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(54) **CHEMICAL REGENERATION METHOD OF OXIDIZED COENZYME NAD (P)+**

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(57) **ABSTRACT**

It discloses a chemical regeneration method of oxidized coenzyme NAD(P)⁺ which is under an oxygen or air atmosphere condition, adding a catalytic amount of bridged flavin, and oxidizing NAD(P)H to obtain NAD(P)⁺. The catalyst for regeneration of cofactor is cheap and easily available small organic molecule having no noble metal; this regeneration system can regenerate NADH and NADPH; this regeneration system has a wide pH range and temperature range, being applicable to various oxidation reactions catalyzed by nicotinamide-dependent oxidoreductase.

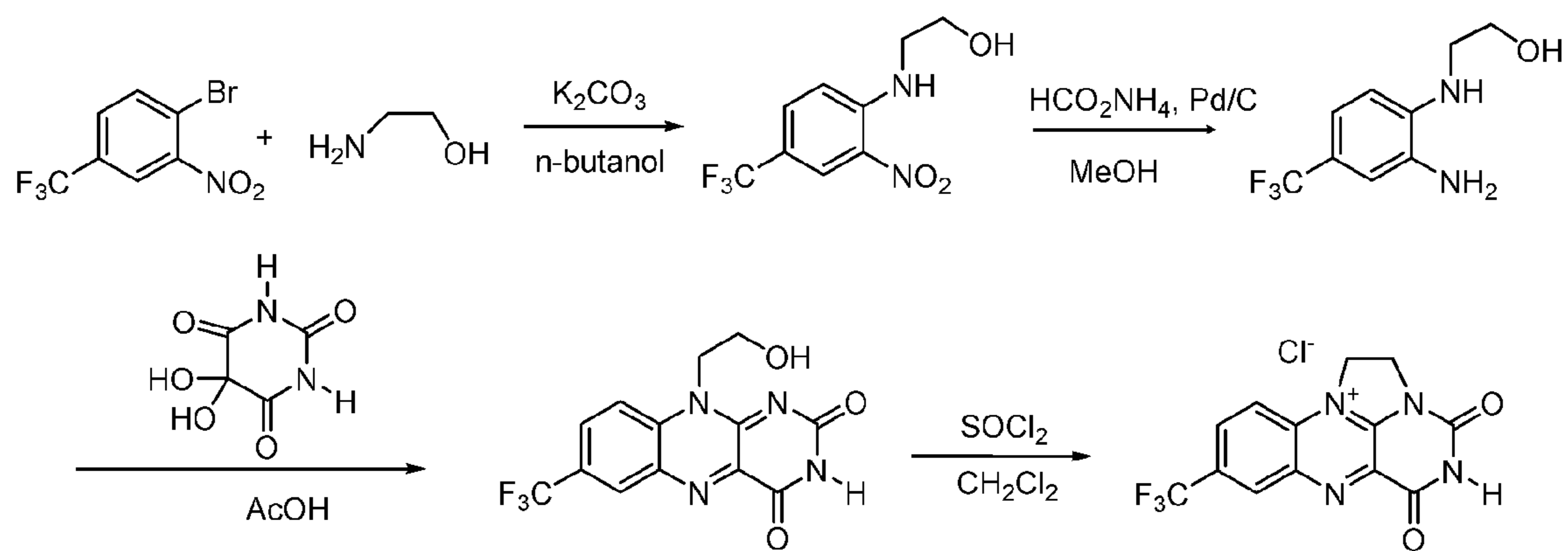


FIG. 1

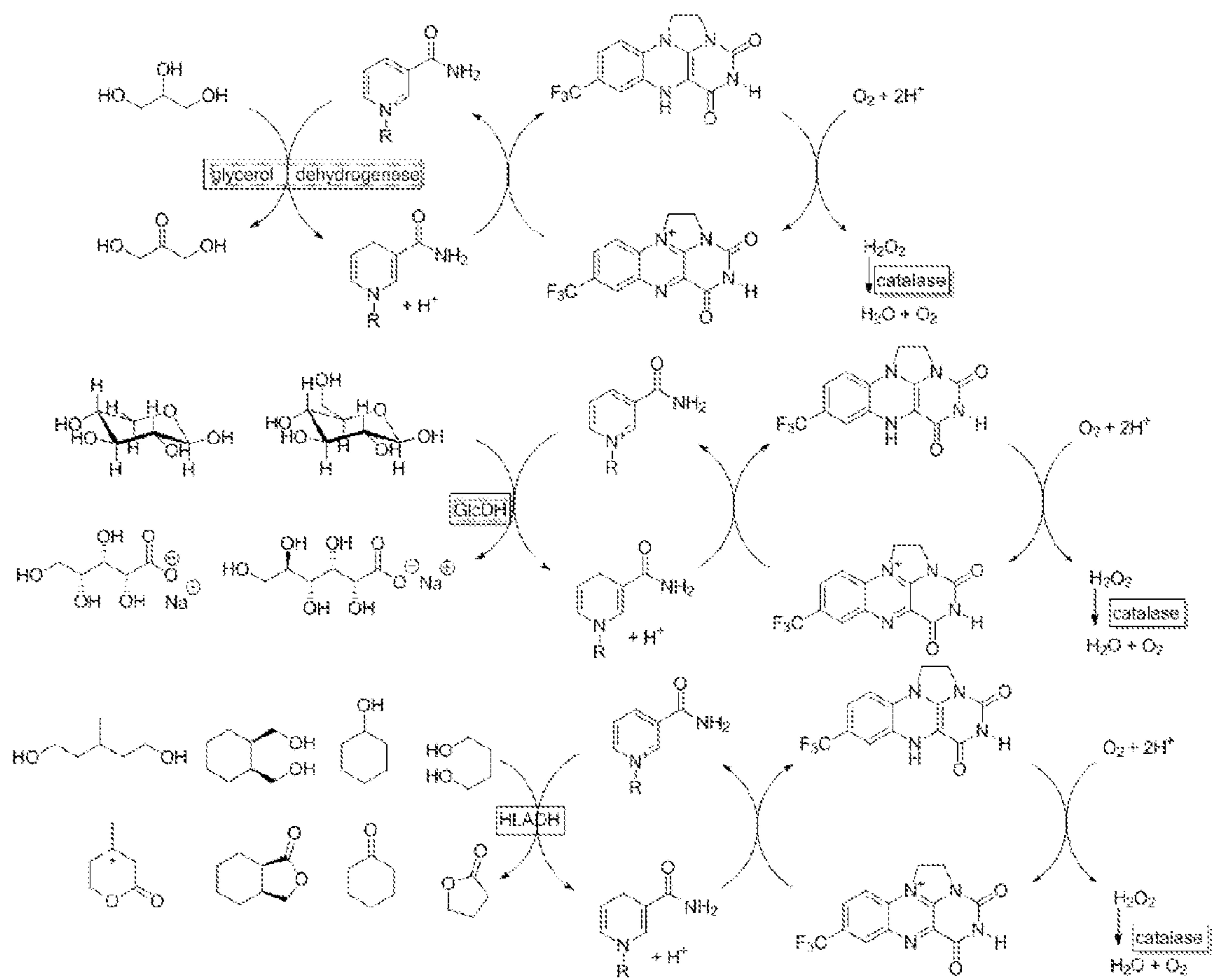


FIG. 2

CHEMICAL REGENERATION METHOD OF OXIDIZED COENZYME NAD(P)⁺

[0001] The application is related to Chinese application Ser. No. CN201610013723.4, filed on Jan. 8, 2016 and entitled "CHEMICAL REGENERATION METHOD OF OXIDIZED COENZYME NAD(P)⁺", which is hereby incorporated herein by reference.

TECHNICAL FIELD

[0002] The present invention relate to a chemical regeneration method of oxidized coenzyme NAD(P)⁺, it belongs to a technical field of coenzyme regeneration.

BACKGROUND OF THE INVENTION

[0003] Oxidoreductase is a kind of enzyme being most widely used after hydrolase, most of the oxidation-reduction reactions catalyzed by it needs NAD(P)H/NAD(P)⁺ to provide reduced hydrogen or to receive hydrogen, but high price of NAD(P)H/NAD(P)⁺ restricts use of the oxidoreductase in industrial production, requiring conducting in situ regeneration of coenzyme to reduce the cost. At present, the coenzyme regeneration includes five methods of whole-cell method, enzyme method, electrochemical method, photochemical method and chemical method. For the whole-cell method, its cost is low, but product separation is complicated, the method is comparatively primitive. The enzyme method includes two methods of substrate coupling method and enzyme coupling method, wherein the substrate coupling method is the same enzyme catalyze both of two different substrates towards reverse directions, meanwhile achieving conversion of main substrate and co-substrate, forming a coenzyme regeneration circulation, co-substrate and main substrate compete an active center of the enzyme, a part of the enzyme activity is used in regeneration of coenzyme, thus the enzyme activity used in catalyzing the main substrate is reduced; in the enzyme coupling method, the oxidization reaction enzyme system and the reduction reaction enzyme system works in one system in parallel, production of the enzyme catalyzes substrate conversion and consumes coenzyme, regeneration of the enzyme catalyzes regeneration of the coenzyme, this method requires that the substrates in two enzyme systems must be independent to each other, to avoid mutual competition to active center of the same enzyme, and optimum reaction condition for these two enzymes are often inconsistent. For the electrochemical method, the cost of electricity is low, there is no stoichiometric regeneration reagent, but the efficiency is low, often requiring participation of a medium, and this method is incompatible to many biochemical catalytic systems, its selectivity is poor. The chemical method uses a special catalyst to transfer oxid-reduction equivalent of the oxidant and reductant to a redox cofactor NAD(P)H/NAD(P)⁺ thereby forming a cofactor regeneration circulation, the commonly used chemical catalyst is a noble metal organic complex (Rh/Ru/Ir), its price is high and center metal may fall off, the center metal often coordinates with amino acid residue of the enzyme, making the metal catalyst and the enzyme both losing activity. The photochemical regeneration method is often limited by deficiency of efficient photosensitizer, thus generally its efficiency is low.

[0004] The chemical method currently used in regeneration of the oxidized cofactor NAD(P)⁺ often uses two kinds of regeneration catalyst for riboflavin mononucleotide

(FMN) [*J. Am. Chem. Soc.* 1982, 104, 4659-4665] and iron (III) porphyrin [*Angew. Chem. Int. Ed.* 2011, 50, 2397-2400]. When the FMN is used in the regeneration, it is required to use overdose [*J. Am. Chem. Soc.* 1982, 104, 4659-4665] or under radiation of a special light source [*Chem Cat Chem* 2011, 3, 338-342] to conduct catalytic regeneration. The iron porphyrin belongs to metal organic complex, its regeneration efficiency is not high, and tends to coordinate with the enzyme, resulting in deactivation.

SUMMARY OF THE INVENTION

[0005] In consideration of above-described problems, the technical problem to be resolved by the present invention is to provide a new chemical method which uses bridged flavin regeneration NAD(P)⁺, to overcome the shortcomings of other regeneration methods.

[0006] To resolve the above-described technical problem, the technical solution adopted by the present invention is as follows:

[0007] The first technical solution to be protected by the present invention is a chemical regeneration method of oxidized coenzyme NAD(P)⁺.

