## **Supplementary Information**

# **TORC1** Coordinates the Conversion of Sic1 from a Target to an Inhibitor of Cyclin-CDK-Cks1

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### INVENTORY OF SUPPLEMENTARY INFORMATION

#### Supplementary Figures

Figure S1 In vivo and in vitro specificity of the anti-Sic1-pThr<sup>33</sup> antibodies.

**Figure S2** Sic1<sup>T173A</sup>-expressing  $cdc4-2^{ts}$  cells are not defective in the clearance of Cln1, Cln2 or Clb5 when treated with rapamycin.

Figure S3 Cln2-HA<sub>3</sub> and Clb5-HA<sub>3</sub> interact with Cdc28 in vivo.

**Figure S4** Proliferating, Sic1<sup>T173A</sup>-expressing cells exhibit enhanced levels of Cln-/Clb-CDK activity.

#### **Supplementary Tables**

Table S1. Strains used in this study Table S2. Plasmids used in this study Table S3. Antibodies used in this study

**Supplementary References** 



**Figure S1** *In vivo* and *in vitro* specificity of the anti-Sic1-pThr<sup>5</sup>/-pThr<sup>33</sup> antibodies. (**a**) Sic1-pThr<sup>5</sup> and Sic1-pThr<sup>33</sup> levels were determined by immunoblot analyses using respective phospho-specific antibodies and extract of exponentially growing cells that expressed plasmid-encoded versions of myc<sub>13</sub>-tagged Sic1 and Sic1<sup>T5A</sup> (left panels), or myc<sub>13</sub>-tagged Sic1 and Sic1<sup>T33A</sup> (right panels). The levels of the Sic1-myc<sub>13</sub> variants were determined by using polyclonal anti-myc antibodies. The asterisk denotes an unspecific band. (**b**, **c**) Clb5-HA<sub>3</sub>-CDK immunocomplexes from exponentially growing yeast cells were used for *in vitro* kinase assays in which the bacterially-purified, N-terminal parts (encompassing the first 100 amino acids) of Sic1 (WT), Sic1<sup>T5A</sup> (T5A), and Sic1<sup>T33A</sup> (T33A) served as substrates. Sic1-pThr<sup>5</sup> (b) and Sic1-pThr<sup>33</sup> (c) levels were determined by immunoblot analyses using respective phospho-specific antibodies and the indicated GST-Sic1<sup>1-100</sup> variants that have been phosphorylated (+), or not (-), by Clb5-HA<sub>3</sub>-CDK. The input levels of the GST-Sic1<sup>1-100</sup> variants were determined by using polyclonal anti-GST antibodies.



**Figure S2** Sic1<sup>T173A</sup>-expressing  $cdc4-2^{ts}$  cells are not defective in the clearance of Cln1, Cln2 or Clb5 when treated with rapamycin.  $cdc4-2^{ts}$  and  $cdc4-2^{ts}$  sic1<sup>T173A</sup> strains expressing Cln1-myc<sub>13</sub>, Cln2-myc<sub>13</sub>, or Clb5-HA<sub>3</sub> were grown as in Fig. 1A. the levels of the tagged proteins were determined by immunoblot analyses using monoclonal anti-myc or anti-HA antibodies. Adh1 levels served as loading control.



**Figure S3** Cln2-HA<sub>3</sub> and Clb5-HA<sub>3</sub> interact with Cdc28 *in vivo*. Plasmid-expressed, HA<sub>3</sub>-tagged Cln2 or Clb5 were immunoprecipitated (IPed) from extracts of exponentially growing wild-type cells. Cell lysates (Input) and anti-HA immunoprecipitates (IP: anti-HA) were analyzed by immunoblotting with anti-HA (top panels), anti-Cdc28 (panels in the middle), and anti-Sic1 (bottom panels) antibodies. Please note that both Cln2-HA<sub>3</sub> and Clb5-HA<sub>3</sub> interact with Cdc28, while only Clb5-HA<sub>3</sub> is able to bind Sic1 as expected. Neither Cdc28 nor Sic1 were recovered in anti-HA immunoprecipitates from extracts of cells that carried an empty plasmid (-; 3<sup>rd</sup> lanes in all panels).



