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(71) Demandeur/Applicant:
DENTSPLY INTERNATIONAL, INC., US

(72) Inventeurs/Inventors:
KLEE, JOACHIM E., DE;
WALZ, UWE, DE

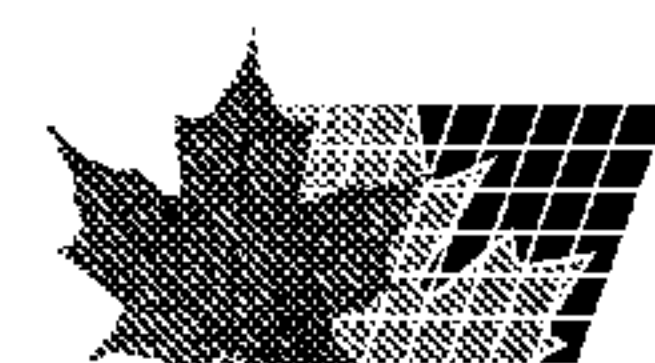
(74) Agent: SMART & BIGGAR

(54) Titre : ADHESIF DENTAIRE SIMPLE A AUTO-AMORCAGE

(54) Title: ONE-PART SELF-PRIMING DENTAL ADHESIVE

(57) Abrégé/Abstract:

A dental adhesive composition for bonding dental restoratives to dentin and enamel provides a one-part self-etching, self-priming dental adhesive composition having hydrolysis stable polymerizable acidic adhesive monomers.



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WALZ, Uwe; Zum Klausenhorn 9, 78465 Konstanz (DE).
- (21) International Application Number: PCT/US02/25005 (74) **Agents:** **HURA, Douglas, J.** et al.; Dentsply International
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WO 03/013444 A1

(54) **Title:** ONE-PART SELF-PRIMING DENTAL ADHESIVE

(57) **Abstract:** A dental adhesive composition for bonding dental restoratives to dentin and enamel provides a one-part self-etching, self-priming dental adhesive composition having hydrolysis stable polymerizable acidic adhesive monomers.

ONE-PART SELF-PRIMING DENTAL ADHESIVE

Technical Field

The invention relates to dental adhesive compositions for bonding dental restoratives to dentin and enamel. More specifically the invention provides a one-part self-etching, self-priming dental adhesive composition comprising hydrolysis stable polymerizable acidic adhesive monomers.

Background of the Invention

Presently, self-etching, self-priming dental adhesives are composed of two-part systems due to stability issues of the polymerizable acidic monomers. The stability issues are due to the hydrolysis of acidic and adhesive monomers in water or water/solvent mixtures. Therefore the acidic and adhesive monomers are stored water-free and mixed with the aqueous part just before application.

Frequently, sulfuric acid ester or phosphorous ester groups are employed in acidic polymerizable adhesive monomers. These acidic groups tend to hydrolyze ester moieties within of the monomers. To overcome these disadvantages polymerizable phosphonic esters were proposed (DE 19918974). However, these monomers still comprise hydrolysable (meth)acrylic ester moieties. Recently, hydrolysis stable monomers with phosphonic acid ester groups based on α -(oxo ethyl) acrylate were claimed (DE 19746708). However, the synthesis of these monomers is rather expensive and cost prohibitive for the envisaged applications.

Two-part self-etching, self-priming dental adhesive systems are either applied sequentially or in one step after mixing the two parts. Both procedures have inherent disadvantages due to clinical complications which might occur inbetween sequential steps (saliva or blood contamination) or due to dosing problems when mixing is required prior to the application of the self-etching adhesive.

In order to overcome these clinical problems it would be advantageous to provide the self-etching adhesive as a one-part system eliminating the need of sequential application or premixing.

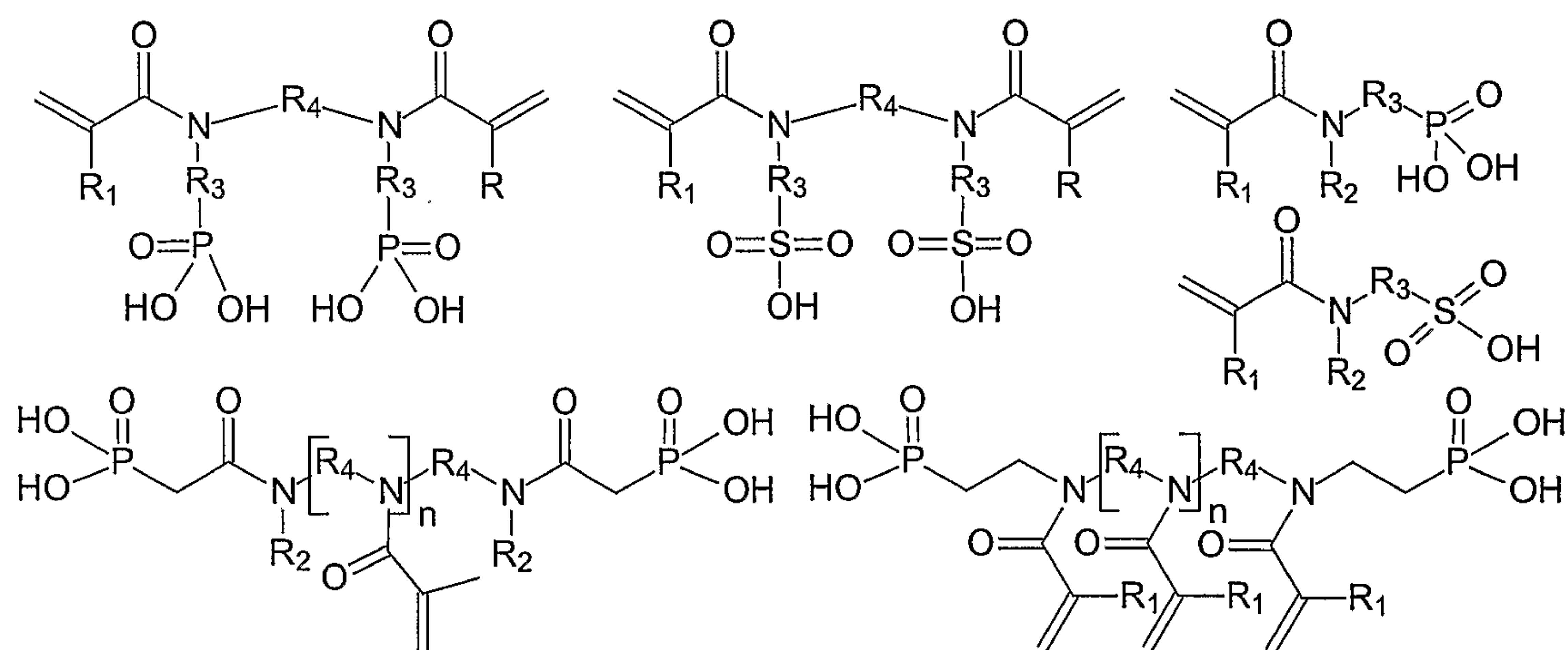
Description of the preferred embodiments

The present invention provides a hydrolysis stable one-part self-etching, self-priming dental adhesive based on (meth)acrylamides and use thereof in polymerizable dental adhesive compositions containing

- i) a polymerizable (meth) acrylamide that comprises at least an organic or inorganic acidic moiety
- ii) a polymerizable monomer
- iii) polymerization initiator, inhibitor and stabilizer.

Preferably the hydrolysis stable one-part self-etching, self-priming dental adhesive comprises at least a carboxylic acid, a phosphoric acid or a sulfuric acid group or most preferably at least a phosphonic or a sulfonic acid moiety.

