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## **Molecular aspects of 5-Fluorouracil and Oxaliplatin activity**

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2010

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Noordhuis, P. (2010). *Molecular aspects of 5-Fluorouracil and Oxaliplatin activity*.

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## Chapter

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### **A non-radioactive sensitive assay to measure 5-fluorouracil incorporation into DNA of solid tumors**

P.Noordhuis, U Holwerda, JAM van Laar, CL van der Wilt, GJ Peters

*Nucleosides Nucleotides & Nucleic acids 23(8&9), 1481-1484, 2004*

### **Abstract**

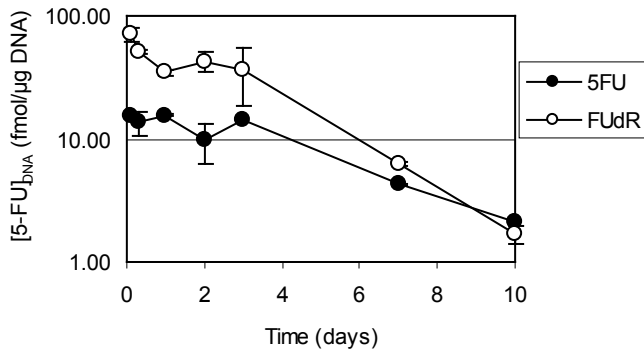
A non-radioactive method to determine 5-Fluorouracil (5FU) incorporation into DNA has been developed. Isolated DNA was enzymatically degraded to bases and the resulting 5FU was measured with standard (gas-chromatography coupled to mass spectrometry (GC-MS) and compared with that of radioactive 5FU in a cell line. Incorporation into DNA of the murine Colon 26-B tumor treated with maximal tolerated doses of 5FU and fluorodeoxyuridine (FdUR) was maximal after 2 hour and was 15.4 and 71.0 fmol/ $\mu$ g DNA, respectively. After a plateau for about 3 days a decrease was observed to  $\pm$  2 fmol/ $\mu$ g DNA after 10 days. The assay is very sensitive and reproducible and can be used in a clinical setting.

### **Introduction**

The mechanism of action of 5FU has been related to inhibition of thymidylate synthase (TS) (1) while the incorporation into RNA and DNA may also play a role in the efficacy of the drug(2). Inhibition of TS results in depletion of dTTP and an increase of dUTP followed by a decreased DNA synthesis. Addition of Leucovorin which acts as precursor of 5,10-methylene-tetrahydrofolate, and is necessary for the formation of the ternary complex with TS and FdUMP, potentiates the inhibition of TS. Incorporation of 5FU into RNA may lead to disturbance of mRNA processing and protein synthesis but could be decreased with UDP-glucose without affecting the efficacy of the drug (3), indicating that incorporation into RNA only plays a minor role in the anti tumor activity. 5FU incorporation into DNA may also play a role in the cytotoxic effect of the drug (4), due to the formation of DNA strand breaks after excision by uracil DNA glycosylase. Repair of these strand breaks is hampered by the imbalance of the deoxynucleotide pools as result of TS inhibition. Incorporation of FdUTP and dUTP into DNA is dependent on the action of dUTPase (5). Since the use of radio labeled 5FU is not practicable in a clinical setting, we developed a non-radioactive assay to determine the 5FU incorporation into DNA.

### **Materials and Methods**

Isolated and purified DNA from murine WiDr colon cancer cells and Colon26-B tumor tissue was dissolved in digestion buffer containing 40 mM Tris, 1 mM MgCl<sub>2</sub>, 0.1 mM ZnCl<sub>2</sub> and 40 mM KH<sub>2</sub>PO<sub>4</sub>, pH=7.4 at 55°C. To 340  $\mu$ l DNA suspension 20  $\mu$ l DNase I (1 mg/ml), 20  $\mu$ l Nuclease P1 (250 U/ml), 10  $\mu$ l Alkaline Phosphatase (1000 U/ml) and 10  $\mu$ l Thymidine Phosphorylase (600 U/ml) were added. Incubation of this

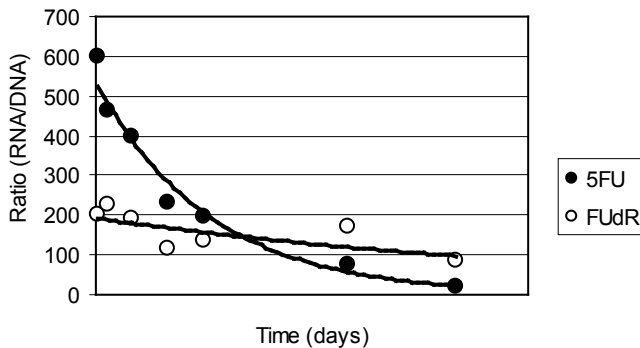


**Figure 1.** Incorporation of 5-FU into DNA of murine Colon-26B tumor tissue at different time points after administration of 80 mg/kg 5-FU (closed circles) and 400 mg/kg FUdR (open circles). Values are means of 2-3 samples  $\pm$  SD

reaction mixture overnight at 37°C resulted in complete degradation of the DNA to bases, including 5FU. After addition of 50  $\mu$ l of the internal standard 5FU-<sup>15</sup>N<sub>2</sub> (0.1  $\mu$ M), 5FU was measured after extraction and derivatization with pentafluorobenzylbromide. GC-MS measurement of 5FU was described before (6).

