Collagen metabolism as a regulator of proline dehydrogenase/proline oxidase-dependent apoptosis/autophagy

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Abstract
Recent studies on the regulatory role of amino acids in cell metabolism have focused on the functional significance of proline degradation. The process is catalysed by proline dehydrogenase/proline oxidase (PRODH/POX), a mitochondrial flavin-dependent enzyme converting proline into Δ1-pyrroline-5-carboxylate (P5C). During this process, electrons are transferred to electron transport chain producing ATP for survival or they directly reduce oxygen, producing reactive oxygen species (ROS) inducing apoptosis/autophagy. However, the mechanism for switching survival/apoptosis mode is unknown. Although PRODH/POX activity and energetic metabolism were suggested as an underlying mechanism for the survival/apoptosis switch, proline availability for this enzyme is also important. Proline availability is regulated by prolidase (proline supporting enzyme), collagen biosynthesis (proline utilizing process) and proline synthesis from glutamine, glutamate, α-ketoglutarate (α-KG) and ornithine. Proline availability is dependent on the rate of glycolysis, TCA and urea cycles, proline metabolism, collagen biosynthesis and its degradation. It is well established that proline synthesis enzymes, P5C synthetase and P5C reductase as well as collagen prolyl hydroxylases are up-regulated in most of cancer types and control rates of collagen biosynthesis. Up-regulation of collagen prolyl hydroxylase and its exhaustion of ascorbate and α-KG may compete with DNA and histone demethylases (that require the same cofactors) to influence metabolic epigenetics.
This knowledge led us to hypothesize that up-regulation of prolidase and PRODH/POX with inhibition of collagen biosynthesis may represent potential pharmacotherapeutic approach to induce apoptosis or autophagic death in cancer cells. These aspects of proline metabolism are discussed in the review as an approach to understand complex regulatory mechanisms driving PRODH/POX-dependent apoptosis/survival.

Keywords Apoptosis · Autophagy · Collagen metabolism · Proline · Proline dehydrogenase/Proline oxidase · Prolidase · Signalling

Introduction
Increasing evidence suggests that proline and collagen metabolism may determine cancer development and progression. Specifically, proline availability determined by collagen biosynthesis/degradation ratio and endogenous synthesis of this amino acid influences redox balance, DNA biosynthesis, epigenetic modifications, apoptosis and autophagy (D’Aniello et al. 2020). Proline degradation by proline dehydrogenase/proline oxidase (PRODH/POX) drives apoptosis, autophagy or survival, depending on the metabolic context. The understanding of the mechanism for the differential PRODH/POX-dependent functions is one of the challenges in cancer cell biology. Proline synthesis enzymes, P5C synthetase and P5C reductase (Ding et al. 2020) as well as collagen prolyl hydroxylases (D’Aniello et al. 2019) are up-regulated in most of cancer types and control rates of collagen biosynthesis. It has been suggested that proline availability and prolyl hydroxylases control progression of cancer cells (Loayza-Puch et al. 2016). Prolyl hydroxylases are Fe+2, ascorbate and α-KG-dependent enzymes. They belong to
the family of dioxygenases that includes collagen prolyl hydroxylase, DNA and histone demethylases and HIF-1α prolyl hydroxylase (D’Aniello et al. 2019). Since collagen prolyl hydroxylase compete with DNA and histone demethylases for ascorbate and α-KG, collagen biosynthesis may influence metabolic epigenetics. Another member of prolyl hydroxylases (α-KG and succinate dependent) was recently implicated in regulation of HIF-1α transcriptional activity by its targeting for proteasomal degradation (Xiong et al. 2018). Interestingly, proline inhibits HIF-1α prolyl hydroxylase and HIF-1α degradation (Surazynski et al. 2008a). The processes may also affect proline availability for PRODH/POX-dependent functions. The knowledge on functional significance of proline and collagen metabolism in complex metabolic regulatory mechanisms may contribute to understanding differential PRODH/POX-dependent functions (apoptosis/autophagy/survival) in cancer cells (Fig. 1).

**Collagen biosynthesis as a regulator of intracellular proline concentration**

Collagen is the most abundant protein in mammals. So far, 29 genetically distinct types of collagen have been described. They differ in terms of their spatial structure, the degree of post-translational modification and the function they perform in various tissues and organs (Ricard-Blum 2011). The most abundant are collagen type I–IV. Their common characteristic feature is the ability to create a trihelical structure composed of identical or different alpha subunits (Brodsky and Ramshaw 1997). The alpha subunits include repeating amino acid triplets containing glycine and often proline and hydroxyproline. The presence of hydroxyproline, an amino acid rarely found in other proteins, is particularly characteristic. It is formed by hydroxylation of prolyl residues in the newly synthesized collagen by prolyl hydroxylase (Myllyharju 2003). It is one of the critical processes in collagen biosynthesis. Any disturbances in this process lead to the inhibition of further stages of collagen biosynthesis contributing to lysosomal degradation of the defective polypeptide chains. It is estimated that about 10–20% of the newly synthesized collagen is degraded intracellularly (Bienkowski 1989). The presence of imino acids, proline and hydroxyproline, constituting about 25% of all collagen amino acids, allows the formation of a triple helix. The trihelical structure allows for the secretion of the molecule outside the cell and makes it resistant to the action of non-specific proteases. However, its function is not limited to tissue support and architecture. It also acts as a ligand for integrin receptors, inducing signalling pathways that regulate cell growth, differentiation and metabolism (Heino 2007). However, the interaction between collagen and integrin receptors is dynamic process. Extracellular collagen is degraded by specific metalloproteinases that cut the molecule into two parts that are further degraded by non-specific proteases (Krane 2008). The resulting short fragments are internalized and degraded intracellularly in the lysosomes to free amino acids, with the exception of imido-dipeptides, e.g., glycyl-proline. Instead, they are degraded by the cytoplasmic prolidase (Surazynski et al. 2008b). It is estimated that in this way the cell recovers about 90% of proline, which can be used for the biosynthesis of new proteins, including collagen (Jackson et al. 1975). However, during last few decades, it has been recognized that proline could play several other functions, e.g., as a stress molecule, regulator

