

PHARMACEUTICAL COMPOSITION, AND PREPARATION METHOD THEREFOR AND APPLICATION THEREOF

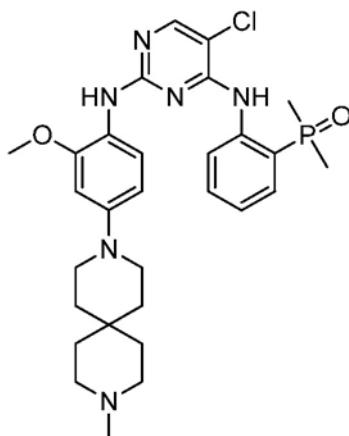
The present application claims priority of Chinese Patent Application No. 202110757316.5, filed before the CNIPA on July 5, 2021, titled “Pharmaceutical composition of spirocyclic aryl phosphorus oxide and preparation method thereof”, and Chinese Patent Application No. 202210660192.3, filed before the CNIPA on June 13, 2022, titled “A pharmaceutical composition, and preparation method therefor and application thereof”, which are incorporated herein by reference in their entirety.

TECHNICAL FIELD

The present invention belongs to the technical field of pharmaceutical chemistry, and relates to a pharmaceutical composition and preparation method and use thereof.

BACKGROUND

Spirocyclic aryl phosphorus oxide, with a structure represented by the following formula, is chemically named as: (2-((5-chloro-2-((2-methoxy-4-(9-methyl-3,9-diazaspiro[5.5]undecan-3-yl)phenyl)amino)pyrimidin-4-yl)amino)phenyl)dimethylphosphine oxide, which has a molecular weight of 569.08, and is a highly selective anaplastic lymphoma kinase (ALK) inhibitor. Its inhibitory effect on the enzyme activity of major kinase targets including ALK wild type, ALK mutant (ALK^{L1196M}, ALK^{C1156Y}) and EGFR mutant EGFR^{T790M/L858R} were determined at molecular level. Spirocyclic aryl phosphorus oxide showed similar or slightly better inhibitory activity to 4 major kinase targets compared with AP26113. Low inhibitory activity to other related serine, threonine and tyrosine proteases except EGFR wild type was observed in the detected concentration range, showing high selectivity to kinase. According to mechanism of action and results of preclinical trials, spirocyclic aryl phosphorus oxide can be used in the treatment of ALK positive non-small cell lung cancer patients with progressive disease after treatment with Crizotinib or unable to tolerate Crizotinib, or ALK positive non-small cell lung cancer patients who have not used ALK inhibitors.



Spirocyclic aryl phosphorus oxide is white or off-white powder with almost no hygroscopicity, and is easily soluble in dichloromethane, sparingly soluble in methanol, slightly soluble in water, ethanol or acetonitrile, and practically insoluble, or insoluble in 1mol/L sodium hydroxide solution. However, spirocyclic aryl phosphorus oxide is easily soluble in 1mol/L hydrochloric acid solution and is a compound of high solubility. The solubility of the drug directly influences the dissolution rate of the composition containing the drug, which in turn influences the bioavailability of the pharmaceutical composition. Therefore, how to improve the dissolution rate of pharmaceutical compositions containing spirocyclic aryl phosphorus oxide in solvents other than hydrochloric acid has become an urgent problem to be solved.

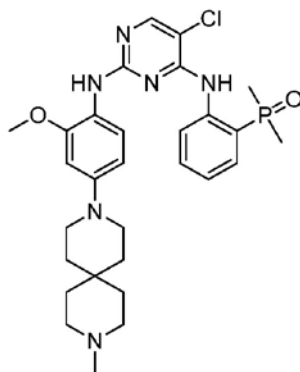
SUMMARY

It is known in the art that the particle size of drugs generally has no significant effect on drug dissolution behavior. However, the inventors of the present application found that although the particle size of spirocyclic aryl phosphorus oxide has a significant effect on the drug dissolution in vitro, too large or too small particle size will lead to a decrease in dissolution of spirocyclic aryl phosphorus oxides. Only by controlling the particle size of spirocyclic aryl phosphorus oxide within a certain range, the drug dissolution can be ensured to meet the requirements. Therefore, in order to improve the dissolution rate of the spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof, the range of particle size should be properly controlled so as to effectively control the dissolution rate of the spirocyclic aryl phosphorus oxide in formulations containing spirocyclic aryl phosphorus oxide, and further improve the bioavailability of the spirocyclic aryl phosphorus oxide in vivo.

In view of this, the object of the present invention is to provide a pharmaceutical composition and preparation method and use thereof. The spirocyclic aryl phosphorus oxide or a

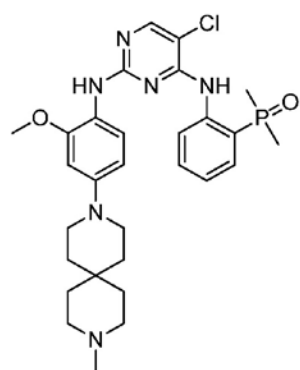
pharmaceutically acceptable salt thereof with a particular range of particle size can improve the dissolution rate of the pharmaceutical composition, and thus improve the bioavailability.

In a first aspect of the present invention, a spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof with a particle size distribution satisfying D90 in a range of 40.3 μm to 79.6 μm is provided, and the spirocyclic aryl phosphorus oxide has the following structure:



In a second aspect of the present invention, a spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof with the particle size distribution satisfying D50 in the range of 13.0 μm to 23.8 μm is provided, or a spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof with the particle size distribution satisfying D50 in the range of 13.0 μm to 18.2 μm is provided.

