


















Ensemble forecasts of COVID-19 activity to support Australia’s pandemic response: 2020–22

Robert Moss¹ Ruarai J. Tobin¹ Mitchell O’Hara-Wild²
Adeshina I. Adekunle³ Dennis Liu⁴ Tobin South⁴ Dylan J. Morris⁴
Gerard E. Ryan^{5,1} Tianxiao Hao^{5,1} Aarathy Babu⁵
Katharine L. Senior^{5,1} James G. Wood⁷ Nick Golding^{5,6}
Joshua V. Ross^{8,9} Peter Dawson³ Rob J. Hyndman² David J. Price^{1,10,*}
James M. McCaw^{1,11,*} Freya M. Shearer^{1,5,*}

¹: Melbourne School of Population and Global Health, The University of Melbourne;
²: Department of Econometrics and Business Statistics, Monash University; ³: Defence Science and Technology Group; ⁴: School of Computer and Mathematical Sciences, The University of Adelaide; ⁵: The Kids Research Institute Australia; ⁶: School of Physics, Mathematics and Computing, University of Western Australia; ⁷: School of Population Health, University of New South Wales; ⁸: Department for Health and Wellbeing, Government of South Australia; ⁹: South Australian Health and Medical Research Institute; ¹⁰: Department of Infectious Diseases, The Peter Doherty Institute for Infection and Immunity; ¹¹: School of Mathematics and Statistics, The University of Melbourne; *: Equal authorship

Abstract

During the COVID-19 pandemic, many countries used real-time data analyses, predictive modelling, and COVID-19 case forecasts, to incorporate emerging evidence into their decisions. In Australia, national and jurisdictional public health responses were informed by weekly ensemble forecasts of daily COVID-19 case counts for each of Australia’s eight states and territories, produced by a consortium of researchers under contract with the Australian Government. We evaluated approximately 100,000 predictions for daily case counts 1–28 days into the future, generated between July 2020 and December 2022, and report here (a) how the ensemble forecasts supported public health responses; (b) how well the ensemble forecast performed, relative to the forecasts produced by each contributing team; and (c) how we refined our reporting and visualisations to ensure that outputs were interpreted appropriately. Similar to COVID-19 forecasting studies in other countries, we found that the ensemble forecast consistently out-performed the individual

NOTE: This preprint reports new research that has not been certified by peer review and should not be used to guide clinical practice.

15 model forecasts, and that performance was lowest when there were rapid changes in
16 the epidemiology, such as periods around epidemic peaks. Our consortium's inter-
17 nal peer-review process allowed us to explain how features of each ensemble forecast
18 related to the design of the individual models, and this helped enable public health
19 stakeholders to interpret the forecasts appropriately. Ultimately, our forecasts pro-
20 vided information that supported public health responses during periods of different
21 policy goals, and over a wide range of epidemic scenarios.

22 1 Introduction

23 The COVID-19 pandemic resulted in major social disruptions around the world [1, 2],
24 as governments implemented a range of measures to curb infections and limit deaths in
25 the face of a novel virus [3–5]. Initial decisions were guided by public health response
26 plans, and as the pandemic unfolded governments increasingly made use of modelling
27 to incorporate emerging evidence into their decisions [6–11]. Australia’s national and
28 jurisdictional public health responses were informed by the results of real-time analyses
29 that were presented to key government advisory committees in weekly situation reports,
30 starting 4th April 2020 [12, 13]. Between July 2020 and December 2023, these reports
31 included ensemble forecasts of daily COVID-19 case counts for each of Australia’s eight
32 states and territories.

33 Beginning on 1st February 2020, the Australian government progressively closed bor-
34 ders to countries with established epidemics, culminating in almost complete cessation of
35 travel from 20th March 2020, and imposed mandatory 14-day hotel quarantine on overseas
36 arrivals from 16th March 2020. These measures were effective in limiting community ex-
37 posure from infected international arrivals. Australia’s initial COVID-19 response focused
38 on preventing local transmission [14] and, despite a steady influx of imported cases, local
39 elimination was achieved for prolonged periods throughout 2020 and 2021, punctuated by
40 several waves of local transmission. Australia’s COVID-19 vaccination program began in
41 February 2021, with the aim of pivoting from preventing local transmission to “reopen-
42 ing” the country (i.e., easing non-pharmaceutical interventions). Vaccination targets and
43 reopening plans were informed by modelling studies [15, 16]. Beginning on 1st November
44 2021, Australian jurisdictions progressively relaxed quarantine requirements for returning
45 fully-vaccinated Australian residents and citizens, and subsequently for international trav-
46 ellers, with quarantine requirements fully removed as of 3rd March 2022. Over the 2022
47 calendar year Australia experienced substantial local transmission of multiple COVID-19
48 variants, with major waves of Omicron BA.1, BA.2, and BA.5, consistent with global
49 epidemiology [17]. Many of these policy decisions were directly informed by our situation
50 reports and the ensemble forecasts contained within.

51 Here we present an analysis of these ensemble forecasts, demonstrating how the ensem-
52 ble consistently out-performed the individual models that were included in the ensemble,
53 and how these forecasts supported public health responses over the 2020–2022 calendar
54 years. These case forecasts also served as an input for COVID-19 hospital bed occupancy
55 forecasts (December 2021 to December 2023) that further supported Australia’s public
56 health responses [18]. A timeline of key events during this period is presented in [Table 1](#).

Year	Month	Event
2020	Jan	First cases imported National goal of preventing local transmission is announced
	Mar	Limited local transmission occurs nationally Hotel quarantine established for returning residents
	Jun	Sustained local transmission occurs in Victoria
	Oct	Local transmission is suppressed in Victoria
	2021	Feb
May		Vaccine roll-out for all adults begins
Jul		Delta variant becomes established in NSW (and subsequently in other jurisdictions)
Aug		National reopening plan targets are announced
Nov		NSW is the first jurisdiction to remove quarantine requirements for returning residents
Dec		Omicron BA.1 becomes established in all jurisdictions except WA
2022	Jan	Omicron BA.1 activity peaks in affected jurisdictions Vaccine roll-out for children aged 5–11 begins Booster doses recommended for at-risk individuals
	Feb	Borders opened to all vaccinated travellers except in WA
	Mar	Omicron BA.2 and BA.5 become established WA is the last jurisdiction to remove quarantine requirements
	Jul	Omicron BA.5 activity peaks
	Sep	Multiple variants become established
	Dec	Multiple variants activity peaks

Table 1: A timeline of key events prior to, and during, the study period (July 2020 to December 2022, inclusive).

57 2 Methods

58 2.1 COVID-19 case data

59 For target data, we used de-identified line lists of reported cases for each Australian state
60 and territory, extracted at each week of the study period from the National Notifiable
61 Disease Surveillance System (NNDSS). The data included the jurisdiction (state or terri-
62 tory), the date of symptom onset (where available), the date when the case notification
63 was received by the jurisdictional health department, and whether the infection was epi-
64 demologically deemed to be acquired locally or overseas. We imputed missing symptom
65 onset dates and estimated reporting delays using a time-varying delay distribution to
66 characterise the duration between symptom onset and case notification [19]. Symptom
67 onset dates were reported for 82% of cases in 2020. As Australia’s response pivoted to
68 “reopening” the country, and daily case counts greatly increased, symptom onset report-
69 ing decreased to 48% of cases in 2021, and 38% in 2022. Nationally aggregated daily case
70 counts are shown in [Figure 1](#).

71 2.2 Forecasting teams and models

72 Four modelling teams contributed forecasts for inclusion in the ensemble, using differ-
73 ent types of model (see [Table 2](#)). This included two stochastic compartmental models,
74 developed by the Defence Science and Technology Group (DST) and the University of
75 Melbourne (UoM), which incorporated different assumptions about effective reproduc-
76 tion numbers and population immunity; a branching process model developed by the
77 University of Adelaide (UoA); and an autoregressive model developed by Monash Uni-
78 versity (MU). We began with three teams contributing to the ensemble (MU, UoA, and
79 UoM) and the DST team began participating in August 2021. After a four-month period
80 where all four teams contributed to the ensemble forecast (2021-11-25 to 2022-03-22, in-
81 clusive), the UoA team stopped participating due to limited staff availability. Throughout
82 this manuscript we will refer to the models, and the forecasts that they produced, using
83 the acronyms listed in [Table 2](#) and used in this paragraph.

