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for a sustainable future

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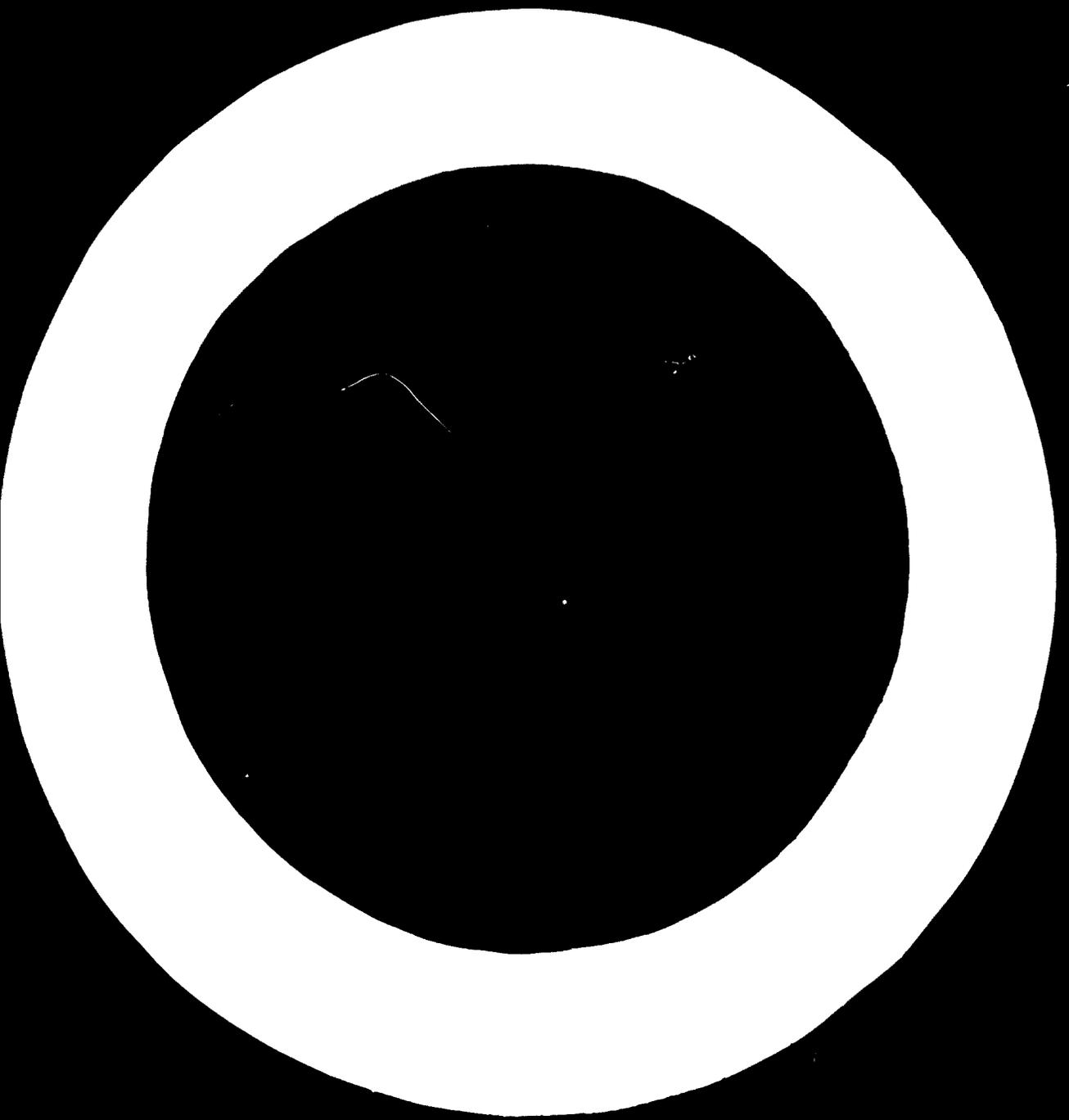
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high temperature and is resistant to dry-cleaning solvents. Due to its inherent property of non-flammability, joined with good processability and a pleasant wool-like hand, the new fibre is highly suitable for a wide range of textile applications.

We regret that some of the pages in the microfiche copy of this report may not be up to the proper legibility standards, even though the best possible copy was used for preparing the master fiche.



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The basic modulus or Young modulus is the ratio, or more exactly the first derivative of stress with respect to strain, $E = \frac{d\sigma}{d\epsilon}$, where σ is the stress and ϵ the strain. In the case of a linear elastic material, $E = \frac{\sigma}{\epsilon}$, where σ is the stress and ϵ the strain. In the case of a non-linear elastic material, E is a function of strain.

It is well known that in the case of a linear elastic material, the modulus of elasticity is a constant. In the case of a non-linear elastic material, the modulus of elasticity is a function of strain. In the case of a non-linear elastic material, the modulus of elasticity is a function of strain.

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The first two of the following observations are made on the basis of a stabilized system, i.e., the system is in a state of equilibrium. In the case of a stabilized system, the following observations are made: (1) The rate of polymerization is independent of the concentration of the monomer, (2) The rate of polymerization is independent of the concentration of the initiator, (3) The rate of polymerization is independent of the concentration of the catalyst, (4) The rate of polymerization is independent of the concentration of the solvent, (5) The rate of polymerization is independent of the concentration of the stabilizer, (6) The rate of polymerization is independent of the concentration of the inhibitor, (7) The rate of polymerization is independent of the concentration of the chain transfer agent, (8) The rate of polymerization is independent of the concentration of the chain extender, (9) The rate of polymerization is independent of the concentration of the chain terminator, (10) The rate of polymerization is independent of the concentration of the chain transfer agent.

The following observations are made on the basis of a non-stabilized system, i.e., the system is in a state of non-equilibrium. In the case of a non-stabilized system, the following observations are made: (1) The rate of polymerization is dependent on the concentration of the monomer, (2) The rate of polymerization is dependent on the concentration of the initiator, (3) The rate of polymerization is dependent on the concentration of the catalyst, (4) The rate of polymerization is dependent on the concentration of the solvent, (5) The rate of polymerization is dependent on the concentration of the stabilizer, (6) The rate of polymerization is dependent on the concentration of the inhibitor, (7) The rate of polymerization is dependent on the concentration of the chain transfer agent, (8) The rate of polymerization is dependent on the concentration of the chain extender, (9) The rate of polymerization is dependent on the concentration of the chain terminator, (10) The rate of polymerization is dependent on the concentration of the chain transfer agent.

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A particular feature of the polymerization process is known for many years, i.e., the dependence of the degree of polymerization on the polymerization temperature. Since the rate of polymerization is lower at lower temperatures, chain transfer agent and inhibitor are less effective at lower temperatures, which is favourable to the formation of polymers with higher molecular weights. Furthermore, the rate of polymerization is lower at lower temperatures, which is favourable to the formation of polymers with higher molecular weights. In the case of PVC, the formation of the radical of the polymer chain is dependent on the

of the dielectric temperature dependence. All these trends, found by means of infrared analysis and other methods, are, in principle, corresponding to the structure of the polymer, according to the corresponding theory. The results of the infrared analysis are given in the following table.

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It is well known that the reaction of vinyl acetate with an oxidizing agent yields a polymer of the type $-(CH_2-CH(O_2CCH_3))_n-$ which is soluble in water. The polymerization of vinyl acetate with an oxidizing agent is a free radical reaction. The reaction is initiated by a free radical, which attacks the double bond of the monomer, forming a radical intermediate. This intermediate then reacts with another monomer molecule, and the process repeats itself until the reaction is terminated.

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Anybody will tell you that the polymerization of vinyl acetate with an oxidizing agent is a free radical reaction. The reaction is initiated by a free radical, which attacks the double bond of the monomer, forming a radical intermediate. This intermediate then reacts with another monomer molecule, and the process repeats itself until the reaction is terminated.

with a fine brush, and a smooth surface, which may be rubbed, if desired, with a fine cloth, and with negative electricity. Several applications of the solution are necessary, and formation of a thin, uniform film is the result. The film is very hard, and is not affected by water, alcohol, ether, benzene, and lime, and is, in fact, insoluble in all these liquids.

The film is very hard, and is not affected by water, alcohol, ether, benzene, and lime, and is, in fact, insoluble in all these liquids. It is also very resistant to acids, and is not affected by dilute solutions of hydrochloric, sulfuric, and nitric acids. It is also very resistant to alkalis, and is not affected by dilute solutions of sodium hydroxide, potassium hydroxide, and ammonia. It is also very resistant to organic solvents, and is not affected by benzene, ether, and alcohol. It is also very resistant to mechanical wear, and is not affected by rubbing with a fine cloth.

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white to yellow coloration of the wool fibers, and of various types, most often during the period of the winter season.

Table 7 reports the results of the tests of the wool fibers. In the course of the tests, the wool fibers were subjected to a number of references in order to determine the effect of the various treatments on the tensile strength of the fibers. The results show that the tensile strength of the fibers is not affected by the various treatments, and that the tensile strength of the fibers is not affected by the various treatments. In fact, the tensile strength of the fibers is not affected by the various treatments, and that the tensile strength of the fibers is not affected by the various treatments.

The above table lists the results of the tests. This determination indicates the tensile strength of the fibers, and the tensile strength values reported for the wool fibers are in the following table.

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Not only does the wool fibers show a decrease in tensile strength relative to the condition of the wool fibers, but the wool fibers also show a considerable loss of weight during the various treatments. In fact, it seems that the wool fibers are not affected by the various treatments, and that the wool fibers are not affected by the various treatments.

1004.34/75
1004.34

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the removal of the residual components of the starting fibre, whereas in the case of synthetic fibres the following effects were obtained: (1) the elimination of the residual components and (2) the increase in the strength of the fibre.

The above-mentioned effects were obtained in all fibres from a range of synthetic fibres, including rayon, viscose, acetate, etc., and in all fibres containing cellulose, including cotton and wool. The above-mentioned effects were obtained in all fibres, including rayon, viscose, acetate, etc., and in all fibres containing cellulose, including cotton and wool. The above-mentioned effects were obtained in all fibres, including rayon, viscose, acetate, etc., and in all fibres containing cellulose, including cotton and wool.

