Evidence that Nek1 does not phosphorylate Rad54-S572 during recovery from IR

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ABSTRACT

A main focus of the work in our lab is on the activity of Tousled Like Kinase 1 (TLK1) in the area of DNA Damage and Repair. As one of its key interactor, TLK1 phosphorylates NIMA related kinase 1 (Nek1), and Nek1 was reported by Spies et al. to phosphorylate and regulate the activity of the key HRR protein Rad54[1], which suggested an intriguing signal transduction pathway: TLK1>Nek1>Rad54. In an effort to confirm such relations, we now report that we have not been able to reproduce key findings from that study. Specifically, we found that Nek1 does not phosphorylate RAD54-S572 as was reported. We generated Nek1-KO mouse NT1 cells[2] and Nek1-Knock-down in Hek293 (same cells as in Spies et al.), and the pRAD54-S572 signal does not change with our custom Ab, with or w/o IR. When we used an Ab from the Lobrich lab, it detected an immunoreactive band of wrong size for RAD54, which also did not change after IR even in synchronized G2 cells, contrary to their report. We also note that their P-assignment was based on guessing a weak consensus Nek1 sequence, and that site-directed mutagenesis of RAD54-S572 failed to yield biological effects in their in in vitro studies[1]. To conclusively establish that S572 is not a site of phosphorylation of Nek1, we carried out a IVK with purified Nek1 and RAD54 followed by MS analysis of the phosphatides, which revealed several but not S572. We also could not reproduce their copurification of Nek1-RAD54 by coIP, calling into question this interaction. Neither we could reproduce their results demonstrating the importance of Nek1 for HRR using the same SceI-mediated DR-GFP conversion assays.

INTRODUCTION

Each day our cells face many exogenous and endogenous challenges that threat the genomic stability. The DNA double strand break is the most lethal form of damage. The major mechanism to maintain the fidelity of genetic code is Homologous recombination repair (HRR). The central enzyme of this pathway is Rad54.[3] Rad54 is a well conserved protein of the Rad52 group, a generally well conserved group in eukaryotes. Rad54 is a motor protein that translocates along dsDNA in an ATP hydrolysis dependent manner.[4] In mammals, the RAD54 gene plays its most important role during early developmental stages.[5] The primary function of homologous recombination in mitotic cells is to repair double-strand breaks (DSBs) that form as a result of replication fork collapse, from processing of spontaneous damage, and from exposure to DNA-damaging agents like topoisomerase poisons.[6] Ideally known as the swiss-army knife of homologous recombination repair, Rad54 functions in different stages of HRR. Rad54 interacts with Rad51 and promotes DNA strand -exchange, translocates along donor template, remodels donor chromatin, promotes Branch-migration of Holliday Junctions.[7-10] With such multi-faceted role in vivo, any post-translational modification of Rad54 may alter its activity. Therefore, it is critical to understand the regulation of this error-free mode of DNA repair in cells.

RESULTS

Given the possibility of a TLK1>Nek1>Rad54 axis and its importance for HRR, we wanted to probe the pattern of pRad54-S572 in cells with perturbed TLK1 activity. In an initial communication with Marcus Lobrich, he stated that they had finished the original antiserum published in the Mol Cell paper but that they would ship us a replacement that was not well-characterized. We tested that antiserum during a time-course of recovery from IR of HEK293 cells, either control or expressing a dominant Nek1 kinase-hypoactive mutant (HEK N5)[11], which was expected to show reduced pRad54-S572 signal (Fig.1); and it was immediately obvious that there were problems. The main immunoreactive band ran on the gel faster than its expected position of ~80kDa for Rad54 (Fig1). The signal was not very clean as there were several other (presumably less specific) prominent bands. Most importantly, the intensity of the pRad54 did not increase after recovery from IR, as it was reported. Given the impossibility of reproducing that part of the work with the Ab they provided, we commissioned our own custom-made pRad54-S572 antiserum from

Thermofisher. After the 3d boost, an ELISA showed that the serum had a high titer (>2X10^5 dilution) and an affinity >100-fold for the phosphopeptide compared to unphosphorylated, even before affinity purification. To establish the specificity of the Ab and its dependence on Nek1, we carried out a siRNA-mediated knockdown of Nek1 in Hek293 cells, similarly to their Fig. 3D experiment. Despite the successful knock-down of Nek1 (Fig.2), we did not observe any change in the signal with our pRad54-S572 antiserum, which detected a band at the correct position for Rad54 and overlapping the signal obtained with a commercial (santa-cruz sc-166370) pan-Rad54 antiserum. This clearly indicated that Nek1 is unlikely to be the kinase responsible for phosphorylation of S572. Considering the possibility that there might still be sufficient Nek1 in the KD-cells to phosphorylate S572, we took advantage of a mouse PCa cell line in which we clonally knocked-out Nek1 via CRISPR/CAS9 to rule out such possibility. Surprisingly, there was no immunoreactive band with pRad54-S572 antiserum in either control or the Nek1-KO clone – note that these cells do express Rad54L and Rad54B (RNAseq data communicated by Xiuping Yu). In contrast, there was a strong band in HEK293, equally in control and Nek1-siRNA-KD cells (Fig. 3). This result suggests that despite the high sequence conservation between mouse and human (Fig. 3B), the pRad54-S572 is not found in mouse, and one should wonder whether such proposed phosphorylation is at all important. To probe this question further, we set out to establish if their report of Rad54-S572 phosphorylation after IR, and particularly in late G2, could be reproduced. We thus, synchronized Hek293 cells following release from G1/S block with HU and irradiated them in early G2 (Fig.4), repeating their experimental conditions. Contrary to what they reported, we found no increase in pRad54-S572 in either asynchronous or cells enriched G2 (Fig.5), for which 8h was reported in their paper as the maximal level of phosphorylation[1].

In order find truly unbiased evidence for the proposed phosphorylation of S572 by Nek1, we carried out an *in vitro* kinase (IVK) and MS phosphopeptides analysis of recombinant (and functional) RAD54 phosphorylated by highly purified Nek1, as described in[12]. Note that a IVK reaction does not imply that RAD54 in a true in vivo substrate of Nek1, but this was done (Fig. 6A) merely to replicate the conditions used by Spies et al. (Fig 3B) in order to assign the putative phosphorylated residue. Note that the only possible interpretation of their Fig 3B is that S572 is the primary, if not the only, phosphor-target of Nek1, since virtually all P32 labeling from the IVK reaction is lost with the GFP-RAD54-S572/A mutant protein. In complete disagreement with their published Fig.3B and their assumed phosphorylation of S572 as the only Nek1 target site, there were several residues phosphorylated in vitro – but not S572 (Fig. 6B). A complete list of all modified the peptides is shown (Sup Table 1), and multiple Ser, Thr, and Tyr were found – note that Nek1 is a dual specificity kinase[13]. Note also that our preparation of Nek1 was fully active, as indicated also by its autophosphorylation at Y315 (MS data not shown), which is diagnostic of its level of activity. In conclusion, their assignment of S572 as the only phosphorylation site by Nek1 cannot be reproduced when analyzed correctly by MS instead of relying on a putative consensus sequence; and their results of Fig.3B remains very difficult to explain.

