Abstract: The present invention relates to a single-pot method for preparing dialkyl carbonates, the method comprises reaction of alkyne oxide with aliphatic or cyclic aliphatic alcohol, using wood ash catalyst, under CO₂ pressure and heating the reaction mixture thereof to obtain dialkyl carbonates.
SINGLE-POT SYNTHESIS OF DIALKYL CARBONATES USING CATALYST FROM NATURAL RESOURCE

FIELD OF THE INVENTION

The invention relates to synthesis of dialkyl carbonates from C0₂, aliphatic/cyclic aliphatic alcohol and alkyene oxide in presence of renewable catalyst in a single step. Particularly this invention relates to a catalyst prepared directly from biomass used in the synthesis of dialkyl carbonates by single-step preparation.

BACKGROUND OF THE INVENTION

C0₂ is a major contributor of climate change and worldwide containment of C0₂ is a major challenge. In current scenario the challenge for decreasing C0₂ in atmosphere needs new ideas and technologies. Generating value added chemicals through green chemistry especially by utilizing greenhouse gas carbon dioxide as raw material is a challenging task. Present fuel oxygenates like ether compounds methyl tert. butyl ether (MTBE), ethyl tert. butyl ether (ETBE), tert. Amyl methyl ether (TAME) and alcohols like methyl alcohol (CH₃OH), ethyl alcohol (C₂H₅OH) are being used as oxygenates. However, aforementioned oxygenates lead to many problems like corrosiveness, environmental harm, and solubility issues. In order to circumvent these problems dialkyl carbonates are alternate future fuel oxygenates exhibiting attractive environmentally benign nature like non-toxic, non-corrosive and low toxic emissions are being explored.

The dialkyl carbonates especially dimethyl carbonate (DMC), diethyl carbonate (DEC), and ethyl methyl carbonate (EMC) have been widely accepted as fuel oxygenates because of excellent gasoline blending properties like high blending octane numbers, low blending Reid vapour pressures (RVP) and lower amounts of toxic emissions compared to ether oxygenates. Several other applications of dialkyl carbonates were explored in green chemical industry for the preparation of polycarbonates, isocyanates, synthetic lubricants, pharmaceutical and agricultural intermediates. In addition, dialkyl carbonates are non-corrosive and highly efficient alkylating agents to replace hazardous phosgene. Due to high volatile value, they are widely used in paint industry.

Scientists have developed several processes for the synthesis of dialkyl carbonates. The conventional process involves usage of hazardous chemicals like phosgene or carbon
monoxide as starting materials. The alternative and non-hazardous process involve utilization of CO₂ as raw material, which is eco-friendly and cost-effective process.

The eco-friendly synthetic route involves reaction of carbon dioxide with alcohol to generate corresponding dialkyl carbonates. Amongst dialkyl carbonates, dimethyl carbonate and diethyl carbonate synthesis has been well recognised and explored. In prior art literature, number of homogeneous catalysts (e.g. tin complexes: distannoxanes) and heterogeneous catalysts (e.g. CeO₂, modified Zirconia, Zeolites) have been investigated and have shown limitations of poor conversions either due to the thermodynamic limitation of reaction or due to catalyst deactivation.

Therefore, finding an alternate route for the synthesis of DMC and DEC from CO₂ involving in situ transesterification of cyclic carbonate (synthesised from epoxide and CO₂) with alcohol in presence of suitable catalyst would be well attempted. Significant efforts have already been devoted in efficient catalyst development for DMC & DEC synthesis from cyclic carbonates. The synthesis of DMC and DEC from epoxide through two step mechanism involving synthesis of cyclic carbonate in 1st step and then transesterifying to DMC and DEC is not economical for commercialisation.

Keeping in view the limitation associated with above processes, one-pot synthesis of DMC/DEC from CO₂, alkylene oxide and alcohol was explored to make the process more viable, eco-friendly and cost-effective. Single-pot synthesis of DMC/DEC from CO₂, alkylene oxide and alcohol in presence of several catalysts mainly metal oxides, inorganic bases/metal oxides, inorganic bases/phosphonium halide on polyethylene glycol, TBAB/Et₃N, ionic liquid, KOH/4Å° MS, biomettalic Cu-Ni/4Å° MS, anion exchange resins, and Mg/ Smacilite etc. have been reported in the literature.

The process conditions in single step synthesis of DEC and DMC from propylene oxide in literature references are reported to involve low catalytic activity, formation of side products, difficulty in catalyst separation and recyclability, and requirement of costly raw materials to synthesise the above catalysts.

Several patents have been published on synthesis of dialkyl carbonates from transesterification of cyclic carbonates with appropriate alcohol in presence of both
homogeneous and heterogeneous catalysts. Few publications have been found on direct synthesis of dialkyl carbonates from alkylene oxide, \( \text{C}_2 \) and alcohol in single-pot reaction.