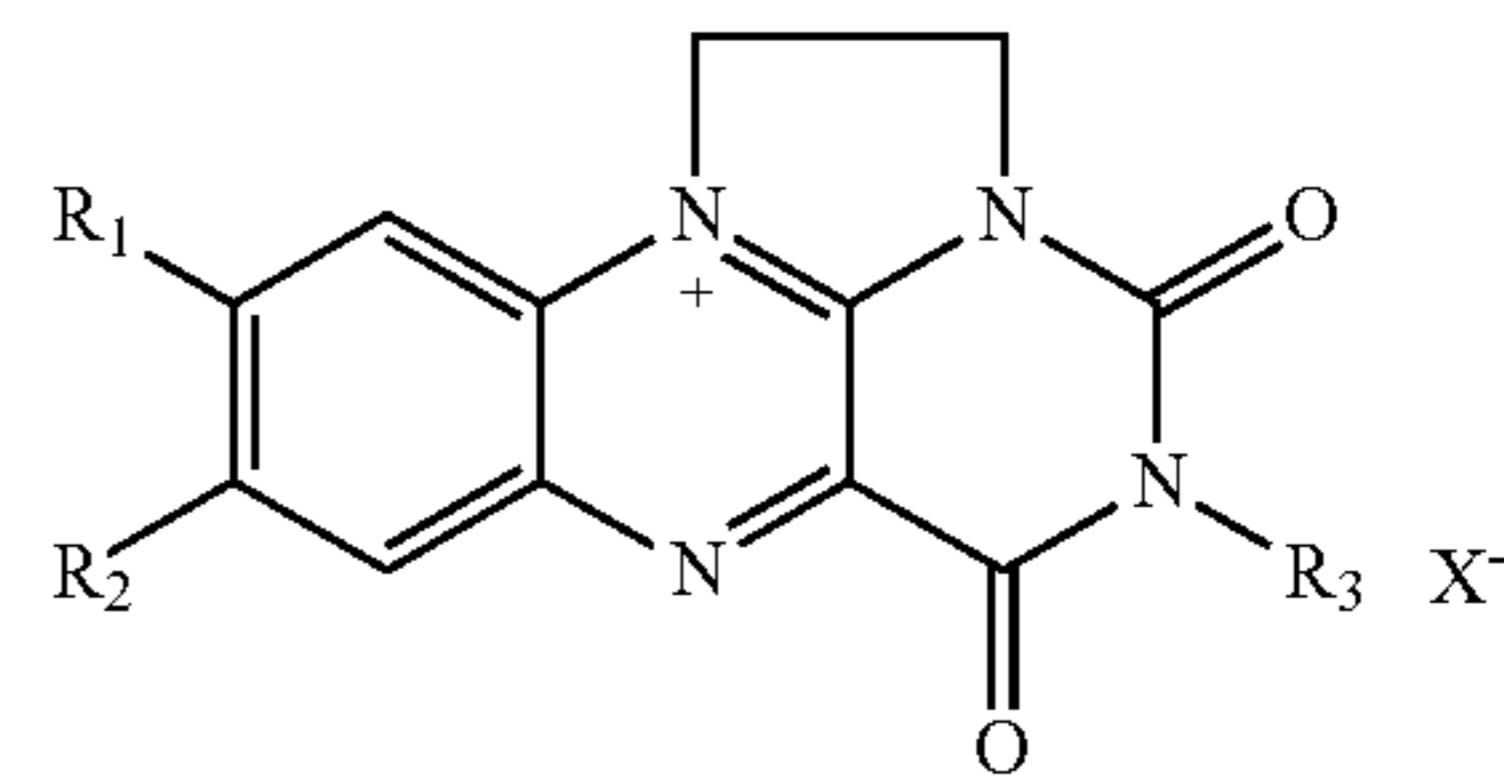
[0008] A chemical regeneration method of oxidized coenzyme NAD(P)⁺, the method is under oxygen or air atmosphere condition, adding a catalytic amount of bridged flavin, oxidizing NAD(P)H to obtain NAD(P)⁺.

[0009] The NAD(P)⁺ of the present invention is NAD⁺ or NADP⁺.

[0010] The NAD(P)H of the present invention is NADH or NADPH.

[0011] Wherein, NADH is oxidized to obtain NAD⁺, and NADPH is oxidized to obtain NADP⁺.

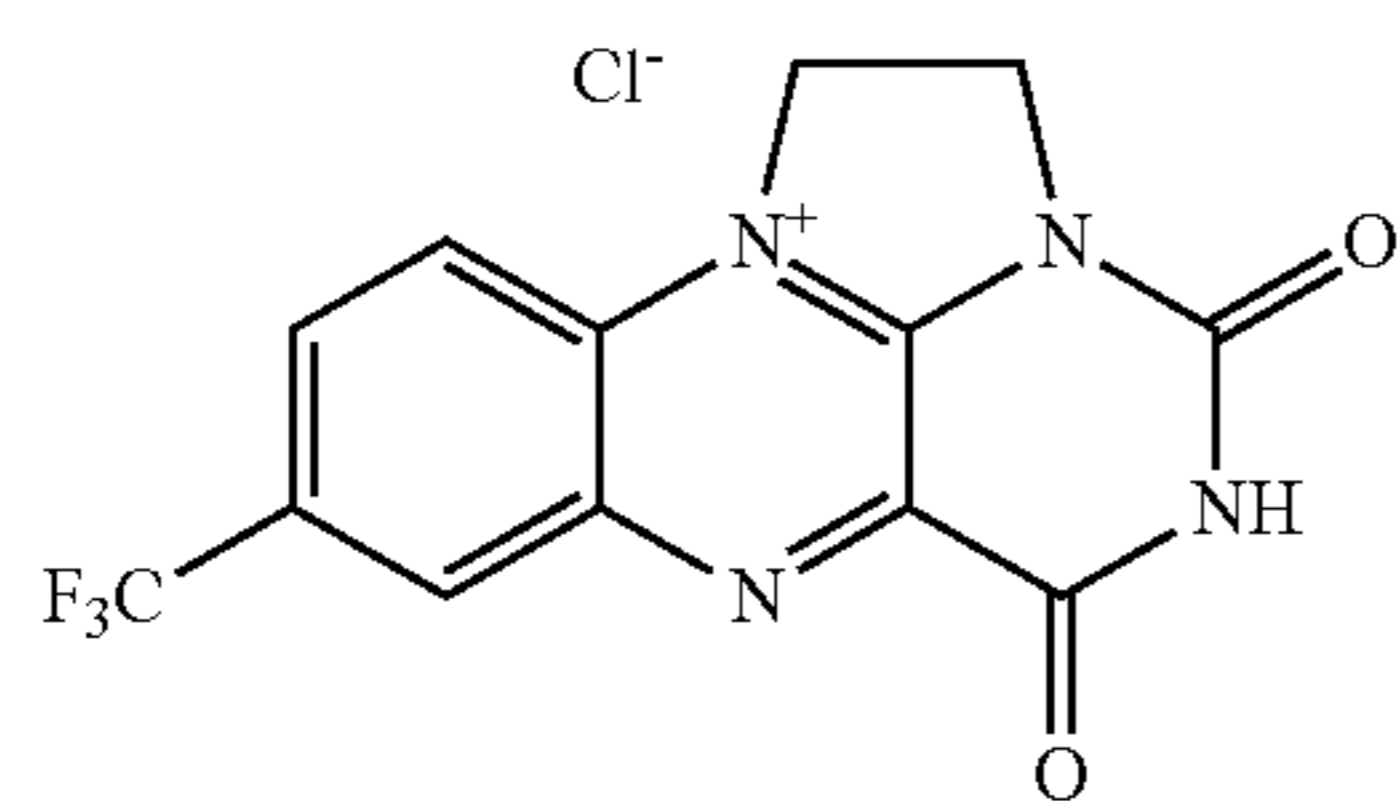
[0012] For the bridged flavin (English name N¹,N¹⁰-ethylene-bridged flavins) of the present invention, its general structure formula is as follow:



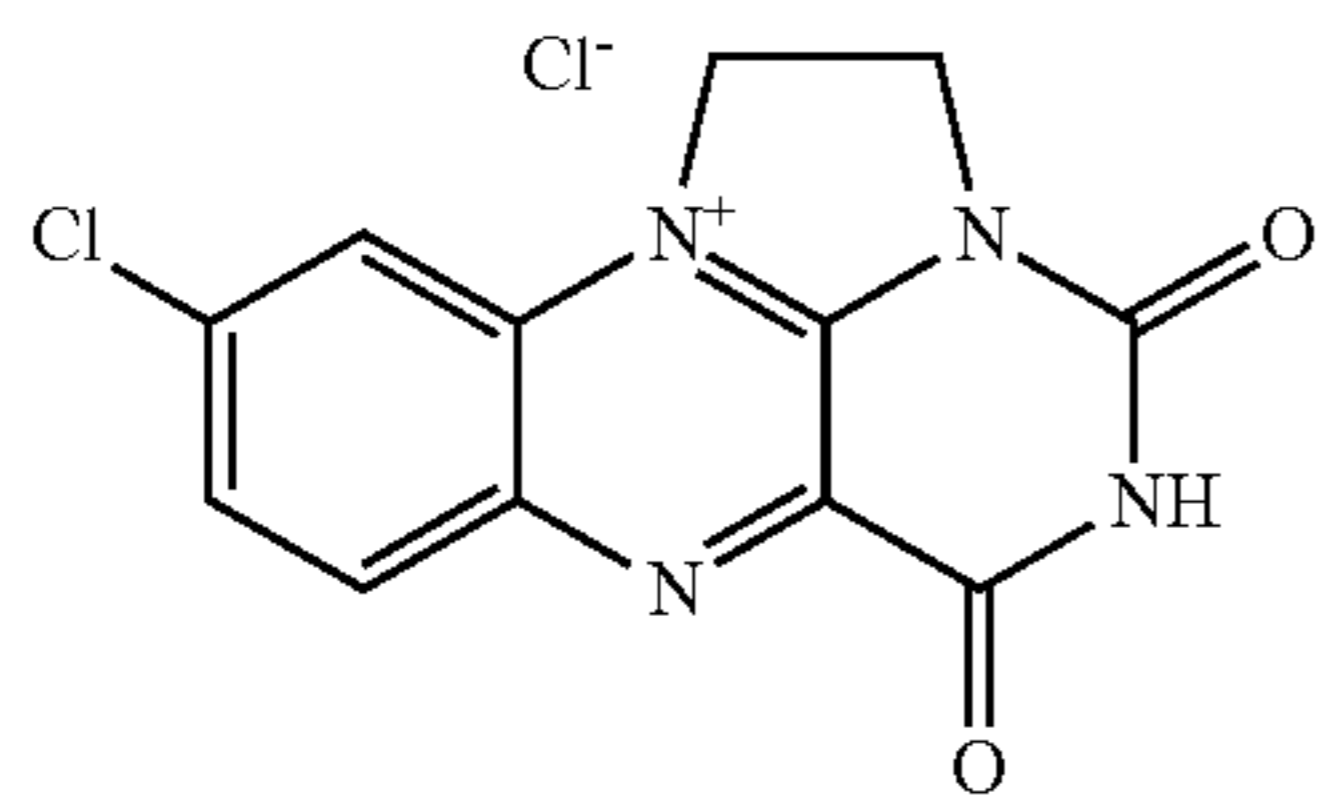
wherein, R₁ and R₂ are independently selected from hydrogen, methyl, trifluoromethyl, methoxyl, halogen atom, nitro, amino; R₃ is selected from hydrogen, C1-C5 alkyl, phenyl, benzyl; X⁻ is selected from halide ion, nitrate radical, trifluoromethanesulfonic acid radical.

[0013] It is preferred that, R₁ includes but not limited to hydrogen, methyl, halogen atom; R₂ includes but not limited to hydrogen, methyl, halogen atom, trifluoromethyl; R₃ includes but not limited to hydrogen, methyl; X⁻ is a halide ion.