![Fig. 1](image_url) Complex regulatory mechanism for PRODH/POX-dependent apoptosis/survival linking glycolysis, TCA, urea cycles, proline synthesis and degradation with collagen biosynthesis and degradation. Hypothesis is provided that up-regulation of prolidase (PEPD) and PRODH/POX with down-regulation of collagen biosynthesis may represent potential pharmacotherapeutic approach to induce apoptosis or autophagic death in cancer cells. ATP adenosine triphosphate, PEPD prolidase, PRO proline, PRODH/POX proline dehydrogenase/oxidase, PYCR 1-pyrroline-5-carboxylate reductase, P5C 1-pyrroline-5-carboxylate, ROS reactive oxygen species, TCA tricarboxylic acid cycle.
of transcription factors and substrate for proline dehydrogenase/proline oxidase (PRODH/POX)-dependent apoptosis/autophagy. Particular attention is focused on the role of proline availability in driving PRODH/POX-dependent functions. In this context, the activity of prolidase (proline supporting enzyme) and collagen biosynthesis (proline utilizing process) could be of critical importance in providing proline for PRODH/POX-induced apoptosis, autophagy or survival.

**PRODH/POX-dependent functions**

Proline dehydrogenase (PRODH, GenBankTM NM_016335) known also as proline oxidase (POX) is flavin-dependent enzyme associated with the inner mitochondrial membrane (Liu and Phang 2012b; Pandhare et al. 2012). The enzyme catalyses conversion of proline into Δ1-pyrroline-5-carboxylate (P5C). During this process, electrons are transported to electron transport chain producing ATP or they directly reduce oxygen, producing reactive oxygen species (ROS). Therefore, PRODH/POX is considered as a tumor suppressor or pro-survival factor, depending on the environmental conditions (Liu et al. 2012a, 2010; Liu and Phang 2012b; Phang and Liu 2012; Phang et al. 2012). Although the mechanism for switching survival/apoptosis mode is unknown, it seems that PRODH/POX activity/expression could be of importance in these processes. During stress situations (e.g., genotoxic, inflammatory, metabolic as oxygen or glucose deficiency), PRODH/POX is up-regulated in cancer cells and evoke pro-survival activity, both in vitro and in vivo (Mock et al. 1990; Myara et al. 1984; Phang 2019; Zareba et al. 2017). However, up-regulation of PRODH/POX expression in cancer cells is counteracted by inhibitory effect of succinate (TCA metabolite) (Burke et al. 2020; Hancock et al. 2016) and lactate (metabolite of Warburg effect) (Chen et al. 2018) on this enzyme. Importantly, a critical factor that mediate adaptation for such metabolic change is AMP-activated protein kinase AMPK (Laderoute et al. 2006; Wang and Guan 2009). This kinase inhibits energy-consuming processes and activates energy-producing processes to restore energy homeostasis during stress situation for the purpose of survival (Hardie 2008; Kato et al. 2002). Therefore, under glucose deficiency, POX generate ATP. Interestingly, under hypoxia PRODH/POX induces ROS production. However, in such conditions, ROS are not related to apoptosis, but rather to protective autophagy (Liu and Phang 2012b; Phang and Liu 2012). Since mammalian target of rapamycin (mTOR) signalling depends on the availability of nutrients (Reiling and Sabatini 2006), it seems that this pathway is important in POX regulation. In fact, inhibition of mTOR by rapamycin was found to upregulate PRODH/POX (Liu and Phang 2012b; Pandhare et al. 2009). However, in conditions of nutrient availability, silencing of PRODH/POX also induced pro-survival phenotype of cancer cells (Zareba et al. 2017). Therefore, it has been suggested that proline availability for PRODH/POX could differentiate the enzyme-dependent functions.

**The role of proline in cell metabolism and collagen biosynthesis**

Studies of last decade provided several lines of evidence for regulatory role of proline availability for PRODH/POX-dependent apoptosis/autophagy in cancer cells (Phang 2019). Large quantity of proline comes from protein degradation. Deregulation of energetic metabolism in cancer cells due to Warburg’s effect facilitates protein degradation as an alternative source of energy. Several studies showed that proline concentration is increased in cancer cells (Catchpole et al. 2011; Hirayama et al. 2009). Both hypoxia (Kakkad et al. 2010) and glucose depletion (Pandhare et al. 2009) were found to induce activity of metalloproteinases, MMP-2 and -9, suggesting the mechanism for the increase in cellular proline concentration. When glucose supply is limited, cancer cells may select proline as an alternative energy source, since proline has an advantage over fatty acids and glutamine, which like glucose requires delivery by the circulation. Therefore, proline may represent energy sense molecule and energy substrate. PRODH/POX expression is often down-regulated in various tumors, limiting mitochondrial proline degradation and PRODH/POX-dependent apoptosis and autophagy. Some studies suggested that critical factor for the process is proline availability that depends on the activity of prolidase (enzyme supporting cytoplasmatic proline level) and the rate of proline utilization in process of collagen biosynthesis (Zareba et al. 2020, 2017; Zareba and Palka 2016).

Most of free proline is released from collagen degradation products by cytoplasmatic prolidase [E.C.3.4.13.9], known also as Peptidase D or Inimopeptidase (Myara et al. 1984). It cleaves imido-dipeptides with C-terminal proline (Mock et al. 1990). The role of prolidase in recycling proline for collagen biosynthesis is well documented (Laderoute et al. 2006; Miltyk et al. 2005; Surazynski et al. 2008b). The enzyme activity as well as collagen biosynthesis were found to be co-ordinately up-regulated by β1-integrin receptor (Ivaska et al. 1999; Palka and Phang 1997) and insulin-like growth factor (IGF-I) signalling (Miltyk et al. 1998). Increase in the enzyme activity is due to its phosphorylation on serine/threonine residues (Surazynski et al. 2005).

