m6A modifications regulate intestinal immunity and rotavirus infection

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31

32 **Abstract**

33 N6-methyladenosine (m6A) is an abundant mRNA modification and affects many
34 biological processes. However, how m6A levels are regulated during
35 physiological or pathological processes such as virus infections, and the *in vivo*
36 function of m6A in the intestinal immune defense against virus infections are
37 largely unknown. Here, we uncover a novel antiviral function of m6A
38 modification during rotavirus (RV) infection in small bowel intestinal epithelial
39 cells (IECs). We found that rotavirus infection induced global m6A modifications
40 on mRNA transcripts by down-regulating the m6A eraser ALKBH5. Mice lacking
41 the m⁶A writer enzymes METTL3 in IECs (*Mettl3* Δ IEC) were resistant to RV
42 infection and showed increased expression of interferons (IFNs) and
43 IFN-stimulated genes (ISGs). Using RNA-sequencing and m6A RNA
44 immuno-precipitation (RIP)-sequencing, we identified IRF7, a master regulator
45 of IFN responses, as one of the primary m6A targets during virus infection. In the
46 absence of METTL3, IECs showed increased *Irf7* mRNA stability and enhanced
47 type I and III IFN expression. Deficiency in IRF7 attenuated the elevated
48 expression of IFNs and ISGs and restored susceptibility to RV infection in
49 *Mettl3* Δ IEC mice. Moreover, the global m6A modification on mRNA transcripts
50 declined with age in mice, with a significant drop from 2 weeks to 3 weeks post
51 birth, which likely has broad implications for the development of intestinal
52 immune system against enteric viruses early in life. Collectively, we
53 demonstrated a novel host m6A-IRF7-IFN antiviral signaling cascade that
54 restricts rotavirus infection *in vivo*.

55

56 **Keywords**

57 Rotavirus, N6-methyladenosine, METTL3, ALKBH5, IFNs

58

59 **Introduction**

N6-methyladenosine (m6A) is the most abundant internal mRNA modification and modulates diverse cellular functions through m6A-related writers, erasers, and readers[1-3]. The m6A modification directly recruits m6A-specific proteins of the YT521-B homology (YTH) domain family[1]. These proteins mediate the m6A-dependent regulation of pre-mRNA processing, microRNA processing, translation initiation, and mRNA decay[1]. In recent works, m6A modifications has been identified in the genomes of RNA viruses and the transcripts of DNA viruses with either a pro-viral or anti-viral role[4-8]. Furthermore, m6A RNA modification-mediated down-regulation of the α -ketoglutarate dehydrogenase (KGDH)-itaconate pathway inhibits viral replication independent of the innate immune response[9]. According to Gao et al. (2020), m6A modification preserves the self-recognition of endogenous transcripts. Deletion of the m6A writer *Mettl3* decreases the m6A modifications in endogenous retrovirus (ERV) transcripts. The accumulation of ERVs activates pattern recognition receptors (e.g. RIG-I) pathways, resulting in a detrimental interferon response in livers of fetal mice[10].

The m6A modification of the enterovirus 71 (EV71) RNA genome is important for viral propagation, and EV71 infection increases the expression of m6A writers *in vitro*[11]. m6A methyltransferase *Mettl3* knockdown reduces whereas m6A demethylase FTO knockdown increases EV71 replication[11]. In addition, human cytomegalovirus can up-regulate the expression of m6A-related proteins[12, 13]. Despite the knowledges about m6A regulation and function during viral infection revealed by these *in vitro* studies, the regulation of m6A modifications and the specific role of m6A in the anti-viral response *in vivo*, especially in the gastrointestinal tract, remains unclear.

Rotavirus (RV), a member of the family *Reoviridae*, is a nonenveloped icosahedral-structured virus with 11 segments of double-stranded RNA. Children under the age of five are at high risk of rotavirus infection, which causes severe diarrhea, dehydration, and death[14]. Rotaviruses encode multiple viral proteins to inhibit innate immune responses by degrading interferon regulatory factors

(IRFs) and mitochondrial antiviral-signaling protein (MAVS), thus facilitating efficient virus infection and replication[15, 16]. The timely induction of an IFN response is key to the host successful control of invading viruses, including RV[17-19]. Here, we found rotavirus infection induced global m6A modifications on mRNA transcripts by down-regulating the m6a eraser ALKBH5. Mice lacking the m⁶A writer enzymes METTL3 in IECs (*Mettl3*ΔIEC) were resistant to RV infection. We identified IRF7, a master regulator of IFN responses[17], as one of the primary m6A targets during virus infection. In the absence of METTL3, IECs showed increased *Irf7* mRNA stability and enhanced type I and III IFN expression. Deficiency in IRF7 attenuated the elevated expression of IFNs and ISGs and restored susceptibility to RV infection in *Mettl3*ΔIEC mice. Collectively, we identified a novel regulation and function of m6A modifications in an enteric viral infection model *in vivo*.

Results

The regulation and function of mRNA m6A modifications during rotavirus infection

RV infections primarily take place in children under the age of five in humans and in neonatal mice younger than 2 weeks-old[14, 20]. Intriguingly, total RNA m6A modifications in the mouse ileal tissues, revealed by a m6A dot blot and mass spectrum (MS) analysis, significantly declined from 2 weeks to 3 weeks post birth (Figure 1a, 1b and 1c), which caused by increased *Alkbh5* expression (Figure 1d, Figure 1-figure supplement 1a and Figure 1-figure supplement 1b) and development of microbiota might be involved (Figure 1-figure supplement 2). Indeed, overexpression of *Alkbh5* in mouse IEC cell line caused decreased m6A level (Figure 1-figure supplement 1c), supporting the role of *Alkbh5* in regulating global m6A levels in intestine. These data indicating a potential link between the resistance to RV infection and the decline of m6A modification during early life development.

Besides, the global m6A RNA modification levels increased in the ileum tissue of suckling mice post RV murine strain EW infection (Figure 1e, 1f and Figure 1-figure supplement 3). As a control, the m6A modification can barely be detected in IECs deficiency with m6A writer *Mettl3* (*Mettl3^{fl/fl} Vil1^{Cre}*, *Mettl3 Δ IEC*) (Figure 1e and 1f). Thus, we hypothesize that RV may induce an enriched cellular m6A modification environment and a weakened innate immune response to facilitate virus replication. To investigate the *in vivo* role of m6A in the anti-RV immunity, we infected *Mettl3 Δ IEC* mice and wild-type (WT) littermates with RV EW strain. The viral RNA load in *Mettl3 Δ IEC* mice ileum tissue was significantly lower than that in the WT mice (Figure 1g and Figure 1-figure supplement 4). Fecal virus shedding was also significantly lower in *Mettl3 Δ IEC* mice (Figure 1h). Genetic knockdown of *METTL3* in HT-29 cells, a human colonic epithelial cell line, by CRISPR-mediated gene silencing, also led to reduced RV replication (Figure 1i), further highlighting the resistance phenotype to RV infection by *METTL3* deficiency.

