Capn3 depletion causes Chk1 and Wee1 accumulation and disrupts synchronization of cell cycle reentry during liver regeneration after partial hepatectomy

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Abstract
Recovery of liver mass to a healthy liver donor by compensatory regeneration after partial hepatectomy (PH) is a prerequisite for liver transplantation. Synchronized cell cycle reentry of the existing hepatocytes after PH is seemingly a hallmark of liver compensatory regeneration. Although the molecular control of the PH-triggered cell cycle reentry has been extensively studied, little is known about how the synchronization is achieved after PH. The nucleolus-localized protein cleavage complex formed by the nucleolar protein Digestive-organ expansion factor (Def) and cysteine proteinase Calpain 3 (Capn3) has been implicated to control wounding healing during liver regeneration through selectively cleaving the tumor suppressor p53 in the nucleolus. However, whether the Def-Capn3 complex participates in regulating the synchronization of cell cycle reentry after PH is unknown. In this report, we generated a zebrafish capn3b null mutant (capn3bΔ19Δ14). The homozygous mutant was viable and fertile, but suffered from a delayed liver regeneration after PH. Mass spectrometry (MS) analysis of nuclear proteins revealed that a number of negative regulators of cell cycle are accumulated in the capn3bΔ19Δ14 liver after PH. Moreover, we demonstrated that Check-point kinase 1 (Chk1) and Wee1, two key negative regulators of G2 to M transition, are substrates of Capn3. We also demonstrated that Chk1 and Wee1 were abnormally accumulated in the nucleoli of amputated capn3bΔ19Δ14 liver. In conclusion, our findings suggest that the nucleolar-localized Def-Capn3 complex acts as a novel regulatory pathway for the synchronization of cell cycle reentry, at least partially, through inactivating Chk1 and Wee1 during liver regeneration after PH.

Keywords: Def, Capn3, Chk1, Wee1, Nucleolus, Cell cycle, Liver regeneration, Partial hepatectomy, Zebrafish

Background
Liver regeneration after PH is an outstanding model to study the synchronization of cell cycle reentry and progression of cell cycle because a seemingly synchronized hepatocyte proliferation is induced after PH (Mao et al. 2014; Kan et al. 2009; Goessling et al. 2008; Michalopoulos 2007; Fausto et al. 2006). Previous reports have shown that in mice, DNA replication peaks at certain time point, and the reentry into mitosis is further controlled by circadian clock, probably through G2-M check point protein kinase Wee1, a downstream effector of Chk1, thus ensures the synchronization of cell cycle progression (Schibler 2003; Matsuo et al. 2003). However, the exact regulatory mechanism of cell cycle synchronization is not well established.

Nucleolus is the subcellular organelle primarily responsible for ribosome synthesis in an eukaryotic cell
(Boisvert et al. 2007; Wang et al. 2012). However, studies have shown that 70% of the nucleolar proteins are unrelated to ribosome biogenesis (Ahmad et al. 2009). These proteins and even some of those involved in ribosome production have been shown to participate in cellular processes such as cell cycle regulation, DNA repair, cell senescence and apoptosis (Diesch et al. 2014). Moreover, increasing evidence has demonstrated that certain proteins tend to accumulate in the nucleoli under stress conditions, indicating the importance of nucleolus in regulating protein homeostasis (Frottin et al. 2019). As an evidence supporting this point, we recently demonstrated that the nucleolar protein Def recruits cysteine protease Capn3 (Capn3b in zebrafish) into the nucleolus to form the Def-Capn3 complex. This complex specifically mediates the cleavage of its downstream substrates including the tumor suppressor p53 to regulate cell cycle progression and liver development (Tao et al. 2013; Guan et al. 2016). Therefore, the turnover of p53 is not only controlled by the proteasome-dependent pathway (Tai and Benchimol 2009) but also a proteasome-independent pathway mediated by the Def-Capn3 complex (Tao et al. 2013; Guan et al. 2016). Interestingly, Def-Capn3 also cleaves the ribosome biogenesis factor Mpp10 to regulate the small subunit assembly (Zhao et al. 2019). However, whether the Def-Capn3 complex regulates cell cycle progression by targeting other key factors is still largely unknown.

We previously reported that def<sup>−/−</sup> mutant liver formed a scar at the amputation site after PH due to the activation of constitutive inflammatory response mediated by p53 and its downstream gene HMGB1 (Zhu et al. 2014). Over-inflammatory response activates the TGF-β signaling which finally leads to fibrosis at the wounding site (Zhu et al. 2014). However, whether the Def-Capn3 complex participates in regulating the cell cycle synchronization after PH is unknown. In this report, based on studying a zebrafish capn3b null mutant (capn3b<sup>A196A114</sup>), we found that the capn3b<sup>A196A114</sup> mutant suffered from a delayed liver regeneration due to disruption of the synchronized cell proliferation after PH. Proteomics data revealed that some negative regulators of cell cycle are accumulated in the capn3b<sup>A196A114</sup> liver after PH. Molecular analysis showed that Chk1 and Wee1 are abnormally accumulated in the nucleoli of the amputated capn3b<sup>A196A114</sup> liver. Biochemical study demonstrated that Chk1 and Wee1 are the substrates of Capn3. These data suggest that the Def-Capn3 complex plays an important role in regulating liver regeneration after PH.

