## **PAPERS AT THE INTERNATIONAL CONFERENCE "INNOVATIVE DIRECTIONS OF DEVELOPMENT OF SCIENCE ON POLYMER FIBER AND COMPOSITE MATERIALS," ST. PETERSBURG, 2020.**

**MATHEMATICAL MODELING AND METHODS OF DETERMINATION OF FUNCTIONAL-USE RELAXATION-RECOVERY PROPERTIES OF POLYMER TEXTILE MATERIALS**

**A. G. Makarov, N. V. Pereborova,** UDC 539.434:677.494 **E. A. Buryak, and A. A. Kozlov**

*Mathematical models and methods of determination of functional-use relaxation-recovery properties of textile industry materials having decisive importance for comparative analysis and qualitative sampling of materials having specific properties are reviewed.*

Functional-use and performance properties of textile industry materials are based primarily on determination of the physicomechanical properties of these materials, to the study of which paramount attention must be paid.

For a comprehensive study and prediction of the functional-use and performance properties of textile industry materials to improve the quality of goods therefrom, it is proposed to conduct investigations of the basic relaxationrecovery and deformation-performance processes, i.e., relaxation and creep, which characterize the key physicomechanical properties of the materials [1-3].

It is expedient to conduct such study making use of mathematical modeling, followed by computer-aided prediction of relaxation and creep.

Although relaxation and creep processes are different in physical nature, they are, in fact, reciprocal processes harmoniously complementing each other. Because of this, the study of relaxation and deformation properties of textile industry materials relating primarily to the class of viscoelastic solids is an essential and, in some cases, an urgent task [4-6].

Relaxation process implies a change in the stress  $\sigma_t$  (or force  $F_t$ ) applied to a material over a time *t* under the action of deformative ε:

$$
\sigma = \sigma(t) = \frac{F(t)}{s}.
$$
 (1)

A key property of relaxation process is relaxation modulus  $E_{et} = \sigma/\varepsilon$ , which has two asymptotic values:modulus of viscoelasticity

$$
E_{\infty} = \lim_{t \to \infty} E_{\varepsilon t} \tag{2}
$$

and modulus of elasticity [7-9]

$$
E_0 = \lim_{t \to 0} E_{\epsilon t}.
$$
\n<sup>(3)</sup>

St. Petersburg State University of Industrial Technologies and Design. E-mail: makvin@mail.ru. Translated from *Khimicheskie Volokna*, No. 3, pp. 3-7, May-June, 2020

0015-0541/20/5203-0135© 2020 Springer Science+Business Media LLC



Fig. 1. Graph depicting experimental "family" of relaxation curves of 33.3 tex Nitron yarn at constant deformation values.

The relaxation modulus  $E_{e}$  can be modeled mathematically using increasing normalized relaxation function  $\varphi_{e}$ that acquires significance in the segment [0.1]:

$$
E_{\varepsilon t} = E_0 - \left(E_0 - E_\infty\right) \varphi_{\varepsilon t}.\tag{4}
$$

Let us take the normalized arc tangent of logarithm (NAL) that characterizes the Cauchy integral distribution as the relaxation function  $\varphi_{\text{cr}}$  [10, 14]:

$$
\varphi_{\varepsilon t} = \frac{1}{2} + \frac{1}{\pi} \arctg\bigg(\frac{1}{b_{n\varepsilon}} \ln \frac{t}{\tau_{\varepsilon}}\bigg),\tag{5}
$$

where  $\tau_{\varepsilon}$  − a characteristic of mean relaxation time and *b<sub>nε</sub>* − a characteristic of relaxation intensity.

For ease of modeling, logarithmic scale of reduced dimensionless time is used.

Instead of the mean relaxation time parameter  $\tau_{\varepsilon}$ , which is defined by the deformation-time function of shears in logarithmic time scale, it is proposed to consider the mean relaxation time  $\bar{\tau}_e$  determined by the equation

$$
\overline{\tau}_{\varepsilon} = \frac{1}{\varepsilon_2 - \varepsilon_1} \int_{\varepsilon_1}^{\varepsilon_1} \tau_{\varepsilon} d\varepsilon , \qquad (6)
$$

where  $\varepsilon_1$ ,  $\varepsilon_2$  – minimum and maximum values from the studied deformation range [15-18].

