



Quantitative Analysis of Vasodilatory Action of Quercetin on Intramural Coronary Resistance Arteries of the Rat *In Vitro*

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Abstract

Background: Dietary quercetin improves cardiovascular health, relaxes some vascular smooth muscle and has been demonstrated to serve as a substrate for the cyclooxygenase enzyme.

Aims: 1. To test quantitatively a potential direct vasodilatory effect on intramural coronary resistance artery segments, in different concentrations. 2. To scale vasorelaxation at different intraluminal pressure loads on such vessels of different size. 3. To test the potential role of prostanoids in vasodilatation induced by quercetin.

Methods: Coronary arterioles (70–240 μm) were prepared from 24 rats and pressurized in PSS, using a pressure microangiometer.

Results: The spontaneous tone that developed at 50 mmHg was relaxed by quercetin in the 10^{-9} moles/lit concentration ($p < 0.05$), while 10^{-5} moles/lit caused full relaxation. Significant relaxation was observed at all pressure levels (10–100 mmHg) at 10^{-7} moles/lit concentration of quercetin. The cyclooxygenase blocker indomethacin (10^{-5} moles/lit) induced no relaxation but contraction when physiological concentrations of quercetin were present in the tissue bath ($p < 0.02$ with Anova), this contraction being more prominent in smaller vessels and in the higher pressure range ($p < 0.05$, Pearson correlation). A further 2–8% contraction could be elicited by the NO blocker L-NAME (10^{-4} moles/lit).

Conclusion: These results demonstrate that circulating levels of quercetin (10^{-7} moles/lit) exhibit a substantial coronary vasodilatory effect. The extent of it is commensurable with that of several other physiological mechanisms of coronary blood flow control. At least part of this relaxation is the result of an altered balance toward the production of endogenous vasodilatory prostanoids in the coronary arteriole wall.

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Data Availability: The authors confirm that all data underlying the findings are fully available without restriction. All relevant data are within the paper and its Supporting Information files.

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Introduction

Dietary polyphenols, with quercetin being a member of this group, are present in substantial amount in various vegetables, fruits, vine and tea, and have beneficial effects on the vessel wall [1–3]. They delay atherosclerotic processes [4] and reduce hypertension [1,5–7]. Quercetin and rutin components of food are easily absorbed, and rutin is transformed into quercetin [8]. Its glucuronide and sulfate conjugates appear in blood plasma [9–11]. These metabolites are cleaved in the peripheral tissues in situ, vessel wall included [12]. A vasorelaxation effect was described both in vivo and in vitro experiments [13–15]. While tissue blood flow is controlled dominantly by resistance arteries, quercetin effects have been studied mainly on larger vessels. Both endothelium dependent [16] and independent [17] mechanisms

of quercetin-induced vasorelaxation have been described. They can be different in different vascular fields. Quercetin was found to be a reducing co-substrate of the second step in the cyclooxygenase (COX) enzymatic reaction [18], thus affecting endogenous prostanoid production [19].

Intramural coronary resistance arteries have a marked spontaneous myogenic tone which is substantially modified by endothelial relaxation and endogenous prostanoid production. Both vasoconstrictor and vasodilator prostanoids are produced by intramural coronary resistance arteries, vasoconstrictors prevailing at higher intraluminal pressures [20].

One of the aims of the present study was to test quantitatively if coronary resistance arteries do show a direct vasodilation in response to physiological concentrations of quercetin. In addition, we investigated the quercetin effect on coronary arterioles of

