Screening metal-organic frameworks for mixture separations in fixed-bed adsorbers using a combined selectivity/capacity metric

Krishna, R.

DOI
10.1039/c7ra07363a

Publication date
2017

Document Version
Final published version

Published in
RSC Advances

License
CC BY

Citation for published version (APA):
Krishna, R. (2017). Screening metal-organic frameworks for mixture separations in fixed-bed adsorbers using a combined selectivity/capacity metric. RSC Advances, 7(57), 35724-35737. https://doi.org/10.1039/c7ra07363a
Screening metal–organic frameworks for mixture separations in fixed-bed adsorbers using a combined selectivity/capacity metric†

Rajamani Krishna DOI: 10.1039/c7ra07363a

1. Introduction

Metal–organic frameworks (MOFs) have considerable potential for the separation of a wide variety of mixtures, offering energy-efficient alternatives to conventional distillation and absorption technologies. For any given separation task, a wide variety of MOFs are on offer; consequently, there is a need for a robust procedure for screening and ranking of MOFs. Particularly noteworthy are recent advances in the large-scale computational screening of MOFs for separation and storage applications, leading in some cases to the discovery of new materials. The potency of computational screening has been demonstrated for CO$_2$ capture, NH$_3$ capture, separation of hexane and heptane isomers, and Xe/Kr mixtures.

Two commonly used metrics for screening are selectivity and capacity. For separation of a binary mixture of components A and B, the adsorption selectivity, $S_{A/B}$, is defined by

$$S_{A/B} = \frac{q_A}{q_B} \frac{y_A}{y_B}$$

where the $q_A$ and $q_B$ represent the molar loadings within the MOF that is in equilibrium with a bulk gas phase mixture with mole fractions $y_A$ and $y_B = 1 - y_A$. The molar loadings, also called gravimetric uptake capacities, are usually expressed with the units mol kg$^{-1}$. The volumetric uptake capacities, are

$$Q_A = \rho q_A; \quad Q_B = \rho q_B$$

where $\rho$ is the crystal framework density of the MOF, expressed say in units of kg m$^{-3}$, or kg L$^{-1}$. The uptake capacities can be calculated using pure components isotherm fits, along with the mixed-gas Langmuir model or the Ideal Adsorbed Solution Theory (IAST) of Myers and Prausnitz for adsorption equilibrium. The screening of MOFs using these two metrics seldom lead to unambiguous choices, as illustrated below for four different mixture separations.
For 20/80 Xe/Kr mixture separations, Fig. 1a shows that the hierarchy of $S_{A/B}$ values is Co-formate > Ag@NiMOF-74 > NiMOF-74 > SBMOF-2 > SAPO-34 > CuBTC. The calculations in Fig. 1 are based on the IAST using the experimental data on the unary isotherms; all calculation details are provided in the ESI,† that also includes the structural details for each of the MOFs mentioned and analyzed in this article. Wang et al.$^{11}$ attribute the highly selective adsorption of Xe in Co-formate to commensurate positioning of Xe within the cages; such commensurate positioning also implies that the intracrystalline diffusivity of Xe will be significantly lower than that of Kr.$^{12-14}$ The introduction of well-dispersed Ag nanoparticles into NiMOF-74, results in stronger interactions of the polarizable Xe with the adsorbent Ag@NiMOF-74. The geometry of the channels of SBMOF-2 provides a better match for the larger Xe atoms rather than smaller Kr, and helps explain the selectivity for Xe over Kr. The hierarchy of Xe uptake capacities is Ag@NiMOF-74 > Co-formate > NiMOF-74 > SBMOF-2 > CuBTC > SAPO-34; this does not go hand-in-hand with the $S_{ads}$ hierarchy.

For 50/50 C$_2$H$_2$/$C_2$H$_4$ mixtures, the $S_{A/B}$ vs. C$_2$H$_2$ uptake capacity plots are shown in Fig. 1b. On the basis of selectivity, the best MOF is SIFSIX-2-Cu/i (2 = 4,4'-dipyridylacetylene, i = interpenetrated); in this case, each C$_2$H$_4$ molecule is bound by two F atoms from different nets.$^{15}$ However, SIFSIX-1-Cu, with the larger pore size, has the higher C$_2$H$_4$ uptake capacity. For separations of 50/50 C$_2$H$_2$/$C_2$H$_6$ mixtures, the highest $S_{ads}$ values are realized with NOTT-300, and PAF-1-SO$_3$Ag; see Fig. 1c. Simultaneous and cooperative hydrogen-bonding, and π⋯π stacking interactions account for the stronger binding of C$_2$H$_4$ in NOTT-300. The π-complexation of the alkenes with Ag(i) ions of PAF-1-SO$_3$Ag account for its high alkene/alkane selectivity. The C$_2$H$_4$ uptakes of NOTT-300 and PAF-1-SO$_3$Ag are, however, significantly lower than that of FeMOF-74, and CoMOF-74; see Fig. 1c.

A MOF that has been customized for C$_3$H$_6$/$C_3$H$_8$ separations is NbOFFIVE-1-Ni (=KAUST-7), belonging to the same class of SIFSIX materials but using pyrazine as the organic linker;$^{16}$ it exhibits the highest selectivity (see Fig. 1d). The tilting of the pyrazine molecule on the linker reduces the pore aperture to about 3 Å, permitting ingress of C$_3$H$_6$ molecules, but essentially excluding C$_3$H$_8$ on the basis of subtle differences in bond lengths and bond angles (see Fig. S19†). On the other hand,
NiMOF-74, with 11 Å hexagonal-shaped channels, has the highest uptake capacity for the unsaturated alkene, but a significantly lower selectivity than KAUST-7.

