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# Substrate specificity in decarboxylation of mixtures of acetate and propionate using oxidized Pt electrodes and galvanic square-wave pulsed electrolysis†

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The decarboxylation (of mixtures) of short-chain carboxylic acids (C<sub>2</sub> and C<sub>3</sub>) on oxidized platinum anodes was investigated using constant current and galvanic square-wave pulse electrolysis. At constant current, a high ethylene to ethane product ratio indicates that propionate is the substrate of preferential decarboxylation in propionate/acetate mixtures, depending on the feed ratio. The specificity of (oxidized) Pt electrodes towards C<sub>3</sub> decarboxylation can be further enhanced by the application of cathodic and anodic pulses. The application of relatively long cathodic pulses and very short anodic pulses has been demonstrated to facilitate the formation of high ethylene to ethane ratio product mixtures, which are higher than those obtained under constant current conditions. In particular, extended cathodic pulses have been observed to enhance the faradaic efficiency towards oxygen and to reduce carboxylate conversion. Based on isotherm and RRDE data, we propose that the selectivity for propionate is attributable to a higher affinity for the oxidized Pt electrode, which is further enhanced by cathodic and anodic pulses. The use of galvanic square wave-pulse electrolysis thus offers a promising pathway for the efficient conversion of bio-derived acids into fuels and chemicals.

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## Introduction

Replacement of fossil resources by biomass for the synthesis of fuels and chemicals can significantly contribute to reaching the climate targets set by the European Commission by 2050.<sup>1</sup> Industrial efforts to optimize the use of biomass-derived carbon are rapidly maturing, resulting in an increased availability of bio-based liquids.<sup>2,3</sup> Although research and development have led to advancement in biomass liquefaction technologies,<sup>4</sup> the quality of these liquids is still fairly poor and requires upgrading before processing in a refinery is feasible. As large scale traditional upgrading of bio-based liquids is energy intensive, alternatively electrochemical processes are developed to improve bioliquid quality and hence establish a circular and bio-based economy.<sup>5</sup> In particular Kolbe electrolysis using Pt electrodes receives increased attention to convert bio-derived

acids and to upgrade pyrolysis oil and make it suitable for the production of fuels and chemicals.<sup>6</sup>

The effect of reaction conditions on performance of (Pt) electrodes has been frequently studied and a broad substrate scope has been evaluated. Nevertheless, a vast majority of studies examines electrolyte solutions containing single (complex) carboxylic acids in often non-aqueous solvents such as acetonitrile, dimethylformamide or methanol, containing supporting electrolytes like sodium sulphate to increase electrical conductivity.<sup>7–11</sup> Utilization of bio-based liquids – such as the water-soluble fraction of pyrolysis oil – in refineries likely requires treatment of concentrated aqueous solutions of mixtures with a high content of short-chain carboxylic acids, with acetic acid (C<sub>2</sub> acid) and propionic acid (C<sub>3</sub> acid) representing up to 30% of the acid content.<sup>12,13</sup>

Recent work from Angulo *et al.* studied the effect of electrolyte composition and operating current density on the product distribution during the electrochemical oxidation of propionic acid in aqueous media using Pt electrodes.<sup>14</sup> They showed that high faradaic efficiency (>50%) towards ethylene was obtained, while butane (the Kolbe product) was not detected being in agreement with earlier work of Levy and Sanderson investigating the electrochemical oxidation of (mixtures of) short-chain *n*-alkanoic acids (C<sub>3</sub>–C<sub>6</sub>) in aqueous media.<sup>15,16</sup> Interestingly, the earlier work of Levy and Sanderson indicated

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that the decarboxylation of  $C_3$  acid results in butane formation exclusively in the presence of hexanoic acid ( $C_6$  acid).<sup>15,16</sup> Klocke *et al.*,<sup>17</sup> studied the influence of carboxylic acid chain length during anodic decarboxylation of 3-oxanonanoic acid and 3-oxapentadecanoic acid in methanol confirming favourable formation of (cross-) coupling products in the presence of a carboxylic acid with a longer chain length. They attributed this phenomenon to enhanced interaction of longer chain acids with the electrode surface. A recent work by Neubert *et al.*<sup>18</sup> investigated the (hetero-) coupling (of mixtures) of  $C_4$ ,  $C_6$  and  $C_8$  acids for bio-fuel production purposes. Overall, the results of the extensive analysis of product distributions and acid conversion suggest a high substrate specificity for  $C_6$  acid when non-equimolar mixtures of  $C_4$ ,  $C_6$  and  $C_8$  are present in the feed (approximately 60% of the total conversion is assigned to  $C_6$  acid decarboxylation). These three studies mainly discuss the product selectivity in mixtures of long chain carboxylic acids, while decarboxylation of mixtures of small molecules ( $C_2$  or  $C_3$ ) was not examined in detail. Moreover, constant-current or constant potential conditions were used to evaluate acid decarboxylation.

