Hepatocytes lacking thioredoxin reductase 1 have normal replicative potential during development and regeneration

MaryClare F. Rollins1,*, Dana M. van der Heide2,*, Carla M. Weisend1, Jean A. Kundert3, Kristin M. Comstock4, Elena S. Suvorova1, Mario R. Capecchi5, Gary F. Merrill6 and Edward E. Schmidt1,7,‡

1Veterinary Molecular Biology, Montana State University, Bozeman, MT 59718, USA
2Biology Department, Oberlin College, Oberlin, OH 44074, USA
3Animal Resources Center, Montana State University, Bozeman, MT 59718, USA
4Biology Department, The College of St Scolastica, Duluth, MN 55811, USA
5Howard Hughes Medical Institute, University of Utah, Salt Lake City, UT 84118, USA
6Department of Biochemistry and Biophysics, Oregon State University, Corvallis, OR 97331, USA
7Center for Reproductive Biology, Washington State University, Pullman, WA 99164, USA

*These authors contributed equally to this work
‡Author for correspondence (eschmidt@montana.edu)

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Summary
Cells require ribonucleotide reductase (RNR) activity for DNA replication. In bacteria, electrons can flow from NADPH to RNR by either a thioredoxin-reductase- or a glutathione-reductase-dependent route. Yeast and plants artificially lacking thioredoxin reductases exhibit a slow-growth phenotype, suggesting glutathione-reductase-dependent routes are poor at supporting DNA replication in these organisms. We have studied proliferation of thioredoxin-reductase-1 (Txnrd1)-deficient hepatocytes in mice. During development and regeneration, normal mice and mice having Txnrd1-deficient hepatocytes exhibited similar liver growth rates. Proportions of hepatocytes that immunostained for PCNA, phosphohistone H3 or incorporated BrdU were also similar, indicating livers of either genotype had similar levels of proliferative, S and M phase hepatocytes, respectively. Replication was blocked by hydroxyurea, confirming that RNR activity was required by Txnrd1-deficient hepatocytes. Regenerative thymidine incorporation was similar in normal and Txnrd1-deficient livers, further indicating that DNA synthesis was unaffected. Using genetic chimeras in which a fluorescently marked subset of hepatocytes was Txnrd1-deficient while others were not, we found that the multigenerational contributions of both hepatocyte types to development and to liver regeneration were indistinguishable. We conclude that, in mouse hepatocytes, a Txnrd1-independent route for the supply of electrons to RNR can fully support DNA replication and normal proliferative growth.

Key words: Ribonucleotide reductase, Liver regeneration, Partial hepatectomy, DNA precursors, Cell cycle, Replication, Proliferation

Introduction
Conversion of ribonucleotides into deoxyribonucleotides by ribonucleotide reductase (RNR) is essential for genomic DNA replication (Arner and Eriksson, 1995; Holmgren, 1981; Nicander and Reichard, 1983; Thelander and Reichard, 1979). Thioredoxin was first discovered as a source of electrons for RNR in E. coli (Laurent et al., 1964) and thioredoxin reductase was identified as the flavin-containing enzyme that used electrons from NADPH to restore the reduced state of oxidized thioredoxin (Moore et al., 1964). Subsequent studies showed that a glutathione-reductase-dependent source of electrons could also support RNR activity and DNA replication. In this pathway, glutathione reductase transfers electrons from NADPH to glutathione, glutathione reduces glutaredoxin, and glutaredoxin donates the electrons to RNR (Holmgren, 1976; Miranda-Vizuete et al., 1994; Miranda-Vizuete et al., 1996).

Similar to bacteria, yeast and plants can survive in the absence of a functional thioredoxin-thioredoxin reductase pathway by using the glutathione-reductase-dependent source of electrons to fuel RNR (Koc et al., 2006; Meyer et al., 2009; Reichheld et al., 2007). Biochemical studies showed that mammalian RNR can also obtain electrons from the glutathione-reductase-dependent pathway (Luthman et al., 1979; Luthman and Holmgren, 1982) and a recent kinetic study showed that either route supports similar RNR enzymatic efficiency in vitro (Avval and Holmgren, 2009). Thus, it appears that most life forms might be able to use either thioredoxin reductase- or glutathione-reductase-dependent pathways to drive RNR-dependent DNA precursor production. However, none of the mammalian biochemical systems used to date have been able to approach the rate of RNR activity that would be required to support S phase replication in vivo (Avval and Holmgren, 2009), and the effectiveness of either source is not necessarily equivalent in all species (Arner, 2009; Fernandes and Holmgren, 2004).

