

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
4 October 2001 (04.10.2001)

PCT

(10) International Publication Number  
**WO 01/73132 A1**

- (51) International Patent Classification<sup>7</sup>: **C12Q 1/68**, G01N 33/53, 33/574, A61K 39/00, 39/38
- (21) International Application Number: PCT/US01/09790
- (22) International Filing Date: 27 March 2001 (27.03.2001)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:  
60/192,229 27 March 2000 (27.03.2000) US
- (71) Applicant (*for all designated States except US*):  
**THOMAS JEFFERSON UNIVERSITY** [US/US];  
Office of Technology Transfer, 1020 Locust Street, Room  
M6, Philadelphia, PA (US).
- (72) Inventors; and
- (75) Inventors/Applicants (*for US only*): **WALDMAN, Scott, A.** [US/US]; 119 Bleddyn Road, Ardmore, PA 19003 (US).  
**PARK, Jason** [US/US]; 925 Latimer Street, Philadelphia,  
PA 19107 (US). **SCHULZ, Stephanie** [US/US]; 117  
Howard Road, West Chester, PA 19380 (US).
- (74) Agent: **DELUCA, Mark**; Woodcock Washburn Kurtz  
Mackiewicz & Norris LLP, 46th floor, One Liberty Place,  
Philadelphia, PA 19103 (US).
- (81) Designated States (*national*): AE, AG, AL, AM, AT, AU,  
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,  
CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM,  
HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK,  
LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX,  
MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL,  
TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (84) Designated States (*regional*): ARIPO patent (GH, GM,  
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian  
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European  
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,  
IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF,  
CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).
- Published:**  
— with international search report
- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

(54) Title: COMPOSITIONS AND METHODS FOR IDENTIFYING AND TARGETING CANCER CELLS OF ALIMENTARY CANAL ORIGIN

(57) Abstract: Screening and diagnostic reagents, kits and methods for primary and/or metastatic stomach or esophageal cancer are disclosed. Compositions for and methods of imaging and treating primary and/or metastatic stomach or esophageal cancer are disclosed. Vaccines compositions and methods of for treating and preventing primary and/or metastatic stomach or esophageal cancer are disclosed.



**WO 01/73132 A1**

## COMPOSITIONS AND METHODS FOR IDENTIFYING AND TARGETING CANCER CELLS OF ALIMENTARY CANAL ORIGIN

### CROSS REFERENCE TO RELATED PATENT APPLICATIONS

This application claims priority to U.S. Provisional Application Number  
5 60/192,229 filed March 27, 2000, which is incorporated herein by reference.

This application is also related to U.S. Patent Number 5,518,888, issued May  
21, 1996, U.S. Patent Number 5,601,990 issued February 11, 1997, U.S. Patent Number  
6,060,037 issued April 26, 2000, U.S. Patent Number 5,962,220 issued October 5, 1999,  
and U.S. Patent Number 5,879,656 issued March 9, 1999, which are each incorporated  
10 herein by reference and U.S. Patent Application Serial Number 09/180,237 filed March 12,  
1997, which is incorporated herein by reference.

### FIELD OF THE INVENTION

The present invention relates to *in vitro* diagnostic methods for detecting  
cancer cells of the alimentary canal, particularly primary and metastatic stomach and  
15 esophageal cancer, and to kits and reagents for performing such methods. The present  
invention relates to compounds and methods for *in vivo* imaging and treatment of tumors  
originating from the alimentary canal, particularly primary and metastatic stomach and  
esophageal tumors. The present invention relates to methods and compositions for making  
and using targeted gene therapy, antisense and drug compositions. The present invention  
20 relates to prophylactic and therapeutic vaccines against cancer cells of the alimentary canal,  
particularly primary and metastatic stomach and esophageal cancer and compositions and  
methods of making and using the same.

## BACKGROUND OF THE INVENTION

There is a need for reagents, kits and methods for screening, diagnosing and monitoring individuals with cancer originating from the alimentary canal, particularly primary and metastatic stomach and esophageal cancer. There is a need for reagents, kits and methods for identifying and confirming that a cancer of unknown origin is originating from the alimentary canal and for analyzing tissue and cancer samples to identify and confirm cancer originating from the alimentary canal and to determine the level of migration of such cancer cells. There is a need for compositions which can specifically target stomach and esophageal cancer cells. There is a need for imaging agents which can specifically bind to stomach and esophageal cancer cells. There is a need for improved methods of imaging stomach and esophageal cancer cells. There is a need for therapeutic agents which can specifically bind to stomach and esophageal cancer cells. There is a need for improved methods of treating individuals who are suspected of suffering from primary and/or metastatic stomach or esophageal cancer. There is a need for vaccine composition to treat stomach and esophageal cancer. There is a need for vaccine composition to treat and prevent stomach and esophageal cancer. There is a need for therapeutic agents which can specifically deliver gene therapeutics, antisense compounds and other drugs to stomach and esophageal cancer cells.

## SUMMARY OF THE INVENTION

The invention further relates to *in vitro* methods of determining whether or not an individual has cancer originating from the alimentary canal, particularly primary and metastatic stomach and esophageal cancer. The present invention relates to *in vitro* methods of examining samples of non-colorectal tissue and body fluids from an individual to determine whether or not GCC, which is expressed by normal colon cells and by colorectal, stomach and esophageal tumor cells, is being expressed by cells in samples other than colon. The presence of GCC protein or of the GCC gene transcript in samples outside the colorectal track is indicative of expression of GCC and is evidence that the individual may be suffering from metastasized colon cancer or primary or metastatic stomach and/or esophageal cancer. In patients suspected of suffering from colorectal cancer, the presence of GCC protein or of the GCC gene transcript in samples outside the

colorectal track is supportive of the conclusion that the individual is suffering from metastatic colorectal cancer. The diagnosis of metastatic colorectal cancer may be made or confirmed. In patients suspected of suffering from stomach or esophageal cancer, the presence of GCC protein or of the GCC gene transcript in samples outside the colorectal  
5 track is supportive of the conclusion that the individual is suffering from primary and/or metastatic stomach or esophageal cancer. The diagnosis of primary and/or metastatic stomach or esophageal cancer may be made or confirmed.

The invention further relates to *in vitro* methods of determining whether or not tumor cells suspected of being stomach or esophageal cancer are stomach or  
10 esophageal in origin. The present invention relates to *in vitro* methods of diagnosing whether or not an individual suspected of suffering from stomach or esophageal cancer is suffering from stomach or esophageal cancer. The present invention relates to *in vitro* methods of examining samples of tumors from an individual to determine whether or not GCC protein, which is expressed by colorectal, stomach or esophageal tumor cells, is being  
15 expressed by the tumor cells. The presence of a GCC protein or of the GCC gene transcript in a sample from a patient suspected of having stomach or esophageal cancer is indicative of expression of GCC and evidence that the individual may be suffering from stomach or esophageal cancer. In tumors which are suspected of being stomach or esophageal tumors, the presence of a GCC protein or of the GCC gene transcript supports  
20 the conclusion that the tumors are of stomach or esophageal cancer and the diagnosis of stomach or esophageal cancer.

The invention further relates to *in vitro* kits for practicing the methods of the invention and to reagents and compositions useful as components in such *in vitro* kits of the invention.

25 The invention further relates to a method of imaging primary and metastatic stomach and esophageal tumors and to methods of treating an individual suspected of suffering from primary and metastatic stomach and esophageal tumors comprising the steps of administering to said individual a pharmaceutical compositions according to the invention, wherein the compositions or conjugated compounds are present in an amount  
30 effective for therapeutic or diagnostic use in humans suffering from primary and/or metastatic stomach or esophageal tumors.

The invention further relates to a method of delivering an active agent to primary and metastatic stomach and esophageal tumor cells comprising the steps of administering to an individual who has primary and/or metastatic stomach or esophageal tumors, a pharmaceutical composition comprising a pharmaceutically acceptable carrier or diluent, and an unconjugated compositions that comprises a liposome that includes GCC ligands on its surface and an active component encapsulated therein.

The invention further relates to killed or inactivated stomach or esophageal tumor cells that comprise a protein comprising at least one epitope of a GCC protein; and to vaccines comprising the same. In some embodiments, the killed or inactivated cells or particles comprise a GCC protein. In some embodiments, the killed or inactivated cells or particles are haptenized.

The invention further relates to methods of treating individuals suffering from stomach or esophageal cancer and to methods of treating individuals susceptible stomach or esophageal cancer. The method of the present invention provides administering to such individuals an effective amount of such vaccines. The invention further relates to the use of such vaccines as immunotherapeutics.

## **DESCRIPTION OF PREFERRED EMBODIMENTS**

### **Definitions**

As used herein, the term "GCC" is meant to refer to the cellular protein expressed by normal colorectal cells, as well as primary and metastasized colorectal, stomach and esophageal cancer cells. In normal individuals, GCC is found exclusively in cells of intestine, in particular in cells in the duodenum, small intestine (jejunum and ileum), the large intestine, colon (cecum, ascending colon, transverse colon, descending colon and sigmoid colon) and rectum.

As used herein, the term "functional fragment" as used in the term "functional fragment of a GCC gene transcript" is meant to refer to fragments of GCC gene transcript which are functional with respect to nucleic acid molecules with full length sequences. For example, a functional fragment may be useful as an oligonucleotide or nucleic acid probe, a primer, an antisense oligonucleotide or nucleic acid molecule or a coding sequence. The nucleotide sequence encoding human GCC protein is disclosed in

F.J. Sauvage *et al.* 1991 *J. Biol. Chem.* **266**:17912-17918, each of which is incorporated herein by reference..

As used herein, the term "functional fragment" as used in the term "functional fragment of a GCC protein" is meant to fragments of GCC protein which function in the same manner as GCC protein with full length sequences. For example, an immunogenically functional fragment of a GCC protein comprises an epitope recognized by an anti-GCC antibody. A ligand-binding functional fragment of GCC comprises a sequence which forms a structure that can bind to a ligand which recognizes and binds to GCC protein.

As used herein, the term "epitope recognized by an anti-GCC protein antibody" refers to those epitopes specifically recognized by an anti-GCC protein antibody.

As used herein, the term "antibody" is meant to refer to complete, intact antibodies, and Fab fragments and F(ab)<sub>2</sub> fragments thereof. Complete, intact antibodies include monoclonal antibodies such as murine monoclonal antibodies, chimeric antibodies and humanized antibodies.

As used herein, the term "GCC ligand" is meant to refer to compounds which specifically bind to a GCC protein. Antibodies that bind to GCC are GCC ligands. A GCC ligand may be a protein, peptide or a non-peptide.

As used herein, the term "active agent" is meant to refer to compounds that are therapeutic agents or imaging agents.

As used herein, the term "radiostable" is meant to refer to compounds which do not undergo radioactive decay; i.e. compounds which are not radioactive.

As used herein, the term "therapeutic agent" is meant to refer to chemotherapeutics, toxins, radiotherapeutics, targeting agents or radiosensitizing agents.

As used herein, the term "chemotherapeutic" is meant to refer to compounds that, when contacted with and/or incorporated into a cell, produce an effect on the cell including causing the death of the cell, inhibiting cell division or inducing differentiation.

As used herein, the term "toxin" is meant to refer to compounds that, when contacted with and/or incorporated into a cell, produce the death of the cell.

As used herein, the term "radiotherapeutic" is meant to refer to radionuclides which when contacted with and/or incorporated into a cell, produce the death of the cell.

As used herein, the term "targeting agent" is meant to refer compounds which can be bound by and or react with other compounds. Targeting agents may be used to deliver chemotherapeutics, toxins, enzymes, radiotherapeutics, antibodies or imaging agents to cells that have targeting agents associated with them and/or to convert or  
5 otherwise transform or enhance co-administered active agents. A targeting agent may include a moiety that constitutes a first agent that is localized to the cell which when contacted with a second agent either is converted to a third agent which has a desired activity or causes the conversion of the second agent into an agent with a desired activity. The result is the localized agent facilitates exposure of an agent with a desired activity to  
10 the cancer cell.

As used herein, the term "radiosensitizing agent" is meant to refer to agents which increase the susceptibility of cells to the damaging effects of ionizing radiation. A radiosensitizing agent permits lower doses of radiation to be administered and still provide a therapeutically effective dose.

15 As used herein, the term "imaging agent" is meant to refer to compounds which can be detected.

As used herein, the term "GCC binding moiety" is meant to refer to the portion of a conjugated compound that constitutes an GCC ligand.

As used herein, the term "active moiety" is meant to refer to the portion of a  
20 conjugated compound that constitutes an active agent.

As used herein, the terms "conjugated compound" and "conjugated composition" are used interchangeably and meant to refer to a compound which comprises a GCC binding moiety and an active moiety and which is capable of binding to GCC. Conjugated compounds according to the present invention comprise a portion which  
25 constitutes an GCC ligand and a portion which constitutes an active agent. Thus, conjugated compounds according to the present invention are capable of specifically binding to the GCC and include a portion which is a therapeutic agent or imaging agent. Conjugated compositions may comprise crosslinkers and/or molecules that serve as spacers between the moieties.

30 As used herein, the terms "crosslinker", "crosslinking agent", "conjugating agent", "coupling agent", "condensation reagent" and "bifunctional crosslinker" are used

interchangeably and are meant to refer to molecular groups which are used to attach the GCC ligand and the active agent to thus form the conjugated compound.

As used herein, the term "colorectal cancer" is meant to include the well-accepted medical definition that defines colorectal cancer as a medical condition  
5 characterized by cancer of cells of the intestinal tract below the small intestine (i.e. the large intestine (colon), including the cecum, ascending colon, transverse colon, descending colon, and sigmoid colon, and rectum). Additionally, as used herein, the term "colorectal cancer" is meant to further include medical conditions which are characterized by cancer of cells of the duodenum and small intestine (jejunum and ileum). The definition of  
10 colorectal cancer used herein is more expansive than the common medical definition but is provided as such since the cells of the duodenum and small intestine also contain GCC.

As used herein, the term "stomach cancer" is meant to include the well-accepted medical definition that defines stomach cancer as a medical condition characterized by cancer of cells of the stomach.

15 As used herein, the term "esophageal cancer" is meant to include the well-accepted medical definition that defines esophageal cancer as a medical condition characterized by cancer of cells of the esophagus.

As used herein, the term "metastasis" is meant to refer to the process in which cancer cells originating in one organ or part of the body relocate to another part of  
20 the body and continue to replicate. Metastasized cells subsequently form tumors which may further metastasize. Metastasis thus refers to the spread of cancer from the part of the body where it originally occurs to other parts of the body.

As used herein, the term "metastasized colorectal cancer cells" is meant to refer to colorectal cancer cells which have metastasized. Metastasized colorectal cancer  
25 cells localized in a part of the body other than the duodenum, small intestine (jejunum and ileum), large intestine (colon), including the cecum, ascending colon, transverse colon, descending colon, and sigmoid colon, and rectum.

As used herein, the term "metastasized stomach cancer cells" is meant to refer to stomach cancer cells which have metastasized. Metastasized stomach cancer cells  
30 localized in a part of the body other than the stomach.



As used herein, the term "metastasized esophageal cancer cells" is meant to refer to colorectal cancer cells which have metastasized. Metastasized esophageal cancer cells localized in a part of the body other than the esophagus.

As used herein, the term "non-colorectal sample" and "extra-intestinal sample" are used interchangeably and meant to refer to a sample of tissue or body fluid from a source other than colorectal tissue. In some preferred embodiments, the non-colorectal sample is a sample of tissue such as lymph nodes. In some preferred embodiments, the non-colorectal sample is a sample of extra-intestinal tissue which is an adenocarcinoma of unconfirmed origin. In some preferred embodiments, the non-

5  
10

As used herein, "an individual suffering from an adenocarcinoma of unconfirmed origin" is meant to refer to an individual who has a tumor in which the origin has not been definitively identified.

As used herein, "an individual is suspected of being susceptible to stomach or esophageal cancer" is meant to refer to an individual who is at a particular risk of developing stomach or esophageal cancer. Examples of individuals at a particular risk of developing stomach or esophageal cancer are those whose family medical history indicates above average incidence of stomach or esophageal cancer among family members and/or those who have already developed stomach or esophageal cancer and have been effectively

15  
20

As used herein, the term "antisense composition" and "antisense molecules" are used interchangeably and are meant to refer to compounds that regulate transcription or translation by hybridizing to DNA or RNA and inhibiting and/or preventing transcription or translation from taking place. Antisense molecules include nucleic acid molecules and derivatives and analogs thereof. Antisense molecules hybridize to DNA or RNA in the same manner as complementary nucleotide sequences do regardless of whether or not the antisense molecule is a nucleic acid molecule or a derivative or analog. Antisense molecules may inhibit or prevent transcription or translation of genes whose expression is linked to cancer.

25

As used herein, the term "GCC immunogen" is meant to refer to GCC protein or a fragment thereof or a protein that comprises the same or a haptenized product thereof,

30

cells and particles which display at least one GCC epitope, and haptenized cells and haptenized particles which display at least one GCC epitope.

As used herein, the term "recombinant expression vector" is meant to refer to a plasmid, phage, viral particle or other vector which, when introduced into an appropriate host, contains the necessary genetic elements to direct expression of the coding sequence that encodes the protein. The coding sequence is operably linked to the necessary regulatory sequences. Expression vectors are well known and readily available. Examples of expression vectors include plasmids, phages, viral vectors and other nucleic acid molecules or nucleic acid molecule containing vehicles useful to transform host cells and facilitate expression of coding sequences.

As used herein, the term "illegitimate transcription" is meant to refer to the low level or background expression of tissue-specific genes in cells from other tissues. The phenomenon of illegitimate transcription thus provides copies of mRNA for a tissue specific transcript in other tissues. If detection techniques used to detect gene expression are sufficiently sensitive to detect illegitimate transcription, the expression level of the transcript in negative samples due to illegitimate transcription must be discounted using controls and/or quantitative assays and/or other means to eliminate the incidence of false positive due to illegitimate transcription. Alternatively, detection of evidence of GCC gene expression in sample is achieved without detecting GCC gene transcript present due to illegitimate transcription. This is accomplished using techniques which are not sufficiently sensitive to detect the GCC gene transcript present due to illegitimate transcription which is present as background.

U.S. Patent Number 5,518,888, issued May 21, 1996, U.S. Patent Number 5,601,990 issued February 11, 1997, U.S. Patent Number 6,060,037 issued April 26, 2000, U.S. Patent Number 5,962,220 issued October 5, 1999, U.S. Patent Number 5,879,656 issued March 9, 1999, and U.S. Patent Application Serial Number 09/180,237 filed March 12, 1997, relate to targeting ST receptors to treat, image, detect and vaccinate against metastasized colorectal cancer. It has now been discovered that primary and metastatic stomach and esophageal cancer cells express ST receptors (GCC). Accordingly, the compositions and methods described in the above-listed patents and application can be used to treat, image, detect and vaccinate against primary and metastatic stomach and

esophageal cancer. The present invention adapts the earlier invention as related to metastasized colorectal cancer to treat, image, detect and vaccinate against primary and metastatic stomach and esophageal cancer.

## GCC

5 Carcinomas derived from the colorectal cells, stomach or esophagus express GCC. The expression of GCC by such tumors enables this protein and its mRNA to be a specific biomarker for the presence of cancer cells in extra-intestinal tissues and blood. Indeed, this characteristic permits the detection of GCC mRNA by RT-PCR analysis to be a diagnostic test to stage patients with colorectal, stomach or esophageal cancer and follow  
10 patients after surgery for evidence of recurrent disease in their blood as well as to detect colorectal, stomach and esophageal cancers. Further, the GCC may be targeted with a ligand conjugated to an active agent in order to deliver the active agent to tumor cells *in vivo*.

U.S. Patent No. 5,518,888 issued May 21, 1996 to Waldman, PCT  
15 application PCT/US94/12232 filed October 26, 1994, U.S. Application Serial No. 08/467,920 filed June 6, 1995, and U.S. Application Serial No. 08/583,447 filed January 5, 1996, which are each incorporated herein by reference, disclose that metastasized colorectal tumors can be targeted for delivery of active compounds by targeting ST receptors (also referred to as guanylin cyclase C or GCC). The presence of ST receptors on  
20 cells outside of the intestinal tract as a marker for colorectal cancer allows for the screening, identification and treatment of individuals with metastasized colorectal tumors. ST receptors may also be used to target delivery of gene therapeutics and antisense compounds to colorectal cells.

U.S. Patent No. 5,601,990 issued February 11, 1997 to Waldman, PCT  
25 application PCT/US94/12232 filed October 26, 1994, and PCT application PCT/US97/07467 filed May 2, 1997, which are each incorporated herein by reference, disclose that detection of evidence of expression of ST receptors in samples of tissue and body fluid from outside the intestinal track indicate metastasized colorectal cancer.

PCT application PCT/US97/07565 filed May 2, 1997, which is incorporated  
30 herein by reference, disclose that immunogens with epitopes that can be targeted by

antibodies that react with ST receptors can be used in vaccines compositions useful as prophylactic and therapeutic anti-metastatic colorectal cancer compositions.

It has been discovered that, in addition to normal colon cells, to primary and to metastasized colon, stomach and esophageal carcinoma cells also express GCC. Normal stomach and esophageal cells do not express GCC. Thus, the present invention provides the use of GCC as a specific molecular diagnostic marker for the diagnosis, staging, and post-operative surveillance of patients with primary and metastasized stomach and esophageal cancer.

Detection of the expression of GCC employing molecular techniques, including, but not limited to, RT-PCR, can be employed to diagnose and stage patients, follow the development of recurrence after surgery and/or remission, and, potentially, screen normal people for the development of colorectal, stomach or esophageal cancer.

GCC is unique in that it is only expressed in normal intestinal cells. Mucosal cells lining the intestine are joined together by tight junctions which form a barrier against the passage of intestinal contents into the blood stream and components of the blood stream into the intestinal lumen. Therefore, the apical location of cells expressing GCC results in the isolation of such cells from the circulatory system so that they may be considered to exist separate from the rest of the body; essentially the "outside" of the body. Therefore, the rest of the body is considered "outside" the intestinal tract. Compositions administered "outside" the intestinal tract are maintained apart and segregated from the only cells which normally express GCC. Conversely, tissue samples taken from tissue outside of the intestinal tract do not normally contain cells which express GCC.

In individuals suffering from colorectal cancer, the cancer cells are often derived from cells that produce and display the GCC and these cancer cells continue to produce GCC. It has been observed that GCC is expressed by colorectal cancer cells. Likewise, GCC is expressed by stomach and esophageal cancer cells.

The expression of GCC by colorectal tumor cells provides a detectable target for *in vitro* screening, monitoring and staging as well as a target for *in vivo* delivery of conjugated compositions that comprise active agents for the imaging and treatment. GCC can also serve as targets for vaccines which may be used to protect against metastasized colorectal cancer or to treat individuals with metastasized colorectal cancer.

The expression of GCC by stomach and esophageal tumor cells provides a detectable target for *in vitro* screening, monitoring and staging as well as a target for *in vivo* delivery of conjugated compositions that comprise active agents for the imaging and treatment. GCC can also serve as targets for vaccines which may be used to protect against  
5 primary and metastatic stomach and esophageal cancer or to treat individuals with primary and metastatic stomach and esophageal cancer.