A third aspect of the present invention provides a pharmaceutical composition comprising a spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof with particular specific particle size, and a pharmaceutically acceptable carrier, wherein the spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof is used as an active ingredient, the particular particle size satisfies D90 in the range of 40.3 μm to 79.6 μm , and the spirocyclic aryl phosphorus oxide has the following structure:



In some embodiments of the present invention, the D50 of the spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof is in the range of 13.0 μm to 23.8 μm .

In some embodiments of the present invention, the D50 of the spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof is in the range of 13.0 μm to 18.2 μm . In some embodiments of the present invention, the spirocyclic aryl phosphorus oxide in the pharmaceutical composition of the present invention is in crystal form A.

In some preferred embodiments, the crystal form A of the spirocyclic aryl phosphorus oxide in the pharmaceutical composition of the present invention has characteristic peaks in the X-ray powder diffraction spectrum at 2θ degrees of $8.5\pm0.20^\circ$, $10.1\pm0.20^\circ$, $12.1\pm0.20^\circ$, $14.6\pm0.20^\circ$, $17.7\pm0.20^\circ$, $18.7\pm0.20^\circ$, $20.4\pm0.20^\circ$ and $21.2\pm0.20^\circ$, using Cu-K α radiation.

In some preferred embodiments, the crystal form A of the spirocyclic aryl phosphorus oxide in the pharmaceutical composition of the present invention has characteristic peaks in the X-ray powder diffraction spectrum at 2θ degrees of $8.5\pm0.20^\circ$, $9.3\pm0.20^\circ$, $10.1\pm0.20^\circ$, $12.1\pm0.20^\circ$, $13.7\pm0.20^\circ$, $14.6\pm0.20^\circ$, $15.7\pm0.20^\circ$, $16.5\pm0.20^\circ$, $17.7\pm0.20^\circ$, $18.7\pm0.20^\circ$, $20.0\pm0.20^\circ$, $20.4\pm0.20^\circ$, $21.2\pm0.20^\circ$, $23.6\pm0.20^\circ$, $25.1\pm0.20^\circ$ and $25.6\pm0.20^\circ$, using Cu-K α radiation.

In some more preferred embodiments, the crystal form A of the spirocyclic aryl phosphorus oxide in the pharmaceutical composition of the present invention has an X-ray powder diffraction spectrum substantially as shown in FIG.4.

In some embodiments, the pharmaceutically acceptable carrier comprises a filler, a disintegrant, an adhesive, a glidant and a lubricant.

In some preferred embodiments, the filler is selected from the group consisting of starch, mannitol, sorbitol, microcrystalline cellulose, lactose, pregelatinized starch and inorganic salts or a combination thereof, preferably a combination of microcrystalline cellulose and pregelatinized starch. The inorganic salts may include, but are not limited to, calcium sulfate (containing two molecules of crystal water), calcium hydrogen phosphate, calcium carbonate, etc.

In some preferred embodiments, the disintegrant is selected from the group consisting of dry starch, sodium carboxymethyl starch, hydroxypropyl starch, low substituted hydroxypropyl cellulose, croscopovidone and croscarmellose sodium or a combination thereof, preferably sodium carboxymethyl starch.

In some preferred embodiments, the adhesive is selected from the group consisting of distilled water, ethanol, starch paste, powdered sugar and syrup, hydroxypropyl methylcellulose,

povidone, hydroxypropyl cellulose, methyl cellulose, ethyl cellulose and sodium carboxymethyl cellulose or a combination thereof, preferably hydroxypropyl methylcellulose.

In some preferred embodiments, the glidant is selected from the group consisting of micronized silica gel and talc or a combination thereof, preferably micronized silica gel.

In some preferred embodiments, the lubricant is selected from the group consisting of magnesium stearate, hydrogenated vegetable oil, polyethylene glycol, dodecyl magnesium sulfate and sodium stearyl fumarate or a combination thereof, preferably magnesium stearate.

In some embodiments, the pharmaceutical composition comprising the spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof of a particular particle size of the present invention is an oral preparation, which is absorbed via gastrointestinal tract through oral administration.

In some preferred embodiments, the pharmaceutical composition of the present invention is a tablet.

In some preferred embodiments, the pharmaceutical composition of the present invention is a film-coated tablet.

In some embodiments, for the pharmaceutical composition of the present invention, the spirocyclic aryl phosphorus oxide or a pharmaceutical acceptable salt thereof accounts for 1.0% to 50.0% by weight in the composition.

In some preferred embodiments, in the pharmaceutical composition of the present invention, the following components each in percentage by weight are: 1% to 50% of spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof, 1% to 90% of the filler, 1% to 10% of the disintegrant, 0.1% to 1% of the glidant and 0.5% to 5% of the lubricant.

In some more preferred embodiments, in the pharmaceutical composition of the present invention, the following components each in percentage by weight are: 5.0% of spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof, 26.0% of microcrystalline cellulose, 59.5% of pregelatinized starch, 5% of sodium carboxymethyl starch, 3.0% of hydroxypropyl methylcellulose, 0.5% of micronized silica gel and 1% of magnesium stearate.

