A novel autoregulatory loop between the Gcn2-Atf4 pathway and L-Proline metabolism controls stem cell identity

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Increasing evidence indicates that metabolism is implicated in the control of stem cell identity. Here, we demonstrate that embryonic stem cell (ESC) behaviour relies on a feedback loop that involves the non-essential amino acid L-Proline (L-Pro) in the modulation of the Gcn2-Elf2α-Atf4 amino acid starvation response (AAR) pathway that in turn regulates L-Pro biosynthesis. This regulatory loop generates a highly specific intrinsic shortage of L-Pro that restricts proliferation of tightly packed domed-like ESC colonies and safeguards ESC identity. Indeed, alleviation of this nutrient stress condition by exogenously provided L-Pro induces proliferation and modifies the ESC phenotypic and molecular identity towards that of mesenchymal-like, invasive pluripotent stem cells. Either pharmacological inhibition of the prolyl-tRNA synthetase by halofuginone or forced expression of Atf4 antagonises the effects of exogenous L-Pro. Our data provide unprecedented evidence that L-Pro metabolism and the nutrient stress response are functionally integrated to maintain ESC identity.

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Naturally occurring amino acids are emerging as key players in the regulation of the phenotypic plasticity of stem cells. Indeed, exogenously provided threonine and methionine, two essential amino acids (EAAs), regulate self-renewal and differentiation of pluripotent stem cells. Moreover, exogenously provided L-Proline (L-Pro), a non-essential amino acid (NEAA), induces mouse ESCs towards an embryonic stem cell-to-mesenchymal-like transition (esMT) that converts compact, adherent ESCs into mesenchymal-like spindle-shaped, highly invasive and metastatic pluripotent stem cells. This fully reversible process resembles the epithelial-to-mesenchymal transition (EMT), which is essential for normal development and contributes to pathological cancer progression.

Interestingly, the Aldh18a1 gene is specifically induced in and marks the Primitive Endoderm (PrE) in the time window when the pluripotent epiblast precursors are specified within the inner cell mass (ICM) of the blastocyst. Since the Aldh18a1 enzyme catalyses the first and rate-limiting step of L-Pro biosynthesis, these findings suggest that L-Pro metabolism may regulate cell lineage segregation in early mammalian embryos. Despite its relevance, the molecular mechanisms underlying L-Pro control of stem cell identity remain largely unknown. This prompted us to investigate the early molecular events regulated by exogenously provided L-Pro in mouse ESCs.

Results

L-Pro modulates the AAR pathway. To provide insights into the earliest molecular events of L-Pro-induced embryonic stem cell-to-mesenchymal-like transition (esMT), we first analysed the transcriptome of ESCs grown at low density under feeder-free condition, at 24 and 48 h +/- L-Pro, in DMEM/FBS/LIF complete medium. Approximately 250 protein-coding genes were deregulated by L-Pro at 24 h (>1.5-fold-change, fdr < 0.0001), and this increased to approximately 900 genes at 48 h (Figures 1a and b; Supplementary Table 1). Gene ontology analysis revealed enrichment in genes involved in amino-acid metabolism at 24 h and in genes involved in focal adhesion and TGFβ signalling at 48 h (Figure 1c). Notably, the mesenchymal-like features became evident only later on, that is, at day 3 of the esMT. Among the genes early downregulated after L-Pro addition (Supplementary Table 1), we focused our attention on the stress-activated transcription factor 4 (Atf4). Interestingly, 77% (14/18) of the genes inhibited by L-Pro (>2-fold change at 24 h) (Supplementary Table 1) are direct targets of Atf4. Atf4 is the main downstream effector of an evolutionarily conserved stress pathway known as the amino acid starvation response (AAR) (Figure 1d), which is induced by uncharged tRNAs that bind to and activate the general stress-activated transcription factor 4 (Atf4).10

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Abbreviations: AAR, amino-acid starvation response pathway; Atf4, stress-activated transcription factor 4; EAAs, essential amino acids; NEAAs, nonessential amino acids; ESCs, embryonic stem cells; esMT, embryonic stem cell-to-mesenchymal-like transition; Elf2α, eukaryotic initiation factor 2; Gcn2, general control nonrepressed 2 protein kinase; HF, halofuginone; L-His, Histidine; ICM, inner cell mass; LC3, microtubule-associated protein light chain 3; L-Pro, L-Proline; PICS, proline-induced cells; PRS, prolyl-tRNA synthetase

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control nonrepressed 2 (Gcn2) protein kinase, leading to phosphorylation of the eukaryotic initiation factor 2 (Eif2α) and to translation of Atf4 mRNA.\textsuperscript{11,12} Accordingly, L-Pro downregulated a set of AAR/Atf4-related genes\textsuperscript{13} involved in non-essential amino acid (NEAA) biosynthesis, amino-acid transport or tRNA loading (Figure 1e). Remarkably, a similar set of genes was found to be upregulated in human T helper (TH17) cells treated with halofuginone (HF) (Figure 1e), a low-molecular weight alkaloid that induces L-Pro starvation by selectively inhibiting prollyl-tRNA synthetase (PRS).\textsuperscript{14,15}

