TFEB VARIANTS AND USES THEREOF

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ABSTRACT

The invention refers to TFEB related molecules, as variants, mutants, truncated proteins, chimeras etc. that are constitutively localized in the nucleus of a eukaryote cell. Such molecules have a therapeutic applicability in all of disorders that need of an induction of the cell autophagic/lysosomal system, as lysosomal storage disorders, neurodegenerative diseases, hepatic diseases, muscle diseases and metabolic diseases.
Fig. 2

A: 

B: 

C: 

D: 

E: 

F: 

H: 

G: 

Legend:

- ctr vs TFEB
- ctr vs SIRNA TFEB
- ctr vs ctr starved
- ctr starved vs SIRNA TFEB starved

Legend:

- normal
- starved
- RNAi TFEB starved

Legend:

- normal medium
- starved medium

Legend:

- Flag
- H3

Legend:

- EBSS
- EGF
- FGF
- insulin
- PMMA
- INF
- LF

Legend:

- nuclear localization (%)

Legend:

- flag
- Akt
- Rap
- MEK
- phosphorylated MEK
- phosphorylated Akt
- tubulin
- H3
- pMEK
- p70S6K
- pAKT

Legend:

- nuclear
- cytoplasm
Fig. 3

(a) WT-TFEB vs S142A

(b) Bar graph showing fold change

(c) Western blot analysis of BAF, LC3II, actin, and Lamp1

(d) Western blot analysis with further labeling

(e) Fluorescence images of cells

(f) Western blot analysis of STARV, ERK2-NA, TFE2-FLAG, IP-FLAG, and blot/ERK

(g) Graphs showing CPM values for TFEB-A-142 and TFEB-S-142

(h) Western blot analysis of ERK, flag, and subcellular localization

Legend:
- CTR
- WT-TFEB
- S142A
- Podnik
- TFE2
- S142A
Fig. 5
Fig. 6
Fig. 8
Fig. 9
Fig. 11
Fig. 12
Fig. 15
Fig 16
MPS-III A MEFs

% of enzyme activity released

SGSH3XF + empty  SGSH3XF + TFEB

Fig. 17
Fig 18

[Image of gel electrophoresis with bands labeled as CYTOSOLIC and NUCLEAR samples. Bands include FLAG\textsuperscript{\textalpha}-TFEB, TUBULIN, and H3 under different conditions: DMSO, CQ, and DMSO, StaA.]
Fig 20

A

<table>
<thead>
<tr>
<th>p-S142-TFEB</th>
<th>TFEB-3xFLAG</th>
<th>TOTAL</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
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</table>

<table>
<thead>
<tr>
<th>H3</th>
<th>Tubulin</th>
<th></th>
</tr>
</thead>
</table>

B

C

D

WT | WT + TORIN1 | S109A | S114A |
---|-------------|-------|-------|
S142A | S211A | Nuclear localization

Phenotype

Conservation

mTOR motif

Less | More
Fig 21
TFEB VARIANTS AND USES THEREOF

CROSS-REFERENCE TO RELATED APPLICATIONS


FIELD OF THE INVENTION

[0002] The invention refers to TFEB related molecules, as variants, mutants, truncated proteins, chimeras etc. that are constitutively localized in the nucleus of a eukaryote cell. Such molecules have a therapeutic applicability in all or many of disorders that need of an induction of the cell autophagic/lysosomal system, as lysosomal storage disorders, neurodegenerative diseases, hepatic diseases, muscle diseases and metabolic diseases.

BACKGROUND OF THE INVENTION

[0003] Autophagy is a catabolic process that relies on the cooperation of two distinct types of cellular organelles, autophagosomes and lysosomes (1). During starvation the cell expands both compartments to enhance degradation and recycling processes.

[0004] The lysosome maintains cellular homeostasis and mediates a variety of physiological processes, including cellular clearance, lipid homeostasis, energy metabolism, plasma membrane repair, bone remodeling, and pathogen defense. All these processes require an adaptive and dynamic response of the lysosome to environmental cues. Indeed, physiologic cues, such as aging and diet, and pathologic conditions, which include lysosomal storage diseases (LSDs), neurodegenerative diseases, injuries and infections may generate an adaptive response of the lysosome (34, 35, 36).

[0005] The understanding of the mechanisms that regulate lysosomal function and underlying lysosomal adaptation is still in an initial phase. A major player in the regulation of lysosomal biogenesis is the basic Helix-Loop-Helix (bHLH) leucine zipper transcription factor, TFEB (2). Among the identified TFEB transcriptional targets are lysosomal hydrolases, which are involved in substrate degradation, lysosomal membrane proteins that mediate the interaction of the lysosome with other cellular structures, and components of the vacuolar H+ ATPase (vATPase) complex, which participate to lysosomal acidification (37, 2).

[0006] WO2010/092112 refers to molecules able to enhance the cellular degradative pathway acting on the so called CLEAR element; among them TFEB is listed.

SUMMARY OF THE INVENTION

[0007] The applicants showed that during starvation the cell activates a transcriptional program that controls major steps of the autophagic pathway, including autophagosome formation, autophagosome-lysosome fusion and substrate degradation. The transcription factor EB (TFEB), a previously identified master gene for lysosomal biogenesis (2), coordinates this program by driving expression of both autophagy and lysosomal genes.

[0008] The applicants found that nuclear localization and activity of TFEB are regulated by specific serine phosphorylations. Similar to starvation, pharmacological or gene mutation based inhibition of specific phosphorylation induces autophagy by activating TFEB. These data unveil a novel, kinase-dependent, mechanism involved in the regulation of the lysosomal-autophagic pathway by controlling the biogenesis and partnership of two cooperating cellular organelles.

[0009] Therefore it is an object of the invention herein disclosed a TFEB variant protein that is constitutively localized in the nucleus of a eukaryote cell. The TFEB variant protein of the invention comprises a substitution or alteration of a serine residue to render the same phosphorylation insensitive. The ordinary skilled in the art would recognize that other amino acid substitutions, other than tyrosine, can be made to render the TFEB variant phosphorylation insensitive. For example the serine residue can be replaced with a natural amino acid, for example a neutral amino acid as alanine, or unnatural amino acid. A TFEB variant protein that is constitutively localized in the nucleus of a eukaryote cell comprises mutants, truncated proteins, chimeras of TFEB.

[0010] In a preferred embodiment the TFEB variant protein consists of an amino acid sequence comprised in Seq. Id No. 2 and wherein the substitution of a serine residue is at SER 142 and/or at SER 211 of Seq. Id No. 2. Preferably the amino acid sequence comprised in Seq. Id No. 2 is from aa. 117 to aa. 166 and the substitution of a serine residue is at SER 142 of Seq. Id No. 2 (Seq Id No. 4). Alternatively the amino acid sequence essentially consists of Seq. Id No. 2 and the substitution of a serine residue is at SER 142 and/or at SER 211. In a most preferred embodiment the substitution(s) at SER 142 and/or SER 211 of Seq. Id. No. 2 are to ALA.

[0011] It is another object of the invention the TFEB variant protein as above disclosed for medical use.

[0012] The TFEB variant protein as above disclosed is advantageously used in the treatment of a disorder that needs of the induction of the cell autophagic/lysosomal system, preferably for use in the treatment of any of the following pathologies: lysosomal storage disorders, neurodegenerative diseases, hepatic diseases, muscle diseases and metabolic diseases.

[0013] Examples of lysosomal storage disorder are: activator deficiency/GM2 gangliosidosis, alpha-mannosidosis, aspartylglucosaminuria, cholesteryl ester storage disease, chronic hexosaminidase A deficiency, cystinosin, Danon disease, Fabry disease, Farber disease, fucosidosis, galactosialidosis, Gaucher disease (including Type I, Type II, and Type III), GM1 gangliosidosis (including infantile, late infantile/juvenile, adult/chronic), 1-cell disease/mucolipidosis II, infantile free sialic acid storage disease/ISOS, juvenile hexosaminidase A deficiency, Krabbe disease (including infantile onset, late onset), metachromatic leukodystrophy, pseudo-Hurler polydystrophy/mucolipidosis IIIA, MPS I Hurler syndrome, MPS I Scheie syndrome, MPS I Hurler-Scheie syndrome, MPS II Hunter syndrome, Sanfilippo syndrome type A/MPS II A, Sanfilippo syndrome type B/MPS IIIB, Morquio type A/MPS IVA, Morquio Type B/MPS IVB, MPS IX hyaluronidase deficiency, Niemann-Pick disease (including Type A, Type B, and Type C), neuronal ceroidlipofuscinoses (including CLN6 disease, atypical late infantile, late onset variant, early juvenile Batten-Spielmeyer-Vogt/juvenile NCL/CLN3 disease, Finnish variant late infantile CLN5, Jansky-Bielschowsky disease/late infantile CLN2/TPP1 disease, Kufs/adult-onset NCL/CLN4 disease, north-
em epilepsy/variant late infantile CLN8, and Santavuori-Hultia infantile/variant late infantile CLN1/PPT disease), beta-mannosidosis, Pompe disease/glycogen storage disease type II, pycnodysostosis, Sandhoff disease/adult onset/GM2 gangliosidosis, Sandhoff disease/GM2 gangliosidosis infantile, Sandhoff disease/GM2 gangliosidosis juvenile, Schindler disease, Salla disease/sialic acid storage disease, Tay-Sachs/GM2 gangliosidosis, Wolman disease, Multiple Sulfatase Deficiency. [0014] Examples of hepatic diseases are: Alpha1 antitrypsin deficiency and Fatty liver disease. [0015] Examples of muscle diseases are: Autophagic Vacuolar Myopathies and X-linked myopathy with excessive autophagy. [0016] Examples of metabolic diseases are: hypercholesterolemia and fatty liver disease. [0017] Examples of neurodegenerative diseases are: Alzheimer’s disease, Parkinson’s disease, Huntington’s disease, Creutzfeldt-Jakob disease, and spinocerebellar ataxia. [0018] It is a further object of the invention a nucleic acid comprising a coding sequence encoding for the TFEB variant protein as above disclosed. Preferably the nucleic acid comprises the sequence of Seq. Id No. 3. [0019] It is a further object of the invention an expression vector comprising under appropriate regulative sequences the nucleic acid as above disclosed. [0020] The expression vector of the invention may advantageously be used for gene therapy. [0021] It is a further object of the invention a method for increasing the production of endogenous or recombinant lysosomal enzymes in an ex vivo cultured cell comprising the steps of:—introducing the nucleic acid according or the expression vector as above disclosed in said cell and—allowing the expression of the encoded TFEB variant protein. [0022] It is a further object of the invention a method of treating a disorder by administering to a subject a therapeutically effective amount of the TFEB variant protein as above disclosed, preferably when the disorder is alleviated by the induction of the cell autophagic/lysosomal system. [0023] More preferably the disorder is selected from the group comprising lysosomal storage disorders, neurodegenerative diseases, hepatic diseases, muscle diseases and metabolic diseases. Examples of such disorders were above provided.

