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Data Article

# Dataset of proteins mapped on HepG2 cells and those differentially abundant after expression of the dengue non-structural 1 protein



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# ABSTRACT

The data supplied in this article are related to the research article entitled "The effect of the dengue non-structural 1 protein expression over the HepG2 cell proteins in a proteomic approach" (K. Rabelo, M.R. Trugillo, S.M. Costa, B.A. Pereira, O.C. Moreira, A.T. Ferreira et al., 2016) [1]. The present article provides the inventory of peptides and proteins mapped in a hepatocyte cell line (HepG2) by mass spectrometry in the presence of the non-structural protein 1 (NS1) of Dengue 2 virus (DENV2). Cells were transfected with pcENS1 plasmid, which encodes the DENV2 NS1 protein, or the controls pcDNA3 (negative control) or pMAXGFP, encoding the

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green fluorescent protein (GFP), a protein unrelated to dengue. Differentially abundant protein lists were obtained by comparing cells transfected with pcENS1 and controls. © 2016 The Authors. Published by Elsevier Inc. This is an open

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## **Specifications Table**

Subject area More specific sub- ject area	Biology Proteomics, Virology
Type of data	Table
How data was acquired	Mass spectrometry
Data format	Raw and analyzed
Experimental factors	HepG2 cells were transfected with plasmids expressing different proteins, lysed, trypsinized and submitted to Orbitrap
Experimental features	All samples were analyzed LTQ-Orbitrap XL mass spectrometer
Data source location	Oswaldo Cruz Foundation, Brazil
Data accessibility	Within this article

# Value of the data

- These data describe the use of quantitative mass spectrometry-based proteomic experiments to assess the biological significance of cell alterations caused by DENV NS1 protein.
- 4756 proteins were mapped and we identify 41 or 81 differentially abundant proteins in the presence of NS1, comparing to controls.
- The data open new perspectives to identify the molecular mechanisms involving DENV NS1 protein in infected cells.

## 1. Data

HepG2 cells were transfected with the plasmids: pcENS1, pcDNA3 and pMAXGFP. To produce accurate data, we used three independent experimental biological replicates and samples were submitted to LTQ-Orbitrap XL (Thermo Scientific). Data analysis, using the PatternLab for Proteomics software, identified 14,138 peptides which mapped to 4756 proteins, from all conditions (HepG2 transfected with the three different plasmids and non-transfected cells) (Supplementary Table S1a–h). Applying the maximum parsimony principle we found 2314 proteins (Supplementary Table S1g). Using the Tfold module we generate the differential abundance distribution when comparing: non-transfected HepG2 x cell transfected with pcDNA3 (Table 1); HepG2 transfected with pcDNA3 x pcENS1 (Table 2) and cells transfected with pMAGFP x pcENS1 (Table 3)[1].

### 2. Experimental design, materials and methods

### 2.1. Cell culture

HepG2 cells (ATCC) were cultivated in Dulbecco's modified Eagle's medium (DMEM) (SIGMA) supplemented with 10% fetal bovine serum (FBS) (Invitrogen). Cells were maintained at 37° C and

Table 1

List of 54 differentially abundant proteins for statistics between HepG2 x pcDNA3.

Locus	Fold Change	pValue	Signal + (pcDNA3)	Signal- (HepG2)	Description
spIP206741 COX5A HUMAN	7.52	0.04600	1.29E-03	1.72E-04	Cytochrome c oxidase subunit 5A, mitochondrial OS=Homo sapiens $GN=COX5A PE=1 SV=2$
spiP39656i OST48_HUMAN	6.45	0.00016	9.18E-04	1.42E-04	Dolichyl-diphosphooligosaccharide-protein glycosyltransferase 48 kDa subunit $OS=Homo$ sapiens $GN=DDOST$ $PE=1$ $SV=4$
spiQ71UI9iH2AV_HUMAN	4.20	0.02071	5.62E-03	1.34E-03	Histone H2A.V OS=Homo sapiens $GN=H2AFV PE=1 SV=3$
spIP0C0S5IH2AZ_HUMAN	4.18	0.02251	5.58E-03	1.34E-03	Histone H2A.Z OS=Homo sapiens $GN=H2AFZ PE=1 SV=2$
spiP02768iALBU_HUMAN	3.40	0.01591	6.34E-04	1.87E-04	Serum albumin $OS=Homo$ sapiens $GN=ALB$ $PE=1$ $SV=2$
tr B4DLR8  B4DLR8_HUMAN	3.32	0.01500	2.00E-03	6.04E-04	NAD(P)H dehydrogenase [quinone] 1 OS=Homo sapiens GN=NQO1 PE=1 SV=1
spIP15559INQO1_HUMAN	3.27	0.01639	1.45E-03	4.45E-04	NAD(P)H dehydrogenase [quinone] 1 OS=Homo sapiens GN=NQO1 PE=1 SV=1: trlH3BNV2 H3BNV2_HU- MAN NAD(P)H dehydrogenase [quinone] 1 OS=Homo sapiens GN=NQO1 PE=1 SV=1
spIP13073I COX41_HUMAN	3.16	0.04534	5.81E-04	1.84E-04	Cytochrome <i>c</i> oxidase subunit 4 isoform 1, mitochondrial OS=Homo sapiens $GN=COX411 PE=1 SV=1$ : trl H3BPG0 H3BPG0_HUMAN Cytochrome <i>c</i> oxidase subunit 4 isoform 1, mitochondrial (Fragment) OS=Homo sapiens $GN=COX411 PE=1 SV=1$
spiQ9UBX3IDIC_HUMAN	3.14	0.02829	6.89E-04	2.19E-04	Mitochondrial dicarboxylate carrier OS=Homo sapiens $GN=SLC25A10 PE=1 SV=2$
triG3V576i G3V576_HUMAN	3.13	0.01342	1.92E-03	6.15E-04	Heterogeneous nuclear ribonucleoproteins C1/C2 OS=Homo sapiens GN=HNRNPC PE=1 SV=1: tr G3V575  G3V575_HUMAN Heterogeneous nuclear ribonucleoproteins C1/C2 (Fragment) OS=Homo sapiens GN=HNRNPC PE=1 SV=1: tr G3V555 G3V555_HUMAN Heterogeneous nuclear ribonucleoproteins C1/C2 (Fragment) OS=Homo sapiens GN=HNRNPC PE=1 SV=1
spiP07910i HNRPC_HUMAN	3.13	0.01342	1.45E-03	4.64E-04	Heterogeneous nuclear ribonucleoproteins C1/C2 OS=Homo sapiens GN=HNRNPC PE=1 SV=4
triB4DY08  B4DY08_HUMAN	3.13	0.01342	1.54E-03	4.93E-04	Heterogeneous nuclear ribonucleoproteins C1/C2 OS=Homo sapiens GN=HNRNPC PE=1 SV=1: trlG3V4W0  G3V4W0_HUMAN Heterogeneous nuclear ribonucleoproteins C1/C2 (Fragment) OS=Homo sapiens GN=HNRNPC PE=1 SV=1: trlG3V251_G3V251_HUMAN Heterogeneous nuclear ribonucleoproteins C1/C2 (Fragment) OS=Homo sapiens GN=HNRNPC PE=1 SV=1: trlG3V3K6G3V3K6_HUMAN Heterogeneous nuclear ribonucleoproteins C1/C2 (Fragment) OS=Homo sapiens GN=HNRNPC PE=1 SV=1: trlG3V5X6I G3V5X6_HUMAN Heterogeneous nuclear ribonucleoproteins C1/C2 (Fragment) OS=Homo sapiens GN=HNRNPC PE=1 SV=1: trlG3V4M8/G3V4M8_HUMAN Heterogeneous nuclear ribonucleoproteins C1/C2 (Fragment) OS=Homo sapiens GN=HNRNPC PE=1 SV=1: trlG3V2H6_HUMAN Heterogeneous nuclear ribonucleoproteins C1/C2 (Fragment) OS=Homo sapiens GN=HNRNPC PE=1 SV=1: trlG3V2H6_HUMAN Heterogeneous nuclear ribonucleoproteins C1/C2 (Fragment) OS=Homo sapiens GN=HNRNPC PE=1 SV=1: trlG3V2H6_HUMAN Heterogeneous
tr G3V2Q1  G3V2Q1_HUMAN	3.13	0.01342	1.46E-03	4.66E-04	Heterogeneous nuclear ribonucleoproteins C1/C2 OS=Homo sapiens GN=HNRNPC $PE=1$ SV=1
tr G3V4C1  G3V4C1_HUMAN	3.13	0.01342	1.52E-03	4.87E-04	Heterogeneous nuclear ribonucleoproteins C1/C2 OS=Homo sapiens GN=HNRNPC $PE=1$ SV=1
spiO151731 PGRC2_HUMAN	3.11	0.00180	7.94E-04	2.55E-04	Membrane-associated progesterone receptor component 2 OS=Homo sapiens $GN=PGRMC2$ PE=1 SV=1

