Estrogen Receptor Alpha as a Key Target of Red Wine Polyphenols Action on the Endothelium

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Abstract

Background: A greater reduction in cardiovascular risk and vascular protection associated with diet rich in polyphenols are generally accepted; however, the molecular targets for polyphenols effects remain unknown. Meanwhile evidences in the literature have enlightened, not only structural similarities between estrogens and polyphenols known as phytoestrogens, but also in their vascular effects. We hypothesized that alpha isoform of estrogen receptor (ERα) could be involved in the transduction of the vascular benefits of polyphenols.

Methodology/Principal Findings: Here, we used ERα deficient mice to show that endothelium-dependent vasorelaxation induced either by red wine polyphenol extract, Provinols™, or delphinidin, an anthocyanin that possesses similar pharmacological profile, is mediated by ERα. Indeed, Provinols™, delphinidin and ERα agonists, 17β-estradiol and PPT, are able to induce endothelial vasodilatation in aorta from ERα Wild-Type but not from Knock-Out mice, by activation of nitric oxide (NO) pathway in endothelial cells. Besides, silencing the effects of ERα completely prevented the effects of Provinols™ and delphinidin to activate NO pathway (Src, ERK 1/2, eNOS, caveolin-1) leading to NO production. Furthermore, direct interaction between delphinidin and ERα activator site is demonstrated using both binding assay and docking. Most interestingly, the ability of short term oral administration of Provinols™ to decrease response to serotonin and to enhance endothelial benefits, via ERα activation, was completely blunted in ERα deficient mice.

Conclusions/Significance: This study provides evidence that red wine polyphenols, especially delphinidin, exert their endothelial benefits via ERα activation. It is a major breakthrough bringing new insights of the potential therapeutic of polyphenols against cardiovascular pathologies.

Introduction

Epidemiological studies have enlightened that women have lower cardiovascular risk than men, and this protection progressively disappears after menopause. These studies (protection in premenopausal women) suggest and experimental studies (prevention of atheroma development in animals) demonstrate a major atheroprotective action of 17β-estradiol (E2) [1, 2]. E2 actions are essentially mediated by two molecular targets: estrogen receptor alpha (ERα) and beta (ERβ), but the former appears to mediate most of the actions of E2 on the cardiovascular system [1,2]. Endothelium represents a well recognized target of E2, which elicits several beneficial actions as increased NO production [1–3] subsequent to activation of endothelial NO synthase (eNOS) via a G-protein [4], and ERK and phosphatidylinositol-3-kinase pathways.

Epidemiological studies reported a greater reduction in cardiovascular risk and greater vascular protection associated with diet rich in polyphenols, including those from red wine [5]. We have previously shown that Provinols™, a polyphenolic extract from red wine, and delphinidin, an anthocyanin pharmacologically active and found in the total extract, induce an increase of endothelial NO dysfunction [12,13].
Evidences in the literature have enlightened, not only structural similarities between estrogens and polyphenols known as phytoestrogens, but also in their vascular effects with regard to endothelial NO production. Indeed, it has been reported that the phytoestrogen genistein produces acute NO-dependent dilation of human forearm vasculature with similar potency to E2 [14]. Also, genistein induces a late but sustained activation of the eNOS system in vitro [15]. Moreover, chronic administration of genistein improves endothelial dysfunction in spontaneously hypertensive rats that involves eNOS, caveolin and calmodulin expression and NADPH oxidase activity [16].

Red wine polyphenols do not contain genistein and no direct evidence for the nature of the receptor triggering the effect of red wine polyphenols receptors in endothelial cells has been demonstrated. Nevertheless, the aim of this work was to investigate the hypothesis that ERα is one of the targets involved in the vasculoprotective effects of Provinols™ and delphinidin. For this, we first studied the endothelium-dependent relaxation to polyphenols in aortas from both ERα Knock-Out (KO) and Wild Type (WT) mice, and then we analyzed molecular pathways associated with NO production in endothelial cells stimulated by Provinols™ and delphinidin by silencing ERα activity or expression either with pharmacological inhibitor or siRNA, respectively. We also studied binding assay and molecular modelling of interaction between delphinidin and ERα. Finally, we tested the physiological relevance of our findings in vivo by testing the effect of short term oral treatment of ERα WT and KO mice with Provinols™ with respect to endothelial NO response.

