UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

Report of Foreign Private Issuer
Pursuant to Rule 13a-16 or 15d-16
of the Securities Exchange Act of 1934

For the month of October 2015

Commission File Number 000-28508

Flamel Technologies S.A.

(Translation of registrant's name into English)

Parc Club du Moulin à Vent
33 avenue du Dr. Georges Levy
69693 Vénissieux Cedex France
(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F.

Form 20-F ☒ Form 40-F ☐

Indicate by check mark whether registrant by furnishing the information contained in this Form is also thereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.

Yes ☐ No ☒

If "Yes" is marked, indicate below the file number assigned to the registrant in connection with Rule 12g3-2(b): 82-_____

Attached hereto and incorporated herein by reference are the following: (i) as Exhibit 1 hereto, the registrant's press release dated October 5, 2015, (ii) as Exhibit 2 hereto, the registrant's press release dated September 9, 2015, (iii) as Exhibit 3 hereto, the registrant's press release dated July 30, 2015, and (iv) as Exhibit 4 hereto, the registrant's press release dated June 29, 2015.

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, thereunto duly authorized.

FLAMEL
TECHNOLOGIES, S.A.

Dated: October 13, 2015

/s/ Michael S. Anderson
Michael S. Anderson
Chief Executive Officer

EXHIBIT INDEX

Exhibit No.	Description
1.	Press Release dated October 5, 2015
2.	Press Release dated September 9, 2015
3.	Press Release dated July 30, 2015
4.	Press Release dated June 29, 2015

Exhibit 1

Flamel Ireland Limited Licenses LiquiTime® Technology to Perrigo for U.S. Over-the-Counter Market

LYON, FRANCE – October 5, 2015 – Flamel Technologies (NASDAQ: FLML) announced today that the Company's Irish subsidiary, Flamel Ireland Limited, has licensed exclusive U.S. rights to the LiquiTime® drug delivery platform to Perrigo's Irish subsidiary, Elan Pharma International Limited, for the U.S. Over-the-Counter (OTC) drug market. LiquiTime® is Flamel's drug delivery platform for modified, extended and controlled release of liquid oral drugs.

Flamel has entered into a multi-product agreement for extended release liquid OTC products in the U.S. market whereby Flamel will receive an upfront payment of $6.0 million and will be eligible for at least $50 million in approval and launch milestones for a minimum of seven products. In addition, Flamel will receive mid-single digit royalties on net sales. The exclusive license includes Flamel's LiquiTime® Ibuprofen and LiquiTime® Guaifenesin oral suspensions.

Flamel and Perrigo believe there is a large market opportunity for other OTC extended release liquid drug formulations, including products containing active ingredient combinations for the US cough/cold market, which analysts have estimated at $6.5 billion annually.

"We are very pleased to partner with Perrigo, one of the premier companies in the US Over-the-Counter consumer goods market. Flamel's LiquiTime® platform meets the unmet medical need for patients who are seeking the convenience of extended release liquid OTC medications. In addition, our LiquiTime® drug delivery platform is patent protected through September 2025 in the US and we will have the ability to apply for product-specific patents to extend those periods of patent protection," said Michael S. Anderson, Chief Executive Officer.

"This licensing agreement with Perrigo is another important validation of the quality and differentiation of Flamel's drug delivery platform technology. Moreover, we are pleased to accomplish one of our major commitments to investors in 2015 by putting these products into the hands of a leading OTC consumer products company that may result in an attractive set of milestone and royalty payments to our Company over time.

