Departure from optimal O₂ level for mouse trophoblast stem cell proliferation and potency leads to most rapid AMPK activation

Yu YANG¹, ², Zhongliang JIANG¹, Alan BOLNICK¹, Jing DAI¹, Elizabeth E PUSCHECK¹ and Daniel A RAPPOLEE¹–⁴)

¹Ob/Gyn, Wayne State University Medical School, Detroit, MI 48201, USA
²Department of Physiology, Wayne State University Medical School, Detroit, MI 48201, USA
³Institutes for Environmental Health Science, Wayne State University School of Medicine, Detroit, MI 48201, USA
⁴Department of Biology, University of Windsor, Windsor, ON N9B 3P4, Canada

Abstract. Previous studies showed that cultured mouse trophoblast stem cells (mTSCs) have the most rapid proliferation, normal maintenance of stemness/potency, the least spontaneous differentiation, and the lowest level of stress-activated protein kinase (SAPK) when incubated at 2% O₂ rather than at the traditional 20% O₂ or hypoxic (0.5% and 0% O₂) conditions. Switching from 2% O₂ induced fast SAPK responses. Here we tested the dose response of AMP-activated protein kinase (AMPK) in its active form (pAMPK) at O₂ levels from 20–0%, and also tested whether pAMPK levels show similar rapid changes when mTSC cultures were switched from the optimal 2% O₂ to other O₂ conditions. There was a delayed increase in pAMPK levels ~6–8 h after switching conditions from 20% to 2%, 0.5%, or 0% O₂. Altering O₂ conditions from 2% to either 20%, 0.5%, or 0% led to rapid increase in pAMPK levels within 1 h, similar to the previously reported SAPK response in mTSC cells removed from 2% O₂. Twelve hours of 0.5% O₂ exposure led to cell program changes in terms of potency loss and suppressed biosynthesis, as indicated by levels of phosphorylated inactive acetyl CoA carboxylase (pACC). Phosphorylation of ACC was inhibited by the AMPK inhibitor Compound C. However, unlike other stressors, AMPK does not mediate hypoxia-induced potency loss in mTSCs. These results suggest an important aspect of stem cell biology, which demands rapid stress enzyme activation to cope with sudden changes in external environment, e.g., from least stressful (2% O₂) to more stressful conditions.

Key words: AMP-activated protein kinase (AMPK), Hypoxia, Stress, Trophoblast stem cell

Stemness can be maintained at an O₂ niche ≤ 5%, and often at 2–3% [12–14]. It was previously established that 2% O₂ is the optimal O₂ level for mTSC in vitro culture by four criteria: lowest stress (SAPK activation) level, lowest expression of differentiation maker mRNAs, highest growth rate and normal maintenance of potency [15]. Stressors force stem cell differentiation, which has been observed in ESCs and induced pluripotent stem cells [16, 17]. Stress-induced differentiation has also been characterized in mTSCs [18]. In screens for the protein kinases that mediate the stress response of mTSCs, many kinases inhibitors were used; it was found that stress-induced differentiation is mediated through SAPK, which does not affect potency, and that AMPK mediates potency loss [5, 19]. SAPK mediates increased levels of Hand1 mRNA, favoring giant cell differentiation and placental lactogen 1 (PL1) expression, and suppressing later chorionic lineages by decreasing levels of Gcm1 mRNA [11, 20]. PL1 is the hormone that mediates maternal recognition of pregnancy in rodents [21]; this makes it the functional equivalent of chorionic gonadotropin in human, and of interferon-like protein in sheep and cattle [22].}

Received: July 26, 2016
Accepted: October 31, 2016
Published online in J-STAGE: November 18, 2016
©2017 by the Society for Reproduction and Development
Correspondence: DA Rappolee (e-mail: drappole@med.wayne.edu)
This is an open-access article distributed under the terms of the Creative Commons Attribution Non-Commercial No Derivatives (by-nc-nd) License <http://creativecommons.org/licenses/by-nc-nd/4.0/>.
to the much slower rates of SAPK activation when O2 levels were switched from 20% to other amounts.

