

Q2 2021 Results

July 28, 2021

Forward Looking Statement and Non-GAAP Financial Information

This presentation contains statements about the Company's future plans and prospects that constitute forward-looking statements for purposes of the safe harbor provisions under the Private Securities Litigation Reform Act of 1995. Actual results may differ materially from those indicated as a result of various important factors, including those discussed in the Company's most recent annual report on Form 10-K and reports on Form 10-Q and Form 8-K. These documents are available on the SEC's website, on the Bristol-Myers Squibb website or from Bristol-Myers Squibb Investor Relations.

In addition, any forward-looking statements represent our estimates only as of the date hereof and should not be relied upon as representing our estimates as of any subsequent date. While we may elect to update forward-looking statements at some point in the future, we specifically disclaim any obligation to do so, even if our estimates change.

This presentation includes certain non-generally accepted accounting principles (GAAP) financial measures that we use to describe our company's performance. The non-GAAP information presented provides investors with additional useful information but should not be considered in isolation or as substitutes for the related GAAP measures. Moreover, other companies may define non-GAAP measures differently, which limits the usefulness of these measures for comparisons with such other companies. We encourage investors to review our financial statements and publicly-filed reports in their entirety and not to rely on any single financial measure. An explanation of these non-GAAP financial measures and a reconciliation to the most directly comparable GAAP financial measure are available on our website at bms.com/investors.

Also note that a reconciliation of certain forward-looking statements, however, is not provided due to no reasonably accessible or reliable comparable GAAP measures for such statements and the inherent difficulty in forecasting and quantifying such statements that are necessary for such reconciliation.

Q2 2021 Results



Giovanni Caforio

Board Chair and
Chief Executive Officer

Q2 2021 Performance

Operational Performance

Strong commercial execution

- Sales of \$11.7B in Q2; +16% YoY, +13% ex-FX
- Strong momentum with launches for I-O and new product portfolio

Pipeline Execution

Significant milestones

- **Solid Tumors:** Opdivo U.S. approval in adj. EC/GEJ & positive CHMP opinion; Opdivo in adj. MIBC - PDUFA Sep 3, 2021
- **Hematology:** Positive data for Breyanzi in 2L TE LBCL; Abecma positive CHMP opinion in 4L+ MM; iberdomide + dex in 4L+ MM data in-house
- **Immunology:** Zeposia U.S. approval in moderate-to-severe UC
- **CV:** Milvexian (FXIa inhibitor) Ph 2 TKR data in-house

Financial Strength

Strong financial results and outlook

- Continued revenue and EPS growth
- Reaffirm 2021 Revenue and Non-GAAP EPS guidance
- Balance sheet strength and strong cash flow generation; debt repayments of ~\$5.7B & executed share repurchases of ~\$3B YTD

Business Development

- Licensed anti-TIGIT bispecific antibody program w/ Agenus & FRα ADC collab w/ Eisai

Execution Scorecard

Financial Expectations

- 2020-2025:
 - Low to mid-single digit revenue CAGR*
 - Low double-digit revenue CAGR for Continuing business*
- Operating margins low to mid 40%^{**}
- ~\$3B of synergies by end of 2022
- \$45B - \$50B of free-cash flow 2021-2023^{**}

On track based on 2021 guidance

2021 Key Milestones

Opdivo (+/- Yervoy)	U.S./EU expected approvals: 1L RCC (9ER) ✓, 1L GC (649, O+Chemo) ✓ adj Eso (577) ✓ adj MIBC (274) ✓
	1L Esophageal (CM-648) ✓
	Opdivo return to annual growth ✓
Relatlimab	1L Melanoma w/ Opdivo Ph3 ✓
Breyanzi	3L+ DLBCL U.S. ✓ / EU approval
	2L TE ✓ and TNE DLBCL
	3L+ CLL
Abecma	4L+ MM U.S. ¹ ✓ / EU approval
Iberdomide + dex	4L+ MM Ph 1b/2a ✓
Deucravacitinib	PsO (2 nd study) Ph3 ✓ & U.S. filing
	UC Ph2 (POC)
Zeposia	UC U.S. ✓ / EU approval
Cendakimab	Initiation of Ph3 ✓
Factor XIa inh.	Total Knee Replacement VTEp Ph2 (POC) ✓
Mavacamten	oHCM U.S. filing ✓ & approval ²

