



## Prescription Pharmaceutical Products

PRODUCT	INDICATION <sup>1,2,4</sup>	TERRITORY <sup>3</sup>						PARTNER
		Canada	Brazil	Argentina	Colombia	Mexico	Others	
<b>Oncology/Hematology</b>								
<b>Tafasitamab</b>	Relapsed or refractory diffuse large B-cell lymphoma (DLBCL)		Approved	Submitted	Submitted	Submitted	Pre-registration	Incyte
<b>Pemigatinib</b>	Metastatic cholangiocarcinoma		Submitted	Submitted	Submitted	Submitted	Pre-registration	Incyte
<b>Akynzeo®</b>	Prevention of chemotherapy-induced acute and delayed nausea and vomiting	Q4-22	Q3-22	Q3-22				Helsinn
<b>Aloxi®</b>	Prevention of acute nausea and vomiting associated with moderately and highly emetogenic cancer chemotherapy	Q4-22						Helsinn
<b>Fostamatinib</b>	Treatment of chronic immune thrombocytopenia		Pre-registration	Pre-registration	Submitted	Submitted		Rigel
<b>Nerlynx®</b>	Extended adjuvant breast cancer and metastatic breast cancer	Q4-19						Puma
<b>Trelstar®</b>	Advanced prostate cancer	Q2-20						Debiopharm
<b>Vidaza®</b>	Myelodysplastic syndrome		Q2-10					Celgene (BMS)
<b>Abraxane®</b>	Metastatic pancreatic cancer		Q4-17					Celgene (BMS)
<b>Halaven®</b>	Metastatic breast cancer and soft tissue sarcoma		Q4-17	Q4-19	Q2-22		Marketed	Eisai
<b>Lenvima®</b>	Differentiated thyroid cancer and unresectable hepatocellular carcinoma		Q4-17		Q1-22		Marketed	Eisai
<b>Lenvima®</b>	Advanced renal cell cancer		Q4-17				Marketed	Eisai
<b>BGx</b>								
<b>Ladevina®</b>	Multiple myeloma; myelodysplastic syndrome			2011	Q3-19		Marketed	Own
<b>Ladevina®</b>	Mantle Cell Lymphoma; follicular lymphoma			2011			Marketed	Own
<b>Zyvalix®</b>	Metastatic prostate cancer			2014	Q2-18		Marketed	Own
<b>Karfib®</b>	Relapsed or refractory multiple myeloma			Q4-19	Submitted		Marketed	Own
<b>Leprid®</b>	Palliative treatment of advanced prostate cancer			2007				Own
<b>Rembre®</b>	Chronic myeloid leukemia			2013	Q1-22		Marketed	Own
<b>Palbocil®, Bapocil®</b>	Breast cancer			Q1-23	Submitted		Approved	Own
<b>Xetrane®</b>	Multiple myeloma			Q2-19	Submitted		Approved	Own
<b>Xetrane®</b>	AIDS-related Kaposi sarcoma			Q2-22				Own

<sup>1</sup> The products in "pre-registration" have not yet been submitted for regulatory review and products in "submitted" are currently under regulatory review. The indication for all products classified as "pre-registration" or "submitted" is the anticipated indication upon regulatory approval.

<sup>2</sup> Refer to the "Products" section below for further details on the indication.

<sup>3</sup> The products with an associated date are currently marketed by Knight in the respective territory. The information provided represents the date when the product was launched by Knight or when it was acquired or in-licensed by Knight if such products had existing sales.

<sup>4</sup> The products in "Approved" have been approved by regulatory authorities but not yet commercially launched.



