






Cite this: *React. Chem. Eng.*, 2020, 5, 51

Determination of fast gas–liquid reaction kinetics in flow†

Jisong Zhang, ^{ab} Andrew R. Teixeira, ^{bc}
Haomiao Zhang ^b and Klavs F. Jensen ^{*b}

We present a flow concept to measure fast gas–liquid reaction kinetics. A tube-in-tube reactor design with semipermeable Teflon AF-2400 tubes is adopted to achieve fast gas–liquid mass transfer without direct contact of gas and liquid. By performing a steady-state flux balance of both gas and liquid flowing into the reactor and developing a mathematical model based on the film theory, the reaction kinetic parameters can be determined with excellent precision with the use of only a single gas flow meter. Reactions of CO₂ with alkanolamines and ozonolysis of organics serve as case studies to assess the ability of the technique to resolve reaction kinetics *in situ*. The proposed strategy presents a new opportunity for the study of fundamental aspects of gas–liquid reactions in a simple, safe, automated and high-throughput manner.

Received 2nd October 2019,
Accepted 8th November 2019

DOI: 10.1039/c9re00390h

rsc.li/reaction-engineering

Introduction

Gas–liquid reactions are ubiquitous chemical transformations that include well known examples such as absorption of CO₂ into aqueous alkanolamines, absorption of H₂S into ferric sulphate solutions, and ozonolysis, hydrogenation, or chlorination of organic substances.^{1,2} A detailed knowledge of the reaction kinetics and thermodynamic characteristics is essential for developing a new and efficient chemical process. For example, to generate an efficient CO₂ capture process, it is highly desirable to study the reaction kinetics of CO₂ with amines as a function of temperature, amine type and concentrations.^{3,4} Ozone is widely used as an oxidant in drinking water and wastewater treatments, and chemical synthesis using ozone has attracted considerable attention.^{5,6} However, it is challenging to perform kinetic studies with ozone due to safety concerns with the strong oxidizer and the facile and rapid decomposition of O₃.⁷ Recent efforts in micro-scale multiphase flow have revealed strategies to understand and minimize interfacial mass transfer limitations in reacting systems.^{8,9} This, in turn, has led to microreactor designs that operate in the absence of transport limitations and approach the desired kinetic rate limit, thus

motivating the need for a precise understanding of intrinsic multiphase kinetics.

The determination of gas–liquid reaction kinetics is often performed with a stirred cell reactor by recording the pressure decay over time with no analysis of the liquid phase.^{10,11} This method is simple to operate and widely applied. However, it can involve a labour-intensive reloading operation and large variations due to its sensitivity to stirring conditions and poorly defined gas–liquid interface.¹² Continuous systems such as laminar jets,¹³ wetted wall columns¹⁴ and microfluidic contactors^{4,15} partially solve this problem, but they are more complex to operate and rippling on the liquid surface could lead to erroneous results. For more details of such techniques, readers are referred to relevant reviews.^{12,16}

Recently, several gas–liquid reactions such as oxidation,^{17,18} hydrogenation,¹⁹ carboxylation,²⁰ methoxycarbonylation²¹ and ozonolysis²² have been applied in a class of membrane reactors named “tube-in-tube” reactors, which are based on a gas-permeable Teflon AF-2400 membrane. In this strategy, gas diffuses through the gas-permeable membrane where it is then equilibrated with a contacting liquid stream. The reactor achieves rapid gas–liquid mass transfer rates (10–30 s)²³ owing to the small diffusion length scales (0.3 mm) while avoiding the direct and chaotic contact of the gas and liquid. The liquid flow in the inner tubing is laminar so the mass transfer process can be precisely described.²⁴ The physical separation of gas and liquid flow by the Teflon AF membrane produces a well-defined gas–liquid interface at the inner tubing wall, which effectively circumvents erroneous surface rippling and hydrodynamically varying mass transfer rates ($k_L a$).

^a The State Key Laboratory of Chemical Engineering, Department of Chemical Engineering, Tsinghua University, Beijing 100084, China

^b Department of Chemical Engineering, Massachusetts Institute of Technology, Cambridge, MA 02139, USA. E-mail: kjensen@mit.edu

^c Department of Chemical Engineering, Worcester Polytechnic Institute 100 Institute Road, Worcester, MA 01609, USA

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c9re00390h



Additionally, this highly stable reactor works well with a wide range of reactive gases and liquids because of the chemically inert fluoropolymer characteristics. Woodley *et al.*²⁵ developed an automated system to collect kinetic data of enzyme-mediated oxidation based on this tube-in-tube reactor demonstrating that the precise control of O₂ concentration in the solution allows accurate studies of oxidation kinetics. Thus, the tube-in-tube reactor is a highly promising tool to study the fundamental aspects of gas-liquid reactions.