Consistent with these findings, L-Pro and HF induced opposite effects on Eif2α phosphorylation and Atf4 protein levels (Figure 1f) and, remarkably, the effect of HF activity was fully counterbalanced by supplemental L-Pro (Figure 1f), suggesting that L-Pro availability regulates AAR in ESCs. We then assessed the specificity of L-Pro and showed that none of the NEAA other than L-Pro either reduced the expression of AAR markers (Figure 1g; Supplementary Figure 1a) or induced TGFβ-related genes (\textit{Lefty1}, \textit{Lefty2}) (Figure 1g; Supplementary Figure 1b). Moreover, none of them

**Figure 1** L-Pro rapidly remodels the ESC transcriptome and modulates the AAR pathway. (a) Genes deregulated (≥1.5-fold; FDR < 0.0001) by L-Pro (0.5 mM) in TBV2 ESCs at 24 and 48 h, as inferred from microarray analysis (Supplementary Table 1). (b) Time course of qPCR analysis of selected genes in L-Pro-treated versus untreated ESCs. Data are presented as fold change compared with control after normalisation to Gapdh. (c) Gene ontology (GO) analysis of L-Pro-treated ESC transcriptome showing gene enrichment in KEGG pathway. (d) Schematic representation of the AAR (Gcn2-Eif2α-ATF4) pathway. (e) Heat-map diagram showing AAR-related genes downregulated (green) in L-Pro-treated ESCs or upregulated (red) by histidinol (HisOH) and halofuginone (HF). (f) Western blot analysis of phospho-(51Ser)-Eif2α and Atf4 in ESCs treated (8 h) with L-Pro (0.5 mM) or HF (8 nM) either alone or in combination. Gapdh was used as a loading control. (g) Effects of different NEAAs on the expression of AAR-related and AAR-unrelated genes. qPCR analysis of \textit{Lefty1} and \textit{Slc1a4} in ESCs treated with individual NEAA (0.5 mM) or left untreated as a control (24 h). (h) Effects of different NEAAs on ESC proliferation (36 h). Proliferation was measured by the CyQuant\textsuperscript{R} assay and expressed as relative fluorescence units (RFU). (i) Effect of different NEAAs on the ESC to PIC transition. Colony-type frequency (domed versus flat) of ESCs +/- individual NEAA as assessed (~300 colonies scored/condition) at day 5 after plating. See Supplementary Figure 1. Data represent the mean ± S.D. from (b, f, g and i) three or (h) five independent experiments, *P < 0.001.
induced ESC proliferation (Figure 1h) or the ESC to PIC phenotypic transition (esMT) (Figure 1i; Supplementary Figure 1c). Finally, as expected by the fact that EEAAs are already present in the culture medium, a further addition of essential amino acids (EEAs), either as a mix or individually, did not induce ESC proliferation (data not shown). We thus concluded that l-Pro is the unique growth-limiting amino acid for ESCs cultured in complete medium (DMEM/FBS/LIF).

It is known that mitochondrial l-Pro catabolism generates reactive oxygen species (ROS), which, in turn, could activate redox-sensitive signalling pathways. We thus evaluated the effect of increasing concentrations of two antioxidants, N-Acetyl-Cysteine (NAC) and Glutathione (GSH) on l-Pro induced ESC proliferation and AAR inactivation. Neither NAC nor GSH counteracted l-Pro activity (Supplementary Figures 1d and e), even at the highest concentration (5 mM) thus ruling out the possibility that l-Pro induced ESC proliferation and AAR inactivation correlate with ROS accumulation.

Finally, we measured the intracellular concentration of free l-Pro and found that it was ~4-fold lower in ESCs compared with PiCs, which in turn was ~10-fold lower than in mouse embryonic fibroblasts (MEFs) (Supplementary Figure 1f). All together, our results indicate that ESCs grown in complete medium experience an intrinsic and specific l-Pro shortage (nutritional stress) that activates AAR, limits their proliferation and prevents the esMT, and that this stress is alleviated by exogenously provided l-Pro.

**Halofuginone antagonises l-Pro-induced proliferation and EsMT.** We have shown that HF prevented l-Pro-dependent AAR inactivation (Figure 1f). We therefore hypothesised that if l-Pro-induced ESC proliferation and esMT relied on AAR neutralization, HF should prevent both effects. Accordingly, HF (6–10 nM) antagonised the pro-proliferative effect of l-Pro (0.2–0.5 mM) in a dose-dependent manner (Figure 2a). In the absence of l-Pro, 2 nM HF inhibited ESC proliferation (Figure 2a). Consistent with this, HF also prevented l-Pro-induced downregulation of AAR-specific markers (Figure 2b; Supplementary Figure 2a). We then assessed the effect of Cycloheximide and Histidinol, which block protein synthesis by different mechanisms. Both inhibitors reduced ESC proliferation and induced AAR-specific markers in a dose-dependent manner (Supplementary Figures 2b and c). However, while exogenously added l-Pro fully counterbalanced the effect of HF (Figure 2c), it failed to antagonise the effect of Cycloheximide and Histidinol on both ESC proliferation and modulation of AAR-specific markers (Supplementary Figures 2b and c). Notably, none of the other NEAA neutralised the anti-proliferative effect of HF (Figure 2c), indicating that HF maintains a high specificity for l-Pro.

