# Out-of-home child care as a risk factor for pre-pandemic endemic human coronavirus infections in a birth cohort of children 0-2 years

Ardythe L. Morrow<sup>1,2</sup> S.C. Conrey<sup>1,2</sup>, L. Niu<sup>1</sup>, A.R. Burrell<sup>1,2</sup>, C.P. Mattison<sup>3</sup>, Z. Teoh<sup>1,2</sup>, M. McMorrow<sup>3</sup>, M. McNeal<sup>2</sup>, D.C. Payne<sup>3</sup>, M.A. Staat<sup>1,2</sup>

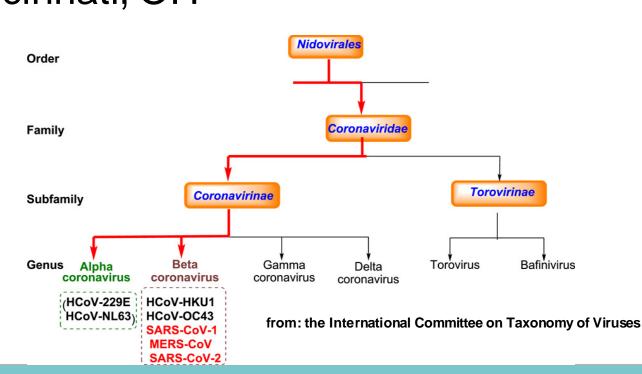
<sup>1</sup>University of Cincinnati College of Medicine, <sup>2</sup>Cincinnati Children's Hospital Medical Center, <sup>3</sup>The Centers for Disease Control and Prevention

Email: morrowa@ucmail.uc.edu



- Four endemic human coronaviruses (HCoV) HKU1, OC43, 229E, NL63 - belong to Coronaviridae, the viral subfamily of SARS-CoV-2. These occur seasonally, routinely circulate worldwide, and typically cause only mild illness
- The COVID-19 pandemic has increased interest in understanding these HCoV for comparison to pandemic coronaviruses
- Group or out of home child care has been a concern during the pandemic as a potential reservoir of infection in the community
- Here we examine the strength of association in out-ofhome child care and other potential risk factors for HCoV infection among children enrolled in PREVAIL, a CDCfunded birth cohort study of healthy, term singleton infants followed in Cincinnati, OH

Figure 1: Taxonomy of Coronaviridae



## Methods

## Data Collection (April 2017 - October 2020)

- Prenatal (3<sup>rd</sup> trimester) enrollment visit
- Weekly postnatal nasal swabs from study children collected, tested by Luminex Respiratory Pathogen Panel
- Risk factor data collected during quarterly study surveys
- Analysis restricted to 91 of 245 children who were the most adherent (≥ 75% of weekly swab collection)

## <u>Definitions</u>

- **HCoV infectious episode** any positive swabs within 30-day window of previous positive
- Out of home child care any care arrangement that does not occur in the child's own home.
- Child care home providers care for small groups of children in a residence that is not the child's
- Child care center group child care in a non-residential setting (licensed child care center)

#### Statistical Analysis

- Anderson-Gill model: risk of any HCoV infection
- Covariates: prior infection, childcare, child sex, race, insurance type, maternal education, maternal age, number of persons in the household, breastfeeding status
- All analysis performed using the R Environment for Statistical Computing<sup>2</sup>