Our hypothesis is that stress induces fast changes in the activity of stress kinases, and that they consequently function to adjust developmental and metabolic programs. Rapid turnover is a feature of many intracellular regulatory and signaling proteins; it enables prompt responses to extracellular or intracellular signals, and rapid cessation of responses upon signal removal. Examples of this include the products of proto-oncogenes, growth factors and inflammatory cytokines [23, 24]. The major regulator of intracellular AMPK activity is the reversible phosphorylation of threonine 172 (Thr172) within the protein’s catalytic α subunit, which activates AMPK [25]. Not surprisingly, AMPK activity also has fast turnover [26]. The level of pAMPK (phosphorylation of AMPKα at Thr172) is often used to indicate AMPK activity [27], and it corresponds with the phosphorylation of its canonical metabolic substrate acetyl CoA carboxylase (ACC Ser79) [28, 29]. ACC catalyzes a rate-limiting phosphorylation of its canonical metabolic substrate acetyl CoA carboxylase (ACC Ser79) [28, 29]. ACC catalyzes a rate-limiting phosphorylation, and that AMPK is responsible for this outcome.

Materials and Methods

Reagents

Fetal bovine serum, RPMI1640, and fibroblast growth factor 4 (FGF4) were from Gibco (Grand Island, NY, USA). Heparin was purchased from Sigma Chemical (St. Louis, MO, USA). Compound C was purchased from EMD Millipore (Cat# 171260; Billerica, MA, USA). The following antibodies used were from Cell Signaling Technology (Danvers, MA, USA): pAMPK (CS 2535), pACC (Ser79) (CS 3661), β-Actin (CS 4790), anti-rabbit HRP-linked antibody (CS 7074), and anti-mouse HRP-linked antibody (CS 7076). Tubulin (Cat# T 9026, St. Louis, MO, USA) antibody came from Sigma Chemical Co. ErrB and ID2 antibody were purchased from Santa Cruz (pAMPK (CS 2535), pACC (Ser79) (CS 3661), β-Actin (CS 4790), anti-rabbit HRP-linked antibody (CS 7074), and anti-mouse HRP-linked antibody (CS 7076). Tubulin (Cat# T 9026, St. Louis, MO, USA) antibody came from Sigma Chemical Co. ErrB and ID2 antibody were purchased from Santa Cruz Biotechnology (SC-489; Dallas, TX, USA), respectively. Anaerobic bags to create 0% O2 were from Hardy Diagnostics (AN010C; Santa Maria, CA, USA).

Cell lines and culture conditions

The mouse trophoblast stem cell isolate was a gift from Dr. Rossant (Samuel Lunenfeld Research Institute, Ontario, Canada). mTSCs were cultured as described previously [20, 33]. Routine culture conditions were 20% O2, with 25 ng/ml FGF4 and 70% embryonic fibroblast conditioned medium. The cells were passaged approximately 24 h before the start of each experiment to allow recovery from passage stress. In the group of experiments where cells were switched from 20% O2 to other O2 levels, post-passage culturing was conducted at 20% O2. Alternately, if cells were planned to be switched from 2% O2 to other O2 levels, they were placed in 2% O2 for 24 h. The starting cell confluence was around 20–30% prior to the switch. After switching, cells were cultured for various length of time and were then lysed for immunoblot analysis. Culture at 20% O2 was conducted in a conventional CO2 incubator; 2% and 0.5% O2 conditions were achieved using commercial gas mixtures containing [2% O2/5% CO2] or [0.5% O2/5% CO2], balanced with N2. All culture media were pre-equilibrated for 24 h, at the specified O2 levels before use.

Results

During the study of SAPK activation, we found that 2% O2 enabled a growth rate that was 2.5-fold higher than that of 20% O2 for mTSC culture in vitro, but the media was very acidic by 24 h [15]. Therefore, in this experiment, the final duration of time after removing cells from 20% O2 was set to 12 h. The switch from 20% to lower O2 levels was designed to emulate the changes that might occur during re-implantation of in vitro cultured embryos...
DEPARTING 2% O₂ RAPIDLY ACTIVATES AMPK

The kinetics of AMPK activation were investigated after changing from 20% O₂ to 2% O₂ (Fig. 1A), to 0.5% O₂ (Fig. 1B), or to 0% O₂ (Fig. 1C). There were consistent increases in pAMPK levels, starting by 6 h, in all O₂ groups. For the 20% to 2% O₂ switch, pAMPK levels fluctuated ~4-fold throughout the 12 h period, after reaching this level by 6 h (Fig. 1A). After switching to 0.5% or to 0% O₂, pAMPK levels continued to increase, with the peak observed at 12 h (Fig. 1B, Fig. 1C). Total AMPK protein levels did not change after the three sets of switch from 20% O₂ during the period studied (Supplementary Fig. 1: online only). Thus, removing cells from 20% O₂ produced consistent increases in pAMPK levels, starting at 6 h independent of what O₂ level the cells were moved into.

