Detection of viral replication in bees

Ivan Meeus¹, Dirk de Graaf², Guy Smagghe¹

¹Lab of Agrozoology, University of Ghent, Coupure 653, 9000 Ghent Belgium,

Email: ivan.meeus@UGent.be

²Laboratory of Zoophysiology, Department of Physiology, Faculty of Sciences, Ghent University,

Krijgslaan 281 (S33), B-9000 Ghent, Belgium

DOI: 10.5073/jka.2012.437.051

Abstract

Recently foraging bees were discovered with pollen loads infected with honey bee viruses uncorrelated with the infection status of the bees itself. This observation has wide implications on the broadly used PCR viral detection techniques. False positives results could be obtained if viral remnants from infected pollen in the bee gut are detected. Integration of the real time PCR technology could help to eliminate these false positive results, however techniques detecting viral replication ultimately prove presence of active viruses. We demonstrated that current minus strand detection methodology often is not selective enough to differentiate between positive RNA strands (inactive virus) and minus RNA strands (replication virus) and provide possible solutions