FORM 1
COMMONWEALTH OF AUSTRALIA
PATENTS ACT 1952

APPLICATION FOR A STANDARD PATENT

We, NATIONAL STARCH AND CHEMICAL CORPORATION
of 10 FINDERNE AVENUE,
BRIDGEWATER, NEW JERSEY
UNITED STATES OF AMERICA

hereby apply for the grant of a standard patent for an invention entitled:

IMPROVED WOODWORKING ADHESIVES BASED ON MULTISTAGE EMULSION POLYMERS.

which is described in the accompanying complete specification

Details of basic application(s):

Number of basic application Name of Convention country in which basic application was filed Date of basic application

343780 US 26 APR 89

My/our address for service is care of GRIFFITH HACK & CO., Patent Attorneys, 601 St. Kilda Road, Melbourne 3004, Victoria, Australia.

DATED this 17th day of April 1990

NATIONAL STARCH AND CHEMICAL CORPORATION
GRIFFITH HACK & CO.

TO: The Commissioner of Patents.
DECLARATION IN SUPPORT OF AN APPLICATION FOR A PATENT

In support of an application made by:

NATIONAL STARCH AND CHEMICAL CORPORATION

for a patent for an invention entitled: IMPROVED WOODWORKING ADHESIVES BASED ON MULTISTAGE EMULSION POLYMERS

I, Herbert J. Baumgarten, Senior Vice President
of National Starch and Chemical Corporation
10 Finderne Ave., Bridgewater, New Jersey 08807 USA

do solemnly and sincerely declare as follows:

1. I am authorised by the above mentioned applicant for the patent to make this declaration on its behalf.

2. The name and address of each actual inventor of the invention is as follows:

Paul R. Mudge, 23 Mulford Lane, Belle Mead, New Jersey 08507
George Hespe, 219 Wescott Road, Neshanic Station, New Jersey 08853

3. The facts upon which the applicant is entitled to make this application are as follows:

The applicant is the assignee of the actual inventors.

4. The basic application(s) as defined by Section 141 of the Act was (were) made as follows:

Country: U.S.A.  on 26th April 1989
in the name(s) Paul R. Mudge & George Hespe

5. The basic application(s) referred to in the preceding paragraph was (were) the first application(s) made in a Convention country in respect of the invention the subject of this application.

Declared at Bridgewater, New Jersey
this 29th day of December 1989

Signed Herbert J. Baumgarten
Senior Vice President
Title
IMPROVED WOODWORKING ADHESIVES BASED ON MULTISTAGE EMULSION POLYMERS

International Patent Classification(s)
C09J 129/04  C09J 133/12

Application No.: 53281/90
Application Date: 17.04.90

Priority Data
Number  Date  Country
343780  26.04.89  US UNITED STATES OF AMERICA

Publication Date: 08.11.90

Applicant(s)
NATIONAL STARCH AND CHEMICAL CORPORATION

Inventor(s)
PAUL R. MUDGE; GEORGE HESPE

Attorney or Agent
GRIFFITH HACK CO. MELBOURNE

Claim
1. A woodworking adhesive comprising a polyvinyl alcohol stabilized emulsion polymer, said polymer being prepared from a two stage polymerization procedure wherein the first stage comprises a polyvinyl acetate copolymer having a Tg within the range of +10 to 40°C, and a second methyl methacrylate stage copolymer having a Tg of +50 to +120°C, with the ratio of the first polymer to the second polymer varying within a range of 10:1 to 10:6.

6. A woodworking adhesive comprising a polyvinyl alcohol stabilized emulsion polymer, said polymer being prepared from a two stage polymerization procedure wherein the first stage comprises a polyvinyl acetate N-methylol acrylamide copolymer having a Tg within the range of +30 to 40°C, and a second stage methyl methacrylate - N-methylol acrylamide - triallyl cyanurate copolymer having a Tg of +100 to +120°C, with the ratio of the first polymer to the second polymer varying within a range of 10:3 to 10:4.
7. A woodworking adhesive comprising a polyvinyl alcohol stabilized emulsion polymer, said polymer being prepared from a two stage polymerization procedure wherein the first stage comprises a polyvinyl acetate copolymer having a Tg within the range of +10 to 40°C, and a second methyl methacrylate stage polymer having a Tg of +50 to +120°C, with the ratio of the first polymer to the second polymer varying within a range of 10:1 to 10:6; the emulsion polymer being formulated with at least one member selected from the class consisting of formaldehyde resins, B-stage phenolic resins and fusion aids.
TO BE COMPLETED BY APPLICANT

Name of Applicant: NATIONAL STARCH AND CHEMICAL CORPORATION

Address of Applicant: 10 FINDERNE AVENUE, BRIDGEBORI, NEW JERSEY UNITED STATES OF AMERICA

Actual Inventor:

Address for Service: GRIFFITH HACK & CO., 601 St. Kilda Road, Melbourne, Victoria 3004, Australia.

