A multicenter phase II trial of neoadjuvant letrozole plus low-dose cyclophosphamide in postmenopausal patients with estrogen receptor-positive breast cancer (JBCRG-07): therapeutic efficacy and clinical implications of circulating endothelial cells

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Abstract
Neoadjuvant endocrine therapy has been reported to decrease tumor size, which leads to increased breast conservation rates. To improve the clinical response, metronomic chemotherapy with endocrine therapy is a promising strategy. A multicenter phase II single-arm neoadjuvant trial with letrozole and cyclophosphamide was conducted. Eligibility criteria included postmenopausal status, T2–4 N0–1, and estrogen receptor-positive breast carcinoma. Letrozole (2.5 mg) plus cyclophosphamide (50 mg) was given orally once a day for 24 weeks. The primary endpoint was the clinical response rate (CRR). To investigate anti-angiogenic effects, circulating endothelial cells (CECs) were quantified using the CellSearch system. From October 2007 to March 2010, 41 patients were enrolled. The CRR was 67.5% (52.0–80.0%), which was above the prespecified threshold (65%). The conversion rate from total mastectomy to breast-conserving surgery was 64% (18/28). Grade 3 or greater nonhematological toxicity was not reported. Clinical response was associated with improved disease-free survival (DFS) ($P = 0.020$). The increase in CEC counts at 8 weeks was observed in nonresponders ($P = 0.004$) but not in responders. Patients with higher CEC counts at baseline or post-treatment showed worse DFS than those with lower counts ($P < 0.001$ at baseline and $= 0.014$ post-treatment). Multivariate analysis showed that post-treatment CEC counts but not pretreatment counts were independently correlated with DFS ($P = 0.046$). In conclusion, neoadjuvant letrozole plus cyclophosphamide showed a good clinical response for postmenopausal patients with estrogen receptor-positive breast carcinoma. CEC quantification is a promising tool for treatment monitoring and prognostic stratification for metronomic therapy following validation of our results in larger studies. Clinical trial registration number: UMIN000001331 Phase II study of neoadjuvant letrozole combined with low-dose metronomic cyclophosphamide for postmenopausal women with estrogen-responsive breast cancer (JBCRG-07).
Neoadjuvant endocrine therapy (NET) is one of the treatment options for postmenopausal patients with endocrine-responsive breast cancer. NET has been reported to result in decreased tumor size and increased breast conservation rates [1–6]. Because endocrine therapy is associated with lower toxicity than chemotherapy, NET is preferable to neoadjuvant chemotherapy, especially in older patients and those with worsening performance status. In order to further improve surgical outcome, it is important to increase the NET response rate without increasing adverse effects. Chemo-endocrine therapy using metronomic chemotherapy is potentially useful in this regard.

Metronomic chemotherapy is the delivery of low doses of cytotoxic drugs at regular frequent intervals to avoid toxic side effects [7, 8]. It has been suggested to act via multiple mechanisms in exerting anticancer effects, including anti-angiogenesis, antitumor immune response, and direct anticancer action [9]. Oral cyclophosphamide is one of the most commonly used metronomic agents and is administered alone or together with other drugs such as capecitabine and methotrexate [10–13]. Because metronomic chemotherapy shows anticancer effects via different mechanisms of action without overt toxic side effects, it is a good candidate in combination with endocrine therapy.

The combined administration of the aromatase inhibitor letrozole with low-dose metronomic cyclophosphamide in elderly patients has been reported earlier [14]. This randomized phase II trial showed an overall response rate (ORR) of 87.7% in patients assigned to receive letrozole plus cyclophosphamide, while letrozole alone showed an ORR of 71.9%. In addition, post-treatment expression of Ki-67 was significantly lower in tumors treated with the combined therapy than in tumors treated with letrozole alone. Thus, the combination of letrozole and oral cyclophosphamide appears effective and promising as neoadjuvant therapy.

We conducted a multicenter phase II single-arm trial of neoadjuvant metronomic chemo-endocrine therapy with letrozole and oral cyclophosphamide in Japan (Japan Breast Cancer Research Group-07 trial: UMIN000001331). To investigate the possible role of anti-angiogenic effects in metronomic chemo-endocrine therapy, circulating endothelial cells (CECs) were quantified prior to and during the neoadjuvant treatment, and their association with treatment response and prognosis was examined.

