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MINERAL ASBESTOS SUBSTITUTES 1'

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United Nations Industrial Development Organization

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SUMMARY

SUBSTITUTE PLASTIC MATERIALS

by

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In Czechoslovakia the field of plastic materials is developing rapidly. The goal of producing 15-20 per cent of the total production of plastics in the country is a realistic one. For this purpose it is necessary to develop substitute plastic materials which are suitable for the manufacture of plastic parts and components. The development of substitute plastic materials is a complex task which requires a detailed study of the properties of the plastics to be replaced and the properties of the substitute materials. The development of substitute plastic materials is a complex task which requires a detailed study of the properties of the plastics to be replaced and the properties of the substitute materials.

This report is a summary of the work done in this field in the Czechoslovak Republic. It is intended to provide information to the members of the United Nations Industrial Development Organization.

...-line can be used in the production of rubber cement coverings, lining plates ...-pressure tubes. It is ... the construction of pressure tubes has not yet been ... it will probably be ... to the ...-line ... the pressure is to be sub- ... of ...

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1. Introduction

I was requested by the Secretariate of UNCTAD to prepare a paper on the problems of substitution of asbestos in the manufacture of asbestos-cement. I have tried to prepare this at our present level of knowledge and as far as possible covering the whole wide field of this problem. However, the too wide field prevented me from going into a more profound and desirable depth. Also some of our regulations, such as a limited capacity prevented me from entering concrete relations and numerical data. Thus in the economic field it was an unusually short period which prevented to carry out thorough analysis on the basis of world prices. For these shortcomings I apologise.

The topics in the paper are concerned with the main results of research projects, which we are just finishing, and are for the substitution of 15 - 20% asbestos mixture in the asbestos-cement by the special fibrous materials. To some, this percentage will seem to be low. Still however, it is technically as well as economically attractive.

About the substitution of mixture of asbestos, I am mentioning only outwordly. It is the problem of future work and in my paper I wanted to confine myself only to the real position, which holds good for the present time.

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 microfiche.

2. Previous attempts about the substitution of asbestos in the asbestos-cement system

2.1 Asbestos is and will long remain the basic raw material for the manufacture of asbestos-cement goods. Significant occurrence of this mineral is very little. From this fact and from the necessity of its laborious separation from asbestos rocks, results the high price in world markets. It is not therefore surprising that attempts have been made relatively long ago either to manufacture artificially asbestos, or at least to partially substitute it by artificial fibres and that too also in the countries where asbestos is in abundance.

2.2 As our country obtains almost all demand of asbestos by its import, it is understandable that periodically we also attempted its partial substitution. However, gradually the whole scale testing of glass and mineral fibres was carried out, but till now really definite success has not been achieved. Once, really serious use of substitution was in the period of second world war, that is use of cellulose pulp (3) in the period of very great shortage of asbestos. For example, GFA-dices were substituted upto 50% of asbestos (1). Cellulose however clearly deteriorated the quality of the products. Their strength gradually reduced, products lost the elasticity, from the effect of high absorbibility of cellulose the frost resistance was reduced and roofing showed strong tendency towards warping and cracking (1,2). Immediately after the war, the addition of cellulose was dropped. Similarly unsuccessfully ended also the experiments with the addition of common glass fibres. All attempted types of fibres were so strongly corroded by alkaline cementing baths, that strength reduction of products for even relatively short period was quite significant (1,2). This revelation was first noticed with fibres, which complied with the model, claimed by Halka for the substitution of fibres (2,4,5), as also with others, which contained low alkali (6,7). Little chemical resistance of commercial glass fibres eliminates the possibility of their use in asbestos-cement manufacture (1,2). Still less

is the prospect of successful substitution by mineral wool, manufactured on the basis of blast furnace slag. Here also low chemical resistance and low strength of fibres eliminates their extensive use (1,4,7).

Mineral fibres manufactured on the basis of some rocks or soils are considerably resistant mainly during short period effect of corrosive medium. For example Bulgarian fibre "Aer-gal" showed on the whole, favorable results during short period tests at the Regional research station for asbestos-cement at Pitra (8). The similar results were also later obtained by Bulgarian workers (9). Due to the lack of long period tests and the varying properties of this mineral wool, its use in mass production was not allowed in our country. In the following chapter we will show, that this restriction was well-founded. Unsuccessfully ended also the attempts of substituting asbestos by fibres with higher content of MgO , which was recommended in our country by Dr. F. Mothmann, owner of the firm MIFA-Chemie at Mannheim (10).

The basalt wool is recommended as the most common substituting fibre for the production of asbestos-cement (1,2,7,10, 11). In our factories, the use of basalt wool as partial substitution of asbestos (10-15% substitution mixture) was started in the middle of fifties. After a very short period however, its use was discontinued. The reason being the shortage of good quality basalt wool and also because, it is dusty and venting of too strong fibres (10-20 μm).

The artificial fibres are used at present in ČSR only in little quantity for improving the filterability of asbestos-cement materials. This follows the increased output in the production links and no substitution of asbestos. About the tests with the new substituting fibres, the mention is being made later on.

2.3 Now at least in short about the problem of how much percentage of asbestos is possible to substitute by mineral fibres, so that properties of the mentioned products of asbestos-cement are not considerably deteriorate. In the lite-

return it is shown, for example, that by basalt wool it is possible to substitute successfully 25 - 30% asbestos (1,11). Other sources are more realistic and show as maximum substitution of 10% (2,12), or only 10-15% (13). In this context we will refer to our experience in the following chapters.

2.4 Partial substitution of asbestos in asbestos-cement goods seems however respectively for waste to care workers. Continuous endeavors are being put in the building industry to make cheaper construction. The development of thin walled and prestressed construction at that time was in the limits of technical possibilities. Qualitative changes are only possible to realize in technical, out of which steel supports were substituted by "microstructure" of glass fibre material. This problem was investigated by persons in USSR (14,15,16). During the experimental production, the supporting was achieved by the pulp free aluminated cement covered by threading or by adding alkali-free premixed "fibrous concrete" with strength in tension of 600-800 kg/cm², with the strength in bending about 250 kg/cm² and with modulus of elasticity of $2 \cdot 10^5$ - $3 \cdot 10^5$ kg/cm². The concrete of such properties for building construction was very attractive. For example large dimensioned cylindrical shell is possible to produce with about 5 times lesser weight than today. Later it was shown, that glass as well as mineral fibre in cement medium, were relatively quickly corroded and products lost strength. The use of lesser aggressive aluminated cement and excessively strong basalt fibres has not helped in any way. Therefore the next technical work was performed in the year 1964 and that till today, when the kind was made of fibres, which are competent to resist the long period aggression of the products of hydration cement (17). No work to protect the surface of fibres from organic films (8,18,19) was reached at realization level. It is understandable, that similar to "microsupport", the organic artificial fibre was also tested and was attractive by special treatment (19) and also retained here during laboratory experiments.

In recent years, the development of glass fibres to resist the effect of cement media was carried out by A. F. Legler (10, 21, 22) with others. The glass ropes for the production of fibres, containing SiO_2 , Al_2O_3 , CaO and MgO in such ratios, that melting corresponds to the region of primary crystallization of amorphous. Although these glasses are not perfectly consistent glass, which was not fatal, due to the composition mainly for good collection of fibres with cement masses.

I apologize for going out of the main problem of the paper. I considered this however to be very interesting.

3. Problems of substituting fibres for asbestos

3.1 History of attempts to substitute asbestos in asbestos-cement products, as stated in the previous chapter, seemed to be rather pessimistic. It is however necessary to emphasize, that these attempts were made with fibres specified for other uses. If we want to manufacture fibres especially for substituting asbestos, we must first of all examine the problems, which such fibres will have in asbestos-cement production. It is unfortunate, that during this we examine the function of asbestos in fresh as well as hard-set asbestos-cement products.

3.2. It is known, that commonly used chrysotile asbestos which consists of strip drawn, regularly arranged, enormously thin and probably hollow elementary particles. Understandably till now it is not possible to pulp these asbestos particles. However the technically obtainable softness is very well known. Fibres are thin, strong, elastic, sufficiently resistant against hydration products of cement, thus forming excellent microstructure cement-paste. Asbestos fibres are in addition "cemented". During stirring, the fibres bind themselves by their absorption forces to the surfaces of the cement particles in the cement bath. Ions Ca^{++} , freed by hydration are attracted to the surface of the fibres and form there crystalline centre for growth and later addition of hydration products. The binding of cement paste with fibres is very strong (25). Lately we have to review our opinion about very high resistant asbestos in cement matrix. From the work of J. Butry (24) it follows, that ions K^{+} disturbs crystalline lattice of chrysotile and frees from it Mg^{2+} . By higher concentration of K_2SO_4 in recirculating water which disturbs the surface of asbestos fibres, reduces their reinforcement capability and can bring about reduced strength of asbestos-cement up to about 30%.

