Halogen-free instinct flame-retardant waterborne polyurethanes: composition, performance, and application

Xuan Yin, Liqi Li, Haosheng Pang, Yunjun Luo and Bing Zhang

Ideal halogen-free instinct flame-retardant waterborne polyurethanes have high flame-retardant efficiency, environmental friendliness, fine compatibility, and good thermostability. Phosphorus flame-retardants are currently widely used in halogen-free instinct flame-retardant waterborne polyurethanes (HIFWPU), especially those with phosphorous–nitrogen co-structures. Phosphorous–nitrogen HIFWPU have become a hotspot because their co-structures provide higher flame-retardance as compared to waterborne polyurethanes. This review introduces three main types of HIFWPU based on composition, performance and application. HIFWPU not only have improved flame-retardance but also satisfy the various requirements for functionality. HIFWPU have been widely developed in textile, furniture, automobile, and aerospace applications.

Introduction

Polyurethanes (PU) have been extensively studied since their discovery in the 1950s by DuPont. With the increasing global concern regarding environmental protection, the application of solvent-based polyurethanes (PU) has been strictly limited due to its emission of volatile organic chemicals (VOCs). Therefore, the development of environmentally friendly PU production has to its emission of volatile organic chemicals (VOCs). Therefore, solvent-based polyurethanes (PU) has been strictly limited due to concern regarding environmental protection. The application of coatings provide high strength, rigidity, and glossiness. For instance, fabric WPU coatings can be modified and controlled for rheological properties. For instance, fabric WPU coatings can be modified and controlled for rheological properties.

Ideal halogen-free instinct flame-retardant waterborne polyurethanes have high flame-retardant efficiency, environmental friendliness, fine compatibility, and good thermostability. Phosphorus flame-retardants are currently widely used in halogen-free instinct flame-retardant waterborne polyurethanes (HIFWPU), especially those with phosphorous–nitrogen co-structures. Phosphorous–nitrogen HIFWPU have become a hotspot because their co-structures provide higher flame-retardance as compared to waterborne polyurethanes. This review introduces three main types of HIFWPU based on composition, performance and application. HIFWPU not only have improved flame-retardance but also satisfy the various requirements for functionality. HIFWPU have been widely developed in textile, furniture, automobile, and aerospace applications.

Introduction

Polyurethanes (PU) have been extensively studied since their discovery in the 1950s by DuPont. With the increasing global concern regarding environmental protection, the application of solvent-based polyurethanes (PU) has been strictly limited due to its emission of volatile organic chemicals (VOCs). Therefore, the development of environmentally friendly PU production has to concern regarding environmental protection. The application of coatings provide high strength, rigidity, and glossiness. For instance, fabric WPU coatings can be modified and controlled for rheological properties. For instance, fabric WPU coatings can be modified and controlled for rheological properties. Ideal halogen-free instinct flame-retardant waterborne polyurethanes have high flame-retardant efficiency, environmental friendliness, fine compatibility, and good thermostability. Phosphorus flame-retardants are currently widely used in halogen-free instinct flame-retardant waterborne polyurethanes (HIFWPU), especially those with phosphorous–nitrogen co-structures. Phosphorous–nitrogen HIFWPU have become a hotspot because their co-structures provide higher flame-retardance as compared to waterborne polyurethanes. This review introduces three main types of HIFWPU based on composition, performance and application. HIFWPU not only have improved flame-retardance but also satisfy the various requirements for functionality. HIFWPU have been widely developed in textile, furniture, automobile, and aerospace applications.

Ideal halogen-free instinct flame-retardant waterborne polyurethanes have high flame-retardant efficiency, environmental friendliness, fine compatibility, and good thermostability. Phosphorus flame-retardants are currently widely used in halogen-free instinct flame-retardant waterborne polyurethanes (HIFWPU), especially those with phosphorous–nitrogen co-structures. Phosphorous–nitrogen HIFWPU have become a hotspot because their co-structures provide higher flame-retardance as compared to waterborne polyurethanes. This review introduces three main types of HIFWPU based on composition, performance and application. HIFWPU not only have improved flame-retardance but also satisfy the various requirements for functionality. HIFWPU have been widely developed in textile, furniture, automobile, and aerospace applications.
The aim of these modifications is to enhance flame-retardance without influencing the WPU’s mechanical and thermal properties. Although one-component IFWPU has many advantages, such as a variable structure and simple fabrication process, it also has many disadvantages, such as low tensile strength, thermostability, and toughness. Two-component IFWPU has gradually begun to replace one-component IFWPU with high transparency, good waterproofness, and stable storage characteristics. Two-component IFWPU consists of flame-retardant waterborne polyol dispersions and a hydrophilic curing agent. For water-dispersing polyols, their structures are easy to control and design. Waterborne polyol dispersions have small particle sizes and good stability. Reactive flame-retardant units can be incorporated into the backbones and branched chains of PU during the synthesis process. Hyperbranched polymers are macromolecules with highly branched structures, which have been applied in adhesives, printing inks, and coatings due to their ecologically friendly, fabricability and stability, which can largely make up for the shortcomings of phosphorus-containing IFWPU. To satisfy the demands of applications, HIFWPU is often designed with many species, which are classified as flame-retardant coatings, flame-retardant coating adhesives, and flame-retardant paints. This paper mainly summarizes the research on HIFWPU based on modifications, compositions, performances, and applications.