The polymerizable (meth) acrylamide that comprises at least a phosphonic or sulfonic acid moiety is characterized by the following formulas:



wherein

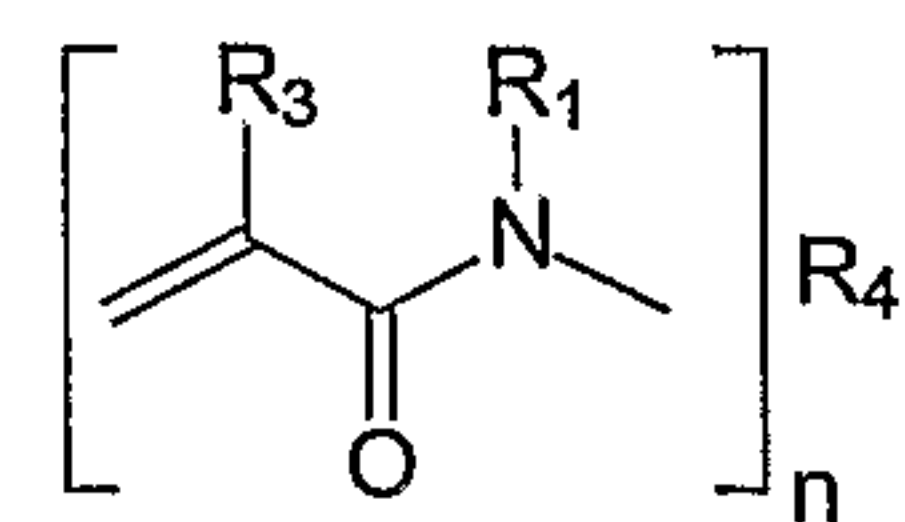
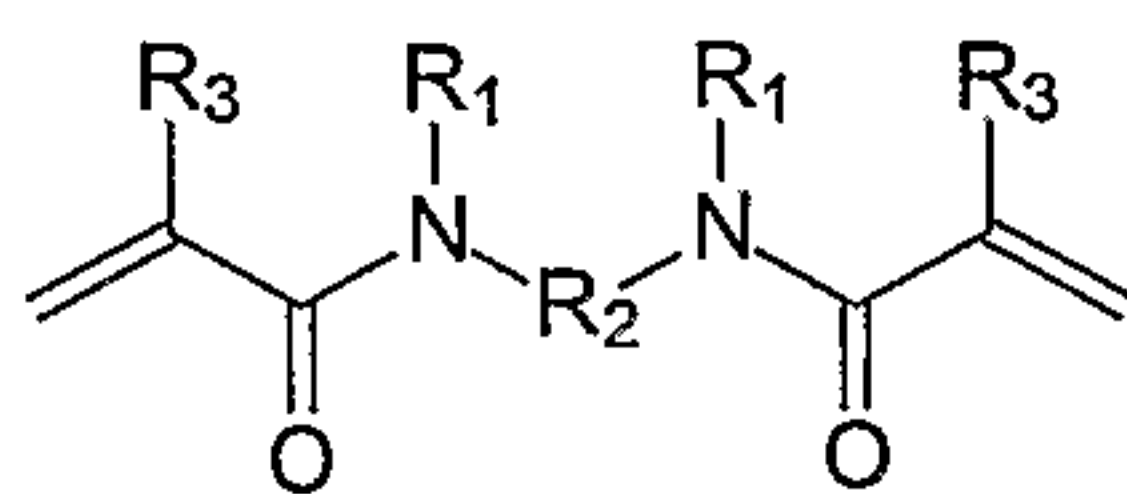
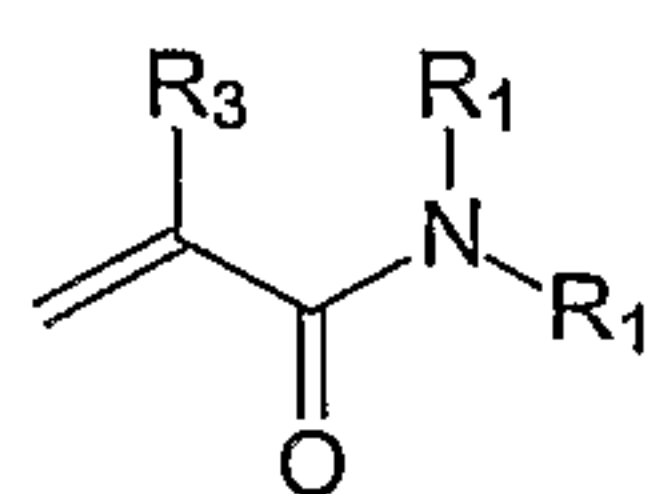
R_1 and R_2 independently are Hydrogen or a substituted or unsubstituted C_1 to C_{18} alkylene, substituted or unsubstituted cycloalkylene, substituted or unsubstituted C_5 to C_{18} arylene or heteroarylene, substituted or

unsubstituted C₅ to C₁₈ alkylarylene or alkylheteroarylene, substituted or unsubstituted C₇ to C₃₀ alkylene arylene,

R₃ and R₂ independently are a difunctional substituted or unsubstituted C₁ to C₁₈ alkylene, difunctional substituted or unsubstituted cycloalkylene, difunctional substituted or unsubstituted C₅ to C₁₈ arylene or heteroarylene, difunctional substituted or unsubstituted C₅ to C₁₈ alkylarylene or alkylheteroarylene, difunctional substituted or unsubstituted C₇ to C₃₀ alkylene arylene,

n is an integers.

In addition to the polymerizable (meth) acrylamide that comprises at least a phosphonic or sulfonic acid moiety polymerizable monomers are applied that also have an improved hydrolysis stability, that are characterized by the following formulas:



wherein

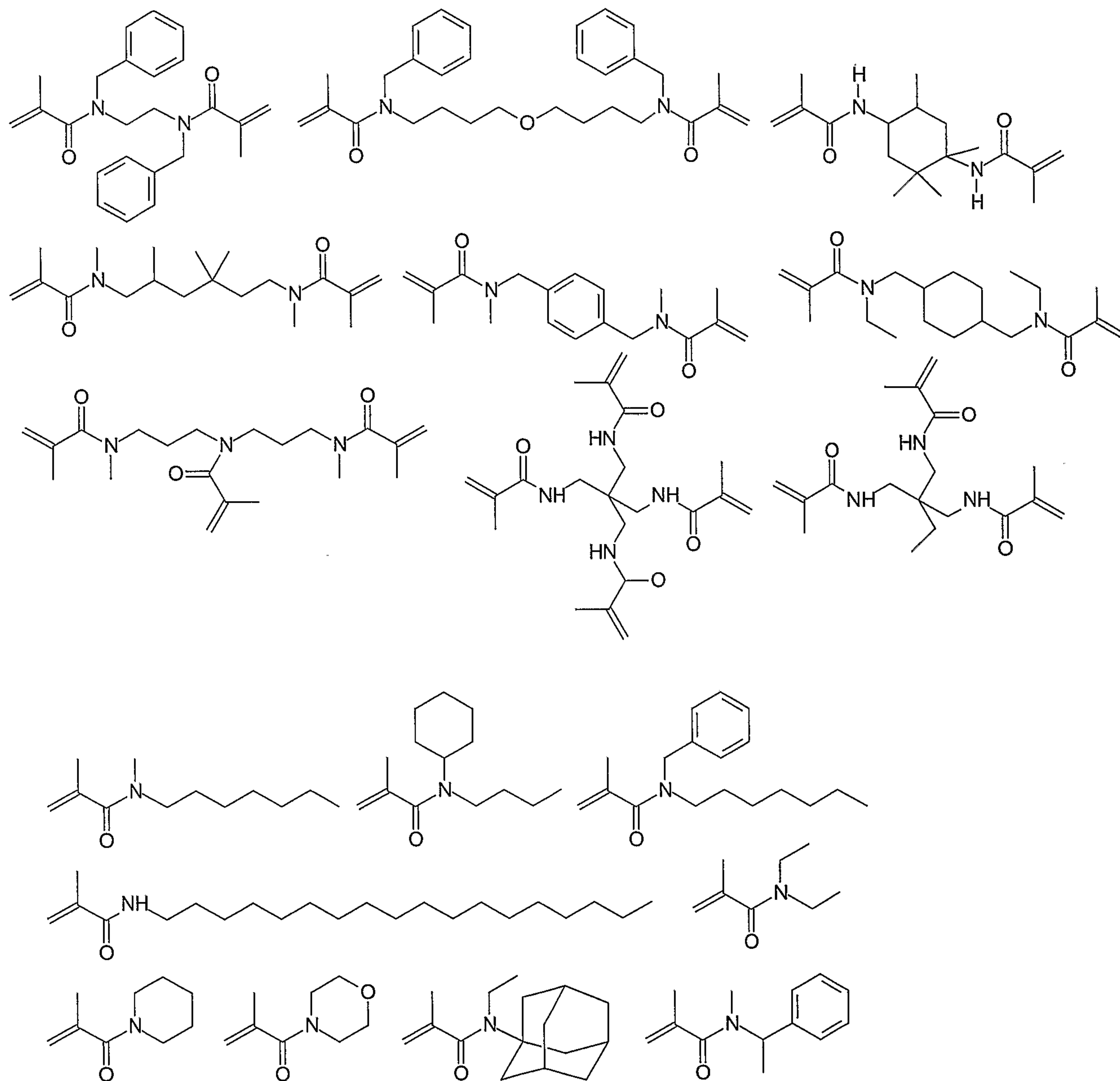
R₁ and R₃ independently are H or a substituted or unsubstituted C₁ to C₁₈ alkylene, substituted or unsubstituted cycloalkylene, substituted or unsubstituted C₅ to C₁₈ arylene or heteroarylene, substituted or unsubstituted C₅ to C₁₈ alkylarylene or alkylheteroarylene, substituted or unsubstituted C₇ to C₃₀ alkylene arylene,

