

## **COVID-19 Weekly Epidemiological Update**

#### Edition 69, published 7 December 2021

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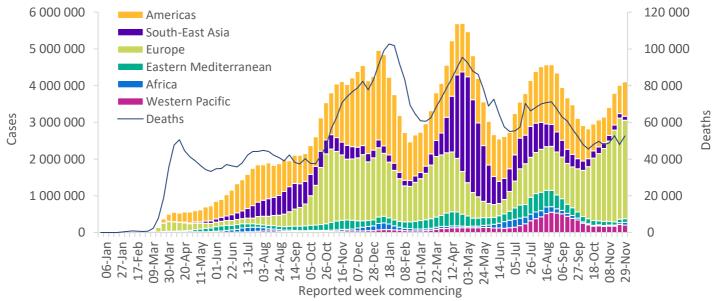
#### **Global overview**

#### Data as of 5 December 2021

Globally, weekly case incidence plateaued this week (29 November - 5 December 2021), with over 4 million confirmed new cases reported, similar to the number reported in the previous week's figures. However, new weekly deaths increased by 10% as compared to the previous week, with over 52 500 new deaths reported. As of 5 December, nearly 265 million confirmed cases and over 5.2 million deaths have been reported globally.

The African Region and the Region of the Americas reported increases in new weekly cases of 79% and 21%, respectively, while the Western Pacific and South-East Asia regions both reported decreases of 10%. The number of new weekly cases reported by the European and Eastern Mediterranean regions were similar to the numbers reported in the previous week. New weekly deaths increased by 49% in the South-East Asia Region and 38% in the Region of the Americas, while the weekly deaths decreased in the African and Eastern Mediterranean Regions by 13% and 8%, respectively. The number of new deaths were similar to those reported in the previous week in both the European and the Western Pacific regions.





<sup>\*\*</sup>See Annex 2: Data, table, and figure notes

The regions reporting the highest weekly case incidence per 100 000 population continue to be the European Region (288.0 new cases per 100 000 population) and the Region of the Americas (91.4 new cases per 100 000 population). Both regions also reported the highest weekly incidence in deaths of 3.1 and 1.3 per 100 000 population, respectively while <1 new death per 100 000 was reported in all other regions.

The highest numbers of new cases were reported from the United States of America (752 394 new cases; a 30% increase), Germany (396 429 new cases; similar to the previous week's figures), the United Kingdom (310 696 new cases; similar to the previous week's figures), France (283 500 new cases; a 49% increase) and the Russian Federation (231 240 new cases; similar to the previous week's figures).

Table 1. Newly reported and cumulative COVID-19 confirmed cases and deaths, by WHO Region, as of 5 December 2021\*\*

WHO Region	New cases in last 7 days (%)	Change in new cases in last 7 days *	Cumulative cases (%)	New deaths in last 7 days (%)	Change in new deaths in last 7 days *	Cumulative deaths (%)
Europe	2 687 257 (65%)	-3%	88 925 399 (34%)	28 990 (55%)	-2%	1 569 599 (30%)
Americas	935 062 (23%)	21%	97 679 255 (37%)	12 987 (25%)	38%	2 360 315 (45%)
Western Pacific	199 495 (5%)	-10%	10 370 429 (4%)	3 220 (6%)	2%	144 204 (3%)
South-East Asia	109 044 (3%)	-10%	44 638 985 (17%)	5 324 (10%)	49%	711 660 (14%)
Eastern Mediterranean	94 724 (2%)	0%	16 846 148 (6%)	1 622 (3%)	-8%	310 727 (6%)
Africa	79 491 (2%)	79%	6 354 835 (2%)	498 (1%)	-13%	153 275 (3%)
Global	4 105 073 (100%)	2%	264 815 815 (100%)	52 641 (100%)	10%	5 249 793 (100%)

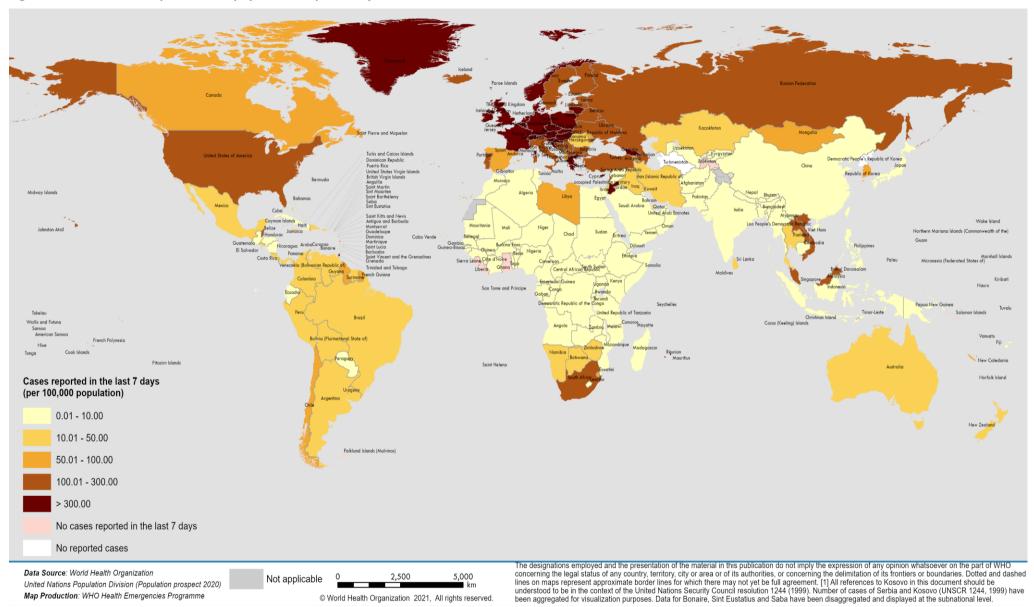
<sup>\*</sup>Percent change in the number of newly confirmed cases/deaths in the past seven days, compared to seven days prior

For the latest data and other updates on COVID-19, please see:

- WHO COVID-19 Dashboard
- WHO COVID-19 Weekly Operational Update and previous editions of the Weekly Epidemiological Update

<sup>\*\*</sup>See Annex 2: Data, table, and figure notes

Figure 2. COVID-19 cases per 100 000 population reported by countries, territories and areas, 29 November - 5 December 2021\*\*



<sup>\*\*</sup>See Annex 2: Data, table, and figure notes

Map Production: WHO Health Emergencies Programme

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Turks and Caicos Islands Deaths reported in the last 7 days (per 100,000 population) 0.01 - 0.50 0.51 - 1.50 1.51 - 3.00 3.01 - 6.00 No deaths reported in the last 7 days No reported cases The designations employed and the presentation of the material in this publication do not imply the expression of any opinion whatsoever on the part of WHO concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted and dashed lines on maps represent approximate border lines for which there may not yet be full agreement. [1] All references to Kosovo in this document should be understood to be in the context of the United Nations Security Council resolution 1244 (1999). Number of cases of Serbia and Kosovo (UNSCR 1244, 1999) have been aggregated for visualization purposes. Data for Bonaire, Sint Eustatius and Saba have been disaggregated and displayed at the subnational level. Data Source: World Health Organization 2,500 5,000 Not applicable United Nations Population Division (Population prospect 2020) Map Production: WHO Health Emergencies Programme © World Health Organization 2021, All rights reserved.