spiP05787iK2C8_HUMAN	3.02	0.01302	8.22E-03	2.72E-03	Keratin, type II cytoskeletal 8 OS=Homo sapiens GN=KRT8 PE=1 SV=7: trlF8VUG2lF8VUG2_HUMAN Keratin, type II cytoskeletal 8 (Fragment) OS=Homo sapiens GN=KRT8 PE=1 SV=1: trlF8VP67/F8VP67_HUMAN Keratin, type II cytoskeletal 8 (Fragment) OS=Homo sapiens GN=KRT8 PE=1 SV=1: trlF8VRG4I F8VRG4 HUMAN Keratin, type II cytoskeletal 8 (Fragment) OS=Homo sapiens GN=KRT8 PE=1 SV=1
spiP61353iRL27_HUMAN	2.98	0.03444	9.43E-04	3.17E-04	60S ribosomal protein L27 OS=Homo sapiens GN=RPL27 PE=1 SV=2: trlK7EQQ9lK7EQQ9_HUMAN 60S ribosomal protein L27 OS=Homo sapiens GN=RPL27 PE=1 SV=1
triK7ELC7i K7ELC7_HUMAN	2.98	0.03444	8.91E-04	2.99E-04	60S ribosomal protein L27 (Fragment) OS=Homo sapiens GN=RPL27 PE=1 SV=1
triE9PCY7  E9PCY7_HUMAN	2.81	0.02643	1.33E-03	4.74E-04	Heterogeneous nuclear ribonucleoprotein H OS=Homo sapiens GN=HNRNPH1 PE=1 SV=1: trlH0YBG7  H0YBG7_HUMAN Heterogeneous nuclear ribonucleoprotein H (Fragment) OS=Homo sapiens GN=HNRNPH1 PE=1 SV=1: trlD6RBM0_HUMAN Heterogeneous nuclear ribonucleoprotein H (Fragment) OS=Homo sapiens GN=HNRNPH1 PE=1 SV=1: trlE5RGH4!E5RGH4_HUMAN Heterogeneous nuclear ribonucleoprotein H (Fragment) OS=Homo sapiens GN=HNRNPH1 PE=1 SV=1: trlD6RIU0]D6RIU0_HUMAN Heterogeneous nuclear ribonucleoprotein H (Fragment) OS=Homo sapiens GN=HNRNPH1 PE=1 SV=1: trlD6RFM3  D6RFM3_HUMAN Heterogeneous nuclear ribonucleoprotein H (Fragment) OS=Homo sapiens GN=HNRNPH1 PE=1 SV=1
triE7EMC6i E7EMC6_HUMAN	2.77	0.00863	5.78E-04	2.08E-04	Annexin OS=Homo sapiens GN=ANXA6 PE=1 SV=1
spiP37108ISRP14_HUMAN	2.71	0.01021	8.50E-04	3.14E-04	Signal recognition particle 14 kDa protein OS=Homo sapiens GN=SRP14 PE=1 SV=2: trlH0YLW0 H0YLW0_HUMAN Signal recognition particle 14 kDa protein OS=Homo sapiens GN=SRP14 PE=1 SV=1
spiQ53GQ0i DHB12_HUMAN	2.42	0.01385	6.34E-04	2.62E-04	Estradiol 17-beta-dehydrogenase 12 OS=Homo sapiens GN=HSD17B12 PE=1 SV=2
sdiP04040iCATA HUMAN	2.42	0.00728	7.06E-04	2.92E-04	Catalase $OS = Homo \text{ sadiens } GN = CAT PE = 1 SV = 3$
triF8VVM2I F8VVM2_HUMAN	2.41	0.00909	8.74E-04	3.62E-04	Phosphate carrier protein, mitochondrial OS=Homo sapiens $GN=SLC25A3$ PE=1 SV=1
tr H0YLA2  H0YLA2_HUMAN	2.30	0.00896	8.54E-04	3.71E-04	Signal recognition particle 14 kDa protein OS=Homo sapiens GN=SRP14 PE=1 SV=1
spiP22087iFBRL_HUMAN	2.21	0.01012	8.83E-04	4.00E-04	rRNA 2'-O-methyltransferase fibrillarin OS=Homo sapiens GN=FBL PE=1 SV=2: trlMOR299IMOR299_HUMAN rRNA 2'-O-methyltransferase fibrillarin (Fragment) OS=Homo sapiens GN=FBL PE=1 SV=1
triH3BNX8i H3BNX8_HUMAN	0.75	0.04600	1.27E-03	1.68E-04	Cytochrome c oxidase subunit 5A, mitochondrial OS=Homo sapiens GN=COX5A PE=1 SV=1
tr H0Y449  H0Y449_HUMAN	- 1.11	0.00212	2.27E-04	2.52E-03	Nuclease-sensitive element-binding protein 1 (Fragment) $OS=Homo$ sapiens $GN=YBX1$ $PE=1$ $SV=1$
tr H3BRN4  H3BRN4_HUMAN	- 1.70	0.00099	4.66E-04	7.91E-04	4-aminobutyrate aminotransferase, mitochondrial OS=Homo sapiens GN=ABAT PE=1 SV=1: spiP80404 GABT_HUMAN 4-aminobutyrate aminotransferase, mitochondrial OS=Homo sapiens GN=ABAT PE=1 SV=3
spIP16401IH15_HUMAN	-1.74	0.04345	2.18E-04	3.79E-03	Histone H1.5 OS=Homo sapiens $GN$ =HIST1H1B PE=1 SV=3
triH3BNQ7I H3BNQ7_HUMAN	-2.14	0.00576	3.69E-04	7.89E-04	4-aminobutyrate aminotransferase, mitochondrial OS=Homo sapiens $GN=ABAT PE=1 SV=1$
triA0A087WYT3I A0A087WYT3_HUMAN	-2.22	0.00267	4.43E-04	9.82E-04	Prostaglandin E synthase 3 OS=Homo sapiens $GN=PTGES3 PE=4 SV=1$
spiQ15185ITEBP_HUMAN	-2.37	0.00333	4.54E-04	1.08E-03	Prostaglandin E synthase 3 OS=Homo sapiens $GN=PTGES3 PE=1 SV=1$
spiO14979i HNRDL HUMAN	-2.55	0.01357	1.96E-04	5.01E-04	Heterogeneous nuclear ribonucleoprotein D-like OS=Homo sapiens GN=HNRNPDL PE=1 SV=3

251

Table 1 (continued)

