

FLAME RETARDANT AND WATER-RESISTANT TREATMENT OF FABRICS

This invention relates to an improved method of treating fabrics to impart flame-retardant and water-resistant properties thereto and to a fabric treated thereby.

A known process for the flame-retardant treatment of fabrics including cellulosic (e.g. cotton) fibres consists of impregnation of the fabric with an aqueous solution of a poly(hydroxyorgano) phosphonium compound, for example, a tetrakis (hydroxyorgano) phosphonium salt. Alternatively, the poly (hydroxyorgano) phosphonium compound may comprise a condensate with a nitrogen - containing compound such as urea. Following impregnation, the fabric is dried and then cured with ammonia to produce a cured, water-insoluble polymer which is mechanically fixed within the fibres of the fabric. After curing, the polymer is oxidised to convert trivalent phosphorus to pentavalent phosphorus and the fabric is washed and dried. Fabrics treated according to the aforesaid process and garments made from such treated fabrics are sold under the Registered Trade Mark PROBAN of Albright & Wilson Limited.

We have now found that the addition of one or more protonated and neutralized amines to the impregnation solution increases the efficiency of fixation of the phosphonium compound within the fibres, improves uniform distribution of the phosphonium compound in the system, and leads to improved flame-retardant and increased water-resistant properties.

Accordingly, the present invention provides a method of treating fabrics to impart flame-retardant and water-resistant properties thereto, said method comprising impregnating the fabric with an aqueous solution including a poly(hydroxyalkyl) phosphonium compound, in which there is added to the impregnating solution one or more primary, secondary or tertiary aliphatic amines having from 12 to 20 carbon atoms, said amines having been protonated and neutralized prior to said addition.

The present invention also provides a flame-retardant and water-resistant fabric treated by the method described in the immediately-preceding paragraph.

The concentration of protonated and neutralized amine in the impregnating solution is suitably in the range 0.05% to 3% by weight, preferably in the range 0.1% to 1% by weight, especially about 0.3% by weight.

In a preferred embodiment of the present invention, the protonated and neutralized amine consists essentially of n-octadecylamine.

In an alternative embodiment of the present invention, the protonated and neutralized amine comprises a mixture of primary aliphatic amines having from 16 to 18 carbon atoms.

Suitably, the poly(hydroxyalkyl) phosphonium compound is a tetrakis (hydroxyalkyl) phosphonium (hereinafter THP) compound, for example a [THP]⁺ salt.

The amines are protonated and neutralized according to the present invention by means of a weak organic acid, for example acetic acid. The protonated and neutralized amine may therefore consist essentially of octadecylamine acetate.

Suitably, the amines may be obtained in an already-protonated and neutralized state.

Alternatively, the amines can simply be mixed with sufficient acetic acid to achieve protonation and neutralization and the so-treated amines added to the impregnation solution.

The present invention will be illustrated, merely by way of example, as follows:

The following fabrics were treated in accordance with the present invention:

<u>Sample Code A</u>	A satin fabric comprising 60% cotton fibres and 40% polyester fibres and having a weight of 280g/m ²
<u>Sample Code B</u>	A twill fabric comprising 60% cotton fibres and 40% polyester fibres and having a weight of 245g/m ²
<u>Sample Code C</u>	A twill fabric comprising 60% cotton fibres and 40% polyester fibres and having a weight of 315g/m ² .
<u>Sample Code D</u>	A plain-weave, pigment-printed fabric comprising 100% cotton fibres and having a weight of 200g/m ²

The fabrics were impregnated with an aqueous solution containing the following percentages by weight of a precondensate of tetrakis (hydroxymethyl) phosphonium chloride and urea, together with protonated and neutralized amines in accordance with the present invention, the ratio of the phosphonium chloride to urea in the condensate being 2:1 molar:

- A: 42.25% by weight
- B: 42.25% by weight
- C: 39% by weight
- D: 32.5% by weight

The impregnated fabrics were squeezed to a wet pick-up in the following ranges based upon the original weight of the fabric:

- A: 80%
- B: 80%
- C: 80%
- D: 90%

The fabrics were then dried at 120°C and kept overnight at ambient temperature to achieve a moisture content in the range 4 to 8 %, preferably 5 to 8%.