About Flamel Technologies - Flamel Technologies SA (NASDAQ: FLML) is a specialty pharmaceutical company utilizing its core competencies in formulation development and drug delivery to develop safer and more efficacious pharmaceutical products, addressing unmet medical needs and/or reducing overall healthcare costs. Flamel currently has approvals for and markets two previously Unapproved Marketed Drugs ("UMDs") in the USA, Bloxiverz® (neostigmine methylsulfate injection) and Vazculep® (phenylephrine hydrochloride injection). The Company intends to add to this branded business by creating additional products, focusing on the development of products utilizing Flamel's proprietary drug delivery platforms, and recently announced FDA acceptance of its third NDA filing with an FDA-assigned PDUFA date of April 30, 2016. Flamel also has several products in development utilizing Micropump® (oral sustained release microparticles platform) along with its tangent technologies, LiquiTime® and Trigger Lock™. The lead project for Micropump® is sodium oxybate. LiquiTime® allows for the extended-release of liquid medicines (such as ibuprofen and guaifenesin) and Trigger Lock™ is an abuse-resistant iteration of Micropump®, designed specifically for long-acting opioids (such as hydromorphone). Additionally, the Company has developed a long acting injectable platform, Medusa™, a hydrogel depot technology currently being studied with exenatide. Flamel's products are targeting high-value molecules and will utilize either the 505(b)(2) approval process for NDAs or biosimilar pathways ultimately approved by FDA and other regulatory authorities. The Company is headquartered in Lyon, France and has operations in St. Louis, Missouri, USA, and Dublin, Ireland. Additional information may be found at www.flamel.com.

Safe Harbor: *This release may include "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995. All statements herein that are not clearly historical in nature are forward-looking, and the words "anticipate," "assume," "believe," "expect," "estimate," "plan," "will," "may," and the negative of these and similar expressions generally identify forward-looking statements. All forward-looking statements involve risks, uncertainties and contingencies, many of which are beyond Flamel's control and could cause actual results to differ materially from the results contemplated in such forward-looking statements. These risks, uncertainties and contingencies include the risks relating to: our dependence on a small number of products and customers for the majority of our revenues; the possibility that our Bloxiverz® and Vazculep® products, which are not patent protected, could face substantial competition resulting in a loss of market share or forcing us to reduce the prices we charge for those products; the possibility that we could fail to successfully complete the research and development for the two pipeline products we are evaluating for potential application to the FDA pursuant to our "unapproved-to-approved" strategy, or that competitors could complete the development of such products and apply for FDA approval of such products before us; our dependence on the performance of third parties in partnerships or strategic alliances for the commercialization of some of our products; the possibility that our products may not reach the commercial market or gain market acceptance; our need to invest substantial sums in research and development in order to remain competitive; our dependence on certain single providers for development of several of our drug delivery platforms and products; our dependence on a limited number of suppliers to manufacture our products and to deliver certain raw materials used in our products; the possibility that our competitors may develop and market technologies or products that are more effective or safer than ours, or obtain regulatory approval and market such technologies or products before we do; the challenges in protecting the intellectual property underlying our drug delivery platforms and other products; our dependence on key personnel to execute our business plan; the amount of additional costs we will incur to comply with U.S. securities laws as a result of our ceasing to qualify as a foreign private issuer; and the other risks, uncertainties and contingencies described in the Company's filings with the U.S. Securities and Exchange Commission, including our annual report on Form 20-F for the year ended December 31, 2014, all of which filings are also available on the Company's website. Flamel undertakes no obligation to update its forward-looking statements as a result of new information, future events or otherwise, except as required by law.*

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Flamel Technologies Announces FDA Acceptance of Third NDA Filing

Lyon, France – September 9, 2015 – Flamel Technologies (NASDAQ: FLML) today announced that it has received a Prescription Drug User Fee Act (PDUFA) date of April 30, 2016 from the U.S. Food and Drug Administration (FDA) for its third New Drug Application (NDA). This is Flamel's third filing to seek FDA approval of an Unapproved Marketed Drug (UMD), consistent with Flamel's successful strategy in this portion of its product portfolio. Based on IMS and other third-party data, the Company estimates that current U.S. market sales of the unapproved versions of this drug are in the range of $70-$80 million per year. Flamel currently has two FDA-approved products on the market, including Bloxiverz® (neostigmine methylsulfate) and Vazculep® (phenylephrine hydrochloride).