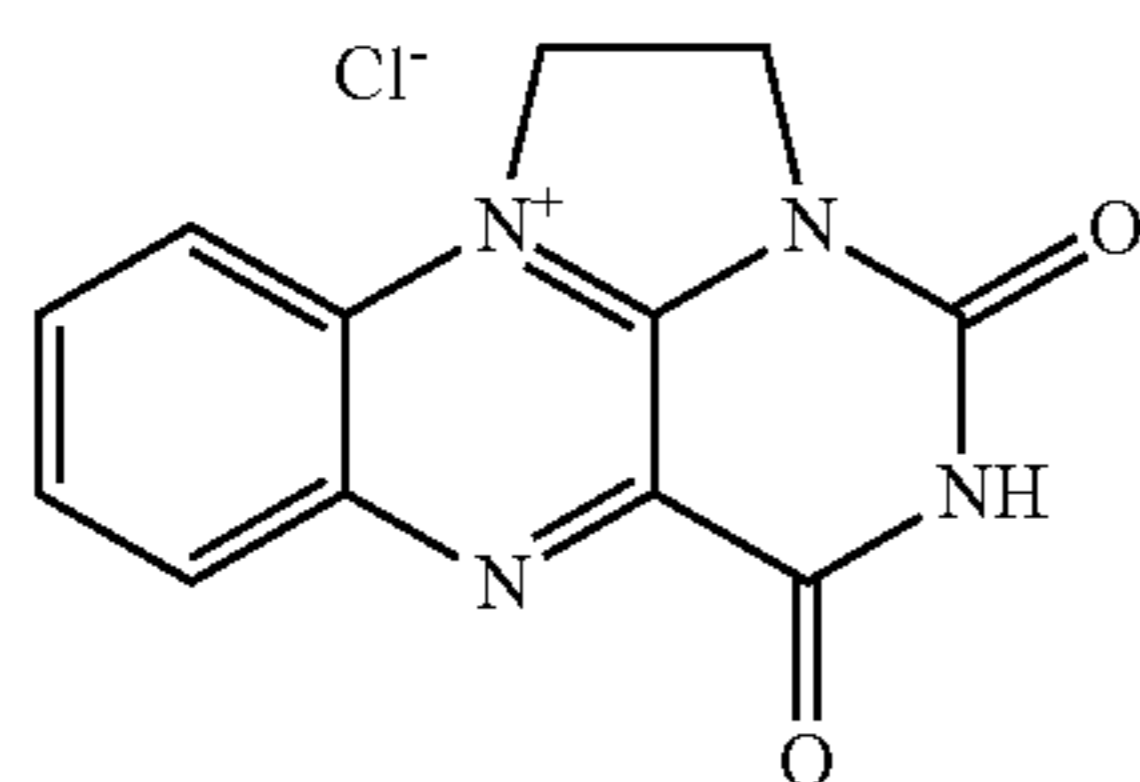
[0014] It is most preferred that, the bridged flavin includes but not limited to 7-trifluoromethyl-1,10-ethyleneisoalloxazine chloride (Compound I), 8-chloro-1,10-ethyleneisoalloxazine chloride (Compound II), or 1,10-ethyleneisoalloxazine chloride (Compound III).



(I)



(II)



(III)

[0015] The bridged flavin of the present invention can be synthesized by referring to existing disclosed literatures, e.g., Literature [*Tetrahedron*, 2001, 57, 4507-4522]; or directly purchased from the market.

[0016] Wherein, the mole percentage of the catalytic amount of the bridged flavin is 0.1-5%, preferably 0.25-2%, most preferably 2% of NAD(P)H.

[0017] Wherein, the condition for oxidizing NAD(P)H to obtain NAD(P)⁺ is pH 4-10, temperature 30-70° C.; preferably the condition for oxidizing NAD(P)H to obtain NAD(P)⁺ is pH 6-8 and temperature 30-40° C.; most preferably the condition for oxidizing NAD(P)H to obtain NAD(P)⁺ is pH 7 and temperature 30° C.

[0018] Wherein, the reaction time for oxidizing NAD(P)H to obtain NAD(P)⁺ is generally based on the reaction being complete or most of the reaction being complete, a person skilled in the art can adopt a ultraviolet spectrophotometry to detect the completion of reaction, because NAD(P)H has a ultraviolet absorption at 340 nm, but NAD(P)⁺ has no absorption at 340 nm, thus the ultraviolet spectrophotometry can be used to detect reaction process, when the sample absorbance after subtracting by blank absorbance of the flavin is reduced to 0, it is regarded that the reaction is complete, when the absorbance approaches 0, it is regarded that most of the reaction is complete, of course a person skilled in the art can also use other existing detecting methods to monitor the reaction progress such that the reaction is complete.

[0019] Wherein, the bridged flavin, as a NAD(P)⁺ regeneration catalyst, can be coupled with the oxidation reaction catalyzed by NAD(P)⁺-dependent oxidoreductase, forming a regeneration circulation system of coenzyme NAD(P)⁺.

[0020] Wherein, the NAD(P)⁺-dependent oxidoreductase includes but not limited to all enzymes of EC1.1.1.X, EC1.2.1.X, EC1.3.1.X, EC1.4.1.X, EC1.5.1.X, EC1.6.1.X, EC1.7.1.X, EC1.8.1.X, EC1.10.1.X, EC1.12.1.X, EC1.13.1.X, EC1.16.1.X, EC1.17.1.X, EC1.18.1.X, EC1.20.1.X, EC1.22.1.X. It is preferred that, the NAD(P)⁺-dependent oxi-

doreductase includes but not limited to horse liver alcohol dehydrogenase, glucose dehydrogenase or glycerol dehydrogenase.

[0021] When the bridged flavin, as NAD(P)⁺ regeneration catalyst, is coupled with the oxidation reaction catalyzed by NAD(P)⁺-dependent oxidoreductase, the amount of the bridged flavin is 0.1-5%, preferably 0.25-2%, most preferably 2% of mole number of the substrate catalyzed by NAD(P)⁺-dependent oxidoreductase.

[0022] The second technical solution to be protected by the present invention is an oxidation reaction.

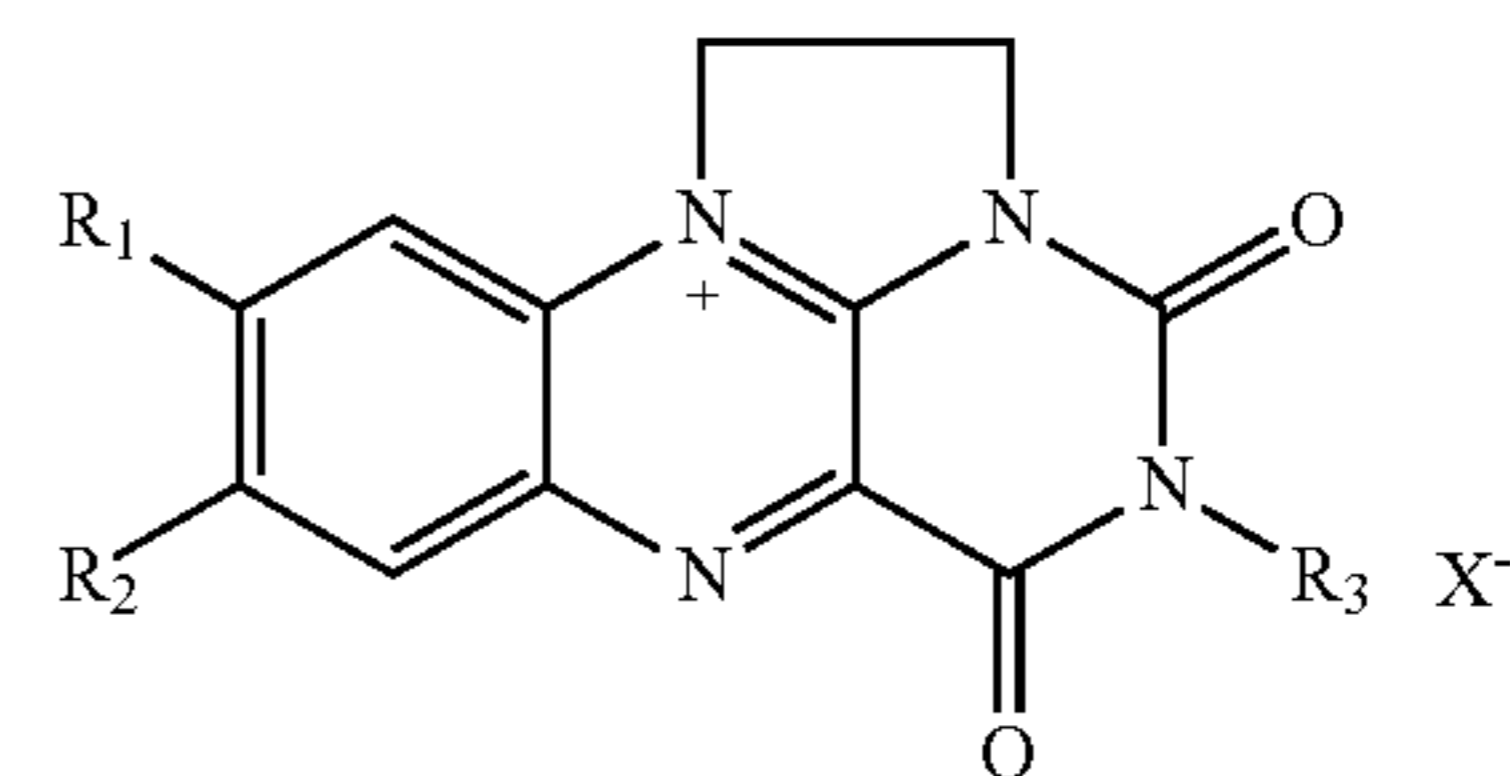
[0023] A oxidation reaction, it uses NAD(P)⁺-dependent oxidoreductase as a catalyst, and the bridged flavin as the NAD(P)⁺ regeneration catalyst, under an oxygen or air atmosphere condition, oxidizing NAD(P)H to obtain NAD(P)⁺, forming a regeneration circulation system of the coenzyme NAD(P)⁺.

[0024] The NAD(P)⁺ of the present invention is NAD⁺ or NADP⁺.

[0025] The NAD(P)H of the present invention is NADH or NADPH.

[0026] Wherein, NADH is oxidized to obtain NAD⁺, and NADPH is oxidized to obtain NADP⁺.