In industrial processing operations, the MOFs are normally used in powder or pelletized form in fixed bed devices; see schematic in Fig. 2. Such units are most commonly operated in a transient mode, with adsorption and desorption cycles; they are called either pressure swing adsorbers (PSA) or temperature swing adsorbers (TSA) depending on the regeneration strategy employed. Most commonly, the separation performance in a fixed-bed adsorber is dictated by mixture adsorption equilibrium; both selectivity and capacity metrics are relevant and important. Other factors that influence separation performance include axial dispersion, intra-particle diffusional limitations, and resistance to diffusional transfer between the bulk gas phase and the surface of the particle.\textsuperscript{17–20} The dispersion and diffusional effects are strongly influenced by the particle size, interstitial fluid velocity, and equipment dimensions. Dispersion and diffusional effects often cause distended breakthrough characteristics and usually lead to diminished separation effectiveness.\textsuperscript{19,21}

Transient breakthrough experiments on laboratory scale fixed-bed equipment have been used for screening MOFs;\textsuperscript{16,22–24} however, this approach is labor intensive and time consuming. An alternative approach, that has gained currency in the literature, is to screen MOFs on the basis of transient breakthrough simulations;\textsuperscript{14,15,18,25–29} in these simulations it is customary to neglect dispersion and diffusional effects but such effects can be also be routinely included, as appropriate.\textsuperscript{21}

The primary objective of this article to develop a simple and robust screening strategy that obviates the need for performing transient breakthrough simulations, or transient breakthrough experiments on a laboratory scale. Towards this end, we define a “new” metric, dubbed the separation potential, $\Delta Q$, that combines the selectivity and capacity metrics in a manner that is truly representative of the separation characteristics of transient fixed bed operations. The value of $\Delta Q$ is calculated using the IAST for mixture adsorption equilibrium. For validation of the combined metric, comparisons are made with the separation performance determined from transient breakthrough simulations for several different mixtures: Xe/Kr, C\textsubscript{2}H\textsubscript{2}/CO\textsubscript{2}, C\textsubscript{2}H\textsubscript{2}/C\textsubscript{2}H\textsubscript{4}, C\textsubscript{2}H\textsubscript{2}/C\textsubscript{6}H\textsubscript{6}, CO\textsubscript{2}/CH\textsubscript{4}, CO\textsubscript{2}/N\textsubscript{2}, CO\textsubscript{2}/H\textsubscript{2}, CO\textsubscript{2}/CO/CH\textsubscript{4}/H\textsubscript{2}, and hydrocarbon isomers using a wide variety of MOFs.

The ESL,\textsuperscript{+} accompanying this publication, provides (a) methodology used for transient breakthrough simulations, (b) analytic solutions to the shock wave model for fixed bed transient operations, (c) structural information on the MOFs investigated, (d) unary isotherm data for each guest/host combination, and (e) details simulation results for each of the investigated systems.

## 2. The separation potential of fixed bed adsorber

For quantifying the separation performance, we neglect axial dispersion effects and consider plug flow of an $n$-component gas mixture through a fixed bed adsorber maintained under isothermal, isobaric conditions; see Fig. 2. The molar concentrations of the constituent species in the bulk fluid phase vary with position $z$ along the bed, and time, $t$. At any position $z$ in the fixed bed, and time $t$, the molar loadings in the adsorbed phase within the pores, varies along the radius of the particle, $r$. The molar concentrations in the gas phase at any position and instant of time are obtained by solving the following set of partial differential equations for each of the species $i$ in the gas mixture.\textsuperscript{30}

\[
\frac{\partial c_i(t, z)}{\partial t} + \frac{\partial (\nu(t, z)c_i(t, z))}{\partial z} + \frac{(1 - \varepsilon)}{\varepsilon} \frac{\partial \bar{q}_i(t, z)}{\partial t} = 0; \quad i = 1, 2, \ldots, n
\]

In eqn (3), $t$ is the time, $z$ is the distance along the adsorber, $\rho$ is the framework density, $\varepsilon$ is the bed voidage, $\nu$ is the interstitial gas velocity, and $\bar{q}_i(t, z)$ is the spatially averaged molar loading within the crystallites of radius $r_c$, monitored at position $z$, and at time $t$. The time $t = 0$, corresponds to the time at which the feed mixture is injected at the inlet to the fixed bed. Prior to injection of the feed, it is assumed that an inert, non-adsorbing, gas flows through the fixed bed.

At any time $t$, during the transient approach to thermodynamic equilibrium, the spatially averaged molar loading within the crystallite $r_c$ is obtained by integration of the radial loading profile

\[
\bar{q}_i(t) = \frac{3}{r_c^2} \int_0^{r_c} q_i(r, t) r^2 dr
\]

For transient unary uptake within a crystal at any position and time with the fixed bed, the radial distribution of molar loadings, $q_i$, within a spherical crystallite, of radius $r_c$, is obtained from a solution of a set of differential equations describing the uptake

![Fig. 2 Schematic of fixed bed adsorber filled with adsorbent particles of radius $r_c$.](image-url)
The molar flux $N_i$ of component $i$ may be described by the simplified version of the Maxwell–Stefan equations in which both correlation effects and thermodynamic coupling effects are considered to be of negligible importance:

$$N_i = -\rho D_i \frac{\partial q_i}{\partial r}$$

(6)