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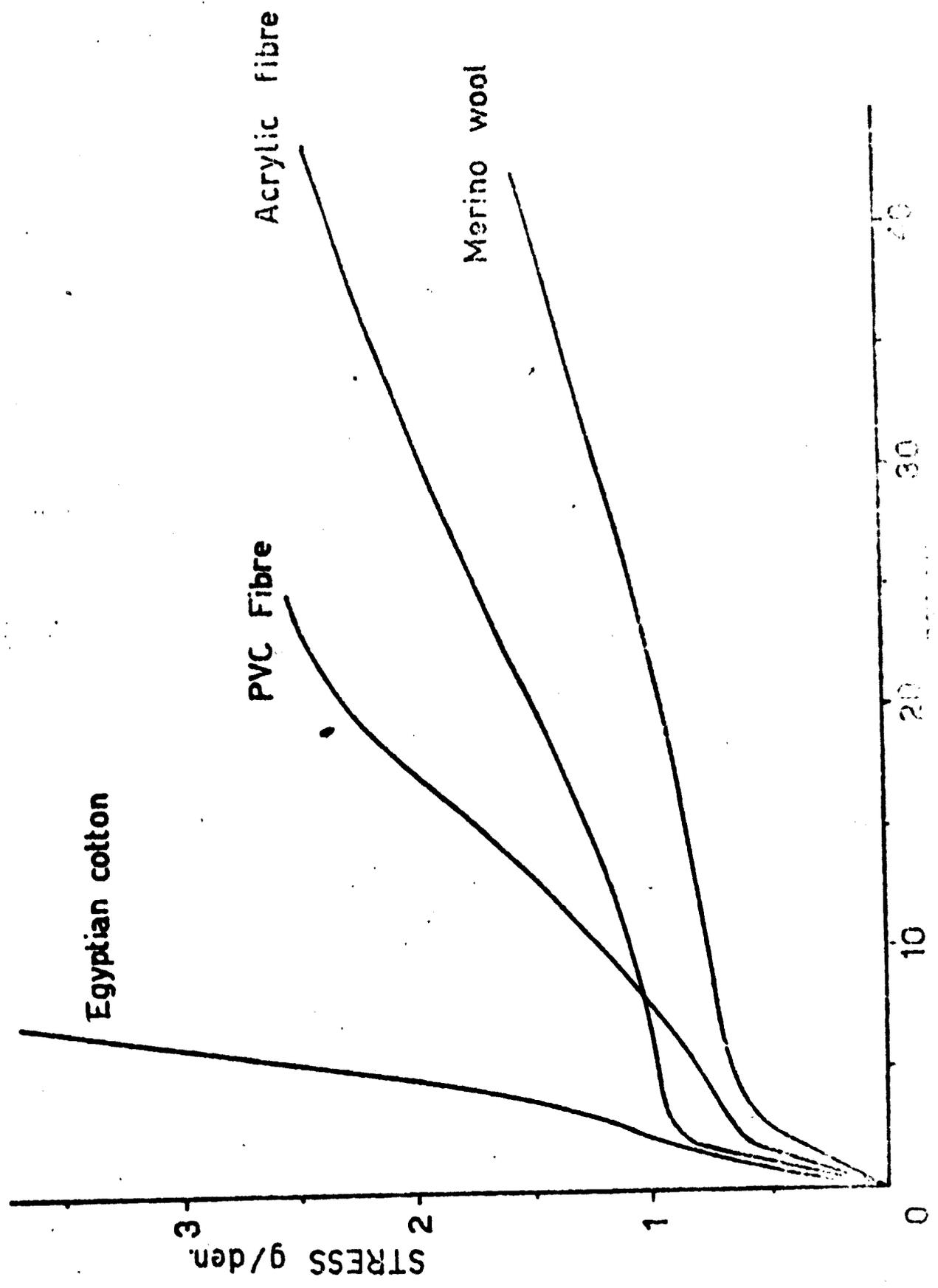
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The use of blends has, involving wool and traditional PVC fibers, such as wool, and is of particular interest because they show how the introduction of new synthetic fibers can be, as a general rule, even superior to those of the already existing materials, in certain cases, provided a way is found. In the production of high-strength fabrics with wool/wool blends, processing is done in a way which allows all the fibers to act in different ways, and a special treatment occurs the shrinkage of the traditional PVC fibers, has been the level of fur, and increased strength and durability - better and more - and thus perfectly imitating the fibers of certain animals. If fibres exist with different characteristics and blends, it becomes possible to develop the stitching by the different application of wool fibres.

Wool fibres have been in production for over one year in an industrial plant of the Textile Institute of Chetillon-lez-Louvain, which consists of 100,000 spindles and provisions for meeting the capacity and already under way.

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STRESS STRAIN CURVES AT 21°C AND 65% R.H. FIG.1



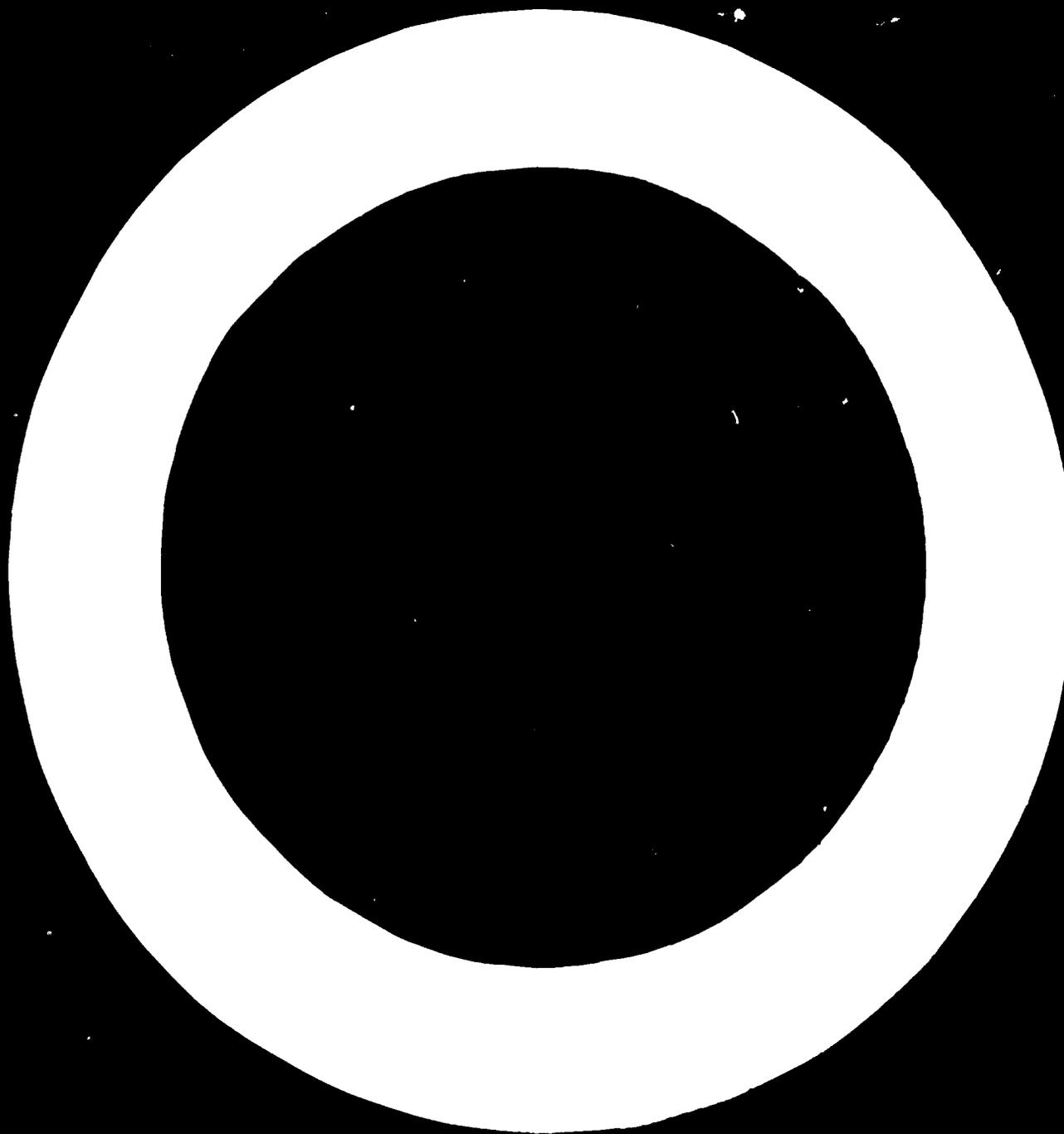
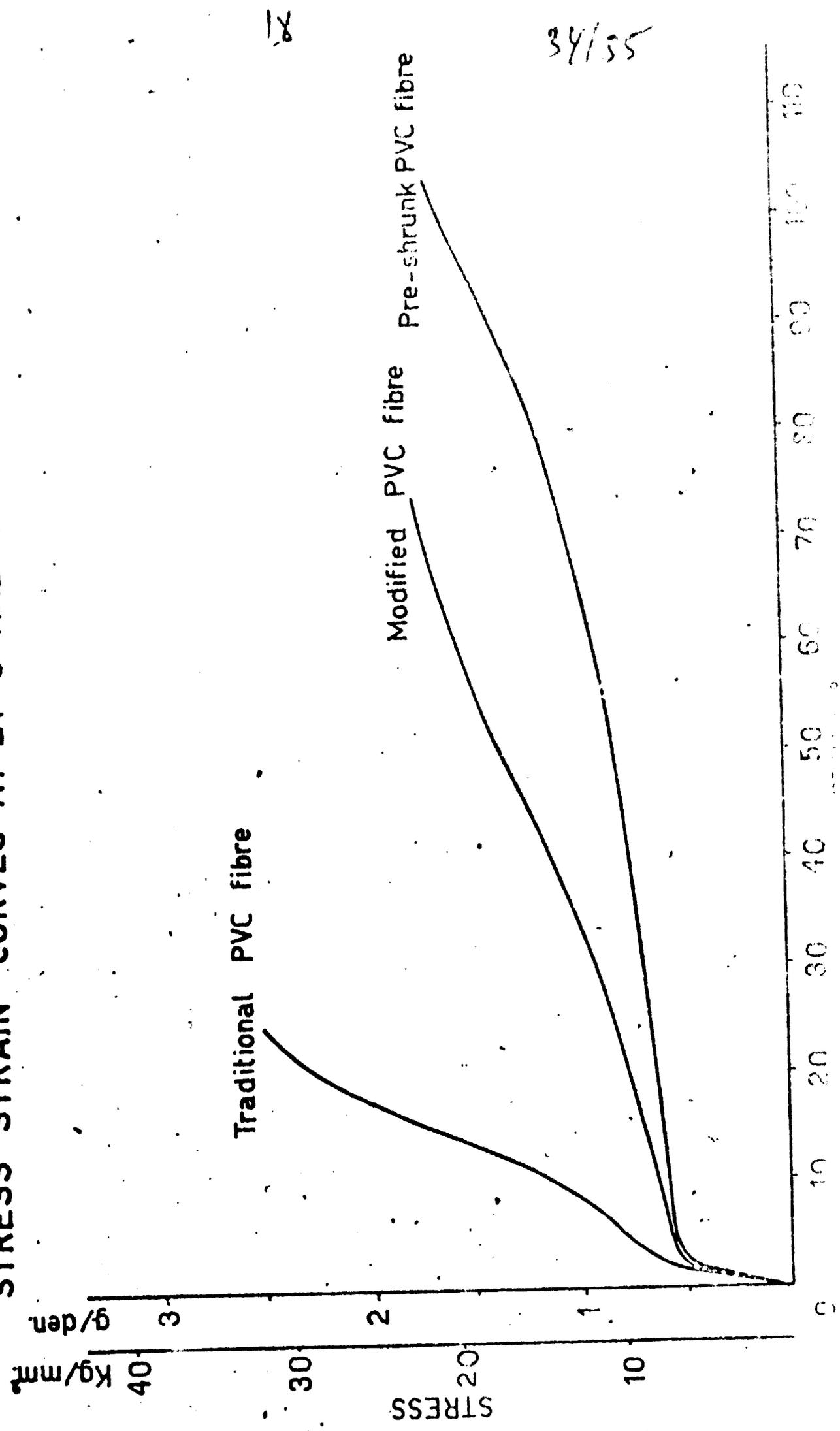
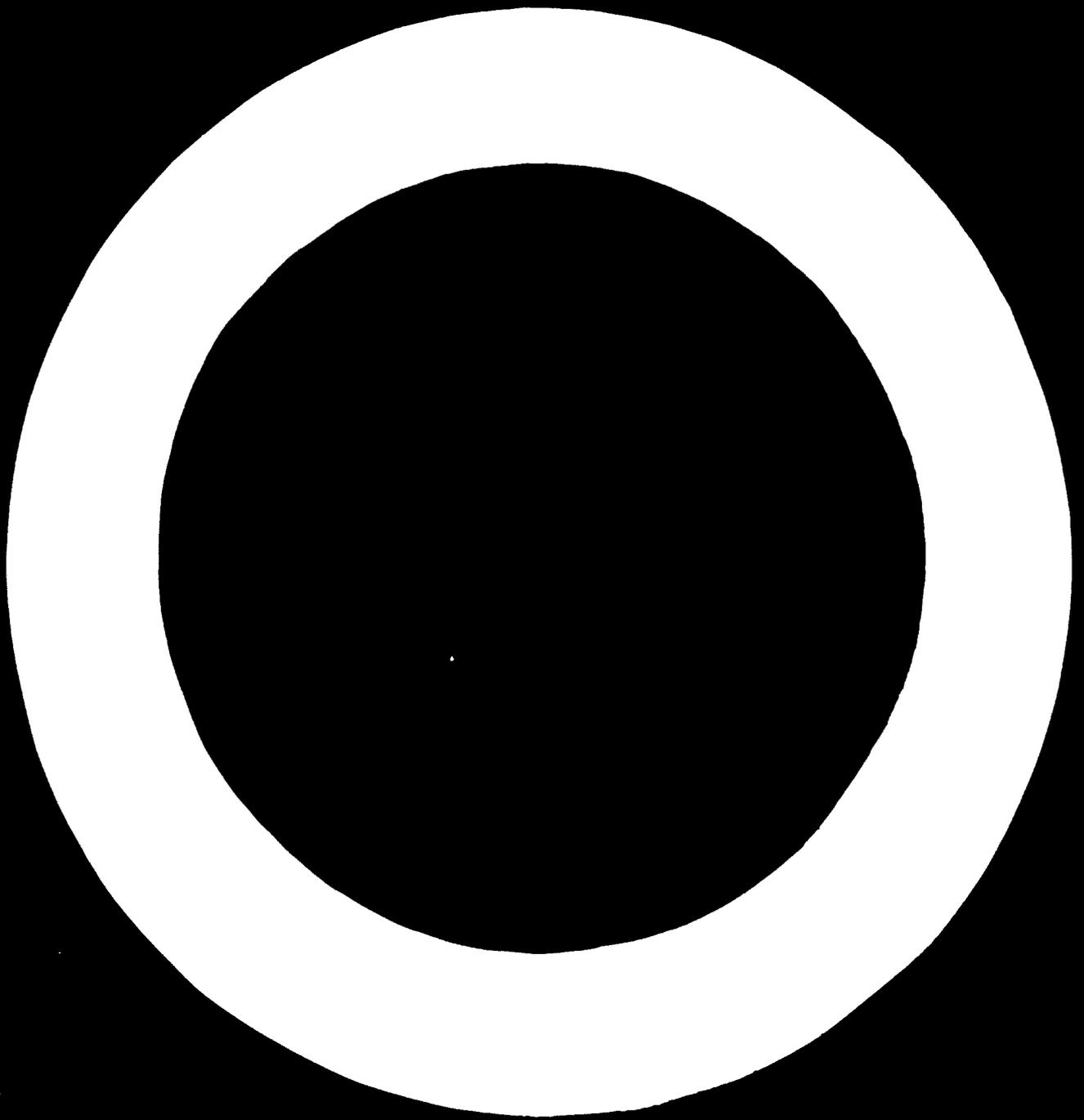


