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# Effects of different ageing methods on the chemical and

## rheological properties of bitumen

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Abstract: Bitumen undergoes ageing, which leads to changes in its chemical and rheological properties, thus 1 2 becoming harder and more brittle in time. This study aims to compare the effects of different laboratory 3 ageing methods on chemistry and rheology of three bitumen types, i.e. a Pen 40/60, a Pen 70/100 and a 4 polymer modified bitumen (PmB). Four ageing protocols were applied: ageing at room temperature, oven 5 ageing, the Pressure Ageing Vessel (PAV), and the Rolling Thin Film Oven Test (RTFOT) combined with 6 PAV ageing. The effects of temperature, pressure, and ageing time were studied using dynamic shear tests 7 and infrared spectroscopy. The results highlight the relationship between chemistry and rheology of bitumen. 8 Bitumen hardening, which was revealed by an increase in complex modulus and a decrease in phase angle, 9 was reflected in the growth of specific chemical functional groups. Among all materials, soft bitumen 10 showed the greater tendency to oxidize. Different behaviour was observed for PmB that presented the 11 highest resistance against oxidation among the studied bitumens, even though the reaction with oxygen 12 caused the deterioration of the added polymer modifiers.

13 Author keywords: Bitumen; Ageing; Chemistry; Rheology; FTIR

16 Introduction

Numerous studies have investigated the sensitivity of asphalt mixtures to the damaging effects of climatic conditions and traffic loading. Especially, the ageing processes that occur during the service life of asphalt pavements lead to obtain mixtures that are more brittle in time and therefore, prone to ravelling and cracking (Molenaar et al. 2010). Various ageing mechanisms are identified in literature, namely physical hardening, loss of volatile components and oxidation. Among all, oxidation is considered to be the most important ageing process that can alter the chemical and rheological properties of bitumen.

The complex mechanical response of pavements is governed by the properties of asphalt mastics (Dondi et al. 2014) and essentially, it depends on the filler properties and on the way they interact with bitumen (Sangiorgi et al. 2016; Vignali et al. 2014; Mazzotta et al. 2016). However, bitumen is an organic material, as it mainly consists of carbon and hydrogen atoms (Lesueur 2009), which reacts with the oxygen present in the atmosphere and undergoes ageing. Actually, it is the ageing susceptibility of bitumen that leads to pavement damage (Lopes et al. 2012).

29 Oxidative ageing is a diffusion-driven phenomenon, which occurs due to photo-oxidation and 30 thermal reaction between bitumen components and atmospheric oxygen (Lu et al. 2008). Oxygen diffuses 31 into bitumen, changes the chemical features of bitumen and consequently, affects its physical properties. In 32 general, the diffusion phenomenon is driven by the internal heat energy and is influenced by several 33 parameters. The average radius of the diffusing molecules and the viscosity of the diffusion medium affect 34 the rate of oxygen diffusion. Furthermore, intermolecular interactions, polarity and temperature influence 35 indirectly this phenomenon (Karlsson and Isacsson 2003; Herrington 2012). Also, the temporal changes in 36 the concentrations of oxygen and reactive bitumen species during the ageing reaction can affect the reaction 37 itself (Herrington 2012).

Corbett (1969), as reported in Lesueur (2009), separated bitumen in four fractions: saturates aromatics, resins and asphaltenes. Each fraction is a continuum of molecules with different size, polarity and aromaticity. Redelius and Soenen (2015) reported that the largest size of bitumen macromolecules is defined by the crude oil source, while its smallest size is determined by the refinery process. Polar molecules influence the elastic behaviour of material by creating a network. On the contrary, non-polar fractions contribute to the viscous response of bitumen (Lerfald 2000). The reaction of bitumen and atmospheric
oxygen leads to the growth of polar and aromatic molecules, which are able to reduce the mobility of
molecules to flow; hence bitumen undergoes hardening and embrittlement.

