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TESTS ON THE MECHANICAL PROPERTIES OF POLYMERS IN THE ASPECT OF AN ATTEMPT TO DETERMINE THE PARAMETERS OF THE MOONEY-RIVLIN HYPERELASTIC MODEL

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Abstract

The article presents testing of the mechanical properties of SIKA[®] polymer adhesives of the type PBM, PMM, PM, and PSM in the aspect of an attempt to determine the parameters of the Mooney-Rivlin hyperelastic model. The article contains a literature review on developed models for hyperelastic materials as well as a description of the author's own results obtained in monaxial tensile and monaxial compression tests conducted on oars and cylindrical samples, respectively. Furthermore, the results of modeling of Mooney-Rivlin hyperelastic model parameters are shown in relation to the value of average parameters for polymers after both a week and a month-and-a-half of ripening.

Keywords: hyperelastic model, hyperelasticity, polymers, adhesive joint, deformation, stresses

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1. INTRODUCTION

Polymers are classified to those chemical substances that are of particular interest in the world of science and industry because of the dynamic nature of technical progress in the area of plastics development (including synthetic polymers) as well as the shaping of flexible joints based on adhesives (including polyurethane and polymer adhesives) [16, 17].

Flexible adhesives based on polyurethane are the materials used for the tests described in this article. These adhesives are primarily used for reinforcements and repairs to masonry and concrete structures. Their main feature is the ability to transfer significant deformations, which is important in the event of an influence on the structure, e.g. strong winds, earthquakes or large temperature gradients [8]. In addition, polyurethane adhesives are characterized by a very low glass-transition temperature owing to the fact that, in a polymer, there is a change of its elastic-brittle or elastic mechanical state with forced elasticity into a highly elastic state [20]. This characteristic indicates that there are very stable mechanical parameters under normal atmospheric conditions, in contrast to, e.g. epoxy resins, which are characterized by a significant reduction in stiffness and strength at temperatures above +40 °C [6].

The objective of the article is an attempt to determine the value of the Mooney-Rivlin hyperelastic model parameters for the tested materials based on the results obtained in monaxial tensile and monaxial compression tests, conducted on polymers in oars and cylindrical samples, respectively, after both a week and a month-and-a-half of ripening.

2. OVERVIEW OF HYPERELASTIC MATERIAL MODELS

Polymer materials may be described using a model of hyperelastic material [7]. The material is referred to as a hyperelastic type for which there exists such a function -W (called elastic potential), which has scalar values and whose argument is a measure of material deformation such that the constitutive compound can be written as the following equation [18]:

$$T = \frac{\partial W}{\partial E} \Leftrightarrow S_{ij} = \frac{\partial W}{\partial E_{ij}} \quad i, j = 1, 2, 3 \tag{2.1}$$

In the case of isotropic materials, the deformation energy functions depend on the so-called invariants of deformation [11].

A detailed review of the models has been described in both foreign [1, 4, 11, 13] and national literature [5, 6, 8, 9, 12]. Jemioło describes in detail the simplest models of isotropic hyperelasticity with division into non-compressible, low

TESTS ON THE MECHANICAL PROPERTIES OF POLYMERS IN THE ASPECT OF AN 3 ATTEMPT TO DETERMINE THE PARAMETERS OF THE MOONEY-RIVLIN HYPERELASTIC MODEL

compressible, and compressible materials, widely implemented in the FEM finite element method.

There are several models of hyperelastic materials, one of the main (first) examples being the Mooney-Rivlin model, most frequently used in practical calculations. It predicts the behavior of isotropic materials similar to rubber material with high accuracy. Other models are, for instance, the Ogden model, the Blatz-Ko model, or the neo-Hookean model, this being Rivlin's previous proposal. For this model, we can distinguish the elastic potential for compressible and non-compressible material. The generalization of the non-compressible neo-Hookean model is the Gent material model, discussed, e.g. in the work [18]. Another model used mainly for rubber-derived non-compressible materials is the Yeoh material model.

In addition, attention should be paid to the dynamic development of numerical methods, which allows describing more and more precisely materials used not only in construction but also in interdisciplinary fields of science such as biomedicine, where a key role is played by the possibility of simulating the behavior of materials under the influence of load, causing significant deformations. Exemplary questions of the hyperelastic models' application in biomechanics, used to describe the material intended for orthopedic liners, can be found in the study [10].

