

Routes of Transmission and principles of transmission interruption

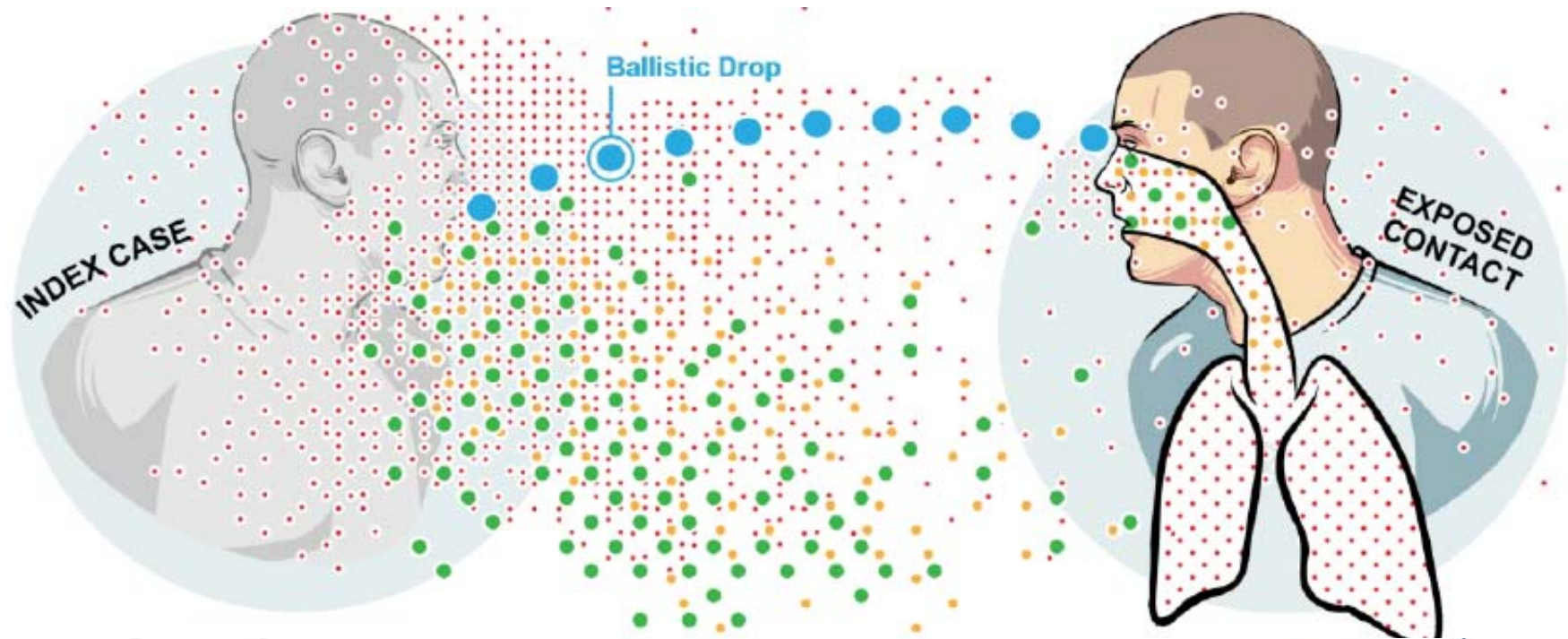
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Transmission of respiratory viruses:

3 routes of transmission

- Contact with secretions loaded with viruses (direct or indirect: fomites), followed by self inoculation
- Large droplets that fall quickly on the ground
- Aerosol sized- droplets that linger in the air and can follow air flow



Aerosols



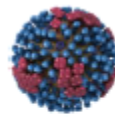
Respirable Aerosol
 ≤ 2.5 to $5\mu\text{m}$



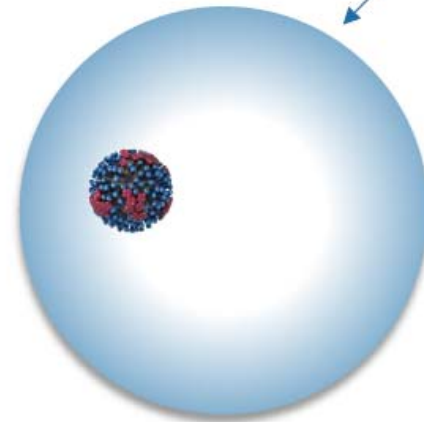
Thoracic Aerosol
 ≤ 10 to $15\mu\text{m}$