### ***In vitro* Diagnostics**

According to some embodiments of the invention, compositions, kits and *in vitro* methods are provided for screening, diagnosing and analyzing patients and patient  
10 samples to detect evidence of GCC expression by cells outside of the intestinal tract wherein the expression of GCC may be suggestive of metastasized colorectal cancer or primary or metastatic stomach or esophageal cancer. In patients suspected of having primary or metastatic stomach or esophageal cancer evidence of GCC expression by cells outside of the intestinal tract is indicative of primary or metastatic stomach or esophageal  
15 cancer and can be used in the diagnosis, monitoring and staging of such patients. Furthermore, the present invention relates to methods, compositions and kits useful in the *in vitro* screening, and analysis of patient and patient samples to detect evidence of GCC expression by tumor cells outside of the intestinal tract wherein the presence of cells that express GCC suggests or confirms that a tumor is of colorectal or stomach or esophageal  
20 cancer origin. In an additional aspect of the invention, compositions, kits and methods are provided which are useful to visualize primary or metastatic stomach or esophageal cancer cells.

*In vitro* screening and diagnostic compositions, methods and kits can be used in the monitoring of individuals who are in high risk groups for stomach or esophageal  
25 cancer such as those who have been diagnosed with localized disease and/or metastasized disease and/or those who are genetically linked to the disease. *In vitro* screening and diagnostic compositions, methods and kits can be used in the monitoring of individuals who are undergoing and/or have been treated for primary stomach or esophageal cancer to determine if the cancer has metastasized. *In vitro* screening and diagnostic compositions,  
30 methods and kits can be used in the monitoring of individuals who are undergoing and/or

have been treated for stomach or esophageal cancer to determine if the cancer has been eliminated. *In vitro* screening and diagnostic compositions, methods and kits can be used in the monitoring of individuals who are otherwise susceptible, i.e. individuals who have been identified as genetically predisposed such as by genetic screening and/or family

5 histories. Advancements in the understanding of genetics and developments in technology as well as epidemiology allow for the determination of probability and risk assessment an individual has for developing stomach or esophageal cancer. Using family health histories and/or genetic screening, it is possible to estimate the probability that a particular individual has for developing certain types of cancer including stomach or esophageal

10 cancer. Those individuals that have been identified as being predisposed to developing a particular form of cancer can be monitored or screened to detect evidence of stomach or esophageal cancer. Upon discovery of such evidence, early treatment can be undertaken to combat the disease. Accordingly, individuals who are at risk for developing stomach or esophageal cancer may be identified and samples may be isolated form such individuals.

15 The invention is particularly useful for monitoring individuals who have been identified as having family medical histories which include relatives who have suffered from stomach or esophageal cancer. Likewise, the invention is particularly useful to monitor individuals who have been diagnosed as having stomach or esophageal cancer and, particularly those who have been treated and had tumors removed and/or are otherwise experiencing

20 remission including those who have been treated for stomach or esophageal cancer.

*In vitro* screening and diagnostic compositions, methods and kits can be used in the analysis of tumors. Expression of GCC is a marker for cell type and suggests the origin of adenocarcinoma of unconfirmed origin suspected of being gastric or esophageal in origin may be stomach or esophageal tumors. Detection of GCC expression can also be

25 used to assist in an initial diagnosis of stomach or esophageal cancer or to confirm such diagnosis. Tumors believed to be stomach or esophageal in origin can be confirmed as such using the compositions, methods and kits of the invention.

*In vitro* screening and diagnostic compositions, kits and methods of the invention can be used to analyze tissue samples from the stomach or esophagus to identify

30 primary stomach or esophageal cancer.

According to the invention, compounds are provided which bind to GCC gene transcript or protein. Normal tissue in the body does not have GCC transcript or protein except cells of the intestinal tract. The expression of GCC is a marker for cell type and is useful in the identification of stomach or esophageal cancer in extra-intestinal  
5 samples.

In some embodiments of the invention, non-colorectal tissue and fluid samples or tumor samples may be screened to identify the presence or absence of GCC protein. Techniques such as ELISA assays and Western blots may be performed to determine whether GCC is present in a sample.

10 In some embodiments of the invention, non-colorectal tissue and fluid samples or tumor samples may be screened to identify whether GCC are being expressed in cells outside of the colorectal tract by detecting the presence or absence of GCC gene transcript. The presence of GCC gene transcript or cDNA generated therefrom can be determined using techniques such as PCR amplification, branched oligonucleotide  
15 technology, Northern Blots (mRNA), Southern Blots (cDNA), or oligonucleotide hybridization.

In some embodiments of the invention, cells of non-colorectal tissue samples or tumor samples may be examined to identify the presence or absence of GCC proteins. Techniques such as immunohistochemistry blots may be performed on tissue sections to  
20 determine whether GCC are present in a sample.

In some embodiments of the invention, cells of non-colorectal tissue samples or tumor samples may be examined to determine whether GCC are being expressed in cells outside of the colorectal track by detecting the presence or absence of the GCC gene transcript. The presence of the GCC gene transcript or cDNA generated therefrom in cells  
25 from tissue sections can be determined using techniques such as *in situ* hybridization.

The presence of GCC in non-colorectal tissue and fluid samples or on cells from non-colorectal tissue samples suggests possible stomach or esophageal cancer. The presence of GCC in a tumor sample or on tumor cells suggests that the tumor may be stomach or esophageal in origin. The presence of the GCC gene transcript in non-  
30 colorectal tissue and fluid samples or in cells from non-colorectal tissue samples suggests

possible stomach or esophageal cancer. The presence of the GCC gene transcript in tumor samples and tumor cells suggests that the tumor may be stomach or esophageal in origin.

Samples may be obtained from resected tissue or biopsy material including needle biopsy. Tissue section preparation for surgical pathology may be frozen and  
5 prepared using standard techniques. Immunohistochemistry and *in situ* hybridization binding assays on tissue sections are performed in fixed cells. Extra-intestinal samples may be homogenized by standard techniques such as sonication, mechanical disruption or chemical lysis such as detergent lysis. It is also contemplated that tumor samples in body fluids such as blood, urine, lymph fluid, cerebral spinal fluid, amniotic fluid, vaginal fluid,  
10 semen and stool samples may also be screened to determine if such tumors are colorectal, stomach or esophageal in origin.

Non-colorectal tissue samples may be obtained from any tissue except those of the colorectal tract, i.e. the intestinal tract below the small intestine (i.e. the large intestine (colon), including the cecum, ascending colon, transverse colon, descending  
15 colon, and sigmoid colon, and rectum) and additionally the duodenum and small intestine (jejunum and ileum). The normal cells of all tissue except those of the colorectal tract do not express GCC. Thus if GCC protein or the GCC gene transcript are detected in non-colorectal samples, the possible presence of colorectal, stomach or esophageal cancer cells is suggested. In some preferred embodiments, the tissue samples are lymph nodes.

20 Tissue samples may be obtained by standard surgical techniques including use of biopsy needles. One skilled in the art would readily appreciate the variety of test samples that may be examined for GCC and recognize methods of obtaining tissue samples.

Tissue samples may be homogenized or otherwise prepared for screening for  
25 the presence of GCC by well known techniques such as sonication, mechanical disruption, chemical lysis such as detergent lysis or combinations thereof.

Examples of body fluid samples include blood, urine, lymph fluid, cerebral spinal fluid, amniotic fluid, vaginal fluid and semen. In some preferred embodiments, blood is used as a sample of body fluid. Cells may be isolated from fluid sample such as  
30 centrifugation. One skilled in the art would readily appreciate the variety of test samples that may be examined for GCC. Test samples may be obtained by such methods as



withdrawing fluid with a syringe or by a swab. One skilled in the art would readily recognize other methods of obtaining test samples.

In an assay using a blood sample, the blood plasma may be separated from the blood cells. The blood plasma may be screened for GCC including truncated proteins  
5 which are released into the blood when one or more GCC are cleaved from or sloughed off from tumor cells. In some embodiments, blood cell fractions are screened for the presence of stomach or esophageal tumor cells. In some embodiments, lymphocytes present in the blood cell fraction are screened by lysing the cells and detecting the presence of GCC protein or the GCC gene transcript which may be present as a result of the presence of any  
10 stomach or esophageal tumor cells that may have been engulfed by the blood cell. In some preferred embodiments, CD34+ cells are removed prior to isolation of mRNA from samples using commercially available immuno-columns.

Aspects of the present invention include various methods of determining whether a sample contains cells that express GCC by nucleotide sequence-based molecular  
15 analysis to detect the GCC gene transcript. Several different methods are available for doing so including those using Polymerase Chain Reaction (PCR) technology, branched oligonucleotide technology, Northern blot technology, oligonucleotide hybridization technology, and *in situ* hybridization technology.

The invention relates to oligonucleotide probes and primers used in the  
20 methods of identifying the GCC gene transcript and to diagnostic kits which comprise such components.

The mRNA sequence-based methods for detect the GCC gene transcript include but are not limited to polymerase chain reaction technology, branched oligonucleotide technology, Northern and Southern blot technology, *in situ* hybridization  
25 technology and oligonucleotide hybridization technology.

The methods described herein are meant to exemplify how the present invention may be practiced and are not meant to limit the scope of invention. It is contemplated that other sequence-based methodology for detecting the presence of the GCC gene transcript in non-colorectal samples may be employed according to the  
30 invention.

A preferred method to detecting the GCC gene transcript in genetic material derived from non-colorectal samples uses polymerase chain reaction (PCR) technology. PCR technology is practiced routinely by those having ordinary skill in the art and its uses in diagnostics are well known and accepted. Methods for practicing PCR technology are disclosed in "PCR Protocols: A Guide to Methods and Applications", Innis, M.A., *et al.* Eds. Academic Press, Inc. San Diego, CA (1990) which is incorporated herein by reference. Applications of PCR technology are disclosed in "Polymerase Chain Reaction" Erlich, H.A., *et al.*, Eds. Cold Spring Harbor Press, Cold Spring Harbor, NY (1989) which is incorporated herein by reference. U.S. Patent Number 4,683,202, U.S. Patent Number 4,683,195, U.S. Patent Number 4,965,188 and U.S. Patent Numbers 5,075,216, which are each incorporated herein by reference describe methods of performing PCR. PCR may be routinely practiced using Perkin Elmer Cetus GENE AMP RNA PCR kit, Part No. N808-0017.

PCR technology allows for the rapid generation of multiple copies of DNA sequences by providing 5' and 3' primers that hybridize to sequences present in an RNA or DNA molecule, and further providing free nucleotides and an enzyme which fills in the complementary bases to the nucleotide sequence between the primers with the free nucleotides to produce a complementary strand of DNA. The enzyme will fill in the complementary sequences adjacent to the primers. If both the 5' primer and 3' primer hybridize to nucleotide sequences on the same small fragment of nucleic acid, exponential amplification of a specific double-stranded size product results. If only a single primer hybridizes to the nucleic acid fragment, linear amplification produces single-stranded products of variable length.

PCR primers can be designed routinely by those having ordinary skill in the art using sequence information. The nucleotide sequence of the GCC gene transcript is set forth in SEQ ID NO:1. To perform this method, RNA is extracted from cells in a sample and tested or used to make cDNA using well known methods and readily available starting materials. Those having ordinary skill in the art can readily prepare PCR primers. A set of primers generally contains two primers. When performing PCR on extracted mRNA or cDNA generated therefrom, if the GCC gene transcript or cDNA generated therefrom is present, multiple copies of the mRNA or cDNA will be made. If it is not present, PCR will

not generate a discrete detectable product. Primers are generally 8-50 nucleotides, preferably about 15-35 nucleotides, more preferably 18-28 nucleotides, which are identical or complementary to and therefor hybridize to the GCC gene transcript or cDNA generated therefrom. In preferred embodiments, the primers are each 15-35 nucleotide, more  
5 preferably 18-28 nucleotide fragments of SEQ ID NO:1. The primer must hybridize to the sequence to be amplified. Typical primers are 18-28 nucleotides in length and are generally have 50% to 60% G+C composition. The entire primer is preferably complementary to the sequence it must hybridize to. Preferably, primers generate PCR products 100 base pairs to 2000 base pairs. However, it is possible to generate products of  
10 50 to up to 10 kb and more. If mRNA is used as a template, the primers must hybridize to mRNA sequences. If cDNA is used as a template, the primers must hybridize to cDNA sequences.

The mRNA or cDNA is combined with the primers, free nucleotides and enzyme following standard PCR protocols. The mixture undergoes a series of temperature  
15 changes. If the GCC gene transcript or cDNA generated therefrom is present, that is, if both primers hybridize to sequences on the same molecule, the molecule comprising the primers and the intervening complementary sequences will be exponentially amplified. The amplified DNA can be easily detected by a variety of well known means. If no GCC gene transcript or cDNA generated therefrom is present, no PCR product will be  
20 exponentially amplified. The PCR technology therefore provides an extremely easy, straightforward and reliable method of detecting the GCC gene transcript in a sample.

PCR product may be detected by several well known means. The preferred method for detecting the presence of amplified DNA is to separate the PCR reaction material by gel electrophoresis and stain the gel with ethidium bromide in order to visual  
25 the amplified DNA if present. A size standard of the expected size of the amplified DNA is preferably run on the gel as a control.

In some instances, such as when unusually small amounts of RNA are recovered and only small amounts of cDNA are generated therefrom, it is desirable or necessary to perform a PCR reaction on the first PCR reaction product. That is, if difficult  
30 to detect quantities of amplified DNA are produced by the first reaction, a second PCR can be performed to make multiple copies of DNA sequences of the first amplified DNA. A

nested set of primers are used in the second PCR reaction. The nested set of primers hybridize to sequences downstream of the 5' primer and upstream of the 3' primer used in the first reaction.

The present invention includes oligonucleotide which are useful as primers  
5 for performing PCR methods to amplify the GCC gene transcript or cDNA generated therefrom.

According to the invention, diagnostic kits can be assembled which are useful to practice methods of detecting the presence of the GCC gene transcript or cDNA generated therefrom in non-colorectal samples. Such diagnostic kits comprise  
10 oligonucleotide which are useful as primers for performing PCR methods. It is preferred that diagnostic kits according to the present invention comprise a container comprising a size marker to be run as a standard on a gel used to detect the presence of amplified DNA. The size marker is the same size as the DNA generated by the primers in the presence of the GCC gene transcript or cDNA generated therefrom. Additional components in some  
15 kits include instructions for carrying out the assay. Additionally the kit may optionally comprise depictions or photographs that represent the appearance of positive and negative results. Positive and negative controls may also be provided.

PCR assays are useful for detecting the GCC gene transcript in homogenized tissue samples and cells in body fluid samples. It is contemplated that PCR on the plasma  
20 portion of a fluid sample could be used to detect the GCC gene transcript.

Another method of determining whether a sample contains cells expressing GCC is by branched chain oligonucleotide hybridization analysis of mRNA extracted from a sample. Branched chain oligonucleotide hybridization may be performed as described in U.S. Patent Number 5,597,909, U.S. Patent Number 5,437,977 and U.S. Patent Number  
25 5,430,138, which are each incorporated herein by reference. Reagents may be designed following the teachings of those patents and that sequence of the GCC gene transcript.

Another method of determining whether a sample contains cells expressing GCC is by Northern Blot analysis of mRNA extracted from a non-colorectal sample. The techniques for performing Northern blot analyses are well known by those having ordinary  
30 skill in the art and are described in Sambrook, J. *et al.*, (1989) *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY.

mRNA extraction, electrophoretic separation of the mRNA, blotting, probe preparation and hybridization are all well known techniques that can be routinely performed using readily available starting material.

The mRNA is extracted using poly dT columns and the material is separated  
5 by electrophoresis and, for example, transferred to nitrocellulose paper. Labeled probes made from an isolated specific fragment or fragments can be used to visualize the presence of a complementary fragment fixed to the paper. Probes useful to identify mRNA in a Northern Blot have a nucleotide sequence that is complementary to the GCC gene transcript. Those having ordinary skill in the art could use the sequence information in  
10 SEQ ID NO:1 to design such probes or to isolate and clone the GCC gene transcript or cDNA generated therefrom to be used as a probe. Such probes are at least 15 nucleotides, preferably 30-200, more preferably 40-100 nucleotide fragments and may be the entire GCC gene transcript.

According to the invention, diagnostic kits can be assembled which are  
15 useful to practice methods of detecting the presence of the GCC gene transcript in non-colorectal samples by Northern blot analysis. Such diagnostic kits comprise oligonucleotide which are useful as probes for hybridizing to the mRNA. The probes may be radiolabeled. It is preferred that diagnostic kits according to the present invention comprise a container comprising a size marker to be run as a standard on a gel. It is  
20 preferred that diagnostic kits according to the present invention comprise a container comprising a positive control which will hybridize to the probe. Additional components in some kits include instructions for carrying out the assay. Additionally the kit may optionally comprise depictions or photographs that represent the appearance of positive and negative results.

25 Northern blot analysis is useful for detecting the GCC gene transcript in homogenized tissue samples and cells in body fluid samples. It is contemplated that PCR on the plasma portion of a fluid sample could be used to detect the GCC gene transcript.

Another method of detecting the presence of the GCC gene transcript by oligonucleotide hybridization technology. Oligonucleotide hybridization technology is  
30 well known to those having ordinary skill in the art. Briefly, detectable probes which contain a specific nucleotide sequence that will hybridize to nucleotide sequence of the

GCC gene transcript. RNA or cDNA made from RNA from a sample is fixed, usually to filter paper or the like. The probes are added and maintained under conditions that permit hybridization only if the probes fully complement the fixed genetic material. The conditions are sufficiently stringent to wash off probes in which only a portion of the probe  
5 hybridizes to the fixed material. Detection of the probe on the washed filter indicate complementary sequences.

Probes useful in oligonucleotide assays at least 18 nucleotides of complementary DNA and may be as large as a complete complementary sequence to the GCC gene transcript. In some preferred embodiments the probes of the invention are 30-  
10 200 nucleotides, preferably 40-100 nucleotides.

One having ordinary skill in the art, using the sequence information disclosed in SEQ ID NO:1 can design probes useful in the invention. Hybridization conditions can be routinely optimized to minimize background signal by non-fully complementary hybridization. In some preferred embodiments, the probes are full length clones. Probes  
15 are at least 15 nucleotides, preferably 30-200, more preferably 40-100 nucleotide fragments and may be the entire GCC gene transcript.

The present invention includes labeled oligonucleotide which are useful as probes for performing oligonucleotide hybridization. The labeled probes of the present invention are labeled with radiolabeled nucleotides or are otherwise detectable by readily  
20 available nonradioactive detection systems.

According to the invention, diagnostic kits can be assembled which are useful to practice oligonucleotide hybridization methods of the invention. Such diagnostic kits comprise a labeled oligonucleotide which encodes portions of the GCC gene transcript. It is preferred that labeled probes of the oligonucleotide diagnostic kits according to the  
25 present invention are labeled with a radionucleotide. The oligonucleotide hybridization-based diagnostic kits according to the invention preferably comprise DNA samples that represent positive and negative controls. A positive control DNA sample is one that comprises a nucleic acid molecule which has a nucleotide sequence that is fully complementary to the probes of the kit such that the probes will hybridize to the molecule  
30 under assay conditions. A negative control DNA sample is one that comprises at least one nucleic acid molecule, the nucleotide sequence of which is partially complementary to the

sequences of the probe of the kit. Under assay conditions, the probe will not hybridize to the negative control DNA sample. Additional components in some kits include instructions for carrying out the assay. Additionally the kit may optionally comprise depictions or photographs that represent the appearance of positive and negative results.

- 5                   Oligonucleotide hybridization techniques are useful for detecting the GCC gene transcript in homogenized tissue samples and cells in body fluid samples. It is contemplated that PCR on the plasma portion of a fluid sample could be used to detect the GCC gene transcript.

- 10                   The present invention relates to *in vitro* kits for evaluating samples of tumors to determine whether or not they are stomach or esophageal in origin and to reagents and compositions useful to practice the same. In some embodiments of the invention, tumor samples may be isolated from individuals undergoing or recovery from surgery to remove tumors in the stomach or esophagus, tumors in other organs or biopsy material. The tumor sample is analyzed to identify the presence or absence of the GCC gene transcript.
- 15                   Techniques such as immunohistochemistry assays may be performed to determine whether GCC are present in cells in the tumor sample. The presence of mRNA that encodes the GCC protein or cDNA generated therefrom can be determined using techniques such as *in situ* hybridization, immunohistochemistry and *in situ* ST binding assay.

- 20                   *In situ* hybridization technology is well known by those having ordinary skill in the art. Briefly, cells are fixed and detectable probes which contain a specific nucleotide sequence are added to the fixed cells. If the cells contain complementary nucleotide sequences, the probes, which can be detected, will hybridize to them.

- 25                   Probes useful in oligonucleotide assays at least 18 nucleotides of complementary DNA and may be as large as a complete complementary sequence to the GCC gene transcript. In some preferred embodiments the probes of the invention are 30-200 nucleotides, preferably 40-100 nucleotides.

- 30                   One having ordinary skill in the art, using the sequence information set forth in SEQ ID NO:1 can design probes useful in *in situ* hybridization technology to identify cells that express GCC. Probes preferably hybridizes to a nucleotide sequence that corresponds to the GCC gene transcript. Hybridization conditions can be routinely optimized to minimize background signal by non-fully complementary hybridization.

Probes preferably hybridize to the full length GCC gene transcript. Probes are at least 15 nucleotides, preferably 30-200, more preferably 40-100 nucleotide fragments and may be the GCC gene transcript, more preferably 18-28 nucleotide fragments of the GCC gene transcript.

5                   The probes are fully complementary and do not hybridize well to partially complementary sequences. For *in situ* hybridization according to the invention, it is preferred that the probes are detectable by fluorescence. A common procedure is to label probe with biotin-modified nucleotide and then detect with fluorescently tagged avidin. Hence, probe does not itself have to be labeled with fluorescent but can be subsequently  
10 detected with fluorescent marker.

                  The present invention includes labeled oligonucleotide which are useful as probes for performing oligonucleotide hybridization. That is, they are fully complementary with mRNA sequences but not genomic sequences. The labeled probes of the present invention are labeled with radiolabeled nucleotides or are otherwise detectable  
15 by readily available nonradioactive detection systems.

                  The present invention relates to probes useful for *in situ* hybridization to identify cells that express GCC.

                  Cells are fixed and the probes are added to the genetic material. Probes will hybridize to the complementary nucleic acid sequences present in the sample. Using a  
20 fluorescent microscope, the probes can be visualized by their fluorescent markers.

                  According to the invention, diagnostic kits can be assembled which are useful to practice *in situ* hybridization methods of the invention are fully complementary with mRNA sequences but not genomic sequences. For example, the mRNA sequence includes different exon sequences. It is preferred that labeled probes of the *in situ*  
25 diagnostic kits according to the present invention are labeled with a fluorescent marker.

                  Immunohistochemistry techniques may be used to identify and essentially stain cells with GCC. Such "staining" allows for analysis of metastatic migration. Anti-GCC antibodies such as those described above of contacted with fixed cells and the GCC present in the cells reacts with the antibodies. The antibodies are detectably labeled or  
30 detected using labeled second antibody or protein A to stain the cells.



The techniques described herein for evaluating tumor sections can also be used to analyze tissue sections for samples of lymph nodes as well as other tissues to identify the presence of cells that express GCC. The samples can be prepared and "stained" to detect expression of GCC.

5               Immunoassay methods may be used in the diagnosis of individuals suffering from stomach or esophageal cancer by detecting presence of GCC in sample of non-colorectal tissue or body fluid from an individuals suspected of having or being susceptible to stomach or esophageal cancer using antibodies which were produced in response to exposure to such GCC protein. Moreover, immunoassay methods may be used to identify  
10 individuals suffering from stomach or esophageal cancer by detecting presence of GCC in sample of tumor using antibodies which were produced in response to exposure to such GCC protein.