Although the electrons supplied to RNR by either the thioredoxin reductase- or the glutathione-reductase-dependent route can support DNA replication, eukaryotic systems studied to date tended to favor the thioredoxin-reductase-dependent pathway (Arner, 2009; Arner and Holmgren, 2006a; Berndt et al., 2007; Lu and Holmgren, 2009). For example, yeast lacking thioredoxins exhibit a reduced rate of proliferation, skewed deoxyribonucleotide pools, and a protracted S phase, consistent with the glutathione-reductase-dependent pathway being a poor provider of DNA precursors (Koc et al., 2006). Measurements of the in vivo redox state of RNR...
verified that thioredoxin is normally the major direct provider of electrons for RNR in yeast (Camier et al., 2007). Similarly, plants lacking thioredoxin reductases show a slow-growth phenotype that is consistent with compromised replication (Reichheld et al., 2007).

The biological world contains two known families of thioredoxin reductase enzymes (Arner, 2009). One, a smaller protein with high substrate fidelity for thioredoxin, is found in Archaea, eubacteria, plants, protists and fungi. The other is a larger enzyme that evolved independently from glutathione reductase, has higher substrate promiscuity, and has been found universally in metazoan animals (Aleksunes and Manautou, 2007; Arner, 2009; Lobanov et al., 2009; Lu and Holmgren, 2009; Meyer et al., 2009). It remains unclear why an ancestral pre-metazoan animal might have discarded its thioredoxin reductase genes (both cytoplasmic and mitochondrial forms of the enzyme) and generated a new lower fidelity family of enzymes from glutathione reductase; however, this event might underlie fundamental differences in the thioredoxin-reductase-pathway functions between animals and other life forms.

Previous studies suggested that, like yeast and plants, metazoans might also preferentially use their thioredoxin-reductase-dependent pathway to fuel RNR. Thus, although mice lacking glutathione reductase are viable (Rogers et al., 2004), mice lacking thioredoxin 1 (Txn1) (Matsui et al., 1996) or thioredoxin reductase 1 (Txnrd1) (Bondareva et al., 2007; Jakupoglu et al., 2005) are embryonic lethal. Lethality, however, does not necessarily implicate an RNR activity deficiency. Txn1 also participates in reducing protein disulfides and supplying electrons to homeostatic and antioxidant systems, including peroxiredoxins, methionine sulf oxide reductases and others (Arner, 2009; Fomenko et al., 2009; Kim and Gladyshev, 2007; Rundlof and Arner, 2004). We recently showed that disruption of Txnrd1 in mouse hepatocytes or fibroblasts results in chronic induction of cytoprotective pathways, which are thought to compensate for the loss of Txnrd1 in homoeostatic antioxidant functions (Suvorova et al., 2009). Embryonic lethality in Txnrd1-deficient mouse embryos occurs only after replication has generated several thousand (Bondareva et al., 2007) or more (Jakupoglu et al., 2005) Txnrd1-deficient cells, again suggesting that functions other than DNA precursor production underlie lethality. Similarly, disruption of the genes encoding either mitochondrial thioredoxin (Txn2) (Nonn et al., 2003) or mitochondrial thioredoxin reductase (Txnrd2) (Conrad et al., 2004) are also embryonic lethal despite these proteins being generally located in a compartment (mitochondria) separate from that of ribonucleotide reduction for genomic DNA replication (cytosol) (Arner, 2009).

Our recent analyses of mice having hepatocyte-specific disruption of the Txnrd1 gene showed strikingly normal liver development, including apparently normal levels of endoreplication and acetykinetic mitosis in mature Txnrd1-deficient hepatocytes (Suvorova et al., 2009). Also, although RNAi knockdown of Txnrd1 in mouse cancer cell cultures inhibits self-sufficient growth and other cancer-related characteristics of the cells, it does not block proliferation (Yoo et al., 2006; Yoo et al., 2007). These studies could not rule out the possibility that replication was supported by Txnrd1 enzyme remaining from maternal (Bondareva et al., 2007; Jakupoglu et al., 2005), pre-hepatocytic (Suvorova et al., 2009) or residual (Yoo et al., 2006; Yoo et al., 2007) sources; however, they provided the first in vivo evidence to suggest that Txnrd1-deficient mammalian cells might be able to undergo genome replication. To determine if this was, indeed, true, we evaluated the requirement of Txnrd1 for replication in hepatocytes.

In the current study, we tested whether hepatocytes in mouse liver were able to proliferate in the absence of Txnrd1. Experimental systems were designed in which cells had been Txnrd1-deficient for extended periods and, in some cases, for many cell cycles, to rule out any possibility that persisting pre-formed Txnrd1 supported replication. Results showed that Txnrd1−/− hepatocytes were able to proliferate and, unlike proliferation in Txnrd-deficient yeast or plants (Koc et al., 2006; Reichheld et al., 2007), hepatocyte proliferative indexes were not measurably affected by loss of Txnrd1. Our results indicate that the Txnrd1-independent route supplying electrons for RNR can support normal replication in a mammalian system. The implications of this finding should be considered in therapies using drugs that inhibit or otherwise affect Txnrd1 activity.

Several different abbreviations are currently used for thioredoxin reductases, including Txnrd, TxnR, TR and TRR. We have used the abbreviations for mouse enzymes recommended by the National Center for Biotechnology Information (NCBI) and the Mouse Genome Informatics database.

