

ACIP COVID-19 Vaccines

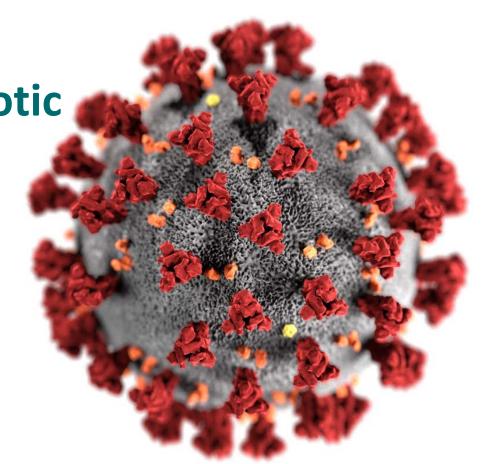
Risk/Benefit assessment of thrombotic thrombocytopenic events after Janssen COVID-19 vaccines:

Applying Evidence to Recommendation Framework





Sara Oliver MD, MSPH ACIP Meeting April 23, 2021



Evidence to Recommendations Framework



Evidence to Recommendations (EtR) Framework

 Structure to describe information considered in moving from evidence to ACIP vaccine recommendations

 Provide transparency around the impact of additional factors on deliberations when considering a recommendation

Evidence to Recommendations (EtR) FrameworkPolicy Question

Should vaccination with the Janssen COVID-19 vaccine (1 dose) be recommended for persons 18 years of age and older under an Emergency Use Authorization?

Evidence to Recommendations (EtR) Framework:

Previous Janssen COVID-19 vaccine Recommendations

EtR Domain	Question			
Public Health Problem	Is the problem of public health importance?			
Benefits and Harms	 How substantial are the desirable anticipated effects? How substantial are the undesirable anticipated effects? Do the desirable effects outweigh the undesirable effects? 			
Values	 Does the target population feel the desirable effects are large relative to the undesirable effects? Is there important variability in how patients value the outcomes? 			
Acceptability	Is the intervention acceptable to key stakeholders?			
Feasibility	Is the intervention feasible to implement?			
Resource Use	Is the intervention a reasonable and efficient allocation of resources?			
Equity	What would be the impact of the intervention on health equity?			

Evidence to Recommendations (EtR) Framework:

Adaptation to Risk/Benefit assessment for Janssen COVID-19 vaccine recommendations

EtR Domain	Question
Public Health Problem	 Recent COVID-19 Epidemiology Thrombosis after COVID-19 Disease Cerebral Venous Sinus Thrombosis (CVST) Heparin Induced Thrombocytopenia (HIT) AstraZeneca COVID-19 vaccines: Available global data
Benefits and Harms	 Benefits of Janssen COVID-19 vaccine Harms of Janssen COVID-19 vaccine: Estimated cases of TTS after Janssen COVID-19 vaccine Benefit/Risk Assessment of COVID-19 vaccines

Evidence to Recommendations (EtR) Framework:

Adaptation to Risk/Benefit assessment for Janssen COVID-19 vaccine recommendations

EtR Domain	Question
Values and Acceptability	 Intent to receive 1-dose COVID-19 vaccine Intent to receive Janssen COVID-19 vaccine over time
Feasibility	 Jurisdictional use of Janssen COVID-19 vaccine Possible impact of Janssen COVID-19 vaccine policy options
Equity	 Possible impact of Janssen COVID-19 vaccine policy options in disproportionately affected populations
Resource Use	No information available

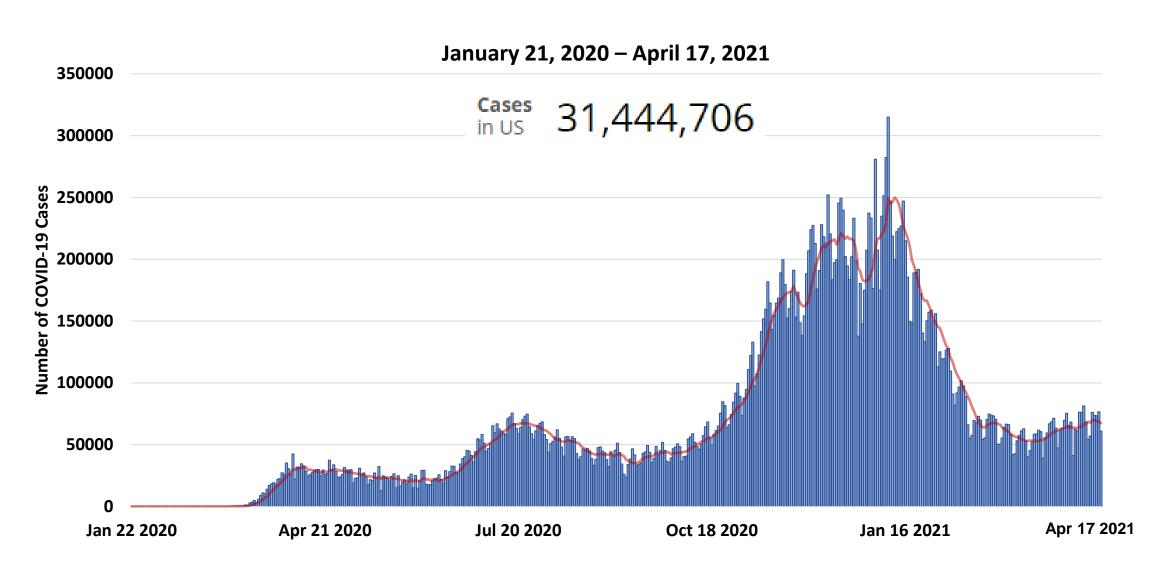
EtR Domain: Public Health Problem



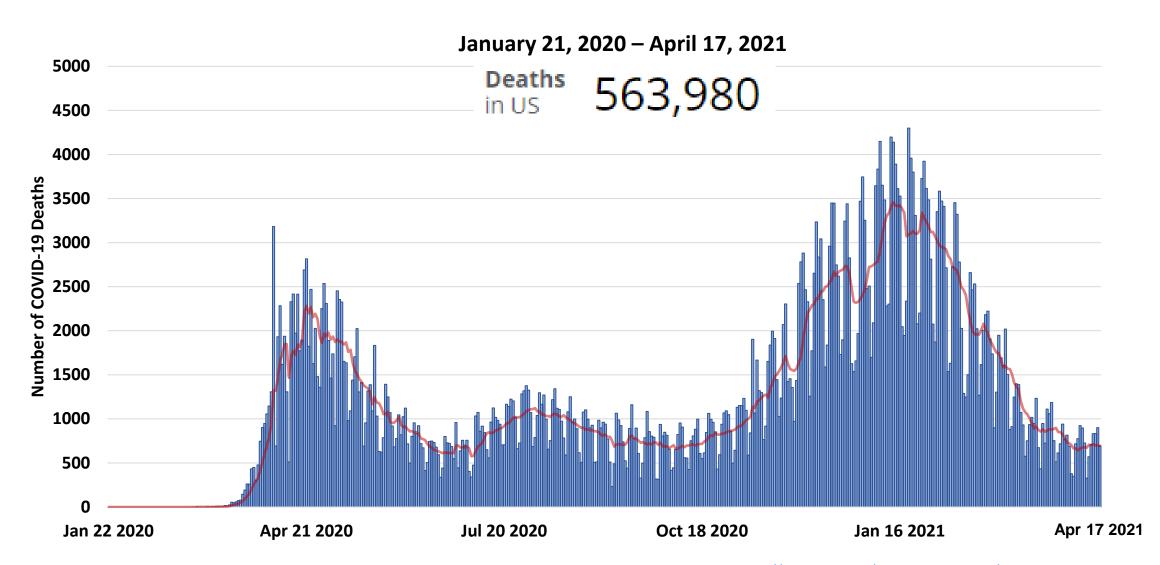
Public Health Problem:

- Recent COVID-19 Epidemiology
 - COVID-19 cases, hospitalization and deaths, stratified by age, sex, race and ethnicity
- Cerebral Venous Sinus Thrombosis (CVST)
- Heparin Induced Thrombocytopenia (HIT)
- Thrombosis after COVID-19 Disease
- AstraZeneca COVID-19 vaccines
 - Review of available global data

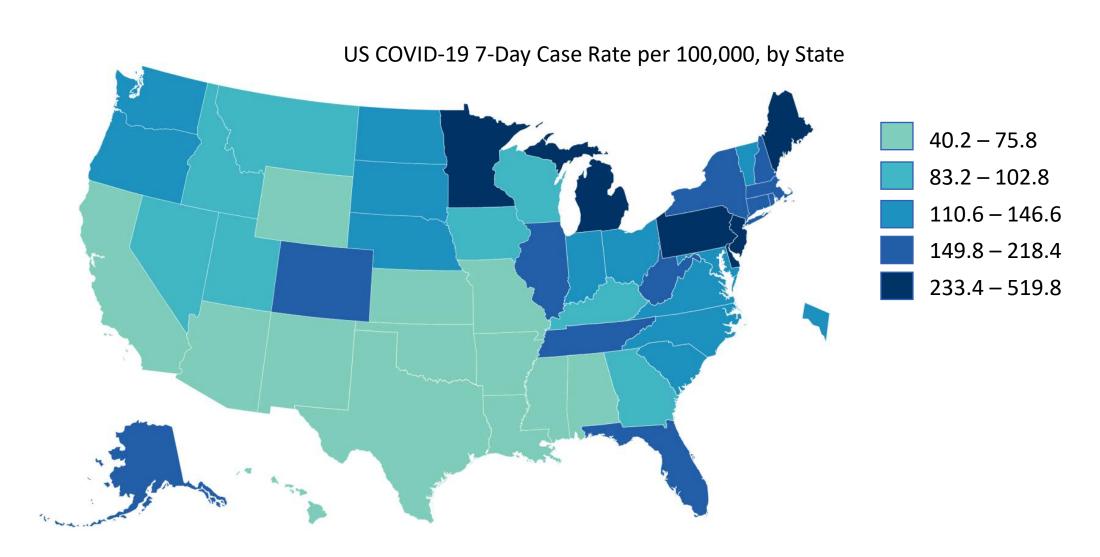
Trends in Number of COVID-19 Cases in the US



