## Review Article

# LITERATURE SURVEY OF FLUE GAS TREATMENT UNIT FOR PACKAGED GASOLINE GENERATORS. 6

### ABSTRACT

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9 A literature survey of Flue Gas Treatment Unit for Packaged Gasoline Generators is reported. Due to shortfall in power supply from National Grid of most developing nations, 10 11 most businesses relied on package generators for electrical power supply for their economic 12 activities. Flue gases containing  $CO, CO_2, SO_2$  and  $NO_x$  were being released into the 13 environment by these fossil fired package generators. The CO is known to be detrimental to 14 health of living beings and at the same time ozone layer depletive. On the other hand, the 15 CO<sub>2</sub> has been contributing to the green house effects. These militating dangers/risks arising 16 from the usage of these package generators strongly informed this review exercise. The global position on legislation, regulation and control of the emission of flue gases into the 17 18 atmosphere has been articulated. Results of in-depth investigations by researchers into the methods/techniques of capturing CO,  $CO_2$ , SO<sub>2</sub> and NO<sub>x</sub> from the flue gases have been 19 highlighted. The incorporation of these capturing methods for CO, CO<sub>2</sub>, SO<sub>2</sub> and NO<sub>x</sub> into 20 21 the design and construction of the package gasoline generators will no doubt go a long way 22 in keeping our environment healthy, safe and secured. 23 KEY WORDS: Flue Gas Treatment Unit, Emission Limit Values, Package Generators. 24 INTRODUCTION By the end of the 20<sup>th</sup> century it was widely accepted that carbon dioxide and several other 25 26 gases are involved in physical and chemical processes in the earth's upper troposphere and 27 stratosphere that may result in global climate change. Arrhenius in 1896 forecasted rising 28 global temperatures as a result of fossil fuel combustion. So-called green house gases

29 (GHG), most importantly Carbon Dioxide (CO<sub>2</sub>), Methane (CH<sub>4</sub>) and Nitrous Oxide (N<sub>2</sub>O) 30 trap the outgoing solar radiation that is reflected by the earth's surface, which leads to global warming.<sup>[1]</sup> Although the most abundant greenhouse gas in the atmosphere is water causing 31 32 approximately 2/3 of the greenhouse effect, the result of increasing concentrations of GHGs 33 that cause the other approximately 1/3 is referred to as the "enhanced greenhouse effect", 34 or, since it is primarily the result of human activities, the "anthropogenic greenhouse 35 effect" <sup>[1]</sup> Consequent on the foregoing therefore, various regulations have been put into place to arrest this green house effect. 36

In order for society to meet up with its energy demand, various forms of energy sources 37 38 have been harnessed by mankind. Most of these power plants are fired by fossil fuels which give CO<sub>2</sub>, CO<sub>3</sub>, SO<sub>2</sub>, NO<sub>x</sub> as by-products of combustion amongst others. CO<sub>2</sub> in particular 39 40 has been identified as the most important green house gas. Various efforts are being made 41 and several actions are being taken by the global regulating bodies to control and limit the 42 emission of CO<sub>2</sub> and other green house gases. These efforts that focus on design /capture 43 concept are not extended to the gasoline package generators. In the developing countries, 44 the package generators which are actually designed as standby generators now play the 45 function of source of main power supply for businesses and homes. The attendant 46 implications are that  $CO_2$ ,  $CO_1$ ,  $SO_2$  and  $NO_x$  are being indiscriminately emitted into the 47 environment to the detriment of health and global warming which could result from the greenhouse effect. Consequent on the foregoing some action plan is required to arrest this 48 indiscriminate pollution of the environment arising from the gasoline package generators' 49 emissions among others. This literature survey therefore seek to articulate the global 50 51 legislation framework in place, research efforts and actions taken so far to control the 52 emission of flue gases into the atmosphere. Here lies the justification for this review.

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### 54 THE PACKAGED GASOLINE GENERATOR AND FLUE GASES

55 Generators range from extreme large structures such as turbo-generators as in thermal 56 power plants, nuclear power plants, hydro power plants, geothermal power plants, Brown 57 gas generator, wood gas generator, wind and tidal power plants to package units used in the 58 home, emergency and leisure as standby sources of power. Generators could be classified 59 according to source of energy hence we have thermal generators, geothermal generators, 60 hydro generators and solar generators. Also generators could be classified based on the 61 source of fuel: such generators include natural gas generators, gasoline generators, diesel 62 generators, brown gas generators and wood gas generators among others. Such package generators are variously referred to as portable generators, portable electric generators, 63 portable power generators, home standby generators, commercial generators and marine 64 generators.<sup>[2]</sup> The power range for a variety of these package generators varies from about 65 800watts to 15 kilowatts.<sup>[2]</sup> These generators could be classified according to manufacturers 66 and marketing outfit. A few of such names include: Honda, Yamaha, Norwall power systems, 67 68 Binatone, Firman, Elemax and Tiger.