A fourth aspect of the present invention provides a preparation method of the pharmaceutical composition of any one in the above embodiments, which comprises the following steps:

1) fully mixing a spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof with a portion of internal excipients including a fillers, an adhesive and a disintegrant, and then granulating;

2) adding other excipients including a filler, a disintegrant and a lubricant, and then fully mixing; and

3) tableting and coating.

In some embodiments of the present invention, a pharmaceutical composition is provided, comprising an effective amount of a spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable carrier, a diluent and an excipient. Pharmaceutical compositions can be formulated for particular administration routes, such as oral administration, parenteral administration, and rectal administration, etc. Oral administration may include, but is not limited to, oral, sublingual and buccal administration, and oral administration may be administered by, but not limited to tablets (film-coated tablets), capsules (including sustained or timed release prescription), pills, powders, granules, elixir, tinctures, suspensions (including nano-suspensions, micron suspensions, spray-dried dispersants), syrup or emulsions; parenteral administration may include, but is not limited to, injection, infusion, and intranasal or other topical administrations. Specifically, it may be administered by, but not limited to, subcutaneous, intravenous, intramuscular or intrasternal injection; infusion techniques, such as aseptic injectable aqueous or non-aqueous solutions or suspensions; intranasal administration, including nasal mucosal administration, such as by inhalation spray; and topical administration, such as in the form of creams or ointments. Rectal administration is for example in the form of suppositories. The above administration routes can be administered separately, but usually with pharmaceutical carriers selected according to the selected administration route and standard pharmaceutical procedures.

A fifth aspect of the present invention provides a pharmaceutical composition of any one in the above embodiments or a pharmaceutical composition prepared by the method of any one in the above embodiments for use in the treatment of cancers. The cancers include non-small cell lung cancer, lymphoma, non-Hodgkin lymphoma, ovarian cancer, cervical cancer, prostate cancer, colorectal cancer, breast cancer, pancreatic cancer, glioma, glioblastoma, melanoma, leukemia, gastric cancer, endometrial carcinoma, lung cancer, hepatocellular carcinoma, gastric carcinoma, gastrointestinal stromal tumor (GIST), acute myeloid leukemia (AML),

cholangiocarcinoma, renal carcinoma, thyroid cancer, anaplastic large cell lymphoma, mesothelioma, multiple myeloma and melanoma.

In some embodiments, the cancer is non-small cell lung cancer.

In some embodiments, the above use is for ALK positive non-small cell lung cancer patients with progressive disease after treatment with Crizotinib or unable to tolerate Crizotinib.

In some embodiments, the above use is for ALK positive non-small cell lung cancer patients who have not used ALK inhibitors.

A sixth aspect of the present invention provides a method of treatment of cancer, which comprises administering a therapeutically effective amount of the pharmaceutical composition of any one in the aforementioned embodiments to the patient. The cancers include non-small cell lung cancer, lymphoma, non-Hodgkin lymphoma, ovarian cancer, cervical cancer, prostate cancer, colorectal cancer, breast cancer, pancreatic cancer, glioma, glioblastoma, melanoma, leukemia, gastric cancer, endometrial carcinoma, lung cancer, hepatocellular carcinoma, gastric carcinoma, gastrointestinal stromal tumor, acute myeloid leukemia, cholangiocarcinoma, renal carcinoma, thyroid cancer, anaplastic large cell lymphoma, mesothelioma, multiple myeloma and melanoma.

In some embodiments, the above cancer is non-small cell lung cancer.

In some embodiments, the above method is used for ALK positive non-small cell lung cancer patients with progressive disease after treatment with Crizotinib or unable to tolerate Crizotinib.

In some embodiments, the above method is used for ALK positive non-small cell lung cancer patients who have not used ALK inhibitors.

In the present invention, D50 refers to the particle size corresponding to the cumulative volume distribution percentage of spirocyclic aryl phosphorus oxides or pharmaceutically acceptable salts thereof reaching 50% measured from a small particle size; D90 refers to the particle size corresponding to the cumulative volume distribution percentage of spirocyclic aryl phosphorus oxides or pharmaceutically acceptable salts thereof reaching 90% measured from a small particle size.

Beneficial technical effects:

The present invention provides a pharmaceutical composition containing a spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof having a particular particle size range, which can increase the dissolution rate of the pharmaceutical composition and improve the bioavailability.

Unless otherwise stated, the following terms and phrases used herein are intended to have the following meanings. A particular term or phrase should not be considered indefinite or unclear in the absence of a particular definition, but should be understood according to its ordinary meaning.

The term “pharmaceutically acceptable” as used herein refers to those compounds, materials, compositions, and/or dosage forms which are, within the scope of sound medical judgment, suitable for contact with tissues of human beings or animals without excessive toxicity, irritation, allergic response, or other problems, or complications, and is commensurate with a reasonable benefit/risk ratio.

