Phosphonic acid: preparation and applications
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Abstract
The phosphonic acid functional group, which is characterized by a phosphorus atom bonded to three oxygen atoms (two hydroxy groups and one P=O double bond) and one carbon atom, is employed for many applications due to its structural analogy with the phosphate moiety or to its coordination or supramolecular properties. Phosphonic acids were used for their bioactive properties (drug, pro-drug), for bone targeting, for the design of supramolecular or hybrid materials, for the functionalization of surfaces, for analytical purposes, for medical imaging or as phosphoantigen. These applications are covering a large panel of research fields including chemistry, biology and physics thus making the synthesis of phosphonic acids a determinant question for numerous research projects. This review gives, first, an overview of the different fields of application of phosphonic acids that are illustrated with studies mainly selected over the last 20 years. Further, this review reports the different methods that can be used for the synthesis of phosphonic acids from dialkyl or diaryl phosphonate, from dichlorophosphine or dichlorophosphine oxide, from phosphonodiamide, or by oxidation of phosphinic acid. Direct methods that make use of phosphorous acid (H₃PO₃) and that produce a phosphonic acid functional group simultaneously to the formation of the P–C bond, are also surveyed. Among all these methods, the dealkylation of dialkyl phosphonates under either acidic conditions (HCl) or using the McKenna procedure (a two-step reaction that makes use of bromotrimethylsilane followed by methanolysis) constitute the best methods to prepare phosphonic acids.

Review
1. Introduction
Phosphonic acid is a functional group featuring two hydroxy moieties, one P=O double bond and one P–C bond. This functional group was incorporated in a broad diversity of molecules and polymers to introduce specific properties including water solubility, coordination or supramolecular properties. Several books, book chapters or reviews have been focused on the construction of the P–C bond [1], on the description of the different classes of phosphorus-containing functional groups [2], on specific applications (hybrid materials [3], surface modification [4], oil industry [5]) or dedicated to a family of compounds
Figure 1: Summary of the synthetic routes to prepare phosphonic acids detailed in this review. The numbers indicate the corresponding sections of this review.

2. Phosphonic acids: properties and applications

In the solid state, a phosphonic acid function possesses one P–O bond which is shorter than the two others and can be attributed to the P=O double bond (as an example for methylphosphonic acid, Figure 2, the P=O bond length is 1.4993(11) Å, the two other P–O bond lengths are 1.5441(11) Å and 1.5443(12) Å and the P–C bond is 1.7586(17) Å). The bond angles at the phosphorus atom are ranging from 103.46(8)° to 112.86(7)° for methylphosphonic acid, indicating a distorted tetrahedral geometry around the phosphorus [9]. This function is very stable but under some oxidative conditions (e.g., Mn(II) with O₂) the rupture of the P–C bond can occur to produce phosphate [10]. The phosphonic acid function possesses two acidic protons featuring, for instance, when R is an aromatic moiety, a first pKₐ ranging from 1.1 to 2.3 and the second acidity which features a pKₐ ranging from 5.3 to 7.2 [11]. The values strongly depend on the electronic properties induced by the substituent R. It must be noted that due to the high polarity of the phosphonic acid function, the purification of phosphonic acids is quite difficult except by recrystallization for the solid samples. The purification by chromatography on silica gel requires very polar eluents (e.g., CHCl₃/MeOH/H₂O 5:4:1, v/v/v) [12] and further purification by HPLC with a C18 grafted column. Due to these difficulties of purification, it must be noted that in many cases the purification occurs on the precursors of the phosphonic acid. As an example, dialkyl phosphonates (the diester derivatives of phosphonic acid) can be easily purified by chromatography on silica gel and diverse clean and efficient methods can be applied to produce phosphonic acid from phosphonates without the need of intense purification in the final step. Indeed, the purification is limited to remove the solvent and volatile reagents that are frequently used in excess (see section 3). Due to the acidity of phosphonic acid this function is deprotonated in water and this property was used to increase the water solubility of organic compounds [13,14], polymers [15,16] or ligands for coordination chemistry [17]. It must be noted that for similar compounds, that only differ by the replacement of a carboxylic acid group with a phosphonic acid moiety, the log PₐW values (partition coefficient between octanol/water solution) are decreased by about 1 log unit (meaning that the phosphonic acid derivatives when placed in an octanol/water mixture the compound’s concentration in the aqueous phase is 10 times the concentration of the corresponding carboxylic acid analogue) [11]. Meanwhile, the phosphonic acid derivatives are more acidic when compared to their carboxylic acid equivalents (1.9 to 2.9 units of pKₐ below

(e.g., aminophosphonic acids [6], organometallic phosphonic acids [7]). However, no recent review was focused on the different methods that can be employed to prepare phosphonic acids which is a function needed to address numerous applications that are summarized in the first part of this review (section 2). Then, we review the principal methods that can be employed to prepare phosphonic acids. The most frequently applied methods start from phosphonates (section 3 and Figure 1). However, other possibilities exist: the hydrolysis of dichlorophosphine or dichlorophosphine oxide (section 4), the hydrolysis of phosphonodiamide (section 5), the direct methods that make use of phosphorous acid (H₃PO₃) to create the P–C bond simultaneously to the formation of phosphonic acid (section 6) or the oxidation of phosphinic acid (section 7). The last section (section 8) includes additional miscellaneous methods to prepare phosphonic acids. Of note, the biosynthesis of phosphonic acid, which is a dynamic field of research aiming to discover new bioactive compounds [8], will not be detailed in this review which is focused on the chemical way of producing phosphonic acids. Of note, this review is not an exhaustive list of all the phosphonic acids synthesized over the last 20 years but a selection of examples that aim to illustrate the most relevant methods that can be employed to produce phosphonic acids.
for the first acidity of phosphonic acid derivatives) [11]. This property was used to design Brønsted acid catalysts that were used for the depolymerization of cellulose [18] and the synthesis of dihydroypyrimidine derivatives [19]. The monosodium salt of phosphonic acids were also employed as organocatalysts for Michael addition [20]. The capacity of phosphonic acids to increase the solubility of organic compounds in water was employed to develop water soluble catalysts [21], or to improve the water solubility of drug-chelate supramolecular assemblies (e.g., calixarene) [22,23]. The water solubility of phosphonic acid is strongly improved when the phosphonic acid is deprotonated (basic media). It is not rare to observe, when preparing NMR tubes with D$_2$O as solvent that phosphonic acid appeared weakly soluble whereas the addition of K$_2$CO$_3$ induced its solubilization. Interestingly the increase of the number of phosphonic acid functional groups on a molecule is not systematically associated with an increase of its water solubility as exemplified with triphenylphosphine functionalized either by two or three phosphonic acid sodium salts; the former being more water soluble than the latter [24]. This behavior is associated to the fact that the trisphosphonic acid sodium salt was already solvated at the solid state.