Furthermore, l-Pro/HF-treated ESCs developed Alkaline Phosphatase-positive (AP+) domed colonies (Figure 2h) and efficiently differentiated in vitro both in cardiomyocytes and in neurons (Figures 2i and j). Finally, HF reverted the esMT inducing the PIC to ESC phenotypic transition, even in the presence of supplemental l-Pro (Supplementary Figure 2h). We thus concluded that HF counteracted l-Pro effects without affecting ESC pluripotency and suggest that the esMT relies on AAR neutralisation.

**L-Pro and AAR are mutually regulated defining a novel autoregulatory loop.** To develop mechanistic insights, we first generated and characterised shRNA-mediated Atf4 knockdown (KD) ESCs (≥80% silencing) (Supplementary Figures 3a and c). Atf4 KD downregulated AAR but not pluripotency-related genes (Supplementary Figure 3d), and it reduced ESC proliferation in the absence of l-Pro (Figure 3a), as described in amino acid-starved cells. Nevertheless, l-Pro, but none of the other NEAA, induced the proliferation of control and Atf4 KD cells at comparable levels (Figures 3a and b). We then evaluated the response of Atf4 KD ESCs to HF (Figure 3c) and found that Atf4 KD increased the susceptibility of ESCs to HF, even when added at sub-lethal concentrations (1–2 nM) (Figure 3d). Most remarkably, l-Pro fully antagonised HF, inducing the proliferation of Atf4 KD ESCs (Figure 3e; Supplementary Figure 3e). Thus, Atf4 is required to improve ESC proliferation/survival. We evaluated whether Atf4 downregulation could affect the esMT. Indeed, similar to the control, Atf4 KD ESCs developed flat colonies only in the presence of l-Pro (Figure 3e), indicating that Atf4 downregulation per se is not sufficient to induce the ESC to PIC transition. We then used a complementary gain-of-function (GOF) approach based on the ROSA-TET-OFF system to assess the effect of Atf4 overexpression. As expected, Atf4 overexpression (Figure 3g; Supplementary Figure 3f) upregulated AAR but not pluripotency markers (Supplementary Figure 3g), and it eventually induced ESC proliferation/survival (Figure 3h; Supplementary Figure 3h), thus leading to the formation of larger colonies even in the absence of supplemental l-Pro (Figure 3i; Supplementary Figure 3i). Remarkably, in the presence of l-Pro, Atf4 GOF ESCs (~Tet) developed domed, compartmented colonies lacking the typical crown of mesenchymal-like cells scattered around the colony core of (~Tet) control cells (Figure 3i; Supplementary Figure 3i). Accordingly, cell motility was impaired in Atf4 GOF cells (Figure 3j). We thus concluded that l-Pro-mediated AAR inactivation/Atf4 downregulation is crucial for the acquisition of the mesenchymal-like motile phenotype.
The opposing effects of Atf4 KD and GOF on ESC proliferation correlate well with the positive control of Atf4 on L-Pro biosynthesis. In mammalian cells, L-Pro is synthesized from L-Glu by a reductive process involving two enzymes, pyrroline-5-carboxylate synthetase and pyrroline-5-carboxylate reductase, encoded by Aldh18a1 and Pycr1, respectively. Indeed, both these genes were early downregulated after L-Pro addition (Figure 1e). Furthermore, HF induced...
Figure 3  Biological effects of L-Pro-dependent modulation of the AAR pathway in ESCs. (a and b) Effect of Atf4 KD on ESC proliferation. FACS-based analysis (EdU incorporation) of Atf4 KD (shATF4) and control (shNT) ESCs proliferation (a) performed at 36 h after plating. The results are expressed as the fold change compared with control. Representative FACS plots of EdU incorporation (b) in Atf4 KD ESC +/- individual NEAAs (0.5 mM) at 36 h after plating. (c) Western blot analysis of Atf4 in untreated or HF-treated Atf4 KD and control ESCs (36 h). The densitometric analysis is expressed in ADU as the Atf4/Gadph ratio. (d) Proliferation of control and Atf4 KD ESCs after 48 h treatment with HF, either alone or with l-Pro, was measured by CyQuant and expressed as RFU. (e) Effect of Atf4 KD on cell colony formation. Representative photomicrographs (e) of colonies generated from Atf4 KD and control ESCs plated +/- l-Pro (0.2 mM) and stained with crystal violet at day 5. Scale bar, 200 μm. Quantification of crystal violet staining of cell colonies generated in the different culture conditions (f). (g-j) Effect of Atf4 overexpression on ESCs behaviour. Immunoblotting analysis of Atf4 protein (g) in Atf4 Tet-OFF ESCs +/- Tet at 48 h. The