

1 *Research Article*

## 2 **Butyl rubber-based composite: Thermal degradation** 3 **and prediction of service life time**

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14 **Abstract:** Butyl rubber-based composite (BRC) is one of the most popular materials for the  
15 fabrication of protective glove against chemical and mechanical risks. However, in many working  
16 places such as metal manufacturing or automotive mechanical services, its mechanical hazards  
17 usually appear together with metalworking fluids (MWFs). The presence of these contaminants,  
18 particularly at high temperature, could modify its properties due to the scission, the plasticization,  
19 the crosslinking of polymer network and thus led to severe modification of mechanical and  
20 physicochemical properties of material. This work aims to determine the effect of temperature and  
21 a metalworking fluid on mechanical behavior of butyl rubber composite dealing with crosslinking  
22 density, cohesion forces and elastic constant of BRC on the based on Mooney-Rivlin's theory. The  
23 effect of temperature with and without MWFs on thermo dynamical properties and morphology of  
24 butyl membranes is also investigated. The prediction of service lifetime is then evaluated from  
25 extrapolation of Arrhenius plot at different temperatures.

26 **Keywords:** Polymer aging, swelling behavior, protective glove materials, elastomers, composites,  
27 rubber

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### 30 **1. Introduction**

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The swelling characteristics of rubbers are related to the deterioration of mechanical properties and may affect to their failure mechanics. Nowadays, there are no models to predict the failure

45 mechanisms in the presence of industrial contaminants. Several investigations have been reported on  
 46 the effect of industrial contaminants on morphology, mechanical properties and structure of gloves  
 47 material. In previous works [15-24], we reported that polymers and composites [16-20, 22, 23, 25-41]  
 48 can be degraded by several aging agents such as heat, water, humidity, oxygen, ozone and various  
 49 other environments factors [16-24]. The scission, crosslinking and plasticization of polymer network  
 50 may be occurred and led to severe deterioration of mechanical and physicochemical properties of  
 51 material [42], which are quite important as regards their applications in various fields of mechanical  
 52 engineering.

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54 Many researchers have been attempted to understand the relationship between morphological  
 55 and property changes with microstructure of rubber material[43], especially with failure mechanism  
 56 of elastomer by using several traditional test methods including tensile test, strain-stress relation, and  
 57 compression measurements [44-46]. Mooney-Rivlin's equation is usually used to evaluate the  
 58 property-structure relationship of material by analysing of tensile stress-strain curves of materials. In  
 59 this model, Rivlin [47, 48] proposed an extension of the Mooney's model by describing  $W$  as a  
 60 polynomial series in  $(I_1 - 3)$  and  $(I_2 - 3)$ :

$$61 \quad W = \sum_{i,j=0}^{\infty} C_{ij} (I_1 - 3)^i (I_2 - 3)^j \quad \text{with } C_{00} = 0 \quad \text{Eq.1}$$

62 During a simple tensile test, we are:

$$63 \quad I_1 = \lambda^2 + \frac{2}{\lambda} \quad I_2 = \frac{1}{\lambda^2} + 2\lambda \quad I_3 = 1 \quad \text{Eq.2}$$

64 The stress  $\sigma_n$  is given by the equation:

$$65 \quad \sigma_n = 2 \left( \lambda - \frac{1}{\lambda^2} \right) \left[ \frac{\partial W}{\partial I_1} + \frac{1}{\lambda} \frac{\partial W}{\partial I_2} \right] \quad \text{Eq.3}$$

66 According to the behavior law, we can still write:

$$67 \quad \sigma_n = \lambda \frac{\partial W}{\partial \lambda_1} + \frac{1}{\lambda} \frac{\partial W}{\partial \lambda_2} \quad \text{Eq.4}$$

68 From which

$$69 \quad \sigma_n = 2 \left( C_1 + \frac{C_2}{\lambda} \right) \left( \lambda - \frac{1}{\lambda^2} \right) \quad \text{Eq.5}$$

70 where  $W$  is the density of deformation energy,  $\lambda$  is elongation of material and  $C_1$  et  $C_2$  are two elastic  
 71 constants associated with network structure and the flexibility of network, respectively. According  
 72 to Eq.5, plot of  $\sigma_n / 2 \left( \lambda - 1/\lambda^2 \right)$  as a function of  $\lambda^{-1}$  should be a linear and the  $C_1$  constant is directly  
 73 calculated by extrapolating of the linear portion of the curve up to  $\lambda^{-1} = 0$  and the  $C_2$  constant is the  
 74 curve slope. In the molecular aspect, this constant  $C_1$  of Mooney-Rivlin's model characterizes by the  
 75 density of crosslinking between elastomer chains [49, 50]. It can be used to evaluate the physical  
 76 manifestation of the degree of crosslinking  $N$  [51, 52].

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$$N = \frac{2C_1}{kT} \quad \text{Eq.6}$$

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The constant  $C_2$  is associated with the cohesion forces (intramolecular bonds and interchain) and reinforcement-matrix interactions[52]. This constant also reflects the flexibility of the structure of the material[53]. It depends on the degree of vulcanization which increases with the increasing of the crosslink density[53]. The values of two constants  $C_1$  and  $C_2$  that can be determined during the tensile tests allow for studying the influence of the crosslinking degree and network structure on the stiffness of material. Gordon [52] is determined the Young's modulus according to  $C_1$  and  $C_2$ :

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$$E = 6(C_1 + C_2) \quad \text{Eq.7}$$

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It reported that the constant  $C_1$  usually remains constant with variation of the swelling degree of polymer material, while the constant  $C_2$  decreases with the increase of swelling degree[54-56]. It lacks a systematically research on the effect of temperature and metalworking on morphology, properties and structure of elastomer materials.

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In this study, a butyl rubber membrane, representing a common material used in protective gloves, has been used to determinate the swelling characteristics to metalworking fluid at various temperatures ranging from 80 °C to 120 °C. The elastic behaviors dealing with elastic constants, crosslinking density and modulus were continuously followed during the contamination procedure to evaluate the dominant parameter on polymer deterioration. The modification of morphology, tensile properties and physicochemical properties were also investigated.

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## 2. Materials and Methods

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### 2.1. Materials

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Butyl rubber based composite (BRC) membranes used in this work are made from butyl rubber with carbon black fillers. These commercial products (thickness of ~1.6mm) are kindly supplied by McMaster-Carr Inc. (Canada). An industrial MWF used in this work, namely Milform 64 SST, is directly purchased from a company in metalworking sector in Quebec, Canada. The chemical composition can be found in our previous publication. The carbon black filler content in composite membranes is estimated to be 42- 44 wt.% (by TGA measurement).

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### 2.2. Contamination procedure

The elastomer membranes having the mass ( $m_0$ ) were immersed in to the liquid contaminants for different periods of time for measuring of swelling index changes according to the ASTM D471-12 standard. They are then delicately wiped by thick wiping papers to remove the excess oil on sample surface and reweighed ( $m_t$ ). The process is repeated until saturated state. The swelling index (M) is calculated according to the relation following:

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$$M(\%) = \frac{m_t - m_0}{m_0} \times 100 \quad \text{Eq.8}$$

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where  $m_0$  and  $m_t$  are respectively the weight of the samples before and after immersion in contaminant.

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### 2.3. Scanning electron microscopy (SEM)

114 Scanning electron microscopy model Hitachi S570 was used to investigate the morphology and  
115 the fractography changes of samples before and after contamination process. These observations  
116 serve to examine the failure modes of morphology changes of elastomer materials.