![Chemical structure of dialkyl phosphonate, phosphonic acid and illustration of the simplest phosphonic acid: methylphosphonic acid.](https://example.com/structure.png)

**Figure 2:** Chemical structure of dialkyl phosphonate, phosphonic acid and illustration of the simplest phosphonic acid: methylphosphonic acid.

The high polarity of the phosphonic acid function and its ability to be ionized in water render this group attractive to design the polar head group of anionic amphiphilic compounds [25,26] or polymeric-based amphiphilic derivatives [27]. This type of compound was used as surfactant to stabilize colloidal solutions of nanocrystals [28], to prepare oil/water emulsions [29], or to prevent corrosion [30-32]. The ionic interactions between phosphonic acid and lipophilic amine produced catanionic supramolecular aggregates that were assessed as HIV inhibitors [26,33].

Many phosphonic acids are present in nature and their biosynthesis involves, as one of the key steps, the isomerization of phosphanenolpyruvate to phosphonopyruvate which is catalyzed by phosphopyruvate mutase [34]. Recent studies indicate that up to 25% of the phosphorus available in the ocean would consist of phosphorus species featuring one P–C bond [35] pointing out the importance of such type of compounds in natural biochemical processes [36]. Phosphonic acid mimics the phosphate group, which is omnipresent in nature, but also the tetrahedral transition-state intermediate encountered for instance during the hydration of carbon dioxide by carbonic anhydrase [37] or during the hydrolysis of amide [38]. The main difference between phosphate and phosphonic acid arises from the higher stability (resistance towards enzymatic degradation) of the P–C bond when compared to the P–O bond present in phosphate. Accordingly, phosphonic acid was used for numerous applications in biology and medicine to mimic the phosphate group leading to antiretroviral drugs (e.g., tenofovir (1)) [39], isoprenoid biosynthesis inhibitors [40,41], antibiotics (e.g., fosfomycin (2)) [42], tyrosine phosphatase inhibitors 3 [43], antimalarial 4 [44], antihypertensive drugs (e.g., K4 5 and K26 6) [45] or the anti-osteoporosis compounds alendronate (7) [46] and zoledronate (8) [47] (Figure 3A). In some cases, the bioactive phosphonic acid is generated in vivo from a phosphonate pre-drug [48] as exemplified by the formation of 10 from 9 which permits to improve the pharmacokinetic properties (Figure 3B) [49]. Some phosphonic acid-containing compounds were also used for their herbicidal properties as exemplified by glyphosate (11) [50] (Figure 3C). Finally, in the field of immunotherapy, γδ T cells, at the difference to αβ T cells, respond to non-peptidic antigen or to the alterations in the expression of normal cell surface components. The subset of the human γδ T cells that express the Vγ9Vδ2 receptor recognize and present non-peptide antigens including prenyl pyrophosphate analogues also identified as phosphoantigens. These Vγ9Vδ2 T cells are involved during bacterial or protozoan infections but play also a role in tumor immunity [51]. The synthesis of prenyl pyrophosphate analogues was assessed as a strategy to activate Vγ9Vδ2 T cells [52]. These analogues are mainly composed of pyrophosphate compounds but some phosphonic acids such as compound 12 was identified as an activator of Vγ9Vδ2 T cells [53,54] (Figure 3D).

Phosphonic acid is a function that is readily involved in hydrogen bonds thus generating auto-assembling supramolecular structures [55] or, when a second organic partner possessing basic groups is present (e.g., amine), mixt supramolecular materials [56-58]. This supramolecular behavior was also involved in the development of organic materials exhibiting proton conduction [59], or organo-gel properties [60]. In the field of analytical chemistry, molecules functionalized with phosphonic acid groups were used to prepare the solid phase for immobilized metal affinity chromatography (IMAC). Such types of solid phases were applied to enrich carbohydrate from extracts [61] or as chiral selectors immobilized on silica for chiral cation exchange chromatography [62] (Figure 4). Porphyrin tetra-functionalized with phosphonic acid groups was used as a chemo-sensor for trinitrotoluene (TNT) [63].
The third important feature of phosphonic acid arises from its coordination properties. Indeed, the three oxygen atoms can be engaged in coordination or ionic-covalent bonds. These coordination properties were used to design molecular hybrid materials [64] also identified as metal organic framework (MOF) or coordination polymers that are synthesized by reaction with a metallic salt (e.g., copper [65], lanthanides [66]) under hydrothermal conditions [67,68] or by insertion of phosphonic acid in low dimensional inorganic materials like layered double hydroxide (LDH) [69] or layered simple hydroxide (LSH) [70]. These materials prepared from phosphonic acids were assessed for numerous applications, including nuclear fuel stewardship and separations of actinides [71], porosity [72-74], bactericidal properties [75], inorganic salt release [76], luminescence [77,78], protonic conduction [79,80], materials with magnetic properties [81,82], or were spread off in polymers to produce nanocomposites [83,84]. Of note, despite the non-chiral nature of compound 13 (Figure 4), it produced, when associated with copper salt, a homochiral non-centrosymmetric crystalline hybrid material [85]. The coordination properties of phosphonic acid were also applied to design tetraazamacrocyclic compounds functionalized with phosphonic acid pendant arms. As an example, ((1,4,7,10-tetraazacyclododecane-1,4,7,10-tetrayl)tetrakis(methylene))tetraphosphonic acid, DOTP, compound 14, Figure 4), was assessed to complex indium [86] or terbium [87] with the aim to develop bone targeting and dosimetry. It was also employed to complex 89Y and applied as pH sensitive NMR probe [88], or as organic precursor of crys-
Figure 4: Illustration of the use of phosphonic acids for their coordination properties and their ability to be involved for the synthesis of hybrid materials or to interact with metal-oxide surfaces.

