PTEN in Chromatin Remodeling

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The tumor suppressor phosphatase and tension homolog (PTEN) is frequently mutated in human cancers, and it functions in multiple ways to safeguard cells from tumorigenesis. In the cytoplasm, PTEN antagonizes the PI3K/AKT pathway and suppresses cellular proliferation and survival. In the nucleus, PTEN is indispensable for the maintenance of genomic stability. In addition, PTEN loss leads to extensive changes in gene expression at the transcriptional level. The linker histone H1, generally considered as a transcriptional repressor, binds to the nucleosome to form a structure named the chromatosome. The dynamics between H1 and chromatin play an important role in determining gene expression. Here, we summarize the current understanding of roles of PTEN in controlling chromatin dynamics and global gene expression, which is crucial function of nuclear PTEN. We will also introduce the recent discovery of the PTEN family members and their functions.

It has been adequately indicated that the tumor suppressor phosphatase and tension homolog (PTEN) antagonizes the PI3K/AKT/mTOR signaling cascade in the cytoplasm, inhibiting cell proliferation and cell survival (Maehama and Dixon 1998; Stambolic et al. 1998). PTEN has long been thought to reside exclusively in the cytoplasm, and the first papers describing its presence in the nuclear compartment were in primary neurons and endothelial cells (Sano et al. 1999), as well as myoepithelial cells of normal breast ducts (Perren et al. 1999). In 2007, Shen et al. reported that PTEN is localized in nucleus and maintains chromosomal integrity (Shen et al. 2007). Since then, functions of nuclear PTEN have been greatly expanded to DNA- and chromatin-related events, including

regulation of genomic stability, DNA replication, DNA repair, and cell-cycle progression (Planchon et al. 2008; Hou et al. 2017). However, molecular mechanisms of how PTEN is shuttled to the nucleus are still not fully understood. Because the PTEN protein lacks either a classical nuclear localization signal (NLS) or nuclear export signal (NES) motif, several mechanisms of its shuttling have been proposed, including simple diffusion (Liu et al. 2005), active shuttling by the RAN GTPase (Gil et al. 2006) or major vault protein (MVP) (Chung et al. 2005), phosphorylation-dependent shuttling (Planchon et al. 2008), and monoubiquitylation-dependent import (Trotman et al. 2007). Recently, SUMOylation has also been reported to control PTEN nuclear localization (Bassi et al. 2013). Nuclear

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PTEN plays an important role in tumor suppressing, and the absence of nuclear PTEN is associated with increased aggressiveness in cancers (Gimm et al. 2000; Baker 2007). Therefore, it is of great importance that we summarize the latest research progress on functions of nuclear PTEN, especially the interplay with histones on chromatin and gene expression resulting in biological and pathological activities.

DYNAMICS OF CHROMATIN STRUCTURE

In eukaryotes, the 2 m of genomic DNA is compactly intertwined and bound to histone proteins, and together they form a fibrous architecture named “chromatin” (Kornberg 1974; Kornberg and Thomas 1974). The fundamental unit of chromatin is named the nucleosome, which consists of ~147 bp of DNA wound around an octamer of core histones (two copies each of H2A,

H2B, H3, and H4) and an additional variable length of linker DNA (Arents and Moudrianakis 1993; Luger et al. 1997). Further packaging of DNA involves a linker histone (H1) bound to the nucleosome with ~10 bp of DNA at the entry/exit sites of the nucleosome core particle, generating a second-level structure called the chromatosome (Simpson 1978). Chromatin can be either densely packed in the form of heterochromatin, which is largely inaccessible to gene transcription and hence harboring inactive genes, or as open and accessible euchromatin, which contains numerous active genes (Fig. 1A). This dynamic, tightly controlled regulation of chromatin structure ensures timely, coordinated, and appropriate gene expression. Chromatin can be regulated by various means, including modifications of DNA (Schübeler 2015), modifications of histones (Tessarz and Kouzarides 2014), and architectural remodeling by protein

