

PAPER

[View Article Online](#)
[View Journal](#) | [View Issue](#)

On the use of molecular-based thermodynamic models to assess the performance of solvents for CO₂ capture processes: monoethanolamine solutions

Charles V. Brand, Edward Graham, Javier Rodríguez, Amparo Galindo, George Jackson and Claire S. Adjiman*

Received 7th March 2016, Accepted 15th April 2016

DOI: 10.1039/c6fd00041j

Predictive models play an important role in the design of post-combustion processes for the capture of carbon dioxide (CO₂) emitted from power plants. A rate-based absorber model is presented to investigate the reactive capture of CO₂ using aqueous monoethanolamine (MEA) as a solvent, integrating a predictive molecular-based equation of state: SAFT-VR SW (Statistical Associating Fluid Theory-Variable Range, Square Well). A distinctive physical approach is adopted to model the chemical equilibria inherent in the process. This eliminates the need to consider reaction products explicitly and greatly reduces the amount of experimental data required to model the absorber compared to the more commonly employed chemical approaches. The predictive capabilities of the absorber model are analyzed for profiles from 10 pilot plant runs by considering two scenarios: (i) no pilot-plant data are used in the model development; (ii) only a limited set of pilot-plant data are used. Within the first scenario, the mass fraction of CO₂ in the clean gas is underestimated in all but one of the cases, indicating that a best-case performance of the solvent can be obtained with this predictive approach. Within the second scenario a single parameter is estimated based on data from a single pilot plant run to correct for the dramatic changes in the diffusivity of CO₂ in the reactive solvent. This parameter is found to be transferable for a broad range of operating conditions. A sensitivity analysis is then conducted, and the liquid viscosity and diffusivity are found to be key properties for the prediction of the composition profiles. The temperature and composition profiles are sensitive to thermodynamic properties that correspond to major sources of heat generation or dissipation. The proposed modelling framework can be used as an early assessment of solvents to aid in narrowing the search space, and can help in determining target solvents for experiments and more detailed modelling.

Department of Chemical Engineering, Centre for Process Systems Engineering, Imperial College London, South Kensington Campus, London SW7 2AZ, UK. E-mail: c.adjiman@imperial.ac.uk; Tel: +44 (0)20 7594 6638

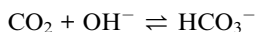
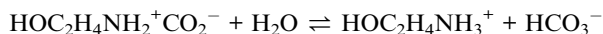
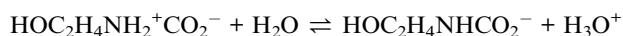
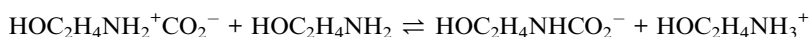
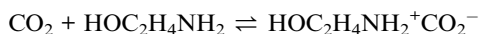


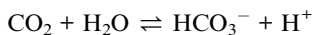
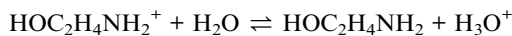
1 Introduction

Carbon dioxide (CO₂) emissions play a major role in climate change and particularly in global warming. Fossil fuel power plants are the major fixed point-source emitters of CO₂. In response to the threat posed by global warming, the Roadmap for 2050 set by the European Commission in 2011 suggested a reduction of greenhouse gas emissions in Europe to 25% by 2020 and to 80% by 2050.¹ In the recent Paris climate conference (COP21), an agreement was made between participating parties to cut greenhouse gas emissions to a level that limits the global average temperature to “well below” 2 °C above pre-industrial levels and to “pursue efforts to limit the temperature increase to 1.5 °C”.² In this context, the development of carbon capture systems must be addressed in the short term, and amine-based post-combustion capture processes are seen as the most promising near-term technology in terms of development and applicability, *e.g.* see Mac Dowell *et al.*³ In this technique, absorption is achieved both physically and chemically, so that significant CO₂ removal can take place even at challenging low partial pressures of the greenhouse gas. The major advantage of this technology is that it can be retrofitted to existing power plants.

There are however several concerns with this technology, in particular the large energy requirements associated with solvent regeneration, the degradation of the solvent, which is exacerbated by the presence of oxygen in the flue gas, and the environmental and health impact that may result from solvent losses and solvent degradation products. These issues are particularly important because of the scale of deployment required to have a meaningful impact on CO₂ emissions. There are significant experimental programmes to identify new solvents^{4–12} and several pilot-plant studies are under way.^{13–21} Modelling studies can play an invaluable and complementary role in addressing some of these issues, including the choice of solvent and operating conditions that yield optimal performance.

A key challenge in realising the benefits of a model-based design approach is the development of models that can accurately predict the behaviour of the CO₂ capture process under different conditions and for a range of solvents. This is particularly difficult in the case of CO₂ absorption due to the complex reaction chemistry that occurs and the large number of ionic species present in the process. For example, in the case of absorption of CO₂ using the most common solvent, an aqueous solution of monoethanolamine (MEA, HOC₂H₄NH₂), the key reactions are:^{22–28}





The main overall reaction products are therefore the zwitterion ($\text{HOC}_2\text{H}_4\text{NH}_2^+\text{CO}_2^-$), the carbamate ($\text{HOC}_2\text{H}_4\text{NHCO}_2^- + \text{HOC}_2\text{H}_4\text{NH}_3^+$), and bicarbonate (HCO_3^-).

Given the complexity of the underlying chemical and physical phenomena, detailed models of the thermodynamics, kinetics, and process units relevant to the absorption systems have been developed for the simulation, optimisation, and design of CO_2 capture processes for a given solvent, as for example in the work of Kucka *et al.*,²⁹ Zhang *et al.*,³⁰ and Kale *et al.*³¹ The elucidation and characterization of the speciation, reaction mechanism, equilibria, and kinetics for mixtures relevant to CO_2 chemisorption are required before detailed models can be developed, and this necessitates extensive experimental investigation. The types of data that are required include data specific to the solvent involved, such as reliable physicochemical (*e.g.* vapour–liquid equilibrium (VLE) and liquid-phase speciation) and kinetic (including reaction rate constants and diffusion coefficients) data, and information regarding the effect of the column specifications, for instance the type of packing material employed. This presents a significant barrier to the rapid development of improved processes for carbon capture.

To overcome this difficulty, the task of identifying solvents that lead to improved CO_2 absorption processes can be subdivided into two main steps. The first is the rapid identification of a list of promising solvents by assessing a wide solvent search space as fully as possible, analyzing key performance indicators that relate closely to process performance, energetics, environmental impact, and solvent degradation. To minimize the reliance on experimental data and accelerate the search for new solvents, models that make it possible to predict physical and chemical properties from molecular structure are highly desirable. The motivation for this first step is to reduce the number of solvents to be considered in more detail, and the models used should therefore offer broad predictive capabilities, which may require making simplifying assumptions in the model development. In the second step, promising solvents can be further analyzed and some of the assumptions made in the first step can be re-assessed. Once a list of candidate solvents is obtained, experiments can be conducted on a subset of these solvents, with the aim of obtaining the information required for a more detailed evaluation of each solvent and the corresponding carbon dioxide process.

To explore the space of possible solvents, there is a need to develop models that offer adequate predictive capabilities without exhaustive reliance on experimental data, and that can provide a quantitative insight into the behaviour of the process; the use of a thermodynamic model to capture the phase and chemical equilibria of mixtures of carbon dioxide, water, and alkanolamine within a process model is investigated in our current paper, as a means of obtaining an estimate of the performance of the absorption. As an initial assessment of the method, we focus on MEA because as a ubiquitous solvent for CO_2 capture there is extensive experimental data available to test the validity of the predictions. Before presenting the model, we first place the proposed approach in the context of other modelling work in the area.



There is an extensive body of literature concerning the modelling and simulation of CO₂ absorption in packed columns. The modelling approaches that have been proposed to date differ in the choice of thermodynamic and kinetic models, and, where appropriate, heat- and mass-transfer models. Most of the effort has been focused on the performance of aqueous monoethanolamine solutions, due to their widespread industrial use and the availability of pilot-plant data, although there have been some models developed for other solvents, notably aminomethylpropanol (AMP).^{16,32–35} In modelling an absorber, the column is usually divided into hypothetical stages, each representing a (sometimes infinitesimal) section of packing in the column.^{36,37} Each stage can be modelled using either an equilibrium or a rate-based model. In an equilibrium model, vapour–liquid equilibrium is assumed at each stage, everywhere on the stage. A rate-based model accounts for heat- and mass-transfer limitations. For chemisorption processes with fast reaction kinetics, as is the case for the process of interest,^{28,38} a rate-based process is more reliable. Indeed, Lawal *et al.*³⁹ have compared the two approaches using the same physical property model and concluded that the rate-based model provides a better description of the pilot-plant¹⁴ temperature profiles where aqueous MEA was used as a solvent. A similar comparison was made by Afkhamipour and Mofarahi³⁵ for CO₂ absorption in AMP solution, and a rate-based model was found to give a better representation of the temperature and composition profiles of the pilot-plant runs.¹³

When using a rate-based approach, an important aspect in model development is the choice of approach to treat heat- and mass-transfer phenomena. The concentration and temperature profiles across hypothetical films in a two-film model can be imposed, taking into account the effect of chemical reactions on mass transfer with an enhancement factor, defined as the ratio of the amount of gas absorbed in a reacting liquid to the amount which would be absorbed if there were no reaction.^{40–42} The enhancement factor varies along the length of the column and can often be adjusted to pilot-plant data. This is the route followed in most models of CO₂ absorption (*e.g.* Tontiwachwuthikul *et al.*,¹³ Sivasubramanian *et al.*,⁴³ Pintola *et al.*,⁴⁴ Pacheco and Rochelle,⁴⁵ Al-Baghli *et al.*,⁴⁶ Tobiesen *et al.*,⁴⁷ Faramarzi *et al.*,⁴⁸ Khan *et al.*,⁴⁹ Neveux *et al.*,⁵⁰ Saimpert *et al.*,⁵¹ Jayarathna *et al.*,⁵² Afkhamipour and Mofarahi⁵³). A comparative review of the rate-based models that have been used to specifically treat CO₂ absorption in aqueous MEA solutions can be found in ref. 54.

As an improvement on the standard film theory, Tobiesen *et al.*⁴⁷ developed a penetration model, where the two films at the interface are described continuously, which was found to describe their own pilot-plant data well. A more rigorous approach is that followed by Kucka *et al.*,²⁹ in which the Maxwell–Stefan formalism is used together with a discretization of the film. This more detailed model leads to better predictions of the concentration and temperature profiles at the pilot-plant scale¹³ than other models, without the need to adjust any parameters to pilot-plant data.²⁹ The Maxwell–Stefan formalism has also been applied by Lawal *et al.*⁵⁵ and Biliyok *et al.*⁵⁶ Kale *et al.*³¹ have investigated a rate-based model with film discretization, and studied the sensitivity of the calculations to several key parameters. They found that good predictions of column profiles can be obtained with a sufficiently fine discretization.

In all rate-based models, empirical mass-transfer correlations are required to account for the type of packing used and the operating conditions. The



correlations of Onda *et al.*^{57,58} and Bravo and Fair⁵⁹ were developed specifically for random packing, whereas the correlations of Rocha *et al.*⁶⁰ were developed for structured packing, but can be applied to random packing by using an equivalence relation linking the random packing characteristics to the structured packing. Correlations developed by Billet and Schultes⁶¹ are also available as they apply to both structured and random packing. In a detailed comparison applied to a model of a CO₂ capture pilot plant presented by Faramarzi *et al.*,⁴⁸ it appears that the main operating conditions to consider when choosing a mass-transfer correlation are the flow rates of the flue gas and the lean solvent.

Another important consideration in modelling CO₂ absorption is the representation of the chemical reactions and fluid-phase equilibria of the mixture of MEA, CO₂ and H₂O. In the earliest absorber models, the thermodynamics of the gas and liquid phases were described with the assumption of ideal gas and ideal solution behaviour. This is the case for example of the model developed by Pandya⁶² and later used by Tontiwachwuthikul *et al.*¹³ However, this model was too simplified to describe the complex interactions between the CO₂ and the solvent, and may not be suitable when transferred to other solvents. Two different approaches are typically followed when developing more accurate models of mixtures exhibiting reaction equilibria: those based on physical theories and those based on chemical theories. Most commonly, a chemical approach (*e.g.* see refs 30 and 63–67) is adopted. In such approaches, all of the reaction species in solution are modelled explicitly, requiring the *a priori* specification of the relevant reaction schemes and their corresponding temperature-dependent equilibrium constants. Hence, the use of a chemical approach requires experimental data on the concentration of species in solution at various temperatures.

Most models proposed to date that specifically treat CO₂–amine–H₂O systems are based on this class of chemical approach; an explicit treatment of the major ionic and non-ionic species formed due to reaction is adopted. Such methods rely on the use of reaction kinetics and equilibrium constants derived from experimental data specific to each reaction.^{29,30,68–73} The methodology of combining the electrolyte-NRTL (eNRTL) approach^{68,74} with a Henry's constant to describe CO₂ solubility has attracted much interest and has been used in recent studies.⁵⁵ The eNRTL approach has also been corrected for inconsistencies by Bollas *et al.*⁷⁵ More recently, Zhang *et al.*³⁰ developed a detailed model of the CO₂–MEA–H₂O amine system using a chemical approach: an eNRTL⁶⁶ model was developed for the treatment of the liquid phase, the perturbed chain statistical associating fluid theory (PC-SAFT)⁷⁶ was used to obtain the vapour phase fugacity coefficients, and Henry's law constants were obtained to provide a full description of the phase equilibria. While this approach arguably provides the most accurate representation of the behaviour of these mixtures to date, a major drawback of these models is that they contain a large number of parameters, requiring ample experimental data for VLE and reaction kinetics, thus making it difficult to use for different conditions and solvents. A similar approach²⁹ is to combine the Soave–Rechlich–Kwong (SRK) equation of state (EoS)⁷⁷ with the eNRTL model. Tobiesen *et al.*⁴⁷ have adapted an activity coefficient model from Hoff⁷⁸ to correlate VLE experimental data, using an experimentally-derived equilibrium constant.⁷⁹ In order to reduce the number of parameters to be determined and the computational effort needed to solve the vapour–liquid equilibrium, Gabrielsen *et al.*⁸⁰ developed their own thermodynamic model to determine the VLE of MEA, CO₂ and H₂O. They



proposed a simple correlation derived from experimental data of CO₂ solubility in aqueous MEA, which is valid for MEA over the conditions considered: in this case CO₂ loading (defined as the number of moles of CO₂ per mole of MEA in the liquid phase) varied from 0 to 0.5 at 313 K and 393 K. As a simplification, a single chemical equilibrium reaction (carbamate formation) was considered, reducing the number of adjustable parameters required.

The combined reaction and activity coefficient models developed to date provide an accurate representation of CO₂–MEA–H₂O systems over a range of conditions. However, extensive parameterization is required and the model parameters obtained are not transferable to other solvents. These models are therefore more suited for their application in step two of the solvent selection process, the detailed quantitative analysis of the performance of the most promising solvents.

An alternative treatment of reactions follows a physical approach. Such approaches have previously been proposed to model the phase and chemical equilibria of mixtures of alkanolamines, CO₂, and H₂O.^{81–84} This methodology is based on a version of the molecular statistical associating fluid theory (SAFT) EoS,^{85,86} in which a square-well intermolecular potential of variable range is used (SAFT-VR).^{87,88} More recently,^{89,90} models for aqueous solutions of CO₂ and primary alkanolamine have been developed within a group contribution reformulation of the SAFT-VR EoS (SAFT- γ SW),^{91,92} where a similar physical treatment of reactions was applied. In these approaches, the reaction products are considered as associated aggregates of the reactants and are not modelled as independent chemical species. Aggregation is driven by short-range association sites incorporated in the molecular models, by means of appropriate site–site interactions. Appropriate association schemes are chosen in order to form the desired reaction products (*e.g.* carbamate, zwitterion, *etc.*). In the SAFT approach, the concentration of each aggregated species in the mixture can be determined from the fraction of association sites that are not bonded.⁹³ In cases where chemical equilibrium can be assumed and under some appropriate approximations (*e.g.* a constant site–site binding energy), such an approach has been shown to be equivalent to the chemical models used to date in which the aggregated chemical species are modelled explicitly.⁹⁴ The equivalence of the chemical and physical approaches means that the parameters within SAFT can be related to the experimental equilibrium constant. The thermodynamic properties of the mixture can thus be determined by specifying the concentration of the reactants, the pressure, temperature, and the parameters describing the interactions between reactant species (including the types, number, energy, and bonding volume of association sites). The physical approach applied to CO₂ and aqueous MEA has been found to provide excellent predictions of the concentration of the key reaction products.^{84,90} Physical approaches are particularly well-suited to the exploration of a wide solvent space in that models can be developed entirely from experimental VLE data (*e.g.* vapour pressure and saturated liquid density) and no data on energetics or speciation are required. In addition, parameters can often be transferred from one compound to another compound on the basis of similarities in molecular chemistry, and the resulting predictions are found to be valid over wide ranges of compositions, temperatures, and pressures. These characteristics make SAFT-VR models, which are based on a physical representation of chemical equilibrium, a suitable platform for the exploration of a large space of



solvent candidates. While the evaluation of potential solvents can be carried out on the basis of thermodynamics alone, it would be more valuable to obtain an evaluation of each solvent based on performance metrics that relate to a specific absorption process; this possibility is investigated in our current work.

The use of the SAFT-VR thermodynamic approach within process models of CO₂ absorption has previously been explored for high-pressure physical absorption in alkanes,⁹⁵ ethers and ether derivatives,⁹⁶ and also for chemisorption. In an early study the modelling of the absorption process was also considered and explored in the context of solvent-blend design.⁹⁷ The absorber model was further developed⁹⁸ showing reliable results for a set of pilot-plant data. Mac Dowell *et al.*⁹⁹ have proposed a dynamic model of a CO₂ absorber, based on the SAFT-VR thermodynamic treatment, which has been used in in-depth studies of the control¹⁰⁰ and economics^{101,102} of post-combustion CO₂ capture processes. Qualitative agreement with data from two pilot-plant runs from ref. 13 was found in these studies.^{99–101} The column profiles obtained by Mac Dowell *et al.*⁹⁹ are somewhat difficult to interpret as the location of the data points reported in the paper does not match the location of the sensors reported in the physical pilot plant. Nevertheless, the findings of this body of work indicate that a physical approach to the modelling the chemical and physical equilibria allows one to capture the process behaviour accurately with a limited set of parameters.

Overall, the models developed to date can be used to adequately represent the general behaviour of the absorber column. However, only the model of Tontiwachwuthikul *et al.*¹³ provides an accurate description of the temperature of the rich solvent at the outlet, and in all of the published models, the bulge in the temperature profile, which is a well-known characteristic of this absorption process,¹⁰³ is reproduced qualitatively but not quantitatively. Existing models reproduce either its magnitude or its location along the column, but not both.

In view of the foregoing discussion, the potential benefits of a novel absorption model in which a SAFT-VR thermodynamic treatment and a rate-based column model are integrated are explored here for CO₂ absorption in aqueous MEA. The proposed approach is based on a two-film model. Because the reactions are treated implicitly with the SAFT-VR EoS, only the key molecular species (*i.e.* MEA, CO₂, H₂O, and N₂) need to be taken into account explicitly at the level of mass and energy balances. The detailed model presented in our current paper includes several modifications over previous work,^{97,104} leading to enhanced model reliability. In Section 2, we describe the SAFT-VR EoS and the heat- and mass-transfer relations used in the rate-based absorber model. The validation of the model, its predictive capabilities, and a sensitivity analysis are presented in Section 3. Two scenarios are investigated, corresponding to different levels of data availability: in the first scenario, it is assumed that no pilot-plant data are available, and the suitability of the model to provide a best-case analysis of process performance is studied; in the second scenario, one pilot-plant run is considered and employed to obtain a more realistic quantification of mass-transfer limitations. The transferability of this analysis is then tested for data at other conditions.