The next sections report the most important and efficient methods to produce phosphonic acids.

3. Preparation of phosphonic acids from phosphonates

3.1. Hydrolysis of phosphonates with hydrochloric acid

The most general method to prepare phosphonic acids from phosphonates is to use concentrated HCl solution (35–37% in water; \( \approx 12 \) M) at reflux for 1 to 12 h as shown in Figure 5. At the end of the reaction the excess of HCl and water can be readily eliminated by distillation and finally, the last traces of water can be removed by azeotrope distillation with toluene. Phosphonic acids, that can be very hygroscopic, can be further dried by leaving the product in a desiccator over P₂O₅ but, usually, they do not require further purification. When further purification is required, purification can be achieved by recrystallization in polar solvents (e.g., acetonitrile \[106\], methanol/acetone \[117\], water \[55\]), or in concentrated HCl \[118\] (acidic media reduces the water solubility of phosphonic acids). When
recrystallized in aqueous solution, the phosphonic acid can co-crystallize with water due to the presence of hydrogen bonds between the phosphonic acid moiety and H$_2$O [119].

As depicted on Figure 6, the selected phosphonic acids prepared by the hydrolysis of phosphonate with concentrated HCl water solution present only few other functional groups. This method was applied to prepare the alkylphosphonic acid $\text{16}$ [120], arylphosphonic acid $\text{17}$ [121] or 18 or block co-polymers possessing pendant arms functionalized with phosphonic acids (compound $\text{19}$ [122]). The hydrolysis with 20% HCl solution ($\approx$6 M) was also occasionally applied to coordination complexes bearing a phosphonate function as exemplified with compound $\text{20}$ [123].

HBr was used more occasionally to hydrolyze phosphonates [124-126]. Regarding the nature of the alkyl chains of dialkyl phosphonates, it must be noted that most of the time these chains are methyl, ethyl, isopropyl or, occasionally, $n$-butyl [127]. This observation is likely explained by the synthetic methods employed to prepare phosphonates that frequently made use of the Arbuzov reaction [128] involving the commercially available trimethyl, triethyl or triisopropyl phosphite [129]. To check the achievement of phosphonate hydrolysis, phosphorus NMR is a method of choice and can also eventually detect side reactions. Indeed, with some substrates it was found that the P–C bond can be cleaved when treated with concentrated HCl as first reported by Redmore et al. [130]. For instance, hydroxynaphthylphosphonate $\text{21}$ when treated with HBr induced partial rupture of the P–C bond to yield phosphoric acid which is difficult to remove from the expected phosphonic acid $\text{22}$ (Figure 7A) [131]. The same side reaction was observed in the course of the acidic hydrolysis of 4-hydroxybenzenephosphonate $\text{23}$ into phosphonic acid $\text{24}$ (Figure 7B). However, the regioisomer diethyl 3-hydroxyphenylphosphonate can be converted to phosphonic acid $\text{25}$ with HCl 35% at reflux without any P–C bond cleavage (unpublished result). The diphosphonic acids $\text{26}$ and $\text{27}$ were also prepared by hydrolysis with HCl 35% without P–C bond cleavage. These results indicated that the mesomeric effect induced by the phenol function likely explained the difference of stability of compounds $\text{23}$ and diethyl 3-hydroxyphenylphosphonate in refluxing HCl 35% solution. Of note, if classical heating at 100 °C is usually employed some reaction under microwave activation was reported to reduce the reaction time up to only 7 minutes [132].

![Figure 5: Hydrolysis of dialkyl phosphonate to phosphonic acid under acidic conditions.](image)

![Figure 6: Examples of phosphonic acids prepared by hydrolysis of dialkylphosphonate with HCl 35% at reflux (16–19) or 20% HCl solution (20).](image)
For the mechanism of hydrolysis of phosphonate by hydrochloric acid, some works dedicated to assess the hydrolysis of trimethyl phosphate are instructive and can likely be extrapolated to phosphonate. When trimethyl phosphate (TMP) is placed in 4 N HCl solution, infrared analysis showed a shift of the P=O band from 1273 cm\(^{-1}\) to 1236 cm\(^{-1}\) [133]. When the HCl concentration was further increased to 12 N, this band disappeared. The titration of trimethyl phosphate with HCl was also monitored by \(^{31}\)P NMR and the results are consistent with a protonation at the oxygen atom that is doubly bonded to the phosphorus atom leading to a phosphorus centered cation (the protonated trimethyl phosphate was characterized by a pK\(_a\) of \(-3.6\)). These results, extrapolated to phosphonate, suggest that the protonation of the phosphonate would occur on the oxygen atom which is doubly bonded to the phosphorus atom. This protonation likely yields the intermediate I (Figure 8) which exists as two mesomeric forms. Then, two competitive mechanisms can occur. Intermediate I can lose a carbocation according to a S\(_{N}1\) mechanism whereas the second way could consist in a nucleophilic substitution (S\(_{N}2\)) involving chloride ions as nucleophilic species to produce the intermediate II. Then, a repetition of this mechanism yields phosphonic acid.

Figure 7: A) and B) Observation of P–C bond breaking during the hydrolysis of phosphonate with concentrated HX water solutions; C) examples of compounds prepared by hydrolysis of dialkyl phosphonate in presence of a 35% HCl–water solution without the observation of a P–C bond cleavage.

Figure 8: Mechanism of the hydrolysis of dialkyl phosphonate with HCl in water.
The preponderant route is likely governed by the stability of the carbocation and the steric hindrance around the electrophilic carbon atom of intermediate I (Figure 8).