The antibodies are preferably monoclonal antibodies. The antibodies are preferably raised against GCC made in human cells. Immunoassays are well known and  
15 there design may be routinely undertaken by those having ordinary skill in the art. Those having ordinary skill in the art can produce monoclonal antibodies which specifically bind to GCC and are useful in methods and kits of the invention using standard techniques and readily available starting materials. The techniques for producing monoclonal antibodies are outlined in Harlow, E. and D. Lane, (1988) *ANTIBODIES: A Laboratory Manual*, Cold  
20 Spring Harbor Laboratory, Cold Spring Harbor NY, which is incorporated herein by reference, provide detailed guidance for the production of hybridomas and monoclonal antibodies which specifically bind to target proteins. It is within the scope of the present invention to include Fabs, recombinant Fabs, F(Ab)<sub>2</sub>s, recombinant F(Ab)<sub>2</sub>s which specifically bind to GCC translation products in place of antibodies.

25               Briefly, GCC protein is injected into mice. The spleen of the mouse is removed, the spleen cells are isolated and fused with immortalized mouse cells. The hybrid cells, or hybridomas, are cultured and those cells which secrete antibodies are selected. The antibodies are analyzed and, if found to specifically bind to the GCC, the hybridoma which produces them is cultured to produce a continuous supply of anti-GCC  
30 specific antibodies.

The antibodies are preferably monoclonal antibodies. The antibodies are preferably raised against GCC made in human cells.

The means to detect the presence of a protein in a test sample are routine and one having ordinary skill in the art can detect the presence or absence of a protein or an antibody using well known methods. One well known method of detecting the presence of a protein is an immunoassay. One having ordinary skill in the art can readily appreciate the multitude of ways to practice an immunoassay to detect the presence of a GCC protein in a sample.

According to some embodiments, immunoassays comprise allowing proteins in the sample to bind a solid phase support such as a plastic surface. Detectable antibodies are then added which selectively binding to the GCC. Detection of the detectable antibody indicates the presence of GCC. The detectable antibody may be a labeled or an unlabeled antibody. Unlabeled antibody may be detected using a second, labeled antibody that specifically binds to the first antibody or a second, unlabeled antibody which can be detected using labeled protein A, a protein that complexes with antibodies. Various immunoassay procedures are described in *Immunoassays for the 80's*, A. Voller et al., Eds., University Park, 1981, which is incorporated herein by reference.

Simple immunoassays may be performed in which a solid phase support is contacted with the test sample. Any proteins present in the test sample bind the solid phase support and can be detected by a specific, detectable antibody preparation. Such a technique is the essence of the dot blot, Western blot and other such similar assays.

Other immunoassays may be more complicated but actually provide excellent results. Typical and preferred immunometric assays include "forward" assays for the detection of a protein in which a first anti-protein antibody bound to a solid phase support is contacted with the test sample. After a suitable incubation period, the solid phase support is washed to remove unbound protein. A second, distinct anti-protein antibody is then added which is specific for a portion of the specific protein not recognized by the first antibody. The second antibody is preferably detectable. After a second incubation period to permit the detectable antibody to complex with the specific protein bound to the solid phase support through the first antibody, the solid phase support is washed a second time to remove the unbound detectable antibody. Alternatively, the

second antibody may not be detectable. In this case, a third detectable antibody, which binds the second antibody is added to the system. This type of "forward sandwich" assay may be a simple yes/no assay to determine whether binding has occurred or may be made quantitative by comparing the amount of detectable antibody with that obtained in a control. Such "two-site" or "sandwich" assays are described by Wide, *Radioimmune Assay Method*, Kirkham, Ed., E. & S. Livingstone, Edinburgh, 1970; pp. 199-206, which is incorporated herein by reference.

Other types of immunometric assays are the so-called "simultaneous" and "reverse" assays. A simultaneous assay involves a single incubation step wherein the first antibody bound to the solid phase support, the second, detectable antibody and the test sample are added at the same time. After the incubation is completed, the solid phase support is washed to remove unbound proteins. The presence of detectable antibody associated with the solid support is then determined as it would be in a conventional "forward sandwich" assay. The simultaneous assay may also be adapted in a similar manner for the detection of antibodies in a test sample.

The "reverse" assay comprises the stepwise addition of a solution of detectable antibody to the test sample followed by an incubation period and the addition of antibody bound to a solid phase support after an additional incubation period. The solid phase support is washed in conventional fashion to remove unbound protein/antibody complexes and unreacted detectable antibody. The determination of detectable antibody associated with the solid phase support is then determined as in the "simultaneous" and "forward" assays. The reverse assay may also be adapted in a similar manner for the detection of antibodies in a test sample.

The first component of the immunometric assay may be added to nitrocellulose or other solid phase support which is capable of immobilizing proteins. The first component for determining the presence of GCC in a test sample is an anti-GCC antibody. By "solid phase support" or "support" is intended any material capable of binding proteins. Well-known solid phase supports include glass, polystyrene, polypropylene, polyethylene, dextran, nylon, amylases, natural and modified celluloses, polyacrylamides, agaroses, and magnetite. The nature of the support can be either soluble to some extent or insoluble for the purposes of the present invention. The support

configuration may be spherical, as in a bead, or cylindrical, as in the inside surface of a test tube or the external surface of a rod. Alternatively, the surface may be flat such as a sheet, test strip, etc. Those skilled in the art will know many other suitable "solid phase supports" for binding proteins or will be able to ascertain the same by use of routine experimentation.

- 5 A preferred solid phase support is a 96-well microtiter plate.

To detect the presence of GCC, detectable anti-GCC antibodies are used. Several methods are well known for the detection of antibodies.

- One method in which the antibodies can be detectably labeled is by linking the antibodies to an enzyme and subsequently using the antibodies in an enzyme
- 10 immunoassay (EIA) or enzyme-linked immunosorbent assay (ELISA), such as a capture ELISA. The enzyme, when subsequently exposed to its substrate, reacts with the substrate and generates a chemical moiety which can be detected, for example, by spectrophotometric, fluorometric or visual means. Enzymes which can be used to detectably label antibodies include, but are not limited to malate dehydrogenase,
- 15 staphylococcal nuclease, delta-5-steroid isomerase, yeast alcohol dehydrogenase, alpha-glycerophosphate dehydrogenase, triose phosphate isomerase, horseradish peroxidase, alkaline phosphatase, asparaginase, glucose oxidase, beta-galactosidase, ribonuclease, urease, catalase, glucose-6-phosphate dehydrogenase, glucoamylase and acetylcholinesterase. One skilled in the art would readily recognize other enzymes which
- 20 may also be used.

- Another method in which antibodies can be detectably labeled is through radioactive isotopes and subsequent use in a radioimmunoassay (RIA) (see, for example, Work, T.S. *et al.*, *Laboratory Techniques and Biochemistry in Molecular Biology*, North Holland Publishing Company, N.Y., 1978, which is incorporated herein by reference). The
- 25 radioactive isotope can be detected by such means as the use of a gamma counter or a scintillation counter or by autoradiography. Isotopes which are particularly useful for the purpose of the present invention are  $^3\text{H}$ ,  $^{125}\text{I}$ ,  $^{131}\text{I}$ ,  $^{35}\text{S}$ , and  $^{14}\text{C}$ . Preferably  $^{125}\text{I}$  is the isotope. One skilled in the art would readily recognize other radioisotopes which may also be used.

- It is also possible to label the antibody with a fluorescent compound. When
- 30 the fluorescent-labeled antibody is exposed to light of the proper wave length, its presence can be detected due to its fluorescence. Among the most commonly used fluorescent

labeling compounds are fluorescein isothiocyanate, rhodamine, phycoerythrin, phycocyanin, allophycocyanin, o-phthaldehyde and fluorescamine. One skilled in the art would readily recognize other fluorescent compounds which may also be used.

Antibodies can also be detectably labeled using fluorescence-emitting metals such as <sup>152</sup>Eu, or others of the lanthanide series. These metals can be attached to the protein-specific antibody using such metal chelating groups as diethylenetriaminepentaacetic acid (DTPA) or ethylenediamine-tetraacetic acid (EDTA). One skilled in the art would readily recognize other fluorescence-emitting metals as well as other metal chelating groups which may also be used.

Antibody can also be detectably labeled by coupling to a chemiluminescent compound. The presence of the chemiluminescent-labeled antibody is determined by detecting the presence of luminescence that arises during the course of a chemical reaction. Examples of particularly useful chemoluminescent labeling compounds are luminol, isoluminol, theromatic acridinium ester, imidazole, acridinium salt and oxalate ester. One skilled in the art would readily recognize other chemiluminescent compounds which may also be used.

Likewise, a bioluminescent compound may be used to label antibodies. Bioluminescence is a type of chemiluminescence found in biological systems in which a catalytic protein increases the efficiency of the chemiluminescent reaction. The presence of a bioluminescent protein is determined by detecting the presence of luminescence. Important bioluminescent compounds for purposes of labeling are luciferin, luciferase and aequorin. One skilled in the art would readily recognize other bioluminescent compounds which may also be used.

Detection of the protein-specific antibody, fragment or derivative may be accomplished by a scintillation counter if, for example, the detectable label is a radioactive gamma emitter. Alternatively, detection may be accomplished by a fluorometer if, for example, the label is a fluorescent material. In the case of an enzyme label, the detection can be accomplished by colorimetric methods which employ a substrate for the enzyme. Detection may also be accomplished by visual comparison of the extent of enzymatic reaction of a substrate in comparison with similarly prepared standards. One skilled in the

art would readily recognize other appropriate methods of detection which may also be used.

The binding activity of a given lot of antibodies may be determined according to well known methods. Those skilled in the art will be able to determine operative and optimal assay conditions for each determination by employing routine experimentation.

Positive and negative controls may be performed in which known amounts of GCC proteins and no GCC protein, respectively, are added to assays being performed in parallel with the test assay. One skilled in the art would have the necessary knowledge to perform the appropriate controls. In addition, the kit may comprise instructions for performing the assay. Additionally the kit may optionally comprise depictions or photographs that represent the appearance of positive and negative results.

GCC may be produced as a reagent for positive controls routinely. One skilled in the art would appreciate the different manners in which the GCC protein may be produced and isolated.

Antibody composition refers to the antibody or antibodies required for the detection of the protein. For example, the antibody composition used for the detection of a GCC in a test sample comprises a first antibody that binds to the GCC as well as a second or third detectable antibody that binds the first or second antibody, respectively.

To examine a test sample for the presence of a GCC, a standard immunometric assay such as the one described below may be performed. A first anti-GCC antibody, which recognizes a specific portion of GCC, is added to a 96-well microtiter plate in a volume of buffer. The plate is incubated for a period of time sufficient for binding to occur and subsequently washed with PBS to remove unbound antibody. The plate is then blocked with a PBS/BSA solution to prevent sample proteins from non-specifically binding the microtiter plate. Test sample are subsequently added to the wells and the plate is incubated for a period of time sufficient for binding to occur. The wells are washed with PBS to remove unbound protein. Labeled anti-GCC antibodies, which recognize portions of GCC not recognized by the first antibody, are added to the wells. The plate is incubated for a period of time sufficient for binding to occur and subsequently

washed with PBS to remove unbound, labeled anti-GCC antibody. The amount of labeled and bound anti-GCC antibody is subsequently determined by standard techniques.

Kits which are useful for the detection of GCC in a test sample comprise a container comprising anti-GCC antibodies and a container or containers comprising  
5 controls. Controls include one control sample which does not contain GCC and/or another control sample which contained the GCC. The anti-GCC antibodies used in the kit are detectable such as being detectably labeled. If the detectable anti-GCC antibody is not labeled, it may be detected by second antibodies or protein A for example which may also be provided in some kits in separate containers. Additional components in some kits  
10 include solid support, buffer, and instructions for carrying out the assay. Additionally the kit may optionally comprise depictions or photographs that represent the appearance of positive and negative results.

The immunoassay is useful for detecting GCC in homogenized tissue samples and body fluid samples including the plasma portion or cells in the fluid sample.

15 Western Blots may be useful in assisting the diagnosis of individuals suffering from stomach or esophageal cancer by detecting presence of GCC of non-colorectal tissue or body fluid. Western blots may also be used to detect presence of GCC in sample of tumor from an individual suffering from cancer. Western blots use detectable anti-GCC - antibodies to bind to any GCC present in a sample and thus indicate the  
20 presence of the receptor in the sample.

Western blot techniques, which are described in Sambrook, J. *et al.*, (1989) *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, which is incorporated herein by reference, are similar to immunoassays with the essential difference being that prior to exposing the sample to the  
25 antibodies, the proteins in the samples are separated by gel electrophoresis and the separated proteins are then probed with antibodies. In some preferred embodiments, the matrix is an SDS-PAGE gel matrix and the separated proteins in the matrix are transferred to a carrier such as filter paper prior to probing with antibodies. Anti-GCC antibodies described above are useful in Western blot methods.

30 Generally, samples are homogenized and cells are lysed using detergent such as Triton-X. The material is then separated by the standard techniques in Sambrook, J. *et*

*al.*, (1989) *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY.

Kits which are useful for the detection of GCC in a test sample by Western Blot comprise a container comprising anti-GCC antibodies and a container or containers  
5 comprising controls. Controls include one control sample which does not contain GCC and/or another control sample which contains GCC. The anti-GCC antibodies used in the kit are detectable such as being detectably labeled. If the detectable anti-GCC antibody is not labeled, it may be detected by second antibodies or protein A for example which may also be provided in some kits in separate containers. Additional components in some kits  
10 include instructions for carrying out the assay. Additionally the kit may optionally comprise depictions or photographs that represent the appearance of positive and negative results.

Western blots are useful for detecting GCC in homogenized tissue samples and body fluid samples including the plasma portion or cells in the fluid sample.

## 15 ***In vivo* Imaging and Therapeutics**

According to some embodiments of the invention, compositions and *in vivo* methods are provided for detecting, imaging, or treating primary and/or metastatic stomach or esophageal tumors in an individual.

When the conjugated compositions of the present invention are administered  
20 outside the intestinal tract such as when administered in the circulatory system, they remain segregated from the cells that line the intestinal tract and will bind only to cells outside the intestinal tract which express GCC. The conjugated compositions will not bind to the normal cells but will bind to primary and/or metastatic stomach or esophageal cells. Thus, the active moieties of conjugated compositions administered outside the intestinal  
25 tract are delivered to cells which express GCC such as primary and/or metastatic stomach or esophageal cancer cells.

Therapeutic and diagnostic pharmaceutical compositions useful in the present invention include conjugated compounds that specifically target cells that express GCC. These conjugated compounds include moieties that bind to GCC which do not bind to cells



of normal tissue in the body except cells of the intestinal tract since the cells of other tissues do not express GCC.

Unlike normal colorectal cells, cancer cells that express GCC are accessible to substances administered outside the intestinal tract, for example administered in the  
5 circulatory system. The only GCC in normal tissue exist in the apical membranes of intestinal mucosa cells and thus effectively isolated from the targeted cancer chemotherapeutics and imaging agents administered outside the intestinal tract by the intestinal mucosa barrier. Thus, primary and/or metastatic stomach or esophageal cancer cells may be targeted by conjugated compounds of the present invention by introducing  
10 such compounds outside the intestinal tract such as for example by administering pharmaceutical compositions that comprise conjugated compounds into the circulatory system.

One having ordinary skill in the art can identify individuals suspected of suffering from primary and/or metastatic stomach or esophageal cancer. In those  
15 individuals diagnosed with stomach or esophageal cancer, it is not unusual and in some cases standard therapy to suspect metastasis and aggressively attempt to eradicate metastasized cells. The present invention provides pharmaceutical compositions and methods for imaging and thereby will more definitively diagnose primary and metastatic disease. Further, the present invention provides pharmaceutical compositions comprising  
20 therapeutic agents and methods for specifically targeting and eliminating primary and/or metastatic stomach or esophageal cancer cells. Further, the present invention provides pharmaceutical compositions that comprise therapeutics and methods for specifically eliminating primary and/or metastatic stomach or esophageal cancer cells.

The pharmaceutical compositions which comprise conjugated compositions  
25 of the present invention may be used to diagnose or treat individuals suffering from primary and/or metastatic stomach or esophageal tumors.

The present invention relies upon the use of a GCC binding moiety in a conjugated composition. The GCC binding moiety is essentially a portion of the conjugated composition which acts as a ligand to a GCC and thus specifically binds to it.  
30 The conjugated composition also includes an active moiety which is associated with the

GCC binding moiety; the active moiety being an active agent which is either useful to image, target, neutralize or kill the cell.

According to the present invention, the GCC binding moiety is the GCC ligand portion of a conjugated composition. In some embodiments, the GCC ligand is an antibody.

In some preferred embodiments, conjugated compounds comprise GCC binding moieties that comprise an anti-GCC antibody.

It is preferred that the GCC ligand used as the GCC binding moiety be as small as possible. Thus it is preferred that the GCC ligand be a non-peptide small molecule or small peptide, preferably less than 25 amino acids, more preferably less than 20 amino acids. In some embodiments, the GCC ligand which constitute the GCC binding moiety of a conjugated composition is less than 15 amino acids. GCC binding peptide comprising less than 10 amino acids and GCC binding peptide less than 5 amino acids may be used as GCC binding moieties according to the present invention. It is within the scope of the present invention to include larger molecules which serve as GCC binding moieties including, but not limited to molecules such as antibodies which specifically bind to GCC.

GCC ligands useful as GCC binding moieties may be identified using various well known combinatorial library screening technologies such as those set forth in Example 1 herein.

An assay may be used to test both peptide and non-peptide compositions to determine whether or not they are GCC ligands or, to test conjugated compositions to determine if they possess GCC - binding activity. Such compositions that specifically bind to GCC can be identified by a competitive binding assay using antibodies known to bind to the GCC. The competitive binding assay is a standard technique in pharmacology which can be readily performed by those having ordinary skill in the art using readily available starting materials.

GCC may be produced synthetically, recombinantly or isolated from natural sources.

Using a solid phase synthesis as an example, the protected or derivatized amino acid is attached to an inert solid support through its unprotected carboxyl or amino group. The protecting group of the amino or carboxyl group is then selectively removed

and the next amino acid in the sequence having the complementary (amino or carboxyl) group suitably protected is admixed and reacted with the residue already attached to the solid support. The protecting group of the amino or carboxyl group is then removed from this newly added amino acid residue, and the next amino acid (suitably protected) is then  
5 added, and so forth. After all the desired amino acids have been linked in the proper sequence, any remaining terminal and side group protecting groups (and solid support) are removed sequentially or concurrently, to provide the final peptide. The peptide of the invention are preferably devoid of benzylated or methylbenzylated amino acids. Such protecting group moieties may be used in the course of synthesis, but they are removed  
10 before the peptides are used. Additional reactions may be necessary, as described elsewhere, to form intramolecular linkages to restrain conformation.

Antibodies against GCC may be routinely produced and used in competition assays to identify GCC ligands or as starting materials for conjugated compounds according to the invention.

15 According to the present invention, the active moiety may be a therapeutic agent or an imaging agent. One having ordinary skill in the art can readily recognize the advantages of being able to specifically target cancer cells with an GCC ligand and conjugate such a ligand with many different active agents.

Chemotherapeutics useful as active moieties which when conjugated to a  
20 GCC binding moiety are specifically delivered to cells that express GCC such as stomach or esophageal cancer cells, are typically small chemical entities produced by chemical synthesis. Chemotherapeutics include cytotoxic and cytostatic drugs. Chemotherapeutics may include those which have other effects on cells such as reversal of the transformed state to a differentiated state or those which inhibit cell replication. Examples of  
25 chemotherapeutics include common cytotoxic or cytostatic drugs such as for example: methotrexate (amethopterin), doxorubicin (adrimycin), daunorubicin, cytosinarabioside, etoposide, 5-4 fluorouracil, melphalan, chlorambucil, and other nitrogen mustards (e.g. cyclophosphamide), *cis*-platinum, vindesine (and other vinca alkaloids), mitomycin and bleomycin. Other chemotherapeutics include: purothionin (barley flour oligopeptide),  
30 macromomycin. 1,4-benzoquinone derivatives and trenimon.

Toxins are useful as active moieties. When a toxin is conjugated to a GCC binding moiety, the conjugated composition is specifically delivered to a cell that expresses GCC such as stomach or esophageal cancer cells by way of the GCC binding moiety and the toxin moiety kills the cell. Toxins are generally complex toxic products of various organisms including bacteria, plants, *etc.* Examples of toxins include but are not limited to: ricin, ricin A chain (ricin toxin), *Pseudomonas* exotoxin (PE), diphtheria toxin (DT), *Clostridium perfringens* phospholipase C (PLC), bovine pancreatic ribonuclease (BPR), pokeweed antiviral protein (PAP), abrin, abrin A chain (abin toxin), cobra venom factor (CVF), gelonin (GEL), saporin (SAP), modeccin, viscumin and volkensin. As discussed above, when protein toxins are employed with GCC binding peptides, conjugated compositions may be produced using recombinant DNA techniques. Briefly, a recombinant DNA molecule can be constructed which encodes both the GCC ligand and the toxin on a chimeric gene. When the chimeric gene is expressed, a fusion protein is produced which includes a GCC binding moiety and an active moiety. Protein toxins are also useful to form conjugated compounds with GCC binding peptides through non-peptidyl bonds.

In addition, there are other approaches for utilizing active agents for the treatment of cancer. For example, conjugated compositions may be produced which include a GCC binding moiety and an active moiety which is an active enzyme. The GCC binding moiety specifically localizes the conjugated composition to the tumor cells. An inactive prodrug which can be converted by the enzyme into an active drug is administered to the patient. The prodrug is only converted to an active drug by the enzyme which is localized to the tumor. An example of an enzyme/prodrug pair includes alkaline phosphatase/etoposidephosphate. In such a case, the alkaline phosphatase is conjugated to a GCC binding ligand. The conjugated compound is administered and localizes at the cancer cell. Upon contact with etoposidephosphate (the prodrug), the etoposidephosphate is converted to etoposide, a chemotherapeutic drug which is taken up by the cancer cell.

Radiosensitizing agents are substances that increase the sensitivity of cells to radiation. Examples of radiosensitizing agents include nitroimidazoles, metronidazole and misonidazole (see: DeVita, V.T. Jr. in *Harrison's Principles of Internal Medicine*, p.68, McGraw-Hill Book Co., N.Y. 1983, which is incorporated herein by reference). The

conjugated compound that comprises a radiosensitizing agent as the active moiety is administered and localizes at the primary and/or metastatic stomach or esophageal cancer cell. Upon exposure of the individual to radiation, the radiosensitizing agent is "excited" and causes the death of the cell.

5 Radionuclides may be used in pharmaceutical compositions that are useful for radiotherapy or imaging procedures.

Examples of radionuclides useful as toxins in radiation therapy include:  $^{47}\text{Sc}$ ,  $^{67}\text{Cu}$ ,  $^{90}\text{Y}$ ,  $^{109}\text{Pd}$ ,  $^{123}\text{I}$ ,  $^{125}\text{I}$ ,  $^{131}\text{I}$ ,  $^{186}\text{Re}$ ,  $^{188}\text{Re}$ ,  $^{199}\text{Au}$ ,  $^{211}\text{At}$ ,  $^{212}\text{Pb}$  and  $^{212}\text{Bi}$ . Other radionuclides which have been used by those having ordinary skill in the art include:  $^{32}\text{P}$  and  $^{33}\text{P}$ ,  $^{71}\text{Ge}$ ,  
 10  $^{77}\text{As}$ ,  $^{103}\text{Pb}$ ,  $^{105}\text{Rh}$ ,  $^{111}\text{Ag}$ ,  $^{119}\text{Sb}$ ,  $^{121}\text{Sn}$ ,  $^{131}\text{Cs}$ ,  $^{143}\text{Pr}$ ,  $^{161}\text{Tb}$ ,  $^{177}\text{Lu}$ ,  $^{191}\text{Os}$ ,  $^{193}\text{Pt}$ ,  $^{197}\text{Hg}$ , all beta negative and/or auger emitters. Some preferred radionuclides include:  $^{90}\text{Y}$ ,  $^{131}\text{I}$ ,  $^{211}\text{At}$  and  $^{212}\text{Pb}/^{212}\text{Bi}$ .