Summing eqn (4) over all $n$ species in the mixture allows calculation of the total average molar loading of the mixture within the crystallite

$$\bar{q}_i(t, z) = \sum_{i=1}^{n} q_i(t, z)$$

(7)

The interstitial gas velocity is related to the superficial gas velocity by

$$v = \frac{\mu}{\epsilon}$$

(8)

In developing the concept of the separation potential, we assume that the value of $\frac{D_i}{r_c^2}$ is large enough to ensure that intra-crystalline gradients are absent and the entire crystallite particle can be considered to be in thermodynamic equilibrium with the surrounding bulk gas phase at that time $t$, and position $z$ of the adsorber

$$q_i(t, z) = q_i(t, z)$$

(9)

The molar loadings $q_i(t, z)$ are calculated using the IAST for mixture adsorption equilibrium using the bulk gas phase partial pressures $p_i$ at that position $z$ and time $t$. It should be noted that the applicability of the IAST is restricted to cases in which there is a homogenous distribution of adsorbate species throughout the microporous framework. The IAST fails for cases in which there is strong hydrogen bonding between the adsorbates, as is the case for water/alcohol, and alcohol/aromatic mixtures. The IAST predictions will also fail when strong segregation effects are present; examples of segregated adsorption include preferential siting of CO$_2$ at the window regions, preferential location at the intersections of network of channels, or within pockets. Other conditions under which IAST calculations are not of adequate accuracy are discussed by Cessford et al.

The adsorber bed, that is operated under isothermal, isobaric conditions, is assumed to be initially free of adsorbates, i.e. we have the initial condition

$$t = 0; q_i(0, z) = 0$$

(10)

Eqn (10) is relevant to the operation of the transient breakthrough experiments on a laboratory scale, but are not truly reflective of industrial operations.

At time, $t = 0$, the inlet to the adsorber, $z = 0$, is subjected to a step input of the $n$-component gas mixture and this step input is maintained till the end of the adsorption cycle when steady-state conditions are reached.

$$t \geq 0; p_i(0,t) = p_{0i}; u(0,t) = u_0$$

(11)

where $u_0 = v_{sf}$ is the superficial gas velocity at the inlet to the adsorber.

As illustration, Fig. 3 presents results of transient breakthrough simulations for the concentrations, and loadings along the length for 20/80 Xe/Kr mixtures in a fixed bed packed with SBMOF-2, for visual appreciation of the traversal of concentration and loading “fronts” along the length of the fixed bed, video animations are available as ESI.† The traversal velocity for the more strongly adsorbed Xe is significantly lower than that of the poorly adsorbed Kr. We define the displacement time interval, $\Delta \tau = \tau_{Xe} - \tau_{Kr}$, as the difference between the breakthrough times of Xe and Kr; during this interval, pure Kr with $<1000$ ppm Xe can be recovered, as demanded by industrial process requirements. From a material balance on the adsorber, the productivity of pure Kr is determined to be 4.46 moles per L of adsorbent MOF that is loaded in the fixed bed. In this article, we consistently choose to express the productivities in volumetric terms because the comparison of different MOFs for the same separation task is on the basis of fixed bed units containing the same volume of MOF adsorbent particles.

It is noteworthy that the spatio-temporal development of concentrations and loadings have distended characteristics, even though axial dispersion and diffusional resistances are ignored in eqn (3); for more detailed discussion of breakthrough characteristics, see the monographs of Ruthven. Such distended characteristics result in diminished productivity of pure Kr. The degree of distention of the concentration fronts is dictated, inter alia, by the adsorption selectivity, $S_{A/B}$; higher selectivity values generally lead to sharper fronts.

In proceeding further, it is necessary to quantify the characteristics of an “ideal” fixed bed adsorber. If the concentration and loading “fronts” were to traverse the column as “shock waves”, the spatio-temporal distributions of gas phase mole fractions, for an A/B mixture, are described by

$$y_A(z, \tau) = y_{A0} - y_{A0} \Phi \left( \frac{z - \tau}{L} \frac{\tau}{\tau_A} \right)$$

$$y_B(z, \tau) = y_{B0} + (1 - y_{B0}) \Phi \left( \frac{z - \tau}{L} \frac{\tau}{\tau_B} \right)$$

(12)

where $\Phi(\zeta) = 1$, $\zeta \geq 0$; $\Phi(\zeta) = 0$, $\zeta < 0$ is the Heaviside function; $y_{A0}$ and $y_{B0} = 1 - y_{A0}$ are the mole fractions entering the bed; $\tau$ is the dimensionless time, $\tau = tu/Ls$; $\tau_A$ and $\tau_B$ are the dimensionless breakthrough times of A and B, with $\tau_A > \tau_B$. The shock wave solutions are indicated by the dotted lines in Fig. 3. The productivity of pure B recovered during the interval $\Delta \tau = \tau_A - \tau_B$ can be determined from an analytic solution to the shock wave model (detailed derivations are provided in the ESI†); the resulting expression is remarkably simple.
The mole fraction $y_B$ in eqn (13) refers to composition of the feed mixture. In this article, the evaluation of the separation performance using the $D_{QX}$ is consistently based on the feed mixture composition entering the fixed bed adsorber, that is initially free of adsorbates (cf. eqn (10)).