Recently pulsed electrolysis emerged as an interesting alternative to improve reactivity and selectivity for many synthetic organic electrochemical reactions.<sup>19–24</sup> In fact, pulsed electrolysis was already used by Hickling and Wilkins to investigate dimer formation during anodic decarboxylation of  $C_2$  acid.<sup>25</sup> Negative current pulses were found to interfere with dimer production, and instead permit oxygen evolution to the expense of an overall lower reactant conversion.<sup>25</sup> Hioki *et al.*<sup>26</sup> used rapid alternating polarity instead of classical direct current measurements highlighting that both product selectivity and reactant conversion for a wide range of carboxylic acids, including biomass-derived acids, are positively affected. The favourable performance was explained by a lower local acidification at the electrode surface resulting in a higher concentration of deprotonated acids (required for the decarboxylation process). Clearly pulsed electrolysis has recently been revealed to provide means to enable electrolysis with unprecedented conversion and selectivity.

In this study, we discuss decarboxylation of mixtures of short-chain carboxylic acids ( $C_2$  and  $C_3$ ) using constant current and galvanic square-wave pulse electrolysis. It is shown that  $C_3$  is the preferred substrate for conversion using (oxidized) Pt electrodes, and that using galvanic square-wave pulsed electrolysis is a means to enhance this substrate specificity of  $C_3$  over  $C_2$ . We will explain the observations by considering adsorption isotherms of (deprotonated) acids on Pt (oxide) surfaces and rotating-ring disk measurements. Methods to enhance the efficient and selective conversion of bio-derived acids *via* mixed Kolbe electrolysis are discussed in relation to its economic viability for the production of fuels and chemicals.

## Experimental

### Materials and chemicals

Platinum foil (0.025 mm thick, 99.9% pure, Alfa Aesar), platinized titanium mesh (Magneto Special Anodes B.V.), and a Pt

ring-disk electrode (Metrohm) were used. Nitric acid (ACS reagent, 70%), acetic acid (glacial, ReagentPlus®, >99%), sodium acetate (ACS reagent, >99.0%), propionic acid (ACS reagent, ≥99.5%), sodium propionate (≥99.0%), hexanoic acid (≥99%) and sodium hydroxide (reagent grade, >97%) were used as purchased from Sigma Aldrich without purification. Ethanol (technical grade, BOOM BV) was used for cleaning purposes. Ultrapure water (18.2 MΩ cm<sup>-1</sup>, from a Millipore, Milli-Q Advantage A10) was used as solvent.

### Electrochemical measurements and product analysis

All measurements were performed in a single compartment three electrode glass cell (100 mL) equipped with a platinum foil working electrode (0.8 cm<sup>2</sup> geometric area), a platinized titanium mesh counter electrode (6 cm<sup>2</sup> geometric area) and a Ag/AgCl (3 M NaCl, ProSense) reference electrode connected to a Biologic VMP3 Potentiostat. A custom-made Teflon electrode holder was used to cover the backside of the working electrode and to establish a defined exposed electrode area. Two needle ports used as gas inlet and outlet, respectively enabled continuous purging of the glass cell and were used for the detection of gaseous products by gas chromatography. Prior to each experiment the glass cell, working-, and counter electrode were cleaned with 10% nitric acid and Milli-Q water to remove organic residues. The electrolyte in mixed Kolbe electrolysis experiments always consisted of a total acid concentration of 1 M. The pH of the electrolyte was adjusted to pH 5. The acid ratio of acetic and propionic acid was varied by adjusting the concentrations of the acids. The freshly prepared electrolyte (70 mL) was deoxygenated using 30 mL min<sup>-1</sup> Helium (>5.0). Electrochemical cleaning of the Pt working electrode was performed using cyclic voltammetry at a scan rate of 200 mV s<sup>-1</sup> after which cyclic voltammograms at 100 and 50 mV s<sup>-1</sup> were recorded. Unless mentioned otherwise, potentials were not corrected for ohmic drop, preventing artefacts caused by overcompensation. If not stated otherwise, results are presented against the RHE scale, calculated according to the following equation:

$$E_{\text{RHE}} = E_{\text{Ag/AgCl}} + 0.059 \text{ pH} + E_{\text{Ag/AgCl}}^0 \quad (1)$$

where  $E_{\text{Ag/AgCl}}$  is the measured potential and  $E_{\text{Ag/AgCl}}^0$  is the Ag/AgCl (3 M NaCl) reference electrode potential (+0.210 V). The stability of the RE was frequently determined using a calibrated master reference electrode (Ag/AgCl; 3 M NaCl, BASi) and a voltmeter. All experiments were carried out at room temperature.

