

Pesticide and/or Herbicide Compositions and Methods of Their Use

The present invention relates to pesticide and/or herbicide compositions including such compositions that are sprayable and contain at least one
5 pesticide and/or herbicide and exhibit improved efficacy test and/or weed control. In the case of systemic herbicides a much greater uptake by plants is observed. Also included are aqueous concentrates that can be used to prepare sprayable compositions. The invention also provides a method of treating a
10 crop area employing the composition of the invention.

It is well known in agriculture to apply various agrochemicals to growing areas by spraying. The growing areas may be crop areas, which can be very large, or smaller growing areas such as those in greenhouses. The agrochemicals applied as sprays include fertilisers, herbicides and pesticides.

15 Herbicides and pesticides can be supplied to the farmer in various forms, for instance as neat liquids, aqueous solutions, aqueous dispersions or slurries of solid herbicide or pesticide. It is normal practice for the manufacturer to supply the farmer with the herbicide or pesticide in the form of a neat liquid or as a high
20 activity solution or slurry. The usual way of applying herbicides or pesticides to an area of land would be by spraying.

Various systems have been devised for convenient dosing of herbicides or pesticides. Spray pumps are well known which spray water from a spray
25 manifold onto the area of land or crop area and which are designed so that concentrated herbicide or pesticide can be dosed into the pump, mixed with water before being sprayed.

During the spraying of herbicides and pesticides it is common to apply anti-drift
30 agents in order to prevent the formation of fine droplets which could be carried beyond the area intended to be treated. Without the use of anti-drift agents, the

spraying of herbicides and pesticides would be inefficient, first of all because there could be inadequate treatment of the land and crop areas intended to be treated and secondly the extraneous spray, if carried beyond the intended treatment zone, could for example be detrimental to other crops, land and water courses.

It is usual to combine the anti-drift agent with either the water which is fed into the spray pumps or to apply it directly into the spray pumps, usually at or shortly after the mixing zone where the water is mixed with the herbicide or pesticide. It is important that the spray drift chemical is metered at the correct dose to ensure that extraneous spray is not formed through under dosing or through overdosing the spray angle is too narrow resulting in uneven distribution of the herbicide or fertiliser.

Polymers of acrylamide and other ethylenically unsaturated monomers have been used as anti-drift agents. It has been generally accepted that polymers which give optimum spray drift control are either non-ionic (eg acrylamide homopolymer) or have relatively low anionic content (e.g. 5 to 30 wt. %) and also have relatively high intrinsic viscosity, for instance above 6 dl/g. Such polymers tend to form viscous aqueous solutions unless used at low concentration. Normal practice is to mix the polymer powder or reverse phase emulsion form with water directly into the spray tank so as to form an aqueous solution of polymer. However, this has the problem that emulsion polymers can be difficult to activate in this situation and polymer powders take a long time to dissolve. It is sometimes necessary to use more polymer as a result of inefficient dissolution of the polymer. Normally in order to minimise the problems with dissolution it would be usual to use polymers of intrinsic viscosity in the range 6 to 15 dl/g. Typically the water containing the pesticide or herbicide would comprise polymer at a concentration in excess of 0.05wt.%.

Canadian 1023264 (Vartiak) describes dilute aqueous sprayable formulations using polymers in the form of reverse phase emulsions to an aqueous system containing pesticides or herbicides and surfactants to prepare sprayable formulations.

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US 6534563 (Bergeron) discloses formulations comprising pesticides or agents for controlling plant growth. The plant protection formulations can be used in the presence of an aqueous solution and conventionally sprayed onto the plant to be treated.

