Rheo-Squeeze Casting of High-Silicon Aluminium Alloy Pipes with Gradient Structures and Their Mechanical Properties

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Received: March 03, 2018; Revised: April 13, 2018; Accepted: May 09, 2018

Three different compositions of high-Si Al alloy pipe, with a gradient structure, were produced through semi-solid rheological squeeze casting, the microstructural characteristics of which were investigated. The mechanism of gradient structure formation and the effect of acicular Fe-rich phases on liquid segregation were revealed. Meanwhile, the crystal structures of the Al FeSi and Al (Fe, Mn)Si phase were elucidated. The relationship between the volume fraction of hard particles in the pipe wall and the macro-hardness was established. Wear resistance tests of the inner surfaces of the pipe walls, and tensile strength tests at different positions around the pipe walls, showed that the δ-AlFeSi phase (acicular Fe-rich phases) reduced the wear resistance and the tensile strength of the alloy. In this study, the gradient distribution of the hard particles caused the outside of the pipe wall to have a higher tensile strength.

Keywords: high-Si Al alloy, rheo-squeeze casting, gradient structure, crystal structure information, mechanical property.

1. Introduction

Al-Si alloys have good casting properties, stable high-temperature performance, and its casting has good air-tightness. Using Al-Si alloys to make fuel engines has become the trend in automobile manufacturing industry since the 1970s. As the content of Si exceeds 12% in Al-Si system alloys, they exhibit excellent wear resistance and heat resistance. So, parts of the engine wear-resistant components are manufactured with high silicon aluminium alloys (Si%>17%), such as: the piston, cylinder liner, etc. Meanwhile, primary Si particles must be modified and Fe, Mn, Cu, and Mg elements should be considered for addition to the alloy system.

Relevant research shows that the application of a high-Si Al cylinder liner could increase engine power and decrease oil consumption in the diesel engines. For improving the wear resistance of the cylinder bore of the engine, additional coating with wear-resistant materials on the surface or casting iron cylinder liner inserts for the cylinder block are effective. The similarity in these approaches is that they establish a wear-resisting functional layer between the cylinder liner and the cylinder block to enhance the efficiency and the service life of the engine.

Considering the cost of manufacture, the high-Si Al alloy cylinder liner, with a gradient structure in the radial direction was produced by the centrifugal casting process. This kind of cylinder liner has a gradient structure containing a Si-rich area on the inner surface and a Si-poor area on the outer surface. The Si-rich area could provide the necessary wear resistance and the Si-poor area on the outer surface has a similar chemical composition to that of the cylinder block. The similar chemical compositions of the cylinder liner and cylinder block benefits the smooth transition of physical and metallurgical properties at the interface between the two.

On the other hand, a high-chromium cast iron pipe with a gradient structure produced by squeeze casting had, like other functionally graded materials prepared through semi-solid squeeze casting processes, linear distributions of the phases or the composition, thus avoiding step-changes therein.

In previous studies, the microstructures of high-Si Al alloys were analysed, involving the technology of alloy slurries preparation and the crystal structures of Fe-rich phases. The present work aimed to squeeze cast the high-Si Al alloys as semi-solid slurries with different microstructural characteristics, to prepare a pipe with a gradient structure in the radial direction. The effect of microstructural characteristics on the distribution of the phases was studied. Crystal structure information about the Fe-rich phases was further enriched. The wear resistance of the inner surface of alloy pipes, and the tensile strength of the alloy pipes, were investigated.