Complete Specification for the invention entitled: IMPROVED WOODWORKING ADHESIVES BASED ON MULTISTAGE EMULSION POLYMERS.

The following statement is a full description of this invention including the best method of performing it known to me:-
The present invention is directed to improved woodworking adhesives prepared from multistage emulsion polymers, the adhesives being characterized by a superior degree of water resistance.

Aqueous emulsions of vinyl acetate homo- or copolymers are widely used in formulating wood adhesives wherein bond strength, fast setting, colorless glue lines and ease of application are desired. In order to improve water resistance and other properties of these adhesives they are generally formulated with other resins such as resorcinol-formaldehyde, urea-formaldehyde, melamine-formaldehyde, phenol-formaldehyde resins and trimethylol phenol oligomer or with fusion aids, such as tetrahydro-furfuryl alcohol or ethylene glycol diacetate. These pre-formulated adhesives may then be catalyzed with various acidic metal salts as is known in the art. These adhesive systems are described, for example, in U.S. Patents No. 3,301,809; 3,563,851; 3,433,701; 3,041,301; 4,474,915; and 4,687,809.

While such formulated vinyl acetate based adhesives provide adequate water resistance for plywood and finger jointed lumber constructions tested at room temperature, under vacuum-pressure soaks and in boiling water, none of the current vinyl acetate adhesives meet ASTM standards for exterior lumber, especially for ASTM D-3110 for edge glued lumber. This is illustrated by the requirements for these tests on pine lumber, as well as the values obtained using conventional formulated crosslinked polyvinyl acetate adhesives (PVA) presented in Table I.
We have now found that superior water resistant emulsions for use in woodworking adhesives may be prepared by a two-stage polymerization procedure wherein a first stage is prepared from a relatively low Tg vinyl acetate polymer and a second stage from a higher Tg methyl methacrylate polymer thereby combining the advantageous wetting or rheological as well as film forming properties of the polyvinyl alcohol stabilized vinyl acetate copolymer with the reduced thermoplasticity and high temperature and water resistance of the higher Tg methyl methacrylate copolymer. The resultant formulated adhesives pass not only the finger jointed lumber requirements but also the ASTM D-3110 requirements for edge glued lumber.

The two stage polymerization utilized herein may be carried out using a variety of specific modifications which are generally referred to as producing "core-shell" or "interpenetrating network" type polymers. Such polymerization procedures are described, for example, in U.S. Pat. Nos. 3,671,610; 3,833,404; and 4,616,057, the disclosures of which are incorporated herein by reference.

More specifically, a vinyl acetate polymer is prepared using conventional batch, semi-batch or continuous emulsion polymerization procedures such as taught in U.S. Pat. Nos. 2,754,280; 2,795,564 and
3,732,184 chosen so as to provide a first-stage polymer having a Tg of 0 to +40°C, preferably 30 to 40°C. While vinyl acetate will comprise the predominant portion of the polymer, up to about 25% of other copolymerizable comonomers may be employed in specific amounts depending on the particular comonomer so as to maintain the required Tg range. Suitable comonomers include, for example, other vinyl esters (e.g., vinyl propionate), vinyl ethers, fumarates, maleates, ethylene, acrylates, etc.

The methacrylate ester which comprises the major portion of the second stage may be copolymerized with minor amounts of acrylate, other methacrylate, (meth)acrylonitrile or styrene comonomers selected to have a Tg within the range of +50 to +125°C, preferably about 80 to 120°C.

Suitable copolymers within this Tg range may be prepared, for example, from copolymers of C₁⁻C₄ acrylates or C₂⁻C₄ methacrylates with methyl methacrylate or other higher Tg methacrylates.

In addition to the vinyl acetate and higher Tg monomers, a pre-crosslinking monomer and/or a post-crosslinking monomer may optionally be present in either or both stages of the polymerization depending upon the specific performance required. If used, it is preferred that the crosslinking monomer be added in the second stage polymerization.

