#### Dr. Leo's "In the Literature"

- Low-dose aspirin increased incident anemia and decline in ferritin in otherwise healthy older adults compared to placebo, independent of major bleeding. 51 events per 1000 person-years compared to 42 events per 1000 person-years.
- Patients with culture-positive or culture-negative empyema who underwent thoracoscopic decortication showed similar short-term and long-term survival outcomes. however, culture positive patients did have longer ICU stays, longer vent usage and longer post-op stay in the hospital.
- 3. No improved clinical outcomes were noted with targeted carbapenem therapy for zosynnonsusceptible/ceftriaxone-susceptible infections
- 4. Cases and severe outcomes associated with SARS-CoV-2 reinfection have increased across the United States since September 2021. Delta wave reinfection rate was 2.7%, and increased to 28% during omicron wave. Reinfection rates leading to hospitalizations and death also increased from delta to omicron substantially.
- 5. In adults with obesity, retatrutide treatment for 48 weeks resulted in substantial reductions in body weight (average of >24% body weight loss.
- 6. RCT for oral semaglutide (Ozempic) 50 mg once per day led to a superior and clinically meaningful decrease in bodyweight compared with placebo

### Annals - 6/2023 - effect of low dose aspirin on incidence of anemia in the elderly.

- 19 114 persons were randomly assigned.
- Anemia incidence in the aspirin and placebo groups was 51.2 events and 42.9 events per 1000 person-years, respectively (hazard ratio, 1.20 [95% CI, 1.12 to 1.29]). Hemoglobin concentrations declined by 3.6 g/L per 5 years in the placebo group and the aspirin group experienced a steeper decline by 0.6 g/L per 5 years (CI, 0.3 to 1.0 g/L). In 7139 participants with ferritin measures at baseline and year 3, the aspirin group had greater prevalence than placebo of ferritin levels less than 45 μg/L at year 3 (465 [13%] vs. 350 [9.8%]) and greater overall decline in ferritin by 11.5% (CI, 9.3% to 13.7%) compared with placebo. A sensitivity analysis quantifying the effect of aspirin in the absence of major bleeding produced similar results
- Low-dose aspirin increased incident anemia and decline in ferritin in otherwise healthy older adults, independent of major bleeding. Periodic monitoring of hemoglobin should be considered in older persons on aspirin
- https://www.acpjournals.org/doi/10.7326/M23-0675

# OFID - 6/2023 - Culture positive and culture negative empyema after thoracoscopic decortication

- A total of 1087 patients with empyema received surgery, and 824 were enrolled after exclusion.
- Among these, <u>366 patients showed positive culture results</u> and <u>458 patients showed negative results</u>.
  - Longer intensive care unit stays (11.69 vs 5.64 days, P < .001),</li>
  - o longer ventilator usage (24.70 vs 14.01 days, P = .002), and
  - o longer postoperative hospital stays (40.83 vs 28.37 days, P < .001) were observed in the culture-positive group.

- However, there was no significant difference in 30-day mortality between the 2 groups (5.2% in culture negative vs 5.0% in culture positive, P = .913). The
- 2-year survival was not significantly different between the 2 groups (P = .236).
- Patients with culture-positive or culture-negative empyema who underwent thoracoscopic decortication showed similar short-term and long-term survival outcomes
- https://doi.org/10.1093/ofid/ofad227

OFID - treatment of zosyn-nonsusceptible/ceftriaxone susceptible infections with carbapenems (CG) vs carbapenem sparing (CSG) regimens.

- Of 1062 patients screened, 200 were included (CG, n = 51; CSG, n = 149). Baseline characteristics, including Charlson Comorbidity Index (CCI; median [interquartile range], 6 [3–9] vs 6 [4–9]; P = .704), were similar between groups, except for more immunocompromised CG patients (29% vs 11%, P = .001).
- The most common infection sources were urinary (31% vs 57%, *P* = .002) and bloodstream (18% vs 17%, *P* = .887). Eighty-eight percent of the CG received meropenem, while 58% of the CSG received ceftriaxone as targeted therapy.
- There was no statistical difference in the primary endpoint between overall groups (27% vs 17%, *P* = .123), nor when stratified by infection source.
- More patients in the CSG switched to oral therapy (15 [29%] vs 100 [67%], P < .001). In multivariate analysis, CCI was an independent predictor of the primary outcome (odds ratio [OR], 1.199 [95% confidence interval, 1.074–1.340]; P = .001), while treatment with carbapenem-sparing therapy was not</li>
- Our study did not find improved clinical outcomes with targeted carbapenem therapy for TZP-NS/CRO-S infections. Carbapenem-sparing agents may be considered to spare carbapenems in noncritically ill patients similar to those included in our cohort
- https://doi.org/10.1093/ofid/ofad262

