- Relative role of community transmission and campus contagion in
- ³ driving the spread of SARS-CoV-2: lessons from Princeton
- 4 University
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17 Abstract

- 18 Mathematical models have played a crucial role in guiding pandemic responses. University
- 19 campuses present a particularly well-documented case for institutional outbreaks, thereby

providing a unique opportunity to understand detailed patterns of pathogen spread. Here,
we present descriptive and modeling analyses of SARS-CoV-2 transmission on the Princeton
University campus—this model was used throughout the pandemic to inform policy decisions
and operational guidelines for the university campus. We demonstrate strong spatiotemporal
correlations in epidemic patterns between the university campus and surrounding communities. These findings are corroborated by our model predictions, which indicate that the
amount of on-campus transmission was likely limited during much of the wider pandemic
until the end of 2021. Finally, we find that a super-spreading event likely played a major
role in driving the recent Omicron variant outbreak on the Princeton University campus
during the spring semester of the 2021–2022 academic year. Despite large numbers of cases
on campus in this period, case levels in surrounding communities remained low, suggesting
that there was little spillover transmission from campus to the local community.

32 Significance

33 Introduction

Predicting and controlling the spread of SARS-CoV-2 has remained a critical public health and scientific question throughout the ongoing SARS-CoV-2 pandemic (Baker et al., 2021). Rapid, asymptomatic transmission of SARS-CoV-2 has hindered intervention efforts, such as contact tracing (Hellewell et al., 2020). Social distancing measures have played major 37 roles in preventing transmission, but can be difficult to maintain for a prolonged period (Galanti et al., 2021). The development of vaccines has provided a safe means of reopening society, but uncertainty remains on their long-term effectiveness in preventing infection and 40 transmission, especially in the face of new viral variants. Mathematical models have played a significant role in guiding these pandemic responses and devising control strategies (Cobey, 2020; Holmdahl and Buckee, 2020; Metcalf et al., 2020; Koelle et al., 2022). Models can help monitor key parameters that govern epidemic dynamics (Kraemer et al., 2021) and retrospectively estimate the impact of intervention measures in reducing transmission (Flaxman et al., 2020). These estimates can further inform projections of future scenarios and allow us to explore the endemicity of SARS-CoV-2 (Kissler et al., 2020; Saad-Roy et al., 2020; Lavine et al., 2021; Saad-Roy et al., 2021). Mathematical models have also been widely deployed in planning campus reopenings. 49 Researchers from various institutions in the US—including Cornell (Frazier et al., 2022), Emory (Lopman et al., 2020), Georgia Institute of Technology (Gibson et al., 2021), and UC Berkeley (Brook et al., 2021)—modeled the feasibility of controlling the epidemic on their campuses and considered mass asymptomatic testing as their main intervention. These modeling efforts helped identify key parameters for control, such as the testing turnaround time, and provided support for implementing similar measures at other institutions. Coupling modeling efforts with real-life implementations in university campuses further provided unique opportunities to directly test model-based predictions of intervention effects in preventing the transmission of SARS-CoV-2 (Frazier et al., 2022)— each university campus offers a relatively well-controlled epidemic setting with a relatively homogeneously behaving population (especially among undergraduate students). Campuses can also offer strong opportunities for control by non-pharmaceutical interventions, such as isolation and mask-wearing; mass asymptomatic testing further provides robust ascertainment for epidemic sizes, allowing for accurate understanding of epidemic patterns.

On the other hand, university campuses also present unique challenges to controlling
an outbreak. A large fraction of asymptomatic infections (due to the young age of university students) and high-density interactions—such as eating in large dining halls and
various social activities—can readily permit rapid transmission. These kinds of contacts
are inherently difficult to keep track of, making contact tracing less effective. The impact
of intervention measures is expected to vary across different university campuses, reflecting
heterogeneity in campus settings such as compliance, resources, community prevalence, as
well as effects of other interventions present on campuses. For example, Duke and Harvard
Universities experienced moderate outbreaks at the beginning of the fall semester in 2021
when in-person classes were allowed, despite high vaccination rates and weekly asymptomatic
testing protocols (Duke University, 2021; Harvard University, 2021), whereas the number of
cases remained low in Princeton University (PU) during the same time period with similar

levels of testing and vaccination. Here, we focused on the dynamics of SARS-CoV-2 on the
PU campus alone to eliminate heterogeneities inherent to such comparisons; we return to
comparisons with other campuses later in the discussion.

We begin with a descriptive analysis of the PU outbreak (Fig. 1), and present modeling analyses of the individual epidemics during 2020-2022. PU is located in Mercer County, New Jersey, USA; the population comprises of 5267 undergraduate students, 2946 graduate students, and around 7000 faculty and staff members. For simplicity, we divided the epidemic into four time periods representing four semesters across two academic years: Fall 2020–2021 (August 24, 2020–January 1, 2021; Fig. 1A), Spring 2020–2021 (January 16, 2021–May 14, 2021; Fig. 1B), Fall 2021–2022 (August 14, 2021–December 31, 2021; Fig. 1C), and Spring 2021–2022 (January 1, 2022–March 18, 2022; Fig. 1D). Throughout the majority of the study period, all students, faculty and staff members who were physically present for more than 8 hours on campus per week were required to participate in asymptomatic testing with varying frequencies. Asymptomatic individuals submitted self-collected saliva samples, from which the presence of SARS-CoV-2 was tested using Reverse Transcription Polymerase Chain Reaction (RT-PCR). Those who tested positive were required to isolate for at least 10 days after symptom onset or test date (whichever was longer) and were released when they had been at least 48 hours with improving or resolving symptoms as per New Jersey Department of Health guidance. PCR positives were exempt from asymptomatic testing for 90 days. Since March 7, 2022, asymptomatic testing frequencies decreased to once a month from once a week for individuals whose vaccine status is up-to-date. This in turn likely reduced the accuracy of surveillance; therefore, we chose to focus on epidemic patterns before this change was implemented. Throughout the study period, contact tracing was also performed for positive cases to alert their close contacts to either quarantine or test more frequently according to the close contacts' vaccination status, and to gather data that could help uncover clusters of transmission or superspreader events. Changes in testing frequency and other intervention measures throughout the study period reflected various factors, including the impact of COVID-19 cases on continuity of operations or continuity of teaching; on severity of disease on campus; the capacity of testing and the healthcare system; and hospitalization rates on campus and in the area. All data used in this analysis are publicly available on the PU COVID-19 Dashboard website: https://covid.princeton.edu/dashboard.

77 Descriptive analysis

During the fall semester of the 2020–2021 academic year, roughly 1000 grad students 108 and 2000 faculty and staff members were present on campus and participated in asymp-100 tomatic testing. All classes were held virtually, and so only a few undergraduate students 110 remained on campus (< 300). Both undergraduate and graduate students were required 111 to get tested twice a week, whereas faculty and staff members were required to get tested 112 once a week. The number of cases remained relatively low throughout the semester with a 113 peak occurring in early December, coinciding with the epidemic trajectory in Mercer County 114 (Fig. 1A). A sudden decrease in the number of cases around Thanksgiving partly reflects the 115 reduced number of tests (3852 and 2972 asymptomatic tests performed on the week ending November 20th and 27th, respectively). The highest number of cases was reported among

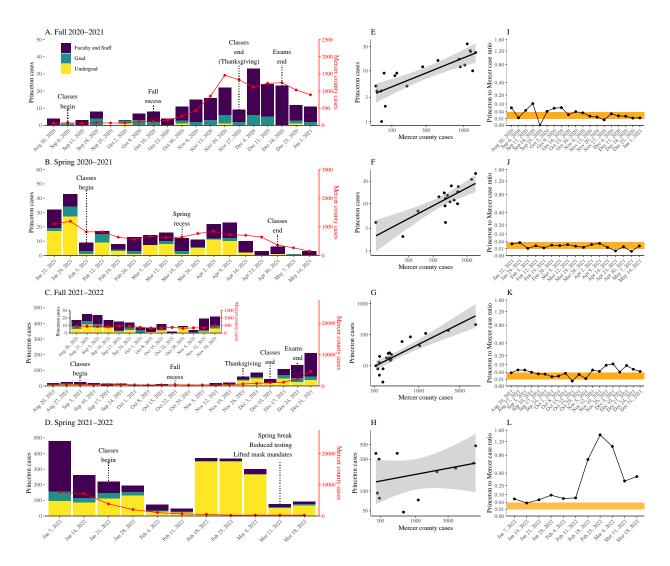


Figure 1: Dynamics of SARS-CoV-2 outbreaks in Princeton University (PU). (A–D) Epidemic trajectories across four semesters: Fall 2020–2021 (A), Spring 2020–2021 (B), Fall 2021–2022 (C), and Spring 2021–2022 (D). Colored bar plots represent the weekly number of cases from both asymptomatic and symptomatic testing in PU. Red lines represent the weekly number of cases in Mercer County. Number of cases in Mercer County is obtained from https://github.com/nytimes/covid-19-data. (E–H) Correlations between the weekly number of cases in PU and in Mercer County. Solid lines and shaded areas represent the estimated linear regression lines and the associated 95% CIs. (I–L) Ratios between weekly numbers of cases reported from PU and Mercer County. The shaded orange area covers the region between 1:100 and 1:25 ratios.

faculty and staff members (= 169), followed by graduate students (= 41), and undergraduate students (= 4).

