

ELITE PHARMACEUTICALS INC /DE/
 Form 4
 November 15, 2006

FORM 4

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

OMB APPROVAL

OMB Number: 3235-0287
 Expires: January 31, 2005
 Estimated average burden hours per response... 0.5

Check this box if no longer subject to Section 16. Form 4 or Form 5 obligations may continue. See Instruction 1(b).

STATEMENT OF CHANGES IN BENEFICIAL OWNERSHIP OF SECURITIES

Filed pursuant to Section 16(a) of the Securities Exchange Act of 1934, Section 17(a) of the Public Utility Holding Company Act of 1935 or Section 30(h) of the Investment Company Act of 1940

(Print or Type Responses)

1. Name and Address of Reporting Person *
BEHL CHARAN

2. Issuer Name and Ticker or Trading Symbol
ELITE PHARMACEUTICALS INC /DE/ [ELI]

5. Relationship of Reporting Person(s) to Issuer
 (Check all applicable)

(Last) (First) (Middle)
C/O ELITE PHARMACEUTICALS INC, 165 LUDLOW AVENUE
 (Street)

3. Date of Earliest Transaction (Month/Day/Year)
11/13/2006

____ Director
 Officer (give title below) _____ 10% Owner
 _____ Other (specify below)
Exec V.P. & Chief Sci Officer

NORTHVALE, NJ 07647

4. If Amendment, Date Original Filed(Month/Day/Year)

6. Individual or Joint/Group Filing(Check Applicable Line)
 Form filed by One Reporting Person
 ____ Form filed by More than One Reporting Person

(City) (State) (Zip)

Table I - Non-Derivative Securities Acquired, Disposed of, or Beneficially Owned

1. Title of Security (Instr. 3)	2. Transaction Date (Month/Day/Year)	2A. Deemed Execution Date, if any (Month/Day/Year)	3. Transaction Code (Instr. 8)	4. Securities Acquired (A) or Disposed of (D) (Instr. 3, 4 and 5)	5. Amount of Securities Beneficially Owned Following Reported Transaction(s) (Instr. 3 and 4)	6. Ownership Form: Direct (D) or Indirect (I) (Instr. 4)	7. Nature of Ownership (Instr. 4)
Common Stock				(A) or (D)	416,000	D	

Reminder: Report on a separate line for each class of securities beneficially owned directly or indirectly.

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SEC 1474 (9-02)

Table II - Derivative Securities Acquired, Disposed of, or Beneficially Owned (e.g., puts, calls, warrants, options, convertible securities)

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1. Title of Derivative Security (Instr. 3)	2. Conversion or Exercise Price of Derivative Security	3. Transaction Date (Month/Day/Year)	3A. Deemed Execution Date, if any (Month/Day/Year)	4. Transaction Code (Instr. 8)	5. Number of Derivative Securities Acquired (A) or Disposed of (D) (Instr. 3, 4, and 5)	6. Date Exercisable and Expiration Date (Month/Day/Year)	7. Title and Amount of Underlying Securities (Instr. 3 and 4)		
				Code	V (A) (D)	Date Exercisable	Expiration Date	Title	Amount of Number of Shares
Option	\$ 2.25	11/13/2006		A	250,000	11/13/2006	11/13/2016	Common Stock	250,000
Option	\$ 2.25	11/13/2006		A	300,000	(1)	(1)	Common Stock	300,000
Option	\$ 2.25	11/13/2006		A	200,000	(2)	(2)	Common Stock	200,000
Warrant	\$ 1.54					10/06/2004	10/06/2010	Common Stock	100,000
Warrant	\$ 3					04/26/2006	12/14/2010	Common Stock	30,000

Reporting Owners

Reporting Owner Name / Address	Relationships			
	Director	10% Owner	Officer	Other
BEHL CHARAN C/O ELITE PHARMACEUTICALS INC 165 LUDLOW AVENUE NORTHVALE, NJ 07647			Exec V.P. & Chief Sci Officer	

Signatures

/s/ Charan Behl 11/15/2006

**Signature of Reporting Person

Date

Explanation of Responses:

* If the form is filed by more than one reporting person, see Instruction 4(b)(v).

** Intentional misstatements or omissions of facts constitute Federal Criminal Violations. See 18 U.S.C. 1001 and 15 U.S.C. 78ff(a).

(1) They vest upon the closing of an exclusive product license for the United States national market, the entire European Union market or the Japan market or product sale transaction of all of the Company's ownership rights in the United States (only once for each individual product) for the Company's first "Non-Generic Opioid Drug" as to 150,000 options and for the Company's second "Non-Generic Opioid Drug" as to 150,000 options.

(2) See remarks for Footnote 2

Remarks:

(2) They vest as follows: (i) upon the commencement of the first Phase III clinical trial relating to the first "Non-Generic Opioid Drug" developed by the Company as to 125,000 options and relating to the second "Non-Generic Opioid Drug" developed by the Company as to 125,000 options.

by the Company as to 75,000 options; (ii) 50,000 options upon the closing of an exclusive product license for the United States national market or product sale transaction of all of the Company's ownership rights (on a product by product basis and only once for each individual product) for each Company drug product, other than the "Non-Generic Opioid Drugs" for which the foregoing "Non-Generic Opioid Drug" were granted under (i) above; (iii) 10,000 options upon the filing by the Company (in the Company's name) with the United States Food and Drug Administration (the "FDA") of either an abbreviated drug application (an "ANDA") or a new drug application (including a NDA filed with the FDA (a "NDA"), for a product not covered by a previous FDA application; (iv) 40,000 options upon the approval by the FDA of any ANDA or NDA (filed in the Company's name) for a product not previously approved by the FDA; (v) 25,000 options upon filing of an application for U.S. patent by the Company (filed in the Company's name); and (vi) 25,000 options upon the granting by U.S. Patent and Trademark Office of a patent to the Company (filed in the Company's name).

Note: File three copies of this Form, one of which must be manually signed. If space is insufficient, *see* Instruction 6 for procedure.

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