Because there was rapid proliferation and metabolic waste accumulation when cells were cultured at 2% O₂ [15], we took advantage of AMPK as a reporter of metabolic stress to evaluate how often the culture medium should be changed when cells are cultured at 2% O₂. The result showed that changing to 2% O₂ did not activate AMPK by 1–3 h, but pAMPK levels became high between 6–8 h, and maintained high levels throughout the 12 h period (Fig. 2). These results suggest that the intake of nutrients and the accumulation of metabolic waste may have already become evident, and been sensed by the cells, after 6–8 h of 2% O₂ culture. Without microfluidic equipment to provide constant nutrient support and waste removal, it would be difficult to routinely culture mTSCs at 2% O₂. Currently in vitro culturing of mTSCs is commonly carried out at 20% O₂ [33], and this does not pose any problems for the isolation, maintenance, or in vivo differentiation capabilities of mTSCs [34].

It should be noted that even though changing cells from 20% O₂ to 2%, 0.5%, and 0% O₂ conditions all produced higher pAMPK levels after 6–8 h, the biological processes underlying these increases are unlikely to be the same. It is likely that the high pAMPK levels observed after 6–8 h of 2% O₂ reflect the metabolic needs initiated by rapid cell proliferation. There was minimal net cell growth in the 0.5% and 0% O₂ groups, as reported previously for mTSCs [15]. Increased pAMPK levels supposedly report other signals raised by hypoxic stress, other than the need of biosynthesis for cell division in these conditions. Further studies are needed to investigate the mechanisms underlying pAMPK level increases in each condition.

To test the hypothesis that moving away from the least stressful 2% O₂ condition would also produce faster AMPK activation (as was observed for SAPK), the kinetics of AMPK activation were studied within a time frame similar to that used in studies of SAPK [15]. Time points of 0.5, 1, 2, 3, and 24 h were chosen. Four early time points (0.5, 1, 2, 3 h) were selected to detect when AMPK activation first occurs. The period of 24 h after removing cells from 2% O₂ was also studied to enable comparisons between our data and the published SAPK data. The results showed that pAMPK levels were near maximum after removing cells from 2% O₂ for 1 h, regardless of the O₂ level that the cells were moved into. The results of switches from 2% O₂ to either 20%, 0.5%, or 0% O₂ are shown in Figs. 3A, 3B, and 3C, respectively. Total AMPK protein levels did not change after the three sets of switch from 2% O₂ during the period studied (Supplementary Fig. 2: online only). Interestingly, the 2% to 20% O₂ switch increased pAMPK levels significantly at 1–3 h, but by 24 h the pAMPK levels returned to baseline (Fig. 3A). This indicates

**Fig. 1.** Switching O₂ conditions from 20% to 2%, 0.5%, or 0% led to increases in pAMPK levels at 6 h. mTSCs were cultured as indicated, lysed, fractionated by SDS-PAGE, and probed with antibodies against pAMPK Thr172. ACTB was used as a loading control. Histograms show the average pAMPK level of three independent biological experiments, with error bars indicating standard error. A) Change from 20% to 2% O₂ led to a delayed increase in pAMPK levels, to ~4-fold over baseline at 6 h; pAMPK remained at these higher levels until 12 h. B) Change from 20% to 0.5% O₂ led to an increase in pAMPK levels, to ~5-fold above baseline at 6 h, and reached a peak of ~10-fold at 12 h. **∗∗** indicates statistical significance compared with time zero at 20% O₂. C) After changing from 20% to 0% O₂, pAMPK levels increased to ~5-fold at 6 h, and reached a peak of ~9-fold over baseline, at 12 h. **∗∗∗** indicates statistical significance compared with time zero at 20% O₂.
that changing cells from the least stressful condition of 2% O2 to the more stressful 20% O2 required a rapid AMPK response. However, unlike in the stressful O2 levels below 2%, cells eventually adapted to 20% O2, and their pAMPK levels decreased.

The maximal stimulation index in the switches from 2% to other O2 levels was only ~2–3 fold, whereas it was ~8–10 fold following the switches away from 20% O2, as shown in Fig. 1. This was the case even when the terminal O2 level following the switch was the same. This is most likely due to the difference in pAMPK levels observed between 2% and 20% O2 at the 0 h baseline, prior to the switch. Because 2% O2 facilitates faster cell proliferation than 20% O2, it is possible that the high metabolic needs of mTSCs cultured at 2% O2 prior to O2 change have already led to a higher pAMPK baseline. As a result, the relative fold changes in pAMPK levels from baseline were smaller after cells were moved away from 2% O2 (Fig. 3).