***Mettl3* deficiency in IECs results in decreased m6A deposition on *Irf7*, and increased interferon responses**

To dissect the underlying mechanism, we performed RNA-sequencing using the IECs from *Mettl3*-deficient mice and littermate controls at steady state. Most of the differentially expressed genes in *Mettl3*-deficient IECs vs WT IECs were enriched in the pathways of “defense response to virus”, “response to interferon-beta”, and “positive regulation of innate immune response” by gene ontology analysis (Figure 2a). Heatmap also showed that a panel of interferon stimulate genes (ISGs) are upregulated in *Mettl3*-deficient IECs compared to WT IECs (Figure 2b). To map potential m6A modification sites on mRNAs of these differentially expressed genes in IECs, we conducted m6A RIP-sequencing based on a previously reported protocol [5]. Metagene plots showed the m6A peak is enriched near the stop codon and 3'-UTR region, which is consistent with previously studies[21, 22](Figure 2-figure supplement 1b). We found that m6A

modified one of the master regulators of IFNs, *Irf7* (Figure 2c), which played a key role in the network of these differential expressed genes in *Mettl3*-deficient IECs compared to WT IECs, analyzed by STRING (Figure 2d). Of note, *Irf7* was the only IRFs that highly expressed in *Mettl3*-deficient IECs, and *Irf7* was the prominently highest expressed *Irf* in IECs (Figure 2e), indicating *Irf7* might be one of the key genes that are modulated by m6A modifications. We also validated our results by m6A RIP-qPCR to examine m6A modification sites in *Irf7* mRNA based on our RIP-sequencing data in *Mettl3* knock-down MODE-K cells and predicted results from the database (<http://rna.sysy.edu.cn>) (Figure 2-figure supplement 1a, Figure 2-figure supplement 1c and Figure 2-figure supplement 1d). It should be noted that our m6A-RIP-seq were performed at steady state. We did not identified previous reported *Ifns*[12, 13], possibly due to the low expression level of *Ifns* in IECs at steady state.

IRF7 is a known master regulator of Type I interferon and Type III interferon-dependent immune responses in respond to virus infection[16, 17, 23] . We reasoned that loss of m6A modification on *Irf7* mRNA is responsible for the increased IFN response and subsequent resistance to RV infection. Thus, we first validated the regulation of *Irf7* mRNA levels by m6A in mice and in cells. We found an increase of *Irf7* mRNA in ileum tissue of *Mettl3* Δ IEC mice compared to that in WT littermate control mice (Figure 2f). Consistently, the expression of *IRF7* mRNA was also higher in *METTL3* knockdown HT-29 and *METTL3* knockout rhesus monkey MA104 cells, suggesting that the regulation of *IRF7/Irf7* expression by m6A is likely conserved across species (Figure 2-figure supplement 2, Figure 2-figure supplement 3). Furthermore, genetic knockdown of *METTL3/Mettl3* in human or mice IEC cell lines also led to increased *IRF7/Irf7* expression and *IFN/Ifn* responses (Figure 2-figure supplement 1f, Figure 2-figure supplement 2). As m6A is known to regulate the mRNA decay, we next sought to determine whether the stability of *IRF7* mRNA is regulated by m6A. We used actinomycin D to block the *de novo* RNA synthesis in HT-29 cells to assess the

RNA degradation by *METTL3* knockdown. The *IRF7* mRNA degraded significantly slower in *METTL3*-knockdown HT-29 cells than the control cell line (Figure 2g, Figure 2-figure supplement 2b).

To directly evaluate the role of m6A in modulating the stability of *Irf7* mRNA, luciferase reporter assays were conducted. In comparison with wild-type *Irf7*-3'UTR (*Irf7*-WT) constructs, the ectopically expressed constructs harboring m6A mutant *Irf7*-3'UTR (*Irf7*-MUT) showed significantly increased luciferase activity (Figure 2h, Figure 2-figure supplement 1e). Further, mutation of m6A modification sites can directly increase the expression of *Irf7* and enhance the antiviral function of *Irf7* in MEF cells (Figure 2-figure supplement 4a and Figure 2-figure supplement 4b). These results suggest that the upregulation of *Irf7* mRNA level in *Mettl3* Δ IEC mice is caused by the loss of m6A modification mediated mRNA decay. To evaluate the potential influence of m6A on *Irf7* transcriptional targets, we also measured the expressions of IFNs and ISGs in rotavirus infected ileum tissue from *Mettl3* Δ IEC mice and littermate WT mice. We found the transcriptional targets of *Irf7*, were all up-regulated in *Mettl3* Δ IEC mice (Figure 2i). Furthermore, we found that the mRNAs of *Irf7* and its transcriptional targets ISGs increased in the ileum of the mice from 1 to 4 weeks, with a dramatic up-regulation from 2 to 3 weeks (Figure 2-figure supplement 5), which was concomitant with the decrease of global m6A modifications (Figure 1a, 1b and Figure 1-figure supplement 1a). These results demonstrated that *METTL3* deficiency in IECs results in decreased m6A deposition on *IRF7*, and increased interferon response.

***IRF7* Deficiency attenuated the increased interferon response and resistance to rotavirus infection in *Mettl3* Δ IEC mice**

To determine whether *Irf7* plays a key role in the resistance phenotype to RV infection in *Mettl3* deficient mice in IECs, we crossed *Irf7*^{-/-} mice to *Mettl3* Δ IEC mice. Following RV oral gavage, the expression of IFNs and ISGs in ileum from

Irf7^{-/-} *Mettl3*ΔIEC mice were significantly lower than those from *Mettl3*ΔIEC mice at 2dpi (Fig 3a-c), and unlike the increased expression of IFNs and ISGs in *Mettl3*ΔIEC mice vs littermate WT controls, by deficiency of IRF7, the expression of IFNs and ISGs in ileum from *Irf7*^{-/-} *Mettl3*ΔIEC mice were not significantly different from those from *Irf7*^{-/-} mice (Fig 3a-c), suggesting that *Irf7* mediates the increased expression of IFNs and ISGs in *Mettl3*ΔIEC mice.

Moreover, the *Irf7*^{-/-} *Mettl3*ΔIEC mice showed significantly higher viral loads in ileum tissue and higher fecal shedding of RV *Mettl3*ΔIEC (Figure 3d-f). Similarly, unlike the much lower fecal viral shedding in *Mettl3*ΔIEC mice vs littermate WT controls, by deficiency of IRF7, the fecal viral shedding in *Irf7*^{-/-} *Mettl3*ΔIEC mice were not significantly different from those in *Irf7*^{-/-} mice (Fig 3d), suggesting that *Irf7* mediates the resistant phenotype of rotavirus infection measured by fecal viral shedding in *Mettl3*ΔIEC mice. Notably, the viral proteins expression difference in ileum from *Irf7*^{-/-} *Mettl3*ΔIEC mice vs that from *Irf7*^{-/-} mice (9.7-fold lower for NSP2 and 9.3-fold lower for VP7), was much lower than the viral proteins expression difference in ileum from *Mettl3*ΔIEC mice vs that from littermate mice (267.1-fold lower for NSP2 and 283.4-fold lower for VP7) (Fig 3e), suggesting that besides the contribution of *Irf7* to the resistant phenotype of rotavirus infection in IECs from *Mettl3*ΔIEC mice, other pathways (e.g. m6A modifications in RV RNA) may also play roles. Therefore, IRF7 is an important mediator of the increased IFNs and ISGs expression in *Mettl3*ΔIEC mice and genetic deletion of *Irf7* restored the resistant phenotype of *Mettl3*ΔIEC mice to RV infection.

Rotavirus suppresses ALKBH5 expression through NSP1 to evade immune defense

We next sought to determine how RV regulates the m6A modifications in IECs. We first measured whether RV infection regulates the m6A-related writer and eraser proteins in the intestine. The protein levels of the methyltransferases

METTL3 and METTL14 and demethylase FTO were not affected by RV infection in ileum tissue (Figure 4a, 4b). In contrast, the protein level of demethylase ALKBH5 was significantly down-regulated by RV infection in the ileum (Figure 4a, 4b). To determine whether ALKBH5 play a role in anti-RV infection since it's suppressed during RV infection, we generated the IEC-specific deletion of ALKBH5 in mice (*Alkbh5^{fl/fl} Vil1^{Cre}, Alkbh5ΔIEC*). The depletion of ALKBH5 in IECs did not affected the anti-RV immune response (Figure 4c), the viral shedding in the feces (Figure 4d), or the viral protein expression in the ileum (Figure 4e), likely due to the suppressed ALKBH5 expression in ileum tissue of WT mice infected by RV.

