

Spectral properties and thermal stability of solar selective absorbing AlNi–Al₂O₃ cermet coating

Yafei Xue, Cong Wang*, Wenwen Wang, Yu Liu, Yongxin Wu, Yuping Ning, Ying Sun

Center for Condensed Matter and Material Physics, Department of Physics, Beihang University, Beijing 100191, PR China

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Abstract

A new solar selective absorbing coating of AlNi–Al₂O₃ cermet multilayer is designed and prepared on Cu or stainless steel (SS) substrate by magnetron sputtering. The coating consists of an IR-reflective metallic layer, an absorption bilayer and an anti-reflection layer from substrate to top. The IR-reflective metallic layer is Cu or Mo. The absorption bilayer consists of a high metal volume fraction (HMF) AlNi–Al₂O₃ cermet sublayer and a low metal volume fraction (LMVF) AlNi–Al₂O₃ cermet sublayer. The anti-reflection layer is Al₂O₃. The optical performance (α/ε) of the coating is 0.94/0.07 on Cu substrate and 0.95/0.078 on SS substrate by optimizing the metal volume fraction and thickness of each layer. The samples deposited on SS substrate were heated at 500 °C for 138 h and 284 h in vacuum to evaluate the optical stability. Experimental results showed that the high temperature stable AlNi intermetallic compound, instead of Mo in the traditional Mo–Al₂O₃ coatings, displayed a good thermal stability at 500 °C. There is no significant change appearing in the optical properties of the coating after the heat treatment. It means that the new coating is a good potential candidate as a high temperature solar selective absorbing coating for parabolic trough concentrated solar power (CSP).

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1. Introduction

Solar spectral selective absorbing coating is a critical component of a solar photo-thermal conversion absorber, which requires a high absorptance in the wavelength 300–2500 nm and a low thermal emissivity at the operating temperature. This is realized by low reflectance (near zero) of an absorber surface in the main solar radiation spectrum and high reflectance (close to one) in the IR region (>2500 nm) (Zhang et al., 1996).

In the last several decades, magnetron sputtering technology has been developed to deposit solar selective coatings for high temperature usage, such as Pt–Al₂O₃ (Nuru et al., 2012) and Cr–Al₂O₃ (Yin et al., 2009) cermet coat-

ings. By computer modeling calculations, Zhang and Mills (1992) predicted a double cermet layer structure. The typical coating structure from surface to substrate consists of an anti-reflection layer composed of a transparent ceramic material that enhances solar transmission, a low metal volume fraction (LMVF) cermet solar absorption layer, a high metal volume fraction (HMF) cermet solar absorption layer, which forms interference absorption bilayer, and an IR-reflective metallic layer that decreases the IR emission. Many experiments have proved that the coating structure has better solar spectral selective performance. Based on above coating structure, a large number of solar selective coating series, such as Mo–SiO₂ (Wang et al., 2011), M–AlN (Zhang, 1998), Ni–NiO (Adsten et al., 2000), W–Al₂O₃ (Antonaia et al., 2010), Ag–Al₂O₃ (Barshilia et al., 2011), Mo–Al₂O₃ (Lanxner and Elgat, 1990; Xinkang et al., 2008) HfMoN/HfON/Al₂O₃ (Selvakumar et al.,

* Corresponding author. Tel./fax: +86 1082338346.

E-mail address: congwang@buaa.edu.cn (C. Wang).

2012) and TiAlN/TiAlON/Si₃N₄ (Biswas et al., 2008; Barshilia et al., 2008; Godinho et al., 2010) have been studied as an absorber surface.

However, parabolic trough concentrated solar power (CSP) requires such kinds of solar selective absorbing coatings working at high temperature (>400 °C). Hence, thermal stability of a coating is becoming a critical problem, which determines the working lifetime of a solar absorber. In the cermet layer of a solar selective coating, single metal particles are usually dispersed in the ceramic matrix layer. However, single metal particles are easy to diffuse, congregate or to be oxidized. This causes deterioration in optical properties of the coating at high temperature. For example, Mo is easy to diffuse in Al₂O₃ ceramic matrix and interfacial boundary at high temperature above 500 °C (Xinkang et al., 2008). If we choose the high temperature stable binary alloy, such as AlNi to disperse in the Al₂O₃ matrix, instead of Mo in Mo–Al₂O₃ coating, maybe it could restrain the diffusion and oxidation, and improve the thermal stability of the absorbing coating at high temperature. We know that the AlNi alloy owns good thermal stability and better anti-diffusion and anti-oxidant effects than single Mo or Ni metal particles (Mahesh et al., 2009). In this paper, we choose AlNi binary alloy to substitute Mo single metal to prepare a new high temperature stabilized AlNi–Al₂O₃ cermet selective absorbing coating. To our knowledge, NiAl–Al₂O₃ solar selective absorbing coating has never been mentioned for parabolic trough CSP applications up to now.