3. MECHANICAL PROPERTIES OF POLYMERS TESTS AND AN ATTEMPT TO DETERMINE THE PARAMETERS OF THE MOONEY-RIVLIN HYPERELASTIC MODEL

3.1. Tested materials

The tests were conducted on samples made of four types of two-component polyurethane adhesives manufactured by Sika[®]: PBM, PMM, PM, and PSM. The adhesives used in the tests (polyurethane-based) are dedicated to adhesive joints in the scope of making flexible joints, and particularly in damaged masonry and concrete structures.

The authors notice that the mechanical properties of polymeric adhesives are closely related to the temperature and speed of deformation of the material over time. Structural changes that occur under the influence of the load affect the mechanical parameters, and this ought to be taken into consideration in the design process of flexible joints. Therefore, in the case of the tests, three deformation speeds at ambient temperature were taken into account.

Tests were also carried out in the aspect of the possibility of using polymeric adhesives for flexible joints in timber structures. For this purpose, the criterion determining a minimum polymer tensile strength not less than 0.5 MPa as well

as the limit of deformability at a level not less than 5% was assumed. This should ensure the strength of the timber-polymer connection at a level not less than 0.5 MPa. The results of these studies are described in the authors' publication [19].

3.2. Description of mechanical properties tests – monaxial testing of unlimited stretching and compression

The first kind of research conducted was a monaxial, unlimited tensile test, which was carried out on oar-shaped fittings in accordance with ASTM standards ASTMD-638-03:2004 Standard test method for tensile properties of plastics [2], PN-EN ISO 527-1:2012 Plastics - Determination of mechanical properties with static stretching - Part 1: General principles [14]. The total length of the sample was increased to 200 mm in accordance with Fig. 1 (presented below) in order to protect the fitting against breaking or sliding in the grips of the testing machine. The sample thickness was 4 mm, while the measuring section length was 50 mm. All samples were checked before testing for squareness, flatness, and edge straightness.

The samples were seasoned in a place with stable humidity of 20%. The test was conducted for two polymer ripening periods, i.e. after 7 days and after 1.5 months from the time of performance.

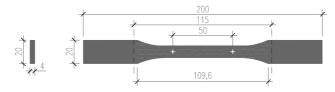


Fig. 1. The sample shape subjected to monaxial unlimited stretching

The test was carried out at ambient humidity and temperature using the following deformation rates:

- 10⁻¹•1/min for 6 elements for each of the polymers tested (PBM, PMM, PM, and PSM),
- $10^{0} \cdot 1/\min$ for 6 elements for each of the polymers tested,
- $10^{1} \cdot 1/\min$ for 6 elements for each of the polymers tested.

During the test, the force F and the corresponding increase in the length of the measuring section, and the distance between the grips of the testing machine were recorded. Tensile stress values were calculated on the basis of the formula [15]:

$$\sigma = \frac{F}{A} \tag{3.1}$$

4

TESTS ON THE MECHANICAL PROPERTIES OF POLYMERS IN THE ASPECT OF AN 5 ATTEMPT TO DETERMINE THE PARAMETERS OF THE MOONEY-RIVLIN HYPERELASTIC MODEL

where: σ – nominal value of stress in MPa, F – registered force value in N, A – initial section area in mm².

Deformation values in stretching were calculated on the basis of the formula [15]:

$$\varepsilon = \frac{\Delta L_0}{L_0} \tag{3.2}$$

where: ε – deformation, expressed in dimensionless-ness or as a percentage, ΔL_0 – increase in the length of the fitting between measuring marks in mm, L_0 – measuring section length of the fitting for the tests in mm.

Another type of research conducted was the monaxial unlimited compression test, which was performed on cylindrical samples with a diameter of 28mm and a height of 28mm, as presented in Fig. 2a. The samples were loaded with a quasi-static load in accordance with ASTM standards D 695-02a: 2002 Standard test method for compressive properties of rigid plastics [3]. The samples were seasoned in a place with stable humidity of 20%. The test was conducted for two polymer ripening periods, i.e. after 7 days and after 1.5 months from the time of performance.

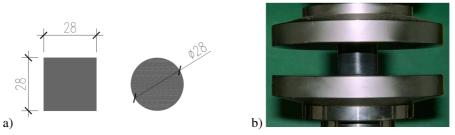


Fig. 2. Sample subjected to monaxial unlimited compression: a) sample shape; b) view of the sample in the machine

The materials were tested in a testing machine at the following deformation speeds:

- $10\%L_0 \cdot 1/\min = 10^{-1} \cdot 1/\min$ for 6 elements for each of the polymers tested (PBM, PMM, PM, and PSM),
- $100\%L_0 \cdot 1/\min = 10^0 \cdot 1/\min$ for 6 elements for each of the polymers tested,
- $1000\%L_0 \cdot 1/\min = 10^1 \cdot 1/\min$ for 6 elements for each of the polymers tested.