In the beginning of the spring semester of the 2020–2021 academic year, the 120 number of cases suddenly increased before classes started (Fig. 1B), reflecting ≈ 3000 undergraduate students returning to campus. Returning students were required to be tested and quarantine for 7 days regardless of their returning location. Most classes remained vir-123 tual, and the testing protocol did not change (twice a week for undergraduate and graduate 124 students, and once a week for faculty and staff members). Some smaller classes were held 125 in-person, but required social distancing (thereby limiting the size of the class) and mask-126 wearing at all times. The number of cases persisted at similar levels to the fall semester and 127 eventually decreased as classes ended and students went home—the decrease in the number 128 of cases in PU also coincided with the decrease in the number of cases in Mercer County. The 129 highest number of cases was reported among faculty and staff members (=111), followed by 130 undergraduate students (=101), and graduate students (=29). 131

For the fall semester of the 2021–2022 academic year, all students and faculty 132 and staff members were required to be vaccinated, with very few medical and religious 133 exemptions. By the beginning of the semesters, 97% of undergraduate students, 96% of 134 graduate students, and 94% of faculty and staff members were vaccinated. Vaccinees were 135 required to be tested once a week, while unvaccinated individuals were required to be tested 136 twice a week. In-person classes and social events fully resumed on campus, though all 137 individuals were required to wear masks indoors with a few exceptions (e.g., when eating or drinking, or when teaching a small class). The number of cases remained similar to previous semesters until November when cases began to increase, primarily among undergraduate students around Thanksgiving (Fig. 1C). In order to prevent transmission, testing frequency was increased to twice a week for undergraduate students on November 27th, 2021; the size
of non-academic gatherings were also limited to 20 people. The number of cases decreased
slightly as classes ended but soon increased again as the Omicron (BA.1) variant began to
spread on campus and in Mercer county.

For the spring semester of the 2021–2022 academic year, all eligible students and 146 faculty and staff members were required to be boosted. By the beginning of the semesters, 65% of undergraduate students, 71% of graduate students, and 82% of faculty and staff 148 members were boosted. Undergraduate students were still required to be tested twice a week 149 to prevent the additional spread of the Omicron variant. The number of cases remained high 150 before classes began but decreased over time, following epidemic patterns in Mercer County 151 (Fig. 1D). Coinciding with the decrease in the campus and local cases numbers, the gathering 152 policy was updated to allow food in events were no longer limited to 20 people; in addition, 153 the testing frequency was reduced to once a week. Following the policy change, a large 154 outbreak occurred on campus with high case numbers persisting until Spring Break (March 155 5th, 2022). The timing of this outbreak also coincided with a rapid turnover of the Omicron 156 subvariant BA.2—the proportion of BA.2 subvariant reached 93.5% (372/398) compared to 157 26.9% (14/52) from the previous week. On March 7, 2022, mask mandates were lifted and 158 testing frequency was reduced to once a month.

60 Comparisons of campus and community transmission

Across the first three semesters, we find strong and significant correlations between the weekly logged numbers of cases from PU and those from Mercer county: fall 2020–2021

 $(\rho = 0.81 \ (95\% \ \text{CI}: 0.55-0.92; \ p < 0.001), \ \text{Fig. 1E}); \ \text{spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ (95\% \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{Spring } 2020-2021 \ (\rho = 0.80 \ \text{CI}: 0.001); \ \text{$ 0.51-0.92; p < 0.001), Fig. 1F); and fall 2021–2022 ($\rho = 0.86$ (95% CI: 0.67–0.94; p < 0.001), 164 Fig. 1G). These correlations are robust even when we stratify cases by the population, except for undergraduate students during Fall 2020, when most were not physically present on campus (Supplementary Figure S1). However, the case patterns in PU were decoupled from those in Mercer county for the spring semester of the 2021–2022 academic year with unclear correlations (Fig. 1H): $\rho = 0.32$ (95% CI: -0.34–0.77; p = 0.33). Stratifying cases by 169 subpopulation shows that case patterns in graduate students and faculty and staff members 170 were still strongly correlated with case patterns in Mercer county, meaning that the campus 171 transmission was limited to undergraduate students (Supplementary Figure S1). We also find 172 strong correlations between the weekly logged numbers of cases from PU and those from 173 other counties in New Jersey (Supplementary Figure S2)—these correlations significantly 174 decreased with distance from Mercer county in both spring 2020–2021 ($\rho = -0.48$ (95% CI: 175 -0.75-0.06; p = 0.03) for all cases and $\rho = -0.51$ (95% CI: -0.77-0.10; p = 0.02) for 176 faculty and staff cases) and fall 2021–2022 ($\rho = -0.68$ (95% CI: $-0.86--0.35, \, p < 0.001$) 177 for all cases and $\rho = -0.74$ (95% CI: -0.89-0.46, p < 0.001) for faculty and staff cases). 178 Across the first three semesters, both the total cases and faculty and staff cases showed 179 similar levels of correlations with local cases. For the spring semester of the 2021–2022 180 academic year, we still find high correlations between faculty and staff cases and local cases 181 throughout other countries ($\rho > 0.8$ across all counties in New Jersey); however, the total 182 cases exhibit considerably weaker correlations due to student-to-student transmission on 183 campus (Supplementary Figure S2).

These correlations likely reflect commuting and contact patterns, and therefore we expect 185 SARS-CoV-2 dynamics on campus to be correlated with those from nearby large cities as 186 well. We find similarly strong correlations with New York City for the first three semesters: fall 2020–2021 ($\rho = 0.74$ (95% CI: 0.44–0.90; p < 0.001), Supplementary Figure S3A); spring 2020-2021 ($\rho = 0.84$ (95% CI: 0.61-0.94; p < 0.001), Supplementary Figure S3B); fall 2021-189 2022 ($\rho = 0.78$ (95% CI: 0.51–0.91; p < 0.001), Supplementary Figure S3C); and spring 2021-2022 ($\rho = 0.33$ (95% CI: -0.34-0.78; p = 0.3), Supplementary Figure S3D). 191 The same picture emerges for Philadelphia except for spring 2020: fall 2020–2021 ($\rho =$ 192 0.83 (95% CI: 0.59-0.93; p < 0.001), Supplementary Figure S4A); spring 2020-2021 ($\rho = 0.29$ 193 $(95\% \text{ CI: } -0.22 - 0.68; p = 0.25), \text{ Supplementary Figure S4B}); \text{ fall } 2021 - 2022 \ (\rho = 0.78 \ (95\% \text{ CI: } -0.22 - 0.68; p = 0.25))$ 194 CI: 0.51-0.91; p < 0.001), Supplementary Figure S4C); and spring 2021-2022 ($\rho = 0.37$ 195 (95% CI: -0.30-0.80; p = 0.27), Supplementary Figure S4D). Including counties from New 196 York and Pennsylvania states into the spatial correlation analysis yields additional insights 197 (Supplementary Figure S5): epidemic dynamics were highly synchronized across all counties 198 in fall 2020–2021 and became less synchronized over time. These correlations significantly 199 decreased with distance in spring 2020–2021 ($\rho = -0.36$ (95% CI: -0.49-0.21; p < 0.001)) 200 and fall 2021–2022 ($\rho = -0.58$ (95% CI: -0.68--0.46; p < 0.001)). These variations likely 201 reflect differences in vaccination levels and the timing of the introduction of the Omicron 202 variant. 203 Finally, mass testing allows us to infer the ratio between the weekly numbers of cases from 204 Princeton and those from Mercer county—we expect this ratio to remain constant over time if the majority of infections on campus is caused by community transmission, provided that

testing patterns remain roughly constant in both places. We find that the ratio between the weekly numbers of cases from Princeton and those from Mercer county stayed between 1:100 208 and 1:25 until the end of the fall semester of the 2021–2022 academic year (Fig. 1I-K). An increase in this ratio at the end of November 2021 was associated with the campus outbreak before Thanksgiving followed by an introduction of the Omicron variant in December—this 211 deviation indicates an increase in the amount of transmission on campus. During the spring 212 semester of the 2021–2022 academic year, the ratio between PU cases and Mercer county 213 cases increased above one, meaning that more cases were reported in PU than elsewhere in 214 Mercer county (Fig. 1L); notably, we did not see an increase in Mercer county cases, meaning 215 that there was little-to-no transmission from campus to local community.