**Figure S4** Proliferating, Sic1<sup>T173A</sup>-expressing cells exhibit enhanced levels of Cln-/Clb5/6-CDK activity. For synchronization, exponentially growing WT cells expressing genomically-tagged Sic1-myc<sub>13</sub> or Sic1<sup>T173A</sup>-myc<sub>13</sub> were treated for 2 h with  $\alpha$ -factor (5  $\mu$ g ml<sup>-1</sup>). Following  $\alpha$ -factor release, samples were collected at the indicated time points. The phosphorylation levels of Thr<sup>5</sup> in Sic1 and Sic1<sup>T173A</sup> were determined by using phosphospecific anti-Sic1-pThr<sup>5</sup> antibodies and normalized with respect to the total levels (quantified by using anti-myc antibodies) of Sic1-myc<sub>13</sub> and Sic1<sup>T173A</sup>-myc<sub>13</sub>, respectively.

## Supplementary Tables

Table S1. Strains used in thi	s study
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Strain	Genotype	Source	Figure	
JK9-3D	MATa, leu2, his4, trp1, ura3, rme1, GAL, HMLa	Ref. (1)	2A/B, 3B, 2C, 4G, S1B/C, S3	
RL343-E1	[JK9-3D] <i>his3</i> , <i>HIS4</i> , <i>cdc28</i> ∆, pRS416- <i>cdc28</i> <sup>as</sup> (F88G)	Ref. (2)	4C	
YMM114	$[JK9-3D] cdc4-2^{ts}::kanMX$	Ref. (3)	1A/C	
YMM118-2D	[JK9-3D] cdc4-2 <sup>ts</sup> ::kanMX, sic1 <sup>T173A</sup> , EMP46::natMX	Ref. (3)	1A/C	
YMM67-1C	$[JK9-3D]$ sic $1\Delta$ ::kanMX	Ref. (3)	1A/B, S1A	
YMM237-1A	[JK9-3D] cdc4-2 <sup>ts</sup> ::kanMX, cdc28A::kanMX, pRS416-cdc28 <sup>as</sup>	This study	1 B/D	
YMM250-10C	$[JK9-3D]$ cdc4-2 <sup>ts</sup> ::kanMX, cdc28 $\Delta$ ::kanMX, pRS416-cdc28 <sup>as</sup> , sic1 <sup>T173A</sup> , EMP46::natMX	This study	1B/D	
YMM246-1C	[JK9-3D] cdc4-2 <sup>ts</sup> ::kanMX, CLB5-HA <sub>3</sub> ::TRP1, CLN1-myc <sub>13</sub> ::kanMX	This study	S2	
YMM247-4B	[JK9-3D] $cdc4-2^{ts}$ :: $kanMX$ , $CLB5-HA_3$ :: $TRP1$ , $CLN1-myc_{13}$ :: $kanMX$ , $sic1^{T173A}$ , $EMP46$ :: $natMX$	This study	S2	
YMM249-2B	[JK9-3D] cdc4-2 <sup>ts</sup> ::kanMX, CLB5-HA <sub>3</sub> ::TRP1, CLN2-myc <sub>13</sub> ::kanMX	This study	S2	
YMM252-2A	[JK9-3D] $cdc4-2^{ts}$ :: $kanMX$ , $CLB5-HA_3$ :: $TRP1$ , $CLN2-myc_{13}$ :: $kanMX$ , $sic1^{T173A}$ , $EMP46$ :: $natMX$	This study	S2	
MJA4090	[JK9-3D] cdc4-2 <sup>ts</sup> ::kanMX, CLB5-HA <sub>3</sub> ::TRP1	This study	S2	