densitometric analysis is expressed in ADU as the Atf4/Gadph ratio. Representative FACS plots of EdU incorporation (h) in Atf4 Tet-OFF ESCs +/- Tet. Representative photomicrographs of colonies (i) generated from Atf4 Tet-OFF ESCs treated with +/- Tet and +/- l-Pro (0.2 mM) and stained with crystal violet at day 5 after plating. Scale bar, 50 μm. Effect of Atf4 overexpression on l-Pro-induced cell motility (j). Control (+Tet) and Atf4 GOF (-Tet) ESCs were treated with +/- l-Pro and migration was assessed at day 5 after plating. The results show the average numbers of cells migrating towards an FBS gradient (1–15%). (k) qPCR analysis of Aldh18a1 and Pycr1 in control (+Tet) and Atf4 GOF (-Tet) ESCs at 24 h and 48 h. (l) The intracellular free L-Pro concentration was measured in control (+Tet) and Atf4 GOF (-Tet) ESCs at day 5 after plating. The results are expressed as the fold change compared with control. (m) Schematic representation of the l-Pro-AAR/Atf4 regulatory feedback loop. Data represent the mean ± S.D. from (a, b, c, f, g, h, j, k, and l) three or (d) five independent experiments. *P<0.001.
**L-Pro starvation induces autophagy in ESCs.** In mammals, amino-acid starvation induces autophagy, a mechanism by which the cells digest their own proteins and organelles. Because ESCs are starved of L-Pro, we evaluated autophagy induction. We first analysed the intracellular distribution of the microtubule-associated protein light chain 3 (LC3), which shifts from diffuse to punctate during autophagosome formation. ESCs stably expressing light chain 3 (LC3), which shifts from diffuse to punctate evaluated autophagy induction. We first analysed the LC3-GFP signal (Supplementary Figure 4c and d), thus indicating that L-Pro starvation in ESCs resembles that of EAA, we used the Histidinol that mimics Histidine starvation and induces autophagy. Interestingly, either Vitamin C or 2i (PD0325901/CHIR99021) inhibitors counteracted the L-Pro-induced apoptosis in cancer cells. We thus reasoned that the L-Pro-dependent downregulation of Atf4 (Figures 1 and 3) could at least partly explain apoptosis during the esMT. Consistent with our hypothesis, apoptotic DNA fragmentation significantly increased in L-Pro-treated Atf4 KD ESCs compared with control (Supplementary Figure 5d, left panel). Moreover, Atf4 overexpression counteracted the pro-apoptotic effect but not the pro-proliferative effect of L-Pro, and reduced autophagy in esMT (Figure 3i and Supplementary Figure 5d, right panel, and Supplementary Figure 5e). Therefore, we assessed the functional relevance of these findings by blocking apoptosis pharmacologically. The pan-caspase inhibitor Z-VAD reduced DNA fragmentation and further increased the survival of L-Pro-treated cells (Figures 5f–h). Remarkably, L-Pro/Z-VAD-treated ESCs generated flat colonies, which showed tight cell–cell contacts rather than the crown of mesenchymal-like cells typical of control colonies (Supplementary Figure 5i). Accordingly, cell migration was significantly impaired in L-Pro/Z-VAD-treated cells and was comparable to that of control ESCs (Figure 5i), suggesting that L-Pro-induced apoptosis is crucial for the acquisition of the motile mesenchymal-like phenotype. Interestingly, either Vitamin C or 2i (PD0325901/CHIR99021) inhibitors counteracted the L-Pro-induced esMT (Supplementary Figure 5g), blocked DNA fragmentation (Supplementary Figure 5h) and increased cell survival (Supplementary Figure 5i). Together, our data provide evidence of a functional link between L-Pro-mediated Atf4 downregulation and the acquisition of the motile mesenchymal-like phenotype in the esMT, which is accompanied by apoptosis induction (Figure 6).
Discussion

Pluripotent stem cells share the tendency to self-renew in vitro as tightly compacted cell aggregates (domed- or spherical-shaped colonies), thus recapitulating their in vivo behaviour. Here, we show that in mouse ESCs this phenotypic feature depends on a highly specific and controlled shortage of the amino acid l-Proline (l-Pro). Indeed, ESCs although cultured in complete rich medium (DMEM/FBS/LIF) experience a specific l-Pro starvation. However, this intrinsic l-Pro shortage is not as extreme as to compromise ESC survival, but it is sufficient to limit their proliferation and influence the gene expression profile. In most cell lines, including ESCs, a condition of amino-acid limitation leads to

