

Research Article

Mechanism of Degradation of the Properties of Recycled Plaster Mixed Aluminate Cement

Zhixin Li , Kaidong Xu, Jina Wang, Jianwu Zhang, Xianwei Ma , and Jishou Niu

School of Material and Chemical Engineering, Henan University of Urban Construction, Pingdingshan 467036, China

Correspondence should be addressed to Zhixin Li; li.zhixin1989@163.com

Received 24 October 2019; Revised 15 January 2020; Accepted 31 January 2020; Published 27 February 2020

Academic Editor: Marco Rossi

Copyright © 2020 Zhixin Li et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

This manuscript investigates the degradation of the properties of recycled plaster-mixed aluminate cement (RAP) and analyzes its degradation mechanism by DSC/TG and SEM. The results showed that the setting time of RAP was shortened due to the fast formation of recycled ettringite (AFt) and its strength was decreased relative to the pure recycled plaster (RP) in the absence of aluminate cement. Different from the properties of RP and RAP, the hydration of commercial plaster was slowed down by the addition of aluminate cement for its low hydration rate, and its strength was increased with respect to the pure commercial plaster (CP) without aluminate cement. Therefore, the properties of RP and RAP could be seen to decrease in relation to CP and commercial plaster mixed aluminate cement (CAP). The SEM and DSC/TG analyses confirm the presence of cluster and fine crystals and noncementing AH_3 in RAP, which demonstrates its degradation of properties.

1. Introduction

At present, commercial gypsum is one of the most commonly used materials in ceramic mold [1, 2] and general construction work [3, 4] because of its characteristics of light weight, stable volume, heat insulation, sound insulation, good fire resistance, and humidity adjustment [5]. Lots of waste gypsum is therefore usually produced after use. According to Suárez et al., the amount of waste gypsum produced each year is as high as 80 million tons, of which about 15 million tons are landfilled [6]. It not only takes up a large amount of arable land and pollutes the environment, but also produces hydrogen sulfide gas when it is mixed with organic matter, which is harmful to humans [7, 8]. Therefore, reutilization of waste gypsum is necessary to protect the environment. However, the current use of waste gypsum is mainly in cement as a retarder [9], as a roadbed material [10, 11], and as a ceramic additive [12]; this utilization of waste gypsum is rare.

It is well known that hydration of hemihydrate gypsum can produce dihydrate gypsum, and the dihydrate gypsum can be calcined to form hemihydrate gypsum, which is the basis of gypsum recycling [13]. Thus, gypsum recycling turns

into feasible by only calcining the waste gypsum to produce a fresh hemihydrate gypsum. Based on this, recycled plaster can be produced by a reversible transformation of dihydrate gypsum into hemihydrate gypsum in an electric furnace at the temperature of 180°C [14]. Ren [15] presented the water/plaster ratio, hydration, and strength of pure recycled plaster without any impurities, and the initial setting time and final setting time were 29 min and 40 min, respectively. The flexural strength and compressive strength were 1.66 MPa and 2.94 MPa, respectively, and the extended hydration and degraded strength of pure recycled plaster could be found in relation to commercial plaster. Zhu et al. [16] analyzed the degradation mechanisms of pure recycled plaster mechanical performance and attributed it to the increased porosity and larger pore size of recycled plaster.

In addition, some scholars have reported the properties of recycled plaster calcined by the waste of hydrated commercial plaster from construction works. Bardella and Camarini [17] described the physical properties and mechanical properties of recycled plaster. Camarini et al. [18] showed the thermal analysis results of recycled plaster. Pinheiro and Camarini [19] reported the chemical results of recycled plaster in different number of cycles. Geraldo

et al. [20] showed that the main phases of recycled plaster did not change with the recycling process. The results showed that the physical, mechanical, thermal, and chemical properties and phases were similar to those achieved from commercial plaster. Indeed, other materials, such as cement, dolomite, quicklime, are always added to commercial plaster to improve their performances. Most of the setting time, water/plaster ratio and strength of recycled plaster may depend on their existence. Therefore, mastering the properties of recycled plaster mixed with other materials is very important and urgent for its use. Ren, Bardella, Camarini, Pinheiro, and Geraldo did not analyze the effect of other existing materials on the properties of recycled plaster.

Camarini et al. investigated the influence of retarder on the workability, setting time, and strength of recycled plaster [21] and found that the citric acid decreased the water/plaster ratio, prolonged the setting time, and reduced the compressive strength of recycled plaster. Heloisa et al. [22] studied the effect of impurities, like dolomite, on the properties of recycled plaster, and concluded that apparent bulk density and bulk density were both decreased, while the setting time, water content to normal consistency, and compressive strength of recycled plaster were all increased. Li et al. [23, 24] studied the degradation of the properties of recycled plaster mixed ordinary Portland cement and quicklime. They attributed the fast hydration to the quick recycling of AFt and increasing of Ca^{2+} in the saturated solution in regard to the Ca^{2+} and SO_4^{2-} ions, respectively. However, little research has been done on the degradation of the properties and action mechanism of recycled plaster mixed aluminate cement (RAP), which severely hampers the use of waste gypsum.

In this paper, the water/plaster ratio, setting time, and strength of recycled plaster mixed aluminate cement (RAP) have been investigated, and its degradation mechanism is also analyzed by using scanning electron microscope (SEM) and Differential Scanning Calorimeter/Thermogravimetric analysis (DSC/TG). The authors hope that this research will help the disposal of waste gypsum.