Endpoints
The primary endpoint was the clinical response rate, assessed using calipers, ultrasound (US), or computed tomography (CT)/magnetic resonance imaging (MRI) during the 24-week neoadjuvant treatment period in the intention-to-treat (ITT) population. Tumor response was evaluated in accordance with RECIST ver. 1.0 [15]. Secondary endpoints included pathological therapeutic effects, breast conservation rate, safety assessed using CTCAE ver. 3.0, disease-free survival (DFS), and overall survival (OS).

The target number of patients in the protocol was set at 40 based on the response rate of 88% in a previous
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Pathological analyses

Pathological analyses were performed in a central laboratory. Tumor biopsy specimens before preoperative therapy were assessed for estrogen receptor, progesterone receptor (PgR), and human epidermal growth factor receptor type 2 (HER2). ER and PgR status were defined as positive for tumors with 10% or more positive tumor cells. HER2 positivity was determined as strong expression (3+) using immunohistochemistry or as HER2:CEP17 ratio >2.2 using fluorescence in situ hybridization [16]. The pathological response was assessed using surgical samples following preoperative therapy. A pathological complete response (pCR) was defined as no residual invasive tumor cells in the mammary gland and lymph nodes. Grade 2 response was defined as reduction in tumor cells by more than two-thirds (66%), and grade 1 was defined as reduction in tumor cells ≤ one-third (33%). The Ki-67 labeling index (LI) using the MIB1 antibody (Dako, Glostrup, Denmark) was calculated by counting positively stained tumor cells per 1000 tumor cells in the hot spots.

Circulating endothelial cells

Blood samples were drawn into CellSave tubes (Veridex, LLC, NJ) prior to, at 8 weeks after treatment initiation, and at completion of the neoadjuvant treatment. Samples were sent to the central laboratory at Kyoto University where they were processed within 72 h after blood sampling. All evaluations were performed without prior knowledge of the patients’ clinical status. The CellSearch system was used for endothelial cell detection, as described previously [17–19]. In brief, magnetic separation was performed using anti-CD146 ferrofluids, followed by labeling with the nuclear stain 4,6-diamidino-2-phenylindole (DAPI), a phycoerythrin-conjugated anti-CD105 antibody, and an allophycocyanin-conjugated anti-CD45 antibody. An additional channel was used for an anti-CD34 antibody conjugated to FITC (clone AC136, Miltenyi, Biotech GmbH, Germany). CECs were defined as CD146+ CD105+ CD45− DAPI− cells in this study. As CD34 is another maker that is positive in circulating endothelial cells, anti-CD34 antibody was added to the additional channel [20]. A gray-scale charge-coupled camera device was used to scan the entire chamber surface, and each captured frame was then evaluated for objects that were potential CEC candidates using image analysis software.

Statistical analysis

Baseline characteristics of patients were summarized as mean (range) for continuous variables and number (%) for categorical variables. The clinical/pathological response rate and the breast conservation rate were calculated at 95% confidence intervals (CIs). AEs during treatment were tabulated based on their CTCAE grades. OS and DFS during follow-up were estimated and compared using the Kaplan–Meier method and log-rank test between groups stratified based on patient characteristics and clinical/pathological outcomes of the neoadjuvant treatment. In biomarker analysis, association of CECs (as continuous variables) with clinical response was evaluated using univariate logistic regression models. The optimal cut-off value for each statistically significant biomarker to predict clinical response was determined using the Youden’s index of the receiver operating characteristics (ROC) curve. Patients were stratified based on the cut-off value into two groups, and the survival rate was compared between them. Multivariate survival analyses were performed using Cox proportional hazards models consisting of statistically significant variables from the survival analyses mentioned above. Multicollinearity was assessed using Spearman’s rank correlation coefficient. To address data sparseness, Firth’s penalized likelihood approach was applied in the regression analyses. A two-sided P-value below 0.05 was considered significant. Statistical analyses were performed using IBM SPSS Statistics 23.0 (IBM Corp., Armonk, NY) and R ver. 3.2.2 (R core team, R Foundation for Statistical Computing, Vienna, Austria).