**One-component HIFWPU**

**Soft segment modification**

Soft segment modification occurs when flame retardants react with other agents to obtain macromolecule polyether polyl- or polyester polyl-containing flame retardant elements. Then, they partially or completely replace the polyether polyl or polyester polyl to react with isocyanate and produce HIFWPU after emulsification. HIFWPU modified via soft segment modification possesses good flame-retardance with little effect on film properties. Soft segment modification can endow HIFWPU with a high content of flame-retardant elements and the high molecular weight of soft segments, which occupy the relatively high ratio of the PU molecular weight. Therefore, PU modified via the partial replacement of flame-retardant soft segments has a greater composition of flame-retardant and relatively unaffected film properties. Chen and Li, from our research group, successfully synthesized HIFWPU modified by organophosphorus soft segments, which improved the limiting oxygen index (LOI) to 32.0%. Two kinds of soft segment flame retardants are usually used: halogen-free polyphosphate (OP550, Clariant AG) and hydroxy-terminated organophosphate oligomer (HFPO, Fyroltex HP). OP550 is a new reactive phosphorus flame retardant that has high flame-retardant effectiveness and little effect on physical and mechanical properties. Its structure is shown in Scheme 1. WPU modified through OP550 is named WPU-OP. Generally, the synthesis uses a common method, as shown in Scheme 2, based on their previous work. Commonly, the stoichiometric ratio of \(-\text{NCO} \to -\text{OH} \ (\text{R})\) is generally from 1.3 to 1.7 because of the preparation technology. A solid content of 20–30% provides good film properties.

Characterizations of the structure and performance of one-component HIFWPU have certain essential aspects. The details are listed in Table 1. Commonly, it is necessary to characterize the structure and the fundamental and functional properties of HIFWPU. In WPU-OP systems, although organophosphorus decreases the decomposition temperature ($T_d$) value and thermostability of HIFWPU (Fig. 2), its carbon yield provides high flame-retardance. According to the carbonization

![Scheme 1](image)

**Scheme 1** Structure of OP550.
theory of Park, organophosphorus will decrease $T_d$ and accelerate the formation of char after being added to PU chains at high temperatures, which can effectively reduce the flammability and heat release rate (HRR) of PU burning. The research by Chen\textsuperscript{88} indicated that the movement of hard segments dispersed in a soft segment phase has a remarkable effect on the microstructure of PU. Therefore, an ethyl side chain enlarged the free volume of the soft segment micro-area after WPU was added to organophosphorus, which aggregated in a more regular short-range ordered structure that needed more energy to be damaged.\textsuperscript{64,88} As a result, the hard segment decomposition temperature ($T_{DH}$) moved to high temperatures as more OP550 was added (Fig. 2). The OP550 soft segment modifier has soft and tough characteristics, which decrease tensile strength and increase flexibility with a large volume of organophosphorus molecular structure via crippling the molecular chain rotation. The high content of organophosphorus in PU always implies high flame-retardance, even to obtain non-combustible material level (the best LOI and UL-94 of WPU-OP were 33.1% and V-0, respectively).\textsuperscript{89}

HFPO is another kind of soft segment modifier (Scheme 3). The synthesis is the same as for WPU-OP, except for the flame retardant. The $T_d$ moves to low temperature and its thermo-stability decreases, but the thermal decomposition rate is reduced to delay thermal decomposition (Fig. 3). In a soft modified system, the lower $T_{cs}$ means higher microphase separation.\textsuperscript{111,112} There are two reasons why $T_{cs}$ decreases with the increase in the HFPO content. One reason is that HFPO is a soft modifier, which reduces the content of hard segments in the PU system and lifts the restraint on the soft segments and enhances mobility at low temperatures.\textsuperscript{113} Another reason is that there is $\text{–O–CH}_3$ in the sidechains, which provides strong chain flexibility. Details of the mechanical properties and flame-retardance are listed in Table 2.

### Hard segment modification

Hard segment flame-retardant modification is like soft segment flame-retardant modification: WPU is incorporated with a modified hard segment flame-retardant, which means a flame-retardant unit is embedded into the hard segment of PU. Hard segment flame-retardant modification is also a simple method. The product not only has high flame-retardance but can also improve other properties like tensile strength. Common halogen-free phosphorus hard segment flame retardants include diethyl-$N,N$-bis(2-hydroxyethyl)amino-methylphosphonate (Fyrol-6) and $2,7$-bis($N,N$-dimethylamine)-1,6,3,8,2,7-octahydrodiazadiazole diphosphate (ODDP), which have reactive units that react with isocyanate. Fyrol-6 is a new reactive phosphorus flame retardant that has good antioxidation and water resistance properties because it can add phosphorus to the hard segments of PU chains via chemical reaction (Scheme 4a). The synthesis is like that in Scheme 2. Compared with soft segment modified systems, in WPU-F6 system, $T_d$ is reduced by Fyrol-6. However, the thermal decomposition rate ($\nu_d$) and char yield of WPU-F6 decreased and increased, respectively, by adding Fyrol-6. Although Fyrol-6, as a hard segment modifier, has high rigidity and strength,