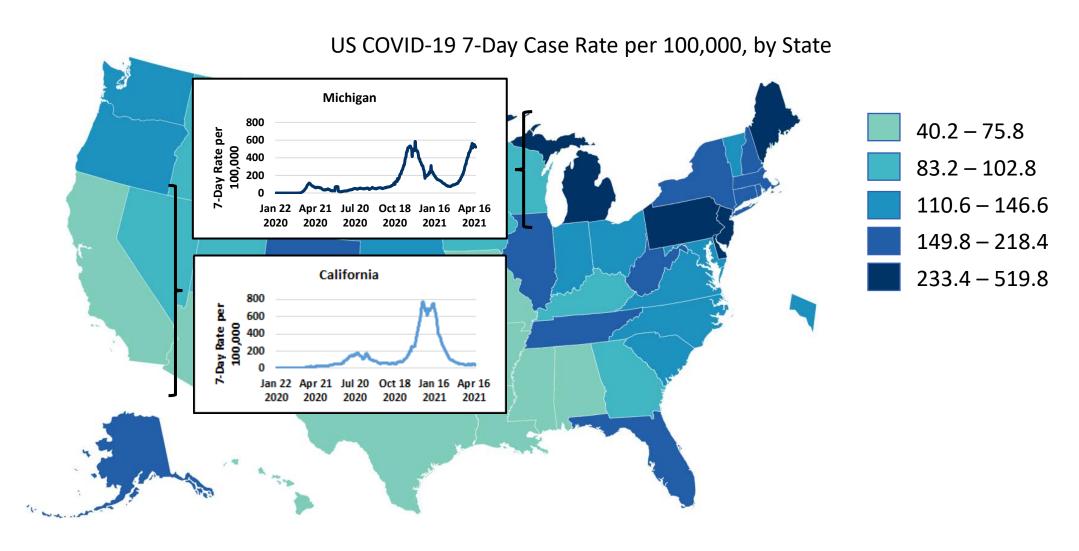
Trends in Number of COVID-19 Deaths in the US



COVID-19 Incidence Rates, by State

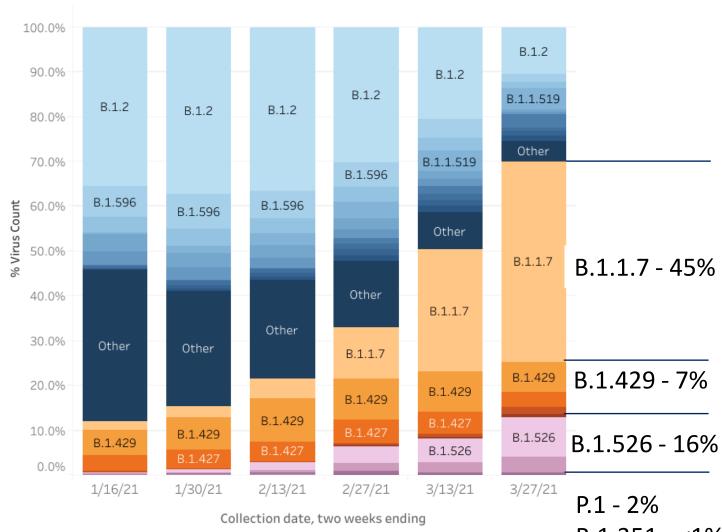


COVID-19 Incidence by State



SARS-CoV-2 Variants Circulating in the United States

Variant Proportions, January 3 - March 27, 2021



B.1.351 - <1%

Lineage	% Total	95%CI	Type	
B.1.1.7	44.7%	41.8-47.5%	VOC	
B.1.2	10.5%	9.5-11.6%		
B.1.526	8.9%	6.9-11.4%	VOI	
B.1.429	6.9%	5.2-9.0%	VOC	
B.1.1.519	4.5%	3.8-5.3%		
B.1.526.1	3.6%	3.0-4.2%	VOI	
B.1.526.2	3.2%	2.6-3.9%		
B.1.427	3.1%	2.4-4.0%	VOC	
B.1	1.6%	1.4-1.9%		
B.1.596	1.6%	1.3-2.0%		
P.1	1.5%	1.1-2.1%	VOC	
R.1	1.1%	0.9-1.4%		
B.1.575	1.1%	0.8-1.4%		
B.1.243	0.7%	0.5-0.9%		
B.1.1	0.7%	0.4-1.0%		
B.1.234	0.5%	0.3-0.6%		
B.1.351	0.7%	0.5-1.1%	VOC	
P.2	0.3%	0.2-0.4%	VOI	
B.1.525	0.3%	0.2-0.4%	VOI	
Other	4.7%	4.0-5.5%		
	B.1.1.7 B.1.2 B.1.526 B.1.429 B.1.526.1 B.1.526.2 B.1.526.2 B.1.427 B.1 B.1.596 P.1 R.1 B.1.575 B.1.243 B.1.243 B.1.1 B.1.234 B.1.351 P.2 B.1.525	B.1.1.7 44.7% B.1.2 10.5% B.1.526 8.9% B.1.429 6.9% B.1.1.519 4.5% B.1.526.1 3.6% B.1.526.2 3.2% B.1.427 3.1% B.1 1.6% B.1.596 1.6% P.1 1.5% R.1 1.1% B.1.575 1.1% B.1.243 0.7% B.1.243 0.7% B.1.234 0.5% B.1.351 0.7% P.2 0.3% B.1.525 0.3%	B.1.1.7 44.7% 41.8-47.5% B.1.2 10.5% 9.5-11.6% B.1.526 8.9% 6.9-11.4% B.1.429 6.9% 5.2-9.0% B.1.1.519 4.5% 3.8-5.3% B.1.526.1 3.6% 3.0-4.2% B.1.526.2 3.2% 2.6-3.9% B.1.427 3.1% 2.4-4.0% B.1 1.6% 1.4-1.9% B.1.596 1.6% 1.3-2.0% P.1 1.5% 1.1-2.1% R.1 1.1% 0.9-1.4% B.1.575 1.1% 0.8-1.4% B.1.243 0.7% 0.5-0.9% B.1.1 0.7% 0.4-1.0% B.1.234 0.5% 0.3-0.6% B.1.351 0.7% 0.5-1.1% P.2 0.3% 0.2-0.4% B.1.525 0.3% 0.2-0.4%	B.1.1.7

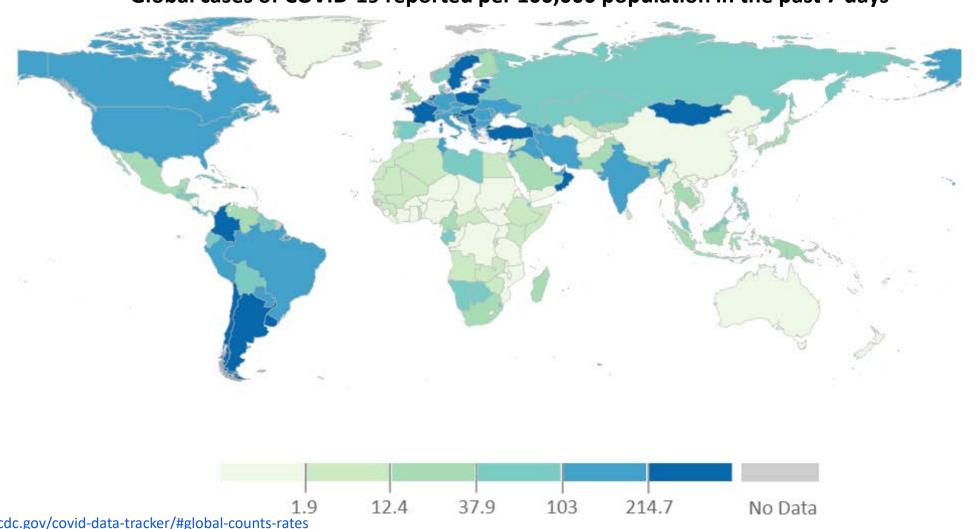
Summary data that appear in the table include specimen collection dates from March 14 through March 27, 2021.

