Corrosion Properties of Ag-Au-Cu-Pd System Alloys Containing Indium

Masayuki Hattori, Teruhiko Tokizaki, Michihiko Matsumoto and Yutaka Oda

Department of Dental Materials Science, Tokyo Dental College, 1-2-2 Masago, Mihama-ku, Chiba 261-8502, Japan

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Abstract

In this study, the corrosion resistance of Ag-Au-Cu-Pd system alloys consisting of 5 or 10 mass% indium was evaluated. Levels of element release and tarnish were determined and electrochemical measurements performed. Results were compared with those for commercial silver-palladium-gold alloy. In terms of electrochemical behavior, the trans-passive potential of these experimental alloys was 168–248mV. Experimental alloys with 25 mass% Au showed similar corrosion resistance to control gold-silver-palladium alloy. Amount of released elements was 14–130 μg/cm² at 7 days, which is in the allowable range for dental alloys. Addition of indium to Ag-Au-Cu-10mass%Pd system alloys was effective in increasing resistance to tarnish and alloys containing 10 mass% of indium showed a minimal decrease in L* values after immersion. These findings indicate that 25Au-37.5Ag-15Cu-10Pd-2Zn-10In-0.5Ir alloy is applicable in dental practice.

Key words: Ag-Au-Cu-Pd system alloy—Corrosion resistance—Tarnish

Introduction

High carat gold alloys with superior corrosion resistance in an oral environment have been widely used as dental alloys, presenting adequate workability and mechanical properties. On the other hand, the gold-silver-palladium alloys covered by the national and social health insurance systems in Japan have also been used over a long period of time. However, palladium allergies have been reported for alloys with a high palladium content[11]. Therefore, the development of alloys with a low palladium content is necessary. Alloys with silver as their main component are considered to have low sulfidation resistance, showing inferior corrosion resistance as well as a tendency to tarnish under oral conditions. Addition of palladium to alloys is quite effective in inhibiting tarnish and corrosion[8]. Therefore, the Japanese Industrial Standard (JIS)[3] has required 20% or more of palladium content.
in gold-silver-palladium alloys. To prevent tarnish and corrosion in alloys containing silver as their main component while reducing the amount of palladium content, an increase in gold or platinum content has been suggested by some reports.\textsuperscript{5,13} Nevertheless, 30 mass% or more of gold content is required in alloys with low palladium content to inhibit tarnish and corrosion. In this situation, increasing the amount of gold or platinum is not a suitable option if one takes into account the effects of price fluctuations in the metal markets.

On the other hand, addition of cadmium or indium to silver alloy was effective in preventing sulfidation by silver.\textsuperscript{2} One study reported that indium was less allergenic than palladium.\textsuperscript{12} Accordingly, in the present study, 6 types of Ag-Au-Cu-Pd system alloy (Au: 20 or 25 mass%, Pd: 10 mass%) containing 5 or 10 mass% indium were produced for evaluation of corrosion and tarnish resistance. Potentiodynamic polarization behavior, tarnish and dissolution of metal elements was determined by immersion tests using sulfide solutions to evaluate the development of a new type of Ag-Au-Cu-Pd system alloy as well as clarify the effect of addition of indium to this alloy system.

### Materials and Methods

#### 1. Alloy preparation

As experimental alloys, 6 types of alloy (100 g each) consisting of 10 mass% palladium, 15 mass% copper, 20 or 25 mass% gold, 5 or 10 mass% indium and 37.5–52.5 mass% silver were produced (Table 1). For fabrication of each alloy, commercially available 99.9% Cu, In, Ag and Zn (Ishifuku Metal Industry, Tokyo, Japan) and 99.99% Au (Ishifuku Metal Industry) were used. As a control, a gold-silver-palladium alloy containing 12% gold (Kinpara S12 (S12): 51Ag-20Pd-14.5Cu-12.0Au and others, mass%, Ishifuku Metal Industry) was tested.

#### 2. Corrosion and tarnish test

1) Electrochemical measurement

Plate type specimens (14 x 14 x 1 mm) were cast with 6 types of experimental alloy and one control alloy. Each specimen was polished using 600 grit waterproof abrasive paper and then ultrasonically cleaned in acetone and distilled water. As a testing solution, 0.9% aqueous NaCl solution (pH = 6) was used. Using the Potentiostat Model 273A & Corrosion software M352C (PARC, Oak Ridge, Tennessee, USA) and an electrolytic cell stored in a thermostatic chamber at 37°C, saturated calomel electrode (SCE) was used as a reference electrode. In addition, a platinum counter electrode was prepared. Each specimen was immersed in an electrolytic cell using a sample holder with 1 cm\(^2\) of the surface area of the specimen exposed.