The dried fabrics were cured with gaseous ammonia to cure the precondensate within the fibres of the fabrics, followed by oxidation with hydrogen peroxide, washing and drying.

TABLE I (below) shows the results of testing for flame-retardant properties according to DIN 66083 s-b:

TABLE I

Sample Code	Direction of test	Ignition time (sec)	Afterflame (sec)	Afterglow (sec)	Char Length (mm)
A	warp	3	0	0	7
		15	0	0	125
		3	0	0	6
		15	0	0	75
		3	0	0	5
		15	0	0	
	weft	3	0	0	7
		15	0	0	87
		3	0	0	8
		15	0	0	75
		3	0	0	7
		15	0	0	75
B	warp	3	0	0	20
		15	0	0	110
		3	0	0	13
		15	0	0	103
		3	5	0	70
		15	-	-	-
	weft	3	0	0	12
		15	0	0	95
		3	0	0	15
		15	0	0	82
		3	0	0	20
		15	0	0	103

Continued

TABLE I - continued

C	warp	3	0	0	5
		15	0	0	112
		3	0	0	5
		15	0	0	88
		3	0	0	5
		15	0	0	100
	weft	3	0	0	5
		15	0	0	86
		3	0	0	5
		15	0	0	98
		3	0	0	5
		15	0	0	71

continued

TABLE I - continued

Sample Code	Direction of test	Ignition time(sec)	Afterflame (sec)	Afterglow (sec)	Char Length (mm)
D	warp	3	0	0	15
		15	0	0	76
		3	0	0	10
		15	0	0	70
		3	0	0	10
		15	0	0	75
		3	0	0	10
		15	0	0	70
	weft	3	0	0	15
		15	0	0	67
		3	0	0	7
		15	0	0	74
		3	0	0	20
		15	0	0	75
		3	0	0	10
		15	0	0	74

TABLE II (below) shows the results of testing for flame-retardant properties according to NFG 07-184 and BS 6249.

TABLE II

Sample Code		-----BS 6249-----			
		NFP 07-184 (damaged area) cm ²	(char length) mm	Afterflame (sec)	Afterglow (sec)
A	warp	25	50	0	0
	weft	26	50	0	0
B	warp	35	82	0	0
	weft	31	62	0	0
C	warp	36	40	0	0
	weft	33	50	0	0
D	warp	29	64	0	0
	weft	24	53	0	0

The results of determination of phosphorus and nitrogen content of the fabrics after 40 washing cycles at 93°C is shown in TABLE III (below).

TABLE III

additive solid* (%)	after NH ₃ cure		as finished		after washing	
	P%	N%	P%	N%	P%	N%
A: 0 (control)	3.66	3.92	2.87	2.64	2.50	2.40
0.3	3.61	3.96	3.46	2.23	3.33	3.01
B: 0 (control)	3.69	4.08	3.15	2.97	2.82	2.60
0.3	3.68	4.29	3.63	3.37	3.24	2.89
C: 0 (control)	3.33	3.40	3.09	2.75	2.89	2.51
0.3	3.42	3.98	3.33	3.14	3.12	2.87
D: 0 (control)	3.21	3.89	2.94	2.94	2.74	2.51
0.3	3.41	4.40	3.31	3.28	3.00	2.84

* octadecylamine acetate

The water-resistance of fabrics treated according to the present invention was determined and the results are shown in TABLE IV below:

TABLE IV

Sample	Water-resistance (cm water)
Untreated fabric (control I)	4
Treatment without protonated amine (control II)	5
Treatment with protonated amine	16

The fabric used in the foregoing tests was Sample Code C (see above).

In another example, the following fabrics were treated in accordance with the present invention:

Sample Code C (As hereinbefore described).

