Phase equilibria in the YFeO$_3$ – YCoO$_3$ system in air

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Abstract

YFe$_{1-x}$Co$_x$O$_3$ solid solutions were prepared by glycerol-nitrate technique. The homogeneity range of solid solutions was studied within the temperature range 1173 – 1573 K. A continues series of solid solution below the decomposition temperature of YCoO$_3$, which was shown to be equal to 1266 ± 6 K, begins to narrow at higher temperatures and becomes equal to 0 ≤ x ≤ 0.1 at 1573 K. The phase diagram of the YFeO$_3$ – YCoO$_3$ system in the “T – composition” coordinates was divided into three fields. Similar to the parent ternary oxides, all single-phase YFe$_{1-x}$Co$_x$O$_3$ solid solutions possess orthorhombically distorted perovskite structure (Pnma space group). Unusual behavior of orthorhombic distortions in YFe$_{1-x}$Co$_x$O$_3$ with temperature was explained by probable changes in spin state of Co$^{3+}$ ions.

1. Introduction

Yttrium ferrite and yttrium cobaltite with the perovskite structure and their mixed derivatives attract much attention due to a set of electrical and magnetic properties [1-12]. Although their structural features and some functional properties were widely studied [1-16], there is lack information concerning phase equilibria and homogeneity ranges of YFe$_{1-x}$Co$_x$O$_3$ solid solutions. It is worth noting that phase equilibria in the Y – Fe – O system as well as thermodynamic characteristics of ternary oxides inside including YFeO$_3$ were studied in detail [17-21]. Also, detailed information available for the Fe – Co – O system [22-24]. Much less information can be found for the Y – Co – O system and the thermal stability of YCoO$_3$ [16, 25]. Thus, the aim of present work was determination of homogeneity ranges of YFe$_{1-x}$Co$_x$O$_3$ solid solutions as a function of temperature in air and establishing of phase equilibria in the YFeO$_3$ – YCoO$_3$ system.

2. Experimental

The samples were prepared via the glycerol-nitrate technique. The starting materials – Y$_2$O$_3$, metallic Co, FeC$_6$O$_{12}$·2H$_2$O – were dissolved in nitric acid and then glycerol was added. Metallic cobalt was prepared from CoO$_4$ by reduction in hydrogen flow at 923 K. The solution was carefully heated to dryness. The obtained residue was slowly heated up and annealed at 1173 K for 20 h. Final annealing of the samples were performed at required temperatures in air for 96 h with grinding after each 12 h. The samples were quenched to room temperature (RT) by removing them from a furnace and placing them to a cold massive copper plate. Phase identification was carried out by means of X-ray powder diffraction (XRD) using a Shimadzu XRD 7000 diffractometer (Cu Kα radiation, 2θ = 20°-90°, 0.02 deg/min, 5 s/point). High temperature XRD measurements were performed using a HTK 1200N (Anton Paar, Austria) high temperature chamber installed at the diffractometer. Unit cell parameters were calculated using Cefref 3 software. The structure was refined by full-profile Rietveld analysis using Fullprof 2017 software.

TGA measurements were performed using a STA 409 PC instrument (Netzsch) within the temperature range of 300 – 1373 K in air.

3. Results and Discussion

First, the parent oxides of the studied system YFeO$_3$ and YCoO$_3$ were prepared and examined. Yttrium ferrite quenched from high temperature within the entire range (1173 – 1473 K) or slowly cooled to RT possesses orthorhombically distorted perovskite structure, which is in good agreement with the results reported earlier [1-4]. Fig. 1 illustrates XRD patterns of YFeO$_3$ prepared at various conditions and evaluated values of unit cell parameters, as an example. The XRD pattern refined by the Rietveld method and structural model of YFeO$_3$ designed...
using the “Diamond 3.2” software is shown in Fig. 2.

Single-phase yttrium cobaltite YCoO$_3$ was obtained only at relatively low temperatures 1173 and 1223 K. Like ferrite it possesses the orthorhombic structure (SG Pnma). The structural parameters of YCoO$_3$ quenched from 1173 and 1223 K refined by the Rietveld method are listed in Table 1.

To preliminarily estimate the decomposition temperature of YCoO$_3$, TGA measurements in a dynamic mode with a heating rate of 3.2 K/min were performed. A sharp drop in the mass of the sample was detected at 1300 K (Fig. 3).

