

1 **Can Soy Methyl Esters Improve Concrete Pavement Joint Durability?**

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31 ABSTRACT

32 While many concrete pavements provide excellent long-term performance, some
33 pavements (primarily in the Midwest) have shown premature deterioration at the joints. This
34 premature deterioration is a concern since it can shorten the life of a pavement that is functioning
35 well otherwise. Previous work has hypothesized that these joints may be susceptible to
36 preferential fluid saturation which can lead to freeze-thaw damage or chemical degradation. This
37 work examines the use of soy methyl ester polystyrene blends (SME-PS) as a method to reduce
38 the rate of fluid ingress into the pore system of the concrete, thereby making the concrete more
39 resistant to deterioration. SME-PS are derived from soy beans and have demonstrated an ability
40 to reduce fluid absorption in concrete when SME-PS is used as a topical treatment. A series of
41 experiments were developed to evaluate the effectiveness of various dosage rates of SME-PS for
42 increasing concrete durability at pavement joints. Experiments show that SME-PS reduces fluid
43 ingress, reduces salt ingress and reduces the potential for freeze-thaw damage. As a result of the
44 positive experimental results, the INDOT is currently conducting field trials that use SME-PS on
45 concrete pavements that are beginning to show signs of premature deterioration with the
46 expectation that SME-PS will extend the life of the joint, thereby reducing maintenance costs
47 and extending the life of the concrete pavement.

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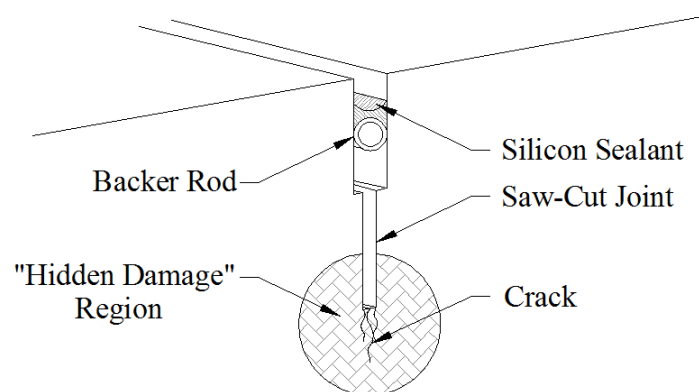
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51 **1.0 BACKGROUND ON JOINTS IN CONCRETE PAVEMENTS**

52 The premature deterioration of concrete pavement joints is a concern since it
53 compromises the performance and potential service life of an otherwise healthy pavement. This
54 damage manifests itself as either the development of cracking parallel to the joint or cracking
55 from a softened region that develops along one side of the bottom of the saw cut to the surface of
56 pavement approximately 4 to 6 inches (100 to 150 mm) from the joint. During field inspections it
57 has been observed that where the joints have shown signs of damage the joint sealant is also
58 damaged and the joint contained standing fluid (the fluid contains water and deicing salts).

59 **Figure 1** illustrates a typical D-1 pavement construction joint that is used in the state of
60 Indiana. A three step process is used to create the joint. First, shortly after the pavement is cast,
61 a saw-cut is placed in the pavement to 1/3 of the pavement depth to control random cracking (1,
62 2). This joint typically causes a crack to form directly below the saw cut. After the first saw-cut
63 is placed, a second cut is placed (approximately 1 inch (25 mm) deep and 1/2 inch (12 mm)
64 wide) to widen the saw-cut near the surface. By allowing the pavement to crack and open before
65 the second cut is placed the joint width can be made more uniform from panel to panel creating a
66 more stable geometry for the joint sealant. The third step involves placing a backer rod in the
67 joint to keep the sealant at the top of the joint. A joint sealant is placed on top of the backer rod
68 and the joint sealant adheres to both sides of the widened saw cut. The sealant is thought to be
69 necessary by some DOT's to keep water and incompressible materials out of the joint; however
70 other DOT's have used unsealed joints (3-8). Silicon is a typical material for sealing joints.



71

72 **FIGURE 1 Typical D-1 contraction joint (9).**

73

74 If the joint sealant remains intact, it will keep water and solutions containing deicing salts
75 out of the joint. However, this sealant frequently becomes damaged and the geometry of

76 pavement joints provides a place for fluid to collect. This fluid will be absorbed by the concrete
77 and can lead to saturation which can result in freeze thaw damage (10). Ironically, while the
78 damaged joint sealant may not prevent fluid from getting into the crack, it will reduce the
79 potential for water evaporation from the joint thereby increasing the potential for saturation.

80

81 **2.0 JOINT SEALANT VS CONCRETE SEALANT**

82 While the previous section describes the conventional concrete joint and sealer used in
83 Indiana, this work also considers the potential for using topical application with a fluid (concrete
84 sealer) that penetrate the pores of the concrete. In this work a distinction will be made between
85 the joint sealant (like a silicon shown in [Figure 1](#)) which forms a physical barrier on the surface
86 of the concrete, with concrete sealants (referred to as topical applications) where a fluid is
87 absorbed into the pore system creates a hydrophobic or blocking layer which inhibits the ingress
88 of water. It should be noted that unlike the joint sealants, the penetrating topical applications do
89 not have the ability to keep incompressible objects out of the joints.