2022/2023 Key Milestones

Opdivo (+/- Yervoy)	Metastatic 1L HCC (CM-9DW)
	Adjuvant Neo-adj Lung EFS (CM-816) Peri-adj Lung (CM-77T)
Bempeg	1L melanoma ³ & 1L renal
Breyanzi	3L+ Follicular lymphoma
Abecma	3L+ MM (KarMMa-3) Ph3
	2L+ MM (KarMMa-2) POC
CC-92480	4L+ MM Ph1/2
CC-93269 (TCE)	Initiation of pivotal trial
Deucravacitinib	PsO U.S./EU approval
	CD & Lupus Ph2 (POC)
Zeposia	CD Ph3
Factor XIa inh.	Secondary Stroke Prevention Ph2 (POC)
Reblozyl	1L MDS (ESA naïve) COMMANDS Ph3
Ph 1/2 Pipeline	>20 POC decisions

To be expanded to include regulatory milestones pending future registrational successes

Continued portfolio renewal into 2H of decade

Launch Portfolio Best / first-in-class assets

Reblozyl[®]
(luspatercept-aamt)
for injection 25mg • 75mg

ONUREG[™]
(azacitidine) tablets
300mg • 200mg

Breyanzi[®]
(isocabtagene maraleucel) SUSPENSION
FOR IV INFUSION

Abecma[™]
(idecabtagene vicleucel) SUSPENSION
FOR IV INFUSION

ZEPOSIA[®]
(ozanimod) 0.92 mg
capsules

Anticipated Launches

deucravacitinib

mavacamten

relatlimab

\$20B-\$25B
NRA revenue
potential
in 2029*

+ relatlimab

Future growth opportunities

- Additional mid-to late-stage pipeline, including:

iberdomide

milvexian

- Diverse early-stage pipeline
- Balance Sheet strength for additional BD opportunities

Q2 2021 Results



David Elkins

Chief Financial Officer

Strong performance in key franchises

Q2 Sales	Net Sales \$ in Billions	Vs. Prior Year
 Revlimid [®] (lenalidomide) capsules	\$3.2	▲ 11%
 Eliquis [™] apixaban	\$2.8	▲ 29%
 OPDIVO [™] (nivolumab) <small>INJECTION FOR INTRAVENOUS USE 50 mg/mL</small>	\$1.9	▲ 16%
 Pomalyst [™] (pomalidomide) capsules	\$0.9	▲ 15%
 ORENCIA [®] (abatacept)	\$0.8	▲ 9%
 SPRYCEL [™] dasatinib 100 mg tablets	\$0.5	▲ 6%
 YERVOY [™] (ipilimumab) <small>injection for intravenous infusion</small>	\$0.5	▲ 38%
 Abraxane [®]	\$0.3	▼ 4%

Q2 2021 Total Sales: \$11.7B, up 16% & 13% ex-FX

YTD Sales	Net Sales \$ in Billions	Vs. Prior Year
 Revlimid [®] (lenalidomide) capsules	\$6.1	▲ 6%
 Eliquis [™] apixaban	\$5.7	▲ 18%
 OPDIVO [™] (nivolumab) <small>INJECTION FOR INTRAVENOUS USE 50 mg/mL</small>	\$3.6	▲ 6%
 Pomalyst [™] (pomalidomide) capsules	\$1.6	▲ 12%
 ORENCIA [®] (abatacept)	\$1.6	▲ 7%
 SPRYCEL [™] dasatinib 100 mg tablets	\$1.0	▼ 2%
 YERVOY [™] (ipilimumab) <small>injection for intravenous infusion</small>	\$1.0	▲ 26%
 Abraxane [®]	\$0.6	- 0%

YTD 2021 Total Sales: \$22.8B, up 9% & 7% ex-FX

Q2 2021 Eliquis performance



Global net sales up 29% in Q2; 18% YTD

US: Continued significant demand growth

- Q2 sales up 26% vs PY; 16% YTD
 - Continued strong underlying demand
 - ~14% TRx growth