Asbestos by now can be substituted in asbestos-cement production only by fibres, which could be cemented, sufficiently and long period resist towards hydration products, would have good length and diameter would be strongest and most elastic possible, did not contain the non-fibrous ingredients, could give the reinforcement with hard-set cement

ness, in case of both filament fibres and staple fibres stirred the individual fibres and could not be hygienically defective. Some caution is taken to avoid possibilities of formation of new mould spots.

3.3. Two possibilities are suggested for the production of fibres with artificial fibres. First, the artificial fibres will have to be suitable for the same spinning conditions as the natural fibres. Second, the artificial fibres will have to be spun in a different way. The substitution of the artificial fibres will depend on the volume and on the production to be achieved. In the case of present-day equipments for production of staple fibres, for example, we consider the upper limit of 50% substitution.

3.4. Long period resistance against oxidation products of cement:

We discuss in detail about this question in the following chapter. Here we will mention only that oxidation of resistance of fibres is usually considered as a problem. It is however necessary to realize, that gradually by oxidation, fibres lose their reinforcement ability. In the case of the products mentioned. It is important to know that of all the long period processes for the fact, that also with seasoned concrete (40-50 years) after descaling, the pH value above 12 was obtained (20).

3.5 Length and diameter of fibres:

Too long fibre creates trouble in production links and too short reinforced little. We consider 30-35 mm as optimum value. From hygienic point of view, too strong fibre creates trouble. If fibre is characteristically resistant, it will be possible to reduce its force to 4-5 kg. Such fibre will be elastic and will not be hygienically defective.

2.6 Strength and cleanness of fibres:

Well known fact is that strength and elasticity of fibres are
or their cleanness from dust fibres etc. During selection
of technology for production of fibres, it is
then necessary to select method, which is to resist as best
as possible the possibility of occurrence of surface defects
of fibres. The defects can then reduce the strength of fibres
and also their cleanness. (see also, for example, [10, 17, 21,
19]).

2.7 Non-fibrous inclusions:

Fibres must not contain particles of diameter more than 5 μ m.
If of the smaller particle content, should be least possible.

2.8 Flotation of mineral wool:

Our tests showed, that with the use of correct procedure for
cleaning mineral wool and gentle hydrophobic cleaning of fibres,
it is possible to obtain with thin fibres also good distri-
bution in electrospinning culture.

2.9 Adhesivity of fibres with cement stones and their hy-
gienic cleanliness:

Reinforcement ability of artificial fibres is possible to
improve by wetly spraying their surfaces, by suitable alkali-
line solution of silicate material. Smaller diameter of fibres
and spraying them simultaneously together solved hygienic problem
during manufacture as also in processing of mineral wool.

4. The loss of usefulness of porous fibres for substituting cations

4.2.4. It is generally, not the criterion for judging the suitability of fibres for substitution of cations is their long period resistance to corrosion in the media of the solvent. Therefore at least briefly, it is necessary to consider the corrosive media, which are used in the practice and about the results of corrosion of the porous type of organic fibres.

4.2.5. With organic fibres, which are subjected to the effect of stronger alkaline medium, brings the attraction and adsorption of OH^- ions on to the silicative group $-\text{Si}-\text{O}-\text{Si}-$. This goes with the rupture of other strong silic bridges and consequently binding the whole group of SiO_2 with surface of fibres. While dealing with corrosion in strong alkaline medium (e.g. NaOH) and further significant surface layer aggressive solution, the bonded hydroxyl silicic groups join with free alkaline ions to form orthosilicates, which goes into the solution. Like this stripping combines of the successive layers of the surface of fibres and corrosion continues.

In the solution of $\text{Ca}(\text{OH})_2$ there can originate also little mobile complexes of hydrosilicates of calcium, which settle down on the surface of fibres. During this period, from these originate new crystalline forms. However, as layers crystallize it does not prevent the successive penetration of aggressive ions to the undisturbed part of fibres. Besides the rising characteristics for corrosion of fibres in alkaline medium for specific conditions and solubility of fibres, other characteristics for corrosion of fibres in acidic medium can also be considered. During which diffusion of substituted cations K^+ and Li^+ , in the surficial layers of fibres

ions H^+ or H_3O^+ takes of ca. heat H. Manual (30) considered these over the range in which there is higher concentration of alkali than in any given solution. For which character of corrosion, the actual overvoltage is determined for both cases of corrosion.

In addition to the basic reactions, the course of corrosion is also affected by other factors, i.e. nature of electrolyte, nature of electrode etc. or fibres.

4.3 Because of the difficulty in determination of fibres directly in cement paste is not easy to get good qualitative results, therefore we tested several indirect methods, such as resistance of fibres in $Ca(OH)_2$ solution, in cement both in $Ca(OH)_2$ solution, and in cement paste of short-term content and nature. After comparing with direct tests in cement paste, we reached the conclusion, that for determining the alkaline resistance of the substituted fibres, it is sufficient to combine the test of long period resistance in $Ca(OH)_2$ solution with short term test in both.

4.4 For test of fibre resistance against $Ca(OH)_2$, we processed a relatively simple method: the polyethylene bottle 70% of fibre was cut, deprived of additional oily arrays and filled with 500 ml of $Ca(OH)_2$ solution of concentration 1060 to 1100 mg CaO/l , practically full bottle is closed airtightly and put on the temperature of $20 \pm 2^\circ C$. Once a week the contents of bottle are stirred for a short time by the slow stirring machine. In agreement with the theoretical consideration about corrosion fibre, it is possible to expect, that reciprocal relation between $Ca(OH)_2$ and fibre will form the difficult soluble hydrated products, containing calcium. Decrease in Ca^{++} ions removed from solution will then determine the degree of corrosion of fibre. The direct determination of Ca^{++} is not hampered by the fact, that fibres with higher content of alkali forms soluble al-

kaline silicates. This is difficult to do, but it is then possible to supplement with microscopic observations of cross-sections of fibres.

By this procedure we tested the tensile resistance of individual fibres of several fibres. Most important results are tabulated in Table 2, given in the end of this report.

Most important results are graphically illustrated in Fig. 1. We see here, that least important result is that of a group of undissolved fibres. Also in first phase with NaOH we see high degree of corrosion. With untreated and treated fibres, the structural corrosion starts disappearing after 2-3 hours, but then quickly continues. The corroded layers of fibres are clearly distinct on microphotographs of basalities (Fig. 2) and splintered (Fig. 3) fibres. Gradually gels grow and after 24 hours distinct ones of the centre grow (Fig. 5). In the later period crystallization slowly continues. In Fig. 4 is also included the effect of average diameter of fibres.

4.5 For determination of resistance of fibres against NaOH we used the method of J. J. Langer and J. F. Rydman (21), modified such, that from it is possible to measure with the fibres, whose diameter is not constant along the whole length. Individual fibres are fastened in the measuring frame by three narrow slots and two around these slots. The force on the fibre is then possible to measure by three force always on the same points. We then heated the frames together with 10-15 fibres at 100°C for 1,5 hours in NaOH . Difference in the diameter before and after leaching, expressed in percentage decrease in original diameter of the fibres, is the measure of its resistance towards NaOH . It is possible to measure fibre force from 6 - 20 gm. Because cleavage resistance of fibres depends strongly on their diameter as well, it is always necessary to state, for which average diameter the

Reasoned now, the whole road, will be that you is the calcu-
lation of regression curves for the data in in three
to the number of data.

In the first case, the regression curve for the fiber of
one of the data points is calculated. The fiber of
the data points is calculated. For example, the figures
of the data points are 10, 20, 30, 40, 50, 60, 70, 80, 90, 100.
The data points are 10, 20, 30, 40, 50, 60, 70, 80, 90, 100.
The data points are 10, 20, 30, 40, 50, 60, 70, 80, 90, 100.

Figure 2. The regression curve for the data points 10, 20, 30, 40, 50, 60, 70, 80, 90, 100.
The data points are 10, 20, 30, 40, 50, 60, 70, 80, 90, 100.

