Activation of innate immunity by mitochondrial dsRNA in mouse cells lacking p53 protein

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**ABSTRACT**

Viral and cellular double-stranded RNA (dsRNA) is recognized by cytosolic innate immune sensors, including RIG-I-like receptors. Some cytoplasmic dsRNA is commonly present in cells, and one source is mitochondrial dsRNA, which results from bidirectional transcription of mitochondrial DNA (mtDNA). Here we demonstrate that Trp53 mutant mouse embryonic fibroblasts contain immune-stimulating endogenous dsRNA of mitochondrial origin. We show that the immune response induced by this dsRNA is mediated via RIG-I-like receptors and leads to the expression of type I interferon and proinflammatory cytokine genes. The mitochondrial dsRNA is cleaved by RNase L, which cleaves all cellular RNA including mitochondrial mRNAs, increasing activation of RIG-I-like receptors. When mitochondrial transcription is interrupted there is a subsequent decrease in this immune-stimulatory dsRNA. Our results reveal that the role of p53 in innate immunity is even more versatile and complex than previously anticipated. Our study, therefore, sheds new light on the role of endogenous RNA in diseases featuring aberrant immune responses.

Keywords: mitochondrial dsRNA; p53; innate immunity; RNase L

**INTRODUCTION**

Type I interferon (IFN) secretion is a first line of defense against viral pathogens in most mammalian cells. IFN production and secretion is activated after host-specific pattern recognition receptors (PRRs) detect pathogen-associated molecular patterns (PAMPs). A wide range of different types of molecules can serve as PAMPs, thus PRRs recognize specific types of ligands. MDA5 (melanoma differentiation-associated protein 5) and RIG-I (retinoic acid-inducible gene I) are two RIG-I-like receptors (RLRs), that detect double-stranded RNA (dsRNA), which is a replication intermediate for RNA viruses (Kang et al. 2002; Yoneyama et al. 2004). Nevertheless, those two receptors recognize different features of dsRNAs: RIG-I selectively binds ssRNA or short blunt-ended dsRNA (less than 1 kb) bearing uncapped 5′-di- or triphosphate whereas MDA-5 binds long dsRNA. Upon activation, both receptors interact via CARD domains with the mitochondrial antiviral signaling protein (MAVS). MAVS recruitment leads to nuclear translocation of the transcription factors IRF3 and NF-κB, and to the production of proinflammatory cytokines, chemokines and type I IFN and later to induction of hundreds of IFN-stimulated genes (ISGs) (Sato et al. 2000).

There are multiple sources of viral dsRNA; viral genomes (dsRNA viruses), viral replication (ssRNA viruses) or viral transcription (DNA viruses). Over the last 10–15 yr, many endogenous dsRNAs have been described, including pre-miRNA, rRNA stem–loops, inverted repeat Alu-elements (IR-Alu) (Chen et al. 2008; Sugimoto et al. 2015) and mitochondrial dsRNA, which is a result of bidirectional transcription of mitochondrial DNA (mtDNA) (Borowski et al. 2013; Dhir et al. 2018). In mice, mtDNA is a ~16.5 kb long, circular DNA molecule, which is organized into mitochondrial nucleoids (Peralta et al. 2012). The mouse mitochondrial genome contains 37 genes coding for 13 proteins, 12S and 16S ribosomal RNAs (rRNAs), and 22 transfer RNAs (tRNAs) (Bibb et al. 1981; Bayona-Bafaluy et al. 2003). The mtDNA genes are arranged mostly on the Heavy (H) strand, which encompasses 12 of the 13 mRNAs, rRNAs and 14 of the 22 tRNAs, while the Light (L) strand codes only for one mRNA and eight tRNAs (Peralta et al. 2012). Among these endogenous dsRNAs, RIG-I and MDA-5 recognize mitochondrial dsRNA using different mechanisms.

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et al. 2012). Almost the entire L strand transcript undergoes rapid decay by the RNA degradosome complex, which includes PNPase and hSuv3 helicase (Borowski et al. 2013). Those two enzymes are extremely important in restricting the levels of mitochondrial dsRNA, as the loss of either of them causes massive accumulation of mitochondrial dsRNA that escapes into the cytoplasm. This mitochondrial dsRNA triggers an MDA5-driven antiviral signaling pathway that results in a type I IFN response and therefore it is another cellular source of dsRNA (Dhir et al. 2018).

Another dsRNA sensing system is the oligoadenylate synthetase (OAS)/RNase L pathway (Li et al. 2017). OAS enzymes (OAS1, OAS2, OAS3, and OASL), are IFN-inducible, and upon sensing dsRNA they produce 2′-5′-oligoadenylates (2–5A) which activates the nuclease RNase L. In mice, there are five Oas genes, Oas1a and Oas1g, Oas2, Oas3 and Oas2 that encode active enzymes and one, Oasl1 that encodes an enzymatically inactive protein (Kakuta et al. 2002; Kristiansen et al. 2011). Oasl1 was initially shown to inhibit the translation of IRF7 mRNA and to act as a negative regulator of type I IFN synthesis (Lee et al. 2013a), but new evidence shows that early in the antiviral response Oasl1 has an opposite role as it promotes RLR signaling by trapping viral RNA in stress bodies (Kang et al. 2018). Active RNase L cleaves all cellular RNAs predominantly in single-stranded regions of UpN dinucleotides (UA and UU > UG) (Silverman and Weiss 2014; Li et al. 2017). In the absence of the RNA editing enzyme, adenosine deaminase acting on RNA 1 (ADAR1), OAS can be activated by self-dsRNA, resulting in RNase L activity and cell death (Li et al. 2017). Recent studies show that RIG-I like receptors are activated by oligoadenylate synthetase-like protein 1 (OASL1). Loss of OASL1 expression reduced RIG-I signaling and enhanced virus replication in human cells. Conversely, Oasl1 expression enhanced RIG-I-mediated IFN induction (Zhu et al. 2014).