90

91 **3.0 EXPERIMENTAL PROGRAM OVERVIEW**

92 Recent research has shown the potential for using plant-oil based products as concrete
93 topical applications (11-13). One of these products is Soy Methyl Ester (SME). SME also has the
94 potential to be both an effective and environmentally friendly topical treatment for concrete.
95 Previous research has shown that when concrete is treated with SME or Soy Methyl Ester-
96 Polystyrene Blends (SME-PS) the water absorption is reduced (14). The focus of this research is
97 evaluating the effectiveness of SME-PS at increasing the durability of concrete, especially in a
98 freeze-thaw environment. This work examines the influence of SME-PS for reducing water
99 absorption, freeze-thaw durability, and the ingress of chlorides ions. To provide a point of
100 reference the SME-PS is compared with other topical applications for concrete that may be used
101 in this type of application. An overview of the testing program can be seen in [Table 1](#) along with
102 the relevant ASTM standards.

103

104

TABLE 1 Testing Program Overview

Test Description	Test Methods	Mixtures*	Topical Application (Concrete Sealant)
Influence of PS on SME Penetration	Volumetric and Visual Observation	C40 & M42	SME SME-PS (M_w 2,400) SME-PS (M_w 44,000) SME-PS (M_w 382,100)
Water Absorption	ASTM C1585	M42	SME – PS (2 doses**) Solvent Based Silane Water Based Silane
Concrete Freeze-Thaw Durability	ASTM C666 (Procedure A)	M45	SME – PS (2 doses**) Solvent Based Silane
Durability of Sealers under Freezing and Thawing Conditions	ASTM C1585 after ASTM C666	C40	SME – PS (2 doses**) Solvent Based Silane Water Based Silane
Chloride Ingress	Visual Observation using $AgNO_3$	M42	SME – PS (2 doses**) Solvent Based Silane Water Based Silane

* Mixture proportions defined in Table 2

** Varying dosages of SME-PS were applied utilizing different SME-PS exposure times

105

106 **3.1 Mixture Proportions**

107 One concrete mixture and two mortar mixtures were used in this study. The concrete was
 108 prepared with a $w/c=0.40$ with 28 % fine aggregate and 38 % coarse aggregate by volume). The
 109 mortars were prepared with 55 % fine aggregate by volume and with water-to-cement ratios
 110 (w/c) of 0.42 and 0.45. A complete list of mixture proportions can be found in [Table 2](#) details on
 111 the constituent materials are available in [\(13\)](#).

112

113

TABLE 2 Mixture Proportions in SSD Condition

Material	C40	M42	M45
Cement (kg/m ³)	316	609	586
Fly Ash – Class C (kg/m ³)	60	-	-
Water (kg/m ³)	150	256	264
Fine Aggregate (kg/m ³)	736	1444	1444
Coarse Aggregate (kg/m ³)	1049	-	-
Air Entraining Admix. (ml/100 kg cem.)	20	-	-
Retarder Admix. (ml/100 kg cem.)	98	-	-
HRWRA (g/100g cem)	0.5	-	-

114

115 **3.2 Topical Treatments for Concrete (Penetrating Sealers)**

116 During this testing program, samples were tested both with and without topical
 117 treatments. Three different topical treatments were tested. The first is Soy Methyl Ester (SME)
 118 blended with 5 % polystyrene (PS) by mass. The results from the SME-PS treated samples were
 119 compared to two commercially available silane sealants. The first is a solvent-based
 120 alkyalkoxysilane sealer (SBS) with greater than 50 % active ingredients. This sealant is a
 121 solution of silane dissolved in an isopropanol solvent. The second commercially available sealer
 122 is a water-based alkyalkoxysilane penetrating sealer (WBS) that consists of 40 % silane. This
 123 sealant is an emulsion of silane in water. The goal of evaluating the SME-PS along with other
 124 topical treatments is to provide a point of reference as this research is early in the understanding
 125 of how SME-PS performs. It should be noted that no attempt was made to optimize any of the
 126 topical treatment applications in this study and more work is likely needed in this area to test
 127 additional topic treatments and additional application approaches to provide the DOTs with a
 128 variety of approaches that can improve pavement joint performance if this concept is successful.

129 **4.0 PENETRATION OF SME-PS INTO CONCRETE**

130 It is important to understand how SME-PS behaves when it is applied to the concrete.
 131 The first step in this study was determining the influence of various factors on the penetrability
 132 of SME. Specifically being studied are the influence of concrete moisture, size of the polystyrene
 133 molecules, and time the SME is allowed to penetrate into the concrete.

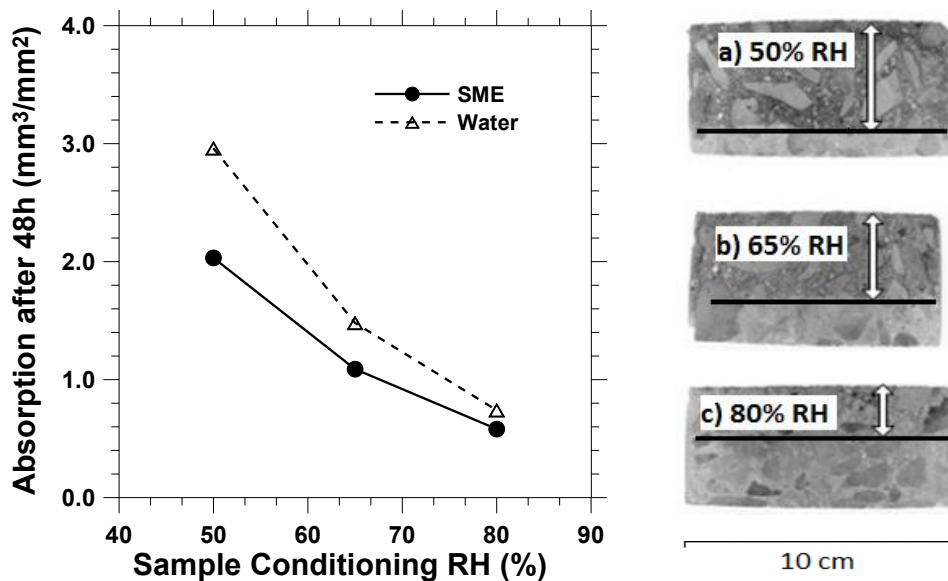
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135 4.1 Influence of Concrete Moisture on SME Penetration