**Methods**

**Zebrafish lines and maintenance**

Zebrafish wild-type (WT) and all relevant mutants used in this study were in the AB background. The zebrafish def<sup>−/−</sup> (def<sup>Δ129</sup>) mutant line was as previously described (Chen et al. 2005). capn3b<sup>A19</sup> mutant was generated using transcription activator-like effector nucleases (TALEN) technique with strand sequences capn3b-TAL(R) GGCGAGAGAAA CAGAAAT and capn3b-TAL(L) AACTCAGAAGCT GCTCCGC. In the capn3b<sup>Δ19</sup> genetic background, CRISPR-Cas9 technology was adopted to generate the capn3b<sup>Δ19Δ14</sup> mutant using a specific gRNA (5′-GAGGTATGTCATCC TGAAGAGG-3′) against the 3rd exon of the capn3b gene. Fish were raised and maintained according to the standard procedure recommended at [http://zfin.org/](http://zfin.org/).

**Plasmid construction, mRNA synthesis, mRNA injection and western blot analysis**

The construction of the HA-p53R143H and Myc-def plasmids has been described previously (Tao et al. 2013). The chk1 and wee1 full length cDNAs derived from the total RNA extracted from 12 h post-fertilization (hpf) zebrafish embryos were cloned into the pcDNA3 vector with an HA-tag, respectively, using primers as listed in Table S1. HA-chk1<sup>A</sup> and HA-wee1<sup>S44A</sup> was constructed through site-directed mutagenesis using the primer pairs listed in Table S1 (Watanabe et al. 2004a). For mRNA injection, mRNAs were synthesized in vitro using the mMESSAGE mMACHINE Kit (Ambion). The methods for protein extraction from zebrafish embryos and western blot analysis were as described previously (Chen et al. 2005; Gong et al. 2015).

**Whole-mount RNA in situ hybridization (WISH) and liver size measurement**

For WISH, Digoxigenin (DIG) (Roche Diagnostics) was used to label the liver fatty acid binding protein (fabp10a) and trypsin probes as previously described (Zhao et al. 2019). WISH performance and liver size measurement were performed as described previously (Chen et al. 2005; Lo et al. 2003).

**PH and measurement of the liver versus body ratio (LBR)**

Fish of different ages were used for PH. The PH and measurement of the LBR were as described previously (Kan et al. 2009). The survival rates of the wild-type (WT) and capn3b<sup>A19Δ14</sup> fish after surgery were greater than 95% after PH in all of the experiments described in this report.

**Cryosectioning and immunofluorescent staining**

Zebrafish cryosections were prepared as described previously (Guan et al. 2016). Antibodies against Fibrillarin (Abcam, ab4566, 1:600), zebrafish Bhat (1:200) (Gao et al. 2018), Capn3b(1:200) (mouse monoclonal antibody, generated by Abmart), Chk1 and Wee1 (1:200) (rabbit polyclonal antibodies, generated by Hangzhou Hua-An Biotechnology Company) were used for
Fig. 1 (See legend on next page.)
immunostaining. After tissue sectioning and storing at –80 °C, the cryosections were rehydrated with two washes of PBS-triton (PBS plus 0.2% Triton X-100) for 5 min (each). For the staining of nucleolar proteins, cryosections were incubated in 0.01 M Sodium Citrate for 5 min at 95-1000C. After cooling down to room temperature, cryosections were washed twice with PBB-triton (PBS-triton plus 0.5% BSA). The sections were then blocked by 20% goat serum in PBB-triton for 1 h (hr) at room temperature. After a brief wash with PBB, the sections were incubated overnight with primary antibody at the desired concentration diluted in PBB-triton 4 °C. After 3 washes with PBB-triton, sections were incubated with secondary antibodies (1:500) and DAPI (1:800) in PBB-triton for 1 h. After 3 washes with PBB-triton, the sections were finally mounted in 80% glycerol and covered with cover-slip for image acquisition under a confocal microscope (Olympus FV1000).

EdU incorporation assay
EdU (5-ethyl-2′-deoxyuridine, 1 nl, 10 mM) was injected into the abdominal cavity of an adult fish (0.5 mg EdU/0.1 g body weight). 24 h after injection, the fish were fixed in 4% PFA for 12 h before proceeding to cryosectioning. Incorporated EdU was detected by Alexa Fluor 488 Azide (Life Technologies, A10266) under a confocal microscope (Olympus FV1000).

Nuclei isolation and nuclear protein extraction
Zebrafish liver dissection and nuclei isolation was as described (Guan et al. 2016; Zhao et al. 2019). The liver tissue of zebrafish was carefully dissected under a pose microscope and put into pre-cooled low permeability solution buffer A (10 mM Hepes, pH 7.9, 10 mM KCl, 1.5 mM MgCl2, 0.5 mM DTT) for 20 min on ice. The liver was then loaded to a 40 μmcell strainer (Sangon Biotech F613461) for removing remaining tissue mass. The collected flow through solution mainly contained nuclei (approximately 90%) judged from examining the solution under a microscope (Olympus U-tv.063XC). The isolated nuclei were centrifuged at 2000 g for 10 min at 4 °C and were ruptured by sonication for fifteen 5-seconcds (5-s) bursts (with 15-s intervals between bursts) at 40% amplitude in IP lysis buffer (containing ‘cell lysis buffer for Western and IP’ and 1 x cOmplete) containing 1.5% SDS. After centrifugation at 12000 g for 15 min, the supernatant was collected and boiled at 100 °C for 10 min and then subjected to mass spectrometry (MS) analysis.

