**Low dose metronomic weekly Docetaxelin previously treated patients with non-small cell lung cancer**

Emad Sadaka and Walid Almorsy

Clinical Oncology Department, Faculty of Medicine, Tanta University, Gharbia, Egypt.

e\_sadaka@hotmail.com,walidaa1@hotmail.com

**Abstract: Background:** Low dose metronomic chemotherapy (LDM) involves administering cytotoxic drugs on a daily or weekly basis at low doses without a long interval of interruption which can modulate the cancer microenvironment and disrupt tumor-associated vascular angiogenesis. The aim of this study is to evaluate the toxicity and efficacy of low dose weekly metronomic docetaxel in previously treated patients with non-small cell lung cancer (NSCLC). **Patients and methods:**Twenty three (23) patients were treated at Tanta University Hospital, Clinical Oncology Department. Patients received weekly 15 mg/m2 of docetaxel intravenously. **Results:** The median age was 59 years. The median number of cycles administered was 15 (range: 5–37). No patients achieved CR; 2 patients showed PR (8.7%); 10 patients had SD (43.5%) with clinical benefit (52.2%) and 11(47.8%) patients showed progressive disease. The median OS time was 10.5 months. The one year survival rate was 43.5%. The median progression-free survival time (PFS) was 4.5 months ranged from 1.5-14 months (95% CI: 2.7–6.3) and the one year PFS rate was 13%. Low dose metronomic docetaxel was well tolerated where no patients experienced grade 4 toxicities and only 2 (8.7%) patients had grade 3 anemia. No patients had high grade (3-4) non-haematological toxicities. **Conclusion:** This study suggested that metronomic low dose weekly Docetaxel was well tolerated and active in patients with previously treated NSCLC.Thus, further investigation of this LDM regimen with larger number of patients is warranted.

[Emad Sadaka and Walid Almorsy.**Low dose metronomic weekly Docetaxel in previously treated patients with non-small cell lung cancer.** *Cancer Biology* 2016;6(2):51-55]. ISSN: 2150-1041 (print); ISSN: 2150-105X (online). <http://www.cancerbio.net>.7. doi:[10.7537/marscbj06021607](http://www.dx.doi.org/10.7537/marscbj06021607).

**Key words**: NSCLC,Low dose metronomic chemotherapy, Taxotere.

**1. Introduction**

Conventional chemotherapy with high doses of a cytotoxic agents followed by a 3-4 weeks rest period is important to balance the drug’s toxic effects on the tumor and on the host. However, recent studies have demonstrated a rapid chemotherapy resistance in certain patient populations after an initial response. **(1)**Also, conventional chemotherapy drugs have been associated with many toxic adverse effects that often impair quality of life and limit continuation of therapy.(2)

A metronomic chemotherapy is characterized by frequent (dose-dense) chemotherapy administration with no interruptions without using the maximal tolerated dose (MTD), no administration of hematopoietic growth factors, preference for oral drugs with low incidence of treatment related toxicity.(3)

The progress of chemotherapy over the past few decades has resulted in survival benefit for patients with advanced non-small cell lung cancer (NSCLC) who have poor prognosis and the use of low-dose metronomic (LDM) chemotherapy might have an antitumor effects. Docetaxel has been recognized as an active drug that can exhibit metronomic activity both in vitro and in vivo.(4, 5, 6)

Yokoi et al*.* (2012) evaluated the toxicity and efficacy of LDM chemotherapy with weekly docetaxel in patients with non-small cell lung cancer (NSCLC). Twenty-seven patients received 15 mg/m2 of docetaxel on a weekly basis. Their results indicate that metronomic docetaxel was well tolerated and active in patients with NSCLC. (7)

Joshi et al*.* (2013) evaluated patients with recurrent and advanced NSCLC were treated with weekly paclitaxel at 80 mg/m 2. They concluded that weekly low-dose continuous metronomic-type scheduling of paclitaxel is safe and effective for those patients. (8)

**2. Patients and methods:**

This study was carried out at Clinical Oncology Department, Tanta University Hospital, between May 2013 and November 2015. Twenty three (23) patients with histologically or cytologically diagnosed NSCLC (clinical stage IIIB or IV at the time of enrollment). Inclusion criteria were: Performance Status (PS) 0–1 according to Eastern Cooperative Oncology Group (ECOG), patients had previously received at least one regimen of platinum based systemic chemotherapy with previous paclitaxel use allowed, adequate baseline bone marrow function (absolute neutrophil count ≥ 1500/ml, platelet count ≥ 100.000/ml and haemoglobin ≥8 gm/dl), adequate renal function (creatinine ≤ 1.5 mg/dl), adequate baseline hepatic function (serum bilirubin ≤ 2.0 mg/d, transaminase aspartate aminotransferase, alanine transaminase ≤ 2.0 the upper limit of normal level).

