

Oxygen surface exchange kinetics of erbia-stabilized bismuth oxide

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Abstract The surface oxygen exchange kinetics of bismuth oxide stabilized with 25 mol% erbia (BE25) has been studied in the temperature and pO_2 ranges 773–1,023 K and 0.1–0.95 atm, respectively, using pulse-response ^{18}O – ^{16}O isotope exchange measurements. The results indicate that BE25 exhibits a comparatively high exchange rate, which is rate determined by the dissociative adsorption of oxygen. Defect chemical considerations and the observed $pO_2^{1/2}$ dependence of the rate of dissociative oxygen adsorption suggest electron transfer to intermediate superoxide ions as the rate determining step in surface oxygen exchange on BE25.

Keywords Isotopic exchange · Surface exchange kinetics · Solid electrolyte · Erbia-stabilized bismuth oxide

Introduction

Stabilized phases of δ - Bi_2O_3 exhibit high oxide ion conductivity, which is related to a high polarizability of the bismuth ion and a high concentration of oxygen vacancies. In the parent cubic fluorite structure of δ - Bi_2O_3 ,

25% of the sites in the oxygen sublattice are vacant. Pure Bi_2O_3 , however, transforms from the δ -phase to the monoclinic α -phase upon cooling below $\sim 1,003$ K [1], showing a discontinuity in the ionic conductivity. The high temperature δ - Bi_2O_3 phase can be stabilized to room temperature by the addition of rare earth oxides [2–6]. Erbia-stabilized bismuth oxide, $(Bi_2O_3)_{0.75}-(Er_2O_3)_{0.25}$, referred to as BE25, shows an ionic conductivity more than two orders of magnitude higher than that of conventional yttria-stabilized zirconia (YSZ), and has been investigated for its possible use as electrolyte for the low-temperature solid oxide fuel cell [5, 6].

Stabilization of δ - Bi_2O_3 with isovalent Er_2O_3 preserves its high vacancy concentration albeit that, in general, a higher Er_2O_3 concentration results in a lower conductivity. At low Er_2O_3 concentrations and when operating at temperatures in the range 973–873 K, the ionic conductivity is found to slowly decay with time due to transformation of the cubic phase into a rhombohedral phase. Kruidhof et al. [7] claimed that, when operating at 973 K, a dopant concentration of at least 27.5% Er_2O_3 is necessary to prevent phase separation. Jiang et al. [8] showed that even at a concentration of 40% Er_2O_3 the conductivity of phase-stabilized δ - Bi_2O_3 is subject to aging due to gradual ordering of the anion sublattice when operating below 873 K. In addition to phase and structural instability, the potential application of bismuth-containing electrolytes in solid oxide fuel cells (SOFCs) is regarded to be limited. The poor redox stability of the bismuth ion easily leads to decomposition (to metallic bismuth) of the bismuth-containing electrolyte under the low oxygen partial pressures at the anode (fuel) side.

Compared with YSZ [9] and gadolinia-doped ceria [10], BE25 shows a remarkably high oxygen surface exchange rate. Current–voltage and impedance measurements of the properties of porous sputtered platinum and gold electrodes

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on BE25 show that the type of electrode material has little effect on the exchange current density, suggesting that for BE25 the entire electrolyte surface is active in the oxygen transfer reaction [11]. Reasonable agreement is found between the exchange current density from current–voltage measurements and the corresponding value calculated from data of ^{18}O – ^{16}O isotopic exchange measurements. The very high oxide ion diffusivity in BE25 relative to the rate of surface exchange is found to create rather flat diffusion profiles in tracer diffusion experiments. For BE25, a critical thickness of 1.7 cm at 773 K and 1 atm $p\text{O}_2$ was deduced by Steele et al. [12] and of 10.3 cm at 923 K at 0.2 atm $p\text{O}_2$ by Capoen et al. [13] from data of ^{18}O – ^{16}O isotopic exchange depth-profiling (IEDP) using secondary ion mass spectrometry (SIMS) analysis. The critical thickness, L_c , reflects the length scale below which the surface exchange is rate limiting, and in tracer diffusion experiments is given by $L_c = D^*/k_s$, where D^* is the tracer-diffusion coefficient and k_s the surface exchange coefficient [14]. The critical thickness at which oxygen diffusion and surface exchange determines equally the rate of oxygen permeation through BE25 was estimated by Bouwmeester et al. [15] to be 0.16 mm at 923 K and 0.09 mm at 1,083 K. Note that the critical thickness deduced from oxygen permeation data is two to three orders magnitude lower than that from the tracer diffusion experiments. While in tracer experiments, D^* governs the diffusion behavior, it is the ambipolar conductivity which controls the rate of oxygen permeation. The ambipolar conductivity in the solid electrolyte BE25 in usual ranges of oxygen partial pressure and temperature is determined by the partial p -type electronic conductivity, following a $p\text{O}_2^{1/4}$ power law dependence.