The values of σ_M stresses and deformations ε_M under compression were calculated using formulas (3.1) and (3.2) respectively, while the deformations in the case of the compression test have the character of relative shortening.

3.3. Mechanical properties tests results – monaxial tests of unlimited stretching and compression

The test results for monaxial unlimited stretching of polymers are depicted in Table 1, in which there are the maximum stress values and maximum deformation values in statistical terms for each of the three test speeds $(10^{-1}, 10^{0}, \text{ and } 10^{1} \cdot 1/\text{min})$. For each type of Sika[®] adhesive tested, the average value of tensile stress (m), standard deviation (s) and coefficient of variation (V) were calculated

Type of polymer		Stre	esses σ_M in N	MPa	Deformation ϵ_M in %			
Test speed		10 ⁻¹ •1/min	10 ⁰ •1/min	10 ¹ •1/min	10 ⁻¹ •1/min	10 ⁰ •1/min	10 ¹ •1/min	
PBM	m	0.3710	0.5040	0.6990	566.30	602.00	586.90	
	s	0.0786	0.0947	0.1300	82.20	28.60	5.30	
	V in %	21.18	18.78	18.59	14.52	4.75	0.91	
	m	0.9980	1.1700	1.5500	196.10	222.30	281.30	
PMM	s	0.0818	0.0908	0.0848	26.30	16.50	26.30	
	V in %	8.19	7.73	5.46	13.44	7.41	9.34	
PM	m	1.2200	1.4400	1.7100	90.90	105.00	145.50	
	s	0.1510	0.0787	0.1920	21.90	4.30	22.10	
	V in %	12.38	5.47	11.23	24.11	4.10	15.19	
PSM	m	2.2000	2.5000	3.0500	155.70	174.30	205.60	
	s	0.1260	0.1790	0.1590	5.90	7.30	22.30	
	V in %	5.73	7.14	5.22	3.77	4.17	10.85	

Table 1. The results for monaxial tensile testing

As presented in Table 1:

- for PSM polymer, the highest stresses were registered $\sigma_M w$ in case of each value test speed (2.2000, 2.5000, and 3.0500 MPa for test speed 10⁻¹, 10⁰, and 10¹·1/min, respectively)
- the smallest variability in nominal stresses appeared to be PSM polymers for test speed 10⁻¹ and 10¹·1/min (V = 5.73 and 5,22%, respectively) and PM for test speed 10⁰·1/min (V = 5.47%),
- the largest variation in nominal stresses results was observed for the PBM polymer regardless of the speed test (V size 21.18, 18.78, and 18.59%, respectively, for test speed 10^{-1} , 10^{0} , and $10^{1} \cdot 1/\text{min}$),
- for the PBM polymer, the largest deformation values ϵ_M were recorded for each test speed value (566.30, 602.00, and 586.90%, respectively, for test speed 10^{-1} , 10^0 and $10^1 \cdot 1/\text{min}$),

TESTS ON THE MECHANICAL PROPERTIES OF POLYMERS IN THE ASPECT OF AN 7 ATTEMPT TO DETERMINE THE PARAMETERS OF THE MOONEY-RIVLIN HYPERELASTIC MODEL

- the smallest variation in the size of deformations appeared to be PSM polymers for the test speed 10^{-1} and $10^{0} \cdot 1/\text{min}$ (V = 3.77 and 4.17%, respectively) and PBM for test speed $10^{1} \cdot 1/\text{min}$ (V = 0.91%),
- the largest variation in deformation results was observed for PM polymers for test speed 10^{-1} and $10^{1} \cdot 1/\min(V = 24.11 \text{ and } 15.19\%$, respectively) and PMM for test speed $10^{0} \cdot 1/\min(V = 7.41\%)$.

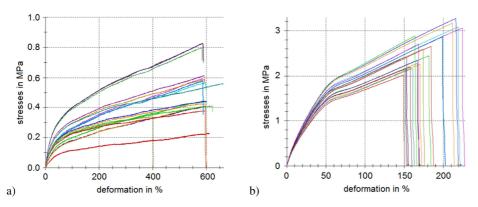


Fig. 3. Charts of stresses dependence on polymers deformation: a) PBM, b) PSM

Fig. 3 shows the resulting curves for unlimited tensile tests for selected PBM and PSM polymers. The charts illustrate the dependencies of stresses σ_M on deformation ϵ_M .