Results

The role of ERα in the endothelium-dependent relaxation to Provinols™ and to delphinidin was evaluated by using vessels taken from ERα WT and KO mice. First, we tested the ability of ERα agonists such as E2, which acts on both ERα and ERβ isoforms, and 1,3,5-tris(4-hydroxyphenyl)-4-propyl-1H-pyrazole (PPT), which is specific for ERα, to activate the endothelium. This was demonstrated by the capacity of the two ERα agonists to induce relaxation in aortas from ERα WT but not from KO mice in the presence of functional endothelium only (Figure 1A and 1B).

The concentration of E2 to elicit maximal relaxation was in accordance with that reported by Li et al [17] in the same vessels. As previously described by our group, red wine polyphenols and delphinidin are able to induce endothelium-dependent relaxation in mice aortas. Interestingly, the vasorelaxant effect of these two polyphenols was found in aortas from ERα WT mice (Figure 1C and 1D), but was completely abolished when ERα is deleted. In ERα deficient mice, a slight contraction to Provinols™ and delphinidin were even detected (Figure 1C and 1D).

These data suggest the involvement of ERα in the endothelium-dependent relaxation in response to the two polyphenols used. Then, we assessed if the endothelium-dependent relaxation evoked by ERα stimulation is due to an increase in NO production. For this we stimulated the human endothelial cell line, EaHy 926, either with Provinols™ or delphinidin for 10 minutes in presence or in absence of Fulvestrant, an ERα pharmacological antagonist, or with a siRNA directed against ERα. Provinols™ and delphinidin were used at maximally active concentration to induce relaxation in the rat aortic rings and to increase cytosolic calcium in endothelial cells, as previously described [7,18]. Both Provinols™ and delphinidin were used at maximally active concentration to induce relaxation in the rat aortic rings and to increase cytosolic calcium in endothelial cells, as previously described [7,18]. Both Provinols™ and delphinidin were able to induce an increase in NO production. However, when ERα was silenced either by Fulvestrant or siRNA, the increase of NO production induced by Provinols™ and delphinidin was completely prevented (Figure 2A). As a positive control for ERα activation-induced

Figure 1. ERα mediates endothelium-dependent relaxation induced by agonists, Provinols™ and delphinidin. Concentration-effect curves to increasing concentrations of PPT (A), 17β estradiol (E2) (B), Provinols™ (C) or delphinidin (D) in aortic rings, with functional endothelium precontracted with U46619, taken from ERα Wild Type (open circles) and Knock-Out (filled circles) mice (n = 5–6). **P<0.01, ***P<0.001.

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NO production, we used PPT and E2. Both agonists were able to enhance NO production, and this effect was abolished when ERα was antagonized by Fulvestrant (Figure 2B). As EaHy 926 are derived from human umbilical vein endothelial cells, we extracted and cultured endothelial cells from aortas taken either from ovariectomized ERα WT or KO mice, or from ovariectomized Swiss mice, using the method described by Kobayashi et al. [19]. Cells were then stimulated with Provinols™, delphinidin and PPT, in absence or in presence of Fulvestrant, using the same protocol as for EaHy 926. Interestingly, in aortic endothelial cells taken either from ERα WT mice or Swiss mice, Provinols™, delphinidin and PPT were all able to induce NO production (Figure 2C and 2E). The ability of the three compounds to stimulate NO release was completely blunted in the aortic endothelial cells extracted from ERα KO mice (Figure 2D) or in aortic endothelial cells taken from ERα WT mice and Swiss mice in the presence of Fulvestrant (Figure 2C and 2E).

