



Isothermal Sections of the Ni-Cr-Ta Ternary System at 1200 °C and 1300 °C

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Abstract: Two isothermal sections of the Ni-Cr-Ta ternary system at 1200 °C and 1300 °C have been determined by using electron probe microanalysis, energy dispersive spectroscopy and differential scanning calorimeter. A Laves phase (Ni, Cr)₂Ta(HT)(C14 structure) with large solid solubility stabilized by the Ni addition was determined in both two isothermal sections. The composition range of this phase was about 25.8–66.0 at.% Cr, 2.5-44.3 at.% Ni, and 24.0-40.0 at.% Ta at 1200 °C, which increased with raising temperature. The melting point of the Ni-Cr alloys decreased with the addition of Ta. No ternary compound was found in both these two isothermal sections. The present work could be significant for practical application of nickel-based alloys and future thermodynamics assessment of the Ni-Cr-Ta ternary system.

Keywords: Ni-Cr-Ta system; isothermal section; electron probe microanalysis

1. Introduction

Nickel-based superalloys have been applied in the aerospace field due to their excellent high-temperature properties, oxidation and corrosion resistance in the extreme harsh environment [1,2]. However, with higher industrial requirements in the structure materials for high-temperature applications in aviation field, materials capable of better mechanical strength, oxidation and corrosion resistance are required. In order to improve the properties of Ni-based alloys, an excellent alternative is to alloy refractory elements [3–6]. Technologically, Cr addition could significantly improve the oxidation and hot-corrosion resistance for the nickel-based alloys by forming a stable oxidation protective layer Cr_2O_3 at elevated temperatures [4,7–9]. Meanwhile, as a solid solution strengthening element, the alloying of Ta also improves the hot-corrosion and oxidation resistance [10,11]. However, the stabilization of topologically close packed (TCP) phase will deteriorate the mechanical properties of the superalloys for excessive addition of Cr and Ta elements [12,13]. Therefore, it is of significant necessity to investigate the phase diagram of the ternary Ni-Cr-Ta system, not only for the future thermodynamics assessment, but also enhancing the potential practical applications.

The Ni-Cr-Ta ternary system consisting of three binary subsystems, Ni-Cr, Ni-Ta, Cr-Ta, is illustrated in Figure 1. In 1986, Nash [14] reviewed the Ni-Cr system with a eutectic reaction at 1345 °C, where an extensive Ni terminal solid solution (face centered cubic) region and a less extensive Cr terminal solid solution (body centered cubic) region were identified. Additionally, Lee [15] and Zhu et al. [16] re-evaluated the Ni-Cr binary system, which is in agreement with the experimental results with the work of Nash.



The Ni-Ta binary system has been investigated by many researchers [17–30]. In 2018, Zhou et al. [30] reviewed the Ni-Ta system, where only four intermetallic compounds Ni₂Ta, Ni₃Ta, NiTa, NiTa₂, and two terminal solid solutions fcc-(Ni), bcc-(Ta) coexisted, and the Ni₈Ta was confirmed to be a metastable phase. There are two eutectic reactions, three peritectic reactions, a peritectoid reaction and a congruent transformation in this system. Two eutectic reactions $L \rightarrow Ni_3Ta + fcc-(Ni)$ and $L \rightarrow Ni_2Ta + Ni_6Ta_7$ occurred at 1330 °C and 1350 °C, respectively. The Ni₆Ta₇ and NiTa₂ phase formed from two peritectic reactions. In addition, several investigators [23–28] have assessed the thermodynamic database of this binary system with CALPHAD method and first-principles calculation.



Figure 1. Binary phase diagrams constituting the Ni-Cr-Ta ternary system [14,30,31].

In 1987, Venkatraman et al. [32] reviewed the Cr-Ta system, in which two terminal solid solutions bcc-(Cr), bcc-(Ta) and intermediate phase Cr_2Ta formed from eutectic reaction occurred at 1760 and 1965 °C, respectively. The intermediate compound Cr_2Ta exhibits two Laves phase modifications. The high-temperature form, Cr_2Ta (HT), has the hexagonal MgZn₂-type (C14) structure, while the low-temperature form, Cr_2Ta (LT), has the cubic MgCu₂-type (C15) structure. In 1996, Okamoto [33] had redrawn this binary phase diagram based on the Venkatraman's work with an adjustment in the form of Cr_2Ta solidus complying with the Gibbs-Konovalov rule. In 1991, Kaufman et al. [23] assessed the Cr-Ta system with CALPHAD approach, then, Dupin et al. [34] re-evaluated the thermodynamic information based on the system using first-principles calculation and re-optimized the phase diagram with CALPHAD method.