P53 controls transcription and is a well-documented tumor suppressor. It is also an ISG, induced by IFN upon viral infection (Takaoka et al. 2003). It is thought that its role in innate immunity is to induce apoptosis, thus preventing the spread of viral infection. P53 that is posttranslationally modified is located in the cytoplasm and enhances the permeability of the mitochondrial outer membrane thus stimulating apoptosis (Green and Kroemer 2009). However, treating Trp53 mutant mouse embryonic fibroblasts (MEFs) with the DNA demethylating agent 5-aza-2′-deoxycytidine (5-aza-dC) (Leonova et al. 2013), was also reported to cause a huge increase in the level of transcripts encoding short interspersed nuclear elements (SINEs) and other species of noncoding RNAs that generated a strong type I IFN response. Thus it appears that another function of p53 in cells is to ensure the silencing of repeats that can accidentally induce an immune response.

Identifying endogenous, immune-stimulating dsRNA is especially important in relation to autoimmune diseases. Here we demonstrate that in the absence of transcription factor p53, an immunogenic, endogenous dsRNA is produced in cells. Surprisingly this endogenous dsRNA does not encode predominantly SINEs or other tandem repeats. Instead we show that this RNA is of mitochondrial origin and is processed by the OAS/RNase L system. Our study therefore sheds new light on the role of endogenous RNA in diseases with aberrant immune responses.

RESULTS

Endogenous dsRNA from Trp53 cells can induce immune responses when transfected into test cells

Our initial hypothesis was that dsRNA from Adar1 mutant cells is unedited and therefore these cells contain higher amounts of immunogenic dsRNAs. This would result in the activation of dsRNA-binding receptors and to subsequent activation of immune pathways leading to type I IFN production. However, MEFs generated from Adar1 mutant embryos where the entire gene is deleted are not viable and require an additional Trp53 mutation eliminating p53 protein expression for viability (Mannion et al. 2014). To validate our hypothesis and investigate if endogenous dsRNA can induce an immune response, total RNA was isolated from Adar1;Trp53 double mutant MEFs and from control wild-type and Trp53 mutant MEFs. DsRNA was isolated from these purified total RNA samples by in vitro immunoprecipitation with J2 antibody specific for dsRNA (Fig. 1A). The quality of the RNA was evaluated by microcapillary electrophoresis (Supplemental Fig. S1). To test the innate immune inducing potential of dsRNA from the different cell types the isolated dsRNA was then transfected back into cultured cells with lipofectamine and innate immune responses were measured; all of the three different MEF cell lines were tested as recipients.

To investigate whether lack of ADAR1 editing in Adar1;Trp53 double mutant MEFs generated dsRNA that can induce an innate immune response after transfection into test cells, we first performed immunoblotting on the test cells after dsRNA transfection to measure the expression of two dsRNA-binding sensors; RIG-I and MDA5 (Fig. 1B; Supplemental Fig. S2). Unexpectedly, the immunoblots revealed that the dsRNA that induced the strongest response was the dsRNA isolated from Trp53 mutant MEFs. DsRNA isolated from Adar1;Trp53 MEFs also increased the expression of RIG-I and MDA5 receptors, but the effect was lower when compared to Trp53 mutant MEFs. This trend was observed for all the three cell lines transfected; WT MEFs, Trp53 MEFs and Adar1;Trp53 MEFs. Nonetheless, quantification relative to α-tubulin level (with ImageJ Software; data not shown), shows higher intensity for cells transfected with dsRNA from Trp53 cells. This experiment was repeated and the same results were observed. The results of
immunoblots demonstrate that the lack of Trp53 in MEF cells generates dsRNA able to activate RIG-I and MDA5 receptors and this effect is independent of the genotype of the recipient cell.

To validate that dsRNA derived from Trp53 mutant MEFs activates an innate immune response in recipient transfected cells, an ELISA to detect the secretion of proinflammatory cytokine IL-6 was performed (Fig. 1C). The highest IL-6 secretion was observed for the cells transfected with dsRNA from Trp53 MEFs, lower for cells transfected with dsRNA from Adar1;Trp53 double mutant MEFs and lowest for dsRNA from WT MEFs. Those results confirmed that this RNA activates the whole pathway, as cytokine secretion is the last step.

