Effects of Alloying Elements on Creep Properties of 9Cr–3.3W–0.5Pd–V, Nb, N, B Steels

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Owing to the fine precipitations of FePd L1₀ type ordered phase $\alpha''$, Pd addition to 9Cr ferritic heat resistant steel improves the creep strength by not only reducing the creep rate but also extending the creep rupture life. In this study, effects of alloying elements on creep properties of Pd added 9Cr ferritic steels have been evaluated with 10 kg-ingots in order to search for substitutional element for the precious metal Pd. Among the metals from VIIA to IB except Fe and Tc, some elements were found to harden the steels aged at around 823 K, which is the precipitation temperature of $\alpha''$. Corresponding to the hardness change, some elements were found to improve the creep properties also. Among such elements, especially Re addition prevented the growth of Laves phase and remarkably reduced the minimum creep rate at lower stress level. From these results, pipe fabrication and welding tests were performed on Pd and Re-added 9Cr steel using 180 kg-ingots. It was found that this steel could easily be welded by gas tungsten arc welding and formed into pipe without harmful defects.

KEY WORDS: ferritic steels; FePd; L1₀ type ordered intermetallic phase; creep; microstructure.

1. Introduction

High Cr ferritic steels such as ASME P91 steel have been successfully used for large diameter and thick section boiler components such as main steam pipe and header in super critical boilers in fossil-fired power plants. Recent trend to utilization of clean energy leading to protection of global environment has been accelerating application of ultra super critical (USC) boilers, which are operated with higher efficiency in power generation than in conventional ones and thus release less amount of carbon dioxide. The USC boilers require new steels with improved creep strength and steam-oxidation resistance at elevated temperatures over 873 K, because of the increase in operating temperature and pressure of the steam. Addition of W to the steel has been found to be effective to increase creep rupture strength at high temperatures and has already been used in some newly developed steels such as P92 and P122 for the USC boilers.

A typical initial microstructure of these steels consist of tempered lath-martensite matrix with dislocation density of the order of $10^{14}/\text{m}^2$ and $\text{M}_2\text{C}_6$ (M: Cr, Fe, Mo, W etc.) type carbides along prior austenite grain boundary as well as packet, block and lath boundaries. MX (M: V, Nb etc. and X: C, N) type carbonitrides are considered to form mainly inside lath, although some are found to form along the boundaries because of the preferential nucleation on the dislocations during tempering. The addition of W to the steels sometimes enhances precipitation of intermetallic phases such as Laves and $\mu$ phases mainly along prior austenite grain boundaries and lath boundaries, and also inside grain during long-term exposure at high temperatures. FePd-based L1₀ type ordered phase $\alpha''$ has recently been found to form coherently inside the grains in the steels with Pd and to be useful to increase the creep resistance like MX carbonitride.

In this study, effects of alloying elements on creep properties of Pd added 9Cr ferritic steels have been evaluated in order to search for substitutional element for the precious metal Pd. Then pipe fabrication and welding tests were performed to confirm the possibility to apply Pd added 9Cr steel to large diameter and thick-wall boiler components.

2. Experimental Procedure

2.1. Fundamental Test

Table 1 shows the chemical compositions of the tested steels. Pd concentrations were varied from 0 to 1.5 mass%: B1 to B4. 0.5% Pd steel: B2 was selected as base steel, and various metals, from VIIA to IB except Fe and Tc, were added with 0.5 mass% to the base steel: B5 to B7, C1 to C6, D1, D2 and D4. 10 kg-ingots were prepared by vacuum induction melting. The ingots were hot forged and rolled into 16 mm square bars. All bars were normalized at 1 373 K for 0.5 h.

Softening behaviors of the steels were examined with tempering from 773 to 1 113 K for 1 to 20 h. After tempering at 1 043 K for 4 h, creep tests were conducted at 923 K with 120 and 100 MPa. The dimension of the creep test specimen was 8 mm in diameter and 42 mm in gauge.
length. Microstructure of the steels in the tempered and the creep-ruptured conditions were examined under scanning electron microscope (SEM). The amount of precipitates for the creep-ruptured specimens was estimated by the quantitative chemical analysis of extracted residues.

2.2. Pipe Fabrication and Welding Test

Table 2 shows the chemical compositions of the steels for this demonstration. From the fundamental test results, 0.5 mass% of Re was added to 9Cr–0.3Mn–9Cr–3W–0.7Pd–V, Nb, N, B steel: MC1 and MC2. Mod. 9Cr–1Mo steel: MA1 and the steel equivalent to B1: MB1 and MB2 were also prepared to the test as the references. Three 180 kg-ingots, MA1, MB1 and MC1, were hot forged and rolled into 25 mm wall thickness plate. Two 180 kg-ingots, MB2 and MC2, were hot forged and hot extruded into pipes. The dimension of the pipe is 84 mm in outside diameter and 12.5 mm in wall thickness. All plates and pipes were normalized at 1373 K for 4 h. MA1, MB1 and MC1 were tempered at 1043 K for 4 h and other were tempered at 823 K for 4 h followed by tempering at 1043 K for 4 h. This double tempering, combination of low temperature aging and high temperature tempering, is reported to give the optimum initial microstructure consisting of fine dispersion of the $\alpha'/$martensite and MX carbonitride inside lath deco-rated with M23C6 along lath, block and packet boundaries of martensite as well as along the prior austenite grain boundaries.12) In order to evaluate the applicability of Pd-bearing steel to the thick-walled pipes of fossil fired power plants, following tests were conducted. High strain rate tensile tests were performed at the plates with a condition of a heating rate of 100 K · min$^{-1}$, a holding time of 30 min, a temperature range from 1273 to 1548 K and a strain rate of 1 sec$^{-1}$. The plates were also welded with AWS ER Ni Cr-3 filler wire by gas tungsten arc welding (GTAW) and the properties of welded joints were examined. Figure 1 shows the cross section of weld groove. The welding conditions are summarized in Table 3.