In this study, the surface exchange kinetics of BE25 is re-investigated using the pulse-response ^{18}O – ^{16}O isotope exchange (PIE) technique. The method recently developed in our laboratory is based upon isotope analysis of an ^{18}O -enriched gas phase pulse after passage of a continuous-flow packed-bed microreactor loaded with the oxide powder [16]. The measurements are carried out under isothermal and iso- $p\text{O}_2$ conditions. Since the method relies on gas phase analysis of the fractions of oxygen isotopomers with masses 36, 34, and 32 ($^{18}\text{O}_2$, $^{16}\text{O}^{18}\text{O}$, and $^{16}\text{O}_2$, respectively) also mechanistic information on the exchange reaction can be extracted from experiment

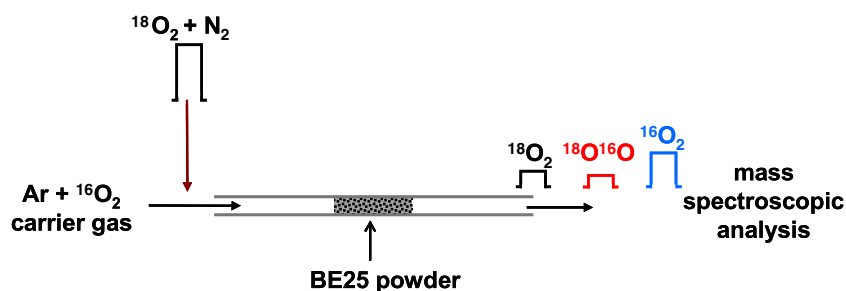
contrary to, for example, data obtained by means of the IEDP-SIMS method.

Experimental

BE25 powder prepared by a conventional solid state reaction was provided by the Fraunhofer-Institut für Keramische Technologien und Systeme, IKTS (Institutsteil Hermsdorf). The as-received powder was uniaxially pressed at 30 MPa into a disk, followed by cold isostatic pressing at 400 MPa. The disk was sintered at 1,123 K for 10 h in air, and crushed into a coarse powder. The fraction passing a 125 μm stainless steel sieve was pre-annealed at 1,073 K for 10 h in air, and sieved again to remove formed agglomerates. Prior to pulse-response isotope exchange measurements, the powders were characterized by X-ray diffraction (Philips pananalytical PW1830), BET surface area (Micromeritics ASAP 2020 M), and particle size (Malvern Mastersizer 2000) measurements.

The pulse-response isotope exchange measurements were performed in the temperature and $p\text{O}_2$ ranges 773–1,023 K and 0.1–0.95 atm, respectively, using a continuous flow packed-bed micro-reactor. During the measurement, approximately 0.08 g of BE25 powder was loaded between two quartz wool plugs in the center of a quartz tubular micro-reactor with inner diameter 2 mm. The length of the packed-bed was typically 13 mm. A $^{16}\text{O}_2/\text{Ar}$ gas mixture was used as carrier gas with a flow rate 50 ml/min (STP). The response to $^{18}\text{O}_2/\text{N}_2$ pulse (500 μl), with the same $p\text{O}_2$ as the carrier gas, passing through the reactor was analyzed by on-line mass spectrometry (Omni Star TM GSD 301 Pfeiffer-Vacuum) at the exit of the reactor as illustrated in Fig. 1. A six-port valve with sample loop was used for injection of the $^{18}\text{O}_2/\text{N}_2$ pulse into the $^{16}\text{O}_2/\text{Ar}$ carrier gas flow stream. Oxygen isotope gas was purchased from Cambridge Isotope Laboratory (>97 atom% $^{18}\text{O}_2$). Nitrogen used as diluent for the $^{18}\text{O}_2$ gas was also used for internal calibration of the mass spectrometer. The mean residence time of the reactor varied between 10 and 35 ms, being a function of flow rate and temperature. Prior to measurements, the BE25 powder packed-bed was pre-treated at 1,073 K for 0.5 h under flowing synthetic air in order to remove possibly adsorbed

Fig. 1 Schematic representation of the pulse-response isotope exchange technique



water and CO₂, and subsequently cooled to room temperature at a rate of 5 K/min. The pulse isotopic exchange measurements were conducted in the $p\text{O}_2$ range 0.1–0.95 atm from room temperature onwards up to 1,023 K. The reactor was equilibrated 0.5 h at fixed T and $p\text{O}_2$ before and after data collection. Blank experiments revealed no exchange activity of the quartz microreactor.

