

The study listed may include approved and non-approved uses, formulations or treatment regimens. The results reported in any single study may not reflect the overall results obtained on studies of a product. Before prescribing any product mentioned in this Register, healthcare professionals should consult prescribing information for the product approved in their country.

<b>Statement of Results status for the study</b>	
<b>Name of Company:</b> GlaxoSmithKline	
<b>Study No:</b> MVC116278 (EudraCT#2011-004435-31)	
<b>Product:</b> lamivudine; maraviroc; zidovudine; zidovudine/lamivudine	
<b>Title:</b> An Expanded Access Protocol for Subjects Who Have Completed Clinical Studies Involving Maraviroc	
<b>Phase:</b> N/A	
<b>Study Period:</b> May 2012 to August 2016	
<b>Centers:</b> 29	
<b>Number of Subjects:</b> 120	
<b>Publications:</b> NA	
<b>Statement on results availability:</b> This is an Expanded Access Program and no participant data is expected to be collected. Thus, no study results are expected in this protocol.	