Review on exsolution and its driving forces in perovskites

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Abstract

Exsolution is a promising method to design metal nanoparticles for electrocatalysis and renewable energy. Metal nanoparticles exsolved from perovskite oxide lattices have been utilized as catalysts in many energy fields because of their high durability and excellent electro-catalytic properties. Although this method has received much attention in recent years, a comprehensive understanding is still lacking because of difficulties in finding a rational combination of driving forces and perovskite supports. Thus, the aim of our work here is to recapitulate the principles of exsolution and collect various exsolution studies by categorizing the driving forces of exsolution and the structural characteristics of perovskite supports. These classifications provide guidelines for selecting suitable materials groups and remodeling existing materials, thereby exploring applications of catalysts using exsolution that are applicable to academic and industrial fields.

1. Introduction

Metal nanoparticles, active sites of electrochemical reactions, have been intensively researched toward the development of more stable and effective catalysts due to their key role in electrochemical energy devices, electronics, optics, and catalysis [1–4]. Since metal nanoparticles are difficult to keep secure and stable, heterogeneous composites that introduce a supporting material to immobilize high-surface-area catalytic particles have been developed. Such composite materials are fabricated by depositing metal nanoparticles on the surface of the supporting material through chemical vapor deposition (CVD) or infiltration methods [5–7]. However, the deposited metal nanoparticles may undergo agglomeration and coarsening during cell operating conditions, leading to fatal performance degradation. Thus, an exsolution technique has been received extensive attention as a promising alternative.

Exsolution is a phenomenon in which metal catalysts are segregated from an oxide lattice to an oxide surface under a reducing atmosphere [8, 9]. Unlike the conventional deposition techniques, exsolution is particularly attractive for its time efficiency because no additional process is required for catalyst addition. Additionally, the exsolved metal nanoparticles have the unique feature of socketing into the oxide surface (figure 1(a)), enhancing cohesion between the metal nanoparticles and the supporting material [10]. Consequently, well-distributed and stable catalysts can be formed on the supporting material surface through the exsolution. The two key factors in producing nanocatalysts through exsolution are: (1) the perovskite supports should have high redox stability and (2) the metal cations should easily come to the surface under a reducing atmosphere. However, the complicated relationship between these factors and the numerous combinations with various perovskite oxides make the phenomena difficult to understand. Therefore, it is essential to identify and organize the exsolution phenomena to choose a rational catalyst.

This review begins with an overview of the general characteristics and driving forces of exsolution in perovskite oxides. Although several articles reviewing the exsolution have been published previously [12, 13], intensive research of exsolution is a recent endeavor, thus it is necessary to recapitulate the principles of the phenomena, and to collect scattered studies. Therefore, we intend to provide guidelines for selecting suitable
Figure 1. (a) Schematic illustration of the particle-substrate interface for deposited and exsolved nickel particles. (b) TEM micrograph (dark field) of a Ni particle exsolved on (110) native surface facet (see figure 1(d)) after ageing (~3% H₂O/5% H₂/Ar, 930 °C, 60 h). (c) Three-dimensional AFM of sockets. (d) TEM image detail (bright field) of the corresponding atomic planes and orientations. (e) AFM image after reduction at 700 °C (15 min, dry H₂). (f) 3D view of figure 1(e). (g) AFM height profile along the line shown in figure 1(e). (a), (b), (c), and (d) Reproduced with permission from [10], Copyright 2015, Nature Publishing Group. (e), (f), and (g) Reproduced with permission from [11], Copyright 2015, American Chemical Society.

2. Characteristics and driving forces of exsolution in perovskite oxides

In this section, we introduce the general characteristics of nanocatalysts produced by exsolution. The general characteristics include morphology, carbon deposition characteristics, reversibility, and exsolution mechanism. Several studies highlight a morphological study of particle-substrate interactions with anchored nanoparticles [10]. As shown in transmission electron microscopy (TEM) analysis (figures 1(b) and (d)), the exsolved Ni particles have a pinned structure that is approximately 30% submerged into the parent oxide. Additionally, the atomic force microscopy (AFM) view of the etched surface (figure 1(c)) demonstrated that nanoparticles were embedded in the oxide matrix to a significant depth. Detailed exsolution morphology is described in a study by Oh et al [11]. As described by this study, exsolution proceeds through the formation of subsurface nucleates, and the movement of metal nucleate to the surface is accompanied by the formation of a pit on the surface where the particle emerges (figures 1(e)–(g)). Socketed structural features resulted in improved thermal stability of the catalyst and significantly reduced coke formation. This can be attributed to carbon fiber growth in exsolved particles occurring as a base-growth, unlike typical carbon fiber that undergoes tip-growth (figure 2(a)). In this case, the base growth of carbon fiber is restrained compared to the tip growth of carbon fiber.

Dopant cations dissolved in the parent oxide lattice matrix are released and then anchored to the surface under a reducing atmosphere, and this process is said to have redox reversibility if it can occur in reverse. It has been speculated that particles generally re-dissolve back into the perovskite oxide under an oxidizing atmosphere. However, as reported by Neagu et al [14], Ni and Co-Ni particles do not re-dissolve back into the underlying perovskite host lattice during the oxidation in La₀.₈Ce₀.₂(₂Ni₀.₄Ti₀.₆)O₃ and La₀.₇Ce₀.₃(Co₀.₃Ni₀.₁)Ti₀.₆O₃ systems (figure 3). In the Co-Ni system, approximately 0.06 exsolved B-site atoms per unit cell, which is thought to make re-dissolution less favorable with the larger amount of 0.2 preferred for A-site vacancy.

The exsolution mechanism has been actively studied in recent years. Exsolution is a chemically driven heterogeneous phase transition, and Gao et al [15] proposed that exsolution arises from the four physical processes of diffusion, reduction, nucleation, and growth (figure 4(a)). Exsolving dopant cations are first
Figure 2. (a) Schematic of possible carbon fiber growth mechanisms. (b) Ni particles (30–100 nm) prepared by vapor deposition on La$_{0.4}$Sr$_{0.4}$TiO$_3$, showing considerable carbon fiber growth. (c) False color micrograph depicting a side view detail of La$_{0.52}$Sr$_{0.28}$Ni$_{0.06}$Ti$_{0.94}$O$_3$ (5% H$_2$, 880 °C, 6 h). Reprinted with permission. (a), (b), and (c) Reproduced with permission from [10] Copyright 2015, Nature Publishing Group.

