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Retinoic Acid Regulates Cell Cycle Genes and Accelerates Normal Mouse Liver Regeneration

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Abstract

All-trans retinoic acid (RA) is a potent inducer of regeneration. Because the liver is the principal site for storage and bioactivation of vitamin A, the current study examines the effect of RA in mouse hepatocyte proliferation and liver regeneration. Mice that received a single dose of RA (25 µg/g) by oral gavage developed hepatomegaly with increased number of Ki67-positive cells and induced expression of cell cycle genes in the liver. DNA binding data revealed that RA receptors retinoic acid receptor β (RAR β) and retinoid x receptor α (RXR α) bound to cell cycle genes Cdk1, Cdk2, Cyclin B, Cyclin E, and Cdc25a in mice with and without RA treatment. In addition, RA treatment induced novel binding of RAR β /RXR α to Cdk1, Cdk2, Cyclin D, and Cdk6 genes. All RAR β /RXR α binding sites contained AGGTCA-like motifs. RA treatment also promoted liver regeneration after partial hepatectomy (PH). RA signaling was implicated in normal liver regeneration as the mRNA levels of RAR β , Aldh1a2, Crabp1, and Crbp1 were all induced 1.5 days after PH during the active phase of hepatocyte proliferation. RA treatment prior to PH resulted in early up-regulation of RAR β , Aldh1a2, Crabp1, and Crbp1, which was accompanied by an early induction of cell cycle genes. Western blotting for RAR β , c-myc, Cyclin D, E, and A further supported the early induction of retinoid signal and cell proliferation by RA treatment. Taken together, our data suggest that RA may regulate cell cycle progression and accelerates liver regeneration. Such effect is associated with an early induction of RA signaling, which includes increased expression of the receptor, binding proteins, and processing enzyme for retinoids.

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Authors' Contributions

Hui-Xin Liu: Performed experiments, analyzed data, generated figures and prepared manuscript.

Irene Ly: Assisted in experiments and prepared manuscript.

Ying Hu: Assisted in experiments and analyzed data.

Yu-Jui Yvonne Wan: Generated idea and supervised overall performance of the project.

Disclosure of potential Conflicts of Interest

The authors declare that there are no known conflicts of interest associated with this publication and there has been no significant financial support for this work that could have influenced its outcome.

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Keywords

retinoic acid receptor; nuclear receptor; hepatocyte; partial hepatectomy; proliferation

1. Introduction

All-trans retinoic acid (RA) has long been recognized as a regeneration-inducing derivative of vitamin A. In amphibians, RA causes “super-regeneration” of naturally regenerative organs, while in mammals, RA induces regeneration of organs that do not normally regenerate, such as the adult mammalian lung (1–3). Interestingly, the liver is the only mammalian organ with highly regenerative properties, making it a prime target for accelerated regrowth through RA. As the principal site for storage and bioactivation of vitamin A in the body, the liver experiences constant RA exposure. Lecithin:retinol acyltransferase-deficient mice, completely lacking in hepatic retinoid stores, have reduced RA levels and impaired liver regeneration in response to partial hepatectomy (PH). Thus, hepatic retinoid storage is required for normal liver regeneration (4).

Despite the essential role of endogenous RA on liver regeneration, there lacks a consensus on the effects of exogenous RA on hepatocyte proliferation *in vivo*, since previous studies have yielded conflicting results (5–8). Early *in vitro* studies reported that RA is a potent inhibitor of DNA synthesis in primary rat hepatocytes (9). Similarly, when administered after PH, RA disrupted rat hepatocyte proliferation *in vivo* by repressing early response genes and increasing the activity of transglutaminase and ornithinedecarboxylase (10–14). In contrast to these findings, other studies have shown that RA enhances hepatocyte proliferation and survival. RA augmented TNF α -stimulated mouse hepatocyte DNA synthesis *in vitro* (15). The direct mitogenic effect of RA has also been demonstrated in normal hepatocyte proliferation of both mice and rats (16, 17). Post-operative administration of RA or synthetic retinoid NIK33 enhanced hepatocyte proliferation in the regenerating rat liver (18).

It is problematic that the existing literature has such contradictory findings since RA is considered a chemotherapeutic agent against certain types of cancer. Despite its anti-proliferative and differentiation effects against malignant cells, it is necessary to acknowledge the role of RA in proliferation and regeneration as well as the clinical implications. Although RA is often used to treat patients with acute promyelocytic leukemia (APL), major complications can manifest in the form of hepatomegaly and hepatotoxicity (19). The conflicting results from previous studies may be attributable to differences in dosage, route of administration, and animal species. Another important factor to consider is the timing of exogenous RA administration. In the previous studies, RA was administered after surgical resection of the liver (10–14, 18). The hepatoprotective role of RA preconditioning has recently been proposed in an ischemic/reperfusion injury model (20), but there is no evidence in the literature that clearly documents the effects of RA treatment prior to PH-induced liver regeneration. Therefore, this study aims to clarify the role of RA in the regenerating liver by demonstrating, for the first time, the effects of RA administration prior to PH.

In this study, RA pretreatment is shown to accelerate PH-induced liver regeneration in mice by increasing the expression of genes encoding Cyclin-Cdk complexes. Moreover, the data indicate that RA pretreatment promoted hepatocyte proliferation in the regenerating mouse liver by modulating cell cycle progression through direct binding by retinoic acid receptor β (RAR β) and retinoid x receptor α (RXR α). Together, these findings emphasize the potential utility of employing retinoids to facilitate liver regrowth following injury.

2. Materials and Methods

2.1 Mice, partial hepatectomy, and sample preparation

Wild type (WT) male mice (3–5 months old) were housed in steel micro-isolator cages (4 mice per cage) at 22°C with a 12-hr/12-hr light/dark cycle. Food and water were provided *ad libitum* throughout the entire study. RA (25 μ g/g) (Sigma-Aldrich Corp., St. Louis, MO) or vehicle control (carboxymethyl cellulose) (Sigma-Aldrich Corp., St. Louis, MO) was administered by oral gavage 48 hours prior to surgery. Standard two-thirds liver resection was performed using the procedure previously described (21–23). Sham-operated mice were included as controls. Surgeries were performed between 9:00 to 11:00 AM. Mice were killed at indicated time points and their liver and body weight recorded at the time of death were used to calculate liver-to-body weight ratio. The data presented were calculated from the mean of three to five mice per time point. Liver tissues were collected and snap frozen in liquid nitrogen and stored at -80°C . A section of each liver was fixed in 10% formalin, embedded in paraffin, and stained with hematoxylin and eosin for histological analysis. All animal experiments were conducted in accordance with the National Institutes of Health (NIH) Guide for the Care and Use of Laboratory Animals and the Institutional Animal Care and Use Committee (IACUC) at the University of California, Davis.

