Role of novel tumor suppressors in colon cancer: Mechanisms and therapeutic opportunities

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Lubna Mehdawi worked as a physician in Benghazi, Libya until 2011 when she moved to Malmö, Sweden for her PhD studies at the Clinical Research Center, Lund University. The main focus of her doctoral thesis is clinical significant and therapeutic opportunities of tumor suppressors in colon cancer patients.
Role of novel tumor suppressors in colon cancer: Mechanisms and therapeutic opportunities

Lubna Mohammed Mehdawi

DOCTORAL DISSERTATION
By due permission of the Faculty of Medicine, Lund University, Sweden.
To be defended at Aulan (Level 3), Kvinnokliniken, Jan Waldenströms gata 47.
On Thursday 27th October 2016 and at 9.15 am.

Faculty opponent:
Prof Marene Landström
Departement of Pathology,
Umeå University
Abstract
Colon cancer is the third most common cancer in the world and the fourth most common cause of cancer related deaths. Inflammation is one of the risk factors for development of colon cancer. Interestingly immune cells as mast cells and inflammatory mediators as LTC₄ play an important role in colon cancer. Genetic predisposition, which might lead to either activation of oncogenes or inhibition of tumor suppressor genes, are risk factors of colon cancer development.

The aim of my thesis was to evaluate the clinical significance of the tumor suppressors 15-PGDH and WNT5A in colon cancer patients. I investigated the underlying mechanisms/signaling triggered by these tumor suppressors in colon cancer cells and whether the re-expression of these tumor suppressors could be an attractive therapeutic strategy for treatment of colon cancer patients.

I found that presence of mast cells in colon cancer tissue was associated with better prognosis of colon cancer patients, and the presence of mast cells in polyps/tumors in a colitis-associated colon cancer mouse model was also beneficial. I found that the tumor suppressor gene 15-PGDH is down-regulated in colon cancer patients as well as in colon cancer cell lines. This down-regulation is often seen in parallel with down-regulated WNT5A, the non-canonical Wnt/β-catenin signaling ligand. I found that down-regulation of both these proteins is associated with poor prognosis for colon cancer patients. My results show that treatment of colon cancer cells with Foxy-5, a WNT5A mimicking peptide, leads to up-regulation of 15-PGDH through JNK/AP-1 pathway. In addition, I also found that the pro-inflammatory mediator LTC₄ via CysLTR2 can induce the expression of 15-PGDH through the JNK pathway, which indicate that LTC₄ via CysLTR2 has an anti-tumor effect.

In conclusion, these findings provide important information for better understanding of the tumor microenvironment and the tumor suppressor genes in colon cancer and might help to identify new therapeutic targets for colon cancer patients.

Key words Colon cancer, Mast cells, 15-PGDH, WNT5A, LTC₄
Role of novel tumor suppressors in colon cancer: Mechanisms and therapeutic opportunities

Lubna Mohammed Mehdawi
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Lund University
Faculty of Medicine
Department of Translational Medicine

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To my parents, my husband, my kids

“And mankind have not been given of knowledge except a little”

Quran, chapter Al-isra, verse no. 85

وَمَا أُوْتِيتُم مِّن الْعِلْمِ إِلَّا قَليلاً
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## Abbreviations

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Full Form</th>
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<tbody>
<tr>
<td>15-PGDH</td>
<td>15-hydroxy prostaglandin dehydrogenase</td>
</tr>
<tr>
<td>5-ASA</td>
<td>5 aminosalicylic acid</td>
</tr>
<tr>
<td>5-FU</td>
<td>5-Fluorouracil</td>
</tr>
<tr>
<td>5-LOX</td>
<td>5-Lipooxygenase</td>
</tr>
<tr>
<td>AA</td>
<td>Arachidonic acid</td>
</tr>
<tr>
<td>AJCC</td>
<td>American Joint Committee on Cancer</td>
</tr>
<tr>
<td>AOM</td>
<td>Azoxymethane</td>
</tr>
<tr>
<td>APC</td>
<td>Adenomatous polyposis coli</td>
</tr>
<tr>
<td>CAC</td>
<td>Colitis associated cancer</td>
</tr>
<tr>
<td>COX-2</td>
<td>Cyclooxygenase-2</td>
</tr>
<tr>
<td>CysLTR1</td>
<td>Cysteinyl leukotriene receptor 1</td>
</tr>
<tr>
<td>CysLTR2</td>
<td>Cysteinyl leukotriene receptor 2</td>
</tr>
<tr>
<td>CysLTs</td>
<td>Cysteinyl leukotrienes</td>
</tr>
<tr>
<td>DSS</td>
<td>Dextran sulphate sodium</td>
</tr>
<tr>
<td>EGFR</td>
<td>Epidermal growth factor receptor</td>
</tr>
<tr>
<td>FAP</td>
<td>Familial adenomatous polyposis</td>
</tr>
<tr>
<td>GALT</td>
<td>Gut associated lymphoid tissue</td>
</tr>
<tr>
<td>GPCRs</td>
<td>G-protein coupled receptors</td>
</tr>
<tr>
<td>GSK-3β</td>
<td>Glycogen synthase kinase 3β</td>
</tr>
<tr>
<td>IBD</td>
<td>Inflammatory bowel disease</td>
</tr>
<tr>
<td>ISC</td>
<td>Intestinal stem cell</td>
</tr>
<tr>
<td>JNK</td>
<td>C-jun N-terminal Kinase</td>
</tr>
<tr>
<td>MCD</td>
<td>Mast cell density</td>
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<tr>
<td>Abbreviation</td>
<td>Description</td>
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<tr>
<td>--------------</td>
<td>-------------</td>
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<tr>
<td>MCS</td>
<td>Mast cells</td>
</tr>
<tr>
<td>Muc-2</td>
<td>Mucin-2</td>
</tr>
<tr>
<td>NSAID</td>
<td>Non steroidal anti-inflammatory drugs</td>
</tr>
<tr>
<td>PCK</td>
<td>Protein kinase C</td>
</tr>
<tr>
<td>PGE$_2$</td>
<td>Prostaglandin E$_2$</td>
</tr>
<tr>
<td>PPAR-α</td>
<td>Peroxisome proliferator activated receptor-α</td>
</tr>
<tr>
<td>SI</td>
<td>Sucrose isomaltase</td>
</tr>
<tr>
<td>SRS</td>
<td>Slow reacting substance</td>
</tr>
<tr>
<td>TAM</td>
<td>Tumor associated macrophages</td>
</tr>
<tr>
<td>TGF-β</td>
<td>Transforming growth factor β</td>
</tr>
<tr>
<td>TMA</td>
<td>Tissue microarray</td>
</tr>
<tr>
<td>TNF-α</td>
<td>Tumor necrosis factor-α</td>
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<tr>
<td>VEGR</td>
<td>Vascular endothelial growth factor</td>
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List of Papers

The following papers are included in this thesis.


II. **Lubna M. Mehdawi**, Chandra Prakash Prasad¹, Roy Ehrnström², Tommy Andersson and Anita Sjölander. Non-canonical WNT5A signaling up-regulates the expression of the tumor suppressor 15-PGDH and induces differentiation of colon cancer cells. 2016. Molecular oncology. Doi: 10.1016.¹

III. **Lubna M. Mehdawi**, Annika Gustafsson, Kent Lundholm, Maria Alvarado-Kristensson and Anita Sjölander. A possible anti-tumor effect of Leukortine C₄ by inducing 15-hydroxyprostaglandin dehydrogenase expression in colon cancer cells. (Manuscript)