2. Experimental Work

Three types of high-silicon aluminium alloys were selected as experimental objects. The chemical composition of these Al22Si-xFe-xMn alloys is listed in Table 1. The Al22Si-2Fe-xMn alloys were prepared from pure Al, 95% Si, 75% Fe, and 75% Mn in an SG-5-12 resistance furnace. Pure Al was heated to 850 °C and held for 15 minutes, and
then rapid melting-Si, which was wrapped in aluminium foil, was added to the Al melt in batches. After stirring the mixtures uniformly, Fe and Mn agents were added and stirred uniformly, and the sample retained for 30 minutes. After cooling of the melt to 800 °C, the C₂Cl₆ was added (to an amount that was 0.5% of the melt by mass) to remove gas and slag. Semi-solid slurries of the alloys were prepared by using self-made segmented cooling plates when the melt was cooled to 780 °C. The liquidus and solidus temperatures, as measured by DSC, were approximately 777 °C and 570 °C (the addition of 2% Fe and 1.46% Mn had little effect on the liquidus and solidus temperatures of these alloys).

| Table 1. Chemical compositions of the experimental alloys (wt. %) |
|-----------------|-------|-----|-----|-----|
| Sample | Si/% | Fe/% | Mn/% | Al/% |
| A₁ | 22–23 | 0 | 0 | Balance |
| A₂ | 22–23 | 2.10 | 0 | Balance |
| A₃ | 22–23 | 2.21 | 1.46 | Balance |

Semi-solid squeeze casting was carried out in a 50 ton-hydraulic machine. The semi-solid slurry was transferred into a mould preheated to 200 °C (Fig. 1). The slurry was pressed at 50 MPa for approximately 10 to 15 s. The squeezed Al alloy pipe was split along the axial direction into two halves (Fig. 2).

The samples were prepared along the longitudinal direction of the squeezed pipe. After pre-grinding, fine-grinding, and polishing, the specimens were etched with 0.5% hydrofluoric acid. Optical microscopy (Leica, Co.) was used to observe the metallographic structure. Measurements of the equivalent diameter (ED) of the hard phases were carried out using image-analysis techniques. ED was determined from the following equation:

\[ D = 2 \sqrt{\frac{A}{\pi}} \]  

Where: \( A \) is the area of the hard phase particle, we can obtain it by the Image-pro plus software suite.

A scanning electron microscope (SEM) (VEGA-3SBH, Tescan Co. & S-3400N, Hitachi Co.) was used to observe the microstructures, fracture morphology, and abrasion morphology. The phase analysis and the structure of Fe-rich phases of the alloys were analysed by using a Focused Ion Beam (FIB) for sampling (Quanta 3D FEG SEM/FIB Dual Beam, FEI Co.) and Transmission Electron Microscopy (TEM) (Tecnai G2-F30-S-TWIN, FEI Co.), respectively.

Five points were chosen along the radial direction to test the macro-hardness of the pipe wall with its gradient structure. The micro-hardness of the Si- and Fe-rich phases was measured using a Vickers hardness tester at 25 gf. Fig. 3(a) shows the sampling positions of the squeeze pipe for the abrasion test and the testing of the tensile strength thereof. Fig. 3(b) shows the schematic diagram of the abrasion test of the worn surface inside the pipes. The tested pin of alloy against Cr13 quenched steel disk with a hardness of 54 to 58HRC was also undertaken. The surface roughness of abrasion samples and the test pin were less than Ra0.8. The experiments were conducted under a 20 N normal load and at a sliding speed of 500 rpm for 1800 s and 7200 s respectively, with a worn radius of 11.5 mm. As a result, the worn distance was 1080 m and 4333.2 m respectively when the sliding speed was 500 rpm. The tensile strength and elongation testing were carried out on the AG-IS10KN electronic universal testing machine at a displacement rate of 0.5 mm/min. A typical test sample is shown in Fig. 3(c).
3. Analysis of the Microstructures of Semi-solid Squeeze Cast Alloy Pipe Walls