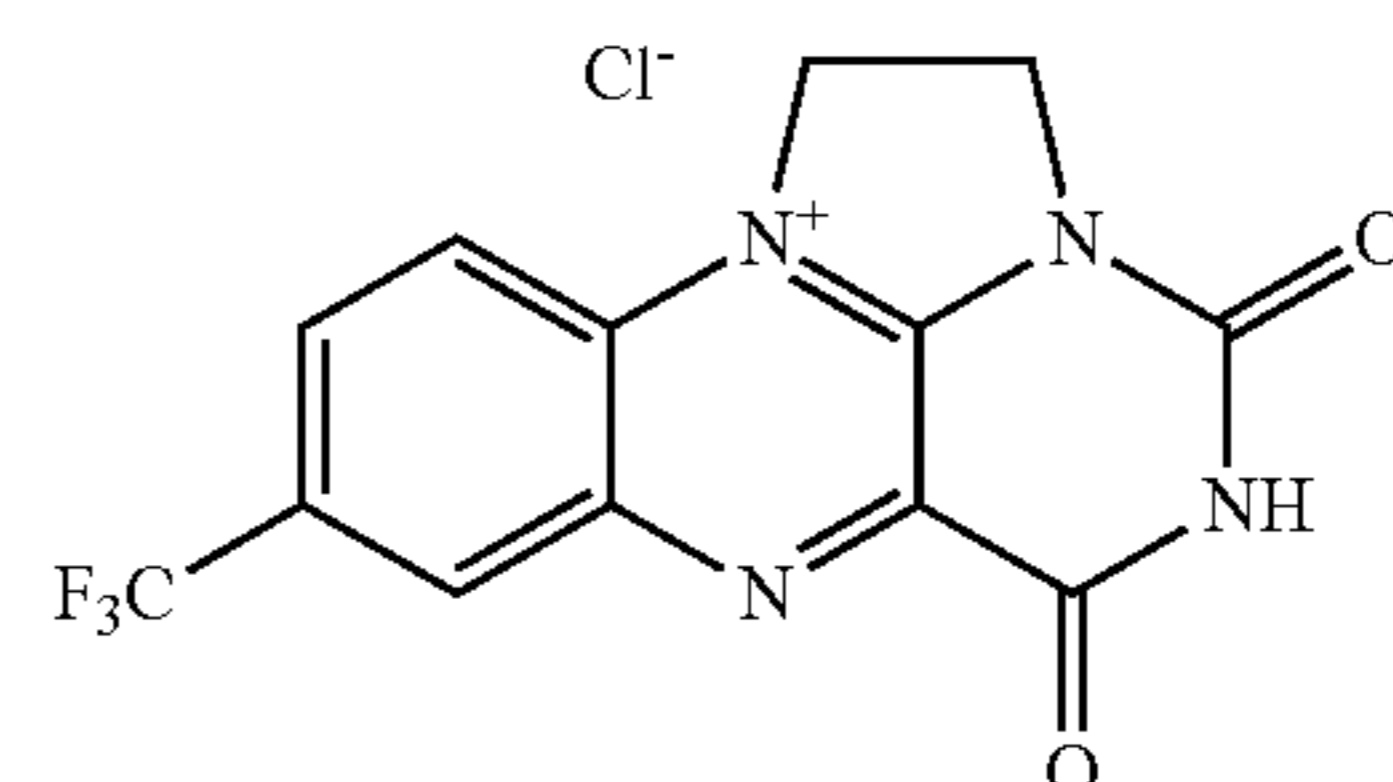
[0027] For the bridged flavin of the present invention (English name N¹,N¹⁰-Ethylene-bridged flavins), its general structure formula is as follow:



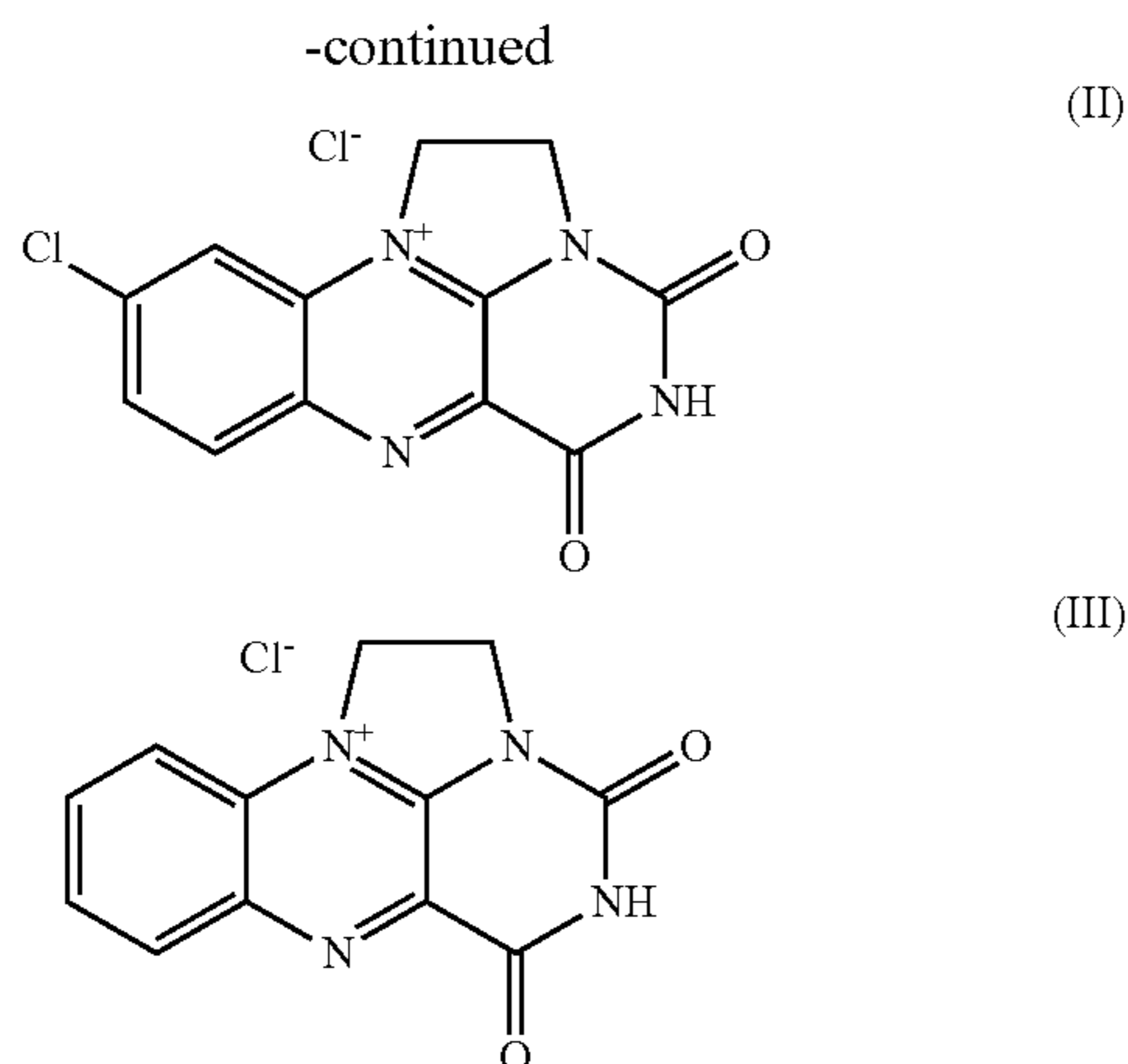
[0028] wherein, R₁ and R₂ are independently selected from hydrogen, methyl, trifluoromethyl, methoxyl, halogen atom, nitro, amino; R₃ is selected from hydrogen, C1-C5 alkyl, phenyl, benzyl; X⁻ is selected from halide ion, nitrate radical, trifluoromethanesulfonic acid radical.

[0029] It is preferred that, R₁ includes but not limited to hydrogen, methyl, halogen atom; R₂ includes but not limited to hydrogen, methyl, halogen atom, trifluoromethyl; R₃ includes but not limited to hydrogen, methyl; X⁻ is halide ion.

[0030] It is most preferred that, the bridged flavin includes but not limited to 7-trifluoromethyl-1,10-ethyleneisoalloxazine chloride (Compound I), 8-chloro-1,10-ethyleneisoalloxazine chloride (Compound II), or 1,10-ethyleneisoalloxazine chloride (Compound III).



(I)



[0031] The bridged flavin of the present invention can be synthesized by referring to existing disclosed literatures, e.g., Literature [*Tetrahedron*, 2001, 57, 4507-4522]; or directly purchased from the market.

[0032] Wherein, mole number catalyzed by the bridged flavin is 0.1-5%, preferably 0.25-2%, most preferably 2% of mole number of the substrate catalyzed by NAD(P)⁺-dependent oxidoreductase.

[0033] Wherein, the condition for oxidizing NAD(P)H to obtain NAD(P)⁺ is pH 3-12 and temperature 30-70° C.; preferably the condition for oxidizing NAD(P)H to obtain NAD(P)⁺ is pH 6-8 and temperature 30-40° C.; most preferably the condition for oxidizing NAD(P)H to obtain NAD(P)⁺ is pH 7 and temperature 30° C.

[0034] Wherein, the NAD(P)⁺-dependent oxidoreductase includes but not limited to all enzymes of EC1.1.1.X, EC1.2.1.X, EC1.3.1.X, EC1.4.1.X, EC1.5.1.X, EC1.6.1.X, EC1.7.1.X, EC1.8.1.X, EC1.10.1.X, EC1.12.1.X, EC1.13.1.X, EC1.16.1.X, EC1.17.1.X, EC1.18.1.X, EC1.20.1.X, EC1.22.1.X. It is preferred that, the NAD(P)⁺-dependent oxidoreductase includes but not limited to horse liver alcohol dehydrogenase, glucose dehydrogenase or glycerol dehydrogenase.

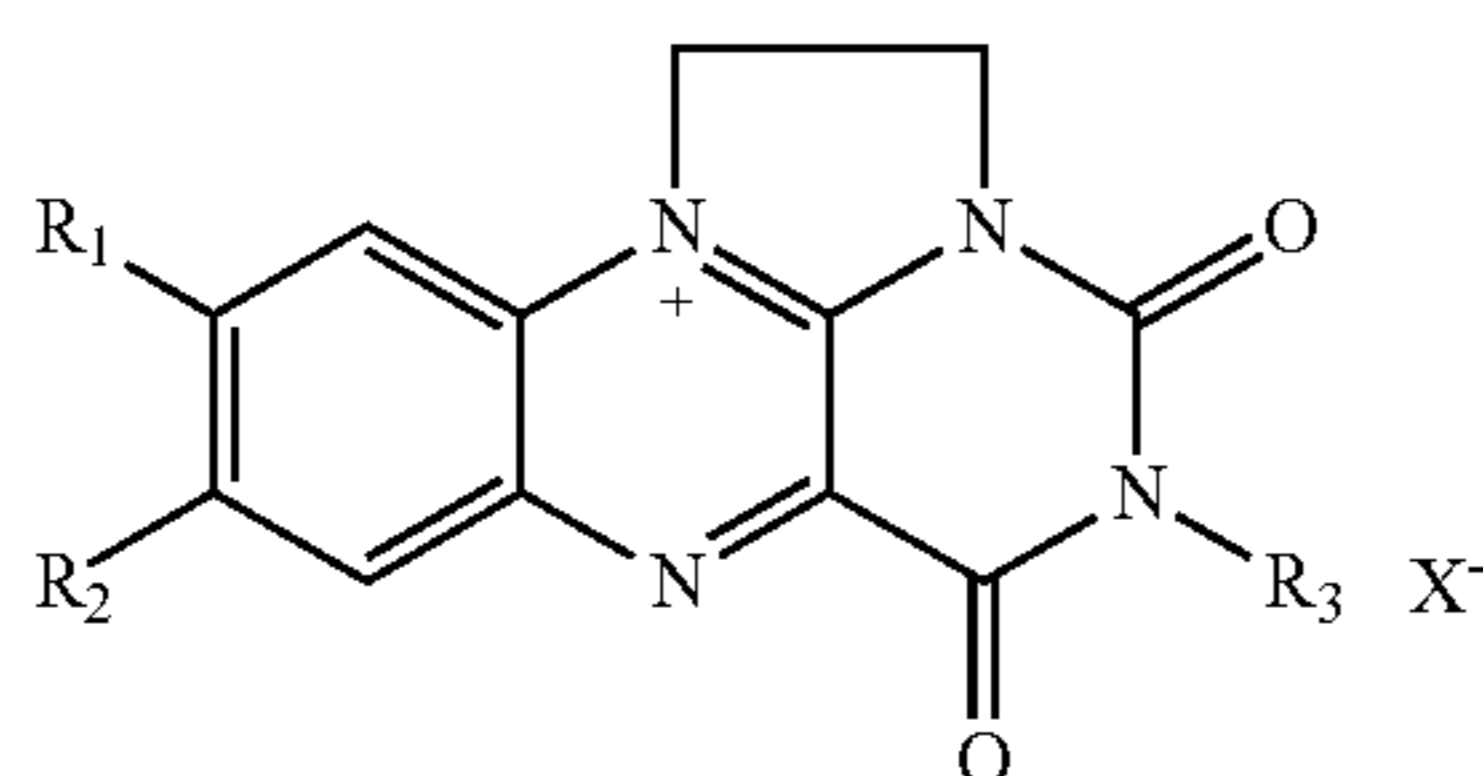
[0035] The third technical solution to be protected by the present invention is use of bridged flavin as a catalyst in oxidizing NAD(P)H to obtain NAD(P)⁺.