217 Mathematical modeling of past outbreaks

We use a discrete-time, individual-based model to simulate the spread of SARS-CoV-2 on
the PU campus. This model was initially developed and used throughout the pandemic
to inform policy decisions in PU, including the frequency of asymptomatic tests and the
number of isolation beds required. We continuously updated the model to reflect changes in
school settings (e.g., students returning back to campus after a virtual semester) as well as
intervention measures (e.g., vaccination in fall 2021 and booster shots with the emergence of
the Omicron variant). Here, we present a generic and parsimonious version that encompasses
sufficient details to characterize the overall spread of SARS-CoV-2 in PU without an overproliferation of parameters. The model consists of four main components simulated on a daily

time scale: (1) infection and transmission dynamics, (2) sampling and testing protocols, (3) isolation protocols, and (4) vaccination dynamics, including waning immunity and booster shots. Previous versions of the model included contact tracing, but we exclude it in this model for simplicity.

Infection processes are modeled based on standard compartmental structures (Supple-231 mentary Figure S6). Once infected, susceptible individuals remain in the exposed stage for $D_e = 2$ days on average, during which they cannot transmit or test positive. Exposed individ-233 uals then enter the presymptomatic stage, during which they can test positive and transmit 234 infections for $D_p = 3$ days on average. Presymptomatic individuals can then either remain 235 asymptomatic with probability $p_a = 0.4$ or develop symptoms with the remaining probabil-236 ity of $1 - p_a = 0.6$; both asymptomatic and symptomatic individuals are assumed to have 237 the same duration of infectiousness ($D_s = 3$) and equal transmission rates. Recovered indi-238 viduals are assumed to be immune to reinfections throughout a semester. Presymptomatic, 239 symptomatic, and asymptomatic infection stages are further divided into two subcompart-240 ments to allow for more realistic and narrower distributions than the exponential distribution 241 (Brett and Rohani, 2020). Transitions between each (sub)compartments are modeled using 242 a Bernoulli process with probabilities that match the assumed means (He et al., 2010): more 243 specifically, transition probabilities are equal to $1 - \exp(-\delta_x)$, where $\delta_x = -log(1 - n/D_x)$ represent the transition rate from stage X and n represents the number of subcompartments. 245 Assumed parameters are broadly consistent with other models of SARS-CoV-2 (Brett and Rohani, 2020; Lavezzo et al., 2020).

Transmission processes are modeled by first setting the contact reproduction number

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 $\mathcal{R}_{\text{contact}}$, which we define as the average number of infectious contacts an infected individual would make throughout the course of their infection; here, infectious contacts refer to 250 contacts that would result in infection when the contacted individual is susceptible to infection. The contact reproduction number implicitly accounts for all intervention measures that we do not model explicitly, such as social distancing and contact tracing—in other 253 words, this contact reproduction number does not account for asymptomatic testing or vaccination, which are modeled separately. We further decompose $\mathcal{R}_{\text{contact}}$ into pre-symptomatic 255 $\mathcal{R}_p = \beta_p D_p$ and (a) symptomatic $\mathcal{R}_s = \beta_s D_s$ reproduction numbers, where β_p and β_s represent the corresponding infectious contact rates during pre-symptomatic and (a)symptomatic 257 stages, respectively. Pre-symptomatic and (a)symptomatic reproduction numbers are calcu-258 lated based on the assumed value of the proportion of presymptomatic transmission $p_p = 0.5$: 259 $\mathcal{R}_p/\mathcal{R}_s = p_p/(1-p_p)$. On each day, all infected individuals who have not yet been isolated 260 then make infectious contacts at random to anyone on campus; the number of infectious 261 contacts are drawn from a negative binomial distribution with a mean of either β_p or β_s 262 and an overdispersion parameter of k = 0.1 to account for the possibility of super-spreading 263 events (Endo et al., 2020). We also rely on cases from Mercer County to crudely capture 264 community dynamics. In particular, we assume that infectious contacts from local or re-265 gional community can be made at random to anyone on campus; these contacts are modeled 266 using a Poisson distribution with a time-varying mean, which is calculated by scaling the 267 daily number of cases by θ and shifting it by 1 week to account for reporting delays. Infectious contacts, whether made by individuals on campus or from outside, result in infection only when the contacted individuals are susceptible; when the contacted individuals are vac-270

cinated, and therefore partially susceptible to infection, they have a reduced probability of infection corresponding to their susceptibility (discussed later).

All individuals on campus are assumed to follow a pre-determined asymptomatic testing 273 plan at a fixed frequency—for example, under weekly testing, one individual can get sampled on days 1, 8, 15, and so forth, while another individual get sampled on days 2, 9, 16, 275 and so forth. We assume that test results come back after one day. Symptomatic individuals can choose to take rapid PCR tests (with results returning on the same day) with a 277 given probability on each day until their symptoms resolve—this probability is set to 1 for 278 simulations presented in the main text. We further assume that symptomatic individuals 270 are isolated immediately when they submit their samples until they receive negative results. 280 All individuals who test positive are required to isolate (following the same isolation rule as 281 described earlier) and are exempt from asymptomatic testing for 90 days. Isolated individ-282 uals are assumed to no longer transmit infections. We assume that PCR tests can detect 283 infections from individuals who are in pre-symptomatic, symptomatic, and asymptomatic 284 stages with 95% sensitivity and 100% specificity. 285

As most students, as well as faculty and staff members, had received two doses of vaccination in the beginning of fall 2021, we do not distinguish the first and second doses. Instead,
we assume that all vaccinated individuals have 90% reduced susceptibility and 20% reduced
transmissibility at the beginning of the semester—these assumptions are consistent with recent estimates by Prunas et al. (2022) that vaccination with BNT162b2 reduces susceptibility by 89.4% (95% CI: 88.7%–90.0%) and infectiousness by 23.0% (95% CI: -11.3%-46.7%)
against the Delta strain. Based on Tartof et al. (2021), vaccine efficacy against susceptibility

is allowed to exponentially wane from 90% to 50% in 20 weeks (and continues to wane at the same rate) for each vaccinated individual; vaccine efficacy against transmissibility is also allowed to wane at the same rate (i.e., from 20% to 11% in 20 weeks).

In this study, we use this model to retrospectively analyze past outbreaks. First, we try to match our model to epidemic patterns seen on campus for the first three semesters, 297 during which there was limited campus transmission, by varying the contact reproduction 298 number $\mathcal{R}_{\text{contact}}$ and the amount of community transmission θ and holding all other param-299 eters constant. For each parameter combination, we simulate 100 epidemic trajectories and 300 calculate the sum of squared differences between the weekly numbers of the observed and 301 predicted positive cases. The population size and testing frequencies (with twice weekly test-302 ing modeled as testing every 3 days) are set to reflect realistic campus settings. Although we 303 account for heterogeneity in the number of individuals in each population group on campus 304 (i.e., undergraduate students, graduate students, and faculty and staff members) and their 305 respective testing patterns (e.g., twice a week for undergraduate and graduate students and 306 once a week for faculty and staff members during fall and spring, 2020), we assume, for sim-307 plicity, that all other parameters are equal across different groups We further assume that 308 the population mixes homogeneously. While these assumptions are most parsimonious, epi-309 demiological parameters and mixing patterns likely differ across groups (e.g., undergraduate 310 students are more likely to infect undergraduate students and also remain asymptomatic). 311 Therefore, our model parameters describe average dynamics across different groups and must be interpreted with care. 313

For fall 2020–2021, we simulate the model assuming 3000 individuals (1000 graduate

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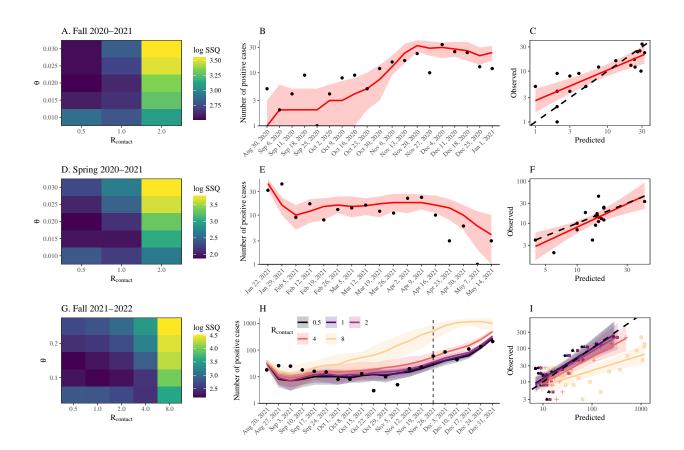


Figure 2: Retrospective analysis of past SARS-CoV-2 outbreaks on Princeton University (PU) campus. (A, D, G) Time series comparisons of model predictions with observed data across ranges of contact reproduction number $\mathcal{R}_{\text{contact}}$ and scaling parameter for community transmission θ . For each parameter combination, we simulate the model 100 times and calculate the sum of squared differences (SSQ) between the reported number of positive cases and the model-predicted number of positive cases. Heat maps represent medians of the logged sum of squared differences. (B, E, H) Model predictions. Solid lines represent median predictions. Shaded areas represent 90% quantiles for the best matching parameter set. Points represent the observed data. (C, F, I) Correlations between model predictions with observed data. Colored solid lines and shaded areas represent the estimated linear regression lines and the associated 95% CIs. Dashed lines represent the one-to-one line.

students and 2000 faculty and staff members) on campus with 1000 of them participating in asymptomatic testing twice a week. We find that a low level of contacts $\mathcal{R}_{\text{contact}} = 0.5$ and a small amount of community transmission $\theta = 0.015$ is most consistent with the observed epidemic dynamics in fall 2020 (Fig. 2A). With these parameters, the model is able to capture the rise and fall in the number of cases with the exception of a sudden decrease in the number of cases around Thanksgiving, which we do not model explicitly (Fig. 2B). The median predictions are positively correlated with the observed dynamics ($\rho = 0.83$; 95% CI: 0.61–0.93; Fig. 2C). Although a wide range of assumptions about the levels of community transmission θ are consistent with the observed dynamics, our simulations preclude high levels of contact, $\mathcal{R}_{\text{contact}} > 2$ (Supplementary Figure S7). Distancing measures on campus and contact tracing efforts likely contributed to lowering contact levels $\mathcal{R}_{\text{contact}}$.