136 The influence of concrete moisture on SME penetration was tested on concrete samples.
 137 A series of 100 mm × 200 mm cylinders were cast of mixture C40. After 24 hours the cylinders
 138 were demolded and sealed in double plastic bags at 23 ± 0.5 °C until the samples reached an age
 139 of 28 d. After 28 d of curing the cylinders were removed from bags and three 50 mm ± 2 mm
 140 thick samples were cut from the central portion of each cylinder with a wet saw. After cutting,
 141 samples were placed in environmental chambers at 23 ± 0.5 °C and at three different relative
 142 humidities (50 ± 1 %, 65 ± 1 % and 80 ± 1 %) for 18 months before testing.

143 After the 18 months, the sides of the sample were sealed with epoxy. A plastic mold was
 144 placed around the top of the sample to prepare a dam. The edge between the plastic mold and the
 145 concrete sample was sealed using silicone, which was allowed to dry for 24 h. The mass of the
 146 samples was then recorded. Approximately 20 g of SME sealant were placed in the dam. After
 147 48 h, the additional sealant was removed from the dam and the mass of the sample was recorded.
 148 [Figure 2](#) shows both the absorption of sealant or water after 48 h. Three samples were used for
 149 each condition. The samples were then cut vertically using a wet saw, for a direct visual
 150 measurement of the sealant penetration. It can be observed that the moisture content of the
 151 samples has a high influence in the penetration of the sealant. Samples conditioned to a lower
 152 relative humidity have more open pore space where water was lost during drying.

153



154 **FIGURE 2 Volumetric and Visual observation after 48 h of SME penetration (from top)**
 155 **into concretes with different levels of saturation. SME penetration is highlighted in black.**
 156 **Sample cross-section is 5 cm by 10 cm.**

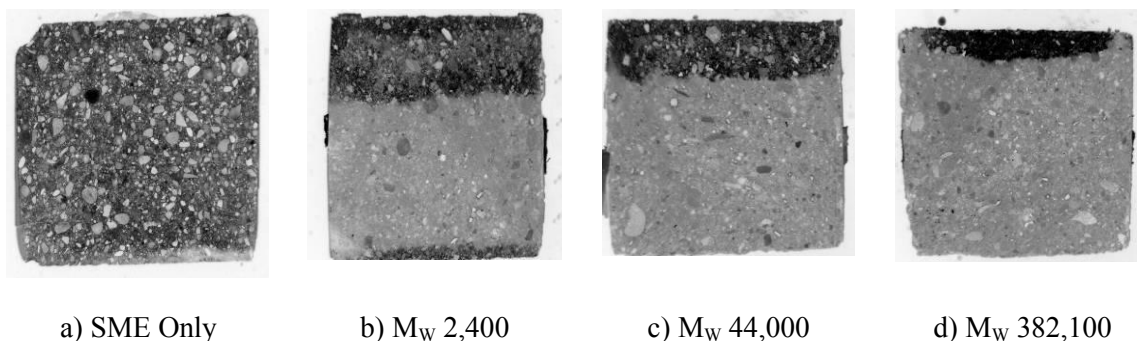
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158 4.2 Influence of Polystyrene Chain Length on SME Penetration

159 Although SME appears to have positive performance by itself, its high solvent capacity
 160 enables it to be blended with polymers such as polystyrene (11). By blending the SME with PS
 161 (or other materials), it may be possible to tailor the properties of the solution for a specific need
 162 (14). Another benefit of using a SME-PS blend is that the waste polystyrene, which otherwise
 163 would be put in a landfill, can be used to extend the material and may remain in the pores as a
 164 'blocking material'. If waste PS is to be used, it is important to understand how different sources
 165 of polymer would affect the final product. In this study, the impact of the polystyrene (PS) chain
 166 length on sealant penetration was tested by varying the molecular weight of the polystyrene.

167 Penetration of SME sealants blended with PS of different size polymer chains was tested
 168 on mixture M42, which were cast on 35 mm x 300 mm cylinder molds, where they were kept
 169 until the age of 28 d. Then, samples were demolded and 50 mm \pm 2 mm thick samples were cut
 170 from the central portion of each cylinder with a wet saw. After cutting, samples were conditioned
 171 by placing them in environmental chambers at 23 \pm 0.5 $^{\circ}$ C and 65 \pm 1 % relative humidity for 8
 172 months to allow them to equilibrate before testing.