[0036] The NAD(P)⁺ of the present invention is NAD⁺ or NADP⁺.

[0037] The NAD(P)H of the present invention is NADH or NADPH.

[0038] Wherein, NADH is oxidized to obtain NAD⁺, and NADPH is oxidized to obtain NADP⁺.

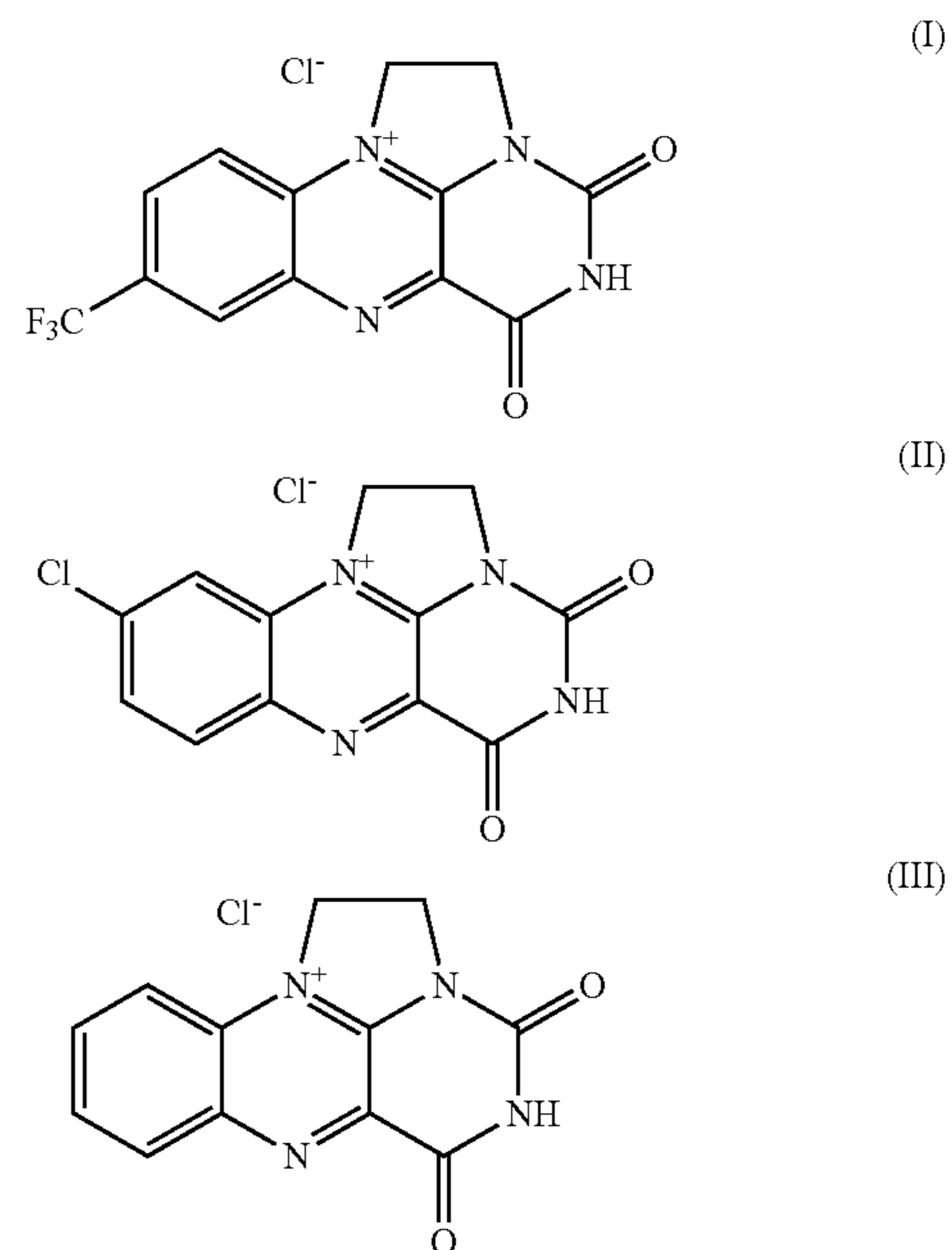
[0039] For the bridged flavin of the present invention (English name N¹,N¹⁰-Ethylene-bridged flavins), its general structure formula is as follow:



wherein, R₁ and R₂ are independently selected from hydrogen, methyl, trifluoromethyl, methoxyl, halogen atom, nitro, amino; R₃ is selected from hydrogen, C1-C5 alkyl, phenyl, benzyl; X⁻ is selected from halide ion, nitrate radical, trifluoromethanesulfonic acid radical.

[0040] It is preferred that, R₁ includes but not limited to hydrogen, methyl, halogen atom; R₂ includes but not limited to hydrogen, methyl, halogen atom, trifluoromethyl; R₃ includes but not limited to hydrogen, methyl; X⁻ is halide ion.

[0041] It is most preferred that, the bridged flavin includes but not limited to 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride (Compound I), 8-chloro-1,10-ethyleneisalloxazine chloride (Compound II), or 1,10-ethyleneisalloxazine chloride (Compound III).



[0042] The bridged flavin of the present invention can be synthesized by referring to existing disclosed literatures, e.g. Literature [*Tetrahedron*, 2001, 57, 4507-4522]; or directly purchased from the market.

[0043] Wherein, the mole number catalyzed by the bridged flavin is 0.1-5%, preferably 0.25-2%, most preferably 2% of mole number of NAD(P)H.

[0044] Wherein, the condition for oxidizing NAD(P)H to obtain NAD(P)⁺ is pH 4-10 and temperature 30-70° C.; preferably the condition for oxidizing NAD(P)H to obtain NAD(P)⁺ is pH 6-8 and temperature 30-40° C.; most preferably the condition for oxidizing NAD(P)H to obtain NAD(P)⁺ is pH 7 and temperature 30° C.

[0045] Wherein, the reaction time for oxidizing NAD(P)H to obtain NAD(P)⁺ is generally based on the react being complete or most the reaction being complete, a person skilled in the art can use ultraviolet spectrophotometry to detect completion of the reaction, because NAD(P)H has an ultraviolet absorption at 340 nm, but NAD(P)⁺ has no absorption at 340 nm, thus ultraviolet spectrophotometry can be used to detect the reaction process, when the absorbance of the sample after subtracting by blank absorbance of

the bridged flavin is reduced to 0, it is regarded that the reaction is complete, when absorbance is close to 0, it is regarded that most of the reaction is complete, of course, a person skilled in the art can also adopt other existing detecting method to monitor the reaction process such that the reaction is complete.

[0046] Beneficial effect: Compared to the prior art, the main advantage of the present invention is: the efficiency of regenerating NAD(P)⁺ by the bridged flavin is very high, the TOF can be up to 19 min⁻¹, and it can completely converting, application extent is wide, the condition is pH 4-10 and temperature 30-70° C. It can be coupled with various oxidation reactions catalyzed by the oxidoreductase under an optimum condition, forming a cofactor regeneration system.

BRIEF DESCRIPTION OF THE DRAWINGS

[0047] FIG. 1 is a synthesis route diagram of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride;

[0048] FIG. 2 is a route diagram of a coupling reaction of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride as a NAD⁺ regeneration catalyst with horse liver alcohol dehydrogenase and glucose dehydrogenase.

DESCRIPTION OF THE EMBODIMENTS

[0049] Based on the following examples, the present invention can be better understood. However, a person skilled in the art will readily understand that, the contents described in the examples are only used to illustrate the present invention, and should not and will not restrict the present invention described in detail in the claims.

Example 1: Synthesis of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride

[0050] 4-bromo-3-nitrotrifluorotoluene (1.76 g, 6.53 mmol) and ethanolamine (1.19 g, 19.59 mmol) were dissolved in 20 mL ethanol, potassium carbonate (1.08 g, 7.83 mmol) was added and stirred under reflux for 5-6 hours, after the reaction solution was cooled, filtered, the filtrate was concentrated under reduced pressure, to obtain an orange solid, and 20 mL saturated brine was added, extracted with ethyl acetate (3×40 mL), the organic phases were combined, washed with a saturated brine, dried on anhydrous sodium sulfate, rotary evaporated, purified by silica gel column chromatography to obtain 4-trifluoromethyl-2-nitro-N-(2-hydroxyethyl)aniline (1.58 g, yield 96%).

[0051] To an anhydrous methanol solution of 4-trifluoromethyl-2-nitro-N-(2-hydroxyethyl)aniline (1.20 g, 4.80 mmol) (5° C.), ammonium formate (1.51 g, 24.0 mmol) and 5% Pd/C (1.60 g) were added, stirred and reacted at 0 to 25° C. for 1 hour, filtered, the filter residue was washed with methanol, the filtrate was concentrated under reduced pressure, producing a red solid, 30 mL saturated brine was added, extracted with ethyl acetate (3×30 mL), the organic phases were combined, dried on anhydrous sodium sulfate, filtered, concentrated under reduced pressure, to obtain a white solid (1.04 g, 94%).

[0052] To a 50° C. acetic acid solution of 4-trifluoromethyl-N-(2-hydroxyethyl)-o-phenylenediamine (1 g, 4.35 mmol) filled with argon, alloxan monohydrate (0.73 g, 4.35 mmol) and boric acid (0.29 g, 4.67 mmol) were added,

reacted for 1 hour, cooled, the obtained yellow solid was filtered, washed with dichloromethane, vacuum dried, the obtained yellow solid was immediately placed into a round bottom flask filled with nitrogen gas, and thionyl chloride (20 mL) was slowly added meanwhile vigorously stirred, reacted for 16 hours under 50° C. nitrogen gas shielding, then cooled and filtered, the obtained solid was washed with dichloromethane, the crude product was dissolved in a minimal amount of 98% formic acid solution, and recrystallized with diethyl ether, to obtain a yellow solid (1.28 g, 95%). ESI-MS⁺: 309; ¹H NMR (CD₃COOD-CF₃COOH, 1:6) δ 4.96 (t, J=8.9 Hz, 2H), 5.58 (t, J=8.9 Hz, 2H), 8.20 (d, J=8.8 Hz, 1H), 8.47 (d, J=8.8 Hz, 1H), 8.81 (s, 1H), ¹³C NMR (CD₃COOD-CF₃COOH, 1:6) δ 46.2, 52.1, 122.6 (q, J=272.6 Hz, CF₃), 118.9, 131.4, 131.7, 133.9, 135.4 (q, J=35.5 Hz, C CF₃), 136.8, 141.1, 145.2, 146.4, 158.8.

Example 2: Synthesis of 1,10-ethyleneisalloxazine chloride

[0053] 2-fluoronitrobenzene (0.92 g, 6.53 mmol) and ethanolamine (1.19 g, 19.59 mmol) were dissolved in 20 mL ethanol, potassium carbonate (1.08 g, 7.83 mmol) was added and stirred under reflux for 5-6 hours, after the reaction solution was cooled, filtered, the filtrate was concentrated under reduced pressure, to obtain an orange solid, and 20 mL saturated brine was added, extracted with ethyl acetate (3×40 mL), the organic phases were combined, washed with a saturated brine, dried on anhydrous sodium sulfate, rotary evaporated, purified by silica gel column chromatography to obtain 2-nitro-N-(2-hydroxyethyl)aniline (0.98 g, yield 82.9%).

[0054] To a anhydrous methanol solution (5° C.) of 2-nitro-N-(2-hydroxyethyl)aniline (0.87 g, 4.80 mmol), ammonium formate (1.51 g, 24.0 mmol) and 5% Pd/C (1.60 g) were added, stirred and reacted at 0-25° C. for 1 hour, filtered, the filter residue was washed with methanol, the filtrate was concentrated under reduced pressure, producing a red solid, and 30 mL saturated brine was added, extracted with ethyl acetate (3×30 mL), the organic phases were combined, dried on anhydrous sodium sulfate, filtered, concentrated under reduced pressure, to obtain a white solid (0.67 g, 92%).