Figure 1. Multiple levels of chromatin assembly. (A) The overlong DNA is highly compacted through association with core histones to form nucleosomes, and then chromatosomes by histone H1 binding. Several aspects show a great impact on chromatin dynamics (interconversion of euchromatin and heterochromatin), such as modifications of DNA, modifications of histones, and other protein complexes that interact with chromatin. (B) Structural domains of the linker histone H1. Although the globular domain (GH1) is the only folded portion of the protein in solution, all three domains are subjected to multiple posttranslational modifications (PTMs). (NTD) amino-terminal domain, (CTD) carboxy-terminal domain.

complexes (Clapier and Cairns 2009). These mechanisms function individually and in concert to modulate genome-wide topology and gene expression, thereby regulating cell differentiation, cell division, and tissue and organism development.

ROLE OF THE LINKER HISTONE H1 IN GENE EXPRESSION AND CHROMATIN DYNAMICS

It is believed that the general role of core histones is to condense DNA and reduce its accessibility, resulting in the inhibition of transcription in vitro (Lorch et al. 1987). However, in vivo studies have revealed that the transcription regulation is largely dependent on the posttranslational modification (PTM) of core histones and have identified specific PTMs of core histones that are related to either transcription activation or repression (Eberharter and Becker 2002; Allis and Jenuwein 2016; Izzo and Schneider 2016). Proteins that add, recognize, and remove these PTMs have also been identified and termed writers, readers, and erasers, respectively (Torres and Fujimori 2015).

In contrast to the knowledge about core histones, much less is known about linker histones in regulation of chromatin structure and function. Histone H1 binds to nucleosomes and shows a dynamic binding affinity for chromatin (Bednar et al. 1998), which allows the modulation of chromatin architecture and gene transcription. Based on findings in *Drosophila*, *Tetrahymena*, and other model organisms, it is appreciated that histone H1 is generally a transcriptional repressor (Ura et al. 1997), and studies of its structure and PTMs have provided hints of how it achieves this function.

Histone H1 species consist of three domains: the central globular domain (GH1, ~80 amino acids), the short amino-terminal domain (NTD, ~35 amino acids), and the carboxy-terminal domain (CTD, ~100 amino acids) (Fig. 1B; Allan et al. 1980). The GH1 domain is the only folded portion of the entire protein in free solution and is proposed to bind symmetrically to DNA on the dyad axis (Ramakrishnan et al. 1993). The NTD is disordered in free solution and does not greatly affect chromatin condensation but is

subject to PTMs that play a regulatory role in H1 function (Allan et al. 1986). The CTD is also disordered in free solution and is assumed to play an important part in chromatin condensation (Hendzel et al. 2004).

The linker histone H1 undergoes various PTMs, including phosphorylation, acetylation, methylation, and ubiquitination. In addition, H1 can undergo various other PTMs, such as citrullination, formylation, denitration, ADP ribosylation, carbonylation, and crotonylation (Fig. 1B; Izzo and Schneider 2016). More than 150 histone-modifying proteins have been identified and their dysregulation can result in inappropriate activation or inactivation of genes (Valencia and Kadoch 2019).

As with core histones H2A and H3, there are also multiple variants of H1 histones in mammalian cells. Although linker histone subtypes are redundant proteins in knockout mice (Fan et al. 2003), suggesting no functional distinction between subtypes, different tissues are characterized by specific subtypes, and transitions in subtypes are observed during embryonic development (Terme et al. 2011; Li et al. 2012). H1.1 to H1.5 and the testis-specific H1t in humans are expressed in a replication-dependent (RD) manner, meaning that these H1 subtypes are synthesized only during S phase, whereas H1.0 and H1x are replication-independent and can be expressed in nonproliferating cells (Fyodorov et al. 2018). Some functional distinctions between the RD subtypes have been observed in gene expression regulation (Alami et al. 2003), and a specific contribution of H1 subtypes to other nuclear events, such as DNA methylation (Fan et al. 2005) and cell-cycle disruption (Sancho et al. 2008), have been shown.