We next asked whether there is redundancy between m6A eraser ALKBH5 and FTO. We first checked the expression of *Alkbh5/ALKBH5* and *Fto/FTO* in our RNA-seq data in mouse IECs, and also a previous reported RNA-seq data in human intestinal enteroid [24]. We found the *ALKBH5* expression is much higher than *FTO* in intestine (Figure 4-figure supplement 1a). We did qPCR analysis in mouse epithelial cell line MODE-K and showed the similar results (Figure 4-figure supplement 1a). Furthermore, we knock down METTL3, ALKBH5 and FTO in MODE-K cell. Through dot blot assay, we found ALKBH5 but not FTO is the dominant m6A eraser in IECs (Figure 4-figure supplement 1). Furthermore, overexpressing of recombinant ALKBH5 increased the IFN/ISGs response and inhibit RRV infection in mouse IEC MODE-K cells (Figure 4-figure supplement 2), suggesting re-expression of ALKBH5 might overcome the immune evasion of RV.

Non-structural protein 1 (NSP1) is a well-established RV-encoded innate immune antagonist that are showed to degrade IRF3 and β -Trcp [25, 26]. To test the potential role of NSP1 in ALKBH5 inhibition, we used the recently developed reverse genetics system and rescued recombinant WT RV SA11 strain and a mutant virus that does not express NSP1 (NSP1-null)[27]. We infected HEK293

cells with WT and NSP1-null RVs, only WT RV reduced ALKBH5 protein levels (Figure 4f), suggesting that the down-regulation of ALKBH5 expression by RV is NSP1-dependent. These results suggest that RV might evade the anti-viral immune response via downregulation of ALKBH5 expression by NSP1.

Discussion

Previous studies reported that m6A modifications on mRNA in mice embryonic fibroblasts or normal human dermal fibroblasts negatively regulate the IFN response by accelerating the mRNA degradation of type I IFNs [12, 13]. However, these studies were mainly conducted *in vitro*, leaving the relationship between m6A and IFN pathway *in vivo* an unexplored territory. Since type I and type III IFNs play a critical role in the antiviral immune response in the GI tract[19], we used an RV infection model, in which IECs are specifically infected, as well as conditional knockout mice with IECs-specific depletion of m6A writer METTL3 or m6A eraser ALKBH5, to study the role of m6A modification in regulating IFN response towards rotavirus in IECs. Using RNA-seq and m6A-seq techniques, we identified IRF7, a key transcription factor upstream of IFNs and ISGs, as one of the m6A modified targets during RV infection, and IRF7 is essential to mediate the elevated anti-rotavirus immune response by METTL3 deficiency. These results identify *Irf7* as an important m6A target, and characterize, for the first time, the regulation of IFN response during RV infection in intestine, enabling a better understanding of how m6A modifications on mRNAs regulate anti-viral innate immune responses.

In addition to directly regulating target genes that involved in innate immune pathways, m6A modification is also known to affect viral gene expression, replication and generation of progeny virions[1, 28]. Some viral RNA genomes are modified by m6A, such as simian virus 40[4, 29], influenza A virus[30], adenovirus[31], avian sarcoma virus[32], Rous Sarcoma virus[33], hepatitis C virus[34], and Zika virus[35]. Through our m6A-RIP-qPCR, Rotavirus RNA was

also found to have m6A modification (Figure 4-figure supplement 3). Of note, genetic deletion of METTL3 in monkey kidney MA104 cell line, which has limited IFN responses[36], also led to reduced RV replication(Figure 2-figure supplement 3). Further, IRF7 deficiency didn't fully restore the suppressed rotaviral infection in *Mettl3* Δ IEC mice (Fig 3e). These results suggest that besides the contribution of *Irf7* to the resistant phenotype of rotavirus infection in IECs from *Mettl3* Δ IEC mice, other pathways (e.g. m6A modifications in RV RNA) may also play roles. The detailed mechanisms warrant further investigations in the future.

Intriguingly, m6A modifications were shown to maintain the genomic stability of mice embryonic stem cells by promoting the degradation of endogenous retrovirus (ERV) mRNA, or by regulating ERV heterochromatin and inhibiting its transcription[2, 3]. The absence of m6A results in the formation of abnormal endogenous dsRNA, which causes an aberrant immune response and necrosis in the hematopoietic system[10]. In the intestine, through immunostaining with J2 antibody, we detected the increase of dsRNA levels in IECs of *Mettl3* Δ IEC mice compare to the littermate WT mice (Figure 4-figure supplement 4). In consistency, the aforementioned RNA-seq data showed that the expression of a set of ISGs including IRF7 was significantly upregulated in *Mettl3* KO IECs. Our data suggest a dual activation model in steady state that in the absence of METTL3, the increase of dsRNA will induce the IFN responses, and the increased stability of *Irf7* mRNA, which serves as a key transcription factor of IFN and ISG expression, will amplify this process. Furthermore, m6A modification of viral RNA genomes affect the activation of innate sensor-mediated signaling[37, 38]. Decreased m6A modification on RV genomes may activate innate sensors directly and induce higher IFN response, which will also be amplified by increase of mRNA stability of *Irf7*.

Although m6A is involved in many important biological processes, the regulation of m6A modifications remain poorly understood. Here, we found RV infection down-regulates the level of m6A eraser ALKBH5 to induce m6A modifications, in an NSP1 dependent manner. The precise mechanism remains to be examined. As a result, ALKBH5 deficiency in IECs results in normal susceptibility to RV infection. In addition, the global m6A modification on mRNA transcripts declines by ages in the intestine, with a significant drop from 2 to 3 weeks post birth, which implicate the drop of RV infectivity in adult mice vs neonatal mice. The dual regulation of m6A levels during RV infection and development provide new insights into the choice of either RV or the host on regulation of m6A modification to achieve either immune evasion or immune surveillance, respectively.

In conclusion, our work shed light into a novel role of m6A modifications in RV infection *in vivo*, and reported a tissue specific regulation of m6A during RV infection or development (Figure 4g). Future studies on tissue-specific regulation of m6A modification by viral infections in other tissues and organs (e.g. lung, liver) will be of interest.

Materials and Methods

Mice

METTL3 conditional knockout mice were generated by inserting two loxp sites into the intron after the first exon and the intron before the last exon of *Mettl3* using CRISPR/cas9 based genome-editing system as previously described[5]. ALKBH5 conditional knockout mice were generated by inserting two loxp sites into the introns flanking the first exon of *Alkbh5* using CRISPR/cas9 based genome-editing system as previously described[39]. Genotyping of *Mettl3*^{fl/fl} mice, *Alkbh5*^{fl/fl} mice, *Vil1*^{Cre} mice (The Jackson Laboratory, Stock No: 021504), and *Irf7*^{-/-} mice (RIKEN BRC, RBRC01420) were confirmed by PCR using primers as list below:

360 *Mettl3*^{fl/fl} mice

361 *Mettl3*-L1+ : CCCAACAGAGAAACGGTGAG

362 *Mettl3*-L2- : GGGTTCAACTGTCCAGCATC

363 *Vil1*^{Cre} mice

364 *Vil1*^{Cre}-182/150-F : GCCTTCTCCTCTAGGCTCGT

365 *Vil1*^{Cre}-182-R : TATAGGGCAGAGCTGGAGGA

366 *Vil1*^{Cre}-150-R : AGGCAAATTTTGGTGTACGG

367 *Irf7*^{-/-} mice

368 RBRC01420-Irf7-WT-F : GTGGTACCCAGTCCTGCCCTCTTTATAATCT

369 RBRC01420-Irf7-Mut-F : TCGTGCTTTACGGTATCGCCGCTCCCGATTTC

370 RBRC01420-Irf7-R : AGTAGATCCAAGCTCCCGGCTAAGTTCGTAC

371 *Alkbh5*^{fl/fl} mice

372 *Alkbh5*-L1+: GCACAGTGGAGCACATCATG

373 *Alkbh5*-L2-: CAGAGGGCAAGCAACCACAC

374 The sex-, age- and background-matched littermates of the knockout or
375 conditional knockout mice were used as the controls in the present study. All
376 mice were on the C57BL/6 background. Mice were maintained in SPF conditions
377 under a strict 12 h light cycle (lights on at 08:00 and off at 20:00). All animal
378 studies were performed according to approved protocols by the Ethics
379 Committee at the University of Science and Technology of China
380 (USTCACUC202101016).

