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22 This is due to the increase in viscosity of the asphalt after heavy oxidation, which makes it difficult for  
23  $C_{20}H_{42}$  to migrate, while the increase in storage temperature can release the potential of the degree of  
24 thermoreversible aging of the asphalt by accelerating the movement of wax molecules. And like  $C_{20}H_{42}$ , the  
25 grade loss of asphalt blended with  $C_{30}H_{62}$  is different at different storage temperatures and oxidation levels,  
26 which proves that for heavily oxidized asphalt, multiple storage temperatures should be selected for testing  
27 its grade loss. Combined with thermal and morphological analysis, the introduction of heavy oxidation not  
28 only increases the amount of wax precipitation, but also significantly increases the thermoreversible aging  
29 of the asphalt by lowering the  $T_g$  of the asphalt and forming more perfect crystals during low temperature  
30 storage. Finally, by comparing the grade loss due to cold storage and extended oxidation, it can be found that  
31 the incorporation of  $C_{20}H_{42}$  will make the asphalt more influenced to physical hardening. Therefore, it is  
32 necessary to study the effect of asphalt wax content on the degree of thermoreversible aging of heavily  
33 oxidized asphalt.

34 **Keywords:** Asphalt binder; Thermoreversible aging; Crystalline wax; Oxidative aging; Physical hardening

## 35 **1. Introduction**

36 Petroleum asphalt is one of the most widely used materials in pavement construction  
37 owing to its good bonding and waterproofing properties. Because of its unique characteristics  
38 such as excellent waterproofing, high damping ratio, good integrity, asphalt has also been  
39 utilized in other applications such as water conservancy, railway engineering. Countries such  
40 as the United States, Germany, Japan, and China has carried out a large number of studies on  
41 asphalt concrete subgrade [1, 2]. Compared with other ordinary subgrade, the advantages of

42 asphalt concrete subgrade mainly include the following five aspects: (1) reducing the stress  
43 effect of upper loads on the subgrade and transferring them to the lower structure; (2) keeping  
44 the moisture content of the subgrade stable as an impermeable layer; (3) acting as a support  
45 layer for ballast and reducing ballast splash; (4) alleviating train vibration and reducing noise;  
46 (5) reducing the whole life cycle cost of the track structure due to the reduction of later  
47 maintenance [3, 4]. The authors' team has systematically studied the damage pattern of full-  
48 section railway asphalt concrete subgrade and found that when asphalt concrete is positioned  
49 at a lower level in the railway subgrade, the train loads tend to be smaller, thus determined that  
50 thermal shrinkage cracks become an important factor affecting the durability of asphalt  
51 concrete [5, 6]. Among the many influencing factors, thermoreversible aging could affect  
52 asphalt concrete subgrade performance in the long run [7-9]. One of the main reasons is that  
53 other forms of aging mainly occur on the pavement surface while thermoreversible aging  
54 occurs in the whole asphalt pavement structure, especially in cold areas.

55 In order to understand the mechanism of thermoreversible aging as an important basis for  
56 the production of high-quality asphalt and solving related pavement distresses, in recent years  
57 researchers have devoted themselves to explain the causes of physical hardening in asphalt.  
58 The term physical hardening was first introduced by Anderson from Shell Laboratories,  
59 Amsterdam, The Netherlands, who attributed the phenomenon to the precipitation of waxes  
60 and the aggregation of asphaltenes, noting that both processes are very slow at low temperature  
61 [10]. Some asphalt will form large microcrystalline wax crystals during long-term low-  
62 temperature storage and are affected by the change of pavement temperature. Therefore,

63 different asphalt could have different allowable wax contents, which is also the reason why  
64 wax is considered to be the main factor in thermoreversible aging [11, 12]. In contrast, studies  
65 have shown that waxes in asphalt are a mixture of alkanes, cycloalkanes and aromatic  
66 compounds and dominated by n-alkanes with carbon numbers of  $C_{15}\sim C_{57}$  [13]. Kovinichi found  
67 that although  $C_{20}H_{42}$  can promote the physical hardening in asphalt, this tendency is affected  
68 by the base asphalt binder and is not determined by the content of  $C_{20}H_{42}$  solely [14].  
69 Subsequent studies found that not all waxes will aggravate the thermoreversible aging of  
70 asphalt. Compared with  $C_{20}H_{42}$ ,  $C_{30}H_{62}$  is significantly less effective in promoting the physical  
71 hardening of asphalt, while squalanecan improve the low temperature performance of asphalt  
72 without aggravating the degree of thermoreversible aging [15, 16].

73         Despite a number of studies have conducted a series of studies on the effects of waxes  
74 with different numbers of carbon molecules on the thermoreversible aging behaviour of asphalt,  
75 the low temperature performance and grade loss of asphalt in most of these studies are  
76 measured after a PAV-20h test using the standard BBR procedure [15, 17, 18]. In recent years  
77 some practitioners have questioned this conventional PAV aging method, as Simon's research  
78 has shown that the PAV-20h test is only able to simulate the condition of asphalt pavements  
79 after 5-6 years of use, which is clearly not suitable to reflect the field aging of the pavement  
80 surface after a long term service [19]. In order to be able to simulate the long-term use of  
81 asphalt pavement, Simon proposed to prolong the aging time of PAV test in another study to  
82 prepare heavily oxidized asphalt, and found that the asphalt under a PAV-40h test can simulate  
83 the condition of asphalt pavements after 10-15 years of use [20]. Other researchers have

84 explored the low-temperature performance distribution of different asphalts after extended  
85 aging. Kilger's research shows that the effect of recycling oils, including bio-oil and REOB,  
86 on the improvement of asphalt's low-temperature performance tends to be weakened after  
87 heavy oxidation compared with asphalt without recycling oils [21]. Ma, on the other hand,  
88 found that the degree of thermoreversible aging of recycled asphalt blended with three different  
89 rejuvenators can only be distinguished after RTFO and PAV aging, and increases with the  
90 extension of PAV aging time [22]. This indicates that like the low-temperature properties of  
91 asphalt, the degrees of thermoreversible aging of different asphalts after severe oxidation will  
92 also be different, and thus it is necessary to investigate the mechanism of the effect of waxes  
93 on the physical hardening properties of heavily oxidized asphalt binders.

94 In view of this, the purpose of this paper is to investigate the interaction between  
95 thermoreversible aging and irreversible aging based on extended bending beam rheometer  
96 (ExBBR) tests, using six different asphalt samples blended with pure wax, and analyzing the  
97 grade loss of heavily oxidized asphalt by introducing three different aging methods, RTFO,  
98 PAV-20h, and PAV-40h test. Based on the results of ExBBR tests, modulated differential  
99 scanning calorimetry (MDSC) and atomic force microscopy (AFM) were used to analyze the  
100 mechanism of the effect of wax on the physical hardening properties of heavily oxidized asphalt.

