The role of chemokine receptor 9/chemokine ligand 25 signaling: From immune cells to cancer cells (Review)

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Abbreviations: CCR, chemokine receptor; CCL25, chemokine ligand 25; IL, interleukin; RA, retinoic acid; P-gp, P-glycoprotein; ERM, Ezrin/Radixin/Moesin; GPR-9-6, G-protein-coupled-receptor-9-6; TECK, thymus-expressed chemokine; Maf, muscular aponeurotic fibrosarcoma; T-ALL, T cell lineage acute lymphocytic leukemia; T-CLL, T cell chronic lymphocytic leukemia; CXCL12, CXC chemokine ligand 12; PE38, Pseudomonas exotoxin 38; HTLV-1, human T lymphotropic virus type 1; GSK-3β, glycogen synthase kinase 3β; FKHR, forkhead in human rhabdomyosarcoma; Wnt5a, wingless-type protein 5a; RhoA, Ras homologue; ROCK, Rho kinase; DLL4, delta-like 4; JAG1, Jagged 1; DLK1, delta-like proteins 1.

Key words: chemokine receptor 9, immune system, tumor suppressor genes

Abstract. Chemokine ligand 25 (CCL25) and chemokine receptor 9 (CCR9) are important regulators of migration, proliferation and apoptosis in leukocytes and cancer cells. Blocking of the CCR9/CCL25 signal has been demonstrated to be a potential novel cancer therapy. Research into CCR9 and CCL25 has revealed their associated upstream and downstream signaling pathways; CCR9 is regulated by several immunological factors, including NOTCH, interleukin 2, interleukin 4 and retinoic acid. NOTCH in particular, has been revealed to be a crucial upstream regulator of CCR9. Furthermore, proteins including matrix metalloproteinases, P-glycoprotein, Ezrin/Radixin/Moesin and Livin are regulated via phosphatidylinositol-3 kinase/ protein kinase B, which are in turn stimulated by CCR9/CCL25. This is a review of the current literature on the functions and signaling pathways of CCR9/CCL25.

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1. Introduction

The cDNA of CCR9 was submitted to Genbank (Genbank no. HSU45982) by Lautens in 1996 as G-protein-coupled-receptor-9-6 (GPR-9-6). The expression and function of GPR-9-6 was investigated by Zaballos et al (1) in 1999. GPR-9-6 was renamed as CCR9, due to its similarity in structure and sequence to chemokine receptor 6 (CCR6), chemokine receptor 7 (CCR7) and STRL33/Bonzo (1). Furthermore, in vitro stimulation of CCR9 with thymus-expressed chemokine (TECK) induced intra-cytoplasmic calcium mobilization and migration of 293 cells (1). TECK, also known as CCL25, was discovered in thymic dendritic cells in 1997 (2), and was subsequently confirmed to be the sole ligand of CCR9 (3). CCR9 expression was revealed to be low in human and murine spleen and lymph nodes but high in thymic tissues (1).

Further studies demonstrated that CCR9 is highly expressed in the colon, small intestine and several other tissues involved...
in the development and maturation of T cells, macrophages and dendritic cells (DCs) (4-9). CCR9 signaling may therefore have an influence on processes, including inflammatory responses and transplantation rejection (10-12). Furthermore, CCR9 has been demonstrated to influence cancer cell migration, proliferation and drug resistance (13-15). This is a review of the current literature on the function of CCR9 in leukocytes and cancer cells.

2. Chemokine receptor 9 in leukocytes

The function of CCR9 in different leukocyte subtypes is relatively comparable (Fig. 1).

CCR9+ DCs migrate to the small intestine and colon to interact with other leukocytes, particularly T cells, to induce local inflammation (4-6). Thymic epithelial cells are the principle source of CCL25 (7); however, thymic DCs also secrete CCL25, which may contribute toward DC-T cell interactions (2).