Fluorescence recovery after photobleaching/fluorescence loss in photobleaching (FRAP/FLIP) experiments show that a large proportion of the H1 exchanges rapidly between sites on euchromatin with an average residence time of ~4 min (Lever et al. 2000). This pattern is very different from that of core histones, for which almost no exchange is observed over the same period of time. Therefore, the conditions for generating a stable in vitro H1-containing chromatosome have not been achieved, and an

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understanding of how H1 is bound to the nucleosome and how chromatin structures are controlled to facilitate gene expression is greatly needed (Crane-Robinson 2016).

Multiple Localizations and Functions of PTEN Family Members

For more than 15 years since its discovery, *PTEN* was thought to be a nonredundant tumor suppressor gene with no variants or isoforms in humans (Keniry and Parsons 2008). Although it is well understood that germline mutations of *PTEN* induce cancer predisposition in various organs, *PTEN* dysfunction could also be an important factor in other diseases, such as diabetes and nervous system diseases (Worby and Dixon 2014). It has long been thought that *PTEN* is so multifunctional that this 403-amino acid (aa) protein plays a role almost everywhere. However, this notion was challenged when a new isoform of *PTEN*, *PTEN*-Long or *PTEN* α , was independently discovered by two groups (Hopkins et al. 2013; Liang et al. 2014). In these studies, the new *PTEN* isoform was found to use a noncanonical translation initiation codon CUG to start translation on the canonical *PTEN* messenger RNA (mRNA), resulting in a 173-aa amino-terminal extended protein. These groups found that with the amino-terminal extended sequences, the localization of the protein was changed; it could be secreted from cells and re-enter adjacent cells to suppress cell proliferation (Hopkins et al. 2013), or it could target mitochondria within cells and regulate energy metabolism (Liang et al. 2014). Soon after that discovery, in 2017, a third *PTEN* family mem-

ber, *PTEN* β , was identified. Similar to *PTEN* α , *PTEN* β used another noncanonical translational initiation codon AUU to start translation and resulted in a 146-aa amino-terminal extended protein that localized predominantly in the nucleolus and regulated pre-ribosomal RNA (rRNA) synthesis (Fig. 2; Liang et al. 2017).

ROLE OF PTEN IN DNA REPLICATION AND REPAIR

PTEN family members can be found in various locations within cells, and both *PTEN* and *PTEN* β show nuclear localization. However, the subnuclear localization of these two proteins is quite different. *PTEN* β is specifically localized in the nucleolus, whereas *PTEN* is mostly localized in the matrix. In the nucleus, *PTEN* is highly interactive with DNA-binding proteins and histones, and is crucial to the processes of DNA replication, DNA repair, and DNA transcription. In 2007, Shen and colleagues reported the intranuclear localization of *PTEN* and showed its colocalization with centromeres in mouse embryonic fibroblasts (MEFs). These researchers found that within the nucleus, *PTEN* physically interacts with centromere-specific binding protein C (CENP-C) to maintain chromosomal stability (Shen et al. 2007). *PTEN* is also involved in pathways of DNA double-strand break (DSB) repair, which is accomplished by homologous recombination (HR) induced by Rad51. *PTEN* also plays an indispensable role in DNA replication via association with many replication-related proteins, such as replication protein A1 (RPA1) on DNA replication forks (Wang et al. 2015) and minichromosome maintenance

Figure 2. Structural and functional summary of the phosphatase and tension homolog (*PTEN*) family proteins. *PTEN* α and *PTEN* β have a 173-aa and 146-aa extension, respectively, at the amino terminus of regular *PTEN*. The differences in amino terminus of the *PTEN* family decides each protein's subcellular location.

complex component 2 (MCM2) (Feng et al. 2015) on replication stress. Replication fork–localized PTEN can physically interact with Rad51 following replication stress for the recovery of stalled forks (Fig. 3; He et al. 2015). Therefore, PTEN guards the genome by controlling multiple processes of chromosome inheritance.