Finally, we were not able to reproduce the reported interaction between Nek1 and Rad54 by coIP, although we cannot rule out that this could be due to a technical failure. Nonetheless, key findings of their paper could not be reproduced. These include a failure to attribute the phosphorylation of S572 to Nek1, its dependency on IR-induced damage, and in general the significance of such Rad54 modification for HRR.

To establish if Nek1 has any significant role in HRR, we carried out SceI-mediated DR-GFP conversion assays in Hek293 cells overexpressing siRNA-resistant wt-Nek1 or the Nek1-T141A dominant negative mutant (HEK N5), similar to the work reported in Fig. 4 of the Spies et al. paper. Again, contrary to their findings, we found that wt-Nek1 did not stimulate HRR and that the N5 overexpressing cells showed insignificant effects on HRR (Fig. 7). Depletion of endogenous Nek1 with siRNA did not change the picture, demonstrating that it is not the endogenous Nek1 that misrepresents the fact that neither wt nor a dominant hypomorphic mutant

of Nek1 has any effect on HRR, in contradiction of the results reported in Fig.4 of the Spies et al paper for the S572A overexpressing mutant. In summary, we were not able to reproduce any of the key results of their paper about the importance of Nek1 for HRR.

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Figure Legends

Figure 1. 2X 10^6 HEK293 cells were seeded 24hrs prior to treatment and then cells were treated with 10Gy of I.R and allowed to recover for indicated times (hrs). 40µg of untreated control (C) and samples were loaded in 8% SDS-PAGE gel and probed with aliquot of antibody sample from Lobrich lab. Tubulin probed as loading control.

Figure 2. 0.3X10^6 HEK293 cells seeded in 6 well plate 24hrs prior to siRNA treatment. Lipofectamine 3000 was used as transfection reagent with indicated amount of siNek1. 40ug of total samples loaded in 8% SDS-PAGE gel and probed with indicated antibodies (Nek1- sc 398813; Rad54- sc 166370; pRad54- Custom generated-Thermofisher; GAPDH- CST #2118)

Figure 3. A) 40µg of cell lysate from NEK1 WT (Cnt), K.O #6 clone (in NT1 cells) and siRNA treated HEK293 cells and control (Cnt) loaded in 8% SDS-PAGE gel and probed for Nek1 and pRad54 (custom generated) antibody. Actin band shown as loading control. B) Human Rad54 and mouse Rad54 protein sequence alignment shown for the conservation of the S572 site.

Figure 4. Cell cycle analysis of HEK293 cells synchronized (asynchronous -Asyn) at G1/S with Hydroxyurea treatment and released for indicated times (as shown in schematic).

Figure 5. Synchronised and aynchronised HEK293 cells were untreated or irradiated (5 Gy) at G2 and allowed to recover for indicated times. 40ug of cell lysate was loaded in 8% SDS-PAGE gel and probed with indicated antibodies. Ku-70 probed for loading control.

Figure 6. Determination of the RAD54 phosphopetides obtained by IVK with purified Nek1. A) Picture of the Coomassie stained gel confirming the purity of the proteins is shown. B) Phospho-peptide map of purified human Rad54 protein in IVK assay with purified Nek1.

Figure 7. A. SceI-mediated DR-GFP conversion assays. Hek293 were stably transfected with a DR-GFP cassette and then with wt-Nek1 or dominant-negative Nek1-T141A (N5) expression constructs. HRR assays were then carried out 48h after transfection of SceI by FACS analysis of the proportion of GFP converted cells. B. Expression of wt-Nek1 or dominant-negative Nek1-T141A. Where indicated, siRNA to endogenous Nek1 was included 24h earlier (demonstration of the efficacy of the knock-down was already published in [11]).