Single-pot synthesis of dialkyl carbonates is reported in US 4,434,105 (EP 000777), which reveals the synthesis of dialkyl carbonates namely dimethyl carbonate and diethyl carbonate from ethylene oxide/propylene oxide, methanol/ethanol and carbon dioxide in the presence of homogeneous as well heterogeneous catalysts in single pot reaction. Dialkyl carbonates mainly DMC and DEC were prepared in the presence of catalysts NaI/T\( \text{I}_2 \)C\( \text{O}_3 \), NaI/TIOH, Nal, T\( \text{I}_2 \)C\( \text{O}_3 \), Imidazole, TBAB, TEAB and organic bases like guanidine, DABCO, triethanol amine and etc. Yields of DMC and DEC are high with ethylene oxide compared to propylene oxide, which may be due to sterically more feasible product formation. The main drawback of this work is expensive catalysts and halogen containing catalysts.

US 5218,135 has revealed two step synthesis of dialkyl carbonate from C\( \text{O}_2 \), alkylene oxide and alcohol in presence of bifuctional catalysts. Initially cyclic carbonate was prepared from alkylene oxide and C\( \text{O}_2 \) in presence of bifuctional catalysts then cyclic carbonate was treated with alcohol in presence of bifuctional catalyst to yield corresponding dialkyl carbonate.

Recently US 2013/0267727 A1 and also US 2006/7084292 B2 have explored the feasibility of integrated process for production of dialkyl carbonates and diols from alkylene oxide, carbon dioxide and aliphatic alcohol. In this integrated process, initially alkylene oxide was reacted with carbon dioxide to produce crude cyclic carbonate in the presence of homogeneous catalyst. Second step involve reaction between crude cyclic carbonate and alcohol in the presence of heterogeneous catalysts like metal oxides to produce dialkyl carbonates.

US 5218,135, US 7491837 B2 and WO 03/000641 A1 also described the integrated process of making dialkyl carbonates for alkyl alcohol, carbon dioxide and alkylene oxide. They initially prepared cyclic carbonate from alkylene oxide and carbon dioxide in presence of homogeneous catalyst. In second process cyclic carbonate reacts with alcohol for producing dialkyl carbonate in the presence of transesterification catalyst.
The catalysts used in the prior art are expensive, non-renewable and non-recyclable. Formation of by-product H₂O during reaction deactivates the catalyst for further use. Also low product yields are reported in references of prior art.

In prior art literature, heterogeneous catalysed single pot synthesis of dialkyl carbonates from propylene oxide, C⁰₂ and methanol in presence of Mg-Smectite gave 32.30% of dimethyl carbonate (Green chemistry, 2003, 5, 71-75).

Present invention discloses a preparation methodology of dialkyl carbonates using wood ash catalyst; involving insertion of C⁰₂ molecule in to an alkylene oxide and transesterification of that intermediate compound with aliphatic alcohol in to dialkyl carbonate in single pot reaction. The inherent property of wood ash catalyst of C⁰₂ insertion along with the property of transestrification was explored for the synthesis of dialkyl carbonates. The present invention addresses one or more such problems of the prior art as discussed above. However, it is contemplated that the invention may prove useful in addressing other problems also in a number of technical areas.

**SUMMARY OF THE INVENTION**

The present invention relates to a single-pot method for preparing dialkyl carbonate, the method comprising dissolving an alkylene oxide in an aliphatic or cyclic aliphatic alcohol, adding a wood ash catalyst, adding C⁰₂ gas under pressure and heating the reaction mixture, cooling the reaction mixture to about room temperature, and depressurizing to obtain dialkyl carbonate. The reaction mixture is filtered to recover the catalyst. The aliphatic or cyclic aliphatic alcohols are Cᵢ to C₁₂ aliphatic alcohols and Cᵢ to C₁₂ cyclic aliphatic alcohols. The aliphatic or cyclic aliphatic alcohol is selected from methanol, ethanol, propanol, butanol, amyl alcohol, hexanol, octanol, cyclohexanol and octahexanol. The alkylene oxide is selected from the group comprising ethylene oxide, propylene oxide, 1,2-epoxy butane, 1,2-epoxy pentane and 1,2-epoxy hexane. The pressure is about 70-90 bar. The heating is carried out at a temperature range of about 100-180°C. The filtered catalyst is washed with an aliphatic alcohol and dried at about 120°C for 24 hours for reuse. The aliphatic alcohol is methanol or ethanol. The catalyst obtained is reused in the process of claim 1.
The present invention also relates to a method of preparing wood ash catalyst comprising washing wood with deionized water; drying the wood until the wood attains constant weight; dry ashing the wood; and calcining the wood to obtain wood ash catalyst.

The wood is selected from \textit{Azadirachta indica} and \textit{Acacia nilotica}. The wood is dried at a temperature of about 60° to 80°C. The dry ashing is carried out at a temperature of 525 ± 25°C. The calcining is carried out at a temperature of about 300°C to about 1200°C.

Our invention presents the wood ash catalyst BWCsoo as heterogeneous catalyst for the synthesis of dimethyl carbonate from propylene oxide, CO\textsubscript{2} and methanol with good yield of 51.11% which is better than reported in literature. Activity of wood ash catalyst BWCsoo was also further evaluated with several alcohols for the synthesis of respective dialkyl carbonates. Amongst, diethyl carbonate was synthesised from propylene oxide, CO\textsubscript{2} and ethanol with highest yield of 53.43%.