R₂ is a difunctional substituted or unsubstituted C₁ to C₁₈ alkylene, difunctional substituted or unsubstituted cycloalkylene, difunctional substituted or unsubstituted C₅ to C₁₈ arylene or heteroarylene, difunctional substituted or unsubstituted C₅ to C₁₈ alkylarylene or alkylheteroarylene, difunctional substituted or unsubstituted C₇ to C₃₀ alkylene arylene,

R_4 is a mono- or polyfunctional substituted or unsubstituted C_1 to C_{18} alkylene, mono- or polyfunctional substituted or unsubstituted cycloalkylene, mono- or polyfunctional substituted or unsubstituted C_5 to C_{18} arylene or heteroarylene, mono- or polyfunctional substituted or unsubstituted C_5 to C_{18} alkylarylene or alkylheteroarylene, mono- or polyfunctional substituted or unsubstituted C_7 to C_{30} alkylene arylene,

n is an integers.

The preferably used bis- and mono (meth) acrylamides are characterized by the following formulas:



The claimed compositions comprise at least a bis- or poly (meth) acrylamide, a polymerizable mono acrylamide, an initiator, a stabilizer, water and/or an organic solvent.

The polymerization initiator is a thermal initiator, a redox-initiator or a photo initiator preferably used is champhor quinone.

To stabilize the dental composition as stabilizer are applied radical absorbing monomers such as hydroquinone monomethylether, 2,6-di-tert.-butyl-p-cresol, tetramethyl piperidine N-oxyl radical, galvanoxyl radical.

Example 1

N,N'-Bis (diethyl ethyl phosphonate)-1.2-Bis (2-aminoethoxy) ethane:

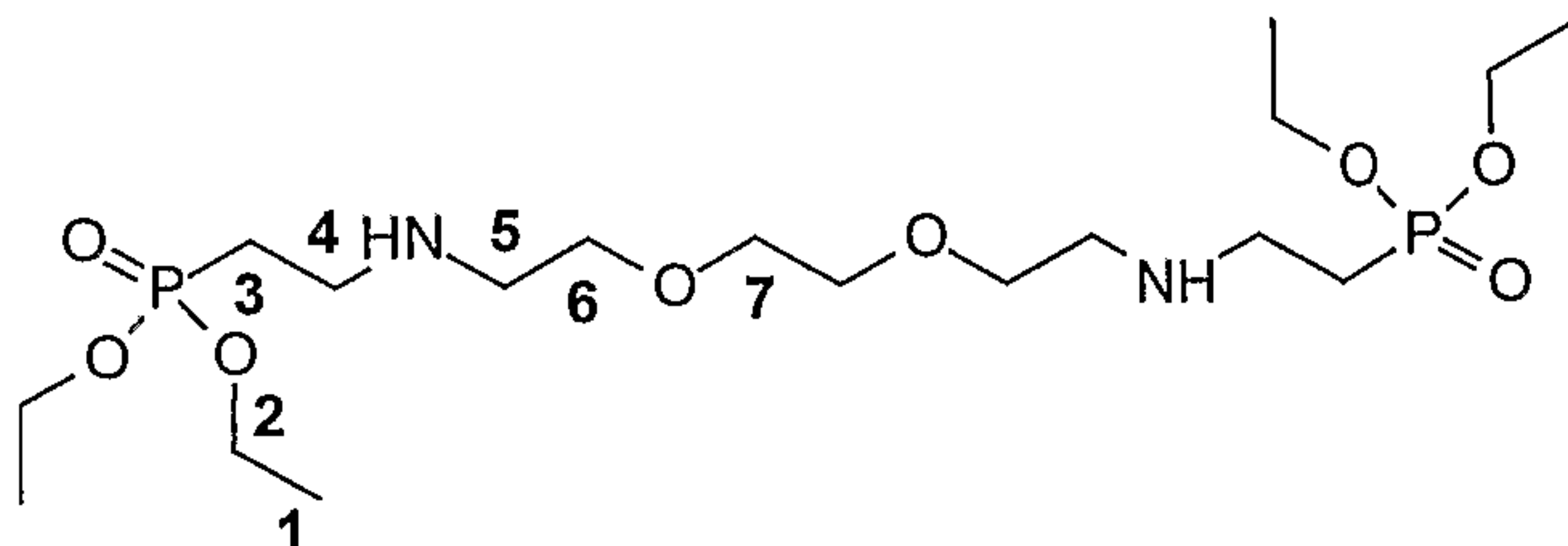
To 11.287 g (0.076 mol) 1.2-Bis (2-aminoethoxy) ethane were added 25.000 g (0.152 mol) Diethyl vinylphosphonate and stirred for 6 hours at 23 °C.

Yield: 36.287 g (100 % d. Th)

(C₁₈H₄₂O₈N₂P₂), 476.49

IR: 3411, 3390 (OH), 2973, 2929, 2885 (CH₂/CH₃), 1390 (CH₂/CH₃), 1078 cm⁻¹ (OH).

¹³C-NMR: 72.1 (6), 70.6 (7), 62.3 (2), 49.9 (5), 34.2 (4), 27.8 (3), 16.4 (1)



N,N'-Bis (diethyl ethyl phosphonate)-N,N'-bismethacrylamido-1.2-Bis (2-aminoethoxy) ethane:

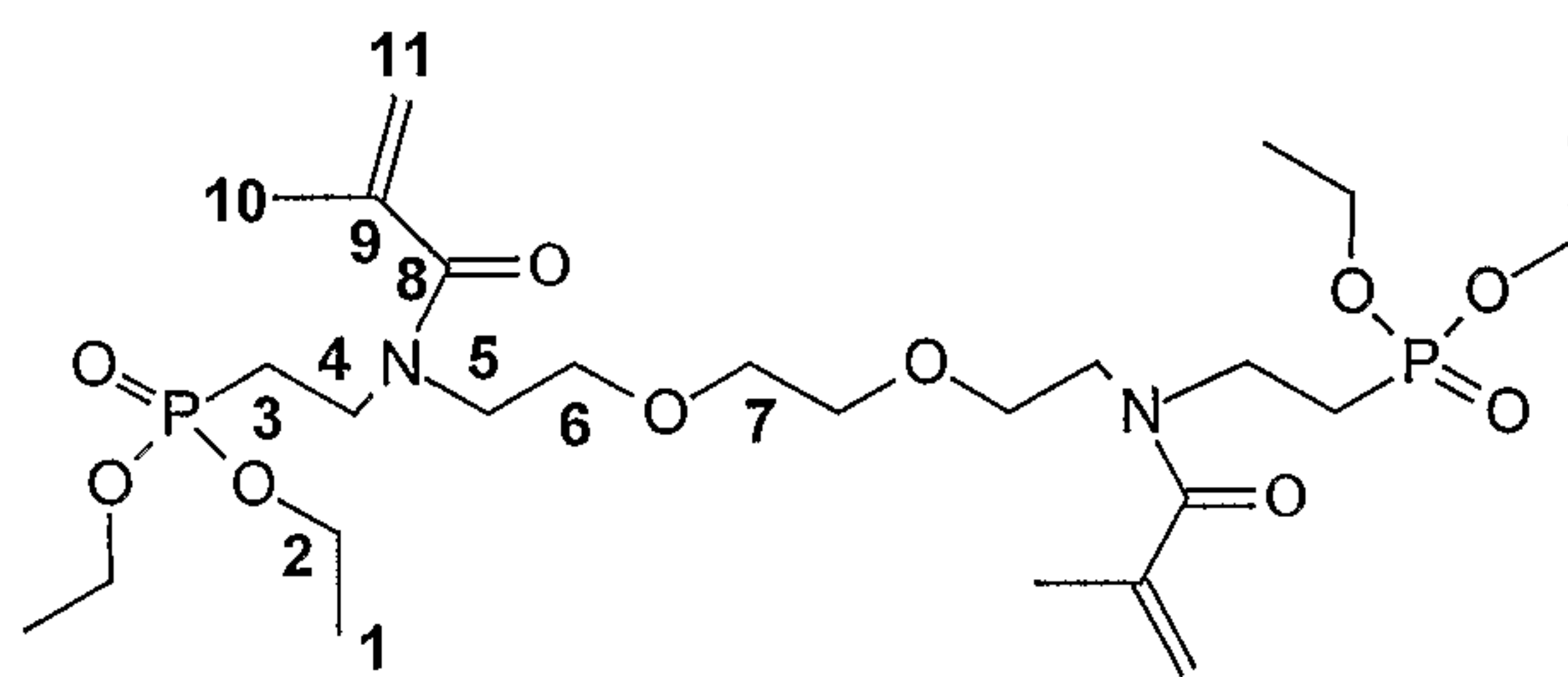
In a 4-necked 1-l-flask equipped with a stirrer, a thermometer and two 50 ml dropping funnels 30.000 g (0.063 mol) of N,N'-Bis (diethyl ethyl phosphonate)-1.2-Bis (2-aminoethoxy) ethane were dissolved in 300 ml of methylene chloride. After cooling to 0-5 °C 13.164 g (0.126 mol) of methacryloyl chloride dissolved in 30 ml of methylene chloride and 5.036 g (0.126 mol) of NaOH dissolved in 15.106 ml of water were added simultaneously under stirring during 1.5 hours so that the temperature remains at 0-5 °C. Thereafter the mixture were stirred at room temperature for additional two hours. Than the reaction mixture were hydrolyzed with 60