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Introduction

Cancer is an ancient disease, with the earliest records dating to ancient Egypt (circa 3000 BC). The Greek physician Hippocrates (460-375 BC) first described the growth of cancer as a crab attaching to the surrounded tissue by its claws. Later, the Roman physician Celsus (25 BC-50 AD), who created the Latin language of medicine, translated the word Crab to Cancer [1]. Cancer became defined as abnormal cell growth, invading surrounding tissues and able to metastasize to other parts of the body. Cancer treatment has improved from surgery in the 19th century to today’s use of adjuvant chemotherapy and radiotherapy. Although progress in cancer treatment is ongoing, it remains a major cause of death worldwide [2]. Various factors, such as chemical carcinogens, ionizing radiation and viruses, can cause non-inherited cancer by inducing genetic damage, leading to the identification of genes that are essential in cancer development. In the 1970s, these genes were classified into two families: proto-oncogenes and tumor suppressor genes. Accumulation of mutations in these genes is believed to cause cellular alterations, and subsequently lead to cancer development [3]. The essential cellular alterations required for neoplastic transformation, referred to as the hallmarks of cancer, include replicative ability, sustained proliferative, evasion of growth suppression and cell death, and induction of angiogenesis and invasion/metastasis [4]. More recently, evasion of immune destruction and reprogramming of energy metabolism have emerged as new hallmarks of cancer. The inflammatory milieu has also been appreciated as one of these hallmarks [5]. Inflammation, particularly chronic inflammation such as in inflammatory bowel disease (IBD), has been established as a connection between inflammatory disease and colorectal cancer [6]. In colon cancer, mutation of oncogenes and tumor suppressor genes leads to activation of several oncogenic pathways and deactivates tumor suppressor pathways [7]. Overexpression of cyclooxygenase-2 (COX-2), which has been shown in the majority of cancers, especially colorectal tumors [8], leads to overproduction of prostaglandin E2, a proinflammatory lipid mediator, which induces proliferation, neovascularization, inhibition of cell death and motility of tumor cells [9]. 15-hydroxyprostaglandin dehydrogenase (15-PGDH) is the catabolic enzyme that leads to degradation of the prostaglandins [10]. It is considered a tumor suppressor gene and is known to be downregulated in colorectal cancer [11].
Here, we investigated the possible mechanisms regulating 15-PGDH in colon cancer cells, and we found that WNT-5A and its mimicking peptide Foxy-5 can induce 15-PGDH expression in colon cancer by activating the JNK/AP-1 pathway and inhibiting the β-catenin pathway. We also found that the proinflammatory mediator LTC₄ could induce the expression of 15-PGDH through CysLTR2 by activating the JNK pathway. However, the presence of mast cells in colon cancer patient material is associated with good prognosis for colon cancer patients.
The intestinal tract and its function

The intestinal tract is one of the most complex and regenerative organs, and it is the largest immunological organ in the human body [12]. It is divided into the small intestine and large intestine. The small intestine is divided into three parts: the duodenum, jejunum and ileum, whereas the large intestine is subdivided into the cecum, ascending, transverse, descending and sigmoid colon (Figure 1).

The intestinal wall is composed of three layers. The outermost layer is the muscular layer, which consists of smooth muscles, the middle layer is a layer of stromal tissue, and the innermost layer is the mucosal layer, which is composed of epithelial cells. Epithelial cells of the intestine are simple, non-ciliated columnar cells, arranged in a layer, that form the luminal surface. Due to the presence of
“gut-friendly” bacteria, the human intestine is normally exposed to a low degree of inflammation [13]. The intestinal epithelium is protected from external pathogens and microorganisms by cells in the lamina propria, which contains many types of immune cells; it is rich in macrophages, lymphoid cells, and dendritic cells, making it the primary location for immune reactions (Figure 2, 3) [14]. Other intestinal defense mechanisms come from the intestine’s own immune system, known as gut-associated lymphoid tissue (GALT). Impairment of this system leads to unbalanced inflammatory responses in the gut and predisposes people to inflammatory bowel disease (IBD) [15]. The absorptive surface of the small intestine is increased because of the presence of the villi, invaginations into the submucosal layer and the crypt of Lieberkun [16]. The epithelial surface of the large intestine is flat, as it lacks villi and contains only crypts (Figure 2).

There are four types of differentiated cells in the intestine. Enterocytes, which are the majority, are absorptive in nature, having brush border microvilli on their apical surface. Goblet cells and Paneth cells secrete mucus and antimicrobial proteins, respectively, providing a substantial physical and biochemical barrier. Enteroendocrine cells secrete hormones such as secretin and pancreozymin. Intestinal epithelial homeostasis is maintained by three mechanisms. First, cell production by the division of the crypt progenitors every 12 h produces 200 cells per crypt. Second, all epithelial cells move continuously upward and reach the top of the villus in approximately 5 days, except for the Paneth cells, which escape the
movement upward. Third, proliferation dictated by the crypt niche is maintained as cells move along the crypt-villus axis [16, 18]. The epithelial cell renewal process within the intestine appears to be totally dependent on a limited number of long-lived intestinal multipotent stem or progenitor cells. Intestinal stem cells (ISC) have the following two properties: the capacity to maintain themselves for long periods of time (self-renewal) and the ability to generate all differentiated cell types, including enterocytes, entero-endocrine, and Paneth cells [19].

Figure 3: Representative intestinal mucosal surface, adapted from Eberl, Gerard, et al., 2005 [20]

Inflammation and cancer

In 1863, Virchow reported the presence of leukocyte infiltration in cancer tissue; he hypothesized that chronic inflammation can lead to cancer. Now, the understanding of the inflammatory microenvironment of cancer tissue supports Virchow’s findings [21]. Tissue injury and inflammation lead to increased cell proliferation; although, it is clear now that proliferation of the cell alone cannot cause cancer. However, DNA damage and sustained cell proliferation in an
inflammatory milieu rich in inflammatory cells and growth factors can promote cancer [22]. The hallmarks of cancer include evading growth suppressors, sustaining proliferative signaling, resisting cell death, enabling explicative immortality, including angiogenesis, and activating invasion and metastasis. In addition, inflammation and genome instability have recently been established as hallmarks of cancer [5] (Figure 4).

Figure 4: Hallmark capability of cancer, adapted from Hanahan, et al., 2011 [5]
Inflammatory bowel disease and colitis associated colon cancer

Inflammatory bowel disease (IBD) defined as chronic intestinal disorder caused by a number of factors that contribute to the development of mucosal inflammation, including environmental and genetic factors as well as a dysregulated immune response [23]. IBD includes Crohn’s disease, which affects any part of the gastrointestinal tract from the mouth to the anus in an irregular pattern, and ulcerative colitis (UC), which affects the rectum and may involve part or all of the colon in a continuous pattern [24]. The inflammation in Crohn's disease is often transmural, whereas it is typically confined to the mucosa in UC [25]. The risk of developing colon cancer is highly increased among these patients, perhaps due to prolonged exposure to inflammation in the bowel. Colitis-associated cancer (CAC) is colon cancer occurs in patients with clinically detectable inflammatory bowel disease [26-29]. The increased risk of developing cancer in IBD patients depends on many factors, such as disease duration and severity and the effectiveness of treatment [30-32]. Treatment of IBD patients with anti-inflammatory drugs such as 5 aminosalicylic acid (5-ASA) has been linked to a decreased risk of developing CAC [33-35], as has a tumor necrosis factor-α (TNF-α) antagonist [36]. In animal studies, treatment with 5-ASA in an AOM/DSS mouse model of CAC shows reduction in dysplastic colon lesions in treated mice [37]

Colon cancer

Colorectal cancer is the second most common cancer in women and the third most common cancer in men worldwide, and the fourth leading cause of cancer-related deaths [38, 39]. Etiological factors of colon cancer include a low fiber and high fat diet and high alcohol intake [40] [41]. Genetic alteration can lead to the activation of oncoproteins as BRAF and KRAS, which are mutated in approximately 50% of CRC [42], inactivation of tumor suppressor genes, such as APC and P53 [43, 44]. The majority (approximately 65-85%) of colon cancer is sporadic; hereditary colon cancer, a genetic disorder and the occurrence of CRC in first- and second-degree relatives, accounts for approximately 10-30% [45, 46]. Familial adenomatous polyposis (FAP) is an autosomal dominant disease characterized by the presence of countless adenomatous polyps throughout the colon and rectum. If not treated, the polyps can progress to cancer at a young age [47, 48]. FAP patients have a germ line mutation in the tumor suppressor gene APC; however, 70% of sporadic colon cancers have somatic mutations in the APC gene [49-51]. APC is a multifunctional protein that participates in many cellular processes, including cell-
cell adhesion, cell migration, and apoptosis; it also regulates β-catenin degradation [52, 53] (Figure 5).