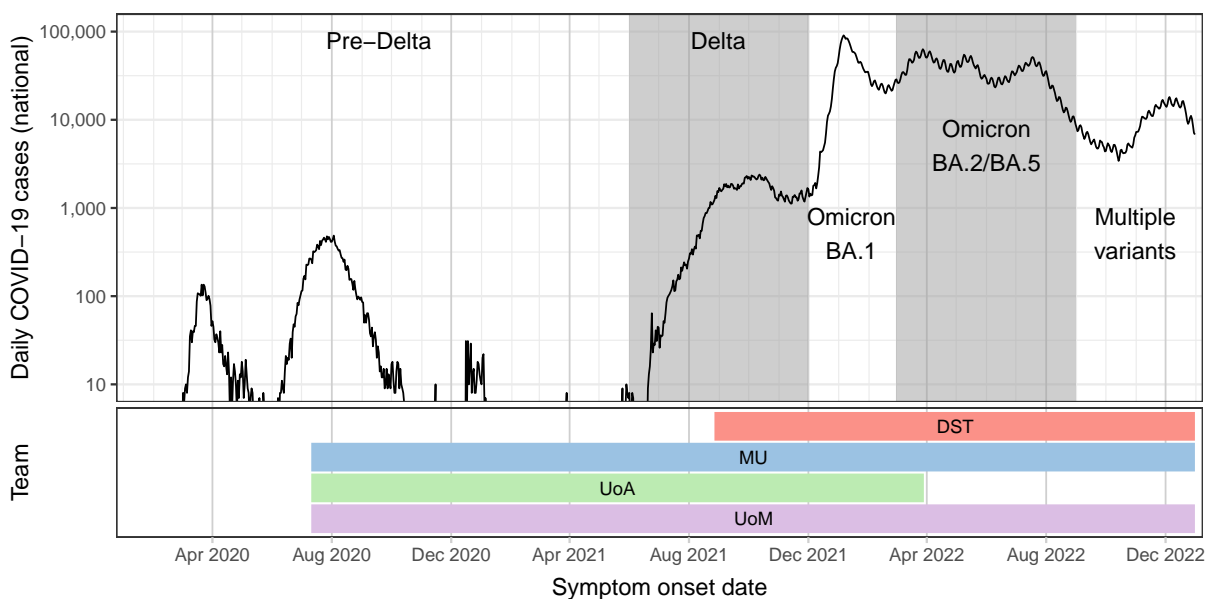


Figure 1: Top: Australian daily confirmed COVID-19 cases for the 2020–2022 calendar years. Dominant strain(s) are identified by alternating backgrounds (white and shaded), as determined by epidemiological assessment in situation reports [13]. Bottom: The periods over which each team contributed to the ensemble forecast. Team names and modelling approaches are listed in Table 2.

Institute	Methodology
Defence Science Technology (DST)	Stochastic SEIR-type compartmental model
Monash University (MU)	Global autoregression model
University of Adelaide (UoA)	Branching process model
University of Melbourne (UoM)	Stochastic SEIR-type compartmental model

Table 2: The teams that contributed to the ensemble forecasts over the study period.

84 The UoM team began contributing national forecasts in April 2020 [20], using a model
85 adapted from a long-running seasonal influenza forecasting project [21–24], while the other
86 teams created new models in response to the emergence of local COVID-19 transmission.
87 All models were subject to ongoing development throughout the study, and these devel-
88 opments are documented in our weekly situation reports [13]. When new model features
89 were introduced, retrospective forecasts were validated against previously-reported data
90 before the new model iteration was considered for inclusion in the ensemble.

91 **2.3 Inclusion/exclusion criteria**

92 If there were known, or strongly suspected, data quality issues for any jurisdiction that
93 could not be resolved with post-processing, we did not include ensemble forecasts for
94 those jurisdictions in the weekly situation report. Models were also excluded when the
95 forecasts produced for that week were assessed to be inappropriately sensitive to trends
96 in the most recent case counts, which were known to be subject to time-varying reporting
97 delays [19]. This decision-making process was facilitated by direct communication with
98 data custodians in the relevant jurisdictions and national public health committees, which
99 enabled rapid awareness of data quality issues.

100 Each week, the modelling teams generated forecasts from the provided data extracts
101 and shared the results for peer review by the broader analysis team (comprising all mod-
102 elling teams and other analysts, which we will refer to as the “consortium” hereafter) in
103 a private online discussion board (in a Slack workspace). Peer review findings and ret-
104 rospective performance results for each model were then tabled for discussion in weekly
105 online consortium meetings, where we collectively reached agreement on which models,

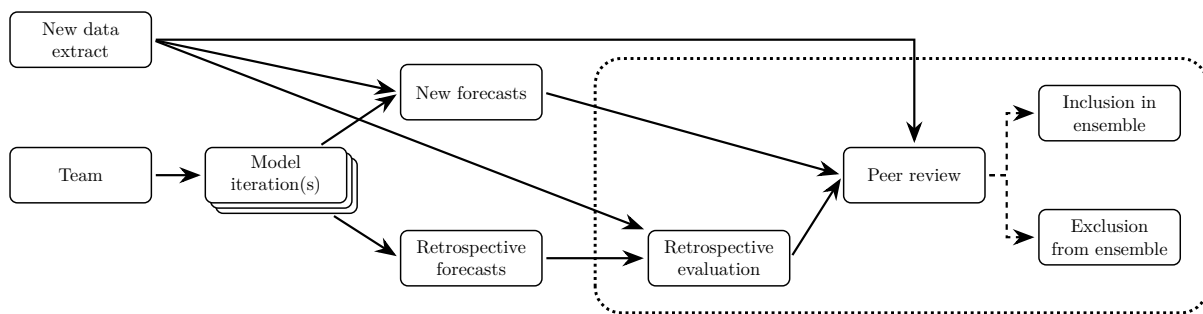


Figure 2: A flowchart of the weekly peer review and inclusion/exclusion decision process, shown here for the model iteration(s) provided by a single modelling team in a given week. The dotted box indicates activities and decisions made by the entire consortium.

106 and model iterations, to include in the ensemble forecast for each jurisdiction that week
107 (see Figure 2). This approach of ongoing discussion and reaching a collective consensus
108 has been shown to improve the performance of human predictions in domains such as
109 geopolitics [25].

110 Models were excluded when their outputs differed dramatically from the consortium’s
111 collective expectations and the current epidemiological assessment, or when a technical
112 issue was identified by the modelling team or by the peer review process. Importantly,
113 each model was evaluated against the most recent data rather than competing against
114 the other models (e.g., via benchmarks such as skill scores), with the purpose of deciding
115 whether the model outputs were likely to be a reliable basis for decision-making [26]. The
116 outcomes of these collective decisions are captured in publicly available reports [13].

117 2.4 Ensemble forecast and evaluation

118 Model forecasts were submitted in the form of 2000 simulated trajectories of daily case
119 counts over the 28-day forecast horizon for each of the eight jurisdictions. We generated
120 equal-weight ensemble forecasts for each jurisdiction by sampling trajectories from each
121 model forecast for that jurisdiction. Notably, weights were allocated to each team rather
122 than to each model iteration. For example, when a team contributed forecasts from
123 multiple model iterations, we considered the iterations as representing a single model over
124 a discrete parameter space. Accordingly, the addition or removal of one model iteration

125 impacted the relative weights of all model iterations from that team, but did not affect
126 the weights allocated to other teams.