Results

Population

From October 2007 to March 2010, 41 patients were enrolled in this study at four medical institutes in Japan (Fig. 1). One patient was excluded from the ITT population because of entry criteria violation (tumor size <2 cm). Six patients were further excluded from the per-protocol set (PPS) due to entry criteria violation in three patients (higher transaminase in one, age less than 60 years in two patients), changing hospitals during the protocol treatment in one patient, and insufficient duration (<90%) of drug administration in two patients. Baseline characteristics of the entire population (safety population), the ITT population, and the PPS population are shown in Table 1.

Clinical and pathological response

The clinical response rate in the ITT population was 67.5% (52.0–80.0%), which was above the prespecified threshold.
(65%). Associations between clinical response and baseline characteristics were assessed. No baseline characteristics including Ki-67 LI were associated with clinical response. Response rates in the HER2-negative and HER2-positive subgroups were 60% and 80%, respectively, which showed no statistically significant difference.

No patients achieved pCR. Seven patients (17.5%) showed grade 2 pathological responses and 28 (70%) showed grade 1 responses.

Changes in Ki-67 LI were assessed based on clinical responses (Fig. 2). Ki-67 LI decreased after treatment in both responders and nonresponders, and no difference in the decrease was observed based on clinical response.

Surgical outcome

Among patients in the ITT population, breast-conserving surgery was performed in 30 patients, and the breast-conserving rate was 75% (30/40). Before treatment, breast-conserving surgery and total mastectomy were anticipated in 12 and 28 patients, respectively. Eighteen patients who were anticipated to receive mastectomy before neoadjuvant treatment received breast-conserving surgery. The conversion rate from total mastectomy to conserving surgery was 64% (18/28).

Safety

Adverse events occurring in all enrolled patients are shown in Table 2. Twenty-two patients (54%) had leukocytopenia, but most (17/22) of them were grade 1. Grade 3 leukocytopenia was observed in one patient. The most common nonhematological AE was arthralgia, which was observed in six patients. One patient was diagnosed with liver cancer 3 months after initiation of the neoadjuvant treatment, and the causal relationship with treatment is unlikely. No

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**Table 1.** Baseline characteristics of patients.

| Patient characteristics | Total enrolled (Safety population) | ITT | PPS |
|-------------------------|----------------------------------|-----|-----|
| Number of patients      | 41                               | 40  | 34  |
| Age                     | Ave. (range)                      | 69.6 (57–82) | 69.9 (57–82) | 69.6 (61–80) |
| Staging                 | T                                 |     |     |     |
| T1                      | 1 (2.4)                           | 0 (0.0) | 0 (0.0) |
| T2                      | 36 (87.8)                         | 36 (90.0) | 31 (91.2) |
| T3                      | 4 (9.8)                           | 4 (10.0) | 3 (8.8) |
| N                       |                                   |     |     |     |
| N0                      | 36 (87.8)                         | 35 (87.5) | 29 (85.3) |
| N1                      | 5 (12.2)                          | 5 (12.5) | 5 (14.7) |
| Receptor status         | ER                                |     |     |     |
| +                       | 41 (100.0)                        | 40 (100.0) | 34 (100.0) |
| –                       | 0 (0.0)                           | 0 (0.0) | 0 (0.0) |
| PgR                     |                                   |     |     |     |
| +                       | 27 (65.9)                         | 26 (65.0) | 20 (58.8) |
| –                       | 14 (34.1)                         | 14 (35.0) | 14 (41.2) |
| HER2                    |                                   |     |     |     |
| +                       | 9 (22.0)                          | 9 (22.5) | 5 (14.7) |
| –                       | 32 (78.0)                         | 31 (77.5) | 29 (85.3) |
| Histological grade      | 1                                 |     |     |     |
| 1                       | 13 (31.7)                         | 13 (32.5) | 9 (26.5) |
| 2                       | 26 (63.4)                         | 25 (62.5) | 24 (70.6) |
| 3                       | 0 (0.0)                           | 0 (0.0) | 0 (0.0) |
| NA                      | 2 (4.9)                           | 2 (5.0) | 1 (2.9) |

ER, estrogen receptor; ITT, intention-to-treat; PPS, per-protocol set.
grade 3 or greater nonhematological toxicity was reported. No patients discontinued the treatment due to AE.

**Survival analysis**

Survival analyses were performed in the PPS. Among 34 patients, postoperative chemotherapy was given to seven patients. The median follow-up period was 68.5 months (range: 18.1–86.5).