#### 117 2.4. Tensile tests

118 Tensile properties of elastomer membranes are measured with the dog bone rectangular  
119 samples. Measurements were performed with an Alliance 2000 (MTS) universal testing machine  
120 equipped with 1000N load cell and operated at a cross-head speed of 500 mm/min according to the  
121 ASTM D412-06 standard test method. For each condition, five replicates are measured.

#### 122 2.5 Thermal gravimetric analysis (TGA)

123 TGA is carried out with original and aged composite samples to evaluate the percentage of filler  
124 content and the effect of aging process on the sample weight loss as a function of time. The TGA is  
125 carried out in nitrogen atmosphere at a heating rate of 20 °C/min using a Parkin Elmer TGA model  
126 4000. Thermograms were recorded from 0 to 800 °C.

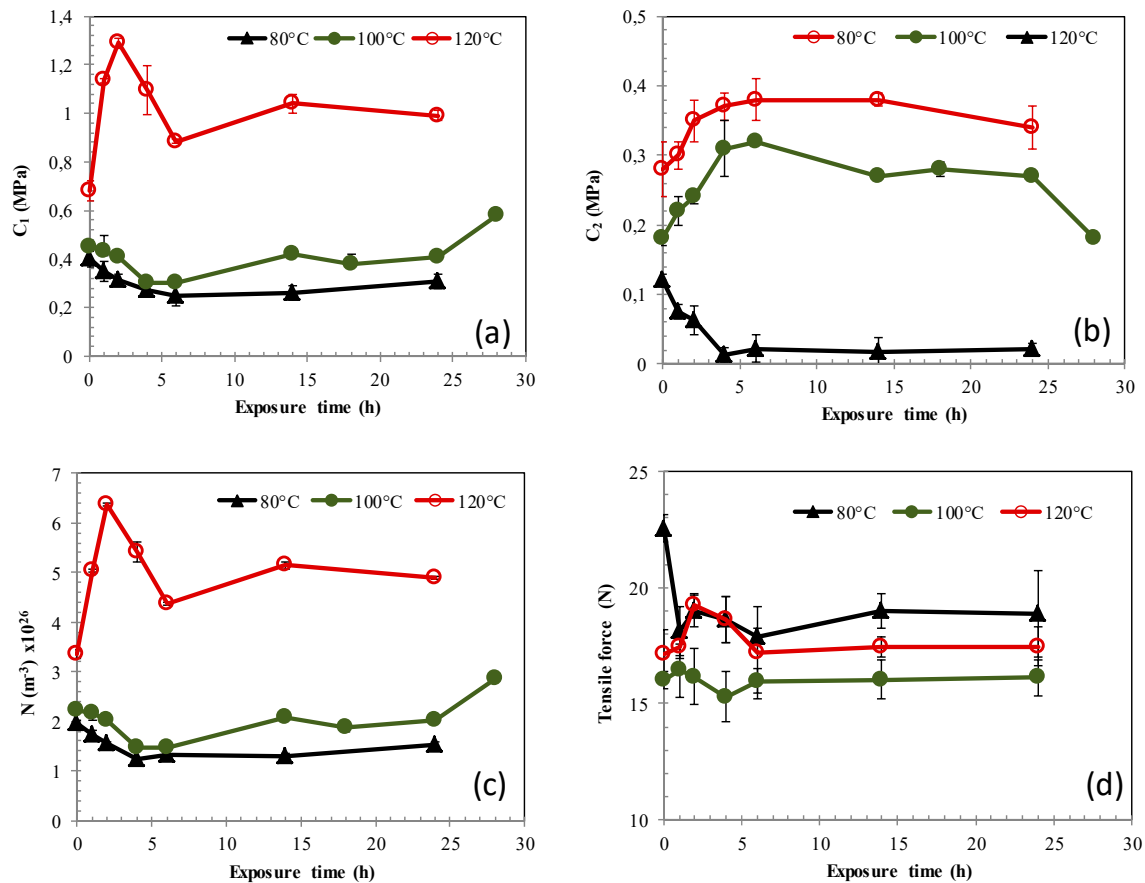
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### 128 3. Results and discussions

#### 129 3.1. Effect of temperature on mechanical properties

130 The influence of temperature on the viscoelastic properties of BRC is investigated at three  
131 different temperatures ranging from 80 and 120°C. Figure 1a shows the relationship between the  
132 constant  $C_1$  and exposure time. It demonstrates that the highest value of constant  $C_1$  is observed at  
133 temperature around of 120°C. At the beginning of exposure ( $t < 4$  h),  $C_1$  undergoes a significant  
134 increase, and then rapidly decreases before achieving a constant value. At lower exposure  
135 temperature (80°C and 100°C), the variation of the constant  $C_1$  is not significant representing by a  
136 slight decrease of  $C_1$  at the first exposure hours then appears to be converged to the similar value to  
137 that observed at 120°C. These results allow confirming that the stiffness of butyl is optionally  
138 increased with temperature, as reported in the case of a neoprene membrane which is exposed to  
139 high temperature [57]. Figure 1b shows the variation of  $C_2$  as a function of exposure time at different  
140 temperatures. The variation of the constant  $C_2$  at 80°C appears to be similar to that at 100°C with  
141 parabolic shape curve. An increase in  $C_2$  at first hours of exposition ( $t < 4$ h) is observed and then it  
142 becomes to slightly decrease with the time. It can be attributed to the increase of cohesive force at the  
143 first hours of exposition time and then decrease [54]. This physical phenomenon is completely  
144 different to that observed at higher exposure temperature (120°C). The constant  $C_2$  decreases  
145 significantly from the first steps of exposure and go down to zero. These results are not surprise and  
146 are in good agreement with that published by Marck [54].

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**Figure 1** Effect of temperature on mechanical properties of BRC as function of MWF exposure time  
 a) constant  $C_1$ ; b) constant  $C_2$ ; c) Crosslinking density ( $N$ ); d) Tensile force.

The synergy between the variation of  $C_1$  and  $C_2$  is also investigated and the results are shown in Table 1. This synergy is determined by calculating the theoretical Young's modulus by using Eq.7 and the experimental Young's modulus from the first linear region of the force-displacement curves (Fig. 1d). The obtained results show that the rigidity of the butyl remains almost unchanged at 80°C. However, a significant increase of Young's modulus with exposure time was recorded at 100°C and 120°C. This can be attributed generally to the increasing of crosslinking density ( $N$ ) that may be occurred at high temperature due to the vulcanization or reticulation of polymer chains. After 24 hrs exposure in MWF, the Young's modulus seems to be increased for all studied temperatures (Fig 1c). Figure 1d demonstrates the tensile force of BRC as a function of exposure time, it can be seen that the rupture properties remain constant during all exposure times whatever the exposure temperature. The variation of tensile strength at break with exposure time is not significant. This implies that the BRC has relative high resistance to temperature for a short time and the later does not affect the mechanical failure behavior.