In our previous work, we have developed a fully automated strategy based on the tube-in-tube reactor geometry for fast *in situ* measurements of gas solubility²¹ and diffusivity²⁴ in liquids. Here we extend the application of this strategy to the determination of reaction kinetics. A mathematical model of gas-liquid mass transfer process in the reactor coupled with chemical reactions is first developed and validated. This model is then used to determine the kinetic reaction rate from gas uptake data at various liquid flow rates. The technique is illustrated with two reaction systems, CO₂ absorption and ozonolysis, demonstrating both good accuracy and efficiency.

Experimental

Carbon dioxide (99.99%, pure) and oxygen (99.994%, pure) were all supplied by Airgas (Salem, NH). Methyl diethanolamine (99%) and propionaldehyde (98%) were obtained from Sigma-Aldrich. Deionized water was obtained from VWR, meeting ASTM type II specifications.

Fig. 1 shows a schematic illustration of the automated system, which builds on our previous flow platforms for measuring gas solubility²³ and diffusivity.²⁴ In brevity, the system is composed of the tube-in-tube reactor, a pump, a gas flow meter and a back pressure regulator. Before experiments were conducted, all liquid feeds were degassed according to literature procedures²⁶ to avoid erroneous

effects of the dissolved air on the kinetic measurements. The tube-in-tube reactor was composed of inner Teflon AF-2400 tubing (O.D. 0.8 mm, I.D. 0.6 mm) and outer PTFE tubing (O.D. 3.18 mm, I.D. 1.59 mm) with a length of 0.7 m. During experiments, liquid solutions were pumped from a syringe pump (Harvard Apparatus PHD 2000), through a PTFE feed tube (O.D. 1.59 mm, I.D. 1.2 mm, length 2 m), into the inner reactor tube. The pressure of liquid flow in this tube-in-tube reactor was controlled by a back pressure regulator placed at the outlet of the liquid flow. The reactive gas was delivered through an impermeable PTFE tube (O.D. 1.59 mm, I.D. 1.2 mm, length 2 m) into the shell side of the tube-in-tube reactor from the regulated cylinder. A shut-off valve (IDEX, P-732) placed at the outlet of the gas flow prevented gas flow out of the reactor during the experiments. Hence, the inlet pressure regulator solely controlled the gas flow pressure.

The gas flow was measured by a thermal mass flow controller (Brooks Instruments, 5850i, 5 sccm O₂) located downstream of the pressure regulator but just upstream of the tube-in-tube reactor inlet. The mass flow controller was set to a fully open state, effectively acting solely as a gas flow meter in the experiments. When using gases other than O₂, the flow meter was used with a gas-specific correction factor provided by Brooks Instruments or calibrated following the method reported in our previous study.²² Two pressure transducers (Omega PX409) at the inlets of the reactor monitored the pressures of the gas and liquid flows separately. The feed tubing and the entire tube-in-tube reactor were all immersed in a stirred water bath with a PID controlled immersion heater to ensure an isothermal operation. LabVIEW software controlled the liquid flow and the temperature of the water bath while acquiring gas flow rates and pressures.

Measurement principles

Most gas-liquid reactions are second-order reactions with first order dependencies on gas A and reactant B,

