

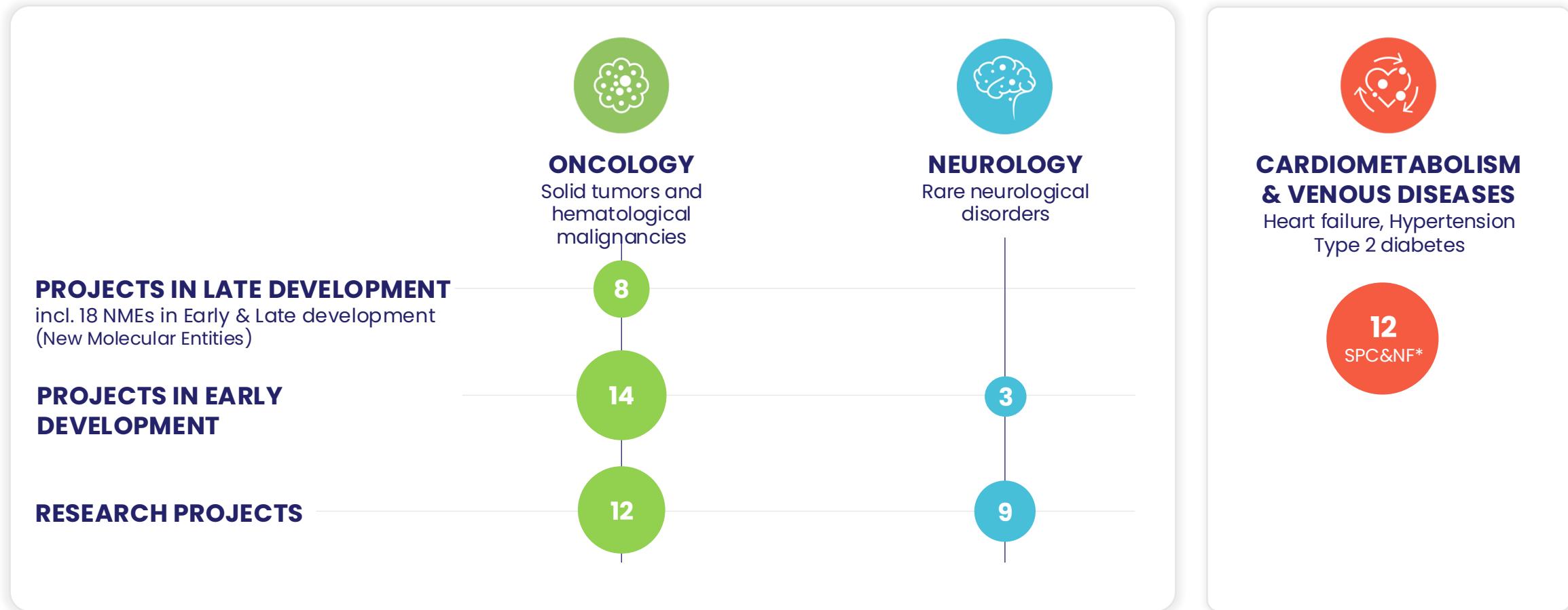
Servier Pipeline

January 2026



SERVIER
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FOCUSED ON EXPANDING OUR PIPELINE



Oncology – Solid tumors

Données à janvier 2026

	Cible	Indication	DPC	Phase I/II	Phase II	Phase III Potential registration	Partenaire
DAROVASERTIB	PKC	Uveal melanoma					IDEA BIOSCIENCES
IVOSIDENIB	IDH1	Chondrosarcoma					
IVOSIDENIB / + Durvalumab +Gemcitabine/Cisplatin		Cholangiocarcinoma					
VORASIDENIB / + temozolamide	IDH1/2	Glioma					
VORASIDENIB / + pembrolizumab		Glioma					
S95018	TIM3						
S95024	CD73	Non-small Cell Lung Cancer + in association					
S95029	NKG2A						
S95035	MAT2A	Gastric cancer					
S241656	RAS/RAF						
S234821	DSC*	Solid tumors					

* Données Scientifiques Confidential

Oncology – Hematological malignancies

	Cible	Indication	DPC	Phase I/II	Phase II	Phase III Potential registration	Partenaire
IVOSIDENIB / + 7+3 (chemotherapy)							
IVOSIDENIB / Azacitidine / Venetoclax	IDH1	Acute Myeloid Leukemia (new indication)					
IVOSIDENIB							
S243249	MENIN	<ul style="list-style-type: none"> Acute Myeloid Leukemia Acute Lymphoblastic Leukemia 					
S236200	DSC*						Vernalis
S247567	DSC*	Hematological malignancies					

	Target	Indication	DPC	Phase I/II	Phase II	Phase III Potential registration
Genetic autism spectre	BK channel	<p>Fragile X syndrome is a genetic disorder caused by a mutation in the FMR1 gene, leading to a deficiency of the fragile X mental retardation protein (FMRP). This protein is crucial for normal brain development and function. The absence of FMRP disrupts synaptic plasticity, which is essential for learning and memory, resulting in cognitive impairments and behavioral challenges.</p> <p>Patients with Fragile X syndrome often exhibit a range of symptoms, including intellectual disability, anxiety, and social difficulties. The condition can also manifest itself through physical features such as an elongated face and enlarged ears.</p>				

	Target	Indication	DPC	Phase I/II	Phase II	Phase III Potential registration
Developmental and epileptic encephalopathies AntiSensOligonucleotid	KCNT1	<p>KCNT1-related Developmental and Epileptic Encephalopathy (DEE) are a group of severe neurological disorders characterized by very early-onset seizures and significant developmental delays. KCNT1-DEE is caused by genetic mutations in the KCNT1 gene that disrupt normal brain development and function. The seizures are most often refractory to standard anti-epileptic medication, complicating treatment.</p> <p>Patients with KCNT1-DEE experience a range of symptoms, including severe cognitive impairments, motor deficits, and behavioral issues. The impact on family dynamics and overall quality of life is profound.</p> <p>More details about ASO technology</p>				

	Target	Indication	DPC	Phase I/II	Phase II	Phase III Potential registration
Developmental and epileptic encephalopathies AntiSensOligonucleotid	ND*	<p>Movement disorders are a diverse group of neurological conditions characterized by abnormal movements that significantly impact daily functioning. These disorders are often due to genetic mutations, neurodegenerative processes, or environmental factors affecting the brain's motor control pathways. Common examples include sustained muscle contractions and abnormal postures, as irregular, rapid movements difficult to control.</p> <p>Patients with rare movement disorders often face challenges in mobility, communication, and social interactions. The unpredictability of symptoms is often associated with emotional distress and a reduced quality of life.</p> <p>More details about ASO technology</p>				

Cardiometabolism & Venous Diseases

As data of January 2026

	Indication	DPC	Phase I/II	Phase II	Phase III Potential registration	Filing
Ivabradine (new formulation)	Heart Failure					
Dapagliflozin/Bisoprolol	Heart Failure					
Bisoprolol/Perindopril/Indapamide/Amlodipine	Hypertension					
Perindopril/Indapamide SR/Amlodipine	Hypertension					
Gliclazide/Metformin	T2 diabetes					
Bisoprolol/Perindopril/Amlodipine	Hypertension					
Dapagliflozin/Gliclazide	T2 diabetes					
+5 ON GOING FEASIBILITY PROJECTS						

