UNITED STATES
SECURITIES AND EXCHANGE COMMISSION

WASHINGTON, D.C. 20549

FORM 8-K

CURRENT REPORT
Pursuant to Section 13 or 15(d)
of the Securities Exchange Act of 1934

Date of Report (Date of Earliest Event Reported): **November 13, 2017**

BEIGENE, LTD.

(Exact name of registrant as specified in its charter)

Cayman Islands	**001-37686**	**98-1209416**
(State or other jurisdiction of incorporation)	(Commission File Number)	(I.R.S. Employer Identification No.)

c/o Mourant Ozannes Corporate Services (Cayman) Limited
94 Solaris Avenue, Camana Bay
Grand Cayman KY1-1108
Cayman Islands
(Address of principal executive offices) (Zip Code)

+1 (345) 949 4123
(Registrant's telephone number, including area code)

Not Applicable
(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

☐ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

☐ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

☐ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

☐ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 or Rule 12b-2 of the Securities Exchange Act of 1934.

Emerging growth company ☒

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☒

Item 2.02 Results of Operations and Financial Condition.

On November 13, 2017, BeiGene, Ltd. (the "Company") announced its financial results for the three and nine months ended September 30, 2017. A copy of the press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K.

The information set forth under this Item 2.02 in this Current Report on Form 8-K and Exhibit 99.1 attached hereto is intended to be furnished and shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such filing.

Item 8.01 Other Events.

On November 13, 2017, the Company issued a press release announcing that it has initiated two new global pivotal clinical trials of BGB-3111, an investigational Bruton's Tyrosine Kinase inhibitor: a Phase 3 trial of BGB-3111 in previously untreated patients with chronic lymphocytic leukemia / small lymphocytic lymphoma and a pivotal Phase 2 trial of BGB-3111 in combination with GAZYVA® (obinutuzumab) in patients with relapsed or refractory follicular lymphoma. The Company also announced completion of patient enrollment in a pivotal trial of BGB-3111 in China in patients with mantle cell lymphoma. The full text of the Company's press release is filed as Exhibit 99.2 to this Current Report on Form 8-K and is incorporated herein by reference.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits .

Exhibit No.	Description
99.1	Press Release titled "BeiGene Reports Third Quarter 2017 Financial Results" issued on November 13, 2017, furnished herewith
99.2	Press Release titled "BeiGene Expands Global Pivotal Program for BTK Inhibitor BGB-3111" issued on November 13, 2017

Exhibit Index

Exhibit No.	Description
99.1	Press Release titled "BeiGene Reports Third Quarter 2017 Financial Results" issued on November 13, 2017, furnished herewith
99.2	Press Release titled "BeiGene Expands Global Pivotal Program for BTK Inhibitor BGB-3111" issued on November 13, 2017

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Date: November 13, 2017

BEIGENE, LTD.

By: /s/ Scott A. Samuels
Name: Scott A. Samuels
Title: Senior Vice President, General Counsel

BeiGene

BeiGene Reports Third Quarter 2017 Financial Results

CAMBRIDGE, Mass. and BEIJING, China, November 13, 2017 (GLOBE NEWSWIRE) -- BeiGene, Ltd. (NASDAQ: BGNE), a commercial-stage biopharmaceutical company focused on developing and commercializing innovative molecularly targeted and immuno-oncology drugs for the treatment of cancer, today reported business highlights and financial results for the third quarter and first nine months of 2017.

"BeiGene has achieved several important milestones since the beginning of the third quarter. Our strategic collaboration with Celgene Corporation has transformed us into a commercial-stage company in China and is expected to enhance the potential of our investigational PD-1 inhibitor, BGB-A317. We have also made important clinical progress with the expansion of the global pivotal program for our BTK inhibitor BGB-3111, the initiation of the first Phase 3 study for BGB-A317 in China, and the completion of patient enrollment of our first China pivotal trials for both BGB-3111 and BGB-A317. In addition, we strengthened our balance sheet with the upfront payment and investment from Celgene as well as our public offering in August," said John V. Oyler, Founder, Chief Executive Officer, and Chairman of BeiGene.