The energy and power delivery to the economies of the developing nations has been dwindling over the years. The quality of power generated, that is supply relative to demand has been on the decline, even in the face of new power stations coming on stream the National Grid. This short fall in meeting the National demand has been so severe that virtually all serious business concerns have to procure one form of generating set or the other in order to guarantee the smooth continuity of their businesses.<sup>[3]</sup>

75 The brand of these generators that are fossil fuel fired release flue gases into the 76 environment. Flue Gas has been defined as gas that exits to the atmosphere via a flue 77 which may be a pipe, channel or chimney for conveying combustion from a fireplace, oven, boiler or steam generator.<sup>[4]</sup>Flue gases are produced when coal, fuel oil, natural gas, wood 78 or any other fuel is combusted in an industrial furnace or boiler, a steam generator in a fossil 79 fuel power plant or other combustion sources.<sup>14</sup>These gases are treated as pollutants. Flues 80 81 can be referred to as "stacks", and they may be found in the form of chimneys, ducts, or 82 simple pipes. Large amount of flue gases are generated around the world on daily basis, 83 with heavy industry and the power industry in particular being responsible for a huge 84 percentage of the total generated. The contents of flue gases are quite variable. The 85 medium being burned can contribute a number of different compounds, and the conditions 86 under which combustion is occurring can also generate more or less emissions. Incomplete 87 combustion at low temperatures or in poorly managed facilities, for example, tends to generate more pollution.<sup>[5]</sup> Components of flue gases include: water vapor, carbon dioxide, 88 nitrogen, particulates, oxygen, carbon monoxide, nitrogen oxide, hydrocarbons, and sulfur 89 90 oxide. Some of these compounds are potentially harmful for the environment, making these gases an issue of concern among environmental advocates. Flue gases can also be 91 92 hazardous for human health, as might occur if they were trapped in an air inversion which 93 pinned them close to the ground for several days, forcing people in the area to inhale hazardous pollutants which could damage their lungs.<sup>[5]</sup> 94

95 There are a number of ways in which flue gases can be controlled, and the processes which produce flue gases are often heavily regulated to force emissions levels down. One of the 96 97 best methods is to avoid generating them at all, either by using alternative technology, 98 improving plant efficiency levels, or studying ways in which operating conditions could be improved to reduce the production of combustion by-products.<sup>[5]</sup> Gases which cannot be 99 prevented can be trapped using filters and scrubbers which clean the air coming out of flues 100 101 so that when it is released into the environment, it contains primarily harmless components. 102 Scrubbing flue gases can even be profitable for a savvy company. For example, the food industry is willing to pay for purified carbon dioxide extracted from flue gases.<sup>[5]</sup> 103 104 LEGISLATIONS/REGULATIONS ON POLLUTION

105 . It is not possible to produce comparative tables with emission limit values for the two 106 regions (that is, EU and USA) because emission limit values are defined in different ways. In

107	the EU, values are defined in terms of mg/nm <sup>3</sup> , whereas In the US values are expressed as
108	nanograms per joule heat input. For the EU, emission limit values are defined in Annex III-VII
109	in the Directive 2001/80/EC on the limitation of emissions of certain pollutants into the air
110	from large combustion plants (the "LCP directive"). Emission limits are defined for $SO_2$ , $NO_x$
111	and dust. Furthermore, there are provisions for combustion plants co-incinerating waste in
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	Annex II of the directive 2000/76/EC of 4 December 2000 on the incineration of waste. For
113	the USA, emission limit values related to combustion plants are given in Title 40 (Protection
114	of Environment), Part 60 (Standards of Performance for New Stationary Sources) in
115	subparts Da, Db, Dc and GG.
116	S60.43 Standard for Sulfur Dioxide (SO <sub>2</sub> )
117	(a) On and after the date on which the performance test required to be conducted by
118	S60.8 is completed, no owner or operator subject to the provision of this subpart
119	shall cause to be discharged into the atmosphere from any affected facility any
120	gases that contain SO <sub>2</sub> in excess of: (1) 340ng/J heat input (0.8lb/MMBtu) derived
121	from liquid fossil fuel or liquid fossil fuel and wood. (2) 520 ng/J heat input (1.2
122	lb/MMBtu) derived from solid fossil fuel or solid fossil fuel and wood residue.
123	(b) When different fossil fuels are burned simultaneously in any combination, the
124	applicable standard (in ng/J) shall be determined by proration using the formula:
125	$P_{SO2} = [y(340) + z(520)]/(y+z)$
126	Where $P_{SO2}$ = prorated standard for SO <sub>2</sub> when burning different fuels simultaneously
127	in ng/J heat input derived from all fossil fuels or from all fossil fuels and wood
128	residue fired.
129	Y = percentage of total heat input derived from liquid fossil fuel and
130	Z = percentage of total heat input derived from solid fossil fuel.
131	Sulfur Content of Liquid Fuels Directive 1999/32/EC on Reduction of Certain
132	Liquid Fuels (26 <sup>th</sup> April 1999).
133	Sulfur content of gas oil not to exceed 0.1 %. Member states shall take all necessary
134	steps to ensure that as from 1 January2003 within their territory heavy fuel oils are
135	not used if their sulfur content exceeds 1.0% by mass. Alternatively, where the
136	emissions of sulfur dioxide from the plant are less than equal to 1700mg/nm <sup>3</sup> at an
137	Oxygen content in the flue gas of 3% by volume on a dry basis. The reference
138	method adopted for determining the sulfur content shall be that determined by:
139	a) ISO Method 8754 (1992) and PrEN ISO 14596 for heavy oil and marine gas oil
140	b) EN Method 24260 (1987) ISO 8754 (1992) and PrEN ISO 14596 for gas oil.
141	The arbitration method will be PrEN ISO 14596. The statistical interpretation of the
142	verification of the sulfur content of the gas oil used shall be carried out in
143	accordance with ISO Standard 4259 (1992). <sup>[6]</sup>
144	CHINA'S "TWELFTH FIVE-YEAR PLAN" ON AIR POLLUTION PREVENTION
145	On 5 <sup>th</sup> December 2012, Ministry of Environmental Protection (MEP), National
145	Development and Reform Commission and Ministry of Finance jointly issued a "12 <sup>th</sup>
	five-year plan on Air Pollution and Control in key regions". The plan was approved
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148	by the China's State Council earlier. The three key regions are: Beijing-Tianjin-
149	Hebei, Yangtze River Delta and Pearl River Delta and 10 city clusters, involving 19
150	provincial level jurisdictions and 117 cities. These areas cover only 14% of the
151	country's land area, but accounts for nearly half (48%) of the country's population,
152	71% of the nations GDP and 52% of the country's coal consumption. The air
153	pollutants emission level is 2.9 to 3.6 times higher than the nation's average.
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157	i) The Plan sets ambient concentration targets for the first time, thus:
	Ambient Air PM <sub>10</sub> SO <sub>2</sub> NO <sub>2</sub> PM <sub>2.5</sub> PM <sub>2.5</sub>