171 **Table 1:** Young Modulus values of a butyl membrane exposed various temperatures

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Exposure time (h)	Exposure temperature					
	80°C		100°C		120°C	
	$E_{theo}$	$E_{exp}$	$E_{theo}$	$E_{exp}$	$E_{theo}$	$E_{exp}$
0	4.08	3.40	3.78	3.00	4.80	4.11
	(0.42)	(0.04)	(0.06)	(0.05)	(0.30)	(0.16)
1	3.90	2.89	3.93	3.15	6.57	6.10
	(0.36)	(0.2)	(0.48)	(0.25)	(0.12)	(1.10)
2	3.99	3.01	3.90	3.20	8.12	7.43
	(0.30)	(0.09)	(0.10)	(0.13)	0.24)	(0.18)
4	3.78	2.99	3.66	2.88	6.65	5.45
	(0.24)	(0.03)	(0.36)	(0.18)	(0.66)	(0.85)
6	3.78	2.98	3.72	3.01	5.44	4.57
	(0.42)	(0.10)	(0.04)	(0.02)	(0.18)	(0.02)
14	3.84	2.98	4.14	3.27	6.35	5.08
	(0.24)	(0.09)	(0.09)	(0.01)	(0.36)	(0.14)
24	3.90	2.96	4.08	3.30	6.08	4.80
	(0.36)	(0.02)	(0.09)	(0.09)	(0.16)	(0.12)

174 Note: between parentheses: standard deviation (SD)

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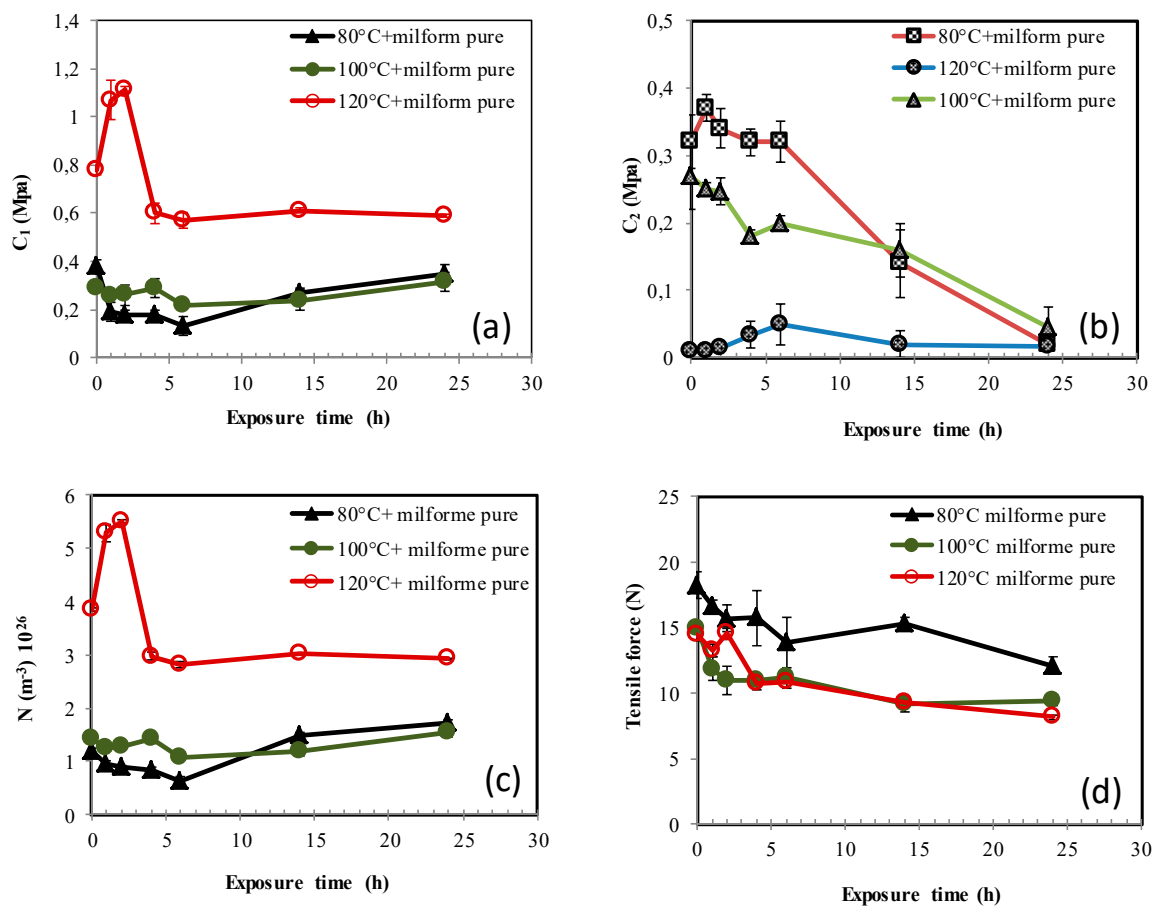
176 *3.2. Effect of simultaneous temperature and metalworking fluid on properties, structure and morphology*

177 In the working condition, glove materials are often in contact with hot MWFs, generated by  
178 friction of machine devices. This factor could be directly affected on the protective performance of  
179 the glove materials. The effect of simultaneous temperature and MWFs on properties of butyl  
180 membranes is investigated. Figure 2a shows the variation of the constant  $C_1$  as a function of expose  
181 temperature in the presence of a metalworking fluid (Milform 64 SST). As shown in Figure 2a, the  
182 changes of the constant  $C_1$  with exposure time at 80°C are substantially similar to that at 100°C, thus  
183 these two curves are approximately superposed. However, at 120°C, the curve tendency is different  
184 when compared to that at lower exposure temperatures, where  $C_1$  values dramatically drop at the  
185 first hours of exposure and then tends to archive a plateau region. Figure 2b shows the effect of  
186 temperature and MWFs on the constant  $C_2$ . It demonstrates that the constant  $C_2$  is rapidly decreased  
187 with the exposure time and approach to zero after 24 hrs of exposition. Therefore, the cohesive  
188 strength of inter-chain eventually became weak [54] and the viscoelastic behavior may be also  
189 affected. The results are in good agreement with that observed by Gumbrell *et al.* [58] and Allen *et*  
190 *al.*[59] in studying the aging process of natural rubber to different solvents.

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192 The calculation of the Young's modulus, in particular at 80 °C and 100 °C, highlights the effect  
193 of simultaneous temperature and contaminants on the rigidity of the material. A significantly  
194 decrease of Young's modulus as a function of exposure time was observed (Table 2). This could be  
195 attributed to the reduction of the cohesive strength of inter-chains. The rigidity of butyl at 120°C is  
196 unstable. The Young's modulus firstly increases, then decreases before achieving a plateau region.

197 This variation is possibly related to the changes in the cross-linking density ( $N$ ) as shown in Figure  
 198 2c. This variation is due to the correlation between the constants  $C_1$  and  $C_2$ . The similar behavior was  
 199 also discussed by Nohilé et al.[4] demonstrating that the swelling of BRC by solvents led to an  
 200 increase of the material rigidity due to the loss of fillers. The later leads to a reduction of the tensile  
 201 strength of BRC as shown in Figures 2d. It can be seen from these figures that the mechanical  
 202 properties of material are decreased whatever exposure temperatures are applied. The longer  
 203 exposure time, the higher loss of mechanical is observed.  
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206 **Figure 2** Effect of temperature and a MWF on mechanical properties of BRC as function of MWF  
 207 exposure time a) constant  $C_1$ ; b) constant  $C_2$ ; c) crosslinking density ( $N$ ); d) tensile force.

