

ABSTRACT:

An acid salt and a crystal form of a pyrazole-containing polycyclic derivative represented by formula (I-a), and a preparation method therefor, a pharmaceutical composition containing a therapeutically effective amount of the crystal form, and an intermediate for preparing the derivative. In particular, the present invention relates to the use of the salt and the crystal form of the compound represented by general formula (I-a) as a P2X3 inhibitor in the preparation of a drug for treating neurogenic diseases.