The Plan sets ambient concentration targets for the first time, thus:					<mark>st time, thus:</mark>	
Ambient	Air	<mark>PM₁₀</mark>	SO <sub>2</sub>	NO <sub>2</sub>	PM <sub>2.5</sub>	PM <sub>2.5</sub>
<b>Quality</b>						<mark>(In 3 Key</mark>

									Regions)
		Concentration	<mark>10%</mark>	<mark>10%</mark>	<mark>7%</mark>	<mark>5%</mark> (	(Expec	ted	6%(Binding
		Targets				Tar			Target)
158	ii)	The Plan sets mor	re ambi	tious er	nissior	targ	ets in	key re	egions than the
159		national tagets							
		Total Emission F				SO <sub>2</sub>	<mark>NO</mark> x		strial PM
		National Targets				<mark>3%</mark>	<mark>10%</mark>		arget
		Targets in Key R	egions	By 201	<mark>5</mark> 1	<mark>12%</mark>	<mark>13%</mark>	<mark>10%</mark>	)
160									
161		If successfully imp							
162 163		reduction of 2.28r tons/year and PM							
164		emission reduction							
165		over \$317 Billion (							pected to bring
166	<b>RESEARCH EF</b>	FORTS TO MINIMI				Joioty	borront	•	
167		ect of SO <sub>2</sub> on select				on (S	CR) of	NO	by NH <sub>3</sub> over a
168		talyst sorbent has							
169		spectroscopy (XPS)							
170		) were used to ev							
171		nd TPR in H <sub>2</sub> sugg							
172	catalyst's oxida	tion ability and inhib	oits NH <sub>3</sub>	oxidati		NO IN	the pr	ocess	of SCR, which
173 174		main promoting effe							
175		ed. Previous work o							
176		can be tuned by mo							
177		another acidic gas f							
178		gned to investigate							
179		t <mark>ies as well as sw</mark>							
180		thermogravimetric							
181		ransform infrared (FT							
182 183		found that the SO <sub>2</sub>							
184		) PPM; P <sub>Tot</sub> = 0.4mp ; (i.e ether and aming							
185		n the other hand,							
186		important tool to ac							
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189		he only feasible teo							
190		such techniques, er							
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193		er these resources of							
195		bstacles related to co							
196		pture and storage (							
197	plants and in st	teel and cement pro	duction	while la	argely e	elimin	ating th	eir CO	O <sub>2</sub> emissions. <sup>[10]</sup>
198	the removal of (	CO <sub>2</sub> from flue gases	is done	with the	e desir	e to re	educe g	reenh	<mark>iouse emissions</mark>
199		ctors. The Econoami							
200		ng an inhibited 30w							
201		% of the CO2 in the							
202 203	primary amines	dilute CO <sub>2</sub> from atm [11]	ospheric	pressu	ne gas	are r	nonoeti	anola	amine and other
200	Printary anniles	•							

- 204 The absorber diameter can be approximated from equation 2:
- 205 Absorber dia,(m) = Asqrt[(te/d)/(%CO<sub>2</sub>)]
- 206 Stripper diameter = 0.13\*sqrt(te/d)

207 Solvent loading and circulation rate (rich solvent circulation rate)=0.71\*(te/d)

- 208 Power consumption  $Kw = (0.4 + 16.4(\%CO_2))^*(te/d)$
- 209 Where A = 0.56 at 3% CO<sub>2</sub> to 0.62 for 13% CO<sub>2</sub>,
- 210  $Te/d = tonne/day CO_2 recovered,$

212  $M = meters.^{[11]}$ 

213 In similar vein, an experimental study on NO<sub>x</sub> removal from a simulated diesel engine was 214 carried out using dielectric barrier discharge (DBD) reactor for various electrode shapes to 215 improve the removal efficiency. The electrode shapes employed in the experiment were a plane, a trench and multipoint configurations. The 0.5mm thickness of alumina 216 217 (Al/sub2/O/sub3/) dielectric barrier was produced on a plane electrode via a powder flame 218 spray. The DBD reactor with multipoint electrodes was applied to 20KVA diesel engine 219 generator exhaust gas treatment. The contained  $NO_x = (NO + NO/sub2/)$  was abated from 70 220 to 40 ppm at no load to the generator and from 340 to 300 ppm at 50A load. The energy for NO<sub>x</sub> removal is obtained to 32g/Kwh.<sup>[12</sup> 221