3.1. Microstructural characteristics of semi-solid squeeze cast alloy pipe walls

The microstructures of semi-solid rheological squeeze casting Al alloy pipe wall are shown in Fig. 4. Fig. 4(a) presents the assembled microstructures of radial regions in the pipe wall, and the colour thereof gradually deepened to dark grey from the outer surface to the inner surface. Through previous research, the grey microstructures are primary Si particles and eutectic Si and the white microstructures are \( \alpha \)-Al phases. The equivalent diameters (ED) of PSPs in regions b to f were 16, 18, 18, 20, 22 ± 2 \( \mu \)m, respectively. As shown in Fig. 4 (b - d), PSPs and eutectic Si have little difference.
in size (regions b - d). Whereas, the ED of PSPs is slightly increased from regions e to f (seen in Fig. 4(e - f)), and the PSPs were largest in the inside region of the pipe wall. The nearer to the inner surface of pipe wall, the smaller the volume fraction (VF) of the α-Al phases: meanwhile, several “segregation islands” (marked by red circles) containing liquid phases appear in the microstructures in regions e and f.

The microstructures of the A2 alloy (with addition of elemental Fe) pipe wall are shown in Fig. 5. Fig. 5 (a) shows the radial assembly microstructures of an A2 alloy pipe wall. The EDs of PSPs in regions b to f (shown in Fig. 5(b - f)) are similar to that in the A1 alloy pipe wall. Compared with the A1 alloy pipe wall, however, the overall trend of the distribution of grey particles in the A2 alloy wall presented a gradual increase, different sized “segregation islands” (Si-poor regions) were present in the radial section of the pipe wall (red circles), in which the distribution of primary Si was discontinuous. It was noteworthy that the number of “liquid segregation islands” was greater than that in A1 alloy, and the acicular Fe-rich phases (Al₄FeSi₂ phases) were distributed around the boundaries of the “liquid segregation island”.

As shown in Fig. 6, no acicular Fe-rich phases were seen in the radial microstructures of the A3 alloy (with its added elemental Fe and Mn) pipe wall, replaced by the blocky Al₁₅(Fe, Mn)₃Si₂ phases. The micro-hardness of the Si particle and blocky Fe-rich phase were measured as 1279 and 276 HV respectively. The two phases could be regarded as hard particles relative to the Al matrix, for which the micro-hardness was 122 HV. Due to the addition of Fe and Mn, the VF of hard phases in each region of A3 alloy wall was higher than that in the A1 and A2 alloys, and the acicular Fe-rich phases were modified into block-like forms by the addition of Mn, and the EDs of the PSPs in each regions are 2 µm higher than that in A1 and A2 alloy pipe walls (calculated from Fig. 6(b - f)). Meanwhile, the number of “liquid segregation islands” in the A3 alloy wall is less than that in the A2 alloy.

As shown in Fig. 7, comparing the hard phases in the regions of each alloy, the EDs of all samples underwent a gradual increase from the outer surface to the inner surface of the three types of alloy pipe walls: with the increase of the VFs of the hard phases from regions b to f, the corresponding macro-hardness of each region in the pipe wall also behaved in a similar fashion, but this phenomenon only occurred in the A1 and A3 alloy walls, whereas, the VF of the hard phases in regions b to f did not increase gradually in the A2 alloy wall. The corresponding volume of the region of increasing hardness in regions b to f did not increase gradually in the A2 alloy wall but exhibited a decreasing trend, especially in regions e and f.

**Figure 4.** Rheo-squeeze casting A1 alloy pipe wall with gradient structure (a) puzzle of A1 alloy pipe wall microstructures; (b - f) microstructures of regions b - f.
Under the influence of the blocky $\text{Al}_{15}(\text{Fe}, \text{Mn})_3\text{Si}_2$ phases, the VF of the hard phases in each region of the A3 alloy wall is higher than that in the corresponding regions of the A1 and A2 alloy walls, which makes the volumes of the higher hardness region of the A3 alloy wall greater than that of the corresponding regions in the A1 and A2 alloy walls. In summary, the macro-hardness of each region in the alloy wall is significantly associated with the VFs of hard phases of each region.