2 Modelling methodology

The development of a rate-based model of a CO₂ absorber is described in this section. In subsection 2.1, the thermodynamic model used for the VLE and the



chemical equilibrium is presented. The non-equilibrium stage approach is introduced in Subsection 2.2.

2.1 Thermodynamic model

The treatment of the chemical-reaction equilibria relevant to the absorption process is a key aspect of the modelling strategy adopted in our work. In the physical approach followed here, the chemical and physical interactions are treated on an equal footing within the SAFT-VR SW EoS.^{87,88} Background information is provided in this section to help the reader better understand the models used and the underlying assumptions.

The SAFT family of equations stems from the first-order thermodynamic perturbation theory (TPT1) of Wertheim.^{93,105–109} In SAFT approaches molecules are modelled as chains of fused spherical segments with embedded short-range association sites incorporated to mediate hydrogen bonding which leads to aggregate formation (speciation). The EoS is developed in terms of the Helmholtz free energy using a perturbation approach, such that the free energy of a chain molecule is obtained with respect to the properties (free energy and radial distribution function) of a reference monomeric (non-bonded) system. The original SAFT^{85,86} EoS has been revisited and modified by a number of researchers so that several versions are now available.^{76,87,88,110–116} A historical account of perturbation theories for polar and associating liquids has recently been presented by Gubbins.¹¹⁷

In the SAFT-VR formulation^{87,88} employed here, a square-well (SW) potential is used to describe the interaction between spherical molecular segments of the reference fluid. The more recent versions of the SAFT-VR EoS incorporate the Mie (generalized Lennard-Jones) potential (SAFT-VR Mie¹¹⁶), and allow for a more reliable representation of the near-critical region and second derivative properties (e.g. heat capacities) which depend mainly on the specific form of the repulsive part of the potential.^{114,116,118,119} Additionally, group contribution (GC)¹²⁰ versions of the SAFT-VR EoSs, namely SAFT- γ SW^{91,92} and SAFT- γ Mie,¹²¹ have also been proposed. These offer additional predictive capabilities in that the properties of a new solvent that has never been synthesized can be predicted (without data specific to that solvent) provided that the interaction parameters for the functional groups appearing in that molecule are available. The assessment of the integration of the molecular-based SAFT-VR SW EoS within an absorption model presented in our current paper can readily be extended to a GC formulation.

In the SAFT-VR SW approach, a molecule i is modelled as a chain of m_i fused homonuclear spherical segments of diameter σ_{ii} . The interactions between two identical segments are described by a square-well potential of range λ_{ii} and depth ε_{ii} (cf. Fig. 1). For each molecule i , the number of site types $N_{s,i}$ must be defined, as well as the number of sites of each type a , $N_{s,ia}$. The sites are characterized by SW site–site energetic $\varepsilon_{ab,ii}^{\text{HB}}$ and range $r_{c,ab,ii}$ parameters (cf. Fig. 1).

In order to model mixtures, combining rules based on the Lorentz–Berthelot form (cf. Haslam *et al.*¹²²) are used to describe the unlike interaction between segments on two different molecules i and j :

$$\sigma_{ij} = \frac{\sigma_{ii} + \sigma_{jj}}{2} \quad (1)$$



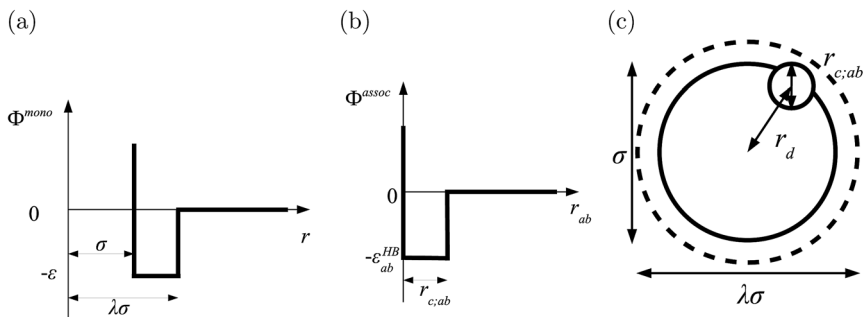


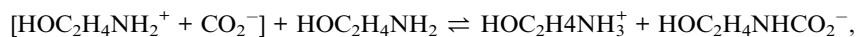
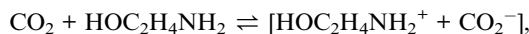
Fig. 1 The square-well potentials for a monomer, Φ^{mono} , and association between the sites, Φ^{assoc} , employed in the SAFT-VR approach. (a) Φ^{mono} is characterized by a hard-core with a diameter σ , range of attraction $\lambda\sigma$ and depth ε . (b) Φ^{assoc} is defined by an off-centre potential of depth $\varepsilon_{ab}^{\text{HB}}$ and of range $r_{c;ab}$. (c) The centre of the site is at a distance r_d from the centre of the segment.

$$\varepsilon_{ij} = (1 - k_{ij}) \sqrt{\varepsilon_{ii} \varepsilon_{jj}} \quad (2)$$

$$\lambda_{ij} = \frac{\sigma_{ii} \lambda_{ii} + \sigma_{jj} \lambda_{jj}}{\sigma_{ij}}, \quad (3)$$

where k_{ij} is an adjustable parameter that characterizes the unlike dispersion attractive interaction. Parameters to describe association between different molecules are estimated using experimental data for the specific mixtures of interest, or for chemically similar mixtures.^{82,84}

In recent work^{82,84,89,90} the reactions involved in aqueous amine solutions of CO_2 are treated implicitly within the SAFT-VR and SAFT- γ frameworks, with the products of the chemical reaction represented as aggregates of the reactant molecules. In the case of MEA the overall set of reactions can be reduced to²⁸



and association sites that mimic the complexation of CO_2 and amine are introduced in the SAFT-VR molecular models. The reaction products can thus be modelled as neutral aggregates of CO_2 and MEA, bonded at association sites as shown in Fig. 2. An accurate overall representation of the vapour-liquid phase equilibria of $\text{MEA} + \text{CO}_2 + \text{H}_2\text{O}$ can be obtained in this manner by estimating the molecular parameters from experimental fluid-phase equilibria data, for both the pure components and mixtures. One important implication of the physical treatment of chemical equilibrium is that there is no need to explicitly specify a reaction scheme or reaction products. The types of products formed (e.g. carbamates or bicarbonates) are dictated by the association scheme chosen (number of sites and strength of their interactions), and the relative extent of formation of the different products depends on the temperature-independent intermolecular parameters that describe the association energies. The fraction of molecules bonded at a given site is an output of the SAFT-VR approach and the



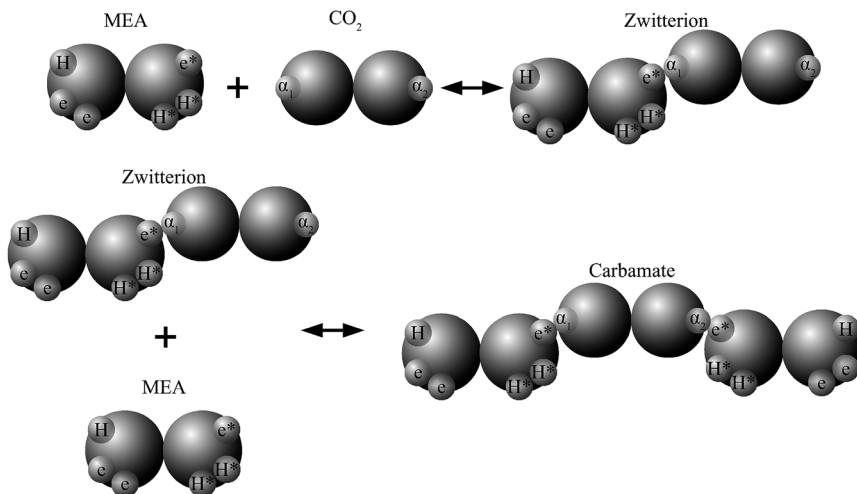


Fig. 2 Schematic representation of the association scheme between MEA and CO₂ (in aqueous media) with a SAFT treatment showing two reaction products.

distribution of reaction products can be determined from a statistical analysis of the values of these fractions at the thermodynamic state of interest.⁹⁴ Thus, although no speciation data are used in deriving the SAFT-VR parameters, the speciation equilibria can nonetheless be predicted successfully.^{84,90}

This type of physical approach greatly reduces the number of parameters needed to describe the mixture compared to explicit approaches such as eNRTL, in which every species must be described as a separate entity and temperature-dependent equilibrium constants must be derived for all the relevant reactions. The SAFT-VR approach has been shown to be applicable to the absorption of CO₂ for a wide range of aqueous alkanolamine solutions.⁸⁴ In many cases, it is possible to transfer parameters from one alkanolamine to another based on molecular similarity, further reducing the need for experimental data.

One key assumption in adopting this type of physical approach is that all reactions are assumed to be at equilibrium, which is only applicable to processes in the physical regime, *i.e.* where mass transfer is the rate limiting process. In the case where a specific treatment of the charged electrolytic species is required, the SAFT-VR approach^{123–128} can be coupled to a chemical approach as appropriate. It is also possible to represent some or all species explicitly where chemical reactions that are not at equilibrium can then be modelled *via* a separate kinetic model. We do not follow this route here, which would fall within the class of chemical approaches, but instead assess the adequacy of the physical approach.

The molecules considered in our study are MEA, H₂O, CO₂, and N₂. A schematic of the molecular models used in the SAFT-VR SW treatment is presented in Fig. 3. The values of all parameters are listed Tables 1–4. The MEA molecule is represented as 2 tangent spherical segments with 2 association sites of type e (electron lone pairs on the oxygen atom), 1 site of type e* (corresponding to the lone pair on the nitrogen atom), 1 site of type H (the hydrogen atom on the hydroxyl group), and two sites of type H* (hydrogen atoms on the amine group).⁸²



The H_2O is represented as 1 spherical segment with 2 e sites and 2 H sites.¹²⁹ The CO_2 model comprises 2 tangent segments and has 1 α_1 site and 1 α_2 site (acceptor sites) that interact only with the e* sites of MEA.⁸⁴ The N_2 molecule is modelled as a fused non-spherical diatomic, with an aspect ratio of $m = 1.4$, and does not have association sites as it is chemically inert and apolar.^{104,130} A site of type e/e* can bond with a site H/H* and *vice versa*. For example, when an e site from MEA and an H site from water come within the cut-off range $r_{\text{c,eH,MEA-H}_2\text{O}}^{\text{HB}}$ of each other, there is a site-site hydrogen-bonding associative interaction of strength $\epsilon_{\text{c,eH,MEA-H}_2\text{O}}^{\text{HB}}$. A representative example of the good overall description of the temperature and pressure dependence of the absorption VLE behaviour of CO_2 in aqueous MEA obtained with the SAFT-VR SW models can be seen in Fig. 4 over several orders of magnitude for the partial pressure of CO_2 . In the standard Wertheim TPT1 treatment at the heart of the SAFT EoS, association into linear-chain, branched-chain, and network aggregates are considered. Association into ring-like structures^{131–134} (and even double bonding¹³⁵ and bond cooperativity¹³⁶) can be taken into account but this is not considered for the systems described in our current work.

2.2 Non-equilibrium absorber model

The absorber is a counter-current vapour–liquid multistage separation column, with a liquid feed at the top stage and a vapour feed at the bottom stage. The vapour product comes off the top stage and the liquid product off the bottom stage. The inside of the column is filled with an inert packing material designed to maximize mass transfer between the vapour and the liquid and to minimize the pressure drop. The vapour and liquid compositions vary continuously with packing height.

The modelling of such a column can be either discrete or continuous in the vertical direction. In a discrete model, the column is divided into hypothetical stages, each of which represents a section of packing in the packed column. For rate-based models, a greater number of stages provides a better description.³⁶ Taken to the limit, the use of an infinite number of stages is equivalent to modelling the column continuously. Although both models could be used; we choose the discrete approach for our study.

The modelling of the stages can be equilibrium or rate-based. In an equilibrium model, it is assumed that vapour–liquid equilibrium is achieved at each stage, everywhere on the stage. A rate-based model accounts for limitations due to

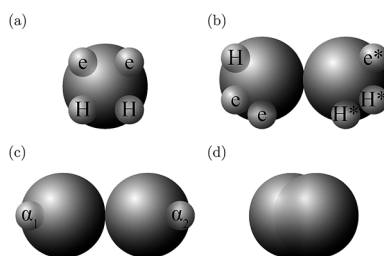


Fig. 3 A schematic of the molecular models used in the SAFT-VR SW approach: (a) H_2O , (b) MEA, (c) CO_2 , and (d) N_2 .



Table 1 The SAFT-VR parameters characterising the pure-component models used in our work: the number of segments m_i , the diameter of the spherical core σ_{ii} , the depth ε_{ii} and range λ_{ii} of the dispersive square-well potential, and the type and the number of hydrogen-bonding sites

i	m_i	σ_{ii} (Å)	ε_{ii}/k (K)	λ_{ii}	e- sites	H- sites	e*- sites	H*- sites	α_1 - sites	α_2 - sites	Source
H ₂ O	1.0	3.03420	250.00	1.78890	2	2	0	0	0	0	Clark <i>et al.</i> ¹²⁹
MEA	2.0	3.57229	305.00	1.58280	2	1	1	2	0	0	Mac Dowell <i>et al.</i> ⁸²
CO ₂	2.0	2.78640	179.27	1.51573	0	0	0	0	1	1	Rodríguez <i>et al.</i> ⁸⁴
N ₂	1.4	3.07357	74.587	1.58795	0	0	0	0	0	0	Mac Dowell ¹⁰⁴

transport phenomena. Due to the complexity of the process modelled in the current system, a rate-based approach is chosen over an equilibrium one in order to capture some of the key physical phenomena.^{137,138}

For the absorber model, the assumption is made that the two phases are distinct, and each phase is perfectly mixed at each stage. A diagram of a non-equilibrium stage is presented in Fig. 5. Vapour from the stage below is brought into contact with liquid from the stage above and the two phases exchange mass and energy through their common interface represented in the diagram by the wavy line. In a rate-based model, separate mass balances are considered for each phase. The two phases are in contact through their interface where material lost by the vapour phase is gained by the liquid phase. The heat transfer is treated in a similar way. There is one energy balance in each phase, and the phases are linked *via* the rate of energy transferred across the phase interface. The compositions of the two phases at the interface are determined assuming equilibrium conditions at the interface.

The behaviour at the phase interface in the rate-based stages is described with a two-film theory.¹³⁷ Each phase is split into two parts: the bulk phase and the film in which the heat and mass transfer occur. In the bulk phase, the concentration, pressure, and temperature are assumed to be uniform, whereas there could be a gradient in composition and temperature in the film. A liquid-gas interface between the two films is sketched in Fig. 6, where the profiles for the composition of component i , temperature, and pressure are represented.

The following assumptions are made:

Table 2 Binary interaction parameters for the mixtures relevant to our work. k_{ij} characterizes the strength of the unlike dispersion interaction between molecules of types i and j , cf. eqn (2)

$i + j$	k_{ij}	Source
MEA + CO ₂	0.4788	Rodríguez <i>et al.</i> ⁸⁴
MEA + H ₂ O	0.01	Mac Dowell <i>et al.</i> ⁸²
MEA + N ₂	0.03	Mac Dowell ¹⁰⁴
CO ₂ + H ₂ O	−0.06	Mac Dowell <i>et al.</i> ⁸²
CO ₂ + N ₂	−0.0599	Mac Dowell ¹⁰⁴
H ₂ O + N ₂	−0.3635	Mac Dowell ¹⁰⁴





Table 3 The site–site association energies $\epsilon_{ab,ij}^{\text{HB}}$ for MEA, H₂O, and CO₂ (cf. Fig. 3). The interaction matrix is symmetrical, i.e. $\epsilon_{e\text{H},ii}^{\text{HB}} = \epsilon_{\text{He},ii}^{\text{HB}}$; the lower diagonal part has been left blank and is implied. The unlike association between sites of the same type is assumed to be symmetric, i.e. $\epsilon_{e\text{H},ij}^{\text{HB}} = \epsilon_{\text{He},jj}^{\text{HB}} = \epsilon_{e\text{H},ji}^{\text{HB}} = \epsilon_{\text{He},ji}^{\text{HB}}$ (ref. 82 and 84)

		$\epsilon_{ab,ij}^{\text{HB}}/k \text{ (K)}$							
		MEA				H ₂ O		CO ₂	
	<i>a/b</i>	e	H	e*	H*	e	H	α_1	α_2
MEA	e	0	2357.79	0	900	0	1780.7121	0	0
	H	—	0	550	0	1780.7121	0	0	0
	e*	—	—	0	960	0	1517.1049	5200	3982.66
	H*	—	—	—	0	1517.1049	0	0	0
H ₂ O	e	—	—	—	—	0	1400.00	0	0
	H	—	—	—	—	—	0	0	0
CO ₂	α_1	—	—	—	—	—	—	0	0
	α_2	—	—	—	—	—	—	0	0

- The model is steady state.
- The bulk phases and films are at chemical equilibrium everywhere (i.e. the reaction rates are faster than the mass transfer rates).^{25,28}
- The interface is at phase and chemical equilibrium.
- The interfacial surface area is the same for both heat and mass transfer.
- The effective area is equal to the wetted area.
- The absorption column is considered to be adiabatic.
- There is no pressure drop along the column.

2.3 Model equations

In this subsection, we present the equations used to model a packed column subsection (stage). The equations are grouped into different categories: energy and mass balances, rate equations, equilibrium equations, mass-transfer

Table 4 The site–site range parameters $r_{c;ab,ij}$ for mixtures containing MEA, H₂O, and CO₂ (cf. Fig. 3). The interaction matrix is symmetrical, i.e. $r_{c;e\text{H},ii} = r_{c;\text{He},ii}$, and so the lower diagonal part has been left blank. The unlike association between sites of the same type is assumed to be symmetric, i.e. $r_{c;e\text{H},ij} = r_{c;\text{He},jj} = r_{c;e\text{H},ji} = r_{c;\text{He},ji}$ (ref. 82 and 84)

		$r_{c;ab,ij} \text{ (Å)}$							
		MEA				H ₂ O		CO ₂	
	<i>a/b</i>	e	H	e*	H*	e	H	α_1	α_2
MEA	e	0	2.08979	0	2.65064	0	2.10763	0	0
	H	—	0	2.65064	0	2.10763	0	0	0
	e*	—	—	0	2.32894	0	2.22626	1.97978	1.96999
	H*	—	—	—	0	2.22626	0	0	0
H ₂ O	e	—	—	—	—	0	2.10822	0	0
	H	—	—	—	—	—	0	0	0
CO ₂	α_1	—	—	—	—	—	—	0	0
	α_2	—	—	—	—	—	—	0	0

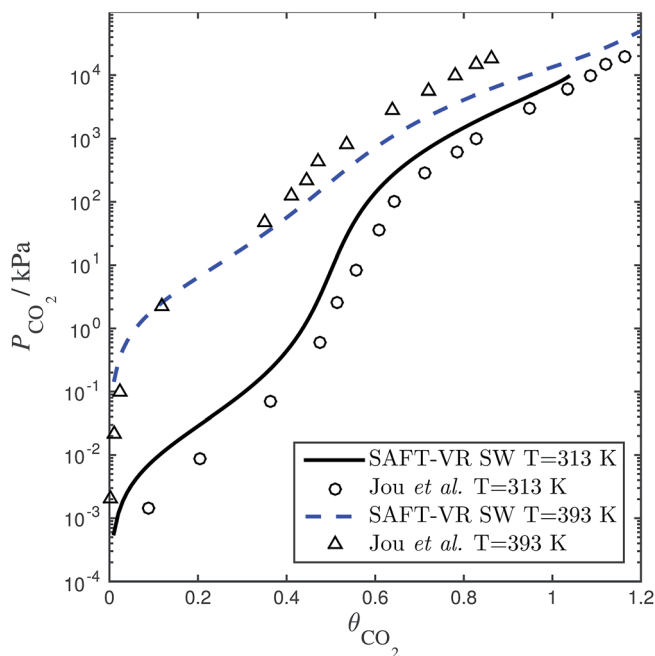


Fig. 4 Solubility of CO₂ in a 30 wt% aqueous solution of MEA expressed as the CO₂ loading (defined as the number of moles of CO₂ per mole of MEA in the liquid solvent) as a function of the partial pressure. The SAFT-VR model predictions (dashed and continuous curves) using parameters from Rodríguez *et al.*⁸⁴ are compared with the experimental data (triangles and circles) from Jou *et al.*¹⁵⁹

correlations, heat-transfer correlations, and diffusion correlations. We list explicitly all of the model equations to clarify the radically different approach followed here. One important model parameter τ is introduced to account for a scaling of the CO₂ diffusivity. This parameter will be estimated from experimental data. In the following equations, the subscript i refers to the i^{th} component, j to the j^{th} stage, c is the total number of components, and N_s is the number of stages in the column. The number of stages is fixed to 50 which is amply

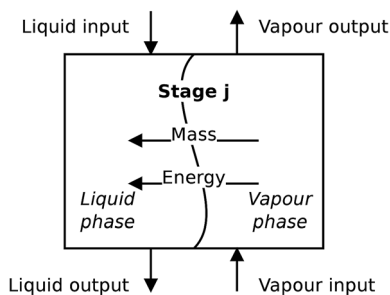


Fig. 5 A schematic of a non-equilibrium stage. This stage represents a section of packing in a packed column.