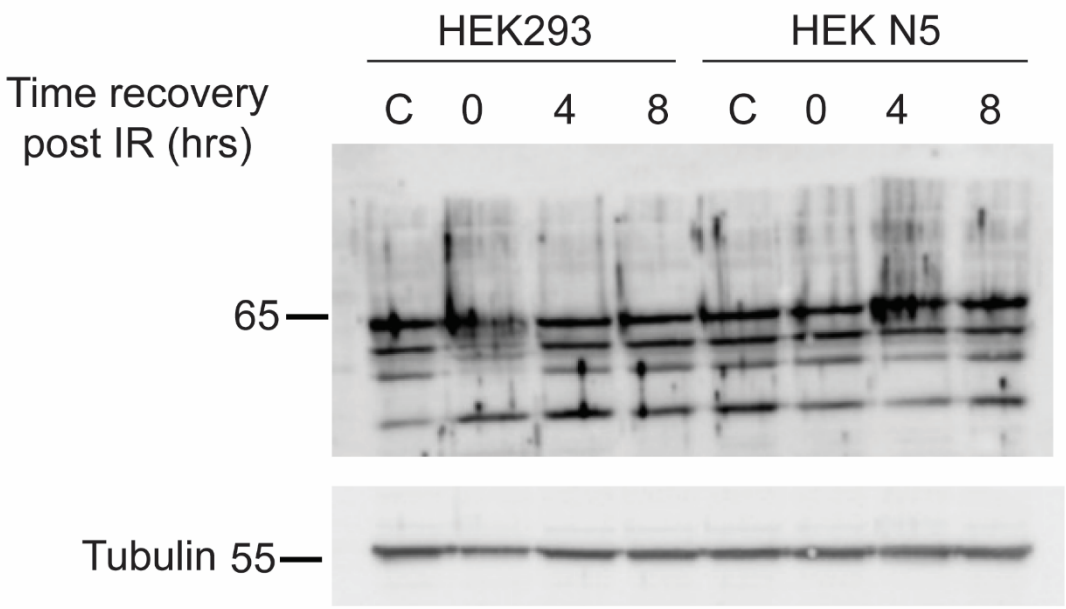

HEK293
HEK N5
Time recovery post IR (hrs)
C 0 4 8 C 0 4 8
65
Tubulin 55

Fig. 1

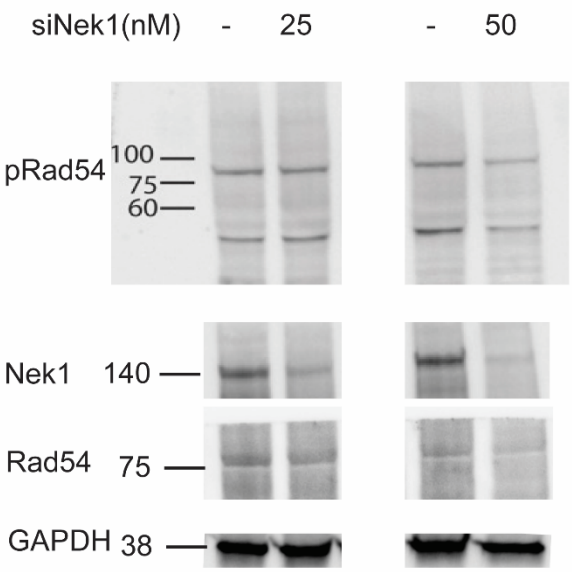

siNek1(nM) - 25 - 50
pRad54
100
75
60
Nek1 140
Rad54 75
GAPDH 38

Fig. 2

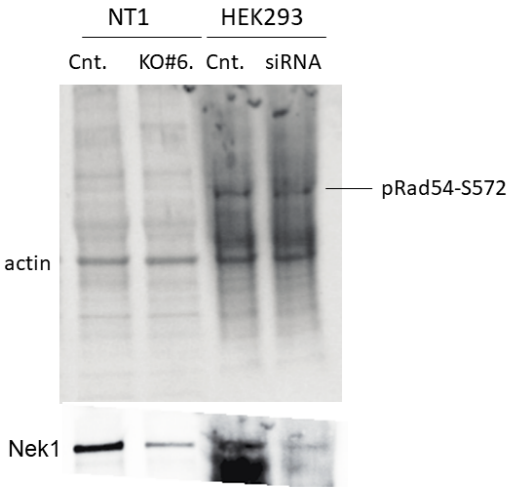
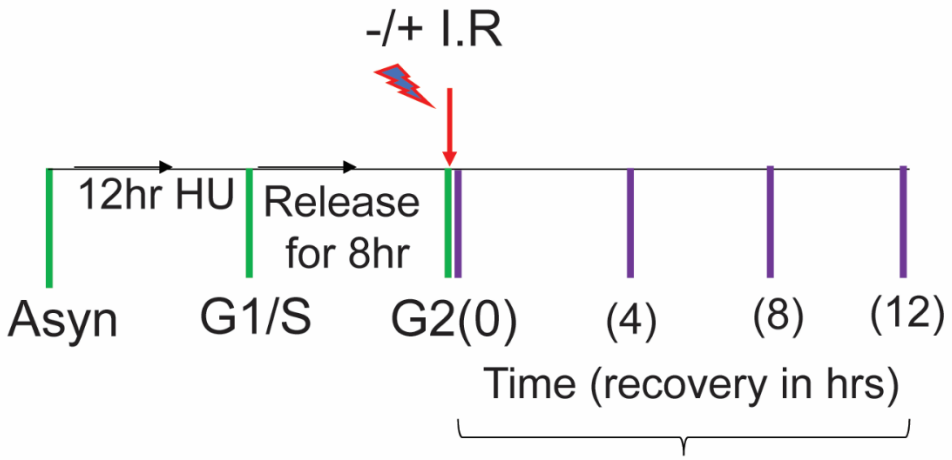


Fig.3A

```
hRad54 VRLDGTMSIKKRAKVVERFNSPSSPDFVFMLSSKAGGCGLNLIGANRLVMFDPDWNPAND 600
mRad54 VRLDGTMSIKKRAKVVERFNSPSSPDFVFMLSSKAGGCGLNLIGANRLVMFDPDWNPAND 600
```

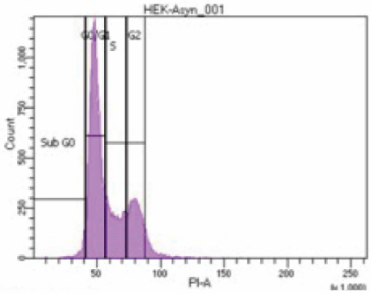
Fig.3B



Asyn

Tube: Asyn_001

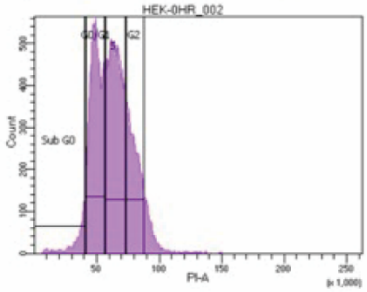
Population	#Events	%Parent	%Total
All Events	29,274	####	100.0
P1	27,829	95.1	95.1
P2	20,774	74.6	71.0
P3	20,564	99.0	70.2
P4	19,942	97.0	68.1
G0/G1	10,577	53.0	36.1
G2	3,469	17.4	11.9
S	3,453	17.3	11.8
Sub G0	758	3.8	2.6



G1/S

Tube: 0HR_002

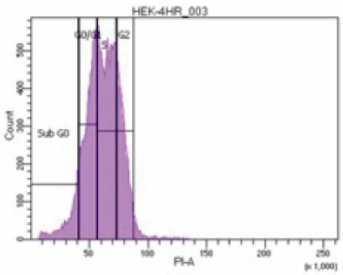
Population	#Events	%Parent	%Total
All Events	25,221	####	100.0
P1	24,400	96.7	96.7
P2	20,271	83.1	80.4
P3	20,083	99.1	79.6
P4	19,887	99.0	78.9
G0/G1	6,536	32.9	25.9
G2	3,417	17.2	13.5
S	7,118	35.8	28.2
Sub G0	918	4.6	3.6



After 4hr release-S phase

Tube: 4HR_003

Population	#Events	%Parent	%Total
All Events	26,999	####	100.0
P1	25,178	93.3	93.3
P2	20,579	81.7	76.2
P3	20,381	99.0	75.5
P4	19,937	97.8	73.8
G0/G1	5,760	28.9	21.3
G2	3,457	17.3	12.8
S	7,536	37.8	27.9
Sub G0	1,676	8.4	6.2



After 8hr release- G2 phase

Tube: 8HR_004

Population	#Events	%Parent	%Total
All Events	27,504	####	100.0
P1	25,393	92.3	92.3
P2	20,421	80.4	74.2
P3	20,234	99.1	73.6
P4	19,923	98.5	72.4
G0/G1	4,704	23.6	17.1
G2	7,138	35.8	26.0
S	4,641	23.3	16.9
Sub G0	1,637	8.2	6.0

