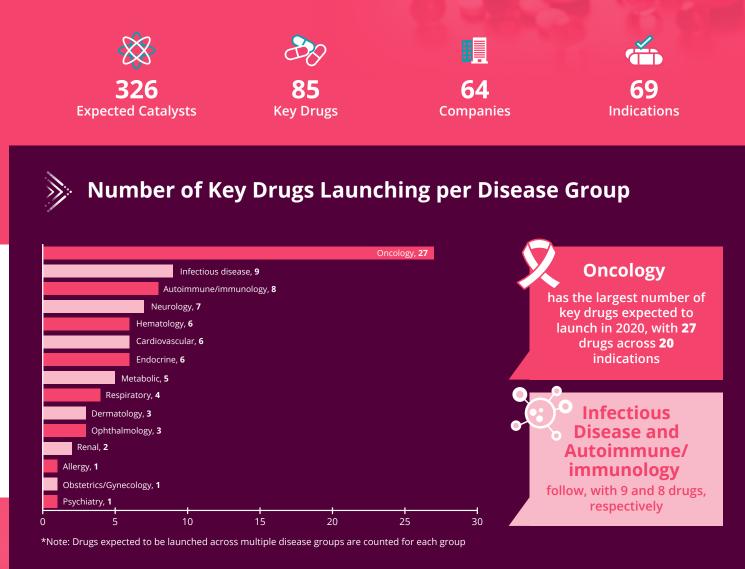
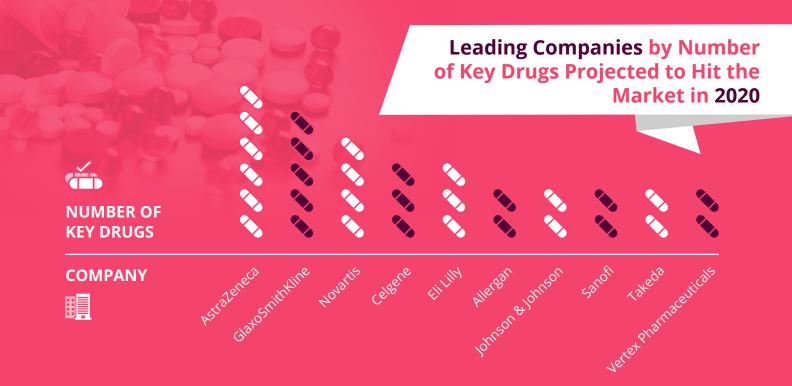
Informa Pharma Intelligence

## Key Potential Drug Launches in 2020

Take a longer-term look at key late-stage drugs projected to hit the market in 2020, including new drug classes, major changes to standards of care, and/or large market opportunities across the wide range of indications covered by Biomedtracker and Datamonitor Healthcare





## AstraZeneca is expected to submit filings for **six** drugs across **seven** indications:

Drug	Indication	Expected Date Range	Filing Type	Current Phase	Likelihood Of Approval
Brilinta	Atherosclerosis	Now-12/31/2019	Supplemental Filings for US, EU, Japan	Ш	52% (5% Above Avg.)
		01/01/2020-12/31/2020	Supplemental China Filing		
Farxiga	Congestive Heart Failure (CHF) and Cardiomyopathies	01/01/2020-12/31/2020	Supplemental Filings for US, Europe, Japan	III	48% (1% Above Avg.)
Roxadustat	Anemia Due to Chronic Renal Failure, Dialysis-Dependent	07/01/2019-12/31/2019	NDA Filing	III	73% (13% Above Avg.)
	Anemia Due to Chronic Renal failure, Dialysis-Independent	07/01/2019-12/31/2019	NDA Filing	III	74% (14% Above Avg.)
Selumetinib	Neurofibromatosis (NF)	07/01/2019-12/31/2019	NDA and MAA Filings	II	17% (Same As Avg.)
Calquence	Chronic Lymphocytic Leukemia (CLL)/Small Cell Lymphocytic Lymphoma (SLL) - NHL	Now-12/31/2019	Supplemental Filings for US and EU – High Risk	III	42% (7% Above Avg.)
	, ,,	01/01/2020-12/31/2020	Supplemental Filings for US and EU – 1L		
Lynparza	Pancreatic Cancer	07/01/2019-12/31/2019	Supplemental Filings for US, EU, Japan	III	45% (10% Above Avg.)

**Brilinta** could experience significant uptake on the market in this novel patient population where optimal long-term therapy with dual anti-platelet therapy is not established and use of aspirin alone is considered effective



**Farxiga** will be the first sodium-glucose cotransporter-2 inhibitor (SGLT-2i) to be approved by the FDA for heart failure with reduced ejection fraction, a significant unmet need affecting approximately 50% of chronic heart failure patients



**Lynparza** is set to become not only the first PARP inhibitor approved for pancreatic cancer, but also the first targeted therapy approved for pancreatic cancer since Tarceva was approved in 2005

## GlaxoSmithKline is expected to submit filings for five drugs across five indications:

Drug	Indication	Expected Date Range	Filing Type	Phase	Likelihood Of Approval
Cabotegravir LAP	HIV / AIDS	Now-06/30/2019	Filing for Approval (Canada)	NDA	94% (6% Above Avg.)
		Now-09/30/2019	MAA Filing - w/Rilpivirine		
Dostarlimab	Uterine (Endometrial) Cancer	07/01/2019-12/31/2019	BLA Filing	I	10% (4% Above Avg.)
GSK2857916	Multiple Myeloma (MM)	07/01/2019-12/31/2019	NDA Filing	II	14% (4% Above Avg.)
Zejula	Prostate Cancer	Now-12/31/2019	Supplemental Filings for US and EU	III	39% (4% Above Avg.)
Trelegy Ellipta	Asthma	10/01/2019-12/31/2019	Supplemental Filings for US and Europe	Ш	71% (3% Above Avg.)



**Cabotegravir**, in a fixed-dose combination with Johnson & Johnson's rilpivirine, is anticipated to be the first long-acting injectable to enter the HIV market, which could generate blockbuster sales by 2024 (if approved)



**GSK2857916**, an antibody-drug conjugate (ADC) directed towards B-cell maturation antigen (BCMA), is a unique drug in the multiple myeloma space where no ADCs or BCMA-directed therapies are currently approved

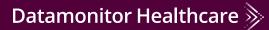


**Zejula** appears to be one of the first PARP inhibitors projected to gain approval for prostate cancer, competing with three other PARP inhibitors in Phase III development

Source: Biomedtracker, Datamonitor Healthcare May 2019







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