According to the present invention, the active moieties may be an imaging agent. Imaging agents are useful diagnostic procedures as well as the procedures used to  
 15 identify the location of cancer cells. Imaging can be performed by many procedures well-known to those having ordinary skill in the art and the appropriate imaging agent useful in such procedures may be conjugated to a GCC ligand by well-known means. Imaging can be performed, for example, by radioscintigraphy, nuclear magnetic resonance imaging (MRI) or computed tomography (CT scan). The most commonly employed radionuclide  
 20 imaging agents include radioactive iodine and indium. Imaging by CT scan may employ a heavy metal such as iron chelates. MRI scanning may employ chelates of gadolinium or manganese. Additionally, positron emission tomography (PET) may be possible using positron emitters of oxygen, nitrogen, iron, carbon, or gallium. Example of radionuclides useful in imaging procedures include:  $^{43}\text{K}$ ,  $^{52}\text{Fe}$ ,  $^{57}\text{Co}$ ,  $^{67}\text{Cu}$ ,  $^{67}\text{Ga}$ ,  $^{68}\text{Ga}$ ,  $^{77}\text{Br}$ ,  $^{81}\text{Rb}/^{81\text{m}}\text{Kr}$ ,  
 25  $^{87\text{m}}\text{Sr}$ ,  $^{99\text{m}}\text{Tc}$ ,  $^{111}\text{In}$ ,  $^{113\text{m}}\text{In}$ ,  $^{123}\text{I}$ ,  $^{125}\text{I}$ ,  $^{127}\text{Cs}$ ,  $^{129}\text{Cs}$ ,  $^{131}\text{I}$ ,  $^{132}\text{I}$ ,  $^{197}\text{Hg}$ ,  $^{203}\text{Pb}$  and  $^{206}\text{Bi}$ .

It is preferred that the conjugated compositions be non-immunogenic or immunogenic at a very low level. Accordingly, it is preferred that the GCC binding moiety be a small, poorly immunogenic or non-immunogenic peptide or a non-peptide. The GCC binding moiety may be a humanized or primatized antibody or a human  
 30 antibody.

GCC ligands are conjugated to active agents by a variety of well-known techniques readily performed without undue experimentation by those having ordinary skill in the art. The technique used to conjugate the GCC ligand to the active agent is dependent upon the molecular nature of the GCC ligand and the active agent. After the GCC ligand and the active agent are conjugated to form a single molecule, assays may be performed to ensure that the conjugated molecule retains the activities of the moieties. The competitive binding assay described above may be used to confirm that the GCC binding moiety retains its binding activity as a conjugated compound. Similarly, the activity of the active moiety may be tested using various assays for each respective type of active agent.

Radionuclides retain their activity, i.e. their radioactivity, irrespective of conjugation. With respect to active agents which are toxins, drugs and targeting agents, standard assays to demonstrate the activity of unconjugated forms of these compounds may be used to confirm that the activity has been retained.

Conjugation may be accomplished directly between the GCC ligand and the active agent or linking, intermediate molecular groups may be provided between the GCC ligand and the active agent. Crosslinkers are particularly useful to facilitate conjugation by providing attachment sites for each moiety. Crosslinkers may include additional molecular groups which serve as spacers to separate the moieties from each other to prevent either from interfering with the activity of the other.

One having ordinary skill in the art may conjugate a GCC ligand to a chemotherapeutic drug using well-known techniques. For example, Magerstadt, M. *Antibody Conjugates and Malignant Disease*. (1991) CRC Press, Boca Raton, USA, pp. 110-152) which is incorporated herein by reference, teaches the conjugation of various cytostatic drugs to amino acids of antibodies. Such reactions may be applied to conjugate chemotherapeutic drugs to GCC ligands, including anti-GCC antibodies, with an appropriate linker. Most of the chemotherapeutic agents currently in use in treating cancer possess functional groups that are amenable to chemical crosslinking directly with proteins. For example, free amino groups are available on methotrexate, doxorubicin, daunorubicin, cytosinarabioside, *cis*-platin, vindesine, mitomycin and bleomycin while free carboxylic acid groups are available on methotrexate, melphalan, and chlorambucil. These functional groups, that is free amino and carboxylic acids, are targets for a variety of

homobifunctional and heterobifunctional chemical crosslinking agents which can crosslink these drugs directly to the single free amino group of an antibody. For example, one procedure for crosslinking GCC ligands which have a free amino group to active agents which have a free amino group such as methotrexate, doxorubicin, daunorubicin, cytosinarabioside, *cis*-platin, vindesine, mitomycin and bleomycin, or alkaline phosphatase, or protein- or peptide-based toxin employs homobifunctional succinimidyl esters, preferably with carbon chain spacers such as disuccinimidyl suberate (Pierce Co, Rockford, IL). In the event that a cleavable conjugated compound is required, the same protocol would be employed utilizing 3,3'- dithiobis (sulfosuccinimidylpropionate; Pierce Co.).

In order to conjugate a GCC ligand that is a peptide or protein to a peptide-based active agent such as a toxin, the GCC ligand and the toxin may be produced as a single, fusion protein either by standard peptide synthesis or recombinant DNA technology, both of which can be routinely performed by those having ordinary skill in the art. Alternatively, two peptides, the GCC ligand peptide and the peptide-based toxin may be produced and/or isolated as separate peptides and conjugated using crosslinkers. As with conjugated compositions that contain chemotherapeutic drugs, conjugation of GCC binding peptides and toxins can exploit the ability to modify the single free amino group of a GCC binding peptide while preserving the receptor-binding function of this molecule.

One having ordinary skill in the art may conjugate a GCC ligand to a radionuclide using well-known techniques. For example, Magerstadt, M. (1991) *Antibody Conjugates And Malignant Disease*, CRC Press, Boca Raton, FLA.; and Barchel, S.W. and Rhodes, B.H., (1983) *Radioimaging and Radiotherapy*, Elsevier, NY, NY, each of which is incorporated herein by reference, teach the conjugation of various therapeutic and diagnostic radionuclides to amino acids of antibodies.

The present invention provides pharmaceutical compositions that comprise the conjugated compounds of the invention and pharmaceutically acceptable carriers or diluents. The pharmaceutical composition of the present invention may be formulated by one having ordinary skill in the art. Suitable pharmaceutical carriers are described in *Remington's Pharmaceutical Sciences*, A. Osol, a standard reference text in this field, which is incorporated herein by reference. In carrying out methods of the present

invention, conjugated compounds of the present invention can be used alone or in combination with other diagnostic, therapeutic or additional agents. Such additional agents include excipients such as coloring, stabilizing agents, osmotic agents and antibacterial agents. Pharmaceutical compositions are preferably sterile and pyrogen free.

- 5                   The conjugated compositions of the invention can be, for example, formulated as a solution, suspension or emulsion in association with a pharmaceutically acceptable parenteral vehicle. Examples of such vehicles are water, saline, Ringer's solution, dextrose solution, and 5% human serum albumin. Liposomes may also be used. The vehicle may contain additives that maintain isotonicity (e.g., sodium chloride,
- 10 mannitol) and chemical stability (e.g., buffers and preservatives). The formulation is sterilized by commonly used techniques. For example, a parenteral composition suitable for administration by injection is prepared by dissolving 1.5% by weight of active ingredient in 0.9% sodium chloride solution.

- The pharmaceutical compositions according to the present invention may be
- 15 administered as either a single dose or in multiple doses. The pharmaceutical compositions of the present invention may be administered either as individual therapeutic agents or in combination with other therapeutic agents. The treatments of the present invention may be combined with conventional therapies, which may be administered sequentially or simultaneously.

- 20                   The pharmaceutical compositions of the present invention may be administered by any means that enables the conjugated composition to reach the targeted cells. In some embodiments, routes of administration include those selected from the group consisting of intravenous, intraarterial, intraperitoneal, local administration into the blood supply of the organ in which the tumor resides or directly into the tumor itself.
- 25 Intravenous administration is the preferred mode of administration. It may be accomplished with the aid of an infusion pump. In addition to an intraoperative spray, conjugated compounds may be delivered intrathecally, intraventrically, stereotactically, intrahepatically such as via the portal vein, by inhalation, and intrapleurally.

- The dosage administered varies depending upon factors such as: the nature of
- 30 the active moiety; the nature of the conjugated composition; pharmacodynamic characteristics; its mode and route of administration; age, health, and weight of the



recipient; nature and extent of symptoms; kind of concurrent treatment; and frequency of treatment.

Because conjugated compounds are specifically targeted to cells with one or more GCC molecules, conjugated compounds which comprise chemotherapeutics or toxins are administered in doses less than those which are used when the chemotherapeutics or toxins are administered as unconjugated active agents, preferably in doses that contain up to 100 times less active agent. In some embodiments, conjugated compounds which comprise chemotherapeutics or toxins are administered in doses that contain 10-100 times less active agent as an active moiety than the dosage of chemotherapeutics or toxins administered as unconjugated active agents. To determine the appropriate dose, the amount of compound is preferably measured in moles instead of by weight. In that way, the variable weight of different GCC binding moieties does not affect the calculation. Presuming a one to one ratio of GCC binding moiety to active moiety in conjugated compositions of the invention, less moles of conjugated compounds may be administered as compared to the moles of unconjugated compounds administered, preferably up to 100 times less moles.

Typically, chemotherapeutic conjugates are administered intravenously in multiple divided doses.

Up to 20 gm IV/dose of methotrexate is typically administered in an unconjugated form. When methotrexate is administered as the active moiety in a conjugated compound of the invention, there is a 10-to 100-fold dose reduction. Thus, presuming each conjugated compound includes one molecule of methotrexate conjugated to one GCC binding moiety, of the total amount of conjugated compound administered, up to about 0.2 - 2.0 g of methotrexate is present and therefore administered. In some embodiments, of the total amount of conjugated compound administered, up to about 200 mg - 2g of methotrexate is present and therefore administered.

To dose conjugated compositions comprising GCC binding moieties linked to active moieties that are radioisotopes in pharmaceutical compositions useful as imaging agents, it is presumed that each GCC binding moiety is linked to one radioactive active moiety. The amount of radioisotope to be administered is dependent upon the radioisotope. Those having ordinary skill in the art can readily formulate the amount of

conjugated compound to be administered based upon the specific activity and energy of a given radionuclide used as an active moiety. Typically 0.1-100 millicuries per dose of imaging agent, preferably 1-10 millicuries, most often 2-5 millicuries are administered. Thus, pharmaceutical compositions according to the present invention useful as imaging agents which comprise conjugated compositions comprising a GCC binding moiety and a radioactive moiety comprise 0.1-100 millicuries, in some embodiments preferably 1-10 millicuries, in some embodiments preferably 2-5 millicuries, in some embodiments more preferably 1-5 millicuries. Examples of dosages include:  $^{131}\text{I}$  = between about 0.1-100 millicuries per dose, in some embodiments preferably 1-10 millicuries, in some 10 embodiments 2-5 millicuries, and in some embodiments about 4 millicuries;  $^{111}\text{In}$  = between about 0.1-100 millicuries per dose, in some embodiments preferably 1-10 millicuries, in some embodiments 1-5 millicuries, and in some embodiments about 2 millicuries;  $^{99\text{m}}\text{Tc}$  = between about 0.1-100 millicuries per dose, in some embodiments preferably 5-75 millicuries, in some embodiments 10-50 millicuries, and in some 15 embodiments about 27 millicuries. Wessels B.W. and R.D. Rogus (1984) *Med. Phys.* 11:638 and Kwok, C.S. *et al.* (1985) *Med. Phys.* 12:405, both of which are incorporated herein by reference, disclose detailed dose calculations for diagnostic and therapeutic conjugates which may be used in the preparation of pharmaceutical compositions of the present invention which include radioactive conjugated compounds.

20 One aspect of the present invention relates to a method of treating individuals suspected of suffering from primary and/or metastatic stomach or esophageal cancer. Such individuals may be treated by administering to the individual a pharmaceutical composition that comprises a pharmaceutically acceptable carrier or diluent and a conjugated compound that comprises a GCC - binding moiety and an active moiety 25 wherein the active moiety is a radiostable therapeutic agent. In some embodiments of the present invention, the pharmaceutical composition comprises a pharmaceutically acceptable carrier or diluent and a conjugated compound that comprises a GCC binding moiety and an active moiety wherein the active moiety is a radiostable active agent and the GCC binding moiety is an antibody. In some embodiments of the present invention, the 30 pharmaceutical composition comprises a pharmaceutically acceptable carrier or diluent and a conjugated compound that comprises a GCC binding moiety and an active moiety

wherein the active moiety is a radiostable therapeutic agent. In some embodiments of the present invention, the pharmaceutical composition comprises a pharmaceutically acceptable carrier or diluent and a conjugated compound that comprises a GCC binding moiety and an active moiety wherein the active moiety is a radiostable active agent

5 selected from the group consisting of: methotrexate, doxorubicin, daunorubicin, cytosinarabioside, etoposide, 5-4 fluorouracil, melphalan, chlorambucil, *cis*-platinum, vindesine, mitomycin, bleomycin, purothionin, macromomycin, 1,4-benzoquinone derivatives, trenimon, ricin, ricin A chain, *Pseudomonas* exotoxin, diphtheria toxin, *Clostridium perfringens* phospholipase C, bovine pancreatic ribonuclease, pokeweed

10 antiviral protein, abrin, abrin A chain, cobra venom factor, gelonin, saporin, modeccin, viscumin, volkensin, alkaline phosphatase, nitroimidazole, metronidazole and misonidazole. The individual being treated may be diagnosed as having metastasized stomach or esophageal cancer or may be diagnosed as having primary stomach or esophageal cancer and may undergo the treatment proactively in the event that there is

15 some metastasis as yet undetected. The pharmaceutical composition contains a therapeutically effective amount of the conjugated composition. A therapeutically effective amount is an amount which is effective to cause a cytotoxic or cytostatic effect on cancer cells without causing lethal side effects on the individual.

One aspect of the present invention relates to a method of treating individuals

20 suspected of suffering from primary and/or metastatic stomach or esophageal cancer. Such individuals may be treated by administering to the individual a pharmaceutical composition that comprises a pharmaceutically acceptable carrier or diluent and a conjugated compound that comprises a GCC binding moiety and an active moiety wherein the active moiety is a radioactive. In some embodiments of the present invention, the

25 pharmaceutical composition comprises a pharmaceutically acceptable carrier or diluent and a conjugated compound that comprises a GCC binding moiety and an active moiety wherein the active moiety is a radioactive and the GCC binding moiety is an antibody. In some embodiments of the present invention, the pharmaceutical composition comprises a pharmaceutically acceptable carrier or diluent and a conjugated compound that comprises a

30 GCC binding moiety and an active moiety wherein the active moiety is a radioactive agent selected from the group consisting of:  $^{47}\text{Sc}$ ,  $^{67}\text{Cu}$ ,  $^{90}\text{Y}$ ,  $^{109}\text{Pd}$ ,  $^{123}\text{I}$ ,  $^{125}\text{I}$ ,  $^{131}\text{I}$ ,  $^{186}\text{Re}$ ,  $^{188}\text{Re}$ ,

<sup>199</sup>Au, <sup>211</sup>At, <sup>212</sup>Pb, <sup>212</sup>B, <sup>32</sup>P and <sup>33</sup>P, <sup>71</sup>Ge, <sup>77</sup>As, <sup>103</sup>Pb, <sup>105</sup>Rh, <sup>111</sup>Ag, <sup>119</sup>Sb, <sup>121</sup>Sn, <sup>131</sup>Cs, <sup>143</sup>Pr, <sup>161</sup>Tb, <sup>177</sup>Lu, <sup>191</sup>Os, <sup>193</sup>Mt, <sup>197</sup>Hg, <sup>32</sup>P and <sup>33</sup>P, <sup>71</sup>Ge, <sup>77</sup>As, <sup>103</sup>Pb, <sup>105</sup>Rh, <sup>111</sup>Ag, <sup>119</sup>Sb, <sup>121</sup>Sn, <sup>131</sup>Cs, <sup>143</sup>Pr, <sup>161</sup>Tb, <sup>177</sup>Lu, <sup>191</sup>Os, <sup>193</sup>Mt, <sup>197</sup>Hg, all beta negative and/or auger emitters. The individual being treated may be diagnosed as having metastasized cancer or may be

5 diagnosed as having localized cancer and may undergo the treatment proactively in the event that there is some metastasis as yet undetected. The pharmaceutical composition contains a therapeutically effective amount of the conjugated composition. A therapeutically effective amount is an amount which is effective to cause a cytotoxic or cytostatic effect on primary and/or metastatic stomach or esophageal cancer cells without

10 causing lethal side effects on the individual. The composition may be injected intratumorally into primary tumors.

One aspect of the present invention relates to a method of detecting primary and/or metastatic stomach or esophageal cancer cells in an individual suspected of suffering from primary or metastasized stomach or esophageal cancer by radioimaging.

15 Individuals may be suspected of having primary stomach or esophageal tumors which diagnosis can be confirmed by administering to the individual, an imaging agent which binds to GCC. Tumors can be imaged by detecting localization at the stomach or esophagus. Individuals may be diagnosed as suffering from metastasized stomach or esophageal cancer and the metastasized stomach or esophageal cancer cells may be

20 detected by administering to the individual, preferably by intravenous administration, a pharmaceutical composition that comprises a pharmaceutically acceptable carrier or diluent and a conjugated compound that comprises a GCC binding moiety and an active moiety wherein the active moiety is a radioactive and detecting the presence of a localized accumulation or aggregation of radioactivity, indicating the presence of cells with GCC. In

25 some embodiments of the present invention, the pharmaceutical composition comprises a pharmaceutically acceptable carrier or diluent and a conjugated compound that comprises a GCC binding moiety and an active moiety wherein the active moiety is a radioactive and the GCC binding moiety is an antibody. In some embodiments of the present invention, the pharmaceutical composition comprises a pharmaceutically acceptable carrier or diluent

30 and a conjugated compound that comprises an GCC binding moiety and an active moiety wherein the active moiety is a radioactive agent selected from the group consisting of:

radioactive heavy metals such as iron chelates, radioactive chelates of gadolinium or manganese, positron emitters of oxygen, nitrogen, iron, carbon, or gallium,  $^{43}\text{K}$ ,  $^{52}\text{Fe}$ ,  $^{57}\text{Co}$ ,  $^{67}\text{Cu}$ ,  $^{67}\text{Ga}$ ,  $^{68}\text{Ga}$ ,  $^{77}\text{Br}$ ,  $^{81}\text{Rb}/^{81\text{M}}\text{Kr}$ ,  $^{87\text{M}}\text{Sr}$ ,  $^{99\text{M}}\text{Tc}$ ,  $^{111}\text{In}$ ,  $^{113\text{M}}\text{In}$ ,  $^{123}\text{I}$ ,  $^{125}\text{I}$ ,  $^{127}\text{Cs}$ ,  $^{129}\text{Cs}$ ,  $^{131}\text{I}$ ,  $^{132}\text{I}$ ,  $^{197}\text{Hg}$ ,  $^{203}\text{Pb}$  and  $^{206}\text{Bi}$ . The individual being treated may be diagnosed as having

5 metastasizing stomach or esophageal cancer or may be diagnosed as having localized stomach or esophageal cancer and may undergo the treatment proactively in the event that there is some metastasis as yet undetected. The pharmaceutical composition contains a diagnostically effective amount of the conjugated composition. A diagnostically effective amount is an amount which can be detected at a site in the body where cells with GCC are

10 located without causing lethal side effects on the individual.

### **Photodynamic imaging and therapy**

According to some embodiments of the invention, GCC binding moieties are conjugates to photoactivated imaging agents or therapeutics. Maier A. et al. Lasers in Surgery and Medicine 26:461-466 (2000) which is incorporated herein by reference

15 disclose an example of photodynamic therapy. QLT, Inc (Vancouver, BC) commercially distribute photosensitive active agents which can be linked to GCC ligands. Such conjugated compounds can be used in photodynamic therapeutic and imaging protocols to activate the GCC-bound conjugated agents which are thus targeted to tumor cells. In some embodiments, the conjugated compounds are applied as an intraoperative spray which is

20 subsequently exposed to light to activate compounds bound to cells that express GCC.

In some embodiments, the photodynamic agent is fluorophore or porphyrins. Examples of porphyrin include: hematoporphyrin derivative (HPD) and porfimer sodium (Photofrin®). A second generation photosensitizers is BPD verteporfin. In some

embodiments the fluorophore is tetramethylrotamine. Lasers are generally the primary

25 light source used to activate porphyrins. Light Emitting Diodes (LEDs) and florescent light sources may also be used in some applications.

In some embodiments, the photodynamic agent is linked to GCC at the GCC N-terminus.

In addition to an intraoperative spray, conjugated compounds may be delivered intrathecally, intraventrically, stereotactically, intrahepatically such as via the portal vein, by inhalation, and intrapleurally.

### **Drug Delivery Targeted To Stomach or Esophageal Cancer Cells Generally**

- 5           Another aspect of the invention relates to unconjugated and conjugated compositions which comprise a GCC ligand used to deliver therapeutic agents to cells that comprise a GCC such as primary and/or metastatic stomach or esophageal cancer cells. In some embodiments, the agent is a drug or toxin such as: methotrexate, doxorubicin, daunorubicin, cytosinarabioside, etoposide, 5-4 fluorouracil, melphalan, chlorambucil,
- 10   *cis*-platinum, vindesine, mitomycin, bleomycin, purothionin, macromomycin, 1,4-benzoquinone derivatives, trenimon, ricin, ricin A chain, *Pseudomonas* exotoxin, diphtheria toxin, *Clostridium perfringens* phospholipase C, bovine pancreatic ribonuclease, pokeweed antiviral protein, abrin, abrin A chain, cobra venom factor, gelonin, saporin, modeccin, viscumin, volkensin, alkaline phosphatase, nitroimidazole, metronidazole and
- 15   misonidazole.. Genetic material is delivered to cancer cells to produce an antigen that can be targeted by the immune system or to produce a protein which kills the cell or inhibits its proliferation. In some embodiments, the GCC ligand is used to deliver nucleic acids that encode nucleic acid molecules which replace defective endogenous genes or which encode therapeutic proteins. In some embodiments, the compositions are used in gene therapy
- 20   protocols to deliver to individuals, genetic material needed and/or desired to make up for a genetic deficiency.

- In some embodiments, the GCC ligand is combined with or incorporated into a delivery vehicle thereby converting the delivery vehicle into a specifically targeted delivery vehicle. For example, a GCC binding peptide may be integrated into the outer
- 25   portion of a viral particle making such a virus a GCC-bearing cell specific virus. Similarly, the coat protein of a virus may be engineered such that it is produced as a fusion protein which includes an active GCC binding peptide that is exposed or otherwise accessible on the outside of the viral particle making such a virus a GCC-bearing cell-specific virus. In some embodiments, a GCC ligand may be integrated or otherwise incorporated into the

liposomes wherein the GCC ligand is exposed or otherwise accessible on the outside of the liposome making such liposomes specifically targeted to GCC-bearing cells.