4.5. Consider the case where the data points are 10, 20, 30, 40, 50, 60, 70, 80, 90, 100.
one during the initial effect of all the data points little
resistant, that the data points are 10, 20, 30, 40, 50, 60, 70, 80, 90, 100.
with
is clear, but the data points are 10, 20, 30, 40, 50, 60, 70, 80, 90, 100.
that the data points are 10, 20, 30, 40, 50, 60, 70, 80, 90, 100.
continue to improve. However, it is not possible to re-
compute the data points are 10, 20, 30, 40, 50, 60, 70, 80, 90, 100.
reading. Figures specified for these curves will have to be
compiled with deliberation.

5. Production of A-fibre

5.1 Development of A-fibre:

Ostrava branch of V. skopny Institute for Construction Materials in Brno is engaged for several years in the research and connected with mineral wool production. In beginning of the year 1957, this branch was transferred to the Ministry of Construction to have right to the possibilities of production of mineral wool able to substitute, particularly the asbestos in asbestos-cement products.

Theoretical study of composition of amorphous fibre, line radius lead to the formation of working hypothesis, according to which heterogeneous production of new type of fibre, e.g. A-fibre was carried out. Positive results of the resistance test of A-fibre in alkaline medium then moved the authors of the hypothesis to apply the patents first in C.S.S.R., later in other countries.

Economic aspect of partial substitution of asbestos was showed favourably, but the enterprise, which is chief producer of mineral fibres in C.S.S.R. supported our institute in the costs of research work, carried in the development of special fibre, able to substitute 10-10% asbestos in asbestos-cement products. These works were started in the year 1960 and are scheduled for completion in middle of the next year, by commencing of mass production of A-fibre.

5.2 Principle of A-fibre.

Theoretical considerations, which are supplementary to the proposed composition of A-fibre extend beyond the time limit of this treatise. In a very simple way we can say, that we came to the well known fact, that majority of oxides with high content of rare elements with strong electropole (Zr, Zn, Ti) show increased resistance towards alkaline lye. First theoretically and then experimentally as well, we derived

the limits of chemical composition in such a way that ions increase the resistance towards Ca(OH)_2 as well, and in addition to it work so intensively that after reaching the required resistance, their very little addition is enough. I apologize, that Czechoslovakian patents registration prevents me before completion of patent procedure, to derive these limits numerically. These are not to life and success depends upon prescribed combination of all chemical components of melting.

We still increase the resistance of the fibres by creating in the surface of fibres a film of suitable organic-silicate material of strength film. The material is selected such, that integrity and force of surface film is possible to separate from the initial protection, used with substituting fibre for asbestos into the perfect protection specified for other uses. Simultaneously, it is possible to make the surface of fibre rough by miniature filament of this material and thus increasing the cohesiveness with cementing paste in first few weeks of their hardening.

5.3 Scheme of production of A-fibre

A-fibre not only fulfilled the requirements prescribed for the successful substituting fibre, but is in addition economically advantageous, is produced from relatively cheaper raw material and that too with suitable process for mass production. One of the possible production scheme is being shown in Fig.7. It is simple and represents the recent state of technic in this field.

Raw material and fuel are fed to the furnace for melting (3) from the storage bunker (1) by skip (2). The granular materials from smaller bunker (4) are gently added direct to melting zone of the furnace via furnace conveyor by special nozzles (Czech.Patent). During charging it is necessary to maintain prescribed ratios of the feed (Patent registration ČSFR, USA etc.). The melt at $1400-1500^\circ\text{C}$ is pulped by four-cylindrical pulping machine (5). Waste from the machine is taken out by

conveyor outside the hall. The fibres formed are conveyed by air through blowing slot from pulping cylinders to the settling tanks (7).

The required compressed air is produced by ventilator. In the middle parts of settling tanks are installed spiral nozzles, by which while the fibres still rise to pulping cylinders, sucking air dilutes and filter of oxide-collectors and fill with tapping catalyst. This fluid is conveyed to nozzles from storage tank by high pressure pump (8). Such modified fibres are conveyed by air vent (9), in on belt conveyor, by which the fibres, formed from fibres elevated from settling tanks. From the elevating belt pass to cleaning table (10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20), where the unremoved impurities are rejected. These are conveyed back to the furnace by means of conveyor to its melting zone (21, 22, 23, 24). Cleaned A-fibre passes between rollers (25) to the conveyor, where it is taken, in direct way in pack to. Undoubtedly, there are also other, and different of production schemes. However, they must guarantee the required amount of A-fibre. At other places I mention about the following important production parts and finish the chapter after considering about the capacity and location for A-fibre production.

5.4 Raw material base:

For the production of A-fibre while using cubic furnace, suitable basic raw materials are those, basic basalt and basaltic rocks with periclase and felds, such as dioritic basalt, tefrity, basalt, and felds, etc. These should have a low percentage of iron and their basic content and their viscosity should be such that it is possible to melt in such type of furnace and pull the melt by controlling mechanisms. For preliminary consideration of suitability of raw materials according to these criteria, we prepared the test

tables. From these we can know how tested raw material will be for use in the production of A-fibre, by complete chemical analysis of raw material with at least 50% reliable estimate. In some cases it is possible to improve the melting index of basic raw material by addition of clays, limestones or dolomites.

Suitability and adaptability of raw materials is possible to estimate in some of the next tested tables. Final decisions are then taken after a few laboratory tests.

At the level of melt in the furnace according to the type of raw material material, generally at least 60% acidic metals, whose ions have strong electric field.

In specific cases, it is possible to use some clays from metallurgy of iron-ferrous metals as well. Also used for batch melting, the suitable new shaft furnace, developed in the field of use of basaltic rocks and besides to use also for some coals, clays or shales. With the use of furnace tank, this development is still wider and quality A-fibre can be used also for other uses than partial substitution of asbestos. While concluding this section I expect, that requirements for chemical composition of A-fibre are precise. Here use can only be of raw material with sufficient chemical composition and during charging in the furnace, their mutual ratios should be maintained.

2.3 Furnace and piping

The melting aggregate shown in technological scheme is cupola. Its economical advantages and technological deficiencies are generally known. For our type of cupola for production of A-fibre is usually notified only by this, that closely below the melting zone provided with three special nozzles, equally placed around circumference of furnace. Nozzles blow the fine granulated active materials to the melting zone. These are conveyed to the blower from small bunker located near the furnace. Charging is by small vibrating feeder having

the 3 - 6 % of total capacity of furnace. Our experience with furnace of 1100 kg melting hour capacity showed, that by this procedure, fibres receive the practically full quantity of active constituent. It is advantageous, if the furnace is provided with the appliance for the periodical removal of reduced iron. About the possibility of use of other types of furnace as well, as mentioned in the foregoing section. For smaller production, there may be more advantages. For pulping the melt, it is essential to use special equipment working on centrifugal principle, and able to produce uniform fibres of even diameter 4,5 - 5 μ m. In during laboratory experiments, as also during semi-operational production, we verified the pulping machine with three to four vertical cylinders. Pulping by sticks is not possible to allow. Fibres produced by these methods have so to say greater quantity of surficial flaws, reduce their shear resistance as also strength. We see this also in Fig. 1, where A-fibre pulped by blowers with 500°C hot medium is Fig. represented by number 5 shows the smaller resistance towards CaOH₂ than the analogous fibres (number 4) produced by pulping machine.

5.6 Settling chamber and surface finishing of fibres.

Settling chamber is of usual construction. With respect to the stickiness of cleaning machine, it is recommended to obtain sufficiently wide regulability of the speed of belt, bringing out the settled mineral wool.

In the central portion of the settling chambers, the produced A-fibre is sprayed by strongly diluted emulsion of organic-silicate material.

For spraying purpose, 4 spiral nozzles are located such, that whole stream of fibres, blown out from rotating cylinders, was covered. Capacity of blowers, size of the spraying tubes of emulsion, dilution of emulsion and operational pressure of the pump should be harmonized so as to get the partial protection as well as roughing of surface of fibres.

The selected organic-silicate material in cement stone is sufficiently long time resistant and its slight remainders which were insufficiently created, function in asbestos-cement production as slipper of carrying waters. Connection slips so slickly, that fibres after leaving cleaning machine does not stick to itself. We solved the problems of combination with spraying the fibre and semi-operationally tested in one of our production of mineral wool.

5.7 Cleaning of A-fibre

As was stated, the A-fibre should not contain higher non-pulped portions and can have only specified maximum length. In the production chain, therefore it is necessary to equip cleaning aggregate. I would not like to prescribe directly the type of cleaning aggregate. However it should be completely clean, as also shortening the function and not damaging too much the surface of fibres. We developed the prototype of the machine, which suits these requirements. Not too big and is possible to move it in production chain within 40 sec. and goes on line this without disturbing the operation links in production of common mineral wool and reverse. The waste from cleaning machine is possible to return to melting zone of the furnace.