DFS at 5 years was 90.9% (95% CI: 48.4–90.4%). Three patients relapsed during follow-up, one with axillary lymph node recurrence, one with chest wall recurrence, and one with lung metastasis. Overall survival at 5 years was 93.9% (95% CI: 74.4–97.0%). Two patients died during follow-up, one with liver cancer and the other with myocardial infarction 3 years after treatment initiation. Baseline factors including T stage, nodal involvement, HER2 status, and types of surgery were not associated with DFS.

Associations of survival with clinical response and AEs were evaluated. Clinical response with US was associated with prognosis; responders showed better DFS than nonresponders ($P = 0.020$) (Fig. 3). Interestingly, leukocytopenia was associated with prognosis; patients with no or mild leukocytopenia (G0 or 1) had better DFS than those

| Table 2. Adverse events. |
|--------------------------|
| Adverse event            | All grade | Grade $\geq$ 3 |
| Diarrhea                 | 1         | 0             |
| Dry mouth/salivary gland | 1         | 0             |
| Stomatitis               | 1         | 0             |
| Infection (cold sore)    | 1         | 0             |
| Periodontal disease      | 1         | 0             |
| Epigastric distress      | 1         | 0             |
| Nausea                   | 1         | 0             |
| Anorexia                 | 3         | 0             |
| Liver cancer             | 1         | 0             |
| Cystitis                 | 2         | 0             |
| Osteoporosis             | 2         | 0             |
| Joint function: hand and finger joint stiffness | 2 | 0 |
| Pain-arthralgia          | 6         | 0             |
| Pain-myalgia             | 1         | 0             |
| Pain-headache            | 1         | 0             |
| Perspiration             | 3         | 0             |
| Postmenopausal syndrome  | 3         | 0             |
| (headache, dizziness)    |           |               |
| Dizziness                | 1         | 0             |
| Leukocytopenia           | 22        | 1             |
| Thrombocytopenia         | 5         | 0             |
| Anemia                   | 7         | 0             |
| Number of patients: 41   |           |               |
with severe leukocytopenia (G2 or 3) \((P = 0.003)\) (Fig. 3). No other factors including pre- and post-treatment Ki-67 LI, changes in Ki-67 LI, or HER2 status were associated with DFS or OS.

**Circulating endothelial cells**

Circulating endothelial cells were quantified prior to and during the neoadjuvant therapy. Their association with clinical response with US and prognosis were evaluated.

In nonresponders, CEC counts were significantly increased at 8 weeks \((P = 0.004)\) compared with pretreatment counts, while in responders, no such increases were observed \((P = 0.35)\) (Fig. 4). Similarly, CD34-positive CEC counts were increased at 8 weeks in nonresponders \((P = 0.003)\) but not in responders \((P = 0.39)\). Baseline counts of CEC and CD34-positive CEC did not correlate with treatment response.

The association between CEC counts and prognosis was evaluated. Cut-off values for CEC and CD34-positive CEC were determined using the Youden’s index of ROC curves. Baseline counts of CEC and CD34-positive CEC were significantly associated with DFS, and patients with higher counts of CEC and CD34-positive CEC showed worse prognosis than those with lower counts \((P < 0.001\) and \(P = 0.004\), respectively) (Fig. 5A). In addition, post-treatment counts of CEC and CD34-positive CEC were also significantly correlated with DFS \((P = 0.014\) and \(P = 0.008\), respectively) (Fig. 5B).

Because clinical response and leukocytopenia were also associated with DFS, multivariate analyses of DFS, including clinical response, leukocytopenia, and pre- and post-treatment counts of CEC, were performed. Interestingly, post-treatment counts of CEC, but not pretreatment counts, were independently correlated with DFS \((P = 0.046)\) (Table 3). A similar result was observed for CD34-positive CEC \((P = 0.043)\) (Table 3).

**Discussion**

In this study, we demonstrated that neoadjuvant metronomic chemo-endocrine therapy with letrozole and cyclophosphamide showed a good response in Japanese postmenopausal women with ER-positive breast cancer, with a conversion rate from mastectomy to breast-conserving surgery of 64% and tolerable toxicity. In addition, increases in CEC counts at week 8 indicated poor response, and post-treatment CEC counts showed a good and independent prognostic value.