Table 2. Chemical compositions of the steels for the demonstrational test. (mass%)

| Mark | C  | Si  | Mn  | Cr  | W  | V  | Nb | N  | B  | Pd | Others |
|------|----|-----|-----|-----|----|----|----|----|----|----|--------|
| MA1  | 0.10 | 0.34 | 0.43 | 8.7 | -  | 0.23 | 0.08 | 0.04 | 0.0001 | -  | Mo: 0.98 |
| MB1  | 0.08 | 0.30 | 0.51 | 9.0 | 2.9 | 0.20 | 0.05 | 0.05 | 0.05 | -  | -      |
| MB2  | 0.08 | 0.30 | 0.51 | 9.0 | 2.9 | 0.20 | 0.05 | 0.05 | 0.05 | -  | -      |
| MC1  | 0.10 | 0.04 | 0.36 | 9.0 | 2.9 | 0.19 | 0.05 | 0.03 | 0.05 | 0.70 | Re: 0.49 |
| MC2  | 0.10 | 0.04 | 0.37 | 9.1 | 2.9 | 0.19 | 0.05 | 0.03 | 0.04 | 0.69 | Re: 0.49 |

Note: MB2 and MC2 were for pipes. Others were for plates.

3. Results and Discussion

3.1. Fundamental Test

Figure 2 shows the tempering parameter dependence of the hardness for Pd-added and Pd-free steels. Pd addition hardened the steel at the temperature around 823 K and the hardness depended on the Pd concentrations. Figure 3 shows the secondary electron image of 1.5% Pd steel: B4 after tempering at 823 K for 20 h. About 100 nm-length precipitates, arranged in specific directions were observed inside martensite lath. This is because Fe–Pd L1$_0$ type ordered phase: $\alpha'$ precipitates coherently and disk-like inside the ferrite matrix: $\alpha$ with the following crystallo-graphic relationship: $\{001\} _{\alpha'} // \{001\} _{\alpha}$, $\langle 100 \rangle _{\alpha'} // \langle 110 \rangle _{\alpha}$. Hardness after 20 h tempering at 823 K of the tested steels is summarized in Fig. 4. 0.5 mass% addition of Mn, Ru,
Rh, Re, Os, Ir, Au or Pt hardened the 0.5% Pd steel to the 1% Pd steel level. These metals were expected to dissolve in $\alpha'$ and to increase the volume fraction of $\alpha'$ comparing to that of 0.5% Pd steel.

Figures 5(a) and 5(b) show the effect of alloying elements on the time to rupture and the minimum creep rate creep at 923 K with 120 and 100 MPa, respectively. Minimum creep rate axis is reversed so that the data correspond to the time to rupture. Corresponding to the changes of the hardness, additions of Ru, Re, Os, Ir, Au or Pt decreased the minimum creep rate and increased the creep rupture life to the level of 1% Pd steel, as shown in Fig. 5(a). Figure 6 shows the time dependence of creep rate for C1, C4 and C5 comparing with B1, B2 and B3. Addition of Re or Os not only decreased the minimum creep rate but also delayed the acceleration creep and prolonged the creep rupture life. Furthermore, C4 showed superior creep resistance to other steels under lower stress condition, as indicated by the smallest minimum creep rate in Fig. 5(b).

While, although increase of the Mn concentration hardened the steel after aging at 823 K, no improvement of creep re-

![Fig. 2. Tempering parameter dependence of the hardness showing the precipitation hardening around 823 K.](image)

![Fig. 3. Secondary electron image of B4 steel after tempering for 20 h showing the Fe–Pd L$_1^0$ ordered phase which is precipitated coherently to the bcc matrix.](image)

![Fig. 4. Comparison of hardness after 20 h tempering at 823 K for the tested steels.](image)

![Fig. 5. Time to rupture and minimum creep rate of the tested steels creep at 923 K with 120 MPa (a) and 100 MPa (b). C4, C5 and D2 in Fig. 5(b) are still testing.](image)
sistance was observed. True strain vs. creep rate curve of C1 steel is shown in Fig. 7. Increase of Mn decreased the creep rate at the early stage of primary creep to the level of B3, but the creep rate came close to B2 at the minimum creep and rapidly increased at the end of the tertiary creep.