Trp53 mutant MEFs were transfected with dsRNA derived from WT, Trp53 or Adar1;Trp53 MEFs and the expression level of mRNA transcripts was measured for Ifit3, Ifit1, Irf7 (three ISGs), IFN-α, Ifih1 (encoding MDA5) and Ddx58 (encoding RIG-I) (Fig. 1D). The results were normalized to mRNA expression in nontransfected cells. The expression of ISG transcripts was significantly up-regulated when cells were transfected with dsRNA from Trp53 MEFs, when compared to cells transfected with dsRNA from Adar1;Trp53 MEFs or WT MEFs. In general, changes in mRNA expression were more modest for Irf7 than for Ifit1 and Ifit3. In the case of IFN-α and Ifih1 transcripts the increase followed the same trend, with the highest increase for cells transfected with dsRNA from Trp53 MEFs. Transcripts encoding the two receptors, Ifih1(MDA5) and Ddx58 (RIG-I), had lower increases in expression in cells transfected with dsRNA from Trp53 MEFs (approximately fivefold) when compared to WT MEFs and Adar1;Trp53 MEFs (approximately twofold) dsRNA. The increase is especially low for Ddx58, which is probably due to its relatively high basal expression in nontransfected cells (as seen on immunoblot, Fig. 1B). Thus, results of immunoblots, ELISA and qPCR assays demonstrate that dsRNA from Trp53 MEFs can induce an immune response when transfected back into Trp53 MEFs. Next, we sought to determine which receptor recognizes endogenous dsRNA from Trp53 mutant cells. RNAi knockdown of transcripts encoding MDA5 and RIG-
I receptors revealed that the immune response induced by this dsRNA is mediated mainly, but not exclusively, via MDA5 receptor (Fig. 2). Immunoblot results show an increase in the expression of MDA5 in cells with RIG-I knockdown. However, when MDA5 is knocked down, the increase in the expression of RIG-I receptor is minor. RIG-I receptor is therefore also involved in this immune response induced by endogenous dsRNA from Trp53 mutant cells, however to a lower extent (Fig. 2). There is variability in this experiment due to the batch difference in poly I:C, but this does not affect the results which is that after knockdown of MDA5 in Trp53 mutant cells the isolated dsRNA is less immune-stimulatory.

For isolation of dsRNA, we used the dsRNA-specific monoclonal antibody J2 that is widely used to detect viral dsRNA in animals and plants (Weber et al. 2006; Dhir et al. 2018). We verified the specificity of J2 for dsRNA in vitro by digesting immunoprecipitated dsRNA from Trp53 MEFs with RNase V1 prior to transfection into Adar1;Trp53 MEFs. RNase V1 is specific for double-stranded helical conformations in RNA (Lowman and Draper 1986; Nilsen 2013). In contrast to undigested and denatured dsRNA, dsRNA digested with RNase V1 was unable to induce immune response measured as expression of two dsRNA-binding sensors; RIG-I and MDA5 (Fig. 3A). This result confirms that the immune-stimulatory effect indeed is mediated by dsRNA. We then determined whether the response is caused by the cytoplasmic or nuclear fraction of dsRNA. For this we performed cellular fractionation with digiton (Supplemental Fig. S4) first; total RNA was then isolated from cytoplasmic and nuclear fractions separately, followed by immunoprecipitation of dsRNA with the J2 antibody. Only dsRNA from the cytoplasmic fraction of Trp53 MEFs was able to induce the immune response (Fig. 3B).

**RNA-seq analysis of dsRNA**

We performed J2 immunoprecipitation-based dsRNA sequencing (dsRNA-seq) to identify the differences between dsRNA derived from wild-type MEF cells and the two mutants cell lines. For sequencing purposes, ribosomal RNA was first depleted from the dsRNA before library preparation; ribosomal RNA-depleted dsRNA was still immune-stimulatory (Supplemental Fig. S5). Chromosome-wise coverage analysis of dsRNA-seq revealed that the mitochondrial genome has the highest ratio of differentially expressed genes in dsRNA from Trp53 mutant versus wild-type cells and from Adar1;Trp53 versus wild-type cells (Fig. 4A). The results were normalized to the total number of protein-coding genes on each chromosome. The analysis of expression levels of individual, differentially expressed genes mapped to the mitochondrial chromosome shows higher fold increase in 12 out of 13 protein-coding genes in Trp53 than in Adar1;Trp53 when compared to wild-type (Fig. 4B). We do not observe an increase in sense and antisense mitochondrial RNA that could form intermolecular duplexes that would activate MDA5. This result is consistent with previous reports showing that MDA5 is not responding to longer stretches of paired sense and antisense strands but instead it recognizes dsRNA that was created as intramolecular duplexes (Runge et al. 2014).

We also analyzed the repetitive sequence content of the J2-immunoprecipitated dsRNA material. For this analysis “RepBase repeat consensus”
for mapping reads representing various types of repetitive elements in rodents was used (Supplemental Fig. S6; Supplemental Table S3). Immunoprecipitated dsRNA obtained from Adar1;Trp53 and from Trp53 MEFs did not show any significant enrichment in repetitive elements when compared to dsRNA from WT MEFs.

**Adar1;Trp53 and Trp53 cells have an elevated immune response**

To investigate whether there is an altered global transcriptional profile of genes involved in innate immune response in the Adar1;Trp53 and Trp53 cells from which the dsRNA was being purified, we performed next generation sequencing of total RNA isolated from these cells and wild-type controls. The resulting data revealed that transcripts of 1427 genes are up-regulated and 2153 down-regulated (at least 1.5 log2 fold and adjusted $P$-value <0.05) in Trp53 MEFs, while transcripts of 922 genes are up-regulated and 1440 down-regulated (at least log 1.5 fold) in Adar1;Trp53 MEFs (Supplemental Table S2; Supplemental Fig. S7).