CCR9 serves an important role in the regulation of T cells. CCR9 (16) and CCR7 (17,18) have been demonstrated to contribute toward the recruitment of T cells to the thymus. T cell development is also regulated by CCR9 (19). CCR9-knockdown skewed T cell subgroup ratios; thymic αβ-T cell development was not affected in CCR9−/− mice, whereas the number of γδ-T cells located in the intestine was increased (19). In addition, CCR9 suppressed differentiation of forkhead box protein 3+ regulatory T cells (20). Several T cell functions were revealed to be CCR9-dependent; CD4+ CCR9+ T helper cells expressed interleukin 21 (IL-21), inducible T cell co-stimulator, transcription factor B-Cell CLL/lymphoma 6 and muscular aponeurotic fibrosarcoma (Maf), and supported B cell antibody production (21). In splenic T cells, blocking CCR9/CCL25 signaling reduced secretion of interferon-γ (IFN-γ) (12). This may serve a role in certain immunological processes, including transplant rejection and the inflammatory response. Anti-CCL25 antibodies decreased the infiltration of cells around skin allografts, which prolonged the survival of the graft (12). Recruitment of T cells to the intestine and colon was demonstrated to be CCR9-dependent (22,23). CCR9+ T cells were recruited to inflammatory bowel disease lesions, and were revealed to be associated with disease activity (10). Similarly, during Aggregatibacter actinomycetemcomitans serotype b infection, CCR9 expression was increased in T cells (11).

Retinoic acid (RA) has been demonstrated to serve a role in DC-T cell interactions. Several studies revealed that CCR9 expression fluctuated at different stages of DC development (9,24). Furthermore, several other studies demonstrated that DCs produced RA, which mediated CCR9 expression in T cells (25) and induced interleukin 10 (IL-10) expression in αβ-T cells (26). T cells were also mediated by RA directly via the RA receptor, which upregulated CCR9 expression (27,28). Therefore, decreased expression of the RA receptor was associated with a reduction in CCR9 expression and an attenuation of graft-versus-host disease (29).

Naïve B cell migration has been demonstrated to be regulated by CCR9 (30). A high CCR9 expression has also been revealed in memory B cells (31). Epstein-Barr virus-infected B cells demonstrated increased CCR9 expression (32). However, CCR9/CCL25 signaling does not appear to be vital for B development, as maturation of B cells was not impaired in CCR9−/− mice (19).

Macrophages migrate to infected peritoneal (33) and acute colitis lesions (6). Macrophages recruited by CCR9, which produced tumor necrosis factor-α (TNF-α), were crucial for liver fibrosis (34). CCR9 expression in these recruited macrophages was regulated by their interaction with hepatic stellate cells (35).

In summary, CCR9 serves a role in the migration, maturation and function of leukocytes.

3. Chemokine receptor 9 in cancer cells

Following the discovery of CCR9/CCL25 signaling pathways in leukocytes, oncology researchers revealed that CCR9 promotes invasion, migration, anti-apoptosis and drug-resistance in several types of tumors (Fig. 1).

CCR9/CCL25 signaling has been rigorously studied in hematological malignancies. In T cell lineage acute lymphocytic leukemia (T-ALL) CD4+ cells, CCR9 was highly expressed, while T cell chronic lymphocytic leukemia (T-CLL) CD4+ cells moderately expressed CCR9, and a low CCR9 expression was demonstrated in normal CD4+ T cells (36). The CCR9+ T-ALL Jurkat, MOLT-4, CEM, SupT1, TALL-1 and DND41 cell lines were revealed to be chemotactic to CCL25 (37). CCL25 stimulation resulted in pseudopodium formation (38), invasion and migration (39). It has been observed that T cells from adult patients with leukemia, that had infiltrated the gastrointestinal tract, were frequently positive for CCR9 (40), which was in line with conclusions from a study that reported that CCR9 expression was associated with gut relapse in pediatric T-ALL (41). Furthermore, CCR9/CCL25 signaling induced P-gp co-localization within the actin cytoskeleton (15). This mechanism led to drug resistance to doxorubicin in the MOLT-4 cell line (15). TNF-α mediated apoptosis was inhibited in CD4+ T-ALL, T-CLL and MOLT-4 cells by CCR9/CCL25 signaling (42). Transformation of gastric mucosa-associated lymphoid tissue to gastric extra-nodal diffuse large B-cell lymphoma was mediated by several chemokines, including CCR9 (43). However, CCR9+ multiple myeloma cells only migrated toward CXC chemokine ligand 12 (CXCL12), indicating a limited function of CCR9 in multiple myeloma (44).

