by the avoided use of natural gas for the production of CO<sub>2</sub> in horticulture. The lower end of the range is not comparable with CCS and that what is currently reported for captured CO<sub>2</sub> added to the OCAP-pipeline.

The future heat supply for Dutch horticulture is unknown, and therefore also the possible sources for CO<sub>2</sub> used in greenhouses. A possibility is the use of biomass for both heat and CO<sub>2</sub> production, but also geothermal heat supply not yielding any CO<sub>2</sub> emissions is an option. Whether or not the application of captured CO<sub>2</sub> aids the shift towards renewable energy and what would be the appropriate reference CO<sub>2</sub> source to consider in the future is a topic that needs further discussion. Therefore the exact carbon footprint reduction after a switch to a renewable heat source in the horticulture sector has been made is uncertain.

### **Utilization in methanol production**

Utilization in methanol production will only lead to net avoided CO<sub>2</sub> emissions when renewable energy is used for methanol and hydrogen production. The net avoided CO<sub>2</sub> emissions will increase when the CO<sub>2</sub> is used in durable products. 'Durable' in this context implies that CO<sub>2</sub> is sequestered for



more than 100 years. In that case, this utilization method could reach net avoided CO<sub>2</sub> emissions of around 700 kg CO<sub>2</sub> per tonne of CO<sub>2</sub> captured. This is comparable to the least efficient type of CCS.

It must be noted that methanol production is not the only possible application of CO<sub>2</sub> in the chemical industry. Other possible utilizations could include the production of polyols for the production polyurethanes.

### **Conclusions on other environmental impacts**

For several reasons, no conclusions could yet be drawn on other environmental impacts and on emissions from the capturing technology:

- additional benefits caused by the additional cleaning of CO<sub>2</sub> containing gas are unknown;
- emissions from degradation of absorbents are unknown.

Because of the lack of data described on the possible environmental benefits and environmental costs of the difference between capture and non-capture at the CO<sub>2</sub> capture location we only describe the environmental impacts relevant for air quality that have to do with the utilization technology.

This study has considered the following environmental impact of the three utilization technology systems; Fine particulate matter formation (PM<sub>2.5</sub> emissions), Terrestrial acidification (SO<sub>2</sub> emissions) and Tropospheric ozone formation (NO<sub>x</sub> emissions). All utilization technology systems lead to a reduction of environmental impacts in these impact categories, even when considering fossil energy use for the capturing technologies. This means that in all cases the production of product that is replaced (natural gas combustion, sand-lime brick and methanol) has higher emissions than the emissions from the electricity used for the utilization technology and in the capturing process.

Possible trade-offs between reduction in CO<sub>2</sub> emissions and acidification and eutrophication exist. The acidification and eutrophication impact of the different CCU routes has not been studied, and in further research it is important to take these into consideration.

### **Interpretation of the conclusions**

The orders of magnitude of CCS and CCU applicability in 2030 are expected to be incomparable. E.g. the potential storage by means of CCS is expected to be much higher than the potential for use of CO<sub>2</sub> in mineralization in the Netherlands. Results must therefore only be seen on a per tonne basis and cannot be extrapolated. The spatial application of the technologies also differ, e.g. CCS can be applied the whole year round while the peak of CO<sub>2</sub> utilization in horticulture is during the growing season and less so in winter.

Because the study carried out is a screening LCA, the drawn conclusions should be seen as indicative figures; they offer an order of magnitude estimation and cannot be seen as representative for individual (industrial) plants present in the Netherlands. Furthermore, because a substitution methodology has been used, the results are not appropriate for consumption-based carbon accounting (see Brander & Wylie, 2011). This means that, when calculating the emissions of a country's total consumption, LCA results that are calculated through the substitution methodology cannot be included. The same holds for using the outcomes for corporate carbon accounting practices.

To make the results applicable to individual CCU routes e.g. CO<sub>2</sub> capture at the AEB MWI in Amsterdam and application of the CO<sub>2</sub> in horticulture in Aalsmeer, a full scale LCA study will need to be conducted.



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# A Life cycle inventory

For this screening LCA, various reports and studies were used for collecting relevant data. This chapter summarizes the data and sources used, based on the previously described system boundaries.

