

METHOD FOR THE PURIFICATION OF TRIORGANOPHOSPHITES BY TREATMENT WITH A BASIC ADDITIVE

CROSS-REFERENCE TO RELATED APPLICATIONS

5 This application claims benefit of priority from Provisional Application No. 60/783,462, filed March 17, 2006. This application hereby incorporates by reference Provisional Application No. 60/783,462 in its entirety.

FIELD OF THE INVENTION

10 The invention relates to the field of preparation and use of triorganophosphite compounds. More specifically, the invention relates to the partial removal of phosphorus acids in triorganophosphite product streams by treatment with a basic additive. Because the phosphorus acids can act as catalysts for triorganophosphite hydrolysis, the treated triorganophosphite
15 product streams can be stabilized against further hydrolysis when exposed to water.

BACKGROUND OF THE INVENTION

Triorganophosphites of the general structure $(R^4O)(R^5O)P(OR^6)$ and
20 $((R^7O)(R^8O)PO)_nA$, where "A" is an optionally substituted or unsubstituted aliphatic, aromatic, or heteroaromatic radical and n is an integer greater than 1, are used in a number of important commercial applications including their use as antioxidants, stabilizers, anti-wear additives and as ligands for various catalytic processes. Generally, triorganophosphites are produced from PX_3 (X = Cl, Br, or
25 I) and the corresponding alcohols (ROH). This reaction occurs stepwise by displacement of X with OR. When X is Cl, the process can form phosphorodichloridite $(R^4O)PCl_2$ and phosphorochloridite $(R^4O)(R^5O)PCl$ intermediates, triorganophosphites $(R^4O)(R^5O)P(OR^6)$ and acid HX.

Several methods for making organophosphites, for example those
30 described in Houben-Weyl, Bd. XXII/2 pages 12-17, G. Thieme Verlag, Stuttgart 1964, and supplement E1, pages 413-421 Stuttgart, New York 1982, are known using readily available PCl_3 and the corresponding alcohols. The acid HX can be

removed by physical separation or by acid-base reaction using organic or inorganic bases. In addition, US Patent No. 6,069,267, and US Patent No. 6,031,120 describe the use of triorganoamines to remove HCl followed by water washing at low temperatures to remove the corresponding ammonium hydrochloride salts from the triorganophosphite mixture.

Houben-Weyl, Bd. XXII/2 Chapter I and pages 30-32, G. Thieme Verlag, Stuttgart 1964 teaches that triorganophosphites are easily hydrolyzed in the presence of water, especially in the presence of acidic compounds, to produce one or more of the compounds diorganohydrogenphosphite $(R^2O)(R^3O)POH$, organodihydrogenphosphite $(R^1O)(HO)PO(H)$, and phosphorous acid H_3PO_3 . Gerard, Hudson and Healy, respectively, (in G.M. Kosolapoff & L. Maier Eds., Organic Phosphorous Compounds, Vol. 5 pages 41-42, Wiley & Sons., New York, 1973; Healy et al. J. Inorg. Nucl. Chem., 1974, 36, 2579) teach that this hydrolysis reaction is autocatalytic due to the acidic properties of diorganohydrogenphosphite $(R^2O)(R^3O)POH$, organodihydrogenphosphite $(R^1O)(OH)PO(H)$ and H_3PO_3 . Hydrolysis under basic conditions was found to be slower than hydrolysis in the presence of acids (Westheimer et al., J. Amer. Chem. Soc. 1988, 110, 183). Without the removal of these acidic hydrolysis products, as a result of hydrolysis, there can be significant degradation and loss of the triorganophosphite product during downstream processing and storage. In order to at least partially address this effect, US Patent No. 3,553,298 teaches that nitrogen-containing compound additives such as amines and magnesium oxide can partially stabilize the triorganophosphites by retarding hydrolysis. However, such additives can cause undesirable effects or be incompatible with the use of the triorganophosphites, for example during preparation of a transition metal-triorganophosphite catalyst or catalyst precursor for reactions such as hydrocyanation and hydroformylation, for example. It would, therefore, be desirable to have a method to stabilize triorganophosphites without using additives that may provide undesirable side effects.

US Patent No. 6,069,267 Example 1 discloses a sequential treatment of a triorganophosphite reaction mixture in an organic solvent with 0.1 N aqueous HCl, 0.1 N aqueous NaOH, followed by distilled water.

US Patent No. 6,844,289 discloses a process for combining a crude ligand mixture, prepared for example by the process of US Patent No. 6,069,267, with divalent nickel compounds and reducing agents to produce a catalyst which is a complex of nickel and a bidentate phosphorus compound. This patent discloses that such crude ligand mixture can contain byproducts which may affect the rate of formation of the nickel-containing catalyst. Disclosed therein are treatment methods which include contacting crude ligand mixture with one or more of, for example, a weakly basic organic resin and a two phase solvent system for liquid-liquid extraction. The patent discloses that various treatments of crude ligand may overcome deleterious rate inhibiting effects of byproduct impurities which are present in the crude ligand.

SUMMARY OF THE INVENTION

As described above, triorganophosphites can be hydrolyzed in basic as well as acidic conditions, with base-catalyzed hydrolysis being slower than acid-catalyzed hydrolysis. In this regard, it has been discovered that hydrolysis products can be removed by treatment with a strong base, so that the overall degradation of triorganophosphite may be decreased.