For spring 2020–2021, we simulate the model assuming 8000 individuals (3000 under-326 graduate students, 2000 graduate students, and 3000 faculty and staff members) on campus 327 with 5000 of them participating in asymptomatic testing twice a week. We further assume 328 that 4000 individuals (3000 undergraduate students and 1000 graduate students) returned 329 to campus over 14 days (January 16, 2021–January 29, 2021); all returning individuals are 330 assumed to be quarantined for 14 days and tested upon returning. Finally, to match the 331 initial influx of cases, we assume that 1% of both returning and on-campus populations are 332 infected at the beginning of simulation (January 16, 2021). 333

A similar set of parameters can capture the observed dynamics in spring 2020–2021. The
best matching parameter predicts a slightly higher levels of community transmission $\theta = 0.02$ (Fig. 2D), but a wide range of parameters are consistent with the observed dynamics as before
(Supplementary Figure S8). Simulations also preclude high $\mathcal{R}_{\text{contact}} > 2$ again, suggesting
that transmission between students were likely limited even though they had returned to
campus—the absence of in-person teaching is likely to have contributed to lowering $\mathcal{R}_{\text{contact}}$.
We also find that initial infections (e.g., from returning students) are required to match

relatively high levels of cases in the beginning of semester (Fig. 2E). Once again, the predicted and the observed numbers of cases are positively correlated ($\rho = 0.62$; 95% CI: 0.20–0.85; Fig. 2F).

For fall 2021–2022, we assume 13000 individuals are present on campus (5000 undergraduate students, 2000 graduate students, and 6000 staff and faculty members) with 98% 345 of them vaccinated—here, vaccine-derived immunity is allowed to wane over time to ask whether the increase in the number of cases around November is consistent with the dynam-347 ics predicted by immunity waning. Vaccinated individuals are tested every week, whereas unvaccinated individuals are tested every 3 days. We further assume 5000 undergraduate 340 students returned to campus over 16 days (August 14, 2021–August 29, 2021). All students 350 were required to test upon return and quarantine until they received a negative test result; 351 for simplicity, we only model the testing process in our simulation (without quarantine) 352 given a short testing delay. Finally, we assume that 0.5% of both returning and on-campus 353 populations are infected at the beginning of simulation (August 14, 2021). We limit our 354 model comparison to November 26th before the Omicron variant was introduced on campus. 355 Even though the numbers of cases during fall 2021 (before a large outbreak) were similar 356 to those during previous semesters, we find that considerably higher levels of community 357 contact θ (≈ 10 fold higher) are required to explain the observed dynamics due to a decreased 358 susceptibility derived from vaccination (Fig. 2G). We note that the parameter θ necessarily 359 depends on our assumed vaccine efficacy against susceptibility, and θ would decrease if we assume a lower vaccine efficacy. Nonetheless, the amount of community contact would still need to be higher than previous semesters as long as the vaccine provides some protection against infection and onward transmission.

While $\theta = 0.15$ and $\mathcal{R}_{\text{contact}} = 0.5$ gives the best matching parameter set with a median 364 logged sum of squared errors of 8.88 (95\% quantile: 6.55-12.6), other parameter sets also give nearly identical fits (Fig. 2H; Supplementary Figure S9): for example, $\theta = 0.1$ and $\mathcal{R}_{\text{contact}} = 1$ gives a median logged sum of squared errors of 8.9 (95\% quantile: 5.79-13.6). Comparing 367 simulations across a wide range of $\mathcal{R}_{\text{contact}}$ (0.5–8) with $\theta = 0.1$ further illustrates that the predicted dynamics are largely insensitive to $\mathcal{R}_{\text{contact}}$ until November 26th (Fig. 2H). All 369 simulations shown in Fig. 2H, except for the $\mathcal{R}_{\text{contact}} = 8$ scenario, are similarly correlated with 370 the observed numbers of cases (Fig. 2G). While the logged sum of squared errors increases 371 with $\mathcal{R}_{\text{contact}}$ (Fig. 2G), these patterns are likely driven by the discrepancy around fall break 372 (week ending October 26th) when the number of cases decreased suddenly, rather than a lack 373 of fit—we did not explicitly model holiday effects for simplicity. Extremely high vaccination 374 rates and frequent testing likely limited transmission on campus, making epidemic dynamics 375 largely insensitive to $\mathcal{R}_{\text{contact}}$ even at a reasonably high value of $\mathcal{R}_{\text{contact}} = 4$. 376 These simulations suggest that an increase in the number of cases in November can be

277 These simulations suggest that an increase in the number of cases in November can be 278 explained by a combination of waning immunity alone without requiring additional changes 279 in transmission dynamics (note we do not allow θ or $\mathcal{R}_{contact}$ to vary over time). When 280 we exclude immune waning from the model, predicted epidemic dynamics exhibit slower 281 growth and require even higher values of campus and community contact rates ($\mathcal{R}_{contact}$ and 282 θ) to qualitatively match the observed dynamics (Supplementary Figure S10)—even so, the 283 logged sum of squared differences are generally higher (with median logged sum of squared 284 differences ranging from 6.9 to 41.3 for the same parameter regime). Thus, combining some amount of immune waning and high campus and community contact rates likely best explains
the epidemic growth near the end of the semester. We note that other factors, such as changes
in behavior, could have also contributed to the increase in the numbers of cases.

Projecting the model beyond November 26th implies that we would have seen a similar 388 growth in the number of cases if conditions remained constant even without the introduction 389 of the Omicron variant. In other words, the Delta strain would have continued to spread 390 on campus at a similar rate if the semester were to (hypothetically) continue until January 391 without additional interventions due to immune waning and growing cases in the community 392 (Fig. 2H). In reality, the situation was more complex: testing frequencies increased and 393 social gatherings were limited in response to an increase in the number of cases. These 394 interventions—as well as students returning back home as classes ended—likely would have 395 reduced contact rates (and therefore transmission of the Delta variant). This reduction in 396 transmission was likely counterbalanced by the introduction of the Omicron variant and its 397 high transmissibility and immune evasion, leading to similar and persistent growth in the 398 number of cases.

The spread of the Omicron variant on campus

Epidemiological conditions and intervention measures changed throughout the spring semester of the 2021–2022 academic year. We therefore extend to model to account for these alterations and focus on the outbreak patterns among undergraduate students. First, based on (Ferguson et al., 2021), we assume that two and three doses of vaccines reduce susceptibility

against the Omicron variant by 10% and 70%, respectively. We also assume that the transmissibility of Omicron is reduced proportionally following the previously assumed 90–to–20 ratio for the Delta variant; in other words, two and three doses of vaccines reduce transmissibility by 2.2% and 15.6%, respectively. The immunity from the third dose is assumed to take 7 days to develop (Moreira Jr et al., 2022) and wane at the same rate as before (in this case, 70% to 39% in 20 weeks). Finally, the isolation period is reduced to 5 days.

Here, we use the extended model to try to understand the drivers of a large campus out-411 break that happened on the week ending February 18, 2022 (Fig. 1D). First, we ask whether 412 changes in testing frequency from biweekly to weekly and an increased reproduction number 413 can explain the outbreak. The increase in the reproduction number can reflect increased 414 contact rates following changes in distancing policy as well as increased transmissibility of 415 the BA.2 subvariant—we do not explicitly distinguish the cause of the increase in the re-416 production number. We do so by simulating the model forward across a range of contact 417 reproduction numbers that are consistent with previous estimates ($\mathcal{R}_{\text{contact}}=2-6$) and intro-418 ducing a 20%–100% increase in the contact reproduction number on February 8, 2022, with 419 changes in the testing frequency. To match the realistic campus setting, we assume that 700 420 students are present on campus as of January 1, 2022, and the remaining 4300 students come 421 back to campus across 28 days. We assume that 99% of students are vaccinated with 60% of them being boosted—we further allow 70 booster shots on each day such that most students 423 will be boosted by the time everyone is back on campus. To match the high numbers of cases on the week ending January 7, 2022, we assume 14\% of the students present on campus are infected as of January 7, 2022 (roughly 100/700). To account for students who were infected with the Omicron variant during the fall semester, we assume that 100 students are already immune to Omicron infection at the beginning of the spring semester—this roughly corresponds to the number of PU cases that were reported in December. Finally, we take the best matching θ value for the previous semester and scale it by the number of undergraduate students relative to the entire population (therefore using $\theta = 0.15 \times 5000/13000$ throughout).