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Flamel Technologies Announces Second Quarter Results of Fiscal Year 2015

Product revenue guidance for 2015 of $170-$185 million reaffirmed

Conference call with management to take place at 10:00 am ET on July 30, 2015

Lyon, France – July 30, 2015 - Flamel Technologies (NASDAQ: FLML) today announced its financial results for the second quarter of fiscal year 2015. Highlights from the quarter include:

- Product revenues of $50 million
- Submission of Unapproved Marketed Drug (UMD) #3 to FDA
- Positive results from two pharmacokinetic (PK) studies in healthy volunteers of abuse-deterrent, extended-release, oral hydromorphone product using Trigger Lock™ technology
- Addition of Sandy Hatten as Senior Vice President of Quality and Regulatory Affairs and Gregory J. Davis as its Vice President of Business and Corporate Development

“We are pleased to report that Bloxiverz® remained strong throughout the quarter with approximately 70% average share during the second quarter following the launch of Fresenius Kabi’s competing product early in early April. For the Company’s second approved marketed drug, Vazculep®, Flamel implemented a price increase in the second quarter for both the 5 and 10 mL vial sizes, for which it now supplies virtually the entire market, and continues to gain share in the 1mL vial size,” said Chief Executive Officer, Mike Anderson. “As we look forward for the full year, we are reaffirming product revenue guidance for 2015 of $170-185 million for both Bloxiverz and Vazculep and will update investors with our third quarter financial report,” added Mr. Anderson.

“The Company continues to focus on expanding its senior management team evident in the appointment of Sandy Hatten to head up Quality and Regulatory Affairs and Greg Davis to focus on Business and Corporate Development, both of whom have outstanding track records in the industry. We submitted our NDA to the FDA for our third unapproved marketed drug in the quarter, consistent with our objective. In our proprietary product pipeline, we were pleased to report positive results from two pilot pharmacokinetic (PK) studies in healthy volunteers of FT227, an abuse-deterrent, extended release oral hydromorphone product using our proprietary Trigger Lock™ drug delivery platform. These results continue a series of positive clinical trial data announcements by Flamel over the past 15 months,” concluded Mr. Anderson.

Flamel's Second Quarter Results

Flamel reported total revenues during the second quarter of 2015 of $49.8 million, an increase of $45.5 million in revenues compared to the prior year period. Total revenues is comprised of net sales of Bloxiverz of $45.5 million, Vazculep of $3.6 million and another product of $0.6 million. Net sales represent units that have been sold through to the hospitals and where the chargeback can be determined, particularly in a period when product prices may be changing. As of June 30, 2015, the Company had Deferred Revenues of $19.3 million, comprised of $17.5 million for Bloxiverz and $1.8 million for Vazculep, compared to $30.7 million and $1.4 million, respectively, in the first quarter of 2015.

Costs of goods and services sold for the second quarter of 2015 were $2.8 million compared to $507,000 in the second quarter of 2014. Research and development costs in the second quarter of 2015 totaled $7.2 million, compared to $3.6 million in the prior year period, an increase of 100%. A substantial portion of this increase is attributed to costs associated with filing of UMD #3 with the U.S. Food and Drug Administration (FDA) and continued investment in the Company's pipeline and other proprietary products. Selling, general and administrative costs were $5.9 million in the second quarter of 2015, an increase of 44% versus $4.1 million in the second quarter of 2014, due to increased stock compensation expense as a result of higher share price and accelerated vesting and costs associated with hiring new personnel. Amortization of R&D assets associated with the development of Bloxiverz and Vazculep was $3.1 million in the second quarter of 2015, consistent with the prior sequential quarter.

Total net interest income was $312,000 in the second quarter of 2015 compared to interest income of $94,000 in the second quarter of 2014. Interest expense has been largely eliminated with the Company's repayment of nearly all of its debt and lines of credit with a portion of the net proceeds from its offering of 12.4 million ADSs in mid-March 2014.

In the second quarter 2015, the Company had foreign exchange loss of ($3.6) million, compared to a foreign exchange gain of $292,000 in the prior year period. While our parent company in France uses the Euro as its functional currency, it holds over $80 million in assets that are U.S. Dollar denominated, which depreciated relative to the Euro over the second quarter.

Operating and net income (loss) includes remeasurement of the fair value of the acquisition liabilities which was an expense of ($32.0) million for the three months ended June 30, 2015 compared to ($12.6) million for the prior year period. These liabilities were incurred as a part of Flamel's acquisition of Éclat Pharmaceuticals in March 2012 and are tied to commercial sales of FDA-approved products as well as other factors described in our Form 20-F. Changes in the fair value of the acquisition liabilities, which are remeasured at each balance sheet date, are recognized in the Company's reported income (loss).