Recently, an increase of NO production via the interaction of a molecular pathway involving the phosphorylation of Src, ERK1/2 and eNOS Ser1177 has been reported with regard to resveratrol [20]. NO pathway was then investigated in order to decipher the molecular mechanisms underlying ERα-associated NO increase and the subsequent vasodilatation induced either by Provinols™, delphinidin or by PPT and E2. We also analysed caveolin-1 expression, a protein that segregates the inactive form of eNOS on the cellular membrane and modulates eNOS activity. We demonstrated that both Provinols™ and delphinidin increased the phosphorylation of Src, ERK1/2 and eNOS Ser 1177, as well as of caveolin-1, in human endothelial cells. Moreover, when ERα was blocked or silenced, the activation of this pathway was completely blunted (Figure 3A to 3D). In addition, the same effects were observed after PPT or E2 treatment in the sense that the two ER agonists increased phosphorylation of Src, ERK1/2, eNOS and caveolin-1. Furthermore, when ERα was antagonized with Fulvestrant, neither PPT nor E2 induced phosphorylation of these enzymes (Figure 3E to 3H), suggesting the involvement of an ERα-dependent mechanism. Even if it has already been shown that both Provinols™ and delphinidin are able to induce NO production in endothelial cells [18], and that estrogens are also able to increase NO production via ERα [21], here we demonstrate for the first time the direct link between the ability of Provinols™ and delphinidin to stimulate NO pathways leading to endothelial NO production through ERα activation.

To verify the direct interaction of red wine polyphenols with ERα and to exclude the implication of other factors, binding assay between delphinidin and ERα was performed. Binding assay showed that delphinidin exerts 73% of specific inhibition against E2 on ERα (Figure 4D). Furthermore, we performed a docking study of delphinidin on ERα. The predicted binding mode of the ligand-binding domain on ERα is relatively similar to that observed in the X-ray structure of the ERα with E2 (Figure 4A and 4B). The three aromatic rings undergo significant apolar contacts with hydrophobic residues located at the centre of the binding site (Leu349, Ala350, Leu384, Leu387, Met388, Leu391,

**Figure 2. ERα activation induces endothelial NO release.** NO production induced by Provinols™, delphinidin, PPT and 17-β estradiol (E2) in endothelial cells and consequences of ERα pathway inhibition with either Fulvestrant or siRNA directed against ERα (siRNA). EaHy 926 cells (A and B) or aortic endothelial cells from either ERα Wild-Type (WT) mice (C), ERα Knock-Out (KO) mice (D) or from Swiss mice (E) were treated for 10 minutes with Provinols™ (10^{-2} g/L), delphinidin (10^{-2} g/L), PPT (10^{-5} M) or E2 (10^{-5} M) and this in the presence or in the absence of either Fulvestrant (30 nM) (hatched bars) or after knocking down the receptor with siRNA (siRNA) (striped bars). Then NO production was assessed by electronic paramagnetic resonance. In panel A, insert shows Western blot for ERα in the control (lane 1) or ERα siRNA (lane 2)-treated EaHy endothelial cells. *P<0.05, **P<0.01, ***P<0.001 vs non treated; #P<0.05, ###P<0.01 vs Provinols™, delphinidin (A, C and D), PPT or E2 alone (B and D), (n=4–6). doi:10.1371/journal.pone.0008554.g002
Phe404, Met421, Ile424, Leu428, Leu525) (Figure 4C). An aromatic edge-to-face interaction is also engaged with the phenyl ring of Phe404, as for E2. Last, a strong H-bond anchors delphinidin and E2 to a polar residue (Glu353) at one end of the binding site. A significant difference with the E2 binding mode is the loss of two H-bonds to Arg394 and His524, which are partly compensated by novel interactions to Leu346 (H-bond to the backbone oxygen atom) and His524 (aromatic interaction). These results indicate that delphinidin, like E2, interact directly with ERα.

