Circadian regulation of SARS-CoV-2 infection in lung epithelial cells

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**ABSTRACT**

The COVID-19 pandemic, caused by SARS-CoV-2 coronavirus, is a global health issue with unprecedented challenges for public health. SARS-CoV-2 primarily infects cells of the respiratory tract, via binding human angiotensin-converting enzyme (ACE2)\(^1,2\), and infection can result in pneumonia and acute respiratory distress syndrome. Circadian rhythms coordinate an organism’s response to its environment and recent studies report a role for the circadian clock to regulate host susceptibility to virus infection\(^3\). Influenza A infection of arhythmic mice, lacking the circadian component BMAL1, results in higher viral replication\(^4\) and elevated inflammatory responses leading to more severe bronchitis\(^5,6\), highlighting the impact of circadian pathways in respiratory function. We demonstrate circadian regulation of ACE2 in lung epithelial cells and show that silencing BMAL1 or treatment with the synthetic REV-ERB agonist SR9009 reduces ACE2 expression and inhibits SARS-CoV-2 entry and RNA replication. Treating infected cells with SR9009 limits viral replication and secretion of infectious particles, showing that post-entry steps in the viral life cycle are influenced by the circadian system. Our study suggests new approaches to understand and improve therapeutic targeting of COVID-19.

Circadian signalling exists in nearly every cell and is primarily controlled by a series of transcription/translation feedback loops. The transcriptional activators BMAL1 and CLOCK regulate thousands of transcripts including their own repressors, REV-ERβ\(\alpha\) and REV-ERβ\(\beta\), that provide a negative feedback loop to control gene expression (Fig.1a). To investigate a role for circadian pathways in the SARS-CoV-2 life cycle we initially assessed whether infection was time-dependent in synchronised Calu-3 lung epithelial cells (Supplementary Fig.1). Infected Calu-3 cells at different circadian times (CT) with SARS-CoV-2 (Fig.1b) showed a rhythmic pattern of viral replication (Fig.1c). To assess whether this was mediated at the level of virus entry we used lentiviral pseudoparticles (pp) expressing the SARS-CoV-2 Spike glycoprotein and showed time-dependent infection (Fig.1d). We showed that SARS-CoV-2pp infection of Calu-3 cells was dependent on Spike-ACE2 interaction by blocking virus uptake with a receptor mimic (ACE2-Fc) or by neutralizing infection with anti-Spike mAb FI-3A (Supplementary Fig.2). Pseudoparticles bearing the vesicular stomatitis virus (VSV) glycoprotein infected the cells with comparable efficiency at all time points (Fig.1d). These data uncover a role for circadian-signalling pathways to regulate Calu-3 epithelial cell susceptibility to SARS-CoV-2 uptake.

Since ACE2 and transmembrane protease serine 2 (TMPRSS2) co-regulate SARS-CoV-2 internalization\(^1,2\) we measured their expression in synchronised Calu-3 cells. ACE2 protein levels varied over 24h, with the trough (CT6) and peak (CT18) of expression associating with SARS-CoV-2pp entry (Fig.2a). In contrast, TMPRSS2 expression was similar at all time points sampled (Fig.2a). The classical model of circadian regulation is one of transcriptional control, however PCR quantification of ACE2 and TMPRSS2 transcripts showed no evidence of a rhythmic pattern in Calu-3 cells (Fig.2b), consistent with post-transcriptional regulation of ACE2. To extend these observations we quantified Ace2 and Tmprss2 transcripts in lung, liver and intestine harvested from light/dark entrained mice and observed limited evidence for a circadian pattern of expression (Supplementary Fig.3). Immunoblotting of ACE2 in the murine lung resulted in multiple bands of varying molecular weight, compromising the interpretation of these experiments. To further explore the role of circadian pathways in regulating ACE2 we shRNA silenced Bmal1, the major circadian transcriptional activator, and showed reduced ACE2 expression but a negligible effect on TMPRSS2 in Calu-3 cells (Fig.2c). To assess the impact of Bmal1 on SARS-CoV-2pp entry we infected the silenced Calu-3 cells and showed a significant reduction in pp infection, whereas
Our studies show circadian regulation of ACE2 in lung epithelial cells and show striking inhibitory effects of BMAL1 silencing and the synthetic REV-ERB agonist SR9009 treatment on ACE2 expression, SARS-CoV-2 entry and virus replication in lung epithelial cells. Of note, SARS-CoV-1 and alpha NL63 also require
ACE2 to enter cells23 and our data support a role for circadian factors in regulating the infection of these related coronaviruses. The human Ace2 promoter encodes putative binding sites for BMAL1/CLOCK and REV-ERB, however, we did not observe binding of these factors by chromatin immunoprecipitation-qPCR in Calu-3 cells (Supplementary Fig.7). In addition, there was no evidence for rhythmic expression of Ace2 transcripts in Calu-3 or in the lung, liver or intestine from entrained mice, consistent with bio-informatic analyses of published diurnal/circadian datasets from baboons24 and mice25. These results are consistent with recent studies showing a role for post-transcriptional and post-translational mechanisms in regulating the circadian clock26-28. There is an emerging role of microRNAs (miRNAs) in modulating the circadian clock29,30 and miRNAs have been reported to regulate ACE2 expression31, providing a possible post-transcriptional mechanism. Our studies show ACE2 to be a rhythmically ‘moving’ target and this could be relevant for evaluating new treatments targeting this step in the viral life cycle.