Figure 3. COVID-19 deaths per 100 000 population reported by countries, territories and areas, 29 November - 5 December 2021\*\*

<sup>\*\*</sup>See Annex 2: Data, table, and figure notes

#### Special Focus: Update on SARS-CoV-2 Variants of Interest and Variants of Concern

WHO, in collaboration with national authorities, institutions and researchers, routinely assesses if variants of SARS-CoV-2 alter transmission or disease characteristics, or impact effectiveness of vaccines, therapeutics, diagnostics or public health and social measures (PHSM) applied by national authorities to control disease spread. Potential Variants of Concern (VOCs), Variants of Interest (VOIs) or Variants Under Monitoring (VUMs) are regularly assessed based on the risk posed to global public health. As evidence becomes available, classifications of variants will be revised to reflect the continuous evolution of circulating variants and their changing epidemiology. Criteria for variant classification, and the current lists of VOCs, VOIs and VUMs, are available on the <a href="https://www.website">WHO Tracking SARS-CoV-2 variants</a> website. National authorities may choose to designate other variants of local interest/concern and are encouraged to investigate and report on the impacts of these variants.

#### Geographic spread and prevalence of VOCs

The current global epidemiology of SARS-CoV-2 is characterized by a predominance of the Delta variant, declining trend in the proportion of Alpha, Beta and Gamma, and the emergence of Omicron which was designated as a <u>Variant of Concern on 26 November</u> (Figures 4 and 5). At present, Omicron cases have been reported in 57 countries across all WHO regions. While most of the cases identified in these countries are currently travel-related, this may change as more information becomes available. Of 899 935 sequences uploaded to <u>GISAID</u> with specimens collected in the last 60 days<sup>i</sup>, 897 886 (99.8%) were Delta, 713 (0.1%) were Omicron, 286 (<0.1%) Gamma, 154 (<0.1%) Alpha, 64 (<0.1%) Beta, and <0.1% comprised other circulating variants (including VOIs Mu and Lambda).

Sub-regional and country-level variation continues to be observed; most notably within some South American countries, where the progression of the Delta variant has been more gradual, and other variants (e.g., Gamma, Lambda, Mu) still contribute a large proportion of reported sequences.

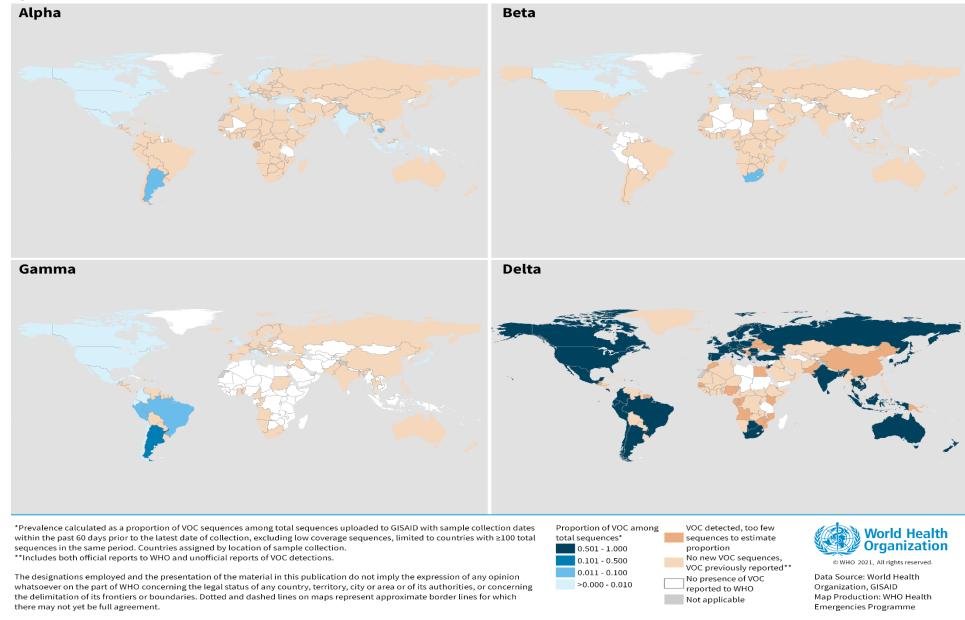
To note, global VOCs distribution should be interpreted with due consideration of surveillance limitations, including differences in sequencing capacities and sampling strategies between countries, as well as delays in reporting.

#### **Additional resources**

- Tracking SARS-CoV-2 Variants
- COVID-19 new variants: Knowledge gaps and research
- Genomic sequencing of SARS-CoV-2: a guide to implementation for maximum impact on public health
- Considerations for implementing and adjusting PHSM in the context of COVID-19

Includes sequences submitted to <u>GISAID</u> with sample collected dates from 3 October to 2 December 2021 (last reported sample at the time of data extraction), excluding low coverage sequences.

Figure 4. Prevalence of Variants of Concern (VOCs) Alpha, Beta, Gamma and Delta in the last 60 days and historic detections, data as of 7 December 2021



Prevalence data based on sequences reported to GISAID, excluding low coverage sequences. See also Annex 1 for reported VOC detections by country/territory/area

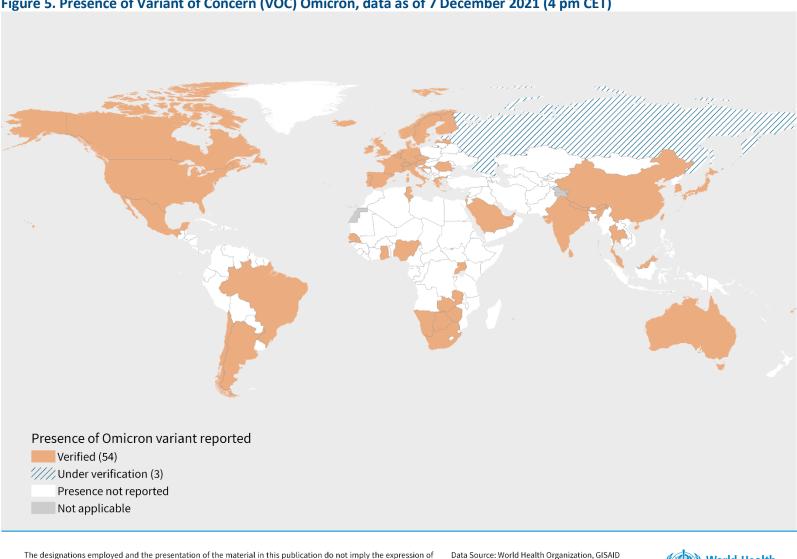


Figure 5. Presence of Variant of Concern (VOC) Omicron, data as of 7 December 2021 (4 pm CET)

The designations employed and the presentation of the material in this publication do not imply the expression of any opinion whatsoever on the part of WHO concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted and dashed lines on maps represent approximate border lines for which there may not yet be full agreement.