The following terms are defined as follows:

\textbf{BW}o: Wood ash without calcination. On using BW\textsubscript{o} as catalyst for the preparation of dialkyl carbonates, the overall yield is 6-7%. The reason for the low yield of dialkyl carbonates is that wood ash contains primarily carbonates of Ca, Mg along with other metal components and less basic in nature. It also does not contain sintered calcium silica phosphate compound. So, the dialkyl carbonate yield is very less in the reaction with BW\textsubscript{o}.

\textbf{BWC500}: Wood ash calcined at temperature 500°C. On using BWC500 as catalyst for the preparation of dialkyl carbonates, the overall yield is 42.46%. The reason for the moderate yield is the partial conversion into active catalyst such as some amount of calcium oxide (CaO) and magnesium oxide (MgO) along with small amount of calcium silica phosphates were observed in XRD studies. Accordingly conversion to dialkyl carbonate is not complete and moderate.

\textbf{BWCsoo}: Wood ash calcined at temperature 800°C. On using BWCsoo as catalyst for the preparation of dialkyl carbonates, the overall yield is 53.43%. The reason for the high yield is that catalyst from wood ash at 800°C contains completely converted magnesium oxide (MgO)
and calcium oxide (CaO) along with sintered calcium silica phosphates which gave highest
dialkyl carbonate yield.

BRIEF DESCRIPTION OF ACCOMPANYING DRAWINGS

Figure 1: IR spectrum of wood ash catalysts;
Figure 2: XRD spectrum of wood ash catalysts; and
Figure 3: TGA analysis of BWcsoo wood ash catalyst.

DETAILED DESCRIPTION OF THE INVENTION

The present invention discloses a preparation methodology of dialkyl carbonates (fuel
oxygenates) by reacting C0₂, aliphatic/cyclic aliphatic alcohol and alkyene oxide in presence
of renewable catalyst in a single step, the said catalyst being prepared from biomass.

An aspect of the invention discloses the preparation methodology of the catalyst; said catalyst
being prepared directly from the biomass. Another aspect of the present invention is the use
of the novel renewable catalyst prepared from biomass in a single step reaction to produce
dialkyl carbonates from aliphatic & cyclic alcohols and propylene oxide by utilizing C0₂.

An aspect of the present invention is that the catalyst prepared from wood ash is novel,
economical and eco-friendly. The wood ash used in accordance with this invention can be
obtained from biomass, including but not necessarily limited to, wood of trees such as
Azadirachta indica, Acacia nilotica. Another aspect of the invention discloses the production
of alkylene glycol as by-product during synthesis of dialkyl carbonates in presence of
renewable catalyst.

The wood ash catalyst is basic in nature. Composition of wood ash catalyst is mixture of
oxides of Ca and Mg along with sintered material calcium silica phosphates. Wood ash also
contains potassium and small quantities of other metal derivatives. The typical combination
of all these compounds in wood ash catalyst makes it a suitable catalyst for various organic
transformations. We explored the inherent properties of wood ash catalyst and utilized for the
single pot synthesis of dialkyl carbonates from C0₂, alkylene oxide and alcohol. The wood
ash catalyst catalyses C0₂ insertion in to alkylene oxide to form cyclic carbonate which is
further converted in situ to dialkyl carbonates in the presence of alcohols. Therefore, wood
ash catalyst helps to convert C0₂ to dialkyl carbonates in the presence of alcohols in a single
pot reaction conditions in comparison to prior art claim of dialkyl carbonates synthesis from CO$_2$ mostly carried out in two separate steps by using two different types of catalysts. The advantage of this invention is wood ash catalyst able to perform synthesis of dialkyl carbonates from CO$_2$ in a single pot reaction conditions.

The preparation methodology according to the present invention involves washing the wood with deionised water and then drying until the wood attains constant weight. The wood was dry ashed separately and calcined. One-pot synthesis of dialkyl carbonates from aliphatic/cyclic aliphatic alcohol, CO$_2$ and alkyne oxide was carried out in a controlled heating system.

In a preferred embodiment, the preparation of dialkyl carbonate involves washing the wood with deionised water and then drying at about 60°-80°C until the wood attains constant weight. The wood was dry ashed separately and calcined, at temperature ranges of about 300°C to about 1200°C. One-pot synthesis of dialkyl carbonates from aliphatic/cyclic aliphatic alcohol, CO$_2$ and alkyne oxide was carried out in an autoclave vessel fixed to stirrer and controlled heating system. In drying step, the wood is heated at a temperature range of 60 to 80°C for 24 hours to remove moisture and impurities present in the wood after washing with deionised water. In the dry ashing step, the wood is burned in a furnace at 525°C ± 25°C (Ref. TAPPI, Ash in wood, pulp, paper and paperboard: combustion at 525°C, T211 om-93, 1993).

