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CONTENTO

Chapter	•	!ma
1.	Introduction and general considerations	1
11.	Water repellent finishes	4
111.	Finishes impacting resistance to micro-organisms	6
17.	Flame retardant finishes	•
٧.	References	14
	Tables 1 through 11	17

1. INTRODUCTION AND GENERAL CONSIDERATIONS

A review of "Duncble Semishes" published by the "mited Nations in 1970(1) summarizes the requerements for functional properties in various end uses as shown in Table 1, and starts that:

"Present levels of finishing technology as regards available chemicals and technical equipment, permit the fulfillment of almost all requirements of consumers."

in fact, the demand for specific functional properties is continually altered by technological and sociological change, and some of the assumptions made just five years ago are no longer valid. Thus, it is appropriate to review again concepts and approaches which apply to the finishing of febrics, specifically cotton-containing fabrics, in the context of centinuing technological progress and of current requirements in the market place.

property they are designed to import or improve. The major areas of concern and of recent progress for durable finishing of cotton-containing fabrics are shown in Table 2. Easy-care finishing has been reviewed and discussed by others at this meeting(2). Finishes which improve release of oily soils are of particular importance for polyester/cotton blend fabrics of high pulyester content, but are not generally required for 100% cotton. This paper will review approaches to durable finishes which are important for imparting water repellency, resistance to micro-organisms, and flame resistance to 100% cotton fabrics in which a high

cotton content preserves the essential performance characteristics of cotton.

The continuing search for improved products and techniques for durable finishing, coupled with new knewledge of the chemistry, structure, morphology and fundamental properties of Fibrous assemblies, has produced a new level of insight into conceptual definitions of finishing problems, of specific approaches, and of meanineful evaluation of results obtained. Thus, the prognatic classisication of functional finishes shown in Table 2 may be supplemented by a classification of Agrable finishes based on the predominent mechanisms of finish-substrate interaction as outlined in Table 3. Since crease-resistance and easy-care performance depend primarily on crosslinking of cellulose chains in the cotton fiber, finishes designed for this use must include polyfunctional molecules, small enough to penetrate the fiber rapidly, and capable of reaction with hydroxy) groups under mild conditions. Soir release and water repellent finishes on the other hand need affect only the surface properties of the fiber (wettability), and can be deposited on fiber surfaces as continuous films or coatings which do not react with the functional groups of the substrate. Resistance to microorganisms is generally attained through the presence of compounds which prevent growth of fungi, mildew or bacteria: specific water insoluble compounds reportedly exhibit adequate durability, and yet diffuse sufficiently to come into contact with the micro-organisms and inhibit their growth. In the case of flame retardant finishes, effectiveness is determined by the chemical composition of the finish and by the presence of specific elements in amounts sufficient to minimize flame propagation in the treated fabric. The distribution of the finish, and the mechanism of insolubilization do not affect flame retardant effectiveness significantly, although these factors

have important secondary effects on durability of the finish and on fabric properties (pulser tree (lambert) to)

The chemical composition of the finish, and the finish-substrate interactions outlined in Table 3 are primary considerations in selecting approaches to the modification of specific functional properties. However, the processing conditions required for application and the methods available for evaluating the effects obtained are also essential components of finishing developments.

Finishing processes traditionally have been based on impregnation from aqueous solution, drying, curing, and--in some cases--after-washing. recent years, a considerable amount of work has been carried out on "nonconventional" processes for fabric finishing. The potential usefulness of non-aqueous systems, where water-sensitive reagents could be employed, has been investigated extensively(3), liquid ammonia has been suggested as a viable medium for several specific finishing processes(4) and treatments in which fabrics are exposed to a gaseous environment containing reagents and/ or catalysts have been evaluated on a commercial scale(5). However, an overwhelming proportion of the yardage produced commercially is finished by conventional processing sequences in which padding from aqueous solution, drying, curing and washing still constitute the essential steps--preferably carried out as parts of a continuous operation. The specialized finishing techniques discussed in this paper have actual or potential commercial significance, do not require extraordinary machinery, instrumentation or devices, and thus can be implemented in textile mill equipment which is either available, or readily adapted to specific processing requirements.