ml of ice-water. The organic phase were separated and the aqueous solution were extracted twice with methylene chloride. The collected organic liquids were washed with 50 ml of 1 n HCl, 50 ml of 1 n NaHCO₃ and sometimes with 50 ml of deionised water until the water shows a pH-value of approximately 7. Than the organic solution was dried over NaSO₄. Thereafter the NaSO₄ was filtered off and to the solution 0.039 g of 2,6-di-tert.-butyl-p-cresol were added. The methylene chloride was removed at 40 °C in vacuum and the bismethacrylamide was dried.

Yield: 32 g (83 % of th.)

(C₂₆H₅₀O₁₀N₂P₂), 612.64

¹³C-NMR: 167.7 (8), 140.2 (9), 121.7 (11), 70.6 (6), 69.4 (7), 62.3 (2), 48.2 (5), 32.5 (4), 25.1 (3), 19.3 (10), 16.4 (1)



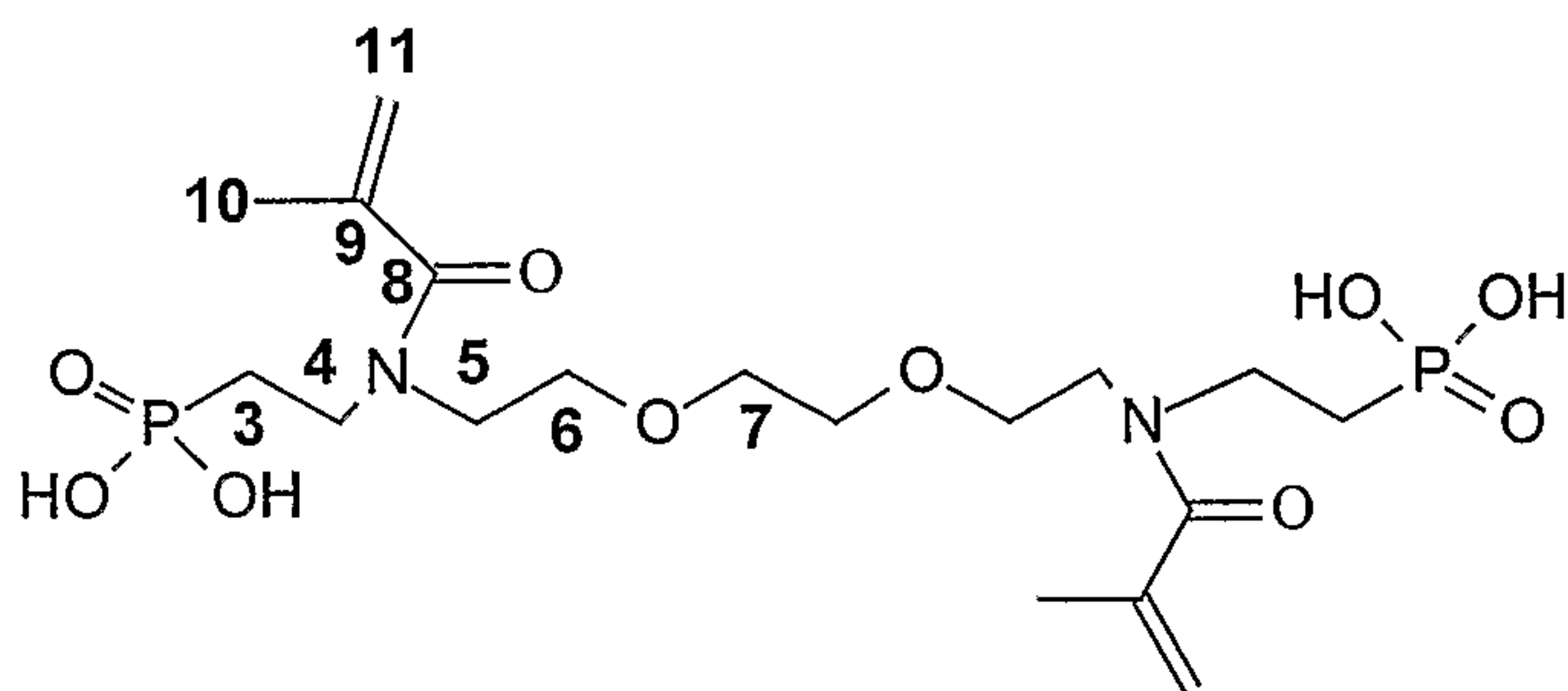
N,N'-Bis (ethyl phosphonic acid)-N,N'-bismethacrylamido-1,2-bis (2-aminoethoxy) ethane:

In a 4-necked 1-l-flask equipped with a stirrer, a thermometer, reflux cooler with CaCl₂-drying tube and 50 ml dropping funnels 30.000 g (0.049 mol) of N,N'-Bis (diethyl ethyl phosphonate)-N,N'-bismethacrylamido-1,2-Bis (2-aminoethoxy) ethane were dissolved in 100 ml of methylene chloride. Then 16.494 g (0.108 mol) Trimethyl bromsilane were added dropwise over an period of 20 minutes under stirring. Thereafter the reaction mixture was stirred for additional 2 hours. By adding of 100 methanol the phosphonic acid silylesters were hydrolyzed. Prior to remove the solvents BHT was added and the product was dried at 40 °C in vacuum.

Yield: 19.11 g (78.0 % d. Th)

(C₁₈H₃₄O₁₀N₂P₂), 500.42

¹³C-NMR: 164.7 (8), 140.2 (9), 121.7 (11), 70.6 (7), 69.4 (6), 48.2 (5), 31.9 (4), 29.5 (3), 19.3 (10)



Example 2

(Bis(3-methacryloylamidopropyl) diethylphosphonic acid ethylester:

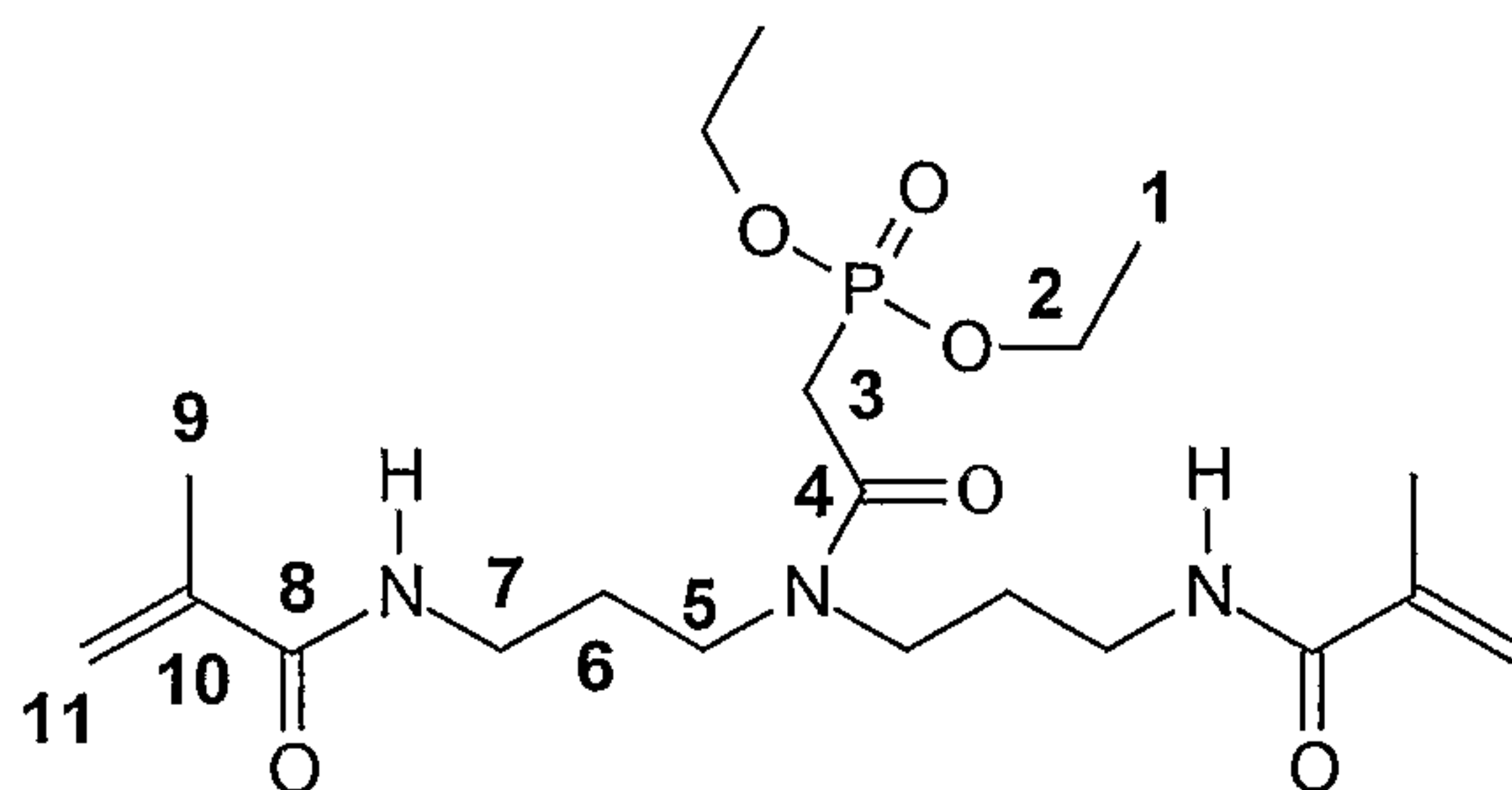
58.073 g (0.443 mol) Bis(3-aminopropyl)amine, 184.454 g (0.894 mol) Dicyclohexyl carbodiimide and 8.651 g (0.071 mol) dimethylamino pyridine were dissolved in a mixture of 250 ml CH_2Cl_2 and 100 ml Acetone. To the cooled mixture ($< 5^\circ\text{C}$) were added 76.200 g (0.885 mol) Methacrylic acid dissolved in 100 ml CH_2Cl_2 so that temperature do not pass 10°C . Then the mixture were stirred for 15 minutes at 0°C and for 20 hours at room temperature. Thereafter the reaction mixture was cooled again and 92.227 g (0.447 mol) Dicyclohexyl carbodiimide dissolved in a mixture of 50 ml CH_2Cl_2 and 50 ml Acetone were added. To this mixture was dropped a solution of 86.804 g (0.443 mol) Diethylphosphonic acid ethylester dissolved in 100 ml Acetone so that temperature do not pass 10°C . Then the mixture were stirred for 15 minutes at 0°C and for 20 hours at room temperature. After this time the precipitated solid was filtered off. To the filtrate were added 0.101 g BHT and the solvent was removed by vacuum distillation. The viscose residue was dissolved in 300 ml CH_2Cl_2 and cooled to 0°C . The precipitating solid was removed and the filtrate was washed twice with 150 ml 1n HCl, 150 ml 1n NaHCO_3 solution and with 150 ml water. Furthermore, the solution was dried over NaSO_4 and the solvent was removed. Than the solid was dissolved in Acetone again and dicyclohexyl urea was filtered of. Prior to remove the solvent, 0.197 g BHT was added, the product was dried in vacuum.

Yield: 155.7 g (79.0 % of th.)

($\text{C}_{20}\text{H}_{36}\text{N}_3\text{O}_6\text{P}$), 445.50

IR: 3010/2933/2856 (CH_2/CH_3), 1695 (CO), 1653/1627 (C=C), 1452 (CH_2/CH_3), 1251 cm^{-1}

^{13}C -NMR: 169.1 (4), 166.5 (8), 140.2 (10), 121.7 (11), 61.6 (2), 44.6 (5), 43.0 (7), 27.9 (6), 27.7 (3), 19.0 (9), 16.4 (1)



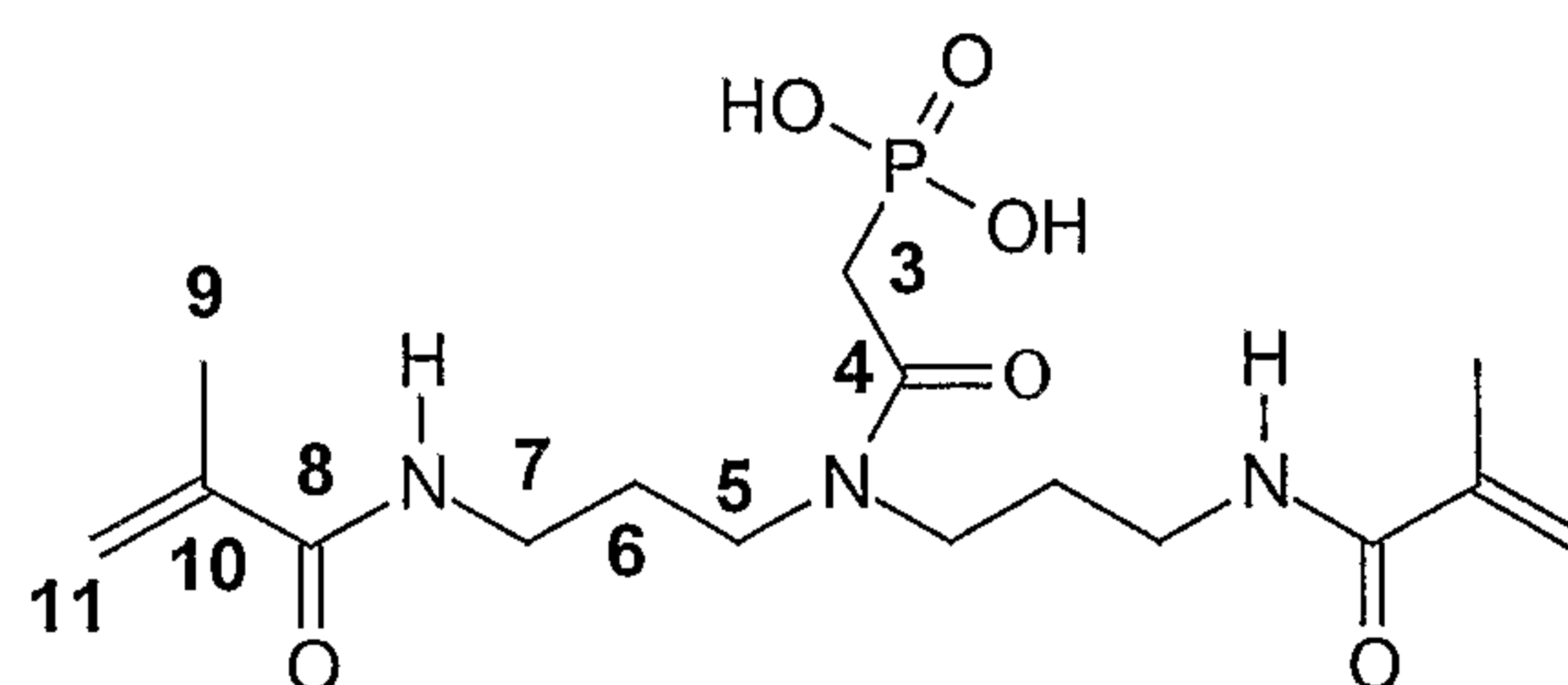
(Bis(3-methacryloylamidopropyl) diethylphosphonic acid:

In a 4-necked 1-l-flask equipped with a stirrer, a thermometer, reflux cooler with CaCl_2 -drying tube and 100 ml dropping funnel 155.700 g (0.350 mol) of (Bis(3-methacryloylamidopropyl) diethylphosphonic acid ethylester were dissolved in 100 ml of methylene chloride. Then 117.77 g (0.769 mol) Trimethyl bromsilane were added dropwise over an period of 60 minutes under stirring. Thereafter the reaction mixture was stirred for additional 2 hours. By adding of 250 methanol the phosphonic acid silylesters were hydrolyzed. Prior to remove the solvents 0.157 g BHT were added and the product was dried at 40 °C in vacuum.