"Looking ahead, we plan to present additional data on BGB-3111 at the 59 th American Society of Hematology Annual Meeting in December. We also look forward to initiating additional pivotal trials for our portfolio assets in the coming months," commented Mr. Oyler.

Third Quarter 2017 and Recent Business Highlights

Clinical Programs:

BGB-3111 , *an investigational small molecule inhibitor of Bruton's tyrosine kinase (BTK)*

- Completed enrollment in the pivotal Phase 2 trial in China of BGB-3111 in relapsed/refractory mantle cell lymphoma;
- Initiated the following trials:
    - Global Phase 3 trial of BGB-3111 compared to bendamustine and rituximab in treatment-naïve chronic lymphocytic leukemia /

small lymphocytic lymphoma patients;

- Global pivotal Phase 2 trial of BGB-3111 in combination with GAZYVA® (obinutuzumab) in relapsed or refractory follicular lymphoma patients; and
- Pivotal Phase 2 trial in China of BGB-3111 in Waldenström's macroglobulinemia.

BGB-A317 , *an investigational humanized monoclonal antibody against the immune checkpoint receptor PD-1*

- Presented data from patients with gastric cancer, esophageal cancer, head and neck squamous cell carcinoma, and ovarian cancer enrolled in the global Phase 1 trial of BGB-A317 in patients with advanced solid tumors at the European Society for Medical Oncology (ESMO) 2017 Congress;
- Presented preliminary Phase 1 data on BGB-A317 in Chinese patients with advanced tumors at the 20 th Annual Meeting of the Chinese Society of Clinical Oncology;
- Completed enrollment in the pivotal Phase 2 trial of BGB-A317 in China in relapsed/refractory classical Hodgkin's lymphoma;
- Completed enrollment in the global Phase 1a/1b trial of BGB-A317 in advanced tumors with a total of over 450 patients;
- Initiated the following trials:
  - Phase 3 trial in China of BGB-A317 as a second- or third-line treatment for patients with advanced lung cancer;
  - Pivotal Phase 2 trial in China of BGB-A317 in previously treated, PD-L1-positive, locally advanced or metastatic urothelial cancer;
  - Phase 2 trial in China of BGB-A317 in combination with chemotherapy as a first-line treatment for patients with advanced lung cancer; and
  - Phase 2 trial in China of BGB-A317 in combination with chemotherapy as a first-line treatment for patients with locally

advanced or metastatic esophageal, gastric, or gastroesophageal junction carcinoma.

BGB-290 , *an investigational small molecule PARP inhibitor*

- Presented updated data from the global Phase 1 trial of BGB-290 in patients with advanced solid tumors at the ESMO 2017 Congress;
- Initiated the following trials:
  - Global Phase 1 trial of BGB-290 in combination with temozolomide in locally advanced or metastatic solid tumors; and
  - Global Phase 1b/2 trial of BGB-290 in combination with radiation therapy and/or temozolomide in glioblastoma.

Corporate Development:

- Closed our global strategic collaboration with Celgene Corporation.

Expected Upcoming Milestones

BGB-3111 (BTK Inhibitor)

- Present additional Phase 1 data for BGB-3111 in non-Hodgkin's lymphoma, updated Phase 1 data for the combination of BGB-3111 and Gazyva® (obinutuzumab), and initial Phase 1 data for the combination of BGB-3111 and BGB-A317 at the 59th American Society of Hematology (ASH) Annual Meeting in Atlanta, GA, December 9-12, 2017.

BGB-A317 (PD-1 Antibody)

- Present initial Phase 1 data for the combination of BGB-3111 and BGB-A317 at the 59th ASH Annual Meeting in Atlanta, GA, December 9-12, 2017; and
- Initiate Phase 3 trials of BGB-A317 in China in the fourth quarter of 2017 or the first quarter of 2018.

BGB-290 (PARP Inhibitor)

- Initiate a pivotal trial in China in the fourth quarter of 2017.