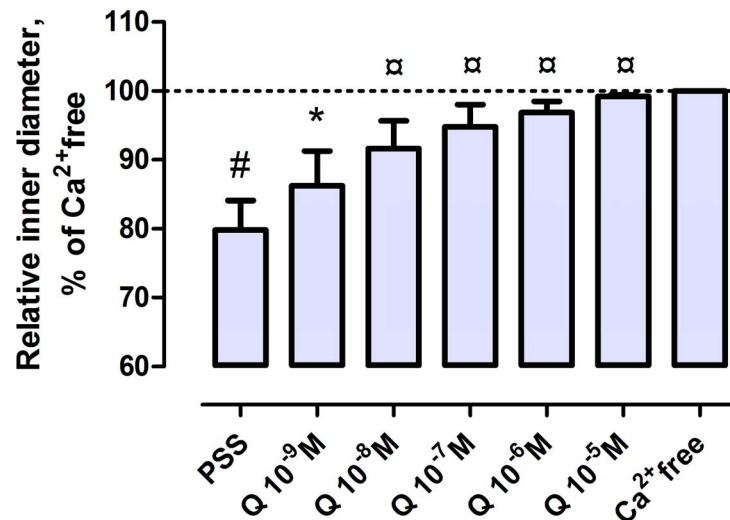


Figure 1. Concentration-response curve of quercetin on coronary resistance arterioles of rat. Intramural coronary arteriole segments were spontaneously contracted, then dilated by quercetin at intraluminal pressure 50 mmHg. (PSS, Krebs-Ringer solution, Ca²⁺-free, Ca²⁺-free solution). #, significantly different from fully relaxed; α, different from spontaneously contracted with ANOVA; * different from spontaneously contracted with the paired t-test.
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different sizes at intact, cylindrical vascular geometry, as according to earlier observations, pharmacological modulation of smaller and larger resistance arteries may differ substantially [21]. After we have found a significant dilation effect, it was assumed that this action could be initiated at least partly through altered production of endogenous vasoactive prostanoids. This hypothesis was tested by indomethacin inhibition of COX activity in vessel segments of different caliber and at different intraluminal pressures.

Materials and Methods

Pressure microangiometry

Male Wistar rats, weighing 350–450 gr were anesthetized by pentobarbital (Nembutal, CEVA, 45 mg/kg body weight i.p.). All procedures conformed the Guide for the Care and Use of

Laboratory Animals (NIH, 1996), the legal and institutional guidelines for animal care and were approved by the Animal Care Committee of the Semmelweis University.

The heart was removed and put in cold oxygenated physiological salt solution (PSS). A terminal branch of the left anterior descending coronary artery (intramural small artery in the rat) was prepared from the ventricular muscle tissue. Vessel segments with no or with limited number of side branches, having outer diameter of 150–250 μm in situ were selected. Preparation was made as described earlier [22]. Excised segments, with lengths over 2.0 mm were cannulated at both ends using microcannulas (outer diameter ~130 μm) and mounted in a glass bottomed tissue bath. Their length was extended to the original in situ value. Segments were pressurized with saline connecting them to servo-controlled pumps (Living Systems, Burlington, VT, US). The temperature of the

