CLAIMS

1. A hydrate of dimethylaminomicheliolide fumarate, characterized in that, the hydrate is in a crystalline form D, the molar ratio of dimethylaminomicheliolide fumarate to water is 1:1, and the hydrate has a molecular formula of $C_{17}H_{27}NO_3 \cdot C_4H_4O_4 \cdot H_2O$; a thermogravimetric analysis shows a weight loss of 3.97%–4.22% before decomposition; a differential scanning calorimetry pattern shows a dehydration endothermic peak at 75±5 °C and a characteristic melting peak at 148±5 °C,

   wherein the hydrate in crystalline form D has characteristic peaks at 2θ angles of 7.8±0.2°, 11.1±0.2°, 11.4±0.2°, 12.6±0.2°, 12.9±0.2°, 14.4±0.2°, 15.3±0.2°, 17.0±0.2°, 18.7±0.2°, 19.7±0.2°, 20.6±0.2°, 21.0±0.2°, 22.5±0.2°, 23.7±0.2°, 24.3±0.2°, 25.5±0.2° and 26.2±0.2° in an X-ray powder diffraction pattern using Cu-Kα radiation, wherein the peak at 7.8±0.2° is an initial peak; the characteristic peak at 20.6±0.2° has a relative intensity of 100%; the crystalline form D is in an orthorhombic crystal system and has a space group of *P*212121, a cell parameter of a = 8.8346(18) Å, b = 14.796(3) Å, c= 16.385(3) Å, α = 90°, β = 90°, and γ = 90°; and a cell volume of 2141.8(8) Å3.

2. The hydrate of dimethylaminomicheliolide fumarate according to claim 1, characterized in that, the hydrate also has characteristic peaks at 2θ angles of 10.5±0.2°, 11.7±0.2°, 12.0±0.2°, 15.6±0.2°, 15.9±0.2°, 16.2±0.2°, 21.3±0.2°, 22.1±0.2°, 23.0±0.2°, 26.4±0.2°, 27.2±0.2°, 28.2±0.2°, 28.6±0.2°, 29.3±0.2°, 30.4±0.2° and 31.1±0.2° in an X-ray powder diffraction pattern using Cu-Kα radiation.

3. A preparation method for the hydrate of dimethylaminomicheliolide fumarate according to claim 1 or 2, characterized in that, the preparation is implemented by reactive crystallization: under the action of stirring, adding dimethylaminomicheliolide and fumaric acid to a mixed solvent system of a solvent S1 and a solvent S2 at a constant temperature of 30 °C–70 °C, with the mass ratio of the solvent S2 to the solvent S1 being (0–3):1 and the molar ratio of dimethylaminomicheliolide to fumaric acid being (1–1.6):1; after 5–10 h of reaction, filtering the reaction mixture and drying the residue at 25–45 °C under normal pressure for 6–10 h to obtain dimethylaminomicheliolide fumarate in a crystalline form D;

   the solvent S1 is a mixed solvent of water and any one of acetone, tetrahydrofuran, 1,4-dioxane, acetonitrile and methyl isobutyl ketone;

   the solvent S2 is a mixed solvent of an ester and an ether;

the ester solvent is selected from any one or two of methyl acetate, ethyl acetate, hexyl acetate and isopropyl acetate;

the ether solvent is selected from any one or two of diethyl ether, methyl ethyl ether, methyl *tert*-butyl ether, dipropyl ether, dibutyl ether, ethylene glycol dimethyl ether, ethylene glycol monomethyl ether, 1,4-dioxane, tetrahydrofuran and 2-methyl tetrahydrofuran;

the mass ratio of any one of acetone, tetrahydrofuran, 1,4-dioxane, acetonitrile and methyl isobutyl ketone to water in the solvent S1 is (5–10):1;

the mass ratio of the ester solvent to the ether solvent in the solvent S2 is (1–3):1;

the mass ratio of the solid starting material dimethylaminomicheliolide to S1 is 1:(6–10).

4. A pharmaceutical composition, which comprises the hydrate of dimethylaminomicheliolide fumarate according to claim 1 or 2 and a pharmaceutically acceptable auxiliary material.

5. The pharmaceutical composition according to claim 4, wherein the pharmaceutical composition is in a tablet, capsule or granule dosage form.

6. Use of the hydrate of dimethylaminomicheliolide fumarate according to claim 1 or 2 in preparing a medicament for the treatment or prevention of a disease or condition, wherein the disease or condition is cancer selected from leukemia, breast cancer, prostate cancer, nasopharyngeal cancer, large intestine cancer, lung cancer, liver cancer, esophageal cancer, gastric cancer, intestinal cancer, renal cancer, oral cancer, Hodgkin's lymphoma, pancreatic cancer, colorectal cancer, cervical cancer, non-Hodgkin's lymphoma, glioma, melanoma, bladder cancer, ovarian cancer, thyroid cancer and Kaposi's sarcoma.