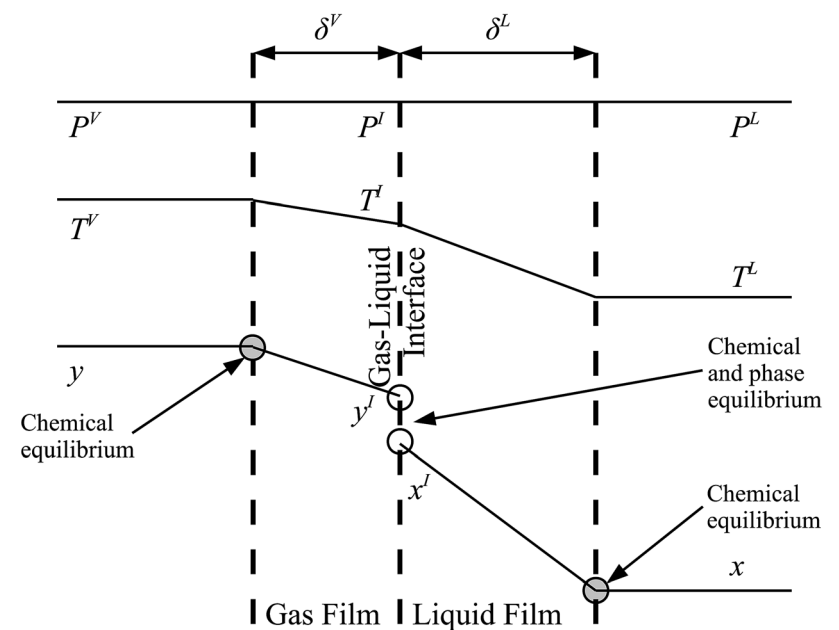


Fig. 6 A schematic of the two-film model of a column stage. δ^V and δ^L represent the thickness of the gas and liquid films, respectively. P^V , P^I , and P^L are the pressure in the bulk vapour phase, at the gas–liquid interface, and in the bulk liquid, respectively. As can be seen an isobaric profile is assumed. The temperatures of the bulk vapour phase, at the gas–liquid interface, and of the bulk liquid are denoted by T^V , T^I , and T^L , respectively. Finally, y_i and x_i are the mole fractions of component i in the bulk vapour and liquid phases, respectively, and y_i^I and x_i^I are the mole fractions of component i at the vapour–liquid interface in the vapour and the liquid phases, respectively.

sufficient to assure numerical convergence without significantly impacting the computational time.

2.3.1 Heat and mass balances. The liquid and vapour phase mole balances for component i on stage j are given by

$$L_{j-1}x_{i,j-1} + N_{i,j}^L = L_jx_{i,j}, \quad i = 1, 2, \dots, c; j = 1, 2, \dots, N_s \quad (4)$$

$$V_{j+1}y_{i,j+1} - N_{i,j}^V = V_jy_{i,j}, \quad i = 1, 2, \dots, c; j = 1, 2, \dots, N_s, \quad (5)$$

where L_j and V_j (mol s^{-1}) are the total liquid and vapour molar flow rate leaving stage j respectively, $x_{i,j}$ and $y_{i,j}$ are the bulk liquid and vapour mole fractions of component i on stage j , $N_{i,j}^L$ (mol s^{-1}) is the net gain of species i in the liquid phase due to interphase transport for stage j , and $N_{i,j}^V$ (mol s^{-1}) is the net loss of species i in the vapour phase due to interphase transport for stage j .

The mole fractions in the streams leaving each stage must sum to unity:

$$\sum_{i=1}^c x_{i,j} = 1 \quad \text{and} \quad \sum_{i=1}^c y_{i,j} = 1 \quad j = 1, 2, \dots, N_s. \quad (6)$$

The energy balances for the liquid and vapour phases respectively are



$$L_{j-1}H_{j-1}^L(T_{j-1}^L, V_{j-1}^L, \underline{x}_{j-1}) + E_j^L = L_jH_j^L(T_j^L, V_j^L, \underline{x}_j), j = 1, 2, \dots, N_s, \quad (7)$$

$$V_{j+1}H_{j+1}^V(T_{j+1}^V, V_{j+1}^V, \underline{y}_{j+1}) - E_j^V = V_jH_j^V(T_j^V, V_j^V, \underline{y}_j), j = 1, 2, \dots, N_s, \quad (8)$$

where H_j^L and H_j^V (J mol⁻¹) are the molar enthalpies of the liquid and gas phases of stage j , respectively. E_j^L (W) is the net gain of energy of the liquid phase through the interface at stage j , and E_j^V (W) is the net loss of energy from the vapour phase through the interface at stage j . The molar enthalpies H_j^L and H_j^V are determined from SAFT-VR as functions of the temperature of the bulk phases, T_j^L and T_j^V (K), the molar volumes of the phases, V_j^L and V_j^V (m³ mol⁻¹), and the composition (vectors) of the phases, $\underline{x}_j$ and $\underline{y}_j$.

There is no accumulation of mass or energy at the vapour-liquid interface:

$$N_{i,j}^L = N_{i,j}^V, i = 1, 2, \dots, c; j = 1, 2, \dots, N_s, \quad (9)$$

$$E_j^L = E_j^V, i = 1, 2, \dots, c; j = 1, 2, \dots, N_s. \quad (10)$$

2.3.2 Rate equations: mass transfer. According to the film model for mass transfer, we define the mass-transfer rates as follows:

$$N_{i,j}^L = k_{i,j}^L a_{T,j} (C_{i,j}^{L,L} - C_{i,j}^{L,V}), i = 1, 2, \dots, c; j = 1, 2, \dots, N_s, \quad (11)$$

$$N_{i,j}^V = k_{i,j}^V a_{T,j} (C_{i,j}^{V,V} - C_{i,j}^{V,L}), i = 1, 2, \dots, c; j = 1, 2, \dots, N_s, \quad (12)$$

where $a_{T,j}$ (m²) is the total interfacial area on stage j available for heat or mass transfer, $k_{i,j}^L$ (m s⁻¹) is the liquid-phase mass-transfer coefficient for component i of stage j , $k_{i,j}^V$ (m s⁻¹) is the vapour-phase mass-transfer coefficient for component i of stage j , $C_{i,j}^{L,L}$ and $C_{i,j}^{V,V}$ (mol m⁻³) are the concentrations of component i in the bulk liquid and vapour phases of stage j , and $C_{i,j}^{L,V}$ and $C_{i,j}^{V,L}$ (mol m⁻³) are the concentrations of component i at the vapour-liquid interface in the liquid and vapour phases of stage j .

The total area for heat and mass transfer is given by the product of the effective specific area and the stage volume:

$$a_{T,j} = a'_j A_{\text{section}} D_z, j = 1, 2, \dots, N_s, \quad (13)$$

where a'_j (m² m⁻³) is the interfacial area density on stage j , A_{section} (m²) is the cross sectional area of the column, and D_z (m) is the stage height given by h_{packing}/N_s , with h_{packing} (m) as the total packing height.

2.3.3 Rate equations: heat transfer. The energy fluxes in the liquid and gas phases at stage j are given by the sum of the conductive and diffusive heat fluxes:

$$E_j^L = Q_{\text{cond},j}^L + Q_{\text{diff},j}^L, j = 1, 2, \dots, N_s, \quad (14)$$

$$E_j^V = Q_{\text{cond},j}^V + Q_{\text{diff},j}^V, j = 1, 2, \dots, N_s, \quad (15)$$

where $Q_{\text{cond},j}^L$ and $Q_{\text{diff},j}^L$ (W) are the conductive and diffusive heat fluxes at the interface into the liquid phase on stage j , and $Q_{\text{cond},j}^V$ and $Q_{\text{diff},j}^V$ (W) are the conductive and diffusive heat fluxes at the interface out of the vapour phase on stage j .



The conductive heat-transfer rate into the liquid phase and out of the vapour phase on stage j are driven by temperature gradients between the bulk phases and the interface, and are given by

$$Q_{\text{cond},j}^{\text{L}} = h_{\text{T},j}^{\text{L}} a_{\text{T},j} (T_j^{\text{I}} - T_j^{\text{I}}), j = 1, 2, \dots, N_s, \quad (16)$$

$$Q_{\text{cond},j}^{\text{V}} = h_{\text{T},j}^{\text{V}} a_{\text{T},j} (T_j^{\text{V}} - T_j^{\text{I}}), j = 1, 2, \dots, N_s, \quad (17)$$

where $h_{\text{T},j}^{\text{L}}$ and $h_{\text{T},j}^{\text{V}}$ ($\text{W m}^{-2} \text{K}^{-1}$) are the heat-transfer coefficients in the liquid and gas phases on stage j , respectively, and T_j^{I} (K) is the temperature at the vapour-liquid interface on stage j .

The diffusive heat fluxes account for the variation in enthalpy in the liquid and vapour streams associated with the transfer of mass from one phase to another. This is obtained based on the individual component fluxes and the corresponding enthalpies of the pure components:

$$Q_{\text{diff},j}^{\text{L}} = \sum_{i=1}^c N_{ij}^{\text{L}} H(T_j^{\text{L}}, V_{ij}^{*\text{L}}, \underline{z}_i) \quad j = 1, 2, \dots, N_s, \quad (18)$$

$$Q_{\text{diff},j}^{\text{V}} = \sum_{i=1}^c N_{ij}^{\text{V}} H(T_j^{\text{V}}, V_{ij}^{*\text{V}}, \underline{z}_i) \quad j = 1, 2, \dots, N_s, \quad (19)$$

where $H(T_j^{\text{L}}, V_{ij}^{*\text{L}}, \underline{z}_i)$ and $H(T_j^{\text{V}}, V_{ij}^{*\text{V}}, \underline{z}_i)$ (J mol^{-1}) are the molar enthalpies of pure component i in the bulk liquid and vapour phase respectively, as calculated with SAFT-VR SW. The variable $\underline{z}_i$ denotes the molar composition vector of a stream consisting of pure component i defined by $z_{k,i} = 1$ if $i = k$ and $z_{k,i} = 0$ otherwise. $V_{ij}^{*\text{L}}$ and $V_{ij}^{*\text{V}}$ ($\text{m}^3 \text{mol}^{-1}$) are the molar volumes of pure component i at stage j in the bulk liquid and vapour phases respectively, obtained from

$$P_j^{\text{L}} = P(T_j^{\text{L}}, V_{ij}^{*\text{L}}, \underline{z}_i), i = 1, 2, \dots, c; j = 1, 2, \dots, N_s, \quad (20)$$

$$P_j^{\text{V}} = P(T_j^{\text{V}}, V_{ij}^{*\text{V}}, \underline{z}_i), i = 1, 2, \dots, c; j = 1, 2, \dots, N_s, \quad (21)$$

where P_j^{L} and P_j^{V} (MPa) are the pressure of the bulk liquid and vapour phases at stage j respectively and $P(T_j^{\text{L}}, V_{ij}^{*\text{L}}, \underline{z}_i)$ and $P(T_j^{\text{V}}, V_{ij}^{*\text{V}}, \underline{z}_i)$ represent evaluations of the pressure using the SAFT-VR SW EoS.

When equating the vapour and liquid fluxes, one obtains an expression which includes the difference between the enthalpies in the vapour phase and in the liquid phase for a pure component i ; this is the enthalpy of vaporization of this component. As we will see, the enthalpy of vaporization of water plays a significant role in the process. We assume that the contribution of N_2 to the change of enthalpy between the two phases is negligible as its mass transfer is very small, and that the enthalpy of vaporization of CO_2 is negligible as the operating conditions are close to or above its critical point. In the case where only one phase is stable for a pure component at the conditions of interest, as is typically the case for CO_2 , one can access the hypothetical phase information by providing an adequate initial guess for the molar volume to the SAFT-VR SW EoS.

2.3.4 Equilibrium relations. Both chemical and phase equilibria are assumed to prevail at the interface so that the conditions of equality of pressure, temperature, and chemical potential must be satisfied:



$$\mu(T_j^I, V_j^{I,V}, \underline{y}_j^I) = \mu(T_j^I, V_j^{I,L}, \underline{x}_j^I) = \mu_{ij}^{I,L} = \mu_{ij}^{I,V}, i = 1, 2, \dots, c; j = 1, 2, \dots, N_s, \quad (22)$$

$$P(T_j^I, V_j^{I,V}, \underline{y}_j^I) = P(T_j^I, V_j^{I,L}, \underline{x}_j^I) = P_j^I, j = 1, 2, \dots, N_s, \quad (23)$$

where $\mu_{ij}^{I,L}$ and $\mu_{ij}^{I,V}$ (J mol^{-1}) are the chemical potentials of component i on stage j in the liquid and vapour phases respectively, $V_j^{I,L}$ and $V_j^{I,V}$ ($\text{m}^3 \text{mol}^{-1}$) are the molar volumes of the liquid phase and the vapour phases at the vapour-liquid interface (I) on stage j , and P_j^I (MPa) is the pressure at the liquid-vapour interface on stage j . The chemical potentials and the molar volume are determined using the SAFT-VR SW EoS.

The mole fractions at the interface must sum to unity:

$$\sum_{i=1}^c x_{ij}^I = 1 \quad \text{and} \quad \sum_{i=1}^c y_{ij}^I = 1 \quad j = 1, 2, \dots, N_s. \quad (24)$$

Each stage is assumed to be in mechanical equilibrium, so that

$$P_j^V = P_j^L = P_j^I, j = 2, \dots, N_s, \quad (25)$$

where

$$P_j^V = P(T_j^V, V_j^V, \underline{y}_j^V), j = 1, 2, \dots, N_s, \quad (26)$$

$$P_j^L = P(T_j^L, V_j^L, \underline{x}_j^L), j = 1, 2, \dots, N_s. \quad (27)$$

Finally, the column pressure drop is assumed to be negligible.

$$P_j^V = P_{j+1}^V, j = 1, 2, \dots, N_s. \quad (28)$$

All pressure terms present in the model are thus equated to the pressure of the gas inlet stream, $P_{N_s+1}^V$. This assumption can easily be removed at a later stage of the model development. Preliminary results have indicated that the pressure drop has a negligible effect on the process used for the validation of our model.

The equations presented in the remainder of this section apply to each stage. The subscript j has been omitted for the purpose of clarity.

2.3.5 Mass-transfer correlations. The interfacial area, the liquid-phase mass-transfer coefficient, and the gas-phase mass-transfer coefficient are obtained using the correlations from Onda *et al.*^{57,58} Other correlations are available, the most commonly used being the ones developed by Rocha *et al.*⁶⁰ and Billet and Schultes.⁶¹ Faramarzi *et al.*⁴⁸ compared the performance of the correlations and found that all three provide a good description; they expressed a slight preference for the correlations of Rocha *et al.*⁶⁰ but could not draw a general conclusion applicable to all operating conditions. The correlations of Rocha *et al.*⁶⁰ apply to structured packing and have to be adapted for use in random packing, whereas those developed by Onda *et al.*^{57,58} have been developed specifically using random Berl saddle packing, the same as the one employed in the pilot plant by Tontiwachuthikul *et al.*¹³ This last set of correlations is sometimes reparameterized when used in commercial software such as Aspen or ProTreat. The correlations developed by Onda *et al.*,^{57,58} as reported by Treybal,¹³⁹ are implemented in our model of the absorber column. The liquid-phase mass-transfer coefficient for component i is given by



$$k_i^L = 0.0051 \left(\frac{\eta^L g}{\rho^L} \right)^{1/3} (\text{Re}^L)^{2/3} (\text{Sc}_i^L)^{-1/2} (a_p L_p)^{0.4} \quad i = 1, 2, \dots, c, \quad (29)$$

where ρ^L (kg m^{-3}) is the density of the liquid phase, η^L ($\text{kg m}^{-1} \text{s}^{-1}$) is the dynamic viscosity of the liquid phase, g (m s^{-2}) is the gravitational acceleration, Re^L is the liquid-phase Reynolds number based on the interfacial area, Sc_i^L is the Schmidt number of component i in the liquid phase, a_p ($\text{m}^2 \text{m}^{-3}$) is the specific surface area of the packing, and L_p (m) is the nominal packing size.

The Reynolds number and the Schmidt number for component i in the liquid phase are

$$\text{Re}^L = \frac{\rho^L u^L}{a' \eta^L}, \quad (30)$$

$$\text{Sc}_i^L = \left(\frac{\eta^L}{\rho^L D_i^L} \right) \quad i = 1, 2, \dots, c, \quad (31)$$

where D_i^L ($\text{m}^2 \text{s}^{-1}$) is the diffusion coefficient of component i in the liquid phase, u^L (m s^{-1}) is the liquid velocity, and a' is the interfacial area density defined in eqn (35).

The vapour-phase mass-transfer coefficient is obtained from

$$k_i^V = 2a_p D_i^V \text{Re}^{V0.7} \text{Sc}_i^{V1/3} (a_p L_p)^{-2}, \quad i = 1, 2, \dots, c, \quad (32)$$

where D_i^V ($\text{m}^2 \text{s}^{-1}$) is the diffusivity of component i in the vapour-phase, Re^V is the vapour-phase Reynolds number, and Sc_i^V is the Schmidt number of component i in the vapour phase.

The Reynolds number and the Schmidt number for the vapour phase are

$$\text{Re}^V = \frac{\rho^V u^V}{a_p \eta^V}, \quad (33)$$

$$\text{Sc}_i^V = \left(\frac{\eta^V}{\rho^V D_i^V} \right) \quad i = 1, 2, \dots, c, \quad (34)$$

where ρ^V (kg m^{-3}) is the density of the vapour phase, η^V ($\text{kg m}^{-1} \text{s}^{-1}$) is the dynamic viscosity of the vapour phase, and u^V (m s^{-1}) is the vapour velocity.

The interfacial area density a' is obtained from^{57,58}

$$\frac{a'}{a_p} = 1 - \exp \left[-1.45 \left(\frac{\sigma_c}{\sigma} \right)^{0.75} (\text{Re}^L)^{0.1} (\text{Fr}^L)^{-0.05} (\text{We}^L)^{0.2} \right], \quad (35)$$

where σ_c (N m^{-1}) is the critical surface tension of the packing material, σ (N m^{-1}) is the vapour-liquid surface tension, Re^L is the liquid-phase Reynolds number based on the specific surface area, Fr^L is the liquid-phase Froude number, and We^L is the liquid-phase Weber number. The interfacial area density, together with the three dimensionless numbers used in eqn (35), are properties of the mixture, not properties of individual components.