Yield: 115.0 g (84.5 % d. Th)

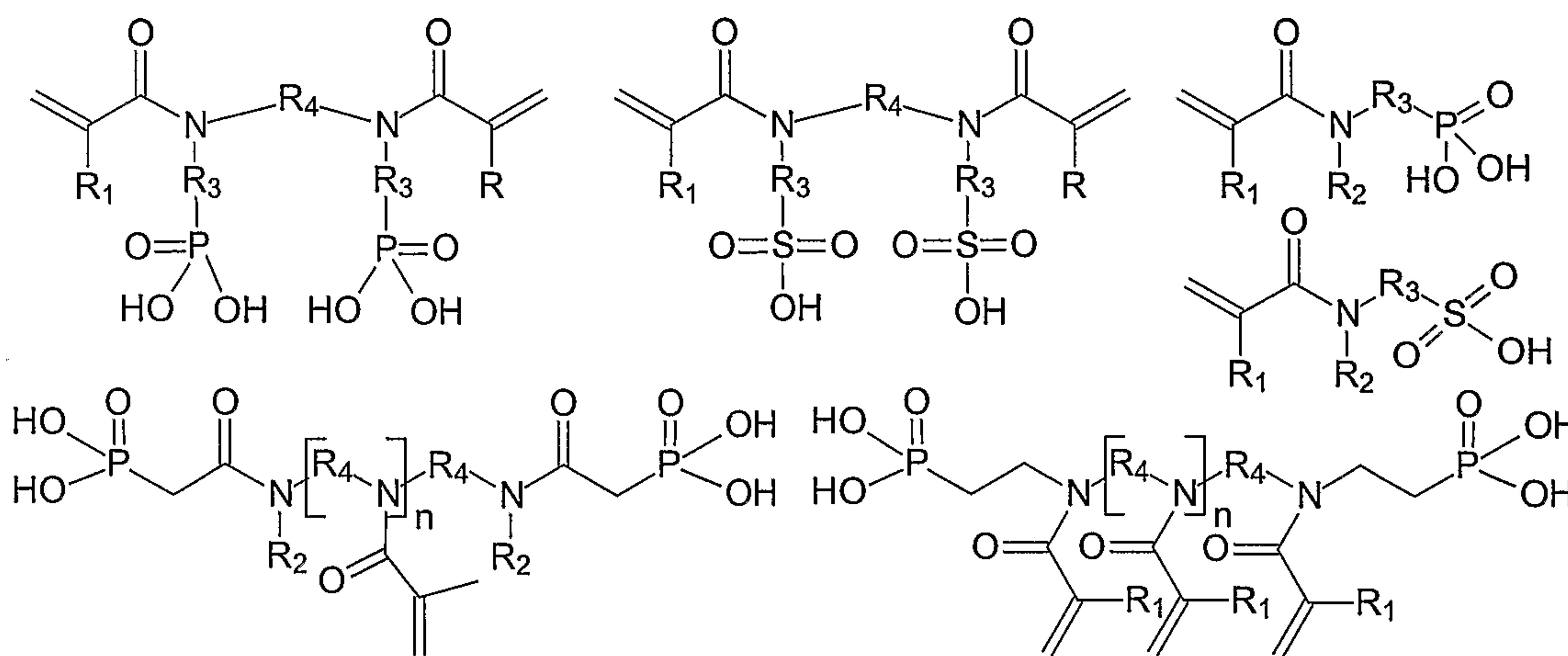
($\text{C}_{16}\text{H}_{28}\text{O}_6\text{N}_3\text{P}$), 389.39

^{13}C -NMR: 169.1 (4), 166.5 (8), 140.2 (10), 121.7 (11), 44.6 (5), 43.0 (7), 27.9 (6), 32.1 (3), 19.0 (9)



We claim

1. Hydrolysis stable one-part self-etching, self-priming dental adhesive comprising
 - (i) an acidic polymerizable (meth) acrylamide that comprises at least an organic or inorganic acidic moiety
 - (ii) a polymerizable monomer
 - (iii) polymerization initiator, inhibitor and stabilizer.
2. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, wherein said acidic polymerizable (meth) acrylamide comprises an organic or inorganic acidic moiety, preferably a carboxylic acid, phosphoric acid, sulfuric acid group and most preferably a phosphonic or sulfonic acid moiety.
3. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, wherein said acidic polymerizable (meth) acrylamide is selected from the group consisting of:



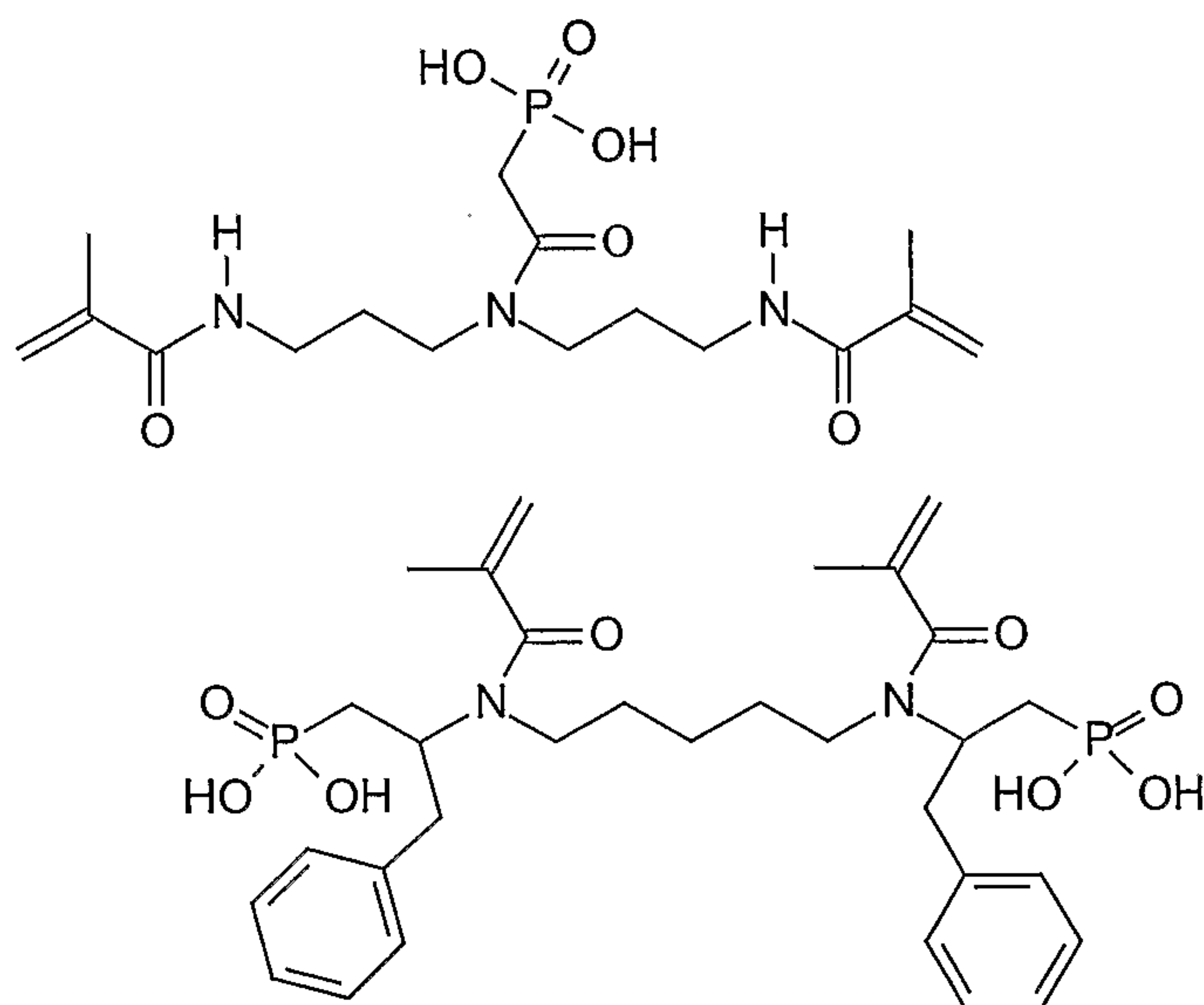
wherein

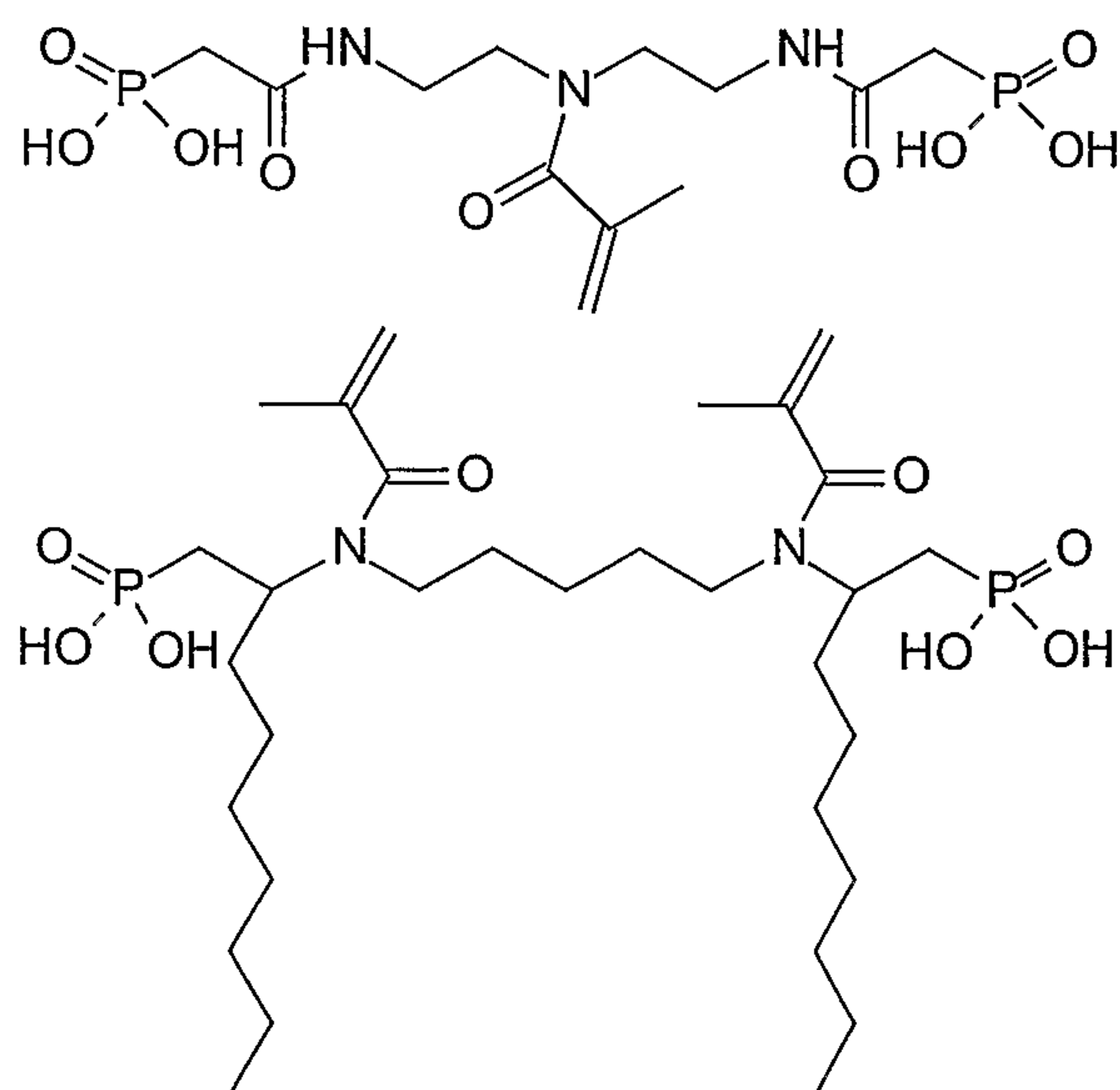
R_1 and R_2 independently are Hydrogen or a substituted or unsubstituted C_1 to C_{18} alkylene, substituted or unsubstituted cycloalkylene, substituted or unsubstituted C_5 to C_{18} arylene or heteroarylene, substituted or unsubstituted C_5 to C_{18} alkylarylene or alkylheteroarylene, substituted or unsubstituted C_7 to C_{30} alkylene arylene,

R_3 and R_2 independently are a difunctional substituted or unsubstituted C_1 to C_{18} alkylene, difunctional substituted or unsubstituted cycloalkylene, difunctional substituted or unsubstituted C_5 to C_{18} arylene or heteroarylene, difunctional substituted or unsubstituted C_5 to C_{18} alkylarylene or alkylheteroarylene, difunctional substituted or unsubstituted C_7 to C_{30} alkylene arylene,

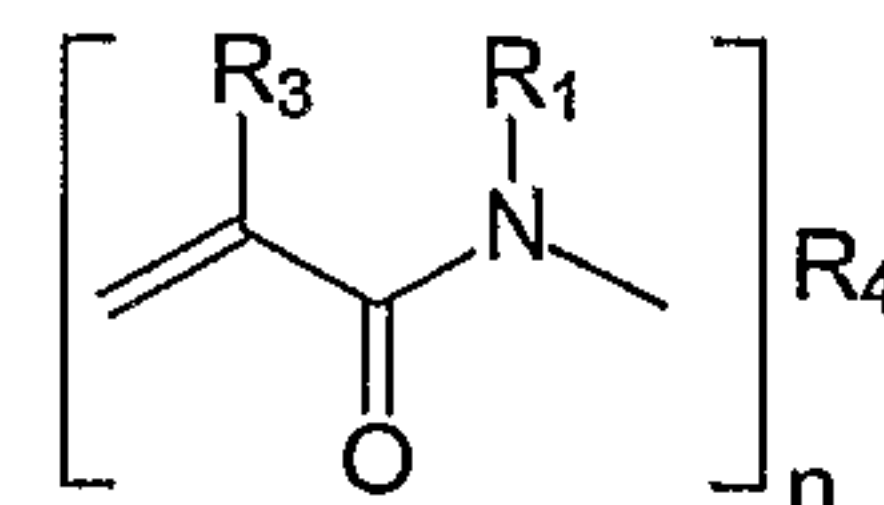
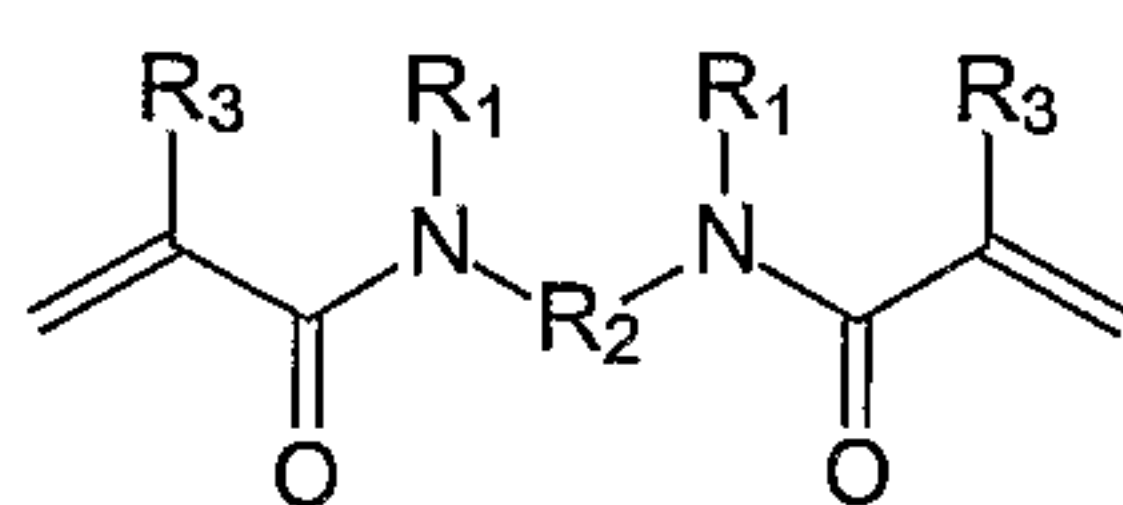
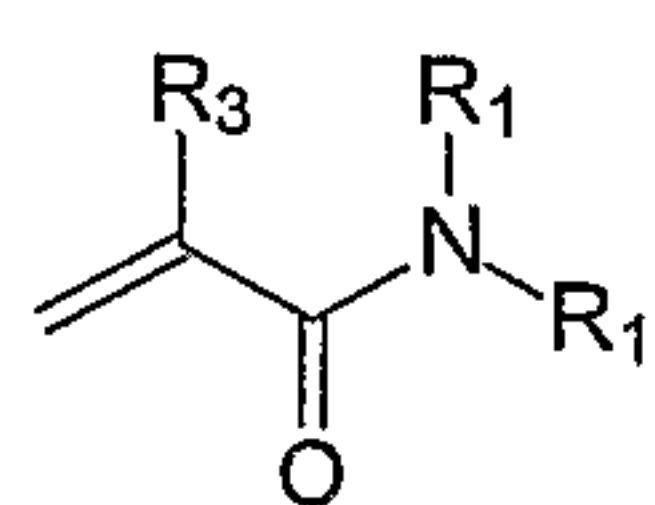
n is an integers.

4. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claims 1 to 3, wherein said acidic polymerizable (meth) acrylamide is selected from the group consisting of:





5. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, wherein polymerizable monomer is selected from the group consisting of:



wherein

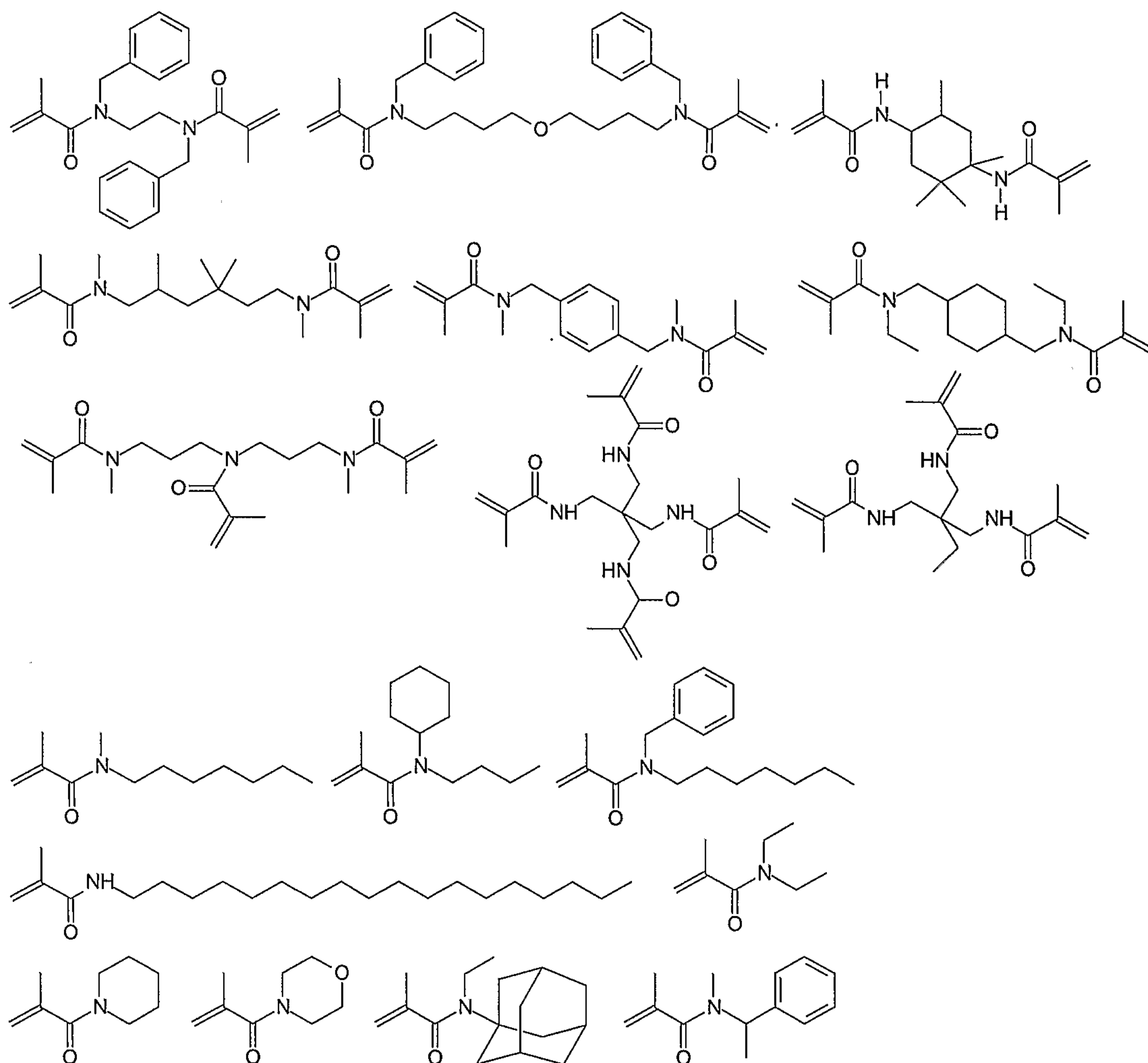
R_1 and R_3 independently are H or a substituted or unsubstituted C_1 to C_{18} alkylene, substituted or unsubstituted cycloalkylene, substituted or unsubstituted C_5 to C_{18} arylene or heteroarylene, substituted or unsubstituted C_5 to C_{18} alkylarylene or alkylheteroarylene, substituted or unsubstituted C_7 to C_{30} alkylene arylene,

R_2 is a difunctional substituted or unsubstituted C_1 to C_{18} alkylene, difunctional substituted or unsubstituted cycloalkylene, difunctional substituted or unsubstituted C_5 to C_{18} arylene or heteroarylene, difunctional substituted or unsubstituted C_5 to C_{18} alkylarylene or alkylheteroarylene, difunctional substituted or unsubstituted C_7 to C_{30} alkylene arylene,

R_4 is a mono- or polyfunctional substituted or unsubstituted C_1 to C_{18} alkylene, mono- or polyfunctional substituted or unsubstituted cycloalkylene,

mono- or polyfunctional substituted or unsubstituted C₅ to C₁₈ arylene or heteroarylene, mono- or polyfunctional substituted or unsubstituted C₅ to C₁₈ alkylarylene or alkylheteroarylene, mono- or polyfunctional substituted or unsubstituted C₇ to C₃₀ alkylene arylene,
n is an integers.

6. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, wherein said polymerizable monomer is a mono-, bis- or poly(meth) acrylamide that is selected from the group consisting of:



7. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, comprised of at least an acidic polymerizable (meth) acrylamide, a polymerizable monomer, an initiator, a stabilizer, pigments, an organic and/or inorganic filler.

8. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, comprising at least an acidic polymerizable (meth) acrylamide, a polymerizable monomer, an initiator, a stabilizer, water or organic solvent.
9. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, comprising of at least an acidic polymerizable (meth) acrylamide, a polymerizable monomer, an initiator, a stabilizer, pigments, an organic and/or inorganic filler, water or/and an organic solvent.
10. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, wherein said polymerizable monomers are applied in a content of 0 to 90 wt-%.
11. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, wherein said polymerization initiator is a thermal initiator, a redox-initiator or a photo initiator.
12. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, wherein said photo initiator preferably is champhor quinone.
13. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, wherein said filler is an inorganic filler and/or an organic filler; preferably the filler is a nanofiller.
14. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, wherein said stabilizer is a radical absorbing monomer such as hydroquinone monomethylether, 2,6-di-tert.-butyl-p-cresol.
15. Hydrolysis stable one-part self-etching, self-priming dental adhesive of claim 1, wherein said organic solvent preferably is acetone, ethanol or tert.-butanol.