3.2. Preliminary analysis of the mechanism underpinning the formation of the gradient structure alloy pipe wall produced through semi-solid squeeze casting

In comparison with Figs. 4 ~ 7, it was found that the EDs and VFs of the hard particles (primary Si and the blocky $\alpha$-$\text{Al}_{15}(\text{Fe}, \text{Mn})_3\text{Si}_2$ phase) gradually increased from the outer surface to the inner surface of the pipe wall. In the first of filling, the driving force of the liquid phase flow during semi-solid rheo-squeeze casting process is the pressure difference in the various parts of the sample. The liquid phase tends to flow towards the lower volume under pressure. At a certain squeeze speed, the slurries of the alloys flow along the surface of the mould core to the bottom of the cavity. Then, in the radial direction, the fine primary Si particles can be carried to the wall of the mould with the liquid phase. Finally, the slurries condensed under pressure and a pipe with a radial gradient of particle forms and sizes was produced.

There are a large number of acicular Fe-rich phases in the A2 alloy, which hinder the flow of the liquid phase and cause flow along the edge of acicular Fe-rich phase particles. This was why a liquid phase flow belt formed and, as can be seen from Fig. 6(c) and (d), the acicular Fe-rich phase was distributed along both sides of the flow belt, and the VF of the liquid difference of both sides of some acicular Fe-rich phase crystal was obvious. This caused liquid segregation of the A2 alloy pipe to be the most significant among the three types of alloy pipe walls tested.

3.3. Analysis of the crystal lattice parameters of different types of Fe-rich phases

Based on previous studies, the $\text{Al}_4\text{FeSi}_2$ phases (acicular Fe-rich phases) obtained under high cooling rates were identified as having a tetragonal structure, and the blocky or Chinese script-$\text{Al}_{15}(\text{Fe}, \text{Mn})_3\text{Si}_2$ phases (precipitated out in the eutectic reaction) was identified as having a hexagonal structure. In this study, the $\text{Al}_4\text{FeSi}_2$ phase and the blocky...
Figure 6. Rheo-squeeze casting A3 alloy pipe wall with gradient structure (a) puzzle of A3 alloy pipe wall microstructures; (b - f) microstructures of regions b - f.

Figure 7. Relationships between the macro-hardness and the hard particles’ VFs of the pipes.

Al_{15}(Fe, Mn)_{3}Si_{2} phase were analysed. Fig. 8 shows the TEM bright-field images and selected area diffraction patterns of the Al_{15}FeSi_{2} phase and the laminar Al_{15}(Fe, Mn)_{3}Si_{2} phase (sampled from the blocky phase by focused ion beam). The analysis showed that the Al_{15}FeSi_{2} phase was in the form of a tetragonal structure with lattice constants $a = 0.60641$ nm and $c = 0.95258$ nm, and the blocky Al_{15}(Fe, Mn)_{3}Si_{2} phase was a body centred cubic structure with lattice constants $a = 0.75198$ nm and $c = 0.77688$ nm.

Figure 8. TEM images of Fe-rich phases (a) bright-field and SAPD of acicular Fe-rich phase; (b) bright-field and SAPD of blocky $\alpha$-Al_{15}(Fe, Mn)_{3}Si_{2} phase sampled by FIB.
4. Effect of Fe-rich Phase Morphology on Wear Resistance and Tensile Strength of the Pipe Wall

It can be seen from Table 2 that the mass loss and wear rate of A2 alloy were the largest in the wear testing of three types of alloy pipe inner surface, while the A3 alloy was the smallest. With increasing wearing test time, the wear rate of the A2 alloy increased most among the three alloys. Therefore, as the data showed, the inner surface of the A3 alloy pipe had the best wear resistance, followed by that of the A1 alloy, with A2 alloy being the worst.

Fig. 9(a) shows the wear morphology of A1 alloy after 1800 s under a load of 20 N at a speed of 500rpm. There were shallow furrows and a lower plastic shear lip in the wear surface. There were fewer spalling pits and less tearing at the top of the plastic shear lip. There was a small amount of fine grinding particles on the wear surface under these test conditions.