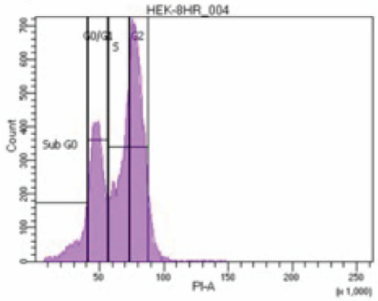


Fig.4

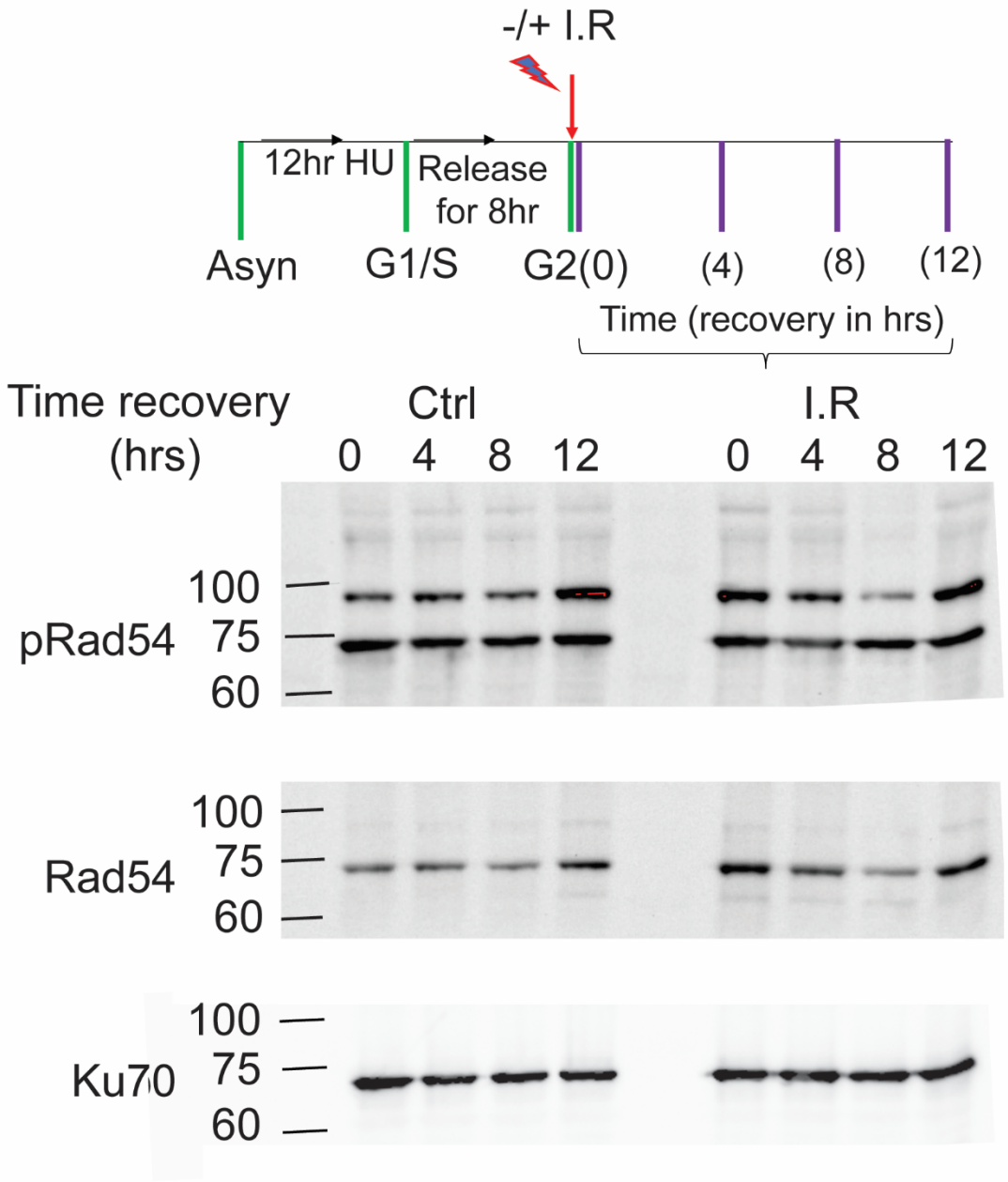

-/+ I.R
12hr HU
Release
for 8hr
Asyn
G1/S
G2(0)
(4)
(8)
(12)
Time (recovery in hrs)
Time recovery
(hrs)
Ctrl
I.R
0
4
8
12
0
4
8
12
100
pRad54
75
60
100
Rad54
75
60
100
Ku70
75
60

Fig. 5

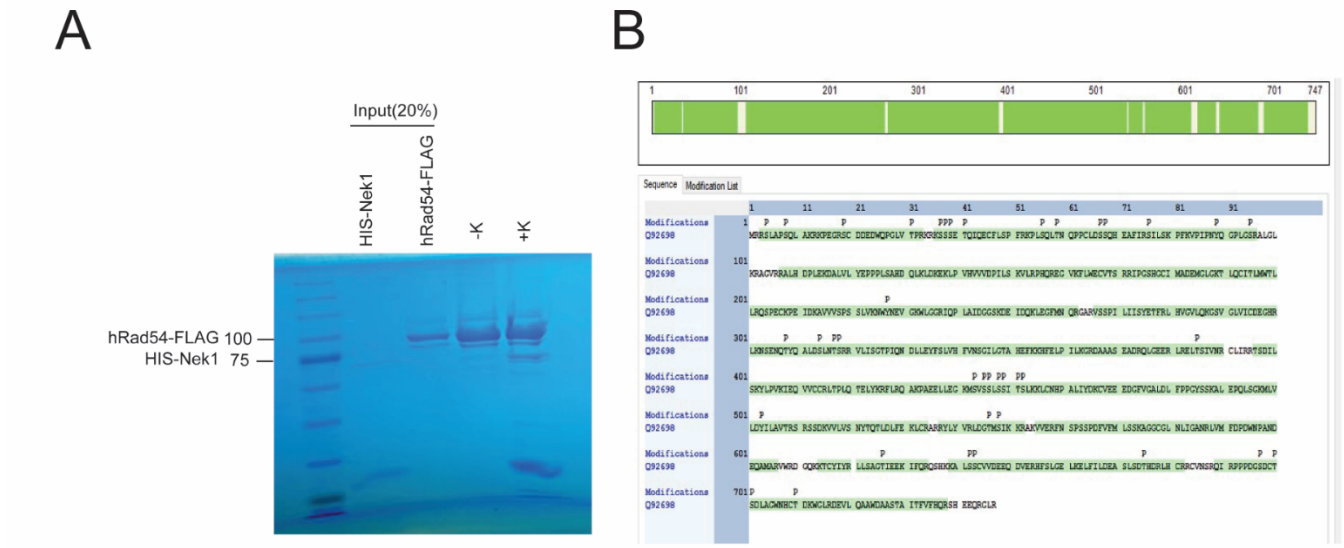

A
B
Input(20%)
HIS-Nek1
hRad54-FLAG
-K
+K
hRad54-FLAG 100
HIS-Nek1 75
Sequence
Modification List