101 **2. Experimental program**

102 **2.1. Materials**

103 **2.1.1 Asphalt binder**

104 According to research, the wax content of asphalt will affect the road performance of  
105 asphalt pavement. In this paper, six kinds of asphalt from different crude oils were selected,  
106 among which the base asphalt from Venezuela came from the Strategic Highway Research  
107 Program (SHRP) Material Reference Library (MRL), which is labeled as ABG, according to  
108 the research there is basically no wax in ABG [23]. While the other asphalt samples are used  
109 for comparison, which are labeled as SWE, CAL, MAO, UNK, and ZNH, whose source and  
110 performance grade (PG) are shown in Table 1. The first three kinds of asphalt binders are  
111 commonly used in other countries while the rest of the asphalt binders usually used in asphalt  
112 pavement construction in China. To compare the oxidative aging with thermoreversible aging,  
113 in this research, the asphalt binders are tested after going through the rolling thin film oven  
114 (RTFO) as well as 20h and 40h of pressure aging vessel (PAV) tests.

115 **Table 1.** Basic properties of asphalt binders.

Sample Code	Source	Limiting high temperature	Limiting low temperature
		PG (°C)	PG (°C)
ABG	Venezuela	65.2	-29.7
SWE	Sweden	63.8	-31.3

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CAL	Caltex, Korea	67.1	-26.5
MAO	Maoming, China	64.9	-28.6
UNK	Unknown, China	65.7	-29.5
ZHH	Zhonghai, China	66.3	-27.7

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116 **2.1.2 Additives**

117 According to previous research on wax in asphalt [15], the waxes in asphalt are mainly  
118 composed of saturated straight-chain hydrocarbons. Since waxes are an important cause of  
119 thermoreversible ageing of asphalt. Therefore, two commercial n-alkanes produced by Sigma-  
120 Aldrich were chosen as pure wax additives in this study, namely n-eicosane ( $C_{20}H_{42}$ ) and n-  
121 dodecane ( $C_{12}H_{26}$ ), with a purity of 99%. Considering that most asphalts for paving have a wax  
122 content of less than 7% [24-26], the percent of wax additives were blended at 1%, 3%, 5% and  
123 7% by mass of asphalt binder. To ensure that the wax can be uniformly distributed in the asphalt,  
124 the base asphalt binder is heated to 165°C during the mixing process and the prepared model  
125 asphalt is sheared for 1h using a shear mixer.

126 **2.2. Test Methods**

127 **2.2.1 Extended bending beam rheometer (ExBBR)**

128 Among different low-temperature evaluation methods developed for asphalt binder in  
129 recent years [27, 28], the most mature and widely used one is the bending beam rheometer  
130 (BBR) test to measure the low temperature performance grade (PG) of asphalt binders.



131 However, it has also been shown that asphalt pavement designed based on PG performance  
132 suffered from severe low-temperature cracking from time to time [29], due to the increase in  
133 stiffness of asphalt binders at low temperatures with increasing storage time [30, 31]. Based on  
134 this, the ExBBR test was developed. Compared to the regular BBR test, the ExBBR test  
135 requires testing the PG performance of the asphalt in six states (3 conditioning time×2 storage  
136 temperature) separately. The grade loss obtained from the ExBBR test can be used to evaluate  
137 the degree of thermoreversible aging of asphalt. According to the requirements of AASHTO  
138 TP122-17, the grade loss of asphalt should be less than 6°C.

### 139 2.2.2 *Modulated differential scanning calorimetry (MDSC)*

140 To understand the thermoreversible aging process of asphalt binders throughout the  
141 service temperature range, the MDSC test was chosen for the study to analyze the thermal  
142 behavior of asphalt. In contrast to conventional thermal analysis tests, the MDSC test allows  
143 the total heat flow signal to be divided into reversing and non-reversing heat flow signals,  
144 which has been favored by many researchers [32, 33]. The procedure is as follows. First, the  
145 sample is heated to 120°C and then left at this temperature for 10min so that to melt all the  
146 crystal structures. Second, the specimen is cooled down to 80°C at a cooling rate of 20°C/min  
147 to avoid overrate oxidation and saving time. Next, the sample is rapidly cooled to -90°C at  
148 10°C/min and kept for 10min. Then, the asphalt sample is heated to 120°C at a heating rate of  
149 600°C/h (10°C/min). Finally, the total heat flow, reversing heat flow, non-reversing heat flow  
150 and the first derivative of reversing heat flow are employed as the four key parameters to  
151 characterize the thermal behavior of asphalt binder.

### 152 2.2.3 Atomic force microscope (AFM)

153 As a visco-elastic-plastic material with significant temperature dependence, asphalt has  
154 been the subject of many studies on the formation and growth mechanisms of the ‘bee-like’  
155 structure in the AFM microstructure parameters. It has been shown that the ‘bee-like’ structure  
156 can be explained based on the wax nucleation theory, leading to the conclusion that its  
157 morphological characteristics are related to the heating history, cooling rate and aging degree  
158 of the sample [34-36]. As this study is focused on the differences in the microscopic  
159 morphology of different asphalts after low-temperature storage, the asphalt is stored at low  
160 temperature for 72h, and the surface of the asphalt is tested in amplitude mode using an AFM  
161 probe. In order to ensure the repeatability of the observation results, three replicates were  
162 carried out for each asphalt sample.