Type of polymer		Stre	esses σ_M in N	ЛРа	Relative shortening ϵ_M in %			
Test speed		10 ⁻¹ •1/min	10 ⁰ •1/min	$10^1 \cdot 1/\min$	$10^{-1} \cdot 1/\min$	10 ⁰ •1/min	$10^1 \cdot 1/\min$	
PBM	m	32.50	32.50	36.80	93.70	93.30	94.00	
	s	0.00879	0.04540	0.66600	0.50	0.40	0.40	
	V in %	0.03	0.14	1.81	0.48	0.39	0.39	
	m	32.50	32.50	35.40	88.70	88.30	88.60	
PMM	s	0.00391	0.05010	0.26400	0.20	0.20	0.10	
	V in %	0.01	0.15	0.75	0.20	0.25	0.07	
PM	m	32.50	32.50	35.10	88.40	87.70	87.60	
	s	0.00840	0.01980	0.13800	0.20	0.30	0.10	
	V in %	0.03	0.06	0.39	0.26	0.37	0.08	
PSM	m	32.50	32.50	34.40	82.50	81.50	81.50	
	s	0.00656	0.04090	0.08100	0.10	0.10	0.40	
	V in %	0.02	0.13	0.24	0.10	0.13	0.44	

Table 2. Results for the monaxial compression test

The test results for monaxial unlimited compression of polymers are shown in Table 2, in which there are the maximum stress values and maximum deformation values in statistical terms (similar to the unlimited tensile test) for each of three test speeds $(10^{-1}, 10^{0}, \text{ and } 10^{1} \cdot 1/\text{min})$. For each type of Sika[®] adhesive tested, the average value of compressive stress (m), standard deviation (s), and coefficient of variation (V) were calculated.

As presented in Table 2:

- for all polymers tested (PBM, PMM, PM, and PSM), the same average value of compressive stress $\sigma_M = 32.50$ MPa for test speed 10^{-1} and $10^{0} \cdot 1/\text{min}$ was observed,
- for the PBM polymer, the largest average stress rate was recorded in the case of the test speed $10^1 \cdot 1/\text{min}$ ($\sigma_M = 36.80$ MPa),
- PMM polymers showed the smallest variation in the size of nominal stresses for test speed $10^{-1} \cdot 1/\text{min}$ (V = 0.01%), PM for test speed $10^{0} \cdot 1/\text{min}$ (V = 0.06%), and PSM for test speed $10^{1} \cdot 1/\text{min}$ (V = 0.24%),
- the largest variety in nominal stresses results was observed for PBM and PM polymers for test speed $10^{-1} \cdot 1/\text{min}$ (V = 0.03%), PMM for test speed $10^{0} \cdot 1/\text{min}$ (V = 0.15%), and PBM for test speed $10^{1} \cdot 1/\text{min}$ (V = 1.81%),
- the maximum variety of nominal stress results was only 1.81%, which indicates a very high consistency of the results recorded during the tests of nominal stresses, with monaxial unlimited compression,
- the largest relative shortening ε_M values were recorded for each test speed value (93.70, 93.30, and 94.00%, respectively, at test 10⁻¹, 10⁰ and 10¹ · 1/min) for the PBM polymer,
- the smallest variability of relative shortening size were PSM polymers for test speed 10^{-1} and $10^{0} \cdot 1/\text{min}$ (V = 0.10 and 0.13%, respectively) and PMM for test speed $10^{1} \cdot 1/\text{min}$ (V = 0.07%),
- the greatest variety of relative shortening results was observed for PBM polymers for test speed 10^{-1} and $10^{0} \cdot 1/\text{min}$ (V = 0.48 and 0.39%, respectively) as well as PSM for test speed $10^{1} \cdot 1/\text{min}$ (V = 0.44%),
- the maximum variety of relative shortening was only 0.48%, which indicates a very high consistency of the results recorded during relative shortening tests with monaxial unlimited compression.

TESTS ON THE MECHANICAL PROPERTIES OF POLYMERS IN THE ASPECT OF AN 9 ATTEMPT TO DETERMINE THE PARAMETERS OF THE MOONEY-RIVLIN HYPERELASTIC MODEL

Fig. 4 presents the result curves for unlimited compression tests for selected PBM and PSM polymers. The charts illustrate the dependencies of stresses σ_M on relative shortening ϵ_M .