The present invention includes a method for separating one or more triorganophosphite components from a crude phosphite mixture containing acidic hydrolysis products by contacting the crude phosphite mixture with a basic additive to produce a second mixture comprising a first phase and a second phase. The first phase comprises the basic additive and one or more components independently selected from the group consisting of $(R^2O)(R^3O)POH$, $(R^1O)(HO)PO(H)$ and H_3PO_3 wherein R^1 , R^2 and R^3 are independently selected from the group consisting of C_1 to C_{18} alkyl, C_6 to C_{18} aryl and hydroxyaryl, and C_3 to C_{18} cycloalkyl and hydroxyalkyl radicals, and wherein R^2 and R^3 can optionally be connected to each other directly by a chemical bond

or through an intermediate divalent group R^9 . R^9 is selected from the group consisting -O-, -S-, and -CH(R^{12})-, wherein R^{12} is selected from the group comprising H, C_6 to C_{18} aryl, and C_1 to C_{18} alkyl.

The second phase comprises one or more triorganophosphite
5 components independently selected from the group consisting of
(R^4O)(R^5O)P(OR^6) and ((R^7O)(R^8O)PO) $_n$ A, wherein R^4 , R^5 , R^6 , R^7 and R^8 are
independently selected from the group consisting of C_1 to C_{18} alkyl, C_6 to C_{18} aryl
and C_3 to C_{18} cycloalkyl radicals, and wherein each R^4 , R^5 and R^6 can optionally
be connected to one or both of the other two directly by a chemical bond or
10 through an intermediate divalent group R^{10} , and R^7 and R^8 can optionally be
connected to each other directly by a chemical bond or through an intermediate
divalent group R^{11} . The intermediate divalent groups R^{10} and R^{11} are
independently selected from the group consisting of -O-, -S-, and -CH(R^{12})-,
wherein R^{12} is selected from the group consisting of H, C_6 to C_{18} aryl, and C_1 to
15 C_{18} alkyl. The substituent "A" is an optionally substituted or unsubstituted
aliphatic, aromatic, or heteroaromatic radical and n is an integer greater than 1.
Examples of A include C_1 to C_{18} aliphatic, C_6 to C_{28} aromatic, or C_3 to C_{28}
heteroaromatic radicals.

The basic additive can comprise at least one compound selected from the
20 group consisting of NaOH, KOH, Na_2CO_3 , K_2CO_3 , $Ca(OH)_2$, NH_4OH , $CaCO_3$, a
strongly basic anion-exchange resin, and combinations thereof. In one
embodiment, the basic additive comprises a strongly basic anion-exchange resin,
for example a strongly basic anion-exchange resin comprising polymer-bound
tetraorgano-ammonium hydroxide groups.

25 In one embodiment of the present invention, the basic additive and the
crude phosphite mixture are contacted at temperatures above about $-25\text{ }^\circ\text{C}$, for
example at temperatures from about $-10\text{ }^\circ\text{C}$ to about $30\text{ }^\circ\text{C}$.

In yet another embodiment of the present invention, the first phase is
maintained at or above a pH of about 13.5 throughout the contacting process.

30 The present invention also relates to a method for separating
triorganophosphite components from a crude phosphite mixture containing acidic

hydrolysis products which further comprises the step of contacting the crude phosphite mixture with water before contacting the crude phosphite mixture with the basic additive.

The present invention also relates to methods for separating
5 triorganophosphite components from a crude phosphite mixture containing acidic hydrolysis products wherein the methods further comprise the steps of separating the first phase from the second phase, and contacting the second phase with the basic additive. In another embodiment the present invention includes contacting the second phase with a brine solution after contacting with
10 the basic additive.

Another embodiment of the present invention further comprises contacting the second phase with a transition metal or a transition metal compound to produce a transition metal-triorganophosphite catalyst or catalyst precursor. Examples of transition metals or transition metal compounds that can be used
15 include $\text{Ni}(\text{COD})_2$ (COD is 1,5-cyclooctadiene), $\text{Ni}[\text{P}(\text{O}-o\text{-C}_6\text{H}_4\text{CH}_3)_3]_3$ and $\text{Ni}[\text{P}(\text{O}-o\text{-C}_6\text{H}_4\text{CH}_3)_3]_2(\text{C}_2\text{H}_4)$, all of which are known in the art. The resulting transition metal-triorganophosphite catalyst or catalyst precursor can be useful for reactions such as hydrocyanation or hydroformylation, for example.

The invention also relates to a method for preparing triorganophosphites,
20 which comprises contacting an alcohol with PCl_3 in the presence of a triorganoamine to produce a first reaction product comprising one or more organophosphites and triorganoamine hydrochloride; removing the triorganoamine hydrochloride from the first reaction product to produce a second reaction product optionally containing one or more triorganophosphites,
25 diorganohydrogenphosphite $(\text{R}^2\text{O})(\text{R}^3\text{O})\text{POH}$, organodihydrogenphosphite $(\text{R}^1\text{O})(\text{HO})\text{PO}(\text{H})$ and H_3PO_3 , wherein R^1 , R^2 and R^3 are independently selected from the group consisting of C_1 to C_{18} alkyl, C_6 to C_{18} aryl and hydroxyaryl, and C_3 to C_{18} cycloalkyl and hydroxyalkyl radicals, and wherein R^2 and R^3 can optionally be connected to each other directly by a chemical bond or through an
30 intermediate divalent group R^9 , wherein R^9 is selected from the group consisting of -O-, -S-, and -CH(R^{12})-, wherein R^{12} is selected from the group consisting of H,

C₆ to C₁₈ aryl, and C₁ to C₁₈ alkyl; and removing at least a portion of one or more of the compounds diorganohydrogenphosphite (R²O)(R³O)POH, organodihydrogenphosphite (R¹O)(HO)PO(H) and H₃PO₃ by contacting the second reaction product with a basic additive.