2.2 Ki67 immunostaining

To assess hepatocyte proliferation, immunostaining was performed with anti-Ki67 antibody (NeoMarkers, Fremont, CA). The number of Ki67-labeled nuclei was counted in at least 15 microscopic fields (20X) for each liver section.

2.3 Chromatin immunoprecipitation followed by sequencing (ChIP-Seq)

ChIP analyses were performed as described previously (5). Briefly, chromatin lysate was precleared before incubation with a ChIP-quality anti-RAR β antibody (Santa Cruz, CA). Anti-IgG (Santa Cruz, CA) and anti-RNA Polymerase II antibodies (Millipore, MA) were used as negative and positive controls, respectively. Target sequences were aligned to the mouse genome based on USC genome browser with Bowtie 0.12.7 (24) followed by peak-calling using Model-based Analysis of ChIP-Seq (MACS) 1.4.1 (25). The called peaks were annotated against the database (GRCm38/mm10) by Peak Analyzer (26). Motif Analysis of Large DNA Datasets (MEME-ChIP) was used to identify potential motifs for target binding.

2.4 Quantification of hepatic mRNA

Hepatic RNA isolated by TRIzol (Invitrogen, Carlsbad, CA) was reverse transcribed to generate cDNA followed by amplification using the ABI Prism 7900HT sequence detection system (Applied Biosystems, Foster City, CA). Hepatic mRNA levels were normalized to

the mRNA expression of glyceraldehyde-3-phosphate dehydrogenase (GAPDH). Primers were designed using Primer3 Input Software (v0.4.0) and sequences are available upon request.

2.5 Western blot

Liver protein (40 μ g) was electrophoresed on SDS-polyacrylamide gels under reducing conditions. Proteins from the gels were transferred to the polyvinylidene fluoride membranes. Anti-Cyclin A, Cyclin D, Cyclin E (Cell Signaling Technology, MA), c-myc, RAR β , and β -Actin (Santa Cruz, CA) antibodies were used for detection of proteins. Protein expression levels were quantified using ImageJ.

2.6 Statistical Analysis

Data are given as mean \pm SD. Statistical analysis was performed using Student's *t* test or one-way analysis of variance. Statistical significance was defined by $p < 0.05$.

3. Result

3.1 RA induces hepatocellular proliferation in mice

WT mice received oral gavage with RA or vehicle 48 hours prior to a sham operation. Liver-to-body weight ratio was significantly higher in RA-treated versus control mice 1 to 7 days after sham operation (Fig. 1A). The RA-induced increase in liver mass correlated with greater hepatocellular proliferation, as shown by increased numbers of Ki67-positive hepatocytes at studied time points (Fig. 1B, C). Together, results from the sham-operated RA treatment group indicate that the administration of RA promotes hepatomegaly and cell proliferation in the normal mouse liver.

3.2 RA enhances RAR β /RXR α binding to cell cycle genes

RAR β is a RA-inducible receptor that has been shown to be up-regulated during the early phase of liver regeneration (27). RAR β forms a heterodimer with RXR α to bind target genes and regulate their transcription. RA-induced hepatomegaly and hepatocyte proliferation were studied on a genomic level by analyzing RAR β and RXR α common DNA binding sites in livers derived from WT mice with or without RA treatment (5, 28). The results suggest that RAR β /RXR α binding mediates the effect of RA on cell cycle regulation. In the vehicle-treated group, 34 out of 129 cell cycle genes obtained from the Kyoto Encyclopedia of Genes and Genomes (KEGG) cell cycle pathway database were bound by RAR β /RXR α heterodimer. In the RA-treated group, there was a significant increase in both the number of cell cycle genes (50 vs. 34) that were bound by RAR β /RXR α as well as the number of peaks (66 vs. 41). These findings are consistent with the observation that RA treatment enhanced cell proliferation. Cell cycle regulatory genes were modulated by RA treatment in a manner supportive of accelerated cell cycle progression. From the KEGG pathway database, the expression of 18 cell cycle genes was studied and 11 of them demonstrated significantly higher mRNA levels following RA treatment (Fig. 2A). Of these 11 genes, five (Cyclin-dependent kinase 1 (Cdk1), Cdk2, Cyclin B, Cyclin E, Cell division cycle 25a (Cdc25a)) shared common RAR β /RXR α binding in both control and RA-treated groups. Additional RAR β /RXR α binding was noted in the Cdk1 and Cdk2 genes after RA treatment. RAR β /

RXR α binding was absent in the Cyclin D and Cdk6 genes in the control group, but was induced in the RA treatment group (Fig. 2B). All aforementioned RAR β /RXR α binding regions contained AGGTCA-like motifs as hormone response elements. However, RAR β /RXR α binding was not found in forkhead box M1 (Foxm1), Cyclin A, Cdc25b, and Cdc25c genes even though RA treatment induced their mRNA levels (Fig. 2A).

3.3 RA treatment accelerated liver regeneration in response to PH

Based on the RA treatment experiments in sham-operated mice, the proliferative effect exerted by exogenous RA in the regenerating liver was studied. WT mice were treated with RA or vehicle control 48 hours prior to the surgery. In agreement with findings from sham-operated groups, RA administration increased the liver-to-body weight ratio in regenerating livers 1 to 5 days after PH (Fig. 3A). In addition, the increase in liver-to-body weight ratio was accompanied by greater hepatocyte proliferation, as demonstrated by the higher number of Ki67-positive cells in RA-treated mouse livers (Fig. 3B, C). Thus, these results indicate that RA administration prior to hepatic resection accelerated regeneration in the mouse liver.