In literature, several studies investigated the changes of bitumen fractions when subjected to ageing. 46 It has been found that aromatics generate resin fractions, which in turn generate asphaltenes. On the other 47 48 hand, the concentration of saturates changes slightly due to their low chemical reactivity (Lesueur 2009). The 49 oxidation of bitumen fractions generates a number of functional groups such as ketones, sulfoxides, 50 dicarboxylic anhydrides, and carboxylic acids. Ketones are formed from specific aromatic molecules, which 51 have two adjacent benzylic carbon moieties. When those molecules are oxidized, first, they result in the 52 formation of dicarboxylic anhydride and then, ketones are made by condensation. Carboxylic acids are 53 produced in small amounts. On the other hand, sulfoxides are the result of organic sulphides oxidation, 54 which represent functional moieties of many types of asphalt (Petersen 2009).

55 The types of oxidation products are consistently the same for bitumen from different sources, but 56 their amount may differ especially between unmodified and polymer modified bitumen (Lu et al. 2008). In 57 reality, the concentration of each product and their tendency to oxidize are influenced by crude oil origin. As 58 sulphides are more reactive, sulfoxides have higher production rate and are formed faster than ketones. After 59 that, ketones are formed, which finally yield anhydrides and carboxylic acids. Lesueur (2009) stated that 60 sulfoxides are thermally unstable and they reach a constant level, which depends on initial sulphur content 61 and oxygen diffusion into the material. In contrast, ketones and carboxylic acids are more stable, but they do not always reach an asymptotic value. Also, Lesueur (2009) indicated that field aged bitumen usually 62 63 achieves a constant value, while an asymptotic value is not reached when bitumen is laboratory aged. On the other hand, Liu et al. (1996) showed that the rate of carbonyl formation becomes constant for all laboratory 64 65 aged bitumen after an initial higher rate period. More recently, Lu et al. (2008) confirmed the increase of carbonyls and sulfoxides molecules after standard short- and long-term ageing protocols, namely after 66 67 Rolling Thin Film Oven Test (RTFOT) and Pressure Ageing Vessel (PAV) ageing. They compared the formation of these functional groups for unmodified and Styrene-Butadiene-Styrene (SBS) modified bitumen 68 69 through infrared spectroscopy. Sulfoxides were observed to grow more in the neat standard bitumen.

Conversely, a greater presence of carbonyl was found for the modified material. The researchers attributed
this increase to both oxidation and degradation of polymers.

72 The rheological and physical characteristics of bitumen are strongly related to the chemical ones. Ageing causes stiffening of the bitumen as shown by an increase of complex modulus and a decrease of 73 74 phase angle (Lu et al. 2008; Molenaar et al. 2010). Moreover, oxidation increases bitumen viscosity, which 75 depends on the type and strength of molecular interactions. Petersen (2009) reported that bitumens with high 76 content of metals, especially vanadium, show major sensitivity to viscosity. In addition, the values of 77 softening point and penetration grade change after oxidation. Lu et al. (2008) reported that the softening point increases for natural bitumen, while the opposite trend was found for polymer modified materials. On 78 79 the contrary, the penetration value decreases with increasing ageing time.

80

#### 81 **Research objectives**

This study aims to compare the effects of different laboratory ageing methods on the chemistry and rheology of bitumen. The influence of temperature, pressure, and ageing time were studied using the Dynamic Shear Rheometer (DSR) tests and Fourier Transform Infrared Spectroscopy (FTIR). Also, the properties of aged bitumens were compared with those of the unaged fresh materials to study the susceptibility of the different bitumen types to ageing.

87

#### 88 Experimental research

To study the effects of different laboratory ageing methods on bitumen chemistry and rheology four ageing protocols were applied: ageing at room temperature, oven ageing, Pressure Ageing Vessel (PAV) and Rolling Thin Film Oven Test (RTFOT) combined with PAV ageing. The effects of temperature, pressure, and ageing time on three bitumens types were studied by means of DSR and FTIR tests. Two replicate samples of each bitumen type were tested at every ageing condition to determine statistically reliable results. In all the cases analysed, similar results were obtained.

