

SUPPLEMENTAL FIG S1

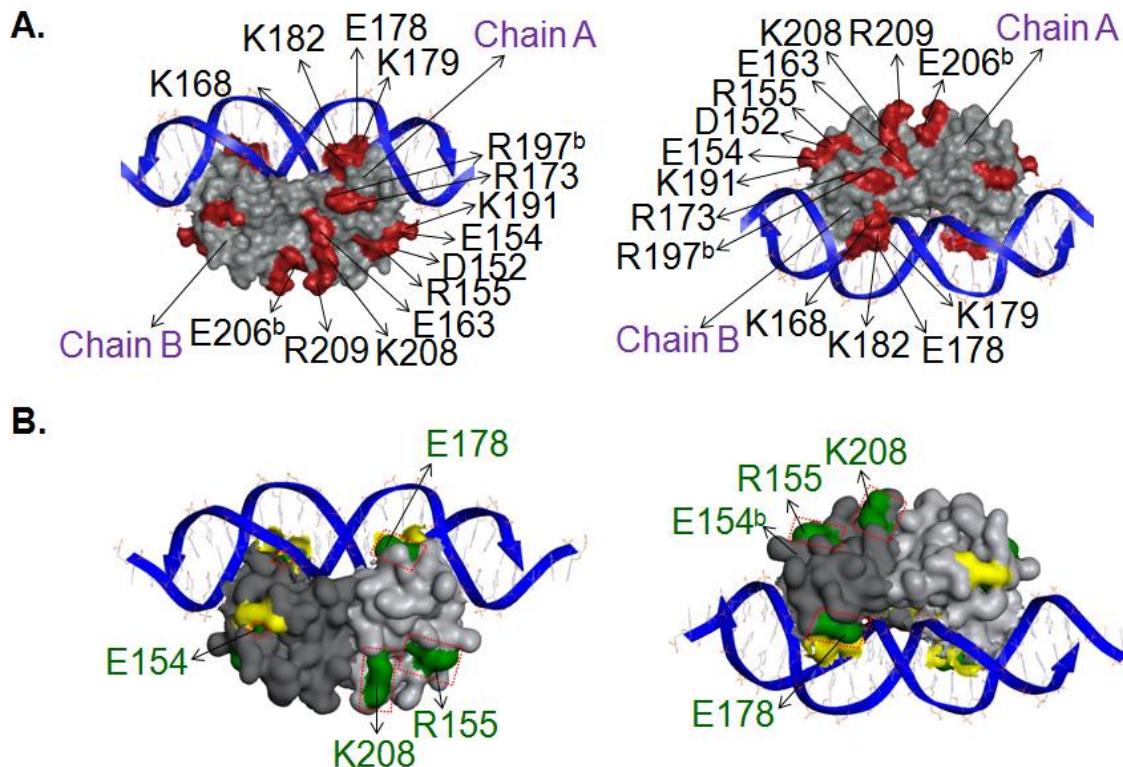
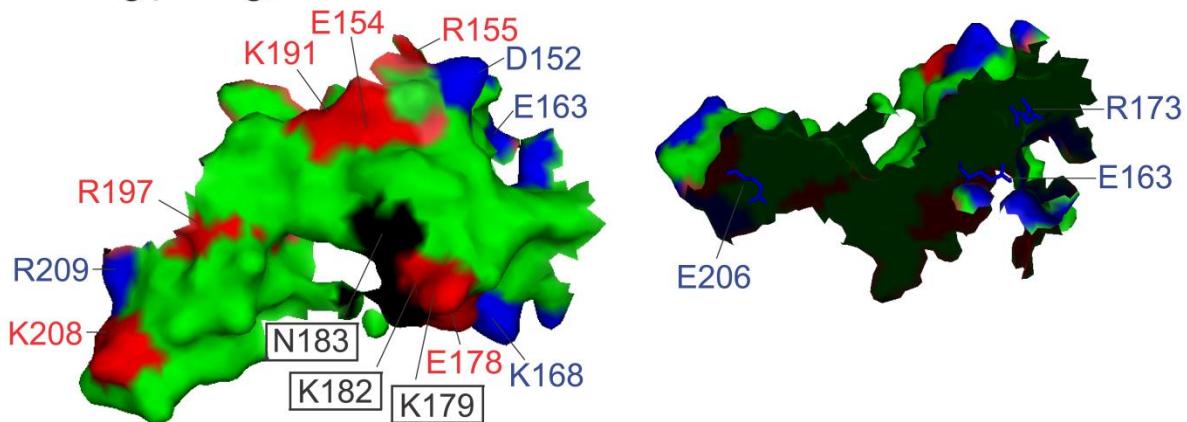