3.4 Regenerating livers in RA-treated mice displayed early induction of RA signaling

Because RA signaling has been implicated in liver regeneration, the expression of genes involved in the RA signaling pathway including RAR β , Aldehyde dehydrogenase family 1 member A2 (Aldh1a2), Cellular retinoic acid binding protein 1 (Crabp1), and Cellular retinol binding protein 1 (Crbp1) were studied in regenerating livers from control and RA-treated mice (29). RAR β is a RA receptor whose expression is inducible by RA (7) while Aldh1a2 is responsible for producing RA (29). Crbp1 and Crabp1 are binding proteins for retinoids with Crabp1 also being a RA target gene (30). The expression of all four genes was transiently induced 1.5 days after PH in control mice. The coordinated induction of all four genes at the same time point suggests the importance of RA signaling in hepatocyte proliferation, which peaked 1.5–2 days after PH (Fig. 4). RA treatment prior to the surgery resulted in early induction of RAR β , Aldh1a2, Crbp1, and Crabp1 1 day after PH. Western blotting showed elevated RAR β protein levels resulting from RA treatment 1 day after PH (Fig. 4E). Moreover, Crbp1 induction was sustained for up to 5 days following surgery. The accelerated liver regeneration in RA-treated mice was accompanied by early activation of RA signaling via increased expression of the receptor, binding proteins, and oxidation enzyme for retinoids.

3.5 RA modulates cell cycle gene expression during liver regeneration

To gain further insights into the effect of RA on cell cycle progression, the expression of cell cycle genes in regenerating mouse livers was studied. Foxm1 and its target genes Cdc25a, b and c as well as the E2f family have been implicated in regulating Cyclin/cdk activity at key cell cycle checkpoints (31). RA treatment facilitated the induction of Foxm1, Cdc25a, Cdc25b, Cdc25c, and E2f transcription factor 1 (E2f1), E2f2, E2f7, E2f8 mRNA levels as early as 1 day after PH, peaking 1.5 days after PH when the hepatocytes are actively proliferating (Fig. 5A–H). RA treatment did not alter the expression of E2f3, 4, 5, and 6 (Fig. 5I–L). In accordance with Ki67 staining data, Proliferating Cell Nuclear Antigen (Pcna) mRNA levels were markedly higher in RA-treated than the control group 1 to 5 days after PH (Fig. 6A). Quantification of cyclins and cdk expression revealed that in the vehicle

control mice, hepatic induction of these key cell cycle genes is observed 1.5 days after PH, a well-established indicator of peak hepatocyte proliferation in regenerating mouse liver (32). In RA-treated mice, hepatic expression of cyclins and cdks was significantly elevated 1 day after PH compared to control (Fig. 6B–H). Consistently, western blotting showed higher protein levels of c-myc, Cyclin D, Cyclin E, and Cyclin A 1 day after PH in RA-treated group relative to the control group (Fig. 7). Taken together, RA treatment accelerated the expression of a panel of cell cycle genes in the regenerating mouse liver.

4. Discussion

RA is well known for its roles in differentiation, anti-proliferation, and regeneration. The complex and pleiotropic effects of RA have yielded conflicting findings on its many functions. Our data showed that oral gavage of RA increased liver size and hepatocyte proliferation. This mitogenic effect was also observed in the regenerating liver when mice were treated with RA prior to PH. Furthermore, new insights into the role of RA and its receptor RAR β *in vivo* were obtained. Although RAR β is generally considered as a tumor suppressor gene, this study demonstrated that hepatic RAR β binding is also implicated in cell cycle signaling. RA treatment accelerates liver cell proliferation and these effects are facilitated by early induction of cell cycle genes, which were accompanied by early activation of RA-mediated signaling.

To determine the molecular basis by which RA promotes hepatocellular proliferation, RAR β /RXR α binding to cell cycle genes was analyzed before and after RA treatment. Within the identified RAR β /RXR α binding sites that contain an AGGTCA-like motif, additional transcription factor binding motifs for SP1, GABPA, and FOXA2 were detected (data not shown). Of particular interest is SP1, which has previously been shown to interact with RAR β and whose binding sites have been identified in genes associated with growth and cell cycle regulation as well as RA signaling. For instance, RA-induced expression of the retinol binding protein gene is regulated by a SP1-RAR β /RXR α complex (33). Moreover, RA can regulate SP1 phosphorylation (34), and SP1 down-regulation leads cell cycle arrest in human cancer cells (35). Specifically, SP1 modulates the activity of checkpoint proteins p21 and p27 in human hepatoma cells (36). Therefore, it appears that the action of RA-RAR β in regulating the cell cycle could be exerted through SP1 or other transcriptional factors. Further study will be necessary to determine whether these additional transcription factors indeed contribute to enhanced hepatocyte proliferation by interacting with RAR β and transcriptionally activating cell cycle genes.

Liver regeneration is a complex process regulated by many signals from the hepatic environment. Different signaling pathways will lead to the activation of transcription factors that either stimulate hepatocyte proliferation or promote cell survival to facilitate liver regrowth (21, 23, 37, 38). The presented data suggest that one such pathway is the RA signaling pathway itself. Our findings indicate that RA signaling is highly orchestrated in the regenerating liver. This coordinated up-regulation of RAR β , ALDH1A2, CRBP1, and CRABP1 peaked earlier in the RA-treated group than the control group, suggesting the important role of RA signaling in promoting and even accelerating hepatocyte proliferation. Since vitamin A is stored in the liver, it is likely that endogenous RA is mobilized in

regenerating liver to up-regulate its direct target genes including CRABP1 and RAR β and their proliferative downstream pathways.