In the absence of changes in testing frequency or an increased reproduction number, 433 the model predicts the number of cases among undergraduate students to continue to de-434 crease over time (Fig. 3). Changes in testing frequency alone have negligible impact on 435 the overall dynamics; when the baseline contact reproduction number $\mathcal{R}_{\text{contact}}$ is sufficiently 436 high ($\mathcal{R}_{\text{contact}} = 6$), changing testing frequency from biweekly to weekly causes the weekly 437 case numbers to stay at a constant level (instead of decreasing). Additional increases in 438 the reproduction number (alongside the changes in testing frequency) can cause the case 439 numbers to further increase, but we are unable to match the observed dynamics even with a 100% increase in the reproduction number. Indeed, a > 10-fold increase in the numbers of 441 cases between the weeks ending February 11 and 18, 2022, would require an unrealistically 442 high increase in the contact reproduction number to explain. These simulations indicate 443 that changes in distancing and testing policies and the increased transmissibility of the BA.2 subvariant alone are unlikely to be the direct causes of the outbreak. 445

Instead, we consider the role of super-spreading events in driving a large Omicron outbreak by simulating 100–300 infections happening on the same day (February 12, 2022, the weekend following the policy change). We still include changes in testing to reflect realistic

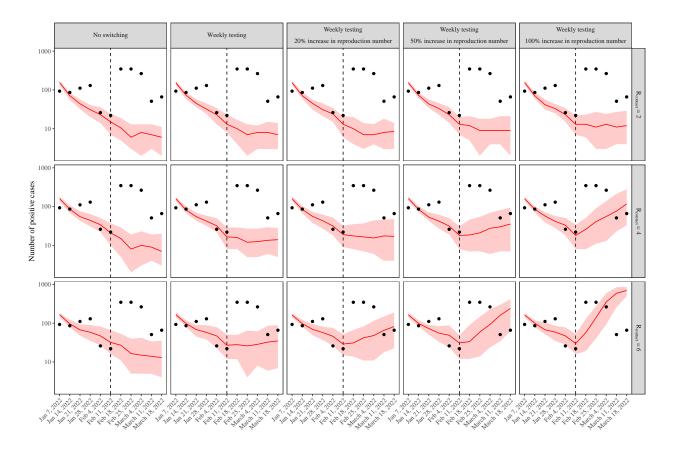


Figure 3: The impact of changes in testing frequency and an increased reproduction number on the spread of the Omicron variant. Solid lines represent median predictions. Shaded areas represent 90% quantiles across 100 simulations. Points represent the observed data. Vertical dashed lines represent the week including February 8, 2022, when distancing and testing policies were updated on PU campus.

settings on campus but do not model the increase in the reproduction number to test the sole effects of super-spreading events. In contrast to previous simulations (Fig. 3), which showed persistent growth in cases following the increase in the reproduction number, an epidemic driven by a super-spreading event plateaus and decays quickly (Fig. 4). In this case, moderate values of baseline reproduction numbers permit a small amount of onward transmission, which can sustain the epidemic for a few weeks, but the reproduction number is not high enough to cause the epidemic to keep growing. Overall, the observed patterns in

cases are more consistent with the epidemic dynamics driven by super-spreading events.

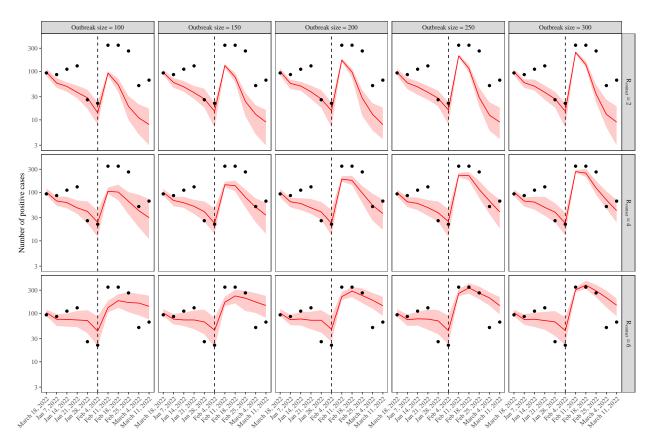


Figure 4: The impact of large super spreading events on the spread of the Omicron variant. Solid lines represent median predictions. Shaded areas represent 90% quantiles across 100 simulations. Points represent the observed data. Vertical dashed lines represent the week including February 8, 2022, when distancing and testing policies were updated on PU campus.

Discussion

- Here, we analyze SARS-CoV-2 outbreaks on the PU campus between fall 2020 and early 2022.
- We demonstrate strong spatiotemporal correlations between the patterns of spread of SARS-
- 460 CoV-2 on campus and those from surrounding communities. These correlations decreased
- with distance from Mercer County in fall 2021–2022, likely reflecting contact and commuting

patterns as the university campus reopened. Mathematical modeling further suggests limited transmission between the university population during fall and spring semesters of the 2020–2021 academic year and an increased frequency of infective community contacts during the fall semester of the 2021–2022 academic year, compared to previous semesters. An increase in the number of cases by the end of November 2021 is consistent with the increase in the levels of community cases and waning immunity. Finally, our analysis highlights the role of super-spreading events in driving the spread of the Omicron variant on the PU campus.

Although previous outbreak reports from other universities primarily focused on within-campus transmission (Wilson et al., 2020; Currie et al., 2021), a few studies identified off-campus infections as an important source of transmission (Fox et al., 2021; Hamer et al.,

470 471 2021). For example, extensive modeling efforts from Cornell University demonstrated an 472 increase in the amount of transmission from outside the university campus during fall 2021 473 and found that community transmissions are the biggest risk for faculty and staff members 474 (Frazier et al., 2022). Our study further extends these findings in demonstrating a strong 475 spatiotemporal correlation in the spread of SARS-CoV-2 between university campuses and 476 surrounding communities; however, when campus transmission is sustained, community cou-477 pling becomes less important. The degree to which community coupling affects campus 478 transmission also depends on the campus. Although Princeton University is located in a small town (Mercer County) with a population of 390,000 (www.census.gov), it is located near large cities, such as New York City and Philadelphia, which can drive infections in smaller cities nearby (Grenfell and Harwood, 1997). For example, contact tracing efforts from Boston University, which is located in a large metropolitan area, found that more than 50% of infections among Boston University affiliates with known exposures could be attributed to sources outside of the university campus (Hamer et al., 2021). In contrast, other university campuses that are far from urban areas may experience weaker community coupling. The degree of coupling will also depend on intervention measures in surrounding communities and on campus. Understanding these heterogeneities is critical for preventing future campus outbreaks.

Our analysis also suggests that comparing the ratios between the cases on university
campuses and neighboring communities can also provide a useful measure for how well a
university campus is controlling the epidemic; however, that this ratio needs to be interpreted
with caution as it is sensitive to changes in testing patterns as well as the numbers of students
on campus. For example, the ratios of cases can suddenly change during holidays when
students are away from campus. Future studies could combine viral phylogenetic data to
better understand spatial patterns of SARS-CoV-2 on campus.

There are several limitations to our analysis. While we demonstrate strong spatiotemporal correlation in the spread of SARS-CoV-2, we are not able to infer the direction of
causality—that is, our analysis does not rule out the possibility that transmission on campus drove infections in nearby communities (as opposed to community transmission driving
on-campus infections). However, seeding from campus is unlikely: intervention measures
on campus (e.g., frequent asymptomatic testing, contact tracing, and virtual classes during
fall and spring semesters of 2020) likely limited onward transmission on campus. In addition, even during periods of large Omicron outbreaks on campus in early 2022, the number
of COVID-19 cases in Mercer County remained low, implying limited transmission from

campus to community. Decreasing patterns in epidemic correlations with distance further highlight the role of spatial spread in driving dynamics of SARS-CoV-2—such patterns are consistent with spatial spread of many other respiratory pathogens (Grenfell et al., 2001; Viboud et al., 2006; Baker et al., 2019).