Fig. 9(b) shows the wear morphology of the A1 alloy after 7200 s under a load of 20 N at a speed of 500rpm. The uneven furrows can be seen in Fig. 9(b), and some of the furrows are deeper. There was evidence of a plastic deformation shear lip on both sides of the furrow (indicated by the arrow in the figure), the top of which had deep spalling pits and tearing. There were a large number of the bigger grinding particles seen on the wear surface. The particles were formed by the spalling of primary Si during repeated rolling; because of the excessive load in the wear process, deformation of the worn surface occurred, and the harder particles acted repeatedly and fractured during the prolonged friction wear process, causing the primary Si particles at the top of the shear lip to fall off and form a spalling pit. The adhesion and rolling action of the two contact friction surfaces in the process of friction wear meant that the larger shear stress tore the plastic shear lip, which was the key fatigue wear mechanism. The falling Si particles gathered in the groove, and played the role of micro-cutting to deepen the groove, which was the key abrasive wear mechanism. The wear of A1 alloy was not a single wear mechanism but a combined action of adhesive wear and abrasive wear under long-term, high-speed friction.

Fig. 9(c) and (d) show the wear morphologies of the inner surface of the A2 alloy pipes. Fig. 9(c) shows the surface morphology of wear samples under a load of 20 N after 1800 s. It can be seen that there are many sheet like fragments falling off the matrix in the wear surface. Some of the fragments were distributed in the top part of the plastic deformation area, and some of the fragments were partly distributed around the bottom of the furrows. There were larger wear particles at the bottom of the furrows. The wear surface morphology of the inner wall of the A2 alloy pipe after wear testing for 7200 s is shown in Fig. 9(d). The surface was badly worn and there were only a few furrows on the surface. The surface of the worn sample spalled and many particles covered the surface abscission zone. Many of these particles had been crushed to form an oxide particle layer. The wear resistance of the inner surface of the A2 alloy pipe was the lowest because the worn surface suffered the spallation of sheet-like fragments during the wear process.

Fig. 9(e) and (f) show the wear morphologies of the inner surface of the A3 alloy pipes. Fig. 9(e) shows the surface morphology of wear samples under a load of 20 N after 1800 s: the plastic shear lip in the figure is obvious, and the top part of the shear lip evinced tearing of the material. There were varying shades and width of furrows in the wear surface. There were a few wear particles at the bottom of the furrows, and there were abrasive particles seen at the edges of the furrows, which were formed by rolling of the detached particles, therefore, under these conditions, the wear mechanism of the inner surface of the A3 alloy pipe was mainly adhesive wear, accompanied by a certain abrasive wear mechanism. Fig. 9(f) shows the surface morphology of wear samples under a load of 20 N load after 1800 s: there were unevenly distributed furrows and wear particles of different sizes on the wear surface. Therefore, under these conditions, the wear mechanism was mainly abrasive wear, accompanied by slight adhesive wear.

Through tensile test, that the tensile strength of the inner sides of the three types of alloy pipes were 96.55 MPa, 76.73 MPa, and 95.96 MPa, respectively (the volume fraction of the hard phase was higher and the average diameter was larger on the inside). The A2 alloy pipe had the lowest tensile strength as it contained a large number of acicular Fe-rich phases: the A1 and A3 alloy tensile strengths were similar. The tensile strengths of the outer part of the three alloys were increased compared with the inner part, which were 125.58 MPa, 83.80 MPa, and 112.54 MPa, respectively. It should be noted that the VF of the hard phase was lower and the ED was smaller in the outer part. It can be concluded that the tensile strength of the inner and outer sides differed due to the VF of hard phases, that is, the higher the VF and the larger the ED, the lower the tensile strength, and vice versa. The appearance of the bulk α-Al_{15}(Fe, Mn)Si_{3} phase in the micro-structure had little effect on the tensile strength of the alloy, but the acicular Fe-rich phase impaired the tensile properties of the alloy.

