

# Neoadjuvant Chemotherapy is Given Prior to a Local Treatment Such as Surgery, and is Designed to Shrink the Primary Tumor

# Lichao Sun<sup>1\*</sup>, Duxin Sun<sup>2</sup>

<sup>1</sup>State Key Laboratory of Molecular Oncology, National Cancer Center/Cancer Hospital, Chinese Academy of Medical Sciences, Beijing, PR China; <sup>2</sup>Department of Pharmaceutical Sciences, University of Michigan, Ann Arbor, MI 48109, USA SHORT COMMUNICATION the most common side-effects of chemotherapy:

Chemotherapy (often abbreviated to chemo and sometimes CTX or CTx) is a type of cancer treatment that uses one or more anticancer drugs (chemotherapeutic agents) as part of a standardized chemotherapy regimen. Chemotherapy may be given with a curative intent (which almost always involves combinations of drugs), or it may aim to prolong life or to reduce symptoms (palliative chemotherapy). Chemotherapy is one of the major categories of the medical discipline specifically devoted to pharmacotherapy for cancer, which is called medical oncology [1-6].

The term chemotherapy has come to connote non-specific usage of intracellular poisons to inhibit mitosis, cell division. The connotation excludes more selective agents that block extracellular signals (signal transduction). The development of therapies with specific molecular or genetic targets, which inhibit growth-promoting signals from classic endocrine hormones (primarily estrogens for breast cancer and androgens for prostate cancer) are now called hormonal therapies. By contrast, other inhibitions of growth-signals like those associated with receptor tyrosine kinases are referred to as targeted therapy. Importantly, the use of drugs (whether chemotherapy, hormonal therapy or targeted therapy) constitutes systemic therapy for cancer in that they are introduced into the blood stream and are therefore in principle able to address cancer at any anatomic location in the body. Systemic therapy is often used in conjunction with other modalities that constitute local therapy (i.e. treatments whose efficacy is confined to the anatomic area where they are applied) for cancer such as radiation therapy, surgery or hyperthermia therapy. Traditional chemotherapeutic agents are cytotoxic by means of interfering with cell division (mitosis) but cancer cells vary widely in their susceptibility to these agents. To a large extent, chemotherapy can be thought of as a way to damage or stress cells, which may then lead to cell death if apoptosis is initiated. Many of the side effects of chemotherapy can be traced to damage to normal cells that divide rapidly and are thus sensitive to anti-mitotic drugs: cells in the bone marrow, digestive tract and hair follicles. This results in

the most common side-effects of chemotherapy: myelosuppression (decreased production of blood cells, hence also immunosuppression), mucositis (inflammation of the lining of the digestive tract), and alopecia (hair loss). Because of the effect on immune cells (especially lymphocytes), chemotherapy drugs often find use in a host of diseases that result from harmful overactivity of the immune system against self (so-called autoimmunity). These include rheumatoid arthritis, systemic lupus erythematosus, multiple sclerosis, vasculitis and many others.

All chemotherapy regimens require that the recipient be capable of undergoing the treatment. Performance status is often used as a measure to determine whether a person can receive chemotherapy, or whether dose reduction is required. Because only a fraction of the cells in a tumor die with each treatment (fractional kill), repeated doses must be administered to continue to reduce the size of the tumor. Current chemotherapy regimens apply drug treatment in cycles, with the frequency and duration of treatments limited by toxicity.

## REFERENCES

- Olowokure O, Qi X. Pancreatic cancer: current standards, working towards a new therapeutic approach. Expert Rev Anticancer Her. 2014; 14: 495-497.
- Goonetilleke KS, Siriwardena AK. Systematic review of carbohydrate antigen (CA 19-9) as a biochemical marker in the diagnosis of pancreatic cancer. Eur J Surg Oncol. 2007; 33: 266-270.
- 3. Tanco S, Zhang X, Morano C, Avilés FX, Lorenzo J, et al. Characterization of the Substrate 6pecLucLt\ of Human Carboxypeptidase A4 and Implications for a Role in Extracellular Peptide Processing. J Biol Chem. 2010; 285: 18385-18396.
- Huang H, Reed CP, Zhang JS, Shridhar V, Wang L, et al. Carboxypeptidase A3 (CPA3): A Novel Gene Highly Induced by Histone Deacetylase Inhibitors during 'L',erentLDtLon of Prostate Epithelial Cancer Cells. Cancer Res. 1999; 59: 2981-2988.
- Kayashima T, Yamasaki K, Yamada T, Sakai H, Miwa N, et al. He novel imprinted carboxypeptidase A4 gene (CPA4) in the 7q32 imprinting domain. Hum Genet. 2003; 112: 220-226.

### Correspondence to:

Received: June 28, 2020; Accepted: July 13, 2020; Published: July 20, 2020

**Copyright:** © 2020 Sun L, et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Citation: Sun L, Sun D (2018) The Potential Diagnostic Value of Serum Cpa4 Level for Pancreatic Cancer. Chemo Open Access 7: 254. doi: 10.4172/2167-7700.1000257

### Sun L, et al.

6. Sun L, Burnett J, Guo C, Xie Y, Pan J, et al. CPA4 is a promising diagnostic serum biomarker for pancreatic cancer. Am J Cancer Res. 2016; 6: 91-96.