The regulation of melanoma cell metastasis by CCR9/CCL25 signaling remains controversial. Certain studies have demonstrated that CCR9 expression is associated with organ-specific metastasis of melanoma cells (45,46). Melanoma cells that had metastasized to the intestine expressed CCR9, whereas cells that had metastasized to other organs did not (47). Besides the intestine, lymph nodes and skin were the sites where melanoma metastasis often occurred; however, CCR9+ melanoma cells were not observed in these locations (47-49). CCR9+ T cells in the melanoma microenvironment have been demonstrated to inhibit metastasis (50). However, certain studies have demonstrated that the function of CCR9 in melanoma was only moderate; the proportion of patients with intestine metastases was low (51), indicating limited organ-specific metastasis (49), which corresponds with the low proportion of CCR9+ cells reported in circulating conditions.
tumor cells (52). Clinical studies demonstrated that CCR7 and chemokine receptor 10 (CCR10), but not CCR9, were associated with a poorer prognosis (53).

Several types of ovarian carcinoma, including serous adenocarcinoma, serous papillary cystadenoma and endometrioid adenocarcinoma, revealed an increased CCR9 expression compared with normal ovarian tissues (54), and demonstrated migration and invasion potential toward chemotactic gradients of CCL25 (55). Intestinal cells, a source of CCL25, were also a frequent metastatic site for ovarian carcinoma (55).

CCR9 expression of two breast cancer cell lines, MDA-MB-231 and MCF7, and benign tissue, were high, medium and low, respectively, which corresponded with their respective aggressiveness (56). Lymph nodes, bone marrow, lung, liver and brain were frequent sites of breast cancer metastasis, and this process might be regulated by CCR9/CCL25 (56-58). CCR9/CCL25 signaling was also revealed to provide a survival advantage to breast cancer cells and inhibited cisplatin-induced apoptosis (59).

In prostate cancer, CCR9 was highly, moderately and lowly expressed in highly invasive LNCaP, moderately invasive PC3 and non-invasive prostatic epithelial cells, respectively (60). CCR9/CCL25 also regulated metastasis (60) and drug-resistance against etoposide (61) in prostate cancer.

In pancreatic cancer, pancreatic intraepithelial neoplasia and pancreatic cancer cells were demonstrated to be CCR9+ (62). The pancreatic cancer PANC-1 cell line was CCR9+, and enhanced invasion was observed following CCL25 treatment (63). Drug-resistance of gemcitabine was stimulated by CCL25 signaling in pancreatic cancer PANC-1, MIAPaCa-2 and AsPC-1 cell lines (14).

In colon cancer, injection of CCR9+ cancer-initiating cells led to formation of gastrointestinal xenograft tumors in mice, whereas blocking CCR9 signaling increased extra-intestinal tumors (13,64).

In the hepatocellular carcinoma cell lines HepG2 and Huh7, CCR9 promoted invasion and migration (65), and might be a marker to predict the prognosis of patients (66).

Similarly, CCR9 could be beneficial in predicting lymph node metastasis and prognosis in lung adenocarcinoma (67). Adenocarcinoma cells revealed a higher migratory and invasive potential in response to CCL25, compared with squamous cell carcinoma cells, which had lower expression of CCR9 and CCL25 (68). Esophageal cancer cells also highly expressed CCR9 and potentially achieved metastasis via CCR9 signaling (69).