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For some applications it is usual to combine two or more agrochemicals. For instance in the application of herbicides, especially systemic herbicides, it is usual to combine the treatment with a fertiliser, such as for instance ammonium sulphate (AMS). The fertiliser stimulates the growth of unwanted plants causing them to take up much more water, together with the herbicide, through the root system. This ensures a more efficient uptake and distribution of herbicide throughout the plant. In this instance the fertiliser may be regarded as an adjuvant in that it increases the efficacy of the herbicide. Thus a fertiliser used in combination with a herbicide is termed a herbicide adjuvant

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Polymeric anti drift agents or anti deposition agents used with herbicides such as glyphosate tend to be added to the spray tank as a separate component to the herbicide formulation. The addition of such polymers directly into a pesticide or herbicide formulation tends to be less favorable for a number of reasons. Generally high concentrations of polymeric materials can be incompatible with both the surfactant and the pesticide or herbicide in aqueous concentrates. Furthermore the polymers may not be sufficiently soluble in the concentrates containing high concentrations of pesticide/herbicide and surfactant. In addition due to the low activity of conventional polymers it is normally necessary to add the polymers at high concentrations and in an aqueous concentrate this generally would not be possible without first reducing

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the pesticide and/or herbicide active level or the surfactant level. Another disadvantage of conventional polymeric anti drift agents or anti deposition agents is that they tend to viscosify the formulations and this may not be acceptable.

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WO-A-00/26160 overcomes some other disadvantages of conventional formulations. This document describes an agrochemical liquid concentrate containing herbicide or pesticide and an inorganic water-soluble compound adjuvant in an amount of at least 10% by weight. The concentrate includes an
10 anti-drift agent which is a water-soluble anionic polymer in an amount up to 1.9 wt.% based on the weight of the concentrate. The concentrate would be diluted in a spray tank usually at least 30 fold, for instance at least 50 fold to form a sprayable composition. The systemic herbicide glyphosate is described as one example of an agrochemical. This composition provides significant
15 improvements in reducing spray drift.

An article by Mickey R. Brigance entitled, "Effect of Polyacrylamide Formulations on Bioefficacy of Glyphosate" (Proceedings of the 7th International Symposium on Adjuvants for Agrochemicals, 8 to 12 November
20 2004 (ISBN No. 1-92 0-01716-X)) describes the evaluation of sprayable herbicidal compositions of glyphosate and the enhancement by polyacrylamides in simple and multifunctional tank mix formulations. The study found that the higher surfactant loaded glyphosate formulations showed smaller increases in efficacy from the polyacrylamide inclusion and that the glyphosate formulations
25 with very low or no surfactants showed the greatest efficacy increase. This study considers drift suppression by reducing fine droplets and reducing droplet bounce for herbicidal formulations.

US 5525575 (Chamberlain) describes an aqueous concentrate that can contain
30 polymers. However, the polymer must have low molecular weight and preferably this is such that its presence in composition does not substantially

affect the spray pattern on the composition.

UK patent application 0503152.1 (internal attorney docket number AG/322371/P1) unpublished at the priority date of the present application,

5 describes a sprayable foliar feed composition comprising, water containing therein,

- i) at least one foliar feed plant nutrient,
- ii) 40 to 1000 ppm of at least one water-soluble polymer, and
- iii) 40 to 5000 ppm of at least one surfactant.

10 This disclosure reveals a composition providing acceptable or improved spray drift and droplet bounce and that the same time provides a significant reduction in droplet runoff.

U.S. 5525575 describes the incorporation of polymers into pesticide or herbicide compositions in order to improve the performance of the actives. It is stated
15 that the polymer may improve transport of the active through the cell wall of the plant. The polymers described all exhibit intrinsic viscosity off below 3 dl/g. Polymers of this molecular weight would not reduce spray drift. Copolymers of acrylamide with dimethyl amino ethyl acrylate (methylchloride quaternary salt)
20 are mentioned but it is stated that the acrylamide content of these materials must be greater than 80%. We have found that these formulations tend to give inadequate droplet bounce control and drift control.

US 6288010 describes the use of anionic polymer in concentrated salts,
25 primarily ammonium sulphate. The polymers mentioned would not be incorporated easily into glyphosate formulations.

U.S. 2004 0058821 refers to the use of anionic and non-ionic polymer either as additives within a tank mix or pesticide formulations. The compatibility of the
30 polymer is improved by co formulating the polymer with a salt, usually ammonium sulphate.