222 In another development, atmospheric  $O_3$ , CO, SO<sub>2</sub> and NO<sub>v</sub>\* were measured at a rural 223 agricultural site in the Yangte Delta of China. Positive correlation between CO and  $NO_v^*$  with 224 a slope  $(\Delta[CO]/\Delta[NO_v^*])$  of 36 (ppbv/ppbv) for the winter and night time measurements. 225 Value for industrialized countries~(10ppbv/ppbv). Where NO<sub>v</sub>\* = (NO + NO<sub>2</sub> + PAN + Organic Nitrates + HNO<sub>3</sub> +  $N_2O_5$  +---). The highest CO/NO<sub>v</sub>\* ratio (30 - 40ppbv/ppbv) 226 occurred in September-December 1999 and June 2000. The good correlation between CO 227 228 and the biomass burning tracer CH<sub>3</sub>CI and the lack of correlation with the industrial tracer 229 C<sub>2</sub>Cl<sub>4</sub> suggest that the burning of biofuels and crop residues is a major source for the 230 elevated CO and possibly for other tracer gases as well. SO<sub>2</sub> to NO<sub>v</sub>\* ratio = 1.37 ppbv/ppbv. 231 High value of ratio is as a result of use of high sulfur coals in China. The measured 232  $SO_2/NO_v^*$  and  $\Delta CO/\Delta NO_v^*$  were compared with the respective ratios from the current 233 emission inventories for the region, which indicated a comparable SO<sub>2</sub>/NO<sub>v</sub>\* emission ratio, 234 but a large discrepancy for CO/NO<sub>x</sub>. The observed CO to NO<sub>v</sub>\* ratio was more than 3 times the emission ratio derived from inventories.<sup>[13]</sup> 235

236 Also, large eddy simulation (LES) of the gas-second order moment (SOM) of particles 237 model based on the two fluid method coupled with the subgrid-scale (SGS) reaction 238 submodel was applied to investigate SO<sub>2</sub> and NO<sub>x</sub> emissions circulating fluidized bed (CFB) 239 combustors. The SGS reaction submodel based on the eddy dissipation concept (EDC) 240 model was developed considering the effect of particles to circulate the SGS reaction rates. 241 The predicted results by the model are in agreement with the experimental data. The flow 242 characteristics including concentration and SOM of particles were analyzed. The simulation 243 results show that the particles including coal and desulfurized particles were sufficiently 244 mixed. The velocity fluctuation of particles was enhanced for the combustion process, and 245 the velocity fluctuation in the axial direction was about twice higher than that in the radial 246 direction. It was observed that the SGS reaction of gas not only improved the homogenous 247 reaction of gas but the heterogeneous reaction also.<sup>[14]</sup>

248 In another investigation, Iron oxide supported on alumina as a catalyst/absorbent for use in 249 the simultaneous removal of  $NO_x$  and  $SO_x$  from power plant stack gases is reported. A dry-250 contacting process underdevelopment would operate under net reducing conditions at 251 temperatures of  $370^{\circ}$  to  $540^{\circ}$ C. Iron oxide is converted to the ferrous state. No is reduced to 252 NH<sub>3</sub> and SO<sub>2</sub> is removed as ferrous sulfide or sulfate. Regeneration with air provides SO<sub>2</sub> 253 and reforms  $Fe_2O_3$ . The reduction of  $SO_2$  by CO and  $H_2$  was studied in fixed bed reactors to 254 determine the effects of temperature and of the other reactive components of flue gas except 255 (flyash) on the rate of reaction and the products formed.  $H_2S$  and COS react with FeO to 256 form FeS. Under readily attainable conditions, virtually complete removal of sulfur