## Figure 2: Demographics of 91 PREVAIL highly adherent children studied

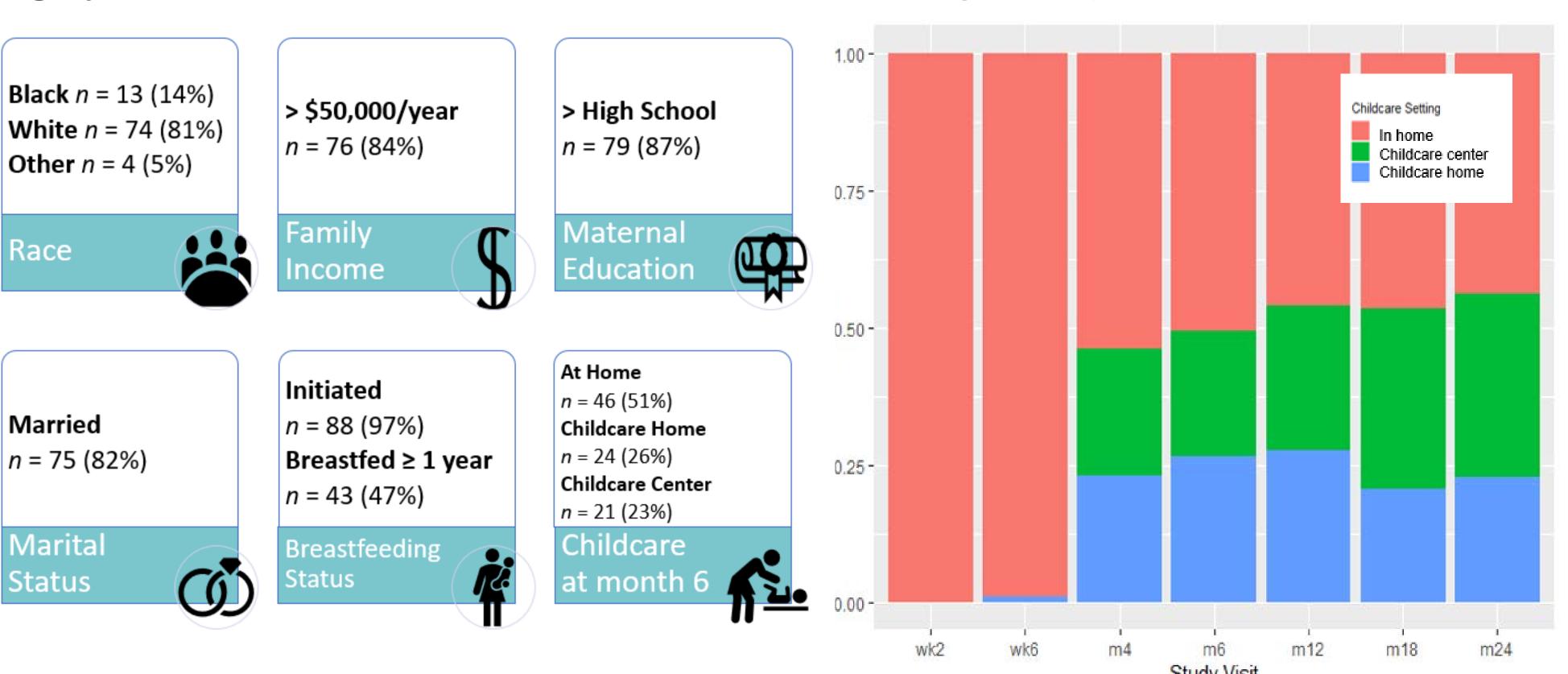


Figure 3: Out-of-home childcare use by