The aliphatic alcohols/cyclic alcohols used according to the present invention include C$_1$ to C$_{12}$ aliphatic alcohols, Ci to C$_{12}$ cyclic aliphatic alcohols like cyclic pentanol, cyclic hexanol, more preferably methanol, ethanol, propanol, butanol, amyl alcohol, hexanol, octanol, cyclohexanol, octahexanol. The alkyne oxides used according to the present invention include ethylene oxide, propylene oxide, 1,2-epoxy butane, 1,2-epoxy pentane, 1,2-epoxy hexane.

The single-pot method for preparing dialkyl carbonate comprises dissolving alkylene oxide in an aliphatic/cyclic aliphatic alcohol, adding wood ash catalyst, adding CO$_2$ gas at a workable pressure range and heating the reaction mixture, cooling the reaction mixture to about room temperature, and depressurizing to obtain dialkyl carbonate. The reaction mixture is filtered
to remove the catalyst. The filtered catalyst was washed with an aliphatic alcohol, preferably methanol or ethanol and dried for reuse.

In a preferred embodiment, the single-pot method for preparing dialkyl carbonate comprises dissolving alkylene oxide in an aliphatic/cyclic aliphatic alcohol, adding wood ash catalyst in the autoclave vessel. The autoclave vessel was pressurised with CO₂ gas. At the workable pressure range of about 70-90 bar and temperature range of about 100-180°C, the reaction mixture was stirred constantly for about 24 hours. Then reaction mixture was cooled to room temperature, depressurised and filtered to remove the catalyst. The filtered catalyst was washed with an aliphatic alcohol, preferably methanol or ethanol and dried at about 120°C for about 24 hours for reuse. The identification and quantification of the components in the reaction mixtures were performed with GC and GC-MS analysis.

Catalyst prepared from wood ash is novel, economical and eco-friendly compared to catalysts reported in literature for the synthesis of dialkyl carbonates. The developed process methodology is single step reaction to produce dialkyl carbonates from aliphatic/cyclic alcohols and propylene oxide by utilizing CO₂. Separation and regeneration of catalyst is performed by low cost techniques.

The CO₂ insertion reaction into propylene oxide requires activation of epoxide and activation of CO₂, which is the most crucial step. This is pull-push of electrons of both the molecules by two different categories of metals. Wood ash catalyst has combination of the above properties due to presence of metal oxides, mixed metal oxides along with alkali earth metal halides. Both alcohol and CO₂ are equally capable to insert into epoxide ring but catalyst should be selective for CO₂ insertion. Wood ash catalyst showed typical interaction with CO₂ and activated it successfully towards epoxide insertion reaction. Prior art literature shows that oxides of Ca and Mg can perform transesterification reactions of propylene carbonate with alcohols (Indian Journal of Chemistry, 52A, 459-466, 2013). Wood ash catalyst able to perform both CO₂ insertion and transesterification reactions simultaneously with high selectivity.

The prepared catalysts were evaluated for their alkalinity (pH), particle size and surface area, results are shown in Table 1. The alkalinity of BW₀ (wood ash without calcination), BWC₅₀ (wood ash calcined at 500°C) and BWC₈₀₀ (wood ash calcined at 800°C) catalysts were found
to be 11.28, 11.68 and 12.02 respectively. It was observed that the calcination temperature affects the alkalinity of the catalysts. The results showed that BWCSoo catalyst calcined at 800°C temperatures has highest alkalinity may be due to thermal decomposition of CaCO$_3$ to CaO which is having higher soluble alkalinity. This observation is in line with that reported in literature, which indicates the increase of alkalinity with the increase of calcination temperature [Biomass and Bioenergy 41 (2012) 94-106]. Further, recovered BWCSoo catalyst alkalinity was determined as 12.04 in order to check the stability of active basic sites present on the catalyst. In addition, XRD pattern of recovered catalyst also confirms the stable catalyst structure after the reaction (Figure 2).

Table 1: Alkalinity, surface area and particles size of the wood ash catalysts

<table>
<thead>
<tr>
<th>Catalyst</th>
<th>Alkalinity (at 28.0 °C)</th>
<th>SA (m$^2$/gm)</th>
<th>APS (µm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>BW$ _0$</td>
<td>11.28</td>
<td>2</td>
<td>20.76</td>
</tr>
<tr>
<td>BWCS500</td>
<td>11.688</td>
<td>1.6</td>
<td>25.65</td>
</tr>
<tr>
<td>BWCS800</td>
<td>12.025</td>
<td>&lt;1</td>
<td>36.26</td>
</tr>
<tr>
<td>BWCS800(Recovered)</td>
<td>12.048</td>
<td>8</td>
<td>56.77</td>
</tr>
</tbody>
</table>

As seen from the Table 1, the surface area of wood ash catalysts decreases with increase of calcination temperature. The surface areas of BW$ _0$, BWCS500 and BWCS800 catalysts were 2.0, 1.6 and < 1 m$^2$/gm respectively. Lower surface area of BWCS800 catalyst than BWCS500 catalyst is due to conversion of CaCO$_3$ into CaO which results in sintering by the formation of calcium phosphate silicate. The observation was further confirmed by XRD study of the catalyst structure (Figure 2).

