

SEAWATER AGEING OF LOW STYRENE EMISSION RESINS FOR MARINE COMPOSITES: MECHANICAL BEHAVIOUR AND NANOINDENTATION STUDIES.

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ABSTRACT

Polyester resins are widely employed for pleasure boat construction. In order to satisfy new environmental legislation on styrene emissions resin suppliers have proposed new formulations, but these show lower failure strains under tensile loading. This paper examines the influence of wet ageing on mechanical properties of low styrene polyester, together with standard polyester and vinyl ester resins. Results are presented from accelerated ageing in natural sea water for 9 months, at 40°C. Heating during accelerated aging results in further cure, which complicates the interpretation of results (simultaneous cross-linking and degradation). Nanoindentation was used to study local changes in the resin after aging. The diffusion kinetics of the low styrene emission resins are similar to those for standard polyester resins. More brittle initially they remain more brittle after aging.

INTRODUCTION

Polyester resins have been employed for pleasure boat construction since the 1950's. However, in recent years concerns over volatile emissions during fabrication have resulted in new environmental legislation on styrene emissions and resin suppliers have proposed new formulations. These are of three types:

- Low styrene content. In this case the molecular weight of the pre-polymer is reduced to keep viscosity low.
- Low styrene emission resins. Here additives are included which block styrene emission during fabrication by creating a surface film.
- Mixed resins, combining low styrene content and emission.

The mechanical properties of glass reinforced polyester laminates depend on a number of factors. The fibre orientation and content are critical but the matrix also plays an important role in maintaining the geometry, providing off-axis strength and protecting the fibres from the environment. The fibre/matrix interface also contributes to these functions. In that study significant differences were noted between the mechanical properties of standard polyester resins and those limiting styrene emissions. For similar resin formulations Perrot et al [2] examined composite delamination resistance and impact behaviour. The low styrene resins showed lower resistance to delamination

propagation under mode I loading (for unidirectional composites), and larger delaminated areas for a given impact energy (for rovimat reinforced composites). These results show the importance of the matrix in the events leading to failure of glass/polyester composites.

The ductility of the matrix resin is a critical parameter which conditions the appearance and progression of damage. The aim of the present presentation is to complete the study of the behaviour of low styrene resins by presenting a study of their ageing behaviour. Many published studies provide data on aged glass reinforced composites, e.g. [3-6], but very few results allow a comparison to be made between the ageing of standard and low styrene materials. This is the main aim of the work described below. Standard resins have been subjected to immersion in sea water for 9 months. Weight gain due to diffusion and the change in mechanical properties have been studied using conventional mechanical tests. Nano-indentation is used on resin samples in order to examine whether this technique can provide additional information on the mechanisms involved in aging.

MATERIALS AND EXPERIMENTAL METHODS

Materials

Four commercial resins have been studied, supplied by the three main suppliers to the French boatbuilding industry: a standard orthophthalic polyester (SO), a low styrene content DCPD (dicyclopentadiene) orthophthalic polyester (LS), a low styrene emission orthophthalic polyester (LES) and a standard vinylester (SV). The two low styrene polyesters (LS, and LES) are commonly used in pleasure boat construction. Table 1 indicates their styrene contents given by suppliers and tensile properties measured in a previous study [1]. The failure strains of the low styrene resins are quite low, it should be remembered that glass fibres fail at strains around 2.8%. All the resins used in this study were mixed in a laboratory at room temperature ($20 \pm 2^\circ\text{C}$) with 1.5 % by weight of methyl ethyl ketone peroxide (MEKP). The cure cycle was 24 hours at room temperature followed by 16 hours at 40°C , which enables a cure state to be reached which is close to that encountered in industrial parts [1]. Unreinforced resin panels 4mm thick were obtained by casting and were then machined to produce dog-bone specimens.