The packed-bed microreactor was designed to approximate ideal plug flow behavior [16]. The isotopic exchange measurements were performed under conditions of negligible uptake of ¹⁸O by the BE25 powder relative to the number of ¹⁶O oxygen present in the oxide. The overall surface exchange rate, \mathfrak{R}_0 (molO m⁻² s⁻¹), was calculated from

$$\mathfrak{R}_0 = \frac{c_g}{S_r F} \ln \left(\frac{f_i^*}{f_e^*} \right) \quad (1)$$

where f_i^* and f_e^* are the ¹⁸O isotope fractions in the pulse at the inlet and exit of the reactor, respectively, S_r is the total BET surface area of the BE25 powder in the reactor, F is the flow rate, and $c_g = 2 p\text{O}_2 / RT$ is the concentration of oxygen in the gas phase. The flow rate F can be equated to that at reference temperature: $F = F_0 \cdot (T/T_0)$. The ¹⁸O isotope gas phase fraction was calculated from the f^{34} and f^{36} mass fractions: $f^* = 0.5 f^{34} + f^{36}$. Average values of three pulse experiments at a given T and $p\text{O}_2$ were taken for evaluation of the surface exchange rate.

Results and discussion

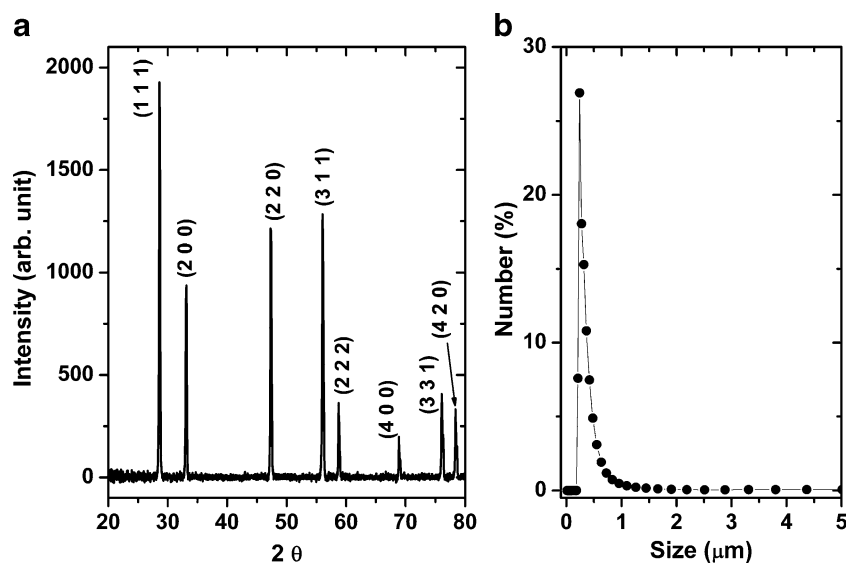
The XRD powder diffractogram (Fig. 2a) confirms the presence of a single cubic phase with a lattice parameter of $a = 5.4736$ (6) Å, which is in agreement with data reported

in literature [17]. No evidence for the formation of second phases is found. The particle size distribution of the crushed BE25 ceramics as measured by laser light scattering is shown in Fig. 2b. The estimated average particle size, $d_{50} = 0.4$ μm, is much smaller than the critical thickness (see “Introduction” section), ensuring that the overall oxygen isotope equilibration is governed by the surface exchange kinetics rather than by bulk diffusion. The associated surface area calculated by the BET method using N₂ (77 K) gas adsorption data amounts to 0.2407 (5) m²/g. The low value reflects the dense nature of the used BE25 powder, and is due to sintering prior to crushing the ceramic disk into a powder. The sintering step is made to avoid diffusion limitations in narrow pores during oxygen isotope equilibration.