Di-tert-butyl phosphonate belongs to the precursors of phosphonic acid that requires milder acidic conditions to be converted into phosphonic acid. The mechanism likely occurs following the $S_N$1 pathway as reported above. This type of di-tert-butyl phosphonate was used for the synthesis of phosphonic acid analogues of peptides like protein-tyrosine kinases (PTKs) [134] or to prepare gadolinium complexes used as magnetic resonance imaging (MRI) contrast agents [135,136]. The introduction of the di-tert-butyl phosphonate was achieved by the reaction between di-tert-butyl phosphate and either benzyl halide [137] (Michaelis–Becker reaction) or aldehyde [134] (Pudovik reaction) or using tris-tert-butyl phosphate (Abramov reaction) [135]. The elimination of the $t$-Bu moieties can be efficiently achieved in presence of trifluoroacetic acid (TFA). As exemplified in Figure 9, the use of TFA induces the formation of the phosphonic acid 29 concomitantly with the formation of carboxylic acid from the tert-butyl ester. The final compounds were purified by removing the excess of TFA (72.4 °C at 1 bar) [138], using HPLC [137] or by precipitation in diethyl ether [136].

All the precedent examples illustrate the hydrolysis of dialkyl phosphonate in phosphonic acid under acidic conditions. Similar protocols (concentrated HCl or HBr) can be used to prepare phosphonic acid from diphenyl phosphonate (Figure 10A). The use of a mixture of HBr in acetic acid was also reported [139,140]. It must be noted that the reaction work-up is usually slightly different since the treatment with propylene oxide that acts as an acid scavenger, can be applied during the purification step. The preparation of phosphonic acid from diphenyl phosphonate was reported to prepare phosphohomocysteine 30 [141], the arginine mimetic 31 that was developed as a potent metallo-aminopeptidase inhibitor [142] or the aminophosphonic acid 32 [143] (Figure 10B).

The mechanism of hydrolysis of diphenyl phosphonate in acidic conditions is likely different to the one occurring with dialkyl phosphonates (Figure 8). For the hydrolysis of diphenyl phosphonate it is likely that after protonation of the phosphonate, water acts as a nucleophile and subsequently phenol is eliminated (Figure 11). The repetition of this sequence produces phosphonic acid.

Of note, the treatment of phosphonate with nucleophilic basic reagents like NaOH [144], LiOH [145] or NaHCO$_3$ [146] is not

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**Figure 9:** Hydrolysis of bis-tert-butyl phosphonate 28 into phosphonic acid 29 [137].

**Figure 10:** A) Hydrolysis of diphenyl phosphonate into phosphonic acid in acidic media. B) Examples of phosphonic acids prepared by this method.
suitable to produce phosphonic acid. Indeed, these conditions yield phosphonic acid mono-esters. This monohydrolysis takes place with dialkyl phosphonate or diphenyl phosphonate [146]. For dimethyl phosphonate, the monohydrolysis can be also achieved using NaI as nucleophile in acetone [147] or butanone as solvent [148].

3.2. Catalytic hydrogenolysis

Dibenzyl phosphonates are readily synthesized using dibenzyl or tribenzyl phosphite. This type of phosphonate offers an alternative to the use of acidic conditions to prepare phosphonic acid since the benzyl moieties can be removed by hydrogenolysis (Figure 12). Palladium on charcoal, which is the most used method to prepare phosphonic acid from dibenzyl phosphonate [149], was the catalyst used to prepare the phosphonic acids 33 [150], 34 [151] and 35 [152] (Figure 12). Of note, mono-hydrolyzed phosphonate can be obtained when a monoalkyl-monobenzyl phosphonate is hydrogenolyzed [150,153]. The synthesis of phosphonic acid from diphenyl phosphonates can be prepared under acidic conditions (see above) but can be also achieved in presence of Adam’s catalyst (PtO₂) and hydrogen as exemplified by the preparation of the compounds 36 [154] and 37 [155] (Figure 13).

It is also worth noticing that phosphonic acid can be prepared by the hydrogenolysis of diallyl phosphonates. For this purpose the use of the Wilkinson catalyst (ClRh(PPh₃)₃) was reported [156].

3.3. McKenna’s method – use of bromotrimethylsilane

As reported above, the hydrolysis in acidic media requires harsh conditions (most of the time concentrated HCl solution in water at reflux) that finally limit its application when the molecules possess sensitive functional groups. The need of a milder method to prepare more functionalized phosphonic acid deriva-
tives was therefore required. In some aspect the selection of benzyl or tert-butyl phosphonate can be suitable since specific mild conditions can be applied to prepare phosphonic acids as discussed above. Nevertheless, as the simplest way to prepare phosphonate involves trialkyl phosphate, the subsequent dealkylation of dialkyl phosphonate under mild conditions was of a great importance. The group of C. E. McKenna reported in 1977 the use of bromotrimethylsilane as a reagent that permitted an efficient transesterification of dialkyl phosphonate to bis-(trimethylsilyl) phosphonate. It is worth noticing that these silylated phosphonates produced quantitatively phosphonic acid derivatives after water or alcohol (methanol, ethanol) treatment [157]. In this initial work, NMR data indicated that silylation of dimethyl or diethyl phosphonate with bromotrimethylsilane was almost quantitative. Other studies indicated that this reaction can be also applied to disopropyl phosphonate, or di-tert-butyl phosphonate leading to the conclusion that all dialkyl phosphonates can be dealkylated by the transesterification followed by a methanolysis or hydrolysis subsequent step. As suggested in the initial works of McKenna, the mechanism occurs by an oxophilic substitution on the silicon atom whereas bromide acts as a leaving group to produce the intermediate I (Figure 14). This intermediate is then dealkylated following a similar path than that occurring with the Arbuzov reaction to produce the intermediate II [158]. The repetition of this mechanism produced de disilylated intermediate III. The hydrolysis of this intermediate III produced phosphonic acid, trimethylsilanol and hexamethyldisiloxane that are two volatile side products. The methanolysis of the intermediate III is even a better choice to transform III into phosphonamic acid since the methoxymethylsilylamine is also volatile and methanol used in excess is also more volatile than water. It must be noted that an experimental proof (using $^{17}$O- and $^{18}$O-enriched diethyl phenylphosphonate) has reported that the terminal oxygen doubly bonded to the phosphorus atom was indeed the nucleophilic atom that attacks the silicon atom [159].

Regarding the mechanism of this reaction, the oxophilic silylation involving the oxygen atom which is doubly bonded to the phosphorus atom, thus producing the intermediate I, is also supported by the works of Bartlett et al. [160] (Figure 15A) that show that the treatment of the phosphonate-thiophosphonate 38 with iodotrimethylsilane produced, after methanolysis, only the phosphonic acid 39. Of note, the thiophosphonate functional group is not hydrolysed. The need of the nucleophilic halide anion is another feature of this reaction. Indeed the treatment of TMSOTf on a phosphonate did not induce the dealkylation likely due to the absence of nucleophilic species [161].