Protein digestion, TMT-labelling, RP-HPLC, LC-MS and data analysis
For MS samples preparation, the lysates (200 μg protein for each sample) were reduced with 100 mM DTT at 37 °C for 1 h. After reduced, the lysates were transferred to the Microcon YM-30 centrifugal filter units ((EMD Millipore Corporation, Billerica, MA), and replaced with 200ul UA (8 M Urea, 100 mM Tris. Cl pH 8.5) twice. After buffer replaced, the proteins were alkylated with 55 mM iodoacetamide (IAA, Sigma-Aldrich, Saint Louis, MO), and digested with sequencing grade trypsin (Promega, Madison, WI) (1:50 (w: w)) at 37 °C overnight. The resultant tryptic peptides were labeled with acetonitrile-dissolved TMT reagents (Thermo Scientific, Rockford, IL) by incubation at room temperature in dark for 2 h. The labeling reaction was stopped by 5% hydroxylamine, and equal amount of labeled samples were mixed together. Offline basic RP-HPLC was performed using a Waters e2695 separations HPLC system coupled with a phenomenex gemini-NX 5u C18 column (250 × 3.0 mm, 110 Å) (Torrance, CA, USA). The sample was separated with a 97 min basic RP-LC gradient as previously described. A flow rate of 0.4 mL/min was used for the entire LC separation. The separated samples were collected into 10 fractions, and completely dried with a SpeedVac concentrator and stored at –
Fig. 2 (See legend on next page.)
20 °C for further analysis. After desalted by StageTip, the peptides were re-suspended in 0.1% formic acid (FA) and analyzed by a LTQ Orbitrap Elite mass spectrometer (Thermo Scientific, Rockford, IL, Waltham, MA) coupled online to an Easy-nLC 1000 in the data-dependent mode. The LC was run with mobile phases containing buffer A (0.1% FA) and buffer B (100% ACN, 0.1% FA). The peptides were separated by a capillary analytic column (length: 25 cm, inner diameter: 150 μm) packed with C18 particles (diameter: 1.9 μm) in a 90-min non-linear gradient (3%–8% B for 10 min, 8%–20% B for 60 min, 20%–30% B for 8 min, 30%–100% B for 2 min, and 100% B for 10 min) with a flow rate of 600 nl/min. The positive ion mode was used for MS measurements, and the spectra were acquired across the mass range of 300–1800 m/z. Higher-energy collisional dissociation (HCD) was used to fragment the fifteen most intense ions from each MS scan.

For analyzing the LC-MS data, the database search was performed for all raw MS files using the software MaxQuant (version 1.6). The Danio rerio proteome sequence database downloaded from Uniprot was applied to searching the data. The parameters used for the database search were set up as follows: The type of search: MS2; The protease used for protein digestion: trypsin; The type of isobaric labels: 6-plex TMT; The minimum reporter parent ion interference (PIF): 0.75; The minimum score for unmodified peptides: 15. Default values were used for all other parameters. Only proteins with no less than two quantified peptides were used for further analysis. DAVID Bioinformatics Database (v6.7) was used to identify proteins for most enriched GO term.

Statistical analysis
For statistic analysis, comparisons were made using the Student’s t-test assuming a two-tailed distribution, with significance being defined as \( p < 0.05 \) (*), \( p < 0.01 \) (**), and \( p < 0.001 \) (***) no significance (NS).

Results
capn3b mutant is defective in liver development under stress conditions
The transcript of the capn3b\(^{Δ19}\) mutant gene (harboring a 19 bp deletion in the exon1) (Fig. 1a) is predicted to encode a 30 amino acids (aa) long peptide containing the N-terminal 8aa of Capn3b and additional 22aa (Fig. 1b). However, there is another ATG after the capn3b\(^{Δ19}\) mutation which might serve as an alternative translation start codon to encode a Capn3b variant with truncation of the N-terminal 93aa (Figure S1B). To avoid the production of this variant (Tao et al. 2013; Sorimachi et al. 1993; Ono et al. 2016), we designed a gRNA targeting the exon3 in the background of capn3b\(^{Δ19}\). The new mutant now carried a 14 bp-deletion in the exon3 in addition to the 19 bp-deletion in the exon1 (designated as capn3b\(^{Δ19Δ14}\) ) (Fig. 1a). The 14 bp-deletion in capn3b\(^{Δ19Δ14}\) disrupted the putative variant (Fig. 1b), (Zhao et al. 2019). Protein analysis showed that both 5dpf-old capn3b\(^{Δ19}\) and capn3b\(^{Δ19Δ14}\) embryos lacked Capn3b (Fig. 1c), (Zhao et al. 2019). In mouse and human, Capn3 is mainly expressed in the muscle (Kramerova et al. 2008; Ono et al. 2006; Richard et al. 1995). Immunostaining of Capn3b protein showed that zebrafish Capn3b was highly expressed in the muscle and was also detected in other organs/tissues in WT. As expected, Capn3b was undetectable in the capn3b\(^{Δ19Δ14}\) embryos at 5dpf (Fig. 1d), demonstrating that capn3b\(^{Δ19Δ14}\) is a null mutation.

