



BRIEF QUESTIONS
AND ANSWERS
ON CURRENT
CLINICAL
CONTROVERSIES

Q: Can angiotensin II receptor blockers be used in patients who have developed a cough or angioedema as a result of taking an ACE inhibitor?

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A: ANGIOTENSIN II RECEPTOR BLOCKERS (ARBs) appear to be an appropriate alternative for patients who have developed an intolerable cough or angioedema while taking an angiotensin-converting enzyme (ACE) inhibitor. The incidence of these adverse effects is much lower with ARBs than with ACE inhibitors; moreover, small clinical studies have shown that ARBs lower blood pressure just as well as ACE inhibitors. Larger clinical trials that are underway should further clarify the efficacy and safety of ARBs compared with ACE inhibitors.

■ COUGH IS COMMON WITH ACE INHIBITORS

ACE inhibitor-induced cough occurs twice as often in women as in men and more frequently in nonsmokers. Although post-marketing surveys indicated that cough occurs in only 0.1% to 3% of patients who take an ACE inhibitor, two randomized controlled trials found the incidence to be much higher.^{1,2} Conservative estimates suggest that about 10% of patients treated with an ACE inhibitor develop this problem.

The cough usually begins within 1 to 2 months after starting therapy. It is dry, persistent, and mostly nonproductive, and it is often described as a “tickling in the throat.” It may occur more frequently when a person reclines, and it usually resolves 5 to 7 days after the drug is discontinued.

The mechanisms responsible for ACE

inhibitor-related cough are not fully understood, but kinins and prostaglandins have been suggested as likely mediators.³ Both bradykinin and prostaglandins are tussive agents, and some prostaglandins have been shown to increase both bronchoconstriction and cough response. These observations seem to mesh with the observed effects of ACE inhibitors, which are associated with an accumulation of bradykinin and substance P and increased prostaglandin production. Evidence suggests that nonsteroidal anti-inflammatory drugs such as sulindac and indomethacin, which inhibit prostaglandin synthesis, can reduce or abolish this adverse effect.^{4,5}

■ ANGIOEDEMA: RARE BUT POTENTIALLY LETHAL

ACE inhibitor therapy is one of the most common causes of angioedema—nonpitting edema of vascular origin that is confined to the head and neck. The edema can affect the floor of the mouth, tongue, larynx, lips, and face.⁶ ACE inhibitor-induced angioedema occurs most frequently in African American patients.

Generally speaking, angioedema is not a common adverse effect of ACE inhibitor therapy. It is nonetheless important because it can be potentially lethal if it develops in the tongue, glottis, or larynx. For that reason, patients should not be switched to another ACE inhibitor if this complication occurs.

■ ARBs AS AN ALTERNATIVE


Angiotensin II receptor blockers, a newer class of drugs, are distinct from the ACE inhibitors in that they have no effect on angiotensin-

ARBs cause fewer side effects than do ACE inhibitors



converting enzyme. Rather, they block the diverse physiologic effects of angiotensin II by specifically blocking angiotensin II tissue receptors.⁷ Currently, there are 6 ARBs approved by the Food and Drug Administration: candesartan (Atacand), eprosartan (Teveten), irbesartan (Avapro), losartan (Cozaar), telmisartan (Micardis), and valsartan (Diovan).

These drugs are not associated with an increase in circulating bradykinins or an increase in prostaglandin production, so theoretically, they should cause less of a problem with cough. In fact, that is what clinical studies of ARBs have found. Overall, the studies showed that the frequency of cough in patients taking ARBs was statistically significantly lower than that in patients taking ACE inhibitors. The frequency was comparable to that observed in the placebo controls.

Although a few cases of angioedema did occur during clinical trials of the six ARBs, there was no clear pattern of occurrence.^{8,9} However, a recent review of case reports documenting ARB-induced angioedema reported that one third of the patients had experienced a prior episode of ACE inhibitor-induced angioedema.¹⁰ In view of these observations, ARBs should be used with caution in patients with a prior history of angioedema. 

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