CYCLISATION-ACYETATION KINETIC OF (R)-(+) -CITRONELLAL BY Zn$^{2+}$-NATURAL ZEOLITE AS SOLID SOLVENT CATALYST

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ABSTRACT

Kinetic in cyclisation-acyetlation of (R)-(+) -citronellal with acetic anhydride was investigated over Zn$^{2+}$-Natural zeolite (Zn$^{2+}$-Natzeo) as a catalyst. (R)-(+) -citronellal has been isolated from citronella oil by fractional distillation under reducing pressure. Enantioselective capillary GC on a Supelco β-DEX 225 column has been used for analysis the enantiomers ratio of citronellal. Catalyst Zn$^{2+}$-Natzeo has prepared through acid activation of natural zeolite from Malang using HF 1% and HCl 6 M, followed by ion-exchange with 3 M NH$_4$Cl and calcination at 450 °C for 1 h under nitrogen to obtained H-natural zeolite (H-Natzeo). H-Natzeo was modified to Zn$^{2+}$-Natzeo by ion exchange with 0.1 M ZnCl$_2$. Cyclisation-acyetlation reaction was carried out by heating (R)-(+) -citronellal (CIT), acetic anhydride (AA), and 1 g catalyst in glass batch reactor with vigorous stirring at 80 °C. Molar ratio CIT/AA that used, i.e. 0.25; 0.5; 1.0; 1.2 and 1.5. As the reaction proceeded, 1 mL sample was taken off at 10; 20; 30; 60; 120; 180 min and extracted using n-hexane for every molar ratio. Structure analysis of product was conducted by GC-MS. Kinetic of the cyclisation-acyetlation reaction was analyzed according to the Langmuir-Hinshelwood mechanism. Increasing molar ratio of CIT/AA will decrease the isopulegyl acetate (IPA) and neo-isopulegyl acetate (NIIPA) formation. Rate constant of cyclisation-acyetlation reaction catalyzed by Zn$^{2+}$-Natzeo was 30.964 - 47.619 mmol(min. g cat)$^{-1}$ at 80 °C, 30 min and the ratio adsorption equilibrium constant $K_{eq}$/K$_A$ was 7.09.

Keywords: Cyclisation-acyetlation, (R)-(+) -citronellal, Zn$^{2+}$-natural zeolite, kinetic

INTRODUCTION

Citronellal is a monoterpen in citronella oil (Cymbopogon winterianus-Java type) that has two optical isomer with molecular weight of 154.25. The reactivity of citronellal is located on carbonyl group, double bond, and the acidity of α-hydrogen. It make possible for citronellal to react with either acid or base. The cyclisation of citronellal over zeolite catalysts and mesoporous materials in the cyclohexane under nitrogen was studied [1-3]. The highest rate of reaction was observed on mesoporous materials and 12 ring zeolite with concentrated Brensted acid, whereas the lowest rate of cyclisation occurred on silica and was not affected by the Brensted acidity. The high selectivity of cyclisation was observed on all catalysts used. The stability of carbocation showed the relationship with stereoselectivity of cyclisation. The analogue stereoselectivity of reaction was observed on the substrate of citronellal racemic mixture and pure enantiomer (+)-citronellal. The racemic mixture produced 8 different isopulegols, and (+)-citronellal only produced 4 pulegols [4-6].

Isopulegyl acetate (IPA) is a fragrance and flavor compound that can synthesized by one step cyclisation-acyetlation method from citronellal or two step cyclisation of citronellal and acetylation of the cyclisation product. Acid-catalyzed in the carboxylic acid esterification and Friedel-Crafts acylation reactions are important synthetic reactions for the preparation of key organic value-added intermediates. Preparation of alkyl esters is of particular importance in the manufacture of intermediates for the fragrance and pharmaceuticals industries. In normal practice, strong mineral acids, such as H$_2$SO$_4$, HF, or supported Lewis-acid catalysts like anhydrous AlCl$_3$/SiO$_2$ and BF$_3$/SiO$_2$, are used for such reactions. However, the use of corrosive mineral acids or Lewis-acid catalysts is associated with a number of environmental problems [7]. A major drawback of such catalytic systems is that they cannot be regenerated, and for Friedel-Crafts acylation more than stoichiometric amounts of catalyst are needed. In addition to this, trouble some work-up steps are also necessary for treating the liberated hazardous and toxic waste products. To avoid major environmental hazards and to satisfy growing stringent