Figure 3. (a)–(b), (d)–(g) SEM images of tracked areas collected at room temperature at highlighted stages of catalytic testing, and corresponding cross-section schematics of the particle-perovskite interface. (c) XPS valence band spectra, guideline showing evolution of Co$_3$O$_4$ phase across selected samples. (h) Schematic of the local appearance of the perovskite surface after cube removal. Reproduced with permission from [14] Copyright 2017, Nature Publishing Group.
diffused from the bulk to the surface and then reduced to metal. Reduced metals are assembled into small nanoparticles that grow in size over the treatment time. Recently, direct observation of particle nucleation and growth during the exsolution has confirmed that the particles grow epitaxially and isotropically while maintaining their original positions [16]. The study also demonstrated that exsolution is limited by the exsolvable ion availability rather than diffusion to the surface. Gao et al [15] studied the effect of reducing time and atmosphere composition on particlegrowth and proposed three analytical models for growth, in which particlegrowth can be limited by strain, reactant amount, and diffusion. The results suggest that both the size-related strain and limited amounts of Ni could be possible mechanisms determining the particle size growth rate. However, the actual exsolution mechanism is likely a much more complex process than the models suggest, as nucleation, growth, and particle interaction need to be considered simultaneously.

Sincetheexsolutionoccursunderareducingatmosphere,ageeneraldrivingforceofexsolutionisanoxygenvacancyformationintheperovskiteoxides. Theoxygenvacancyformationdestabilizesthelatticestoichiometry,causingthesegregationofB-sitecationstomaintainstructuralstability.Severaldrivingforces,suchastemperature,non-stoichiometrybyA-site defect,voltagebiasing,strain,phasetransition,andtopotacticionexchange,havebeenintroducedtoformtheoxygenvacancy(figure 5).However,thedegreeofoxygenvacancyformationdiffersforeachperovskitestructure.Thus,thefollowingsectionsdiscussvariousexsolutionphenomenaandstructuralpropertiesintheperovskiteoxidestodevelopamoredetailedunderstanding.Alistofmaterialsbystructureanddrivingforceisgivenintable 1.

2.1. Exsolution on stoichiometric single perovskite
Perovskite oxides (ABO₃) and derived structures (e.g. double perovskite (A₂B₂O₆₋₁) and Ruddlesden–Popper (RP, Aₙ₊₁B₂O₃₋₂n₊₁)-based materials have been mainly studied as the support materials for exsolution due to their excellent structural and compositional flexibility (figure 6). In typical perovskite oxides, it is possible to withstand some lattice mismatch between A–O and B–O bonds, thus allowing doping with different types of cations at each site [60, 61]. This flexibility of the perovskite structure enables the combination of a wide range of constituent ions, providing a variety of catalytic properties. In particular, studies have been conducted on redox couples such as Ti⁴⁺/Ti³⁺, Mn⁴⁺/Mn³⁺, Ce⁴⁺/Ce³⁺, Mo⁶⁺/Mo⁵⁺, Cr³⁺/Cr²⁺, Nb⁴⁺/Nb³⁺, and V⁴⁺/V³⁺, which can be stable under a redox atmosphere.

EarlyexsolutionconceptswereappliedtostoichiometricperovskitewithA/B = 1, in which only readily reducible cations (e.g. Ni²⁺, Ru²⁺, Rh⁴⁺, Pd⁴⁺, and Pt⁴⁺) could be exsolved to metal nanoparticles [62]. In such a case, most cations remain in the bulk, thus the number of nanoparticles available on the surface is limited. Since LaFe₀.₅₇Co₀.₃₆Pd₀.₀₉O₃ [9] opened the possibility of in situ growth catalysts, perovskite oxides (ABO₃) have been employed as supporting frameworks. In early research, noble metals (e.g. Pd [9, 63], Ru [19, 20, 64], Rh, and Pt) were doped into B-site in perovskite lattice and partly exsolved under a reducing atmosphere. The Pd-perovskite catalyst [9] lasted about 100 h in an aging test under an atmosphere consisting of engine exhaust. The mol percentage of Pd was increased to 0.2 in La₀.₃Sr₀.₇Co₀.₃Pd₀.₃O₃.₅ [17], and the elimination of Pd nanoparticles upon oxidation and their re-nucleation upon reduction were observed. Meanwhile, Hamada et al [65] proposed a possible mechanism for the self-regeneration of Pd catalysts in LaFe₁₋ₓPdₓO₃. On the basis of the density functional theory (DFT) calculations, oxygen vacancies stabilized the surface segregation of Pd in the perovskite supporting material, forming LaPdOₓₐ in the
| Classification          | Catalysts                                                                 | Nature of materials | Environmental conditions for exsolution | Catalytic applications                        | Stability (hrs) |
|------------------------|---------------------------------------------------------------------------|---------------------|----------------------------------------|-----------------------------------------------|-----------------|
| Stoichiometric single perovskite | LaFeO$_{0.52}$CoO$_{0.35}$PdO$_{0.03}$O$_3$ [9]                        | Noble metal in B-site | Temperature and gas condition         | Automotive emissions control                  | 100             |
|                        | LaFeO$_{0.95}$PdO$_{0.05}$O$_3$ [9]                                      | Noble metal in B-site |                                        | Automotive emissions control                  | 100             |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Pd$_x$O$_3$ [17]                         | Noble metal in B-site |                                        | SOFC anode                                    | 40              |
|                        | La$_{0.57}$Sr$_{0.43}$MnO$_2$                                              | Noble metal in B-site |                                        | SOFC anode                                    | 100             |
|                        | La$_{0.38}$Sr$_{0.62}$Cr$_{0.5}$Ru$_{0.5}$O$_{1.5}$ [19]                 | Noble metal in B-site |                                        | SOFC anode                                    | 60              |
|                        | La$_{0.05}$Sr$_{0.95}$Fe$_{0.05}$O$_{1.5}$                                | Noble metal in B-site |                                        | SOFC anode                                    | -               |
|                        | LaCoO$_{0.95}$NbO$_{0.05}$                                               | Noble metal in B-site |                                        | Refining catalysts                             | 72              |
|                        | La$_{0.2}$Sr$_{0.8}$Co$_{0.7}$Ni$_{0.3}$O$_{1.5}$                         | Ni metal in B-site   |                                        | Reforming catalysts                           | -               |
|                        | La$_{0.4}$Sr$_{0.6}$Fe$_{0.5}$CoO$_{1.5}$                               | Ni metal in B-site   |                                        | Reforming catalysts                           | -               |
|                        | (La$_{0.2}$Sr$_{0.8}$)Cr$_{0.5}$Fe$_{0.5}$O$_{1.5}$                       | Ni metal in B-site   |                                        | Reforming catalysts                           | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Co$_{0.5}$Mn$_{0.5}$O$_{1.5}$                       | Ni metal in B-site   |                                        | Reforming catalysts                           | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Fe$_{0.5}$CoO$_{1.5}$                               | Ni metal in B-site   |                                        | Reforming catalysts                           | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Cr$_{0.5}$CoO$_{1.5}$                               | Ni metal in B-site   |                                        | Reforming catalysts                           | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Co$_{0.5}$FeO$_{1.5}$                               | Ni metal in B-site   |                                        | Reforming catalysts                           | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Ti$_{0.8}$CoO$_{1.5}$                               | Co-Fe metal in B-site|                                        | Refining catalysts                             | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Ti$_{0.8}$CoO$_{1.5}$                               | Co-Fe metal in B-site|                                        | Refining catalysts                             | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Ti$_{0.8}$Co$_{0.5}$Fe$_{0.5}$O$_{1.5}$              | Co-Fe metal in B-site|                                        | Refining catalysts                             | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Ti$_{0.8}$Co$_{0.5}$Mn$_{0.5}$O$_{1.5}$              | Co-Fe metal in B-site|                                        | Refining catalysts                             | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Ti$_{0.8}$Co$_{0.5}$FeO$_{1.5}$                      | Co-Fe metal in B-site|                                        | Refining catalysts                             | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Ti$_{0.8}$Co$_{0.5}$MnO$_{1.5}$                     | Co-Fe metal in B-site|                                        | Refining catalysts                             | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Ti$_{0.8}$Co$_{0.5}$MnO$_{1.5}$                     | Co-Fe metal in B-site|                                        | Refining catalysts                             | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Ti$_{0.8}$Co$_{0.5}$FeO$_{1.5}$                      | Co-Fe metal in B-site|                                        | Refining catalysts                             | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Ti$_{0.8}$Co$_{0.5}$MnO$_{1.5}$                     | Co-Fe metal in B-site|                                        | Refining catalysts                             | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Ti$_{0.8}$Co$_{0.5}$FeO$_{1.5}$                      | Co-Fe metal in B-site|                                        | Refining catalysts                             | -               |
|                        | La$_{0.3}$Sr$_{0.7}$Ti$_{0.8}$Co$_{0.5}$MnO$_{1.5}$                     | Co-Fe metal in B-site|                                        | Refining catalysts                             | -               |
| Non-stoichiometric single perovskite | La$_{0.57}$Sr$_{0.43}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [10]            | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [31]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.57}$Sr$_{0.43}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [32]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [33]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.57}$Sr$_{0.43}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [34]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [35]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [36]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [37]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [38]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [39]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [40]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [41]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [42]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [43]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [44]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [45]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
|                        | La$_{0.37}$Sr$_{0.63}$CoO$_{1-x}$Ni$_{0.3}$O$_{1.5}$ [46]              | A-site deficiency   |                                        | SOEC cathode condition                        | 10              |
Table 1. Continued.