The particle sizes of BW$ _0$, BWCS500 and BWCS800 were determined as 20.76, 25.65 and 36.26 µm respectively. The result reveals that particle size of the catalysts has increased with increasing calcination temperature [Biomass Bioenergy 4 (1993) 103-106]. The possible reason may be the increase of sintering with increase of temperature, leads to increase in particle size as reported in earlier study [Biomass and Bioenergy 41 (2012) 94-106]. Recovered BWCS800 catalyst particle size is 56.77 µm may be due to the formation of some amounts of calcium hydroxide and carbonates however this does not has any effect on
catalyst performance on subsequent reuse. These results were further supported by XRD analysis as shown in Figure 2.

**IR analysis**

The FTIR spectra of wood ash catalysts are given in Figure 1. The spectra of BWo catalyst shows the presence of carbonates, C0₃²⁻ (the peaks at 1795, 1427, 875 and 711 cm⁻¹) and phosphates, P0₄³⁻ components (peaks at 1112, 1047 and 617 cm⁻¹), confirms the presence of metal carbonates (mainly CaC0₃) and metal phosphates. The IR spectrum of BWC₅₀₀ catalyst shows similar spectral features. After calcination at 800°C, BWC₈₀₀ catalyst indicates the presence of metal oxides (peak at 3425, 2960, 1462, 1408, 921 and 516 cm⁻¹) along with the calcium phosphates silicate (peak at 2856, 1388 and 1118 cm⁻¹). The results reveal that the calcination at higher temperature results in carbonates decomposition to metal oxides.

IR analysis of recovered BWC₈₀₀ catalyst reveals that structure of the catalyst is remain same after reaction however formation of very small quantity of calcium hydroxide (3645 cm⁻¹) was observed.

**XRD analysis**

The XRD pattern of BW0, BWC₅₀₀, BWC₈₀₀ and recovered BWC₈₀₀ catalyst is shown in Figure 2. The results indicate that BWo catalyst is mainly composed of CaC0₃ along with small amount of KC1 and SiO₂. After calcination at 500°C, the pattern of crystalline phases remains same in BWC₅₀₀ catalyst with additional appearance of CaO crystalline phase.

On further calcination at 800°C, BWC₈₀₀ catalyst shows the presence of metal oxides mainly CaO and MgO along with KC1 and mixed metal phosphates Ca₂SiO₄₀.05Ca₃(PO₄)₂. These results were further confirmed by elemental analysis.

XRD analysis of recovered BWC₈₀₀ catalyst show same structure as of fresh BWC₈₀₀ catalyst. However some decreasing intensity of crystalline phases of CaO, MgO and mixed metal phosphates is due to formation of calcium hydroxide is confirmed by increasing of pH value and IR analysis.

**Elemental Analysis**

The elemental analysis of BWC₈₀₀ catalyst shows the presence of Ca, K and Mg in higher
amounts as 32.30, 10.91 and 5.79 mass fraction percentage respectively. While the Na, P and Al are in notable amounts as 1.42, 1.40 and 1.90 mass fraction percentage respectively, the transition metals present in BWCsoo are mainly Fe, Mn and Zn in trace quantities. The presence of the minor quantities of Silicon was also confirmed by XRF analysis.

TGA Analysis

In heterogeneous catalysts, stability of active sites on catalyst surface is very important. BWCsoo catalyst stability was measured by TG analysis carried out on TG model 2950 Hi Resolution modulated TGA, with heating rates 10°C/min, temperature ramp up to 800°C.

The results given in the Figure 3 shows the weight loss of 2.19 wt. % is observed at 198°C corresponding to the removal of loosely absorbed water on the surface of catalyst. Further the decomposition of calcium carbonate at 425°C and 660°C producing calcium oxide associated with 1.809 wt. % weight loss. Additionally, the stability of BWC80 catalyst is confirmed by metal analysis of carbonate product produced with XRF technique and found all the metal <1 mg/Kg. Above findings also confirm the non-leaching behaviour of the catalyst and are in agreement with the earlier study [Biomass and Bioenergy 41 (2012) 94-106].

These dialkyl carbonates are extensively studied as fuel oxygenates in the literature (Energy & Fuels 1997, 11, 2-29). The octane number, RVP and toxic emissions from dialkyl carbonates are compared with ether oxygenates and reported in the literature (Energy Fuels 2010, 24, 4812-4819). However, synthesis of these dialkyl carbonates reported through phosgene route which is corrosive and environmentally not favourable. We have developed environmentally safe method for preparation of dialkyl carbonates from CO2 as feed stock and carrying out the preparation in single pot in presence of wood ash as novel catalyst.