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Fig 1. Cyclisation-acetylation of (R)-(+-)citronellal isopulegyl acetate isomers

global environmental regulations, it is imperative to develop a truly catalytic process with minimal waste production. Based on the formation of isopulegyl stereoisomers from citronellal cyclisation, assumed that the cyclisation-acetylation of racemic mixture of citronellal produced 8 stereoisomers of IPA, and enantiomer (+)-citronellal only produced 4 stereoisomers of IPA [9] (Fig. 1).

We have studied transformation of (R)-(+-)citronellal from Indonesian Citronella oil to produce stereoisomers of IPA by Lewis acid and modified natural zeolite catalysts. Isolation and identification of the product showed that the main compound is the mixture of IPA and its stereoisomer neo-isopulegyl acetate (NIPA) [8-10]. Stereoselectivity of citronellal cyclisation-acetylation will describe by study of kinetic and mechanism reaction. The aim of this publication is to identification, understanding, and quantification of "solid solvent" effects when modified natural zeolite is applied as catalyst for cyclisation-acetylation of citronellal in the liquid phase.

Derozane et al. [11] enounced and quantified in the late 1980s the principles of molecular confinement in the intracrystalline volume of zeolites and showed that zeolites can indeed behave like solid solvents, as described and discussed in several recent papers. They provide a new framework for the understanding and use of zeolitic heterogeneous catalysts in organic synthesis, i.e., under conditions where reactions are usually operated at rather low temperature and/or in the liquid phase.

Derozane et al. [12] concluded that a Langmuir–Hinshelwood model describes best the kinetics of reaction catalysed by zeolites in the liquid phase [13-14], e.g., Friedel–Crafts reactions [15], as it accounts for the competition of the reactants and products for the occupancy of the intracrystalline volume of the zeolite, where reaction occurs. This model is preferred to an Eley–Rideal model. Indeed, although only one reactant may be activated at the catalytic sites, the other reactant(s) and product(s) will block its access to the catalytic sites considering the molecular size of the zeolite (micropores, channels, and cavities).

Consider to rate equation that proposed by Derozane et al. [12], the rate of cyclisation-acetylation of citronellal defined as the molar of reaction product per minute and per g of catalyst. The rate equation becomes:

\[
R = \frac{k_{CIT}K_{AA}[[CIT]] [AA]}{(1 + k_{CIT}[[CIT]] + K_{AA}[[AA]] + K_P [P])^2}
\]  

where \(([[CIT]], [AA] and [P])\) are molar of citronellal, acetic anhydride, and product respectively. \(k\) being the kinetic rate constant (inclusive of the number of catalytic sites), and \(k_{CIT}, K_{AA}\), and \(K_P\) representing the adsorption constants of the reactants and product \((P)\). As the intracrystalline volume of the zeolite is most likely to be saturated by the reactants and products for a reaction operated in liquid phase at low temperature, the factor "1" in the denominator can be neglected relative to the other terms, and Eq. (1) becomes

\[
R = \frac{k_{CIT}K_{AA}[[CIT]] [AA]}{(K_{CIT}[[CIT]] + K_{AA}[[AA]] + K_P [P])^2}
\]

which can be converted into a quadratic equation

\[
y = \frac{1}{R} \left( \frac{[[CIT]]}{[AA]} \right) - \frac{k_{CIT}[[CIT]]^2}{kK_{AA}} + \frac{2[[CIT]]}{K_{CIT}} + \frac{k}{k_{CIT}}
\]

or, in a simplified form,

\[
y = ax^2 + bx + c
\]

with

\[
y = \frac{1}{R} \left( \frac{[[CIT]]}{[AA]} \right) - \frac{2a}{b} \text{ and } k = \frac{2}{b} \text{ or } (a/c)^{1/2}
\]