The term “pharmaceutically acceptable salt” refers to the derivatives prepared from the compounds of the present invention and relatively non-toxic acids or bases. These salts can be prepared during synthesis, separation and purification of compounds, or can be prepared solely by using the purified compound in the free form to react with suitable acids or bases. When the compound contains relatively acidic functional groups, it reacts with alkali metals, alkaline earth metal hydroxides or organic amines to obtain alkali addition salts, including cations based on alkali and alkaline earth metals, such as sodium, lithium, potassium, calcium, magnesium, etc., and non-toxic ammonium, quaternary ammonium and amine cations, including, but not limited to, ammonium, tetramethylammonium, tetraethylammonium, methylamine, dimethylamine, trimethylamine, triethylamine, ethylamine, etc. Pharmaceutically acceptable salt also covers salts of amino acids, such as arginate, gluconate, galacturonate, etc. When a compound contains relatively basic functional groups, it reacts with organic or inorganic acids to obtain acid addition salts, including inorganic salts such as hydrochloride, hydrobromide, nitrate, carbonate, bicarbonate, phosphate, monophosphate, dihydrogen phosphate, sulfate, hydrogen sulfate, hydroiodate, hydrobromide, metaphosphate and pyrophosphate, or organic salts including, for example, acetate, propionate, octanoate, isobutyrate, oxalate, malonate, succinate, suberate, sebacate, fumarate, maleate, mandelate, benzoate, chlorobenzoate, methyl benzoate, dinitrobenzoate, naphthoate, benzene sulfonate, toluenesulfonate, phenylacetate, citrate, lactate, maleate, tartrate, methanesulfonate, etc.

The term “pharmaceutically acceptable carrier” refers to a medium generally acceptable in the art for the delivery of bioactive agents to animals, particularly mammals, according to the administration and property of formulation, including, for example, adjuvants, excipients or vehicles, such as fillers, disintegrants, adhesives, glidants, lubricants, diluents, preservatives, flow modulators, wetting agents, emulsifiers, suspending agents, sweeteners, flavors, fragrances, antibacterial agents, antifungal agents, lubricants and dispersants. Pharmaceutically acceptable carriers are formulated according to a quantity of factors within the knowledge of those skilled in the art. Such factors include, but are not limited to, the types and properties of the active agents to be formulated, the subjects to be administered with the composition containing such active agents, the expected administration routes of the composition, and the indications of target treatment. Pharmaceutically acceptable carriers include both aqueous and non-aqueous medium, as well as a variety of solid and semi-solid formulations. In addition to active agents, such carriers include many different ingredients and additives, and such additional ingredients included in prescriptions for a variety of reasons (such as stabilizing active agents, adhesives, etc.) are well known to those skilled in the art. The term “excipient” generally refers to the carrier, diluent, and/or medium required for the formulation of an effective pharmaceutical composition.

The term “therapeutically effective amount” refers to a sufficient amount of the pharmaceutical composition of the present invention to treat disorders in a reasonable effect/risk ratio suitable for any medical treatment. However, it should be recognized that the total daily dosage of the composition of the present invention must be determined by the attending physician within the scope of reliable medical judgment. For any specific patient, the specific therapeutically effective dose level shall depend on a variety of factors, including the disorder to be treated and the severity of the disorder; the activity of the specific compound used; the specific composition used; age, weight, general health condition, gender and diet of the patient; the administration time, route and excretion rate of the specific compound used; treatment duration; drugs used in combination with or simultaneously with the specific compounds used; and similar factors known in the medical field.

BRIEF DESCRIPTION OF THE DRAWINGS

In order to explain the examples of the present invention and the technical solutions of the prior art more clearly, the following briefly introduces the drawings used in the examples and the

prior art. Obviously, the drawings in the following description are only some examples of the present invention, and for those skilled in the art, other drawings may be obtained based on these drawings without any creative efforts.

FIG. 1 shows the X-ray powder diffraction spectrum of the remaining solid spirocyclic aryl phosphorus oxide of samples 1-8 in Table 2 in suspensions during solubility test;

FIG. 2 shows the comparison of the dissolution curves of examples 1-4 and comparative examples 1-4 in phosphate buffer at pH 6.8;

FIG. 3 shows the comparison of the dissolution curves of examples 1-4, and comparative examples 1-4 in purified water with 0.5% Tween 20 added;

FIG. 4 shows the X-ray powder diffraction spectrum of crystal form A of spirocyclic aryl phosphorus oxide.

DETAILED DESCRIPTION

In order to make clear the purpose, technical solutions, and advantages of the present invention, the present invention is further described in detail with reference to the drawings and examples. It is obvious that the described examples are only a portion of the examples of the present invention but not all of them. Based on the examples of the present invention, all other examples obtained by those skilled in the art without creative effort will fall within the scope of the protection of the present invention.

In the example of the present invention, the crystal form of spirocyclic aryl phosphorus oxide is crystal form A, which is a known and commercially available raw material, or can be synthesized by or in accordance with a method known in the art.

Determination of solubility of spirocyclic aryl phosphorus oxide

The solubilities of spirocyclic aryl phosphorus oxides in buffers at different pHs were determined. Nine test samples of 4 mg were each weighed and added to sample bottles of 1.5mL, and then different solvents (0.1 mol/L HCl, 0.01 mol/L HCl, pH3.5 acetate buffer, pH4.5 acetate buffer, pH5.5 phosphate buffer, pH6.5 phosphate buffer, pH6.8 phosphate buffer, pH7.5 phosphate buffer and water) of 1 mL were added. According to the dissolution, the raw material compounds were continuously added to form a saturated solution. A magneton was added to the saturated solution and then stirred on a magnetic agitator (temperature was 37 °C, away from

light). After stirring for 24 hours, the samples were taken and centrifuged, and the residual solid samples in the lower layer were tested by X-ray powder diffraction (XRPD). The upper sample was filtered by a filter membrane, the final pH value of the filtrate was measured, and the saturated solubility of the compound was determined by high performance liquid chromatography (HPLC). The results of the determination are shown in Table 1, and the results of the XRPD are shown in FIG. 1.