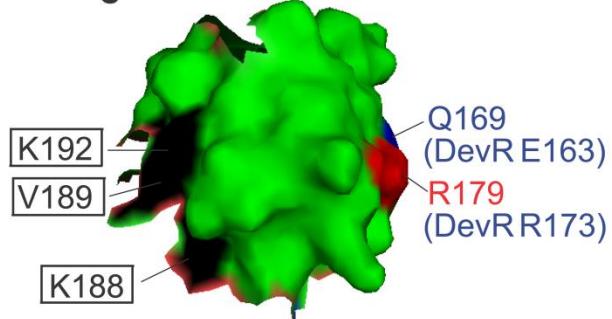
FIG S1 Location of amino acids important for DevR-mediated activation. A. All amino acids mutated in this study are colored in red. **B.** pc mutants (pc) are in dark green and boxed in red rectangles. Alanine substitution mutants defective in DNA binding are in yellow (K191, R197 and K179). The molecular surface representation of both chains of DevR_C (chains A and B in light grey and dark grey) was created from data in PDB file 3C3W (Wisedchaisri *et al.*, 2008) using a marching cubes algorithm of Discovery Studio 3.1 and probe radius of 1.4 Å. Individual surfaces of amino acid atoms were depicted by custom colors to highlight the mutations studied. The blue arrows represent the backbone of double stranded DNA, ‘b’ suffix for some residues (e.g. E154^b) indicates that this residue is located at the back.

SUPPLEMENTAL FIG S2

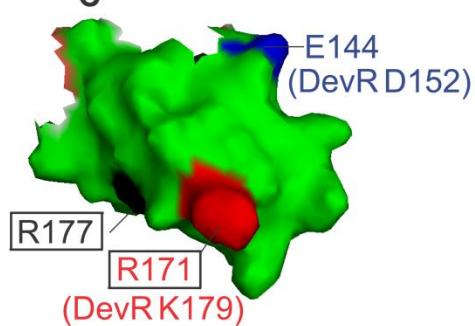
A. DevR_C (DosR_C)



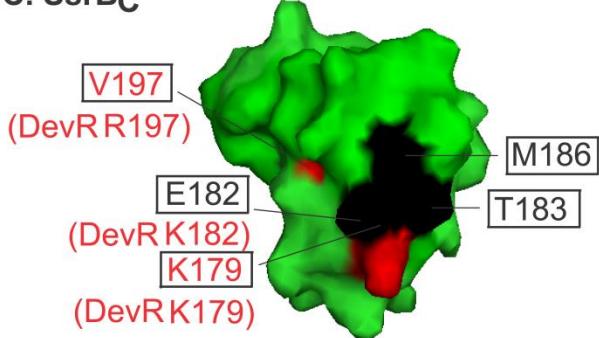
B. NarL_C



D. FixJ_C



C. SsrB_C



E.

$\alpha 7$

DevR 152	DQ <u>ER</u> TLLGLLS	162
NarL 158	P <u>RE</u> R <u>DILKLIA</u>	168
FixJ 144	E <u>ER</u> QVLSAVV	154
SsrB 152	L <u>RE</u> QVLKLID	162

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DevR 178	E <u>KTVK</u> NYVSRLLA <u>K</u>	191
NarL 184	E <u>STVK</u> HVK <u>HM</u> LKK	197
FixJ 170	P <u>RTVEVH</u> RANVMA <u>K</u>	183
SsrB 178	I <u>KTV</u> E <u>THRMNMMR</u> <u>K</u>	191

FIG S2 Comparison of DevR_C/DosR_C with NarL family members. A. Left panel: DevR_C/DosR_C structure was generated from PDB file 3C3W (Wisedchaisri *et al.*, 2008) using PyMol. Amino acid mutations that are proficient (in blue) and defective (in red) in transcription

activation are indicated. Right panel: Back view of DevR_C/DosR_C. **B-D.** The amino acids in the DevR orthologs corresponding to amino acids with known phenotypes in DevR are indicated using the same color code as in panel A. The residues with probable or confirmed DNA binding function are boxed. PDB files 3C3W, 1RNL, 1X3U and 2JPC were used to generate protein structures for DosR/ DevR, NarL, FixJ and SsrB, respectively using PyMol. **E.** Protein sequence alignment of DevR and orthologs for α 7 and α 9 helices. Three residues that are conserved among all regulators and display a pc phenotype (E, R and K) in DevR are in bold. The residues implicated in binding to DNA are underlined.

