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Pressurized CO₂ as a carboxylating agent for the biocatalytic *ortho*-carboxylation of resorcinol†

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The utilization of gaseous carbon dioxide instead of bicarbonate would greatly facilitate process development for enzyme catalyzed carboxylations on a large scale. As a proof-of-concept, 1,3-dihydroxybenzene (resorcinol) was carboxylated in the *ortho*-position using pressurized CO₂ (~30–40 bar) catalyzed by *ortho*-benzoic acid decarboxylases with up to 68% conversion. Optimization studies revealed tight pH-control and enzyme stability as the most important determinants.

Introduction

Great efforts are currently undertaken to utilize the inexpensive, non-toxic and abundantly available waste gas CO₂ as a C₁ carbon source for the syntheses of valuable chemicals, materials or fuels.¹ However, despite the fact that photosynthetic CO₂-fixation mediated by RuBisCO is one of the most dominant reactions in nature, which binds ~10¹¹ tons CO₂ p.a.,² the chemical activation of CO₂ remains challenging due to the high energy input required for substrate activation.¹ Recently, the catalytic carboxylation of epoxides using salen complexes, zinc salts and double metal cyanide catalysts opened access to poly(ether)-carbonates for the production of polyurethanes³ and is at the threshold of industrial implementation. Further carboxylation strategies using (transition) metal⁴ or organocatalysts⁵ have been developed to broaden

the usability of carbon dioxide, but these methods are still in their infancy regarding commercialization.

The harsh reaction conditions (~90 bar, 120–300 °C), varying *ortho/para*-selectivity and incomplete yields are major issues in the large-scale production of salicylic acids *via* the chemical carboxylation of phenolates using pressurized CO₂ gas (Kolbe–Schmitt reaction).⁶ Although improved by microwave-heating using a bicarbonate-based ionic liquid, the process still suffers from moderate selectivity and yields.⁷

Biocatalytic methods have been explored as alternatives for the carboxylation of electron-rich (hetero)aromatic compounds to yield the corresponding carboxylic acids.⁸ Mild reaction conditions, exquisite regioselectivity and excellent yields (*e.g.* 95% for the bio-carboxylation of resveratrol)⁹ emphasize the power of bio-carboxylation processes.

However, in the majority of biocatalytic carboxylation protocols reported so far, bicarbonate is used as a CO₂ source, which needs to be applied at elevated concentrations (~3 M) to shift the equilibrium towards the thermodynamically unfavored carboxylation.¹⁰ In practice, excess bicarbonate is not only wasteful, but also creates problems during work-up (foaming) upon acidification. In contrast, the use of alternative CO₂ sources, such as pressurized or sub/supercritical CO₂ for biocatalytic carboxylation is not well investigated. So far, biocatalytic carboxylations were only successful when additional HCO₃[−] (2–3 M) was applied.¹¹ In order to develop an operationally simple protocol amenable to scale-up, the use of pressurized CO₂ gas was investigated in the carboxylation of 1,3-dihydroxybenzene (**1**, resorcinol, Fig. 1a) as a test substrate using 2,3-dihydroxybenzoic acid decarboxylase from *Aspergillus oryzae* (2,3-DHBD_Ao),¹² 2,6-dihydroxybenzoic acid decarboxylase from *Rhizobium* sp. (2,6-DHBD_Rs)¹³ and salicylic acid decarboxylase from *Trichosporon moniliiforme* (SAD_Tm),^{10a} which are highly active in the presence of bicarbonate.^{9–11} Special emphasis was devoted to pressure and pH effects on enzyme stability.

The exposure of enzymes to scCO₂‡ pressure has an impact on activity, stability or selectivity,¹⁴ which is either due to con-

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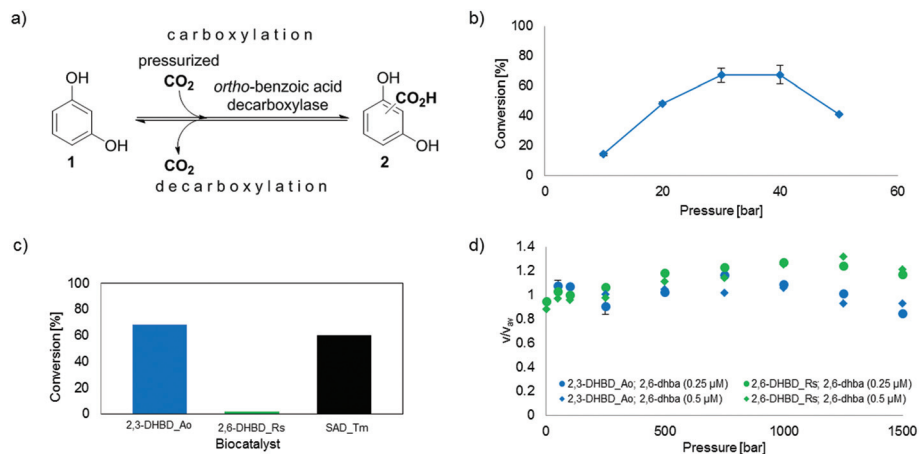