Net loss from Continuing Operations for the second quarter of 2015 was ($17.4) million versus net loss from Continuing Operations of ($20.2) million in the year-ago period. Loss per share from Continuing Operations was ($0.43) on both a basic and diluted level in the second quarter of 2015 versus loss per share from Continuing Operations (both basic and diluted) of ($0.53) in the second quarter of 2014.

Flamel is disclosing non-GAAP financial measures when providing financial results, including adjusted net loss and adjusted loss per share. Flamel believes that an evaluation of its ongoing operations (and comparison of current operations with historical and future operations) would be difficult if the disclosure of its financial results were limited to financial measures prepared only in accordance with generally accepted accounting principles (GAAP) in the U.S. In addition to disclosing its financial results in accordance with GAAP, Flamel is disclosing certain non-GAAP results that exclude fair value remeasurements, impairment of intangible assets, amortization expense of intangible assets, effects of accelerated reimbursement of certain debt instruments and unrealized foreign exchange gains and losses on assets and liabilities denominated in foreign currency and include operating cash flows associated with the acquisition liabilities and Royalty Agreements, in order to supplement investors' and other readers' understanding and assessment of the Company's financial performance. The Company's management uses these non-GAAP measures internally for forecasting, budgeting and measuring its operating performance. Investors and other readers are encouraged to review the related GAAP financial measures and the reconciliation of non-GAAP measures to their most closely applicable GAAP measure set forth below and should consider non-GAAP measures only as a supplement to, not as a substitute for or as a superior measure to, measures of financial performance prepared in accordance with GAAP.

Adjusted net income (non-GAAP) for the second quarter of 2015 was $13.9 million versus adjusted net loss (non-GAAP) of ($5.2) million in the second quarter of 2014. Adjusted earnings (non-GAAP) per share (diluted) was $0.34 in the second quarter of 2015 versus adjusted loss (non-GAAP) per share (diluted) of ($0.13) in the prior year period.

The Company's cash position as of June 30, 2015 was $116.1 million compared to $113.2 million as of March 31, 2015.

	Three months ended June 30,				Six months ended June 30,			
	2014		2015		2014		2015	
GAAP Net income (loss) and diluted earnings (loss) per share…...……	$ (21,073)	$ (0.55)	$ (17,400)	$ (0.43)	$ (47,711)	$ (1.43)	$ (5,753)	$ (0.14)
Fair value remeasurement of acquisition liabilities	12,607		32,000		27,233		37,254	
Fair value remeasurement of royalty agreement	1,079		2,726		1,235		2,985	
Amortization of Intangible R&D Assets	2,938		3,139		5,875		6,282	
Accelerated reimbursement of acquisition note	-		-		3,013		-	
Accelerated reimbursement of facility agreements	-		-		4,741		-	
Tax effects of the above items	-		-		(2,338)		-	
Earn-out acquisition payment payable	(383)		(9,379)		(994)		(15,175)	
Royalty payable	(54)		(1,270)		(141)		(2,116)	
Unrealized foreign exchange (gain)/loss	(274)		4,045		(481)		(5,205)	
Adjusted Net Income (Loss) and adjusted diluted earnings (loss) per share,	$ (5,161)	$ (0.13)	$ 13,861	$ 0.34	$ (9,569)	$ (0.29)	$ 18,272	$ 0.45

A conference call to discuss these results and other updates is scheduled for **10:00 AM ET on Thursday, July 30, 2015.** A question and answer period will follow management's prepared remarks. To participate in the conference call, investors are invited to dial 888-684-1282 (U.S. and Canada) or +1-913-312-1513 (international). The conference ID number is 1557700. The conference call webcast may be accessed at the investors section of www.flamel.com. The archived webcast of the conference call will be available for 90 days on Flamel's website.

