Wnt5b-associated exosomes promote cancer cell migration and proliferation

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Wnts are secretory proteins that are conserved evolutionally and regulate at least two intracellular signaling pathways: β-catenin-dependent and -independent. Among 19 human or mouse Wnt family members, Wnt1 and Wnt3a activate the β-catenin-dependent pathway, whereas Wnt5a and Wnt11 activate the β-catenin-independent pathway. Wnt5a is one of the most extensively studied Wnts and has been shown to be essential for embryonic development. Wnt5a is also involved in several post-natal diseases, including cancer and inflammatory diseases. Wnt5b is a Wnt5a family member (80% identity) in mammals; one Wnt5 also exists in Drosophila and zebrafish (pipetail). Wnt5b is essential for convergent extension in zebrafish, whereas in the mouse, Wnt5a, but not Wnt5b, is essential for development.

Evidence has accumulated that Wnt5b expression is involved in mouse chondrocyte maturation and is associated with human diabetic mellitus and adipogenesis. Similar to Wnt5a, it is likely that Wnt5b is involved in cancer progression. Periostin, interferon-induced transmembrane protein 1, and Wnt5b were reported to be highly expressed in head and neck squamous cell carcinomas; their expression promoted the invasive ability of head and neck squamous cell carcinomas by inducing MMP-10. Wnt5b was also highly expressed in breast cancer with high invasive activity, and Wnt5b expression in basal-like cancers was associated with brain metastasis. In breast cancers and mammary stem cells, secreted MMP-3 interacted with and inactivated Wnt5b, thereby enhancing the β-catenin-dependent pathway. Lung cancer and pancreatic cancer cells produced Wnt5b, which promoted invasion activity, when they were treated with transforming growth factor-β (TGF-β). However, the biochemical properties and cancer-associated activities of Wnt5b have not been well characterized compared with Wnt5a because Wnt5b has not yet been purified.

Wnt proteins are post-translationally glycan- and lipid-modified secretory proteins that are ready to self-aggregate and bind heparin sulfate proteoglycans. Therefore, Wnts are not easily diffusible in the extracellular milieu. Several possible mechanisms have been proposed to explain the long-distance movements of Wnts. Lipoprotein particles are large, globular complexes composed of a central core of hydrophobic lipids that are associated with apoproteins and surrounded by a monolayer of membrane phospholipids. In Drosophila, the Wingless (Wg) protein derived from wing discs was found to...
be copurified with lipoprotein particles. Larvae with reduced lipoprotein particles showed a narrow expression of Distalless, a target gene of Wnt signaling, in wing porch. In mammals, Wnt3a was released from mouse fibroblasts by the presence of lipoproteins in the culture medium, specifically through high-density lipoprotein particles. A Wnt3a mutant

### Table 1. Purification of Wnt5b

| Volume, mL | Protein concentration, mg/mL | Total protein, mg | Wnt5b concentration, µg/mL | Wnt5b, µg |
|------------|---------------------------|------------------|---------------------------|-----------|
| Wnt5b CM  | 20 000 (in total)         | 4.67             | 93 400                    | 0.0288    | 576.0     |
| Blue Sepharose | 625 (in total)       | 23.70†          | 14.800                    | 0.3620    | 226.5     |
| Gel filtration | 80 (in total)        | 32.30†          | 2.650                     | 2.2100    | 176.8     |
| Heparin Sepharose | 3.0                | 52.00†          | 0.156                     | 50.0000   | 150.0     |

Concentration of Wnt5b protein in the conditioned medium was determined by comparing its signal intensity on a Wnt5b immunoblot to that of serial dilutions of a known amount of purified Wnt5b protein. Concentration in µg/mL.