Transcripts of some genes were up-regulated in both cell types, including classical proinflammatory and type I IFN-dependent genes encoding proteins of the Oas family (Oas3, Oasl1), IFN-induced protein with tetra triopeptide repeats 1 (Ifit1) and 3 (Ifit3), Irf7, Ifi205, and Isg15 (Fig. 5A). Two of the genes involved in immune responses, interleukin-1 receptor antagonist (Il1rn) and interleukin-4 receptor (Il4ra) were down-regulated in both cell lines. Interestingly, immunoglobulin-like domain-containing receptor 2 (Ildr2), which is a novel negative regulator for T cells involved in autoimmune response (Hecht et al. 2018), was highly down-regulated (over 10-fold) in Trp53 cells, but slightly up-regulated in Adar1;Trp53 cells.

Sequencing results were confirmed by RT-qPCR comparing mRNA transcripts for Oas3, Ifit3 and Isg15 (Fig. 5B). Not all qPCR results correspond to sequencing results. For example, Oasl1 mRNA transcripts were higher in Adar1;Trp53 than in Trp53 MEFs (data not shown). Collectively, this finding suggests that, both Adar1;Trp53 and Trp53 MEFs when compared to wild-type cells, have altered transcriptional profiles of genes involved in innate immune responses, however transcripts of more genes are elevated in Trp53 MEFs. Additionally, Trp53 MEFs show a fivefold increase in Ifn-α mRNA transcript, compared to wild-type cells (Fig. 5B), whereas Adar1;Trp53 MEFs
show an over fivefold increase in lfn-β mRNA transcript, compared to wild-type cells (Fig. 5B). Interestingly, chromosome-wise coverage analysis of total RNA-seq of differentially expressed genes again revealed that the mitochondrial genome has the highest ratio of differentially expressed genes in RNA from Trp53 versus wild-type cells and Adar1;Trp53 versus wild-type cells (Fig. 5C). The results were normalized to the total number of protein-coding genes on each chromosome. All genes with an adjusted P-value <0.05 were considered as differentially expressed. Data are average of three independent experiments.

**Immunogenic dsRNA is a mitochondrial RNA cleaved by RNase L**

RNA-seq of total RNA revealed that one of the most up-regulated genes in both Adar1;Trp53 and Trp53 MEFs, was Oas3 (in comparison to WT MEFs) which is a member of the OAS family. Oas3 upon binding to dsRNA, activates RNase L that cleaves cellular RNA into small fragments (Kakuta et al. 2002; Ibsen et al. 2014). Among all Oas proteins, Oas3 shows a dominant role in RNase L activation, with a higher affinity for dsRNA than either Oas1 or Oas2 (Li et al. 2016). We therefore hypothesized, that dsRNA from Trp53 MEFs that induces an innate immune response in transfected recipient cells, could be a product of RNase L cleavage. We chose to test this hypothesis using dsRNA from Trp53 MEFs as the mRNA encoding RNase L is slightly up-regulated in Trp53 MEFs whereas it is not up-regulated and remains at the same level in the WT cells as in Adar1;Trp53 MEFs (Fig. 6A). RNase L was knocked down in Trp53 MEF with esiRNAs (Supplemental Fig. S8), followed by total RNA extraction and immunoprecipitation of dsRNA (Fig. 6B). The J2 antibody only recognizes dsRNA that is greater than 40 perfectly paired bases so esiRNAs, which are 21 bp long, are not immunoprecipitated with this antibody. Isolated dsRNA was then transfected into Trp53 MEFs and the immune response was measured in the transfected cells. As a control, cells were transfected with dsRNA derived from Trp53 MEFs treated with a general, nonspecific siRNA.

The expression of three chosen ISG mRNA transcripts, Ifit3, Ifit1, and Irf7, was significantly less induced when reporter cells were transfected with dsRNA derived from cells with RNase L knockdown, in comparison to cells transfected with dsRNA from Trp53 MEFs (Fig. 6C). Cells transfected with dsRNA obtained from Trp53 MEFs treated with negative control siRNAs show similar induction of ISGs to cells transfected with control dsRNA from Trp53 MEFs. This demonstrates that the ability of dsRNA to induce immune response relies on the presence of RNase L. The same reduced induction was observed for mRNA transcripts encoding IFN-α, Ifih1 and Ddx58, with significant decrease in expression in reporter cells transfected with dsRNA from cells with silenced RNase L (Fig. 6C).

To further confirm the effect of RNase L knockdown upon dsRNA immunogenicity, an ELISA assay was used to detect the secretion of proinflammatory cytokine IL-6 (Fig. 6D). The highest IL-6 secretion was observed from the cells transfected with dsRNA from Trp53 MEFs, lower
from cells transfected with dsRNA from Trp53 MEFs with negative siRNAs and the lowest for dsRNA from Trp53 MEFs with RNase L knocked down. Taken together with qPCR results, this shows that the dsRNA from Trp53 MEFs that can induce an innate immune response may in-deed be the product of RNase L cleavage, as the silencing of RNase L in those cells decreases the immunogenicity of dsRNA significantly.

Finally, to further support the idea that the dsRNA that triggers immune response is of mitochondrial origin, we inhibited mitochondrial transcription with low concentrations of ethidium bromide prior to dsRNA isolation (Fig. 7). Ethidium bromide causes decreased mitochondrial transcription without significantly affecting nuclear transcription (Hayakawa et al. 1998). In ethidium bromide–treated cells, analysis of transcripts by qPCR shows decreased mitochondrial transcription compared to untreated cells (Fig. 7B). DsRNA isolated from EtBr-treated cells was transfected into Adar1;Trp53 MEFs; immunoblot with RIG-I and MDA5 antibodies on proteins from transfected cells show that dsRNA isolated from ethidium bromide–treated cells triggers a weaker immune response (Fig. 7A). This result unambiguously demonstrates that in Trp53 MEFs there is an increase in mitochondrial dsRNA reaching the cytoplasm that can increase the hazard of triggering an innate immune response.