Faradaic efficiencies were determined using galvanostatic conditions at a current density of 250 mA cm<sup>-2</sup> to ensure Kolbe electrolysis occurs.<sup>27</sup> Throughout the measurements the solution was continuously stirred by a magnetic stirring bar at a stirring rate of 900 rpm. Liquid analysis to determine the amount of liquid products and the  $C_2/C_3$  conversion was performed after a constant charge of 5000 C had passed.

Pulsed electrolysis experiments using fast galvanic square-wave pulses were performed in the three-electrode electrochemical cell described above using a VersaSTAT 3 potentiostat (Princeton Applied Research). Anodic pulses were performed



similarly to the constant current experiments at  $250 \text{ mA cm}^{-2}$ , while negative galvanic pulses were performed at  $-1 \text{ mA cm}^{-2}$ . Experiments were performed with an anodic pulse length of 10 or 200 ms, while the duration of the cathodic pulse was varied from 50 to 500 ms. Galvanic square-wave pulses were performed for a total duration of 2 hours irrespectively of the anodic and cathodic pulse duration.

Product analysis was performed by gas chromatography (GC, Interscience CompactGC, the Netherlands) and high-performance liquid chromatography (HPLC, Agilent technology 1200 series, Agilent). Online detection of gaseous products by GC was performed at 10 minutes intervals with a He ( $>5.0$ ) purge at a constant flow rate of  $30 \text{ mL min}^{-1}$ . Light gases ( $\text{H}_2$ ,  $\text{O}_2$ ,  $\text{CO}_2$ ) were detected with a ShinCarbon micropacked column (ST 80/100 2 m, 0.53 mm at  $90 \text{ }^\circ\text{C}$ ) connected to a thermal conductivity detector (TCD) operating at  $110 \text{ }^\circ\text{C}$ . Hydrocarbons (C1–C4) were detected using a Rt Q BOND PLOT (0.32 mm ID,  $10 \mu\text{m}$ , 15 m, at  $60 \text{ }^\circ\text{C}$ ) column connected to a flame ionization detector (FID) operating at  $150 \text{ }^\circ\text{C}$ . Liquid products were measured by injecting liquid aliquots of the electrolyte into a HPLC equipped with a refractive index detector. Acetic acid, propionic acid and ethanol were detected on a Hi-Plex H column which was heated to  $65 \text{ }^\circ\text{C}$ . The mobile phase consists of  $5 \text{ mM H}_2\text{SO}_4$  (flow rate of  $0.6 \text{ mL min}^{-1}$ ).  $\text{H}_2\text{SO}_4$  converted the acetate or propionate present in the sample to acetic or propionic acid, which was also detected as such.

Temkin isotherms were constructed from the underpotential deposition of hydrogen ( $H_{\text{upd}}$ ) determined by cyclic voltammetry measurements performed at a scan rate of  $100 \text{ mV s}^{-1}$  unless mentioned otherwise. As supporting electrolyte, a  $100 \text{ mM}$  acetic acid/sodium acetate buffer with a pH of 5 was used and different concentrations of propionic acid (or hexanoic acid for comparison) were added to the electrolyte, while maintaining a pH of 5. The charge (the integral of current over time) related to  $H_{\text{upd}}$  was determined from both adsorption and desorption after subtracting the capacitance current. The plotted values represent the charge associated with  $H_{\text{upd}}$  desorption, which generally matches the  $H_{\text{upd}}$  charge from adsorption. The adsorption isotherms were determined by fitting the coverages at various concentrations in OriginPro using the Temkin adsorption model, as this model resulted in the best fit.<sup>28</sup>

Rotating ring disc electrode (RRDE) experiments were performed using a Metrohm Autolab RRDE with a Pt disk used as working electrode and a Pt ring electrode for *in situ* detection of products. The counter electrode (CE) was a  $2 \text{ cm}^2$  Pt flag electrode. The collection efficiency of the Pt ring electrodes was determined to be 24%.