![Image](image-url)

Fig. 1. Determination of the glycan profiles of Wnt5b. (a) Coomassie Brilliant Blue (CBB) staining of an SDS polyacrylamide gel containing fractions from all chromatographies in the purification process. CM, conditioned medium. (b) Purified Wnt5a and Wnt5b proteins were stained with CBB and probed with anti-Wnt5a and anti-Wnt5b antibodies. (c) Wnt5b glycopeptides obtained from nano-flow liquid chromatography were separately analyzed by MALDI mass spectrometry (MS). (d) MALDI mass spectrum of a peptide containing Ser223 of Wnt5b. (e) MALDI-MS/MS spectrum from the ion at m/z 1455.9 in (d). The sequences from the N- and C-termi and the position of the modified residue were read out based on y', and b' ions.
that lacked palmitate was not associated with lipoproteins. The lipid modification of Wnt3a could be important for targeting Wnt3a to the phospholipids of lipoproteins. These results suggest that lipoprotein particles act as machinery for the efficient intercellular movement and stabilization of Wnt3a/Wg. In addition, it has been reported that some proteins, including swim and afamin, bound to and solubilized Wg and Wnts. (31,32) It is also possible that Wnt is secreted and spread by being attached to membranous vesicles. (33) A substantial fraction of Wnts were found on extracellular vesicles known as exosomes. (34,35) As intraluminal vesicles that originate in large multivesicular bodies (MVBs), exosomes are released in the extracellular milieu following fusion of MVBs with cell surface membranes. (36,37) In larval neuromuscular junctions, Wg was released with Evi/Wntless-containing extracellular vesicles and traveled from one cell to another at the synaptic cleft. (38) Subsequent electron microscopy and proteomics studies revealed that these vesicles are exosomes. (39) In addition, Wnt3a and Wg are observed on exosomes released from mammalian L and HEK293 and Drosophila S2 cells. (40,41) Fibroblasts secreted exosomes, which were internalized by breast cancer cells (BCCs) and associated with Wnt11. Exosome-associated Wnt11, in turn, promoted BCC protrusion activity and motility to drive invasive behavior. (42) Thus, exosome-associated Wnt ligands could be involved in intercellular communication. However, whether Wnt5b is secreted with exosomes is unclear; if so, whether Wnt5b-associated exosomes show some activity remains to be clarified.

Here, we characterized purified Wnt5b and found that Wnt5b is secreted with exosomes in a cell context. We also showed that Wnt5b-associated exosomes promote cancer cell migration and proliferation in a paracrine manner.

### Materials and Methods

#### Purification of Wnt5b

Wnt5b was purified to near homogeneity from Wnt5b conditioned medium (CM) through three successive column chromatography, including Blue Sepharose HP.
Table 3. Summary of observed and theoretical masses of glycopeptides of human Wnt5b and the relative abundance of each glycoform observed in MALDI mass spectrometry

| Observed molecular mass | Theoretical molecular mass | Sequence | Modification | Position | Relative abundance (%) |
|------------------------|-----------------------------|----------|--------------|----------|------------------------|
| 2652.19                | 2652.06                     | WNCSTADNASVFGR | HexNAc2Hex4 | N93      | 4.3                    |
| 2814.26                | 2814.12                     |                      | HexNAc2Hex5 |          | 2.6                    |
| 2976.32                | 2976.17                     |                      | HexNAc2Hex6 |          | 12.0                   |
| 3138.38                | 3138.22                     |                      | HexNAc2Hex7 |          | 41.2                   |
| 3300.46                | 3300.28                     |                      | HexNAc2Hex8 |          | 35.4                   |
| 3462.54                | 3462.33                     |                      | HexNAc2Hex9 |          | 4.4                    |
| 2301.15                | 2300.96                     | NESTGSLGTQGR        | HexNAc3Hex3 | N291     | 5.3                    |
| 2463.12                | 2463.01                     |                      | HexNAc3Hex4 |          | 9.9                    |
| 2625.19                | 2625.07                     |                      | HexNAc3Hex5 |          | 16.6                   |
| 2828.22                | 2828.14                     |                      | HexNAc4Hex5 |          | 7.3                    |
| 2990.32                | 2990.20                     |                      | HexNAc4Hex6 |          | 23.9                   |
| 3152.40                | 3152.25                     |                      | HexNAc4Hex7 |          | 43.0                   |
| 3187.64                | 3187.34                     | STGSLGTQGRLCNKTS‡  | HexNAc2Hex6 | N305     | 15.1                   |
| 3349.77                | 3349.40                     |                      | HexNAc2Hex7 |          | 37.4                   |
| 3511.81                | 3511.45                     |                      | HexNAc2Hex8 |          | 37.4                   |
| 3673.95                | 3673.50                     |                      | HexNAc2Hex9 |          | 10.2                   |