## Results

The assay was first validated with in WiDr Colon cancer cells as described for 5FU incorporation into RNA (6). The incorporation of 25  $\mu$ M 5FU into DNA after 2 hr and 4 hr was  $0.15 \pm 0.02$  and  $0.35 \pm 0.09$  pmol/ $\mu$ g DNA as determined with GC-MS (means  $\pm$  SEM). At 50  $\mu$ M the 5FU incorporation after 2 and 4 hr was  $0.35 \pm 0.097$  and  $0.57 \pm 0.141$  pmol/ $\mu$ g DNA, respectively. This was in the same range as described previously with radioactive 5FU in this cell line,  $0.63 \pm 0.11$  pmol/hr/ $10^6$  cells at 15  $\mu$ M 5FU (7). Subsequently incorporation into DNA of Colon26-B tumors was determined. Incorporation into DNA after treatment with 5FU (80 mg/kg) or FUdR (400 mg/kg) was maximal at 2 hours after drug treatment and was 15.4 and 71.0 fmol/ $\mu$ g DNA, respectively. The level of incorporation remained high for about 3 days after which a decrease was observed to  $\pm 2$  fmol/ $\mu$ g DNA after 10 days both for 5FU and FUdR. In contrast to RNA, DNA incorporation did not correlate with the intra-tumoral concentration of 5FU, neither for 5FU or FUdR (8)



**Figure 2.** Ratio of incorporation of fluoropyrimidines into RNA and DNA. For each separate tumor the incorporation into RNA and DNA after treatment with 5FU or FUdR was calculated and plotted as  $(\text{pmol}/\mu\text{g RNA})/(\text{fmol}/\mu\text{g DNA})$ . The RNA data have been published earlier (From Ref 8.)

## Discussion

The higher initial incorporation into DNA is in agreement with a higher formation of FdUMP from FUdR compared to 5FU(8). Apparently potential breakdown of FdUTP by dUTPase did not prevent the incorporation into DNA. The relatively long retention of 5FU in DNA might be due to the favorable incorporation conditions because of dTTP depletion as result of TS inhibition but also to a rather inefficient repair by Uracil DNA Glycosylase. The data give strong evidence that DNA incorporation of fluoropyrimidines contributes to their antitumor activity since the better antitumor activity of FUdR was associated with a higher extent of incorporation of FUdR into DNA as compared to 5FU.

This method may provide the possibility to evaluate the role of 5FU incorporation into DNA in tumor tissue during treatment of patients and the relative role of 5FU incorporation into DNA compared to RNA incorporation and the inhibition of TS. Furthermore this approach may help to evaluate potential changes in the mechanisms when 5FU is combined with other drugs such as irinotecan, oxaliplatin and raltitrexed.

## References

1. Peters GJ, K. C. Fluoropyrimidines as antifolate drugs. In: J. AL (ed.), *Antifolate Drugs in Cancer Therapy*, pp. 101 - 145. Totowa: Humana Press Inc, 1999.
2. Sobrero, A. F., Aschele, C., and Bertino, J. R. Fluorouracil in colorectal cancer--a tale of two drugs: implications for biochemical modulation. *J Clin Oncol*, 15: 368-381, 1997.
3. Codacci-Pisanelli, G., Van der Wilt, C. L., Smid, K., Noordhuis, P., Voorn, D., Pinedo, H. M., and Peters, G. J. High-dose 5-Fluorouracil with uridine-diphosphoglucose rescue increases thymidylate synthase inhibition but not 5-Fluorouracil incorporation into RNA in murine tumors. *Oncology*, 62: 363-370, 2002.
4. Schuetz, J. D., Wallace, H. J., and Diasio, R. B. 5-Fluorouracil incorporation into DNA of CF-1 mouse bone marrow cells as a possible mechanism of toxicity. *Cancer Res*, 44: 1358-1363, 1984.
5. Caradonna, S. J. and Cheng, Y. C. The role of deoxyuridine triphosphate nucleotidohydrolase, uracil-DNA glycosylase, and DNA polymerase alpha in the metabolism of FUdR in human tumor cells. *Mol Pharmacol*, 18: 513-520, 1980.
6. Peters, G. J., Noordhuis, P., Komissarov, A., Holwerda, U., Kok, M., Van Laar, J. A., Van der Wilt, C. L., Van Groeningen, V., and Pinedo, H. M. Quantification of 5-fluorouracil incorporation into RNA of human and murine tumors as measured with a sensitive gas chromatography-mass spectrometry assay. *Anal Biochem*, 231: 157-163, 1995.
7. Peters, G. J., Laurensse, E., Leyva, A., Lankelma, J., and Pinedo, H. M. Sensitivity of human, murine, and rat cells to 5-fluorouracil and 5'-deoxy-5-fluorouridine in relation to drug-metabolizing enzymes. *Cancer Res*, 46: 20-28, 1986.
8. van Laar, J. A., van der Wilt, C. L., Rustum, Y. M., Noordhuis, P., Smid, K., Pinedo, H. M., and Peters, G. J. Therapeutic efficacy of fluoropyrimidines depends on the duration of thymidylate synthase inhibition in the murine colon 26-B carcinoma tumor model. *Clin Cancer Res*, 2: 1327-1333, 1996.

