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Aeterna Zentaris Inc.  
Form 6-K  
August 13, 2008

FORM 6-K  
SECURITIES AND EXCHANGE COMMISSION  
Washington, D.C. 20549

REPORT OF FOREIGN ISSUER

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

For the month of August 2008

Commission File No. 000-30752

AETERNA ZENTARIS INC.  
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1405, boul. du Parc-Technologique  
Quebec, Quebec  
Canada, G1P 4P5  
(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 /X/ Form 40-F / /

Indicate by check mark whether the 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 /X/

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

DOCUMENTS INDEX

DOCUMENTS DESCRIPTION

1. Press Release dated August 12, 2008: Aeterna Zentaris Reports Second  
Quarter 2008 Financial and Operating Results

[AETERNA ZENTARIS LOGO]

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PRESS RELEASE  
For immediate release

AETERNA ZENTARIS REPORTS SECOND QUARTER 2008 FINANCIAL AND OPERATING RESULTS

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ALL AMOUNTS ARE IN U.S. DOLLARS

QUEBEC CITY, CANADA, AUGUST 12, 2008 - AETerna Zentaris Inc. (NASDAQ: AEZS, TSX: AEZ), a global biopharmaceutical company focused on endocrinology and oncology, today reported financial and operating results for the second quarter ended June 30, 2008.

### SECOND QUARTER 2008 HIGHLIGHTS

- Advancement of cetrorelix Phase 3 program in BPH;
  - First efficacy trial: patient enrollment completed;
  - Second efficacy trial: patient enrollment ongoing / completion planned for third quarter of 2008 remains on target;
  - Safety trial: first patient dosing initiated.
  
- Completion of sale of Quebec City building for \$7.1 million.

Juergen Ernst, Chairman, Interim President and CEO at AETerna Zentaris commented, "During the quarter, we achieved our key objectives as our Phase 3 program in BPH with our lead compound cetrorelix met all recruitment goals and remains on track, with first results expected in the third quarter of 2009. Furthermore, we monetized our Quebec City building which provided additional non-dilutive funding. Over the next few months, we will focus on advancing our Phase 3 program in BPH with cetrorelix, while we endeavour to conclude additional non-dilutive transactions and strategic partnerships."

### CONSOLIDATED RESULTS FOR THE SECOND QUARTER ENDED JUNE 30, 2008

CONSOLIDATED SALES AND ROYALTIES increased to \$8.2 million for the three-month period ended June 30, 2008, compared to \$7.7 million for the same period in 2007. The increase in sales and royalties for the three-month period ended June 30, 2008 is related primarily to additional sales of Cetrotide(R), partly offset by the exclusion of sales from Impavido(R) in the second quarter of 2008.

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LICENSE FEES REVENUES decreased to \$2.2 million for the three-month period ended June 30, 2008, compared to \$3.9 million for the same period in 2007. The decrease for the three-month period ended June 30, 2008, is mainly attributable to the termination of the Company's licensing agreement with Solvay in 2007, which triggered additional amortization of upfront payments in the second quarter of 2007.

CONSOLIDATED R&D COSTS, NET OF TAX CREDITS AND GRANTS were \$17.3 million for the three-month period ended June 30, 2008 compared to \$7.8 million for the same period in 2007. Additional R&D expenses for the three-month period ended June 30, 2008, are mainly related to the advancement of our Phase 3 program in BPH with our lead product, cetrorelix.

CONSOLIDATED SELLING, GENERAL AND ADMINISTRATIVE (SG&A) EXPENSES were \$6.6 million for the three-month period ended June 30, 2008 compared to \$4.5 million for the same period in 2007. The increase in SG&A expenses for the three-month period ended June 30, 2008 is primarily due to non-recurring corporate expenses related to organizational changes, including severance paid to the former President and CEO as well as to the Senior Vice President and CBO that were implemented in the second quarter of 2008.

CONSOLIDATED NET LOSS for the three-month period ended June 30, 2008 was \$20.6 million or \$0.39 per basic and diluted share, compared to \$4.8 million or \$0.09

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per basic and diluted share for the same period in 2007. The increase in net loss for the three-month period ended June 30, 2008, compared to the same period in 2007, is primarily attributable to the increased R&D costs related to the advancement of cetrorelix into our Phase 3 program for the treatment of BPH and non-recurring SG&A corporate costs.

The consolidated cash and short-term investments were \$24.8 million as at June 30, 2008.