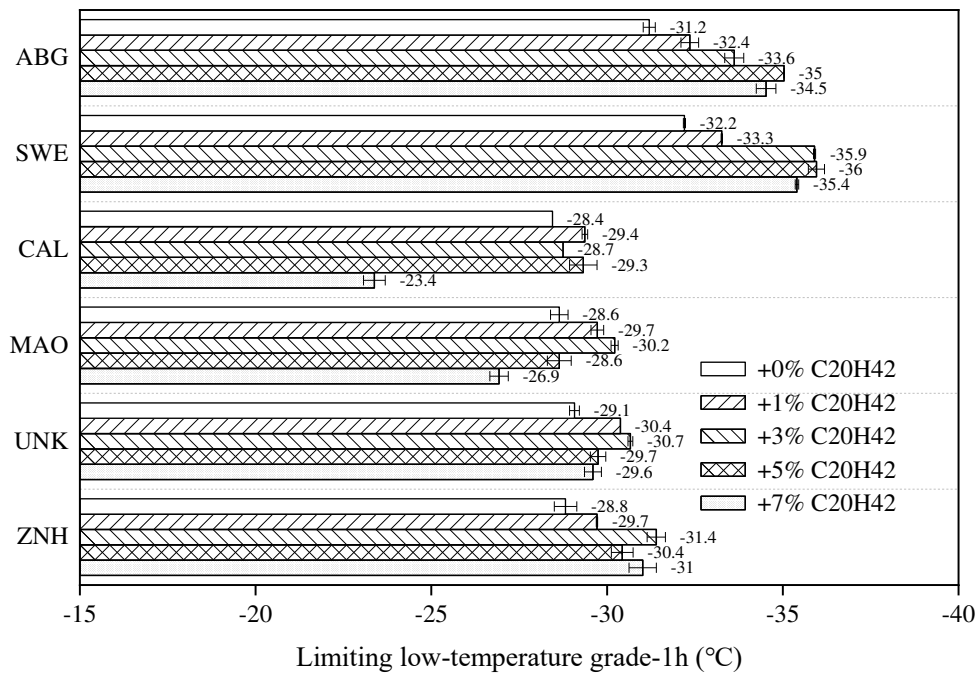
## 163 3. Results and discussion

### 164 3.1 Oxidation effects on limiting grade of asphalt binder

165 Fig. 1 shows the oxidation effects on limiting grade of asphalt binder based on ExBBR  
166 test. As can be seen, the addition of  $C_{20}H_{42}$  to different RTFO aged asphalts effectively  
167 decreased the limiting grade of asphalt in the beginning. In fact, the addition of 5% of  $C_{20}H_{42}$   
168 to ABG and SWE exhibited the best improvement in terms of low-temperature performance  
169 (limiting grade was reduced by 4°C), followed by MAO, UNK and ZNH (limiting grade was  
170 reduced by 1°C~2°C) while CAL had the least improvement (limiting grade was reduced by  
171 less than 1°C). However, the limiting grade of most asphalts began to increase after adding 3%

172 of C<sub>20</sub>H<sub>42</sub> while for ABG and SWE the turning point was at 5%. Combined with previous  
 173 research results [37, 38], this may be due to the different wax contents of asphalts from different  
 174 origins which asphalt with low wax content can accommodate more C<sub>20</sub>H<sub>42</sub> before saturation.

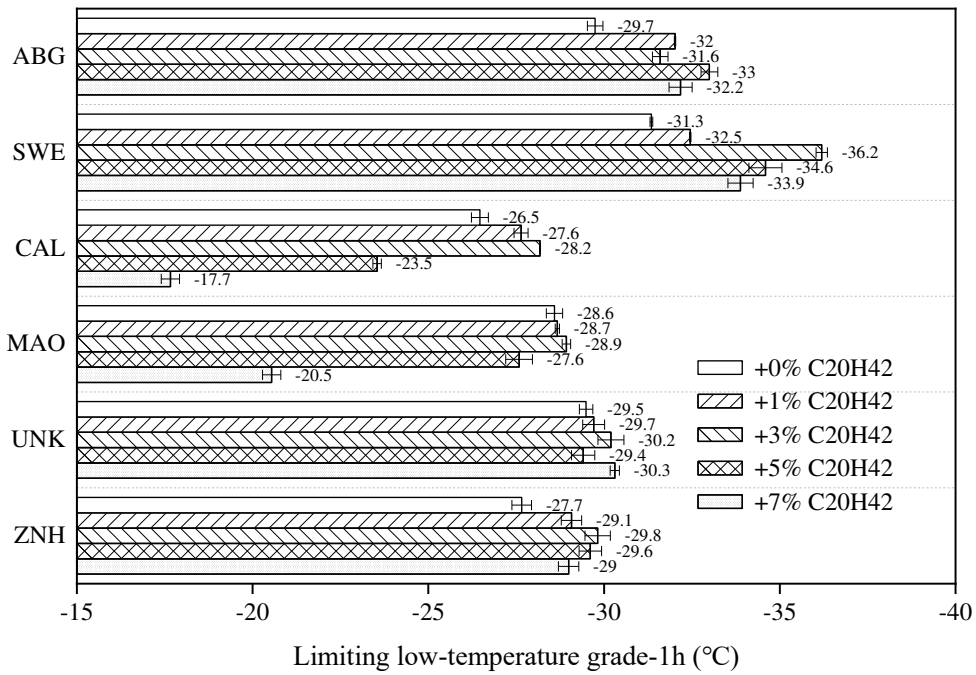
175 By comparing the limiting grade of asphalt at different aging stages: 40h PAV>20h  
 176 PAV>RTFO, for most asphalts compared to RTFO the limiting grade of asphalt after 40h PAV  
 177 has increased by 1°C~2°C. Meanwhile, it is worth noting that the limiting grade of ABG and  
 178 SWE after 40h PAV starts to increase with more than 3% of C<sub>20</sub>H<sub>42</sub>, which indicates that the  
 179 introduction of heavy oxidation will not only lead to the deterioration of asphalt low  
 180 temperature performance, but also lead to the decrease in the capacity for n-alkanes in asphalt.



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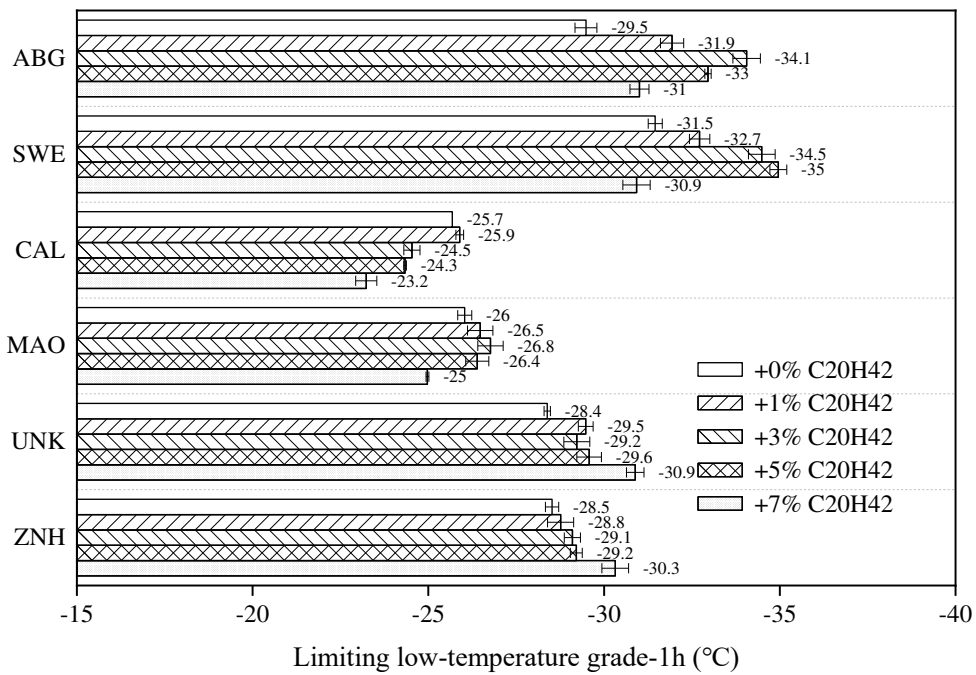
(a) RTFO



183

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(b) 20h PAV



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(c) 40h PAV

187

Fig. 1. Oxidation effects on limiting grade of asphalt binder.