The active agent in the conjugated or unconjugated compositions according to this aspect of the invention is a drug, toxin or nucleic acid molecule. The nucleic acid  
5 may be RNA or preferably DNA. In some embodiments, the nucleic acid molecule is an antisense molecule or encodes an antisense sequence whose presence in the cell inhibits production of an undesirable protein. In some embodiments, the nucleic acid molecule encodes a ribozyme whose presence in the cell inhibits production of an undesirable protein. In some embodiments, the nucleic acid molecule encodes a protein or peptide that  
10 is desirably produced in the cell. In some embodiments, the nucleic acid molecule encodes a functional copy of a gene that is defective in the targeted cell. The nucleic acid molecule is preferably operably linked to regulatory elements needed to express the coding sequence in the cell.

Liposomes are small vesicles composed of lipids. Genetic constructs which  
15 encode proteins that are desired to be expressed in GCC-bearing cells are introduced into the center of these vesicles. The outer shell of these vesicles comprise an a GCC ligand. *Liposomes* Volumes 1, 2 and 3 CRC Press Inc. Boca Raton FLA, which is incorporated herein by reference, disclose preparation of liposome-encapsulated active agents which include antibodies in the outer shell. In the present invention, a GCC ligand such as for  
20 example an anti-GCC antibodies is associated with the in the outer shell. Unconjugated compositions which comprise a GCC ligand in the matrix of a liposome with an active agent inside include such compositions in which the GCC ligand is preferably an antibody.

In one embodiment, the delivery of normal copies of the p53 tumor suppressor gene to the cancer cells is accomplished using GCC ligand to target the gene  
25 therapeutic. Mutations of the p53 tumor suppressor gene appears to play a prominent role in the development of many cancers. One approach to combating this disease is the delivery of normal copies of this gene to the cancer cells expressing mutant forms of this gene. Genetic constructs that comprise normal p53 tumor suppressor genes are incorporated into liposomes that comprise a GCC ligand. The composition is delivered to  
30 the tumor. GCC ligands specifically target and direct the liposomes containing the normal gene to correct the lesion created by mutation of p53 suppressor gene. Preparation of

genetic constructs is with the skill of those having ordinary skill in the art. The present invention allows such construct to be specifically targeted by using the GCC ligands of the present invention. The compositions of the invention include a GCC ligand such as an anti-GCC antibody associated with a delivery vehicle and a gene construct which

5 comprises a coding sequence for a protein whose production is desired in the cells of the intestinal tract linked to necessary regulatory sequences for expression in the cells. For uptake by cells of the intestinal tract, the compositions are administered orally or by enema whereby they enter the intestinal tract and contact cells which comprise GCC. The delivery vehicles associate with the GCC by virtue of the GCC ligand and the vehicle is

10 internalized into the cell or the active agent/genetic construct is otherwise taken up by the cell. Once internalized, the construct can provide a therapeutic effect on the individual.

### **Antisense**

The present invention provides compositions, kits and methods which are useful to prevent and treat stomach or esophageal cancer cells by providing the means to

15 specifically deliver antisense compounds to stomach or esophageal cancer cells and thereby stop expression of genes in such cells in which undesirable gene expression is taking place without negatively effecting cells in which no such expression occurs.

The conjugated compositions of the present invention are useful for targeting cells that express GCC including stomach or esophageal cancer cells. The conjugated

20 compositions will not bind to non-colorectal derived cells. Non-colorectal cells, lacking GCC, do not take up the conjugated compositions. Normal colorectal cells do have GCC and will take up the compositions. The present invention provides compositions and methods of delivering antisense compositions to stomach or esophageal cancer cells.

The present invention provides a cell specific approach in which only normal

25 and cancerous colorectal cells and primary and/or metastatic stomach or esophageal cancer cells are exposed to the active portion of the compound and only those cells are effected by the conjugated compound. The GCC binding moiety binds to normal and cancerous colorectal cells and primary and/or metastatic stomach or esophageal cancer cells. Upon binding to these cells, the conjugated compound is internalized and the delivery of the

30 conjugated compound including the antisense portion of the molecule is effected. The



presence of the conjugated compound in normal colorectal cells has no effect on such cells because the cancer-associated gene for which the antisense molecule that makes up the active moiety of the conjugated compound is complementary is not being expressed. However, in colorectal cancer cells, the cancer gene for which the antisense molecule that  
5 makes up the active moiety of the conjugated compound is complementary is being expressed. The presence of the conjugated compound in colorectal cancer cells serves to inhibit or prevent transcription or translation of the cancer gene and thereby reduce or eliminate the transformed phenotype.

The invention can be used to combat primary and/or metastasized colorectal,  
10 stomach or esophageal cancer as well as to prevent the emergence of the transformed phenotype in normal colon cells. Thus the invention can be used therapeutically as well as prophylactically.

One having ordinary skill in the art can readily identify individuals suspected of suffering from stomach or esophageal cancer. In those individuals diagnosed with  
15 stomach or esophageal cancer, it is standard therapy to suspect metastasis and aggressively attempt to eradicate metastasized cells. The present invention provides pharmaceutical compositions and methods for specifically targeting and eliminating primary and/or metastatic stomach or esophageal cancer cells. Further, the present invention provides pharmaceutical compositions that comprise therapeutics and methods for specifically  
20 eliminating primary and/or metastatic stomach or esophageal cancer cells.

The present invention relies upon the use of a GCC binding moiety in a conjugated composition. The GCC product binding moiety is essentially a portion of the conjugated composition which acts as a ligand to the GCC and thus specifically binds to these receptors. The conjugated composition also includes an active moiety which is  
25 associated with the GCC binding moiety; the active moiety being an antisense composition useful to inhibit or prevent transcription or translation of expression of genes whose expression is associated with cancer.

According to the present invention, the active moiety is an antisense composition. In particular, the antisense molecule that makes up the active moiety of a  
30 conjugated compound hybridizes to DNA or RNA in a stomach or esophageal cancer cell and inhibits and/or prevents transcription or translation of the DNA or RNA from taking

place. The antisense compositions may be a nucleic acid molecule, a derivative or an analogs thereof. The chemical nature of the antisense composition may be that of a nucleic acid molecule or a modified nucleic acid molecule or a non-nucleic acid molecule which possess functional groups that mimic a DNA or RNA molecule that is complementary to  
5 the DNA or RNA molecule whose expression is to be inhibited or otherwise prevented. Antisense compositions inhibit or prevent transcription or translation of genes whose expression is linked to stomach or esophageal cancer, i.e. cancer associated genes.

Point mutations insertions, and deletions in K-ras and H-ras have been identified in many tumors. Complex characteristics of the alterations of oncogenes HER-  
10 2/ERBB-2, HER-1/ERBB-1, HRAS-1, C-MYC and anti-oncogenes p53, RB1.

Chemical carcinogenesis in a rat model demonstrated point mutations in fos, an oncogene which mediates transcriptional regulation and proliferation. See: Alexander, RJ, *et al.* Oncogene alterations in rat colon tumors induced by N-methyl-N-nitrosourea. *American Journal of the Medical Sciences*. **303**(1):16-24, 1992, Jan. which is hereby  
15 incorporated herein by reference including all references cited therein which are also hereby incorporated herein by reference.

Chemical carcinogenesis in a rat model demonstrated point mutations in the oncogene abl. See: Alexander, RJ, *et al.* Oncogene alterations in rat colon tumors induced by N-methyl-N-nitrosourea. *American Journal of the Medical Sciences*. **303**(1):16-24,  
20 1992, Jan.

MYC is an oncogene that plays a role in regulating transcription and proliferation. A 15-base antisense oligonucleotide to myc complementary to the translation initiation region of exon II was incubated with colorectal cancer cells. This antisense molecule inhibited proliferation of colorectal cancer cells in a dos-dependent  
25 fashion. Interestingly, the uptake of this oligonucleotide was low (0.7%). Also, transfer of a normal chromosome 5 to colorectal cancer cells resulted in the regulation of myc expression and loss of proliferation. These data suggest that a tumor suppressor gene important in the regulation of myc is contained on this chromosome.

A novel protein tyrosine phosphatase, G1, has been identified. Examination  
30 of the mRNA encoding this protein in colorectal tumor cells revealed that it undergoes point mutations and deletions in these cells and may play a role in proliferation

characteristic of these cells. Takekawa, M. *et al.* Chromosomal localization of the protein tyrosine phosphatase G1 gene and characterization of the aberrant transcripts in human colon cancer cells. *FEBS Letters*. 339(3):222-8, 1994 Feb. 21; which is hereby incorporated herein by reference including all references cited therein which are also

5 hereby incorporated herein by reference.

Gastrin regulates colon cancer cell growth through a cyclic AMP-dependent mechanism mediated by PKA. Antisense oligodeoxynucleotides to the regulatory subunit of a specific class of PKA inhibited the growth-promoting effects of cyclic AMP in colon carcinoma cells. See: Bold, RJ, *et al.* Experimental gene therapy of human colon cancer.

10 *Surgery*. 116(2):189-95; discussion 195-6, 1994 Aug. and Yokozaki, H., *et al.* An antisense oligodeoxynucleotide that depletes RI alpha subunit of cyclic AMP-dependent protein kinase induces growth inhibition in human cancer cells. *Cancer Research*. 53(4):868-72, 1993 Feb 15, which are both hereby incorporated herein by reference including all references cited therein which are also hereby incorporated herein by

15 reference.

CRIPTO is an epidermal growth factor-related gene expressed in a majority of colorectal cancer tumors. Antisense phosphorothioate oligodeoxynucleotides to the 5'-end of CRIPTO mRNA significantly reduced CRIPTO expression and inhibited colorectal tumor cell growth *in vitro* and *in vivo*. Ciardiello, F. *et al.* Inhibition of CRIPTO

20 expression and tumorigenicity in human colon cancer cells by antisense RNA and oligodeoxynucleotides. *Oncogene*. 9(1):291-8, 1994 Jan. which are both hereby incorporated herein by reference including all references cited therein which are also hereby incorporated herein by reference.

Many carcinoma cells secrete transforming growth factor alpha. A 23

25 nucleotide antisense oligonucleotide to TGF alpha mRNA inhibited both DNA synthesis and proliferation of colorectal cancer cells. Sizeland, AM, Burgess, AW. Antisense transforming growth factor alpha oligonucleotides inhibit autocrine stimulated proliferation of a colon carcinoma cell line. *Molecular Biology of the Cell*. 3(11):1235-43, 1992 Nov. which is hereby incorporated herein by reference including all references cited therein

30 which are also hereby incorporated herein by reference.

Antisense compositions including oligonucleotides, derivatives and analogs thereof, conjugation protocols, and antisense strategies for inhibition of transcription and translation are generally described in: *Antisense Research and Applications*, Crooke, S. and B. Lebleu, eds. CRC Press, Inc. Boca Raton FLA 1993; *Nucleic Acids in Chemistry and*  
5 *Biology* Blackburn, G. and M.J. Gait, eds. IRL Press at Oxford University Press, Inc. New York 1990; and *Oligonucleotides and Analogues: A Practical Approach* Eckstein, F. ed., IRL Press at Oxford University Press, Inc. New York 1991; which are each hereby incorporated herein by reference including all references cited therein which are hereby incorporated herein by reference.

10           The antisense molecules of the present invention comprise a sequence complementary to a fragment of a colorectal cancer gene. See Ullrich et al., *EMBO J.*, 1986, 5:2503, which is hereby incorporated herein by reference.

Antisense compositions which can make up an active moiety in conjugated compounds of the invention include oligonucleotides formed of homopyrimidines can  
15 recognize local stretches of homopurines in the DNA double helix and bind to them in the major groove to form a triple helix. See: Helen, C and Toulme, JJ. Specific regulation of gene expression by antisense, sense, and antigene nucleic acids. *Biochem. Biophys Acta*, 1049:99-125, 1990 which is hereby incorporated herein by reference including all references cited therein which are hereby incorporated herein by reference. Formation of  
20 the triple helix would interrupt the ability of the specific gene to undergo transcription by RNA polymerase. Triple helix formation using myc-specific oligonucleotides has been observed. See: Cooney, M, *et al. Science* 241:456-459 which is hereby incorporated herein by reference including all references cited therein which are hereby incorporated herein by reference.

25           Antisense oligonucleotides of DNA or RNA complementary to sequences at the boundary between introns and exons can be employed to prevent the maturation of newly-generated nuclear RNA transcripts of specific genes into mRNA for transcription.

Antisense RNA complimentary to specific genes can hybridize with the mRNA for tat gene and prevent its translation. Antisense RNA can be provided to the cell  
30 as "ready-to-use" RNA synthesized in vitro or as an antisense gene stably transfected into cells which will yield antisense RNA upon transcription. Hybridization with mRNA results

in degradation of the hybridized molecule by RNase H and/or inhibition of the formation of translation complexes. Both result in a failure to produce the product of the original gene.

Antisense sequences of DNA or RNA can be delivered to cells. Several  
5 chemical modifications have been developed to prolong the stability and improve the function of these molecules without interfering in their ability to recognize specific sequences. These include increasing their resistance to degradation by DNases, including phosphotriesters, methylphosphonates, phosphorothioates, alpha-anomers, increasing their affinity for their target by covalent linkage to various intercalating agents such as  
10 psoralens, and increasing uptake by cells by conjugation to various groups including polylysine. These molecules recognize specific sequences encoded in mRNA and their hybridization prevents translation of and increases the degradation of these messages.

Conjugated compositions of the invention provide a specific and effective means for terminating the expression of genes which cause neoplastic transformation.  
15 GCC undergo ligand-induced endocytosis and can deliver conjugated compounds to the cytoplasm of cells.

GCC - binding moieties are conjugated directly to antisense compositions such as nucleic acids which are active in inducing a response. For example, antisense oligonucleotides to MYC are conjugated directly to an anti-GCC antibody. This has been  
20 performed employing peptides that bind to the CD4 receptor. See: Cohen, JS, ed. *Oligodeoxynucleotides: Antisense Inhibitors of Gene Expression. Topics in Molecular and Structural Biology*. CRC Press, Inc., Boca Raton, 1989. which is hereby incorporated herein by reference including all references cited therein which are hereby incorporated herein by reference. The precise backbone and its synthesis is not specified and can be  
25 selected from well-established techniques. Synthesis would involve either chemical conjugation or direct synthesis of the chimeric molecule by solid phase synthesis employing Fmoc chemistry. See: Haralambidis, J, *et al.* (1987) *Tetrahedron Lett.* 28:5199-5202, which is hereby incorporated herein by reference including all references cited therein which are hereby incorporated herein by reference. Alternatively, the peptide-  
30 nucleic acid conjugate may be synthesized directly by solid phase synthesis as a peptide-peptide nucleic acid chimera by solid phase synthesis. Nielsen, PE, *et al.* (1994) Sequence-

specific transcription arrest by peptide nucleic acid bound to the DNA template strand. *Gene* 149:139-145, which is hereby incorporated herein by reference including all references cited therein which are hereby incorporated herein by reference.

In some embodiments, polylysine can be complexed to conjugated  
 5 compositions of the invention in a non-covalent fashion to nucleic acids and used to enhance delivery of these molecules to the cytoplasm of cells. In addition, peptides and proteins can be conjugated to polylysine in a covalent fashion and this conjugate complexed with nucleic acids in a non-covalent fashion to further enhance the specificity and efficiency of uptake of the nucleic acids into cells. Thus, GCC ligand is conjugated  
 10 chemically to polylysine by established techniques. The polylysine-GCC-1 translation product ligand conjugate may be complexed with nucleic acids of choice. Thus, polylysine-orosomucoid conjugates were employed to specifically plasmids containing genes to be expressed to hepatoma cells expressing the orosomucoid receptor. This approach can be used to delivery whole genes, or oligonucleotides. Thus, it has the  
 15 potential to terminate the expression of an undesired gene (eg. MYC, ras) or replace the function of a lost or deleted gene (eg. hMSH2, hMLH1, hPMS1, and hPMS2).

According to a preferred embodiment, Myc serves as a gene whose expression is inhibited by an antisense molecule within a conjugated composition. GCC binding moieties are used to deliver a 15-based antisense oligonucleotide to myc  
 20 complementary to the translation initiation region of exon II. The 15-base antisense oligonucleotide to MYC is synthesized as reported in Collins, JF, Herman, P, Schuch, C, Bagby GC, Jr. *Journal of Clinical Investigation*. 89(5):1523-7, 1992 May. In some embodiments, the conjugated composition is conjugated to polylysine as reported previously. Wu, GY, and Wu, CH. (1988) Evidence for ed gene delivery to Hep G2  
 25 hepatoma cells in vitro. *Biochem*. 27:887-892 which is incorporated herein by reference.

Conjugated compositions may be synthesized as a chimeric molecule directly by solid phase synthesis. pmolar to nanomolar concentrations for this conjugate suppress MYC synthesis in colorectal cancer cells *in vitro*.

Antisense molecules are preferably hybridize to, i.e. are complementary to, a  
 30 nucleotide sequence that is 5-50 nucleotides in length, more preferably 5-25 nucleotides and in some embodiments 10-15 nucleotides.

In addition, mismatches within the sequences identified above, which achieve the methods of the invention, such that the mismatched sequences are substantially complementary to the cancer gene sequences are also considered within the scope of the disclosure. Mismatches which permit substantial complementarity to the cancer gene sequences will be known to those of skill in the art once armed with the present disclosure. The oligos may also be unmodified or modified.

Therapeutic compositions and methods may be used to combat stomach or esophageal cancer in cases where the cancer is localized and/or metastasized. Individuals are administered a therapeutically effective amount of conjugated compound. A therapeutically effective amount is an amount which is effective to cause a cytotoxic or cytostatic effect on cancer cells without causing lethal side effects on the individual. An individual who has been administered a therapeutically effective amount of a conjugated composition has a increased chance of eliminating stomach or esophageal cancer as compared to the risk had the individual not received the therapeutically effective amount.

To treat localized stomach or esophageal cancer, a therapeutically effective amount of a conjugated compound is administered such that it will come into contact with the localized tumor. Thus, the conjugated compound may be administered orally or intratumorally. Oral and rectal formulation are taught in Remington's Pharmaceutical Sciences, 18th Edition, 1990, Mack Publishing Co., Easton PA. which is incorporated herein by reference.

The pharmaceutical compositions according to the present invention may be administered as either a single dose or in multiple doses. The pharmaceutical compositions of the present invention may be administered either as individual therapeutic agents or in combination with other therapeutic agents. The treatments of the present invention may be combined with conventional therapies, which may be administered sequentially or simultaneously.

The present invention is directed to a method of delivering antisense compounds to normal and cancerous colorectal cells and to stomach or esophageal cancer cells and inhibiting expression of cancer genes in mammals. The methods comprise administering to a mammal an effective amount of a conjugated composition which

comprises a GCC binding moiety conjugated to an antisense oligonucleotide having a sequence which is complementary to a region of DNA or mRNA of a cancer gene.

The conjugated compounds may be administering to mammals in a mixture with a pharmaceutically-acceptable carrier, selected with regard to the intended route of administration and the standard pharmaceutical practice. Dosages will be set with regard to weight, and clinical condition of the patient. The conjugated compositions of the present invention will be administered for a time sufficient for the mammals to be free of undifferentiated cells and/or cells having an abnormal phenotype. In therapeutic methods treatment extends for a time sufficient to inhibit transformed cells from proliferating and conjugated compositions may be administered in conjunction with other chemotherapeutic agents to manage and combat the patient's cancer.

The conjugated compounds of the invention may be employed in the method of the invention singly or in combination with other compounds. The amount to be administered will also depend on such factors as the age, weight, and clinical condition of the patient. See Gennaro, Alfonso, ed., Remington's Pharmaceutical Sciences, 18th Edition, 1990, Mack Publishing Co., Easton PA.

### **Therapeutic and Prophylactic Vaccines**

The invention relates to prophylactic and therapeutic vaccines for protecting individuals against primary and/or metastatic stomach or esophageal cancer cells and for treating individuals who are suffering from primary and/or metastatic stomach or esophageal cancer cells.

According to the present invention, GCC serves as targets against which a protective and therapeutic immune response can be induced. Specifically, vaccines are provided which induce an immune response against GCC. The vaccines of the invention include, but are not limited to, the following vaccine technologies:

- 1) DNA vaccines, i.e. vaccines in which DNA that encodes at least an epitope from an GCC is administered to an individual's cells where the epitope is expressed and serves as a target for an immune response;
- 2) infectious vector mediated vaccines such as recombinant adenovirus, vaccinia, *Salmonella*, and BCG wherein the vector carries genetic information that encodes



at least an epitope from an GCC protein such that when the infectious vector is administered to an individual, the epitope is expressed and serves as a target for an immune response;

3) killed or inactivated vaccines which a) comprise either killed cells or  
5 inactivated viral particles that display at least an epitope from an GCC protein and b) when administered to an individual serves as a target for an immune response;

4) haptenized killed or inactivated vaccines which a) comprise either killed cells or inactivated viral particles that display at least an epitope from an GCC protein, b) are haptenized to be more immunogenic and c) when administered to an individual serves  
10 as a target for an immune response;

5) subunit vaccines which are vaccines that include protein molecules that include at least an epitope from an GCC protein; and

6) haptenized subunit vaccines which are vaccines that a) include protein molecules that include at least an epitope from an GCC protein and b) are haptenized to be  
15 more immunogenic.

The present invention relates to administering to an individual a protein or nucleic acid molecule that comprises or encodes, respectively, an immunogenic epitope against which an therapeutic and prophylactic immune response can be induced. Such epitopes are generally at least 6-8 amino acids in length. The vaccines of the invention  
20 therefore comprise proteins which are at least, or nucleic acids which encode at least, 6-8 amino acids in length from GCC protein. The vaccines of the invention may comprise proteins which are at least, or nucleic acids which encode at least 10 to about 1000 amino acids in length. The vaccines of the invention may comprise proteins which are at least, or nucleic acids which encode at least, about 25 to about 500 amino acids in length. The  
25 vaccines of the invention may comprise proteins which are at least, or nucleic acids which encode at least, about 50 to about 400 amino acids in length. The vaccines of the invention may comprise proteins which are at least, or nucleic acids which encode at least, about 100 to about 300 amino acids in length.

The present invention relates to compositions for and methods of treating  
30 individuals who are known to have primary and/or metastatic stomach or esophageal cancer cells. Primary and/or metastatic stomach or esophageal cancer may be diagnosed

by those having ordinary skill in the art using the methods described herein or art accepted clinical and laboratory pathology protocols. The present invention provides an immunotherapeutic vaccine useful to treat individuals who have been diagnosed as suffering from primary and/or metastatic stomach or esophageal cancer. The

5 immunotherapeutic vaccines of the present invention may be administered in combination with other therapies.

The present invention relates to compositions for and methods of preventing primary and/or metastatic stomach or esophageal cancer in individual is suspected of being susceptible to stomach or esophageal cancer. Such individuals include those whose family

10 medical history indicates above average incidence of stomach or esophageal cancer among family members and/or those who have already developed stomach or esophageal cancer and have been effectively treated who therefore face a risk of relapse and recurrence. Such individuals include those which have been diagnosed as having stomach or esophageal cancer including localized only or localized and metastasized stomach or esophageal

15 cancer which has been resected or otherwise treated. The vaccines of the present invention may be to susceptible individuals prophylactically to prevent and combat primary and metastatic stomach or esophageal cancer.

The invention relates to compositions which are the active components of such vaccines or required to make the active components, to methods of making such

20 compositions including the active components, and to methods of making and using vaccines.

The amino acid and nucleotide sequences of the GCC is set forth as SEQ ID NO:1.

The present invention relates to recombinant vectors, including expression

25 vectors, that comprise the GCC gene transcript or a fragment thereof. The present invention relates to recombinant vectors, including expression vectors that comprise nucleotide sequences that encode a GCC protein or a functional fragment thereof.