![Figure 5: β-catenin destruction complex, adapted from Li, Vivian, et al., 2012 [54]](image)

**Adenoma-carcinoma sequence**

Adenomas are important precursor lesions for carcinomas. The adenoma-carcinoma sequence describes the stepwise progression from normal to dysplastic epithelium to carcinoma, with the accumulation of specific mutations at specific stages [55]. These mutations affect genes and pathways that are important for the regulation of cell growth and differentiation [56]. APC mutations occur early during the progression from adenoma to carcinoma [57, 58]. KRAS mutations also occur early in the adenoma-carcinoma sequence; however, KRAS mutations are less common in small adenomas. P53, SMAD2, and SMAD4 are tumor suppressors; mutations in these genes can lead to colon cancer progression [55].
Colon cancer classification and staging

In general, cancer staging provides critical information about the extent of the disease, helping in treatment decisions and prognosis determination. In 1932, Dukes proposed a classification of colorectal cancer based on the extent of the disease, which is evaluated by the degree of the tumor infiltration through the wall of the bowel and the presence or absence of lymph node involvement [59]. In 1987, the American Joint Committee on Cancer (AJCC) and the International Union Against Cancer introduced the cancer staging system based on the size and the extent of the tumor (T), number and degree of lymph node involvement (N), and presence or absence of distant metastasis (M) [60] (Table 1). This staging system of CRC was updated in 2009 and is the most common staging system used in clinical practice [61, 62].

Tumor, node, metastasis (TNM) staging of colorectal cancer

**T - primary tumor**

TX - primary tumor cannot be assessed  
T0 - No evidence of primary tumor  
Tis - Carcinoma in situ: intraepithelial or invasion of lamina propria  
T1 - Tumor invades the submucosa  
T2 - Tumor invades the muscularis propria  
T3 - Tumor invades through the muscularis propria into the subserosa or non-peritonealized pericolic or perirectal tissues  
T4 - Tumor directly invades other organs or structures and/or perforated visceral peritoneum

**N - regional lymph nodes**

NX - Regional lymph node cannot be assessed  
N0 - No regional lymph node metastasis  
N1 - Metastasis in one to three regional lymph nodes
N2 - Metastasis in four or more regional lymph nodes

M- distant metastasis

MX - Distant metastasis cannot be assessed

M0 - No distant metastasis

M1 - Distant metastasis

**Table 1. Comparison of TNM and Duckes classification**

<table>
<thead>
<tr>
<th>Stage</th>
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<td>Tis</td>
<td>M0</td>
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<tr>
<td>Stage I</td>
<td>T1, T2</td>
<td>M0</td>
</tr>
<tr>
<td></td>
<td>N0</td>
<td>M0</td>
</tr>
<tr>
<td>Stage II</td>
<td>T3, T4</td>
<td>M0</td>
</tr>
<tr>
<td></td>
<td>N0</td>
<td>M0</td>
</tr>
<tr>
<td>Stage III</td>
<td>T1, T2</td>
<td>M0</td>
</tr>
<tr>
<td></td>
<td>N1 or N2</td>
<td>M0</td>
</tr>
<tr>
<td>Stage IV</td>
<td>Any T</td>
<td>M1</td>
</tr>
<tr>
<td></td>
<td>Any N</td>
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</table>

**Treatment of colorectal cancer**

In general, CRC treatment depends on many factors, such as the age of the patient and disease stage. The four key approaches currently used for CRC treatment are surgery, which is the main approach, chemotherapy, radiotherapy, which is mostly for rectal cancer, and targeted therapies [63]. To reduce symptoms and increase survival in metastatic CRC, chemotherapy and targeted therapies are given [64]. The most commonly used chemotherapy is 5-fluorouracil (5-FU); antibodies targeting EGFR (epithelial growth factor receptor) or VEGF (vascular endothelial growth factor) are commonly used [65]. Colorectal cancer patients with mutant KRAS have no benefit with anti-EGFR therapy, which make it a promising predictive biomarker for personalized treatment [66].
Signaling pathways in colon cancer

Wnt signaling pathway

Wnt proteins are a family of secreted cysteine-rich glycoproteins approximately 40 kDa in size [67]. The name Wnt is derived from a fusion of wingless, which is a *Drosophila* segment polarity gene, and integrated or int-1, the vertebrate homolog [68, 69]. Wnt protein signaling pathways play an important role in many cellular processes, including cell fate, cell polarity, cell motility, primary axis formation and organogenesis; most recently, this pathway has been shown to play a role in stem cell renewal [70, 71].

To induce and regulate intracellular transduction signaling, Wnt proteins bind to cell surface G protein coupled receptors, called frizzled receptors (FZ), in the presence or absence of the co-receptors LRP 5/6, ROR or RYK [72-75]. Wnt malformation can lead to various diseases, including cancer and degenerative diseases [76]. The Wnt signaling pathway can be divided into the canonical signaling pathway (Wnt/β-catenin) and the non-canonical signaling pathways (Wnt/Ca²⁺ signaling, planar cell polarity (PCP)) [74, 77].

In canonical Wnt/β-catenin signaling, in the absence of a Wnt ligand, such as Wnt1 and Wnt3a, the free cytoplasmic β-catenin binds to a destruction complex consisting of Axin, APC (adenomatous polyposis coli), and GSK-3β (glycogen synthase kinase-3β-catenin), leading to β-catenin phosphorylation, ubiquitination and degradation by proteasomes. Thus, the Wnt/β-catenin target genes remain inactive (Figure 6a). When Wnt ligand binds to frizzled receptors, a cascade of events involving Dishevelled (Dvl) leads to the phosphorylation of the co-receptor LRP5/6 and the recruitment of Axin complex to the receptor. These events lead to the inhibition of Axin-mediated β-catenin phosphorylation, whereby β-catenin is stabilized, accumulates in the cytoplasm and translocates to the nucleus where it activates Wnt target genes, such as COX-2, C-myc and Cyclin-D1 (Figure 6b) [78-80].
Non-canonical Wnt signaling (β-catenin independent Wnt signaling) is an umbrella term for all β-catenin independent pathways. In contrast to the canonical Wnts, non-canonical Wnts, such as Wnt-4, Wnt-11 and WNT5A, are non-transforming, and they are involved in processes such as polarized cellular movements [82-84].

The Wnt/Ca$^{2+}$ pathway, a non-canonical Wnt signaling pathway, involves the binding of ligands, such as Wnt5a and Wnt-11, to the FZ receptor, a G-protein coupled receptor, leading to activation of phospholipase C (PCL), which subsequently triggers the intracellular release of calcium ions [85, 86]. Calcium activates protein kinase C (PKC) [87], which regulates cellular processes such as cell adhesion and tissue separation during gastrulation via Cdc42. Intracellular accumulation of Ca$^{2+}$ also activates calmodulin-dependent protein kinase II [88], which in turn can antagonize canonical Wnt signaling by activating TAK1 (TGFβ activated kinase) and NLK (Nemo-like kinase) [89, 90].

The planar cell polarity pathway (PCP) is another non-canonical pathway, first described in Drosophila, that regulates cytoskeletal polarization. The Wnt/jun N-terminal Kinase (JNK) pathway in vertebrates is similar to the PCP pathway in Drosophila. Binding of a non-canonical Wnt ligand, such as Wnt-11, to the FZ receptor activates Dvl, which mediates cytoskeletal rearrangements through activation of small GTPases, such as Rho and Rac [84, 91].
WNT5A

WNT5A is one of the most extensively studied non-canonical Wnts, and it is a representative of β-catenin-independent, non-canonical Wnt signaling. WNT5A plays essential roles in organ development, cell proliferation, tissue orientation and cell migration [92]. In addition to signaling through non-canonical pathways [82, 93], WNT5A can also activate or inhibit canonical Wnt signaling depending on the presence or absence of certain receptors/co-receptors [89, 94, 95]. Recent studies have shown that any misregulation of WNT5A signaling results in cancerous growth of certain tissues, but it is debatable whether WNT5A has a tumor suppressor or tumor promoting effect; this depends on the type of tissue and additional factors, regulators and receptors available [82, 93]. In many studies, Wnt5a has a tumor suppressor effect, as it is downregulated in many cancers, including colorectal cancer [96], invasive ductal breast carcinomas [97, 98], neuroblastoma [99], hepatocellular carcinoma and leukemia [100, 101]. In colon cancer, loss of WNT5A is associated with poor outcomes in stage Dukes B colon cancer patients [96].

Eicosanoids

Eicosanoids are a class of lipid mediators derived from 20-carbon polyunsaturated fatty acids (the Greek word eicosa means 20). The main eicosanoid precursor in the mammalian system is arachidonic acid (AA), which belongs to the ω-6 family of polyunsaturated fatty acids [102]. Three main enzymatic pathways metabolize arachidonic acid. Eicosanoids and prostanoids (prostaglandins and thromboxanes) are synthesized via the COX pathway, leukotrienes and hydroxyl eicosa-tetraenoic acids (HETEs) are generated via the lipooxygenase (LOX) pathway, and epoxides are generated via the cytochrome P-450 epoxygenase pathway [103] (Figure 7).

