Cilia in autophagy and cancer

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Abstract
Cancer cells are distinguished from normal cells by increased proliferation and metabolism, loss of polarity control, and the potential to invade other tissues of the body. As hubs of signaling transduction, primary cilia have been linked to diverse developmental and degenerative disorders. Interestingly, loss of cilia has been observed in multiple malignant tumors, suggesting a potential suppressive role of cilia in cancer development. More recently, emerging studies began to unveil the bidirectional interaction of cilia and autophagy, a basic cellular clearance and recycling mechanism to regulate cell homeostasis. Here, we summarize the interplay between cilia and autophagy and discuss the roles of cilia in both autophagy and cancer.

Keywords: Cilia, Cell cycle, Cilia-related signaling, Autophagy, Cancer

Background
Cilia/flagella are eukaryotic cell organelles protruding from cell surface into environment. Most vertebrate cells assemble a single primary cilium, when they exit from the cell cycle into differentiated or quiescent status [1]. This ∼5 μm tall, microtubule-based protrusion is essential for multiple signaling transductions [2, 3]. Autophagy is a destructive cellular process to degrade disordered cell organelles and protein aggregates, and maintain cellular homeostasis. More recently, cilia are attracting interests as structures having bidirectional interaction with autophagy. This review will discuss the relationship between cilia and autophagy and emphasize the function of cilia in cancer development.

The connections between cilia and cancer
Primary cilia possesses an axoneme consisting of nine doublet microtubules, which is surrounded by a specialized membrane [4]. In interphase, cillum biogenesis is initiated by the attachment of a Golgi-derived membrane vesicle on the distal end of mother centriole [5, 6]. Subsequently, the nucleated axoneme buds from mother centriole and bends the cell membrane to form the structure [5, 6]. Considering that centrosomes direct spindle formation in mitosis, cilia must be disassembled before mitosis to liberate the captive centriole and to promote the formation of spindle [7–10]. The presence of cilia can suppress abnormal cell growth by restricting cell cycle (Fig. 1). Although the ciliary membrane is continuous with cell membrane, the lipid and protein compositions of ciliary membrane are different from cell membrane compositions [11–15]. The specialized ciliary membrane makes cilia capable of transducing multiple cellular signaling [2, 16–21].

Since primary cilia have the ability to influence cell cycle and modulate cilia-related signaling transduction, dysfunction of cilia has long been proposed as a prerequisite step of cancer development [7, 22]. In contrast to normal cells, cilia are lost in multiple cancer types [22]. Clinical data also show that cilia formation is compromised in multiple human cancers including breast cancer, cholangiocarcinoma, melanoma, pancreatic cancer, prostate cancer, and renal cell carcinoma [17, 22–30]. These observations suggest that cilia play a suppressive role in cancer development. Although defective cilia and cancer are always associated, a direct role of cilia in tumorigenesis is still elusive.

The negative correlation between cilia and cell cycle has been discovered for many decades. The studies in Snell’s group provided the first molecular link between cilia disassembly and cell cycle progression [7, 31]. They found that the disassembly of Chlamydomonas cilia requires protein CALK, a member of Aurora kinase family, which promotes cell cycle [31]. In 2007, Golemis's
group optimized a system to study the mechanisms of cilia disassembly [32]. Briefly, cells were treated by serum starvation to induce cilia formation. Serum was added into the medium to trigger cilia disassembly and cell cycle re-entry. Like the observation in Chlamydomonas, they found that mammalian cells also shorten their cilia through Aurora kinase-dependent pathway. Upon serum stimulation, HEF1 activated Aurora A, which phosphorylated HDAC6 to promote cilia disassembly during cell cycle re-entry [32, 33]. According to these findings, a straightforward question will be whether cell cycle progression is blocked with defective cilia disassembly? Tsiokas and colleagues found that knockdown of Nde1, a mother centriole-localized protein, led to elongated cilia in mammalian cells [34]. Interestingly, G0 cells with longer cilia by depletion of Nde1 delayed cell cycle re-entry after serum addition. To confirm the delayed re-entry was caused by cilia, the group co-knockdown Nde1 with either Ift88 or Ift20, two essential genes for ciliogenesis, and found that the inhibitory effect was reversed [34]. Concomitantly, Sung’s group showed that Tctex-1 localizes to transition zone after phosphorylation at Thr94, where it promotes cilia disassembly before S-phase entry [35]. Consistently, depletion of Tctex-1 resulted in delayed cilia disassembly along with delayed cell cycle re-entry in ciliated cells, but not in non-ciliated cells [35]. Both of the studies indicate a suppressive role of cilia in cell cycle progression (Fig. 1), raising the possibility that loss of cilia promotes unrestricted cell cycle progression in cancer cells.