[0055] In a 50° C. acetic acid solution of N-(2-hydroxyethyl)-o-phenylenediamine (0.66 g, 4.35 mmol) filled with argon, alloxan monohydrate (0.73 g, 4.35 mmol) and boric acid (0.29 g, 4.67 mmol) were added, reacted for 1 hour, cooled, the resulting yellow solid was filtered, washed with dichloromethane, vacuum dried, the yellow solid obtained was immediately placed into a round bottom flask filled with nitrogen gas, and thionyl chloride (20 mL) was slowly added meanwhile vigorously stirred, reacted for 16 hours under 50° C. nitrogen gas shielding, then cooled and filtered, the solid obtained was washed with dichloromethane, the crude product was dissolved in a minimal amount of 98% formic acid solution, and recrystallized with diethyl ether, to obtain a yellow solid (0.88 g, 73%). ESI-MS⁺: 240.8, ¹H NMR (CD₃COOD-CF₃COOH, 1:6) δ 4.94 (t, J=8.9 Hz, 2H), 5.53 (t, J=8.9 Hz, 2H), 8.04-8.57 (4H); ¹³C NMR (CD₃COOD-CF₃COOH, 1:6) δ 45.9, 51.9, 117.2, 130.3, 131.9, 133.2, 134.2, 142.1, 142.2, 144.1, 146.7, 159.4.

Example 3: Synthesis of 8-chloro-1,10-ethyleneisalloxazine chloride

[0056] 2,4-dichloronitrobenzene (1.25 g, 6.53 mmol) and ethanolamine (1.19 g, 19.59 mmol) were dissolved in 20 mL

ethanol, potassium carbonate (1.08 g, 7.83 mmol) was added and stirred under reflux for 5-6 hours, after the reaction solution was cooled, filtered, the filtrate was concentrated under reduced pressure, to obtain an orange solid, and 20 mL saturated brine was added, extracted with ethyl acetate (3×40 mL), the organic phases were combined, washed with saturated brine, dried on anhydrous sodium sulfate, rotary evaporated, purified by silica gel column chromatography to obtain 5-chloro-2-nitro-N-(2-hydroxyethyl)aniline (1.26 g, yield 89%).

[0057] To a 20 mL water solution (100° C.) containing 5-chloro-2-nitro-N-(2-hydroxyethyl)aniline (1.04 g, 4.80 mmol) and Sn (1.71 g, 14.40 mmol), 9 mL concentrated hydrochloric acid was slowly dripped in, then the reaction was placed at 5° C. and cooled for 30 minutes, neutralized with a 50% NaOH water solution, extracted with ethyl acetate (3×30 mL), the organic phases were combined, dried on anhydrous sodium sulfate, filtered, concentrated under reduced pressure, to obtain a white solid (0.47 g, 52%).

[0058] To a 50° C. acetic acid solution of 4-trifluoromethyl-N-(2-hydroxyethyl)-o-phenylenediamine (0.37 g, 2.0 mmol) filled with argon, alloxan monohydrate (0.32 g, 2.0 mmol) and boric acid (0.13 g, 2.14 mmol) were added, and reacted for 1 hour, cooled, the solid formed was filtered, washed with dichloromethane, vacuum dried, the solid obtained was immediately placed into a round bottom flask filled with nitrogen gas, and thionyl chloride (10 mL) was slowly added meanwhile stirred vigorously, reacted for 16 hours under 50° C. nitrogen gas shielding, then cooled and filtered, the solid obtained was washed with dichloromethane, the crude product was dissolved in a minimal amount of 98% formic acid solution, and recrystallized with diethyl ether, to obtain a solid (0.43 g, 69%). ESI-MS+: 274.9, ¹H NMR (CD₃COOD-CF₃COOH, 1:6) δ 4.92 (t, J=8.9 Hz, 2H), 5.49 (t, J=8.9 Hz, 2H), 8.06 (d, J=8.4 Hz, 2H), 8.50 (d, J=8.4 Hz, 1H); ¹³C NMR (CD₃COOD-CF₃COOH, 1:6) δ 45.9, 51.9, 117.2, 130.9, 131.8, 134.2, 135.1, 140.6, 144.6, 146.6, 150.1, 159.1.

Example 4:

7-trifluoromethyl-1,10-ethyleneisalloxazine chloride oxidizing NADH to NAD⁺

[0059] At 30° C., in a 10 mL 50 mM potassium phosphate buffer of pH 7, the initial concentration of NADH was 0.2 mM, the concentration of 7-trifluoromethyl 1,10-ethyleneisalloxazine chloride was 4 μM, the reaction solution was connected with the outside air. The change of the light absorption value of the reaction system at 340 nm was detected every 2 minutes, the conversion rate was calculated, and the results were shown in Table 1.

TABLE 1

Conversion rate of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride oxidizing NADH with time		
time (min)	light absorption value at 340 nm	conversion rate (%)
0	1.244	0.00
1	0.5991	51.84
2	0.2815	77.37
3	0.1425	88.55
4	0.0676	94.57
5	0.0313	97.48
6	0.0137	98.90

TABLE 1-continued

Conversion rate of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride oxidizing NADH with time		
time (min)	light absorption value at 340 nm	conversion rate (%)
7	0.0056	99.55
8	0.002	99.84
9	0	100

Example 5:

7-trifluoromethyl-1,10-ethyleneisalloxazine chloride oxidizing NADPH to NADP⁺

[0060] At 30° C., in a 10 mL 50 mM potassium phosphate buffer of pH 7, the initial concentration of NADPH was 0.2 mM, the concentration of 7-trifluoromethyl 1,10-ethyleneisalloxazine chloride was 4 μM, the reaction solution was connected with the outside air. Change of light absorption value of the reaction system at 340 nm was detected every 2 minutes, the conversion rate was calculated, and the results are shown in Table 2.

TABLE 2

Conversion rate of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride oxidizing NADPH with time		
time (min)	light absorption value at 340 nm	conversion rate (%)
0	1.244	0.00
1	0.5862	52.88
2	0.204	83.60
3	0.0983	92.10
4	0.0215	98.27
5	0.0098	99.21
6	0.0033	99.73
7	0	100

Example 6: 8-chloro-1,10-ethyleneisalloxazine chloride oxidizing NADH to NAD⁺

[0061] At 30° C., in a 10 mL 50 mM potassium phosphate buffer of pH 7, the initial concentration of NADH was 0.2 mM, the concentration of 8-chloro-1,10-ethyleneisalloxazine chloride was 4 μM, the reaction solution was connected with the outside air. Change of the light absorption value of the reaction system at 340 nm was detected every 2 minutes, the conversion rate of NADH was calculated, and the results are shown in Table 3.

TABLE 3

Conversion rate of 8-chloro-1,10-ethyleneisalloxazine chloride oxidizing NADH with time		
time (min)	light absorption value	conversion rate (%)
0	1.244	0.00
1	0.721	42.04
2	0.5612	54.89
3	0.4401	64.62
4	0.3471	72.10
5	0.2751	77.89
6	0.2192	82.38
7	0.1752	85.92

TABLE 3-continued

Conversion rate of 8-chloro-1,10-ethyleneisalloxazine chloride oxidizing NADH with time		
time (min)	light absorption value	conversion rate (%)
8	0.1406	88.70
9	0.1133	90.89
10	0.092	92.60
11	0.0751	93.96
12	0.0615	95.06
13	0.0506	95.93
14	0.042	96.62
15	0.0352	97.17
16	0.0299	97.60
17	0.0255	97.95
18	0.0219	98.24
19	0.0192	98.46
20	0.0169	98.64

Example 7: 1,10-ethyleneisalloxazine chloride oxidizing NADH to NAD⁺

[0062] At 30° C., in a 10 mL 50 mM potassium phosphate buffer of pH 7, the initial concentration of NADH was 0.2 mM, the concentration of 1,10-ethyleneisalloxazine chloride was 4 μM, the reaction solution was connected with the outside air. Change of the light absorption value of the reaction system at 340 nm was detected every 2 minutes, the conversion rate of NADH was calculated, and the results are shown in Table 4.

TABLE 4

Conversion rate with time of 1,10-ethyleneisalloxazine chloride oxidizing NADH		
time (min)	light absorption value	conversion rate (%)
0	1.244	0.00
1	0.7525	39.51
2	0.616	50.48
3	0.5059	59.33
4	0.4166	66.51
5	0.3436	72.38
6	0.2839	77.18
7	0.2354	81.08
8	0.1957	84.27
9	0.163	86.90
10	0.1363	89.04
11	0.1145	90.80
12	0.0964	92.25
13	0.0816	93.44
14	0.0693	94.43
15	0.0591	95.25
16	0.0506	95.93
17	0.0436	96.50
18	0.038	96.95
19	0.0334	97.32
20	0.0295	97.63

Example 8:

7-trifluoromethyl-1,10-ethyleneisalloxazine chloride oxidizing NADH at different pHs

[0063] At 30° C., respectively in 10 mL 50 mM potassium phosphate buffer of pH 4-10, the initial concentration of NADH was 0.2 mM, the concentration of 7-trifluoromethyl 1,10-ethyleneisalloxazine chloride being added was 0.5 μM, the reaction solution was connected with the outside air, after being reacted for 10 minutes, the light absorption value

at 340 nm was detected, the concentrations of NADH and NAD⁺ were calculated, TON and TOF at different pHs were calculated, and the results are shown in Table 5, when pH<7, TON and TOF were increased with increase in pH, when pH>7, TON and TOF were reduced with increase in pH.

TABLE 5

TON and TOF of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride oxidizing NADH at different pHs		
pH	TON	TOF (min ⁻¹)
4	36.01	3.60
5	38.91	3.89
6	53.06	5.31
7	95.18	9.52
8	72.67	7.27
9	49.84	4.98
10	19.29	1.93

Example 9:

7-trifluoromethyl-1,10-ethyleneisalloxazine chloride oxidizing NADH at different temperatures

[0064] In a 50 mM potassium phosphate buffer of pH 7, the initial concentration of NADH was 0.2 mM, the concentration of 7-trifluoromethyl 1,10-ethyleneisalloxazine chloride was 0.5 μM, the reaction solution was connected with the outside air, after being reacted at 30° C., 40° C., 50° C., 60° C., and 70° C. respectively for 10 minutes, the light absorption value at 340 nm was detected, the concentrations of NADH and NAD⁺ were calculated, TON and TOF at different temperatures were calculated, and the results are shown in Table 6, when T<40° C., changes of TON and TOF with temperature was not large, when T>40° C., TON and TOF were reduced with increase in temperature, the reason is that solubility of air in water is reduced with increase in temperature.

TABLE 6

TON and TOF of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride oxidizing NADH at different pHs		
T (° C.)	TON	TOF (min ⁻¹)
30	96.78	9.68
40	96.14	9.61
50	56.27	5.63
60	48.23	4.82
70	42.77	4.28

Example 10: Substituted derivative of 1,10-ethyleneisalloxazine chloride oxidizing NADH to NAD⁺

[0065] At 30° C., in a 50 mM potassium phosphate buffer of pH 7, the initial concentration of NADH was 0.2 mM, the concentrations of 7-trifluoromethyl-3-methyl-1,10-ethyleneisalloxazine chloride (Compound A), 7-methyl-1,10-ethyleneisalloxazine chloride (Compound B), 8-methyl-1,10-ethyleneisalloxazine chloride (Compound C), 7,8-dimethyl-1,10-ethyleneisalloxazine chloride (Compound D), 3-benzyl-1,10-ethyleneisalloxazine chloride (Compound E) were 0.5 μM, the reaction solution was connected with the outside air, after being reacted for 10 minutes, the

light absorption value at 340 nm was detected, the concentrations of NADH and NAD⁺ were calculated, TON and TOF at different temperatures were calculated, and the results are shown in Table 7.