Sample Code E A twill fabric comprising 60% cotton fibres and 40% polyester fibres and having a weight of 240 g/m².

The fabrics were impregnated with an aqueous solution containing the following percentages by weight of a precondensate of tetrakis (hydroxymethyl) phosphonium chloride and urea, together with protonated and neutralized amines in accordance with the present invention, the ratio of the phosphonium chloride to urea in the condensate being 2:1 molar:

C: 40.95% by weight

E: 37.05% by weight

The impregnated fabrics were squeezed to a wet pick-up in the following ranges based upon the original weight of the fabric:

C: 77%

E: 99%

The fabrics were then dried at 120°C to achieve a fabric moisture content of between 14-18%.

The dried fabrics were cured with gaseous ammonia in the following manners:

C1: In one step

C2: In two stages, one after the other

E1: In one step

E2: In two stages, one after the other

This was followed by oxidation with hydrogen peroxide, washing and drying.

Table V (below) shows the results of testing for flame-retardant properties according to DIN 66083 s-b:

TABLE V

Sample Code	Direction of test	Ignition time (sec)	Afterflame (sec)	Afterglow (sec)	Char length (mm)
C1	warp	3	1	0	7
		15	0	0	110
		3	1	0	9
		15	0	0	70
	weft	3	0	0	5
		15	0	0	70
		3	0	0	5
		15	0	0	75
C2	warp	3	0	0	5
		15	0	0	65
		3	1	0	5
		15	0	0	60
	weft	3	1	0	7
		15	0	0	60
		3	1	0	5
		15	0	0	55
E1	warp	3	1	0	11
		15	0	0	65
		3	2	0	11
		15	0	0	70
	weft	3	1	0	11
		15	0	0	65
		3	0	0	8
		15	0	0	75
E2	warp	3	1	0	8
		15	0	0	65
		3	0	0	7
		15	0	0	72
	weft	3	0	0	5
		15	0	0	70
		3	1	0	8
		15	0	0	85

Table VI (below) shows the results of testing for flame-retardant properties according to NFG 07-184.

TABLE VI

Sample Code	Direction of test	Damaged Area (cm ²)
C1	warp	21
	weft	23
C2	warp	21
	weft	22
E1	warp	27
	weft	25
E2	warp	24
	weft	22

The results of determinations of the phosphorus and nitrogen content of the fabrics before and after 40 washing cycles at 90°C with a detergent containing 5% perborate is shown in Table VII (below).

TABLE VII

Sample Code	After NH ₃ Cure		As finished		After washing	
	P%	N%	P%	N%	P%	N%
C1	3.53	3.92	3.47	3.23	3.28	3.10
C2	3.52	4.42	3.53	3.39	3.53	3.43
E1	4.01	4.68	3.66	3.44	3.65	3.59
E2	3.98	5.00	3.86	3.70	3.85	3.76

In yet another example the fabrics, coded C and E, were padded with the standard mixture and dried at 120°C to a fabric moisture content of between 9-12%. The fabrics were cured with gaseous ammonia in a one step manner, followed by heat curing at 130°C. The fabrics were then oxidised with hydrogen peroxide, followed by washing and drying. (Sample Codes were designated as C3 and E3 respectively).

The fabric (coded C) was also treated under the above conditions in large quantities in the plant (sample coded CM).

Table VIII shows the results of testing for flame-retardant properties according to DIN 66083.

TABLE VIII

Sample Code	Direction of test	Ignition time (sec)	Afterflame (sec)	Afterglow (sec)	Char length (mm)
C3	warp	3	0	0	5
		15	0	0	90
		3	0	0	5
		15	0	0	95
	weft	3	0	0	5
		15	0	0	75
		3	0	0	5
		15	0	0	90
CM	warp	3	1	0	5
		15	0	0	110
		3	0	0	5
		15	0	0	76
	weft	3	1	0	5
		15	0	1	50
		3	1	0	5
		15	0	1	55
E3	warp	3	0	0	5
		15	0	0	70
		3	0	0	5
		15	0	0	75
	weft	3	0	0	5
		15	0	0	70
		3	0	0	5
		15	0	0	98

Table IX (below) shows the results of testing for flame-retardant properties according to NFG 07-184.