Supplemental Table S1. Primers used in this study

Primer name	Sequence (5'-3')	Application
hsp60BstBI f	CCGTTCGAAGGTGACCACAACGACGCGCCCGC	Cloning of <i>hsp60</i> promoter into pJFR19
hsp60NdeI r	CCGCATATGTGCGAAGTGATTCTCCGGATCG	(This study)
FLNdeI f	GCCCCATATGGTAAAGGTCTTCTGGTCGAT	Cloning of <i>devR</i> into plasmid pUS P _{hsp60}
FLNdeI r	CCGCATATGCTATCATGGTCCATCACCGGGTGG	(This study)
pJFRNew f	TCTTTGACTGAGCCTTCG	Orientation check of <i>hsp60</i> promoter and <i>devR</i> cloned in pJFR19 (This study)
pJFR-r	CCCCTGATTCTGTGGATAACCGTATTACC	
UGSTdevR f	<u>GCCGGATCC</u> CATGGTAAAGGTCTTCTGGTC	Cloning of <i>devR</i> or <i>devR</i> _{mut} into pGEX4T1
UGSTdevR r	<u>CCGGGATCC</u> CATGGTCCATCACCGG	(This study)
FL100B f	GCGCTGGATCCGATGGTAAAGGTCTTCTGGTC GAT	<i>devR</i> coding region cloned at BamHI and ClaI sites in pUAB100 as <i>devR</i> -mDHFR F[1,2] fusion, under <i>hsp60</i> promoter
FL100B r	GCGCTCATCGATTCTGGTCCATCACCGGGTGG CCG	
SigA200 f	GCGCTCCAATTGTTGTGGCAGCGACCAAAGCAA GCA	SigA coding region cloned at MfeI and ClaI sites in pUAB200 as <i>sigADHFR</i> F[3] fusion, under <i>hsp60</i> promoter
SigA200 r	GCGCTCATCGATTCCGTCCAGGTAGTCGCGCAG GAC	
D152 (<u>GAC</u> to <u>GCC</u>)	D152A-f TCAGGCCTTACCG <u>CCC</u> CAGGAGCGGACG D152A-r CGTCCGCTCTGG <u>GG</u> CGGTAAGGCCCTGA	Site Directed Mutagenesis [SDM] (This study)
E154 (<u>GAG</u> to <u>GCG</u>)	E154A-f CTTACCGACCAGG <u>CG</u> CGGACGCTACTG E154A-r CAGTAGCGCTCCG <u>CG</u> CTGGTCGGTAAG	SDM (This study)
R155 (<u>CGG</u> to <u>GCG</u>)	R155A-f ACCGACCAGGAG <u>CG</u> ACGCTACTGGCCTG R155A-r CAGGCCAGTAGCG <u>CG</u> CTGGTCGGTGGT	SDM (This study)
E163 (<u>GAG</u> to <u>GCG</u>)	E163A-f GGCCTGTTAGCG <u>CG</u> GGGCTGACCAAC E163A-r GTTGGTCAGGCC <u>CG</u> CTAACCAACAGGCC	SDM (This study)
K168 (<u>AAG</u> to <u>GCG</u>)	K168A-f GGCCTGACCAAC <u>CG</u> CGAGATCGCCGACCGA K168A-r TCGGTGGCGATCTG <u>CG</u> GGTTGGTCAGGCC	SDM (This study)
R173 (<u>CGA</u> to <u>GCA</u>)	R173A-f CAGATGCCGAC <u>CG</u> AAATGTTCTAGCCAA R173A-r TTGGCTAGGAACAT <u>CG</u> GTGGCGATCTG	SDM (This study)
E178 (<u>GAA</u> to <u>GCA</u>)	E178A-f ATG TTCCTAGCC <u>CG</u> AAAGACGGTGAAG E178A-r CTTCACCGTCTT <u>CG</u> GGCTAGAACAT	SDM (This study)
K179 (<u>AAG</u> to <u>GCG</u>)	K179A-f TTCTAGCCGA <u>AG</u> CGACGGTGAAGAACTAC K179A-r GTAGTTCTCACCGT <u>CG</u> CTTGGCTAGGAA	SDM (This study)
K191 (<u>AAG</u> to <u>GCG</u>)	K191A-f CGGTTGCTGGCC <u>CG</u> CTGGGATGGAACGT K191A-r ACGTTCCATGCCAG <u>CG</u> GGCCAGCAACCG	SDM (This study)
R197 (<u>CGG</u> to <u>GCG</u>)	R197A-f GGCATGGAACGT <u>CG</u> ACGCAAGCCGCGTA R197A-r TACCGCGGTTGCGT <u>CG</u> ACGTTCCATGCC	SDM (This study)
E206 (<u>GAG</u> to <u>GCG</u>)	E206A-f GTATTGCGAC <u>GG</u> CGTTGAAGCGCTCG E206A-r CGAGCGCTCAAC <u>CG</u> CGTGCAGAATAC	SDM (This study)
K208 (<u>AAG</u> to <u>GCG</u>)	K208A-f GCGACGGAGTTGG <u>CG</u> CGCTCGGCCACCC K208A-r GGGTGGCCCGAG <u>CG</u> CCAACCTCCGTGC	SDM (This study)
R209 (<u>CGC</u> to <u>GCC</u>)	R209A-f ACGGAGTTGAAG <u>GG</u> CTCGCGGCCACCCGGT R209A-r ACCGGGTGGCCCGAG <u>GG</u> CTTCAACTCCGT	SDM (This study)
D215 (<u>GAT</u> to <u>GCT</u>)	D215A-f CGGCCACCCGGT <u>CG</u> TCGACCATGACAA D215A-r TTGTCATGGTCA <u>CG</u> ACCGGGTGGCCG	SDM (This study)
LH1	CGAGTCGACAGAGCACGAAGGTCTGCCAGCGGAGGACCTTT GGCCCTGCGTCGACCGA	Gel shift assays (P+S box) (Chauhan <i>et al.</i> , 2009)
LH2	TCGGTCGACCCAGGGCAAAGGTCTCCGCTGGCGAGCCTTC GTGCTCTGGTCGACTCG	