Map Production: WHO Health Emergencies Programme



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Presence of the Omicron variant is based on information reported to WHO. It includes countries/territories/areas reporting the detection of VOCs among travellers (e.g., imported cases detected at points of entry), or local cases (detected in the community). See also Annex 1 for reported VOC detections by country/territory/area.

## **Update on Variant of Concern Omicron**

### **Background**

Omicron is the fifth SARS-CoV-2 variant to be designated as a Variant of Concern (VOC) by WHO, following the designation of the Alpha, Beta, Gamma and Delta variants. The first known laboratory-confirmed case of Omicron was identified from a specimen collected on 9 November 2021 in South Africa, with the variant (Pango nomenclature B.1.1.529) first reported to WHO on 24 November. In consultation with the Technical Advisory Group on Virus Evolution (TAG-VE), WHO designated B.1.1.529 as a VOC on 26 November in view of the potential for enhanced transmissibility and/or a degree of immune escape, given the number of mutations (26-32) in the spike protein, as well as concerning initial epidemiological reports from South Africa, including signals of increased risk of reinfection. Here, we present an update on the current situation in terms of the epidemiology and transmissibility, clinical severity, risk of reinfection and the potential impact on diagnostics, vaccines and therapeutics.

As more data are analysed and understood about the potential implications that Omicron may have on the epidemiology, transmissibility, clinical severity, prevention and treatment of SARS-CoV-2 infection, our understanding of this variant will continue to evolve, and we will issue updates as further evidence becomes available.

## **Epidemiology**

In South Africa, where Omicron was first reported, the case incidence of COVID-19 has continued to rise since the second week of November, with 62 021 new cases reported between 29 November and 5 December, a 111% increase compared to the previous week. An increase in the test positivity rate (TPR) has also been seen from 1.2% the week beginning the 7 November, to 22.4% the week beginning the 2 December. An initial increase in incidence in Gauteng province in mid-November was thought in part, to be due to a cluster of cases among students at a university. Very large increases in the weekly incidence of cases have also been seen in some countries neighbouring South Africa including: Eswatini (1990%); Zimbabwe (1361%); Mozambique (1207%), Namibia (681%) and Lesotho (219%). These other countries have very low vaccination coverage ranging from 12.1% of the total population fully vaccinated in Namibia to 26.7% in Lesotho. In South Africa 25.2% of the total population is fully vaccinated. While drivers of these increases remain unknown, it is plausible that spread of Omicron in combination with enhanced testing following the declaration of a VOC, play a role, together with the relaxation of public health and social measures (PHSMs) and sub-optimal immunization coverage.

Among countries in other regions reporting increasing spread of the Omicron variant, hundreds of cases of this variant have now been reported from countries in other regions.<sup>3</sup> Since the last update published on 30 November, additional countries across all six WHO Regions have reported confirmed cases of the Omicron variant. As of 7 December 2021, the Omicron variant has been confirmed in 57 countries. However, given the predominant circulation of the Delta variant in many countries, particularly in countries in the European Region and in the United States of America, it is too early to draw any conclusions about the impact of Omicron will have on the global epidemiology of COVID-19.

## Transmissibility

While there seems to be evidence that the Omicron variant may have a growth advantage over other circulating variants it is unknown whether this will translate into increased transmissibility. Based on several assumptions about

the growth advantage, timing of introduction in the European Region, and population mixing and public health and social measures (PHSM) implementation, the European Centre for Disease Prevention and Control,<sup>4</sup> forecasted that if 1% of SARS-CoV-2 infections are due to the Omicron variant, it will become dominant in Europe, comprising >50% of the new infections, by 1 January 2022, with a growth advantage of >120%; and by 1 March 2022 with a growth advantage of >30%.

Ongoing and planned epidemiological studies, including detailed cluster investigations, contact-tracing and household transmission studies, coupled with neutralization studies from people previously vaccinated or infected and studies of vaccine effectiveness will help improve our understanding of the interplay between increased transmissibility and immune escape as drivers of increased transmission.

### **Clinical severity**

Currently only limited data are available, making it challenging to assess any changes in disease severity with the Omicron variant. As of 6 December, all of the 212 confirmed cases identified in 18 European Union countries for which there was information available on severity were asymptomatic or mild.<sup>3</sup> While South Africa saw an 82% increase in hospital admissions due to COVID-19 (from 502 to 912) during the week 28 November – 4 December 2021, it is not yet known the proportion of these with the Omicron variant.<sup>5</sup> Even if the severity is equal or potentially even lower than for Delta variant, it is expected that hospitalizations will increase if more people become infected and that there will be a time lag between an increase in the incidence of cases and an increase in the incidence of deaths. Further information is needed to fully understand the clinical picture of those infected with the Omicron variant and WHO encourages countries to contribute to the collection and sharing of hospitalized patient data through the WHO COVID-19 Clinical Data Platform.

#### **Risk of reinfection**

Preliminary analysis suggests that the mutations present in the Omicron variant may reduce neutralising activity of antibodies resulting in reduced protection from natural immunity. This may explain why the variant seems to be spreading rapidly in a highly immune population such as South Africa, in which current vaccination coverage in adults is about 35%, but in which seroprevalence levels are estimated to be as high as 60-80% due to past infections, according to recent epidemiological studies and modelling.<sup>6</sup>

A modelling study (pre-print) based on data from nearly three million individuals in South Africa with a laboratory confirmed infection at least 90 days prior found an increase in the risk of re-infection during November 2021 compared to time periods earlier in the pandemic (estimated relative hazard ratio for reinfection versus primary infection of 2.39 (95%CI 1.88-3.11 from 1-27 November 2021 compared to wave 1 (June – September 2020), corresponding with the emergence of the Omicron variant. This information provides an initial assessment of the risk of re-infection however, further studies are needed to confirm this, including the ability of the Omicron variant to infect or re-infect those who have been vaccinated, as well as to determine the severity of these breakthrough or re-infections.<sup>7</sup>

## **Impact on diagnostics**

SARS-CoV-2 infection can be diagnosed using either molecular tests (NAAT, PCR) or antigen-detection assays. Interim guidance on diagnostic testing for SARS-CoV-2 can be found <a href="here">here</a> and on the use of antigen-detection tests can be found <a href="here">here</a>. PCR tests that include multiple gene targets are unlikely to be affected and should continue to be used

to detect SARS-CoV-2 infection, including the Omicron variant. This has been confirmed by statements issued by suppliers as well as the US FDA, based on sequence analysis.