Fig. 10 shows the SEM micrograph of the tensile fracture surface of the inner part of the A1, A2, and A3 alloy pipes. The tensile fracture surface of alloy A1 had a large number of dimples of different depth (arrow points) and conical protrusions, as shown in Fig. 10(a). The edges of the dimples were sharp. At the same time, many planes similar to a cleavage platform could be seen on the fracture surface, therefore, the fracture surface of the specimen was deemed to have arisen from brittle fracture. Fig. 10(b) shows the tensile fracture of the A2 alloy. There were many dimple structures and conical protrusions on the surface of the alloy,
Figure 9. Worn morphologies of the inner faces of pipes’ walls: (a, c, e) the worn surfaces of A1, A2, A3 alloy pipe walls for 1800s; (b, d, f) the worn surfaces of A1, A2, A3 alloy pipe walls for 7200s.

Table 2. Wear test data of the inner face of the A1, A2 and A3 alloy pipes

| Load and wear speed | Wear time/s | A1 alloy | A2 alloy | A3 alloy |
|---------------------|-------------|----------|----------|----------|
| 500r/min, 20N       | 1800s       | 13       | 0.46%    | 30       | 1%       | 6        | 0.48%    |
|                     | 7200s       | 62       | 2.14%    | 90       | 3.2%     | 58       | 2.03%    |
and a smooth tearing section can be seen at the position indicated by the arrow. Fig. 10(c) shows the tensile fracture of the A3 alloy. There were many cleavage platforms and a large number of dimples on the surface, therefore, the A3 alloy fracture was also deemed to have been a brittle fracture.

By means of Fig.10, it can be deduced that there were significant stresses acting in the specimens during the tensile testing of the A1, A2, and A3 alloy pipes and that these were mainly parallel to the direction of applied tensile load. Many micro-cracks were seen to be distributed on the surface of the primary Si particles and blocky α-Al\textsubscript{15}(Fe, Mn)\textsubscript{3}Si\textsubscript{2} phase, and the stress concentration was the most obvious at the tip of the crack. The main cracks formed at these micro-cracks and propagated under the action of the applied tensile force. The crack would have continued to propagate along the acicular Fe-rich phase boundary and the edge of the lamellar structure when the main cracks propagated to the acicular Fe-rich phase or the lamellar eutectic Si structure. A fracture was formed perpendicular to the direction of application of tensile force after crack confluence.

5. Conclusions

High-Si Al alloy pipe with gradient structure can be produced through semi-solid squeeze casting. During the process of semi-solid slurry filling, the slurry flow along the surface of the mould core to the bottom of the cavity. Then, in the radial direction, the fine primary Si particles are carried to the wall of the mould with the liquid phase. However, acicular Fe-rich phases (Al\textsubscript{4}FeSi\textsubscript{2} phases) hinder the flow of the liquid phase and cause liquid segregation in the microstructures of the pipe wall. Thereby, this effects the gradient varying of the macro-hardness at the pipe wall, and the Al\textsubscript{4}FeSi\textsubscript{2} phases even worsen the wear resistant and tensile properties of alloy. The blocky Fe-rich phases (Al\textsubscript{15}(Fe, Mn)\textsubscript{3}Si\textsubscript{2} phases) are in favour of the wear resistant property, and have little impact on tensile properties of alloy wall. Meanwhile, the TEM analysis showed that the Al\textsubscript{4}FeSi\textsubscript{2} phase was in the form of a tetragonal structure with lattice constants \(a = 0.60641\) nm and \(c = 0.95258\) nm, and the blocky Al\textsubscript{15}(Fe, Mn)\textsubscript{3}Si\textsubscript{2} phase was a body centred cubic structure with lattice constants \(a = 0.75198\) nm and \(c = 0.77688\) nm.

6. Acknowledgement

The authors acknowledge funding for this research from National Natural Science Foundation of China (51505205), Science Foundation of Kunming University of Science and Technology (KKZ3201532005). This work is supported by the National and Local Joint Engineering Laboratory of Advanced Metal Solidification Forming and Equipment Technology, Kunming University of Science and Technology. Kunming, China.
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