In the 1930s, the vasodepressor and smooth muscle-stimulating activities of prostaglandins were discovered, although the origin and the structures of these compounds were not yet reported. In 1982, Samuelson and Bergström demonstrated the 5-LOX pathway and the discovery of cysteinyl leukotrienes as the slow-reacting substance (SRS) of anaphylaxis. These mediators were responsible for the contraction of the smooth muscles of the guinea pig lung after treatment with cobra venom. They were also found to be mediators in asthma and other immediate hypersensitivity reactions [104, 105]. In addition to their role in acute systemic inflammation [103], the role of eicosanoids in the maintenance of the hemostasis of the intestine [106] and their dysregulated functions in
pathological conditions, such as chronic inflammation and cancer, are well established [107, 108].

![Diagram of Arachidonic acid metabolism and cysteinyi leukotriene signaling](Image)

**Figure 7:** Arachidonic acid metabolism and cysteinyi leukotriene signaling. Adapted from Savari et al., 2014 [109]

**The COX pathway**

Cyclooxygenase (COX) is the key enzyme required to convert AA to prostanoid. The prostanoid-synthesizing enzyme COX exists in two isoforms, COX-1 and COX-2.

COX-1, which is expressed by most cells, is considered to have a housekeeping function in cells via the production of homeostatic prostanoids. COX-2, which is induced by stress and cytokines, is the source of prostanoid formation in
inflammation and cancer [110]. Prostanoids include prostaglandins and thromboxane A₂. In addition to their role in inflammation, prostaglandins are involved in many physiological functions, including blood clotting, bone metabolism, wound healing kidney function and immune response [111]. Prostaglandins include PGE₂, PGF₂α, PGD₂ and PGI₂, which mediate their action via the activation of receptors, including EP1-4 for PGE₂, prostaglandin receptor F₂α for PGF₂α, DP₁ and DP₂ for PGD₂, and prostaglandin I₂ (IP receptor) for PGI₂ [112]. Thromboxane A₂ is an activating and recruiting mediator for platelets, mediating its action through the specific G-protein coupled receptor TBXA2R [113].

Elevated levels of COX-2 have been identified in colon [114, 115], breast [116], lung [117], pancreatic, esophageal and ovarian cancers [118-120]. PGE₂ is the most common prostaglandin found in many human malignancies, including colon and breast cancer, and it is associated with poor prognosis [107]. COX-2 produces PGE₂, the only prostanoid demonstrated to induce proliferation, angiogenesis, cell death inhibition and motility of tumor cells [9, 121], and more recently, to expand the number of colon cancer stem cells [122]. 15-Hydroxyprostaglandin dehydrogenase (15-PGDH) is the enzyme responsible for the degradation of PGE₂, converting it into its inactive metabolites [123]. 15-PGDH is highly expressed in normal colon mucosa but is lost in many CRCs [124], and its decreased expression has also been implicated in many other cancers, such as lung, bladder, pancreatic and gastric cancer [125-128]. Deletion of the 15-PGDH gene increases colonic PGE₂ levels and enhances tumorigenesis in vivo [124], while treatment of colon cancer cells with indomethacin (NSAID) decreases the expression of COX-2 and induces 15-PGDH up-regulation in colon cancer cells [129].

The 5-Lipoxygenase pathway (5-LOX)

Leukotrienes (LTs)

In contrast to prostaglandins, which are produced by most cells, LTs are mainly produced by leukocytes, such as mast cells, macrophages, basophils and eosinophils. The term leuko- is given according to the cell type (leukocytes) where they were originally identified, and triene refers to the presence of three conjugated double bonds [130]. The 5-LOX pathway is involved in the biosynthesis of leukotrienes, it interacts with 5-LOX activating protein (FLAP) converting AA to LTA₄. Thereafter, LTA₄ is metabolized to biologically active LTB₄ or to cysteiny1 leukotrienes (CysLTs): LTC₄, LTD₄ and LTE₄ [107, 109] (Figure 7). In addition to its role in inflammatory diseases such as asthma [131],
LTB₄ has chemotactic activity and pro-tumorigenic effects in breast cancer and melanoma [132, 133].

Leukotrienes, particularly, CysLTs, mediate their actions through the following two cysteinyi leukotriene receptors (CysLTRs): CysLTR1, which is sensitive to classical antagonists, and CysLTR2, which is not inhibited by classical antagonists. Both receptors belong to the G-protein coupled receptor (GPCR) family [134, 135]. Binding of CysLT ligand to the receptor induces conformational changes leading to G-protein activation, converting GDP to GTP; GTP hydrolysis and many intracellular events occur, such as increased concentrations of Ca²⁺ in the cytosol [135]. CysLTR1 is the high affinity CysLT receptor and has a higher affinity for LTD₄ than LTC₄ [136], whereas CysLTR2 has a lower overall affinity but an equal affinity for LTD₄ and LTC₄ [137]. The link between alteration of CysLTRs and cancer has been documented. High expression of CysLTR1 has been reported in many cancers, including the brain, bladder, prostate, breast, and colon [138-141]. High expression of CysLTR1 is associated with poor prognosis of breast and CRC patients. Patients expressing low CysLTR1 and high CysLTR2 have a good prognosis, which is linked to higher survival of CRC patients [142]. CysLTR1 has been shown to induce the accumulation of free β-catenin through LTD₄, thereby inducing proliferation and migration by activating target genes associated with carcinogenesis, such as CyclinD1, C-myc and COX-2 [143]. Moreover, we found that LTD₄ and PGE₂ can induce the promotion of cancer stem cells in both colon cancer cells and xenograft mouse models [144, 145]. In contrast, a CysLTR1 antagonist can prevent the growth of colon cancer xenograft via reducing proliferation and inducing apoptosis of tumor cells [146]. Cysltr1⁻/⁻ mice have significantly reduced tumor formation compared to wild type mice, which indicates the role of CysLTR1 in tumorigenesis [147]. However, LTC₄ acting through CysLTR2 does not induce proliferation of colon cancer cells; it is involved in the terminal differentiation of colon cancer cells, implicating the anti-tumorigenic activity of CysLTR2 compared to CysLTR1 [148].

Inflammatory cells and colorectal cancer

Colorectal cancers, like other solid tumors, are infiltrated by different types of immune cells. Innate immune cells, such as mast cells, macrophages, natural killer cells, neutrophils and dendritic cells, have been found in these tumors [149]. Adaptive immune system cells, such as T-lymphocytes, are also detected in colorectal tumors, where they are either pro- or antitumorigenic. T cells are important for inflammation, cancer development and progression, and also for
anticancer immunity [150-152]. Macrophages, especially tumor-associated macrophages (TAM), are also found in solid tumors and are believed to have an M2-like phenotype [153]. In colon cancer, we found that TAMs promote colon cancer cell migration and subsequent metastasis [154]. In my thesis, I have focused on the role of mast cells in colon cancer.

**Mast cells (MCs)**

Mast cells (MCs) were first discovered by the German scientist Ehrlich in 1878, who observed connective tissue cells that appeared “well fed”, referring to the presence of filled secretory granules that he called “Mastzellen” [155]. However, Kitamura and co-workers reported that MCs are long-lived, hematopoietic cells derived from pluripotent precursors of the bone marrow [156, 157]. MCs are divided into mucosal and connective tissue mast cells. They normally leave the bone marrow as immature cells that circulate through vascular system as immature progenitors; they complete their development either within mucosal or connective tissues [157]. MC granules contain different types of mediators, such as histamine, serotonin, tryptase, chymase and tumor necrosis factor (TNF-α) [158]. In humans, MCs have been categorized into two groups depending on their protease content; one mainly contains tryptase or (more rarely) chymase, and the other group has both proteases. In mice, MCs are divided into connective tissue mast cells, which contain heparin, histamine, tryptase and chymase in their granules, and mucosal mast cells, which contain chymase and small amounts of histamine [159] (Figure 8).