It is also necessary to mention, that during A-fibre production, it is not possible to use cleaning system working with pneumatic transportation of fibres. Little non-pulped portions, present in mineral wool are sprayed so much on the surface of the fibre that its chemical resistance and strength is reduced considerably. This is documented by the photograph from electron microscope, which we have shown in Fig. 9.

5.8 Packing and dispatch

A-fibre is possible to bag by any sufficiently precise equipment.

Dispatch in packets is not too advantageous for any reason. If the customer is unable to pay, the size of bag is impossible to select, so that it will be difficult to predict the production of the above-mentioned.

6. Properties of A-fibre

So far we have mentioned only about the properties of A-fibre in general terms. I will state very briefly, those in the table below for the sake of comparison necessarily.

6.1. General properties

6.1.1. Resistance to acids and alkalis. It is known that the A-fibre is highly resistant to acids and alkalis in asbestos- and asbestos-cement products. It is also resistant to acids.

6.1.2. Resistance to alkalis. Towards alkalis products of cement:

Resistance of A-fibre to produce A-fibre towards $Ca(OH)_2$ and $NaOH$ is shown in the table below in G. Fig. 1 and Table 1, 2. In Fig. 2 then we give the microphotograph of A-fibre produced by process (in general it is designated by the symbol ϕ) in the state in which it is saturated $Ca(OH)_2$ solution. We see in this, that fibres are practically not corroded and only some are isolated cases of gel-like drops on the surface.

Defect resistance is also shown by the fibre produced conventionally by industrial equipments. In Fig. 10 are given the average values and 95% reliability some of 33-40% increase in concentration of $Ca(OH)_2$ during the testing of A-fibre from our first semi-industrial experiment. Corresponding numerical data are shown in Table 1. These are fibre without surface finish. First results of test of surface finish A-fibre from second big experiment are also good.

Results of test for resistance of A-fibre direct in cement are also good. During increase in examination of thin sections from testing, (100% asbestos + 50% A-fibre + cement), put on the terrace of our workshop for 1 1/2 years, no corrosion was found on A-fibre. No change was also found on A-fibre in envelopes dates, heated for 100 hours at 28-30°C. In the reverse after 10 years heating in autoclave in series

at 170°C and 5 atm, the 4-fibre was distinctly corroded at the point of contact with asbestos. Keeping in mind the effect of pressure and, on the other hand, the fact that during autoclave tests the 4-fibre was not corroded.

5.4. Kinetics of the corrosion process

During our practical experiences with routine substitution of asbestos, 4-fibre of average diameter of 2.5 microns was used. Substitution of length of fibre was possible to determine experimentally. It is possible to determine the rate of corrosion with the help of the relation to the practical use, showing the effect of small 4-fibre. In determining the length of 4-fibre, therefore we use only our experience.

5.5. Strength and elasticity of fibres

The relation between diameter and strength of fibre is a relation for the mineral fibres. This approximately holds good for 4-fibre as well. On the other hand, it is possible to use a fibre of 2 microns diameter, strength of at least 100 kpa, whose elasticity of such thin fibres is sufficient.

6.0. Corrosiveness of 4-fibre with cementing mass and hygienic staff wear.

As was stated, the surface of the fibre is covered by organic-silicate material. This covering is of course only of sub-microscopic size. We can only examine this by electron microscopy. For example, in the case of 4-fibre using stereoscopic transmission electron microscope with substitution of asbestos, the same can be seen.

In connection with our experience, we must take into attention the fact that the hygienic conditions of working environment. Just during the production of 4-fibre is almost strongly exposed to the dust of spraying in settling chamber. Only when working machine, we are planning to have a sensitive place. This will be especially in the central

de-dusting system of production.

In the manufacture of rayon-cement goods, we did not had the trouble with dust of β -fibre during our tests. Sprayed filter also showed in this direction enough effectiveness. In addition to this still to be desired: keep in the β -fibre specific remaining moisture. This is so little that it does not spoil the type in which β -fibre is packed, even during long storage. Utilizing sufficiently reduced the chain diameter and elasticity of the β -fibre.

7. Use of A-fibre - partial substitution of asbestos

7.1. As it was known, that whenever the substituting fiber was introduced in any country at the request of one of the undertakings producing asbestos, the latter were requested to develop such a fiber, which would be able to substitute 15-20% asbestos in asbestos-cement roofing and lining sheets, quality of products with A-fibre should be prepared as all the standards applied for these products. Simultaneous use of A-fibre in asbestos-cement products should not interfere with the management of existing technology.

7.2. In connection with this work, several laboratory experiments, field tests were carried out the mentioned tests as well as industrial experiment, working on 141-142 system. Their purpose was to verify the possibilities of A-fibre not only from the point of view of properties of finished products, but also from the view of nature of auxiliary arrangements, productivity, etc. and this always during several plant operations.

First test which was carried out in August 1966 was short period (7 hours) of initial trial composed of a of preliminary verification of effect of substituting asbestos of production process. In this, clear A-fibre without surface finish was used. During experimental operation usual production sequence was maintained. Ratio of asbestos and A-fibre was changed such, that in asbestos mixture, 10, 15, 20% of asbestos was gradually substituted by the same quantity of A-fiber. In all such production was over 1000 m² of corrugated sheets. Tests on finished products showed that substituting frost resistance and strength in tension while remaining agreed with the requirements of "CS. 70 1401 "asbestos-cement corrugated sheets". Against usual production, the strength of corrugated sheets (compared with 100% substitution by A-fiber) was reduced at some points. The mixture of 85% asbestos and 15% A-fibre showed after 10 days 240 kg/cm² and after 100 days 255 kg/cm².

center of strength and absorbability of products were a bit lower than usual practice.

During second half of production in June 1957 some technological procedures were produced and used for surface finishing of fibre. This consisted of production of extremely upto 40 hours so that during production of 100% fibre in production of air covered by 100% of A-fibre, and a elastic mixture was applied to the surface of the fibre. Out of the whole quantity of fibre produced, in the first phase of the experiment, Dk substitution was 100% quantity of A-fibre by weight, in the second phase Dk was substituted for 100% substitution aspecton 1:125 then greater quantity of A-fibre during which a substitution was made from 100% of A-fibre to 100% of biological strength. On the contrary, some of the influences were favourable: gentle drying of softening water showed the probability of the dispersion station in case 1 production. The filterability of a suspension of aspecton was improved due to 1:125 fibre. This was shown by gradual decrease of sediment content in softening water and by this, that gently reduced number of revolutions of circular sieve, necessary to reel one sheet (from about 12 to 10). Fibre material continuously maintained sufficient plasticity and not even during 40% substitution, on the edges of the wool there was not cracks. The dispersion of A-fibre in asbestos-cement suspension was so nice, that it was not possible to distinguish the center of artificial fibre in the products. Increased filterability of the suspension, partly increased also the output of the production arrangement and in total 28 1/2 shift against the planned 24 1/2 shift. In all, in this experiment 12,000 m² of corrugated sheets were produced, from which 400 pieces were selected for laboratory tests. After 28 d./3 storage in a dry cellar, following strength in tension while bending were recorded (practically corresponding with usual production):

with Dk substitution: $T = 220 \text{ kg/cm}^2$; $s = 12,5 \text{ kg/cm}^2$
with 10% substitution: $T = 150 \text{ kg/cm}^2$; $s = 15,5 \text{ kg/cm}^2$.

Absorbability as well as from the nature of the products agreed with 311. It is also being referred about the results of this agreement, which would be a really expected to be more effective in giving ideas for the production, that is, the production of high-strength fibers of polyester in the next few years.

In addition to the economic advantages, it also has the technological advantages.

The advantages of production of high-strength fibers, developed with the use of the new technology, are the production of high-strength fibers. During the last few years, in the year 1968, in the production of high-strength fibers, the production of high-strength fibers, which are of high strength, are being produced by pure polyester mixtures.

7.2. Our work in this field is not finished yet. By the end of this year we will be able to add the addition of fiber in the next 4 years the production of high-strength fibers and that too also for some other purposes. We are planning to do this as a separate project. Next year we will carry out the basic work with material, containing 50% polyester and 50% glass fiber.