- U.S. 2001 0034304 refers to the addition of polymeric deposition agents within a pesticide. Specifically mentioned are cationic guar gum and/or polyacrylamides which are exemplified as anionic polymers of acrylamide. The example describes compositions comprising high levels of polymer and employing salts such as ammonium sulphate in order to achieve a stable formulation. The exemplified formulations appear to be saturated with little possibility for the glyphosate formulation to include additional surfactant. U.S. 2004 0211234 describes similar compositions.
- 10 US 6210696 is concerned with low molecular weight copolymers of dimethyl amino ethyl acrylate with water in soluble monomers such as alkyl acrylates to stabilize dispersions of oil. The polymers described would be unsuitable as anti drift agents due to their low water solubility and low molecular weight.
- 15 There still exists a need to provide a pesticide or herbicide formulation that provides significantly improved spray drift control. It would also be desirable to provide a composition that enables improved deposition and penetration of the active whilst at the same time improving spray drift control.
- 20 Furthermore, prior art formulations generally cannot be formulated at concentrations that give control of droplet bounce and that the same time adequately controlling deposition. Normally high levels of surfactant need to be included in the formulation to provide any balance of droplet bounce and control of deposition.
- 25 According to the present information we provide a pesticide or herbicide composition that is an aqueous concentrate which is dilutable to form a sprayable formulation comprising, water containing therein,
- 30 i) at least one pesticide or herbicide active and

- ii) a spray drift control agent comprising a spray drift control agent comprising at least one water-soluble cationic polymer that has been formed from ethylenically unsaturated monomers and
- iii) at least one cationic surfactant.

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In a further aspect of the present invention we also provide a method of treating a crop area or an area of land with a pesticide or herbicide which comprises providing an aqueous concentrate, said aqueous concentrate comprises water and containing therein,

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- i) at least one pesticide or herbicide active and
- ii) a spray drift control agent comprising at least one water-soluble cationic polymer that has been formed from ethylenically unsaturated monomers and
- iii) at least one cationic surfactant,

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diluting the aqueous concentrate in a spray tank to form a sprayable formulation and then distributing the sprayable formulation over the crop area or area of land.

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We have found that the inclusion of the water-soluble cationic polymers into the pesticide or herbicide formulation significantly reduces the losses of spray formulation. In addition the composition reduces droplet bounce and maximizes deposition of the pesticide or herbicide, especially when surfactants are included in formulation. This leads to improved efficacy of the pesticide or

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herbicide. We have found that even low levels of said cationic polymer optionally with low levels of surfactant provide excellent control of droplet bounce.

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In general the herbicide or pesticide composition of the present invention is provided as an aqueous concentrate. This aqueous concentrate is then diluted to form the sprayable formulation which can then be sprayed on to a crop area

or an area of land to be treated. Generally the aqueous concentrate is diluted from 20 to 100 fold to give the desired concentration of pesticide and/or herbicide and also the desired concentration of cationic polymer.

- 5 Alternatively the herbicide or pesticide may initially be provided as an aqueous concentrate, optionally containing surfactant, and this is diluted to form the sprayable formulation into which the cationic polymer is combined prior to spraying. The cationic polymers may be added directly to the spray tank, either as an aqueous stock solution, inverse emulsion or as a partially dehydrated
10 polymer dispersion in oil (liquid dispersion product).

- In the composition at least one surfactant is included. The exact choice of surfactant will depend upon the particular pesticide or herbicide used, types of plant, choice of polymer and the location. Effective results are obtained when
15 the surfactant is selected from the group consisting of tallow amine ethoxylates, alkyl phenol ethoxylates, alcohol ethoxylates and alkyl poly glucosides. The at least one surfactant should be cationic or at least a free base amine that can be rendered cationic. A particular preferred surfactant is a tallow amine ethoxylate. Usually the tallow amine ethoxylate is 15 to 20 mole ethoxylate units, although 5
20 mole ethoxylate may also be used. Although tallow amines may be regarded as non-ionic in some circumstances, they will be cationic in the aqueous concentrates of the present invention and diluted aqueous sprayable formulations formed therefrom.