Fig. 6

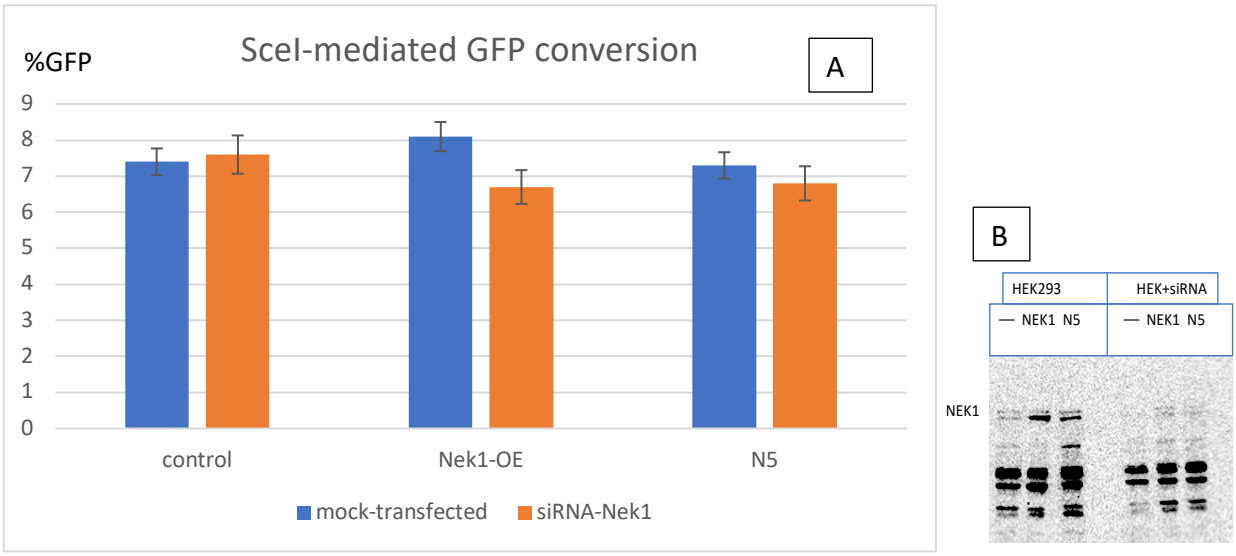

%GFP
SceI-mediated GFP conversion
A
9
8
7
6
5
4
3
2
1
0
control
Nek1-OE
N5
mock-transfected
siRNA-Nek1
B
HEK293
HEK+siRNA
— NEK1 N5
— NEK1 N5
NEK1

Fig. 7

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Supplemental Table 1

Confidence	Sequence	Modificatic	Qvality PEP	Qvality q-va	# Protein G	# Proteins	# PSMs	Master Pro
High	LVMFDPDV	2xOxidatio	3.95E-05	0.004739	1	1	79	Q92698
High	IEQVVCCR	2xCarbami	0.000465	0.004739	1	1	40	Q92698
High	TLQCITLMV	2xCarbami	0.103627	0.004739	1	1	1	Q92698
High	KLCNHPALI	2xCarbami	2.21E-07	0.004739	1	1	59	Q92698
High	LCNHPALIY	2xCarbami	4.37E-07	0.004739	1	1	97	Q92698
High	QIRPPPDGS	2xCarbami	4.81E-06	0.004739	1	1	5	Q92698
High	QIRPPPDGS	2xCarbami	9.26E-11	0.004739	1	1	38	Q92698
High	YLPVKIEQV	2xCarbami	0.0002	0.004739	1	1	58	Q92698
High	MLVLDYILA	1xPhospho	0.186682	0.004739	1	1	1	Q92698
High	NWYNEVGK	1xPhospho	0.025781	0.004739	1	1	4	Q92698
High	LLSAGTIEEK	1xPhospho	0.0006	0.004739	1	1	1	Q92698
High	LRELTSIVNF	1xPhospho	0.128289	0.004739	1	1	1	Q92698
High	ELFILDEASL	1xPhospho	0.005986	0.004739	1	1	2	Q92698
High	SLAPSQLAK	1xPhospho	0.0191	0.004739	1	1	6	Q92698
High	RSLAPSQLA	1xPhospho	0.00741	0.004739	1	1	2	Q92698
High	SILSKPFK	1xPhospho	0.001382	0.004739	1	1	3	Q92698
High	VPIPNYQGF	1xPhospho	0.025547	0.004739	1	1	4	Q92698
High	SILSKPFKVF	1xPhospho	0.000224	0.004739	1	1	16	Q92698
High	LDGTMSIK	1xPhospho	0.010477	0.004739	1	1	3	Q92698
High	LKNSENQTY	1xPhospho	0.003824	0.004739	1	1	9	Q92698
High	LKNSENQTY	1xPhospho	0.011858	0.004739	1	1	8	Q92698
High	MSVSSLSSI	1xPhospho	0.003098	0.004739	1	1	12	Q92698
High	NSENQTYQ	1xPhospho	0.000117	0.004739	1	1	5	Q92698
High	NSENQTYQ	1xPhospho	0.000167	0.004739	1	1	16	Q92698
High	LVMFDPDV	1xOxidatio	2.01E-05	0.004739	1	1	90	Q92698
High	LDGTMSIK	1xOxidatio	0.001572	0.004739	1	1	39	Q92698
High	LDGTMSIK	1xOxidatio	0.00119	0.004739	1	1	15	Q92698
High	LEGFMNQR	1xOxidatio	0.003824	0.004739	1	1	27	Q92698
High	IQPLAIDGG	1xOxidatio	1.81E-06	0.004739	1	1	82	Q92698
High	MLVLDYILA	1xOxidatio	4.03E-06	0.004739	1	1	108	Q92698
High	MSVSSLSSI	1xOxidatio	4.98E-05	0.004739	1	1	22	Q92698
High	MSVSSLSSI	1xOxidatio	2.69E-06	0.004739	1	1	57	Q92698
High	FLRQAKPAE	1xOxidatio	0.090318	0.004739	1	1	13	Q92698
High	FNSPSSPDF	1xOxidatio	0.000102	0.004739	1	1	52	Q92698
High	DEIDQKLEG	1xOxidatio	0.003576	0.004739	1	1	8	Q92698
High	ALEPQLSGK	1xOxidatio	0.035362	0.004739	1	1	1	Q92698
High	EGVKFLWE	1xCarbami	0.000689	0.004739	1	1	3	Q92698
High	RIPGSHGCI	1xCarbami	7.72E-07	0.004739	1	1	48	Q92698
High	RIPGSHGCI	1xCarbami	2.86E-07	0.004739	1	1	69	Q92698
High	GSVGLVICD	1xCarbami	0.000188	0.004739	1	1	198	Q92698
High	GSVGLVICD	1xCarbami	0.00034	0.004739	1	1	1	Q92698
High	RIPGSHGCI	1xCarbami	6.88E-08	0.004739	1	1	110	Q92698
High	IPGSHGCIM	1xCarbami	0.000937	0.004739	1	1	22	Q92698
High	IPGSHGCIM	1xCarbami	6.54E-06	0.004739	1	1	37	Q92698
High	IPGSHGCIM	1xCarbami	1.5E-08	0.004739	1	1	19	Q92698
High	KALSSCVVD	1xCarbami	1.58E-05	0.004739	1	1	14	Q92698
High	ALSSCVVDE	1xCarbami	0.002291	0.004739	1	1	2	Q92698