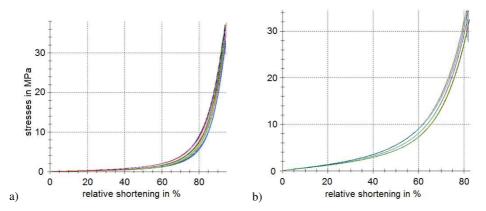


Fig. 4. Charts of stresses dependence on polymers relative shortening: a) PBM, b) PSM

3.4. Mooney-Rivlin hyperelastic model parameterization

The multinominal form of constitutive equations takes the following form:

$$W = \sum_{i+j=1}^{N} C_{ij} (\bar{I}_1 - 3)^i (\bar{I}_2 - 3)^j + \sum_{i=1}^{N} \frac{1}{D_i} (J_{el} - 1)^{2i}$$
(3.3)

For the parameter N = 1, the equation takes the form of the deformation potential energy of the Mooney-Rivlin form (M-R) with the following value:

$$W^{M-R} = C_{10}(\bar{I}_1 - 3) + C_{01}(\bar{I}_2 - 3) + \frac{1}{D_1}(J_{el} - 1)^2$$
(3.4)

where: C_{10} , C_{01} , and D_1 are constant materials which are dependent on temperature, I_1 and I_2 are the first and second invariants of the deformation matrix deviator, respectively, and J_{el} is the elastic increase in volume as a result of thermal expansion.

Assuming a constant material temperature, the elastic energy potential of the Mooney-Rivlin model is described by the equation:

$$W^{M-R} = C_{10}(\bar{I}_1 - 3) + C_{01}(\bar{I}_2 - 3)$$
(3.5)

Based on the work [9] for the model of hyperelastic material (3.3), a simplified formula of the equation (3.6) can be determined which shows the relationship

between the nominal stress and the multinomial model N = 1 (Mooney-Rivlin equation) for incompressible material.

The procedure for determining the Mooney-Rivlin model constants is based on the algorithm for minimizing the mean square error for equation (3.7), transformed from equation (3.6) as follows [20]:

$$\frac{P}{A_0} = 2 \cdot (1 - \frac{1}{\lambda^3}) \cdot (\lambda \cdot C_{10} + C_{01})$$
(3.6)

$$\sigma(\lambda) = \frac{P}{A_0} = C_{10} \cdot (2\lambda - \frac{2}{\lambda^2}) + C_{01} \cdot (2 - \frac{2}{\lambda^3})$$
(3.7)

where: P is the value of the compressive force, A_0 is the cross-sectional area of the sample, and λ is the stretching. C_{10} and C_{01} material constants are the parameters determined from monaxial tension or compression tests. After transformations, we obtain a straight line equation, explained in detail in the study [20].

Based on the results of our research conducted and described in chapter 3.3, the parameters of the Mooney-Rivlin hyperelastic model were determined for each of the polymers (PBM, PMM, PM, and PSM). The results are presented in graphic form, separately for two polymer ripening periods, i.e. 7 days and 1.5 months after the samples were made, taking into consideration three different deformation rates.

Fig. 5 illustrates the sizes of parameters obtained from experimental results in the monaxial tensile test as well as the average parameters for the Mooney line for an exemplary deformation speed which is $10^{-1} \cdot 1/\min$ for the 7-day period of polymer ripening.

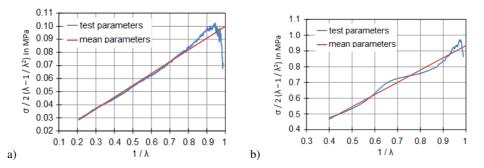


Fig. 5. Mooney graph for the polymer after 7 days of ripening: a) PBM, b) PSM

TESTS ON THE MECHANICAL PROPERTIES OF POLYMERS IN THE ASPECT OF AN 11 ATTEMPT TO DETERMINE THE PARAMETERS OF THE MOONEY-RIVLIN HYPERELASTIC MODEL

Fig.6 illustrates the values of parameters obtained from experimental results in the monaxial compression test as well as the average parameters for the Mooney line for an exemplary deformation speed of $10^{-1} \cdot 1/\text{min}$ for a 7-day period of polymer ripening.

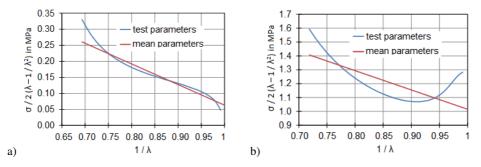


Fig. 6. Mooney graph for the polymer after 7 days of ripening: a) PBM, b)PSM

Corresponding sets of Mooney charts were obtained for the polymers tested in the 1.5-month period of polymer ripening with both monaxial tension and monaxial compression.