Resin	Styrene content (wt. %)	Young's modulus (MPa)	Failure stress (MPa)	Failure strain (%)
SO	42	3077 ± 192	49 ± 7	$2,1 \pm 0,5$
LS	35 – 38	3120 ± 188	30 ± 7	$1,2 \pm 0,4$
LES	37 – 39	3295 ± 54	25 ± 3	$0,9 \pm 0,1$
SV	48 – 51	2901 ± 125	60 ± 2	$3,2 \pm 0,2$

Table 1: Initial resin properties [1]



Figure 1: Nanoindentation machine

Aging conditions

Directly after curing resins were placed in continuously circulating natural sea water containers for up to 9 months. The seawater is pumped directly from the Brest Estuary and is available on tap at the IFREMER centre. The only filtering is to remove sediment particles. The resins were aged at 40°C. Mass gain was followed by periodic weighing on a *Sartorius LA 310 S* balance (precision 0.1 mg). Samples were removed from the aging containers and wiped with paper towels before weighing to dry the surfaces. The mass M of each sample at time t is expressed as a percentage as :

$$M = \frac{M_t - M_0}{M_0} * 100 \quad (1)$$

Mechanical tests

An *Instron 4302* test machine was used to perform tensile tests on specimens at a loading rate of 2 mm/min. Resins were tested in the form of dog-bone specimens according to the ISO standard 5276. An *Instron* extensometer 2620-602 was used to measure strains with a gauge length of 25 mm. All samples were tested wet on removal from the aging containers, after a delay of 30 minutes to allow the specimen temperature to reach room temperature. At least four specimens were tested for each aging condition. Dynamic mechanical analysis (DMA) was also performed in three point flexure on resins after 9 months in water at 40°C, using a *TA Instruments DMA 2980*. The standard ISO 6721 procedure was applied (1 Hz, 3°C/min distance between supports 50 mm). The glass transition temperatures (T_g) quoted here correspond to the change in slope of the plots of storage modulus versus temperature.

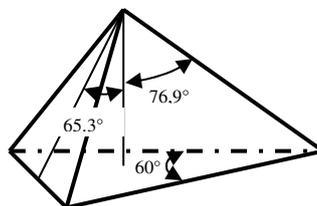


Figure 2: Berkovich indenter

Nanoindentation



Figure 3: Example of indent on resin surface.

Tests were performed on resin samples using an *MTS XP* machine (Figure 1) with a Berkovitch indenter (Figure 2). The tip displacement is controlled during indentation with a loading frequency of 70Hz (dynamic / continuous stiffness measurement method unless noted otherwise). Figure 3 shows the form of the indents. The loads are measured during indentation and unloading. Figure 4 shows a typical force-displacement plot ($P=f(h)$) for a polyester resin.

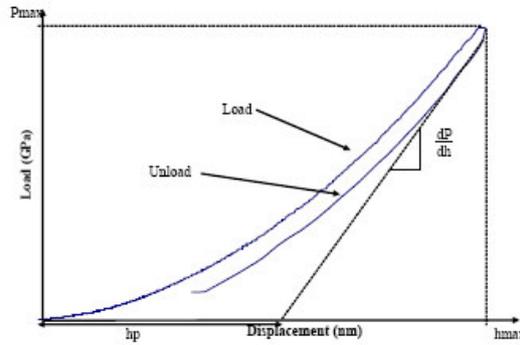


Figure 4: Example of force-displacement recording from nano-indentation test on polyester resin

Contact pressure is determined from the plastic penetration depth. The imprint area, A , is calculated as [7]:

$$A = 3 \sqrt{3} h_p^2 \tan^2(\Theta) \quad (2)$$

With $\theta = 65.3^\circ$ for the Berkovitch tip this simplifies to $A = 24.5h_p^2$. From the contact depth (h_p), the maximum depth (h_{max}) and the maximum force (P_{max}), a contact stiffness ($S = dP/dh$) is calculated. For this tip the geometrical factor $\varepsilon = 0.75$.

$$h_p = h_{max} - \varepsilon \frac{P_{max}}{S} \quad (3)$$

The contact stiffness allows a reduced modulus (E_r) to be determined as a function of the geometry of the tip using the method proposed by Oliver and Pharr [8]. A is the contact area, and ε is a geometrical constant, $\beta = 1.034$ for the Berkovitch tip.

