Wet regeneration of waste artificial sand used in sand casting using chemical solutions

Eun Yeong Ko1,2, Kyeong Ho Kim1, Jae Ho Baek1, Inseong Hwang2, Man Sig Lee1,3†

1Ulsan Division, Korea Institute of Industrial Technology (KITECH), Korea
2Department of Civil and Environmental Engineering, Pusan National University, Korea
3Green Process & System Engineering Department, University of Science & Technology (UST), Korea

ABSTRACT
Natural sand and organic binders have been the main materials used in sand casting. However, inorganic binders are used in novel casting technologies, and research on artificial sand as a substitute of natural sand has been conducted. Herein, the wet regeneration of waste artificial sand was performed using a sodium silicate binder. Acidic, neutral, and basic solutions were used for wet regeneration. The effects of binder removal and reusability of the regenerated sand were investigated by comparing the characteristics and strengths of the artificial and regenerated sand. It was demonstrated that the basic solution effectively removed the binder from the surface of the waste artificial sand. The Si-O-Si bond cleavage in the binder occurred due to the high concentration of hydroxide ions in the basic solution. The strength of the regenerated sand treated with the basic solution and that of the artificial sand were similar. Moreover, the strengths of regenerated sands treated with the acidic or neutral solutions were lower than that of artificial sand due to the presence of residual binder. These results indicated that the basic solution was suitable for removing the sodium silicate binder. The quality of the regenerated sand was similar to that of artificial sand.

Keywords: Artificial sand, Sodium silicate binder, Waste foundry sand, Wet regeneration

1. Introduction

Natural sand is mainly used for making molds in sand casting, and waste foundry sand (WFS) from the casting process is discarded on landfills [1, 2] or reused as concrete and ceramic materials [3–7]. WFS disposal on landfills may cause not only an increase in operational costs but also environmental pollution. WFS contains toxic heavy metals and particulate matters that can leach into the ground and result in soil and ground water pollution [1, 2]. Moreover, this leads to additional disposal costs due to the need for remediation and complying regulations. In addition, landfill sites are becoming insufficient due to increasing WFS volumes, so they are not currently an effective disposal method. The foundry industry must comply with environmental protection and resource conservation regulations [8]. Therefore, it is necessary to develop an appropriate regeneration method to reuse WFS in the casting process.

Several studies have been conducted using artificial sand to overcome the limitations of natural sand [9–11]. The use of artificial sand for making molds is advantageous for several reasons. First, high-durability artificial sand cannot be easily crushed and produces less dust [9]. Second, spherical artificial sand requires less amounts of binder for a specific mold volume, which decreases the amount of gas generated during casting and results in better casting quality [10]. Despite these advantages, the high cost of artificial sand mainly limits its applications [11]. Therefore, efficient regeneration methods should be developed to compensate that limitation and to reduce its disposal cost after casting.

Typically, there are three different methods to regenerate WFS: dry, thermal, and wet regeneration [12]. Dry regeneration removes the binder on the sand surface using friction, making this process significantly simple compared to other methods. However, the co-produced dust during dry regeneration negatively affects the working environment and the quality of the regenerated sand [12, 13]. Thermal regeneration removes the binder using a high-temperature treatment (> 600°C), which is energy costly and causes the emission of harmful gases [13]. In contrast, wet regeneration uses water and chemical solutions for binder removal, and produces...
better-quality reused sand than other methods [12, 14].

The waste sand used herein was prepared from water glass binder on artificial sand, and it was recycled through wet regeneration. The purpose of this study was to determine the effect of the chemical solutions on binder removal and to evaluate the reusability of the regenerated sand. Typically, representative acidic and basic solutions are used for regeneration. The use of oxalic acid [15] and hydrochloric acid [16] as the acidic solutions has been reported, while sodium hydroxide [17] and potassium hydroxide have been used as basic solutions. Because the silica component on the sand surface cannot be removed with acid solutions [18], most studies used basic solutions. Herein, potassium hydroxide, being one constituent of the binder, was chosen among the base solutions for the experiments. Hydrochloric acid as the acidic solution based on our prior research [19], and deionized water was utilized as a pH reference material. The effect of the solution type on binder removal was confirmed by comparing the resulting characteristics in the regenerated sand, based on chemical composition, surface and particle size analysis, and flexural strength.

