



# An evaluation of the thermal fatigue performance of three alloys for casting mould applications

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Paper written on project work carried out in partial fulfilment of B. Eng. (Metallurgical Engineering)

## Synopsis

A petrochemical company experiences premature thermal fatigue failure of the casting moulds used in catalyst production. The aim of the project was to find an alternative alloy that would outperform the current low-alloy cast steel used for the moulds. Based on their thermo-fatigue properties, 3CR12 ferritic stainless steel and H11 tool steel were chosen for testing and comparison with the currently used BS3100 B7 cast steel. Samples of each material were subjected to temperature cycling in a Gleeble 1500™ thermo-mechanical processing simulator, followed by surface analyses. The main parameters derived from the test work were the total true strain, the hot strength of the materials, and the number of cycles to failure. Additionally, the coefficient of thermal expansion for each material was measured using a Bähr dilatometer. H11 tool steel yielded the best performance by way of having the fewest surface cracks, the lowest total true strain per cycle, the most cycles to failure, the highest hot strength, and the lowest coefficient of thermal expansion.

## Keywords

thermal fatigue, fatigue failure, casting moulds, thermal expansion.

## Introduction

The Catalyst Manufacturing Plant (CMP) is a dusty environment with temperatures higher than ambient. Molten mill scale (magnetite,  $\text{Fe}_3\text{O}_4$ ) is tapped from the electric arc furnace into the casting moulds at temperatures close to 1600°C. After 10 seconds the magnetite is quenched with a water spray for 50 seconds to a minimum temperature of 400°C. Sometimes there is residual water in the pans from the cooling fog process (water sprays) in the empty mould. In a typical plant, the factory produces 40–44 batches a day containing between 150–300 moulds, depending on the time required for the specific batch (Buchholz, 2014). The typical batch in the CMP usually weighs about 6 t. The thickness of the tap, speed of the casting belt, and tilting angle all affect the number of moulds per batch.

One of the main problems faced during the molten mill scale casting is the premature failure of the casting moulds. Cracks initiate on the surface of the mould, which is in contact with the hot catalyst, and propagate into the interior of the mould. Cracks are generally concentrated in the middle area of the mould where the catalyst makes first contact with the mould, and which experiences

the highest temperature fluctuations. This suggests that the cracks are caused by thermal fatigue during the casting process.

In order to prolong the life of the mould and improve the operational efficiency, a recommendation for a new alloy or an improvement of the existing casting mould had to be made.

In many instances, there are no service life records for specific moulds to track their life span, as well as no records regarding the surface temperatures of the mould during the heating and cooling cycle. The operating conditions and safety concerns make it impossible to measure the surface temperatures. Therefore, the temperature fluctuations can only be estimated.

On average, the casting moulds last about 6 months in this operating environment. One of the primary indications that the mould has reached the end of its life is the sticking of the catalyst to the mould due to the enlarged surface area caused by the surface cracks, as seen in Figure 1. This type of surface is called the 'elephant skin' and is caused by thermal fatigue.

## Thermal fatigue failure

All materials of construction are affected by thermal fatigue. This is a result of cyclic stresses caused by variations in temperature. The damage occurs under repeated thermal cycling in the form of cracking in the metal, specifically where relative movement (differential expansion) is constrained (American Petroleum Institute, 2011).

The failure of the moulds in the current investigation can be classified as a low-cycle high-strain fatigue problem, typically with  $N < 10^4$  cycles. The number of cycles to failure is

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Figure 1—Thermo-fatigue surface of a BS3100 B7 mould

fewer than in typical high-frequency low-cycle fatigue. In these casting moulds, the thermal fatigue failure is controlled by a cyclic *plastic strain* rather than a cyclic stress. The cyclic plastic strain results from the thermal stresses, which are a consequence of the temperature fluctuations during the casting process. The number of cycles to failure,  $N_f$ , for this type of fatigue failure can be predicted by the Coffin-Manson equation.

This approach is similar to what is known as the Paris equation for the stable crack growth regime. The Paris equation for high-frequency low-cycle fatigue is

$$\frac{da}{dN} = C_2 \Delta \varepsilon^m \quad [1]$$

where

$C_2$  = Constant

$\Delta \varepsilon$  = Cyclic strain amplitude.

