

A MODIFIED DIRECT PCR PROTOCOL FOR DVI APPLICATIONS USING MICROFLOQ™ DIRECT SWABS FROM DECOMPOSING HUMAN REMAINS COUPLED WITH THE QIAGEN INVESTIGATOR 24PLEX GO! KIT

Coral Loockerman¹, MS, Sheree Hughes-Stamm^{1,2}, PhD, [Rachel Houston¹, PhD](#).

¹Department of Forensic Science, Sam Houston State University

²School of Biomedical Sciences, University of Queensland

Disaster victim identification (DVI) relies on rapid identification of decomposing human remains, often in remote areas without access to storage facilities. Collection of biological material using swabs may prove easier, more efficient, and more amenable to storage in harsh conditions. microFLOQ™ direct swabs have been identified as a potential alternative for more rapid collection and processing of DNA in forensic and DVI situations.

4N6FLOQSwabs™: Genetics and microFLOQ™ direct swab were used to collect DNA from red muscle via an incision in the arm or leg of a decomposing human cadaver. Traditional DNA processing with the Genetics swabs was compared to a direct amplification strategy using the microFLOQ™ swab coupled with the Investigator 24plex QS GO! Kit. Additionally, both swabs types were evaluated for their ability to store DNA for up to three months. The direct amplification strategy was optimized by pre-treating the swab (washing, vortexing, lysis) prior to amplification and slightly modifying the cycling parameters. As an alternate method, the microFLOQ™ swabs were used to sub-sample DNA stored on the Genetics swabs.

Results indicate that both swab types were able to store DNA at room temperature. STR success rates of traditional and direct PCR method were comparable but were highly dependent on the stage of decomposition and the sample location. Up to day 10, full profiles were obtained using both processing methods with samples taken from the leg. Full profiles were obtained from day 13 and day 20 using traditional methods, while partial profiles were obtained on day 13 using microFLOQ™ swabs and subsampling. The Quality Sensor (QS) markers were used to assess sample quality. Interestingly, the QS markers indicated that even after pre-washing the microFLOQ™ swabs, there was still inhibition present in the amplification for microFLOQ™ swabs that were dried overnight. However, when the microFLOQ™ swabs were processed within hours of swabbing or used to sub-sample, there was less to no inhibition indicated by the QS markers and profile completeness improved. Overall, microFLOQ™ swabs in conjunction with the GO! Kit facilitated direct processing in the laboratory from decomposing remains.