When determination only of qualitative properties of the materials is needed, switch from functional relationship to a constant is justified for mathematical modeling of relaxation-recovery properties. Such switch greatly simplifies the mathematical model, which is of considerable importance for investigating qualitative viscoelastic properties. Note that in more detailed studies of viscoelastic-relaxation processes, for example, from the spectral analysis premises, such switch to a simplified mathematical model is unjustified [19-21].

Thus, equation (4) is a mathematical model of polymer material relaxation process.

The choice of the function NAL as the foundation of the mathematical model of relaxation is not accidental because the Cauchy probability distribution, the integral function of distribution of which it is, possesses a unique property, i.e., the sum of the characteristics distributed in accordance with the Cauchy probability law also has by its distribution the Cauchy probability distribution. For textile industry materials obeyance to this law is extremely important because any complex textile object is an aggregate of simpler textile objects (yarns consist of fibers, fabrics consist of yarns, etc.). So, if the parameters of simpler textile materials obey the Cauchy probability distribution law, the parameters of more complex textile materials will also obey this distribution law [22-25].

The method of determination of functional-use relaxation-recovery properties of textile and light industry materials is based on numerical treatment of experimental "family" of relaxation curves for values of deformation constant  $\epsilon$  = const obtained on a stress relaxometer.

Name	Linear density, tex	Breaking stress, GPa	<b>Breaking</b> deformation, %
Capron-91	91	1.08	13.2
Capron-149	149	1.10	17.0
Laysan-15.6	15.6	0.52	24.3
Lavsan-114	114	0.83	11.5
Capron-187	187	0.78	15.5
Capron-189	189	0.91	16.8
Nitron $-33.3$	33.3	0.92	12.8
Capron-410	410	0.68	23.3

Table 1. Technical Characteristics of Studied Textile Yarns.

An example of the graph of "family" of relaxation curves for Nitron polymer yarn with a linear density of 33.3 tex at constant values of deformation ε is given in Fig. 1.

Introducing the formula for the studied normalized relaxation function (5)

 $\overline{\phantom{a}}$ 

$$
\varphi_{\varepsilon t} = \frac{1}{2} + \frac{1}{\pi} \arctg(W_{\varepsilon t}) = \varphi(W_{\varepsilon t})
$$
\n(7)

with the argument

$$
W_{\varepsilon t} = \frac{1}{b_{n\varepsilon}} \ln \frac{t}{\tau_{\varepsilon}} = \frac{1}{b_{n\varepsilon}} \left[ \ln \left( \frac{t}{t_1} \right) + \ln \left( \frac{t_1}{\tau_{\varepsilon}} \right) \right]
$$
(8)

and differentiating (4) further we get the expression for the derivative from the relaxation modulus:

$$
E_{\varepsilon t}^{\prime} = \frac{\partial E_{\varepsilon t}}{\partial \ln \frac{t}{t_1}} = -(E_0 - E_{\infty})\overline{r}_{\varepsilon t} = -\frac{1}{b_{n\varepsilon}} \cdot \frac{1}{\pi} \cdot (E_0 - E_{\infty}) \frac{1}{1 + W_{\varepsilon t}},\tag{9}
$$

which contains the relaxation core  $\bar{r}_{ei}$ :

$$
\overline{r}_{\varepsilon t} = \frac{\partial \varphi_{\varepsilon t}}{\partial \ln t} = \frac{1}{b_{n\varepsilon}} \cdot \frac{1}{\pi} \cdot \frac{1}{1 + W_{\varepsilon t}^2}.
$$
\n(10)

The quantity  $t_1$  means the value of the "base" time, which we will assume for convenience as  $t_1 = 60$  sec. The base time value is needed so that a dimensionless quantity lay under the sign of the logarithm.

