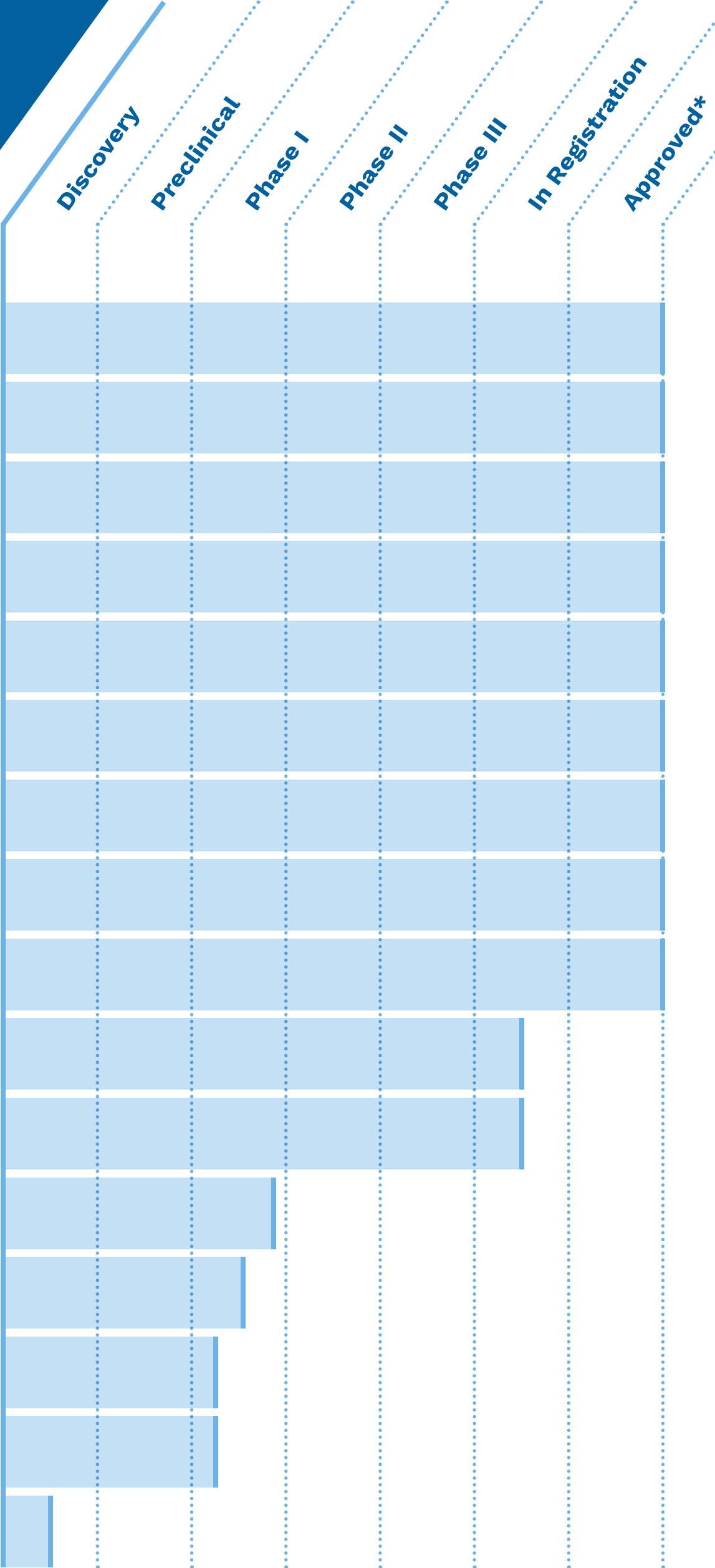


ONCOLOGY PORTFOLIO - Approved Products & Pipeline

INN /COMPOUND	CLASS	TARGET & MODALITY	INDICATION OR DISEASE
Vinorelbine	Chemotherapy	<div>Vinca Alkaloid</div> <div></div>	Advanced BREAST CANCER as monotherapy or in combination with other agents
		<div>Vinca Alkaloid</div> <div></div>	Advanced NSCLC as monotherapy or in combination with other chemotherapies
		<div>Vinca Alkaloid</div> <div></div>	Adjuvant treatment of NSCLC in combination with platinum-based chemotherapy
Vinflunine	Chemotherapy	<div>Vinca Alkaloid</div> <div></div>	Advanced or metastatic transitional-cell carcinoma of the UROTHELIAL tract
Neratinib ¹	Targeted therapy	<div>Pan-HER inhibitor</div> <div></div>	HER2 amplified / overexpressed / HR+ early BREAST CANCER (extended adjuvant post trastuzumab based therapy)
Tabelecleucel ²	Cell therapy	<div>Allogeneic T-cell immunotherapy</div> <div></div>	Epstein-Barr virus associated post-transplant lymphoproliferative disease (EBV + PTLD)
Encorafenib ³ / Binimetinib ⁴ (± other agents)	Targeted therapy	<div>BRAF inhibitor + MEK inhibitor</div> <div></div>	Unresectable or metastatic <i>BRAF</i> ^{V600} -mutant MELANOMA
		<div>BRAF inhibitor</div> <div></div>	Metastatic <i>BRAF</i> ^{V600E} -mutant CRC after prior systemic therapy
		<div>BRAF inhibitor + MEK inhibitor</div> <div></div>	Metastatic <i>BRAF</i> ^{V600E} -mutant NSCLC ⁵
		<div>BRAF inhibitor</div> <div></div>	Previously treated metastatic <i>BRAF</i> ^{V600E} -mutant CRC in Chinese population - NAUTICAL study
		<div>BRAF inhibitor + MEK inhibitor</div> <div></div>	Metastatic <i>BRAF</i> ^{V600E} -mutant NSCLC in Chinese population - OCEAN study
Exarafenib ⁶	Targeted therapy	<div>Pan-RAF small molecule inhibitor</div> <div></div>	<i>NRAS</i> mutant MELANOMA and other <i>BRAF</i> / <i>NRAS</i> mutant solid tumors
PFL-721 / STX-721 ⁷	Targeted therapy	<div>Mutant-selective EGFR-exon20 small molecule inhibitor</div> <div></div>	NSCLC with <i>EGFR</i> exon 20 insertion mutation
PFL-241 / STX-241 ⁷	Targeted therapy	<div>Mutant-selective 4th generation EGFR small molecule inhibitor</div> <div></div>	NSCLC with <i>EGFR</i> exon 19/21 + C797S mutations
PFL-002 / VERT-002 ⁸	Targeted therapy	<div>Anti-cMET antibody degrader</div> <div></div>	NSCLC / solid tumors with mutations or amplification of <i>MET</i>
Not-disclosed ⁹	Targeted therapy	<div>Multiple targets</div> <div></div>	ONCOLOGY



*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 Biophama (partnered with Pfizer since 2020). 5.Clinical development under Pfizer sponsorship. 6.Originated by Kinnate Biopharma Inc. Investigational Pan-RAF Inhibitor, Exarafenib, acquired by Pierre Fabre Laboratories. 7.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. 8.Originated by Vertical Bio AG, acquired by Pierre Fabre Laboratories. 9.Discovery partnership with Vernalis Lt and RedRidge Bio.

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