Metabolic labeling of RNA uncovers the contribution of transcription and decay rates on hypoxia-induced changes in RNA levels

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ABSTRACT

Cells adapt to environmental changes, including fluctuations in oxygen levels, through the induction of specific gene expression programs. However, most transcriptomic studies do not distinguish the relative contribution of transcription, RNA processing, and RNA degradation processes to cellular homeostasis. Here we used metabolic labeling followed by massive parallel sequencing of newly transcribed and preexisting RNA fractions to simultaneously analyze RNA synthesis and decay in primary endothelial cells exposed to low oxygen tension. We found that changes in transcription rates induced by hypoxia are the major determinant of changes in RNA levels. However, degradation rates also had a significant contribution, accounting for 24% of the observed variability in total mRNA. In addition, our results indicated that hypoxia led to a reduction of the overall mRNA stability from a median half-life in normoxia of 8.7 h, to 5.7 h in hypoxia. Analysis of RNA content per cell confirmed a decrease of both mRNA and total RNA in hypoxic samples and that this effect is dependent on the EGLN/HIF/TSC2 axis. This effect could potentially contribute to fundamental global responses such as inhibition of translation in hypoxia. In summary, our study provides a quantitative analysis of the contribution of RNA synthesis and stability to the transcriptional response to hypoxia and uncovers an unexpected effect on the latter.

Keywords: hypoxia; RNA decay; transcription; HIF

INTRODUCTION

Eukaryotic cells show a remarkable regulatory flexibility that allows them to adjust to a constantly varying environment. Modulation of gene expression, consequence of changes at the transcriptional and post-transcriptional levels, plays a central role in the adaptation to environmental changes and stresses. However, quantitative information about the relative contribution of transcription to the changes in gene expression in response to specific signals is still scarce.

The response to hypoxia, defined as the situation where oxygen consumption exceeds its supply, is a paradigmatic

example of how gene expression shapes cell physiology to cope with environmental fluctuations. The hypoxia inducible factors (HIFs) are transcription factors that play a fundamental role in this response. HIF is a heterodimeric transcription factor conserved in all metazoans (Iyer et al. 1998) that is composed of an oxygen-regulated alpha subunit (HIF α) and a constitutive expressed beta subunit (HIF β) also known as aryl hydrocarbon receptor nuclear translocator (ARNT). There are three genes encoding HIF α subunits: HIF1A, EPAS1 (also known as HIF2A), and HIF3A. All these proteins belong to the PER-ARNT-SIM (PAS) subfamily of the basic helix-loop-helix (bHLH) family of transcription

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factors and they have a similar domain organization. HIF α and HIF β subunits share an amino terminus bHLH domain that mediates the DNA binding and a central region or PAS domain that facilitates the heterodimerization (Jiang et al. 1997). Only the α subunits contain an oxygen dependent degradation domain (ODD) with proline residues (Pro 402/Pro 564 in HIF1A, Pro 405/Pro 531 in EPAS1, and pro 492 in HIF3A) that are essential for the oxygen-induced HIF stabilization. Furthermore, HIF1A and EPAS1 present two transactivation domains NTAD and CTAD that can bind transcriptional coregulators such as CBP/p300 and the cyclin dependent kinase 8 (CDK8) (Arany et al. 1996; Kasper et al. 2005; Galbraith et al. 2013; Choudhry et al. 2014; Dengler et al. 2014). The regulation of HIF α stability is mediated by a family of enzymes called EGL-nine homologs (EGLNs) that, in normoxic conditions, use molecular oxygen to hydroxylate specific proline residues in HIF α . The hydroxylation in one of these residues is sufficient for the recognition of HIF α by the von Hippel-Lindau tumor suppressor protein (pVHL) that is the substrate recognition subunit of an E3-ubiquitin ligase complex. Thus, pVHL binding leads to HIF α ubiquitination, targeting it for degradation via proteasome (Maxwell et al. 1999; Ivan et al. 2001; Jaakkola et al. 2001). In hypoxic conditions, HIF is stabilized and binds the RCGTG motif present in the regulatory regions of its targets genes and interacts with the transcriptional machinery to regulate their transcription.

The global gene expression pattern induced by hypoxia has been widely studied (Hu et al. 2003; Scheurer et al. 2004; Elvidge et al. 2006; Warnecke et al. 2008; Xia and Kung 2009; Mimura et al. 2012; Weigand et al. 2012; Salama et al. 2015; Galbraith et al. 2017; Chen et al. 2018; Moreau et al. 2018; Chakraborty et al. 2019; Smythies et al. 2019). However, these studies relied on RNA-seq and microarrays that detect mRNA steady state levels and therefore cannot distinguish between regulation of transcription and other mechanisms of post-transcriptional regulation. In this regard, apart from transcription, other processes such as RNA maturation and decay are under an exhaustive control and contribute to changes in gene expression. The relative importance of each process to determine the steady state expression levels of proteins has been studied in a few cellular systems, including the determination of basal levels of expression in fibroblasts (Schwanhauser et al. 2011) and the response of dendritic cells to lipopolysaccharide (Rabani et al. 2011). The conclusion of these and related studies is that transcription rate explains most of the observed variability in gene expression (Rabani et al. 2011; Schwanhauser et al. 2011, 2013; Li et al. 2014). However, the generality of this conclusion and more specifically, to which extent the transcriptional changes mediated by HIF affect mRNA expression changes observed under hypoxia, is unknown. In this regard, it has been demonstrated that in addition to transcription, regulation of half-life of certain mRNAs has fundamental roles in

hypoxia. For instance, the mRNA level of vascular endothelial growth factor A (VEGFA), a potent angiogenic factor, is regulated by hypoxia through both HIF-mediated transcriptional up-regulation, as well as HIF-independent post-transcriptional regulation. VEGFA mRNA is an unstable transcript with a short half-life (60–180 min) in normoxia. However, under hypoxia VEGFA mRNA is stabilized by the binding of different RNA-binding proteins, such as HuR or hRNP-L, to AU-rich elements (AREs) present in its 3'-UTR (Levy et al. 1996, 1998; Shih and Claffey 1999; Goldberg-Cohen et al. 2002). Other examples of transcripts whose half-lives are regulated by hypoxia through AREs include erythropoietin (EPO) (Goldberg et al. 1991; Rondon et al. 1991; Ho et al. 1995) and tyrosine hydroxylase (TH) mRNAs (Czyzyk-Krzeska et al. 1994). In addition, hypoxia has an important role in the regulation of miRNAs (Voellenkle et al. 2012; Nallamshetty et al. 2013; Camps et al. 2014), which in turn regulate mRNA stability and translation. Altogether, these results suggest a pervasive effect of hypoxia on mRNA stability, therefore, a global study of how hypoxia modulate mRNA half-life is needed to fully understand how cells adapt to low oxygen situations.

Herein, we determined the relative contribution of RNA transcription and stability to gene expression changes induced by hypoxia in primary human endothelial cells (human umbilical vein endothelial cells, HUVEC) that are central actors in the induction of angiogenesis, a key adaptation response to hypoxia. Our results show that *de novo* transcription explains 72% of gene expression changes observed upon exposure to low oxygen conditions, while half-life regulation underlies most of the remaining variability (24%). In addition, we found that the EGLN/HIF/TSC2 axis increases global RNA decay rates, leading to a 30% reduction of cellular mRNA as well as rRNA content. Thus, our work reveals an unexpected role of hypoxia on global RNA stability and provides quantitative data to understand the effect of hypoxia on gene expression.