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BRIEF DESCRIPTION OF THE FIGURE

FIG. 1 is a graphical representation showing the fraction of hydrolysis products remaining in the organic phase after contact with a control (brine) as compared to contact with a basic additive.

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DETAILED DESCRIPTION OF THE INVENTION

The present invention includes methods for purifying and stabilizing triorganophosphites by removing acidic impurities from the phase containing the triorganophosphites. For example, acidic impurities such as diorganohydrogenphosphite (R²O)(R³O)POH, organodihydrogenphosphite (R¹O)(HO)PO(H), and H₃PO₃ can occur as a result of aqueous workup of crude triorganophosphites, such as (R⁴O)(R⁵O)P(OR⁶), and ((R⁷O)(R⁸O)PO)_nA, and from unconverted phosphorodichloridite (R⁴O)PCl₂ and phosphorochloridite (R⁴O)(R⁵O)PCl intermediates used as components for the making of triorganophosphites. Furthermore, small amounts of the above-described acidic impurities can increase in concentration during storage of triorganophosphite materials by autocatalytic hydrolysis upon exposure to moisture, causing yield loss and undesirable effects with the use of triorganophosphites in applications such as catalysis. At least one of these disadvantages may be overcome by removal of acidic impurities by means of extraction with a basic additive.

In at least one embodiment falling within the scope of the present invention, a crude phosphite mixture comprising at least one triorganophosphite selected from the group consisting of (R⁴O)(R⁵O)P(OR⁶), and ((R⁷O)(R⁸O)PO)_nA; and one or more of the compounds diorganohydrogenphosphite (R²O)(R³O)POH, organodihydrogenphosphite (R¹O)(HO)PO(H), and H₃PO₃ is contacted with a basic additive to produce a second mixture. The second mixture

comprises two phases - a first phase and a second phase. The first phase comprises the basic additive and at least a portion of one or more of the components consisting of diorganohydrogenphosphite $(R^2O)(R^3O)POH$, organodihydrogenphosphite $(R^1O)(HO)PO(H)$, and H_3PO_3 . The second phase
5 comprises at least a portion of the organophosphites. The first phase of the second mixture can optionally be separated from the second phase, using general separation methods known by those skilled in the art. For example, if the basic phase is liquid, such as aqueous NaOH, a conventional mixer-settler can be used to contact the crude phosphite mixture and aqueous NaOH then
10 separate the second phase comprising the triorganophosphites from the first phase comprising the acidic impurities. If, instead, a basic ion-exchange resin is deployed, a decanter or filter may be used.

R^1 , R^2 , and R^3 are independently selected from the group consisting of C_1 to C_{18} alkyl, C_6 to C_{18} aryl and hydroxyaryl, and C_3 to C_{18} cycloalkyl and
15 hydroxyalkyl radicals; R^2 and R^3 can optionally be connected to each other directly by a chemical bond or through an intermediate divalent group R^9 . R^9 is selected from the group consisting of -O-, -S-, and -CH(R^{12})-, wherein R^{12} is selected from the group consisting of H, C_6 to C_{18} aryl, or C_1 to C_{18} alkyl.

R^4 , R^5 , R^6 , R^7 , and R^8 are independently selected from the group
20 consisting of C_1 to C_{18} alkyl, C_6 to C_{18} aryl, and C_3 to C_{18} cycloalkyl radicals. Each of R^4 , R^5 , and R^6 can optionally be connected to one or both of the other two directly by a chemical bond or through an intermediate divalent group R^{10} . R^7 and R^8 can optionally be connected to each other directly by a chemical bond or through an intermediate divalent group R^{11} . R^{10} and R^{11} are independently
25 selected from the group consisting of -O-, -S-, and -CH(R^{12})-, wherein R^{12} is selected from the group consisting of H, C_6 to C_{18} aryl, and C_1 to C_{18} alkyl. The substituent "A" is an optionally substituted or unsubstituted aliphatic, aromatic or heteroaromatic radical where n is an integer greater than 1.

The basic additive employed in the present invention can comprise one or
30 more compositions comprising at least one basic compound and/or material. The basic additive may, for example, comprise at least one compound selected

from the group consisting of NaOH, KOH, Na₂CO₃, K₂CO₃, Ca(OH)₂, NH₄OH, CaCO₃, a strongly basic anion-exchange resin, and combinations thereof. Examples of suitable strongly basic ion-exchange resins are resins comprising polymer-bound tetraorgano-ammonium hydroxide groups.

5 Since hydrolysis rates are dependent on temperature, the extraction temperature should ideally be as low as possible only restricted by physical limits. In the case of liquid-liquid extractions, a physical limit impacting the range of suitable operation temperatures is, for example, the freezing point of the liquid phases. Since under commercial operation the temperature range achievable is
10 subject to the limits of the equipment available, temperatures above about -25 °C are generally preferred. An example of a non-limiting temperature range for this process is from about -10 °C to about 30 °C.

 Ideally, the pH of the basic phase should remain at or above about 13.5 throughout the extraction process. This can be accomplished by using a 1 normal
15 solution of a strong base in water or a strongly basic anion-exchange resin of equivalent strength. If the production of the triorganophosphite generates a salt constituting a weak acid such as a triorganoamine hydrochloride, the weak acid should be removed first to allow for an effective extraction process of the acidic diorganohydrogenphosphite (R²O)(R³O)POH, organodihydrogenphosphite
20 (R¹O)(HO)PO(H), and H₃PO₃ impurities with the basic additive. Triorganoamine hydrochloride salt removal can be achieved by filtration and/or by water extraction prior to contacting with the basic additive as, for example, described in US Patent Nos. 6,069,267, and 6,031,120.