PTEN INTERACTS WITH HISTONE H1 AND CONTROLS CHROMATIN CONDENSATION

Several lines of evidence have shown that the PTEN status has an influence on global gene expression (Matsushima-Nishiu et al. 2001; Li et al. 2006), but the molecular mechanisms were not well understood. More recently, Chen et al. discovered an interplay between PTEN and the linker histone H1 as potential means of PTEN

impact on gene expression (Chen et al. 2014). In this study, PTEN loss in MEFs leads to cumulative errors of heterochromatin protein 1 α (HP1 α) foci formation, indicating impaired structural organization of chromatin in the nucleus. Further investigation using FLAG-hemagglutinin (FH) tandem affinity purification with ectopic PTEN followed by SDS-PAGE and mass spectrometry analysis identified several subtypes of histone H1 as PTEN-interacting proteins and indicated that the C2 domain of PTEN and the CTD of H1 are responsible for the interaction. Because the CTD of histone H1 accounts for much of its binding affinity to chromatin, PTEN loss leads to histone H1 dissociation from chromatin and chromatin decondensation. Moreover, PTEN and histone H1 were found to act cooperatively to repress

Figure 3. Role of phosphatase and tension homolog (PTEN) in DNA replication and repair. Nuclear PTEN safeguards the genome in multiple ways. PTEN is involved in maintenance of centromere stability, DNA replication progression, and DNA repair via homologous recombination through interaction with or regulation of relative proteins, either dependent or independent of its protein kinase activity. (ssDNA) single-stranded DNA.

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histone H4 acetylation at lysine 16, which is an epigenetic marker for chromatin activation. Microarray analysis of Pten-deficient MEFs revealed a number of up-regulated genes, including the oncogenes Kras, Braf, and Akt1; the survival factors Bcl2 and Nfkb2; the estrogen receptor Esr1; and the tumor invasion factor Cd44 (Fig. 4A). This mechanism appears to be a part of the cellular machinery that controls global gene expression profiles through regulating DNA accessibility and hierarchical levels of chromatin organization (Chen et al. 2014).

NUCLEAR PTEN BINDS TO HP1 α AND MAINTAINS HETEROCHROMATIN STRUCTURE

In 2015, a study by Gong and colleagues revealed that PTEN could bind directly to heterochromatin and to the resident heterochromatin protein HP1 α . In PTEN knockout in cells, there was a significant reduction in histone H3 trimethyl Lys 9 (H3K9me3) foci intensity, a hallmark of chromatin decondensation. Consistent with this, Gong and colleagues found, using quantitative real-time PCR (RT-qPCR), overexpression of satellite DNA following PTEN loss in both mouse and human cells (Gong et al. 2015). This research, together with Chen's study, reveals the existence of a complex regulatory network based on the relationship between PTEN, histones, and heterochromatin status, which is integral to the control of gene expression (Fig. 4A).

PTEN REGULATES GLIOBLASTOMA ONCOGENESIS THROUGH CHROMATIN-ASSOCIATED COMPLEX OF DAXX AND HISTONE H3.3

In addition to its association with the linker histone and HP1 α , PTEN was recently shown to form a complex with the histone chaperone DAXX and the core histone H3.3 (Benitez et al. 2017). DAXX binds to the unfolded PTEN-hinge domain (amino acids 186–202), whereas PTEN binds to the histone-binding domain of DAXX. The DAXX-H3.3 interaction influences gene expression in several different

ways (Puto and Reed 2008; Wethkamp and Klempnauer 2009; Wong et al. 2009; Goldberg et al. 2010). Using RT-qPCR and anti-H3.3 chromatin immunoprecipitation (ChIP) approaches, four genes with an inverse correlation between gene expression and H3.3 enrichment in PTEN-deficient cells compared with that in PTEN-WT cells were identified (CCND1, MYC, FOS, and BCL2) (Benitez et al. 2017), suggesting that PTEN represses oncogene expression by recruiting H3.3 to chromatin. Additionally, DAXX inhibition in glioblastoma patient-derived glioma neurosphere (GBM-PDX) cultures suppressed tumor growth and increases survival, specifically in a PTEN-deficient background (Benitez et al. 2017). Therefore, the PTEN-DAXX-H3.3 complex regulates gene transcription and can be a new therapeutic target to revert tumorigenesis caused by the loss of PTEN function in GBM (Fig. 4B).