Prolidase plays also important role in regulation of transcription factors. Overexpression of prolidase in colorectal cancer cells contributed to increase in nuclear hypoxia-inducible factor (HIF-1α) level and HIF-1α-dependent gene products, e.g., vascular endothelial growth factor (VEGF) and glucose transporter-1 (Glut-1) (Surazynski et al. 2008a). The activity of HIF-1α is regulated primarily at the level...
of its degradation (Jaakkola et al. 2001). The hydroxylation of specific proline residues in the oxygen-dependent domain (ODD) by prolyl hydroxylase (PHD) targets HIF-1α for ubiquitination and proteasomal degradation. It has been found that proline inhibits the degradation of HIF-1α, up-regulating its transcriptional activity. Another transcription factor affected by prolidase is NF-κB. Inhibition of prolidase activity by Cbz-pro contributed to up-regulation of NF-κB expression in fibroblasts (Wang and Guan 2009). On the other hand, transfection of colorectal cancer cells with prolidase vector was found to inhibit NF-κB expression (Wang and Guan 2009). The link between collagen biosynthesis and NF-κB is that the transcription factor is a potent inhibitor of gene expression for α1 and α2 subunits of type I collagen (Kouba et al. 1999; Miltyk et al. 2007; Rippe et al. 1999).

PRODH/POX down-regulates HIF-1α transcriptional activity (Liu et al. 2009a). When proline (inhibitor of HIF-1α degradation) is utilized by PRODH/POX, then HIF-1α undergoes proteasomal degradation limiting its pro-survival functions. PRODH/POX, the only proline degrading enzyme, initiates proline conversion to glutamate and α-ketoglutaric acid that inhibits transcriptional activity of HIF-1, since it is co-substrate of PHD, an enzyme, which hydrolyses specific prolyl residues of HIF-1α. Increase in PHD activity contributes to increase in HIF-1α degradation and a decrease in HIF-1 dependent gene expressions (Liu et al. 2009a; Verma 2006).

The above data suggest that proline plays important role in cellular metabolism. Therefore, collagen biosynthesis, the main utilization of proline, may represent regulatory mechanism for proline-dependent functions. In fact, collagen biosynthesis is tightly regulated by growth factors, hormones and integrin receptor ligands at both transcriptional and post-transcriptional levels (Sienkiewicz et al. 2004).

Several lines of evidence suggest that inhibition of collagen biosynthesis increases intracellular proline concentrations, making it available as a substrate for PRODH/POX-dependent functions. An example is inhibition of collagen biosynthesis by betulin derivative contributing to PRODH/POX-dependent apoptosis in endometrial adenocarcinoma cells (Szoka et al. 2017). Similar effect on collagen biosynthesis and apoptosis was found in leiomyoma cells treated with 2-methoxyestradiol (Salama et al. 2006) or in adenocarcinoma endometrial cells treated with PPARγ ligands (Fan et al. 2018; Surazynski et al. 2009). The mechanism for PPARγ-dependent inhibition of collagen biosynthesis was found at the level of NF-κB stimulation (Karna et al. 2006). As mentioned above, NF-κB is potent inhibitor of collagen gene expression. However, the PPARγ-induced functions are dependent on the status of estrogen receptor (Bonofiglio et al. 2005; Fan et al. 2018). It has been suggested that ERβ evokes pro-apoptotic effects, while ERα anti-apoptotic (Lu and Katzenellenbogen 2017). Probably, the mechanism of the process is a result of interaction between ERα and PPARγ (Bonofiglio et al. 2005; Kociecka et al. 2010). In view of estrogen-dependent modulation of collagen biosynthesis (Karna et al. 2006; Surazynski et al. 2009), it cannot be excluded that ERα/PPARγ cross-talk regulate proline availability for PRODH/POX-dependent functions.

High cytoplasmic proline concentration could be also considered as a pro-survival and pro-inflammatory since it contributes to up-regulation of transcriptional activity of HIF-1α that induces expression of COX-2, VEGF, TNF-α, TGF-β, IL-1, NF-κB (Surazynski et al. 2008a). Some of the cytokines stimulate collagen biosynthesis (e.g., COX-2, TNF-α, TGF-β) others inhibit the process (e.g., IL-1, NF-κB). Up-regulation of collagen biosynthesis removes proline from cytoplasm, limiting its availability for PRODH/POX-dependent functions. However, it may also contribute to tissue fibrosis, usually accompanying prolonged inflammation in cancer tissues. In view of cancer therapy, severe tissue fibrosis can impair drug delivery to the tissue. This suggests that enhanced collagen biosynthesis is not beneficial both from the point of view of molecular mechanism (POX-dependent functions), tissue function (fibrosis) and pharmacotherapy (fibrosis and inflammation). Although the complex regulatory mechanisms driving PRODH/POX-dependent functions are not well understood, in general, the above data suggest that inhibition of collagen biosynthesis in cancer cells contributes to increase in PRODH/POX-induced apoptosis and the process (collagen biosynthesis) could be considered a target for cancer treatment.

**Cell metabolic processes regulate proline availability for PRODH/POX-dependent function**

PRODH/POX cooperate with P5C reductase (P5CR) participating in proline turnover between mitochondria and cytoplasm. The conversion of proline to P5C that is shuttled between mitochondria and cytosol is coupled to glucose metabolism by pentose phosphate pathway (Pandhare et al. 2009; Phang et al. 2012, 2008). It is vital in maintenance of redox balance in a cell due to participation of NADPH/NADH in conversion of P5C to proline. Moreover, P5C is converted by P5C dehydrogenase (P5CDH) to glutamate, which is a precursor of α-ketoglutaric acid—a component of TCA cycle. PRODH/POX and ornithine aminotransferase (OAT) could also transform proline into ornithine that enters urea cycle (Liu and Phang 2012b). In view of the inhibitory role of PRODH/POX in tumor progression (Donald et al. 2001; Liu et al. 2005; Maxwell and Rivera 2003), all those metabolic cycles are of great importance in the PRODH/POX-dependent functions in neoplastic cells.