2. Materials and Methods

2.1. Materials. The original gypsum (OG) comes from Yuhuan gypsum factory in Yingcheng, Hubei, China. The aluminate cement is produced by Guizhou Xifeng Rendu Building Materials Co., Ltd., CA-50. Their chemical compositions were measured by X-ray fluorescence (XRF). The chemical composition analysis of OG is displayed in Table 1, and its main chemical components are SO_3 and CaO. Its main mineralogical composition is dihydrate ($\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$). The specific surface area and chemical composition analysis are displayed in Table 2. The content of CaO and Al_2O_3 are 34% and 53.85%, respectively, and the main mineralogical composition is $\text{CaO} \cdot \text{Al}_2\text{O}_3$ and $\text{CaO} \cdot 2\text{Al}_2\text{O}_3$.

2.2. Recycling Process. The recycling process is shown in Figure 1, and it contains the processes of crushing,

TABLE 1: Chemical composition analysis of original gypsum.

Original gypsum	SO_3	CaO	SiO_2	Al_2O_3	Fe_2O_3	K_2O	SrO
Chemical composition (%)	47.57	38.73	1.15	0.42	0.13	0.07	0.04

TABLE 2: Specific surface area and chemical composition analysis of aluminate cement.

Specific surface area (m^2/kg)	Loss	CaO	Al_2O_3	SiO_2	MgO	SO_3	Fe_2O_3	TiO ₂	
CA-50	335	0.25	34.00	53.85	5.85	0.53	0.06	1.89	2.75

calcination, and hydration. First, OG was crushed with a jaw crusher, and grounded with a laboratory steel ball mill for 2 minutes. Second, the powdered OG was calcined at the temperature of 180°C for 3 hours in an electric oven. Commercial plaster (CP) was obtained and added into aluminate cement of 10%. Third, the mixture was mixed with water to obtain the hardened commercial plaster-mixed aluminate cement (CAP), and the water to plaster ratio was 0.60. Finally, the hardened CAP was dried till mass stabilization at 45°C , and the recycled plaster-mixed aluminate cement (RAP) and hardened RAP were achieved as the same process above. Specially, the water to plaster ratio of RAP was 0.86 when the same fluidity was reached. When the aluminate cement was not added in the CP, recycled plaster (RP), hardened CP, and hardened RP were obtained following the same steps as before, and the water to plaster ratio of CP and RP were 0.63 and 0.85, respectively. The properties tests were conducted based on the Chinese standard GB 9776-2008 [25]. The test methods for w/p (consistency), setting time, and strength have been described in our previous research [24]. The sample dimensions for mechanical properties determination was $40\text{ mm} \times 40\text{ mm} \times 160\text{ mm}$. And the compressive strength and flexural strength were measured after curing for 14 days. For flexural strength, 3 independent samples were determined and the average value was calculated. While for the compressive strength values, the average of six independent samples were taken.

2.3. Experimental Methods. Partial replacement of commercial plaster by aluminate cement was adopted to investigate degradation law and analyze action mechanism of recycled plaster-mixed aluminate cement, and the dosage of aluminate cement was 10 wt.%. The experiment was carried out by adding aluminate cement to commercial plaster and then crushing after hydration and calcining into recycled plaster mixed aluminate cement.

A program was created during the test to help understand the progress of the processes, which is shown in Figure 2. In brief, the water to plaster ratio of powdered CP, CAP, RP, and RAP were tested first, and then the setting time of CP, CAP, RP, and RAP were measured based on their water to plaster ratios. Afterwards, the flexural strength of

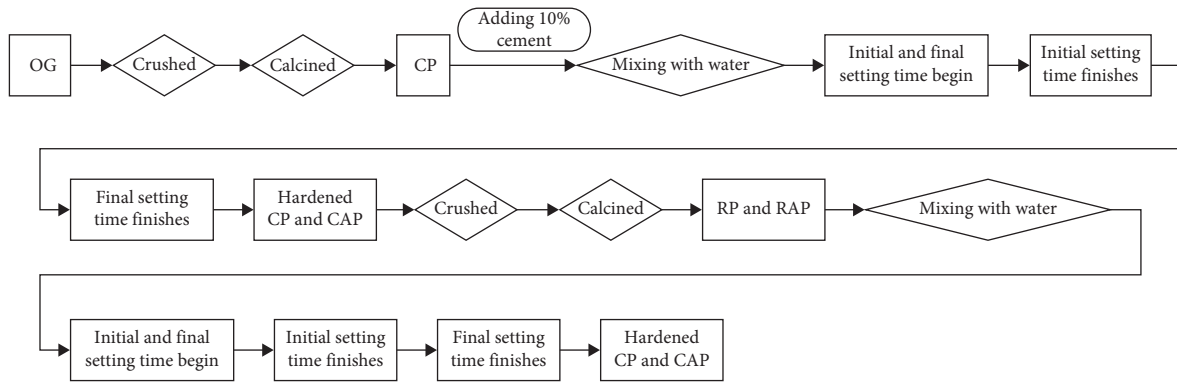


FIGURE 1: The recycling process.