In general, MCs are best known for their involvement in allergic reactions. When mast cells are activated, either through recognition receptors or associated with IgE receptor, they degranulate and release their contents into the microenvironment [160] (Figure 8). Activating MCs can also produce inflammatory mediators de novo, including cysteinyl leukotrienes (LTC₄, LTD₄, and LTE₄) [161]. In addition to their role in allergy and anaphylaxis, MCs also have a role in tissue remodeling, angiogenesis and immune modulation [162].
The role of MCs in cancer is debatable, including whether they have tumor suppressor effects or tumor promoting effects [163]. MCs are long-lived cells; therefore, they can influence the microenvironment for a long time. Many studies have reported the correlation of the presence of MCs and tumor development with poor patient outcome [164]. High MC density (MCD) together with angiogenesis was associated with poor prognosis in patients with colon [165-167], lung [168] and pancreatic cancer [169]. However, many other studies have shown a protective role of MCs in human cancers. A multivariate analysis of colorectal cancer patients, including Dukes stage, sex, age, perioperative blood transfusion, tumor location, and counts of MCs, predicted good survival [163]. MCs activate nuclear peroxisome proliferator-activated receptor-γ (PPAR-γ), which is associated with better clinical outcome in colon cancer [170]. Low MC density is associated with worse prognosis in non-small lung cancer patients [171, 172]. A positive correlation between MC density and better clinical outcomes in prostate cancer has been documented [173]. In our study, we have also shown that high MCD is associated with good prognosis in colon cancer patients [174].
Aim

The general aim of the thesis is to evaluate the clinical significance of various tumor suppressors in colon cancer patients. Additionally, we investigated the underlying mechanisms or signaling triggered by these tumor suppressors in colon cancer cells, and whether the re-expression of these tumor suppressors could be an attractive therapeutic strategy for the treatment of colon cancer patients.

1. To explore the relevance of mast cells for the overall survival of colon cancer patients and a colitis-associated colon cancer mouse model
2. To investigate the effect of WNT-5A and its mimicking peptide Foxy-5 on 15-PGDH expression in colon cancer
3. To investigate the effect of LTC4, through its receptor CysLT2, on 15-PGDH expression in colon cancer cell lines and colon cancer tissue
Methodology

Cell lines

All cell lines used in my projects were from ATCC. HT-29 colon cancer cells (ATCC# HTB-38™), grown in McCoy’s 5A medium with glutamine, have a wild-type \textit{KRAS} gene and mutant carboxy-truncated \textit{APC} gene. Caco-2 colon cancer cells (ATCC# HTB-37™) have a wild-type \textit{KRAS} gene, wild-type \textit{BRAF} gene and mutated \textit{APC} gene. MDA-MB-468 (ATCC# HTB-132) breast cancer cells, grown in DMEM, have mutated \textit{P53} and \textit{BR1} genes. The media was supplemented with 10% fetal bovine serum and 100 μg/ml penicillin/streptomycin. Cells were maintained at 37°C in a humidified atmosphere containing 5% CO\textsubscript{2}. The cells were routinely screened for the absence of mycoplasma contamination.

Western blotting and cell fractionation

After washing the cells twice with ice-cold PBS, whole cell lysates were prepared with lysing in lysis buffer. The resulting lysates were homogenized by 10 passages through a syringe and then centrifuged at 10,000 × g for 10 min at 4°C. The supernatant was considered to represent the whole cell lysate. For nuclear fractions, a nuclear extraction kit from Thermo Scientific was used according to the manufacturer’s instructions. Laemmlle buffer (4X) was added to the samples. The samples were adjusted to contain equal amounts of protein and boiled for 10 min in sample buffer (0.5 M Tris-base, 10% SDS, glycerol, bromophenol blue, 15 mg/ml DTT) before being loaded onto 10% SDS polyacrylamide gels and analyzed by electrophoresis (SDS-PAGE). After separation, the proteins were transferred to a PVDF membrane (Bio-Rad, Hercules, CA, USA). The membranes were blocked for 1 h with 3% BSA/PBS at room temperature, followed by incubation with a primary antibody overnight at 4°C. The membranes were washed extensively and incubated with the corresponding secondary antibody for 1 h at RT. The membranes were washed and incubated with Immobilon™ Western Chemiluminescent HRP Substrate (Immobilon™ Western, Merck Millipore, Billerica, MA, USA), and the proteins were detected using the Bio-Rad ChemiDoc...
XRS+ system. The densitometric analyses were conducted using Image Lab 3.0 software.

Patients

All patients underwent surgery for colorectal cancer at Malmö hospital in 1990. The tumor material was retrospectively collected, and no matching or stratification was performed. All patients had a pathologically confirmed diagnosis of adenocarcinoma before surgery. All tumors with available slides or paraffin blocks were histologically re-evaluated on hematoxylin and eosin (H&E) stained slides. Clinical and histopathological data were retrieved from patient charts and pathology records. Tumors were staged according to TNM classification for malignant tumors, and tumor grade was determined as high, medium or low. The follow up started at the time of diagnosis and ended at death, emigration or December 31, 2000. The study was approved by the ethical committee of Lund University. In project II, tumor tissues and corresponding mucosa from 23 colon cancer patients were from the Department of Clinical Science, Sahlgrenska Academy, Gothenburg, Sweden. Tumors and large bowel tissue samples were collected down to the serosa level, kept fresh frozen in liquid nitrogen, and stored at -80°C until analysis. A certified pathologist staged all tumors. Tumor samples contained approximately 70-80% tumor cells based on visual inspection [175].

Tumor tissue microarray (TMA) and immunohistochemistry

From the tumor tissue, two to three 1.5 mm tissue cores were placed in a new paraffin block using an automated Beecher Micro-Arrayer (Beecher Instruments, Sun Prairie, WI, USA). The normal areas were chosen from the distal parts of the safe margins. Immunohistochemical staining was performed as previously described [142]. Briefly, the formalin-fixed and paraffin-embedded sections were cut into 1 μm sections, dried, de-paraffinized and rehydrated, then stained with H&E for re-evaluation by a pathologist (RE). Tumor sections were then stained for 15-PGDH or WNT5A, as previously described. All immunohistochemical procedures were performed using a Dako automatic slide stainer (Dako) according to the manufacturer’s instructions.
Real-time qPCR

Total RNA from the different cell samples was isolated using Qiagen RNeasy Plus Mini Kits (Qiagen GmbH, Hilden, Germany). The cDNA synthesis was performed using RevertAid H Minus M-MuLV reverse transcriptase (ThermoFisher Scientific, USA). The following primers were used: HPRT1 (HS99999909), HPGD (15-PGDH) HS00168359_m1), SI (SI, HS00356112_m1), MUC2 (mucin-2, HS00159374_m1) and CCND1 (cyclin D1, HS00765553_m1). Amplifications were performed in an Mx3005P system (Agilent Technologies, Inc., CA, USA). The reactions were normalized to the housekeeping gene HPRT1 and analyzed with MxPro qPCR software (Agilent Technologies, Santa Clara, CA, USA).

Luciferase assay

A Dual-Luciferase Reporter Assay System (Promega, Madison, WI, USA) was used. 15-PGDH promoter plasmids [141] (a gift from Professor Birgit Gellersen, University of Hamburg, Germany) at a final concentration of 1 μg/ml, together with a control Renilla luciferase reporter plasmid of 50 ng/ml, were used for the transfections. The DNA plasmids were allowed to form complexes with PolyFect Transfection Reagent (Qiagen) (ratio 4:1) in Opti-MEM (Gibco/Thermo Fisher Scientific, Waltham, MA, USA), and the cells were treated with the DNA-PolyFect mixture at 37°C for 24 h. The medium was changed to a serum-containing medium, and the cells were allowed to recover for 24 h. Thereafter, the cells were incubated for 2 h with a serum-free medium before treatments. The cells were washed twice with PBS and lysed by the addition of the passive lysis buffer from the Dual-Luciferase Reporter Assay System. The lysed samples were cleared by centrifugation at 1000 x g for 5 min according to the manufacturer’s instructions. Firefly and the control Renilla luminescence were measured on a MiniLumat LB 9506 luminometer (Berthold Technologies GmbH, Dusseldorf, Germany) according to the protocol, and the ratio was calculated. Triplicate samples were prepared in every set of experiments.

Immunofluorescence

In project II, cells were seeded to grow on coverslips for three days, washed with PBS, and fixed in 4% paraformaldehyde for 15 min. Next, cells were washed with
PBS, permeabilized in 0.1% Triton X-100 in PBS for 5 min, and washed six times in PBS. Cells were thereafter incubated at RT in 3% goat serum in PBS for 45 min and washed, followed by 1 h incubation with anti-β-catenin antibody (1:500) in 1% goat serum/PBS. Cells were then stained with DAPI for 3 min (1:1000). The coverslips were washed and mounted in Zeiss fluorescent mounting medium. Fluorescent images were taken using the Zeiss LMS 700 confocal microscope (Carl Zeiss Microscopy GmbH, Jena, Germany). In project III, cells were cultured on coverslips for 3 days. Then, cells were fixed and permeabilized in methanol/acetone (1:1; -20°C for 4 min). Cells were washed with PBS and incubated for 30 min in PBS staining buffer (PBS, 1% FCS, 0.5% BSA) to prevent non-specific antibody binding. Cells were incubated 1 h with primary antibody, washed, and incubated 1 h with Alexa 480 or Alexa 568 secondary antibody (Invitrogen). Cover slips were washed and mounted on glass slides with a fluorescent mounting medium. Fluorescence images were captured using a Zeiss LSM 700 confocal microscope (Carl Zeiss).