Table 1 Determination results of solubility of spirocyclic aryl phosphorus oxides in buffers with different pH values

Solvents	Status (24 hrs)	Solubility (mg/mL)	Dose/solubility (mL, in 60mg specifications)	Conclusions	XRPD
0.1 mol/L HCl (pH 1.02)	yellow suspension	53.4	1.12	soluble	no change
0.01 mol/L HCl (pH2.02)	white suspension	6.9	8.69	slightly soluble	no change
pH 3.5 buffer (pH 3.53)	dark yellow solution	279.4	0.22	easily soluble	no change
pH 4.5 buffer (pH 4.49)	yellow suspension	108.0	0.56	easily soluble	no change
pH 5.5 buffer (pH 5.47)	white suspension	7.1	8.45	slightly soluble	no change
pH 6.5 buffer (pH 6.45)	white suspension	11.1	5.41	sparingly soluble	no change
pH 7.5 buffer (pH 7.53)	white suspension	0.85	70.59	very slightly soluble	no change
pH 6.8 buffer (pH 6.83)	white suspension	3.6	16.67	slightly soluble	no change
water (pH 6.33)	white suspension	0.06	>250	practically insoluble	no change

Note: the easily soluble, soluble, sparingly soluble, slightly soluble and very slightly soluble in Table 1 mean that 1g of spirocyclic aryl phosphorus oxides can be dissolved in 1 ml to less than 10 ml, 10 ml to less than 30 ml, 30 ml to less than 100 ml, 100 ml to less than 1000 ml,

1000 ml to less than 10000 ml of solvent, respectively, and practically insoluble means that 1g of spirocyclic aryl phosphorus oxides can not be completely dissolved in 10000 ml of solvent.

It can be seen from Table 1 that the solubility change of spirocyclic aryl phosphorus oxide in buffers at different pHs is related to pH, and the spirocyclic aryl phosphorus oxide can be soluble in 0.1mol/L hydrochloric acid solution, easily soluble in buffer at pH3.5 and buffer at pH4.5, slightly soluble in 0.01mol/L hydrochloric acid solution, buffer at pH5.5 and buffer at pH6.8, sparingly soluble in buffer at pH6.5, very slightly soluble in buffer at pH7.5, and practically insoluble in water. It can be seen from FIG. 1, the crystal form of spirocyclic aryl phosphorus oxide does not change in different solvents.

Preparation and process of pharmaceutical compositions containing spirocyclic aryl phosphorus oxide with different particle sizes

The content uniformity and in vitro dissolution rate of pharmaceutical compositions containing the spirocyclic aryl phosphorus oxide with different particle sizes were tested.

Table 2: Spirocyclic aryl phosphorus oxide samples with different particle sizes

Particle size	Sample 1	Sample 2	Sample 3	Sample 4	Sample 5	Sample 6	Sample 7	Sample 8
D90 (μm)	11.6	27.1	40.3	62	69	79.6	122	193
D50 (μm)	5.4	10.1	13.0	23.8	16.1	18.2	47.9	63.1
D10 (μm)	0.7	1.3	1.6	5.0	1.8	2.3	6.8	7.8

Table 3 Solid composition of spirocyclic aryl phosphorus oxide

Prescription	
Unit	mg/tablet
spirocyclic aryl phosphorus oxide	60.0
microcrystalline cellulose PH101	46.8
pregelatinized starch	110.4
sodium carboxymethyl starch	12.0
hydroxypropyl methylcellulose	7.2
silicon dioxide	1.2
magnesium stearate	2.4
Total	240

Examples 1-4

Preparation process:

Spirocyclic aryl phosphorus oxide samples 3-6 with different particle sizes in Table 2 were prepared according to the prescription in Table 3 and the following preparation steps to obtain different pharmaceutical compositions sample 3A, sample 4A, sample 5A and sample 6A:

adding spirocyclic aryl phosphorus oxides with different particle sizes, microcrystalline cellulose, pregelatinized starch, hydroxypropyl methylcellulose and half the prescription amount of sodium carboxymethyl starch to High shear mixer to mix for 5 min; and adding purified water to granulate with stirring for 8 min;

performing wet granulation by granulator, drying in fluidized bed and sieving by granulator;

adding the remaining sodium carboxymethyl starch and silicon dioxide to the particles and mixing; adding magnesium stearate, sieving, mixing, and tableting.

Comparative examples 1-4

Except that spirocyclic aryl phosphorus oxides were successively replaced with samples 1, 2, 7 and 8 shown in Table 2, the rest were the same as in example 1.

Determination of content uniformity and dissolution rate of pharmaceutical compositions in various examples and comparative examples

1) Detection of content uniformity

The content uniformity test of the pharmaceutical composition of each example and comparative example refers to the “Appendix XE content uniformity test method” of Chinese Pharmacopoeia 2015 edition, and the results are shown in Table 4.

Table 4 Content Uniformity

		Average content (%)	Content Uniformity (A+2.20S≤15.0)
Example 1	Sample 3A	100.0	4.7
Example 2	Sample 4A	100.1	6.1
Example 3	Sample 5A	98.8	5.9
Example 4	Sample 6A	99.7	4.5
Comparative example 1	Sample 1A	98.9	5.2

Comparative example 2	Sample 2A	99.8	4.6
Comparative example 3	Sample 7A	99.4	5.4
Comparative example 4	Sample 8A	99.0	4.3

According to the results of content uniformity, the content uniformity of the pharmaceutical compositions can meet the requirements of quality standards when the particle size of raw materials is not more than 200 μm and not less than 10 μm .