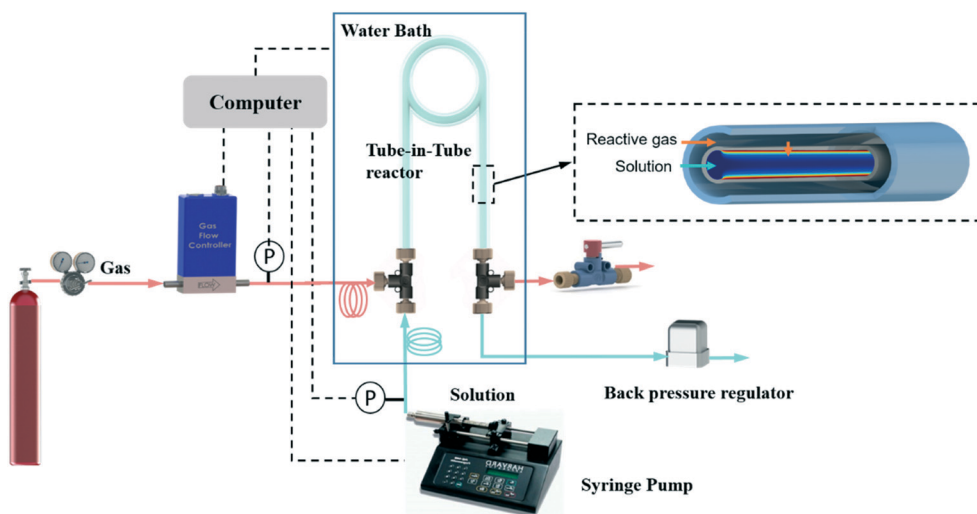


Fig. 1 Schematic of the automated flow system for gas-liquid reaction kinetics determination.



respectively. To simplify the kinetic model, reaction kinetics are usually measured under pseudo-first-order regimes whereby an excess liquid reagent is present and reaction rates are measured at low conversion. To describe the gas-liquid mass transfer, the well-established two-film theory²⁷ is usually adopted, describing the mass transfer as molecular diffusion through a stagnant gas/liquid layer of thickness δ and the bulk gas/liquid is well mixed as depicted in Fig. 2.

The Hatta number, Ha , describes the ratio of reaction to diffusion rates in the thin film:

$$Ha = \sqrt{kC_B D_A} / k_L \quad (1)$$

When the reaction is in the pseudo-first-order regime, the concentration of reactant B in the liquid film is considered constant and equal to the bulk liquid concentration (Fig. 2). In that case, eqn (1) transforms to:

$$Ha = \sqrt{k' D_A} / k_L \quad (2)$$

where k' is the pseudo-first-order kinetic constant, $k' = kC_B$. To ensure the reaction is in the pseudo-first-order regime, Ha should fulfil the following conditions:¹⁶

$$2 < Ha \ll E_i \quad (3)$$

where E_i is the enhancement factor for an irreversible instantaneous reaction.

$$E_i = 1 + \frac{D_B C_B}{\nu_B D_A C_A^*} \quad (4)$$

where ν_B is the stoichiometric coefficient of species B in the reaction.

An illustration relating the mathematical model to determine reaction kinetics is included in Fig. 3. In brevity, the gas mass transfer flux, N_A , can be directly measured by transforming the gas flow rate.

$$N_A = \frac{F_G}{V_s 2\pi RL} \quad (5)$$

According to the Danckwerts mass transfer model,²⁸ the mass transfer flux N_A in the fast pseudo first order regime is proportional to the liquid phase equilibrium concentration:

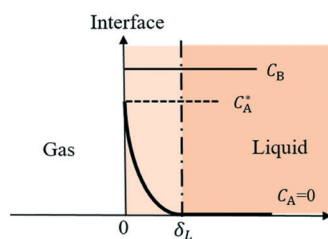


Fig. 2 Concentration profiles in the liquid film for gas absorption with chemical reactions in pseudo-first-order regime.

$$\left[\begin{array}{l} N_A = KC_A^* = KHP_A \\ N_A = \frac{F_G}{V_s 2\pi RL} \\ \text{One } F_G \text{ data point} \end{array} \right] \left[\begin{array}{l} \text{Known } k_m \\ 1/K = R/k_m R_m + 1/k_L' \end{array} \right] \left[\begin{array}{l} \text{Known } k_L, D_A \\ k_L' \\ k_L' = k_L \sqrt{1 + Ha^2} \\ Ha = \sqrt{k' D_A} / k_L \end{array} \right] \left[\begin{array}{l} k' \\ k' = kC_B \end{array} \right] \left[\begin{array}{l} k \end{array} \right]$$

Fig. 3 Determination of gas-liquid reaction kinetics based on the model and experimental data.

$$N_A = KC_A^* \quad (6)$$

According to Henry's law ($C_A = HP_A^*$), the following equation is obtained:

$$N_A = KHP_A \quad (7)$$

By equating eqn (5) and (7), the overall mass transfer coefficient, K , can be directly determined from the measured gas flow, F_G , as the only factor requiring experimental determination. This overall mass transfer coefficient is a weighted average of the respective liquid-side and membrane-side mass transfer coefficients. As the mass transfer across the membrane k_m is known, the mass transfer coefficient in liquid with chemical reaction, k_L^* , can now be deconvoluted from the overall mass transfer coefficient according to the resistance-in-series model^{29,30} as follows:

$$\frac{1}{K} = \frac{R}{k_m R_m} + \frac{1}{k_L^*} \quad (8)$$

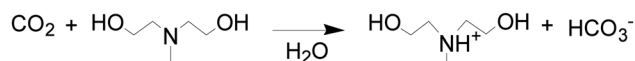
The Ha value is then determined based on the following relationship.³¹

$$k_L^* = k_L \sqrt{1 + Ha^2} \quad (9)$$

Finally, the value of k' and corresponding reaction rate constant, k , can be calculated. It should be noted that the thermodynamic and transport characteristics (gas solubility and diffusivity in liquids) are also required for the determination of reaction kinetics. Both can be obtained in this platform according to the aforescribed methods.^{23,24} In addition, the reactor should operate in a laminar flow regime. The maximum Reynolds number in the current experiments is only 96.