DISCUSSION

The complex role of p53 in the immune system is increasingly appreciated (Muñoz-Fontela et al. 2016). Cells expressing p53 show p53-dependent apoptosis in response to viral infection (Turpin et al. 2005). As expected, mice deficient in p53 are prone to infection with viruses, including influenza A virus, probably due to the lack of an apoptotic response (Yan et al. 2015). Additionally,
p53 directly activates transcription of a set of immune-responsive genes, including Tlr3, and IFN regulatory factors 5 (Irf5) and 7 (Irf7) (Mori et al. 2002; Taura et al. 2008; Yan et al. 2015). In recent years, it has become evident that p53 is associated with the development of autoimmune diseases and suppresses the aberrant expression of proinflammatory factors (Takatori et al. 2014). Defective p53 is also now linked to the development of rheumatoid arthritis, SLE and dermatomyositis/polymyositis (Kovacs et al. 1997; Chauhan et al. 2004; Mimura et al. 2007). P53 directly inhibits the production of numerous cytokines by inhibiting signal transducer and activator of transcription 1 (STAT1) (Youlyouz-Marfak et al. 2008), and p53 deficiency in macrophages increases the production of proinflammatory cytokines, such as IL-1, IL-6, IL-12, and TNF-α (Komarova et al. 2005; Gudkov et al. 2011).

Here, we show that dsRNA from Trp53 mutant murine cell lines induces an aberrant innate immune response in transfected reporter cells. This is related to the Trp53 mutation, as dsRNA isolated from wild-type MEF cells is unable to induce this immune response. Interestingly, the innate immune inducing effect of the dsRNA appears to be increased by the presence of ADAR1 in those cells, and decreased when ADAR1 is absent. The response to transfected dsRNA is mediated mainly, but not exclusively, via the MDA5 receptor; the RIG-I receptor is also involved in this immune response, however to a lesser extent. Cellular sensing of this endogenous dsRNA through MDA5 and RIG-I leads to their interaction with MAVS signaling protein, translocation of transcription factor IRF7, and finally to the type I IFN and IL-6 secretion and up-regulation of ISGs.

We are confident that the observed effect relies on endogenous dsRNA, and not on dsDNA or other nucleic acids. Immuno precipitation with an antibody specific for dsRNA was described in multiple publications (most recently in [Dhir et al. 2018]), which together with our control experiments prove the dsRNA specificity of the antibody. We can also control the dsRNA quality prior to transfection by analysis with microcapillary electrophoresis. In addition, the response is eliminated when we treat with a dsRNA-specific RNase.

The results presented here were surprising to us initially because we had expected that dsRNA not edited by ADAR1 would be more immune-stimulatory. One of the biological roles of ADAR1 is to negatively regulate the IFN response by editing endogenous dsRNA. This was observed in Adar1 deficient mice that die by embryonic day E12.5 with severe effects of massive interferon production, liver disintegration and widespread apoptosis (Hartner et al. 2004, 2009; Wang and Carmichael 2004; Mannion et al. 2014). Additionally, Adar1p150 mutant MEFs also show abnormal type I IFN response (Ward et al. 2011) and are not viable. The Adar1 null mutant we use here bears a deletion of exon 2 to 13, removing most of the open reading frame of the protein. Other Adar1 mutant alleles still contain dsRNA-binding domains and viable Adar1 MEFs can be generated from them (Wang et al. 2004). It is only after generating Adar1;Trp53 double mutant MEFs that we were able obtain viable cells. These cells were characterized by elevated immune responses, which were reduced after transfection of inosine containing dsRNA (Vitali and Scadden 2010; Mannion et al. 2014). These results demonstrated that the cell uses inosine to help discriminate between self and non-self dsRNA. If inosine is present in dsRNA it binds to RLRs and prevents activation of the innate immune response. However, if no inosine is present then the cell cannot discriminate “self” from “non-self” and treats the dsRNA as being of viral origin and activates a type I IFN response.

We found that endogenous dsRNA can indeed induce an immune response and this effect was associated to the p53 deficiency. To examine this dsRNA in more detail, we performed dsRNA-seq. The results revealed the mitochondrial chromosome as the chromosome with the highest proportion of genes differentially expressed between Trp53 and wild-type cells, and between Adar1;Trp53 and wild-type cells. The analysis of expression levels of individual, differentially expressed genes mapped to the mitochondrial chromosome, shows higher fold increase in 12 out of 13 protein-coding genes, in Trp53 compared to wild-type than in Adar;Trp53 compared to wild-type, with the only exception being the mitochondrial encoded cytochrome C oxidase III (MT-CO3). In contrast to recently published results (Borowski et al. 2013; Dhir et al. 2018), most of the detectable cellular dsRNA is not encoded by the mitochondrial genome; however, our results still indicate that the ability of endogenous dsRNA to induce immune response relies mostly on the mitochondrial fraction and not on dsRNA in general. We also find that in whole transcripomes of both Trp53 MEF and Adar1;Trp53 MEF cell lines the highest percentage of differentially expressed genes was identified in the mitochondrial genome. In addition, when mitochondrial transcription was decreased by growing the cells in low concentrations of ethidium bromide, the dsRNA isolated from these cells was less immune-stimulatory when transfected into Adar1;Trp53 MEFs. This demonstrates that the endogenous dsRNA that we isolated with dsRNA antibodies is of mitochondrial origin and is not encoding SINEs or other RNAs with tandem repeats as was found when the DNA-demethylating agent 5-aza-dC was used to treat Trp53MEFs (Leonova et al. 2013).