^{*} Other represents >200 additional lineages, which are each circulating at <1% of viruses

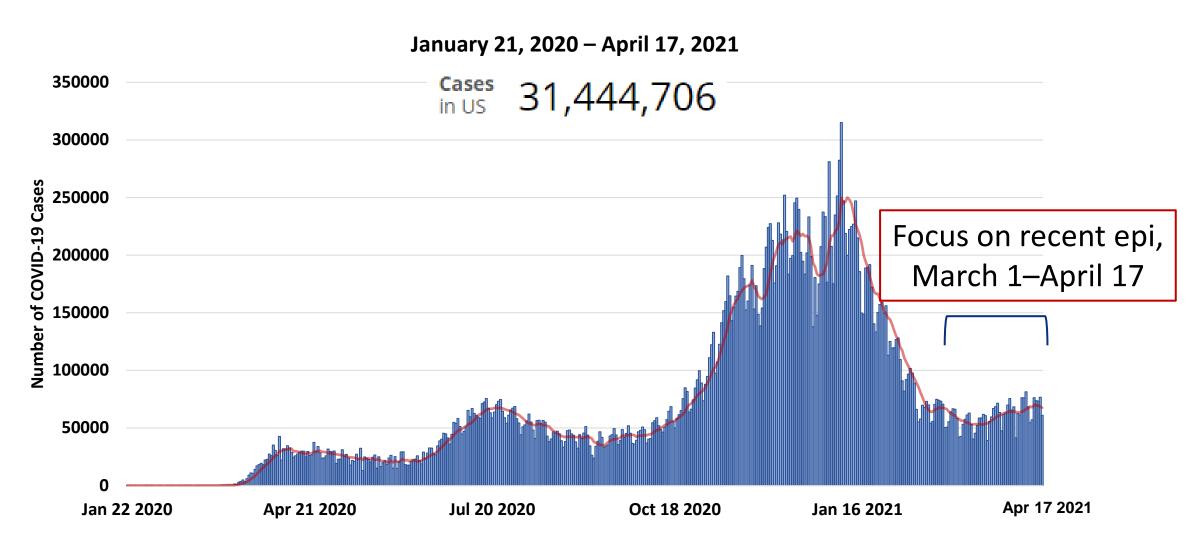
^{**} Most recent data are subject to change as samples from that period are still being processed.

Global COVID-19 Incidence Rates

Global cases of COVID-19 reported per 100,000 population in the past 7 days

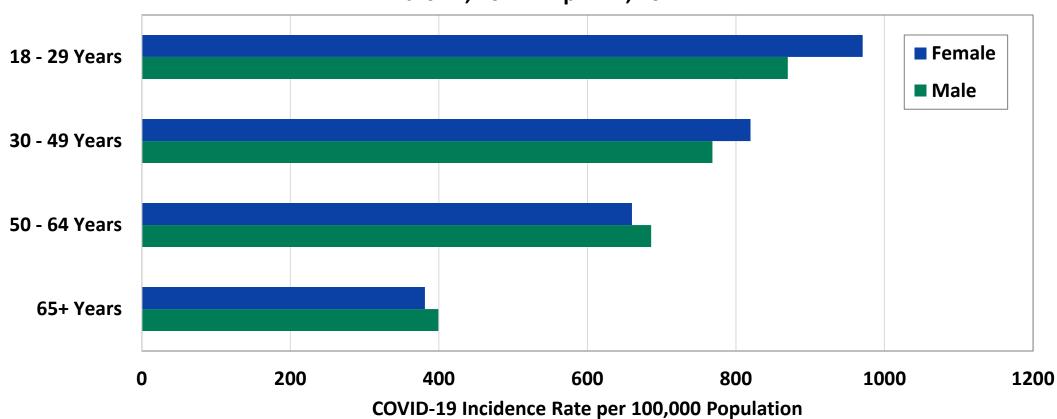


Trends in Number of COVID-19 Cases in the US



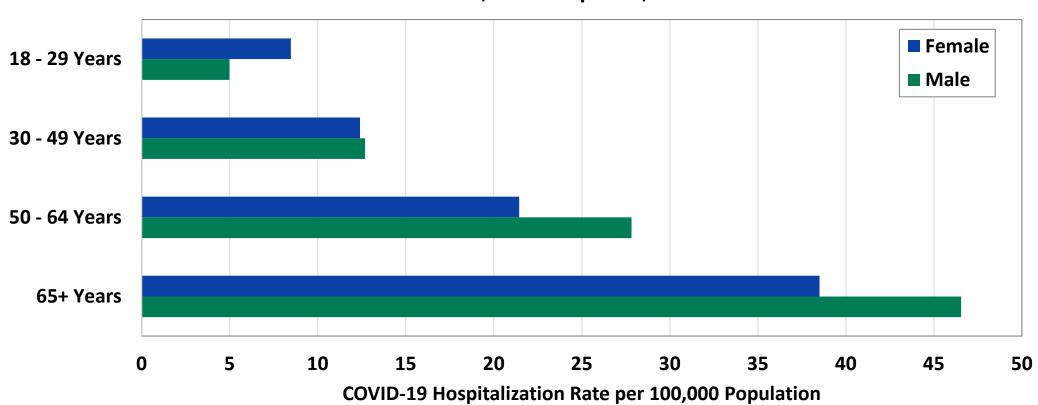
COVID-19 Incidence Rates, by Age Group and Sex

COVID-19 Incidence Rate per 100,000 Population, by Age Group and Sex March 1, 2021 – April 17, 2021



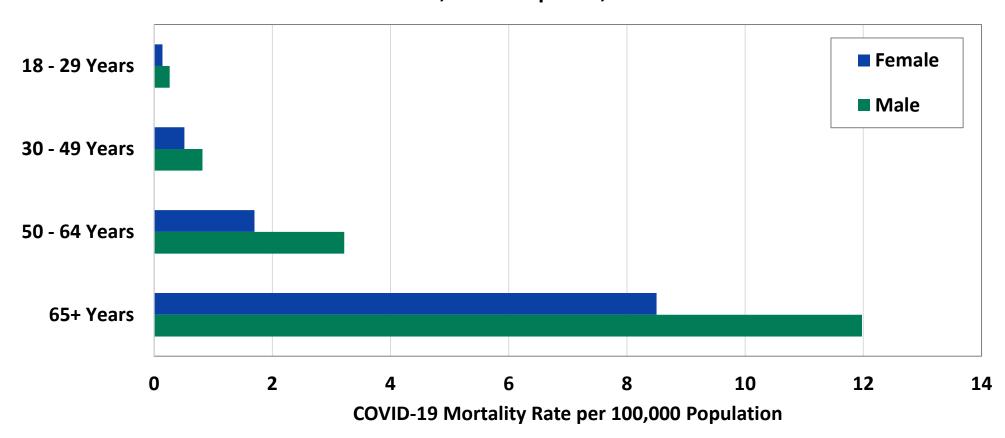
COVID-19 Hospitalization Rates, by Age Group and Sex

COVID-19 Hospitalization Rate per 100,000 Population, by Age Group and Sex March 1, 2021 – April 17, 2021

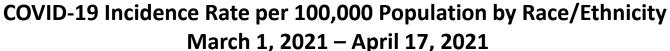


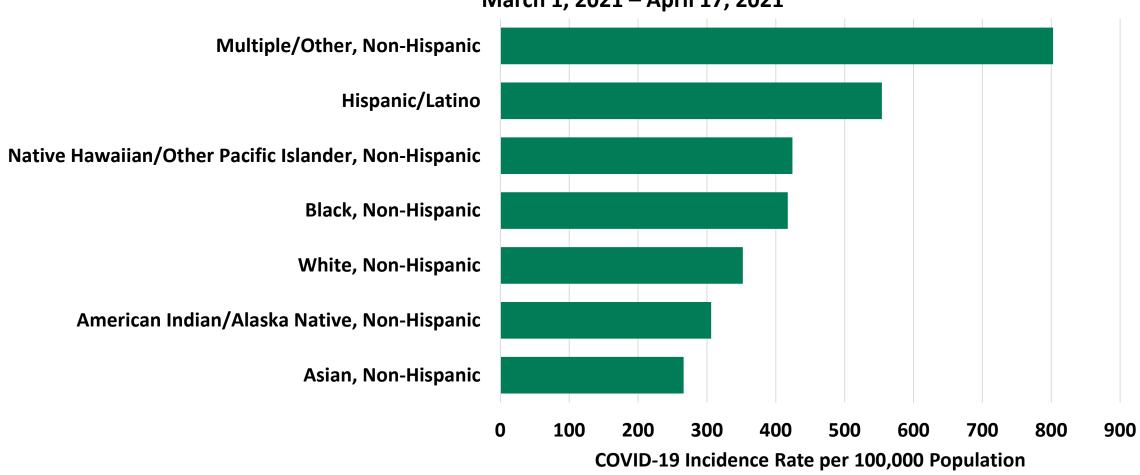
COVID-19 Mortality Rates, by Age Group and Sex

COVID-19 Mortality Rate per 100,000 Population, by Age Group and Sex March 1, 2021 – April 17, 2021