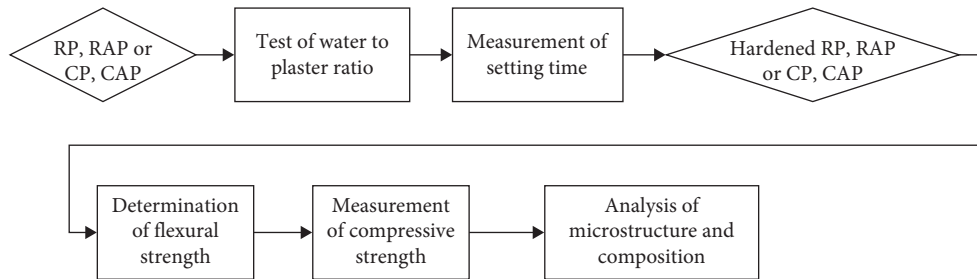


FIGURE 2: The testing program.

hardened CP, CAP, RP, and RAP was determined before the measurement of the compressive strength. Finally, the microstructure and composition of hardened CP, hardened CAP, hardened RP, and hardened RAP were analyzed by SEM and DSC/TG.

2.4. Materials Characterization. The compositions of hardened CP, hardened CAP, hardened RP, and hardened RAP were determined by simultaneous DSC-TG device (SMP/PF7548/MET/400 W) and X-ray diffraction analysis (XRD), and the DSC-TG was from room temperature up to 600°C, using a heat rate of 10°C/min. The morphological investigations of hardened CP, hardened CAP, hardened RP, and hardened RAP were made by SEM (TESCAN VEGA III LMH).

3. Results and Discussions

3.1. Water/Plaster Ratio and Setting Time of the Powdered CP, CAP, RP, and RAP. Figure 3 shows the water to plaster ratios of the composites. As can be seen, the water to plaster ratio of CAP does not reach the pure CP and is 5% lower than that of CP. This result is expected due to the low water requirement of aluminate cement. This is consistent with the research of Zhao et al. [26]. On the contrary, the water-plaster ratio of RAP and pure RP is 35% and 43% higher with respect to that of CAP and CP, and the water to plaster ratio of RAP exceeds the pure RP.

The setting time of plaster and plaster mixed aluminate cement is presented in Figure 4. As can be seen, the initial setting time of RP is 33 minutes and its final setting time is 13

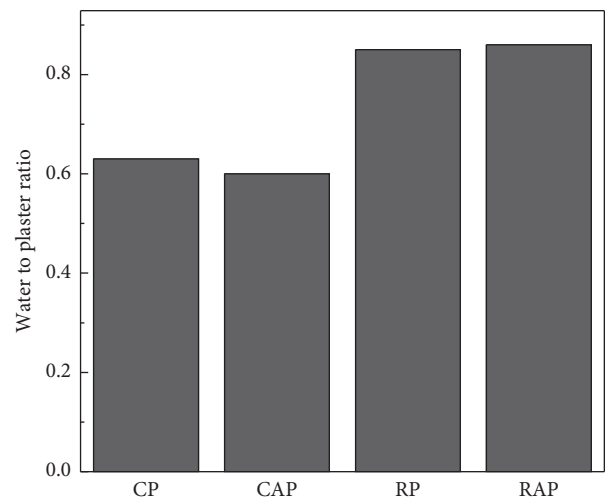


FIGURE 3: Results of water/plaster ratio.

minutes longer than the initial setting time, a result that agrees with the findings of Ren [15]. However, aluminate cement significantly shortens the setting time of RAP. The initial and final setting time are reduced to 5.5 minutes and 10.5 minutes, respectively, which are 83% and 77% lower than that for the initial setting time and final setting time of RP. This is due to the rapid recycling of ettringite (AFt) (Figure 5). Unlike the recycled products, the initial setting time of CAP is increased from 8.5 minutes to 13.5 minutes in comparison to CP and is 59% longer than that of CP, and the final setting time is raised from 13.5 minutes to 17.5 minutes, which is 30% longer than that of CP. The opposite trend in setting time can be found between RAP and CAP.

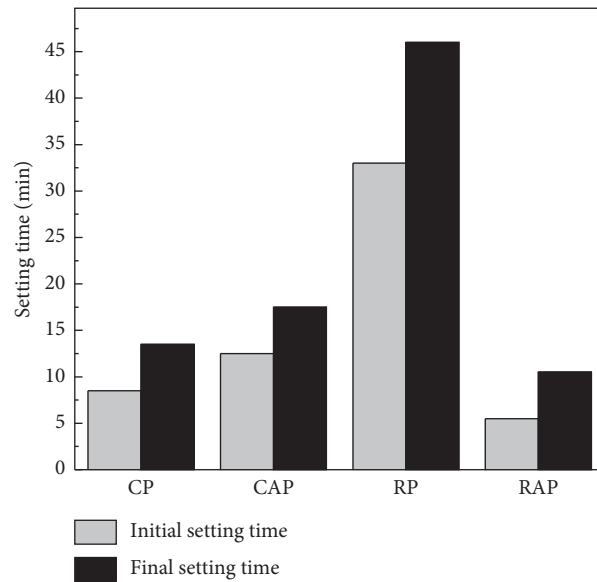


FIGURE 4: Results of setting time.