Locus	Fold Change	pValue	Signal+ (pcDNA3)	Signal- (HepG2)	Description
triA0A087WUK2i A0A087WUK2_HUMAN	-2.55	0.01357	2.27E-04	5.79E-04	Heterogeneous nuclear ribonucleoprotein D-like OS=Homo sapiens GN=HNRNPDL PE=4 SV=1
spiP01009iA1AT_HUMAN	-2.74	0.02442	2.45E-04	6.73E-04	Alpha-1-antitrypsin OS=Homo sapiens GN=SERPINA1 PE=1 SV=3: trlG3V2B9lG3V2B9_HUMAN Short peptide from AAT (Fragment) OS=Homo sapiens GN=SERPINA1 PE=1 SV=1: trlG3V544[G3V544_HUMAN Short peptide from AAT (Fragment) OS=Homo sapiens GN=SERPINA1 PE=1 SV=1: trlG3V387lG3V387_HUMAN Short peptide from AAT (Fragment) OS=Homo sapiens GN=SERPINA1 PE=1 SV=1: trlG3V588lG3V588_HUMAN Short peptide from AAT (Fragment) OS=Homo sapiens GN=SERPINA1 PE=1 SV=1: trlG3V488lG3V588_HUMAN Short peptide from AAT (Fragment) OS=Homo sapiens GN=SERPINA1 PE=1 SV=1: trlG3V487lG3V487lG3V588_HUMAN Short peptide from AAT (Fragment) OS=Homo sapiens GN=SERPINA1 PE=1 SV=1: trlG3V487lG3V487lG3V487lG3V487lG3V488_HUMAN Short peptide from AAT (Fragment) OS=Homo sapiens GN=SERPINA1 PE=1 SV=1: trlG3V487lG3V487lG3V487lG3V488_HUMAN Short peptide from AAT (Fragment) OS=Homo sapiens GN=SERPINA1 PE=1 SV=1: trlG3V487lG3V487lG3V487lG3V487lG3V488_HUMAN Short peptide from AAT (Fragment) OS=Homo sapiens GN=SERPINA1 PE=1 SV=1: trlG3V487lG3V487lG3V488_HUMAN Short peptide from AAT (Fragment) OS=Homo sapiens GN=SERPINA1 PE=1 SV=1: trlG3V487lG3V488_HUMAN Short peptide from AAT (Fragment) OS=Homo sapiens GN=SERPINA1 PE=4 SV=1
spiP05455iLA_HUMAN	-2.88	0.02516	2.26E-04	6.52E-04	Lupus La protein OS=Homo sapiens GN=SSB PE=1 SV=2: trlE7ERC4lE7ERC4_HUMAN Lupus La protein (Fragment) OS=Homo sapiens GN=SSB PE=1 SV=1: trlE9PGX9lE9PGX9_HUMAN Lupus La protein (Fragment) OS=Homo sapiens GN=SSB PE=1 SV=1
spIP10599ITHIO_HUMAN	-3.01	0.04302	2.14E-04	6.44E-04	Thioredoxin $OS$ = Homo sapiens $GN$ = TXN $PE$ = 1 $SV$ = 3
triE7EMB3I E7EMB3_HUMAN	-3.13	0.04674	6.16E-04	1.93E-03	Calmodulin OS=Homo sapiens GN=CALM2 PE=1 SV=1
triH0Y7A7i H0Y7A7_HUMAN	- 3.13	0.04674	6.45E-04	2.02E-03	Calmodulin (Fragment) OS=Homo sapiens GN=CALM2 PE=1 SV=1
triE7ETZ0I E7ETZ0_HUMAN	- 3.13	0.04674	8.05E-04	2.52E-03	Calmodulin OS=Homo sapiens GN=CALM1 PE=1 SV=1
spIP62158ICALM_HUMAN	- 3.16	0.04206	8.10E-04	2.56E-03	Calmodulin OS=Homo sapiens GN=CALM1 PE=1 SV=2: trlQ96HY3lQ96HY3_HUMAN CALM1 protein OS=Homo sapiens GN=CALM3 PE=1 SV=1: trlG3V361lG3V361_HUMAN Calmodulin (Fragment) OS=Homo sapiens GN=CALM1 PE=1 SV=1
triJ3QQX2  J3QQX2_HUMAN	-3.54	0.00702	2.84E-04	1.00E-03	Rho GDP-dissociation inhibitor 1 OS=Homo sapiens GN=ARHGDIA PE=1 SV=1: trlJ3KTF8JJ3KTF8_HUMAN Rho GDP-dissociation inhibitor 1 (Fragment) OS=Homo sapiens GN=ARHGDIA PE=1 SV=3: trlJ3KS60J J3KS60_HUMAN Rho GDP-dissociation inhibitor 1 OS=Homo sapiens GN=ARHGDIA PE=1 SV=1
spiP52565i GDIR1_HUMAN	-3.59	0.00515	3.27E-04	1.17E-03	Rho GDP-dissociation inhibitor 1 OS=Homo sapiens GN=ARHGDIA PE=1 SV=3
spiP39687i AN32A_HUMAN	-3.84	0.01529	1.68E-04	6.45E-04	Acidic leucine-rich nuclear phosphoprotein 32 family member A OS=Homo sapiens GN=ANP32A PE=1 SV=1
spIP25787IPSA2_HUMAN	-4.00	0.02378	3.39E-04	1.36E-03	Proteasome subunit alpha type-2 OS=Homo sapiens $GN=PSMA2 PE=1 SV=2$
spiP17174iAATC_HUMAN	-4.13	0.01587	2.65E-04	1.09E-03	Aspartate aminotransferase, cytoplasmic $OS=Homo$ sapiens $GN=GOT1$ $PE=1$ $SV=3$
triK7ER90i K7ER90_HUMAN	-4.13	0.03401	1.33E-04	5.49E-04	Eukaryotic translation initiation factor 3 subunit G (Fragment) OS=Homo sapiens GN=EIF3G PE=1 SV=1
spiP06748INPM_HUMAN	-4.20	0.02188	8.93E-04	3.75E-03	Nucleophosmin OS=Homo sapiens GN=NPM1 PE=1 SV=2: trlE5RI98/E5RI98_HUMAN Nucleophosmin (Fragment) OS=Homo sapiens GN=NPM1 PE=1 SV=1
triJ3QLC8i J3QLC8_HUMAN	-4.24	0.01244	1.73E-04	7.34E-04	60S ribosomal protein L17 OS=Homo sapiens GN=RPL17 PE=1 SV=1
	-5.72	0.01114	6.04E-04	3.45E-03	Histidine triad nucleotide-binding protein 1 OS=Homo sapiens GN=HINT1 PE=1 SV=2

spIP497731 HINT1\_HUMAN splQ86SX6I -6.140.04372 2.79E-04 1.71E-03 Glutaredoxin-related protein 5, mitochondrial OS=Homo sapiens GN=GLRX5 PE=1 SV=2 GLRX5\_HUMAN spIQ15181IIPYR\_HUMAN -6.550.03416 8.78E-05 5.75E-04 Inorganic pyrophosphatase OS=Homo sapiens GN=PPA1 PE=1 SV=2 tr|H0YCY6| - 7.66 0.00032 7.86E-05 6.02E-04 FAD-AMP lyase (cyclizing) (Fragment) OS=Homo sapiens GN=DAK PE=1 SV=1 H0YCY6\_HUMAN

### Table 2

List of 41 differentially abundant proteins for statistics between pcDNA3 x pcENS1.

Locus	Fold Change	pValue	Signal + (pcENS1)	Signal- (pcDNA3)	Description
spiQ9UK22IFBX2_HUMAN triF2Z2V0I	2.887204251 2.448891458	0.00294 0.01885	0.000565646 0.000415618	0.000195915 0.000169717	F-box only protein 2 OS=Homo sapiens GN=FBXO2 PE=1 SV=2 Copine-1 (Fragment) OS=Homo sapiens GN=CPNE1 PE=1 SV=1
spiP05386iRLA1_HUMAN spiP43243i MATR3_HUMAN	2.20861757 2.106781667	0.01819 0.00112	0.002338141 0.000626087	0.001058645 0.000297177	60S acidic ribosomal protein P1 OS=Homo sapiens GN=RPLP1 PE=1 SV=1 Matrin-3 OS=Homo sapiens GN=MATR3 PE=1 SV=2: trlB3KM87lB3KM87_HUMAN Matrin-3 OS=Homo sapiens GN=MATR3 PE=1 SV=1
triA8MXP9 A8MXP9 HUMAN	2.106781667	0.00112	0.000592509	0.000281239	Matrin-3 OS=Homo sapiens GN=MATR3 $PE=1$ SV=1
spiP51149iRAB7A_HUMAN	1.938939554	0.00429	0.000768084	0.000396136	Ras-related protein Rab-7a OS=Homo sapiens GN=RAB7A PE=1 SV=1: trlC9J592lC9J592_HUMAN Ras-related protein Rab-7a (Fragment) OS=Homo sapiens GN=RAB7A PE=1 SV=1
spiQ9P0351 HACD3_HUMAN	1.842287267	0.00587	0.00044159	0.000239697	Very-long-chain (3R)-3-hydroxyacyl-CoA dehydratase 3 OS=Homo sapiens GN=PTPLAD1 PE=1 SV=2
spiP52926i HMGA2_HUMAN	-0.002597139	0.02916	0.000932861	0.002422769	High mobility group protein HMGI-C OS=Homo sapiens $GN$ =HMGA2 PE=1 SV=1
tr F5H2A4  F5H2A4_HUMAN	-0.002597139	0.02916	0.00086171	0.002237981	High mobility group protein HMGI-C OS=Homo sapiens $GN$ =HMGA2 PE=1 SV=1
tr F5H6H0  F5H6H0_HUMAN	-0.002597139	0.02916	0.000691713	0.001796475	High mobility group protein HMGI-C OS=Homo sapiens $GN=HMGA2 PE=1 SV=1$
triJ3QRW1  J3QRW1_HUMAN	-0.018702242	0.00638	0.000248201	0.000464192	26S protease regulatory subunit 8 (Fragment) OS=Homo sapiens $GN=PSMC5 PE=1 SV=1$
spiP00338iLDHA_HUMAN	- 0.267759962	0.03438	0.000722573	0.001934761	> L-lactate dehydrogenase A chain OS=Homo sapiens GN=LDHA PE=1 SV=2: trlF5GXY2  F5GXY2_HUMAN L-lactate dehydrogenase A chain (Fragment) OS=Homo sapiens GN=LDHA PE=1 SV=3
spiP02765iFETUA_HUMAN	-0.475079452	0.0123	0.000111731	0.00053081	Alpha-2-HS-glycoprotein OS=Homo sapiens GN=AHSG PE=1 SV=1
spIP08238IHS90B_HUMAN	- 1.523033786	0.0036	0.002236377	0.003406078	Heat shock protein HSP 90-beta OS=Homo sapiens $GN=HSP90AB1 PE=1 SV=4$
spiP02545iLMNA_HUMAN	- 1.560530275	0.00312	0.000879213	0.001372039	Prelamin-A/C OS=Homo sapiens GN=LMNA PE=1 SV=1
spiP39023iRL3_HUMAN	-2.033045505	0.00327	0.000226732	0.000460956	60S ribosomal protein L3 OS=Homo sapiens GN=RPL3 PE=1 SV=2: trlG5E9G0lG5E9G0_HUMAN 60S ribosomal protein L3 OS=Homo sapiens GN=RPL3 PE=1 SV=1: trlB5MCW2lB5MCW2_HU- MAN 60S ribosomal protein L3 (Fragment) OS=Homo sapiens GN=RPL3 PE=1 SV=1
spIP02768 ALBU_HUMAN	-2.302973671	0.00479	0.000275409	0.000634261	Serum albumin OS=Homo sapiens GN=ALB PE=1 SV=2
spiP26373iRL13_HUMAN	-2.352456304	0.0203	0.000324859	0.000764216	60S ribosomal protein L13 OS=Homo sapiens GN=RPL13 PE=1 SV=4: trlH3BUK8 H3BUK8_HU- MAN 60S ribosomal protein L13 (Fragment) OS=Homo sapiens GN=RPL13 PE=1 SV=1: trlJ3QSB4  J3QSB4_HUMAN 60S ribosomal protein L13 (Fragment) OS=Homo sapiens GN=RPL13 PE=1 SV=1
tr A0A087X1S2  A0A087X1S2_HUMAN	-2.401479196	0.00478	0.000370535	0.000889831	Nuclease-sensitive element-binding protein 1 OS=Homo sapiens GN=YBX1 PE=4 SV=1
tr A0A087WZH7  A0A087WZH7_HUMAN	-2.408424932	0.00411	0.000303789	0.000731652	Myristoylated alanine-rich C-kinase substrate OS=Homo sapiens GN=MARCKS $PE=4$ SV=1