Potentiodynamic polarization was performed under the following conditions: potential range, between −1,000 mV and +300 mV; scanning rate, 0.33 mV/s. The
relationship between potential and current density was plotted. Transpassive potential, passive current density and corrosion potential at the potentiodynamic polarization curves were determined according to ISO10271 and JIS T 6002.  

2) Tarnish test  

After casting experimental and control alloys into plates (10 × 10 × 1 mm), the specimens were embedded in epoxy resin. Thereafter, mirror polishing was performed using the standard metallographic procedure followed by ultrasonic cleaning with acetone and distilled water. The testing was conducted according to ISO08891. A cyclic dipping device (Nihon Void, Tokyo, Japan) was used to repeat specimen soaking in 0.1 mol/liter sodium sulfide hydrate solution (pH ≈ 12.5; 10–15 sec) and air (45–50 sec) for 72 hr. Subsequently, differences in color were measured with a color meter (MCR-A, Minolta, Tokyo, Japan). Color difference (ΔE*ab) between at before and after cyclic immersion was determined using the following equation:

\[ \Delta E^{*}_{ab} = \left[ (\Delta L^*)^2 + (\Delta a^*)^2 + (\Delta b^*)^2 \right]^{1/2} \]

3) Static immersion test  

After casting experimental and control alloys into plates (10 × 10 × 1 mm), the surfaces of the specimens were polished using 600 grit waterproof abrasive paper. The specimens were ultrasonically cleaned in ethanol and distilled water. After hanging specimens by a nylon string in a sample bottle, they were stored in 4 ml aqueous solution (pH ≈ 2.3) comprising 0.1 mol/liter lactic acid and 0.1 mol/liter NaCl for 7 days at 37°C using a thermostatic chamber following the method of ISO10271. Elements dissolved into the testing solution after 7 days immersion were analyzed by inductively coupled plasma-atomic emission spectroscopy (Vista-MPX, SII, Chiba, Japan). In this test, seven elements consisting of Au, Ag, Cu, Pd, Zn, In and Ir were analyzed. The lower detection limit of each element was 0.07 µg. The released quantity of each element was determined by obtaining the average of 3 specimens in each group.  

3. Statistics  

For the statistical analysis of released elements and color differences, a one-way analysis of variance (ANOVA) and the Scheffé multiple comparison test were conducted.  

Results  

1. Corrosion resistance and tarnish  

The potentiodynamic polarization curves of the experimental alloys containing 25%
gold (#2500, #2505, #2510) are shown in Fig. 1. In terms of corrosion potential, which ranged between $-200$ and $-150$ mV, the #2500 experimental alloy showed the minimal value at $-200$ mV; while #2505 exhibited the highest value at $-150$ mV. All the alloys in this group were passivated at around $-50$ to $0$ mV. In terms of passive current densities at $100$ mV, which ranged from 5 to $30 \mu$A/cm$^2$, #2510 exhibited the highest current density. Thereafter, a rapid increase in current densities at potentials of above $200$ mV was recorded.

The potentiodynamic polarization curves of the experimental alloys containing 20% gold (#2000, #2005, #2010) are shown in Fig. 2. As with the experimental alloy containing 25% gold, corrosion potential ranged between $-200$ and $-150$ mV, with #2000 experimental alloy showing the minimal value at $-200$ mV, while #2010 showed the highest value at $-150$ mV. All the alloys in this group were passivated at around $-50$ to $0$ mV. In terms of passive current densities at $100$ mV, which ranged from 5 to $10 \mu$A/cm$^2$, #2010 showed the highest current density. A rapid increase in the current densities at potentials of between 150 and 200 mV was recorded.

The corrosion potential of S12 as a control at $-100$ mV was the highest value. It was passivated at a potential of around $-30$ mV; the passive current density at $100$ mV was approximately $3 \mu$A/cm$^2$. A rapid increase in the current density at around $250$ mV was observed.

The potentials with a rapid increase in current densities are presented as transpassive potentials in Fig. 3. The transpassive potentials of the experimental alloys ranged from 168 to 248 mV (vs. SCE). In this test, #2500 showed the highest value at $248 \pm 11$ mV, while #2010 showed the minimum value, $168 \pm 5$ mV. The transpassive potential of S12 as a control was $247 \pm 4$ mV.

The color values ($L^*$) of the experimental alloys at before and after immersion are
shown in Fig. 4. Before immersion, \( L^* \) values of all testing materials ranged at almost the same level (80–84). The value of S12 (80) as a control was slightly lower than that of the other alloys. After immersion, \( L^* \) values of all testing materials ranged between 50 and 72, with #2510 and #2000 showing the maximum (72) and minimum (50) values, respectively. S12 as a control presented a relatively higher value (68) than the experimental groups except #2510.