As for the Ni-Cr-Ta ternary system, Nash et al. [35] investigated the phase equilibria in the Ni-rich portion of this system at 1000 °C and 1250 °C to establish a ternary Laves phase NiCrTa (the lattice parameter, a = 4.844 Å, c = 7.89 Å, annealed at 1250 °C, and a = 4.885 Å, c = 7.888 Å, annealed at 1000 °C) with hexagonal MgZn₂-type (C14) structure using electron microprobe and X-ray diffraction analysis. In 1985, Schittny et al. [36] reconfirmed the ternary compound NiCrTa in the partial isothermal section

at 1000 °C with concentration range of 0–40 at. % Ta. However, there is no ternary compound except for a Laves phase Cr_2Ta (hexagonal, MgZn₂-type, a = 4.844 Å, c = 7.9091 Å) in the isothermal section at 1100 °C, according to Nikolaev's et al. experimental phase diagram [37]. Additionally, Dupin et al. assessed the thermodynamic database of the Ni-Cr-Ta system [38]. The stable phases in the ternary Ni-Cr-Ta system are listed in the Table 1.

System	Phase	Pearson Symbol	Prototye	Space Group	Strukturberid	cht Ref.
Ni-Cr	fcc	cF4	Cu	Fm-3m	A1	[14]
	bcc-(Cr)	cI2	W	Im-3m	A2	[14]
	Ni ₂ Cr	oP6	Pt ₂ Mo			[14]
Ni-Ta	fcc	cF4	Cu	Fm-3m	A1	[30]
	bcc-(Ta)	cI2	W	Im-3m	A2	[30]
	Ni ₃ Ta	tI8	TiAl ₃	I4/mmm	D0 ₂₂	[30]
		mP16	NbPt ₃	P2 ₁ /m		[30]
		oP8	Cu ₃ Ti	Pmmn	$D0_a$	[30]
	Ni ₂ Ta	tI6	MoSi ₂	I4/mmm	C11 _b	[30]
	Ni ₆ Ta ₇	hP13	Fe ₇ W ₆	R-3m	D85	[30]
	NiTa ₂	tI12	Al ₂ Cu	I4/mcm	C16	[30]
Cr-Ta	bcc-(Cr)	cI2	W	Im-3m	A2	[31]
	bcc-(Ta)	cI2	W	Im-3m	A2	[31]
	Cr ₂ Ta(HT)	hP12	MgZn ₂	P63/mmc	C14	[31]
	$Cr_2Ta(LT)$	cF24	MgCu ₂	Fd-3m	C15	[31]

Table 1. The stable solid phases in the Ni-Cr-Ta ternary system.

2. Experimental Details

High purity metals nickel (99.9 wt. %), chromium (99.9 wt. %) and Tantalum (99.9 wt. %) were used as our raw material to obtain alloys. All the metals were well cleaned to avoid the input of impurity surface oxidation before melting. All alloys were displayed in the form of atomic ratios (at. %). The ingots, around 20 g, were re-melted at least four times to get the uniformity with less than 0.5 wt. % weight loss. The alloys were melted in a high purity argon atmosphere arc furnace with a non-consumable tungsten electrode on a water-cooled copper platform. Then, all specimens were individually sealed in silica capsules with high purity argon, annealed at 1200 °C for 35 days and 1300 °C for 15 days, respectively. Additionally, in order to prevent oxidation, we put some pure yttrium fillings in the quartz capsules. Some alloys with liquid phase at 1300 °C were wrapped in the pure tantalum foil to avoid contact reaction with quartz.