95

#### 96 Materials and sample preparation

97 Three bitumen types were adopted throughout this study, two standard or neat bitumens and one polymer 98 modified bitumen. The standard materials varied with respect to their penetration grade; a Q8 Pen 40/60 99 bitumen and a Q8 Pen 70/100 bitumen were used. The Q8 polymer modified bitumen with a grade QmB 100 45/80 - 65 was obtained by the addition of Styrene-Butadiene-Styrene (SBS) polymers, which are often used 101 to enhance the durability and resistance of bitumen (Dondi et al. 2016). Table 1 lists the properties of the 102 materials.

103 Cylindrical bitumen samples were prepared according to the European Standard NEN-EN 104 12594:2014. The diameter of the samples was equal to 27.50 mm and their thickness was 2 mm. Bitumen, in 105 quantity lower than one liter, was heated in the oven for 120 minutes at temperature that did not exceed the 106 softening point of each specific bitumen type more than 100°C. The liquid bitumen was stirred and poured in 107 a silicon mould to cool down. Then, the mould was placed in a refrigerator for about 15 minutes. The 108 samples were finally demoulded and were ready for ageing in the lab.

109

#### 110 Ageing methods

Four ageing methods were applied, namely ageing at room temperature, oven ageing, ageing by means of the Pressure Ageing Vessel, and an ageing protocol that included the combination of Rolling Thin Film Oven Test and PAV ageing.

114 Ageing at room temperature was performed at five different time intervals of 5, 10, 15, 20 and 25 115 days. The average room temperature was 24.6°C. During room ageing the samples were not homogenized, as 116 this would require the application of heat that could affect the results. Oven ageing was carried out at 135°C 117 for 60 hours, and 5, 10, 15 days. A standard quantity of bitumen ( $90\pm3$  g) was heated and poured into 118 aluminium containers forming a 1 cm thick layer. During ageing, bitumen was manually stirred, rotating the 119 trays clockwise and counter-clockwise once a day in order to encourage the homogeneous ageing of the samples. The Pressure Ageing Vessel test was performed at standard conditions, i.e. ageing at 100°C 120 temperature and 2.1 MPa pressure for 20 hours, following the NEN-EN 14769:2005 European Standard. 121 122 Moreover, PAV test was realized at the same temperature and pressure but for a double ageing time (40 123 hours). Finally, the short- and long-term standard ageing procedures were combined; first the samples were

aged using the RTFOT according to the NEN-EN 12607-1:2014 European Standard followed by PAV atstandard conditions. The aforementioned ageing methods are listed in Table 2.

126

#### 127 Test methods

The effects of ageing were evaluated by means Dynamic Shear Rheometer tests and Fourier Transform 128 129 Infrared Spectrometry. The Perkin Elmer Spectrum 100 FTIR spectrometer was used in the Attenuated Total 130 Reflectance (ATR) mode to identify the chemical functional groups of the bitumens. The wavelength of the 131 reflected beam is characteristic of each element and indicates the presence of specific functional groups. Using the ATR technique, sample preparation was simple. A small sample of material was taken using a 132 spatula and placed under the FTIR anvil. The FTIR spectrum was obtained in the spectral range between 133 134 4,000 and 600 cm<sup>-1</sup> with a scanning resolution of 4 cm<sup>-1</sup> averaging five scans for each measurement in order 135 to increase the signal-to-noise ratio of the measurements. Then the spectrum was normalized to allow the 136 quantitative analysis of the results.

137 The oxidation of bitumen was evaluated at specific bands of wavelengths, which can reveal the 138 formation of the characteristic oxidation products. Researchers have identified four specific wavelengths 139 where the changes due to ageing can be observed. Due to the complexity of the chemical composition of 140 bitumen a fixed peak cannot be found for bitumens especially if the origin of materials is different (Van der 141 Bergh 2011). For this reason, in this study, the effects of ageing were analyzed considering specific bands of 142 wavelength, as defined by Lamontagne et al (2001), and the corresponding area under those bands (Van der 143 Bergh 2011). The peak areas were evaluated using semi-quantitative analysis and then the four ageing 144 indices, namely the aromaticity, long chain, carbonyl and sulfoxide were determined, as presented in Table 145 3.