The majority of Omicron variant sequences reported include a 69-70 deletion mutation in the Spike protein. There are some public sequences lacking this mutation and, at the present time, it is unclear if this reflects true sequence diversity or is a sequencing artifact. Presence of the 69-70 deletion causes dropout of some S-gene targets in PCR assays, such as the TaqPath COVID-19 Combo Kit and TaqPath COVID-19 CE-IVD RT-PCR Kit (Thermo Fisher Scientific). This S-gene target failure (SGTF) can be used as a marker suggestive of Omicron. However, confirmation should be performed by sequencing the sample, as this deletion is found in other VOCs (e.g., Alpha and subsets of Gamma and Delta) currently circulating at low levels globally, but possibly circulating at higher levels locally.

All four WHO emergency use listing (EUL) approved antigen-detection rapid diagnostic tests (Ag-RDTs), listed <a href="here">here</a>, target the Nucleocapsid protein of SARS-COV-2. The vast majority of Omicron sequences reported to date include the G204R and R203K mutations in the Nucleocapsid protein, which are present in many other variants currently in circulation. This has not been reported to affect the accuracy of Ag-RDTs to detect SARS-CoV-2. In addition, the majority of Omicron sequences contain a 3 amino acid deletion at positions 31-33 and the P13L mutation in the Nucleocapsid protein. The specific impact of these mutations on the performance of Ag RDTs is currently unclear.

Official statements from several Ag-RDT suppliers, including two with EUL-approved assays, indicate that based on sequence analysis, the performance of their tests is not impacted by the Omicron variant. Preliminary laboratory evidence is emerging that independently confirms that Ag-RDTs can accurately diagnose infection with the Omicron variant. To date, there have been no reported misdiagnoses (false negative results) for any WHO EUL approved diagnostic product in relation to Omicron.

#### Impact on vaccines

There is a need for more data to assess whether the mutations present on the Omicron variant may result in reduced protection from vaccine derived immunity and data on vaccine effectiveness, including the use of additional vaccination doses. WHO will continue to work with partners to monitor and evaluate these data once they become available. Vaccine effectiveness studies are vital to understand how vaccines protect against infection, symptomatic and severe disease, and death. WHO guidance on best practices to conduct these types of studies can be found on our website.

#### Impact on treatments

WHO continues to work with researchers to understand the effectiveness of treatments against the Omicron variant; however, Interleukin-6 Receptor Blockers and corticosteroids are expected to continue to be effective in the management of patients with severe disease.

#### **Conclusions**

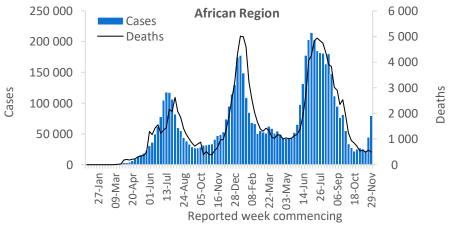
Whilst many questions about the Omicron variant remain unanswered, more information will continue to emerge in the coming weeks, with the TAG-VE and other groups reviewing and analysing data as it becomes available. It is important to continue to accelerate access to vaccines globally and for countries to continue to enhance surveillance, reporting initial cases or clusters to WHO and share genome sequences on publicly available databases such as GISAID. We recommend the public continue to prevent the spread of SARS-CoV-2 by improving ventilation of indoor spaces, wearing well-fitted masks, avoiding crowded spaces, practising hand hygiene and keeping an appropriate physical distance from others.

# WHO regional overviews Epidemiological week 29 November – 5 December 2021

## **African Region**

The case incidence in the African Region continues to increase with over 79 000 new cases reported during the week of 29 November to 5 December, a 79% increase. However, weekly deaths have continued to decrease, with just under 500 new deaths reported in the past week, a 13% decrease. Twenty-one of the 49 countries in the region (43%) reported an increase of >10% in new cases as compared to the previous week, with the highest numbers of new cases reported from South Africa (62 021 new cases; 104.6 new cases per 100 000; a 111% increase), Zimbabwe (4572 new cases; 30.8 new cases per 100 000; a 1361% increase) and Réunion (2140 new cases; 239.0 new cases per 100 000; a 14% increase). However, proportionally, very large increases in the incidence of cases were also seen in Eswatini (1900%), Mozambique (1207%) and Namibia (681%).

Six of the 49 countries in the Region reported an increase of over 10% in the number of new weekly deaths, with the highest numbers of new deaths reported from South Africa (174 new deaths; <1 new death per 100 000; a 21% decrease), Mauritius (126 new deaths; 9.9 new deaths per 100 000; an 31% increase), and Ethiopia (58 new deaths; <1 new death per 100 000; a 9% decrease).

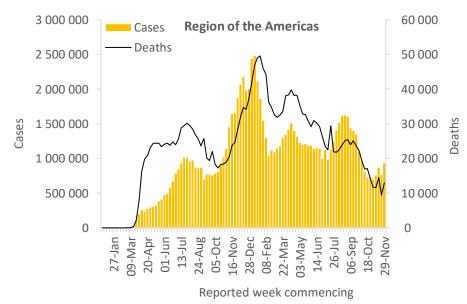


Updates from the African Region

## **Region of the Americas**

The Region of the Americas reported a 21% increase in case incidence in the past week, with over 935 000 new cases reported. This trend is largely driven by the increase in the incidence of cases reported in the United States of America (752 394 new cases; 227.3 new cases per 100 000; a 30% increase). Twenty-seven percent (15/56) of countries in the region reported increases of over 10%. In addition to the United States of America, countries reporting the highest numbers of cases included Brazil (61 779 new cases; 29.1 new cases per 100 000; similar to the previous week's figures) and Canada (20 188 new cases; 53.5 new cases per 100 000; similar to the previous week's figures).

The incidence of deaths also increased with just under 13 000 new deaths reported, a 38% increase compared to the previous week. The highest numbers of new deaths were reported from the United States of America (8527 new deaths; 2.6 new deaths per 100 000; a 56% increase), Brazil (1443 new deaths; <1 new death per 100 000; a 9% decrease) and Mexico (1002 new deaths; <1 new death per 100 000; a 55% increase).

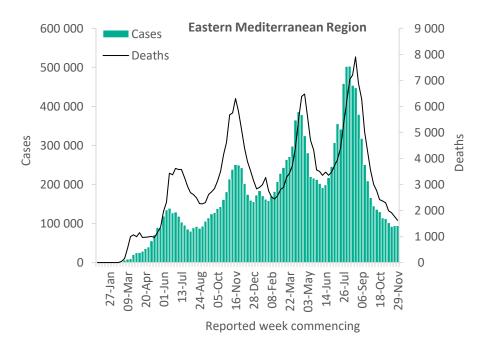


Updates from the Region of the Americas

#### **Eastern Mediterranean Region**

The weekly incidence of cases in the Eastern Mediterranean Region remained stable with over 94 000 new cases reported. The weekly incidence of deaths decreased by 8%, with over 1600 reported. However, nearly half (10/22) of countries in the region reported a >10% increase in weekly incidence of cases. Most cases continued to be reported from three countries: Jordan (32 108 new cases; 314.7 new cases per 100 000; a 15% increase), the Islamic Republic of Iran (26 255 new cases; 31.3 new cases per 100 000; an 18% decrease), and Lebanon (10 406 new cases; 152.5 new cases per 100 000; an 11% increase).