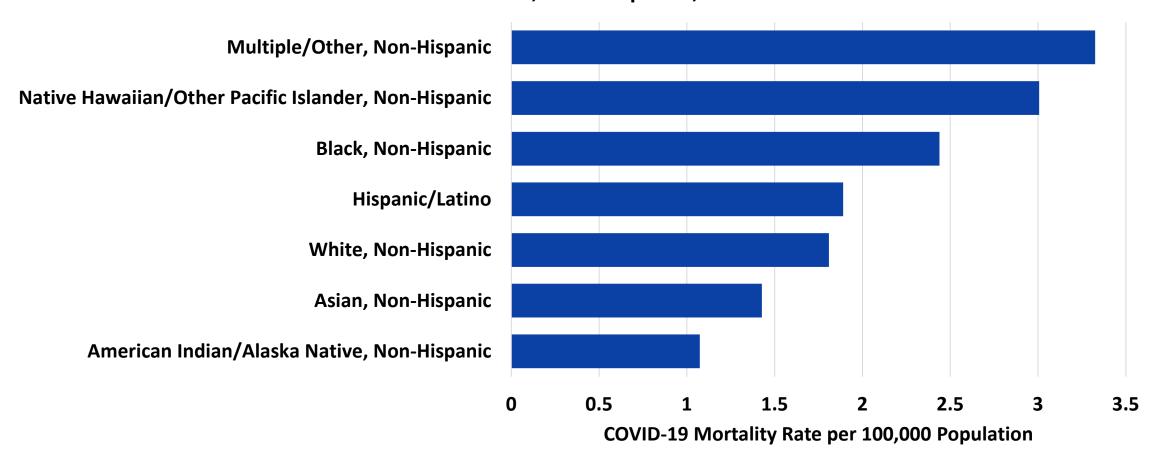
COVID-19 Incidence Rates, by Race/Ethnicity





COVID-19 Mortality Rate, by Race/Ethnicity

COVID-19 Mortality Rate per 100,000 Population by Race/Ethnicity March 1, 2021 – April 17, 2021



Summary of the COVID-19 Epidemiology

March 1-April 17, 2021

Incidence

- Cumulative incidence rate for adults: 710.9 per 100,000 population
- Younger females (18-29 years) have the highest incidence of new infections

Hospitalization

- Cumulative hospitalization rate for adults: 20.6 per 100,000 population
- Most hospitalizations still occur in persons aged ≥ 65 years
 - Proportion of hospitalizations occurring in persons aged ≥65 years declining

Mortality

- Cumulative mortality rate: 3.0 per 100,000 population
- Most COVID-19 deaths still occur in persons aged ≥ 65 years
 - Proportion of deaths occurring in persons aged ≥65 years declining

Epidemiology of Cerebral Venous Sinus Thrombosis (CVST) and Splanchnic Vein Thrombosis (SVT)

- Cerebral Venous Sinus Thrombosis (CVST) incidence: 14.5–28.5 per million U.S population
 - Incidence increasing in recent years (4% annually)
 - Higher in women aged 18–49 years
 - Risk factors (e.g. hereditary thrombophilia, oral contraceptives, obesity) identified in up to 85% of cases
 - Mortality ~5-10%
- Splanchnic Vein Thrombosis (SVT) incidence: 84–179 per million U.S. population
 - Incidence higher among men
 - Risk increases with age
- Incidence with thrombocytopenia much lower than without thrombocytopenia
 - CVST with thrombocytopenia: 0.7–1.6 per million U.S. population

Heparin-Induced Thrombocytopenia with Thrombosis (HITT)

- Heparin-induced thrombocytopenia (HIT) occurs in 0.5% to 1% of patients exposed to unfractionated heparin for medical and surgical indications
 - Incidence: 23–45 per million total U.S. population*
- Of patients with HIT, thrombosis occurs in about 20%–64% (called HITT)
- Immune mediated antibodies against platelet factor 4 (PF4) & heparin
- Risk factors for developing thrombosis
 - Genetic polymorphisms
 - Lower platelet count (and earlier fall in count)
 - Higher titer of anti-heparin/PF4 antibodies
 - Prior surgery (cardiac, orthopedic, trauma)
 - Cardiovascular disease
- Limited case series published on HIT occurring after COVID-19

^{*} Source: HCUP NIS 2018 and Marketscan (Continuously-enrolled Commercial Insurance and Medicaid) for 2019, unable to distinguish autoimmune HIT vs heparin-induced HIT Arepally et al. 2021. https://www.ahajournals.org/doi/epub/10.1161/ATVBAHA.120.315445; Nand et al. 1998 <a href="https://doi.org/10.1002/(SICI)1096-8652(199709)56:1<12::AID-AJH3>3.0.CO;2-5">https://doi.org/10.1002/(SICI)1096-8652(199709)56:1<12::AID-AJH3>3.0.CO;2-5; Fabris et al 2002. https://onlinelibrary.wiley.com/doi/epdf/10.1046/j.1365-2796.2002.01021.x; Greinacher et al. 2005. https://www.thieme-connect.com/products/ejournals/abstract/10.1160/TH04-12-0825

CVST associated with COVID-19

- Systematic review, meta-analysis of CVST among patients hospitalized for COVID-19
 - Estimates between 0.03% and 0.08% of hospitalized COVID-19 patients
- Estimated risk of 5–6 cases of CVST per million SARS-COV-2 infections*
- CVST + thrombocytopenia in COVID-19 patients is extremely rare
- Pathology appears different than TTS after COVID-19 vaccines
 - PF4/heparin specific antibodies negative by ELISA or platelet functional assay for confirmed COVID-19 patients (n=222), including 10 with thromboembolic complications

^{*} Data source: Premier Healthcare Database, January 2020-January 2021

Acronyms: Cerebral Venous Sinus Thrombosis (CVST), Thrombosis with Thrombocytopenia Syndrome (TTS)

Katsanos et al. (2020) https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7753413/; Baldini et al. (2021) https://doi.org/10.1111/ene.14727; Mowla et al. (2020) https://doi.org/10.1016/j.jns.2020.117183; Greinacher et al. Research Square preprint (Apr 9, 2021): https://www.researchsquare.com/article/rs-404769/v1

CVST associated with COVID-19

- Recent study using electronic health records (81m patients, mostly U.S.) to estimate CVST & portal vein thrombosis (PVT) incidence in 513,284 COVID-19 cases
 - Highest baseline CVST incidence observed over any 2-week period: **0.41** per million people
 - CVST incidence 2 weeks after COVID-19 diagnosis (among hospitalized): 39 per million
 - CVST incidence higher after COVID-19 diagnosis than after mRNA vaccines (4.1 per million) or after influenza disease (0.0 per million)
 - Mortality was 20% for CVST and 19% for PVT

Limitations

- Unable to assess incidence of CVST after adenovirus vector vaccines (Janssen, AstraZeneca)
- Did not provide rates of CVST + thrombocytopenia
 - Limited ability to directly compare to rates of CVST + thrombocytopenia after vaccines reported in US or Europe

Thrombosis with Thrombocytopenia Syndrome (TTS) after AstraZeneca vaccine in Europe

- As of 4 April 2021, **169** cases of CVST & **53** cases of splanchnic vein thrombosis reported to EudraVigilance. ~34 million people vaccinated in EEA & UK by this date.
 - EU ~10 cases per million (1 case per 100,000) vaccinated adults
 - Higher in younger adults compared to older adults
 - Most of cases in women aged <60 years within 2 weeks of receiving 1st vaccine dose
- European Medicines Agency concluded benefit/risk ratio still favorable to use vaccine
 - Causal association plausible
 - Unable to identify definitive cause, but possibly similar to heparin-induced thrombocytopenia
 - No specific risk factors to date (epidemiology may be related to vaccine delivery)
 - Added unusual blood clots with low platelets to the label as very rare side effect

UK decision on use of AstraZeneca vaccine — April 14, 2021

- Through April 14: 168 reports of blood clotting with low platelets
 - 77 CVST with thrombocytopenia; 91 in other major veins with thrombocytopenia
 - 93 women, 75 men, aged 18–93 years
 - 32 deaths
 - Most cases occurred after first vaccine dose; one case occurred after second dose
- Rate: 7.9 per million (21.2 million AZ doses given)
- Benefits continue to outweigh risks stronger evidence for a link of vaccine to extremely rare blood clots with lower platelets, but more work needed
 - Careful consideration be given to those at higher risk of blood clots because of medical conditions or pregnancy
 - Continue to give second doses, except to those with blood clots and low platelets after first dose
- Recommended ages 18–29 years at low risk of infections be offered other vaccines

EMA's safety committee (PRAC) meeting on U.S. TTS cases after Janssen vaccine, April 20, 2021

- Reviewed evidence on 8 U.S. reports of serious cases of unusual blood clots with low levels of blood platelets, one of which had a fatal outcome. Over 7 million U.S. people had received Janssen vaccine, as of 13 April 2021
- Cases reviewed very similar to cases occurring with AstraZeneca COVID-19 vaccine
- Concluded that a warning about unusual blood clots with low blood platelets should be added to the product information for Janssen COVID-19 vaccine
- Very rare event, and the overall benefits of Janssen COVID-19 vaccine in preventing COVID-19 outweigh the risks of side effects
- Emphasized importance of healthcare provider awareness

Public Health Problem:

COVID-19

<u>Hospitalization</u>: 200 per million population

<u>Death</u>: 30 per million population

CVST after COVID-19

5-6 per million SARS-COV-2 infections

HIT

23–45 per million population

CVST

14.5–28.5 per million population

CVST + Thrombocytopenia

0.7–1.6 per million population

TTS after AZ vaccine

10 per million vaccinated population

<u>EU</u>:

<u>UK</u>:

7.9 per million vaccinated population

EtR Domain: Benefits and Harms



Benefits and Harms:

- Benefits of Janssen COVID-19 vaccine
 - Prevention of COVID-19 cases, hospitalizations and deaths
- Harms of Janssen COVID-19 vaccine
 - Estimated cases of TTS after Janssen COVID-19 vaccine, by age and gender
- Benefit and Risk summary