Figure 3 shows the fractions of oxygen isotopomers in the gas phase volume associated with the pulse at the exit of the reactor as measured at different temperatures. It is seen that under the conditions maintained during the experiments, the exchange reaction starts around 773 K. Above this temperature, the ¹⁸O₂ fraction decreases profoundly with increase of temperature, i.e., more ¹⁸O is taken up by the sample, which is exchanged for ¹⁶O as evidenced by the simultaneous increase of the ¹⁶O₂ fraction. The ¹⁸O¹⁶O fraction remains comparatively low at all temperatures. The latter provides evidence that oxygen surface exchange on BE25 is limited by the dissociative adsorption of oxygen, as is further discussed below. The highest temperature of measurement (1,023 K) was chosen 50 K lower than the temperature of the pre-annealing in order to prevent microstructural changes to the powder.

The temperature dependence of the surface exchange rate, \mathfrak{R}_0 , calculated using Eq. (1) is well described by an Arrhenius equation, as shown in Fig. 4. Also shown in this figure are results from previous studies [11–13, 18], showing good agreement with those from the present study. The

Fig. 2 a XRD pattern and b particle size distribution of BE25 powder used for the PIE experiments



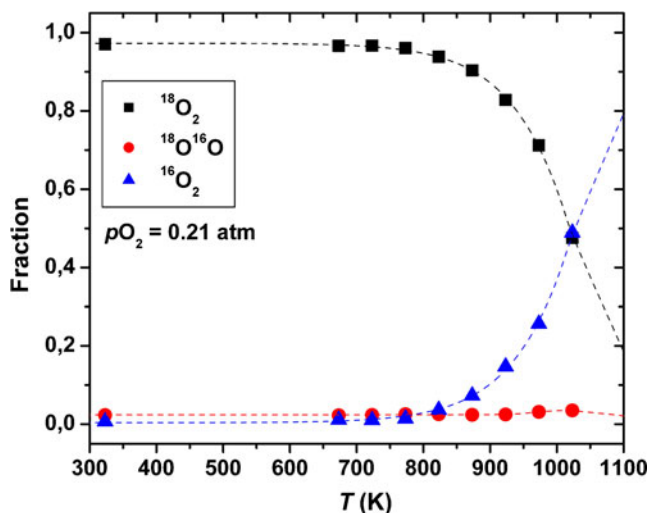


Fig. 3 Oxygen isotope fractions ($^{18}\text{O}_2$, $^{18}\text{O}^{16}\text{O}$, and $^{16}\text{O}_2$) as a function of temperature from PIE measurements at a nominal pressure of 0.21 atm $p\text{O}_2$. The drawn line is from model calculations, assuming constant activation energies for dissociative adsorption and incorporation over the entire temperature range. Error bars (for the 95% confidence interval of the mean) are smaller than the size of the symbols

calculated activation energy of 111 (3) kJ mol^{-1} is in reasonable agreement with 130(5) kJ mol^{-1} calculated from data obtained by monitoring the time-dependent ^{18}O – ^{16}O isotopic equilibration on a polished BE25 dense ceramic disk in a closed volume [11, 18]. The surface exchange rates extracted from data of the depth-profiling studies displayed in Fig. 4 were recalculated using $\mathfrak{R}_0 = k_s c_o$, where k_s (cm s^{-1}) is the surface exchange coefficient and c_o the concentration

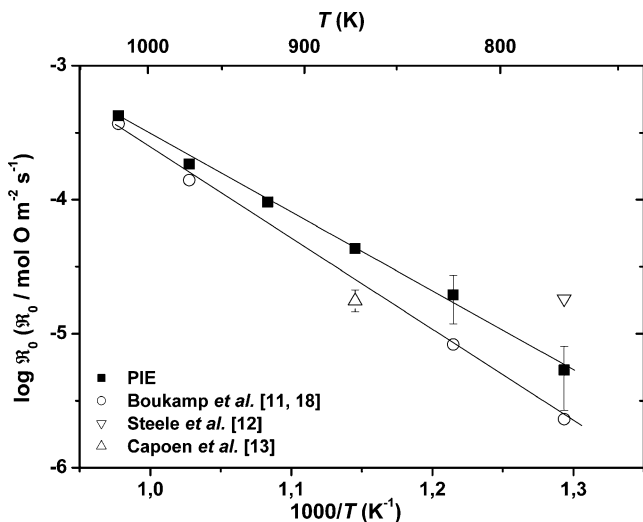


Fig. 4 Overall surface oxygen exchange rate for BE25 as a function of inverse temperature at a nominal pressure of 0.21 atm $p\text{O}_2$. Also shown are data from prior studies on BE25, obtained using time-dependent ^{18}O – ^{16}O gas phase equilibration (Boukamp et al. [11, 18]) and ^{18}O – ^{16}O isotope exchange followed by depth profiling (Steele et al. [12] and Capoen et al. [13]) methods. Error bars represent the 95% confidence interval of the mean; when not shown, these are smaller than the symbol size

