

AUTHORIZATION FORM FOR RELEASE OF HEALTH INFORMATION

PATIENT NAI	BLAYK, BONZE 2 05/01/56		MAI	DEN OR OTHER NAME	
DATE OF BIR'I		95770	COR	D #:	
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DAY PHONE:					
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I specifically authorize Most recent an All visits since Pap Smear	nual exam			ED: DAT	ES
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TURPOSE OF RELI	EASE OF INFORMATIO			ty of care	
To ensure that our	records are current and	l accurate, please com	olete & retur	n this form promptly to	the clinic site checked
<i>below. This will end</i> □ Corning I35 Walnut St.	able us to provide our ci	lient with proper media ☐ Hornell 174 Main St. (607)324-112	cal care. The A 3 4 (ank you for your coope ☑ Ithaca 314 W. State St. 607)273-1513 607)273-8776 fax	
		CONDITIONS OF A	UTHORIZAT	TION	
. This Authorizati	on will expire on (inser	rt date or event):	1 0 1		
the date notified e	except to the extent that	PP of the Southern Fir	ger Lakes ha	s already acted upon su	g, and it will be effective on ach Authorization.
B. Information used	or disclosed pursuant to ral privacy regulations.	this Authorization mag	y be subject t	o re-disclosure by the r	ecipient and no longer
	is release of information	, my healthcare and pa	yment for my	y healthcare will not be	affected if I do not sign thi
	ed a copy of this signed.	Authorization form.			
SIGNATURE OF	ank og	1 (1014)	VT/LEGAL GU	ARDIAN/AUTHORIZED PE	ERSON DATE
	PATIENT DA) (
WITNESS	DA DA	9/21/12	AUTHORI	TY TO ACT ON BEHALF (OF PATIENT
WITNESS DATE REQUEST FILLED	0 11 12 C	ATE FOR OFFICE I		TY TO ACT ON BEHALF	OF PATIENT



14121 PP OF THE SOUTHERN FINGER LAKES, INC. ITHACA CENTER 314 WEST STATE ST. ITHACA, NY 14850

PATIENT INFORMATION

NAME: BL

BLAYK, BONZE ANNE ROSE

PATIENT ID#: 95770

ACCESSION #: 014121031432

DOB:

05/01/56

SPECIMEN INFORMATION

COLLECTED:

09/21/12 09/24/12

RECEIVED: REPORTED:

09/24/12

COLLECTED BY:

LEIA RAPHAELIDIS

PERFORMING LAB: C

CDD

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TEST PERFORMED

IN RANGE

OUT OF RANGE

REF. RANGE

Syphilis

Negative by EIA

NON-REACTIVE

Amplified CT - Urine

Negative

NEGATIVE

Amplified GC - Urine

Negative

NEGATIVE

Comment: Positive nucleic acid amplification test (NAAT) results are considered presumptive evidence of infection. Clinical correlation is necessary to determine if a positive test result is actually due to infection (true positive result), as this likelihood is decreased in low risk asymptomatic patients. A negative result does not preclude infection since results are dependent on specimen adequacy, absence of interfering substances, and a detectable number of organisms.

The analytical performance characteristics of this assay have been determined by CENTER FOR DISEASE DETECTION. METHODOLOGY: Strand Displacement Amplification (SDA).

This information has been disclosed to you from confidential records which are protected by law. Privacy laws prohibit you from making any further disclosure of this information without the specific written consent of the person to whom it pertains, or otherwise permitted by law. Any unauthorized further disclosure in violation of the law may result in a fine or jail sentence or both. A general authorization for the release of medical or other information is not, except in limited circumstances set forth in this part, sufficient authorization for further disclosure.



☐Corning 135 Walnut St. (607)962-4686

(607)962-7520 fax

□Elmira 755 E. Church St. (607)734-3313 (607)734-3392 fax □Hornell 111 Seneca St. (607)324-1124 (607)324-2666 fax □Ithaca 314 W. State St. (607)273-1513 (607)273-8776 fax □Watkins Glen 106 W. 4th St. (607)535-0030 (607)535-5040 fax

Rapid HIV Antibody Test Result

Patient Name: Bouze Anne Pose Blayk C	Collection Date: 9 121 1(2					
Counselor Initials: 56						
Patient's Date of Birth:/ Race:	W Gender:					
Name of Authorizing Physician : Joyce Leslie, MD						
The HIV-1/2 antibody result from the Rapid HIV Antibody Test is:						
Non-Reactive Reactive (Negative) (Preliminary Positive) □	L					
Type of Specimen: Rapid Test, oral fluid						
Meaning of the test result:						
A non-reactive (negative) test result means that no antibodies to HIV-1/2 have been detected. HIV antibodies may be absent during the "window period" of infection. Follow-up testing may be necessary if indicated by risk factors.						
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A **reactive** (**preliminary positive**) test result suggests that antibodies to HIV-1/2 may be present in the oral fluid or blood. A specimen will have to be sent to a laboratory for a Western Blot test prior to confirmation of the reactive test result. Precautions should be taken to avoid the chance of spreading HIV.

An **invalid** test result resulting from a rapid test can be due to a problem running the test or an interfering substance in the specimen. A specimen using other testing technology should be collected and submitted to a laboratory for HIV testing.

Questions: If you have any questions about the rapid test result, please contact the Authorizing Physician.

Confidentiality and Disclosure:

This information has been disclosed to you from confidential records which are protected by state law. State law prohibits you from making any further disclosure of this information without the specific written consent of the person to whom it pertains, or as otherwise permitted by law.