The highest numbers of new deaths continued to be reported from the Islamic Republic of Iran (575 new deaths; <1 new death per 100 000; an 18% decrease), Egypt (377 new deaths; <1 new death per 100 000; a 13% decrease), and Jordan (200 new deaths; 2.0 new deaths per 100 000; a 19% increase).

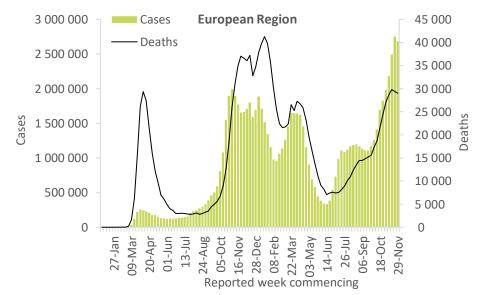


Updates from the Eastern Mediterranean Region

### **European Region**

Following an increase in the incidence of cases from mid-October, the weekly number of new cases in the European Region plateaued this week with just over 2.6 million new cases reported. The incidence in deaths also remained stable compared to the previous week, with over 29 000 new deaths reported. Fewer countries (11/61, 18%) reported an increase in new weekly cases of over 10% compared to the previous week. Germany and the United Kingdom continue to report the highest number of new cases with 396 429 new cases (476.7 new cases per 100 000; similar to the previous week's figures) and 310 696 new cases (457.7 new cases per 100 000; similar to the previous week's figures), respectively, with France reporting the third highest number of new cases (283 500 new cases; 435.9 new cases per 100 000; a 49% increase).

The highest numbers of new deaths continued to be reported from the Russian Federation (8523 new deaths; 5.8 new deaths per 100 000; similar to the previous week's figures); Ukraine (3163 new deaths; 7.2 new deaths per 100 000; an 18% decrease) and Poland (2636 new deaths; 6.9 new deaths per 100 000; a 19% increase).

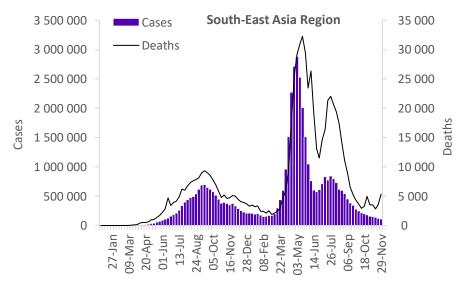


Updates from the European Region

#### **South-East Asia Region**

Since July 2021, the incidence of cases in the South-East Asia Region has continued to decline with over 109 000 new cases reported this week, a 10% decrease as compared to the previous week. Only one country reported an increase of over 10%, Timor-Leste (6 new cases; <1 case per 100 000; a 100% increase). The highest number of new cases continued to be reported from India (60 732 new cases; 4.4 new cases per 100 000; similar to the previous week's figures), Thailand (34 428 new cases; 49.3 new cases per 100 000; an 18% decrease) and Sri Lanka (5162 new cases; 24.1 new cases per 100 000; a 12% decrease).

The number of new weekly deaths however, increased by 49% as compared to the previous week, with over 5300 new deaths reported this week, the majority being reported from India (4772 new deaths; <1 new death per 100 000; a 65% increase). While Thailand and Sri Lanka reported the second and third highest number of deaths this past week, both countries reported a decline (237 new deaths; <1 new death per 100 000; a 26% decrease, and 156 new deaths; <1 new death per 100 000; a 12% decrease, respectively).



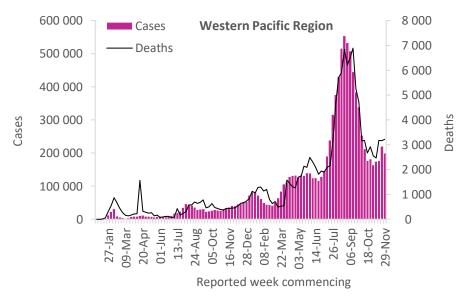
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Updates from the South-East Asia Region

### **Western Pacific Region**

Following an increase in the weekly case incidence in the Western Pacific Region from early November, in the past week a 10% decrease was seen with just under 200 000 new cases reported. However, five of the 27 countries in the region reported an increase in case incidence of >10%, including French Polynesia (3976%), China (147%), the Northern Mariana Islands (75%), the Republic of Korea (26%) and Fiji (20%). The highest number of new cases continued to be reported from Viet Nam (97 374 new cases; 100.0 new cases per 100 000; a 14% decrease), Malaysia (34 897 new cases; 107.8 new cases per 100 000; an 8% decrease) and the Republic of Korea (32 142 new cases; 62.7 new cases per 100 000; a 26% increase).

The region reported over 3200 new deaths this week, similar to the previous week's figures. Two of the three countries reporting the highest numbers of new deaths showed an increasing trend: Viet Nam (1369 new deaths; 1.4 new deaths per 100 000; a 36% increase) and the Republic of Korea (304 new deaths; <1 new death per 100 000; an 11% increase), while the Philippines reported a decrease (1025 new deaths; <1 new death per 100 000; a 21% decrease).



Updates from the Western Pacific Region

### **Summary of the COVID-19 Weekly Operational Update**

The <u>Weekly Operational Update</u> is a report provided by the COVID-19 Strategic Preparedness and Response Plan (SPRP) Monitoring and Evaluation team, which aims to update on the ongoing global progress against the <u>COVID-19 SPRP 2021</u> framework, and to highlight country-level actions and WHO support to countries. In this week's edition published on 7 December, highlights include the following:

- Kenya increases uptake and equity for COVID-19 vaccinations
- Supporting Omicron variant detection and COVID-19 response in southern Africa
- New oxygen production stations to be constructed in Yemen
- "Data for action": WHO/Europe pilots enhanced Emergency Response Information Management System (ERIMS) with Azerbaijan
- Home-based care for COVID-19 patients begins with community engagement in Lao People's Democratic Republic
- Progress on a subset of indicators from the SPRP 2021 Monitoring and Evaluation Framework
- Updates on WHO's financing to support countries in SPRP 2021 implementation and provision of critical supplies

## **Technical guidance and other resources**

- WHO technical guidance
- WHO COVID-19 Dashboard
- WHO Weekly Operational Updates on COVID-19
- WHO COVID-19 case definitions
- COVID-19 Supply Chain Inter-Agency Coordination Cell Weekly Situational Update
- Research and Development
- Open WHO courses on COVID-19 in official UN languages and in additional national languages
- WHO Academy COVID-19 mobile learning app
- <u>The Strategic Preparedness and Response Plan</u> (SPRP) outlining the support the international community can provide to all countries to prepare and respond to the virus
- Recommendations and advice for the public:
  - o Protect yourself
  - Questions and answers
  - o <u>Travel advice</u>
- EPI-WIN: tailored information for individuals, organizations, and communities