127 As described in the previous section, we held weekly meetings in which we reviewed
128 individual model forecasts and the ensemble forecast prior to including these forecasts in
129 our weekly situation reports to government. Forecast performance was evaluated visually
130 (using daily credible intervals) against previous weeks' forecasts and the most recent data,
131 and quantitatively using Continuous Ranked Probability Scores (CRPS). The weekly situ-
132 ation reports included plots of the ensemble forecast for each jurisdiction, and comparisons
133 of the previous forecasts (made 1–4 weeks ago) against the most recent data for each ju-
134 risdiction. These comparisons included visualisations for the ensemble, and also for each
135 individual model. In this study we also report forecast bias $B(t)$ as per Funk et al. [27]:

$$B(t) = 1 - [P(Y_t \leq y_t) + P(Y_t \leq y_t - 1)]$$

136 2.5 Data availability

137 The analysis code, summary outputs, figures, and supplementary materials are pro-
138 vided in a public git repository: [https://gitlab.unimelb.edu.au/rgmoss/analysis-](https://gitlab.unimelb.edu.au/rgmoss/analysis-australian-covid19-ensemble-forecasts-results)
139 [australian-covid19-ensemble-forecasts-results](https://gitlab.unimelb.edu.au/rgmoss/analysis-australian-covid19-ensemble-forecasts-results).

140 Note that this repository *does not include* the input cases files and output ensemble
141 forecasts (approximately 2.3 Gb). These are provided in a separate data repository:
142 [doi:10.26188/29434298](https://doi.org/10.26188/29434298).

143 **3 Results**

144 We begin with an overview of the model inclusion/exclusion outcomes for each ensemble
145 forecast. We then present an overview of the ensemble forecast performance, and
146 demonstrate that the ensemble forecasts were more reliable than the forecasts generated
147 by each individual model. We conclude this section by detailing how the ensemble fore-
148 casts supported public health activities during each phase of the study period (defined by
149 predominant circulation of different COVID-19 variants, as illustrated in [Figure 1](#)) and
150 highlight ways in which we improved forecast visualisations and reporting over the study
151 period.

152 **3.1 Models included in each ensemble forecast**

153 As shown in [Table 3](#), all models were regularly included in the ensemble forecasts. The
154 DST and UoM teams often contributed forecasts generated from multiple model itera-
155 tions, and when models were excluded from the ensemble it was most often *a single model*
156 *iteration* from either team. The most common reason for exclusion was for a model to
157 exhibit large and systematic biases in the forecasts produced in previous weeks (i.e., retro-
158 spective posterior predictive checks). And in several instances the UoA team encountered
159 computational issues and were unable to provide forecasts in a timely manner.

160 **3.2 Summary of forecast performance**

161 Ensemble forecasts for the months of March, April, and May 2022 are shown in [Figure 3](#)
162 for four of the eight Australian jurisdictions (forecasts for the remaining jurisdictions are
163 shown in figures S3 and S4 in appendix S1). These example forecasts demonstrate key
164 features that were exhibited by the ensemble forecasts over the entire study period (2020–
165 2022). In brief, the ensemble was generally in good agreement with the data. Consistent
166 with findings from COVID-19 forecasting studies in other countries [[6–11](#), [28](#), [29](#)], it was
167 an ongoing challenge to predict the timing of change points. The forecasts exhibited
168 a tendency to overshoot peaks, and to undershoot the onset and ending of individual

Strain	Forecasts	DST	MU	UoA	UoM	# Models
Pre-Delta	302	—	100.0%	93.7%	100.0%	2–3 (2.99)
Delta	183	4.4%	100.0%	100.0%	100.0%	3–4 (3.04)
Omicron BA.1	94	84.0%	100.0%	57.4%	100.0%	2–4 (3.41)
Omicron BA.2/BA.5	208	96.2%	96.2%	7.7%	100.0%	2–4 (3.00)
Multiple variants	89	100.0%	100.0%	—	98.9%	2–3 (2.99)
	876	376	868	536	875	2–4 (3.03)

Table 3: The percentage of ensemble forecasts for which each model was included, and the number of models included in each ensemble forecast (minimum, maximum, and mean), reported separately for each period in which particular strain(s) dominated. Note that models could be excluded from the ensemble for various reasons, from being assessed as inappropriately sensitive to trends in the most recent data, or otherwise producing suspect predictions, to being unavailable due to technical issues and/or delays in model development (see [subsection 2.3](#)).

169 waves. In particular, the forecasts tended to predict substantially larger peaks than were
 170 ultimately observed. However, as shown in the 4-week forecasts in [Figure 3](#), the observed
 171 peaks still fell within the forecast credible intervals. The extremely broad intervals for
 172 the Victorian forecast for 15 March 2022 arise from a single model, which was assessed as
 173 highly uncertain but consistent with the data, and included in the ensemble that week.

174 Ensemble forecast coverage (shown in [Figure 4](#)) was reasonable across most pandemic
 175 waves. Coverage is the probability that a credible interval will contain the true value,
 176 and ideally an X% credible interval will include the true value X% of the time. The best
 177 coverage was observed during the “Pre-Delta” and “Delta” periods where, for example,
 178 almost exactly 50% of the ensemble 50% credible intervals included the true value. During
 179 this time, Australian jurisdictions responded rapidly to local disease activity in pursuit
 180 of strong suppression [[14](#)], and while the precise impact of these interventions was chal-
 181 lenging to predict, there was only limited local COVID-19 transmission and case counts
 182 remained low. Australia then pivoted from pursuing strong suppression to reopening and
 183 accepting substantial levels of local COVID-19 transmission. In the subsequent “Omicron
 184 BA.1” period, the models failed to anticipate the massive surge and rapid decline in daily

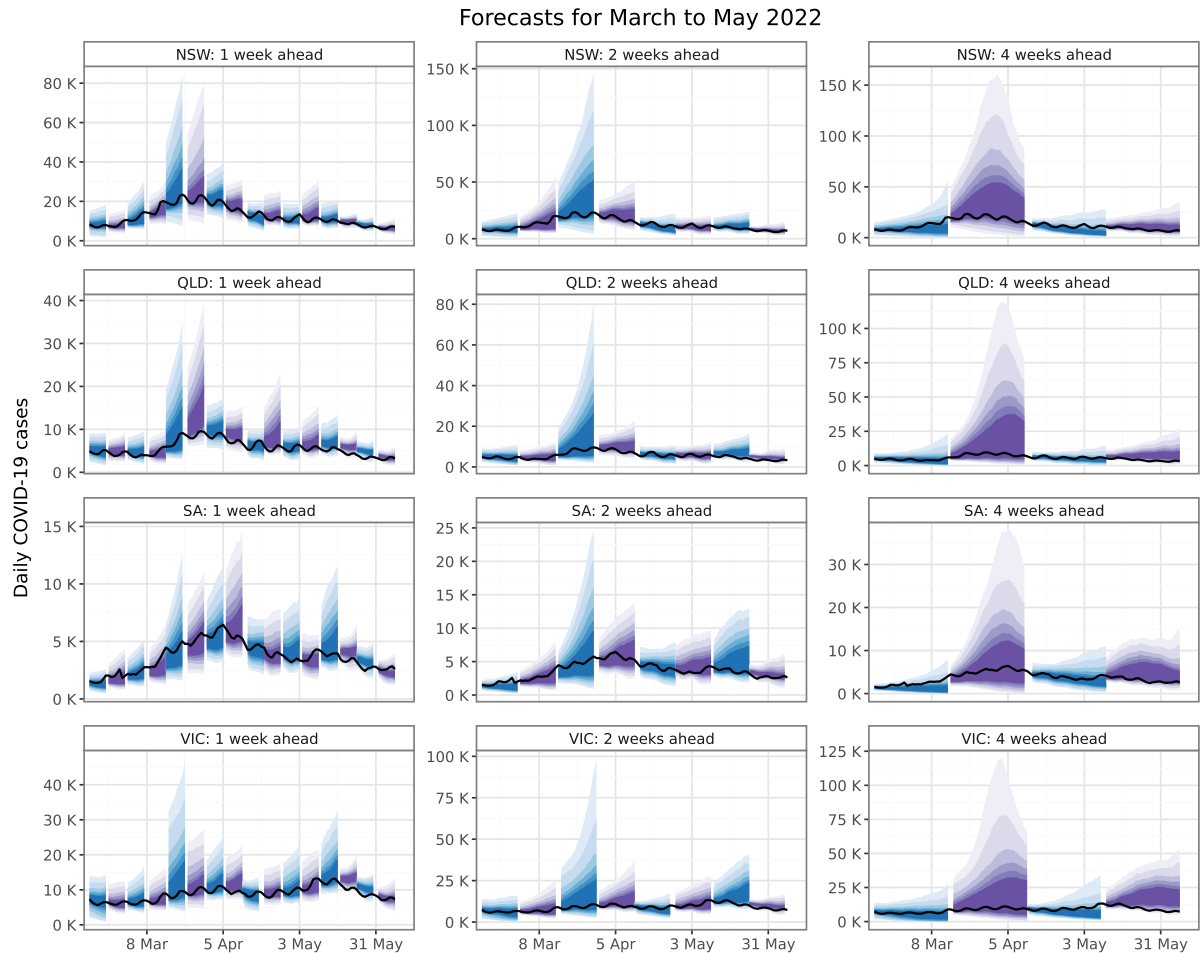


Figure 3: Ensemble forecasts over the months of March, April, and May 2022, shown for four Australian jurisdictions: New South Wales (NSW); Queensland (QLD); South Australia (SA); and Victoria (VIC). The left columns shows 1-week forecasts for each ensemble forecast, the middle column shows 2-week forecasts for every second ensemble forecast, and the right column shows 4-week forecasts for every fourth ensemble forecast. Shaded regions illustrate the 50%, 60%, 70%, 80%, 90%, and 95% credible intervals, and black lines depict the data as reported at the end of the study period.

