

Patient ID SA00056606	Patient Name SAMPLEREP, NKCP	Birth Date 1966-06-10	Gender F	Age 46
Order Number SA00056606	Client Order Number SA00056606	Ordering Physician Client, Client	Report Notes	
Account Information C7028846 DLMP Rochester		Collected 16 Apr 2013 08:07		

N.K. Cytotoxicity Profile

N.K. Cytotoxicity Result

% Cyto. E:T 100	MCR	% Cyto. E:T 12.5	MCR
52 %		19 %	
% Cyto. E:T 50	MCR	% Cyto. E:T 6.25	MCR
46 %		10 %	
% Cyto. E:T 25	MCR	% Cyto. E:T 3.13	MCR
30 %		5 %	

NKC Control Tube

Comment MCR

Normal NK cytotoxic function with normal NK cell absolute counts in whole blood.

ADDITIONAL INFORMATION

Lytic activity diminishes with time in heparinized blood specimens regardless of storage conditions. Specimens >24 hours old may yield falsely low results. Approximately 25% of healthy individuals present with abnormally decreased NK lytic activity. At least 3 consecutive results of reduced NK cytotoxic function over a period of approximately 3 months is required before a definitive assessment of impaired NK cytotoxicity can be attributed to a patient. Clinical correlation recommended.

T- and B-Cell QN by Flow Cytometry

CD45 Lymph Count, Flow	MCR	% CD19 (B Cells)	MCR
2.06 thou/mcL	Reference Value 0.82-2.84	16 %	Reference Value 6-24
% CD3 (T Cells)	MCR	% CD16+CD56 (NK cells)	MCR
69 %	Reference Value 58-86	14 %	Reference Value 4-28

Performing Site Legend

Code	Laboratory	Address
MCR	Mayo Clinic Dept. of Lab Med and Pathology	200 First Street SW, Rochester, MN 55905



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% CD4 (Helper Cells)

50 %

MCR
Reference Value
32-64

CD16+CD56 (NK cells)

293 cells/mcL

MCR
Reference Value
59-513

% CD8 (Supp'r Cells)

18 %

MCR
Reference Value
13-40

CD4 (Helper Cells)

1041 cells/mcL

MCR
Reference Value
365-1437

CD3 (T Cells)

1417 cells/mcL

MCR
Reference Value
550-2202

CD8 (Supp'r Cells)

364 cells/mcL

MCR
Reference Value
145-846

CD19 (B Cells)

336 cells/mcL

MCR
Reference Value
70-409

H/S Ratio

2.9

MCR
Reference Value
≥0.9

Received: 16 Apr 2013 08:14

Reported: 22 Apr 2013 16:10

Laboratory Notes

- 1 Analyte Specific Reagent: This test was developed and its performance characteristics determined by Mayo Clinic. It has not been cleared or approved by the U.S. Food and Drug Administration.

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Code	Laboratory	Address
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