Third Quarter 2017 Financial Results

Cash, Cash Equivalents, and Short-Term Investments were $757.44 million as of September 30, 2017, compared to $407.43 million as of June 30, 2017 and $368.17 million as of December 31, 2016. The increase in the quarter was primarily attributable to the Celgene strategic collaboration. Funding from Celgene includes upfront licensing fees of $263.00 million, $92.05 million of which was received as of September 30, 2017 and the rest of which is payable in the fourth quarter of 2017, and an equity investment of $150.00 million. In addition, net proceeds from our August 2017 follow-on public offering contributed $188.52 million, after deducting underwriting discounts and offering-related expenses. These were partially offset by cash used in operating activities and for capital expenditures during the three months ended September 30, 2017.

The Company consolidates the BeiGene Biologics joint venture in its consolidated financial statements. As of September 30, 2017, cash, cash equivalents and short-term investments included $141.64 million of cash held by BeiGene Biologics.

Cash provided by operations for the three months ended September 30, 2017 was $6.60 million, compared to a use of cash of $24.28 million for the same period in 2016. The increase in cash flow from operating activities was primarily attributable to upfront licensing fees of the Celgene collaboration, which offset increased research and development (R&D) and selling, general and administrative (SG&A) expenses in the period. Capital expenditures for the quarter ended September 30, 2017 were $18.79 million, including $13.94 million attributable to BeiGene Biologics, compared to $6.68 million for the same period in 2016. The increase was primarily attributable to increased investment in our manufacturing facilities in Guangzhou and Suzhou.

Revenue for the three months ended September 30, 2017 was $220.21 million, compared to nil in the same period in 2016, and was comprised of $8.82 million of net product revenue and $211.39 million of collaboration revenue. The product revenues represent net product sales of Abraxane and

Revlimid in China following the effective date of the Celgene transaction, August 31, 2017. The collaboration revenue recognized in the period relates to the upfront fee allocated to the BGB-A317 license element of the arrangement.

R&D Expenses for the three months ended September 30, 2017 were $87.66 million, compared to $30.11 million in the same period in 2016. The increase was primarily attributable to increased spending on clinical activities and manufacturing for BGB-3111, BGB-A317, and BGB-290 due to expansion of ongoing clinical programs and increased employee compensation-related expenses as a result of increased headcount to support a broader clinical program. R&D-associated share-based compensation expense was $10.38 million for the three months ended September 30, 2017, compared to $2.14 million for the same period in 2016, primarily due to increased headcount, higher share price, and increased expense attributable to non-employee consultant awards.

SG&A Expenses for the three months ended September 30, 2017 were $15.64 million compared to $4.72 million in the same period in 2016. The increase was primarily attributable to increased employee compensation-related expenses as a result of increased headcount, including personnel costs for the employees transferred from Celgene China and higher share price as well as higher professional service fees, including legal, finance and accounting fees related to the Celgene transaction, patent prosecution activities and costs to support our growing operations. In addition, SG&A-associated share-based compensation expense was $2.95 million for the three months ended September 30, 2017, compared to $0.64 million for the same period in 2016.

Net Income attributable to BeiGene, Ltd. for the three months ended September 30, 2017 was $117.39 million compared to a net loss of $35.49 million in the same period in 2016.

Financial Summary

Select Consolidated Balance Sheet Data (U.S. GAAP)
(Amounts in thousands of U.S. Dollars)

	September 30, 2017 (unaudited)	December 31, 2016 (audited)
Cash, cash equivalents and short-term investments	$ 757,435	$
Unbilled receivable	170,950	—
Prepaid expenses and other current assets	33,945	
Property and equipment, net	55,322	
Goodwill and other intangible assets	9,421	—
Total assets	1,049,059	405,813
Accounts payable	35,168	
Accrued expenses and other payables	46,991	
Deferred revenue	38,609	—
Bank loan	18,036	
Shareholder loan	140,311	—
Noncontrolling interest	14,349	—
Total equity	$ 764,489	$