Suitable pre-crosslinking agents include alkylene glycol diacylates and methacrylates, such as ethylene glycol diacylate, 1,3-butylen glycol diacylate, propylene glycol diacylate, triethylene glycol dimethacrylate etc., 1,3-glycerol dimethacrylate, 1,1,1-trimethylol propane dimethacrylate, 1,1,1-trimethylol ethane diacylate, pentaerythritol trimethacrylate, sorbitol pentamethacrylate, methylenebisacrylamide, methylene bismethacrylamide, divinyl benzene, vinyl methacrylate, vinyl crotonate, vinyl acrylate, divinyl adipate; also di- and tri-allyl compounds, such as
triallyl cyanurate, triallyl isocyanurate, diallyl phthalate, allyl methacrylate, allyl acrylate, diallyl maleate, diallyl fumarate, diallyl itaconate, diallyl malonate, diallyl carbonate, triallyl citrate, triallyl aconitate; also divinyl ether, ethylene glycol divinyl ether and the like.

The amount of active crosslinking agent in each stage of the polymer emulsion of the present invention may vary from about 0.01 to 0.5 percent, preferably from about 0.05 to 0.25 percent by weight of the polymer.

Examples of post-crosslinking agents are: N-alkyldlamides of alpha, beta ethylenically unsaturated carboxylic acids having 3-10 carbons, such as N-methylol acrylamide, N-ethanol acrylamide, N-propanol acrylamide, N-methylol methacrylamide, N-ethanol methacrylamide, N-methylol maleamide, N-methylol maleimide, N-methylol maleamic acid, N-methylol maleamic acid esters; the N-alkyldlamides of the vinyl aromatic acids, such as N-methylol-p-vinylbenzamide and the like; also N-(alkoxymethyl) acrylates and methacrylates, where the alkyl group has from 1-8 carbon atoms, such as N-(methoxymethyl) acrylamide, N-(butoxymethyl) acrylamide, N-(methoxymethyl) methacrylamide, N-(butoxymethyl) allyl carbamate and N-(methoxymethyl) allyl carbamate, and mixtures of these monomers with allyl carbamate, acrylamide or methacrylamide. If present, these monomers are used in amounts of 0.1 to 6% by weight.

Olefinically unsaturated acids may also be employed in either stage of the polymerization to improve adhesion and to contribute additional heat resistance. These acids include the alkenoic acids having from 3 to 6 carbon atoms, such as acrylic acid, methacrylic acid, crotonic acid; alkenedioic acids, e.g., itaconic acid, maleic acid or fumaric acid or mixtures thereof in amounts sufficient to provide up to about 4 parts, preferably 0.5 to 2.5 parts, per 100 parts of major monomer.
In addition, certain copolymerizable monomers which assist in the stability of the copolymer emulsion, e.g., vinyl sulfonic acid and 2-acrylamido-2-methylpropene sulfonic acid are used herein as latex stabilizers. These stabilizers are added in amounts of from about 0.2 to 3% by weight of the monomer mixture.

Conventional batch, semi-batch or continuous emulsion polymerization procedures may be utilized herein.

Suitable as polymerization catalysts are the water-soluble free-radical-formers generally used in emulsion polymerization, such as hydrogen peroxide, sodium persulfate, potassium persulfate and ammonium persulfate, as well as tert-butyl hydroperoxide, in amounts of between 0.01 and 3% by weight, preferably 0.01 and 1% by weight based on the total amount of the emulsion. They can be used alone or together with reducing agents such as sodium formaldehyde-sulfoxylate, ferrous salts, sodium dithionite, sodium hydrogen sulfite, sodium sulfite, sodium thiosulfate, as redox catalysts in amounts of 0.01 to 3% by weight, preferably 0.01 to 1% by weight, based on the total amount of the emulsion.

The free-radical-formers can be charged with the protective colloid or be added during the polymerization in doses.

