

22-fb-9_11

Änd. Ind. 2

CoA
Certificate of Analysis

17-OH Progesterone

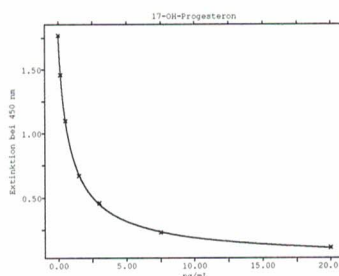
REF : EIA-1292**LOT** : 22K082-4
 : 2023 - 08 - 31
(yyyy - mm - dd)

Controls	Acceptance Range ng/mL	Observed concentration ng/mL	Standards (Calibrators)		OD at 450 nm (620 nm Ref)	Acceptance Range Max. OD
Name			ng/mL			
22KL042	0.34 – 0.88	0.52	S0	0.0	2.46	≥ 1.20
22KH042	3.72 – 9.82	7.32	S1	0.15	1.97	
			S2	0.5	1.44	
40411	0.44 – 1.15	0.58	S3	1.5	0.81	
40412	1.38 – 3.63	1.97	S4	3.0	0.56	
40413	4.28 – 11.3	6.42	S5	7.5	0.29	
			S6	20.0	0.15	

All materials of human origin have been tested for the presence of HBs antigen and antibodies to HIV 1, HIV 2 and HCV, and were found to be nonreactive by licensed tests in accordance with standard EN 13641.

Incubation times	60 min at RT / 30 min at RT
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Standard Curve



KIT COMPONENTS	VOLUME	LOT	yyyy-mm-dd
Microtiterwells	96 Wells	22W062-2	2024 - 06 - 30
Enzyme Conjugate	1 x 25 mL	22D082-3	2023 - 08 - 31
Standards	7 x 1 mL	22S042	2024 - 04 - 30
Control Low	1 x 1 mL	22KL042	2024 - 04 - 30
Control High	1 x 1 mL	22KH042	2024 - 04 - 30

KIT COMPONENTS	VOLUME	LOT	yyyy-mm-dd
Wash Solution	1 x 30 mL	TWP072	2024 - 07 - 31
Substrate Solution	1 x 25 mL	TMBSX13	2024 - 06 - 13
Stop Solution	1 x 14 mL	SD072	2024 - 07 - 31
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A sample of this product lot has been tested and found to meet all product specifications. This lot is released.

2022 - 08 - 26
Test Date (yyyy-mm-dd)

Quality Control Manager

Freigegebenes Originaldokument siehe Ablage QMB/QA:
„QMH Kapitel 9“

Seite 1 von 1