The role of interconversion of proline, P5C, ornithine and glutamate in PRODH/POX-dependent functions was described earlier (Huynh et al. 2020; Karna et al. 2020).
However, some cells do not express PRODH/POX, e.g., fibroblasts (Downing et al. 1977). The main function of the cells is biosynthesis of extracellular matrix constituents, including collagen. Therefore, PRODH/POX-dependent degradation of proline is undesirable in these cells. Proline derived from protein degradation is involved directly in process of collagen biosynthesis. It has been documented in several cell culture conditions (Karna et al. 2020). However, the process of collagen synthesis requires glutamine metabolism, particularly the synthesis of proline from glutamate and PSC (Hamanaka et al. 2019). During conversion of glutamate into proline, two molecules of NADPH are oxidized, one in mitochondria and one in cytosol (Li and Wu 2018). Probably, it is required for maintenance of redox potential, alternatively proline bearing reducing potential is utilized in collagen synthesis. In this way, collagen biosynthesis and glutamine metabolism is coupled to maintain redox balance. It has been proved in recent studies showing that proline biosynthesis supporting collagen biosynthesis is a vent for TGF-β-induced mitochondrial redox stress (Schworer et al. 2020).

In cancer cells, collagen biosynthesis is not a priority process as in fibroblasts. Probably in cancer cells, collagen biosynthesis modulates proline availability for PRODH/POX. In these cells, proline for collagen biosynthesis comes preferentially from P5C (e.g., generated from glutamine or ornithine). P5C conversion into proline is catalysed by three isoforms of P5C reductase: mitochondrial PYCR1/2 responsible for proline synthesis from glutamate and PYCRL, responsible for proline synthesis from ornithine (De Ingenis et al. 2012). It seems that intracellular proline derived from collagen degradation at first has to be processed by PRODH/POX yielding P5C in mitochondria (that is mixed with P5C from other sources) which in cytoplasm is converted back to proline by PYRCL. Proline produced from P5C derived from different sources could be a substrate for collagen biosynthesis or, when the process of collagen biosynthesis is inhibited, it could be again degraded by PRODH/POX. The shuttling proline-P5C between mitochondria and cytoplasm, called “proline cycle” (Phang et al. 1981) may work as an engine for regulation of oxidative potential. The intensity of proline cycle depends probably on complex regulatory mechanism for cell metabolism among which, the rate of collagen biosynthesis that utilizes substrate for PRODH/POX seems to play important role. The conversion of P5C to proline requires the presence of NADPH that is oxidized to NADP+. The generation of NADP+ is favoured when collagen biosynthesis is increased. The process is coupled with pentose phosphate pathway that utilizes NADP+ into NADPH in reaction catalysed by glucose 6-phosphate dehydrogenase, contributing to synthesis of purine nucleotides for DNA biosynthesis. In this context, collagen biosynthesis could reflect DNA biosynthesis. However, when collagen biosynthesis is up-regulated, it decreases proline availability for PRODH/POX, limiting proline cycle. In such a case, P5C is produced from glutamate and ornithine. However, production of proline from ornithine mediates a transfer of reducing potential from cytosolic NADPH into mitochondrial NAD+ limiting NADPH pool for P5C reductase and NADP+ for glucose 6-phosphate dehydrogenase. Therefore, collagen biosynthesis may play critical role not only in regulation of proline availability for PRODH/POX-dependent functions but also in metabolism of glutamine/glutamate and urea cycle (Huynh et al. 2020).

Based on the above divagations, the question arises what is the role of prolidase (as an enzyme supporting free proline) in regulation of PRODH/POX-dependent functions. It seems that in the mechanism of PRODH/POX-induced apoptosis or survival, proline availability and PRODH/POX expression are equally important. Previously, we have found correlation between PRODH/POX and prolidase in MCF-7 cells (Zareba et al. 2020, 2017). PRODH/POX silencing contributed to drastic increase in prolidase activity, while overexpression of prolidase attenuated PRODH/POX expression contributing in both conditions to pro-survival phenotype of the cells. PRODH/POX silencing was also correlated to collagen biosynthesis inhibition. Although the mechanism of this process requires further studies, it seems that collagen biosynthesis inhibition may result from down-regulation of prolyl hydroxylase, an important enzyme in collagen biosynthesis. It has been documented that free proline inhibits prolyl hydroxylase (Surazynski et al. 2008a) suggesting a mechanism for proline-dependent attenuation of collagen biosynthesis. On the other hand, increase in prolidase activity creates conditions for proline availability for PRODH/POX-dependent functions.

In view of the data, it seems that the extent of PRODH/POX expression may represent another player in complex regulatory processes mediated by proline metabolism. Whether up-regulation of PRODH/POX at proline availability in cancer cells could contribute to apoptosis requires to be explored. Therefore, the mechanisms for regulation of PRODH/POX are of great importance.