Con a similar note, Iron titanate catalyst (F <sub>a</sub> T,O <sub>a</sub> ) is a potential candidate for the substitution of constraints (Correction (SCR) of NO <sub>a</sub> with NH <sub>b</sub> because of its high SCR activity and N <sub>a</sub> . <sup>151</sup> Also, an Iron titanate catalyst with a crystalline phase, prepared by a co-precipitation method showed excellent activity, stability, selectivity and SO <sub>2</sub> H <sub>2</sub> O durability in the selective catalytic reduction (SCR) of NO <sub>a</sub> with NH <sub>b</sub> because of its high SCR activity and N <sub>a</sub> . <sup>151</sup> Also, an Iron titanate catalyst with a crystalline phase, prepared by a co-precipitation method showed excellent activity, stability, selectivity and SO <sub>2</sub> H <sub>2</sub> O durability in the selective catalytic reduction of NO with NH <sub>b</sub> in the medium temperature range <sup>117</sup> Furthermore, a cyclic process is described for removing the oxides of Sulfur from flue gas by absorption at 625°F and then converting the oxides to elemental sulfur by reduction of the spent absorbent. Pilot plant experiments are reported in which a solid absorbent of alkalized alumina in free and affied fall is used to remove SO <sub>2</sub> and SO <sub>3</sub> from the combustion gas of a pulverized coal- fired furnace. The absorbent is then regenerated by heating with hydrogen or with steam- reformed natural gas. Fly ash does not interfer and attrition is negligible. <sup>118</sup> On the other hand, the effectiveness of limestone and dolomite in removing SO <sub>2</sub> from flue gas can be temperature range where furnace conditions and reaction kinetics are likely to be favorable. <sup>119</sup> on the other hand, one of the most important steps in the wet limestone gypsum flue gas desulfurization (WFGD) process is CaCO <sub>3</sub> dissolution, which provides the dissolved atkinity necessary for SO <sub>2</sub> absorption. Accurately evaluating the CaCO <sub>3</sub> dissolution rate is important in the design and efficient operation of WFGD plants. In the present work, the dissolution of limestone from different sources in South Artica has been studied in a pH-stat apparatus under conditions similar to those encountered in wet FGD processes. The i	257	compounds was achieved for gas-phase residence times of about 1 to 10ms. NO and NO <sub>2</sub>
conventional V <sub>2</sub> O <sub>2</sub> -WO <sub>2</sub> (M <sub>2</sub> O <sub>2</sub> )T <sub>1</sub> O <sub>2</sub> and F <sub>2</sub> /C <sub>2</sub> -Zeolite catalysts for the selective catalytic reduction (SCR) of NO <sub>2</sub> with NH <sub>3</sub> because of its high SCR activity and N <sub>2</sub> . <sup>[16]</sup> Also, an Iron titanate catalyst with a crystalline phase, prepared by a co-precipitation method showed excellent activity, stability, selectivity and SO <sub>2</sub> /H <sub>2</sub> O durability in the selective catalytic reduction of NO with NH <sub>3</sub> in the medium temperature range <sup>117</sup> Furthermore, a cyclic process is described for removing the oxides of Sulfur from flue gas by absorption at 625 <sup>6</sup> F and then converting the oxides to elemental sulfur by reduction of the spent absorbent. Pliof plant experiments are reported in which a solid absorbent of alkalized alumina in free and baffield fail is used to remove SO <sub>2</sub> and SO <sub>3</sub> from the combustion gas of a pulverized coal- fired funce. The absorbent is then regenerated by heating with hydrogen or with steam- reformed natural gas. Fly ash does not interfere and attrition is negligible. <sup>[18]</sup> On the other hand, the effectiveness of limestone and dolomite in removing SO <sub>2</sub> from flue gas can be estimated by thermo-chemical reactions. Both lime and magnesia are shown to be suitable for removing SO <sub>2</sub> from tof thu gas, with lime more suitable than magnesia over the favorable. <sup>[19]</sup> on the other hand, one of the most important steps in the wel limestone gypsum flue gas desulfurzation (WFCD) process is CaCO <sub>2</sub> dissolution, which provides the dissolved alkalinity necessary for SO <sub>2</sub> absorption. Accurately evaluating the CaCO <sub>2</sub> dissolution rate is important in the design and efficient operation of WFGD plants. In the present work, the dissolution of limestone from different sources in South Africa has been studied in a pH-stat apparatus under conditions similar to those encountered in wel FGD processes. The influence of various parameters such as the reaction temperature, (30 < T <70°C), CaCO <sub>2</sub> particle size (25 < d g < 50 × 00, gradiculate control, mercury control and others ( VOCCs, dissolution rate in	258	were also reduced. <sup>[15]</sup>
reduction (SCR) of NO, with NH <sub>5</sub> because of its high SCR activity and N <sub>2</sub> . <sup>169</sup> Also, an iron titanate catalyst with a crystalline phase, prepared by a co-precipitation method showed excellent activity, stability, selectivity and SO,/H <sub>2</sub> O durability in the selective catalytic reduction of NO with NH <sub>5</sub> in the medium temperature rang <sup>10-17</sup> Eurthermore, a cyclic process is described for removing the oxides of Sulfur from flue gas by absorption at 625°F and then converting the oxides to elemental sulfur by reduction of the spent absorbent. Pilof plant experiments are reported in which a solid absorbent of alkalized alumina in free and affield fall is used to remove SO <sub>2</sub> and SO <sub>3</sub> from the combustion gas of a pulverized coal- fired furnace. The absorbent is then regenerated by heating with hydrogen or with steam- reformed natural gas. Fly ash does not interfere and attrition is negligible. <sup>118</sup> On the other hand, the effectiveness of limestone and dolomite in removing SO <sub>2</sub> from flue gas can be estimated by thermo-chemical reactions. Both lime and magnesia are shown to be suitable for removing SO <sub>2</sub> from hot flue gas, with lime more suitable than magnesia over the temperature range where furnace conditions and reaction kinetics are likely to be favorable. <sup>119</sup> on the other hand, one of the most important steps in the wet limestone gypsum flue gas desulfurization (WFGD) process is CaCO <sub>2</sub> dissolution, which provides the dissolved alkalinity necessary for SO <sub>2</sub> absorption. Accurately evaluating the CaCO <sub>3</sub> dissolution rate is inportant in the design and efficient operation of WFGD plants. In the present work, the dissolution of limestone from different sources in South Africa has been studied in a pH-stat apartule size (25 < d p < 63µm), solution acidity (4 < pH < 6) and chemical composition were studied in order to determine the kinetics of CaCO <sub>3</sub> dissolution. The results obtained indicate that the dissolution rate increased with a decrease in the particle size and increase in temperature. <sup>120</sup> In a similar v	259	On a similar note, Iron titanate catalyst ( $F_eT_iO_4$ ) is a potential candidate for the substitution of
titanate catalyst with a crystalline phase, prepared by a co-precipitation method showed excellent activity, stability, selectivity and SO <sub>2</sub> /H <sub>2</sub> O durability in the selective catalytic reduction of NO with NH <sub>3</sub> in the medium temperature range <sup>117</sup> Furthermore, a cyclic process is described for removing the oxides of Suffur from flue gas by absorption at 625°F and then converting the oxides to elemental suffur by reduction of the spent absorbent. Pilot plant experiments are reported in which a solid absorbent of alkalized alumina in free and baffield fall is used to remove SO <sub>2</sub> and SO <sub>3</sub> from the combustion gas of a puberized coal- fired furnace. The absorbent is then regenerated by heating with hydrogen or with steam- reformed natural gas. Fly ash does not interfere and attrition is negligible. <sup>181</sup> On the other hand, the effectiveness of lineace conditions and reaction kinetics are likely to be estimated by thermo-chemical reactions. Both lime and magnesia are shown to be suitable for removing SO <sub>2</sub> from hot flue gas, with lime more suitable than magnesia over the estimated by thermo-chemical reactions. Both lime and magnesia were the favorable. <sup>190</sup> on the other hand, one of the most important steps in the wel limestone gypsum flue gas desulfurization (WFGD) process is CaCO <sub>3</sub> dissolution, which provides the dissolved alkalinity necessary for SO <sub>2</sub> absorption. Accurately evaluating the CaCO <sub>3</sub> dissolution rate is important in the design and efficient operation of WFGD plants. In the present work, the dissolution of limestone from different sources in South Africa has been studied in a pH-stat apparatus under conditions similar to those encountered in wet FGD processes. The influence of various parameters such as the reaction temperature, ( $30 \le 1 \le 70^\circ$ C), CaCO <sub>3</sub> particle size ( $25 \le d \ne 53^{SD}$ M), solution actific 4 s pH 4 6) and chemical composition were studied in order to determine the kinetics of CaCO <sub>3</sub> dissolution. The results obtained indicate that the dissolution rate increased with a decrease	260	
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	309	devices. <sup>[25]</sup> A performance report is documented on the economic, technical and