The color difference values (\( \Delta E^{ab} \)) after immersion of the experimental alloys are shown in Fig. 5. \( \Delta E^{ab} \) values ranged between 11 and 36. #2510 showed the highest value (36); #2005 showed the minimum value (11). The \( \Delta E^{ab} \) value of S12 as a control (12) was relatively lower.

The quantity of release for each element and the total released amount from the experimental alloys are shown in Table 2. Dissolution of Ag, Cu, Pd, Zn and In from each experimental alloy was found, but release of Au and Ir was below the lower detection limit (0.07 \( \mu g/cm^2/7 \) days). The minimum value of total dissolution (14.61 ± 10.15 \( \mu g/cm^2/7 \) days) was observed in #2500; the maximum value (130.65 ± 9.75 \( \mu g/cm^2/7 \) days) was found in #2010. The value of the control group (S12) was 4.47 ± 1.53 \( \mu g/cm^2/7 \) days.

### Discussion

1. **Alloy composition**

   Studies regarding indium as an additive element in dental alloys have been conducted over a number of years. Indium is a relatively
low allergenic element among additive elements for dental alloys\textsuperscript{10}. Matsumoto\textsuperscript{6} reported that addition of 20–30\% indium to silver alloys improved resistance to tarnish. In the present study, the effects of addition of 5 or 10 mass\% indium to Ag-Au-Cu-Pd system alloys were investigated in terms of resistance to corrosion and tarnish.

Yoshida et al.\textsuperscript{13} and Matsumoto et al.\textsuperscript{5} conducted studies on Ag-Au-Cu-Pd system alloys containing 5 mass\% palladium, and found that 30–40 mass\% gold was needed in addition to 5 mass\% palladium to obtain a sufficient amount of resistance to corrosion and tarnish. The palladium content of commercially available gold-silver-palladium alloys adopted for national insurance is regulated to 20 mass\%.

In the present study, to reduce the amount of gold as much as possible, these amounts were fixed at 20 and 25 mass\% and palladium content at 10 mass\%.

2. Corrosion and tarnish resistance

Although each experimental alloy presented similar potentiodynamic polarization behavior, the experimental alloys with 25 or 20 mass\% gold showed different levels of transpassive potential (Figs. 1, 2). A transpassive potential of more than 200 mV existed in the former, whereas less than 200 mV transpassive potential was found in the latter, indicating the influence of the gold content. On the other hand, a higher passive current density was found in the experimental alloys with 10\% indium (#2510, #2010) than the alloys with 5\% indium (#2505, #2005). This suggests that increase in indium content causes lowering of corrosion resistance. According to the comparison of potentiodynamic polarization curves against commercially available gold-silver-palladium alloy as a control, all the experimental alloys with 25 mass\% gold were smaller than in those with 20 mass\% gold, indicating the effect of gold content in alloys. In addition, the $\Delta E_{ab}$ values of the experimental alloys with 10 mass\% of indium were lower than those with 5 mass\% indium. As with the aforementioned mechanism, this result indicates the preventive effects of indium against sulfidation by silver. Only experimental #2510 showed the same level of color difference with the control alloy (commercially available gold-silver-palladium alloy).

Large amounts of Cu, In and Zn were released in the solutions by the specimens in this study. Although the zinc content in all the experimental alloys was relatively low (2 mass\%), released amounts of Zn were relatively high. This result agrees with those of previous reports showing that released amount of Zn was not in proportion to that of other alloy components\textsuperscript{14}; it is suggested that base metals
such as Cu, In and Zn on the specimen surfaces are selectively released in solutions. Released amounts of alloy elements from dental alloys should be kept as low as possible. The total released amount from dental alloys as determined by the ISO 22674 standard is 200µg/cm²/7 days or less. The total released amounts in all the experimental alloys showed higher values than the control gold-silver-palladium alloy; however, all the values were within the limited range of the ISO standard. Therefore, the released amounts in the experimental alloys in the present study are considered to be within the allowable range for dental alloys.

In conclusion, addition of indium to Ag-Au-Cu-10mass%Pd system alloys is effective in increasing tarnish resistance. These findings suggest that 25Au-37.5Ag-15Cu-10Pd-2Zn-10In-0.5Ir alloy is applicable in dental practice from the viewpoint of corrosion and tarnish resistance.

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References


Reprint requests to:
Dr. Masayuki Hattori
Department of Dental Materials Science,
Tokyo Dental College,
1-2-2 Masago, Mihama-ku,
Chiba 261-8502, Japan
E-mail: hattori@tdc.ac.jp