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Dutch Chamber of Commerce
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Subject

Results posting of 2007-001089-33: "Infusion of a single dose of erythropoietin to Prevent Injury in an Ischemia Reperfusion forearm model - A randomised cross-over study to evaluate if infusion of a single dose of EPO protects against ischemia-reperfusion injury in man"

Dear EMA,

On behalf of the Prof. dr. G.A.P.J.M. Rongen as the 2nd Sponsor of this study, I hereby confirm that the screening and inclusion of subjects in the following clinical trial protocol: "Infusion of a single dose of erythropoietin to Prevent Injury in an Ischemia Reperfusion forearm model - A randomised cross-over study to evaluate if infusion of a single dose of EPO protects against ischemia-reperfusion injury in man" with EudraCT number 2007-001089-33 (favourable Ethics Committee opinion on Sep 23rd 2008), **has never started**.

The reason for this is that the study product, Annexin A5, which was required to measure the primary study outcome, **did not come available**.

The Declaration End of Trial Form B7 reported "Completed", but this should be "**Prematurely Ended**".

We hope this information is satisfactory to finalize results posting.

With kind regards,

Arjan Nooteboom,
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cc Prof.dr. G.A.P.J.M Rongen