environmental feasibility tests on running a 20MW diesel generator on vegetable oil (palm
oil). The main constraint identified in this report is the availability and cost of the vegetable
(palm) oil. <sup>[26]</sup> The Malaysian AM cash market prices for palm oil stood at US \$612 per
metric Tonne.<sup>[27]</sup> Also global petrol price in USA stood at US\$0.97 per litre, UK at US\$2.18
and Nigeria at US\$0.58 per litre respectively.<sup>[28]</sup> In the same vein World market gas prices
averaged US\$3.41 per gallon.<sup>[29]</sup> While Butane gas is priced £16.75 (or \$27.3) per 4.5kg
cylinder and Propane gas £16 (or \$26.08) per 3.9kg cylinder.<sup>[30]</sup> Also, gasoline consumption

317 for a 10KVA package gasoline generator stands at18 litres in 4.5 hours.<sup>[31]</sup>

### 319 SUMMARY OF LITERATURE REVIEW

### **Table 1: Summary of Legislation/ Regulation Review.**

		EU	USA	
EMISSION LIMIT VALUES (ELV)		mg/NM <sup>3</sup>	ng/J Heat	ppbv/ppbv
	ELV defined in	Annex III-VII in the Directive 2001/80/EC. Directive 1999/32/EC	Title 40 Part 60 Subparts Da, Db, Dc and GG.	"12 <sup>TH</sup> 5-year Plan" on Pollution Prevention
<mark>SO₂</mark>	Liquid fossil fuel	0.1 by mass	<mark>340</mark>	10%. National target 2015 = 8%. Target region = 12%
	<mark>Solid fossil</mark> fuel	1.0% for heavy fuel oils Or for <u>&lt;</u> 1700mg/Nm <sup>3</sup> @O₂ content in flue gas of 3% by volume.	<mark>520</mark>	
	Combined	For heavy oil and marine gas oil: ISO method 8754 (1992) For gas oil; EN 2426O (1987)	P <sub>SO2</sub> (see equation 1)	
	NO <sub>2</sub>			7% National target 2015 = 10% Target region = 13%

 SOURCE: Summary Of Legislation/Regulation Review above.

Table 2 Summary of capture methods from flue gas

S/No. COMPONENTS REDUCTION METHOD MATERIALS AUTHOR'S

				REFERENCE NUMBER
1	SO₂	Ambient content measured at rural agricultural site	Gas analyzers	[13]
2		Large eddy simulation (LES), Gas-second order moment (SOM) and CFB combustors coupled with sub-grid scale (SGS). (two-fluid method)	Coal and desulfurized particles	[14]
<mark>3</mark>		A dry contacting process for simultaneous removal of $NO_x$ and $SO_x$ from plant stack.	Iron oxide supported on alumina as catalyst	[15]
<mark>4</mark>		Cyclic process of removing SO <sub>2</sub> and SO <sub>3</sub> by absorption	Absorbent, alkalized alumina in free and baffle fall.	[18]
<mark>5</mark>		Thermo-chemical reactions using limestone and dolomite	Limestone, dolomite and magnesia	[19]
<mark>6</mark>		Limestone gypsum absorption	limestone	[20]
7	NO <sub>x</sub>	Selective catalytic reduction (SCR) presence of SO <sub>2</sub> weakens NH <sub>3</sub> oxidation to NO	NH <sub>3</sub>	[8]
8		Dielectric barrier discharge (DBD) reactor applied to 20KVA diesel engine generator	Al/sub2/O/sub3/ Alumina dielectric barrier	[12]
<mark>9</mark>		Ambient content measured at rural agricultural site	Gas analyzers	[13]
<mark>10</mark>		Large eddy simulation (LES) gas-second order moment (SOM) and CFB combustors coupled with sub-grid (SGS). (two-fluid method)	Coal and desulfurized particles	[14]
<mark>11</mark>		A dry-contacting process for simultaneous removal of NO <sub>x</sub> and SO <sub>x</sub> from plant stack.	Iron oxide supported on alumina as catalyst	[15]
<mark>12</mark>		Iron titanate catalyst+ $V_2O_5$ - $WO_3$ and Fe/Cu-Zeolite catalyst for SCR of NO <sub>x</sub> with NH <sub>3</sub> .	Iron titanate, Zeolite and NH₃	[16]
<mark>13</mark>		Catalytic reduction of NO with NH <sub>3</sub>	SO <sub>2</sub> /H <sub>2</sub> O. iron titanate as	[17]

			catalyst.	
14	CO2	Nanoparticle organic Hybrid materials (NOHMs) absorption of $CO_2$ was affected by the presence of $SO_2$ in the flue gas, that is reduction in $CO_2$ capture.	Nanoparticle organic Hybrid materials (NOHMs)	[9]
<mark>15</mark>		Absorption or adsorption	sorbent	<mark>[10]</mark>
<mark>16</mark>		Econoamine FG (Absorbents)	MEA	[11]
<mark>17</mark>	CO	Ambient content measured at rural agricultural site	Biomass burning tracer, CH <sub>3</sub> Cl; Industrial tracer, $C_2Cl_4$ .	[13]

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330 SOURCE: Summary from literature review above.