One of the advantages of neoadjuvant treatment is an increase in breast-conserving rate. The IMPAKT trial, which compared anastrozole, tamoxifen, and both in combination in the neoadjuvant setting, showed that the conversion rates from mastectomy to breast-conserving surgery were 44%, 31%, and 24%, respectively [6]. Thus, the conversion rate achieved in this study (64%) was higher than the rate in any endocrine treatment group of the IMPAKT trial. Another neoadjuvant endocrine study, the PROACT trial, compared anastrozole and tamoxifen [1]. Although the conversion rate was not reported, the improvement in surgery including the conversion from mastectomy to breast-conserving surgery was observed in 38.1% and 29.9% of the patients who received anastrozole and tamoxifen, respectively. Altogether, these results suggest that the combination of letrozole and cyclophosphamide would give a higher conversion rate than endocrine therapy alone.

In nonresponders, CEC counts increased at week 8, while in responders, such an increase was not observed. In our previous study, we showed that CEC counts done with the CellSearch system increased during neoadjuvant

![Figure 3](image-url) Disease-free survival according to clinical response and leukocytopenia. Better clinical response and milder leukocytopenia were associated with a better disease-free survival \((P = 0.020\) and \(0.003\), respectively).
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Chemotherapy, especially during therapy involving taxane-based regimens [17]. Such increases have been suggested to contribute to angiogenesis and neovascularization in order to repair damaged tissues, including normal and cancerous tissues [21–23]. Metronomic chemotherapy is expected to prevent such a vascular rebound in neovascularization, especially in tumor tissues, which is one of the suggested mechanisms for its anticancer effect. Therefore, it is conceivable that prevention of neovascularization due to metronomic chemotherapy led to maintained CEC counts in responders, while failure of such prevention resulted in increased CEC counts in nonresponders.

In our study, although CEC counts at both baseline and post-treatment showed prognostic value, only post-treatment CEC counts had independent prognostic power. Poor prognosis in patients with high post-treatment CEC counts may be a result of insufficient anti-angiogenic response with metronomic chemo-endocrine therapy. This result seems consistent with the prognostic value of post-treatment Ki-67 LI in NET, which showed better prognostic power than pretreatment Ki-67 LI [24–27]. Our results along with other reports suggest that biological responses, such as the antiproliferative response indicated by Ki-67 LI and anti-angiogenic response indicated by CEC counts after metronomic therapy, show more precise prognostic value than the baseline biology of tumors.

**Figure 4.** Change in circulating endothelial cell count and clinical response. Nonresponders showed an increase in circulating endothelial cell count at 8 weeks \((P = 0.004)\), while responders did not \((P = 0.35)\).

**Figure 5.** (A) Disease-free survival according to pretreatment CEC and CD34-positive CEC count. Higher counts of CEC and CD34-positive CEC showed a worse prognosis than lower counts \((P < 0.001\) and \(= 0.004\), respectively). (B) Disease-free survival according to post-treatment CEC and CD34-positive CEC count. Higher counts of post-treatment CEC and CD34-positive CEC showed a worse prognosis \((P = 0.014\) and \(0.008\), respectively).
Leukocytopenia was associated with prognosis in this study. Severe leukocytopenia (G2 or G3) was associated with worse DFS. This seems contradictory to results reported with conventional chemotherapy in adjuvant settings for early-stage breast cancer [28–30]. These previous studies indicated that severe myelosuppression was associated with better prognosis in patients with breast cancer receiving CMF (cyclophosphamide, methotrexate, and 5-fluorouracil) or CAF (cyclophosphamide, doxorubicin, and 5-fluorouracil), suggesting that hematological toxicity due to conventional chemotherapy may represent biological activity of the drugs, resulting in improved prognosis. However, metronomic chemotherapy has been suggested to exert anticancer effects via different mechanisms of action compared to conventional chemotherapy, one of which is activation of antitumor immune response. Indeed, low-dose cyclophosphamide has been implicated in activation of innate immunity [31–33]. Therefore, myelosuppression during metronomic treatment may lead to insufficient immune activation, which might result in poor treatment efficacy in patients with severe leukocytopenia.