In human cells, mitochondrial RNA can induce immune response to a similar extent as bacterial RNA (Dhir et al. 2018). Mitochondrial dsRNA is particularly dangerous as it can lead to the activation of potent innate immune defense mechanisms that have evolved to protect vertebrates against microbial and viral attack. This may result in autoimmune disorders. Under the normal
circumstances, this RNA is strictly controlled by the degradosome components, mitochondrial RNA helicase SUV3 and polynucleotide phosphorylase (PNPase). Loss of either of those enzymes results in massive accumulation of mitochondrial dsRNA that escapes into the cytoplasm and drives type I IFN response (Dhir et al. 2018). Another recent publication has demonstrated that Protein kinase RNA-activated (PKR) also binds mitochondrial dsRNA which can regulate its phosphorylation and signaling (Kim et al. 2018).

We found that Trp53 mutant MEFs, both with and without concurrent knockout of Adar1, have elevated levels of multiple genes involved in the immune response. This was observed in cells not stimulated with dsRNA. Since Trp53 cells have fivefold higher IFN-α secretion than wild-type and Adar1;Trp53 cells, it was then crucial to identify the main transcripts differing between the Trp53 mutant and the other two lines. Among the most up-regulated genes in both Trp53 and Adar1;Trp53 cells, when compared to wild-type, were members of the Oas family, Oasl1 and Oas3. Oas family enzymes catalyzes the synthesis of oligoadenylates of the general structure ppp(A2′p)nA (2′–5′), which upon binding, activate the endoribonuclease RNase L. When activated, RNase L catalyzes the degradation of both viral and cellular RNAs (Hovanessian and Justesen 2007). Oas1-3 upon binding to dsRNA, activate RNase L, with Oas3 having the dominant role in this process (Ibsen et al. 2014; Li et al. 2016). The role of mouse Oasl1 however is still puzzling. Mouse Oasl1 gene is the orthologue of the human OASL gene and is enzymatically inactive (Kristiansen et al. 2011). Until recently, the only reported role of Oasl1 was the negative regulation of IFN response (Kristiansen et al. 2011; Lee et al. 2013b; Oh et al. 2016). However, a recent report demonstrates that Oasl1 plays context-dependent roles in the antiviral response (Kang et al. 2018). In early stages of viral infection, Oasl1 forms stress granules trapping viral RNAs and promoting efficient RLR signaling. Stress granule formation is dependent on the RNA-binding activity of Oasl1. However, in the late stages of infection, Oasl1 inhibits translation resulting in down regulation of IFN production (Kang et al. 2018). These results demonstrate that Oasl1 has a more complicated role in response to viral dsRNA, than previously appreciated.

FIGURE 6. Immunogenic dsRNA is a product of RNase L cleavage. (A) RNaseL mRNA expression in WT, Trp53, and Adar1;Trp53 MEFs. (B) Schematic representation of the experiment. Prior to isolation of endogenous dsRNA from Trp53 MEFs, RNaseL was knocked down in those cells with esiRNA. Isolated dsRNA was transfected back into Trp53 MEFs. (C) RT-qPCR analysis of Ifit3, Ifit1, Irf7, IFN-α, Ddx58, and Ifih1 mRNA in Trp53 MEFs transfected with dsRNA from Trp53, RNaseL knocked down MEFs and control dsRNA. Results are normalized to mRNA expression in nontransfected cells. (D) ELISA showing the mean levels of IL-6 in cell-culture supernatants of Trp53 MEFs transfected with dsRNA from Trp53, RNaseL knocked down MEFs and control dsRNA. Results are normalized to cytokine expression in nontransfected cells. Data are mean ± SD of three independent experiments. (*) P ≤ 0.05, (**) P ≤ 0.01, (***) P ≤ 0.001.
The level of RNaseL transcript in unstimulated cells is higher in Trp53 cells than in wild-type or Adar1;Trp53. Therefore, we studied the role of the OAS/RNase L system in the production of immunogenic dsRNA in this cell line. Nevertheless, RNase L exists in the cell in inactive form and is activated by dimerization, which occurs upon 2-5A binding. Knocking down RNase L in Trp53 MEFs results in the loss of the immune-stimulating ability of the endogenous dsRNA. This can be seen on multiple levels of the type I IFN response in transfected reporter cells, starting with the lowered expression of MDA5 and RIG-I receptors, through smaller production of IFN-α and IL-6, and finally, much lower expression of ISGs. Active RNase L cleaves all cellular RNAs predominantly in single-stranded regions at UpN dinucleotides (UA and UU > UG), however the cleaved RNA may have single- and double-stranded regions (Wreschner et al. 1981). The J2 antibody used for immunoprecipitation recognizes continuous duplex structures of at least 40 bp in length (Bonin et al. 2000).