CONCLUDING REMARKS

Nuclear PTEN maintains genomic stability through multiple mechanisms. PTEN-null cells show chromatin instability (CIN) and have many manifestations of CIN, including centromere breakage (Shen et al. 2007), spontaneous elevation of replication stress (Wang et al. 2015), failure of stalled fork recovery (Feng et al. 2015; He et al. 2015), checkpoint dysfunction throughout all phases of the cell cycle (Weng et al. 1999; Puc et al. 2005), genome-wide copy number alterations (Sun et al. 2014), and global chromatin decondensation (Chen et al. 2014; Gong et al. 2015). The interplay of PTEN with histone H1, HP1 α , H3.3, and heterochromatin sheds light on how this powerful tumor suppressor influences expression of thousands of genes, as well as the extent of chromatin condensation. Despite this knowledge, much remains to be elucidated regarding the structural basis behind these interactions. Although the structure of the nucleosome core particle at atomic resolution was solved more than a decade ago (Richmond and Davey 2003), determination of the structures of the chromatin-containing linker histones has been very challenging. However, important progress has been made in resolving

Figure 4. Interplay between phosphatase and tension homolog (PTEN) and histones. (A) Interplay between PTEN, H1, and heterochromatin protein 1α (HP1α). PTEN forms a complex with H1 and HP1α on chromatin, which condenses the genome and represses oncogenic gene transcription. When PTEN is depleted in cells, H1 and HP1α fall off the chromatin, leading to acetylation of H4 at K16 and chromatin decondensation, which causes oncogenic gene transcription. (B) Interplay between PTEN and H3.3. PTEN interacts with H3.3 and its chaperone DAXX, and anchors the complex on the chromatin to repress oncogenic gene transcription. On PTEN deletion, the DAXX-H3.3 complex dissociates the chromatin, resulting in gene activation. DAXX inhibition suppresses tumor growth through global H3.3 distribution changes on chromatin leading to up-regulation of tumor suppressor genes and down-regulation of oncogenes. (mRNA) messenger RNA.

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higher-order chromatin structures, including the crystal structure of the nucleosome-linker histone complex at near-atomic resolution (Zhou et al. 2015), the cryo-electron microscopic (cryo-EM) structure of the 30 nm chromatin fiber (Song et al. 2014), and the nuclear positioning of topologically associating domains (TADs) (Geeven et al. 2015), revealing detailed interactions between the linker histone and DNA in the nucleosome. The nucleosome forms a transcriptional barrier to RNA polymerase II (RNAPII) progression, resulting in transcriptional inhibition. A very recent study solved the cryo-EM structure of nucleosome-transcribing RNAPII elongation complexes (ECs), revealing that transcription elongation factors Elf1 and Spt4/5 reshape the downstream edge of the EC and intervene between RNAPII and the nucleosome. These mediators facilitate RNAPII progression through superhelical location SHL(-1) and suppress RNAPII pausing at SHL(-5), providing a platform for various chromatin activities (Ehara et al. 2019). In addition, “assay for transposase-accessible chromatin using sequencing” (ATAC-seq) has recently emerged as one of the most powerful techniques for studying genome-wide chromatin accessibility profiling (Buenrostro et al. 2013). By using this approach to study open chromatin under the regulation of PTEN and histones, we will better understand how PTEN regulates gene expression at the chromatin level. In summary, combining our increasing knowledge of the structure and principles of histone and nucleosome organization and techniques to evaluate whole genome-wide open chromatin regions, the specific role of PTEN in this elaborate and complex process will be revealed and understood.

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