The octane number of prepared dialkyl carbonates viz. DMC, DEC blend with gasoline is studied by CFR (Combustion Fuel Research)* engine test method, the results are as follows in Table 2:

<table>
<thead>
<tr>
<th>S. No:</th>
<th>FUEL TYPE</th>
<th>RON by CFR engine test ASTM D2699</th>
<th>Improved RON value</th>
<th>RVP values (KPa)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Blending carried out to keep overall oxygen content in gasoline to 2.7% by mass as per IS 2796-2008 motor gasoline BS IV specifications.

These dialkyl carbonates are extensively studied as fuel oxygenates in the literature (Energy & Fuels 1997, 11, 2-29). The octane number, RVP and toxic emissions from dialkyl carbonates were compared with ether oxygenates and reported in the literature (Energy Fuels 2010, 24, 4812-4819). The synthesis of these dialkyl carbonates through phosgene route is corrosive and environmentally not favourable. We have developed environmentally safe method from CO₂ as feed stock and single pot reaction in presence of wood ash as catalyst.

Having described the basic aspects of the present invention, the following non-limiting examples illustrate specific embodiment thereof.

**EXAMPLES**

**Example 1: One-pot synthesis of DEC from Ethanol, CO₂ and Propylene oxide using BWo as catalyst**

In a 700 ml autoclave, propylene oxide (10.17 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash catalyst BWo (7.0 gm, 10% wt/wt) were charged. The autoclave vessel was pressurised with CO₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring for reaction time 24 hours. Then reaction mixture was cooled to room temperature, depressurised and filtered to remove the catalyst. The filtered catalysts was washed with ethyl alcohol and dried at 120°C for 24 hours before reuse. The reaction mixture was subjected to distillation. First fraction contains mixture of excess ethanol and DEC and second fraction contains mixture of propylene glycol and propylene carbonate. The
identification and quantification of the components in the fractions were performed with GC and GC-MS analysis.

Analytical analysis of ethanol fraction reveals that formation of DEC is 1.26 gm (2.20 mmol). The calculated yield of DEC corresponding to propylene oxide is 6.06%.

Example 1 explains that reaction mixture was cooled to room temperature after 24 hours, depressurised and filtered to remove the catalyst. The filtered catalyst was washed with methanol (20 ml) dried at 120°C for 24 hours for reuse.

**Example 2: Synthesis of DEC from Ethanol, CO₂ and Propylene oxide using BWC 500 as catalyst**

Propylene oxide (10.17 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash catalyst BWC 500 (7.0 gm, 10% wt/wt) were charged into 700 ml autoclave vessel. The autoclave vessel was pressurised with CO₂ pressure, heated to 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 1.

The quantitative analysis of ethanol fraction reveals that formation of DEC is 8.79 gm (74.47 mmol). The calculated yield of DEC with respect to propylene oxide is 42.46%.

**Example 3: Synthesis of DEC from Ethanol, CO₂ and Propylene oxide using 10 wt % BWC 500 as catalyst**

700 ml autoclave was charged with propylene oxide (10.17 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash catalyst BWCsoo (7.0 gm, 10% wt/wt). The autoclave vessel was pressurised with CO₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Then reaction mixture was cooled to room temperature, depressurised and filtered to remove the catalyst. The filtered catalysts was washed with ethyl alcohol and dried at 120°C for 24 hours before reuse. The reaction mixture was subjected to distillation. First fraction contains mixture of excess ethanol and DEC and second fraction contains mixture of propylene glycol and propylene carbonate. The identification and quantification of the components in the fractions were performed with GC and GC-MS analysis.
The quantitative analysis of ethanol fraction reveals that formation of DEC is 11.06 gm (93.67 mmol). The calculated yield of DEC from propylene oxide is measured as 53.43%.

Example 4: Synthesis of DEC from Ethanol, C0₂ and Propylene oxide using BWC₉₀₀ as catalyst at 70 bar C0₂ pressure

In a 700 ml autoclave, propylene oxide (10.17 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash catalyst BWC₉₀₀ (7.0 gm, 10% wt/wt) was charged. The autoclave vessel was pressurised with C0₂ pressure, heated to 150°C temperature and 70 bar pressure under constant stirring for reaction time 24 hours. Reaction mixture work up was carried out as described in example 3.

The analytical analysis of ethanol fraction reveals the formation of DEC is 10.23 gm (86.65 mmol). The calculated yield of DEC corresponding to propylene oxide is 49.42%.

Example 5: Synthesis of DEC from Ethanol, C0₂ and Propylene oxide using BWC₉₀₀ as catalyst at 90 bar C0₂ pressure

Propylene oxide (10.17 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash catalyst BWC₉₀₀ (7.0 gm, 10% wt/wt) were charged in to 700 ml autoclave vessel. The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature and 90 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 3.

Analytical analysis of ethanol fraction reveals that formation of DEC is 8.78 gm (74.38 mmol). The measured yield of DEC from propylene oxide is 42.41%.

Example 6: Synthesis of DEC from Ethanol, C0₂ and Propylene oxide using BWC₉₀₀ as catalyst at 80 bar C0₂ pressure and 120 °C temperature.