Genetic designations used in this paper are as follows. Gene loci are italicized and separated by a semicolon. Alleles are designated as superscript and separated by a slash. Transgenes are followed by a superscript to indicate whether they are homozygous present (‡), hemizygous (†) or absent (†). For example, Txnrd1+/−;AlbCre‡ is heterozygous for a null (−) allele of Txnrd1 and does not carry an AlbCre transgene.

**Results**

**Txnrd1-deficient hepatocyte populations expand and replicate DNA during liver development**

We previously established a mouse (Mus musculus, strain C57Bl/6J) system in which all differentiated hepatocytes lack Txnrd1 (Suvorova et al., 2009). Briefly, female animals that are homozygous for the functionally wild-type yet Cre-inactivatable (‘conditional’) Txnrd1cond/− allele (Bondareva et al., 2007) and for a two-color Cre-responsive reporter gene, ROSA²mG (Muzumdar et al., 2007) (genotype Txnrd1cond/−;ROSAmG/+;AlbCre‡) were mated with males that were heterozygous for the null allele of Txnrd1 (Bondareva et al., 2007) and carried two copies of an albumin promoter/enhancer-driven Cre transgene (Postic et al., 1999) (genotype Txnrd1+/−;AlbCre‡). Offspring were either Txnrd1+/−;ROSAmG/+;AlbCre‡ (hereafter referred to as experimental) or Txnrd1+/−;ROSAmG/+;AlbCre† (control). The AlbCre transgene drives Cre expression in all differentiated hepatocytes (Postic and Magnuson, 2000; Postic et al., 1999) including those that arise in fetal stages (Weisend et al., 2009). The ROSA²mG allele, in its non-recombined state, drives strong membrane-targeted accumulation of the red fluorescent protein tdTomato in all cell types (Muzumdar et al., 2007). Exposure to Cre deletes the tdTomato cistron and uncovers a cistron that yields a membrane-associated modified green fluorescent protein (GFP). Thus, in experimental animals, all cells are Txnrd1cond/− and red fluorescent, except differentiated hepatocytes, which are Txnrd1+/− and green fluorescent (Suvorova et al., 2009). Previous studies showed that livers from these mice expressed almost no functional Txnrd1 mRNA or protein, suggesting all hepatocytes completely lacked Txnrd1 (Suvorova et al., 2009). In control animals all cells are red fluorescent and Txnrd1cond/− except differentiated hepatocytes, which are green-fluorescent and Txnrd1+/− (Suvorova et al., 2009). In these studies as well as in previous studies, we have been unable to measure a difference in phenotype between
**Contributions of Tnfrd1-deficient hepatocytes versus Tnfrd1-containing progenitor cells to liver regeneration**

To induce global proliferation of differentiated hepatocytes, we performed partial hepatectomies on adult mice. Following surgical removal of two-thirds of normal rodent liver, most differentiated hepatocytes rapidly enter the cell cycle and proliferate to regenerate liver mass (Higgins and Anderson, 1931; Martins et al., 2008; Mitchell and Willenbring, 2008; Watanabe, 1970). DNA synthesis peaks around 36 hours post-hepatectomy as the first semi-synchronous wave of hepatocytes transits S phase, and proliferation remains elevated until regeneration is complete, after 7-10 days (Mitchell and Willenbring, 2008). At hepatectomy, experimental liver masses were ~24% greater than control liver masses; rates of liver mass accumulation following a two-thirds hepatectomy were similar in experimental and control mice (Fig. 3A), suggesting that Tnfrd1-deficient livers were not compromised in regenerative tissue growth.

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**Fig. 1. Hepatocyte-specific allelic conversion in mouse liver by AlbCre.**

(A) Low-magnification micrograph showing merged fluorescence and DAPI staining of a cryosection of liver from an adult (P42) animal of genotype Tnfrd1cond/cond;ROSA^tm2mG/AlbCre. All hepatocytes in this image, identified as large and generally cuboidal cells with abundant cytoplasm with large heterogeneous weakly DAPI-stained nuclei, have green fluorescent membranes, which indicates that AlbCre efficiently directed Cre-mediated allelic conversion in hepatocytes. All non-hepatocyte cells, in particular the endothelial cells, identified as very small irregularly shaped cells with small strongly DAPI-stained nuclei, which form tubular networks surrounding blood vessels and capillaries, exhibited red fluorescence. Note that the intensity of green fluorescence varies zonally as a function of the relative distance from portal (P; arterial) versus venous (V) circulation. Because the intensity of red fluorescence in non-hepatocyte cells did not vary similarly, the peri-venous zones appear darker and redder. We are uncertain of the causes of this hepatocyte-specific zonal fluorescence variation, although it might reflect zonal differences in metabolic activity of hepatocytes resulting from natural gradients in oxygen tension or nutrient availability. Careful examination revealed that nearly all hepatocytes in all zones had exclusively green membranes.