173 After the 8 months, the sides of the sample were sealed with epoxy. After the epoxy had
 174 hardened, aluminum tape was placed around the sample to prepare a dam. The edge between the
 175 aluminum tape and the mortar samples was sealed using silicone, which was allowed to dry for
 176 24 h. The mass of the samples was then recorded. 5 g of sealant was placed on the dam and
 177 allowed to absorb. In this experiment four SME-PS blends were tested: No PS (pure SME), M_w
 178 2,400 PS, M_w 44,000 PS, and M_w 382,100 PS. Each of these molecular weights corresponds to
 179 the chain length of the polystyrene molecules. Each of the SME-PS blends was prepared with 5
 180 % PS by mass. After 48 h, the sealant was carefully removed and samples were cut vertically
 181 using a wet saw, for a direct visual measurement of the sealant penetration (Figure 3). Results
 182 show that the presence of PS influenced the ability of the sealer to penetrate the samples. As the
 183 molecular weight of the PS increases the penetration depth of the sealer decreases.



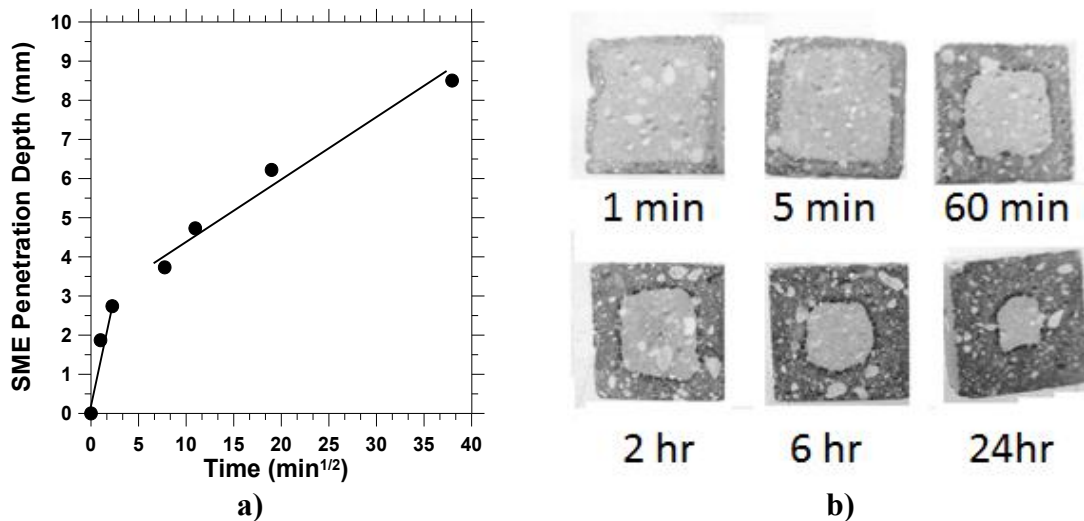
184 **FIGURE 3 Penetration depth of SME-PS blends highlighted in black (penetrating from**
 185 **top surface only). Each sample was treated with a SME-PS blend prepared with**
 186 **polystyrene of various chain lengths. Samples are 35 mm by 50 mm in cross-section.**

187

188 **4.3 Influence of Time on SME Penetration**

189 This test investigated the effect of role of time on the penetration depth of SME-PS. To
 190 perform this study, six 2.5 cm x 2.5 cm x 10 cm prisms of mixture M42 were prepared. These
 191 samples were demolded at 24 h and allowed to dry at $23 \pm 1^\circ\text{C}$ and $50 \pm 2\%$ RH for seven days
 192 prior to being submerged in SME-PS containing a fluorescent dye (0.5% by mass). At certain
 193 time intervals the samples were removed from the solution. These samples were cut with a small
 194 wet saw and photographed under an ultraviolet light to determine the penetration depth of the
 195 SME-PS. Both images of the SME-PS penetration and a graphical display of penetration depth
 196 over time can be seen in Figure 4. The penetration depth shows a square root of time
 197 dependence with two rates, an initial faster absorption rate which appears to slow over time.

198



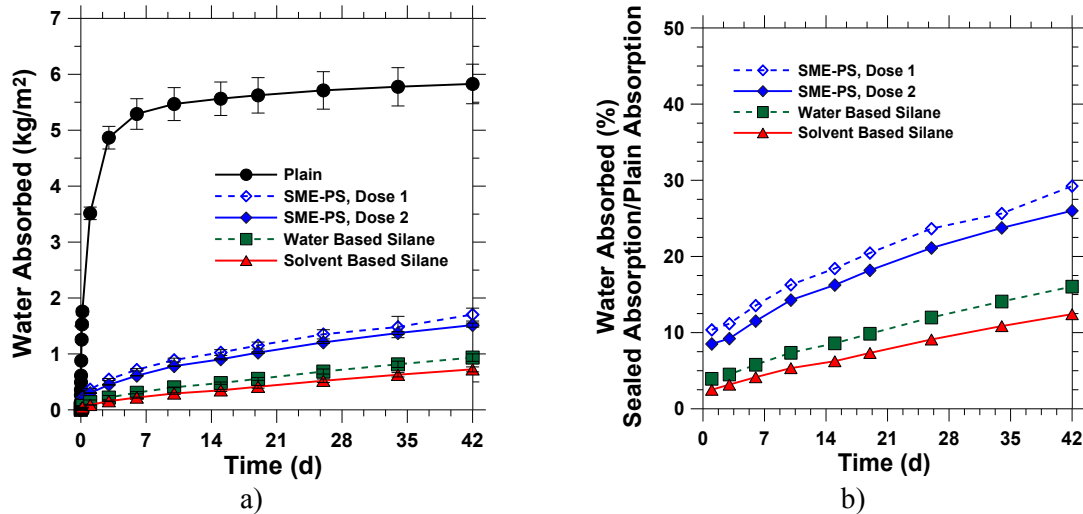
199 **FIGURE 4 Penetration of SME into 2.5 cm square samples: a) penetration depth vs. time,**
 200 **b) SME-PS highlighted in black as a function of time.**