Experimentally, a constant rate of liquid flow was applied and allowed to achieve well-developed steady flow in the inner tube; the corresponding constant gas flux was measured at the inlet flow meter capturing the flux through the membrane into the liquid at steady state. For kinetics measurements in the gas-liquid reaction system, a relatively high liquid flow rate is required to achieve the low conversions needed to accurately measure the reaction rate constant. This original strategy overcomes the slow sampling and poor interfacial control present in classical pressure decay and microfluidic approaches while retaining the





Scheme 1 Chemical absorption of CO₂ into aqueous MDEA.

advantages of easy operation, fast transport time and improved safety.

Results and discussion

To demonstrate the accuracy and efficiency of this automated platform for reaction kinetics determination, we considered two case studies: absorption of CO₂ into aqueous methyl diethanolamine (MDEA) (Scheme 1) and ozonolysis of propionaldehyde (Scheme 2).

CO₂ absorption kinetics

MDEA is a tertiary amine, which can provide a higher capacity, smaller reaction enthalpy with CO₂ and lower vapour pressure compared with traditionally used primary and secondary amines.³² Therefore, the reaction kinetics of CO₂ with aqueous MDEA are well described in the literature using different techniques.^{33–35} To ensure that the reaction is in the pseudo-first-order regime ($2 < \text{Ha} \ll E_i$), the following conditions were chosen: temperature: 20, 40 and 60 °C; MDEA concentration: 10 and 20 wt%.

Fig. 4 shows the measured transient gas flow rate as the liquid flow rate is stepped from 0.2 to 2.7 ml min^{−1} during the reaction. At low liquid flow rates, F_G increases with increasing F_L before becoming nearly constant as F_L is increased above 1 mL min^{−1}. However, according to eqn (5) and (7), F_G should be independent of F_L across the entire flow window for a reaction-controlled process. This behaviour occurs at low flow rates because the large absorption rate of CO₂ leads to a decrease of MDEA concentration in the liquid phase, which invalidates the pseudo-first-order approximation that MDEA is in excess and not changing substantially over the course of the experiment. When the liquid flow rate is sufficiently large, the decrease of MDEA concentration is negligible and the measured gas flow rate is constant. Therefore, a relatively high liquid flow rate was used to ensure that the conversion of MDEA remained less than 2%. The reaction enthalpy of absorption of CO₂ in MDEA solution is only in the range of 20–30 kJ mol^{−1}.³⁶ As a result, the temperature rise due to the heat of reaction is negligible. In addition, there is no increase in the liquid phase molar volume from the dissolution of gas into the liquid because of the small amount of gas absorption. The typical measurement process progressed by recording the gas flow rate at a large liquid flow rate and then the kinetic rate

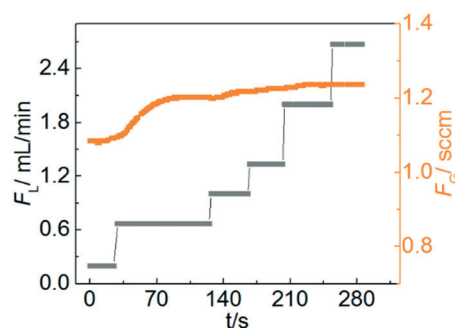


Fig. 4 Measured gas flow rate as a function of set liquid flow rate for the CO₂–MDEA system. Gas pressure: 1.1 bar; temperature: 40 °C; MDEA concentration: 20 wt%.

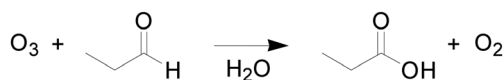
constant was calculated. It is recommended that at least two liquid flow rates are tested to verify that the measured gas flow rate is constant in the experiments. To improve the accuracy, the flow rate at several pressures can be recorded and linear regression of gas flow rate *versus* pressure should be performed to extract the overall mass transfer coefficient in accordance with eqn (5) and (7).