| Classification | Catalysts | Nature of materials | Environmental conditions for exsolution | Catalytic applications | Stability (hrs) |
|----------------|-----------|---------------------|-----------------------------------------|------------------------|----------------|
| Double perovskite | PrBaMn$_{2.2}$O$_{5.5+6}$ [47] | Phase transition | Temperature and gas condition | SOFC anode | |
| | PrBaMn$_{2.2}$Co$_{0.3}$O$_{5.5+6}$ [47] | Phase transition | | | |
| | PrBaMn$_{2.2}$Ni$_{0.3}$O$_{5.5+6}$ [47] | Phase transition | | | |
| | PrBaMn$_{1.7}$Fe$_{0.3}$O$_{5.5+6}$ [47] | Phase transition | | | |
| | NdBaMn$_{2}$O$_{5.5+6}$ [48] | Phase transition | | | |
| | PrBaMn$_{1.7}$Co$_{0.4}$Ni$_{0.3}$O$_{5.5+6}$ [49] | Phase transition | | | |
| | PrBaMn$_{1.7}$Co$_{0.4}$Ni$_{0.3}$O$_{5.5+6}$—Fe infiltration [50] | Phase transition and topotactic exchange | | | |
| | PrBaMn$_{1.8}$Co$_{0.2}$O$_{5.5+6}$ [51] | Phase transition | | | |
| | (Pr$_{0.4}$Sr$_{0.6}$)$_3$(Fe$_{0.85$}Nb$_{0.15}$)$_2$O$_{7}$ [52] | Phase transition | | | |
| | Sr$_3$FeMoO$_{7.6}$ [53] | Phase transition | | | |
| | Pr$_{0.8}$Sr$_{1.2}$Co$_{0.4}$Fe$_{0.8}$Nb$_{0.2}$O$_{4+6}$ [54] | Phase transition | | | |
| | La$_{1.2}$Sr$_{0.8}$Mn$_{0.4}$Fe$_{0.6}$O$_{4}$ [55] | Phase transition | | | |
| | Sr$_3$FeMoO$_{6.5}$ [56] | Phase transition | | | |
| | La$_{0.8}$Sr$_{1.2}$Fe$_{0.8}$Co$_{0.1}$O$_{4+6}$ [57] | Phase transition | | | |
| RP perovskite | PrBaMn$_{1.7}$Co$_{0.2}$O$_{5.5+6}$ [51] | Phase transition | Temperature and gas condition | SOFC anode and SOEC cathode | |
| | (Pr$_{0.4}$Sr$_{0.6}$)$_3$(Fe$_{0.85$}Nb$_{0.15}$)$_2$O$_{7}$ [52] | Phase transition | | | |
| | Sr$_3$FeMoO$_{7.6}$ [53] | Phase transition | | | |
| | Pr$_{0.8}$Sr$_{1.2}$Co$_{0.4}$Fe$_{0.8}$Nb$_{0.2}$O$_{4+6}$ [54] | Phase transition | | | |
| | La$_{1.2}$Sr$_{0.8}$Mn$_{0.4}$Fe$_{0.6}$O$_{4}$ [55] | Phase transition | | | |
| | Sr$_3$FeMoO$_{6.5}$ [56] | Phase transition | | | |
| | La$_{0.8}$Sr$_{1.2}$Fe$_{0.8}$Co$_{0.1}$O$_{4+6}$ [57] | Phase transition | | | |
| | La$_{1.2}$Sr$_{0.8}$Co$_{0.2}$Mn$_{0.8}$O$_{4}$ [58] | Phase transition | | | |
| | SrGdNi$_{0.2}$Mn$_{0.8}$O$_{4+6}$ [59] | Phase transition | | | |
| | La$_{1.2}$Sr$_{0.8}$Co$_{0.2}$Mn$_{0.8}$O$_{4}$ [58] | K$_2$NiF$_2$-type structural perovskite | | CO$_2$ electrolysis | |
| | SrGdNi$_{0.2}$Mn$_{0.8}$O$_{4+6}$ [59] | K$_2$NiF$_2$-type structural perovskite | | | |
Figure 5. Driving forces to promote the exsolution in perovskite oxides: gas/temperature, cation deficiency, biasing, strain, phase transition, and topotactic exchange. In the driving forces, the green spheres indicate exsolved nanoparticle. In the middle circle, blue, green, and red spheres indicate A-site cation, B-site cation, and oxygen, respectively.