Consolidated Statements of Operations (U.S. GAAP)
(Amounts in thousands of U.S. Dollars, except for number of American Depositary Shares (ADSs) and per ADS data) (unaudited)

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2017	2016	2017	2016
Revenue				
Product revenue, net	$ 8,822	$ —	$ 8,822	$ —
Collaboration revenue	211,391	—	211,391	1,070
Total revenues	220,213	—	220,213	1,070
Expenses:				
Cost of sales – products	(1,944)	—	(1,944)	—
Research and development	(87,660)	(30,106)	(177,678)	(69,100)
Selling, general and administrative	(15,641)	(4,722)	(35,187)	(11,760)
Amortization of intangible assets	(63)	—	(63)	—
Total expenses	(105,308)	(34,828)	(214,872)	(80,860)
Income (loss) from operations	114,905	(34,828)	5,341	(79,790)

Interest (expense) income, net	(1,785)	(75)	(3,581)	336
Changes in fair value of financial instruments	—	—	—	(1,514)
(Loss) gain on sale of available-for-sale securities	—	(137)	10	(1,077)
Other income (expense), net	1,103	(327)	1,531	732
Income (loss) before income taxes	114,223	(35,367)	3,301	(81,313)
Income tax benefit (expense)	3,061	(127)	2,680	(306)
Net income (loss)	$ 117,284	$ (35,494)	$ 5,981	(81,619)
Less: Net loss attributable to noncontrolling interest	(102)	—	(237)	—
Net income (loss) attributable to BeiGene, Ltd.	$ 117,386	$ (35,494)	$ 6,218	(81,619)
Net income (loss) attributable to common shareholders per ADS, basic	$ 2.79	$ (1.08)	$ 0.15	(2.77)
Net income (loss) attributable to common shareholders per ADS, diluted	$ 2.54	$ (1.08)	$ 0.14	(2.77)
Weighted-average number of ADS outstanding - basic	42,118,973	32,933,655	40,563,845	29,497,875
Weighted-average number of ADS outstanding - diluted	46,200,975	32,933,655	43,172,139	29,497,875

Consolidated Statements of Comprehensive Income (Loss) (U.S. GAAP)

(Amounts in thousands of U.S. Dollars) (unaudited)

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2017	2016	2017	2016
Net income (loss)	$ 117,284	$ (35,494)	$ 5,981	$ (81,619)
Other comprehensive income (loss), net of tax of nil:				
Foreign currency translation adjustments	341	377	985	(13)
Unrealized holding gain, net	51	121	58	857
Comprehensive income (loss)	117,676	(34,996)	7,024	(80,775)
Less: Comprehensive loss attributable to noncontrolling interests	(70)	—	(178)	—
Comprehensive income (loss) attributable to BeiGene, Ltd.	$ 117,746	$ (34,996)	$ 7,202	$ (80,775)

About BeiGene

BeiGene is a global, commercial-stage, research-based biotechnology company focused on molecularly targeted and immuno-oncology cancer therapeutics. With a team of over 700 employees in China, the United States, and Australia, BeiGene is advancing a pipeline consisting of novel oral small molecules and monoclonal antibodies for cancer. BeiGene is also working to create combination solutions aimed to have both a meaningful and lasting impact on cancer patients. BeiGene markets ABRAXANE® (nanoparticle albumin–bound paclitaxel), REVLIMID® (lenalidomide), and VIDAZA® (azacitidine) in China under a license from Celgene Corporation. [i]

Forward-Looking Statements

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995 and other federal securities laws, including statements regarding BeiGene's financial condition; results of operations and business outlook; the momentum of its business, as well as the advancement of, and anticipated clinical development and regulatory milestones and plans related to BeiGene's drug candidates and clinical trials, including commencing new trials and providing data readouts and updates for its drug candidates. Actual results may differ materially from those indicated in the forward-looking statements as a result of various important factors, including risks related to BeiGene's ability to demonstrate the efficacy and safety of its drug candidates; the clinical results for its drug candidates, which may not support further development; actions of regulatory agencies, which may affect the initiation, timing and progress of clinical trials; BeiGene's ability to achieve market acceptance in the medical community necessary for commercial success; BeiGene's ability to obtain and maintain protection of intellectual property for its technology and drugs; BeiGene's reliance on third parties to conduct preclinical studies and clinical trials and to manufacture its products and product candidates; BeiGene's limited operating history and BeiGene's ability to obtain additional funding for operations and to complete the development and commercialization of its drug candidates, as well as those risks more fully discussed in the section entitled "Risk Factors" in BeiGene's most recent quarterly report on Form 10-Q, as well as discussions of potential risks, uncertainties, and other important factors in BeiGene's