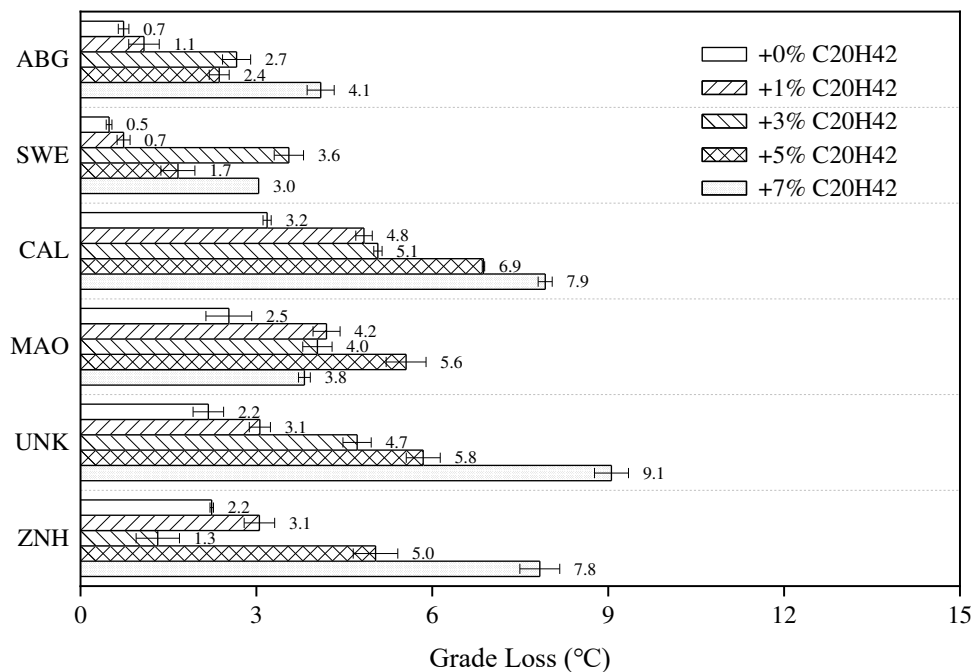
### 188 *3.2 Oxidation effects on grade loss of asphalt binder*

189 If only based on the limiting grade of the asphalt binder, it can be easily concluded that  
190 for most asphalt the addition of  $C_{20}H_{42}$  is an effective solution to improve the low temperature  
191 performance of asphalt and that this improvement is almost independent of the effects of  
192 oxidative aging. However, some studies in recent years have shown that with the extension of  
193 the storage time of asphalt in low-temperature environment, the wax and asphaltene in the  
194 asphalt begin to slowly aggregate and exhibit physical hardening, which indirectly affects the  
195 low-temperature performance of the asphalt [25, 39].

196 According to the result of grade loss of asphalt blended with  $C_{20}H_{42}$  in low temperature  
197 environment given in Fig. 2, it can be observed that there is no significant difference in the  
198 grade loss of six asphalt without wax in RTFO aging stage, while the grade loss of six asphalt  
199 after blending with  $C_{20}H_{42}$  can be easily distinguished. Among the six asphalt, CAL, MAO,  
200 UNK and ZNH blended with high content of  $C_{20}H_{42}$  all showed significant hardening behavior  
201 with grade loss higher than  $6^{\circ}C$ . In contrast, the degree of thermoreversible aging of ABG and  
202 SWE, on the other hand, is not significantly affected by n-alkanes, and the maximum grade  
203 loss is only about  $4^{\circ}C$ . Referring to the research of Ding [23], this is due to the fact that ABG  
204 contains almost no wax and thus the asphalt can accommodate more wax before saturation.

205 Compared to the RTFO aging stage, the grade loss of most asphalt has increased after  
206 PAV. However, for ABG and SWE blended with  $C_{20}H_{42}$ , the grade loss of asphalt after 20h  
207 PAV remained almost unchanged. In contrast, after the introduction of 40h PAV, the grade loss

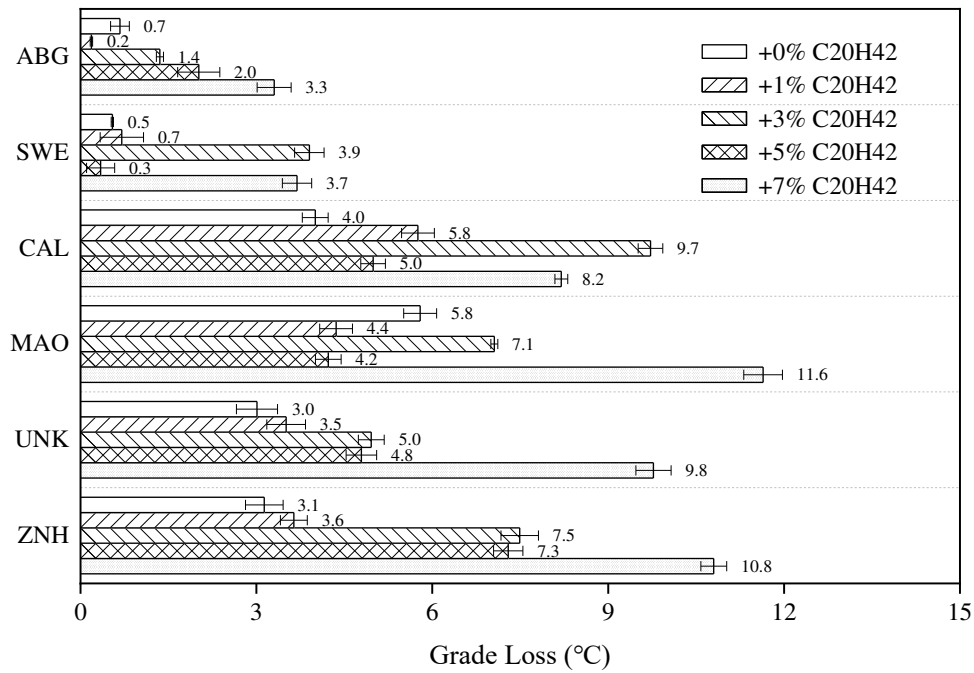
208 of the above two asphalt after heavy oxidation has increased to around 6°C. This can be  
 209 explained using the findings of Tabatabaee [40, 41], that is, the presence of waxes will promote  
 210 the phase separation of asphaltene components in asphalt. As the degree of oxidation increases,  
 211 the increase of asphaltene content in asphalt will accelerate the rate of phase separation,  
 212 resulting in a rapid increase in the hardening tendency of asphalt. However, this theory is not  
 213 sufficient to explain the effect of the thermo-oxidative aging on the degree of thermoreversible  
 214 aging of the other four types of asphalt. Taking MAO+3% C<sub>20</sub>H<sub>42</sub> as an example, the grade loss  
 215 of asphalt at the 20h PAV and 40h PAV stages is 7.1°C and 7.6°C, respectively. While for  
 216 UNK+3% C<sub>20</sub>H<sub>42</sub>, the grade loss of heavily oxidized asphalt decreased from 5.0°C at the 20h  
 217 PAV stage to 3.2°C at the 40h PAV stage.