In addition to the circadian regulation of ACE2-mediated viral entry, we observed a marked suppression of SARS-CoV-2 RNA and genesis of infectious particles following SR9009 treatment and in Bmal1 silenced cells. Our bio-informatic analysis suggests that 30% of SARS-CoV-2 host factors are potentially BMAL1/REV-ERB regulated, highlighting a role for circadian-signalling to influence multiple steps in the SARS-CoV-2 life cycle. Further work is needed to characterise the circadian-dependent mechanisms of SARS-CoV-2 repression. A key finding from our study is the potential application of chronomodifying drugs for the treatment of COVID-1932. Dexamethasone is one of the few drugs that can reduce the severity of COVID-1933 and is known to synchronise circadian pathways34-36. Over the last decade, a number of compounds that target core clock proteins have been developed37, including REV-ERB38,39 and RORs40,41 agonists that have been shown to inhibit hepatitis C virus and HIV replication42,43. A report demonstrating REV-ERB dependent and independent effects of SR900944 suggests some additional off-target effects. We cannot exclude the possibility of additional pathways contributing to SR9009 antiviral activity; however, our use of genetic targeting approaches confirms a role for BMAL1 in regulating SARS-CoV-2 replication. REV-ERBα agonists can also impact the host immune response by suppressing IL-645 and so may offer a ‘two pronged’ approach to reduce viral replication and adverse host immune responses.

Circadian clocks can also impact on the pharmacokinetics and pharmacodynamics of drug responses46. Epidemiological47-49 and molecular evidence50 shows that night shift workers suffer circadian disruption and are at an increased risk of developing chronic inflammatory diseases. Identifying whether shift work is a risk factor for acquiring SARS-CoV-2 infection or developing moe severe disease could inform public health policy. Our observations raise questions as to how our in vitro studies translate to humans, where the time of exposure to SARS-CoV-2 may impact on the likelihood of infection, the host response, virus shedding, transmission and disease severity and are worthy of further investigation.
FIGURE LEGENDS

Figure 1. SARS-CoV-2 is circadian regulated.
(a) Circadian transcription–translation feedback loops control daily rhythmic expression of circadian genes. The molecular clock generates a cycle of 24-hour periodicity where a heterodimeric transcription factor BMAL1-CLOCK activates the transcription from E-box sequences in target gene promoters. Gene products feedback to repress the transcriptional activity of the heterodimeric activator. Ligand activation of REV-ERB represses the transcription of Bmal1. (b) Circadian infection protocol. (c) Calu-3 cells were synchronized by serum shock, inoculated with SARS-CoV-2 at an MOI of 0.1 or 0.01 for 1h and viral RNA measured after 24h. RNA levels are expressed relative to CT0; mean ± S.E.M., n = 2. (d) Calu-3 cells were serum shocked and synchronised cells inoculated with SARS-CoV-2pp or VSV-Gpp for 2h with luciferase activity measured 24h later. Data expressed relative to CT0; mean ± S.E.M., n = 6-12, Kruskal–Wallis test with Dunn’s multiple comparisons.

