



# Characterization of NiAl with cobalt produced by combustion synthesis

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## ABSTRACT

In this study, as an alloying element the effect of cobalt addition by 2.5 wt.% and 5 wt.% on the NiAl intermetallic compound produced by pressure-assisted combustion synthesis method was investigated. As starting materials aluminum powder with 15  $\mu\text{m}$  size, carbonyl-nickel 4–7  $\mu\text{m}$  size and cobalt powders 10–44  $\mu\text{m}$  size having 99%, 99.8% and 99.9% purity, respectively were used. The formation temperature of intermetallic compound for three different powders mixture determined by DSC analysis was approximately 654 °C. The production of NiAl was carried out in electrical resistance furnace in open air with a uniaxial pressure of 150 MPa at 1050 °C for 60 min. Optical and SEM studies showed that intermetallic compounds have low porosity and phase transformation has been completed. The distribution of alloying elements was confirmed by EDS analysis. The presence of NiAl and CoAl phases was determined by XRD analysis. The relative density of pure NiAl, NiAl + 2.5 wt.% Co and NiAl + 5 wt.% Co materials was 99.65%, 99.48% and 99.30% and the microhardness of materials was about 368 HV<sub>1.0</sub>, 398 HV<sub>1.0</sub> and 425 HV<sub>1.0</sub>, respectively.

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## 1. Introduction

The Ni–Al system is of particular interest. The strongly ordered B2 intermetallic NiAl has a number of unique physical and mechanical properties that make it attractive for use at high temperature and in aggressive environments due to its high-melting point (1640 °C), low density ( $\rho = 5.89 \text{ g/cm}^3$ ), excellent corrosion and oxidation resistance, combined with its ability to retain strength and stiffness at elevated temperatures [1,2]. However, poor room temperature strength is a significant obstacle to the alloy's structural use. Alloying behaviors of various elements in NiAl can be summarized the effects of transition metals on NiAl. The classified those elements as three types. Ti, Zr, Hf, Nb and Ta (type A) have low solubility in NiAl and exist in the form of ternary intermetallic compounds such as Heusler ( $\text{Ni}_2\text{Al}_x$ ) phase and Laves ( $\text{NiAl}_x$ ) phase. These ternary compounds can increase NiAl's creep strength dramatically. V, Cr, Mo and W (type B) can form pseudo-binary eutectic systems with NiAl and the eutectic structure improves NiAl's room temperature toughness. Fe, Co and Cu (type C) have high solubility in NiAl. These additions can be used to generate a second ductile phase to improve the NiAl alloy's room temperature ductility [3,4].

The characteristic features of the Ni–Al–Co system are complete solid solution series between B2–NiAl and B2–CoAl and between Ni and Co [5]. Co atoms replace the Ni lattice sites because exceptionally small formation energy is required for Co inserted into

Ni sublattice its also easier to create vacancies in the Ni sublattice than in Al sublattice [6]. The martensitic phase transformation from the ferromagnetic state in Co–Ni–Al  $\beta$  alloy system has been found by the author's group [7–9]. The shape memory alloys are widely noticed because of their good hot/cold workability.  $\beta$  base Co–Ni–Al alloys have the B2  $\rightarrow$  L1<sub>0</sub> phase transformation, the Curie and the martensitic transition temperatures of the  $\beta$  phase increase and decrease with increasing the Co content, respectively [7,8]. Chen and Han [10] have added Co into NiAl and show that the compressive strength and compressive ductility of the stoichiometric NiAl have been greatly improved by adding proper amount of Co.

NiAl intermetallics have several potential applications including turbochargers, high-temperature dies and moulds, furnace fixtures, rollers in steel slab heating furnaces, hydroturbines, cutting tools, pistons and valves and various components within gas turbines [11]. There are many ways in order to produce intermetallic compounds such as self-propagating high-temperature synthesis, powder processing, casting and combustion synthesis [12]. One of the simple, fast and economic production process combustion synthesis (CS) process is based on the concept that an exothermic wave propagating in a self-sustained manner through a heterogeneous medium, is used to synthesize different advanced materials [13–17]. There are two modes by which CS can occur: self-propagating high-temperature synthesis (SHS) and volume combustion synthesis (VCS). In both cases, reactants may be pressed into a pellet, typically cylindrical in shape. The samples are then heated by an external source (e.g., tungsten coil, laser) either locally or uniformly to initiate an exothermic reaction [17,18]. On the other hand, the problem such as porosity has prohibited the

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wide spread use of combustion method [12,19]. The formation of porosity can be eliminated by careful control over the reaction and/or through the application of external pressure during CS [20].

The main aim of present study is to investigate the effect of cobalt on properties of NiAl intermetallics manufactured by pressure-assisted combustion synthesis.