381

382 **Cell culture**

383 The MA104 cell line was obtained from the Cell Resource Center, Peking Union
384 Medical College (which is the headquarter of National Infrastructure of Cell Line
385 Resource). HEK293T (ATCC CRL-3216), HT-29 (ATCC HTB-38D™) were obtained
386 from the American Type Culture Collection (ATCC). MODE-K cell were obtained
387 from Shanghai HonSun Biological Technology Co., Ltd.

388 The identity of these cell lines were authenticated with STR profiling

389 (FBI, CODIS). All of these cells were cultured in Dulbecco's modified Eagle's

medium (DMEM) (Hyclone) supplemented with 10% fetal bovine serum (FBS) (Clark); All cells were cultured at 37°C in 5% CO₂. All cells were tested for eliminating the possibility of mycoplasma contamination.

Poly(I:C) transfection

Poly(I:C) (OKA, A55994) was transfected at a concentration of 2 µg/ml using Lipofectamine 3000 (Invitrogen) according to the manufacturer's protocol. Cells were analyzed at indicated hours.

Plasmids and SgRNAs

pSIN-m*Alkbh5*-3xflag and pLVX-m*Alkbh5*-puro was constructed by cloning mouse *Alkbh5* CDS into pSIN-3xflag and pLVX-puro plasmid.

pLVX-m*Irf7*-wt-puro was constructed by cloning mouse *Irf7* CDS and 3'UTR into pLVX-puro.

pLVX-m*Irf7*-Mut-puro was constructed by cloning mutated mouse *Irf7* CDS and 3'UTR into pLVX-puro. (The potential m6A modification sites (1400A, 2213A, 2693A, 2927A, 3446A, 3475A, 3491A, 3590A and 3620A of mice *Irf7*(NC_000073)) were predicted on SRAMP website. The A to G mutation in these sites did not change the amino acids)

All gene silencing was done using a CRISPR-cas9 system, with lentiCRISPR v2 plasmid (Addgene no. 52961). The following sgRNAs were cloned downstream of the U6 promoter:

Human, rhesus and mouse METTL3: 5' -GGACACGTGGAGCTCTATCC-3';

Mouse ALKBH5: CCTCGTAGTCGCTGCGCTCG;

Mouse Fto: GAAGCGGTCCAGACCGCGG;

Lentiviruses were generated by co-transfection of lentiCRISPR v2 constructs and packaging plasmids (psPAX2, Addgene no. 12260 and pMD2.G, Addgene no. 12259), using PEI DNA transfection reagent (Shanghai maokang biotechnology), into HEK293T cells, according to the manufacturer's instructions. At 48h post transfection, supernatants were collected and filtered through a 0.22µm

polyvinylidene fluoride filter (Millex). To induce gene silencing, cells were transduced with lentivirus expressing sgRNA and were puromycin selected (2µg/ml) for 4–5 days. The depletion of target proteins was confirmed by immunoblot analysis.

Virus infections

Rhesus and simian RV strains, including RRV (Rhesus), SA11-4F (simian), SA11-NSP1null (simian) were propagated in MA104 cells as previously described[26]. Viruses were activated by trypsin (5 µg/ml) at 37°C for 30 min prior to infection. Cells were washed with PBS three times and incubated with RV at different MOIs at 37°C for 1 hr. After removal of RV inoculum, cells were washed with PBS, cultured in serum-free medium (SFM) and harvested for qPCR and western blot analysis.

EW stock virus was prepared by infecting 5-days-old C57BL/6J mice, and harvesting crude centrifugation-claried intestinal homogenate as previously described[15].

For all rotavirus infection except indicated elsewhere, 8-day-old wild-type mice, or genetically deficient mice were orally inoculated by gavage with RV EW virus as previously described[15]. Mice were sacrificed, stool and small intestinal tissue were collected at indicated time points post infection. Viral loads in intestinal tissues and feces were detected by RT-qPCR.

Quantitative analysis of m6A level

The m6A analysis by LC-MS/MS was performed by Metware Biotechnology Co., Ltd (Wuhan, China). In brief, 1 µg purified RNA was digested into single nucleosides with 1 U nuclease P1 (Takara) and 1 U rSAP (Takara) and incubated at 37 °C. The digested RNA was injected into a LC-MS/MS which includes the ultra-performance liquid chromatography with a C18 column. The positive ion multiple reaction-monitoring (MRM) mode was adopted to detect m6A

abundance. m6A levels were calculated from the standard curve which was generated from pure nucleoside standards.

RT-qPCR

For cells and tissues, total RNA was extracted with TRNzol Universal reagent (Tiangen) in accordance with the manufacturer's instructions. Real-time PCR was performed using SYBR® Premix Ex Taq™ II (Tli RNaseH Plus) (Takara) and complementary DNA was synthesized with a PrimeScript™ RT reagent Kit with gDNA Eraser (Takara). The target genes were normalized to the housekeeping gene (*Gapdh*, *HPRT*) shown as $2^{-\Delta Ct}$. The used primers are as follows:

Primers detect mouse genes:

Mettl3-F: ATTGAGAGACTGTCCCCTGG
Mettl3-R: AGCTTTGTAAGGAAGTGCGT
Mettl14-F: AGACGCCTTCATCTCTTTGG
Mettl14-R: AGCCTCTCGATTT CCTCTGT
Fto-F: CTGAGGAAGGAGTGGCATG
Fto-R: TCTCCACCTAAGACTTGTGC
Alkbh5-F: ACAAGATTAGATGCACCGCG
Alkbh5-R: TGTCCATTTCCAGGATCCGG
Wtap-F: GTTATGGCACGGGATGAGTT
Wtap-R: ATCTCCTGCTCTTTGGTTGC
Gapdh-F: TGAGGCCGGTGCTGAGTATGTCG
Gapdh-R: CCACAGTCTTCTGGGTGGCAGTG
Hprt-F: ACCTCTCGAAGTGTTGGATACAGG
Hprt-R: CTTGCGCTCATCTTAGGCTTTG
Irf7-F: ATGCACAGATCTTCAAGGCCTGGGC
Irf7-R: GTGCTGTGGAGTGCACAGCGGAAGT
Isg15-F: GGTGTCCGTGACTAACTCCAT
Isg15-R: TGGAAAGGGTAAGACCGTCCT

479 Oas1a-F: GCCTGATCCCAGAATCTATGC
480 Oas1a-R: GAGCAACTCTAGGGCGTACTG
481 Ifnb-F: TCCGAGCAGAGATCTTCAGGAA
482 Ifnb-R: TGCAACCACCACTCATTCTGAG
483 Ifnl3-F: AGCTGCAGGCCTTCAAAAAG
484 Ifnl3-R: TGGGAGTGAATGTGGCTCAG
485 Ifit1-F: GAACCCATTGGGGATGCACAACCT
486 Ifit1-R: CTTGTCCAGGTAGATCTGGGCTTCT