Figure 4 l-Pro regulates autophagy in ESCs and is counteracted by HF. (a) Time-course analysis of LC3 puncta formation in ESCs. Representative fluorescent photomicrographs (left panel) of LC3-GFP wild-type (LC3_wt) or GFP-LC3G120A mutant (LC3_mut) ESCs plated on gelatin +/- l-Pro (0.2 mM). Scale bar, 25 μm. The number of cells with punctate LC3-GFP (right panel) was measured as the percentage (~400 cell scored/condition) of the total number of cells. (b and c) l-Pro supplementation prevents LC3-GFP re-localisation in a specific and dose-dependent manner. Punctate LC3-GFP ESCs were quantified after 36 h of treatment with increasing l-Pro concentrations (b) or with NEAAs (c). (d) Representative fluorescent photomicrographs (left panels) of LC3-GFP localisation in control and ESCs treated with l-Pro (0.2 mM) either alone or with HF (6 and 8 nM). The quantification was performed 36 h after plating (right panel). (e) Representative microphotographs of acidic vesicular organelles (AVO) stained with toluidine blue dye (left panels, red arrows) and red-fluorescent acidic dye (Lysotracker Red, right panels, yellow arrows) in control (- l-Pro; upper panel) and l-Pro (0.2 mM; lower panel)-treated ESCs at day 5 of the esMT. Representative confocal image of autophagolysosomes. Scale bar, 25 μm. (f) Transmission electron microscopy (TEM) analysis of control and l-Pro (0.2 mM)-treated ESCs at day 5 after plating. Representative images of the electron-dense bodies delimited by the double membrane, with a typical narrow empty (electron lucent) space between the two sheets (orange arrowheads). Data represent the mean ± S.D. from (a, b, c, and d) three independent experiments, *P<0.001, **P<0.05

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the induction of the amino-acid starvation response (AAR/Atf4) pathway and autophagy, and, if protracted and uncompensated, this condition leads to apoptotic cell death. Accordingly, AAR and autophagy, but not apoptosis, are progressively induced in feeder-free growing ESCs and are specifically counteracted by exogenously added L-Pro in a dose-dependent manner. Moreover, L-Pro starvation exerts similar effects to that of pharmacological-induced starvation of an essential amino acid (EAA), such as Histidine (L-His). It is known that EAAs cannot be synthesised by the cells and must be exogenously provided; conversely, NEAAs, such as L-Pro, can be produced by the cells, either de novo or from other amino acids. Thus, the finding that an intrinsic limitation of L-Pro occurs in ESCs cultured in complete rich medium was totally unexpected. Interestingly, although it was not specifically discussed, a previous report showed that only the deprivation of L-Pro among all the NEAAs limits the in vitro colony formation ability of E14Tg2A ESCs. Of note, mouse fibroblasts release L-Pro, which likely explain why feeder cells protect ESCs from this nutrient starvation response.

