Glyco-Polypeptides (Comosain) and Chimeric White Blood Cell Therapy in Late-Stage Refractory Solid Carcinoma of Lung, Prostatic and Bladder Cancers Review of 35 Cases

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Purpose
Administered of oral Glyco-polypeptides (Comosain) in cancer treatment in nonclinical trials has been reported as early in 1968 by Wolf M, & Ransberger k [1]. In vitro and animal studies have suggested of anti-metastatic effect for Glyco-polypeptides (Comosain). Batkin & Taussig in 1988 reported that orally administered Glyco-polypeptides (Comosain) reduced the incidence of pulmonary metastasis in Lewis lung cancer cells in mice. In recent years, 1988 Batkin & Taussig suggested the antitumor mechanisms are due to fibrinolytic effect in Glyco-polypeptides (Comosain) [2-4]. Taussig & Batkin in 1988 discovered that Glyco-polypeptides (Comosain) has anti-Platelet aggregation effects [5]. Taussig and Batkin in 1985 also discovered Inhibition the growth of tumor cells such as Lewis lung carcinoma, V-8 lymphoma, MC1-1 acites, KATO-gastric carcinoma cells. Maurer & Hozumi, in 1994 Discovered Glyco- polypeptides (Comosain) Induced Differentiation in leukemic cells. Hale, & Haynes in 1992 and Cantrell et. in 1996 had suggested that due to Major Histocompatibility Protein Kinases, such as MMAPK (Major Mitogen Activating Protein Kinase) and TPK (Tyrosine Phosphorylation Kinase) inhibitors were activated by Glyco-polypeptides (Comosain) [6,7]. T-cell activation and cascade production of Interleukin 2, 6, 8, and TNF-a (Tumor Necrotizing Factors) via CD-2, CD-3 surface antigen of WBC.

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From the experimental studies above, we conclude that activation of Glyco-polypeptides (Comosain) in lymphocytes and T-cells have anti-metastatic effects both in vitro and in vivo.

In the present study, we compared the modulation of low dose cohort and high dose cohort of Comosain administration to the patients with stage 3, and stage 4, refractory solid tumors, which including various types of carcinoma of lung, prostate and bladder. All patients failed previously on at least two regimens of chemotherapy and /or failed on radiation therapy, The treatment were carried out for at least 24 to 28 weeks, the complete blood count, liver, renal function, hematopoietic elements, tumor markers were evaluated at an interval of every 2 to 4 weeks, the computerized tomography scan were performed at an interval of every 3 to 4 months. The size of tumors were measured, the tumor markers were recorded for the evaluation of complete response (CR), partial response (PR), stable disease (SD), and progressive disease (PD) according to the Standard Response Criteria of National Cancer Institute (NCI). The Common toxicity were recorded by using NCI’s Standard Toxicity Criteria. The results of CR and PR were promising and astonishing when Comosain were administered in patients of high dose group.

Materials and Methods

Glyco-polypeptides (Comosain) were purchased from Natural Organics Laboratories, Amityville, N.Y., Capsules to contain Glyco-polypeptides (Comosain) were purchased from Capusugel Co. Greenwood, North Carolina. Comosain were analysed by using SDS-Polyacryl- Amide Gel Electrophoresis (SDS-PAGE), Cation Exchange Chromatography (CEC), Florescense High Performance Liquid Chromatography (FPLC) to determinate the purity and separation of Glyco-polypeptides (Comosain) fraction of F1, F2, F3, F4, F5, F6, F9 in stem.

Glyco-polypeptides (Comosain) were detected by Amperometric detection. Monosaccharides fraction are L-fructose, D-galactosamine, D-glucosamine, D-xyllose, D-mannose, D-glucose, D-galactose, D-fructose, and Deoxyribose [9].

Clinical Application and Study protocol;

I. Patients Eligibility and Selection (Total number of patients: 35)

(I) Patients with stage III and IV solid cancer of lung, prostate and bladder with tissue proof of well-documented malignancies, whether by tissue biopsies and have not been helped by conventional radiation therapy and/or chemotherapies for at least two separated regimens are eligible for this study.