The expressions for the dimensionless numbers are

$$\text{Re}^L = \frac{\rho^L u^L}{\eta^L a_p}, \quad (36)$$



$$\text{Fr}^{\text{L}} = \frac{a_{\text{p}} u^{\text{L}^2}}{g}, \quad (37)$$

$$\text{We}^{\text{L}} = \frac{\rho^{\text{L}} u^{\text{L}^2}}{a_{\text{p}} \sigma}. \quad (38)$$

The velocities of each phase are

$$u^{\text{L}} = \frac{\dot{v}^{\text{L}}}{A_{\text{section}}}, \quad (39)$$

$$u^{\text{V}} = \frac{\dot{v}^{\text{V}}}{A_{\text{section}}}, \quad (40)$$

where $\dot{v}^{\text{L}}$ and $\dot{v}^{\text{V}}$ ($\text{m}^3 \text{s}^{-1}$) are the volumetric flow rates of the liquid and gas phase, respectively.

2.3.6 Heat-transfer correlations. For the gas and liquid heat-transfer coefficients we use the correlations presented in Treybal's *Mass Transfer Operations*.¹³⁹ Once again these expressions apply to each stage. The expression for the liquid-phase heat-transfer coefficient is

$$\frac{h_{\text{T}}^{\text{L}} d_{\text{s}}}{\lambda_{\text{T}}^{\text{L}}} = 25.1 \left(\frac{d_{\text{s}} L_{\text{spec}}}{\eta^{\text{L}}} \right)^{0.45} (\text{Pr}^{\text{L}})^{0.45}, \quad (41)$$

where h_{T}^{L} ($\text{W m}^{-2} \text{K}^{-1}$) is the liquid-phase heat-transfer coefficient, d_{s} (m) is the diameter of a sphere of the same surface area as a single packing particle (not the same as a_{p}), $\lambda_{\text{T}}^{\text{L}}$ ($\text{W m}^{-1} \text{K}^{-1}$) is the liquid thermal conductivity, L_{spec} (kg s^{-1}) is the specific liquid flow rate, and Pr^{L} is the liquid-phase Prandtl number. The latter is obtained from the following expression:

$$\text{Pr}^{\text{L}} = \frac{C_{\text{p}}^{\text{L}} \eta^{\text{L}}}{\lambda_{\text{T}}^{\text{L}}}, \quad (42)$$

where C_{p}^{L} ($\text{J kg}^{-1} \text{K}^{-1}$) is the specific isobaric heat capacity of the liquid phase. The vapour-phase heat-transfer coefficient is given by

$$\frac{h_{\text{T}}^{\text{V}}}{C_{\text{p}}^{\text{V}} V_{\text{spec}}} \text{Pr}^{\text{V}^{2/3}} = 1.195 \left(\frac{d_{\text{s}} V_{\text{spec}}}{\eta^{\text{V}} (1 - \varepsilon_{\text{L}_0})} \right)^{-0.36}, \quad (43)$$

where h_{T}^{V} ($\text{W m}^{-2} \text{K}^{-1}$) is the vapour-phase heat-transfer coefficient, C_{p}^{V} ($\text{J kg}^{-1} \text{K}^{-1}$) is the specific isobaric heat capacity of the vapour phase, Pr^{V} is the Prandtl number for the gas phase, V_{spec} (kg s^{-1}) is the specific vapour flow rate, and ε_{L_0} is the operating void space in the packing (assumed to be equal to the void fraction ε). The Prandtl number for the gas phase is obtained from

$$\text{Pr}^{\text{V}} = \frac{C_{\text{p}}^{\text{V}} \eta^{\text{V}}}{\lambda_{\text{T}}^{\text{V}}}, \quad (44)$$

where $\lambda_{\text{T}}^{\text{V}}$ ($\text{W m}^{-1} \text{K}^{-1}$) is the vapour thermal conductivity.

2.3.7 Vapour-phase diffusion coefficient. The vapour-phase diffusion coefficient used in eqn (34) is obtained from the expression of Fuller and



co-workers^{140–142} as reported by Poling *et al.*⁷⁷ It is assumed that the components are diffusing through air:

$$D_i^V = 10^{-4} \frac{0.00143 T^{V1.75}}{10 P^V M_{i-\text{air}}^{1/2} \left[\Sigma_{V_i}^{1/3} + \Sigma_{V_{\text{air}}}^{1/3} \right]} \quad i = 1, 2, \dots, c, \quad (45)$$

where Σ_v (Å) is the atomic diffusion volume, and $M_{i-\text{air}}$ is given by

$$M_{i-\text{air}} = 2[(1/M_{\text{air}}) + (1/M_i)]^{-1}, \quad i = 1, 2, \dots, c, \quad (46)$$

where M_i (g mol⁻¹) is the molar mass of component i .

2.3.8 Liquid-phase diffusion coefficient. The liquid-phase binary diffusion coefficients are described with different correlations. The mutual diffusion coefficients of CO₂ at very low concentration in pure MEA and pure H₂O are derived from the Wilke–Chang correlation:^{77,143}

$$D_{\text{CO}_2,k}^\circ = \frac{7.4 \times 10^{-8} (\phi_k M_k)^{1/2} T^L}{\eta_k^L \left(V_{\text{CO}_2}^m \right)^{0.6}} \quad k = \text{H}_2\text{O}, \text{MEA}, \quad (47)$$

where ϕ_k is the “association factor” of solvent k , and $V_{\text{CO}_2}^m$ (cm³ mol⁻¹) is the molar volume of CO₂ at its normal boiling temperature.

These mutual diffusion coefficients are used in the correlation of Takahashi *et al.*¹⁴⁴ to determine the diffusivity of CO₂ in a liquid mixture of H₂O and MEA:

$$D_{\text{CO}_2}^L = \tau \times 10^{-4} \left(\frac{V^L}{\eta^L} \right)^{1/3} \sum_{k=\text{H}_2\text{O}, \text{MEA}} x_k D_{\text{CO}_2,k}^\circ \left(\frac{\eta_k^L}{V_k^{*L}} \right)^{1/3}, \quad (48)$$

where we have introduced the scaling prefactor τ to the correlation. We refer to eqn (48) as the effective diffusivity (ED) correlation. The presence of this scaling factor provides an effective approach to modelling several effects that are not accounted for due to the use of a chemical approach and the consequent implicit treatment of the reaction products: the acceleration of mass-transfer due to the depletion of CO₂ *via* chemical reactions;⁴⁰ the reduction in mass transfer due to the ion pairs formed and their electrostatic interaction with other species;⁴⁰ and the reduction in the mass transfer due to CO₂ being present in larger aggregated products (bicarbonate and carbonate species). The latter effect was observed by Han *et al.*¹⁴⁵ in a molecular-dynamics study of the transport properties of CO₂ in MEA, indicating a significant decrease in the diffusion coefficient with increasing CO₂ loading. The value of the parameter τ and its impact on model predictions are discussed in Section 3. Finally, the Perkins and Geankoplis¹⁴⁶ method is used to calculate the diffusivity of H₂O, MEA, and N₂ in the liquid phase:

$$D_i^L = 10^{-4} \times \frac{7.4 \times 10^{-8} (\phi M)_i^{1/2} T}{1000 \eta^L (V_i^m)^{0.6}} \quad i = \text{H}_2\text{O}, \text{MEA}, \text{N}_2, \quad (49)$$

where V_i^m (cm³ mol⁻¹) is the molar volume of component i at its normal boiling temperature, and the term $(\phi M)_i$ (g mol⁻¹) represents the “association factor” of component i in the liquid mixture which is derived from

$$(\phi M)_i = \sum_{j=1, j \neq i}^n x_j \phi_j M_j \quad i = \text{H}_2\text{O}, \text{MEA}, \text{N}_2. \quad (50)$$



Table 5 Correlations used in the process model of the absorber

Physical property	Component	Source
Liquid-phase viscosity	H ₂ O	Westmeier ¹⁶⁰
	MEA	Leibush and Shorina ¹⁶¹
	Mixture	Maham <i>et al.</i> ¹⁶²
Vapour-phase viscosity		Kestin <i>et al.</i> ¹⁶³
Vapour–liquid surface tension		Vazquez <i>et al.</i> ¹⁵²
Heat conductivity		Haynes ¹⁶⁴
Liquid heat capacity	H ₂ O	Laliberte ¹⁶⁵
	MEA	Mundhwa and Henni ¹⁶⁶
Vapour heat capacity	H ₂ O	Riddick <i>et al.</i> ¹⁶⁷
	N ₂	Vargaftik ¹⁶⁸
	CO ₂	Bender <i>et al.</i> ¹⁶⁹
	CO ₂	Takahashi <i>et al.</i> ¹⁴⁴
Liquid-phase diffusion coefficient	H ₂ O, MEA, N ₂	Perkins and Geankoplis ¹⁴⁶
Vapour-phase diffusion coefficient	CO ₂ , H ₂ O, MEA, N ₂	Poling <i>et al.</i> ⁷⁷

The other correlations of experimental data used in our model are listed in Table 5.

3 Results

The model described in Section 2.2 is implemented in gProms¹⁴⁷ and an in-house implementation of the SAFT-VR SW EoS¹⁴⁸ is accessed *via* a Foreign Object Interface.

In this section, we investigate the predictive capabilities of the proposed model by thorough comparisons with the pilot-plant data obtained by Tontiwachwuthikul *et al.*¹³ The inputs required to model these data and relevant assumptions are discussed in Section 3.1. As discussed in Section 3.2, we first assume that there are no pilot-plant data available for the solvent in question and analyze the suitability of the model for the provision of a best-case analysis of the process performance. It is then assumed that only one pilot-plant run is available and the corresponding concentration profiles are used to estimate the single parameter τ , related to mass-transfer limitations, as summarized in Section 3.3. The transferability of this parameter is assessed by comparing model predictions against data for other pilot-plant runs. Finally, in Section 3.4, the sensitivity of the model to several parameters is explored, providing insights into the behaviour of CO₂ absorption columns.

3.1 Model inputs

The absorption column studied by Tontiwachwuthikul *et al.*¹³ has an internal diameter of 0.1 m and a total packed height of 6.6 m; care has to be taken not to mistake the total column height of 7.2 m with the packed height, which is the relevant dimension in modelling the absorber. The absorption column internals are randomly packed 12.7 mm ceramic Berl saddles. The gas inlet stream is assumed to be free of MEA and the liquid solvent inlet stream free of N₂. The input values used are listed in Table 6 and the characteristics of the column and the packing are listed in Table 7. Some of the data needed for the simulation of



Table 6 Inputs to absorber column model for Runs T13 to T22 (ref. 13)

Input variable	T13	T14	T15	T16	T17	T18	T19	T20	T21	T22
Inlet gas flow (mol m ⁻² s ⁻¹)	17.54	17.54	18.4	17.51	17.54	18.29	16.72	18.32	13.72	18.3
Inlet gas pressure (MPa)	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Inlet gas temperature (K)	292.15	292.15	292.15	292.15	293.15	293.15	292.15	292.15	292.15	292.15
Inlet gas CO ₂ mole fraction	0.153	0.156	0.195	0.155	0.156	0.191	0.115	0.192	0.191	0.191
Inlet gas H ₂ O mass fraction	0.0058	0.0058	0.0058	0.0058	0.0058	0.0058	0.0058	0.0058	0.0058	0.0058
Inlet liquid flow (m ³ m ⁻² h ⁻¹)	13.5	13.5	13.5	9.5	13.5	9.5	13.5	9.5	9.5	9.5
Inlet liquid temperature (K)	292.15	292.15	292.15	292.15	293.15	293.15	292.15	292.15	292.15	292.15
Inlet MEA concentration (kmol m ⁻³)	2.00	2.00	2.03	2.08	3.08	2.00	2.00	2.55	2.00	3.00
Lean loading θ_{CO_2}	10 ⁻⁴	0.118	10 ⁻⁴	10 ⁻⁴	0.237	10 ⁻⁴	10 ⁻⁴	10 ⁻⁴	10 ⁻⁴	10 ⁻⁴

the process were not explicitly reported by Tontiwachwuthikul *et al.*¹³ in particular, the temperature of the flue gas, and the concentration of water in the flue gas. Close inspection of an earlier paper detailing the experimental apparatus¹⁴⁹ reveals that the temperature of the flue gas is controlled to be that of the lean solvent with a thermostatic bath. The inlet vapour stream is composed of ambient air and bottled CO₂. Typical values of the relative humidity of air (defined as the mole fraction of water vapour divided by the mole fraction of air saturated with water at the same temperature and pressure) lie between 20% to 70%, which, at 20 °C and 1 bar, corresponds to a mass fraction of H₂O $\omega_{\text{H}_2\text{O}}^{\text{V}}$ in the flue gas of between 0.003 and 0.01.¹⁵⁰ A sensitivity analysis of the extent of humidity indicates that the variation of $\omega_{\text{H}_2\text{O}}^{\text{V}}$ between 0 and 0.072 (corresponding to a mole fraction 0.12) has a negligible impact on the temperature and composition profiles. The effect of the amount of water in the flue gas is discussed in more detail in Section 3.3.2. For our working model, the nominal amount of water in the inlet flue gas is fixed to $\omega_{\text{H}_2\text{O}}^{\text{V}} = 0.0058$, corresponding to 50% humidity at 20 °C and 1 bar.

Table 7 Characteristics of the column and the packing

Name	Value	Source
A_{section} (m ²)	0.00785	Tontiwachwuthikul <i>et al.</i> ¹³
a_{p} (m ² m ⁻³)	466	Treybal ¹³⁹
σ_{c} (N m ⁻¹)	61	Ref. 170 pg. 18–34, Tables 18–11
L_{p} (m)	0.0127	Tontiwachwuthikul <i>et al.</i> ¹³
ε	0.63	Ref. 139 pg. 198, Table 6.3
D_{z} (m)	0.132	Tontiwachwuthikul <i>et al.</i> ¹³
d_{s} (m)	0.31622	Ref. 139 pg. 206, Table 6.5



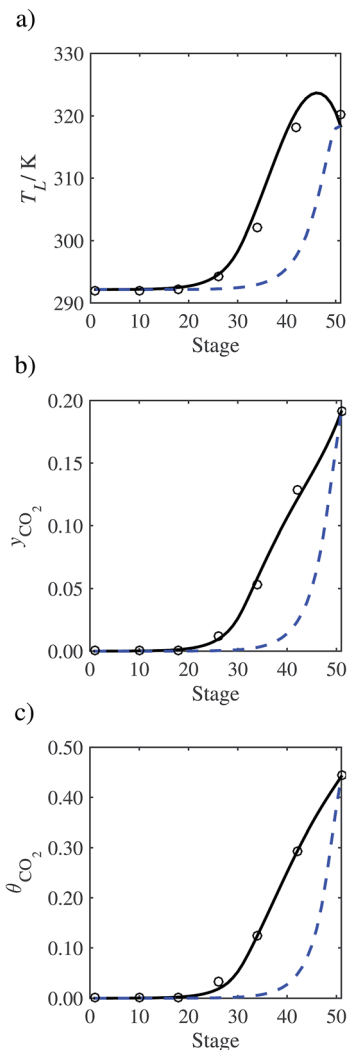


Fig. 7 Comparison of the pilot-plant data (circles)¹³ and the result of our model (curves) for Run T22 for the absorption of CO₂ in an 18 wt% solution of MEA. The dashed curves represent the results obtained without scaling the CO₂ diffusivity ($\tau = 1$) and the continuous curves the results obtained with the ED correlation (eqn (48)) by scaling the CO₂ diffusivity in the liquid phase to 4.1% of its original value ($\tau = 0.041$). (a) Temperature profile for the liquid phase, (b) gas-phase CO₂ concentration profile, and (c) liquid-phase CO₂ loading (defined as the number of moles of CO₂ per mole of MEA in the liquid phase). Stage 50 corresponds to the bottom of the column.

3.2 Scenario 1

In the absence of pilot-plant data, the scaling factor τ for the diffusion coefficient of CO₂, *cf.* eqn (48), is set to 1. The predictions with $\tau = 1$ are presented in Fig. 7 and Fig. 9–17 as dashed curves. In all but one case (Run T18, Fig. 14), the rate of absorption of CO₂ throughout the column is found to be over-predicted as the estimates for the gas-phase CO₂ concentration profiles y_{CO_2} all fall under the



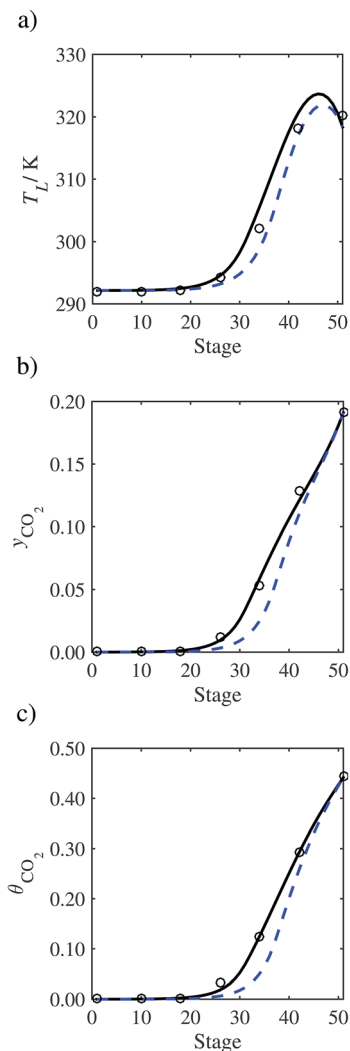


Fig. 8 Sensitivity analysis of the mass transfer correlation on the profiles predicted with our model compared with the experimental pilot-plant data of Run T22¹³: mass-transfer correlations from Onda *et al.*^{57,58} (continuous curve); mass-transfer correlations from Rocha *et al.*^{60,151} (a) Temperature profile for the liquid phase, (b) gas phase CO_2 concentration profile, and (c) liquid phase CO_2 loading. Stage 50 corresponds to the bottom of the column.

measured values. With the exception of Run T18, the temperature profiles T_L and the liquid-phase CO_2 loading profiles θ_{CO_2} are similarly under-predicted. Complete absorption occurs at approximately stage 30, whilst an analysis of the pilot-plant data suggests that this occurs between stages 10 to 20. The size of the column required to achieve maximum absorption is therefore underestimated based on these predictions alone.

In Run T18 (Fig. 14), there are two competing effects: the overestimation of the absorption of CO_2 tends to increase the temperature of the liquid phase, which in



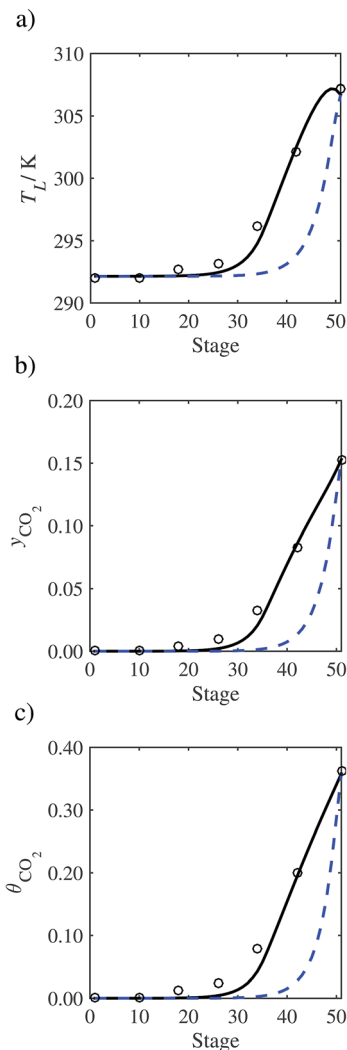


Fig. 9 Comparison of the pilot-plant data (circles)¹³ and the result of our model (curves) for Run T13. The dashed curves represent the results obtained without scaling the CO_2 diffusivity ($\tau = 1$) and the continuous curves the results obtained with the ED correlation (eqn (48)) by scaling the CO_2 diffusivity in the liquid phase to 4.1% of its original value ($\tau = 0.041$). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading. Stage 50 corresponds to the bottom of the column.

turn tends to decrease the extent of absorption of CO_2 in the liquid. Since this is the only run in which the CO_2 in the inlet gas is not completely absorbed within the column, the cooling effect of the liquid feed is not as apparent as in other runs. Given the relatively low recovery of CO_2 and the fact that equilibrium is not reached at the bottom of the column, our findings indicate that the model is best used as an indicator of process performance by modelling a sufficiently large absorber to achieve equilibrium.