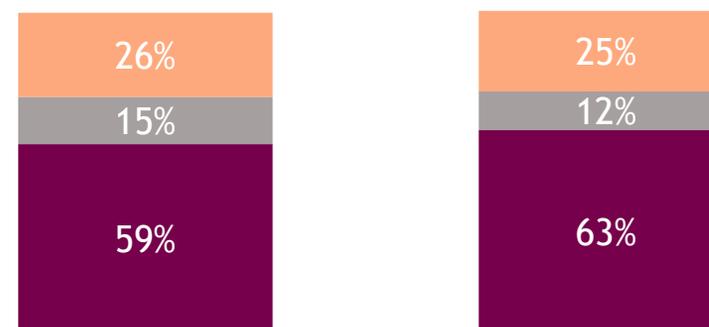
International: Strong demand growth

- Continues to be #1 OAC in key markets with future growth

Significant future growth opportunity

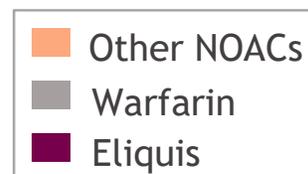
- Expect to continue to grow share within an expanding class

NBRx Share - US

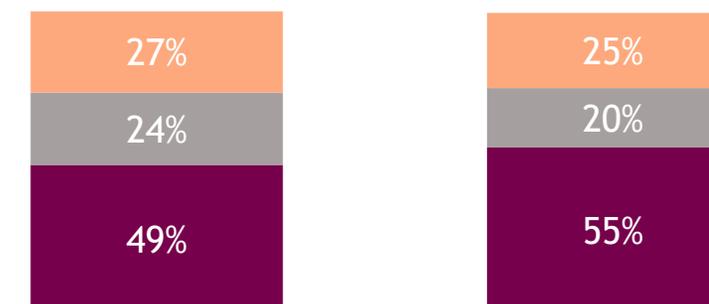


Q2 2020

Q2 2021



TRx Share - US



Q2 2020

Q2 2021

Q2 2021 Opdivo performance



Global net sales up 16% in Q2

U.S.

- Return to growth (+13% vs. PY)
- 1L lung* shares in low double-digits
- Leadership position in 1L renal enabled by O+Cabo
- Strong early launches in Upper GI

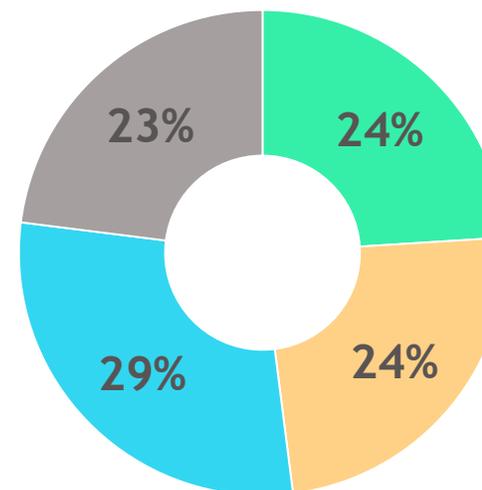
International

- Momentum from new launches, FX, improved COVID dynamics vs. PY

Near term growth drivers

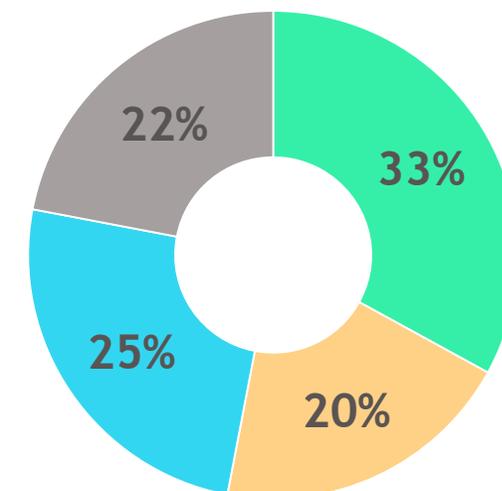
- Momentum from recent launches
- Potential next launches: adjuvant bladder (CM-274), 1L esophageal (CM-648)