2. Experimental

2.1. Preparation of the Waste Artificial Sand

The artificial sand (Simat International Limited Company, Hong Kong) and 2 wt% of sodium silicate-based binder [20] were mixed at 220 rpm for 120 s (device from Youngjin Machinery Co., Ltd, Korea). The mixed sand was formed into a core (device from Youngjin Machinery Co., Ltd, Korea) at a pressure of 4 bar. Hot air (150°C) was blown into the core for 30 s to harden the binder. The produced 22.4 × 22.4 × 170 mm (W × L × H) core was used as the waste artificial sand. The chemical compositions of laboratory-made and waste site-generated artificial sand were almost equivalent based on previous reports, making valid to carry out the experiments using laboratory-made artificial sand.

2.2. Regeneration process

Three different chemical solutions (based on hydrochloric acid, deionized water, and potassium hydroxide) were used separately to act as the chemical solution (Fig. 1) for wet regeneration of the waste artificial sand, and the effects of these solutions were compared. The concentrations of the acidic and basic solutions were 0.2 M hydrochloric acid (HCl; purity: 35%, grade: EP, Daejung Chemical & Metals Co., Ltd, Korea) and 0.2 M potassium hydroxide (KOH; purity 85%, grade: EP, Daejung Chemical & Metals Co., Ltd, Korea), respectively. Deionized (DI) water was used as the neutral solution for wet regeneration.

Fig. 1 shows the schematic of the regeneration process. The same process was repeated three times using DI water for neutralizing the pH of the sand. The regenerated sand was dried at 105°C for 2 h using an oven.

The three regenerated sand samples were denoted according to the regeneration media used as sand treated with the basic solution (B1), sand treated with the neutral solution (N1), and sand treated with the acid solution (A1), respectively. N1 and A1 were re-treated with the KOH solution according to the regeneration process and subsequently labelled as N2 and A2.

2.3. Analysis Methods

X-ray fluorescence spectroscopy (XRF, Zetium, PANalytical, Netherlands) was used to compare the elemental compositions of the regenerated sand and artificial sand. Zeta potential measurements (ZetaPALS, Brookhaven, USA) were used to confirm the surface charge on the sand. 0.5 g of each sand sample was dispersed in 1 mL of DI water. The data for all samples were analyzed from five replicates.

Field emission scanning electron microscopy (FE-SEM, SU 8020, Hitachi, Japan) and energy dispersive X-ray spectroscopy (EDS, X-Max50 011, Horiba, Japan) were used for surface analysis of the sand samples.

Particle size distributions were obtained by mechanical sieving as follows: dried sand (100 g) was passed through a series of sieves stacked up in increasing mesh order (850, 600, 425, 300, 212, 150, 105, 75, 53, and 0 μm). The sieves were shaken for 3 min using a sieve shaker (CO-211-6, Chunggye, Korea), and the sample weights in each sieve were recorded. The particle distributions...
Table 1. Chemical Compositions of Artificial Sand, Waste Artificial Sand, and Regenerated Sand

| Element   | Artificial sand (wt%) | Waste artificial sand (wt%) | Regenerated sand (wt%) | Re-treated sand (wt%) |
|-----------|-----------------------|-----------------------------|------------------------|-----------------------|
| Na2O      | 0.08                  | 1.90                        | 0.07                   | 0.06                  |
| Al2O3     | 67.50                 | 60.48                       | 67.44                  | 66.31                 |
| SiO2      | 18.02                 | 23.54                       | 17.96                  | 20.01                 |
| K2O       | 1.85                  | 2.08                        | 1.85                   | 1.87                  |
| TiO2      | 5.32                  | 5.08                        | 5.40                   | 5.14                  |
| Fe2O3     | 5.11                  | 4.63                        | 5.13                   | 4.55                  |
| others    | 2.12                  | 2.29                        | 2.15                   | 2.06                  |

The strength was tested using a core (prepared as the same as the waste artificial sand) as the specimen. The specimens were stored under different conditions. Under normal conditions, they were stored at a temperature of 20°C and relative humidity (RH) of 30% for 1 min and 60 min. Under humid conditions, they were stored at a temperature of 38 °C and RH of 65% for 120 min. The specimens for analysis under the humidity condition were stored at a temperature of 20°C and relative humidity (RH) of 30% for 1 min and 60 min. Under humid conditions, they were stored at a temperature of 38 °C and RH of 65% for 120 min.