## Results and discussion

### Chronopotentiometry for faradaic efficiency analysis

Product selectivity and substrate specificity in (mixtures of)  $\text{C}_2$  and  $\text{C}_3$  acid were determined using chronopotentiometry, online-GC measurements to detect gaseous products, and analysis of liquid products at the end of the experiments using HPLC. Using electrolyte mixtures with different  $\text{C}_2$  to  $\text{C}_3$  acid

ratios, the faradaic efficiency (%) towards the different reaction products and the conversion (mmoles) of  $\text{C}_2$  and  $\text{C}_3$  acid were determined during electrolysis at  $250 \text{ mA cm}^{-2}$  on a Pt electrode after a charge of 5000 C passed the cell. Electrolysis of pure  $\text{C}_2$  acid electrolyte exclusively yields carbon dioxide and ethane (the Kolbe product), in agreement with earlier work.<sup>27,29,30</sup> Carbon dioxide is not included in the product distribution in Fig. 1 as it is considered a byproduct resulting from the decarboxylation of carboxylic acid. The electron transfer that occurs prior to decarboxylation is already accounted for in the calculation of the faradaic efficiency of the final products (Kolbe, Hofer-Moest, non-Kolbe, disproportionation, and ester products, see Fig. S1† in ESI for the reaction mechanism). A conversion of approximately 47 mmol of  $\text{C}_2$  acid was determined which results in a selectivity of 92% for  $\text{C}_2$  acid decarboxylation (the maximum possible conversion considering the charged passed was 51 mmol, calculated by dividing the transferred charge by the Faraday constant and number of electrons required for acid decarboxylation). This is also in agreement with previous literature on Kolbe electrolysis of  $\text{C}_2$  acid.<sup>27,29–31</sup>

Electrolysis of the  $\text{C}_3$  acid resulted in ethylene and ethanol formation with a  $\text{FE}_{\text{ethylene}}$  of  $\sim 60\%$  and a  $\text{FE}_{\text{ethanol}}$   $\sim 27\%$ , respectively. These values are in excellent agreement with data of earlier work on the electrochemical decarboxylation of  $\text{C}_3$  acid, which reported product yields of 61% and 31% towards ethylene and ethanol, respectively, in a  $2 \text{ M}$   $\text{C}_3$  acid electrolyte with similar pH.<sup>15</sup> We also identified formation of ethane, butane and ethyl propionate, in minor quantities. In total, the sum of the individual FE remains short of 100%, which is likely explained by alternative oxidation processes such as the oxidation of ethanol to carbon dioxide, the formation of hydrogen peroxide *via* water oxidation, or Pt dissolution during electrolysis.<sup>32</sup> Interestingly, the conversion of only 28 mmol of  $\text{C}_3$  in

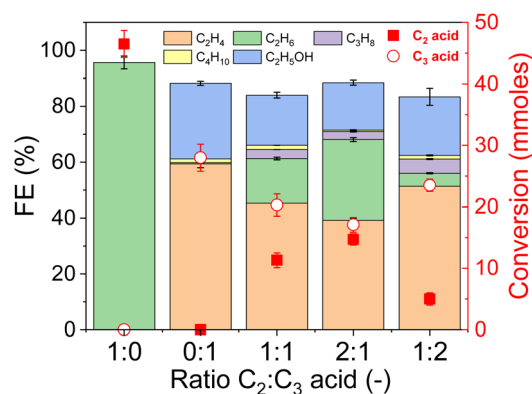


Fig. 1 Kolbe electrolysis of mixtures of acetic acid ( $\text{C}_2$ ) and propionic acid ( $\text{C}_3$ ) with different ratios (left y-axis). Faradaic efficiency (%) to ethylene, ethane, propane, butane and ethanol (right y-axis). Conversion (in mmol) of  $\text{C}_2$  acid (circle) and  $\text{C}_3$  acid (square). Conversion was determined after a charge of 5000 C passed through the cell. The electrolyte pH was approximately the  $\text{pK}_a$  of both acids ( $\text{pH} \sim 5$ ), the total acid concentration was  $1 \text{ M}$  and measurements were performed with an applied current density of  $250 \text{ mA cm}^{-2}$ . The error margins are based on  $N = 2$ .



comparison to 47 mmol of  $C_2$  acid, suggests significant differences in reaction mechanism of decarboxylation of  $C_3$  acids compared to  $C_2$  acid.  $C_3$  acids yield primarily products generated *via* a two-electron pathway, whereas for  $C_2$  acid oxidation, product formation usually requires only one electron transfer step.