Considering that the extreme value of the derivative from the relaxation modulus  $E_{st}$  is attained at  $W_{st} = W_t = 0$ , let us determine its corresponding characteristic value of relaxation modulus  $E_{\tau}$ :

$$
E_{\tau} = E_0 - (E_0 - E_{\infty}) \cdot 0.5 = \frac{(E_0 + E_{\infty})}{2},
$$
\n(11)

which enables determination of the width of the band of relaxation modulus values:

$$
2\Delta E_{\tau} = 2 \cdot (E_0 - E_{\infty}) = E_0 - E_{\infty} = 2 \cdot (E_{\tau} - E_{\infty}),
$$
\n(12)

whence we get the elasticity modulus value

$$
E_0 = E_\tau + \Delta E_\tau \tag{13}
$$

and the viscoelasticity modulus value

$$
E_{\infty} = E_{\tau} - \Delta E_{\tau} \,. \tag{14}
$$

137

Name	Linear density, tex (warp/weft)	Width, mm	Breaking stress, GPa	Breaking deformation, %
$CT-2$	16/16	2	1.53	18
$CT-13$	16/16	13	1.69	20
$CT-15$	29/29	15	1.82	25
$CT-16$	29/29	16	1.91	26
$CT-25$	29/29	25	4.41	25
$CT-26$	29/29	26	5.-89	20
<b>ShK</b>	303/14	5	1.96	30

Table 2. Technical Characteristics of Studied Capron Tapes.

Table 3. Technical Characteristics of Studied Technical Fabrics.

Name	Width, mm	Composition, $%$	Linear density, $text(x \alpha)$	Breaking stress, GPa	<b>B</b> reaking deformation
<b>TF-11</b>	140	Lavsan-60, Capron 40	6.7/18.8	2.9	18
$TF-13$	100	Capron-100	3.3/5.0	3.7	22
<b>TF-14</b>	140	La vsan- $100$	7.1/16.3	3.4	19
TF-16	150	Lavsan-80, Capron -20	6.9/14.3	3.6	21
<b>TF-17</b>	100	$Capron-100$	5.0/5.0	3.1	18

Hence, under the condition of attainability of the extremum at the midpoint of the band at  $W<sub>z</sub> = 0$  we can determine using expression (9) the characteristic value for the derivative from the relaxation modulus:

$$
E_{\tau} = -\frac{1}{b_{n\epsilon}} \cdot \frac{1}{\pi} \cdot (E_0 - E_{\infty}) = -\frac{1}{b_{n\epsilon}} \cdot \frac{2}{\pi} \cdot \Delta E_{\tau} \,. \tag{15}
$$

Further, from expressions (9) and (15) we get

$$
E_{\text{et}} = E_{\tau} \cdot \frac{1}{1 + W_{\text{et}}^2},\tag{16}
$$

or

$$
W_{\text{et}} = \pm \sqrt{\frac{E_{\tau}^{'} }{E_{\text{et}}^{'}} - 1} \,. \tag{17}
$$

Subtracting equation (4) from equation (11) and taking account of (12) we get:

l.

$$
E_{\rm et} - E_{\tau} = (E_0 - E_{\infty}) \cdot (0.5 - \varphi_{\rm et}) = 2\Delta E_{\tau} (0.5 - \varphi_{\rm et}), \tag{18}
$$

or

$$
\Delta E_{\tau} = -0.5 \frac{(E_{\varepsilon t} - E_{\tau})}{\varphi(W_{\varepsilon t}) - 0.5},\tag{19}
$$

whence we get the characteristic value of the relaxation intensity  $1/b_{n\epsilon}$ .

$$
\frac{1}{b_{\varepsilon t}} = -\frac{\pi}{2} \cdot \frac{E_{\tau}'}{\Delta E_{\tau}}.
$$
\n(20)

The characteristic of the mean relaxation time  $\tau_{\varepsilon}$  is obtained as a parameter of time shift of the relaxation curve obtained for the deformation ε value until coincidence with the generalized relaxation curve obtained by equation (4) [26-28].

138

Name	$E_0$ GPa	$E_{\infty}$ , GPa	$1/b_{n\epsilon}$	$\overline{\tau}_e$ , sec
Capron -91	3.9	1.2	0.43	193
Capron-149	2.8	1.5	0.41	98
Lavsan- $15,6$	13.9	6.5	0.13	138
Lavsan-114	14.0	4.0	0.09	132
Capron-187	2.4	1.5	0.16	142
Capron-189	3.1	1.2	0.42	139
Nitron $-33.3$	5.9	0.8	0.30	104
Capron $-410$	3.9	1.8	0.32	207

Table 4. Calculated Relaxation-Recovery Properties of Textile Yarns.