Finally, we show that short term oral administration of ProvinolsTM decreased contraction to serotonin (5-HT) in the presence of functional endothelium (Figure 5A) and enhanced the sensitivity to acetylcholine endothelium-dependent relaxation in aorta from ERαWT mice (Figure 5C) in association with increased NO production (Figure 5E) and reduced superoxide anions in mesenteric arteries (Figure 5F). All of these effects of oral administration of ProvinolsTM were completely blunted in ERα deficient mice (Figure 5B, 5D–5F). These results suggest that ERα triggers the in vivo effects of ProvinolsTM and demonstrate for the first time the physiological relevance of this receptor in triggering the vascular protection induced by red wine polyphenols.

Red wine contains a wide variety of polyphenols, which derive mainly from grape solids (skin and seeds) and can be divided in two classes, flavonoids and non-flavonoids. Although, the non-flavonoid, resveratrol has been reported to trigger some of the beneficial effects of red wine polyphenols including the activation of sirtuin and NO pathway, we have focused our attention on the flavonoid components of the red wine polyphenols such as delphinidin. Indeed, our previous study looking at the possible active principles that support the endothelial NO-dependent relaxation produced by red wine polyphenols, including ProvinolsTM, demonstrate that anthocyanins and oligomeric-condensed tannins exhibit a pharmacological profile comparable to the original extract, the most potent being delphinidin [6]. Beside, delphinidin has been reported to induce endothelial NO release via an increase of cytosolic Ca2+ in endothelial cells [7] and protects against endothelial cell apoptosis acting through NO pathway [22].

The major aim of this work was to investigate the hypothesis that ERα is one of the targets involved in the vasculoprotective effects of ProvinolsTM and delphinidin. Firstly, we report the existence of an endothelium-dependent relaxation associated with ERα-stimulation, c-Src/ERK1/2-mediated activation of eNOS, with consequent endothelial NO release via a non-genomic mechanism at the same range of concentration than that reported

**Figure 3.** ERα activation induces phosphorylation of NO signaling proteins. Activation of NO pathway induced by ProvinolsTM, delphinidin, PPT and 17-β estradiol (E2) in endothelial cells and consequences of ERα pathway inhibition with either Fulvestrant or siRNA directed against ERα (siRNA). EaHy 926 cells were treated for 10 minutes with ProvinolsTM (10−7 g/L), delphinidin (10−5 g/L), PPT (10−5 M) or E2 (10−5 M) and this in the presence or in the absence of either Fulvestrant (30 nM) (hatched bars) or after knocking down the receptor with siRNA (siRNA) (striped bars). Then cells were lysed to perform Western Blot analysis, *P<0.05, **P<0.01, ***P<0.001 vs non treated; #P<0.05, ##P<0.01 vs ProvinolsTM, delphinidin (A, B, C and D), PPT or E2 alone (E, F, G and H), (n = 4–6).

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by Li et al. [17]. Secondly, the most important novel observation is that the endothelium-dependent relaxation to Provinols TM and delphinidin found in aortas from ER α WT is completely abolished when ER α is deleted. Interestingly, both compounds elicit contraction in endothelium-denuded arteries taken from ER α deficient mice suggesting that the molecular targets of the two compounds on the smooth muscle are different to ER α (data not shown). Moreover, the ability of the two compounds to enhance the rapid release of NO (10 min) from cultured endothelial cells associated with phosphorylation of Src, ERK1/2, eNOS and caveolin-1 is blunted after silencing ER α either by Fulvestrant or siRNA. Finally, the capacity of Provinols TM and delphinidin to increase NO in mouse aortic endothelial cells was not only abolished in the presence of Fulvestrant and most likely when these cells were taken from ER α deficient mice. Altogether, these data demonstrate the direct link between the ability of Provinols TM and delphinidin on ERα triggering the in vivo effects of Provinols TM and demonstrate for the first time the physiological relevance of this receptor. One can advance the hypothesis that ERα might by one of the molecular target(s) triggering the beneficial effects of dietary supplementation of Provinols TM on obesity-associated alterations with respect to metabolic disturbances and cardiovascular functions recently reported in Zucker fatty (ZF) rats [24]. Further studies should be conducted in order

