

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 Discontinuation of the study</b>	
<b>Name of Company:</b> GlaxoSmithKline	
<b>Study No:</b> 111846 (EudraCT# 2008-003318-81)	
<b>Product:</b> SB208136	
<b>Title:</b> A phase IV, open, randomized, controlled study to demonstrate the non-inferiority of co-administration of GSK Biologicals' live attenuated measles-mumps-rubella-varicella vaccine and Baxter's Neisseria meningitidis C conjugate vaccine versus separate administration of each of the vaccines in healthy children aged 12 through 23 months	
<b>Phase:</b> IV	
<b>Study Period:</b> NA	
<b>Centres:</b> NA	
<b>Number of Subjects:</b> 0	
<b>Publications:</b> NA	
<b>Statement on discontinuation of the study:</b> Discontinued by GlaxoSmithKline during preparation of the trial. No patient entered the study; therefore, no results / data are available.	