triA0A087WWU8i A0A087WWU8 HUMAN	-2.440187743	0.00449	0.000253765	0.000619234	Tropomyosin alpha-3 chain OS=Homo sapiens $GN=TPM3 PE=4 SV=1$
spiP29966 MARCS HUMAN	-2.684863748	0.00191	0.00026381	0.000708295	Myristoylated alanine-rich C-kinase substrate $OS=Homo$ sapiens $GN=MARCKS$ $PE=1$ $SV=4$
splQ9GZT3ISLIRP_HUMAN	-2.730197953	0.00514	0.000172275	0.000470346	SRA stem-loop-interacting RNA-binding protein, mitochondrial OS=Homo sapiens GN=SLIRP $PE=1$ SV=1
triG3V2S9i G3V2S9_HUMAN	-2.730197953	0.00514	0.000151436	0.000413449	SRA stem-loop-interacting RNA-binding protein, mitochondrial OS=Homo sapiens GN=SLIRP $PE=1$ SV=1
tr H0YJ40 H0YJ40_HUMAN	-2.730197953	0.00514	0.000195604	0.000534038	SRA stem-loop-interacting RNA-binding protein, mitochondrial (Fragment) OS=Homo sapiens GN=SLIRP PE=1 SV=1
tr A0A087WUN7  A0A087WUN7_HUMAN	-2.730197953	0.00514	0.000204109	0.000557257	SRA stem-loop-interacting RNA-binding protein, mitochondrial OS=Homo sapiens GN=SLIRP $PE=4$ SV=1
tr G3V4X6  G3V4X6_HUMAN	-2.730197953	0.00514	0.000191612	0.00052314	SRA stem-loop-interacting RNA-binding protein, mitochondrial OS=Homo sapiens GN=SLIRP $PE=1$ SV=1
spIP53004IBIEA_HUMAN	-2.899141374	0.03897	0.000143578	0.000416253	Biliverdin reductase A OS=Homo sapiens GN=BLVRA PE=1 SV=2: triC9J1E1/C9J1E1_HUMAN Biliverdin reductase A (Fragment) OS=Homo sapiens GN=BLVRA PE=1 SV=1
tr H3BT36  H3BT36_HUMAN	-3.082880329	0.02293	0.000536524	0.001654038	Proteasome subunit alpha type-2 OS=Homo sapiens $GN=PSMA2 PE=4 SV=1$
spiP62269iRS18_HUMAN	- 3.304625932	0.02144	0.000280286	0.000926242	40S ribosomal protein S18 OS=Homo sapiens GN=RPS18 PE=1 SV=3: triJ3JS69JJ3JS69_HUMAN 40S ribosomal protein S18 OS=Homo sapiens GN=RPS18 PE=1 SV=1
tr B7WNR0  B7WNR0_HUMAN	-3.482472238	0.0473	0.000142247	0.00049537	Serum albumin OS=Homo sapiens GN=ALB PE=1 SV=1
tr D6RHD5  D6RHD5_HUMAN	-3.482472238	0.0473	0.000153093	0.000533143	Serum albumin OS=Homo sapiens GN=ALB PE=1 SV=1
tr H0YA55  H0YA55_HUMAN	-3.482472238	0.0473	0.000154779	0.000539015	Serum albumin (Fragment) OS=Homo sapiens GN=ALB PE=1 SV=1
tr H7C367  H7C367_HUMAN	-3.818005689	0.03869	0.000206727	0.000789284	Non-POU domain-containing octamer-binding protein (Fragment) OS=Homo sapiens $GN$ =NONO PE=1 SV=3
tr A0A087X0X3  A0A087X0X3_HUMAN	- 3.895589738	0.04525	0.000131893	0.000513803	Heterogeneous nuclear ribonucleoprotein M OS=Homo sapiens GN=HNRNPM PE=4 SV=1
spiQ5VTE0IEF1A3_HUMAN	-4.17671178	0.00362	0.001520654	0.006351333	Putative elongation factor 1-alpha-like 3 OS=Homo sapiens GN=EEF1A1P5 PE=5 SV=1
	-4.228640752	0.01546	0.000141501	0.000598356	Profilin-2 US = Homo  sapiens  GN = PrN2 PE = 1 SV = 1
	-4.228040752	0.01546	0.00011706	0.000495003	Prolitin $OS = Homo Sapiens GN = PFN2 PE = 1 SV = 1$
O5TA01 HUMAN	-4.574529519	0.01927	0.000230100	0.001052855	Glutatinone S-transferase onlega-1 (riaginent) $OS = Horito Saplens GN = GSTOT PE = 1 SV = 1$
spiQ9H9B4i	-4.423436439	0.01596	0.000148649	0.000657539	Sideroflexin-1 OS=Homo sapiens GN=SFXN1 PE=1 SV=4: trlD6RFI0ID6RFI0_HUMAN Side-
SFXN1_HUMAN					roflexin-1 (Fragment) OS=Homo sapiens GN=SFXN1 PE=1 SV=3
spiP25786iPSA1_HUMAN	-9.640254654	0.00328	4.97959E-05	0.000480045	Proteasome subunit alpha type-1 OS=Homo sapiens GN=PSMA1 PE=1 SV=1: triF5GX11  F5GX11_HUMAN Proteasome subunit alpha type-1 OS=Homo sapiens GN=PSMA1 PE=1 SV=1

Table 3

List of 81 differentially abundant proteins for statistics between pMAXGFP x pcENS1.