In Fig. 4A, we superimposed the dynamics and magnitude of pAMPK levels, based on data represented in Fig. 1 and Fig. 3. This shows average pAMPK levels after 12 h of different O2 treatment compared to the 0 h baseline. pAMPK levels were lowest at 20% O2 and highest at 0.5–0% O2, with an approximate 10-fold change. pAMPK levels at 2% O2 were between those of the 20% and 0.5–0% O2 conditions, which produced an S-shaped curve. The speed of pAMPK (Fig. 4A) activation after moving cells away from 20% or 2% O2 to other O2 levels mimicked the pSAPK response observed in a similar experimental condition (Fig. 4B), which was cited with permission from a previous publication [11]. Sudden changes in O2 environments, from the least stressful 2% to either higher or lower O2 levels, induced rapid AMPK and SAPK activation at 1 h. When cells were moved from 20% O2 to 0% O2, the levels of pAMPK and pSAPK increased at 6–8 h.

The stem cell state of mTSCs is characterized by the expression of potency factors. Loss of potency predisposes mTSCs to differentiation. We investigated the biological consequences of hypoxic exposure at 0.5% O2 on the levels of two mTSC potency factors: inhibition of differentiation 2 (ID2), and estrogen-related receptor beta (ERRB) (Fig. 5A). The time point of 12 h was chosen because, in 0.5% O2 conditions, pAMPK reached peak levels at 12 h. In addition, to study whether inhibition of AMPK could reverse potency loss, we needed to choose a time point when potency loss had already occurred. In the 0.5% O2 culture, there was no appreciable potency loss in the two time points (4 h and 8 h) prior to 12 h (Supplementary Fig. 3: online only). mTSC incubated at 0.5% O2 for 12 h led to a significant
DEPARTING 2% O₂ RAPIDLY ACTIVATES AMPK

AMPK mediates profound changes in mTSC metabolic regulation and stemness/differentiation balance [11]. It is not surprising that AMPK was activated rapidly when cultured mTSCs were from optimal 2% O₂ conditions to hyperoxic 20% O₂ or hypoxic 0.5% O₂. The highest stimulation index of both pAMPK and pSAPK occurred at 0–0.5% O₂ (Fig. 4), suggesting that hypoxia at O₂ conditions below 2% is more stressful for mTSCs than that at the 20% O₂ condition. Because the stimulus response of an enzyme is a product of speed (i.e., the “direness” index) and magnitude (i.e., the “stimulation” index), the most powerful AMPK response occurred when mTSCs were moved from 2% to below 2% O₂. It is interesting that switching mTSCs from the less ideal 20% O₂ environment induced slower activation of AMPK at 6–8 h, while switching them from the least stressful 2% O₂ environment induced fast AMPK activation. We call the similar kinetic patterns of the relatively faster AMPK and SAPK responses a “direness response” when cells were moved out of the optimal 2% O₂ environment. The speed of change in kinase activation is based on AMPK sensing the dissimilar cellular state at starting 2% or 20% O₂, not the final AMPK activity state at 0% or 0.5% O₂ (which is reflected by the similar stimulation index). The speed with which stress enzymes were activated after cells were moved out of the least stressful 2% O₂ condition may reflect the profound stress initiated by deviating from this ideal cellular condition. In times of noxious environmental stimuli, cells must quickly change programs to adapt, or else they die. AMPK and SAPK play important roles in sensing environmental cues and determining cell fate [35, 36].

Although both AMPK and SAPK are stress kinases, they do not always respond to stress in the same way [37]. SAPK is ubiquitously expressed, and is activated by multiple types of stress, including UV radiation, hyperosmolality, ischemia/reperfusion injury, and stimulation by TNF-α [11]. SAPK participates in intracellular signaling pathways that control cell proliferation, differentiation, apoptosis, cytoskeletal integrity, and other functions, as has been reviewed in [35]. By contrast, AMPK is first and foremost a kinase — with the primary role of maintaining ATP balance by regulating anabolic and catabolic metabolism [1]. The relatively higher level of pAMPK observed at 2% O₂ may be due to the depletion of energy substrates and/or the accumulation of acidic, metabolic wastes; alternatively, it may have occurred in anticipation of the energy requirements of rapid cell proliferation [38]. At 2% O₂, the increase in pACC level was not accompanied by the loss of potency, supporting the new state that the cells must respond to; this is in contrast with the outcome for cells in 0.5% O₂. Further studies are needed to elucidate the mechanism of AMPK activation that occurs when cells are moved into the least stressful 2% O₂ environment.