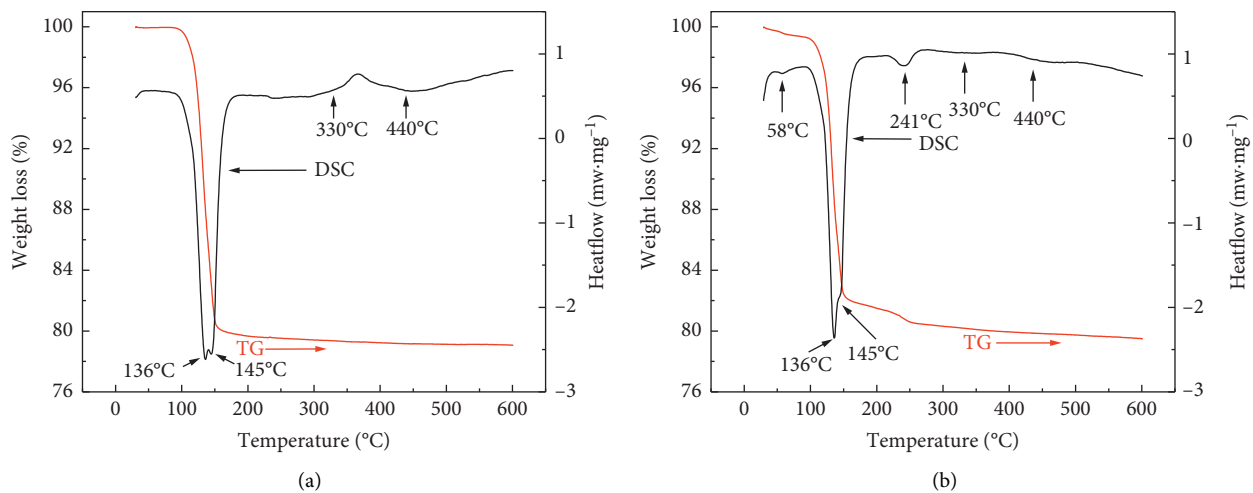


FIGURE 5: DSC/TG curves of hardened RP (a) and hardened RAP (b)

3.2. Strength of Hardened CP, Hardened CAP, Hardened RP, and Hardened RAP. Figure 6 presents the strength of hardened CP, hardened CAP, hardened RP, and hardened RAP, from which we can see that hardened RP and hardened RAP exhibit low strength, while hardened CP and hardened CAP display steadily high strength. The addition of aluminate cement increases the flexural strength (FS) of the hardened CAP from 5.29 MPa to 5.89 MPa, and the compressive strength (CS) increases from 10.08 MPa to 13.54 MPa. The rate of increase in strength can be seen in Figure 7(a). In contrast, the flexural strength and compressive strength of the hardened RAP are reduced from 2.70 MPa to 2.40 MPa and from 4.80 MPa to 4.59 MPa, respectively. The strength reduction rate is shown in Figure 7(b). Simultaneously, compared to hardened CP and hardened CAP, the strength of hardened RP and hardened RAP is very low. In the absence of aluminate cement, the

strength reduction rate of hardened RP is about 50% with respect to the pure hardened CP, the number rises swiftly in the presence of aluminate cement (Figure 8), as the strength decrease rate of hardened RAP is as high as 65%. Therefore, it can be concluded that the opposite strength development trend can be seen between hardened RP, hardened RAP and hardened CP, hardened CAP. Moreover, the rate of strength reduction of hardened RAP is much higher than that of hardened RP without aluminate cement.

3.3. SEM Images. Figure 9 shows that aluminate cement significantly changes the microstructure of hardened plaster. The hardened CP has a needle-like crystal structure. After the addition of aluminate cement, the crystals of the hardened CAP have changed, and a large number of gelatinous crystals have appeared. The hydrated products of

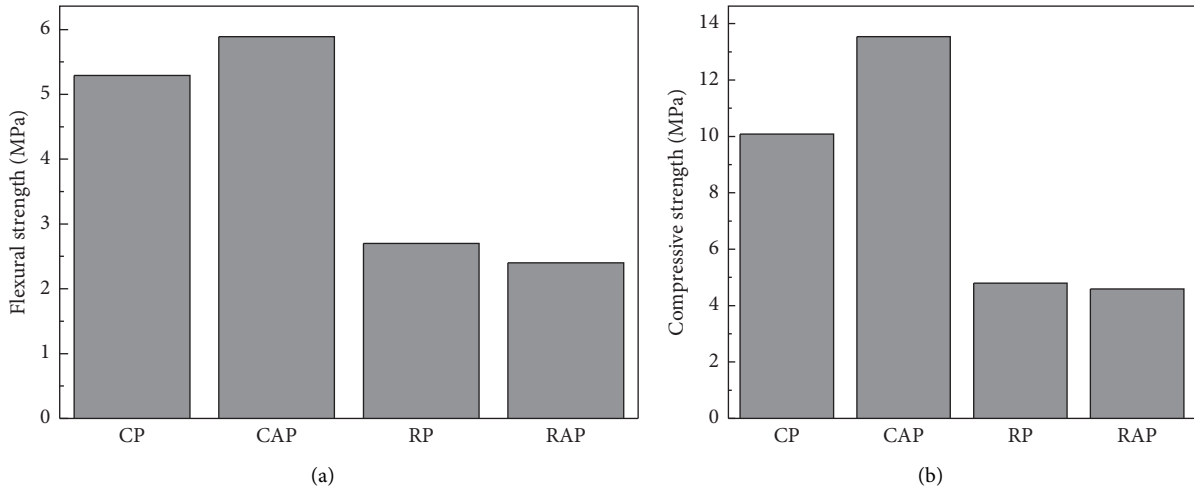


FIGURE 6: Results of strength.

