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VIRAL HEPATITIS

#P31 - Fibrosis is not regressed 5 years after HCV and HBV treatment, MAFLD can be the cause

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Background & Aims

Fibrosis index-4 (FIB-4) has been established as non-invasive test for identifying severe fibrosis or high risk of liver-related event in patients with chronic viral hepatitis. In metabolic associated fatty liver disease (MAFLD), FIB-4 had better diagnostic accuracy for estimation of liver fibrosis among serum markers. We aimed to assess fibrosis regression after viral eradication to understand fibrogenesis.

Methods

This partially retrospective partially prospective cohort was conducted on 100 patients (50 were chronic hepatitis C «CHC» received and finished direct acting anti-viral drugs and achieved sustained virologic response and 50 were chronic hepatitis B «CHB» started and still on nucleos(t) ide analogues with undetected viremia). Demographic, clinical, laboratory, radiological data, treatment regimens and virus status were recoreded.

Results

Sixty five patients were male, mean age 49+/-12.08 years, mean duration after starting treatment 5.09+/-1.55 years. CHC patients, 84% received sofosbuvir and daclatasvir for 12 weeks, CHC patients, 78% receiving tenofovir. Thirty six were dyslipidemic, 7% diabetic and 4% hypertensive. Clinically, 59% were obese according to waist hip ratio (WHR). Mean waist circumference (WC) and WHR were 101.46+/-13.09 and 0.89+/-0.05 cm respectively. Mean alpha feto protein (AFP) pre treatment and at time of enrollment 50.99+/-277.38 and 24.15+/-126+/-72 ng/ml respectively. FIB-4 before treatment and at time of enrollment was 2.16+/-2.66 and 3.03+/-4.05 respectively. Liver by ultrasound examination was homogenous in texture and bright echopatteren in 95% of patients. CHC group were more diabetic with higher AFP and FIB-4 both before and at time of enrollment compared with CHB group. In CHC group, 34% dyslipidemic, 38% obese and FIB-4 before and at time of enrollment was 2.38+/-2.63 and 3.86+/-5.13 respectively. In CHB group, 38% dyslipidemic, 66% obese and FIB-4 before and at time of enrollment was 1.93+/-2.68 and 2.20+/-2.32 respectively.

According to FIB-4, patients with mild to significant fibrosis increased from 30 to 34% and patients with advanced fibrosis increased from 14 to 22% before and 5 years of treatment. Patients with progressed fobrosis fulfilling MAFLD criteria.

Conclusions

Fibrogenesis is a dynamic process and can be progressed in patients with chronic viral hepatitis even after viral eradication and suppression. MAFLD can be the cause.

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