Colitis-associated colon cancer (CAC) mouse model

The chemically induced CAC model was generated using 6- to 8-week-old female mice during a 90-day period (n=5 mice/genotype). The CAC model was generated through an intraperitoneal injection of the carcinogen azoxymethane (10 mg/kg body weight), followed by three cycles of the irritant dextran sodium sulfate in the drinking water, provided ad libitum, for five days. Mice with a targeted Cysltr1 gene disruption (Cysltr1<sup>-/-</sup>) on the C57BL/6N background were a kind gift from Professor Frank Austen (Harvard Medical School, Brigham and Women’s Hospital, Boston, MA, USA) [176]. Cysltr1<sup>-/-</sup> mice and their control littermates were bred and maintained at Lund University Animal Facility in Malmö according to ethical permit M262-12, which was approved by the Regional Ethical Committee for Animal Research at Lund University, Sweden. The breeding colony was genotyped for the presence of the Cysltr1 allele by PCR assays using the following three primers: Cysltr1<sup>sense</sup>: 5’-AAAACAATGACGTGCACTATAAAG-3’, Cysltr1<sup>antisense</sup>: 5’-AATCATGTATACTTGGAAGAAGGCTGA-3’, and Neo<sup>antisense</sup>: 5’-ATCTTTGTCAATGGCCGATCCCAT-3’, generating a product of 284 bp for the wild type and 333 bp for the null Cysltr1 allele.

All mouse colons were first rinsed with a physiological salt solution and subsequently fixed in formalin using the “swiss roll” method, followed by embedding in paraffin.
MDA-MB-468 cell xenografts

The generation of MDA-MB-468 breast cancer cells stably expressing a WNT5A vector or an empty control vector was previously described by Prasad et al [177]. Four- to five-week-old female athymic nude mice were purchased from Harlan Laboratories Inc. (Boxmeer, Netherlands). The cells (2 x 10^6 cells in 100 µl of serum-free medium) were inoculated subcutaneously into both flanks of the mice. The animals were sacrificed 24 days after they had been inoculated, and the tumors were removed, measured and weighed. The primary tumors were dissected, fixed in 4% paraformaldehyde and embedded in paraffin. Sections (4 µm) were cut and subsequently stained with the 15-PGDH-specific antibody. The stained tissue sections were scanned with the Scanscope CS System (Aperio, Bristol, UK) and analyzed with the Aperio Image Scope software. These animal experiments were performed at Pharmatest Services Ltd (Turku, Finland) under the ethical permission no. 3257/04.10.07/2014.

Statistical analysis

GraphPad Prism software 5.0 (San Diego, CA, USA) was used for the statistical analyses. The differences between groups of data were considered statistically significant if P ≤ 0.05 by the two-tailed Student’s t-test. All means were calculated based on data from at least three different experiments. Overall survival (OS) was the primary endpoint. Kaplan-Meier estimates were used to illustrate survival, and log rank test was used to assess for the equality of survival curves. Hazard ratios were estimated using a Cox proportional hazards model for OS in uni- and multivariate analyses. SPSS version 19.0 (SPSS, IBM, Armonk, NY, USA) was used for the statistical analyses of all the immunostaining data.
Results and discussion

Paper I

Chronic inflammation is a risk factor for colon cancer. The key players in the inflammation–neoplasia relationship are the immune cells and the products released by the innate immune system. Chronic inflammation is dominated by the presence of cells, macrophages and MCs at the site of inflammation [178]. MCs induce T-lymphocyte activation and recruitment through the production of chemotactic cytokines such as C-C motif chemokine ligand 5 (CCL5), which bind to G-protein coupled receptor C-C chemokine receptor type 5 (CCR5). MCs, eosinophils and basophils also produce cytokines such as interleukin 4 (IL-4), which induce the apoptosis of tumor cells. Macrophages and MCs secrete tumor necrosis factor α (TNF-α), which induces both tumor cell death and leukocyte infiltration [179]. MCs are a primary source of intratumoral prostaglandin D2 (PGD2), which limits tumor growth through the suppression of angiogenesis and proliferation and promotes apoptosis [107]. Although MCs have a clear role in inflammation and allergic reaction, their role in cancer is not yet clear. Several studies have shown that MCs can promote tumor development and progression. MCs have been shown to play a role in primary breast cancer angiogenesis [180] and lung cancer angiogenesis [181]. Furthermore, other studies have shown that high MCD was correlated with low overall survival of colon cancer patients [165, 167]. In contrast, still other studies have reported the correlation between high MCD and longer survival of colon cancer patients [163, 166, 182]. The aim of my study was to explore the relevance of mast cells in CRC.

In our study, tumor samples from 72 colon cancer patients were evaluated for the presence of MCs by immunohistochemistry in normal and cancer tissues using anti-tryptase and anti-chymase antibodies. MCD was evaluated in normal and cancer tissue. We found that in normal tissue, the mean MCD±SEM was 15±1.6 for tryptase-positive MCs and 101±11.5 for chymase-positive MCs; in cancer tissue, the mean MCD was 8±1.1 tryptase-positive and 52±7.4 chymase-positive MCs. For both tryptase- and chymase-positive cells, the MCD was significantly higher in normal tissue compared to cancer tissue (P=<<0.001). For survival analysis, we found that colon cancer patients with high MCD have significantly longer survival (n=70, P=0.031). We observed the same result in patients without
distant metastasis at the time of diagnosis (n=56, P=0.004) and in a subgroup of patients younger than 75 years (n=33, P=0.015). We preformed univariate and multivariate Cox regression analysis. In univariate analysis, we found that high MCD in cancer tissue significantly reduced the risk of death by 46% (HR 0.539; 95% CI 0.302-0.961); in multivariate analysis, our data showed that MCD was independently associated with longer overall survival in colon cancer patients (HR 0.380; 95% CI 0.202-0.713). Previously, we have shown that patients with few M2 macrophages in their cancer tissues have a better prognosis[154]; here, we found that patients having low M2 and high MCD in cancer tissue had statistically significantly longer survival compared to patients with high M2 and low MCD.

We have previously shown that low CysLTR1 expression in CRC patients is associated with better overall survival [142]. In this study, we investigated the correlation between MCs and cytoplasmic CysLTR1. We found a significant negative correlation between the number of MCs in cancer tissue and CysLTR1 expression (Pearson’s correlation, r = 0.3; P = 0.028). Based on this result, we investigated the presence of MCs in normal and tumor tissue of wild type and Cysltr1-/- mice in a colitis associated colon cancer (CAC) model, a model known to have fewer polyps/tumors [147]. Our result showed that Cysltr1-/- mice had a significantly higher number of tryptase-positive MCs in polyps/tumors compared with wild-type mice (P = < 0.05). Taken together, these results indicate that mast cells have an anti-tumor effect, possibly by creating an anti-tumor environment in colon cancer [174].

**Paper II**

PGE$_2$ plays an important role in cancer, especially colorectal cancer, where its levels are elevated [183], and can influence many of the hallmarks of cancer [5, 107] (see figure 4). This elevated level of PGE$_2$ is due to the up-regulation of COX-2, the enzyme responsible for converting arachidonic acid into PGH2, which is subsequently converted to prostaglandins via prostaglandin synthase [184]. In normal tissue, PGE$_2$ is degraded by 15-hydroxyprostaglandin dehydrogenase (15-PGDH) by converting it into a biologically less active metabolite. 15-PGDH is downregulated in colon cancer, which suggests that 15-PGDH could be a tumor suppressor gene [185]. Elevated levels of PGE$_2$ were previously attributed to COX-2 up-regulation, but more recently to the down-regulation of 15-PGDH. The WNT signaling pathway plays an important role in the homeostasis of the intestinal epithelium, and it is activated in 90% of sporadic CRC [186]. Canonical Wnt/β-catenin signaling has been shown to interact with the 15-PGDH promoter and repress 15-PGDH expression in colon cancer [123]. The aim of the present study
was to investigate the effect of non-canonical Wnt/β-catenin signaling, via WNT5A, on the expression of 15-PGDH.