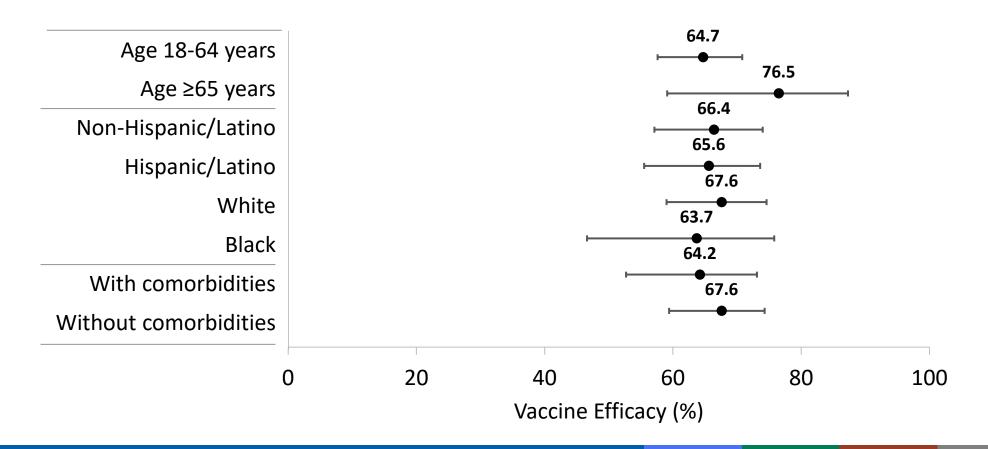
Benefits of the Janssen COVID-19 vaccine

- The clinical trial demonstrated efficacy against symptomatic, laboratory-confirmed COVID-19. The overall efficacy was **66.3%** (95% CI: 59.9%, 71.8%)
- Vaccine efficacy against COVID-19 associated hospitalization was 93% (95% CI: 71%, 98%)
- Higher efficacy against severe outcomes than for any symptomatic COVID-19
 - VE against deaths due to COVID-19: 100%
- Efficacy against severe disease[†] remained high across world regions (73-82%*), suggesting protection against severe illness with variant strains

†Definition: Respiratory Rate ≥ 30, Heart Rate ≥125, SpO2≤ 93% on room air at sea level or PaO2/FIO2< 300 mm Hg; OR respiratory failure or Acute Respiratory Distress Syndrome (ARDS), defined as needing high-flow oxygen, non-invasive or mechanical ventilation, or ECMO; OR evidence of shock (systolic blood pressure <90mmHg, diastolic BP<60mmHg or requiring vasopressors); OR significant acute renal, hepatic or neurologic dysfunction; OR admission to an intensive care unit or death *Assessed ≥ 14 days post vaccination

Benefits of the Janssen COVID-19 vaccine

Similar efficacy for across age, sex, race, and ethnicity categories, and those with underlying medical conditions at ≥14 days post-vaccination



Benefits of the Janssen COVID-19 vaccine

- Vaccine shipment and storage (3 months) at refrigerator temperatures (2-8°C)*
 - Refrigerator-stable vaccine could facilitate the availability of the Janssen COVID-19 vaccine in many community settings and mobile sites
- Single-dose series
- Easier to reach some disproportionately affected groups such as: homeless, rural residents, justice-involved, disabled, homebound, or with no/limited access to healthcare

* Long-term storage at standard freezer temperatures (-20°C)

Potential Harms of the Janssen COVID-19 vaccine

- 7.98 million vaccine doses administered*and 15 confirmed TTS cases as of April 21, 2021
 - Additional potential TTS cases under review, including potential male cases

		Femal	es	Males		
Age group	Cases	Doses admin	Reporting rate [†]	Cases*	Doses admin	Reporting rate [†]
18-49 years old	13	1,866,294	7.0 per million	0	1,977,330	0 per million
50+ years old	2	2,125,239	0.9 per million	0	2,010,144	0 per million

^{*} Source of doses administered: https://covid.cdc.gov/covid-data-tracker/#vaccinations; Some age- and sex-specific doses administered data were imputed

[†] Reporting rate = TTS cases per 1 million Janssen COVID-19 vaccine doses administered

^{*} One TTS case occurred in the Phase 3 trial in a male aged 18-49 years.

Summary of Different Risk-Benefit Analyses

Population Level Risk-Benefit Analysis

Type of analysis Objective 1. Population (6-month period) Quantify COVID-19 infections, hospitalizations, and deaths averted under different assumptions about resumption of Janssen vaccination Quantify population-level, age-specific benefits and harms of resuming vaccination with Janssen COVID-19 vaccine

Individual Level Risk-Benefit Analysis

Type of analysis	Objective		
2. Direct (1-month period)	 Quantify direct age and sex-specific benefits and harms, per million Janssen vaccine doses 		

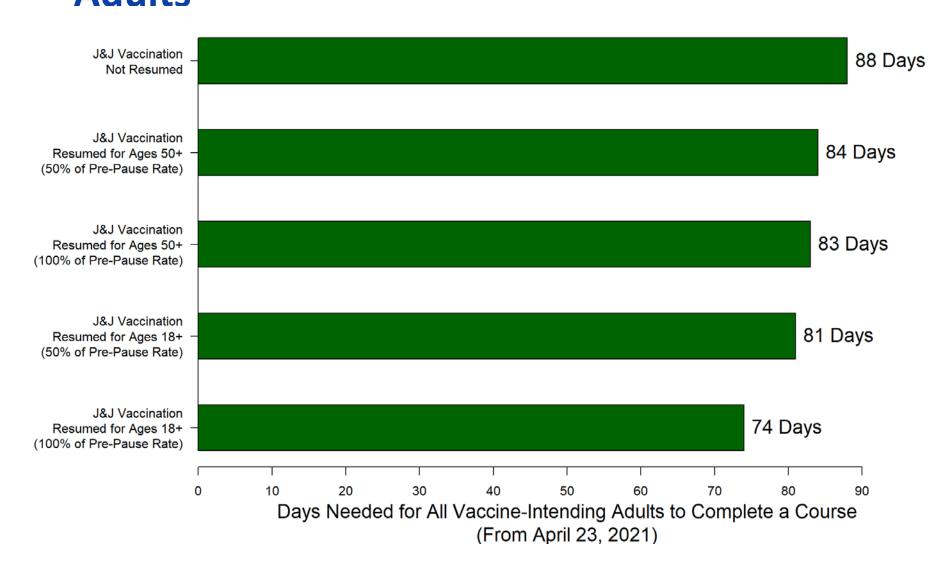
Modeled Janssen COVID-19 Vaccine Scenarios

- Objective: Quantify COVID-19 infections, hospitalizations, and deaths under different assumptions about resumption of Janssen vaccination
- In all scenarios we assume continued use of mRNA vaccines
- Evaluated the following scenarios:
 - Janssen vaccination not resumed
 - Janssen vaccination resumed on April 24th for all adults aged 18+ years
 - **50%** of pre-pause J&J administration rate (in 18+ age group)
 - 100% of pre-pause J&J administration rate (in 18+ age group)
 - Janssen vaccination resumed on April 24th in adults aged 50+ years only
 - 50% of pre-pause J&J administration rate (in 50+ age group)
 - 100% of pre-pause J&J administration rate (in 50+ age group)

Model Overview & Assumptions

- Compartmental model used to simulate incident infections, hospitalizations, and deaths over the course of the epidemic in the US
 - Stratified by age, essential-worker status, and underlying conditions
 - Calibrated to observations through Spring 2021. Future activity based on external scenarios.
- Two vaccine types: mRNA and Janssen (VE informed by RCTs)
 - Immunity develops 14 days after administration
 - 28 days between mRNA doses
- No loss to follow-up or delays in administration
- No waning of immunity over time

Modeled Time to Complete Vaccination for All Intending Adults



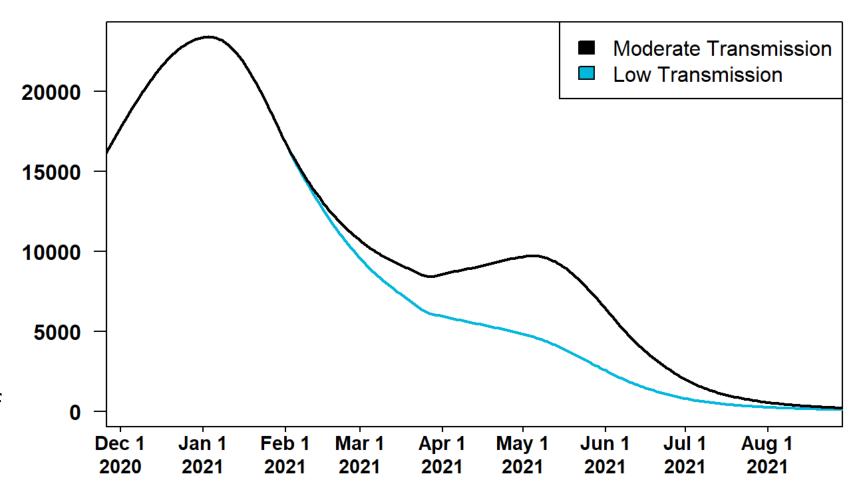
Multiple assumptions:

- Vaccine intent in each age group remains same
- Administration of mRNA vaccines continues at the same rate as before pause
- In practice, pace of administration may slow as those experiencing barriers to access and/or greater hesitancy comprise a greater share of the unvaccinated

