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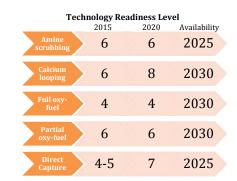
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1 Carbon capture in the cement industry: technologies, progress and retrofitting

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9 **TOC**



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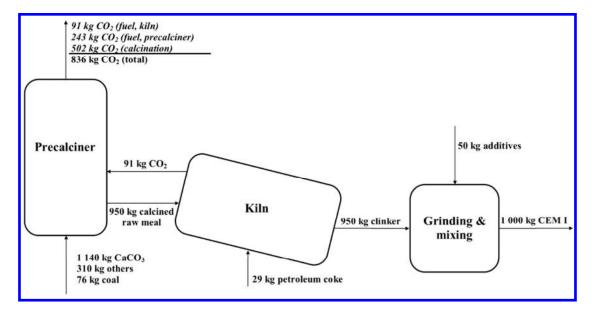
11 Abstract

Several different carbon capture technologies have been proposed for use in the cement industry. This paper reviews their attributes, the progress that has been made towards their commercialisation and the major challenges facing their retrofitting to existing cement plants. A Technology Readiness Level (TRL) scale for carbon capture in the cement industry is developed. For application at cement plants, partial oxy-fuel combustion, amine scrubbing and calcium looping are the most developed (TRL 6 = pilot system demonstrated in relevant environment), followed by direct capture (4 – 5 = component/system validation at lab-scale in relevant environment) and full oxy-fuel combustion (4 = component/system validation at lab-scale

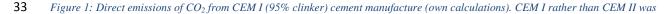
- 19 in lab environment). Our review suggests that advancing to TRL 7 (demonstration in plant environment)
- seems to be a challenge for the industry, representing a major step up from TRL 6.
- 21 The important attributes that a cement plant must have in order to be 'carbon capture ready' for each capture
- 22 technology selection is evaluated. Common requirements are space around the preheater/precalciner section,
- 23 access to CO_2 transport infrastructure and a retrofittable preheater tower. Evidence from the electricity
- 24 generation sector suggests that carbon capture readiness is not always cost-effective. The similar durations of
- 25 cement plant renovation and capture plant construction suggests that synchronising these two actions may
- save considerable time and money.

27 Introduction

- 28 Unlike most industrial processes, almost two-thirds (64%) of the CO₂ emissions emanating from the Portland
- 29 cement industry come from process chemistry rather than from fuel combustion¹. As shown in Figure 1,
- around 880 kg CO_2 is generated per tonne of clinker in a typical (1 Mtpa, 3 000 tpd) cement plant², which
- 31 produces CEM I (95% clinker).



32



34 chosen for comparisons in this paper because of its smaller range of composition than CEM II (95 – 100% clinker by weight versus 35

35 – *94%*).

36 The cement industry is likely to play a role in reducing greenhouse gas emissions to combat anthropogenic 37 climate change. Many decarbonisation pathways suggest that direct specific emission levels of around 350 – 410 kg CO₂/t cement will be required^{1,3}. However, increasing clinker substitution, alternative fuel use and 38 thermal energy efficiency¹ can only lead to specific emissions per tonne of cement falling from 730 kg CO_2/t 39 40 cement in 2009 to about $540 - 590 \text{ kg CO}_2/t$ cement in 2050. Alternative, lower CO₂-intensity cements have been suggested but uptake is not expected to be anywhere near the levels required if the sector is to meet these 41 42 targets⁴. Many NGO-based analysts, such as the IPCC and IEA, agree that the main technology group able to achieve the remaining required emission reductions is carbon capture and storage (CCS)^{1,5}, owing to the 43 relatively high concentration of CO₂ in the flue gas from these large, point-source emitters⁶. Estimates suggest 44 45 that the Spanish cement industry could reduce its specific direct emissions by only 21% between 2010 and 2050 without CCS⁷, and that UK cement sector absolute CO₂ emissions could be reduced by 66% in the 1990 46 to 2050 period if CCS is not available but by 81% if it is⁸. 47 48 However, none of the 45 large-scale CCS projects in design, construction or operation involves the cement

industry⁹. Most operating carbon capture plants are in natural gas processing⁹, but by 2050 seven industrial 49 sectors could account for about half of CO₂ emissions avoided by CCS¹⁰. Commercial-scale application of the 50 51 technology in the cement industry is seen by most as being five to ten years away at best and that few, if any, carbon capture plants will exist before 2030^{11-17} . Little research into the practicalities of installing the capture 52 plant at a cement plant, particularly in the case of retrofitting, has been published^{18–20}. A lack of effective 53 54 policy drivers - such as a substantial carbon price, effective strategies to address carbon leakage and promotion of access to capital – is limiting progress and impeding commercial-scale demonstration^{11,19,21}. An 55 56 estimate that failure to develop CCS for industrial applications could increase climate policy costs globally by 221 bn \in_{2013} /y by 2050¹³ illustrates the importance of the technology to the cement sector and other energy-57 58 intensive sectors.

59 This paper starts by developing a new Technology Readiness Level (TRL) methodology for carbon capture at 60 cement plants. The paper then describes the five following promising carbon capture processes: amine 61 scrubbing, calcium looping, full oxy-fuel combustion, partial oxy-fuel combustion and direct capture, before 62 assessing them according to several criteria including the TRL methodology. Based on current R&D efforts, 63 the TRL of each capture technology in 2020 and a date for commercial availability is predicted. Finally, some 64 of the changes to a cement plant required to enable construction and operation of each carbon capture

technology are identified and compared; the most important issues to take into consideration when designing a

66 cement plant which is likely to require retrofitting with CCS in the future are highlighted. It should be noted

67 that this paper focusses on carbon capture technologies, not the complete chain of capture, transport and

68 storage.

69 Evaluation of carbon capture technologies for cement plants

70 Technology Readiness Levels (TRLs)

71 TRLs are used for determining how close to operational deployment a technology is and this approach has been extensively used across CCS literature related to electricity generation^{22,23}. In Table 1, we modified 72 73 electricity generation-specific methodologies from the US Department of Energy Clean Coal Research Program²³ and the GCCSI²² to be relevant to cement manufacture. The original US DoE TRL specification 74 75 included two quantitative measures for many of the levels: the size of the process as a percentage of final size 76 of the power station, and a volumetric flow rate of flue gas. This concept has been retained. The flue gas and 77 production rates at each level are equivalent. 'Commercial-scale' is assumed to be a minimum of 1 000 tpd 78 (tonnes of clinker per day), and a demonstration cement plant is assumed to have a capacity at least 250 tpd.

79 Promising technologies for carbon capture at cement plants

80 Five promising carbon capture technologies for use at cement plants are described and discussed below. A

summary, including costs, is presented in Table 2. For comparison, global average thermal energy

consumption in 2012 was 3 530 MJ/t clinker, down from 3 750 MJ/t clinker in 2000². Average electrical

83 consumption was 74 kWh_e/t clinker and 99 kWh_e/t cement in 2012^2 . Typical investment costs for a cement

plant in Europe are $250 \notin_{2013}/(\text{tpa})^{20}$. A 3 000 tpd (1 Mtpa) cement plant produces approximately as much CO₂

85 as a 125 MW_e coal-fired power station.