Electrolysis of mixtures of  $C_2$  and  $C_3$  acids was performed either using equimolar amounts of the acids or excess of either  $C_2$  or  $C_3$  in a 1 : 2 ratio. Electrolysis of a 1 : 1 mixture of  $C_2$  and  $C_3$  acid revealed a clear substrate specificity towards conversion of the  $C_3$  acid with the main products being ethylene and ethanol (with a summed FE of  $\sim 64\%$ ), while ethane formation only accounted for a FE of 16%. Additionally, the conversion of  $C_3$  acid of 21 mmol closely resembled the conversion of a pure  $C_3$  acid electrolyte. Decarboxylation of the  $C_2$  acid amounted to 11 mmol, in agreement with the lower electron demand for the Kolbe dimer formation, ethane. Even for electrolytes containing an excess of  $C_2$  acid (ratio  $C_2 : C_3 = 2 : 1$ ),  $C_3$  acid was still favourably oxidized. This is evident from a high contribution of ethylene and ethanol (total FE: 56%) in the product composition and a slightly higher conversion of  $C_3$  compared to  $C_2$  acid (17 mmol *vs.* 15 mmol). The preferred conversion of  $C_3$  acid is likely associated with preferred (stronger) adsorption characteristics.

To verify this hypothesis, mixtures of  $C_3$  and  $C_6$  acid (having an even higher adsorption strength than  $C_3$  acid) were electrolysed (Fig. S2†). Contrasting electrolyte mixtures of  $C_2$  and  $C_3$  with a ratio of 2 : 1, the presence of  $C_6$  acid significantly decreased the conversion of  $C_3$  acid by a factor of four from 28 mmol to 7 mmol, most likely at the expense of  $C_6$  acid conversion as oxygen (formed *via* competitive water oxidation) was not detected. Even in a 6 : 1 excess of  $C_3$  acid, the conversion of  $C_3$  acid appeared to be significantly reduced (18 mmol) in comparison to electrolytes containing only  $C_3$  acid. Furthermore in the presence of  $C_6$  acid, the product distribution of  $C_3$  acid oxidation products was altered and a significant contribution of butane of 29% (2 : 1) and 13% (6 : 1) was observed in agreement with the work of Levy and Sanderson.<sup>15</sup> Though a quantitative analysis of the different (cross coupled) oxidation products of  $C_6$  acid was not feasible,<sup>15,33</sup> these results support the conclusion that substrate specificity of oxidized Pt electrodes is determined by differences in surface affinity of adsorbates.

### Pulsed electrolysis

Galvanic square-wave pulsed electrolysis was performed in a 1 : 1 mixture of acetic acid and propionic acid using positive current pulses ( $J_a = 250 \text{ mA cm}^{-2}$ ) with a fixed duration ( $t_a$  of 10 or 200 ms), while negative current pulses ( $J_c = -1 \text{ mA cm}^{-2}$ ) with varying durations ( $t_c = 50\text{--}500$ ) (see Fig. S3† in the ESI† for a schematic representation) were used. In the following primarily the ratio of the major products  $C_2H_4$  and  $C_2H_6$  (the major products of Kolbe oxidation of  $C_3$  and  $C_2$  acid respectively) is considered to analyse the impact of the perturbation profile (see Fig. 2).

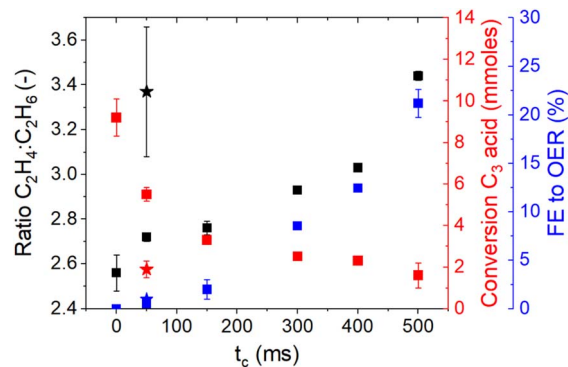


Fig. 2 Kolbe electrolysis performed with a 1 : 1 mixture of  $C_2$  and  $C_3$  acid (1 M, pH 5) using pulsed electrolysis. The ratio of ethylene and ethane, the  $C_3$  acid conversion and the faradaic efficiency towards oxygen were determined during alternating anodic (pulse length of 10 ms and 200 ms at  $250 \text{ mA cm}^{-2}$ ) and cathodic (at  $-1 \text{ mA cm}^{-2}$ ; pulse length indicated by the x-axis). Experiments performed at an anodic pulse length of 10 ms are shown using star symbols and anodic pulse was fixed with a duration of 200 ms are shown by squares.