Accelerated regeneration was also associated with elevated expression of cell cycle regulators and increased hepatocyte DNA replication and mitosis. RA-mediated activation of Foxm1 facilitates a coordinated induction of several common target genes through RAR β binding. Foxm1 also regulates Cdc25, Cyclin E/Cdk2, and Cyclin B/Cdk1, making it a critical gatekeeper of G1/S and G2/M cell cycle transitions, as well as mitotic spindle assembly (39). The E2f transcription factor family plays an important role in regulating cellular proliferation by activating a panel of genes involved in progression through the G1 phase as well as DNA replication (23). The E2f family includes transcription activators and repressors. Activators such as E2f1-3 promote cell cycle progression, while repressors (E2f4-8) inhibit cell cycle. E2f7-8 can inhibit the action of E2f1 via a negative feedback loop (40). Cyclins and Cdks are E2f transcriptional targets (41). RA treatment increased the induction of Foxm1, E2f1, and E2f2 expression. Furthermore, RA treatment elevated the expression of Cyclin D-Cdk 4/6, Cyclin E-Cdk2, and Cyclin A-Cdk2 kinase complexes suggesting increased hepatocellular G1/S phase entry activities, all of which are associated with enhanced hepatocyte DNA replication. Finally, RA treatment induced up-regulation in Cyclin B-Cdk1 complex and Cdc25s mRNA levels in regenerating livers, which are indicative of mitotic entry.

Our findings run contrary to the hallmark anti-proliferative characteristics of RA, which has been used as a chemotherapeutic treatment against certain forms of cancer (42–44). Among the RAR isoforms (α , β , γ), RAR β is the predominant receptor mediating the inhibitory effects of RA on cancer cell proliferation and is considered as a tumor suppressor gene (45, 46). However, the application of RA and other retinoids does not always demonstrate efficacy in preventing or treating cancer (47–54). On the contrary, RA and other retinoids can promote rather than inhibit cancer cell survival and growth as well as increase the incidence of certain cancers in mouse models and human subjects (46, 47, 49, 53). Among the most notable examples are large scale trials on the incidence of lung cancer that found no benefit and even potentially adverse effects of RA or retinoid supplementation (55, 56). Similarly, retinoid chemoprevention did not have a significant effect on lung cancer patients (47). Furthermore, the presence of RAR β actually promotes the growth of mammary carcinoma in the stromal cell compartment (57). The conflicting data on the proliferative effect of RA underscores the need to exercise caution when evaluating RA as a chemotherapeutic agent. Nevertheless, this does not discount the potentially beneficial applications of RA, particular in the context of liver regeneration. The accelerated liver regeneration previously demonstrated sheds additional light on a promising approach to protect the liver during transplants or when challenged with acute or chronic injury. Because efficient liver regeneration is vital for improving surgical outcomes, the pivotal role of RA in enhancing hepatocyte proliferation makes it a strong candidate for reducing post-operative hepatic failure.

In summary, RAR β is implicated as a key component in RA-induced stimulation of cell cycle signaling pathways that promote hepatocyte proliferation. We anticipate that future studies will employ a RAR β liver-specific knockout mouse model as well as use receptor

specific ligands to further characterize the role of hepatic RAR β in cell cycle progression. A better mechanistic understanding of the proliferative effects exerted by RA may lead to the development of therapeutics to promote liver regeneration in the clinical setting.

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Abbreviations

Aldh1a2	aldehyde dehydrogenase family 1 member A2
Cdk1	Cyclin-dependent kinase 1
Cdc25	cell division cycle 25
ChIP-seq	chromatin immunoprecipitation followed by next generation sequencing
Crabp1	cellular retinoic acid binding protein 1
Crbp1	cellular retinol binding protein 1
E2f	E2f transcription factor
Foxm1	forkhead box M1
KEGG	Kyoto encyclopedia of Genes
Pcna	proliferating cell nuclear antigen
PH	partial hepatectomy
qRT-PCR	real-time quantitative reverse transcription PCR
RA	all-trans retinoic acid
RAR	retinoic acid receptor
RXR	retinoid X receptor
WT	wild type

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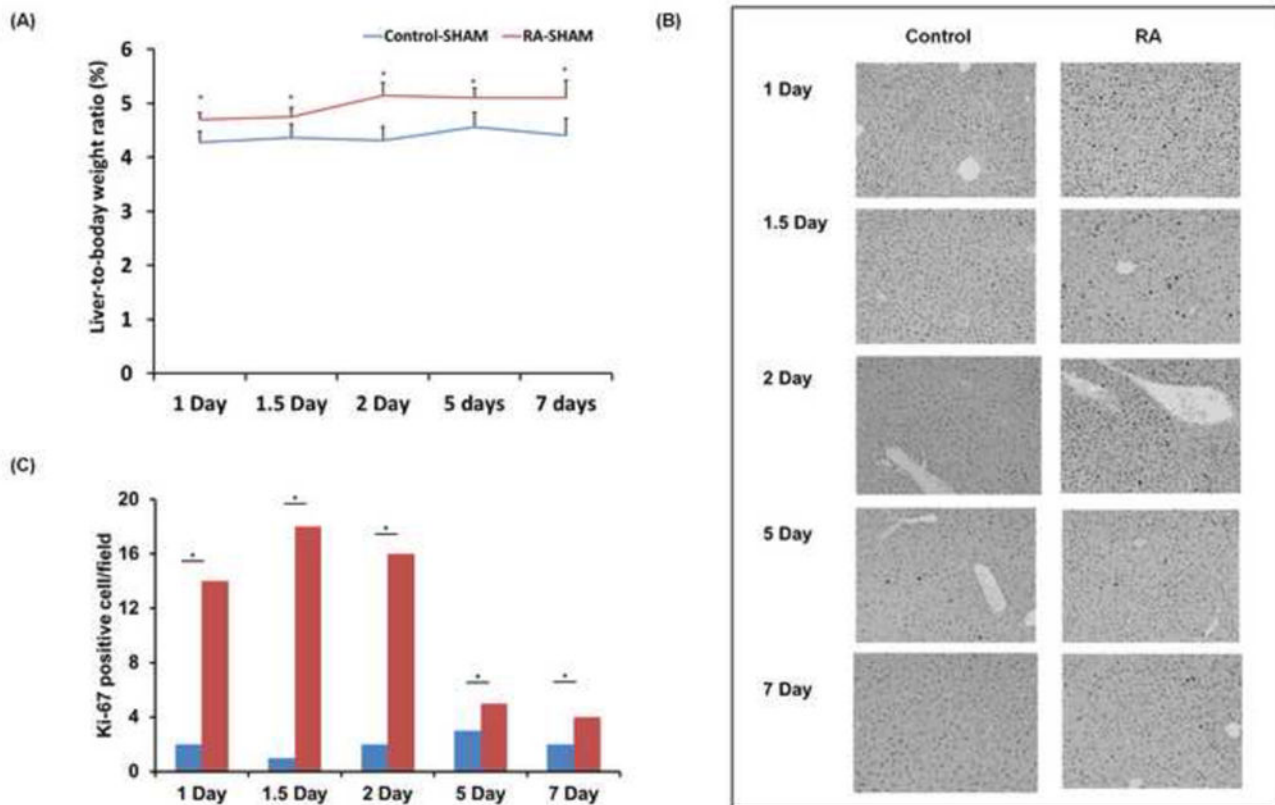