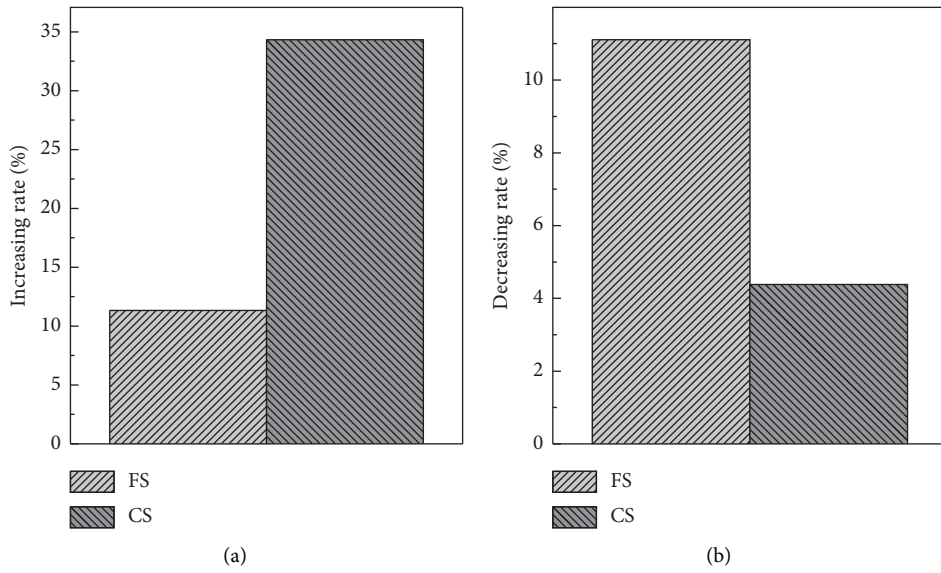


FIGURE 7: The strength changing rate of commercial plaster and recycled plaster aluminate cement mixes. (a) The increasing rate of CAP. (b) The decreasing rate of RAP.

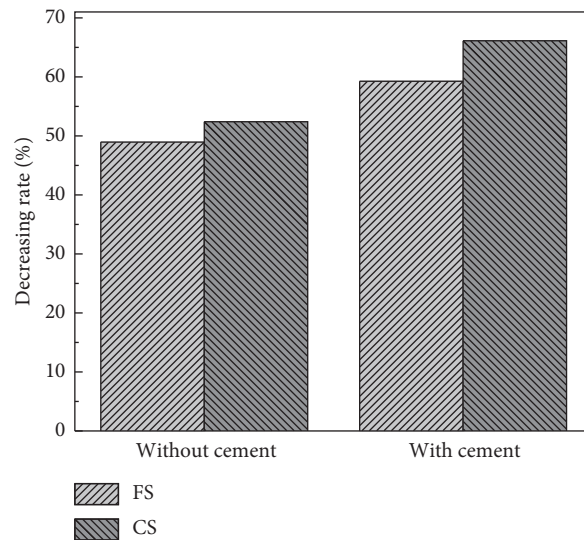


FIGURE 8: The strength decreasing rate of recycled plaster before and after adding aluminate cement.

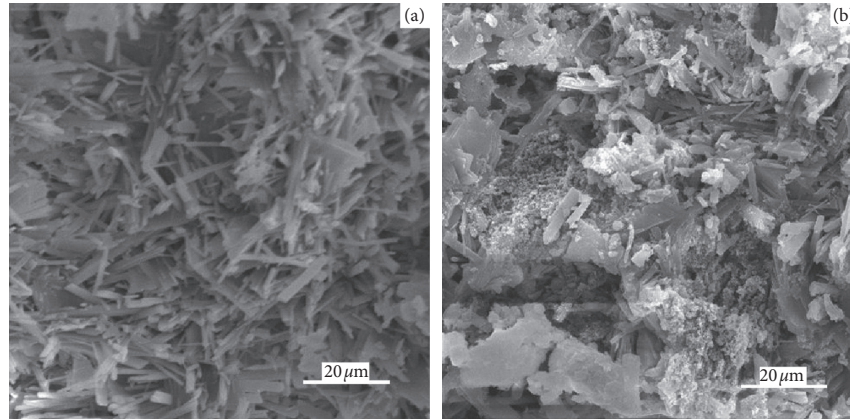


FIGURE 9: Microstructure of hardened CP (a) and hardened CAP (b).