Figure 2. Circadian regulation of ACE2 dependent SARS-CoV-2 infection.
(a) Synchronised Calu-3 cells were assessed for ACE2, TMPRSS2 and BMAL1 expression together with housekeeping β-actin by western blotting; data are representative of two experiments. Densitometric analysis quantified ACE2 and TMPRSS2 in individual samples and normalized to their own β-actin loading control. SARS-CoV-2pp infection values from Fig.1d were plotted as a dotted line to illustrate concordance with ACE2 expression levels. (b) Synchronized Calu-3 cells were assessed for ACE2, TMPRSS2 and BMAL1 mRNA by qRT-PCR and expressed relative to CT0; data are the average of two independent experiments. (c) Calu-3 cells were transduced with lentivirus encoding shBmal1 or control and ACE2, TMPRSS2, BMAL1 and β-actin protein expression assessed by western blotting after 24 h. (d) Control or shBmal1 treated cells were infected with lentiviral pseudotypes expressing wild type (WT) or mutant (D614G or Furin KO) Spike variants or with VSV-Gpp. Viral entry (luciferase activity) was measured 24h later and data expressed relative to WT (mean ± S.E.M., n = 4-6, Mann–Whitney test).

Figure 3. REV-ERB agonist reduces ACE2 dependent SARS-CoV-2 entry and cell-cell fusion.
(a) Calu-3 cells were treated with SR9009 (5 or 10 µM) for 24h and assessed for ACE2 and TMPRSS2 expression together with housekeeping β-actin by western blotting. (b) Control or shBmal1 silenced Calu-3 cells were treated with SR9009 for 24h followed by infection with SARS-CoV-2pp or VSV-Gpp. After 24h viral entry was assessed by measuring luciferase activity and expressed relative to untreated cells (mean ± S.E.M., n = 3, Kruskal–Wallis ANOVA with Dunn’s test). (c) Control or SR9009 treated Calu-3 cells were infected with SARS-CoV-2 wild type (WT), D614G or furin mutant pp. Viral entry was expressed relative to untreated cells 24h later (mean ± S.E.M., n = 4-6, Kruskal–Wallis ANOVA with Dunn’s test). (d) Control or SR9009 treated Calu-3 cells were infected with SARS2-S-VSVpp and 24h later luciferase activity measured and data expressed relative to untreated cells (mean ± S.E.M., n = 3). Stats not needed for dose curves (e) Primary Bronchial Epithelial Cells (PBECs) were treated with SR9009 (20 µM) for 24h prior to SARS-CoV-2pp infection. Viral entry was measured as luciferase activity and data presented as mean ± S.E.M of triplicate wells from three independent donors relative to untreated control cells, Paired two-tailed t test. (f) SR9009 treated Huh-7 cells were assessed for ACE2 expression together with the housekeeping gene β-actin. (g) Huh-7 target cells expressing a split rLuc-GFP reporter (8-11) were treated overnight with SR9009 at the indicated concentrations or with DMSO vehicle (Ctrl.) before culturing with Spike expressing HEK-293T cells with the split rLuc-GFP (1-7) and SR9009 added.
for the indicated times. A representative image of SARS-CoV-2 Spike induced syncytia at 3 days post co-culture is shown (left panel). GFP-positive syncytia were quantified every 4h using an IncuCyte real-time imaging platform (right panel). Five fields of view were obtained per well at 10x magnification and GFP expression quantified by calculating the total GFP area using the IncuCyte analysis software.