(B) High-magnification fluorescence micrographs of a DAPI-stained liver cryosection from an aged adult (P280) animal of genotype Tnfrd1cond/cond;ROSA^tm2mG/AlbCre. The red channel (left) shows endothelial membranes surrounding capillaries. The green channel (middle) shows hepatocyte membranes. The merged channel (right) shows that hepatocyte-endothelial interfaces have both red and green membranes (white arrows), whereas hepatocyte-hepatocyte interfaces have only green membranes (yellow arrows). Scale bars: 50 μm.
Liver growth can occur by either proliferation of differentiated hepatocytes or proliferation of pre-hepatocytic progenitor cells followed by differentiation into hepatocytes (Oertel and Shafritz, 2006). Because allelic conversion was driven by AlbCre, which is induced coincident with hepatocyte differentiation (Postic et al., 1999; Weisend et al., 2009), proliferating progenitor cells would contain Txnrd1. To distinguish contributions of Txnrd1-containing pre-hepatocytic progenitor cells from proliferation of Txnrd1-deficient differentiated hepatocytes, we developed a novel ‘ten-day chronometer’ of hepatocyte lineage age based on conversion of the \textit{ROSA}^{\text{mTmG}} allele by AlbCre (S. V. Iverson, K.M.C., C.M.W., J.A.K. and E.E.S., unpublished data). Briefly, activation of the AlbCre transgene during hepatocyte differentiation (Postic et al., 1999; Weisend et al., 2009) will simultaneously convert both the \textit{Txnrd1}^{\text{cond}} allele and the \textit{ROSA}^{\text{mTmG}} allele (Suvorova et al., 2009; Weisend et al., 2009). Cre-mediated recombination of \textit{ROSA}^{\text{mTmG}} initiates a progressive conversion in membrane fluorescence from red to green as GFP accumulates and tdTomato protein decays. Fluorescent marker conversion in hepatocyte membranes required roughly 10 days to complete (Muzumdar et al., 2007) and data not shown). Therefore, 48 hours after partial hepatectomy, hepatocytes arising from pre-hepatocytic progenitor cells (0- to 48-hours post-onset of Cre expression) will exhibit low ratios of green/red fluorescence and could be readily distinguished from the purely green hepatocytes arising from proliferation of preexisting hepatocytes. At this time point, nearly all hepatocyte membranes exhibited purely green fluorescence (Fig. 3B). At low incidences, fields of view in some peri-portal regions could be found with newly differentiated hepatocytes [i.e. retaining some red membrane fluorescence; Fig. 3C, orange arrows denote younger (redder) hepatocytes than the light-green arrows], verifying that contributions of progenitor cells to liver regeneration could be detected using this system. Importantly, frequencies of newly differentiated hepatocytes were low during regeneration (~1% of total hepatocytes) and did not differ significantly between experimental and control livers (Fig. 3C,D). Most hepatocytes 2 days after partial hepatectomy arose from cell lineages that had differentiated into hepatocytes prior to hepatocyte; pre-hepatocytic progenitor cells were only minor contributors.

Thioredoxin-deficient yeast are able to proliferate; however they exhibit a protracted S phase and a prolonged cell cycle (Koc et al., 2006). Static frequencies of S phase hepatocytes cannot, alone, indicate either differences in cell cycle duration or in S phase duration; however, by measuring multiple static proliferative indexes representing different cell cycle phases, one can more accurately determine whether the cell cycle is perturbed (supplementary material Fig. S1). To test whether disruption of \textit{Txnrd1} in hepatocytes affected cell cycle progression, we measured the proliferative index by PCNA staining, the S phase index by BrdU labeling, and the M phase index by staining for PHH3 (Fig. 4A). Results showed no measurable differences in any of these proliferative indexes between hepatocytes within regenerating control and experimental livers (Fig. 4B).

Based on previous studies in lower eukaryotes (Koc et al., 2006; Reichheld et al., 2007), it was surprising that mouse hepatocyte proliferation was not measurably affected by disruption of \textit{Txnrd1}. As a control, we confirmed that the RNR inhibitor hydroxyurea entirely prevented regenerative replication of both normal and \textit{Txnrd1}-deficient hepatocytes (Fig. 5A). Also, in vivo post-hepatectomy metabolic labeling showed that thymidine incorporation during liver regeneration did not differ between normal and \textit{Txnrd1}-deficient livers (Fig. 5B). In combination, the data above show that proliferative liver growth was driven primarily by hepatocyte proliferation, not by proliferation of \textit{Txnrd1}-containing progenitor cells, and they indicate that the ability of RNR to support hepatocyte DNA replication was not measurably compromised by disruption of \textit{Txnrd1}.