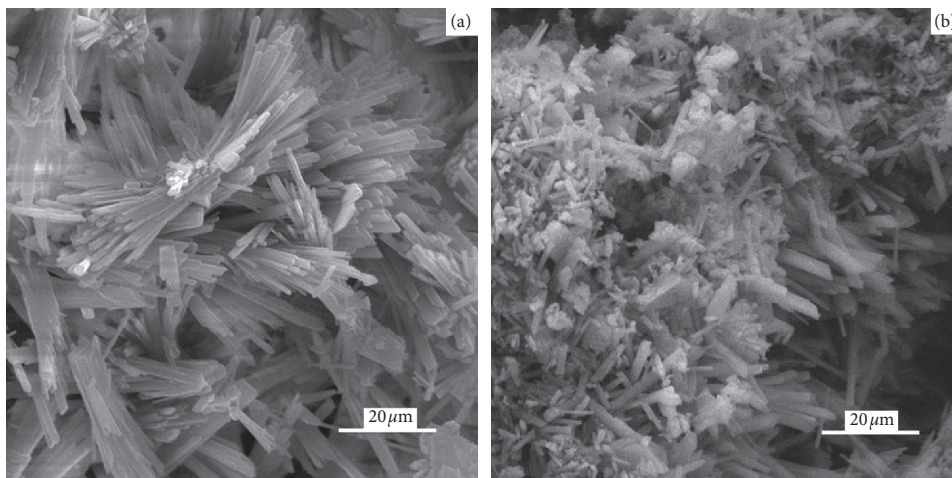


FIGURE 10: Microstructure of hardened RP (a) and hardened RAP (b).

aluminate cement, AFt, and AH_3 , are dispersed in the crystals, reducing the voids of the hardened CAP and enhancing its stability. Therefore, the strength of the hardened CAP is improved.

The SEM images of hardened RP and RAP are exhibited in Figure 10. It is evident that the morphology and habits of the crystals have changed. The crystals of the hardened RP are obviously coarsened and arranged into parallel beams; but it is still interconnected with each other. As for the hardened RAP, the crystals contain not only cluster crystals similar to those of hardened RP, but also a large number of fine crystals, which are dispersed and not filled into the plaster voids, thereby weakening their overlapping. Furthermore, the presence of AH_3 of noncementing reduces the strength of hardened RAP as well. Thus, the strength of the hardened RAP is decreased.

3.4. DSC/TG. Figure 11 shows the DSC/TG curves for hardened CP and hardened CAP. Figure 11(a) obviously reveals that endotherms at 139°C and 152°C occur as a result of the dehydration of dihydrate gypsum to hemihydrate gypsum and hemihydrate gypsum to anhydrite III, and the endotherms near 340°C and 450°C are due to the

transformation of anhydrite III to anhydrite II and lattice rearrangement, illustrating the existence of dihydrate gypsum, which agrees well with Tong [27] and Zhao et al. [28]. And it is also confirmed by XRD (Figure 11(c)). However, other peaks are emerged in the hardened CAP (Figure 11(b)) as well. For example, the occurrence of an endothermic peak at 80°C is due to the decomposition of AFt into monosulfate (AFm). The small endothermic peak at 248°C is caused by the endothermic peak of AH_3 , which is consistent with Zhao et al. [26]. Moreover, the endotherms at 139°C and 152°C are reduced relative to the hardened CP. The above indicates that AFt and AH_3 are present and the quantity of gypsum is low in the hardened CAP. It is fully described that the strength of the hardened CAP is obtained through the hydration products of AFt and AH_3 , which agrees with the previous finding [26].

In Figure 5(a), there are two consecutive endothermic peaks near 136°C and 145°C due to the dehydration of dihydrate gypsum to hemihydrate gypsum and hemihydrate gypsum to anhydrite III, and the endotherms around 330°C and 440°C are due to the transformation of anhydrite III to anhydrite II and lattice rearrangement, which is similar to the endothermic peaks of hardened CP, consistent with Tong [27] and Zhao et al. [28], and the endotherm near 330°C is

