

**SEVERE PULMONARY DISEASE ASSOCIATED
WITH E-CIGARETTE USE OUTBREAK**

NATIONAL CASE REPORT FORM (CDC)

October 31, 2019

Lung Injury Associated with E-cigarette Use or Vaping | National Case Report Form

CDC is investigating cases of unexplained lung injury associated with electronic cigarette use or vaping as detailed in CDC's Health Advisory (<https://emergency.cdc.gov/han/han00421.asp>). Local and state health departments should complete this form for any probable or confirmed case patient (see [case definition](#)) and transmit data to CDC using DCIPHER or by contacting CDC State Points of Contact.

Case ID Number _____ Medical Record Number _____
Case status Probable Confirmed Died? Yes No If yes, date of death _____ (see clinical section)
Was patient hospitalized? Yes No If yes, hospitalization date _____ Discharge date _____
Date reported to public health department _____ Name of Public Health Department _____
Person completing form _____ Contact phone number _____

PART I: PATIENT DEMOGRAPHICS AND EXPOSURES

Patient Demographics

County _____ State _____ Gender Male Female Other Age _____ years
Race White Black American Indian/Alaska Native Asian Native Hawaiian or Other Pacific Islander Other
Ethnicity Hispanic Non-Hispanic

Patient Substance Use in the Past 3 Months (90 days)

Any e-Cigarette use or vaping (e.g., vaping, dabbing)? Yes No Refused to answer
If yes, substance(s) vaped or dabbed in past 3 months?
 Nicotine Marijuana, THC oil, THC concentrates, hash oil, wax Cannabidiol (CBD) Synthetic Cannabinoids Flavors alone
 Other substances, specify _____ Unknown
Any combustible tobacco smoking (e.g., cigarettes, cigars)? Yes No Any other tobacco products (e.g., smokeless tobacco)? Yes No
Any combustible marijuana smoking (i.e., any non-vape marijuana)? Yes No Any other marijuana products (e.g., edibles)? Yes No
Any nicotine e-cigarette use or vaping reported? Yes No Unknown Date last used _____
If yes, what is the frequency of use? Daily A few times per week, specify: _____ A few times per month, specify _____
 Monthly or less [Skip logic: On average, how many times per day? _____]
Did patient report vaping flavoured nicotine in e-Cigarette and/or vape product(s)? Yes No
How many brands of nicotine containing products vaped or dabbed in the past 3 months? _____ [enter whole number]
Where was the nicotine e-Cigarette(s) or vaping product(s) purchased or obtained? Check all that apply
 Recreational dispensary Vape or smoke shop Pop-up shop Grocery store/drugstore/Convenience store Family or friend
 Dealer Online Other, describe _____
What kind of device(s) were used with this substance? Select all that apply
 Disposable e-cigarette or vaping device E-cigarettes with pre-filled or refillable cartridges (e.g., using battery pens, Ego, EVO, Ooze pen, Caliplug, 510 battery) E-cigarette with tank that you refill with liquids (including sub-ohm, mod or modifiable systems)
 E-cigarettes with pre-filled or refillable "pods" or pod cartridges (e.g. JUUL, Suorin) Other, describe _____
Was this a mod device (a device that allows user to choose higher and/or variable temperatures)? Yes No Unknown
Did patient modify, or add a substance to, the device(s) that was not intended by the manufacturer? Yes No Unknown
If yes, explain _____
Does patient know anyone else who became ill from vaping nicotine? Yes No
If yes, were nicotine products or devices shared with that person? Yes No
Product sample sent for testing? Yes No If yes, where was sample tested _____ Product sample ID number(s) _____
Any THC e-cigarette use or vaping reported? Yes No Unknown Date last used _____
If yes, what is the frequency of use? Daily A few times per week, specify: _____ A few times per month, specify _____
 Monthly or less [Skip logic: On average, how many times per day? _____]
Did patient report vaping flavoured THC in e-Cigarette and/or vape product(s)? Yes No
How many brands of THC containing products vaped or dabbed in the past 3 months? _____ [enter whole number]
What was the purpose of THC product(s) use? medical purposes nonmedical (recreational) purposes other, specify _____
Which THC substance(s) were used in an e-cigarette, vaping device, vaporizer, or dab rig? Select all that apply
 Marijuana herb THC oils Butane hash oil THC concentrate (e.g., wax, batter/budder, crumble, shatter, pull and snap)
 THC powder (e.g., dry sift) Other, describe _____
Where was the THC e-Cigarette(s) or vaping product(s) purchased or obtained? Check all that apply
 Medical dispensary Recreational dispensary (retail cannabis/marijuana shop) Vape or smoke shop Pop-up shop
 Grocery store/Drugstore/Convenience store Family or friend Illicit dealer Online Other, describe _____
What kind of device(s) were used with this substance? Select all that apply
 Disposable device Device with pre-filled cartridges Device with tank that you refill with liquids (e.g., mods)
 Device with pre-filled or refillable "pods" or pod cartridges (e.g. JUUL, Suorin) Dab rig Vaporizer (for dry herbs, etc.) Other _____
What brand of THC cartridge(s) were used with device(s): Rove Dank Vapes Golden Gorilla Smart Cart Other _____
Was this a mod device (a device that allows user to choose higher and/or variable temperatures)? Yes No Unknown
Did patient modify, or add a substance to, the device(s) that was not intended by the manufacturer? Yes No Unknown
If yes, explain _____
Does patient know anyone else who became ill from vaping THC? Yes No
If yes, were THC products or devices shared with that person? Yes No
Product sample sent for testing? Yes No If yes, where was sample tested _____ Product sample ID number(s) _____