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Condensed Consolidated Statements of Operations
(Amounts in thousands, except per share data)

	Three months ended June 30,		Six Months Ended June 30,	
	2014	2015	2014	2015
Revenue:				
License and research revenue	$1,819	-	$2,417	-
Product sales and services	$2,483	49,765	6,443	$82,476
Other revenues	$16	30	36	45
Total revenue	4,318	49,795	8,896	82,521
Costs and expenses:				
Cost of goods and services sold	(507)	(2,756)	(1,280)	(6,386)
Research and development	(3,600)	(7,204)	(7,544)	(13,226)
Selling, general and administrative	(4,065)	(5,873)	(7,581)	(10,336)
Fair value remeasurement of acquisition liabilities, incl. related parties	(12,607)	(32,000)	(27,233)	(37,254)
Amortisation of intangible R&D assets	(2,938)	(3,139)	(5,875)	(6,282)
Acquisition note expenses, incl. related parties	-	-	(3,013)	-
Total	(23,717)	(50,972)	(52,526)	(73,484)
Profit (loss) from continuing operations	(19,399)	(1,177)	(43,630)	9,037
Interest income	.. 94	653	139	1,451
Interest expense	-	(341)	(5,552)	(482)
Interest expense on debt related to the royalty agreement with related parties	(1,079)	(2,726)	(1,235)	(2,985)
Foreign exchange gain (loss)	292	(3,565)	471	7,936
Other income (loss)	29	(2)	81	5
Income (loss) before income taxes from continuing operations	(20,063)	(7,158)	(49,726)	14,962
Income tax benefit (expense)	(137)	(10,242)	2,665	(20,715)
Net income (loss) from continuing operations	($20,200)	($17,400)	($47,061)	($5,753)
Net income from discontinued operations	($873)	$0	($650)	$0
Net income (loss)	($21,073)	($17,400)	($47,711)	($5,753)
Earnings (loss) per ordinary share (Basic):				
Continuing operations	($0.53)	($0.43)	($1.41)	($0.14)
Discontinued operations	($0.02)	$0.00	($0.02)	$0.00
Net income (loss)	($0.55)	($0.43)	($1.43)	($0.14)
Weighted average number of shares outstanding (in thousands) :				
Basic	38,438	40,353	33,403	40,281
Diluted	38,438	40,353	33,403	40,281

Flamel Technologies Announces Positive Results from First Clinical Trials with Trigger Lock™ Hydromorphone Abuse-Deterrent Product

Meeting with FDA will be requested before the end of 2015

Lyon, France – June 29, 2014 - Flamel Technologies (NASDAQ: FLML) today announced positive results from two pilot pharmacokinetic (PK) studies in healthy volunteers of FT227, an abuse-deterrent, extended-release, oral hydromorphone product using its proprietary Trigger Lock™ drug delivery platform. Flamel's Trigger Lock™ allows the development of abuse-deterrent extended release formulations of opioids and other drugs susceptible to abuse. Hydromorphone is used for relief of moderate to severe pain in patients requiring continuous around-the-clock opioid treatment for an extended period of time.

The PK studies were intended to provide sufficient data for the Company to select a preferred prototype formulation to move forward into pivotal studies. The studies compared three FT227 prototypes to the comparator product Jurnista© (sold as Exalgo© in the United States) in both fasted and fed conditions at a dose of 32mg.

Under fasted conditions, comparing the area under the curve (AUC) and the peak plasma concentration (C_{max}) of FT227 to Jurnista© in 16 subjects, the results identified a FT227 formulation that met the bioequivalence criteria for both parameters. Under fed conditions (14 subjects), the same formulation was bioequivalent in terms of AUC to Jurnista© but outside of the C_{max} bioequivalence criterion at the lower confidence interval level. Comparing the effect of food on the PK parameters of the FT227 prototypes across the two studies, no notable difference is seen in either AUC or C_{max} in fed and fasted conditions. This suggests that administration of FT227 will not be subject to a clinically relevant food effect.

In both studies FT227 was well tolerated and no serious adverse events were reported.

In addition, Flamel has generated substantial in vitro data comparing the abuse deterrence properties of FT227 compared to other marketed abuse-deterrent opioid products. The Company is confident that Trigger Lock™ is a robust platform for opioids that will set a high standard in terms of abuse deterrence. Further abuse deterrence data are being generated by an independent contract research organization and will be subject to further announcements. Flamel is planning to meet with the U.S. Food and Drug Administration (FDA) before the end of 2015 to discuss the remainder of the development plan for FT227. The product is designed to be filed as a 505(b)(2) New Drug Application (NDA). Based on current expectations, the selected formulation will be scaled-up over the coming months and the Company plans to begin pivotal registration studies by mid-2016.