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219 **Table 2:** Young Modulus values of BRC membrane exposed at different temperatures in the presence of  
 220 metalworking fluid  
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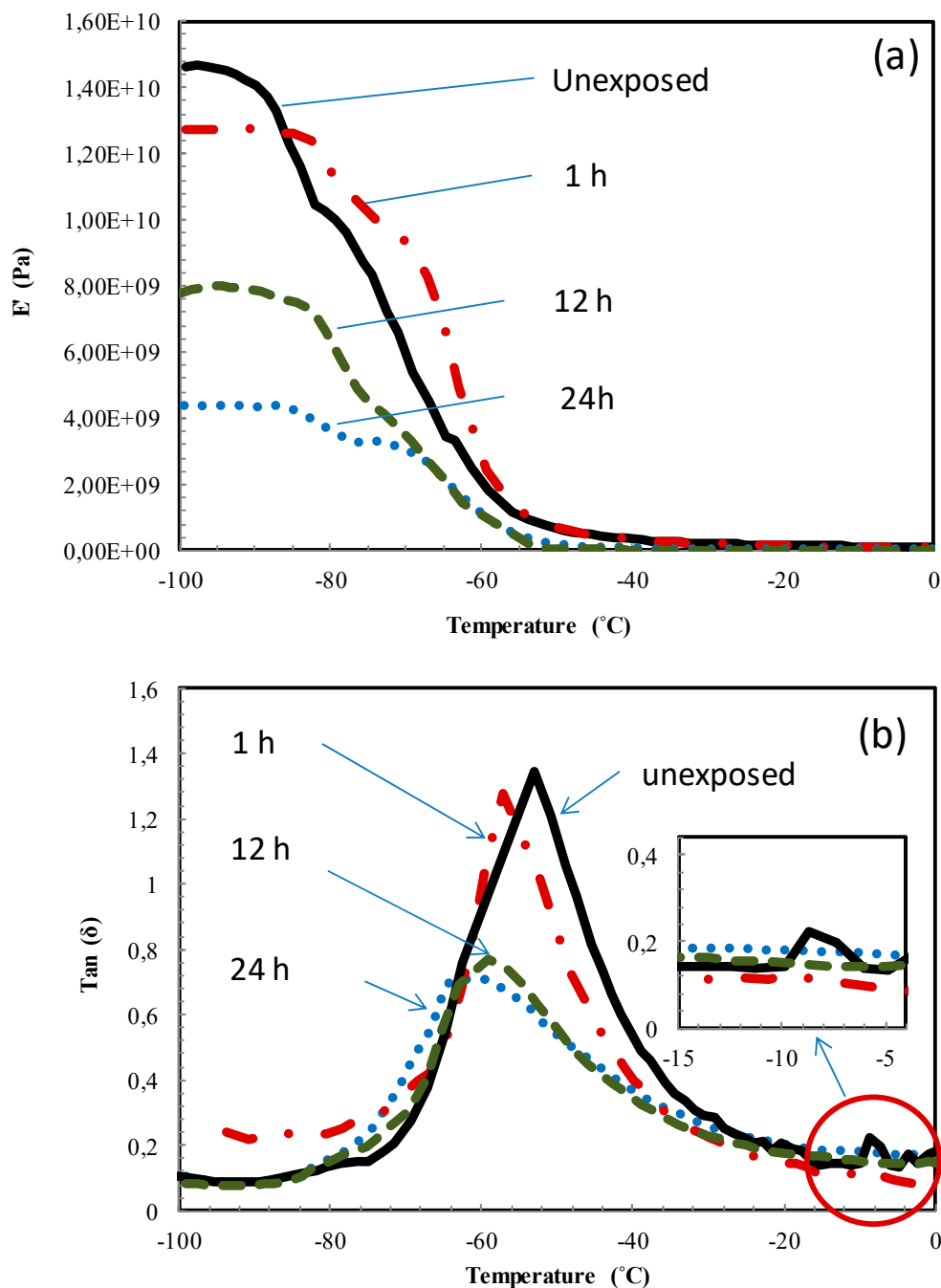
t (h)	Exposure temperatures					
	80°C		100°C		120°C	
	E <sub>theo</sub>	E <sub>exp</sub>	E <sub>theo</sub>	E <sub>exp</sub>	E <sub>theo</sub>	E <sub>exp</sub>
0	3.72 (0.36)	2.87 (0.20)	3.36 (0.36)	2.70 (0.22)	4.74 (0.15)	4.26 (0.28)
1	3.36 (0.36)	2.64 (0.16)	3.03 (0.12)	2.51 (0.09)	6.76 (0.54)	6.28 (0.17)
2	3.12 (0.30)	2.43 (0.08)	3.03 (0.20)	2.52 (0.01)	6.78 (0.09)	6.46 (0.01)
4	3.00 (0.24)	2.41 (0.05)	2.82 (0.30)	2.33 (0.11)	3.804 (0.36)	3.26 (0.18)
6	2.70 (0.42)	2.30 (0.03)	2.52 (0.12)	2.18 (0.03)	3.72 (0.36)	3.41 (0.03)
14	2.46 (0.24)	2.31 (0.05)	2.40 (0.48)	2.01 (0.05)	3.66 (0.18)	3.15 (0.04)
24	2.22 (0.36)	1.98 (0.02)	2.16 (0.42)	1.92 (0.06)	3.64 (0.04)	3.06 (0.05)

Note: between parentheses: standard deviation (SD)

### 222 3.3. Effect of temperature and MWF on dynamical Thermo-mechanical properties

223 Dynamical thermo-mechanical analysis (DMTA) provides access to the elastic modulus and  
 224 tangent values of materials. Figure 3a shows the example of Young modulus of BRC membrane  
 225 before and after aging at 100 °C during 24 h in MWF (Milform 64SST). The curves are recorded from  
 226 temperature -120 °C to 70°C. It shows that thermal degradation of membrane in metalworking fluid  
 227 significantly reduces its Young modulus. It is clearly seen that the Young modulus is decreased from  
 228  $1.45 \times 10^7$  to  $4 \times 10^6$  MPa at -120 °C after aging at 100 °C for 24 h of exposure.  
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235 **Figure 3** Evolution of storage modulus (a) and tangent delta values (b) of butyl rubber composite  
 236 membrane before and after exposure in metalworking fluid (Milform 64 SST) at 100  $^{\circ}\text{C}$  during 24 h.

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238 For an elastomer material, the glass transition temperature ( $T_g$ ) is important factor that directly  
 239 involves to the final properties of materials. This parameter is directly determined by DMTA curves,  
 240 corresponding to the maximum peak of  $\tan(\delta)$ . Figure 3b shows the example of the variation of  $\tan$   
 241 ( $\delta$ ) of aged and un-aged samples at 100  $^{\circ}\text{C}$  with different aging times. It can be seen that the curve  
 242 has a transition located around at  $-45$   $^{\circ}\text{C}$ , due to the glass transition temperature of butyl rubber.  
 243 Another peak at a higher temperature at  $-8^{\circ}\text{C}$  was also observed. This may be associated to movement  
 244 of local molecular relaxations side [60, 61]. These local molecular movements not only affect the

245 viscoelastic response, but also the mechanical properties (elastic modulus, plastic deformation) and  
246 also the diffusion of solvents in polymer matrix.

247 After aging at 100 °C in a metalworking fluid (Milform 64SST) for 24 h, a significant increase of  
248 the glass transition temperature value from - 45 °C to - 63 °C temperature was observed. This result  
249 is in good agreement with that in the literature in showing that there was a decrease of  $T_g$  of BRC  
250 during thermal aging in silicone oil [62]. The intensity of tan peak ( $\delta$ ) of aged sample is observed to  
251 be lower than that of the un-aged sample. The disappearance of the peak corresponding to the  
252 secondary transition temperature associated with the local molecular movement is observed in the  
253 aged sample. It can be attributed by the presence of MWF in elastomer matrix and leads to a better  
254 "molecular lubrication" due to the plasticization affect [63]. Regard to the reduction of  $T_g$ , since the  
255 glass transition temperature varies linearly with the molar mass and the molecular loss lead to a shift  
256 of the  $T_g$  to lower temperature areas [64]. The decrease of  $T_g$  observed this present study, is may be  
257 due to the thermal and chemical degradation of material by chains scissions that lead to a reduction  
258 of the molecular mass, that can explain the deterioration of mechanical properties of the material as  
259 shown in the previous section.