†Cys (O) is an S-(propionamide) cysteine. ‡Peptide was obtained from the Glu-C digest of Wnt5b.

Results

Purification and characterization of Wnt5b. To characterize Wnt5b biochemically, we first purified Wnt5b from CM of L cells that stably expressed Wnt5b (L/Wnt5b cells) by the use of three successive conventional column chromatographies (Table 1, Fig. 1a). Using peptides based on different amino acid sequences between Wnt5a and Wnt5b as antigens, anti-Wnt5 antibodies that specifically recognized Wnt5a and Wnt5b were generated (Fig. 1b). Purified Wnt5b induced Dvl2 phosphorylation in NIH3T3 cells (Fig. S1a) and activated Rac in HeLaS3 cells to a similar extent as purified Wnt5a (Fig. S1b).

Wnt5b has four possible N-glycosylation sites, namely Asn−X−Ser/Thr, including N93, N99, N291, and N305. Wnt5b peptides, which were digested with trypsin or Glu-C, were subjected to nano-flow liquid chromatography (LC) followed by MALDI mass spectrometry (MS)/MS and nano-flow LC/electrospray ionization MS (LC/ESI-MS). Approximately 70% of the entire Wnt5b amino acid sequence was covered (Table 2). Among the four possible glycosylation sites, N93, N291, and N305 were glycosylated with high-mannose-, hybrid-, and high-mannose-type oligosaccharides, respectively (Fig. 1c). The MALDI-MS/MS analysis of the peptide containing N93 and N99 observed at m/z 3300.46 revealed that N99 is not glycosylated (Fig. S1c). Further analyses of digested peptides using nano-flow reversed-phase LC and MALDI-MS/MS confirmed that glycans were attached to N93, N291, and N305 of Wnt5b (Table 3).

A peptide containing S223, observed at m/z 1455.9 (Fig. 1d), was identified as a peptide with a monounsaturated fatty acid (C16:1, palmitoleic acid) (Fig. 1e), indicating that Wnt5b was modified with palmitoleic acid as well as Wnt3a, Wnt5a, and Wnt11.46,47

Wnt secretion modes vary among cell lines. Wnts are secreted into the extracellular space through secretory vesicles or extracellular microvesicles. In cultured cells, Wnts were reported to be secreted with extracellular microvesicles, such as exosomes, which were recovered in the 100 000 g centrifugation precipitation fractions from CM. These fractions contained clathrin and tumor susceptibility gene 101 (TSG101), which play roles in the trafficking of exosomes.48,49 Wnts released by the fusion of secretory vesicles with cell surface membranes were detected in supernatant fractions. In L cells stably expressing Wnts, most Wnt5b, Wnt5a, and Wnt3a proteins were recovered in supernatant fractions; Wnts were not detected in precipitation fractions, or only a small fraction was detected (Fig. 2a). In MDCK cells expressing Wnts, almost all Wnt1, Wnt5a, and Wnt11 proteins were recovered in supernatant fractions (Fig. S2a). Some cancer cell lines, including HeLaS3 cervical cancer, A549 lung adenocarcinoma, and KKLs gastric...
cancer cells, expressed Wnt5a endogenously, and most Wnt5a secreted in CM was present in supernatant fractions (Fig. 2b). Therefore, in these cells, Wnt1, Wnt3a, Wnt5a, and Wnt11 might not be associated with extracellular microvesicles prepared by 100,000 g centrifugation, but it is possible that there are microvesicles associated with other Wnts in supernatant fractions when exosomes are prepared by other methods.

