

## Assay Performance Data

<b>Naam assay:</b>	Cardiolipine IgG & IgM As panel in serum [LOINC 24319-6]																		
<b>Methode, fabrikant:</b>	Fluorescence Enzyme Immuno Assay (FEIA) – EliA op Phadia 250, ThermoFisher Scientific																		
<b>Traceerbaarheid:</b>																			
Gekalibreerd naar	<p>Afkomstig uit de bijsluiter:  The calibration curve is obtained with EliA IgG/IgM Calibrators. The IgG/IgM Calibrators are traceable via an unbroken chain of calibrations to the International Reference Preparation (IRP) 67/86 of Human Serum Immunoglobulins A, G and M from World Health Organization (WHO).  The standardization of the assay is adjusted to a set of established standard sera (Harris et al., 1987). Results are expressed in GPL-U/ml and MPL-U/ml (1 GPL-Unit or MPL-Unit corresponds to the binding activity of 1 µg/ml of cardiolipin IgG or IgM antibody that was purified from the standard serum by affinity chromatography).</p>																		
<b>Referentie-interval of afkapgrenzen</b> (gebruik UCUM-eenheden)																			
Herkomst referentiewaarden	<p>Afkomstig uit de bijsluiter:  The ranges (negative, weak positive, positive) recommended for the evaluation of the results are:</p> <table border="1"> <thead> <tr> <th>Test</th> <th>Unit</th> <th>negative</th> <th>weak positive</th> <th>positive</th> </tr> </thead> <tbody> <tr> <td>EliA Cardiolipin IgG</td> <td>GPL- U/mL</td> <td>&lt;10</td> <td>10 -40</td> <td>&gt;40</td> </tr> <tr> <td>EliA Cardiolipin IgM</td> <td>MPL- U/ml</td> <td>&lt;10</td> <td>10 - 40</td> <td>&gt;40</td> </tr> </tbody> </table>				Test	Unit	negative	weak positive	positive	EliA Cardiolipin IgG	GPL- U/mL	<10	10 -40	>40	EliA Cardiolipin IgM	MPL- U/ml	<10	10 - 40	>40
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<b>Stabiliteit monster</b>																			
Kamertemperatuur 4 °C -20 °C	<p>Ja/Nee*, termijn: 2 dagen  Ja/Nee*, termijn: 2 weken  Ja/Nee*, termijn: 12 maanden</p>																		
<b>Detectielimieten</b> (gebruik UCUM-eenheden)																			
LoD (Limit of Detection) LoQ (Limit of Quantitation)	N.v.t. bij autoantistofbepalingen																		
<b>Imprecisie</b> (gebruik UCUM-eenheden)																			
Concentratie (level invullen)	<p>IgG: Level interne controle: 33 U/mL, CV (%): 12,2%  IgM: Level interne controle: 28 U/mL, CV (%): 8,3%</p>																		
<b>Meetbereik</b> (gebruik UCUM-eenheden)																			
Meetbereik	<p>IgG: 0.5 - 418 U/mL  IgM: 0.8 – 472 U/ml</p>																		
<b>Extern QC programma</b>																			
(Inter)nationaal extern QC programma	Euroimmun Phospholipids																		

[\[<sup>^</sup>Top\]](#)

Indien niet beschikbaar, alternatieve aanpak	
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\* Doorhalen wat niet van toepassing is.

Ingevuld door: Caroline Roozendaal	Datum: 14-12-2018
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