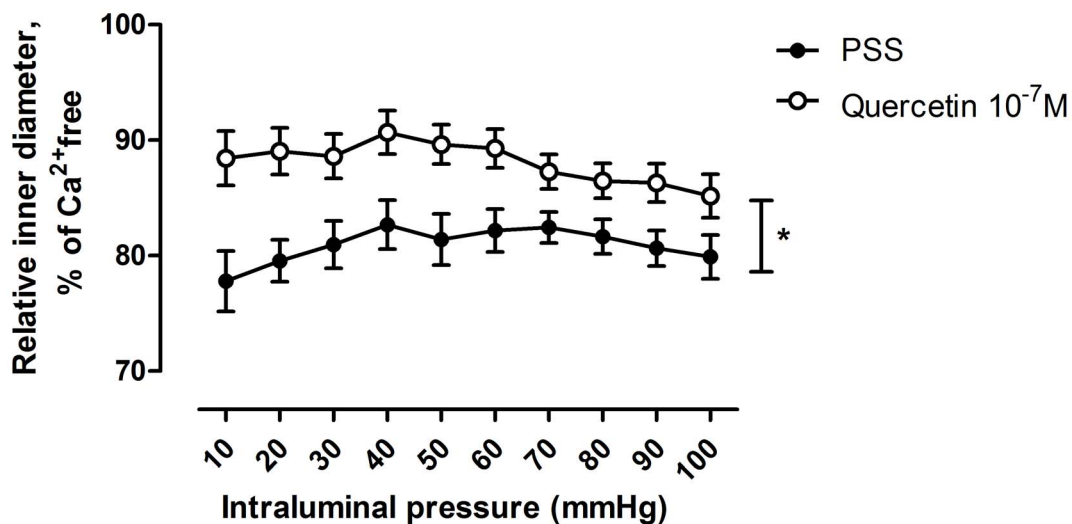


Figure 2. Quercetin dilation on spontaneously contracted small coronary arterioles at different pressures. Full circles represents diameters in PSS, empty circles are diameters in quercetin 10⁻⁷ moles/lit. * p<0.05 with ANOVA
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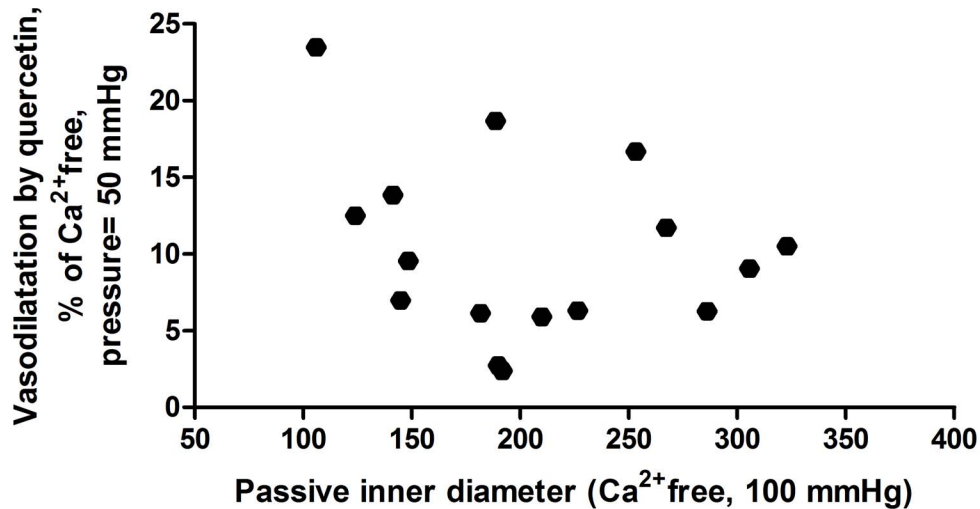


Figure 3. Quercetin induced dilatation as function of morphological lumen size. We found no correlation ($p > 0.05$ with Pearson's correlation).

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bath was thermostated at 37°C, and bubbled with a gas mixture of 5% CO₂, 20% O₂ and 75% of N₂, keeping pH at 7.4. A continuous superfusion velocity of 2.8 ml/min was ensured. The bath was positioned on the stage of an inverted microscope (Leica), the pictures of the segments were taken by a digital camera (Leica DFC 320) and an image analyzing software (Leica Qwin). Outer and inner diameters of segments were measured off-line on frozen pictures. Calibration was made using a micrometer etalon (Wild).

Materials

All drugs, including quercetin, were purchased from Sigma-Aldrich. The composition of the PSS (Krebs-Ringer solution) we used: NaCl 119, KCl 4.7, NaH₂PO₄ 1.2, MgSO₄ 1.17, NaHCO₃ 24, CaCl₂ 2.5, glucose 5.5, and EDTA 0.034 (in mmol/lit). The Ca²⁺-free PSS contained: NaCl 92, KCl 4.7, NaH₂PO₄ 1.18,

MgCl₂ 20, MgSO₄ 1.17, NaHCO₃ 24, glucose 5.5, EGTA 2, and EDTA 0.025 (in mmol/lit).

In vitro protocols

In the first series of experiments, concentration-response curves with quercetin were taken on coronary resistance artery segments from 8 rats. To do that, segments were pressurized at 50 mmHg. This pressure was continuously kept throughout the measurement. Such segments develop substantial spontaneous tone under similar circumstances. After 40 minutes of equilibration, superfusion was changed to PSS containing increasing concentrations of quercetin from 10⁻⁹ to 10⁻⁵ moles/lit. (Physiological concentrations in humans are reported to be 10⁻⁷ moles/lit [7].) 20 min equilibration was left for all steps, at the end of which stable diameters were registered. Reversibility was checked by repeated administration of