6. Economic effectiveness of production and use of A-fibre

6.1. Joint efforts in the field of the extent of consumption of A-fibre.

In order to reduce the consumption of A-fibre, it is necessary to create conditions for the substitution of A-fibre with other types of fibre. It is necessary to work on the substitution of A-fibre with other types of fibre. If the consumption of A-fibre is 0,5 kg substitutes of A and B categories, then the consumption of A-fibre is 0,5 kg substitutes of A and B categories. If the substitution, that means, the consumption of 4 000 tons of A-fibre. And when the substitution is possible, the consumption for production of A-fibre will be increasing, and it will be increasing quantity, and it will be increasing quantity. That is, the A-fibre will be produced at one place in our country, and from there it will be delivered to consumers. In addition, the production of A-fibre in our country is not near to the world production, the time of 1 000 - 5 000 tons yearly. It is, in our country has not exceeded 20% of the price of A-fibre as of 1970.

6.2. To provide the calculation of savings in production costs

While calculating savings in production costs, we will take the assumption, that the production of A-fibre is not roofing, Canadian A-fibre was used of the consumption 20% category 5 - 20% category 4 - 20% category 3.

Net price of the A-fibre is about 200 Cdn \$. Also, the calculation of savings in production costs of the A-fibre is based on the countries importing aspects, we can assume that price of 1 ton of A-fibre does not exceed the price of 1 ton of A-fibre substitutes. From which it is about the 200 Cdn \$ for one ton of A-fibre. It is possible to save 200 000 Cdn \$. For example, with difficult or positive foreign trade balance, it is then possible to obtain fact in foreign exchange savings. In such cases, that need not be 4 000

tons asbestos in Canada, follows and can save yearly 1 300 000 \$.

In the foregoing paragraphs I have stated I understandably took real preliminary assumptions verified by research.

As we have seen, that our ideas concerning industry is placed in world scale only in very low percentage category. It is possible to assume, that substitution of substituting fibre would be seen much higher in many countries. How approximately it looks, the rising consumption of A-fibre and with it total savings in material costs is shown in Table 4 in enclosures.

8.3. Orientational data of A-fibre production

As was stated, total production of A-fibre is on the whole obtained by the usual arrangements known from the production of mineral wool. Required supplementary equipments (mainly cleaning machine) are given in chapter 2. Their price form 5 - 15% of the price of usual production chain for mineral wool. It was also indicated, that for production of A-fibre, it is possible to modify relatively simply the large production of mineral wool as by its owner can have sufficiently modern pulping equipment. Because it is possible without disturbing the operation to pass from production of A-fibre to common production of mineral wool, the whole year's capacity would not be used for A-fibre. Nevertheless I give some informative figures.

We have plants for production of mineral wool with yearly capacity :

2 000 tons - 5 000 tons - 15 000 tons

All three types of plants can be used for producing A-fibre.

For these following number of workers are counted:

Manufacture with production of 3 000 tons/yearly ... 38 workers
5 000 tons/yearly ... 49 workers
15 000 tons/yearly ... 82 workers

Dealing with the worker's productivity. Their distribution is shown in Table 2.

Corresponding to the above statistics for production of 4-11-60 is the data in Table 3.

When the worker's output is positively influenced by specific conditions in the plant, it is possible to assume, that with a view to increasing the economic categories (wages, discipline, productivity, turnover etc.) report expresses the low quality of the plant's economic effectiveness.

9. Summary

In USSR was developed special fibre (i.e. A-fibre), able to substitute 15-20% of asbestos in asbestos-cement products. Basic raw materials for its manufacture are possible to find according to our opinion in variety of the countries. A-fibre is produced in the equipment, used for manufacture of mineral wool. Degree of mechanization and also the actual costs of production chain are to a specific measure mainly in the region of control (as with raw materials in finished products) possible to change according to the specific ratios of production countries. Price of supplementary equipment does not exceed usually 15% of price of production chain. In no case, however the production costs of A-fibre will exceed double the costs of usual mineral wool production. That means, with except special cases, price of A-fibre does not exceed 50% of the price of average mixture with imported asbestos. Hence the profits are here, in addition to the savings in costs of basic material in the case by importing asbestos as well as considerable savings in foreign exchange. A-fibre is possible to use for manufacture of asbestos-cement roofing, lining sheets and pressure free cylinders. Use for the manufacture of pressure cylinders is not yet tried. It is likely, that A-fibre will not be possible to use for other products produced through autoclaving process.

Partial substitution of A-fibre in asbestos-cement manufacture does not require any restrictions for technological equipment. Addition of A-fibre does not cause any trouble in technological process. Movement of production on the contrary increases. Quality of products with 15-20% substitution of asbestos by A-fibre corresponds to the standards for asbestos-cement goods. That means, that these products can be used everywhere in case of classical asbestos-cement.

Production of A-fibre is possible to set up, or adapted from the constructed set up for production of mineral wool wherever there is easily able to be of 2 000 tons A-fibre, that is where at least 1 500 000 kg of asbestos-cement is produced

poorly. With smaller scale it is necessary to manage the manufacture even that would produce alternative A-fibre and carbon fibre. For isolating purpose, manufacture should be advantageously situated in the centre of the near of the area. With respect to the relatively low specific gravity of A-fibre (1.4 - 1.5) by ρ , it is economically advantageous to treat root materials rather than finished products. In developing countries that are producing A-fibre could stock it for earlier countries of same countries. Low specific gravity of A-fibre could undoubtedly reduce cost of its transport by sea.

Work, which was ordered to us is near the completion. It is however likely that in the future we will continue to work and try perhaps for higher, upto 50% substitution of asbestos by A-fibre. These products however will probably have some technical problems different from asbestos-cement. For example, while using in climatically exacting regions, it would be necessary to calculate with care, however efficient protection of surfaces of finished products (this problem in CCSR is being approached for study). High percentage of these products could be their advantage. High percentage of substituting asbestos will also contribute to the solution of problems of silicosis diseases, which occurs sometime in asbestos-cement industry. All these however are the subject of future work.

Table no.1 Resistance of fibre towards $Ca(OH)_2$.

Basic raw material	Production thickness in mm	Increase in concentration of $Ca(OH)_2$ in days		
		30	60	120
Basalt 1	21,0	5,1	1,0	10,2
Basalt 2	17,9	2,0	3,2	41,8
Basalt 3	190	3,1	3,2	15,4
Mergel	27,3	1,7	4,5	42,1
Slag 1	1,7	1,7	2,5	34,7
Slag 2	4,4	11,4	19,5	48,2
Slag 3	8,6	26,8	40,7	44,8
Glass	23,2	11,4	37,4	45,3
A-fibre	5,2	3,0	2,3	5,0
A-fibre	5,0	1,0	1,0	1,0

Horizontal cylinder, S=4-cylindrical pulping machine, J=blower, A=other method.

Table no.2 Resistance of fibre towards NaOH.

Type of fibre	original diameter in	reduced diameter after leaching
Wool	14,7	28
Cellulose	12,7	43
G - 1	14,7	42
A - Fibre	12,7	26
PAGE	12,7	22
G - 2	14,7	5

Table no.3 Resistance of industrially manufactured A-fibre towards $\text{Ca}(\text{OH})_2$

Leopase in concentration $\text{Ca}(\text{OH})_2$	30	90	180	360	days
Arithmetic average \bar{x}	1,0	2,7	4,0	6,1	6
Standard deviation s	0,0	0,4	0,0	1,0	7
$u - 2$	0,4	1,9	3,7	4,3	6
$u + 2$	1,6	3,5	5,7	11,5	6

Table no.4 Increase in savings with respect to increase in production.

Yearly production of roofing 1000 sq/ft, 48 cm	Yearly consumption of asbestos in tons	Yearly consumption of Ar-fibre in tons	Savings on Ar-fibre in Rs.	Yearly saving on costs in Rs.	Yearly saving on overhead costs in Rs.
25 000	21 000	4 200		770 000	1 310 000
50 000	42 000	8 400	160	1 344 000	2 593 000
75 000	64 000	12 800	160	2 018 000	4 104 000
160 000	85 000	17 000		2 720 000	9 450 000

Table no 5 Survey of number of workers and their distribution in A-fibre manufacture.