Propylene oxide (10.17 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash catalyst BWC₉₀₀ (7.0 gm, 10% wt/wt) were charged in to 700 ml autoclave vessel. The autoclave vessel was pressurised with C0₂ pressure, heated to reach 120°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 3.
Analytical analysis of ethanol fraction reveals that formation of DEC is 5.08 g (43.05 mmol). The calculated yield of DEC corresponding to propylene oxide is 24.54%.

**Example 7: Synthesis of DEC from Ethanol, C0₂ and Propylene oxide using 5 wt % BWCsoo as catalyst**

In a 700 ml autoclave vessel added propylene oxide (10.17 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash catalyst BWCsoo (3.5 gm, 5% wt/wt). The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 3.

Analytical analysis of Ethanol fraction had DEC of 5.49 gm (54.86 mmol). The formation yield of DEC with respect to propylene oxide is 26.52%.

**Example 8: Synthesis of DEC from Ethanol, C0₂ and Propylene oxide using 15 wt % BWCsoo as catalyst**

Propylene oxide (10.17 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash catalyst BWCsoo (10.5 gm, 15% wt/wt) were charged into 700 ml autoclave vessel. The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 3.

The analytical analysis of ethanol fraction revealed that formation of DEC is 5.27 gm (44.66 mmol). The calculated yield of DEC corresponding to propylene oxide is 25.45%.

**Example 9: Synthesis of DEC from Ethanol, C0₂ and Propylene oxide using 10 wt % of recovered BWCsoo from example 3 as catalyst**

Propylene oxide (10.17 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash recovered catalyst BWCsoo from example 3 (10.5 gm, 15% wt/wt) were charged into 700 ml autoclave vessel. The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 3.
The analytical analysis of ethanol fraction reveals that formation of DEC is 9.85 gm (83.50 mmol). The measured yield of DEC from propylene oxide is 47.6%.

Example 9 shows the recovered catalyst used second time for the monitoring DEC synthesis under standardised reaction conditions. The DEC yield with second time recovered catalyst is slightly lower 47.6% than the previous results. Third cycle usage of wood ash catalyst gave 46.5% of DEC yield.

**Example 10: Synthesis of DEC from Ethanol, C0₂ and Propylene oxide using BWC₈₀₀ as catalyst at 80 bar C0₂ pressure and 180°C temperature**

In a 700 ml autoclave, propylene oxide (10.17 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash catalyst BWC₈₀₀ (7.0 gm, 10% wt/wt) was charged. The autoclave vessel was pressurised with C0₂ pressure, heated to 180°C temperature and 80 bar pressure under constant stirring for reaction time 24 hours. Reaction mixture work up was carried out as described in example 3.

The analytical analysis of ethanol fraction reveals the formation of DEC is 9.86 gm (83.56 mmol). The calculated yield of DEC corresponding to propylene oxide is 47.86%.

**Example 11: Synthesis of DEC from Ethanol, C0₂ and Ethylene oxide using BWC₈₀₀ as catalyst at 80 bar C0₂ pressure and 150°C temperature**

700 ml autoclave was charged with ethylene oxide (7.7 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash catalyst BWC₈₀₀ (7.0 gm, 10% wt/wt). The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 1. The quantitative analysis of ethanol fraction reveals that formation of diethyl carbonate is 10.33 gm (87.54 mmol). The calculated yield of diethyl carbonate from ethylene oxide is measured as 50.07%.

**Example 12: Synthesis of DMC from Methanol, C0₂ and Ethylene oxide using BWC₈₀₀ as catalyst at 80 bar C0₂ pressure and 150°C temperature**

Ethylene oxide (7.7 gm, 0.175 mol), methanol (44.80 gm, 1.40 mol) and wood ash catalyst BWC₈₀₀ (5.5 gm, 10% wt/wt) were taken in to 700 ml autoclave vessel and closed tightly. The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature...
and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 1. The quantitative analysis of methanol fraction reveals that formation of dimethyl carbonate is 8.20 gm (91.11 mmol). The calculated yield of dimethyl carbonate from ethylene oxide is measured as 52.13%.

Example 13: Synthesis of DEC from Ethanol, C0₂ and 1,2-epoxyhexane using BWCsoo as catalyst at 80 bar C0₂ pressure and 150°C temperature
The reactants mixture of 1,2-epoxyhexane (17.53 gm, 0.175 mol), ethanol (64.70 gm, 1.40 mol) and wood ash catalyst BWC₈₀₀ (8.25 gm, 10% wt/wt) was charged into autoclave vessel and lid was closed. The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 1. The quantitative analysis of ethanol fraction reveals that formation of diethyl carbonate is 4.35 gm (36.86 mmol). The calculated yield of diethyl carbonate from epoxyhexane is measured as 21%.

Example 14: Synthesis of DMC from Methanol, C0₂ and 1,2-epoxyhexane using BWCsoo as catalyst at 80 bar C0₂ pressure and 150°C temperature
1,2-epoxyhexane (17.53 gm, 0.175 mol), methanol (44.80 gm, 1.40 mol) and wood ash catalyst BWC₈₀₀ (6.23 gm, 10% wt/wt) were charged into 700 ml autoclave vessel and tightly closed. The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 1. The quantitative analysis of methanol fraction reveals that formation of dimethyl carbonate is 4.42 gm (49.00 mmol). The calculated yield of dimethyl carbonate from epoxyhexane is measured as 28.06%.