(II) Or patients must have no available therapy known to provide clinical benefit. For example, the lung, prostate and bladder cancer patients must have failed at least 2 chemotherapy regimens in the metastatic setting.

(III) Additionally

Patient’s age is between 18 and 95+ years, not taking anticoagulants, have no history of abdominal fistula, gastro-enteral perforation, peptic ulcer diseases, or intra-abdominal abscess within 4 months prior to study enrollment, and patient has not had major surgery within 4 weeks prior to study enrollment, and other requirements are same as NCI’s criteria. Also, Patient does not have uncontrolled hypertension, diabetes, or cardiac arrhythmia, and not allergic to Glyco-polypeptides (Comosain) -containing products, not pregnant or breastfeeding. Patient’s WBC count < 3K/ul, hemoglobin < 9.0 g/dL, platelet counts must be <100,000/ul, and INR < 1.5 have no significant abnormal hepatic and/or renal function.

Patient’s tumors are measurable between 0.2 – 10+ cm in size and number between 1–15+. All measurable tumors that have spread to the bones, liver, lung, kidney, and abdomen will be included in the data analysis.

Patients who are eligible for this study will be randomly assigned to either the low dose cohort or the high dose cohort by a coin toss. Each study subject will be assigned a patient number for the purpose of this study.

Methods of Study

The Dose of Comosain at 50 mg /kg/day is extrapolated from in vivo animal studies and determined to be safe by a Safety study on healthy human subjects.

The High Dose cohort will be given Comosain at 50 mg /kg/day (at a body weight of 50- 60 kg) to a maximum of 2400 mg /day and divided into 2 doses/day of 1200 mg/dose, and taken with meals. Low dose Cohort patients will be given Comosain at 10mg/kg/day, that is 500 mg /day, divided into two doses of 250 mg / dose and taken with meals. High Dose Cohort – The number of patients will be at least 21. Low dose Cohort--The number of patients will be at least 14.

(A) Blood/laboratory tests will be scheduling every 2-4 weeks, which include CBC, Chemistry- 7, Chemistry-24, liver and renal function, CEA, CA125, CA153, CA199, PSA, TSH, alfa-Feto- Protein and other tumor markers.

(B) Radiological tests will be assessed every 3-4 months. Each patient will be also assessed every 4 weeks for any side effects that they may have experienced.

(C) Using NCI standard toxicity criteria for hematology, renal, and hepatic system evaluation.

(D) Adverse events, serious adverse events reporting information also using NCI criteria

Duration and Route of Administration

The patients will be evaluated by blood tests and/or CT scans at the end of each 6 weeks cycle and at six months for signs of disease progression. If the disease did not progress, then treatment will continue, and the patient will be evaluated every six months thereafter until the investigator determines otherwise. If the disease did progress, then the patient will be taken off the study. On the Humanitarian base, the low dose group patients will be transferred to the high dose group due to lack of efficacy in the treatment.

Results

At the end of six months, each patient will be determined whether or not to continue with this therapy and assess the efficacy of the therapy by using NCI Standard response Criteria:

1. Evaluation of Target Lesions

A. Complete Response (C R): Disappearance of all Target lesions, and lymph nodes must be reduced < 10mm.

B. Partial Response (P R): At least a 30 % decrease in the sum of the diameters of target lesions compared with the baseline sum diameters.

C. Progressive Disease (P D): At least a 20% increase in the sum of the diameters of target lesions compared with the smallest sum on study. In addition, the sum must demonstrate an absolute increase of at least 5 mm. The appearance of new lesions is also considered progressions.

D. Stable Disease (S D): Neither sufficient shrinkage to qualify
for PR nor sufficient increase to qualify for PD, taking as reference to the smallest sum diameters while on study.

II). Evaluation of Non-Target Lesions
A. Complete Response (CR): Disappearance of all non-target lesions and normalization of tumor marker level. All lymph nodes must be < 10mm short axis.
B. Non-CR/ Non-PD: Persistence of one or more non-target lesion(s) and/or maintenance of tumor marker level above the normal limits.
C. Progressive Disease (PD): Appearance of one or more new lesions and/or unequivocal progression of existing non-target lesions. Unequivocal progression should not normally trump target lesion status.