 The total amount of molar equivalents of base to be used is, at least in
25 part, a function of the total amount of molar equivalents of acidic diorganohydrogenphosphite (R²O)(R³O)POH, organodihydrogenphosphite (R¹O)(HO)PO(H), and H₃PO₃ impurities that are present prior to the treatment with the basic additive. For example, a 10- to 1000-fold excess of the base can be suitable, provided that, in the case of a solution extraction, a minimum pH of
30 about 13.5 of the basic solution is maintained throughout the extraction process, or, in the case of a resin treatment, an equivalent basicity is maintained.

The volume of the basic phase can be determined by practical limitations, such as reactor size and separation capacity, by the pH limits, and by molar equivalent ratios of base to acidic impurities. For instance, in the case of an extraction using aqueous NaOH, the volume of the basic first phase can, for example, be between one-tenth and one-third of the product second phase volume. The amount of NaOH or polymer-bound tetraorgano-ammonium hydroxide can, for example, be between about 0.05 and about 1 equivalent of the total amount of phosphorus in the triorganophosphite solution phase.

The contact time is, at least in part, a function of scale, mixing equipment, and hydrolysis rates of a specific organophosphite under conditions of basic extraction, and can, for example, be from about 15 minutes to about 5 hours.

For many processes using a basic additive, one or two extractions using a 10-fold excess of base to acid at a pH greater than 13 for about 15 to about 30 minutes should be sufficient. In the case of a treatment with a strongly basic anion-exchange resin comprising polymer-bound tetraorgano-ammonium hydroxide groups, the phase containing the phosphorus compounds can be mixed with a predetermined amount of the resin or the foresaid phase can be passed through a column containing the resin. The amount of resin used is, at least in part, a function of the amount of acidic impurities to be removed. For example, when using a strongly basic anion-exchange resin, a one time treatment with an amount of resin equivalent to a 6-10 fold excess of base to acid may be sufficient. The resin may be regenerated using a basic aqueous solution of NaOH as known by those skilled in the art.

Typically, the extraction process is relatively insensitive to pressure and, in this regard, is usually limited only by practical considerations. For practical reasons, preferred reaction pressure ranges can, for example, be from about 10 psia (69 kPa) to about 50 psia (345 kPa).

In the case of a liquid-liquid extraction, the basic solution may contain a soluble salt, such as NaCl, to aid the separation process.

To avoid further triorganophosphite hydrolysis, contact with the basic additive should ideally be done soon after the production of the crude phosphite

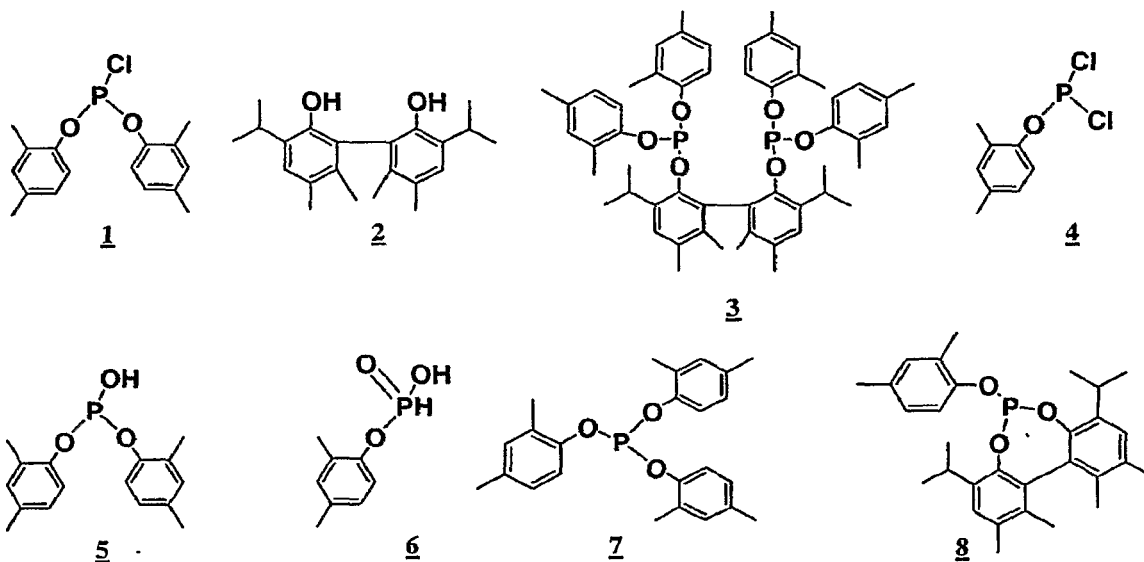
mixture and preferably before further concentration and or heat exposure of the crude phosphite product mixture.

Acidic compounds to be removed may include, but are not restricted to, diorganohydrogenphosphite ($R^2O)(R^3O)POH$, organodihydrogenphosphite
 5 ($R^1O)(HO)PO(H)$, H_3PO_3 , aromatic alcohols, and ammoniumhydrochlorides and ammoniumhydrobromides.

Examples

The following non-limiting examples further illustrate embodiments of the
 10 present invention.

