News & Events

May 2010

AlphaBeta Pharma will be present at "Accelerating Early Clinical Development Conference"

On -13th – 14th October 2010, Maritim proArte Hotel Berlin, Berlin Germany

AlphaBeta Pharma is Leading the Way to Maximise Value and Reduce Timelines with Efficient Decision Making, Effective Phase I – II Transition and Driving Earlier Attrition

AlphaBeta Pharma will be sponsoring "Optimising efficiency in meeting regulatory authority requests" Workshop on 15th October.

Registration 08:00 – Start 08:30 – Lunch 12:00 – Afternoon Start 13:00 – Finish 15:30

Why delegates should attend this workshop:

Delays in clinical trials can often occur when striving to meet regulatory authority requests. Crucial to moving as quickly as possible through your early stage trials, is dealing with the regulatory authorities in an effective manner.

This workshop will provide you with:

- Practical Strategies
- Case Studies
- Real-life Examples

On how to successfully and efficiently meet requests from the regulatory authorities

What this workshop will cover:

- How to design a clinical development programme accepted by the Regulatory Authorities
- How to effectively communicate with the Regulatory Authorities
- How to reduce timelines on submissions and approval
- Exploring the discrepancies between National versus EMA advice
- How to successfully employ the Scientific Advice
- To what extent can SA help to speed/improve quality of drug development in early phases?
- Critical CMC submissions and interactions with Regulatory Authorities to avoid approval delays

If you have any questions or queries, please email them to AlphaBeta Pharma info@alphabetapharma.com

Ends

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