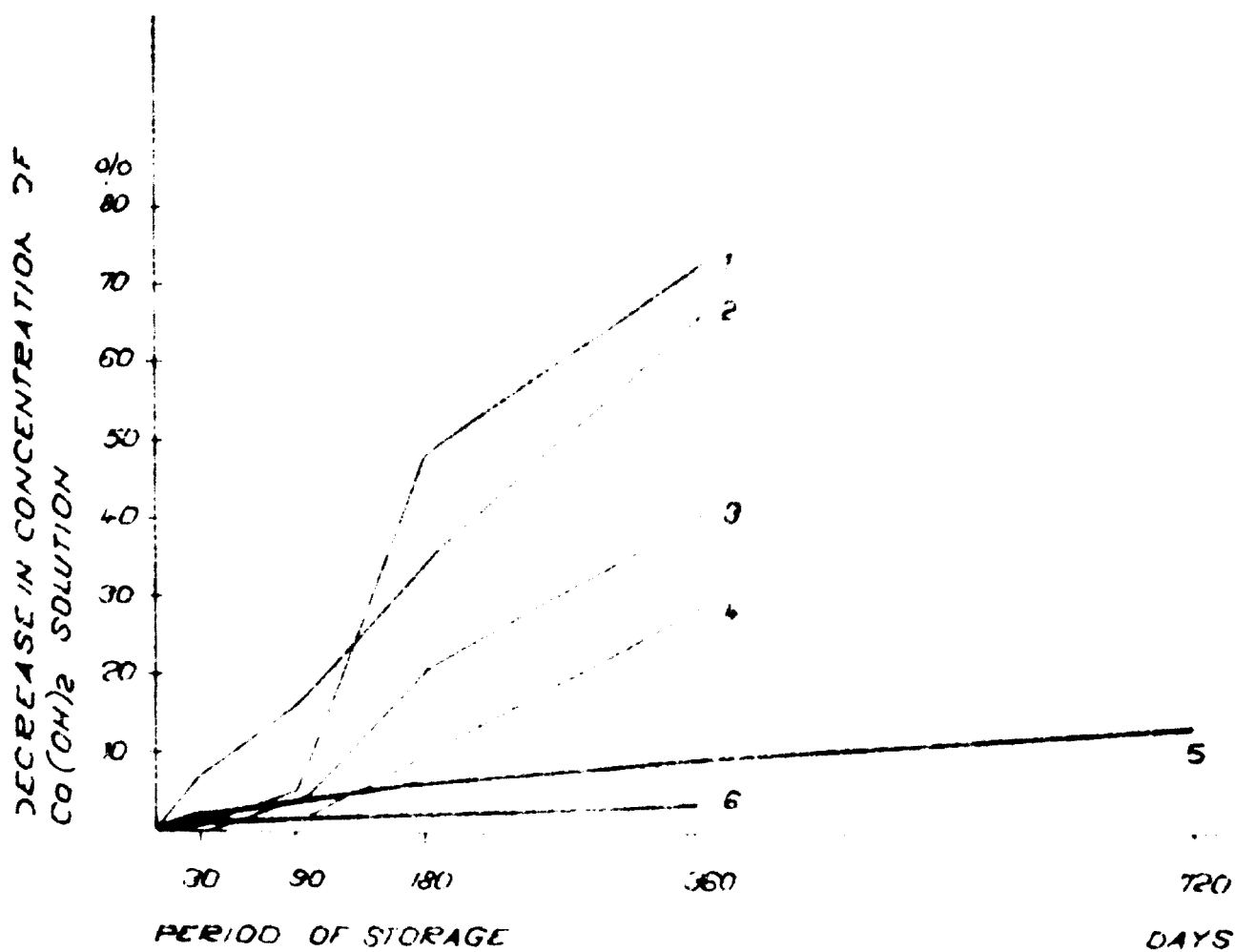
Working place	Yearly production				Number of shifts	Number of shifts	Yearly production 1000 t	Number of shifts	Yearly production 1000 t
	I	II	III	IV					
Storage for raw materials	2	1	1	1	3	3	2	3	2
Feeding appliance	3	1	1	1	1	3	2	2	2
Melting and pulping	3	2	2	2	2	3	3	2	2
Supervision at production line	-	-	-	1	1	1	3	1	1
Packing	3	3	3	4	4	13	3	6	6
Expedition	1	2	-	2	-	2	2	1	3
Storage auxiliary for materials	1	-	-	1	-	1	1	1	1
Installation of line	-	-	-	1	1	3	3	1	1
Maintenance	3	1	1	1	1	4	3	2	2
Maintenance of electric plant	3	1	1	1	1	4	3	1	1
Laboratory	1	1	-	1	-	1	1	1	1
Removal in storage	3	2	2	2	2	6	3	3	3
Works siding	2	1	1	1	-	2	2	2	2

16 12 10 10 40 21 15 13 10 50 30 30 10 10 50

Table no.6 Consumption of raw materials and fuel for A-fibre
manufacture.

	Yearly production
3 000 t	15 000 t
4 700 t	22 000 t
Netolunporosi coke	1 550 t
6070 koalase	4 950 t

FIG. 1 RESISTANCE OF FIBRES IN $CO(OH)_2$ SOLUTION



FIBRES : 1 - FROM CLAY, 2 - FROM BLAST FURNACE
 SLAG AND BASALT, 3 - FROM BASALT (15 μm),
 4 - FROM BASALT (21 μm), 5 AND 6 A - FIBRE



Fig.2: Microphotograph of basaltic fibre after 8 months storage in $\text{Ca}(\text{OH})_2$ solution, 620 x magnification.

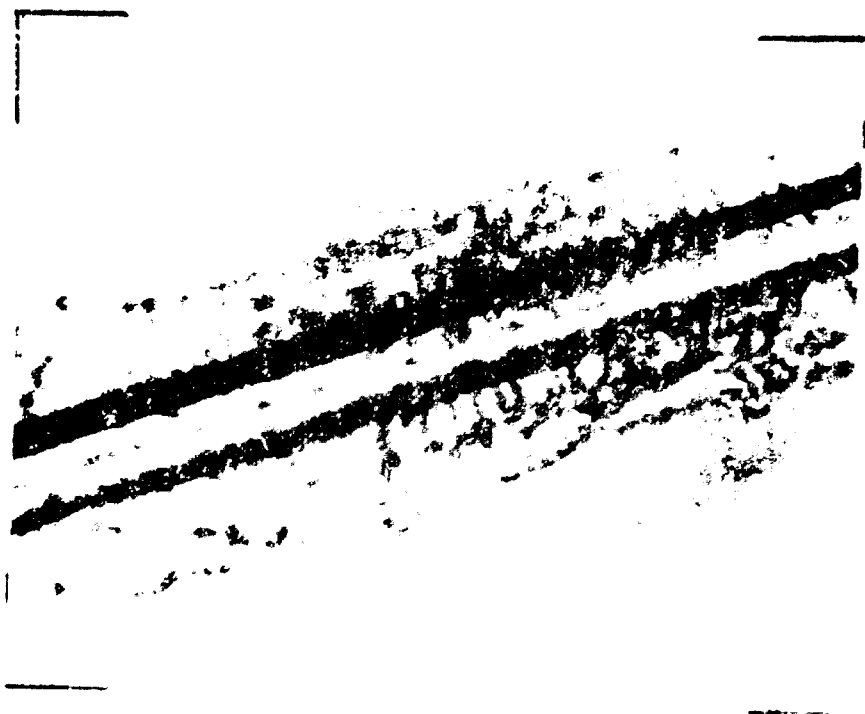


Fig.3: Microphotograph of fibre from clay after 8 months storage in $\text{Ca}(\text{OH})_2$ solution, 620 x magnification.

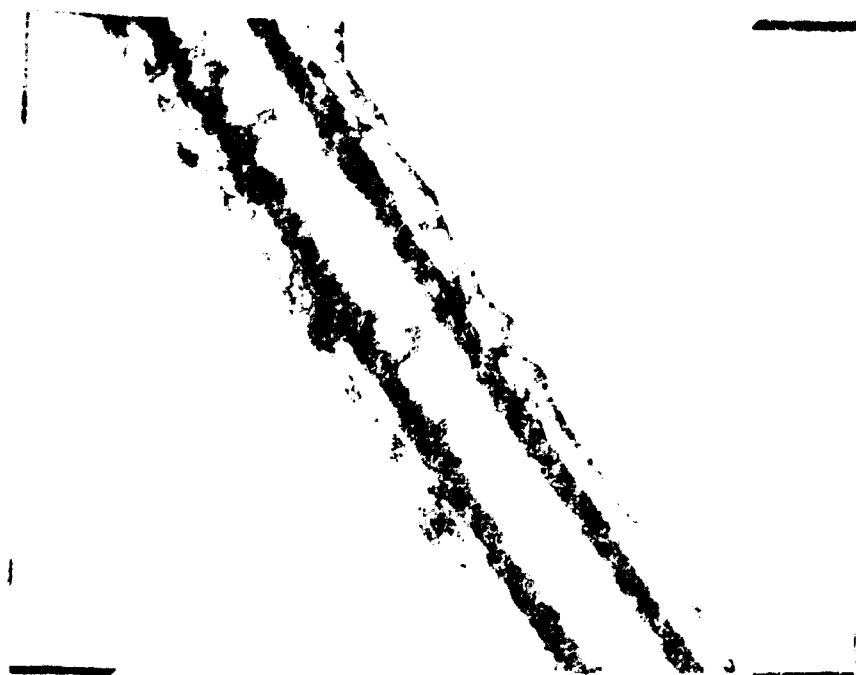


Fig.4: Microphotograph of fibre from clay after 15 months storage in $\text{Ca}(\text{OH})_2$ solution, 620 x magnification. Direct light.