Although the objective response rate (67.5%) in our study appears a little lower than that (87.7%) in a previous report by Bottini et al. [14], some differences exist between the two studies. Bottini’s study included only elderly patients, and thus, median patient age in our study was lower in comparison. More than half of the patients in Bottini’s study had histological grade 3 tumors, while none of the patients in our study had grade 3 tumors. The clinical response was assessed using calipers in Bottini’s study, while it was assessed with calipers, US, and CT/MRI in our study. These differences might have contributed to different response rates in the two studies.

This study was limited in terms of some parameters. One of its biggest limitations was its small sample size. Because this was a phase II trial investigating clinical efficacy and tolerability of combined treatment with letrozole and low-dose cyclophosphamide, the sample size was set at 40. In order to validate the clinical utility of the treatment, a larger study is warranted. It is also important to interpret the results including the prognostic analysis with this sample size cautiously. To confirm the prognostic value of CEC, it is necessary to conduct a larger study in which CECs are serially measured. The definition of ER and PgR positivity is another issue. In 2010, the American Society of Clinical Oncology/College of American Pathologists recommended that ER and PgR assays be considered positive if there are at least 1% positive tumor nuclei in the sample on testing with appropriate controls [34]. Because this study started in 2007, the old criteria of ER and PgR were used. Thus, future studies to validate our results should be conducted with the new definition of ER and PgR positivity. Another limitation was that this was a single-arm study in which chemo-endocrine therapy was not compared with either endocrine therapy alone or metronomic chemotherapy alone. It is, therefore, not clear whether combined administration of letrozole with metronomic cyclophosphamide resulted in a better outcome than letrozole or cyclophosphamide alone would have in this population. A randomized controlled study would be required for such a comparison in a larger confirmative study.

In conclusion, metronomic chemo-endocrine therapy with letrozole plus cyclophosphamide showed a good response and was tolerated in Japanese postmenopausal patients with ER-positive breast cancer. An increase in CEC counts during the treatment was associated with poor response, and post-treatment CEC counts as well as clinical response were independent prognostic factors. The combination of letrozole and cyclophosphamide could be an option for postmenopausal women with ER-positive breast cancer. CEC quantification would be a promising tool for treatment monitoring and prognostic stratification following validation of our results in larger prospective studies.

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Conflict of Interest

T Ueno has received honoraria from Chugai Pharmaceutical Co., Ltd., Eisai Co., Ltd., Novartis Pharma K.K. N Masuda has received honoraria from Chugai Pharmaceutical Co., Ltd., Astra Zeneca K.K. S Morita has received honoraria from Chugai Pharmaceutical Co., Ltd. M Toi has received research funding from Taiho Pharmaceutical Co., Ltd., Chugai Pharmaceutical Co., Ltd., Eisai Co., Ltd., Shimadzu Corporation, C & C, Japan Breast Cancer Research Group, Astra Zeneca K.K., AFI Technology, Daiichi-Sankyo Co., Ltd.

References

1. Cataliotti, L., A. U. Buzdar, S. Noguchi, J. Bines, Y. Takatsuka, K. Petrakova, et al. 2006. Comparison of anastrozole versus tamoxifen as preoperative therapy in postmenopausal women with hormone receptor-positive breast cancer: the Pre-Operative “Arimidex” Compared to Tamoxifen (PROACT) trial. Cancer 106:2095–2103.

2. Ellis, M. J., and C. Ma. 2007. Letrozole in the neoadjuvant setting: the P024 trial. Breast Cancer Res. Treat. 105(Suppl. 1):33–43.

3. Ellis, M. J., V. J. Suman, J. Hoog, L. Lin, J. Snider, A. Prat, et al. 2011. randomized phase II neoadjuvant comparison between letrozole, anastrozole, and exemestane for postmenopausal women with estrogen receptor-rich stage 2 to 3 breast cancer: clinical and biomarker outcomes and predictive value of the baseline PAM50-based intrinsic subtype–ACOSOG Z1031. J. Clin. Oncol. 29:2342–2349.

4. Toi, M., S. Saji, N. Masuda, K. Kuroi, N. Sato, H. Takei, et al. 2011. Ki67 index changes, pathological response and clinical benefits in primary breast cancer patients treated with 24 weeks of aromatase inhibition. Cancer Sci. 102:858–865.

5. Masuda, N., Y. Sagara, T. Kinoshita, H. Iwata, S. Nakamura, Y. Yanagita, et al. 2012. Neoadjuvant anastrozole versus tamoxifen in patients receiving goserelin for premenopausal breast cancer (STAGE): a double-blind, randomised phase 3 trial. Lancet Oncol. 13:345–352.