**Contributions of \textit{Txnrd1}-containing versus \textit{Txnrd1}-deficient cell lineages to development and regeneration of genetically chimeric livers**

For a potentially more sensitive comparison of relative proliferative potentials, we generated fetally induced genetically chimeric mice in which individual livers contained both red-fluorescent-tagged...
normal and green-fluorescent-tagged Tnrd1-deficient hepatocytes. Using these mice, we evaluated the contributions of both hepatocyte types to developmental and regenerative liver growth. By this approach, the size of the adult liver can be seen as a target niche that the two types of hepatocytes will compete to populate. This results in a ‘multi-generational relay-race’ in which cells with a subtle genetic growth advantage will pass this advantage to their daughter cells, thus amplifying the proliferative difference each generation. To produce the mosaics, we used the Tamoxifen-inducible ROSACreER allele (Badea et al., 2003). Dams of genotype Tnrd1cond/cond;ROSAmT-mG/mT-mG were mated to sires of genotype Tnrd1cond/cond;ROSACreER/CreER. In this mating, the dams themselves lack the ROSA<sup>CreER</sup> allele and do not exhibit Cre activity in response to Tamoxifen; however, all progeny are genotype Tnrd1cond/cond;ROSAtmG/CreER so that fetal cells exposed to Tamoxifen can convert to Tnrd1<sup>−/−</sup> and from red to green fluorescence. A time and dose study established that intraperitoneal (IP) administration of 0.5 mg of 4-hydroxytamoxifen (4OHT) to pregnant dams at embryonic day 14.5 (E14.5) resulted in 100% survival of pups to term (C.M.W., J.A.K. and E.E.S., unpublished data). By this dosage protocol, pups uniformly exhibited 30-50% green (Tnrd1<sup>−/−</sup>) hepatocytes as neonates (Fig. 6A and not shown), whereas frequencies of green cells in other organs were much lower (ranging from a low of <0.1% in brain to a high of ~10% in pancreas, with most organs ~1%; C.M.W. and E.E.S., unpublished data).

Following maternal administration of 4OHT at E14.5, cryosections of livers harvested from neonatal [postnatal day (P) 6], juvenile (P14-P28), young adult (P35) and older adult (P102) offspring showed similar mosaic distributions of ~50-70% red
replication of Txnrd1-deficient hepatocytes during liver regeneration. Adult (P42) mice of genotypes $\text{Txnrd1}^{\text{cond/+}};\text{AlbCre}^1$ (left panels, controls) or $\text{Txnrd1}^{\text{cond/-}};\text{AlbCre}^1$ (right panels, experimentals) were two-thirds hepatectomized. For PCNA (upper panels) and PHH3 (middle panels) staining, livers were harvested 48 hours post-hepatectomy when the first ‘wave’ of proliferative hepatocytes would be around G2-M in the cell cycle. The abundant hepatocytes with diffuse cytoplasmic PCNA staining (upper panel) are presumably in M or early G1 phase, after nuclear membrane dissolution allows PCNA protein to redistribute into the cytoplasm. Hepatocytes with nuclear or cytoplasmic staining were scored as proliferative, as both types allows PCNA protein to redistribute into the cytoplasm. For BrdU staining (lower panel, pink arrow) were found in Txnrd1-deficient (green) hepatocytes, verifying that Txnrd1-deficient hepatocytes had proliferative potential.

To more accurately estimate how many consecutive cell cycles hepatocytes had completed, we quantified the cellular dynamics of liver development. Fresh whole livers harvested from E14.5 fetal mice from our colony weighed 0.0315 ± 0.0032 g (mean ± s.e.m., $n=6$). Previously, we showed that E14.5 liver is composed of <10% differentiated hepatocytes by mass (Weisend et al., 2009). In adults, the liver weighs ~1.6 g (Fig. 3A) and is composed of ~95% differentiated hepatocytes by mass (Schmidt and Schibler, 1995; Weisend et al., 2009). Thus, there is a roughly 500-fold increase in mass of total liver hepatocytes during development.

Although a portion of this is accounted for by the increase in average liver cell size that occurs during rodent development (~fivefold during post-natal development in rats (Schmidt and Schibler, 1995)), we estimate that there is also a 30- to 100-fold developmental increase in the number of hepatocyte genomes in mice, which accounts for the majority of this growth. This increase in hepatocyte genomes, in the absence of any cell death, would require five to seven consecutive cell doublings ($2^{5}=32; 2^{7}=128$). Since relative representation of both the E14.5-converted $\text{Txnrd1}^{-/-}$ (green) and the non-converted $\text{Txnrd1}^{\text{cond/cond}}$ (red) hepatocytes was similar in mosaic juvenile and adult livers (Fig. 6A,B), we concluded that, by adulthood the cell lineages leading to both hepatocyte types had undergone a minimum of five to seven consecutive replicative cycles since their Cre-mediated genetic divergence without showing a measurable difference in proliferative potential.
Next, we tested whether adult Txnrd1-deficient hepatocytes that arose from lineages that had been converted at E14.5 retained the same capacity as normal hepatocytes to contribute to the rapid proliferative burst that follows a two-thirds hepatectomy (i.e. 1–1.5 more replicative cycles). As modeled in Fig. 6C, we predicted that, if the Txnrd1-deficient hepatocytes could contribute to regeneration, then the ratio of red to green hepatocytes would be similar before and after hepatectomy; if not, we would observe an effective dilution of green cells. Results showed that the relative representation of red and green cells was similar before and after regeneration (Fig. 6D). Moreover, at 72 hours post-hepatectomy, mitotic figures were found in Txnrd1-deficient (green) cells, verifying these cells were engaged in the cell cycle (Fig. 6D, white arrows). We conclude that normal and Txnrd1-deficient mouse hepatocytes have equivalent developmental and regenerative proliferative potentials.