The bridges were mainly formed by the Si-O-Si bonds, whereas sodium was not bonded to the structure or present on the silica or sand surface in its ionic state [21, 23]. Because sodium ions do not participate in binder bridges formation, the sodium content in the samples was easily removed through the regeneration process [24]. The Na2O contents for B1, N1 and A1 were 0.07%, 0.06%, and 0.06%, respectively. The Na2O contents for all the regenerated sand samples were lower than that of the waste artificial sand (1.90%) and similar to that of the artificial sand (0.08%). In addition, SiO2 contents for B1, N1 and A1 were 17.96%, 20.01%, and 21.27%, respectively. The SiO2 content (which was the binder's main component) was used to determine whether the binder was removed or not. Considering the SiO2 content of the artificial sand as 18.02%, it was confirmed that most binders were removed for B1. However, N1 and A1, having higher SiO2 content than that of the artificial sand, were considered to have remaining binders.

These results can be explained through the binder solubility dependence on the pH of the chemical solution. For the basic solution, Si-O-Si bonds cleavage in the binder occurred by chemical reaction with hydroxyl groups present in the basic solution (Eq. (3)) [25]. When using the acidic solution, silica was formed as SiO2 through the reaction with the acid, and then remained on the surface of the sand particles (Eq. (4)) [26].

\[
\text{Dehydration} = \text{Si} - \text{O}^- + \text{Na}^+ + \text{H}_2\text{O} \rightarrow \text{Si} - \text{OH} + \text{Na}^+ + \text{OH}^- 
\]

\[
\text{Condensation} = \text{Si} - \text{OH} + \text{OH}^- + \text{Si} \equiv \rightarrow \text{Si} - \text{O}^- + \text{Si} + \text{H}_2\text{O}
\]

\[
\text{Na}_2\text{SiO}_3 + 2\text{HCl} \rightarrow 2\text{NaCl} + \text{H}_2\text{O} + \text{SiO}_2
\]

3.2. Surface Analysis of Sand Particles

Fig. 2 shows the scanning electron microscopy (SEM) images of the artificial sand, waste artificial sand, and regenerated sand. The artificial sand particles (Fig. 2(a)) were spherical and exhibited smooth surfaces. As shown in Fig. 2(c), a smooth surface with
no impurities were observed for B1 and its morphology was similar to that of the artificial sand. In addition, the surface of N1 was partially covered with impurities (Fig. 2(d)), while most of the surface of A1 had several impurities (Fig. 2(e)). The surface impurities for N1 and A1 were similar to that of the waste artificial sand (Fig. 2(b)).

Fig. 3 shows the EDS results for the artificial sand and regenerated sand. The EDS was conducted to detect Al, Si, and Na compounds, because these are the main components of the sand and the binder. The EDS mapping images in Figs. 3(a) and (b) showed that Al and Si compounds were uniformly distributed on both the surfaces of the artificial sand particles and B1. Moreover, their elemental contents were similar. As shown in Figs. 3(c) and (d), the Si composition was densely concentrated on the impurities of the sand surface for N1 and A1. Hence, the impurities were the products of the residual binder, which was not removed by the neutral and acidic solutions. The spectra in Fig. 3 revealed that the Si component gradually decreased in the following order: B1, N1 and A1.

3.3. AFS and Particle Size Distributions

Typically, sand strength is affected by the particle size, size distribution, shape, and storage humidity as well as the amount of binder and the pH value on the sand surface [28–30]. Fig. 4 shows the particle size distributions of the artificial sand and the regenerated sand. All samples had narrow particle size distributions, in the 150–425 μm mesh range. The calculated AFS values for the artificial sand sample, B1, N1 and A1 were 46, 45, 45, and 44, respectively. It was assumed that this slight difference did not affect the strength directly.

3.4. Flexural Strength

The low core flexural strengths cannot be managed by operators and may be damaged during assembly, storage, and transport, and also may collapse the mold during casting [20]. Accordingly, the strength of the regenerated sand was tested to determine if it could be used as a substitute of artificial sand.