Table 1 The structural parameters of YCoO$_3$ quenched from 1173 and 1223 K refined by the Rietveld method

| T, K | 1173 | 1223 |
|------|------|------|
| a, Å | 5.41856(5) | 5.41784(5) |
| b | 7.36230(7) | 7.36195(7) |
| c, Å | 5.13591(5) | 5.13604(5) |
| V, (Å$^3$) | 204.811(1) | 204.855(3) |
| x(Y) | 0.4316(2) | 0.4315(2) |
| z(Y) | 0.0183(3) | 0.0184(3) |
| x(01) | 0.5287(1) | 0.5285(2) |
| z(01) | 0.5956(2) | 0.5972(1) |
| x(02) | 0.2002(1) | 0.1988(1) |
| y(02) | 0.0487(8) | 0.0487(8) |
| z(02) | 0.3051(2) | 0.3047(1) |
| d$_{10-101}$, Å | 2.234(8) | 2.227(7) |
| d$_{10-111}$, Å | 2.437(6) | 2.439(6) |
| d$_{11-202}$, Å | 2.266(6) | 2.263(6) |
| d$_{11-212}$, Å | 2.558(6) | 2.557(6) |
| d$_{00-200}$, Å | 1.911(2) | 1.913(2) |
| d$_{00-202}$, Å | 1.939(6) | 1.933(6) |
| d$_{00-210}$, Å | 1.941(7) | 1.949(6) |
| R$_{wp}$, % | 10.7 | 13.2 |
| R$_{exp}$, % | 12.3 | 15.5 |
| R$_{exp}$, % | 9.18 | 11.3 |
| R$_{exp}$, % | 3.58 | 3.10 |
| R$_{exp}$, % | 6.85 | 3.11 |
| $\chi^2$ | 1.783 | 1.877 |

Fig. 1 XRD patterns of YFeO$_3$ quenched from various temperatures in air: (a) 1373 K; (b) 1173 K and (c) slowly cooled to room temperature and unit cell parameters evaluated by the Rietveld refinement

To refine the decomposition temperature in TGA measurements, a static mode was used. The following protocol was used: a single-phase sample was heated at a rate of 1 K/min to 1110 K and equilibrated at this temperature for 8 h. Then the temperature was increased in a step of 20 K and the sample was kept at a fixed temperature until a constant mass was established. No significant mass changes were detected at $T \leq 1260$ K. The next step to 1280 K results in a dramatic weight loss.

XRD analysis of the sample quenched after annealing at 1273 K, which was originally a single-phase YCoO$_3$, showed the presence of significant amounts of yttrium oxide and cobalt oxide (II) as secondary phases (Fig. 4). It should be mentioned that small amount of Co$_3$O$_4$ forms due to partial oxidation of CoO while cooling since latter is thermodynamic stable form of cobalt oxide at 1273 K in air.

Thus, one can conclude that YCoO$_3$ decomposes according the reaction

$$YCoO_3 = \frac{1}{2} Y_2O_3 + CoO + \frac{1}{4} O_2 \quad (1)$$

within the range 1260 < $T_{dec}$, K < 1273. This allows us to evaluate $T_{dec}(YCoO_3) = 1266 \pm 6$ K.

Fig. 2 XRD pattern refined by the Rietveld method and structural model of YFeO$_3$ along the b axis designed using the “Diamond 3.2” software.

Fig. 3 TGA curve for single-phase YCoO$_3$ in air measured in a dynamic mode with a heating rate of 3.2 K/min.
Prolonged annealing of the sample with nominal composition corresponding to YCoO$_3$ at 1373 K reveals coexistence of two binary simple oxides Y$_2$O$_3$ and CoO.

Since YCoO$_3$ is only stable below 1266 K, it is likely that a continuous series of YFe$_{1-x}$Co$_x$O$_3$ solid solutions cannot be obtained at higher temperatures. Indeed, continuous series of YFe$_{1-x}$Co$_x$O$_3$ solid solutions in the range of $0 \leq x \leq 1$ was obtained at 1173 K. The homogeneity range at 1273 K was evaluated as $0 \leq x \leq 0.9$. The sample with $x=0.95$ contained together with perovskite phase also Y$_2$O$_3$ and enriched by cobalt Fe$_{1-x}$Co$_x$O with the rock salt structure (Fig. 5). Further increase of temperature leads to a decrease in Co content in the limiting YFe$_{1-x}$Co$_x$O$_3$ solid solution (Table 2).