For 20/80 Xe/Kr separation with SBMOF-2, with $p_A = 20$ kPa, $p_B = 80$ kPa, IAST calculations yield $Q_A = 1.52$ mol L$^{-1}$, $Q_B = 0.66$ mol L$^{-1}$, $S_{ads} = 9.25$, $\Delta Q = 5.4$ mol L$^{-1}$, somewhat higher than the Kr productivity value of 4.46 mol L$^{-1}$, determined from breakthrough simulations. Analogous calculations for the other five MOFs considered in Fig. 1a, enable the construction of a plot of the productivity of pure Kr versus the values of $\Delta Q_{Kr}/Q_{Xe} = Q_{Kr}/Q_{Xe} - 0.33$, calculated from eqn (13); see Fig. 4a. The two sets of data are linearly inter-related, leading us to conclude that the use of the metric $\Delta Q_{Kr}/Q_{Xe}$ leads to the same hierarchy of Kr productivities as that obtained from transient breakthrough simulations.

Also plotted as right $y$-axis in Fig. 4a is the time interval $\Delta t = t_{Kr} - t_{Xe}$, that is also found to be linearly related to the Kr productivity in the fixed bed. For screening MOFs on the basis of experimental breakthroughs, the appropriate and convenient metric would be $\Delta t = t_{Kr} - t_{Xe}$; see Fig. S25 and S26.$^\dagger$

Fig. 4b compares the Xe capture capacity, determined from breakthrough simulations for the six different MOFs, with the IAST calculated values of Xe uptake $Q_{Xe}$, calculated from the IAST; the inter-dependence is precisely linear; the actual productivity values are slightly lowered below $Q_{Xe}$. Furthermore, the Xe capture capacity is also found to be linearly related to the breakthrough time for Xe, $t_{Xe}$ plotted as right $y$-axis in Fig. 4b. It is also interesting to note that the same hierarchy of separation performance is realized for the six MOFs, whether compared on the basis of Kr productivity or Xe capture capacity.$^{14}$

For CO$_2$ capture from flue gases and natural gas streams, the appropriate metric for screening and ranking MOFs is the CO$_2$
Furthermore, it is noteworthy that the hierarchy dictated by six MOFs is the same as that found in Fig. 4; see Fig. S13.

hierarchy of productivities of pure Xr (with <1000 ppm Kr) of the phase exiting the desorption bed. Breakthrough simulations for presents that productivity of pure A that can be recovered in the gas uptake capacity, Q_{CO2}. This is precisely equivalent to screening on the basis of the breakthrough time for CO2, \( \tau_{CO2} \), in view of the linear inter-dependence of \( \tau_{CO2} \) and Q_{CO2}. Indeed, Xiang et al.\textsuperscript{27} have used the metric \( \tau_{CO2} \) for screening MOFs and zeolites for CO2 capture applications.

The corresponding expression for the separation potential for the desorption cycle is \( \Delta Q_{A/B} = Q_A - y_A Q_B - y_B \) and represents that productivity of pure A that can be recovered in the gas phase exiting the desorption bed. Breakthrough simulations for the desorption cycle of 20/80 Xe/Kr separations show that the hierarchy of productivities of pure Xr (with <1000 ppm Kr) of the six MOFs is the same as that found in Fig. 4; see Fig. S13.\textsuperscript{†} Furthermore, it is noteworthy that the hierarchy dictated by \( \Delta Q_{A/B} \) does not follow that of \( S_{X/B} \), as can be verified by comparison of Fig. 4 with Fig. 1a.

Analogous results to those presented in Fig. 4a and b are obtained for separations of the five different binary mixtures: C2H2/CO2, n-pentane (nC5)/2-methylbutane (2MB), C2H4/C2H6, C2H4/C3H6, and C2H2/C2H4; the obtained results are summarized in Fig. 5a–f. In all cases, the productivity of the less-strongly adsorbed component in fixed beds containing identical volumes of different MOFs, is linearly related with separation potential, \( \Delta Q_{B/A} = Q_A y_B - Q_B \), determined from IAST. This linear relationship also holds for separation of 50/50 CO2/CH4, 15/85 CO2/N2, and 20/80 CO2/H2 mixtures; see the results presented in Fig. 6. Some important conclusions to be drawn from the results in Fig. 5 and 6 are discussed below.

For C2H2/C2H4 separations, the purity requirement of C2H4 is stringent (<40 ppm C2H2) and the MOF with the highest C2H4 productivity depends on feed mixture composition. For 50/50 mixture, the best performance is obtained with SIFSIX-1-Cu, that has the highest C2H2 uptake capacity (cf. Fig. 1b). The MOF with the highest selectivity, SIFSIX-2-Cu-i suffers from capacity limitations for 50/50 mixtures. For separation of 1/99 C2H2/C2H4 mixture separations, the best MOF is SIFSIX-2-Cu-i; in this case, capacity considerations are of significantly lesser importance. The separation potential \( \Delta Q_{A/B} \) appropriately combines the separate influences of uptake, selectivity, and mixture composition.

For C2H2/C2H4 separations, the highest C2H4 productivity is realized with FeMOF-74,\textsuperscript{43} even though this MOF has a significantly lower selectivity than NOTT-300 (cf. Fig. 1c). Corresponding simulations of the desorption phase (presented in Fig. S20)\textsuperscript{†} shows that FeMOF-74 also shows the highest productivity of 99.95% pure C2H4 product, that is demanded in industry as feedstock for polymerization reactors.

For 50/50 C2H6/C3H8 separations, KAUST-7, the MOF with the highest selectivity, ranks the lowest in the hierarchy of 99.95% C2H4 productivities. The highest C2H4 productivity is realized with NiMOF-74, that possesses the right balance between selectivity and uptake capacity. For 50/50 alkene/alkane separations, capacity considerations overcome any selectivity disadvantages associated with M-MOF-74 adsorbents.