Inhalable Droplets
 $\leq 100\mu\text{m}$

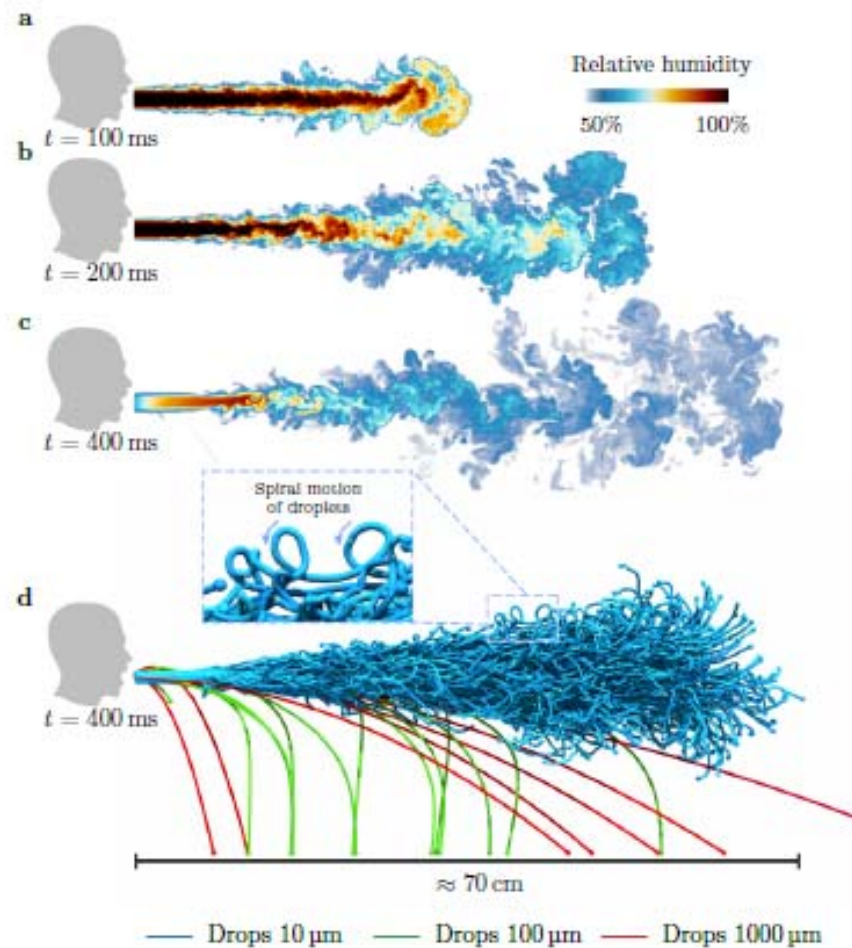


$0.1\ \mu\text{m}$



$0.5\ \mu\text{m}$
 $(0.2-100\ \mu\text{m})$

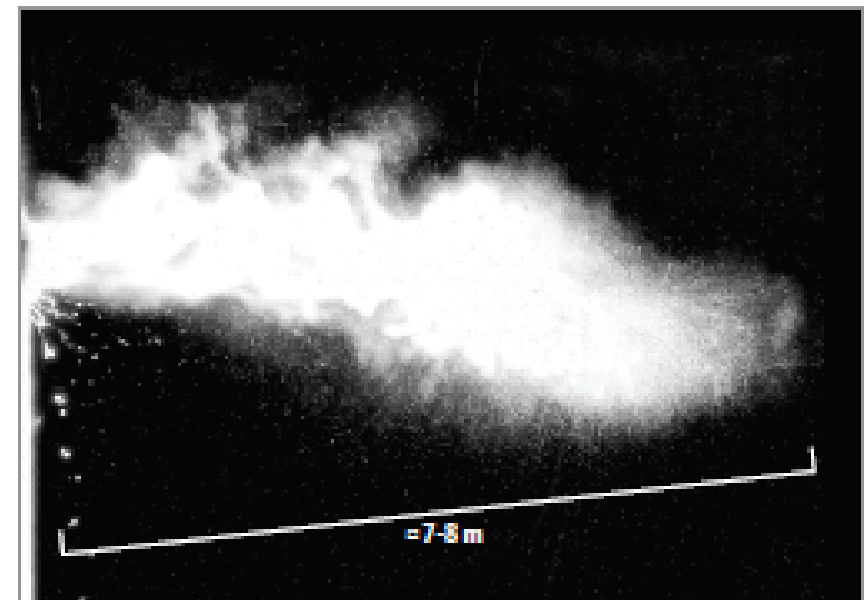
Aerodynamic diameter of particle	Settling time in still air (3 m fall)
100 μm	10 sec
40 μm	1 min
20 μm	4 min
10 μm	17 min
5 μm	67 min
3 μm	3.1 hours
1 μm	27.8 hours



Cheong KL et al Extended lifetime of respiratory droplets in a turbulent vapour puff and its implication on airborne disease transmission Preprint August 2020

Horizontal distance traveled on an horizontal jet emitted at height of 2m; RH 50% (NaCl 0.9%)
Xie et al Indoor Air 2007; 17: 211-225

Figure. Multiphase Turbulent Gas Cloud From a Human Sneeze



JAMA Insights

Turbulent Gas Clouds and Respiratory Pathogen Emissions
Potential Imp... Lydia Bourouiba, PhD Reducing Transmission of COVID-19