### Table 1 Essential characterizations for one-component HIFWPU

| Performance or structure                          | Characterization                        | Reference |
|--------------------------------------------------|-----------------------------------------|-----------|
| Structure                                        | FTIR, $^1$H NMR, GPC                     | 90–93     |
| Particle size and emulsion stability             | Nanosizer                                | 94 and 95 |
| Viscosity and rheological properties             | Viscometer                               | 94 and 96 |
| Tensile strength and elongation at break         | Lecronic and mechanical testing machine  | 97–99     |
| Thermal properties                               | DSC, DMA, TGA, TGA-FTIR                 | 80 and 100–103 |
| Flame-retardance                                | SEM, EDS, UL-94, LOI, FTIR, cone calorimeter | 104–107  |
| Water resistance                                | Water absorption, contact angle measurement | 108–110  |
Table 2 Tensile strength and flame-retardance of soft segment modification, hard segment modification, and soft-hard segment modification systems

| Properties (content of flame retardant) | WPU-OP | WPU-HF | WPU-F6 | WPU-OD | WPU-OPF6 | WPU-HFF6 | WPU-OPOD |
|---------------------------------------|--------|--------|--------|--------|----------|----------|----------|
| LOI (%)                               | 33.1   | 34.2   | 30.2   | 30.6   | 30.4     | 36.6     | 31.4     |
| (15 wt%)                              | 23.4   | 29.1   | 26.4   | 29.6   | 29.4     | 31.7     | 30.4     |
| (5 wt%)                               | V-0    | V-0    | V-0    | V-1    | V-0      | V-0      | V-0      |
| UL-94                                 | 2.1    | 10.4   | 12.6   | 9.3    | 14.8     | 7.5      | 5.0      |
| (15 wt%)                              | 12.6   | 17.8   | 8.2    | 2.1    | 16.1     | 12.0     | 9.3      |
| (5 wt%)                               |        |        |        |        |          |          |          |

The synthesis of HFPO is like that of OP550/ODDP system (WPU-OPOD) (5 wt%), which makes it a nonflammable material.

The synthesis of OP550, HFPO and Fyro-6 are all industrial products. ODDP (Scheme 4b) is a laboratory-made flame retardant, fabricated by Luo’s group led by Gu. The synthesis is like those of the above three systems. In the WPU modified by the ODDP (WPU-OD) system, the viscosity and particle size were larger but were still 1.5 mPa s and 220 nm, respectively, and the char yield of modified samples was 2.3-times that of non-modified samples. This means that the soft and hard segment co-modification modifier plays a more obvious role in the material carbonization during high-temperature decomposition. Adding OP550 also decreased the tensile strength and elongation at break, which is like the case for WPU-OP. The LOI reached 30.4%, making it a nonflammable material, which is a little more than that of WPU-F6 but less than those of WPU-OP and WPU-HF.

Fig. 3 TGA thermograms for the poly(ECH-co-BO) series and poly-(FCGE)-BO copolymer series.© 2014 Springer Science+Business Media, LLC.

Details of the mechanical properties and flame-retardance are listed in Table 2.

Soft-co-hard segment modification

To further improve flame-retardance without affecting mechanical properties, soft and hard segment co-modification was developed. Three modified systems are introduced in this section: the OP550/Fyro-6 system (WPU-OPF6), HFPO/Fyro-6 system (WPU-HFF6), and OP550/ODDP system (WPU-OPOD). The synthesis of co-modification is different from that of single modification (Scheme 2). R generally ranges from 1.3 to 1.7 because of the preparation technology. The solid content is often 20–30% to obtain good film properties. As a result, the variation of the co-modification system is like that of the single system. $T_d$ decreased with the addition of OP550 but the char yield of modified samples was 2.3-times that of non-modified samples. This means that the soft and hard segment co-modification modifier plays a more obvious role in the material carbonization during high-temperature decomposition. Adding OP550 also decreased the tensile strength and elongation at break, which is like the case for WPU-OP. The LOI reached 30.4%, making it a nonflammable material, which is a little more than that of WPU-F6 but less than those of WPU-OP and WPU-HF.

The synthesis of WPU-HFF6 is like that of WPU-OPF6. The $T_{max}$ was reduced remarkably with the addition of HFPO. Meanwhile, HFPO delayed $v_d$ but reduced thermostability. The char yield increased sharply from 0.4% to 14.0% due to the addition of organophosphorus. Adding HFPO also decreased the tensile strength and elongation at break, which is like the case for WPU-HF. This was because of the large content and volume of soft segments. The best LOI was 37.1%, which is characteristic of a non-combustible material. This value is much higher than those of WPU-HF and WPU-F6, which means that the soft segment flame-retardant modifier plays an
important role in improving flame-retardance. The synthesis of the final co-modification system is also like that of WPU-OPF6, and the variation in thermal properties is also like those of the above two systems. The char yield increased drastically from 4.84 to 14.03%. OP550 and ODDP also delayed $\nu_d$ in HIFWPU systems, as shown in Fig. 5. The LOI can reach 31.6%, which is 1.0% larger than that of WPU-OD. However, the modification efficiency of ODDP is a little lower than that of Fyrol-6. Fyrol-6 has more organophosphorus than ODDP at the same additive amount.

Summary

To clearly show the key values of each system, a list is provided in Table 2. Although the hard segment flame-retardant modifier can improve the tensile strength, the elongation at break also decreases. In terms of flame-retardance, the LOI of soft segment modified systems is commonly larger than that of hard segment modified systems. Meanwhile, the LOI of soft and hard segment co-modified systems are also higher than that of hard segment modified systems. These indicate that the soft segment flame-retardant modifier plays an important role in improving flame-retardance because it has more organophosphorus than the hard segment flame-retardant modifier. The P-C from the soft and hard segment flame-retardant modifiers produces stable polyphosphoric acid, which forms a thermal insulation layer that protects the internal material to reduce weight loss in PU at high temperatures. Usually, more organophosphorus is beneficial for forming P-C layers to obtain better flame-retardance.