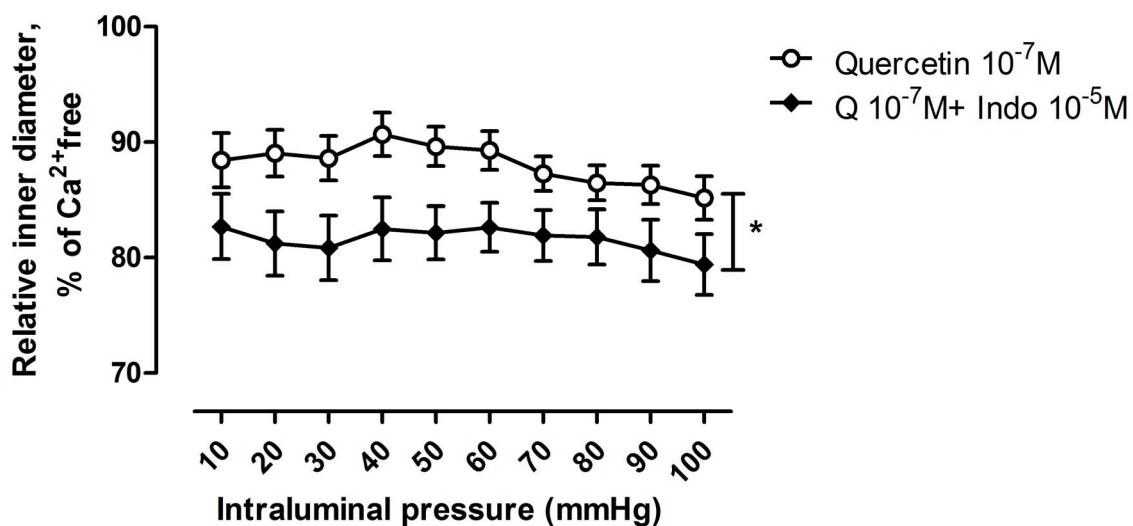


Figure 4. Effect of indomethacin on inner diameter of quercetin-treated coronary arteriole segments. Note contraction with indomethacin ($p < 0.05$ with ANOVA), in contrast the expected vasodilation, indicating the presence of vasodilatory prostanoids with the polyphenol. Indomethacin concentration was 10⁻⁵ moles/lit, represented by full circles, quercetin was in 10⁻⁷ moles/lit concentration, represented by empty circles.

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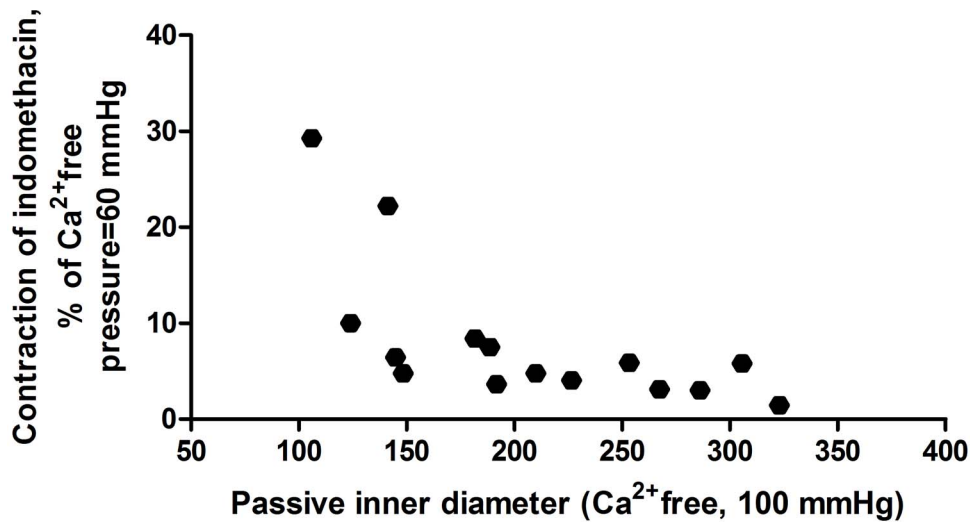


Figure 5. Correlation of indomethacin induced contraction on quercetin-dilated segments with the morphological lumen size. Note a more prominent contraction in smaller vessels ($p < 0.05$ with Pearson's correlation). doi:10.1371/journal.pone.0105587.g005

PSS. Finally, calcium-free PSS was superfused (20 minutes) and relaxed diameters were measured. Concentration-response curves were determined. Relaxation from spontaneous tone was expressed in percents of fully relaxed diameter.

In a second series, pressure-radius curves were taken using 16 similar vessel segments. Pressure was raised in 10 mmHg steps from 10 up to 100 mmHg. Then 10^{-7} mol/lit quercetin containing PSS solution was superfused, and the pressure-radius curves were repeatedly taken. Next, the cyclooxygenase blocker indomethacin was applied (10^{-5} moles/lit) with the quercetin still in the bath. NO effect was tested by additional application of the NO blocker L-NAME (10^{-4} moles/lit). Reproducibility was controlled by repeated measurements with 10^{-7} moles/lit

quercetin alone and then with PSS without any drug added (spontaneous tone). The experiment was terminated by superfusion with Ca^{2+} -free PSS (full relaxation). Contractions and relaxations were expressed in % of inner diameter measured at the same pressure in Ca^{2+} -free PSS.