# TABLE 3 – ALTERNATIVE/ENHANCEMENT SOURCES TO GASOLINE GENERATORS S/No. ALTERNATIVE/ENHANCEMENT FUEL LIMITATIONS REFERENCE

<u>5/140.</u>	SOURCES TO GASOLINE GENRATORS		LIMITATIONS	
<mark>1</mark>	Micro Gas Turbo Generator at	<mark>Mixture of</mark>	High fuel	<mark>24</mark>
	University of Roma	Butane 8 1	consumption	
		and		
		Propane Propane		
		gas		
<mark>2</mark>	20MW Diesel Generator test	Vegetable	Cost and	<mark>26</mark>
		<mark>(Palm) oil</mark>	availability	
<mark>3</mark>	Mathematical Modeling of Biomass	<mark>coal</mark>	Flue gas	<mark>23</mark>
	Gasification		treatment not	
			squarely	
			addressed as	
			primary	
			objective	
<mark>4</mark>	Power Quality Management	-	Addressed the	<mark>25</mark>
			<mark>efficiency in</mark>	
			power usage	
			but silent on flue	
			gas managment	

# TABLE 4 – FUEL PRICES

<mark>S/No.</mark>	FUEL	PRICE	REFERENCE
<mark>1</mark>	<mark>Gasoline</mark>	USA = \$0.97/Litre; UK = \$2.18/Litre and	<mark>[28]</mark>
		Nigeria = \$0.58/Litre.	
<mark>2</mark>	gas	US\$3.41 per gallon	[29]
<mark>3</mark>	Palm oil (vegetable oil)	US\$612 per Metric Tonne	[27]

<mark>4</mark>	Butane Gas	<mark>4.5kg = £16.75 (or \$27.3)</mark>	<mark>[30]</mark>
	Propane Gas	<mark>3.9kg = £16 (or \$26.08)</mark>	

1 SOURCE: SUMMARY FROM LITERATURE REVIEW ABOVE

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344	

### RELATING RESEARCH EFFORTS TO LEGISLATION

Legislations, Standards and Regulations from the EU, USA and China as detailed in 345 Table 1 above are guite comprehensive in the regulating and control of flue gas 346 emissions into the atmosphere. On the other hand, the research and investigations into 347 the process, methodology and strategy for the removal of  $SO_x$ ,  $NO_x$ , CO and  $CO_2$  from 348 flue gases as detailed in Table 2 has been thorough, detailed and thought provoking. 349 References [8] to [20] have demonstrated various ways in capturing SO2, NOX, CO and CO<sub>2</sub> from flue gases. On the other hand, Reference [24] addressed mathematical model 350 351 of biomass gasification while Reference [26] investigated Power Quality Management 352 Protocol in residential buildings with co-generative capability, with the view of optimization of power usage. This report will be of immense benefit to developing 353 354 economies that are grappling and tinkering with the possibility of providing adequate power. In another development, References [24] and [26] investigated alternative fuel 355 356 regime (that is butane + propane gas and palm oil respectively) and generation methods 357 to gasoline generators. According to Reference [24], the fuel (that is butane + Propane) consumption by the 10KW micro gas turbo generator at Roma = 0.01kg/s, that is 358 359 0.01x60x60kg/hr = 36kg/hr. 360 From Table 4, 4.5kg of Butane = US\$27.3. 361 Therefore 0.5kg of Butane = (27.3/4.5)x0.5 = US\$3.03 362 Therefore 0.5kg of Propane = (26.08/3.9)x0.5 = US\$3.34. 363 Hence 1kg of gas (that is 0.5kg Butane + 0.5kg Propane) = US\$3.03 + US\$3.34 = 364 US\$6.37. 365 But the 10KW micro turbo generator consumes 36kg of mixed gas per hour. Hence the 366 cost of running the 10KW micro turbo generator per hour = 36XUS = US = 229.32. 367 On the other hand, according to Reference [31], a 10KVA package Gasoline Generator consumes 18litres of petrol in 4.5hours. This implies that this package generator 368 369 consumes (18litres)/4.5hrs = 4litres/hr. 370 But from Table 4, Reference [28] posits that a litre of petrol in Nigeria cost US\$0.58. 371 Hence the cost of running this 10KVA Package gasoline generator = 4x US\$0.58 = 372 US\$2.32.

Comparing the US\$2.32/hr for running this 10KVA package gasoline generator with the US\$229.32/hr for running the 10KW micro turbo generator at Roma, this translates to a cost ratio of 1 to 98.85. The outcome of the Roma project is therefore a very inefficient result against the balance of economics and the background of already available operating generator efficiency in the global market. Consequent on the foregoing, the outcome of the Roma project will not be of immediate benefit as a substitute for package gasoline generators in the developing economies.

