NEAT TRIAL — RANDOMISATION FORM

Patient Surname: ..... **Forename:** .....

Date of Birth: _____/_____/_____ **Hospital Number:** .....

Referring Consultant: ..... **Hospital:** .....

ELIGIBILITY CHECKLIST

1 Histological diagnosis of invasive breast cancer ☐
2. Clinically early stage disease ☐
3. Complete excision of tumour ☐
4. Definite indication for chemotherapy, in the opinion of the responsible clinician ☐
5. Patient is fit to receive treatment according to either of the study arms ☐
6. Patient has given written informed consent ☐
7. Patient has not received previous radiotherapy or chemotherapy ☐
8. Adequate renal, hepatic and bone marrow function ☐
9. No longer than 6 weeks has elapsed since surgery ☐
10. No previous cancer except basal cell carcinoma or carcinoma in situ ☐
11. Non-pregnant and no risk of pregnancy during chemotherapy ☐

INFORMATION ESSENTIAL FOR RANDOMISATION

Age: ≤ 50 ☐ > 50 ☐

Node -ve ☐ **Node +ve** → 1-3 ☐ ≥4 ☐

Radiotherapy to be given? No ☐

Yes ☐ → intend to randomise into **SECRAB** ☐

→ **with** CMF ☐ *OR* **after** CMF ☐

INFORMATION REQUESTED AT RANDOMISATION

Tumour size: ____.____ x ____.____ cm

Tumour grade: 1 (well diff.) ☐ 2 (mod. diff.) ☐ 3 (poorly diff.) ☐

ECOG Performance status ☐

How do you intend to give Cyclophosphamide? oral ☐ ***OR*** *i.v.* ☐

Date to start chemotherapy: ____/____/____

Menopausal status: Pre ☐ Post ☐ Peri ☐ Unknown ☐

Oestrogen Receptors: Neg ☐ Pos ☐ Unknown ☐

Is patient on, or to go on tamoxifen? No ☐ Yes ☐ Unknown ☐

If Yes: Will it be concurrent with chemotherapy? → Start Date ___/___/___

***Or*, after completion of chemotherapy?** → Proposed Start Date ___/___/___

Has patient agreed to complete Q of L questionnaires? Yes ☐ No ☐

TREATMENT ALLOCATION

Epirubicin + CMF ☐

Classical CMF ☐

TRIAL NUMBER ☐

Signed: ..... **Date of Randomisation : ____/____/___**