

Description

Alkoxylated dendrimers and use thereof as biodegradable demulsifiers

- 5 The present invention relates to the use of alkoxylated dendrimers for breaking water-oil emulsions, in particular in the production of crude oil.

During its recovery, crude oil is produced as an emulsion with water. Before the crude oil is further processed, these crude oil emulsions must be
10 broken into the oil fraction and the water fraction. For this purpose, use is generally made of petroleum demulsifiers. Petroleum demulsifiers are surface-active polymeric compounds which are able to effect the required separation of the emulsion constituents within a short time.

- 15 Petroleum demulsifiers disclosed in US-4 321 146 are alkylene oxide block copolymers and in US-5 445 765 are alkoxylated polyethyleneimines. These can be used as individual components, in mixtures with other demulsifiers, or else as crosslinked products. Crosslinkings are carried out, for example, by means of reactions with alkoxylated low molecular weight
20 alcohols (such as, for example, glycerol or pentaerythritol) or alkoxylated alkylphenol formaldehyde resins with bifunctional compounds such as diepoxides or diisocyanates. Such crosslinked compounds are disclosed in US-5 759 409 and US-5 981 687.

- 25 US-4 558 120 and US 4 737 550 describe narrowly meshed star-shaped polymers (amidoamine dendrimers) which can be used as water-soluble demulsifiers (more accurately: deoilers) for crude oil/water emulsions. These products were neither alkoxylated nor used for breaking water/crude oil emulsions.

- 30 The use of alkoxylated, alkylated and esterified dendritic polyesters (in particular Boltorns® from Perstorp) is described in US-6 114 458. The products are reportedly suitable, inter alia, as curing agents for heat-curing resins, as viscosity modifiers for linear polymers or as stabilizers for
35 suspensions and emulsions.

The varying properties (e.g. asphaltene, paraffin and salt content, chemical composition of the natural emulsifiers) and proportions of water in different

crude oils make it imperative to further develop the existing petroleum demulsifiers. In particular, a low concentration and broad applicability of the petroleum demulsifiers to be used as well as the higher effectiveness which is to be strived for is of prime importance from an economic and ecological point of view. In addition, demulsifiers are increasingly required which have good biodegradability and low bioaccumulation in order to replace the controversial products based on alkylphenol.

The object was thus to develop novel alkylphenol-free petroleum demulsifiers which are superior in their effect to the products already known, can be used in an even lower concentration and have better biodegradability.

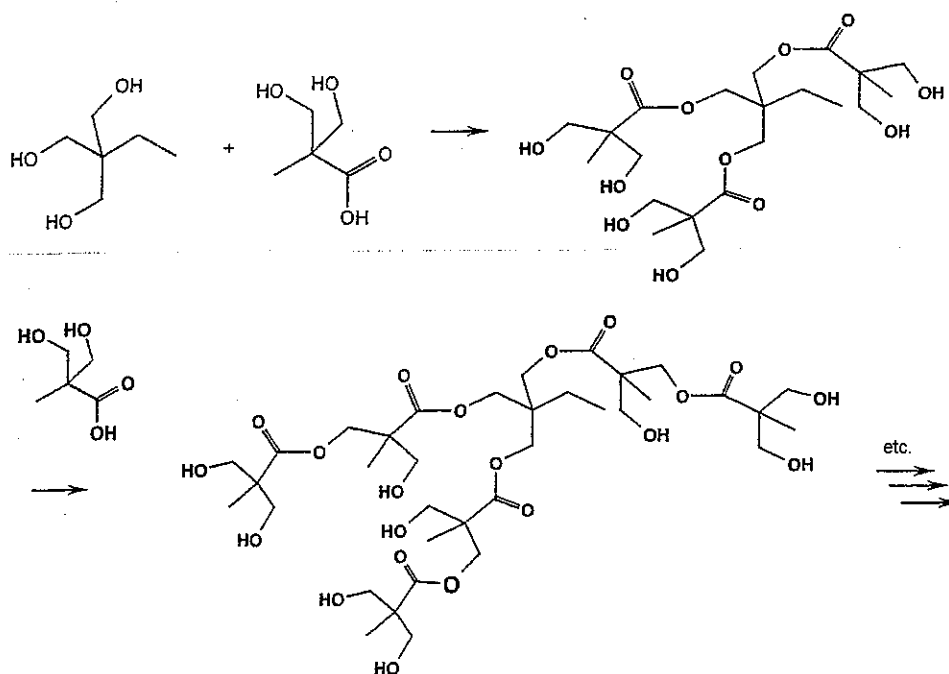
Surprisingly, it has been found that alkoxyated dendritic polyesters (dendrimers) exhibit an excellent effect as petroleum demulsifiers even at a very low concentration. In addition, they have exhibited significantly better biodegradabilities in accordance with OECD 306 compared to many conventional commercial demulsifiers.