The other important function of primary cilium is its ability to regulate multiple signaling pathways, the dysfunctions of which are associated with a number of cancers [22]. As a cilia-dependent pathway (Fig. 2a), Sonic Hedgehog (Shh) signaling has important functions in guiding embryonic development by regulating cell differentiation and proliferation [20, 36–39]. In the absence of Shh ligand, membrane proteins Patched and Gpr161 are localized to cilia. On the other hand, the most majority of Smoothened is excluded from cilia, though a basal level of protein is thought to be traffic inhibitors of cell cycle progression. Shh promotes cilia disassembly and cell cycle re-entry, while its absence leads to cilia elongation and cell cycle arrest. This antagonistic effect suggests that cilia disassembly is crucial for cell cycle progression, providing a potential therapeutic target for cancer treatment.
through cilia as well [37, 39]. Shh transcriptional factors 
Gli2 and Gli3 are cleaved to Gli2R and Gli3R repressor forms 
and inhibit Shh downstream transcriptions [40]. 
Upon the Shh ligand binding, Patched and Gpr161 are 
moved out from cilia, but Smoothened is accumulated 
in cilia. Consequently, the stabilized Gli2 and Gli3 can 
be activated as Gli2A and Gli3A, which trigger the tran-
scription of Gli1 and other Shh target genes [39, 40]. 
In several types of cancers, the abnormal activations of 
Shh are observed [22, 41–43]. For instance, dysregulated 
activation of Shh contributes to basal cell carcinoma 
and medulloblastoma development [41, 43]. It is inter-
esting that primary cilia are either positive or negative 
regulators of Shh-related oncogenesis, depending on 
the initiating oncogenic mutations [41, 43]. Wnt signal-
ing is critical to animal development and homeostasis 
[44]. Upregulation of Wnt signaling has also been linked 
to tumorigenesis [22, 42, 44]. Although the functions 
of cilia in Wnt signaling are still controversial, it seems 
that both canonical and non-canonical Wnt signaling 
can be regulated by cilia (Fig. 2b, c). Down regulation 
or loss of ciliary proteins, including BBS1, BBS4, Kif3a, 
IFT88, and OFD1, leads to accumulation of β-catenin, 
which subsequently increases the transcription of Wnt 
target genes in Zebrafish embryos and mouse cells and 
embryos [21, 44, 45]. Interestingly, Ift88, Ift172, and 
Kif3a mutant mice, which also lack functional primary 
cilia, failed to show any phenotype caused by upregu-
lated Wnt signaling [46]. Considering that Wnt signaling 
is strictly regulated in specific developmental stage and 
tissue, these results may still reflect the ability of cilia to 
regulate Wnt signaling. In contrast to canonical Wnt, 
non-canonical signaling is β-catenin independent and 
involved in the regulation of cell polarity [47–49]. Cystic 
diseases are well-established models of human diseases 
causd by dysfunctional cilia. Similar with tumor cells, 
a typical symptom of cystic organs is a loss of cell polarity 
[17, 50–52]. Consistent with these observations, pla-
nar cell polarity (PCP) mutant phenotypes, including 
open eyelids and disorganized stereocilia, are found in 
Bbs1, Bbs4, and Bbs6 defective mice [53]. Loss of two 
cilia-related proteins, Ivn/NPHP2 and OFD1, also leads 
to PCP-regulated convergent extension defects in ver-
tebrates [48, 54]. All of these studies demonstrated that 
cilia are capable of regulating both canonical and non-
canonical Wnt signaling [18, 44, 46, 55–57].

Including Shh and Wnt, the list of cilia-related signal-
ings is growing fast. TGF-beta, Hippo, and notch signal-
ing are also connected with cilia [17, 58–61]. In addition 
to the function of receiving and transducing signaling, 
recent studies in Chlamydomonas showed cilia can also 
release signaling active vesicles and act as signaling trans-
mitting organelles to regulate the behavior of other cells

[62–64]. All these studies show a strong link between 
cilia and cancer development. However, if and how cilia 
function in tumorigenesis remains unclear. Tumors are 
highly heterogeneous tissues and consisted of different 
cell types, including tumors cells and tumor-associated 
fibroblasts, endothelial cells, and immune cells [65–69]. 
These cells and extra-cellular matrix constitute tumor 
stroma [66, 69, 70]. Functioning as sensing organelles, 
loss of cilia may alter the signaling network and cell–cell 
communications inside tumor stroma. Involved in mul-
tiple signaling, the function of cilia in tumorigenesis will 
be far more complex than it appears now and will not be 
limited to the cell cycle regulation and polarity control. If 
and how cilia contribute to cancer development remains

an important question to be addressed.