It is worth noticing that the nucleophilic attack of bromide only occurs on alkyl chains as exemplified by a work of Pohjala et
al. [162] (Figure 15B) The authors showed that the diphosphate 40 featuring both phenyl and alkyl substituents treated with chlorotrimethylsilane and sodium iodide in CH$_3$CN yielded, after methanolysis and treatment with NaOH, compound 41 resulting from the hydrolysis of the diethyl phosphonate moiety. Finally, it must be noted that BrSiMe$_3$ selectively induced the silylation of phosphonate without affecting ketone, amide, halogenoalkane or alkyne [163]. The preparation of phosphonic acid from phosphonate with bromotrimethylsilane was also applied to methylene-tris-phosphonic acids [164], and to substituted hexaethyl 1,1,1-tris phosphonate as illustrated with the reactivity of compound 42. This reaction occurred in presence of collidine to prepare, after treatment with sodium hydroxide, the collidinium salt 43 [165] (Figure 15C).

The mechanism of dealkylation with bromotrimethylsilane, that produces a bis-silylated phosphonate intermediate, thus incited to use tris(trimethylsilyl) phosphite as nucleophilic phosphorus species to produce after methanolysis phosphonic acid [166] or hydroxymethylene bis-phosphonic acid as illustrated in Figure 16 with the synthesis of the hydroxymethylenebisphosphonic acid 45 [167].

The trans-esterification of phosphonate with bromotrimethylsilane to produce silylated phosphonate requires at least two equivalents of BrSiMe$_3$ per phosphonate function. Usually an excess is engaged in the reaction to guarantee a full conversion of dialkyl phosphonate into the disilylated phosphonate. The same reaction is also observed with the use of iodosotrimethylsilane (Me$_3$SiI) [168]. However, with chlorotrimethylsilane the reaction is not really efficient as indicated in the initial work of McKenna since only a partial conversion was observed after several days of reaction. Nevertheless, the use of a mixture of chlorotrimethylsilane with NaI in acetonitrile can be used to achieve the silylation of dialkyl phosphonates that can be then transformed in phosphonic acids after hydrolysis or methanolysis. Following these experimental conditions, the efficacy is likely explained by the in situ generation of iodosotrimethylsilane that result from a halogen exchange reaction [169]. This method (ClSiMe$_3$ + NaI in acetonitrile) is currently less employed likely
due to the need to remove NaI from the final phosphonic acid after the step of hydrolysis or methanolysis. It is worth noticing that when NaI, LiBr or KI was used alone in anhydrous solvents (acetone, MeCN or butanone) under heating (80–100 °C) a selective monodeprotection of the dialkyl phosphonates (alkyl = methyl or ethyl) was observed. The sodium or lithium salts being formed in high yields (87–97% yields) [170]. This procedure was applied to prepare a molecular receptor of lysine residue [171]. Following this procedure, the monodealkylation can be explained by the formation of a sodium salt that due to electronic factors prevent the second dealkylation. However, the protonation of this anionic compound 46 by ion exchange procedure yields 47 that can then be fully dealkylated with NaI in acetonitrile to produce the phosphonic acid disodium salt 48 [172] (Figure 17).

Bromotrimethylsilane is nowadays the gold standard reagent to produce phosphonic acid from dialkyl phosphate under mild conditions (usually at room temperature). This reaction can be achieved in a non-protic solvent including CH₂Cl₂ [173], acetonitrile [174], chloroform [175], DMF [176], pyridine [177] or collidine [178]. The use of BrSiMe₃ simultaneously as reagent and solvent was also reported [120]. The direct production of sodium salts of phosphonic acid is readily achieved by adding 2 N NaOH/water solution to the silylated phosphonate [178]. This two-step sequence: 1. bromotrimethylsilane; 2. methanolysis or hydrolysis was used for the synthesis of numerous phosphonic acids including heterocyclic compounds that are too sensitive to be prepared by the transformation of phosphonates under acidic conditions. As an illustration, the thiophene diphosphonic acid 49 [173], the pyridine oxide 50 [179], the furan phosphonic acid 51 [174], the bipyridine bis thiophene phosphonic acid 52 [180] or the α-aminophosphonic acid 53 that was assessed as inhibitor of the human farnesyl pyrophosphate synthase (hFPPS) [181] were prepared from their corresponding diethylphosphonates (Figure 18). Nucleotide analogues were also prepared by this methodology as exemplified by the compound 54 [176] or 55 [182]. For these last two examples, the bromotrimethylsilane induced the silylation of diethyl phosphate but also the phosphoramidate and the phosphinate functional groups. Other phosphonic acids possessing different functionalities including phosphine 56 [183], trimethylsilyl 57 [184], diazo 58 [185] or styrene 59 [186] moieties were also prepared efficiently using bromotrimethylsilane followed by methanolysis.

This two-step methodology (1. BrSiMe₃; 2. MeOH or H₂O) was also applied to prepare macromolecules functionalized with phosphonic acid functional groups. For instance, a cyclodextrine derivative (compound 60 [177], Figure 19), amphiphilic compounds that were used for self-assembled monolayers (e.g., compound 61 [187]), polymers (PEG derivative 62 [188] or functionalized polyethylene 63 [189]) were reported. The McKenna’s procedure was also applied to prepare polyphosphonic acids as exemplified by the compounds 64 [120], 65 [190], 66 [191] and the dendrimers 67 [192].

The use of BrSiMe₃ was also applied to diverse organometallic compounds as exemplified in Figure 20. This procedure was applicable to compounds featuring CpFe(CO)₂ 68 [193] or ferrocenyl 69 [194,195] moieties but also to palladium or platinum pincer complexes 70 [196] or arene–chrome carbonyl platinium pincer complexes 71 [197]. It must be noted that when the organometallic complex possessed halogen–metal moiety, then Me₃SiBr can induce halogen exchange [198].