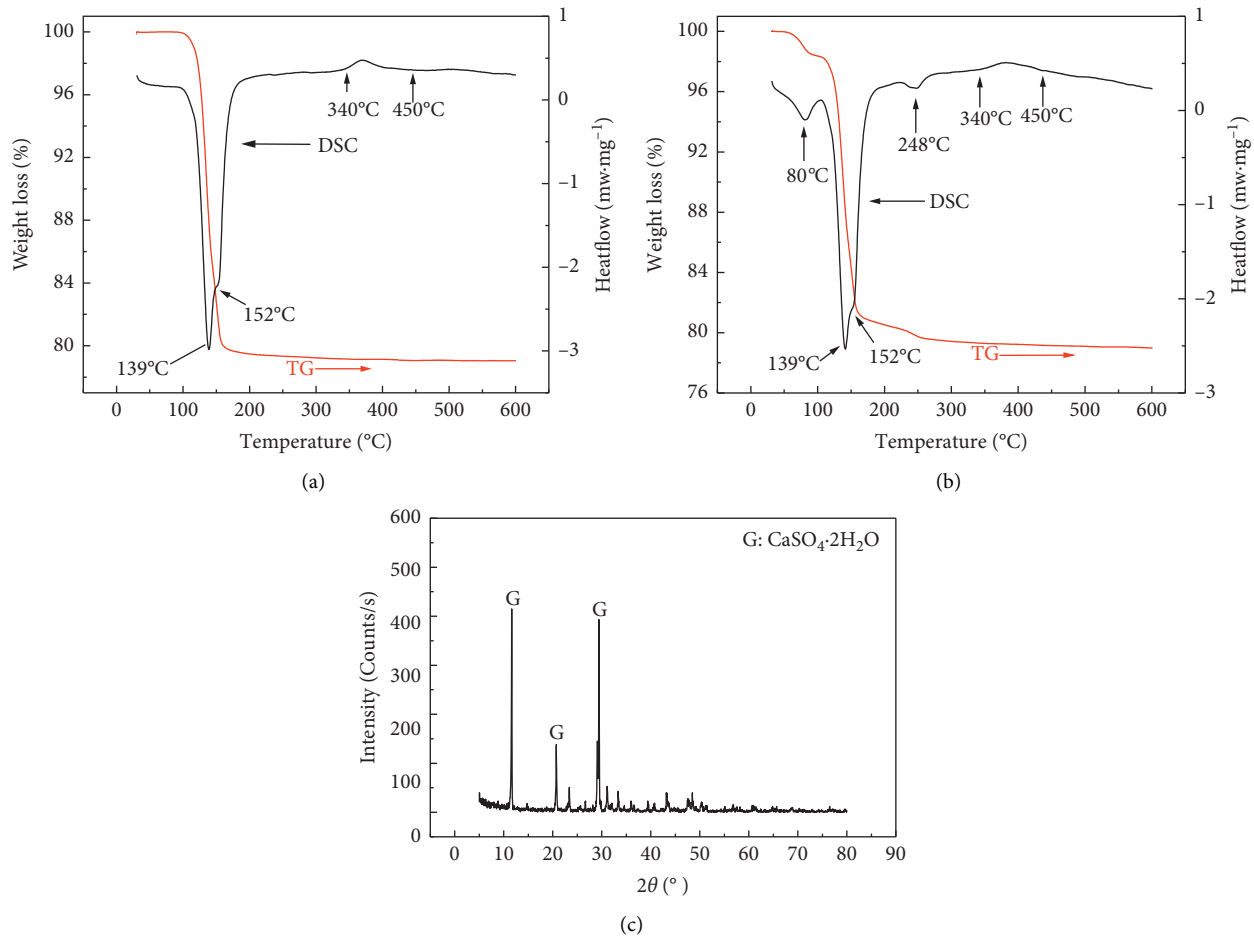


FIGURE 11: DSC/TG curves of hardened CP (a) and hardened CAP (b) and XRD patterns of hardened CP (c).

also confirmed by Geraldo et al. [20]. To our surprise, the DSC/TG curves of the hardened RAP have changed significantly, and the endothermic peaks of the hardened RAP at 136°C and 145°C are reduced due to the addition of aluminate cement (Figure 5(b)). This is owing to the low quantity of gypsum and presence of undecomposed AH_3 and recycled AFt, which is identical to Li et al. [23]. It manifests that the hydration products of aluminate cement are resided in the hardened CAP and hardened RAP, but the roles of AH_3 played in the hardened CAP and hardened RAP are different, and their formation process are not exactly the same. As a result, it can be concluded that the compressive and flexural strengths of hardened RAP are decreased by the presence of noncemented AH_3 and low-strength recycled AFt, while a high-strength hardened CAP is obtained by its hydration products of AFt and AH_3 .

4. Conclusions

In this paper, properties and degradation mechanism of recycled plaster mixed aluminate cement are studied. Obviously, the setting time of RAP is shortened by the rapid recycling of AFt, while the hydration of CAP becomes slow for the low hydration rate of aluminate cement. Surprisingly, the strength of RAP is reduced, which is not aligned with the

routine situation of aluminate cement for increasing the strength of commercial plaster. The decreased strength of RAP is caused by the noncemented AH_3 and low-strength recycled AFt and the fine crystals of the hardened RAP, which was confirmed by SEM and DSC/TG analysis. In principle, this research on recycled plaster mixed aluminate cement will lay a theoretical foundation for the use of recycled plaster.

Data Availability

The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

Acknowledgments

The authors sincerely acknowledge the Key Program of Higher Education of Henan (20A560005), and Youth Key Teachers Funding Program of Henan University of Urban Construction (YCJQNGGJS201908), and Henan Provincial Solid Waste Comprehensive Disposal and Ecological

Utilization Collaborative Innovation Center Open Project Funded Project.