86 Amine scrubbing

87 This is an end-of-pipe technology; it only involves the flue gas and so does not directly affect the cement
88 manufacture process except, for example, energy management strategies and start up and shut down

89 procedures. Capture rates are expected to be $\ge 90\%^{24}$ but some studies have examined lower rates¹⁸.

The thermal energy demand of amine scrubbing is very high (at least 2 GJ/t CO_2)²⁴ and it generally has to be provided via CHP and/or waste heat recovery. Owing to the paucity of low-grade heat at most cement plants, it may be significantly cheaper to capture only a proportion (up to 50%) of the CO₂ from the plants and not invest in extra heat generation capacity²⁵. Furthermore, the flue gas clean-up required increases plant footprint and capital and operating costs²⁶. As with all capture technologies, there are knock-on environmental effects from using amine scrubbing²⁷.

- 96 With respect to electricity generation, the technology is at TRL $8 9^{21}$. For cement production, the pilot plant 97 in Brevik, Norway is the most developed, and with a flue gas flow rate of approximately 125 L/s its TRL is 5 98 $- 6^{28}$. We are not aware of any plans for larger-scale pilot projects in the short- to medium-term. A preliminary 99 estimate of commercial availability is 2025 – 2030, significantly later than the IEA's estimate of 2020¹.
- 100 Full oxy-fuel combustion

101 Oxy-fuel uses a mixture of oxygen (separated from air) and recycled CO_2 as the combustion gas, reducing the 102 CO_2 separation plant's complexity and size²⁰. The capture rate is expected to be > 90%²⁹.

103 Although energy efficiency³⁰ and clinker throughput³¹ are expected to improve in an oxy-fuel cement plant, an

air separation unit (ASU) using up to 60 kWh_e/t clinker is required to produce pure oxygen for the process²⁹.

105 Alternative processes for oxygen production are being developed which could reduce the energy penalty 24 .

106 Unlike the other four technologies, full oxy-fuel combustion will affect the whole cement plant. The design of 107 virtually every unit is different from a traditional cement plant to take account of different gas properties and to minimise gas ingress or egress from the units²⁰. This is likely to be technically achievable but expensive; on 108 this basis we agree with others¹⁹ that retrofitting full oxy-fuel capture to an existing cement plant is unlikely to 109 110 be an attractive proposition. New-build full oxy-fuel cement plants are expected to cost around 220 - 290 111 112 cost alone is equivalent to 22.2 – 29.2 €/t cement. Similar numbers calculated for the other technologies are 113 given in parentheses after their capital costs.

Full oxy-fuel is seen by some²⁰ as the best technology for new-build low-carbon cement manufacture, but
development is difficult because the next stage is the construction of a whole, albeit small, cement plant. Its

116 TRL is 4 and until the ECRA's \in 50M, 500 tpd pilot plant is funded³⁴ this is not expected to increase; however,

118 organisations (such as universities) and to the authors' knowledge no company has announced any intention to

such progress could raise full oxy-fuel's TRL to 8. This step seems to be without the remit of most research

119 fund such a pilot plant in the near- or medium-term. An estimate of commercial availability is 2030 – 2040.

120 Partial oxy-fuel combustion

117

121 The difficulties with applying full oxy-fuel combustion have led to a 'partial oxy-fuel' approach where the 122 preheaters and precalciner are oxy-fuelled and the kiln and cooler are air-fuelled (i.e. conventional). It is expected that the capture rate could be as high as $70\%^{20,35}$. The preheaters and precalciner would have to be 123 124 redesigned and made gas-tight, but retrofitting is expected to be easier than for full oxy-fuel because the kiln 125 and cooler would not change. Since 75% of the fuel is burned in the precalciner it is assumed that a partial oxy-fuel ASU would require about 45 kWh_c/t clinker²⁰. A partial oxy-fuel retrofit is expected to cost around 126 85 €/(tpa)³² (8.6 €/t) whilst new-builds are expected to be in the region of 225 - 275€/(tpa)^{20,35} (22.7 - 27.7) 127 128 €/t).

A 30 – 50 tpd pilot plant has been built by a consortium including Air Liquide, FLSmidth and Lafarge, and a
feasibility and cost exercise regarding retrofitting partial oxy-fuel to a cement plant undertaken³⁶. Its TRL is
therefore 6, but without the next step of a full FEED study it is unlikely to increase soon³⁷. A preliminary
estimate of commercial availability is 2025 – 2035, similar to the IEA's estimate of 2025¹.

133 Calcium Looping (CaL)

134 Calcium looping (CaL) involves chemical reactions between CO_2 and calcium oxide sorbent in a pair of

135 circulating fluidised beds. There are energetic and waste benefits that can be achieved by integrating CaL with

136 cement manufacture from using CaCO₃ as a sorbent precursor and operating at > $600^{\circ}C^{38}$. High-grade 'waste

137 heat' from the process can be used to generate additional electricity; this should be about the same as the

amount required by the cement, capture and CO₂ compression plants combined.

139 An ASU using about 20 kWh_e/t clinker would be required to produce oxygen for the calciner. Fuel

140 consumption would increase by about 50% but the CO₂ avoidance rate is expected to be $\ge 90\%^{35}$. The

141 preheaters would need altering to take into account the diversion of limestone from the usual raw meal entry

142 point at the first preheater to the CaL calciner; Ozcan et al.³⁹ assume that the waste CaO sorbent would be

mixed with the rest of the raw meal between the precalciner and kiln (the 'diversion' design). The flue gases
would flow into the CaL carbonator between the third and second preheaters. Alternatively, the CaL calciner
could replace the precalciner (the 'replacement' design)³⁵. Another rather different design ('HECLOT', by
ITRI) uses a rotary kiln calciner; this could encounter the same issues surrounding gas-tightness as full oxyfuel combustion^{40,41}.

- 148 The largest project so far is HECLOT in Taiwan, which captures 1 tCO₂/h from 3.1 t/h flue gas using the
- rotary kiln calciner⁴⁰. Thus, CaL in the cement industry is at TRL 6. ITRI is planning to build a larger plant in
- 150 2017 which, if successful, will raise the TRL to 8^{40} . There are no known plans to build a cement-based CaL
- 151 pilot plant with a fluidised bed calciner. A preliminary estimate of commercial availability is 2025 2030.

152 *Direct Capture*

Direct capture only captures emissions coming from the calcination of limestone, which account for about 64% of the CO_2 generated at a typical cement plant¹. This process is being developed by Calix, an Australian company. Most of the information in this section comes from the company directly, via the website and from discussions with employees⁴².

Direct capture occurs in a vertical shell-and-tube heat exchanger known as a direct capture unit (DCU). Raw meal and steam pass down the tubes and are heated and calcined by heat transferred from flue gases from a combustion process flowing through the shell. Because no external gases enter the tubes, the gas coming out of them is a virtually pure CO_2 /steam mix. After steam knock-out, the CO_2 should be suitable for compression⁴².