**Regulation of PRODH/POX expression**

The most potent inducer of PRODH/POX activity is tumor suppressor p53 (Phang et al. 2008; Polyak et al. 1997). Transcriptional regulation of PRODH/POX by p53 was found in the PRODH/POX promoter, containing a p53-response element (Maxwell and Kochevar 2008). PRODH/POX expression is also activated by peroxisome proliferator-activated gamma receptor (PPARγ), ligand-dependent transcription factor (Willson et al. 2000). Also, in this case PPARγ response element (PPRE) was found in the PRODH/POX promoter (Pandhare et al. 2006).
Thiazolidinedione (TZD, as rosiglitazone, troglitazone, and pioglitazone) are well-known agonist of PPARγ (Fumery et al. 2017). They have been also shown to evoke anticancer potential, especially for obesity-related cancers as prostate, breast, colon, thyroid, lung, and pituitary cancers (Kazberuk et al. 2020; Wang et al. 2017). These properties have been linked to the anti-inflammatory activity of PPARs. PPARγ was found to upregulate PRODH/POX expression (Pandhare et al. 2006), accompanied by ROS generation in several cancer cell lines (Kim et al. 2007; Liu and Phang 2012a; Pandhare et al. 2006; Wang et al. 2011). PPARγ-dependent up-regulation of PRODH/POX expression was linked to inhibition of tumour necrosis factor α (TNF-α) as well as interleukin 1β (IL-1β) (Han et al. 2017). Several studies have also suggested a relationship between PRODH/POX and cyclooxygenase II activity (COX-2), showing that high expression of PRODH/POX inhibited COX-2 expression (Liu et al. 2009b). The up-regulation of COX-2 has been found in many cancers and correlated with a worse prognosis in several types of malignant neoplasms (Coussens and Werb 2002). The anticancer potential of non-steroidal anti-inflammatory drugs (NSAIDs) is well-known phenomenon (Fujii et al. 2014; Harris et al. 2012; Toloczko-Iwaniuk et al. 2020). Moreover, NSAIDs are ligands of PPARγ (Kazberuk et al. 2020; Seetha et al. 2020). However, the mechanism of anticancer activity of NSAIDs is still unknown.

Knocking down PRODH/POX inhibited the apoptotic response to PPARγ ligands (Liu et al. 2012b; Phang et al. 2008). Therefore, it was suggested that TZD-dependent apoptosis undergoes through PRODH/POX-dependent ROS generation. Among the most potent down-regulators of PRODH/POX is miR-23b* (Liu et al. 2010) and oncogenic transcription factor c-MYC (Liu et al. 2012b). Overexpression of miR-23b* attenuated PRODH/POX activity, ROS generation, apoptosis and augmented HIF-1 signalling (Liu et al. 2010; Surazynski et al. 2008a) contributing to oncogenesis and tumor progression. Of great importance is finding that MYC inhibited PRODH/POX expression through up-regulation of miR-23b* attenuating production of ROS and apoptosis (Liu et al. 2012b). The factors involved in this process are outlined in Fig. 2

**Concluding remark**

The above data suggest that collagen metabolism (synthesis and degradation) through modulation of proline availability for PRODH/POX is involved in regulation of metabolism of glutamine, TCA and Urea cycles. Since collagen metabolism...
biosynthesis and more specifically, collagen prolyl hydroxylase competes with DNA and histone demethylases for ascorbate and α-KG, the inhibition of collagen biosynthesis could attenuate epigenetically cancer progression, while proline (saved in this process) could serve as an inhibitor of prolyl hydroxylase and substrate for PRODH/POX-dependent apoptosis. These findings allow to provide hypothesis that up-regulation of prolidase and PRODH/POX with down-regulation of collagen biosynthesis may represent potential pharmacotherapeutic approach to induce apoptosis or autophagic death in cancer cells. Such a possibility is currently under investigation.

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Conflict of interest The authors declare that they have no competing interests.

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References

Bienkowski RS (1989) Intracellular degradation of newly synthesized collagen. Reviv Biol Cellar 21:423–443

Bonofilio D et al (2005) Estrogen receptor alpha binds to peroxisome proliferator-activated receptor response element and negatively interferes with peroxisome proliferator-activated receptor gamma signaling in breast cancer cells. Clin Cancer Res 11:6139–6147. https://doi.org/10.1158/1078-0432.CCR-04-2453

Brodsky B, Ramshaw JA (1997) The collagen triple-helix structure. Matrix Biol 15:545–554. https://doi.org/10.1016/S0945-053x(97)90030-5

Burke L, Guterman I, Palacios Gallego R, Britton RG, Burschowsky D, Tufarelli C, Rufini A (2020) The Janus-like role of proline metabolism in cancer. Cell Death Discov 6:104. https://doi.org/10.1038/s41420-020-00341-8

Catchpole G et al (2011) Metabolic profiling reveals key metabolic features of renal cell carcinoma. J Cell Mol Med 15:109–118. https://doi.org/10.1111/j.1582-4934.2009.00939.x

Chen Y et al (2018) 3-Bromopyruvate sensitizes human breast cancer cells to TRAIL-induced apoptosis via the phosphorylated AMPK-mediated upregulation of DR5. Oncol Rep 40:2435–2444. https://doi.org/10.3892/or.2018.6644

Cousens LM, Werb Z (2002) Inflammation and cancer. Nature 420:860–867. https://doi.org/10.1038/nature01322

D’Aniello C et al (2019) Collagen prolyl hydroxylation-dependent metabolic perturbation governs epigenetic remodeling and mesenchymal transition in pluripotent and cancer cells. Cancer Res 79:3235–3250. https://doi.org/10.1158/0008-5472.CAN-18-2070

D’Aniello C, Patriarca EJ, Phang JM, Minchiotti G (2020) Proline metabolism in tumor growth and metastatic progression. Front Oncol 10:776. https://doi.org/10.3389/fonc.2020.00776

De Ingenis J et al (2012) Functional specialization in proline biosynthesis of melanoma. PLoS ONE 7:e45190. https://doi.org/10.1371/journal.pone.0045190

Ding Z et al (2020) Metabolic pathway analyses identify proline biosynthesis pathway as a promoter of liver tumorigenesis. J Hepatol 72:725–735. https://doi.org/10.1016/j.jhep.2019.10.026

Donald SP, Sun XY, Hu CA, Yu J, Mei JM, Valle D, Phang JM (2001) Proline oxidase, encoded by p53-induced gene-6, catalyzes the generation of proline-dependent reactive oxygen species. Cancer Res 61:1810–1815

Downing SJ, Phang JM, Kowaloff EM, Valle D, Smith RJ (1977) Proline oxidase in cultured mammalian cells. J Cell Physiol 91:369–376. https://doi.org/10.1002/jcp.1040910306