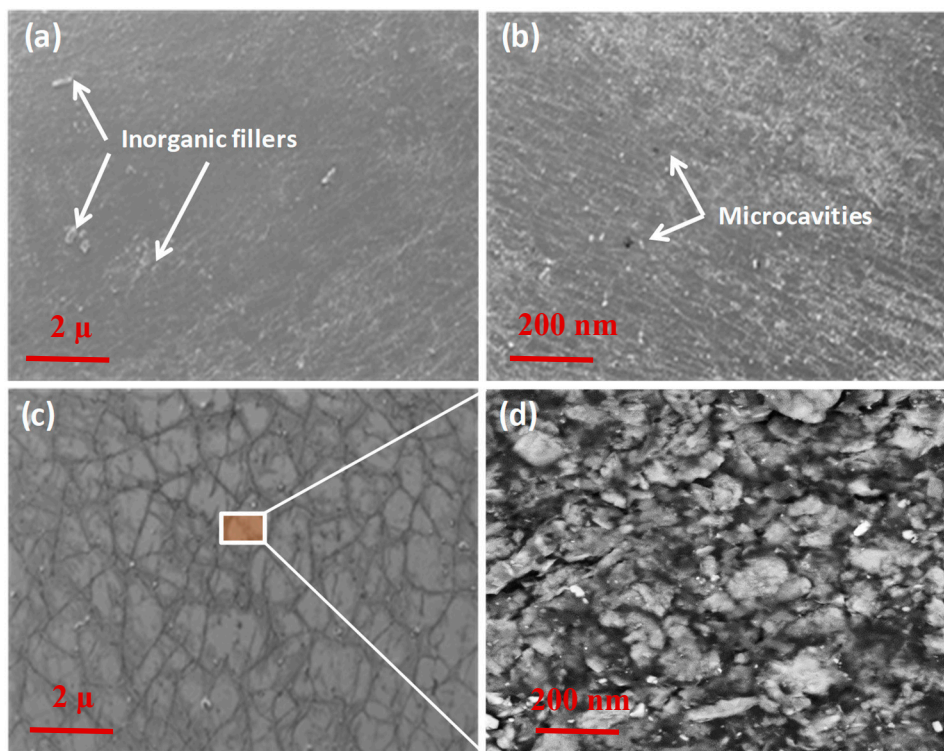
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261 3.4. *Effect of temperature and MWF on morphology, thermal properties and service lifetime of butyl*  
262 *composites*

263 The scanning electron microscope was used to study the changes of the BRC membrane  
264 morphology during aging process. Figure 4 shows the SEM images of a sample of butyl before and  
265 after exposure in MWF. Figure 4 d shows that in the case of unexposed sample, a large distribution  
266 of fillers in the elastomer matrix is observed. However, these reinforcements are missing at the  
267 surface of exposed sample. The appearance of holes on the surface of the aged sample is also  
268 observed, this can be due to the removal of fillers that replaced by metalworking fluid (Fig. 4b). In  
269 other studies, the same damage was observed with bromobutyl rubber when they are undergone a  
270 thermal aging process [65]. The presence of MWF at elevated temperature, various micro cracks on  
271 the exposed sample (Fig. 4c) can be found due to the loss of reinforcements at the sample surface,  
272 generated by thermo-oxidation aging and plasticization effect. This observation is in good agreement  
273 the literature [65]. The appearance of micro-cracks on rubber materials is believed to derive from the  
274 nucleation effect of the dissolved vapor and gas in the low molecular weight domains of the rubber  
275 [66]. The loss of reinforcement is confirmed by thermogravimetric analysis (TGA) under argon  
276 atmosphere conditions. The thermal stability of BRC membrane was evaluated by thermo  
277 gravimetric analysis at the temperature range from 30 to 800°C (Figure 5). The TGA curves show that  
278 the decomposition BRC evolves in the same way for both aged and un-aged samples. It found that  
279 the onset decomposition temperature of an original sample is about 205 °C with a residue of 52 % at  
280 460 °C while the onset decomposition temperature for the aged one is about 228 °C with a residue of  
281 42 % at 460 °C. In other words, the aged process leads to a reduction of carbon black filler from 52 %  
282 for original rubber to about 42 % for aged sample.

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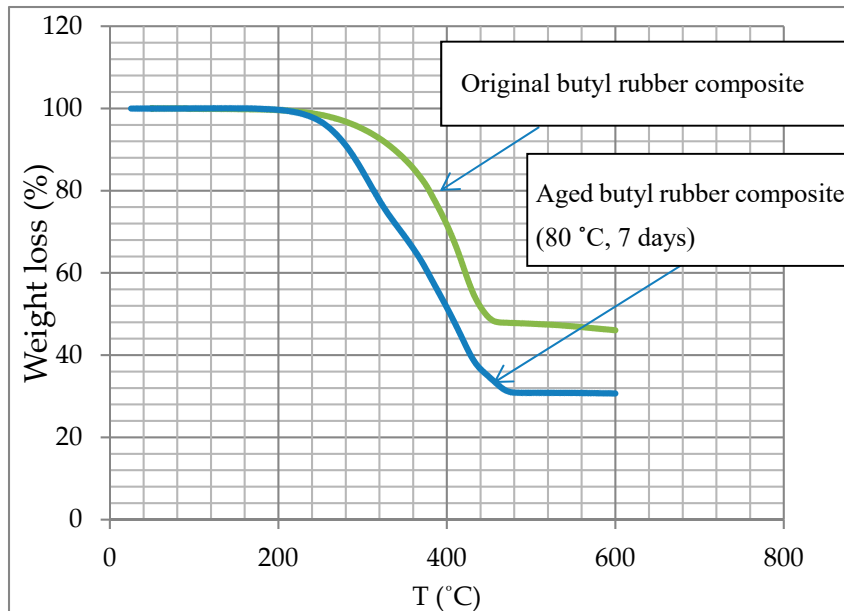
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**Figure 4** SEM images of butyl rubber composite membranes: a) non-aged sample, b) aged at 100 °C, c) aged at 100 °C with the presence of a metalworking fluid (Milform 64 SST) during 24 hrs, d) higher magnification of image 4c, showing the rugosity of sample surface.



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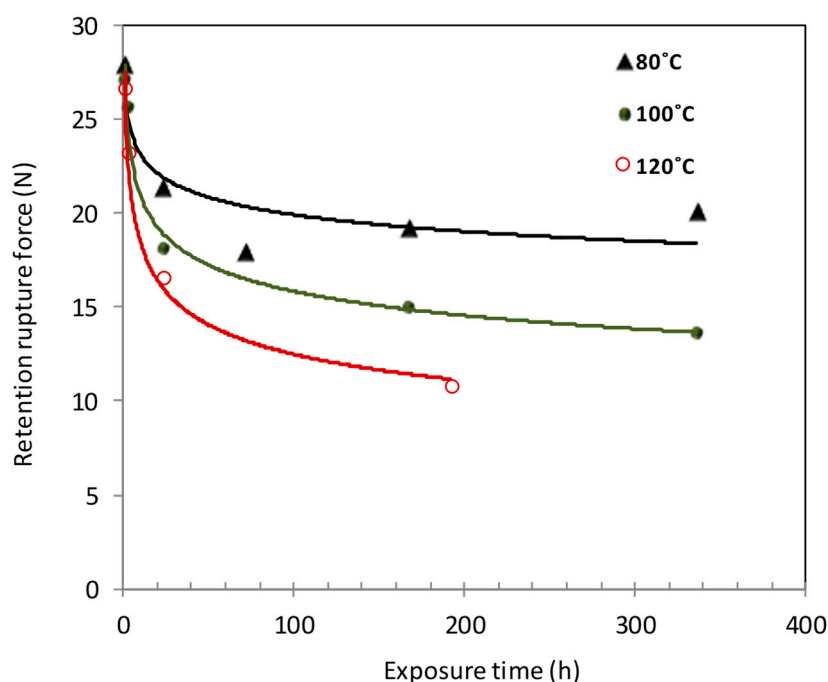
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**Figure 5** Thermogravimetric curves of butyl rubber composite before and after aged in Milform 64 SST at 80 °C during a week.