BRIEF DESCRIPTION OF THE DRAWINGS

[0024] FIG. 1 TFEB induces autophagy. (A) HeLa cells stably overexpressing TFEB were transfected with a GFP-LC3 plasmid and treated as indicated. Approximately 100 cells were analyzed in triplicate for each experiment. The graph shows quantification of GFP-positive vesicles. (B-F) Western blot analysis of LC3 in (B) TFEB-3xFlag stable overexpressing (+) and control cells (-). The graph represents the quantification using imageJ software analysis of LC3II expression (relative to actin) from three independent blots; (C) TFEB stable overexpressing cells, which were serum and amino acid-starved (Starv) for the indicated time (h-hours), (D-F) cellular lysates isolated from TFEB-RNAi and control cells treated with scrambled RNAi (ctr) cultured in (D) normal media, (E) starved media, or (F) starved media supplemented with bafilomycin (4 h; 400 nM). The graph represents the quantification of LC3II expression (relative to actin) from three independent blots and band intensities were quantified using imageJ software analysis. (G) TFEB mRNA levels were analyzed by qPCR using cDNAs prepared from cells transfected with 3 different siRNA oligos targeting TFEB (oligo #1, #2, #3), or with a scrambled siRNA oligo (ctr). (H) Representative confocal images of fixed HeLa cells stably expressing GFP-mRFP-LC3 transfected with empty (control) or TFEB vector. A minimum of 2000 cells was counted and the values represent the average number of vesicles (relative to the control, %) obtained from three independent experiments. AL (autolysosomes)—mRFP positive/GFP negative vesicles; total: mRFP positive vesicles. (All error bars represent standard deviations. T-test (unpaired) p value (*<0.05, (**)<0.01)

[0025] FIG. 2 Starvation regulates TFEB nuclear translocation and activity. (A) Scatter Plot graphs displaying the logarithmic value of the fold change differences in the relative expression levels of 51 autophagy-related genes in HeLa cells cultured in different conditions. X-axis—control group. Y-axis—treated group. Circles represent genes with increased (red) or decreased (green) fold change. Comparisons were as indicated. (B) Chromatin immunoprecipitation (ChiP) analysis. The histogram shows the amount of immunoprecipitated DNA as detected by qPCR assay. Values were normalized to the input and plotted as relative enrichment over a mock control. Experiments were performed in triplicate. (C) qPCR analysis of TFEB-target gene expression in normal, starved, and in TFEB-siRNA starved cells. GAPDH and HPRT represents housekeeping genes, while ATG10, ATG9A and ATG4D represent control genes (non-TFEB target genes). (D-F) HeLa cells stably overexpressing TFEB were left untreated or nutrient starved for 4 h. (D) Five fields containing 50-100 cells each were analyzed for TFEB nuclear localization. P value (<=0.01. (E) Cells were subjected to nuclear/cytosolic fractionation and blotted with Flag antibody. H3 and tubulin were used as nuclear and cytosolic markers, respectively. (F) Nuclear fractions were blotted with Flag and H3 (loading control) antibodies. (G) Western blot analysis of Flag, tubulin and H3 in nuclear extracts prepared from normal, starved and starved/stimulated with normal media cells for 1 h (normal) or pretreated with AP-2 (AKT inhibitor), Rapamycin (mTOR inhibitor) and U0126 (MEK inhibitor) 1 h prior to media stimulation. Total extracts were used to verify the efficiency of the inhibitors. (H) qPCR analysis of lysosomal and autophagic genes in TFEB siRNA or TFEB-scrambled control cells transfected with either a constitutive active MEK (caMEK) plasmid or with an empty vector. Stimulation was performed where indicated. (All error bars represent standard deviations. T-test (unpaired) p value (*<0.05, (**)<0.01))

[0026] FIG. 3 Serine phosphorylation regulates TFEB activation. (A) TFEB subcellular localization in HeLa cells expressing mutated versions of TFEB-3xFlag, immunostained with Flag antibody. Five fields from three independent experiments, containing 50-100 cells each were analyzed. (B) qPCR analysis of TFEB target gene expression 24 h post-transfection with empty, normal and mutant TFEB plasmids. (C,D) Western blot analysis of LC3 and Lamp1 (C) and protein extracts from HeLa cells transfected with equal amounts of empty (pcDNA), TFEB-3xFlag or TFEBsi142A-3xFlag vectors. Bafilomycin was added where indicated. Westerns were done in triplicate and the quantification of proteins levels were normalized to actin levels. (E) Analysis of autolysosomes (AL—mRFP positive/GFP negative) in HeLa cells stably expressing GFP-mRFP-LC3 and transfected with either pcDNA, Tfeb or Ser-Tfeb for 24 h. Quantification as reported in FIG. 1H. (F) Western blot analy-
sis using anti-Erk antibody on HeLa cells transfected with HA-Erk2 and/or TFEB-3xFlag, kept in full serum or nutrient starved for 4 h and immunoprecipitated with anti-Flag antibody. Lysates were immunoprecipitated with anti-
Flag and blotted with an anti-Erk antibody. (G) In vitro kinase assay. Recombinant kinases were incubated in the presence of ATP γS 32P and a peptide spanning from amino acid 120 to 170 of TFEB protein (TFEB-S-142) or with a similar peptide in which serine 142 was substituted with alanine (TFEB-A-142). Phosphorylation efficiency (“phosphorylation sensitivity”) was measured as the amount of radioactivity incorporated by the peptides. (H) HeLa stable clones overexpressing TFEB were transfected with siRNA oligonucleotides specific for ERK1/2 or with control siRNA. 48 h later cells were left untreated, serum starved or serum and amino acid (a.a.) starved for 4 h, harvested and subjected to nuclear isolation and Flag immunoblotting. Total lysates were probed with ERK antibody. All error bars represent standard deviations. P value (*)<0.05.

[0027] FIG. 4 In vitro analysis of TFEB-mediated induction of autophagy. (A) Immunofluorescence analysis of GFP-positive vesicles in fed, 16 h-fasted, and 24 h-fasted mice. Quantification of vesicles is shown in the graph. (B) qPCR analysis of TFEB target gene expression in liver samples from fed and fasted animals (n=3; Error bars represent standard deviations, P value (*)<0.05). Gapdh and Hprt were used as reference genes. (C,D) Analysis of TFEB subcellular localization in two-mouse old wild type mice infected with AAV-9Tcfe6-11A and fasted 16 h prior to sacrifice. (C) HA-immunofluorescence analysis. The graph shows quantification of nuclear HA signal. 100 transduced cells were counted for each liver. n=3 mice/group. *p<0.001. (D) Western blot analysis of HA, Tubulin and H3 in liver specimens subjected to nuclear fractionation. Total liver lysates were probed with an HA antibody to verify comparable transgene expression between fed and fasted animals. (E) Western blot analysis of LC3, actin, p-ERK1/2 and ERK1/2 in liver extracts from mice injected AAV-9Tcfe6-11A. (F) Western blot analysis of GFP and DAPI staining in cryopreserved liver slices from 2-month old GFP-LC3-transgenic mice injected with either AAV- 
Tcfe6-HA or with saline solution (control group) and fed ad libitum or fasted for 24 h prior sacrifice. Quantification of GFP-positive vesicles is shown in the graph. (G) qPCR analysis of both autophagic and lysosomal TFEB-target gene expression in liver samples isolated from conditional Tcfe6-
3xFLAG transgenic mice (Tcfe6-3xFlag;AlbCRE), in which transgene expression is driven by a liver-specific CRE recombinase (i.e. Albumin-CRE). (H) Western blot analysis of LC3 and actin in liver protein extracts from Alb-CRE, Tcfe6-3xFlag and Tcfe6-3xFlag;Alb-CRE mice.