Docetaxel was administered intravenously (15 mg/m2) diluted with 5% glucose (250 mL) over 60 minutes each week without interval. Treatment was continued as long as there is no disease progression or unacceptable toxicity (e.g., grade3 or worse hematological/ non-hematological toxicity, not appetite loss, constipation, and nausea/vomiting). Treatment response was evaluated using chest CT scan every 8 weeks during the treatment period and when indicated.

**Statistical analysis:**

Version 21 of SPSS software was used. The OS was defined as the time from the start of the LDM chemotherapy to the time of death from any cause or to the date of last visit. At same time, PFS was estimated from date of start of metronomic chemotherapy till the date of progression and/or change of therapy due to any reason (e.g. toxicity, patient request) or death. Treatment-related adverse effects were evaluated using National Cancer Institute Common Toxicity Criteria, version 2.0.9. (9) Objective tumor response was evaluated using the Response Evaluation Criteria in Solid Tumors (RECIST), version 1.0.10 (10). The OS was estimated with the Kaplan–Meier product method.

**3. Results:**

Twenty three (23) patients were evaluated in this study from May 2013 to November 2015. All patients were treated at Tanta University Hospital, Clinical Oncology Department. Table (1) showed the patients’ characteristics. The median age was 59 years (range: 46 –71).Nineteen patients (82.6%) were males and 5 patients (17.4%) were females. Adenocarcinoma and squamous cell carcinoma were presented in 15 (65.2%) and 8 patients (34.8%), respectively. Stage IIIB presented in 8 patients (34.8%), while 15 patients (65.2%) had stage IV disease. The performance status (PS) according to Eastern Cooperative Oncology Group (ECOG) was 1 in 6 patients (26.1%) and 2 in 17 patients (73.9%). Eighteen patients received chemotherapy alone and 5 patients received concomitant chemotherapy and thoracic irradiation (CCRT). Seven patients received one chemotherapy regimen and 16 patients received two regimens. The median number of cycles administered was 15 (range: 5–37). Treatment was discontinued in patients developed disease progression, unacceptable toxicity, deterioration of PS, and/or patient refusal to continue the treatment. These patients offered the best supportive care.

|  |
| --- |
| **Table (1): Patients characteristics** |
| **Characters** | **No (%)** |
| **Age (years)** | MedianRangeMean ±SD | 5946-7159.09±6.9 |
| **Gender** | MaleFemale | 19 (82.6%)4 (17.4%) |
| **Performance Status (ECOG)** | 12 | 6 (26.1%)17 (73.9%) |
| **Pathology** | AdenocarcinomaSCC | 15 (65.2%)8 (34.8%) |
| **Tumor Stage** | IIIbIV | 8 (34.8%)15 (65.2%) |
| **Previous Treatment** | ChemotherapyCCRT | 18 (78.3%)5 (21.7%) |
| **No. of prior CTh regimens** | 12 | 7 (30.4%)16 (69.6%) |
| **No. of administered cycles** | TotalMedianRange | 411155-37 |
| SSC: Squamous cell carcinoma; CCRT: Concurrent chemo-radiation therapy |
|  |

Table (2) showed the treatment response where no patients achieved CR; 2 patients showed PR (8.7%); 10 patients had SD (43.5%); and 11 patients (47.8%)showed progressive disease. Seventeen patients died, and 6 patients were still alive at end of the study. The Kaplan–Meier survival curve of OS is shown in Figure 1. The median OS time was 10.5 months (95% CI: 8.2–12.8). The one year survival rate was 43.5%. The median progression-free survival time (PFS) was 4.5 months ranged from 1.5-14 months (95% CI: 2.7–6.3). The one year PFS rate was 13%. At the end of the study, 19 (82.6%) patients had disease progression while 4 (17.4%) patients were free of progressive disease (2 patients with SD and 2 patients with PR). Regarding treatment related haematological toxicities, table (3) showed that low dose metronomic docetaxel was well tolerated where no patients experienced grade 4 toxicities and only 2 (8.7%) patients had grade 3 anemia. No patients had high grade (3-4) non haematological toxicities (nausea, vomiting, diarrhea, hepatic, renal, and neurotoxicity).