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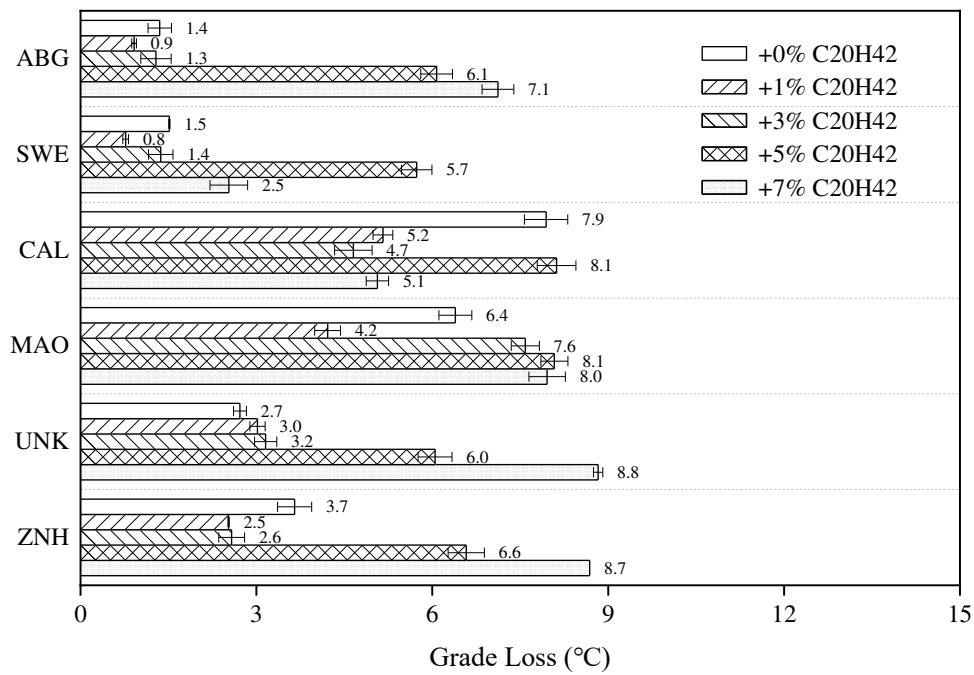
(a) RTFO



220

221

(b) 20h PAV



222

223

(c) 40h PAV

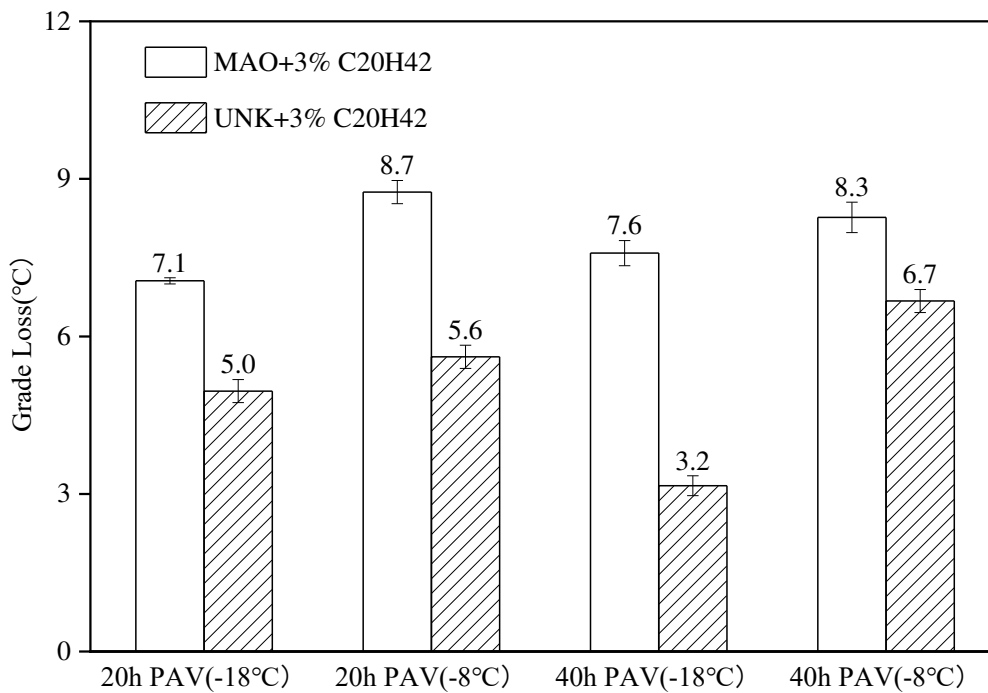
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**Fig. 2.** Oxidation effects on grade loss of asphalt binder.

225 **3.3 Storage temperature effects on grade loss of asphalt binder**

226 To further explain the effect of heavy oxidation on the grade loss of MAO+3% C<sub>20</sub>H<sub>42</sub>  
227 and UNK+3% C<sub>20</sub>H<sub>42</sub>, this study chose -8°C as the control group based on the original storage  
228 temperature (-18°C) to investigate the effect of storage temperature on the grade loss of heavily  
229 oxidized asphalt, as shown in Fig. 3. It can be seen from the figure that when the storage  
230 temperature is increased from -18°C to -8°C, the grade loss of asphalt also shows an increasing  
231 tendency. But for the two different asphalt, the increase in grade loss differs. For MAO+3%  
232 C<sub>20</sub>H<sub>42</sub>, the degree of thermoreversible aging of the asphalt after 20h PAV and 40h PAV  
233 remains almost the same after increasing the storage temperature, whereas the difference in  
234 grade loss between each is only 0.4°C. As for UNK+3% C<sub>20</sub>H<sub>42</sub>, the increase in storage  
235 temperature unleashed the potential for the degree of thermoreversible aging of heavily  
236 oxidized asphalt, and at a storage temperature of -8°C, the grade loss of asphalt after 40h PAV  
237 increased by 3.5°C, surpassing the sample after 20h PAV.





238

239

**Fig. 3.** Grade loss of asphalt binders under different cold storage temperature.

240

Such phenomenon can be explained according to the phase separation theory [42-45],

241

according to which asphalt can be viewed as a binary mixture consisting of bitumen and wax,

242

a typical binary mixture is shown in Fig. 4. As can be seen in the figure, the distribution states

243

of the phases of the asphalt differ at different temperatures. When the temperature is high

244

enough ( $T_1$ ), the mixture is stable at any binary ratio. And as the temperature decreases, the

245

mixture may go through a metastable state ( $T_2$ ) and eventually reach an unstable state ( $T_3$ ).