Fig. 1 (a) Enzyme-catalyzed de/carboxylation of resorcinol (**1**). Carboxylated product **2** is a mixture of regio-isomeric 2,6- (2,6-dhba, **2a**) and 2,4-dihydroxybenzoic acid (2,4-dhba, **2b**) with a ratio of 3 : 4;^{10b} (b) CO₂ pressure dependence of the carboxylation of **1** using 2,3-DHBD_Ao; (c) carboxylation activity of decarboxylases with CO₂ (30 bar) using **1** as a substrate; (d) stopped-flow measurements of the decarboxylation of 2,6-dhba (**2a**) with pressure-pretreated (≤ 1.5 kbar) 2,3-DHBD_Ao and 2,6-DHBD_Rs.

formational changes in their secondary and tertiary structure¹⁵ or due to the chemical modification of basic amino acid residues (e.g. Lys, Arg, His) by *N*-carboxylation forming carbamates.¹⁶ The most prominent is the carboxylation of lysine residues (e.g. in RuBisCO,¹⁷ urease¹⁸), which is required for structural reasons (e.g. ligand for binding of metal ions in RuBisCO)¹⁷ or even mandatory for catalysis (e.g. β -lactamase OXA-10 from *Pseudomonas aeruginosa*,¹⁹ biotin-dependent enzymes²⁰). In contrast to these rare beneficial effects, the scCO₂ treatment of enzymes was reported to cause a decrease or complete loss of enzyme activity due to enforced conformational changes (e.g. horseradish peroxidase,^{15b} lipase,^{15a} tyrosinase^{15d}).

Experimental

General

Resorcinol (**1**) and 2,6-dihydroxybenzoic acid (**2a**) were purchased from Sigma Aldrich and 2,4-dihydroxybenzoic acid (**2b**) was obtained from Fluka Analytical. SYPRO orange was purchased from Invitrogen. The pressure reactor system (DigiCAT-system from the HEL Group, volume 16 mL) was equipped with a gas inlet and a magnetic stirrer. A HisTrapFF column with a Ni-NTA Matrix and a PD10 desalting column were obtained from GE Healthcare and Vivaspin 20 (30 kDa) was obtained from Sartorius AG. CO₂ gas (3.0 = 99.9% purity) was obtained from the Linde Group. High pressure stopped-flow measurements were performed with a Hi-Tech Scientific HPSF-56 high pressure stopped-flow spectrophotometer from TgK Scientific. A thermal cycler and a CFX real time system for fluorescence measurements were from Bio Rad and a WebQC calculator was used for pH-calculations.²¹ 2,3-Dihydroxybenzoic acid decarboxylase from *Aspergillus oryzae* (2,3-DHBD_Ao), 2,6-dihydroxybenzoic acid decarboxylase from

Rhizobium sp. (2,6-DHBD_Rs) and salicylic acid decarboxylase from *Trichosporon moniliiforme* (SAD_Tm) were cloned and overexpressed as previously described.²²

General procedure for biotransformation under pressurized CO₂ gas

Lyophilized whole cells (90 mg *E. coli* host cells containing the corresponding overexpressed enzyme with an activity of 5.7 ± 0.7 U mg⁻¹ 2,3-DHBD_Ao and 38.1 ± 0.8 U mg⁻¹ 2,6-DHBD_Rs, respectively) were rehydrated in TRIS-HCl buffer (2850 μ L, pH 9.0, 100 mM) for 30 min. The substrate **1** [10 mM final concentration, dissolved in 150 μ L MeOH (5% v/v)] was added to the enzyme solution (3 mL final volume) which was transferred into a pressure reactor. After CO₂ gas (30 bar) was applied *via* an additional gas inlet for ~ 1 h, the reaction mixture was stirred at 50 rpm for 24 h at 30 °C in the tightly sealed pressure reactor. After 24 h the reaction was stopped by taking 100 μ L of the reaction mixture and diluting it in 900 μ L of H₂O/MeCN/trifluoroacetic acid (TFA, 50 : 50 : 3) to precipitate the enzyme, which was removed by centrifugation (10 min, 14 000 rpm). The supernatant was directly used for measurements on a reversed-phase HPLC system.