PART II: CLINICAL INFORMATION

Symptoms at Initial Presentation to Medical Care

Chief complaint _____ Date symptom(s) started _____

GI symptoms? Yes No Unknown If yes, describe _____

Respiratory symptoms? Yes No Unknown If yes, describe _____

Constitutional symptoms? Yes No Unknown If yes, describe _____
(e.g., fever, chills, malaise)

Weight loss during current illness? Yes No Unknown If yes, amount (lb) _____

Medical History

Chronic respiratory disease (including asthma, COPD, etc.)? Yes No If yes, specify type of disease _____

Heart disease? Yes No If yes, specify type of disease _____

Anxiety? Yes No

Depression? Yes No

Other chronic illness? Yes No If yes, specify type of chronic illness _____

Pregnant? Yes No Unknown If yes, trimester First Second Third Unknown

Imaging

CT performed Yes No If yes, location of abnormal findings Bilateral Right Left Normal (no findings)
If yes, infiltrates/opacities present Yes No Subpleural sparing Yes No Unknown

Chest X-ray performed Yes No If yes, location of abnormal findings Bilateral Right Left Normal (no findings)
If yes, infiltrates/opacities present Yes No

Specify other abnormal chest imaging findings (e.g., pneumothorax) _____

Infectious Disease Testing

Respiratory viral panel	<input type="checkbox"/> Positive (specify _____)	<input type="checkbox"/> Negative	<input type="checkbox"/> Pending	<input type="checkbox"/> Not done
Influenza	<input type="checkbox"/> Positive (specify _____)	<input type="checkbox"/> Negative	<input type="checkbox"/> Pending	<input type="checkbox"/> Not done
Blood cultures	<input type="checkbox"/> Positive (specify organisms _____)	<input type="checkbox"/> Negative	<input type="checkbox"/> Pending	<input type="checkbox"/> Not done
Legionella urinary antigen	<input type="checkbox"/> Positive	<input type="checkbox"/> Negative	<input type="checkbox"/> Pending	<input type="checkbox"/> Not done
Strep pneumoniae urinary antigen	<input type="checkbox"/> Positive	<input type="checkbox"/> Negative	<input type="checkbox"/> Pending	<input type="checkbox"/> Not done
Mycoplasma pneumoniae	<input type="checkbox"/> Positive (specify _____)	<input type="checkbox"/> Negative	<input type="checkbox"/> Pending	<input type="checkbox"/> Not done
Other	<input type="checkbox"/> Positive (specify _____)	<input type="checkbox"/> Negative	<input type="checkbox"/> Pending	<input type="checkbox"/> Not done