Almost all Wnts were recovered in precipitation fractions when Wnt5b, Wnt5a, and Wnt3a were transiently expressed in Caco-2 colon cancer cells (Fig. 2c). However, Wnt5b NQ (N93Q, N99Q, N291Q, and N305Q) mutant that lacks glycosylation and Wnt5b S223A mutant that lacks lipidation were hardly recovered in precipitation and supernatant fractions of CM (Fig. S2b,c), suggesting that glycosylation and lipidation are essential for Wnt5b secretion and its association with exosomes.

To analyze endogenous Wnt5b on exosomes, the expression levels of Wnt5b mRNA were quantified in various cancer cell lines. Among various types of cancer cells, Wnt5b mRNA was highly expressed endogenously in PANC-1 cells (Fig. S2d). Wnt5b mRNA expression was the highest in PANC-1 cells compared with other Wnt mRNA (Fig. S2d). Approximately 55% of endogenous Wnt5b released from PANC-1 cells was recovered in precipitation fractions, which also contained CD63, CD81, and TSG101 that are known to be present on exosomes (Fig. 2d). Therefore, whether secreted Wnt5b and exosome markers, such as CD81 and TSG101, were not co-sedimented, was unclear, but it might be due to Wnt5b not only being associated with exosomes containing CD81 and TSG101, but also other exosomes.
Development of a reporter assay system to measure Wnt5a and Wnt5b signaling. As Wnt5b-associated exosomes prepared from 50 mL CM did not show Dvl2 phosphorylation or Rac activation activities (data not shown), we developed a highly sensitive reporter system to measure Wnt5a and Wnt5b signaling using CHO cells. As shown in a previous report,\(^{54}\) the N-terminal region of Ror2 was fused with the transmembrane and intracellular region of low-density lipoprotein receptor-related protein 6 (LRP6) to thereby construct a Ror2-LRP6 chimera (Fig. 3a). The chimera construct was stably expressed in CHO cells that had been already transfected with a TOPflash gene, which is a luciferase reporter plasmid that contains wild-type T-cell factor binding regions (CHO/Ror2-LRP6-TOP cells; Fig. 3a).

Using CHO/Ror2-LRP6-TOP cells, Wnt5a activity was measured as control. Wnt5a CM induced the phosphorylation of the Ror2-LRP6 chimera under the same conditions used in the induction of the phosphorylation of endogenous LRP6 by Wnt3a CM (Fig. 3b). Although Wnt5a CM did not increase luciferase activity without Ror2-LRP6 chimera expression, chimera expression strongly promoted a Wnt5a-dependent increase in luciferase activity (Fig. 3c). Knockdown of endogenous Frizzled 2 (Fz2) or the addition of secreted Frizzled-related protein-2 sFRP2 \(^{6,55}\) reduced the Wnt5a-dependent luciferase activity (Fig. 3d,e). Therefore, in this reporter system, Wnt5a activates transcription factor T-cell factor activity through the Ror2-LRP6 chimera and endogenous Fz2. Similar to Wnt5a, purified Wnt5b increased luciferase activity in a dose-dependent manner (Fig. 3f). Such a reporter system is therefore useful for quantitatively measuring Wnt5a or Wnt5b activity.