All alloys were water quenched after heat treatment and well prepared for metallographic analysis. The equilibrium compositions of phases in the specimens were determined by electron probe microanalysis (EPMA) with 20 kV accelerating voltage and 1.0×10^{-8} A probe current. Additionally, the equilibrium compositions of liquid phases in some alloys annealed at 1300 °C were measured by energy dispersive spectroscopy (DSC) with 20 kV accelerating voltage and 2.0×10^{-9} A probe current. The crystal structure was identified by a Phillips Panlytical X-pert diffractometer using Cu-K α radiation with 40 kV voltage and 40 mA current. The results were measured in the range of 2 θ from 20° to 90° with a step interval of 0.015308° and a count time of 0.3 s per step. The melting points of some alloys were determined by differential scanning calorimeter (DSC) with a heating and cooling rate of 10 °C/min.

3. Results and Discussion

3.1. Microstructure

The phase relationship of the Ni-Cr-Ta ternary system at 1200 °C was established from 33 alloys annealed for 35 days. The nominal compositions of alloys and compositions of different phases at equilibrium are displayed in the Table 2. Meanwhile, the microstructure and XRD results of typical alloys annealed at 1200 °C for 35 days are presented in Figures 2 and 3, respectively.

Table 2.	Equilibrium	compositions of e	each phase in t	the Ni-Cr-Ta ter	rnary alloys an	nealed at 1200 °	°C for
35 days.							

	Phase Equilibrium	Composition (at. %)						
Nominal Alloys (at. %)		Phase 1		Phase 2		Phase 3		
1110ys (uu /s)	Phase 1/Phase 2/Phase 3	Cr	Та	Cr	Ta	Cr	Ta	
Ni54Cr43.5Ta2.5	fcc	43.4	53.9					
Ni _{80.5} Cr ₇ Ta _{12.5}	fcc/Ni ₃ Ta	8.5	10	3.3	19.9			
Ni ₅₉ Cr ₃₃ Ta ₈	fcc/Ni ₃ Ta	41.5	5.0	3.0	21.7			
Ni ₁₄ Cr ₆₅ Ta ₂₁	bcc-(Cr)/(Ni, Cr) ₂ Ta(HT)	97.0	0.4	50.8	30.0			