The final measurement results including reaction rate constants at three temperatures and two MDEA concentrations are summarized in Table 1, while the detailed calculations are outlined in Table S1.† Fitting k values *versus* temperature with an Arrhenius relationship, the activation energy is determined to be 52.9 ± 4.4 kJ mol^{−1}, in good agreement with the literature value (stirred cell reactor) of 49.2 kJ mol^{−1}.³⁴ The uncertainty in the activation energy results from the standard error of linear fitting.

Ozonolysis kinetics

There are two obstacles to determining ozonolysis kinetics: 1) ozone gas from an ozone generator is not pure O₃, but rather a mixture of O₃ and O₂. The current experimental setup is not suitable for the mixed gas because the outlet of gas flow is blocked and the unreacted gas would accumulate as the reaction progresses. This would decrease the partial pressure of reacted gas in the gas phase of the reactor and thus would decrease the corresponding O₃ gas flux. 2) There is no reported literature on the Teflon AF 2400 O₃ permeability, which is needed for the above analysis.

To measure gas–liquid reaction kinetics involving a mixed gas stream, the system required some modifications (Fig. S1†) to prevent the accumulation of diluent gases present in the reacting gas mixture from the ozone generator. A back pressure regulator was placed on the gas flow outlet instead of the shut-off valve. The gas flowing into the reactor was controlled by a thermal gas flow controller at a constant rate and a gas flow meter was placed downstream of the reactor before the back pressure regulator to measure the gas flow rate exiting the reactor. The corresponding gas flux into the liquid was calculated with the difference of these two gas flows. To account for the decreased driving force for O₃ into



Scheme 2 Ozonolysis of propionaldehyde.



Table 1 Determination of reaction rates of the CO₂-MDEA system^a

| <i>T</i> (°C) | <i>C</i> _{MDEA} (mol L ⁻¹) | <i>F</i> _G / <i>P</i> _A (sccm bar ⁻¹) | <i>k'</i> (s ⁻¹) | <i>k</i> _{average} (L mol ⁻¹ s ⁻¹) |
|---------------|---|---|------------------------------|--|
| 20 | 0.85 | 1.13 | 1.6 | 2.0 ± 0.1 |
| | 1.71 | 1.19 | 3.4 | |
| 40 | 0.84 | 1.12 | 5.9 | 6.7 ± 0.3 |
| | 1.69 | 1.17 | 11.2 | |
| 60 | 0.84 | 1.08 | 15.7 | 26.8 ± 0.5 |
| | 1.68 | 1.15 | 45.1 | |

^a Liquid flow rate: 2.7 mL min⁻¹.

the liquid, the length of the tube-in-tube reactor was increased to 2.1 m to increase the value of the measured gas flow rate.

For O₃ permeability, we proposed a simple method to determine it by leveraging the automated system as shown in Fig. 1. A pressure larger than atmospheric pressure at the gas phase of the reactor was set by the upstream pressure regulator. The liquid inlet was sealed to prevent liquid-flow into the system and gas-flow out from it. The back pressure on the inner tube was set to atmospheric pressure with the outlet directly vented. In this way, the gas flow through the Teflon AF membrane can be directly determined with the gas flux through the flow meter. Thus, the gas permeability can be calculated in a similar fashion to established approaches in the literature.³⁷

To determine the permeability of pure O₃, the O₂ flow rate through the membrane was first measured and then that of the O₃ mixture. The permeability of pure O₃ could then be determined with the difference of the two permeances. Table 2 summarizes the results for the O₃ permeability determination. The measured permeability of O₂ was 7.2 × 10⁻⁶ mL (STP) cm cm⁻² s⁻¹ bar⁻¹, which is in good agreement with the established literature value of 7.4 × 10⁻⁶ mL (STP) cm cm⁻² s⁻¹ bar⁻¹.³⁷ Furthermore, the permeability of O₃ was substantially lower, consistent with the size-dependent permeability through Teflon AF. Note that the reading of steady gas flow of the O₃ mixture must be taken immediately after flow stabilization to ensure that the concentration of O₃ between the inner and outer tubes does not drift due to the changes in the permeabilities of O₂ and O₃ through the membrane.

The ozonolysis of propionaldehyde (Scheme 2) was chosen to further verify the ability to extract kinetic rate parameters from a multiphase reaction system with a diluent gas. To suppress the decomposition of O₃ in aqueous solution, pH control (<7) and scavenger addition are typically required during the kinetics determination of ozonolysis.⁷ In our

system, the overall residence time of liquid in the reactor is less than 40 s, during which negligible O₃ decomposition occurs,⁶ so the solution requires no pH control (pH = 7) or scavenger addition. To ensure that the reaction was in the pseudo-first-order regime, the following conditions were chosen: temperature: 20, 30 and 40 °C; propionaldehyde concentration: 1 mol L⁻¹.

