PiPS
Probiotic in Preterm babies Study

Site Training Log

Study Name: PiPS - A randomised controlled trial of Probiotics in Preterm Infants

Principal Investigator: _____ **Hospital:** _____

A Informed consent **B Randomisation** **C Stool sample collection** **D Data collection form completion**
E Intervention preparation **F All of the above** **G Other** *(Please specify - Insert additional responsibilities as required)*

It is the Principal Investigator's responsibility to ensure that every person delegated to perform a trial activity is suitably trained for the role they are given and that it is correctly recorded in both the Training and Delegation Logs. A training visit from a PiPS research nurse can be requested at any time by contacting the Trial Office or a local member of the PiPS trial team (who has already received the appropriate PiPS training) may train a member of staff in their proposed role. Training material can be found in the PiPS Documentation Box, Site and Pharmacy Files or on the PiPS website (www.npeu.ox.ac.uk/pips).

Trainee Name	Job Title	Training topic(s)	Trained by	Date: *(dd/mm/yy)*	Trainer Signature

Please continue overleaf

To be completed by the trainer

To be filed in the PiPS Site File

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To be completed by the trainer

To be filed in the PiPS Site File

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