Figure 6. Crystal structures of (a) single perovskite (ABO$_3$), (b) double perovskite (AA'B$_2$O$_{5+δ}$), and (c) RP perovskite (An+1BnO$_3$ n+1).

vicinity of the LaFe$_{1-x}$Pd$_x$O$_3$ surface. The LaPdO$_{3-δ}$ layer on the surface allows Pd atoms to move in and out of the perovskite lattice without long-distance diffusion through the bulk during the redox cycles.

In the case of Pd exsolution, co-exsolution with Fe has been reported in B-site excessive stoichiometric La$_{0.8}$Sr$_{0.2}$Fe$_{0.9}$Nb$_{0.1}$Pd$_{0.04}$O$_{3-δ}$ [18]. Fe$^0$ and Pd$^0$ metallic nanoparticles were confirmed to exsolve on the surface of the perovskite anode during operation under a hydrogen atmosphere. The introduction of Pd increased electrical conductivity and H$_2$-adsorption. In the study on exsolving Ru [19] from a host lattice, increasing the electrochemical kinetics of Ru had a more pronounced effect than increasing that of Ni. The exsolved Ru nanoclusters were smaller (≤ 5 nm) than those of Ni (~10 nm), and there were more Ru
nanoclusters than Ni nanoclusters. An attempt was made to dope Ir on the B-site of LaCrO$_3$-based perovskite [21]. The CH$_4$ conversion of the dry reforming reaction reached 81% at 750 °C and was maintained for about 72 h without coke formation on the catalyst surface.

Among transition metals, early research on producing Ni nanoparticles was carried out based on La$_{1-x}$Sr$_x$Cr$_{1-y}$Ni$_y$O$_3$ [22]. The catalytic properties improved with the formation of Ni nanoparticles under a reducing atmosphere, however, catalytic activity performance was not proportional to the amount of Ni.

La$_{0.5}$Sr$_{0.5}$Ti$_{0.75}$Ni$_{0.25}$O$_{3+\delta}$ containing a significant amount of Ni was also successfully synthesized and Ni nanoparticles were exsolved in a hydrogen atmosphere. As an anode of solid oxide fuel cell (SOFC), La$_{0.75}$Sr$_{0.25}$Cr$_{0.5}$Mn$_{0.3}$Ni$_{0.2}$O$_{2.8}$ [26] was studied and the limit of the solid solution of Ni was 20% in B site. The formation of Ni nanoparticles during the reduction process could facilitate the water steam dissociation. Gao et al [66] recently studied the energetics of Ni segregation in a SrTiO$_3$-based perovskite framework. They demonstrated that Ni preferentially segregates toward the (100)-oriented and SrTiO$_3$-terminated surfaces. The Ni surface segregation is energetically promoted by the presence of A-site and O-site vacancies, whereas the substitution of Sr$^{3+}$ with La$^{3+}$ hinders surface segregation. Sun et al reported on the production of coking-tolerant anodes by co-doping Ni and Ce [25]. Exsolved metallic Ni provides a strong active site for fuel oxidation reactions, while Ce species with redox couples provide abundant oxygen ions. Moreover, a SOFC equipped with a Ni/Ge co-doped lanthanum strontium titanate anode also showed good electrochemical performance and stability in both 5000 ppm H$_2$/H$_2$ and dry CH$_4$.

Alloy nanocatalysts among transition metals have also been reported in stoichiometric perovskite. La$_{0.5}$Sr$_{0.5}$Fe$_{0.8}$Ni$_{0.2}$O$_{3+\delta}$ [27] showed excellent current density during CO$_2$ electrolysis and it was found that the material gained redox stability from the symmetry between reduction and re-oxidation (figure 7). In (La$_{0.75}$) (Sr$_{0.25}$) (Cr$_{0.5}$Fe$_{0.35}$Ni$_{0.15}$)O$_3$ [28], it was demonstrated that the exsolution of bimetallic Fe-Ni alloy nanoparticles occurred on the surface of the perovskite. For strontium ferrites based perovskites, Liu et al [29] revealed that Fe-Ni nanoparticles can be exsolved from Sr$_3$Fe$_{1.4}$Ni$_{0.6}$Mn$_{0.3}$O$_6$-δ perovskite. They examined the redox reversibility of the exsolved Fe-Ni nanoparticles which were oxidized to transition metal oxides (Ni,Fe)O instead being reincorporated into the parent perovskite oxide at 800 °C in air. Fe-Cu alloy nanocatalysts were also exsolved in SrFe$_{0.8}$Cu$_{0.2}$O$_{3-\delta}$ [67]. The high conductivity in 5% H$_2$/Ar was achieved because of the exsolved Fe (or Fe-Cu alloy). Co-Fe nanocatalysts on La$_{0.5}$Sr$_{0.5}$Cr$_{0.3}$Fe$_{0.6}$Co$_{0.1}$O$_{3-\delta}$ [30] showed a redox process at 800 °C where the exsolved particles were completely reincorporated into the perovskite lattice.

2.2. Exsolution on non-stoichiometric single perovskite

For these stoichiometric compositions, most cations are present in the oxide matrix under a reducing atmosphere, which is not available for gas-phase catalysis. Additionally, the compositions undergo difficulties associated with phases involving undesirable A-site cations (equation (1)).

$$\text{ABO}_3^\gamma \xrightarrow{\text{Exsolution}} (1 - \alpha) \text{ABO}_3^\gamma' + \alpha \text{AO} + \alpha \text{B}$$  

(1)

$$\text{A}_{1-\alpha}\text{BO}_3^\gamma \xrightarrow{\text{Exsolution}} (1 - \alpha) \text{ABO}_3^\gamma' + \alpha \text{B}$$  

(2)