Similar to capn3 knockout (KO) mice (Kramerova et al. 2004), capn3b\(^{Δ19Δ14}\) mutant were viable and fertile. We compared the liver development between WT and capn3b\(^{Δ19Δ14}\) in different rearing density by WISH using the liver marker fabp10a (Ribas et al. 2017). When grown at a lower density (60 embryos/9 cm-diameter dish), the size of the capn3b\(^{Δ19Δ14}\) liver was significantly smaller than that of WT at both 3dpf and 5dpf (Fig. 2a, left panel). Surprisingly, when grown at a higher density (120 embryos/9 cm-diameter dish), the size of the capn3b\(^{Δ19Δ14}\) liver was significantly larger than that of WT at both 3dpf and 5dpf (Fig. 2a, right panel). Interestingly, it appeared that the development of the WT liver but not the capn3b\(^{Δ19Δ14}\) liver was more sensitive to the condition of higher rearing density (Fig. 2b).

Next, we grew WT and capn3b\(^{Δ19Δ14}\) zebrafish embryos under a higher temperature (34.5 °C) from 12hpf. WISH using fabp10a and exocrine pancreatic marker trypsin showed that the development of both liver and exocrine pancreas in the capn3b\(^{Δ19Δ14}\) embryos were severely retarded at 2dpf, however, were surprisingly recovered at 3dpf (Fig. 2c). Moreover, at 3dpf, high
Fig. 3 (See legend on next page.)
Fig. 3 Delayed hepatocyte proliferation in capn3bΔ19Δ14 at the resection site after PH. a, b Comparison of LBRs at 3-, 7- and 30-dpH in WT and capn3bΔ19Δ14 fish operated at the age of 8-months-old (a) or 16-months-old (b). Sham control: cut through the abdominal skin only. c-f Co-staining of EdU and Bhmt (liver marker) at the wounding site of the liver in WT and capn3bΔ19Δ14 fish at 3dpH (c) and 7dpH (e). Red dashed line: outlining the cutting site. DAPI: staining nuclei. Scale bar: 50 μm. Corresponding statistical data of the EdU-positive cells were provided (d, 3dpH; f, 7dpH). EdU was injected 24 h before harvesting at 3dpH or 7dpH. Three sections per fish and total three fish were evaluated for each genotype in each case. Student’s T-test for statistical analyses, *, p < 0.05; **, p < 0.01

Fig. 4 Delayed hepatocyte proliferation in capn3bΔ19Δ14 in the deeper region after PH. a, b Co-staining of EdU and Bhmt (hepatocyte marker) for analyzing proliferating hepatocytes in the deeper region away from the amputated site in WT and capn3bΔ19Δ14 at 3dpH (a) and 7dpH (b). DAPI: staining nuclei
temperature-treated $capn3b^{Δ19Δ14}$ embryos had curved body (Fig. 2d), which is likely associated with the role of Capn3 in the muscle (Richard et al. 1995; Kramerova et al. 2004).

capn3b mutant exhibits shortened body length and alteration in the liver to body ratio

We noticed that the $capn3b^{Δ19Δ14}$ adult fish is smaller than the WT control fish. Indeed, measurement of the
**A** Venn diagram for down regulated proteins

- Sham Unique: 63
- PH Unique: 155
- Shared: 29

**B** Most enriched GO term for biological process

- Oxidation reduction process
- Lipid metabolic process
- Cellular response to estrogen stimulus
- Fatty acid biosynthetic process
- Metabolic process
- Steroid biosynthetic process
- Lipid transport
- Fatty acid metabolic process
- Very long-chain fatty acid biosynthetic process
- Sphingolipid biosynthetic process

**C** Protein level of ribosomal proteins

- Log10 (fold change: MU/WT)

Legend:
- sham
- PH

See legend on next page.
body lengths showed that the WT fish continued to grow from 6 months (average 31.38 mm) to 8 months (average 32.04 mm) and to 12 months (average 33.71 cm) (Fig. 2e). In contrast, the growth of capn3bΔ14 fish was obviously retarded from 6 months (30.58 mm) to 8 months (29.58 mm) and 12 months (29.59 mm) (Fig. 2e). LBR is relatively constant in zebrafish (Kan et al. 2009; Zhu et al. 2014). We checked the LBR of adult WT and capn3bΔ14 fish. LBR in WT did not show significant difference between 6 and 12 months (Fig. 2f). In contrast, the capn3bΔ14 fish exhibited a higher LBR at 12 months than 6 months (Fig. 2f), probably due to shortened body length or failure in maintaining the liver homeostasis. Taken together, these results showed that, as the capn3-KO mice (Kramerova et al. 2004), the growth of capn3bΔ14 fish is sensitive to environmental stresses.