Wnt5b-associated exosomes from PANC-1 cells are active. To examine endogenous Wnt5b activity in PANC-1 cells, the WNT5B gene was successfully knocked out by a CRISPR/Cas9 system (Fig. 4a); Wnt5b protein expression was completely lost (Fig. 4b). WNT5B knockdown (KO) PANC-1 cells showed reduced migration and invasion activity (Fig. 4c). Transforming growth factor-β increased the expression of Snail and vimentin mRNA, and decreased E-cadherin mRNA expression in PANC-1 cells (Fig. 4d). In WNT5B KO PANC-1 cells, TGF-β-dependent increases in Snail and vimentin mRNAs and the decrease in E-cadherin mRNA were attenuated (Fig. 4d), suggesting the involvement of Wnt5b signaling in epithelial–mesenchymal transition induced by TGF-β. However, WNT5B KO did not affect PANC-1 cellular proliferation (Fig. 4e). These results are consistent with previous observations using Wnt5b siRNA.\(^{25}\)

Using CHO/Ror2-LRP6-TOP cells, Wnt5b-associated exosome activity was measured. Precipitation fractions from CM of PANC-1 cells increased luciferase activity in a dose-dependent manner, but those from WNT5B KO PANC-1 cells decreased luciferase activity dramatically (Fig. 4f). To clarify the biological significance of paracrine mechanisms of Wnt5b-associated exosomes, we used A549 cells as target cells, because A549 cells were reported to migrate and proliferate in response to Wnt5a.\(^{20}\) Wnt5a was knocked down to reduce endogenous Wnt5a effects in A549 cells (A549/shWnt5a cells). Exosomes prepared from wild-type PANC-1 cells, but not those from Wnt5b KO PANC-1 cells, stimulated cell migration of, and increased 5-ethyl-2′-deoxyuridine (EdU) incorporation into A549/shWnt5a cells (Fig. 4g,h). Therefore, Wnt5b secreted with exosomes is active.

Fig. 3. Development of reporter assay to measure Wnt5a and Wnt5b signaling. (a) Schema of the reporter system used to measure Wnt5a/b signaling including CHO cells stably expressing a Ror2–low-density lipoprotein receptor-related protein 6 (LRP6) chimera and TOP-flashing (CHO/Ror2-LRP6-TOP cells). GSK3-β, glycogen synthase kinase 3β; TCF/LEF, T-cell factor/lymphoid enhancer factor. (b) CHO/Ror2-LRP6-TOP cells were stimulated with 1 mL control, Wnt3a, or Wnt5a conditioned medium (CM) for 3 h, and cell lysates were probed with anti-phospho-LRP6 and anti-LRP6 antibodies. (c) CHO/Ror2-LRP6-TOP cells and parent CHO cells stably expressing only TOP-flashing were stimulated with control or Wnt5a CM for 8 h. A luciferase assay was carried out and the activity was expressed as a fold increase compared with samples without Wnt5a and Ror2-LRP6. Results are shown as means ± SD of three independent experiments. *P < 0.05. (d) CHO/Ror2-LRP6-TOP cells, which had been transfected with control or Frizzled-2 (Fz2) siRNAs, were stimulated with control or Wnt5a CM for 8 h. (e) CHO/Ror2-LRP6-TOP cells were stimulated with Wnt5a CM for 8 h in the presence of the indicated concentrations of secreted Frizzled-related protein-2 sFRP2. (f) CHO/Ror2-LRP6-TOP cells were stimulated with the indicated concentrations of purified Wnt5b for 8 h.

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To confirm that Wnt5b-associated exosomes are taken up by cells, exosomes prepared from PANC-1 cells were labelled with PKH26. Exosome uptake in CHO cells was observed in a time-dependent manner at 37°C, but not at 4°C (Fig. 5a,b). Internalization of labeled exosomes was also detected as merged images of vertical sections by confocal stacks (Fig. 5c). To observe that Wnt5b is indeed associated with exosomes, we generated a plasmid vector expressing HA-tagged Wnt5b (HA-Wnt5b), by introducing HA tag sequence at position Arg36, located in a weakly conserved surface region opposite the Frizzled receptor binding site.56 HA-Wnt5b was active comparable to non-tagged Wnt5b in CHO/Ror2-LRP6-TOP cells (Fig. 5d). Immunoelectron microscopy revealed that HA-Wnt5b is present on exosomes isolated from PANC-1 cells expressing HA-Wnt5b (Fig. 5e).