Discussion

Txnrd1 disruption, selenium assimilation, and the hepatocyte selenoproteome

Mammalian Txnrs are selenoproteins, meaning they contain the atypical amino acid selenocysteine (Sec) (Gladyshev and Hatfield, 1999; Lobanov et al., 2009; Lu and Holmgren, 2009). Only 25 mammalian selenoproteins are known or predicted, all of which are oxidoreductases and all of which have Sec participating in active site electron transfer (Fomenko et al., 2007; Lobanov et al., 2007; Lobanov et al., 2009; Lu and Holmgren, 2009). Production of Sec, selenide is either acquired from organic dietary sources or as inorganic selenate and selenite. Dietary selenite is reduced to selenide by either the glutathione reductase (Gsr)-reduced glutathione-glutaredoxin or the Txnrd-Txn systems, whereas the mechanisms of selenate reduction remain uncertain (Lu et al., 2009; Lu and Holmgren, 2009). We have not tested whether disruption of Txnrd1 has global effects on the selenoproteome within individual cells, although the apparent redundancy of sources and reductase systems for mobilization of dietary selenium suggests Sec availability would not become limiting in Txnrd1-deficient cells.

Alternative pathways of supplying electrons to RNR

In all life forms studied to date, RNR activity is essential for DNA replication. RNR activity, in turn, is dependent on a source of electrons. Genetic malleability of bacterial, yeast and plant systems allowed identification of both thioredoxins and glutaredoxins as the electron donors for RNR in these systems; thioredoxins and glutaredoxins ultimately obtain the electrons from NADPH via thioredoxin reductase or via glutathione reductase and glutathione, respectively (Avval and Holmgren, 2009; Meyer et al., 2009).
Previously, it had not been possible to test the relative contributions of these two pathways in vivo in mammals. It has long been known that mammalian glutaredoxin can provide electrons to RNR in vitro (Luthman et al., 1979; Luthman and Holmgren, 1982) and a recent study showed that both Txn1 and glutaredoxin are similarly efficient at supplying electrons to RNR in vitro (Avval and Holmgren, 2009). None of these biochemical assays could approach rates of deoxyribonucleotide production required during S phase, however, and no previous study definitively established that mammalian cells could replicate DNA and proliferate in the absence of Txnrd1. The survival and relatively normal physiology of mice lacking Gsr indicates that the Gsr-dependent pathway is not necessary in mammals (Rogers et al., 2004). Conversely, embryonic lethality of mice lacking either Txn1 (Matsui et al., 1996) or Txnrd1 (Bondareva et al., 2007; Jakupoglu et al., 2005) indicates that some aspect of the Txnrd1-dependent pathway is essential in mammals.

Although mice having a disrupted Txnrd1-Txn1 system are not viable, several lines of evidence suggested Txnrd1 is not essential for cell proliferation. First, cells in Txnrd1-deficient mouse embryos underwent at least twelve consecutive doublings prior to embryonic arrest (Bondareva et al., 2007; Jakupoglu et al., 2005). Second, knockdown of Txnrd1 in cell cultures did not affect cell proliferation (Yoo et al., 2006; Yoo et al., 2007). Third, mice lacking Txnrd1 in all differentiated hepatocytes have normal or slightly larger than normal sized livers (Fig. 3A) (Suvorova et al., 2009). It was possible that proliferation in these systems used cryptic sources of Txnrd1 (i.e. maternally derived in embryos, progenitor cell-derived in hepatocytes or residual following knockdown). Alternatively, a Txnrd1-independent pathway might have been supporting replication.

The most obvious alternative pathway for supporting RNR activity in Txnrd1-deficient hepatocytes would be the Gsr-dependent system that has been described in other eukaryotes and demonstrated in vitro. Alternatively, however, the mitochondrial Txnrd2 enzyme might be compensating for loss of Txnrd1. Although ribonucleotide reduction for genome replication occurs in the cytosol and there is no evidence of a mitochondrial source of electrons for this reaction (Arner, 2009; Arner and Holmgren, 2006a; Meyer et al., 2009), the Txnrd2 gene contains two alternative promoters and first exons capable of issuing mRNAs that would each encode cytosolic isoforms of Txnrd2 (Turanov et al., 2006). Accumulation of endogenous cytosolic Txnrd2 has not, to our knowledge, been shown. However, naturally occurring mRNAs encoding these isoforms have been described, and overexpression of mimgenes encoding these isoforms fused to GFP in cultured cells results in cytosolic accumulation of the fluorescent Txnrd2-GFP fusion proteins (Turanov et al., 2006). Moreover, in vitro, recombinant Txnrd2 can reduce Txn1 (Turanov et al., 2006) and diminution of Txnrd1 activity in cell cultures by RNAi or inhibitory drugs does not lead to Txn1 oxidation (Watson et al., 2008).