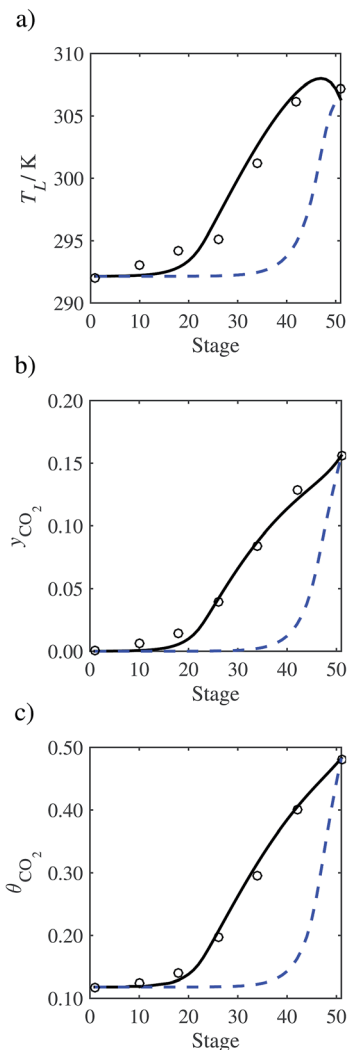


Fig. 10 Comparison of the pilot-plant data (circles)¹³ and the result of our model (curves) for Run T14. The dashed curves represent the results obtained without scaling the CO_2 diffusivity ($\tau = 1$) and the continuous curves the results obtained with the ED correlation (eqn (48)) by scaling the CO_2 diffusivity in the liquid phase to 4.1% of its original value ($\tau = 0.041$). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading. Stage 50 corresponds to the bottom of the column.

In general, an optimistic prediction of process performance is thus obtained with the model when no pilot-plant data are used ($\tau = 1$). Indeed, although the depletion of CO_2 through chemical reactions has an accelerating effect on mass transfer, the reduced diffusivities of the product species lead to an overall reduction in mass transfer rates. This suggests that the proposed predictive model could be used to obtain a preliminary assessment of novel solvents in the absence of pilot-plant data: if their best-case ($\tau = 1$) performance is found to be



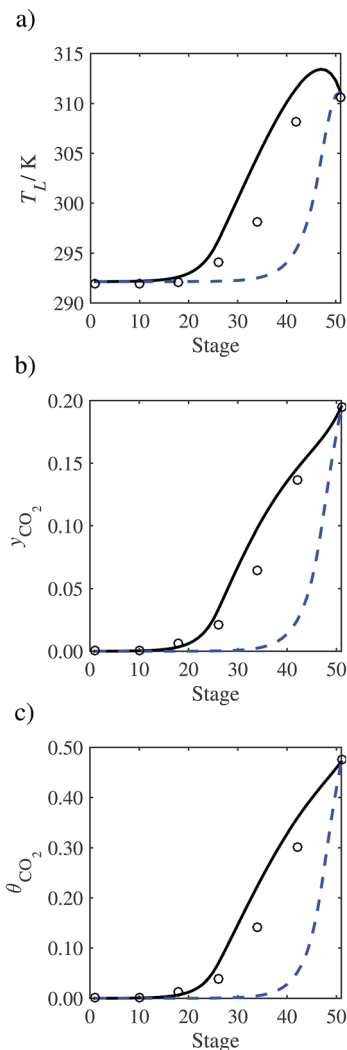


Fig. 11 Comparison of the pilot-plant data (circles)¹³ and the result of our model (curves) for Run T15. The dashed curves represent the results obtained without scaling the CO_2 diffusivity ($\tau = 1$) and the continuous curves the results obtained with the ED correlation (eqn (48)) by scaling the CO_2 diffusivity in the liquid phase to 4.1% of its original value ($\tau = 0.041$). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading. Stage 50 corresponds to the bottom of the column.

significantly less than that of MEA or another suitable benchmark, these solvents would be eliminated from further consideration without undertaking an experimental programme.

3.3 Scenario 2

3.3.1 Estimation of the scaling factor. In order to investigate whether the proposed model can be used to provide quantitative agreement with the



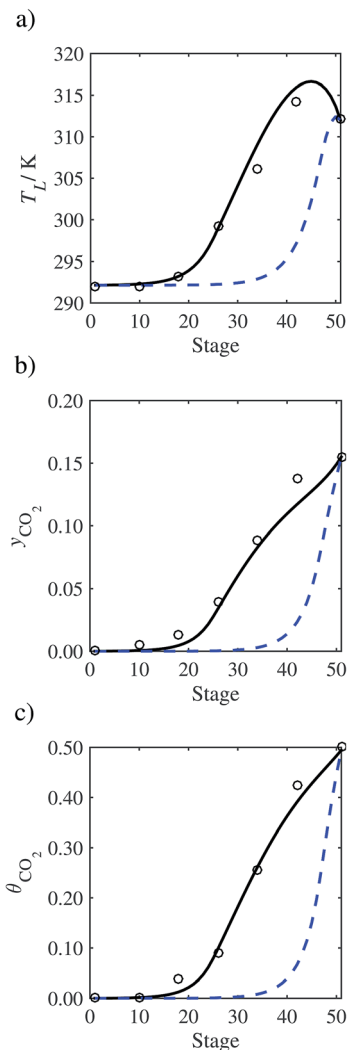


Fig. 12 Comparison of the pilot-plant data (circles)¹³ and the result of our model (curves) for Run T16. The dashed curves represent the results obtained without scaling the CO_2 diffusivity ($\tau = 1$) and the continuous curves the results obtained with the ED correlation (eqn (48)) by scaling the CO_2 diffusivity in the liquid phase to 4.1% of its original value ($\tau = 0.041$). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading. Stage 50 corresponds to the bottom of the column.

pilot-plant runs, the value of τ is now estimated by fitting to selected pilot-plant data. In keeping with our requirement to rely on only a limited amount of experimental data in the first step of the solvent selection process (and the exploration of a large space of solvents), the value τ is estimated from only one pilot-plant run (Run T22 of Tontiwachwuthikul *et al.*¹³). Additionally, this value is determined by minimizing the absolute deviation between the model predictions and the gas and liquid composition profiles, y_{CO_2} and θ_{CO_2} . The resulting



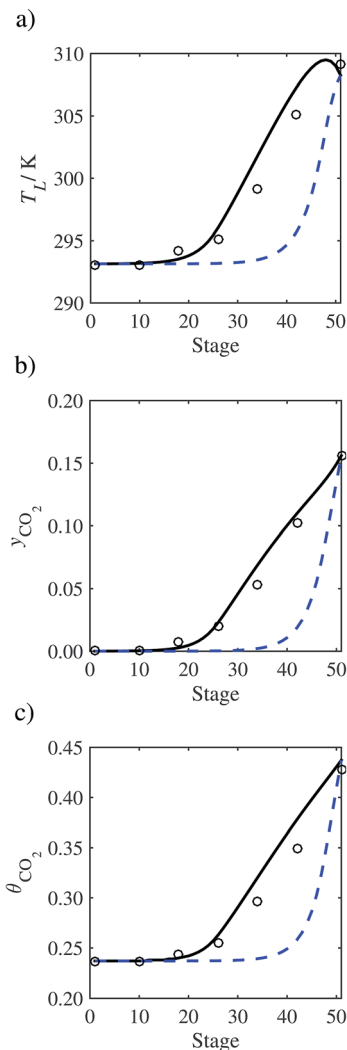


Fig. 13 Comparison of the pilot-plant data (circles)¹³ and the result of our model (curves) for Run T17. The dashed curves represent the results obtained without scaling the CO_2 diffusivity ($\tau = 1$) and the continuous curves the results obtained with the ED correlation (eqn (48)) by scaling the CO_2 diffusivity in the liquid phase to 4.1% of its original value ($\tau = 0.041$). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading. Stage 50 corresponds to the bottom of the column.

temperature profile for Run T22 is hence predicted rather than correlated. The value of τ in the ED correlation (eqn (48)) that results in the minimum deviation from the compositional pilot-plant data corresponds to a scaling of the liquid-phase diffusivity to 4.1% of its original value (*i.e.*, $\tau = 0.041$). The liquid-phase temperature, gas-phase CO_2 concentration, and liquid-phase CO_2 concentration profiles for Run T22, determined with and without the scaling factor for the diffusivity of CO_2 , are represented in Fig. 7 as continuous and dashed curves, respectively.



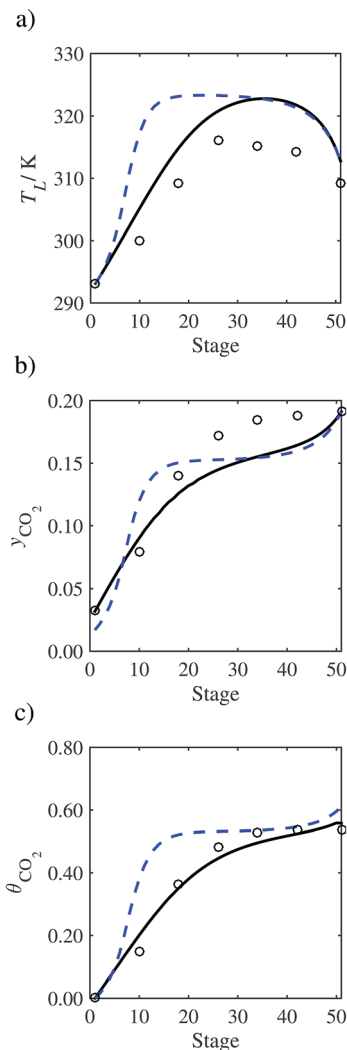


Fig. 14 Comparison of the pilot-plant data (circles)¹³ and the result of our model (curves) for Run T18. The dashed curves represent the results obtained without scaling the CO_2 diffusivity ($\tau = 1$) and the continuous curves the results obtained with the ED correlation (eqn (48)) by scaling the CO_2 diffusivity in the liquid phase to 4.1% of its original value ($\tau = 0.041$). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading. Stage 50 corresponds to the bottom of the column.

As the parameter τ is estimated from pilot-plant data, this may be correcting any potential errors in the chosen mass-transfer correlation rather than the diffusivity alone. To the best of our knowledge, there is no published experimental data for the diffusion of the carbamate product in aqueous MEA. In order to assess the impact of the choice of mass-transfer correlation on the value of τ , we also implement the mass-transfer correlations of Rocha *et al.*^{60,151} in our current model of the absorber with the same scaling factor of $\tau = 0.041$. The liquid-phase

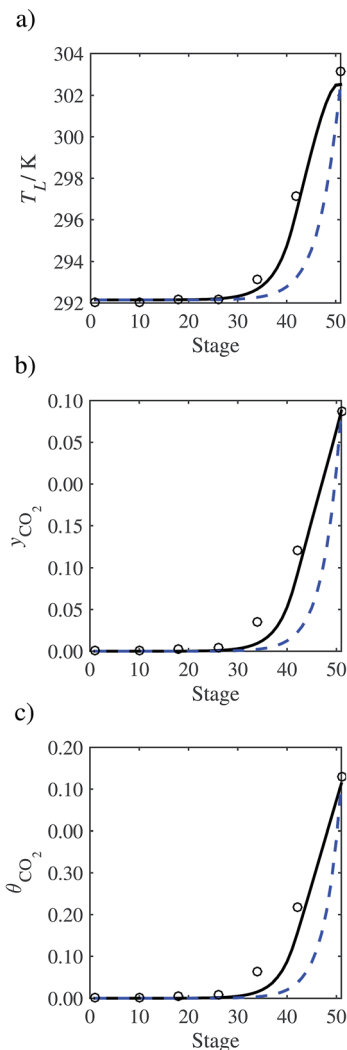


Fig. 15 Comparison of the pilot-plant data (circles)¹³ and the result of our model (curves) for Run T19. The dashed curves represent the results obtained without scaling the CO_2 diffusivity ($\tau = 1$) and the continuous curves the results obtained with the ED correlation (eqn (48)) by scaling the CO_2 diffusivity in the liquid phase to 4.1% of its original value ($\tau = 0.041$). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading. Stage 50 corresponds to the bottom of the column.

temperature, gas-phase CO_2 concentration, and liquid-phase CO_2 concentration profiles for Run T22 obtained with the two different mass transfer correlations are compared in Fig. 8. It can be seen that the profiles obtained are very similar: with the correlation of Rocha *et al.*^{60,151} there is a slight underestimation of the composition of CO_2 in the liquid and gas phases, though one should bear in mind that the diffusivity parameter is determined from the data using the correlation of Onda *et al.*^{57,58} Both mass-transfer correlations predict the experimental



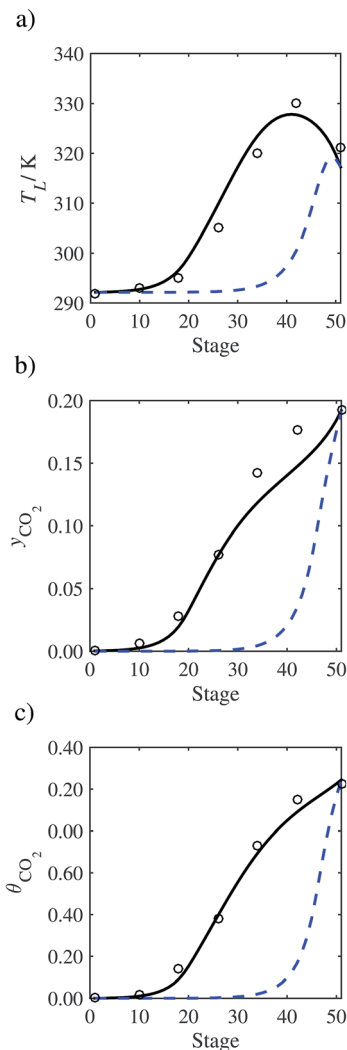


Fig. 16 Comparison of the pilot-plant data (circles)¹³ and the result of our model (curves) for Run T20. The dashed curves represent the results obtained without scaling the CO₂ diffusivity ($\tau = 1$) and the continuous curves the results obtained with the ED correlation (eqn (48)) by scaling the CO₂ diffusivity in the liquid phase to 4.1% of its original value ($\tau = 0.041$). (a) Temperature profile for the liquid phase, (b) gas-phase CO₂ concentration profile, and (c) liquid-phase CO₂ loading. Stage 50 corresponds to the bottom of the column.

temperature profiles equally well. In conclusion, the value of τ is found to be essentially independent of the choice of the mass-transfer correlation (at least for this set of process conditions), confirming our hypothesis that a scaling of the diffusivity is necessary to accurately model the process.

3.3.2 Model predictions. Other runs with the MEA absorption pilot plant can now be simulated in a predictive manner with the ED correlation (eqn (48)) using the value of the diffusivity parameter ($\tau = 0.041$) obtained based on Run T22. The



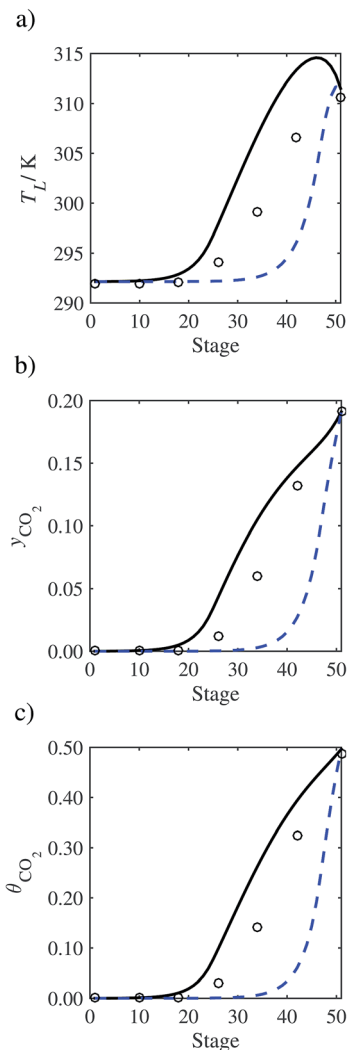


Fig. 17 Comparison of the pilot-plant data (circles)¹³ and the result of our model (curves) for Run T21. The dashed curves represent the results obtained without scaling the CO_2 diffusivity ($\tau = 1$) and the continuous curves the results obtained with the ED correlation (eqn (48)) by scaling the CO_2 diffusivity in the liquid phase to 4.1% of its original value ($\tau = 0.041$). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading. Stage 50 corresponds to the bottom of the column.

predictions for the profiles of the temperature in the liquid phase, the gas-phase CO_2 concentration, and liquid-phase CO_2 loading for the different Runs T13 to T21 are represented in Fig. 9–17 with continuous curves. These runs represent a variety of operating conditions in terms of amine concentration, inlet liquid-phase CO_2 loading, inlet gas-phase CO_2 concentration, and gas-to-liquid flow rate ratio (see Table 6).

Good agreement is found between the model predictions and the pilot-plant data for Runs T13, T14, T16, T17, T19, and T20 in relation to the liquid-phase



temperature and the liquid- and gas-phase compositions along the entire length of the column. There is a slight over-prediction of the temperature and the composition profile for Runs T15 and T21, though the compositions at the top and at the bottom of the column are accurately described. The outlet liquid temperature for Run T18 is overestimated by about 12 K; there is a good match between the model predictions and the experimental values for the composition profiles. A good representation of the temperature bulges for Runs T16 and T20 can be seen in terms of their location along the column, and the amplitude is predicted accurately for both of these runs. To the best of our knowledge, no other published model provides a description of the temperature bulge to this level of fidelity.^{29,33,48,99}

In all runs except Run T18, flat profiles are observed toward the top of the column. This plateau means that a maximum in the absorption is reached, indicating that equilibrium is achieved not only at the vapour–liquid interface but also between the bulk liquid and the bulk vapour phases. As a result the profiles are not affected by making the column any higher. This plateau is depicted in Fig. 18 for Run T22. It is apparent that the temperatures of the liquid phase, the vapour phase, and the vapour–liquid interface are all equal between stages 1 and 15 (*i.e.* at the top of the column). Similarly, the CO₂ gas-phase composition in the bulk vapour phase and at the vapour interface are equal between stages 1 and 15, as are the CO₂ loadings in the liquid phase and at the interface. The profiles for the temperature of the liquid phase and the temperature at the interface are found to be almost identical, suggesting that all the heat-transfer resistance is in the vapour film.

Run T18 stands out from this set of runs as it does not exhibit a plateau in the profiles as found for the other operating conditions, meaning that the whole length of the column is used for absorption. The lack of a plateau region is clearly apparent from Fig. 19. From Table 6, one can see that Run T18 corresponds to the lowest amount of MEA in the lean solvent and the highest amount of CO₂ in the flue gas compared to the other runs. As a result the whole column is required for absorption. In other runs, the totality of CO₂ is absorbed as the gas travels up the column between stages 50 and 15, so that lean solvent flowing at the top of the absorber (stages 1 to 15) has essentially no CO₂ left to absorb. Run T18 is the only run where there is still CO₂ in the gas stream leaving the absorber, so the lean solvent starts absorbing CO₂ as soon as it enters the column at the top. These differences explain why a larger discrepancy is observed between the predicted and measured profiles for Run T18.