Locus	Fold	pValue	Signal +	Signal-	Description
-	Change		(pcENS1)	(pMAXGFP)	•
triE9PIZ4iE9PIZ4_HUMAN	4.27	0.02884	5.1564100956E-	1.2067638747E-	Cysteine and histidine-rich domain-containing protein 1 OS=Homo sapiens GN=CHORDC1 PE=1
			04	04	SV=1
tr H0YMI6	2.63	0.04622	3.6248916762E-	1.3785223116E-	Proteasome subunit alpha type (Fragment) $OS=Homo$ sapiens $GN=PSMA4$ $PE=1$ $SV=1$
H0YMI6_HUMAN			04	04	
spiP60900iPSA6_HUMAN	2.51	0.01497	6.5820719273E-	2.6200153475E-	Proteasome subunit alpha type-6 OS=Homo sapiens $GN=PSMA6 PE=1 SV=1$
			04	04	
trlG3V295I	2.51	0.01497	7.9763039119E-	3.1749939680E-	Proteasome subunit alpha type $OS=Homo$ sapiens $GN=PSMA6$ $PE=1$ $SV=1$
G3V295_HUMAN			04	04	
tr G3V3I1	2.51	0.01497	1.0940470906E-	4.3548903750E-	Proteasome subunit alpha type $OS=Homo$ sapiens $GN=PSMA6$ $PE=1$ $SV=1$
G3V3I1_HUMAN			03	04	
triG3V3U4	2.51	0.01497	1.5132613964E-	6.0235866869E-	Proteasome subunit alpha type $OS=Homo$ sapiens $GN=PSMA6$ $PE=1$ $SV=1$
G3V3U4_HUMAN			03	04	
trlG3V5Z7I	2.51	0.01497	6.4253559290E-	2.5576340297E-	Proteasome subunit alpha type $OS=Homo$ sapiens $GN=PSMA6$ $PE=1$ $SV=1$
G3V5Z7_HUMAN			04	04	
spiP61626iLYSC_HUMAN	2.46	0.00250	8.0788085952E-	3.2813609299E-	Lysozyme C OS=Homo sapiens $GN=LYZ PE=1 SV=1$
			04	04	
spIP62081IRS7_HUMAN	2.07	0.00203	4.7837342096E-	2.3069960621E-	40S ribosomal protein S7 OS=Homo sapiens GN=RPS7 PE=1 SV=1
			04	04	
spi043175iSERA_HUMAN	2.05	0.02052	3.5415580623E-	1.7288143062E-	> D-3-phosphoglycerate dehydrogenase OS=Homo sapiens GN=PHGDH PE=1 SV=4
			04	04	
trIM0R210	1.88	0.03982	1.8803502747E-	1.0008461759E-	40S ribosomal protein S16 OS=Homo sapiens $GN=RPS16 PE=1 SV=1$
M0R210_HUMAN			03	03	
spIP62249IRS16_HUMAN	1.84	0.03469	1.6295554871E-	8.8430929241E-	40S ribosomal protein S16 OS=Homo sapiens GN=RPS16 PE=1 SV=2
			03	04	
triA0A087WZ27i	1.84	0.03469	1.6295554871E-	8.8430929241E-	Zinc finger protein 90 OS=Homo sapiens $GN=ZNF90 PE=4 SV=1$
A0A087WZ27_HUMAN			03	04	
spIP46782IRS5_HUMAN	1.82	0.04115	1.3188765458E-	7.2344215749E-	40S ribosomal protein S5 OS=Homo sapiens GN=RPS5 PE=1 SV=4: trlM0R0F0lM0R0F0_HUMAN
			03	04	40S ribosomal protein S5 (Fragment) OS=Homo sapiens $GN=RPS5$ $PE=1$ $SV=1$
spiP50395iGDIB_HUMAN	1.82	0.04377	7.3807276416E-	4.0635192954E-	Rab GDP dissociation inhibitor beta OS=Homo sapiens GN=GDI2 PE=1 SV=2: trlV9GYF8
			04	04	V9GYF8_HUMAN Rab GDP dissociation inhibitor beta (Fragment) OS=Homo sapiens GN=GDI2
					PE=1 SV=1
spIP62136IPP1A_HUMAN	1.81	0.04937	7.1055348515E-	3.9154015764E-	Serine/threonine-protein phosphatase PP1-alpha catalytic subunit OS=Homo sapiens
			04	04	GN = PPP1CA PE = 1 SV = 1
tr K7ERG4	1.79	0.03148	9.9775810350E-	5.5628189049E-	Small nuclear ribonucleoprotein Sm D2 OS=Homo sapiens $GN=SNRPD2 PE=1 SV=1$
K7ERG4_HUMAN			04	04	
trIM0R0R2I	1.79	0.04224	1.1751143601E-	6.5592088945E-	40S ribosomal protein S5 OS=Homo sapiens GN=RPS5 PE=1 SV=1
MOROR2_HUMAN			03	04	

spiP47755	1.76	0.00907	3.8503391537E-	2.1892360270E-	F-actin-capping protein subunit alpha-2 OS=Homo sapiens $GN=CAPZA2 PE=1 SV=3$
CAZA2_HUMAN			04	04	
tr A8MXQ1  A8MXQ1_HUMAN	1.58	0.00171	1.8271019599E- 03	1.1545048441E- 03	Pituitary tumor-transforming gene 1 protein-interacting protein $OS=Homo$ sapiens $GN=PTTG1IP$ PE=4 $SV=1$
spiP432431 MATR3 HUMAN	1.56	0.00335	6.2608664580E- 04	4.0083578369E- 04	Matrin-3 OS=Homo sapiens GN=MATR3 PE=1 SV=2: tr\B3KM87\B3KM87\HUMAN Matrin-3 OS=Homo sapiens GN=MATR3 PE=1 SV=1
triA8MXP9	1.56	0.00335	5.9250881452E-	3.7933844557E-	Matrin-3 OS=Homo sapiens GN=MATR3 PE=1 SV=1
triE9PSD5i	0.43	0.02884	4.9845297591E-	1.1665384122E-	Cysteine and histidine-rich domain-containing protein 1 OS=Homo sapiens GN=CHORDC1 PE=1
E9PSD5_HUMAN			04	04	SV=1
spIP18085IARF4_HUMAN	0.18	0.01873	1.0283317157E- 03	5.6312821438E- 04	ADP-ribosylation factor 4 OS=Homo sapiens GN=ARF4 PE=1 SV=3: trlC9JPM4lC9JPM4_HUMAN ADP-ribosylation factor 4 (Fragment) OS=Homo sapiens GN=ARF4 PE=1 SV=1
tr A0A087X1Z3  A0A087X1Z3 HUMAN	-0.20	0.00919	2.9662475890E- 04	5.9412009618E- 04	Proteasome activator complex subunit 2 OS=Homo sapiens GN=PSME2 PE=4 SV=1
triH0YM70I H0YM70_HUMAN	-0.20	0.00919	3.3045038930E- 04	6.6187063346E- 04	Proteasome activator complex subunit 2 OS=Homo sapiens GN=PSME2 PE=1 SV=1: trlHOYKU2  HOYKU2_HUMAN Proteasome activator complex subunit 2 (Fragment) OS=Homo sapiens GN=PSME2 PE=1 SV=1
tr H3BT71  H3BT71 HUMAN	-0.22	0.03725	3.6164761545E- 04	7.8828832196E- 04	RNA-binding motif protein, X chromosome, N-terminally processed OS=Homo sapiens GN=RBMX PE=1 SV=1
triH0YMF4 H0YMF4_HUMAN	-0.24	0.01281	4.5610249693E- 04	1.0891334963E- 03	60S ribosomal protein L28 OS=Homo sapiens GN=RPL28 PE=1 SV=1
spiQ92928i RAB1C_HUMAN	-0.33	0.00379	1.7941339393E- 04	5.9502701116E- 04	Putative Ras-related protein Rab-1C OS=Homo sapiens GN=RAB1C PE=5 SV=2
spiQ9H0U4i RAB1B HUMAN	-0.33	0.00379	1.7941339393E- 04	5.9502701116E- 04	Ras-related protein Rab-1B OS=Homo sapiens GN=RAB1B PE=1 SV=1
spiP08238i HS90B HUMAN	- 1.19	0.00037	2.2363769567E- 03	2.6541184171E- 03	Heat shock protein HSP 90-beta OS=Homo sapiens $GN=HSP90AB1 PE=1 SV=4$
spiP15531i NDKA HUMAN	- 1.38	0.00005	1.9216269100E- 03	2.6508648328E- 03	Nucleoside diphosphate kinase A OS=Homo sapiens GN=NME1 PE=1 SV=1
spiP22570iADRO_HUMAN	- 1.40	0.00127	4.9680105400E- 04	6.9510914113E- 04	NADPH:adrenodoxin oxidoreductase, mitochondrial OS=Homo sapiens GN=FDXR PE=1 SV=3
spiP02545i LMNA HUMAN	- 1.41	0.00262	8.7921299898E- 04	1.2356451265E- 03	Prelamin-A/C OS=Homo sapiens GN=LMNA PE=1 SV=1
spiQ8NBS9i	- 1.55	0.01047	4.3710044382E-	6.7551841613E-	Thioredoxin domain-containing protein 5 OS=Homo sapiens GN=TXNDC5 PE=1 SV=2
trik7ES89 K7ES89 HUMAN	- 1.57	0.00284	3.3391802783E- 04	5.2383721696E- 04	Dual-specificity protein phosphatase 3 (Fragment) OS=Homo sapiens GN=DUSP3 PE=1 SV=1
spiP60953i CDC42_HUMAN	- 1.59	0.01599	5.7952351298E- 04	9.2218098533E- 04	Cell division control protein 42 homolog OS=Homo sapiens GN=CDC42 PE=1 SV=2: trlQ5JYX0  Q5JYX0_HUMAN Cell division control protein 42 homolog (Fragment) OS=Homo sapiens GN=CDC42 PE=1 SV=1
triE3W990i E3W990 HUMAN	- 1.74	0.00295	2.0678393968E- 04	3.6029249723E- 04	Sequestosome-1 (Fragment) OS=Homo sapiens GN=SQSTM1 PE=1 SV=1
	- 1.79	0.01080	-	-	