Results
Age distribution, all the patients are mainly in their 6 decade and above. The disease classification and distribution are as following: lung carcinoma account for 45.7% (16/35), in prostate carcinoma the incidence is 42.8% (15/35), in bladder carcinoma the incidence are 11.4% (4/35).

The overall clinical response rate in high dose group patient:

|        | CR  | PR  | SD   | PD  |
|--------|-----|-----|------|-----|
| Lung CA| 75% (12/16) | 25% (4/16) | 0% (0/32) | 0% (0/32) |
| Prostate CA | 80% (12/15) | 13.3% (2/15) | 0% (0/15) | 6.7% (1/15) |
| Bladder CA | 50% (2/4) | 25% (1/4) | 0% (0/4) | 25% (1/4) |

The tumor markers such as CEA, CA-125, CA-199, PSA, and alpha-feto-protein are been monitored, their values are corresponding to the tumor masses, they return to normal value when tumor have CR, and when the tumor progress the tumor marker value are elevated.

The serious adverse effect in toxicity in both groups are not observed, there were no serious hematopoietic or hepato-renal toxicity, no anaphylactic reaction or life threaten events. There were rarely minor side effects such as nausea, vomiting, diarrhea, palpitation, headache, insomnia, pruritus, urticaria, and skin rash. We conclude that Glyco-polypeptides (Comosain, Ananas) administered in an amount of 2500 to 3000 mg/day to the patients with average body weight are effective and non-toxic.

Case-1: Lung cancer pre and post treatments:

Case-2: Lung cancer pre and post treatments:

Case-3: Lung cancer pre and post treatments:

Lung Cancer 4th Stage Metastasis to Brain, bone, and liver - 4.5 Months Treatment

Lung Cancer 4th Stage Met to Brain, bone, and liver - 4.5 Months Treatment

Prostate Cancer 4th Stage Mets - 4 Months Treatment
Case 1: Breast cancer pre and post treatment (Kidney mets)

Case 2: Breast cancer pre and post treatment (Bone Mets)

Case 3: Breast cancer pre and post treatment (Liver Mets, 9.2cm shrinkage to 2.6cm)

Case 4: Ovarian cancer pre and post treatment

Conclusion
In summary, Glyco-polypeptides (Comosain) administration in double-blind study showed effectiveness only in patients with high dose cohort of 50 mg/kg/day regimen. The low dose cohort showed no efficacy at all. Both groups did not show serious adverse effects such as leukopenia, anemia, hepatorenal toxicity, anaphylactic reaction, and life-threaten events. Minor adverse effects such as nausea, vomiting, diarrhea, urticaria, insomnia, palpitation, pruritus, and headache occurred rarely.

The remarkable cancercidal effects probably due to massive production of Interleukin-II, VI, VIII, and tumor necrotizing factors from CD-2, CD-3 in monocytes and lymphocytes. The fibrinolytic effects on tumor surface antigens of CD-44, CD-44V, CD-44S, CD-45, and CD-47 which induce dehydration, necrosis, and possible calcification in the tumor cells. This action mechanism of Glyaco-polypeptides (Comosain) is mainly attributed to inhibition of 2 kinases: Major Mitogen Activating Protein Kinases and Tyrosine Phosphorylation Kinases. In the WBC culture test with concentration of Glyco-polypeptides (Comosain) in an amount of 1 mg/ml will increase the production of Interleukin II by 400 times/10^6 WBC, Interleukin-6 by 650 times/10^6 WBC, and the TNF by 42 times/10^6 WBC.

The results in the high dose group patients showed remarkable CR rates of 74%, PR rate of 20%, SD rate of 0%, PD rate of 5.7%. Dr. HR Maurer in his complimentary tumor therapy also showed Glyco-polypeptides (Comosain) in an amount of 1000-to-3000 mg/day for the period of 1 to 3 years has no severe side effects nor any life threaten events. (Bromalain in der komplementaren Tumortherapie, Cancer Journal 31: pp.66-73, 1999) [10-14].