487

488 **Primers detect human genes:**

489 hGAPDH-F: ATGACATCAAGAAGGTGGTG
490 hGAPDH-R: CATACCAGGAAATGAGCTTG
491 hIRF7-F: CGAGACGAAACTTCCCGTCC
492 hIRF7-R: GCTGATCTCTCCAAGGAGCC
493 hIFNL3-F: TAAGAGGGCCAAAGATGCCTT
494 hIFNL3-R: CTGGTCCAAGACATCCCCC
495 hCXCL10-F: TGGCATTCAAGGAGTACCTC
496 hCXCL10-R: TTGTAGCAATGATCTCAACACG
497 hIFIT1-F: CAACCATGAGTACAAATGGTG
498 hIFIT1-R: CTCACATTTGCTTGGTTGTC

499

500 **Primers detect rhesus genes:**

501 Rhesus-GAPDH-F: ATGACATCAAGAAGGTGGTG
502 Rhesus-GAPDH-R: CATACCAGGAAATGAGCTTG
503 Rhesus-IFIT1-F: CAACCATGAGTACAAATGGTG
504 Rhesus-IFIT1-R: CTCACACTTGCTTGGTTGTC
505 Rhesus-IRF7-F: GTTCGGAGAGTGGCTCCTTG
506 Rhesus-IRF7-R: TCACCTCCTCTGCTGCTAGG
507 Rhesus-IFNL1-F: ACTCATACGGGACCTGACAT
508 Rhesus-IFNL1-R: GGATTCGGGGTGGGTTGAC

509 Rhesus-IFNb-F: GAGGAAATTAAGCAGCCGCAG
510 Rhesus-IFNb-R: ATTAGCAAGGAAGTTCTCCACA

511

512 **Primers detect virus genes:**

513 Rotavirus EW-NSP2-F: GAGAATGTTCAAGACGTACTCCA

514 Rotavirus EW-NSP2-R: CTGTCATGGTGGTTTCAATTTC

515 Rotavirus EW-VP4-F: TGGCAAAGTCAATGGCAACG

516 Rotavirus EW-VP4-R: CCGAGACACTGAGGAAGCTG

517 Rotavirus EW-VP7-F: TCAACCGGAGACATTTCTGA

518 Rotavirus EW-VP7-R: TTGCGATAACGTGTCTTTCC

519 RRV VP7-F: ACGGCAACATTTGAAGAAGTC

520 RRV VP7-R: TGCAAGTAGCAGTTGTAACATC

521 RRV NSP2-F: GAGAATCATCAGGACGTGCTT

522 RRV NSP2-R: CGGTGGCAGTTGTTTCAAT

523 RRV NSP5-F: CTGCTTCAAACGACCCACTCAC

524 RRV NSP5-R: TGAATCCATAGACACGCC

525

526 **m6A dot blot assay**

527 Total RNA was isolated from mice ileum using TRNzol Universal Reagent
528 (Tiangen, Lot#U8825) according to the manufacturer's instructions. RNA
529 samples were quantified using UV spectrophotometry and denatured at 95 °C for
530 3 min. The m6A-dot-blot was performed according to a published work[40]. In
531 brief, the primary rabbit anti-m6A antibody (1:5000, Synaptic System, #202003)
532 or the primary rabbit anti-m6A antibody (1:1000, Sigma-Aldrich,
533 ABE572-I-100UG) was applied to the Amersham Hybond-N+ membrane (GE
534 Healthcare, USA) containing RNA samples. Dot blots were visualized by the
535 imaging system after incubation with secondary antibody HRP-conjugated Goat
536 anti-rabbit IgG (Beyotime, A0208).

537

538 **Western blot**

Briefly, cells and tissue were lysed with RIPA buffer (Beyotime Biotechnology) supplemented with PMSF (Beyotime Biotechnology) and protease inhibitor cocktail (Roche). Mettl3 (abcam, ab195352, 1:2000), Mettl14 (sigma, HPA038002, 1:2000), ALKBH5 (sigma, HPA007196, 1:2000), FTO (abcam, ab92821), NSP1 and VP6 (gift from Harry B. Greenberg lab), Gapdh (proteintech), Tubulin (proteintech), beta-actin (proteintech), Phospho-IRF-7 (Ser437/438) (D6M2I) (CST), Phospho-TBK1/NAK (Ser172) (D52C2) (CST), TBK1/NAK (D1B4) (CST) and IRF-7 (D8V1J) (CST) antibodies were used in accordance with the manufacturer's instructions. After incubation with the primary antibody overnight, the blotted PVDF membranes (Immobilon, IPVH00010) were incubated with goat anti-rabbit IgG-HRP (Beyotime, A0208) or goat anti-mouse IgG-HRP (Beyotime, A0216) and exposed with BIO-RAD ChemiDoc™ Imaging System for a proper exposure period.

RNA degradation assay

The stability of targeted mRNA was assessed as previously described[5]. In brief, *Mettl3* knock down HT-29 and control cell were plated on 24-well plate. Actinomycin-D (MCE, HY17559) was added to a final concentration of 5μM, and cells were harvested by indicated time points after actinomycin-D treatment. The RNA samples are processed and qPCR was used to measure the mRNA transcripts, all data were normalized to that of t=0 time point.

Dual-luciferase assay

pmirGLO (Firefly luciferase, hRluc) vector of the Dual-luciferase Reporter assay system (Promega, E1910) was used to determine the function of m6A modification within the 3'UTR of *Irf7* transcripts. The potential m6A modification sites (3446A, 3475A, 3491A, 3590A and 3620A of mice *Irf7*(NC_000073) were predicted on SRAMP website. The mutants harbor the A to G mutations at these sites. The assay was performed according to the manufacture's instruction of Promega Dual-Luciferase Reporter Assay

System(E1910): Briefly, 300 ng of pmirGLO vector containing Irf7-3'UTR or m6A-mutant Irf7-3'UTR were transfected into HEK293T cells in triplicate wells. The relative luciferase activity was accessed 36 h post-transfection.

Isolation of IECs in the intestine

Small intestines were excised and flushed thoroughly three times with PBS. They were turned inside out and cut into ~1 cm sections then transferred into RPMI with 2 mM EDTA, and shaken for 20 min at 37 °C. Supernatants were collected through a 100-mm cell strainer to get single-cell suspensions. Cells were collected as the IEC fraction which contains both epithelial cells (~90%) and lymphocytes (IEL, ~10%). Single-cell suspension was used for further analysis.

RNA-Seq

IECs from *Mettl3* Δ IEC mice as well as the wild-type littermate control mice were isolated as described in previous section. Total RNAs were extracted with TRNzol universal RNA Reagen kits. Berrygenomics (Beijing, China) processed the total RNA and constructed the mRNA libraries, and subject them to standard illumine sequencing on Novaseq 6000 system, and obtained > 40 million Pair-end 150 reads for each sample. Raw RNA-sequencing reads were aligned to the mouse genome (mm10, GRCm38) with STAR (v2.5.3a). Gene expression levels and differential analysis was performed with edgeR(v3.29.2). Genes were considered significantly differentially expressed if showing ≥ 1.5 -fold change and FDR < 0.05. Gene set analysis was performed and enriched pathways were obtained through online bioinformatics tools (metascape) and GSEA (v4.0.3). Pathway plot were gene-rated with R package 'ggplot2'[5].

m6A RNA-IP-qPCR & m6A RNA-IP-Seq

m6A RNA-IP-Seq was carried out according to a previously published protocol[5]. In brief, total cellular RNA extracted from WT C57 mice IEC was fragmented by

ZnCl₂ followed by ethanol precipitation. Fragmented RNA was incubated with an anti-m⁶A antibody (Sigma Aldrich ABE572-I) or IgG IP Grade Rabbit polyclonal antibody (abcam, lot: 934197). The eluted RNA and input were subjected to high-throughput sequencing using standard protocols (Illumina, San Diego, CA, USA) or processed as described in 'RT-qPCR' section, except that the data were normalized to the input samples. The m⁶A RIP-Seq data were analyzed as described previously[5].