The measurement results are given in Table 3 with a detailed calculation process outlined in Table S2.† The decrease of gas flow rate ($F_G = F_G^{\text{in}} - F_G^{\text{out}}$) was used to derive the O₃ absorption rate ($F_{O_3} = F_{O_3}^{\text{in}} - F_{O_3}^{\text{out}}$) by assuming that oxygen is not absorbed and the O₂ flow rate is constant. With the method described in the Measurement principle section, the reaction rate constants were determined. The obtained reaction rate constant (0.83 ± 0.10 L mol⁻¹ s⁻¹ at 20 °C) is lower than that in the literature (2.5 ± 0.4 L mol⁻¹ s⁻¹ at 20 °C),³⁸ which could result from different reaction conditions (pH and scavengers) or errors from the O₃ solubility and diffusivity data. The activation energy is determined to be 48.9 ± 2.1 kJ mol⁻¹, at the upper end of the range of 35–50 kJ mol⁻¹ for ozonolysis of most organic compounds.³⁸

The strategy is demonstrated to be a simple and fast (<2 min) method to measure fast gas-liquid reaction kinetics in multiphase systems. In addition, it can provide direct tabulation determination of reaction kinetics under different conditions such as liquid phase composition, temperature and pressure. Similar to the gas solubility and gas diffusivity determination, the system should be operated below the glass transition temperature of Teflon AF-2400 (240 °C).³⁷ To improve the accuracy of this method, the gas flow rate obtained from the gas flow meter should be large, which is usually larger than 0.1 sccm. As a result, a high pressure (>5 bar) is adopted during the gas solubility and diffusivity determination. During the reaction kinetics determination, low pressures (<5 bar) prove to be accurate due to the enhancement of chemical reactions in terms of mass transfer

Table 2 Parameters for the determination of O₃ permeability^a

| Temperature | <i>F</i> _{O₂} (sccm) | <i>F</i> _{O₃} ^b (sccm) | <i>κ</i> _{O₂} (mL (STP) cm cm ⁻² s ⁻¹ bar ⁻¹) | <i>κ</i> _{O₃} (mL (STP) cm cm ⁻² s ⁻¹ bar ⁻¹) |
|-------------|--|---|---|---|
| 20 | 3.01 | 3.29 | 7.23 × 10 ⁻⁶ | 1.209 × 10 ⁻⁵ |
| 30 | 3.00 | 3.28 | 7.21 × 10 ⁻⁶ | 1.207 × 10 ⁻⁵ |
| 40 | 2.99 | 3.27 | 7.19 × 10 ⁻⁶ | 1.206 × 10 ⁻⁵ |

^a Gas pressure difference across the membrane: 1.72 bar, inner tube length: 2.1 m. ^b O₃ gas concentration: 14%, O₂ gas concentration: 86%.

Table 3 Determination of reaction rates of the O₃-propionaldehyde system^a

| <i>T</i> (°C) | <i>F</i> _G ⁱⁿ – <i>F</i> _G ^{out} (sccm) | <i>F</i> _{O₃} ⁱⁿ – <i>F</i> _{O₃} ^{out} (sccm) | <i>k</i> (L mol ^{–1} s ^{–1}) |
|---------------|---|---|---|
| 20 | 0.38 | 0.330 | 0.83 ± 0.10 |
| 30 | 0.43 | 0.342 | 1.55 ± 0.21 |
| 40 | 0.47 | 0.357 | 3.01 ± 0.41 |

^a Propionaldehyde concentration: 1 mol L^{–1}, liquid flow rate: 2.7 mL min^{–1}, gas pressure: 2.23 bar, O₃ concentration: 14%.

rate. Notably, this automated system can work at even higher pressures (<7 MPa) without causing mechanical strain on the Teflon tubing and the fittings. Also, as with gas diffusivity determination,²⁴ the mass transfer coefficient across the membrane *k*_m should be close to or much larger than the mass transfer coefficient in liquid flow *k*_L to ensure the accuracy of measurement. This should be confirmed before performing the reaction kinetics measurement for other reactants or temperatures.