### CONFERENCE CALL

Management will be hosting a conference call for the investment community beginning at 4:30 p.m. Eastern Time today, Tuesday, August 12, to discuss second quarter 2008 results. To participate in the live conference call by telephone, please dial 416-644-3430, 514-807-8791 or 800-814-4862. Individuals interested in listening to the conference call on the Internet may do so by visiting [www.aezsinc.com](http://www.aezsinc.com). A replay will be available on the Company's Web site for 30 days.

### ABOUT AETERNA ZENTARIS INC.

Aeterna Zentaris Inc. is a global biopharmaceutical company focused on endocrine therapy and oncology with proven expertise in drug discovery, development and commercialization.

News releases and additional information are available at [www.aezsinc.com](http://www.aezsinc.com).

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### FORWARD-LOOKING STATEMENTS

This press release contains forward-looking statements made pursuant to the safe harbor provisions of the U.S. Securities Litigation Reform Act of 1995. Forward-looking statements involve known and unknown risks and uncertainties, which could cause the Company's actual results to differ materially from those in the forward-looking statements. Such risks and uncertainties include, among others, the availability of funds and resources to pursue R&D projects, the successful and timely completion of clinical studies, the ability of the Company to take advantage of business opportunities in the pharmaceutical industry, uncertainties related to the regulatory process and general changes in economic conditions. Investors should consult the Company's quarterly and annual filings with the Canadian and U.S. securities commissions for additional information on risks and uncertainties relating to the forward-looking statements. Investors are cautioned not to rely on these forward-looking statements. The Company does not undertake to update these forward-looking statements. We disclaim any obligation to update any such factors or to publicly announce the result of any revisions to any of the forward-looking statements contained herein to reflect future results, events or developments except if we are requested by a governmental authority or applicable law.

### CONTACTS

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ATTACHMENT: Financial summary

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(IN THOUSANDS OF US DOLLARS, EXCEPT SHARE  
AND PER SHARE DATA)

CONSOLIDATED RESULTS UNAUDITED	THREE MONTHS ENDED JUNE 30,		2009
	2008	2007	
	\$	\$	\$
REVENUES			
Sales and royalties	8,250	7,698	10,948
License fees	2,207	3,853	6,060
	10,457	11,551	17,008
OPERATING EXPENSES			
Cost of sales	4,758	3,075	7,833
R&D costs, net of tax credits and grants	17,345	7,815	25,160
Selling, general and administrative	6,606	4,517	11,123
Depreciation and amortization			
Property, plant and equipment	397	392	789
Intangible assets	876	928	1,804
	29,982	16,727	46,706
LOSS FROM OPERATIONS	(19,525)	(5,176)	(24,701)
OTHER REVENUES (EXPENSES)			
Interest income	311	300	611
Interest expense	(53)	(53)	(106)
Foreign exchange (loss) gain	(502)	(637)	(1,139)
Loss on disposal of long-lived assets held for sale	(810)	-	(810)
	(1,054)	(390)	(1,444)
LOSS BEFORE INCOME TAXES	(20,579)	(5,566)	(26,145)
INCOME TAX RECOVERY	-	731	731
NET LOSS FROM CONTINUING OPERATIONS	(20,579)	(4,835)	(25,416)
NET (LOSS) EARNINGS FROM DISCONTINUED OPERATIONS	-	(11)	(11)
NET LOSS FOR THE PERIOD	(20,579)	(4,846)	(25,427)
NET LOSS PER SHARE FROM CONTINUING OPERATIONS			
Basic and diluted	(0.39)	(0.09)	(0.48)
NET LOSS PER SHARE			
Basic and diluted	(0.39)	(0.09)	(0.48)

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WEIGHTED AVERAGE NUMBER OF SHARES

Basic and diluted 53,187,470 53,179,470 53,18

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(IN THOUSANDS OF US DOLLARS)  
 CONSOLIDATED BALANCE SHEETS  
 UNAUDITED

	JUNE 30, 2008	Decemb 200
	\$	\$
Cash and short-term investments	24,827	41
Other current assets	18,018	18
	-----	-----
Long-term assets	42,845	59
	52,704	63
	-----	-----
Total assets	95,549	123
	=====	=====
Current liabilities	24,046	22
Deferred revenues	3,112	3
Long-term payable	235	
Employee future benefits	10,337	9
	-----	-----
Shareholders' equity	37,730	34
	57,819	88
	-----	-----
Total liabilities and shareholders' equity	95,549	123
	=====	=====

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SIGNATURE

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.

AETERNA ZENTARIS INC.

Date: August 13, 2008

By: /s/Dennis Turpin

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 Dennis Turpin  
 Senior Vice President, Chief Financial Officer