

## **Prematurely ended - Statement**

<b>EudraCT Number:</b>	2008-000653-36
<b>Full title of the study:</b>	Efficacy and tolerability of Docetaxel-Gemcitabin in patients with advanced non-small cell lung cancer and an ECOG-Performance-Index of 2. A multicenter phase II-trial.
<b>Sponsor-Code:</b>	DOCE-GEM
<b>Sponsor:</b>	Charité – Universitätsmedizin Berlin
<b>Principal Investigator:</b>	Medizinische Klinik m. S. Infektiologie und Pneumologie Dr. med. Daniel Binder
<b>Product:</b>	DOCETAXEL, 80 mg (Taxotere EU/1/95/002/001) GEMCITABINHYDROCHLORID, 200 mg (Gemzar Zul. Nr. :52222.00.00)
<b>Date of early termination:</b>	2010/07/08
<b>Statement:</b>	<p>The trial will be terminated because of insufficient recruiting. The planned number of recruited patients can not be achieved in an appropriate period of time.</p> <p>At present, there are no patients receiving any study medication</p> <p>The early termination occurs at a time point when only few of the planned patients were recruited. Thus, it is not possible to get reliable results/conclusions regarding primary and secondary endpoints. The trial will be terminated without a statement on the study endpoints.</p>