Two-component HIFWPU

Lu (one of Luo’s colleagues) and the authors used methoxy-polyethylene glycols (MPEG) and hexamethylene diisocyanate (HDI) trimer to prepare hydrophilic curing agents containing ether bonds.119 Lu further used toluene diisocyanate (TDI), DMPA, and dibromoneopentyl glycol (DBNPG) to synthesize waterborne flame-retardant hydroxyl prepolymers that were then reacted with a hydrophilic curing agent to obtain two-component HIFWPU. The hydrophilic curing agent improved the thermostability, tensile strength, and water resistance of the two-component WPU. Meanwhile, the LOI was over 28.0% with the addition of 10 wt% DBNPG. In our previous research, various kinds of two-component HIFWPU and hydrophilic curing agents have been investigated, which have been introduced in the review.123,124 The mixed emulsion was subjected to high-speed shear mixing or ultrasonic vibration to gain a two-component HIFWPU (Scheme 5).

Both Fyrol-6 and OP550 promote the degradation of WPUs and increase the production of carbon residue. The compositions of HIFWPU modified by Fyrol-6 and OP550 have been displayed in ref. 34. For consistency with previous research, the results of 2K-WPU1 (without being modified by flame-retardant hydrophilic curing agent) and 2K-WPU2 (modified by flame-retardant hydrophilic curing agent) are compared in Table 3.125,126 The flame-retardance and thermostability also improved because the flame-retardant hydrophilic curing agent decomposes to emit SO$_2$. TGA-FTIR analysis showed that the SO$_2$ combines with phosphoric acid and polyphosphoric acid to play a concerted catalytic role that improves the flame-retardance of the two-component WPU during heating. Subsequently, a functional two-component HIFWPU was researched by adding nano-TiO$_2$ to further improve flame retardance. Consequently, the LOI and UL-94 reached 30.5% and V-0, respectively. TGA and TGA-FTIR indicated that nano-TiO$_2$ can restrain thermal decomposition to improve flame-retardance and combine with phosphoric acid and polyphosphoric acid to provide synergistic flame-retardance.125,126 Meanwhile, nano-TiO$_2$ also provides ultraviolet radiation resistance and visible light-shielding effects because of its light-influencing characteristics.127–130

Araç, from Turkey, prepared two-component HIFWPU with various contents of phosphorous compound and N3300 curing agent.130 He found that the LOI could reach 29.0%. The Universitat Rovira i Virgili in Spain prepared two-component HIFWPU via Fyrol-6;23 researchers also synthesized an epoxy system IHPO-DGEBA via IHPO and DGEBA. Northwest Agricultural and Forestry University also prepared a two-component HIFWPU via Fyrol-6.a A series of hydroxy-terminated PU dispersions were prepared through 4,4-diphenyl-methylene-diisocyanate, polypropylene glycol, and hydrophilic-modified polyisocyanate curing agent. The LOI and UL-94 were 28.1%...
and V-2, respectively, with a 12.0% content of Fyrol-6 (PU3). The burning rate was restrained after adding Fyrol-6 with an increasing char yield of 19.9% (Fig. 6). Besides, the tensile strength, modulus, and impact strength were all improved due to the addition of Fyrol-6.

Zhang from Sichuan University prepared a flame-retardant chain extender and flame-retardant polyols,75,110 which were applied in a two-component HIFWPU. To balance the flame-retardance, thermostability, and mechanical properties, FRWPU was synthesized using the cyclic phosphoramidate lateral group bearing diol (PNMPD) and a silane coupling agent. He found that the tensile strength increased to 6.1 MPa with the addition of 12.0% PNMPD and 60% silane coupling agent. Meanwhile, the LOI and UL-94 were 27.7% and V-0, respectively. Furthermore, the thermal release rate peak (PHRR), total heat release (THR), smoke production rate (SPR), and total suspended particulates (TSP) decreased to 36.0%, 42.9%, 40.1%, and 35.4%, respectively. According to thermostability and flame-retardance mechanisms, the condensed phase flame-retardance obtained from phosphorus and silane chains could restrain the emissions of heat and inflam mable gas and accelerate the formation of carbon layers containing –P=O–O–Si–. Zhang further used PNMPD to act as a chain extender to prepare PNWPU. He found that the thermostability and water resistance were improved but not the flame-retardance. The LOI and UL-94 were 27.2% and V-0, respectively, while the PHRR, THR, SPR, and TSP decreased to 44.7%, 39.0%, 42.9%, and 36.1%, respectively. Additionally, the surface of the condensed phase was covered with rugged and intumescent carbon layers containing abundant polyphosphoric acid, which could effectively impede the conduction of heat, oxygen gas, and volatile matter from the flame-retardant and shielding areas.