## **Annexes**

Annex 1. List of countries/territories/areas reporting variants of concern as of 7 December 2021

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Afghanistan	•	-	•	-	-
Albania	•	-	0	-	-
Algeria	•	-	•	-	-
Andorra	0	0	0	-	-
Angola	•	•	•	•	-
Anguilla	•	-	•	-	-
Antigua and Barbuda	•	•	•	•	-
Argentina	•	•	•	•	•*
Armenia	•	-	•	-	-
Aruba	•	•	•	•	-
Australia	•	•	•	•	•*
Austria	•	•	•	•	•*
Azerbaijan	•	-	0	-	-
Bahamas	•	-	•	•	-
Bahrain	•	•	•	•	-
Bangladesh	•	•	•	0	-
Barbados	•	-	•	•	-
Belarus	•	-	0	-	-
Belgium	•	•	•	•	•*
Belize	•	-	•	•	-
Benin	•	•	•	•	-
Bermuda	•	•	•	-	-
Bhutan	•	•	•	-	-
Bolivia (Plurinational State of)	•	-	•	•	-
Bonaire	•	-	•	•	-
Bosnia and Herzegovina	•	•	0	•	-
Botswana	0	•	•	-	•*
Brazil	•	•	•	•	•*

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
British Virgin Islands	•	-	•	•	-
Brunei Darussalam	•	•	•	-	-
Bulgaria	•	•	•	-	-
Burkina Faso	•	-	•	-	-
Burundi	•	•	•	-	-
Cabo Verde	•	-	•	-	-
Cambodia	•	•	•	-	-
Cameroon	•	•	•	•*	-
Canada	•	•	•	•	•*
Cayman Islands	•	•	•	•	-
Central African Republic	•	•	•	-	-
Chad	•	-	-	-	-
Chile	•	•	•	•	•*
China	•	•	•	•	•*
Colombia	•	-	•	•	-
Comoros	-	•	•	-	-
Congo	•	•	•	•	-
Costa Rica	•	•	•	•	-
Croatia	•	•	0	•	•*
Cuba	•	•	•	-	-
Curaçao	•	•	•	•	-
Cyprus	•	•	0	-	-
Czechia	•	•	•	•	•*
Côte d'Ivoire	•	•	0	-	-
Democratic Republic of the Congo	•	•	•	-	-
Denmark	•	•	•	•	•*
Djibouti	•	•	•*	-	-

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Dominica	•	-	•	-	-
Dominican Republic	•	-	•	•	-
Ecuador	•	-	•	•	-
Egypt	•	-	•	-	-
El Salvador	•	-	•	•	-
Equatorial Guinea	•	•	•	-	-
Estonia	•	•	0	0	0*
Eswatini	0	•	•	-	-
Ethiopia	•	•	•	-	-
Falkland Islands (Malvinas)	•	•	-	-	-
Faroe Islands	•	-	-	•	-
Fiji	0	-	•	-	•*
Finland	•	•	•	•	•*
France	•	•	•	•	•*
French Guiana	•	•	•	•	-
French Polynesia	•	•	•	•	-
Gabon	•	•	•	-	-
Gambia	•	-	•	-	-
Georgia	•	0	•	-	-
Germany	•	•	•	•	•*
Ghana	•	•	•	•	•*
Gibraltar	•	-	0	-	-
Greece	•	•	•	•	•*
Greenland	-	-	•	-	-
Grenada	•	-	•	•	-
Guadeloupe	•	•	•	•	-
Guam	•	•	•	•	-
Guatemala	•	•	•	•	-

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Guinea	•	•	•	-	-
Guinea-Bissau	•	•	•	-	-
Guyana	-	-	•	•	-
Haiti	•	-	•	•	-
Honduras	•	-	•	•	-
Hungary	•	0	0	•	-
Iceland	•	•	•	•	•*
India	•	•	•	•	•*
Indonesia	•	•	•	-	-
Iran (Islamic Republic of)	•	•	•	-	-
Iraq	•	•	•	•	-
Ireland	•	•	•	•	•*
Israel	•	•	•	•	•*
Italy	•	•	•	•	•*
Jamaica	•	-	•	-	-
Japan	•	•	•	•	•*
Jordan	•	•	•	•	-
Kazakhstan	•	0	•	-	-
Kenya	•	•	•	-	-
Kosovo[1]	•	0	0	-	-
Kuwait	•	•	•	-	-
Kyrgyzstan	•	•	•	-	-
Lao People's Democratic Republic	•	-	•	-	-
Latvia	•	•	0	•	•*
Lebanon	•	-	•	-	-
Lesotho	-	•	0	-	-
Liberia	•	•	•	-	-
Libya	•	•	-	-	-
Liechtenstein	•	-	0	0	-

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Lithuania	•	•	0	•	-
Luxembourg	•	•	•	•	•*
Madagascar	•	•	-	-	-
Malawi	•	•	•	-	-
Malaysia	•	•	•	-	•*
Maldives	•	-	•	-	•*
Mali	-	-	•	-	-
Malta	•	0	0	•	-
Martinique	•	•	•	•	-
Mauritania	•	•	•	-	-
Mauritius	•	•	•	-	-
Mayotte	•	•	0	-	-
Mexico	•	•	•	•	•*
Monaco	•	•	•	-	-
1ongolia	•	-	•	-	-
1ontenegro	•	-	0	0	0*
Montserrat	•	-	•	•	-
Morocco	•	•	•	-	-
Лоzambique	•	•	•	-	-
Лyanmar	•	-	•	-	-
Namibia	•	•	•	•	•*
Nepal	•	-	•	-	•*
Netherlands	•	•	•	•	•*
New Caledonia	•	-	•	-	-
New Zealand	•	•	•	•	-
Nicaragua	•	•	•	•	-
Niger	•	-	•	-	-
Nigeria	•	•	•	-	•*
North Macedonia	•	•	0	-	-

Omicron	Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
*	Northern Mariana Islands (Commonwealth of the)	0	-	•	-	-
	Norway	•	•	•	•	•*
	Occupied Palestinian Territory	•	•	•	-	-
*	Oman	•	•	•	-	-
*	Pakistan	•	•	•	•	-
	Panama	•	•	•	•	-
	Papua New Guinea	-	-	•	-	-
	Paraguay	•	-	•	•	-
	Peru	•	-	•	•	-
	Philippines	•	•	•	•	-
	Poland	•	0	•	•	-
*	Portugal	•	•	•	•	•*
	Puerto Rico	•	•	•	•	-
	Qatar	•	•	•	-	-
*	Republic of Korea	•	•	•	•	•*
	Republic of Moldova	•	-	•	-	-
	Romania	•	•	•	•	•*
	Russian Federation	•	•	•	0	0*
	Rwanda	•	•	•	-	-
*	Réunion	•	•	0	•	•*
*	Saba	-	-	•	-	-
*	Saint Barthélemy	•	-	•	-	-
	Saint Kitts and Nevis	-	-	•	-	-
	Saint Lucia	•	-	•	-	-
	Saint Martin	•	•	•	-	-
	Saint Pierre and Miquelon	-	-	•	-	-
*	Saint Vincent and the Grenadines	-	-	•	•	-
	Sao Tome and Principe	•	-	0	-	-
	Saudi Arabia	•	•	•	-	•*