Low-cycle fatigue test data are often represented as a plot of the plastic strain range *versus* cycles to failure. When plotted on log-log coordinates, a straight line is obtained, which is represented by the Coffin-Manson relation:

$$\Delta \varepsilon_p = C_2 (N_f)^\beta \quad [2]$$

where  $\Delta \varepsilon_p$  is the plastic strain range,  $N_f$  is the number of cycles to failure (fatigue life),  $C_2$  is a constant and  $\beta$  is the fatigue ductility exponent (generally  $-0.5 > \beta > -0.7$ ) (Coffin Jr., L.F., 1954; Manson, S.S., 1954)

### Experimental procedure

The physical properties of the three steels that were investigated are given in Table I. The selection of 3CR12 and H11 steels was based on their melting points, coefficients of thermal expansion, and thermal conductivities. The chemical compositions of the steels are given in Tables II.

The thermal fatigue simulation was carried out in a Gleeble 1500™ thermo-mechanical processing simulator. This apparatus enables the specimen to be subjected to either isothermal compression or tensile tests at elevated temperature. However, in this investigation, the specimens were subjected to temperature cycling without constraint.

Three samples, 10 mm in diameter and 116.5 mm in length, were prepared of each material (Figure 2). The test zone at the centre of each sample was milled to a thickness of 7 mm for entire gauge length of 50 mm so as to create sharp corners simulating the edges of the mould. Two K-type

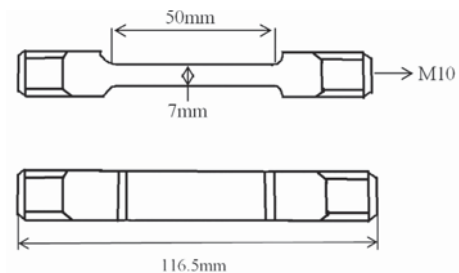


Figure 2—Schematic illustration of sample for testing

Table I

### Physical properties of the three steels

Material	Melting temperature (°C)	Thermal expansion coefficient ( $\mu\text{m}/\text{m}\cdot^\circ\text{K}$ )	Thermal conductivity ( $\text{W}/\text{m}\cdot^\circ\text{K}$ )
BS 3100 B7	-	11.9	42
H11 tool steel	1427	13.3	28.6
3CR12 ferritic stainless steel	1430–1510	12.8	40

Table II

### Alloy compositions\*, %

	C	Cr	Mn	Mo	Ni	Cu	Co	Si	Ti	Al	V	P	N	Nb	Ca	S	W	B
BS 3100 B7 <sup>a</sup>	0.17	2.25	0.75	1.00	0.4	-	-	0.35	-	-	-	0.025	-	2	3	0.02	-	-
3CR12 <sup>b</sup>	0.019	11.01	0.81	0.03	-	0.1	0.02	0.72	0.13	0.017	0.08	0.022	0.012	0.001	-	-	-	0.001
H11 <sup>c</sup>	0.37	4.95	0.36	1.26	0.22	0.13	0.01	1.08	-	0.019	0.95	0.023	-	-	0.0014	0.003	0.04	-

\*Balance Fe

<sup>a</sup>Buchholz, 2014

<sup>b</sup>ASTM A240 and EN 10088-2 (Columbus Stainless, 2015)

<sup>c</sup>Du Toit, 2015

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thermocouples were spot-welded onto each specimen: one in the middle and one further away from the centre of the specimen, to measure the temperature. Each test was run for 100 cycles over approximately 3 hours. In a case where the cracks appeared before 100 cycles were completed, the experiment was stopped prematurely. The temperature was fluctuated between 400 and 1200°C. In the first 50 seconds, the specimen was heated to 1200°C at a heating rate of 20°C/s, followed by another 10 seconds of soaking at the elevated temperature. The specimen was then cooled to 400°C at a cooling rate of 20°C/s. This cycle was repeated 100 times.

After completion of the test, the surface of each specimen was examined for cracks or other irregularities. Photographs of the surface were taken, before and after cleaning with Hibatex solution to remove oxides, using a stereo microscope at magnifications of 12× and 15×. The data generated was used to calculate the total plastic strain in 100 cycles. Three tests were conducted for each steel, and an average value for the total plastic strain after 100 cycles was obtained. Equation 3 was used to calculate the total plastic strain,  $\epsilon$ , after 100 cycles.

$$\epsilon = \ln\left(\frac{\Delta L}{L_o}\right) \quad [3]$$

where  $\Delta L$  is the change in length (m) and  $L_o$  is the gauge length of the specimen (m).

The hot strength of the steels was determined at 1100°C through an isothermal hot compression test, see Figure 7. The strain was limited to 0.8 to avoid barrelling effects.