The behaviour of Runs T15, T18, and T21 could be represented more accurately by estimating specific values of τ for these runs. However, this would not be in keeping with our objective to develop a predictive modelling platform to support solvent-design activities. The good overall quantitative agreement achieved with a unique value of τ indicates that the scaling factor can be applied in a transferable manner at different operating conditions (at least for similar types of column packing).

An analysis of the deviation between the column profiles presented here and those obtained when a different run is chosen to estimate τ is also undertaken. The same method is applied to estimate τ with the ED correlation (eqn (48)) based on pilot plant Runs T13–T21. The values of τ range between 0.027 (for Run T19) and 0.076 (for Run T21). These extreme values are then used to predict the column



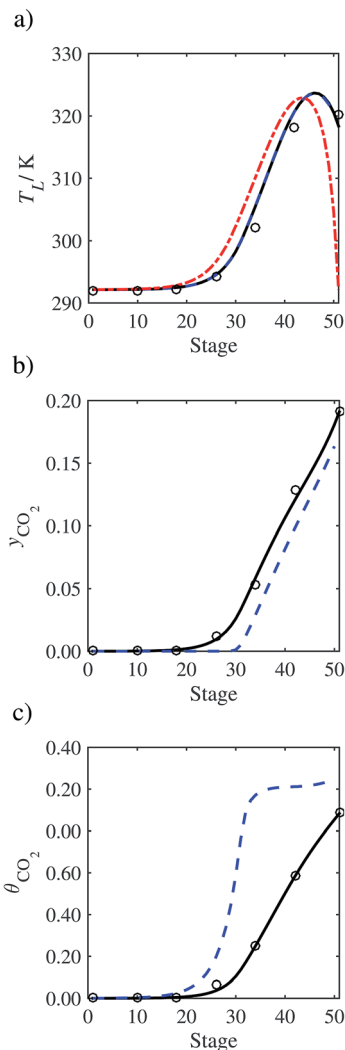


Fig. 18 Pilot plant data for Run T22 of Tontiwachwuthikul *et al.*¹³ Comparison of the predictions of our model for: (a) the temperature of the bulk liquid phase (continuous curve), the temperature at the vapour–liquid interface (dashed curve indistinguishable from the continuous curve), and the temperature of the bulk vapour phase (dot-dashed curve); (b) the gas-phase CO_2 concentration in the bulk vapour phase (continuous curve) and at the interface (dashed curve); (c) the liquid-phase CO_2 loading in the bulk liquid phase (continuous curve) and at the interface (dashed curve).

profiles (for T_L , y_{CO_2} , and θ_{CO_2}). The absolute errors between the values for each variable obtained with $\tau = 0.041$ (for Run T22) and the values obtained from the extreme values of τ are calculated, and averaged over all of the stages and column profiles. The calculated mean errors are 4.35 K for T_L , 0.080 for θ_{CO_2} and 0.023 for y_{CO_2} . These values provide an indication of the error bounds for the profiles presented in our current work, based on choosing any single pilot plant run arbitrarily to estimate the value of τ .



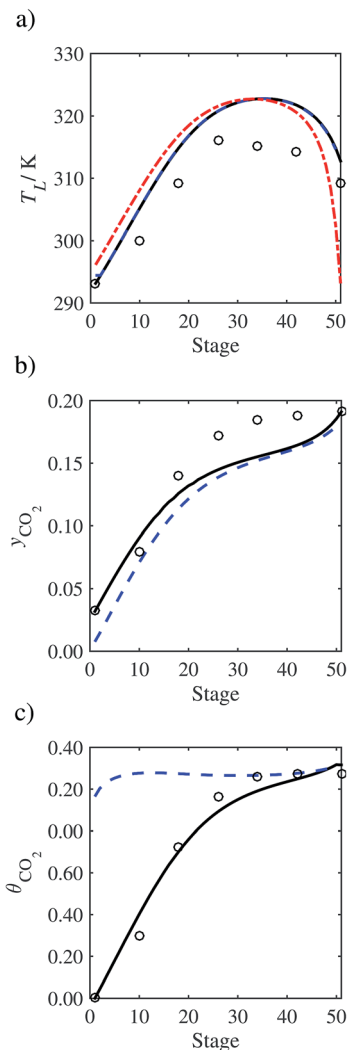


Fig. 19 Pilot plant data for Run T18 of Tontiwachwuthikul *et al.*¹³ Comparison of the predictions of our model for: (a) the temperature of the bulk liquid phase (continuous curve), the temperature at the vapour–liquid interface (dashed curve indistinguishable from the continuous curve), and the temperature of the bulk vapour phase (dot-dashed curve); (b) the gas-phase CO_2 concentration in the bulk vapour phase (continuous curve) and at the interface (dashed curve); (c) the liquid-phase CO_2 loading in the bulk liquid phase (continuous curve) and at the interface (dashed curve).

3.4 Sensitivity analysis

A sensitivity analysis is now employed to assess the relative importance of selected mass- and heat-transfer parameters. Different key parameters in the model are altered to observe their impact on the predicted profiles. The properties considered are listed in Table 8.

3.4.1 Mass transfer. In the model, mass transfer is mediated *via* mass-transfer coefficients.^{57,58} These coefficients are highly dependent on the viscosity,



the diffusivity, and to a certain degree, the surface tension of the fluid considered; these properties are thus considered as key parameters in the model and their influence on the temperature and composition profiles are assessed. The values of the parameters assessed in the sensitivity analysis are varied within physically realistic ranges. We should note that though marked variations are expected for the diffusivity and viscosity for a reacting system of this type, the change in the vapour–liquid surface tension is expected to be somewhat less dramatic. The values of the vapour and liquid diffusivities are doubled and halved, and the vapour–liquid surface tension is varied within $\pm 10 \text{ mN m}^{-1}$ from the nominal value. Varying the diffusivity or viscosity in the gas phase has no discernable effect on the temperature profile of the liquid or the composition profiles. However, varying these properties in the liquid phase results in a significant variation in the profiles. The effect on the profiles due to the variation of the liquid viscosity and the diffusivity in the liquid phase is represented in Fig. 20 and 21. Varying the vapour–liquid interfacial tension produces a less significant variation in the profiles as can be seen in Fig. 22. The data that are originally considered for the surface-tension correlation in our model are for a mixture of only H_2O and MEA.¹⁵² Jayarathna *et al.*¹⁵³ have recently published experimental data of the surface tension of mixtures of H_2O , MEA and CO_2 , finding an increase in the vapour–liquid interfacial tension on absorption of CO_2 . A new correlation from these data is also implemented in our process model. A comparison of the temperature and composition profiles obtained with the two different correlations is shown in Fig. 22. The difference in the profiles is negligible so changing the surface tension correlation is not considered necessary in this case. This also suggests that the process model is less sensitive to realistic changes in the interfacial tension than for corresponding changes in the diffusivity and viscosity. We should note that the vapour–liquid surface tension can also be determined from the SAFT-VR EOS within a density-functional treatment.^{84,154–156} We do not

Table 8 Sensitivity analysis of key properties in the model

Parameter description	Variation	Effect
Vapour diffusivity	Doubled Halved	Negligible effect
Liquid diffusivity	Doubled Halved	Significant effect See Fig. 20
Vapour viscosity	Doubled Halved	Negligible effect
Liquid viscosity	Doubled Halved	Significant effect See Fig. 21
Surface tension	$+ 10 \text{ mN m}^{-1}$ $- 10 \text{ mN m}^{-1}$	Negligible effect see Fig. 22
Heat transfer in the liquid	$\times 10$ $\div 10$	Negligible effect
Heat transfer in the vapour	$\times 10$ $\div 10$	Limited effect See Fig. 23
Heat of vaporization of water	Doubled Halved	Significant effect See Fig. 24
Mass fraction of water in inlet flue gas	Varied between 0 and 0.072	Negligible effect See Fig. 25



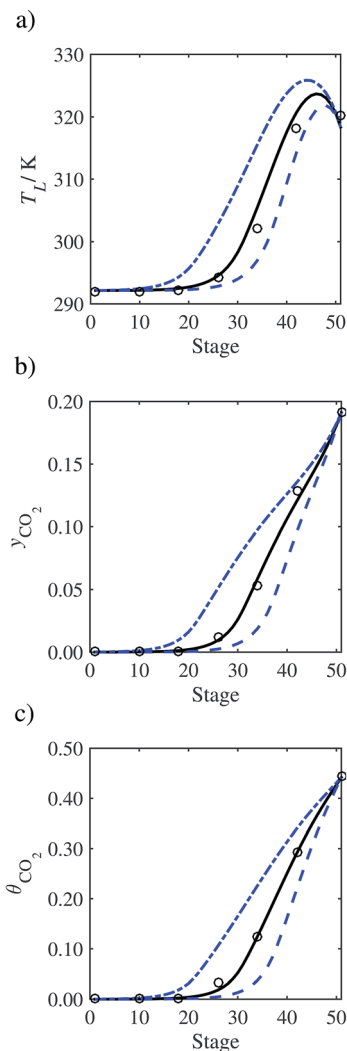


Fig. 20 Sensitivity analysis of the CO_2 diffusivity in the liquid phase on the profiles predicted with our model in terms of the scaling parameter τ compared with the experimental pilot-plant data of Run T22 of Tontiwachwuthikul *et al.*:¹³ $\tau = 0.041$ (continuous curve), $\tau = 0.082$ (dashed curve), and $\tau = 0.021$ (dot-dashed curve). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading.

however pursue this approach here as the full DFT of the ternary H_2O –MEA– CO_2 mixture is computationally challenging, and as has just been discussed the use of the correlations for the surface tension of the H_2O –MEA binary system provides a good representation of the pilot-plant data for the full mixture. The use of SAFT-VR DFT will be considered in future studies.

It should be noted that the values of the mole fractions and temperatures at the outlets are not affected by the variations of the surface tension or the viscosity and diffusivity in the liquid phase. It would be safe to assume that under these operating conditions, where the residence time is large enough for the system to reach



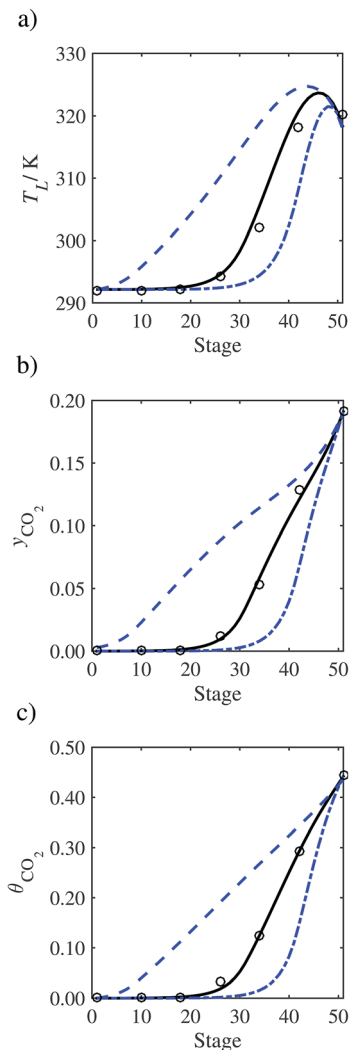


Fig. 21 Sensitivity analysis of the liquid viscosity on the profiles predicted with our model compared with the experimental pilot-plant data of Run T22 of Tontiwachwuthikul *et al.*:¹³ nominal value (continuous curve), viscosity doubled (dashed curve), and viscosity halved (dot-dashed curve). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading.

equilibrium, the outlet values are determined solely by thermodynamic considerations while the composition profiles are mostly dependent on mass transfer.

3.4.2 Heat transfer. The major source of heat in the absorption process is a consequence of the exothermic reaction between MEA and CO_2 , which occurs in the liquid phase. This heat can then be transferred to the vapour phase. Another major element in the energy balance is the heat associated with the transfer of water from one phase to the other. In the model, the heat transfer is mediated *via* heat-transfer coefficients.¹³⁹ Two notable thermal parameters can be identified in the model: the heat-transfer coefficient for the liquid phase, and the heat-transfer



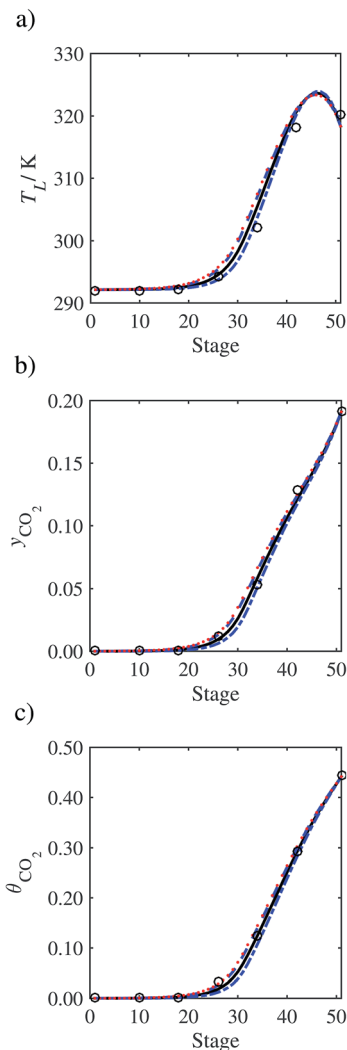


Fig. 22 Sensitivity analysis of the vapour–liquid surface tension on the profiles predicted with our model compared with the experimental pilot-plant data of Run T22 of Tontiwachwuthikul *et al.*¹⁵ nominal value (continuous curve),¹⁵² +10 mN m^{-1} (dashed curve), –10 mN m^{-1} (dot-dashed curve), and the surface tension correlation for the loaded MEA solution obtained from Jayarathna *et al.*¹⁵³ (dotted curve). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading.

coefficient for the gas phase. In addition, the magnitude of the enthalpy changes as the components are transferred from one phase to another or as the temperature changes in the gas and liquid streams affects the overall temperature profile. For instance, Kvamsdal and Rochelle¹⁰³ and Faramarzi *et al.*⁴⁸ have identified heat capacities as sensitive properties for the temperature profiles. Here, we consider the impact of the heat of vaporization of water on the model predictions. Although it is a well characterized thermodynamic quantity, its variation provides insights into the behaviour of the column.



The value of the heat of vaporization of water is doubled and halved relative to the value predicted with the SAFT-VR EoS, and the values of the heat transfer coefficients are multiplied by 10 and divided by 10 (Table 8). The variation of the liquid heat-transfer coefficient has no visible effect on the liquid-phase temperature and composition profiles. The effects on the profiles resulting from varying the heat-transfer coefficients and the heat of vaporization of water are presented in Fig. 23 and 24, respectively.

As can be seen from Fig. 23, despite a variation of one order of magnitude in the vapour heat-transfer coefficient, the effect on the temperature profile is very

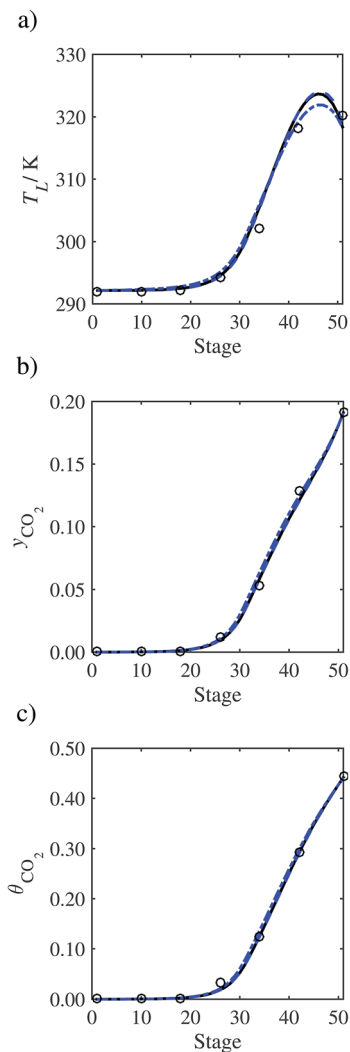


Fig. 23 Sensitivity analysis of the vapour-phase heat-transfer coefficient on the profiles predicted with our model compared with the experimental pilot-plant data of Run T22 of Tontiwachwuthikul *et al.*:¹³ Nominal value (continuous curve), coefficient value multiplied by 10 (dashed curve), and coefficient value divided by 10 (dot-dashed curve). (a) Temperature profile for the liquid phase, (b) gas-phase CO₂ concentration profile, and (c) liquid-phase CO₂ loading.



limited. It can be concluded that for the operating conditions under consideration, the model is insensitive to the heat-transfer coefficients and there is no need to investigate heat-transfer correlations further.

The variation of the heat of vaporization of water is found to have a significant impact on the liquid phase temperature profile when its value is doubled, and a lesser impact on the composition profiles; it has a limited effect on the end-point values. An increase in the heat of vaporization magnifies the amplitude of the temperature bulge, whereas a reduction eliminates the bulge. This suggests that the heat of vaporization of H₂O is responsible for the rate at which the liquid heats up

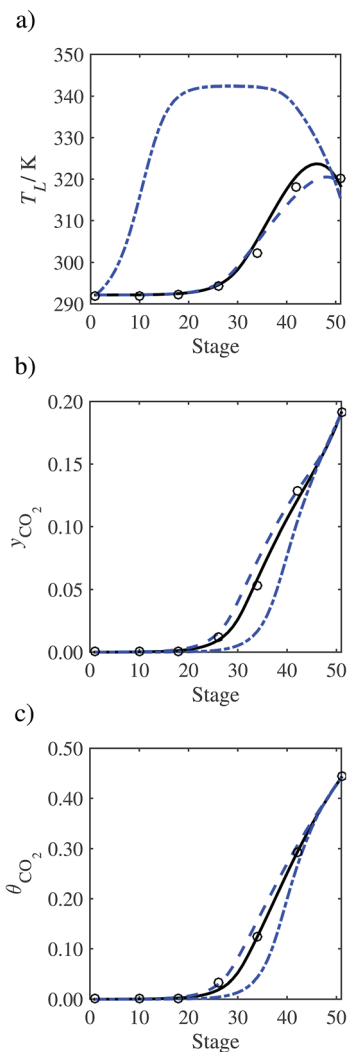


Fig. 24 Sensitivity analysis of the enthalpy of vaporization of water on the profiles predicted by our model compared with the experimental pilot-plant data of Run T22 of Tontiwachwuthikul *et al.*:¹³ Nominal value (continuous curve), enthalpy doubled (dashed curve), and enthalpy halved (dot-dashed curve). (a) Temperature profile for the liquid phase, (b) gas-phase CO₂ concentration profile, and (c) liquid-phase CO₂ loading.



and cools down. A similar effect is observed when a sensitivity analysis is carried out on the heat of absorption of CO_2 , however the analysis is not presented here because the adjustment of both properties leads to similar (but opposite) corrections to the energy balance. The key elements to predict the temperature bulge accurately are therefore the enthalpy of absorption of CO_2 and the enthalpy of vaporization of water. The absorption of CO_2 releases energy that heats both phases. As the gas temperature increases upon entering the column, water is transferred from the liquid phase to the gas phase through evaporation to maintain saturation; this results in an absorption of energy, and a consequent decrease in temperature.