- 25 The surfactant may be added to the formulation in order to improve the efficacy of the pesticide or herbicide. Typically the surfactant improves the penetration of the active ingredient through the leaf surface which is important for systemic herbicides. The surfactant also aids deposition.
- 30 Generally where surfactant is included in the composition, in the aqueous concentrate it would generally be present in an amount of up to 15% by weight

of total composition. Preferably however surfactant will be present in the concentrate at between 4 and 8 or 9%. On dilution the sprayable formulation may contain surfactant in an amount of up to 1100 ppm and preferably between 300 and 750 ppm.

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In order to function effectively as a spray drift control agent the at least one cationic polymer used in the present invention will generally be of a molecular weight sufficient to exert an effect on the spray distribution of the sprayable formulation. That is a polymer of sufficiently high molecular weight to give spray drift control properties and is not a low molecular weight material which would act as a dispersant. Typically the polymer will have a high molecular weight, for instance at least 5 or 6 million and as much as 20 or 30 million or higher. Preferably the cationic polymer will exhibit an intrinsic viscosity of at least 4 dl/g. More preferably the polymer has an intrinsic viscosity of at least 6 dl/g. In this specification intrinsic viscosity is measured by suspended level viscometer at 20°C in 1 M sodium chloride buffered to pH 7.

Intrinsic viscosity of polymers may be determined by preparing an aqueous solution of the polymer (0.5-1% w/w) based on the active content of the polymer. 2 g of this 0.5-1% polymer solution is diluted to 100 ml in a volumetric flask with 50 ml of 2M sodium chloride solution that is buffered to pH 7.0 (using 1.56 g sodium dihydrogen phosphate and 32.26 g disodium hydrogen phosphate per litre of deionised water) and the whole is diluted to the 100 ml mark with deionised water. The intrinsic viscosity of the polymers are measured using a Number 1 suspended level viscometer at 25°C in 1M buffered salt solution.

Preferably IV is at least 8 dl/g, more preferably at least 9 dl/g. It may be up to for instance 30 dl/g but generally it is found that the optimum combination of low viscosity of the composition and spray drift control performance is given by polymers having IV not more than about 20 or 18 dl/g. Preferably IV is not more

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than 16, more preferably not more than 15 dl/g. Particularly preferred IV ranges are from 9 to 13 dl/g, especially 10 to 12 dl/g.

In the present invention the water-soluble polymer is preferably substantially
5 dissolved. That is, the polymer is taken into the solution such that substantially
no visible solid material remains. The polymer may be prepared by
polymerisation of a water soluble monomer or water soluble monomer blend. By
water soluble we mean that the water soluble monomer or water soluble
monomer blend has a solubility in water of at least 5g in 100 ml of water. The
10 polymer may be prepared conveniently by any suitable polymerisation process.

Desirably the polymer may be prepared by reverse phase emulsion
polymerisation, optionally followed by dehydration under reduced pressure and
temperature and often referred to as azeotropic dehydration to form a
15 dispersion of polymer particles in oil. Alternatively the polymer may be provided
in the form of beads by reverse phase suspension polymerisation, or as a
powder by aqueous solution polymerisation followed by comminution, drying
and then grinding. The polymers may be produced as beads by suspension
polymerisation or as a water-in-oil emulsion or dispersion by water-in-oil
20 emulsion polymerisation, for example according to a process defined by EP-A-
150933, EP-A-102760 or EP-A-126528.

The polymer may be branched but in general it is substantially linear and is not
cross-linked.

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The polymer is typically formed from water soluble monomer or monomer blend,
usually water soluble ethylenically unsaturated monomer. The cationic content
i.e. the proportion of cationic monomer in the monomer blend used to form the
polymer, is variable and may be up to 100 wt. %, although more desirably high
30 cationicity polymers are preferred. Typically the cationic polymer would

comprise at least 50 wt. % cationic monomer, more preferably between 60 and 90 wt. %.

5 The high cationicity polymers may be defined in terms of their cationic charge density. Generally it is preferred that the polymers exhibit a cationic charge density of at least 3 mEq/g.