Based on these findings, researchers attempted to design CCR9-specific therapies, including CCR9 antagonists, monoclonal antibodies against CCR9, and RNAi of CCR9. Computational modeling of CCR9 antagonists revealed several compounds, one of which inhibited proliferation and invasion of pancreatic cancer cell lines and interacted synergistically with gemcitabine (14). In vivo models revealed that a CCR9 monoclonal antibody increased apoptosis, necrosis of tumor tissue, complement-dependent cytotoxicity and antibody-dependent cytotoxicity by natural killer cells. The antibody was also demonstrated to decrease proliferation and tumor vascularization in MOLT-4 cell lines (70). CCR9 inhibition by RNAi facilitated T cell-associated immunotherapy of breast and pancreatic cancer cell lines (71). Another trial used CCL25 fused with Pseudomonas exotoxin 38 (PE38) toxin, a truncated derivative of Pseudomonas exotoxin A, which induced apoptosis in MOLT-4 cell lines (72).

In summary, CCR9/CCL25 signaling is an important mediator of malignant behaviors in a number of cancer cells. Blocking of CCR9/CCL25 signaling appears to be a potential novel strategy for cancer therapy.

Figure 1. Function of CCR9/CCL25 signaling in immune cells and cancer cells. Blocks in orange indicate positive regulation, while blue indicates negative regulation by CCR9/CCL25 signaling. CCR9, chemokine receptor 9; CCL25, chemokine ligand 25; FoxP3+, Forkhead box P3; IL-21, interleukin 21; ICOS, inducible T cell co-stimulator; INF-γ, interferon-γ; DCs, dendritic cells; T-ALL, T-cell acute lymphoblastic leukaemia.
4. Regulation of chemokine receptor 9

Several immunological factors have been demonstrated to upregulate CCR9, including human T-lymphotropic virus type 1 (HTLV-1)-encoded transcriptional activator Tax (40), RA (25) and Epstein-Barr virus (32). Others, including DAPT (37) and co-stimulation of IL-2 with IL-4 (36), have been revealed to downregulate CCR9. The NOTCH pathway has been revealed to mediate CCR9 expression; however, its function differs in T-ALL and colon cancer (Fig. 2).

NOTCH1 mutations were frequently identified in T-ALL; certain mutations resulted in 6l ligand-independent NOTCH signaling or lengthened the half-life period of NOTCH1 signaling. Suppression of NOTCH1 by DAPT or RNAi, downregulated CCR9 in the T-ALL cell lines Jurkat, MOLT-4 and CEM, SupT1, TALL-1 and DND41 (37).

However, in CCR9+ colon cancer cells, NOTCH signaling was revealed to be reduced compared with that in CCR9- colon cancer cells. Since no significant difference in CCR9 mRNA levels was revealed between CCR9+ and CCR9- cells, it was confirmed that NOTCH lowered CCR9 expression levels by increasing proteasomal degradation (13).

5. Downstream signaling of chemokine receptor 9/chemokine ligand 25

CCL25 binding led to interactions between CCR9 Gβγ and PI3K, which resulted in activation of Akt (73). In this study, glycogen synthase kinase 3β (GSK-3β) and forkhead in human rhabdomyosarcoma (FKHR) were demonstrated to be downstream targets of Akt (73), which further influenced the migration, invasion and drug resistance of cancer cells. An alternative stimulation of PI3/Akt by wingless-type protein 5a (Wnt5a) following CCR9/CCL25 interaction was revealed in another study (74), indicating a complex cross-talk between these signaling pathways (Fig. 2).