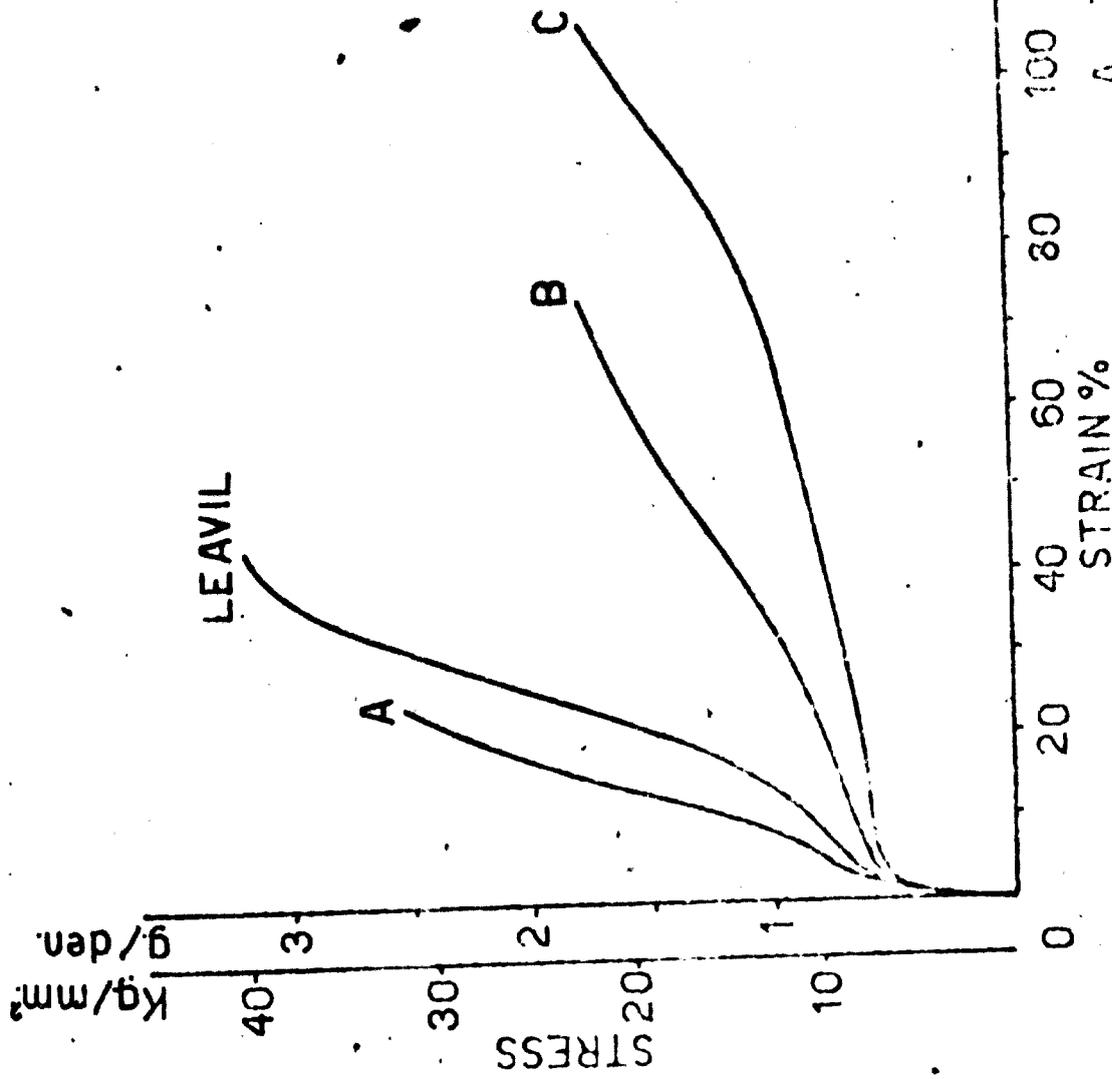
FIG. 2

STRESS STRAIN CURVES AT 21°C AND 65% R.H.

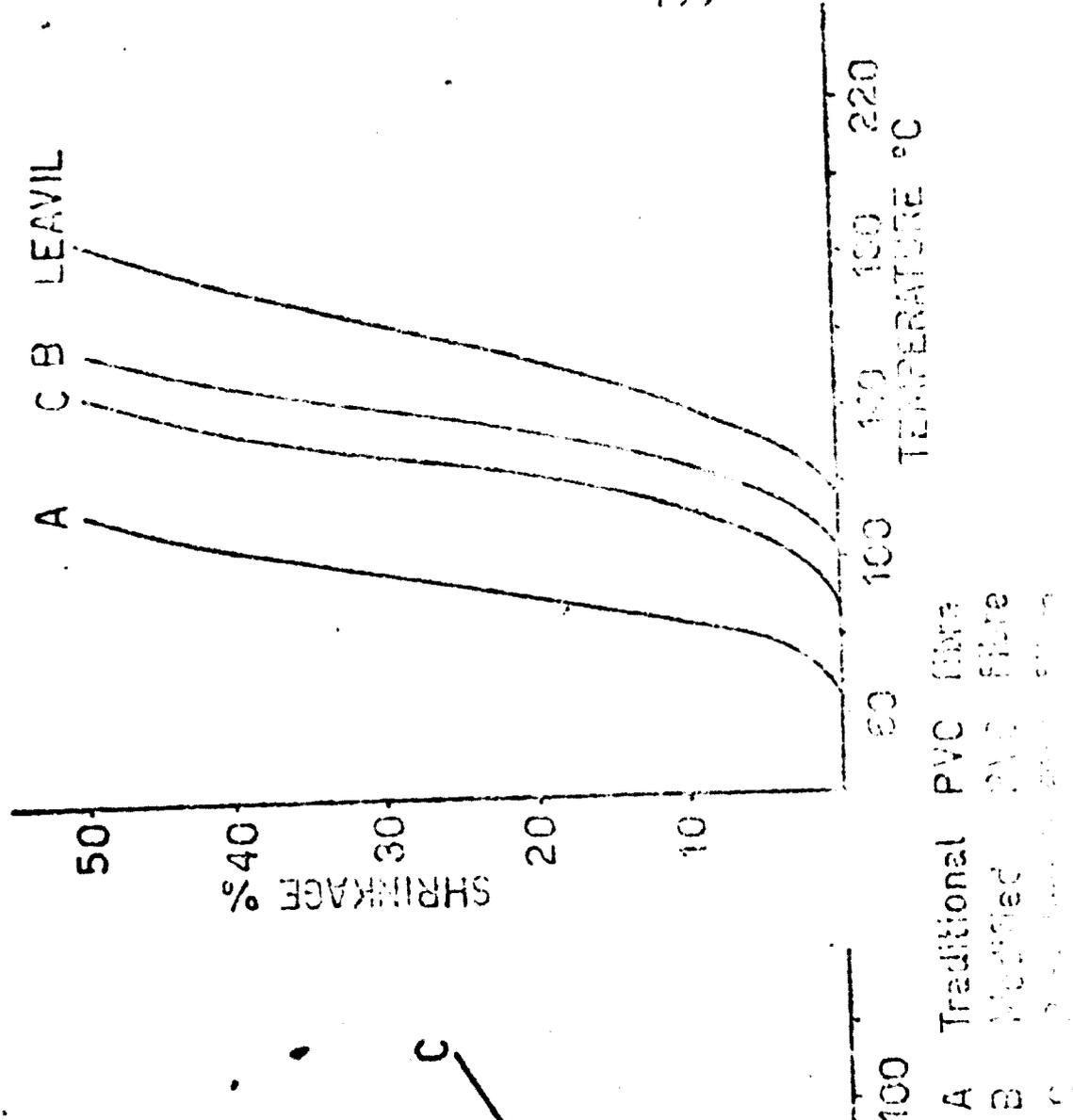




STRESS STRAIN CURVES
AT 21°C AND 65% R.H.