Desirably the cationic polymers may be prepared from at least one cationic ethylenically unsaturated monomer and optionally one or more non-ionic
10 ethylenically unsaturated monomers. Suitable cationic monomers include dialkyl amino alkyl (meth) acrylates (including acid addition salts and quaternary ammonium salts), dialkyl amino alkyl (meth) acrylamides (including acid addition salts and quaternary ammonium salts), di allyl di alkyl ammonium halides. Particularly suitable cationic monomers are for instance the methyl chloride
15 quaternary salt of dimethyl amino ethyl acrylate, the methyl chloride quaternary salt of di methyl amino ethyl methacrylate, acrylamido propyl tri methyl ammonium chloride, methacrylamido propyl tri methyl ammonium chloride and di allyl di methyl ammonium chloride. Nonionic monomers include acrylamide, methacrylamide, 2-hydroxy ethyl acrylate and N-vinylpyrrolidone. One
20 particularly preferred polymer is the copolymer of acrylamide with the methyl chloride quaternary ammonium salt of di methyl amino ethyl acrylate.

When the composition of the present invention is an aqueous concentrate at least one cationic polymer may be present in an amount of between 0.1 and 2%
25 by weight of total composition. Preferably though particularly effective results are found when the amount of cationic polymer or polymers are included in the aqueous concentrate is between 0.5 and 0.9% by weight. When the formulation is diluted to form a sprayable formulation typically the amount of polymer present will be in the range of between 25 and 150 ppm by weight.

Any suitable pesticide or herbicide may be used in the present invention.

Preferred herbicide compositions include at least one systemic herbicide active such as glyphosate or glufosinate. Other examples of herbicides include phenoxy based herbicides such as 2,4-D amine salt. Preferably the systemic
5 herbicide is glyphosate. Generally glyphosate will be present as the iso propyl amine, potassium, ammonium or other salts.

Typically the at least one pesticide or herbicide will be present in the aqueous concentrate in an amount of between 200 and 700 g/l. In the diluted to
10 sprayable formulation the pesticide or herbicide will typically be present in an amount of between 4 and 35 g/l.

Particularly preferred aqueous concentrates comprise glyphosate as the active herbicide ingredients together with water-soluble cationic polymers of
15 acrylamide with cationic monomers such as di methyl amino ethyl acrylate quaternized with methyl chloride. The glyphosate is usually included as the iso propyl amine salt in aqueous solution. The preferred formulation includes surfactant and this is preferably tallow amine ethoxylate (usually 15 to 20 or mole ethoxylate).

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The following examples illustrate invention.

Example 1

A cationic polymer is tested in combination with TAE 20 mole ethoxylate surfactant (AU391) supplied by Adjuvants Unlimited Inc.

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Polymer A is a bead copolymer prepared by suspension polymerisation from a monomer blend containing 80% di methyl amino ethyl acrylate (DMAEA) quaternised with Methyl Chloride and 20% acrylamide with IV 7 dl/g.

- 10 Four test formulations are prepared according to the general recipe below in order to determine the effect of the various constituents:

	Glyphosate IPA technical grade (62% active)	= 66.0 g
	Polymer A	= 0 or 0.75 g
15	Surfactant (TAE 20 mole ethoxylate)	= 0 or 7 g
	Water	= to 100 g

The test formulations above are diluted to a rate of 1% v/v for spraying. This gives an equivalent concentration of active to the rate of use in the field.

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Reduced droplet bounce is demonstrated using the following method.

- A Hewlett Packard 214B Pulse generator connected to a piezo-electric disc connected to a flat tipped glass nozzle with an aperture of 500 μm is used to
25 generate droplets of $\sim 1000 \mu\text{m}$ in diameter. A glass nozzle tip is mounted 20cm above the leaf surface, which is set at an angle of 45°.

- Each droplet is allowed to fall 20cm before impacting with a Pea leaf (variety used is Lincoln pea leaves). The distance that each droplet bounces on impact
30 is recorded and this number is used to determine the percentage of droplets that would be deflected from an average sized leaf (2.5 cm in length). Fifty

droplets are tested for each test solution using 5 individual leaves (10 drops per leaf).