Fig. 5 shows the core strengths under different conditions. The flexural strengths of the artificial sand for 1 and 60 min were 218 and 325 N/cm², respectively. The initial strength for 1 min was high (218 N/cm²), which indicated that it could be easily managed by an operator without causing any damage. The core was exposed to the normal condition at 150 °C, and the binder was sufficiently hardened. The flexural strength after 60 min was higher than that obtained after 1 min. The flexural strengths of B1 after 1 and 60 min were 220 and 320 N/cm², respectively. The strength of B1 was quite similar to that of the artificial sand. However, the flexural strengths of N1 and A1 after 60 min were 198 and 121 N/cm², respectively. The strength of N1 and A1 was considerably lower than that of the artificial sand and B1. This can be explained because N1 and A1 had a large amount of residual binder, which hindered the combination of regenerated sand and new binder. When the core was set to the humidity condition, its strength gradually decreased. The flexural strength of the artificial sand after 60 min under the normal condition was 325 N/cm²; however, the flexural strength of the artificial sand after 120 min under the humidity condition was 220 N/cm². The flexural strengths of B1, N1 and A1 after 120 min were 194, 168, and 105 N/cm², respectively.

The core strength increased with increasing amounts of binder. However, because the characteristics of the residual binder in the
Fig. 3. SEM and EDS analysis results of samples: (a) artificial sand, (b) B1, (c) N1, and (d) A1.

Fig. 4. Particle size distributions of artificial sand and regenerated sand (B1, N1, and A1).

Fig. 5. Flexural strengths of artificial sand and regenerated sand.
waste artificial sand changed, it was considered as an impurity. This was confirmed comparing the DSC curves of the new and cured binders (Fig. 6). The endothermic peak of the new binder (Fig. 6(b)) at 123°C confirmed the dehydration condensation reaction between the silanol groups of sodium silicate [31]. The binder cured at 150°C (Fig. 6(a)) exhibited no such a peak. This indicated that the dehydration condensation reaction did not occur, even when the cured binder was reheated, i.e., no bridges were formed between the sand particles, similar to the new binder. Therefore, because the residual binder on the regenerated sand became an impurity, the core strength of the regenerated sand with residual binder was comparatively lower [30].

3.5. Re-treated N1 and A1

Large amounts of remaining binder were present on the surface of the regenerated N1 and A1 sands. B1 (obtained using potassium hydroxide as the cleaning solution) effectively removed the entire binder. Therefore, N1 and A1 were regenerated once more using potassium hydroxide as the cleaning solution to determine its effect.

Table 1 compares the chemical compositions of the artificial sand, N2 and A2 samples. The Na2O content in artificial sand, N2 and A2 was 0.08%, 0.06%, and 0.06%, respectively. The SiO2 contents for N2 and A2 were similar to that of the artificial sand. The Al2O3 content in N2 and A2 was slightly higher than that of the artificial sand.

Fig. S1 shows the SEM images of re-treated N1 and A1 samples. N1 and A1 exhibited several impurities on the surface (Fig. 2). In contrast, the surfaces of N2 and A2 (samples re-treated with the basic solution) were smooth and had no impurities.

Fig. 7 shows the flexural strengths of the re-treated N1 and A1 samples, which were compared with those of the artificial sand and B1. The flexural strengths of N2 and A2 after 1 min were 248 and 272 N/cm², respectively, and the flexural strengths of N2 and A2 after 60 min were 337 and 374 N/cm², respectively. The flexural strengths of N1 and A1 with remaining binder on the surface were lower than that of the artificial sand, but the flexural strengths of N2 and A2 were higher than that of the artificial sand.