The present invention relates to host cells which comprise such vectors and to methods of making GCC protein using such recombinant cells.

The present invention relates to the isolated GCC gene transcript and to the isolated GCC proteins and to isolated antibodies specific for such protein and to hybridomas which produce such antibodies.

The present invention relates to the isolated GCC and functional fragments thereof. Accordingly, some aspects of the invention relate to isolated proteins that  
5 comprise at least one epitope of an GCC.

Some aspects of the invention relate to the above described isolated proteins which are haptenized to render them more immunogenic. That is, some aspects of the invention relate to haptenized proteins that comprise at least one GCC epitope.

10 Accordingly, some aspects of the invention relate to isolated nucleic acid molecules that encode proteins that comprise at least one GCC epitope.

Naked DNA vaccines are described in PCT/US90/01515, which is incorporated herein by reference. Others teach the use of liposome mediated DNA transfer, DNA delivery using microprojectiles (U.S. Patent No. 4,945,050 issued July 31,  
15 1990 to Sanford et al., which is incorporated herein by reference), and DNA delivery using electroporation. In each case, the DNA may be plasmid DNA that is produced in bacteria, isolated and administered to the animal to be treated. The plasmid DNA molecules are taken up by the cells of the animal where the sequences that encode the protein of interest are expressed. The protein thus produced provides a therapeutic or prophylactic effect on  
20 the animal.

The use of vectors including viral vectors and other means of delivering nucleic acid molecules to cells of an individual in order to produce a therapeutic and/or prophylactic immunological effect on the individual are similarly well known.

Recombinant vaccines that employ vaccinia vectors are, for example, disclosed in U.S.  
25 Patent Number 5,017,487 issued May 21, 1991 to Stunnenberg *et al.* which is incorporated herein by reference.

In some cases, tumor cells from the patient are killed or inactivated and administered as a vaccine product. Berd *et al.* May 1986 *Cancer Research* 46:2572-2577 and Berd *et al.* May 1991 *Cancer Research* 51:2731-2734, which are incorporated herein  
30 by reference, describes the preparation and use of tumor cell based vaccine products. According to some aspects of the present invention, the methods and techniques described

in Berd *et al.* are adapted by using stomach or esophageal cancer cells instead of melanoma cells.

The manufacture and use of isolated translation products and fragments thereof useful for example as laboratory reagents or components of subunit vaccines are well known. One having ordinary skill in the art can isolate the GCC gene transcript or the specific portion thereof that encodes GCC or a fragment thereof. Once isolated, the nucleic acid molecule can be inserted it into an expression vector using standard techniques and readily available starting materials.

The recombinant expression vector that comprises a nucleotide sequence that encodes the nucleic acid molecule that encodes GCC or a fragment thereof or a protein that comprises the GCC or a fragment thereof. The recombinant expression vectors of the invention are useful for transforming hosts to prepare recombinant expression systems for preparing the isolated proteins of the invention.

The present invention relates to a host cell that comprises the recombinant expression vector that includes a nucleotide sequence that encodes GCC protein or a fragment thereof or an GCC or a fragment thereof. Host cells for use in well known recombinant expression systems for production of proteins are well known and readily available. Examples of host cells include bacteria cells such as *E. coli*, yeast cells such as *S. cerevisiae*, insect cells such as *S. frugiperda*, non-human mammalian tissue culture cells chinese hamster ovary (CHO) cells and human tissue culture cells such as HeLa cells.

The present invention relates to a transgenic non-human mammal that comprises the recombinant expression vector that comprises a nucleic acid sequence that encodes the proteins of the invention. Transgenic non-human mammals useful to produce recombinant proteins are well known as are the expression vectors necessary and the techniques for generating transgenic animals. Generally, the transgenic animal comprises a recombinant expression vector in which the nucleotide sequence that encodes GCC or a fragment thereof or a protein that comprises GCC or a fragment thereof operably linked to a mammary cell specific promoter whereby the coding sequence is only expressed in mammary cells and the recombinant protein so expressed is recovered from the animal's milk.

In some embodiments, for example, one having ordinary skill in the art can, using well known techniques, insert such DNA molecules into a commercially available expression vector for use in well known expression systems such as those described herein.

The expression vector including the DNA that encodes a GCC or a functional  
5 fragment thereof or a protein that comprises a GCC or a functional fragment thereof is used to transform the compatible host which is then cultured and maintained under conditions wherein expression of the foreign DNA takes place. The protein of the present invention thus produced is recovered from the culture, either by lysing the cells or from the culture medium as appropriate and known to those in the art. The methods of purifying the GCC  
10 or a fragment thereof or a protein that comprises the same using antibodies which specifically bind to the protein are well known. Antibodies which specifically bind to a particular protein may be used to purify the protein from natural sources using well known techniques and readily available starting materials. Such antibodies may also be used to purify the protein from material present when producing the protein by recombinant DNA  
15 methodology. The present invention relates to antibodies that bind to an epitope which is present on one or more GCC-1 translation products or a fragment thereof or a protein that comprises the same. Antibodies that bind to an epitope which is present on the GCC are useful to isolate and purify the protein from both natural sources or recombinant expression systems using well known techniques such as affinity chromatography. Immunoaffinity  
20 techniques generally are described in Waldman *et al.* 1991 *Methods of Enzymol.* **195**:391-396, which is incorporated herein by reference. Antibodies are useful to detect the presence of such protein in a sample and to determine if cells are expressing the protein. The production of antibodies and the protein structures of complete, intact antibodies, Fab fragments and F(ab)<sub>2</sub> fragments and the organization of the genetic sequences that encode  
25 such molecules are well known and are described, for example, in Harlow, E. and D. Lane (1988) *ANTIBODIES: A Laboratory Manual*, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY. which is incorporated herein by reference.

In some embodiments of the invention, transgenic non-human animals are generated. The transgenic animals according to the invention contain nucleotides that  
30 encode GCC or a fragment thereof or a protein that comprises the same under the regulatory control of a mammary specific promoter. One having ordinary skill in the art

using standard techniques, such as those taught in U.S. Patent No. 4,873,191 issued October 10, 1989 to Wagner and U.S. Patent No. 4,736,866 issued April 12, 1988 to Leder, both of which are incorporated herein by reference, can produce transgenic animals which produce GCC or a fragment thereof or a protein that comprises the same. Preferred

5 animals are goats and rodents, particularly rats and mice.

In addition to producing these proteins by recombinant techniques, automated peptide synthesizers may also be employed to produce GCC or a fragment thereof or a fragment thereof or a protein that comprises the same. Such techniques are well known to those having ordinary skill in the art and are useful if derivatives which

10 have substitutions not provided for in DNA-encoded protein production.

In some embodiments, the protein that makes up a subunit vaccine or the cells or particles of a killed or inactivated vaccine may be haptenized to increase immunogenicity. In some cases, the haptenization is the conjugation of a larger molecular structure to GCC or a fragment thereof or a protein that comprises the same. In some

15 cases, tumor cells from the patient are killed and haptenized as a means to make an effective vaccine product. In cases in which other cells, such as bacteria or eukaryotic cells which are provided with the genetic information to make and display a GCC or a fragment thereof or a protein that comprises the same, are killed and used as the active vaccine component, such cells are haptenized to increase immunogenicity. Haptenization is well

20 known and can be readily performed.

Methods of haptenizing cells generally and tumor cells in particular are described in Berd *et al.* May 1986 *Cancer Research* 46:2572-2577 and Berd *et al.* May 1991 *Cancer Research* 51:2731-2734, which are incorporated herein by reference. Additional haptenization protocols are disclosed in Miller *et al.* 1976 *J. Immunol.*

25 117(5:1):1591-1526.

Haptenization compositions and methods which may be adapted to be used to prepare haptenized GCC immunogens according to the present invention include those described in the following U.S. Patents which are each incorporated herein by reference: U.S. Patent Number 5,037,645 issued August 6, 1991 to Strahilevitz; U.S. Patent Number

30 5,112,606 issued May 12, 1992 to Shiosaka *et al.*; U.S. Patent Number 4,526,716 issued July 2, 1985 to Stevens; U.S. Patent Number 4,329,281 issued May 11, 1982 to

Christenson *et al.*; and U.S. Patent Number 4,022,878 issued May 10, 1977 to Gross. Peptide vaccines and methods of enhancing immunogenicity of peptides which may be adapted to modify GCC immunogens of the invention are also described in Francis *et al.* 1989 *Methods of Enzymol.* **178**:659-676, which is incorporated herein by reference. Sad *et al.* 1992 *Immunology* **76**:599-603, which is incorporated herein by reference, teaches methods of making immunotherapeutic vaccines by conjugating gonadotropin releasing hormone to diphtheria toxoid. GCC immunogens may be similarly conjugated to produce an immunotherapeutic vaccine of the present invention. MacLean *et al.* 1993 *Cancer Immunol. Immunother.* **36**:215-222, which is incorporated herein by reference, describes conjugation methodologies for producing immunotherapeutic vaccines which may be adaptable to produce an immunotherapeutic vaccine of the present invention. The hapten is keyhole limpet hemocyanin which may be conjugated to a GCC immunogen.

Vaccines according to some aspects of the invention comprise a pharmaceutically acceptable carrier in combination with a GCC immunogen.

Pharmaceutical formulations are well known and pharmaceutical compositions comprising such proteins may be routinely formulated by one having ordinary skill in the art. Suitable pharmaceutical carriers are described in *Remington's Pharmaceutical Sciences*, A. Osol, a standard reference text in this field, which is incorporated herein by reference. The present invention relates to an injectable pharmaceutical composition that comprises a pharmaceutically acceptable carrier and a GCC immunogen. The GCC immunogen is preferably sterile and combined with a sterile pharmaceutical carrier.

In some embodiments, for example, GCC or a fragment thereof or a fragment thereof or a protein that comprises the same can be formulated as a solution, suspension, emulsion or lyophilized powder in association with a pharmaceutically acceptable vehicle.

Examples of such vehicles are water, saline, Ringer's solution, dextrose solution, and 5% human serum albumin. Liposomes and nonaqueous vehicles such as fixed oils may also be used. The vehicle or lyophilized powder may contain additives that maintain isotonicity (e.g., sodium chloride, mannitol) and chemical stability (e.g., buffers and preservatives). The formulation is sterilized by commonly used techniques.

An injectable composition may comprise the GCC immunogen in a diluting agent such as, for example, sterile water, electrolytes/dextrose, fatty oils of vegetable

origin, fatty esters, or polyols, such as propylene glycol and polyethylene glycol. The injectable must be sterile and free of pyrogens.

The vaccines of the present invention may be administered by any means that enables the immunogenic agent to be presented to the body's immune system for  
5 recognition and induction of an immunogenic response. Pharmaceutical compositions may be administered parenterally, i.e., intravenous, subcutaneous, intramuscular.

Dosage varies depending upon known factors such as the pharmacodynamic characteristics of the particular agent, and its mode and route of administration; age, health, and weight of the recipient; nature and extent of symptoms, kind of concurrent treatment,  
10 frequency of treatment, and the effect desired. An amount of immunogen is delivered to induce a protective or therapeutically effective immune response. Those having ordinary skill in the art can readily determine the range and optimal dosage by routine methods.

The following examples are illustrative but are not meant to be limiting of the present invention.

## 15 EXAMPLES

As stated above, a GCC binding moiety is a GCC ligand that may be an antibody, a protein, a polypeptide, a peptide or a non-peptide. Peptides and non-peptide GCC ligands may be identified using well known technology.

Over the past 10 years, it has become recognized that the specific high-  
20 affinity interaction of a receptor and a ligand, for example a GCC and an anti-GCC antibody, has its basis in the 3-dimensional conformational space of the ligand and the complimentary 3-dimensional configuration of the region of the molecule involved in ligand binding. In addition, it has become recognized that various arrays of naturally-occurring amino acids, non-natural amino acids, and organic molecules can be organized in  
25 configurations that are unrelated to the natural ligands in their linear structure, but resemble the 3-dimensional structure of the natural ligands in conformational space and, thus, are recognized by receptors with high affinity and specificity. Furthermore, techniques have been described in the literature that permit one of ordinary skill in the art to generate large libraries of these arrays of natural amino acids, non-natural amino acids  
30 and organic compounds to prospectively identify individual compounds that interact with



receptors with high affinity and specificity which are unrelated to the native ligand of that receptor. Thus, it is a relatively straightforward task for one of ordinary skill in the art to identify arrays of naturally occurring amino acids, non-natural amino acids, or organic compounds which can bind specifically and tightly to the GCC, which bear no structural  
5 relationship to an anti-GCC antibody.

To identify GCC ligands that are peptides, those having ordinary skill in the art can use any of the well known methodologies for screening random peptide libraries in order to identify peptides which bind to the GCC. In the most basic of methodologies, the peptides which bind to the target are isolated and sequenced. In some methodologies, each  
10 random peptide is linked to a nucleic acid molecule which includes the coding sequence for that particular random peptide. The random peptides, each with an attached coding sequence, are contacted with a GCC and the peptides which are unbound to the GCC are removed. The nucleic acid molecule which includes the coding sequence of the peptide that binds to the GCC can then be used to determine the amino acid sequence of the  
15 peptide as well as produce large quantities of the peptide. It is also possible to produce peptide libraries on solid supports where the spatial location on the support corresponds to a specific synthesis and therefore specific peptide. Such methods often use photolithography-like steps to create diverse peptide libraries on solid supports in which the spatial address on the support allows for the determination of the sequence.

20 The production of organic compound libraries on solid supports may also be used to produce combinatorial libraries of non-peptide compounds such as oligonucleotides and sugars, for example. As in the case of peptide libraries on solid supports, the spatial location on the support corresponds to a specific synthesis and therefore specific compound. Such methods often use photolithography-like steps to create  
25 diverse compound libraries on solid supports in which the spatial address on the support allows for the determination of the synthesis scheme which produced the compound. Once the synthesis scheme is identified, the structure of the compound can become known.

Gallop *et al.* 1994 *J. Medicinal Chemistry* 37:1233, which is incorporated herein by reference, provides a review of several of the various methodologies of screening  
30 random peptide libraries and identifying peptides from such libraries which bind to target proteins. Following these teachings, GCC specific ligands that are peptides and that are

useful as GCC specific binding moieties may be identified by those having ordinary skill in the art.

Peptides and proteins displayed on phage particles are described in Gallop *et al. Supra*. Random arrays of nucleic acids can be inserted into genes encoding surface proteins of bacteriophage which are employed to infect bacteria, yielding phage expressing the peptides encoded by the random array of nucleotides on their surface. These phage displaying the peptide can be employed to determine whether those peptides can bind to specific proteins, receptors, antibodies, etc. The identity of the peptide can be determined by sequencing the recombinant DNA from the phage expressing the peptide. This approach has the potential to yield vast arrays of peptides in a library (up to  $10^9$  unique peptides). This technique has been employed to identify novel binding peptides to the fibrinogen receptor on platelets, which bear no sequence homology to the natural occurring ligands of this receptor (Smith *et al.*, 1993 *Gene* **128**:37, which is incorporated herein by reference). Similarly, this technique has been applied to identify peptides which bind to the MHC class II receptor (Hammer *et al.*, 1993 *Cell* **74**:197, which is incorporated herein by reference) and the chaperonin receptor (Blond-Elguindi *et al.*, 1993 *Cell* **75**:717, which is incorporated herein by reference).

Peptides displayed on plasmids are described in Gallop *et al. Supra*. In this approach, the random oligonucleotides which encode the library of peptides can be expressed on a specific plasmid whose expression is under the control of a specific promoter, such as the lac operon. The peptides are expressed as fusion proteins coupled to the Lac I protein, under the control of the lac operon. The fusion protein specifically binds to the lac operator on the plasmid and so the random peptide is associated with the specific DNA element that encodes it. In this way, the sequence of the peptide can be deduced, by PCR of the DNA associated with the fusion protein. These proteins can be screened in solution phase to determine whether they bind to specific receptors. Employing this approach, novel substrates have been identified for specific enzymes (Schatz 1993).

A variation of the above technique, also described in Gallop *et al. Supra*, can be employed in which random oligonucleotides encoding peptide libraries on plasmids can be expressed in cell-free systems. In this approach, a molecular DNA library can be constructed containing the random array of oligonucleotides, which are then expressed in a

bacterial in vitro transcription/translation system. The identity of the ligand is determined by purifying the complex of nascent chain peptide/polysome containing the mRNA of interest on affinity resins composed of the receptor and then sequencing following amplification with RT-PCR. Employing this technique permits generation of large  
5 libraries (up to  $10^{11}$  recombinants). Peptides which recognize antibodies specifically directed to dynorphin have been identified employing this technique (Cull *et al.*, 1992 *Proc. Natl. Acad. Sci. USA* **89**:1865, which is incorporated herein by reference).

Libraries of peptides can be generated for screening against a receptor by chemical synthesis. For example, simultaneous preparation of large numbers of diverse  
10 peptides have been generated employing the approach of multiple peptide synthesis as described in Gallop *et al. Supra*. In one application, random peptides are generated by standard solid-phase Merrifield synthesis on polyacrylamide microtiter plates (multipin synthesis) which are subsequently screened for their ability to compete with receptor binding in a standard competitive binding assay (Wang *et al.*, 1993 *Bioorg. Med. Chem. Lett.* **3**:447, which is incorporated herein by reference). Indeed, this approach has been  
15 employed to identify novel binding peptides to the substance P receptor (Wang *et al. Supra*). Similarly, peptide libraries can be constructed by multiple peptide synthesis employing the "tea bag" method in which bags of solid support resin are sequentially incubated with various amino acids to generate arrays of different peptides (Gallop *et al.*  
20 *Supra*). Employing this approach, peptides which bind to the integrin receptor (Ruggeri *et al.*, 1986 *Proc. Natl. Acad. Sci. USA* **83**:5708, which is incorporated herein by reference) and the neuropeptide Y receptor (Beck-Sickinger *et al.*, 1990 *Int. J. Peptide Protein Res.* **36**:522, which is incorporated herein by reference) have been identified.

In general, the generation and utility of combinatorial libraries depend on (1)  
25 a method to generate diverse arrays of building blocks, (2) a method for identifying members of the array that yield the desired function, and (3) a method for deconvoluting the structure of that member. Several approaches to these constraints have been defined.

The following is a description of methods of library generation which can be used in procedures for identifying GCC ligands according to the invention.

30 Modifications of the above approaches can be employed to generate libraries of vast molecular diversity by connecting together members of a set of chemical building

blocks, such as amino acids, in all possible combinations (Gallop *et al. Supra*) In one approach, mixtures of activated monomers are coupled to a growing chain of amino acids on a solid support at each cycle. This is a multivalent synthetic system.

Also, split synthesis involves incubating the growing chain in individual  
5 reactions containing only a single building block (Gallop *et al. Supra*). Following attachment, resin from all the reactions are mixed and apportioned into individual reactions for the next step of coupling. These approaches yield a stochastic collection of  $n^x$  different peptides for screening, where  $n$  is the number of building blocks and  $x$  is the number of cycles of reaction.

10 Alternatively, arrays of molecules can be generated in which one or more positions contain known amino acids, while the remainder are random (Gallop *et al. Supra*). These yield a limited library which is screened for members with the desired activity. These members are identified, their structure determined, and the structure regenerated with another position containing defined amino acids and screened. This  
15 iterative approach ultimately yields peptides which are optimal for recognizing the conformational binding pocket of a receptor.

In addition, arrays are not limited to amino acids forming peptides, but can be extended to linear and nonlinear arrays of organic molecules (Gordon *et al.*, 1994 *J. Medicinal Chemistry* 37:1385, which is incorporated herein by reference). Indeed,  
20 employing this approach of generating libraries of randomly arrayed inorganic building blocks, ligands which bound to 7-transmembrane receptors were identified (Zuckermann *et al.*, 1994 *J. Med. Chem.* 37:2678, which is incorporated herein by reference).

Libraries are currently being constructed which can be modified after synthesis to alter the chemical side groups and bonds, to give "designer" arrays to test for  
25 their interaction with receptors (Osteresh *et al.*, 1994 *Proc. Natl. Acad. Sci. USA* 91:11138, which is incorporated herein by reference). This technique, generating "libraries from libraries", was applied to the permethylation of a peptide library which yielded compounds with selective antimicrobial activity against gram positive bacteria.

Libraries are also being constructed to express arrays of pharmacological  
30 motifs, rather than specific structural arrays of amino acids (Sepetov *et al.*, 1995 *Proc. Natl. Acad. Sci. USA* 92:5426, which is incorporated herein by reference). This technique

seeks to identify structural motifs that have specific affinities for receptors, which can be modified in further refinements employing libraries to define structure-activity relationships. Employing this approach of searching motif libraries, generating "libraries of libraries", reduces the number of component members required for screening in the early  
5 phase of library examination.

The following is a description of methods of identifying GCC ligands according to the invention from libraries of randomly generated molecules.

Components in the library which interact with receptors may be identified by their binding to receptors immobilized on solid support (Gordon *et al. Supra*).

10 They may also be identified by their ability to compete with native ligand for binding to cognate receptors in solution phase (Gordon *et al. Supra*).

Components may be identified by their binding to soluble receptors when those components are immobilized on solid supports (Gordon *et al. Supra*).

Once a member of a library which binds receptors has been identified, the  
15 structure of that member must be deconvoluted (deduced) in order to identify the structure and generate large quantities to work with, or develop further analogs to study structure-activity relationships. The following is a description of methods of deconvolution for deducing the structure of molecules identified as potential GCC ligands according to the invention.

20 Peptide libraries may be expressed on the surface of bacteriophage particles (Gallop *et al. Supra*). Once the peptide interacting with the receptor has been identified, its structure can be deduced by isolating the DNA from the phage and determining its sequence by PCR.

Libraries expressed on plasmids, under the control of the Lac operon can be  
25 deconvoluted since these peptides are fused with the lac I protein which specifically interacts with the lac operon on the plasmid encoding the peptide (Gallop *et al. Supra*) The structure can be deduced by isolating that plasmid attached to the lac I protein and deducing the nucleotide and peptide sequence by PCR.

Libraries expressed on plasmids can also be expressed in cell-free systems  
30 employing transcription/translation systems (Gallop *et al. Supra*). In this paradigm, the

protein interacting with receptors is isolated with its attached ribosome and mRNA. The sequence of the peptide is deduced by RT-PCR of the associated mRNA.

Library construction can be coupled with photolithography, so that the structure of any member of the library can be deduced by determining its position within  
5 the substrate array (Gallop *et al. Supra*). This technique is termed positional addressability, since the structural information can be deduced by the precise position of the member.

Members of a library can also be identified by tagging the library with identifiable arrays of other molecules (Ohlmeyer *et al.*, 1993 *Proc. Natl. Acad. Sci. USA*  
10 90:10922, which is incorporated herein by reference, and Gallop *et al. Supra*). This technique is a modification of associating the peptide with the plasmid of phage encoding the sequence, described above. Some methods employ arrays of nucleotides to encode the sequential synthetic history of the peptide. Thus, nucleotides are attached to the growing peptide sequentially, and can be decoded by PCR to yield the structure of the associated  
15 peptide. Alternatively, arrays of small organic molecules can be employed as sequencable tags which encode the sequential synthetic history of the peptide. Thus, nucleotides are attached to the growing peptide sequentially, and can be decoded by PCR to yield the structure of the associated peptide. Alternatively, arrays of small organic molecules can be employed as sequencable tags which encode the sequential synthetic history of the library  
20 member.

Finally, the structure of a member of the library can be directly determined by amino acid sequence analysis.