## A.1 Carbon capture from CO<sub>2</sub> source and preparation for injection into Smart CO<sub>2</sub> grid

### Municipal waste incineration (MWI) plant

A study has been conducted on the application of the Procede Gas Treating technology for capture of CO<sub>2</sub> specifically for injection of CO<sub>2</sub> into the OCAP pipeline. This study gives a description of the consumables for the quench unit, the capture plant and the compression needed for injection. Because of the difference in compression assumed needed for the CO<sub>2</sub> Smart Grid, we have only used the data on the quench unit and the capture plant. See Table 11 for the overview of the inputs needed as obtained from (Monteiro, et al., 2015).

Table 11 - Net inputs for CO<sub>2</sub> capture and preparation for injection into the Smart CO<sub>2</sub>-grid at a MWI

	Conventional MWI	MWI with capture	Net	Net per tonne captured CO <sub>2</sub>
Electricity for capture	-	1,693 MWh/year	1,693 MWh/year	0.10 GJ/tonne CO <sub>2</sub>
Steam for capture	-	57,596 MWh/year	57,596 MWh/year	3.48 GJ/tonne CO <sub>2</sub>
Cooling water for capture	-	9,384,595 m <sup>3</sup> /year	9,384,595 m <sup>3</sup> /year	157 m <sup>3</sup> /tonne CO <sub>2</sub>
CO <sub>2</sub> capture		59,600 tonne/year	59,600 tonne/year	

Data source: (Monteiro, et al., 2015).

### Reduction in electricity production

There is one aspect that is not included in the process described by (Monteiro, et al., 2015); a reduction in electricity production because of the steam/heat used by the capture technology. According to the AVR (MWI of Rotterdam) the reduction in electricity production is approximately 0.25 MWe per MW heat extracted. This means that per tonne CO<sub>2</sub> captured the electricity production decreased with approximately 0.87 GJe.

The reduction of electricity production can be seen as an electricity input needed for the CO<sub>2</sub> capture and is taken into consideration as electricity input from the Dutch electricity grid.

### Compression

Monteiro, et al., (2015) do not mention the exact pressure of the produced CO<sub>2</sub> stream. We therefore assume that it is produced at a 1 bar(a) pressure. This means that the stream still needs to be compressed to 40 bar(a). Further compression has been based on the operational conditions of a compressor given in (Geological Survey of the Netherlands, 2009). Based on this source to get to 40 bar(a) from 1 bar(a) an approximate 295 MJ/tonne CO<sub>2</sub> captured is needed.



Monteiro, et al.,(2015) also do not mention the exact purity of the CO<sub>2</sub> gas stream produced. However, since their study refers to producing CO<sub>2</sub> to be injected in the OCAP line, the purity is likely to be 93% (see Section 3.2).

The off gas of MWIs needs to reach a high purity level; there is stringent emission regulation in the Netherlands. The CO<sub>2</sub> capture unit is placed after the conventional purification steps. This might lead to a further reduction in emissions, but this is not taken into consideration in this screening LCA because of a lack of data.

## Blast furnace gas from the blast furnace process

This study looks at an amine-based capture method for the blast furnace process. This technology is listed by the IEA as one of the primary technologies for CO<sub>2</sub> capture in iron production (IEA, 2013). The net inputs for this technology are given by (IEA, 2013) and shown in Table 12.

**Table 12 - Net inputs for CO<sub>2</sub> capture from blast furnace gas and preparation for injection into the Smart CO<sub>2</sub> grid**

	Conventional iron production	Iron production with capture	Net	Net per tonne captured CO <sub>2</sub>
Electricity for capture	-	572,622,619 kWh year	572,622,619 kWh/year	0.6 GJ/tonne CO <sub>2</sub>
Steam for capture	-	8,082,495 GJ/year	8,082,495 GJ/year	2.35 GJ/tonne CO <sub>2</sub>
MDEA make-up for capture	-	688 tonne/year	688 tonne/year	0.2 kg/tonne CO <sub>2</sub>
MDEA disposal for capture	-	688 tonne/year	688 tonne/year	0.2 kg/tonne CO <sub>2</sub>
Raw water for capture	-	10,557,185 m <sup>3</sup> /year	10,557,185 m <sup>3</sup> /year	3 m <sup>3</sup> /year
CO <sub>2</sub> capture	-	3,439,360 tonne/year	3,439,360 tonne/year	-

Data source: (IEA, 2013).

## Reduction in electricity production

There is one difference between the technology as described by the IEA and the likely application of the technology for Tata IJmuiden: the blast furnace gas from Tata IJmuiden is currently fed to two different power plants (Velsen 25 and IJmond 1) where it is being incinerated to produce electricity. In case these two plants are not operational a third plant (Velsen 24) is used. Of the three Velsen 25 has the largest capacity of 375 MW. Applying the CO<sub>2</sub> capture at the Velsen 25 plant leads to a reduction in electricity production of Velsen 25.