4. Conclusions

This study investigated the effect of binder removal and the reusability of regenerated sand according to the chemical solution used in the wet regeneration process. Comparing the characteristics of the artificial sand and the regenerated sand, using the basic solution was the most effective, among the three types of solutions, to remove the residual binder from waste artificial sand. The hydroxyl groups in the basic solution caused Si-O-Si cleavage that formerly bridged the sand particles; this was confirmed by XRF and zeta potential analysis. When the strengths of the artificial sand and the regenerated sand were compared, it was observed that the strength of regenerated sand treated with the basic solution (B1) was similar to that of the artificial sand. Moreover, the regenerated sand treated with neutral and acidic solutions (N1 and A1) had lower strength than the artificial sand. When N1 and A1 were re-treated with the basic solution to remove residual binder, the strengths of the regenerated sand (N2 and A2) increased. These results indicated that the residual binder on the surface of regenerated sand decreased the strength of regenerated sand, because it hindered the formation of binder bridges due to complete curing reaction of the silanol groups of the residual binder. This was demonstrated through the DSC curve of the binder.

Therefore, the potassium hydroxide solution was most suitable in the wet regeneration process of waste artificial sand using inorganic binders.

Several studies were conducted to compare the results of dry and wet regeneration of waste sand by using inorganic binders [14, 17]. The characteristics of regenerated sand using wet regeneration compared to dry regeneration were similar to those of the raw sand and in this study, and the properties of regenerated sand using basic solution were similar to those of the raw sand.
However, wet regeneration can lead to problems related to wastewater treatment, although studies have been reported regarding biological treatment [32]. Further research on the treatment of the wastewater generated from the experiments described in this study will be conducted.

Acknowledgements

This study was supported by the Technology Innovation Program (200000126 and 10067386) funded by the Ministry of Trade, Industry & Energy (MOTIE, Korea) and the Korea Institute of Industrial Technology (KITECH) through the Research and Development (KITECH EO-20-0140) grant.

Author Contributions

E.Y.K. (M.D.) conducted all the experiments and wrote the manuscript with input from all authors. K.H.K. (M.D., researcher) planned the experiments and revised the manuscript. J.H.B. (Ph.D., Senior researcher) elaborated the discussion of the experiments’ results. I.S.H. (Professor) revised the manuscript. M.S.L. (Ph.D., Principal researcher) wrote and revised the manuscript.