Approx. U.S. Sales Mix



- NSCLC
- RCC
- Melanoma
- All others

Approx. Ex-U.S. Sales Mix



Note: percentages approximate based on tumor ranges

Q2 2021 Multiple Myeloma performance

Global sales growth of 11%

- US sales growth of 6%
 - Increased use of triplet regimens & longer treatment duration
 - Recovery to pre-COVID levels in NRx
- International sales growth of 24%
 - Demand from triplets & maintenance use
 - Improved COVID dynamics & FX

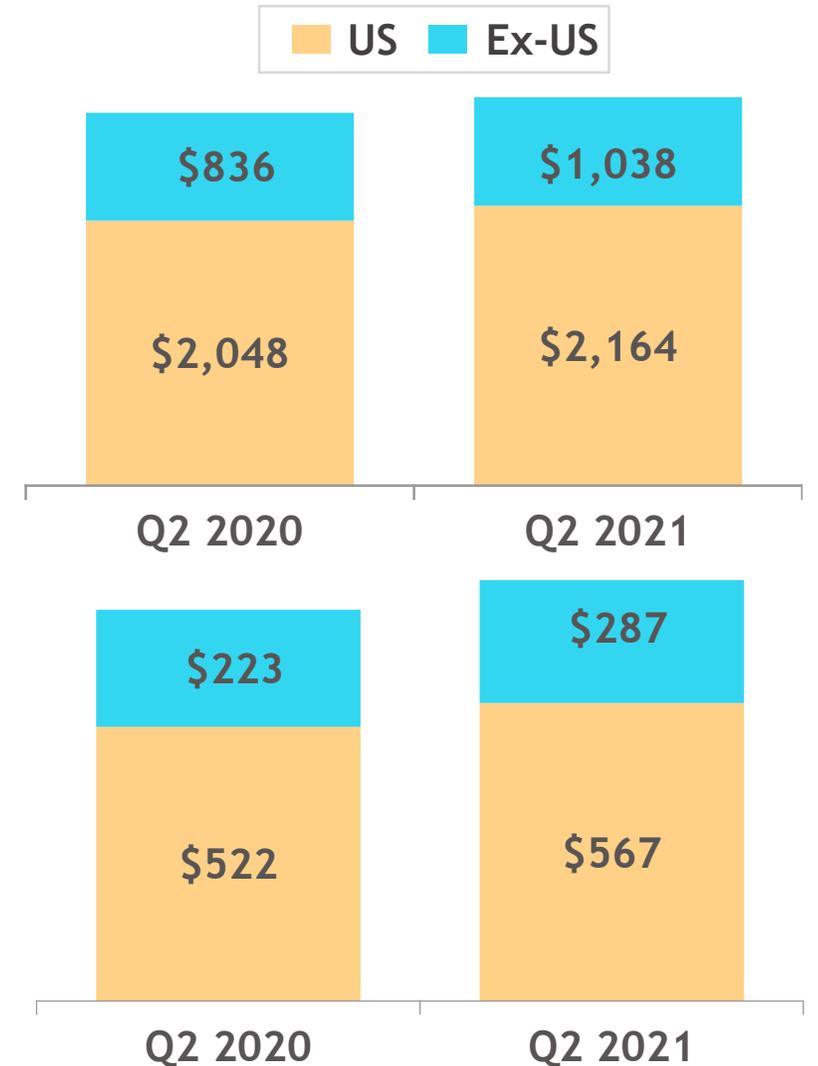


Global sales growth of 15%

- Demand growth from new triplet regimens and use in earlier lines



Global Net Sales



Advancing new product launches



- Continued transition from initial bolus to underlying incidence demand
- Remain focused on new patients earlier in their treatment journey & dose optimization
- Expansion in global markets