It is worth noting that the values of \(a, b,\) and \(c\) in Eq. (4) are related to each other. For a perfect fit of the data, \(2(a/c)^{1/2} = b\) should be equal to 1. Fitting the experimental data using Eq. (3–5) enables the evaluation of \(k_{CIT}/K_{AA}\) and \(k\). The quality of the fit is determined mostly by the values of the \(a\) and \(b\) coefficients. The coefficient \(c\) corresponds to the ordinate at the origin.

**EXPERIMENTAL SECTION**

**Materials**

Citronella oil (Cymbopogon winterianus-Java type) was obtained from Yogyakarta and the natural zeolite was obtained from Malang-East Java. Zinc chloride,
Table 1. Experimental conditions for the cyclisation-acetylation of (R)-(−)-citronellal by acetic anhydride: effect of the reactant ratio

<table>
<thead>
<tr>
<th>No</th>
<th>Parameter</th>
<th>Exp 1</th>
<th>Exp 2</th>
<th>Exp 3</th>
<th>Exp 4</th>
<th>Exp 5</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>CIT/AA</td>
<td>0.25</td>
<td>0.50</td>
<td>1</td>
<td>1.25</td>
<td>1.50</td>
</tr>
<tr>
<td>2</td>
<td>Citronellal (mmol)</td>
<td>19.20</td>
<td>38.40</td>
<td>38.40</td>
<td>47.52</td>
<td>57.12</td>
</tr>
<tr>
<td>3</td>
<td>Acetic anhydride (mmol)</td>
<td>76.17</td>
<td>76.17</td>
<td>38.08</td>
<td>38.08</td>
<td>38.08</td>
</tr>
<tr>
<td>4</td>
<td>Catalyst Zn²⁺-Natzeo (g)</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
</tbody>
</table>

acetic anhydride, n-hexane, sodium sulphate anhydrous were provided by Merck.

Instrumentation

A set of fractional distillation under reduced pressure was used for isolation of citronellal from citronella oil. A set reflux glass ware with cooler and filled with nitrogen was used as reactor. Composition of citronella oil, distillate, and crude products determined by Gas Chromatography (Hewlett Packard 5890 Series II). A 30 m×0.25 mm i.d.×0.25 μm film thickness Supelco β-DEX 225 capillary column was used for the enantioselective gas chromatography with FID detector. Structure elucidation of citronellal and products conducted by Gas Chromatography–Mass Spectroscopy (Shimadzu QP 5000), FTIR spectrophotometer (Shimadzu FTIR 8201 PC) and ¹H NMR spectrometer (JNM PMX 50 NMR).

Procedure

Isolation of (R)-(−)-Citronellal

(R)-(−)-Citronellal was isolated from citronella oil by fractional distillation under reduced pressure. Enantioselective capillary GC on a Supelco β-DEX 225 column have been used for analysis the enantiomers ratio of citronellal. Structure identification of citronellal was carried out by GC-MS, IR, and ¹H NMR.

Preparation of Zn²⁺-Natzeo

Catalyst Zn²⁺-natural zeolite (Zn²⁺-Natzeo) has been prepared through acid activation of natural zeolite from Malang using HF 1% and HCl 6 M, followed by ion-exchange with 3 M NH₄Cl and calcination at 450 °C for 1 h under nitrogen to obtain H-natural zeolite (H-Natzeo).

Zn²⁺-Natzeo was obtained from H-Natzeo by ion-exchange with 0.1 M ZnCl₂.