The coefficient of thermal expansion was determined using the Bahr dilatometer. This test was done in accordance with ASTM E831-14 specifications. The thermal expansion coefficient  $\alpha$  is a material constant. It is the amount of expansion or contraction a material undergoes per kelvin during heating or cooling and is given by the following equation:

$$\alpha = \frac{dL}{LdT} \quad [4]$$

where

$dL$  = Change in length ( $\mu\text{m}$ )

$dT$  = Change in temperature respectively ( $^{\circ}\text{K}$ )

$L$  = Total length of the specimen (m)

$\alpha$  = Thermal expansion coefficient ( $\mu\text{m}/\text{m}\cdot^{\circ}\text{K}$ )

The thermal expansion coefficient should be as low as possible when the material is used to make moulds used in the casting process.

The specimens were 5 mm in diameter and 10 mm long. Three of these specimens were machined for each alloy. The sample was heated to 1220°C (above  $A_{c3}$ ) at a heating rate of 0.25°C/s. The specimen was then soaked for 60 seconds and afterwards cooled to room temperature at a cooling rate equivalent to that of the heating rate. Lastly,  $A_{c3}$  is the temperature at which ferrite completes its transformation to austenite during the heating up cycle.

### Results and discussion

#### Surface evaluation

Figures 3, 4, and 5 show stereo micrographs for BS3100 B7, 3CR12, and H11 after 100 thermal cycles between 400 and 1200°C.

As expected, the surface of BS3100 B7 was heavily oxidized. After removing the oxides, cracks were exposed and were visible in the centre area in both longitudinal and transverse directions (Figure 3). 3CR12, in contrast, exhibited less oxidation than BS3100 B7. However, cracks were visible in the transverse direction in Figure 4. Strain marks (material flow lines) in the lengthwise direction were also present. No visible cracks or irregularities like strain markings or necking were observed in H11 (Figure 5). The specimen was still in sound condition after 100 cycles.

Figure 6 shows a summary of the total plastic strain for each steel after 100 thermal fatigue cycles. The total true plastic strain per 100 cycles for BS3100 B7, 3CR12, and H11 was 0.148, 0.182, and 0.058 respectively. The total plastic strain represents the measure of resistance to thermal fatigue.



Figure 3—BS3100 B7 specimen subjected to thermal cycling after cleaning, showing longitudinal and transverse cracks



Figure 4—3CR12 specimen subjected to thermal cycling after cleaning, showing transverse cracks



Figure 5—H11 specimen subjected to thermal cycling after cleaning

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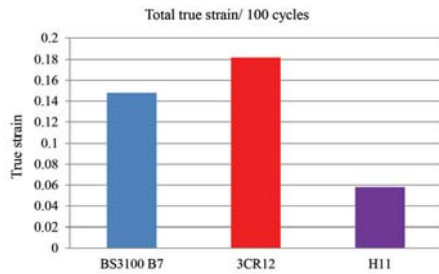


Figure 6—Comparison of the results of the total true strain per 100 cycles

## Estimated number of cycles to failure ( $N_f$ )

The average number of cycles to failure for BS3100 B7, based on the industrial data, is 11 340 *i.e.*  $N_f$  (BS3100 B7) = 11 340 cycles. The cycles to failure for 3CR12 and H11 are unknown. However, the ratios of the cycles to failure for BS3100 B7:3CR12 and BS3100 B7:H11 can be determined based on the total true strain per 100 cycles. The number of cycles to failure for 3CR12 and H11 can then be calculated by multiplying the respective ratios by  $N_f$  (BS3100 B7). The results are shown in Table III.

Although H11 tool steel looks promising, it is more expensive than BS3100 B7. Therefore, before H11 can be recommended as an alternative material, a comprehensive cost benefit analysis would be required. Based on a rough calculation regarding the raw material cost, the cost of H11 is 3.5 times higher than that of BS3100 B7 (Table IV).

## Hot-strength: high-temperature steady-state stress

Figure 7 shows the Von Mises true stress *versus* true strain for the three steels. H11 exhibited the highest hot strength of 31 MPa. The hot strengths of 3CR12 and BS3100 B7 were lower than H11, and similar *i.e.* 16 and 18 MPa respectively.

## Coefficient of thermal expansion

Figure 8 shows a typical dilatometer-generated plot of change in length *versus* temperature. The  $\alpha$  values for BS3100 B7, 3CR12, and H11 were found to be 15.8, 13.1, and 12.8  $\mu\text{m}/\text{m}^\circ\text{C}$  respectively.