Ni ₄₆ Cr ₃₃ Ta ₂₁	bcc-(Cr)/(Ni, Cr) ₂ Ta(HT)/Ni ₃ Ta	81.5	0.9	33	23.4	3.4	22.4	
Ni ₆₅ Cr ₅ Ta ₃₀	Ni ₃ Ta/Ni ₂ Ta/(Ni, Cr) ₂ Ta(HT)	0.6	25.1	1.6	37.3	26.5	31.9	
Ni ₄₀ Cr ₂₂ Ta ₃₈	Ni ₆ Ta ₇ /(Ni, Cr) ₂ Ta(HT)	10.8	45	27.8	34.8			
Ni ₃₀ Cr ₃₁ Ta ₃₉	Ni ₆ Ta ₇ /(Ni, Cr) ₂ Ta(HT)	20.7	45.2	36.3	35.8			
Ni ₂₁ Cr ₃₃ Ta ₄₆	Ni ₆ Ta ₇ /(Ni, Cr) ₂ Ta(HT)	28.4	48	44.8	38.3			
Ni ₃₉ Cr ₄ Ta ₅₇	Ni ₆ Ta ₇ /NiTa ₂	6.5	52.7	0.2	64.5			
Ni ₁₃ Cr ₇ Ta ₈₀	NiTa ₂ /bcc-(Ta)	12	62.2	4.0	95.2			
Ni ₃₃ Cr ₃₄ Ta ₃₃	$(Ni, Cr)_2 Ta(HT)$	34.5	33.0					
Ni ₇₈ Cr ₉ Ta ₁₃	fcc/Ni ₃ Ta	11.4	8.3	1.9	21.9			
Ni53Cr37Ta10	fcc/Ni ₃ Ta/bcc-(Cr)	41.9	4.6	3.6	22.5	80.3	0.6	
Ni ₃₈ Cr ₃₂ Ta ₃₀	$(Ni, Cr)_2 Ta(HT)$	31.9	29.5					
Ni ₂₅ Cr ₃₇ Ta ₃₈	$Ni_6Ta_7/(Ni, Cr)_2Ta(HT)$	15.5	48.5	38.6	35.6			
Ni ₆₄ Cr ₇ Ta ₂₉	Ni ₃ Ta/Ni ₂ Ta/(Ni, Cr) ₂ Ta(HT)	0.5	24.9	1.8	37.1	26.6	32.1	
Ni53Cr8Ta39	Ni ₆ Ta ₇ /Ni ₂ Ta/(Ni, Cr) ₂ Ta(HT)	9.6	46.1	0.8	33.5	26.8	33.2	
Ni ₂ Cr ₇₅ Ta ₂₃	bcc-(Cr)/(Ni, Cr) ₂ Ta(HT)	98.6	0.7	66.7	30.8			
Ni ₉ Cr ₄₈ Ta ₄₃	Ni ₆ Ta ₇ /(Ni, Cr) ₂ Ta(HT)	40.1	50	54.1	39.6			
Ni ₂₇ Cr ₄₁ Ta ₃₂	$(Ni, Cr)_2 Ta(HT)$	40.9	32.1					
Ni ₂₆ Cr ₅₇ Ta ₁₇	Ni ₆ Ta ₇ /bcc-(Ta)	19.1	53.8	6.2	93.7			
Ni ₃₅ Cr ₅₃ Ta ₁₂	bcc-(Cr)/(Ni, Cr) ₂ Ta(HT)/Ni ₃ Ta	81.3	0.9	33.5	23.7	3.2	22.5	
Ni ₂₇ Cr ₅₃ Ta ₂₀	bcc-(Cr)/(Ni, Cr) ₂ Ta(HT)	92.8	0.2	36.9	25.9			
Ni ₂₃ Cr ₅₃ Ta ₂₄	bcc-(Cr)/(Ni, Cr) ₂ Ta(HT)	96.3	0.5	45.3	28.9			
Ni ₂₂ Cr ₄₅ Ta ₃₃	$(Ni, Cr)_2 Ta(HT)$	45.2	32.9					
Ni ₁₃ Cr ₅₃ Ta ₃₄	$(Ni, Cr)_2 Ta(HT)$	53.1	34					
Ni ₁₆ Cr ₃₈ Ta ₄₆	Ni ₆ Ta ₇ /(Ni, Cr) ₂ Ta(HT)	33.3	49.8	51	38.9			
Ni ₁₂ Cr ₃₃ Ta ₅₅	Ni ₆ Ta ₇ /bcc-(Ta)	35.6	52.1	6.8	93.1			
Ni ₁₅ Cr ₁₉ Ta ₆₆	Ni ₆ Ta ₇ /bcc-(Ta)	24.4	55.1	6.4	93.5			
Ni ₃₆ Cr ₅₈ Ta ₆	fcc/Ni ₃ Ta/bcc-(Cr)	41.8	4.5	3.4	22.3	80.1	0.5	
Ni ₅ Cr ₄₀ Ta ₅₅	Ni ₆ Ta ₇ /(Ni, Cr) ₂ Ta(HT)/bcc-(Ta)	39.2	51.5	58.7	38.5	6.3	93.5	
$Ni_{10}Cr_{25}Ta_{65}$	Ni ₆ Ta ₇ /bcc-(Ta)	30.5	52.4	5.4	93.2			