Fan P, Abderrahman B, Chai TS, Yerrum S, Jordan VC (2018) Targeting peroxisome proliferator-activated receptor gamma to increase estrogen-induced apoptosis in estrogen-deprived breast cancer cells. Mol Cancer Ther 17:2732–2745. https://doi.org/10.1158/1535-7163.MCT-18-0088

Fuji R et al (2014) Restoration of E-cadherin expression by selective Cox-2 inhibition and the clinical relevance of the epithelial-to-mesenchymal transition in head and neck squamous cell carcinoma. J Exp Clin Cancer Res 33:40. https://doi.org/10.1186/1756-9966-33-40

Fumery M et al (2017) Peroxisome proliferator-activated receptor gamma (PPARgamma) regulates lactase expression and activity in the gut. EMBO Mol Med 9:1471–1481. https://doi.org/10.15252/emmm.201707795

Hamanaka RB et al (2019) Glutamine metabolism is required for collagen protein synthesis in lung fibroblasts. Am J Respir Cell Mol Biol 61:597–606. https://doi.org/10.1165/rcmb.2019-0008OC

Han Q, Yuan Q, Meng X, Huo J, Bao Y, Xie G (2017) 6-Shogaol attenuates LPS-induced inflammation in BV2 microglia cells by activating PPAR-gamma. Oncotarget 8:42001–42006. https://doi.org/10.18632/oncotarget.16719

Hancock CN, Liu W, Alvord WG, Phang JM (2008) Co-regulation of mitochondrial respiration by proline dehydrogenase/oxidase and succinate. Amino Acids 48:859–872. https://doi.org/10.1007/s00726-015-2134-7

Hardie DG (2008) AMPK and Raptor: matching cell growth to energy supply. Mol Cell 30:263–265. https://doi.org/10.1016/j.molcel.2008.04.012

Harris RE, Beebe J, Alshafee GA (2012) Reduction in cancer risk by Cox-2 inhibition and the clinical relevance of the epithelial-to-mesenchymal transition. J Exp Oncol 10:776. https://doi.org/10.3389/ fonc.2020.00776

Han J, Schedule Z, Wang Q, Bao Y, Xie G (2017) 6-Shogaol attenuates LPS-induced inflammation in BV2 microglia cells by activating PPAR-gamma. Oncotarget 8:41928–41936. https://doi.org/10.18632/oncotarget.17566

Hieino J (2007) The collagen family members as cell adhesion proteins. Clin Exp Pharmacol Physiol 34:1001–1010. https://doi.org/10.1111/j.1522-1502.2007.00939.x

Hirayama A et al (2009) Quantitative metabolome profiling of colon and stomach cancer microenvironment by capillary electrophoresis time-of-flight mass spectrometry. Cancer Res 69:4918–4925. https://doi.org/10.1158/0008-5472.CAN-08-4806