For CO₂ pressure studies 10, 20, 30, 40 and 50 bar CO₂ gas was applied.

For buffer concentration studies 100, 250 and 500 mM TRIS-HCl buffer was applied.

The CO₂ pressure pretreatment experiments with 2,6-DHBD_Rs were performed under the same conditions as described above at 10, 40 and 50 bar CO₂ gas (30 mg whole cells, 950 μ L TRIS-HCl buffer, pH 9.0, 100 mM), however, without the addition of the substrate. The pressure pretreated enzyme was then used for the decarboxylation of 2,6-dhba **2a** [final concentration 10 mM, dissolved in 50 μ L MeOH (5% v/v)] in a glass vial (1 mL final volume). The vials were tightly



sealed with screw caps and samples were shaken for 24 h, 120 rpm at 30 °C.

For the determination of the enzymatic activity, the lyophilized whole cells of 2,3-DHBD_Ao and 2,6-DHBD_Rs (10 mg mL⁻¹) were rehydrated in TRIS-HCl buffer (950 μL, pH 9.0, 100 mM) for 30 min. The substrate **2a** (10 mM final concentration) was added to the enzyme solution (1 mL final volume). The vials were shaken at 30 °C with 120 rpm for 0, 1, 2, 4, 6, 8, 10, 12, 15 and 20 min.

All screening experiments were carried out at least in triplicates and all reactor experiments at least in duplicates.

High pressure stopped-flow system

High pressure stopped-flow measurements were performed under a pressure of 1 bar–1.5 kbar using a purified enzyme (0.5 μM 2,3-DHBD_Ao and 2,6-DHBD_Rs, respectively) in TRIS-HCl buffer (pH 9.0, 100 mM) with two different 2,6-dhba concentrations (0.25 and 0.5 μM **2a**, dissolved in 5% v/v MeOH) over 1 min at 30 °C. Spectral changes of the reaction were monitored at 320 nm. All screening experiments were carried out at least in triplicates.

Thermostability experiments

For differential scanning fluorimetry, protein solution [10 μL, 0.2 g L⁻¹ in 5 mM MES (pH 6), 150 mM NaCl], SYPRO orange (10 μL, 1:500 diluted in sterile ultrapure water) and multi-component buffer (pH 4 to 9) (10 μL, 1:2:2 molar ratio of L-malic acid, MES and TRIS; 1 M total concentration)²³ were mixed in 96 well plates. Using a C1000 thermal cycler, the solution was heated at 1.2 °C per minute, from 25 °C to 95 °C. Fluorescence was measured every 0.3 °C, using channel 2 of a CFX real time system. For the smaller step size experiment between pH 4 and 5, sodium citrate buffer (100 mM) was used and the temperature range extended from 10 °C to 95 °C. The melting temperature T_m was calculated as the minimum of the first derivative of the fluorescence *vs.* the temperature. All experiments were carried out in triplicates.

Analytics

HPLC analysis. HPLC/UV experiments were performed on a HPLC Agilent 1260 Infinity system with a diode array detector and a reversed phase Phenomenex Luna column C18 (100 Å, 250 × 4.6 mm, 5 μm, column temperature 24 °C). Conversions were determined by comparison with calibration curves for products and substrates prepared with an authentic reference material. All compounds were spectrophotometrically detected at 263 nm. The method was run over 22 min with H₂O/TFA (0.1%) as the mobile phase (flow rate 1 mL min⁻¹) and a MeCN/TFA (0.1%) gradient (0–2 min 5%, 2–15 min 5–100%, 15–17 min 100%, 17–22 min 100–5%).

HR-MS analysis. HR-MS analysis was performed on a nanoHPLC (Ultimate 3000 RSLCnano system – Dionex) system coupled with q-TOF Maxis II-ETD with an ESI-ionisation in positive mode.

Results and discussion

In spite of the previous reports that carboxylation of phenols requires significant concentrations of bicarbonate (~0.5 M minimum, levelling off at ~1 M to reach a flat plateau at ~3 M) to achieve appreciable conversion levels,¹¹ but fails with CO₂ (gas) alone,^{11b} we tested the use of commercial sparkling water as a medium for the carboxylation of resorcinol (**1**) using 2,3-DHBD_Ao. Surprisingly, conversions of 23% and 7% were measured in water samples containing only 55 mM or 4 mM HCO₃⁻, respectively, which favorably compares to 22% obtained under standard conditions (3 M bicarbonate) (for details see the ESI†).