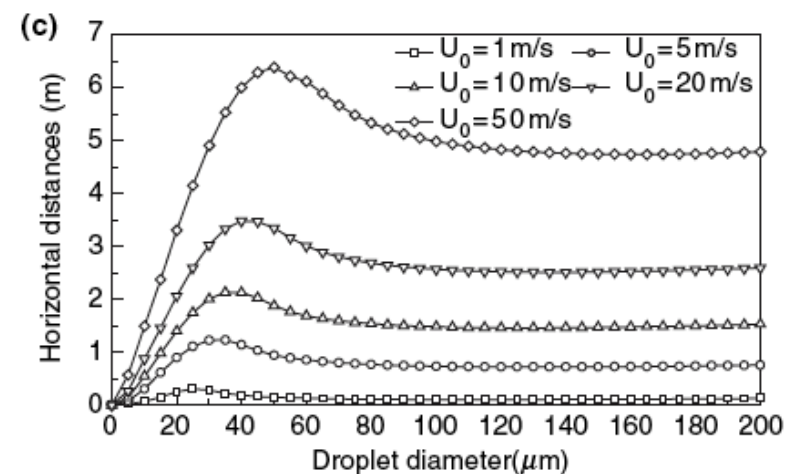


TABLE I. Aerosol Administration of Influenza A2/Bethesda/10/63 to Volunteers.

Inhaled virus (TCID ₅₀)	Vol #	Illness	Virus recovery (days after inoc)	Neutralization antibody	
				Before inoculation	28 days after
126	1	*	*	1280	2560
	2			2560	2560
	3			640	1280
78	4			160	160
	5			320	320
	6			320	320
59	7			40	1280
	8			80	80
	9			80	80
1	10	+	3-7	<5	80
2	11			<5	<5
	12			<5	<5
	13			<5	<5
5	14		4-7	<5	320
	15			40	40
	16			80	80
	17			40	40
	18			10	1280
	19			20	1280
	20			5	5120
	21	+	2-6	<5	<5
	22			<5	640
	23			<5	<5

* Blank spaces indicate no response.

Comparison of human infectious dose of influenza virus by aerosol or intranasal route

- Aerosol:
 $\text{HID}_{50} = 0.6 \text{ to } 3 \text{ TCID}_{50}$
- Intranasal (large droplet)
 $\text{HID}_{50} = 127 \text{ to } 320 \text{ TCID}_{50}$

(Douglas R.G. Influenza in Man. Pp 375-447 in
The Influenza Viruses and Influenza, Kilbourne E.D. ed,
Academic Press, New York 1975.)

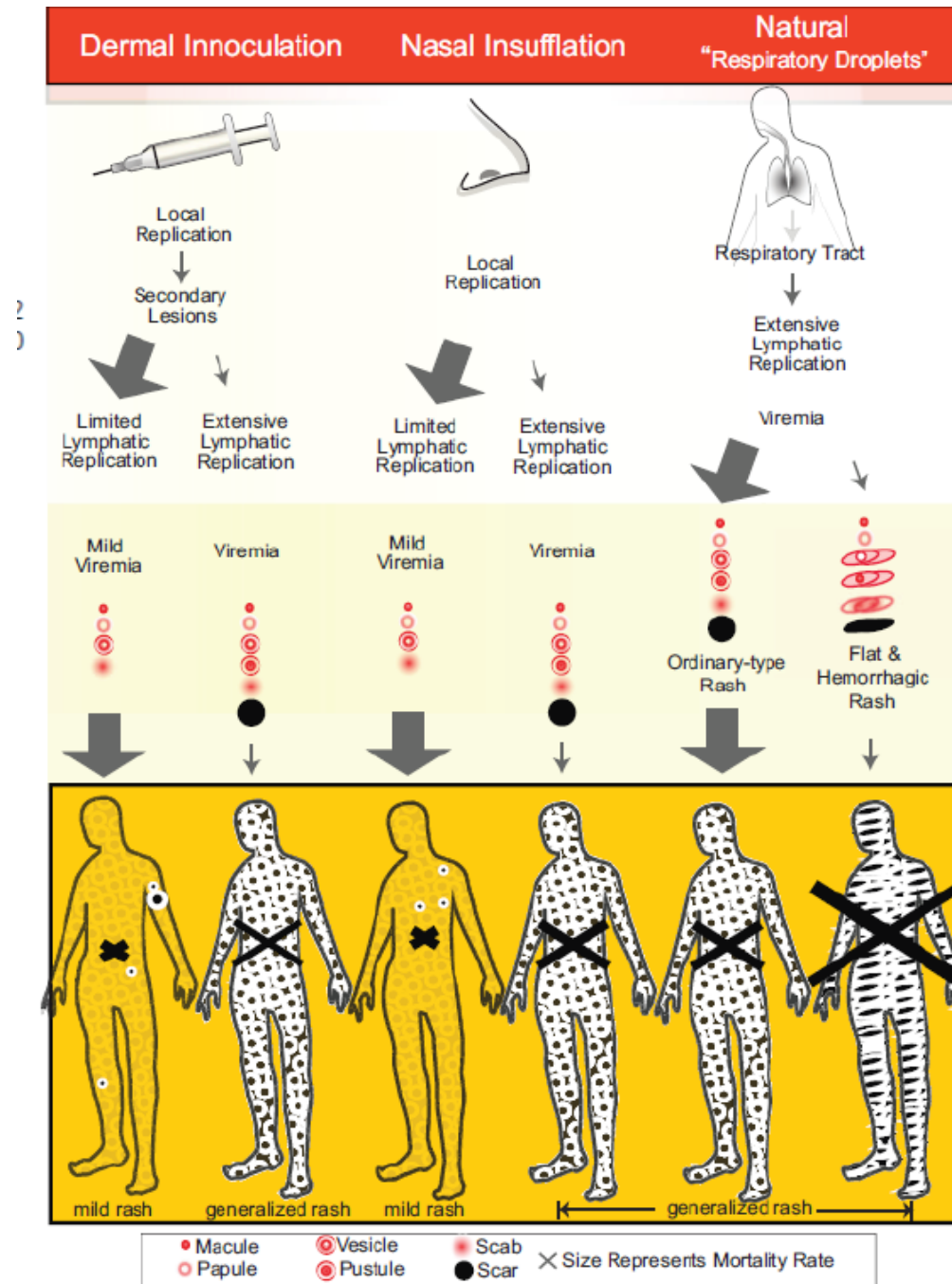
Diseases caused by experimental influenza infections in humans