of oxygen ions in the oxide. It should be mentioned that it cannot be excluded that different preparation routes and pre-annealing temperatures, and departures from the targeted composition and/or contaminations may influence the magnitude of the surface exchange rate. Figure 5 shows the $p\text{O}_2$ dependence of the overall surface exchange rate at different temperatures. At all four different temperatures, \mathfrak{R}_0 is found approximately proportional to $(p\text{O}_2)^n$ with $n=0.5$, in nice agreement with results published previously [18].

The above results demonstrate the usefulness of PIE as a technique for acquisition of the oxygen surface exchange rate of fast oxygen-ionic conducting solids. The ^{18}O -containing gas phase pulse passes through a ‘sea of ^{16}O ’ captured by the oxide powder bed, ensuring in this way that ^{16}O is returned to the gas phase upon every act of exchange. Measurements of the surface exchange rate are enabled as long as the effective time constant associated with isotopic exchange is of similar order as the time scale of the experiment, i.e., the residence time of the packed-bed reactor. These are determined by oxide characteristics, such as the powder surface area, and by design and operating conditions, such as the length of the packed-bed reactor, gas flow rate, and temperature, as can be inferred from Eq. (1). The design of the packed bed reactor to approximate plug flow conditions ensures that the distribution of residence times is small, and its average value simply can be calculated from the void volume of the reactor bed and the volumetric flow rate.

An advantage of methods based upon gas phase analysis to monitor ^{18}O – ^{16}O isotope equilibration between the gas

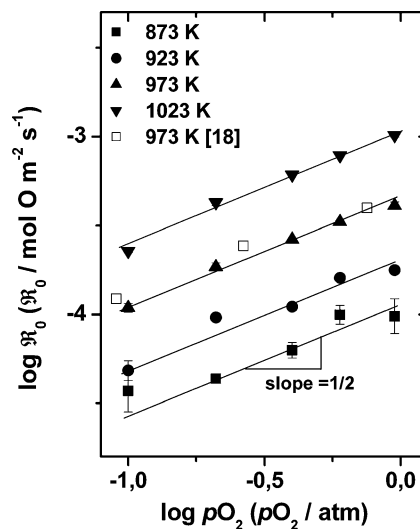


Fig. 5 Oxygen partial pressure dependence of the overall exchange rate on BE25 at different temperatures using PIE (filled symbol). Lines with slope 1/2 are given to guide the eye. Error bars represent the 95% confidence interval of the mean; when not shown, these are smaller than the symbol size. The data from time-dependent ^{18}O – ^{16}O gas phase equilibration on BE25 dense ceramic at 973 K reported by Boukamp et al. [18] (empty square symbol) is also included