CONTRIBUTION FRANCISHS

Fabrics to ared with water repelle it finishes are no readily wet by water--but retain their perposity, because and appearance. The surface of individual fibers is ambified by the finish. while interstices between Tibers and marweak yaras rumain, in predciple, essentially unchanged. (By contrast, the effectiveness of water reperient coatings depends on the formation of a continuous hydrophotic surface which aliens the porosity of the fabric, and its permeability to air and moisture.) Several classes of durable water repellent finishes have attained commercial success. These are summerized in Table 4. Excivatives of long chain fatty acids (such as steeramidementhy) by ild inium chienthe) were developed many years ago, and were thought or as reactive finisher, capable of covariant bending with the hydroxyl groups of collulate withinguis. In the light of recent knowledge of the fiber (cotton) summittee and reactivity, the effects and durability obtained with this type we dompower and now indespreted at resulting from deposition of hydrophobic, valor hasoluble compounds on liber surfaces rather than -- the formation of covalence bonds. The silicone and fluorochemical polymens form a hydrophobic film on fiber surfaces. Effect iveness depends on the continuity of the film. and on the absence of conteminents which enhance wettings by water. Since the polymers are generally applied from aqueous emulsions by padding, drying and curing. the presence of residual enulisifiers on the finished fabrics can pose problems. Herufacturers of the silicone and Pubrochemical emulsions have developed formulations containing "non-reweiting emulsifiers" specifically selected to avoid this shortcoming. The amount of water repellent applied is generally 376% (on the weight of fabric treated) for the fatty compounds, 1-2% for the

silicones and 0.5% or lower for the fluorochemical. In part, these differences retriect the countries economies of the compounds, and in part also the high chemical cost of the lilicone and of the fluorochemical polymers - Formulations have also been developed and synergistic effects have been claimed for combinations of faity acid derivatives and fluorochemicals(6). Advantages in efficiency have been claimed for the application of silicones and of fluorochemicals from chlorinated solvents(7) but it is not known whether this approach is used commercially. The details of specific formulations and procedures used for the application of water repellent finishes are generally considered proprietary, and are dependent on the fiber composition and construction of the fabric treated. Construction is particularly important to determining the required amount of water repailent, and the optimum conditions of processing. The level of water repellency obtained is evaluated by standard test procedures, and standards for metting and water penetration of fabrics have been defined by the American National Standards Institute(a). From the results of these tests, industry defines the performance of a given fabric as "water repellent" (resists wetting), "shower resistant" (protects against water penetration during a brief shower), "rain resistant" (protects against water penetration in moderate rain), and "storm resistant" (protects atainst water penetration in heavy rain).

TITL LINESPEC IMPARISING PROSECTABLE TO MICRO OPGANISMS

fungi) can, in principle, be grouped according to several approaches summarized in Table 5. The fiber may be chemically modified to resist attack: for example, conversion of reliables to diacetate or triacetate yields a fiber which does not support mold growth, and even acetyl degrees of substitution lower than 2 have a significant effect. This approach essentially implies a chemical transformation of polymer and fiber which has far reaching consequences on the properties and performance of the products.

In a second approach, a resin which provides an impermeable barrier to bacteria and fungi is added to the fabric. This approach is generally used for industrial fabrics, where considerations of porosity, permeability and appearance are secondary.

been obtained on cotton in conjunction with the application of crosslinking agents, of fire retardant finishes, and of other finishing agents insolubilized in situ. The most effective approach to not resistance through application of resins has been the polymerization of N-methylol-melamine derivatives in water-swollen fabric(9) which is known as the ARIGAL(R) process (Ciba Geigy). Cotton fabric treated in this manner is reported to retain 100% of its strength after soil burial for up to 15 weeks. The mechanism by which polyfunctional N-methylol compounds and other finishes capable of generating formaldchyde inhibit the growth of micro-urganisms in treated cotton may entail restricted diffusion of moisture into the cross-linked fibrous substrate, as well as slow controlled release of formaldchyde from treated fabric or from the resin network formed in or on fibers in the

course of application.