Figure 1. Increased hepatocyte proliferation in RA-treated mouse liver

WT mice were treated with RA or vehicle by oral gavage 48 hours prior to a sham operation. (A) Mice were sacrificed and weighed 1, 1.5, 2, 5 and 7 days after surgery to calculate differences in liver to body weight ratio between the groups. (B) Representative photomicrographs of Ki67 immunohistochemical staining of liver sections from WT mice with and without RA treatment at indicated time points (n = 4). (C) Ki67-positive cells in the livers of WT mice with and without RA pre-treatment after sham operation. The number of proliferating hepatocytes was determined by counting Ki67-positive hepatocytes in at least 15 microscopic fields (20X) per liver sample. Liver sections from all mice were used for analyses. Means \pm SD are graphed with * indicating $p < 0.05$.

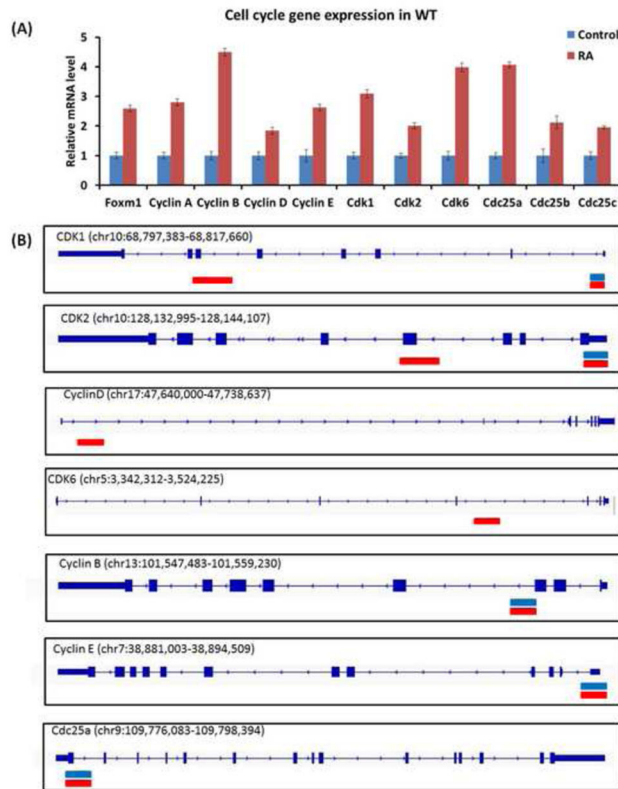


Figure 2. Differential RAR β /RXR α binding in cell cycle genes after RA treatment

(A) Quantitative RT-PCR analysis of a panel of cell cycle genes derived from the KEGG pathway database. From a selection of 18 cell cycle genes, 11 showed significantly increased expression following RA treatment. (B) The effects of RA were studied on a genomic level by analyzing RAR β and RXR α common DNA binding sites with AGGTCA-like motifs from mouse livers with or without RA treatment. Of the 11 cell cycle genes that were up-regulated after RA treatment, five genes (Cdk1, Cdk2, Cyclin B, Cyclin E, Cdc25a) had common RAR β /RXR α binding in both the control and RA treatment groups, as demonstrated by the concurrent blue (control) and red (RA treatment) bars. Novel RAR β /RXR α binding was observed on Cdk1 and Cdk2 genes after RA treatment. Unique RAR β /RXR α binding was found on Cyclin D and Cdk6 genes in the RA-treated group.