Springer
Huynh TYL, Zareba I, Baszanowska W, Lewoniewska S, Palka J (2020) Understanding the role of key amino acids in regulation of proline dehydrogenase/proline oxidase (prodh/pox)-dependent autophagy/apoptosis as an approach to targeted cancer therapy. Mol Cell Biochem. https://doi.org/10.1007/s11010-020-03685-y
Ivaska J, Reunanen H, Westermarck J, Koivist o L, Kähräi VM, Heino J (1999) Integrin alpha2beta1 mediates isofor m-specific activation of p38 and upregulation of collagen gene transcription by a mechanism involving the alpha2 cytoplasmic tail. J Cell Biol 147:401–416
Jaakkola P et al (2001) Targeting of HIF-alpha to the von Hippel-Lindau ubiquitilyation complex by O2-regulated prolyl hydroxylatos. Science 292:468–472. https://doi.org/10.1126/science.1059796
Jackson SH, Dennis AW, Greenberg M (1975) I minodipeptiduria: a mechanism involving the alpha2 cytoplasmic tail. J Cell Biol 147:401–416
Karna E, Miltyk W, Pałka JA, Jarzabek K, Wołczyński S (2006) Hyaluronic acid counteracts interleukin-1-induced inhibition of collagen biosynthesis in cultured human chondrocytes. Pharmacol Res 54:275–281. https://doi.org/10.1016/j.phrs.2006.06.002
Karna E, Szoka L, Huynh TYL, Palka JA (2020) Proline-dependent regulation of collagen metabolism. Cell Mol Life Sci 77:1911–1918. https://doi.org/10.1007/s00018-019-03363-3
Kato K, Ogura T, Kishimoto A, Minegishi Y, Nakajima N, Miyazaki K (2004) Plasma prolidase activity: a possible index of collagen catabolism in chronic liver disease. Clin Chem 30:211–215
Kazberuk A, Zareba I, Surazynski A (2020) A novel plausible mechanism of NSAIDs-induced apoptosis in cancer cells: the implication of proline oxidase and peroxisome proliferator-activated receptor. Pharmacol Rep 72:1152–1160. https://doi.org/10.1007/s43440-020-00140-z
Kim KY, Ahn JH, Cheon HG (2007) Apoptotic action of peroxisome proliferator-activated receptor-gamma activation in human non small-cell lung cancer is mediated via proline oxidase-induced reactive oxygen species formation. Mol Pharmacol 72:674–685. https://doi.org/10.1124/mol.107.035584
Kociecka B, Surazynski A, Miltyk W, Palka J (2010) The effect of Tel-misaran on collagen biosynthesis depends on the status of estrogen activation in breast cancer cells. Eur J Pharmacol 628:51–56. https://doi.org/10.1016/j.ejphar.2009.11.043
Koubja DJ et al (1999) Nuclear factor-kappa B mediates TNF-alpha inhibitory effect on alpha 2(I) collagen (COL1A2) gene transcription in human dermal fibroblasts. J Immunol 162:4226–4234
Krane SM (2008) The importance of proline residues in the structure, stability and susceptibility to proteolytic degradation of collagens. Amino Acids 35:703–710. https://doi.org/10.1007/s00726-008-0073-2
Laderoute KR et al (2006) 5’-AMP-activated protein kinase (AMPK) is induced by low-oxygen and glucose deprivation conditions found in solid-tumor microenvironments. Mol Cell Biol 26:5336–5347. https://doi.org/10.1128/MCB.00166-06
Li P, Wu G (2018) Roles of dietary glycine, proline, and hydroxyproline in collagen synthesis and animal growth. Amino Acids 50:29–38. https://doi.org/10.1007/s00726-017-2490-6
Liu W, Phang JM (2012a) Proline dehydrogenase (oxidase) in cancer. BioFactors 38:398–406. https://doi.org/10.1002/biof.1036
Liu W, Phang JM (2012b) Proline dehydrogenase (oxidase), a mitochondrial tumor suppressor, and autophagy under the hypoxia microenvironment. Autophagy 8:1407–1409. https://doi.org/10.4161/auto.21152
Liu Y et al (2005) MnSOD inhibits proline oxidase-induced apoptosis in colorectal cancer cells. Carcinogenesis 26:1335–1342. https://doi.org/10.1093/carcin/bgi083
Liu W et al (2009a) Berberine reduces fibronectin and collagen accumulation in rat glomerular mesangial cells cultured under high glucose condition. Mol Cell Biochem 325:99–105. https://doi.org/10.1007/s11010-008-0024-y
Liu Y, Borchert GL, Donald SP, Diwan BA, Anver M, Phang JM (2009b) Proline oxidase functions as a mitochondrial tumor suppressor in human cancers. Cancer Res 69:6414–6422. https://doi.org/10.1158/0008-5472.CAN-09-1223
Liu W et al (2010) miR-23b targets proline oxidase, a novel tumor suppressor protein in renal cancer. Oncogene 29:4914–4924. https://doi.org/10.1038/onc.2010.237
Liu W, Glunde K, Bhuwalia ZM, Raman V, Sharma A, Phang JM (2012a) Proline oxidase promotes tumor cell survival in hypoxic tumor microenvironments. Cancer Res 72:3677–3686. https://doi.org/10.1158/0008-5472.CAN-12-0080
Liu W, Le A, Hancock C, Lane AN, Dang CV, Fan TW, Phang JM (2012b) Reprogramming of proline and glutamine metabolism contributes to the proliferative and metabolic responses regulated by oncogenic transcription factor c-MYC. Proc Natl Acad Sci U S A 109:8983–8988. https://doi.org/10.1073/pnas.1203241109
Loayza-Puch F et al (2016) Tumour-specific proline vulnerability uncovered by differential ribosome codon reading. Nature 530:490–494. https://doi.org/10.1038/nature16982
Lu W, Katzenellenbogen BS (2017) Estrogen receptor-beta modulation of the ERalpha-p53 loop regulating gene expression in liver cancer cells. Autophagy 8:1407–1409. https://doi.org/10.1002/auto.21152
Maxwell SA, Rivera A (2003) Proline oxidase induces apoptosis in tumor cells, and its expression is frequently absent or reduced in renal carcinomas. J Biol Chem 278:9784–9789. https://doi.org/10.1074/jbc.M210012200
Maxwell SA, Kochevar GJ (2008) Identification of a p53-response element in the promoter of the proline oxidase gene. Biochem Biophys Res Commun 369:308–313. https://doi.org/10.1016/j.bbrc.2008.01.171
Miltyk W, Karna E, Wolczyński S, Palka J (1998) Insulin-like growth factor I-dependent regulation of proline oxidase activity in cultured human skin fibroblasts. Mol Cell Biochem 193:177–183.
Miltyk W, Surazynski A, Kasprzak KS, Fivash MJ, Buzard GS, Phang JM (2005) Inhibition of proline oxidase activity by nickel causes decreased growth of proline auxotrophic CHO cells. J Cell Biochem 94:1210–1217. https://doi.org/10.1002/jcb.20384
Miltyk W, Karna E, Phang JA (2007) Prolidase-independent mechanism of camptothecin-induced inhibition of collagen biosynthesis in cultured human skin fibroblasts. J Biol Chem 141:287–292. https://doi.org/10.1093/jcb/mvm022
Mock WL, Green PC, Boyer KD (1990) Specificity and pH dependence for acylproline cleavage by prolidase. J Biol Chem 265:19600–19605
Myara I, Myara A, Mangeot M, Fabre M, Charpentier C, Lemonnier A (1984) Plasma prolidase activity: a possible index of collagen catabolism in chronic liver disease. Clin Chem 30:211–215
Myllyharju J (2003) Prolyl 4-hydroxylases, the key enzymes of collagen biosynthesis. Matrix Biol 22:15–24. https://doi.org/10.1016/s0945-053x(03)00006-4
Palka JA, Phang JM (1997) Prolidase activity in fibroblasts is regulated by interaction of extracellular matrix with cell surface integrin receptors. J Cell Biochem 67:166–175
Pandhare J, Cooper SK, Phang JM (2006) Proline oxidase, a proapoptotic gene, is induced by troglitazone: evidence for both peroxisome proliferator-activated receptor gamma-dependent
Collagen metabolism as a regulator of proline dehydrogenase/proline oxidase-dependent... 1925

and independent mechanisms. J Biol Chem 281:2044–2052. https://doi.org/10.1074/jbc.M507867200

Pandhare J, Donald SP, Cooper SK, Phang JM (2009) Regulation and function of proline oxidase under nutrient stress. J Cell Biochem 107:759–768. https://doi.org/10.1002/jcb.22174

Phang JM (2019) Proline metabolism in cell regulation and cancer biology: recent advances and hypotheses. Antioxid Redox Signal 30:635–649. https://doi.org/10.1089/ars.2017.7350

Phang JM, Liu W (2012) Proline metabolism and cancer. Front Biosci (Landmark Ed) 17:1835–1845

Phang JM, Yeh GC, Hagedorn CH (1981) The intercellular proline cycle. Life Sci 28:53–58. https://doi.org/10.1016/0024-3205(81)90365-9