The conditional hepatocyte-specific Txnrd1-deficient system provided a model in which prolonged temporal separation of gene disruption and proliferation could be achieved in vivo, thus eliminating the possibility that persistent pre-formed Txnrd1 was supporting replication. In the current study, we used this system to show that disruption of Txnrd1 had no measurable effect on the proliferative potential of hepatocytes in mice. Our results, combined with the normal physiology of Gsr-null mice (Rogers et al., 2004), lead us to conclude that, similar other life forms, animal cells can probably proliferate using either Txnrd- or Gsr-derived electrons to fuel RNR. Moreover, unlike yeast (Camier et al., 2007; Koc et al., 2006) and perhaps plants (Reichheld et al., 2007), we detected no evidence that the Txnrd1-dependent system was a preferential source of electrons. It is possible that this difference between species relates to the distinct evolutionary origins of metazoan compared with other eukaryotic thioredoxin reductases. Further studies will be required to distinguish the possible respective roles of Gsr versus cytosolic forms of Txnrd2 in supporting replication; however, intriguingly, the transcriptions of both Txnrd1-deficient embryos (Bondareva et al., 2007) and livers (Suvorova et al., 2009) do not show any compensatory upregulation of mRNAs encoding either Gsr or Txnrd2. Thus, whatever pathway is supporting RNR activity during hepatocyte replication it might be sufficiently abundant and robust to fully support genomic DNA replication without further induction, and thus might be a major, as opposed to an alternative or compensatory, source of reducing potential for RNR. Although there might be some cell types or circumstances with different requirements, we find that either the Txnrd1-dependent or Txnrd1-independent route is probably equivalent in terms of supporting physiological RNR activity in hepatocytes.

Other roles and functions of the Txnrd1-Txn1 system

In addition to providing electrons to RNR, the Txnrd1-Txn1 system is a major protein thiol reductase; it provides reducing potential to enzymes including peroxiredoxins and methionine sulfoxide reductase, and it is generally considered to be a major participant in cellular antioxidant defense and redox regulation (Arner, 2009; Arner and Holmgren, 2006a; Meyer et al., 2009). We previously showed that disruption of Txnrd1 in embryos or hepatocytes resulted in a strong cytoprotective response, including induction of drug metabolism genes and upregulation of Nrf2 stress-response pathway (Bondareva et al., 2007; Suvorova et al., 2009). It is unclear how these responses might compensate for loss of Txnrd1 in cells. However, in the mutant livers where biochemical analyses could be performed, we observed no evidence of oxidative stress, suggesting this response was, indeed, compensatory (Suvorova et al., 2009). We are aware of no evidence that any of the cytoprotective pathways induced in Txnrd1-deficient hepatocytes could directly supply electrons to RNR.

Implications for Txnrd- and Txn-targeted pharmaceuticals

Considerable attention has been given to possible roles for Txnrd1, both favorable and deleterious, in cancer initiation and progression (Arner, 2009). Whereas Txnrd1 activity is thought to protect against the initiation of cancers (Arner, 2009; Arner and Holmgren, 2006b; Ganther, 1999), many aggressive cancers exhibit increased expression of Txnrd1 (Berggren et al., 1996; Nakamura et al., 2000; Rundlof and Arner, 2004), suggesting Txnrd1 activity supports cancer growth and progression (Arner, 2009; Arner and Holmgren, 2006a). Txnrd1 has been extolled as a ‘drugable target’ for cancer therapies (Arner, 2009; Arner and Holmgren, 2006a; Bindoli et al., 2009; Gandin et al., 2009). Numerous cancer drugs or drug candidates affect Txnrd1 activity, either as a primary target or, as in the case of cisplatin and other platinum-containing compounds (Sasada et al., 1999; Witte et al., 2005), as a collateral target. Some of these include arsenic trioxide (Lu et al., 2007), various organo-gold compounds (Bindoli et al., 2009; Casini et al., 2009; Coronnello et al., 2005; Gandin et al., 2009; Marzano et al., 2007; Rigobello et al., 2009), some mercury-containing compounds (Carvalho et al., 2008), and certain quinols (Chew et al., 2008),...
Materials and Methods