One common underlying feature of all the results presented in Fig. 4–6 is that the actually productivity values are lower than the corresponding \( \Delta Q_{A/B} \) values; the reasons for the diminished performance are investigated in the following section.

3. Investigation of the relative influences of selectivity and uptake capacity

To gain deeper insights into the relative importance of selectivity and capacity metrics, we investigated the separation of binary A/B mixtures using 20 different “hypothetical” HypMOFs. The unary adsorption isotherms for each HypMOF is described by the 1-site Langmuir isotherm.
We assume the saturation capacity of both A and B to be identical to each other; this allows the use of the mixed-gas Langmuir model to calculate mixture adsorption equilibrium:

\[
q = \frac{q_{\text{sat}} b_p}{1 + b_p p}
\]

(14) \hspace{1cm} \frac{q_A}{q_{\text{sat}}} = \frac{b_{\text{A}p_A}}{1 + b_{\text{A}p_A} + b_{\text{B}p_B}} \quad ; \quad \frac{q_B}{q_{\text{sat}}} = \frac{b_{\text{B}p_B}}{1 + b_{\text{A}p_A} + b_{\text{B}p_B}}

(15)

Twenty different “hypothetical” HypMOFs are “constructed” by choice of the three parameters \(q_{\text{sat}}, b_A, b_B\); ten of the HypMOFs have \(q_{\text{sat}} = 2.5 \text{ mol kg}^{-1}\); for the other ten MOFs,
The values of \( b_A, b_B \) are chosen to realize selectivities of 1.5, 2, 4, 6, 8, 10, 20, 10^2, 10^3, and 10^4. In all cases, the Langmuir parameters are chosen such that species A is more strongly adsorbed.

Transient breakthrough simulations for equimolar (i.e., \( y_A = 0.5 = y_B = 1 - y_A \)) A/B mixtures in a fixed bed packed with twenty different HypMOFs were undertaken. As illustration, Fig. 7a shows transient breakthrough simulations for HypMOF-9 and HypMOF-11; HypMOF-9 has a saturation capacity \( q_{\text{sat}} = 2.5 \) mol kg\(^{-1}\) and selectivity \( S_{A/B} = b_A/b_B = 1000. \) HypMOF-11 has a higher saturation capacity, \( q_{\text{sat}} = 5 \) mol kg\(^{-1}\), but a significantly lower selectivity \( S_{A/B} = b_A/b_B = 1.5. \) For both HypMOFs, the more strongly adsorbed A is the component that elutes last; the less strongly adsorbed B breaks through earlier. For HypMOF-9, with the higher selectivity, the component B breaks through practically at the start of the breakthrough “experiment”. We demand a purity of 99.95% B in the product gas; there is a finite time interval, \( \Delta t \), during which pure 99.95% pure B can be recovered. The larger value of \( \Delta t \), the higher is the productivity of pure B.

Fig. 7b presents a plot of the number of moles of B produced per liter of HypMOF in the packed bed, as a function of the selectivity \( S_{A/B} \); higher productivities are realized with MOFs having the higher saturation capacity. Increase of selectivities above 100 does not lead to a concomitant increase in the amount of pure B that can be recovered; this implies that the total exclusion of any component, i.e., “infinite” selectivities, may not be the fruitful strategy to adopt for MOF development. To appreciate the influence of selectivity of the productivities, we combine eqn (1) and (13) to obtain

\[
\Delta Q_{B/A} = Q_A \frac{y_B}{1 - y_B} \left( \frac{1}{S_{A/B}} \right)
\]

If \( S_{A/B} \gg 1, \) \( \Delta Q_{B/A} = Q_A \frac{y_B}{1 - y_B} \), and the productivity of pure B is not influenced by selectivity but is purely dictated by the uptake capacity of the more strongly adsorbed component A. Conversely, for “difficult” separations, with values of \( 1 < S_{A/B} < 10 \), there is a sharp increase in productivities with increased \( S_{A/B} \).

In consonance with the results in Fig. 4, and 5, Fig. 7c shows that the productivity of pure B is linearly correlated with the separation potential \( \Delta Q_{B/A}. \) The dotted line is the parity line, underscoring the fact that the actual productivities are lower than \( \Delta Q_{B/A}. \) The % deviation of the actual productivities from \( \Delta Q_{B/A} \) are plotted in Fig. 7d; the higher the selectivity the
smaller is the deviation from $\Delta Q_{BA}$. The reasoning is that higher selectivities lead to sharper concentration fronts, and closer approach to the shock wave “idealization” of breakthroughs. For a selectivity value of $10^4$, there is only a 10% deviation from $\Delta Q_{BA}$. Put another way, high values of selectivity allows the separation potential to be reached more closely, but the potential itself is largely influenced by saturation capacity and component uptakes.

The foregoing discussions help us appreciate why NOTT-300 and KAUST-7 do not exhibit high productivities in alkene/alkane separations in fixed bed units (cf. Fig. 5e and f), despite possessing high selectivities: both these MOFs suffer severe capacity limitations.