The reduction of electricity production can be seen as an electricity input needed for the CO<sub>2</sub> capture. Figure 16 shows the configuration of the Velsen 25 plant. The steam produced by the turbine is 540°C and 180 bar(a), the steam is fed into the High Pressure (HP) turbine after which it is fed back into the boiler where it is reheated to 540°C and 40 bar(a). The steam is then used in the Intermediate Pressure (IP) turbine where after which it reaches Low Pressure (LP) turbines. The condensing occurs at 24 degrees and 30 mbar(a).

Figure 17 shows the Velsen 25 plant with CO<sub>2</sub> capture assuming no changes in the boiler efficiency and no net change in parasitic power consumption. The steam for the MDEA reboiler (2.35 GJ/tonne CO<sub>2</sub> as given by (IEA, 2013)) is supplied from the outlet of the IP turbine. The outlet of the IP turbine is the most logical place to tap steam from since it has the least influence on electricity production (latest



possible stage) and still reaches the required 120°C required for the MDEA reboiler. The loss in electricity production therefore occurs at the LP turbine.

The overall electric efficiency of the Velsen 25 plant is 43%. The HP turbine has the highest efficiency and the LP turbine the lowest efficiency. The approximate efficiency of the LP turbine is 27%. Assuming the turbine runs on full load, the reduction of efficiency is minimal due to the steam extraction. This therefore leads to a reduction in production of 0.65 GJe per tonne of CO<sub>2</sub> captured.

Figure 16 - Velsen 25 plant without CO<sub>2</sub> capture, per 0.48 tonne CO<sub>2</sub>

