GLAXOSMITHKLINE PLC Form 6-K May 06, 2009

#### 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 period ending May 2009

GlaxoSmithKline plc (Name of registrant)

980 Great West Road, Brentford, Middlesex, TW8 9GS (Address of principal executive offices)

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

Form 20-F x Form 40-F

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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

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Securities Exchange Act of 1934.

Yes No x

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Issued: Wednesday 6 th May 2009

Research Triangle Park

NC

# GlaxoSmithKline to Divest US Rights for Wellbutrin XL

(R)

#### to Biovail for \$510 Million

GlaxoSmithKline plc (GSK) today announced that it has entered into an agreement to divest full commercial rights to Wellbutrin XL

VVC

in the United States to Biovai I International Laboratories SRL

, a subsidiary of Biovail Corporation (NYSE, TSX:

**BVF** 

), for \$510 million (£340 million).

The agreement is subject to Hart-Scott-Rodino regulatory clearance in the United States

Generic competition to

Wellbutrin XL

began at the end of 2006 for the 300mg tablet and during the second quarter of 2008 for the 150mg tablet.

US

s

ales of

Wellbutrin XL

in the first quarter of 2009 were £45 million (-70%).

"We are actively reshaping our

US

business and managing the transition occurring in our product portfolio

," said

Deirdre Connelly, President North American Pharmaceuticals

, GlaxoSmithKline

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This transaction is one

of a series of actions we are taking

to maximi

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e the value of our current assets and

to enable us to r

esource and invest

in

new products and upcoming launches."

Under the terms of the agreement, GSK will transfer the US NDA and license the

Wellbutrin XL

trademark to Biovail for use in the

US

. GSK will retain existing rights to

Wellbutrin XL

(excluding

Canada

) for countries outside the

US

Sales of

Wellbutrin XL

outside the

US

were £7 million in the first quarter of 2009.

GSK expects to record a pre-tax gain of approximately £340 million in Other Operating Income as a result of

this

divestment. The company

now expects the combined total of Other Operating Income and profit on disposal of interests in associates

to be around £700 million in 2009.

Wellbutrin XL

(bupropion hydrochloride extended-release tablets) is indicated for the treatment of major depressive disorder and seasonal affective disorder. It was developed by Biovail and has been distributed by GSK in the

**United States** 

since September 2003.

#### GlaxoSmithKline

- one of the world's leading research-based pharmaceutical and healthcare companies - is committed to improving the quality of human life by enabling people to do more, feel better and live longer. For further information please visit

www.gsk.com

S M Bicknell Company Secretary

06 May 2009

#### GlaxoSmithKline

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## Cautionary statement regarding forward-looking statements

Under the safe harbor provisions of the U.S. Private Securities Litigation Reform Act of 1995, GSK cautions investors that any forward-looking statements or projections made by GSK, including those made in this announcement, are subject to risks and uncertainties that may cause actual results to differ materially from those projected. Factors that may affect GSK's operations are described under 'Risk Factors' in the 'Business Review' in the company's Annual Report on Form 20-F for 2008.

#### **Trademarks**

Brand names appearing in italics throughout this document are trademarks of GSK.

#### **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 authorised.

GlaxoSmithKline plc (Registrant)

Date: May 06, 2009

By: VICTORIA WHYTE

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Victoria Whyte Authorised Signatory for and on behalf of GlaxoSmithKline plc