Therefore, A-site deficient perovskites (A/B < 1) have been proposed to alleviate these problems [8, 11, 68, 69]. In the systems, the exsolution reverts perovskite to a more stable ‘defect-free’ ABO$_3$ stoichiometry, thus inhibiting A-site segregation (equation (2)). Moreover, harder-to-reduce cations can also be exsolved and exsolution preferentially emerges on the surface [69]. For example, in A-site deficient La$_{0.4}$Sr$_{0.6}$TiO$_3$, extremely stable Ti cations can also be exsolved as TiO$_2$ on the surface of the perovskite [69]. Based on this finding, much research on A-site deficient perovskites has been conducted to create more metallic nanoparticles on stable supporting materials in the past few years. Neagu et al [8] systematically reported on the role of A-site deficiency in B-site exsolution. In an A-site stoichiometric sample, a smooth surface exhibits no exsolution, whereas nanoparticles are exsolved on the surface in the case of A-site deficient perovskite under reducing condition. When reducing A-site deficient perovskite A$_{1-\alpha}$BO$_3^\gamma$, oxygen vacancies are introduced as oxygens are removed. The presence of oxygen vacancy can cause spontaneous the exsolution of B-site species, thus destabilizing the perovskite lattice and re-establishing the stoichiometry. The exsolved nanocatalyst is well anchored on the perovskite substrate and stable against aglomeration. Gao et al [15] reported Ni exsolution on a Sc-based A-site deficient perovskite material La$_{0.4}$Sr$_{0.6}$Sc$_{0.2}$Ni$_{0.1}$O$_{3-\delta}$. The reduced samples showed high electrochemical catalytic activity in symmetric-cell measurements with an area-specific resistance of 0.055 Ω cm$^2$ at 800 °C in a humid hydrogen atmosphere. For LaMnO$_3$ based perovskite, it has been reported that 20 mol% Ni-doped LaMnO$_3$, La$_{0.8}$Mn$_{0.2}$O$_2$, O$_3$ [37] showed resistance to carbon fiber formation due to their strong interfacial bonding to the substrate. La$_{0.6}$Sr$_{0.8}$Cr$_{0.85}$Ni$_{0.15}$O$_{3-\delta}$
Figure 7. Thermogravimetric analysis testing. (a) Weight loss and differential thermal analysis of LSFN powders in 5% H₂/N₂ reducing atmosphere. (b) Redox cycling ability test of LSFN powders, conducted first in a 5% H₂/N₂ reducing atmosphere in the ascending temperature range. After it was cooled, the analysis environment was switched to air flow in the same ascending temperature range. Reproduced with permission from [27]. Copyright 2016, American Chemical Society.

[38] exhibited enhanced electrochemical performance and attractive redox stability in a 5000 ppm H₂S-H₂. A-site deficient perovskite also showed high performance in solid oxide electrolyzer cell (SOEC) cathodes [39]. Ni-anchored La₀.₅Sr₀.₅TiO₃₋δ showed an approximately 20% better current efficiency than a bare cathode under an applied voltage of 2.0 V at 800 °C. For titanate co-doped with Mn and Ni, Ni metal can be exsolved to grow on the Mn-doped titanate [40]. This active catalyst showed the improved electrocatalytic activity and Faradaic efficiency for CO₂ electrolysis. Additionally, electrical conductivity is enhanced by the presence of metallic Ni. A great deal of research has been conducted recently on the exsolution for room temperature electrolysis. La₀.₄Sr₀.₄Ti₀.₉O₃₋δ [41] perovskites decorated with Ni nanoparticle have been reported to have catalytic activity in hydrogen evolution reaction (HER). This catalyst can work as a highly efficient HER electrocatalyst in basic media. As a bifunctional effect in the catalyst, the La₀.₄Sr₀.₄Ti₀.₉O₃₋δ perovskite promotes the dissociation of water and generation of hydrogen intermediates (H_ads), and the exsolved Ni nanoparticles facilitate the adsorption of hydrogen intermediates and generation of H₂. It was also possible to exsolve bimetallic nanoparticles such as Ni-Fe in A-site deficient perovskites. Sun et al [24] attempted to exsolve Ni-Fe bimetallic nanoparticles to improve the performance of a SOFC’s monometallic anode, and they found that adding Fe helped to create more oxygen vacancies, and promoted the material reducibility. Zhu et al [46] revealed that A-site deficiency is important to avoid B-site deficiency after exsolution. In that study, the oxide phase in reduced Sr₀.₉₅Ti₀.₃Fe₀.₆₃Ni₀.₀7O₃₋δ showed a well-ordered perovskite structure, whereas the greater B-site deficiency in reduced SrTi₀.₃Fe₀.₆₃Ni₀.₀7O₃₋δ resulted in a highly disordered and strained structure. Among transition metals other than Ni, Cu also can be exsolved. For example, (La₀.₇₅Sr₀.₂₅)₀.₉(Cr₀.₅Mn₀.₅)₀.₉Cu₀.₁O₃₋δ [42] can be used in CO₂ electrolysis by exsolving Cu on the surface of the perovskite. They observed that the current efficiencies of 85% were obtained with
Cu-decorated nanocatalysts for direct CO₂ electrolysis. Cu metal nanocatalysts were also effective in steam electrolysis [43]. The polarization resistance of (La₀.₂Sr₀.₈)₀.₅Ti₀.₅Cu₀.₁O₃₋₀.₆ [43] was improved to 1.5 Ω cm² at 800 °C in hydrogen compared to 3 Ω cm² for the bare sample. In addition, the current efficiency of the Cu-anchored cathode is approximately 20% better than that of the bare cathode. Research on exposition on Ag was carried out to develop SOFC cathode materials. Zhu et al reported that the exsolved Sr₀.₉₅Ag₀.₀₅Nd₀.₁Cu₀.₉O₃₋₀.₆ [44] electrode was very active for oxygen reduction reaction (ORR), achieving a very low area-specific resistance (≈0.214 Ω cm² at 500 °C). In addition to high-temperature applications, Ag has excellent catalytic activity in other energy storage and conversion applications. An Ag nanoparticle modified (La₀.₈Sr₀.₂)₀.₉₅MnO₃₋₀.₆ [45] was used as a cathode in dye-sensitized solar cells (DSSCs), displaying enhanced activity for the electrocatalytic triiodide (I⁻) reduction reaction (IRR) in DSSCs as compared to the Ag/perovskite hybrids synthesized by physical mixing and impregnation methods.

In addition to the non-stoichiometric approach for exsolution, studies on other variables have been carried out to control B-site exsolution. These include altering factors such as lattice strain, electrical polarization, and gas environment. Strain effect plays an important role in determining the degree of exsolution. In this study of strain effect, a model system using epitaxial thin film is used. Tuning surface polarization, and gas environment. Strain effect plays an important role in determining the degree of high electrical conductivity [46].

Electrocatalysts exhibit excellent catalytic properties, such as fast surface oxygen exchange, rapid oxygen ion diffusion, and cathode for CO₂. Because of its oxygen vacancy channel and structural characteristics, the double perovskite (AA’B₂O₅₋₀.₅) exhibits excellent catalytic properties, such as fast surface oxygen exchange, rapid oxygen ion diffusion, and high electrical conductivity [70, 71]. As shown in figure 5(b), the A and A’ sites are ordered along the c-axis, and oxygen vacancy is mainly formed in the A-site layer. In this regard, Sengodon et al reported PrBaMn₂O₅₋₀.₃ (PBMO) layered oxygen-deficient double perovskite as ceramic anode for direct hydrocarbon SOFC [1]. The PBMO double perovskite was fabricated through a two-step sintering process. Pr₀.₅Ba₀.₅MnO₃₋₀.₅ single perovskite with a mixture of cubic and hexagonal phases was synthesized in the air at 950 °C and then annealed in H₂ at 800 °C to fabricate the PBMO. The differential thermal analysis curve indicated an exothermic peak at approximately 400 °C in the reducing atmosphere, indicating that the phase transition from a single perovskite to a double perovskite occurred during the annealing process.