The direct of the second the base where he at the envert resistance to micro-organisms as the deposition of active compound, which are effective bactericides or fungionnes. These must be afficient, non-discoloring, durable to washing and, above all, non-roxid to bumon. The balance between activity, which implies diffusivity in the sy tem, and durability to washing, which implies lack of solubility and resistance to leaching in water is particularly difficult to attain. Few among the numerous claims found in the patent literature describe compound, which are in commercial use. Tables 6 and 7 show compounds which have been used commercially for fabric finishing: those listed in Table 6 have varying degrees of solubility, and generally limited resultance to leaching. For the organometallic compounds shown in Table 7, activity depends on the presence of heavy metal. Copper compounds are both fungistatic and bacteriostatic, and extensively used for preservation of tent convas and sandbags, even though they impart a green color to the treated fabrics. Tin compounds have the advantage of being colorless, but they exhibit varying degrees of toxicity. Zinc compounds have lower activity, but lower toxicity as well. Mercury compounds (not shown in Table 7), have been essentially ruled out by the Environmental Protection Agency due to potential cumulative toxicity effects. Generally speaking, compounds designed to inhibit mildew and not are used primarily for military and institutional fabrics. They are effective in low concentrations and applied by pad/dry/techniques from emulsions or organic solvents. Durability to leaching depends on the solubility of the specific compound, and durability to laundering is at best limited. The relative effectiveness of some of the compounds, shown in Tables 6 and 7, and the concentrations used in fabric finishing, vary greatly: for example, for

the highly effective (and now forbidden) phenyl mercuric acetate, a 0.01% concentration as additional to give 50° corength resolution after ten days' soil burnel. For copper -8 -quinolimetate, the needed concentration is 0.05%; and for salicylamilide, 0.5%.

IV. FLAME RETARDANT FINISHES

Research and development work on flame retardant finishing has been greatly stimulated by recent legislation in the U.S., and in other countries. At this time, this field is perhaps the most important, and certainly the most active area of investigation in textile chemistry.

Durable flame retardant finishes for cotton-containing fabrics may be defined as those which impart self-extinguishing behavior, and are not removed in laundering. A fabric which is described as "self-extinguishing" when placed in a vertical position and ignited at the bottom, will not continue to burn after the source of ignition is removed. The conditions of testing (specimen size and mounting, source, site and time of ignition, etc.) must be carefully defined when indicating self-extinguishing behavior, but, in the U.S., the term is generally used with reference to the conditions specified for the test of the U.S. children's sleepwear standard (23) which is a modification of previously developed vertical flammability tests.

Finishing of cotton and cotton blend fabrics to impart self-extinguishing behavior and flame resistance durable to laundering, involves problems which are far more complex than those encountered in other finishing processes. Firstly, the amount of insolubilized or fixed finish required is considerably greater than for other functional finishes: the added non-fibrous material tends to impair the aesthetics and performance properties

formity and durability must be regorously controlled since deviations can have disest has consequences in litigation or product liability suits.

Lastly, results of tests for the evaluation of flarmability are critically dependent on a large number of variables and laboratory testing of finish fabric becomes an exacting and costly part of finishing development, and quality control.

Interest in the application of flame retardant compounds to cotton fabrics dates back to the seventeenth century(24), when clay and plaster paris were applied to canvas used for theater curtains as flame retardant. From that time until World War II, investigations of textile flame retard ants were limited to water soluble salts, even though in 1913, William Newry Perkin defined the requirements for flame retardant finishes in woll which still apply today(25):

the feel or durability of the cloth, or cause it to go damp as so many chemicals do, and it must not make it dusty. It must not affect the colors or the design when into the cloth or dyel or printed upon it; nothing (such as arsenic, antimony or lead) of a poisonous nature or in any way deleterious to the skin may be used and the fireproofing must be permanent, that is to say, it must not be removed even in the case of a garment which may possibly be washed 50 times or more.

Furthermore, in order that it may have a wide application, the processmust be cheap.

Progress from Perkin's time to World War II is well documented in a book by Little(26), which remains a classic cornerstone of our current knowledge on flame retardant finishing techniques for cotton fabrics. In

this book, important generalizations are set forth concerning the chemistry of fire retaidants for cotton. The outstanding effectiveness of phosphorus is recognized; and the foundation is laid for most subsequent work on the mechanism for inhibiting combustion in cotton fabrics, and on approaches to the development of durable finishes.