Statistical procedures

Values are expressed as means \pm SEM. Statistical analysis was performed using paired t-test, or two-way repeated measures ANOVA, followed by Tukey post-hoc test. Values of $p < 0.05$ were considered statistically significant.

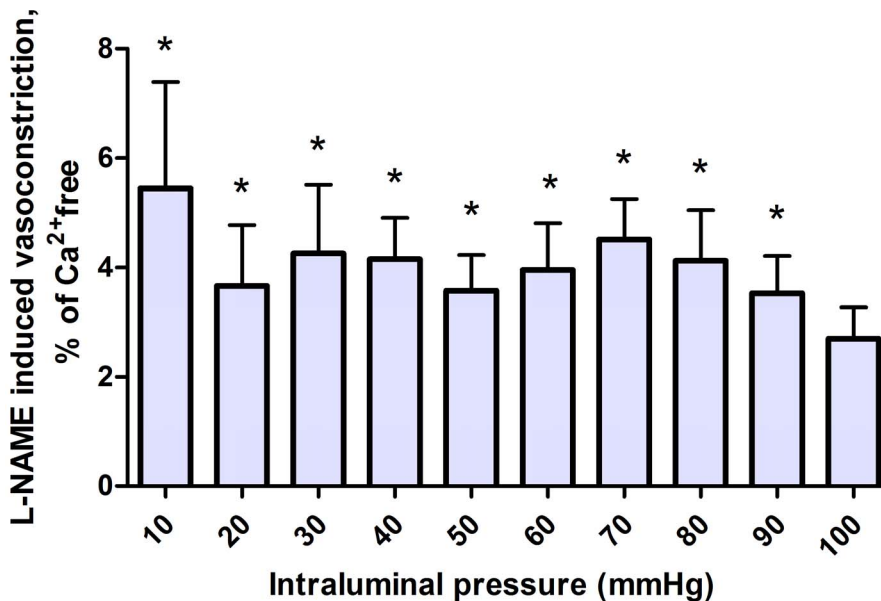


Figure 6. Vasoconstriction induced by L-NAME in coronary arteriole segments pretreated with quercetin and indomethacin. Concentrations: L-NAME: 10^{-4} moles/lit, quercetin: 10^{-7} moles/lit, and indomethacin: 10^{-5} moles/lit. * $p < 0.05$ with ANOVA doi:10.1371/journal.pone.0105587.g006

Results

In the first series, the vessel segments incubated in oxygenized PSS exhibited a substantial spontaneous tone. At 50 mmHg intravascular pressure, the average inner diameter was $127 \pm 25 \mu\text{m}$, which corresponds to a $20.2 \pm 4.3\%$ contraction from fully relaxed state. Increasing concentrations of quercetin induced increasing relaxations. In 10^{-9} moles/lit concentration, statistically significant relaxation was seen, while 10^{-5} moles/lit fully relaxed the segments. This relaxation (reduction of spontaneous tone) was found to be reversible, as original tone returned upon washing off the substance (Fig. 1).

In the second series of experiments, 10^{-7} mol/lit quercetin concentration was further tested on pressure-radius curves in the pressure range of 10–100 mmHg. A massive dilation was found at all pressure levels studied (Fig. 2). There was no significant correlation between the extent of quercetin-dilation and the morphological diameter of the segments (Fig. 3).

The COX blocker indomethacin, in a concentration of 10^{-5} moles/lit induced a significant contraction of all quercetin treated segments at all pressure levels (Fig. 4). This effect seems to depend on morphological lumen size: a statistically significant negative correlation between morphological diameter (Ca^{2+} -free solution, 100 mmHg pressure) and indomethacin induced contraction at higher pressures (60–100 mmHg) was found (Fig. 5).

Finally, the NO-blocker L-NAME was given (10^{-4} moles/lit) to test the level of endothelial vasodilation. Significant contraction was found, but its extent varied only between 2–8% of relaxed diameter (Fig. 6).