Under constant current conditions ( $t_c = 0$ ), a  $C_2H_4 : C_2H_6$  ratio of 2.5 was obtained (compare also Fig. 1). Using alternating pulses (using  $t_a = 200$  ms) the ratio is clearly increasing, with the duration of the cathodic pulse. At the longest cathodic pulse duration  $t_c = 500$  ms, a maximum ratio of  $C_2H_4 : C_2H_6$  of  $\sim 3.5$  is obtained. The favourable formation of ethylene, *i.e.* the product of  $C_3$  acid decarboxylation is accompanied by a significant decrease in the total conversion of  $C_3$  acid and a significant increase in oxygen evolution as revealed by a  $FE_{\text{oxygen}}$  of approximately 22% at the longest cathodic pulse durations. It is noteworthy that the use of shorter anodic pulses ( $t_a = 10$  ms at  $J_c = -1 \text{ mA cm}^{-2}$ , as indicated by star symbols in Fig. 2) enables the decarboxylation process to be conducted at a favourable  $C_2H_4 : C_2H_6$  ratio. The ratio of the products was approximately 3.4, while the oxygen evolution reaction (OER) was suppressed. However, variations in the cathodic current density ( $J_c = -5 \text{ mA cm}^{-2}$  or  $J_c = 0.1 \text{ mA cm}^{-2}$ ) did not result in any favourable product composition.

### Voltammetry and adsorption isotherms

Rotating-ring disk measurements (Pt disc and Pt ring electrode) were used to further evaluate the electrochemical behaviour of  $C_2$  and  $C_3$  carboxylic acids (Fig. 3). While linear sweep voltammetry (LSV) was performed at the Pt disc (solid lines,  $J_{\text{disc}}$ ), chronoamperometry was performed at the Pt ring at 0.1 V *vs.* RHE allowing for oxygen detection by the oxygen reduction reaction (dashed lines,  $J_{\text{ring}}$ ).

Independent of the carboxylic acid used, below an applied disk potential of 1.7 V *vs.* RHE, no significant disc or ring current was detected and only from  $\sim 1.7$  V *vs.* RHE an anodic current was observed at the disk independent of the acid used.

Considering that  $J_{\text{ring}}$  is simultaneously increasing, oxygen evolution occurs at the disk.<sup>27,30</sup> In acetic acid (black traces in Fig. 3),  $J_{\text{disc}}$  reaches a plateau in current density of  $17 \text{ mA cm}^{-2}$  at 2.4–2.5 V *vs.* RHE, which is commonly known as the inflection



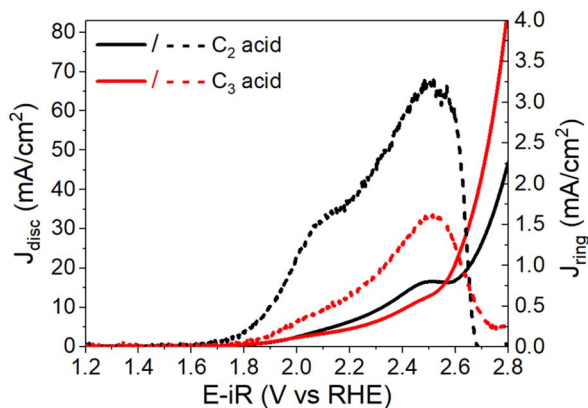


Fig. 3 Rotating-ring disk measurements (3000 rpm) of 0.5 M  $C_2$  (black curves) acid and 0.5 M  $C_3$  (red curve) acid (both at pH 5) using a Pt disk (solid lines) and Pt ring (dashed lines) electrode. Linear sweep voltammetry ( $10 \text{ mV s}^{-1}$ ) was performed at the disk while the ring potential was  $E_{\text{ring}} = +0.1 \text{ V}$  vs. RHE. The ring currents have been corrected for the collection efficiency and inverted to allow for comparison to the  $J_{\text{disc}}$ .

zone, and according to literature coincides with a transition between OER at lower potentials to hydrocarbon production (Kolbe reaction) at higher potentials.<sup>30</sup> Although less pronounced, an inflection zone is also visible at approximately 2.5 V when using  $C_3$  acids as electrolyte. In line with  $J_{\text{disc}}$  also the  $J_{\text{ring}}$  increases from  $\sim 1.7 \text{ V}$  to more positive potentials, confirming the formation of oxygen. At potentials above the inflection zone,  $J_{\text{ring}}$  drops to (almost) zero, indicating that oxygen evolution was (largely) suppressed at these high anodic potentials. In addition to the slight shift in applied Pt disk potential, a clear difference in  $J_{\text{ring}}$  between  $C_2$  (black traces in Fig. 3) and  $C_3$  (red traces in Fig. 3) acid was observed. Despite similar disk currents obtained in the plateau region, a maximum  $J_{\text{ring},C_2}$  of  $\sim 3.2 \text{ mA cm}^{-2}$  at 2.4 V was obtained for  $C_2$  acids, while for  $J_{\text{ring},C_3}$  only a maximum current density of  $\sim 1.6 \text{ mA cm}^{-2}$  was obtained. The difference in  $J_{\text{ring}}$  is assigned to suppression of the OER with increasing carbon chain length of the carboxylic acid used.