**Figure 5** Apoptotic cell death occurs during L-Pro-induced esMT. (a) Representative photomicrographs of control and L-Pro (0.5 mM)-treated ESCs at day 5 after plating (left panel). Arrowheads indicate floating cells and debris. Scale bar, 100 μm. Quantification of dead cells by the trypan blue exclusion assay (right panel) in adherent and total (floating plus adherent) cell populations. (b) Representative frames of time-lapse series of L-Pro-treated ESCs. Arrowheads indicate cells located at the periphery of a flat colony exploding into a cascade of apoptotic-like bodies. Scale bar, 25 μm. (c) Representative fluorescent photomicrographs of Hoechst-stained nuclei. Apoptotic figures, including a pyknotic nucleus (py), nuclear fragmentation (nf) and apoptotic bodies (ab), were distinguished from mitotic figures, prophase (p), pro-metaphase (p-m), metaphase (m), anaphase (a) and telophase (t). Data show the percentage of apoptotic nuclei (~900 nuclei scored/condition) in control and L-Pro-treated ESC cultures (day 5). (d) Quantitative determination of DNA laddering by a photometric immunoassay. (e) FACS analysis of Propidium Iodide (PI)-stained control or L-Pro-treated ESCs. The fraction of cells in SubG1 phase is indicated. (f–h) Effect of Z-VAD on L-Pro-induced esMT and cell motility. Quantitative analysis of DNA fragmentation (f) in control or L-Pro (0.2 mM)-treated ESCs either alone or in the presence of Z-VAD (0.1 mM). Representative photomicrographs of the colonies (g) showing round (ESC) or flat (PiC) phenotype. Scale bar, 200 μm. (h) Quantification of crystal violet staining of cell colonies generated in the different culture conditions. Effect of Z-VAD on cell motility (i). ESCs were treated with L-Pro +/- Z-VAD for 5 days or left untreated as a control, and cell migration was assessed and expressed as the average number of cells migrating towards an FBS gradient (1–15%). Data represent the mean ± S.D. from three or five independent experiments, *P < 0.001.
AAR/Atf4, and AAR/Atf4 under the control of L-Pro availability, Aldh18a1 and Pycr1 between AAR/Atf4 and L-Pro biosynthetic pathway might be reduced/inefficient in ESCs. This finely tuned interplay protein collagens. The Atf4 ↔ Aldh18a1 and Pycr1 (pyrroline ring) structure it is synthesised by highly specific unique among the NEAAs because, due to its atypical cyclic identity. The ESC identity depends on a highly specific intrinsic nutrient stress. Alleviation of this stress (L-Pro fullness/AAR inactivation) initially increased L-Pro intake. It suggests that they may be at least partially rescued by approaches) or pharmacologically (HF, Z-VAD) modulates esMT cells. Targeting of the L-Pro ↔ AAR/Atf4 axis genetically (AAR/LOF and GOF approaches) or pharmacologically (HF, Z-VAD) modulates esMT How L-Pro shortage stress is generated in ESCs? L-Pro is unique among the NEAAs because, due to its atypical cyclic (pyrroline ring) structure it is synthesised by highly specific enzymes encoded by Aldh18a1 and Pycr1 genes.35 The generation of Atf4 GOF and LOF cell lines allowed us to identify a L-Pro-AAR/Atf4-Aldh18a1/Pycr1 autoregulatory loop in ESCs. Indeed, upon Atf4 overexpression Aldh18a1/Pycr1 expression increases and ESCs are less dependent on exogenous L-Pro; conversely, Atf4 knockdown reduces Aldh18a1/Pycr1 expression and increases ESC requirement for L-Pro to proliferate. Thus, by placing the expression of L-Pro biosynthesis genes (Aldh18a1 and Pycr1) under the control of AAR/Atf4, and AAR/Atf4 under the control of L-Pro availability, this loop maintains L-Pro as a growth limiting metabolite in ESCs. Notably, L-Glu and L-Orn, two metabolic precursors of L-Pro, when added to the culture medium are less efficient than L-Pro in (i) inactivating the AAR and (ii) inducing ESC proliferation, but later on it induces the esMT, which is accompanied by a huge induction of macroautophagy and apoptosis. The esMT, ESCs acquire mesenchymal-like features, becoming highly motile and invasive pluripotent stem cells. Targeting of the L-Pro ↔ AAR/Atf4 axis genetically (Atf4 LOF and GOF approaches) or pharmacologically (HF, Z-VAD) modulates esMT...
Briefly, ESCs were seeded at low density (50–500 cells/cm²) onto gelatin-coated plates and grown in the presence of L-Pro (500 μM–1 mM) for 5 days. Medium was changed at day 3 with addition of fresh L-Pro. PiCs were harvested using accutase (Sigma-Aldrich, St. Louis, MO, USA) and cultured in the presence of L-Pro (500 μM). Both ESCs and PiCs were cultured in high glucose Dulbecco's modified Eagle's medium (Invitrogen, Life Technologies, Eugene, OR, USA) supplemented with 15% ES-screened fetal bovine serum (FBS, Euroclone, Milan, Italy), 0.1 mM β-mercaptoethanol (Sigma-Aldrich), 1 mM sodium pyruvate, 2 mM glutamine, 100 U/ml penicillin/streptomycin all from GIBCO (Carlsbad, CA, USA) and 100 U/ml recombinant LIF (ESGRO, Millipore, Darmstadt, Germany).

For Halofugone (HF) treatments, HF (Sigma-Aldrich) was dissolved in DMSO (2 mM) and used at the indicated concentrations (1–20 mM). Amino acids including L-Pro, NEAs (l-Gly, l-Ala, l-Ser, l-Asp, l-Glu and l-Asn) and l-Orn were purchased from Sigma-Aldrich. Cycloheximide, Histrindin, NAC and GSH (Sigma-Aldrich) were dissolved in water and used at the indicated concentrations (0.15–0.6 μg/ml; 0.6–2.4 mM, 0.625–5 mM and 0.04–5 mM, respectively).

For cardiac differentiation, E7SCs were induced to form embryoid bodies (EBs) in hanging drops (500 cells/drop) and were further differentiated without addition of growth factors, as described.46 Neuronal differentiation was performed as previously described.46

Generation of Atf4 KD and GOF ES cell lines. To generate Atf4 KD ESCs, 1 x 10^6 undifferentiated TBV2 ESCs were infected with either non-targeting control or Atf4 shRNA lentiviruses (pLKO1 vectors, Open Biosystems, GE Dharmacon, Lafayette, CO, USA). Twenty-four hours after infection, cells were subjected to puromycin selection (Sigma-Aldrich, 2 μg/ml) for 2 days and subsequently grown for additional 4–5 days, until clones appeared. Clones were pooled together and silencing efficacy was evaluated at RNA and protein levels. Four independent shRNA viral vectors, targeting different Atf4 mRNA sequences, were used and one shRNA vector was selected on the basis of the highest silencing efficacy. Atf4 Tet-Off ESCs (Atf4 GOF) were generated as previously described.46 Briefly, mouse Atf4 cDNA (Origene; cat. n. MR29597) was cloned in the exchange vector pTHC/MCS247 and then targeted to the Rosa26 locus of the E83 ESCs, as described.48 A PCR-based assay on genomic DNA was used to identify the positive clones.19 Two independent clones were tested for Atf4 mRNA and protein induction upon tetracycline (Sigma-Aldrich, 1 μg/ml) removal.