subsequent filings with the U.S. Securities and Exchange Commission. All information in this press release is as of the date of this press release, and BeiGene undertakes no duty to update such information unless required by law.

Investor/Media Contact

Lucy Li, Ph.D.
+1 781-801-1800
ir@beigene.com
media@beigene.com

i ABRAXANE®, REVLIMID®, and VIDAZA® are registered trademarks of Celgene Corporation.

BeiGene,Ltd.

BeiGene Expands Global Pivotal Program for BTK Inhibitor BGB-3111

- Initiates global Phase 3 trial of BGB-3111 compared to bendamustine and rituximab (BR) in treatment-naïve chronic lymphocytic leukemia / small lymphocytic lymphoma (CLL/SLL) patients
- Initiates global pivotal Phase 2 trial of BGB-3111 in combination with GAZYVA® (obinutuzumab) in relapsed or refractory follicular lymphoma (FL) patients
- Completes enrollment of pivotal Phase 2 trial in China in mantle cell lymphoma (MCL) patients

CAMBRIDGE, Mass., and BEIJING, China, November 13, 2017 (GLOBE NEWSWIRE) -- BeiGene, Ltd. (NASDAQ:BGNE), a commercial-stage biopharmaceutical company focused on developing and commercializing innovative molecularly targeted and immuno-oncology drugs for the treatment of cancer, today announced the initiation of two new global pivotal clinical trials of BGB-3111, an investigational Bruton's Tyrosine Kinase (BTK) inhibitor, including a Phase 3 trial of BGB-3111 in previously untreated patients with CLL/SLL and a pivotal Phase 2 trial of BGB-3111 in combination with GAZYVA® (obinutuzumab) in patients with relapsed or refractory FL. Along with a global Phase 3 trial comparing BGB-3111 to ibrutinib in Waldenström's macroglobulinemia (WM), initiated in early 2017, BGB-3111 is now being evaluated in global pivotal trials in three distinct indications. Additionally, BGB-3111 is being evaluated in a broad pivotal clinical development program in China, including ongoing pivotal Phase 2 trials in MCL, CLL, and WM, which was initiated in August 2017. BeiGene also announced today that enrollment in the pivotal Phase 2 trial of BGB-3111 in China in MCL patients was completed in September 2017.

"The initiation of two additional pivotal trials expands our global registration-directed clinical development of BGB-3111 to additional indications, including patients with

follicular lymphoma, a common B cell malignancy for which BTK inhibitors are not yet approved. We look forward to continuing the development of BGB-3111 as a potentially best-in-class BTK inhibitor for patients worldwide who suffer from hematological malignancies," commented John V. Oyler, Founder, Chief Executive Officer, and Chairman of BeiGene.

"We believe that these two new pivotal trials are supported by our growing clinical experience with BGB-3111, in which over 600 patients have been dosed to date. With the newly initiated Phase 3 CLL/SLL trial, we aim to investigate whether BGB-3111 could be an effective treatment option for a broad population of CLL/SLL patients requiring initial treatment. The initiation of the pivotal trial in follicular lymphoma is an effort to determine whether the combination of BGB-3111 and obinutuzumab represents an effective treatment option for a high-unmet-need population of relapsed or refractory patients to potentially support the pursuit of accelerated or conditional approval of this regimen," commented Jane Huang, M.D., Chief Medical Officer, Hematology.