- On the other hand, according to Reference [26], the testing of 20MW Diesel Generator 380 381 using vegetable oil (that is palm oil) as substitute has some merit. According to 382 Reference [27] from Table 4, the price of palm oil is US\$612per metric tonne. This is 383 approximately US\$0.612 per litre. Also, according to Reference [29], the price of diesel 384 is US\$3.41 per gallon. This is approximately US\$0.85 per litre. Since the price of palm 385 oil is lower than that of diesel, it makes economic sense to investigate the viability of palm oil as a substitute fuel for the diesel generator. However, the price of petrol in 386 387 Nigeria according to Reference [28] is US\$0.58 per litre. Against this backdrop, the use 388 of palm oil as a substitute fuel in Nigeria will amount to a bad economic decision.
- 389

### 390 It Can Generally Be Summarized That The Following Holds:

391 1. Generators are devices that convert mechanical energy to electrical energy.

- 3922. Power supply from National grid of most developing nations (Nigeria inclusive) is393 insufficient for economic activities.
- Consequent on above, most businesses rely on package generators for electrical
   power delivery.
- 396 4. All generators that are fossil fuel fired release flue gases into the environment.
- 397 5. Most flue gases contain SO<sub>2</sub>, NO<sub>X</sub>, CO and CO<sub>2</sub>. The CO is detrimental to health of
  398 living men and animals and at the same time depletes the ozone layer. On the
  399 other hand the CO<sub>2</sub> contributes to the greenhouse effect.
- 400 6. Consequent on the above, World Health and Environmental regulating bodies have
  401 taken initiatives to regulate and reduce the emissions of these gases to as low as
  402 reasonably possible (ALARP).
- 403 7. Efforts are being made to develop purifying/capturing methods for these green
  404 house gases while at the same time seeking alternative renewable energy sources
  405 to replace energy sources that generates the green house gases.
- 406 8. Most of the improvement efforts on energy sources that are safe and efficient in
  407 usage have been geared towards grid equipment such as turbo-generators, boilers,
  408 Industrial furnaces, geothermal, hydro and solar energy.
- 409 9. Identified methods of capture of CO and CO<sub>2</sub> include the absorption of the gases
  410 using either the wet or dry methods.
- 411 10. Different storage and discarding methods were identified as means of managing412 the captured gases.
- 413 The efforts and improvements made in the capture of  $SO_2$ ,  $NO_x$ , CO and  $CO_2$  were 414 never targeted at the gasoline package generators. This appears to be an oversight in 415 the sense that the package generators in the first instance were not actually designed as 416 full time generators. Rather, they were designed to serve as standby and emergency 417 units. But the acute deficit of power supply to economic businesses in developing nations and coupled with poor finances, have compelled the businesses to resort to 418 419 usage of package generators as main power source. This has led to ample generation of 420  $SO_2$ ,  $NO_x$ , CO and CO<sub>2</sub>, the ODs and GHGs abundantly in the developing countries. 421 Since the use of fossil fuel will remain with us for a long time to come, the need therefore 422 arises for improvement action to be taken in minimizing the emission of the OD and 423 GHGs from the package generators into the environment.
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# WHAT FURTHER IMPROVEMENTS NEED TO BE DONE?

From the foregoing, it is evident that much has been achieved in the Legislations, Regulation 426 427 and Control of flue gases. Much of the emphasis has been directed to the large combustion 428 plants (the "LCP directive"). The volume of research into the capture of  $SO_2$ ,  $NO_3$ , CO and  $CO_2$  from flue gas has been extensive. The implementation of the research outcomes have 429 430 been so far directed mainly to the LCPs to the detriment of the package generators. The 431 package generators will continue to be relevant in the scheme of economic and domestic 432 activities in the developing economies in as much as the national power grids continue to be 433 epileptic in their service delivery. Therefore, the attendant noise and flue (CO and CO<sub>2</sub> 434 inclusive) pollution will be with the society for a long time to come. The immediate and urgent 435 thing for now is to mitigate and embark on some control measures to limit this pollution to an 436 acceptable statutory level. Hence the need is here now more than ever to incorporate the 437 identified capture technique for SO<sub>2</sub>, NO<sub>x</sub>, CO and CO<sub>2</sub> into the package gasoline generators. Hence in the design and fabrication of the SO<sub>2</sub>, NO<sub>3</sub>, CO and CO<sub>2</sub> capture unit, 438 439 emphasis should be placed on the absorption of the gases using either the wet or dry 440 methods. Some form of sequencing should be done to align capturing process to discarding 441 process in order to optimize the overall system. The outcome of this project effort will keep 442 the environment safer and put more money in the pockets of business units, as the 443 environment becomes cleaner, neater and safer hence minimized environmental and 444 pollution litigations fines and expenses. Also the health and medical bills will drop.

### 446 CONCLUSION

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The impact of high concentration of CO, CO<sub>2</sub>, SO<sub>2</sub> and NO<sub>X</sub> to human life is lethal and 447 448 should not be treated with kid gloves. This severity index is more than enough justification 449 for action to be taken to minimize the attendant scenario that leads to the over generation of 450 these gases. The global position on legislation, regulation and control of Emission Limit 451 Values (ELV) of flue gases has been articulated. All efforts directed towards limiting the indiscriminate and uncontrolled generation of CO, CO<sub>2</sub>, SO<sub>2</sub> and NO<sub>x</sub> is a welcome 452 453 development. Various researchers have experimented on the means and 454 methods/techniques to capture and trap these gases. The incorporation of these capturing 455 methods in the package gasoline generators will go a long way in reducing the pollution of 456 the environment through flue gases. The suggested alternative sources of power as 457 substitute for the package gasoline generator by some of the researchers were found to be economically unviable. The review acknowledged the hazard levels of CO, CO<sub>2</sub>, SO<sub>2</sub> and 458 459 NO<sub>X</sub>.

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