6. Smith, I. E., M. Dowsett, S. R. Ebbs, J. M. Dixon, A. Skene, J. U. Blohmer, et al. 2005. Neoadjuvant treatment of postmenopausal breast cancer with anastrozole, tamoxifen, or both in combination: the Immediate Preoperative Anastrozole, Tamoxifen, or Combined with Tamoxifen (IMPACT) multicenter double-blind randomized trial. J. Clin. Oncol. 23:5108–5116.

7. Browder, T., C. E. Butterfield, B. M. Kraling, B. Shi, B. Marshall, M. S. O’Reilly, et al. 2000. Antiangiogenic scheduling of chemotherapy improves efficacy against experimental drug-resistant cancer. Cancer Res. 60:1878–1886.

8. Klement, G., S. Baruchel, J. Rak, S. Man, K. Clark, D. J. Hicklin, et al. 2000. Continuous low-dose therapy with vinblastine and VEGF receptor-2 antibody induces sustained tumor regression without overt toxicity. J. Clin. Invest. 105:R15–R24.

9. Andre, N., M. Carre, and E. Pasquier. 2014. Metronomics: towards personalized chemotherapy? Nat. Rev. Clin. Oncol. 11:413–431.

10. Bojko, P., G. Schimmel, D. Bosse, and W. Abenhardt. 2012. Metronomic oral cyclophosphamide in patients with advanced solid tumors. Onkologie 35:35–38.

11. Gebbia, V., H. Boussen, and M. R. Valerio. 2012. Oral metronomic cyclophosphamide with and without methotrexate as palliative treatment for patients with metastatic breast carcinoma. Anticancer Res. 32:529–536.

12. Wang, Z., J. Lu, S. Leaw, X. Hong, J. Wang, Z. Shao, et al. 2012. An all-oral combination of metronomic cyclophosphamide plus capecitabine in patients with anthracycline- and taxane-pretreated metastatic breast cancer: a phase II study. Cancer Chemother. Pharmacol. 69:515–522.

13. Yoshimoto, M., S. Takao, M. Hirata, Y. Okamoto, S. Yamashita, Y. Kawaguchi, et al. 2012. Metronomic oral combination chemotherapy with capecitabine and cyclophosphamide: a phase II study in patients with HER2-negative metastatic breast cancer. Cancer Chemother. Pharmacol. 70:331–338.

14. Bottini, A., D. Generali, M. P. Brizzi, S. B. Fox, A. Bersiga, S. Bonardi, et al. 2006. Randomized phase II trial of letrozole and letrozole plus low-dose metronomic oral cyclophosphamide as primary systemic treatment in elderly breast cancer patients. J. Clin. Oncol. 24:3623–3628.

15. Eisenhauer, E. A., P. Therasse, J. Bogaerts, L. H. Schwartz, D. Sargent, R. Ford, et al. 2009. New response evaluation criteria in solid tumours: revised RECIST guideline (version 1.1). Eur. J. Cancer 45:228–247.

16. Wolff, A. C., M. E. Hammond, J. N. Schwartz, K. L. Hagerty, D. C. Allred, R. J. Cote, et al. 2007. American Society of Clinical Oncology/College of American Pathologists guideline recommendations for human epidermal growth factor receptor 2 testing in breast cancer. J. Clin. Oncol. 25:118–145.
17. Ali, A. M., T. Ueno, S. Tanaka, M. Takada, H. Ishiguro, A. Z. Abdellah, et al. 2011. Determining circulating endothelial cells using Cell Search system during preoperative systemic chemotherapy in breast cancer patients. Eur. J. Cancer 47:2265–2272.

18. Johnston, S., J. Jr Pippen, X. Pivot, M. Lichinitser, S. Sadeghi, V. Dieras, et al. 2009. Lapatinib combined with letrozole versus letrozole and placebo as first-line therapy for postmenopausal hormone receptor-positive metastatic breast cancer. J. Clin. Oncol. 27:5538–5546.

19. Di Leo, A., G. Jerusalem, L. Petruzelka, R. Torres, I. N. Bondarenko, R. Khasanov, et al. 2010. Results of the CONFIRM phase III trial comparing fulvestrant 250 mg with fulvestrant 500 mg in postmenopausal women with estrogen receptor-positive advanced breast cancer. J. Clin. Oncol. 28:4594–4600.