Jurisdiction	4 weeks	3 weeks	2 weeks	1 week
ACT	-	76.0%	78.4%	86.9%
NSW	-	69.4%	77.9%	83.5%
NT	-	74.1%	73.9%	78.0%
QLD	-	66.8%	73.4%	76.5%
SA	-	71.2%	74.3%	76.4%
TAS	-	75.2%	76.8%	83.7%
VIC	-	67.6%	70.4%	81.5%
WA	-	71.4%	75.9%	81.1%
Mean CRPS:	0.501	0.395	0.288	0.184

Table 4: The improvement in ensemble forecast CRPS for a given observation, as the lead time becomes shorter (i.e., as the forecasting date approaches the observation date). Each cell shows the percentage of predicted distributions for which the CRPS decreased, relative to the predictions made one week earlier. The bottom row shows the mean ensemble forecast CRPS by lead time (calculated for all forecasts, not only those where CRPS decreased).

185 case counts; forecasts were overly confident and exhibited a tendency to first undershoot
186 and then overshoot the true case counts. Forecast coverage improved markedly for the
187 “Omicron BA.2/BA.5” and “Multiple variants” periods, with a slight tendency towards
188 being overly uncertain. For periods where the individual models exhibited marked dif-
189 ferences in forecast coverage, the ensemble forecast coverage was similar to, if not better
190 than, the coverage of the best individual model.

191 As might be expected, observations that lay further into the future were harder to
192 predict. Forecast performance was highest for short lead times, and steadily decreased as
193 the lead time increased. This was true not only on aggregate, but also for the majority
194 of individual observations. As shown in [Table 4](#), for a given observation the forecast
195 performance consistently improved from the initial prediction (when the observation was
196 22–28 days ahead) to the final prediction (when the observation was 1–7 days ahead).

197 Forecasts exhibited substantial biases around epidemic peaks. This is illustrated in
198 [Figure 5](#), which shows the bias for each model and the ensemble for forecasts generated in
199 the 4 weeks leading up to, and the 4 weeks after, each observed peak. Forecasts generated

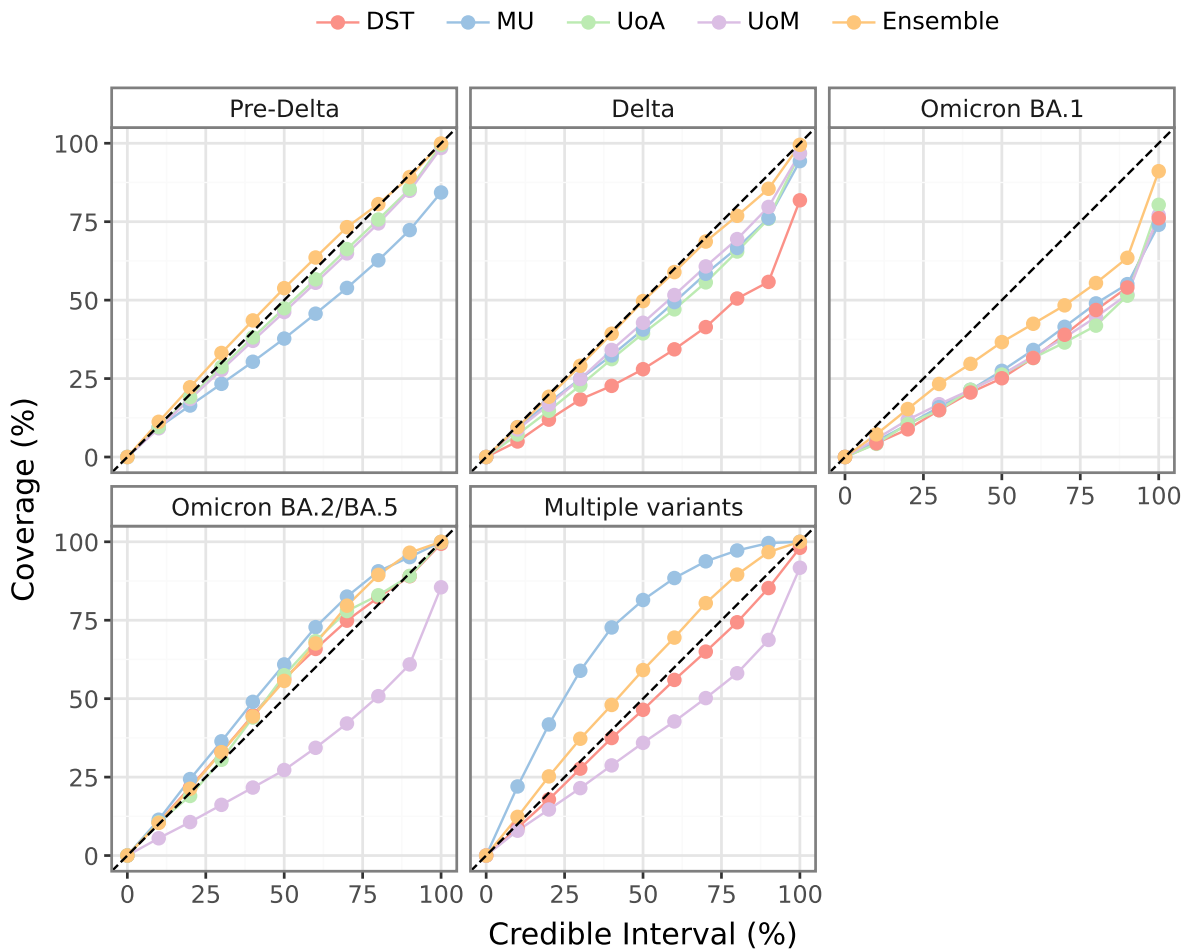


Figure 4: Observed coverage of the forecast credible intervals for each individual model, and for the ensemble, with respect to the ground truth case counts as reported after the end of the study period. Results are reported with respect to each dominating strain (refer to Figure 1). Dashed lines indicate perfect coverage. Points below the dashed lines indicate overly certain forecasts, points above the dashed lines indicate overly uncertain forecasts.

200 2–4 weeks prior to observed peaks undershot the data (negative bias) more often than
201 they overshot the data (positive bias). In contrast, forecasts generated from 1 week prior
202 to 1 week after the observed peaks had a tendency to overshoot the data (positive bias),
203 which highlights the challenge of identifying peaks in real-time when surveillance data are
204 subject to ascertainment biases and reporting delays [12].

205 No model consistently produced the least biased forecasts for weeks preceding or fol-
206 lowing the observed peaks. When there was a wide spread in bias, the ensemble forecast
207 was often less biased than most, if not all, of the individual model forecasts. Similar
208 trends are evident in individual model and ensemble forecast performance (see appendix
209 S5); no model consistently produced the most accurate forecasts for weeks preceding or
210 following the observed peaks, and the ensemble tended to mostly out-perform individual
211 models.

212 While the ensemble forecasts struggled to accurately predict peak sizes and timing
213 (evident in the measure of forecast bias, discussed above), in most weeks preceding or
214 following an observed peak, a proportion of the forecast trajectories were positively corre-
215 lated with the pre-peak and post-peak case counts (appendix S5). This indicates that the
216 forecasts could at least partially characterise the qualitative trends in future case counts
217 (i.e., whether case counts would increase, decrease, or remain stable).