[0028] FIG. 5 TFEB transient overexpression induces autophagy. (A) HeLa cells were transiently transfected with a plasmid encoding for tagged TFEB protein. 48h after transfection cells were collected, lysed and 10 mg of protein samples were analyzed for LC3, Flag and actin immunoreactivity. Experiments were performed in triplicate and band intensities were quantified using imageJ software analysis (Error bars represent standard deviations. p value (**)<0.05) (B) COS-7 cells were transiently transfected with an empty vector or with a TFEB-3xFlag vector. 24h later cells were treated for 4h with lysosomal inhibitors (pepsatin/E64, 10 μg/ml, SIGMA). 10 μg of cell lysates were subjected to LC3 and actin immunoblotting.

[0029] FIG. 6 Induction of autophagy in Tcfe6 overexpressing MEFS. (A,B) Electron micrograph of MEFS infected with lentivirus expressing Tcfe6 and control cells. (a) Autophagic structures were observed upon Tcfe6 expression, including autophagosomes (AV) and autolysosomes (AL). (B) Formation of early autophagosomes. Isolation membrane (arrows) surrounding electron-dense cytoplasmatic material. (C) Quantitation of number of autophagic structure (AV and AL) and (D) of early autophagosomes. At least 30 cells/group were analyzed. Error bars represent SEM; p value (*)<0.05, (***)<0.0001.

[0030] FIG. 7 TFEB promotes autophagosome formation. (A) Control and stable TFEB-overexpressing cells were treated with baflomycin (12h 400 nM) harvested and subjected to LC3II, Flag and actin immunoblotting. (B) Control and TFEB-overexpressing cells were left untreated or treated with 10 μg/ml lysosomal inhibitor pepstatin/E64 for 4 h, lysed and subjected to LC3, Flag and actin immunoblotting. Experiments were performed in triplicate and band intensities were quantified using imageJ software analysis (Error bars represent standard deviations. p value (**)<0.05).

[0031] FIG. 8 TFEB increases autophagic proteolysis. Rate of long-lived protein degradation in Tcfe6-overexpressing, Tcfe6-depleted and control cells in either normal or starved condition. 3-methyl adenine (SMA) was added where indicated (Error bars represent standard deviations. p value (**)<0.05).

[0032] FIG. 9 Distribution of the TFEB putative binding elements in the promoter regions of a subset of autophagy genes. Numbers indicate the distance of the binding element from the transcription start site (TSS).

[0033] FIG. 10 Starvation enhances TFEB activity. Luciferase reporter assay using a construct carrying four tandem copies of TFEB binding sites. Both normal and TFEB-overexpressing HeLa cells were transfected with an artificial promoter with TFEB binding sites. Both cells types displayed increased transactivation potential when cultured in starved conditions. (Error bars represent standard deviations p (**)<0.05)

[0034] FIG. 11 Starvation induces TFEB nuclear translocation through MAPK. (A) Starvation induces cytosolic TFEB mobility shift and nuclear translocation. Normal medium; starved medium (4 h); starved+normal, indicates that cell were cultured in starved medium (4 h) and supplemented with normal medium 1 h prior to harvesting. Cytosolic and nuclear fractions were subjected to Flag immunoblotting. (B) Analysis of TFEB cellular localization by immunofluorescence in HeLa cells treated as indicated in FIG. 2G. The graph shows percentage of cells that display TFEB nuclear localization. Error bars represent standard deviations. P value (**)<0.05

[0035] FIG. 12 TFEB nuclear traslocation is dependent on S142 phosphorylation. (A) HeLa cells expressing Tcfe6-3xFlag, S142A-3xFlag, S332-3xFlag or S423-3xFlag proteins were subjected to nuclear protein isolation. Equal amounts of nuclear proteins were verified by ponceau staining. (B) HeLa cells expressing Tcfe6-3xFlag, S142A-3xFlag and S142D- 
3xFlag proteins were subjected to nuclear protein isolation in normal and in starved conditions. (C) Flag immunoblotting of cytosolic protein isolated from HeLa cells expressing Tcfe6-
3xFlag and Tcfe6-S142A-3xFlag showing that in normal media S142A migrates as lower MW band compared to WT TFEB while this shift is not evident anymore in starved conditions. (D) Flag immunoblotting of cytosolic protein iso-
lated from starved HeLa cells expressing TFEB-3xFlag, S142A-3xFlag and S142D-3xFlag showing a reduced shift of TFEB-S142D.

**[0036]** FIG. 13 S142A TFEB mutant displays enhanced activity. HeLa cells stably overexpressing GFP-LC3 were transfected with equal amounts of empty, TFEB-3xFlag or S142A-TFEB-3xFlag plasmids and the number of autophagosomes was quantified. At least ten fields (containing 4-10 cells) were analyzed for each point. Experiments were performed in triplicate. Error bars represent standard deviations. p value (*p<0.05).

**[0037]** FIG. 14 Multiple sequence alignment of TFEB-human S142 phosphorylation site with TFEB paralogues, MITF and relevant TFEB-related family members. TFEB-human homologs were identified by BLAST (2.2.17) search against UniProtKB database at ExPASy Proteomics Server. The applicants removed the hits with “putative”, “uncharacterized” and “cDNA” keywords and hits without gene names. Next, the applicants authors aligned the remaining homologs with ClustalW (1.82). The multiple sequence alignment was generated by Seaview. The figure shows only a 20 amino acid-long segment of TFEB_HUMAN sequence aligned with other proteins from TFEB, MITF, TCFEB, TF3 and TCFE3 families. “sp” stands for SwissProt entry, while “tr” denotes Tremble entry. P19498 is a UniProtKB accession code. TFEB_B_HUMAN indicates gene name and species name respectively.

**[0038]** FIG. 15 Strategy for TcFEb overexpression in vivo. (A) Representative images of cryopreserved liver slices immunostained with anti-HA antibody (to verify viral transduction efficiency). (B) Liver protein extracted from Tcfeb-HA injected and control mice were immunoblotted HA and actin antibodies. (C) Generation of a transgenic mouse line for TcFEB conditional overexpression. The map of the transgene vector, before and after CRE recombinase is illustrated at the top. Representative genotypes of littermates are shown on the left, while the correspondent liver-specific TcFEB overexpression in mouse n4 is shown on the right.

**[0039]** FIG. 16 TFEB overexpression increases the release of lysosomal enzymes in the culture medium of MEFs, NSCs, HeLa, and COS-7 cells. Activities of lysosomal enzymes acid phosphatase, beta-galactosidase, and beta-nicotinamide were determined in the culture medium and in cells transfected with either an empty vector or with a TFEB-expression vector. HeLa, Cos 7 cells and mouse embryonic fibroblasts from mouse models of MLIV (S7), MPSIII A (S7), and MSD were transfected using PolyFect Transfection Reagent (Qiagen) or lipofectamine 2000 Reagent (Invitrogen), according to the manufacturer’s protocols. TFEB-3xFLAG HeLa stable cell lines (CF7) was previously described (2). The figure shows percentages of enzyme activities released compared to total activities.

**[0040]** FIG. 17 TFEB exerts a positive control on lysosomal exocytosis. MPSIIIA MEF Cells were maintained in DMEM supplemented with 10% FBS and penicillin/streptomycin (normal culture medium). Sub-confluent cells were transfected using Lipofectamine™ 2000 (Invitrogen) according to manufacturer’s protocols. MPS-IIIA MEFs were co-transfected with a plasmid encoding a tagged sulfamidase (SGSH3xFlag) and either an empty plasmid or a plasmid encoding TFEB. One day after transfection the medium was replaced with DMEM 0.5% FBS. Two days after transfection the conditioned medium and the pellet were collected for sulfamidase activity measurement and the percentage of the enzyme released in the medium calculated.

**[0041]** FIG. 18 Lysosomal stress induces TFEB nuclear translocation. Immunoblotting of proteins extracted from HeLa cells that express TFEB-3xFlag treated with chloroquine (CQ) or Salicylilalamide A (SalA), subjected to nuclear/cytosolic fractionation and blotted with antibody against FLAG to detect TFEB. Histone 3 (H3) and tubulin were used as nuclear and cytosolic markers, respectively. Blots are representative of triplicate experiments.