Our mathematical model relies on simplifying assumptions. For example, we assume 510 conservatively that the entire university populations mix homogeneously and have identical 511 campus and community contact rates (captured by $\mathcal{R}_{\text{contact}}$ and θ , respectively). In reality, 512 increases in cases were often associated with specific transmission clusters, suggesting het-513 erogeneity in transmission patterns. Contact levels also likely differ between different groups: 514 for example, faculty and staff members are more likely to interact with community members 515 than undergraduate students and would be at a higher risk for community infections (Frazier 516 et al., 2022); therefore, our homogeneous mixing assumption is conservatively pessimistic. 517 We also do not account for changes in behavior; instead, we assume constant values for θ 518 and $\mathcal{R}_{\text{contact}}$ throughout a semester. While we cannot rule out the possibility that changes 519 in behavior (and therefore transmission rates) could have contributed to various epidemics 520 (e.g., the Omicron wave beginning in the fall semester of the 2021–2022 academic year), we 521 were able to capture the majority of epidemic patterns without modeling them. We also do 522 not explore parameter uncertainty, which can lead to underestimation of overall uncertainty (Elderd et al., 2006). We also note that intervention measures that were introduced to PU 524 may not necessarily be applicable in other institutions.

Despite the simplicity of the analysis, our study provides important lessons for controlling
SARS-CoV-2 outbreaks on university campuses in general. First, our analysis highlights the

power of mass asymptomatic testing for epidemic measurement and planning. Combining other interventions measures, such as social distancing, mask wearing, and vaccination, can help provide a safe means of reopening university campuses—but the extent to which these interventions are implemented will necessarily depend on resource availability. Second, we expect immune waning and superspreading to continue to play important roles in driving 532 campus transmission—keeping vaccine statuses up-to-date within the campus community 533 will be critical moving forward. In addition, preventing large gatherings can help prevent 534 large superspreading events in the midst of a rising epidemic. Third, the safe reopening 535 of a university campus must consider the spread of SARS-CoV-2 within the surrounding 536 community as they can both potentially drive transmission in each other—however, the 537 degree to which infections spread from campus to community remains uncertain. Finally, 538 intervention measures placed on campuses must continue to adapt and change to reflect 539 changes in epidemiological conditions. 540

The emergence of new variants—in particular, their ability to evade prior immunity and transmit better—continues to add uncertainty to the future controllability of the ongoing SARS-CoV-2 pandemic. Nonetheless, as population-level immunity increases (either due to infection or vaccination), we are (hopefully) transitioning to an endemic phase, during which COVID-19 is expected to become less severe (Lavine et al., 2021). Many institutions have so far sought to minimize infections on their campuses early in the pandemic, thereby implementing as many interventions as possible—but such measures can be difficult to maintain for prolonged periods both economically and societally. As the Omicron variant began to spread, many campuses—including the PU campus—opted in for less intense interventions,

reflecting difficulties in controlling the spread and a lack of severe cases among the majority of vaccinated students. As we continue to transition to future phases of the pandemic,
the expectations for reopening campuses (e.g., whether to minimize infections on campuses)
need to be re-evaluated, accounting not only for changes in epidemic dynamics but also for
our perception of the pathogen. The answers to these questions ultimately depend on the
landscape of SARS-CoV-2 immunity and its future evolutionary dynamics (Saad-Roy et al.,
2020; Baker et al., 2021).

Data availability

All data and code are stored in a publicly available GitHub repository (https://github.

com/parksw3/university-covid).

560 References

Baker, R. E., A. S. Mahmud, C. E. Wagner, W. Yang, V. E. Pitzer, C. Viboud, G. A. Vecchi,
C. J. E. Metcalf, and B. T. Grenfell (2019). Epidemic dynamics of respiratory syncytial
virus in current and future climates. *Nature communications* 10(1), 1–8.

Baker, R. E., S. W. Park, C. E. Wagner, and C. J. E. Metcalf (2021). The limits of SARSCoV-2 predictability. *Nature Ecology & Evolution* 5(8), 1052–1054.

Brett, T. S. and P. Rohani (2020). Transmission dynamics reveal the impracticality

- of COVID-19 herd immunity strategies. *Proceedings of the National Academy of Sciences* 117(41), 25897–25903.
- Brook, C. E., G. R. Northrup, A. J. Ehrenberg, J. A. Doudna, M. Boots, I. S.-C.-. T.
- 570 Consortium, et al. (2021). Optimizing COVID-19 control with asymptomatic surveillance
- testing in a university environment. *Epidemics 37*, 100527.
- ⁵⁷² Cobey, S. (2020). Modeling infectious disease dynamics. Science 368(6492), 713–714.
- ⁵⁷³ Currie, D. W., G. K. Moreno, M. J. Delahoy, I. W. Pray, A. Jovaag, K. M. Braun, D. Cole,
- T. Shechter, G. C. Fajardo, C. Griggs, et al. (2021). Interventions to Disrupt Coronavirus
- Disease Transmission at a University, Wisconsin, USA, August-October 2020. Emerging
- infectious diseases 27(11), 2776.
- 577 Duke University (2021). Important COVID updates for undergrads, Aug. 30.
- 578 Elderd, B. D., V. M. Dukic, and G. Dwyer (2006). Uncertainty in predictions of disease
- spread and public health responses to bioterrorism and emerging diseases. *Proceedings of*
- the National Academy of Sciences 103(42), 15693–15697.
- Endo, A., Centre for the Mathematical Modelling of Infectious Diseases COVID-19 Working
- Group, S. Abbott, A. Kucharski, and S. Funk (2020). Estimating the overdispersion in
- 553 COVID-19 transmission using outbreak sizes outside China [version 3; peer review: 2
- approved]. Wellcome Open Research 5(67).
- Ferguson, N., A. Ghani, A. Cori, A. Hogan, W. Hinsley, and E. Volz (2021). Report 49:
- Growth, population distribution and immune escape of Omicron in England.

- Flaxman, S., S. Mishra, A. Gandy, H. J. T. Unwin, T. A. Mellan, H. Coupland, C. Whittaker,
- H. Zhu, T. Berah, J. W. Eaton, et al. (2020). Estimating the effects of non-pharmaceutical
- interventions on COVID-19 in Europe. *Nature* 584 (7820), 257–261.
- Fox, M. D., D. C. Bailey, M. D. Seamon, and M. L. Miranda (2021). Response to a COVID-19
- outbreak on a university campus—Indiana, August 2020. Morbidity and Mortality Weekly
- Report 70(4), 118.
- Frazier, P. I., J. M. Cashore, N. Duan, S. G. Henderson, A. Janmohamed, B. Liu, D. B.
- Shmoys, J. Wan, and Y. Zhang (2022). Modeling for COVID-19 college reopening deci-
- sions: Cornell, a case study. Proceedings of the National Academy of Sciences 119(2).
- Galanti, M., S. Pei, T. K. Yamana, F. J. Angulo, A. Charos, D. L. Swerdlow, and J. Shaman
- (2021). Social distancing remains key during vaccinations. Science 371 (6528), 473–474.
- Gibson, G., J. S. Weitz, M. P. Shannon, B. Holton, A. Bryksin, B. Liu, M. Sieglinger,
- A. R. Coenen, C. Zhao, S. J. Beckett, et al. (2021). Surveillance-to-Diagnostic Testing
- Program for Asymptomatic SARS-CoV-2 Infections on a Large, Urban Campus in Fall
- 2020. Epidemiology 33(2), 209–216.
- 602 Grenfell, B. and J. Harwood (1997). (meta) population dynamics of infectious diseases.
- Trends in ecology & evolution 12(10), 395-399.
- 664 Grenfell, B. T., O. N. Bjørnstad, and J. Kappey (2001). Travelling waves and spatial hier-
- archies in measles epidemics. Nature 414 (6865), 716–723.

- Hamer, D. H., L. F. White, H. E. Jenkins, C. J. Gill, H. E. Landsberg, C. Klapperich,
- K. Bulekova, J. Platt, L. Decarie, W. Gilmore, et al. (2021). Assessment of a COVID-19
- control plan on an urban university campus during a second wave of the pandemic. JAMA
- Network Open 4(6), e2116425-e2116425.
- Harvard University (2021). Increase in COVID-19 cases, Take Steps to Protect Yourself & Our Community.
- He, D., E. L. Ionides, and A. A. King (2010). Plug-and-play inference for disease dynamics:
- measles in large and small populations as a case study. Journal of the Royal Society
- Interface 7(43), 271–283.
- Hellewell, J., S. Abbott, A. Gimma, N. I. Bosse, C. I. Jarvis, T. W. Russell, J. D. Munday,
- A. J. Kucharski, W. J. Edmunds, F. Sun, et al. (2020). Feasibility of controlling COVID-19
- outbreaks by isolation of cases and contacts. The Lancet Global Health 8(4), e488–e496.
- Holmdahl, I. and C. Buckee (2020). Wrong but useful—what COVID-19 epidemiologic
- models can and cannot tell us. New England Journal of Medicine 383(4), 303-305.
- 620 Kissler, S. M., C. Tedijanto, E. Goldstein, Y. H. Grad, and M. Lipsitch (2020). Project-
- ing the transmission dynamics of SARS-CoV-2 through the postpandemic period. Sci-
- $ence\ 368(6493),\ 860-868.$
- Koelle, K., M. A. Martin, R. Antia, B. Lopman, and N. E. Dean (2022). The changing
- epidemiology of SARS-CoV-2. Science 375 (6585), 1116–1121.