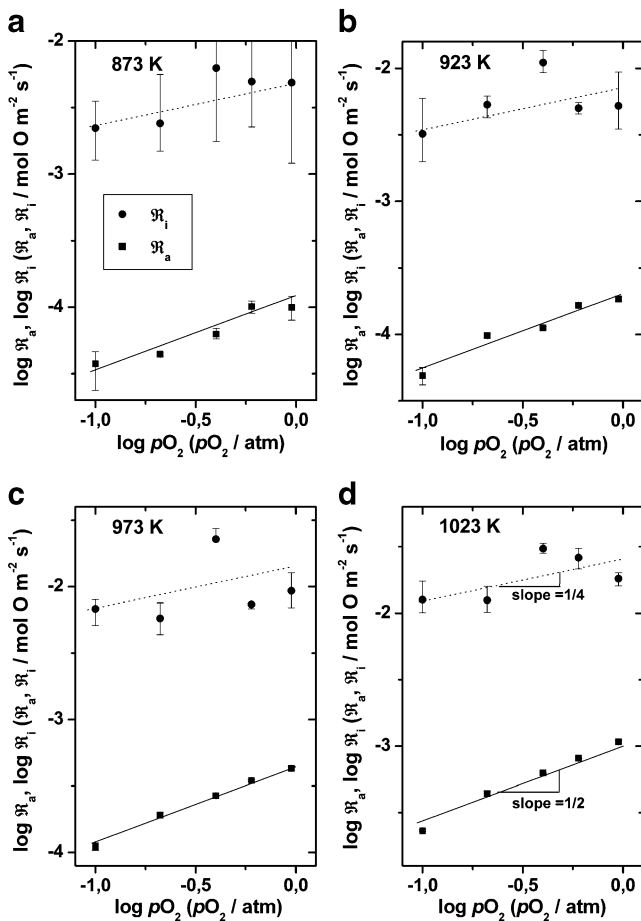


Fig. 6 The oxygen partial pressure dependence of the rates of dissociative adsorption (R_a , filled square) and oxygen incorporation (R_i , filled circle) at different temperatures. Trendlines with slopes 1/2 and 1/4 are given to guide the eye. Error bars represent the 95% confidence interval of the mean; when not shown, these are smaller than the symbol size

phase and the oxide over that of depth profiling methods is that kinetic information can be extracted from the mass isotopomer distribution resulting from isotopic scrambling. The isotopic exchange reaction can be modeled as a sequence of two steps, dissociative adsorption occurring with a rate R_a , and subsequent incorporation of oxygen in the oxide lattice with a rate R_i ,



and includes oxygen adatoms O_{ad} on the oxide surface as reaction intermediates. The ^{18}O isotope fraction at the oxide surface is balanced by the reversible rates R_a and R_i in conjunction with the isotopic compositions of the gas phase and oxide. The validity of the two-step scheme for modeling isotopic exchange relies on the assumption that

isotopic scrambling can only occur after dissociative adsorption of oxygen on the oxide surface, excluding for example, the possibility that exchange may occur via a three-atom intermediate. In a quasi steady-state approximation for the isotopic exchange reaction, it follows

$$\frac{1}{R_0} = \frac{1}{R_a} + \frac{1}{R_i} \quad (4)$$

The overall exchange process can be conceptually viewed as a series of resistances. Expressions for the corresponding molar gas phase fractions of $^{18}\text{O}_2$, $^{16}\text{O}^{18}\text{O}$, and $^{16}\text{O}_2$ after exchange for a given time within the framework of the two-step scheme are presented elsewhere [16, 18]. Finally, it should be noted that the surface exchange reaction may involve a series of steps with several possible intermediates, e.g., O_2 , O_2^- , and O^- . Both steps in the given two-step scheme still may represent a lumped reaction sequence with, for example, charge transfer and surface diffusion of atoms or ions as possible rate determining steps.

Figure 6a–d show the $p\text{O}_2$ dependences of R_a and R_i for BE25 at different temperatures. The results indicate that the surface exchange reaction on BE25 is mainly limited by the rate of oxygen dissociative adsorption R_a . This result can be deduced intuitively from the very small molar $^{18}\text{O}^{16}\text{O}$ fraction observed during isotopic exchange, as illustrated in Fig. 3. At all temperatures, R_a appears proportional to $p\text{O}_2^{1/2}$, in obvious correspondence with the $p\text{O}_2$ dependence of R_0 , and found one to two orders of magnitude lower than R_i .

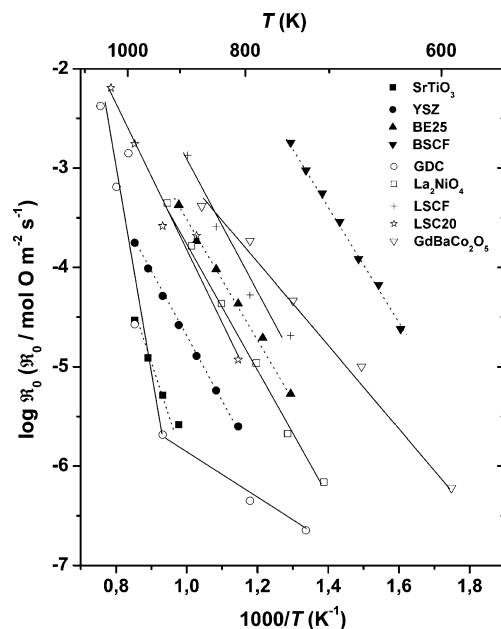
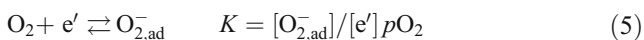