Bitumen rheology was characterized through DSR tests. The Anton Paar EC-Twist 502 device was used to analyse the material response over a wide range of temperatures and frequencies that occur during the pavement service conditions. First, the Linear Viscoelastic Range (LVE) of each bitumen type was determined by means of amplitude sweep tests. The maximum strain value within the linear viscoelastic range was 0.9% for Pen 40/60, 1.4% for Pen 70/100 and 1.0% for PmB. These values were chosen to be the strain amplitude values applied during the frequency sweep tests for each bitumen. The tests were performed using the parallel plate configuration in a strain controlled mode, applying an oscillatory shear stress on a sample of 1 mm thickness. The tests were performed at a range of frequencies from 10 Hz to 0.1 Hz and at five temperatures, specifically at 10°C, 20°C, 30°C, 40°C and 50°C.

155

#### 156 **Results and discussion**

#### 157 Fourier Transform Infrared Spectroscopy analysis

158 The FTIR spectra were obtained in a wavelength range from 4,000 to 600 cm<sup>-1</sup>. However, the changes due to 159 ageing can be found at wavelengths less than 1800 cm<sup>-1</sup>, as shown in Fig. 1.

160 Fig. 2 shows the aromaticity and long chain indices for all bitumen types after the application of the various ageing methods. The aromaticity index denotes the aromatization of bitumen and relates to the 161 162 increase of viscosity (Nivitha et al. 2016). The results show that the aromaticity index does not vary 163 significantly after ageing at room temperature for none of the three bitumen types. In contrast, oven ageing seems to affect the aromatic fraction. The value of the aromaticity index increases in time for the polymer 164 165 modified bitumen, especially after five days of oven ageing. For the neat bitumens, the aromatic fraction 166 shows an increase after five days of oven ageing similar to the PmB, but then remains relatively constant. 167 The standard PAV method does not influence significantly the value of the aromaticity index compared to its value before ageing. The application of the extended PAV protocol results in an increase of the aromaticity 168 169 index similar to this of prolonged oven ageing.

Nivitha et al. (2016) observed a decrease of the long chain index with increasing ageing time.
Similarly, Fig. 2 shows that the long chain index decreases with extended oven ageing for all bitumens.
Depending on the bitumen type, the index seems to decrease with ageing time (for PmB and Pen 70/100) or
reaches a steady state after five days of oven ageing (Pen 40/60). The evolution of the long chain index does
not show a specific trend with time when aged at room temperature.

On the contrary, Fig. 3 shows that the values of the carbonyl and sulfoxide indices increase during ageing; the extent of increase depends on the ageing method applied to the bitumen samples. The carbonyl index denotes the formation of ketones, dicarboxylic anhydrides and carboxylic acids due to the reaction between carbon and oxygen (C=O), whereas the sulfoxide index denotes the S=O stretching due to the 179 oxidation of sulphides. Both indices are identified as key parameters when studying bitumen ageing, as the180 carbonyl groups and sulfoxides are the most important oxidation products.

181 It is interesting to note that the carbonyl index has zero value after room ageing, regardless of the 182 duration of ageing and the bitumen type. On the other hand, ageing in the oven at 135°C causes an increase 183 of the carbonyl index. The duration of oven ageing influenced the formation of carbonyl compounds, as 184 indicated by the growth of the carbonyl index. After 60 hours of oven ageing, the carbonyl index of both neat 185 bitumens remains zero. After 5 days of oven ageing, a slight increase is observed for the harder Pen 40/60 186 bitumen, whereas a higher value is obtained for the Pen 70/100. However, after 10 days of oven ageing the 187 value of the carbonyl index increases substantially for both bitumen types and becomes one order of 188 magnitude higher than its value at five days. Overall, the formation rate of carbonyl compounds is higher for 189 Pen 70/100 as indicated by the absolute values of the carbonyl index, which are one order of magnitude 190 higher than those of the harder bitumen. The formation of carbonyl compounds for the PmB occurs after 10 191 days of oven ageing, unlike for the unmodified bitumens. The delay in carbonyl formation can be attributed 192 to the inherently low susceptibility of polymer modified bitumens to higher temperatures.