Hyperbranched HIFWPU

To further develop the HIFWPU, Luo’s group synthesized hyperbranched HIFWPU and their blocking products via blending halogen-free flame-retardants and hyperbranched WPU. Tian synthesized acetic anhydride waterborne hyperbranched PU (WHBPU) via IPDI, DMPA, and diethanol amine (DEOA).111 Acetyl-terminated waterborne hyperbranched PU (AWHBPU) and WHBPU have been successfully synthesized after acetic anhydride and stearic acid blocking, as proven via FTIR, 13C-NMR, GPC, and functional calculations.112 The viscosity and particle size of three kinds of hyperbranched WPU were all <7.0 mP s and 100 nm, respectively. The $T_d$ of WHBPU and AWHBPU was 183 °C, but that of stearic acid-terminated waterborne hyperbranched PU (SWHBPU) was 205.7 °C because the bond energy of stearic acid is lower.113 The $T_d$ of AWHBPU and SWHBPU were lower than that of SWHBPU due to acetic anhydride and stearic acid blocking WHBPU, which decreased the constraint of the hydrogen bonds from the hydroxyl groups. Therefore, three kinds of hyperbranched HIFWPU were prepared through the blending of hyperbranched WPU with its blocking products. As a result, the LOI remained at over 30.0% and even the LOI of SWHBPU was 32.3%. AWHBPU and SWHBPU can promote phosphorus migration to the film surface to improve flame-retardance during the curing progress. Meanwhile, adding hyperbranched WPU can expand microphase separation to increase tensile strength, especially elongation at break in AWHBPU and SWHBPU.111

Tian prepared three kinds of hyperbranched WPU/organophosphorus hard segment HIFWPU (FWPU, FAPU, and FS PU). The variation in their viscosities was like that of non-mixed hyperbranched WPUs (Fig. 7). Although the thermostability was increased by adding hyperbranched WPU, the $T_d$ of soft segments was higher. Meanwhile, the decomposition was delayed by adding AWHBPU and SWHBPU. Besides, microphase separation and carbon yield were increased due to phosphorus enrichment by adding AWHBPU and SWHBPU. Their flame-retardance increased with the flame-retardant effectiveness. Additionally, Tian prepared three kinds of hyperbranched WPU/organophosphorus soft segment HIFWPU (OWPU, OAPU, and OSPU). The variation in their viscosities was like that of non-mixed hyperbranched WPU (Fig. 7). Although the thermostability was increased by adding hyperbranched WPU, the $T_d$ of soft segments was higher. Meanwhile, the decomposition was delayed by adding AWHBPU and SWHBPU. Besides, the crosslinking degree and carbon yield were higher due to the phosphorus enrichment of WHBPU and SWHBPU. For comparison with the six kinds of hyperbranched WPU/organophosphorus soft segment HIFWPU (FWPU, FAPU, and FS PU), the variation in their viscosities was like that of non-mixed hyperbranched WPU (Fig. 7). Although the thermostability was increased by adding hyperbranched WPU, the $T_d$ of soft segments was higher. Meanwhile, the decomposition was delayed by adding AWHBPU and SWHBPU. Besides, the crosslinking degree and carbon yield were higher due to the phosphorus enrichment of WHBPU and SWHBPU.
HIFWPU above, Tian synthesized hyperbranched WPU/compound HIFWPU (TWPU, TAPU, and TSPU). The variations in their viscosity, mechanical properties, and water resistance were like those in non-mixed hyperbranched WPU. The flame retardance was maintained at a high level, and WBHPU and SWHBPU can maintain flame retardance in different systems.

In summary, WBHPU and SWHBPU can maintain flame retardance in various systems. The variations in viscosity, mechanical properties, and water resistance of coated systems were consistent with those of the non-coated polyester Oxford tent cloth. The flame-retardance and other properties of hyperbranched HIFWPU are largely dependent on the structure of hyperbranched WPU and the type of flame retardant used.

Other types of HIFWPU

Besides one-component HIFWPU, two-component HIFWPU, and hyperbranched HIFWPU, there are other types of HIFWPU prepared through special techniques. Feng from Luo’s group prepared cross-linked prepolymers using polyester triols, polyether glycol, a hydrophilic chain extender, and diisocyanate. Then, Feng134 used OP550 and an oleophilic chain extender to modify the cross-linked prepolymer to obtain high-hydrostatic HIFWPU. The emulsion had a good appearance and stability after adding 12 wt% of OP550. Although the $T_d$ decreased, the carbon yield increased because the phosphate ester binds with low energy.38,103 The high hydrostatic HIFWPU had good compatibility, and a tensile strength and LOI of 26.5 MPa and 28.4%, respectively. Besides, the flame-retardant mechanism of high hydrostatic HIFWPU was similar to that of two-component HIFWPU.74 The particle size and zeta potential of the post-chain extension of HIFWPU were below 100 nm and ~30 μm, respectively. Meanwhile, the thermostability decreased because brominated polyphenyl ether (BPPO) accelerated the decomposition of PU chains. BPPO was decomposed to produce phosphoric acid, which catalyzed the decomposition of soft and hard segments; BPPO also increased the carbon yield and compatibility. The tensile strength and LOI were improved to 26.0 MPa and 28.8%, respectively, by adding BPPO. PHTP and THR were reduced to 65% and 26.3%, respectively. The increasing magnesium hydroxide sulfate hydrate (MHS) played the role of “diluent” in fire to enhance the fire performance index (FPI).135 The gas-phase and coagulation-phase flame-retardance of BPPO are key to promoting the formation of carbon and the release of nonflammable gases (Fig. 8).