The bidirectional interplay between cilia and autophagy

Autophagy has essential functions in multiple physiologi-
ical processes [71]. The relationship of cilia and autophagy 
has been missing for a long period. A set of recent stud-
ies established the relationship of cilia and autophagy 
(Fig. 3) [72–74]. In contrast to cells in vivo, most in vitro 
cultured cells do not express cilia. Early studies showed 
that serum withdrawal leads to cell cycle exit and induces 
ciliogenesis. Interestingly, serum starvation can also 
trigger autophagy. The most natural question to ask is if

![Fig. 3 Illustrative model of the links between cilia, autophagy, and cancer. Autophagy has dual roles in ciliogenesis by selectively degrading cilia essential protein IFT20 to suppress cilia formation, or degrading suppressive protein OFD1 to promote cilia formation. Inversely, cilia can enhance autophagy through cilia-dependent Shh signaling. Both cilia and autophagy are proposed as regulators for cancer development. The cross-talk between cilia and autophagy may provide new applications for cancer drug discovery](image)
these two concurrent events are related? Tang and colleagues demonstrated that OFD1, a ciliopathy protein, was degraded by autophagy to promote cilia formation upon serum starvation [72, 75]. OFD1 is localized to two cilia-related subcellular structures, distal end of centrioles and centriolar satellites [72, 76–78]. The centriolar OFD1 is thought to maintain centriolar length and integrity, which is required for cilia formation [76]. However, the function of satellite pool was unclear. Tang showed that autophagy largely eliminated the satellite OFD1 but not centriolar OFD1. Inhibition of autophagy attenuated satellite OFD1 degradation and led to lower ciliogenesis rate and shorter cilia. Consistently, depletion of OFD1 by RNA interference dramatically increased cilia formation in murine embryonic cells and restored ciliogenesis in MCF7 cells, a breast cancer cell line originally lacking cilia [72]. All these data demonstrated a suppressive role of satellite OFD1 in cilia formation and suggested a positive role of autophagy in ciliogenesis. In contrast to stimulated autophagy, Cuervo’s group showed that basal level autophagy acts as a negative regulator for ciliogenesis by degrading IFT20, ciliary essential protein [74]. The switch of basal autophagy and stimulated autophagy may potentiate autophagy positively or negatively controlling cilia formation in response to environmental changes [79]. However, a surprising aspect in these two studies is that the ratio of ciliated cells to ciliary length in ATG5−/− autophagy-defective MEF cells are quite different [72, 74]. Given that the confluence of cells has strong influence on cilia formation, one possible explanation of the observations might be attributed to the different status of cell density in these two studies. MTOR is a known negative regulator of autophagy [80]. Wang et al. showed that MTOR activity is upregulated in cilia-suppressed cells, also suggesting that lower autophagy level is associated with attenuated cilia formation [81]. Consistent with the observation, they also showed that upregulating autophagy activity prompted cilia elongation and down-regulating autophagy activity led to shortened cilia in kidney cells [81]. Taken together, these studies demonstrated that autophagy can serve as a dual-role regulator of ciliogenesis by alternatively eliminating ciliary essential protein(s) or its suppressive protein(s) [75, 79]. Understanding of the mechanisms that controls the autophagy switch to turn on/off of cilia formation will be an important question for future studies.