246

After entering the metastable state or even the unstable state, the wax phase in the asphalt starts

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to separate from the asphalt phase, causing the aggregation of wax molecules and eventually

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leading to the crystalline precipitation of wax in the asphalt. Combined with the findings of Lei

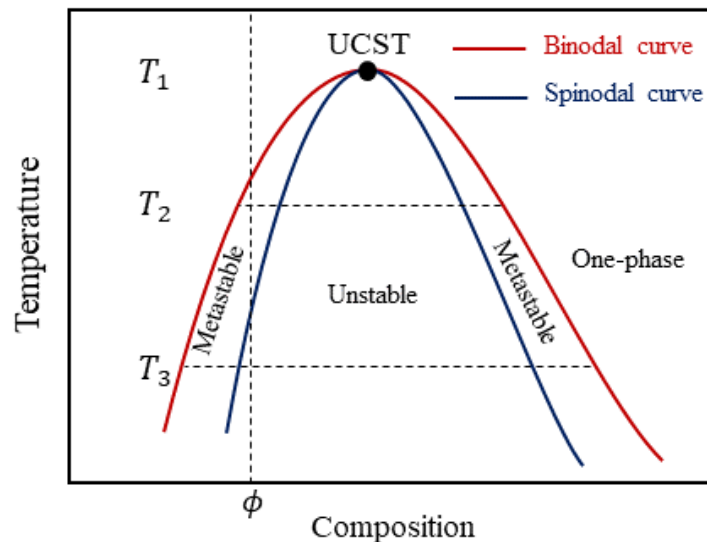
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[46], it can be explained that the viscosity of asphalt increases after heavy oxidation, which

250

makes it difficult for wax molecules to migrate and thus inhibits wax precipitation. Therefore,

251 even if heavily oxidized asphalt has reached an unstable state in a low temperature environment,  
252 there may not be significant thermoreversible aging inside the asphalt. In this case, the increase  
253 in storage temperature will effectively accelerate the movement rate of wax molecules, and the  
254 degree of thermoreversible aging of asphalt will be increased.

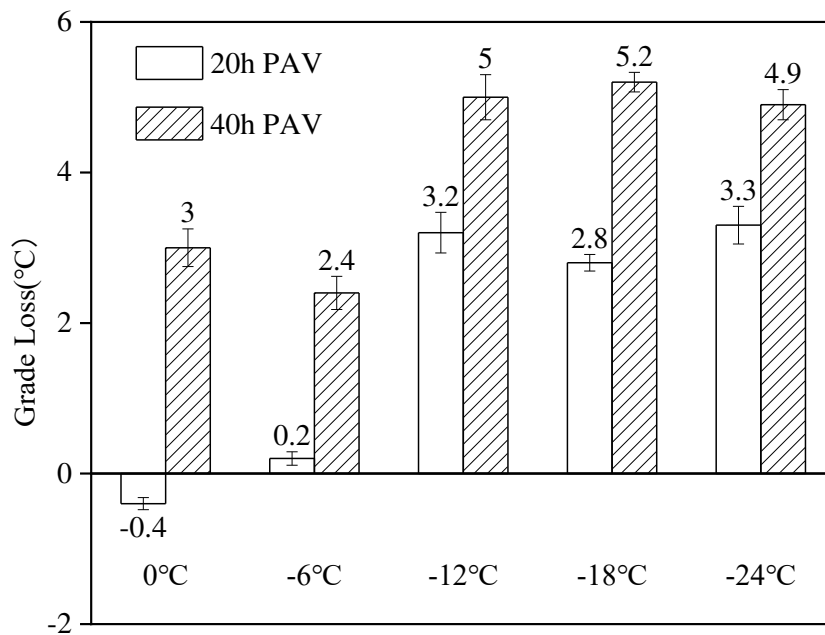


255

256 **Fig. 4.** Possible phase diagram of the asphalt at different temperature.

257 In order to be able to confirm whether the storage temperature will affect the precipitation  
258 and crystallization of n-alkanes other than  $C_{20}H_{42}$  in asphalt. In this study, ABG with almost no  
259 wax is selected as the base asphalt, and  $C_{30}H_{62}$  is used as the additive to study the effect of  
260 storage temperature on the grade loss of PAV aged ABG+5%  $C_{30}H_{62}$ , and the results are shown  
261 in Fig. 5. As shown in the figure, if only following the traditional test method, the grade loss  
262 of oxidized asphalt ( $-18^{\circ}C$ , 20h PAV) is  $2.8^{\circ}C$ . By changing the storage temperature and  
263 oxidation degree of the tested asphalt, the grade loss is only  $-0.4^{\circ}C$  under the optimal storage  
264 conditions ( $0^{\circ}C$ , 20h PAV), while the grade loss of asphalt increased to  $5.2^{\circ}C$  under the worst  
265 storage conditions ( $-18^{\circ}C$ , 40h PAV). This indicates that, like  $C_{20}H_{42}$ , the effect of  $C_{30}H_{62}$  on

266 the degree of thermoreversible aging of asphalt is quite different at different storage  
 267 temperatures and oxidation degrees. Therefore, in order to better characterize the  
 268 thermoreversible aging of heavily oxidized asphalt, multiple storage temperatures should be  
 269 selected to test the grade loss of the asphalt.



