Poster

HEPATITIS C VIRUS INFECTION AMONG PATIENTS AT HOSPITAL TENGKU AMPUAN AFZAN, KUANTAN: GENOTYPES DISTRIBUTION, RNA SEQUENCE VARIATION, RISK FACTORS AND CO-MORBIDITIES

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Hepatitis C virus (HCV) genotyping is very important for the clinical management of HCV-infected patients. HCV strains demonstrate a high degree of genetic heterogeneity. The 5' un-translated region (5'UTR) of HCV is the most highly conserved region of the virus genome, making this region ideal target for HCV RNA amplification-based detection assays. Additional target to determine the genotypes of HCV is the non-structural (NS) 5B region of the viral genome. The aim of this study was to determine HCV genotypes and identify sequence variances in common genotypes as well as their association with certain factors. Nucleotide sequence analysis of 5'UTR and NS5B region were performed to identify genotypes of 40 HCV infected patients attending Gastroenterology Clinic, HTALL Kuantan Pahang. Of the 40 samples, 39 (97.5%) tested positive for HCV RNA RNA-positive isolates were subjected to direct sequencing and the results were compared with reference sequences from NCBI homepage. Based on the analysis of both of the above regions, 31 (77.5%) isolates were successfully classified into their genotypes and subtypes; 3a (57.5%), 1a (12.5%), 3b (2.5%) and 1b (2.5%) No genotype 2, 4, 5 and 6 were found in this study. However, there was one mixed genotype (3a/1a) HCV infection. Sequence variations were analysed in both regions, NS5B region demonstrated greater genetic variability in its sequence man 5'UTR. No significant association was found between HCV genotypes with patients demographic data, risk factors, alcohol consumption and co-infection with HBV == HIV. In conclusion, the use of the 5'UTR and NS5B-based assays proved to be a sensitive and reliable method for HCV RNA detection and genotyping. Genotype followed by genotype 1 were the common HCV genotypes found in this state population. The base sequences of both genotypes varied at certain locations both the 5'UTR and NS5B regions however NS5B region sequences displaced higher heterogeneity. Furthermore, no risk factors and co-morbidities were linked to HCV genotypes.

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