4. The separation potential for multicomponent mixtures

Many industrially processes involve separations of mixtures containing more than two components, and the shock wave model can be readily extended to $n$-component mixtures, as detailed in the ESL† For separation of a quaternary A/B/C/D mixture in which component D, with the lowest adsorption strength, needs to be recovered in pure form from the exiting gas stream from the fixed bed, the expression for the separation potential for D/CBA separation is

$$\Delta Q_{D/CBA} = (Q_A + Q_B + Q_C) \frac{y_D}{1-y_D} - Q_D$$

(17)

For recovery of pure E from a 5-component A/B/C/D/E mixture, expression for the separation potential is an intuitive extrapolation of eqn (17):

$$\Delta Q_{E/D/CBA} = (Q_A + Q_B + Q_C + Q_D) \frac{y_E}{1-y_E} - Q_E$$

(18)

In the event the desired product exiting the fixed bed is the combination (E + D), the appropriate separation potential for the ED/CBA “cut” is
\[ \Delta Q_{ED/CHA} = (Q_a + Q_b + Q_c) \begin{bmatrix} y_D + y_E \\ 1 - y_D - y_E \end{bmatrix} - (Q_D + Q_E) \]  \hspace{1cm} (19) 

The use of eqn (17)-(19) will be illustrated for H\textsubscript{2} purification and the separation of hydrocarbon isomers.

Pressure swing adsorption is the most commonly used technology for recovery of pure hydrogen from a gaseous streams containing 60–90 mol% H\textsubscript{2}.\textsuperscript{44} Steam-methane reformer off-gas, after it has been further treated in a water–gas shift reactor, is a commonly used feed gas stream, with typical compositions 70–80% H\textsubscript{2}, 15–25% CO\textsubscript{2}, 3–6% CH\textsubscript{4}, and 1–3% CO.\textsuperscript{28,44,45} The production of pure H\textsubscript{2} from steam-methane reformer off-gas requires operation of PSA units at pressures reaching about 7 MPa. Typically, H\textsubscript{2} is required with 99.95% + purity.\textsuperscript{44} There is no unique and unambiguous definition of adsorption selectivity that is appropriate for H\textsubscript{2} purification processes. As illustration, Fig. 8a shows the component mole fractions in the exit gas from a fixed bed packed with CuTDPAT, fed with 40/5/50 CO\textsubscript{2}/CO/CH\textsubscript{4}/H\textsubscript{2} mixtures at 298 K and 6 MPa.\textsuperscript{46} The sequence of breakthroughs is H\textsubscript{2}, CH\textsubscript{4}, CO, and CO\textsubscript{2}. The purity of H\textsubscript{2} in the product gas exiting the adsorber is primarily dictated by the presence of CO, and CH\textsubscript{4}, rather than CO\textsubscript{2}, that is the main impurity in the feed gas mixture. The CO/H\textsubscript{2} and CH\textsubscript{4}/H\textsubscript{2} adsorption selectivities are far more relevant than the CO\textsubscript{2}/H\textsubscript{2} selectivity. There is a finite time interval during which pure H\textsubscript{2}, of the required 99.95% + purity, can be recovered from the gases exiting the fixed bed.

Fig. 8b presents a comparison of productivities of four different adsorbents, plotted as a function of the separation potential, \( \Delta Q_{H_2/(CO+CH_4+CO_2)} \) from IAST calculations of eqn (17); the linear inter-dependence con-dered from the gases exiting the fixed bed packed with CuTDPAT, fed with 40/5/50 CO\textsubscript{2}/CO/CH\textsubscript{4}/H\textsubscript{2} mixtures at 298 K and 6 MPa.\textsuperscript{46} The sequence of breakthroughs is H\textsubscript{2}, CH\textsubscript{4}, CO, and CO\textsubscript{2}. The purity of H\textsubscript{2} in the product gas exiting the adsorber is primarily dictated by the presence of CO, and CH\textsubscript{4}, rather than CO\textsubscript{2}, that is the main impurity in the feed gas mixture. The CO/H\textsubscript{2} and CH\textsubscript{4}/H\textsubscript{2} adsorption selectivities are far more relevant than the CO\textsubscript{2}/H\textsubscript{2} selectivity. There is a finite time interval during which pure H\textsubscript{2}, of the required 99.95% + purity, can be recovered from the gases exiting the fixed bed.

The separation of hexane isomers, \( n \)-hexane (\( n \)C6), 2-methylpentane (2MP), 3-methylpentane (3MP), 2,2 dimethylbutane (22DMB), and 2,3 dimethylbutane (23DMB) is required for production of high-octane gasoline. The values of the Research Octane Number (RON) increases with the degree of branching; the RON values are: \( n \)C6 = 30, 2MP = 74.5, 3MP = 75.5, 22DMB = 94, 23DMB = 105. Therefore, di-branched isomers are preferred products for incorporation into the high-octane gasoline pool.\textsuperscript{21,47-48} Typically, we aim to product a product with a RON value higher than 92. As illustration, Fig. 9a shows the transient development of exit gas compositions from a bed packed with MgMOF-74 with an equimolar 5-component feed mixture. Product gas with 92+ RON can be recovered from the displacement interval in which the di-branched isomers 22DMB and 23DMB are eluted, as indicated. When the mono-branched isomers break through, there is a sharp decrease in the RON of the product gas. The desired separation is between (22DMB + 23DMB), and (\( n \)C6 + 2MP + 3MP). The appropriate expression for the separation potential is

\[ \Delta Q_{(22DMB+23DMB)/(nC6+2MP+3MP)} = (Q_{nC6} + Q_{2MP} + Q_{3MP}) \begin{bmatrix} y_{22MB} + y_{23MB} \\ 1 - y_{22MB} - y_{23MB} \end{bmatrix} - (Q_{22DMB} + Q_{23DMB}) \]  \hspace{1cm} (20) 