**[0042]** FIG. 19 mTORC1 regulates TFEB. (A) Lysosomal stress inhibits mTOR signalling. Immunoblotting of protein extracts isolated from HeLa cells treated overnight, as indicated. Membranes were probed with antibodies for p-T202/ Y204-ERK1/2, ERK1/2, p-T389-S6K, and S6K to measure ERK and mTORC1 activities. (B) Torin 1 induces TFEB dephosphorylation and nuclear translocation. FLAG immunoblotting of cytosolic and nuclear fractions isolated from TFEB-3xFLAG HeLa cells cultured in amino acid-free media and subsequently stimulated as indicated for at least 3 h. Correct subcellular fractionation was verified with H3 and tubulin antibodies. (C, D) Effects and dose-response curves of ERK and mTOR inhibitors on TFEB nuclear translocation. TFEB-GFP HeLa cells were seeded in 384-well plates, incubated for 12 h, and treated with 10 different concentrations of the ERK inhibitor U0126 or the mTOR inhibitors Rapamycin, Torin 1 and Torin 2 ranging from 2.54 nM to 50 µM. After 3 h at 37°C in RPMI medium containing one of the compounds, the cells were washed, fixed, and stained with DAPI and photographed by using confocal automated microscopy (Opera high content system, Perkin Elmer). (C) Representative images of test concentrations for each compound. Scale bars represent 30 µm. (D) The graph shows the percentage of nuclear translocation at the 10 different concentrations of each compound (in log of the concentration). The EC50 for each compound was calculated using Prism software (see Materials and methods for details). (E) Amino acids induce TFEB molecular weight shift. Immunoblotting of protein extracts isolated from HEK-293T cells transfected either TFEB-3xFLAG or with an empty vector were nutrient starved and stimulated for 50 min with amino acids (a.a.). Antibody used were p-T389-S6K, S6K and FLAG. (F) Rag knockdown induces TFEB nuclear translocation. HeLa cells stably expressing Flag-3xTFEB were infected with lentiviruses encoding a Short hairpin (Sh-) RNA targeting lenseurine (control) or RagC and RagD mRNAs. In all, 96 h post infection, cells were left untreated (N=normal media), starved (S=starved media) or treated with Torin 1 (T=Torin 1) for 4 h and then subjected to nuclear/cytosolic fractionation. TFEB localization was detected with a FLAG antibody, whereas tubulin and H3 were used as controls for the cytosolic and nuclear fraction, respectively; levels of S6K phosphorylation were used to test RagC and RagD knockdown efficiency. (G) mTORC2 does not affect TFEB phosphorylation. Mouse embryonic fibroblasts (MEF) isolated from Sin1−/− or control embryos (E14.5) were infected with a retrovirus encoding TFEB-3xFLAG; 48 h post infection, cells were treated with Torin 1 (T) for 4 h, where indicated, subjected to nuclear/ cytosolic fractionation and immunoblotted for FLAG, tubulin, and H3.

**[0043]** FIG. 20 mTORC1 phosphorylates TFEB at serine 142 (S142). (A) Torin 1 induces S142 dephosphorylation. HeLa cells were treated as indicated and total and nuclear extracts were probed with a TFEB p-S142 phospho-antibody.
and with anti-FLAG antibody. (B) Schematic representation of TFEB protein structure with the predicted mTORC1 phosphorylation sites and their conservation among vertebrates. Numbering is according to human isoform 1. (C) Sequence conservation scores of the phosphorylation sites and quantitative agreement between mTOR consensus motif and the sequence around the phosphorylation sites of TFEB. (D) S142 and S211 regulate TFEB localization. Flag immunostaining of TFEB subcellular localization in HeLa cells expressing serine-to-valine mutated versions of TFEB-3×Flag. Nuclei were stained with DAPI. Values are means of five fields containing at least 50 transfected cells. Student’s t-test (unpaired) ***P<0.001. Scale bars represent 30 p.m.

**Fig. 21** The lysosome regulates gene expression by TFEB. (A) Chloroquine treatment inhibits mTORC1 activity in primary hepatocytes. Primary hepatocytes isolated from 2-month-old Tcfebfloxb/fox (control) and Tcfebfloxb/flox; Alb-Cre(Tcfeb−/−) mice were left untreated or treated overnight with Torin 1, U0126, or Chloroquine. Subsequently, cells were lysed and protein extracts were immunoblotted with the indicated antibodies. (B, C) TFEB mediates the transcriptional response to chloroquine and Torin 1. Quantitative PCR (qPCR) of TFEB target genes in primary hepatocytes from control (floxb/fox) and Tcfeb−/− (floxb/flox; alb-Cre) mice. Cells were treated with Chloroquine (left) and Torin 1 (right). The expression levels were shown as % increased expression of the treated versus the corresponding untreated samples. Values represent means ±d of three independent hepatocyte preparations (three mice/genotype). Student’s t-test (two tailed) *P-values<0.05.

**DETAILED DESCRIPTION OF THE INVENTION**

Materials and Methods

Cell Culture and Media and Drugs and Cellular Treatment

[0045] HeLa and COS and HEK-293T cells were purchased from ATCC. Cells were cultured in the following media: (normal) DMEM high glucose supplemented with 10% FBS; (starvation) HBSS media with Ca and Mg supplemented with 10 mM HEPES; (Serum) EBSS supplemented with 20% FBS; (amino acid media) Glucose and serum free DMEM; Drugs treatment: Rapamycin (2.5 mg/ml, SIGMA) 2-4 h otherwise indicated; Bafilomycin, (400 mg, SIGMA) 2-4 h; Insulin (100 mg/ml SIGMA) for 2 h; EGF, FGF (BD biosciences); LIF (100 mg/ml SIGMA); Millipore) 2 h; PMA (1 mg/ml) 2 h. U0126 (MEKI) were used at 25 mM (Cell Signaling). AIP1 (AKT inhibitor) were used at 1 mM. Lysosomal inhibitors were pepstatin and E64 (10 mg/ml 4 h SIGMA). The following drugs were used in the experiments of FIGS. 18-2: Rapamycin (2.5 μM 5 μM, otherwise indicated) from SIGMA; Torin1 (250 nM 250 nM, otherwise indicated) from TOCRIS; U0126 (50 μM 50 μM) from Cell Signaling technology; Chloroquine (100 μM 100 μM) from SIGMA; Salicylhalamide A (2 μM 2 μM) was a kind gift from Jeff De Brabander (UT Southwestern).

[0046] Primary hepatocytes were generated as follow: 2-month-old mice were deeply anesthetized with Avertin (240 mg/kg) and perfused first with 25 ml of HBSS (Sigma H6648) supplemented with 10 mM HEPES and 0.5 mM EGTA and after with a similar solution containing 100 U/ml of Collagenase (Wako) and 0.05 mg/ml of Trypsin inhibitor (Sigma). Liver was dissociated in a petri dish, cell pellet was washed in HBSS and plated at density of 5x10⁶ cells/35 mm dish and cultured in William’s medium E supplemented with 10% FBS, 2 mM glutamine, 0.1 mM Insulin, 0.1 mM Dexamethasone and pen/strep. The next day, cells were treated as described in the text. Sin 1- and control MEFs were generated as previously described(46) and maintained in DMEM supplemented with 10% FBS, glutamine and pen/strep.

Generation of a Tcfeb™ mouse line

[0047] The applicants used publicly available embryonic stem (ES) cell clones (http colont double forward slash www dot eucomm dot org trailing slash) in which Tcfeb was targeted by homologous recombination at exons 4 and 5. The recombinant ES cell clones were injected into blastocysts, which were used to generate a mouse line carrying the engineered allele. Liver-specific KO was generated crossing the Flox/Flox mice with a transgenic line expressing the CRE under the Albumin promoter (ALB-CRE) obtained from the Jackson laboratory. All procedures involving mice were approved by the Institutional Animal Care and Use Committee of the Baylor College of Medicine.

Transfection, Plasmids and siRNA

[0048] Both plasmids and siRNA were transfected with lipofectamine LTX (Invitrogen) using a reverse transfection protocols. siRNA-transfected cells were collected after 48 or 72 h. siRNA TFEB were used at 50 nM (Dharmacon), siRNA ERK1/2 were used at 100 nM (Cell Signaling).

[0049] Cells were transiently transfected with DNA plasmids pPR5-5mycPA1, pCEP4-TFEB-bis, pCMV2-TFEB, and p3xFLAG-CMV-TFEB using lipofectamine2000 or LTX (Invitrogen) according to the protocol from manufacturer. Site-direct mutagenesis was performed according to the manufacturer instructions (Stratagene) verifying the correct mutagenesis by sequencing.

Western Blotting

[0050] Cells or tissues were solubilized in RIPA buffer supplemented with protease (ROCHE) and Phosphatase (SIGMA) inhibitors. From 10 to 30 micrograms were loaded on 4-12% Bis-Tris gel (NUPAGE, Invitrogen), transferred to PVDF membranes and analyzed by western blot using the ECL method (Pierce). The following antibodies were used: LC3 (Novus Biological), FLAG, b-ACTIN, TUBULIN (SIGMA), HA (Covance), H3, ERK1/2, p-ERK1/2, p-AKT, p-70S6K (Cell Signaling), ERK2 (Santa Cruz). Protein levels were quantified by using ImageJ software analysis.

Nuclear/Cytosolic Fractionation

[0051] Cells were seeded at 50% of confluence in 6 well dishes and serum starved overnight (ON). Normal medium was added the following day either in presence of DMSO or kinase inhibitors. Subcellular fractionation was carried out as previously reported. Briefly, cells were lysed in 0.5 Triton X-100 lysis buffer (50 mM Tris-HCl, 0.5% triton, 137.5 mM NaCl, 10% glycerol, 5 mM EDTA supplemented with fresh protease and phosphatase inhibitors. Supernatant represented cytosolic fraction while nuclear pellet was washed twice and lysed in 0.5 Triton X-100 buffer 0.5% SDS and sonicated.

Degradation of Long-Lived Proteins

[0052] Sub-confluent cells were incubated with L-U14C-serine for 20 h and chased for 1 h with cold media to degrade short-lived proteins. Subsequently cells were incubated with either normal media or starvation media (eventually in the presence of 3-MA) for 4 h. The rate of long-lived protein
degradation was calculated from the ratio of soluble radioactivity in the media to that insoluble in the acid-precipitable cell pellet.

