## **ONCOLOGY PORTFOLIO - Approved Products & Pipeline**

										e Fabre
INN /Compound	Class	Target & Modality	Indication or Disease	Discovery	Preclinical	<b>Phase I</b>	Phase II	Phase III	In Registration	Approved
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						i i	
		Vinca Alkaloid	Adjuvant treatment of <b>NSCLC</b> in combination with platinum-based chemotherapy				1 1 1	1		
Vinflunine	Chemotherapy	Vinca Alkaloid	Advanced or metastatic transitional-cell carcinoma of the <b>UROTHELIAL</b> tract							
Neratinib <sup>1</sup>	Targeted therapy	Pan-HER inhibitor	HER2 amplified / overexpressed / HR+ early <b>BREAST CAN</b> (extended adjuvant post trastuzumab based therapy)	CER						
Tabelecleucel <sup>2</sup>	Cell therapy	Allogeneic T-cell immunotherapy	Epstein-Barr virus associated post-transplant lymphoproliferative disease <b>(EBV + PTLD)</b>							
Encorafenib³ / Binimetinib⁴ (± other agents)	Targeted therapy	BRAF inhibitor + MEK inhibitor	Unresectable or metastatic BRAF <sup>v600</sup> -mutant <b>MELANOMA</b>							
		BRAF inhibitor	Metastatic BRAF <sup>V600E</sup> -mutant <b>CRC</b> after prior systemic therapy				1 1	1 1 1		
		BRAF inhibitor + MEK inhibitor	Metastatic BRAF <sup>V600E</sup> -mutant <b>NSCLC</b> <sup>5</sup>				I I I	1 1 1		
		BRAF inhibitor + MEK inhibitor	Metastatic BRAF <sup>V600E</sup> -mutant <b>NSCLC</b> in Chinese population - OCEAN study							
		BRAF inhibitor	Previously untreated metastatic BRAF <sup>V600E</sup> -mutant <b>CRC</b> <sup>6</sup> - ANCHOR CRC study							
		BRAF inhibitor	Previously treated metastatic BRAF <sup>V600E</sup> -mutant <b>CRC</b> in Chinese population - NAUTICAL study							
Exarafenib <sup>7</sup>	Targeted therapy	Pan-RAF small molecule inhibitor	NRAS mutant <b>MELANOMA</b> and other BRAF / NRAS mutant solid tumors							
PFL-721/STX-7218	Targeted therapy	Mutant-selective EGFR-exon20 small molecule inhibitor	NSCLC with EGFR exon 20 insertion mutation				       			
PFL-241/STX-2418	Targeted therapy	Mutant-selective 4 <sup>th</sup> generation EGFR small molecule inhibitor	NSCLC with EGFR exon 19/21 + C797S mutations							
PFL-002/VERT-0029	Targeted therapy	Anti-cMET antibody degrader	• <b>NSCLC</b> / solid tumors with mutations or amplification of MET				1 1 1 1 1			
Not-disclosed <sup>10</sup>	Targeted therapy	Multiple targets	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 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. 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 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