Conclusions

In this paper, we have developed a flow strategy to measure fast gas-liquid reaction kinetics based on the tube-in-tube reactor geometry using gas-permeable Teflon AF-2400. The reaction rate constant can be determined with only a measured gas flow rate within 2 min. Using the reaction of CO₂ with an alkanolamine and the challenging ozonolysis of organics as the examples, we demonstrate the viability of this technique for rapid determination of reaction rate constants for multiphase, multicomponent gas/liquid systems. The determined kinetic parameters using this technique agree well with the literature values from classical batch analyses. The proposed strategy opens a new opportunity for the studies of kinetics of fast gas-liquid reactions in a simple, automated and high-throughput manner.

Nomenclature

- C*_B Liquid phase molar species B concentration (mol L^{–1})
- C*_A^{*} Equilibrium concentration of gas in the liquid, determined by Henry's law (*C*_A^{*} = *H**P*_A, in which *H* is Henry's constant) (mol L^{–1})
- D*_A Gas A diffusivity in the liquid (m² s^{–1})
- F*_G Gas flow rate into the system measured by the gas thermal mass flow meter (sccm)
- F*_L Liquid flow rate into/out of the system, user-defined by the pump (mL min^{–1})
- H* Henry's constant (mol L^{–1} bar^{–1})
- K* Overall mass transfer coefficient (m s^{–1})
- k*_m Mass transfer coefficient across the membrane (m s^{–1})
- k*_L Mass transfer coefficient in liquid flow (m s^{–1})
- k*_L^{*} Mass transfer coefficient in liquid flow with chemical reaction (m s^{–1})

- k* Kinetics rate constant, (L mol^{–1} s^{–1})
- k*' Pseudo-first-order kinetic constant, *k*_B (s^{–1})
- L* Length of the tube-in-tube reactor (m)
- N*_A Gas mass transfer flux (mol m^{–2} s^{–1})
- V*_s Volume occupied by 1 mole gas under standard conditions (22.414 L mol^{–1})
- R* Inner radius of the inner tube (m)
- R*_o Outer radius of the inner tube (m)
- R*_m Logarithmic mean radius of the inner tube (m)
- P*_L^V Vapour pressure of liquid (bar)
- P*_L Liquid pressure (bar)
- P*_G Gas pressure (bar)
- P*_A Partial pressure of gas A in the gas phase, *P*_G – *P*_L^V (bar)
- ν* Stoichiometric coefficient of species *j* in reaction
- κ* Gas permeability (mL (STP) cm^{–1} cm^{–2} s^{–1} bar^{–1})

Dimensionless numbers

- Ha* Hatta number, *Ha* = √(*kC*_B*D*_A)/*k*_L
- E*_i Enhancement factor for an irreversible instantaneous reaction, *E*_i = 1 + 1 / (ν*D*_i*C*_i^{*} / *D*_j*C*_j)

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Financial support from the Novartis-MIT Centre for Continuous Manufacturing is acknowledged.

Notes and references

- G. T. Rochelle, *Science*, 2009, 325, 1652–1654.
- M. Brzozowski, M. O'Brien, S. V. Ley and A. Polyzos, *Acc. Chem. Res.*, 2015, 48, 349–362.
- M. Abolhasani, A. Gunther and E. Kumacheva, *Angew. Chem., Int. Ed.*, 2014, 53, 7992–8002.
- W. Li, K. Liu, R. Simms, J. Greener, D. Jagadeesan, S. Pinto, A. Gunther and E. Kumacheva, *J. Am. Chem. Soc.*, 2012, 134, 3127–3132.
- M. D. Lundin, A. M. Danby, G. R. Akien, P. Venkitasubramanian, K. J. Martin, D. H. Busch and B. Subramaniam, *AIChE J.*, 2017, 63, 2819–2826.
- J. L. Sotelo, F. J. Beltran, F. J. Benitez and J. Beltran-Heredia, *Ind. Eng. Chem. Res.*, 1987, 26, 39–43.
- T. T. Dang, P.-F. Biard and A. Couvert, *Ind. Eng. Chem. Res.*, 2016, 55, 8058–8069.
- A. Gavrilidis, A. Constantinou, K. Hellgardt, K. K. Hii, G. J. Hutchings, G. L. Brett, S. Kuhn and S. P. Marsden, *React. Chem. Eng.*, 2016, 1, 595–612.
- C. Yang, A. R. Teixeira, Y. Shi, S. C. Born, H. Lin, Y. Li Song, B. Martin, B. Schenkel, M. Peer Lachegurabi and K. F. Jensen, *Green Chem.*, 2018, 20, 886–893.
- B. Lu, X. Wang, Y. Xia, N. Liu, S. Li and W. Li, *Energy Fuels*, 2013, 27, 6002–6009.