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Senegal	•	•	•	-	•*
Serbia	•	-	•	-	-
Seychelles	•	•	•	-	-
Sierra Leone	-	•	•	-	-
Singapore	•	•	•	•	•*
Sint Maarten	•	•	•	•	-
Slovakia	•	•	•	-	-
Slovenia	•	•	•	•	-
Somalia	•	•	•	-	-
South Africa	•	•	•	0	•*
South Sudan	•	•	•	-	-
Spain	•	•	•	•	•*
Sri Lanka	•	•	•	-	•*
Sudan	•	•	-	•	-

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Suriname	•	•	•	•	-
Sweden	•	•	•	•	•*
Switzerland	•	•	•	•	•*
Thailand	•	•	•	•	•*
Timor-Leste	•	-	•	-	-
Togo	•	•	•	•	-
Trinidad and Tobago	•	-	•	•	-
Tunisia	•	•	•	-	•*
Turkey	•	•	•	•	-
Turks and Caicos Islands	•	-	•	•	-
Uganda	•	•	•	-	•*
Ukraine	•	0	0	-	-
United Arab Emirates	•	•	•	•	•*
United Kingdom	•	•	•	•	•*

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
United Republic of Tanzania	-	•	-	-	-
United States Virgin Islands	•	•	•	•	-
United States of America	•	•	•	•	•*
Uruguay	•	•	•	•	-
Uzbekistan	•	•	0	-	-
Vanuatu	-	-	•	-	-
Venezuela (Bolivarian Republic of)	•	-	•	•	-
Viet Nam	•	•	•	-	-
Wallis and Futuna	•	-	-	-	-
Yemen	•	•	-	-	-
Zambia	•	•	•	-	•*
Zimbabwe	•	•	•	-	•*

<sup>\*</sup>Newly reported in this update. "•" indicates that information for this variant was received by WHO from official sources. "o" indicates that information for this variant was received by WHO from unofficial sources and will be reviewed as more information become available. \*\*Includes countries/territories/areas reporting the detection of VOCs among travellers (e.g., imported cases detected at points of entry), or local cases (detected in the community). Excludes countries, territories, and areas that have never reported the detection of a variant of concern. See also Annex 2: Data, table, and figure notes

#### Annex 2. Data, table, and figure notes

Data presented are based on official laboratory-confirmed COVID-19 case and deaths reported to WHO by country/territories/areas, largely based upon WHO <u>case definitions</u> and <u>surveillance guidance</u>. While steps are taken to ensure accuracy and reliability, all data are subject to continuous verification and change, and caution must be taken when interpreting these data as several factors influence the counts presented, with variable underestimation of true case and death incidences, and variable delays to reflecting these data at the global level. Case detection, inclusion criteria, testing strategies, reporting practices, and data cut-off and lag times differ between countries/territories/areas. A small number of countries/territories/areas report combined probable and laboratory-confirmed cases. Differences are to be expected between information products published by WHO, national public health authorities, and other sources.

Due to public health authorities conducting data reconciliation exercises that remove large numbers of cases or deaths from their total counts, negative numbers may be displayed in the new cases/deaths columns as appropriate. When additional details become available that allow the subtractions to be suitably apportioned to previous days, graphics will be updated accordingly. A record of historic data adjustment made is available upon request by emailing <a href="mailto:epi-data-support@who.int">epi-data-support@who.int</a>. Please specify the countries of interest, time period, and purpose of the request/intended usage. Prior situation reports will not be edited; see <a href="mailto:covid19.who.int">covid19.who.int</a> for the most up-to-date data. COVID-19 confirmed cases and deaths reported in the last seven days by countries, territories, and areas, and WHO Region (reported in previous issues) are now available at: <a href="https://covid19.who.int/table">https://covid19.who.int/table</a>.

'Countries' may refer to countries, territories, areas or other jurisdictions of similar status. The designations employed, and the presentation of these materials do not imply the expression of any opinion whatsoever on the part of WHO concerning the legal status of any country, territory, or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted and dashed lines on maps represent approximate border lines for which there may not yet be full agreement. Countries, territories, and areas are arranged under the administering WHO region. The mention of specific companies or of certain manufacturers' products does not imply that they are endorsed or recommended by WHO in preference to others of a similar nature that are not mentioned. Errors and omissions except, the names of proprietary products are distinguished by initial capital letters.

[1] All references to Kosovo should be understood to be in the context of the United Nations Security Council resolution 1244 (1999). In the map, the number of cases of Serbia and Kosovo (UNSCR 1244, 1999) have been aggregated for visualization purposes.

#### References

- 1. COVID-19 Epidemiology Update 2nd December 2021 SA Corona Virus Online Portal. Accessed December 7, 2021. https://sacoronavirus.co.za/2021/12/03/covid-19-epidemiology-update-2nd-december-2021/
- 2. COVID-19-Weekly-Epidemiology-Brief-week-46-2021.pdf. Accessed December 6, 2021. https://www.nicd.ac.za/wp-content/uploads/2021/11/COVID-19-Weekly-Epidemiology-Brief-week-46-2021.pdf
- 3. Epidemiological update: Omicron variant of concern (VOC) data as of 6 December 2021 (12.00). European Centre for Disease Prevention and Control. Published December 6, 2021. Accessed December 7, 2021. https://www.ecdc.europa.eu/en/news-events/epidemiological-update-omicron-variant-concern-voc-data-6-december-2021
- 4. Threat Assessment Brief: Implications of the further emergence and spread of the SARS CoV 2 B.1.1.529 variant of concern (Omicron) for the EU/EEA first update. European Centre for Disease Prevention and Control. Published December 2, 2021. Accessed December 6, 2021. https://www.ecdc.europa.eu/en/publications-data/covid-19-threat-assessment-spread-omicron-first-update
- 5. Datcov19\_National\_Export-20211203.pdf. Accessed December 6, 2021. https://www.nicd.ac.za/wp-content/uploads/2021/12/Datcov19\_National\_Export-20211203.pdf
- 6. SACMC-Fourth-wave-report-17112021-final.pdf. Accessed December 7, 2021. https://www.nicd.ac.za/wp-content/uploads/2021/11/SACMC-Fourth-wave-report-17112021-final.pdf
- 7. Pulliam JRC, van Schalkwyk C, Govender N, et al. *Increased Risk of SARS-CoV-2 Reinfection Associated with Emergence of the Omicron Variant in South Africa*. Epidemiology; 2021. doi:10.1101/2021.11.11.21266068



