# EQUIPMENT ASSESSED AND EVALUATION CENTRE INFORMATION

1. DETAILS OF EQUIPMENT ASSESSED AND CENTRE

| | | |
|---|---|---|
| 1.1 | Equipment model | |
| 1.2 | Manufacturer | |
| 1.3 | Supplier | |
| 1.4 | Serial number(s) | |
| 1.5 | Evaluation centre | |
| 1.6 | Breast screening centre project leader (including telephone number) | |

2. INSTALLATION

| | | |
|---|---|---|
| 2.1 | Date of start of installation | |
| 2.2 | All adjustments made to suit local radiographic requirements by the installation engineer should be recorded. The engineer should confirm that all adjustments made conform with the manufacturer's installation protocol | |
| 2.2.1 | Adjustments to suit local radiographic requirements | |
| 2.2.2 | Comment by engineer on adjustments made | |
| 2.3 | Date of acceptance for clinical use | |
| 2.4 | Date of start of clinical evaluation | |
| 2.5 | Date of completion of clinical evaluation | |

3. FILM AND FILM HANDLING EQUIPMENT (Note: It is important that these are not changed during the evaluation period)

| | | |
|---|---|---|
| 3.1 | Manufacturer and type of film used during the evaluation | |
| 3.2 | Manufacturer and model of film processor | |
| 3.3 | Processing time dry to dry | |
| 3.4 | Developer temperature | |
| 3.5 | Manufacturer and type of processing chemicals | |
| 3.6 | Make and type of cassettes and screens used during the evaluation | |
| 3.7 | Total number of clinical films taken during the evaluation period | |
| 3.8 | Total number of sensitometry films taken during the evaluation period | |

4. NUMBER OF EXAMINATIONS UNDERTAKEN

| | | |
|---|---|---|
| 4.1 | Number of women screened | |
| 4.2 | Number of women assessed | |
| 4.3 | Number of women examined with magnification | |
| 4.4 | Number of stereotactic examinations | |