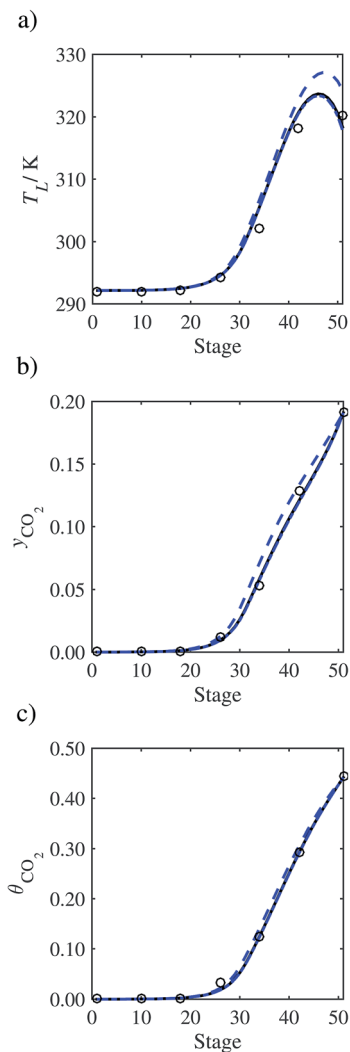


Fig. 25 Sensitivity analysis of the amount of water in the inlet flue gas on the profiles predicted with our model compared with the experimental pilot-plant data of Run T22 of Tontiwachwuthikul et al.¹³ Nominal value (continuous curve), $y_{\text{H}_2\text{O}} = 0.072$ (dashed curve), and $y_{\text{H}_2\text{O}} = 0$ (dot-dashed curve). (a) Temperature profile for the liquid phase, (b) gas-phase CO_2 concentration profile, and (c) liquid-phase CO_2 loading.



In conclusion, the competition between these two thermal effects (the absorption of CO₂ increases the temperature and the vaporization of water decreases the temperature) is responsible for the observed and predicted temperature bulge. This interpretation is consistent with that reported in Mac Dowell *et al.*⁹⁹

3.4.3 Effect of humidity in the flue gas. The amount of water in the flue gas is considered to be important in determining the position of the temperature bulge in the absorber.^{103,157} In the case of the pilot plants examined by Tontiwachwuthikul *et al.*¹³ and Dugas,¹⁴ the inlet flue gas is not saturated in water, while the flue gas is saturated in the pilot-plant studies of Tobiesen *et al.*⁴⁷ and Gabrielsen *et al.*¹⁶ In order to assess the impact of humidity in our model of the absorber, we vary the amount of water in the inlet flue gas from $\omega_{\text{H}_2\text{O}}^{\text{V}} = 0$ to 0.072 (corresponding to a mole fraction of 0.12) (water-rich). The effect on the profiles is represented in Fig. 25. It is apparent that varying the amount of water in the flue gas does not have a significant impact on the composition profiles. The effect is visible only for the liquid-phase temperature profile, where the outlet temperature value is higher by approximately 5 K for the saturated gas $\omega_{\text{H}_2\text{O}}^{\text{V}} = 0.072$ than for the dry gas $\omega_{\text{H}_2\text{O}}^{\text{V}} = 0$. Biliyok *et al.*⁵⁶ also find that increasing the moisture content of the flue gas affects the absorber temperature profile, however the increase in temperature is more significant in their study because a larger change in moisture content is considered ($\omega_{\text{H}_2\text{O}}^{\text{V}}$ varies between 0.015 and 0.15, where the latter value represents a two-phase aqueous system). The negligible effect on the composition profiles is in contrast to the work of Mac Dowell *et al.*⁹⁹ who found a large effect of the humidity of the inlet gas on the flux profiles in the mass transfer zone.

4 Conclusions

An absorber model for CO₂ capture is developed with the aim of being as predictive as possible, in order to support solvent-design activities prior to extensive experimental investigations. The heat and mass transfer are described with rate-based equations, in common with many other process models. Unusually, however, a physical approach is taken to model the chemical reactions taking place in the absorber. Both vapour-liquid equilibrium and chemical equilibrium are treated within the SAFT-VR thermodynamic framework, ensuring a consistent and accurate representation of the physical interactions in the system under the assumption that the reaction kinetics are not rate-determining. This approach lends itself to straightforward extension to other solvents, as a consequence of the transferable nature of the SAFT molecular models and the relatively small number of parameters and data required to develop them.

Without making use of pilot-plant data in model development, we find that the proposed model can generally be used to obtain a best-case performance of the solvent in question. This modelling approach is valuable for narrowing the solvent search space as solvents may be quickly rejected by comparing their performance in such a test. With very limited pilot-plant data we find that by adjusting a single parameter in the ED correlation for the mass transfer, that corrects for the diffusivity of CO₂ in the liquid phase, the model can be used to predict with quantitative accuracy a variety of different operating conditions. Excellent predictions are obtained for the liquid-phase temperature profiles and the liquid- and gas-phase compositions along the column in most cases, with



moderate deviations in some instances. The comparison of the absorption performance of different solvents *via* this method may further aid in the narrowing of the solvent search space, and then a more quantitative comparison could be carried out.

Following accurate predictions of the column profiles, a careful sensitivity analysis is conducted. We find that the liquid viscosity and diffusivity are key properties for the prediction of the composition profiles. The column profiles are also shown to be sensitive to the thermodynamic properties that are major sources of heat generation or dissipation.

The main benefit of the proposed modelling framework, which is based on the physical modelling of the underlying chemical reactions, is the ability to assess new solvents for which there may be limited data. This can be further enhanced through the adoption of a group-contribution EoS, which makes use of the same physical concepts, such as the SAFT- γ Mie EoS.¹²¹ The modification of the proposed model to use this group-contribution approach is straightforward and offers an additional predictive capability as new solvents may be analyzed for which no experimental data are available. Additionally, it is clear that an extension of the work presented here to the desorption process and the coupling of the absorber and desorber is required.¹⁵⁸ This would allow for a more comprehensive predictive assessment of new solvents and for the rapid evaluation of many alternative multifunctional amines for the optimal capture of CO₂ from flue gas.

Nomenclature

$\dot{V}^L$	Volumetric flow rate of the liquid phase ($\text{m}^3 \text{s}^{-1}$)
$\dot{V}^V$	Volumetric flow rate of the vapour phase ($\text{m}^3 \text{s}^{-1}$)
ε	Void fraction (-)
ε_{L_0}	Operating void space in the packing (-)
η^L	Dynamic viscosity of the liquid phase ($\text{kg m}^{-1} \text{s}^{-1}$)
η^V	Dynamic viscosity of the vapour phase ($\text{kg m}^{-1} \text{s}^{-1}$)
λ_T^L	Liquid thermal conductivity ($\text{W m}^{-1} \text{K}^{-1}$)
λ_T^V	Vapour thermal conductivity ($\text{W m}^{-1} \text{K}^{-1}$)
λ_{ij}	Parameter of the attractive range of the intermolecular potential between two segments i and j (-)
$\mu_{i,j}^L$	Chemical potential of component i in the liquid phase on stage j (J mol^{-1})
$\mu_{i,j}^V$	Chemical potential of component i in the gas phase on stage j (J mol^{-1})
$\omega_{\text{H}_2\text{O}}^V$	Mass fraction of water in the vapour phase
ϕM	Association factor for the liquid mixture (-)
ϕ_j	Association factor of solvent j (-)
ρ^L	Density of the liquid phase (kg m^{-3})
ρ^V	Density of the vapour phase (kg m^{-3})
σ	Vapour–liquid surface tension (N m^{-1})
σ_c	Critical surface tension of the packing material (N m^{-1})
Σ_v	Atomic diffusion volume ($\text{\AA}$)
σ_{ii}	Diameter of segments forming molecule i ($\text{\AA}$)
τ	Scaling factor for the liquid diffusivity of CO ₂ in the solvent (-)
θ_{CO_2}	CO ₂ loading in the liquid phase (-)
$\underline{x}_j$	Vector of liquid mole fractions on stage j (-)



y_j	Vector of vapour mole fractions on stage j (-)
$\underline{z}_i$	Molar composition vector of a stream consisting of pure component i (-)
$\varepsilon_{ab,ij}^{\text{HB}}/k$	Depth of the interaction potential between association site a on molecule i and site b of molecule j (K)
ε_{ij}/k	Depth of the intermolecular potential between two segments i and j (K)
a_j^*	Interfacial area density on stage j ($\text{m}^2 \text{m}^{-3}$)
a_p	Specific surface area of the packing ($\text{m}^2 \text{m}^{-3}$)
A_{section}	Cross-sectional area of the column (m^2)
$a_{T,j}$	Total interfacial area available for heat or mass transfer on stage j (m^2)
c	Total number of components
$C_{i,j}^{\text{L}}$	Concentration of component i in the liquid phase on stage j (mol m^{-3})
$C_{i,j}^{\text{V}}$	Concentration of component i in the vapour phase on stage j (mol m^{-3})
$C_{i,j}^{\text{L,L}}$	Concentration of component i at the liquid–vapour interface in the liquid phase (mol m^{-3})
$C_{i,j}^{\text{L,V}}$	Concentration of component i at the liquid–vapour interface in the vapour phase (mol m^{-3})
C_p^{L}	Specific isobaric heat capacity of the liquid phase ($\text{J kg}^{-1} \text{K}^{-1}$)
C_p^{V}	Specific isobaric heat capacity of the vapour phase ($\text{J kg}^{-1} \text{K}^{-1}$)
D_i^{L}	Diffusion coefficient of component i in the liquid phase ($\text{m}^2 \text{s}^{-1}$)
D_i^{V}	Diffusion coefficient of component i in the vapour phase ($\text{m}^2 \text{s}^{-1}$)
$D_{i,j}^{\circ}$	Mutual diffusion coefficient of solute i at very low concentrations in solvent j ($\text{cm}^2 \text{s}^{-1}$)
d_s	Diameter of a sphere of the same surface as a single packing particle (m)
D_z	Stage height (m)
E_j^{L}	Net gain of energy of the liquid phase on stage j (W)
E_j^{V}	Net loss of energy from the vapour phase on stage j (W)
Fr^{L}	Liquid-phase Froude number (-)
g	Gravitational acceleration (m s^{-2})
H_j^{L}	Enthalpy of the liquid phase on stage j (J mol^{-1})
H_j^{V}	Enthalpy of the vapour phase on stage j (J mol^{-1})
h_{packing}	Total packing height (m)
$h_{T,j}^{\text{L}}$	Liquid-phase heat-transfer coefficient on stage j ($\text{W m}^{-2} \text{K}^{-1}$)
$h_{T,j}^{\text{V}}$	Vapour-phase heat-transfer coefficient on stage j ($\text{W m}^{-2} \text{K}^{-1}$)
k	Boltzmann constant (J K^{-1})
k_j^{L}	Liquid-phase mass-transfer coefficient on stage j (m s^{-1})
k_j^{V}	Vapour-phase mass-transfer coefficient on stage j (m s^{-1})
k_{ij}	Binary interaction parameter used to compute the strength of the interactions between segments i and j (-)
L_j	Liquid molar flow rate leaving stage j (mol s^{-1})
L_p	Nominal packing size (m)
L_{spec}	Specific liquid flow rate (kg s^{-1})
M_i	Molecular weight of component i (g mol^{-1})
m_i	Number of segments in the molecule i
$N_{i,j}^{\text{L}}$	Net gain of species i in the liquid phase due to interphase transport on stage j (mol s^{-1})
$N_{i,j}^{\text{V}}$	Net loss of species i in the vapour phase due to interphase transport on stage j (mol s^{-1})
N_s	Number of stage in the column
$N_{s,ia}$	Number of sites of type a on molecule i
$N_{s,i}$	Number of site types for molecule i



P_j^I	Pressure at the vapour–liquid interface on stage j (MPa)
P_j^L	Pressure in the liquid phase on stage j (MPa)
P_j^V	Pressure in the vapour phase on stage j (MPa)
Pr^V	Prandtl number for the vapour phase (-)
$Q_{cond,j}^L$	Conductive heat flux in the liquid phase on stage j (W)
$Q_{diff,j}^L$	Diffusive heat flux in the liquid phase on stage j (W)
$Q_{cond,j}^V$	Conductive heat flux in the vapour phase on stage j (W)
$Q_{diff,j}^V$	Diffusive heat flux in the vapour phase on stage j (W)
$r_{c;ab,ij}$	Attractive range of the interaction potential between association site a on molecule i and site b of molecule j (Å)
Re^L	Liquid-phase Reynolds number based on the interfacial area (-)
Re^V	Vapour-phase Reynolds number (-)
Sc_i^L	Schmidt number of component i in the liquid phase (-)
Sc_i^V	Schmidt number of component i in the vapour phase (-)
T_j^I	Temperature at the vapour–liquid interface on stage j (K)
T_j^L	Temperature of the liquid phase on stage j (K)
T_j^V	Temperature of the vapour phase on stage j (K)
u^L	Liquid velocity (m s ⁻¹)
u^V	Vapour velocity (m s ⁻¹)
V_j^L	Molar volume of the liquid phase on stage j (cm ³ mol ⁻¹)
V_j^V	Molar volume of the vapour phase on stage j (cm ³ mol ⁻¹)
$V_{i,j}^{*L}$	Molar volume of the pure component i in the bulk liquid phase on stage j (cm ³ mol ⁻¹)
$V_{i,j}^{*V}$	Molar volume of the pure component i in the bulk vapour phase on stage j (cm ³ mol ⁻¹)
$V_j^{I,L}$	Molar volume of the liquid phase at the vapour–liquid interface on stage j (cm ³ mol ⁻¹)
$V_j^{I,V}$	Molar volume of the gas phase at the vapour–liquid interface on stage j (cm ³ mol ⁻¹)
V_i^m	Molar volume of solute i at its normal boiling temperature (cm ³ mol ⁻¹)
V_j	Vapour molar flow rate leaving stage j (mol s ⁻¹)
V_{spec}	Specific vapour flow rate (kg s ⁻¹)
We^L	Liquid-phase Weber number (-)
Re^L	Liquid-phase Reynolds number based on the specific surface area (-)

Acknowledgements

We are very grateful to Leila Faramarzi and Georgios Kontogeorgis for helpful discussions regarding details of the pilot-plant and mass-transfer correlations. C. V. B. and E. G. thank the Department of Chemical Engineering of Imperial College London and Engineering and Physical Sciences Research Council (EPSRC) of the UK for the award of PhD studentships. Additional funding to the Molecular Systems Engineering Group from the EPSRC (grants GR/T17595, GR/N35991, EP/E016340, EP/J003840/1, and EP/J014958) is also gratefully acknowledged.

References

- 1 The European Commission, *A Roadmap for Moving to a Competitive Low Carbon Economy in 2050*, <http://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:52011DC0112&from=EN>, 2011.



- 2 UNFCCC, *Adoption of the Paris Agreement*, <https://unfccc.int/resource/docs/2015/cop21/eng/l09r01.pdf>, 2015.
- 3 N. Mac Dowell, N. Florin, A. Buchard, J. Hallett, A. Galindo, G. Jackson, C. S. Adjiman, C. K. Williams, N. Shah and P. S. Fennell, *Energy Environ. Sci.*, 2010, **3**, 1645–1669.
- 4 S. Paul, A. K. Ghoshal and B. Mandal, *Chem. Eng. J.*, 2008, **144**, 352–360.
- 5 H. P. Mangalapally, R. Notz, S. Hoch, N. Asprion, G. Sieder, H. Garcia and H. Hasse, *Energy Proc.*, 2009, **1**, 963–970.
- 6 A. Bardow, K. Steur and J. Gross, *Ind. Eng. Chem. Res.*, 2010, **49**, 2834–2840.
- 7 F. Barzagli, M. Di Vaira, F. Mani and M. Peruzzini, *ChemSusChem*, 2012, **5**, 1724–1731.
- 8 H. P. Mangalapally, R. Notz, N. Asprion, G. Sieder, H. Garcia and H. Hasse, *Int. J. Greenhouse Gas Control*, 2012, **8**, 205–216.
- 9 J. Li, C. You, L. Chen, Y. Ye, Z. Qi and K. Sundmacher, *Ind. Eng. Chem. Res.*, 2012, **51**, 12081–12088.
- 10 Y. K. Salkuyeh and M. Mofarahi, *Int. J. Energy Res.*, 2013, **37**, 973–981.
- 11 S. Chen, S. Chen, X. Fei, Y. Zhang and L. Qin, *Ind. Eng. Chem. Res.*, 2015, **54**, 7212–7218.
- 12 P. V. Kortunov, M. Siskin, L. S. Baugh and D. C. Calabro, *Energy Fuels*, 2015, **29**, 5940–5966.
- 13 P. Tontiwachwuthikul, A. Meisen and C. J. Lim, *Chem. Eng. Sci.*, 1992, **47**, 381–390.
- 14 R. E. Dugas, PhD thesis, The University of Texas at Austin, 2006.
- 15 F. A. Tobiesen and H. F. Svendsen, *Ind. Eng. Chem. Res.*, 2006, **45**, 2489–2496.
- 16 J. Gabrielsen, H. F. Svendsen, M. L. Michelsen, E. H. Stenby and G. M. Kontogeorgis, *Chem. Eng. Sci.*, 2007, **62**, 2397–2413.
- 17 H. Habaki, J. M. Perera, S. E. Kentish, G. W. Stevens and W. Fei, *Sep. Sci. Technol.*, 2007, **42**, 701–716.
- 18 H. R. Godini and D. Mowla, *Chem. Eng. Res. Des.*, 2008, **86**, 401–409.
- 19 A. J. Cottrell, J. M. McGregor, J. Jansen, Y. Artanto, N. Dave, S. Morgan, P. Pearson, M. I. Attalla, L. Wardhaugh, H. Yu, A. Allport and P. H. M. Feron, *Energy Proc.*, 2009, **1**, 1003–1010.
- 20 R. Notz, H. P. Mangalapally and H. Hasse, *Int. J. Greenhouse Gas Control*, 2012, **6**, 84–112.
- 21 M. Akram, U. Ali, T. Best, S. Blakey, K. Finney and M. Pourkashanian, *Int. J. Greenhouse Gas Control*, 2016, **47**, 137–150.
- 22 G. Astarita, *Mass Transfer with Chemical Reaction*, Elsevier, 1967.
- 23 H. Hikita, S. Asai, H. Ishikawa and M. Honda, *Chem. Eng. J.*, 1977, **13**, 7–12.
- 24 P. V. Danckwerts, *Chem. Eng. Sci.*, 1979, **34**, 443–446.
- 25 S. S. Laddha and P. V. Danckwerts, *Chem. Eng. Sci.*, 1981, **36**, 479–482.
- 26 G. Astarita, D. W. Savage and A. Brito, *Gas Treating with Chemical Solvents*, John Wiley & Sons, 1983.
- 27 D. E. Penny and T. J. Ritter, *J. Chem. Soc., Faraday Trans. 1*, 1983, **79**, 2103–2109.
- 28 P. M. M. Blauwhoff, G. F. Versteeg and W. P. M. Van Swaaij, *Chem. Eng. Sci.*, 1984, **39**, 207–225.
- 29 L. Kucka, I. Muller, E. Y. Kenig and A. Górak, *Chem. Eng. Sci.*, 2003, **58**, 3571–3578.
- 30 Y. Zhang, H. Que and C.-C. Chen, *Fluid Phase Equilib.*, 2011, **311**, 67–75.