Examples of the invention were prepared from crude phosphite mixtures containing one or more components having the following chemical structures: diorganohydrogenphosphite 5, organodihydrogenphosphite 6, monodentate
 15 triorganophosphites 7 and 8, phosphorochloridite 1, phosphorodichloridite 4, biphenol 2, and bidentate triorganophosphite 3.



The following example shows that acidic diorganohydrogenphosphite 5,
 20 organodihydrogenphosphite 6, and H_3PO_3 representing primary hydrolysis products of triorganophosphites 7, phosphorochloridite 1, and

phosphorodichloridite 4 can be significantly reduced or quantitatively removed from a solution containing the monodentate and bidentate triorganophosphites 7, 8, and 3 by extraction with an aqueous solution of a strong base. Results are summarized in Table 1 and reported graphically in FIG. 1. For quantitative NMR
5 analysis, 0.250 mL of the organic phase and 0.600 mL of a 0.01 molar triphenylphosphine oxide (TPPO) in C₆D₆ were combined and analyzed by ³¹P NMR (T1 = 5 sec.). Unless stated otherwise, the term brine refers to a saturated aqueous NaCl solution (about 26 wt% NaCl).

10 **Example A**

Synthesis of a Crude Phosphorochloridite 1 Containing Reaction Mixture:

A temperature controlled 2000 mL baffled flask equipped with an overhead stirrer was charged with 400 mL of 1.0 molar PCl₃ in toluene and 4.0
15 mL 2.0 molar triethylamine in toluene. Under vigorous stirring, a solution of 411 mL 2.0 molar triethylamine in toluene and a solution of 411 mL 2.0 molar 2,4-xylenol in toluene were added separately and concurrently via peristaltic pump at 2.3 mL/min. During the addition period, the reaction temperature was maintained at 5 °C. The ³¹P NMR analysis exhibited transformation to the corresponding
20 phosphorochloridite 1 (³¹P NMR chemical shift 162 ppm) in 90% selectivity. Other phosphorus containing components were triorganophosphite 7 (9%) and phosphorodichloridite 4 (1%) representing the crude solution of phosphorochloridite 1 used in the Water Extraction (Control Experiment) as well as Examples A1 and A2.

Water Extraction (Control Experiment):

Under nitrogen, 25 mL of the crude solution of phosphorochloridite 1 in toluene (0.56 molar) was combined with 14 mmol of triethylamine (Et₃N) and 10 mL of water (H₂O). The mixture was stirred vigorously for 5 minutes before the
30 two phases were separated. ³¹P NMR analysis of the organic phase indicated complete hydrolysis of phosphorochloridite 1 (162 ppm) to a mixture of the

corresponding diorganohydrogenphosphite 5, organodihydrogenphosphite 6, and traces of H_3PO_3 (^{31}P NMR chemical shifts -2 to + 2 ppm). Subsequently, the organic phase was contacted with two successive portions of 10 mL 13 wt% brine and one portion of 10 mL 26 wt% brine by vigorous mixing for 5 minutes each, and subsequent separation of the two phases. Quantitative ^{31}P NMR analysis of the organic phase was conducted after each treatment with brine, but no significant reduction of hydrolysis products in the organic phase was observed at any time.

Example A1 (Aqueous KOH Treatment):

Under nitrogen, 25 mL of a 0.56 molar solution of phosphorochloridite 1 in toluene was combined with 14 mmol of Et_3N and 10 mL of H_2O . The mixture was stirred vigorously for 5 minutes before the two phases were separated. ^{31}P NMR analysis of the organic phase indicated complete hydrolysis of phosphorochloridite 1 (^{31}P NMR chemical shift 162 ppm) to a mixture of the corresponding diorganohydrogenphosphite 5, organodihydrogenphosphite 6, and traces of H_3PO_3 (^{31}P NMR chemical shift -2 to + 2 ppm). Subsequently, the organic phase was contacted with two successive portions of 10 mL 1 molar aqueous KOH and one portion of 10 mL 26 wt% brine by vigorous mixing for 5 minutes each, and subsequent separation of the two phases. Quantitative ^{31}P NMR analysis of the organic phase after each treatment with aqueous KOH was conducted indicating a significant reduction of hydrolysis products by the aqueous KOH treatment.

Example A2 (Aqueous NaOH Treatment):

Under nitrogen, 25 mL of a 0.56 molar solution of phosphorochloridite 1 in toluene were combined with 14 mmol of Et_3N and 10 mL of H_2O . The mixture was stirred vigorously for 5 minutes before the two phases were separated. ^{31}P NMR analysis of the organic phase indicated complete hydrolysis of phosphorochloridite 1 to a mixture of the corresponding diorganohydrogenphosphite 5, organodihydrogenphosphite 6, and traces of

H₃PO₃ (³¹P NMR chemical shift -2 to + 2 ppm). Subsequently, the organic phase was contacted with two successive portions of 10 mL 1 molar aqueous NaOH and one portion of 10 mL 26 wt% brine by vigorous mixing for 5 minutes each, and subsequent separation of the two phases. Quantitative ³¹P NMR analysis of the organic phase after each treatment was conducted indicating quantitative removal of hydrolysis products by the aqueous NaOH treatment.

Table 1

Fraction of Acidic Hydrolysis Products Diorganohydrogenphosphite 5, Organodihydrogenphosphite 6, and H₃PO₃ Remaining in the Organic Phase as Determined by ³¹P NMR.