It is now established that phosphorus-containing compounds exert their flame retardant action by decomposing to species which alter thermal degradation reactions in the substrate and decrease the concentration of combustible products while enhancing dehydration reactions (27). It has also been shown that phosphorus containing flame retardants are more effective when used in conjunction with nitrogen-containing compounds (28). In a 1006 cotton fabric, the presence of a sufficient amount of phosphorus (2% to 4%, depending on fabric construction), preferably in conjunction with nitrogen (2% to 6%) effectively imparts self-extinguishing behavior. When durability of the flame retardant finish is required, the phosphorus and mitrogen must be insolubilized either in substitutent groups covalently bonded to cellulose hydroxyls, or in a crosslinked polymer network formed in or on fibers. Phosphorus and nitroger thus must be part of molecules capable of forming a three-dimensional polymer "in situ" and/or of reacting with the fabric substrate during the finishing process. The amount of finish required to attain the desired phosphorus content in the treated fabric depends primarily on the phosphorus content of the "active" flame retardant species, and on the insolubilization yield (efficiency) in the finishing process. The chemical stability of the insolubilized products determines the amount of finish retained in laundering (durability). Phosphorus compounds used in durable flame retardent finishes must must the

requirements summarized in Table 8 and, in addition, compatibility with nitrogenous co-reactants and with additives must be considered. The essential requirements for the finishing process include absence of toxic compounds, and, preferably, the use of conventional equipment and procedures in the finishing plant. The overall objectives of satisfactory (or adequate) fabric performance (including but not limited to flame resistance), aesthetics, and economics must of course be met.

Numerous organophosphorus compounds and formulations have been evaluated for flame retardent finishing of cotton fabrics over a period of 20 or 30 years. Many have failed to meet one or more among the critical requirements outlined, and thus have not attained commercial status. At this time, only three chemical systems are used commercially in the U.S. The summiry presented in Table 9 indicates the organophosphorus compound, the other essential components of the finish (if any), and the principal mechanism of finish insolubilization for these systems. Salient information on the finishing processes is summarized in Table 19.

Performance properties of the finished cotton fabrics are considered acceptable, with advantages in fabric hand or strength retention claimed for specific fabric/finish combinations by individual chemical manufacturers or textile mill processors. However in all instances, the finish stiffens the fabric somewhat, causes tensile strength losses of about 20-30%, and tear strength losses of about 35-45%, depending on the fabric construction and finish add-on. Treated fabrics are self-extinguishing according to the vertical flammability test specified in the children's sleepwear standard (23), initially and after 50 launderings with high phosphate detergents.

Accumulation of calcium and magnesium salts (which may occur on laundering in hard water without adequate amounts of phosphate) impairs flame

by rinsing with acid. Eleaching with hypochlorite, on the other hand, accelerates less of finish and flame resistance is irreversibly lost. In brief, nonmercial flame retardant finishes for cotton fulfill the requirements outlined (Table 8) only in part. More efficient organophosphorus compounds are needed in order to decrease the amount applied and, therefore, side effects and cost.. Research to develop new reagents is continuing with this objective in mind. For example, results obtained in the laboratory with amides of chlaromethyl phosphonic acid and of methyl phosphonic acid (33), (34) demonstrate considerable progress towards the goal of importing self-extinguishing behavior at low weight gains.

Finishing of polyester/cotton blend fabrics is far less advanced. The problem is complex, since the two fibers differ in their physical response. and in the course of their chemical degradation at elevated temperature. Flame retardant finishes that are effective on 100% cotton do not necessorily decrease the flammability of polyester/cotton blends--at least not in terms of the results obtained in the vertical ilammability test for selfextinguishment(23). Finishes applied to blends from aqueous solutions tend to penetrate cotton fibers preferentially, and the concentration of reactive compounds in the cotton becomes exceedingly high. Furthermore, the concentration of hydroxyl groups in the total substrate is lower in the case of blends (in proportion to the polyester content), and the stoichiemetry of the insolubilization reactions is inevitably affected by these factors. Extensive research activity on durable flame retardant finishes for polyester/cotton blends during the last few years has resulted in considerable progress(35), and some promising approaches are emerging even though comm cially finished blend fabrics containing 50% or more polyester are not

evailable. Table 11 shows some finishing systems claimed to be effective on polyester/conton blends. Frame netardancy is obtained through the presence of phosphorus, or brownine, or combinations of these elements. In these finishes, compounds containing phosphorus and/or brownine are not reacted with the substrate. Adequate durability to washing is related to the solubility characteristics of the flame retardant compounds, and to the use of resin binders in the finish formulation. Large amounts of the finishes are needed to impart self-extinguishing behavior to the fabric and fabric stiffmess is a major problem in most instances.

A brownne-containing polyester (Dacron 900F $^{(R)}$), a copolymer in which part of the ethylene glycol is replaced by the brownnated glycol below)

"conventional" polyester fibe in polyester/cotton blend fabrics is replaced by this copolymen fiber (which contains approximately 6% Bromine), or by an equivalent modified polyester fiber, lower amounts of flame retardant finish are required to meet a given flammability test and undesirable effects on febric properties are reduced. The major problem then becomes one of cost.