Improved mass deposition/reduced spray drift of the formulations is

- 5 demonstrated by measuring the mass of the composition that is lost when the solution is sprayed under standard conditions.

Test solutions are sprayed into a wind tunnel (dimensions length 3 m, height 2 m, width 2.5 m). A spray head containing a 110° flat fan nozzle is mounted
 10 60cm above the floor. Wind at 6mph (measured at the centre of the tunnel) is generated using a fan placed at a distance of 10 cm behind the nozzle. A 2.5 by 3 m collection device is positioned centrally under the spray head such that a 2.8 m length of the collection device is down wind from the nozzle. The solution to be tested is placed in a brass spray canister attached to the spray head. The
 15 canister, tubing and spray head are then weighed. A source of compressed air is used to expel the test solution through the nozzle at a pressure of 3 bar onto a pre-weighed collection device. The test solutions are sprayed for 135 seconds. The canister and spray head are then reweighed in order to determine the weight of the solution expelled. The collection device is carefully
 20 folded and reweighed. The mass of the solution collected is then calculated and used to determine the loss of spray due to drift. The results are shown in Table 1.

Table 1

Formulation	Bounce (% deflected)	Spray Drift (% loss)
Glyphosate IPA alone	100	8.3
Glyphosate IPA and polymer A	54	7.6
Glyphosate IPA and TAE 20	58	9.8
Glyphosate IPA, TAE 20 and Polymer A	12	7.9

The results show that

- the polymer and the surfactant reduce droplet bounce to a similar extent
- the combination of the surfactant and polymer virtually eliminates droplet bounce
- the inclusion of the surfactant alone increases drift and the mass of active lost
- the inclusion of polymer decreases the amount of spray drift and increases the mass of active deposited

Example 2

Two further polymers are tested as co-formulants within glyphosate formulations either alone or in combination with TAE 20.

Polymer B is a powder grade copolymer prepared by solution polymerisation of a monomer blend containing 80% di methyl amino ethyl acrylate (DMAEA) quaternised with Methyl Chloride and 20% acrylamide with IV 7 dl/g.

Polymer C is a powder grade copolymer containing 90% di methyl amino ethyl acrylate (DMAEA) quaternised with Methyl Chloride and 10% acrylamide with IV 8.5 dl/g.

The test methods described in example 1 are used. The polymers are included at a rate of 0.75% and the surfactant is included at a rate of 7%. All formulations are tested at a dilution rate of 1% v/v. The results are shown in Table 2.

Table 2

Formulation	Bounce (% deflected)	Spray Drift (% loss)
Glyphosate IPA and polymer B	88	7.5
Glyphosate IPA, TAE 20 and Polymer B	6	8
Glyphosate IPA and polymer C	42	8.5
Glyphosate IPA, TAE 20 and Polymer C	2	7.5

The results show that:

- 5
 - the combination of the surfactant and polymer virtually eliminates droplet bounce
 - the inclusion of polymer decreases the amount of spray drift and increases the mass of active deposited

10 Example 3

Two further surfactants are tested in combination with polymer A.

Tallow amine ethoxylate (TAE) 5 mole ethoxylate supplied by Adjuvants

15 Unlimited Inc (AU393)

Alkyl poly glycoside (APG) surfactant supplied by Cognis Inc (Agrimul 2069)

20 The test methods described in example 1 are used. The polymer is included at a rate of 0.75% and the TAE surfactant was included at a rate of 7%, the APG surfactant was included at a rate of 4% on solids (8% as supplied as a 50% concentrate). The results are shown in table 3.

Table 3

Formulation	Bounce (% deflected)	Spray Drift (% loss)
Glyphosate IPA and TAE 5	100	7.8
Glyphosate IPA, TAE 5 and Polymer A	74	6.8
Glyphosate IPA and APG	24	9.5
Glyphosate IPA, APG and Polymer A	8	8.9

The results show that:

- 5
 - the combination of the surfactant and polymer reduces droplet bounce
 - the inclusion of polymer decreases the amount of spray drift and increases the mass of active deposited.