The following patents, which are each incorporated herein by reference, describe methods of making random peptide or non-peptide libraries and screening such  
25 libraries to identify compounds that bind to target proteins. As used in the present invention, GCC can be the targets used to identify the peptide and non-peptide ligands generated and screened as disclosed in the patents.

U.S. Patent Number 5,270,170 issued to Schatz *et al.* on December 14, 1993, and U.S. Patent Number 5,338,665 issued to Schatz *et al.* on August 16, 1994, which are  
30 both incorporated herein by reference, refer to peptide libraries and screening methods which can be used to identify GCC ligands.

U.S. Patent No. 5,395,750 issued to Dillon et al. on March 7, 1995, which is incorporated herein by reference, refers to methods of producing proteins which bind to predetermined antigens. Such methods can be used to produce GCC ligands.

U.S. Patent No. 5,223,409 issued to Ladner et al. on June 29, 1993, which is  
5 incorporated herein by reference, refers to the directed evolution to novel binding proteins. Such proteins may be produced and screened as disclosed therein to identify GCC ligands.

U.S. Patent No. 5,366,862 issued to Venton et al. on November 22, 1994, which is incorporated herein by reference, refers to methods for generating and screening useful peptides. The methods herein described can be used to identify GCC ligands.

10 U.S. Patent No. 5,340,474 issued to Kauvar on August 23, 1994 as well as U.S. Patent No. 5,133,866, U.S. Patent No. 4,963,263 and U.S. Patent No. 5,217,869, which are each incorporated herein by reference, can be used to identify GCC ligands.

U.S. Patent No. 5,405,783 issued to Pirrung et al. on April 11, 1995, which is incorporated herein by reference, refers to large scale photolithographic solid phase  
15 synthesis of an array of polymers. The teachings therein can be used to identify GCC ligands.

U.S. Patent No. 5,143,854 issued to Pirrung et al. on September 1, 1992, which is incorporated herein by reference, refers to a large scale photolithographic solid phase synthesis of polypeptides and receptor binding screening thereof.

20 U.S. Patent No. 5,384,261 issued to Winkler et al. on January 24, 1995, which is incorporated herein by reference, refers to very large scale immobilized polymer synthesis using mechanically directed flow patterns. Such methods are useful to identify GCC ligands.

U.S. Patent No. 5,221,736 issued to Coolidge et al. on June 22, 1993, which  
25 is incorporated herein by reference, refers to sequential peptide and oligonucleotide synthesis using immunoaffinity techniques. Such techniques may be used to identify GCC ligands.

U.S. Patent No. 5,412,087 issued to McGall et al. on May 2, 1995, which is incorporated herein by reference, refers to spatially addressable immobilization of  
30 oligonucleotides and other biological polymers on surfaces. Such methods may be used to identify GCC ligands.

U.S. Patent No. 5,324,483 issued to Cody et al. on June 28, 1994, which is incorporated herein by reference, refers to apparatus for multiple simultaneous synthesis. The apparatus and method disclosed therein may be used to produce multiple compounds which can be screened to identify GCC ligands.

5 U.S. Patent No. 5,252,743 issued to Barrett et al. on October 12, 1993, which is incorporated herein by reference, refers to spatially addressable immobilization of anti-ligands on surfaces. The methods and compositions described therein may be used to identify GCC ligands.

U.S. Patent No. 5,424,186 issued to Foder et al. on June 13, 1995, which is  
10 incorporated herein by reference, refers to a very large scale immobilized polymer synthesis. The method of synthesizing oligonucleotides described therein may be used to identify GCC ligands.

U.S. Patent No. 5,420,328 issued to Campbell on May 30, 1995, which is incorporated herein by reference, refers to methods of synthesis of phosphonate esters.  
15 The phosphonate esters so produced may be screened to identify compounds which are GCC ligands.

U.S. Patent No. 5,288,514 issued to Ellman on February 22, 1994, which is incorporated herein by reference, refers to solid phase and combinatorial synthesis of benzodiazepine compounds on a solid support. Such methods and compounds may be  
20 used to identify GCC ligands.

As noted above, GCC ligands may also be antibodies and fragments thereof. Indeed, antibodies raised to unique determinants of these receptors will recognize that protein, and only that protein and, consequently, can serve as a specific targeting molecule which can be used to direct novel diagnostics and therapeutics to this unique marker. In  
25 addition, these antibodies can be used to identify the presence of GCC or fragments thereof in biological samples.



**CLAIMS**

1. An *in vitro* method of screening an individual who is suspected of having primary and/or metastatic stomach or esophageal cancer for primary and/or metastatic stomach or esophageal cancer cells comprising the steps of examining a sample of  
5 extraintestinal tissue and/or body fluids from an individual to determine whether GCC is being expressed by cells in said sample wherein expression of said GCC indicates a possibility of primary and/or metastatic stomach or esophageal cancer cells in said sample.
2. The method of claim 1 wherein expression of said GCC by said cells is determined by detecting the presence of GCC gene transcription product.
- 10 3. The method of claim 1 wherein expression of said GCC by said cells is determined by polymerase chain reaction wherein said sample is contacted with primers that selectively amplify GCC gene transcript or cDNA generated therefrom.
4. The method of claim 1 wherein expression of said GCC by said cells is determined by immunoassay wherein said sample is contacted with antibodies that  
15 specifically bind to GCC gene translation product.
5. The method of claim 1 wherein said sample is body fluid.
6. The method of claim 1 wherein said sample is blood.
7. The method of claim 1 wherein said sample is lymphatic tissue and/or fluid.
8. The method of claim 1 wherein said sample is a lymph node sample.
- 20 9. The method of claim 1 wherein the individual has previously been diagnosed with having stomach or esophageal cancer.

10. The method of claim 1 wherein the individual has previously been diagnosed with and treated for stomach or esophageal cancer
11. An *in vitro* method of screening an individual suspected of having primary and/or metastatic stomach or esophageal cancer for primary and/or metastatic stomach or  
5 esophageal cancer cells comprising the steps of examining a sample of extraintestinal tissue and/or body fluids from an individual to determine whether GCC gene transcription or translation product is present in said sample wherein the presence of GCC gene transcription or translation product in said sample indicates that the individual may have primary and/or metastatic stomach or esophageal cancer cells in said sample.
- 10 12. The method of claim 10 comprising the steps of examining a sample of extraintestinal tissue and/or body fluids from an individual to determine whether GCC gene transcription product is present in said sample.
13. The method of claim 12 wherein the presence of GCC gene transcription product is determined by polymerase chain reaction wherein said sample is contacted with  
15 primers that selectively amplify GCC gene transcript or cDNA generated therefrom.
14. The method of claim 11 wherein the presence of GCC gene translation product is determined by immunoassay wherein said sample is contacted with antibodies that specifically bind to GCC gene translation product.
15. The method of claim 11 wherein said sample is body fluid.
- 20 16. The method of claim 11 wherein said sample is blood.
17. The method of claim 11 wherein said sample is lymphatic tissue and/or fluid.
18. The method of claim 11 wherein said sample is a lymph node sample.

19. The method of claim 11 wherein the individual has previously been diagnosed with having stomach or esophageal cancer.
20. The method of claim 11 wherein the individual has previously been diagnosed with and treated for stomach or esophageal cancer
- 5 21. An *in vitro* method of confirming that a tumor cell removed from a patient suspected of having stomach or esophageal cancer cells is a stomach or esophageal tumor cell comprising the step of determining whether a tumor cell expresses GCC wherein expression of GCC indicates that the tumor cell is a stomach or esophageal tumor cell.
22. The method of claim 21 wherein expression of GCC by said tumor cell is  
10 determined by detecting the presence of GCC gene transcription product.
23. The method of claim 21 wherein expression of GCC by said tumor cell is determined by polymerase chain reaction wherein mRNA from said tumor cell or cDNA generated therefrom is contacted with primers that selectively amplify GCC gene transcript or cDNA generated therefrom.
- 15 24. The method of claim 21 wherein expression of GCC by said tumor cell is determined by immunoassay wherein protein from said tumor cell is contacted with antibodies that specifically bind to GCC gene translation product.
25. A method of diagnosing an individual who has stomach cancer comprising the steps of examining a sample of stomach tissue to detect the presence of GCC transcript  
20 or translation product wherein the presence of GCC transcript or translation product in a stomach sample indicates stomach cancer.
26. The method of claim 25 comprising the steps of examining said sample of stomach tissue to determine whether GCC gene transcription product is present in said sample.

27. The method of claim 26 wherein the presence of GCC gene transcription product is determined by polymerase chain reaction wherein said sample is contacted with primers that selectively amplify GCC gene transcript or cDNA generated therefrom.
28. The method of claim 26 wherein the presence of GCC gene translation product is determined by immunoassay wherein said sample is contacted with antibodies that specifically bind to GCC gene translation product.
29. A method of diagnosing an individual who has esophageal cancer comprising the steps of examining a sample of esophagus tissue to detect the presence of GCC transcript or translation product wherein the presence of GCC transcript or translation product in an esophageal sample indicates esophageal cancer.
30. The method of claim 29 comprising the steps of examining said sample of esophageal tissue to determine whether GCC gene transcription product is present in said sample.
31. The method of claim 30 wherein the presence of GCC gene transcription product is determined by polymerase chain reaction wherein said sample is contacted with primers that selectively amplify GCC gene transcript or cDNA generated therefrom.
32. The method of claim 29 wherein the presence of GCC gene translation product is determined by immunoassay wherein said sample is contacted with antibodies that specifically bind to GCC gene translation product.
33. A kit for diagnosing an individual who has stomach and/or esophageal cancer comprising either:
- a) a container comprising polymerase chain reaction primers that selectively amplify GCC gene transcript or cDNA generated therefrom;
  - and one or more of:

5 a container comprising a positive PCR assay control sample,  
 a container comprising a negative PCR assay control sample,  
 instructions for obtaining and/or processing a sample,  
 instructions for performing a PCR diagnostic assay, and  
 photographs or illustrations depicting a positive result and/or a  
 negative result of a PCR diagnostic assay; or

b) a container comprising antibodies that specifically bind to GCC gene  
 translation product;

and one or more of:

10 a container comprising a positive immunoassay control  
 sample,

a container comprising a negative immunoassay control  
 sample,

15 instructions for obtaining and/or processing a sample,  
 instructions for performing an immuno diagnostic assay, and  
 photographs or illustrations depicting a positive result and/or a  
 negative result of an immuno diagnostic assay.

34. A method of treating an individual suspected of suffering from primary  
 and/or stomach or esophageal cancer comprising the steps of administering to said  
 20 individual a therapeutically effective amount of a composition comprising:

- i) an ST receptor ligand; and
- ii) an active agent.

35. The method of claim 34 wherein the ST receptor ligand is conjugated to the  
 active agent.

25 36. The method of claim 34 wherein said an active agent is selected from the  
 group consisting of: methotrexate, doxorubicin, daunorubicin, cytosinarabioside,  
 etoposide, 5-4 fluorouracil, melphalan, chlorambucil, *cis*-platinum, vindesine, mitomycin,  
 bleomycin, purothionin, macromomycin, 1,4-benzoquinone derivatives, trenimon, ricin,

ricin A chain, *Pseudomonas* exotoxin, diphtheria toxin, *Clostridium perfringens*  
 phospholipase C, bovine pancreatic ribonuclease, pokeweed antiviral protein, abrin, abrin  
 A chain, cobra venom factor, gelonin, saporin, modeccin, viscumin, volkensin, alkaline  
 phosphatase, nitroimidazole, metronidazole, misonidazole,  $^{47}\text{Sc}$ ,  $^{67}\text{Cu}$ ,  $^{90}\text{Y}$ ,  $^{109}\text{Pd}$ ,  $^{123}\text{I}$ ,  $^{125}\text{I}$ ,  
 5  $^{131}\text{I}$ ,  $^{186}\text{Re}$ ,  $^{188}\text{Re}$ ,  $^{199}\text{Au}$ ,  $^{211}\text{At}$ ,  $^{212}\text{Pb}$ ,  $^{212}\text{B}$ ,  $^{32}\text{P}$  and  $^{33}\text{P}$ ,  $^{71}\text{Ge}$ ,  $^{77}\text{As}$ ,  $^{103}\text{Pb}$ ,  $^{105}\text{Rh}$ ,  $^{111}\text{Ag}$ ,  $^{119}\text{Sb}$ ,  
 $^{121}\text{Sn}$ ,  $^{131}\text{Cs}$ ,  $^{143}\text{Pr}$ ,  $^{161}\text{Tb}$ ,  $^{177}\text{Lu}$ ,  $^{191}\text{Os}$ ,  $^{193\text{M}}\text{Pt}$ ,  $^{197}\text{Hg}$ ,  $^{43}\text{K}$ ,  $^{52}\text{Fe}$ ,  $^{57}\text{Co}$ ,  $^{67}\text{Cu}$ ,  $^{67}\text{Ga}$ ,  $^{68}\text{Ga}$ ,  $^{77}\text{Br}$ ,  
 $^{81}\text{Rb}/^{81\text{M}}\text{Kr}$ ,  $^{87\text{M}}\text{Sr}$ ,  $^{99\text{M}}\text{Tc}$ ,  $^{111}\text{In}$ ,  $^{113\text{M}}\text{In}$ ,  $^{123}\text{I}$ ,  $^{125}\text{I}$ ,  $^{127}\text{Cs}$ ,  $^{129}\text{Cs}$ ,  $^{131}\text{I}$ ,  $^{132}\text{I}$ ,  $^{197}\text{Hg}$ ,  $^{203}\text{Pb}$  and  $^{206}\text{Bi}$ .

37. A method of radioimaging primary and/or stomach or esophageal cancer  
 cells comprising the steps of administering to an individual a composition comprising an  
 10 ST receptor ligand linked to a detectable agent.

38. The method of claim 37 wherein said detectable agent is selected from the  
 group consisting of:  $^{47}\text{Sc}$ ,  $^{67}\text{Cu}$ ,  $^{90}\text{Y}$ ,  $^{109}\text{Pd}$ ,  $^{123}\text{I}$ ,  $^{125}\text{I}$ ,  $^{131}\text{I}$ ,  $^{186}\text{Re}$ ,  $^{188}\text{Re}$ ,  $^{199}\text{Au}$ ,  $^{211}\text{At}$ ,  $^{212}\text{Pb}$ ,  $^{212}\text{B}$ ,  
 $^{32}\text{P}$  and  $^{33}\text{P}$ ,  $^{71}\text{Ge}$ ,  $^{77}\text{As}$ ,  $^{103}\text{Pb}$ ,  $^{105}\text{Rh}$ ,  $^{111}\text{Ag}$ ,  $^{119}\text{Sb}$ ,  $^{121}\text{Sn}$ ,  $^{131}\text{Cs}$ ,  $^{143}\text{Pr}$ ,  $^{161}\text{Tb}$ ,  $^{177}\text{Lu}$ ,  $^{191}\text{Os}$ ,  $^{193\text{M}}\text{Pt}$ ,  
 $^{197}\text{Hg}$ ,  $^{43}\text{K}$ ,  $^{52}\text{Fe}$ ,  $^{57}\text{Co}$ ,  $^{67}\text{Cu}$ ,  $^{67}\text{Ga}$ ,  $^{68}\text{Ga}$ ,  $^{77}\text{Br}$ ,  $^{81}\text{Rb}/^{81\text{M}}\text{Kr}$ ,  $^{87\text{M}}\text{Sr}$ ,  $^{99\text{M}}\text{Tc}$ ,  $^{111}\text{In}$ ,  $^{113\text{M}}\text{In}$ ,  $^{123}\text{I}$ ,  $^{125}\text{I}$ ,  
 15  $^{127}\text{Cs}$ ,  $^{129}\text{Cs}$ ,  $^{131}\text{I}$ ,  $^{132}\text{I}$ ,  $^{197}\text{Hg}$ ,  $^{203}\text{Pb}$  and  $^{206}\text{Bi}$ .

## SEQUENCE LISTING

<110> Thomas Jefferson University  
Waldman, Scott A.  
Park, Jason  
Schulz, Stephanie

<120> Compositions And Methods For Identifying And Targeting Cancer Cells Of Alimentary Canal Origin

<130> TJU2469

<150> 60/192,229

<151> 2000-03-27

<160> 2

<170> PatentIn version 3.0

<210> 1

<211> 3787

<212> DNA

<213> Homo sapiens

<220>

<221> CDS

<222> (118)..(3336)

<400> 1

```

tggagtgggc tgaggggactc cactagaggc tgtccatctg gattccctgc ctccctagga      60
gcccaacaga gcaaagcaag tgggcacaag gagtatgggtt ctaacgtgat tgggggtc      117
atg aag acg ttg ctg ttg gac ttg gct ttg tgg tca ctg ctc ttc cag      165
Met Lys Thr Leu Leu Leu Asp Leu Ala Leu Trp Ser Leu Leu Phe Gln
1          5          10          15
ccc ggg tgg ctg tcc ttt agt tcc cag gtg agt cag aac tgc cac aat      213
Pro Gly Trp Leu Ser Phe Ser Ser Gln Val Ser Gln Asn Cys His Asn
          20          25          30
ggc agc tat gaa atc agc gtc ctg atg atg ggc aac tca gcc ttt gca      261
Gly Ser Tyr Glu Ile Ser Val Leu Met Met Gly Asn Ser Ala Phe Ala
          35          40          45
gag ccc ctg aaa aac ttg gaa gat gcg gtg aat gag ggg ctg gaa ata      309
Glu Pro Leu Lys Asn Leu Glu Asp Ala Val Asn Glu Gly Leu Glu Ile
          50          55          60
gtg aga gga cgt ctg caa aat gct ggc cta aat gtg act gtg aac gct      357
Val Arg Gly Arg Leu Gln Asn Ala Gly Leu Asn Val Thr Val Asn Ala
65          70          75          80
act ttc atg tat tcg gat ggt ctg att cat aac tca ggc gac tgc cgg      405
Thr Phe Met Tyr Ser Asp Gly Leu Ile His Asn Ser Gly Asp Cys Arg
          85          90          95
agt agc acc tgt gaa ggc ctc gac cta ctc agg aaa att tca aat gca      453
Ser Ser Thr Cys Glu Gly Leu Asp Leu Leu Arg Lys Ile Ser Asn Ala
          100          105          110
caa cgg atg ggc tgt gtc ctc ata ggg ccc tca tgt aca tac tcc acc      501
Gln Arg Met Gly Cys Val Leu Ile Gly Pro Ser Cys Thr Tyr Ser Thr
          115          120          125

```

ttc cag atg tac ctt gac aca gaa ttg agc tac ccc atg atc tca gct Phe Gln Met Tyr Leu Asp Thr Glu Leu Ser Tyr Pro Met Ile Ser Ala 130 135 140	549
gga agt ttt gga ttg tca tgt gac tat aaa gaa acc tta acc agg ctg Gly Ser Phe Gly Leu Ser Cys Asp Tyr Lys Glu Thr Leu Thr Arg Leu 145 150 155 160	597
atg tct cca gct aga aag ttg atg tac ttc ttg gtt aac ttt tgg aaa Met Ser Pro Ala Arg Lys Leu Met Tyr Phe Leu Val Asn Phe Trp Lys 165 170 175	645
acc aac gat ctg ccc ttc aaa act tat tcc tgg agc act tcg tat gtt Thr Asn Asp Leu Pro Phe Lys Thr Tyr Ser Trp Ser Thr Ser Tyr Val 180 185 190	693
tac aag aat ggt aca gaa act gag gac tgt ttc tgg tac ctt aat gct Tyr Lys Asn Gly Thr Glu Thr Glu Asp Cys Phe Trp Tyr Leu Asn Ala 195 200 205	741
ctg gag gct agc gtt tcc tat ttc tcc cac gaa ctc ggc ttt aag gtg Leu Glu Ala Ser Val Ser Tyr Phe Ser His Glu Leu Gly Phe Lys Val 210 215 220	789
gtg tta aga caa gat aag gag ttt cag gat atc tta atg gac cac aac Val Leu Arg Gln Asp Lys Glu Phe Gln Asp Ile Leu Met Asp His Asn 225 230 235 240	837
agg aaa agc aat gtg att att atg tgt ggt ggt cca gag ttc ctc tac Arg Lys Ser Asn Val Ile Ile Met Cys Gly Pro Glu Phe Leu Tyr 245 250 255	885
aag ctg aag ggt gac cga gca gtg gct gaa gac att gtc att att cta Lys Leu Lys Gly Asp Arg Ala Val Ala Glu Asp Ile Val Ile Ile Leu 260 265 270	933
gtg gat ctt ttc aat gac cag tac ttg gag gac aat gtc aca gcc cct Val Asp Leu Phe Asn Asp Gln Tyr Leu Glu Asp Asn Val Thr Ala Pro 275 280 285	981
gac tat atg aaa aat gtc ctt gtt ctg acg ctg tct cct ggg aat tcc Asp Tyr Met Lys Asn Val Leu Val Leu Thr Leu Ser Pro Gly Asn Ser 290 295 300	1029
ctt cta aat agc tct ttc tcc agg aat cta tca cca aca aaa cga gac Leu Leu Asn Ser Ser Phe Ser Arg Asn Leu Ser Pro Thr Lys Arg Asp 305 310 315 320	1077
ttt cgt ctt gcc tat ttg aat gga atc ctc gtc ttt gga cat atg ctg Phe Arg Leu Ala Tyr Leu Asn Gly Ile Leu Val Phe Gly His Met Leu 325 330 335	1125
aag ata ttt ctt gaa aat gga gaa aat att acc acc ccc aaa ttt gct Lys Ile Phe Leu Glu Asn Gly Glu Asn Ile Thr Thr Pro Lys Phe Ala 340 345 350	1173
cat gcc ttc agg aat ctc act ttt gaa ggg tat gac ggt cca gtg acc His Ala Phe Arg Asn Leu Thr Phe Glu Gly Tyr Asp Gly Pro Val Thr 355 360 365	1221
ttg gat gac tgg ggg gat gtt gac agt acc atg gtg ctt ctg tat acc Leu Asp Asp Trp Gly Asp Val Asp Ser Thr Met Val Leu Leu Tyr Thr 370 375 380	1269
tct gtg gac acc aag aaa tac aag gtt ctt ttg acc tat gat acc cac	1317