The OAS system is localized in multiple cellular compartments. Whereas RNase L can be either cytoplasmic or mitochondrial (Le Roy et al. 2007; Kjær et al. 2014), OAS can be localized in mitochondria, cytoplasm, but additionally also in the ER and nucleus (Lin et al. 2009). The cellular localization of the OAS/RNase L system may also depend on the stage of cell life. Activation of RNase L results in the degradation of all RNA within the cell, viral and endogenous, which leads to apoptosis of mammalian cells in a caspase-dependent manner. At the beginning of apoptosis, RNase L and OAS are localized in the mitochondria and cytosol fractions, while at the onset of apoptosis both enzymes are largely in mitochondria (Domingo-Gil and Esteban 2006). Based on the obtained results it is not possible to confirm with full certainty whether isolated mitochondrial dsRNA was cleaved by mitochondrial or cytoplasmic RNase L. Both possibilities seem plausible. Nevertheless, we are confident that the immunoprecipitated RNA, and therefore RNA that can induce immune response, is mitochondrial and that it was processed by the OAS/RNase L system.

Overall, our results demonstrate a role of p53—OAS axis in mitochondrial RNA processing and preventing self-nucleic acid such as dsRNA from aberrantly activating innate immune responses. The lack of p53 transcription factor causes activation of the OAS/RNase L system in the absence of a danger signal such as viral infection.

MATERIALS AND METHODS

Cell culture

Mouse embryonic fibroblasts (MEF) were derived from mouse embryos of the same genetic background (C57BL/6J) as previously described (Mannion et al. 2014). Cells were grown in DMEM high glucose medium (BioSera) in the presence of 10% fetal bovine serum (FBS). To silence RNase L, MEF Trp53 cells were transfected with 30 nM of 21 bp esiRNA, targeting mouse RNase L (MISSION, Sigma), with Lipofectamine 3000 for 48 h. The silencing effectiveness was measured by qPCR, with primers specific for RNase L (see Supplemental Table S1). The silencing specificity of siRNA was assessed by parallel transfection with 30 nM of siRNA.
Universal Negative Control (MISSION, Sigma). To silence RIG-I and MDA5 receptors, Adar1;Trp53 MEFs were transfected with 30 nM of esiRNA targeting mouse Ifih1 or 30 nM of esiRNA targeting mouse Ddx58 (MISSION, Sigma), with Lipofectamine 3000 for 72 h.

**RNA extraction**

Total RNA was obtained from cells by phenol/chloroform extraction with an overnight isopropanol precipitation. RNA samples were treated with 1 µL DNase I (Qiagen) per 20 µg of total RNA and then precipitated in 70% isopropanol with 150 mM sodium acetate. A 2 µg aliquot of total RNA was reverse transcribed into cDNA with a reverse transcriptase with 100 µM Oligo(dT)15 primer (GeneAll). Samples were incubated for 1 h at 55°C and the reaction terminated by heating for 5 min at 70°C.

**Quantitative RT-PCR**

For all qPCR experiments, the QuantStudio 12K Flex Real-Time PCR System (Thermo Scientific) was used. PCR was performed with FastStart Universal SYBR Green Master (Rox) (Sigma) with gene-specific primers (see Supplemental Table S1). Results were normalized to the mRNA expression of β-actin or Gapdh. To analyze relative quantification of genes, the comparative CT Method (ΔΔCT) was used. Samples were analyzed in technical duplicates and biological triplicates.

**Immunoprecipitation of dsRNA**

Forty micrograms of total RNA was isolated as described above and incubated with the J2 anti-dsRNA IgG2a monoclonal antibody (Scicons) in the presence of IP buffer (50 mM Tris-HCl pH 8, 150 mM NaCl, 1% Triton X-100, and 1 mM EDTA) at 4°C for 16 h. Next, 70 µL of Protein A-Sepharose 4B Fast Flow beads (Sigma) was added to the RNA-J2 mix and incubated for another 4 h at 4°C. The beads coupled with J2-RNA complexes were then washed gently three times at 4°C with IP buffer. DsRNA was isolated from the beads with the standard phenol–chloroform RNA extraction protocol. The concentration and quality of the isolated RNA was measured by TapeStation (Agilent 2200) according to the manufacturer’s instructions.

**Transfection with dsRNA**

In the experiments, three cell lines, WT MEF, Trp53 MEF, and Adar1;Trp53 MEF, were transfected with dsRNA (also derived from WT MEF, Trp53 MEF, and Adar1;Trp53 MEF). Always, the same amount of immunoprecipitated RNA was transfected into cells with Lipofectamine 3000. This was 2 µg transfected into ~1.6 × 10^5 of MEFs in 2.3 mL of media.

**ELISA**

Cell culture media was stored at −80°C and thawed at room temperature prior to ELISA. The amount of IL-6 secreted by MEF cells (confluent 4 cm^2 in 1 mL of medium) was measured with Mouse IL-6 ELISA Set (BD OptEIA) according to the manufacturer instructions. Samples were analyzed in technical duplicates and biological triplicates.

**Immunoblotting**

Cells (1–1.2 × 10^6) were lysed in 50 mM Tris-HCl pH 8, 150 mM NaCl, 1% Triton X-100, and 1 mM EDTA with the addition of protease inhibitors (cOmplete, Roche). Protein samples were separated at 100 V by 10% SDS-PAGE electrophoresis and blotted on a nitrocellulose membrane. Membranes were incubated with specific antibodies: mouse anti-α-tubulin (1:8000, Sigma-Aldrich), rabbit anti-MDA5 (1:800, Cell Signaling), rabbit anti-RIGI (1:800, Cell Signaling), rabbit anti-fibrillarin (1:10,000, Abcam). HRP-conjugated anti-rabbit (1:80,000) and anti-mouse (1:5000) antibodies were purchased from Sigma-Aldrich. Proteins were revealed with Pierce ECL Western Blotting Substrate (Thermo Fisher Scientific). All immunoblots were performed in triplicate and the quantification of the protein bands from all three experiments is shown in each figure.