- by the aerosol route: disease is very similar to that seen in natural infections.
- by intranasal drops (large droplets): disease is milder, with a longer incubation period and usually no involvement of the lower respiratory tract.

1) Douglas R.G. Influenza in Man. Pp 375-447 in *The Influenza Viruses and Influenza*, Kilbourne E.D. ed, Academic Press, New York 1975

2) Little J.W. et al *J Med Virol* 3: 177-188, 1979.

3) Knight V.pp. 175-182 in: Hers JF, Winkles KC, eds. *Airborne Transmission and Airborne Infections VIth International Symposium on Aerobiology*. New York: Wiley; 1973



Milton DK Cellular
and Infection Microbiology
published: 29 November 2012
doi: 10.3389/fcimb.2012.00150

ORIGINAL ARTICLE

Evidence of Airborne Transmission of the Severe Acute Respiratory Syndrome Virus

Ignatius T.S. Yu, M.B., B.S., M.P.H., Yuguo Li, Ph.D., Tze Wai Wong, M.B., B.S.,
Wilson Tam, M.Phil., Andy T. Chan, Ph.D., Joseph H.W. Lee, Ph.D.,
Dennis Y.C. Leung, Ph.D., and Tommy Ho, B.Sc.

N Engl J Med 2004;350:1731-9.

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Clin Infect Dis. 2016 Aug 1; 63(3): 363–369.

PMCID: PMC7108054

Published online 2016 Apr 18. doi: 10.1093/cid/ciw239: 10.1093/cid/ciw239

PMID: [27090992](#)

Extensive Viable Middle East Respiratory Syndrome (MERS) Coronavirus Contamination in Air and Surrounding Environment in MERS Isolation Wards

[Sung-Han Kim](#),  [So Young Chang](#), [Minki Sung](#), [Ji Hoon Park](#), [Hong Bin Kim](#), [Heeyoung Lee](#), [Jae-Phil Choi](#), [Won Suk Choi](#), and [Ji-Young Min](#)

OPEN

Aerosol and surface contamination of SARS-CoV-2 observed in quarantine and isolation care

Joshua L. Santarpia^{1,2,✉}, Danielle N. Rivera², Vicki L. Herrera¹, M. Jane Morwitzer¹, Hannah M. Creager², George W. Santarpia², Kevin K. Crown², David M. Brett-Major¹, Elizabeth R. Schnaubelt^{1,3}, M. Jana Broadhurst¹, James V. Lawler^{1,2}, St. Patrick Reid¹ & ...

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ARTICLE

<https://doi.org/10.1038/s41467-020-16670-2>

OPEN

Detection of air and surface contamination by SARS-CoV-2 in hospital rooms of infected patients

Po Ying Chia^{1,2,3,11}, Kristen Kelli Coleman^{4,11}, Yian Kim Tan^{5,11}, Sean Wei Xiang Ong^{1,2,11}, Marcus Gum⁵, Sok Kiang Lau⁵, Xiao Fang Lim⁵, Ai Sim Lim⁵, Stephanie Sutjipto^{1,2}, Pei Hua Lee^{1,2}, Than The Son⁴, Barnaby Edward Young^{1,2,3}, Donald K. Milton⁶, Gregory C. Gray^{4,7,8}, Stephan Schuster⁹, Timothy Barkham^{2,10}, Partha Pratim De^{2,3}, Shawn Vasoo^{1,2,3}, Monica Chan^{1,2}, Brenda Sze Peng Ang^{1,2,3,10}, Boon Huan Tan⁵, Yee-Sin Leo^{1,2,3,10}, Oon-Tek Ng^{1,2,3,12,✉}, Michelle Su Yen Wong^{5,12}, Kalisvar Marimuthu^{1,2,10,12,✉} & for the Singapore 2019 Novel Coronavirus Outbreak Research Team*

