

크레메진이 만성 콩팥병 진행에 미치는 영향

국립중앙의료원 내과¹, 연세대학교 의과대학 내과학교실², 고려대학교 안산병원 내과학교실³
가천의과대학교 길병원 내과학교실⁴, 가톨릭대학교 의정부성모병원 내과학교실⁵, 보라매병원 내과학교실⁶
인제대학교 일산백병원 내과학교실⁷, 가톨릭대학교 서울성모병원 내과학교실⁸, 분당서울대학교병원 내과학교실⁹
한림대학교 평촌성심병원 내과학교실¹⁰, 건국대학교병원 내과학교실¹¹, 서울대학교병원 내과학교실¹²

차란희¹, 강신욱², 차대룡³, 장재현⁴, 윤선애⁵, 임춘수⁶, 한상엽⁷
박철휘⁸, 나기영⁹, 김성균¹⁰, 박정환¹¹, 김연수¹²

The Effect of AST-120 (Kremezin[®]) on the Progression of Chronic Kidney Disease in Korea

Ran-hui Cha¹, Shin Wook Kang², Dae Ryong Cha³, Jae Hyun Chang⁴, Sun Ae Yoon⁵, Chun Soo Lim⁶
Sang Yeop Han⁷, Cheol-whee Park⁸, Ki Young Na⁹, Sung Gyun Kim¹⁰, Jung Hwan Park¹¹, Yon Su Kim¹²

Department of Internal Medicine¹, National Medical Center
Department of Internal Medicine², Yonsei Medical School Severance Hospital
Department of Internal Medicine³, Korea University Ansan Medical Center
Department of Internal Medicine⁴, Gachon University Gil Hospital
Department of Internal Medicine⁵, St. Mary's Hospital, EuijungBu
Department of Internal Medicine⁶, Seoul National University Boramae Hospital
Department of Internal Medicine⁷, Inje University Ilsan Hospital
Department of Internal Medicine⁸, St. Mary's Hospital, Seoul
Department of Internal Medicine⁹, SNUH Bundang Hospital
Department of Internal Medicine¹⁰, Hallym University Pyoung Chon Hospital
Department of Internal Medicine¹¹, Konkuk University Hospital
Department of Internal Medicine¹², Seoul National University Hospital

Background: AST-120 is an oral adsorbent excreting uremic toxins including indoxyl sulfate into feces. And AST-120 was effective in reducing the renal function decline and proteinuria as well as in improving the pathologic changes of renal diseases. However, AST-120 did not prove the capability to delay the initiation of renal replacement therapy in chronic kidney disease (CKD) patients. We aimed to find the long-term effect of AST-120 on the renal progression (doubling of SCr, creatinine clearance (CCr) decrease more than 50%, or initiation of renal replacement therapy) in patients with moderate to severe CKD.

Methods: We prospectively recruited 579 patients (CKD stage 3 and 4) from 11 medical centers in Korea and randomized them into AST-120 and control arm through the mixed block randomization as well as being stratified according to both the gender and the cause of CKD. A total of 6 gram of AST-120 in 3 divided doses was given to participants in AST-120 arm as well as standard conventional treatment. They were followed up every 3 months up to 36 months.

Results: A total of 465 patients were evaluated. Medication compliance of AST-120 was 89.9±11.82% (median 93.9%). Mean SCr and estimated CCr level was 2.81±0.666 mg/dl and 26.79±7.263 ml/min/1.73m², respectively. There was no significant difference in the occurrence of composite primary outcomes between two treatment arms (Log-rank p=0.330). Diabetic nephropathy and renal dysfunction were risk factors for the occurrence of primary outcomes and both additively affected (Log-rank p<0.001). The velocity of estimated CCr decline was less in AST-120 arm and this was especially profound in patients with diabetic nephropathy. The slope of 1/SCr significantly decreased in both two treatment arms and the value of AST-120 arm was less than that of control arm (p<0.05). Urinary protein excretion rate was decreased from 2.04±1.980 g/g Cr to 1.79±2.023 g/g Cr in control arm and from 1.97±2.053 g/g Cr to 1.234±1.198 g/g Cr in AST-120 arm (p<0.05).

Conclusions: AST-120 slowed the renal function decline although it did not affect on the occurrence of primary outcome. Longer period of observation of the participants with or without AST-120 is needed.

Key Words: 만성콩팥병, 크레메진, 효과

Chronic kidney disease, Kremezin, Effect