Mice and animal care

All animal procedures were approved by the Montana State University Institutional Animal Care and Use Committee (MSU-IACUC). All renewable resources in this study are available on request for unrestricted non-profit research unless regulated by another party. All mice were housed with specialized care conditions including sterilized food, water and bedding in HEPA-filtered air-conditioned (Tecniplast) under a 14 hour:10 hour light:dark cycle. Feed (PicoLab mouse diet 5058, which contains selenium derived as a natural component of the grains and supplemented to 0.05 ppm with selenium) and water were unrestricted. All mutant alleles were followed by PCR analyses of genomic DNA samples using the primers and conditions described previously (Bondareva et al., 2007; Suvorova et al., 2009; Weisend et al., 2009). Mouse lines (Gt(Rosa)26Sortm4(ACTB-tdTomato,-EGFP)Luo/+) (here abbreviated ‘Rosa26CreERT2’) (Muzumdar et al., 2007), B6;129-Gt(Rosa)26Sortm4(ACTB-tdTomato,-EGFP)Luo/+(‘Rosa26CreERT2’) (Badea et al., 2003), B6.Cg-Tg(Alb-Cre)22MgnJ (‘AlbCre’) (Postic et al., 1999), and C57Bl/6J were obtained from Jackson Labs (stock numbers 007576, 004847, 003574 and 000664, respectively) or from a Nikon Eclipse E800 or an Olympus BX60 microscope. For all immunofluorescences, the whole area shown was uniformly adjusted using the ‘Autolevels’ function in Photoshop CS3 software. Some micrographs were electronically enlarged or reduced in Photoshop CS3 software. Scale bars in figures were set by photographing a non-standard slide rule under each microscope used and simply noted to the same electronic enlargement or reduction as the biological images shown. For non-merged immunofluorescences, the whole area shown was adjusted using the ‘Autocontrast’ function in Photoshop CS3. With the exception of the images in Fig. 3C, all micrographs were representative of the whole organ. In the case of Fig. 3C, newly differentiated hepatocytes were clustered around widely scattered peri-portal regions. Images were taken to show these clusters, as arbitrarily chosen fields generally contained no red hepatocytes (e.g. Fig. 1A and Fig. 3B). All adjustments and manipulations made to images were performed uniformly to the entire image. None of these adjustments qualitatively or quantitatively affect the interpretations or conclusions arising from the data, but instead serve useful aesthetic functions and ease visualization and enumeration of important biological characteristics of each sample.

Nuclei isolation and DNA extraction for thymidine labeling studies

Following thyidine labeling, animals were killed by CO2 asphyxiation and livers were harvested into ice-cold PBS. Nuclei were isolated using protocols described previously (Schmidt and Schibler, 1995). Briefly, liver was blotted, weighed, minced and homogenized under final conditions of 10% (w/v) 1.8 M sucrose/20% (v/v) glycerol, 10 mM HEPES, pH 7.8, 15 mM KC1, 0.15 mM spermine, 0.5 mM spermidine, 1 mM EDTA, in a motor-driven Teflon and glass homogenizer. Lysate was layered on 0.5 ml cushions of 2.0 M sucrose, 10% glycerol, 10 mM HEPES, pH 7.8, 15 mM KC1, 0.15 mM spermine, 0.5 mM spermidine, 1 mM EDTA and were centrifuged at 116,000 g, 26,000 rpm in a SW-41 Ti rotor, 4°C, for 1 hour in an SW-41 rotor. Nuclei were resuspended in nucleus resuspension buffer (10% glycerol, 10 mM HEPES, pH 7.8, 15 mM KC1, 0.15 mM spermine, 0.5 mM spermidine, 1 mM EDTA), total nuclear acid content was determined, and concentrations were adjusted to 10 mg/ml nuclear acid with nuclear resuspension buffer. Samples equivalent to 0.0 (no input nuclei), background controls (0.1, 0.3 and 1.0 mg of nuclear acid were adjusted to 200 µl with nuclear resuspension buffer, boiled for 3 minutes, and cooled to room temperature. RNAses A and T1 were added to final concentrations of 10 µg/ml and 350 U/ml, respectively, and samples were incubated at 42°C for 30 minutes. Twenty percent SDS (10 µl) and 0.5 µl of 200 U/ml Papain/seminase K (Sigma) were then incubated at 50°C for 15 minutes. Samples were diluted with 800 µl of water, and high molecular mass DNA was precipitated by adding 120 µl of 100% (w/v) trichloroacetic acid, and the samples were mixed, and incubated on ice 30 minutes. Precipitated DNA was collected by vacuum filtration onto 0.5 cm glass-fiber filter discs. Discs were washed sequentially with a large volume of 10% trichloroacetic acid followed by 70% ethanol, and were assayed by liquid scintillation counting.

Statistical analyses

Statistical analyses used at least three biological replicates (i.e. from different animals) for each condition. Data in graph panels were presented as mean ± standard error of the mean (s.e.m.). For proliferative indexes, significance was determined using the Student’s t-test with a minimum P-value of 0.05. For [3H]thymidine incorporation (Fig. 6B), each sample (three input nuclei
concentrations from three different animals from each genotype) was counted for four 1-minute periods by liquid scintillation counting. Counts per minute values for each sample were averaged, and the average background (scintillation fluid only, no input nucleii) was subtracted. Data were plotted and a best-fit line was assigned by least-squares regression; line fit was evaluated by the coefficient of determination, $r^2$. Slope represents thymidine incorporation per input nucleus.

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References


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normal cell cycle

protracted G1

protracted S

M index             10                                      6.7    6.7
S index              50            30   67
M/S ratio             0.2 0.2      0.1