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nature

<https://doi.org/10.1038/s41586-020-2271-3>

Accelerated Article Preview

Aerodynamic analysis of SARS-CoV-2 in two Wuhan hospitals

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Accelerated Article Preview Published online 27 April 2020

Yuan Liu, Zhi Ning, Yu Chen, Ming Guo, Yingliu, Nirmal Kumar Gali, Li Sun, Yusen Duan, Jing Cai, Dane Westerdahl, Xinjin Liu, Ke Xu, Kin-fai Ho, Haidong Kan, Qingyan Fu & Ke Lan

cdc Centers for Disease Control and Prevention

Volume 26, Number 7—July 2020

Dispatch

Aerosol and Surface Distribution of Severe Acute Respiratory Syndrome Coronavirus 2 in Hospital Wards, Wuhan, China, 2020

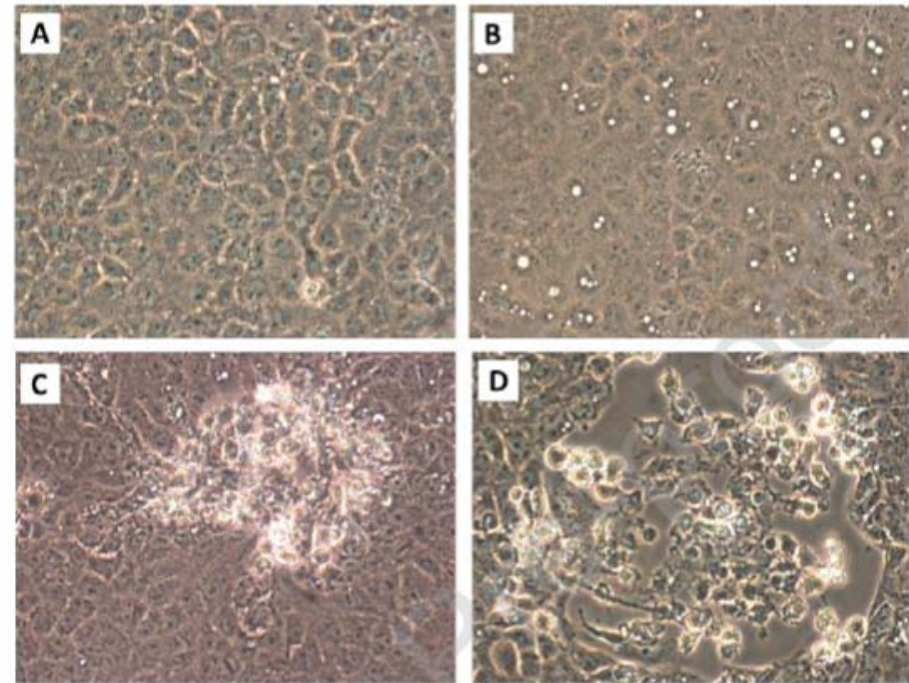
Zhen-Dong Guo¹, Zhong-Yi Wang¹, Shou-Feng Zhang¹, Xiao Li, Lin Li, Chao Li, Yan Cui, Rui-Bin Fu, Yun-Zhu Dong, Xiang-Yang Chi, Meng-Yao Zhang, Kun Liu, Cheng Cao, Bin Liu, Ke Zhang, Yu-Wei Gao✉, Bing Lu✉, and Wei Chen✉

Viable SARS-CoV-2 in the air of a hospital room with COVID-19 patients

John A. Lednicky, Michael Lauzardo, Z. Hugh Fan, Antarpreet Jutla, Trevor B. Tilly, Mayank Gangwar, Moiz Usmani, Sripriya Nannu Shankar, Karim Mohamed, Arantza Eiguren-Fernandez, Caroline J. Stephenson, Md. Mahbubul Alam, Maha A. Elbadry, Julia C. Loeb, Kuttinchantran Subramaniam, Thomas B. Waltzek, Kartikeya Cherabuddi, J. Glenn Morris Jr., Chang-Yu Wu

PII: S1201-9712(20)30739-6

DOI: <https://doi.org/10.1016/j.ijid.2020.09.025>



medRxiv preprint doi: <https://doi.org/10.1101/2020.07.13.20041632>; this version posted July 21, 2020. The copyright holder for this preprint (which was not certified by peer review) is the author/funder, who has granted medRxiv a license to display the preprint in perpetuity. It is made available under a CC-BY-NC-ND 4.0 International license.

