INTRODUCTION

Radiocontrast-induced nephropathy (RCIN) is considered to be the third most common cause of in-hospital acquired acute renal failure, and is associated with in-hospital and long term high morbidity and mortality. RCIN is usually defined as acute deterioration of the renal function that occurs within 72 hr after intravascular administration of contrast agents, and is characterized by an increase in serum creatinine of 25% or > 0.5 mg/dl from the baseline level, in the absent of another etiology. Strategies to diminish the incidence involve hydration, pharmacological manipulation with renal...
vasodilators, endothelin receptor antagonists or antioxidants, and even prophylactic continuous hemofiltration or hemodialysis\textsuperscript{1,6,8,10}. Some studies have shown that human atrial natriuretic peptide (hANP) has renal protective effects\textsuperscript{2,11-13}, but the beneficial effects for RCIN prevention have not been clearly shown. We treated a diabetic patient with these strategies for preventing RCIN and compare the results.

**CASE REPORT**

A diabetic, hypertensive 66-year-old man was admitted to the hospital for evaluation of angina pectoris. Treadmill exercise test showed ST-segment depression in leads $\textcircled{a}, \textcircled{b}$, $\textcircled{aR}, \textcircled{RA}$, $\textcircled{F}, \textcircled{4}$, $\textcircled{6}$. Laboratory test showed hemoglobin A\textsubscript{1c} of 4.5\% while taking glibenclamide 2.5 mg/day. Serum blood urea nitrogen was 22 mg/dl and serum creatinine was 1.26 mg/dl. Estimated creatinine clearance was 55 ml/min. Coronary angiography revealed a long diffuse stenosis of 90\% in the left anterior descending artery segments 6 to 7. Percutaneous coronary intervention (PCI) was scheduled 6 days after diagnostic catheterization. The preventive strategy for RCIN used hANP (0.025 g/kg/min) co-administered with 0.9% saline (60 ml/hr) starting 2 hr prior to the PCI and continued for 24 hr. During the intervention, the rotablator was required to remove the diffusely calcified plaque and diagonal branch protection was performed with left anterior descending artery stenting. A total of 450 ml of non-ionic contrast medium was used. The serum creatinine level increased to 27\% above the baseline 48 hr later, but decreased to the previous level rapidly (Fig. 1). The clinical course was good and the patient was discharged without any symptoms.

Four months later, the patient came to our hospital with recurrent chest pain. On admission, blood urea nitrogen was 20 mg/dl, and serum creatinine was 1.71 mg/dl with estimated creatinine clearance 40 ml/min. Diagnostic catheterization was planned with possible on-site PCI. Reno-protection used only 0.9% saline hydration; starting 2 hr before the procedure at 100 ml/hr, then reduced to 60 ml/hr after PCI and continued for 24 hr.

The finding was a diffuse in-stent restenosis with 99\% occlusion (Fig. 2). Plain balloon angioplasty was successfully performed as treatment. The whole procedure (coronary angiography, left ventriculography and PCI) required 220 ml of contrast medium. A great increase in the serum creatinine from baseline level was observed with a maximum peak at 4th day (6.38 mg/dl, 273\%), taking 2 weeks to return to the previous level (Fig. 1).

Unfortunately, 3 months later, the patient came again to the emergency room with increasing chest pain with dyspnea. Electrocardiography showed 2 mm ST elevation in $\textcircled{1}, \textcircled{2}$. Troponin T was positive but creatine kinase-MB was within the normal range. Blood urea nitrogen was 28 mg/dl, serum creatinine was 1.63 mg/dl and creatinine clearance was 35 ml/day. Echocardiography revealed severe
deterioration of left ventricular function. At that time, more aggressive renal protection was considered necessary. Continuous hemofiltration with fluid replacement of 1,000 ml/hr starting 2 hr before the PCI and continued for 18 hr was performed. Coronary angiography showed 99% in-stent restenosis and cutting balloon angioplasty was performed. Volume of contrast media was 180 ml. Serum creatinine increased to 1.95 mg/dl (20% from baseline) at 72 hr after intervention and returned to the previous level after 6 days (Fig. 1).

DISCUSSION

Guidelines to prevent RCIN emphasize the importance of identifying patients at high risk before intervention\(^{14,15}\). Contrast volume should be reduced and volume expansion with saline infusion should be considered to assure a good renal perfusion. Moreover, the use of nephrotoxic drugs should be avoided\(^{14}\).

The most popular method to prevent RCIN is volume expansion with saline infusion (1 ml/kg-body weight/hr) starting before the use of contrast medium and continued afterward\(^{15,16}\). However, the best benefits are observed only in patients with mild renal impairment and this method is not suitable in all patients.

Some studies demonstrated that hANP has a protective effect for various etiologies of renal dysfunction\(^{10,17,18}\). hANP has natriuretic effects, increasing glomerular filtration rate by dilating afferent arterioles while constricting the efferent arterioles. hANP also antagonizes the secretion of renin, aldosterone and vasopressin. Animal experiments have suggested its usefulness in RCIN\(^{21}\). However, clinical trials could not show substantial effects, with conflicting results especially in diabetic patients\(^{9,10,18}\). This discrepancy may be due to the presence of heterogeneity in the response to hANP for each patient.

The present patient was at high risk of developing RCIN, and underwent coronary intervention repetitively, so we had the chance to assess the effects of different strategies. Saline + hANP protocol was apparently better than only saline, considering the lower increase of serum creatinine and rapid recovery to the previous level. But there are some limitations for the comparison: the renal function pre-interventional statuses were not exactly the same, and the amount of contrast media used in the hANP protocol (450 ml) was almost twice as much as that in only saline. In spite of these conflicting factors, we believe that the big difference in the increase of serum creatinine after intervention between two strategies is strongly indicative.

At the third intervention, we selected prophylactic continuous hemofiltration to obtain effective renal protection without volume overload. We had a relative low increase of serum creatinine (19%) with return to the previous level within 1 week. Continuous hemofiltration starting before PCI and continued for 24 hr afterward reduced the incidence of RCIN\(^{13}\). The main feature of the protective effect was attributed to the high volume of fluid administered, without fluid loss or volume overload, and maintaining hemodynamic stability. This also leads to maintenance of renal perfusion and rapid dilution of contrast medium. However, the complexity and high cost involved in this method is the biggest disadvantage as a routine way of preventing RCIN.

Comparing all distinct variables at each intervention, we think that hANP infusion with saline hydration is an effective method to prevent RCIN and was as effective as hemofiltration in our patient. With further studies, we should clarify what kind of patient would most benefit from hANP infusion.
要約

冠動脈形成術による造影剤腎症に対しヒト心房性ナトリウム利尿ペプチドが有用であった1例

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経皮の冠動脈形成術施行後に腎機能はしばしば悪化する。糖尿病やうっ血性心不全を合併している腎障害例では、その頻度が著しい。造影剤が急性肾障害を軽減すべき種々の方法が試みられているが、その効果は一様でない。我々は糖尿病性腎症を合併した66歳、男性に対し、冠動脈形成術を施行し、おのおのの治療方法により腎障害軽減効果を比較することができた。結果は、生理的食塩水のみによる輸液は効果が少なかったのに対し、ヒト心房性Na利尿ペプチド + 生理的食塩水による輸液および血液濾過が良好な腎保護効果を示した。

References