Many kinases demonstrate early and late activation, and mediate active metabolism rather than pathologic hypoxia. This was because, unlike that observed in cells cultured at 0.5% O₂ after 12 h of 2% O₂ treatment, the levels of potency factors were comparable to those of the 20% O₂ control group. The increase in pACC levels observed at 0.5% O₂ was reversed by ~50% via introduction of the AMPK inhibitor compound C, while the increase in pACC level observed at 2% O₂ was reversed completely by compound C.

Discussion

Fig. 4. O₂ stresses induced an S-shaped curve for pAMPK levels, and a U-shaped curve for pSAPK levels, but activation of both enzymes was most rapid when O₂ was changed from the least stressful condition at 2% O₂. A) Summary of the pAMPK dose and kinetic responses to changes in O₂ levels, based on the data presented in Figs. 1–3. The tail of the arrow is the O₂ level at time zero before the change in O₂ conditions, and the head of the arrow is the level of O₂ into which the cells were moved. mTSCs responded to culture at 20%, 2%, 0.5%, and 0% O₂ with an S-shaped pAMPK dose-response curve, with a maximal increase of ~10-fold over baseline after 12 h of culture. B) Summary of the SAPK dose and kinetic responses to changes in O₂ levels, based on a published figure we cite with permission [11]. The tail of the arrow is the O₂ level at time zero before the change in O₂ conditions, and the head of the arrow is the level of O₂ into which the cells were moved. pSAPK levels were lowest at 2% O₂ and increased rapidly within 1 h, when the cells were removed from 2% O₂. The change from 20% to 0% O₂ produced a slower activation of SAPK.

(-50%) loss of ERRB and a not significant (-35%) loss of ID2. AMPK inhibitor compound C did not reverse the loss of ERRB caused by hypoxic stress. Thus, the hypothesis that stress induces potency loss was supported. However, unlike potency loss caused by other stressors, inhibiting AMPK cannot reverse potency loss induced by hypoxic stress. Consistent with a previous report [15], mTSCs maintained their potency at 2% O₂ (Fig. 5B). After 12 h of 2% O₂ culture, levels of ERRB and ID2 were comparable to those observed in the 20% control group at 0 h. Unexpectedly, adding 10 µM compound C to 2% O₂ culture significantly decreased the level of ID2 by approximately 45%. There were significant increases (~4-fold and 2.7-fold) in the levels of ACC (Ser79) phosphorylation after 12 h of 0.5% and 2% O₂ culture, respectively. These increases in pACC are consistent with the increases in pAMPK shown in Fig. 2, which are more likely to reflect the needs of cells engaged in
distinct downstream events at different activation times [39–42]. During a microarray study of mTSC responses to hyperosmolar stress, we found that the early stress response (within 30 min of sorbitol treatment) is to downregulate highly changing mRNAs (all 31 genes with significant change were downregulated). However, by 24 h, 158 genes were upregulated and 130 were downregulated, including genes involved in the cell cycle, apoptosis, macromolecular synthesis, and differentiation [43]. The direct effects of AMPK were not investigated in the microarray study. Because AMPK is activated under hyperosmolar stress [4], it is likely to have a role in the hyperosmolar stress-induced changes that cells undergo. If rapid and early responses to stress have been successful, cells may regain their balance after the stress subsides; or if the stress persists, larger-scale changes in transcription, cell cycle, and differentiation may follow, leading to irreversible changes in cell program. In mTSCs, mESCs, and early mouse embryos under hyperosmotic and genotoxic stress, as well as to certain drugs or diet supplements, AMPK downregulates cellular potency factors [4, 5] and predisposes the cells to differentiation. The fact that moving cells from 2% O2 to 20% O2 induced early increases in pAMPK levels that eventually fell back to baseline is informative. This suggests that cells are capable of coping with sudden environmental changes and that if the new environment is not too stressful, they regain their balance.