In order to evaluate the usability of CO₂ (gas) for carboxylation, the influence of various levels of CO₂ pressure (10–50 bar) on the conversion of resorcinol (**1**, 10 mM, TRIS-HCl buffer 100 mM, pH 9.0) using 2,3-DHBD_Ao¹² was determined (Fig. 1b). A bell-shaped curve of the CO₂ pressure with an optimum between 30 and 40 bar was found corresponding to a maximum conversion of 68% of carboxylated product (**2**). The conversion was very low below ≤10 bar and dropped significantly at 50 bar. A time study proved that under these conditions equilibrium was reached at ~24 h (see ESI, Fig. S3†).

In order to examine whether pressurized CO₂ gas (30 bar) is also accepted by other decarboxylases, 2,6-DHBD_Rs¹³ and SAD_Tm^{10a} were tested (Fig. 1c). While SAD_Tm yielded similar results obtained with 2,3-DHBD_Ao (66% and 60% conv., respectively), 2,6-DHBD_Rs did not lead to an appreciable amount of carboxylated product (**2**, conv. <2%). This result corroborates a previous observation, that 2,6-DHBD_Rs is inactive under 50–80 bar of CO₂.^{11b}

To answer the question whether pressure *per se* (a physical consequence) or pressurized carbon dioxide (a chemical effect) is responsible for the inactivation of 2,6-DHBD_Rs, high pressure stopped-flow experiments were performed. For reasons of simplicity, the activity of (hydrostatic) pressure-pretreated 2,6-DHBD_Rs was determined in the (energetically favored) decarboxylation direction with **2a** as a substrate (Fig. 1d). The fairly constant velocity ($v/v_{av} = 0.8–1.3$) of substrate consumption (monitored by a decrease of absorbance at 320 nm) of both enzymes pretreated with up to 1.5 kbar reveals their general pressure stability (Fig. 1d, see also ESI, Fig. S5–S7†). Consequently, the inactivation of 2,6-DHBD_Rs can be explicitly assigned to the action of pressurized CO₂.

In order to determine whether the CO₂ dependent inactivation of 2,6-DHBD_Rs is reversible, the biocatalyst was pretreated with CO₂ pressure (10, 40 and 50 bar, respectively) before measuring its decarboxylation activity (Fig. 2a). The sharp drop in conversion between pretreatments with 40 and 50 bar CO₂ (92% *versus* 40% conv.) clearly indicates that 2,6-DHBD_Rs is irreversibly deactivated beyond ~40 bar CO₂.

Since carbamate formation *via* the carboxylation of lysine residues is a prime suspect for enzyme deactivation, HR-MS measurements were performed. However, no difference in mass between the native and CO₂ pressure (50 bar) treated 2,6-DHBD_Rs was detected, thus inactivation is most likely not