The polymerization is carried out at a pH of between 2 and 7, preferably between 3 and 5. In order to maintain the pH range, it may be useful to work in the presence of customary buffer systems, for example, in the presence of alkali metal acetates, alkali metal carbonates, alkali metal phosphates. Polymerization regulators, like mercaptans, aldehydes, chloroform, ethylene chloride and trichloroethylene, can also be added in some cases.
The polymerization is carried out using a protective colloid as the stabilizing agent. Generally, the colloid chosen is a standard grade of polyvinyl alcohol typically employed in such polymerizations. In general, this colloid is used at levels of 0.05% to 4% by weight based on the total emulsion. The polyvinyl alcohol may be added, in its entirety, to the initial charge to the polymerization zone or a portion of the emulsifier, e.g. from 90 to 25 percent thereof, can be added continuously or intermittently during polymerization.

The polymerization reaction is generally continued until the residual vinyl acetate monomer content is below 1%. The completed reaction product is then allowed to cool to about room temperature, while sealed from the atmosphere.

To the above described vinyl acetate copolymer is added a second monomeric system comprising the methyl methacrylate monomer together with any optional pre- and post-crosslinking agents which may be utilized. The pre- and post-crosslinking agents optionally used in the second stage polymerization are the same classes of monomers described previously. For convenience of cure, it may be desirable to use the same crosslinking agents in both stages of the polymerization, however, it is not required and different monomers may be employed.

The ratio of the first stage polymer to the second stage polymer may vary from about 10:1 to 10:6 and is preferably in the range of 10:3 to 10:4.

All of the second monomeric mixtures may be added directly to the first polymer emulsion and the second polymerization begun or the second monomeric mixture may be added gradually as the polymerization reaction proceeds.
The latices are produced and used at relatively high solids contents, e.g. up to about 60%, although they may be diluted with water if desired. The preferred latices will contain about from 45 to 55, and, most preferred about 50% weight percent solids.

The emulsions may be used in any conventional woodworking adhesive compositions and formulated with such additives as are commonly used in the production of these adhesives. Such additives include formaldehyde resins such as resorcinol formaldehyde, urea formaldehyde, melamine formaldehyde and phenol formaldehyde; as well as B-stage phenolic resins, such as trimethylol phenol oligomer, prepared by means of any conventional phenolaldehyde condensation reaction which may be added in an amount of 1 to 20%, by weight, based on total emulsion, preferably 3 to 10%, by weight.

The adhesives may also be formulated with 1 to 7%, by weight, polymer solids of a fusion aid such as tetrahydrofurfuryl alcohol, ethylene glycol diacetate, and ethylene glycol monoethyl ether acetate as taught in U.S. Patent No. 4,474,915, the disclosure of which is incorporated herein by reference.

The addition of acidic, metal salt curing agents may also be desired in order to accelerate the cure of the formulated adhesive. The preferred curing agents for use herein comprise acidic, metal salts selected from the group consisting of chromic nitrate, chronic perchlorate, aluminum nitrate, aluminum chloride, zinc nitrate, and para-toluene sulfonic acid. The proportion of acidic, metal salt curing agent which is added will depend upon the rate of cure which is desired in the final product but a practical range has been found to be from about 0.003 to 0.12 gram equivalents of curing agent (anhydrous basis) for each 100 grams of the total weight of the blend of the adhesive composition.
If fusion aids are used in the formulation, it may also be desirable to add cross-linking inhibitors such as are described in U.S. Pat. No. 4,085,074 issued Apr. 18, 1978 to M.C. Woo and re-issued as U.S. Pat. Re. No. 30,576. Typical inhibitors include ammonia, alkyl amines (e.g. triethyl amine), and alkylol amines (e.g. triethanol amine and N-methylethanol amine). When used, they are added ordinarily in amounts such that the mole ratio of the inhibitor to curing agent ranges from 0.1 to 1.7. They may be added to the vessel at the end of the polymerization reaction or they may be post added prior to or with the curing agent. Their addition improves the storage stability of the adhesives.

In preparing the adhesive compositions of this invention, it is merely necessary first to combine, by mixing, the aqueous emulsion poly. with the additive(s).

The following examples are given to illustrate the present invention, but it will be understood that they are intended to be illustrative only and not limitative of the invention. In the examples, all parts are by weight and all temperatures in degrees Celsius unless otherwise noted.

**EXAMPLES 1 – 18**

**Stage 1**

The following ingredients were charged to a four neck two liter flask equipped with a mechanical stirrer, condenser, addition funnel, thermometer and N₂ gas inlet.