"We are very pleased with these initial data on FT227 Trigger Lock™ and we are highly confident in our Trigger Lock™ abuse deterrent platform. We look forward to meeting with the FDA and to moving FT227 into pivotal studies in 2016," said Mike Anderson, Chief Executive Officer.

As of December 2014, Trigger Lock™ is protected by seven Flamel patent application families, which expire between November 2025 and December 2033.

About Flamel Technologies - Flamel Technologies SA (NASDAQ:FLML) is a specialty pharmaceutical company utilizing its core competencies in formulation development and drug delivery to develop safer and more efficacious pharmaceutical products, addressing unmet medical needs and/or reducing overall healthcare costs. Flamel currently has approvals for and markets two previously Unapproved Marketed Drugs ("UMDs") in the USA, Bloxiverz™ (neostigmine methylsulfate injection) and Vazculep™ (phenylephrine hydrochloride injection). The Company intends to add to this branded business by creating additional products, focusing on the development of products utilizing Flamel's proprietary drug delivery platforms. Flamel currently has several products in development utilizing Micropump® (oral sustained release microparticles platform) along with its tangent technologies, LiquiTime® and Trigger Lock™. The lead project for Micropump is Sodium Oxybate. LiquiTime allows for the extended-release of liquid medicines (such as Ibuprofen and Guaifenesin) and Trigger Lock is an abuse-resistant iteration of Micropump, designed specifically for long-acting opioids. Additionally, the Company has developed a long acting injectable platform, Medusa™, a hydrogel depot technology currently being studied with Exenatide. Flamel's products are targeting high-value molecules and will utilize either the 505(b)(2) approval process for NDAs or biosimilar pathways ultimately approved by FDA and other regulatory authorities. The Company is headquartered in Lyon, France and has operations in St. Louis, Missouri, USA, and Dublin, Ireland. Additional information may be found at www.flamel.com.

Safe Harbor: *This release includes statements concerning our operating results (including product sales), financial condition and product development milestones, which are "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995. All statements herein that are not clearly historical in nature are forward-looking, and the words "anticipate," "assume," "believe," "expect," "estimate," "plan," "will," "may," and the negative of these and similar expressions generally identify forward-looking statements. All forward-looking statements involve risks, uncertainties and contingencies, many of which are beyond Flamel's control and could cause actual results to differ materially from the results contemplated in such forward-looking statements. These risks, uncertainties and contingencies include the risks relating to: our dependence on a small number of products and customers for the majority of our revenues; the possibility that our Bloxiverz® and Vazculep™ products, which are not patent protected, could face substantial competition resulting in a loss of market share or forcing us to reduce the prices we charge for those products; the possibility that we could fail to successfully complete the research and development for the two pipeline products we are evaluating for potential application to the FDA pursuant to our "unapproved-to-approved" strategy, or that competitors could complete the development of such products and apply for FDA approval of such products before us; our dependence on the performance of third parties in partnerships or strategic alliances for the commercialization of some of our products; the possibility that our products may not reach the commercial market or gain market acceptance; our need to invest substantial sums in research and development in order to remain competitive; our dependence on certain single providers for development of several of our drug delivery platforms and products; our dependence on a limited number of suppliers to manufacture our products and to deliver certain raw materials used in our products; the possibility that our competitors may develop and market technologies or products that are more effective or safer than ours, or obtain regulatory approval and market such technologies or products before we do; the challenges in protecting the intellectual property underlying our drug delivery platforms and other products; our dependence on key personnel to execute our business plan; the possibility that we may cease to qualify as a foreign private issuer, which would increase the costs and expenses we incur to comply with U.S. securities laws; and the other risks, uncertainties and contingencies described in the Company's filings with the U.S. Securities and Exchange Commission, including our annual report on Form 20-F for the year ended December 31, 2014, all of which filings are also available on the Company's website. Flamel undertakes no obligation to update its forward-looking statements as a result of new information, future events or otherwise, except as required by law.*

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