The production of sulfoxides does not follow the same trend. Sulfoxides are formed to some extent when the samples are aged at room temperature, indicating that the natural content of sulphur reacts with atmospheric oxygen even at lower temperatures. It is, therefore, shown that the sulfoxides (S=O) have a higher production rate than the carbonyls. This finding verifies the higher rate production of sulfoxides than ketones reported by Lesueur (2009).

198 Sulfoxides in Pen 70/100 bitumen are also formed at a higher rate compared to Pen 40/60 as 199 indicated by the sulfoxide index. The value of the sulfoxide index for Pen 70/100 changed by 33%-40% after 200 ageing at room temperature and 52-62% after oven ageing compared to its value at unaged conditions. In 201 contrast, the sulfoxide index for Pen 40/60 increased only by 12% after room ageing and 11-16% after oven 202 ageing compared to the reference unaged value. These findings indicate that the soft bitumen is more 203 susceptible to oxidative ageing than the harder Pen 40/60 bitumen. For the unmodified bitumens, the 204 application of the standard PAV method, combined or not with the standard RTFOT method, results in the 205 growth of both the sulfoxide and carbonyl index. From a comparison between the O60h protocol and the 206 PAVS protocol, it can be observed that the application of pressures higher than the atmospheric pressure

207 promotes the formation of carbonyl compounds, even though the PAVS ageing was performed for less time 208 and at lower temperature than O60h ageing after which no carbonyls were produced.

209 On the other hand, the sulfoxide index for the PmB did not vary considerably with ageing time at room temperature, whereas oven ageing caused an increase in its value after 5 days of ageing. The 210 211 application of high temperature and pressure (PAVS, PAVM and RTFOT+PAVS ageing methods) affected 212 the growth of sulfoxides, but did not increase significantly the amount of carbonyls when using the two 213 standard ageing protocols. It was observed that the application of PAVS and RTFOT+PAVS ageing resulted 214 in a minor increase of the carbonyl index for the PmB, while the extended PAVM protocol had a significant 215 effect on the carbonyl index, possibly due to the prolonged ageing time. The presence of styrene and 216 butadiene copolymers in the PmB is detected at 700 cm<sup>-1</sup> and 968 cm<sup>-1</sup> wavelength, respectively (BRRC 217 2013). The polymer peaks after the application of each ageing method are listed in Table 3. The results show 218 that the styrene copolymer degrades with ageing, especially after oven ageing and the PAV ageing protocols.

219 Among the various ageing methods the PAVM and the prolonged oven ageing were the most 220 effective ageing protocols, as indicated by the changes in the carbonyl and sulfoxide indices. Both the 221 carbonyl and sulfoxide indices were found to increase during ageing. However, it has been observed that the 222 formation of carbonyl compounds was mainly the result of applying high temperature. It is therefore 223 suggested that the carbonyl index and the relative increase of the sulfoxide index (compared to a reference 224 unaged value) are considered together when characterizing ageing in bituminous mixtures; especially in the 225 case of bitumen recovered from asphalt pavements, where such high temperatures do not usually take place. 226 using only the carbonyl index as ageing indicator may lead to incorrect conclusions with regard to the 227 bitumen susceptibility to ageing. The graph in Fig. 4 compares the development of oxidation products after three ageing methods, namely after 25 days of room ageing, 15 days of oven ageing and the application of 228 229 PAVM ageing. The formation of oxidation products varies with the ageing method for all bitumens. 230 Nevertheless, the results confirm the greater sensitivity of Pen 70/100 bitumen to oxidize.

231

#### 232 Dynamic shear rheometer test results

Oxidative ageing is known to alter the rheological properties of bitumens, causing an increase in complex
 modulus and a decrease in phase angle. Every bitumen type responds in a different way to ageing, and

therefore the changes in physical properties can be different, mainly due to its dependency on oxidation
kinetics (Juristyarini et al. 2011). Fig. 5 and Fig. 6 show the master curves of Pen 40/60 and Pen 70/100
bitumens, respectively, at unaged state and after the application of the R25d, O15d, RTFTO+PAVS and
PAVM ageing protocols.