Clinical Course of Lung Injury

Is this the first time patient is presenting for clinical care for these symptoms? Yes No If yes, is a follow-up visit scheduled? Yes No

Was patient hypoxemic (<95) at any outpatient visit or hospitalization? Yes No If yes, date(s) _____ Lowest value: _____

Outpatient visit #1 Yes No If yes, date of visit _____ Outpatient visit #2 Yes No If yes, date of visit _____
Were there additional outpatient/clinic visits? Yes No If yes, specify number of additional visits _____

Urgent care visit #1 Yes No If yes, date of visit _____ Urgent care visit #2 Yes No If yes, date of visit _____
Were there additional urgent care visits? Yes No If yes, specify number of additional visits _____

Emergency Department (ED) visit #1 Yes No If yes, date of visit _____ ED visit #2 Yes No If yes, date of visit _____
Were there additional ED visits? Yes No If yes, specify number of additional visits _____

If hospitalized, was patient re-hospitalized at a later date? Yes No If yes, hospitalization date _____ Discharge date _____
Were there additional hospitalizations? Yes No If yes, specify number of additional hospitalizations _____

ICU Admission Yes No If yes, ICU admission date _____ ICU duration (in days) _____

Treated with steroids? Yes No If yes, medication(s): _____ dose: _____ start date: _____ duration: _____ Taper

Treated with antibiotics? Yes No If yes, medication(s): _____ dose: _____ start date: _____ duration: _____

Treated with antivirals? Yes No If yes, medication(s): _____ dose: _____ start date: _____ duration: _____

Required respiratory support? Yes No Intubated (duration _____) BiPAP/CPAP/High flow Supplemental oxygen

Required ECMO (Extracorporeal membrane oxygenation)? Yes (duration _____) No

Clinical specimens

Bronchoalveolar lavage performed? Yes, date of sample _____ No If yes, where tested _____ Specimen ID _____
If yes, lipid staining Yes No
If yes, lipid-laden macrophages seen Yes No

Blood sample testing performed? Yes, date of sample _____ No If yes, where tested _____ Specimen ID _____

Urine sample testing performed? Yes, date of sample _____ No If yes, where tested _____ Specimen ID _____

Lung biopsy performed? Yes, date of sample _____ No If yes, where tested _____ Specimen ID _____
If yes, lipid staining? Yes No
If yes, lipid-laden macrophages seen? Yes No
If yes, findings consistent with acute lung injury? Yes No If no, specify findings _____
If yes, other significant findings _____

Death Information

Died Yes No If yes, specify location _____ Date of death _____

Immediate cause of death _____ Contributing causes of death _____

Autopsy performed? Yes No If yes, autopsy sample collected Yes No If yes, where tested _____ Specimen ID _____
If yes, lipid staining performed on autopsy lung tissue? Yes No If yes, lipid-laden macrophages seen? Yes No
If yes, findings consistent with acute lung injury? Yes No If no, specify findings _____
If yes, other significant autopsy findings _____

Specimen Manifest Form

CDC Case ID (note: this is used by state epidemiologists when submitting case data to CDC)	State Case ID	Sample ID	Matrix (urine, whole blood, serum, plasma, BAL, etc)	Box # or ID	Position in Box	Volume (mL)	Collection Date	Any pertinent comments (hemolyzed sample, clotted sample, etc)

CDC estimates the average public reporting burden for this collection of information as 10 minutes per response, including the time for reviewing instructions, searching existing data/information sources, gathering and maintaining the data/information needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Information Collection Review Office, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30333; ATTN: PRA (0920-1011).