Chem-physical flame retardants are always reactive inorganic nanomaterials (such as nano-montmorillonite, nanometer silicon dioxide, and nanosilicate) modified by chemical reactive groups.116 Usually, reactive inorganic nanomaterials provide characteristics like rigidity, heat resistance, and chemical resistance to WPU. Besides, some reactive inorganic nanomaterials also provide flame-retardance to WPU. So far, intumescent flame retardants have attracted more attention for use in high-transparency coatings because of the non-addition of another carbon source, which can reduce solid addition.137–140 Ammonium polyphosphate (APP-II) is a halogen-free inorganic flame retardant with N and P, which has a high polymerization degree, good thermostability, low hygroscopicity, and is green.67 However, APP-II has many disadvantages; for example, flame-retardant coated textiles prepared via APP-II show whitening, frosting, and low strength. Zhou added ethyl orthosilicate and methyltrimethoxysilane as precursors to cover an APP-II carrier to obtain synergistic flame-retardance through the sol–gel method. Consequently, the improved flame-retardant coated textile did not appear to have frosting and had good application prospects. Zeng et al. also used APP-II to prepare a flame-retardant coating adhesive.141 Besides phosphorus-containing polymers, antimony oxide and chlorocyclic triphosphoride are popular in the synthesis of intumescent HIFWPU. This kind of HIFWPU has many functional properties in addition to flame-retardance, like a matte appearance, water resistance, and zero formaldehyde release. The flame-retardant mechanism of intumescent HIFWPU is as follows. Tripolycyanamide and its salt belong to nitrogen or N–P flame retardants, which mainly consist of an air source and acid salt. Most current intumescent flame retardants contain N or P as a flame-retardant activity component.142 The advantages of intumescent flame retardants are low toxicity, corrosion, and smoke production. Most flame retardants containing tripoly- cyanamide and its salt have intumescent char formation and release CO$_2$, N$_3$, and NH$_2$ without halogen hydrate during heating. Meanwhile, the $T_o$ of the flame retardant is >250 °C. Halogen-free intumescent HIFWPU has N-P synergy flame-retardance143 and a condensed-gas phase because phosphate is the main part of the intumescent flame retardant.

Applications

Currently, HIFWPU are mainly applied as coatings for textiles, leather, and furniture to provide flame-retardance and other functional properties. In this section, we introduce application examples for textiles and wood (most is research from Luo’s group). Chen prepared a terylene textile coated with HIFWPU modified through HFPO. As a result, the flame-retardance of the textile coating was improved because the LOI increased and the UL-94 reached B1.71 Although the color change was reduced to 1.87, the color fastness was maintained at level-1. Stiffness slightly increased although the wrinkle recovery angle slightly decreased. The wash resistance of the coated terylene textile was good in soap testing. Li also prepared a terylene textile coated with HIFWPU modified with Pyrol-6. The flame-retardance of
the textile coating maintained its primary color and soft hand-feel.\textsuperscript{144-146} The LOI increased from 14.6% to 20.3% and the UL-94 reached $B_1$ (Fig. 9a). There was no dark smoke and little smoke production during heating. The carbon layer of the nonmodified textile had large amounts of fluffy cavities after heating (Fig. 9b).\textsuperscript{147-149} The carbon layer of the modified textile had no cavities after heating and formed a smooth and dense surface to isolate $O_2$ and heat (Fig. 9c).

Gu found that polyester Oxford tent cloth modified with ODDP had a good adhesive force but a hard hand-feel.\textsuperscript{149} The LOI and UL-94 were 23.7% and $B_2$, respectively. Further, Gu found that the LOI of polyester Oxford tent cloth modified with Pyror-6 was 21.6%. Polyester Oxford tent cloth modified with OP550 and Pyror-6 had outstanding flame-retardance, with LOI = 25.2% and UL-94 = $B_1$.\textsuperscript{57} Furthermore, the polyester Oxford tent cloth modified with Pyror-6 had a good adhesive force and soft hand-feel, even though it had a C-level antistatic property.\textsuperscript{150,151} Zhu\textsuperscript{152} incorporated anthraquinone chromophore (DV26) and OP550 to synthesize flame-retardant and colorful WPU (DOWPU) coatings. He obtained a LOI of 29.7%. Meanwhile, the dry rubbing, wet rubbing and washing color-fastness of modified polyester fabrics were 4–5, 4–5 and 4, respectively (Fig. 10). Moreover, the LOI of polyester fabrics coated with DOWPU also reached 27.1%, which displayed excellent application properties of DOWPU. Ding modified montmorillonite (MMT) with OP550 to obtain high-quality and monolayered nanosheets.\textsuperscript{153} The high-quality and monolayered nanosheets were then added to the \textit{in situ} polymerization to prepare environment-friendly WPU (OPWPU) coating\textsuperscript{154} and low gloss WPU (OWPU) coatings.\textsuperscript{155} The 60° gloss of coated polyvinyl chloride (PVC) leather was below 0.3 GU and 4.6 GU, respectively. Meanwhile, the LOI was over 18% after being coated. Feng used high hydrostatic HIFWPU to coat the Oxford cloth and then washed the Oxford cloth. Feng found that the LOI and UL-94 were 23.4% and $V-1$, respectively, after soaping.\textsuperscript{156} The water pressure resistance increased from 5.9 MPa to 9.7 MPa after soaping, which showed brilliant water pressure resistance in the textile. The stiffness, color change, and wrinkle recovery angle of the Oxford cloth were improved. A wood WPU varnish modified with BPPO was prepared by Wu. As a result, table-dry performance, water resistance, solvent resistance, abrasion resistance, and hardness were all improved.\textsuperscript{157} Then, Wu used wood WPU varnish modified with BPPO to coat the surface of polyester Oxford tent cloth. The LOI and UL-94 were 26.3% and $B_1$, respectively. Smoke production and toxic pollutants decreased during heating. Color change, stiffness, and breaking strength were improved by adding BPPO.