**capn3b** mutant suffers from delayed liver regeneration due to delayed cell proliferation after PH

Adult zebrafish liver comprises two dorsal lobes and one ventral lobe (Kan et al. 2009; Zhu et al. 2014). Previous reports have shown that, in zebrafish, hepatocyte proliferation starts in 2-3 days after resecting almost the entire ventral lobe (~40% of the liver mass) and that the liver weight can be fully restored within 7 days post hepatectomy (dpH) (Kan et al. 2009; Zhu et al. 2014). We conducted PH on the ventral lobe (applied in all PH in this work) on 8-months-old adult WT and capn3bΔ14 fish. The sham group only had their abdominal skin cut through. LBR was used to evaluate liver regeneration. We examined zebrafish at 3dpH, 7dpH and 30dpH and found that LBR was recovered within 7dpH in the WT group (Fig. 3a, left panel). In contrast, the LBR failed to be fully restored at 7dpH but was restored to the normal at 30dpH in capn3bΔ14 fish (Fig. 3a, right panel). We also performed PH in 16-months-old fish and found that the LBR in WT was largely restored at 7dpH and was fully restored to the normal at 30dpH (Fig. 3b, left panel). For the 16-months-old capn3bΔ14 fish, as for the 8-months-old fish, the LBR was not restored at 7dpH but regained their LBR at 30dpH (Fig. 3b, right panel). Taken together, the data demonstrated that capn3bΔ14 fish was delayed in liver regeneration after PH.

We examined cell proliferation in the amputated region by abnormally injecting EdU into the amputated WT and capn3bΔ14 fish at 2dpH and harvested the liver at 3dpH for staining of EdU and Bhmt (marker for hepatocytes). We calculated the EdU-positive cells out of the total DAPI-positive cells. The results showed that, as reported previously (Kan et al. 2009; Zhu et al. 2014), WT hepatocytes were actively undergoing proliferation (14.28% EdU-positive cells) at 3dpH (Fig. 3c, d). In contrast, the capn3bΔ14 liver had very few EdU-positive cells (0.81%) at 3dpH (Fig. 3c, d). We also found that the proliferation of hepatocytes was active in the deeper region (away from the amputated site) in WT but not in capn3bΔ14 PH-liver at 3dpH (Fig. 4a). Therefore, cell cycle reentry in both amputated and deeper region in capn3bΔ14 fish was arrested/delayed at 3dpH.

Next, we injected EdU at 6dpH and harvested the liver at 7dpH. There were almost no proliferating hepatocytes in WT both in the amputated (1.86%) (Fig. 3e, f) and deeper region (Fig. 4b), demonstrating the completion of liver regeneration at 7dpH. On the other hand, in capn3bΔ14 fish, proliferation of hepatocytes seemingly has just begun in the amputated region (15.88% EdU-positive cells) (Fig. 3e, f). Intriguingly, in the deeper region, cell proliferation seemingly has not yet fully activated in capn3bΔ14 fish (Fig. 4b). Whether cell proliferation would be activated later in the deeper region remains unknown. Therefore, cell cycle reentry was not only delayed but also unsynchronized in capn3bΔ14 after PH.

**Proteome analysis reveals the involvement of cell cycle negative regulators in early regeneration of capn3bΔ14 liver after PH**

We previously showed that the Def-Capn3 complex cleaves proteins such as p53 and Mpp10 to regulate cell cycle (Tao et al. 2013; Guan et al. 2016; Zhao et al. 2019). To determine the proteins involved in the delay of capn3bΔ14 liver regeneration, we isolated liver nuclei from WT sham, capn3bΔ14 sham, WT PH, and capn3bΔ14 PH fish (each with three repeats, each repeat containing 8 male fish) at 3dpH for protein extraction. All nuclear protein samples were enriched with nucleolar proteins Def and Mpp10 when compared with the total protein extract and cytoplasmic protein fraction from WT livers (Fig. 5a). Total nuclear proteins were subjected to MS analysis (Table S2 and S3). Heat map clustering analysis of the MS data showed that three WT sham samples were clearly distinctive from three capn3bΔ14 sham samples (Fig. 5b, left panel). The MS data from two WT PH samples (WT-1P and WT-3P)
**A** Venn diagram for upregulated proteins

- Sham: 62
- Share: 19
- PH: 325

**B** Most enriched GO term for biological process

- Proteolysis involved in cellular protein catabolic process
- rRNA processing
- Ubiquitin-dependent protein catabolic process
- mRNA processing
- Maturation of LSU-rRNA
- Proteasome-mediated ubiquitin-dependent protein catabolic process
- Positive regulation of RNA polymerase II transcriptional preinitiation
- Ribosome biogenesis
- Protein catabolic process
- mRNA splicing, via spliceosome

**C** Proteasome subunits

| Log10(fold change: MU/WT) |
|---------------------------|
| 0.1 | 0.2 | 0.3 |
| psmb1 | psmb2 | psmc1b, psmc1a |
| psmc5 | psmc4 | psmc3 |
| psmc6 | psmc5 | psmc4 |
| psmc1b | psmc1a | psmc3 |
| psmc2 | psmc1 | psmc2 |
| psmc3 | psmc4 | psmc3 |
| psmc6 | psmc5 | psmc6 |
| psmc3 | psmc4 | psmc3 |
| psmc6 | psmc5 | psmc6 |
| psmc3 | psmc4 | psmc3 |
| psmc6 | psmc5 | psmc6 |

**D** Protein level of cell cycle repressors

| Protein | Sham | PH |
|---------|------|----|
| ppm1g   |      |    |
| H1.2    |      |    |
| krt18a.2|      |    |
| ddb2    |      |    |
| chd1l   |      |    |
| mre11a  |      |    |
| bcclip  |      |    |
| parp2   |      |    |
| hmgα2   |      |    |
| hmgα1b  |      |    |
| xrc5    |      |    |
| xrc6    |      |    |