(a)

(c

Ni₂Ta





Figure 2. The microstructure of typical alloys in the Ni-Cr-Ta system annealed at 1200 °C for 35 days. (a) Ni₅₃Cr₃₇Ta₁₀; (b) Ni₆₄Cr₇Ta₂₉; (c) Ni₅₃Cr₈Ta₃₉; (d) Ni₅₉Cr₃₃Ta₈; (e) Ni₂₇Cr₅₃Ta₂₀; (f) Ni₁₂Cr₃₃Ta₅₅.



Figure 3. The XRD patterns of typical alloys in the Ni-Cr-Ta system annealed at 1200 °C for 35 days. (a) Ni₆₄Cr₇Ta₂₉; (b) Ni₅₃Cr₈Ta₃₉; (c) Ni₁₂Cr₃₃Ta₅₅; (d) Ni₁₃Cr₅₃Ta₃₄.

As presented in Figure 2a-c, the microstructure of three-phase regions was detected in these alloys. Figure 2a showed the three-phase equilibrium, two terminal solid solutions fcc, bcc-(Cr) and a compound Ni₃Ta, bright regions in the microstructure of the Ni₅₃Cr₃₇Ta₁₀ alloy. Figure 2b showed the three-phase equilibrium of the Ni₆₄Cr₇Ta₂₉ alloy, in which the white precipitated (Ni, Cr)₂Ta(HT) phase was uniformly distributed in the Ni₃Ta and Ni₂Ta phase. Moreover, the three-phase equilibrium state was identified by the XRD result in Figure 3a. Three-phase region, Ni₂Ta, Ni₆Ta₇ and (Ni, Cr)₂Ta(HT) was found in the Ni₅₃Cr₈Ta₃₉ alloy after annealed at 1200 °C for 35 days. Additionally, the XRD analysis in Figure 3b just confirmed the microstructure. As can be seen from Figure 2d–f, three two-phase regions were identified in these three alloys. Figure 2d showed the equilibrium of the gray matrix fcc and white Ni₃Ta phase in the Ni₅₉Cr₃₃Ta₈ alloy annealed at 1200 °C for 35 days. The phase relation of the Ni₂₇Cr₅₃Ta₂₀ alloy, a terminal solid solution bcc-(Cr) and a compound (Ni, Cr)₂Ta(HT), was described in Figure 2e. Furthermore, there was a two-phase section of white bcc-(Ta) and gray Ni₆Ta₇ phase in the Ni₁₂Cr₃₃Ta₅₅ alloy as illustrated in Figure 2f. Additionally, the crystal structure of the Ni₁₂Cr₃₃Ta₅₅ alloy was identified by the XRD result displayed in Figure 3c. Figure 3d showed the XRD result of a single (Ni, Cr)₂Ta(HT) phase in the Ni₁₃Cr₅₃Ta₃₄ alloy and the microstructure was displayed in the Figure 3e.

In the experiment, several alloys were designed to investigate phase relation of the Ni-Cr-Ta system at 1300 °C. Table 3 listed the alloys compositions and phase equilibrium compositions of the alloys annealed at 1300 °C. In addition, the microstructure and XRD patterns of typical alloys were presented in the Figures 4 and 5, respectively. Figure 4a,c showed two three-phase equilibriums with liquid phase. As presented in Figure 4a, the microstructure of three-phase region, gray liquid phase, black fcc phase and white Ni₃Ta phase, was determined in the Ni₅₉Cr₃₃Ta₈ alloy annealed at 1300 °C for 3 h. Figure 4c displayed the three-phase microstructure of liquid phase, oval-shaped bcc-(Cr) and (Ni, Cr)₂Ta(HT) phase in the Ni₃₅Cr₅₃Ta₁₂ alloy annealed at 1300 °C for 3 h. There was a three-phase section, black matrix Ni₃Ta, white precipitated Ni₂Ta and gray (Ni, Cr)₂Ta(HT) identified in the Ni₆₅Cr₅Ta₃₀ alloy as shown in Figure 4b. Figure 4d, e illustrated two two-phase equilibrium. The Ni_6Ta_7 and (Ni, Cr)₂Ta(HT) phase were determined in the Ni₁₆Cr₃₈Ta₄₆ alloy described in Figure 4d, and the two-phase equilibrium was supported by the XRD result in Figure 5a. The microstructure of the $Ni_{12}Cr_{33}Ta_{55}$ alloy in Figure 4e was confirmed as bcc-(Ta) and Ni_6Ta_7 by the XRD pattern presented in Figure 5b. Figure 4f showed the three-phase region, bcc-(Ta), Ni₆Ta₇ and (Ni, Cr)₂Ta(HT), of the Ni₅Cr₄₀Ta₅₅ alloy, and the result was confirmed by the XRD pattern in Figure 5c. As shown in Figure 4d–e, the cracks and holes were observed in the brittle Laves phase Ni₆Ta₇.