In semmery, the goal of imparting durable flame resistance to 106% sotten fabrics without significantly altering aesthetics and performance properties is now within reach. For polyester/cotton blends, research must continue for some time to come and new approaches must be developed before a comparable status is reached.

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Table 1

	CREASE R'SISTANT (WASH AND WEAR)	WATER RCPEL- ENT	OLEO- PHOBIC	FLAME- PROOF	RACTERIO- STATIC & FUNGISTATIC
MOVEN (OUTERWEAR)	0	(L)		(E)	•
MOVEN (LININGS AND UNDERWEAR)	0	\bigcirc	\bigcirc		(
WITTED (OUTERMEAR)	()	(D)	•	•	•
INSTITED (LIMINGS AND UNDERWEAR)	()	\bigcirc	\circ	(②
DED LINEN	O	\bigcirc	0		•
UPWELSTERY (CURTAINS, GRAPETS, ETC.)	(3)	O	©	(1)	•
INTUSTRIAL FAURICS (AMNINGS, TYVIS, ETC.)	0	0	(b)		•



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Table 2. Durable finishes for cottna-containing februs

TYSTRED PROPERTY	CURPENT TECHNOLOGICAL SPPROACHES		
EASY CARE (LUMBLE PRESS)	MEACTION OF COTTON WISH CROSSLINKING AGENTS		
FREASE OF CILY JUILS	(1) MYDERDERIC IC. ET AR POLYMERS. (3) COLYMPIS COMPAINING PERFLUORDALKYL GROUPS AND PYLHORPILIC SEGMENTS.		
MATER REPELLENCY	(#) COMPOUNDS CONTAINING LONG CHAIN (>C ₁₆) FERME GROUPS. (8) POLYSILOXANES. (C) FOLYMERS CONTAINING PERMEDOMOREKYE GROUPS.		
RESISTANCE TO MICRO-ORGANISMS (HILBEN, ROT, BACTERIA)	(A) ORG/ NOME CALLID COMPOUNDS (B) MALOGENATED PHEMOLS (C) ANILIDES (D) SUBSTRATE MODELICATION		
FLAME RESISTANCE	(A) DROUMOPHUSPHORUS COMPHUMOS (B) HALDREMATED COMPONINS		

Table 3. Finish-substrate interactions in Burable finishes for cotton-containing fabrics

FINESH-SUBSTRATE SEPERACTIONS	GASY CARE (CROSS- LINKING)	SOIL RELEASE	MATER REPEL- LENCY	MESTSY- ANCE TO MICHO- ORGANISMS	FLAME RESIST- AMEE
PROTECT OF THE PROTEC	•	•	•	•	•
THESE MESCHALIZATION T. STATEM MESTIGN (MEACTION WITH GM)	•				•
2. W STYS POLYNERS-					•
S. SHOPLUFICITATION OF PREFERED POLINER		•	•		•
TO SEPTEMBLY OF SEPTEMBLE OF SE			•	•	•

Table 4. Burable water repellent finishes

CHEMICAL TYPE	PROTOTYPE	SUBABILITY TO WASHING DRY CLEANOUS	
DERIVATIVES OF LONG CHAIN FATTY ACIDS	[CPBN31camx - CHTNCPNP]C1_	eres rest	
PULYSILONNES (SILICONES)	CN ₃ CN ₃ CN ₃ CN ₃ CN ₃	HOSENARE GOOD	
POLYPENPLUSING- ALRYL ACRYLATES (I'L USINGCHEROCAL)	Coly - Col}-	6800 6880	

Table 5. Approaches to antibacterial and antifungel finishes

MANAN	PRINCIPLE	ENOPLES
CHEMICAL MODEF (CATION	CHANGE IN CHEMICAL STRUCTURE OF SUBSTRATE	A. AGETYLATION B. CYANGETHYLATION
RESIN CARDION	COATING FIBERS WITH SUB- STRUCE WHICH IS IMPER- VIOUS TO MICHO-CHOMISMS	••
IN SITY FORMETION OF CROSSLINNED RESIN	CONTROLLED RELEASE OF PONIAL DENY DE AND/OR CELLULOSE CROSS- LÍNKING	POLYCONDENSATION PRODUCTS OF A. POLYMETHYLOL MELANGHE B. TETRAKIS MYRADHY METHYL PHOOPHO- MINH CHLORIDE (THPC)
ACTIVE DESIGNACIONE OF PRODESIGN CONTRACTION	CONTROLLED RELEASE OF ACTIVE HUMEDIENT FROM TREATED SUBSTRATE	A. SURPACE ACTIVE AGENTS D. PHENOLS C. ANILISES D. GREANWETALLIC

