

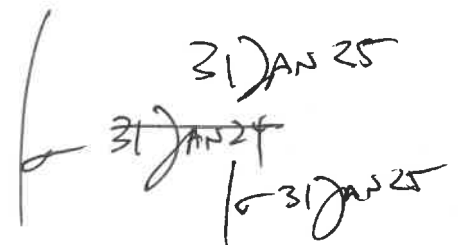
Order Details		Ordering Professional	Request / Accession	R-P61063673
Donor ID	14244	C4041 - San Diego Sperm Bank	Received	12/19/2024 10:40
DOB	01/19/1995	8950 Villa La Jolla Dr B214	Final Report	12/21/2024 09:44
Gender	M	La Jolla, CA 92037	Report Generated	12/21/2024 09:44
First Collected	12/18/2024		Time Zone	US Mountain Standard Time

Flagged Results

Test	Completed	Results	Ref. Range
CMV IgM using GSDX Automation	12/21/2024 09:32	Positive	Negative
CMV Total Ab	12/20/2024 14:17	Positive	Negative

Results

Test	Completed	Results	Ref. Range
Chlamydia trachomatis - Panther Platform	12/20/2024 05:24	Negative	Negative
Hepatitis B Core Total Ab	12/20/2024 09:57	Non Reactive	Non Reactive
Hepatitis B Surface Ag	12/20/2024 10:02	Non Reactive	Non Reactive
Hepatitis C Virus Ab	12/20/2024 10:00	Non Reactive	Non Reactive
HIV-1/HIV-2 Plus O	12/20/2024 09:54	Non Reactive	Non Reactive
HTLV I/II Ab	12/20/2024 09:55	Non Reactive	Non Reactive
Neisseria gonorrhoeae - Panther Platform	12/20/2024 05:24	Negative	Negative
Syphilis Screening – Nontreponemal (Automated)	12/20/2024 09:32	Non Reactive	Non Reactive
Ultrio Elite HBV	12/21/2024 08:52	Non Reactive	Non Reactive
Ultrio Elite HCV	12/21/2024 08:52	Non Reactive	Non Reactive
Ultrio Elite HIV-1/2	12/21/2024 08:52	Non Reactive	Non Reactive
WNV	12/20/2024 06:44	Non Reactive	Non Reactive



 31 JAN 25

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Sample Reference(s)

Collection Time Zone: Pacific Standard Time

Lab Sample ID	Type	Collected	Test(s)
S-P61063673A <i>Donor Type: Living</i>	Serum Red Top - 17mm	12/18/2024 11:30	2771, 2772, 2773, 31050
S-P61063673B <i>Donor Type: Living</i>	Plasma EDTA - 13mm	12/18/2024 11:30	32110, 32140, 32210, 35210, 35460, 3991
S-P61063673C <i>Donor Type: Living</i>	Plasma EDTA - 13mm	12/18/2024 11:30	35030
S-P61063673D <i>Donor Type: Living</i>	Plasma EDTA - 13mm	12/18/2024 11:30	2722
S-P61063673E <i>Donor Type: Living</i>	Aptima Urine Tube	12/18/2024 11:30	3674, 3675

Test Reference(s)

Code	Name	Methodology Description
3674	Chlamydia trachomatis - Panther Platform	The APTIMA COMBO 2 Assay is a target amplification nucleic acid probe test for the in vitro detection of rRNA from Chlamydia trachomatis using the PANTHER System.
31050	CMV IgM using GSDX Automation	Gold Standard Diagnostics CMV IgM Ab (EIA) Test.
35030	CMV Total Ab	Immucor Capture-CMV solid phase red- cell adherence system for the detection of IgG and IgM Ab to CMV.
32110	Hepatitis B Core Total Ab	ORTHO® HBc ELISA Test System is an FDA approved qualitative assay for the detection of total antibody to hepatitis B virus core antigen (anti-HBc) in living human serum or plasma indicated for the screening of blood and blood products intended for transfusion and as an aid in the diagnosis of ongoing or previous hepatitis B virus infection.
32140	Hepatitis B Surface Ag	BioRad Genetic Systems HBsAg EIA 3.0 is FDA approved for living donor screening.
32210	Hepatitis C Virus Ab	ORTHO HCV Version 3.0 ELISA is FDA approved for living donor screening.
35210	HIV-1/HIV-2 Plus O	BioRad GS HIV-1/HIV-2 Plus O EIA is FDA approved for living donor screening. This is a screening assay; it is neither a diagnostic assay nor a confirmatory test for the presence of HIV.
35460	HTLV I/II Ab	The Avioq HTLV I/II Microelisa System is FDA approved for screening living donors and heart-beating organ donors.
3675	Neisseria gonorrhoeae - Panther Platform	The APTIMA COMBO 2 Assay is a target amplification nucleic acid probe test for the in vitro detection of rRNA from Neisseria gonorrhoeae using the PANTHER System.

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- | | | |
|------|--|---|
| 3991 | Syphilis Screening – Nontreponemal (Automated) | The ASI RPR Test for Syphilis, tested on the ASI Evolution automated analyzer, is a qualitative flocculation test for the detection of nontreponemal antibodies in human serum and plasma to aid in the diagnosis of syphilis. FDA-licensed test for cadaveric and living-donor screening. |
| 2773 | Ultrio Elite HBV | The Procleix Ultrio Elite Assay is FDA licensed for living and cadaveric donor screening by Nucleic Acid Test by Transcription Mediated Amplification for the detection of HBV. |
| 2772 | Ultrio Elite HCV | The Procleix Ultrio Elite Assay is FDA licensed for living and cadaveric donor screening by Nucleic Acid Test by Transcription Mediated Amplification for the detection of HCV. |
| 2771 | Ultrio Elite HIV-1/2 | The Procleix Ultrio Elite Assay is FDA licensed for living and cadaveric donor screening by Nucleic Acid Test by Transcription Mediated Amplification for the detection of HIV-1 and HIV-2. |
| 2722 | WNV | The Procleix WNV assay is a qualitative in vitro Nucleic Acid Test by Transcription Mediated Amplification for the detection of WNV RNA in plasma specimens. This is a screening assay and is not intended for use as an aid in the diagnosis of WNV. This assay is FDA licensed for both living and cadaveric donor screening. |

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