The DCU will replace the precalciner and receive hot raw meal from the preheaters. Modelling by Calix suggests that the energy penalty after heat integration will be $\pm 2\%$ of the thermal energy requirement of the cement plant⁴³. Retrofitting should be relatively easy because it requires the replacement of only the preheaters and precalciner.

A pilot plant has operated with an equivalent capacity of 160 tpd clinker. The lack of information about theimpurities in the raw meal limits its TRL to 4. Calix is planning to build a 320 tpd pilot plant at a European

168 cement plant before 2020 and successful operation will raise the TRL to 7. A preliminary estimate of

169 commercial availability is 2025 - 2030.

170 Prospects for further development and technology champions

If there were great pressure to commercialise cement CCS as soon as possible, amine scrubbing would likely
be the first available, but the lack of such pressure offers other technologies the chance to catch up. Amine
scrubbing's main problem is its cost (see Table 2); a cheaper alternative at a similar level of development
would stand a good chance of supplanting it. However, no technology is likely to be widely available before
2025.

176 Direct Capture and Calcium Looping seem to be progressing fastest and possibly could reach TRL 7 by 2020;

177 no other technology is expected to reach this level soon although partial oxy-fuel combustion could overtake

178 them if the AL/Lafarge/FLS consortium decides to progress with trials.

Scale-up can require significant investment; the six-tenths 'rule'⁴⁵ suggests that increasing the scale of a
process by an order of magnitude will quadruple capital investment costs. Building the confidence of potential
investors or developers is critical for carbon capture projects because most of the technologies are developed
by a sequence of organisations on the path to commercialisation.

183 In this context, TRL 7 seems to be the major obstacle for capture processes in the cement industry. This may 184 be because it is the point at which traditional university-led research is too small-scale to develop the 185 technology further. Companies or larger research institutions acting as a 'champion' for a specific technology 186 are generally more suited to carry on development beyond TRL 6. Such organisations are Calix (direct 187 capture) and ITRI (calcium looping). The ECRA, as a research collaboration of several cement manufacturers, 188 does not necessarily have the independence and resources to develop a pilot-scale oxy-fuel plant. Although the 189 AL/Lafarge/FLS consortium (partial oxy-fuel) would appear to have massive financial and technological 190 resources, it is likely that limited funds and scope prevent it from continuing development. Amine scrubbing 191 has many champions but whether much of their focus is on the cement industry is debatable. The absence of 192 commercial reasons to invest in a decade-long development & demonstration programme makes TRL 7 193 virtually impossible for technologies currently championed by universities and small research institutes.

Of particular interest to plant owners may be technologies that can be installed, if not operated, at a low extra cost. Designing a process to be easily convertible to partial oxy-fuel (e.g. more air-tight preheaters) may help to reduce costs in the long-run; this is discussed below. Furthermore, direct capture theoretically offers 50 – 60% capture for very little added cost for new-builds. It is possible that such a plant could be built and run competitively until the rest of the CCS chain is available.

199 Technologies such as amine scrubbing, which are already in use in other industries, have the benefit of

200 learning within those industries as well as design and equipment suppliers with relevant experience. Oxy-fuel

201 systems should not suffer too much in this respect; oxygen production is similar across industries and although

202 changes to all major process units are required, these should be well within the competencies of cement plant

203 manufacturers. Direct capture and CaL are quite process-specific so are unlikely to benefit in this respect.

204 Early indications are that retrofitting a cement plant with some form of carbon capture (except amine

scrubbing) will have a capital cost in the region of 100 €/(tpa) (10.1 €/t) compared with a reference new-build

cement plant cost of approximately 250 €/(tpa)⁴⁶ (25.2 €/t). A new-build cement plant with carbon capture is

207 expected to cost in the region of 300 €/(tpa) (30.3 €/t). Costs of CO₂ avoided are around 20 – 80 €/t CO₂, again

208 excepting amine scrubbing. It is more difficult to gain a clear picture here because of the different discount

rates used across the literature which range from 6% to 16% but tend to cluster around the 8-10%

210 region^{23,24,35,47}.

211 The range of capital costs for amine scrubbing varies wildly, and this is in part due to assumptions about the

source of the extra energy for stripping the CO_2 from the solvent^{18,32}. Most studies focus on MEA

solvent^{18,20,30,48}; it is more likely that more advanced amines would be used, reducing both the capital and
operating costs.

Any capture process must allow the cement plant to continue to produce in-spec cement. Amine scrubbing should not have a significant effect beyond affecting the energy management on site unless waste heat recovery is installed on the kiln. Cycling calcium oxide (or all the raw meal) through a calcium looping system will affect the physical properties of the solids, something which could have an effect on cement quality and is currently being studied batch-wise in laboratories. Direct capture's DCU could also have an effect on the properties of the calcined raw meal, and the pilot plant planned for construction by 2017 should

221 produce relevant data to evaluate possible effects. In-spec cement was created during full oxy-fuel laboratory

studies. It can be expected that by 2020 the quality of cement made in a plant with any of these capture

223 process attached will have been tested and hopefully confirmed to be within relevant standards such as EN

224 197⁴⁹.

225 Retrofitting cement plants with carbon capture technology

At some point it may be necessary to attach carbon capture facilities to an existing cement plant, a process known as retrofitting. The IEA assumes that the retrofitting of existing point-source emitters with carbon capture is likely to be necessary from 2020 in order to reach emission targets³. Retrofitting is generally seen to be more difficult and expensive than applying CCS to new-builds because there may be issues surrounding access, plant footprint and management of fuels and other resources. The plant must also be shut down for the installation of the new equipment. Only a few sources in the literature have discussed these issues³². A contribution to this topic is provided below.

233 Shutdown time

Fixed costs represent approximately 40% of total costs of operation⁵⁰ so closing down a plant for an extended
period leads to significant financial repercussions. Any overruns in construction and commissioning would
add yet more costs, with fixed costs alone being in the order of €3M per month for a typical 1 Mtpa clinker
plant⁵¹.

The first significant retrofit of a power station with CCS was of Boundary Dam Unit 3 with amine scrubbing,
which started operation in October 2014. Putting aside the testing and commissioning time, the construction
took thirteen months although it should be noted that the power station unit was refurbished at the same
time⁵².

242 Cement plants undergo various shutdowns for repairs, maintenance and improvement. These range from short

annual shutdowns of around a month to longer shutdowns performed maybe once in a generation;

244 modernisation of complete plants can take more than a year. This can be compared with the construction of a

new cement plant, which takes around 18 - 24 months. (These durations come from promotional material, so

cannot be assumed to be representative of the industry as a whole 5^{53} .)

Thus, the time periods for refurbishment of cement plants and installation of carbon capture at power stations are similar. This suggests that applying carbon capture during a cement plant refurbishment may be the most convenient strategy, in a manner similar to Boundary Dam Unit 3. Changes to virtually all process units will mean the shutdown period for full oxy-fuel combustion is likely to be long. By contrast, connecting a preconstructed amine scrubbing plant to the preheater exhaust may be possible within the period of an annual shutdown (about a month). The other technologies will likely fall somewhere in between.