<table>
<thead>
<tr>
<th>Ingredients</th>
<th>Amounts (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Water</td>
<td>345</td>
</tr>
<tr>
<td>Ammonium persulfate</td>
<td>0.45</td>
</tr>
<tr>
<td>Ammonium hydroxide (28% ol)</td>
<td>0.05</td>
</tr>
<tr>
<td>Vinol 523 (88% PVAc-medium molecular weight)</td>
<td>16.5</td>
</tr>
<tr>
<td>Vinyl acetate</td>
<td>10</td>
</tr>
</tbody>
</table>
The temperature was raised under a nitrogen atmosphere to 72°C.

After 5-10 minutes at 72°C the following ingredients were added as a pre-emulsified monomer mixture over 3 hrs at 72-75°C.

<table>
<thead>
<tr>
<th>Ingredients</th>
<th>Amounts (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vinyl acetate</td>
<td>450</td>
</tr>
<tr>
<td>N-methylol acrylamide (48% sol)</td>
<td>31</td>
</tr>
<tr>
<td>Vinol 523</td>
<td>0.5</td>
</tr>
<tr>
<td>Water</td>
<td>142.5</td>
</tr>
</tbody>
</table>

At the end of the monomer slow addition the temperature was allowed to exotherm to 90°C, held for 20 minutes then cooled to 60°C for stage 2.

**Stage 2 - Slow Addition Process used in Examples 1-17**

The emulsion from stage 1 was cooled to 60°C and a mixture of 15g Triton X305 surfactant in 170g water was added with stirring. Then a monomer mixture of 150g methyl methacrylate, 4.5g of isobutoxy methacrylamide and 0.75 triallyl cyanurate was added over a period of 90 minutes in a uniform addition mode. Concurrent with the monomer addition 1.5g of tert-butylhydroperoxide in 25g of water and 1g of sodium formaldehyde sulfoxalate in 25g of water were added over 120 minutes. At the finish of these additions the emulsion was cooled to room temperature.

**Stage 3 - Batch Equilibration Process used in Example 18**

The stage 1 emulsion was cooled to 60°C and then a mixture of 15g Triton X305 surfactant in 170g of water was added with stirring. Secondly, a monomer mixture of 150g methyl methacrylate and 0.75g diallyl maleate was added over 15 minutes with continual agitation followed by an equilibration period of 60 minutes at 60°C. After the equilibration period, the addition of 1.5g tert-butylhydroperoxide in 25g water and 1g
sodium formaldehyde sulfoxalate in 25g water was started and completed over 60 minutes. Once the initiator additions were finished the emulsion was cooled to room temperature.

The resulting emulsions were then tested using the following procedures:

Test Procedures

In all cases the polymer emulsions were catalysed with 5% of a 32° Be aluminum chloride solution. The construction was edge glued pine bonded in a radio frequency (RF) press. After aging for 7 days, the construction was cut into individual specimens, and put through one of the following durability cycles as required by ASTM D-3110.

Room Temperature - Specimens were broken in compression shear with no further aging.

Vacuum-Pressure Soaks - Specimens were submerged in water. A 15 MM vacuum was drawn for 30 minutes, followed by 75 psi hydrostatic water pressure for 30 minutes. Specimens were broken in compression shear while still wet.

Boils - Specimens were submerged in boiling water for 4 hours, then oven dried for 20 hours, then boiling water for another 4 hours. They were then cooled to room temperature in cold water and broken while wet.

The results of the testing, shown in Tables 3 and 4, indicate the superior results achieved using the adhesives of the invention. Thus, the results presented in Table 4 clearly show that the adhesives described herein pass the commercial requirements defined in ASTM D-3110 reproduced in Table 1 above. Thus, in addition to readily passing the finger jointed lumber standards, as do the currently utilized polyvinyl acetate adhesives, these adhesives also meet the far more stringent edge-glued requirements.
Further, the results presented in Examples 1, 2 and 3 of Table 3 show improvements achieved using various levels of MMA/IBMA/TAC in the second stage with 30 parts MMA on 100 parts of first stage being optimum for use on pine.

Example 2 shows that NMA in first stage is more beneficial than the use of IBMA only in the second stage as in Example 7.

Examples 4, 5, 6 show the effect of lower Tg second stage (1/1 ratio of EA/MMA) with only 50 parts second stage showing improvement over the control crosslinked polyvinyl acetate. The results indicate that lower levels of higher Tg second stage are more effective than higher levels of lower Tg second stage (MMA = +100°C Tg; 50/50 EA/MMA = +60 Tg; VA = +30 Tg)

Examples 8 and 9 also illustrate the effect of the lower Tg ethyl acrylate (EA) monomer in the first stage which is less effective in that it detracts from the overall performance even with high levels of second stage reinforcement.