child age/study visit

Figure 4: Prevalence map of HCoV infection in PREVAIL

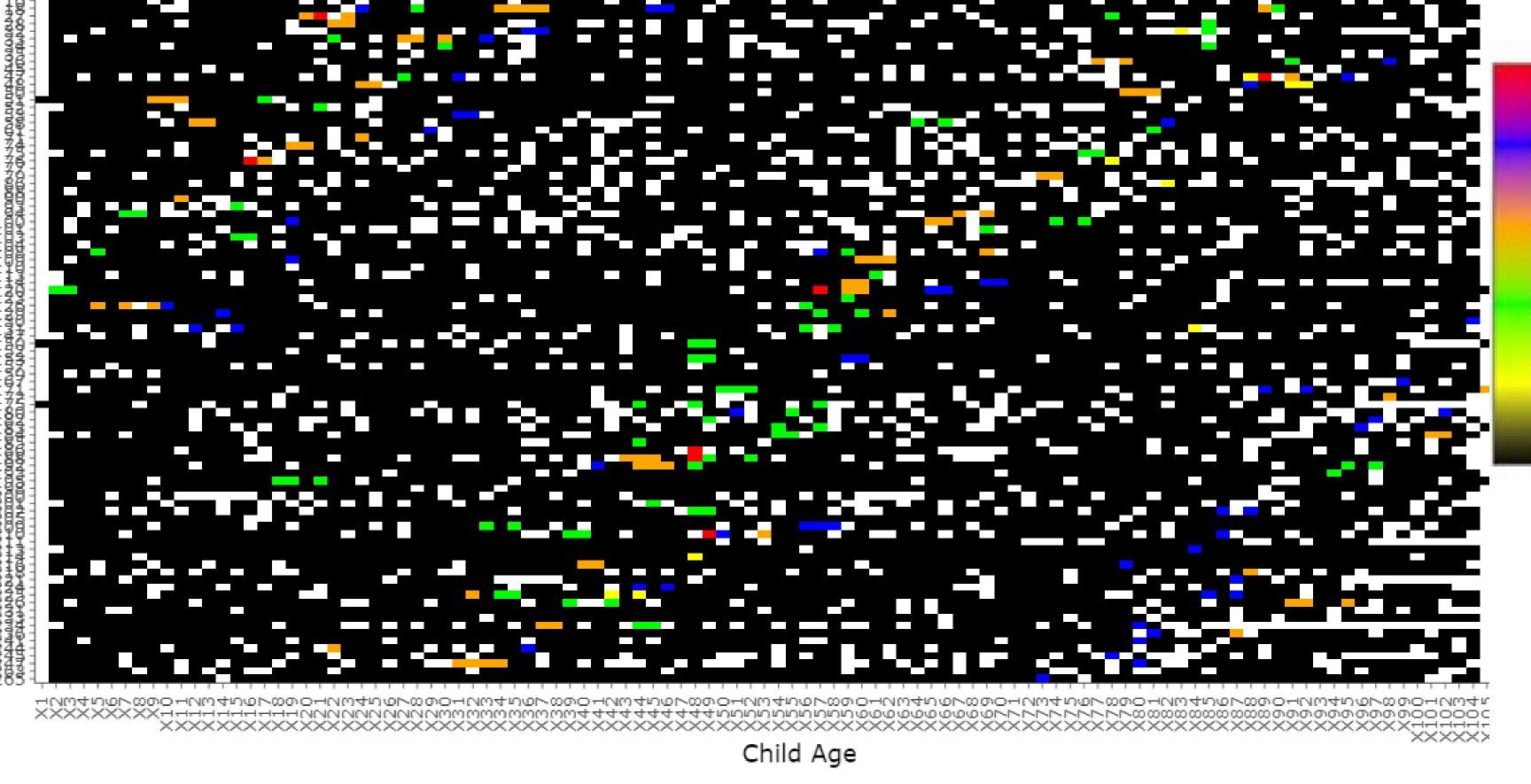
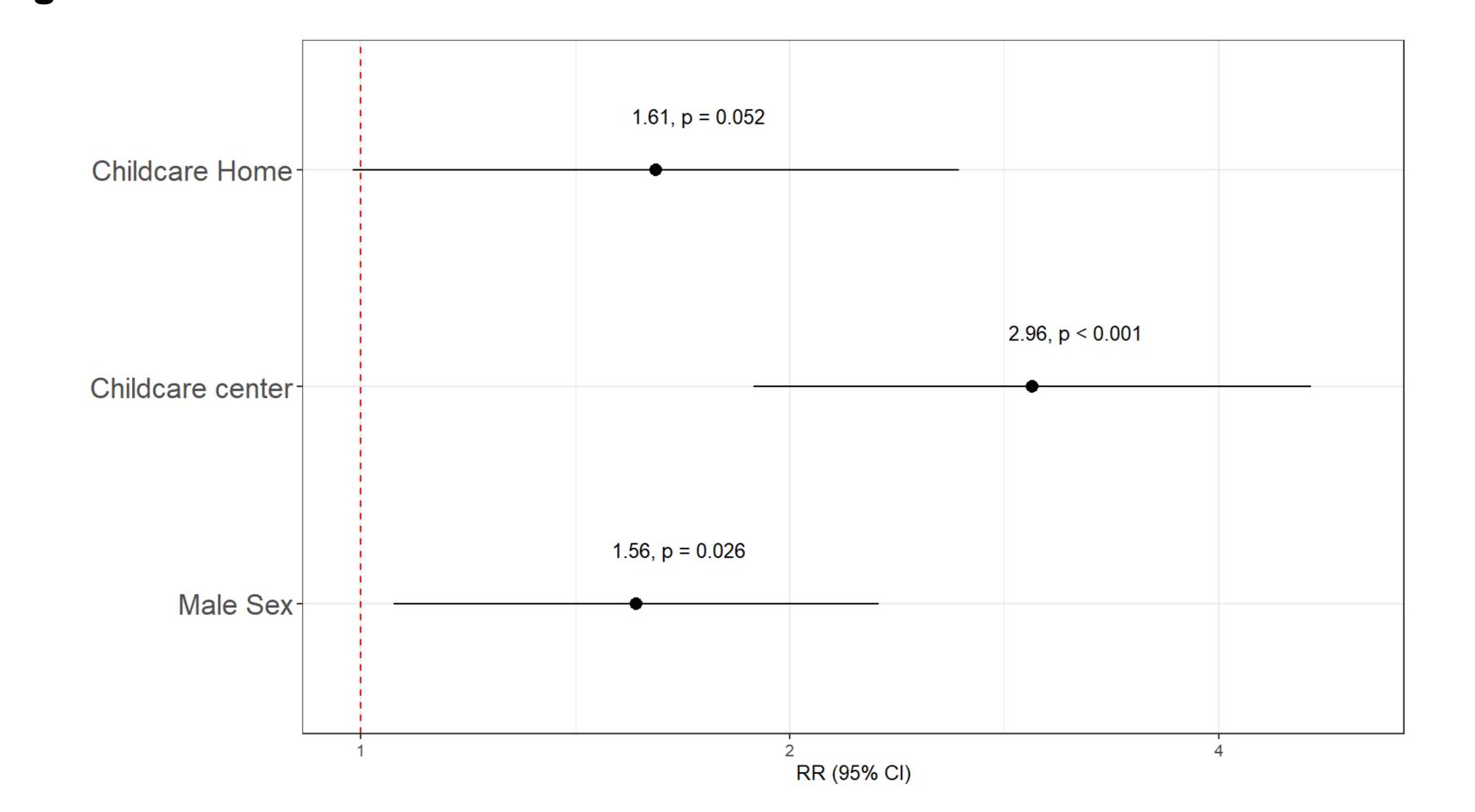