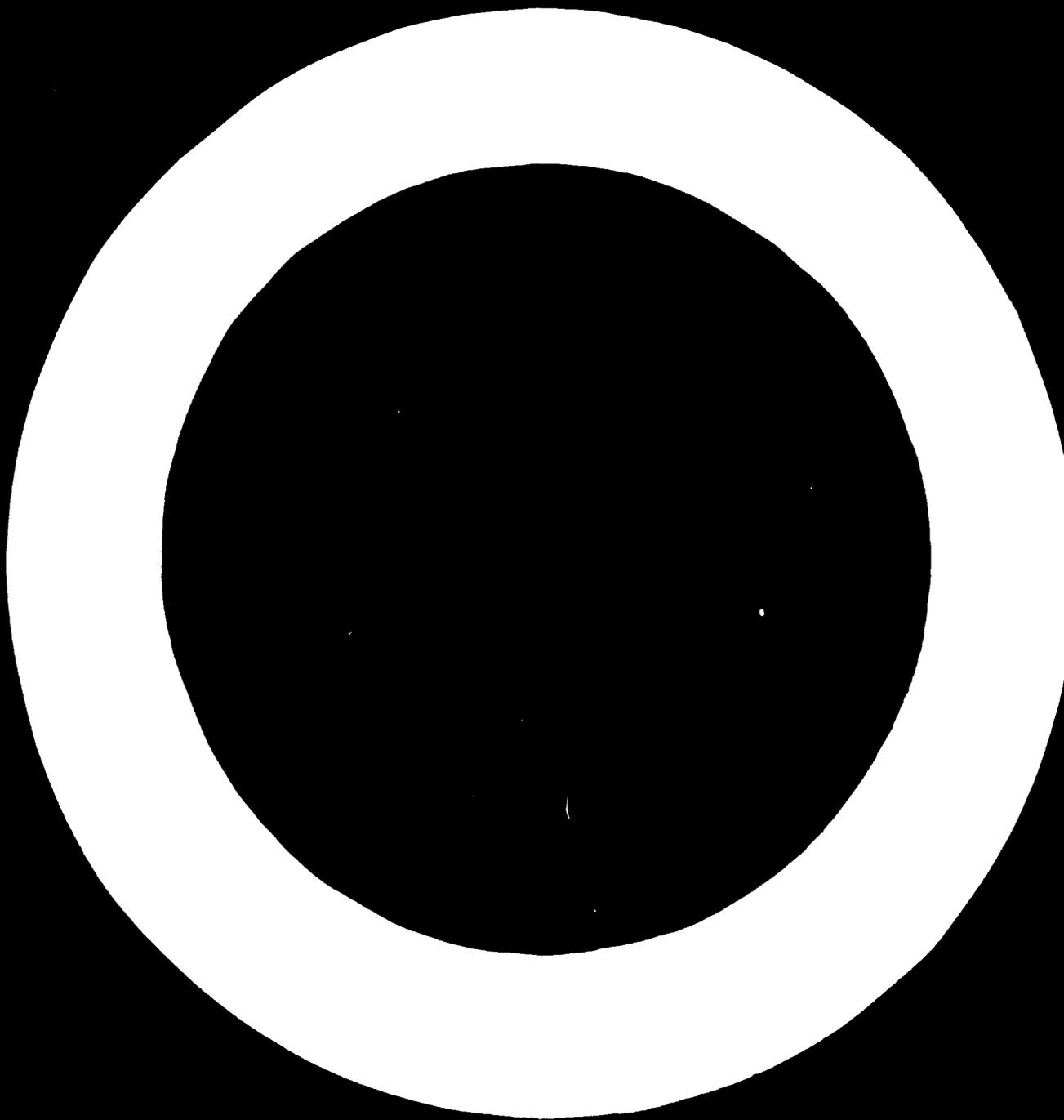
EMPIRICAL RELATIONSHIPS
OF PVC FIBRES IN AN INERT
MEDIUM (SILICON OIL)



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A Traditional PVC fibre
B Modified PVC fibre
C ...