The use of bromotrimethylsilane to prepare phosphonic acid from dialkyl phosphate is rarely associated with undesired side reactions. Nevertheless, G. David et al. have reported in a study dedicated to the synthesis of methacrylate monomers functionalized with phosphonic acids the occurrence of side compounds [199]. The treatment of the bis-phosphonic 72 with TMS-Br followed by methanolysis, induces a cleavage of the carboxylic esters simultaneously with the formation of the phosphonic acid 75 (Figure 21). The authors have shown, using model compounds, that the hydrolysis is triggered by the acidity of the phosphonic acid. Interestingly, the addition of aqueous ammonia during methanolysis of the disilylated phosphate produced the expected phosphonic acid 73 without any cleavage of the carboxylic ester function.

A second example reported by Pailloux et al. shows that depending on the reaction time for the hydrolysis of the silyl-
Figure 18: Phosphonic acid synthesized by the sequence 1) bromotrimethylsilane 2) methanolysis or hydrolysis. The notation R indicates the nature of the alkyl chains present on the phosphonate precursors of these phosphonic acids.

lated phosphonate 76 (Figure 22) either the expected phosphonic acid 77 or the phosphonic acid 78 resulting from the opening of the benzoxazole fragment (after 3 days in contact with water), were produced [200]. This side reaction is likely explained by the sensitivity of the product to acidic condition coming from the acidity of the phosphonic acid function. During the hydrolysis the precipitation of the phosphonic acid 77 occurred; a rapid filtration (5 minutes) led to isolate the phosphonic acid 77 in 70% yield. However, a prolonged contact time with water (3 days) yields the phosphonic acid 78 that features a benzoxazole ring opening. It can be concluded form these studies (Figure 21 and Figure 22) that the side reactions is not due to bromotrimethylsilane but to the sensitivity of the product to acidic conditions that are induced by the acidity of the phosphonic acid functional group.

3.4. Dealkylation with boron reagents

Dealkylation of dialkyl phosphonates could also be performed using C_6H_11BCl_2 [201] or BBr_3 [202] under mild and efficient conditions as reported by Mortier et al. The reaction can be applied to dialkyl phosphonates (R = Me, Et, iPr, t-Bu) and proceed first at −30 °C and then at 70 °C in an aprotic solvent (toluene) for 6 hours (Figure 23). The authors reported the formation of a primary adduct R_3B=O=PR(OR')_2 which was subsequently decomposed as borophosphonates oligomers.
through a bis-bromodealkylation [203]. The phosphonic acids were finally obtained after methanolysis with quantitative conversion and yields up to 95% Interestingly, this procedure is selective to P–O dealkylation and compatible with the presence of other groups such as allyl, ketone, primary alcohol, phthalimide, ester or thioether as illustrated with the transformation of the phosphonate 79 to the phosphonic acid 80 (Figure 23). The best results were obtained when 0.9 equivalents of BBr$_3$ per phosphonate function was used.

3.5. Other conditions for dealkylation of phosphonate

The dealkylation of diethyl phosphonate groups present in amino acids or peptides requires mild conditions to avoid the reactivity with the other functional groups. If Me$_3$SiBr can be used to dealkylate peptides functionalized with diethyl phosphonate groups [204], other mild conditions, developed in the context of peptide chemistry, were reported. It is based on the use of trimethylsilyl trifluoromethanesulfonate (TMS-OTf) as
Figure 20: Examples of organometallic complexes functionalized with phosphonic acids that were prepared by the hydrolysis of dialkyl phosphonate according to the McKenna’s method. The notation R indicates the nature of the alkyl chains present on the phosphonate precursors of these phosphonic acids.

Figure 21: Side reaction observed during the hydrolysis of methacrylate monomer functionalized with phosphonic acid.

Figure 22: Influence of the reaction time during the hydrolysis of compound 76.

Figure 23: Dealkylation of dialkyl phosphonates with boron tribromide.

silylating agent, trifluoroacetic acid (TFA) as acidic reagent, and dimethylsulfide (DMS) and m-cresol to avoid side reactions (from S_N1 or S_N2 mechanism) [204] that can involve the functional groups present in the amino acid or peptide. These conditions were applied to prepare the compound 82 from 81 [205,206] (Figure 24).
4. From dichlorophosphine (R–PCl₂) or dichlorophosphine oxide (R–P=OCl₂)

Aryl- and to a lesser extent alkylphosphonic acids have been prepared by the hydrolysis under mild conditions of arylidichlorophosphate or arylidichlorophosphate oxide. The hydrolysis is often performed under basic conditions using aqueous sodium hydroxide (5% NaOH); the phosphonic acid being lately isolated by acidification with hydrochloric acid. For example, the sterically hindered phenylphosphonic acid \( 85 \) was prepared in a two-step procedure starting from the dichlorophenylphosphine \( 83 \). Ph–PCl\(_2\) was first oxidized into the phenylphosphine oxide \( 84 \) with sulfuryl chloride in CCl\(_4\) and then hydrolysis occurred with aqueous NaOH to afford the corresponding arylphosphonic acid \( 85 \) in good yield (65% yield, 2 steps) as shown in Figure 25 [207].

More recently, the synthesis of \( \alpha \)-trifluoromethylbenzenephosphonic acid \( 87 \) was prepared by hydrolysis of the phenylidichlorophosphate oxide \( 86 \) with NaOH in acetonitrile (75% yield, Figure 26). The hydrolysis in acidic conditions (10% HCl) was also reported and produced the phosphonic acid with similar yield (75–85%). However, hydrolysis under acidic conditions required an extended reaction time [208].

5. Phosphonic acid from phosphonodiamide \( \text{RP=O} (\text{NR}_2)_2 \)

Phosphonodiamides are commonly used as precursors of phosphonic acids especially for the synthesis of aminophosphonic acids (\( \alpha \) [209,210], \( \beta \) [211] or \( \gamma \) [212]) and also for the synthesis of nucleoside analogues [213]. The use of these intermediates is likely explained by the possibility to use chiral diamine that can act as a chiral auxiliary to control the chirality of the carbon atom in \( \alpha \)-position of the phosphorus atom as exemplified by the alkylation of the phosphonodiamide \( 88 \) (Figure 27). Furthermore, phosphonodiamide can be purified by different methods including chromatography and their transformation to phosphonic acid is easily achieved by hydrolysis under acidic conditions as illustrated by the reaction of compound \( 89 \) that produced the \( \alpha \)-amino-phosphonic acid \( 90 \) after acidic hydrolysis [210] (Figure 27). Similar acidic hydrolysis was used to produce the imidazolyl phosphonic acid \( 91 \) [214], \( 92 \) [215], the chiral phosphonic-carboxylic acid \( 93 \) [216] or the cyclopropylphosphonic acid \( 94 \) that was assessed as inhibitor of the glutamate metabotropic receptors [217].