The invention therefore provides for the use of alkoxyated dendrimers having a molecular weight of from 2400 to 100 000 g/mol which have been alkoxyated with C₂-C₄-alkylene oxide groups or a mixture of such alkylene oxide groups such that the alkoxyated dendrimer has a degree of alkoxylation of from 1 to 100 alkylene oxide units per free OH group, for breaking oil/water emulsions, in amounts of from 0.0001 to 5% by weight, based on the oil content of the emulsion to be broken.

These alkoxyated dendrimers are preferably obtainable from dendrimers by alkoxylation of the free OH groups with a C₂-C₄-alkylene oxide or a mixture of such alkylene oxides in a molar excess so that the alkoxyated dendrimer has the specified degree of alkoxylation. Dendrimers are also commercially available. Particular preference is given to using Boltorn® H2O and H310 (Perstorp).

The preparation and the molecular structures of the dendrimers used according to the invention are described comprehensively in US-5 418 301. These are dendritic polyesters which are constructed from a central starting molecule (generally a diol or polyol) and subsequent successive multiple

esterification with a carboxylic acid having two further reactive groups (such as, for example, dimethylolpropanoic acid). Fig. 1 shows the reaction principle.



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Fig. 1: Synthesis in principle of dendritic polyesters by reaction of trimethylolpropane and 2,2-dimethylolpropanoic acid

- 10 The dendrimers used for the alkoxylation are dendritic polyesters, in general based on a mono-, di- or polyfunctional starting alcohol and a carboxylic acid which has as least two hydroxyl groups as dendritic growth component. The starting alcohol used is preferably bis(trimethylolpropane), bis(trimethylolpropane), dipentaerythritol, pentaerythritol, alkoxyated
- 15 pentaerythritol, trimethylolpropane, trimethylolpropane, alkoxyated trimethylolpropane, glycerol, diglycerol, triglycerol, polyglycerol, neopentyl glycol, dimethylolpropane, sorbitol or mannitol.

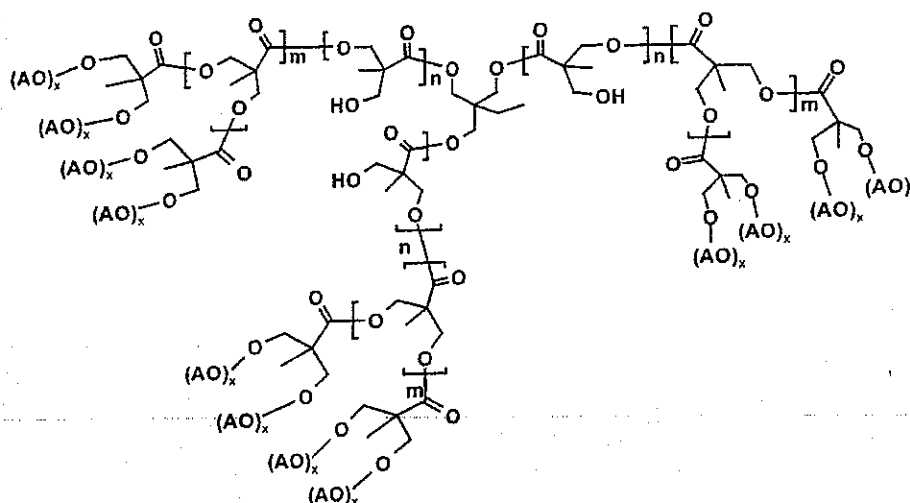
20 The carboxylic acid which leads to the dendritic chain growth is preferably dimethylolpropanoic acid, α,α -bis(hydroxymethyl)butanoic acid, α,α,α -tris(hydroxymethyl)ethanoic acid, α,α -bis(hydroxymethyl)pentanoic acid, α,α -bis(hydroxy)propanoic acid or 3,5-dihydroxybenzoic acid.