|  |
| --- |
| **Table (2): Response to treatment.** |
| **Response****--------------------------------**Complete Response | **No (%)****---------------**0 (0%) |
| Partial Response (PR) | 2 (8.7%) |
| Stable Disease (SD) | 10 (43.5%) |
| Progressive Disease (PD) | 11 (47.8%) |

|  |
| --- |
| **Table (3): Treatment-related toxicities** |
| **Toxicity** | **Grade I****No (%)** | **Grade II****No (%)** | **Grade III****No (%)** | **Grade IV****No (%)** |
| Anemia | 14 (60.9%) | 7 (30.4%) | 2 (8.7%) | **-** |
| Neutropenia | 2 (8.7%) | 3 (13.1%) | - | **-** |
| Thrombocytopenia | 2 (8.7%) | 3 (13.1%) | - | **-** |
| Nausea | 5 (21.7%) | 1 (4.3%) | - | **-** |
| Vomiting | 4 (17.4%) | 1 (4.3%) | - | **-** |
| Diarrhea | 3 (13.1%) | 1 (4.3%) | - | **-** |
| Hepatic | 13 (56.5%) | 2 (8.7%) | - | **-** |
| Renal | 3 (13.1%) | 1 (4.3%) | - | **-** |
| Neuropathy | 4 (17.4%) | 1 (4.3%) |  |  |

|  |  |
| --- | --- |
| 1**Fig (1): Overall survival for all patients** | 2**Fig (2): Progression free survival for all patients** |

**4. Discussion:**

Low dose metronomic chemotherapy is an emerging form of chemotherapy with distinct mechanisms of action from conventional chemotherapy (e.g., antiangiogenesis). Although developed to overcome resistance to conventional chemotherapy, metronomic chemotherapy is subject to resistance on its own. Metronomic chemotherapy may overcome chemotherapy resistance and control tumors with low side effects.(11-15)

Docetaxel as a metronomic chemotherapy has also been shown to decrease the micro vessel density of tumors and inhibit the mobilization of circulating endothelial precursors. Thus, these findings have been the basis for several clinical studies on the use of LDM chemotherapy for NSCLC treatment.(16, 17)

Among 23 patients were evaluated in our study, No patients achieved CR; 2(8.7%) patients showed PR; 10 (43.5%) patients had SD; and 11(47.8%) patients showed progressive disease. Seventeen patients died, and 6 patients were still alive at end of the study. The median OS time was 10.5 months (95% CI: 8.2–12.8) and the one year survival rate was 43.5%. The median progression-free survival time (PFS) was 4.5 months ranged from 1.5-14 months (95% CI: 2.7–6.3). The one year PFS rate was 13%. At the end of the study, 19 (82.6%) patients had disease progression while 4 (17.4%) patients were free of progressive disease (2 patients with SD and 2 patients with PR).

Regarding treatment related haematological toxicities, the low dose metronomic docetaxel was well tolerated where no patients experienced grade 4 toxicities and only 2 (8.7%) patients had grade 3 anemia. No patients had high grade (3-4) non haematological toxicities (nausea, vomiting, diarrhea, hepatic, renal, and nurotoxicity).

Joshi et al*.* (2013) evaluated the administration of weekly paclitaxel in relapsed and refractory NSCLC or in patients not eligible for platinum-based chemotherapy. The complete remission was (2.7%), partial remission (32.4%), stable disease (32.4%), and progressive disease (27%). The median PFS was four months, and the estimated median OS was seven months. Paclitaxel was well tolerated. The most frequent grade 3 toxicities included anemia in 8%, neutropenia in 5.4%, and sensory neuropathy in 8%.They concluded that low dose continuous weekly paclitaxel is safe and effective in recurrent and refractory NSCLC. (8)

Yokoiet al*.* (2012) evaluated LDM chemotherapy with weekly low-dose docetaxel for previously treated twenty-seven patients diagnosed with non-small cell lung cancer (NSCLC). Patients received 15 mg/m2 of docetaxel intravenously weekly without interval. Eleven patients were stage IIIB, and 16 patients were stage IV. The performance status was 0 or 1 according to eastern Cooperative Oncology Group. There was no severe hematological adverse effect (neutropenia and/or thrombocytopenia). One patient (3.7%) showed complete remission, one patient (3.7%) shoed partial response, 12 patients (44.4%) had stable disease and 13 patients (48.1%) had progressive disease. The median survival time was 16.4 months and one year survival rate was 58.8%. The median progression free survival time was 6.5 months. Their results suggested that metronomic docetaxel was active and well tolerated in patients with NSCLC. (7)