Table 5. Calculated Relaxation-Recovery Properties of Studied Capron Tapes.

Name	$E_0$ , GPa	$E_{\infty}$ , GPa	$1/b_{nE}$	$\bar{\tau}_{e}$ , sec
$CT-2$	65.0	28.5	0.213	942
$CT-13$	45.7	14.8	0.313	832
$CT-15$	43.8	15.7	0.334	984
$CT-16$	42.2	16.2	0.249	791
$CT-25$	40.4	18.8	0.356	718
$CT-26$	38.2	19.2	0.241	637
<b>ShK</b>	49.8	21.5	0.312	587

Table 6. Calculated relaxation-recovery properties of studied technical fabrics.



The relaxation characteristics  $E_o$ ,  $E_o$ ,  $\tau_e$ , and  $b_{ne}$  of textile industry materials obtained by the method of modeling of the functional-use relaxation-recovery properties of these materials are proposed to be used further for evaluation of their quality. Note as well that the relaxation properties obtained by mathematical modeling using NAL function obey Cauchy probability distribution, which is particularly important for materials that have a complex (composite) macroscopic structure.

The method of determination of functional-use relaxation-recovery properties of textile industry materials is applicable for a variety of textile materials, such as textile yarns (technical characteristics listed in Table 1), Capron tapes (Table 2), and technical fabrics (Table 3).

The calculated relaxation-recovery properties of textile yarns are listed in Table 4, of Capron tapes, in Table 5, and of technical fabrics, in Table 6.

A preliminary evaluation of the functional-use relaxation-recovery properties of the studied materials can be made from the calculation parameters of the functional-use relaxation-recovery properties of textile industry materials, namely, elasticity modulus  $E_0$ , viscoelasticity modulus  $E_{\omega}$ , relaxation process intensity  $1/b_{n\varepsilon}$ , and mean relaxation time  $\bar{\tau}_{\varepsilon}$  [29-32].

For example, among the textile yarns, Nitron yarn having a linear density of 33.3 tex is the one whose elastic properties are restored most fully after deformation because its viscoelasticity modulus is the lowest  $(E = 0.8 \text{ GPa})$ . The elastic properties of Lavsan yarn having a linear density of 15.6 tex are least restored because its viscoelasticity modulus is the highest  $(E_0 = 6.5 \text{ GPa})$ .

At the same time, the restoration is most intense for Capron yarn having a linear density of 91 tex because its intensity parameter value is the highest ( $1/b_{n\epsilon} = 0.43$ ). The restoration process is least intense for Lavsan yarn because its intensity parameter has the lowest value  $(1/b<sub>ne</sub> = 0.09)$ .

If the textile yarns are compared in terms of mean relaxation time, it can be stated here that among the textile yarns the elastic properties of Capron yarn having a linear density of 149 tex are restored fastest because its mean relaxation time is the least ( $\bar{\tau}_{\epsilon}$  = 98 sec) and the elastic properties of Capron yarn with a linear density of 410 tex are restored the slowest because its mean relaxation time is the longest ( $\overline{\tau}_e = 207$  sec).

Thus, application of the mathematical model and the method of determination of functional-use relaxationrecovery properties of textile industry materials is tested on a representative group of textile materials, for which predictable relaxation-recovery parameters and properties, which have a decisive importance for comparative analysis and qualitative sapling of materials possessing specific properties, were obtained.

*This work was financed within the ambit of execution of the state assignment of the Ministry of Science and Higher Education of the Russian Federation, Project No. FSEZ-2020-0005.*