$$E_r = \frac{S \sqrt{\pi}}{2 \beta \sqrt{A}} \quad (4)$$

The reduced modulus does not take into account the strain in the diamond tip. In order to obtain the sample modulus this component must be removed (using indenter modulus $E_i=1140$ GPa and indenter Poisson's ratio $\nu_i = 0.07$) [6].

$$E = \frac{(1 - \nu^2)}{\frac{1}{E_r} - \frac{(1 - \nu_i^2)}{E_i}} \quad (5)$$

Nanoindentation was performed on the aged resin tensile specimens described above. Before each series of tests a calibration indentation was performed according to the standard test method described in [7]. Each test then consisted of placing the tip in contact with the sample and loading. Indentation depth was 2000 nm and the rate of loading and unloading was 0.05 s^{-1} . This depth was chosen following preliminary studies using different depths, which indicated larger variability of modulus values for small depths. The load was maintained for 60s. Then the sample was unloaded at the same rate to 10% of the maximum load and the thermal drift of the test was calculated. First, in order to measure surface modulus, a series of measurements was made directly after aging, at different points (50 measurements) to obtain a representative measurement of elastic modulus. In this case the as-aged surface was indented. A second series of measurements was made subsequently, this time through the specimen thickness in order to examine stiffness profiles. In this case specimens had been left for at least 9 months in air before indentation, in order to allow water desorption. These samples were cut and polished with diamond paste. Through thickness measurements were made at mid-length of the tensile specimens. Indentations were made along lines of 100 equi-distant indentations through the sample thickness, Figure 5

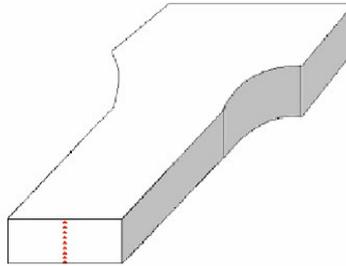


Figure 5. Position of nanoindentations,

RESULTS and DISCUSSION

Mass gain of resins Figure 6 shows how the masses of resins change with square root of time during immersion at 40°C . The three polyesters show similar behaviour, close to Fickian though a stable saturation plateau is not reached. The vinylester shows a different behaviour with much lower mass gain. Although the standard and low styrene polyesters have different chemical compositions they appear to absorb similar amounts of water. However it should be noted that mass gain depends on many factors, including the degree of cross-linking, interfaces and defects such as voids.

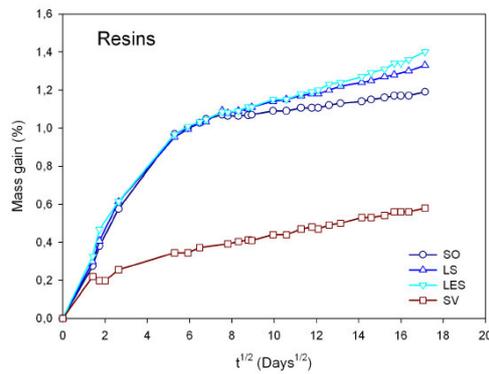


Figure 6: Resins mass gains in sea water at 40°C

Tensile properties of resins after aging

Figure 7 presents typical load-displacement plots and Figure 8 shows tensile properties of the four resins before and after aging for 9 months in seawater at 40°C. The trends are similar for all four materials, an increase in modulus (up to 16% for the polyesters and nearly 30% for the vinyl ester) and a large drop in failure strain (35% loss for the LES polyester, which is the most brittle initially, and between 50 and 60% for the other materials). The post-cure applied here (16h at 40°C) is used to obtain materials equivalent to industrial components left for 3 months at 20°C without postcure but does not result in complete cure [1]. Complete post-curing of samples would be possible, and might simplify interpretation of the results, but it is never found in service so has little practical significance. The thermal energy of seawater heated to 40°C enables cure to continue as clearly indicated by DMA results in Table 3. Values of T_g after 9 months at 40°C are 15 to 20°C higher than initial values. Similar increases in T_g and storage modulus are noted for the three polyesters. This evolution of cure makes the interpretation of the results more difficult, but the drop in strain to failure is very large and cannot be explained solely by higher cross-linking [1]. There is also a degradation of the resins. The low styrene resins lose proportionally a little less of their ductility after aging but as they are very brittle initially they are even more brittle after aging. It should also be noted that in service boat structures made with these resins will continue to cure, and temperatures of 40°C are not extreme. There will be a drop in T_g as water enters the material, typically 10°C per percent of water, but for an aging temperature of 40°C the material T_g will still be above the bath temperature.