218 **3.3 Relative model performance**

219 Recall that at each week of the study period, sample trajectories of daily case counts were
220 provided over four-week horizons for each Australian jurisdiction. For each individual
221 daily case count prediction, we ranked the individual models and the ensemble using
222 CRPS on log-transformed values [30]. The results for days where at least one case was
223 reported are shown in Figure 6. Collectively, the individual models were more likely
224 than the ensemble to be the top-ranked forecast, but no single model dominated the top
225 ranking. The strength of the ensemble was that it was *most often the best or second-*
226 *best* forecast. Skill scores for each model, relative to the ensemble, further reinforce the
227 finding that no model consistently out-performed the ensemble (Figure 7). The model

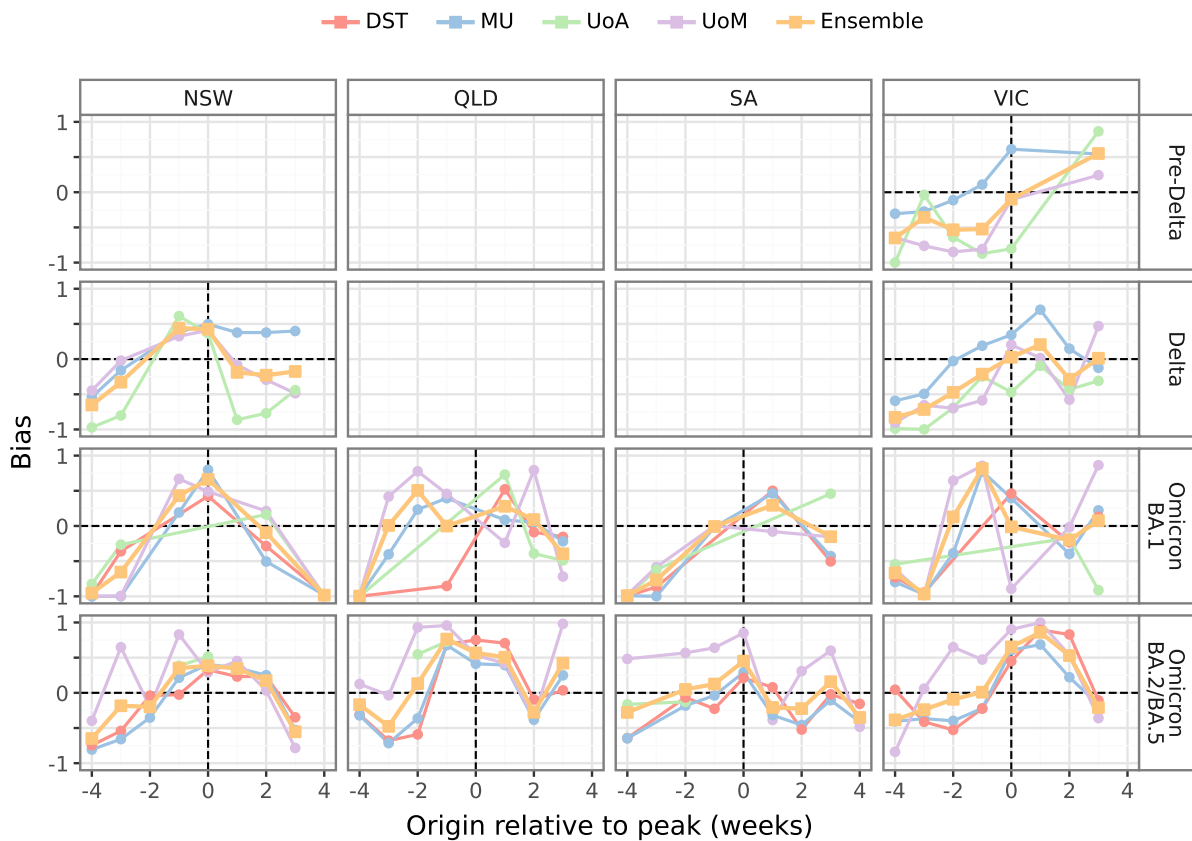


Figure 5: Forecast bias for each model and the ensemble, shown for forecast origins within 4 weeks either side of the largest peak (minimum 200 cases) observed in four jurisdictions (columns) for each dominant strain (rows). Positive bias indicates a tendency to overshoot the data, negative bias indicates a tendency to undershoot the data. The “Multiple variants” period is not included in the figure, because case counts were generally flat and the observed peaks occurred towards the end of the study period. See appendix S5 for forecast bias around these peaks over all eight jurisdictions.

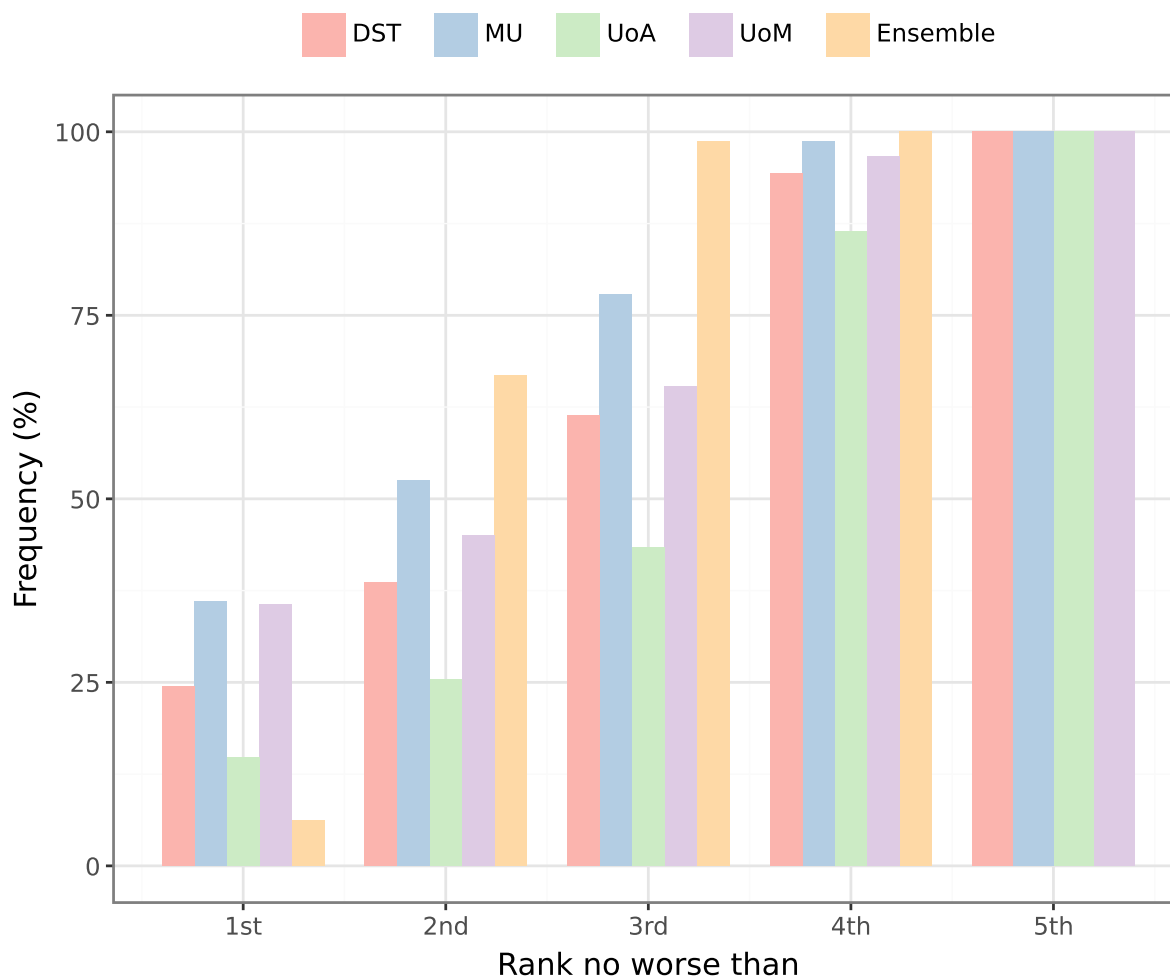


Figure 6: Cumulative rankings for each individual model and for the ensemble, for days where at least one case was reported, calculated using CRPS on log-transformed values. While the ensemble was rarely the top-ranked model, it was *at least* 2nd-best approximately two-thirds of the time (66.7%), and was *almost always* in the top three (98.7%).