253 Carbon Capture Readiness (CCR)

The length (and cost) of shutdown periods for installation of the different technologies may ultimately become a major determinant of which of them, if any, are competitive. A way to reduce this time and expense could be by designing the cement plant to be 'carbon capture ready' from the outset. Although CCS is not currently viable in the cement sector, plant owners may wish to ensure that they can install it with minimal disturbance once it is. Alterations to the original design of the site and the cement plant itself to make them CCR could reduce time and cost during retrofitting for a small up-front investment.

Published work on CCR in the cement sector has focussed on amine scrubbing. Liang & Li¹⁸ provide a list of 260 261 21 criteria split into six categories for assessing the potential to retrofit cement plants with amine scrubbing: 262 extra space on site, access to storage capacity, water supply, sufficient electricity & steam, cement production technology and flue gas properties. The IEA GHG³⁰ states that the four main requirements for amine 263 264 scrubbing retrofitting are land, electricity import, steam production and removal of certain gases from the flue 265 gas. The first is simple to understand – the new units require space – but this may not be so easy in practice, as 266 cement plants are often surrounded by land which is unsuitable or that belongs to another entity. Electricity 267 can either be imported from the grid or produced on site, but again this will require space and/or money. 268 Amine scrubbing requires low concentrations of NO_2 , SO_2 and O_2 in the flue gas so a pre-treatment stage will 269 be necessary; this is not an insurmountable challenge.

To better understand the requirements of each technology for CCR, the changes to each relevant unit in the cement manufacturing process are compared in Table 3. Some site-wide considerations, and those concerning new units, have also been identified. The preheaters usually need to be replaced because they will have to

273 handle a gas mix with different properties (full oxy-fuel) and/or a different mass flow rate (CaL, direct capture

274 & partial oxy-fuel). Oxy-fuel systems also require more air-tight units. Attaching amine scrubbing could

change the operating conditions of the preheaters because a large enough pressure gradient will be required to

ensure the gases flow from the preheaters to the capture plant. Preheaters at a 'diversion' design CaL plant

will require tie-in locations where the gases can be diverted to the capture plant and back again.

278 The precalciner will require changes in all cases except amine scrubbing and 'diversion' calcium looping; in a

279 'replacement' design it will be replaced by the CaL calciner. In direct capture the precalciner will be replaced

280 with the direct capture unit (DCU) which will require a larger area and a new raw meal conveyance system

between the preheaters and DCU may be required. In oxy-fuel combustion, the design of the precalciner will

need to change slightly to take into account the altered gas and flame properties but it should be possible to fit

it in roughly the same area as an air-fuel precalciner.

The kiln and coolers will only require alterations in full oxy-fuel, and in this case full replacement is likely to be the most practical option, with new, air-tight designs being installed. A two-stage cooler will be required, in which the first stage uses recycled CO_2 and the second stage air to cool the clinker²⁰.

Since none of the carbon capture technologies is yet available, cement plant owners may not wish to invest in CCR based on one technology. However, there are several common requirements across all or most of the technologies. By identifying these and considering whether they merit investment up front, the plant owner can reduce retrofitting costs without locking himself in to one technology. Some major considerations for each technology are shown in Table 3 and the ones in common are discussed below.

292 Critical issues for CCR

276

The availability of land for expansion is already a concern at many sites and may be the factor which prevents or delays roll-out of CCS at some of them. Plant layout is related to this issue; all capture technologies require space at specific locations around the cement plant so ensuring that existing units do not have to be moved a few metres to make room for others could greatly reduce shut-down time. Setting aside space solely to facilitate easier construction and access on-site during retrofitting could also reduce shut-down costs. In all cases, a CO_2 compression and temporary storage facility will require space. In general, relatively large zones should be reserved for the capture plant close to the preheater tower and precalciner/kiln connection.

300 Cement plants tend to be located on limestone deposits; although some researchers have suggested that plants 301 are built within the region of a CCS cluster¹⁹, it is unlikely that this will happen except where the cluster is 302 located upon a suitable geological formation. Limestone is not suitable for CO_2 storage so there is likely to be 303 a need for significant and reliable CO_2 transport between plant and storage site. Purchasing, or having an option to purchase, the storage capacity is also extremely important³². Discussions with local authorities on 304 305 planning applications for capture plants and CO₂ pipelines at the time of cement plant construction could 306 increase the chance that the plant and pipeline can be built when required. These issues are not unique to the 307 cement sector and so are not discussed in more detail here.

308 Other important issues for CCR

309 Some items may be relatively cheap to construct when building the original cement plant, but difficult or 310 expensive to alter later on. For example, if some or all of the major pipe-runs for the capture plant are installed 311 at the same time as those for the cement plant itself, fewer changes are likely to be required later and perhaps a 312 shorter shut-down will be possible. Several of the technologies would benefit from the preheater tower being 313 adaptable to house the new preheaters and/or precalciner. However, care should be taken in choosing to apply CCR without assessment of the benefits. For example, Bohm et al⁵⁴ determined that CCR costing 4% of the 314 315 total cost of the plant made little difference to the economics of IGCC power stations. Lucquiaud et al. suggest that making a pulverised coal power station CCR could cost less than 1% of capital costs⁵⁵, and Liang et al. 316 317 determine that such power stations in China are up to 10% less likely to close early⁵⁶. Rohlfs & Madlener 318 calculated that it was usually more cost-effective to close a modern, unabated power station and replace it with a completely new abated power station⁵⁷. Discounted cash-flow analysis can identify whether the extra capital 319 320 expenditure for particular items is financially attractive or more extensive rebuilding or replacement at a later 321 date is more suitable. This is not applicable for some particular items such as land – if the plant does not have 322 room to build the capture facilities on existing land or expand into adjacent areas, the capture plant may never 323 be built regardless of the profitability.

In conclusion, carbon capture in the cement is several years away but timely consideration of the challenges which lie ahead, such as retrofitting and ensuring cement plant/capture plant compatibility, will reduce their complexity in the long run. The lack of large-scale (> 50 tpd) pilot plants in the cement industry is currently the biggest impediment to further capture technology development and commercialisation.