RESULTS

Transcription rate is the major determinant of hypoxic mRNA levels

The transcriptional response to hypoxia has been studied in many different model systems at the genome scale (Hu et al. 2003; Scheurer et al. 2004; Elvidge et al. 2006; Warnecke et al. 2008; Xia and Kung 2009; Mimura et al. 2012; Weigand et al. 2012; Salama et al. 2015; Galbraith et al. 2017; Chen et al. 2018; Moreau et al. 2018; Chakraborty et al. 2019; Smythies et al. 2019). However, in all these works mRNA levels were determined by means of RNA-sequencing or microarrays and hence they could only assess the effect of hypoxia on steady state mRNA levels, which are determined by the combination of the relative rates of transcription and decay. Aiming to determine

the contribution of these two processes in the response to hypoxia, we pulse-treated HUVEC for 2 h with 4-thiouridine (4sU) to label nascent RNA and then characterized the pattern of newly transcribed mRNAs by affinity capture of the labeled transcripts followed by high-throughput sequencing (Fig. 1A). Incorporation of 4sU into mRNA after a short pulse can be used as a proxy for transcription rate and comparison of changes in this nascent fraction to those in total mRNA levels allows quantification of the role of mRNA stability on gene expression (Dolken et al. 2008; Schwanhauser et al. 2011). Hence, to determine the contribution of transcription to gene expression, we compared expression changes induced by hypoxia in the total mRNA fraction to those observed in the newly synthesized fractions (Fig. 1A). As shown in Figure 1B, taking a relatively relaxed threshold of $FDR < 0.05$, only about one third of the genes found differentially expressed in the total mRNA-

fraction showed altered transcription (27% and 35% of the down- and up-regulated genes, respectively). However, this percentage gradually increased up to ~70% in both fractions when, in addition to the FDR, we used increasingly restrictive fold-change thresholds to consider genes as differentially expressed in hypoxia (Fig. 1C). This result suggests that transcription is mostly responsible for large expression changes in the total mRNA fraction, whereas changes of small magnitude are likely due to variations of mRNA half-lives. In addition, we observed a significant positive linear correlation between changes in gene expression observed in total and newly transcribed mRNA fractions (Fig. 1D). By focusing on genes whose expression was significantly altered by hypoxia ($FDR < 0.01$) in either total or newly transcribed mRNA fractions, we found that the transcriptional component (changes in newly synthesized mRNA) accounted for 72% of the observed

FIGURE 1. Hypoxia-induced changes in gene expression are predominantly controlled at the level of transcription. (A) Schematic diagram depicting the experimental procedure and the information provided by each RNA fraction. (B) Venn diagrams showing the overlap between the genes found differentially overexpressed ($FDR < 0.05$) in the total (Total Up) and newly transcribed (Newly Up) fractions and the differentially down-regulated ($FDR < 0.05$) in the total (Total Down) and newly transcribed (Newly Down) fractions. (C) The graph represents the proportion of genes in the total mRNA fraction that were also differentially expressed in the newly transcribed fraction for: all regulated genes (black line), significantly down-regulated genes (purple line), and significantly up-regulated genes (orange line). Genes were considered differentially expressed if, in addition to being significantly regulated ($FDR < 0.05$), showed an absolute value of induction above the indicated threshold of $\log_2FC$ (x-axis). The magnitude of change induced by hypoxia was calculated for each gene as the absolute value of the logarithm of the ratio of the expression level in normoxia over hypoxia ($\log_2FC$). (D) The magnitude of change ($\log_2$ -fold change, $\log_2FC$) induced by hypoxia in 4sU-labeled genes ("Newly tr mRNA") was represented against the changes in the total mRNA fraction ("Total mRNA"). Genes significantly ($FDR < 0.01$) regulated by hypoxia in either fraction are labeled in green. Regression analysis of significantly regulated genes showed that total mRNA level increased significantly with transcription ($F_{(1,2225)} = 5756$, P -value < 0.01).

variability in total mRNA (regression analysis, $F_{(1,2225)} = 5756$, $P < 10^{-16}$, $R^2_{2225} = 0.7212$). In sum, these results suggest that transcriptional mechanisms largely determine gene expression changes in hypoxia.

RNA stability is significantly affected by hypoxia

In spite of the major role of transcription, this process cannot explain around 28% of the variability in total mRNA levels. To further explore the contribution of nontranscriptional processes, we investigated the effect of hypoxia on mRNA half-lives by sequencing newly synthesized mRNA (4sU-labeled) in parallel with preexisting (nonlabeled) and total mRNA fractions from the same RNA sample (Fig. 1A). This analysis allowed us to determine individual mRNA half-lives under normoxia and hypoxia (Supplemental Table S1; Dolken et al. 2008; Schwanhauser et al. 2011). First of all, we compared the half-lives values obtained in our analysis with the half-lives values determined for mouse-fibroblasts (NIH3T3) by the same technique in two independent works (Dolken et al. 2008; Schwanhauser et al. 2011). We found that the distribution of the half-lives of normoxic and hypoxic HUVEC mRNAs (median half-life of 8.7 and 5.7 h, respectively) fell between the range of the mRNA half-lives determined by Dolken et al. and Schwanhauser et al. (median half-life of 5 and 9 h, respectively), suggesting that our HUVEC determinations were comparable in magnitude to those reported for mouse fibroblasts (Fig. 2A). In addition, half-life of selected mRNAs were determined by measuring mRNA stability after inhibition of transcription by actinomycin D. For this analysis, mRNAs were selected according to the 4sU half-life determinations to include a wide representation of short-lived and medium-lived mRNAs (Supplemental Tables S1, S2). The mRNA half-life values estimated by this method were in good agreement with the values derived from the 4sU experiment (Fig. 2B; Pearson's $R_{48} = 0.65$, $P < 0.001$, linear regression $R^2 = 0.77$). Therefore, our determinations based on sequencing of the distinct mRNA fractions allowed for an accurate determination of mRNA half-lives. Next, we investigated whether hypoxia-induced changes on mRNA stability could account for the remaining variability in total mRNA levels that are not explained by transcription. A linear regression model including mRNA half-lives as an additional explanatory variable accounted for up to 96% of changes in total mRNA induced by hypoxia (multiple linear regression analysis, $F_{(2,1851)} = 26110$, $P < 0.0001$, adjusted $R^2_{1851} = 0.9658$). Thus, changes in mRNA half-life explain an additional 24% of the observed effects on gene expression and both variables, transcription and stability, jointly explain most of the variability in mRNA levels (Supplemental Movie S1).

In spite of its relatively minor role in the gene expression changes induced by hypoxia, the analysis of the effects of hypoxia on mRNA half-life revealed a modest but signifi-

cant ($P < 0.01$ paired t-test) reduction of the overall mRNA stability, from a median half-life in normoxia of 8.7 h to a median half-life in hypoxia of 5.7 h (Fig. 2A,C). We confirmed this result by estimation of individual mRNA levels over time upon inhibition of transcription with actinomycin D under normoxia or hypoxia (Fig. 2D). For these analyses we selected a set of mRNAs whose changes in stability in response to hypoxia, according to half-lives determined in our 4sU analysis (Supplemental Tables S1, S2), varied in magnitude. This selection included those genes represented in Figure 2B. Although the vast majority of mRNAs showed reduced half-life under hypoxia, our analysis also identified a few transcripts that were stabilized. Notably, among the latter was VEGFA, a transcript known to be stabilized in hypoxia (Levy et al. 1996, 1998; Levy 1998; Shih and Claffey 1999; Goldberg-Cohen et al. 2002). The pervasive effect on mRNA decay rates, suggested a general mechanism of RNA degradation rather than a sequence-specific mechanism such as those mediated by miRNAs. Therefore, we explored if the effect of hypoxia on mRNA decay rates was also observed in unfractionated RNA; most of which consists in rRNA. With this objective, we pulse-labeled HUVEC cells with tritiated uridine and determined the decay of radioactive label in total RNA. As shown in Figure 2E, the remaining radioactivity on total RNA after 16 h of culture was higher in normoxia than in hypoxia, suggesting increased rRNA decay rates in hypoxia. Thus, hypoxia induces a global decay in RNA, rather than a specific effect on protein coding genes.

In addition to increased decay, a general reduction on transcription rates could also potentially contribute to decreased RNA levels in hypoxic cells. However, we did not observe a decrease in 4sU incorporation in HUVEC cells exposed to 1% oxygen for 8 h compared to normoxic controls (Fig. 2F). In fact, mRNA synthesis was slightly increased in hypoxia, suggesting very similar global transcription rates in both situations. In agreement, determination of the quantity of the 47S precursor ribosomal RNA (pre-rRNA) after 16 h of hypoxic exposure, which correlates with the rate of rRNA transcription (Cui and Tseng 2004), did not show significant differences between normoxia and hypoxia (Fig. 2G).