## 2. Experimental details

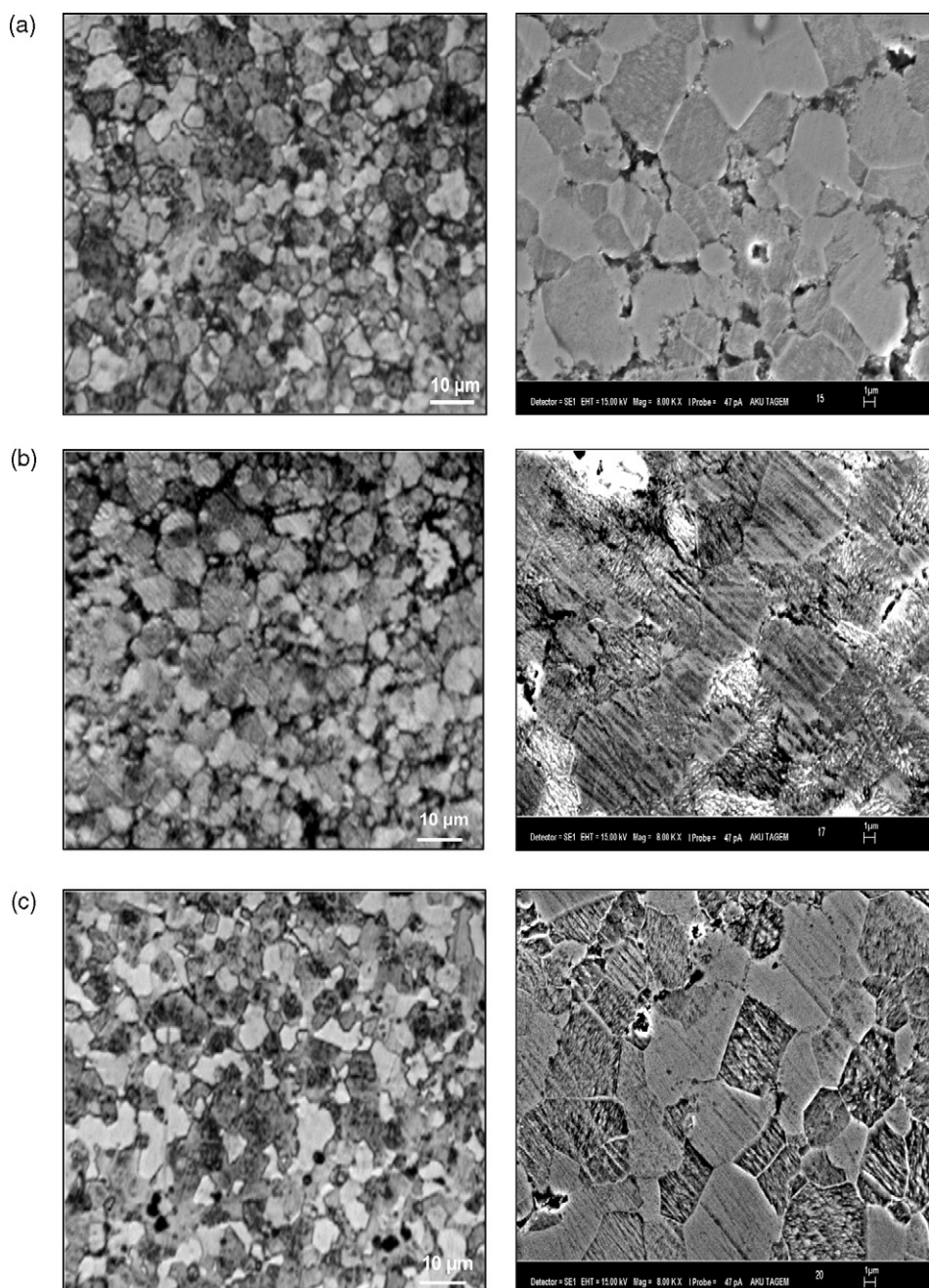
### 2.1. Production of test materials

In order to manufacture intermetallic materials as starting materials gas-atomized aluminum powder with 15  $\mu\text{m}$  size, carbonyl-nickel powder 4–7  $\mu\text{m}$  size and cobalt powder 10–44  $\mu\text{m}$  size having 99%, 99.8% and 99.9% purity, respectively were used. Ni and Al powders were mixed in stoichiometric ratio corresponding to the NiAl intermetallic phase, in a molar proportion of 1:1. The powders including 2.5 wt.% Co and 5 wt.% Co as alloying element were mixed in a ball mill for 10 min. in Ar + 3% H<sub>2</sub> gas medium with the addition of small amount of 0.1 ethanol. Prior

to sintering, the mixture was cold pressed into a cylindrical compact in a metal die coated with a thin layer of boron nitride under a uniaxial pressure of 150 MPa. The diameter and height of the compact samples were 15 mm and 5 mm, respectively. The compacts were synthesized and simultaneously consolidated in open atmosphere. Then the samples were heated at a heating rate of 20 °C min<sup>-1</sup> to sintering temperature, 1050 °C, and hold at this temperature for 60 min. Process was carried out without using vacuum and inert gas under a uniaxial pressure of 150 MPa. The samples were removed from the furnace having 900 °C in normal atmosphere then cooled to room temperature.

### 2.2. Characterization

Thermal analysis was performed by differential scanning calorimeter (DSC) to determine the reaction sequence and the ignition point in nitrogen atmosphere and at a heating rate of 20 °C min<sup>-1</sup> on consisting of three different powders mixture. The relative and bulk densities of the synthesized samples were measured in terms of Archimedes' method. The samples were polished and etched using a solution of HNO<sub>3</sub> (33.3 vol.%), CH<sub>3</sub>COOH (33.3 vol.%) and HCl (33.3 vol.%). The presence of NiAl, NiAl + 2.5 wt.% Co and NiAl + 5 wt.% Co materials was determined using Rigaku X-



**Fig. 1.** Optical (left side) and SEM (right side) views of (a) NiAl, (b) NiAl + 2.5 wt.% Co, and (c) NiAl + 5 wt.% Co.

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