The parameters of the Mooney-Rivlin model for all tested polymers and the full scope of deformation for all three deformation growth rates $(10^{-1}, 10^{0}, \text{ and } 10^{1} \cdot 1/\text{min})$ are presented in tabular form. Based on the C₀₁ and C₁₀ material constants constituting the parameters of the linear function, the initial shear modulus G₀ and the initial modulus of longitudinal elasticity E₀ were estimated using the following dependencies:

$$G_0 = 2(C_{10} + C_{01}) \tag{3.8}$$

$$E_0 = 3G = 6(C_{10} + C_{01}) \tag{3.9}$$

Obtained values of the Mooney-Rivlin model parameters for the 7-day polymer ripening period are presented in Table 3.

Type of polymer and		Tensile test				Compression test			
test speed		C ₀₁	C ₁₀	E ₀	G_0	C ₀₁	C ₁₀	E ₀	G_0
PBM	$10^{-1} \cdot 1/\min$	0.090	0.010	0.600	0.200	-0.563	0.619	0.334	0.111
	10 ⁰ •1/min	0.069	0.020	0.536	0.179	-0.541	0.693	0.910	0.303
	10 ¹ •1/min	0.105	0.026	0.781	0.260	-0.670	0.881	1.266	0.422
PMM	10 ⁻¹ •1/min	0.384	0.042	2.559	0.853	-0.290	0.857	3.401	1.134
	10 ⁰ •1/min	0.376	0.057	2.597	0.866	-0.452	1.022	3.419	1.140
	10 ¹ •1/min	0.405	0.082	2.920	0.973	-0.543	1.139	3.579	1.193
PM	10 ⁻¹ •1/min	8.172	-4.661	21.065	7.022	0.560	0.380	5.640	1.880
	10 ⁰ •1/min	7.854	-4.128	22.355	7.452	0.020	0.930	5.660	1.890
	10 ¹ •1/min	7.888	-3.931	23.741	7.914	-0.150	1.150	6.050	2.020
PSM	10 ⁻¹ •1/min	0.777	0.157	5.604	1.868	-0.290	0.857	3.401	1.134
	10 ⁰ •1/min	0.697	0.185	5.293	1.764	-1.456	2.547	6.550	2.183
	10 ¹ •1/min	0.726	0.242	5.803	1.934	-1.653	2.832	7.072	2.357

Table 3.Parameter values for the Mooney-Rivlin model (7-day polymer ripening period)

4. CONCLUSIONS

Based on the results obtained from the monaxial tensile test (Table 3), it can be concluded that the PBM, PMM, PM, and PSM polymers are characterized by parameters characteristic for hyperelastic materials as the experimental curves have been satisfactorily adjusted to Mooney's straight line.

On the basis of the result values for nominal stresses σ_M and deformations ε_M in the monaxial tensile test (Table 1), the following relationship was observed between the maximum values of strain during stretching as well as deformation on the speed of deformation. With the increase of deformation speed, the value of maximum strain and deformation increased during the test. A deviation from this rule was observed only for deformation of the PBM polymer, the value of which was the highest at a speed of $10^{0} \cdot 1/\text{min}$.

In terms of the possibility to apply polymers in susceptible timber-polymer joints, it has been observed in accordance with the assumed criteria that all polymers except PBM meet the adopted criterion concerning a minimum tensile strength of 0.5 MPa.

What is more, it was observed that the obtained average tensile strength values exceeded the minimum value of 0.5 MPa and the limit deformability above 5%, with the exception of the PBM polymer, which did not reach the required level for the test speed of 10^{-1} ·1/min, and for the speed 10^{0} ·1/min, the result was just on the minimum limit.

However, the axial compression test did not confirm the positive results of the monaxial tensile tests. Based on the results obtained, it is very difficult to

TESTS ON THE MECHANICAL PROPERTIES OF POLYMERS IN THE ASPECT OF AN 13 ATTEMPT TO DETERMINE THE PARAMETERS OF THE MOONEY-RIVLIN HYPERELASTIC MODEL

classify polymers to a group of hyperelastic materials. A satisfactory result of adjusting the experimental curve from the tests to Mooney's straight line in the full scope of deformation was achieved only for the PBM polymer. Thus, it may be concluded that the tested materials should be better described by the Hooke linear-elastic model under compression.

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