The star-shaped monodisperse dendrimers are alkoxyated with one or more C₂-C₄-alkylene oxides, preferably ethylene oxide (EO) or propylene oxide (PO). The alkoxyating agent is used in molar excess. As is known in the prior art, the alkoxylation takes place by reacting the dendrimers with
5 an alkylene oxide under elevated pressure of generally 1.1 to 20 bar at temperatures of from 50 to 200°C. The alkoxylation takes place at the free OH groups of the dendrimers. The amount of alkylene oxide used is such that the average degree of alkoxylation is between 1 and 100 alkylene oxide units per free OH group. Average degree of alkoxylation is
10 understood here as meaning the average number of alkoxy units which is attached to each free OH group. It is preferably 2 to 70, in particular 5 to 50, specifically 20 to 40.

Preferably, the alkoxylation is carried out firstly with PO and then with EO. The ratio of EO to PO in the alkoxyated dendrimer is preferably between
15 1:1 and 1:10, in particular 1:2 to 1:10. According to the invention, however, the alkoxylation can also take place in the reverse order, firstly EO, then PO, or with a mixture of PO and EO.

The dendrimer obtained following alkoxylation preferably has a molecular
20 weight of from 2400 to 80 000 units, in particular from 10 000 to 50 000 units, specifically 15 000 to 30 000.

The alkoxyated dendrimers prepared by the described process are reproduced by way of example on the basis of a dendrimer of
25 trimethylolpropane and 2,2-dimethylolpropanoic acid by the following structure (formula 1):



(AO)_x-O radicals are the alkoxyated OH groups, in which AO is a C₂-C₄-alkylene oxide unit and x is the degrees of alkoxylation. n and m are the corresponding degrees of condensation of 2,2-dimethylolpropanoic acid and are described in detail in US-5 418 301.

A preferred subject-matter of the present invention is the use of the alkoxyated dendrimers as demulsifiers for oil/water emulsions in the recovery of petroleum.

According to the invention, the alkoxyated dendrimers for increasing the molecular weight and thus for improving the demulsifying properties can be reacted with multifunctional crosslinkers as is known in the prior art.

The following crosslinkers are particularly preferred:

bisphenol A diglycidyl ether, butane-1,4-diol diglycidyl ether, hexane-1,6-diol diglycidyl ether, ethylene glycol diglycidyl ether, cyclohexanedimethanol diglycidyl ether, resorcinol diglycidyl ether, glycerol diglycidyl ether, glycerol triglycidyl ether, glycerol propoxylate triglycidyl ether, polyglycerol polyglycidyl ether, p-aminophenol triglycidyl ether, polypropylene glycol diglycidyl ether, pentaerythritol tetraglycidyl ether, sorbitol polyglycidyl ether, trimethylolpropane triglycidyl ether, castor oil triglycidyl ether, diaminobiphenyl tetraglycidyl ether, soya oil epoxide, adipic acid, maleic acid, phthalic acid, maleic anhydride, succinic anhydride, dodecylsuccinic anhydride, phthalic anhydride, trimellitic anhydride, pyromellitic anhydride, dimethoxydimethylsilane, diethoxydimethylsilane, tetraalkoxysilanes, toluene diisocyanate, diphenylmethane diisocyanate.

The specified crosslinkers and chemically related compounds are preferably used in the range from 0.1-10% by weight, particularly preferably 0.5-5% by weight and specifically 1.0-2.5% by weight, based on the alkoxyated dendrimer.