2) Dissolution detection

The dissolution performance of the pharmaceutical composition in each example and comparative example was tested according to the dissolution determination method (Chinese Pharmacopoeia 2015 edition IV 0512), and specifically, the rotational speed was 50 rpm, maintained at 37 ± 0.5 °C. The dissolution curve of the pharmaceutical composition was investigated, and the dissolution data are shown in Tables 5 and 6 below.

Table 5 Dissolution data of spirocyclic aryl phosphorus oxide solid composition in phosphate buffer at pH6.8

	Sample number	Comparison of dissolution curves (%)					
		5min	10min	15min	30min	45min	60min
Example 1	Sample 3A	40.5	80.7	91.2	96.8	98.2	99.4
Example 2	Sample 4A	42.3	82.9	92.6	97.6	99.7	99.7
Example 3	Sample 5A	43.6	83.1	94.7	97.8	99.2	99.6
Example 4	Sample 6A	42.7	82.6	93.4	96.1	98.7	99.3
Comparative example 1	Sample 1A	19.7	41.7	62.4	86.9	91.5	91.2
Comparative example 2	Sample 2A	26.9	52.5	71.2	88.6	93.4	92.1
Comparative example 3	Sample 7A	29.4	69.7	81.2	90.4	93.4	94.7
Comparative example 4	Sample 8A	37	66.5	75.9	86.1	89	91

Table 6 Dissolution data of spirocyclic aryl phosphorus oxide solid composition in purified water + 0.5% Tween 20

	Sample number	Comparison of dissolution curves (%)						
		5min	10min	15min	30min	45min	60min	90min
Example 1	Sample 3A	14.2	35.2	51	65.4	74.2	76.5	82.7
Example 2	Sample 4A	18.5	43.3	55.7	67.8	74.1	77.8	83
Example 3	Sample 5A	19.7	42.6	56.8	66.3	72.5	78.4	84
Example 4	Sample 6A	15.2	32.4	48.6	63.1	69.8	74.2	78.7
Comparative example 1	Sample 1A	12	29.2	46.7	68.3	74.8	77.7	82.4
Comparative example 2	Sample 2A	11.7	30.1	46.7	65.9	72.4	76.5	81.9
Comparative example 3	Sample 7A	11.5	30	39.3	50.6	55.8	59.2	64.1
Comparative example 4	Sample 8A	19.8	34.4	43	55.4	61.3	65.6	72.2

The dissolution results of Table 5 and FIG. 2 show that in phosphate buffer at pH6.8, when the D90 of spirocyclic aryl phosphorus oxide is in the range of 40.3 μm to 79.6 μm and the D50 is in the range of 13.0 μm to 23.8 μm , the dissolution results are substantially the same, and there is no significant difference. Compared with dissolution data of other particle sizes, spirocyclic aryl phosphorus oxides with D90 in the range of 40.3 μm to 79.6 μm and D50 in the range of 13.0 μm to 23.8 μm are dissolved out most rapidly from pharmaceutical compositions. When the particle size of D90 is less than 40.3 μm or D50 is less than 13.0 μm , the dissolution gradually becomes slower, and when D90 is more than 79.6 μm or D50 is more than 23.8 μm , the dissolution is also slower.

The dissolution results of Table 6 and FIG. 3 show that in the medium of purified water + 0.5% Tween 20, the dissolution rates of spirocyclic aryl phosphorus oxide are substantially the same when the D90 is in the range of 11.6 μm to 79.6 μm and the D50 is in the range of 5.4 μm to 23.8 μm , and the dissolution rate decreases when the particle size D90 of raw material is more

than 100 μm or the D50 is more than 40 μm . Referring to Tables 5 and 6, spirocyclic aryl phosphorus oxides have good dissolution performance in the medium of phosphate buffer at pH6.8 and purified water + 0.5% Tween 20 when the D90 is in the range of 40.3 μm to 79.6 μm and the D50 is in the range of 13.0 μm to 23.8 μm .

Investigation on the stability of pharmaceutical composition

1) Influencing factor test

The pharmaceutical composition prepared in Example 2 of the present invention was selected to investigate the generation of total impurities in the sample under the conditions of light, high temperature (60 °C or 40 °C) and high humidity (Rh90% $\pm$ 5% or Rh75% $\pm$ 5%). The calculation of total impurities (%) is obtained by determining the peak area ratio of the samples to be tested and the reference samples using LC.

Table 7 Data of influencing factors of spirocyclic aryl phosphorus oxide tablets

Influencing factors	Index	0 day	5 days	10 days	30 days
Light	Total impurity (%)	none detected	0.15	0.08	0.26
High temperature (60 °C)	Total impurity (%)	none detected	0.16	0.19	0.29
High temperature (40 °C)	Total impurity (%)	none detected	0.11	0.14	0.18
High humidity test (RH90% $\pm$ 5%)	Total impurity (%)	none detected	0.10	0.15	0.11
	Loss on drying (%)	3.3	7.8	7.8	8.7
High humidity test (RH75% $\pm$ 5%)	Total impurity (%)	none detected	0.15	0.10	0.11
	Loss on drying (%)	3.3	6.9	7.0	7.3

According to the results in the table, the related ingredients in the pharmaceutical composition in Example 2 increased slightly under the conditions of light and high temperature, but the content was low. Although the moisture increased a lot under high humidity, the related

ingredients remained substantially unchanged. Therefore, the pharmaceutical composition provided by the present application has good stability under the conditions of high temperature and high humidity.