Experiment	Step / Treatment	Remaining fraction of acidic hydrolysis products in organic phase (% mol)
Water Extraction(control)	10 mL water, 8 mmol	100%
	Et ₃ N	
	10 mL 13 wt% brine	99%
	10 mL 13 wt% brine	98%
	10 mL 26wt% brine	100%
Example A1	10 mL water, 8 mmol	100%
	Et ₃ N	
	10 mL 1 M KOH	46%
	10 mL 1 M KOH	7%
	10 mL 26 wt% brine	5%
Example A2	10 mL water, 8 mmol	100%
	Et ₃ N	
	10 mL 1 M NaOH	34%
	10 mL 1 M NaOH	4%
	10 mL 26 wt% brine	0%

FIG. 1 illustrates the results from Table 1 in graphical form.

The following Example and Table 2 show the benefit of strongly basic anion-exchange resin treatment for the purity and stability of the crude bidentate triorganophosphite **3** under exposure to water over an extended period of time, for example, under certain storage conditions.

AmberlystTM A26-OH (Rohm and Haas Company) is a strong base, type 1, anionic, macroreticular polymeric resin based on crosslinked styrene divinylbenzene copolymer containing quaternary ammonium hydroxide groups; d = 675 g/L, concentration of active sites = 0.80 equiv/L.

Example B

Synthesis of a Crude Bidentate Triorganophosphite **3 Reaction Mixture:**

A temperature controlled 500 mL baffled flask equipped with an overhead stirrer was charged with 200 mL of 1.0 molar PCl_3 in toluene. Under vigorous stirring, a solution of 203.6 mL 2.0 molar triethylamine in toluene and a solution of 203.6 mL 2.0 molar 2,4-xyleneol in toluene were added separately and concurrently via peristaltic pump at 2.9 mL/min. During the addition period, the reaction temperature was maintained at 5 °C. The ^{31}P NMR analysis exhibited transformation to the corresponding phosphorochloridite **1** (^{31}P NMR chemical shift 162 ppm) in 90% selectivity. Under vigorous stirring, 252.8 mmol of Et_3N was added at 20 °C, followed by a solution of 84.6 mL of 1.0 molar biphenol **2** via a peristaltic pump at 2 mL/min. The reaction mixture was vigorously contacted with 150 mL H_2O . Quantitative ^{31}P NMR analysis of the organic phase revealed a phosphorus distribution of 81 mol% bidentate triorganophosphite **3**, 4 mol% monodentate triorganophosphite **8**, 12 mol% monodentate triorganophosphite **7**, and hydrolysis products **5**, **6**, and H_3PO_3 equaling 3 mol % of the total phosphorus content. This solution represents the crude bidentate triorganophosphite **3** reaction mixture used in the Brine Treatment (Control Experiment) and Example B1.

Brine Treatment (Control Experiment):

About 200 mL of the crude bidentate triorganophosphite 3 reaction mixture representing 67 mmol phosphorus was contacted with 50 mL of 26 wt% brine. Approximately 20% of the solvent was removed in vacuo at 40 °C over 30
5 minutes. The remaining solvent was removed in vacuo at 55 °C over one hour. Quantitative ^{31}P NMR analysis of the organic phase was conducted after the contact with brine, and of the concentrated sample, but no significant reduction of hydrolysis products in the organic phase was observed at any time.

The crude mixture was diluted with toluene to about 80 mL resulting in a
10 33 wt% mixture. To 40 mL of this mixture, representing about 34 mmol phosphorus, was added 0.4 mL of water and the mixture stirred vigorously for 6 days while monitoring the content of phosphorus compounds by ^{31}P NMR. Quantitative ^{31}P NMR analysis of the organic layer was conducted after 2 hours, 1 day, 2 days, 3 days, and 6 days. Results are summarized in Table 2 and
15 indicate relatively fast hydrolysis of all the triorganophosphites.

Example B1 (Resin Treatment):

About 200 mL of the crude bidentate triorganophosphite 3 reaction mixture representing 67 mmol phosphorus was stirred with 11.5 g (13.6 mmol hydroxy
20 anion equivalent) of AmberlystTM A26-OH resin for two days. After the resin was removed, approximately 20% of the solvent was evaporated in vacuo at 40 °C over 30 minutes. The remaining solvent was removed in vacuo at 55 °C over one hour. Quantitative ^{31}P NMR analysis of the organic layer was conducted after treating with resin for two hours and two days. The results summarized in
25 Table 2 indicate that all hydrolysis products were removed quantitatively after 2 hours of treatment, and no hydrolysis of the triorganophosphites was observed after prolonged treatment with resin over 2 days.

The crude mixture was diluted with toluene to about 80 mL resulting in a
30 33 wt% mixture. To 40 mL of this mixture, representing about 34 mmol phosphorus, was added 0.4 mL of water and the mixture stirred vigorously for 13 days. Quantitative ^{31}P NMR analysis of the organic layer was conducted after 2

hours, 1 day, 2 days, 3 days, 6 days and 13 days. In contrast to the control experiment without AmberlystTM A26-OH resin treatment, no significant increase in hydrolysis products was observed at any time.

Table 2

Fraction of Acidic Hydrolysis Products Diorganohydrogenphosphite 5,
Organodihydrogenphosphite 6, and H_3PO_3 Compared to Bidentate
Triorganophosphite 3 Remaining in the Organic Phase
as Determined by ^{31}P NMR.