270

271

**Fig. 5.** Grade loss of ABG+5%C30 caused by isothermal conditioning.

272

### **3.4 Influence mechanism of oxidative aging on physical hardening of asphalt binder**

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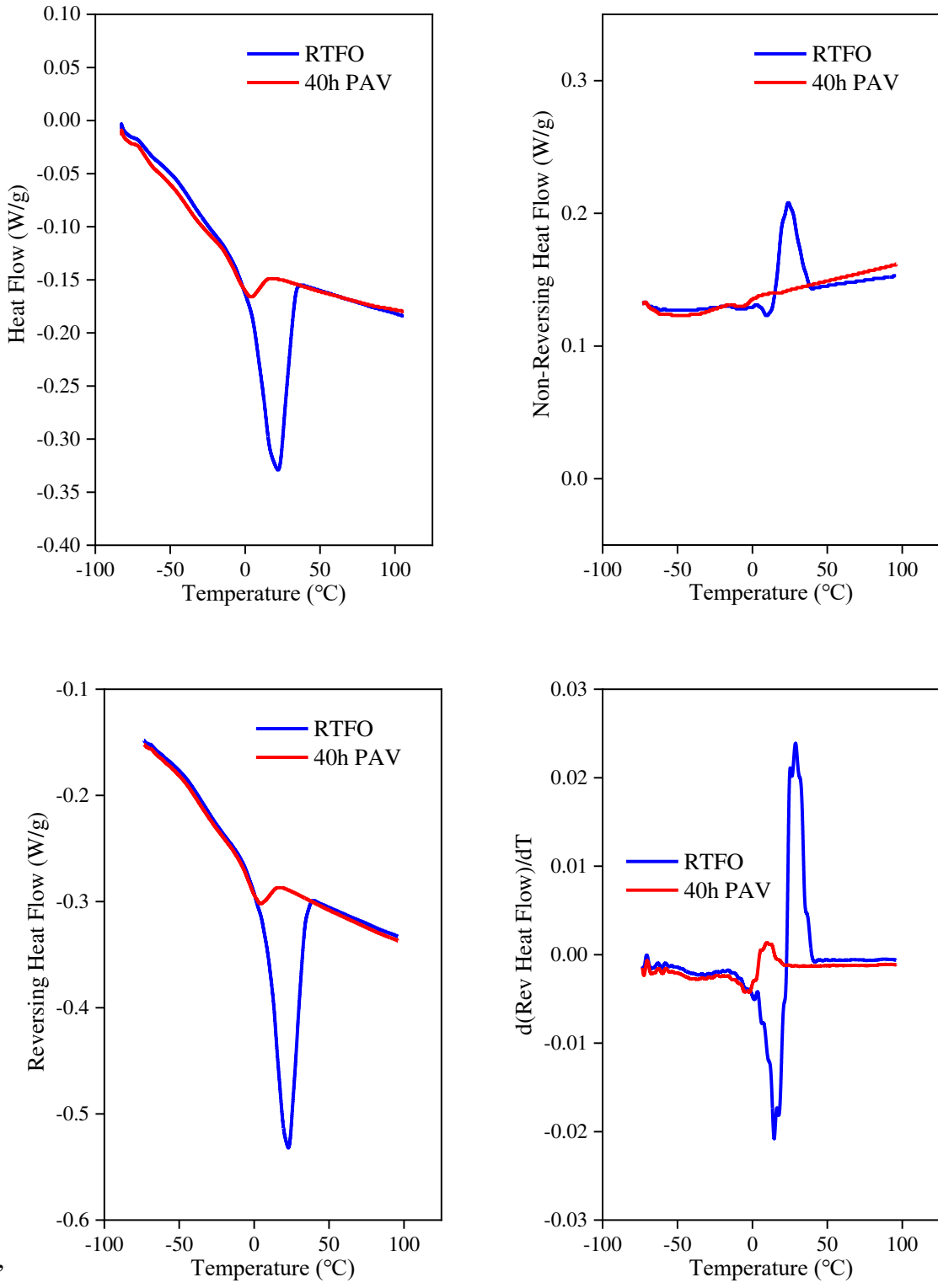
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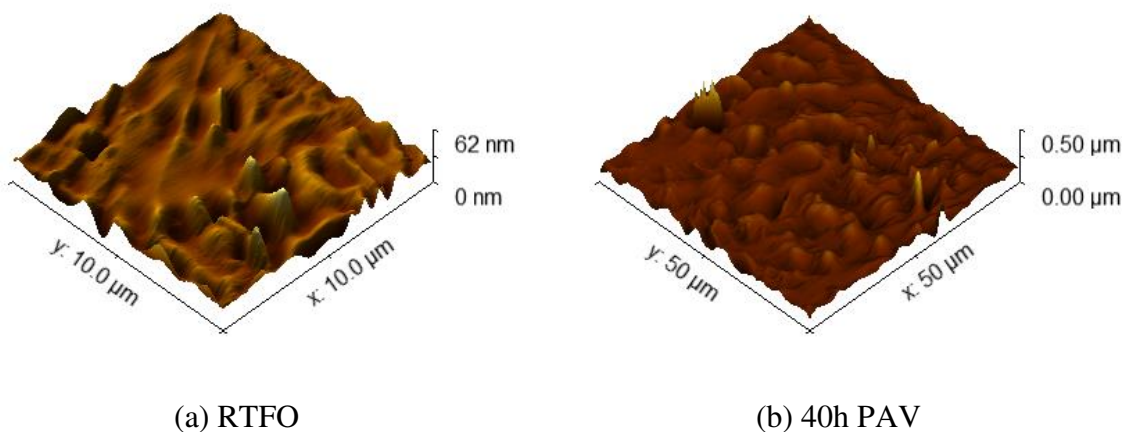
In order to further explain the influence mechanism of oxidative aging on the physical hardening of asphalt, two aging stages of ABG+5% C<sub>20</sub>H<sub>42</sub> are selected for the study, RTFO and 40h PAV, with a grade loss of 2.4°C and 6.1°C, respectively, and MDSC is used to investigate the thermal behavior of ABG+5% C<sub>20</sub>H<sub>42</sub> at different oxidative aging stages, as shown in Fig. 6. Based on the non-reversing heat flow curves of the asphalt, it can be seen that there is an endothermic peak in the asphalt at both aging stages, while the area of the

279 endothermic peak reflects the amount of crystal fraction (CF) in the asphalt, which is usually  
280 caused by the crystallization or precipitation of long-chain alkane compounds [47, 48].  
281 Compared with RTFO, the temperature corresponding to the endothermic peak of asphalt after  
282 40h PAV decreased from 9.6°C to -6.0°C, with an area increase of 12.3%. As for the reversing  
283 heat flow of asphalt, significant difference in the shape of the DSC curves of asphalt at different  
284 aging stages can be observed, indicating that phase separation occurs in asphalt after heavy  
285 oxidation. The first derivative of reversing heat flow curves of the asphalt at the RTFO and 40h  
286 PAV aging stages have an maximum value at 28.3°C and 9.5°C, respectively, which can be  
287 regarded as the glass transition temperature ( $T_g$ ) of the asphalt [9]. The decrease of  $T_g$  of asphalt  
288 after heavy oxidation will lead to a larger proportion of  $C_{20}H_{42}$  in high-elastic state in the early  
289 cooling stage, and this part of  $C_{20}H_{42}$  will be changed from high-elastic state to glassy state  
290 with higher molecular chain stiffness after long-term low-temperature storage. This means that  
291 the heavy oxidation will not only cause more waxes to crystallize out of the asphalt, but will  
292 also significantly aggravate the degree of thermoreversible aging of the asphalt by lowering the  
293  $T_g$  of the asphalt to a more perfect crystal during low temperature storage.



**Fig. 6.** Thermal behavior of ABG+5% C<sub>20</sub>H<sub>42</sub> under different oxidation degree.

295 And the surface micromorphology images of ABG+5% C<sub>20</sub>H<sub>42</sub> under two different aging  
296 states in AFM amplitude mode are shown in Fig. 7. From Fig. 7, it can be seen that there are a  
297 large number of peaks like protrusions on the asphalt surface after RTFO, while the number of  
298 peaks on the surface increases after 40h PAV and its peak value also grows to 0.5μm. Such  
299 peak like protrusion is related to the crystallization of microcrystalline wax and waxy  
300 molecules when cooled to the test temperature [49]. To quantitatively evaluate the effect of  
301 peak like protrusions on the surface morphology of the asphalt, the average roughness ( $R_a$ ) and  
302 root mean square roughness ( $R_q$ ) are calculated by Gwyddion software to describe the  
303 roughness of asphalt, the values of which are shown in Table 2. The results show that after  
304 heavy oxidation, the  $R_a$  and  $R_q$  of the asphalt increased by 5.58 mm and 8.28 nm, respectively,  
305 indicating that the increase in oxidation degree will lead to an increase in the surface roughness  
306 of asphalt, that is, more wax will precipitate from asphalt crystals, leading to an increase in the  
307 degree of thermoreversible aging of asphalt.