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332 Table 1: Technology Readiness Levels for CCS in the cement industry

Basic principles observed and reported	Lowest level of technology readiness. Scientific research begins to be translated into applied R&D. Examples include
basic principles observed and reported	
	desktop studies of a technology's basic properties.
Fechnology concept and/or application	Invention begins. Once basic principles are observed, practical applications can be invented. Applications are speculative
ormulated	and there may be no proof or detailed analysis to support the assumptions. Examples are still limited to analytic studies.
Analytical and experimental critical	Active R&D is initiated. This includes analytical and laboratory-scale studies to physically validate the analytical
function and/or characteristic proof of	predictions of separate elements of the technology (e.g., individual technology components have undergone laboratory-
concept	scale testing using bottled gases to simulate major flue gas species at a scale of < 0.5 L/s, and simulated raw materials).
Component and/or system validation in	A bench-scale prototype has been developed and validated in the laboratory environment. Prototype is defined as < 1 tpd
a laboratory environment	(e.g., complete technology process has undergone bench-scale testing using synthetic flue gas composition at a scale of <
	20 L/s, and simulated raw materials).
Laboratory-scale similar-system	The basic technological components are integrated so that the system configuration is similar to (matches) the final
validation in a relevant environment	application in almost all respects. Prototype is defined as < 1 tpd clinker scale (e.g., complete technology has undergone
	testing using actual flue gas composition at a scale of < 20 L/s and actual raw materials).
Engineering/pilot-scale prototypical	Engineering-scale models or prototypes are tested in a relevant environment. Pilot or process-development-unit scale is
ystem demonstrated in a relevant	defined as 1 – 50 tpd (e.g., complete technology has undergone small pilot-scale testing using actual flue gas
environment	composition at a scale equivalent to $0.04 - 1 \text{ Nm}^3$ /s and actual raw materials).
	rmulated nalytical and experimental critical naction and/or characteristic proof of oncept omponent and/or system validation in laboratory environment aboratory-scale similar-system lidation in a relevant environment ngineering/pilot-scale prototypical stem demonstrated in a relevant

7	System prototype demonstrated in a	This represents a major step up from TRL 6, requiring demonstration of an actual system prototype in a relevant				
	plant environment	environment. Final design is virtually complete. Pilot or process-development-unit demonstration of a 50 – 250 tpd				
		clinker scale (e.g., complete technology has undergone large pilot-scale testing using actual flue gas composition at a				
		scale equivalent to approximately $1 - 4.5 \text{ Nm}^3$ /s and actual raw materials).				
8	Actual system completed and qualified	The technology has been proven to work in its final form and under expected conditions. In almost all cases, this TRL				
	through test and demonstration in a	represents the end of true system development. Examples include start-up, testing, and evaluation of the system within a				
	plant environment	\geq 250 tpd plant with CCS operation (e.g., complete and fully integrated technology has been initiated at full-scale				
		demonstration including start-up, testing, and evaluation of the system using actual flue gas composition at a scale				
		equivalent to $\ge 4.5 \text{ Nm}^3$ and actual raw materials).				
9	Actual system operated over the full	The technology is in its final form and operated under the full range of operating conditions. The scale of this technology				
	range of expected conditions	is expected to be \geq 1000 tpd plant with CCS operations (e.g., complete and fully integrated technology has undergone				
		full-scale demonstration testing using actual flue gas composition at a scale equivalent to \geq 18 Nm ³ and actual raw				
		materials).				

Table 2: CO₂ capture from the cement industry: technology comparisons

Attribute	Amine scrubbing*	Calcium looping	Full oxy-fuel	Partial oxy-fuel	Direct capture
Capital cost (€2013)	213 M for 2 Mtpa RF	269 M NB (inc cement plant	291 M for 1 Mtpa NB ³²	97 – 107 M for 1 Mtpa RF ³⁵	Unknown.
	(China) ¹⁸	cost) for 1 Mtpa ³⁵	104 M for 1 Mtpa RF ³²	85 M for 1 Mtpa RF ³²	
	440 – 540 for 1 Mtpa NB ³²	125 M NB (capture plant		275 M for 1 Mtpa NB ³²	
	245 – 350 for 1 Mtpa RF ³²	only) for 1 Mtpa ³⁵			

Overall cost, avoided	46 - 57 NB @ DR $6 - 16$ % ¹⁸	75 – 85 RF @ DR 10 % ¹⁵	39 NB @ DR 8 % ³²	49 NB @ DR 8 % ³²	Unknown
(€ ₂₀₁₃ /t CO ₂)	51 NB @ DR 7 $\%^{48}$	18 NB ³⁵	41 RF @ DR 8 % ³²	54 RF @ DR 8 % ³²	
	107 NB @ DR 10 $\%^{58}$	31 NB ⁵⁹		12 NB ³⁵	
	52 – 104 @ DR 8 % ³²			$54 - 69 \ \mathrm{RF}^{37}$	
	143 – 187 RF @ DR 10 % ¹⁵			58 RF ³²	
	172 – 333 (short-term)			62 RF ³⁶	
	86 (long-term) ⁴⁷				
	53 RF ¹⁸				
Typical capture rate	> 90 %	> 90 %	> 90 %	65 %	60 %
Complexity	Low: mature end-of-pipe	Medium: integration should	High: Increased design	Medium: Increased design and	Low: Operational knowledge of
	technology, but extensive FG	be simple but fluidised bed	and maintenance	maintenance complexity	direct capture in cement industry
	clean-up is required before	combustor operation is outside	complexity; operation of	(although less than full oxy-	currently non-existent except for
	capture	cement industry knowledge	the plant changes,	fuel); operation of the plant	one company but kiln/cooler
			especially in kiln and	should be relatively similar to	section identical to before.
			cooler. Kiln stop likely if	unabated cement	
			O ₂ supply fails.		
Major changes to	None	Precalciner replaced with dual	New preheaters and	New preheaters and precalciner	Precalciner replaced with direct
cement process		fluidised beds (or, for	precalciner necessary.	necessary.	capture unit (DCU) tower.

		HECLOT, one fluidised bed	Changes to kiln burner		
		and a rotary kiln), steam cycle	and cooler designs		
		and associated equipment	necessary. False air flow		
			reduction requires altered		
			designs of units		
Capture plant	Large because of installation	Possibly slightly larger than	Relatively large - air	Medium (0.5 ha) – air	Small. DCU tower likely to be
footprint	of SCR & FGD systems as	partial oxy-fuel but smaller	separation, waste heat	separation, waste heat recovery,	shorter but wider than a preheater
	well as capture plant ²⁶	than full oxy-fuel. CO ₂	recovery and CO ₂	FG recycling and CO ₂	tower; gas treatment plant will be
		processing unit required to	processing units will take	processing units will take up	small due to low capture rate and
		remove chlorides & water. A	up space.	space, but lower capture rate	inherent purity of CO ₂ (only
		steam cycle will need to be		and O_2 demand means they will	water removal necessary)
		installed.		be smaller than full oxy-fuel	
Cement quality	No change expected	No change observed at lab	No change observed at lab	No change observed at lab scale	Unknown
		scale	scale		
Retrofittability	Easy, since few changes to the	'Diversion' and	Technically possible but	Relatively easy. Precalciner and	Relatively easy. Probably similar
	cement plant itself are	'Replacement' designs:	doubts about practicality	preheater replacement will	to partial oxy-fuel as both require
	required. Physical connection	Possible, but prolonged	remain. Long shutdown	require a lengthy shutdown, but	preheater & precalciner
	to cement plant probably	shutdown likely while dual	expected for installation	length (and risks) not as great as	replacement. Modular nature of
	possible in annual shutdown	FBCs installed. Space may be	of new equipment and	for full oxy-fuel.	capture technology should enable