201 **5.0 WATER ABSORPTION**

202 The ability of the topical treatment to minimize water absorption into concrete was
 203 measured using ASTM C1585 (15). The topical treatments being tested are SME-PS (2 dosages),
 204 SBS, and WBS. Samples were prepared as 100 mm x 200 mm cylinders of mixture M42.
 205 Samples were demolded at 24 h and sealed in a double plastic bag for 14 d. At this time each
 206 cylinder was cut into three 5 cm tall discs. The top and bottom 2.5 cm of the cylinder was
 207 discarded. The cylinders were allowed to condition at $23 \pm 1^\circ\text{C}$ and $50 \pm 2\%$ RH for 6 month
 208 prior to the application of the sealants. SME-PS was applied to cylinders by submerging the
 209 exposed face in the SME-PS for 6h (dose 1) and 24 h (dose 2). The SBS and WBS were applied

210 evenly with a paint brush on the exposed surface. Results of the water sorption test can be found
 211 in Figure 5.

212



213 **FIGURE 5 Water absorption experimental results: a) water absorbed (Error bars**
 214 **represent +/-1 standard deviation), b) reduction in water absorption. (Average of three**
 215 **samples)**

216

217 The application of each of the topical treatments tested resulted in a substantial reduction
 218 in the amount of water absorbed into the samples. Both silane sealants were more effective at
 219 reducing water absorption than the SME-PS.

220

221 6.0 FREEZE-THAW DURABILITY

222 When concrete contains water the concrete can be susceptible to reaching a critical level
 223 of saturation which can lead to freeze-thaw damage (10, 16). This potential for degradation may
 224 be increased when the water contains aggressive ions such as chlorides (17). A common method
 225 for improving the freeze-thaw durability of concrete is through the use of air entraining
 226 admixtures (AEA) (16). One alternative method for improving freeze/thaw durability of concrete
 227 may be to use topical treatments which would help to reduce the potential for saturation by
 228 reducing the rate of water penetration (Figure 5). It should be noted that a sealer only protects
 229 the concrete from fluid ingress from the surface, however it does not prevent possible water
 230 coming from under the slab.

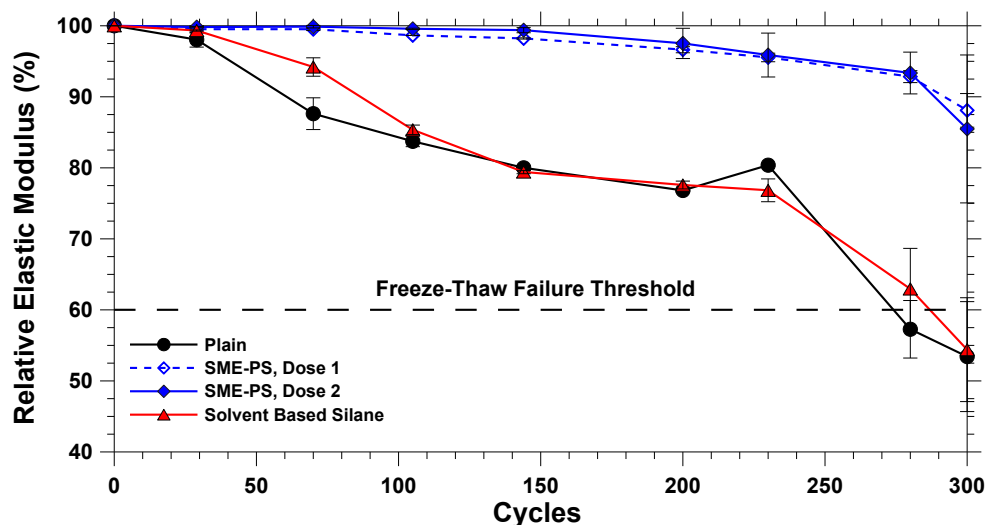
231 6.1 Cold Weather Behavior of SME-PS

232 SME is a solution of different fatty acid methyl esters (FAME) which begin to lose their
 233 solubility and come out of solution at a critical temperature. This results in a second solid phase
 234 forming in the solution. For SME these solids are conglomerations of waxy crystals that will give

235 the solution a cloudy appearance. The temperature at which this first occurs is known as the
 236 cloud point. For SME, this value is typically accepted as 0 °C (18, 19). The cloud point of SME-
 237 PS with 5 % and 10 % PS was measured to be 5 °C according to ASTM D2500 (20). As the
 238 temperature continues to drop below the cloud point, more of the FAMEs will come out of
 239 solution and precipitate into waxy solids. Eventually there will be enough solidification that the
 240 solution will lose its ability to flow like a liquid. At this point, known as the pour point, SME
 241 becomes a gel-like substance. This temperature is typically accepted as -4°C (19). These cold
 242 flow properties of SME are a major limitation for using these materials as an alternative fuel
 243 source (18) since the waxy solidifications in the SME will clog filters within the fuel system. In
 244 terms of concrete, the solidified SME would clog the pores of the concrete, presumably further
 245 reducing fluid ingress.