228 rankings for each ensemble forecast (appendix S2) also demonstrate that there were no
229 obvious predictors of which model would perform the best for a given time period or
230 epidemiological context.

231 3.4 Pre-Delta: May to October 2020

232 We now highlight how the forecasts supported public health activities in each phase of
233 the study period, beginning with the Pre-Delta phase. This first period of sustained local
234 COVID-19 transmission began in mid-2020, where the majority of cases occurred in the

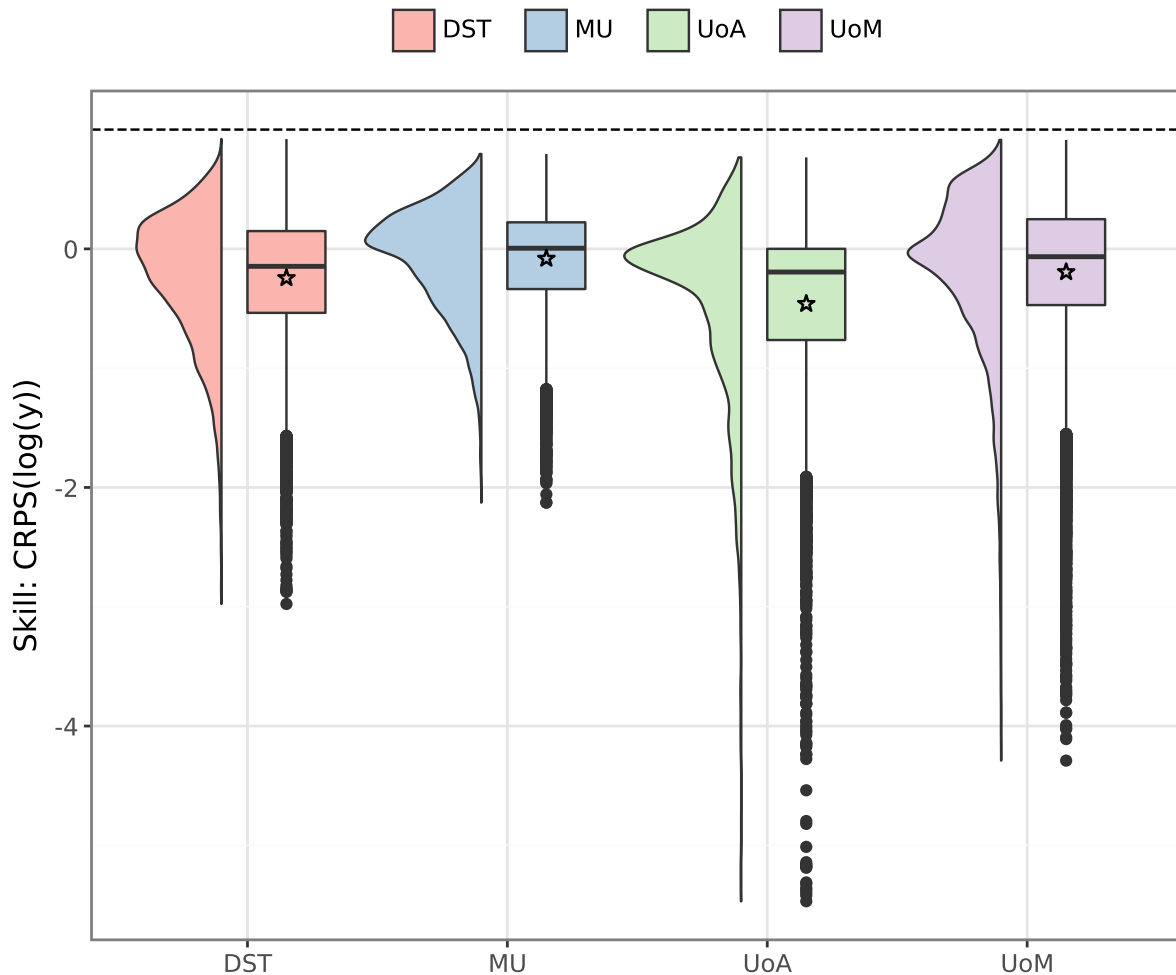


Figure 7: Distribution of skill scores for each individual model, measured against the ensemble, for days where at least one case was reported (around 53% of all observations). Skill scores were calculated using CRPS on log-transformed values [30]. Stars indicate mean skill scores. The MU model performed better than the other models, with a median skill score of 0.002. Mechanistic models performed well for extended periods of zero cases (median skill scores of 0.75–0.97, not shown here) but were otherwise outperformed by the ensemble, with median scores of -0.07 (UoM), -0.14 (DST), and -0.19 (UoA).

235 state of Victoria. Ensemble forecasts for this period are presented in appendix S1. The
236 performance of the UoM forecasts over this second wave has already been reported [12],
237 and we begin by comparing the ensemble forecasts to the UoM forecasts over this wave.

238 In the weeks prior to the peak in daily cases (3 August 2020), the upper bounds of the
239 UoM forecasts substantially overshoot the true peak, and predicted sustained exponential
240 growth in cases. This exponential growth was tempered in the ensemble, which averaged
241 over three models (MU, UoA, and UoM). Accordingly, the ensemble outperformed the
242 UoM forecasts (which had a mean skill score of -0.37, see appendix S4), even though the
243 ensemble forecasts tended to undershoot the data. By late July, we identified that case
244 ascertainment and reporting delays had substantially increased, and our right-truncation
245 adjustments were under-estimating the true case counts in the most recent days prior to
246 each forecast. In the weeks after the peak, the UoM forecasts confidently predicted a
247 sustained decrease in cases, while the MU and UoA models exhibited broader credible
248 intervals and tended to overshoot the data. As a result, the UoM forecasts performed
249 better than the ensemble forecasts (mean skill score of 0.31, see appendix S4).

250 Daily case incidence in Victoria steadily decreased from the August peak. In early
251 September the Victorian government announced a gradual easing of restrictions, subject
252 to reaching specific 14-day moving average case thresholds. Because the ensemble forecast
253 comprised individual trajectories from each contributing model, we were able to calculate
254 the proportion of all trajectories that satisfied a given 14-day case threshold on each date
255 in the forecast horizon, and reported this as the daily probability of achieving each tar-
256 get threshold. These predictions were provided to the Victorian government throughout
257 September and October.

258 **3.5 Delta: June to December 2021**

259 The onset of the Delta variant in June 2021 marked the first instance of sustained local
260 COVID-19 transmission for most Australian jurisdictions, and public health responses
261 focused on preventing local transmission. Accordingly, daily case counts remained very
262 low in most jurisdictions, only exceeding 25 cases per day in the Australian Capital

263 Territory (peak of 51), New South Wales (peak of 1,495), and Victoria (peak of 1,955).
264 Ensemble forecasts for this period are presented in appendix S1.

265 Perhaps the single greatest value of the forecasts in this period was to demonstrate
266 how rapidly case counts could increase if local transmission was left unchecked.

267 In May 2021, New South Wales Health set a target of achieving 80% coverage of the
268 adult population with two vaccination doses by the end of December 2021 [31]. This was
269 followed in August 2021 by an interim target of 70% coverage of the adult population
270 with two doses, to encourage vaccine uptake and to begin easing restrictions for fully
271 vaccinated people [31]. As a consequence, there was substantial vaccine roll-out over the
272 4-week forecast horizons from August onward. This was accompanied by rapid model
273 development from the DST and UoM teams, with the effects of vaccination incorporated
274 into these models in September 2021. The statistical MU model did not require such
275 adjustments.

276 The forecasts were generally in very good agreement with the data in each jurisdiction.
277 In particular, the mean CRPS values for New South Wales and Victoria were consistently
278 lower than for the Pre-Delta wave in Victoria (see appendix S5).