Wnt5b-associated exosomes from Caco-2 cells are active. Caco-2 cells expressed Wnt5a mRNA more highly than Wnt5b mRNA (see Fig. S2d). To reduce endogenous Wnt5a effects, Wnt5a was knocked down in Caco-2 cells stably expressing GFP (Caco-2/GFP cells) or Wnt5b (Caco-2/Wnt5b cells) (Fig. S3a,b). Knockdown of Wnt5a did not affect cell migration and expression of Wnt5b did not affect it either, regardless of Wnt5a expression (Fig. S3c). Therefore, Wnt5b could not be involved in Caco-2 cell motility. However, Wnt5b expression promoted cell proliferation compared with GFP expression, and knockdown of Wnt5b inhibited Wnt5b-dependent cellular proliferation (Figs 6a, S4a).

TSG101 was knocked down by two siRNAs (#1 and #2). Knockdown of TGS101 by siRNA#1 suppressed Wnt5b-associated exosomes from PANC-1 cells and relative migration and invasion activities of A549/shWnt5a cells were expressed as percentages of those of wild-type PANC-1 cells. Results are shown as means ± SD of three independent experiments. *P < 0.05. (d) WT and WNT5B KO PANC-1 cells were stimulated with 5 ng/mL transforming growth factor-β (TGF-β) for 48 h and mRNA levels of Snail, vimentin, and E-cadherin (E-cad) were measured by quantitative real-time PCR. The results were expressed as fold-changes compared with WT cells without TGF-β stimulation. n.s., not significant. (e) WT and WNT5B KO PANC-1 cells were subjected to cell proliferation assay. (f) CHO/Ror2-LRP6-TOP cells were treated with the P100 fraction recovered from the indicated volumes of PANC-1 conditioned medium (CM) for 8 h. (g) A549/shWnt5a cells were subjected to migration assay in the presence of the exosomes isolated from CM of WT and WNT5B KO PANC-1 cells and relative migration activity of A549/shWnt5a cells were expressed as percentages of those of A549/shWnt5a cells in the absence of the exosomes. (h) A549/shWnt5a cells were treated for 48 h with the exosomes isolated from CM of WT and WNT5B KO PANC-1 cells, and then the cells were incubated with 5-ethynyl-2'-deoxyuridine (EdU) for 1 h before fixation and stained with DRAQ5. EdU-positive cells are expressed as percentages of positively stained cells compared with total DRAQ5 stained cells per field (n = 200–300). Scale bar = 100 μm.
secretory modes depend on cell types, with active Wnt5b-associated exosomes.

Discussion

In this study, we purified Wnt5b for the first time and revealed its biochemical properties and activities. Purified Wnt5b induced Dvl2 phosphorylation, Rac activation, and luciferase activity at comparable levels to purified Wnt5a. At least in vitro, little difference exists between the biochemical activities of purified Wnt5a and Wnt5b; the large difference noted in KO mouse phenotypes may be attributable to their expression patterns in vivo.

The profiles of glycans attached to Wnt5b were determined by ESI-MS/MS. N93 and N305 of Wnt5b, which correspond to N114 and N326 of Wnt5a, were modified with high-mannose-type glycans. At least one of these asparagine residues is conserved in 17 Wnt members, including Wnt3a and Wnt11, among 19 human Wnts. In addition, Wnt5b was modified with a hybrid-type glycan at N291, which corresponds to N312 of Wnt5a. Although the glycosylation sites are unique in Wnt5b and Wnt5b, the role of glycosylation at asparagine residues (N291 of Wnt5b and N312 of Wnt5a) remains to be clarified. Wnt5b was also modified with palmitoleic acid at S223, along with serine residues at the same positions of Wnt3a, Wnt5a, and Wnt11. The lipidation of Wnt5b could allow it to be inserted into the groove of the extracellular domain of Frizzled. Consistent with the results of other Wnts, Wnt5b NQ and Wnt5b S223A, indeed, failed to be secreted.