Reference	E' at 30°C (GPa)		T _g (°C)	
	unaged	aged	unaged	aged
SO	3.5	3.8	58	73
LS	3.9	4.1	58	78
LES	3.5	3.6	58	78

Table 2: DMA results for polyesters before and after 9 months in seawater at 40°C.

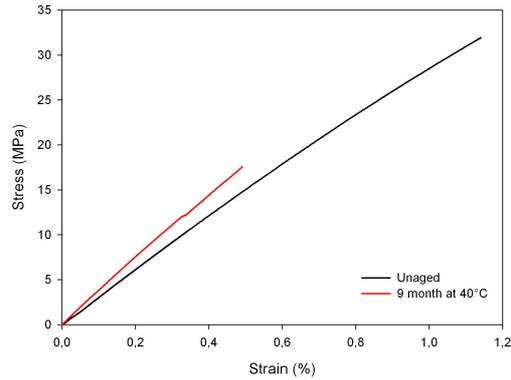


Figure 7: Typical tensile test stress-strain plots, LS resin

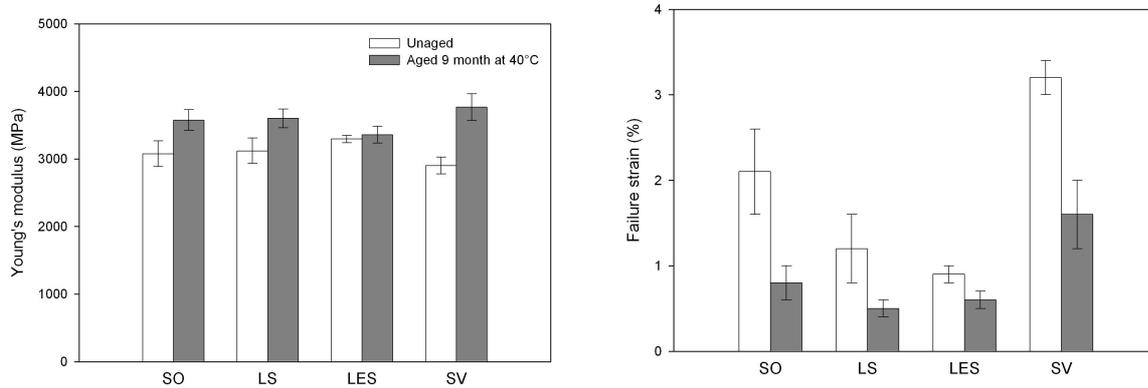


Figure 8 : Influence of 9 months' sea water ageing at 40°C on resin properties: Young's modulus and failure strain.

Nanoindentation.

The tests described above are traditionally used to study this type of material and provide a global indication of properties. In order to examine how properties evolve on a local level nanoindentation was used on three of the resins (SO, LS and SV). This is a very powerful technique but requires certain precautions to validate the procedures. A large number of preliminary tests allowed the test conditions to be defined. Modulus was independent of frequency over the range studied so 70 Hz was used for indentation through the sample thickness. At this frequency there is little influence of parasite frequencies. Nanoindentation is sensitive to surface modifications and moisture, but due to the long drying period any changes noted are caused by irreversible changes in the materials.