RNA Extraction, Reverse Transcription, ChIP and Quantitative PCR

Total RNA was extracted from tissues using TRIzol (Invitrogen) or from cells using RNAasy column (Qiagen). Reverse transcription was performed using TaqMan reverse transcription reagents (Applied Biosystems). Lysosomal and autophagic gene specific primers were previously reported. Autophagy gene primers and mouse primers were purchased from SABioscience. Fold change calculations were calculated using SABioscience's online data analysis website (http colon double forward slash www dot sabiosciences dot com forward slash per forward slash analyze dot analysis dot php) which uses the DDC method. In brief, the average of the most stable housekeeping genes (GAPDH, ACTB, B2M, RPL13A, HPRT and Cyclophilin) were used as "normalizer" genes to calculate the DDC value. Next, the DDC value is calculated between the "control" group and the "experimental" group. Lastly, the fold change is calculated using $2^{(-\Delta\Delta CT)}$. Biological replicates were grouped to allow calculating the fold change values. Unpaired T-Test was used to calculate statistical significance. Asterisks in the graph indicate that the P-value was $<0.05$.

Protein Kinase Prediction

Applicants used five methods including CrPhos0.8, GPS-2.1, PhosphoMotifFinder, Networkin and PhosphoID using the default parameters (15-19). They further filtered CrPhos0.8 and GPS-2.1 predictions according to their confidence scores. For the former, we took into account the predictions with a false positive rate (FPR) equals or less than 30%. For the latter, they considered the predictions with score equals or higher than 5. GPS-2.1 scores were calculated as the difference between actual score and threshold values. We took all the predictions from other three methods in the case of Networkin, we combined predictions from both Networkin and Networkin 2. Each method describes the kinases associated by S142 site in a different kinase classification, which simply involves four hierarchical levels: kinase family, kinase subfamily and kinase itself. To obtain a general consensus in each hierarchical level, we classified each prediction in these four hierarchical levels, if the predictions were not already classified in that manner, they searched for the missing classifications at the http colon double forward slash slack kinase dot org forward slash slack kinase database under vertebrate Glade and human species. Consensus in each classification is found according to the majority vote in each classification.

In Vitro Kinase Assay

TFEB-S-142: aa. o 117-166 of Seq Id No. 2:

PPPAAPPGVRAGHVLSSAGNSAPNPS-
MAMLHIGSNPERELDDVIDNIMR and TFEB-A-142: Seq Id No. 4, corresponding to aa. of 117-166 of Seq Id No. 2 where Ser 142 was substituted with Ala (bold):

were synthesized by GENESCRIPT corp. The test peptides TFEB-A-142 and TFEB-S-142 were made up to 1 mM in 50 mM HEPES pH7. There appeared to be no issue with dissolution. The kinase assay was performed at room temperature for 40 minutes at 200 μM ATP and 100 μM of each peptide, using Millipore’s standard radiometric assay. All protein kinases were used at their standard Kinase FLUORER assay concentration. Following incubation, all assays were stopped by the addition of acid and an aliquot spotted onto P30 and Filtermat A to separate products. All tests were carried out in triplicate, and the usual substrate for each protein kinase included as a control.

In Vivo Gene Delivery

The mice were housed in the transgenic mouse facility of Baylor College of Medicine (Houston, Tex., USA). GFP-LC3 transgenic mice were a kind gift of N. Mizushima. C57B6 female mice (4 weeks old) were used, if not otherwise specified. The AAV vector was produced by the TIGEM AAV Vector Core Facility. Briefly, the mouse TFEB (TeFEB) coding sequence was cloned into the pAAV2.1-CMV-GFP plasmid by replacing the GFP sequence and fused in frame with a HA tag. The resulting pAAV2.1-CMV-TeFEB-HA was then triple transfected in sub-confluent 293 cells along with the pAd-Helper and the pack2/9 packaging plasmids. The recombinant AAV2/9 vectors were purified by two rounds of CsCl. Vector titers, expressed as genome copies (GC/mL), were assessed by both PCR quantification using TaqMan (Perkin-Elmer, Life and Analytical Sciences, Waltham, Mass.) and dot blot analysis. Each mouse was retro-orbital injected with $1.25 \times 10^{11}$ viral particle and sacrificed after 3 weeks. Starved mice were fed-deprived for 16 h when analyzed for gene expression, or for 24 h when analyzed for GFP-LC3 dots number.

Histology and Immunofluorescence

Liver samples were collected and fixed overnight in 4% paraformaldehyde in PBS. After cryoprotection in 10 and 30% sucrose in PBS, the specimens were frozen in OCT (Sakura Finetech, Torrance, Calif.) and sectioned 30 μm thick. Images were taken on an AxioImager2 (Zeiss, Thornwood, N.Y.). For immunofluorescence, slices were blocked for 2 h at RT in 2.5% BSA in PBS+0.1% Triton X-100. After blocking, specimens were incubated for 20 h with the primary antibody and, after 3x washes in PBS+0.05% TX-100, for 3 h with secondary antibodies conjugated either with Alexafluor 488 or Alexafluor 555 (Invitrogen). For immunohistochemistry analyses of HA the avidin-biotin complex (ABC) method was used (Vectastain Elite ABC kit). Anti-GFP was from Abcam; (dilution 1:500).

Electron Microscopy

Control and TFEB-overexpressing cells were washed with PBS, and fixed in 1% glutaraldehyde dissolved in 0.2 M Hepes buffer (pH 7.4) for 30 min at room temperature. The cells were then postfixed for 2 h in OsO4. After dehydration in graded series of ethanol, the cells were embedded in Epon 812 (Fluka) and polymerized at 60° C. for 72 h. Thin sections were cut at the Leica EM UC6, counterstained with uranyl acetate and lead citrate. EM images were acquired from thin sections using a Philips Tecnai-12 electron microscope equipped with an ULTRA VIEW CCD digital camera (Philips, Eindhoven, The Netherlands). Quantifica-
tion of vacuolization was performed using the AnalySIS software (Soft Imaging Systems GmbH, Munster, Germany). Selection of cells for quantification was based on their suitability for stereologic analysis, i.e., only cells sectioned through their central region (detected on the basis of the presence of Golgi membranes) were analyzed.

Animal Models
[0059] All procedures involving mice were approved by the Institutional Animal Care and Use Committee of the Baylor College of Medicine. GFP-LC3 transgenic line was described previously. Tissue specific overexpression of Tcfeb was generated as follows: Tcfeb-Lc3xFlag cDNA was inserted after a CAGCAT cassette [chicken aminprotease (CAG)] followed by chloramphenicol acetyltransferase (CAT) cDNA flanked by 2 loxP sites and used to generate transgenic mice (Baylor College of Medicine transgenic core). Mice were then crossed with Albumin-CRE (obtained from the Jackson laboratory) line. For 48 Starvation protocol the mice were food deprived for 22 h, subsequently were fed for 2 h and fasted again for 24 h prior sacrifice.

Enzymatic Activities
[0060] Lysoosomal enzymes acid phosphatase, beta-galactosidase, and beta- hexosaminidase activities were measured using the appropriate fluorometric or colorimetric substrates. SODH activity was measured following protocols described in Fraldi et al., Hum Mol Gen 2007 (33).

Immunoblotting and Antibodies
[0061] The mouse anti-TFEB monoclonal antibody was purchased from My Biosource catalogue No. MBS 120432. To generate anti-pS142 specific antibodies, rabbits were immunized with the following peptide coupled to KLH: AGNSAPN[pSer]PMAMHLIC. Following the fourth immunization, rabbits were sacrificed and the serum was collected. Non-phosphospecific antibodies were depleted from the serum by circulation through a column containing the non-phosphorylated antigen. The phosphospecific antibodies were subsequently purified using a column containing the phosphorylated peptide.

Cells were lysed with M-PER buffer (Thermo) containing protease and phosphatase inhibitors (Sigma); nuclear/cytosolic fractions were isolated as above described. Proteins were separated by SDS-PAGE (Invitrogen; reduced NuPAGE 4-12% Bis-tris Gel MES SDS buffer). If needed, the gel was stained using 20 ml Imperial Protein Stain (Thermo Fisher) at room temperature for 1 h and destained with water. Immunoblot analysis was performed by transferring the protein onto a nitrocellulose membrane with an I-Blot (Invitrogen). The membrane was blocked with 5% non-fat milk in TBS-T buffer (TBS containing 0.05% Tween-20) and incubated with primary antibodies anti-FLAG and anti-TUBULIN (Sigma; 1:2000), anti-H3 (Cell Signaling; 1:10 000) at room temperature for 2 h whereas the following antibodies were incubated ON in 5% BSA: anti-TFEB (My Biosource; 1:1000), anti-P TFEB (1:1000) ERK1/2, p-ERK1/2, p-P70S6K, P70S6K (Cell Signaling; 1:1000). The membrane was washed three times with TBS-T buffer and incubated with alkaline phosphatase-conjugated IgG (Promega; 0.2 mg/ml) at room temperature for 1 h. The membrane was washed three times with TBS buffer and the expressed proteins were visualized by adding 10 ml Western Blue Stabilized Substrate (Promega).