The Infectious Nature of Patient-Generated SARS-CoV-2 Aerosol

Joshua L. Santarpia^{1,2*}, Vicki L. Herrera¹, Danielle N. Rivera², Shanna Ratnesar-Shumate¹, St. Patrick Reid^{1*}, Paul W. Denton³, Jacob W.S. Martens³, Ying Fang⁴, Nicholas Conoan¹, Michael V. Callahan⁵, James V. Lawler¹, David M. Brett-Major¹ and John J. Lowe^{1*}

1. University of Nebraska Medical Center

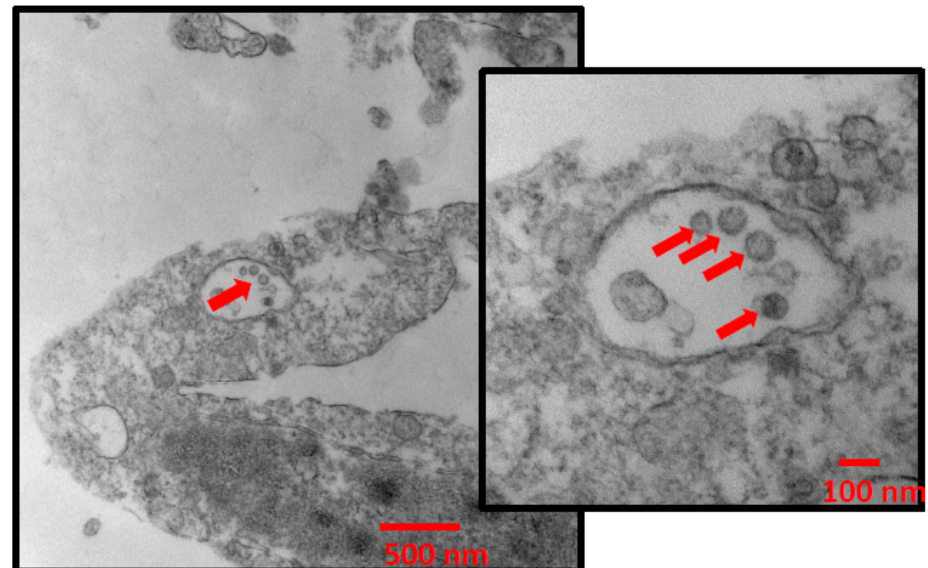


Figure 3. Electron micrographs of SARS-CoV-2 virions cultivated from the sub-micron filter from Room 5C.

Isolation of SARS-CoV-2 from the air in a car driven by a COVID patient with mild illness

John A. Lednicky,^{1,2} Michael Lauzardo,^{1,3} Md. M. Alam,^{1,2} Maha A. Elbadry,^{1,2} Caroline J.

Stephenson,^{1,2} Julia C. Gibson,^{1,2} and J. Glenn Morris, Jr.^{1,3*}

ABSTRACT

We used a Sioutas personal cascade impactor sampler (PCIS) to screen for SARS-CoV-2 in a car driven by a COVID-19 patient. SARS-CoV-2 was detectable at all PCIS stages by PCR and was cultured from the section of the sampler collecting particles in the 0.25 to 0.50 μm size range.

Title: Development of a Coronavirus Disease 2019 Nonhuman Primate Model Using Airborne Exposure

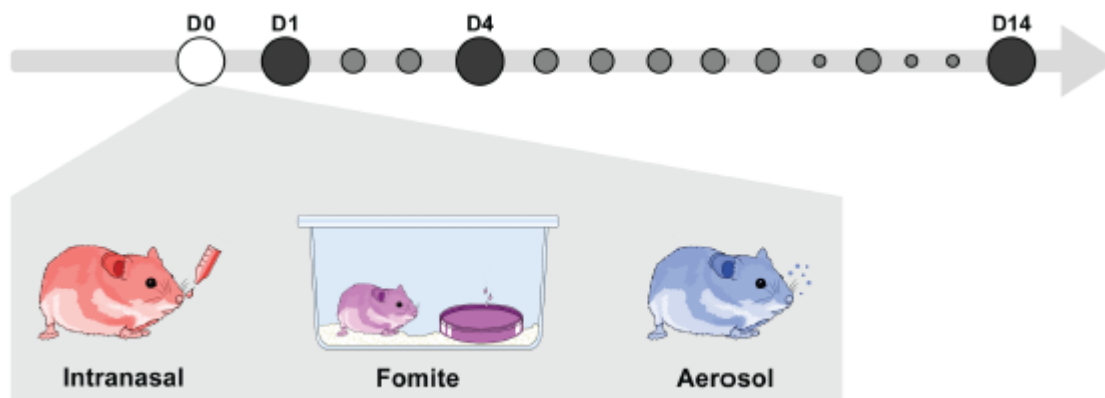
Authors: Sara C. Johnston^{1*}, Alexandra Jay², Jo Lynne Raymond³, Franco Rossi⁴, Xiankun Zeng³, Jennifer Scruggs³, David Dyer⁴, Ondraya Frick⁴, Joshua Moore², Kerry Berrier⁴, Heather Esham⁴, Joshua Shamblin², Willie Sifford², Jimmy Fiallos², Leslie Klosterman², Stephen Stevens², Lauren White², Philip Bowling², Terrence Garcia⁴, Christopher Jensen⁴, Jeanean Ghering⁴, David Nyakiti⁴, Stephanie Bellanca⁴, Brian Kearney⁵, Wendy Giles², Nazira Alli⁴, Fabian Paz², Kristen Akers⁵, Denise Danner⁵, James Barth⁵, Joshua A. Johnson⁵, Matthew Durant⁵, Ruth Kim⁵, Margaret LM Pitt⁷, Aysegul Nalca^{6*}