For use as petroleum demulsifiers, the alkoxyated dendrimers are added to the water/oil emulsions, which preferably takes place in solution. Preferred solvents for the alkoxyated dendrimers are paraffinic or aromatic solvents. The crosslinked alkoxyated dendrimers are used in amounts of

from 0.0001 to 5% by weight, preferably 0.0005 to 2% by weight, in particular 0.0008 to 1% by weight and specifically 0.001 to 0.1% by weight, based on the oil content of the emulsion to be broken.

5 General experimental procedures for the alkoxylation

Ethylene oxide

10 The dendrimers specified in the examples below were introduced into a 1 l glass autoclave and the pressure in the autoclave was adjusted to about 0.2 bar above atmospheric pressure with nitrogen. It was slowly heated to 140°C and, after reaching this temperature, the pressure was again adjusted to 0.2 bar above atmospheric pressure. Then, at 140°C, the desired amount of EO (see examples) was metered in, during which the
15 pressure should not exceed 4.5 bar. When the EO addition was complete, the mixture was left to after-react for a further 30 minutes at 140°C.

Propylene oxide

20 The dendrimers specified in the examples below were introduced into a 1 l glass autoclave and the pressure in the autoclave was adjusted to about 0.2 bar above atmospheric pressure with nitrogen. It was slowly heated to 130°C and, after reaching this temperature, the pressure was again adjusted to 0.2 bar above atmospheric pressure. Then, at 130°C, the
25 desired amount of PO was metered in (see examples), during which the pressure should not exceed 4.0 bar. When the PO addition was complete, the mixture was left to after-react for a further 30 minutes at 130°C.

30 The degree of alkoxylation was determined by means of ^{13}C NMR. This determination method should always be used for the purposes of the present invention.

Boltorn® H20 and Boltorn® H310 are dendrimers of a dihydroxycarboxylic acid and a polyol.

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Examples

Example 1:

Boltorn[®] H20 + 10 mol PO/OH + 5 mol EO/OH
(average molar mass according to GPC about 11 500 g/mol)

Example 2:

- 5 Boltorn[®] H20 + 20 mol PO/OH + 10 mol EO/OH
(average molar mass according to GPC about 22 900 g/mol)

Example 3:

- 10 Boltorn[®] H20 + 10 mol EO/OH + 10 mol PO/OH
(average molar mass according to GPC about 15 200 g/mol)

Example 4:

- 15 Boltorn[®] H20 + [10 mol EO/OH + 10 mol PO/OH] mixture
(average molar mass according to GPC about 15 600 g/mol)

Example 5:

Boltorn[®] H20 + 40 mol PO/OH + 20 mol EO/OH
(average molar mass according to GPC about 47 500 g/mol)

20 Example 6:

Boltorn[®] H310 + 20 mol PO/OH + 10 mol EO/OH
(average molar mass according to GPC about 39 600 g/mol)

Example 7:

- 25 Boltorn[®] H310 + [20 mol PO/OH + 10 mol EO/OH] mixture
(average molar mass according to GPC about 40 200 g/mol)

Example 8:

- 30 Boltorn[®] H310 + 20 mol PO/OH + 20 mol EO/OH
(average molar mass according to GPC about 53 500 g/mol)

Example 9:

Boltorn[®] H310 + 10 mol EO/OH + 10 mol PO/OH
(average molar mass according to GPC about 27 100 g/mol)

Example 10:

Crosslinking the alkoxyated dendrimer from Example 1 with bisphenol A diglycidyl ether

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In a 250 ml three-necked flask with contact thermometer, stirrer and reflux condenser, 100 g of alkoxyated dendrimer were heated to 80°C with gentle nitrogen flushing. At this temperature, 2.5 g of bisphenol A diglycidyl ether (80% strength solution in an aromatic solvent) were quickly added dropwise. The reaction temperature was then increased to 120°C and the reaction mixture was stirred for 8 h until unreacted diglycidyl ether could no longer be detected by means of titration of the epoxy number. The product was evaporated to dryness on a rotary evaporator (yield: 101.9 g) and the molar mass was analyzed by means of GPC ($M^* \approx 18\,200$ g/mol, standard polyethylene glycol).