2) Accelerated stability test

Sample 1A, sample 3A and sample 4A of pharmaceutical compositions prepared in Comparative Example 1, Example 1 and Example 2 of the present invention were selected to investigate the formation of related ingredients (single impurity and total impurity) in the samples at 40°C and Rh75% respectively.

Table 8 Accelerated stability data

Time	Sample 1A (with desiccant)			Sample 1A (without desiccant)			Sample 3A (with desiccant)			Sample 4A (with desiccant)		
	Impurity 1(%)	Other single impurity (%)	Total impurity (%)	Impurity 1(%)	Other single impurity (%)	Total impurity (%)	Impurity 1(%)	Other single impurity (%)	Total impurity (%)	Impurity 1(%)	Other single impurity (%)	Total impurity (%)
0 day	0.01	0.03	0.08	0.55	None detected	0.55	0.03	0.04	0.13	0.02	0.03	0.11
Accelerate for 1 month exposure	/	/	/	/	/	/	0.11	0.05	0.24	0.07	0.04	0.2
Accelerate for 1 month seal	0.01	0.03	0.13	0.56	0.05	0.61	0.07	0.04	0.23	0.06	0.04	0.16
Accelerate for 2 months	0.05	0.03	0.13	0.57	None detected	0.57	0.08	0.04	0.18	0.08	0.06	0.22
Accelerate for 3 months	0.09	0.04	0.17	0.58	None detected	0.58	0.1	0.05	0.2	0.09	0.05	0.21
Accelerate for 6 months	0.1	0.05	0.19	0.58	0.07	0.58	0.14	0.07	0.22	0.14	0.08	0.24

Note: the “/” in Table 8 indicates that there is no corresponding parameter.

Sample 1A (with desiccant), sample 3A (with desiccant) and sample 4A (with desiccant) were all added with desiccant in the process of product stability placement. During the process, the stability was substantially the same, and the growth trend of related ingredients was substantially the same. Sample 1A (without desiccant) was not added with desiccant during the placement process, the related ingredients were more in 0 day, but did not change much in the later stage. Compared with the stability data of sample 1A with and without desiccant, the results showed that the total impurity content was significantly lower after adding desiccant.

Clinical trials

1) Prescription

Raw materials	Sample 9A	Sample 10A
Unit	mg/tablet	mg/tablet
spirocyclic aryl phosphorus oxide (D90 = 62 μm , D50 = 23.8 μm , D10 = 5.0 μm)	30	60
microcrystalline cellulose	23.4	46.8
pregelatinized starch	55.2	110.4
sodium carboxymethyl starch	6.0	12.0
hydroxypropyl methylcellulose	3.6	7.2
micronized silica gel	0.6	1.2
magnesium stearate	1.2	2.4
weight of plain tablet	120	240
Opadry coating powder (white)	3.6	7.2

2) The preparation processes of pharmaceutical composition samples 9A and 10A is shown in Example 1.

3) Sample 9A and sample 10A were taken for clinical trials according to the following dosages. The data are shown in Table 9 below.

Table 9 Clinical trial data of spirocyclic aryl phosphorus oxide tablets

Administration dose	60mg	90mg	120mg	180mg
Administration mode	60mg/1 tablet	30mg/1 tablet + 60mg/1 tablet	60mg/2 tablets	60mg/3 tablets
C _{max} (nM)	274.32	356.47	489.22	369.00
T _{max} (hr)	1.328	1.685	1.808	2.288
T _{1/2} (hr)	25.56	21.70	21.87	23.53

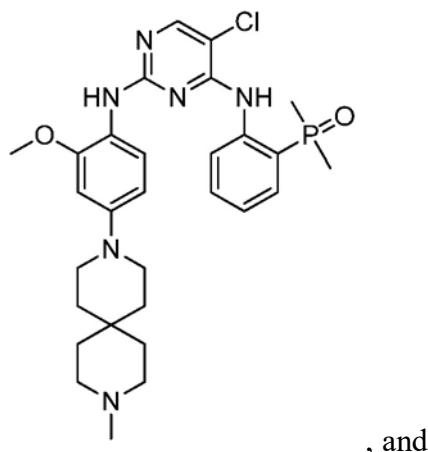
AUC _{0-t} (nM.hr)	3377.43	4852.29	7134.80	5835.07
AUC _{0-∞} (nM.hr)	3778.77	5575.31	7437.80	6463.01
MRT _{0-∞} (hr)	28.00	23.79	25.20	27.44

According to the analysis of clinical data, AUC_{0-∞} substantially increased linearly when the dosage was increased from 60 mg to 120 mg, and increased by less than 1.5 times when the dosage was increased from 60 mg to 90 mg. Relatively speaking, the bioavailability of 60 mg was slightly higher in vivo. However, there is no significant difference in bioavailability between 60 mg and 120 mg dosages, and there is no statistically significant differences. However, the bioavailability of 180 mg dose group was lower than that of 120 mg dose group, indicating that the bioavailability in vivo did not increase with the increase of dosage.

The method of the present invention has been described through the preferred embodiments, and relevant personnel can obviously change or appropriately modify and combine the method and application described herein within the content, spirit and scope of the present invention to realize and apply the present invention. It is particularly pointed out that although specific embodiments have been illustrated and described, all similar substitutions and modifications are obvious to those skilled in the art, and they are regarded as included in the present invention.