- 625 Kraemer, M. U., O. G. Pybus, C. Fraser, S. Cauchemez, A. Rambaut, and B. J. Cowling
- 626 (2021). Monitoring key epidemiological parameters of SARS-CoV-2 transmission. *Nature*
- medicine 27(11), 1854-1855.
- Lavezzo, E., E. Franchin, C. Ciavarella, G. Cuomo-Dannenburg, L. Barzon, C. Del Vecchio,
- 629 L. Rossi, R. Manganelli, A. Loregian, N. Navarin, et al. (2020). Suppression of a SARS-
- 630 CoV-2 outbreak in the Italian municipality of Vo'. Nature 584 (7821), 425–429.
- Lavine, J. S., O. N. Bjornstad, and R. Antia (2021). Immunological characteristics govern
- the transition of COVID-19 to endemicity. Science 371 (6530), 741–745.
- Lopman, B., C. Y. Liu, A. Le Guillou, T. L. Lash, A. P. Isakov, and S. M. Jenness (2020).
- A model of COVID-19 transmission and control on university campuses. *MedRxiv*.
- 635 Metcalf, C. J. E., D. H. Morris, and S. W. Park (2020). Mathematical models to guide
- pandemic response. *Science* 369 (6502), 368–369.
- Moreira Jr, E. D., N. Kitchin, X. Xu, S. S. Dychter, S. Lockhart, A. Gurtman, J. L. Perez,
- 638 C. Zerbini, M. E. Dever, T. W. Jennings, et al. (2022). Safety and efficacy of a third dose
- of BNT162b2 COVID-19 vaccine. New England Journal of Medicine.
- Prunas, O., J. L. Warren, F. W. Crawford, S. Gazit, T. Patalon, D. M. Weinberger, and
- V. E. Pitzer (2022). Vaccination with BNT162b2 reduces transmission of SARS-CoV-2 to
- 642 household contacts in Israel. Science, eabl4292.
- Saad-Roy, C. M., S. E. Morris, C. J. E. Metcalf, M. J. Mina, R. E. Baker, J. Farrar, E. C.
- Holmes, O. G. Pybus, A. L. Graham, S. A. Levin, et al. (2021). Epidemiological and

- evolutionary considerations of SARS-CoV-2 vaccine dosing regimes. *Science* 372(6540),
- 646 363–370.
- Saad-Roy, C. M., C. E. Wagner, R. E. Baker, S. E. Morris, J. Farrar, A. L. Graham, S. A.
- Levin, M. J. Mina, C. J. E. Metcalf, and B. T. Grenfell (2020). Immune life history,
- vaccination, and the dynamics of SARS-CoV-2 over the next 5 years. Science 370(6518),
- 650 811-818.
- Tartof, S. Y., J. M. Slezak, H. Fischer, V. Hong, B. K. Ackerson, O. N. Ranasinghe, T. B.
- Frankland, O. A. Ogun, J. M. Zamparo, S. Gray, et al. (2021). Effectiveness of mRNA
- 653 BNT162b2 COVID-19 vaccine up to 6 months in a large integrated health system in the
- USA: a retrospective cohort study. The Lancet 398(10309), 1407–1416.
- Viboud, C., O. N. Bjørnstad, D. L. Smith, L. Simonsen, M. A. Miller, and B. T. Gren-
- fell (2006). Synchrony, waves, and spatial hierarchies in the spread of influenza. sci-
- ence 312(5772), 447-451.
- 658 Wilson, E., C. V. Donovan, M. Campbell, T. Chai, K. Pittman, A. C. Seña, A. Pettifor, D. J.
- Weber, A. Mallick, A. Cope, et al. (2020). Multiple COVID-19 clusters on a university
- campus—North Carolina, August 2020. Morbidity and Mortality Weekly Report 69(39),
- 661 1416.

662 Supplementary Figures

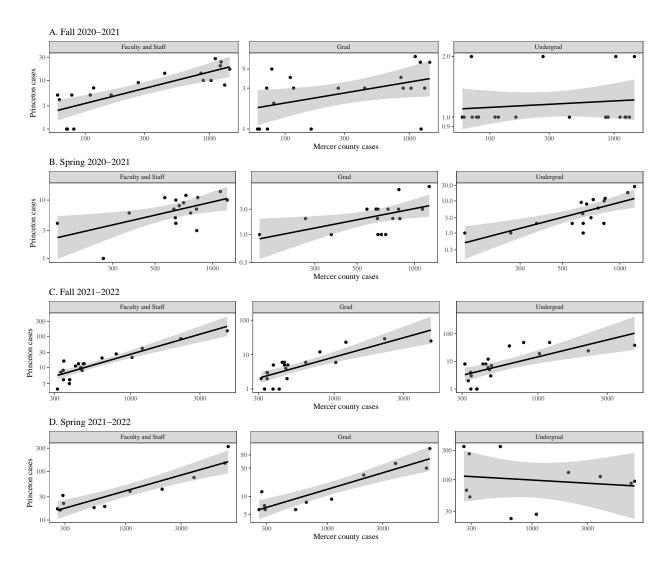


Figure S1: Correlations between the weekly number of cases in PU and in Mercer County stratified by subpopulations. Points represent the number of reported cases. Solid lines and shaded areas represent the regression line and the associated 95% confidence intervals.

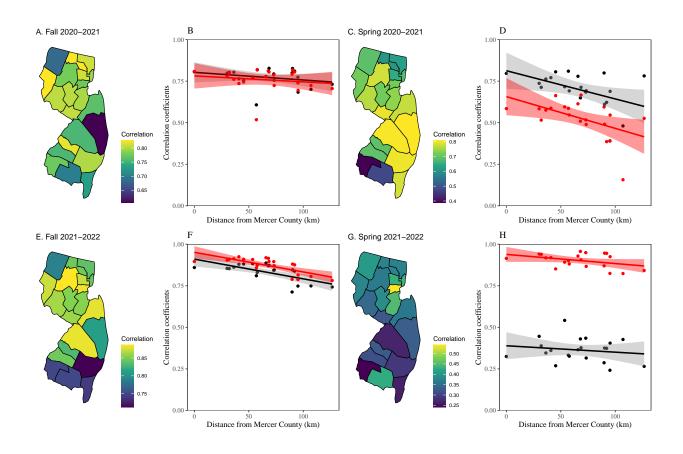


Figure S2: Correlations between the weekly number of cases in PU and in counties in New Jersey. (A–C) Map of correlations between the weekly number of cases in PU and in counties in New Jersey. (D–F) Relationship between case correlations and distance from Mercer County. Points represent the estimated correlation coefficients. Solid lines and shaded areas represent the regression line and the associated 95% confidence intervals. Black points and lines represent correlations based on all cases in PU. Red points and lines represent correlations based on cases among faculty and staff members in PU.

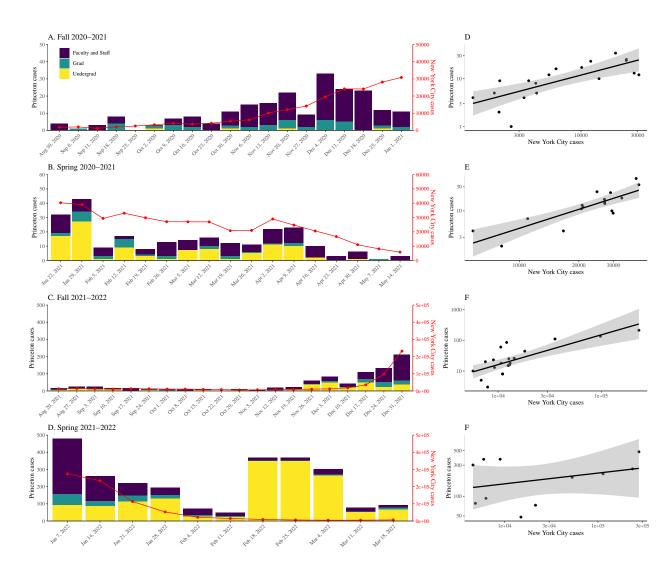


Figure S3: Dynamics of SARS-CoV-2 outbreaks in PU and New York City. (A–C) Epidemic trajectories across three semesters: Fall 2020 (A), Spring 2020 (B), and Fall 20201 (C). Colored bar plots represent the weekly number of cases from both asymptomatic and symptomatic testing in PU. Red lines represent the weekly number of cases in New York City. (D–F) Correlations between the weekly number of cases in PU and in New York City. Solid lines and shaded areas represent the estimated linear regression lines and the associated 95% CIs.