	period. Space for capture plant	a constraint.	alteration of existing units		some prefabrication and reduce
	may be an issue on many sites.	'HECLOT': replacement of			construction times on site
		kiln will cause a long			
		shutdown. As with full oxy-			
		fuel, practicality of gas-tight			
		rotary kilns must be			
		demonstrated			
Current Technology	6	6	4	6	4-5
Readiness Level	0.125 Nm ³ /s real FG	3.1 tph FG (0.7 Nm ³ /s FG)	Lab-scale tests	2 – 3 tph RM (1.3 – 2 tph) pilot	One-tube (10 tph RM, 6.6 tph/160
(TRL) with respect to	scrubbed ²⁸ (ca. 0.2% of full	HECLOT PP in operation in	undertaken, but no PP	plant in Denmark operated	tpd) tests undertaken, but not at a
cement manufacture	size)	Taiwan ⁵⁹ but results not yet	built yet ²⁰	successfully ³⁷	cement plant with only with high-
		published (1.2% of full size)			purity RM. Heat integration not
					tested ⁴³ .
TRL expected in	6	8	4	6	7
2020 assuming	No new amine scrubbing PP	ITRI plans to build a 30 MW_t	ECRA plans to build a 2	Consortium not progressing	20 tph RM (ca. 13 tph/320 tpd
successful completion	projects in cement sector are	(11 Nm ³ /s, 20% of full size)	tph PP seem to be on hold	with FEED because of lack of	clinker, 10 % of full size) PP to
of current plans	currently known	HECLOT PP in 2017 ⁵⁹	so unlikely to be	viable business model ³⁷	be built in 2018 – 2020.
			completed by 2020		
Time until wide	10 – 15 years	10 – 15 years	15 – 25 years	10 – 20 years	10 – 15 years
availability					

334

- 335 $\mathbf{RF} = \text{retrofit. Includes only cost of capture plant. NB} = \text{New-build. Includes cost of cement plant (usually about 150 M<math>\in$ in Europe). $\mathbf{DR} = \text{discount rate. RM} = \text{raw meal. FG}$
- = flue gas. **PP** = pilot plant. **Full size** = 3 000 tpd clinker (1 Mtpa), or 55 Nm^3/s flue gas. *Includes the cost of CHP for heat provision.

337

338

Table 3: Technology-specific considerations for designing capture-ready cement plants

Aspect of	Amine scrubbing	Calcium looping	Direct capture		Oxy-fuel
plant				Partial	Full
Raw materials	If a CHP plant is to be built,	More fuel (ca. 50%) will be required on	If necessary, a source	A larger electric	ity grid connection should be
& fuel	the fuel supply should be	site so storage & handling facilities could	of purer (i.e. low-Cl)	installed so that enot	ugh electricity can be imported to
handling;	considered. This may include	be designed to accommodate this from	raw materials should	run the ASU a	nd other capture equipment
utility	a natural gas pipeline	the start. Combustion of alternative fuels	be identified		
connections	connection.	in a CFB may be difficult so coal			
		facilities may be the most important to			
		over-size.			
	Cooling and process w	vater connections will be necessary			
Preheaters		The ability to connect the flue gas exhaust	to the gas clean-up syste	em should be included	
	The exhaust from the	The tower should be built to a specificat	ion whereby it can accom	nmodate the new desig	gn of preheaters required in the
	preheaters will go to the	capture plant.			
	FGD plant. Enough pressure	Tie-in locations for connection to the	The preheaters		
	will have to be present to let	CaL calciner should be designed and	should be at a height		
	it flow; this may affect plant	included ('diversion' design)	to allow good		

	design or require the		connection between		
	installation of an extra fan		them, the DC		
			calciner and the kiln.		
Precalciner	No action necessary.	The connections between the calciner and	Sufficient space for	The calciner he	busing design must be able to
		the kiln and preheaters should be	the larger direct	accommodat	te the post-retrofit calciner.
		appropriate for re-connection to the CaL	capture calciner is		
		calciner ('replacement' and 'HECLOT'	necessary.		
		designs)			
Kiln		No action necessary.			The kiln should be as airtight as
					possible. The region around the
					burner, including the air supply
					should be suitable for
					retrofitting with the new burner
					and gas supply. The kiln must
					be compatible with the
					refractory required for oxy-fuel
					combustion
Cooler		No action necessary.			The cooler, or at least the site of
					the cooler, should be adaptable

					for oxy-fuel operation. This
					may include building a two-
					stage cooler, which is likely to
					be larger than a standard cooler.
Plant footprint	A very large amount of land	The cement plant may require a different	A small amount of	A significant amou	nt of land will be required for an
	will be required to build the	layout to ensure that a CaL system can be	land will be required	ASU and the recir	rculation loop. Land for the gas
	capture facilities. This	fitted between the preheaters and kiln or	to accommodate the	clean-up plant shou	ld be made available close to the
	should be close to the	within the preheater train. Space for the	DCU and flash	р	reheater tower.
	preheater exhaust. The CHP	ASU and steam cycle should be provided	condenser.		
	plant should be built close by	relatively close to the CaL plant location,			
	to reduce the distance that	and gas clean-up and compression should			
	the steam has to be	not be too far away from the calciner.			
	transported				
Other	Gypsum will be produced			Purification &	
	on-site from the FGD plant;			compression plant	
	disposal or sale of this			for partial oxy-fuel	
	should be considered			plant (1 Mtpa)	
				would require 0.5	
				ha.	

340 **References**

341	(1)	Elzinga, D.; Bennett, S.; Best, D.; Burnard, K.; Cazzola, P.; D'Ambrosio, D.; Dulac, J.; Fernandez
342		Pales, A.; Hood, C.; LaFrance, M.; McCoy, S.; Mueller, S.; Munuera, L.; Poponi, D.; Remme, U.;
343		Tam, C.; West, K.; Chiavari, J.; Jun, F.; Qin, Y. Energy Technology Perspectives 2015: Mobilising
344		Innovation to Accelerate Climate Action; International Energy Agency: Paris, 2015.
345	(2)	WBCSD Cement Sustainability Initiative. "Getting the Numbers Right" (GNR)
346		http://www.wbcsdcement.org/index.php/key-issues/climate-protection/gnr-database.
347	(3)	Diczfalusy, B.; Wråke, M.; Breen, K.; Burnard, K.; Cheung, K.; Chiavari, J.; Cuenot, F.;
348		D'Ambrosio, D.; Dulac, J.; Elzinga, D.; Fulton, L.; Gawel, A.; Heinen, S.; Ito, O.; Kaneko, H.;
349		Koerner, A.; McCoy, S.; Munuera, L.; Remme, U.; Tam, C.; Trigg, T.; Trudeau, N.; Yamada, H.
350		Energy Technology Perspectives 2012: Pathways to a Clean Energy System (ETP 2012);
351		International Energy Agency: Paris, 2012.
352	(4)	Heidrich, C.; Sanjayan, J.; Berndt, M. L.; Foster, S.; Sagoe-Crentsil, K. Pathways and Barriers for
353		Acceptance and Usage of Geopolymer Concrete in Mainstream Construction; Nashville, 2015.
354	(5)	Edenhofer, O.; Pichs-Madruga, R.; Sokona, Y.; Farahani, E.; Kadner, S.; Seyboth, K.; Adler, A.;
355		Baum, I.; Brunner, S.; Eickemeier, P.; others. Climate Change 2014: Mitigation of Climate
356		Change: Contribution of Working Group III to the Fifth Assessment Report of the
357		Intergovernmental Panel on Climate Change. Camb. Univ. Press Camb. U. K. N. Y. NY USA
358		2014 , <i>1</i> .
359	(6)	Napp, T.; Gambhir, A.; Florin, N.; Fennell, P. S. Reducing CO2 Emissions from Heavy Industry: A
360		Review of Technologies and Considerations for Policy Makers; Grantham Briefing Paper 7;
361		Grantham Institute for Climate Change, Imperial College London: London, 2012.
362	(7)	García-Gusano, D.; Herrera, I.; Garraín, D.; Lechón, Y.; Cabal, H. Life Cycle Assessment of the
363		Spanish Cement Industry: Implementation of Environmental-Friendly Solutions. Clean
364		Technol. Environ. Policy 2014 , 17 (1), 59–73.