**E** DNA Repair Related Proteins

| Protein | Sham | PH |
|---------|------|----|
| nbn     |      |    |
| lig3    |      |    |
| int3    |      |    |
| int5    |      |    |
| ercc3   |      |    |
| ddb2    |      |    |
| cul4a,b; cul4a | |    |
| chd1l   |      |    |
| aptx    |      |    |
| waca; wacb |     | |
| nrb12   |      |    |
| nrb15   |      |    |
| mre11a  |      |    |
| h2afx   |      |    |
| bcclip  |      |    |
| parp1   |      |    |
| parp2   |      |    |
| lig3    |      |    |
| polb    |      |    |
| nth1    |      |    |
| neil1   |      |    |
| hmgα1a  |      |    |
| hmgα2   |      |    |
| hmgα1b  |      |    |
| apex1   |      |    |
| xrc1    |      |    |
| xrc5    |      |    |
| xrc6    |      |    |

Fig. 7 (See legend on next page)
were clearly distinctive from two \textit{capn3b}}^{Δ19Δ14} \textit{PH} samples (3b-1P and 3b-2P) (Fig. 5b, right panel). We chose the best fit clustering groups (for sham samples, WT-1S, WT-3S, 3b-1S and 3b-2S were used, for PH samples, WT-1P, WT-3P, 3b-1P and 3b-2P were used) for further analysis (Fig. 5c).

For the down-regulated proteins (cutoff value: WT verse \textit{capn3b}}^{Δ19Δ14} ≥ 1.45 folds), a total of 92 and 184 down-regulated proteins were identified for the \textit{capn3b}}^{Δ19Δ14} sham (Table S4) and PH samples (Table S5), respectively, while 29 proteins (Table S6) were shared between \textit{capn3b}}^{Δ19Δ14} sham and PH samples (Fig. 6a). Gene ontology (GO) analysis of the 155 proteins uniquely downregulated in \textit{capn3b}}^{Δ19Δ14} PH samples showed that most of the proteins were involved in metabolic and lipid biosynthesis processes in the category of biological process (Fig. 6b). Majority of ribosomal proteins were also down-regulated in \textit{capn3b}}^{Δ19Δ14} PH samples (Fig. 6c). Since active proliferating cells demand higher energy supply and ribosome biogenesis, these results support the observation that cell cycle reentry is not activated in \textit{capn3b}}^{Δ19Δ14} at 3dpH (Fig. 3, Fig. 4).

For the upregulated proteins (cutoff value: \textit{capn3b}}^{Δ19Δ14} verse WT ≥ 1.45 folds), a total of 71 and 344 upregulated proteins were identified in \textit{capn3b}}^{Δ19Δ14} sham (Table S7) and PH samples (Table S8), respectively, while 19 proteins (Table S9) were shared (Fig. 7a), suggesting that different proteomes were mobilized between WT and \textit{capn3b}}^{Δ19Δ14} PH samples. For 325 uniquely upregulated proteins in \textit{capn3b}}^{Δ19Δ14} PH samples, GO analysis showed that many proteolysis-related proteins fell into the biological process category (Fig. 7b). Further analysis of 32 26S proteasome subunits in the MS data showed that there was no significant difference between WT and mutant sham samples, however, 31 subunits were significantly accumulated in \textit{capn3b}}^{Δ19Δ14} PH samples (Fig. 7c). GO analysis also revealed that rRNA processing related proteins were upregulated in \textit{capn3b}}^{Δ19Δ14} PH samples (Fig. 7b), probably due to requirement of extra number of ribosomes for protein biosynthesis to cope with the activation of proteolysis activity in \textit{capn3b}}^{Δ19Δ14} PH fish. Alternatively, depletion of Capn3 might have led to accumulation of its direct or indirect targets that might act as a cue to trigger the activation of the 26S proteasome pathway.

Interestingly, several negative regulators of cell cycle, including PPM1G (2.63 folds), DDB2 (1.87 folds) and H1.2 (linker histone) (2.20 folds), were found to be upregulated in \textit{capn3b}}^{Δ19Δ14} PH samples (Fig. 7d). PPM1G, the phosphatase of p27, functions to stabilize p27 through dephosphorylating p27 at T198 site thus to inhibit cell cycle progression (Sun et al. 2016). DDB2 is able to facilitate IR-induced phosphorylation of Chk1, thus causing cell cycle arrest (Zou et al. 2016). H1.2 augments global association of pRb with chromatin, enhances transcriptional repression by pRb, and facilitates pRb-dependent cell-cycle arrest (Munro et al. 2017). In addition, many DNA-repair related proteins were upregulated in \textit{capn3b}}^{Δ19Δ14} PH samples (Fig. 7e). We failed to identify cyclins and CDKs in our MS data, probably due to relatively low expression of these proteins. Therefore, the delay of cell cycle reentry in \textit{capn3b}}^{Δ19Δ14} PH fish is likely due to up-regulation of proteins involved in repressing cell cycle and down-regulation of proteins involved in metabolic activities.