Cyclisation-acetylation of citronellal

The activated catalyst (1 g), citronellal (CIT) and acetic anhydride (AA) reactant mixture, with molar ratio in the range CIT/AA 0.25–1.50 (Table 1) were filled in a glass batch reactor. The reaction was conducted at 80 °C with vigorous stirring under nitrogen. As the reaction proceeded, 0.5 mL samples were taken from the system at 10, 20, 30, 60, 120 and 180 min and extracted using n-hexane, analysis was achieved by gas chromatography. Reaction rates are calculated based on the number of moles of IPA-NIPA produced per minute and per g of catalyst.

RESULT AND DISCUSSION

Isolation of (R)-(−)-Citronellal

The maximum concentration of Citronellal (97.3%) was obtained from 250 mL citronella oil (contain about 47% of citronellal) by fractional distillation under reduced pressure at 5 cmHg and 110–120 °C. Asymmetrical GC method has been used to characterize and quantify enantiomers of citronellal. GC analysis using β-DEX 225 showed that isolated citronellal contains 88.21% ee of (R)-(−)-citronellal. FTIR spectrum of isolated citronellal showed that υmax: 1724 cm⁻¹ (s, C=O aldehyde), 2870 and 2715 cm⁻¹ (w, C-H aldehyde), 2924 cm⁻¹ (s, C–H sp³), 1643 cm⁻¹ (w, C=C), 1450 cm⁻¹ (m, –CH₂–), 1381 cm⁻¹ (m, –CH₃), ¹H-NMR (80 MHz, TMS) δ=9.75 ppm (t, 1 H aldehyde proton), δ=5.08 ppm (t, 1 H, C=C olefin proton), δ=2.36 ppm (t, 1 H, H₃), δ=1.88-2.30 ppm (m, 2 H –CH₂– proton bonded to C=C olefin), δ=1.68 ppm (s, 6 H =C(CH₃)₂ isopropylidene proton), δ=1.32 ppm (m, 1H, asymmetric C proton), and δ=1.06 ppm (d, -CH₃ proton). GC-MS m/z = 154, 139, 121, 95, 83, 69, 55, and 41 (base peak).

Cyclisation-acetylation kinetic of (R)-(−)-citronellal with acetic anhydride/Zn²⁺-Natzeo

Table 2 showed the result of cyclisation-acetylation of (R)-(−)-citronellal with acetic anhydride/Zn²⁺-Natzeo. Analysis crude product by GC-MS
showed that fragmentation of IPA and NIPA are similar with m/z = 136 (M⁻-CH₂COOH), 121, 107, 93, 81, 67, 43 (base peak), and 41. Conversions are expressed with respect to the reactant in default, i.e., acetic anhydride. Conversions are thus simply defined as \( X(\%) = \text{moles of IPA-NIPA produced at time } t / \text{moles of acetic anhydride at } t = 0 \)

The increasing of CIT/AA molar ratio showed decreasing yield of IPA-NIPA (Fig. 2). Stereoslectivity to produce IPA not change by variation of CIT/AA.

Decreasing of acetic anhydride showed decreasing of acetyl cation concentration in the reaction system and reaction rate (Fig 3).

Calculation of rate constant and adsorption equilibrium constant ratio of \((R)-(+)-\text{citronellal to acetic anhydride on catalyst surfaces } \text{Zn}^{2+}\text{-Natzeo was followed Langmuir-Hinshelwood equation. The equation stated few assumption that 1) active sites onto catalyst surfaces were uniform and had the same energy level, 2) there was only one reactant molecule which was adsorbed onto active sites, 3) adsorbed reactant molecules will compete to get the active sites and 4) reaction between } (R)-(+)-\text{citronellal with acetic anhydride occurred after both of those adsorbed onto catalyst surfaces [12].}"

Based on reaction rates data which, the quadratic equations were obtained by making graph of \((1/R) ([\text{CIT}] / [\text{AA}]) \text{ vs } [\text{CIT}] / [\text{AA}]\). The quadratic equations in Table 3.