In summary, these results suggest that hypoxia results in modest increase of mRNA and rRNA degradation and, although no global changes in transcription are detected (Fig. 2F,G), changes in transcription of specific mRNAs largely contributes to the observed changes in mRNA expression during hypoxia (Fig. 1D).

Hypoxia-induced RNA decay is HIF-dependent

The HIF axis is the best characterized signaling pathway in the response to hypoxia. Thus, as a first attempt to explore the molecular mechanism by which RNA degradation was

FIGURE 2. Decrease of mRNA and total RNA half-lives under hypoxic conditions. (A) Distribution of mRNA half-lives, determined by 4sU pulse-labeling as indicated in Materials and Methods, in normoxic (blue) and hypoxic (red) HUVEC. The values published in Dolken et al. (2008) (Dolken, dotted line) and (Schwanhauser et al. 2011) (Schw, black line) are overlapped for comparison. (B) Comparison of individual half-life values obtained by the high-resolution gene expression profiling experiment and the data derived from the determination of mRNA half-life upon inhibition of RNA-pol with actinomycin D. ($F_{(1,49)} = 168$, $P < 0.001$, linear regression $R^2 = 0.77$) (C) Distribution of changes in decay rates induced by hypoxia, as determined in the high-resolution profiling experiments, represented as the logarithm of the half-life in hypoxia over normoxia. (D) The graph represents the half-life of individual genes, calculated upon inhibition of RNA-pol with actinomycin D, in HUVEC cells in normoxia versus hypoxia as $\log_2$ of the ratio. Each dot represents the mean of three independent experiments. The red dot represents *VEGFA*. (E) Pulse-labeling of total RNA determined in HUVEC. The graph represents the logarithm of the ratio of the hypoxic (16 h) remaining radioactivity normalized by normoxic values. The comparison of the observed ratios was significantly higher than expected from a sample with a mean $\log_2$ -ratio of 0 (single sample Student's *t*-test, P -value = 0.0122). (F) The amount of mRNA in the newly synthesized mRNA fraction (4sU-labeled) from the experiment described in Figure 1A was determined by spectrophotometry and represented as the $\log_2$ of the ratio of the values obtained in hypoxia over normoxia. Each point represents data from an independent experiment. The comparison of the observed ratios was significantly higher than expected from a sample with a mean $\log_2$ -ratio of 0 (single sample Student's *t*-test, P -value = 0.049). (G) 47S rRNA precursor levels were determined by RT-qPCR in cells exposed overnight to normoxia or hypoxia and values normalized to 28S rRNA content. Each dot represents the ratio hypoxia over normoxia in an independent experiment. The observed ratios were not significantly different to the expected value of 0 (single sample Student's *t*-test, $P = 0.8185$).

increased in hypoxia, we investigated the effect of EGLN inhibitor Dimethylxalylglycine (DMOG) on total RNA mass. The EGLN family of enzymes acts as cellular oxygen sensors and functions as the most apical elements of the HIF axis suppressing its activity in the presence of oxygen, thus DMOG prevents HIF α degradation in normoxia. As is shown in Figure 3A, mRNA content was reduced in hypoxic conditions as well as in DMOG-treated cells compared with normoxia. Since HIF is the major target of EGLNs and a key player in the response to hypoxia, we next examined its involvement in this process by silencing the constitutive ARNT subunit. ARNT knockdown resulted in 36%–15% of control ARNT expression values and blocked the

induction of the HIF target *EGLN3* (Fig. 3B,C). Additionally, we performed an ARNT western blot to confirm ARNT silencing (Fig. 3D). Importantly, ARNT knockdown significantly attenuated the RNA decay promoted by hypoxia (Fig. 3E), suggesting that HIF activity is necessary in this process. To further address the involvement of HIFs, we silenced *HIF1A* or *EPAS1* expression to 30%–35% and 30%–15% of their control levels, respectively (Fig. 3F–H). Then, silenced and control HUVEC were pulse-labeled with tritiated uridine and exposed to normoxia or hypoxia for 16 h. After measuring the radioactivity present in total RNA, we observed that silencing of *HIF1A* or *EPAS1* partially prevent RNA decay (Fig. 3I).

FIGURE 3. (Legend on next page)

To analyze if HIF stabilization was sufficient to promote RNA degradation, we infected HUVEC with lentiviral vectors that overexpressed constitutively active forms of *EPAS1* (EPASPP) or *HIF1A* (HIFPP) that had their oxygen-sensitive prolines residues substituted by alanine to prevent hydroxylation, rendering these isoforms constitutively active even in normoxia (Kondo et al. 2002). Additionally, we included mutant versions of these constructs whose bHLH motif was altered to prevent their capacity to bind DNA (EPASPPBH, HIFPPBH) (Kondo et al. 2002). As expected, the expression of EPASPP in normoxia resulted in a strong induction of *ADM* mRNA, a *EPAS1*-target gene, whereas the expression of EPASPPBH had no effect

(Fig. 3J,K). Similarly, the HIF1A-target *BNIP3* was strongly induced by overexpression of HIF1PP with intact bHLH domain (Fig. 3L,M). We also analyzed by western blot the HIFPP/HIFPPBH and EPASPP/EPASPPBH overexpression (Fig. 3N). As is shown in Figure 3O,P, transduction of the constitutively active forms of both *EPAS1* and *HIF1A* decreased RNA content per cell in normoxia. In contrast, the overexpression of the mutants lacking a functional DNA binding motif did not reduce RNA content. Therefore, these results suggest that HIF transcriptional activity is necessary and sufficient to induce RNA decay. Furthermore, the dependency on an intact bHLH domain suggested that the increased decay rate might be