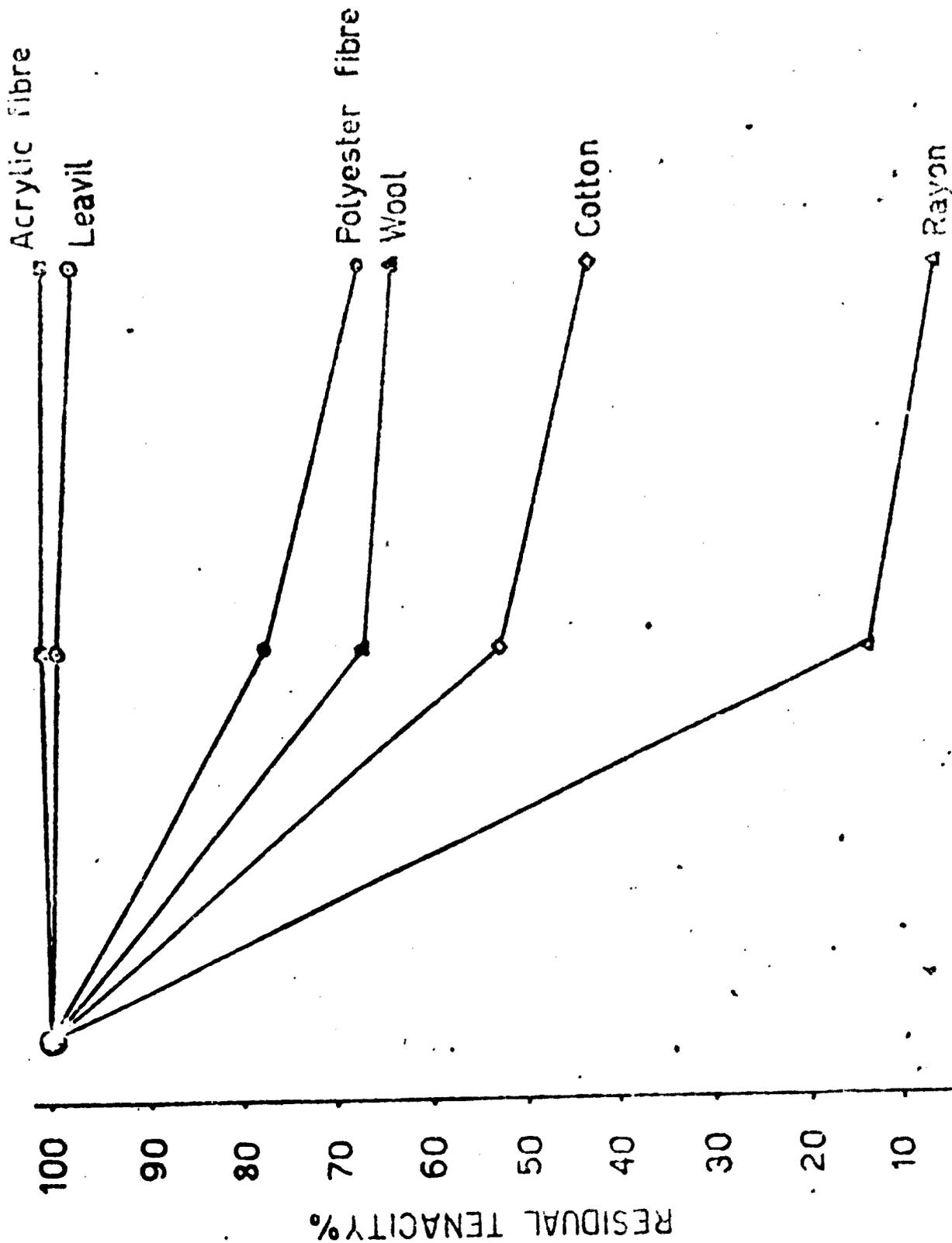


SUNLIGHT FASINESS

10000

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MONTHS OF SUNLIGHT

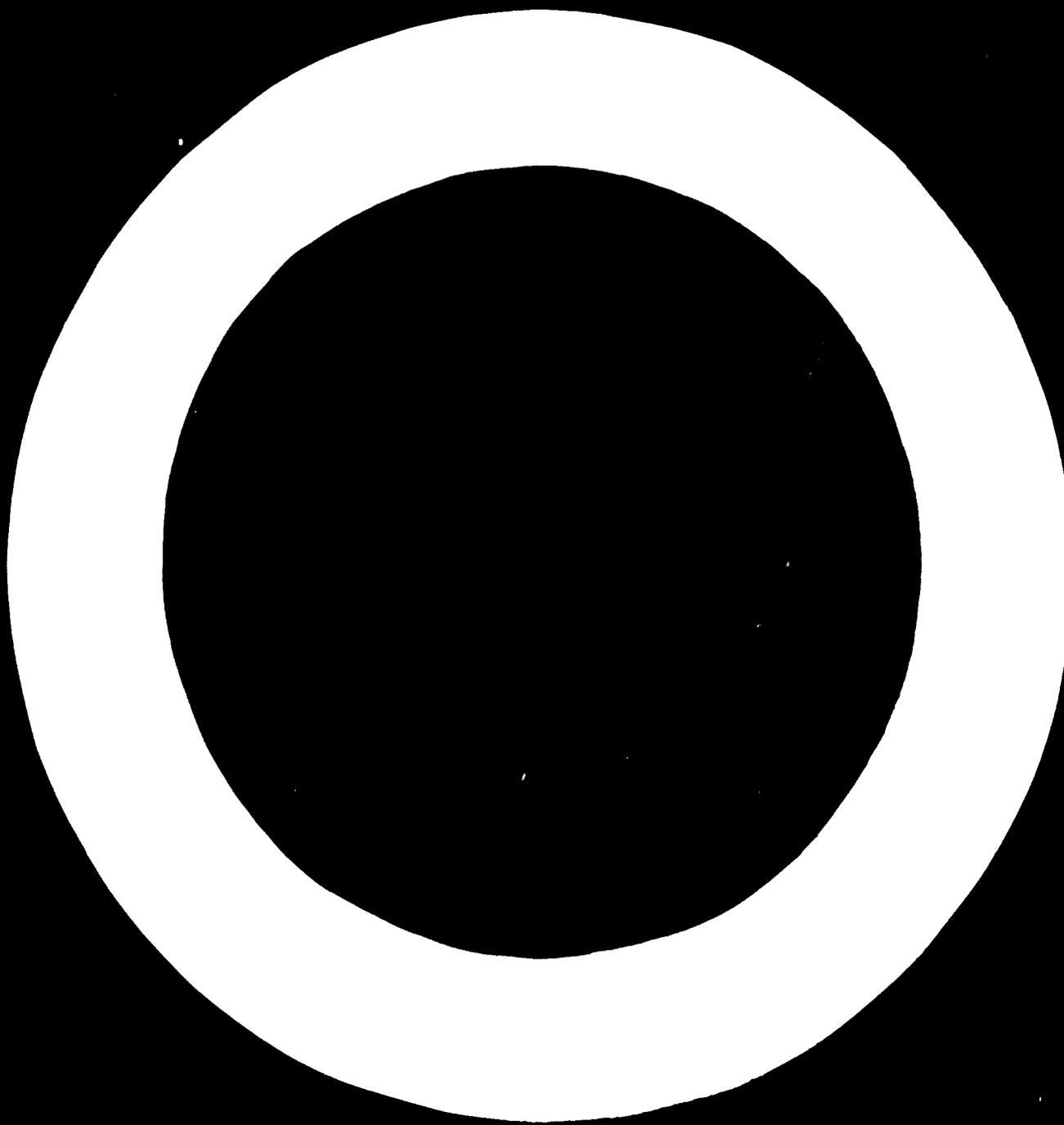
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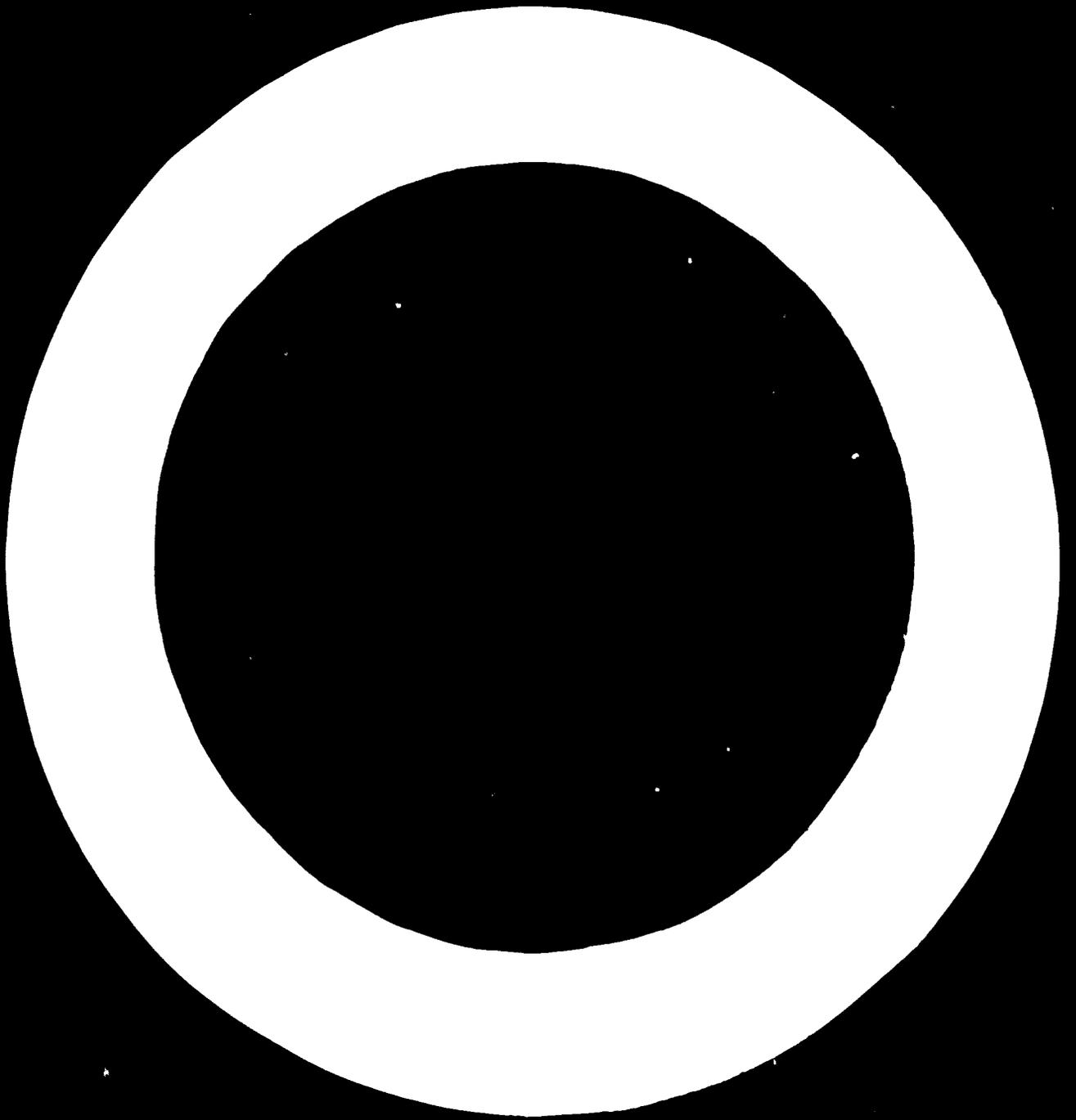


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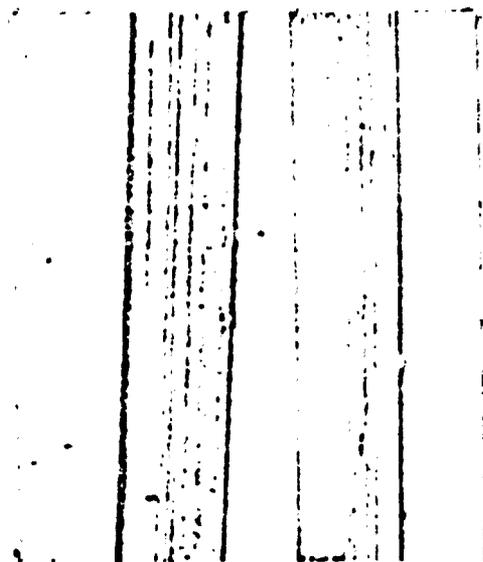
Fig. 5 - HEAVN resistance to chemical agents

Solution	Test conditions	Weight loss %	Tensile strength p/dia	Elong. %	Charitic Modulus p/dia	Color
-	as such	-	3.3	40	30	-
2% HNO ₃	168 hours at 20°C	0.45	2.85	50	30	-
2% H ₂ SO ₄	168 " " 20°C	0.20	3.3	50	29	-
2% HCl	168 " " 20°C	0.00	3.3	47	30	-
2% NaOH	168 " " 20°C	0.20	3.2	49	27	-
2% H ₂ SO ₄	1 hour to boil	1.5	3.3	49.5	28	-
30% H ₂ SO ₄	1 " "	1.5	3.25	50	30	-
2% NaOH	1 " "	0.8	3.2	50	27	Yellow
30% NaOH	1 " "	0.8	2.8	50	30	Yellow