The synthesis of phosphonic acid via phosphonodiamide intermediates started by the use of the nucleophilic bis(dialkylamino)chlorophosphine. This possibility is illustrated by the recent works reporting the reaction of bis(diethylamino)chlorophosphine \( 95 \) with the acetal \( 96 \) in the presence of a Lewis acid to yield the phosphonodiamide \( 97 \). Then the nucleophilic addition of adenine and the hydrolysis of the phosphonodiamide function in phosphonic acid produce compound \( 98 \) [213] (Figure 28A). A second example corresponds to the nucleophilic addition of \( N \)-heterocyclic phosphine \( 99 \) to a nitroalkene (phospho-Michael reaction) as shown in Figure 28B. The thiourea unit in \( 99 \) plays a crucial role by assuming intramolecular interaction with the nitro function that finally led to the forma-
Figure 27: A) Illustration of the synthesis of chiral phosphonic acids from phosphonodiamides. B) Examples of phosphonic acid prepared by acidic hydrolysis of phosphonodiamides.

Figure 28: A) Illustration of the synthesis of the phosphonic acid 98 from phosphonodiamide 97. B) Use of cyclic phosphonodiamide 100 to prepare phosphonic acid. C) Illustration of the use of lithiated bis(diethylamino)phosphine borane complex 102 to prepare phosphonic acid.

The reduction of the nitro function and the hydrolysis under acidic conditions of the phosphonodiamide produced the phosphonic acid 101 [211]. As a last example, lithiated dianimophosphine borane 102 was added to imine to produce the bis(diethylamino)phosphine borane 103 (Figure 28C). Then acidic hydrolysis conditions and likely air oxidation produced the phosphonic acid 104 [209].

Phosphonodiamide, which is more robust than phosphonate towards some nucleophilic species like a phosphide anion (R₂P-Li), was used to prepare triarylphosphine functionalized with three phosphonic acid groups (Figure 29) [24]. The sequence requires the preparation of the 4-fluorophenylphosphonodiamide 107 which is prepared by nucleophilic addition of an aryllithium intermediate (from 105) on chlorophosphadiamide 106. In the last step, the tris(phenylphosphonodiamide)phosphine 108 was hydrolysed into the tris-phosphonic acid 109 using water and HCl up to pH 1. Of note, the same reaction that engaged diethyl 4-fluorophenylphosphonate instead of the 4-fluorophenylphosphonodiamide 107 is inefficient (3% yield) due to the monohydrolysis of the phosphonate.

6. Direct method from phosphorous acid H₃PO₃
The direct methods correspond to the methods where the formation of the P-C bond yields simultaneously phosphonic acid.
These reactions mainly use phosphorous acid (H$_3$PO$_3$) as source of phosphorus. The main difficulty with such type of reaction is the purification of the product since crystallization, precipitation and dialysis constitute the possible methods of purification. The purification by chromatography requires, due to the high polarity of phosphonic acids, reversed-phase chromatography and is therefore limited to preparative RP-HPLC [218]. Despite these difficulties, some reactions are very efficient as reported below.

6.1. Moedritzer–Irani reaction

The reaction of an amine (primary or secondary amine), formaldehyde (aqueous or solid paraformaldehyde) and phosphorous acid in acidic media produces amino-bis(methylenephosphonic acid, Figure 30A,B). This reaction, is also known as the Moedritzer–Irani reaction since it was first reported by these authors in 1966 [219]. When primary amine is used, the reaction produces a bis-phosphonic acid since the secondary amine that is generated in situ is more reactive than the starting primary amine [220]. In consequence, the stoichiometry must be adapted and 2 equivalents of both formaldehyde and phosphorous acid must be used. The reaction is also applicable to secondary amine and produce mono-aminomethylene phosphonic acid (Figure 30B). The yield of this reaction is variable as illustrated in Figure 30C. The compound 110 was obtained in 20% from diaminocyclohexane [221]. The compound 111 was prepared from a bis(α-aminomethylene diphosphonic acid) and was isolated in 42% yield. This compound was assessed as a scale inhibitor [222]. As a last example, the compound 112 was prepared from diaminooctane using microwave heating. The final product was isolated in 63% yield [223].

The Moedritzer–Irani reaction was also applied to introduce phosphonic acid functions on polymer or dendrimers possessing reactive amine. Accordingly, phosphonic acids were introduced on polyethylene glycol 113 [188], polyethylene imine 114 [224] or chitosan 115 [225]. The functionalization of polyacrylamide obtained by reversible addition-fragmentation chain transfer (RAFT) polymerization was also recently reported to produce 116 (Figure 31). However, the conditions of the Moedritzer–Irani reaction induced the hydrolysis of the polymers side chains [226].
6.2. Reactivity of phosphorous acid (H$_3$PO$_3$) with imine, nitrile and carbonyl groups

The phosphorous acid function reacts with imine in the absence of solvent to produce α-amino-phosphonic acid as exemplified by the synthesis of the α-amino-phosphonic acid 118 from the imine 117 (Figure 32) [227]. It is worth noting that the reaction of H$_3$PO$_3$ with some enamine (e.g., 1-morpholinocyclohexene) gives a reduction of the enamine to amine. This is likely why this reaction was seldomly used.

The reaction of phosphorous acid on nitrile in presence of methanesulfonic acid followed by the addition of POCl$_3$ or PCl$_3$ [222] is a method that produces in one step the aminomethylene bis-phosphonic acid [228] (Figure 33A). This methodology was applied to prepare the compound 119 [229] that was assessed as HIV-1 integrase inhibitor. Compound 120 [230] was prepared simply by the refluxing acetonitrile with phosphorous acid at 130 °C for 12 hours. 120 was a member of a series of compounds developed as a potential inhibitor of the farnesyl pyrophosphate synthase. The reaction also occurs with amide (Figure 33B) as exemplified for the synthesis of compounds 121 [231], 122 [232] or 123 [231].