Ser 385	Val	Asp	Thr	Lys	Lys 390	Tyr	Lys	Val	Leu	Leu 395	Thr	Tyr	Asp	Thr	His 400	
gta	aat	aag	acc	tat	cct	gtg	gat	atg	agc	ccc	aca	ttc	act	tgg	aag	1365
Val	Asn	Lys	Thr	Tyr 405	Pro	Val	Asp	Met	Ser 410	Pro	Thr	Phe	Thr	Trp 415	Lys	
aac	tct	aaa	ctt	cct	aat	gat	att	aca	ggc	cgg	ggc	cct	cag	atc	ctg	1413
Asn	Ser	Lys	Leu	Pro	Asn	Asp	Ile	Thr 425	Gly	Arg	Gly	Pro	Gln	Ile	Leu	
			420										430			
atg	att	gca	gtc	ttc	acc	ctc	act	gga	gct	gtg	gtg	ctg	ctc	ctg	ctc	1461
Met	Ile	Ala	Val	Phe	Thr	Leu	Thr	Gly 440	Ala	Val	Val	Leu	Leu	Leu	Leu	
		435										445				
gtc	gct	ctc	ctg	atg	ctc	aga	aaa	tat	aga	aaa	gat	tat	gaa	ctt	cgt	1509
Val	Ala	Leu	Leu	Met	Leu	Arg	Lys	Tyr	Arg	Lys	Asp 460	Tyr	Glu	Leu	Arg	
	450					455										
cag	aaa	aaa	tgg	tcc	cac	att	cct	cct	gaa	aat	atc	ttt	cct	ctg	gag	1557
Gln	Lys	Lys	Trp	Ser	His	Ile	Pro	Pro	Glu	Asn	Ile	Phe	Pro	Leu	Glu	
465					470					475					480	
acc	aat	gag	acc	aat	cat	gtt	agc	ctc	aag	atc	gat	gat	gac	aaa	aga	1605
Thr	Asn	Glu	Thr	Asn	His	Val	Ser	Leu	Lys 490	Ile	Asp	Asp	Asp	Lys	Arg	
				485										495		
cga	gat	aca	atc	cag	aga	cta	cga	cag	tgc	aaa	tac	gtc	aaa	aag	cga	1653
Arg	Asp	Thr	Ile	Gln	Arg	Leu	Arg	Gln	Cys	Lys	Tyr	Val	Lys	Lys	Arg	
			500					505					510			
gtg	att	ctc	aaa	gat	ctc	aag	cac	aat	gat	ggt	aat	ttc	act	gaa	aaa	1701
Val	Ile	Leu	Lys	Asp	Leu	Lys	His	Asn	Asp	Gly	Asn	Phe	Thr	Glu	Lys	
		515					520					525				
cag	aag	ata	gaa	ttg	aac	aag	ttg	ctt	cag	att	gac	tat	tac	acc	cta	1749
Gln	Lys	Ile	Glu	Leu	Asn	Lys	Leu	Leu	Gln	Ile	Asp	Tyr	Tyr	Thr	Leu	
	530					535					540					
acc	aag	ttc	tac	ggg	aca	gtg	aaa	ctg	gat	acc	atg	atc	ttc	ggg	gtg	1797
Thr	Lys	Phe	Tyr	Gly	Thr	Val	Lys	Leu	Asp	Thr	Met	Ile	Phe	Gly	Val	
545					550					555					560	
ata	gaa	tac	tgt	gag	aga	gga	tcc	ctc	cgg	gaa	gtt	tta	aat	gac	aca	1845
Ile	Glu	Tyr	Cys	Glu	Arg	Gly	Ser	Leu	Arg	Glu	Val	Leu	Asn	Asp	Thr	
				565					570					575		
att	tcc	tac	cct	gat	ggc	aca	ttc	atg	gat	tgg	gag	ttt	aag	atc	tct	1893
Ile	Ser	Tyr	Pro	Asp	Gly	Thr	Phe	Met	Asp	Trp	Glu	Phe	Lys	Ile	Ser	
			580					585					590			
gtc	ttg	tat	gac	att	gct	aag	gga	atg	tca	tat	ctg	cac	tcc	agt	aag	1941
Val	Leu	Tyr	Asp	Ile	Ala	Lys	Gly	Met	Ser	Tyr	Leu	His	Ser	Ser	Lys	
		595					600					605				
aca	gaa	gtc	cat	ggc	cgt	ctg	aaa	tct	acc	aac	tgc	gta	gtg	gac	agt	1989
Thr	Glu	Val	His	Gly	Arg	Leu	Lys	Ser	Thr	Asn	Cys	Val	Val	Asp	Ser	
	610					615					620					
aga	atg	gtg	gtg	aag	atc	act	gat	ttt	ggc	tgc	aat	tcc	att	ttg	cct	2037
Arg	Met	Val	Val	Lys	Ile	Thr	Asp	Phe	Gly	Cys	Asn	Ser	Ile	Leu	Pro	
625					630					635					640	
cca	aaa	aag	gac	ctg	tgg	aca	gct	cca	gag	cac	ctc	cgc	caa	gcc	aac	2085
Pro	Lys	Lys	Asp	Leu	Trp	Thr	Ala	Pro	Glu	His	Leu	Arg	Gln	Ala	Asn	

645	650	655	
atc tct cag aaa gga gat gtg tac agc tat ggg atc atc gca cag gag Ile Ser Gln Lys Gly Asp Val Tyr Ser Tyr Gly Ile Ile Ala Gln Glu 660 665 670			2133
atc att ctg cgg aaa gaa acc ttc tac act ttg agc tgt cgg gac cgg Ile Ile Leu Arg Lys Glu Thr Phe Tyr Thr Leu Ser Cys Arg Asp Arg 675 680 685			2181
aat gag aag att ttc aga gtg gaa aat tcc aat gga atg aaa ccc ttc Asn Glu Lys Ile Phe Arg Val Glu Asn Ser Asn Gly Met Lys Pro Phe 690 695 700			2229
cgc cca gat tta ttc ttg gaa aca gca gag gaa aaa gag cta gaa gtg Arg Pro Asp Leu Phe Leu Glu Thr Ala Glu Glu Lys Glu Leu Glu Val 705 710 715 720			2277
tac cta ctt gta aaa aac tgt tgg gag gaa gat cca gaa aag aga cca Tyr Leu Leu Val Lys Asn Cys Trp Glu Glu Asp Pro Glu Lys Arg Pro 725 730 735			2325
gat ttc aaa aaa att gag act aca ctt gcc aag ata ttt gga ctt ttt Asp Phe Lys Lys Ile Glu Thr Thr Leu Ala Lys Ile Phe Gly Leu Phe 740 745 750			2373
cat gac caa aaa aat gaa agc tat atg gat acc ttg atc cga cgt cta His Asp Gln Lys Asn Glu Ser Tyr Met Asp Thr Leu Ile Arg Arg Leu 755 760 765			2421
cag cta tat tct cga aac ctg gaa cat ctg gta gag gaa agg aca cag Gln Leu Tyr Ser Arg Asn Leu Glu His Leu Val Glu Glu Arg Thr Gln 770 775 780			2469
ctg tac aag gca gag agg gac agg gct gac aga ctt aac ttt atg ttg Leu Tyr Lys Ala Glu Arg Asp Arg Ala Asp Arg Leu Asn Phe Met Leu 785 790 795 800			2517
ctt cca agg cta gtg gta aag tct ctg aag gag aaa ggc ttt gtg gag Leu Pro Arg Leu Val Val Lys Ser Leu Lys Glu Lys Gly Phe Val Glu 805 810 815			2565
ccg gaa cta tat gag gaa gtt aca atc tac ttc agt gac att gta ggt Pro Glu Leu Tyr Glu Glu Val Thr Ile Tyr Phe Ser Asp Ile Val Gly 820 825 830			2613
ttc act act atc tgc aaa tac agc acc ccc atg gaa gtg gtg gac atg Phe Thr Thr Ile Cys Lys Tyr Ser Thr Pro Met Glu Val Val Asp Met 835 840 845			2661
ctt aat gac atc tat aag agt ttt gac cac att gtt gat cat cat gat Leu Asn Asp Ile Tyr Lys Ser Phe Asp His Ile Val Asp His His Asp 850 855 860			2709
gtc tac aag gtg gaa acc atc ggt gat gcg tac atg gtg gct agt ggt Val Tyr Lys Val Glu Thr Ile Gly Asp Ala Tyr Met Val Ala Ser Gly 865 870 875 880			2757
ttg cct aag aga aat ggc aat cgg cat gca ata gac att gcc aag atg Leu Pro Lys Arg Asn Gly Asn Arg His Ala Ile Asp Ile Ala Lys Met 885 890 895			2805
gcc ttg gaa atc ctc agc ttc atg ggg acc ttt gag ctg gag cat ctt Ala Leu Glu Ile Leu Ser Phe Met Gly Thr Phe Glu Leu Glu His Leu 900 905 910			2853

cct ggc ctc cca ata tgg att cgc att gga gtt cac tct ggt ccc tgt 2901  
 Pro Gly Leu Pro Ile Trp Ile Arg Ile Gly Val His Ser Gly Pro Cys  
 915 920 925  
 gct gct gga gtt gtg gga atc aag atg cct cgt tat tgt cta ttt gga 2949  
 Ala Ala Gly Val Val Gly Ile Lys Met Pro Arg Tyr Cys Leu Phe Gly  
 930 935 940  
 gat acg gtc aac aca gcc tct agg atg gaa tcc act ggc ctc cct ttg 2997  
 Asp Thr Val Asn Thr Ala Ser Arg Met Glu Ser Thr Gly Leu Pro Leu  
 945 950 955 960  
 aga att cac gtg agt ggc tcc acc ata gcc atc ctg aag aga act gag 3045  
 Arg Ile His Val Ser Gly Ser Thr Ile Ala Ile Leu Lys Arg Thr Glu  
 965 970 975  
 tgc cag ttc ctt tat gaa gtg aga gga gaa aca tac tta aag gga aga 3093  
 Cys Gln Phe Leu Tyr Glu Val Arg Gly Glu Thr Tyr Leu Lys Gly Arg  
 980 985 990  
 gga aat gag act acc tac tgg ctg act ggg atg aag gac cag aaa ttc 3141  
 Gly Asn Glu Thr Thr Tyr Trp Leu Thr Gly Met Lys Asp Gln Lys Phe  
 995 1000 1005  
 aac ctg cca acc cct cct act gtg gag aat caa cag cgt ttg caa 3186  
 Asn Leu Pro Thr Pro Pro Thr Val Glu Asn Gln Gln Arg Leu Gln  
 1010 1015 1020  
 gca gaa ttt tca gac atg att gcc aac tct tta cag aaa aga cag 3231  
 Ala Glu Phe Ser Asp Met Ile Ala Asn Ser Leu Gln Lys Arg Gln  
 1025 1030 1035  
 gca gca ggg ata aga agc caa aaa ccc aga cgg gta gcc agc tat 3276  
 Ala Ala Gly Ile Arg Ser Lys Pro Arg Arg Val Ala Ser Tyr  
 1040 1045 1050  
 aaa aaa ggc act ctg gaa tac ttg cag ctg aat acc aca gac aag 3321  
 Lys Lys Gly Thr Leu Glu Tyr Leu Gln Leu Asn Thr Thr Asp Lys  
 1055 1060 1065  
 gag agc acc tat ttt taaacctaaa tgagggtataa ggactcacac aaattaaaat 3376  
 Glu Ser Thr Tyr Phe  
 1070  
 acagctgcac tgaggccagg caccctcagg tgcctgaaa gcttactttc ctgagacctc 3436  
 atgaggcaga aatgtcttag gcttggtgc cctgtttgga ccatggactt tctttgcatg 3496  
 aatcagatgt gttctcagtg aaataactac ctccactct ggaaccttat tccagcagtt 3556  
 gttccaggga gcttctacct ggaaaagaaa agaatttcat ttattttttg tttgtttatt 3616  
 tttatcgttt ttgtttactg gctttccttc tgtattcata agatttttta aattgtcata 3676  
 attatatttt aaatacccat cttcattaaa gtatatttaa ctcataattt ttgcagaaaa 3736  
 tatgctatat attaggcaag aataaaagct aaagggtttcc caaaaaaaaa a 3787

<210> 2  
 <211> 1073  
 <212> PRT  
 <213> Homo sapiens

&lt;400&gt; 2

Met Lys Thr Leu Leu Leu Asp Leu Ala Leu Trp Ser Leu Leu Phe Gln  
 1 5 10 15

Pro Gly Trp Leu Ser Phe Ser Ser Gln Val Ser Gln Asn Cys His Asn  
 20 25 30

Gly Ser Tyr Glu Ile Ser Val Leu Met Met Gly Asn Ser Ala Phe Ala  
 35 40 45

Glu Pro Leu Lys Asn Leu Glu Asp Ala Val Asn Glu Gly Leu Glu Ile  
 50 55 60

Val Arg Gly Arg Leu Gln Asn Ala Gly Leu Asn Val Thr Val Asn Ala  
 65 70 75 80

Thr Phe Met Tyr Ser Asp Gly Leu Ile His Asn Ser Gly Asp Cys Arg  
 85 90 95

Ser Ser Thr Cys Glu Gly Leu Asp Leu Leu Arg Lys Ile Ser Asn Ala  
 100 105 110

Gln Arg Met Gly Cys Val Leu Ile Gly Pro Ser Cys Thr Tyr Ser Thr  
 115 120 125

Phe Gln Met Tyr Leu Asp Thr Glu Leu Ser Tyr Pro Met Ile Ser Ala  
 130 135 140

Gly Ser Phe Gly Leu Ser Cys Asp Tyr Lys Glu Thr Leu Thr Arg Leu  
 145 150 155 160

Met Ser Pro Ala Arg Lys Leu Met Tyr Phe Leu Val Asn Phe Trp Lys  
 165 170 175

Thr Asn Asp Leu Pro Phe Lys Thr Tyr Ser Trp Ser Thr Ser Tyr Val  
 180 185 190

Tyr Lys Asn Gly Thr Glu Thr Glu Asp Cys Phe Trp Tyr Leu Asn Ala  
 195 200 205

Leu Glu Ala Ser Val Ser Tyr Phe Ser His Glu Leu Gly Phe Lys Val  
 210 215 220

Val Leu Arg Gln Asp Lys Glu Phe Gln Asp Ile Leu Met Asp His Asn  
 225 230 235 240

Arg Lys Ser Asn Val Ile Ile Met Cys Gly Gly Pro Glu Phe Leu Tyr  
 245 250 255

Lys Leu Lys Gly Asp Arg Ala Val Ala Glu Asp Ile Val Ile Ile Leu  
 260 265 270  
 Val Asp Leu Phe Asn Asp Gln Tyr Leu Glu Asp Asn Val Thr Ala Pro  
 275 280 285  
 Asp Tyr Met Lys Asn Val Leu Val Leu Thr Leu Ser Pro Gly Asn Ser  
 290 295 300  
 Leu Leu Asn Ser Ser Phe Ser Arg Asn Leu Ser Pro Thr Lys Arg Asp  
 305 310 315 320  
 Phe Arg Leu Ala Tyr Leu Asn Gly Ile Leu Val Phe Gly His Met Leu  
 325 330 335  
 Lys Ile Phe Leu Glu Asn Gly Glu Asn Ile Thr Thr Pro Lys Phe Ala  
 340 345 350  
 His Ala Phe Arg Asn Leu Thr Phe Glu Gly Tyr Asp Gly Pro Val Thr  
 355 360 365  
 Leu Asp Asp Trp Gly Asp Val Asp Ser Thr Met Val Leu Leu Tyr Thr  
 370 375 380  
 Ser Val Asp Thr Lys Lys Tyr Lys Val Leu Leu Thr Tyr Asp Thr His  
 385 390 395 400  
 Val Asn Lys Thr Tyr Pro Val Asp Met Ser Pro Thr Phe Thr Trp Lys  
 405 410 415  
 Asn Ser Lys Leu Pro Asn Asp Ile Thr Gly Arg Gly Pro Gln Ile Leu  
 420 425 430  
 Met Ile Ala Val Phe Thr Leu Thr Gly Ala Val Val Leu Leu Leu Leu  
 435 440 445  
 Val Ala Leu Leu Met Leu Arg Lys Tyr Arg Lys Asp Tyr Glu Leu Arg  
 450 455 460  
 Gln Lys Lys Trp Ser His Ile Pro Pro Glu Asn Ile Phe Pro Leu Glu  
 465 470 475 480  
 Thr Asn Glu Thr Asn His Val Ser Leu Lys Ile Asp Asp Asp Lys Arg  
 485 490 495  
 Arg Asp Thr Ile Gln Arg Leu Arg Gln Cys Lys Tyr Val Lys Lys Arg  
 500 505 510

Val Ile Leu Lys Asp Leu Lys His Asn Asp Gly Asn Phe Thr Glu Lys  
 515 520 525

Gln Lys Ile Glu Leu Asn Lys Leu Leu Gln Ile Asp Tyr Tyr Thr Leu  
 530 535 540

Thr Lys Phe Tyr Gly Thr Val Lys Leu Asp Thr Met Ile Phe Gly Val  
 545 550 555 560

Ile Glu Tyr Cys Glu Arg Gly Ser Leu Arg Glu Val Leu Asn Asp Thr  
 565 570 575

Ile Ser Tyr Pro Asp Gly Thr Phe Met Asp Trp Glu Phe Lys Ile Ser  
 580 585 590

Val Leu Tyr Asp Ile Ala Lys Gly Met Ser Tyr Leu His Ser Ser Lys  
 595 600 605

Thr Glu Val His Gly Arg Leu Lys Ser Thr Asn Cys Val Val Asp Ser  
 610 615 620

Arg Met Val Val Lys Ile Thr Asp Phe Gly Cys Asn Ser Ile Leu Pro  
 625 630 635 640

Pro Lys Lys Asp Leu Trp Thr Ala Pro Glu His Leu Arg Gln Ala Asn  
 645 650 655

Ile Ser Gln Lys Gly Asp Val Tyr Ser Tyr Gly Ile Ile Ala Gln Glu  
 660 665 670

Ile Ile Leu Arg Lys Glu Thr Phe Tyr Thr Leu Ser Cys Arg Asp Arg  
 675 680 685

Asn Glu Lys Ile Phe Arg Val Glu Asn Ser Asn Gly Met Lys Pro Phe  
 690 695 700

Arg Pro Asp Leu Phe Leu Glu Thr Ala Glu Glu Lys Glu Leu Glu Val  
 705 710 715 720

Tyr Leu Leu Val Lys Asn Cys Trp Glu Glu Asp Pro Glu Lys Arg Pro  
 725 730 735

Asp Phe Lys Lys Ile Glu Thr Thr Leu Ala Lys Ile Phe Gly Leu Phe  
 740 745 750

His Asp Gln Lys Asn Glu Ser Tyr Met Asp Thr Leu Ile Arg Arg Leu  
 755 760 765

Gln Leu Tyr Ser Arg Asn Leu Glu His Leu Val Glu Glu Arg Thr Gln

770

775

780

Leu Tyr Lys Ala Glu Arg Asp Arg Ala Asp Arg Leu Asn Phe Met Leu  
 785 790 795 800

Leu Pro Arg Leu Val Val Lys Ser Leu Lys Glu Lys Gly Phe Val Glu  
 805 810 815

Pro Glu Leu Tyr Glu Glu Val Thr Ile Tyr Phe Ser Asp Ile Val Gly  
 820 825 830

Phe Thr Thr Ile Cys Lys Tyr Ser Thr Pro Met Glu Val Val Asp Met  
 835 840 845

Leu Asn Asp Ile Tyr Lys Ser Phe Asp His Ile Val Asp His His Asp  
 850 855 860

Val Tyr Lys Val Glu Thr Ile Gly Asp Ala Tyr Met Val Ala Ser Gly  
 865 870 875 880

Leu Pro Lys Arg Asn Gly Asn Arg His Ala Ile Asp Ile Ala Lys Met  
 885 890 895

Ala Leu Glu Ile Leu Ser Phe Met Gly Thr Phe Glu Leu Glu His Leu  
 900 905 910

Pro Gly Leu Pro Ile Trp Ile Arg Ile Gly Val His Ser Gly Pro Cys  
 915 920 925

Ala Ala Gly Val Val Gly Ile Lys Met Pro Arg Tyr Cys Leu Phe Gly  
 930 935 940

Asp Thr Val Asn Thr Ala Ser Arg Met Glu Ser Thr Gly Leu Pro Leu  
 945 950 955 960

Arg Ile His Val Ser Gly Ser Thr Ile Ala Ile Leu Lys Arg Thr Glu  
 965 970 975

Cys Gln Phe Leu Tyr Glu Val Arg Gly Glu Thr Tyr Leu Lys Gly Arg  
 980 985 990

Gly Asn Glu Thr Thr Tyr Trp Leu Thr Gly Met Lys Asp Gln Lys Phe  
 995 1000 1005

Asn Leu Pro Thr Pro Pro Thr Val Glu Asn Gln Gln Arg Leu Gln  
 1010 1015 1020

Ala Glu Phe Ser Asp Met Ile Ala Asn Ser Leu Gln Lys Arg Gln  
 1025 1030 1035

Ala Ala Gly Ile Arg Ser Gln Lys Pro Arg Arg Val Ala Ser Tyr  
1040 1045 1050

Lys Lys Gly Thr Leu Glu Tyr Leu Gln Leu Asn Thr Thr Asp Lys  
1055 1060 1065

Glu Ser Thr Tyr Phe  
1070



# INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/09790

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : C12Q 1/68; G01N 33/53, 33/574; A61K 39/00, 39/38

US CL : 435/ 6, 7.1, 7.23, 975; 424/ 184.1, 193.1

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 435/ 6, 7.1, 7.23, 975; 424/ 184.1, 193.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)  
STN, WEST, MEDLINE

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	LONDON et al. Siganl transduction pathways via guanylin and uroguanylin in stomach and intestine. Amer.J.Physio. 1997, Vol. 273, pages G93-G105, especially pages G93-94, and Fig. 5.	1-28,33
Y	US 5,601,990 A (WALDMAN) 11 February 1997 (11.02.1997), cols 3-4, 7-8, 7 and 20.	1-28,33
Y	US 5,879,656 A (WALDMAN) 09 March 1999 (09.03.1999), entire article, especially column 8, lines 3-10, and lines 47-49, and claims.	1-28, 33-38
A	LI et al. Peptide-regulated guanylate cyclase pathways in rat colon: in situ localization of GCA, GCC, and guanylin mRNA. Amer.J.Physio. 1993, Vol. 265, pages G394-G402, entire article, especially page G394.	1-38

☐ Further documents are listed in the continuation of Box C.

☐ See patent family annex.

\* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

02 June 2001 (02.06.2001)

Date of mailing of the international search report

12 JUL 2001

Name and mailing address of the ISA/US

Commissioner of Patents and Trademarks  
Box PCT  
Washington, D.C. 20231

Facsimile No. (703)305-3230

Authorized officer

Gary B. Nickol, Ph.D.

Telephone No. 703-308-0196

专利名称(译)	用于鉴定和靶向消化道来源的癌细胞的组合物和方法		
公开(公告)号	<a href="#">EP1272665A1</a>	公开(公告)日	2003-01-08
申请号	EP2001922739	申请日	2001-03-27
[标]申请(专利权)人(译)	托马斯杰弗逊大学		
申请(专利权)人(译)	托马斯杰弗逊大学		
当前申请(专利权)人(译)	托马斯杰弗逊大学		
[标]发明人	WALDMAN SCOTT A PARK JASON SCHULZ STEPHANIE		
发明人	WALDMAN, SCOTT, A. PARK, JASON SCHULZ, STEPHANIE		
IPC分类号	G01N33/53 A61K31/196 A61K31/407 A61K31/4164 A61K31/475 A61K31/513 A61K31/519 A61K31/7034 A61K31/704 A61K31/7048 A61K31/7068 A61K31/711 A61K33/24 A61K38/00 A61K38/43 A61K38/46 A61K39/00 A61K45/00 A61K47/48 A61K48/00 A61K51/00 A61P1/00 A61P35/00 A61P35/04 C12N15/09 C12Q1/02 C12Q1/48 C12Q1/6886 G01N33/566 G01N33/574 C12Q1/68 C12P19/34 C07H21/04		
CPC分类号	A61K31/7034 A61K31/7048 A61P1/00 A61P35/00 A61P35/04 A61P37/04 A61K47/6871 C12Q1/6886 C12Q2600/158 G01N33/57407 G01N33/57419 G01N33/57446 A61K9/127 A61K39/0011 A61K39/001102 A61K51/1075 C07K16/3046 C07K16/40 C12N15/113 G01N2333/988		
优先权	60/192229 2000-03-27 US		
其他公开文献	EP1272665A4 EP1272665B1		
外部链接	<a href="#">Espacenet</a>		

#### 摘要(译)

公开了初级和/或转移性胃或食管癌的筛选和诊断试剂，试剂盒和方法。  
公开了用于成像和治疗原发性或/或转移性胃或食管癌的组合物和方法。  
公开了用于治疗 and 预防原发性或/或转移性胃或食管癌的疫苗组合物和方法。