LEAVII. fibre cross-section



LEAVII. fibre surface

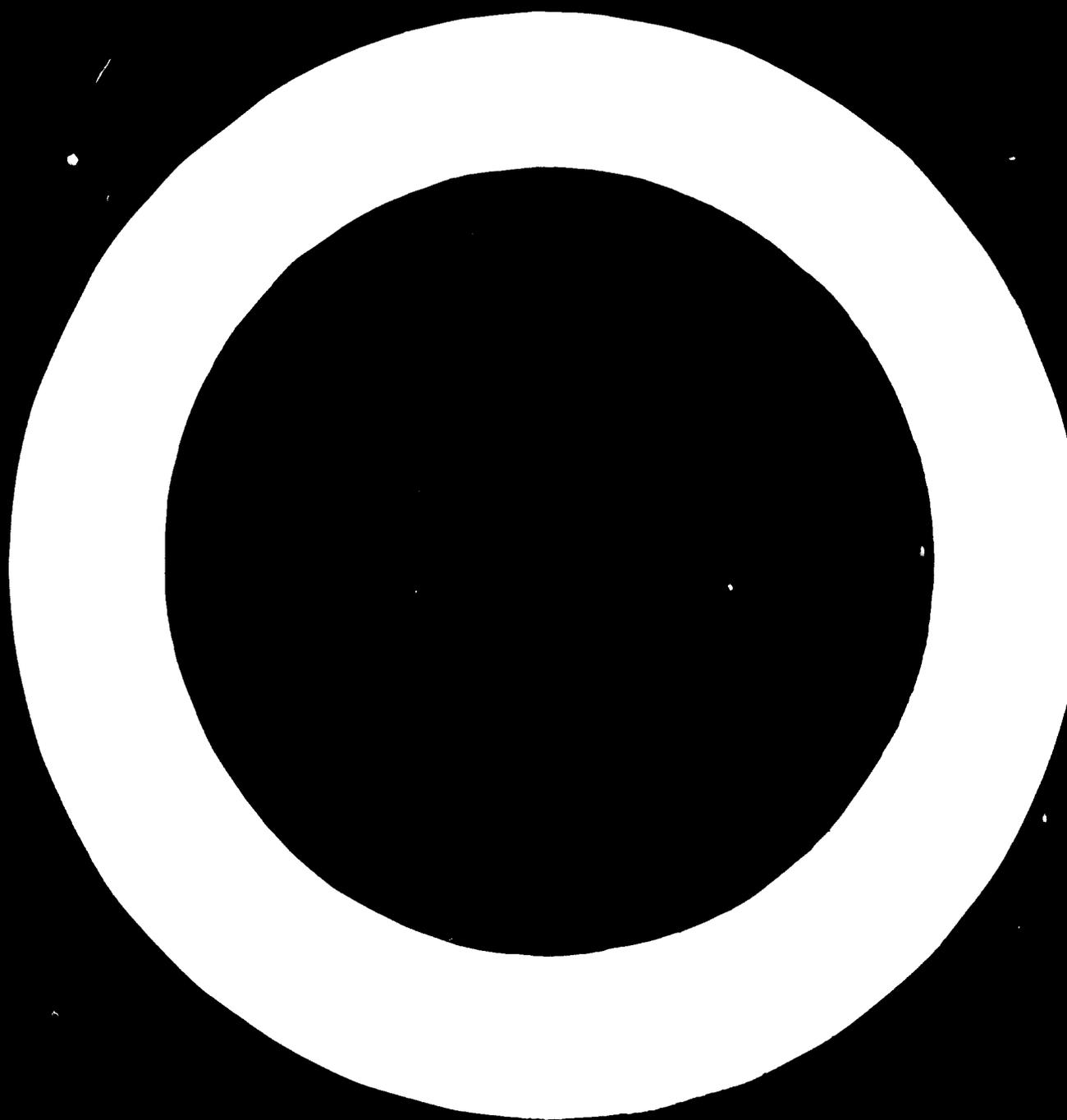
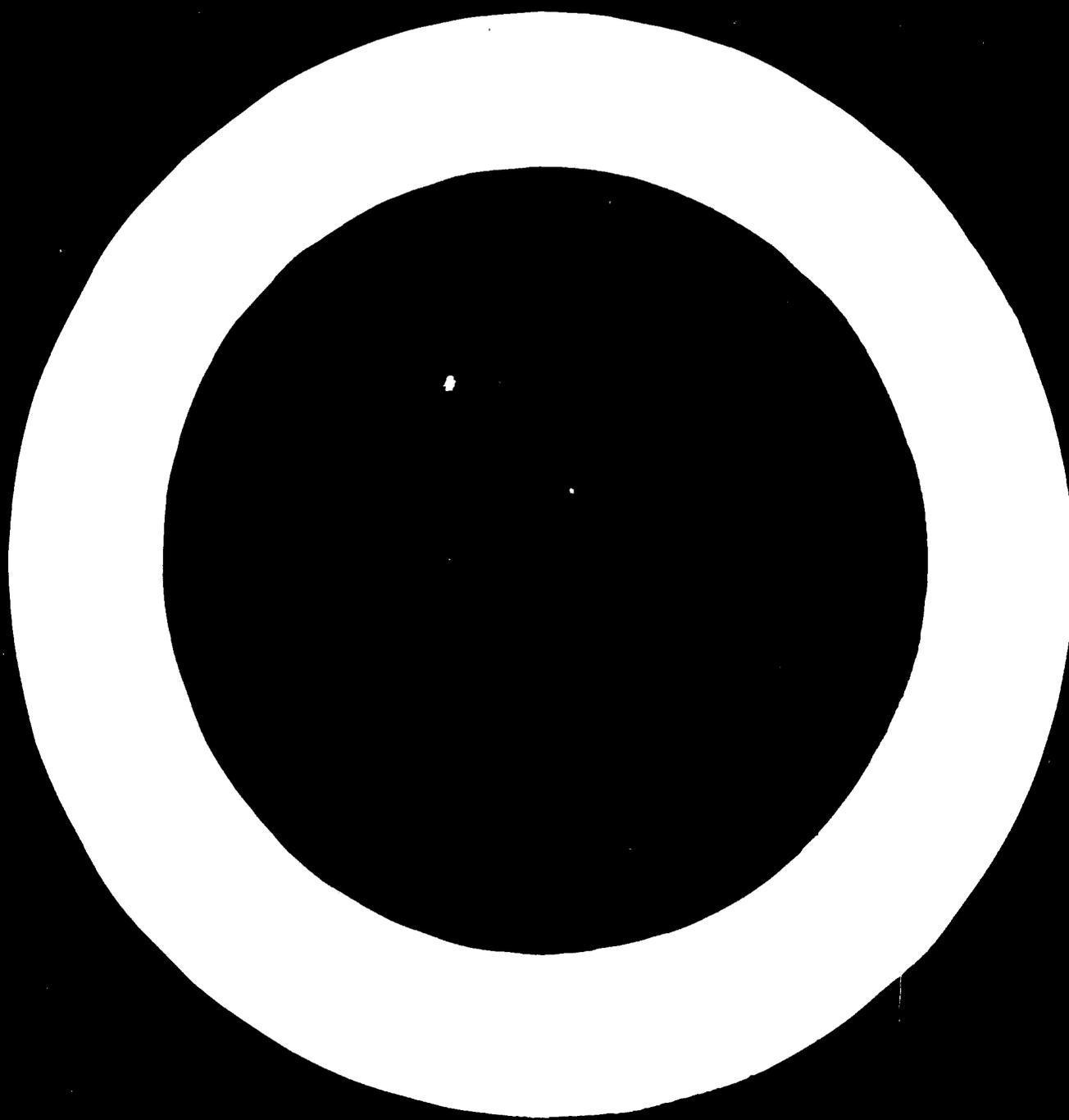


Fig. 7 - LENVIN fibre properties (commercial deniers)

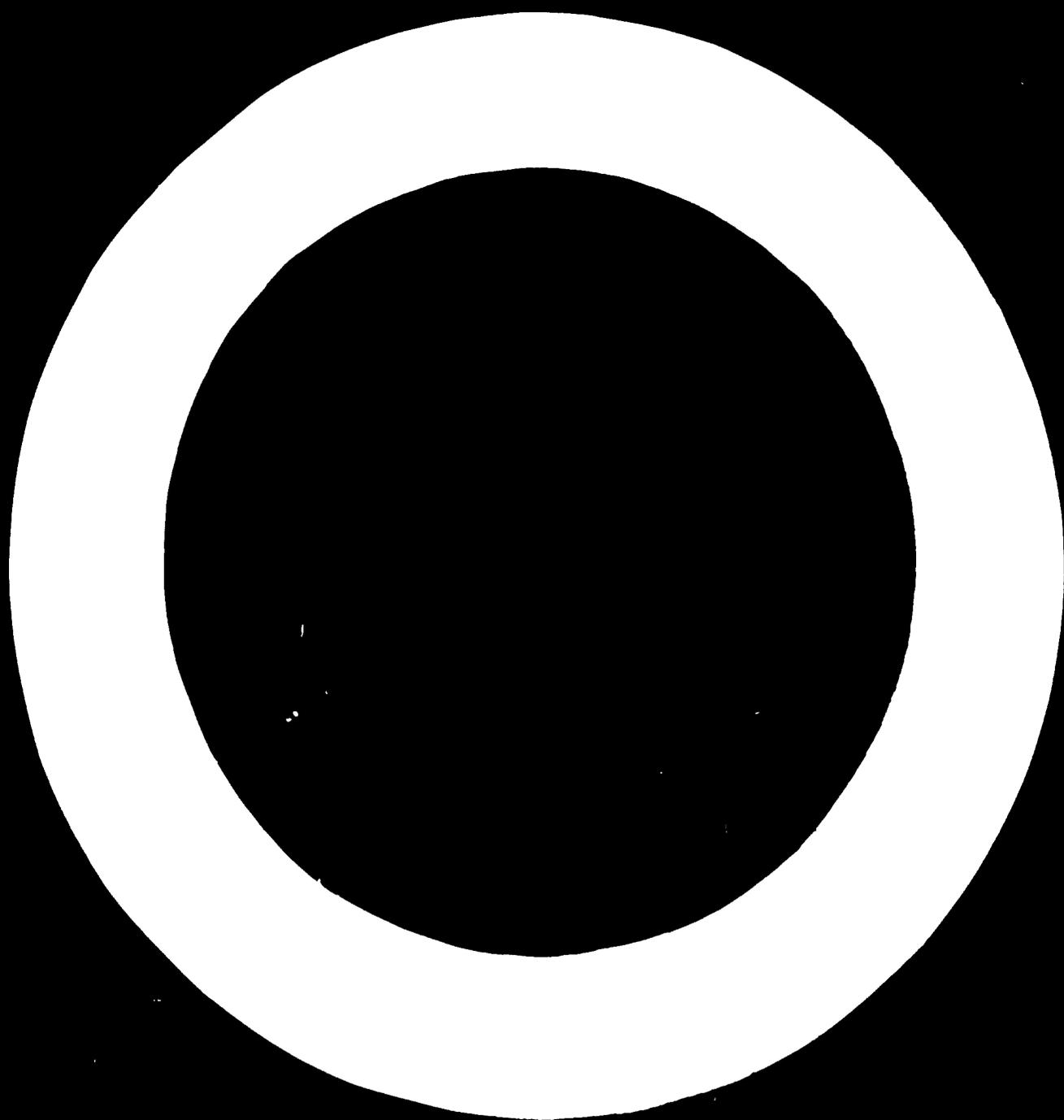
Denier	Tenacity g/den	Elongation %	Elastic Modulus g/den	Yield Point g/ten	Yield Tenacity g/den	Stress No.
1.5	3.30	36	30	0.60	2.8	85
2.0	3.20	38	29	0.55	2.7	80
3.0	3.10	40	28	0.55	2.5	80
5.0	2.9	46	27	0.50	2.1	76
8.0	2.8	50	27	0.50	1.9	70
15.0	2.6	60	27	0.50	1.7	66
25.0	2.3	70	24	0.50	1.4	63