**Subcellular fractionation**

Fractionation protocol was adapted from Holden and Horton (2009) and Liu and Fagotto (2011). Trp53 MEFs cells were grown on 15 cm dishes until they reached ~50% confluency. Cells on the dish were washed with cold 1× PBS and permeabilized with 6 mL of digitonin solution (1× NEH Buffer containing 45 µg/mL digitonin, 10 mM DTT, and 10 mM MgCl2) by gentle rocking at 4°C for 10 min. (10× NEH Buffer: 1500 mM NaCl, 2 mM EDTA, 200 mM HEPES-NaOH pH 7.4). The released cytoplasmic fraction was collected and cleared by centrifugation at 500g, 4°C for 10 min. Cytoplasmic RNA was isolated from clear supernatant with TruPure Reagent (Sigma-Aldrich) as per manufacturer’s instructions. Cell residues on the dish were washed with cold 1× PBS and collected with 3 mL of Buffer 2 (150 mM NaCl, 50 mM HEPES-NaOH pH 7.4, 1% NP40) by scraping, followed by 30 min incubation on ice. Nuclei were pelleted by centrifugation at 7000g, 4°C for 15 min. The supernatant containing other organelles was discarded and the pelleted nuclei were washed with cold 1× PBS. The nuclear pellet from one 15 cm dish was disrupted by incubating for 30 min at 37°C with 10 U of TURBO DNase (Thermo Fisher) in 10× Reaction Buffer with occasional pipetting using 100 µL tips. Afterward, nuclear RNA was isolated with TruPure Reagent as per manufacturer’s instructions. A small part of both cytoplasmic and nuclear fraction was saved for immuno-blotting to verify the fractions’ purity.

**Ethisiode bromide treatment of MEFs to block mitochondrial transcription**

Trp53 MEFs were seeded in 10 cm plates and grown until they reached ~80% confluency. Then, ethidium bromide (ApplChem) was added to the cell culture medium at a final concentration of 0.05 µg/mL (Hayakawa et al. 1998; Surovtseva et al. 2011). Cells were grown in ethidium bromide-supplemented medium for 24 h. Cells were washed 3× with PBS, then total RNA was
isolated and treated with DNase as described above. Expression of mitochondrial genes was analyzed by quantitative RT-PCR as described above with RNA from untreated Trp53 MEFs as a control. Random hexamers were used for reverse transcription instead of oligo(dT)15. Cycling conditions for qPCR: one cycle of 95°C for 10 min, then 40 cycles of 95°C for 10 sec, 55°C for 20 sec, and 72°C for 8 sec. Total RNA from ethidium bromide–treated cells was also used for dsRNA immunoprecipitation using the same protocol as described above.

**NGS library preparation**

Total RNA was isolated from cells using the phenol–chloroform extraction method and treated with DNase I (Ambion). RNA was depleted of rRNA with the Ribocop rRNA Depletion Kit (Lexogen) and used for library preparation using the SENSE Total RNA-seq Library Prep Kit (Lexogen). For the sequencing of the dsRNA, immunoprecipitation was performed with J2 (dsRNA-specific antibody), then 1 µg per sample was depleted of the dsRNA, immunoprecipitation was performed with J2 (dsRNA-specific antibody), then 1 µg per sample was depleted of rRNA with Ribocop rRNA Depletion Kit (Lexogen). The rRNA depleted dsRNA was used for library preparation with SENSE Total RNA-seq Library Prep Kit (Lexogen). The libraries were prepared with TruSeq Illumina adapters. Sequencing was performed on the NextSeq 500/550 sequencer.

**Differential gene expression**

The raw data generated during sequencing were quality checked using FastQC and preprocessed with Trimmomatic. Adapter sequences and low-quality ends were trimmed (Phred score <3; both 5’ and 3’ ends). Alignment was performed by STAR to the mouse reference genome–GRCm38 primary assembly; Ensembl release 86 (NGS of total RNA) and release 91 (NGS of dsRNA). Raw gene counts were counted only from uniquely mapped reads by featureCounts (NGS of total RNA). Estimated raw gene counts were counted from both uniquely and multimapped reads by RSEM (NGS of total RNA). Strandedness of the sequencing was considered during the counting. Differential gene expression analysis was performed by DESeq2 Bioconductor package and the raw *P*-values were adjusted for multiple testing error using Benjamini–Hochberg method. Samples were analyzed in biological triplicates.

**Analysis of dsRNA sequences immunoprecipitated with J2 antibody**

Raw alignment coverage was calculated using DeepTools and normalized to counts-per-million (CPM). Signals of each of the strands were calculated separately. All positions that did not have coverage of at least 0.5 CPM were filtered out. Only signals with a continuous length of at least 150 bp (S9 A) or 500 bp (S9 B) were kept. The regions were selected only if they were present in two out of three replicates for each condition (WT, Trp53 and Adar1;Trp53). Visualization of the strands was done in ggplot2 R package.

**SUPPLEMENTAL MATERIAL**

Supplemental material is available for this article.

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