Figure 5: Anderson Gill model: Risk estimates for HCoV infection







## Results

- Demographics in line with Greater Cincinnati region in terms of race, income; higher than region maternal education, married status<sup>3</sup>, and breastfeeding initiation
- By 6 months, ½ of children were in out-of-home childcare
- Use of childcare homes & childcare centers evenly split
- Peer group size was > in childcare centers than child care homes (med. = 9 *vs* 4/group; *p*<0.001)
- HCoV prevalence in first two years of life
- 131 HCoV infectious episodes detected
  - 70 (77%) children had ≥1 HCoV infection
  - 40 (44%) had ≥ 2 HCoV infections
- Anderson Gill survival model:
- Use of childcare centers (RR 3.0, 95%Cl 1.9, 4.6) and male sex (1.6, 95%Cl 1.1, 2.3) increased risk for HCoV infection
- Use of childcare homes borderline significant (RR 1.6, 95%CI 1.0, 2.6)
- Not significant: race, insurance type, maternal education, maternal age, number of persons in the household, breastfeeding status

## Conclusions

HCoV HKU1

HCoV NL63

HCoV OC43

HCoV 229E

No infection

- HCoV infections were highly prevalent for all children
- The majority of children used out-of-home childcare after 6 months of age
- Relative risk of HCoV was higher in both child care homes and centers.
- While confidence bands overlap between these two child care settings and home care, the trend suggested < risk for child care homes (RR=1.6) compared to child care centers (RR=3.0), consistent with differences in peer group size
- Our findings are limited to endemic HCoV, but may be relevant to pandemic HCoV as well

## **Strengths**

- Ability to assess time-varying and stable risk factors
- High sample adherence = ability to detect infections

#### Limitations

- Small sample size, lack of SARS-CoV2 for comparison
- Compliance restricted diversity, higher SES than region

## **Future Directions**

- Add serologic-identified infections
- Compare symptoms, severity

## References

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