CMs have been investigated as a potential model for SARS-CoV-2, with variable disease findings noted following either IT/IN or IT/IN/ocular inoculation (11, 22). Importantly, fever, which represents a prominent human disease sign and which was consistently noted for CM on the present study, was not noted for any CMs on these prior studies.

(1 -3 μm)

SARS-CoV-2 disease severity and transmission efficiency is increased for airborne but not fomite exposure in Syrian hamsters.

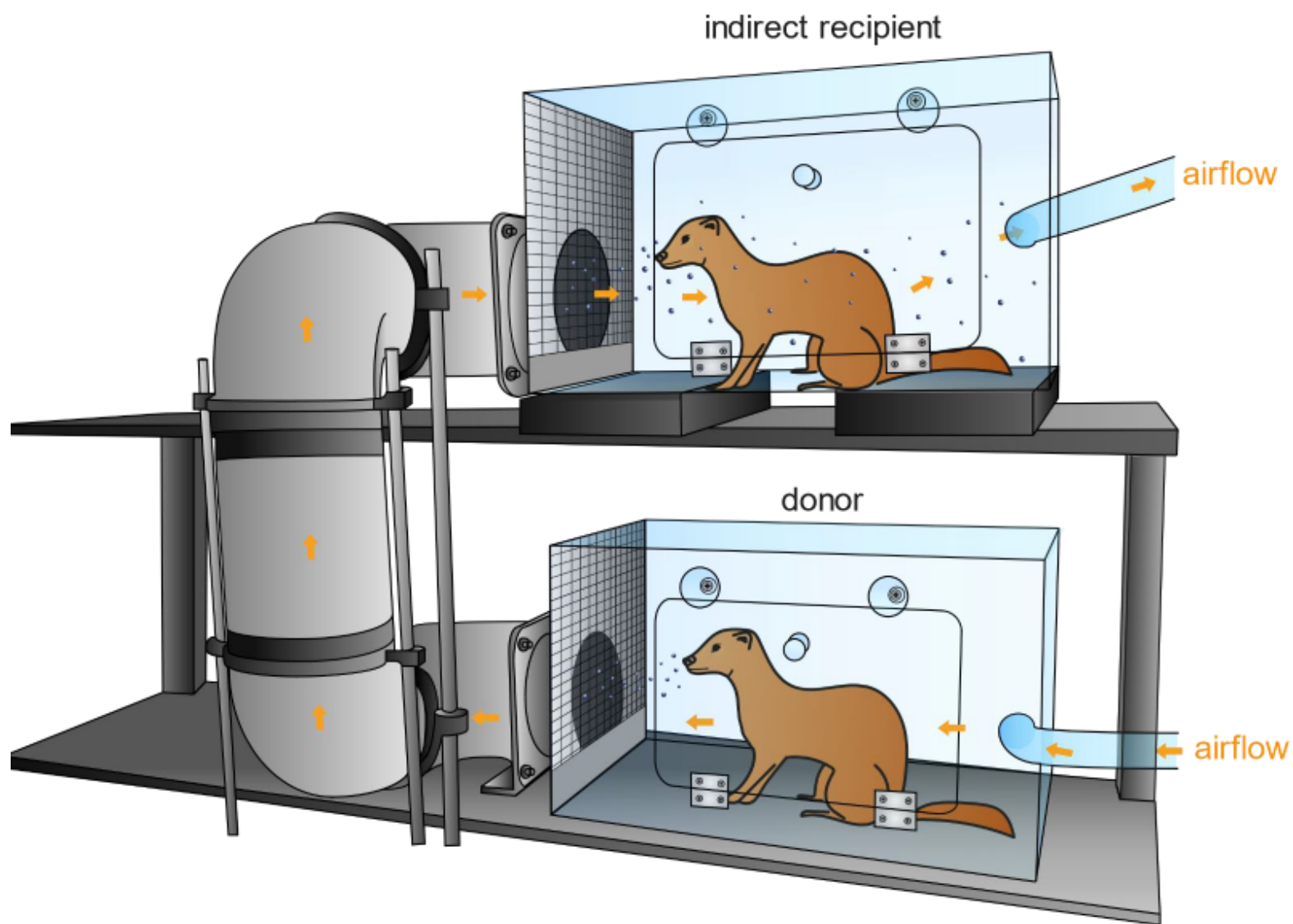
Julia R. Port^{1*}, Claude Kwe Yinda^{1*}, Irene Offei Owusu¹, Myndi Holbrook¹, Robert Fischer¹, Trenton Bushmaker^{1,2}, Victoria A. Avanzato¹, Jonathan E. Schulz¹, Neeltje van Doremalen¹, Chad S. Clancy³, Vincent J. Munster^{1#}

