The organisers wish to thank GlaxoSmithKline and AstraZeneca for their sponsorship of this meeting and the NMRDG for their help and support with the organisation.

AstraZeneca

NMRDG Spring Meeting

Advances in NMR on Small Amounts of Material

Thursday 26th Feb 2004

AstraZeneca R&D Charnwood, Loughborough, UK

9.45 Coffee

10.20 Introduction and Welcome (Mike Bernstein, AstraZeneca)

Session 1 Introduction to the Techniques

Chair: Richard Lewis (AstraZeneca)

- 10.30 Steve Coombes (Pfizer, Sandwich)

 Cryo-flow probes: Fixed and demountable flow cells
- 10.45 Martin Sandvoss (GSK)

 LC-SPE-NMR, separating the chromatography from the NMR
- 11.00 Dave Russell (Pfizer, Kalamazoo)

 Optimising one's sample for cryogenic NMR probes
- 11.15 Ian Jones (AstraZeneca R&D Macclesfield)

 Experience with the 1mm probe in a pharmaceutical development environment
- 11.30 Chris Sleigh (AstraZeneca R&D Charnwood)

 Evaluation of a CapNMR and CapLCNMR system
- 11.45 Paul Tan (Lilly)

 Automated sample preparation of small batches of compound

Buffet Lunch

Session 2 A Glimpse of the Future

Chair: Martin Sandvoss (GSK)

1.00 Klaus Albert (Institut für Organische Chemie, Universität Tübingen, Germany) LC-NMR – Present status and future

- 1.30 David Strand (Protasis)

 Dynamic Field Gradient Focusing: A new separation engine for NMR
- 1.45 Manfred Spraul (Bruker)

 Methods to increase NMR performance on small sample amounts
- 2.00 Iain Green (Varian Inc)
 Small molecules, small samples, big applications, a vendor perspective
- 2.15 Peter Meadows (Jeol)
 Who says NMR is Insensitive?

2.30 Coffee

Session 3 Industry Workflow

How can we make these technologies work in a high-throughput environment? *Chair: Adrian Davis (Pfizer)*

- 3.00 Mark O'Neil-Johnson, (Sequoia Sciences Inc., San Diego, CA)

 From HPLC to NMR structure elucidation, What can I do with twenty five micrograms?
- 3.25 Russell Mortishire-Smith (Merck Sharp and Dohme)
 Fish or cut bait? Workflow and metabolite identification by NMR
- 3.50 Gary Sharman (AstraZeneca R&D Macclesfield)
 Impurity and degradation product identification in pharmaceutical development

4.15 Discussion

Chair: Duncan Farrant (GSK)

4.45 Closing Remarks

Followed by informal discussion.