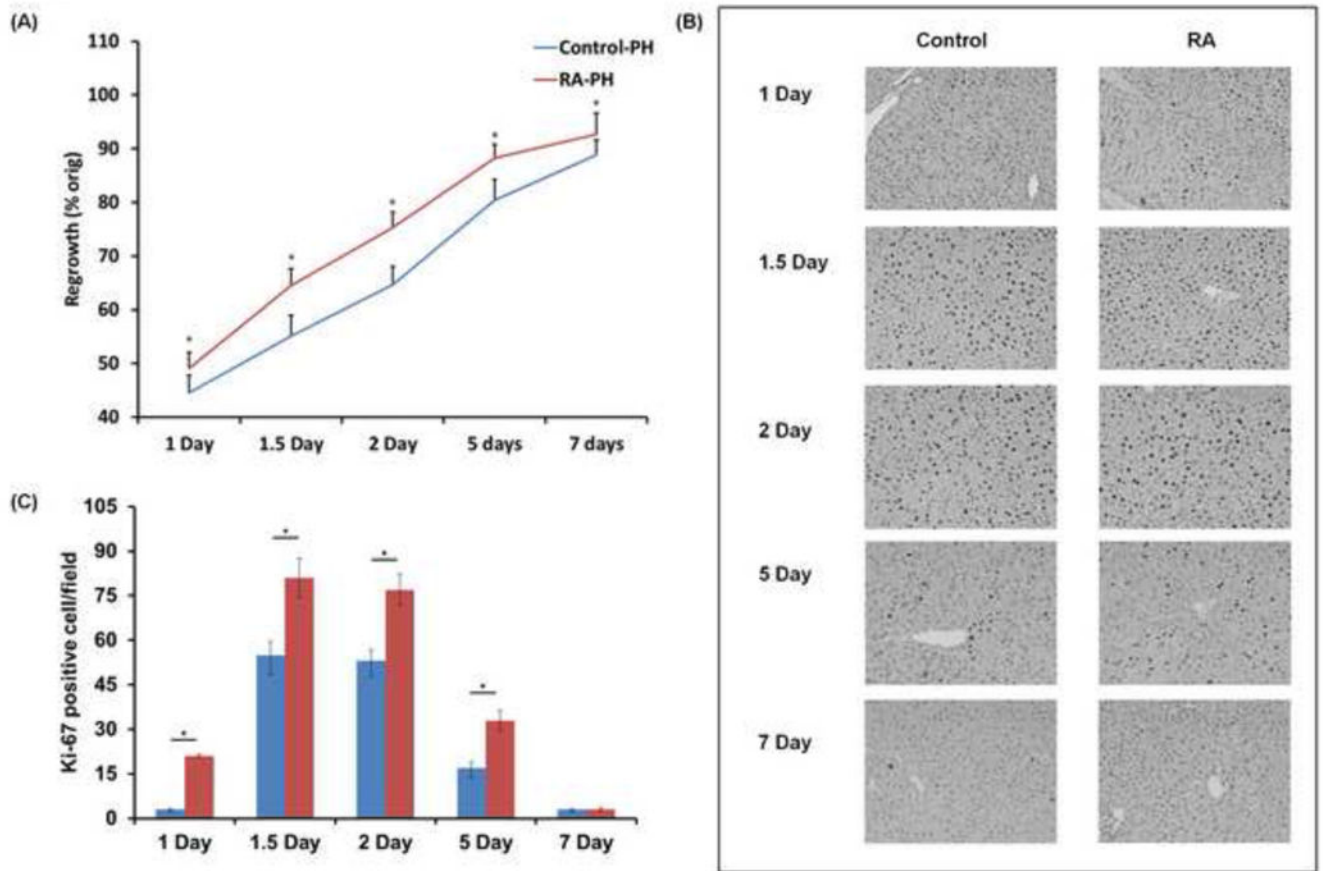


Figure 3. RA treatment accelerated hepatocyte proliferation in regenerating mouse liver

WT mice were treated with RA or vehicle by oral gavage 48 hours prior to partial hepatectomy. (A) Mice were sacrificed and weighed at 1, 1.5, 2, 5 and 7 days after surgery to calculate differences in liver to body weight ratio between treatment groups. (B) Representative photomicrographs of Ki67 immunohistochemical staining of liver sections from wild type mice with and without RA treatment at indicated time points (n = 4). (C) Ki67-positive cells in the livers of wild type mice with and without RA pre-treatment after PH. The number of proliferating hepatocytes was determined by counting Ki67-positive hepatocytes in at least 15 microscope fields (20X) per liver sample. Liver sections from all mice were used for analyses. Means \pm SD are graphed with * indicating $p < 0.05$.