High Content Nuclear Translocation Assay
[0062] TFEB-GFP cells were seeded in 384-well plates, incubated for 12 hours, and treated with ten different concentrations (5000 nM, 16666.6 nM, 5555.5 nM, 1851.85 nM, 617.28 nM, 205.76 nM, 68.58 nM, 22.86 nM, 22.86 nM, 7.62 nM, and 2.54 nM) of ERK inhibitor U0126 (Sigma-Aldrich) and mTOR inhibitors Rapamycin (Sigma-Aldrich), Torin 1 (Biomarin), and Torin 2 (Biomarin). After 3 hours at 37°C in RPMI medium cells were washed, fixed, and stained with DAPI. For the acquisition of the images, ten pictures per each well of the 384-well plate were taken by using confocal automated microscopy (Opera high content system, Perkin Elmer). A dedicated script was developed to perform the analysis of TFEB localization on the different images (Acapella software, Perkin Elmer). The script calculates the ratio value resulting from the average intensity of nuclear TFEB-GFP fluorescence divided by the average of the cytosolic intensity of TFEB-GFP fluorescence. The results were normalized using negative (RPMI medium) and positive (HBSS starvation) control samples in the same plate. The data are represented by the percentage of nuclear translocation at the different concentrations of each compound using Prism software (GraphPad software). The EC50 for each compound was calculated using non-linear regression fitting (Prism software).

Results
TFEB Induces Autophagy
[0063] (Macro)autophagy is an evolutionarily conserved mechanism that targets intracytoplasmic material to lysosomes, thus providing energy supply during nutrient starvation (1, 3). Autophagy activation during starvation is regulated by mTOR, whose activity is dependent on cellular energy needs. [0064] As autophagy is the result of a tight partnership between autophagosomes and lysosomes (1), applicants tested whether TFEB, a transcription factor that controls lysosomal biogenesis, regulated autophagy. As TFEB exerts a positive control on lysosomal biogenesis and function (2) and on lysosomal exocytosis (FIGS. 16 and 17), one would expect that TFEB overexpression should decrease the number of autophagosomes due to their increased degradation by the lysosomes. Surprisingly, stable TFEB overexpression in HeLa cells increased significantly the number of autophagosomes, as determined by using the LC3 marker, which specifically associates with autophagosomes (4-7) (FIGS. 1a,b). Similar data were obtained by transient overexpression of TFEB in HeLa and Cos cells (FIG. 5). An increase in the number of autophagosomes was also detected by electron microscopy on mouse embryonic fibroblast (MEFs) infected with a lentivirus overexpressing TFEB (FIG. 6). This increase persisted in cells treated with lysosomal inhibitors of autophagosome/LC3II degradation bafilomycin and pepstatin/E64 (8), indicating that TFEB activates the formation of autophagosomes (FIG. 1a and FIG. 7). Nutrient starvation did not further increase the number of autophagosomes in TFEB-overexpressing cells (FIG. 1a,c), suggesting a saturating effect of TFEB overexpression on autophagy and raising the possibility that TFEB may be an important mediator of starvation-induced autophagy.

[0065] Consistent with these findings, RNA interference (RNAi) of TFEB in HeLa cells resulted in decreased levels of LC3II both in normal and starved conditions, either in the
presence or absence of bafilomycin (FIGS. 1d-f). Notably, the decrease of LC3II correlated with the levels of TFEB down-regulation achieved by the different RNAi oligos, demonstrating the specificity of the assay (FIG. 1g). These gain and loss of function data suggest that the biogenesis of autophagosomes and lysosomes are co-regulated by TFEB. Applicants next measured the rate of delivery of autophagosome to lysosome using an RFP-GFP tandem tagged LC3 protein (9), which discriminates early autophagic organelles (GFP-positive/mRFP-negative) from acidified autolysosomes (GFP-negative/mRFP-positive), as the GFP signal (but not the mRFP) is quenched inside acidic compartments (9). They found that the number of autophagolysosomes was higher in TFEB overexpressing cells compared to control cells, indicating that TFEB promotes autophagosome-lysosome fusion, thus enhancing the autophagic flux (FIG. 1h). Functional evidence of TFEB role in the regulation of autophagy came from the observation that degradation of long-lived proteins was enhanced by TFEB overexpression, and reduced by TFEB knock-down. This enhancement was abolished by the autophagy inhibitor 3-methyl adenine (3-MA) (10) (FIG. 8).

[0066] To test whether TFEB regulated the expression of autophagy genes, applicants analyzed the mRNA levels of a group of 51 genes reported to be involved in several steps of the autophagic pathway (1, 12, 13). They observed that the enhancement of the expression levels of autophagy genes in cells overexpressing TFEB was very similar to the one obtained during starvation (HeLa cells 4 h in EBSS media) (Pearson correlation: r value =-0.42; p value =0.001), while they were downregulated after TFEB silencing (FIG. 2a and Tables 1 and 2). Among them the expression of UVRAG, WIPI, MAP1LC3B, SQSTM1, VPS11, VPS18 and ATG9B was most significantly affected by TFEB overexpression (Tables 1 and 2). These genes are known to play a role in different steps of autophagy and appeared to be direct targets of TFEB, as they carry at least one CLEAR site (2) in their promoters (FIG. 9). Interestingly, VPS11, VPS18 and UVRAG play roles in autophagosome delivery to lysosomes (14), consistent with the observation of a significant enhancement of lysosome-autophagosome fusion in cells overexpressing TFEB.

[0067] These data indicate that TFEB is involved in the transcriptional regulation of starvation-induced autophagy. This conclusion is strongly bolstered by the following observations. First, the luciferase reporter assay (2) showed that starvation enhanced the effects of TFEB on target gene transcription (FIG. 10). Second, the expression of TFEB direct targets was upregulated in starved cells and this upregulation was inhibited by TFEB silencing (FIGS. 2a-c).

Starvation Regulates TFEB Nuclear Translocation and Activity

[0068] To identify the mechanism of starvation-induced activation of TFEB applicants analyzed its subcellular localization and post-translational modifications in starved cells. In normal conditions TFEB is localized to the cytoplasm (2). They observed that nutrient starvation (EBSS media) rapidly induced TFEB nuclear translocation (FIGS. 2d,e), and that cytosolic TFEB from starved cells appeared to have a lower molecular weight compared to that of normally fed cells, as revealed by western blot analysis (FIG. 11a). This molecular weight shift occurred rapidly but transiently and was abolished within 1 h after re-adding normal media to starved cells, concomitant with a decrease of nuclear TFEB (FIG. 11a). By supplementing EBSS media either with serum, amino acids or growth factors (i.e. insulin or EGF) applicants observed a significant inhibition of TFEB nuclear translocation compared to starved media alone (FIG. 2f). Almost no effect was observed when EBSS was supplemented with cytokines (i.e. INF or LIF) (FIG. 2f), suggesting that activation of TFEB is a process regulated by a signaling mechanism, which is sensitive to nutrient and growth factors. Applicants stimulated starved cells with normal medium supplemented with drugs inhibiting the mTOR (Rapamycin), PI3K-AKT (Tricirbin) and MEK (U0126) kinases. MEK inhibition resulted in TFEB nuclear localization, at level similar to starvation, while AKT and mTOR inhibition had no effect (FIG. 2g and FIG. 11b). These data suggest that TFEB activity is regulated by MAP kinase, uncovering an unexpected role of this signaling pathway in the regulation of starvation-induced autophagy. Furthermore, the expression of a constitutively active MEK (caMEK) in HeLa cells resulted in downregulation of TFEB target gene expression during starvation, thus mimicking the effect of TFEB knockdown (FIG. 2h), while caMEK overexpression in TFEB-depleted cells had no effect on the expression of TFEB target genes (FIG. 2h).

Serine Phosphorylation Regulates TFEB Activation

[0069] To analyze more in detail the relationship between MAPK signaling and TFEB applicants performed a mass-spectrometry analysis and identified at least three serines (i.e. S142, S332, and S402) that were phosphorylated in nutrient rich medium but not in starved medium. They mutated each of these three serines to alanines to abolish phosphorylation. Mutant TFEB proteins were individually expressed into HeLa cells and TFEB nuclear translocation analyzed. The TFEB(S142A) mutant showed a significantly increased nuclear localization compared to TFEB WT, TFEB(S332A) and TFEB(S402A) (FIG. 3a and FIG. 12a). Conversely the phospho-mimetic mutant (TFEB(S142D)) was unable to translocate into the nucleus upon nutrient starvation (FIG. 12b). The S142A TFEB mutant migrates at lower molecular weight in normal but not in starved medium, while the S142D mutant displayed a reduced shift during starvation compared to WT TFEB (FIG. 12c,d), further demonstrating that S142 is phosphorylated in normal but not in starved media. The expression of TFEB(S142A) resulted in increased expression levels of TFEB target genes compared to TFEB WT, TFEB(S332A) and TFEB(S402A) (FIG. 3b). Consistently, TFEB(S142A) caused a stronger induction of the autophagic/lysosomal system, compared to wt TFEB, as demonstrated by the increased number of autophagosomes (FIG. 3c and FIG. 13), lysosomes (FIG. 3d) and autophagolysosomes (FIG. 3e). Thus, TFEB nuclear translocation and activation are regulated by the phosphorylation of serine 142.