CLAIMS

1. A pharmaceutical composition, comprising a spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof as an active ingredient, and a pharmaceutically acceptable carrier, wherein, the spirocyclic aryl phosphorus oxide has the following structure:



the D90 of the spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof is in a range of 40.3 μm to 79.6 μm .

2. The pharmaceutical composition according to claim 1, wherein, the D50 of the spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof is in a range of 13.0 μm to 23.8 μm .

3. The pharmaceutical composition according to claim 1, wherein, the D50 of the spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof is in a range of 13.0 μm to 18.2 μm .

4. The pharmaceutical composition according to any one of claims 1-3, wherein, the spirocyclic aryl phosphorus oxide is in crystal form A.

5. The pharmaceutical composition according to any one of claims 1-3, wherein, the pharmaceutically acceptable carrier includes a filler, a disintegrant, an adhesive, a glidant and a lubricant.

6. The pharmaceutical composition according to claim 5, wherein, the filler is selected from the group consisting of starch, mannitol, sorbitol, microcrystalline cellulose, lactose, pregelatinized starch and inorganic salts or a combination thereof.

7. The pharmaceutical composition according to claim 6, wherein, the filler includes a combination of microcrystalline cellulose and pregelatinized starch.

8. The pharmaceutical composition according to claim 5, wherein, the disintegrant is selected from the group consisting of dry starch, sodium carboxymethyl starch, hydroxypropyl starch, low substituted hydroxypropyl cellulose, crospovidone and croscarmellose sodium or a combination thereof.

9. The pharmaceutical composition according to claim 8, wherein, the disintegrant includes sodium carboxymethyl starch.

10. The pharmaceutical composition according to claim 5, wherein, the adhesive is selected from the group consisting of distilled water, ethanol, starch paste, powdered sugar and syrup, hydroxypropyl methylcellulose, povidone, hydroxypropyl cellulose, methyl cellulose, ethyl cellulose and sodium carboxymethyl cellulose or a combination thereof.

11. The pharmaceutical composition according to claim 10, wherein, the adhesive includes hydroxypropyl methylcellulose.

12. The pharmaceutical composition according to claim 5, wherein, the glidant is selected from the group consisting of micronized silica gel and talc or a combination thereof.

13. The pharmaceutical composition according to claim 12, wherein, the glidant includes micronized silica gel.

14. The pharmaceutical composition according to claim 5, wherein, the lubricant is selected from the group consisting of magnesium stearate, hydrogenated vegetable oil, polyethylene glycol, dodecyl magnesium sulfate and sodium stearyl fumarate or a combination thereof.

15. The pharmaceutical composition according to claim 14, wherein, the lubricant includes magnesium stearate.

16. The pharmaceutical composition according to any one of claims 1-3, wherein, the pharmaceutical composition is a tablet.

17. The pharmaceutical composition according to claim 16, wherein, the pharmaceutical composition is a film-coated tablet.

18. The pharmaceutical composition according to any one of claims 1-3, wherein, the components each in percentage by weight are: 1% to 50% of spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof, 1% to 90% of the filler, 1% to 10% of the disintegrant, 0.1% to 1% of the glidant and 0.5% to 5% of the lubricant.

19. The pharmaceutical composition according to claim 18, wherein, the components each in percentage by weight are: 5.0% of spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof, 26.0% of microcrystalline cellulose, 59.5% of pregelatinized starch, 5% of sodium carboxymethyl starch, 3.0% of hydroxypropyl methylcellulose, 0.5% of micronized silica gel and 1% of magnesium stearate.

20. A method for preparing the pharmaceutical composition according to claim 1, comprising the following steps:

1) mixing spirocyclic aryl phosphorus oxide or a pharmaceutically acceptable salt thereof with internal excipients including a filler, an adhesive and a disintegrant, and then granulating;

2) adding other excipients including a filler, a disintegrant and a lubricant, and then mixing; and

3) tableting and coating.

21. Use of the pharmaceutical composition according to any one of claims 1-19 or the pharmaceutical composition prepared according to the method of claim 20 in treating cancer.

22. The use according to claim 21, wherein the cancer includes non-small cell lung cancer, lymphoma, non-Hodgkin lymphoma, ovarian cancer, cervical cancer, prostate cancer, colorectal cancer, breast cancer, pancreatic cancer, glioma, glioblastoma, melanoma, leukemia, gastric cancer, endometrial carcinoma, lung cancer, hepatocellular carcinoma, gastric carcinoma, gastrointestinal stromal tumor, acute myeloid leukemia, cholangiocarcinoma, renal carcinoma, thyroid cancer, anaplastic large cell lymphoma, mesothelioma, multiple myeloma, and melanoma.

23. The use according to claim 22, wherein the cancer is non-small cell lung cancer.

24. A method for treating cancer, comprising administering a therapeutically effective amount of the pharmaceutical composition according to any one of claims 1-19 to a patient, wherein, the cancer includes non-small cell lung cancer, lymphoma, non-Hodgkin lymphoma, ovarian cancer, cervical cancer, prostate cancer, colorectal cancer, breast cancer, pancreatic cancer, glioma, glioblastoma, melanoma, leukemia, gastric cancer, endometrial carcinoma, lung cancer, hepatocellular carcinoma, gastric carcinoma, gastrointestinal stromal tumor, acute myeloid leukemia, cholangiocarcinoma, renal carcinoma, thyroid cancer, anaplastic large cell lymphoma, mesothelioma, multiple myeloma and melanoma.

25. The method according to claim 24, wherein the cancer is non-small cell lung cancer.