Experiment	Step / Treatment	Acidic hydrolysis Products (mol % Phosphorus)	Bidentate Triorganophosphite <u>3</u> (mol % Phosphorus)
Brine Experiment (control)	H ₂ O wash	3	81
	Brine treatment	3	81
	After concentrating	3	81
	2 hours with H ₂ O	4	81
	1 day with H ₂ O	5	80
	2 days with H ₂ O	11	75
	3 days with H ₂ O	64	16
	6 days with H ₂ O	66	15
Example B1	H ₂ O wash	3	81
	Resin 2 hours	0	84
	Resin 2 days	0	84
	2 hours with H ₂ O	0	85
	1 day with H ₂ O	0	85
	2 days with H ₂ O	0	85
	3 days with H ₂ O	0	85
	6 days with H ₂ O	0	85
	13 days with H ₂ O	0	85

Additional Examples

The following Examples demonstrate the removal of 3 mol% of total phosphorus and 6 mol% of total phosphorus, respectively, of acidic hydrolysis products diorganohydrogenphosphite 5, organodihydrogenphosphite 6, and H_3PO_3 by treatment with aqueous NaOH during the workup of a crude bidentate triorganophosphite 3.

Example 1

Synthesis of a Crude Bidentate Triorganophosphite 3 Reaction Mixture:

A temperature controlled 2000 mL baffled flask equipped with an overhead stirrer was charged with 400 mL of 1.0 molar PCl_3 in toluene and 4.0 mL 2.0 molar triethylamine in toluene. Under vigorous stirring, a solution of 411 mL 2.0 molar triethylamine in toluene and a solution of 411 mL 2.0 molar 2,4-xilenol in toluene were added separately and concurrently via a peristaltic pump at 2.3 mL/min. During the addition period, the reaction temperature was maintained at 5 °C. The ^{31}P NMR analysis exhibited transformation to the corresponding phosphorochloridite 1 (^{31}P NMR chemical shift 162 ppm) in 90% selectivity. Under vigorous stirring, 600 mmol of Et_3N were added at 20 °C followed by a solution of 87.9 mL of 1.0 molar biphenol 2 via a peristaltic pump at a rate of 1.2 mL/min.

Workup:

After the completion of reaction, the reaction mixture was contacted with 300 mL of 13 wt% brine under vigorous stirring to remove triethylamine hydrochloride salts. Quantitative ^{31}P NMR of the organic phase (Table 3) indicated hydrolysis products (^{31}P NMR chemical shift -2 to +2 ppm), equivalent to 3 mol % of the total phosphorus content. After separating, the organic phase was contacted with 300 mL of 20 wt% brine containing 8 mmol of NaOH (pH 12) followed by 300 mL of 26 wt% brine. Quantitative ^{31}P NMR analysis of the organic phase indicated hydrolysis products equivalent to 3 mol % of total phosphorus (^{31}P NMR chemical shift -2 to +2 ppm) indicating that the amount of base used was not sufficient to remove the acidic impurities. Subsequently, the organic phase was contacted successively with 300 mL of 1 molar NaOH (pH 14) and 300 mL 26 wt% brine. Quantitative ^{31}P NMR analysis of the organic phase indicated quantitative removal of the hydrolysis products. The molar distribution of phosphorus in the organic phase was 82% bidentate triorganophosphite 3 and 18 % monodentate triorganophosphites 7 and 8 at ^{31}P NMR chemical shifts of 133, 132, and 136 ppm, respectively.

Table 3

Phosphorus Distribution During Aqueous Workup with NaOH (Example 1)

Step / Treatment	Acidic Hydrolysis Products (mol % Phosphorus)	Bidentate Triorganophosphite <u>3</u> (mol % Phosphorus)
13 % wt. brine	3	78
20 % wt. Brine + 8 mmol NaOH followed by 26 % wt. brine	2	79
1 M NaOH followed by 26 % wt. brine	0	82

5 Example 2**Synthesis of a Crude Bidentate Triorganophosphite 3 Reaction Mixture:**

A temperature controlled 500 mL baffled flask equipped with an overhead stirrer was charged with 100 mL of 1.0 molar PCl_3 in toluene and 1.0 mL 2.0 molar triethylamine in toluene. Under vigorous stirring, a solution of 100 mL 2.0 molar triethylamine in toluene and a solution of 100 mL 2.0 molar 2,4-xylenol in toluene were added separately and concurrently via a peristaltic pump at 2.9 mL/min. During the addition period, the reaction temperature was maintained at 45 °C. The ^{31}P NMR analysis exhibited transformation to the corresponding phosphorochloridite 1 (^{31}P NMR chemical shift 162 ppm) in 89% selectivity. Under vigorous stirring, 130 mmol of Et_3N was added at 20 °C followed by a solution of 37.5 mL of 1.0 molar biphenol 2 via a peristaltic pump at 2 mL/min.

Workup:

The reaction mixture was vigorously contacted with 75 mL H_2O . Quantitative ^{31}P NMR analysis of the organic phase indicated hydrolysis products equaling 6 mol % of the total phosphorus content (Table 4). After separating the aqueous phase, the organic phase was contacted successively with two portions of 75 mL 1 molar aqueous NaOH followed by 75 mL of 26 wt% brine. Quantitative ^{31}P NMR analysis of the organic phase indicated quantitative removal of hydrolysis products. The molar distribution of phosphorus in the

organic phase was 76 % bidentate triorganophosphite 3 and 23 % monodentate triorganophosphites 7 and 8 at ^{31}P NMR chemical shifts of 133, 132, and 136 ppm, respectively.