Examples 10, 11, 12 show effect of post crosslinking in the second stage which, although not essential for strength (PSI at break), does aid in wood failure with NMA and IBMA being similar. Examples 13 and 14 show that pre-crosslinking in either the first or in the second stage also is not necessary for these improvements.

Table 4 shows the edge glued testing results for formulated adhesives. In this table the addition to the control of 6% trimethylol phenol oligomer (TMP) which is a plasticizer and post crosslinker helps in boiling water resistance (from 200 to 300 psi), however as shown in Example 2, the TMP effect is much larger giving data which passed the D3110 specification for edge glued lumber. The control emulsion plus TMP does not reach performance of two-stage without TMP.
In contrast, Example 17 shows that the simplest polymer system with methyl methacrylate homopolymer in the second stage, when formulated with TMP, passes the D3110 specification for edge glued lumber unlike the control emulsion. Furthermore, improved results would be achieved utilizing the two stage polymerization procedure without crosslinking monomer in either stage over similar standard adhesives prepared using a one stage polymerization.
<table>
<thead>
<tr>
<th>Example</th>
<th>First Stage of Monomers</th>
<th>Major Monomers</th>
<th>Pre and Post Xlinking Monomers</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>% of First Stage</td>
<td>% on Second Stage Total</td>
</tr>
<tr>
<td></td>
<td>ViAc</td>
<td>NMA</td>
<td>EA</td>
</tr>
<tr>
<td>5</td>
<td>Control</td>
<td>100</td>
<td>3.4</td>
</tr>
<tr>
<td>10</td>
<td>5</td>
<td>100</td>
<td>3.4</td>
</tr>
<tr>
<td>6</td>
<td>15</td>
<td>100</td>
<td>3.4</td>
</tr>
<tr>
<td>7</td>
<td>8</td>
<td>100</td>
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</tr>
<tr>
<td>8</td>
<td>9</td>
<td>80</td>
<td>3.4</td>
</tr>
<tr>
<td>15</td>
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<td>100</td>
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<td>3.4</td>
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<td>3.4</td>
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<tr>
<td>15</td>
<td>16</td>
<td>100</td>
<td>3.4</td>
</tr>
<tr>
<td>17</td>
<td>18</td>
<td>100</td>
<td>3.4</td>
</tr>
</tbody>
</table>

Control = commercially utilized PVA stabilized NMA-crosslinked polyvinyl acetate

ViAc = Vinyl acetate       TAC = Triallyl cyanurate
NMA = N-methylol acrylamide DAM = Diallyl maleate
EA = Ethyl acrylate        n-IBMA = N-(Isobutoxymethyl) acrylamide
MMA = Methyl methacrylate
### TABLE 3

UNFORMULATED ADHESIVES - ASTM D3110 EXTERIOR EDGE GLUED PINE CONSTRUCTIONS

<table>
<thead>
<tr>
<th>Example</th>
<th>Room Temperature</th>
<th>Pressure Soaks</th>
<th>Boils</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>PSI</td>
<td>%WF</td>
<td>PSI</td>
</tr>
<tr>
<td>5 Pine Wood Control</td>
<td>1232</td>
<td>100</td>
<td>-</td>
</tr>
<tr>
<td>1</td>
<td>1411</td>
<td>97</td>
<td>483</td>
</tr>
<tr>
<td>2</td>
<td>1376</td>
<td>100</td>
<td>590</td>
</tr>
<tr>
<td>3</td>
<td>1404</td>
<td>96</td>
<td>599</td>
</tr>
<tr>
<td>4</td>
<td>1427</td>
<td>94</td>
<td>576</td>
</tr>
<tr>
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<td>485</td>
</tr>
<tr>
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<td>1252</td>
<td>99</td>
<td>485</td>
</tr>
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<td>92</td>
<td>611</td>
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<td>15</td>
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<td>547</td>
</tr>
<tr>
<td>18</td>
<td>1252</td>
<td>97</td>
<td>571</td>
</tr>
</tbody>
</table>

Control = commercially utilized PVA stabilized NMA-crosslinked polyvinyl acetate
**TABLE 4**