246 6.2 Concrete Freeze-Thaw Durability

247 The freeze-thaw of the topical treatments was tested using ASTM C666A (freezing and
 248 thawing in water) (21). Samples were prepared from mixture M45. The samples were demolded
 249 after 24 h and sealed in double plastic bags for 60 d. After this time the samples were opened and
 250 allowed to dry at 23 °C and 50 % RH for 10 d. At this time sealants were applied. The SME-PS
 251 was applied by submerging the samples in the sealant for 6 h (dose 1) and 24 h (dose 2). The
 252 solvent based silane was applied in an even coating with a paint brush. Prior to starting testing,
 253 sealants were allowed to cure for 7 d and then samples were submerged in saturated lime water
 254 for 48 h. The relative elastic modulus can be seen in Figure 6.

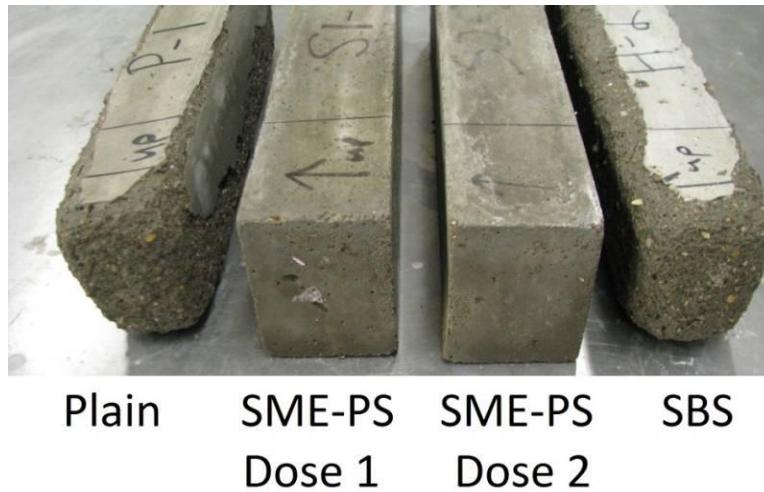


255
256 **FIGURE 6 Freeze-Thaw experimental results, relative elastic modulus.**

257
258 Use of the solvent based silane sealant slowed the effects of freeze-thaw damage during
 259 the first 100 cycles. Prior to 100 cycles, samples treated with the SBS received 50 % less damage
 260 than untreated samples. After 100 cycles the both the SBS treated and untreated samples were at

261 the same level of damage. After 75 cycles the cover of the untreated and SBS began to spall and
 262 its mass was reduced (Figure 7). The use of SME-PS sealant was less susceptible to damage
 263 from freezing and thawing. After 300 cycles of freezing and thawing the untreated and SBS
 264 treated samples had a relative modulus of approximately 55 % (below the ASTM C666 limit of
 265 60 %) while the SME-PS samples had a relative modulus of 85 %.

266



267

268 **FIGURE 7 Freeze-Thaw samples after 280 cycles. Untreated (plain) samples and SBS**
 269 **treated samples show significant cover loss. SME-PS treated samples show no visual**
 270 **damage.**

271

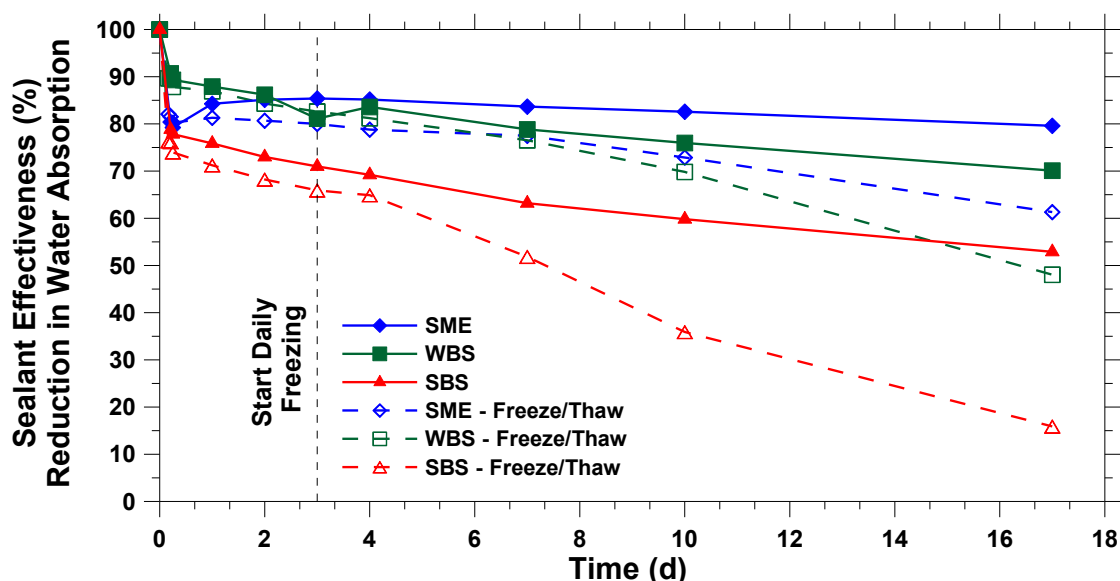
272 6.3 Sealant Freeze-Thaw Durability

273 Concrete samples treated with the solvent based silane sealant showed a reduction in
 274 water ingress of approximately 95 % (14). However, when these same materials were tested for
 275 freeze-thaw durability (21), the solvent based silane sealant did not show the same benefits. This
 276 section begins to explore the reasons for this difference in behavior.

