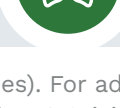


# ONCOLOGY PORTFOLIO - Approved Products & Pipeline

| INN /COMPOUND                          | CLASS            | TARGET & MODALITY                                                         | SETTING                                                                                                                                                                                                                                     | Discovery | Preclinical | Phase I | Phase II | Phase III | In Registration | Approved* |
|----------------------------------------|------------------|---------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------|-------------|---------|----------|-----------|-----------------|-----------|
| PFL-002 <sup>1</sup>                   | Targeted therapy | Anti-c-MET antibody degrader                                              |  <b>NSCLC</b> / solid tumors with MET mutations or amplification                                                                                         |           |             |         |          |           |                 |           |
| PFL-241 <sup>2</sup>                   | Targeted therapy | Mutant-selective 4 <sup>th</sup> generation EGFR small-molecule inhibitor |  <b>NSCLC</b> with <i>EGFR</i> exon 19/21 + C797S mutations                                                                                              |           |             |         |          |           |                 |           |
| PFL-721 <sup>2</sup>                   | Targeted therapy | Mutant-selective EGFR-exon 20 small-molecule inhibitor                    |  <b>NSCLC</b> with <i>EGFR</i> exon 20 insertion mutation                                                                                                |           |             |         |          |           |                 |           |
| Exarafenib <sup>3</sup>                | Targeted therapy | Pan-RAF small-molecule inhibitor                                          |  <i>NRAS</i> mutant <b>MELANOMA</b> and other <i>BRAF</i> / <i>NRAS</i> mutant solid tumors                                                              |           |             |         |          |           |                 |           |
| Not disclosed <sup>4</sup>             | Targeted therapy | Multiple targets                                                          |  <b>ONCOLOGY</b>                                                                                                                                         |           |             |         |          |           |                 |           |
| Encorafenib <sup>5</sup>               | Targeted therapy | BRAF inhibitor + other agents                                             |  Metastatic <i>BRAF</i> <sup>V600E</sup> -mutant <b>CRC</b> in first-line treatment in combination with cetuximab and FOLFOX <sup>6</sup>                |           |             |         |          |           |                 |           |
|                                        |                  | BRAF inhibitor + other agents                                             |  Metastatic <i>BRAF</i> <sup>V600E</sup> -mutant <b>CRC</b> after prior systemic therapy in combination with cetuximab                                   |           |             |         |          |           |                 |           |
| Encorafenib + binimetinib <sup>5</sup> | Targeted therapy | BRAF inhibitor + MEK inhibitor                                            |  Unresectable or metastatic <i>BRAF</i> <sup>V600</sup> -mutant <b>MELANOMA</b>                                                                         |           |             |         |          |           |                 |           |
|                                        |                  | BRAF inhibitor + MEK inhibitor                                            |  Advanced <i>BRAF</i> <sup>V600E</sup> -mutant <b>NSCLC</b>                                                                                            |           |             |         |          |           |                 |           |
| Tabelecleucel <sup>7</sup>             | Cell therapy     | Allogeneic T-cell immunotherapy                                           |  Relapsed or refractory Epstein-Barr virus positive post-transplant lymphoproliferative disease ( <b>EBV+ PTLD</b> ) after at least one prior therapy  |           |             |         |          |           |                 |           |
| Neratinib <sup>8</sup>                 | Targeted therapy | Pan-HER inhibitor                                                         |  HER2 amplified / overexpressed / HR+ early <b>BREAST CANCER</b> after adjuvant trastuzumab-based therapy less than one year ago                       |           |             |         |          |           |                 |           |
| Vinflunine                             | Chemotherapy     | Vinca Alkaloid                                                            |  Advanced or metastatic transitional cell carcinoma of the <b>UROTHELIAL</b> tract as monotherapy after failure of a prior platinum-containing regimen |           |             |         |          |           |                 |           |
| Vinorelbine                            | Chemotherapy     | Vinca Alkaloid                                                            |  Advanced <b>BREAST CANCER</b> as monotherapy or in combination with other agents                                                                      |           |             |         |          |           |                 |           |
|                                        |                  | Vinca Alkaloid                                                            |  Advanced <b>NSCLC</b> as monotherapy or in combination with other chemotherapies                                                                      |           |             |         |          |           |                 |           |
|                                        |                  | Vinca Alkaloid                                                            |  Adjuvant treatment of <b>NSCLC</b> in combination with platinum-based chemotherapy                                                                    |           |             |         |          |           |                 |           |

\*The settings refer to the therapeutic indications approved in the EU countries where the product is registered (it may differ in non EU countries). For adult patients only, except Tabelecleucel which is indicated for paediatric patients 2 years of age and older.  
1.Originated by Vertical Bio AG, acquired by Pierre Fabre Laboratories. 2.Originated by Scorpion Therapeutics acquired by Pierre Fabre Laboratories. 3.Originated by Kinnate Biopharma Inc. Investigational Pan-RAF Inhibitor, Exarafenib, acquired by Pierre Fabre Laboratories. 4.Discovery partnership with Vernalis Lt and RedRidge Bio. 5.Developed and launched in partnership with Array Biophama (partnered with Pfizer since 2020). 6.Clinical development and sponsorship done solely by Pfizer. 7.Commercialized in partnership with Atara Biotherapeutics. 8.Commercialized in partnership with Puma Biotechnology.

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