The author also prepared halogen-free two-component WPU varnishes by adding Pyror-6 or OP550. The results are listed in Table 4, which shows that the pencil hardness was significantly improved. In particular, the pencil hardness reached 2H by adding Pyror-6. Additionally, square impact, backswing, flexibility, adhesion, and solvent resistance were maintained at the highest level according to Chinese standards.\textsuperscript{158}

Hyperbranched HIFWPU can improve the hydrostatic pressure resistance, stiffness, tongue-tearing strength, and wrinkle-recovery angle of textiles.\textsuperscript{132} The after-flame time of the coated textile was still 0 s. The coating textile had good soap-to-wash resistance after adding hyperbranched HIFWPU. Tian prepared three kinds of hyperbranched WPU/organophosphorus hard segment HIFWPU (FWPU, FAPU, and FSPU), which were used to coat polyester Oxford tent cloth; the flame-retardance was slightly affected. WBHPU and SWHBPU can maintain flame-retardance in Oxford tent cloth. Additionally, Tian prepared three kinds of hyperbranched WP/organophosphorus soft segment HIFWPU (OWPU, OAPU, and OSPU), which were also used to coat polyester Oxford tent cloth. The flame retardance was slightly affected, and WBHPU and SWHBPU maintained flame retardance in the systems. Finally, FWPU, FAPU, and FSPU were used to coat polyester Oxford tent cloth. The flame-retardance was affected slightly.

Xu used intumescent flame retardants to prepare flame-retardant coating adhesives. The flame-retardant coating adhesive was a novel, halogen-free, green, flame-retardant coating adhesive, which had low smoke and toxicity and did not form melting drops during heating. Pan synthesized an APP-II-coated siloxane flame-retardant coating agent. The LOI and after-flame time of the coated textile were 44.3% and 0 s, respectively, after baking at 180 °C. Jiang prepared a flame-retardant coating adhesive by adding a cationic phosphorus flame retardant, antimony trioxide, and melamine.\textsuperscript{159} The

![Fig. 9](image_url) Images and SEM of flame-retardant Dacron after vertical burning: (a) textile after vertical burning; and SEM images of the surfaces of the (b) nonmodified Dacron and (c) modified Dacron.\textsuperscript{145,146} 1994–2016 China Academic Journal Electronic Publishing House.

![Fig. 10](image_url) Images of coated polyester fabrics.\textsuperscript{157} 2019 Elsevier B.V.
flame-retardant coating adhesive solved the problems of a non-durable flame-retardant effect, heavy losses, and easy color change during burning. Furthermore, the flame-retardant effect, smoke density, and smoke toxicity of the flame-retardant coating adhesive all reached the standard of non-combustibility, so it could be applied as seating fabric in aircraft and high-speed trains.

HIFWPU not only has good flame-retardancy and environmental protection but also has many functionalities such as luminescence, dyeing and so on. However, the versatility of HIFWPU urgently needs to be improved and its applications expanded in textile, wood, automotive and other fields. This is based on the in-depth discussion on HIFWPU, in order to better face the applications. The next step can use different physical or chemical methods to give HIFWPU more functionalities, which can balance mechanical properties, thermostability and flame-retardancy. Details are as follows: (1) to expand the synthesis method, the effective flame-retardancy of HIFWPU will be improved by designing and synthesizing flame retardants with different structure types. (2) HIFWPU can be endowed with properties such as anti-corrosion, super hydrophobicity, biological anti-bacterial and biodegradability, to meet the needs of broader application fields. (3) The thermostability and mechanical properties of HIFWPU can be improved by adding a matrix containing fluorine and a rigid large ring.

Conclusions

Different materials and applications have different requirements for HIFWPU, which means the matching demands for collecting halogen-free flame retardants. The ideal halogen-free flame-retardant waterborne PU has high flame-retardant efficiency, environmental friendliness, fine compatibility, and good thermostability and photostability. It also has a simple fabrication process, readily-available raw materials, and low cost. Hardly any HIFWPU can satisfy all the above conditions. Generally, the properties of WPU are reduced by adding too many nanometre-sized inorganic fillers. Although HIFWPU modified with P-N flame retardants are non-toxic, low smoke, and non-polluting, they are too expensive. Intumescent flame retardants can endow WPU with excellent properties but their surfaces need to be modified to prepare stable emulsions. Nowadays, HIFWPU not only have improved flame-retardance but also satisfy various requirements, which endow them with good potential for development. In the future, HIFWPU will occupy the main market in the surface finishing field and will be able to replace traditional products such as halogen-containing FWPU, solvent-based PU and blending-FWPU. Moreover, HIFWPU will be further developed due to their controllable molecular chain structure and excellent biocompatibility, which facilitate creative synthesis, novel modification and high-performance products.

Author contributions

X. Y.: visualization, conceptualization, writing-reviewing and editing. L. L. investigation. H. P.: writing-original draft preparation. Y. L.: project administration. B. Z.: supervision.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work is funded by the National Natural Science Foundation of China under Grant No. 51905295, Talent Introduction Program of BUCT under Grant No. buctrc202101 and the National Key Research and Development Program of China under Grant No. 2016YFC0204400.

