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## Biomanufacturing Innovation Group

### Overcoming the Challenges in compliant manufacture of ATMPs

**Monday 4<sup>th</sup> July 2011**  
**Agenda**

Time	Item	
09.30 - 10.00	<b>Registration and Coffee</b>	
10.00 - 10.30	Introduction and Welcome	Mark Bustard, HealthTech and Medicines KTN  Angela Osborne, eXmoor pharma  Glyn Stacey - NIBSC and UKSCB
10.30 - 11.00	Cell banking challenges and solutions	Lyn Healy, NIBSC
11.00 - 11.30	Regulatory challenges and ensuring GMP compliance	Ian Rees, MHRA
11.30 - 12.00	<b>Coffee BREAK</b>	
12.00 - 12.30	Regulatory challenges and opportunities in moving manufacturing sites	Paul Kemp, Intercytex
12.30 - 13.00	Obtaining an IMP license – approach, challenges and solutions	Gill Lewis, eXmoor pharma
13.00 - 14.00	<b>LUNCH - including short tour of UK SCB labs in groups of 8</b>	
14.00 - 14.30	Outsourcing cell therapy product manufacture – opportunities and challenges	Kenny Pollock, ReNeuron
14.30 - 15.00	GMP issues in moving from research to manufacture - a case study	Sarah Moyle, CBF Oxford
15.00 - 15.30	The Biotherapeutics Development Unit: rationale, description and lessons learnt	Heike Lentfer, CR-UK
15.30	<b>Coffee – including tour of CR-UK BDU in groups of 16</b>	
16.30	<b>WRAP UP and CLOSE</b>	

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