High	ALSSCVVDE	1xCarbami	7.16E-06	0.004739	1	1	25	Q92698
High	ALSSCVVDE	1xCarbami	0.000106	0.004739	1	1	3	Q92698
High	FLWECVTSF	1xCarbami	6.98E-05	0.004739	1	1	162	Q92698
High	FLWECVTSF	1xCarbami	0.033493	0.004739	1	1	1	Q92698
High	QSPECKPEI	1xCarbami	9.62E-05	0.004739	1	1	36	Q92698
High	QSPECKPEI	1xCarbami	6.08E-05	0.004739	1	1	17	Q92698
High	TLQCITLMV	1xCarbami	8.25E-05	0.004739	1	1	119	Q92698
High	AGGCGLNL	1xCarbami	2.13E-05	0.004739	1	1	22	Q92698
High	AGGCGLNL	1xCarbami	0.008985	0.004739	1	1	1	Q92698
High	TLQCITLMV	1xCarbami	3.74E-06	0.004739	1	1	152	Q92698
High	KLCNHPALI	1xCarbami	8.56E-06	0.004739	1	1	68	Q92698
High	KTCYIYR	1xCarbami	0.000716	0.004739	1	1	9	Q92698
High	SCDDEDWC	1xCarbami	0.01714	0.004739	1	1	6	Q92698
High	LCNHPALIY	1xCarbami	4.64E-05	0.004739	1	1	131	Q92698
High	SCDDEDWC	1xCarbami	2.62E-06	0.004739	1	1	20	Q92698
High	TCYIYR	1xCarbami	0.002952	0.004739	1	1	29	Q92698
High	FNSPSSPDF	1xCarbami	0.00245	0.004739	1	1	2	Q92698
High	FNSPSSPDF	1xCarbami	2.5E-06	0.004739	1	1	6	Q92698
High	CVEEEDGFV	1xCarbami	3.49E-06	0.004739	1	1	154	Q92698
High	ELFILDEASL	1xCarbami	0.00092	0.004739	1	1	2	Q92698
High	ELFILDEASL	1xCarbami	0.000216	0.004739	1	1	11	Q92698
High	VVLVSNYTC	1xCarbami	0.000245	0.004739	1	1	3	Q92698
High	MSVSSLSSIT	1xCarbami	6.57E-06	0.004739	1	1	1	Q92698
High	KPLSQLTNC	1xCarbami	0.116337	0.004739	1	1	1	Q92698
High	KPLSQLTNC	1xCarbami	5.2E-07	0.004739	1	1	27	Q92698
High	KPLSQLTNC	1xCarbami	1.19E-09	0.004739	1	1	119	Q92698
High	KSSSETQIQE	1xCarbami	0.000282	0.004739	1	1	11	Q92698
High	KSSSETQIQE	1xCarbami	2.31E-06	0.004739	1	1	43	Q92698
High	SSSETQIQEC	1xCarbami	5.36E-05	0.004739	1	1	24	Q92698
High	SSSETQIQEC	1xCarbami	4.94E-06	0.004739	1	1	60	Q92698
High	ALEPQLSGK		0.004906	0.004739	1	1	35	Q92698
High	ALEPQLSGK		0.000164	0.004739	1	1	6	Q92698
High	ALEPQLSGK		0.00071	0.004739	1	1	7	Q92698
High	ALHDPLEK		0.004028	0.004739	1	1	13	Q92698
High	ALHDPLEKD		4.14E-08	0.004739	1	1	285	Q92698
High	ALHDPLEKD		1.71E-09	0.004739	1	1	153	Q92698
High	AVVVSPSSL		7.95E-05	0.004739	1	1	44	Q92698
High	AVVVSPSSL		2.94E-05	0.004739	1	1	3	Q92698
High	DAAASEADF		0.001249	0.004739	1	1	1	Q92698
High	DAAASEADF		0.000106	0.004739	1	1	23	Q92698
High	DAAASEADF		0.026255	0.004739	1	1	1	Q92698
High	DALVLYEPP		5.86E-05	0.004739	1	1	68	Q92698
High	DALVLYEPP		1.32E-06	0.004739	1	1	27	Q92698
High	DALVLYEPP		5.06E-07	0.004739	1	1	10	Q92698
High	DEIDQKLEG		6.32E-05	0.004739	1	1	11	Q92698
High	DEVLQAAW		4.45E-07	0.004739	1	1	4	Q92698
High	DRPSVNSIL		0.00165	0.004739	1	1	1	Q96PY6
High	EKLPVHVVV		5.14E-06	0.004739	1	1	72	Q92698