RIP-ptpn4-F: CCTCCCATCCCGGTCTCCACC

RIP-ptpn4-R: GGCTGCCCATCTTCAGGGGT

RIP-RPS14-F: ACCTGGAGCCCAGTCAGCCC

RIP-RPS14-R: CACAGACGGCGACCACGACG

RIP-Tlr3-F: TGCTCAGGAGGGTGGCCCTT

RIP-Tlr3-R: CGGGGTTTGCGCGTTTCCAG

m⁶A-IRF7-F1: GACAGCAGCAGTCTCGGCTT

m⁶A-IRF7-R1: ACCCAGGTCCATGAGGAAGT

m⁶A-IRF7-F2: GGCAAGACTTGTCAGCAGGG

m⁶A-IRF7-R2: TAGACAAGCACAAGCCGAGAC

m⁶A sites on RV-EW RNA were predicted on <http://www.cuilab.cn/sramp> website, and m⁶A-RIP-qPCR primer were designed on NCBI primer blast according to the predicted m⁶A sites.

m⁶A-RIP-qPCR primer:

RIP-EW-VP1-F: ACGAAATGCTTGTTGCTATGAGT

RIP-EW-VP1-R: AACCTGTCCGTCAACCATTC

RIP-EW-VP2-F: GGCCAGAACAGGCTAAACAAC

RIP-EW-VP2-R: CGCAGTTCTCTTTGCGCCATTT

RIP-EW-VP3-F: CGATGACAGCACAAAAGTCGG

RIP-EW-VP3-R: CGTGTCTCTTGCGAAGTC

RIP-EW-VP4-F: TCAGCAGACGGTTGAGACTG

RIP-EW-VP4-R: GGCTGAGATGTCATCGAAGTT

RIP-EW-NSP1-F: CCTCACATCTCTGCTACATGAACT

628 RIP-EW-NSP1-R: TGCTGGTTGGACATGGAATGA
629 RIP-EW-VP6-F: CTGCACTTTTCCCAAATGCTCA
630 RIP-EW-VP6-R: GAGTCAATTCTAAGTGTCTAGTCCG
631 RIP-EW-NSP3-F: CTTGACGTGGAGCAGCAAC
632 RIP-EW-NSP3-R: AATGTTTCAATGTCGTCCAACG
633 RIP-EW-NSP2-F: TCCACCACTCTAAAGAACTACTGC
634 RIP-EW-NSP2-R: TCCGCTGTCATGGTGGTTTC
635 RIP-EW-VP7-F: TCGGAACCTGCAGACTTGAT
636 RIP-EW-VP7-R: GCTTCGTCTGTTTGCTGGTA
637 RIP-EW-NSP4-F: TGCACTGACTGTTCTATTTACGA
638 RIP-EW-NSP4-R: GGGAAGTTCGCATTGCTAGT
639 RIP-EW-NSP5/6-F: GGACACCGCAAGGTCAAAAA
640 RIP-EW-NSP5/6-R: TCGTCTGAGTCTGATTCTGCTT

641

642 **J2 Immunofluorescent staining**

643 IECs from *Mettl3* Δ IEC mice as well as from the wild-type littermate control mice
644 were isolated as described in previous section. Isolated IEC were centrifuged
645 onto glass slides and fixed with 4% Paraformaldehyde for 30 min at room
646 temperature. Subsequently, permeabilized and blocked with PBS containing
647 0.1% Triton-X-100 and 5% bovine serum albumin for 1 h at room temperature.
648 Double-stranded RNA (dsRNA) was labeled by a mouse monoclonal antibody J2
649 (Scicons) for 2 h at room temperature, followed by incubation with anti-mouse
650 IgG Alexa Fluor 594-conjugated antibody (Invitrogen) for 1 h, and cells nuclei
651 were visualized with 4,6-diamidino-2-phenylindole (DAPI, Invitrogen). All
652 fluorescence images were analyzed via confocal imaging using Zeiss LSM880.

653

654 **Statistical analysis**

655 Statistical analysis was performed with the GraphPad Prism 8.0 (GraphPad, Inc.,
656 USA). Experiments were independently repeated for indicated times listed in the
657 figure legend. Representative data was exhibited as the means $\pm$ SEM.

Quantitative data was compared using Student's t test. In addition, correlational analysis of gene expression was conducted with linear regression. P-values for every result were labeled on figures, and $P < 0.05$ was reckoned as statistically significant (* $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$, **** $P < 0.0001$, NS., not significant).

Data availability statement

The data that support the findings of this study were stored in Dryad, Dataset, <https://doi.org/10.5061/dryad.p2ngf1vr8>. RNA sequencing data are available from the SRA database with accession numbers PRJNA713535.

Author Contribution:

A.W. designed, performed and interpreted most of the experiments. W.T. designed and analyzed the RNA sequencing data and m6A-Seq data. J.T. did the m6A-Seq experiment. J.W., J.G., G.Z., X.R., R.L., and D.W. helped with animal experiments and cellular experiments, S.D., R.A.F., K.Z., and H.B.L. provided critical comments and suggestions; A.W. and S.Z. wrote the manuscript. S.D. and W.T. edited the manuscript. S.Z. and W.P. supervised the project.

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Figure Legends

Figure 1. Rotavirus infection increases global m6A modifications, and *Mettl3* deficiency in IECs results in increased resistance to rotavirus infection.

(a) m6A dot blot analysis of total RNA in ileum tissues from mice with different ages. Methylene blue (MB) staining was the loading control.

(b) Quantitative analysis of (a) (mean $\pm$ SEM), statistical significance was determined by Student's t-test (**P < 0.005, ***P < 0.001, NS., not significant). The quantitative m6A signals were normalized to quantitative MB staining signals.

(c) mass spectrum (MS) analysis of m6A level in ileum tissue from mice with different ages. (mean $\pm$ SEM), Statistical significance was determined by Student's t-test (*P < 0.05, NS., not significant).

(d) qPCR analysis of indicated genes in ileum tissues from mice with different ages (mean $\pm$ SEM). Statistical significance was determined by Student's t-tests between groups (*P < 0.05, ***P<0.001, NS., not significant).

(e) WT and *Mettl3* Δ IEC mice were infected by rotavirus EW strain at 8 days post birth. m6A dot blot analysis of total RNA in ileum IEC at 2 dpi. Methylene blue (MB) staining was the loading control.

(f) Quantitative analysis of (e) (mean $\pm$ SEM). Statistical significance was determined by Student's t-test (*P < 0.05, ***P<0.001, NS., not significant). The quantitative m6A signals were normalized to quantitative MB staining signals.

(g-h) *Mettl3* Δ IEC mice and littermate controls were infected by rotavirus EW strain at 8 days post birth. qPCR analysis of RV viral loads in ileum tissue (g) or fecal samples (h) from *Mettl3* Δ IEC mice and littermate controls at 4 days post infection (dpi) (littermate WT n=4, *Mettl3* Δ IEC n=4, mean $\pm$ SEM). Statistical significance was determined by Student's t-tests between genotypes (*P < 0.05).

(i) qPCR analysis of indicated genes in Rhesus rotavirus (RRV)-infected HT-29

cells transduced with *Mettl3* sgRNA or control sgRNA, at indicated hours post infection (hpi) (mean $\pm$ SEM), statistical significance was determined by Student's t-test (*P < 0.05, ***P < 0.001, NS., not significant).

Experiments in (a, d-f, and i) are repeated twice, (g and h) are repeated four times.

Figure 2. *Mettl3* deficiency in IECs results in decreased m6A deposition on *Irf7*, and increased interferon responses.

(a) Gene ontology (GO) analysis of differentially expressed genes in IECs from *Mettl3* Δ IEC mice vs IECs from littermate WT mice.

(b) Heat map of a subset of up-regulated ISGs in IECs from *Mettl3* Δ IEC mice vs IECs from littermate WT mice, as revealed by RNA-seq (normalized data).