Example 4.

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Two polymers are tested in field trials in combination with a TAE surfactant.

Formulations One and Two contained 7.0% w/w TAE (approximately half the standard commercial inclusion rate for TAE) and 15% w/w TAE (the standard full commercial rate of TAE) respectively, and are tested for comparative purposes.

Formulation Two contains polymer D, a solid grade polyacrylamide homo polymer with an anionic content of 0% and an intrinsic viscosity of 16 dl/g, at a rate of 0.13% w/w with TAE at 7.0% w/w in the final formulation. The level of the polymer in the formulation is limited by the compatibility of the various components in the concentrated herbicide formulation. Results are shown for comparative purposes.

25 Formulation Four contains polymer B at a rate of 0.7% w/w with TAE at 7.0%

w/w in the final formulation.

The above formulations contain 480 g/l glyphosate, present as the isopropyl amine salt. A random plot design of four replicates is treated with 1 pint/acre of each 480 g/l test glyphosate formulation. The plots are assessed visually for the percent control of two weeds: velvet leaf and morning glory. The assessment for morning glory is carried out 20 days post treatment. The plots receive a second treatment of Glyphosate 43 days after the initial treatment in order to control the velvet leaf. The visual assessment for the velvet leaf is carried out 14 days after the second treatment. The results of the trial are summarised below:

Table 4

Formulation	% control		
	Morning Glory	Velvet Leaf	Average
One	64	90	77
Two	71	93	82
Three	71	86	79
Four	69	95	82

The results show that

- The efficacy of the low TAE content formulation is lower than the high TAE content formulation in the absence of polymer
- Polymer D improved the performance of the herbicide over TAE alone at low rate; however the formulation was not as effective as the high rate of TAE
- Polymer C in combination with a low rate of TAE gave similar control to the TAE at full rate

Claims

1. A pesticide or herbicide composition that is an aqueous concentrate which is dilutable to form a sprayable formulation comprising, water containing therein,
 - i) at least one pesticide or herbicide active and
 - ii) a spray drift control agent comprising at least one water-soluble cationic polymer that has been formed from ethylenically unsaturated monomers and
 - iii) at least one cationic surfactant,in which the at least one cationic polymer exhibits an intrinsic viscosity of at least 4 dl/g and the cationic polymer exhibits a cationic charge density of at least 3 mEq/g and/or is formed from at least 50 wt.% cationic monomeric repeating units.
2. A composition according to claim 1 in which the at least one surfactant is present in an amount of up to 15% by weight of total composition.
3. A composition according to claim 1 or 2 in which the cationic polymer is formed from at least one ethylenically unsaturated cationic monomer and optionally a nonionic monomer.
4. A composition according to any of claims 1 to 3 which is an aqueous concentrate in which at least one cationic polymer is present in an amount of between 0.1 and 2% by weight of total composition.
5. A composition according to any of claims 1 to 3 which is a sprayable formulation in which at least one cationic polymer is present in an amount of between 25 and 150 ppm by weight of total composition.
6. A composition according to any of claims 1 to 5 comprising at least one systemic herbicide active, preferably glyphosate.
7. A composition according to any of claims 1 to 4 or 6 in which at least one pesticide or herbicide is present in an amount of between 50 and 700 g/l.

8. A method of treating a crop area or an area of land with a pesticide or herbicide which comprises providing an aqueous concentrate said aqueous concentrate comprises water and containing therein,

- i) at least one pesticide or herbicide active and
- ii) a spray drift control agent comprising at least one water-soluble cationic polymer that has been formed from ethylenically unsaturated monomers and
- iii) at least one cationic surfactant,

diluting the aqueous concentrate in a spray tank to form a sprayable formulation and then distributing the sprayable formulation over the crop area or area of land, in which the at least one cationic polymer exhibits an intrinsic viscosity of at least 4 dl/g and the cationic polymer exhibits a cationic charge density of at least 3 mEq/g and/or is formed from at least 50 wt.% cationic monomeric repeating units.