FORMULATED ADHESIVES - ASTM D3110 EXTERIOR EDGE GLUED PINE CONSTRUCTIONS

<table>
<thead>
<tr>
<th>Example</th>
<th>Formulation</th>
<th>Room Temp.</th>
<th>Pressure Soaks</th>
<th>Boils</th>
</tr>
</thead>
<tbody>
<tr>
<td>5 Pinewood Control (A)</td>
<td>-</td>
<td>1232 100</td>
<td>-</td>
<td>508 100</td>
</tr>
<tr>
<td>10 Control</td>
<td>-</td>
<td>1411 97</td>
<td>483 22</td>
<td>205 12</td>
</tr>
<tr>
<td>2</td>
<td>+6% TMP</td>
<td>1478 96</td>
<td>479 21</td>
<td>303 27</td>
</tr>
<tr>
<td>2</td>
<td>-</td>
<td>1404 96</td>
<td>509 77</td>
<td>408 49</td>
</tr>
<tr>
<td>10</td>
<td>+3% TMP</td>
<td>1271 100</td>
<td>585 60</td>
<td>509 79</td>
</tr>
<tr>
<td>2</td>
<td>+6% TMP</td>
<td>1382 100</td>
<td>595 66</td>
<td>513 85</td>
</tr>
<tr>
<td>2</td>
<td>+3% THFA</td>
<td>1572 97</td>
<td>616 57</td>
<td>404 24</td>
</tr>
<tr>
<td>2</td>
<td>+6% THFA</td>
<td>1327 98</td>
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<td>+3% EGDA</td>
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<td>1513 94</td>
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<td>17</td>
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<td>1464 94</td>
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TMP = Trimethylolphenolic oligomer
THFA = Tetrahydrofurfuryl alcohol
EGDA = Ethylene Glycol Diacetate

Control = commercially utilized PVA stabilized NMA-crosslinked polyvinyl acetate
THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A woodworking adhesive comprising a polyvinyl alcohol stabilized emulsion polymer, said polymer being prepared from a two stage polymerization procedure wherein the first stage comprises a polyvinyl acetate copolymer having a Tg within the range of +10 to 40°C, and a second methyl methacrylate stage copolymer having a Tg of +50 to +120°C, with the ratio of the first polymer to the second polymer varying within a range of 10:1 to 10:6.

2. The adhesive of Claim 1, wherein the stage two polymer has a Tg of 80 to 120°C, and wherein the ratio of the first stage polymer to second stage polymer is in the range of 10:3 to 10:4.

3. The adhesive of Claim 1, wherein the polyvinyl alcohol is present in an amount of 0.05 to 4% by weight of the total emulsion.

4. The adhesive of Claim 1, wherein there is additionally present in either or both of stage one or two a pre- or post-crosslinking monomer.

5. The adhesive of Claim 1 wherein there is additionally present in the emulsion polymer up to 4 parts by weight of an alkenoic or alkenedioic acid having from 3 to 6 carbon atoms per 100 parts of the monomers of the emulsion polymer.
alkenedioic acids, e.g., itaconic acid, maleic acid or fumaric acid or mixtures thereof in amounts sufficient to provide up to about 4 parts, preferably 0.5 to 2.5 parts, per 100 parts of major monomer.

6. A woodworking adhesive comprising a polyvinyl alcohol stabilized emulsion polymer, said polymer being prepared from a two stage polymerization procedure wherein the first stage comprises a polyvinyl acetate N-methylol acrylamide copolymer having a Tg within the range of +30 to 40°C, and a second stage methyl methacrylate - N-methylol acrylamide - triallyl cyanurate copolymer having a Tg of +100 to +120°C, with the ratio of the first polymer to the second polymer varying within a range of 10:3 to 10:4.

7. A woodworking adhesive comprising a polyvinyl alcohol stabilized emulsion polymer, said polymer being prepared from a two stage polymerization procedure wherein the first stage comprises a polyvinyl acetate copolymer having a Tg within the range of +10 to 40°C, and a second methyl methacrylate stage polymer having a Tg of +50 to +120°C, with the ratio of the first polymer to the second polymer varying within a range of 10:1 to 10:6; the emulsion polymer being formulated with at least one member selected from the class consisting of formaldehyde resins, B-stage phenolic resins and fusion aids.

8. The woodworking adhesive of Claim 7, wherein there is also present an acid metal salt curing agent in an amount of from 0.003 to 0.12 gram equivalents per 100 grams of the adhesive.