[0070] To identify the specific kinase responsible for the phosphorylation of serine 142, applicants performed bioinformatic analyses using methods that are based on computational models built upon a set of experimentally validated phosphorylation sites (15-19) (see methods for details). Consistently with previous results, they identified the serine-specific Extracellular Regulated Kinases (ERKs) as the top-ranking candidates for the phosphorylation of serine 142 (Table 3). Interestingly, serine 142 is highly conserved in other members of the H.L.H-leucine zipper gene family, such as the Microphthalmia Transcription Factor (MITF) (FIG. 14), where it was found to be phosphorylated by ERK2 (20). Further evidence of ERK2-mediated TFEB phosphorylation
came from ERK2-TFEB co-immunoprecipitation (FIG. 3) in normal but not in starved media. Furthermore siRNA-mediated knock-down of ERK1/2 proteins induced TFEB nuclear translocation to a similar extent as nutrient starvation (FIG. 3).

In Vivo Analysis of TFEB-Mediated Induction of Autophagy

Applicants analyzed the physiological relevance of TFEB-mediated control of the lysosomal/autophagic pathway in vivo in GFP-LC3 transgenic mice (11). They focused studies on the liver, due to the reported autophagic response observed in liver upon nutrient depletion. In liver, the number of GFP-positive vesicles started to increase after 24 hrs of fasting, and peaked at 48 hrs (see mat and methods for 48 h starvation protocol) (FIG. 4a), while the transcriptional induction of both autophagic and lysosomal TFEB target genes was evident after 16 hrs of fasting (FIG. 4b). Therefore, transcriptional activation precedes autophagosome formation in vivo. Importantly, at 16 hrs of fasting the sub-cellular localization of TFEB was completely nuclear (FIGS. 4c, d) and the level of ERK phosphorylation was reduced compared to fed animals (FIG. 4e), indicating that starvation regulates TFEB activity in vivo, similarly to what was observed in cultured cells.

Applicants evaluated if TFEB was sufficient to induce autophagy in vivo using both viral- and transgene-mediated TFEB overexpression. GFP-LC3 transgenic mice (11) were injected systemically with an adenovirus containing the murine Tcefb cDNA tagged with an HA epitope (AAV 2/9-Tcefb-HA) (FIGS. 15a, b). Liver specimens from Tcefb-injected animals showed a significant increase in the number of GFP positive vesicles, and this increase was further enhanced by starvation (FIGS. 4e, f). In addition, liver samples from conditional Tcefb-3xFLAG transgenic mice, in which transgene expression is driven by a liver-specific CRE recombinase (i.e. Albumin-CRE) (FIG. 15c), displayed a significant increase in the expression of lysosomal and autophagic genes and in the number of autophagosomes compared to control littermates (FIGS. 4g, h). Together, these data point to an important role of TFEB in the transcriptional regulation of starvation-induced autophagy.

TORC1 Regulates TFEB Subcellular Localization

TFEB subcellular localization was then analysed in HeLa and HEK-293T cells transiently transfected with a TFEB-3xFLAG plasmid and treated overnight with inhibitors of lysosomal function. These treatments included the use of chloroquine, an inhibitor of the lysosomal pH gradient, and Salicylhydroxilamide A (Sa1A) a selective inhibitor of the v-ATPase (38). Immunoblotting performed after nuclear/cytoplasmic fractionation revealed that also lysosomal stress induced nuclear translocation of exogenously expressed TFEB and that again TFEB nuclear accumulation was associated with a shift of TFEB-3xFLAG to a lower molecular weight, suggesting that lysosomal stress may affect TFEB phosphorylation status (FIG. 18).

Based on the observation that mTORC1 resides on the lysosomal membrane and its activity is dependent on both nutrient and lysosomal function (39, 40), applicants postulated that the effects of lysosomal stress on TFEB nuclear translocation may be mediated by mTORC1. Consistent with this idea, chloroquine or Sa1A inhibited mTORC1 activity as measured by level of p-P70S6K, a known mTORC1 substrate (FIG. 19A), (40). The involvement of mTOR appears in contrast with our previous observation that Rapamycin, a known mTOR inhibitor, did not affect TFEB activity. However, recent data indicate that Rapamycin is a partial inhibitor of mTOR, as some substrates are still efficiently phosphorylated in the presence of this drug (41). Therefore, applicants used kinase inhibitors Torin 1 and Torin 2, which belong to a novel class of molecules that target the mTOR catalytic site, thereby completely inhibiting mTOR activity (41, 47, 48).

Applicants stimulated starved cells, in which TFEB is dephosphorylated and localized to the nucleus, with an amino-acid rich medium supplemented with Torin 1 (250 nM), Rapamycin (2.5 μM) or ERK inhibitor U0126 (50 μM). Stimulation of starved cells with nutrients alone induced a significant TFEB molecular weight shift and re-localization to the cytoplasm (FIG. 19B). Nutrient stimulation in the presence of the ERK inhibitor U0126 at a concentration of 50 μM induced only a partial TFEB molecular weight shift, suggesting that phosphorylation by ERK partially contributes to TFEB cytoplasmic localization. Treatment with 2.5 μM Rapamycin also resulted in a partial molecular weight shift but did not affect TFEB subcellular localization (FIG. 19B). However, Torin 1 (250 nM) treatment entirely prevented the molecular weight shift induced by nutrients and, in turn, resulted in massive TFEB nuclear accumulation. These data were confirmed in a cell-based high content assay using stable HeLa cells overexpressing TFEB fused to the green fluorescent protein (TFEB-GFP). In the assay imaging of treated cells is acquired by an automated confocal microscope (OPERA system) and the analysis of those images with Acapella image software calculates the ratio of the average of fluorescence intensity of TFEB-GFP between the cytosol and nucleus of the cell (see Materials and methods for details) (FIGS. 19 C and D). As Torin 1 inhibits both mTORC1 and mTORC2 complexes, applicants next evaluated the contribution of each complex to TFEB regulation. Three main observations suggest that TFEB is predominantly regulated by mTORC1: (1) stimulation of starved cells with amino acids, which activate mTORC1 but not mTORC2, induced an extensive TFEB molecular weight shift, which is highly suggestive of a phosphorylation event (FIG. 19E); (2) knockdown of RagC and RagD, which mediate amino-acid signals to mTORC1, caused TFEB nuclear accumulation even in cells kept in full nutrient medium (FIG. 19F); (3) in cells with disrupted mTORC2 signalling (Sin 1−/− mouse embryonic fibroblasts (MEFs)) (49, 50, 46) TFEB underwent a molecular weight shift and nuclear translocation upon Torin 1 treatment that were similar to control cells (FIG. 19G).

mTORC1 Controls TFEB Subcellular Localization Via the Phosphorylation of S142

To test whether mTORC1 phosphorylates TFEB at S142, applicants generated a phosphospecific antibody that recognizes TFEB only when phosphorylated at S142. Using this antibody, applicants authors observed that TFEB was no longer phosphorylated at S142 in HeLa cells stably overexpressing TFEB-3xFLAG and cultured in nutrient-depleted media, consistent with applicants’ results above reported (FIG. 20A).

Subsequently, they analysed the levels of S142 phosphorylation in starved cells supplemented with normal media with or without either Torin 1 or Rapamycin. While Torin 1 clearly blunted nutrient-induced S142 phosphorylation, rapamycin did not, suggesting that S142 represents a
rapamycin-resistant mTORC1 site (FIG. 20A). These results clearly demonstrate that TFEB is an mTOR substrate and that S142 is a key residue for the phosphorylation of TFEB also by mTOR.

[0078] Recent findings suggest that mTORC1 phosphorylates its target proteins at multiple sites (42, 43, 44). To identify additional serine residues that may be phosphorylated by mTOR, applicants searched for consensus phosphoacceptor motif for mTORC1 (42) in the coding sequence of TFEB (FIGS. 20 B and C). They mutagenized all TFEB amino-acid residues that were putative mTORC1 targets into alanines. Then they tested the effects of each of these mutations on TFEB subcellular localization and found that, similarly to S142A, a serine-to-alanine mutation at position 211 (S211A), resulted in a constitutive nuclear localization of TFEB (FIG. 20 D). Mutations of the other serine residues behaved similarly to the wild-type TFEB (FIG. 20D).

[0079] Together, these data indicate that, other than S142, S211 also plays a role in TFEB subcellular localization and suggest that S211 represents an additional target site of mTORC1.

The Lysosome Regulates Gene Expression in TFEB

[0080] As the interaction of TFEB with mTORC1 controls TFEB nuclear translocation, applicants tested whether the ability of TFEB to regulate gene expression was also influenced by this interaction. The expression of several lysosomal/autophagic genes that were shown to be targets of TFEB (37) was tested in primary hepatocytes from a conditional knockout mouse line in which TFEB was deleted in the liver (Tfedeflox/flox; alb-CRE), and in a control mouse line (Tfedeflox/lox). Cells were treated with either chloroquine or Torin 1, or left untreated. These treatments inhibited mTOR as measured by the level of p-S6K or ERK were unaffected (FIG. 21A). Primary hepatocytes isolated from TFEB conditional knockout mice cultured in regular medium did not show significant differences in the expression levels of several TFEB target genes compared with control hepatocytes. However, while the expression of TFEB target genes was upregulated in hepatocytes from control mice after treatment with chloroquine, this upregulation was significantly blunted in hepatocytes from TFEB conditional knockout mice (FIG. 21B). Similarly, the transcriptional response upon Torin 1 treatment was significantly reduced in hepatocytes from TFEB conditional knockout mice (FIG. 21C). Together, these results indicate that TFEB plays a key role in the transcriptional response induced by the lysosome via mTOR.