365 (8) Industrial Decarbonisation and Energy Efficiency Roadmaps to 2050 - Publications - GOV.UK 366 https://www.gov.uk/government/publications/industrial-decarbonisation-and-energy-367 efficiency-roadmaps-to-2050 (accessed Jun 5, 2015). 368 (9) Global CCS Institute. Large-Scale CCS Projects Database 369 http://www.globalccsinstitute.com/projects/large-scale-ccs-projects#map (accessed Nov 19, 370 2015). 371 (10) International Energy Agency. Technology Roadmap: Carbon Capture and Storage, 2013 372 Edition; Energy Technology Perspectives; Paris, 2013. 373 (11)International Energy Agency. Global Action to Advance Carbon Capture and Storage: A Focus 374 on Industrial Applications - Annex to Tracking Clean Energy Progress 2013; International 375 Energy Agency: Paris, 2013. 376 (12) Barker, D. Global Technology Roadmap for CCS in Industry: Sectoral Assessment: Cement; 377 UNIDO, 2010. 378 (13) Kober, T.; de Coninck, H.; Mikunda, T.; Bazilian, M. CCS in Industry: The Case for an 379 Undervalued Mitigation Option; CATO2-WP2.3-D09; CATO-2: Utrecht, Netherlands, 2012. 380 (14)Leese, R. MPA Cement GHG Reduction Strategy: Technical Document; Mineral Products 381 Association: London, 2013. 382 (15) Element Energy; Carbon Counts; PSE; Imperial College; University of Sheffield. Appendix to 383 "Demonstrating CO2 Capture in the UK Cement, Chemicals, Iron and Steel and Oil Refining 384 Sectors by 2025: A Techno-Economic Study"; UK Departments of Energy and Climate Change 385 and Business, Innovation and Skills: London, 2014. 386 (16)Harland, K.; Pershad, H.; Slater, S.; Cook, G.; Watt, J. Potential for the Application of CCS to UK 387 Industry and Natural Gas Power Generation for the Committee on Climate Change; Element 388 Energy, 2010. 389 (17)World Business Council for Sustainable Development; International Energy Agency. Cement 390 Technology Roadmap 2009: Carbon Emissions Reductions up to 2050; 2009.

391	(18)	Liang, X.; Li, J. Assessing the Value of Retrofitting Cement Plants for Carbon Capture: A Case
392		Study of a Cement Plant in Guangdong, China. Energy Convers. Manag. 2012, 64, 454–465.
393	(19)	Li, J.; Tharakan, P.; Macdonald, D.; Liang, X. Technological, Economic and Financial Prospects
394		of Carbon Dioxide Capture in the Cement Industry. <i>Energy Policy</i> 2013 , <i>61</i> , 1377–1387.
395	(20)	Hoenig, V.; Hoppe, H.; Koring, K.; Lemka, J. ECRA CCS Project – Report on Phase III; TR-ECRA-
396		119/2012; European Cement Research Academy: Duesseldorf, Germany, 2012.
397	(21)	Global CCS Institute. The Global Status of CCS: 2014; Melbourne, Australia, 2014.
398	(22)	Global CCS Institute; Electric Power Research Institute; WorleyParsons. Strategic Analysis of
399		the Global Status of Carbon Capture and Storage. Report 4: Existing Carbon Capture and
400		Storage Research and Development Networks around the World; Strategic Analysis Series; 4;
401		GCCSI: Melbourne, Australia, 2009.
402	(23)	Office of Fossil Energy. 2012 Technology Readiness Assessment - Carbon Capture, Utilization
403		and Storage (CCUS); United States Department of Energy, 2012.
404	(24)	Boot-Handford, M. E.; Abanades, J. C.; Anthony, E. J.; Blunt, M. J.; Brandani, S.; Dowell, N. M.;
405		Fernández, J. R.; Ferrari, MC.; Gross, R.; Hallett, J. P.; Haszeldine, R. S.; Heptonstall, P.;
406		Lyngfelt, A.; Makuch, Z.; Mangano, E.; Porter, R. T. J.; Pourkashanian, M.; Rochelle, G. T.;
407		Shah, N.; Yao, J. G.; Fennell, P. S. Carbon Capture and Storage Update. Energy Environ. Sci.
408		2013 , 7 (1), 130–189.
409	(25)	Graff, O. CCS in Aker Solutions with a Focus on Cement Industry, Norcem International CCS
410		Conference, Langesund Norway, May 20 – 21 2015.
411	(26)	Florin, N.; Fennell, P. S. Assessment of the Validity of "Approximate Minimum Land Footprint
412		for Some Types of CO2 Capture Plant"; United Kingdom Department for Energy and Climate
413		Change, 2010.
414	(27)	Volkart, K.; Bauer, C.; Boulet, C. Life Cycle Assessment of Carbon Capture and Storage in

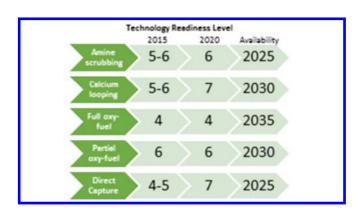
415 Power Generation and Industry in Europe. *Int. J. Greenh. Gas Control* **2013**, *16*, 91–106.