The phase transition under the reducing conditions leads to spontaneous exsolution without external control such as A-site defects. Additionally, the oxygen vacancy formation on the [PrO₂] layer also causes the exsolution of B-site transition metals [51]. Following the DFT using the Vienna ab initio simulations package (VASP) code, the double perovskite structure (−3.62 eV) has a much lower energy barrier for the exsolution of B-site atoms than a single perovskite (−4.82 eV). Therefore, the crystal reconstruction of perovskite is a favorable process for B-site cations exsolution. Indeed, MnO is known as having hard-to-reduce cations yet is also exsolved from NdBaMn₂O₅₋₀.₃ double perovskite under reducing conditions [48].

Furthermore, Kwon et al reported the contribution of various transition metals toward exsolution on a PrBaMn₁.₇T₀.₃O₅₋₀.₃ (T = Mn, Co, Ni, and Fe) double perovskite [47]. In their experimental results, Co, Ni,
Figure 8. (a) Schematic image of co-segregation in the layered perovskite. Pr, Ba, Mn, T (Mn, Co, Ni, and Fe) and O atoms are shown as grey, green, dark blue, purple and red, respectively. The inset red boxes indicate the co-segregation of B-cation with an oxygen vacancy. (b) Comparison of the co-segregation energy with the dopant materials. (a) and (b) Reproduced with permission from [47] Copyright 2017, Nature Publishing Group.

and Mn were exsolved from the double perovskite, but Fe showed no exsolution. Transition metals generally have different degrees of reduction, however, the reducibility is not enough to explain the degrees of exsolution. Because the exsolution is a complex phenomenon that occurs from the inside of the lattice to the surface, the parent materials characteristics should be considered. The exsolution mechanism in the double perovskite is best described as the formation of a Schottky-type defect, which can be expressed as:

$$T_{\text{Mn}}^{X} + O_{\text{O}}^{X} \leftrightarrow TO + V_{\text{O}}^{\text{oo}} + V_{\text{Mn}}^{\text{co}}$$

(3)

where $T_{\text{Mn}}^{X}$ is the B-site dopant in the Mn site with a zero net charge, $O_{\text{O}}^{X}$ is oxygen in the oxygen site with a zero net charge, $V_{\text{O}}^{\text{oo}}$ is the oxygen ion vacancy with a net charge of +2, and $V_{\text{Mn}}^{\text{co}}$ represents the cation vacancy in the Mn site with a net charge of −2.

Therefore, the co-segregation energy of B-site transition metals with oxygen vacancies was introduced to verify the exsolution trends in the double perovskite. As shown in the schematic image (figure 8(a)), the B-site cations co-segregate with nearby oxygen toward the surface. Based on the Mn used as host cation, Co and Ni have higher co-segregation energy, and Fe has lower co-segregation energy, which is well supported by experimental results (figure 8(b)). Meanwhile, PrBaMn$_{1.7}$Co$_{0.3}$O$_5$+$\delta$ and PrBaMn$_{1.7}$Ni$_{0.3}$O$_5$+$\delta$ with exsolved metal nanoparticles show higher electrochemical performance than PrBaMn$_2$O$_5$+$\delta$ and PrBaMn$_{1.7}$Fe$_{0.3}$O$_5$+$\delta$ with exsolved oxide nanoparticles. This is because the exsolved metal nanoparticles have excellent catalytic activity for fuel oxidation [15, 72]. Additionally, no significant degradation was observed at a constant voltage of 0.6 V at 700 °C in C$_3$H$_8$ fuel, meaning that the double perovskite with exsolved metal nanoparticles has a high carbon coking tolerance in a hydrocarbon atmosphere.

Various catalyst design studies have been conducted based on the exsolution trends in the PBMO double perovskite. First, Co and Ni were co-doped on the Mn site of PBMO to investigate an alloy exsolution [49]. Since Co and Ni have higher co-segregation energies than Mn, both were expected to exsolve in the PrBaMn$_{1.7}$Co$_{0.1}$Ni$_{0.2}$O$_5$+$\delta$ (PBMCNO). As shown in the high-angle annular dark field (HAADF) image of PBMCNO with energy-dispersive x-ray spectroscopy elemental mapping images (figure 9(a)), Co and Ni coexist on the surface of PBMCNO, which indicates that Co-Ni alloy nanoparticles are exsolved after reduction at 850 °C in H$_2$. The lattice distance of the alloy nanoparticle is 0.127 nm (figure 9(d)), which is consistent with the lattice spacing of the (220) planes of the Co-Ni metal alloy.

It is reasonable to think that to form exsolved Co-Ni alloy nanoparticles, the aggregation of two doped B-site cations and the segregation to the surface should occur under a reducing atmosphere. Thus, two possible mechanisms, ‘bulk alloy formation’ and ‘surface alloy formation’, were considered to understand the alloy exsolution mechanism. During bulk alloy formation, Co–O$_{\text{O}}$ (oxygen vacancy)–Ni aggregation occurs in the bulk of PBMCNO, after which Co–O$_{\text{O}}$–Ni segregates toward the surface. On the contrary, in the surface alloy formation, after Co–O$_{\text{O}}$–Co and Ni–O$_{\text{O}}$–Ni segregate to the surface, two metals with oxygen vacancy aggregate on the surface of PBMCNO. The Gibbs energies of aggregation in the bulk (bulk alloy
formation) and at the surface (surface alloy formation) are 0.02 eV and $-0.01\ eV$, respectively, demonstrating that surface alloy formation is thermodynamically more favorable. The in-situ x-ray diffraction (XRD) results also confirmed that after the metallic Co located at 44.25° and the metallic Ni located at 44.7° were observed at 700 °C under an H$_2$ atmosphere, the two peaks combined to form a single peak at a temperature above 750 °C. To illustrate the notable improvement in the catalytic activity of the exsolved Co–Ni alloy, the dry reforming of methane was conducted at 800 °C–900 °C in a fixed-bed reactor. The CO$_2$ conversion of PBMCNO with Co–Ni alloy nanoparticles is almost four times higher than that of PBMO with MnO nanoparticles at 900 °C, which verifies that the exsolved Co–Ni metal alloy improves the catalytic activity for hydrocarbon oxidation.