Equations on Table 3 could be used for determination of reaction rate constants \((k_{\text{CIT/AA}})\) and adsorption equilibrium constant ratio, \(K_{\text{CIT/AA}}\), if the requirement was fulfill as \(2(a.c)^{1/2} / b = 1.54\) for kinetic analysis. Reaction rate constant was determined by using \(k_{\text{CIT/AA}} = (2/b)\) or \(k_{\text{CIT/AA}} = (a.c)^{1/2}\). The value of \(k_{\text{CIT/AA}} = (2/b)\) equation was 47.619 mmol (min. g cat)\(^{-1}\), whereas the value of \(k_{\text{CIT/AA}} = (a.c)^{1/2}\) was 30.964 mmol (min. g cat)\(^{-1}\). Adsorption equilibrium constant ratio, \(K_{\text{CIT/AA}} = (2a/b)\) was 7.09. This calculation showed that the reaction rate constant of cyclisation-acetylation \((R)-(+)-\text{citronellal with } \text{Zn}^{2+}\text{-Natzeo/acetic anhydride at reaction time 30 min and 80 °C was 30.964-47.619 mmol (min. g cat)\(^{-1}\).}"

The 7.09 of \(K_{\text{CIT/AA}}\) value showed that adsorption of \((R)-(+)-\text{citronellal to } \text{Zn}^{2+}\text{-Natzeo catalyst was stronger than that of acetic anhydride. The presence of carbonyl group and double bond of C=C (ene group) at}
citronellal molecule was assumed as a controlling factor for tightening the substrate adsorption to catalyst surfaces. The difference of the substrate adsorption abilities was not determined only by reagent polarity, but also by pore sizes and catalyst polarity. The increasing of citronellal adsorption onto catalyst surfaces was inhibited the adsorption of acetic anhydride, even if the excess amount could fill the catalyst pores.

Based on mechanism which was proposed by Deraoune et al. about acetylation of anisole [12], the rate determining step was formation of acetyl-citronellal complexes. According to Freese et al. [16] and Smith et al. [17], an acyl cation produced by reaction of the acetyating agent with the $\text{Zn}^{2+}$-Natzeo was proposed to be the critical species reacting with the adsorbed citronellal as a substrate, implying that reaction of the acyl cation-$\text{Zn}^{2+}$-Natzeo complex with the citronellal as a substrate was the rate-determining step. However, no evidence was provided for the formation of an acyl cation-$\text{Zn}^{2+}$-Natzeo complex, it was demonstrated that the rate constant for acylation was more dependent on the nature of the substrate rather than on the nature of the acetylating agent. Zeolite catalyzed reaction into liquid phase was not dependent only by reactant ratio, but also pores size and zeolite polarity.

The proposed mechanism of $(R)$-$(+)$-citronellal cyclisation-acetylation with acetic anhydride/$\text{Zn}^{2+}$-Natzeo describes in Fig. 4.

CONCLUSION

The major products of one pot cyclisation-acetylation of $(R)$-$(+)$-citronellal with $\text{Zn}^{2+}$-Natzeo/acetic anhydride was IPA and NIPA. According to Langmuir-Hinshelwood mechanism, kinetic analysis of cyclisation-acetylation of $(R)$-$(+)$-citronellal with $\text{Zn}^{2+}$-Natzeo/acetic anhydride showed that the rate constant $k = 47.619 \text{mmol (min.g cat)}^{-1}$, the value of $k_{\text{OT/AA}} = 30.964 \text{mmol (min. g catalyst)}^{-1}$, and the adsorption equilibrium constant ratio $K_{\text{OT/AA}} = 7.09$. Adsorption ability of citronellal was much stronger than that of acetic anhydride onto catalyst surfaces, the greater amount of citronellal adsorbed onto catalyst was decreased the reaction rate of citronellal cyclisation-acetylation.

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REFERENCES