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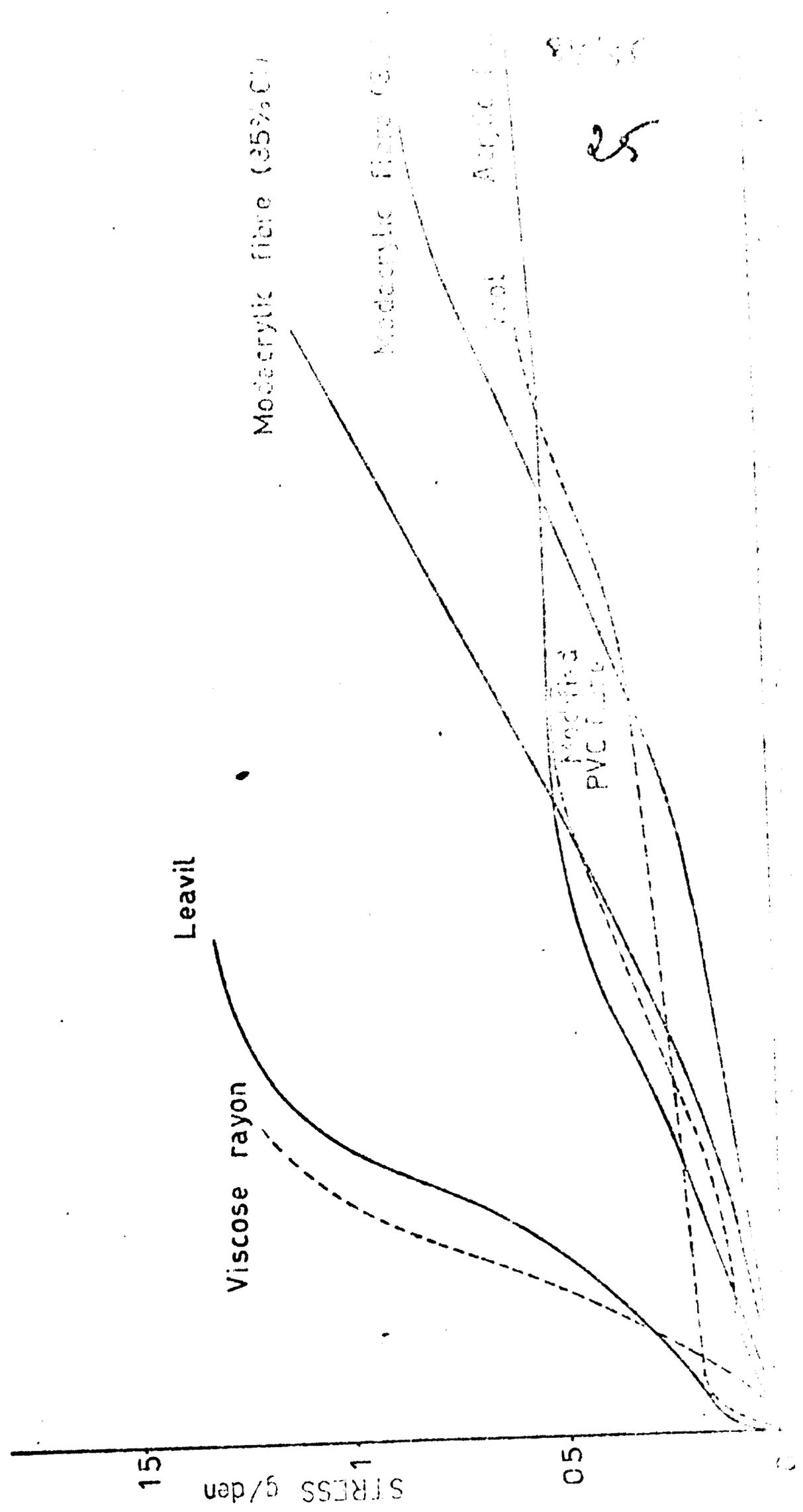
Fig. 8 - Properties of LEAVIL and other fibres under standard conditions and in water.

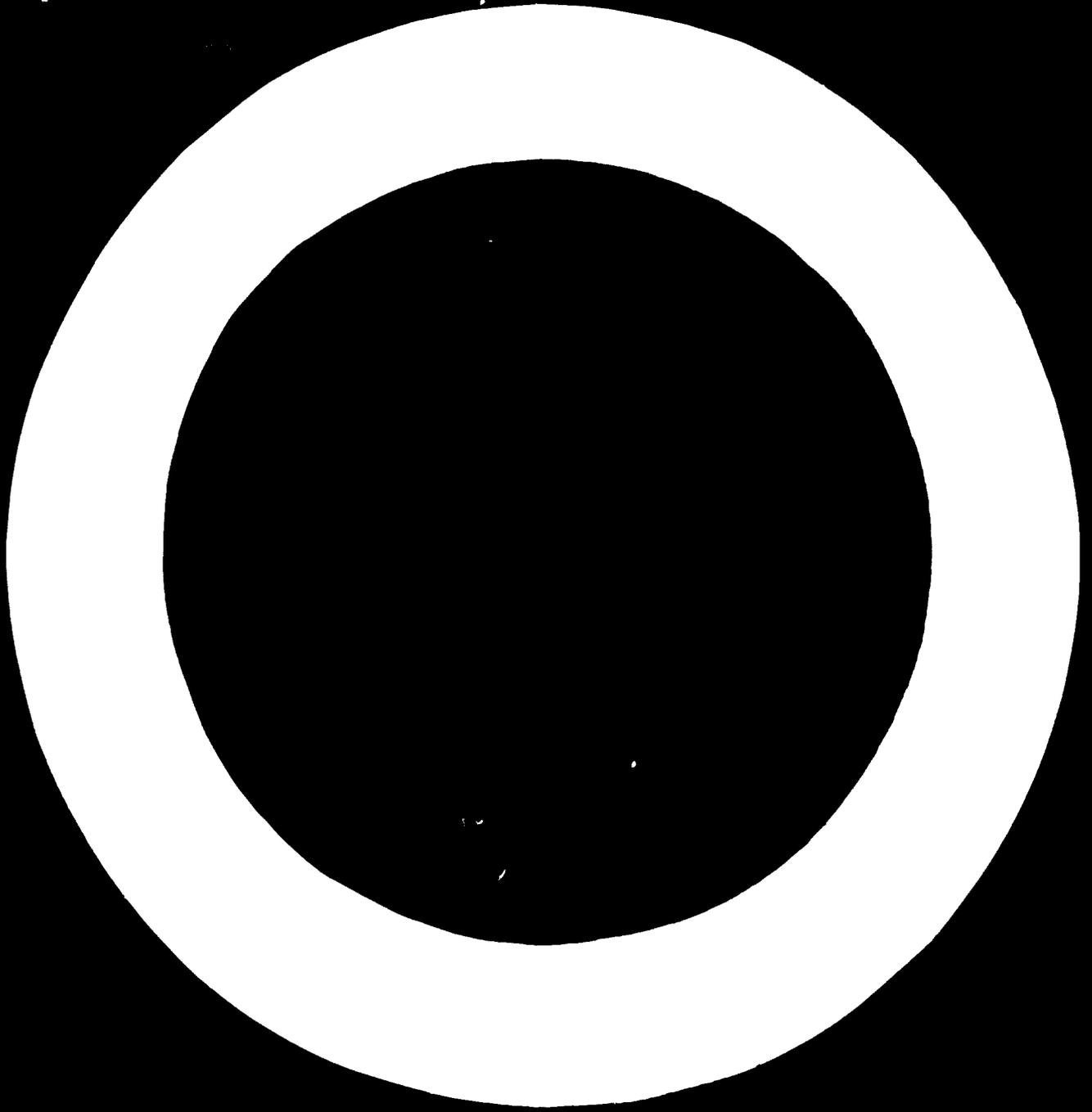
FIBRE	21°C, 65% R.H.		21°C in water		90°C in
	Tenacity g/den	Elong. %	Tenacity g/den	Elong. %	Tenacity g/den
Leavil	3.15	41	3.1	45	1.25
Traditional IVC fibre	2.5	24	2.3	27	0.8
Pre-shrunk IVC fibre	1.7	105	1.6	105	0.7
Modified IVC fibre	1.7	72	1.6	74	0.5
Modacrylic fibre (35% chlorine)	2.5	50	2.3	60	1.1
Modacrylic fibre (30% chlorine)	2.9	44	2.3	47	0.8
Acrylic fibre	2.4	43	1.8	46	0.6
Viscose rayon	2.5	18	1.4	24	1.2
Wool	1.3	39	1.2	57	0.5