FIGURE 3. Hypoxia-induced mRNA decay is HIF-dependent. (A) mRNA content per cell from cells grown for 16 h in normoxia, hypoxia (Hyp) or DMOG (DMOG). Each dot represents the ratio of mRNA levels in hypoxia or DMOG over normoxia in an independent experiment. The observed ratios were significantly lower than expected from a sample with a mean ratio of 1 (single sample Student's *t*-test, *P*-value = 0.033 and *P*-value = 0.041, respectively). (B) *ARNT* mRNA levels in HUVEC cells transduced with lentiviral particles expressing a shRNA to silence *ARNT* expression versus control (NS). The *ARNT* expression was on average 21% and 25% of that in control cells in normoxia and hypoxia, respectively, and significantly different to the expected value of 1 (single sample Student's *t*-test, *P* = 0.006 and *P* = 0.007 for cells under normoxia and hypoxia, respectively). (C) *EGLN3* mRNA levels in HUVEC cells transduced with lentiviral particles expressing a shRNA to silence *ARNT* expression versus control (NS). The effect of hypoxia on *EGLN3* induction was significantly attenuated in *ARNT* knockdown cells (*P*-value = 0.017; paired Student's *t*-test). (D) HUVEC cells transduced with lentiviral particles expressing a shRNA to silence *ARNT* (sh*ARNT*) or a control shRNA (NS) or mock infected (UT) were exposed to normoxia (Nx) or hypoxia (Hyp) for 16 h. The level of *ARNT* protein and tubulin were determined by immunoblot. (E) Ratio of hypoxic versus normoxic total RNA content per cell from control cells (NS) and cells whose *ARNT* expression was silenced (sh*ARNT*). Each dot represents a determination from a single experiment and matched observations (derived from the same experiment) are represented joined by a line. The effect of hypoxia on RNA level was significantly attenuated in *ARNT* knockdown cells (*P*-value = 0.014; paired Student's *t*-test). (F) *EPAS1* mRNA levels in HUVEC cells transduced with lentiviral particles expressing a shRNA to silence *EPAS1* expression versus control (NS). *EPAS1* expression was on average 30% and 15% of that in control cells in normoxia and hypoxia, respectively, and significantly different to the expected value of 1 (single sample Student's *t*-test, *P* = 0.0021 and *P* = 0.0002 for cells under normoxia and hypoxia, respectively). (G) *HIF1A* mRNA levels in HUVEC cells transduced with lentiviral particles expressing a shRNA to silence *HIF1A* expression versus control (NS). *HIF1A* expression was on average 30% and 35% of that in control cells in normoxia and hypoxia, respectively and significantly different to the expected value of 1 (single sample Student's *t*-test, *P* = 0.0009 and *P* = 0.0002 for cells under normoxia and hypoxia, respectively). (H) HUVEC cells transduced with lentiviral particles expressing a shRNA to silence *HIF1A* (sh*HIF1A*), *EPAS1* (sh*EPAS1*) or a control shRNA (NS) or mock infected (UT) were exposed to normoxia (Nx) or hypoxia (Hyp) for 16 h. The levels of *HIF1A*, *EPAS1* and β -actin were determined by immunoblot. (I) Pulse-labeling of total RNA determined in control HUVEC cells (NS) and cells whose *HIF1A* or *EPAS1* expression was silenced (sh*HIF1A* and sh*EPAS1*). The graph represents the logarithm of the ratio of the hypoxic (16 h) remaining radioactivity normalized by normoxic values. Each dot represents a determination from a single experiment. The effect of hypoxia on RNA level showed clear tendency to be attenuated in sh*HIF1A* and sh*EPAS1* but did not reach statistical significance (sh*HIF1A* *P* = 0.14; paired Student's *t*-test; sh*EPAS1* *P* = 0.08; paired Student's *t*-test). (J) *EPAS1* mRNA levels of HUVEC cells transduced with a constitutively stable form of *EPAS1* (EPASPP) or a stable but transcriptionally inactive form (EPASPPBH) compared with the control (PRRL). The expression of mutant forms of *EPAS1* was similar (*P* = 0.43, paired Student's *t*-test). (K) *ADM* mRNA levels of HUVEC cells transduced with a constitutively stable form of *EPAS1* (EPASPP) or a stable but transcriptionally inactive form (EPASPPBH) compared with the control (PRRL). The effect of EPASPP on *ADM* expression was significantly higher than that of EPASPPBH (*P* = 0.023, paired Student's *t*-test). (L) *HIF1A* mRNA levels of HUVEC cells transduced with a constitutively stable form of *HIF1A* (HIFPP) or a stable but transcriptionally inactive form (HIFPPBH) compared with the control (PRRL). The expression of HIFPPBH was higher than that of HIFPP but the difference did not reach statistical significance (*P* = 0.34, paired Student's *t*-test). (M) *BNIP3* mRNA levels of HUVEC cells transduced with a constitutively stable form of *HIF1A* (HIFPP) or a stable but transcriptionally inactive form (HIFPPBH) compared with the control (PRRL). The effect of HIFPP on *BNIP3* expression was higher than that of HIFPPBH, but the difference did not reach statistical significance (*P* = 0.079, paired Student's *t*-test). (N) Western blot to determine the overexpression of the HIFPP, HIFPPBH, EPASPP, EPASPPBH, or empty vector (PRRL) in samples in normoxia (Nx) or hypoxia (Hyp) for 16 h. (O,P) Ratio of hypoxic versus normoxic total RNA content per cell in HUVEC cells overexpressing a constitutively active *EPAS1* (EPASPP) or *HIF1A* (HIFPP) and a constitutively active *EPAS1* or *HIF1A* whose bHLH motive has been mutated (EPASPPBH, HIFPPBH). Each dot represents a determination from a single experiment and matched observations (derived from the same experiment) are represented joined by a line. The expression of EPASPP resulted in RNA level significantly reduced as compared to control cells, (*P*-value = 0.025, single sample Student's *t*-test) and the effect of each *EPAS1* mutant on RNA level was significantly different (*P*-value = 0.025, paired Student's *t*-test). The expression of HIFPP resulted in RNA level reduced as compared to control cells, but did not reach statistical significance (*P*-value = 0.08, single sample Student's *t*-test) and the effect of each HIF mutant on RNA level was not significantly different (*P*-value = 0.23, paired Student's *t*-test). (Q) Ratio of remaining radioactivity present in hypoxic samples over normoxia in untreated cells (UT) and cells treated with actinomycin D (ACTD), cycloheximide (CHX) or puromycin (PURO) after pulse-labeling with tritiated uridine. Each dot represents a determination from a single experiment. A single factor ANOVA showed a significant difference among the four treatment groups UT, ACTD, CHX, and PURO ($F_{(3,8)} = 8.487$, *P*-value = 0.007) and a posteriori Tukey test showed that all drug treatment means were significantly different than the control mean (*P*-value = 0.006, *P*-value = 0.022, and *P*-value = 0.047 for ACTD, CHX, and PURO, respectively).

mediated by a HIF-target(s). To test this possibility, HUVEC cells were pulsed during 2 h with tritiated uridine, and then treated with actinomycin (ACTD), cycloheximide (CHX) or puromycin (PURO) and exposed to hypoxia or normoxia for 8 additional hours. After these treatments, RNA was purified from samples and the remaining radioactivity was quantified by scintillation counting. As is shown in Figure 3Q, inhibition of transcription with actinomycin D or inhibition of translation with cycloheximide or puromycin completely blocked RNA degradation in hypoxia. All these results suggest that a HIF target(s), whose transcription is induced by hypoxia, is responsible for the increased RNA decay rates.

Inactivation of mTOR through TSC2 induces RNA decay in hypoxia

It is well established that during hypoxia, the oxygen-deprived cells diminish protein synthesis through the inhibition of multiple key regulators of translation, including eIF2a and the mammalian target of rapamycin (mTOR), in an attempt to save energy (Liu et al. 2006; Wouters and Koritzinsky 2008). In addition, RNA stability is intimately coupled to translation (Garneau et al. 2007), therefore, it could be possible that the decreased RNA half-life observed in hypoxia was a consequence of blunted translation. To test this possibility, we first determined the effect of hypoxia on translation in HUVEC. To that end, we used the Click-iT Plus OPP Alexa Fluor 647 Protein Synthesis Assay Kit to detect protein synthesis in normoxic, DMOG-treated cells and cycloheximide-treated control cells. As is shown in Figure 4A, DMOG induces a significant decrease in the protein translational rates compared with normoxic cells. The effect of hypoxia and DMOG on protein synthesis was independently confirmed using metabolic pulse-labeling with ³⁵S-Methionine (Fig. 4G). However, while silencing of ARNT prevented RNA decay (Fig. 3E), it had no effect on the inhibition of protein synthesis induced by DMOG (Fig. 4B,C), suggesting that RNA decay is not a mere consequence of blunted translation.

It has been described that hypoxia triggers HIF-dependent and independent mechanisms to suppress translation. Thus, some of the HIF-dependent pathways could be coupled to RNA degradation. One of the former mechanisms involves the TSC2-dependent inactivation of mTOR by the HIF-target *REDD1*, resulting in the inhibition of the cap-dependent translation (Brugarolas et al. 2004; Wouters and Koritzinsky 2008). In agreement, we observed decreased phosphorylation of the ribosomal protein S6 (pS6), a direct target of the kinase mTOR, upon treatment with hypoxia (Fig. 4D) or DMOG (Fig. 4E). Accordingly, interference of *EPAS1* prevented the suppression of mTOR activity by hypoxia in HUVEC (Fig. 4F). Thus, we next tested the effect of knocking down TSC2 expression on protein synthesis and RNA stability. Hypoxia

still repressed protein synthesis in TSC2 silenced cells (Fig. 4G), in spite of efficient reduction of TSC2 expression to 11%–7% of the control levels (Fig. 4H). Despite its lack of effect on translation, silencing of TSC2 significantly prevented RNA decay induced by hypoxia (Fig. 4I). Altogether, these results suggest that pathway EGLN/HIF/TSC2 regulates RNA stability independently of its effect on protein synthesis.