TABLE 7

TON and TOF of several bridged flavins catalytically oxidizing NADH		
Compound	TON	TOF (min ⁻¹)
A	89.34	8.93
B	70.53	7.05
C	69.98	7.00
D	56.88	5.69
E	75.27	7.53

Example 11:

7-trifluoromethyl-1,10-ethyleneisalloxazine chloride as a catalyst for regenerating NAD⁺ coupled with horse liver alcohol dehydrogenase catalyzing lactonization of 1,4-butanediol to γ -butyrolactone

[0066] At 30° C., in a 1 mL 50 mM potassium phosphate buffer (pH8) of 1,4-butanediol 20 mM, NAD⁺ 0.1 mM, 7-trifluoromethyl-1,10-ethyleneisalloxazine 0.05 mM, catalase 50 U/mL, and horse liver alcohol dehydrogenase 20 U/mL, the reaction solution was connected with the outside air. 200 μ L reaction solution were taken out respectively at 1 h, 2 h, 4 h, 8 h of reaction, and 200 μ L ethyl acetate was added for extraction, the reaction process was detected by gas chromatography, and the results are shown in Table 8:

TABLE 8

Conversion rate with reaction time of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride coupled with horse liver alcohol dehydrogenase catalyzing 1,4-butanediol	
time (h)	conversion rate (%)
1	38
2	71.18
4	93
8	100

Example 12:

7-trifluoromethyl-1,10-ethyleneisalloxazine chloride as a catalyst for regenerating NAD⁺ coupled with horse liver alcohol dehydrogenase catalyzing oxidation of cyclohexanol to cyclohexanone

[0067] At 30° C., in a 1 mL 50 mM potassium phosphate buffer (pH8) of cyclohexanol 20 mM, NAD⁺ 0.1 mM, 7-trifluoromethyl-1,10-ethyleneisalloxazine 0.05 mM, catalase 50 U/mL, horse liver alcohol dehydrogenase 20 U/mL, the reaction solution was connected with the outside air connect. 200 μ L reaction solution were taken out respectively at 1 h, 2 h, 4 h, 8 h, 12 h, 24 h of the reaction, 200 μ L ethyl acetate was added for extraction, the reaction process was detected by gas chromatography, and the results are shown in Table 9:

TABLE 9

Conversion rate with reaction time of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride coupled with horse liver alcohol dehydrogenase catalyzing cyclohexanol	
time (h)	conversion rate (%)
1	31
2	39.23
4	51.36
8	62.3
24	90.72

Example 13:

7-trifluoromethyl-1,10-ethyleneisalloxazine chloride as a catalyst for regenerating NAD⁺ coupled with horse liver alcohol dehydrogenase catalyzing oxidation and lactonization of 3-methyl-1,5-pentanediol to 3-methylvalerolactone

[0068] At 30° C., in a 1 mL 50 mM potassium phosphate buffer (pH8) of 3-methyl-1,5-pentanediol 20 mM, NAD⁺ 0.1 mM, 7-trifluoromethyl-1,10-ethyleneisalloxazine 0.05 mM, catalase 50 U/mL, horse liver alcohol dehydrogenase 20 U/mL, the reaction solution was connected with the outside air. 200 μ L reaction solution were taken out respectively at 1 h, 2 h, 4 h, 8 h, 12 h, 24 h of the reaction, 200 μ L ethyl acetate was added for extraction, the reaction process was detected by gas chromatography, and the results are shown in Table 10.

TABLE 10

Conversion rate with reaction time of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride coupled with horse liver alcohol dehydrogenase catalyzing 3-methyl-1,5-pentanediol	
time (h)	conversion rate (%)
1	59.83
2	71.18
4	82.76
8	93.77
12	98.64
24	100

Example 14:

7-trifluoromethyl-1,10-ethyleneisalloxazine chloride as a catalyst for regeneration NAD⁺ coupled with glucose dehydrogenase catalytically oxidizing glucose to gluconic acid

[0069] At 30° C., in a 10 mL reaction system of glucose 25 mM, NAD⁺ 0.5 mM, 7-trifluoromethyl-1,10-ethyleneisalloxazine 0.5 mM, NaCl 50 mM, catalase 50 U/mL, glucose dehydrogenase 120 U/mol (glucose), the reaction solution was connected with the outside air. 200 μ L reaction solution were taken out respectively at 1 h, 2 h, 4 h, 6 h of the reaction, 800 μ L water was added for dilution, the reaction process was detected by liquid chromatography, and the results are shown in Table 11.

TABLE 11

Conversion rate with reaction time of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride coupled with glucose dehydrogenase catalyzing glucose	
time (h)	conversion rate (%)
1	38.1
2	55.0
4	89.6
6	100

Example 15:

7-trifluoromethyl-1,10-ethyleneisalloxazine chloride as a catalyst for regenerating NAD⁺ coupled with glucose dehydrogenase catalytically oxidizing xylose to xylonic acid

[0070] At 30° C., in a 10 mL reaction system of xylose 25 mM, NAD⁺ 0.5 mM, 7-trifluoromethyl-1,10-ethyleneisalloxazine 0.5 mM, NaCl 50 mM, catalase 50 U/mL, glucose dehydrogenase 120 U/mol (xylose), the reaction solution was connected with the outside air. 200 μ L reaction solution were taken out respectively at 1 h, 2 h, 4 h, 6 h, 12 h, 24 h of the reaction, 800 μ L water was added for dilution, the reaction process was detected by liquid chromatography, and the results are shown in Table 12.

TABLE 12

Conversion rate with reaction time of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride coupled with glucose dehydrogenase catalyzing xylose	
time (h)	conversion rate (%)
1	17.5
2	32.3
4	52.1
6	65.9
8	76.8
12	82.9
24	100

Example 16:

7-trifluoromethyl-1,10-ethyleneisalloxazine chloride as a catalyst for regenerating NAD⁺ coupled with glycerol dehydrogenase catalytically oxidizing glycerol to 1,3-dihydroxyacetone

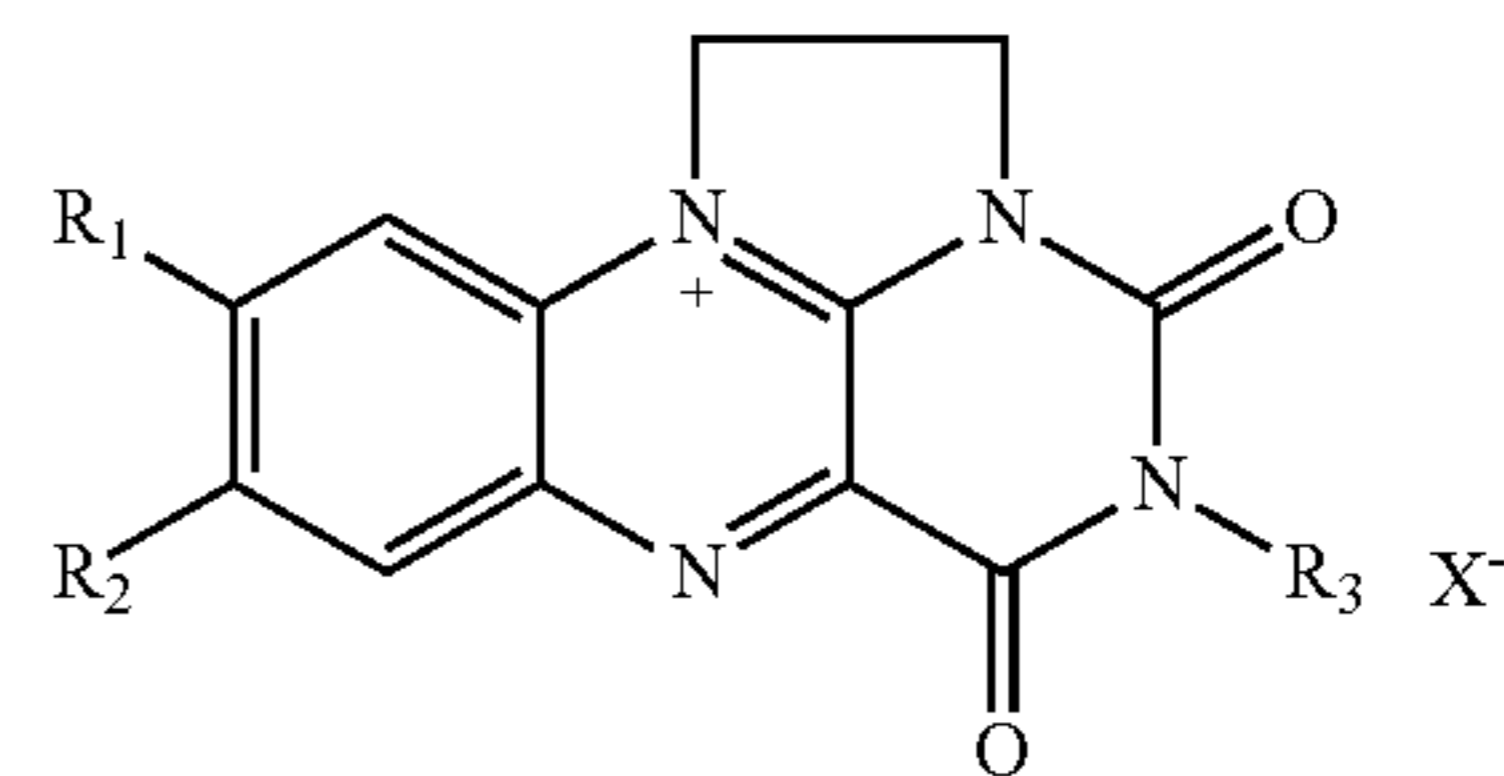
[0071] At 30° C., in 10 mL reaction system (potassium phosphate buffer of 50 mM and pH 10) of glycerol 25 mM, NAD⁺ 0.5 mM, 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride 0.5 mM, atalase 50 U/mL, glycerol dehydrogenase 120 U/mol, the reaction solution was connected with the outside air. 200 μ L reaction solution were taken out respectively at 1 h, 2 h, 4 h, 8 h, 12 h, 24 h of the reaction, 800 μ L of water was added for dilution, the reaction process detected by liquid chromatography, and the results are shown in Table 13.

TABLE 13

Conversion rate with reaction time of 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride coupled with glycerol dehydrogenase catalyzing glycerol	
time (h)	conversion rate (%)
1	235
2	56.3
4	51.7
8	72.6
12	86.5
24	100

What is claimed is:

1. A method for regenerating chemically an oxidized coenzyme NAD(P)⁺, characterized in that, under an oxygen or air atmosphere condition, adding a catalytic amount of a bridged flavin, and oxidizing NAD(P)H to obtain NAD(P)⁺.
2. The method according to claim 1, characterized in that, the bridged flavin has a formula as follow:



wherein, R₁ and R₂ are independently selected from a group consisting of hydrogen, methyl, trifluoromethyl, methoxyl, halogen atom, nitro and amino; R₃ is selected from a group consisting of hydrogen, C1-C5 alkyl, phenyl and benzyl; X⁻ is selected from a group consisting of halide ion, nitrate radical and trifluoromethanesulfonic acid radical.

3. The method according to claim 1, characterized in that, the bridged flavin is 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride, 8-chloro-1,10-ethyleneisalloxazine chloride, or 1,10-ethyleneisalloxazine chloride.

4. The method according to claim 1, characterized in that, the mole number of the catalytic amount of the bridged flavin is 0.1-5% of the mole number of the NAD(P)H.

