

## Life Sciences Reporting Summary

Nature Research wishes to improve the reproducibility of the work that we publish. This form is intended for publication with all accepted life science papers and provides structure for consistency and transparency in reporting. Every life science submission will use this form; some list items might not apply to an individual manuscript, but all fields must be completed for clarity.

For further information on the points included in this form, see [Reporting Life Sciences Research](#). For further information on Nature Research policies, including our [data availability policy](#), see [Authors & Referees](#) and the [Editorial Policy Checklist](#).

Please do not complete any field with "not applicable" or n/a. Refer to the help text for what text to use if an item is not relevant to your study. For final submission: please carefully check your responses for accuracy; you will not be able to make changes later.

### ▶ Experimental design

#### 1. Sample size

Describe how sample size was determined.

Sample size of n=3 was chosen for experiments where statistical analyses were performed (RaPID Western Blots, qPCR, CLIP-qPCR)

#### 2. Data exclusions

Describe any data exclusions.

No data was excluded

#### 3. Replication

Describe the measures taken to verify the reproducibility of the experimental findings.

Experiments were reliably reproduced.

#### 4. Randomization

Describe how samples/organisms/participants were allocated into experimental groups.

N/A

#### 5. Blinding

Describe whether the investigators were blinded to group allocation during data collection and/or analysis.

N/A

Note: all in vivo studies must report how sample size was determined and whether blinding and randomization were used.

#### 6. Statistical parameters

For all figures and tables that use statistical methods, confirm that the following items are present in relevant figure legends (or in the Methods section if additional space is needed).

- |                          |  |
|--------------------------|--|
| n/a                      | Confirmed  |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> The <u>exact sample size</u> ( <i>n</i> ) for each experimental group/condition, given as a discrete number and unit of measurement (animals, litters, cultures, etc.)   |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> A description of how samples were collected, noting whether measurements were taken from distinct samples or whether the same sample was measured repeatedly   |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> A statement indicating how many times each experiment was replicated   |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> The statistical test(s) used and whether they are one- or two-sided<br><i>Only common tests should be described solely by name; describe more complex techniques in the Methods section.</i>                       |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> A description of any assumptions or corrections, such as an adjustment for multiple comparisons  |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> Test values indicating whether an effect is present<br><i>Provide confidence intervals or give results of significance tests (e.g. P values) as exact values whenever appropriate and with effect sizes noted.</i> |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> A clear description of statistics including <u>central tendency</u> (e.g. median, mean) and <u>variation</u> (e.g. standard deviation, interquartile range)  |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> Clearly defined error bars in <u>all</u> relevant figure captions (with explicit mention of central tendency and variation)  |

See the web collection on [statistics for biologists](#) for further resources and guidance.

## ► Software

Policy information about [availability of computer code](#)

### 7. Software

Describe the software used to analyze the data in this study.

Prism 7 was used to plot data and perform statistics. R code used to process and analyze is included in the paper. Crapome was used to analyze spectral count data from RaPID-MS. DAVID v6.7 was used to analyze proteins bound by viral RNA.

For manuscripts utilizing custom algorithms or software that are central to the paper but not yet described in the published literature, software must be made available to editors and reviewers upon request. We strongly encourage code deposition in a community repository (e.g. GitHub). *Nature Methods* [guidance for providing algorithms and software for publication](#) provides further information on this topic.

## ► Materials and reagents

Policy information about [availability of materials](#)

### 8. Materials availability

Indicate whether there are restrictions on availability of unique materials or if these materials are only available for distribution by a third party.

No restrictions

### 9. Antibodies

Describe the antibodies used and how they were validated for use in the system under study (i.e. assay and species).

CELF1 (Santa Cruz Biotechnology, sc-20003) <https://www.scbt.com/scbt/product/cug-bp1-antibody-3b1>  
 IREB2 (Santa Cruz Biotechnology, sc-33682) <https://www.scbt.com/scbt/product/irp-2-antibody-7h6>  
 RC3H1 (Abcam, ab70196) <http://www.abcam.com/rc3h1-antibody-ab70196.html>  
 HA (Cell Signaling, C29F4) <https://www.cellsignal.com/products/primaryantibodies/ha-tag-c29f4-rabbit-mab/3724>  
 Streptavidin IRDye800 (Licor, 925-32230) <https://www.licor.com/bio/products/reagents/irdye/streptavidin/>  
 Streptavidin 800CW Goat-anti Mouse IgG (Licor, 925-32210)  
 Streptavidin 800CW Goat-anti Rabbit IgG (Licor, 925-32211)  
 anti-PAX6 (Biolegend 901301), anti-NCad (BD Biosciences 610920), anti-Tuj1 (Abcam ab78078), anti-NeuN (Abcam ab177487), anti-QKI (Abcam ab126742).

All antibodies were validated by source and cited in multiple papers.

### 10. Eukaryotic cell lines

a. State the source of each eukaryotic cell line used.

293T cells (Clontech), Huh7 (gift from Glenn lab, Stanford), U87 (ATCC), Human primary keratinocytes isolated from surgical specimens (Stanford)

b. Describe the method of cell line authentication used.

Cell lines not authenticated

c. Report whether the cell lines were tested for mycoplasma contamination.

Cell lines were tested for mycoplasma contamination every 2 weeks.

d. If any of the cell lines used are listed in the database of commonly misidentified cell lines maintained by [ICLAC](#), provide a scientific rationale for their use.

No commonly misused cell line used.

## ► Animals and human research participants

Policy information about [studies involving animals](#); when reporting animal research, follow the [ARRIVE guidelines](#)

### 11. Description of research animals

Provide all relevant details on animals and/or animal-derived materials used in the study.

No animals used

Policy information about [studies involving human research participants](#)

### 12. Description of human research participants

Describe the covariate-relevant population characteristics of the human research participants.

Study did not involve human subjects