Table 6. Effective compounds in finishing for resistance to micro-organisms

TYPE	TRUCTURE (EXAMPLES)	REF.	APPLICATIONS AND LIMITATIONS
SURFACE ACTIVE AGENTS	CH2-N-CH3 CH3 CH3 CH3 CH3 R = C ₁₂ H ₂₅ - , C ₁₄ 14 ₂₉ - QUATERNARY AMMONIUM COMPOUNDS	(10)	BACTERICIDES. WATER SOLUBLE (LIMITED DURABILITY TO LAUNDERING)
PHENOLS	C1 C1 OR C1 C1 R = H , -COC ₁₁ H ₂₃ PENTACHLOROPHENOL (AMD ESTERS)	(11), (12)	ROTPROOFING AGENTS: SODIUM SALT IS WATER SOLUBLE AND LEACHED EASILY. (R=Na) LAURATE ESTER (R= COC11N23) LESS EFFICIENT BUT MORE LEACH-RESISTANT
	OH OH CH2-C1 C1 R.2'NETHYLENEBIS 4-CHLOROPHENOL	(13), (14)	ROTPROOFING AGENT. APPLIED FROM ORGANIC SOLVENT OR EMULSION.
MILIBES	OH CONH -	(15)	HACTERICIDE AND ROTPROOFING. APPLIED FROM ORGANIC SOLVENT.
	X = C1,Br MALOBENATED SALICYLANILIDES	(16)	HALOGENATED DERIVATIVES NAVE SHOWN SOME SIDE EFFECTS (CONTACT SENSITIZING).

Table 7. Organometallic compounds in finishing for resistance to micro-organisms

STRUCTURE (EXAMPLES)	REF.	APPLICATIONS AND LIMITATIONS
COPPER-8-QUINOL INOLATE	(17). (18)	ROTPROOFING OF OUTDOOR FABRICS, SOLUBILIZED WITH OLEICACID OR CASTOR OIL FOR APPLICATION FROM SOLVENT. IMPARTS GREEN COLOR.
ZINC NAPHTHENATE	(19)	MOLD INHIBITOR, FUNGISTATIC ACTIVITY.
ZINC-2-MERCAPTO DENZOTHIAZOLE	(20)	FUNGISTATIC ACTIVITY
[(CH3)NCS- 2n++ S 2 ZINC DINETHYL DITHIO- CANDANATE	(20)	FUNGISTATIC ACTIVITY. CAN BE PRECI- PITATED IN SITU FROM SODEUM SALT AND Zu SULFATE.
(C _q N _g) _g Sn-X X = CN _g COO, C _g N _g COO TRESUTY, TIN ACETATE, ORNEOATE	(21), (22)	FUNGISTATIC AND BACTERIOSTATIC EFFECTS ON FABRIC AT VERY LOW CONCENTRATION.

Table 8. Phosphorus compounds used in durable flame retardant finishes

REQUIREMENT	COJECTIVE
CAPABLE OF INSOLUBILIZATION BY POLYMERIZATION WITH COMMINERS AND/OR BY REACTION WITH ON	SMRADILITY OF FINISM TO LANGERING
NJOH PHOSPHONUS CONTENT	LAMEST POSSIBLE ABBED WEIGHT IN TREATED FABRIC
LON CARBON CONTENT	LOWEST POSSIBLE AGGED FUEL IN TREATED FAGRIC
AGGENCE OF IGNIC SITES (AFTER INSOLUBILIZATION)	AVOID ION EXCHANGE WITH NETALS IN LAUNGERING