- 416 (28) Bjerge, L.-M.; Brevik, P. CO2 Capture in the Cement Industry, Norcem CO2 Capture Project
- 417 (Norway). *Energy Procedia* **2014**, *63*, 6455–6463.
- 418 (29) Zeman, F. Oxygen Combustion in Cement Production. *Energy Procedia* 2009, 1 (1), 187–194.
- 419 (30) Barker, D. J.; Holmes, D.; Hunt, J.; Napier-Moore, P.; Turner, S.; Clark, M. CO2 Capture in the
- 420 *Cement Industry*; IEAGHG: Cheltenham, UK, 2008.
- 421 (31) Bhatty, J. I.; Miller, F. M.; Kosmatka, S. H.; Bohan, R. P. Innovations in Portland Cement
- 422 *Manufacturing*; Portland Cement Association: Skokie, IL, 2011.
- 423 (32) Koring, K.; Hoenig, V.; Hoppe, H.; Horsch, J.; Suchak, C.; Klevenz, V.; Emberger, B. *Deployment*424 of CCS in the Cement Industry; 2013/10; IEAGHG: Cheltenham, UK, 2013.
- 425 (33) Moya, J. A.; Pardo, N.; Mercier, A. The Potential for Improvements in Energy Efficiency and
- 426 CO2 Emissions in the EU27 Cement Industry and the Relationship with the Capital Budgeting
- 427 Decision Criteria. J. Clean. Prod. **2011**, *19* (11), 1207–1215.
- 428 (34) Schneider, M. ECRA's Oxyfuel Project, Norcem International CCS Conference, Langesund
 429 Norway, May 20 21 2015.
- 430 (35) Rodríguez, N.; Murillo, R.; Abanades, J. C. CO2 Capture from Cement Plants Using Oxyfired
- 431 Precalcination And/or Calcium Looping. *Environ. Sci. Technol.* **2012**, *46* (4), 2460–2466.
- 432 (36) Gale, J. A Global Perspective on CO2 Capture Developments, 2014.
- 433 (37) Davison, J. Pilot Plant Trial of Oxy-Combustion at a Cement Plant; Information Paper 2014-IP7;
- 434 IEAGHG: Cheltenham, UK, 2014.
- 435 (38) Dean, C. C.; Blamey, J.; Florin, N. H.; Al-Jeboori, M. J.; Fennell, P. S. The Calcium Looping Cycle
- 436 for CO2 Capture from Power Generation, Cement Manufacture and Hydrogen Production.
- 437 *Chem. Eng. Res. Des.* **2011**, *89* (6), 836–855.
- 438 (39) Ozcan, D. C.; Ahn, H.; Brandani, S. Process Integration of a Ca-Looping Carbon Capture
- 439 Process in a Cement Plant. *Int. J. Greenh. Gas Control* **2013**, *19*, 530–540.

- 440 (40) Industrial Technology Research Institute. R & D Achievements on Carbon Capture and Storage
- 441 in ITRI, Taiwan http://ccs.tw/sites/default/files/datashare/pdf/2014-10-06-0923-
- 442 gong_yan_yuan_ccsyan_fa_cheng_guo_ying_wen_.pdf (accessed Nov 22, 2015).
- 443 (41) Chang, M.-H.; Huang, C.-M.; Liu, W.-H.; Chen, W.-C.; Cheng, J.-Y.; Chen, W.; Wen, T.-W.;
- 444 Ouyang, S.; Shen, C.-H.; Hsu, H.-W. Design and Experimental Investigation of Calcium Looping
- 445 Process for 3-kWth and 1.9-MWth Facilities. *Chem. Eng. Technol.* **2013**, *36* (9), 1525–1532.
- 446 (42) Calix Ltd. Direct Separation Technology for Low Emissions Intensity Lime and Cement
- 447 http://www.calix.com.au/cement-and-lime.html (accessed Mar 10, 2015).
- 448 (43) Sceats, M. Direct Capture for the Cement Industry, 2015.
- 449 (44) Tokheim, L.-A. Benchmark Study Commercial Scale Perspective, Norcem International CCS
- 450 Conference, Langesund Norway, May 20 21 2015.
- 451 (45) Green, D. W.; Perry, R. H. *Perry's Chemical Engineers' Handbook, Eighth Edition*, 8 edition.;
 452 McGraw-Hill Professional: New York, 2007.
- 453 (46) Schneider, M.; Hoenig, V. Development of State of the Art Techniques in Cement
- 454 *Manufacturing: Trying to Look Ahead (CSI/ECRA Technology Papers)*; Cement Sustainability
 455 Initiative, 2009.
- 456 (47) Kuramochi, T.; Ramírez, A.; Turkenburg, W.; Faaij, A. Comparative Assessment of CO2 Capture
- 457 Technologies for Carbon-Intensive Industrial Processes. Prog. Energy Combust. Sci. 2012, 38
- 458 (1), 87–112.
- 459 (48) Ho, M. T.; Allinson, G. W.; Wiley, D. E. Comparison of MEA Capture Cost for Low CO2
- 460 Emissions Sources in Australia. *Int. J. Greenh. Gas Control* **2011**, *5* (1), 49–60.
- 461 (49) British Standards Institute. Cement. Composition, Specifications and Conformity Criteria for
- 462 *Common Cements*; BS EN 197-1:2011; 2011.
- 463 (50) Lafarge SA. Lafarge Annual Report 2007; Paris.
- 464 (51) Cochez, E.; Nijs, W. *ETSAP: Cement Production*; Technology Brief 103; IEA ETSAP, 2010.

- 465 (52) Carbon Capture Storage project in Estevan takes another step forward
- 466 http://www.estevanmercury.ca/news/city/carbon-capture-storage-project-in-estevan-takes-
- 467 another-step-forward-1.1450820 (accessed Jun 4, 2015).
- 468 (53) FLSmidth Highlights Archive http://www.flsmidth.com/en-US/eHighlights/Highlights+archive
 469 (accessed Jun 5, 2015).
- 470 (54) Bohm, M. C.; Herzog, H. J.; Parsons, J. E.; Sekar, R. C. Capture-Ready Coal plants—Options,
- 471 Technologies and Economics. Int. J. Greenh. Gas Control **2007**, 1 (1), 113–120.
- 472 (55) Lucquiaud, M.; Chalmers, H.; Gibbins, J. Capture-Ready Supercritical Coal-Fired Power Plants
 473 and Flexible Post-Combustion CO2 Capture. *Energy Procedia* 2009, *1* (1), 1411–1418.
- 474 (56) Liang, X.; Reiner, D.; Gibbins, J.; Li, J. Assessing the Value of CO2 Capture Ready in New-Build
- 475 Pulverised Coal-Fired Power Plants in China. Int. J. Greenh. Gas Control **2009**, 3 (6), 787–792.
- 476 (57) Rohlfs, W.; Madlener, R. Assessment of Clean-Coal Strategies: The Questionable Merits of
- 477 Carbon Capture-Readiness. *Energy* **2013**, *52*, 27–36.
- 478 (58) Barker, D. J.; Turner, S. A.; Napier-Moore, P. A.; Clark, M.; Davison, J. E. CO2 Capture in the
- 479 Cement Industry. *Energy Procedia* **2009**, *1* (1), 87–94.
- 480 (59) Global CCS Institute. ITRI Calcium Looping Pilot (accessed Mar 10, 2015).

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TOC 84x47mm (96 x 96 DPI)

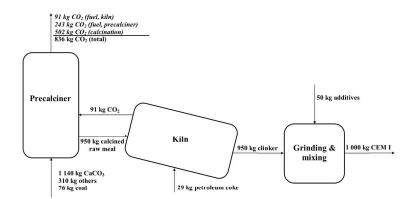


Figure 1: Direct emissions of CO2 from CEM I (95% clinker) cement manufacture (own calculations). CEM I rather than CEM II was chosen for comparisons in this paper because of its smaller range of composition than CEM II (95 – 100% clinker by weight versus 35 – 94%). 209x147mm (300 x 300 DPI)