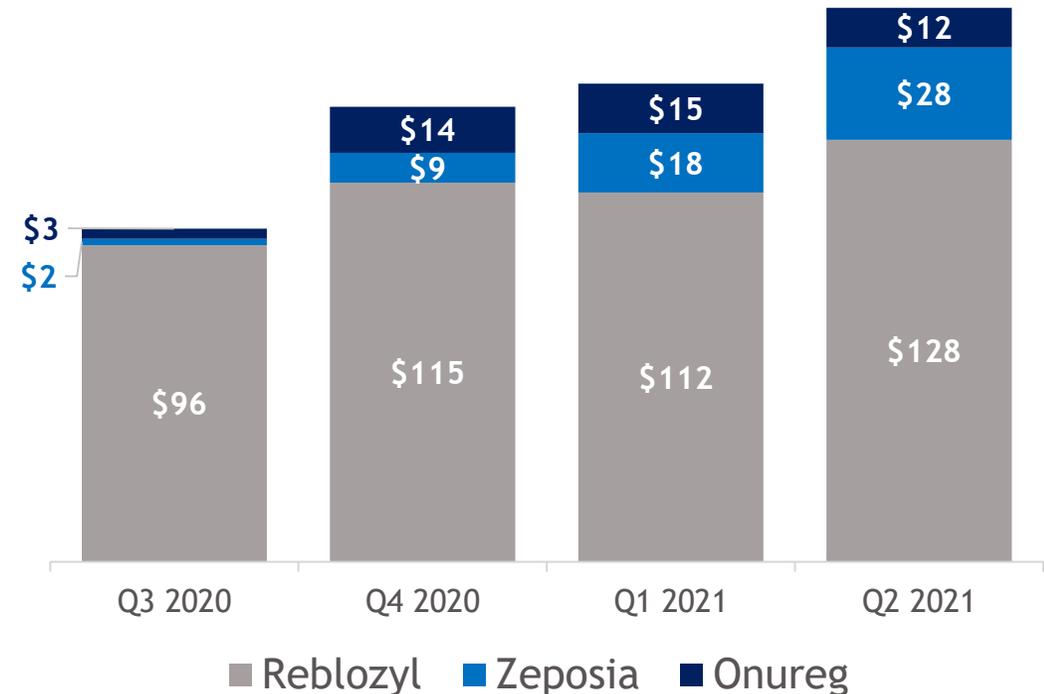


- #1 S1P modulator in written Rx in MS
- Acceleration of Rx to commercial demand
- UC approved in the U.S. - positive physician receptivity
- MAA under review



- Strong demand growth sequentially
- Establishing profile in 1L response maintenance setting in AML
- Approved by EC June 2021

