

Item 7.01 Regulation FD Disclosure

On October 21, 2009, Antigenics Inc. announced that the Committee for Medicinal Products for Human Use (CHMP), of the European Medicines Agency, verbally informed the company at an oral meeting to anticipate a negative opinion on the marketing authorization application for Oncophage (vitespen) in early-stage, localized renal cell carcinoma (kidney cancer). Antigenics will evaluate its options, including an appeal of this decision, after the CHMP has formally adopted an opinion at the November 2009 plenary meeting.

The full text of the press release issued in connection with the announcement is being furnished as Exhibit 99.1 to this current report on Form 8-K.

Item 9.01 Financial Statements and Exhibits

(d) Exhibits

The following exhibit is furnished herewith:

99.1 Press Release dated October 21, 2009

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.

ANTIGENICS INC.

Date: October 21, 2009 By: /s/ Shalini Sharp

Shalini Sharp
CFO

EXHIBIT INDEX

Exhibit No. Description of Exhibit

99.1	Press Release dated October 21, 2009
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