239 It can be observed that the most noticeable changes in complex modulus due to ageing were recorded 240 at low frequencies as the bitumen was in the rubbery flow region, whereas minor changes were observed at 241 high frequencies (glassy transition region). Similar to the results obtained from the FTIR measurements, it 242 was observed that oven ageing, at 135°C for 15 days, affected the rheology of bitumens more than the combined effects of high temperature and pressure. Again, ageing at room temperature for 25 days did not 243 244 result in significant differences with respect to the rheological response of fresh (unaged) bitumen. It seems 245 that oxidation took place, but in a very slow rate, confirming that bitumen reacts slowly with atmospheric 246 oxygen (Herrington 2012). The phase angle of the unmodified binders decreased considerably with ageing, 247 indicating an increase in elasticity of bitumen and a reduction of its flow characteristics. Oven ageing, in 248 particular, caused a great increase in phase angle especially at low frequencies (high temperatures), which in 249 combination to the increase of complex modulus, is an indication of embrittlement of the material.

Fig. 7 shows the complex modulus and phase angle master curves for the PmB. The complex modulus of PmB increased with ageing, but to a less extent if compared to the unmodified bitumens. It was observed that after 15 days oven ageing the complex modulus, at low frequencies, increased by one order of magnitude. Overall, the rheological response of the PmB after oven ageing did not differ significantly with its response after the application of the PAVM and PAVS+RTFOT protocols, in contrast to what was observed for the unmodified binders. This possibly occurs due to the high temperature susceptibility of the polymer modified bitumen, which can undergo higher temperatures for prolonged ageing times.

The phase angle master curve at unaged condition shows a characteristic plateau in the intermediate frequency area. This feature actually shows the presence of polymer into the bitumen chemical structure. This characteristic plateau in the phase angle master curves is maintained after 25 days of room temperature ageing. However, the application of high temperature for prolonged time (O15d) or combinations of high temperature and pressure (PAVM and RTFOT+PAVS) caused a smoothening of the curve, as shown in Fig. 10. This result may be explained by the fact that degradation of the polymer modifier can occur after the specific ageing protocols. The FTIR measurements support this hypothesis as it was observed that the styrene copolymer peak degraded with ageing, especially after the application of the O15d and PAVM and RTFOT+PAVS protocols. In general, based on the rheological properties, the PmB was found to have greater resistance to the oxidation processes, while Pen 70/100 bitumen was highly sensitive to ageing.

267

#### 268 Conclusions

This study aimed to compare the effects of different laboratory ageing methods on the chemical and rheological properties of unmodified and SBS modified bitumens. The evolution of the chemical properties during ageing was monitored using the indices that correspond to aromaticity and long chains, as well as to the formation of carbonyls and sulfoxides. Moreover, the characteristic peaks of the modifiers were considered. The changes in rheology were evaluated on the basis of complex modulus and phase angle master curves.

275 The influence of high temperature was clearly observed for unmodified bitumens when compared to 276 the polymer modified bitumen in terms of both thermo-sensitivity reduction (Vignali et al. 2016) and ageing 277 response. The ageing indexes and the rheological response of both neat bitumen types, Pen 40/60 and Pen 278 70/100, demonstrate that high temperatures applied for a prolonged time ( $\geq 10$  days) influence more the 279 bitumen response than the combined effects of high temperature and pressure due to PAV and RTFOT. For 280 the polymer modified bitumen, though, oven ageing at high temperature did not result in large differences in 281 its rheological response when compared to the response after the application of the PAVM and 282 PAVS+RTFOT protocols, possibly due to the high temperature susceptibility of the polymer modified 283 bitumen. Nevertheless, the characteristic plateau in the phase angle master curve at unaged condition, 284 denoting the presence of polymer modifiers, was smoothened out after the application of high temperature for prolonged time or combinations of high temperature and pressure. It was hypothesized that the 285 286 aforementioned ageing conditions degrade the polymer modifier; the evolution of the characteristic FTIR 287 peaks of the modifiers confirmed this hypothesis.