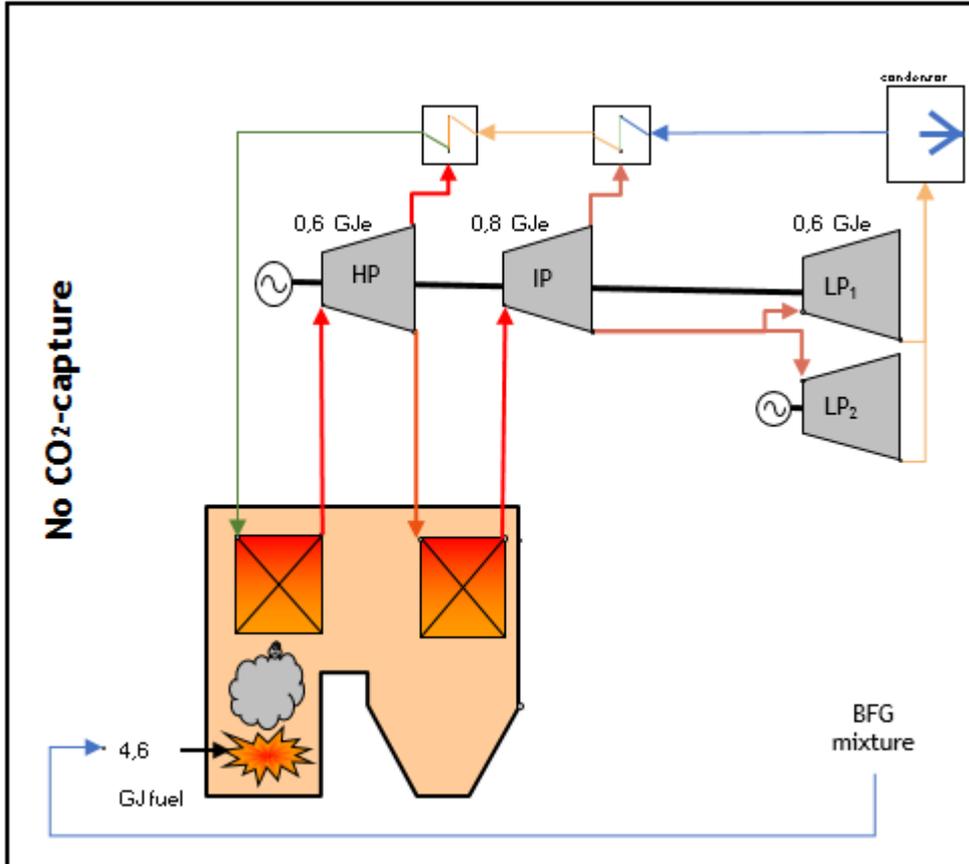
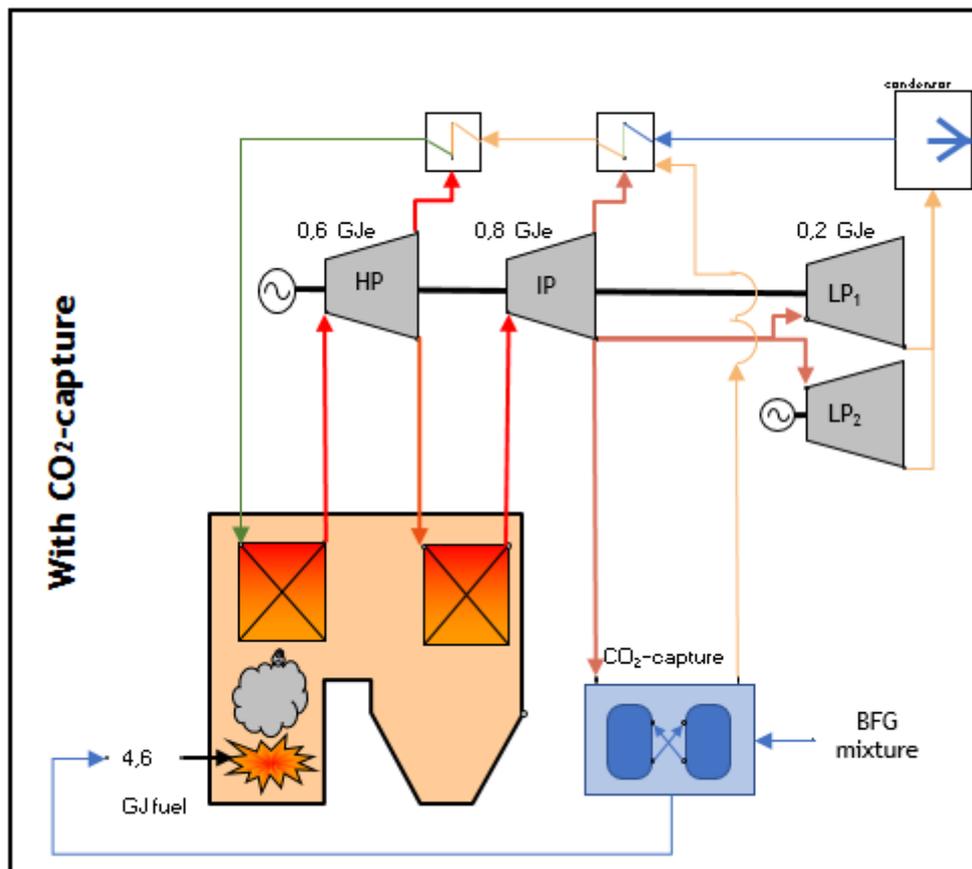


Figure 17 - Velsen 25 plant with CO<sub>2</sub> capture, per 0.48 tonne CO<sub>2</sub>



### Increased temperature blast furnace gas

Due to the capture technology the lower heating value of the blast furnace gas increases (Zhang, et al., 2013). This means that a higher electricity production could be achieved when supplying the blast furnace gas after CO<sub>2</sub> capture to the boiler. Since the exact influence of an increase in lower heating value for the Velsen 25 plant is unknown the result of the increased temperature blast furnace gas is not included in this study.

### Compression

The capture technology as described by (IEA, 2013) produces CO<sub>2</sub> with a purity of 99.9% at a pressure of 110 bar(a). To be able to meet the specifications of 40 bar(a) for the CO<sub>2</sub> Smart Grid much less compression is needed. Compression energy has been estimated based on the operational conditions of a compressor given in (Geological Survey of the Netherlands, 2009). Based on this source the energy needed to get from 40 bar(a) to 110 bar(a) is approximately 100 MJ / tonne CO<sub>2</sub> captured. This energy use is subtracted from the total energy use for the capture at the blast furnace.

## Emissions

The blast furnace gas from Tata Steel is used to produce energy from at the Velsen 23 plant. The gas is incinerated here, removing a number of harmful substances. However this plant currently still emits fine particulate matter, H<sub>2</sub>S and COS. When adding a CO<sub>2</sub> capture unit after the Velsen 23 plant there is a possibility that the emissions of fine particulate matter and H<sub>2</sub>S decrease.

Since no concrete data is available, however, on the exact impact of installing a CO<sub>2</sub> capture unit these possible benefits are not taken into consideration. The use of MDA as an absorbent has in the past lead to the production of aerosols. The MDEA absorbent is less prone to degeneration than MDA is but the exact emissions because of the use of this absorbent are unknown.