## **REFERENCES**

- 1. A. G. Makarov, *Izv. Vuzov. Tekhnol. Text. Prom.*, No. 2, 12-16 (2000).
- 2. A. G. Makarov, *Izv. Vuzov. Tekhnol. Text. Prom.*, No. 2, 13-17 (2002).
- 3. A. G. Makarov, N. V. Pereborova, et al., *Khim. Volokna*, No. 5, 44-47 (2013).
- 4. V. V. Golovina, P. P. Rymkevich, et al., *Khim. Volokna*, No. 6, 33-40 (2013).
- 5. P. P. Rymkevich, A. A. Romanova, et al., *J. Macromol. Sci.*, *Part B: Physics*, 52, No. 12, 1829-1847 ( 2013).
- 6. A. G. Makarov, G. Y. Slutsker, and N. V. Drobotun, *Techn. Phys.*, 60, No. 2, 240-245 (2015).
- 7. A. G. Makarov, G. Y. Slutsker, et al., *Fiz. Tv. Tela*, 58, No. 4, 814-820 (2015).
- 8. A. G. Makarov, A. V. Demidov, et al., *Khim. Volokna*, No. 6, 60-67 (2015).
- 9. A. G. Makarov, N. V. Pereborova, et al., *Ibid.*, 68-72.
- 10. A. G. Makarov, N. V. Pereborova, et al., *Izv. Vuzov. Tekhnol. Text. Prom.*, No. 5 (359), 48-58 (2015).
- 11. A. G. Makarov, A. V. Demidov, et al., *Izv. Vuzov. Tekhnol. Text. Prom.*, No. 6 (360), 194-205 (2015).
- 12. A. G. Makarov, N. V. Pereborova, et al., *Khim. Volokna*, No. 1, 37-42 (2016).
- 13. A. G. Makarov, A. V. Demidov, et al., *Khim. Volokna*, No. 2, 52-58 (2016).
- 14. A. V. Demidov, A. G. Makarov, et al., *Izv. Vuzov. Tekhnol. Text. Prom.*, No. 1 (367), 250-258 (2017).
- 15. A. G. Makarov, N. V. Pereborova, et al., *Izv. Vuzov. Tekhnol. Text. Prom.*, No. 2 (368), 309-313 (2017).
- 16. A. G. Makarov, N. V. Pereborova, et al., *Izv. Vuzov. Tekhnol. Text. Prom.*, No. 4 (370), 287-292 (2017).
- 17. A. G. Makarov, N. V. Pereborova, et al., *Khim. Volokna*, No. 1, 69-73 (2017).
- 18. A. G. Makarov, N. V. Pereborova, et al., *Khim. Volokna*, No. 2, 59-63 (2017).
- 19. A. V. Demidov, A. G. Makarov, et al., *Khim. Volokna*, No. 4, 46-51 (2017).
- 20. N. V. Pereborova, A. V. Demidov, et al., *Khim. Volokna*, No. 2, 36-39 (2018).
- 21. A. G. Makarov, N. V. Pereborova, et al., *Khim. Volokna*, No. 3, 94-97 (2018).
- 22. N. V. Pereborova, A. G. Makarov, et al., *Khim. Volokna*, No. 4, 54-56 (2018).
- 23. A. G. Makarov, N. V. Pereborova, et al., *Khim. Volokna*, No. 4, 117-120 (2018).
- 24. N. V. Pereborova, A. G. Makarov, et al., *Khim. Volokna*, No. 5, 89-92 (2019).
- 25. N. V. Pereborova, A. G. Makarov, et al., *Khim. Volokna*, No. 6, 3-6 (2018).
- 26. N. V. Pereborova, A. G. Makarov, et al., *Khim. Volokna*, No. 6, 87-90 (2018).
- 27. N. V. Pereborova, A. V. Demidov, et al., *Izv. Vuzov. Tekhnol. Text. Prom.*, No. 2 (374), 251-255 (2018).
- 28. N. V. Pereborova, A. G. Makarov, et al., *Izv. Vuzov. Tekhnol. Text. Prom.*, No. 3 (375), 253-257 (2018).
- 29. N. V. Pereborova, A. G. Makarov, et al., *Khim. Volokna*, No. 5, 68-70 (2019).
- 30. N. V. Pereborova, A. G. Makarov, et al., *Khim. Volokna*, No. 5, 71-73 (2019).
- 31. N. V. Pereborova, A. V. Demidov, et al., *Izv. Vuzov. Tekhnol. Text. Prom.*, No. 2 (380), 192-198 (2019).
- 32. N. V. Pereborova, A. V. Demidov, et al., *Izv. Vuzov. Tekhnol. Text. Prom.*, No. 3 (381), 242-247 (2019).