257

Locus	Fold Change	pValue	Signal + (pcENS1)	Signal- (pMAXGFP)	Description
spiQ14847  LASP1_HUMAN			5.3129711808E- 04	9.5247256550E- 04	LIM and SH3 domain protein 1 OS=Homo sapiens GN=LASP1 PE=1 SV=2: trlC9J9W2 C9J9W2_HUMAN LIM and SH3 domain protein 1 (Fragment) OS=Homo sapiens GN=LASP1 PE=1 SV=1
tr G3V1V0  G3V1V0_HUMAN	- 1.80	0.01301	4.6887694121E- 04	8.4429729734E- 04	Myosin, light polypeptide 6, alkali, smooth muscle and non-muscle, isoform CRA_c OS=Homo sapiens GN=PDE6H PE=4 SV=1: trlF8W1R7IF8W1R7_HUMAN Retinal cone rhodopsin-sensitive cGMP 3',5'-cyclic phosphodiesterase subunit gamma OS=Homo sapiens GN=PDE6H PE=4 SV=1
spiP60660iMYL6_HUMAN	- 1.80	0.01301	4.9992839427E- 04	9.0021102564E- 04	Myosin light polypeptide 6 OS=Homo sapiens $GN=MYL6 PE=1 SV=2$
tr B7Z6Z4  B7Z6Z4_HUMAN	- 1.80	0.01301	3.1718146023E- 04	5.7114228937E- 04	Retinal cone rhodopsin-sensitive cGMP 3',5'-cyclic phosphodiesterase subunit gamma OS=Homo sapiens GN=PDE6H PE=2 SV=1
tr F8VPF3  F8VPF3 HUMAN	- 1.80	0.01301	5.8068605796E- 04	1.0456297298E- 03	Retinal cone rhodopsin-sensitive cGMP 3',5'-cyclic phosphodiesterase subunit gamma (Fragment) OS=Homo sapiens GN=PDE6H PE=4 SV=1
triG8JLA2i G8ILA2 HUMAN	- 1.80	0.01301	4.9663939168E- 04	8.9428858468E- 04	Retinal cone rhodopsin-sensitive cGMP 3',5'-cyclic phosphodiesterase subunit gamma OS=Homo saniens GN=PDF6H PE=4 SV=1
triJ3KND3 I3KND3 HUMAN	- 1.80	0.01301	4.9663939168E- 04	8.9428858468E- 04	Retinal cone rhodopsin-sensitive cGMP 3',5'-cyclic phosphodiesterase subunit gamma OS=Homo saniens CN=PDF6H PF-4 SV-1
spiP00338iLDHA_HUMAN	- 1.91	0.04007	7.2257279358E- 04	1.3809825096E- 03	> L-lactate dehydrogenase A chain OS=Homo sapiens GN=LDHA PE=1 SV=2: trlF5GXY2/ F5GXY2_HUMAN L-lactate dehydrogenase A chain (Fragment) OS=Homo sapiens GN=LDHA PE=1 SV=3
spi095292IVAPB_HUMAN	- 1.92	0.02424	2.7412268027E- 04	5.2623836772E- 04	Vesicle-associated membrane protein-associated protein B/C OS=Homo sapiens GN=VAPB PE=1 SV=3
tr H7C2l1  H7C2l1_HUMAN	- 1.99	0.01036	2.1368459062E- 04	4.2559598943E- 04	Protein arginine N-methyltransferase 1 OS=Homo sapiens GN=PRMT1 PE=1 SV=1: splQ99873 ANM1_HUMAN Protein arginine N-methyltransferase 1 OS=Homo sapiens GN=PRMT1 PE=1 SV=2: trlE9PKG1_E9PKG1_HUMAN Protein arginine N-methyltransferase 1 OS=Homo sapiens GN=PRMT1 PE=1 SV=1
spiQ9UL46i PSME2_HUMAN	-2.00	0.00919	3.1524137556E- 04	6.3140796832E- 04	Proteasome activator complex subunit 2 OS=Homo sapiens $GN=PSME2 PE=1 SV=4$
tr A0A087WZH7  A0A087WZH7 HUMAN	-2.05	0.01540	3.0378860599E- 04	6.2309294752E- 04	Myristoylated alanine-rich C-kinase substrate $OS=Homo$ sapiens $GN=MARCKS$ $PE=4$ $SV=1$
spiQ9NX63i MIC19 HUMAN	-2.06	0.04956	2.1744009773E- 04	4.4818711141E- 04	MICOS complex subunit MIC19 OS=Homo sapiens GN=CHCHD3 PE=1 SV=1
triC9JRZ6i C9IRZ6 HUMAN	-2.06	0.04956	2.1275388873E- 04	4.3852790643E- 04	MICOS complex subunit MIC19 OS=Homo sapiens GN=CHCHD3 PE=1 SV=1
spiQ92688i AN32B_HUMAN	-2.16	0.04445	2.5741410440E- 04	5.5640940806E- 04	Acidic leucine-rich nuclear phosphoprotein 32 family member B OS=Homo sapiens GN=ANP32B PE=1 SV=1: trlQ5T6W8lQ5T6W8_HUMAN Acidic leucine-rich nuclear phosphoprotein 32 family member B (Fragment) OS=Homo sapiens GN=ANP32B PE=1 SV=1
tr H0Y6E7  H0Y6E7_HUMAN	-2.18	0.03725	3.6660169237E- 04	7.9908679213E- 04	RNA-binding motif protein, X chromosome, N-terminally processed (Fragment) OS=Homo sapiens GN=RBMX PE=1 SV=2