Q2 2021 Global Net Sales



Cell Therapy Performance

Two differentiated CAR T products,
now in the marketplace



Best-in-class CD19 CAR T
Global net sales: \$17M

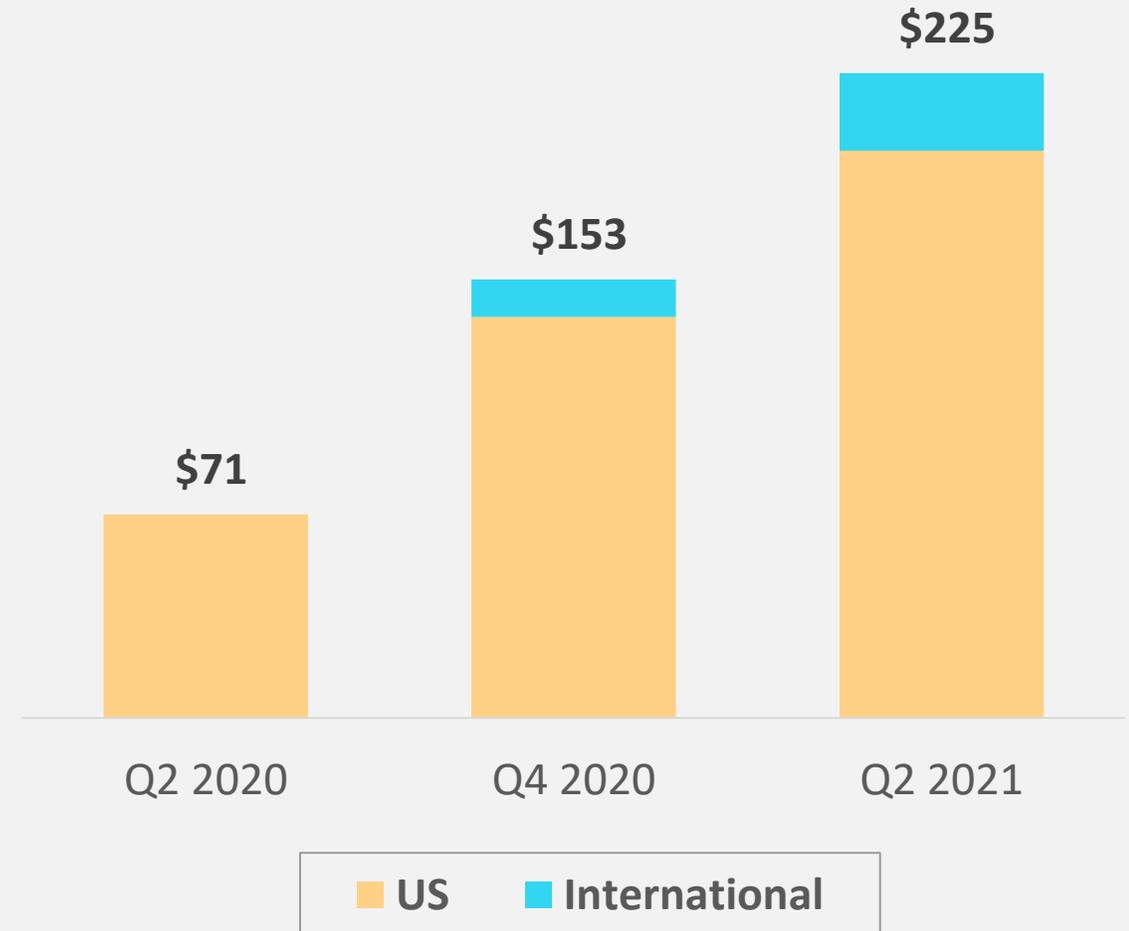


First-in-class BCMA CAR T
Global net sales: \$24M

-
- Rapid site activation (>65 sites)
 - Strong demand and physician awareness

New Launch Sales Performance

- Contributed \$225M in Q2'21
- Approaching ~\$1B annual run rate
- Strong outlook for future growth



Q2 2021 Financial Performance

<i>\$ in billions, except EPS</i>	US GAAP		Non-GAAP	
	Q2 2021	Q2 2020	Q2 2021	Q2 2020
Total Revenues, net	11.7	10.1	11.7	10.1
Gross Margin %	79.0%	73.4%	79.8%	80.5%
MS&A	1.9	1.6	1.9	1.6
R&D	3.3	2.5	2.3	2.2
Effective Tax Rate	31.7%	104.9%	16.9%	13.8%
Diluted EPS	0.47	(0.04)	1.93	1.63
Diluted Shares Outstanding <i>(# in millions)</i>	2,252	2,263	2,252	2,297

Significant financial flexibility to support a balanced approach to capital allocation

\$B	Q2 2021
Total Cash**	\$13.1B
Total Debt	\$45.2B
Net Debt Position	\$32.0B

Future innovation through business development

- Strategically Aligned
 - Scientifically Sound
 - Financially Attractive
-

Committed to reducing debt

- ~\$5.7B in debt reduction YTD
 - Maintain strong investment-grade credit ratings
-

Returning capital to shareholders

- Continued dividend growth*
- \$3B - \$4B total share repurchase planned; executed ~\$3B YTD & remain opportunistic in 2H

2021 Guidance Details

	GAAP	Non-GAAP
Net Sales	High single-digit increase	High single-digit increase
Gross Margin %	~79%	~80%
MS&A Expense	In line with 2020	Low single-digit increase
R&D Expense	Low single-digit decrease	Mid single-digit increase
Tax Rate	~23%	~16%
Diluted EPS	\$2.77-\$2.97	\$7.35 - \$7.55