DISCUSSION

In the present work, we aimed to achieve a comprehensive and quantitative understanding of the gene expression changes that occur under adaptation of primary endothelial cells to hypoxic conditions. Previous studies focused on the analysis of the steady state mRNA levels, which are dictated by transcription and decay rates. Thus, we used a metabolic-labeling strategy to study the relative contribution of these processes on mRNA levels in response to hypoxia. In agreement with previous results on general mRNA regulation (Dolken et al. 2008; Rabani et al. 2011; Schwanhauser et al. 2011), we found that, although the relative contribution of transcription and decay rates is different for each individual gene, on average, transcription plays a dominant role on the changes induced by hypoxia in total mRNA fraction (Fig. 1D). Specifically, hypoxia-induced changes in transcription rates explain up to 72% of the variability in total mRNA levels. Moreover, the fraction of differentially expressed genes shared by the total and newly transcribed fractions gradually decreases with more lenient thresholds of fold-change (Fig. 1C). This result suggests that transcription is mostly responsible for large expression changes in the total mRNA fraction, whereas changes of small magnitude are mainly due to variations of mRNA half-lives. Given the role of miRNAs in fine-tuning expression levels at the post-transcriptional level and the impact of hypoxia on the expression of noncoding RNAs (Choudhry et al. 2014), it is tempting to speculate that the effects of hypoxia on mRNA half-lives could be mediated by miRNAs. Alternatively, this observation may indicate a higher false positive rate when calling differentially expressed genes using lenient thresholds. Finally, we also found genes whose changes in transcription did not correlate with changes in the total RNA fraction. One reason for this discrepancy is that the relative long half-life of mRNAs obscure the detection of changes in transcription by the determination of total expression levels.

Despite the major role of transcription, changes in half-life also contribute to gene regulation by hypoxia. According to our linear model, half-life explains nearly 24% of the variability on gene expression levels. To identify biological processes associated to genes whose expression was preferentially modulated at the level of transcription or stability, we applied gene set

enrichment analysis (GSEA) to the list of genes expressed in HUVEC sorted according to their hypoxia-induced change in transcription or decay rates, respectively. We found an enrichment in gene ontology (GO) terms related to metabolism, in particular monosaccharide catabolism, and to response to oxygen levels for genes whose tran-

scription was up-regulated in response to hypoxia. However, we could not find any statistically significant overrepresentation of GO-terms for decay-rate sorted genes. In addition, GSEA did not find enrichment of targets of any specific miRNA in the sets of genes whose half-life was decreased by hypoxia. This suggests that

FIGURE 4. (Legend on next page)

either mRNA decay is controlled by the combined effect of the many independent miRNAs, that are induced by hypoxia (Kulshreshtha et al. 2007; Camps et al. 2014) and the effect of different RNA binding proteins, such as AUBPs, or that it is not mediated by miRNAs. It is important to emphasize that the relative contribution of transcription and stability to the hypoxia-induced changes in mRNA levels vary for individual genes, 72% and 24% being the contribution of these two processes on average. Moreover, the increased expression of individual mRNAs can be the result of the coordinated effect of hypoxia on their transcription and decay rates. This is exemplified by *VEGFA*, a prototypical hypoxia-responsive gene, whose induction involves up-regulation of transcription and decrease of decay rate (Shima et al. 1995; Levy et al. 1996). Changes in *VEGFA* mRNA stability allow for a fast response to hypoxia; while transcriptional regulation contributes as a later mechanism. On the other hand, the half-life of *SLC23A2* (that encodes for the major transporter of ascorbate in endothelial cells) is dramatically reduced under hypoxia from 7.6 to 3 h (Supplemental Table S1) in parallel with a reduction in its transcription. Interestingly, ascorbate has been described to prevent HIF1A stabilization (Kuiper et al. 2010, 2014), thus the rapid mRNA destabilization of *SLC23A2* could facilitate HIF induction under hypoxia.

A potential caveat of our analyses is that our determinations of transcription and decay rates derive from the same 4sU-data sets and thus both variables are not independent, as formally required in multivariate linear regression.

Nonetheless, the concordance of half-life values derived from the 4sU-labeling experiments and those derived from an independent technique based on inhibition of transcription with actinomycin D support the validity of our half-life estimations.

In spite of the relatively minor role of mRNA decay on regulation of gene repression, we found that hypoxia resulted in modest (median reduction of 34%–36%) but statistically significant reduction of the overall mRNA stability (Fig. 2A,C). This effect was confirmed by the determination of mRNA decay rates upon transcription inhibition with actinomycin-D (Fig. 2D). We also found that, not only mRNA, but rRNA content per cell was also decreased in hypoxia (Fig. 2E). This result has an important implication, although hypoxia induces the transcription of a large number of miRNAs (Kulshreshtha et al. 2007; Camps et al. 2014), the global effect on both mRNA and rRNA makes miRNAs unlikely mediators of this process, and suggests a general mechanism of RNA degradation rather than a sequence-specific mechanism such as those mediated by miRNAs, in agreement with the result of GSEA mentioned above. In search for a potential mechanism, we found that EGLN/HIF1A-EPAS1 axis was involved in this effect (Fig. 3). Silencing of *ARNT*, *HIF1A* or *EPAS1* prevented the decrease in RNA content by hypoxia (Fig. 3E,I), while overexpression of active forms of HIF1A and EPAS1 dramatically decrease RNA content per cells (Fig. 3O,P). Thus, this group of experiments suggest that HIF is necessary and sufficient to induce RNA decay in hypoxia. The fact

FIGURE 4. Inactivation of mTOR through TSC2 induces RNA decay in hypoxia. (A) Ratio of the integrated density of the 647 fluorescence divided by the integrated density of the DAPI fluorescence determined by Click-iT Plus OPP Alexa Fluor 647 Protein Synthesis Assay Kit in HUVEC cells grown 16 h with cycloheximide (CHX), DMOG (DMOG), or untreated (UT). The 647 fluorescence of CHX and DMOG treatment was significantly lower than that of untreated cells (UT) (CHX P -value < 0.0001, DMOG P -value < 0.0001. Unpaired Student's t -test). The experiment was repeated in duplicate, a representative experiment is shown. Each dot represents an individual cell. (B) Images of the fluorescence determined by Click-iT Plus OPP Alexa Fluor 647 Protein Synthesis Assay Kit shown in Figure 4C. (C) Ratio of the integrated density of the 647 fluorescence divided by the integrated density of the DAPI fluorescence determined by Click-iT Plus OPP Alexa Fluor 647 Protein Synthesis Assay Kit in HUVEC cells grown 16 h with cycloheximide (CHX), DMOG (DMOG), or untreated (UT). HUVEC cells transduced with lentiviral particles expressing a shRNA to silence *ARNT* (shARNT) or control shRNA (NS). The 647 fluorescence of CHX and DMOG treatment was significantly lower than that of untreated cells (UT) in both conditions: in shARNT cells and in control cells. (****) Adjusted P -value < 0.0001; [*] adjusted P -value = 0.0104, two-way ANOVA followed by Tukey's multiple comparison test). Hypoxia did not have a different effect in cells whose *ARNT* expression was silenced, compared with the control cells ([ns] adjusted P -value > 0.999; two-way ANOVA followed by Tukey's multiple). Each dot represents an individual cell. The graph represents a representative experiment that was performed in duplicate. (D) HUVEC were exposed to hypoxia for the indicated periods of time and the level of HIF1, EPAS1, phospho-S6 Ribosomal Protein (rpS6), total rpS6 and lamin B1 were determined by immunoblot. (E) HUVEC were treated with DMOG (DMOG) or vehicle (Nx) for 16 h and the levels of phospho-S6 Ribosomal Protein (rpS6), total rpS6 and lamin B1 were determined by immunoblot. (F) HUVEC cells transduced with lentiviral particles expressing a shRNA to silence *HIF1A* (shHIF1a) or *EPAS1* (shEPAS1) expression, a control shRNA (NS) or mock infected (UT) were exposed to normoxia (Nx) or hypoxia (Hyp) for 16 h. The level of HIF1, EPAS1, phospho-S6 Ribosomal Protein (rpS6), total rpS6, and lamin B1 were determined by immunoblot. (G) Ratio of the radioactivity incorporated into proteins precipitated with TCA over total label in cell extracts of hypoxic or DMOG-treated HUVEC cells compared with normoxic extracts. HUVEC cells were transduced with lentiviral particles expressing a shRNA to silence *TSC2* (shTSC2) or control shRNA (NS). Each dot represents a single experiment. Silencing of *TSC2* expression did not result in any statistical difference with the control cells (P -value = 0.1632 and P -value = 0.44 for Hyp and DMOG treated cells, respectively. Paired Student's t -test). (H) *TSC2* mRNA levels in HUVEC cells transduced with lentiviral particles expressing a shRNA to silence *TSC2* expression versus control (NS). *TSC2* expression was on average 11% and 7% of that in control cells in normoxia and hypoxia, respectively, and significantly different to the expected value of 1 (single sample Student's t -test, P < 0.0001 for cells under normoxia and hypoxia). (I) Pulse-labeling of total RNA determined in control HUVEC cells (NS) and cells whose *TSC2* expression was silenced (shTSC2). The graph represents the logarithm of the ratio of the hypoxic (16 h) remaining radioactivity normalized by normoxic values. Each dot represents a determination from a single experiment. The effect of hypoxia on RNA is significantly reduced by silencing *TSC2* expression (P -value = 0.0288; paired Student's t -test).