tr C9J4S4 C9J4S4_HUMAN	-2.19	0.00627	4.4976713953E- 04	9.8605217655E- 04	Ras-related protein Rab-7a OS=Homo sapiens GN=RAB7A PE=1 SV=1
spi015143I ARC1B_HUMAN	-2.20	0.00773	1.6888176270E- 04	3.7175544430E- 04	Actin-related protein 2/3 complex subunit 1B OS=Homo sapiens GN=ARPC1B PE=1 SV=3: trlC9JFG9 C9JFG9_HUMAN Actin-related protein 2/3 complex subunit 1B (Fragment) OS=Homo sapiens GN=ARPC1B PE=1 SV=3: trlC9J6C8C9J6C8_HUMAN Actin-related protein 2/3 complex subunit 1B (Fragment) OS=Homo sapiens GN=ARPC1B PE=1 SV=3: trlC9JQM82HUMAN Actin-related protein 2/3 complex subunit 1B (Fragment) OS=Homo sapiens GN=ARPC1B PE=1 SV=3: trlC9JEY11 (C9JEY1_HUMAN Actin-related protein 2/3 complex subunit 1B (Fragment) OS=Homo sapiens GN=ARPC1B PE=1 SV=1: trlF8VXW2IF8VXW2_HUMAN Actin-related protein 2/3 complex subunit 1B OS=Homo sapiens GN=ARPC1B PE=1 SV=2: trlC9J4Z7lC9J4Z7_HUMAN Actin-related protein 2/3 complex subunit 1B (Fragment) OS=Homo sapiens GN=ARPC1B PE=1 SV=1: trlC9K0571 C9K057_HUMAN Actin-related protein 2/3 complex subunit 1B (Fragment) OS=Homo sapiens GN=ARPC1B PE=1 SV=1: trlC9JB7lC9JB7_HUMAN Actin-related protein 2/3 complex subunit 1B (Fragment) OS=Homo sapiens GN=ARPC1B PE=1 SV=1: trlC9K0571 C9K057_HUMAN Actin-related protein 2/3 complex subunit 1B (Fragment) OS=Homo sapiens GN=ARPC1B PE=1 SV=1: trlC9JB7lC9JB7_HUMAN Actin-related protein 2/3 complex subunit 1B (Fragment) OS=Homo sapiens GN=ARPC1B PE=1 SV=1: trlC9JT6_HUMAN Actin-related protein 2/3 complex subunit 1B (Fragment) OS=Homo sapiens SN=ARPC1B PE=1 SV=3
tr K7ELC7  K7ELC7 HUMAN	-2.22	0.03550	3.7971620398E- 04	8.4465213197E- 04	60S ribosomal protein L27 (Fragment) OS=Homo sapiens GN=RPL27 PE=1 SV=1
spIP29966I MARCS_HUMAN	-2.30	0.00963	2.6381044628E- 04	6.0621816617E- 04	Myristoylated alanine-rich C-kinase substrate $OS=Homo$ sapiens $GN=MARCKS$ $PE=1$ $SV=4$
tr H3BT36  H3BT36_HUMAN	-2.45	0.00347	5.3652366733E- 04	1.3139040782E- 03	Proteasome subunit alpha type-2 OS=Homo sapiens $GN=PSMA2 PE=4 SV=1$
spiP46779iRL28_HUMAN	-2.59	0.00655	3.4415463721E- 04	8.9038650796E- 04	60S ribosomal protein L28 OS=Homo sapiens GN=RPL28 PE=1 SV=3
tr H0YKD8  H0YKD8_HUMAN	-2.59	0.00655	2.7734814881E- 04	7.1754677406E- 04	60S ribosomal protein L28 OS=Homo sapiens GN=RPL28 PE=1 SV=1
tr H0YLP6  H0YLP6_HUMAN	-2.59	0.00655	5.2976612693E- 04	1.3705949617E- 03	60S ribosomal protein L28 OS=Homo sapiens GN=RPL28 PE=1 SV=1
spi000483i NDUA4_HUMAN	-2.74	0.03035	3.7754188433E- 04	1.0331761575E- 03	Cytochrome c oxidase subunit NDUFA4 OS=Homo sapiens GN=NDUFA4 PE=1 SV=1
spiQ15274i NADC_HUMAN	-2.75	0.00691	2.0572201432E- 04	5.6661625682E- 04	Nicotinate-nucleotide pyrophosphorylase [carboxylating] OS=Homo sapiens GN=QPRT PE=1 SV=3: trlC9JCJ5lC9JCJ5_HUMAN Uncharacterized protein (Fragment) OS=Homo sapiens PE=4 SV=5
tr H7BZ11  H7BZ11_HUMAN	-3.07	0.00710	2.2555444718E- 04	6.9288128636E- 04	Protein RPL36A-HNRNPH2 OS=Homo sapiens GN=RPL36A-HNRNPH2 PE=3 SV=2
trij3KQN4  J3KQN4_HUMAN	- 3.07	0.00710	1.8743256878E- 04	5.7577459007E- 04	60S ribosomal protein L36a OS=Homo sapiens GN=RPL36A PE=3 SV=1: splP83881 RL36A_HU- MAN 60S ribosomal protein L36a OS=Homo sapiens GN=RPL36A PE=1 SV=2: tr R4GN19  R4GN19_HUMAN 60S ribosomal protein L36a OS=Homo sapiens GN=RPL36A PE=4 SV=1
spiQ969Q0i RL36L HUMAN	-3.07	0.00710	2.5108891290E- 04	7.7132067726E- 04	60S ribosomal protein L36a-like OS=Homo sapiens GN=RPL36AL PE=1 SV=3
tr H0Y5B4  H0Y5B4 HUMAN	-3.07	0.00710	2.3763772114E- 04	7.2999992670E- 04	60S ribosomal protein L36a OS=Homo sapiens GN=RPL36A PE=3 SV=2
triD6RHD5i D6RHD5 HUMAN	- 3.10	0.01904	1.5309327240E- 04	4.7403637068E- 04	Serum albumin OS=Homo sapiens GN=ALB PE=1 SV=1
0	- 3.10	0.01904			Serum albumin OS=Homo sapiens GN=ALB PE=1 SV=1

Table 3 (continued)

Locus	Fold Change	pValue	Signal + (pcENS1)	Signal- (pMAXGFP)	Description
triB7WNR0  B7WNR0_HUMAN			1.4224658306E- 04	4.4045079786E- 04	
tr H0YA55  H0YA55_HUMAN	- 3.10	0.01904	1.5477932166E- 04	4.7925703555E- 04	Serum albumin (Fragment) OS=Homo sapiens $GN=ALB PE=1 SV=1$
spiQ9HAV7i GRPE1_HUMAN	-3.24	0.02180	1.2662540785E- 04	4.1001496166E- 04	GrpE protein homolog 1, mitochondrial OS=Homo sapiens $GN=GRPEL1 PE=1 SV=2$
spiP62269iRS18_HUMAN	-3.24	0.01564	2.8028634126E- 04	9.0884462029E- 04	40S ribosomal protein S18 OS=Homo sapiens GN=RPS18 PE=1 SV=3: trlJ3JS69JJ3JS69_HUMAN 40S ribosomal protein S18 OS=Homo sapiens GN=RPS18 PE=1 SV=1
triC9JQB3i C9IOB3 HUMAN	-3.26	0.01641	1.1866193463E- 04	3.8718514857E- 04	Ras-related protein Ral-B (Fragment) OS=Homo sapiens $GN=RALB PE=1 SV=1$
triE9PLD0 E9PLD0 HUMAN	-3.70	0.04512	1.5238542031E- 04	5.6381042429E- 04	Ras-related protein Rab-1B OS=Homo sapiens GN=RAB1B PE=3 SV=1
spiQ9UN86i G3BP2_HUMAN	-3.81	0.00228	1.1344218542E- 04	4.3192164871E- 04	Ras GTPase-activating protein-binding protein 2 OS=Homo sapiens $CN=G3BP2 PE=1 SV=2$ : trl DGRB17/DGRB17_HUMAN Ras GTPase-activating protein-binding protein 2 (Fragment) OS=Homo sapiens $CN=G3BP2 PE=1 SV=1$ : trlDGRAC7/DGRAC7_HUMAN Ras GTPase-activating protein-binding protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=1$ : trlDGRGJ4/ DGRGJ4_HUMAN Ras GTPase-activating protein-binding protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=1$ : trlDGRBW8/DGRBW8_HUMAN Ras GTPase-activating protein-binding protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=3$ : trlDGRBW8/DGRBW8_HUMAN Ras GTPase-activating protein-binding protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=3$ : trlDGRBND/DGRBR0_HUMAN Ras GTPase-activating protein-binding protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=3$ : trlDGRBND/DGRBR0_HUMAN Ras GTPase-activating protein-binding protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=3$ : trlDGRBM9/HUMAN Ras GTPase-activating protein-binding protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=1$ : trlDGRBM9/HUMAN Ras GTPase-activating protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=1$ : trlDGRBM9/HUMAN Ras GTPase-activating protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=3$ : trlDGR9A4/DGR9A4_HUMAN Ras GTPase-activating protein-binding protein-binding protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=3$ : trlDGR9A4/DGR9A4_HUMAN Ras GTPase-activating protein-binding protein-binding protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=3$ : trlDGR9A4/DGR9A4_HUMAN Ras GTPase-activating protein-binding Protein-binding protein-binding protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=3$ : trlDGR9A4/DGR9A4_HUMAN Ras GTPase-activating protein-binding Protein-binding Protein-binding protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=3$ : trlDGR9A4/DGR9A4_HUMAN Ras GTPase-activating protein-binding Protein-binding Protein 2 (Fragment) OS=Homo sapiens $GN=G3BP2 PE=1 SV=3$ : trlDGR9A5/HUMAN Ras GTPase-activating protein-binding Protein DS=HOMO Sapiens $GN=G3BP2 PE=1 SV=3$
tr M0QXU7  M0QXU7_HUMAN	- 3.82	0.03844	9.5943708277E- 05	3.6629063166E- 04	Mitochondrial import inner membrane translocase subunit TIM44 (Fragment) OS=Homo sapiens GN=TIMM44 PE=1 SV=1
spiQ9H9B4i SFXN1_HUMAN	-4.04	0.00033	1.4864899148E- 04	6.0096890241E- 04	Sideroflexin-1 OS=Homo sapiens GN=SFXN1 PE=1 SV=4: triD6RFI0ID6RFI0_HUMAN Side- roflexin-1 (Fragment) OS=Homo sapiens GN=SFXN1 PE=1 SV=3
trIJ3KRE2I J3KRE2_HUMAN	-4.63	0.02210	1.0372838583E- 04	4.8012684663E- 04	Rho GDP-dissociation inhibitor 1 OS=Homo sapiens $GN=ARHGDIA PE=1 SV=1$
spiP04179 SODM_HUMAN	- 5.14	0.00047	1.1732492605E- 04	6.0335127872E- 04	Superoxide dismutase [Mn], mitochondrial OS=Homo sapiens $GN=SOD2 PE=1 SV=2$

under humid atmosphere with 5% CO<sub>2</sub>. In all experiments, cells were transfected between 86 and 89 cell passages with 70–80% confluence.

