
Provisional Authorisation for COVID-19 Tests

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| <i>Applicant</i> | EUROIMMUN Anti-SARS-CoV-2 NCP ELISA (IgG) |
| <i>Name of test</i> | EUROIMMUN Anti-SARS-CoV-2 NCP ELISA (IgG) |
| <i>Intended purpose (As per manufacturer's information for use)</i> | <i>The enzyme immunoassay (ELISA) provides semiquantitative in vitro determination of human antibodies of the immunoglobulin class IgG against SARS-CoV-2 NCP in serum, EDTA, heparin or citrate plasma or dried blood spots (DBS) to support the diagnosis of SARS-CoV-2 infection and constitutes a supplement to the direct pathogen detection. Moreover, serology can be applied to collect epidemiological data. The product is designed for use as IVD and can optionally be processed on fully automated equipment. The format 96 x 20 has been specially designed for processing on the EUROLabWorkstation ELISA.</i> |
| <i>Date of Provisional Authorisation</i> | 5 August 2020 |