Phang JM, Pandhare J, Liu Y (2008) The metabolism of proline as microenvironmental stress substrate. J Nutr 138:2008S-2015S

Phang JM, Liu W, Hancock C, Christian KJ (2012) The proline regulatory axis and cancer. Front Oncol 2:60. https://doi.org/10.3389/fonc.2012.00060

Poljak K, Xia Y, Zweier JL, Kinzler KW, Vogelstein B (1997) A model for p53-induced apoptosis. Nature 389:300–305. https://doi.org/10.1038/389300a0

Reiling JH, Sabatini DM (2006) Stress and mTORTure signaling. Oncogene 25:6373–6383. https://doi.org/10.1038/sj.onc.1209889

Ricard-Blum S (2011) The collagen family Cold Spring. Harb Perspect Biol 3:a004978. https://doi.org/10.1101/cshperspect.a004978

Rippe RA, Schrum LW, Stefanovic B, Solis-Herruzo JA, Brenner DA (1999) NF-kappaB inhibits expression of the alpha1(I) collagen gene. DNA Cell Biol 18:751–761. https://doi.org/10.1089/104459990134890

Salama SA, Nasr AB, Dubey RK, Al-Hendy A (2006) Estrogen metabolite 2-methoxyestradiol induces apoptosis and inhibits cell proliferation and collagen production in rat and human leiomyoma cells: a potential medicinal treatment for uterine fibroids. J Soc Gynecol Investig 13:542–550. https://doi.org/10.1016/j.sgi.2006.09.003

Schworer S et al (2020) Proline biosynthesis is a vent for TGFbeta-independent mitochondrial redox stress. EMBO J 39:e103334. https://doi.org/10.15252/embj.2019103334

Seetha A, Devaraj H, Sudhandiran G (2020) Indomethacin and juglone inhibit inflammatory molecules to induce apoptosis in colon cancer cells. J Biochem Mol Toxicol 34:e22433. https://doi.org/10.1002/jbt.22433

Sienkiewicz P, Palka M, Palka JA (2004) Oxidative stress induces IGF-I receptor signaling disturbances in cultured human dermal fibroblasts. A possible mechanism for collagen biosynthesis inhibition. Cell Mol Biol Lett 9:643–650

Surazynski A, Liu Y, Miltyk W, Phang JM (2005) Nitric oxide regulates prolidase activity by serine/threonine phosphorylation. J Cell Biochem 96:1086–1094. https://doi.org/10.1002/jcb.20631

Surazynski A, Miltyk W, Palka J, Phang JM (2008b) Prolidase-dependent regulation of collagen biosynthesis. Amino Acids 35:731–738. https://doi.org/10.1007/s00726-008-0051-8

Surazynski A, Donald SP, Cooper SK, Whiteside MA, Salnikow K, Liu Y, Phang JM (2008a) Extracellular matrix and HIF-1 signaling: the role of prolidase. Int J Cancer 122:1435–1440. https://doi.org/10.1002/ijc.23263

Surazynski A, Jarzabek K, Miltyk W, Wolczynski S, Palka J (2009) Estrogen-dependent regulation of PPAR-gamma signaling on collagen biosynthesis in adenocarcinoma endometrial cells. Neoplasma 56:448–454

Szoka L, Karna E, Hlebowicz-Sarat K, Karaszewski J, Boryczka S, Palka JA (2017) Acetylenic derivative of betulin induces apoptosis in endometrial adenocarcinoma cell line. Biomed Pharmacother 95:429–436. https://doi.org/10.1016/j.biopha.2017.08.104

Toloczko-Iwanicki N et al (2020) Proline-dependent induction of apoptosis in oral squamous cell carcinoma (OSCC)-the effect of celecoxib. Cancers (Basel). https://doi.org/10.3390/cancers12010136

Verma A (2006) Oxygen-sensing in tumors. Curr Opin Clin Nutr Metab Care 9:366–378. https://doi.org/10.1097/01.mco.0000232895.28674.79

Wang W, Guan KL (2009) AMP-activated protein kinase and cancer. Acta Physiol (Oxf) 196:55–63. https://doi.org/10.1111/j.1748-1716.2009.01980.x

Wang J, Lv X, Shi J, Hu X, Du Y (2011) Troglitazone induced apoptosis via PPAR gamma activated POX-induced ROS formation in HT29 cells. Biomed Environ Sci 24:391–399. https://doi.org/10.3967/0895-3988.2011.04.010

Wang Q, Imam MU, Yida Z, Wang F (2017) Peroxisome proliferator-activated receptor gamma (PPARgamma) as a target for concurrent management of diabetes and obesity-related cancer. Curr Pharm Des 23:3677–3688. https://doi.org/10.2174/1381612823666170704125104

Willson TM, Brown PJ, Sternbach DD, Henke BR (2000) The PPARs: from orphan receptors to drug discovery. J Med Chem 43:527–550

Xiong G et al (2018) Collagen prolyl 4-hydroxylase 1 is essential for HIF-1alpha stabilization and TNBC chemoresistance. Nat Commun 9:4456. https://doi.org/10.1038/s41467-018-06893-9

Zareba I, Palka J (2016) Prolidase-proline dehydrogenase/proline oxidase-collagen biosynthesis axis as a potential interface of apoptosis/autophagy. BioFactors 42:341–348. https://doi.org/10.1111/biof.12837

Zareba I, Surazynski A, Chrusciel M, Miltyk W, Doroszko M, Rahman N, Palka J (2017) Functional consequences of intracellular proline levels manipulation affecting PRODH/POX-dependent pro-apoptotic pathways in a novel in vitro cell culture model. Cell Physiol Biochem 43:670–684. https://doi.org/10.1016/j.biopha.2017.08.104

Zareba I et al (2020) Overexpression of prolidase induces autophagic death in MCF-7 breast cancer. Cells Cell Physiol Biochem 54:875–887. https://doi.org/10.33594/00000275

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