Figure 4. REV-ERB agonist inhibits SARS-CoV-2 replication post-entry.
(a) Overlap between SARS-CoV-2 host factors and BMAL1 or REV-ERB regulated genes. Sequence analysis of promoters of genes encoding SARS-CoV-2 host genes with HOMER (Hypergeometric Optimization of Motif EnRichment tool) identifies a canonical E-box motif ‘CANNTG’ in 144 of the 153 genes and an ROR response element ‘RCGTCA’ in 80. (b) Bmal1 silencing reduces SARS-CoV-2 RNA. Calu-3 cells were transduced with lentivirus encoding shBmal1 or a control followed by SARS-Cov2 infection. Viral RNA was measured and expressed relative to control cells (mean ± S.E.M., n = 4-6, Mann–Whitney test). (c) Calu-3 cells were treated with 20 µM SR9009 either 24h before infection or from 2h post-inoculation with SARS-CoV-2. Cells were incubated for 22h, intracellular viral RNA quantified by qPCR and expressed relative to the untreated control. Infectivity of the extracellular viral particles was assessed by plaque assay using VERO cells. Data are presented as mean ± S.E.M. from n=3 independent biological replicates. Statistical significance was determined using Kruskal–Wallis ANOVA with Dunn’s test.
STAR METHODS

Materials. All reagents and chemicals were obtained from Sigma-Aldrich (now Merck) unless stated otherwise. REV-ERB agonist SR9009 was purchased from Calbiochem, US, dissolved in dimethyl sulfoxide (DMSO) and cytotoxicity determined by a Lactate dehydrogenase (LDH) assay (Promega, UK) or MTT assay (Sigma, UK). The BMAL1 promoter was amplified from genomic DNA using forward primer: 5’-CCGCTCGAGGGGACAACGGCGAGCTCGCAG-3’ with reverse primer: 5’-CCCAAGCTTCTGGCCCGCCCGCGCGCAAGTC-3’ and cloned into the pGL3 luciferase reporter vector (Promega, UK). The lenti-shBmal1 plasmid was purchased from Abmgood (UK). Recombinant ACE2-Fc was previously reported51.

Cell culture. Calu-3, Huh-7, HEK293T and VERO E6 cells were cultured in DMEM supplemented with 10% fetal bovine serum (FBS), 2mM L-glutamine, 100 U/ml penicillin and 10µg/ml streptomycin (all reagents from Life Technologies/Thermo Fisher). VERO E6/TMPRSS2 cells (a VERO E6 cell line stably overexpressing the TMPRSS2 gene, kindly provided by Dr Makoto Takeda at Department of Virology III, National Institute of Infectious Diseases52) were cultured in DMEM supplemented with 10% fetal bovine serum, 10 units/mL penicillin, 10 mg/mL streptomycin, 10 mM HEPES (pH 7.4), and 1 mg/mL G418 (Life Technologies, UK). All cell lines were maintained at 37°C and 5% CO2 in a standard culture incubator. Human PBECs were obtained using flexible fibreoptic bronchoscopy from healthy control volunteers under light sedation with fentanyl and midazolam. Participants provided written informed consent. The study was reviewed by the Oxford Research Ethics Committee B (18/SC/0361). Airway epithelial cells were taken using 2mm diameter cytology brushes from 3rd to 5th order bronchi and cultured in Airway Epithelial Cell medium (PromoCell, Heidelberg, Germany) in submerged culture.

SARS-CoV-2 pseudoparticle genesis and infection. SARS-CoV-2 lentiviral pp were generated by transfecting HEK-293T cells with p8.91 (Gag-pol), pCSFW (luciferase reporter) and a codon optimised expression construct pcDNA3.1-SARS-CoV-2-Spike, as previously reported53. The Furin cleavage site mutant was generated by mutagenesis of a pcDNA3.1 based clone expressing a C-terminally flag-tagged SARS-CoV-2 Spike protein (Wuhan-Hu-1 isolate; MN908947.3). The polybasic cleavage site TNSPRRA in SARS-CoV-2 Spike was replaced with the corresponding SARS-CoV variant sequence SLL. The pNBF SARS-CoV2 FL D614G mutant was a kind gift from Dr. Daniel Watterson and Dr. Naphak Modhiran at the University of Queensland. Supernatants containing viral pp were harvested at 48 and 72h post-transfection, frozen and stored at -80°C. As a control pp were generated that lacked a viral envelope glycoprotein (No Env) and were included in all infection experiments to control for non-specific uptake. This control was included in all pp experiments and the luciferase values obtained subtracted from values acquired with the SARS-CoV-2pp. To confirm spike-dependent infection, SARS-CoV-2pp were incubated with the anti-S-mAb FI-3A (1µg/mL)54 for 30 min prior to infection for all experiments.

