

ONCOLOGY PORTFOLIO - Approved Products & Pipeline



*Approved at least by EMA. Vinorelbine was approved by National Health Authorities (before EU centralized procedure implementation).

1.Commercialized in partnership with Puma Biotechnology. 2.Commercialized in partnership with Atara Biotherapeutics. 3.Encorafenib + cetuximab is approved for BRAF^{V600E}-mutant mCRC, after prior systemic therapy. Encorafenib + binimetinib is approved for unresectable or metastatic BRAF-mutant melanoma, in adult patients. 4.Developed and launched in partnership with Array Biopharma (partnered with Pfizer since 2020). 5.Clinical development under Pfizer sponsorship. 6.ANCHOR CRC study status: completed. 7.Originated by Kinnate Biopharma Inc. Investigational Pan-RAF inhibitor, Exarafenib, acquired by Pierre Fabre Laboratories. 8.Co-development in partnership with Scorpion Therapeutics Inc., Boston, USA. The sponsor of these studies could be Pierre Fabre Medical Care or Scorpion Therapeutics Inc. 9.Originated by Vertical Bio AG, acquired by Pierre Fabre Laboratories. 10.Discovery partnership with Vernalis Ltd.

This display includes ongoing clinical trials for both approved and investigational compounds. Inclusion in this display does not imply regulatory approval for these compounds or all indications. Products should be used in accordance with their prescribing information. Information about the trials can be found at www.ClinicalTrials.gov