279 **3.6 Omicron BA.1: January to March 2022**

280 This wave coincided with a national pivot from pursuing strong suppression to reopening
281 and management of substantial levels of local COVID-19 transmission. Ensemble forecasts
282 for this period are presented in appendix S1.

283 The forecasts struggled to predict the massive surge and subsequent decline in daily
284 case counts, as demonstrated by decreased coverage (Figure 4), increased bias (Figure 5,
285 appendix S3), and worse calibration than other waves (appendix S6).

286 This was due, at least in part, to mechanistic models failing to account for significant
287 reduction in vaccine protection against Omicron, rapid changes in case ascertainment [32],
288 and reduced mixing due to school holiday effects over December and January. Also in
289 January, the vaccine roll-out began for children aged 5–11 years, and booster doses were
290 recommended for at-risk individuals [33].

291 A primary concern in all jurisdictions was the timing of the epidemic peak, because the
292 substantial increase in cases was impacting workforce capacity and causing disruptions to
293 food supply chains, which in turn prompted panic buying [34]. The forecasts consistently
294 predicted that the peaks would occur around 2 weeks later than they actually occurred
295 (and for the mechanistic models, this was robust to adjustments such as relaxing assump-
296 tions regarding case ascertainment and immunity). Despite this inaccuracy, the forecast
297 predictions that case activity would peak and begin to decrease in a matter of weeks
298 remained an important and useful message for government.

299 **3.7 Omicron BA.2/BA.5: April to August 2022**

300 This period saw the gradual replacement of Omicron BA.1 with Omicron BA.2 and BA.5,
301 and case counts in all jurisdictions steadily decreased after the large Omicron BA.1 peaks.
302 Ensemble forecasts for this period are presented in appendix S1.

303 The forecasts reported on 11 June, generated from daily case counts up to 31 May
304 (inclusive), predicted downwards trends in all jurisdictions, consistent with the trends in
305 the most recent case counts. On 16 June we received additional genomic data from New
306 South Wales that allowed us to estimate the transmission advantage of Omicron BA.5,
307 relative to Omicron BA.1 and BA.2. By incorporating this transmission advantage into
308 the mechanistic models, our next forecasts predicted upwards trends in all jurisdictions
309 *despite no such trend in the reported aggregate data*. Consistent with these predictions,
310 daily case counts began to increase later in June, and all jurisdictions experienced a peak
311 in July (see figure S4).

312 By predicting this inflection point before it occurred, and by doing so based on appro-
313 priate interpretation of relevant local data, these forecasts helped convince public health
314 stakeholders that the current downwards trend in cases would not be sustained.

315 **3.8 Multiple variants: September to December 2022**

316 Following the peak and decline of Omicron BA.5, Australia experienced circulation of
317 numerous COVID-19 variants, with no single variant becoming dominant. Data from

318 international contexts provided evidence that many of these variants had growth advan-
319 tages over Omicron BA.1, BA.2, and BA.5, although it was unclear how these estimates
320 might translate to Australia’s immune landscape. Ensemble forecasts for this period are
321 presented in appendix S1.

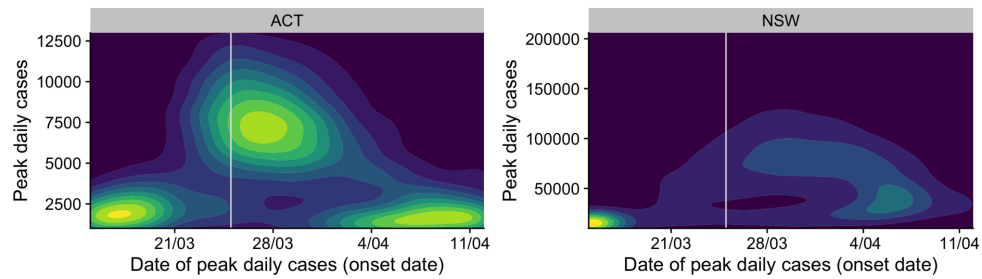
322 Relative to the preceding Omicron BA.2/BA.5 period, the forecasts exhibited similarly
323 good coverage (Figure 4) and performance (appendix S8). When case counts began to
324 increase in November, the forecasts were confident that the epidemic peaks would be
325 similar in size, or smaller than, the Omicron BA.5 peaks. This was a reassuring message
326 that would prove to be accurate, and brings us to the end of our study period.

327 **3.9 Communication of outputs**

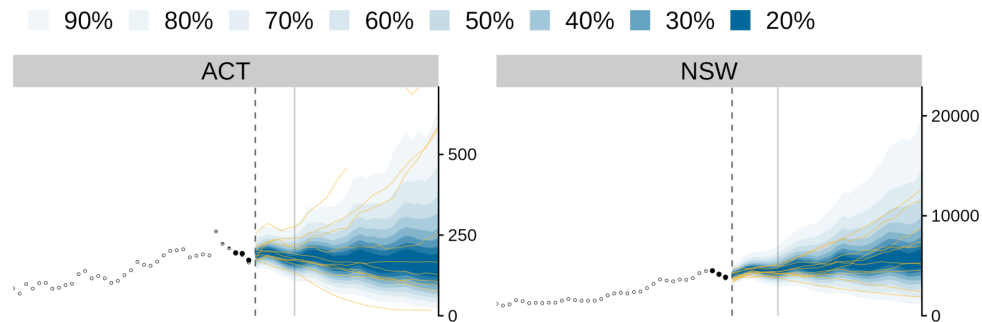
328 Over the 3.5 years of work summarised here, an ongoing concern was to ensure that model
329 predictions and other quantitative outputs were reported in such a way that our stakehold-
330 ers (spanning state, territory, and national governments and public health committees)
331 would be able to interpret them appropriately and accurately act on and communicate
332 their implications. While we did not have the capacity to undertake formal assessments of
333 how the 152 weekly situation reports that we produced over this study period were inter-
334 preted, as described in McCaw and Plank [35] we did have open communication channels
335 with our stakeholders and were able to explore and refine our reporting and visualisations
336 in an iterative manner.

337 Figure 8 shows several examples of visualisations included in these reports. For exam-
338 ple, our primary form of communicating the forecasts was daily credible intervals (see, e.g.,
339 Figure 3). However, this form can obscure the predicted size and timing of an epidemic
340 peak, and so we explored the use of density plots to extract these features from individual
341 forecast trajectories (Figure 8a). Another simple, yet surprisingly useful, modification
342 to the daily credible interval figures was to overlay several randomly-selected forecast
343 trajectories (Figure 8b), which helped to avoid the credible interval contours being mis-
344 interpreted as case count trajectories. Finally, plotting past forecasts against the most
345 recent data (Figure 8c) was a simple way of communicating recent forecast performance
346 that provided stakeholders with a qualitative basis for deciding how much trust to place

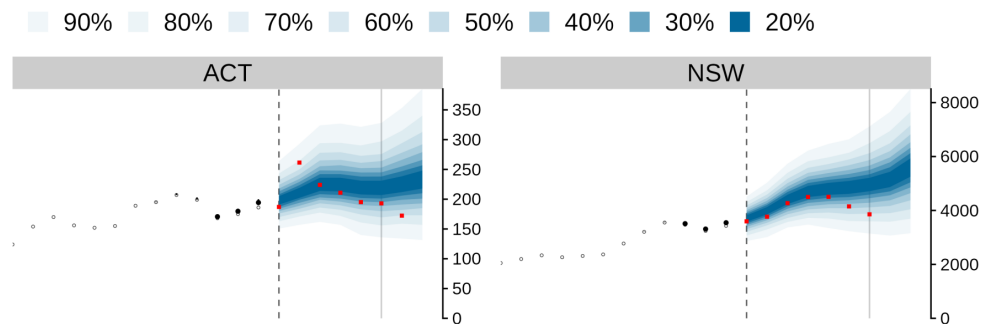
347 in the current forecasts.



(a) Density plots showing predicted peak timing and peak case incidence, from the report produced on 24 March 2022.



(b) Ensemble forecasts of new daily local cases, showing credible intervals and sample trajectories, from the report produced on 25 November 2022.



(c) Retrospective evaluation of ensemble forecasts from the previous week against new data (red points), from the report produced on 25 November 2022.

Figure 8: Example figures from weekly situation reports.

