

Intrahepatic Arterial Injection of Holmium166-Chitosan Complex in the Hepatoma Treatment: A New Therapeutic Modality

NC Yoo^{1,2,5}, JT Lee³, JD Lee⁴, HC Chung^{1,2}, JH Kim^{1,2}, JK Roh^{1,2},
JS Min¹, YM Moon², KH Kim⁵, B S Kim¹, KB Park⁶

Yonsei Cancer Center/Res Inst¹, Depts of Int Med², Diag Radiol¹, Nucl Med², Clin Pharmacol¹, Yonsei Univ Coll of Med,
Korea Atomic Energy Res Inst³, Seoul, Korea

About 80% of hepatoma patients are diagnosed as an advanced inoperable stage. At present the patients are dependent only on hepatic artery embolization or intra-arterial chemotherapy. Holmium-166 is mainly a beta-ray emitter (E_{max} : 1.84MeV, $t_{1/2}$: 26.8hr) with 90% absorption in 2.3mm of tissue and maximum 8mm in depth. Holmium166-chitosan complex (Ho166-C) is less permeable. For small hepatoma (<3cm in diameter), we obtained 86% response rate by percutaneous Ho166-C injection. For a larger tumor (>3cm), multiple injections are needed, which resulted in decreased distribution homogeneity and responsiveness. We tried intrahepatic arterial Ho166-C injection in 45 nodular type hepatoma. Mean dose of Ho166-C was 103.25mCi. The pharmacokinetics after injection of 100, 150 and 200mCi Holmium166-C were: $C_{max}(nCi)$ 31.69, 63.91, 75.03; $AUC(nCi)$ 122.27, 284.52, 219.62; $Cl(L/min)$ 0.187, 0.527, 0.911, respectively, and T_{max} were 0.25(hr) in all dose. Among them, 34 patients(75.5%) showed successful response. Five (13.3%) had no response and one(2.2%) were stable. Reversible bone marrow suppression was main complication, which was evident in non-responsive patients. These results suggest that intrahepatic arterial injection of Ho166-C could be a highly effective and safe treatment of advanced hepatoma. Further comparative randomized trials are needed.