In the general exsolution process, significant amounts of cations remain in the host bulk during the reduction process due to the limited diffusion rate of metal cations [62, 73]. In addition, the exsolution can cause structural instability because of excessive cation loss in the host material [74]. A recent study was conducted to increase the number of exsolved nanoparticles and stabilize the bulk using the exsolution trend and topotactic ion exchange [50]. The topotactic ion exchange is a solid-state reaction that reorganizes the cation locations [75, 76]. As mentioned previously, exsolution in stoichiometric double perovskite (AA'B$_{2-x}$C$_x$O$_{5+\delta}$) leaves B-site vacancies. In contrast, topotactic ion exchange maintains stoichiometric double perovskite without B-site vacancies after exsolution, which is illustrated by the following equation.

$$\text{AA}'\text{B}_{2-x}\text{C}_x\text{O}_{5+\delta} + x\text{G} \rightarrow \text{AA}'\text{B}_{2-x}\text{G}_x\text{O}_{5+\delta} + x\text{C}$$  \hspace{1cm} (4)

where AA'B$_{2-x}$C$_x$O$_{5+\delta}$ represents the stoichiometric double perovskite, xG represents the guest cation, AA'B$_{2-x}$G$_x$O$_{5+\delta}$ represents a new double perovskite without B-site vacancy, and xC represents the exsolved metal.

In the study of topotactic ion exchange, PrBaMn$_{1.7}$Co$_{0.3}$O$_{5+\delta}$ and Fe were selected as the host oxide and the guest cation, respectively. This is because Co ($-0.55\ eV$) had higher co-segregation energy and Fe ($-0.15\ eV$) had lower co-segregation energy than Mn ($-0.47\ eV$) [47]. As a result of the topotactic ionic exchange between Co and Fe during the reduction process (figure 10), Fe entered the host material and Co exsolution occurred without B-site vacancies. To demonstrate the topotactic ion exchange process between host cation (i.e. Co and Mn) and guest cation (i.e. Fe), DFT calculation was conducted. The calculation of incorporation energy between B-site cations indicated that Co$\leftrightarrow$Fe ($-0.41\ eV$) is thermodynamically more favorable than Mn$\leftrightarrow$Fe ($0.22\ eV$). Therefore, as the amount of Fe guest cations increased, more Co cations were exsolved to the surface of the host material. The number of exsolved nanoparticles was thus verified to
increase with the amount of guest Fe cations owing to the topotactic ion exchange, which greatly improved the electrochemical performance of the SOFC anode. The topotactic ion exchanged anode showed a maximum power density of 1.834 W cm\(^{-2}\) in humidified H\(_2\), achieving the highest performance among the various recently developed SOFC anodes using exsolution [52, 77, 78].

2.4. Exsolution on Ruddlesden–Popper (RP) type perovskite

RP type perovskites have been widely researched as catalysts that can provide interesting physical and electrochemical properties because of their dimensionality effects and unique structure [79–82]. The RP structure consists of alternating perovskite and rock salt layers, which leads to fast ionic and electronic transport [83]. However, the conventional methods for fabricating RP perovskites require relatively high sintering temperatures (≥ 1350 °C), which reduces specific surface area and catalytic activity. To decrease the sintering temperature, Yang et al utilized the phase transition of perovskite in a reducing atmosphere [52]. As a result of the reduction process, RP type perovskite (Pr\(_{0.4}\)Sr\(_{0.6}\)Fe\(_{0.85}\)Nb\(_{0.15}\)O\(_7\)) with a Co-Fe alloy was formed from Pr\(_{0.4}\)Sr\(_{0.6}\)Co\(_{0.2}\)Fe\(_{0.7}\)Nb\(_{0.1}\)O\(_{3-\delta}\) cubic perovskite. As with the double perovskite, exsolution occurs simultaneously with phase transition from the single perovskite to the RP type perovskite under the reduction process [54, 56, 84, 85].

Furthermore, Du et al reported the phase transition of perovskite with the exsolution in detail [53]. As shown in figure 11, the Sr\(_2\)FeMo\(_{0.65}\)Ni\(_{0.35}\)O\(_{6-\delta}\) (SFMNi) perovskite is reduced to a Sr\(_3\)FeMoO\(_7\) \(_\delta\) RP phase, a Sr\(_{2}\)Fe\(_{1.7}\)Mo\(_{0.3}\)O\(_{3-\delta}\) cubic phase, and a Ni-Fe metallic alloy. The RP phase is a Sr\(_3\)FeMoO\(_7\) \(_\delta\) layered perovskite (A\(_1\)B\(_2\)O\(_7\)), consisting of the two layers of Sr\(_{2}\)Fe\(_{0.7}\)Mo\(_{0.3}\)O\(_{3-\delta}\) cubic perovskite (ABO\(_3\)) between the rock salt (AO) layers. The phase transition of the SFMNi can be expressed as:

\[
\text{Sr}_2\text{FeMo}_{0.65}\text{Ni}_{0.35}\text{O}_{6-\delta} \rightarrow \frac{1}{2}\text{SrFe}_{0.7}\text{Mo}_{0.3}\text{O}_{3-\delta} + \frac{1}{2}\text{Sr}_3\text{FeMoO}_{7-\delta} + \frac{1}{8}\text{Fe}_{1.2}\text{Ni}_{1.8} + \frac{1}{2}\text{O}_2 (g) \tag{5}
\]
Figure 11. Schematic images for structural transformation of $\text{Sr}_2\text{FeMo}_{0.65}\text{Ni}_{0.35}\text{O}_{6-\delta}$ in reduction process. Reproduced with permission from [53], Copyright 2016 American Chemical Society.

| Techniques                        | Depth resolutions | Description                                  |
|-----------------------------------|-------------------|----------------------------------------------|
| Low-energy ion scattering (LEIS)  | 1–2 atomic layers | Elemental composition and lattice arrangement at the outmost layer |
| Auger electron spectroscopy (AES) | Several nanometers | Elemental composition of material surface |
| Scanning electron microscopy (SEM) | Several nanometers | Microstructure and morphology |
| X-ray photoelectron spectroscopy (XPS) | Several nanometers | Local elemental composition and electronic state at the surface |
| Transmission electron microscopy (TEM) | Atomic-scale | Local elemental distribution with lateral resolution of the surface |
| Atomic force microscopy (AFM)     | Several nanometers | Surface morphology |

Table 2. Characterization techniques for the study of exsolution.

The transformed SFMNi anode with exsolved Ni-Fe alloy exhibited high catalytic activity toward the $\text{H}_2$ and $\text{CH}_4$ oxidation. In addition, a single-cell with an SFMNi anode showed no detectable degradation under a constant current density of 0.2 A cm$^{-2}$ in $\text{CH}_4$, indicating good carbon coking tolerance of the ceramic anode.

The RP perovskites have excellent phase stability and reversibility under a redox cycle [55, 57–59]. Chung et al [55] reported that RP structure of $\text{La}_{1.2}\text{Sr}_{0.8}\text{Mn}_{0.4}\text{Fe}_{0.6}\text{O}_4$ (RPLSMF) and $\text{in situ}$ B-site exsolution of Fe metal was obtained by $\text{in situ}$ reduction of $\text{La}_{0.6}\text{Sr}_{0.4}\text{Mn}_{0.2}\text{Fe}_{0.8}\text{O}_3$ (LSMF) single perovskite. In the XRD results, the RPLSMF with Fe metal returned to the LSMF with no detectable secondary phase after re-oxidation in air. When the re-oxidized material was annealed again under the hydrogen atmosphere, the RPLSMF of the same composition as the first reduced sample was reconstituted, confirming the reversibility of phase transition during the redox cycle.