Despite the function of autophagy in ciliogenesis, cilia and cilia-related Shh signaling are potential regulators of autophagy. Several components of autophagic machinery localize around ciliary or periciliary region [74, 79]. ATG16L1, AMBRA1, LC3, GABARAP, and VPS15 staining showed discrete puncta along cilia [74, 79]. ATG14, VPS34, ATG7, and ATG5 are found at the basal body region [74, 79]. Regarding the vesicular activity of ciliary pocket, the presence of autophagic initiating molecules around cilia suggests ciliary area as a possible novel origin of autophagosome formation to activate autophagy. Consistently, lower autophagy activity is observed in IFT20 knockdown MEF cells and IFT88 knockout kidney epithelial cells, both of which have defects in ciliogenesis [74, 79]. Interestingly, the activation of Shh rescued the defective autophagy flux, indicating that cilia upregulate autophagy through cilia-dependent Shh signaling [74, 79]. In another study, cilia-suppressed cells also showed repressed autophagy, which might be resulted from enhanced MTOR activity [81]. In ciliated neuron and smooth muscle cells, autophagy activation was also observed after the upregulation of Shh by its ligand binding [82, 83]. One argument against the positive role of cilia in promoting autophagy comes from the studies of Pancreatic ductal adenocarcinoma (PDAC). PDAC are malignant tumors with high-level autophagy [84]. However, cilia are absent in human and mouse PDAC tissues compared to highly ciliated normal tissues [27]. In other word, cilia loss fails to downregulate autophagy in these malignant cells. Although cilia and cilia-dependent Shh have emerged as possible regulators of autophagy, more insightful mechanisms of the regulating system remain to be elucidated.

**Does the crosstalk between cilia and autophagy influence cancer development?**

The first link between autophagy and cancer is from the studies of Beclin 1, an essential protein for autophagy initiation [85, 86]. Unlike other tumor suppressors, *Beclin 1* is characterized as a haploid-insufficient tumor suppressor gene, monoallelic mutations of which lead to defective function [85, 86]. Interestingly, only premalignant tumors, but not malignant tumors, are observed in autophagy-deficient mice by knockout of *atg5* or *atg7*, two autophagy essential genes, suggesting a suppressive role of autophagy in cell transformation [87]. In contrast to the function of autophagy in transformation, high level of autophagy is required in malignant tumors, including PDAC and non-small cell lung cancer, to maintain the high level of metabolism [84, 88, 89]. These data suggest a dual role of autophagy in cancer development (Fig. 3). One possible explanation is that autophagy plays different roles in cell transformation and transformed cell progression. At the early stage of cancer development, autophagy can degrade harmful factors, including aggregated proteins and aged mitochondria, to prevent cells from accumulating genomic mutations [89–92]. After cancer cell transformation, autophagy can provide substrates for high-level metabolism and prevent toxic product accumulation, both of which promote cancer survival and
and autophagy is emerging as a new field for future studies. Interestingly, the bidirectional interplay between cilia serve as negative regulator of cancer development by providing additional checkpoint of cell cycle progression.

In ciliated cells, the presence of cilia might positively regulate autophagy, preventing metabolic waste accumulation and constitutive cellular damage, which is a potent factor in inducing cancer development [74, 92]. However, in malignant cells, why the loss of cilia and upregulated autophagy are associated together is still poorly understood. Reversely, autophagy has dual roles in ciliogenesis by degrading essential or suppressive cilia-related proteins. If autophagy plays a role in cancer through cilia, how cells modulate the switch to turn on/off cilia expression will be an important point to be addressed. Although the interplay between cilia and autophagy has emerged, recent studies may have just begun to touch a small tip of a giant iceberg. Future studies will hopefully provide more evidences to reveal the complicated connections between cilia and autophagy.

Conclusion
The list of cilia’s functions is growing fast. As discussed above, cilia can restrict mitosis and inhibit abnormal cell proliferation [7, 17, 22]. In this way, theoretically, cilia serve as negative regulator of cancer development by providing additional checkpoint of cell cycle progression. Interestingly, the bidirectional interplay between cilia and autophagy is emerging as a new field for future studies. Autophagy selectively turns on/off cilia formation by alternatively degrading ciliary essential protein, IFT20, or suppressive protein, OFD1. The mechanisms modulating this switch are still unknown. Involved in autophagy initiation, cilia enhance autophagy flux through cilia-related Shh. Whether and how other cilia-related signaling(s) participate(s) autophagy regulation remains unclear. Given the broad functions of cilia and autophagy in the regulation of cell proliferation and metabolism, discovery of drugs specifically targeting these two regulators will provide a wide therapeutic approach for cancer and other diseases.

Abbreviations
Shh: Sonic Hedgehog signaling; IFT: intraflagellar transport; OFD1: oral-facial-digital syndrome 1; PCP: planar cell polarity; PDAC: pancreatic ductal adenocarcinoma.

Authors’ contributions
MC and QZ wrote the paper. Both authors read and approved the final manuscript.

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Competing interests
Both the authors declare that they have no competing interests.

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