

<p>DL 5.4.8. Review adherence to case definition by site and draft report of corrective actions</p>	<p>The contractor shall review site-specific adherence to the syndromatic case definitions, report on and implement corrective actions.</p> <p>Comprehensive review of consistent adherence to the agreed ILI case definition, and the syndromatic definition for COVID-19 cases. The format of the reports shall be agreed with ECDC.</p> <p>The contractor will be responsible for the implementation of the study according to the study protocol. Datasets containing the collected data will be regularly shared with ECDC. The contractor will be responsible for the timely delivery of technical documents, of which the format will be agreed in advance with ECDC. The contractor shall ensure that all study sites are performing the study activities according to the agreed plan and will immediately report to ECDC in case of any issue or challenge in the implementation of the study.</p>	<p>Once during year 2</p>	<p>Word - ECDC template</p>
<p>DL 5.4.9. Regular laboratory results summary table</p>	<p>The contractor shall provide a quarterly laboratory results summary table to ECDC specifying the number of laboratory specimens processed and tested and the proportion of positive samples by study site. All study should be tested according to the EU/EEA protocol and country-specific SOPs with a validated laboratory assay (such as polymerase chain reaction, PCR) and report regularly overview results.</p> <p>This deliverable includes all costs of sample processing, transport, laboratory data management and the diagnostic assay and refers to cost of performing the molecular diagnostic (i.e. RT-PCR) and serology. The national public health authorities have access to ECDCs support for sequencing and antigenic characterisation of both SARS-CoV-2 and seasonal influenza viruses, which should be utilised, where relevant. ECDC will decide which countries to be offered support.</p> <p>It is expected that the contractor shall ensure that the laboratory components of the study will be implemented in each study site according to the study protocol, and that the epidemiological and clinical data will be collected using standard</p>	<p>Every four months</p>	<p>Word or a suitable electronic platform</p>