## CDC - 6/2023 - Trends in COVID reinfections and hospitalizations between 9/2021 and 12/2022

- Cases and severe outcomes associated with SARS-CoV-2 reinfection have increased across the
  <u>United States since September 2021.</u> CDC recommends staying up to date with COVID-19
  vaccinations and receiving early antiviral treatment, if eligible, to reduce the risk for severe
  COVID-19—associated outcomes
- As a percentage of all infections, <u>reinfections increased substantially</u> from the
  - o <u>Delta (2.7%)</u> to the
  - o Omicron BQ.1/BQ.1.1 (28.8%) periods; during the same periods,
- increases in the percentages of reinfections among COVID-19—associated hospitalizations
  - o (from 1.9% [Delta] to
  - 17.0% [Omicron BQ.1/BQ.1.1]) and
- deaths
  - o (from 1.2% [Delta] to
  - o 12.3% [Omicron BQ.1/BQ.1.1]) were also substantial.
- Percentages of all COVID-19 cases, hospitalizations, and deaths that were reinfections were consistently higher across variant periods among adults aged 18–49 years compared with those among adults aged ≥50 years. The median interval between infections ranged from 269 to 411

days by week, with a steep decline at the start of the BA.4/BA.5 period, when >50% of reinfections occurred among persons previously infected during the Alpha variant period or later. To prevent severe COVID-19 outcomes, including those following reinfection, CDC recommends staying up to date with COVID-19 vaccination and receiving timely antiviral treatments, when eligible.

https://www.cdc.gov/mmwr/volumes/72/wr/mm7225a3.htm?s cid=mm7225a3 x

### NEJM - 6/2023 - retratrutide for obesity - phase 2 trial

- Retatrutide (LY3437943) is an <u>agonist of the glucose-dependent insulinotropic polypeptide</u>, <u>glucagon-like peptide 1, and glucagon receptors</u>
- We enrolled 338 adults, 51.8% of whom were men. The least-squares mean percentage change in body weight at 24 weeks in the retatrutide groups was −7.2% in the 1-mg group, −12.9% in the combined 4-mg group, −17.3% in the combined 8-mg group, and −17.5% in the 12-mg group, as compared with −1.6% in the placebo group.
- At 48 weeks, the least-squares mean percentage change in the retatrutide groups was -8.7% in the 1-mg group, -17.1% in the combined 4-mg group, -22.8% in the combined 8-mg group, and -24.2% in the 12-mg group, as compared with -2.1% in the placebo group.
- At 48 weeks, a weight reduction of 5% or more, 10% or more, and 15% or more had occurred in 92%, 75%, and 60%, respectively, of the participants who received 4 mg of retatrutide; 100%, 91%, and 75% of those who received 8 mg; 100%, 93%, and 83% of those who received 12 mg; and 27%, 9%, and 2% of those who received placebo.
- The most common adverse events in the retatrutide groups were gastrointestinal; these events were dose-related, were mostly mild to moderate in severity, and were partially mitigated with a lower starting dose (2 mg vs. 4 mg). Dose-dependent increases in heart rate peaked at 24 weeks and declined thereafter
- In adults with obesity, retatrutide treatment for 48 weeks resulted in substantial reductions in body weight
- https://www.nejm.org/doi/full/10.1056/NEJMoa2301972?query=featured home

#### LANCET - 6/2023 - oral ozempic RCT

- The estimated mean bodyweight change from baseline to week 68 was −15·1% (SE 0·5) with oral semaglutide 50 mg versus −2·4% (0·5) with placebo (estimated treatment difference −12·7 percentage points, 95% CI −14·2 to −11·3; p<0·0001)
- In adults with overweight or obesity without type 2 diabetes, oral semaglutide 50 mg once per day led to a superior and clinically meaningful decrease in bodyweight compared with placebo
- https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(23)01185-6/fulltext