MJA4091	MJA4091 $[JK9-3D]$ cdc4-2 <sup>ts</sup> ::kanMX, CLB5-HA <sub>3</sub> ::TRP1, sic1 <sup>T173A</sup> , EMP46::natMX			
YMM91	[JK9-3D] sic1 <sup>T173A</sup> , EMP46::natMX	Ref. (3)	2A/B, 3B,	
MJA490	[JK9-3D] CKS1-HA <sub>3</sub> ::kanMX	This study	2D-F, 4A	
MJA491	[JK9-3D] CKS1-HA <sub>3</sub> ::kanMX, sic1 <sup>T173A</sup> , EMP46::natMX	This study	2D-E, 4A	
MJA524-2B	$[JK9-3D] sic I^{R262A/L264A} - myc_{13} :: kanMX$	This study	3A	
YMM63	[JK9-3D] <i>SIC1-myc</i> <sub>13</sub> :: <i>kanMX</i>	Ref. (3)	3A, 4B, S4	
YMM98	[JK9-3D] sic1 <sup>T173A</sup> -myc13::kanMX, EMP46::natMX	Ref. (3)	3A, S4	
MJA523	[JK9-3D] sic1 <sup>R262A/L264A</sup> , EMP46::natMX	This study	3B	
MJA536	$[JK9-3D]$ cln1 $\Delta$ ::kanMX, cln2 $\Delta$ ::kanMX	This study	3B	
MJA528	[JK9-3D] sic1 <sup>T173A/R262A/L264A</sup> , EMP46::natMX	This study	3B	
YMM232-6A	[JK9-3D] <i>clb5∆::kanMX, clb6∆::kanMX, sic1<sup>T173A</sup>, EMP46::natMX</i>	This study	3B	
MJA544-2B	$[JK9-3D]$ $cln1\Delta$ :: $kanMX$ , $cln2\Delta$ :: $kanMX$ , $sic1^{T173A}$ , $EMP46$ :: $natMX$	This study	3B	
YMM253-11A	[JK9-3D] <i>clb5∆::kanMX, clb6∆::kanMX</i>	This study	3B	
MJA547	$[JK9-3D]$ clb5 $\Delta$ ::kanMX, cln1 $\Delta$ ::kanMX, cln2 $\Delta$ ::kanMX, sic1 <sup>T173A</sup> , EMP46::natMX	This study	3B	
MJA545	[JK9-3D] CKS1-HA <sub>3</sub> ::kanMX, CLB5-myc <sub>13</sub> ::kanMX, sic1 <sup>T173A</sup> , EMP46::natMX	This study	3D	
MJA546	[JK9-3D] CKS1-HA3::kanMX, CLB5-myc13:kanMX	This study	3D	
MJA531	[JK9-3D] CKS1-HA <sub>3</sub> ::kanMX, CLN2-myc <sub>13</sub> ::kanMX, sic1 <sup>T173A</sup> , EMP46::natMX	This study	3E	
MJA530	[JK9-3D] CKS1-HA <sub>3</sub> ::kanMX, CLN2-myc <sub>13</sub> ::kanMX,	This study	3E	
YMM231-1A	[JK9-3D] CKS1-HA <sub>3</sub> ::kanMX, mpk1 $\Delta$ ::kanMX	This study	4A	
YMM230-8A	[JK9-3D] CKS1-HA <sub>3</sub> ::kanMX, cdc55 <i>A</i> ::natMX	This study	4A	
YMM233-3A	[JK9-3D] CKS1-HA <sub>3</sub> ::kanMX, cdc55∆::natMX, sic1 <sup>T173A</sup> , EMP46::natMX	This study	4A	
MJA518	$[JK9-3D] cdc28\Delta, pRS416-cdc28^{as}$	This study	4C	
MJA519	$[JK9-3D]$ cdc28 $\Delta$ , pRS416-cdc28 <sup>as</sup> , rim15 $\Delta$ ::kanMX	This study	4C/H	
YMM58-1B	[JK9-3D] rim154::kanMX	This study	4E/F	
MM3D	[JK9-3D] $cdc28\Delta$ , pRS416- $cdc28^{as}$ , $rim15\Delta$ :: $kanMX$ , $LEU2$ :: $CYC1p-HHF2$ - $tDimer$	This study	4D	