Based on the double cermet layer structure, we prepared high metal volume fraction AlNi–Al₂O₃ layer (HMF), low metal volume fraction AlNi–Al₂O₃ layer (LMF) and Al₂O₃ anti-reflection layer from bottom to top on Cu substrate as solar absorbing coating by DC and RF magnetron sputtering. Here Cu works as an IR-reflective metallic layer which owns good infrared reflectance. We also deposited the coating on SS substrate and a Mo layer was deposited between SS and HMF as IR-reflective metallic layer. The optical properties and thermal stability of the coating were investigated.

2. Experimental

Metallic Mo, alloy AlNi and ceramic Al₂O₃ in the coating were deposited using DC and RF magnetron sputtering, respectively. Mo (99.99% purity) metallic target, Al₂O₃ (99.99% purity) ceramic target and AlNi (99.99% purity) alloy target are mechanically clamped to three planar sputter sources which are mounted horizontally on the

base of the vacuum chamber. All of the Mo, Al₂O₃ and AlNi targets have a diameter of 60 mm and thickness of 4 mm. The vacuum chamber was pumped to a base pressure of 4×10^{-3} Pa, and then putting Ar gas into the chamber, the sputtering pressure was changed by adjusting the valve of the molecular pump. The parameters for the deposition of Mo, AlNi and Al₂O₃ are listed in Table 1. Prior to the deposition of the multilayer films, Cu or SS substrate (25 mm × 25 mm × 1 mm) was rinsed in acetone and ethanol baths for 15 min. For the layers of HMF and LMF, the expected change of the metal volume fraction is adjusted by the sputtering time of AlNi and Al₂O₃ when the substrate rotates facing the two targets alternately. There are many AlNi minilayers (2–6 nm) and Al₂O₃ minilayers (3–8 nm) alternately deposited, forming the AlNi–Al₂O₃ absorption layer, i.e. HMF or LMF sublayer. For example, when we deposit AlNi minilayer with the thickness of 6 nm and Al₂O₃ minilayer with 4 nm for six times alternately, then the metal volume fraction and thickness of the absorption layer are 60% and 60 nm, respectively.

The deposition rates of the Mo, AlNi alloy and ceramic Al₂O₃ were derived by measuring the thickness of the films using a Tencer P-10 α -step surface profiler. The optical reflectance was measured by UVPC3100 spectrophotometer in the wavelength range of 0.2–2.6 μ m and Shimadzu IR-470 Infrared Spectrophotometer in the range of 2.6–25 μ m. The nearly normal emissivity was measured by NEM-80 normal emissivity tester at 80 °C. The structure of the films was investigated by X-ray diffraction (XRD) on a Dmax diffractometer with Cu K α (40 kV, 20 mA, $\lambda = 0.15406$ nm) radiation with the 2θ angles in the range 20–90° in the scanning step of 0.02°. Surface morphology was observed by a JEM-100CX scanning electron microscopy (SEM). The evaluation of thermal stability was carried out by putting the coating sample at 500 °C for 12 h, 138 h and 284 h, respectively, in a vacuum quartz tube with a pressure of 5×10^{-2} Pa.

3. Results and discussion

3.1. Optimization and characterization of AlNi single layer

Figs. 1 and 2 show XRD patterns of the single AlNi layer prepared under different sputtering powers on Cu and SS substrates, respectively. When the target power density was below 2.93 W/cm², only the diffraction peaks from Cu or SS substrate were observed. It is only when the target power density was increased above 2.93

Table 1
The parameters for the deposition of Mo, AlNi and Al₂O₃ layers.

Phase	Sputtering method	Ar gas (sccm)	Total pressure (Pa)	Sputtering distance (mm)	Voltage (V)	Current (A)	Power density (W/cm ²)
Mo	DC	50	0.75	60	240	0.7	5.94
AlNi	DC	50	1.0	60	460	0.18	2.93
Al ₂ O ₃	RF	50	1.0	60	1100	0.3	5.66

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