Table 9. Commercial chamicals for flame retardant finishing of cotton fabrics

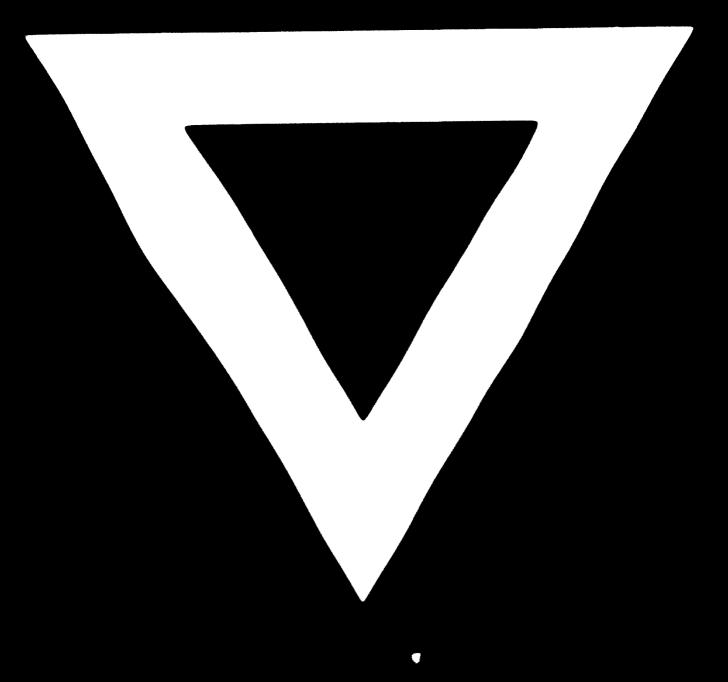
CONTOURS CONTOURS	COREACTANT(S) REQUIRED	INSOLUBILIZA- TION MECHANISM	TRADE NAME (MANUFACTURERS)	RCF.
(CH30)2P-CH2CH2CH2CH	NCH ₂ OH -	REACTION WITH CELLULOSE OH	PYROVATEX CP (CIBA-GEIGY)	(29)
[(MBCM ₂) ₄ P]+ X- X-C1, 6M, etc.	m ₂ com ₂ , etc.	IN SITU POLY- MERIZATION WITH COREACTANT (POLYCONDENSA- TION)	THPC, THPOH (HOOKER CHEMI- CAL, AMERICAN CYANAMID, ALBRIGHT AND WITSON)	(25), (30), (31)
CH - CH ⁵ (-0-6-601 ⁵ CH ⁵)	CIF-CHCOINCHFOR	IN SITU POLY- MERIZATION (FREE RADICAL)	FYROL 76 (STAUFFER CHEMICAL CO.)	(32)

Table 10. Flame retardant finishing processes for cotton fabrics

CHEMICAL SYSTEM (see Table 9)	PROCEDURE (Essential steps)	S FINS	SM (RAMSE)	FINISHED PARTIC (resea)
PYRONATEX CP+ N-NETNYLOL MELAMINE+ +ACID CATALYST	PAD/DRY; CURE (320-350°F); WASH.	39-40	29-30	2-2.5
THPSH (neutralized THPC)	PAD; BRY TO 10% MOISTURE; EXPOSE TO NM3 GAS; OXIDIZE/MASH.	39-40	25-35	3-6
FYREL 76 + N-HETHYLOL ACRYLANTOE + + PREE MADICAL CATALYST	PAD/DRY; CURE (300-350°F); WASH.	25-36	20-30	2-4

Table 11. Durable flame retardant finishes for polyester/cotton blend fabrics

FLAME RETARDANT COMPOUNDS IN FINISH	INSOLUBILIZATION ON FABRIC	REF.	PROBLEMS IN TREATED FABRIC
CONDENSATION PRODUCTS OF TETRAKISHYDROXYLMETHYL PHOSPHONIUM CHLORIDE (THPC) (PHOSPHONIUM OLIGOMER)	IN SITU POLYMERIZATION	(36)	HIGH % P NEEDED; STIFFENING; HIGH COST.
THPC + N-METHYLCL CUMPOUNDS + TRIS-2.3-DIBROMO-PROPYL PHOSPHATE (TBPP)	TBPP "TRAPPED" BY POLYMER FORMED IN SITU	(37)	STIFFENING; LIMITED DURABILITY OF BROSSINE
TWPC + N-METHYLOL COMPOUNDS + POLYVINYL BROWLDE (PVB)		(36)	STIFFENING; DISCOLORATION
DECABRONO DIPHENYL OXIDE + + ANTINONY OXIDE + ACRYLIC BINDER (P. 44)	BINDING OF SOLID INSOLUBLE FLAME RE- TARDANT COMPOUNDS.	(39)	STIFFENING; DUSTING, WHITEMING; LIMITED MESISTANCE TO ADRASIVE MEAR.



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