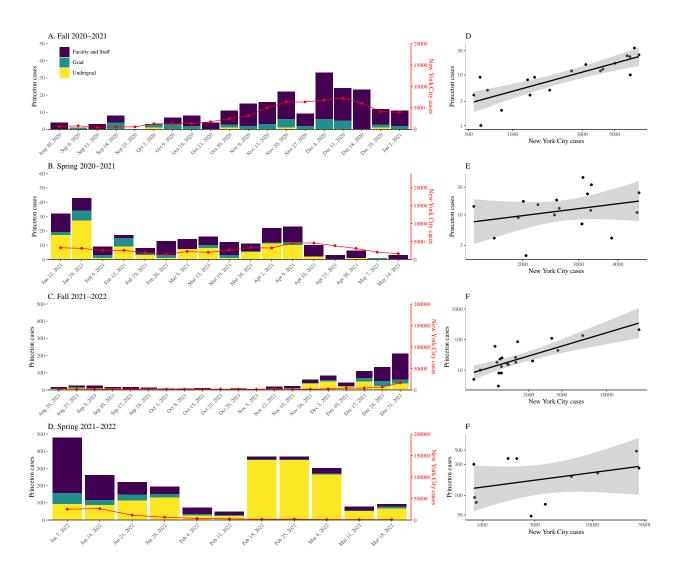


Figure S4: Dynamics of SARS-CoV-2 outbreaks in PU and Philadelphia. (A–C) Epidemic trajectories across three semesters: Fall 2020 (A), Spring 2020 (B), and Fall 20201 (C). Colored bar plots represent the weekly number of cases from both asymptomatic and symptomatic testing in PU. Red lines represent the weekly number of cases in Philadelphia. (D–F) Correlations between the weekly number of cases in PU and in Philadelphia. Solid lines and shaded areas represent the estimated linear regression lines and the associated 95% CIs.

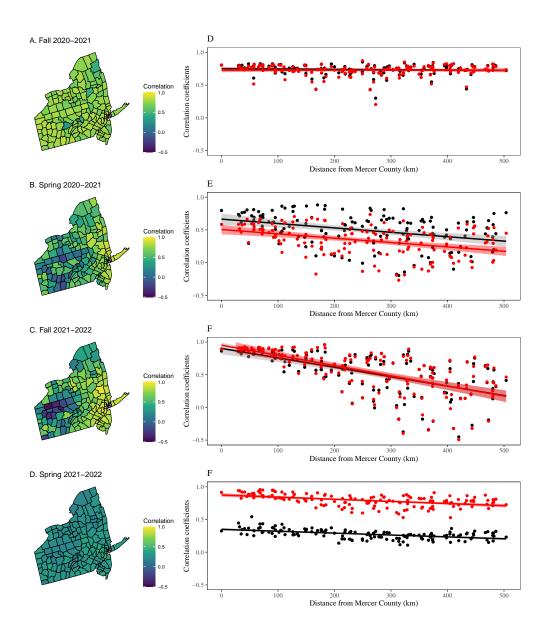


Figure S5: Correlations between the weekly number of cases in PU and in counties in New Jersey, Pennsylvania, and New York State. (A–C) Map of correlations between the weekly number of cases in PU and in counties in New Jersey, Pennsylvania, and New York State. (D–F) Relationship between case correlations and distance from Mercer County. Points represent the estimated correlation coefficients. Solid lines and shaded areas represent the regression line and the associated 95% confidence intervals. Black points and lines represent correlations based on all cases in PU. Red points and lines represent correlations based on cases among faculty and staff members in PU. New York City is excluded from this analysis as the data provided by New York Times are not further stratified by county levels.

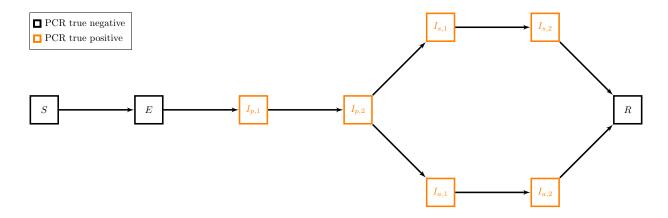


Figure S6: Compartmental diagram of the individual-based model. Each compartment represents a stage of infection: susceptible S, exposed S, pre-symptomatic I_p , symptomatic I_s , asymptomatic I_a , and recovered R. Pre-symptomatic, symptomatic, and asymptomatic stages are further divided into two subcompartments. Individuals in pre-symptomatic, symptomatic, and asymptomatic stages can test positive with 95% sensitivity.

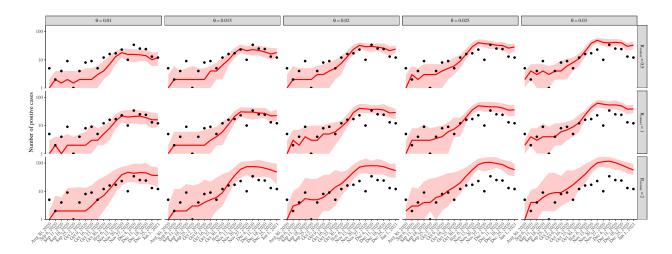


Figure S7: Comparisons between model predictions and the observed numbers of cases for fall 2020. Points represent the weekly number of reported cases in PU. Red lines and shaded areas represent median model predictions and 90% quantiles across 100 simulations. See figure 2 in the main text for details.

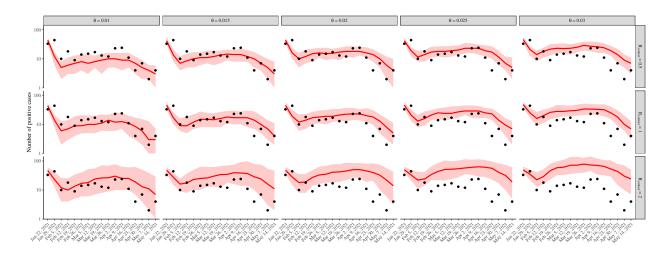


Figure S8: Comparisons between model predictions and the observed numbers of cases for spring 2020. Points represent the weekly number of reported cases in PU. Red lines and shaded areas represent median model predictions and 90% quantiles across 100 simulations. See figure 2 in the main text for details.

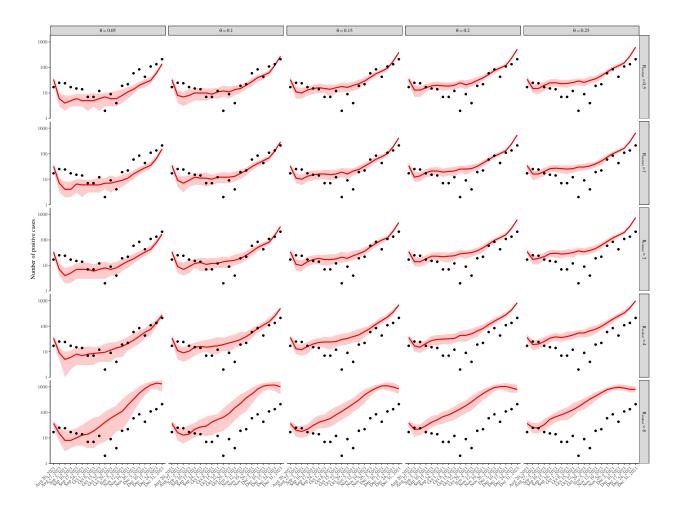


Figure S9: Comparisons between model predictions and the observed numbers of cases for fall 2021. Points represent the weekly number of reported cases in PU. Red lines and shaded areas represent median model predictions and 90% quantiles across 100 simulations. See figure 2 in the main text for details.

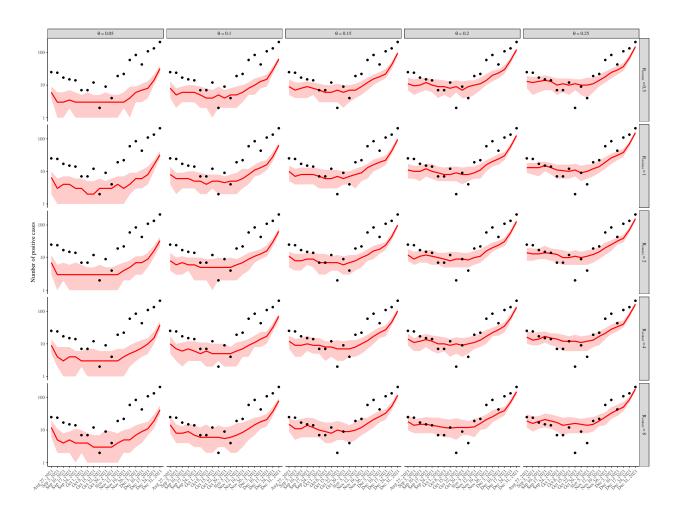


Figure S10: Comparisons between model predictions and the observed numbers of cases for fall 2021 without immunity waning. Points represent the weekly number of reported cases in PU. Red lines and shaded areas represent median model predictions and 90% quantiles across 100 simulations. See figure 2 in the main text for details.