5. The method according to claim 1, characterized in that, the condition for oxidizing NAD(P)H into NAD(P)⁺ is in a pH 4-10 and at a temperature 30-70° C.

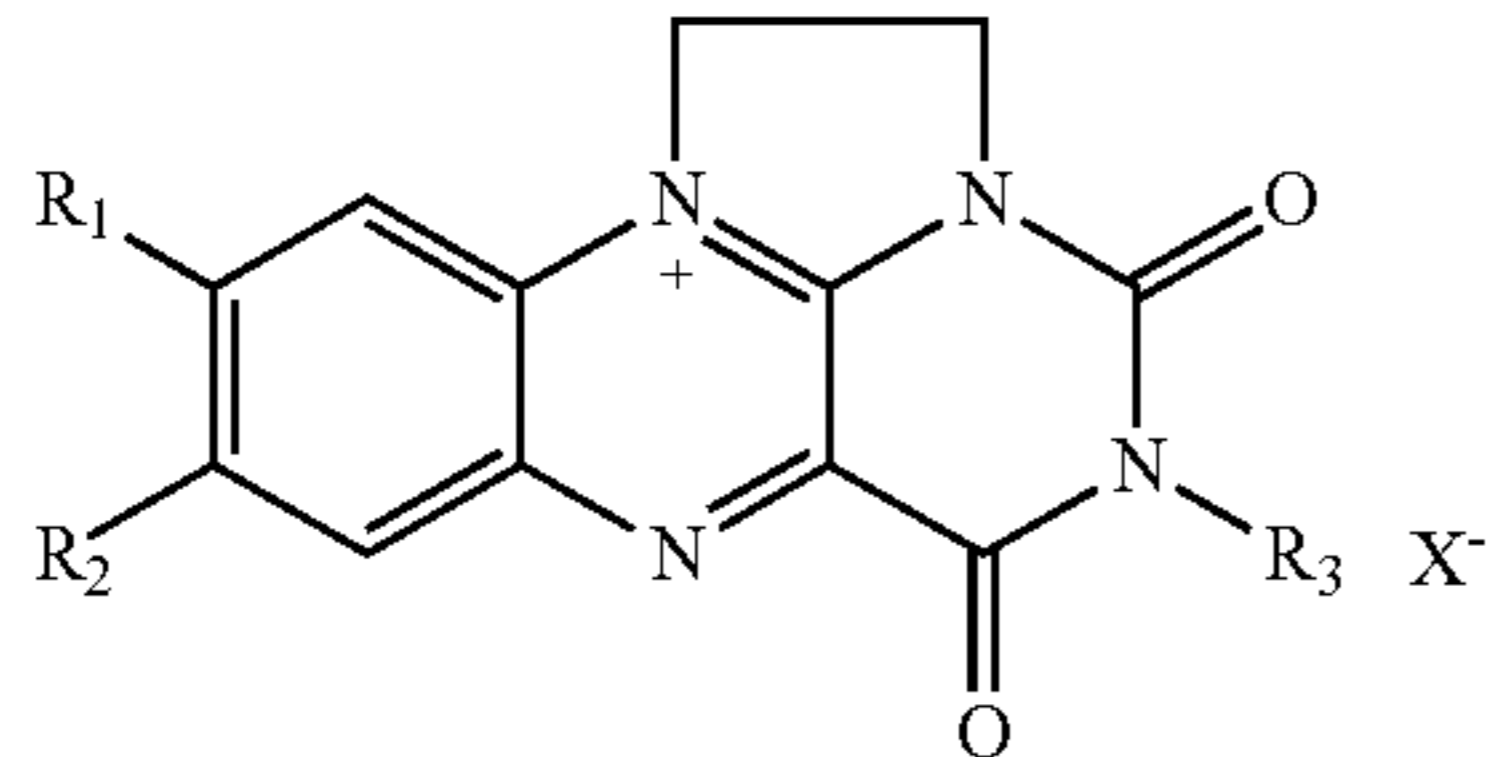
6. The method according to claim 1, characterized in that, the bridged Flavin is coupled with an oxidation reaction catalyzed by a NAD(P)⁺-dependent oxidoreductase, forming a regeneration circulation system of the coenzyme NAD(P)⁺.

7. The method according to claim 6, characterized in that, the NAD(P)⁺-dependent oxidoreductase is one or more enzymes selected from a group consisting of EC1.1.1.X, EC1.2.1.X, EC1.3.1.X, EC1.4.1.X, EC1.5.1.X, EC1.6.1.X, EC1.7.1.X, EC1.8.1.X, EC1.10.1.X, EC1.12.1.X, EC1.13.1.X, EC1.16.1.X, EC1.17.1.X, EC1.18.1.X, EC1.20.1.X and EC1.22.1.X.

8. The method according to claim 7, characterized in that, the NAD(P)⁺-dependent oxidoreductase is horse liver alcohol dehydrogenase, glucose dehydrogenase or glycerol dehydrogenase.

9. A process for an oxidation reaction by using a NAD(P)⁺-dependent oxidoreductase as a catalyst, characterized in that, using a bridged flavin as a NAD(P)⁺ regeneration catalyst, under oxygen or air atmosphere condition, oxidizing NAD(P)H into NAD(P)⁺, forming a regeneration circulation system of coenzyme NAD(P)⁺.

10. The process according to claim **9**, characterized in that, the bridged flavin has a formula as follow:



wherein, R₁ and R₂ are independently selected from a group consisting of hydrogen, methyl, trifluoromethyl, methoxyl, halogen atom, nitro and amino; R₃ is selected from a group consisting of hydrogen, C1-C5 alkyl, phenyl and benzyl; X⁻ is selected from a group consisting of halide ion, nitrate radical and trifluoromethanesulfonic acid radical.

11. The process according to claim **10**, characterized in that, the bridged flavin is 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride, 8-chloro-1,10-ethyleneisalloxazine chloride, or 1,10-ethyleneisalloxazine chloride.

12. The process according to claim **9**, characterized in that, the mole number catalyzed by the bridged flavin is 0.1-5% of the mole number of a substrate catalyzed by an enzyme.

13. The process according to claim **9**, characterized in that, the condition for oxidizing NAD(P)H into NAD(P)⁺ is in pH 4-10 and at temperature 30-70° C.

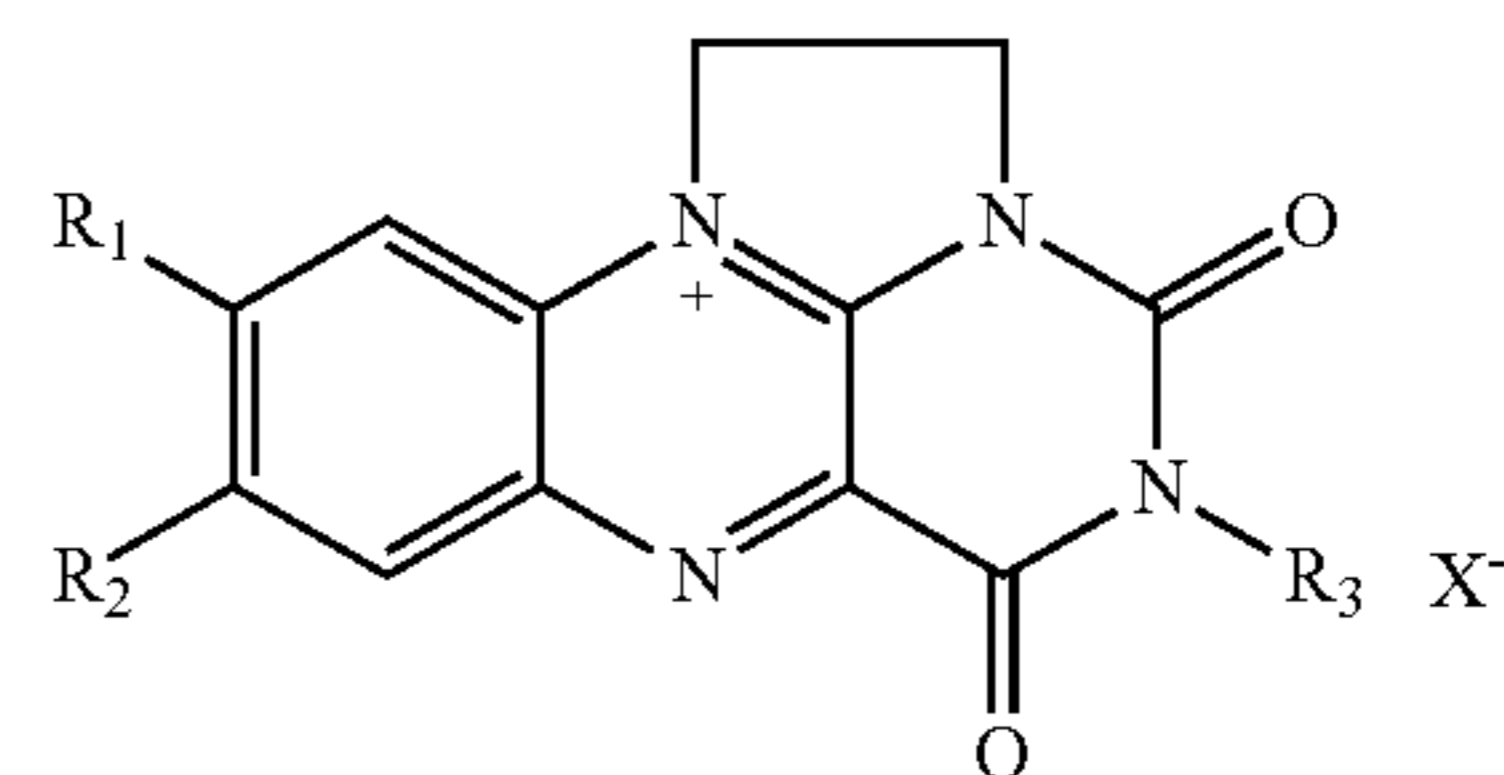
14. The process according to claim **9**, characterized in that, the NAD(P)⁺-dependent oxidoreductase is one or more enzymes selected from a group consisting of EC1.1.1.X, EC1.2.1.X, EC1.3.1.X, EC1.4.1.X, EC1.5.1.X, EC1.6.1.X,

EC1.7.1.X, EC1.8.1.X, EC1.10.1.X, EC1.12.1.X, EC1.13.1.X, EC1.16.1.X, EC1.17.1.X, EC1.18.1.X, EC1.20.1.X and EC1.22.1.X.

15. The process according to claim **14**, characterized in that, the NAD(P)⁺-dependent oxidoreductase is horse liver alcohol dehydrogenase, glucose dehydrogenase or glycerol dehydrogenase.

16. A method for using a bridged flavin as a catalyst for oxidizing NAD(P)H into NAD(P)⁺.

17. The method according to claim **16**, characterized in that, the bridged flavin has formula as follow:



wherein, R₁ and R₂ are independently selected from a group consisting of hydrogen, methyl, trifluoromethyl, methoxyl, halogen atom, nitro and amino; R₃ is selected from a group consisting of hydrogen, C1-C5 alkyl, phenyl and benzyl; X⁻ is selected from a group consisting of halide ion, nitrate radical and trifluoromethanesulfonic acid radical.

18. The method according to claim **17**, characterized in that, the bridged flavin is 7-trifluoromethyl-1,10-ethyleneisalloxazine chloride, 8-chloro-1,10-ethyleneisalloxazine chloride, or 1,10-ethyleneisalloxazine chloride.

19. The method according to claim **16**, characterized in that, the mole number catalyzed by the bridged flavin is 0.1-5% of the mole number of NAD(P)H.

20. The method according to claim **16**, characterized in that, the condition for oxidizing NAD(P)H into NAD(P)⁺ is in pH 4-10 and at temperature 30-70° C.

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