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Example 11

Crosslinking the alkoxyated dendrimer from Example 1 with dodecylsuccinic anhydride

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In a 250 ml three-necked flask with contact thermometer, stirrer and water separator, 100.0 g of alkoxyated dendrimer, 1.5 g of alkylbenzenesulfonic acid and 3.5 g of dodecylsuccinic anhydride were initially introduced at room temperature. The reaction mixture was then heated to 165°C and stirred for a further 8 h at this temperature until no more water of reaction formed in the water separator (reaction control: acid number). The product was evaporated to dryness on a rotary evaporator (yield: 102.9 g) and the molar mass was analyzed by means of GPC ($M^* \approx 19\,800$ g/mol, standard polyethylene glycol).

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Example 12

Crosslinking the alkoxyated dendrimer from Example 1 with toluene 2,4-diisocyanate

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In a 250 ml three-necked flask with contact thermometer, stirrer and reflux condenser, 100.0 g of alkoxyated dendrimer were heated to 60°C with gentle nitrogen flushing. Then, at this temperature, 3.0 g of toluene 2,4-diisocyanate were slowly added dropwise. The reaction temperature

was increased to 100°C and the reaction mixture was stirred for a further 8 h at this temperature (reaction control: isocyanate number). The product was evaporated to dryness on a rotary evaporator (yield: 102.8 g) and the molar mass was analyzed by means of GPC ($M^* \approx 21\,400$ g/mol, standard polyethylene glycol).

Determining the breaking effectiveness of petroleum demulsifiers

To determine the effectiveness of a demulsifier, the water separation from a crude oil emulsion per time, and also the dewatering and desalting of the oil were determined. For this, demulsifying glasses (tapered, graduated glass bottles with screw lids) were charged in each case with 100 ml of the crude oil emulsion, in each case a defined amount of the demulsifier was metered in just below the surface of the oil emulsion using a micropipette, and the demulsifier was mixed into the emulsion by intensive shaking. The demulsifying glasses were then placed into a conditioning bath (30°C and 50°C) and the water separation was monitored.

During demulsification and after it had finished, samples were taken from the oil from the upper section of the demulsifying glass (so-called top oil), and the water content was determined in accordance with Karl Fischer and the salt content was determined conductometrically. In this way, it was possible to assess the novel demulsifiers according to water separation and also dewatering and desalting of the oil.

Breaking effect of the demulsifiers described

Origin of the crude oil emulsion: Holzkirchen sonde 3, Germany

Water content of the emulsion: 46%

Salt content of the emulsion: 5%

Demulsification temperature: 50°C

Table 1:
Effectiveness of alkoxyated dendrimers as demulsifiers compared to standard products (concentration 20 ppm)

Water separation [ml] per time [min]	5	10	20	30	45	60	90	120	180	Water in the top oil [%]	Salt in the top oil [ppm]
Product from 1	0	0	5	11	27	33	38	42	44	0.79	152
Product from 2	1	4	10	16	35	42	45	45	45	0.40	98
Product from 3	2	5	12	18	36	42	45	45	46	0.32	73
Product from 4	2	5	13	19	37	43	45	45	45	0.38	68
Product from 5	4	9	18	27	38	43	45	46	46	0.14	25
Product from 6	1	5	12	18	34	42	44	44	45	0.58	94
Product from 7	1	6	13	19	34	43	45	45	45	0.51	81
Product from 8	2	8	17	28	39	44	45	46	46	0.27	31
Product from 9	5	10	23	32	40	45	46	46	46	0.21	20
Product from 10	4	9	23	34	42	45	45	46	46	0.27	32
Product from 11	2	5	13	25	35	42	42	43	43	0.59	78
Product from 12	3	10	24	32	40	45	45	45	46	0.15	22
Standard 1: Dissolvan 1952 (comparison)	0	0	3	6	10	17	23	28	32	1.59	420
Standard 2: Dissolvan 4738 (comparison)	0	0	0	4	10	24	33	39	39	0.92	205