The interaction and adsorption strength of the different acids with the Pt surface were obtained from measurements of the underpotential deposition of hydrogen ( $H_{\text{upd}}$ ) in the presence of different concentrations of the acids using cyclic voltammetry (CV)<sup>28</sup> (Fig. S4†) and adsorption isotherms constructed (Fig. S5†) thereof. A buffer of  $C_2$  acid was used to exclude the introduction of foreign anions and to maintain a stable concentration of deprotonated carboxylic acid. Examples of the cyclic voltammetry (CV) measurements performed with a Pt electrode in a  $C_2$  acid buffer (100 mM) with 50 mM of  $C_3$  or  $C_6$  acid are shown in Fig. S4.†

In the  $C_2$  acid buffer, hydrogen adsorption and desorption on the different facets of Pt is observed at  $\sim 0.25$  and  $\sim 0.1 \text{ V}$  and  $\sim 0.05$  and  $0.2 \text{ V}$  when sweeping towards negative and positive potentials, respectively.<sup>34,35</sup> The current associated with hydrogen underpotential deposition ( $H_{\text{upd}}$ ) is proportional to the amount of hydrogen ad- or desorbed from the Pt surface and

can therefore be used to reveal the fraction of H sites that are occupied by adsorbed (foreign) molecules.<sup>28</sup> Thus, the decrease in  $H_{\text{upd}}$  peak current density upon addition of  $C_3$  acid to the  $C_2$  acid buffer (red trace in Fig. S4a†), is directly related to the adsorption of  $C_3$  acid with the Pt surface and agrees well with reports on adsorption of other organic molecules such as phenol in acidic media.<sup>28,36</sup> Integrating the  $H_{\text{upd}}$  current obtained for different concentrations of  $C_3$  allows to determine the total charge, *i.e.*, the concentration of adsorbed hydrogen. Comparison to the  $H_{\text{upd}}$  charge determined for the Pt surface in the  $C_2$  acid buffer only allows to estimate the fraction of  $H_{\text{upd}}$  that was inhibited by the presence of the  $C_3$  acid. The fraction of  $H_{\text{upd}}$  inhibited *vs.* the logarithmic concentration of  $C_3$ , reflecting the Temkin isotherm, results in a linear correlation (Fig. S5b†) that allows to estimate the heat of adsorption or Temkin constant of the  $C_3$  carboxylic acid from the slope of a linear fit to be  $\alpha = 0.365$ .<sup>37</sup>

The Temkin constant for the adsorption of hexanoic acid was determined for comparison. As shown in Fig. S5b,† a higher  $\alpha$  ( $\alpha = 0.403$ ) for  $C_6$  acid, and hence a higher heat of adsorption indicates a stronger interaction of  $C_6$  acid with the platinum electrode surface compared to  $C_3$  acid. The adsorption data of  $C_2$  acid on a Pt electrode surface suggest an  $\alpha$  value of only 0.08.<sup>38</sup> Overall the RRDE measurements and adsorption isotherms support the hypothesis of a stronger adsorption with increasing carboxylic acid chain length and thus the product distribution of Kolbe electrolysis of mixtures of carboxylic acids will likely depend on the adsorption strength of the individual components and their concentrations.

## Discussion

To explain the effect of a longer cathodic duration in square-wave pulse experiments (Fig. 2), we assume the following: during a negative current pulse, the established carboxylate layer will likely desorb and the double layer structure will thus rearrange as a function of the cathodic pulse duration ( $t_c$ ) and hence on the amount of passed cathodic charge ( $Q_c$ ). With increasing  $Q_c$ , adsorbed carboxylate anions will desorb from the Pt surface, yet it is reasonable to assume that desorption of  $C_2$  occurs preferentially due to the significantly lower adsorption strength, as demonstrated by the Temkin isotherms. During the positive pulse, the stronger adsorption of  $C_3$  acid will lead to a successive change in the ratio of adsorbed  $C_3$  to  $C_2$  anions at the Pt surface and thus to a preferred conversion of  $C_3$  acid compared to  $C_2$  acid, consequently increasing the share of  $C_2H_4$  in the product distribution.