TABLE 1. Primers used in this study

Application	Name	Forward primer	Reverse primer
qPCR	<i>ACTB</i>	AAGGCCAACCGCGAGAAG	ACAGCCTGGATAGCAACGTAC
qPCR	<i>EGLN3</i>	ACACGAAGTGCAGCCCTCTT	TCTTCAGCATCAAAGTACCAGACAGT
qPCR	<i>VEGFA</i>	CCTTGCTGCTCTACCTCCAC	ATGATTCTGCCCTCCTCCTT
qPCR	<i>ZNF800</i>	CACGCCGCAATGAAGAGC	AAGGAGGATCTCCAGTTCCA
qPCR	<i>ZNF281</i>	ATGACCACCATGGCACTGAG	TCTGGCTTTGGCCTTTTTGC
qPCR	<i>ZNF496</i>	AGGAAGAGAATGTGCGGGAC	CCCCGTAATCCTCCCCAATG
qPCR	<i>POLR3D</i>	CCCTCGGGGGAGTCAAGA	TCTCTGTCCTTTACAGCTTC
qPCR	<i>EHMT1</i>	GGGAGCCTCAGCAGGATTG	CTTCATCGGCAGCCATAGGT
qPCR	<i>TMEM167B</i>	CCGCCGCTATTACCACTGAA	AGCCAGGTTTGTAGACGAGG
qPCR	<i>SCYL2</i>	GGACACTACTCTAGGACCGGG	TTCCAAGCTAGCCCATTGCC
qPCR	<i>EID2</i>	TGCTCCCGGCAGAATACAAG	CACCCTGTGAGGATTTGCT
qPCR	<i>TERF2IP</i>	CGGATAGCGGGGAACAC	AATCAGGAGGGCTCTCATCCA
qPCR	<i>DCAF15</i>	CCGCTACGTCCTCTCTACA	TCGTCCTGGAATAGCCGAAC
qPCR	<i>DCAF16</i>	CTCTGAGAAGGGAGACGACG	ACTTCAGAGGTGCTTGTGGAA
qPCR	<i>MCM5</i>	CACGTTGGATGCTGCCTTG	CCGGGTATTTCTGCTTGGTG
qPCR	<i>ACE</i>	GTGTGGAACGAGTATGCCGA	GGGTGTGGTTGGCTATTTGC
qPCR	<i>PPRC1</i>	CAGTGTTGGGGAAGTCGAA	TTCCAGGTCCTTGTCCCTCA
qPCR	<i>MCM2</i>	ATCTACGCCAAGGAGAGGGT	GTAATGGGGATGCTGCCTGT
qPCR	<i>MCM3</i>	AAAGACGGCAGACTCACAGG	GGCGATTATGCGGATTGAC
qPCR	<i>MCM6</i>	TCCAAGACCTGCCTACCAGA	ACAAGCTCTGGGTGAAGTGG
qPCR	<i>DKC1</i>	CGCGGTGCAGGGTAACAT	GGCAATGACTCCGCTCCTTT
qPCR	<i>HMGCS1</i>	GGCGTCCCCTCCAAATGAT	GGTACTTTCTTGGCAGGGCT
qPCR	<i>NOP58</i>	AAGCATGCAGCTTCTACCGT	GGTTTTGGCTGCCAGCATT
qPCR	<i>PGM1</i>	AGCTATGAGAAGGACGTTGCC	TGGAGGGACGTACCACATCA
qPCR	<i>DNAJB6</i>	GGACGGGGTCTGTTTTCTCT	GAGAATGAAGTGAGGCCCCCC
qPCR	<i>HTATIP2</i>	CCAAAGTCACGCTCATTGGC	GTGGTACCCAGGCAACAGAA
qPCR	<i>SEMA6C</i>	AACCACGGATTGCGAACTCA	GGGCTTGAGGGGGATAGGAA
qPCR	<i>TMEM44</i>	TCCTTCGGCCTGTGGATCTG	GGTCCTGTCTGGGTTTCTGT
qPCR	<i>ENO2</i>	TGGCTTTGCCCCCAATATCC	TCTTTCCGTGTAGCCAGCC
qPCR	<i>KLF7</i>	CAAATGCAACCACTGCGACA	CATGGCAACTCTGGCCTTTC
qPCR	<i>NARF</i>	ATCTTCAGACATGCGGCCAA	CAGCAAAGCGTAACACCACC
qPCR	<i>P4HA2</i>	TGGTACAACCTCTTGCGGAG	TTGACATGGGCTGAAGGACC
qPCR	<i>CACNB3</i>	CTAGTGAAAGAGGGCGGGGA	CCGCCTGCTTTTGTCTCTG
qPCR	<i>DLG4</i>	AGCAGGAGTTCACAGAGTGC	CTCGATGACACGCTTACCT
qPCR	<i>ZNF581</i>	ACACCCAGAGGTCCCCTTCT	TCTCAACGGAGGAAATGCCA
qPCR	<i>BNIP3</i>	GGTCAAGTCGGCCGAAAAT	TGGAGGTTGTGACAGCGCTT
qPCR	<i>ARNT</i>	GAGAATTCAGGAATAGTGGCT	CGAGTCTTAGCAGTAGCCTGG
qPCR	<i>Pre-RNA</i>	CCTGCTGTTCTCTCGCGCTCCGAG	AACGCCTGACACGCACGGCAGGAG
qPCR	<i>ADM</i>	GGGCTTGGACTTCGGAGTTT	GTACATCAGGGCGACGGAAA
qPCR	<i>HIF1A</i>	Hs 00936368 - m1 HIF1A	
qPCR	<i>EPAS1</i>	Hs 01026149 - m1 EPAS1	

that the active HIF isoforms lacking the bHLH motif do not lead to RNA decay together with the dependence on de novo transcription and translation (Fig. 3O–Q) point to a common target of HIF1A and EPAS1 as an effector for inducing RNA decay whose expression is induced in response to hypoxia.

REDD1 (official name DDIT4) is a target of HIF, and its induction favors the dissociation of TSC2 from the inhibito-

ry 14-3-3 proteins, leading to the inhibition of mTOR (Brugarolas et al. 2004; Liu et al. 2006; Deyoung et al. 2008; Wouters and Koritzinsky 2008). Our results show that, in parallel to RNA decay, HUVEC cells inactivate mTOR in a HIF-dependent manner (Figs. 3E, 4F). In addition, silencing of *TSC2* prevents RNA decay induced by hypoxia (Fig. 4I), suggesting that the axis EGLN/HIF/REDD1/TSC2/mTOR regulates RNA stability in hypoxia

in HUVEC cells. However, we have not formally proved that the effect of HIF on mTOR is mediated through *REDD1* and *TSC2*. In fact, there are other mechanisms by which mTOR is regulated by hypoxia. First, AMP-activated protein kinase (AMPK) regulates mTOR in a HIF-independent way, through *TSC1-TSC2* (Liu et al. 2006). Second, the HIF-dependent induction of *BNIP3* (BCL2/adenovirus E1B 19 kDa protein-interacting protein 3) regulates mTOR by direct binding to RHEB (Li et al. 2007). In addition, the promyelocytic leukemia tumor suppressor (PML) can bind to mTOR during hypoxia and inactivate it by sequestration in nuclear bodies (Bernardi et al. 2006). Thus, further work will be required to determine the specific candidates that play a role in mTOR inactivation by hypoxia in HUVEC cells.