References

- [1] R. Reixach, R. Del Rey, J. Alba, G. Arbat, F. X. Espinach, and P. Mutjé, "Acoustic properties of agroforestry waste orange pruning fibers reinforced polypropylene composites as an alternative to laminated gypsum boards," *Construction and Building Materials*, vol. 77, pp. 124–129, 2015.
- [2] O. Gencel, J. J. del Coz Diaz, M. Sutcu, F. Koksall, F. P. Álvarez Rabanal, and G. Martínez-Barrera, "A novel lightweight gypsum composite with diatomite and polypropylene fibers," *Construction and Building Materials*, vol. 113, pp. 732–740, 2016.
- [3] H.-C. Wu, Y.-M. Xia, X.-Y. Hu, and X. Liu, "Improvement on mechanical strength and water absorption of gypsum modeling material with synthetic polymers," *Ceramics International*, vol. 40, no. 9, pp. 14899–14906, 2014.
- [4] N. B. Singh and B. Middendorf, "Calcium sulphate hemihydrate hydration leading to gypsum crystallization," *Progress in Crystal Growth and Characterization of Materials*, vol. 53, no. 1, pp. 57–77, 2007.
- [5] C. Y. Wang, *Civil Engineering Materials*, Peking University Press, Beijing, China, 2013.
- [6] S. Suárez, X. Roca, and S. Gasso, "Product-specific life cycle assessment of recycled gypsum as a replacement for natural gypsum in ordinary Portland cement: application to the Spanish context," *Journal of Cleaner Production*, vol. 117, pp. 150–159, 2016.
- [7] A. López and A. Lobo, "Emissions of C&D refuse in landfills: a European case," *Waste Management*, vol. 34, no. 8, pp. 1446–1454, 2014.
- [8] T. Raghavendra and B. C. Udayashankar, "Engineering properties of controlled low strength materials using flyash and waste gypsum wall boards," *Construction and Building Materials*, vol. 101, pp. 548–557, 2015.
- [9] C. Chandara, K. A. M. Azizli, Z. A. Ahmad, and E. Sakai, "Use of waste gypsum to replace natural gypsum as set retarders in Portland cement," *Waste Management*, vol. 29, no. 5, pp. 1675–1679, 2009.
- [10] T. Kamei, A. Ahmed, and T. Shibi, "Effect of freeze-thaw cycles on durability and strength of very soft clay soil stabilised with recycled Bassanite," *Cold Regions Science and Technology*, vol. 82, pp. 124–129, 2012.
- [11] T. Kamei, A. Ahmed, and T. Shibi, "The use of recycled bassanite and coal ash to enhance the strength of very soft clay in dry and wet environmental conditions," *Construction and Building Materials*, vol. 38, pp. 224–235, 2013.
- [12] A. P. Godinho-Castro, R. C. Testolin, L. Janke, A. X. R. Corrêa, and C. M. Radetski, "Incorporation of gypsum waste in ceramic block production: proposal for a minimal battery of tests to evaluate technical and environmental viability of this recycling process," *Waste Management*, vol. 32, no. 1, pp. 153–157, 2012.
- [13] J. H. Peng, *Study on Mechanism of Building Gypsum Water Reducer and Retarder*, Chongqing University, Chongqing, China, 2004.
- [14] Y. Kojima and T. Yasue, "Synthesis of large plate-like gypsum dihydrate from waste gypsum board," *Journal of the European Ceramic Society*, vol. 26, no. 4-5, pp. 777–783, 2006.
- [15] L. N. Ren, *Reclaimed Gypsum Properties Change Law and Mechanism*, Chongqing University, Chongqing, China, 2014.
- [16] C. Zhu, J. Zhang, W. Yi, W. Cao, J. Peng, and J. Liu, "Research on degradation mechanisms of recycled building gypsum," *Construction and Building Materials*, vol. 173, pp. 540–549, 2018.
- [17] P. S. Bardella and G. Camarini, "Recycled plaster: physical and mechanical properties," *Advance Material Research*, vol. 374–377, pp. 1307–1310, 2011.
- [18] G. Camarini, S. M. M. Pinheiro, and K. Tannous, "Thermal analysis of recycled gypsum from construction and demolition waste," *Applied Mechanic Material*, vol. 260–261, pp. 977–980, 2012.
- [19] S. M. M. Pinheiro and G. Camarini, "Characteristics of gypsum recycling in different cycles," *International Journal of Engineering and Technology*, vol. 7, no. 3, pp. 215–218, 2015.
- [20] R. H. Geraldo, S. M. M. Pinheiro, J. S. Silva et al., "Gypsum plaster waste recycling: a potential environmental and industrial solution," *Journal of Cleaner Production*, vol. 164, pp. 288–300, 2017.
- [21] G. Camarini, M. C. C. Pinto, A. G. D. Moura, and N. R. Manzo, "Effect of citric acid on properties of recycled gypsum plaster to building components," *Construction and Building Materials*, vol. 124, pp. 383–390, 2016.
- [22] H. C. F. Heloisa, F. C. Cagnoni, and F. F. S. Ferreira, "Comparison of physical and mechanical properties of civil construction plaster and recycled waste gypsum," *Journal of Building Engineering*, vol. 22, pp. 504–512, 2019.
- [23] Z. X. Li, K. D. Xu, J. H. Peng, J. N. Wang, X. W. Ma, and J. S. Niu, "Study on hydration and mechanical property of portland cement-blended recycled plaster materials," *Advances in Materials Science and Engineering*, vol. 2018, Article ID 2692347, 8 pages, 2018.
- [24] Z. X. Li, K. D. Xu, J. H. Peng, J. N. Wang, X. W. Ma, and J. S. Niu, "Study on hydration and mechanical property of quicklime blended recycled plaster materials," *Construction and Building Materials*, vol. 202, pp. 440–448, 2019.
- [25] J. G. Zheng, Z. M. Wang, Y. F. Yuan et al., *GB/T 9776-2008, Calcined Gypsum*, Standardization Administration of China, Beijing, China, 2008.
- [26] M. Zhao, J. H. Peng, M. T. Zhang, Z. X. Li, and D. L. Zhu, "Performance improvement and mechanism of model-gypsum by aluminate gypsum," *Journal of Southeast University (Natural Science Edition)*, vol. 44, no. 9, pp. 1030–1036, 2014.
- [27] S. T. Tong, "Study on analysis for crystal species of gypsum by applying differential scanning calorimetry (DSC)," *Journal of Wuhan University of Science and Technology (Natural Science Edition)*, vol. 24, no. 3, pp. 243–256, 2001.
- [28] M. Zhao, J. H. Peng, M. T. Zhang, Z. X. Li, and H. X. Zhao, "Mechanisms for the enhanced ceramics gypsum model with ordinary Portland cement," *Journal of Hunan University (Natural Sciences)*, vol. 41, no. 12, pp. 46–52, 2014.