Nominal	Phase Equilibrium	Composition (at. %)					
Alloys (at%)	Phase 1/Phase 2/Phase 3	Phase 1		Phase 2		Phase 3	
	1 huse 1/1 huse 2/1 huse 5	Cr	Та	Cr	Ta	Cr	Ta
Ni ₅₄ Cr _{43.5} Ta _{2.5}	fcc	46.0	2.4				
Ni59Cr33Ta8 *	fcc/Ni ₃ Ta/L	34.3	6.6	3.7	22.4	37.5	11.7
Ni ₁₄ Cr ₆₅ Ta ₂₁	bcc-(Cr)/(Ni, Cr) ₂ Ta(HT)	95.5	0.6	51.6	29.6		
Ni ₄₆ Cr ₃₃ Ta ₂₁ *	L/(Ni, Cr) ₂ Ta(HT)/Ni ₃ Ta	35.1	15.9	33.0	24.6	3.3	23.8
Ni ₆₅ Cr ₅ Ta ₃₀	Ni ₃ Ta/Ni ₂ Ta/(Ni, Cr) ₂ Ta(HT)	0.6	23.6	1.0	31.3	22.4	31.9
Ni ₄₀ Cr ₂₂ Ta ₃₈	Ni ₆ Ta ₇ /(Ni, Cr) ₂ Ta(HT)	11.6	45.5	22.9	35.2		
Ni ₃₀ Cr ₃₁ Ta ₃₉	Ni ₆ Ta ₇ /(Ni, Cr) ₂ Ta(HT)	18.9	46.7	36.5	35.6		
Ni ₂₁ Cr ₃₃ Ta ₄₆	Ni ₆ Ta ₇ /(Ni, Cr) ₂ Ta(HT)	27.7	48.3	45.1	38.2		
Ni ₃₉ Cr ₄ Ta ₅₇	Ni ₆ Ta ₇ /NiTa ₂	0.5	65.5	7.2	55		
Ni ₁₃ Cr ₇ Ta ₈₀	NiTa ₂ /bcc-(Ta)	12.0	58.2	4.7	94.5		
Ni33Cr34Ta33	$(Ni, Cr)_2 Ta(HT)$	34.5	33.1				
Ni ₇₈ Cr ₉ Ta ₁₃	fcc	9.2	12.3				
Ni53Cr37Ta10 *	fcc/L	38.4	6.5	36.9	13.5		
Ni ₃₈ Cr ₃₂ Ta ₃₀	$(Ni, Cr)_2 Ta(HT)$	32.2	30.2				
Ni ₂₅ Cr ₃₇ Ta ₃₈	$Ni_6Ta_7/(Ni, Cr)_2Ta(HT)$	19.6	47.6	39.3	36.6		
Ni ₆₄ Cr ₇ Ta ₂₉	Ni ₃ Ta/Ni ₂ Ta/(Ni, Cr) ₂ Ta(HT)	0.7	23.4	1.2	31.1	22.3	31.6
Ni53Cr8Ta39	Ni ₆ Ta ₇ /Ni ₂ Ta/(Ni, Cr) ₂ Ta(HT)	8.5	45.4	0.2	32.7	17.4	36.5
Ni ₂ Cr ₇₅ Ta ₂₃	bcc-(Cr)/(Ni, Cr) ₂ Ta(HT)	98.1	0.8	65.2	32.3		
Ni ₉ Cr ₄₈ Ta ₄₃	$Ni_6Ta_7/(Ni, Cr)_2Ta(HT)$	42.3	50.2	54.3	40.6		
Ni ₂₇ Cr ₄₁ Ta ₃₂	$(Ni, Cr)_2 Ta(HT)$	41.0	31.8				
Ni ₂₆ Cr ₅₇ Ta ₁₇	Ni ₆ Ta ₇ /bcc-(Ta)	19.8	54.0	5.1	94.3		
Ni35Cr53Ta12 *	bcc-(Cr)/(Ni, Cr) ₂ Ta(HT)/L	82.8	0.9	37.5	24.4	39.8	12.4
Ni ₂₇ Cr ₅₃ Ta ₂₀	bcc-(Cr)/(Ni, Cr) ₂ Ta(HT)	89.2	0.5	43.6	27.0		
Ni ₂₃ Cr ₅₃ Ta ₂₄	bcc-(Cr)/(Ni, Cr) ₂ Ta(HT)	93.5	0.5	45.5	26.8		
Ni ₂₂ Cr ₄₅ Ta ₃₃	$(Ni, Cr)_2 Ta(HT)$	45.3	32.5				
Ni ₁₃ Cr ₅₃ Ta ₃₄	$(Ni, Cr)_2 Ta(HT)$	53.5	33.7				
Ni ₁₆ Cr ₃₈ Ta ₄₆	Ni ₆ Ta ₇ /(Ni, Cr) ₂ Ta(HT)	35.7	48.8	49.8	39.4		
Ni ₅₁ Cr ₂₈ Ta ₂₁ *	Ni ₃ Ta/(Ni, Cr) ₂ Ta(HT)/L	3.1	23.4	33.0	24.5	34.8	16.2
Ni ₁₂ Cr ₃₃ Ta ₅₅	Ni ₆ Ta ₇ /bcc-(Ta)	36.9	51.0	5.4	94.1		
$Ni_{15}Cr_{19}Ta_{66}$	Ni ₆ Ta ₇ /bcc-(Ta)	26.8	53.3	5.3	93.8		
Ni ₃₆ Cr ₅₈ Ta ₆ *	fcc/bcc-(Cr)/L	40.1	6.1	80.7	0.5	39.7	11.8
Ni ₇₈ Cr ₁₂ Ta ₁₀	fcc/Ni ₃ Ta	13.4	9.7	3.2	21.5		
Ni ₅ Cr ₄₀ Ta ₅₅	Ni ₆ Ta ₇ /(Ni, Cr) ₂ Ta(HT)/(Ta)	41.6	51.1	53.5	42.1	6.3	93.5
Ni ₁₀ Cr ₂₅ Ta ₆₅	Ni ₆ Ta ₇ /bcc-(Ta)	29.8	54.6	6.8	93.1		