mTOR has been involved in the suppression of translation during hypoxia (Liu et al. 2006; Wouters and Koritzinsky 2008). However, to our knowledge, it has not been implicated in RNA decay in hypoxia, in spite of the well-documented relation between translation and RNA stability. When mTOR is inactivated, it decreases its serine/threonine kinase activity leading to dephosphorylation of its targets such as S6K and 4E-BP1. 4E-BP1 dephosphorylated sequesters eIF4E protein favoring its dissociation from the cap of the mRNAs, inhibiting CAP-dependent translation and leaving the mRNA unprotected. Translation is a highly energy-demanding process that consumes a lot of energy from the cells, thus there are divergent regulatory pathways that converge to ensure the blockage of translation. Additionally to the mTOR axis, there are other mechanisms by which hypoxia regulates translation independently of mTOR such as the regulation of ternary complex availability through the phosphorylation of the alpha subunit of eIF2 α . This occurs by the unfolded protein response (UPR) in the endoplasmic reticulum in a HIF-independent manner through PERK activation (Koumenis et al. 2002). The existence of these mTOR-independent pathways to block translation can explain why we do not observe a recovery in translation by silencing *TSC2* or *HIF* (Fig. 4C,G). In this context the increased RNA decay in hypoxia, instead of being a mere consequence of the inhibition of translation, could be contributing to the repression of protein synthesis in hypoxia by decreasing mRNA availability.

We observed rRNA degradation in parallel with mRNA degradation in hypoxia that could not be explained just by the dissociation of eIF4e (Fig. 2E). An attractive hypothesis is that in hypoxic cells mRNAs that are not being translated, together with the ribosomes, which are no longer needed for the translation of the hypoxic cell, are targeted for degradation; resulting in decreased half-life rates of both mRNA and rRNA. mTOR is fundamental for cell growth and cell survival as it regulates important processes such as mRNA translation, autophagy, or metabolism (Kim

and Guan 2019) and has been involved in ribosomal biogenesis (Mayer and Grummt 2006; Iadevaia et al. 2012). Thus, mTOR could also regulate rRNA decay. In this regard, it has been described that *S. cerevisiae* treated with rapamycin rapidly decreases its ribosomic content by 40% to 60% (Pestov and Shcherbik 2012). In addition, mTOR inhibition also increases ribophagy by random engulfment of cytosolic ribosomes (An and Harper 2018, 2020; Wyant et al. 2018). Thus, mTOR inactivation by hypoxia could mediate rRNA as well as mRNA degradation by hypoxia; however, further work is needed to determine the mechanisms by which mTOR inactivation favors rRNA decay.

In addition to HUVEC, we have also observed decreased RNA content in hypoxia in other types of endothelial cells, such as human microvascular endothelial cells (HMVEC), but not in other cell types like Hela, HEK293T, human fibroblasts or mouse embryonic fibroblasts (data not shown). Thus, RNA degradation could be a specific response of endothelial cells to hypoxia. Alternatively, it could be a general response that is triggered at different hypoxic thresholds depending on the cell type. In agreement, Hepa 1–6 (Johnson et al. 2008), C2C12, and NIH3T3 cells (Mekhail et al. 2006) but not HUVEC, blunt their RNA synthesis in response to very low oxygen concentrations or anoxia. Different cell types could diverge in the mechanisms used to decrease RNA levels in hypoxia to inhibit translation.

In eukaryotes, the majority of mRNAs undergo decay through a pathway that is initiated by the shortening of the poly(A) tail by deadenylases (PAN2–PAN3, CCR4–NOT, and PARN).

Therefore, by targeting some of the enzymes that are involved in the RNA-decay machinery, global RNA half-life could potentially be regulated. However, we did not observe hypoxic-induction of any of the genes encoding general RNA turnover proteins or endonucleases. In fact, we found decreased expression of several components of the exosome complex (*EXOSC2*, *EXOSC4*, *EXOSC6*, and *EXOSC9*) required for general RNA degradation (data not shown). Nevertheless, we cannot rule out that hypoxia alters the protein level or activity of the enzymes involved in general RNA degradation.

In summary, our work supports that transcription is the major determinant of the relative abundance of transcripts in response to hypoxia. Notwithstanding, RNA stability significantly contributes to the changes in gene expression during adaptation to decreased oxygenation. In addition, we also uncovered an unexpected effect of hypoxia on total RNA content, by a EGLN/HIF/TSC2 mediated increase in global decay rates that could potentially contribute to fundamental global responses such as inhibition of translation. Finally, our data provides the first quantitative description of the effects of hypoxia on mRNA half-lives at global scale.

MATERIALS AND METHODS

Cell culture and treatments

Human umbilical vein endothelial cells (HUVEC) from pools of donors were purchased from Lonza (Lonza, C2519A) and grown in Endothelial Medium Bullet Kit (EGM-2, Lonza, CC-3162). For viral production, HEK 293T were maintained in Dulbecco's modified Eagle's medium (Gibco, 41966052) supplemented with 50 U/mL penicillin (Gibco, 15140122), 50 µg/mL streptomycin (Gibco, 15140122), 2 mM glutamine (Gibco, 25030123), and 10% (v/v) fetal bovine serum (Gibco, 10270106). All cells were grown at 37°C and 5% CO₂ in a humidified incubator and tested regularly for mycoplasma contamination. For hypoxia treatment, cells were grown at 37°C in a 1% O₂, 5% CO₂, 94% N₂ gas mixture in a Whitley Hypoxystation H35 (Don Whitley Scientific). Alternatively, hypoxic conditions were mimicked using 500 µM Dimethylxylglycine (DMOG, Sigma, D3695). Actinomycin, cycloheximide, puromycin and rapamycin purchased from Sigma, were used at 5 µg/mL, 10 µg/mL, 2 µg/mL, and 20 nM, respectively.

Metabolic labeling with 4-thiouridine and purification of newly synthesized mRNA

We used the protocol described by Dolken et al. (2008) and Schwanhauser et al. (2011) that is represented in Figure 1A. Briefly, exponentially growing HUVEC cells were exposed to 21% (normoxia) or 1% (hypoxia) oxygen for 8 h and pulse labeled with 4-thiouridine (400 µM, 4sU, Sigma, T4509) during the last 2 h of treatment. After treatment, total cellular RNA was isolated from cells using TRI-reagent (Ambion, AM9738). An amount of 100 µg of total RNA was subjected to a biotinylation reaction to label the newly transcribed RNA containing the 4sU moiety (Pierce, EZ-Link Biotin-HPDP, 21341). Then, RNA was purified using Ultrapur Phenol:Chloroform:Isoamylalcohol (Invitrogen, 15593 031) and labeled RNA was isolated from the total RNA by affinity chromatography using streptavidin coated magnetic beads (µMacs Streptavidin Kit; Miltenyi, 130074 101).

High-throughput sequencing and bioinformatics analysis

Libraries of total RNA, labeled RNA and unlabeled RNA were prepared using the standard protocol for messenger RNA sequencing (Illumina, TruSeq Stranded mRNA) and sequenced on HiSeq2000 instrument (Illumina) according to the manufacturer's protocol. The resulting reads were aligned to the GRCh37/hg19 assembly of the human genome using TopHat (Trapnell et al. 2009) with the default parameters. Finally, the gene expression level was calculated as the number of reads per gene, computed using HTSeq (Anders et al. 2015) and gene features as defined in the GRCh37.75 release of the human genome (gtf file). Differential gene expression analysis was performed with the Bioconductor (Gentleman et al. 2004) edgeR package (Robinson et al. 2010) for the R statistical Software (<http://www.R-project.org/>). The raw reads and processed data derived from RNA-seq experiments published in Tiana et al. (2018) are available at NCBI's Gene Expression Omnibus (Barrett et al.