Notes and references

1 K. L. Noble, Prog. Org. Coat., 1997, 32, 131–136.
2 W. J. Macknight and M. Yang, J. Polym. Sci., Polym. Symp., 1973, 42, 817–832.
3 D. E. Fiori, Prog. Org. Coat., 1997, 32, 65–71.
4 C. E. Tas, E. Berksun, D. Koken, S. Unal and H. Unal, ACS Appl. Polym. Mater., 2021, 3, 3929–3940.
5 N. Ballard, ACS Appl. Polym. Mater., 2020, 2, 4045–4053.
6 D. A. Ley, D. E. Fiori and R. J. Quinn, Prog. Org. Coat., 1999, 35, 109–116.
7 T. Hisakazu, S. Yasuyuki and Y. Fumio, Colloids Surf., A, 1999, 153, 597–601.
8 R. Pires and H. J. Laas, Surf. Coat. Int., Part B, 2002, 85, 169–242.
9 O. L. Figovsky, L. Shapovalov and O. Axenov, Surf. Coat. Int., Part B, 2004, 87, 71–148.
10 C.-H. Yang, L.-W. Chong, L.-M. Huang, Y.-L. Lee and T.-C. Wen, Mater. Chem. Phys., 2005, 91, 154–160.
11 Y. Wang, J. Deng, J. Zhao and H. Shi, Prog. Org. Coat., 2020, 148, 105845.
12 Y. Wang, J. Deng, J. Zhao and H. Shi, Polym. Adv. Technol., 2020, 32, 153–164.
130 K. Lee, K. Sato and A. R. Mohamed, Mater. Lett., 2016, 163, 240–243.
131 X. Tian, J. Li and Y. Luo, Trans. Beijing Inst. Technol., 2012, 32, 1195–1199.
132 X. Tian, J. Li, Y. Luo and X. Li, Polym. Mater. Sci. Eng., 2013, 29, 19–22.
133 F. Bao and W. Shi, Prog. Org. Coat., 2010, 68, 334–339.
134 J. Feng, Z. Ge, C. Chai, S. Wang, D. Yu, G. Wu and Y. Luo, Prog. Org. Coat., 2016, 97, 91–98.
135 S. Fang, Y. Hu, L. Song, J. Zhan and Q. He, J. Mater. Sci., 2007, 43, 1057–1062.
136 J. W. Seo and B. K. Kim, Polym. Bull., 2005, 54, 123–128.
137 R. G. Puri and A. S. Khanna, J. Coat. Technol. Res., 2016, 14, 323–331.
138 H. Li, Z. Hu, S. Zhang, X. Gu, H. Wang, P. Jiang and Q. Zhao, Prog. Org. Coat., 2015, 78, 318–324.
139 Z. Wang, E. Han and W. Ke, Can. Metall. Q., 2013, 46, 89–94.
140 W. H. Awad and C. A. Wilkie, Polym. Adv. Technol., 2011, 22, 1297–1304.
141 J. Zeng, H. Chen, Y. Luo, J. Luo and S. Wang, Polym. Mater. Sci. Eng., 2010, 26, 121–124.
142 J. Wang, S. Yang, G. Li and J. Jiang, J. Funct. Polym., 2003, 16, 238–242.
143 G. Camino, L. Costa and E. Casorati, J. Appl. Polym. Sci., 1988, 35, 1863–1876.
144 F. Li, Y. Luo, X. Li and J. Li, Mater. Sci. Technol., 2011, 19, 36–40.
145 F. Li, Y. Luo, X. Li, X. Li and J. Li, Dyeing Finish., 2013, 39, 13–15.
146 F. Li, Y. Luo, X. Li and J. Li, J. Chem. Eng. Chin. Univ., 2018, 26, 716–720.
147 H. Yang and C. Q. Yang, Polym. Degrad. Stab., 2005, 88, 363–370.
148 S. Giraud, S. Bourbigot, M. Rochery, I. Vroman, L. Tighzert, R. Delobel and F. Poutch, Polym. Degrad. Stab., 2005, 88, 106–113.
149 E. D. Weil, R. H. Hansen and N. Patel, Prospective Approaches to More Efficient Flame-Retardant Systems, American Chemical Society, Washington, DC, 1990.
150 L. Gu and Y. Luo, Polym. Mater. Sci. Eng., 2015, 31, 1–5.
151 L. Gu and Y. Luo, Trans. Beijing Inst. Technol., 2016, 36, 435–440.
152 J. Zhu, J. Li, W. Cai and Y. Luo, Prog. Org. Coat., 2020, 140, 105525.
153 Z. Ding, J. Li, B. Zhang and Y. Luo, Nanoscale, 2020, 12, 17083–17092.
154 Z. Ding, J. Li, W. Xin and Y. Luo, Appl. Clay Sci., 2020, 198, 105798.
155 Z. Ding, J. Li, W. Xin, G. Zhang and Y. Luo, Prog. Org. Coat., 2019, 136, 105273.
156 J. Feng, Z. Ge, C. Chai, S. Wang, D. Yu, G. Wu and Y. Luo, Prog. Org. Coat., 2016, 97, 91–98.
157 G. Wu, J. Li, C. Chai, Z. Ge, J. Lin and Y. Luo, RSC Adv., 2015, 5, 97710–97719.
158 D. A. Ley, D. E. Fiori and R. J. Quinn, J. Coat. Technol., 2000, 72, 63–69.
159 Z. Jiang, F. Ge, Z. Cai, L. Zeng, G. Chen and G. Chen, Dyeing Finish., 2008, 22, 6–9.