Modeling Transmission Scenarios

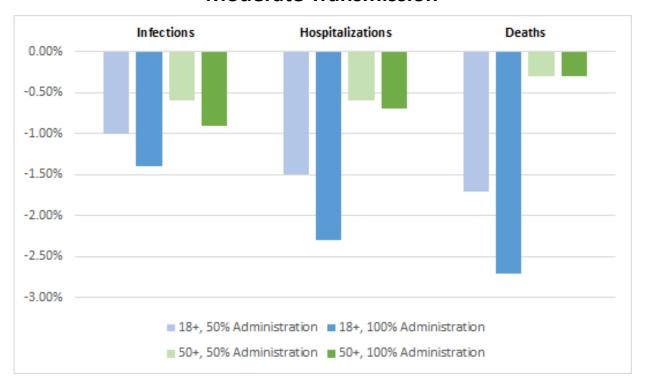
Incident Daily Hospitalizations

- Calibrated assuming no interruption of Janssen vaccination under two scenarios reflecting continued level of nonpharmaceutical interventions (i.e., low transmission and moderate transmission)
- Calibrated to Round 4 of the COVID-19 Scenario Hub



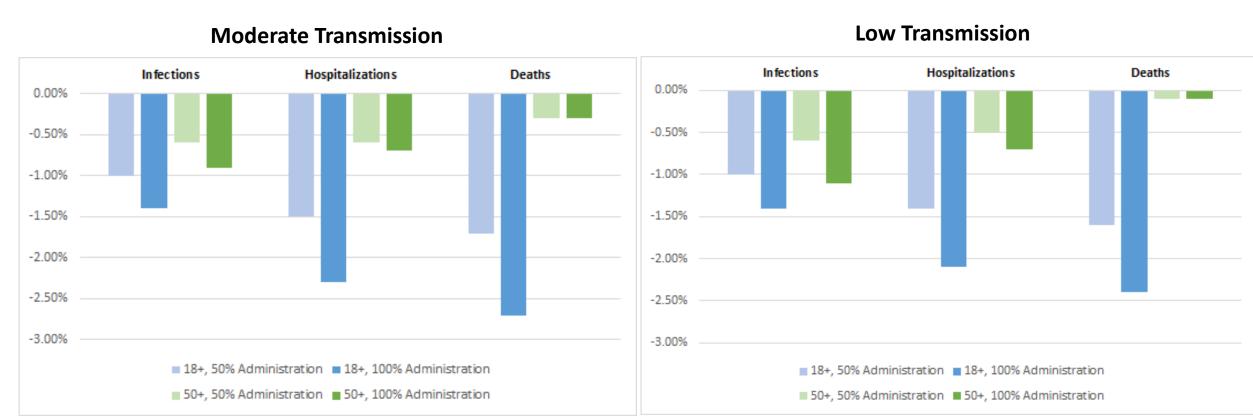
Percent Change* in Outcomes 6 Months Post-Pause, by Janssen Resumption Strategy

Moderate Transmission



^{*} compared to a scenario in which administration of the J&J vaccine does not resume

Percent Change* in Outcomes 6 Months Post-Pause, by Janssen Resumption Strategy



There is a benefit of resuming Janssen vaccination in terms of infections, hospitalizations and deaths prevented under different epidemiologic assumptions

^{*} Compared to a scenario in which administration of the J&J vaccine does not resume

Population Risk-benefit assessment

- Quantify age-specific risks and benefits of resuming vaccination with Janssen COVID-19 vaccine
 - Risks: Number of TTS
 - Benefits: Prevention of COVID-19-related hospitalizations,
 ICU admissions, and deaths

Population Risk-Benefit Inputs

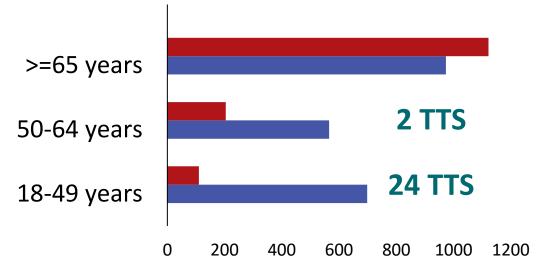
- Estimates from compartmental model
 - Numbers of hospitalizations and deaths due to COVID-19
 - Number of persons estimated to receive Janssen COVID-19 vaccine after pause
 - Conditions: Low and moderate transmission; 50% and 100% of prior vaccination rate; Resuming in ages 18+ and 50+, vs no resumption
 - 6-month time horizon
- ICU admissions per hospitalization (CDC Pandemic Planning Scenarios)
- Counts of TTS cases by age (CDC data)
- Numbers vaccinated with Janssen COVID-19 vaccine, by age, before pause (CDC data)

Benefits and harms of resuming vaccination for ages ≥18 years vs. ≥50 years over 6-month period

Moderate transmission; Vaccination resumed at 50% of rate before pause



Resume vaccination: age 18+



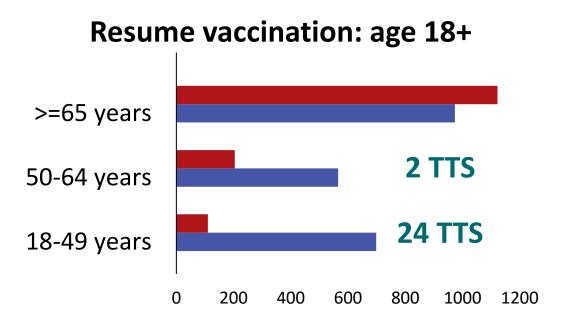
26 TTS in 9.8M vaccinations
Prevent 1,435 deaths, 2,236 ICU admissions

¹ Based on observed cases adjudicated as of 4/21/2021 NOTE: in Phase III RCT, one male in 18-49 year age group experienced TTS; not included in this analysis Acronyms: Thrombosis with Thrombocytopenia Syndrome (TTS)

Benefits and harms of resuming vaccination for ages ≥18 years vs. ≥50 years over 6-month period

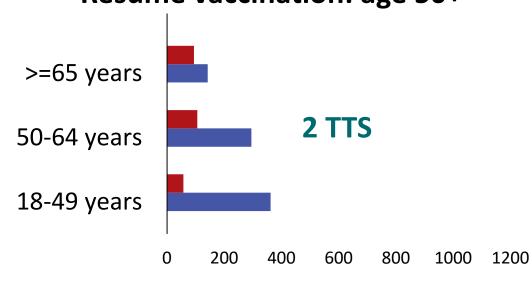
Moderate transmission; Vaccination resumed at 50% of rate before pause

Deaths averted



Resume vaccination: age 50+

ICU admissions averted



2 TTS in 3.6M vaccinations
Prevent 257 deaths, 779 ICU admissions

26 TTS in 9.8M vaccinations

Prevent 1,435 deaths, 2,236 ICU admissions

¹Based on observed cases adjudicated as of 4/21/2021

NOTE: in Phase III RCT, one male in 18-49 year age group experienced TTS; not included in this analysis

Acronyms: Thrombosis with Thrombocytopenia Syndrome (TTS)

Summary of population-level risks and benefits by recommendation, all scenarios

Recommendation for all persons aged 18+

- Risks: Expect 26–45 TTS cases,
 depending on uptake
- Benefits: Depend on uptake, amount of transmission
 - 800–3,500 fewer ICU admissions
 - 600–1,400 fewer deaths

Recommendation for all persons aged 50+

Risks: Expec**2–3 TTS**cases, depending on uptake

Benefits: Depend on uptake, amount of transmission

300–1000 fewer ICU

admissions

40-250 fewer deaths

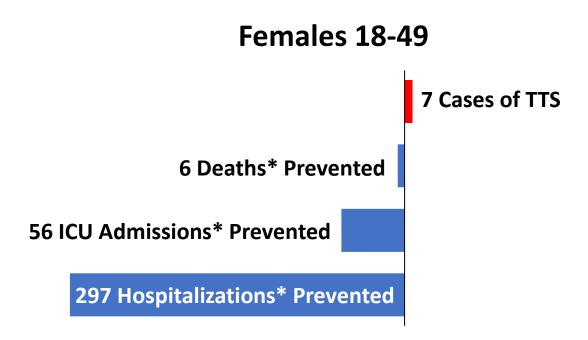
Note: Benefits of vaccination apply to the whole population over a 6-month period, and result from direct and indirect effects.