348 4 Discussion

349 4.1 Principal findings

350 Similar to findings reported in COVID-19 forecasting studies in other countries [6–11, 28,
351 29], the use of an ensemble to combine forecasts from multiple models improved forecast
352 performance and reliability (Figures 4–6). Our ensemble forecasts provided information
353 that supported public health responses throughout the study period, from the early peri-
354 ods of preventing local transmission, to the later periods of sustained local transmission
355 and concerns about the burden placed on healthcare systems. Notable outcomes during
356 the study period include:

- 357 • Pre-Delta: we used the forecasts to predict when 14-day average case targets would
358 be achieved, which would trigger the gradual easing of mobility restrictions in the
359 state of Victoria.
- 360 • Delta: this was the first instance of sustained local transmission for most jurisdic-
361 tions, and the forecasts were useful for illustrating how rapidly case counts could
362 increase if transmission was not curtailed.
- 363 • Omicron BA.1: the peak occurred 1–2 weeks earlier than the forecasts predicted,
364 but forecast predictions that activity would peak and then decline in a matter of
365 weeks was reassuring in the face of workplace absenteeism that disrupted supply
366 chains.
- 367 • Omicron BA.2/BA5: the forecasts were able to correctly predict an increase in
368 cases while the current case counts were decreasing, and this was a particularly
369 useful message for health protection committees.
- 370 • Multiple variants: the forecasts accurately predicted that the peaks in late 2022
371 would not be larger than the Omicron BA.5 peaks.

372 4.2 Study strengths

373 The ensemble approach allowed us to readily incorporate forecasts contributed by teams
374 across Australia, and provided a framework for rapid evaluation of new model iterations
375 for potential inclusion in the ensemble. The weekly performance evaluations and model
376 inclusion/exclusion decisions relied on measures of forecast error (using CRPS) under the
377 assumption that existing policies would persist for the entire forecast horizon. However, as
378 we have described above, the ensemble forecasts influenced policy decisions throughout the
379 study period, and some of these decisions directly impacted local COVID-19 transmission
380 and/or case ascertainment. Accordingly, when policy decisions were understood to have
381 influenced the case data reported over a forecast horizon, we factored these effects into
382 our analyses. In particular, where the difference between forecasts and reported data were
383 consistent with the likely effects of the policy decisions, we considered this a successful
384 outcome [36].

385 We deliberately chose to use equal-weight ensemble forecasts and, as we explain in the
386 methods section, each model was evaluated against the most recent data rather than com-
387 peting against the other models for inclusion in the ensemble [26]. As we have discussed in
388 the results section, a model's ranking in recent weeks was not a reliable predictor of that
389 model's ranking in future weeks. The strongest finding we have regarding the rankings
390 is that the ensemble was either the best, or second-best, more often than any individ-
391 ual model. Both findings are consistent with ensemble COVID-19 forecast evaluations in
392 other countries [9, 10].

393 Finally, we note here that the ensemble forecasts presented here also served as an input
394 for COVID-19 hospital bed occupancy forecasts (December 2021 to December 2023) that
395 further supported Australia's public health responses [18].

396 4.3 Study limitations

397 While pursuing the prevention of local transmission, Australian jurisdictions maintained
398 high testing levels and the proportion of infected persons that were identified as cases was
399 likely to be both very high, and to remain relatively constant. The ensemble forecasts

400 performed very well during this period (the “Pre-Delta” and “Delta” phases).

401 When Australia transitioned to re-opening, case ascertainment was substantially re-
402 duced, and was challenging to estimate in near-real-time [32]. By February 2022, more
403 than 94% of people over the age of 16 were fully vaccinated [37], and while the mechanistic
404 models were subject to numerous adaptations and refinements to account for vaccination
405 coverage, booster vaccinations, and waning immunity, throughout 2022 it became more
406 challenging to estimate population susceptibility to the emerging variants of concern. This
407 was due to an increasingly complex population immune landscape, resulting from hetero-
408 geneous levels of vaccination and immunising exposure, and challenges in estimating the
409 immunogenicity of emerging variants.

410 However, despite these challenges, the inclusion of both mechanistic and statistical
411 models in the ensemble substantially improved the forecast performance and reliability
412 across the entire study period. The only period where the forecast did not capture the case
413 data was the rapid emergence and decline of Omicron BA.1, which occurred as Australia
414 pivoted from pursuing strong suppression to reopening and accepting substantial levels of
415 local transmission — a transition that was inherently challenging to predict.

416 **4.4 Meaning and implications**

417 Our ensemble COVID-19 forecasts were produced under contract with the Australian
418 Government Department of Health and Aged Care, and participation was limited to a
419 small number of institutes. In contrast, the United States of America and Europe both es-
420 tablished public COVID-19 forecast hubs that had open submission policies and included
421 all submissions that complied with the hubs’ technical requirements in their ensemble
422 forecasts [9, 10]. This open nature resulted in large numbers of participating teams, with
423 67 teams contributing US forecasts (July 2020 to December 2021 [10]), and 48 teams
424 contributing European forecasts (March 2021 to March 2022 [9]). Both hubs reported
425 that this open nature also limited the possibility to understand drivers of forecast per-
426 formance, with many teams participating at different times, participating intermittently,
427 and providing varied and/or limited descriptions of their methods [9, 10].

428 Forecast evaluations from both hubs reported findings that are broadly consistent with
429 those reported here. Forecasts performed well in periods of stable behaviour, but struggled
430 at longer horizons around inflection points, and individual models varied widely in their
431 ability to account for new COVID-19 variants. The ensemble forecasts were consistently
432 among the best-performing forecasts across all horizons and locations, and weighting
433 individual models by their past performance did not improve the ensemble performance.
434 The US hub reported that forecast prediction intervals were generally over-confident and
435 had low coverage, particularly when case numbers were changing rapidly [10], similar to
436 our findings for the Omicron BA.1 wave (Figure 4).

437 The ensemble forecasts presented here played an important role in supporting Aus-
438 tralian public health decision-making over a wide range of epidemiological and policy con-
439 texts. Consistent with reflections from modelling and data analysis communities around
440 the world [35, 38–40], frequent communication between modellers and public health stake-
441 holders — and the mutual understanding and trust that this fostered — was integral to
442 the utility of these ensemble forecasts. Being organised as a consortium, with several
443 members sitting on key public health committees, meant that there were direct lines of
444 communication between decision makers and modellers, and this in turn expedited the
445 prototyping and development of effective analyses and communications.

446 More broadly, the collective role of our consortium as a means for weekly peer-review
447 of model forecasts, and the weekly decision of which model iteration(s) to include, directly
448 supported rapid model development and evaluation, while also ensuring that only “known
449 good” model iterations were contributing to the ensemble forecast. Having common
450 performance targets and evaluation processes for all models, conducted openly within
451 the consortium each week, fostered mutual collaboration. This collaborative structure
452 also helped to avoid duplication of effort, which was extremely beneficial at times of high
453 stress, urgent delivery schedules, and unsustainable workloads. Such an approach may be
454 challenging to adapt to the scale of the US and European forecast hubs.

455 A final benefit of this organisation, and the small size of our consortium relative to
456 similar groups overseas, was retaining the human element in selecting which model itera-

457 tion(s) to include in the ensemble, rather than using an arbitrary quantitative threshold or
458 simply including all available forecasts in the ensemble. This allowed us to explain features
459 of each ensemble forecast, and the underlying rationale for these features in terms of the
460 contributing models and their assumptions, which helped our public health stakeholders
461 to interpret the forecasts appropriately.

462 5 Ethics statement

463 The study was undertaken as urgent public health action to support Australia's COVID-
464 19 pandemic response. The study obtained data under the National Health Security
465 Agreement for the purposes of national communicable disease surveillance. Contractual
466 obligations established strict data protection protocols, as agreed between the University
467 of Melbourne and sub-contractors and the Australian Government Department of Health
468 and Aged Care. Oversight and approval for use in supporting Australia's pandemic re-
469 sponse, and for publication, were provided by the data custodians (represented by the
470 Communicable Diseases Network of Australia, CDNA). The ethics of the use of these
471 data for these purposes, including publication, was agreed by the Department of Health
472 and Aged Care with CDNA. All methods were carried out in accordance with the relevant
473 guidelines and regulations.

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