20. Bertolini, F., Y. Shaked, P. Mancuso, and R. S. Kerbel. 2006. The multifaceted circulating endothelial cell in cancer: towards marker and target identification. Nat. Rev. Cancer 6:835–845.

21. Bertolini, F., S. Paul, P. Mancuso, S. Monestiroli, A. Gobbi, Y. Shaked, et al. 2003. Maximum tolerable dose and low-dose metronomic chemotherapy have opposite effects on the mobilization and viability of circulating endothelial progenitor cells. Cancer Res. 63:4342–4346.

22. Shaked, Y., A. Ciarrocchi, M. Franco, C. R. Lee, S. Man, A. M. Cheung, et al. 2006. Therapy-induced acute recruitment of circulating endothelial progenitor cells to tumors. Science 313:1785–1787.

23. Shaked, Y., E. Henke, J. M. Roodhart, P. Mancuso, M. H. Langenberg, M. Colleoni, et al. 2008. Rapid chemotherapy-induced acute endothelial progenitor cell mobilization: implications for antiangiogenic drugs as chemosensitizing agents. Cancer Cell 14:263–273.

24. Chang, J., T. J. Powles, D. C. Allred, S. E. Ashley, A. Makris, R. K. Gregory, et al. 2000. Prediction of clinical outcome from primary tamoxifen by expression of biologic markers in breast cancer patients. Clin. Cancer Res. 6:616–621.

25. Dowsett, M., I. E. Smith, S. R. Ebbs, J. M. Dixon, A. Skene, R. A’Hern, et al. 2007. Prognostic value of Ki67 expression after short-term presurgical endocrine therapy for primary breast cancer. J. Natl Cancer Inst. 99:167–170.

26. Ellis, M. J., Y. Tao, J. Luo, R. A’Hern, D. B. Evans, A. S. Bhatnagar, et al. 2008. Outcome prediction for estrogen receptor-positive breast cancer based on postneoadjuvant endocrine therapy tumor characteristics. J. Natl Cancer Inst. 100:1380–1388.

27. Kenny, F. S., P. C. Willsher, J. M. Gee, R. Nicholson, S. E. Pinder, I. O. Ellis, et al. 2001. Change in expression of ER, bcl-2 and MIB1 on primary tamoxifen and relation to response in ER positive breast cancer. Breast Cancer Res. Treat. 65:135–144.

28. Cameron, D. A., C. Massie, G. Kerr, and R. C. Leonard. 2003. Moderate neutropenia with adjuvant CMF confers improved survival in early breast cancer. Br. J. Cancer 89:1837–1842.

29. Mayers, C., T. Panzarella, and I. F. Tannock. 2001. Analysis of the prognostic effects of inclusion in a clinical trial and of myelosuppression on survival after adjuvant chemotherapy for breast carcinoma. Cancer 91:2246–2257.

30. Saarto, T., C. Blomqvist, P. Rissanen, A. Auvinen, and I. Elomaa. 1999. Haematological toxicity: a marker of adjuvant chemotherapy efficacy in stage II and III breast cancer. Br. J. Cancer 75:301–305.

31. Chen, C. S., J. C. Doloff, and D. J. Waxman. 2014. Intermittent metronomic drug schedule is essential for activating antitumor innate immunity and tumor xenograft regression. Neoplasia 16:84–96.

32. Doloff, J. C., C. S. Chen, and D. J. Waxman. 2014. Anti-tumor innate immunity activated by intermittent metronomic cyclophosphamide treatment of 9L brain tumor xenografts is preserved by anti-angiogenic drugs that spare VEGF receptor 2. Mol. Cancer. 13:158.

33. Wu, J., and D. J. Waxman. 2014. Metronomic cyclophosphamide schedule-dependence of innate immune cell recruitment and tumor regression in an implanted glioma model. Cancer Lett. 353:272–280.

34. Hammond, M. E., D. F. Hayes, M. Dowsett, D. C. Allred, K. L. Hagerty, S. Badve, et al. 2010. American Society of Clinical Oncology/College Of American Pathologists guideline recommendations for immunohistochemical testing of estrogen and progesterone receptors in breast cancer. J. Clin. Oncol. 28:2784–2795.