Estimation of direct benefits and risks to vaccinated persons

- Evaluate direct benefits and risk, per million Janssen vaccine doses
- Used to visualize sex differences in risk and benefits
- Calculations based on recent hospitalization incidence, VE, Janssen vaccinations to date, number of persons already vaccinated
- 30-day period

Risks and benefits by for females, by age group

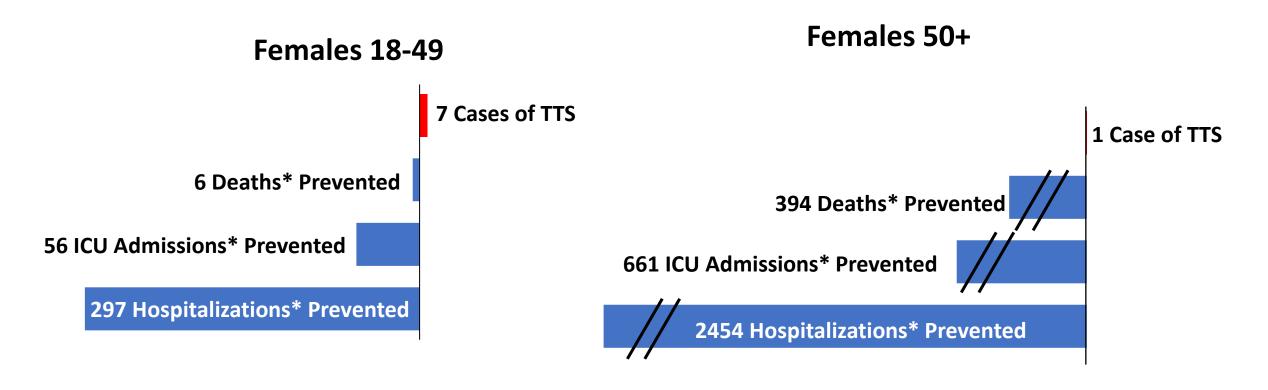
For every **1 million** doses of vaccine given with current US exposure risk¹



^{*} Deaths, ICU admissions, and deaths due to COVID-19 Acronyms: Thrombosis with Thrombocytopenia Syndrome (TTS)

Risks and benefits females, by age group

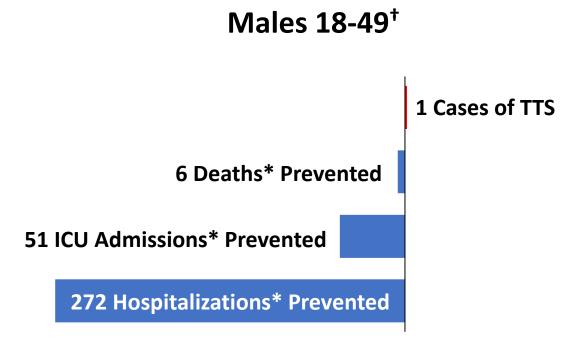
For every **1 million** doses of vaccine given with current US exposure risk¹



^{*} Deaths, ICU admissions, and deaths due to COVID-19 Acronyms: Thrombosis with Thrombocytopenia Syndrome (TTS)

Risks and benefits males, by age group

For every 1 million doses of vaccine given with current US exposure risk1



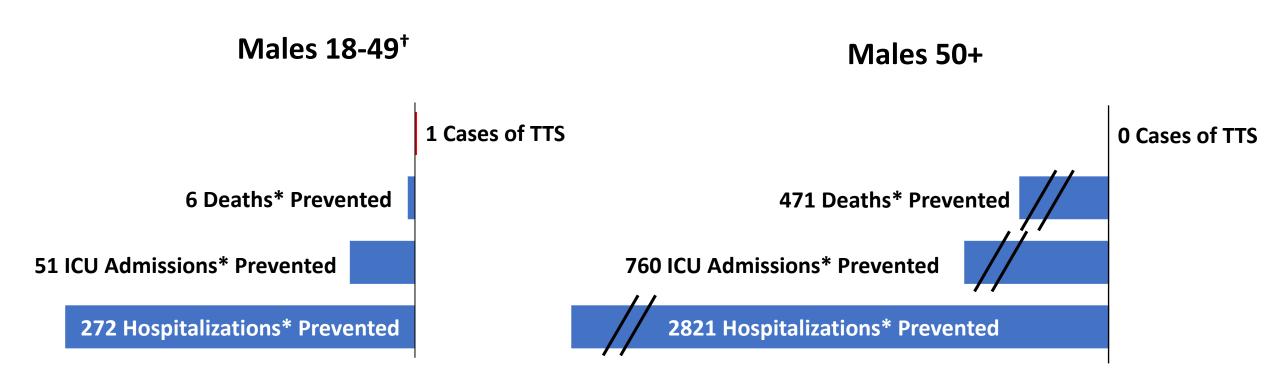
Acronyms: Thrombosis with Thrombocytopenia Syndrome (TTS)

[†]Analyses incorporated one TTS case that occurred in the Phase 3 trial in a male aged 18-49 years.

^{*}Deaths, ICU admissions, and deaths due to COVID-19

Risks and benefits males, by age group

For every 1 million doses of vaccine given with current US exposure risk1



[†]Analyses incorporated one TTS case that occurred in the Phase 3 trial in a male aged 18-49 years.

^{*}Deaths, ICU admissions, and deaths due to COVID-19

Risk-benefit interpretations

- Population
 - Takes into account direct and indirect (herd) effects of vaccination
 - Incorporates availability of different vaccines
 - Simulates incidence, hospitalizations, and deaths over course of pandemic
 - 6-month time horizon
- Shows large population benefit of vaccination relative to rare TTS

- Direct
 - Considers individual benefits of vaccination vs. individual risks
 - Only considers getting Janssen vaccine vs. not getting a vaccine
 - Short, 1-month time horizon
- Shows positive balance for benefits vs. risks for all age and sex groups
- Balance of risks and benefits varies by age and sex

EtR Domains: Values and Acceptability



Values and Acceptability:

- Intent to receive 1-dose COVID-19 vaccine
 - Data stratified by age, sex, race/ethnicity and income
- Intent to receive Janssen COVID-19 vaccine over time
- Effect on overall vaccine confidence

Values and Acceptability: Intent to receive 1-dose COVID-19 vaccine

- Among unvaccinated respondents, if both a <u>2-dose and 1-dose</u> COVID-19 vaccine were available, which would you choose? Assume that both types of vaccines are safe.
 - Respondents could choose 1-dose, 2-dose, either, neither
 - Data collected February 2021
- Examined by age, sex, race/ethnicity, and income
 - Overall, 6% exclusively preferred 1-dose vaccine
 - No differences in proportion that preferred 1-dose by age, sex, or income
 - Significantly more Hispanic than White respondents (11%) preferred 1-dose

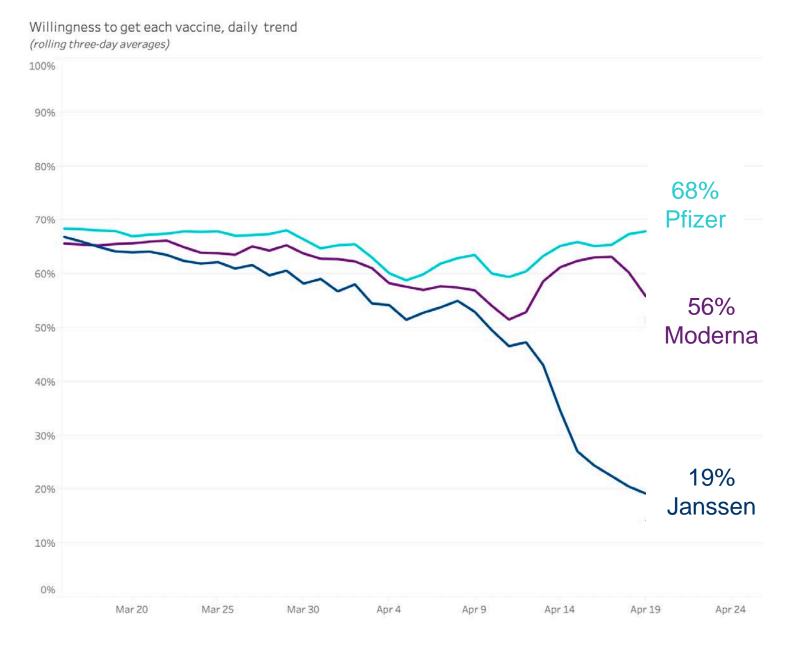
Values and Acceptability:

Intent to receive Janssen COVID-19 vaccine over time

- Only 37% of respondents called the Janssen COVID-19 vaccine safe after the pause was announced¹
 - Drop of 15% in two to three days
- Americans now much less likely to prefer the Janssen COVID-19 vaccine²
 - 13% decline in preference for the Janssen COVID-19 vaccine
 - Declined 9% to 25% across age and race categories

Willingness to receive the J&J vaccine dropped immediately as a result of the pause

(Results among those who say "yes" they plan to get vaccinated but haven't yet)









Values and Acceptability:

Effect on overall vaccine confidence

- Drop in vaccine confidence does not appear to extend to the Pfizer-BioNTech and Moderna COVID-19 vaccines¹
 - 59% consider them safe
 - 19% feel they are unsafe
- Recent poll did not suggest reduction in intent to be vaccinated²
 - 40% more likely to receive COVID-19 vaccine compared to one month ago
 - 36% report no change in intent
- A different survey found half of the unvaccinated are less inclined to receive COVID-19 vaccine after the pause, regardless of brand³

^{1. &}lt;a href="https://today.yougov.com/topics/politics/articles-reports/2021/04/15/johnson-johnson-vaccine-confidence">https://today.yougov.com/topics/politics/articles-reports/2021/04/15/johnson-johnson-vaccine-confidence

^{2.} deBeaumont Foundation Poll, April 15-16, 2021. Vaccine Confidence Grows Despite J&J Pause

^{3.} CVS Health Survey- COVID-19 Vaccine Brand Preferences and Hesitancy Post J&J Pause

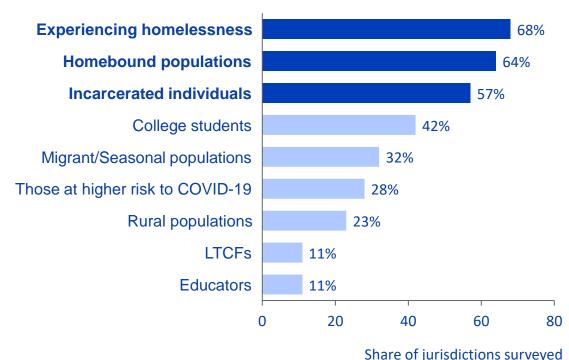
EtR Domain: Feasibility



Feasibility: Jurisdictions' pre-pause use of Janssen vaccine

Populations: Focus on reaching those experiencing homelessness, homebound or currently incarcerated

Q: Prior to the pause in administration of the Janssen vaccine, which populations had you focused on vaccinating with this product?