SARS-CoV-2 VSV pp were generated as previously reported1 using reagents provided by Stefan Pöhlmann (Infection biology unit, German Primate Center, Göttingen, Germany). Briefly, HEK-293T cells were transfected with an expression construct encoding a c-terminal truncated version of SARS-CoV-2-S for assembly into VSV pp (pCG1-SARS-CoV-2-S-ΔC)55. After 24h, the transfected cells were infected with a replication-incompetent VSV (VSV*ΔG56) containing GFP and firefly luciferase reporter genes, provided by Gert Zimmer (Institute of Virology and Immunology, Mittelhäusern, Switzerland). After 1h, the cells
were washed with PBS before medium containing a VSV-G antibody (I1, mouse hybridoma supernatant from CRL-2700, ATCC) and supernatants harvested after 24h. The VSV*ΔG used for generating the pps was propagated in BHK-21 G43 cells stably expressing VSV-G. Viral titres were determined by infecting Calu-3 cells and measuring cellular luciferase after 48h.

**SARS-CoV-2 propagation and infection.** For infection experiments, the SARS-CoV-2 (Victoria 01/20 isolate) obtained from Public Health England was propagated in Vero E6 cells. Naïve Vero E6 cells were infected with SARS-CoV-2 at an MOI of 0.003 and incubated for 48-72h until visible cytopathic effects were observed. Culture supernatants were harvested, clarified by centrifugation to remove residual cell debris and stored at -80°C before measuring the infectious titre by plaque assay. Briefly, Vero E6 cells were inoculated with serial dilutions of SARS-CoV-2 stocks for 2h followed by addition of a semi-solid overlay consisting of 3% carboxymethyl cellulose (SIGMA). Cells were incubated for 72h, plaques enumerated by fixing cells using amido black stain and plaque-forming units (PFU) per mL calculated.

For infection of Calu-3 cells, cells were plated 24h before infecting with the stated MOI. Cells were inoculated with virus for 2h after which the unbound virus was removed by washing three times with with phosphate buffered saline (PBS). Unless otherwise stated, infected cells were maintained for 24h before harvesting for downstream applications.

**SARS-CoV-2 cell-cell fusion assay.** The SARS-CoV-2 cell-cell fusion assay was performed as described previously. Briefly, HEK-293T Lenti rLuc-GFP 1-7 (effector cells) and Huh-7.5 Lenti rLuc-GFP 8-11 (target cells) cells were seeded separately at 7.5x10^5 per well in a 6 well dish in 3mL of phenol-red free DMEM, supplemented with 10% FBS, 1% sodium pyruvate and 1% penicillin/streptomycin (10,000 U/mL) before culturing overnight at 37°C, 5% CO2. The effector cells were transfected with a plasmid expressing SARS-CoV-2 Spike or a blank vector. Meanwhile, target cells were diluted to 2x10^5/mL and 100µL seeded into a clear, flat-bottomed 96 well plate and mock-treated or treated with SR9009 (3µM or 7µM). The following day, the effector cells were harvested, diluted to 2x10^5/mL and 100µL cultured with the drug/target cell mix, with SR9009 concentrations maintained at 3µM or 7µM. To quantify GFP expression, cells were imaged every 4h using an IncuCyte S3 live cell imaging system (Essen BioScience). Five fields of view were obtained per well at 10x magnification and GFP expression determined by calculating the total GFP area using the IncuCyte analysis software.