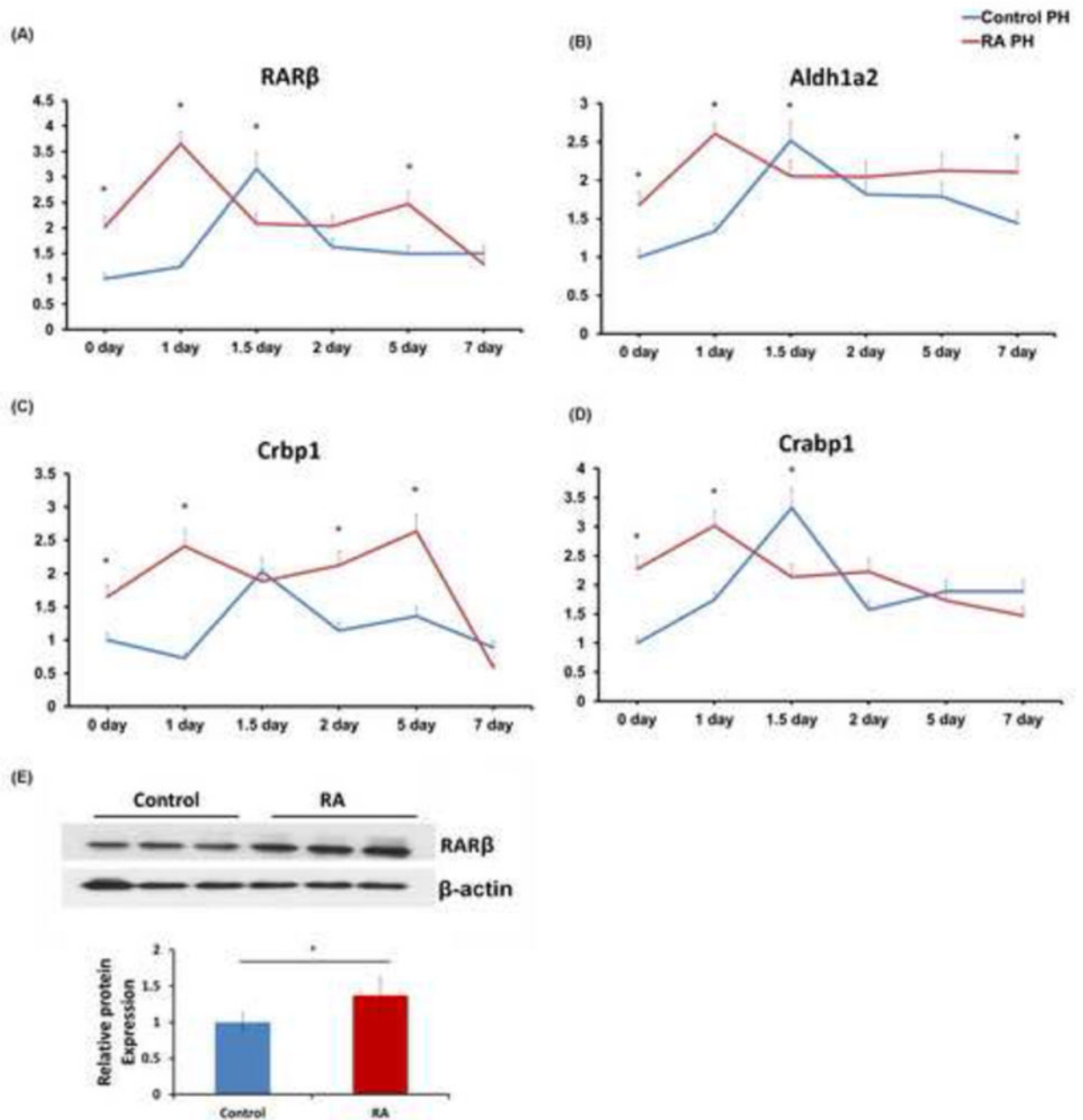


Figure 4. Early induction of RA signaling in RA-treated regenerating mouse livers
 (A–D) RAR β , Aldh1a2, Crbp1, and Crabp1 are genes encoding the RA receptor, retinoid oxidation enzyme, and binding proteins, respectively. mRNA levels of these genes were quantified at indicated time points after PH in control and RA treatment groups. The expression of all four genes was transiently induced 1.5 days after PH in the control treatment group. The introduction of RA treatment 48 hours prior to surgery led to an earlier induction of these RA signaling genes. (E) Western blotting analysis of RAR β protein levels in regenerating liver 1 day after PH from control and RA treatment groups. Means \pm SD are graphed with * indicating $p < 0.05$.

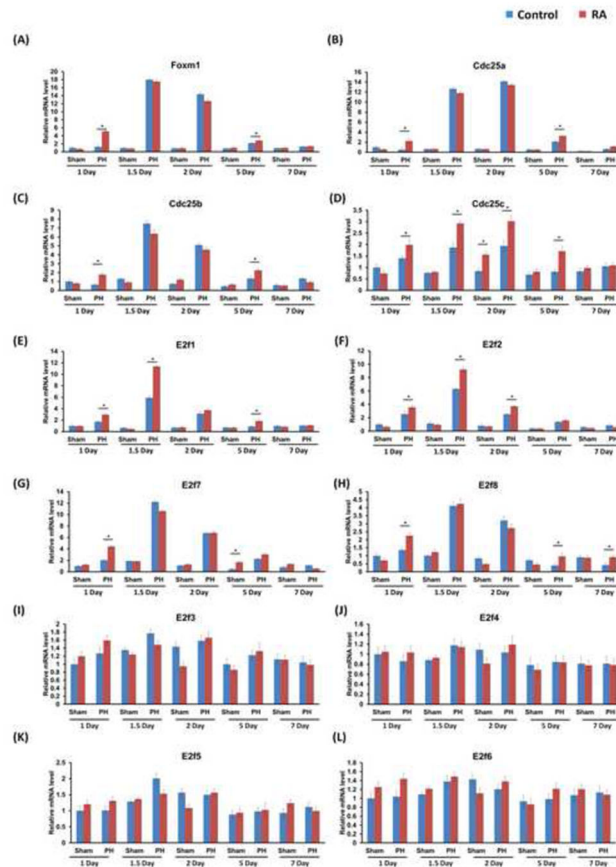


Figure 5. RA modulated the expression of cell cycle regulators during liver regeneration WT mice with or without RA treatment received PH and were killed at indicated time points. The expression of cell cycle regulating genes was studied. These genes include transcriptional regulators (A) Foxm1, (B–D) Cdc25a, b, c, and (E–H) the E2f family of cell cycle regulators. The peaks in gene expression between 1.5 and 2 days after PH in control mice correspond with the peak in hepatocyte proliferation in regenerating livers. RA treatment facilitated early induction of these cell cycle genes 1 day after PH compared with the control group. (I–L) E2f3–6 did not demonstrate a significant change in gene expression levels with RA treatment. Means \pm SD are graphed with * indicating $p < 0.05$.