Fig. 7 Surface oxygen exchange rates for different oxides as a function of inverse temperature. SrTiO_3 [21], 8 mol% yttria-stabilized zirconia (YSZ) [16], BE25 (this study), $\text{Ba}_{0.5}\text{Sr}_{0.5}\text{Co}_{0.8}\text{Fe}_{0.2}\text{O}_{3-\delta}$ (BSCF) [16], 10 mol% gadolinia-doped CeO_2 (GDC) [22], La_2NiO_4 [19], $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.2}\text{Fe}_{0.8}\text{O}_{3-\delta}$ (LSCF) [23], $\text{La}_{0.8}\text{Sr}_{0.2}\text{CoO}_{3-\delta}$ (LSC20) [24], and $\text{GdBaCo}_2\text{O}_5$ [25]

The dissociative adsorption of oxygen on the surface can be represented by the successive reaction steps



where K is the mass action constant of reaction (5). The observed $p\text{O}_2^{1/2}$ dependence of \mathfrak{R}_a can be explained by assuming that electron transfer to intermediate superoxide ions, O_2^- , according to reaction (6), is the rate determining step in oxygen dissociative adsorption. Noting that the isotopic exchange experiment is carried out under conditions of equilibrium, $\mathfrak{R}_a = \overrightarrow{\mathfrak{R}}_a = \overleftarrow{\mathfrak{R}}_a$, where $\overrightarrow{\mathfrak{R}}_a = \overrightarrow{k}[\text{O}_{2,\text{ad}}^-][e']$ and $\overleftarrow{\mathfrak{R}}_a = \overleftarrow{k}[\text{O}_{\text{ad}}^-]^2$ are the balanced forward and backward reaction rates, respectively, and \overrightarrow{k} and \overleftarrow{k} the corresponding rate constants. In deriving these kinetic equations, a low surface coverage of adsorbed oxygen species is assumed. To infer the $p\text{O}_2$ dependence of \mathfrak{R}_a , those of the concentrations entering these kinetic equations must be derived from preceding or succeeding equilibria. Under the experimental conditions, the bulk concentration of oxygen vacancies in BE25 is invariant with $p\text{O}_2$. From defect chemical considerations, it follows that for BE25 the concentration of electrons varies according to $[e'] \propto p\text{O}_2^{-1/4}$. Substitution in the expression for $\overrightarrow{\mathfrak{R}}_a$ ($= \overrightarrow{k}[\text{O}_{2,\text{ad}}^-][e'] = K\overleftarrow{k}[e']^2p\text{O}_2$) shows that in accord with the experimental observations \mathfrak{R}_a is predicted to follow a $p\text{O}_2^{1/2}$ dependence. Reaction (6) may proceed via formation of a transient peroxide intermediate, O_2^{2-} , which is known to be very unstable, before actual splitting of the adsorbed oxygen molecule into oxygen adatoms O_{ad}^- .

As seen from Fig. 6a–d, the rate of oxygen incorporation \mathfrak{R}_i shows a weak dependence on $p\text{O}_2$. The trend lines in these figures with a slope of 1/4 should be considered with care given the large scatter in the obtained values of \mathfrak{R}_i . \mathfrak{R}_i may either refer to a successive reaction step or to diffusion of O_{ad}^- or electron holes to the reaction site. More research is, however, needed for clarification.

Finally, Fig. 7 shows a graphical collection of data obtained for the surface oxygen exchange rate observed on various oxides using different techniques. It is seen that BE25 shows a comparatively high surface exchange rate, much higher than well-known solid electrolytes YSZ and gadolinia-doped ceria, and even higher than La_2NiO_4 , which has been proposed as a cathode material for intermediate-temperature SOFCs [19, 20].

Conclusions

The results obtained from this study indicate that in the temperature and $p\text{O}_2$ ranges investigated BE25 exhibits a

comparatively high exchange rate, which is rate determined by the dissociative adsorption of oxygen. Electron transfer to intermediate superoxide ions is proposed as the rate determining step. The results further demonstrate the usefulness of pulse-based measurements of ^{18}O – ^{16}O isotope exchange, which method employs a continuous flow packed bed microreactor, for kinetic and mechanistic studies of surface oxygen exchange of fast ionic conducting solids.

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