297 Changes in tensile force are observed at all studied temperatures (Figure 6). Indeed, after 24 hrs  
 298 of exposure, the tensile force drops by about 23 % at 80 °C. Whereas it decreases by about 41 % at 120  
 299 °C. The results of tensile tests of BRC at different temperatures helps to give a prediction of the service  
 300 life out by applying the Arrhenius model to the time-temperature data (Eq.9). In fact, the variation of  
 301 rate of overall aging processes can be calculated from mentioned above equation (Eq.9). Where E is  
 302 activation energy, R is universal constant (8.314 J/mol.K), T is absolute temperature and A is the pre-  
 303 exponential factor. The curves describing the relationship between  $\ln(t)$  vs  $1/T$  should be linear and  
 304 thus the activation energy and service life time can be calculated from this curve.

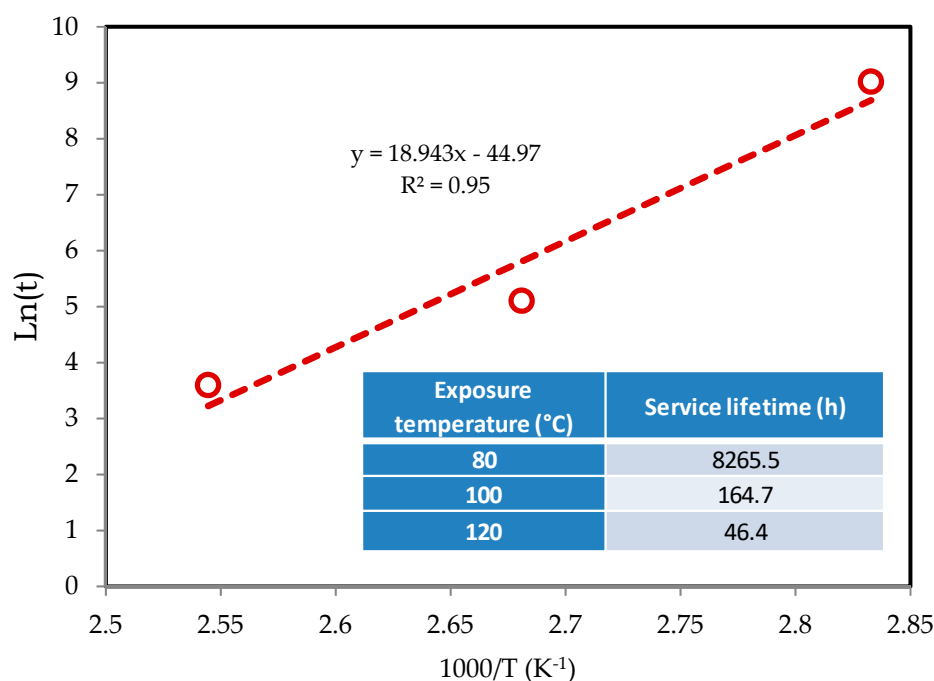
$$K = A \exp(-E/RT) \quad \text{Eq.9}$$



309  
 310 **Figure 6** Variation of tensile force of BRC at different temperatures as a function of exposure time.  
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312 The definition of the end-used service lifetime of the material is still a subject in question in the  
 313 literature. However, the assumption proposed by Rachid *et al.*[67] in which the material is considered  
 314 run out when there is a reduction of about 50 % of tensile force seems to be acceptable. Here, were  
 315 calculated the service life time for each aging temperature, using an empirical regression proposed  
 316 method By El Aidani *et al.* [16, 67]. These lifetime values were exploited from Arrhenius curve, as  
 317 shown in Figure 7, by plotting the logarithm of the lifetimes as a function of the inverse of the exposed  
 318 temperatures. This Figure shows a good agreement with the *Arrhenius* model with high linear  
 319 correlation coefficient ( $R^2 = 0.95$ ). This indicates that the effect of thermal treatment between 80 and  
 320 120 °C on tensile force of BRC can be satisfactorily described by the Arrhenius model, which can  
 321 therefore be used to predict the lifetime in this range of temperatures. The results of service life time  
 322 of BRC can be found in Figure 6. An activation energy value of 157 kJ.mol<sup>-1</sup> was obtained from the

323 Arrhenius curve. That means, at 80 °C, the mechanical properties of BRC remains acceptable till 344  
 324 days while it drops to about 2 days at 120 °C.  
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326  
 327 **Figure 7** Extrapolation of Arrhenius plot of BRC membrane based on tensile force. Insert:  
 328 Predicted values of service lifetime, calculated from Arrhenius plot based on tensile force at different  
 329 temperature.

#### 330 4. Conclusions

331 The effect of temperature with and without MWFs on the evolution of mechanical and physico-  
 332 chemical descriptors as well as structure of BRC is investigated. The Mooney-Rivlin's theory is used  
 333 to determine the changes of physical and chemical bonds and the viscoelastic behavior of BRCs when  
 334 they are in contact with a MWFs at high temperature via the determination of the elastic constant  $C_1$   
 335 and  $C_2$  at different temperatures ranging from 80 °C to 120 °C. The results show that temperature and  
 336 MWFs lead to the decrease of elastic constant  $C_2$ , suggesting a decrease of the force of cohesion of  
 337 polymer network. When BRC membranes are in contact with MWFs at high temperature, the  
 338 irreversible behavior, related to intrinsic property of elastomers, is missing and significant changes  
 339 in mechanical properties of material are observed.

340 The temperature is found to be accelerated the degradation process and the loss of mechanical  
 341 properties of material was more pronounced at high temperature. The decrease of elasticity and  
 342 reduction of the glass transition temperature of BRC is also observed, implying that BRC is not  
 343 appropriated to use in oily working conditions, particularly in high temperature. In this case, gloves  
 344 making from BRC must be changed more frequently to ensure an adequate protection level against  
 345 mechanical and chemical risks. This work provides also useful information for rubber manufacturers  
 346 to better understand the physicochemical degradation phenomena of their products under oily  
 347 working conditions and thus improve the products quality.

348 **Author Contributions:** P.N.T has planned, supervised the works and written the manuscript. E.T has realized  
 349 the experiments and participated to explain results. T.A.N. participated to edit the manuscript.

350 **Funding:** This research was funded by the *Institut de Recherche Robert-Sauvé en Santé et en Sécurité du Travail*,  
351 Quebec, Canada (Project R-983).

352 **Acknowledgments:** Thanks also to Prof. Jaime Lara (university of Montreal) for his constant support during this  
353 project.

354 **Conflicts of Interest:** The authors declare no conflict of interest.