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Table 4

Phosphorus Distribution During Aqueous Workup with NaOH (Example 2)

Step / Treatment	Acidic Hydrolysis Products (mol % Phosphorus)	Bidentate Triorganophosphite <u>3</u> (mol % Phosphorus)
H ₂ O	6	71
1 st 1M NaOH	4	74
2 nd 1M NaOH	0	76

Many modifications and other embodiments of the inventions set forth herein will come to mind to one skilled in the art to which these inventions pertain having the benefit of the teachings presented in the foregoing descriptions and the associated Figure. Therefore, it is to be understood that the inventions are not to be limited to the specific embodiments disclosed and that modifications and other embodiments are intended to be included within the scope of the appended claims.

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CLAIMS

What is claimed is:

1. A method for separating one or more triorganophosphite components from a crude phosphite mixture containing acidic hydrolysis products, said method comprising:

contacting said crude phosphite mixture with a basic additive to produce a second mixture comprising a first phase and a second phase, wherein said first phase comprises the basic additive and one or more components independently selected from the group consisting of $(R^2O)(R^3O)POH$, $(R^1O)(HO)PO(H)$ and H_3PO_3 , wherein R^1 , R^2 and R^3 are independently selected from the group consisting of C_1 to C_{18} alkyl, C_6 to C_{18} aryl and hydroxyaryl, and C_3 to C_{18} cycloalkyl and hydroxyalkyl radicals, and wherein R^2 and R^3 can optionally be connected to each other directly by a chemical bond or through an intermediate divalent group R^9 ; and

said second phase comprises one or more triorganophosphite components independently selected from the group consisting of $(R^4O)(R^5O)P(OR^6)$ and $((R^7O)(R^8O)PO)_nA$, wherein R^4 , R^5 , R^6 , R^7 and R^8 are independently selected from the group consisting of C_1 to C_{18} alkyl, C_6 to C_{18} aryl and C_3 to C_{18} cycloalkyl radicals, wherein each of R^4 , R^5 and R^6 can optionally be connected to one or both of the other two directly by a chemical bond or through an intermediate divalent group R^{10} , wherein R^7 and R^8 can optionally be connected to each other directly by a chemical bond or through an intermediate divalent group R^{11} , wherein A is an optionally substituted or unsubstituted aliphatic, aromatic or heteroaromatic radical, wherein n is an integer greater than 1; and

wherein R^9 , R^{10} , and R^{11} are independently selected from the group consisting of $-O-$, $-S-$, and $-CH(R^{12})-$, wherein R^{12} is selected from the group consisting of H , C_6 to C_{18} aryl, and C_1 to C_{18} alkyl.

2. The method of claim 1 wherein the basic additive comprises at least one compound selected from the group consisting of NaOH, KOH, Na₂CO₃, K₂CO₃, Ca(OH)₂, NH₄OH, CaCO₃, a strongly basic anion-exchange resin, and combinations thereof.
3. The method of claim 1 wherein the basic additive comprises a strongly basic anion-exchange resin.
4. The method of claim 3 wherein said strongly basic anion-exchange resin comprises polymer-bound tetraorgano-ammonium hydroxide groups.
5. The method of claim 1 wherein said basic additive and said crude phosphite mixture are contacted at a temperature of from about -10 °C to about 30 °C.
6. The method of claim 1 wherein said first phase is maintained at or above a pH of about 13.5.
7. The method of claim 1 further comprising the step of contacting said crude phosphite mixture with water before contacting the crude phosphite mixture with said basic additive.
8. The method of claim 1 further comprising the steps of:
 - b) separating said first phase from said second phase; and
 - c) contacting said second phase with the basic additive.
9. The method of claim 8 further comprising the step of:
 - d) contacting the second phase with a brine solution.

10. The method of claim 1 further comprising contacting the second phase with a transition metal or a transition metal compound to produce a transition metal-triorganophosphite catalyst or catalyst precursor.

11. A method for preparing triorganophosphites, said method comprising:

a) contacting an alcohol with PCl_3 in the presence of a triorganoamine to produce a first reaction product comprising one or more organophosphites and triorganoamine hydrochloride;

b) removing the triorganoamine hydrochloride from the first reaction product to produce a second reaction product optionally containing one or more triorganophosphites, diorganohydrogenphosphite $(\text{R}^2\text{O})(\text{R}^3\text{O})\text{POH}$, organodihydrogenphosphite $(\text{R}^1\text{O})(\text{HO})\text{PO}(\text{H})$ and H_3PO_3 , wherein R^1 , R^2 and R^3 are independently selected from the group consisting of C_1 to C_{18} alkyl, C_6 to C_{18} aryl and hydroxyaryl, and C_3 to C_{18} cycloalkyl and hydroxyalkyl radicals, and wherein R^2 and R^3 can optionally be connected to each other directly by a chemical bond or through an intermediate divalent group R^9 , wherein R^9 is selected from the group consisting of $-\text{O}-$, $-\text{S}-$, and $-\text{CH}(\text{R}^{12})-$, wherein R^{12} is selected from the group consisting of H , C_6 to C_{18} aryl, and C_1 to C_{18} alkyl; and

c) removing at least a portion of one or more of the compounds diorganohydrogenphosphite $(\text{R}^2\text{O})(\text{R}^3\text{O})\text{POH}$, organodihydrogenphosphite $(\text{R}^1\text{O})(\text{HO})\text{PO}(\text{H})$ and H_3PO_3 by contacting the second reaction product with a basic additive.

12. The method of claim 11, wherein the basic additive comprises at least one compound selected from the group consisting of NaOH , KOH , Na_2CO_3 , K_2CO_3 , $\text{Ca}(\text{OH})_2$, NH_4OH , CaCO_3 , a basic ion-exchange resin, and combinations thereof.

13. The method of claim 11, wherein step c) is repeated.