High	ELFILDEASL		1.5E-06	0.004739	1	1	222	Q92698
High	ELTSIVNR		0.001595	0.004739	1	1	10	Q92698
High	EVAVLANM		0.142523	0.004739	1	1	3	Q96PY6
High	FNSPSSPDF		2.17E-06	0.004739	1	1	133	Q92698
High	GRDAAASE		2.46E-06	0.004739	1	1	17	Q92698
High	HFELPILK		3.79E-05	0.004739	1	1	176	Q92698
High	HFELPILKG		0.004021	0.004739	1	1	4	Q92698
High	HFSLGELK		4.79E-05	0.004739	1	1	207	Q92698
High	HFSLGELKE		4.38E-08	0.004739	1	1	139	Q92698
High	HPNIVQYR		0.004413	0.004739	1	1	2	Q96PY6
High	KHFELPILK		0.001088	0.004739	1	1	120	Q92698
High	LDGTMSIK		0.000439	0.004739	1	1	37	Q92698
High	LDGTMSIK		0.002048	0.004739	1	1	9	Q92698
High	LDGTMSIK		0.038011	0.004739	1	1	1	Q92698
High	LDKEKLPV		2.99E-08	0.004739	1	1	177	Q92698
High	LEGFMNQR		0.001825	0.004739	1	1	35	Q92698
High	IGEGSFGK		0.003307	0.004739	1	1	1	Q96PY6
High	LHVGVLQK		0.0009	0.004739	1	1	88	Q92698
High	LKNSENQT		7.35E-06	0.004739	1	1	36	Q92698
High	LKNSENQT		0.000206	0.004739	1	1	11	Q92698
High	LLSAGTIEEK		0.000128	0.004739	1	1	151	Q92698
High	LLSAGTIEEK		6.68E-05	0.004739	1	1	3	Q92698
High	LPVHVVVD		6.57E-06	0.004739	1	1	167	Q92698
High	LPVHVVVD		0.000305	0.004739	1	1	4	Q92698
High	IQPLAIDGG		0.001598	0.004739	1	1	8	Q92698
High	IQPLAIDGG		8.82E-06	0.004739	1	1	89	Q92698
High	IQPLAIDGG		1.11E-07	0.004739	1	1	165	Q92698
High	LRELTSIVN		0.000748	0.004739	1	1	183	Q92698
High	LTPLQTELY		0.000356	0.004739	1	1	40	Q92698
High	LTPLQTELY		2.78E-05	0.004739	1	1	94	Q92698
High	LTPLQTELY		0.219871	0.004739	1	1	1	Q92698
High	LVMFDPDV		5.77E-06	0.004739	1	1	114	Q92698
High	IVQNILGNE		0.024054	0.004739	1	1	2	Q96PY6
High	MLVLDYILA		5.63E-06	0.004739	1	1	98	Q92698
High	MSVSSLSSI		0.000209	0.004739	1	1	19	Q92698
High	MSVSSLSSI		7.73E-06	0.004739	1	1	119	Q92698
High	NSENQTYQ		7.47E-07	0.004739	1	1	47	Q92698
High	NSENQTYQ		0.000207	0.004739	1	1	13	Q92698
High	NWYNEVGK		0.005236	0.004739	1	1	88	Q92698
High	NWYNEVGK		0.000543	0.004739	1	1	5	Q92698
High	QAKPAEELL		5.04E-05	0.004739	1	1	119	Q92698
High	QAKPAEELL		0.00054	0.004739	1	1	1	Q92698
High	QAKPAEELL		0.001735	0.004739	1	1	2	Q92698
High	RALHDPLEK		0.006919	0.004739	1	1	4	Q92698
High	RALHDPLEK		9.54E-09	0.004739	1	1	236	Q92698
High	RSLAPSQLA		9.69E-05	0.004739	1	1	10	Q92698
High	RSLAPSQLA		0.121563	0.004739	1	1	1	Q92698
High	RVLISGTPI		0.001506	0.004739	1	1	32	Q92698

High	RYLYVR		0.005017	0.004739	1	1	82	Q92698
High	SLAPSQLAK		0.002679	0.004739	1	1	38	Q92698
High	SLAPSQLAK		0.013897	0.004739	1	1	3	Q92698
High	SILSKPFK		0.001511	0.004739	1	1	30	Q92698
High	SILSKPFKVF		1.49E-07	0.004739	1	1	129	Q92698
High	SSDKVVLVS		1.28E-06	0.004739	1	1	265	Q92698
High	TSDILSK		0.000624	0.004739	1	1	11	Q92698
High	TSDILSKYLF		0.001429	0.004739	1	1	4	Q92698
High	VLISGTPIQN		9.12E-09	0.004739	1	1	38	Q92698
High	VLISGTPIQN		1.38E-08	0.004739	1	1	70	Q92698
High	VPIPNYQGF		4.87E-05	0.004739	1	1	70	Q92698
High	VSSPILIISYE		4.33E-06	0.004739	1	1	209	Q92698
High	VSSPILIISYE		0.023489	0.004739	1	1	46	Q92698
High	VVERFNSPS		0.028862	0.004739	1	1	1	Q92698
High	VVLVSNYTC		4.09E-06	0.004739	1	1	179	Q92698
High	WGLRDEVL		0.009857	0.004739	1	1	1	Q92698

# Missed Cl	Theo. MH+	Confidence	Percolator	Percolator	XCorr (by Search Engine): Sequest HT
0	2251.975	High	0.000146	1.1E-12	2.59
0	1063.502	High	0.000146	3.86E-10	2.22
1	2960.504	High	0.000146	0.000185	0.78
2	3868.856	High	0.000146	6.31E-16	4.02
1	3740.761	High	0.000146	6.31E-16	4.44
0	2807.155	High	0.000146	7.31E-15	6.22
0	2727.189	High	0.000146	6.31E-16	7.75
1	1663.866	High	0.000146	5.17E-11	2.79
0	1486.774	High	0.000146	0.000872	0.64
0	1089.44	High	0.000146	5.74E-06	1.33
0	1140.555	High	0.000146	7.07E-10	2.32
1	1280.672	High	0.000146	0.000322	1.97
0	1940.864	High	0.000146	1.7E-07	2.53
0	994.4969	High	0.000146	2.78E-06	1.89
1	1150.598	High	0.000146	2.83E-07	3.12
0	999.5275	High	0.000146	5.15E-09	1.93
0	1477.72	High	0.000146	5.61E-06	2.71
1	2378.263	High	0.000146	6.78E-11	4.92
0	944.4159	High	0.000146	6.5E-07	2.09
1	2262.04	High	0.000146	5.83E-08	3.36
2	2418.141	High	0.000146	8.77E-07	4.91
1	1547.775	High	0.000146	3.53E-08	2.31
0	2020.861	High	0.000146	1.45E-11	4.73
1	2176.962	High	0.000146	3.37E-11	5.02
0	2235.98	High	0.000146	2.2E-13	4.2
0	880.4445	High	0.000146	7.02E-09	1.84
1	1008.539	High	0.000146	3.62E-09	2.37
0	1010.472	High	0.000146	5.83E-08	2.12
2	2818.404	High	0.000146	7.81E-16	4.53
0	1422.803	High	0.000146	4.82E-15	2.59
0	1355.709	High	0.000146	1.9E-12	3.38
1	1483.804	High	0.000146	1.84E-15	4.28
2	3065.655	High	0.000146	0.00013	1.63
0	1805.841	High	0.000146	1.06E-11	3.26
1	1738.806	High	0.000146	4.96E-08	3.2
1	2346.31	High	0.000146	1.24E-05	0.87
1	1610.8	High	0.000146	9.81E-10	2.93
1	1960.904	High	0.000146	6.31E-16	5.95
1	1944.909	High	0.000146	6.31E-16	5.9
0	1398.679	High	0.000146	4.5E-11	3.78
1	1639.858	High	0.000146	1.83E-10	2.43
1	1928.914	High	0.000146	6.31E-16	5.66
0	1804.803	High	0.000146	2.05E-09	2.87
0	1788.808	High	0.000146	1.52E-14	3.88
0	1772.813	High	0.000146	6.31E-16	5.49
1	1863.875	High	0.000146	1.23E-13	5.14
0	1815.747	High	0.000146	1.72E-08	3.29