Fig. 9b compares 92+ RON productivities for nine different adsorbent materials: ZnMOF-74, MgMOF-74, Co(BDP), Fe\textsubscript{2}(BDP)\textsubscript{3}, MFI, BEA, ZIF-8, Zn(bdc)\textsubscript{3}dabco, and ZIF-77, plotted as a function of the separation potential \( \Delta Q_{(22DMB+23DMB)/(nC6+2MP+3MP)} \); the interdependence is linear and we conclude that the best performing MOF for this separation duty is Fe\textsubscript{2}(BDP)\textsubscript{3}; this conclusion is in line with the earlier work of Herm et al.\textsuperscript{49}
The production of \( p \)-xylene involves the separation of 4-component equimolar \( o\)-xylene(1)/\( m\)-xylene(2)/\( p\)-xylene(3)/ethylbenzene(4) mixtures. The volumetric separation potential for preferential adsorption of \( p\)-xylene, and rejection of \( o\)-xylene, \( m\)-xylene, and ethylbenzene is

\[
\Delta Q_{(O\text{-x}+M\text{-x}+\text{EthBz})/p\text{-x}} = \frac{(Q_{O\text{-x}} + Q_{M\text{-x}} + Q_{\text{EthBz}})(1 - y_{O\text{-x}} - y_{M\text{-x}} - y_{\text{EthBz}}) - (Q_{O\text{-x}} + Q_{M\text{-x}} + Q_{\text{EthBz}})}{\rho_i \Delta \gamma_{p\text{-x}}}
\]

(21)

Screening adsorbents using the combined metric \( \Delta Q_{(O\text{-x}+M\text{-x}+\text{EthBz})/p\text{-x}} \) shows that the best separation performance is achieved with DynaMOF-100 that represents a Zn(II)-based dynamic coordination framework, \([\text{Zn}_4\text{O}(\text{L})_3]\) where the ligand \( \text{L} = 4,4'-(1,2\text{-phenylene})\text{bis}(\text{oxy})\text{dibenzoate} \), whose synthesis has been reported by Mukherjee et al.52,53 The framework of DynaMOF-100 gets transformed in such a manner as to allow optimal packing of \( p\)-xylene within the cavities. The superiority of DynaMOF-100 over the commercially used \( \text{BaX} \) zeolite needs to be established by further experimental investigations.

5. Limitations of the concept of separation potential

The expression developed for the combined metric \( \Delta Q \) is applicable to fixed beds that are initially free of adsorbates; consequently, the application of this metric is restricted to screening purposes, and not to process modelling. Another key assumption in deriving the simple expression for \( \Delta Q \) is that intra-crystalline gradients are absent and the entire crystallite particle can be considered to be in thermodynamic equilibrium with the surrounding bulk gas phase at that time \( t \) and position \( z \) of the adsorber; see eqn (9). The validity of this assumption depends on the value of the parameter \( \frac{D_i}{R_e^2} \). For low values of diffusivities, and/or large sizes of crystals, the influence of intra-crystalline diffusion cannot be ignored. Herm et al.49 have screened MOFs for separation of hexane isomers by detailed transient breakthrough simulations including intra-crystalline diffusion effects; they demonstrate that inclusion of such effects alters the separation hierarchy.

There are instances of diffusion-selective operations in which diffusional effects over-ride the influence of mixture
adsorption equilibrium and is the prime driver for separations. Examples of diffusion-selective separations include: (1) selective uptake of \( \text{N}_2 \) from \( \text{N}_2/\text{CH}_4 \) mixtures using LTA-4A zeolite and Ba-ETS-4, \( 57-59 \) and (2) selective uptake of \( \text{O}_2 \) from \( \text{O}_2/\text{N}_2 \) mixtures using LTA-4A zeolite and CMS. \( 55,60-63 \) The concept of separation potential cannot be directly applied to screen MOFs for diffusion-selective separations; a completely different approach is required.

Another important limitation of screening MOFs on the basis of \( \Delta Q \) is that this metric does not take account of the regeneration costs, that are dictated largely by the isosteric heat of adsorption, \( Q_{\text{st}} \), of the more strongly adsorbed component. The extension of the expression for \( \Delta Q \) to include the \( Q_{\text{st}} \) parameter is a fruitful area for further research.

### 6. Conclusions

A comparative evaluation of MOFs for separating mixtures of \( \text{Xe/Kr} \), \( \text{C}_3\text{H}_8/\text{CO}_2 \), \( \text{C}_2\text{H}_6/\text{C}_2\text{H}_4 \), \( \text{C}_2\text{H}_6/\text{C}_2\text{H}_4\text{H}_2 \), \( \text{C}_2\text{H}_6/\text{C}_2\text{H}_6\text{H}_2 \), \( \text{CO}_2/\text{CH}_4 \), \( \text{CO}_2/\text{N}_2 \), \( \text{CO}_2/\text{H}_2 \), \( \text{CO}_2/\text{CO/CH}_4/\text{H}_2 \), and hydrocarbon isomers in fixed beds has been carried out using a combination of transient breakthrough simulations, and the IAST for mixture adsorption equilibrium. The following major conclusions emerge from this study.

1. The separation performance in fixed bed devices is governed by a combination of adsorption selectivity, and uptake capacity. Increase of selectivity values above 100, may not lead to a corresponding increase in productivity.

2. Low uptake capacities diminish the separation performance of MOFs with high selectivities. A good illustration is KAUST-7 that has the highest selectivity for \( \text{C}_3\text{H}_6/\text{C}_3\text{H}_8 \) separations but is severely capacity limited; consequently, the capability of this MOF to produce 99.95% pure product is the lowest amongst the investigated adsorbent materials.