**Oligonucleotides:**

| qPCR Primers (5’-3’) |  |
|----------------------|--|
| Human ACE2 forward: GGGATCAGAGATCGGAAGAAGAAA |  |
| Human ACE2 reverse: AGGAGGTCTGAACATCATCAGTG |  |
| Human TMPRSS2 forward: AGGTGAAAGCGGGTGTGAGG |  |
| Human TMPRSS2 reverse: ATAGCTGGTGACCTGACCTGAG |  |
| Human B2M forward: CTACACTGAATTCACCCCCACTG |  |
| Human B2M reverse: ACCTCCATGATGCTGCTTACATG |  |
| Human BMAL1: TaqMan Gene Expression Assay, ThermoFisher, Hs00154147 |  |
| Mouse Ace2 forward: AATTCACACTGAAGCTGGGAAGAA |  |
| Mouse Ace2 reverse: CCATCCAGTGTTCCACCCCA |  |
| Mouse Tmprss2 forward: CTCAGGCAACGGTACC |  |
| Mouse Tmprss2 reverse: GTCTCTGGTGAGGGCATCTA |  |
RT-qPCR. Cells were washed in PBS then lysed using Tri-reagent (Sigma), and mRNA extracted by phase separation. Equal amounts of cDNA were synthesised using the High Capacity cDNA Kit (Applied Biosystems) and mRNA expression determined using Fast SYBR master mix in a StepOne thermocycler (Applied Biosystems) using the ΔΔCt method. The lung tissues were lysed in TRI Reagent (Sigma) using a gentleMACS dissociator and RNA extracted using the manufacturers protocol. cDNAs were synthesized using the Maxima H Minus cDNA Synthesis Kit (Thermo Scientific) and mRNA quantified using iTaq Universal SYBR Green Supermix (Biorad) and specific primers on a QuantStudio 3 RT-PCR system (Applied Biosystems).

Immunoblotting. Cell lysates were prepared by washing cells with PBS, then lysed in Igepal lysis buffer (10mM Tris pH 7.5, 0.25M NaCl, 0.5% Igepal) supplemented with protease inhibitor cocktail (Roche Complete™) at 4°C for 5 min, followed by clarification by centrifugation (3 min, 12,000 rpm). Supernatant was mixed with Laemmli sample buffer, separated by SDS-PAGE and proteins transferred to polyvinylidene difluoride membrane (Immobilon-P, Millipore). Membranes were blocked in 5% milk in PBS/0.1% Tween-20, then incubated with anti-ACE2 (Abcam ab108252), anti-TMPRSS2 (SCBT sc-515727), anti-BMAL1 (Abcam Ab93806) or Anti-β-actin (Sigma A5441) primary antibodies and
appropriate HRP-conjugated secondary antibodies (DAKO). Chemiluminescence substrate (West Dura, 34076, Thermo Fisher Scientific) was used to visualize proteins using a ChemiDoc XRS+ imaging system (BioRad). Anti-β-actin-HRP conjugate (Abcam ab49900) and/or Coomassie brilliant blue staining was used to verify equal protein loading and densitometric analysis performed using ImageJ software (NIH).

**Bioinformatics.** The published 153 SARS-CoV-2 host factors were converted to Entrez gene names\(^\text{19}\). BMAL1 regulated genes were obtained from the published BMAL1 ChIP-seq in the mouse liver\(^\text{21}\). REV-ERB regulated genes were defined from published liver-specific loss of the REV-ERB paralogues\(^\text{22}\). Promoters (-1kb from TSS) of genes encoding SARS-CoV-2 host factors were analyzed with the HOMER (Hypergeometric Optimization of Motif EnRichment) tool for motif discovery (E-box motif CANNTG; RORE motif RGGTCA).