To further understand the biological consequences of hypoxic stress and AMPK activation, we studied the effects of an AMPK inhibitor, and the levels of two potency factors, ID2 [44] and ERRB [34, 45], in cells cultured at traditional 20% O2, 0.5% O2, or 2% O2 conditions. ID2 is an mTSC stemness marker, and is a key potency-maintenance gene. Forced expression of proteins in the ID family inhibits the differentiation of human cytotrophoblasts [44]. ERRB is also a mTSC stemness marker, and it is involved in chorionic lineage specification and potency maintenance following implantation [45]. With FGF4 removal, normally differentiated mTSCs lose expression of ID2 and ERRB, which is consistent with their role as potency factors [20, 34]. Hypoxic stress at 0.5% O2 also drove ID2 and ERRB downregulation, despite the presence of the potency-maintaining growth factor FGF4; however, AMPK inhibition did not reverse potency loss at 0.5% O2. Unlike the minimal effects that compound C had on the levels of ERRB and ID2 in cells cultured at 20% O2 and 0.5% O2, both potency factors were reduced by compound C in cells at 2% O2 (although the decrease in ERRB did not reach statistical significance). We know that AMPK activity was inhibited by compound C at 2% O2, because pACC (Ser79) level was significantly reduced. ACC carries the classical AMPK substrate motif [46], and the level of ACC phosphorylation at Ser79 indicates the activity of AMPK. However, in addition to being an AMPK inhibitor, compound C is known to have AMPK-independent effects on multiple cellular processes, such as cell cycle progression.

Fig. 5. Hypoxia at 0.5% O2 significantly increased the pACC levels, and decreased the levels of potency factor ERRB. mTSCs were cultured for 12 h at 20% O2, 0.5% O2, or 2% O2, with or without 10 μM compound C. Increased pACC levels due to 0.5% O2 or 2% O2 exposure were mitigated by compound C; however, the loss of ERRB protein at 0.5% O2 was not reversed. “*” indicates statistical significance compared with time zero at 20% O2.
irreversible loss of their potency factor proteins. Occasional stressors exist, and AMPK functions are needed. Knockdown of AMPK catalytic α subunits leads to reduced cell culture, and AMPK sits at the center of cellular energy regulation by affecting lipid synthesis [11]. The unique feature of hypoxic stress is a generalized phenomenon, and future studies are needed to test these hypotheses.

**Limitations of this study, and future directions**

Here, we studied the dynamics of AMPK activation and found that moving cells out of 20% O₂ conditions activated AMPK at a slower rate than when they were moved out of the least stressful conditions at 2% O₂. We did not study the upstream events that mediate the early (1 h) and late (6–8 h) AMPK response, nor did we study how the 2% O₂ environment differs from that of 20% O₂ in causing these changes. Because 2% O₂ is associated with faster mTSC proliferation (~7 h doubling time), anabolic processes such as transcription, translation, and DNA replication should also be more active at 2% O₂. Bacterial studies showed that rapidly growing cells operate close to their optimal energy efficiency [51], which is necessary to support the high rates of biosynthesis during cell proliferation [52, 53]. We speculate that the demands of high energy turnover that are associated with rapid cell growth may make mTSCs more susceptible to perturbations in the environment. An alternate hypothesis is that healthier cells, cultured in the least stressful environment, are inherently more capable of sensing stress and mounting a rapid adaptive response. Different cell types and stimuli should be used to gain better understanding of whether this is a generalized phenomenon, and future studies are needed to test these hypotheses.

**Acknowledgments**

Thanks to Drs Husam Abu-Soud and Awniyi Awnouga for analysis and comments on the manuscript. This research was supported by grants to DAR from NIH (1R03HD061431 02) and from the Office of the Vice President for Research at Wayne State University.

**References**


15. Zhou S, Xie Y, Puscheck EE, Rappolee DA. Oxygen levels that optimize TSC culture are identified by maximizing growth rates and minimizing stress. Placenta 2011; 32: 475–481. [Medline] [CrossRef]


45. Xie Y, Abdallah ME, Awonuga AO, Slater JA, Puscheck EE, Rappolee DA. Benzo(a) pyrene causes PRKAAR1/2-dependent ID2 loss in trophoblast stem cells. Mol. Reprod. Dev. 2010; 77: 533–539. [Medline] [CrossRef]


52. Susa M, Olivier AR, Fabbaro D, Thomas G. EGF induces biphasic S6 kinase activation: late phase is protein kinase C-dependent and contributes to mitogenesis. Cell. 1989; 57: 817–824. [Medline] [CrossRef]


64. Maitra A, Dill KA. Bacterial growth laws reflect the evolutionary importance of energy efficiency. Proc. Natl. Acad. Sci. USA 2015; 112: 406–411. [Medline] [CrossRef]