L-Pro measurement. The intracellular concentration of free L-Pro in ESCs, PiCs and mouse embryonic fibroblasts (MEFs) was quantified as previously described.48

Colony assay, alkaline phosphatase and crystal violet staining. For colony phenotype assay, ESCs were plated at low density (50–250 cells/cm²) in complete medium (DMEM/15%FBS/LIF) on gelatin-coated plates and colonies were grown for 5 days in the presence/absence of L-Pro (250–500 μM), with a medium change at day 3.

Alkaline phosphatase activity was assessed using the AP staining kit (System Biosciences, Mountain View, CA, USA) following the manufacturer's instructions.

Staining with crystal violet was performed as previously described. Briefly, cells were washed twice with phosphate-buffered saline (PBS) and fixed/stained with a solution of 6% glacial acetic acid and crystal violet. After 30 min at RT, cells were carefully washed with tap water and dried for further analysis. For quantification, crystal violet was dissolved with 30% acetic acid (1 ml) for 15 min at RT and absorbance was read at 540 nm, using the Synergy H1 Microplate Reader (BioTek, Winooski, VT, USA).

Proliferation assays and viable cell count. For the cell proliferation assay, ESCs were plated at 1.5 x 10^5 cells/cm² on gelatin-coated plates in ESC medium with or without L-Pro and cell viability was measured at 36 h, using the colorimetric CyQUANT® cell proliferation assay (Invitrogen), following the manufacturer's instructions. Briefly, triplicate samples were washed with PBS and stored at −80 °C for 2 h. After thawing, cells were incubated with a mix of cell lysis buffer and CyQUANT® dye. Absorbance was analysed at 490–520 nm, using the Fluroskan Ascent FL Microplate Fluorometer and Luminometer (Thermo Fisher Scientific, Waltham, MA, USA).

For the proliferation assay, the Click-iT Edu Flow Cytometry Assay (Invitrogen) was used. Briefly, cells were incubated with 5-ethyl-1H-2-deoxyuridine (EdU) (10 μM; overnight at 37 °C), dissociated, fixed and permeabilized, following the manufacturer's instructions. Samples were analysed at FACSCanto using the DivaTM software (BD Biosciences, San Jose, CA, USA).

Viable count was performed on triplicate samples; briefly, cells were counted on the Cellometer Auto T4 (Nexcelom Bioscience, Lawrence, MA, USA) and % viability was simultaneously calculated by trypan blue exclusion.

Cell migration. Migration assays were performed using polycarbonate (5 μm pore, Costar, Tewksbury, MA, USA) or PET membrane transwells (8 μm pore; BD Biosciences). Cells were detached with accutase (Sigma-Aldrich) plated (2 x 10^5) transwell and incubated for 20–22 h as previously described.5

Time lapse experiments. ESCs were seeded on gelatin-coated plates at 50 cells/cm² and grown in complete medium supplemented with appropriate compounds. Images were captured (x20) every 5 min for 24 h using a Leica DMi6000B inverted microscope equipped with a microscope incubator (Oikolab, Naples, Italy).

Real-time PCR. Total RNAs were isolated using RNeasy mini kit (Qiagen, Hilden, Germany) and retro-transcribed using the QuantiTect Reverse Transcription kit (Qiagen). SYBR Green PCR master mix (FluoCycle II SYBR, Euroclone) was used for quantitative real-time PCR. Primers were in Supplementary Table 2. For microarray experiments, RNA samples were analysed at the microarray facility of the Ohio State University as previously described.4 Statistical data analysis was performed at the Bioinformatics Core Facility (http://bioinformatics.tigem.it) of TIGEM, Italy. The relative signal intensities of each probe across the six samples assayed for each cell line were evaluated selecting a false discovery rate (FDR) of < 0.001 to assess significant gene differential expressions. Genes were classified based on Gene Ontology terms (DAVID Bioinformatics Resources; http://david.abcc.ncifcrf.gov).

