



Patient **Jane Doe**
 Accession ACN-17-04-00726

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CLIA ID: 05D2068632
 LAB ID: GENEX-LPC

Respiratory Panel Report **Abnormal**

PATIENT	Name	Jane Doe	ACCESSION	Accession	ACN-17-04-00726	PROVIDER	Name	Doctor Doctor
	DOB	Feb 12, 1970 Age(47)		Report Date	Apr 19, 2017 16:09		Phone	
	Gender	Female		Report Version	Final Report		Fax	
	MRN			Specimen Used				
	Diagnosis Codes			Specimen : Nasopharyngeal Swab				
	J06.9			Specimen ID: 595				
J21.9		Collected: Apr 01, 2017						
		Received: Apr 02, 2017						

Lab Results

Final Results **Positive for viral Pathogen, " Influenza A/H3."**

Panel	Test	Result	
●	Viruses	Adenovirus	Not Detected
●		Coronavirus 229E	Not Detected
●		Coronavirus HKU1	Not Detected
●		Coronavirus NL63	Not Detected
●		Coronavirus OC43	Not Detected
●		Human Metapneumovirus	Not Detected
●		Human Rhinovirus/Enterovirus	Not Detected
●		Influenza A	Not Detected
●		Influenza A/H1	Not Detected
●		Influenza A/H1-2009	Not Detected
●		Influenza A/H3	Detected
●		Influenza B	Not Detected
●		Parainfluenza 1	Not Detected
●		Parainfluenza 2	Not Detected
●		Parainfluenza 3	Not Detected
●		Parainfluenza 4	Not Detected
●		Respiratory Syncytial Virus	Not Detected
●	Bacteria	Bordetella pertussis	Not Detected
●		Chlamydomphila pneumoniae	Not Detected
●		Mycoplasma pneumoniae	Not Detected

Methodology The FilmArray Respiratory Pathogen Panel incorporates multiplex polymerase chain reaction (mPCR) with FilmArray system for the simultaneous qualitative detection and identification of multiple respiratory viral and bacterial nucleic acids. For each



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prepared using SimpleLims.



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sample, Nucleic acid purification occurs by using magnetic bead technology which is then combined with a preheated master mix to initiate the reverse transcription (RT) step. The effect of this stage of the PCR is to enrich for the target nucleic acids present in the sample which is then followed by targeting the specific nucleic acid sequences from each of the pathogens detected by using individual specific primers for different assays. After this stage the temperature is slowly increased and fluorescence in each well of the array is monitored and analyzed to generate a melting curve. The temperature at which a specific PCR product melts is consistent and predictable and the FilmArray software automatically evaluates the data from replicate wells for each assay to report results. The Data is then analyzed to provide a report summarizing which pathogens are present. Analyte targets (virus and bacteria nucleic acid sequences) may persist in vivo, independent of virus or bacteria viability. Detection of analyte target(s) does not guarantee that the corresponding live organism(s) is present, or that the corresponding organism(s) is the causative agent for clinical symptoms. This test has been validated at Genex Laboratory Professional Corporation according to guidelines.

Limitations

The results of this test should not be used as the sole basis for diagnosis, treatment, or other management decisions. Negative results in the setting of a respiratory illness may be due to infection with pathogens that are not detected by this test or, lower respiratory tract infection that is not detected by a nasopharyngeal swab specimen. Positive results do not rule out co-infection with other organisms: the agent(s) detected by the FilmArray RP may not be the definite cause of disease. Additional laboratory testing (e.g. bacterial and viral culture, immunofluorescence, and radiography) may be necessary when evaluating a patient with possible respiratory tract infection.

References

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Electronically Signed By:

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Molecular Genetic Pathologist

On Apr 19, 2017 16:09:34



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