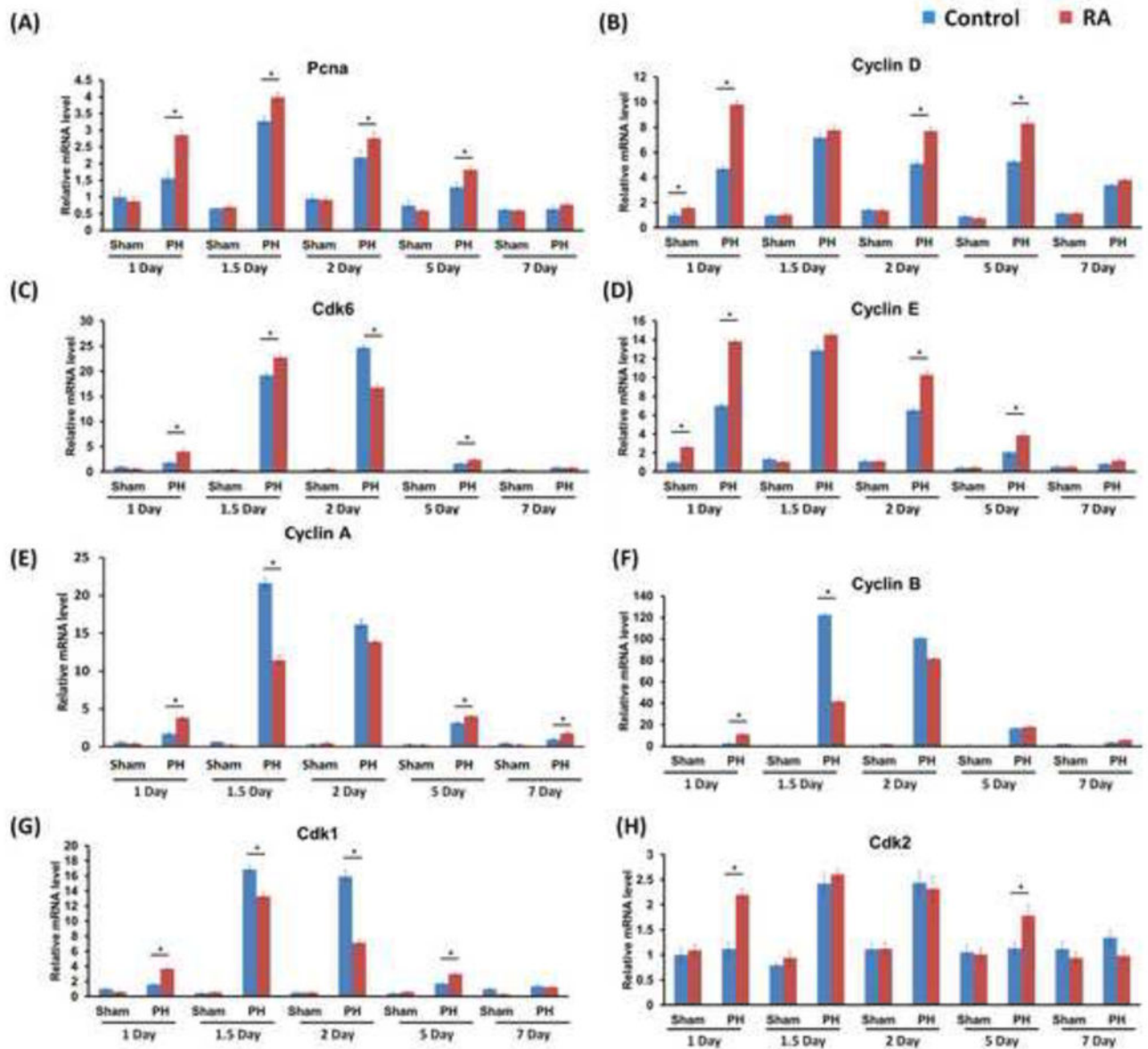


Figure 6. RA up-regulated cell cycle gene expression during liver regeneration
 WT mice with or without RA treatment received PH and were killed at indicated time points. The expression of cell cycle genes was studied. (A) mRNA levels of *Pcna*, an indicator of DNA synthesis, were consistently higher in the RA treatment group compared to the control group from 1 to 5 days after PH. (B–G) mRNA levels of Cyclins and Cdks were measured by quantitative RT-PCR. In the control group, characteristic induction of these key cell cycle genes is observed between 1.5–2 days after PH. RA treatment led to an earlier induction of cyclins and cdks 1 day after PH compared to control. Means \pm SD are graphed with * indicating $p < 0.05$.

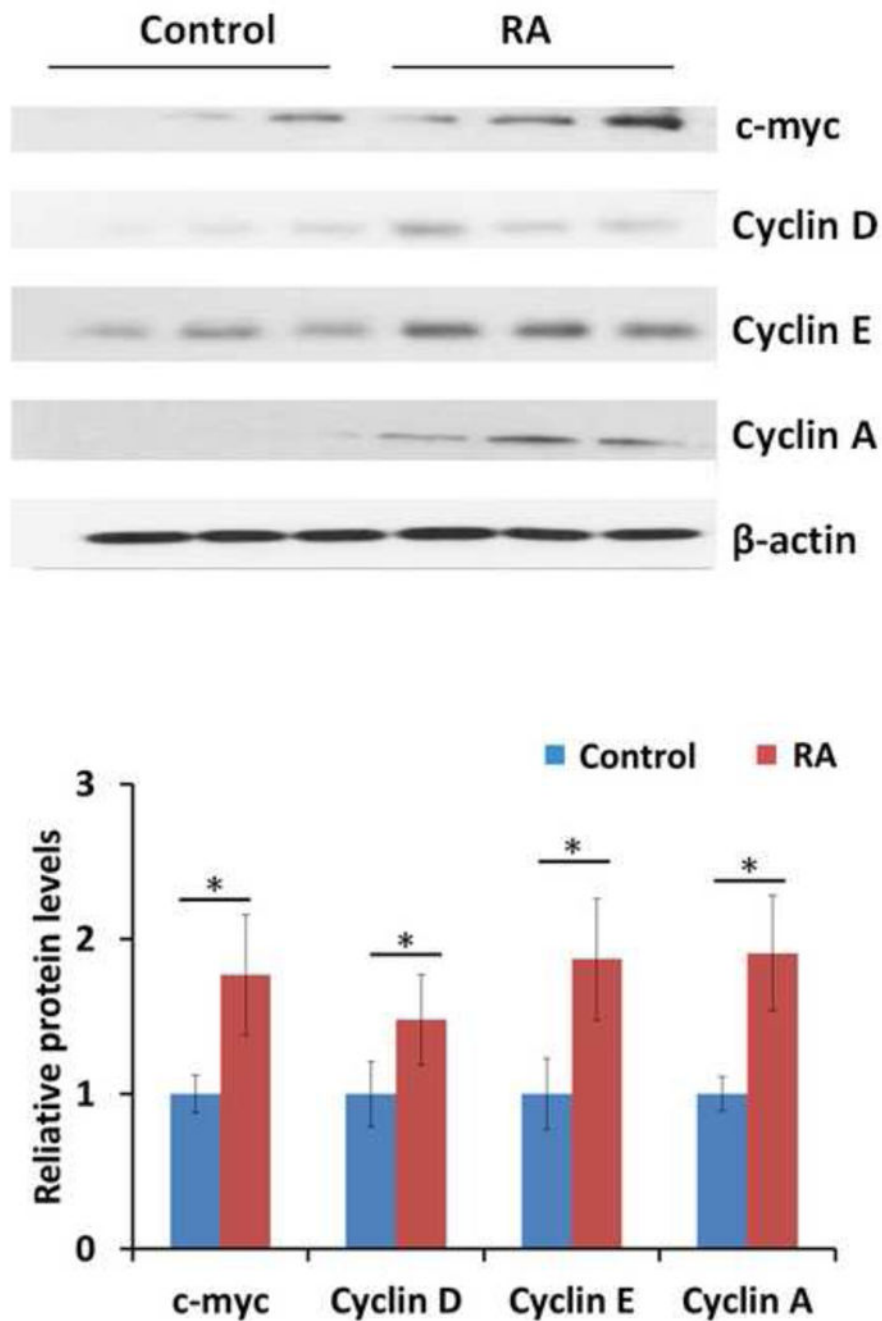


Figure 7. Effect of RA on cell cycle protein expression during liver regeneration

WT mice with or without RA treatment received PH and were killed 1 day after surgery. Western blots showed hepatic protein levels of c-myc, Cyclin D, Cyclin E, and Cyclin A in both groups. The protein levels were quantified by densitometry. Values for individual mouse were normalized to β -actin levels and were expressed relative to untreated mice. RA treatment led to higher hepatic induction of c-myc, Cyclin D, Cyclin E, and Cyclin A 1 day after PH compared to control. Means \pm SD are graphed with * indicating $p < 0.05$.