The aromaticity index and the long chains index did not vary significantly after ageing at room temperatures; nevertheless, the application of high temperatures and pressure resulted in a decrease of the long chain index and an increase of the aromaticity index. In some cases, no specific trend was observed with respect to ageing time or temperature. In contrast, both the carbonyl and sulfoxide indices increased with ageing. Carbonyls formed at a slower rate than the sulfoxides. Ageing at room temperature did not stimulate carbonyl formation for any of the bitumens; the carbonyl index remained equal to zero. After 60 hrs of oven ageing at high temperature for the unmodified bitumens (or five days for the polymer modified bitumen) carbonyl compounds were formed, as indicated by the growth of the carbonyl index.

Similarly, ageing at room temperature did not influence the rheological properties of the bitumens. Overall, bitumen hardening, which was revealed by an increase in complex modulus and a decrease in phase angle, was reflected in the growth of specific chemical functional groups. The polymer modified bitumen showed a highest resistance against the oxidation, thus against stiffening and embrittlement, than neat standard bitumens. Among the unmodified bitumens, the soft Pen 70/100 bitumen exhibited the greater tendency to stiffen compared to the harder Pen 40/60 bitumen.

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### **Table 1.** Bitumen Specifications

Property	Unit	Pen 40/60	Pen 70/100	PmB
Penetration @25°C	mm	40/60	70/100	45/80
Softening Point	°C	49	44	65
Fraass Breaking Point	°C	-7	-14	-15
Dynamic viscosity @60°C	Pa∙s	175	160	-
Complex shear modulus 1.59Hz - @60°C	kPa	3.6	1.8	12
Phase angle 1.6Hz - @60°C	0	87	88	69

## **Table 2**. Applied Ageing Methods

Ageing method	Ageing time	Abbreviation
Fresh (unaged)	-	F
Room @24.6°C	5 days	R5d
	10 days	R10d
	15 days	R15d
	20 days	R20d
	25 days	R25d
Oven @135°C	60 hours	O60h
	5 days	O5d
	10 days	O10d
	15 days	O15d
PAV @100°C & 2.1MPa	20 hours	PAVS
	40 hours	PAVM
RTFOT+PAV	Standard conditions	RTFOT+PAVS

#### **Table 3.** Indices calculated for all bitumens

Long chain A	A <sub>734-710</sub> /A <sub>1376</sub> +A <sub>1460</sub>	
Carbonyl		
Curoonyi	A <sub>1753-1660</sub> / $\sum$ A	
Sulfoxide A	A <sub>1047-995</sub> /\_A	
$\sum A = A_{(2953,2862)} + A_{1700} + A_{1600} + A_{1460} + A_{1376} + A_{1030} + A_{864} + A_{814} + A_{743} + A_{724}$		

Ageing method	Styrene peak [700 cm-1]	Butadiene peak [968 cm-1]
fresh (unaged)	0.3436	0.2102
R5d	0.2995	0.1954
R10d	0.3069	0.2067
R15d	0.3001	0.1862
R20d	0.2726	0.1859
R25d	0.2877	0.1890
O60h	0.3219	0.2137
O5d	0.3156	0.2173
O10d	0.3006	0.2147
O15d	0.3085	0.2150
PAVS	0.3097	0.2179
PAVM	0.3036	0.2233
RTFOT+PAVS	0.3251	0.2334

**Table 4.** Polymer Peaks of Polymer Modified Bitumen obtained by FTIR

















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Fig. 1. Detail of the FTIR spectra.

**Fig. 2.** Aromaticity and long chain indices for all bitumen types after the application of the different ageing methods.

Fig. 3. Carbonyl and sulfoxide indices for all bitumen types.

Fig. 4. Growth of C=O and S=O oxidation products of laboratory aged bitumens.

Fig. 5. Master curves of complex modulus and phase angle of Pen 40/60 at different ageing conditions.

Fig. 6. Master curves of complex modulus and phase angle of Pen 70/100 at different ageing conditions.

Fig. 7. Master curves of complex modulus and phase angle of PmB at different ageing conditions.