Table 2:

Biodegradability of alkoxyated dendrimers (*closed bottle test* in accordance with OECD 306) compared to standard products

Biodegradability [%] after	14 days	28 days
Product from 1	25.7	46.8
Product from 6	32.9	55.6
Product from 10	30.7	49.5
Product from 12	45.7	62.5
Standard: Dissolvan 1952 (comparison)	4.0	9.3
Standard: Dissolvan 4738 (comparison)	19.8	26.3
Reference (sodium benzoate) (comparison)	61.7	82.4

Claims

1. The use of alkoxyated dendrimers having a molecular weight of from 2400 to 100 000 g/mol which have been alkoxyated with C₂-C₄-alkylene oxide groups or a mixture of such alkylene oxide groups such that the alkoxyated dendrimer has a degree of alkoxylation of from 1 to 100 alkylene oxide units per free OH group, for breaking oil/water emulsions, in amounts of from 0.0001 to 5% by weight, based on the oil content of the emulsion to be broken.
2. The use as claimed in claim 1, where the dendrimer is a dendritic polyester based on a mono-, di- or polyfunctional starting alcohol and a carboxylic acid as dendritic growth component which has at least two hydroxyl groups.
3. The use as claimed in claim 1 and/or 2, where the starting alcohol used is bis(trimethylolpropane), bis(trimethylolethane), dipentaerythritol, pentaerythritol, alkoxyated pentaerythritol, trimethylolethane, trimethylolpropane, alkoxyated trimethylolpropane, glycerol, diglycerol, triglycerol, polyglycerol, neopentyl glycol, dimethylolpropane, sorbitol or mannitol.
4. The use as claimed in one or more of claims 1 to 3, where the carboxylic acid to the dendritic chain growth is dimethylolpropanoic acid, α,α -bis(hydroxymethyl)butanoic acid, α,α,α -tris(hydroxymethyl)ethanoic acid, α,α -bis(hydroxymethyl)pentanoic acid, α,α -bis(hydroxy)propanoic acid or 3,5-dihydroxybenzoic acid.
5. The use as claimed in one or more of claims 1 to 4, where the alkoxyated dendrimers have a molecular weight of from 10 000 to 50 000 g/mol.
6. The use as claimed in one or more of claims 1 to 5, in which the average degree of alkoxylation is between 1 and 70 alkylene oxide units per free OH group.
7. The use as claimed in one or more of claims 1 to 6, in which the alkylene oxide is ethylene oxide or propylene oxide.

8. The use as claimed in one or more of claims 1 to 7, in which a mixed alkoxylation with ethylene oxide and propylene oxide in the ratio from 1:2 to 1:10 is present.

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9. The use as claimed in one or more of claims 1 to 8, where the alkoxyated dendrimers are crosslinked using bisphenol A diglycidyl ether, butane-1,4-diol diglycidyl ether, hexane-1,6-diol diglycidyl ether, ethylene glycol diglycidyl ether, cyclohexanedimethanol diglycidyl ether, resorcinol diglycidyl ether, glycerol diglycidyl ether, glycerol triglycidyl ether, glycerol propoxylate triglycidyl ether, polyglycerol polyglycidyl ether, p-aminophenol triglycidyl ether, polypropylene glycol diglycidyl ether, pentaerythritol tetraglycidyl ether, sorbitol polyglycidyl ether, trimethylolpropane triglycidyl ether, castor oil triglycidyl ether, diaminobiphenyl tetraglycidyl ether, soya oil epoxide, adipic acid, maleic acid, phthalic acid, maleic anhydride, succinic anhydride, dodecylsuccinic anhydride, phthalic anhydride, trimellitic anhydride, pyromellitic anhydride, dimethoxydimethylsilane, diethoxydimethylsilane, toluene diisocyanate, diphenylmethane diisocyanate.

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