Why Do We Need a Selective Angiotensin II Type 2 Receptor Agonist?

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Editorial Commentary

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Nevertheless, there is accumulating evidence that the commonly accepted scheme with a balance between a constrictor effect of AT1R and a dilator effect of AT2R is too simplistic and does not always reflect experimental evidence. The work performed by Verdonk et al,1 published in this issue of Hypertension, on the AT2R agonist C21 further highlights the complexity of this system, in addition to bringing key information on the mechanism of action of this new drug. Although C21 binds to AT2R with high affinity, this study also shows that C21 induces a puzzling combination of vasodilatation and vasoconstriction. Surprisingly, C21-mediated vasodilation was independent of AT2R and of the endothelium. Indeed, the authors have shown that C21 has a direct inhibitory effect on calcium influx into smooth muscle cells, thus leading to relaxation. C21 was tested in rat and mouse arteries, as well as human coronary arteries, and C21-dependent vasodilatation through calcium entry inhibition was also observed in mice lacking the gene encoding for AT2R. Furthermore, in the isolated perfused heart, C21 induces an initial AT1R-dependent contraction followed by AT2R-independent dilatation. The constrictor effect of AT1R was more pronounced in hypertensive rats, as shown previously.6

This article brings up 2 major issues requiring further discussion. First, C21-dependent dilatation relies mainly on calcium entry blockade without interfering with the RhoA-Rho-kinase pathway, which has been shown to be inhibited after AT1R stimulation. This finding requires reanalysis of the studies showing protective effects of C21. Indeed, the organ-protective effect of C21 is compatible with the effects of the dihydropyridine calcium channel blockers, that is, reduction of proliferation, inflammation, fibrosis, and vasoconstriction. The authors also show that low doses C21 are needed to selectively block AT2R. Lower doses of C21 might be used after AT1R blockade, which induces a large increase in AT2R expression level. In this condition, lower doses of C21 might be sufficient to more selectively block AT2R. Lower-dose C21 could, thus, be useful in resistant hypertension, although in this case its effect on calcium currents might also be advantageous. Low-dose C21 might also provide additional protection against organ damage in kidney and heart diseases in patients treated with angiotensin receptor blockers. Nevertheless, no data are yet available to support this assumption. It should be noted that more and more patients, not only hypertensive patients, receive angiotensin receptor blockers. In these conditions, a better knowledge of the effect of AT2R stimulation is essential, especially in the different organs affected by cardiovascular diseases. Second, this work further confirms with straightforward experiments...
the complexity of AT1R physiopathology. That AT1R stimulation induces dilatation or contraction depending on the type of artery or on the disease makes it difficult to use tools targeting this receptor.

Indeed, the findings of this article raise several important questions, the answers to which await further investigation. First, is there enough AT2R to be stimulated in the absence of AT1R blockade or in the absence of other factors increasing AT1R expression level and, second, is the effect of AT1R stimulation enough predictable. Although AT1R do not seem to desensitize, its stimulation does not induce important dilatation, and, in some cases, vasoconstriction may occur as observed in hypertensive6 or in aged rats.7 Furthermore, AT2R-dependent dilatation may be stronger and perhaps more persistent in female than in male rats.8 However, in the present study, coronary arteries from 2 men and 3 women were used without obvious difference attributed to gender, so more attention should be given in future studies to a possible sex difference in the effects of AT1R stimulation. Indeed, AT2R may have a role, other than the estrogen receptors, in the protection against cardiovascular events provided by female hormones before menopause. Of course, this very hypothetical issue remains to be further investigated.

That the effect of AT2R stimulation varies so much according to tissue type, age, sex, or diseases remains puzzling. Nevertheless, a more direct interaction between receptors may provide, at least in part, an explanation. Indeed, AT2R-AT1R dimerization might occur and affect the consecutive signal transduction. Although this concept still lacks functional evidence, receptor dimerization should be taken into account in future studies, especially because it may affect receptor internalization and intracellular signaling. Internalization of AT2R does not seem to occur, whereas AT1R internalization follows rapidly its stimulation.9 Consequently AT2R-AT1R heterodimerization might affect, and eventually prevent, AT1R intracellular signaling. Finally, although the role of internalized AT1R is not yet fully understood, angiotensin II receptor, AT1R, AT2R, and AT7R, have been identified in the nucleus and in mitochondria.10 Once again, the AT2R differs from the other receptors because it lacks the canonical nuclear localization sequence. Nevertheless, in isolated nuclear or mitochondrial fractions, AT2R stimulation with CGP4211A induces NO production, which is suppressed by PD123319 and by Nω-nitro-L-arginine methyl ester.10 Although the role of this nuclear renin-angiotensin system remains mainly unknown, it may add to the diversity of the effects of AT2R stimulation and hopefully help in better understanding its physiology.

Thus, reinterpretation of recently published articles using C21 in animal models of cardiovascular diseases is required. Nevertheless, it remains that C21 has beneficial effects against organ damage, although the precise mechanism of action is now less clear. Indeed, unlike C21, many commonly prescribed drugs are not well known, and their mechanism of action remains a matter of debate. Most importantly, much remains to be discovered concerning the role of AT2R and the mechanisms involved in the cardiovascular system.

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