![Figure 4. Binding assay and docking show direct interaction between delphinidin and ERα activator site.](image)

In the present study, direct interaction of delphinidin with ERα excluding the implication of other factors is demonstrated by the capacity of this compound to exert 73% of specific inhibition against E2 on ERα. Furthermore, we perform a docking study of delphinidin on ERα. The predicted binding mode of the ligand-binding domain on ERα is relatively similar to that observed in the X-ray structure of the ERα with E2.

These data strongly suggest that ERα triggers the in vivo effects of Provinols TM and demonstrate for the first time the physiological relevance of this receptor. One can advance the hypothesis that ERα might by one of the molecular target(s) triggering the beneficial effects of dietary supplementation of Provinols TM on obesity-associated alterations with respect to metabolic disturbances and cardiovascular functions recently reported in Zucker fatty (ZF) rats [24]. Further studies should be conducted in order
to evaluate the role of other pathways that may be involved in cardiovascular effects induced by red wine polyphenols including ERβ or cyclooxygenase pathways.

The findings that vascular protection induced by red wine polyphenols, and in particular by delphinidin, requires ERα activation is a major breakthrough in understanding the therapeutic potential of polyphenols in cardiovascular pathologies. These properties of red wine might explain the prevention of ischemic heart disease [24,25], stroke [10] and metabolic diseases [24], in different experimental models.

**Methods**

Provinols™ was obtained from Société Française des Distilleries (Vallon Pont d’Arc, France) and delphinidin was purchased from Extrasynthese (Genay, France). The university of Angers ethical committee approved the present protocol. All animal studies were carried out using approved institutional protocols and were conformed the Guide for the Care and Use of Laboratory Animals published by US National Institutes of Health (NIH Publication No. 85–23, revised 1996). Methods for vascular reactivity performed in mice [26,27] and endothelial cells extraction and culture were set up as previously described [19]. Methods for RNA interference and transient transfection to silence ERα were adapted from Agouni et al. [26]. NO and O₂⁻ production in mesenteric arteries from ERα WT or KO mice receiving either standard diet or diet containing Provinols™ (*P<0.05 vs control diet; #P<0.05 KO vs WT; n = 5).

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Figure 5. Involvement of ERα in vascular effects induced by an oral treatment with Provinols™. Concentration-effect curves to increasing concentrations of serotonin (5-HT) in aortas, with endothelium, from ERα Wild Type (WT) (A) or Knock-Out (KO) (B) mice treated either with control diet (open circles) or diet containing Provinols™ for 2 weeks at 20 mg/kg/day (filled circles, ***P<0.001, n = 6). Concentration-effect curves to increasing concentrations of acetylcholine (Ach) in aortas with endothelium, pre-contracted at 80% of the maximal contraction with U46619 from ERα WT (C) or KO (D) mice treated either with control diet (open circles) or diet containing Provinols™ for 2 weeks at 20 mg/kg/day (filled circles) (*P<0.05, n = 6). Quantification of NO (E) and O₂⁻ production (F) in mesenteric arteries from ERα WT or KO mice receiving either standard diet or diet containing Provinols™ (*P<0.05 vs control diet; #P<0.05 KO vs WT; n = 5).
Supporting Information

Methods S1
Found at: doi:10.1371/journal.pone.0008554.s001 (0.04 MB DOC)

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References


Author Contributions

Conceived and designed the experiments: MC AT RA. Performed the experiments: MC AT DR. Analyzed the data: MC AT DR RA. Contributed reagents/materials/analysis tools: MC AT MCM JFA. Wrote the paper: MC AT MCM JFA RA.