Non-GAAP EPS reaffirmed

Q&A



Giovanni Caforio, M.D.
Board Chair,
Chief Executive Officer



Chris Boerner, Ph.D.
Executive VP,
Chief Commercialization Officer



David Elkins
Executive VP,
Chief Financial Officer



Samit Hirawat, M.D.
Executive VP,
Chief Medical Officer,
Global Drug Development

2021 News Flow

Asset	Timing
Opdivo Approval in 1L renal (CM-9ER)	Approved in U.S. & EU ✓
Opdivo Approval in 1L gastric (CM-649)	Approved in U.S. ✓
Opdivo Approval in adjuvant esophageal (CM-577)	Approved ✓ MAA under review
Opdivo Approval in adjuvant MIBC (CM-274)	PDUFA date September 3, 2021
Opdivo CM-648 in 1L esophageal	Positive topline ✓ April 2021
Relatlimab + Opdivo vs Opdivo mono 1L Melanoma CA224-047	Positive topline ✓ March 2021
Breyanzi Approval in 3L+ LBCL	Approved in U.S. ✓ MAA under review
Breyanzi 2L DLBCL	Positive topline ✓ June 2021
3L+ CLL	2H 2021

Asset	Timing
Abecma Approval in 4L+ MM	Approved in U.S. ¹ ✓ MAA under review
Zeposia Approval in UC	Approved in U.S. ✓ MAA under review
iberdomide + dex	Data in-house ² ✓
deucravacitinib Psoriasis Ph3 POETYK PSO-2 (IM011-047)	Positive topline ✓ Feb 2021
deucravacitinib Ph2 POC in UC	2H 2021
milvexian (FX1a inhib) POC in VTEp for total knee replacement	Data in-house ² ✓
mavacamten Obstructive HCM	PDUFA date January 28, 2022 ✓

Active Clinical Development Portfolio

Phase 1

Phase 2

Phase 3

Marketed

Oncology

AHR Antagonist (Ikena)**	Anti-NKG2A	Anti-TIM3	motolimod
Anti-CCR8	Anti-OX40	AR LDD	NLRP3 Agonist
Anti-CTLA-4 NF-Probody	Anti-SIRPα*	CD3xPSCA (GEMoAB)**	STING Agonist
Anti-IL8	TIGIT Bispecific (Agenus)**	IL-12 Fc	TGFβ Inhibitor

Anti-CTLA-4 NF	BET Inhibitor* (CC-90010)
Anti-CTLA-4 Probody	
Anti-Fucosyl GM1	
Anti-TIGIT	
	FRα ADC
	LSD1 Inhibitor*

bempegal- desleukin
linrodostat
subcutaneous nivolumab
relatlimab*



Hematology

A/I CELMoD (CC-99282)	BCMA NKE	ROR1 CAR T	CD22 ADC (TriPhase)**
CK1α CELMoD	BCMA TCE	BCMA NEX T	CD3xCD33 (GEMoAB)**
GSPT1 CELMoD (CC-90009)	BCMA CAR T (bb21217)	CD19 NEX T	CD33 NKE
BCMA ADC	GPRC5D CAR T	BET Inhibitor* (CC-95775)	CD47xCD20

A/I CELMoD (CC-92480)
BET Inhibitor (BMS-986158)
iberdomide



Cardiovascular

FXIa Inhibitor	FPR-2 Agonist	Cardiac Myosin Inhibitor	ROMK Inhibitor
----------------	---------------	-----------------------------	----------------

danicamtiv
FA-Relaxin

milvexian (FXIa Inhibitor)

mavacamten



Immunology

Anti-CD40	IL2 Mutein	MK2 Inhibitor	TLR 7/8 Inhibitor	TYK2 Inhibitor (Nimbus)**
IL2-CD25	Imm. Tolerance (Anokion)**	S1PR1 Modulator	TYK2 Inhibitor	

branebrutinib
iberdomide

deucravacitinib
cendakimab



Fibrosis

NME 1

HSP47	LPA ₁ Antagonist
JNK Inhibitor	pegbelfermin

Neuroscience

Anti-Tau (Prothena)**	BTK Inhibitor	FAAH/MGLL Dual Inhibitor
--------------------------	---------------	-----------------------------

COVID-19

SARS-CoV-2 mAb Duo

Data as of July 21, 2021