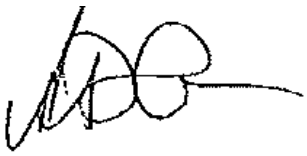
18th Nov 2013

To whom it may concern,

I hereby testify that the work carried out by David Fairlamb, in connection with Clinical Evaluations for certain Synergy Health products, was of a very satisfactory standard, in full compliance with the Medical Device Directive requirements.

David's work was consistently provided in timely fashion.

The evaluations in question covered quite a broad range of our product portfolio, including CE marked biocides, various procedure packs and plain & x-ray detectable gauzes & swabs. The classifications of such products ranged from Class I, non sterile to Class IIb, sterile.



Alexander Thomson

Quality Manager

Synergy Health

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