Gorn *et al.*(2008) studied21 patients with stage IV disease NSCLC to evaluate the use of docetaxel and oral trofosfamide as a metronomic second line treatment of patients with metastatic non-small cell lung cancer (NSCLC). Patients received docetaxel 25 mg/m2 on days 1, 8, and 15 every 4 weeks plus trofosfamide 50 mg per day. The overall response rate was 19%, median overall survival was 6.9 months, the median progression-free survival 2.9 months, the one year survival rate was 28.6%, and the 2-year survival rate 7.1%. No grade IV toxicity was observed. They suggested that, the combination of docetaxel and trofosfamide as a metronomic regimen is active and well tolerable in patients with metastatic NSCLC. (16)

**Conclusion:**

This study suggested that in patients with previously treated NSCLC, metronomic low dose weekly Docetaxel showed activity and was well tolerated.Thus, LDM regimen may be considered as an alternative choice in those patients. Further investigation with larger number of patients is warranted.

**Corresponding Author**

EmadSadaka

e\_sadaka@hotmail.com

**References**

1. Agarwal R, Kaye SB. Ovarian cancer: strategies for overcoming resistance tochemotherapy .Nat Rev Cancer 2003; 3:502–16.
2. Gasparini G. Metronomic scheduling: the future of chemotherapy? Lancet Oncol 2001; 2: 73 3–40.
3. Mross K, Steinbild S. Metronomic anti-cancer therapy: an ongoing treatment option for advanced cancer patients. Journalof Cancer Therapeutics & Research 2012, doi: 10.7243/2049-7962-1-32.
4. Hahnfeldt P, Folkman J, Hlatky L. Minimizing long-term tumor burden: the logic formetronomic chemotherapeutic dosing and its antiangiogenic basis. J Theor Biol. 2003; 220(4):545–554.
5. Shaked Y, Emmenegger U, Man S, etal*.*Optimal biologic dose of metronomic chemotherapy regimens is associated with maximum antiangiogenic activity. Blood. 2005; 106(9):3058–3061.
6. Kamat AA, Kim TJ, Landen CN Jr,etal*.*Metronomic chemotherapy enhances the efficacy of antivascular therapy in ovarian cancer. Cancer Res. 2007; 67(1):281–288.
7. Yokoi T, Tamaki T, Shimizu T, *et al.*A pilot study of a metronomic chemotherapy regimen with weekly low-dose docetaxel for previously treated non-small cell lung cancer .Lung Cancer: Targets and Therapy 2012; 3:15-20.
8. Joshi A, Noronha V, Patil VM, *et al.*Efficacy and safety of metronomic administration of paclitaxel for advanced recurrent non-small-cell lung cancerIndian Journal of Cancer, Vol. 50, No. 2, April-June, 2013, pp. 122-127.
9. Oken MM, Creech RH, Tormey DC, *etal.*Toxicity and response criteria of the Eastern Cooperative Oncology Group. Am J Clin Oncol. 1982; 5(6):649–655.
10. Therasse P, Arbuck SG, Eisenhauer EA, *et al.*New guidelines to evaluate the response to treatment in solid tumors. European Organization for Research and Treatment of Cancer, National Cancer Institute of the United States, National Cancer Institute of Canada. J Natl Cancer Inst. 2000; 92 (3):205–216.
11. C, [Wong A](http://www.ncbi.nlm.nih.gov/pubmed/?term=Wong%20A%5BAuthor%5D&cauthor=true&cauthor_uid=23728939), [Francia G](http://www.ncbi.nlm.nih.gov/pubmed/?term=Francia%20G%5BAuthor%5D&cauthor=true&cauthor_uid=23728939), *et al.*Preclinical analysis of resistance and cross-resistance to low-dose metronomic chemotherapy. [Invest New Drugs.](http://www.ncbi.nlm.nih.gov/pubmed/23728939) 2014 Feb;32(1):47-59.
12. André N, Banavali S, Snihur Y, *etal.*Has the time come for metronomics in low-income and middle-income countries? Lancet Oncol 2013;14:e239-48.
13. Pasquier E, Kavallaris M, Andre N. Metronomic chemotherapy: New rationale fornew directions. Nat Rev ClinOncol 2010; 7: 455-65.
14. Patil V, Noronha V, Krishna V, et al*.*Oral metronomic chemotherapy in recurrent, metastatic and locally advanced head and neck cancers .ClinOncol 2013; 25: 388.
15. Patil V, Noronha V, D'cruzAK,etal*.*Metronomic chemotherapy inadvanced oral cancers. J Cancer Res Ther 2012; 8: S106-10.
16. Gorn M, Habermann CR, Anige M, *et al.*A pilot study of docetaxel and trofosfamide as second-line “metronomic” chemotherapy in the treatment of metastatic non-small cell lung cancer (NSCLC).Onkologie.2008; 3(4): 185–189.

5/18/2016