**Chromatin immuno-Precipitation (ChIP) and quantitative PCR:** 1x10^7 Calu-3 cells were harvested from 80% confluent 15cm plates and fixed with 1% formaldehyde (Sigma Aldrich 47608) for 10 min at room temperature before quenching with 125 mM glycine. Cells were washed twice with ice cold PBS, pelleted (800rpm, 10 min 4°C) and lysed in 500µL of Nuclear Extraction buffer (10 mM Tris pH 8.0, 10 mM NaCl, 1% NP-40) supplemented with a protease inhibitor cocktail (Roche Complete™). Samples were diluted 1:1 in ChIP Dilution Buffer (0.01% SDS, 1.1% Triton, 0.2mM EDTA; 16.7 mM Tris pH8.1, 167mM NaCl) and pulse sonicated using a Bioruptor R sonicator (Diagenode, U.K.) at high power for 15 min at 4°C (15 sec on, 15 sec off). Sonicated lysates were clarified by centrifugation at 1300 rpm for 10 min and precleared with Protein A agarose beads (Millipore, 16-156). Samples were immunoprecipitated with primary anti-BMAL1 (ChIP grade, Abcam Ab3350) or anti-REV-ERBα (ChIP grade, Abcam Ab181604) or IgG control mAb and precipitated with Protein A agarose beads. Precipitates were washed in low salt buffer (0.1% SDS, 1% Triton, 2 mM EDTA, 20 mM Tris pH8.1, 150mM NaCl), high salt buffer (0.1% SDS, 1% Triton, 2mM EDTA, 20 mM Tris pH 8.1, 500 mM NaCl), LiCl Buffer (1% Igepal, 1mM EDTA, 10 mM Tris pH 8.1, 250 mM LiCl, 1% sodium deoxycholate) and finally twice in TE wash buffer (10 mM Tris pH8.0, 1mM EDTA) before being eluted from the beads in 240µL of elution buffer (0.1 M NaHCO₃, 1% SDS). Complexes were reverse crosslinked in a heated shaker at 65°C overnight, 1400 rpm, in the presence of 200 mM NaCl. Eluates were treated with Proteinase K (SIGMA) and RNaseA (SIGMA) before cleanup using MiniElute PCR Purification columns (Qiagen). Samples were analysed on a LightCycler96 (Roche) using SYBR green qPCR mastermix (PCR Biosystems, UK). Fold enrichment was calculated for each sample relative to their own IgG controls.

**Animals.** Mouse experiments were carried out at the Institute of Viral and Liver Disease animal facility (approval number E-67-482-7). C57BL/6J male mice were purchased from Charles River and housed in individually ventilated cages under a 12/12 dark/light cycle with a ZT0 corresponding to 7am. After two weeks of acclimatisation, nine-week-old mice were sacrificed at different time points (ZT0, ZT4, ZT8, ZT12, ZT16, ZT20; n = 5/time point). Organs were harvested after exsanguination by intracardiac puncture, frozen in liquid nitrogen and kept at -80°C until further processing.

**Statistical Analysis.** All data are presented as mean values ± SEM. P values were determined using Mann-Whitney testing (two group comparisons) or with a Kruskal–Wallis ANOVA (multi group comparisons) using PRISM version 9. In the figures * denotes p < 0.05, ** < 0.01, *** <0.001, **** <0.0001, n.s. denotes non-significant.
DATA AVAILABILITY
The authors declare that all data supporting the findings of this study are available within the article and its Supplementary Information files or are available from the authors upon request.

AUTHOR CONTRIBUTIONS
XZ designed and conducted experiments and co-wrote MS; ST designed and conducted experiments and co-wrote MS; FW designed and conducted experiments; PACW designed and conducted experiments; HB conducted experiments and analysed data; JMH ran bio-informatic analysis; SBM provided PBECs; LM provided resources; LH and CB conducted experiments; CS designed and analyzed experiments; NZ conducted experiments; CC conducted experiments; HS designed and conducted experiments; LH conducted experiments; CB conducted experiments; AA conducted experiments; SW conducted analysis; TKH provided reagents; LS provided reagents; K-YAH provided reagents; KW provided experimental data; TSCH provided reagents; AJ provided experimental data; DB designed and conducted experiments and provided reagents; TFB designed and analyzed experiments, provided reagents and co-wrote MS; JAM designed the study and co-wrote the MS.

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CONFLICTS OF INTEREST
The other authors declare no financial interests.
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Figure 1. SARS-CoV-2 infection is circadian.
Figure 2. Circadian regulation of ACE2 dependent SARS-CoV-2 entry.
Figure 3. REV-ERB agonist reduces ACE2 dependent SARS-CoV-2 entry and cell-cell fusion.
Figure 4. REV-ERB agonist inhibits SARS-CoV-2 replication post-entry.