\textbf{Chek1 and Wee1 are substrates of the Def-Capn3 complex}

Western blot analysis showed that p53, a key negative regulator of cell cycle, was accumulated in the \textit{capn3b}}^{Δ19Δ14} PH samples at 3dpH (Fig. 9a). However, we could not identify p53 protein signatures in our MS data, likely due to its low expression beyond the detection limit of MS. To overcome this limitation, we sought to identify Capn3 substrates based on predicting Capn3 recognition motif \textit{[LIMV]} X (Goessling et al. 2008; Michalopoulos 2007) \textit{[LIMV]} X (Kan et al. 2009) \textit{[LIMV]} \textit{[DE]} among a group 15 of selected cell cycle related proteins (Fig. 8a, left panel; Table S10). We found that Chk1 contained one and Wee1 protein contained two Capn3 recognition motifs which are highly conserved among human, mouse and zebrafish (Fig. 8a, middle and right panels). Chk1 and Wee1 are important G2-M transition check-point proteins which can be activated by phosphorylation upon internal or external stresses (Barnum and O’Connell 2014). Previous studies of the turnover of these two proteins have mostly focused on the proteasome degradation pathway (Zhang et al. 2005; Watanabe et al. 2004b). To determine whether Chk1 and Wee1 are indeed the substrates of the Def-Capn3 complex.
Fig. 8 (See legend on next page.)
complex, we co-injected different combinations of mRNA into WT embryos at the one-cell stage and extracted total proteins at 8hpf. Protein analysis showed that, as expected (Tao et al. 2013; Guan et al. 2016), the p53 mutant protein p53R143H is resistant to Def-mediated degradation. In contrast, co-injection of chek1 or wee1S44A mRNA with def mRNA greatly downregulated the protein levels of both Chk1 and Wee1 (Fig. 8a). We then generated a plasmid expressing a Chk1 mutant protein (Chk1Δ) in which the Capn3 recognition motif was deleted. Co-injection experiment showed that the Chk1Δ mutant protein was resistant to Def-mediated degradation (Fig. 8c).

Next, we examined Chk1 and Wee1 in the defΔ mutant at 5dpf and found that both proteins were up-regulated (Fig. 8d), as did the p53 protein (Tao et al. 2013). Co-immunostaining of Fibrillarin (Fib) (a nucleolar marker) with Chk1 or Wee1 revealed that while both Chk1 and Wee1 were distributed in the nucleoplasm and nucleolus in WT these two proteins were mainly enriched in the nucleoli in the defΔ mutant (Fig. 8e, f), as did the p53 protein (Tao et al. 2013).

**Capn3b depletion enhances Chk1 and Wee1 accumulation after PH**

To determine the status of Chk1 in capn3bΔ19Δ14, we first examined the levels of Chk1 proteins in capn3bΔ19Δ14 embryos at 3.5dpf and 5dpf grown at 30°C and 34.5°C, respectively. Although no obvious difference was observed between WT and capn3bΔ19Δ14 embryos at 3dpf Chk1 protein was greatly accumulated in capn3bΔ19Δ14 embryos at 5dpf at both rearing conditions (Fig. 9b).

Next, we examined Chk1 protein in the liver of WT and capn3bΔ19Δ14 PH fish at 3dpH. The result showed that the expression levels of Chk1 were lower in both WT and capn3bΔ19Δ14 sham groups than in the livers of PH fish (Fig. 9c), suggesting that PH promotes the accumulation of Chk1. However, the liver of capn3bΔ19Δ14 PH fish accumulated much more Chk1 proteins than did the WT PH fish (Fig. 9c). Finally, we performed immunostaining experiment to determine the status of Wee1 protein in regenerating liver at 3dpH. We found that, in WT, Wee1 protein was distributed in the nucleus with certain degree of enrichment in the nucleolus (Fig. 9d, upper panels). In contrast, Wee1 protein was mainly accumulated in the nucleus in the capn3bΔ19Δ14 mutant liver at 3dpH (Fig. 9d, lower panels). These results indicate that both Chk1 and Wee1 might be, at least in part, responsible for the delayed liver regeneration in capn3bΔ19Δ14 PH-fish.

**Discussion**

We have previously proposed that the Def-Capn3 complex might serve as a nucleolar checkpoint for cell proliferation by selective inactivation of cell cycle-related substrates such as p53 during early organogenesis (Guan et al. 2016). In this report, we strengthened the above point by demonstrating that the Def-Capn3b complex is capable to cleave Chk1 and Wee1, two well-defined negative regulators of cell cycle.

In the context of liver regeneration after PH, we found that depletion of Capn3 delayed liver regeneration by disrupting the synchronization of cell cycle reentry, possibly through accumulating Chk1, Wee1, p53, PPM1G and other negative regulators. Notably, PPMG1 is known to stabilize Chk1 protein and Wee1 is a downstream effector of Chk1 (Sun et al. 2016), therefore, it is reasonable to propose that Chk1 and Wee1 likely play a key role in delaying cell proliferation during the regeneration of the capn3bΔ19Δ14 PH-liver. The MS data showed that the levels of 31 subunits of the 26S proteasome complex were elevated in the capn3bΔ19Δ14 PH-liver. It is imaginable that the activated proteasome pathway could gradually degrade the accumulated negative regulators of cell cycle in the capn3bΔ19Δ14 PH-liver although the synchronization of cell proliferation was probably compromised. This might explain why the capn3bΔ19Δ14 PH-liver finally regained its mass at 30dpH.