B4 steel showed the degradation of creep properties by excess addition of Pd, as shown in Fig. 5(b). Figure 8 shows the time–strain curves of B2, B3 and B4 crept at 923 K with 100 MPa. A sudden rupture at the early stage of tertiary creep was observed in B4 steel. It was suggested that too much addition of fine precipitation such as $\alpha'$ and MX carbonitride deteriorated the creep ductility resulting in the reduction of creep rupture life.

Paying attention to C1 and C4, precipitation behavior of these steels was evaluated by quantitative analysis of extracted residue and microstructural observation by SEM. Figure 9 shows the amount of precipitates of B1, B2, B3, C1 and C4 after crept at 923 K and 120 MPa. The amount of Mn in precipitates depended on that of Pd in precipitates, suggesting the Mn dissolution in $\alpha'$. The Pd to Mn ratio in precipitates was about 0.5 so that the amount of Mn in precipitates was limited to only 0.2 mass%, which was less than the concentration of Mn in B2 steel, in the case of 0.5% Pd steel. It was suggested that the Mn addition promoted the precipitation of $\alpha'$, but in the saturated condition the amount of $\alpha'$ for C1 was almost equivalent to that for B2. As a result, C1 showed superior creep resistance to B2 in the early stage of primary creep but came close to B2 on the way to the minimum creep.

Figure 10 shows the back scattered electron images of ruptured specimen crept at 923 K with 120 MPa for B2, B3, C1 and C4. Prior austenite, packet and block boundaries of B2 and B3 were decorated with M23C6 (gray precipitates in Fig. 10) and Laves phase (bright one in Fig. 10). And there was no clear difference in boundary decoration between B2 and B3. Poor boundary decoration was observed in Mn increased steel. On the contrary, continuous precipitation along boundaries and fine Laves phase inside lath were observed in Re-added steel. It was assumed that the growth of the precipitations in C1 accelerated the tertiary creep, as shown in Fig. 7, and the continuous precipitations in C4 delayed the transition to tertiary creep, as shown in Fig. 6, by stabilizing the boundaries. It was also assumed that the fine Laves precipitates in C4 reduced the minimum creep rate.

![Fig. 6. Typical time vs. creep rate curves of the tested steel crept at 923 K and 120 MPa.](image)

![Fig. 7. True strain vs. creep rate curve of C1 steel comparing with B2 and B3 steels.](image)

![Fig. 8. Effects of Pd concentration on time–strain curves crept at 923 K and 100 MPa. Reduction of area is also indicated in the figure.](image)

![Fig. 9. Amount of precipitates for ruptured specimens crept at 923 K and 120 MPa.](image)
remarkably, as shown in Fig. 5(b).

3.2. Pipe Fabrication and Welding Test

From the fundamental test results, C4 steel was selected for this test. Chemical composition of MC1 and MC2 was slightly modified from C4, as shown in Table 2, concerning the improvement of creep properties. N was decreased to 0.03 mass% to decrease fine precipitation of MX instead of $\alpha'$ in order to increase the creep ductility. Mn was decreased slightly to prevent the growth of precipitations such as Laves phase. C, Si and W were also changed slightly to get the fully martensitic structure.

First, high temperature ductility was evaluated to judge the possibility of pipe fabrication by hot forging and extrusion process. Figure 11 shows the high strain rate tensile test results for MA1, MB1 and MC1. Reduction of area for MC1 kept more than 70% around hot forming temperature and the tensile strength was not remarkably higher than that of modified 9Cr–1Mo steel. In hot rolling and hot extrusion process, surface defects such as cracking occur, when the reduction in area of the steel is less than 60%.13) From these results, pipe fabrication was done and succeeded without harmful defects.

Then, the plates were welded by gas tungsten arc welding. All plates were also welded successfully without defects. Figure 12 shows the optical microstructure of the welded joints. The formation of $\delta$-ferrite was observed in near fusion line of MB1 joint. While MC1 joint showed fully martensitic structure even along the fusion line. The Charpy impact properties of the welded joints at 273 K are shown in Fig. 13. Although the values of absorbed energies of the MC1 joint were lower than that of modified 9Cr–1Mo steel joint, the values still fulfilled the requirements for the practical use.
4. Conclusion

In order to search for substitutional element for the precious metal Pd, effects of alloying elements on creep properties of Pd-added 9Cr ferritic steels have been evaluated. Following results were acquired:

1) Addition of Ru, Re, Os, Ir, Au or Pt hardened the 0.5Pd steel after tempering at precipitation temperature of $\alpha'$ and also improve the creep property to the 1% Pd steel level.

2) Increase of Mn concentration promoted the precipitation of $\alpha''$ but did not increase the amount of $\alpha'$, resulting in no improvement of creep properties.

3) Re addition prevented the growth of precipitation such as M$_2$C$_6$ and Laves phase, resulting in improving the creep properties at lower stress level.

From the above fundamental test results, demonstrational pipe fabrication and welding tests were performed for Pd and Re added steel. Pd and Re added steel was successfully formed into pipe and welded without harmful defects.

Detailed creep properties of fabricated plates and pipes will be reported in near future.

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