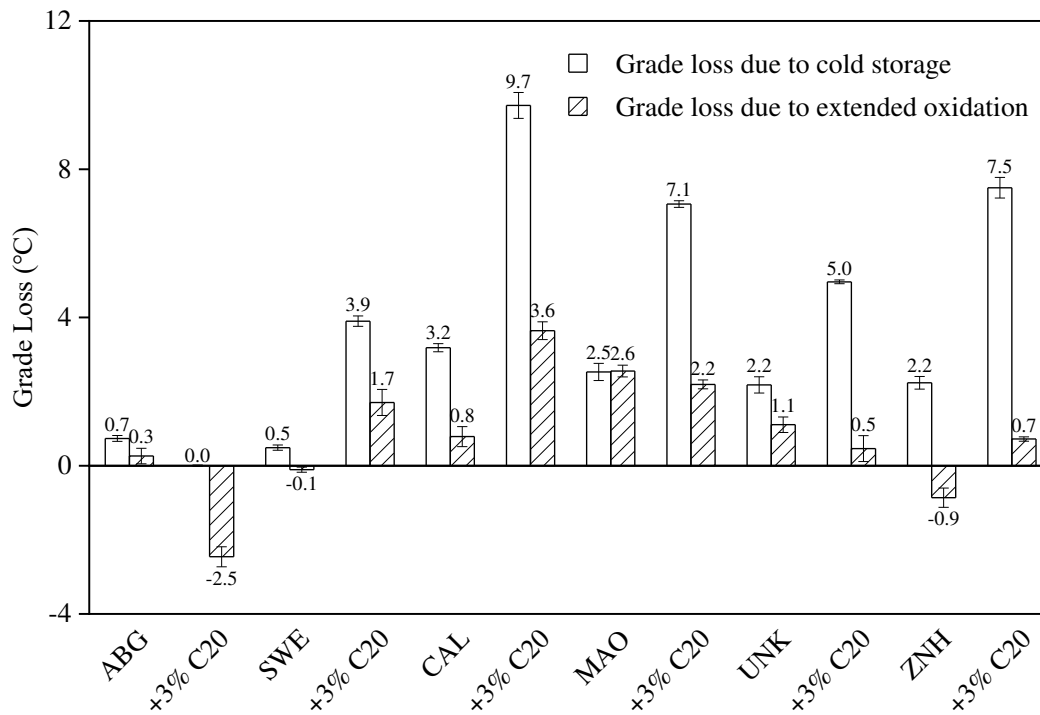
308 **Fig. 7.** AFM images of ABG+5% C<sub>20</sub>H<sub>42</sub> under different oxidation degree.

309 **Table 2.** Surface roughness statistics of ABG+5% C<sub>20</sub>H<sub>42</sub>.

Oxidation degree	$R_a$ (nm)	$R_q$ (nm)
RTFO	10.68	15.17
40h PAV	16.26	23.38

310 **3.5 Comparison between thermoreversible aging and oxidation**

311 Fig. 8 compares the grade loss due to cold storage and extended oxidation. As shown, for  
312 most of the asphalts, physical hardening has a stronger impact on grade loss than oxidation.  
313 For the asphalt samples not blended with n-alkanes, there is no significant dominance of  
314 physical hardening on grade loss, and the difference between the two is less than half a 3°C. In  
315 contrast, with asphalt blended with C<sub>20</sub>H<sub>42</sub>, the grade loss due to cold storage increased  
316 significantly, and the grade loss gap caused by cold storage and extended aging grew to 6°C,  
317 which indicates that the grade loss of wax-blended asphalt is determined by physical hardening.  
318 Therefore, it is necessary to study the effect of asphalt wax content on the degree of  
319 thermoreversible aging of heavily oxidized asphalt.



320

321

**Fig. 8.** Grade loss of asphalt binder due to cold storage and extended oxidation.

322

#### 4. Conclusions

323

In this paper, the effects of cold storage temperature on the thermoreversible aging characteristics of asphalt binder samples doped with wax additives, especially heavily oxidized asphalts, were studied through a series of laboratory experiments. The following conclusions can be drawn from the results:

327

(1) With the appropriate addition of  $C_{20}H_{42}$ , the limiting grade of asphalt can be reduced,

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while due to the different sources of asphalt crude oil, some asphalt with lower wax content

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can accommodate more n-alkanes. While the introduction of heavy oxidation will not only lead

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to the deterioration of asphalt low temperature performance, but also lead to the decrease in the

331

capacity for n-alkanes in asphalt.



332 (2) Among the six asphalt binders, for ABG and SWE blended with  $C_{20}H_{42}$ , the maximum  
333 grade loss of both asphalt binders was only about  $4^{\circ}C$  during the RTFO stage due to the  
334 extremely low wax content, while the grade loss of asphalt after 20h PAV was almost  
335 unchanged. However, after 40h PAV, the grade loss of asphalt grows to  $6^{\circ}C$ , which indicates  
336 that asphalt with good grade loss under low aging conditions may also show significant  
337 thermoreversible aging behavior after heavy oxidation.

338 (3) For the  $C_{20}H_{42}$ -doped asphalt with high viscosity, there is a tendency to decrease the  
339 grade loss of asphalt in low temperature storage after heavy oxidation instead. This is due to  
340 the viscosity of asphalt increases after heavy oxidation, which makes it difficult for wax  
341 molecules to migrate, thus inhibiting the precipitation of wax. In this case, the increase in  
342 storage temperature will effectively accelerate the movement of wax molecules, thus releasing  
343 the potential of the degree of thermoreversible aging of heavily oxidized asphalt.

344 (4) Similar to  $C_{20}H_{42}$ , for heavy oxidized asphalt blended with  $C_{30}H_{62}$ , the effect of  $C_{30}H_{62}$   
345 on the degree of thermoreversible aging of asphalt is quite different at different storage  
346 temperatures and oxidation levels, and the grade loss distribution of asphalt varies from  $-0.4^{\circ}C$   
347 to  $5.2^{\circ}C$ . Therefore, in order to better characterize the thermoreversible aging of heavily  
348 oxidized asphalt, multiple storage temperatures should be selected to test the grade loss of  
349 asphalt.

350 (5) The results of MDSC and AFM show that heavy oxidation will not only cause more  
351 wax to crystallize out of the asphalt, which leads to an increase in surface roughness, but also

352 significantly increases the thermoreversible aging of the asphalt by lowering the  $T_g$  of the  
353 asphalt and forming more perfect crystals during low temperature storage.

354 (6) By comparing the grade loss due to cold storage and extended oxidation, it can be  
355 observed that for most asphalt, physical hardening has a stronger impact on grade loss than  
356 oxidation. Especially for asphalt blended with  $C_{20}H_{42}$ , the grade loss gap caused by cold storage  
357 and extended oxidation grows from 3°C to 6°C. Therefore, it is necessary to study the effect of  
358 asphalt wax content on the degree of thermoreversible aging of heavily oxidized asphalt.

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