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Development of vaccine for "hepatitis C" (Papaya mosaic virus as an expression vector for foreign epitopes)

Varsha Raja* and MG AbouHaidar

Address: Laboratory of Virology, Department of Botany, University of Toronto, Toronto, Canada

* Corresponding author

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This work focuses on the use of Papaya Mosaic Virus (PMV), a plus-sense RNA potexvirus that forms rod-shaped particles, for high yield production of Hepatitis C virus (HCV) epitopes. We produced PMV clones in which an epitope from the HCV core gene was inserted into the N-terminus of the PMV capsid gene. Furthermore the infectivity and functionality of these clones and whether they retained HCV sequences post infection was assayed. The preliminary results indicate that the clones are weakly infectious and in those cases that infections do occur and lesions are produced, it seems that the HCV sequence is lost. This could be due to homologous recombination in clones containing two copies of the capsid gene. Future work will focus on optimization of foreign inserts through further work on PMV clones containing one copy of the capsid gene in which the HCV sequence has been inserted.