#### 2.2. Plasmids

The recombinant plasmid pcENS1 was previously constructed in our laboratory [2], using the pcDNA3 mammalian expression vector (Invitrogen). It contains the sequence of 63 nucleotides that encodes 21 amino acids from the C-terminal portion of the DENV2 envelope (E) protein and the full length DENV2 *ns1* gene. The vector pcDNA3 was used as a negative control, while the plasmid pMAXGFP (Amaxa), which encodes the green fluorescent protein (GFP) from *Pontellina plumata* copepod, was used as a control for expression of a DENV non-related protein.

#### 2.3. Transfection

Transfection was performed by nucleofection with the Nucleofector V<sup>TM</sup> kit (Amaxa), according to manufacturer's recommendation. Briefly, HepG2 cells were seeded on 75 cm<sup>2</sup> bottles, harvested after 4 or 5 days with the aid of cell scrapes in 3 ml of CMF solution (8 g/L of NaCl; 0.4 g/L of KCl; 0.1 g/L of Na<sub>2</sub>SO<sub>4</sub>; 0.39 g/L of Na<sub>2</sub>HPO<sub>4</sub>.12H<sub>2</sub>O; 0.15 g/L of KH<sub>2</sub>PO<sub>4</sub>; 1.1 g/L of glucose; 0.0025 g/L of phenol red, pH 7.4), centrifuged at 500 g for 5 min and suspended in the nucleofection solution (Amaxa). Cell suspension with 5 µg of DNA plasmids (10<sup>6</sup> cells/100 µl/cuvettes) was submitted to an electric shock in the Nucleofector 6 equipment (Amaxa), using the T-28 program. Nine cuvettes were used for each sample (pcDNA3, pcENS1 or pMAXGFP). After shock, cells received 500 µL of DMEM with 10% FBS and were immediately transferred to microcentrifuge tubes containing another 500 µL DMEM with 10% FBS. Cells were seeded on 25 cm<sup>2</sup> flasks, incubated in humid atmosphere with 5% CO<sub>2</sub> at 37 °C for 24 h.

### 2.4. Proteomic sample preparation

Cells were centrifuged at 500 g for 10 min and suspended in 50 mM ammonium bicarbonate buffer containing 0.2% of RapiGest<sup>TM</sup> SF (Waters). The protein concentration was determined using Qubit 2.0<sup>®</sup> kit (Invitrogen) following the manufacturer's instructions. A total of 50 µg protein was used for each sample. Samples were treated with 5 µL of 100 mM dithiothreitol for reduction, incubated for 3 h at 37 °C. After reaching room temperature, samples were alkylated with 5 µL of 400 mM iodoacetamide for 15 min, in the dark. Trypsin (Promega) was added in the ratio 1:50 enzyme/substrate and digestion was performed for 20 h, at 37 °C. The reaction was stopped after adding formic acid to final concentration of 1%. Aliquots from this digestion were desalted by using POROS R2 C8–18 resin (Invitrogen), packaged in micropipette tips (Millipore) and equilibrated in TFA 1%. After washing with 0.1% TFA, peptides were eluted in 0.1% TFA with 70% acetonitrile and completely dried in the vacuum centrifuge.

### 2.5. Isoelectric focalization of peptides (OFFGEL)

Twenty five micrograms of peptides were solubilized in 1.8 mL of 0.01% ampholytes (OFFGEL buffer pH 3–10) containing 4% ( $\nu/\nu$ ) glycerol and was submitted to the 3100 OFFGEL Fractionator with the OFFGEL Low Res Kit pH 3–10 (Agilent Technologies) immobilized pH gradient (IPG) DryStrips, following the Agilent's instructions. The peptides were also separated according to the manufacturer's instruction and optimized as described in Hubner et al. [3]. Twelve well fractionations were focused for 20 kV with a maximum current of 50 mA and power of 200 mW for 24 h. Each fraction was separately desalted as previously described and suspended in 40  $\mu$ L of 1% formic acid. All fractions were analyzed on 10 cm reversed phase (RP) column coupled to an LTQ-Orbitrap XL mass spectrometer.

#### 2.6. Liquid chromatography-tandem mass spectrometry (LC-MS/MS) analysis

Desalted peptides fractions were loaded separately onto a 10 cm RP column coupled to the mass spectrometer by using a Proxeon easy-nLC-System (Thermo Scientific Easy-nLC II). Four microliters

were initially applied to a 2 cm long (100  $\mu$ m internal diameter) trap column packed with 5  $\mu$ m, 200  $\stackrel{\text{P}}{=}$ Magic C18 AQ matrix (Michrom Bioresources) followed by separation on a 10 cm long (75 µm internal diameter) separation column packed with the same matrix directly on a self-pack 5–15 µm Tip empty column (New Objective). Samples were loaded onto the trap column at 2 µL/min while chromatographic separation occurred at 200 nL/min. Mobile phase A consisted of 0.1% formic acid in water while mobile phase B consisted of 0.1% formic acid in acetonitrile. Peptides were eluted with a gradient of 2–40% of B over 32 min followed by up to 80% B in 4 min, maintaining at this concentration for 2 min more, before column equilibration. The HPLC system was coupled to the LTQ-Orbitrap XL via a nanoscale LC interface (Thermo Scientific). Source voltage was set to 1.9 kV, the temperature of heated capillary was set to 200 °C and tube lens voltage to 48 and 100 V, respectively. The target precursor specters were acquired in ion trap full scan MS with 60,000 while FWHM full AGC target was set to 500,000. MS1 spectra were acquired on the Orbitrap analyzer (300–1700 m/z) at a 60,000 resolution (for m/z 445.1200). For each spectrum, the 10 most intense ions were submitted to CID fragmentation (minimum signal required of 10,000; isolation width of 2.5; normalized collision energy of 35.0; activation Q of 0.25 and activation time of 30 s, followed by MS2 acquisition on the linear trap quadrupole analyzer. Dynamic exclusion option was enabled and set with the following values for each parameter: repeat count = 1; repeat duration = 30 s; exclusion list size = 500; exclusion duration = 45 s and exclusion mass width = 10 ppm. Data were acquired in technical triplicates using the Xcalibur software (version 2.0.7).

### 2.7. Protein identification

The raw data files were processed and quantified using PatternLab for Proteomics software v 3.2 [4] (available at: http://max.ioc.fiocruz.br/mtrugilho/RabeloK2016/). Peptide sequence matching (PSM) was performed using the Comet algorithm [5] against the UniProt database (http://www.uni prot.org/) with human proteins entries downloaded January 2015, plus a FASTA file containing Dengue virus and GFP sequences, retrieved from the NCBI database. A target-reverse strategy was employed for increased confidence in protein identifications [6]. The search considered tryptic and semi-tryptic peptide candidates. The cysteine carbamidomethylation and oxidation of methionine were considered as fixed and variable modifications, respectively. The Comet search engine considered a precursor mass tolerance of 40 ppm and bins of 1.0005 for the MS/MS. The validity of the peptide spectrum matches were assessed using PatternLab's Search Engine Processor (SEPro) module [7]. Briefly, identifications were grouped by charge state (+2 and > +3) and then by tryptic status (i.e., tryptic or semi-tryptic), resulting in four distinct subgroups. For each result, the XCorr, DeltaCN and Secondary Score values were used to generate a Bayesian discriminator. SEPro then automatically established a cutoff score to accept a false-discovery rate (FDR) of 1% based on the number of decoys, independently performed on each data subset, resulting in a false-positive rate that was independent of tryptic status or charge state [7]. Additionally, a minimum sequence length of 6 amino acid residues was required. Then, only PSMs with less than 5 ppm were considered to compose a final list of proteins supported by at least three independent evidences (e.g., identification of a peptide in different charge states, modified and non-modified version of the same peptide, or different peptides). All identification results are reported with less than 1% FDR, both peptide and protein level, by PatternLab's SEPro module. Spectral counting for estimation of protein copy number was accomplished using the normalized spectral abundance factor (NSAF) [8]. These conditions generate 14,138 peptides which mapped to 4756 proteins, from all samples (HepG2 transfected with the three different plasmids and non-transfected cells) (Supplementary Table S1a-h). Applying the maximum parsimony principle we found 2314 proteins (Supplementary Table S1g). Differentially abundant proteins were pinpointed (Table 2a-c) using PatternLab's TFold module with a Benjamini-Hochberg *q*-value of 0.05 [9].

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#### Transparency document. Supporting Material

Transparency data associated with this article can be found in the online version at http://dx.doi. org/10.1016/j.dib.2016.11.083.

## Appendix A. Supporting Material

Supplementary data associated with this article can be found in the online version at http://dx.doi. org/10.1016/j.dib.2016.11.083.

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