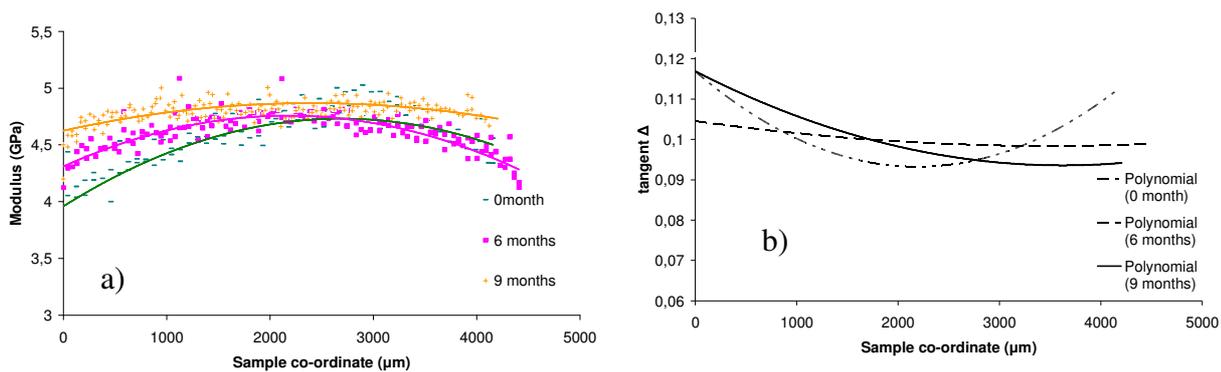


Figure 9 : Change in (a) modulus and (b) tangent Δ through the thickness, with aging time, resin SO

Figure 9 shows the profiles of E' and tangent Δ for the standard polyester. The first observation, Figure 9a, is that the modulus is not uniform through the specimen thickness. This is the case for all aging periods. The modulus is lower at the specimen edges and reaches a maximum at the specimen centre. Tangent Δ shows the opposite trend, Figure 9b, being highest at the edges and lowest in the centre. This is noted for all three resins but the kinetics of the changes during aging vary with resin type. The vinyl ester (not shown) shows similar behaviour. The properties of the central region hardly evolve during aging, while the properties of the faces (in contact with warm water) increase to reach those of the centre. This suggests that the dominant mechanism is a post-cure. For the first 6 months the modulus at the centre increases with no change to the outer region, while the tangent Δ decreases. After 6 months the central modulus stabilizes (E' and E''). Between 6 and 9 months the properties of the outer regions change until uniform behaviour is observed. This suggests that for these two resins the aging effects are also minor compared to the post-curing effects (increase in E' and decrease in $\tan \Delta$). The reasons for the evolution in properties from the centre to the outer surfaces are complex. Exothermic reactions may result in higher degree of curing at the centre initially, particularly as fabrication involves casting into a steel mould, which may result in temperature gradients from the surface to the centre. The subsequent contact with water at 40°C allows cure to continue, though chemical effects such as leaching out of small molecular weight species at the surface may also affect the final resin state.

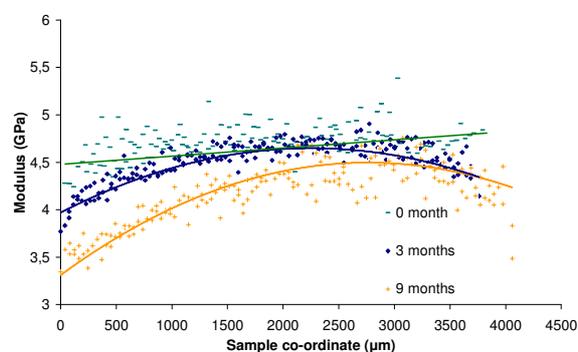


Figure 10: Change in modulus with aging time, resin LS.

