Corrigendum AIHTA Project Report No.: 172

It was brought to our attention that we misclassified the telemonitoring devices used in Austria by the company Telbiomed. According to the Medical Device Regulation, we initially reported that the medical device in use in Austria is a Class IIb medical device, while a Notified Body already classified the device as Class IIa. Hence, we corrected the necessary passages in the report, and we provide an updated version.