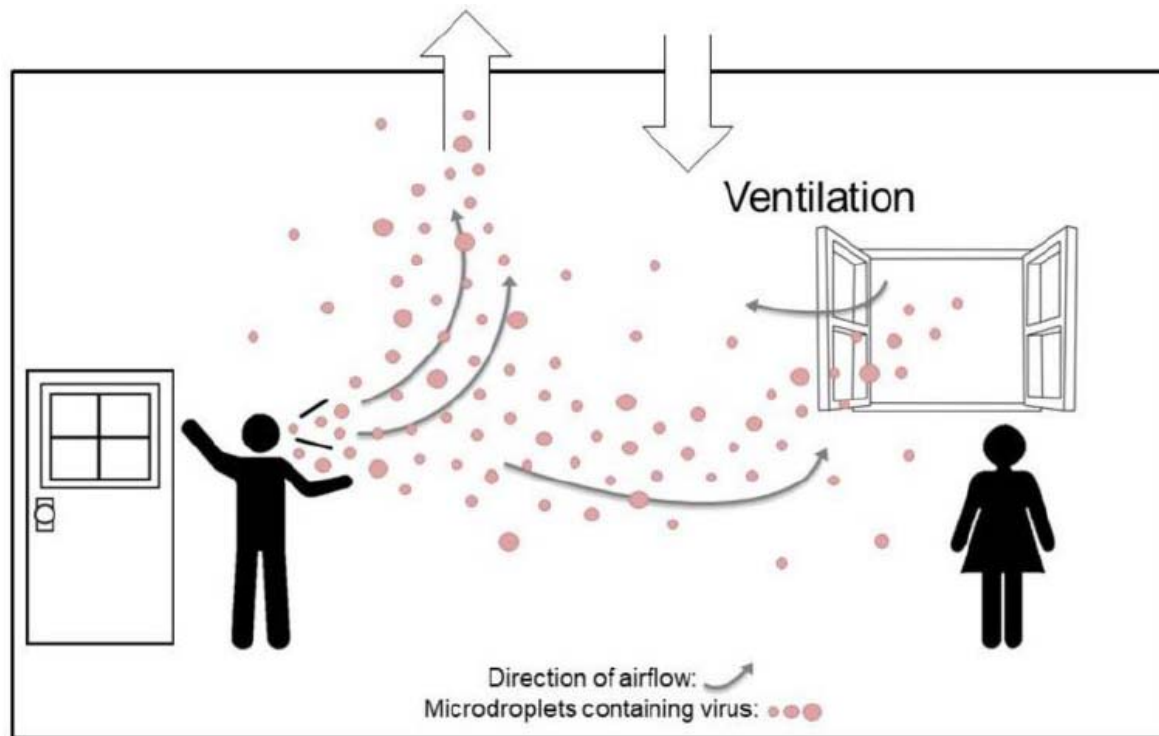


Aerosols of 1 -5 μm

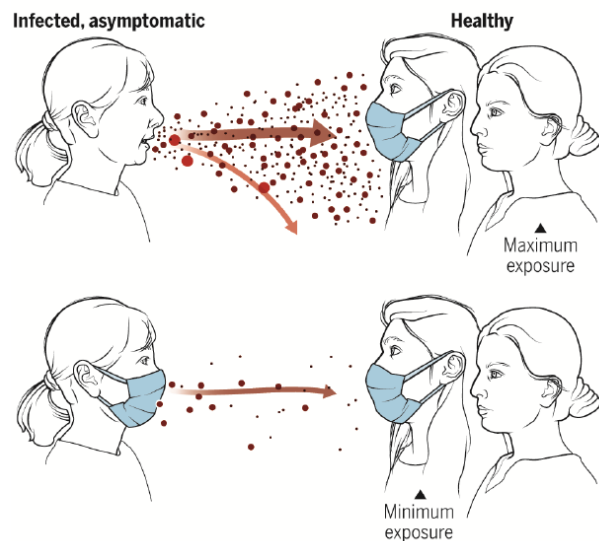
“Intranasal and aerosol inoculation caused more severe respiratory pathology, higher virus loads and increased weight loss. Fomite exposure led to milder disease manifestation characterized by an anti-inflammatory immune state and delayed shedding pattern.”

Figures





Morawska L
Milton DK
CID July 6 2020



Prather KA et al Science May 27 2020

3M

<https://blogs.cdc.gov/niosh-science-blog/2017/07/06/elastomerics/>