References

1. Siddiquea R, Kaur G, Rajor A. Waste foundry sand and its leachate characteristics. Resour. Conserv. Recycl. 2010;54:1027-1036.
2. Bhardwaj B, Kumar P. Waste foundry sand in concrete: A review. Constr. Build. Mater. 2017;156:661-674.
3. Guney Y, Sarı YD, Tunçan A, Donmez S. Re-usage of waste foundry sand in high-strength concrete. Waste Manag. 2010;30:1705-1713.
4. Lin DF, Luo HL, Lin JD, Zhuang ML. Characterizations of temperature effects on sintered ceramics manufactured with waste foundry sand and clay. J. Mater. Cycles. Waste. Manag. 2018;20:127-136.
5. Venkatesan M, Zabi Q, Shah IH, Park HS. Optimum utilization of waste foundry sand and fly ash for geopolymer concrete synthesis using D-optimal mixture design of experiments. Resour. Conserv. Recycl. 2019;148:114-123.
6. Mavroulidou M, Lawrence D. Can waste foundry sand fully replace structural concrete sand? J. Mater. Cycles. Waste. Manag. 2019;21:594-605.
7. Basar HM, Aksoy ND. Pre-treatment of Waste Foundry Sand Via Solidification/Stabilization. CLEAN - Soil, Air, Water. 2013;41:94-101.
8. Li Y lei, Wu G hua, Liu W cai, Chen W cai, Zhang L, Wang Y xin. Effect of reclaimed sand additions on mechanical properties and fracture behavior of furan no-bake resin sand. China Foundry. 2017;14:128-137.
9. Mizuki T, Kanno T. Establishment of Casting Manufacturing Technology by Introducing an Artificial Sand Mold with Furan Resin and Realizing a Clean Foundry. Int. J. Met. 2018;12:772-778.
10. Wen S, Shen Q, Wei Q, et al. Material optimization and post-processing of sand moulds manufactured by the selective laser sintering of binder-coated Al2O3 sands. J. Mater. Process. Technol. 2015;225:93-102.
11. Kosuge K, Sunaga M, Goda R, Onodera H, Okane T. Cure and collapse mechanism of inorganic mold using spherical artificial sand and water glass binder. Mater. Trans. 2018;59:1784-1790.
12. Ying-Min L, Tian-Shu W, Wei-Hua L. Research on regeneration methods of animal glue waste sand for foundry. R. Soc. Open. Sci. 2018;5:172-270.
13. Park CL, Kim BG, Yu Y. The regeneration of waste foundry sand and residue stabilization using coal refuse. J. Hazard. Mater. 2012;203-204:176-182.
14. Wang L, Jiang W, Liu F, Fan Z. Investigation of parameters and mechanism of ultrasound-assisted wet reclaimation of waste sodium silicate sands. Int. J. Cast. Met. Res. 2018;31:169-176.
15. Hefei City Tenver Precision Casting Co., Ltd., Regeneration treatment method of clay wet type used sand. 105903894 (2016.08.31), China Patent
16. Jang JE, Jung ES, Manufacturing method of recycling sand using waste foundry sand. 10-0760646, Korea Patent
17. Fan Z, Huang N, Wang H, Dong X. Dry recycling and wet reclaimation of used sodium silicate sand. China. Foundry. 2005;2:38-43.
18. Yang X, Roonasi P, Holmgren A. A study of sodium silicate in aqueous solution and sorbed by synthetic magnetite using in situ ATR-FTIR spectroscopy. J. Colloid. Interface. Sci. 2008;328:41-47.
19. Ko EY, Kim KH, Baek JH, Hwang IS, Lee MS. Characteristics of the chemical regeneration for used artificial sand according to the pH value of chemical solution (in Korean). Korea Foundry Society Spring meeting. 2019.
20. Bae MA, Kim MH, Park JW, Lee MS. Effect of additives on the sand burning of inorganic binder in Al-Si7Mg alloy casting. J. Korean. Inst. Met. Mater. 2018;56:327-333.
21. Lucas S, Tognonvi MT, Gelet JL, Soro J, Rossignol S. Interactions between silica sand and sodium silicate solution during consolidation process. J. Non. Cryst. Solids. 2011;357:1310-1318.
22. Fan ZT, Huang NY, Dong XP. In house reuse and reclamation of used foundry sands with sodium silicate binder. Int. J. Cast. Met. Res. 2004;17:51-56.
23. Lim HM, Lee J, Jeong JH, Oh S-G, Lee S-H. Comparative Study of Various Preparation Methods of Colloidal Silica. Engineering 2010;02:998-1005.
24. Adelman JG, Eloutakit S, Demopoulos GP. Investigation of sodium silicate-derived gels as encapsulants for hazardous materials - The case of scorodite. J. Hazard. Mater. 2015;292:108-117.
25. Kouassi SS, Tognonvi MT, Soro J, Rossignol S. Consolidation mechanism of materials obtained from sodium silicate solution and silica-based aggregates. J. Non. Cryst. Solids. 2011;357:3013-3021.
26. Yang X, Roonasi P, Holmgren A. A study of sodium silicate in aqueous solution and sorbed by synthetic magnetite using in situ ATR-FTIR spectroscopy. J. Colloid. Interface. Sci. 2008;328:41-47.
27. Hartmut P, Bernhard ES. Inorganic binders : for mould and core production in the foundry. Berlin : Fachverlag Schiele & Schön; 2014.
28. Holtzer M, Górny M, Dańko R. Microstructure and Properties of Ductile Iron and Compacted Graphite Iron Castings. SpringerBriefs in Materials; 2015. p. 27-43.

29. Parappagoudar MB, Pratihar DK, Datta GL. Modeling and Analysis of Sodium Silicate-Bonded Moulding Sand System Using Design of Experiments and Response Surface Methodology. J. Manuf. Sci. Prod. 2012;11:1-14.

30. Khan MM, Singh M, Jadhav GN, Mahajani SM, Mandre S. Characterization of Waste and Reclaimed Green Sand Used in Foundry Processing. Silicon. 2020;12:677-691.

31. Zhou X, Yang J, Su D, Qu G. The high-temperature resistant mechanism of α-starch composite binder for foundry. J. Mater. Process. Technol. 2009;209:5394-5398.

32. Wang H, Fan Z, Yu S, Liu F, Li X. Wet reclamation of sodium silicate used sand and biological treatment of its wastewater by Nitzchia palea. China. Foundry. 2012;9:34-38.