## Fossil oil refining

The description of the CO<sub>2</sub> capture plant at the hydrogen facility has been obtained from (IEA, 2017). This report describes the energy balance for a conventional hydrogen facility as well as the energy balance of the plant with several different CO<sub>2</sub> capture technologies. We have calculated the difference between the conventional hydrogen plant and the plant using a cryogenic capture technology (including membranes) as described in Case 2B to get to an energy consumption per tonne CO<sub>2</sub> captured (see Table 13).

**Table 13 - Net inputs for CO<sub>2</sub> capture and preparation for injection into the Smart CO<sub>2</sub> grid at a hydrogen plant**

	Conventional hydrogen plant	Hydrogen plant with capture	Consumption for capture	Net per tonne CO <sub>2</sub> captured
Electricity to grid	9.9 MWh	0.3 MWh	9.6 MWh	0.22 MWh
Natural gas consumption (Feedstock)	1,219.7 GJ/h	1,219.7 GJ/h	0 GJ/h	0 GJ/h
Natural gas consumption (Fuel)	201.4 GJ/h	198.3 GJ/h	3.2 GJ/h	- 0.075 GJ
CO <sub>2</sub> captured	0 tonne/h	42.89 tonne/h	-	

Note: Conventional hydrogen plant based on the base case and hydrogen plant based on case 2B from (IEA, 2017). Figures might not add up due to rounding.

The capture technology as described by (IEA, 2017) produces CO<sub>2</sub> with a purity of 99.64% at a pressure of 110 bar(a). Other components in the CO<sub>2</sub> stream include 0.27 vol% CH<sub>4</sub> and 0.07 vol% CO. To be able to meet the specifications of 99.9% vol% for the CO<sub>2</sub> Smart Grid a further treatment step is needed. Further purification would naturally happen with the CATOX technology in which the CO<sub>2</sub> stream is combined with O<sub>2</sub> along a catalytic bed. No energy is needed for this process. High purity O<sub>2</sub> is needed, but only a small amount per tonne of CO<sub>2</sub>. Therefore the production of O<sub>2</sub> is disregarded.

Compression energy has been estimated based on the operational conditions of a compressor given in (Geological Survey of the Netherlands, 2009). Based on this source the energy needed to get from 40 bar(a) to 110 bar(a) is approximately 100 MJe/tonne CO<sub>2</sub> captured. This energy use is subtracted from the total energy use for the capture at the hydrogen plant.

The emissions from a conventional hydrogen plant after fossil oil refining are limited to water vapour and CO<sub>2</sub>. When capturing the CO<sub>2</sub> in a CO<sub>2</sub> capture plant therefore no other emissions are captured in the process.



## A.2 CO<sub>2</sub> utilization

### Horticulture

As explained previously, the reference case for the utilization of captured CO<sub>2</sub> in horticulture is using a gas burner for the generation of heat and CO<sub>2</sub>. Little quantitative information on the reduction of CO<sub>2</sub> emissions through the delivery of captured CO<sub>2</sub> in horticulture is available. In a previous (confidential) study by CE Delft, the quantity of gas used exclusively for the production of CO<sub>2</sub> is said to be 7 m<sup>3</sup> gas/m<sup>2</sup>. Using a CO<sub>2</sub> emission factor of 2.04 kg CO<sub>2</sub>/m<sup>3</sup> for the incineration of natural gas, this amounts to 14.28 kg CO<sub>2</sub> emission/m<sup>3</sup>. This is equal to 490 m<sup>3</sup> gas avoided per tonne CO<sub>2</sub> added to the greenhouse.

This quantity is specific for the addition of CO<sub>2</sub> during the summer period. Over the entire year the same confidential study by CE Delft shows that in the OCAP-pipeline when one tonne of CO<sub>2</sub> is added to a greenhouse approximately 0.5 a tonne of CO<sub>2</sub> from natural gas burners are avoided. We show both of these cases.