277 To begin it should be noted that the silane sealants used in this study are not 100 % active
 278 ingredients. The silane was transported by an inactive carrier. The silane in SBS was dissolved in
 279 isopropanol. The WBS was an emulsion of silane and water. As the sealant cures, the carrier
 280 fluid evaporates leaving behind the silane forming a protective membrane on the surface of the
 281 concrete. As the membrane is exposed to freezing temperatures it is susceptible to thermal
 282 contraction. If the forces become large enough, the membrane could begin to crack and loss
 283 effectiveness. In contrast the SME never “cures” or hardens. When SME is applied to concrete, it
 284 is absorbed into the pores and remains fluid. Any thermal contraction would not result in the
 285 development of tensile stresses in the sealant.

286 In order to test this hypothesis, mixture C40 was treated with the sealants and exposed to
 287 freezing and thawing cycles. After freeze-thaw testing water absorption was evaluated using
 288 ASTM C1585 (15). For this series of experiments the samples were conditioned at 23 ± 1 °C
 289 and $50 \pm 2\%$ RH for 12 months prior to the application of the sealants. The sealants were applied
 290 in the same manner as described in the water absorption. Prior to testing, the samples labeled
 291 “Freeze/Thaw” were exposed to 7 freezing-thawing cycles between 5 °C and -18 °C. After 3
 292 days of testing these samples are exposed to a daily freezing cycle (without being removed from
 293 the water). The efficiency of the sealants, reduction in water absorption compared to untreated
 294 samples, can be seen in Figure 8.

295



296

297

FIGURE 8 Sealant effectiveness after freezing and thawing.

298

299 Slight differences (up to 5 %) were observed in the absorption behavior between the
 300 samples that underwent freezing and those that were kept at 23 °C within the first three days of
 301 testing. From this it can be implied that none of the sealants tested were vulnerable to freeze-
 302 thaw damage when the concrete was dry. After three days of testing, samples were exposed to
 303 daily freeze-thaw cycles (12 hours at -18 °C and 12 hours at 5 °C) while in water. Following the
 304 additional 7 freeze-thaw cycles all of the sealants tested lost some of their efficiency. The most
 305 vulnerable sealant type was the SBS, which absorbed 85% of the volume of water absorbed by
 306 untreated samples following freezing. The least vulnerable sealant was the SME-PS. It only lost
 307 about 20 % efficiency after freezing and thawing. Please note that the WBS only lost 23 % of its
 308 effectiveness during this testing. It is believed that the emulsifying agents left in the concrete
 309 allowed for some movement of silane as the concrete was being damaged. This will lead initial
 310 effectiveness of the sealant under these conditions.

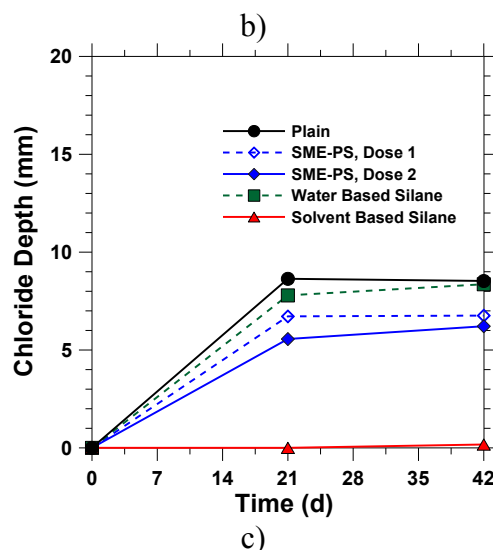
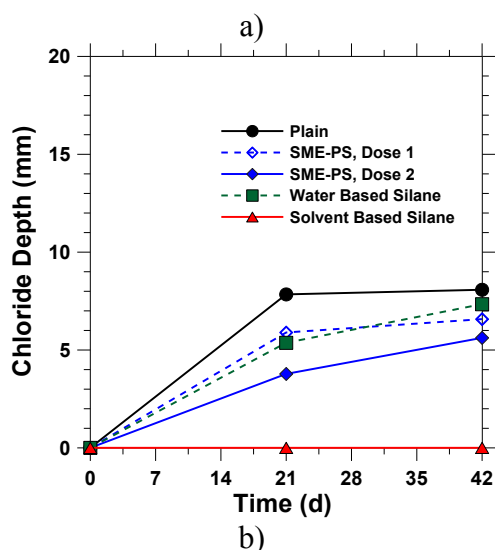
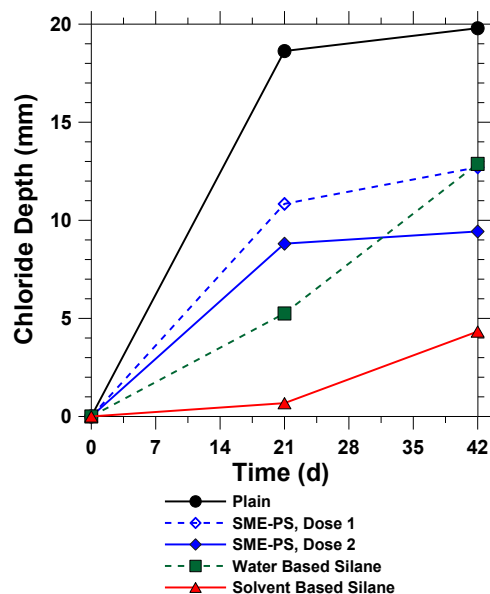
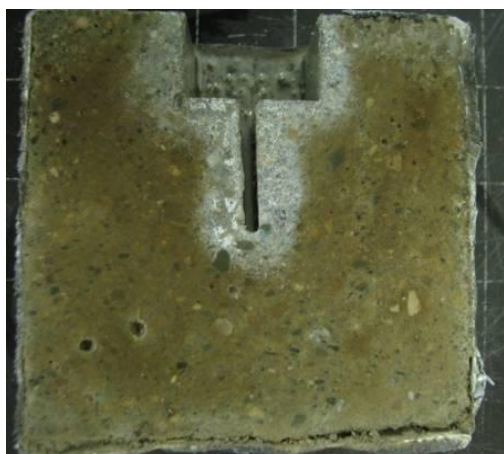
311 7.0 CHLORIDE INGRESS