Question: What are the most effective treatments in late stage of gynecological cancers?
Finding: Since our patients have failed chemotherapy and radiation therapy, we are looking for new method to treat those clinical situations. We used Comosain in cancer treatment through our experience and literature evidence, which showed Comosain with remarkable effectiveness against most tumors.
Meaning: In our clinical trial, we entered 14 patients into low dose cohort and 21 patients into high dose group. We transferred low dose group into high dose cohort due to ineffectiveness after 6 weeks. Total number of 35 high dose group patients were discussed and analyzed.

Our findings in this study were correspondence with Dr. Maurer, Dr. Eckert, Dr. Harrach (5, 9,11,12, 21, 31, 41) and many other authors. Glyco-polypeptides in treating various type of cancers only high dose are effective. There were no hematological, hepato-renal toxicities.

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References
1. T Harrach, K Eckert, H R Maurer, I Machleidt, W Machleidt (1998) Isolation and characterization of two forms of an acidic bromelain stem proteinase an acidic Bromelain Proteinase. Journal of Protein Chemistry 17: 351-361.
2. Barnes, S (1195) Effect of genistein on in vitro and in vivo models of cancer: Journal of Nutrition 125: 77S-783S.
3. Batkin, S, TaussigSJ, Szekeresz J (1988) Antimetastatic effect of bromelain with or without its proteolytic and anticoagulant activity, Journal Cancer Res. Clinical Oncology.
4. Birch M, Miychell S, Hart IR (1991) Isolation and characterization of human melanoma cell variants expressing high and low levels of CD 44. Cancer Research 51: 6660-6667.

5. Cantrell (1996) Micosomal study of Bromelain and its intracellular signal transduction namely, T-cell receptor (TCR)/CD3 signaling and Interleukin II (IL-2) production, which are activated by Major Histocompatibility Complex (MHC) expressed on antigen presenting cells (APC).; Ann. Review Immunology 14: 259-274.

6. Castillo, MH, Perkins E (1989) The effect of Bio-flavonoid-Quercetin on squamous cell carcinoma of head and neck origin, American Journal of Surgery 158: 351-355.

7. Cooreman W (1978) Bromelain Pharmacological Enzymes-Properties and assay method, (pp 07-12; Ruyssen R. and Lauwers A. Story-Scientia Scientific Publishing Co.

8. Gallatin, W M, E.A. Wayner, P A Hoffman (1989) Structure homology between lymphocyte receptor for high endothelium and class III extracellular matrix. Proc. National Acad. Sci. USA86, 4654, 1989.

9. Desser, L, Rehberger A, Paukovits, W (1993) Proteolytic enzymes and amylase induce cytokine production in human peripheral blood mononuclear cells in vitro, Cancer Biotherapy 9: 253-263.

10. Eckert K, Grunberg E, Garbin F, Maurer HR (1997) Preclinical studies with prothymosin a-1 on mononuclear cells from tumor patients, International Journal Immuno pharmacology 19: 493-500.

11. Eckert Klaus, Grabowska Edyta, Strange Rainer, Schneider Ulrike, Maurer H.Raine, et al. (1999) Effects of oral bromelain administration on the impaired immunocytotoxicity of mononuclear cells from mammary tumor patients, Oncology Report 6: 1191-1199.

12. Harrach T, Gebauer F, Eckert K, Kunze R, Maurer H R (1994) Bromelain proteinases modulate the CD44 expression on human Molt 4/8 leukemia and SK-Mel28 melanoma cells in vitro. International Journal of Oncology 5: 485-488.

13. Kandaswami C, Perkins E (1991) Anti-proliferative effects of citrus flavonoids on human squamous cell carcinoma in vitro: Cancer Lett 56: 147-152.

14. Matsumoto G, Nghiem MP, Nozaki N, Schmits R, Penninger JM (1998) Cooperation between CD44 and LFA-1/CD11a adhesion receptors in lymphokine-Activated killer cell cytotoxicity, Journal of Immunology 160: 5781-589.