The nitrile group can be replaced in this reaction with a carboxylic acid function. In that case the final product is a hydroxymethylenebis-phosphonic acid (Figure 34). This reaction was optimized to produce, for instance, compounds 124 [233,234] or 125 [229]. This reaction is not further detailed herein as it was recently reviewed [235].

7. Preparation of phosphonic acid by oxidation of phosphinic acid

Phosphinic acid derivatives (also identified as phosphonous acid) are prepared by reaction of hypophosphorous acid (Figure 35) on alkene or alkyne (hydrophosphonation) [236], by its addition on aldehyde or imine [237] or by hydrolysis of alkyl or aryl dichlorophosphine (RPCl$_2$) [238] (for a review see reference [239]). Then, these phosphinic acids constitute a suitable precursor to produce phosphonic acids by oxidation (Figure 35).

The oxidation of phosphinic acid is readily achieved in the presence of DMSO [26] and a catalytic amount of iodide as exemplified by the synthesis of compound 127 by oxidation of 126 [240] (Figure 36A). This method was also applied to prepare 1-aryl-1-hydroxymethylphosphonic acid [241]. HgCl$_2$ [242] or bromine water are also efficient for the oxidation of phosphinic acid. As an example, the phosphinic acid 128 (Figure 36B) which is prepared by the addition of hypophosphorous acid on...
imine, was converted quantitatively in α-amino phosphonic acid 129 with bromine water [243] (Figure 36B). HgCl$_2$, despite its toxicity and environmental hazard is nevertheless an efficient reagent for the oxidation of phosphinic acid [244]. Diiodide in the presence of acid (HI) in water/ethanol solution yields the oxidation of the phosphinic acid 130 without oxidizing the thioether function [245] (Figure 36C). The Atherton–Todd conditions [246] (CCl$_4$, NEt$_3$, H$_2$O) in the presence of water was also applied for the oxidation of phosphonic acid [247]. The oxidation of the phosphinic acid into phosphonic acid can be achieved with air or by a catalytic process involving the presence of palladium salts [236,248]. For instance, Kafarski et al. reported an efficient procedure that consisted in transforming the H-phosphinic acid into trivalent trimethylsilyl esters by reaction with hexamethyldisilazane followed by the oxidation with air followed by methanolysis [249]. A similar method that used a silylated intermediate and oxygen as oxidant was reported by Piettre et al. to produce alkyl-α,α-difluorophosphonic acid [250]. Ozone was also reported as oxidizing agent for the synthesis of phosphonic acid as exemplified by the synthesis of the diphosphonic acid 133 that was prepared from the phosphinic acid 132 [251] (Figure 36D).

### 8. Miscellaneous

If the most frequently used methods to prepare phosphonic acid have been reported above, others exist. In a non-exhaustive way of presentation, few of them are presented below.

Barton et al. reported a procedure to prepare phosphonic acids from carboxylic acids that made use of white phosphorus (P$_4$)
[252]. First, the carboxylic acid is esterified with $N$-hydroxy-2-thiopyridone in the presence of DCC to produce 134 (Figure 37). This compound 134 was added to $P_4$ solubilized in THF and was stirred for 30 minutes before replacing THF by DME. Then, H$_2$O$_2$ was added portion-wise and the solution was heated at reflux to produce the phosphonic acid 135. For some compounds, the reaction with H$_2$O$_2$ occurred at rt and then SO$_2$ is added to complete the oxidation reaction. The work-up is, however, tedious because some excess of $P_4$ must be removed without any contact with oxygen. This procedure was applied to natural compounds including lipophilic carboxylic acid (e.g., linoleic acid) or amino acid (e.g., L-2-amino-4-phosphonobutyric acid).

Red phosphorus is much easier to handle due to its polymeric nature that renders this compound much stable but also less reactive. Its reaction with benzaldehyde was reported to produce an $\alpha$-hydroxy-phosphinic acid intermediate that was converted to phosphonic acid in the presence of HI in an aqueous organic media at reflux (Figure 38). However, the benzylphosphonic acid (136) was formed simultaneously with phosphoric acid thus requiring a purification step [253].

More recently, the phosphonation of graphite was reported. First, a mechanochemical cracking yielded carboradical intermediates that were reacted with red phosphorus and then oxidized in the presence of air to produce graphene phosphonic acid 137 [116] (Figure 39). This material which is water soluble was used as a non-toxic flame-retardant.
Conclusion

Phosphonic acid is a functional group of interest for many current fields of research that include the development of bioactive compounds, medical imaging, material sciences or surface chemistry.

The most frequently used method to prepare phosphonic acids is stirring dialkyl phosphonates with concentrated HCl in aqueous solution at reflux. Despite these drastic conditions, this protocol is a method of choice to prepare phosphonic acids that are stable within acidic media and that are thermally stable. The advantage of this methodology is that it can be applied on large scale and, as exemplified in this review, was applied to a huge variety of compounds. According to this method, the purification step is rendered simple by the fact that the excess of reagents and side products (alkyl halide) are easily removed under vacuum. For the molecules featuring acid sensitive functionalities, the McKenna’s method, that starts by the reaction of dialkyl phosphonate with bromotrimethylsilane to produce a bis-silylated phosphonate under mild conditions (usually at 20 °C), is a method of choice. This intermediate (silylated phosphonate) is then quantitatively converted into phosphonic acid through hydrolysis or methanolysis. The reaction has a very broad variety of compounds. According to this method, the purification is also simple since the excess of reagent (phosphonic acid). According to the McKenna’s method the purification is also simple since the excess of reagent can be easily removed under vacuum.

Beside these two main methods to prepare phosphonic acids other direct methods can be selected. In that case the P–C bond is formed simultaneously with the production of a phosphonic acid functional group. In this way, the Moedritzer–Irani reaction is a well-documented and an efficient procedure. The limitation of this procedure can arise from the purification step which is mainly limited to crystallization.

Other methods are available as reported in this review; however, their use has a less broad scope but can be applied to prepare some specific phosphonic acids.

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