* Indicated that the alloy was annealed at 1300 $^\circ C$ for 3 h.



Figure 4. The microstructure of typical alloys in the Ni-Cr-Ta system annealed at $1300 \degree C$. (a) $Ni_{59}Cr_{33}Ta_8$; (c) $Ni_{35}Cr_{53}Ta_{12}$ alloys annealed at $1300 \degree C$ for 3 h. (b) $Ni_{65}Cr_5Ta_{30}$; (d) $Ni_{16}Cr_{38}Ta_{46}$; (e) $Ni_{12}Cr_{33}Ta_{55}$; (f) $Ni_5Cr_{40}Ta_{55}$ alloys annealed at $1300 \degree C$ for 15 days.



Figure 5. The XRD patterns of typical alloys in the Ni-Cr-Ta system annealed at 1300 °C for 15 days. (a) Ni₁₆Cr₃₈Ta₄₆; (b) Ni₁₂Cr₃₃Ta₅₅; (c) Ni₅Cr₄₀Ta₅₅.

The measured equilibrium compositions of the Ni-Cr-Ta system at 1200 °C and 1300 °C were listed in the Tables 2 and 3, respectively. According to the experimental results, the isothermal sections of the Ni-Cr-Ta ternary system at 1200 °C and 1300 °C were established in Figures 6 and 7. At 1200 °C, there are eight stable solid phases, fcc, Ni₃Ta, Ni₂Ta, Ni₆Ta₇, NiTa₂, bcc-(Ta), bcc-(Cr), and (Cr)₂Ta(HT). Meanwhile, five three-phase regions, fcc + Ni₃Ta + bcc-(Cr), Ni₃Ta + bcc-(Cr) + (Ni, Cr)₂Ta(HT), Ni₃Ta + Ni₂Ta + (Ni, Cr)₂Ta(HT), Ni₂Ta + Ni₆Ta₇ + (Ni, Cr)₂Ta(HT) and Ni₆Ta₇ + bcc-(Ta) + (Ni, Cr)₂Ta(HT), were experimentally determined and marked as triangle with solid lines in present work. However, the other three three-phase equilibria, $Ni_6Ta_7 + NiTa_2 + bcc-(Ta)$, $bcc-(Cr) + (Cr)_2Ta(LT) + (Ni, Cr)_2Ta(HT)$ and bcc-(Ta) + $(Cr)_2Ta(LT)$ + $(Ni, Cr)_2Ta(HT)$, at 1200 °C were inferred and presented as triangle with dash lines. A single-phase region (Ni, Cr)₂Ta(HT) with hexagonal MgZn₂-type (C14) structure was detected with a large composition range of about 25.8-66.0 at. % Cr, 2.5-44.3 at. % Ni, and 24.0-40.0 at. % Ta, at 1200 °C. It should be noted that the (Ni, Cr)₂Ta(HT) appears in two isothermal sections of the Ni-Cr-Ta system in spite of the non-existence in Cr-Ta subsystem at two temperatures. Distinctly, the Ni addition to the Cr-Ta binary system stablizes the (Ni, Cr)₂Ta(HT) phase at low temperature. The Ni₃Ta and Ni₂Ta just dissolves a little of Cr, while the solubility of Cr in the Ni₆Ta₇ and NiTa₂ phase reaches up to 40.1 and 13.1 at. %, respectively. Despite the same crystal structure of Ta and Cr, the solubility of Ta/Cr in two bcc terminal solid solution is quite small.