## 355 5. References

- 356  
357 1. Youssf, O., et al., *Cyclic Performance of Steel–Concrete–Steel Sandwich Beams with Rubcrete and LECA*  
358 *Concrete Core*. Journal of Composites Science, 2019. **3**(1): p. 5.
- 359 2. Youssf, O., et al., *Influence of Mixing Procedures, Rubber Treatment, and Fibre Additives on Rubcrete*  
360 *Performance*. Journal of Composites Science, 2019. **3**(2): p. 41.
- 361 3. Aguilar-Bolados, H., et al., *Removal of Surfactant from Nanocomposites Films Based on Thermally Reduced*  
362 *Graphene Oxide and Natural Rubber*. Journal of Composites Science, 2019. **3**(2): p. 31.
- 363 4. Nohilé, C., P.I. Dolez, and T. Vu-Khanh, *Mechanical and Chemical Effects of Solvent Swelling on Butyl*  
364 *Rubber*. 2010: p. 527-534.
- 365 5. Ebnesajjad, S., *8 - Characteristics of Adhesive Materials*, in *Handbook of Adhesives and Surface Preparation*,  
366 S. Ebnesajjad, Editor. 2011, William Andrew Publishing: Oxford. p. 137-183.
- 367 6. G. Pape, P., *15 - Adhesion Promoters*, in *Handbook of Adhesives and Surface Preparation*, S. Ebnesajjad,  
368 Editor. 2011, William Andrew Publishing: Oxford. p. 369-386.
- 369 7. Rabilloud, G., *12 - Adhesives for Electronics*, in *Handbook of Adhesives and Surface Preparation*, S.  
370 Ebnesajjad, Editor. 2011, William Andrew Publishing: Oxford. p. 259-299.
- 371 8. Rabilloud, G., *9 - Heat-Resistant Adhesives*, in *Handbook of Adhesives and Surface Preparation*, S.  
372 Ebnesajjad, Editor. 2011, William Andrew Publishing: Oxford. p. 185-220.
- 373 9. Fabris, H.J. and W.G. Knauss, *Synthetic Polymer Adhesives*. 1989: p. 131-177.
- 374 10. Haworth, J.P. and F.P. Baldwin, *Butyl Rubber Properties and Compounding*. Industrial & Engineering  
375 Chemistry, 1942. **34**(11): p. 1301-1308.
- 376 11. Ford, F.P. and A.M. Gessler, *Some Properties of Butyl Rubber-Carbon Black Systems*. Industrial &  
377 Engineering Chemistry, 1952. **44**(4): p. 819-824.
- 378 12. Yu, B.-C., et al., *A new approach for recycling waste rubber products in Li–S batteries*. Energy &  
379 Environmental Science, 2017. **10**(1): p. 86-90.
- 380 13. Xu, W. and S.S. Que Hee, *Permeation of a Metalworking Fluid Through a Latex Glove Under Field Use*  
381 *Conditions*. Bulletin of Environmental Contamination and Toxicology, 2009. **84**(1): p. 5.
- 382 14. Xu, W. and S.S. Que Hee, *Swelling of four glove materials challenged by six metalworking fluids*. Arch  
383 Environ Contam Toxicol, 2008. **54**(1): p. 1-8.
- 384 15. Nguyen-Tri, P., et al., *Swelling behavior of polymeric membranes to metalworking fluids*. Journal of Applied  
385 Polymer Science, 2018. **135**(3): p. 45717.
- 386 16. El Aidani, R., et al., *Photochemical aging of an e-PTFE/NOMEX® membrane used in firefighter protective*  
387 *clothing*. Polymer Degradation and Stability, 2013. **98**(7): p. 1300-1310.
- 388 17. Nguyen-Tri, P., et al., *Chemical ageing of a polyester nonwoven membrane used in aerosol and drainage*  
389 *filter*. Polymer Degradation and Stability, 2014. **101**: p. 71-80.
- 390 18. Nguyen, T.V., et al., *Accelerated degradation of water borne acrylic nanocomposites used in outdoor protective*  
391 *coatings*. Polymer Degradation and Stability, 2016. **128**: p. 65-76.

- 392 19. Nguyen, T.P., *Nanoscale analysis of the photodegradation of Polyester fibers by AFM-IR* Journal of  
393 Photochemistry and Photobiology A: Chemistry, 2018. **Accepted**.
- 394 20. Nguyen, T.V., et al., *Stability of acrylic polyurethane coatings under accelerated aging tests and natural*  
395 *outdoor exposure: The critical role of the used photo-stabilizers*. Progress in Organic Coatings, 2018. **124**: p.  
396 137-146.
- 397 21. Nguyen-Tri, P., et al., *Nanocomposite Coatings: Preparation, Characterization, Properties, and Applications*.  
398 International Journal of Corrosion, 2018. **2018**: p. 1-19.
- 399 22. Nguyen-Tri, P. and R.E. Prud'homme, *Nanoscale analysis of the photodegradation of polyester fibers by*  
400 *AFM-IR*. Journal of Photochemistry and Photobiology A: Chemistry, 2019. **371**: p. 196-204.
- 401 23. Nguyen-Tri, P., et al., *Recent progress in the preparation, properties and applications of superhydrophobic*  
402 *nano-based coatings and surfaces: A review*. Progress in Organic Coatings, 2019. **132**: p. 235-256.
- 403 24. Ouellet-Plamondon, P.N.T.S.R.C., *Nanomaterials-Based Coatings : Fundamentals and Applications*. Vol. 1.  
404 2019: Elsevier. 474.
- 405 25. Tri Phuong, N., V. Gilbert, and B. Chuong, *Preparation of Recycled Polypropylene/ Organophilic Modified*  
406 *Layered Silicates Nanocomposites Part I: The Recycling Process of Polypropylene and the Mechanical*  
407 *Properties of Recycled Polypropylene/Organoclay Nanocomposites*. Journal of Reinforced Plastics and  
408 Composites, 2008. **27**(18): p. 1983-2000.
- 409 26. Nguyen Tri, P. and V. Gilbert, *Non-isothermal Crystallization Kinetics of Short Bamboo Fiber-reinforced*  
410 *Recycled Polypropylene Composites*. Journal of Reinforced Plastics and Composites, 2010. **29**(17): p. 2576-  
411 2591.
- 412 27. Nguyen Tri, P., C. Sollogoub, and A. Guinault, *Relationship between fiber chemical treatment and*  
413 *properties of recycled pp/bamboo fiber composites*. Journal of Reinforced Plastics and Composites, 2010.  
414 **29**(21): p. 3244-3256.
- 415 28. Boukehili, H. and P. Nguyen-Tri, *Helium gas barrier and water absorption behavior of bamboo fiber*  
416 *reinforced recycled polypropylene*. Journal of Reinforced Plastics and Composites, 2012. **31**(23): p. 1638-  
417 1651.
- 418 29. Nguyen Tri, P., A. Guinault, and C. Sollogoub, *Élaboration et propriétés des composites polypropylène*  
419 *recyclé/fibres de bambou*. Matériaux & Techniques, 2012. **100**(5): p. 413-423.
- 420 30. Triki, E., et al., *Investigation of tearing mechanisms of woven textile*. Polymer Composites, 2012. **33**(9): p.  
421 1578-1585.
- 422 31. Triki, E., et al., *Mechanics and mechanisms of tear resistance of woven fabrics*. Theoretical and Applied  
423 Fracture Mechanics, 2012. **61**: p. 33-39.
- 424 32. Tri, P.N., et al., *Crystallization behavior of poly(lactide)/poly( $\beta$ -hydroxybutyrate)/talc composites*. Journal of  
425 Applied Polymer Science, 2013. **129**(6): p. 3355-3365.
- 426 33. Triki, E., et al., *Combined puncture/cutting of elastomer membranes by pointed blades: Characterization of*  
427 *mechanisms*. Journal of Applied Polymer Science, 2015. **132**(26): p. n/a-n/a.
- 428 34. Azizi, S., et al., *Electrical and thermal phenomena in low-density polyethylene/carbon black composites near*  
429 *the percolation threshold*. Journal of Applied Polymer Science, 2018: p. 47043.
- 430 35. Azizi, S., et al., *Electrical and thermal conductivity of ethylene vinyl acetate composite with graphene and*  
431 *carbon black filler*. Polymer Testing, 2018. **72**: p. 24-31.
- 432 36. Nguyen Tri, P. and R.E. Prud'homme, *Crystallization and Segregation Behavior at the Submicrometer Scale*  
433 *of PCL/PEG Blends*. Macromolecules, 2018. **51**(18): p. 7266-7273.