(c) m6A-RIP-seq analysis of *Irf7* and *Gapdh* mRNA in the ileum of WT mice.

(d) Gene regulation network of a subset of up-regulated genes including IRF7.

(e) Heat map of Interferon regulatory factors (*Irfs*) in IECs from *Mettl3* Δ IEC mice vs IECs from littermate WT mice, as revealed by RNA-seq (RPKM).

(f) *Mettl3* Δ IEC mice and littermate controls were infected by EW at 8 days post birth. qPCR analysis of the *Irf7* expression in ileum and jejunum from *Mettl3* Δ IEC mice and littermate control at 2 dpi (littermate WT n=4, *Mettl3* Δ IEC mice n=3, mean $\pm$ SEM). Statistical significance was determined by Student's t-test (*P < 0.05, **P < 0.005).

(g) q-PCR analysis of *Irf7* mRNA in *Mettl3* knockdown HT-29 cells or control cells in indicated time points post actinomycin D treatment (n=3, mean $\pm$ SEM). Statistical significance was determined by Student's t-test (*P < 0.05, ***P < 0.001, NS., not significant).

(h) Relative luciferase activity of sgEV and sg*Mettl3* HEK293T cells transfected

with pmirGLO-*Irf7*-3'UTR (*Irf7*-WT) or pmirGLO-*Irf7*-3'UTR containing mutated m6A modification sites (*Irf7*-MUT). The firefly luciferase activity was normalized to Renilla luciferase activity (n=3, mean $\pm$ SEM). Statistical significance was determined by Student's t-tests between groups (*P < 0.05, NS., not significant).

(i) *Mettl3* Δ IEC mice and littermate control and were infected by EW at 8 days post birth. qPCR analysis of selected *IFNs* and *ISGs* in ileum tissue at 2 dpi (littermate WT, n=4, *Mettl3* Δ IEC mice, n=3, mean $\pm$ SEM). Statistical significance was determined by Student's t-tests between genotypes (*P < 0.05, **P < 0.005).

Experiments in (f, h and i) are repeated three times, (g) are repeated twice.

Figure 3. IRF7 Deficiency attenuated the increased interferon response and resistance to rotavirus infection in *Mettl3* Δ IEC mice

(a-c) WT control mice, *Mettl3* Δ IEC mice, *Irf7*^{-/-} mice and *Mettl3* Δ IEC *Irf7*^{-/-} mice are all littermates. They were infected by RV EW at 8 days post birth. qPCR analysis of selected *IFNs* (a), *ISGs* (b), or *Irf7* (c) expression in ileum from indicated groups of mice at 2 dpi (littermate WT n=7, *Mettl3* Δ IEC n=5, *Irf7*^{-/-} n=3, *Mettl3* Δ IEC *Irf7*^{-/-} n=6, mean $\pm$ SEM). Statistical significance was determined by Student's t-tests between genotypes (*P < 0.05, **P < 0.005, ***P < 0.001, ****P < 0.0001, NS., not significant).

(d) qPCR analysis of fecal rotaviral shedding in indicated groups of mice at 4 dpi (littermate WT n=5, *Mettl3* Δ IEC n=5, *Irf7*^{-/-} n=3, *Mettl3* Δ IEC *Irf7*^{-/-} n=4, mean $\pm$ SEM). Statistical significance was determined by Student's t-tests between genotypes (*P < 0.05, **P < 0.005, NS., not significant).

(e-f) qPCR analysis of RV proteins expression (e) or *Mettl3* and *Irf7* (f) in ileum from indicated groups of mice at 4 dpi (littermate WT n=7, *Mettl3* Δ IEC n=5, *Irf7*^{-/-} n=3, *Mettl3* Δ IEC *Irf7*^{-/-} n=6, mean $\pm$ SEM). Statistical significance was determined by Student's t-tests between genotypes and (*P < 0.05, **P < 0.005, ***P < 0.001,

NS., not significant).

Experiments in (a-f) are repeated twice.

Figure 4. Rotavirus suppresses ALKBH5 expression through NSP1 to evade immune defense

(a) WT mice were infected by RV EW at 8 days post birth. Immunoblotting with antibodies target ALKBH5, FTO, METTL14 and METTL3 in ileum tissue from mice infected with RV EW at 2 dpi or treated with PBS.

(b) Quantitative analysis of (a) (mean $\pm$ SEM), Statistical significance was determined by Student's t-test (*P < 0.05).

(c-e) *Alkbh5* Δ IEC mice and littermate controls were infected by RV EW at 8 days post birth. qPCR analysis of indicated genes expression in ileum (c), viral shedding in feces (d), and viral proteins expression in ileum (e), from *Alkbh5* Δ IEC mice or littermate controls at 4 days post infection (littermate WT n=6, *Alkbh5* Δ IEC n=5, mean $\pm$ SEM). Statistical significance was determined by Student's t-tests between genotypes (*P < 0.05, NS., not significant).

(f) Immunoblotting with antibodies target ALKBH5, NSP1, VP6 and GAPDH in HEK293 cells infected by SA11-4F and SA11-NSP1null (MOI=1) for 24h.

(g) Graphical abstract illustrating the functions and molecular mechanisms of m6A modifications on *Irf7* in anti-RV infection.

Experiments in (a-e) are repeated three time, (f) are repeated twice.

Figure 1-figure supplement 1.

ALKBH5 regulates total RNA m6A modification level in intestine.

(a) Immunoblotting with antibodies target ALKBH5 and TUBULIN in ileum

tissues from mice with different ages.

(b) Quantitative analysis of (a)(mean $\pm$ SEM), Statistical significance was determined by Student's t-test (*P < 0.05, NS., not significant).

(c) Immunoblotting with antibodies target ALKBH5 and TUBULIN in MODE-K cells transfected with pSIN-EV or pSIN-m*Alkbh5*-3xFlag for 24h. m6A dot blot analysis of total RNA in indicated samples. Methylene blue (MB) staining was the loading control.

Experiments in (a and b) are repeated twice, (c) are repeated three times.

Figure 1-figure supplement 2.

Dot blot analysis of total RNA m6A modification level in the ileum of germ free mice.

(a) m6A dot blot analysis of total RNA in ileum tissues from germ free mice with different ages. Methylene blue (MB) staining was the loading control.

(b) Quantitative analysis of (a) (mean $\pm$ SEM), statistical significance was determined by Student's t-test (NS., not significant). The quantitative m6A signals were normalized to quantitative MB staining signals.

Figure 1-figure supplement 3.

Mass spectrum (MS) analysis of total RNA m6A modification level in RV infected ileum.

(a) WT and *Mettl3* Δ IEC mice were infected by rotavirus EW strain at 8 days post birth. MS analysis of m6A level in ileum tissue from mice at 2 dpi (mean $\pm$ SEM). Statistical significance was determined by Student's t-test (**P < 0.005).

Figure 1-figure supplement 4. Schematic design of RV infection model.

Figure 2-figure supplement 1.

Characterization of m6A modifications on *Irf7* mRNA.

(a) Putative m6A sites of mouse or human *Irf7*/*IRF7* on the genomes.

(b) Metagene plots of m6A modified sites.

(c) m6A-RIP-qPCR confirms *Irf7* as an m6A-modified gene in IECs. RNA of sgEV and sg*Mettl3* MODE-K cells was incubated with an anti-m6A antibody (Sigma Aldrich, ABE572-I). The eluted RNA and input were processed as described in 'RT-qPCR' section, the data were normalized to the input samples (n=3, mean $\pm$ SEM, Statistical significance was determined by Student's t-test (*P < 0.05, **P < 0.005, NS., not significant). *Tlr3* and *Rps14* were measured with m6A sites specific qPCR primers as positive control and negative control, *Irf7* was measured with predicted m6A sites specific qPCR primer.