3. The separation potential \( \Delta Q \), that is calculable on the basis of IAST provides a simple and convenient metric to screen and rank the separation capability of MOFs. The value of \( \Delta Q \) defines the upper limit to the achievable separations in fixed bed units. The actual separations in fixed bed adsorbers will be lower than the IAST-calculated \( \Delta Q \) values because of distended breakthroughs. Use of \( \Delta Q \) for screening purposes obviates the need for performing transient breakthrough calculations.

4. The concept of \( \Delta Q \) is particularly advantageous for multicomponent separations because it combines several selectivities and uptake capacities into one combined metric that quantifies the desired separation task in hand.

5. The concept of the separation potential can be straightforwardly incorporated into large-scale computational screening of MOFs. \( 1-3 \)

An important limitation of the treatment presented in this work is that the combined metric \( \Delta Q \) describes the productivity for conditions in which diffusional limitations are absent. When diffusional effects are of significant importance, the simple concept of separation potential is inadequate for gauging separation performance, and detailed simulations including diffusional effects are necessary. \( 19,21 \)

In using \( \Delta Q \) for screening, trace impurities, such as water vapor, that may be present in the mixtures have been ignored; the limitations of this assumption have been detailed by Burtch et al. \( 64 \) Furthermore, other aspects such as manufacturing costs for the adsorbents and regeneration costs need to be taken into consideration in the evaluation of MOFs.

### Notation

- \( b \) Langmuir constant, \( \text{Pa}^{-1} \)
- \( c_i \) Molar concentration of species \( i \) in gas mixture, \( \text{mol m}^{-3} \)
- \( c_{i0} \) Molar concentration of species \( i \) in gas mixture at inlet to adsorber, \( \text{mol m}^{-3} \)
- \( D_i \) Maxwell– Stefan diffusivity, \( \text{m}^2 \text{s}^{-1} \)
- \( L \) Length of packed bed adsorber, \( \text{m} \)
- \( n \) Number of species in the mixture, dimensionless
- \( N_i \) Molar flux of species \( i \), \( \text{mol m}^{-2} \text{s}^{-1} \)
- \( p_i \) Partial pressure of species \( i \) in mixture, \( \text{Pa} \)
- \( p_t \) Total system pressure, \( \text{Pa} \)
- \( q_i \) Molar loading of species \( i \), \( \text{mol kg}^{-1} \)
- \( q_{i, \text{sat}} \) Molar loading of species \( i \) at saturation, \( \text{mol kg}^{-1} \)
- \( Q \) Volumetric loading in adsorbed phase, \( \text{mol m}^{-3} \)
- \( \Delta Q \) Separation potential, \( \text{mol m}^{-3} \)
- \( Q_{\text{st}} \) Isosteric heat of adsorption, \( \text{J mol}^{-1} \)
- \( r \) Radial coordinate, \( \text{m} \)
- \( r_c \) Radius of crystallite, \( \text{m} \)
- \( S_{\text{A/B}} \) Adsorption selectivity, dimensionless
- \( t \) Time, \( \text{s} \)
- \( T \) Absolute temperature, \( \text{K} \)
- \( u \) Superficial gas velocity in packed bed, \( \text{m s}^{-1} \)
- \( v \) Interstitial gas velocity in packed bed, \( \text{m s}^{-1} \)
- \( y \) Gas phase mole fraction, dimensionless
- \( z \) Distance along the adsorber, \( \text{m} \)

### Greek letters

- \( \varepsilon \) Voidage of packed bed, dimensionless
- \( \rho \) Framework density, \( \text{kg m}^{-3} \)
- \( \tau \) Time, dimensionless

### Subscripts

- \( A \) Referring to component \( A \)
- \( B \) Referring to component \( B \)
- \( i \) Referring to component \( i \)
- \( t \) Referring to total mixture

### References

1. Y. J. Colón and R. Q. Snurr, *Chem. Soc. Rev.*, 2014, **43**, 5735–5749.
2. C. E. Wilmer, M. Leaf, C. Y. Lee, O. K. Farka, B. G. Hauser, J. T. Hupp and R. Q. Snurr, *Nat. Chem.*, 2012, **4**, 83–89.
56 J. Kärger, D. M. Ruthven and D. N. Theodorou, *Diffusion in Nanoporous Materials*, Wiley - VCH, Weinheim, 2012.

57 H. W. Habgood, *Can. J. Chem.*, 1958, 36, 1384–1397.

58 S. J. Bhadra and S. Farooq, *Ind. Eng. Chem. Res.*, 2011, 50, 14030–14045.

59 B. Majumdar, S. J. Bhadra, R. P. Marathe and S. Farooq, *Ind. Eng. Chem. Res.*, 2011, 50, 3021–3034.

60 S. Farooq, M. N. Rathor and K. Hidajat, *Chem. Eng. Sci.*, 1993, 48, 4129–4141.

61 S. Farooq, *Gas Sep. Purif.*, 1995, 9, 205–212.

62 Y. D. Chen, R. T. Yang and P. Uawithya, *AIChE J.*, 1994, 40, 577–585.

63 S. Sircar and A. L. Myers, in *Handbook of Zeolite Science and Technology*, ed. S. M. Auerbach, K. A. Carrado and P. K. Dutta, Marcel Dekker, New York, 2003, pp. 1063–1104.

64 N. C. Burtch, H. Jasuja and K. S. Walton, *Chem. Rev.*, 2014, 114, 10575–10612.