

# Global Health Cast 50 September 20, 2023







Prof. Dr. Joe Schmitt

@Prof\_Schmitt

# **Every Week**

12.00 noon - CET



## 50th Podcast

# **Check our GHP Website:**

www.ID-EA.org

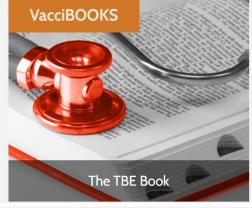






FDA / EMA / WHO

**VacciPROFILES** 



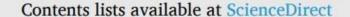


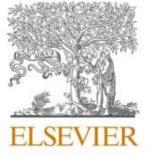




## What we talk about today

- ➤ Intelligent people get their COVID-19 vaccines first
- > Past COVID infection, side effects top reasons for not getting booster
- > 166 Legionellosis cases and 23 deaths reported in Poland
- ➤ Is your vaccination status up to date?
- ➤ Adverse events following COVID-mRNA dosing in subjects ≥65 years





### Journal of Health Economics

JOURNAL OF HEALTH ECONOMICS

journal homepage: www.elsevier.com/locate/jhealeco



# Cognitive ability, health policy, and the dynamics of COVID-19 vaccination

Mikael Elinder<sup>a</sup>, Oscar Erixson<sup>b,\*</sup>, Mattias Öhman<sup>b</sup>

# TL,DR: Smart people first in line for COVID-19 vaccines

#### ARTICLE INFO

Keywords: Intelligence Vaccination COVID-19 Nudge

Administrative data

Twin-design

#### ABSTRACT

We examine the relationship between cognitive ability and prompt COVID-19 vaccination using individual-level data on more than 700,000 individuals in Sweden. We find a strong positive association between cognitive ability and swift vaccination, which remains even after controlling for confounding variables with a twin-design. The results suggest that the complexity of the vaccination decision may make it difficult for individuals with lower cognitive abilities to understand the benefits of vaccination. Consistent with this, we show that simplifying the vaccination decision through pre-booked vaccination appointments alleviates almost all of the inequality in vaccination behavior.



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#### Vaccine

Available online 4 September 2023

In Press, Corrected Proof (?) What's this?





# Understanding low COVID-19 booster uptake among US adults

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#### **Conclusions**

Improvement in booster uptake is necessary for optimal public health in the United States. The development of vaccines against SARS-CoV-2 occurred at an unprecedented speed, but vaccine uptake remains among the greatest current public health challenges as updated boosters continue to be developed and made available to the public. Interventions to improve vaccination rates require a variety of approaches.



# Poland launches probe as Legionella confirmed in city water supply

Intelligence bosses are looking into whether the outbreak in Rzeszów is a result of tampering.





## Eleven people die from Legionnaires' disease outbreak in Polish town of Rzeszow





# Transmission of Legionella Bacteria Legionella aerosolize and are inhaled (or aspirated) from human-made or natural water systems

Susceptible person

can become infected

Legionella grow

in human-made water systems

## World Health Organization

"As of 11 September 2023, a total of 166 cases of legionellosis, including 23 deaths, have been reported from Poland."

Levofloxacin or azithromycin for 7–10 days is recommended in cases of moderate to severe Legionella pneumonia. For immunocompromised hosts, a 21-day course of levofloxacin or a 10-day course of azithromycin is usually recommended

https://www.ncbi.nlm.nih.gov/pmc/articles/PMC9124264/



Legionella found in the

natural environment

## **Current Vaccines**

Maternal Immunization (5)

Infant, toddlers (≥11+4)

School Entry/ Adolescents (6) Adults (>3)

≥65 yrs (9)

- TdaP
- Influenza
- COVID-19
- RSV

(Recommended, not licensed)

- DTaP-Hib-IPV-HBV
  - PCV
- Rotavirus
- Influenza
- MenACWY
- MenB
- TBE (>1 yr)
- MMR-V

- Boosters TdaP
- Influenza
- COVID-19
- MenACWY
- MenB
- HPV
- TBE

- Boosters
- Tdap-IPV
- Influenza
- HPV
- TBE
- "Workplace Vx"

- Boosters
- TdaP
- TBE
- COVID-19
- Influenza
- PCV
- Zoster
- MenACWY
- MenB













# Medical Need Based on sepcial host/exposure

#### **Traveler**

## **Boosters**

- TBE (FSME)
- HAV
- **HBV**
- Influenza
- MenACWY
- MenB
- JE
- **Typhoid**
- Rabies
- Yellow fever
- Cholera
- Dengue

### **Underlying** diseases

- Influenza
- Boosters
- (risk-based Vx)

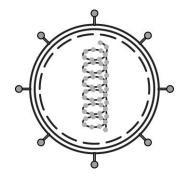






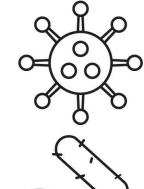
### **B-&T-cell Defect/** Cancer\*

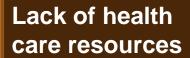
- Influenza
- PCV
- MenACWY
- MenB
- H. influenzae b
- **HBV**
- Boosters



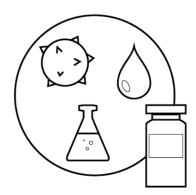
### **Emerging** infections

- COVID-19 VOCs
- Avian influenza
- Cholera
- Smallpox
- Monkeypox





Monkeypox Ebola





## Potential AE after COVID-19 mRNA vaccination in subjects >65 years and older: Two self-controlled studies in the U.S

*Background:* Our near-real-time safety monitoring of 16 adverse events (AEs) following COVID-19 mRNA vaccination identified potential elevation in risk for six AEs following primary series and monovalent booster dose administration. The crude association with AEs does not imply causality. Accordingly, we conducted robust evaluation of potential associations.

*Methods:* We conducted two self-controlled case series studies of COVID-19 mRNA vaccines (BNT162b2 and mRNA-1273) in U.S. Medicare beneficiaries aged  $\geq$  65 years. Adjusted incidence rate ratio (IRRs) and 95 % confidence intervals (CIs) were estimated following primary series doses for acute myocardial infarction (AMI), pulmonary embolism (PE), immune thrombocytopenia (ITP), disseminated intravascular coagulation (DIC); and following monovalent booster doses for AMI, PE, ITP, Bell's Palsy (BP) and Myocarditis/Pericarditis (Myo/Peri).

Results: The primary series study included 3,360,981 individuals who received 6,388,542 primary series doses; the booster study included 6,156,100 individuals with one monovalent booster dose. The AMI IRR following BNT162b2 primary series and booster was 1.04 (95 % CI: 0.91 to 1.18) and 1.06 (95 % CI: 1.003 to 1.12), respectively; for mRNA-1273 primary series and booster, 1.01 (95 % CI: 0.82 to 1.26) and 1.05 (95 % CI: 0.998 to 1.11), respectively. The hospital inpatient PE IRR following BNT162b2 primary series and booster was 1.19 (95 % CI: 1.03 to 1.38) and 0.86 (95 % CI: 0.78 to 0.95), respectively; for mRNA-1273 primary series and booster, 1.15 (95 % CI: 0.94 to 1.41) and 0.87 (95 % CI: 0.79 to 0.96), respectively. The studies' results do not support that exposure to COVID-19 mRNA vaccines elevate the risk of ITP, DIC, Myo/Peri, and BP.

Conclusion: We did not find an increased risk for AMI, ITP, DIC, BP, and Myo/Peri and there was not consistent evidence for PE after exposure to COVID-19 mRNA primary series or monovalent booster vaccines. These results support the favorable safety profile of COVID-19 mRNA vaccines administered in the U.S. elderly population.



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