- 434 37. Satyabrata, M.T., Anh Nguyen; Phuong, Nguyen-Tri, *Noble Metal-Metal Oxide Hybrid Nanoparticles*.  
435 Vol. 1. 2018: Elsevier.
- 436 38. Duc Chinh, V., et al., *Synthesis of Gold Nanoparticles Decorated with Multiwalled Carbon Nanotubes (Au-*  
437 *MWCNTs) via Cysteaminium Chloride Functionalization*. Scientific Reports, 2019. **9**(1): p. 5667.
- 438 39. Nguyen Tri, P., et al., *Antibacterial Behavior of Hybrid Nanoparticles*. 2019: p. 141-155.
- 439 40. Nguyen Tri, P., et al., *Methods for Synthesis of Hybrid Nanoparticles*. 2019: p. 51-63.
- 440 41. Nguyen-Tri, P., V. Nguyen, and T. Nguyen, *Biological Activity and Nanostructuration of Fe<sub>3</sub>O<sub>4</sub>-Ag/High*  
441 *Density Polyethylene Nanocomposites*. Journal of Composites Science, 2019. **3**(2): p. 34.
- 442 42. Ben Hassine, M., et al., *Time to failure prediction in rubber components subjected to thermal ageing: A*  
443 *combined approach based upon the intrinsic defect concept and the fracture mechanics*. Mechanics of  
444 Materials, 2014. **79**: p. 15-24.
- 445 43. Syed, I.H., et al., *Nonlinearity in the Mechanical Response of Rubber as Investigated by High-Frequency*  
446 *DMA*. Polymers (Basel), 2019. **11**(4).
- 447 44. Xiang, K., et al., *Investigation on the thermal oxidative aging mechanism and lifetime prediction of butyl*  
448 *rubber*. Macromolecular Research, 2013. **21**(1): p. 10-16.
- 449 45. Haghtalab, A. and S. Rahimi, *Study of viscoelastic properties of nanocomposites of SiO<sub>2</sub>-acrylonitrile-*  
450 *butadiene-styrene*. Journal of Applied Polymer Science, 2013. **127**(6): p. 4318-4327.
- 451 46. Lewicki, J.P., et al., *Degradative Thermal Analysis and Dielectric Spectroscopy Studies of Aging in*  
452 *Polysiloxane Nanocomposites*. 2009. **1004**: p. 239-254.
- 453 47. Rivlin, R.S., *Large Elastic Deformations of Isotropic Materials. IV. Further Developments of the General*  
454 *Theory*. Philosophical Transactions of the Royal Society A: Mathematical, Physical and Engineering  
455 Sciences, 1948. **241**(835): p. 379-397.
- 456 48. Rivlin, R.S., *Large Elastic Deformations of Isotropic Materials. I. Fundamental Concepts*. Philosophical  
457 Transactions of the Royal Society A: Mathematical, Physical and Engineering Sciences, 1948. **240**(822):  
458 p. 459-490.
- 459 49. Arimoto, H.,  *$\alpha$ - $\gamma$  Transition of nylon 6*. Journal of Polymer Science Part A: General Papers, 1964. **2**(5): p.  
460 2283-2295.
- 461 50. Slichter, W.P., *Molecular motion in polyamides*. Journal of Polymer Science, 1959. **35**(128): p. 77-92.
- 462 51. Hagen, R., L. Salmén, and B. Stenberg, *Effects of the type of crosslink on viscoelastic properties of natural*  
463 *rubber*. Journal of Polymer Science Part B: Polymer Physics, 1996. **34**(12): p. 1997-2006.
- 464 52. Gordon, M., *The Physics of Rubber Elasticity (Third Edition)*. L. R. G. Treloar, Clarendon Press, Oxford. 1975  
465 pp. xii + 370. Price: £14.00. British Polymer Journal, 1976. **8**(1): p. 39-39.
- 466 53. Sombatsompop, N., *Investigation of Swelling Behavior of NR Vulcanisates*. Polymer-Plastics Technology  
467 and Engineering, 1998. **37**(1): p. 19-39.
- 468 54. Mark, J.E., *Experimental Determinations of Crosslink Densities*. Rubber Chemistry and Technology, 1982.  
469 **55**(3): p. 762-768.
- 470 55. Priss, L.S., *Molecular origin of constants in the theory of rubber-like elasticity considering network chains*  
471 *steric interaction*. Pure and Applied Chemistry, 1981. **53**(8): p. 1581-1596.
- 472 56. Mark, J.E., *The Constants 2C<sub>1</sub> and 2C<sub>2</sub> in Phenomeno-Logical Elasticity Theory and Their Dependence on*  
473 *Experimental Variables*. Rubber Chemistry and Technology, 1975. **48**(3): p. 495-512.
- 474 57. Ha-Anh, T. and T. Vu-Khanh, *Prediction of mechanical properties of polychloroprene during thermo-*  
475 *oxidative aging*. Polymer Testing, 2005. **24**(6): p. 775-780.



- 476 58. Gumbrell, S.M., L. Mullins, and R.S. Rivlin, *Departures of the elastic behaviour of rubbers in simple*  
477 *extension from the kinetic theory*. Transactions of the Faraday Society, 1953. **49**(0): p. 1495-1505.
- 478 59. Allen, G., et al., *Thermodynamics of rubber elasticity at constant volume*. Transactions of the Faraday  
479 Society, 1971. **67**(0): p. 1278-1292.
- 480 60. Dutta, N.K., D. Khastgir, and D.K. Tripathy, *The effect of carbon black concentration on the dynamic*  
481 *mechanical properties of bromobutyl rubber*. Journal of Materials Science, 1991. **26**(1): p. 177-188.
- 482 61. García, M., I. Garmendia, and J. García, *Influence of natural fiber type in eco-composites*. Journal of  
483 Applied Polymer Science, 2008. **107**(5): p. 2994-3004.
- 484 62. Ali, S., et al., *Thermal and Thermo-Mechanical Behavior of Butyl based Rubber Exposed to Silicon Oil at*  
485 *Elevated Temperature*. Journal of the Chemical Society of Pakistan, 2013. **35**(6): p. 1437-1444.
- 486 63. Barrere, C. and F. Dal Maso, *Résines époxy réticulées par des polyamines : structure et propriétés*. Revue de  
487 l'Institut Français du Pétrole, 2006. **52**(3): p. 317-335.
- 488 64. Fox, T.G. and P.J. Flory, *Second-Order Transition Temperatures and Related Properties of Polystyrene. I.*  
489 *Influence of Molecular Weight*. Journal of Applied Physics, 1950. **21**(6): p. 581-591.
- 490 65. G K Kannan, L.V.G., L Nirmala and N S Kumar, *Thermal ageing studies of bromo-butyl rubber used in*  
491 *NBCpersonal protective equipment*. Journal of Scientific & Industrial Research, 2010. **69**: p. 841-849.
- 492 66. Grandcoin, J., A. Boukamel, and S. Lejeunes, *A micro-mechanically based continuum damage model for*  
493 *fatigue life prediction of filled rubbers*. International Journal of Solids and Structures, 2014. **51**(6): p. 1274-  
494 1286.
- 495 67. El Aidani, R., P.I. Dolez, and T. Vu-Khanh, *Effect of thermal aging on the mechanical and barrier properties*  
496 *of an e-PTFE/Nomex® moisture membrane used in firefighters' protective suits*. Journal of Applied Polymer  
497 Science, 2011. **121**(5): p. 3101-3110.