0	1735.78	High	0.000146	1.88E-14	3.73
1	2647.267	High	0.000146	1.15E-11	3.56
0	1197.572	High	0.000146	4.24E-12	1.96
1	1353.673	High	0.000146	1.09E-05	1.5
0	1330.631	High	0.000146	9.08E-12	2.86
1	2397.269	High	0.000146	3.04E-12	2.78
0	1664.886	High	0.000146	6.31E-12	2.56
0	1272.648	High	0.000146	2.52E-13	3.46
1	3473.615	High	0.000146	4.5E-07	1.76
0	1648.891	High	0.000146	4.04E-15	2.95
1	1471.773	High	0.000146	2.88E-14	4.49
1	1003.503	High	0.000146	1.08E-09	2.42
0	1854.736	High	0.000146	2.13E-06	3.13
0	1343.678	High	0.000146	1.61E-12	3.91
0	1774.77	High	0.000146	1.73E-15	4.38
0	875.408	High	0.000146	3.14E-08	1.09
1	3059.471	High	0.000146	2.02E-08	1.18
1	3043.476	High	0.000146	1.55E-15	3.72
0	2416.101	High	0.000146	3.4E-15	4.99
1	2507.138	High	0.000146	1.96E-09	4.92
1	2427.172	High	0.000146	6.2E-11	3.93
1	2298.216	High	0.000146	8.44E-11	3.95
2	2792.469	High	0.000146	1.53E-14	5.08
0	2826.268	High	0.000146	0.00025	1.22
0	2746.302	High	0.000146	6.31E-16	5.13
0	2666.336	High	0.000146	6.31E-16	7.06
1	2123.947	High	0.000146	1.17E-10	4.61
1	2043.98	High	0.000146	1.27E-15	5.02
0	1995.852	High	0.000146	2.26E-12	4.04
0	1915.885	High	0.000146	7.81E-15	3.28
0	942.5255	High	0.000146	1.06E-07	1.93
1	2330.315	High	0.000146	3.26E-11	2.8
2	2573.448	High	0.000146	1.05E-09	2.82
0	922.4993	High	0.000146	6.6E-08	2.47
1	2909.541	High	0.000146	6.31E-16	7.56
2	3265.747	High	0.000146	6.31E-16	7
0	1085.656	High	0.000146	5.78E-12	2.36
1	2076.112	High	0.000146	5.45E-13	4.28
0	905.3959	High	0.000146	4.05E-09	1.99
1	1617.746	High	0.000146	1.16E-11	3.81
2	1886.931	High	0.000146	6E-06	2.18
0	2006.059	High	0.000146	2.8E-12	3.17
1	2362.265	High	0.000146	6.31E-16	3.91
2	2619.403	High	0.000146	6.31E-16	5.69
1	1722.812	High	0.000146	3.35E-12	1.94
0	2476.226	High	0.000146	6.31E-16	5.07
0	1257.68	High	0.000146	7.87E-09	2.52
1	1673	High	0.000146	8.58E-15	3.32

0	1860.897	High	0.000146	6.31E-16	4.19
0	931.5207	High	0.000146	7.25E-09	1.99
0	974.5339	High	0.000146	0.000425	1.23
0	1789.847	High	0.000146	1.1E-15	2.37
2	1830.869	High	0.000146	1.49E-15	3.79
0	996.5877	High	0.000146	9.92E-13	2.6
1	1209.71	High	0.000146	6.58E-08	2.99
0	930.5043	High	0.000146	1.74E-12	2.6
1	2772.384	High	0.000146	6.31E-16	5.33
0	1026.548	High	0.000146	8.22E-08	1.74
1	1124.683	High	0.000146	2.92E-09	2.18
0	864.4495	High	0.000146	3.37E-10	2.05
1	992.5445	High	0.000146	1.32E-08	1.66
2	1148.646	High	0.000146	1.49E-05	1.31
2	2029.206	High	0.000146	6.31E-16	7.2
0	994.4775	High	0.000146	9.99E-09	2.04
0	794.4043	High	0.000146	4.12E-08	1.3
0	893.5567	High	0.000146	1.86E-09	2.08
1	2182.073	High	0.000146	2.01E-14	5.76
2	2338.175	High	0.000146	5.57E-11	4.39
0	1060.588	High	0.000146	1.79E-11	1.45
1	1604.901	High	0.000146	3.82E-12	3.8
0	1415.862	High	0.000146	1.54E-14	4.2
1	2302.387	High	0.000146	1.41E-10	2.19
0	1098.615	High	0.000146	7.29E-09	2.17
1	1826.949	High	0.000146	3.09E-14	3.5
2	2802.409	High	0.000146	6.31E-16	5.64
1	1200.706	High	0.000146	1.19E-09	3.32
0	1205.678	High	0.000146	2.05E-10	1.75
1	1361.779	High	0.000146	4.74E-13	3.24
2	1778.032	High	0.000146	0.001359	1.3
0	2219.985	High	0.000146	1.13E-14	3.4
0	1877.003	High	0.000146	4.86E-06	1.35
0	1406.808	High	0.000146	1.07E-14	2.84
0	1339.714	High	0.000146	5.74E-11	2.95
1	1467.809	High	0.000146	2.26E-14	4.31
0	1940.894	High	0.000146	6.31E-16	3.55
1	2096.996	High	0.000146	5.61E-11	4.6
0	1009.474	High	0.000146	1.24E-07	1.75
1	1578.781	High	0.000146	5.6E-10	2.26
0	1312.711	High	0.000146	1.96E-12	3.42
1	2633.407	High	0.000146	5.51E-10	3.62
2	2761.502	High	0.000146	8.85E-09	3.66
1	1078.6	High	0.000146	2.41E-07	2.31
2	3065.642	High	0.000146	6.31E-16	5.9
1	1070.632	High	0.000146	9.28E-12	2.74
2	1226.733	High	0.000146	0.000281	1.13
1	3915.08	High	0.000146	6.32E-09	1.88

1	869.4992	High	0.000146	1.12E-07	1.39
0	914.5306	High	0.000146	2.5E-08	1.7
2	1637.945	High	0.000146	1.28E-06	2.07
0	919.5611	High	0.000146	6.38E-09	2.21
1	2298.297	High	0.000146	6.31E-16	4.59
1	2286.186	High	0.000146	6.31E-16	3.75
0	763.4196	High	0.000146	7.78E-10	1.89
1	1363.783	High	0.000146	5.6E-09	3.34
0	3758.979	High	0.000146	6.31E-16	7.77
1	3887.074	High	0.000146	6.31E-16	7.48
0	1397.754	High	0.000146	1.8E-12	2.17
0	1624.894	High	0.000146	5.7E-15	2.46
1	2499.433	High	0.000146	4.57E-06	0.98
1	2273.127	High	0.000146	7.56E-06	3.16
0	1869	High	0.000146	4.99E-15	4.39
1	2988.512	High	0.000146	5.62E-07	1.46