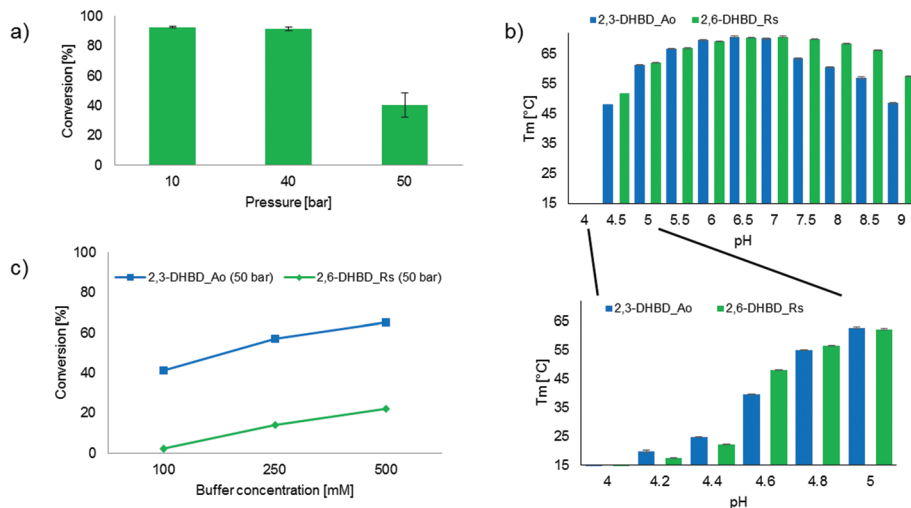


Fig. 2 (a) Decarboxylation activity of CO₂ pressure pretreated (10, 40 and 50 bar) 2,6-DHBD_Rs using **2a** as a substrate; (b) DSF comparison of the pH dependent melting temperature of 2,3-DHBD_Ao and 2,6-DHBD_Rs in multicomponent buffer (pH 4 to 9) and citrate buffer (pH 4 to 5); (c) Influence of TRIS-HCl buffer concentration on the carboxylation of **1** using 2,6-DHBD_Rs and 2,3-DHBD_Ao under 50 bar CO₂ pressure.

caused by the carboxylation of basic amino acid residues (see ESI, Fig. S4†).

Carbon dioxide is readily dissolved in the aqueous reaction medium leading to a drop in pH due to the dissociation of H₂CO₃.²⁴ This effect was applied by Hofland *et al.*²⁵ using CO₂ gas as a ‘volatile acid’ within a range of pH 4–9 to precipitate proteins. To evaluate whether differences in pH-dependent structural stability between 2,3-DHBD_Ao and 2,6-DHBD_Rs could explain their disparate activity, differential scanning fluorimetry (DSF) experiments were performed with both proteins. A first experiment using a multi-component buffer system (L-malic acid, MES, TRIS)²³ shows a broad pH-window from pH 4 to 9, while a second run using citrate buffer and smaller increments reveals details within the pH range of 4 to 5. Overall, 2,3-DHBD_Ao and 2,6-DHBD_Rs behave similarly over the whole pH range. Both are thermally most stable between pH 6–7 and show a continuous decrease in denaturation temperature upon higher or lower pH levels (Fig. 2b). Both enzymes are unstable already at room temperature when the pH of the medium reaches below 4.6.

Given that this pH is likely reached in water in a CO₂ pressurized system (30 bar CO₂ in 0.1 M TRIS-buffer corresponds to a calculated pH of 4.6),²¹ the influence of the buffer capacity was investigated. An increase of buffer concentration/capacity (TRIS-HCl buffer, 100, 250 and 500 mM) to compensate for acidification due to H₂CO₃ dissociation and product formation considerably improved the conversion of the carboxylation of **1** with both enzymes (2,3-DHBD_Ao ~1.5-fold increase; 2,6-DHBD_Rs ~10-fold increase) (Fig. 2c). These results as well as the DSF analysis clearly indicate that the pH value in the pressure chamber is at the edge of the operational pH-window for both enzymes, with 2,3-DHBD_Ao performing slightly better.

Table 1 Comparison of the atom economy of various carboxylation methods of resorcinol (**1**)

Method	Yield [%]	Atom economy [%]	Ref.
Biocatalytic (CO ₂ gas)	68	100	This work
Biocatalytic (HCO ₃ ⁻)	22	73	22
Chemical (Kolbe-Schmitt)	56	52	6b
Chemical (Kolbe-Schmitt)	47	61	26
Chemical (microwave-assisted Kolbe-Schmitt)	62	55	7

Since the economic usage of resources constitutes an important parameter, the atom economy of various *o*-carboxylation systems was compared (Table 1, see the ESI†). An excellent atom efficiency of 100% combined with a good yield (68%) verifies the benefit of the biocatalytic approach using CO₂ (gas). By way of comparison, the biocatalytic alternative using high amounts of bicarbonate shows a significant drop in atom efficiency (73%), which further drops in the case of traditional chemical (52% and 61%, respectively) or microwave-assisted methods (55%).

Conclusion

In summary, we have demonstrated that pressurized carbon dioxide can be used directly as a carboxylating agent in the enzyme catalyzed *o*-carboxylation of a phenol as an alternative to the high concentration of bicarbonate. Two enzyme candidates (2,3-DHBD_Ao and SAD_Tm) readily accepted the alternative CO₂ source for the carboxylation of the model substrate resorcinol. In contrast, 2,6-DHBD_Rs was inapplicable under



CO₂ pressure due to irreversible inactivation, which was correlated to a decrease in pH caused by the dissociation of H₂CO₃.

Overall, the use of pressurized CO₂ gas significantly improves the efficiency of biocatalytic carboxylations and facilitates downstream-processing of this benign and sustainable approach in using CO₂ as a carbon feedstock for the synthesis of organic acids.

Conflicts of interest

There are no conflicts of interest to declare.

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‡scCO₂ = supercritical carbon dioxide.

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