Figure 6. Experimentally determined isothermal section of the Ni-Cr-Ta system at 1200 °C.



Figure 7. Experimentally determined isothermal section of the Ni-Cr-Ta system at 1300 °C.

The isothermal section at 1300 °C is shown in Figure 7, where with existence of liquid phase, two three-phase regions fcc + Ni₃Ta + bcc-(Cr) and Ni₃Ta + bcc-(Cr) + (Ni, Cr)₂Ta(HT) disappear and are replaced by four confirmed three-phase equilibria, fcc + bcc-(Cr) + L, fcc + Ni₃Ta + L, bcc-(Cr) + (Ni, Cr)₂Ta(HT) + L and Ni₃Ta + (Ni, Cr)₂Ta(HT) + L. The other three-phase regions are similar to the isothermal sections at 1200°C, where the three-phase equilibria, Ni₃Ta + Ni₂Ta + Ni₂Ta + (Ni, Cr)₂Ta(HT), Ni₂Ta + Ni₆Ta₇ + (Ni, Cr)₂Ta(HT) and Ni₆Ta₇ + bcc-(Ta) + (Ni, Cr)₂Ta(HT), were determined and the other three were inferred. With the temperature rising to 1300 °C, a liquid phase was identified near the Ni-Cr side alloys. It is evident that the Ta addition to the Ni-Cr alloys decreases the melting point of the Ni-Cr eutectic alloys in consideration of the eutectic reaction L \rightarrow Ni₄₄ + Cr₅₆ at 1345 °C.

3.3. The Liquid Region

As can be observed from the microstructure in Figure 4c and isothermal section at 1300 °C, a liquid phase was confirmed at 1300 °C. However, according to the experimental results in the three subsystems, no liquid phase exists at 1300 °C. In order to confirm the experimental results in the present work, the DSC analysis was conducted to obtain the melting point of the related alloys. On the basis of DSC result and microstructure in Figure 8, the bcc-(Cr) phase transformed to liquid phase as the temperature increased from 1200 °C to 1285 °C and the melting point of the bcc-(Cr) phase in the Ni₅₁Cr₂₈Ta₂₁ alloy was measured to be about 1237 °C. Owing to the liquid phase that appeared near the Ni-Cr side, we supposed that the Ta addition in Ni-Cr alloys decreases the temperature of the eutectic reaction, $L \rightarrow fcc + bcc-(Cr)$. The corresponding results indicate that the Ta addition reduces the melting point of the Ni-Cr alloys.

16

12

8

4

0

1180

Endothermic mW/mg



1260

1

1

1280 1290

Figure 8. The heating curve of the Ni₅₁Cr₂₈Ta₂₁ alloy and microstructure annealed at (**a**) 1200 °C for 35 days; (**b**) 1285 °C for 3 h.

Temperature / °C

1240

T_m(1237°C)

1220

Heating curve

1200

4. Conclusions

In the present work, the isothermal sections of the Ni-Cr-Ta ternary system at 1200 °C and 1300 °C were experimentally established. The corresponding results are shown as follows:

- (1) The solubility of Cr in Ni₆Ta₇ phase was about 41.6 at. % at 1300 °C, and no ternary compound was found at two sections.
- (2) The high temperature (Ni, Cr)₂Ta(HT) (MgZn₂-type) phase with a large composition range was determined at both two temperatures, which was stabilized by the Ni addition to Cr–Ta alloys against low temperature, and its solubility increased as temperature raise from 1200 °C to 1300 °C.
- (3) A small liquid region was confirmed at 1300 °C, while it disappeared at 1200 °C. The results indicate that the addition of Ta reduced the melting point of the Ni–Cr alloys.

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