- 11 L. Kucka, J. Richter, E. Y. Kenig and A. Górak, *Sep. Purif. Technol.*, 2003, **31**, 163–175.
- 12 P. D. Vaidya and E. Y. Kenig, *Chem. Eng. Commun.*, 2007, **194**, 1543–1565.
- 13 N. Ramachandran, A. Aboudheir, R. Idem and P. Tontiwachwuthikul, *Ind. Eng. Chem. Res.*, 2006, **45**, 2608–2616.
- 14 Q. Li, Y. Wang, S. An and L. Wang, *Energy Fuels*, 2016, **30**, 7496–7502.
- 15 C. Zheng, B. Zhao, K. Wang and G. Luo, *AIChE J.*, 2015, **61**, 4358–4366.
- 16 H. Kierzkowska-Pawlak, *Ecol. Chem. Eng. S.*, 2012, **19**, 175–196.
- 17 G. Wu, A. Constantinou, E. Cao, S. Kuhn, M. Morad, M. Sankar, D. Bethell, G. J. Hutchings and A. Gavriilidis, *Ind. Eng. Chem. Res.*, 2015, **54**, 4183–4189.
- 18 T. P. Petersen, A. Polyzos, M. O'Brien, T. Ulven, I. R. Baxendale and S. V. Ley, *ChemSusChem*, 2012, **5**, 274–277.
- 19 M. O'Brien, N. Taylor, A. Polyzos, I. R. Baxendale and S. V. Ley, *Chem. Sci.*, 2011, **2**, 1250.
- 20 A. Polyzos, M. O'Brien, T. P. Petersen, I. R. Baxendale and S. V. Ley, *Angew. Chem., Int. Ed.*, 2011, **50**, 1190–1193.
- 21 P. Koos, U. Gross, A. Polyzos, M. O'Brien, I. Baxendale and S. V. Ley, *Org. Biomol. Chem.*, 2011, **9**, 6903–6908.
- 22 M. O'Brien, I. R. Baxendale and S. V. Ley, *Org. Lett.*, 2010, **12**, 1596–1598.
- 23 J. Zhang, A. R. Teixeira, H. Zhang and K. F. Jensen, *Anal. Chem.*, 2017, **89**, 8524–8530.
- 24 J. Zhang, A. Teixeira, H. Zhang and K. F. Jensen, *Anal. Chem.*, 2019, **91**, 4004–4009.
- 25 R. H. Ringborg, A. Toftgaard Pedersen and J. M. Woodley, *ChemCatChem*, 2017, **9**, 3285–3288.
- 26 R. Battino, M. Banzhof, M. Bogan and E. Wilhelm, *Anal. Chem.*, 1971, **43**, 806–807.
- 27 W. K. Lewis and W. G. Whitman, *Industrial & Engineering Chemistry*, 1924, **16**, 1215–1220.
- 28 J.-C. Charpentier, *Adv. Chem. Eng.*, 1981, **11**, 1–133.
- 29 S. Atchariyawut, C. Feng, R. Wang, R. Jiratananon and D. T. Liang, *J. Membr. Sci.*, 2006, **285**, 272–281.
- 30 A. Mansourizadeh and A. F. Ismail, *J. Hazard. Mater.*, 2009, **171**, 38–53.
- 31 M. M. Sharma and P. V. Danckwerts, *Br. Chem. Eng.*, 1970, 522–528.
- 32 P. D. Vaidya and E. Y. Kenig, *Chem. Eng. Technol.*, 2007, **30**, 1467–1474.
- 33 A. Jamal, A. Meisen and C. Jim Lim, *Chem. Eng. Sci.*, 2006, **61**, 6590–6603.
- 34 H. Kierzkowska-Pawlak and A. Chacuk, *Ecol. Chem. Eng. S.*, 2010, **17**, 463–475.
- 35 N. J. M. C. Penders-van Elk, P. W. J. Derks, S. Fradette and G. F. Versteeg, *Int. J. Greenhouse Gas Control*, 2012, **9**, 385–392.
- 36 M. Gupta, E. F. da Silva, A. Hartono and H. F. Svendsen, *J. Phys. Chem. B*, 2013, **117**, 9457–9468.
- 37 I. Pinnau and L. G. Toy, *J. Membr. Sci.*, 1996, **109**, 125–133.
- 38 J. Hoigné and H. Bader, *Water Res.*, 1983, **17**, 173–183.