3. Characterization techniques of exsolution

The advanced characterization techniques are essential to identify the origin of exsolution and control its properties since exsolution occurs within a few nanometers. In recent years, the technological development of the analysis tool has been intensively improved, so in this part, a brief overview of the tool analyzing the exsolution is provided and summarized in table 2.

Low-energy ion scattering (LEIS) spectroscopy is a technique that can detect the composition and structure of the outmost surface. The observation range of LEIS reaches one or two atomic layers of the surface, which is one of the most precise surface analysis methods. Using the primary ion beam, the information on composition, element, and lattice arrangement can be obtained from the energy difference with the reflected primary ion beam. For example, Kühner’s group studied surface chemistry of perovskite-based materials using LEIS [86]. They found that removal of the Sr-rich surface layer from $\text{La}_{0.6}\text{Sr}_{0.4}\text{CoO}_{3-\delta}$ could reduce the polarization resistance. Comprehensive studies have shown that LEIS appears to be a very useful characterization tool for analyzing the phenomenon of exsolution by providing profiling of elemental distribution in the near-surface regions of the perovskite oxides [86, 87].

Auger electron spectroscopy (AES) is a surface analysis technique that uses a high-energy electron beam to probe the composition of the material surface. This analysis technique collects information by detecting electrons emitted from higher energy levels called Auger electrons. Since this information includes the average of several atomic layers, exsolution information can be obtained in combination with depth profiling. Also, scanning electron microscopy (SEM) has been widely used to analyze surface morphology. This is one of the most intuitive devices to view particles after forming exsolved particles. Joo et al analyzed
by comparing the change in the number of exsolved particles of the SEM images to examine the correlation between the number of exsolved particles and electrochemical performance [50].

X-ray photoelectron spectroscopy (XPS) is a surface-sensitive quantitative spectroscopic technique that measures elemental composition, empirical formula, chemical state, and electronic state. The basic use is to obtain the oxidation state of B-site cation. Therefore, the metal nanoparticles exsolved on the surface and the cation state bound to oxygen in the bulk can be distinguished, thereby enabling quantitative analysis such as metal–cation ratio [88]. For example, it has been found that Ni is not present in the outermost atomic layer in the low-index (110) and (001) faces of a single crystal of La$_{1.67}$Sr$_{0.33}$NiO$_4$ + $\delta$ through a combination of analytical techniques with LEIS and XPS [87]. Recently, research on surface chemistry has been conducted through in situ XPS. Shao-Horn’s group analyzed the surface of La$_{1-x}$Sr$_x$CoO$_{3-\delta}$ thin film during heating and cooling cycles and found that the surface structure and chemistry are closely related to temperature [89].

Transmission electron microscopy (TEM) is an optical device where the electron beam passes through a specimen to create an image. Since the specimen is fixed on the grid and has a very thin thickness of 100 nm or less, electrons pass through interaction with the specimen. Energy-dispersive x-rays spectroscopy (EDS) is used for elemental and chemical analysis of samples. The combined use of TEM and EDS can provide information on local structures and compositions on surfaces, which can be very powerful tools for studying exsolution.

Atomic force microscopy (AFM) is a scanning probe microscope that uses the force between a probe and a sample surface atom. The surface image of the sample is formed by monitoring the change in the reflected laser caused by the tip bending as the tip scratches the surface. AFM does not require the conductivity of the sample nor the pretreatment of the sample which are required in SEM, therefore it can be used for analyzing various materials. Dragos et al have analyzed the dimension of exsolved particles by etching the sample surface [10].

4. Applications of exsolution

The exsolved nanoparticles on perovskite oxides have been intensively studied in the area of catalysis, electrocatalysis, and photocatalysis owing to their high catalytic properties and durability [39, 42, 50]. In particular, since the exsolution occurs in the reducing atmosphere at high temperatures, it has been widely used to design electrode materials in solid oxide cell (SOC) operated at 700 °C–850 °C [47, 49]. Exsolved metallic nanoparticles provide an excellent catalytic active site for fuel oxidation reaction in the SOC [47] and CH$_4$ conversion in the dry reforming reaction [50]. Besides, the exsolved composite oxides act as a superior catalyst for the HER, ORR, and oxygen evolution reaction (OER) [41, 90, 91]. Jiang et al reported that the improvement in electrochemical performance is mainly caused by the strong interaction and the electron transfer between the host perovskite oxide and exsolved metal nanoparticle [91]. In the photocatalysis, the exsolved metal nanoparticles are plasmonically active under visible light illumination, enhancing photocatalytic activity and the visible light absorption [92–94]. Excited free electrons within metal nanoparticles can be directly injected into the conduction band of perovskite oxide because of intimate contact between the metal and oxide, which participates in water reduction reactions to produce hydrogen [94].

5. Conclusion and outlook

Transition metal exsolution is a promising technique for designing catalysts in electrocatalysis and renewable energy. In the exsolution process, the metal cations exsolv from the parent oxide lattice to the oxide surface under a reducing atmosphere. Exsolved metal nanoparticles have high durability because of their unique feature of being socketed into the parent oxide surface. To understand the exsolution phenomena, parent oxide characteristics should be considered because the degree of exsolution is closely related to the oxide structure. Perovskite oxides (ABO$_3$) based materials have been mainly researched as the parent oxides for exsolution because of their excellent compositional structural flexibility. There are various structures in the perovskite oxides, each of which has a different main driving force for exsolution. In a single perovskite, harder-to-reduce cations can be exsolved by A-site deficiency of the perovskite, whereas in double and RP perovskites, phase transition under a reducing atmosphere spontaneously leads to exsolution without additional control. Those phenomena are demonstrated by advanced characterization techniques such as LEIS spectroscopy, AES, XPS, SEM, AFM, and TEM.

Until now, the exsolution phenomena in perovskite oxides have been widely studied, but there are still some fundamental open questions, such as how to maintain stable structure while the composition of the perovskite oxide changes from the stoichiometric to non-stoichiometric in the exsolution; what is the depth at which the exsolution occurs; how to achieve real time observation of the exsolution under operando...
conditions (e.g., the SOC operating atmosphere). Therefore, in-depth understanding of exsolution and well-controlled experimental conditions with the advanced characterization techniques are required to unravel the scientific questions. These studies may provide significant insights into fundamental understanding of exsolution as well as the exploration of perovskite-based catalysts at various operating conditions of energy devices.

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