(d) Knock down efficiency of METTL3 in MODE-K cells.

(e) Knock down efficiency of METTL3 in 293t cells used for luciferase assay.

(f) Western blot analysis of sgEV and sg*Mettl3* MODE-K cells transfected with 2 μ g/ml poly I:C at indicated hours, at least three replicate experiments were performed.

Experiments in (c-f) are repeated twice.

Figure 2-figure supplement 2.

METTL3 knockdown in HT-29 cells results in increased IFN response.

(a) qPCR analysis of IFN/ISGs in Rhesus rotavirus-infected METTL3 WT and KD HT-29 cells at indicated hours post infection (hpi) (mean $\pm$ SEM), statistical

significance was determined by Student's t-test (*P < 0.05, **P < 0.005, ***P < 0.001, NS., not significant).

(b) Knock down efficiency of METTL3 in HT-29 cells.

Experiments in **(a and b)** are repeated three times.

Figure 2-figure supplement 3.

METTL3 deficiency in MA104 cells results in increased resistance to Rhesus rotavirus infection.

(a) qPCR analysis of viral RNAs, IFNs, and ISGs in Rhesus rotavirus-infected *METTL3* WT and KO MA104 cells at 24 hours post infection (hpi) (WT control n=4, *METTL3* ko n=3, mean ± SEM), statistical significance was determined by Student's t-test (**P < 0.005, ***P < 0.001, NS., not significant).

(b) Knock out efficiency of METTL3 in MA104 cells.

Experiments in **(a and b)** are repeated three times.

Figure 2-figure supplement 4.

m6A modification on *Irf7* mRNA regulates its expression.

(a) WT and *Irf7*^{-/-} MEF cell were transduced with lentivirus expressing pLVX-EV-puro, pLVX-*Irf7*-WT-puro or pLVX-*Irf7*-Mut-puro containing mutated m6A modification sites. qPCR analysis of genes in different cells at indicated hours post infection (hpi) (mean ± SEM). Statistical significance was determined by Student's t-tests between groups (*P < 0.05, **P < 0.005, ***P < 0.001, NS., not significant).

(b) Western blot analysis of IRF7 expression level in different MEF cells.

Experiments in **(a and b)** are repeated three times.

Figure 2-figure supplement 5.

Expression of *Irf7* and *ISGs* in ileum from mice during early life development.

(a) qPCR analysis of indicated genes in the ileum (mean $\pm$ SEM). Statistical significance was determined by Student's t-tests between ages (*P < 0.05, **P < 0.005, NS., not significant).

Experiments in (a) are repeated twice.

Figure 4-figure supplement 1.

A non-redundant role of *Alkbh5* in regulating m6A modification in IECs.

(a) Relative expression level of *ALKBH5/Alkbh5* and *FTO/Fto* in human intestinal enteroid, mice IEC and mice IEC cell MODE-K.

(b) m6A dot blot analysis of total RNA in different MODE-K cells. Methylene blue (MB) staining was the loading control.

Experiments in (b) are repeated twice.

Figure 4-figure supplement 2.

ALKBH5 over-expression in MODE-K cells results in enhanced IFN response and increased resistance to Rhesus rotavirus infection.

(a) MODE-K cells were transduced with Lentivirus (expressing pLVX-EV-puro or pLVX-m*Alkbh5*-puro). qPCR analysis of virus genes and *Ifn/ISGs* in Rhesus rotavirus-infected EV and m*Alkbh5* over-expressing cells at indicated hours post infection (hpi) (mean $\pm$ SEM), statistical significance was determined by Student's t-test (*P < 0.05, **P < 0.005, NS., not significant).

(b) Expression level of ALKBH5 in EV and m*Alkbh5* over-expressed MODE-K cells at indicated hours post infection.

Experiments in (a and b) are repeated twice.

Figure 4-figure supplement 3.

m6A-RIP-qPCR analysis of the predicted m6A sites on Rotavirus RNA.

(a) Total RNA was extracted from IECs from 1-week-old WT mice infected by RV for 2 days. Fragmented RNA was incubated with an anti-m6A antibody (Sigma Aldrich ABE572) and IgG IP Grade Rabbit polyclonal antibody (abcam, lot: 934197). The eluted RNA and input were processed as described in 'RT-qPCR' section, the data were normalized to the input samples (n=2, mean $\pm$ SEM), *Rps14* was chosen as a negative control.

Figure 4-figure supplement 4.

***Mettl3* deficiency leads to aberrant dsRNA formation in isolated IECs.**

(a) IECs from 6-weeks-old *Mettl3* Δ IEC mice as well as the WT littermate controls were isolated. Double-stranded RNA (dsRNA) was labeled by immunostaining with a mouse monoclonal antibody J2 (Scicons), cells nuclei were visualized with 4,6-diamidino-2-phenylindole (DAPI, Invitrogen). All fluorescence images were analyzed via confocal imaging using Zeiss LSM880.

Experiments in (a) are repeated twice.

Titles of Source data file

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Figure 1. Rotavirus infection increases global m6A modifications, and *Mettl3* deficiency in intestinal epithelial cells results in increased resistance to rotavirus infection.

Figure 2. *Mettl3* deficiency in intestinal epithelial cells results in decreased m6A deposition on *Irf7*, and increased interferon responses.

Figure 3. IRF7 Deficiency attenuated the increased interferon response and resistance to rotavirus infection in *Mett13* Δ IEC mice.

Figure 4. Rotavirus suppresses ALKBH5 expression through NSP1 to evade immune defense.

Figure 1-figure supplement 1. ALKBH5 regulates total RNA m6A modification level in intestine.

(a) Immunoblotting with antibodies target ALKBH5 and TUBULIN in ileum tissues from mice with different ages.

(b) Quantitative analysis of (a) (mean $\pm$ SEM), Statistical significance was determined by Student's t-test (*P < 0.05, NS., not significant).

(c) Immunoblotting with antibodies target ALKBH5 and TUBULIN in MODE-K cells transfected with pSIN-EV or pSIN-mAlkbh5-3xFlag for 24h. m6A dot blot analysis of total RNA in indicated samples. Methylene blue (MB) staining was the loading control. Experiments in (a and b) are repeated twice, (c) are repeated three times.

a **b**

Figure 1-figure supplement 2. Dot blot analysis of total RNA m6A modification level in the ileum of germ free mice.

(a) m6A dot blot analysis of total RNA in ileum tissues from germ free mice with different ages. Methylene blue (MB) staining was the loading control.

(b) Quantitative analysis of (a) (mean $\pm$ SEM), statistical significance was determined by Student's t-test (NS., not significant).

The quantitative m6A signals were normalized to quantitative MB staining signals.

a

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(a) WT and *Mettl3* Δ IEC mice were infected by rotavirus EW strain at 8 days post birth. MS analysis of m6A level in ileum tissue from mice at 2 dpi (mean $\pm$ SEM). Statistical significance was determined by Student's t-test (**P < 0.005).

a

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Experiments in (c-f) are repeated twice.

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(b) Knock down efficiency of *METTL3* in HT-29 cells.

Experiments in (a and b) are repeated three times.

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(a) qPCR analysis of viral RNAs, IFNs, and ISGs in Rhesus rotavirus-infected *Mettl3* WT and KO MA104 cells at 24 hours post infection (hpi) (WT control n=4, *Mettl3* ko n=3, mean $\pm$ SEM), statistical significance was determined by Student's t-test (**P < 0.005, ***P < 0.001, NS., not significant).

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Experiments in **(a)** and **(b)** are repeated three times.

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Experiments in (b) are repeated twice.

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(b) Expression level of ALKBH5 in EV and m*Alkbh5* over-expressed MODE-K cells at indicated hours post infection.

Experiments in **(a)** and **(b)** are repeated twice.

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a

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