TABLE IX

Sample Code	Direction of test	Damaged Area (cm ²)
C3	warp	28
	weft	26
CM	warp	27
	weft	25
E3	warp	27
	weft	26

The results of determination of phosphorus and nitrogen content of the fabrics after 40 washing cycles at 93°C is shown in Table X (below).

TABLE X

Sample Code	After heat Cure		As finished		After washing	
	P%	N%	P%	N%	P%	N%
C3	3.82	4.04	3.54	3.21	3.31	2.91
CM	3.53	3.57	3.24	2.88	3.07	2.69
E3	4.10	4.50	3.73	3.62	3.43	3.18

Fabrics treated according to the present invention may suitably consist essentially of cellulosic fibres, e.g. cotton fibres.

Alternatively, the fabrics may comprise both cellulosic and non-cellulosic fibres, for example polyamide fibres, acrylic fibres, aramid fibres, polyester fibres or polybenzimidazole fibres.

Suitably, the maximum content of non-cellulosic fibres in such a fabric is 70%, e.g. the fabric may comprise 60% cotton fibres and 40% polyester fibres.

A suitable weight range for the fabrics treated according to the present invention is from 0.05 to 1.0 kg/m².



GB CLAIMS

1. A method of treating fabrics to impart flame-retardant and water-resistant properties thereto, said method comprising impregnating the fabric with an aqueous solution including a poly(hydroxyalkyl) phosphonium compound, in which there is added to the impregnating solution one or more primary, secondary or tertiary aliphatic amines having from 12 to 20 carbon atoms, said amines having been protonated and neutralized prior to said addition.
2. A method according to Claim 1, in which the concentration of said protonated and neutralized amine in said solution is in the range 0.05% to 3% by weight.
3. A method according to Claim 2, in which the concentration of said amine is in the range 0.1% to 1% by weight.
4. A method according to Claim 2 or 3, in which the concentration of said amine is about 0.3% by weight.
5. A method according to any one of Claims 1 to 4, in which said protonated and neutralized amine consists essentially of n-octadecylamine.
6. A method according to any one of Claims 1 to 4, in which said protonated and neutralized amine comprises a mixture of primary aliphatic amines having from 16 to 18 carbon atoms.
7. A method according to any one of Claims 1 to 6, in which the poly(hydroxyalkyl) phosphonium compound is a tetrakis (hydroxyalkyl) phosphonium compound.



8. A method according to Claim 7, in which the tetrakis (hydroxyalkyl) phosphonium compound is a tetrakis (hydroxymethyl) phosphonium salt.
9. A method according to any one of the preceding claims, in which the amines are protonated and neutralized by means of a weak organic acid.
10. A method according to Claim 9, in which the acid is acetic acid.
11. A method according to Claim 10, in which said protonated and neutralized amine consists essentially of octadecylamine acetate.
12. A method for treating fabrics to impart flame-retardant and water-resistant properties thereto, substantially as hereinbefore described with reference to any one of the Examples.
13. A flame-retardant and water-resistant fabric treated by a method according to any one of the preceding claims.
14. A fabric according to Claim 13, said fabric consisting essentially of cellulosic fibres.
15. A fabric according to Claim 14, said fabric consisting essentially of cotton fibres,
16. A fabric according to Claim 13, said fabric comprising both cellulosic and non-cellulosic fibres.
17. A fabric according to Claim 16, in which said non-cellulosic fibres are polyamide fibres, acrylic fibres, aramid fibres, polyester fibres or polybenzimidazole fibres.
18. A fabric according to Claim 16 or 17, in which the maximum content of said non-cellulosic fibres is 70%.
19. A fabric according to Claim 16, 17 or 18, said fabric comprising 60% cotton fibres and 40% polyester fibres.