Trial Design

The Phase 3 trial in CLL/SLL is designed to compare BGB-3111 to BR and will be conducted in North America, Europe, Australia, New Zealand, and Asia. The study will enroll previously untreated CLL/SLL patients ineligible for intensive chemo-immunotherapy (i.e., fludarabine, cyclophosphamide, and rituximab), who will be divided into two cohorts. The first cohort is designed to include 420 patients without a 17p deletion (del17p), who will be randomized in a 1:1 ratio to receive either BGB-3111 until progression or six cycles of BR. Crossover will be allowed in the BR arm upon progression. The primary endpoint will be progression-free survival (PFS), and secondary endpoints include overall response rate (ORR), duration of response (DOR), overall survival (OS), and patient-reported outcomes. Patients with del17p will be enrolled in a second cohort to receive BGB-3111 until progression and will be assessed for response and safety.

The pivotal Phase 2 trial in FL is designed to evaluate BGB-3111 in combination with obinutuzumab in patients who have had at least two prior lines of therapy and who progressed within 12 months of their last treatment or were refractory to their last treatment. The primary endpoint will be ORR and obinutuzumab monotherapy will be included as a comparator, in order to evaluate the contribution of BGB-3111. The trial is expected to enroll approximately 210 patients in North America, Europe, Australia, and New Zealand who will be randomized 2:1 to receive either BGB-3111 with obinutuzumab or obinutuzumab alone. Patients in the obinutuzumab arm will have the option to add BGB-3111 after 12 months if a response has not been achieved. Secondary endpoints of the study include DOR, PFS, OS, and time to response.

About BGB-3111

BGB-3111 is a potent and highly selective investigational small molecule inhibitor of BTK. BGB-3111 has demonstrated higher selectivity against BTK than ibrutinib, a BTK inhibitor currently approved by the U.S. Food and Drug Administration and the European Medicines Agency, based on biochemical assays, higher exposure than ibrutinib based on their respective Phase 1 experience in separate trials, and sustained 24-hour BTK occupancy in both the peripheral blood and lymph node compartments.

About BeiGene

BeiGene is a global, commercial-stage, research-based biotechnology company focused on molecularly targeted and immuno-oncology cancer therapeutics. With a team of over 700 employees in China, the United States, and Australia, BeiGene is advancing a pipeline consisting of novel oral small molecules and monoclonal antibodies for cancer. BeiGene is also working to create combination solutions aimed to have both a meaningful and lasting impact on cancer patients. BeiGene markets ABRAXANE® (nanoparticle albumin–bound paclitaxel), REVLIMID® (lenalidomide), and VIDAZA® (azacitidine) in China under a license from Celgene Corporation. [i]

Forward-Looking Statements

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995 and other federal securities laws, including statements regarding the encouraging clinical data of BGB-3111 and BeiGene's advancement of, and anticipated clinical development and regulatory milestones and plans related to BGB-3111. Actual results may differ materially from those indicated in the forward-looking statements as a result of various important factors, including BeiGene's ability to demonstrate the efficacy and safety of its drug candidates; the clinical results for its drug candidates, which may not support further development; actions of regulatory agencies, which may affect the initiation, timing and progress of clinical trials; BeiGene's ability to achieve market acceptance in the medical community necessary for commercial success; BeiGene's ability to obtain and maintain protection of intellectual property for its technology and drugs; BeiGene's reliance on third parties to conduct preclinical studies and clinical trials; BeiGene's limited operating history and BeiGene's ability to obtain additional funding for operations and to complete the development and commercialization of its drug candidates, as well as those risks more fully discussed in the section entitled "Risk Factors" in BeiGene's most recent quarterly report on Form 10-Q, as well as discussions of potential risks, uncertainties, and other important factors in BeiGene's subsequent filings with the U.S. Securities and Exchange Commission. All information in this press release is as of the date of this press release, and BeiGene undertakes no duty to update such information unless required by law.

Investor/Media Contact

Lucy Li, Ph.D.
+1 781-801-1800
ir@beigene.com
media@beigene.com

i ABRAXANE®, REVLIMID®, and VIDAZA® are registered trademarks of Celgene Corporation.