2009) and are accessible through the GSE89831 GEO Series accession number. All computations were performed using R software package (<http://www.R-project.org/>).

mRNA half-life determination

Individual mRNA half-lives were calculated as described in Dolken et al. (2008). For each mRNA half-life was computed according to the following equation:

$$t_{1/2} = -\frac{t_L \times \ln(2)}{\ln\left(1 - \frac{e}{a} \times C_e\right)},$$

where e and a represent the amount of newly transcribed RNA and total mRNA, respectively, expressed as the normalized counts from the RNA-seq experiment. t_L is the labeling time and C_e is a correction factor estimated from slope of the linear regression fit of the u/a and e/a values for all the genes expressed in HUVEC according to the formula:

$$\frac{u}{a} = \frac{1}{C_u} - \frac{e C_e}{a C_u}.$$

In this equation u represents the amount of preexisting mRNA expressed as the normalized counts from the RNA-seq experiments. The C_e and C_u correction factors can be determined by the estimate of the slope and intercept of the linear fit of the expression level of all three fractions (total, newly transcribed and preexisting). All computations were performed using R software package.

RNA extraction and qPCR

RNA was extracted and purified using the RNeasy Mini Kit (Qiagen; 74106) following manufacturer's instructions. For quantitative-RT PCR analysis, 1 µg of total RNA of each sample was reversed-transcribed to cDNA (Transcriptor First Strand cDNA Synthesis kit, Roche 04379012001), cDNA was diluted 1:20 and used as template for amplification reactions, carried out with the Power SYBR green PCR Master Mix (RNeasy Mini Kit Applied Biosystems; 4367659) following manufacturer's instructions. PCR amplifications were carried out in a StepOne Realtime PCR System (Applied Biosystems; 4376357). Data were analyzed with StepOne software and expression levels were calculated using $\Delta\Delta CT$, using β -actin as reference. Primers sequences used for qPCR analysis are provided in Table 1. All RT-qPCR data are available in Supplemental Table 3

Individual mRNA half-life calculation using actinomycin D

HUVEC cells were seeded in 6 cm² plates at a density of 4×10^5 cells/well 8 h prior to experiments. To determine the effect of hypoxia on mRNA decay rates, cells were exposed to normoxia or hypoxia for 16 h. Then, treated with 5 µg/mL actinomycin D (Sigma; A9415), a general transcription inhibitor, and RNA was isolated 0, 1, 2, 4, 8, and 12 h after transcription inhibition. Cells were kept at the same oxygen tension for the whole duration of the experiment. The relative amount of specific mRNAs were determined by quantitative RT-PCR. Their decay constant (k_{decay})

was obtained from the slope of the regression line obtained by the representation of the fraction of mRNA remaining at each time point:

$$\ln\left(\frac{N_t}{N_0}\right) = -K_{\text{decay}} \times t.$$

N_t/N_0 is the remaining fraction of mRNA at time point t after actinomycin D addition. Finally, the half-life for each mRNA was calculated from its decay constant as:

$$t_{1/2} = \frac{\ln(2)}{K_{\text{decay}}}.$$

Pulse-labeling of HUVEC cells with tritiated uridine

HUVEC cells were seeded in 6 cm² plates (4×10^5 cells/well) and 8 h after plating were pulsed for 2 h with tritiated uridine (Uridine [5,6-³H]; PerkinElmer; NET367001MC) at final concentration of 2 µCi/mL. After labeling, cells were washed twice with phosphate buffered-saline (PBS), and incubated in nonradioactive complete culture medium under normoxia or hypoxia for 16 h. The radioactivity present in total RNA fractions was measured using Liquid Scintillation (Optiphase Hisafe II; PerkinElmer, 1200-436) and the Liquid Scintillation Analyzer (PerkinElmer; TriCarb 2800TR).

rRNA synthesis determination

The determination of the 47S precursor ribosomal RNA (pre-RNA), was performed by qPCR and normalized by 28S using the primers described in Table 1.

Calculation of total RNA and mRNA mass per cell

HUVEC cells were seeded in 6 cm² plates at a density of 4.8×10^5 cells per plate and were exposed to normoxia, hypoxia or normoxia in the presence of 500 µM DMOG for 16 h. Then DNA and RNA were isolated from each sample and determined by spectrophotometry (NanoDrop ND-1000 instrument, Thermo Scientific). We calculated RNA mass per cell considering that human cells contain 6.6 pg of DNA per cell.

Lentiviral production

Lentiviral vector pGIPz (Open Biosystems) was used to silence the mRNA expression of *ARNT* (V3LHS_393533), *HIF1A* (V2LHS_132150), *EPAS1* (V2LHS_113753) and *TSC2* (V3LHS_642078), as well as a nonsilencing lentiviral shRNA Control (NS; RHS4346). For the overexpression of HIF mutants stabilized in normoxia, we used PRRL lentiviral vectors containing *HIF1A* and *EPAS1* mutant sequences under cytomegalovirus (CMV) promoter (Kondo et al. 2002). These vectors contain the ODD prolines residues converted to alanine for maintaining HIF constitutively active in normoxic conditions (pRRL-HIF1A-P402A; P564A-IRES-eGFP and pRRL-EPAS1-P405A;P531A-IRES-eGFP). In addition, we used other HIF overexpressing vectors whose bHLH motif had been mutated to prevent DNA-binding (pRRL-HIF1A-P402A;P564A-bHLH* -IRES-eGFP and pRRL-EPAS1-P405A;P531A-bHLH* -IRES-eGFP). Lentivirus were produced and

titrated in HEK293T cells as previously described (Punzon et al. 2004). For transduction of HUVEC cells, lentiviruses were used at a multiplicity of infection (MOI) of 1–2 for 8 h, resulting in more than 95% transduced (GFP-positive) cells 72 h after infection.

Western blot

Cells were harvested in RIPA buffer (50 mM Tris-HCl pH 8, 150 mM NaCl, 0.02% NaN₃, 0.1% SDS, 1% NP-40, 1% Sodium Deoxycholate) containing protease inhibitors (Complete ULTRA tablet; Roche; 06538304001). Protein concentration was quantified using BioRad DC protein assay (BioRad, 5000112) and 40 µg of each sample was resolved in 10% SDS-polyacrilamide gels. Proteins were then transferred to Polyvinylidene difluoride (PVDF) membrane (Immobilon-P, Millipore, IPVH100010). Membranes were blocked with 5% nonfat dry milk in TBS-T (50 mM Tris-HCl pH 7.6, 150 mM NaCl, 0.1% Tween-20) and incubated with the corresponding antibody (*HIF1A*: abcam; ab2185; *EPAS1*: Novus; NB100-122, *LaminB1*: Santa Cruz, sc-6216, *Phospho-S6 Ribosomal Protein* [Ser235/236], Cell Signaling, 4858; and *S6 Ribosomal Protein*, Cell signaling, 2217).

Methionine S-35 pulse-labeling of HUVEC cells

HUVEC cells were seeded in 6 cm² plates at a density of 4.8×10^5 cells per plate and exposed to normoxia, hypoxia or normoxia in the presence of 500 µM DMOG for 16 h. Then, cells were washed with DMEM lacking methionine and containing 10% of dialyzed FBS to eliminate the excess of methionine and 25 mM Hepes. Medium was removed and 0.5 ml of DMEM with ³⁵S-35 methionine (200 µCi/mL) was added, then cells were incubated at 37°C 30 min. Cells were washed with cold PBS and recovered by cell-scraping and centrifugation and precipitated with 10% TCA in glass fiber filter disks. To calculate the ratio of TCA-precipitated label relative onto total radioactivity, we measured the amount of radioactivity of each disk by scintillation counting and compared with the total radiolabeled cell suspension fraction obtained before precipitation.

Click-iT Plus OPP Alexa Fluor 647 Protein Synthesis Assay Kit

HUVEC cells were seeded over coverslips in 24 multiwell plates at a density of 50,000 cells per well, and treated with cycloheximide or DMOG for 16 h. Then we used the Click-iT Plus OPP Alexa Fluor 647 Protein Synthesis Assay Kit following manufacturer's instructions. The analysis of the images was performed by measuring individual cell integrated density of the 647 channel divided by the DAPI integrated density.

SUPPLEMENTAL MATERIAL

Supplemental material is available for this article.

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