Immunoblotting. Whole-cell lysates were prepared either in 20 mM Tris pH 7.9, 120 mM KCl, 5 mM MgCl₂, 0.2% Nonidet P-40, 5 mM EDTA, 10% glycerol (for PARP1 and Casp3 analysis) or in 100 mM Tris pH 8, 140 mM NaCl, 20 mM EDTA, 0.2%, SDS, 1% Nonidet P-40 lysis buffer, resolved on SDS-PAGE gels and transferred onto PVDF membranes using the iBlot dry Transfer System (Life Technologies, Carlsbad, CA, USA). Blocked membranes were incubated with the following primary antibodies: Atf4 (1: 500); eFlx2 (1: 1000); phospho-eFlx2 (1: 500); PARP1 (1: 1000), Cell Signaling Technology, CST, Danvers, MA, USA), which detects the full-length protein (116 kDa) as well as the large caspase-cleaved fragment (89 kDa); anti-Casp3 (1: 1000) all from CST, and anti-LC3 (2 χg/ml, Novus Biologics, Littleton, CO, USA) anti-Gapdh (1: 10000, Dako, Glostrup, Denmark). Detection was performed with ECL reagents (Pierce, Thermo Scientific, Waltham, MA, USA). The ImageJ software (public source; http://imagej.nih.gov/) was used for densitometric quantification.

Preparation of cytospin samples. Cells (1–1.5 x 10^5) were dissociated with accutase for 5 min at 37 °C and resuspended in 15% FBS/1x PBS. Cell samples were centrifuged at 900 r.p.m. for 15 min onto glass slides (2 spots of 1 x 10^4 cells each) using a Thermo Shandon Cytocentrifuge (CytoSpin 4, Thermo Fisher Scientific). Specimens were directly analysed or fixed for further analysis.

Immunofluorescence analysis. Differentiated EBs and monolayer cultures were fixed with 4% paraformaldehyde (PFA) and permeabilized with 0.1% Triton X-100 at RT. The following primary antibodies were used: MF20 (1: 50), Developmental Studies Hybridoma Bank; βIII-Tubulin (1: 400, Sigma-Aldrich). After washing in 0.5% Tween-1x PBS, cells were incubated with secondary antibodies (1: 200, Alexa Fluor, Molecular Probes Inc., Life Technologies).

For autophagy detection, cytospin samples (see above) were stained with Lysotracker Red DND-99 dye. Briefly, cells (50 000 cells/slot) were washed with PBS 1 x, stained with Lysotracker Red DND-99 dye (Molecular Probes Inc.) for 1 min at room temperature. Cells were gently washed and fixed in 4% PFA for 30 min, washed three times with PBS, and stained with membrane stain WGA-Alexa Fluor 488 Conjugate (Invitrogen) following the manufacturer's instructions. Cell nuclei were counterstained with Hoechst 33342 (Invitrogen). Images were obtained using the DMi6000B microscope and the DFC 350FX BW digital camera (Leica, Solms, Germany). Leica FW4000 and AF6000 software were used for image acquisition/ elaboration. Confocal images were acquired at x63 magnification on a LSM710 confocal fluorescence microscope (Carl Zeiss Inc., Jena, Germany) using the ZEN 2008 software (Carl Zeiss Inc.).
Toluidine blue staining. Samples were fixed with 4% PFA, washed three times with PBS and stained with toluidine blue (1% in water) for 10 min at RT. The samples were observed using a DM6000B upright microscope and the LAS V.4.0 software (Leica) was used for images acquisition.

Nuclear morphology analysis. Cells (floating and adherent) were cytopsised (see above), fixed in 4% PFA and the nuclei were stained with Hoechst 33342 (Invitrogen) in 1 x PBS with 0.1% Triton X-100. Fluorescent labelling was visualised using the DM6000B microscope (Leica) and images acquired with the DFC 350FX B/W camera.

Apoptosis assays. DNA laddering associated with apoptotic cell death was quantified by spectrophotometric detection of cleaved DNA/histone complexes, using the Cell Death Detection ElisaPlus kit (Roche, Welwyn Garden City, UK), following the manufacturer’s instructions. The immunohasay was performed on duplicate samples by measuring at 405 nm against a blank solution (with a 490-nm reference wavelength).

For subG1 determination, floating and adherent sub-populations (at day 5) were collected, permabilized and stained in 0.1% Na citrate, 0.1% Triton X-100, 0.2 µg/ml propidium iodide (PI), 55 µg/ml DNease-free RNase.

Transmission electron microscopy. Cells (ESCs and PiCs) were fixed in 2.5% glutaraldehyde in cacodylate buffer, post-fixed in osmium tetra oxide, 2.5% glutaraldehyde in cacodylate buffer, post-fixed in osmium tetra oxide, and contrasted with uranyl acetate and lead citrate. Grids were imaged using a Tecnai F20 equipped with a Gatan imaging filter equipped with a Gatan Multiscan camera.

For detection of membrane asymmetry, intact cells were labelled with Alexa Fluor 488 Annexin V and PI using the Dead Cell Apoptosis Kit (Invitrogen) following the manufacturer’s instructions. Samples were analysed at 405 nm against a blank solution (with a 490-nm reference wavelength).

Statistical analysis. Statistical significance was determined by a two-tailed paired Student’s t-test. P-values < 0.01 were considered as statistically significant. Error bars show mean ± S.D. unless otherwise indicated.

Conflict of Interest

The authors declare no conflict of interest.

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