At a high  $Q_c$ , the anodic pulse charge ( $Q_a$ ) is likely not sufficient to fully restore Kolbe conditions. As a result, oxygen evolution is no longer suppressed by the formation of a (dense) carboxylate layer resulting in the exponential increase in  $O_2$  formation at  $t_c > 150 \text{ ms}$  (see Fig. 3). Despite variations in electrolyte composition and pulse conditions, an exponential increase in oxygen evolution was also observed in the work of Hickling *et al.* when cathodic pulses were applied.<sup>25</sup> They investigated the effect of cathodic pulses using aqueous acetic acid solutions and found that when  $Q_a$  was roughly 75 times



higher than  $Q_c$ , oxygen was formed with a CE of  $\sim 20\%$ , which closely resembles the OER FE obtained here. It is important to note that a shift in product selectivity and substrate specificity as a result of the galvanic pulses as well as the implications of changes in pulse duration were not revealed previously.<sup>25</sup>

Based on the presented results it is expected that by applying galvanic square-wave pulses in mixtures with other carboxylic acids, substrate specificity will also shift to the stronger adsorbing acid. Thus, utilization of galvanic square-wave pulses to influence product selectivity and substrate selectivity in mixed Kolbe electrolysis poses opportunities in the field of electrosynthesis using bio-based liquids and potentially allows for the formation of high-value products. For example, selective production of ethylene from propionic acid can contribute to the production of bio-based fuels or plastics. Furthermore, besides manipulation of acid specificity of the oxidized Pt surface, using galvanic square-wave pulses might also prevent undesired oxidation of other compounds naturally present in bio-based liquids, *e.g.* alcohols. Nevertheless, further research is necessary to gain a deeper understanding of these potential opportunities and likely additional means to suppress the undesired occurrence of the OER might be essential. Examples of additives which have been used previously to suppress the OER, include sodium dodecyl sulphate, which forms a hydrophobic layer on the Pt anode and improves selectivity to conversion of hydrocarbons.<sup>39</sup> Alternatively, the platinum anode could be modified by a PTFE-layer to increase the hydrophobic nature of the electrode to similarly suppress OER.<sup>40</sup>

Despite the opportunities that pulsed electrolysis in mixed Kolbe electrolysis can offer, it is important that the results obtained in this work are approached with caution. Firstly, the results do not account for the differences in diffusion constants of the carboxylic acids. Changes in the thickness of the diffusion layer can lead to variations in acid conversion and product selectivity. To provide more insight in this regard, measurements using forced transport such as rotating disk electrode (RDE) measurements or measurements in flow cell configurations could be employed. Secondly, the precise composition of the adsorbed layer on the platinum surface is still not known and detailed understanding of the structure of the adsorbed layer during pulsed electrolysis is required. *In situ* techniques such as surface-enhanced infrared spectroscopy (SEIRAS) or surface-enhanced Raman spectroscopy (SERS) could reveal the composition of the adsorbed layer in (mixed) Kolbe electrolysis. Nevertheless, this work clearly emphasizes the potential of pulsed electrolysis as a strategic tool to modulate product distributions and substrate specificities beyond mixed Kolbe electrolysis of  $C_2$  and  $C_3$  acid, *i.e.*, for electrolysis of more complex technical feeds.

## Conclusions

Kolbe electrolysis of  $C_2$  and  $C_3$  acid as well as mixtures thereof was investigated using Pt electrodes. The reactant specificity and product selectivity were analyzed and it is highlighted that  $C_3$  acid is preferably oxidized over  $C_2$  acid. Conversion of  $C_3$  was further improved using a galvanic square-wave pulse

electrolysis, but at the expense of lower conversion and formation of oxygen at extended cathodic pulse durations which was efficiently suppressed by modulation of the anodic pulse duration. The preferential conversion of  $C_3$  acids is explained by adsorption properties of the acids on the (oxidized) Pt surfaces, confirmed by Temkin adsorption isotherms measurements of  $C_3$  and  $C_6$  acid. Galvanic square-wave electrolysis is a means to manipulate the substrate specificity towards the acid with the strongest affinity for the electrode surface and can be used to advance the production of fuels and chemicals from bio-based resources.

## Data availability

Data for this article, including gas chromatography, cyclic voltammetry and chronopotentiometry data will be made available at Zenodo at <https://doi.org/10.5281/zenodo.14515195>.

## Author contributions

M. Olde Nordkamp: conceptualization, data curation, formal analysis, investigation, methodology, visualization, writing – original draft. T. Ashraf: conceptualization, data curation, formal analysis, methodology. G. Mul: conceptualization, funding acquisition, resources, supervision, writing – review & editing. B. Mei: formal analysis, methodology, conceptualization, supervision, writing – review & editing.

## Conflicts of interest

There are no conflicts to declare.

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