Based on our data, we propose that PH will activate the cell cycle reentry program in hepatocytes but meanwhile will impose a restriction on cell cycle progression by triggering the expression of negative regulators of cell cycle such as p53, Chk1 and Wee1. Although these proteins are subjected to proteasome pathway mediated degradation the synchronization of cell cycle reentry is achieved by the Def-Capn3b pathway which functions to...
Fig. 9 Hepatic accumulation of Chk1 and Wee1 in capn3bΔ19Δ14 at 3dpH. a Elevation of p53 protein level in capn3bΔ19Δ14 after PH. Western blot analysis of p53 at different time point after PH as indicated. Total proteins were extracted from the liver tissues of sham, 3dpH, 5dpH, 7dpH, 10dpH and 14dpH, respectively. GAPDH, loading control. b Western blot of Capn3b and Chk1 in WT and capn3bΔ19Δ14 mutant embryos at 3.5dpf and 5dpf grown at 30 °C and 34.5 °C, respectively. The embryos were shifted to 34.5 °C at 12hpf till the time of harvesting. β-Actin: loading control. c Western blot of Chk1 in WT and capn3bΔ19Δ14 sham and PH groups at 3dpH group. Fibrillarin: loading control. d Co-immunostaining of Wee1 and Fibrillarin in the liver of WT and capn3bΔ19Δ14 mutant fish at 3dpH. DAPI: staining nuclei.
inactivate these proteins more promptly. Since the Def-Capn3b pathway mediated substrate cleavage is independent of the ubiquitination pathway we believe the Def-Capn3b pathway plays a unique function in the regulation of cell cycle progression during liver regeneration after PH. Based on the fact that the Def-Capn3 pathway also operates in human cells (Tao et al. 2013; Guan et al. 2016), it is envisaged that the Def-Capn3 pathway likely plays a role in regulating liver regeneration in mammals.

Zebrafish genome contains 19 calpain homologous genes (Tao et al. 2013; Ma et al. 2019). These calpain homologs might compensate for the loss-of-function of Capn3b in capn3bΔ19Δ14, which nicely explains why the capn3bΔ19Δ14 fish is viable and fertile (Ma et al. 2019). However, the growth of capn3bΔ19Δ14 fish is retarded, including a small liver at the fry stage and short body length at the adult stage. The growth retardation is exaggerated in response to environmental stresses such as high rearing intensity or high temperature. These observations suggest that gene compensation in the capn3bΔ19Δ14 fish is conditional. This is consistent with the observation that Def mainly recruits Capn3b but not its closer homolog Capn3a to the nucleolus (Tao et al. 2013). It would be interesting to explore the extent of gene compensation for different genes under different conditions in the future. Interestingly, the capn3bΔ19Δ14 fish displayed an LBR largely similar to the WT fish, suggesting that the delay of liver regeneration in capn3bΔ19Δ14 fish after PH is genuinely attributed to the depletion of Capn3b. Future work is required to generate capn3 Δ KO mouse for determining whether this is also the case in mammals after PH.

Supplementary information

Supplementary information accompanies this paper at https://doi.org/10.1186/s13619-020-00049-1.

Additional file 1: Table S1. List of primers for plasmid construction.

Additional file 2: Table S2. Full MS data for six sham protein samples.

Additional file 3: Table S3. Full MS data for six PH protein samples.

Additional file 4: Table S4. 92 proteins downregulated in capn3ΔΔ1419 sham samples.

Additional file 5: Table S5. 184 downregulated in capn3ΔΔ1419 PH samples.

Additional file 6: Table S6. 29 downregulated proteins shared in capn3ΔΔ1419 sham and PH samples.

Additional file 7: Table S7. 71 upregulated proteins in capn3ΔΔ1419 sham samples.

Additional file 8: Table S8. 344 upregulated proteins in capn3ΔΔ1419 PH samples.

Additional file 9: Table S9. 19 upregulated proteins shared in capn3ΔΔ1419 sham and PH samples.

Additional file 10: Table S10. Predicting Capn3 recognition sites in cell-cycle related proteins.

Abbreviations

Capn3: Calpain3; Chk1: Check-point kinase 1; DBD2: DNA damage binding protein 2; Def: Digestive-organ expansion factor; ddp: days-post-fertilization; dph: days-post-hepatectomy; GO: Gene ontology; fabp10a: liver fatty acid binding protein; MS: Mass spectrometry; PH: Partial hepatectomy; PPM1G: Protein phosphatase 1G; TALEN: Transcription activator-like effector nucleases; WT: Wild-type

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Authors’ contributions

FC, YCW and JRP designed the experiments. FC, DLH, HS and CG performed the zebrafish experiments. YCW performed the MS experiment and FC and JRP analyzed the data and wrote the manuscript. All read and approved the final manuscript.

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Availability of data and materials

Dataset described in this work can be downloaded from the supplementary Tables S1, S2, S3, S4, S5, S6, S7, S8, S9 and S10 from the journal website. Materials request should be addressed to the corresponding author: pengj@zju.edu.cn.

Ethics approval and consent to participate

All animal experiments were conducted according to the guidelines of “Regulation for the Use of Experimental Animals in Zhejiang Province” with a permit issued by the Animal Ethics Committee in Zhejiang University (ETHICS CODE Permit NO. ZJU2011-1-11-009).

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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