Vaccination settings: Three core settings used by jurisdictions to administer Janssen vaccine

Mobile vaccination

Temporary PODs and mobile vans able to reach transient, rural and

homebound individuals

Emergency departments

- Provided at discharge from urgent care or ER departments
- Particularly for 'safety-net' hospitals reaching transient groups

Student health centers

 On-campus vaccination centers with ambition to vaccinate students unable or less likely to return for second dose at end of semester

Feasibility: Impact if Janssen recommended <u>for specific</u> populations

Jurisdictions may need to reconfigure some vaccination sites, update scheduling tool, and have difficulty serving disproportionately affected populations



Changes to vaccination sites and schedulers might include...

Providers may need to carry multiple vaccines, if recommendation is restricted by sex.

Challenging to set-up dedicated community PODs if Janssen recommended for specific groups.

Would require IT systems update to internal scheduling and pre-screening tools.



Health depts. expressed concern about communicating change

Concerns about difficulty of communicating rationale for specific groups to public

Potential need to revise public-facing comms materials, provider training collateral, alongside re-training staff.

Expect low uptake on Janssen vaccine given negative publicity.



Greater difficulty serving disproportionately affected populations

Likely to see drop in completed series for those "at risk of loss to follow-up" e.g., experiencing homelessness, seasonal workers.

More challenging to reduce gap in vaccine disparities for racial and ethnic minorities through mobile vaccination.

Would increase barriers to access in rural and hard-to-reach areas.

Feasibility: Impact if Janssen were <u>no longer</u> recommended

Jurisdictions are particularly concerned about 2nd dose management and equity





Janssen provided flexibility to jurisdictions to...

Avoid additional second dose management, particularly for transient and hard-to-reach populations

Run mobile vaccination clinics without need for return visits

Reduce administrative burden on providers

Fully vaccinate college students before end of school year





Convenience of single dose appeals to many recipients

Some individuals hesitant about receiving an mRNA vaccine

Possibility of second dose side effects causes some to favor Janssen

Some providers with lower volumes of patients have preference for single dose vaccine



Greater difficulty serving disproportionately affected populations

Increased challenge to reach homebound, transient, and rural populations because of need to administer second dose

Less flexibility to use mobile vaccination units

Reduced ability to vaccinate upon ED/hospital discharge

Decreased vaccine supply from loss of Janssen could harm vaccine access

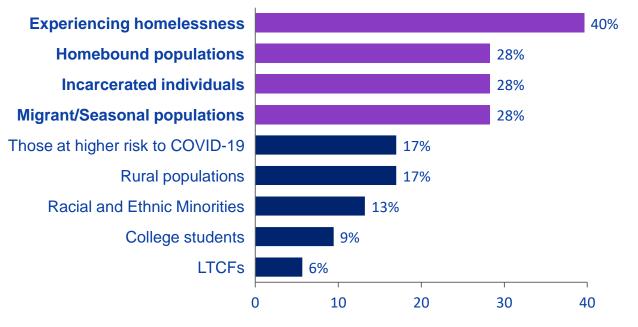
EtR Domain: Equity



Equity: Jurisdictions concerned revised recommendations would disproportionately affect several populations

Jurisdictions frequently raised four populations at risk of disproportionate impact

Q: Which, if any, populations would be disproportionally impacted if Janssen vaccine was no longer recommended or recommended for only a subset of the population?



Share of jurisdictions surveyed

Examples raised

Region 6 jurisdiction: "Most concerned about sub-populations that are difficult to reach e.g., people exp. homelessness, people who are working, as well as difficult geographic areas."

Region 2 jurisdiction: "The hardest to reach transient populations such as the homeless and those moving through substance use treatment programs, mental health treatment programs, etc. would be most impacted, given that they are among the most difficult individuals with whom to connect to provide second dose vaccination"

Region 10 jurisdiction: "...not having this vaccine would have an impact on trying to reduce the gap in vaccine disparities. We have community partners lined up to host events using the Janssen vaccine that are working with vulnerable populations in the state."

Policy Options



Policy Options for Janssen Policy Recommendations

Do **not** recommend use of Janssen vaccine

Recommend use of Janssen vaccine in **all adults**≥18 years of age

Recommend use of Janssen/J&J COVID-19 vaccine in **some** populations

Policy Options for Janssen Policy Recommendations

- Recommend against use for all persons
- Reaffirm recommendations for all age and sex
 - FDA to include warning statement with EUA
- Recommend vaccination only for adults ≥50 years of age
- Reaffirm recommendations for use; women aged <50 years should be aware
 of the increased risk of TTS, and may choose another COVID-19 vaccine
 (i.e. mRNA vaccines)

	Pros	Cons
Recommend against use in all persons	 No further cases of CVST/TTS after Janssen vaccine 	 Would remove choice from individuals Could lead to excess COVID-19 cases & deaths Could disproportionately impact at-risk populations with barriers to access or difficulty returning for 2nd dose
	Allow for flexibility/choiceAllow for use of the vaccine in	 Burden on individual to understand risk; health dept/providers to convey risk May lead to more cases of TTS At-risk populations for COVID-19 likely at risk for barriers to TTS identification and treatment

	Pros	Cons
Recommend vaccination only for adults ≥50 years of age	 Would remove vaccine from most at-risk population (reduce TTS cases) Clear to communicate 	 Difficult to implement (vaccination sites could need stock two vaccines) Would remove an option in a population with lower risk (young men) Could disproportionately impact at-risk populations
Reaffirm recommendations for use; women <50 should be aware of increased risk, and may choose another COVID-19 vaccine	 Allow for flexibility/choice Allow for use of the vaccine in harder to reach populations, while still acknowledging risk in young women 	 Could be difficult to implement (vaccination sites could need stock two vaccines) Could be difficult to communicate

Policy Options for Janssen Policy Recommendations Work Group Summary

- Detailed discussion of risk/benefit balance difficult in many current vaccination settings
- Recommendations that require vaccination sites to require two types of vaccines would be difficult to implement
- Access to vaccines for hard-to-reach populations remains important
- Risk/benefit balance may change as the pandemic evolves and risk of COVID-19 disease changes

Policy Options for Janssen Policy RecommendationsWork Group Summary

- Work Group discussed benefits and concerns with all policy options for Janssen COVID-19 vaccine
- No single policy option as clear choice by Work Group
- However, many on Work Group appreciated flexibility of a broader recommendation for use, but acknowledgement of elevated risk in women <50 years of age

Previous Janssen vote:

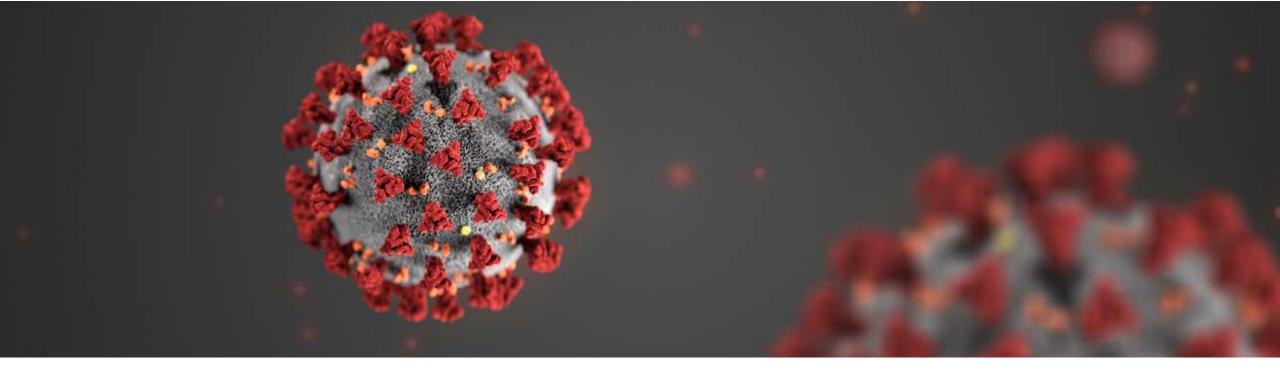
The Janssen COVID-19 vaccine is recommended for persons 18 years of age and older in the U.S. population under the FDA's Emergency Use Authorization

Question for ACIP to discuss:

• Given the review of the benefits and risks, what recommendation does ACIP feel is appropriate for use of the Janssen COVID-19 vaccine?

Policy Options for Janssen Policy Recommendations

- Recommend against use for all persons
- Reaffirm recommendations for all age and sex
 - FDA to include warning statement with EUA
- Recommend vaccination only for adults ≥50 years of age
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For more information, contact CDC 1-800-CDC-INFO (232-4636)
TTY: 1-888-232-6348 www.cdc.gov

Thank you

The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the Centers for Disease Control and Prevention.

