REFERENCES - TOWN HALL 91 - 10-22-2023

A PAPER IN *Nature* demonstrated the utility of nasal swabs for assessing mucosal immune responses to SARS-CoV-2.

https://www.nature.com/articles/s41598-023-44989-5

A RANDOMIZED CONTROLLED TRIAL PUBLISHED IN *THE NEW ENGLAND JOURNAL OF MEDICINE* SHOWED THAT TREATMENT WITH INHALED FLUTICASONE FUROATE FOR 14 DAYS DID NOT RESULT IN A SHORTER TIME TO RECOVERY THAN PLACEBO AMONG OUTPATIENTS WITH COVID-19.

https://www.nejm.org/doi/full/10.1056/NEJMoa2209421?query=featured coronavirus

A JAMA NETWORK OPEN STUDY DEMONSTRATED THAT HIGH-FLOW NASAL OXYGEN AND NONINVASIVE VENTILATION APPEAR NOT TO BE AEROSOL-GENERATING PROCEDURES.

https://jamanetwork.com/journals/jamanetworkopen/fullarticle/2810485?guestAccessKey=7c3b5876-0b0d-4e3c-9934-

3dc5e8d276f9&utm_source=silverchair&utm_medium=email&utm_campaign=jama_network&utm_content=covid_weekly_highlights&adv=001602730367

A LARGE JAMA NETWORK OPEN RETROSPECTIVE COHORT STUDY FOUND THAR COVID-19 WAS ASSOCIATED WITH A SUBSTANTIAL RISK FOR AUTOIMMUNE AND AUTOINFLAMMATORY CONNECTIVE TISSUE DISORDERS.

https://jamanetwork.com/journals/jamanetworkopen/fullarticle/2810259?utm source=silverch air&utm medium=email&utm campaign=article alert-

jamanetworkopen&utm_content=wklyforyou&utm_term=100623?adv=001602730367

A JAMA Internal Medicine study of US Veterans found that COVID-19 survivors had no clinically significant excess hazard of death greater than comparators among those who survived at least 6 months after infection.

https://jamanetwork.com/journals/jamainternalmedicine/fullarticle/2808237?guestAccessKey=f00cadc8-f47b-4832-9a0e-

4b110f8fb089&utm source=silverchair&utm medium=email&utm campaign=jama network&ut m content=covid weekly highlights&adv=001602730367

A JAMA STUDY FOUND THAT THAT LONG-TERM SUPPORT FOR FAMILY MEMBERS OF ICU PATIENTS WITH COVID-19 ARDS SHOULD BE THE SAME AS FOR RELATIVES OF PATIENTS WITH OTHER CAUSES OF ARDS.

https://jamanetwork.com/journals/jama/fullarticle/2809191

A LANCET PUBLIC HEALTH PAPER PROVIDED A RETROSPECTIVE ASSESSMENT OF COVID SURVEILLANCE SYSTEMS USED IN ENGLAND DURING THE PANDEMIC, CONCLUDING THAT DEPLOYING A SUITE OF MONITORING SYSTEMS IS OPTIMAL.

https://www.thelancet.com/journals/lanpub/article/PIIS2468-2667(23)00219-0/fulltext

Another Lancet paper reported on the comparative effectiveness of nirmatrelvir/ritonavir versus sotrovimab for preventing severe COVID-19 outcomes in non-hospitalized high-risk patients during Omicron waves, finding approximately equivalent outcomes for both.

https://www.thelancet.com/journals/lanepe/article/PIIS2666-7762(23)00160-6/fulltext

An opinion piece in *The Journal of Infectious Diseases* presents evidence that so-called 'hybrid immunity' (i.e., vaccination plus infection) produces more robust immunity than either alone.

https://academic.oup.com/jid/advance-

article/doi/10.1093/infdis/jiad353/7245175?searchresult=1

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A PAPER IN *CLINICAL INFECTIOUS DISEASES* FOUND THAT, IN A HIGHLY IMMUNE ADULT POPULATION, MEDIAN SARS-COV-2 VIRAL LOADS PEAKED AROUND THE FOURTH DAY OF SYMPTOMS AND THAT INFLUENZA A VIRAL LOADS PEAKED SOON AFTER SYMPTOM ONSET.

https://academic.oup.com/cid/advance-article/doi/10.1093/cid/ciad582/7285011?searchresult=1

A SHORT *CLINICAL INFECTIOUS DISEASES* OPINION PIECE WRITTEN BY A FELLOW IN INFECTIOUS DISEASES DESCRIBES HIS GRIEVING PROCESS DURING THE PANDEMIC IN PART STIMULATED BY HIS STRUGGLES WITH CLOSE FAMILY MEMBERS WHO ROUTINELY REPORTED THEIR ANTI-SCIENCE AND CONSPIRACY THEORIES ABOUT **COVID**.

https://academic.oup.com/cid/advance-article/doi/10.1093/cid/ciad578/7295834

A PAPER IN *Infection Control and Hospital Epidemiology* found that an immediate, substantial, and sustained increase of healthcare-associated respiratory viral infections occurred after the institution discontinued universal masking.

https://www.cambridge.org/core/journals/infection-control-and-hospital-epidemiology/article/healthcareassociated-respiratory-viral-infections-after-discontinuing-universal-masking/E3B1E21AFB9D9BA4C535F7BB810A3D1C

THE *U.S. FOOD AND DRUG ADMINISTRATION* AMENDED THE EMERGENCY USE AUTHORIZATION (EUA) FOR THE NOVAVAX COVID-19 VACCINE, ADJUVANTED TO INCLUDE THE 2023-2024 FORMULA FOR ANYONE 12 YEARS OF AGE AND OLDER.

https://www.fda.gov/vaccines-blood-biologics/coronavirus-covid-19-cber-regulated-biologics/novavax-covid-19-vaccine-adjuvanted

THE NIAID IS BEGINNING A CLINICAL TRIAL OF A "UNIVERSAL" INFLUENZA VACCINE.

https://www.nih.gov/news-events/news-releases/nih-clinical-trial-universal-flu-vaccine-candidate-begins