Mediation of cell invasion and migration by CCR9/CCL25 signaling may involve matrix metalloproteinases (MMPs) (55,56,60), Ras homologue (RhoA)Rho kinase (ROCK)-myosin light chain (MLC) signal (39), Wnt5a (74) and ERM protein family (38). In the ovarian cancer OVCAR-3 and CAOV-3 cell lines, CCR9/CCL25 selectively regulated certain MMPs, which degraded the cell matrix to promote invasion (55). Different patterns of MMP regulation were observed in each cell line; for example, in the OVCAR-3 cell line, expression of MMP-2, 3, 8, 9, 10, 11 and 13 was increased, while MMP-1, 2, 3, 8 and 10 expression levels were elevated in the CAOV-3 cell line (55). A similar mechanism was observed in prostate cancer cell lines (60). CCL25 treatment increased MMP-1 and 2 expression in LNCaP, and MMP-2, 11 and 13 in PC3 cells (60). RhoA-ROCK-MLC signaling, which is associated with cell motility and morphology was stimulated by CCL25 in the MOLT-4 cell line (39). Another pathway has been demonstrated to promote the invasion and migration of MOLT-4 cells; CCL25 induced Wnt5a activation by promoting protein kinase C expression and activation in MOLT-4 cells (74). Activated Wnt5a further induced PI3K/Akt signaling and enhanced cell migration and invasion (74). Invasion-related ERM protein was revealed to be translocated from the cytoplasm to the cell membrane following CCL25 treatment, which contributed to metastasis (38).

Stimulation of drug-resistance by CCL25 might be achieved via P-gp (15), inhibition of GSK-3β and FKHR (59), Livin protein (42), anti-apoptosis protein (PI3K, AKT, extracellular-regulated kinase 1/2 (ERK1/2) and GSK-3β) and caspase-3 (61). P-gp was revealed to promote the extrusion of the antitumor agent, doxorubicin, from the cytoplasm to the extracellular space, resulting in multidrug resistance. This resistance was dependent on interactions between ERM and the actin cytoskeleton following CCL25 stimulation (15). In breast cancer, inhibition of CCR9/CCL25 downstream signaling via GSK-3β or FKHR, significantly improved the chemotherapeutic efficacy of cisplatin (59). Alongside upregulation of anti-apoptosis proteins, including PI3K, Akt, ERK1/2 and GSK-3β, CCL25 also induced drug resistance via downregulation of caspase-3 activity (61). CCL25 suppressed TNF-α-mediated apoptosis by increasing expression of Livin, a member of the inhibitor of apoptosis protein family, via c-jun-NH2-kinase 1 kinase activity (42).

6. Hypothesis and conclusion

NOTCH and several other signaling proteins regulate CCR9, and CCR9/CCL25 signaling mediates certain downstream...
proteins to promote metastasis and drug-resistance in cancer cells. Based on the findings that healthy intestine and colon cells physiologically produce CCL25 (2,6,7,5), we hypothesized that CCL25 induces chemotaxis and cell survival signaling in leukocytes and cancer cells. For extra-intestinal cancer, the CCL25 concentration in healthy intestinal micro-environment is higher than that in the primary tumor micro-environment; thus, CCR9+ cancer cells tend to metastasize to the intestine. However, the micro-environment of colon cancer is rich in CCL25 and therefore, CCR9 may be downregulated by NOTCH to reduce chemotaxis and promote metastasis. Investigation of CCL25 levels in the micro-environment of primary and metastatic lesions would further test this hypothesis. Another possible mechanism is based on the seed-earth hypothesis, which states that CCL25 promotes the survival of circulating tumor cells that have metastasized to the intestine.

NOTCH signaling has an opposite function in T-ALL and colon cancer. This contradiction might be explained by the diversity in ligands and downstream signaling. In T-ALL, it is a ligand of NOTCH1 (37), while delta-like 4 (DLL4), Jagged 1 (JAG1) and delta-like proteins 1 (DLK1) are the most probable ligands of NOTCH in colon cancer (13).

In summary, CCR9/CCL25 signaling mediated the migration, invasion and drug resistance of cancer cells. Further studies should focus on elucidating the mechanisms of associated upstream and downstream signaling of CCR9.

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Competing interests
The authors declare that they have no competing interests.

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