Evidence has accumulated that cancer cells, along with epithelial cells and neurons, secrete exosomes that originate from MVBs in vitro and in vivo. As tumor-derived exosomes, it has been proposed that they exhibit anti-immune effects to promote tumor aggressiveness and antitumor responses, and to facilitate cancer cell migration to future metastatic sites by educating bone marrow progenitors. Such diverse roles of exosomes could depend on their composition in terms of proteins, lipids, and nucleic acids. Some Wnts were associated with exosomes that are secreted from human and Drosophila cell lines. Fibroblast-derived exosomes are associated with Wnt11 in BCCs and exosome-associated Wnt11 promotes cancer cell metastatic activity through a β-catenin-independent pathway.

Although a definitive protocol to isolate exosomes has not yet been established, one of the acceptable methods is sedimentation by ultracentrifugation at 100 000 g after successive centrifugations at increasing speeds (2000 and 10 000 g). Using this method, we found that the association of Wnts with exosomes varies among cell types, and that Wnt5b is secreted with exosomes from PANC-1 and Caco-2 cells. The
association of Wnt5b with exosomes was confirmed by immunoelectron microscopy.

To examine whether Wnt5b-associated exosomes are active, we developed a reporter assay system for measuring Wnt5a and Wnt5b signaling using CHO cells stably expressing a Ror2-LRP6 chimera and a TOP-flash gene. In this system, when Wnt5a or Wnt5b binds to the extracellular domain of Ror2 of the chimera with endogenous Fz, the signal induces the expression of luciferase through the intracellular domain of LRP6. As PANC-1 cells express high levels of Wnt5b relative to other Wnt ligands, the luciferase activity induced by exosomes could be derived from Wnt5b. Wnt5b-associated exosomes from Caco-2/Wnt5b cells also increased the luciferase activity. However, exosomes prepared from a large amount of CM from WNT5B KO PANC-1 cells still showed slight luciferase activity. Thus, the possibility that other Wnt ligands, which bind to endogenous LRP6 in CHO cells, also activate luciferase expression cannot be excluded. The presence of active Wnt5b-associated exosomes from PANC-1 and Caco-2 cells was also supported by results showing that the exosomes promote cell migration and proliferation of A549 cells. The results also suggest that Wnt5b-associated exosomes act in a paracrine manner. However, exosomes from PANC-1 and Caco-2 cells did not affect their migration and proliferation in an autocrine manner. This might be due to the difference of the sensitivity for Wnt5b-associated exosomes.

Almost all Wnt5b secreted from Caco-2 cells was associated with exosomes. Wnt5b secretion and Wnt5b-dependent proliferation was suppressed by TSG101 knockout. TSG101 belongs to endosomal sorting complex required for transport components and is required for the secretion of Wnt3a with exosomes in HEK293 cells. However, as no individual protein or class of proteins has been shown to be universally required for exosome production in all systems, the trafficking steps that target Wnt to exosomes may vary among cancer cell lines. Further studies are necessary for a full understanding of the actions of Wnt5b and Wnt5b-associated exosome in cancer cells.

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Disclosure Statement

The authors have no conflict of interest.
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Supporting Information

Additional Supporting Information may be found online in the supporting information tab for this article:

**Fig. S1.** Characterization of Wnt5b protein.

**Fig. S2.** Wnt secretion in MDCK cells and expression of Wnt5b in cancer cell lines.

**Fig. S3.** Wnt5b is not involved in Caco-2 cell migration.

**Fig. S4.** TSG101 knockdown suppresses the secretion of Wnt5b-associated exosomes.

**Table S1.** Forward and reverse primers for real-time RT-PCR used in this study.

**Table S2.** Small interfering RNA used in this study.

**Data S1.** Supplementary materials and methods.