Plasmid	Genotype	Source	Figure
pRS415	CEN, <i>LEU2</i>	Ref. (5)	
pMJA2881	[pRS415] ADH1p-SIC1-myc13	This study	S1A
pMJA3173	[pRS415] ADH1p-sic1 <sup>T5A</sup> -myc <sub>13</sub>	This study	S1A
pMJA3174	[pRS415] <i>ADH1p-sic1</i> <sup>T33A</sup> -myc <sub>13</sub>	This study	S1A
pMJA2995	[pRS415] <i>ADH1p-SIC1</i> <sup>150-285</sup> -myc <sub>13</sub>	This study	2F
pMJA2996	[pRS415] <i>ADH1p-sic1</i> <sup>150-285 - T173A</sup> -myc <sub>13</sub>	This study	2F
pRS416	CEN, URA3	Ref. (5)	S3
pMJA3038	[pRS416] ADH1p-CLB5-HA3	This study	2B/C, 3A, 4F/G, S3, S1B/C, S3
YCplac33	CEN, URA3	Ref. (6)	
JCE456	[YCplac33] ADH1p-CLN2-HA <sub>3</sub>	Ref. (7)	2A/C, 4D, 4F/G, S3
pAS2654	[YCplac33] ADH1p-LST7-HA3	Ref. (8)	4F
pGEX	GST	Ref. (9)	4G, S1B/C
pMMT2629	[pGEX] GST-SIC1	This study	2C
pMMT2630	[pGEX] GST-sic1 <sup>T173A</sup>	This study	2C
pMJA3029	[pGEX] GST-sic1 <sup>R262A/L264A</sup>	This study	2C
pMJA3037	[pGEX] GST-SIC1 <sup>1-100</sup>	This study	2C, 3A, 4G, S1B/C
pMJA3219	[pGEX] <i>GST-sic1</i> <sup>T5A(1-100)</sup>	This study	S1B/C
pMJA3220	[pGEX] GST-sic1 <sup>T33A(1-100)</sup>	This study	S1B/C
pVW995	[pGEX] GST-RIM15944-1149	Ref. (10)	4G
pVW827	CEN, LEU2, ADH1p-GST-RIM15	Ref. (10)	4F
pVW904	2µ, LEU2, TDH3p-RIM15-myc <sub>13</sub>	Ref. (10)	4H
pVW910	2μ, LEU2, TDH3p-rim15 <sup>T1075A</sup> -myc <sub>13</sub>	Ref. (10)	4H
pFD1008	CEN, TRP1, ADH1p-rim15 <sup>K823Y</sup> -GFP	Ref. (10)	4D/E

Table S2. Plasmids used in this study

Table S3. Antibodies used in this study

Name	Dilution	Source
Anti-Sic1	1:1'000	sc-50441 Santa Cruz
Anti-Adh1	1:200'000	Calbiochem
Anti-Sic1-pThr <sup>173</sup>	1:1'000	GenScript
Anti-Sic1-pThr <sup>5</sup>	1:1'000	GenScript
Anti-Sic1-pThr <sup>33</sup>	1:1'000	GenScript
Anti-myc	1:3'000	9E10; sc-40; Santa Cruz
Anti-HA	1:1'000	Enzo Life Sciences
Anti-GST	1:3'000	Bethyl Laboratories
Anti-Igo1-pSer <sup>64</sup>	1:1'000	GenScript
Anti-Igo1	1:1'000	Eurogentec
Anti-Cln2	1:1'000	Santa Cruz
Anti-Cdc28	1:300	Santa Cruz
Anti-Clb5	1:1'000	Santa Cruz
Anti-Sch9-pThr <sup>737</sup>	1:1'0000	GenScript
Anti-Sch9	1:1'000	GenScript
Anti-Rim15-pThr <sup>1075</sup>	1:10'000	Eurogentec
Goat anti-rabbit IgG HRP	1:3'000	Biorad
Goat anti-mouse HRP	1:3'000	Biorad
Donkey anti-goat HRP	1:5'000	Abcam
Goat anti-mouse IgG-Fcy HRP	1:5'000	Jackson Immunoresearch
Goat anti-mouse IgG, light chain HRP	1:5'000	Jackson Immunoresearch

#### **Supplementary References**

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