312 In addition to studying water penetration, this research evaluated the potential ingress of
313 chloride ions that may come from deicing salts. Saw-cut joint specimens of mixture M42 were
314 created. These specimens, shown in 10, were 10 cm x 10 cm x 10 cm mortar cubes. On the top
315 face of this cube a 2.5 cm x 2.5 cm well was cast in place (Figure 9a). 24 hour after casting, the
316 specimens were demolded and a 25 mm deep saw cut was cut into the middle of the well. At this
317 time the specimens were sealed for 14 days and then allowed to dry in a 50 % RH chamber for
318 an additional 90 days. The sides of the specimen were sealed to form a reservoir to hold the salt
319 solutions. This specimen geometry was selected as it attempts to mimic the behavior of a saw cut
320 joint in the field (11).

321 After the construction of the epoxy dams, the samples were treated with the topical
322 sealants. The topical treatments being tested are SME-PS (2 dosages), Solvent Based Silane, and
323 Water Based Silane. For SME-PS dose 1, SBS, and WBS the sealant was pooled in the reservoir
324 for 6 h. For SME-PS dose 2, the material was pooled for 24 h. The sealants were allowed to cure
325 for 7 d prior to the addition of the salt solutions.

326 For this series of tests three salt solutions were used. The three solutions are all
327 commercially available de-icing solutions of 23 % Sodium chloride (NaCl), 30 % Magnesium
328 chloride (MgCl₂), and 32 % Calcium chloride (CaCl₂). These salts solutions were pooled in the
329 reservoir for 21 d and 42 d. At this time the samples were saw cut and 0.1M Silver nitrate
330 (AgNO₃) was applied to samples. A white precipitate (AgCl) forms on the sample in the
331 presence of chloride ions. This reaction will occur when the chloride ion concentration is over
332 165 ppm (22). These samples were photographed and the images were analyzed to determine the
333 chloride penetration depth using image analysis (13). The depth of chloride ingress for this
334 experiment can be seen in Figure 9 b) to d).

335



336 **FIGURE 9 Chloride ingress depth: a) typical chloride ingress sample (100 mm x 100 mm)**
 337 **sprayed with silver nitrate, b) sodium chloride (23% by mass), c) magnesium chloride**
 338 **(30% by mass) , and d) calcium chloride (32% by mass).**

339
 340 The SBS sealant was the most effective of the topical treatments at reducing chloride
 341 ingress completely eliminating any ingress of chlorides for the $MgCl_2$ and $CaCl_2$ solutions and
 342 reduced the penetration depth by 80 % at 42 d for the $NaCl$ solution. When using SME-PS, the
 343 dosage rate of sealant was directly related to its effectiveness. The samples were treated with a
 344 larger dosage of SME-PS resulted in a reduction of chloride depth about 10 % greater than that
 345 of the smaller dosage. The least effective sealant for preventing chloride ingress was the WBS.
 346 After 42 d of ponding, the chloride penetration depth was reduced by 35 %, 10 %, and 0 % for
 347 the $NaCl$, $MgCl_2$ and $CaCl_2$ solutions respectively.

348 **8.0 CONCLUSIONS**

349 This paper has reported results of experiments to evaluate the use of Soy Methyl Ester
350 Polystyrene (SME-PS) as a topical treatment for concrete pavement. Based in the results of
351 these experiments it is hypothesized that SME-PS may increase the durability of concrete
352 pavement joints. It was observed that the penetration depth of SME-PS is dependent on concrete
353 moisture level, size of PS molecules, and time. As the concrete moisture level increases, the
354 amount of SME-PS that can be absorbed will decrease. As the chain length of PS increases, the
355 amount of SME-PS absorbed will decrease. SME-PS reduced damage from freezing and
356 thawing. After 300 cycles, the untreated samples had a relative elastic modulus of 55 %
357 compared to the 85 % of the SME-PS treated samples. Field trials have begun to evaluate the
358 performance of using SME-PS in pavement joints during new construction as well as in areas
359 where the concrete has begun to show signs of premature deterioration.

360

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369 **10.0 REFERENCES**

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