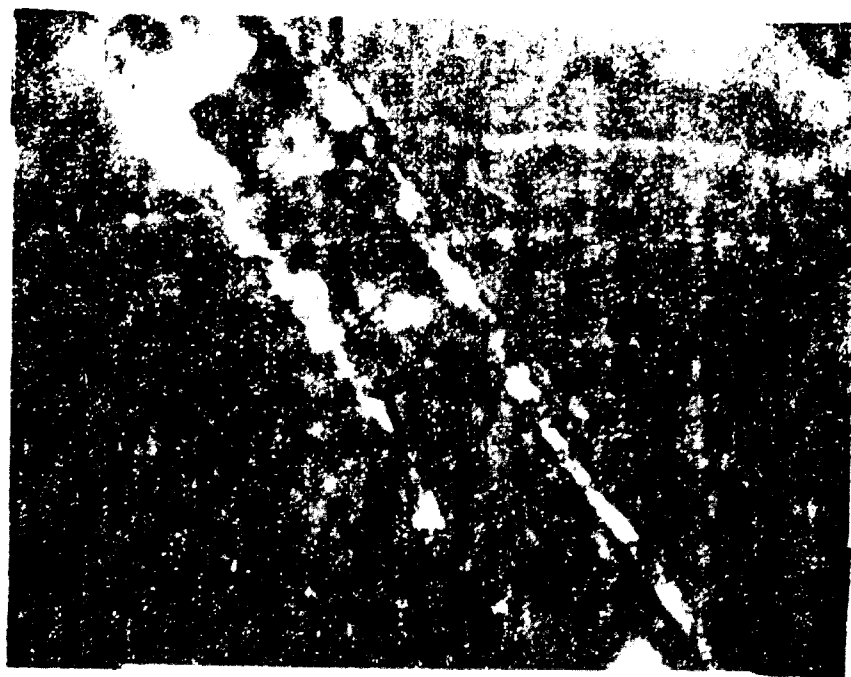
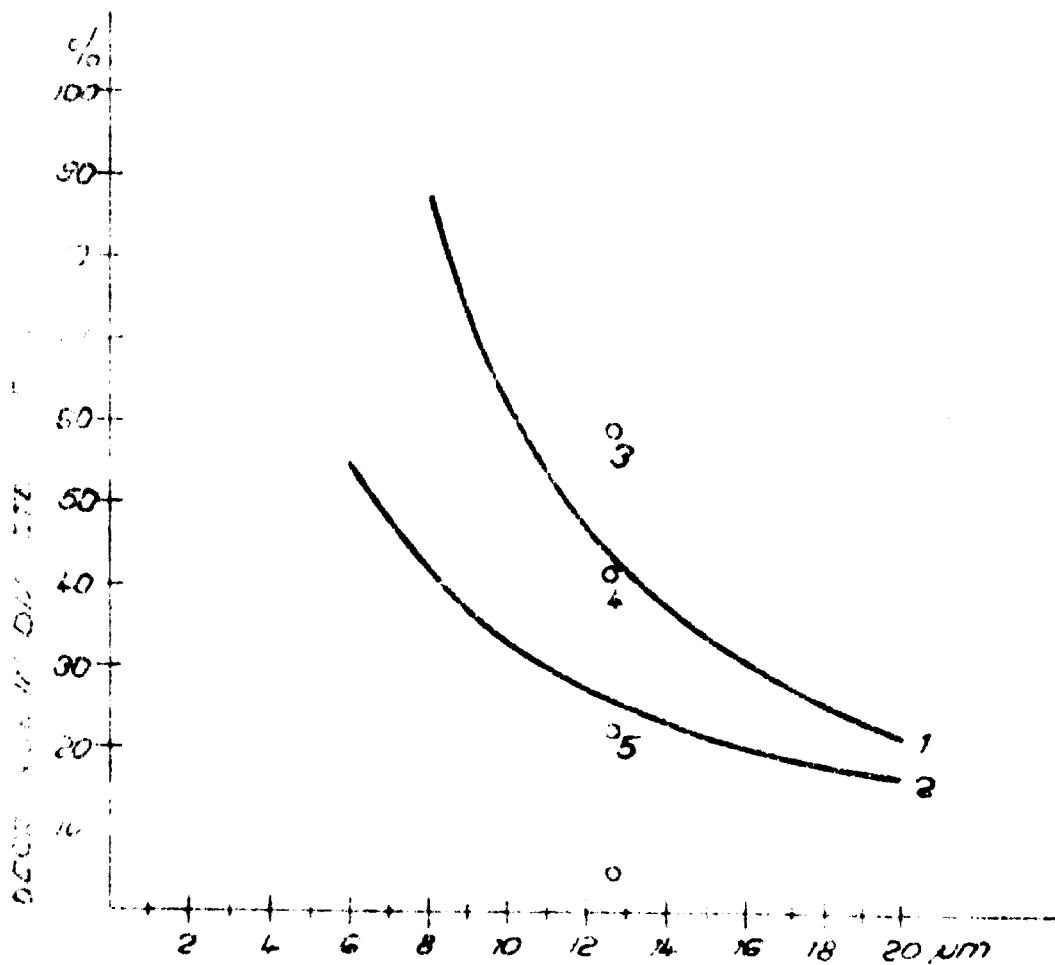


Fig.5: Same photo as in Fig.4. Polarised light, crossed nicols.