## Conclusions

The thermal fatigue behaviours of BS3100 B7, 3CR12, and H11 were compared under similar conditions.

H11 had the lowest total true strain of 0.058 per 100 cycles, indicating that it had the highest resistance to thermal fatigue compared with 3CR12 and BS3100 B7. The mould life would be longer than for the other two steels, and was estimated to be about 35 478 cycles.

However, before H11 tool steel can be recommended as an alternative material, a comprehensive cost analysis is required. Based on a rough calculation regarding the raw material cost, the cost of H11 is 3.5 times higher than that of BS3100 B7.

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Table III

## Estimated number of cycles to failure for 3CR12 and H11 using the total true strain per 100 cycles

Steel type	Ratio based on the total strain ( $\Delta\epsilon_p$ ) per 100 cycles	$N_f$ predicted
3CR12	1:1.3	13 931
H11	1:3.1	35 478

Table IV

## Raw material costs for BS3100 B7 (van Aarde, 2015) and H11 tool steel (Alibaba Group, 2015)

Steel type	Raw material cost (R/ton)
BS3100 B7	9 500
H11	32 550

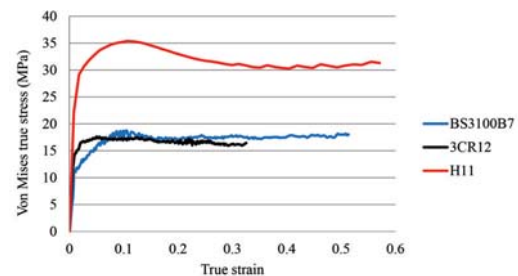


Figure 7—Von Mises true stress vs. true strain curves for BS3100 B7, 3CR12, and H11 at 1100°C and strain rate of 0.001/s

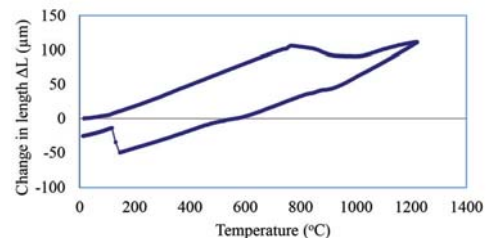


Figure 8—Change in length vs temperature curve for BS3100 B7 for the determination of  $\alpha$

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# The effect of particle size on the rate and depth of moisture evaporation from coal

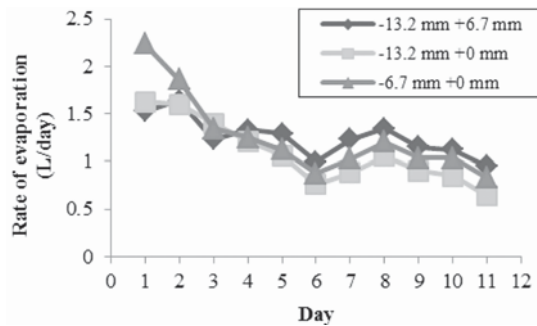


Figure 5—Daily rates of evaporation from each size range

It can be seen that all coal samples reached the same moisture content by the end of the experimental run. The coarse coal sample reached the final moisture content first.

To compare the effect of particle size on the rate of evaporation, the daily rates of evaporation were calculated for each size range as shown in Figure 5. It can be seen that moisture initially evaporates faster from the fine coal sample (-6.7 mm) than from the other coal samples. After two days, the surface of the fine coal sample was dry and the rate of evaporation decreased. The same phenomenon was seen in the experiments that investigated the depth to which evaporation occurs. This is attributed to the porous structure of the coarse sample.

During the investigation, it was found that the initial moisture content had an influence on the rate of evaporation. A lower initial moisture content resulted in a lower maximum rate of evaporation.

## Conclusions

This study showed that bed particle size affects the extent of moisture evaporation, and that the porous structure of a coarse coal stockpile aids the process. Experimental results indicate that moisture drainage takes place within the first two days of exposure to environmental conditions. Evaporation of moisture can be seen up to the fourth day, but this is a function of the initial moisture content of the coal. Coal with a higher initial moisture content will take a longer time to lose all surface moisture.

It was shown that water can evaporate from underneath the surface of coal stockpiles. Increased void spacing has a positive influence on the effectiveness of evaporation for coal stockpile drying. The results obtained in this study can be used to manage stockpiles efficiently. Up to 0.4 m of coal can be reclaimed from the stockpile surface and the remaining coal can be left to dry for another four days – if no rain occurs.

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