**Table 14 - Inventory horticulture case**

	Amount	Reference
Gas use of greenhouses (reference case)	7 m <sup>3</sup> /m <sup>2</sup> of greenhouse space.	(CE Delft, 2017); confirmed by LTO glaskracht/OCAP
Avoided burning of natural gas (summer)	490 m <sup>3</sup> /tonne CO <sub>2</sub> added to greenhouse	(CE Delft, 2017); confirmed by LTO glaskracht/OCAP;
Avoided burning of natural gas (OCAP)	245 m <sup>3</sup> /tonne CO <sub>2</sub> added to greenhouse	(CE Delft, 2017)

### Mineralisation

For the utilization of the captured CO<sub>2</sub> in mineralisation in Compensatiesteen, the developer RuwBouw Group was contacted. RuwBouw Group provided data on the electricity use, the amount of Compensatiesteen per tonne CO<sub>2</sub> in, the input of stainless steel slags, and the amount of avoided production of sand-lime brick.

**Table 15 - Inventory mineralisation case**

	Amount/tonne CO <sub>2</sub> in	Reference
Electricity	82.05 kWh	Interview with developer
Compensatiesteen	2.05 m <sup>3</sup> (= 4 tonne)	Interview with developer
Input of stainless steel slag (max.)	3.75 ton	Interview with developer
Avoided production of sand-lime brick (max.)	3.75 ton	Interview with developer

The avoided sand-lime brick is modelled as German sand lime brick production. Electricity used in the process is assumed to be medium-voltage.

No data has been obtained on the cleaning of the stainless steel slags before utilization in the Compensatiesteen process.

## Methanol

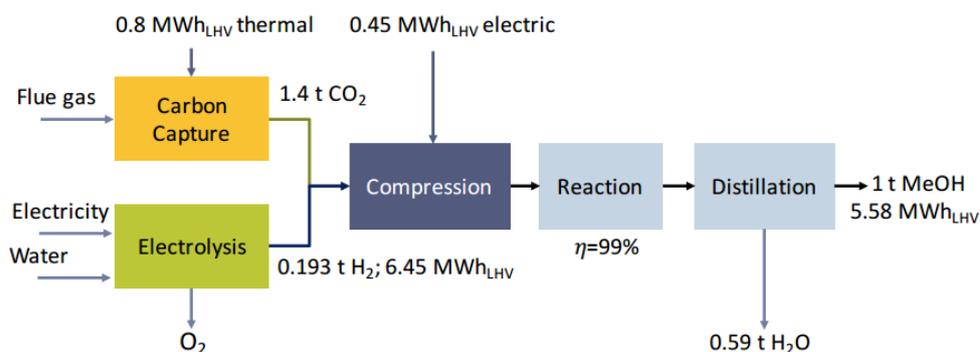
Data on the production of methanol using captured CO<sub>2</sub> was retrieved from a mass energy balance as presented in (Stefansson, 2015). Figure 18 shows the readily available data on the production process including the ratios of weight between raw materials and desired products. Additionally, data from (Rocha, et al., 2017) was used for the electricity use of the electrolysis step.

**Table 16 - Inventory methanol case**

	Amount	Reference
Electricity hydrogenation CO <sub>2</sub>	0.32 MWh/tonne CO <sub>2</sub> in	(Stefansson, 2015)
Electricity electrolysis H <sub>2</sub>	51.20 MWh/tonne H <sub>2</sub> produced	(Rocha, et al., 2017)
Hydrogen (in)	0.14 tonne/tonne CO <sub>2</sub> in	(Stefansson, 2015)
Water (out)	0.42 tonne/tonne CO <sub>2</sub> in	(Stefansson, 2015)
Methanol (out)	0.71 tonne/tonne CO <sub>2</sub> in	(Stefansson, 2015)

The hydrogen used in the process is assumed to be produced through chlor-alkali electrolysis, using a diaphragm cell. For grey electricity, medium voltage Dutch average electricity is used. For the sensitivity case, in which green electricity is used, this is assumed to be derived from a >3MW onshore wind turbine.

**Figure 18 - Mass balance and energy balance for CRI CO<sub>2</sub> to methanol technology**



Bron: (Stefansson, 2015).

### A.3 Carbon Capture and Storage (CCS)

The reference technology of CCS is mainly based on data retrieved from (Koornneef, et al., 2008). Furthermore extra compression from 40 to 130 bar(a) is determined based on (Geological Survey of the Netherlands, 2009). This case is kept very simple, and no infrastructure is taken into account.

**Table 17 - Inventory CCS**

	Amount/tonne CO <sub>2</sub> in	Reference
Electricity for compression from 40 bar(a) to 130 bar(a)	100 MJe	(Geological Survey of the Netherlands, 2009)
Electricity for injection - compression energy	7 kWh	(Koornneef, et al., 2008)
Fugitive CO <sub>2</sub> emissions from compressor	0.0003	(Koornneef, et al., 2008)