We have investigated the expression of both 15-PGDH and WNT5A using tissue microarrays (TMA) with human colon cancer tissues and matched normal tissues. The paired samples were stained with both 15-PGDH and WNT5A specific antibodies. Both proteins were expressed at high levels in normal mucosa and significantly downregulated in cancer tissues. For further analysis, tumor samples from 84 patients with primary colon cancer were used to generate TMAs, which were stained with 15-PGDH for immunohistochemical evaluation. Out of 84 patients, 74 (88%) had low expression and 10 (12%) had high expression of 15-PGDH. Of the 84 patient samples analyzed for 15-PGDH, 66 were analyzed for the expression of WNT5A, and our results showed 26 (39%) of these samples have no or low expression of WNT5A, based on the positive correlation found between 15-PGDH and WNT-5A (Pearson correlation r = 0.3; P = 0.027). Kaplan-Meier survival analysis shows that patients with high levels of 15-PGDH tended to have better survival compared to those expressing low levels (P = 0.08). For WNT5A, patients expressing high levels of WNT5A had statistically significantly better survival compared to the patients with low expression levels (P = 0.002). All 26 patients with low expression WNT5A levels also had low expression of 15-PGDH, indicating that loss of WNT5A protein expression in CRC correlated with loss of 15-PGDH protein expression. Moreover, patients expressing low levels of both proteins had reduced overall survival compared to patients with high levels of both proteins (P = 0.005).

We next investigated whether WNT5A signaling can lead to re-expression of 15-PGDH protein in colon cancer cells (HT-29 and Caco-2) that are negative for WNT5A. After treating cells with either recombinant WNT5A or its mimicking peptide Foxy-5, the cells showed significantly increased expression of 15-PGDH at both the protein and mRNA levels. To investigate the signaling pathway by which WNT5A mediates 15-PGDH expression, we analyzed the c-jun N-terminal kinase (JNK) pathway, a well-studied target of WNT5A signaling. We observed an up-regulation of phospho-JNK after stimulating the cells with Foxy-5. We also observed down-regulation of 15-PGDH promoter activity in the presence of a JNK inhibitor after stimulation with both rWNT5A or Foxy-5 in cells transfected with the 15-PGDH promoter. Furthermore, we also found that JNK inhibition down-regulates 15-PGDH mRNA levels. These results indicate that WNT5A mediates 15-PGDH expression through the JNK pathway. We further investigated whether WNT5A suppressed canonical Wnt/β-catenin signaling, and whether that suppression could affect the expression of 15-PGDH. We found that rWNT5A or Foxy-5 reduced the non-phospho-active form of β-catenin and induced the phosphorylation of β-catenin, which reduced the total level of β-catenin in both HT-29 and Caco-2 colon cancer cell lines. Moreover, we investigated whether
inhibition of canonical Wnt/\(\beta\)-catenin signaling could influence 15-PGDH expression. To this end, we used XAV-939, an inhibitor of Wnt/\(\beta\)-catenin signaling that stabilizes axin, leading to increased degradation of \(\beta\)-catenin. Our results showed that XAV-939 can induce 15-PGDH expression and reduce non-phospho-active \(\beta\)-catenin and the total level of \(\beta\)-catenin to a similar level as rWNT5A. We also observed that in the presence of rWNT5A and XAV-939, the increase of 15-PGDH expression and reduction of active form or total \(\beta\)-catenin were more pronounced. Our finding showed that non-canonical WNT5A can inhibit canonical Wnt/\(\beta\)-catenin signaling, and as a result, an up-regulation of 15-PGDH was observed. In breast cancer cells, we noticed the up-regulation of 15-PGDH after stimulating MDA-MB-468 cells with rWNT5A or Foxy-5. In mice, we injected WNT5A-transfected MDA-MB-468 cells into mice flanks; as a control, we injected MDA-MB-468 cells transfected with empty vector (control). We found a significant reduction in tumor growth and tumor volume in mice with WNT5A-transfected cells compared to the control mice. The tumors were immunostained with anti-15-PGDH antibody, and we observed a significant increase in 15-PGDH intensity in tumors with WNT5A-transfected cells compared to the control. These results confirm our previous finding that WNT5A induces the expression of 15-PGDH in both colon and breast cancer cells in vivo and in vitro. Cancer cells are proliferative cells in their nature, and they are poorly differentiated. To investigate whether our finding could affect the differentiation of colon cancer cells, we studied the effect of WNT5A on differentiation by measuring the level of two differentiation markers, mucin-2 (Muc-2) and sucrose isomaltase (SI). A significant up-regulation in both markers was observed in both cell lines after treatment with WNT5A and Foxy-5. This up-regulation could be inhibited in the presence of both a JNK inhibitor and the GSK-3\(\beta\) inhibitor (CHIR-99021), indicating that WNT5A could induce differentiation in colon cancer cells via activating the JNK pathway and inhibiting Wnt/\(\beta\)-catenin signaling. We believe that this could be due to the ability of WNT5A to induce 15-PGDH expression.

In conclusion, our findings reveal a novel interaction between two tumor suppressor proteins, 15-PGDH and WNT5A, through two different pathways. The benefit of this finding is that the level of 15-PGDH in tumor tissue could be used as biomarker in future clinical studies with WNT5A-mimicking peptide Foxy-5 [187].
Paper III

We previously showed that high expression of CysLTR2 in colon cancer patients is associated with better prognosis, which indicates that CysLTR2 has an anti-tumor effect [148]. Based on these data, the aim of this study was to investigate the effect of LTC₄, via CysLTR2, on the tumor suppressor 15-PGDH. We found that tumor tissue from 23 colon cancer patients expressed low levels of both CysLTR2 and 15-PGDH. Next, we examined whether LTC₄ through CysLTR2 can induce the expression of 15-PGDH. Our results showed that after treatment with LTC₄, a significant induction of 15-PGDH expression was observed in HT-29 and Caco-2 cells. This up-regulation was inhibited by pretreatment with CysLTR2 antagonist. Our results showed that LTC₄ via CysLTR2 could induce the expression of 15-PGDH. The 15-PGDH promoter has binding sites for AP-1 [188]. Therefore, we elucidated whether LTC₄ can induce AP-1 activity, and which pathway is involved in this process. We transfected HT-29 cells with a 15-PGDH promoter construct with different binding sites for AP-1. The cells were pretreated with different pathway inhibitors, including MAPK inhibitor, PI3K inhibitor, JNK1 inhibitor and CysLTR2 antagonist. We found that LTC₄ induced 15-PGDH promoter activation, which was inhibited by the CysLTR2 antagonist. We also found that except for the JNK1 inhibitor, none of the other inhibitors could inhibit the promoter activity of 15-PGDH. Furthermore, phosphorylation of JNK by LTC₄ was detected by Western blot in HT-29 cells. We speculated that LTC₄, via CysLTR2, acted through JNK/AP-1 pathway to induce the expression of 15-PGDH in colon cancer cells.

Independently, both CysLTR2 and 15-PGDH were previously shown to inhibit the proliferation of colon cancer cells [148, 189], but their role in inducing the differentiation of colon cancer cells is not clear. Here, we investigated their effect on differentiation by studying two differentiation markers, sucrase isomaltase (SI) and mucin-2 (Muc-2). After stimulating the cells with LTC₄, we found significant up-regulation of Muc-2 mRNA levels. This up-regulation was inhibited in the presence of both the CysLTR2 antagonist and the JNK1 inhibitor. Furthermore, we confirmed the results by confocal microscopy. We also observed a significant up-regulation of SI at the protein and mRNA levels. These results indicated that LTC₄ signaling through a CysLTR2/JNK dependent pathway could induce differentiation of colon cancer cells.
Conclusions

The conclusions of my thesis are as follows:

- The presence of mast cells in the tumor microenvironment is associated with longer survival of CRC patients by creating an antitumor microenvironment.

- Re-expression of the tumor suppressor gene 15-PGDH by WNT5A or its mimicking peptide Foxy-5 leads to better and longer survival of CRC patients.

- CysLTR2 shows antitumor effects by re-expressing 15-PGDH in combination with a more differentiated colon cancer cell.