Thermal decomposition of Co-saturated solid solution with the temperature increase can be shown by following reaction:

$$\text{YFe}_{1-x}\text{Co}_x\text{O}_3 \rightarrow \text{YFe}_{1-x}\text{Co}_x\text{O}_3 + \text{Co}_{1-y}\text{Fe}_y\text{O} + Y_2\text{O}_3 + 2\text{O}_2$$

(2)

were $x' > x''$ and $y'$ corresponds to the Fe-saturated solid solution at a fixed temperature. It is worth noting that the process described by scheme (2) differs significantly from the one occurs according to equation (1). The latter corresponds to a nonvariant thermodynamic equilibrium, when all participating phases coexist at fixed $T$ and $P_0$. In contrast, scheme (2) represents the situation when Co-saturated single-phase YFe$_{1-x}$Co$_x$O$_3$ solid solutions in the left-hand side exist at $T'$, and an increase of temperature to $T'' = T' + \Delta T$ causes a depletion of cobalt in solid solution and displacement of its composition YFe$_{1-x'}$Co$_{x''}$O$_3$ as well as appearance of two secondary phases, namely, Y$_2$O$_3$ and Co$_{1-y'}$Fe$_{y'}$O. The left-hand side and the right-hand side in the scheme (2) represent the phase composition in the system at different temperatures, $T'$ and $T'' = T' + \Delta T$, respectively. Thus, scheme (2) describes nonequilibrium process that occurs due to the change in thermodynamic parameter, in this case it is temperature. A similar process can take place at fixed temperature due to the decrease in $P_0$.

Based on the results of phase composition of all studied samples the “T-composition” phase diagram of the YFeO$_3$ – YCoO$_3$ system in air was drawn (Fig. 6).

All single-phase YFe$_{1-x}$Co$_x$O$_3$ solid solutions quenched from all studied temperatures possess the orthorhombic structure, like parent ternary oxides. The influence of temperature on the crystal structure of YFe$_{1-x}$Co$_x$O$_3$ ($x = 0.35$ and $0.45$) was studied by in situ high temperature (HT) XRD measurements. The structural parameters of YFe$_{1-x}$Co$_x$O$_3$ ($x = 0.35$ and $0.45$) at various temperatures refined by the Rietveld method are listed in Tables 3 and 4.

Temperature dependencies of the unit cell parameters and unit cell volume demonstrate visible non-linearity (Fig. 7). As a rule, the distortions of crystal structure tend to decrease with the temperature rise; however, the ortho-

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**Table 2** The homogeneity range value for the YFe$_{1-x}$Co$_x$O$_3$ solid solutions at various temperatures

| $T$, K  | Homogeneity range | $T$, K  | Homogeneity range |
|---------|-------------------|---------|-------------------|
| 1173    | $0 \leq x \leq 1$ | 1373    | $0 \leq x \leq 0.9$ |
| 1273    | $0 \leq x \leq 0.9$ | 1423    | $0 \leq x \leq 0.3$ |
| 1223    | $0 \leq x \leq 0.68$ | 1473    | $0 \leq x \leq 0.2$ |

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**Fig. 4** XRD patterns of the samples with nominal composition of YCoO$_3$ fired and quenched from various temperatures in air: (a) 1373 K; (b) 1323 K; (c) 1273 K; (d) 1223 K; (e) 1173 K.

**Fig. 5** XRD patterns for YFe$_{1-x}$Co$_x$O$_3$ ($x=0.9$, 0.95) equilibrated at 1473 K and quenched to RT.

**Fig. 6** Phase diagram of the YFeO$_3$ – YCoO$_3$ system in air.

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Table 4 The structural parameters of YFe₆₋₁ₓCoₓO₁₉ at various temperatures refined by the Rietveld method using HT-XRD measurements

| Table 4 | The structural parameters of YFe₆₋₁ₓCoₓO₁₉ at various temperatures refined by the Rietveld method using HT-XRD measurements |
|---|---|
| **SG Pnma : Y(4c)(x; 0.25; z); Fe/Co(4a)(x; 0; o); O₁(4c)(x; 0.25; z); O₂(8d)(x; y; z)** |
| **T, K** | 298 | 473 | 623 | 673 | 973 | 1273 | 1373 |
| **a, Å** | 5.51229(9) | 5.52600(9) | 5.53528(9) | 5.55559(9) | 5.60219(9) | 5.61552(9) |
| **b, Å** | 7.48356(13) | 7.51456(13) | 7.52654(13) | 7.54977(13) | 7.60614(13) | 7.62717(13) |
| **c, Å** | 5.21895(9) | 5.23834(9) | 5.24821(9) | 5.26589(9) | 5.30639(9) | 5.32121(8) |
| **V, (Å³)** | 215.218(2) | 217.524(4) | 218.648(7) | 220.869(6) | 226.111(7) | 227.910(7) |
| **x(Y1)** | 0.43423(4) | 0.4433(4) | 0.4343(3) | 0.4345(3) | 0.4349(3) | 0.4356(3) |
| **y(Y1)** | 0.0178(4) | 0.0170(4) | 0.0168(5) | 0.0166(4) | 0.0164(5) | 0.0158(5) |
| **z(O1)** | 0.3528(18) | 0.3519(18) | 0.3504(18) | 0.3516(18) | 0.3532(18) | 0.3530(19) |
| **z(O2)** | 0.6004(19) | 0.5999(20) | 0.5984(20) | 0.6031(20) | 0.6030(20) | 0.602(2) |
| **Rₚ, %** | 11.8 | 12.2 | 12.5 | 12.9 | 13.4 | 13.8 | 14.0 |
| **Rwp, %** | 15.0 | 15.2 | 15.2 | 15.7 | 15.8 | 16.0 | 16.2 |
| **Rp, %** | 10.6 | 10.6 | 10.7 | 10.7 | 11.0 | 11.3 | 11.4 |
| **R1, %** | 4.25 | 4.83 | 4.98 | 4.91 | 5.01 | 5.38 | 5.09 |
| **R₁, %** | 3.43 | 3.87 | 4.05 | 3.89 | 4.26 | 4.80 | 4.91 |
| **χ²** | 1.985 | 2.057 | 2.036 | 2.148 | 2.043 | 2.002 | 2.019 |
rhombic distortion parameters calculated by the formula [26, 27]:

$$D_{\text{orth}} = \frac{1}{3} \sum_{i=1}^{3} |a_i - \bar{a}| \times 100\%$$  \hspace{1cm} (3)

where $D_{\text{orth}}$ is the orthorhombic distortion parameter, %; $\alpha_1 = a_1$; $\alpha_2 = b$; $\alpha_3 = c/\sqrt{2}$ and $\bar{a} = (a \times b \times c/\sqrt{2})^{1/3}$, exhibit visible anomalies within 600 – 1200 K (Fig. 8). Such behavior could not be explained by the change in oxidation states of 3d metals. TGA analysis of $\text{YFe}_{1-x}\text{Co}_x\text{O}_3$ ($x = 0.35$ and 0.45) reveals tiny mass changes while heating, which means that both oxides possess almost stoichiometric oxygen content within entire temperature range. Thus, the oxidation state of 3d atoms is equal to 3+ and remains unchanged with the increase of temperature. Another reason that can induce structural transformations can be changes in spin states. Such temperature-induced changes in the spin state of Co$^{3+}$ ion in LnCoO$_3$ perovskites were reported by Raccach and Goodenough [28] and in later publications [7, 29–31]. Low-spin ($t^6_{2g}$) state can transforms into intermediate-spin ($t^4_{2g}e^1_{g}$) state and finally into high-spin ($t^4_{2g}e^2_{g}$) state. Although the possibility of spin state changes for Fe$^{3+}$ ion is still questionable and its high-spin state is more favorable [2, 5], possible spin-state transition of Co$^{3+}$ ion can cause the observed anomalies. However, this needs to be checked by further independent experiments.

It is worth noting that Co substitution for Fe reveals much stronger effect on changes in the unit cell volume due to the size effect ($r_{\text{Fe}^{3+}(\text{HS})} = 0.645\, \text{Å}$, $r_{\text{Co}^{3+}(\text{LS})} = 0.545\, \text{Å}$, $r_{\text{Co}^{3+}(\text{HS})} = 0.61\, \text{Å}$ [32]) rather than temperature. The decrease in the unit cell volume of $\text{YFe}_{1-x}\text{Co}_x\text{O}_3$ with the cobalt content (Fig. 9) is much more significant in comparison with the temperature dependence. 

![Fig. 7](image_url) The unit cell parameters and unit cell volume of $\text{YFe}_{1-x}\text{Co}_x\text{O}_3$ ($x = 0.35$ and 0.45) versus temperature

![Fig. 8](image_url) The pseudo-cubic unit cell parameter ($a_p$) and the orthorhombic distortion parameter ($D$) for $\text{YFe}_{1-x}\text{Co}_x\text{O}_3$ ($x = 0.35$ and 0.45) versus temperature

![Fig. 9](image_url) The unit cell volume of $\text{YFe}_{1-x}\text{Co}_x\text{O}_3$ versus Co content ($x$)
4. Conclusions

The homogeneity range and crystal structure of YFe$_x$Co$_{1-x}$O$_3$ solid solution have been studied within the entire composition range (0 ≤ x ≤ 1) at temperatures up to 1000 K. Partial substitution of Co for Fe has not changed the orthorhombic perovskite structure. Possible change in the spin state of Co$^{2+}$ ions is a reasonable assumption for the unusual behavior of orthorhombic distortions in YFe$_x$Co$_{1-x}$O$_3$ (0.35 ≤ x ≤ 0.45) with temperature.

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