PRODUCT	INDICATION <sup>1,2</sup>	TERRITORY <sup>3</sup>						PARTNER
		Canada	Brazil	Argentina	Colombia	Mexico	Others	
<b>Infectious Diseases</b>								
<b>Ambisome®</b>	Invasive fungal infection		1997					Gilead
<b>Cresemba®</b>	Invasive fungal infection		Q2-20	Q3-19	Q3-19	Q2-19	Marketed	Basilea
<b>Impavido®</b>	Leishmaniasis						Marketed	Own
<b>BGx</b>								
<b>Dolufevir®</b>	HIV infection			Q2-21				Own
<b>Other Specialty</b>								
<b>Exelon®</b>	Symptomatic treatment of mild to moderately severe dementia in people with Alzheimer's and Parkinson's disease	Q2-21	Q2-21	Q2-21	Q2-21	Q2-21	Marketed	Own
<b>Ibsrela®</b>	IBS-C	Q1-21						Ardelyx
<b>Salofalk®</b>	Ulcerative colitis			2007	Pre-2019		Marketed	Dr. Falk
<b>Ursofalk®</b>	Primary biliary cirrhosis			2007	Pre-2019		Marketed	Dr. Falk
<b>Imvexxy®</b>	Moderate-to-severe dyspareunia	Approved						TXMD
<b>Bijuva®</b>	Moderate-to-severe vasomotor symptoms due to menopause	Approved						TXMD
<b>BGx</b>								
<b>Fibroner®</b>	Idiopathic pulmonary fibrosis			2017			Marketed	Own
<b>Toliscriin® DPI</b>	Pseudomonas aeruginosa lung infection in patients with cystic fibrosis			2017			Marketed	Own
<b>Toliscriin® 1-2</b>	Severe acute or resistant chronic infections due to colistin sensitive strains of gram-negative pathogenic bacilli			2017			Marketed	Own
<b>Tobradosa Haler®</b>	Chronic lung infections due to Pseudomonas aeruginosa			2018			Marketed	Own

<sup>1</sup> The products in "pre-registration" have not yet been submitted for regulatory review and products in "submitted" are currently under regulatory review. The indication for all products classified as "pre-registration" or "submitted" is the anticipated indication upon regulatory approval.

<sup>2</sup> Refer to the "Products" section below for further details on the indication.

<sup>3</sup> Products with dates represent products currently marketed by Knight. The information provided represents the date at which the product was launched by Knight or date at which product with existing sales was acquired or in-licensed by Knight.

<sup>4</sup> The products in "Approved" have been approved by regulatory authorities but not yet commercially launched.



## Products Pipeline

PRODUCT	INDICATION OR THERAPEUTIC AREA <sup>1,2,4</sup>	TERRITORY <sup>3</sup>						EXPECTED LAUNCH YEAR
		Canada	Brazil	Argentina	Colombia	Mexico	Others	
<b>Oncology/Hematology</b>								
<b>Tafasitamab</b>	Relapsed or refractory diffuse large B-cell lymphoma (DLBCL)		Approved	Submitted	Submitted	Submitted	Pre-registration	2024 -2025
<b>Pemigatinib</b>	Metastatic cholangiocarcinoma		Submitted	Submitted	Submitted	Submitted	Pre-registration	2025-2026
<b>Fostamatinib</b>	Treatment of chronic immune thrombocytopenia		Pre-registration	Pre-registration	Submitted	Submitted		2025-2026
<b>Imvexxy®</b>	Moderate-to-severe dyspareunia	Approved						2024
<b>Bijuva®</b>	Moderate-to-severe vasomotor symptoms due to menopause	Approved						2024
<b>Palbocil®, Bapocil®</b>	Breast Cancer				Submitted		Approved	2025
<b>Xetrane®</b>	Multiple myeloma				Submitted		Approved	2025
<b>Karfib®</b>	Relapsed or refractory multiple myeloma				Submitted			2025
<b>Rembre®</b>	Chronic myeloid leukemia						Submitted	2024
<b>Undisclosed Molecule</b>	Oncology/ Hematology			Development				2025
<b>Undisclosed Molecule</b>	Oncology/ Hematology			Development				2025
<b>Undisclosed Molecule</b>	Oncology/ Hematology		Development		Development	Development		2026 - 2027
<b>Undisclosed Molecule</b>	Oncology/ Hematology		Development					2027
<b>Other Specialty</b>								
<b>Undisclosed Molecule</b>	Other Specialty		Development	Development	Submitted		Development	2025 - 2026
<b>Undisclosed Molecule</b>	Other Specialty		Development		Development	Development		2026 - 2027

<sup>1</sup> The products in "pre-registration" have not yet been submitted for regulatory review and products in "submitted" are currently under regulatory review. The indication for all products classified as "pre-registration" or "submitted" is the anticipated indication upon regulatory approval.

<sup>2</sup> Refer to the "Products" section below for further details on the indication.

<sup>3</sup> Products with dates represent products currently marketed by Knight. The information provided represents the date at which the product was launched by Knight or date at which product with existing sales was acquired or in-licensed by Knight.

<sup>4</sup> The products in "Approved" have been approved by regulatory authorities but not yet commercially launched.