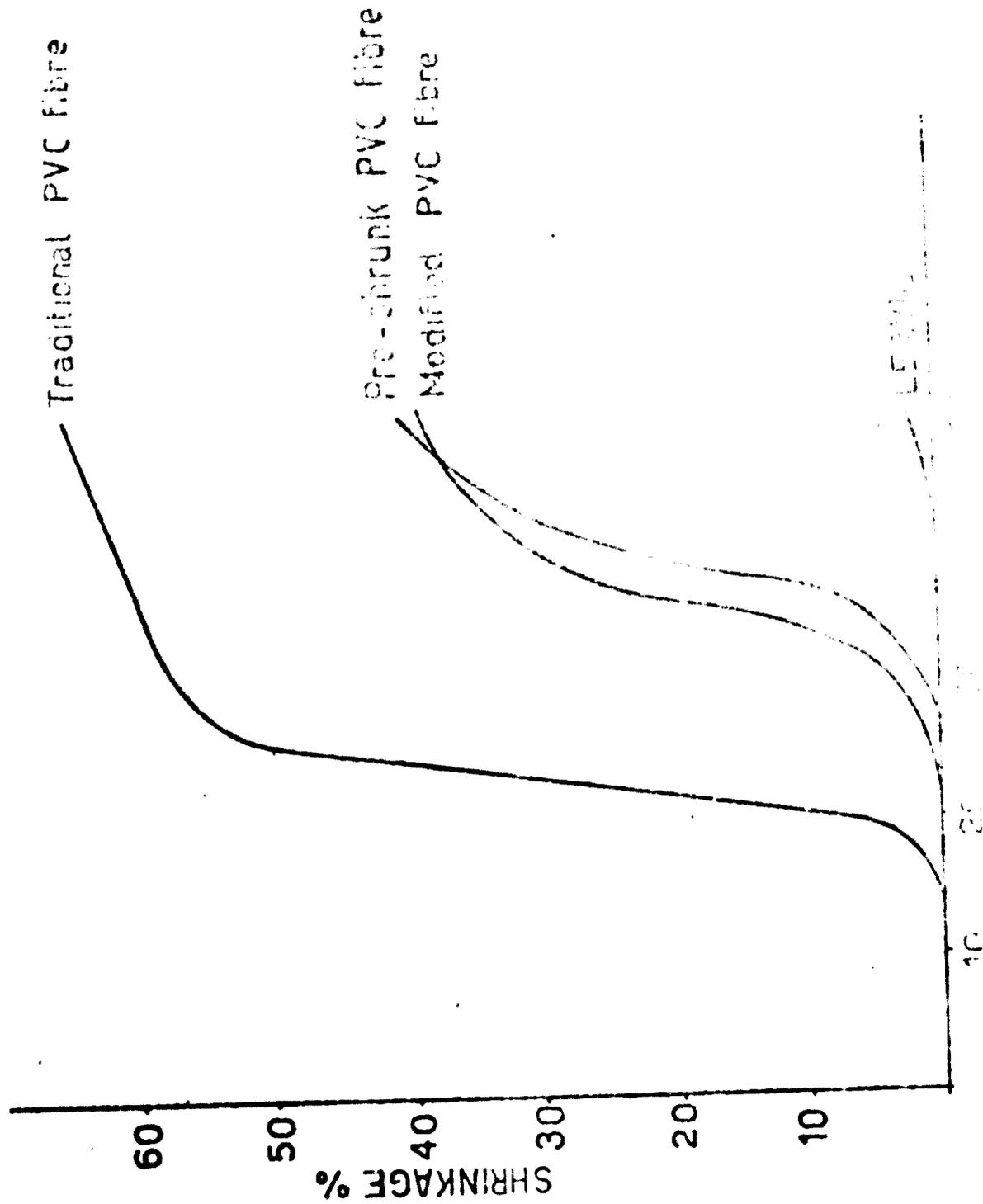
STRESS STRAIN CURVES IN WATER AT 20°C

FIG. 9





DIMENSIONAL STABILITY OF PVC FIBRES IN TRICHLOROETHYLENE FIG.10



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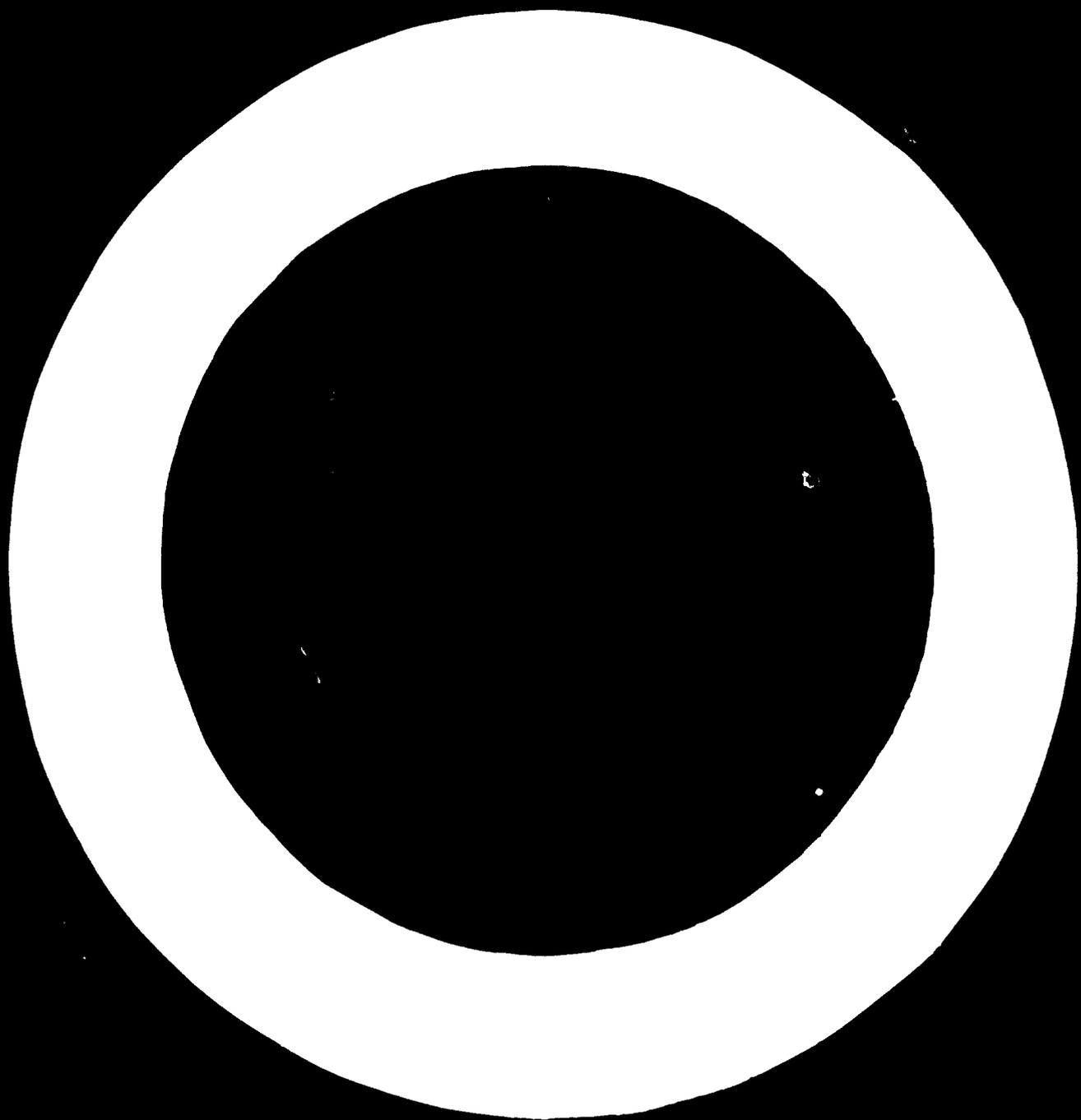
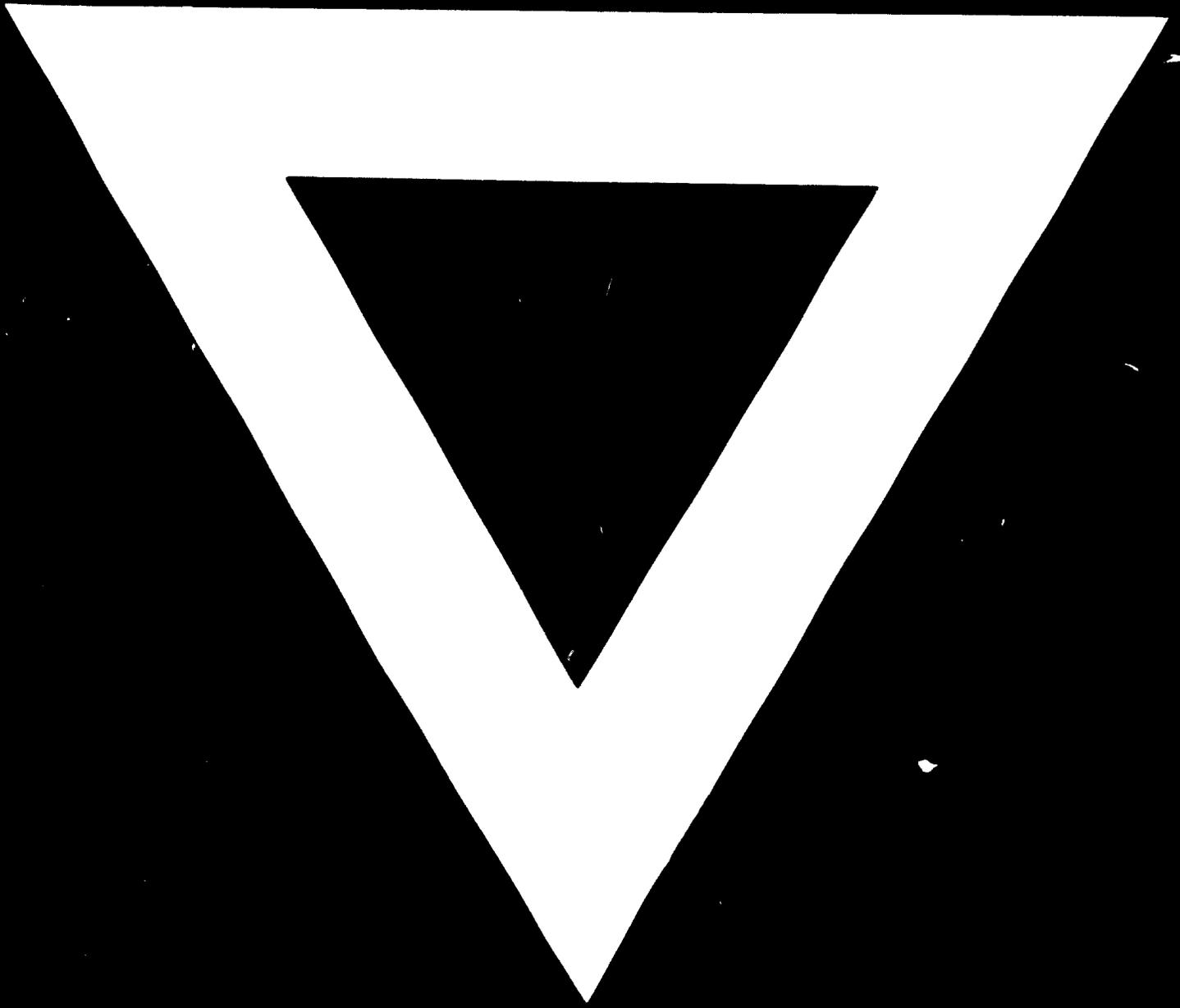


Fig. 12 - Dispersed dyes selected for dyeing UPMIL fibre

Acetaminon Light Yellow 37	(FRANCOLOR)	Resolin Red 33	(BAYCO)
Artisil Yellow 200 Peron 22-201	(SANDOZ)	Resolin Scarlet 301	(BAYCO)
Artisil Brilliant Yellow 201 Peron 27-6001	(SANDOZ)	Tersotile Redine 71	(AGNA)
Artisil Yellow 21 Peron 27-71	(SANDOZ)	Artisil Violet 200 Peron Brilliant 7-2001	(SANDOZ)
Palanil Yellow 30	(BASF)	Palanil Brilliant Violet 4701	(BASF)
Palanil Yellow 37	(BASF)	Samaron Violet 4001	(HOECHST)
Palanil Blue Yellow 20	(BASF)	Resolin Blue-Violet 201	(BAYCO)
Samaron Yellow 4001	(HOECHST)	Peron Brilliant Violet 21	(SANDOZ)
Samaron Brilliant Yellow 401	(HOECHST)	Artisil Violet 71	(SANDOZ)
Resolin Yellow 20	(BAYER)	Tersotil Violet 71	(AGNA)
Resolin Yellow 21	(BAYER)	Esterequinon Light Blue 31	(FRANCOLOR)
Resolin Yellow 21	(BAYER)	Peron Blue 2-11	(SANDOZ)
Spectyl Yellow P-501	(GHEG)	Palanil Brilliant Blue 301	(BASF)
Microsetile Yellow 37	(AGNA)	Palanil Blue 7	(BASF)
Samaron Blue Yellow 20	(GHEG)	Palanil Blue 701	(BASF)
Samaron Brilliant Orange 201	(HOECHST)	Samaron Blue 401	(HOECHST)
Nibcol Orange 40	(GIBA)	Samaron Blue 200	(HOECHST)
Tersotil Light Orange 301 Extra	(FRANCOLOR)	Peron Dark Blue 7-201	(SANDOZ)
Tersotil Light Orange 301 Extra	(FRANCOLOR)	Peron Dark Blue 7-201	(SANDOZ)
Palanil Orange 3	(BASF)	Resolin Blue 201	(BAYCO)
Samaron Orange 200	(HOECHST)	Resolin Blue 201	(BAYCO)
Samaron Orange 20	(HOECHST)	Spectyl Blue P-50	(GHEG)
Samaron Brilliant Orange 200	(HOECHST)	Spectyl Blue P-5 conc.	(GHEG)
Acetaminon Light Pink 101	(FRANCOLOR)	Microsetile Blue 20	(AGNA)
Palanil Brilliant Pink 201	(BASF)	Peron Yellow-Brown S-201	(SANDOZ)
Samaron Pink 201	(HOECHST)	Palanil Yellow-Brown 201	(BASF)
Peron Brilliant Red S-01	(SANDOZ)	Tersotile Yellow-Brown 201	(AGNA)
Palanil Brilliant Red 301	(BASF)	Tersotile Brown 301	(AGNA)
Palanil Red 37	(BASF)	Peron Brown S-301	(SANDOZ)
Peron Red 2-201	(SANDOZ)	Peron Brown S-201	(SANDOZ)
Peron Brilliant Red 7-201	(SANDOZ)	Peron Black S-201	(SANDOZ)
Resolin Brilliant Red 33	(BAYCO)	Peron Black S-201	(SANDOZ)
		Tersotile Black 21	(AGNA)





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