

CLAIMS

1. Co-crystal of Gabapentin, Ketoprofen and Lysine wherein the molar ratio of the components is 1:1:1.
2. The co-crystal as claimed in claim 1, characterized by the following XRPD diffraction peaks: 3.6, 9.5, 9.6, 18.5 and 20.0 degrees 2-theta $\pm$ 0.2 degrees 2-theta, preferably further characterized by the following XRPD diffraction peaks: 15.4, 17.8, 21.0, 21.8 and 24.2 degrees 2-theta $\pm$ 0.2 degrees 2-theta.
3. The co-crystal as claimed in claim 1 or 2, further characterized by one or more of the following:
 - a DSC thermogram with the endothermic peak corresponding to the melting point at about 141.4°C as reported in Figure 2,
 - a TGA thermogram as reported in Figure 5,
 - FT Raman and FT-IR absorption bands as reported in Tables 6 and 7,
 - solution ^1H -NMR signals as reported in Figure 8 and Table 8, and
 - solid state ^{13}C CPMAS signals as reported in Figures 9 to 11 and in Table 9.
4. The co-crystal as claimed in any one of claims 1 to 3, wherein said Ketoprofen is (S)-Ketoprofen.
5. The co-crystal as claimed in any one of claims 1 to 4, wherein said Lysine is (S)-Lysine.
6. The co-crystal as claimed in any one of claims 1 to 5 for use as a medicament.
7. The co-crystal as claimed in any one of claims 1 to 6 for use in the prevention, reduction or treatment of pain and/or inflammation.
8. The co-crystal for use as claimed in claim 7, wherein said pain is acute or chronic pain.
9. The co-crystal for use as claimed in claims 7 or 8, wherein said pain is selected from headache, toothache, menstrual pain, muscle pain, neuropathic pain, pain associated to neuroinflammation, diabetic neuropathy, cancer pain, osteoarthritis, low back pain, sciatalgia, fibromyalgia, trigeminal neuralgia; post-surgical and post-operative pain, post herpetic neuralgia, rheumatoid arthritis, ankylosing spondylitis, frozen shoulder, phantom limb pain or HIV pain.

10. A pharmaceutical composition comprising a co-crystal as claimed in any one of claims 1 to 5 and a least one pharmaceutically acceptable excipient.
11. The pharmaceutical composition as claimed in claim 10, which contains 0.5-60% by weight of the co-crystal and 40-99.5% by weight of one or more pharmaceutically acceptable excipients.
12. The pharmaceutical composition as claimed in claims 10 or 11 which is an oral solid composition.
13. Process for the preparation of a co-crystal as claimed in any one of claims 1 to 5, which comprises:
- a) suspending Gabapentin, Ketoprofen and Lysine in a suitable solvent,
- b) dissolving Gabapentin, Ketoprofen and Lysine, optionally by heating the suspension, optionally under stirring, till a clear solution is obtained,
- c) subsequently cooling the solution, and
- d) optionally adding an anti-solvent.
14. Process as claimed in claim 13, wherein said Ketoprofen and Lysine in step a) are in form of a Ketoprofen Lysine salt or a co-crystal.
15. Process as claimed in claims 13, wherein said Ketoprofen is the free acid and/or said Lysine is in neutral form.
16. Process as claimed in any one of claims 13 to 15, wherein said Gabapentin is in neutral form (zwitterionic internal salt).
17. Process as claimed in any one of claims 13 to 16 wherein in step a) the molar ratio of Gabapentin vs Ketoprofen is between 1:1 and 1.5:1, preferably between 1:1 and 1.2:1, more preferably is about 1:1.