- These findings provide an important clue to better understand the tumor microenvironment and the tumor suppressor genes in colon cancer, and can help to identify new therapeutic targets for colon cancer patients.
Cancer is uncontrolled cell growth with the ability to invade or spread to other parts of the body. colorectal cancer (CRC) is cancer of colon and rectum. CRC is the third most common cancer and the fourth leading cause of cancer-related deaths worldwide. CRC is a multifactorial disease, with environmental causes such as excessive consumption of alcohol, diets rich in red meat or low in fiber, and low physical activity, as well as genetic causes, which involve either the activation of oncogenes, the genes that have the potential to cause cancer, or the inhibition of the tumor suppressor genes, the genes that protect the cell from cancer. CRC is closely linked to chronic inflammation. Patients with inflammatory bowel disease (IBD) have an increased risk of developing CRC. Many studies have shown that host immune cells that have infiltrated the tumor can enforce a massive inflammatory response that can kill tumor cells and inhibit growth of the tumor. The milieu which contains cellular infiltration surrounding the tumor is called the tumor microenvironment.

The aim of this thesis was to evaluate the clinical significance of the tumor microenvironment, including mast cells (MCs), and tumor suppressor genes, including 15-hydroxyprostaglandin dehydrogenase (15-PGDH) and WNT5A, in colon cancer, and whether the re-expression of these tumor suppressors could be an attractive therapeutic strategy for the treatment of colon cancer patients.

Mast cells (MCs) are immune cells derived from bone marrow. They are well known in allergic reactions, but their role in cancers, including colon cancer, is still under debate. Past studies have shown that MCs in the tumor microenvironment could have tumor promoting effects, whereas other studies showed that MCs have tumor suppressor effects. In our study, we found that the presence of MCs in the tumor microenvironment is beneficial for colon cancer patients; patients with more MCs have a better prognosis and longer survival, indicating that MCs have an antitumor effect.

15-PGDH is a tumor suppressor known to be downregulated in colon cancer. In our study, we found that colon cancer patients expressing higher levels of 15-PGDH have good prognosis compared to those who have low expression. We also found that WNT5A, another tumor suppressor, interacts with 15-PGDH. Patients expressing high levels of both proteins have even better prognosis and longer survival than patients expressing low levels of both proteins. We also found that
WNT5A and its mimicking peptide, Foxy-5, can suppress β-catenin, which is often mutated in colon cancer. We also found that the eicosanoid LTC₄, which is an inflammatory mediator present in the tumor microenvironment, has an anti-tumor effect by inducing the expression of 15-PGDH and inducing the differentiation of colon cancer cells by changing the phenotype of cancer cells to more normal-like cells.

In summary, these new findings provide a better understanding of both the tumor microenvironment and tumor suppressor genes and may help to identify new therapeutic targets for colon cancer treatment.

Syftet med min avhandling var att utvärdera den kliniska betydelsen av tumören omgivande miljö, som mastceller, och tumörsuppressorgener, som 15-hydroxyprostaglandin dehydrogenas (15-PGDH) och WNT5A i tjocktarmscancer, och om re-expression av dessa tumörsuppressorer skulle kunna vara en attraktiv terapeutisk strategi för behandling av tjocktarmscancer.

Mastceller är immunceller som produceras i benmärgen. De är välkända i allergiska reaktioner, men deras roll i cancer, inklusive koloncancer, är fortfarande inte klarlagda. Tidigare studier har visat att mastceller i tumören mikromiljö kan ha tumörbefrämjande effekterna, medan andra studier visade att mastceller har tumörhämmande effekter. I vår studie fann vi att närvaron av mastceller i tumören mikromiljön är fördelaktigt för koloncancerpatienter; patienter med fler mastceller har en bättre prognos och längre överlevnad, vilket indikerar att mastceller har en antitumöreffekt.

15-PGDH är en tumörsuppressor som ofta är nedreglerad i tjocktarmscancer. I vår studie fann vi att tjocktarmscancerpatienter som uttrycker högre nivåer av 15-PGDH har god prognos jämfört med patienter som har lågt uttryck. Vi fann också att WNT5A en annan tumörsuppressor, interagerar med 15-PGDH. Patienter som uttrycker höga nivåer av båda proteinerna har ännu bättre prognos och längre
överlevnad än patienter som uttrycker låga nivåer av dessa proteiner. Vi fann också att WNT5A eller dess peptid, Foxy-5, kan undertrycka β-catenin, som ofta är förändrad i tjocktarmscancer. Vi fann också att leukotrien C4, som är en inflammatorisk mediator som finns i tumörens mikromiljö, har en anti-tumöreffekt genom att öka mängden av tumörsuppressort proteinet 15-PGDH samt att inducera differentiering av koloncancerceller genom att ändra fenotypen av cancerceller till en mer ”normal-liknande cell”.

Sammanfattningsvis, kan dessa nya rön ge en bättre förståelse för både tumörens mikromiljö och tumörsuppressort proteinerna 15-PGDH och WNT5A som kan bidra till att identifiera nya terapeutiska mål för behandling av tjocktarmscancer.
دور الجينات الكابحة لسرطان القولون، آلية وفرص العلاج

السرطان هو نمو الخلايا غير المنضبط مع القدرة على الغزو أو الإنتشار إلى أجزاء أخرى من الجسم. سرطان القولون هو ثالث أكثر أنواع السرطان شيوعاً في العالم، وهو مرض متعدد العوامل والأسباب، منها أسباب بيئية مثل الإستهلاك المفرط للكحول، والوجبات الغنية باللحوم الحمراء أو منخفضة الألياف، بالإضافة إلى انخفاض النشاط البدني، وفضلاً عن الأسباب الوراثية، والتي ترتبط إما بتعديل الجينات السرطانية أو الجينات التي يمكن أن تسبب السرطان، أو تثبيط الجينات الكابحة للورم.

ويرتبط سرطان القولون ارتباطاً وثيقاً بالإنتفاخ المزمن والإنتفاخ الرئيسي.

كما أن المرضا الذين يعانون من الاستهلاك المفرط للخواص هم خطر الإصابة بسرطان القولون. وقد أظهرت العديد من الدراسات أن الأورام السرطانية تحتوي على خلايا مناعية ذات تأثير التهابي واسع النطاق مع خاصية منع أي نمو غير طبيعي والقدرة على القضاء على الخلايا السرطانية.

وكان الهدف من هذه الرسالة تقييم الأهمية السريرية للخلايا المناعية والجينات الكابحة لورم القولون، بغية استخدامها كاستراتيجية علاجية لمرضي سرطان القولون مستقبلاً.

توصلت الدراسة إلى أن وجود هذه الخلايا المناعية والجينات الكابحة مفيد لمريضي سرطان القولون من الناحية السريرية فهي ذات تأثير مضاد للورم. مما يعزز فرص البقاء على قيد الحياة مدة أطول.

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References


161. Kanaoka Y, Boyce JA. Cysteinyl leukotrienes and their receptors: cellular
distribution and function in immune and inflammatory responses. Journal of

162. Maltby S, Khazaie K, McNagny KM. Mast cells in tumor growth: angiogenesis,
tissue remodelling and immune-modulation. Biochimica et biophysica acta.

163. Nielsen HJ, Hansen U, Christensen IJ, Reimert CM, Brunner N, Moesgaard F.
Independent prognostic value of eosinophil and mast cell infiltration in colorectal

164. Khazaie K, Blatner NR, Khan MW, Gounari F, Gounaris E, Dennis K, et al. The
significant role of mast cells in cancer. Cancer and Metastasis Reviews.

165. Gulubova M, Vlaykova T. Prognostic significance of mast cell number and
microvascular density for the survival of patients with primary colorectal cancer.

166. Yodavudh S, Tangjitgamol S, Puangsa-art S. Prognostic significance of microvessel
density and mast cell density for the survival of Thai patients with primary colorectal
cancer. Journal of the Medical Association of Thailand = Chotmaihet thangphaet.

and mast cell density in the prognostic assessment of colorectal carcinomas.
Digestive and liver disease: official journal of the Italian Society of
Gastroenterology and the Italian Association for the Study of the Liver.

168. Imada A, Shijubo N, Kojima H, Abe S. Mast cells correlate with angiogenesis and
poor outcome in stage I lung adenocarcinoma. The European respiratory journal.

Crosstalk between mast cells and pancreatic cancer cells contributes to pancreatic
tumor progression. Clinical cancer research: an official journal of the American

expression of peroxisome proliferator-activated receptor gamma (PPARG,
PPARgamma) is associated with good prognosis. Gastroenterology.

cell phenotypes and microvessels in non-small cell lung cancer and its prognostic

172. Welsh TJ, Green RH, Richardson D, Waller DA, O'Byrne KJ, Bradding P.
Macrophage and mast-cell invasion of tumor cell islets confers a marked survival


Role of novel tumor suppressors in colon cancer: Mechanisms and therapeutic opportunities

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