[0081] Both transcriptional-dependent (24, 25) and independent mechanisms regulating autophagy have been described (26, 27). The study identifies novel, kinase-dependent, regulatory circuits that control multiple crucial steps of the autophagic pathway such as autophagosome formation, autophagosome-lysosome fusion and lysosome-mediated degradation of the autophagosomal content. Interestingly, applicants observed that the transcriptional induction of the autophagic/lysosomal genes precedes autophagosome formation. It could be envisaged that such transcriptional-dependent mechanism ensures a more prolonged and sustained activation of autophagy.

[0082] Autophagy dysfunction has been linked to several genetic disorders (28-30), by contrary previous studies showed that enhancement of autophagy has a therapeutic effect in animal models of neurodegenerative diseases and hepatic fibrosis (29, 31, 32).

[0083] The discovery of a novel mechanism that controls, at the transcriptional level, the lysosomal-autophagic pathway suggests novel approaches to modulate cellular clearance in these diseases. Furthermore, it provides a spin-off for therapeutic approaches based on lysosomal enzymes, suggesting new strategies for increasing the productivity of cell lines producing endogenous or recombinant lysosomal enzymes (FIGS. 16 and 17). Moreover, TFEB overexpression was able to promote substrate clearance and to rescue cellular vacuolization in LSDs (45); thus, the identification of a phosphorylation-mediated mechanism that regulates TFEB activity offers a new tool to promote cellular clearance in health and disease.

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Pearson product-moment correlation coefficient (PMCC) was obtained by comparing the gene expression profiles shown, i.e. TFEB stable overexpression vs. gene expression profiles of starved HeLa cells.

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**TABLE 4**

Prediction of S142 phosphorylation using different methods

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Results of the prediction of phosphorylation of S142 using five different methods. Methods are given in the first column. The second column indicates confidence score cutoff as described in methods, when available. The third column shows the actual format of prediction obtained by the corresponding method. The next four columns show the prediction in the kinase group, kinase family, kinase subfamily and kinase protein classifications, respectively.

REFERENCES


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Ann Met Ala Glu Leu Ala Gln Gin Val Val Lys Gin Glu Leu Pro Ser

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Glu Glu Gly Pro Gly Glu Ala Leu Met Leu Gly Ala Val Pro Asp Glu Pro Leu Pro Ala Leu Pro Pro Gin Ala Leu Pro Leu Pro
-continued

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acc cag cca cca tcc cca tcc cat cac ctg gac ttc agc cac aac ctg
Thr Gin Pro Pro Ser Pro Phe His His Leu Asp Phe Ser His Ser Leu
385   390   395   400

agc ttt ggg ggc agg gag gac ggt ccc cgc gac tcc ccc gag ccc
Ser Phe Gly Gly Arg Glu Asp Gly Glu Pro Pro Gly Tyr Pro Glu Pro
405   410   415

cgg ggc cgg cat gcc ttc cca ttc ccc agc ctg tcc aag aag gat
Leu Ala Pro Gly His Gly Ser Pro Phe Pro Ser Leu Ser Lys Lys Asp
420   425   430

cgg ggc cgg cat gcc ttc cca ttc ccc agc ctg tcc aag aag gat
Leu Ala Pro Gly His Gly Ser Pro Phe Pro Ser Leu Ser Lys Lys Asp
420   425   430

cgg agc agc ttc agt gag gac gag gag gac gag gag gac gag gag gac
Arg Ser Ser Phe Ser Met Glu Gly Asp Val Leu
465   470

<210> SEQ ID NO 2
<211> LENGTH: 476
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens

<400> SEQUENCE: 2
Met Ala Ser Arg Ile Gly Leu Arg Met Gin Leu Met Arg Glu Glu Ala
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Gln Gin Glu Glu Gin Arg Glu Arg Met Gin Gin Gin Ala Val Met His
20   25    30
Tyr Met Gin Gin Gin Gin Gin Gin Gin Gin Gin Leu Gly Gly Pro
35   40    45
Pro Thr Pro Ala Ile Asn Thr Pro Pro His Phe Gin Ser Pro Pro Pro
50   55    60
Val Pro Gly Glu Val Leu Lys Val Gin Ser Tyr Leu Glu Asp Pro Thr
65   70    75    80
Ser Tyr His Leu Gin Gin Gin Ser Gin Gin Gin Lys Val Arg Glu Tyr Leu
95   100   105   110
Ser Glu Thr Tyr Gly Asn Lys Phe Ala Ala His Ile Ser Pro Ala Gin
120   125
Gly Ser Pro Lys Pro Pro Pro Pro Pro Pro Arg Ala Gly
130   135   140
His Val Leu Ser Ser Ser Ala Gly Asn Ser Ala Pro Asn Ser Pro Met
145   150   155   160
 Ala Met Leu His Ile Gly Ser Asn Pro Glu Arg Glu Leu Asp Asp Val
165   170   175
Ile Asp Asn Ile Met Arg Leu Asp Asp Val Leu Gly Tyr Ile Asn Pro
180   185   190
Glu Met Gin Met Pro Asn Thr Leu Pro Leu Ser Ser Ser His Leu Asn
195  200  205
Val Tyr Ser Ser Asp Pro Gin Val Thr Ala Ser Leu Val Gly Val Thr
210  215  220
Ser Ser Cys Pro Ala Asp Leu Thr Gin Lys Arg Glu Leu Thr Asp
225  230  235  240
 Ala Glu Ser Arg Ala Leu Ala Lys Glu Arg Gin Lys Lys Asp Asn His
250  255  260
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<210> SEQ ID NO 3
<211> LENGTH: 150
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<222> OTHER INFORMATION: DNA encoding for a variant fragment of TPEB
<220> FEATURE:
<221> NAME/KEY: CDS
<222> LOCATION: (1)...(150)

<400> SEQUENCE: 3

ccc cca cca gcc gcc tcc cca ggg gtt gca gct gsa cac gtt ctc ccc
Pro Pro Ala Ala Ser Pro Gly Val Arg Ala Gly His Val Leu Ser
1 5 10 15

tcc tcc gct gcc aac agt gtt ccc aat gcc ccc atg gcc atg ccc
Ser Ser Ala Gly Asn Ser Ala Asn Pro Arg Ala Asn Leu His
20 25 30

att ggc tcc aac cct gag agg gag tgg gat gct att gag aac att
Ile Gly Ser Asn Pro Gly Arg Val Leu Asp Asp Val Ile Asn Ile
35 40 45

atg cgt
Met Arg
50
1. A nucleic acid comprising a coding sequence encoding for a transcription factor EB (TFEB) variant protein, (a) wherein the encoded TFEB variant protein comprises SEQ ID NO: 2, wherein Ser is replaced by a non-serine amino acid residue at positions 142 and/or 211 of SEQ ID NO: 2; or (b) wherein the encoded TFEB variant protein consists of amino acid residues 117 to 166 of SEQ ID NO: 2, wherein Ser is replaced by a non-serine amino acid residue at a position corresponding to position 142 of SEQ ID NO: 2; and wherein said variant protein induces autophagy.

2. The nucleic acid according to claim 1, wherein the non-serine amino acid residue of the encoded TFEB variant protein is Ala.

3. The nucleic acid according to claim 1, wherein the encoded TFEB variant protein consists of SEQ ID NO: 2, wherein Ser is replaced by a non-serine amino acid residue at amino acid residue positions 142 and/or 211.

4. The nucleic acid according to claim 1, wherein the encoded TFEB variant protein is according to claim 1, part (a).

5. The nucleic acid according to claim 1, wherein the encoded TFEB variant protein is according to claim 1, part (b).

6. The nucleic acid of claim 1 consisting of a coding sequence encoding for a transcription factor EB (TFEB) variant protein, (a) wherein the encoded TFEB variant protein comprises SEQ ID NO: 2, wherein Ser is replaced by a non-serine amino acid residue at positions 142 and/or 211 of SEQ ID NO: 2; or (b) wherein the encoded TFEB variant protein consists of amino acid residues 117 to 166 of SEQ ID NO: 2, wherein Ser is replaced by a non-serine amino acid residue at a position corresponding to position 142 of SEQ ID NO: 2; and wherein said variant protein induces autophagy.

7. The nucleic acid according to claim 6, wherein the non-serine amino acid residue of the encoded TFEB variant protein is Ala.

8. An expression vector comprising, under appropriate regulative sequences, the nucleic acid according to claim 1.

* * * * *

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**Protein Sequence:**

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Pro Pro Pro Ala Ala Ser Pro Gly Val Arg Ala Gly His Val Leu Ser
1  5  10  15
Ser Ser Ala Gly Asn Ser Ala Pro Asn Ala Pro Met Ala Met Leu His
20 25 30
Ile Gly Ser Asn Pro Glu Arg Glu Leu Asp Asp Val Ile Asp Asn Ile
35 40 45
Met Arg
50
```