RT16S f	ATGACGGCCTTCGGGTTGAA	Real Time RT PCR (Gautam <i>et al.</i> , 2011b)
RT16S r	CGGCTGCTGGCACGTAGTTG	
RT3134c f	CTGGCTGGGTCGGCCCTTA	Real Time RT PCR (Gautam <i>et al.</i> , 2011b)
RT3134c r	GCTGACCTGGGAGGTTGTCG	
devR f4	CCGATCTGCCGTGTCGATC	Real Time RT PCR (Taneja <i>et al.</i> , 2010)
devR r3	GTCCAGCCCCACATCTTT	
RTnarK2 f	CGGTTTGACGGTGGTCCGC'	Real Time RT PCR (Gautam <i>et al.</i> , 2011a)
RTnarK2 r	TCACGAAGCACGACCATGGCC	
RT1738 f	CGACGAACACGAAGGATTGA	Real Time RT PCR (Gautam <i>et al.</i> , 2011a)
RT1738 r	ACACCCACCAATTCCCTTTCC	
fdxA f	TGTCCGGTCGACTGTATCTATGA	Real Time RT PCR (Gupta <i>et al.</i> , 2011)
fdxA r	GGCAGG CCG GTTTGC	
RT2031c f	CGCACCGAGCAGAAGGA	Real Time RT PCR (Gautam <i>et al.</i> , 2011a)
RT2031c r	ACCGTGCGAACGAAGGAA	
RTTgsl f	CAGTGATTGGCTCGCTACAG	Real Time RT PCR (Gautam <i>et al.</i> , 2011b)
RTTgsl r	ACATCATTGATGGTGACGTCG	

GCT, GCC, GCA, GCG codes for Alanine; SDM- site directed mutagenesis; BamHI sites are underlined; Bold and underline represents nucleotide substitution; Bold represents substituted codon.