Example 15: Synthesis of Dicyclohexyl Carbonate from cyclohexanol, C0₂ and 1,2-epoxyhexane using BWC₈₀₀ as catalyst at 80 bar C0₂ pressure and 150°C temperature
700 ml autoclave was charged with 1,2-epoxyhexane (17.53 gm, 0.175 mol), cyclohexanol (140.81 gm, 1.40 mol) and wood ash catalyst BWC₈₀₀ (15.8 gm, 10% wt/wt). The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 1. The quantitative analysis of reaction mixture
reveals that formation of dicyclohexyl carbonate is 7.16 gm (31.54 mmol). The calculated yield of dicyclohexyl carbonate from epoxyhexane is measured as 18%.

Example 16: Synthesis of DMC from methanol, C0₂ and propylene oxide using BWC₈₀₀ as catalyst at 80 bar C0₂ pressure and 150°C temperature

The mixture of propylene oxide (10.17 gm, 0.175 mol), methanol (44.80 gm, 1.40 mol) and wood ash catalyst BWC₈₀₀ (5.5 gm, 10% wt/wt) was taken into 700 ml autoclave vessel and closed with lid. The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 1. The quantitative analysis of methanol fraction reveals that formation of dimethyl carbonate is 8.05 gm (89.5 mmol). The calculated yield of dimethyl carbonate from propylene oxide is measured as 51.11%.

Example 17: Synthesis of didecyl carbonate from n-decanol, C0₂ and propylene oxide using BWC₈₀₀ as catalyst at 80 bar C0₂ pressure and 150°C temperature

700 ml autoclave was charged with propylene oxide (10.17 gm, 0.175 mol), n-decanol (221.60 gm, 1.40 mol) and wood ash catalyst BWC₈₀₀ (23.0 gm, 10% wt/wt). The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 1. The quantitative analysis of reaction mixture reveals that formation of didecyl carbonate is 19.80 gm (57.89 mmol). The calculated yield of didecyl carbonate from propylene oxide is measured as 33.0%.

Example 18: Synthesis of dicyclohexyl carbonate from cyclohexanol, C0₂ and propylene oxide using BWC₈₀₀ as catalyst at 80 bar C0₂ pressure and 150°C temperature

Propylene oxide (10.17 gm, 0.175 mol), cyclohexanol (140.81 gm, 1.40 mol) and wood ash catalyst BWC₈₀₀ (15.0 gm, 10% wt/wt) were taken into 700 ml autoclave vessel and closed tightly. The autoclave vessel was pressurised with C0₂ pressure, heated to reach 150°C temperature and 80 bar pressure under constant stirring during the reaction for 24 hours. Reaction mixture work up was carried out as described in example 1. The quantitative analysis of reaction mixture reveals that formation of dicyclohexyl carbonate is 9.55 gm (42.0 mmol). The calculated yield of dicyclohexyl carbonate from propylene oxide is measured as 24%.
We Claim:

1. A single-pot method for preparing dialkyl carbonate, the method comprising:
   (a) dissolving an alkylene oxide in an aliphatic or cyclic aliphatic alcohol,
   (b) adding a wood ash catalyst,
   (c) adding CO₂ gas under pressure and heating the reaction mixture,
   (d) cooling the reaction mixture to about room temperature, and depressurizing to obtain dialkyl carbonate.

2. The method as claimed in claim 1, wherein the reaction mixture is filtered to recover the catalyst.

3. The method of claim 1, wherein the aliphatic or cyclic aliphatic alcohols are C₁ to C₂ aliphatic alcohols and C₆ to C₁₂ cyclic aliphatic alcohols.

4. The method of claim 2, wherein the aliphatic or cyclic aliphatic alcohol is selected from methanol, ethanol, propanol, butanol, amyl alcohol, hexanol, octanol, cyclohexanol and octahexanol.

5. The method of claim 1 where the alkylene oxide is selected from the group comprising ethylene oxide, propylene oxide, 1,2-epoxy butane, 1,2-epoxy pentane and 1,2-epoxy hexane.

6. The method of claim 1 wherein the pressure is about 70-90 bar.

7. The method of claim 1 wherein the heating is carried out at a temperature range of about 100-180°C.

8. The method of claim 2 wherein the filtered catalyst is washed with an aliphatic alcohol and dried at about 120°C for 24 hours for reuse.

9. The method of claim 8 wherein the aliphatic alcohol is methanol or ethanol.

10. The method of claim 8, wherein the catalyst obtained is reused in the process of claim 1.
Figure 3
INTERNATIONAL SEARCH REPORT

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

C07C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, CHEM ABS Data, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

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Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:

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Date of the actual completion of the international search: 2 April 2015

Date of mailing of the international search report: 16/04/2015

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