- 31 C. Kale, A. Górak and H. Schoenmakers, *Int. J. Greenhouse Gas Control*, 2013, **17**, 294–308.
- 32 A. Aboudheir, P. Tontiwachwuthikul and R. Idem, *Ind. Eng. Chem. Res.*, 2006, **45**, 2553–2557.
- 33 J. Gabrielsen, M. L. Michelsen, E. H. Stenby and G. M. Kontogeorgis, *AIChE J.*, 2006, **52**, 3443–3451.
- 34 W.-J. Choi, J.-B. Seo, S.-W. Cho, S.-W. Park and K.-J. Oh, *Korean J. Chem. Eng.*, 2009, **26**, 705–710.
- 35 M. Afkhamipour and M. Mofarahi, *Int. J. Greenhouse Gas Control*, 2013, **15**, 186–199.
- 36 R. Taylor and R. Krishna, *Multicomponent Mass Transfer*, John Wiley & Sons, 1993.
- 37 F. M. Khoury, *Multistage Separation Processes*, CRC Press, 2005.
- 38 J. Ying and D. A. Eimer, *Ind. Eng. Chem. Res.*, 2013, **52**, 2548–2559.
- 39 A. Lawal, M. Wang, P. Stephenson and H. Yeung, *Fuel*, 2009, **88**, 2455–2462.
- 40 P. V. Danckwerts, *Gas-Liquid Reactions*, McGraw-Hill, 1970.
- 41 W. J. DeCoursey, *Chem. Eng. Sci.*, 1982, **37**, 1483–1489.
- 42 W. P. M. van Swaaij and G. F. Versteeg, *Chem. Eng. Sci.*, 1992, **47**, 3181–3195.
- 43 M. S. Sivasubramanian, H. Sardar and R. H. Weiland, *Oil Gas J.*, 1985, **83**, 133–136.
- 44 T. Pintola, P. Tontiwachwuthikul and A. Meisen, *Gas Sep. Purif.*, 1993, **7**, 47–52.
- 45 M. A. Pacheco and G. T. Rochelle, *Ind. Eng. Chem. Res.*, 1998, **37**, 4107–4117.
- 46 N. A. Al-Baghli, S. A. Pruess, V. F. Yesavage and M. S. Selim, *Fluid Phase Equilib.*, 2001, **185**, 31–43.
- 47 F. A. Tobiesen, H. F. Svendsen and O. Juliussen, *AIChE J.*, 2007, **53**, 846–865.
- 48 L. Faramarzi, G. M. Kontogeorgis, M. L. Michelsen, K. Thomsen and E. H. Stenby, *Ind. Eng. Chem. Res.*, 2010, **49**, 3751–3759.
- 49 F. Khan, V. Krishnamoorthi and T. Mahmud, *Chem. Eng. Res. Des.*, 2011, **89**, 1600–1608.
- 50 T. Neveux, Y. L. Moullec, J.-P. Corriou and E. Favre, *Ind. Eng. Chem. Res.*, 2013, **52**, 4266–4279.
- 51 M. Saimpert, G. Puxty, S. Qureshi, L. Wardhaugh and A. Cousins, *Chem. Eng. Sci.*, 2013, **96**, 10–25.
- 52 S. A. Jayarathna, B. Lie and M. C. Melaaen, *Comput. Chem. Eng.*, 2013, **53**, 178–189.
- 53 M. Afkhamipour and M. Mofarahi, *Int. J. Greenhouse Gas Control*, 2014, **25**, 9–22.
- 54 M. Llano-Restrepo and E. Araujo-Lopez, *Int. J. Greenhouse Gas Control*, 2015, **42**, 258–287.
- 55 A. Lawal, M. Wang, P. Stephenson, G. Koumpouras and H. Yeung, *Fuel*, 2010, **89**, 2791–2801.
- 56 C. Biliyok, A. Lawal, M. Wang and F. Seibert, *Int. J. Greenhouse Gas Control*, 2012, **9**, 428–445.
- 57 K. Onda, E. Sada and H. Takeuchi, *J. Chem. Eng. Jpn.*, 1968, **1**, 62–66.
- 58 K. Onda, H. Takeuchi and Y. Okumoto, *J. Chem. Eng. Jpn.*, 1968, **1**, 56–62.
- 59 J. L. Bravo and J. R. Fair, *Ind. Eng. Chem. Process Des. Dev.*, 1982, **21**, 162–170.
- 60 J. A. Rocha, J. L. Bravo and J. R. Fair, *Ind. Eng. Chem. Res.*, 1996, **35**, 1660–1667.
- 61 R. Billet and M. Schultes, *Chem. Eng. Res. Des.*, 1999, **77**, 498–504.



- 62 J. D. Pandya, *Chem. Eng. Commun.*, 1983, **19**, 343–361.
- 63 F. Dolezalek, *Z. Phys. Chem.*, 1908, **64**, 727–747.
- 64 A. Fredenslund, R. L. Jones and J. M. Prausnitz, *AIChE J.*, 1975, **21**, 1086–1099.
- 65 J. M. Prausnitz, R. N. Lichtenthaler and E. G. de Azevedo, *Molecular Thermodynamics of Fluid-phase Equilibria*, Pearson Education, 1998.
- 66 Y. Song and C.-C. Chen, *Ind. Eng. Chem. Res.*, 2009, **48**, 7788–7797.
- 67 A. Fredenslund, *Vapor-liquid equilibria using UNIFAC: a group-contribution method*, Elsevier, 2012.
- 68 D. M. Austgen, G. T. Rochelle, X. Peng and C. C. Chen, *Ind. Eng. Chem. Res.*, 1989, **28**, 1060–1073.
- 69 D. M. Austgen, G. T. Rochelle and C. C. Chen, *Ind. Eng. Chem. Res.*, 1991, **30**, 543–555.
- 70 L. Kucka, E. Y. Kenig and A. Górak, *Ind. Eng. Chem. Res.*, 2002, **41**, 5952–5957.
- 71 C. Noeres, E. Y. Kenig and A. Górak, *Chem. Eng. Process.*, 2003, **42**, 157–178.
- 72 E. Y. Kenig, L. Kucka and A. Górak, *Chem. Eng. Technol.*, 2003, **26**, 631–646.
- 73 L. Kucka, J. Richter, E. Y. Kenig and A. Górak, *Sep. Purif. Technol.*, 2003, **31**, 163–175.
- 74 C.-C. Chen and L. B. Evans, *AIChE J.*, 1986, **32**, 444–454.
- 75 G. M. Bollas, C. C. Chen and P. I. Barton, *AIChE J.*, 2008, **54**, 1608–1624.
- 76 J. Gross and G. Sadowski, *Ind. Eng. Chem. Res.*, 2001, **40**, 1244–1260.
- 77 B. E. Poling, J. M. Prausnitz and J. P. O'Connell, *The Properties of Gases and Liquids*, McGraw-Hill, 5th edn, 2001.
- 78 K. A. Hoff, PhD thesis, Norges Teknisk-Naturvitenskapelige Universitet, 2003.
- 79 R. H. Weiland, M. Rawal and R. G. Rice, *AIChE J.*, 1982, **28**, 963–973.
- 80 J. Gabrielsen, M. L. Michelsen, E. H. Stenby and G. M. Kontogeorgis, *Ind. Eng. Chem. Res.*, 2005, **44**, 3348–3354.
- 81 J. Button and K. Gubbins, *Fluid Phase Equilib.*, 1999, **158**, 175–181.
- 82 N. Mac Dowell, F. Llovel, C. S. Adjiman, G. Jackson and A. Galindo, *Ind. Eng. Chem. Res.*, 2010, **49**, 1883–1899.
- 83 N. Mac Dowell, F. E. Pereira, F. Llovel, F. J. Blas, C. S. Adjiman, G. Jackson and A. Galindo, *J. Phys. Chem. B*, 2011, **115**, 8155–8168.
- 84 J. Rodríguez, N. Mac Dowell, F. Llovel, C. S. Adjiman, G. Jackson and A. Galindo, *Mol. Phys.*, 2012, **110**, 1325–1348.
- 85 W. G. Chapman, K. E. Gubbins, G. Jackson and M. Radosz, *Fluid Phase Equilib.*, 1989, **52**, 31–38.
- 86 W. G. Chapman, K. E. Gubbins, G. Jackson and M. Radosz, *Ind. Eng. Chem. Res.*, 1990, **29**, 1709–1721.
- 87 A. Gil-Villegas, A. Galindo, P. J. Whitehead, S. J. Mills, G. Jackson and A. N. Burgess, *J. Chem. Phys.*, 1997, **106**, 4168–4186.
- 88 A. Galindo, L. A. Davies, A. Gil-Villegas and G. Jackson, *Mol. Phys.*, 1998, **93**, 241–252.
- 89 A. Chremos, E. Forte, V. Papaioannou, A. Galindo, G. Jackson and C. S. Adjiman, *Chem. Eng. Trans.*, 2013, **35**, 427–432.
- 90 A. Chremos, E. Forte, V. Papaioannou, A. Galindo, G. Jackson and C. S. Adjiman, *Fluid Phase Equilib.*, 2016, **407**, 280–297.
- 91 A. Lymperiadis, C. S. Adjiman, A. Galindo and G. Jackson, *J. Chem. Phys.*, 2007, **127**, 234903.
- 92 A. Lymperiadis, C. S. Adjiman, G. Jackson and A. Galindo, *Fluid Phase Equilib.*, 2008, **274**, 85–104.



- 93 G. Jackson, W. G. Chapman and K. E. Gubbins, *Mol. Phys.*, 1988, **65**, 1–31.
- 94 I. G. Economou and M. D. Donohue, *AIChE J.*, 1991, **37**, 1875–1894.
- 95 F. E. Pereira, E. Keskes, A. Galindo, G. Jackson and C. S. Adjiman, *Comput. Chem. Eng.*, 2011, **35**, 474–491.
- 96 J. Burger, V. Papaioannou, S. Gopinath, G. Jackson, A. Galindo and C. S. Adjiman, *AIChE J.*, 2015, **61**, 3249–3269.
- 97 N. Mac Dowell, A. Galindo, G. Jackson and C. S. Adjiman, *Comput.-Aided Chem. Eng.*, 2010, **28**, 1231–1236.
- 98 C. V. Brand, J. Rodríguez, A. Galindo, G. Jackson and C. S. Adjiman, *Comput.-Aided Chem. Eng.*, 2012, **31**, 930–934.
- 99 N. Mac Dowell, N. J. Samsatli and N. Shah, *Int. J. Greenhouse Gas Control*, 2013, **12**, 247–258.
- 100 A. Arce, N. Mac Dowell, N. Shah and L. F. Vega, *Int. J. Greenhouse Gas Control*, 2012, **11**, 236–250.
- 101 N. Mac Dowell and N. Shah, *Int. J. Greenhouse Gas Control*, 2013, **13**, 44–58.
- 102 A. Alhajaj, N. Mac Dowell and N. Shah, *Int. J. Greenhouse Gas Control*, 2016, **44**, 26–41.
- 103 H. M. Kvamsdal and G. T. Rochelle, *Ind. Eng. Chem. Res.*, 2008, **47**, 867–875.
- 104 N. Mac Dowell, PhD thesis, Imperial College London, 2010.
- 105 M. S. Wertheim, *J. Stat. Phys.*, 1984, **35**, 19–34.
- 106 M. S. Wertheim, *J. Stat. Phys.*, 1984, **35**, 35–47.
- 107 M. S. Wertheim, *J. Stat. Phys.*, 1986, **42**, 459–476.
- 108 M. S. Wertheim, *J. Stat. Phys.*, 1986, **42**, 477–492.
- 109 W. G. Chapman, G. Jackson and K. E. Gubbins, *Mol. Phys.*, 1988, **65**, 1057–1079.
- 110 S. H. Huang and M. Radosz, *Ind. Eng. Chem. Res.*, 1990, **29**, 2284–2294.
- 111 S. H. Huang and M. Radosz, *Ind. Eng. Chem. Res.*, 1991, **30**, 1994–2005.
- 112 F. J. Blas and L. F. Vega, *Mol. Phys.*, 1997, **92**, 135–150.
- 113 J. Gross and G. Sadowski, *Ind. Eng. Chem. Res.*, 2002, **41**, 5510–5515.
- 114 T. Lafitte, D. Bessi eres, M. M. Pi eiro and J.-L. Daridon, *J. Chem. Phys.*, 2006, **124**, 024509.
- 115 T. Lafitte, M. M. Pi eiro, J.-L. Daridon and D. Bessi eres, *J. Phys. Chem. B*, 2007, **111**, 3447–3461.
- 116 T. Lafitte, A. Apostolakou, C. Avenda o, A. Galindo, C. S. Adjiman, E. A. M ller and G. Jackson, *J. Chem. Phys.*, 2013, **139**, 154504.
- 117 K. E. Gubbins, *Fluid Phase Equilib.*, 2016, **416**, 3–17.
- 118 S. Dufal, T. Lafitte, A. Galindo, G. Jackson and A. J. Haslam, *AIChE J.*, 2015, **61**, 2891–2912.
- 119 S. Dufal, T. Lafitte, A. J. Haslam, A. Galindo, G. N. I. Clark, C. Vega and G. Jackson, *Mol. Phys.*, 2015, **113**, 948–984.
- 120 V. Papaioannou, C. S. Adjiman, G. Jackson and A. Galindo, Group contribution methodologies for the prediction of thermodynamic properties and phase behaviour in mixtures, in *Molecular Systems Engineering*, ed. C. S. Adjiman and A. Galindo, Wiley-VCH, Germany, 2010, vol. 6, ch. 4, pp. 135–172.
- 121 V. Papaioannou, T. Lafitte, C. Avenda o, C. S. Adjiman, G. Jackson, E. A. M ller and A. Galindo, *J. Chem. Phys.*, 2014, **140**, 054107.
- 122 A. J. Haslam, A. Galindo and G. Jackson, *Fluid Phase Equilib.*, 2008, **266**, 105–128.



- 123 A. Galindo, A. Gil-Villegas, G. Jackson and A. N. Burgess, *J. Phys. Chem. B*, 1999, **103**, 10272–10281.
- 124 A. Gil-Villegas, A. Galindo and G. Jackson, *Mol. Phys.*, 2001, **99**, 531–546.
- 125 B. H. Patel, P. Paricaud, A. Galindo and G. C. Maitland, *Ind. Eng. Chem. Res.*, 2003, **42**, 3809–3823.
- 126 B. Behzadi, B. Patel, A. Galindo and C. Ghotbi, *Fluid Phase Equilib.*, 2005, **236**, 241–255.
- 127 P. Paricaud, L. Tazi and J. M. Borgard, *Int. J. Hydrogen Energy*, 2010, **35**, 978–991.
- 128 J. M. A. Schreckenberger, S. Dufal, A. J. Haslam, C. S. Adjiman, G. Jackson and A. Galindo, *Mol. Phys.*, 2014, **112**, 2339–2364.
- 129 G. N. I. Clark, A. J. Haslam, A. Galindo and G. Jackson, *Mol. Phys.*, 2006, **104**, 3561–3581.
- 130 P. Paricaud, A. Galindo and G. Jackson, *Ind. Eng. Chem. Res.*, 2004, **43**, 6871–6889.
- 131 R. P. Sear and G. Jackson, *Phys. Rev. E: Stat. Phys., Plasmas, Fluids, Relat. Interdiscip. Top.*, 1994, **50**, 386–394.
- 132 D. Ghonasgi, V. Perez and W. G. Chapman, *J. Chem. Phys.*, 1994, **101**, 6880–6887.
- 133 R. P. Sear and G. Jackson, *Mol. Phys.*, 1996, **87**, 517–521.
- 134 A. Galindo, S. J. Burton, G. Jackson, D. P. Visco Jr and D. A. Kofke, *Mol. Phys.*, 2002, **100**, 2241–2259.
- 135 R. P. Sear and G. Jackson, *Mol. Phys.*, 1994, **82**, 1033–1048.
- 136 R. P. Sear and G. Jackson, *J. Chem. Phys.*, 1996, **105**, 1113–1120.
- 137 R. Krishnamurthy and R. Taylor, *AIChE J.*, 1985, **31**, 449–455.
- 138 R. Krishnamurthy and R. Taylor, *AIChE J.*, 1985, **31**, 456–465.
- 139 R. E. Treybal, *Mass-Transfer Operations*, McGraw-Hill, 3rd edn, 1981.
- 140 E. N. Fuller and J. C. Giddings, *J. Chromatogr. Sci.*, 1965, **3**, 222–227.
- 141 E. N. Fuller, P. D. Schettler and J. C. Giddings, *Ind. Eng. Chem.*, 1966, **58**, 18–27.
- 142 E. N. Fuller, K. Ensley and J. C. Giddings, *J. Phys. Chem.*, 1969, **73**, 3679–3685.
- 143 C. Wilke and P. Chang, *AIChE J.*, 1955, **1**, 264–270.
- 144 M. Takahashi, Y. Kobayashi and H. Takeuchi, *J. Chem. Eng. Data*, 1982, **27**, 328–331.
- 145 B. Han, Y. Sun, M. Fan and H. Cheng, *J. Phys. Chem. B*, 2013, **117**, 5971–5977.
- 146 L. R. Perkins and C. J. Geankoplis, *Chem. Eng. Sci.*, 1969, **24**, 1035–1042.
- 147 Process Systems Enterprise, gPROMS v4.0, 1997–2015, <http://www.psenderprise.com/gproms>.
- 148 N. M. Kakalis, A. I. Kakhu and C. C. Pantelides, *Proc. 6th International Conf. on Foundations of Computer Aided Process Design*, CACHE Publications, 2004, pp. 537–40.
- 149 P. Tontiwachwuthikul, A. Meisen and C. J. Lim, *Can. J. Chem. Eng.*, 1989, **67**, 602–607.
- 150 ASHRAE, *2011 ASHRAE Handbook*, ASHRAE, 2011.
- 151 J. A. Rocha, J. L. Bravo and J. R. Fair, *Ind. Eng. Chem. Res.*, 1993, **32**, 641–651.
- 152 G. Vazquez, E. Alvarez, J. M. Navaza, R. Rendo and E. Romero, *J. Chem. Eng. Data*, 1997, **42**, 57–59.
- 153 S. A. Jayarathna, A. Weerasooriya, S. Dayarathna, D. A. Eimer and M. C. Melaen, *J. Chem. Eng. Data*, 2013, **58**, 986–992.



- 154 G. J. Gloor, G. Jackson, F. J. Blas, E. M. del Río and E. de Miguel, *J. Chem. Phys.*, 2004, **121**, 12740–12759.
- 155 G. J. Gloor, G. Jackson, F. J. Blas, E. M. del Río and E. de Miguel, *J. Phys. Chem. C*, 2007, **111**, 15513–15522.
- 156 F. Llorell, A. Galindo, F. J. Blas and G. Jackson, *J. Chem. Phys.*, 2010, **133**, 024704.
- 157 H. M. Kvamsdal, J. Hetland, G. Haugen, H. F. Svendsen, F. Major, V. Kårstad and G. Tjellander, *Int. J. Greenhouse Gas Control*, 2010, **4**, 613–622.
- 158 C. V. Brand, J. Rodriguez, A. Galindo, G. Jackson and C. S. Adjiman, *Energy Procedia*, 2013, **37**, 1566–1571.
- 159 F. Y. Jou, A. E. Mather and F. D. Otto, *Can. J. Chem. Eng.*, 1995, **73**, 140–147.
- 160 S. Westmeier, *Chem. Tech. (Leipzig, Ger.)*, 1977, **29**, 218–222.
- 161 A. G. Leibush and E. D. Shorina, *Zh. Prikl. Khim.*, 1947, **20**, 69–76.
- 162 Y. Maham, C. N. Liew and A. E. Mather, *J. Solution Chem.*, 2002, **31**, 743–756.
- 163 J. Kestin, Y. Kobayashi and R. T. Wood, *Physica*, 1966, **32**, 1065–1089.
- 164 W. N. Haynes, *Handbook of Chemistry and Physics*, CRC Press, 2011.
- 165 M. Laliberte, *J. Chem. Eng. Data*, 2009, **54**, 1725–1760.
- 166 M. Mundhwa and A. Henni, *J. Chem. Eng. Data*, 2007, **52**, 491–498.
- 167 J. A. Riddick, W. B. Bunger and T. K. Sakano, *Organic Solvents*, Wiley, New York, 4th edn, 1986.
- 168 N. B. Vargaftik, *Moskva*, 1972, 433–434.
- 169 R. Bender, K. Bier and G. Maurer, *Ber. Bunsen-Ges. Phys. Chem.*, 1981, **85**, 778–784.
- 170 R. H. Perry and D. W. Green, *Perry's Chemical Engineers' Handbook*, McGraw Hill, 6th edn, 2008.