Discussion

In this study it was proved that the polyphenol quercetin elicits a direct concentration-dependent vasodilation on intramural coronary resistance arteries at physiologically realistic concentrations. In the very low concentration of 10^{-9} moles/lit a slight but significant relaxation was found in the intrinsic spontaneous tone. Equivalent to the physiological concentration range of 10^{-7} moles/lit in humans [7] resulted in $10.1 \pm 1.5\%$ elevation in the inner diameter. Close to the therapeutic concentrations of 10^{-6} moles/lit quercetin [7], the spontaneous vascular tone almost fully diminished. Quercetin, it proves, can be listed among potent known coronary vasodilators. A direct vasodilatory action of quercetin has been demonstrated until now exclusively on ring preparations of larger vessels, on major coronary arteries [13–17,23,24], as well as on resistance size mesenteric [17], and coronary arteries [25]. The pressure microangiometry method we applied was able to demonstrate vasodilatory actions of quercetin at physiologically and pharmacologically realistic concentrations, at real intraluminal pressures and inducing hemodynamically substantial alterations in inner diameter.

In addition, we found, that relaxation caused by 10^{-7} moles/lit quercetin on the small coronary arterioles does not depend on vessel diameter in the range of 70–250 μm , all samples relaxed by appr. 10 percents. It means that the spontaneous tone was reduced to its half in each segment simply by the presence of physiological concentrations of quercetin in the tissue bath. As smaller coronary resistance arteries have larger tone, the vasorelaxation induced on smaller vessels were more potent. Considering Hagen-Poiseuille's law, this might induce approximately 22–28% increase in flow at

larger arterioles (over 150 μm diameter), and 42–50% increase at smaller coronaries (under 150 μm diameter).

Our measurements provide some further insight into the possible mechanism of this vasodilatory action. Quercetin treated coronary arteriole segments contracted at all transmural pressures studied in response to the cyclooxygenase (COX) blocker indomethacin, proving the presence and activity of vasodilatory endogenous prostanoids in the vascular wall under the polyphenol effect. According to earlier observations, without quercetin, similar preparations produce mostly vasoconstrictor prostanoids when challenged with increasing intraluminal pressures [26]. This points to one potential mechanism of the vasodilation: altered balance toward vasodilatory prostanoids in the coronary resistance artery wall. As to the molecular mechanism, quercetin was reported to be a modulator in the COX enzymatic reaction [19], in this way affecting the endogenous prostanoid production [26,27]. Our results are in good agreement with these observations. Pharmacological modulation of vasoactive endogenous prostanoid production in the vessel wall has been demonstrated earlier with our contribution [26,27].

Addition of the NO blocker L-NAME to the tissue bath induced vasoconstriction, but its extent (2–8% at different pressures) does not indicate a substantial NO-dependent vasorelaxing effect of quercetin. Namely, it is not larger than the L-NAA induced contractions of similar segments with indomethacin in the bath without the presence of quercetin [26]. It is well-known that quercetin exhibits antioxidant [28,29] and ROS scavenging properties [30]. This can help to preserve NO, and provide longer half-life time, to express vasodilatation [31].

In conclusion, these experiments suggest that physiological circulating levels of quercetin continuously keep activated a coronary vasodilatory system, of which significance and extent is comparable with those of several other physiological mechanisms of coronary resistance control. Our observations suggest that production of endogenous vasodilatory prostanoids depends on the presence of such reducing agents as the polyphenols. Here, we can cite the original view of the Nobel-prize winner Hungarian biochemist Albert Szentgyörgyi, who suggested to recognize dietary polyphenols as vitamins [32] (“vitamin P” he called them). This view was challenged later because of absence of manifest deficiency symptoms [33]. We assume that the prevailing massive metabolic control (hypoxic vasodilation) of cardiac tissue perfusion easily overrides the missing polyphenol-relaxation, this way masking deficiency symptoms. However, if under pathological circumstances, the balance of vasoconstriction-relaxation is disturbed, it might reduce remaining control capacity. Maintaining a physiological quercetin concentration in the blood plasma may help correct the balance toward vasorelaxation in coronary arterioles.

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Author Contributions

Conceived and designed the experiments: AM-K EM GyLN. Performed the experiments: AM-K. Analyzed the data: AM-K GyLN. Contributed reagents/materials/analysis tools: EM GyLN. Contributed to the writing of the manuscript: AM-K EM GyLN.

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