The response of the low styrene content resin to aging is rather different, Figure 10. In this case the initial properties (E' , E'' and $\tan \Delta$) appear quite uniform. During aging

the modulus decreases significantly at the edges, while the tangent Δ starts to increase. This suggests that degradation related to water ingress is occurring. The main mechanisms which are observed during aging of polyesters are plasticization, which is generally reversible when water is removed, differential swelling, residual stress relaxation, and hydrolysis. The nanoindentation results presented here do not allow us to separate these phenomena, but they do clearly illustrate the heterogeneous nature of both the materials and the changes during aging. To complete this section a series of longitudinal indentations was performed on samples aged for nine months, to check that the changes observed were not confined to one section. Very similar profiles are noted for all three materials. It is important to note that nanoindentation and tensile tests do not measure exactly the same properties. Generally nanoindentation modulus values are higher than results from standard tensile tests. This is the case here, the former are from 10 to 50% higher than the latter as shown in Figure 11, which shows average values across the indented section. Several reasons have been invoked to explain this:

- The type of loading is compressive in nanoindentation, not tensile,
- The test frequencies are quite different (70Hz for nanoindentation, compared to a much lower frequency in the quasi-static tensile test) ,
- Hydrostatic pressure is generated below the Berkovitch indenter [10],
- Surface effects, the initial part of the force-depth curve is affected by surface roughness, local oxidation [11] and other surface phenomena.

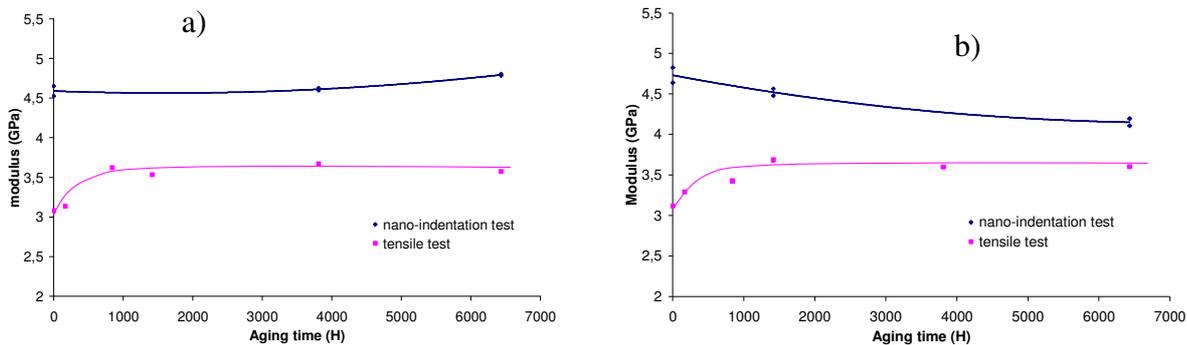


Figure 11 : Comparison between tensile test modulus and values from nanoindentation
a) SO (b) LS

While the trends in tensile test modulus and nanoindentation modulus appear similar for the standard materials SV and SO, which have mainly evolved due to postcuring, the low styrene resin results are contradictory. This may be related to the fact that while the tensile tests were performed directly after aging, on specimens containing water, nanoindentation was performed on specimens which had dried slowly over a period of several months. Relaxation and even hydrolysis may occur during the drying period. Rapid desorption by increasing temperature could have been used to minimize the influence of water during that period but this introduces other changes.

CONCLUSION

Glass fibre reinforced polyesters are widely used in boat-building, and low styrene resins satisfying environmental regulations are increasingly replacing standard

polyesters, both for contact moulding and infusion. In this study resin properties and the influence of aging in seawater have been examined at both the macroscopic and microscopic scale. This has shown that the macroscopic behaviour of low styrene resins is very similar to that of standard orthophthalic polyesters (same mass gain and property evolution). However, brittle initially, the low styrene grades are even more brittle after aging. Nanoindentation has revealed the heterogeneity of these materials and provides additional information on the complex changes which occur when post-curing and degradation occur simultaneously. The low styrene resins appear to lose strength more slowly than the standard resins but after 9 months similar losses are noted. It is tempting to suggest that changing from a standard polyester to a low styrene resin does not affect the aging behaviour of these marine composites, but further studies to examine coupling between damage and water ingress, and to quantify changes in damage tolerance are necessary before concluding.

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