FIG. 6 RESISTANCE OF FIBRE IN NO OH SOLUTION

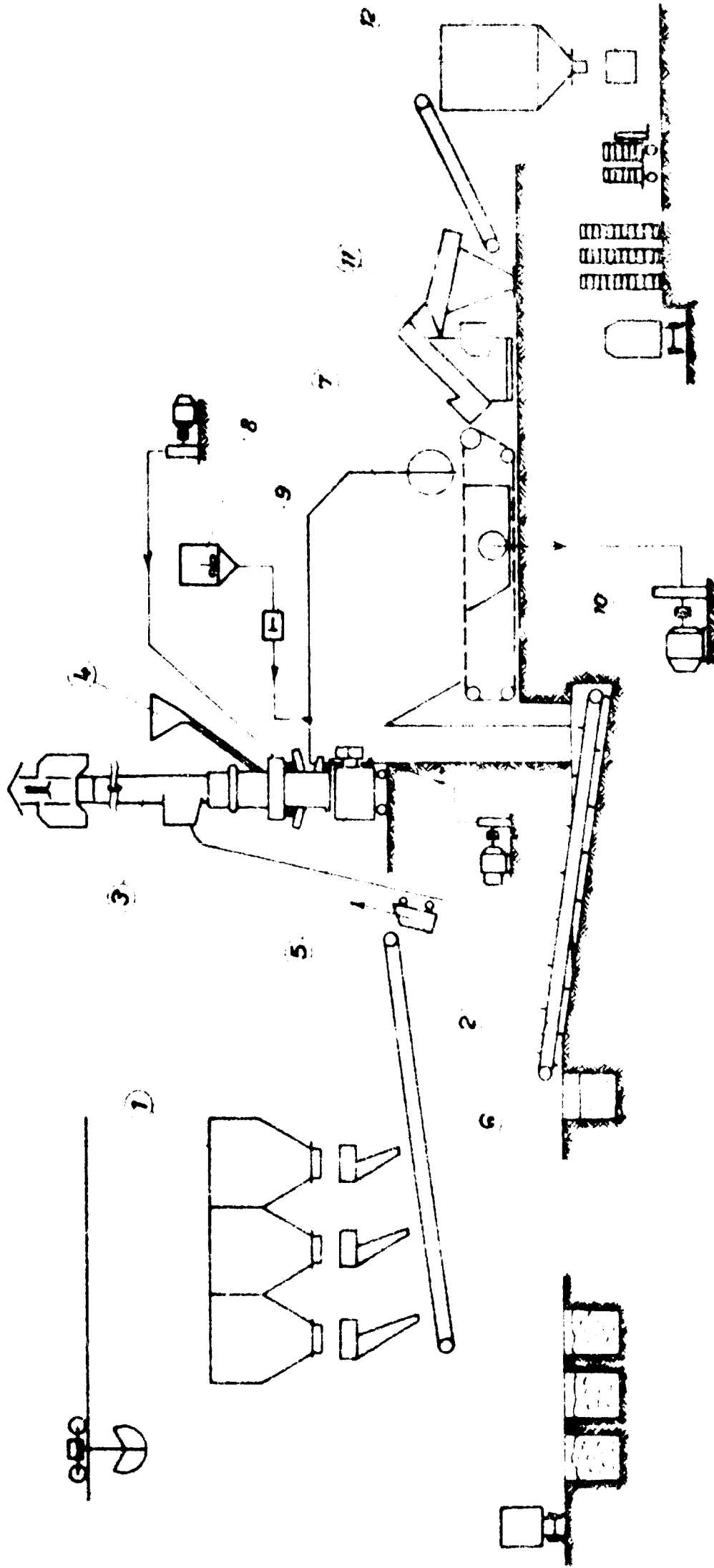


ORIGINAL DIAMETER OF FIBRES

1-BASALTIC FIBRE, 2-A-FIBRE, 3-FIBRE FROM GLASS

4-G4, 5-FIBRE FROM PREFORMED PYREX

FIG. 7 : PRODUCTION SCHEME OF A - FIBRE MANUFACTURE



- 1 STORAGE FOR RAW MATERIALS AND COKE
- 2 FEEDING APPLIANCE
- 3 CUPOLA FURNACE
- 4 AUXILIARY STORAGE WITH CHARGER
- 5 PULPING MACHINE
- 6 SELF CONVEYOR
- 7 SPINNING MACHINES

- 8 STORAGE FOR EMULSION
- 9 HIGH PRESSURE PUMP
- 10 VENTILATOR
- 11 GASKETING MACHINE
- 12 AUXILIARY STORAGE
- 13 PACKING MACHINE

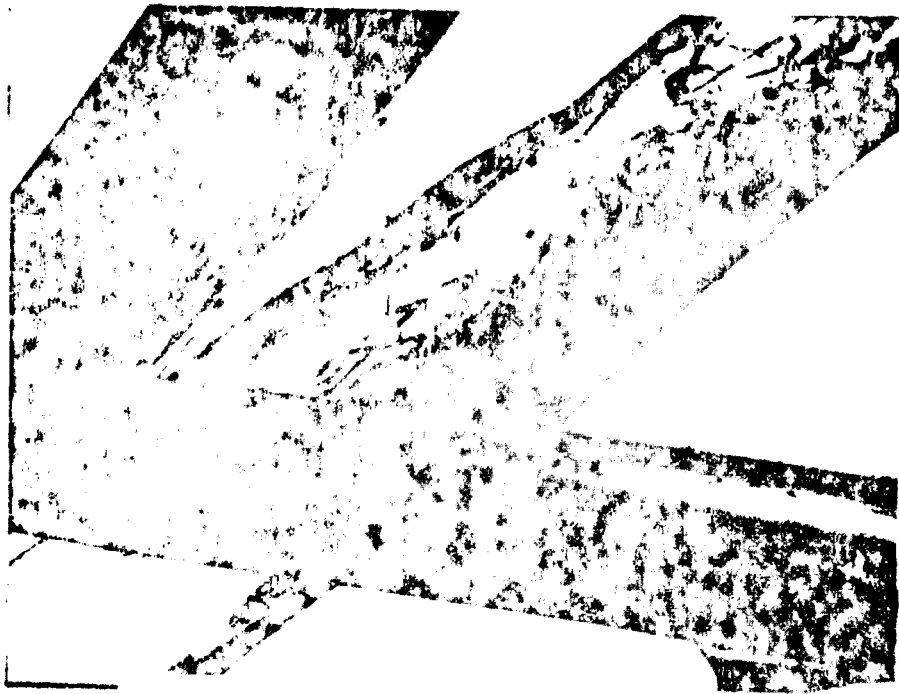


Fig.8: Photo of fibre surface, sprayed during pneumatic transport. Electron microscope. Magnification 8 400 x.



Fig.9: Microphotograph of A-fibres after 15 months of storage in $\text{Ca}(\text{OH})_2$ solution. Magnification 620 x.

FIG 10 RESISTANCE OF A - FIBRE IN CO(OH)₂ SOLUTION

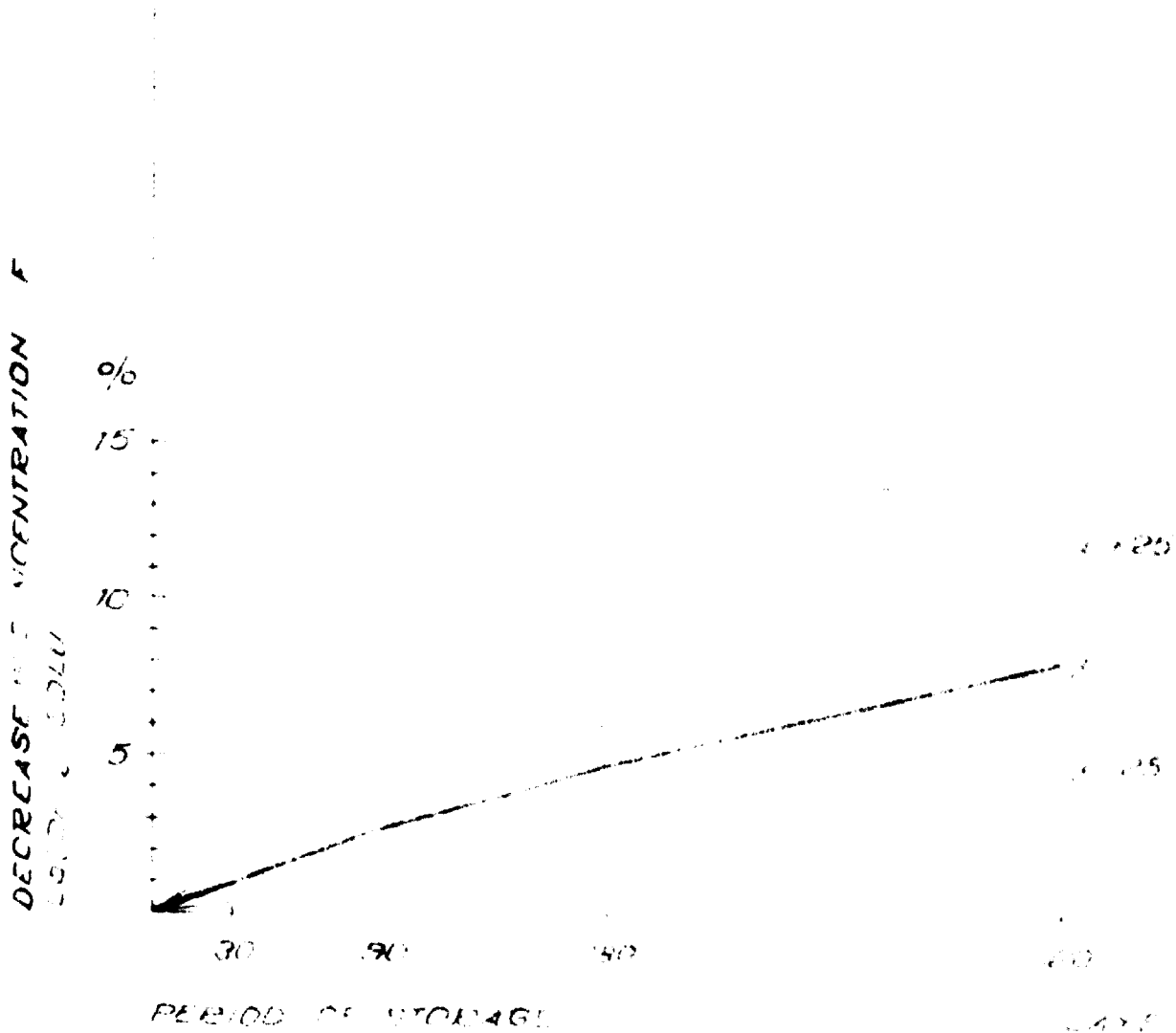




Fig.11: Microphotograph of α -titane in asbestocement
after 18 months storage. Magnification 1500x.



Fig.12: Surface of rock face. Photographed.
Magnification 7.5x.



Fig.13: Surface of α -form of streptocyan.
Magnification is 140x.

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