## **COVID-19 Weekly Epidemiological Update**

#### Edition 70, published 14 December 2021

#### In this edition:

- Global overview
- Special focus: Update on SARS-CoV-2 Variants of Interest and Variants of Concern
- WHO regional overviews
- Summary of the Weekly Operational Update

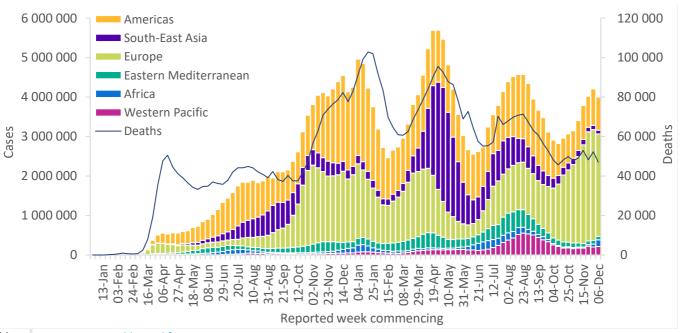
#### **Global overview**

#### Data as of 12 December 2021

Globally, the weekly incidence of both cases and deaths declined during the past week (6-12 December 2021), with decreases of 5% and 10% respectively, as compared to the previous week. Nonetheless, this still corresponded to over 4 million new confirmed cases and just under 47 000 new deaths. As of 12 December, nearly 269 million confirmed cases and nearly 5.3 million deaths have been reported globally.

The African Region reported the largest increase in new cases last week (111%) followed by and the Western Pacific Region which reported an increase of 7%. The Region of the Americas and South-East Asia Region both reported decreases of 10% and the European Region reported a 7% decrease. The number of new weekly cases reported by the Eastern Mediterranean Region was similar to the numbers reported in the previous week. New weekly deaths decreased by 50% in the South-East Asia Region (due to an artificial increase in deaths from batch reporting in the previous week) and 14% in the Region of the Americas, while the number of weekly deaths in all other regions remained similar to those reported in the previous week.

Figure 1. COVID-19 cases reported weekly by WHO Region, and global deaths, as of 12 December 2021\*\*



<sup>\*\*</sup>See Annex 3: Data, table, and figure notes

The regions reporting the highest weekly case incidence per 100 000 population continue to be the European Region (277.9 new cases per 100 000 population) and the Region of the Americas (81.9 new cases per 100 000 population). Both regions also reported the highest weekly incidence in deaths of 3.0 and 1.0 per 100 000 population, respectively while <1 new death per 100 000 was reported in all other regions.

The highest numbers of new cases were reported from the United States of America (674 019 new cases; 9% decrease), Germany (351 738 new cases; 11% decrease), the United Kingdom (350 340 new cases; 13% increase), France (335 972 new cases; 19% increase), and the Russian Federation (215 283 new cases; 7% decrease).

Table 1. Newly reported and cumulative COVID-19 confirmed cases and deaths, by WHO Region, as of 12 December 2021\*\*

WHO Region	New cases in last 7 days (%)	Change in new cases in last 7 days *	Cumulative cases (%)	New deaths in last 7 days (%)	Change in new deaths in last 7 days *	Cumulative deaths (%)
Europe	2 593 221 (65%)	-7%	91 631 852 (34%)	28 362 (60%)	-3%	1 598 688 (30%)
Americas	837 345 (21%)	-10%	98 521 311 (37%)	10 562 (22%)	-14%	2 371 246 (45%)
Western Pacific	213 915 (5%)	7%	10 584 344 (4%)	3 335 (7%)	4%	147 539 (3%)
Africa	167 682 (4%)	111%	6 522 517 (2%)	491 (1%)	-1%	153 766 (3%)
South-East Asia	98 021 (2%)	-10%	44 737 006 (17%)	2 643 (6%)	-50%	714 303 (13%)
Eastern Mediterranean	90 633 (2%)	-4%	16 936 781 (6%)	1 568 (3%)	-3%	312 295 (6%)
Global	4 000 817 (100%)	-5%	268 934 575 (100%)	46 961 (100%)	-10%	5 297 850 (100%)

<sup>\*</sup>Percent change in the number of newly confirmed cases/deaths in the past seven days, compared to seven days prior

For the latest data and other updates on COVID-19, please see:

- WHO COVID-19 Dashboard
- WHO COVID-19 Weekly Operational Update and previous editions of the Weekly Epidemiological Update

<sup>\*\*</sup>See Annex 3: Data, table, and figure notes

Cases reported in the last 7 days (per 100,000 population) 0.01 - 10.00 10.01 - 50.00 50.01 - 100.00 100.01 - 300.00 > 300.00 No cases reported in the last 7 days No reported cases The designations employed and the presentation of the material in this publication do not imply the expression Data Source: World Health Organization, of any opinion whatsoever on the part of WHO concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted and dashed lines on maps represent approximate border lines for which there may not yet be full agreement. Data for Bonaire, Sint United Nations Population Division (population prospect 2020), EuroStat

Map Production: WHO Health Emergencies Programme Eustatius and Saba have been disaggregated and displayed at the subnational level.

Figure 2. COVID-19 cases per 100 000 population reported by countries, territories and areas, 6 December - 12 December 2021\*\*

<sup>\*\*</sup>See Annex 3: Data, table, and figure notes

Deaths reported in the last 7 days (per 100,000 population) 0.01 - 0.50 0.51 - 1.50 1.51 - 3.00 3.01 - 6.00 > 6.00 No deaths reported in the last 7 days No reported cases Data Source: World Health Organization, maps represent approximate border lines for which there may not yet be full agreement. Data for Bonaire, Sint Eustatius and Saba have been disaggregated and displayed at the subnational level. © World Health Organization 2021, All rights reserved. Map Production: WHO Health Emergencies Programme

Figure 3. COVID-19 deaths per 100 000 population reported by countries, territories and areas, 6 December - 12 December 2021\*\*

<sup>\*\*</sup>See Annex 3: Data, table, and figure notes

#### Special Focus: Update on SARS-CoV-2 Variants of Interest and Variants of Concern

WHO, in collaboration with national authorities, institutions and researchers, routinely assesses if variants of SARS-CoV-2 alter transmission or disease characteristics, or impact effectiveness of vaccines, therapeutics, diagnostics or public health and social measures (PHSM) applied by national authorities to control disease spread. Potential Variants of Concern (VOCs), Variants of Interest (VOIs) or Variants Under Monitoring (VUMs) are regularly assessed based on the risk posed to global public health. As evidence becomes available, classifications of variants will be revised to reflect the continuous evolution of circulating variants and their changing epidemiology. Criteria for variant classification, and the current lists of VOCs, VOIs and VUMs, are available on the <a href="https://www.website">WHO Tracking SARS-CoV-2 variants</a> website. National authorities may choose to designate other variants of local interest/concern and are encouraged to investigate and report on the impacts of these variants.

#### Geographic spread and prevalence of VOCs

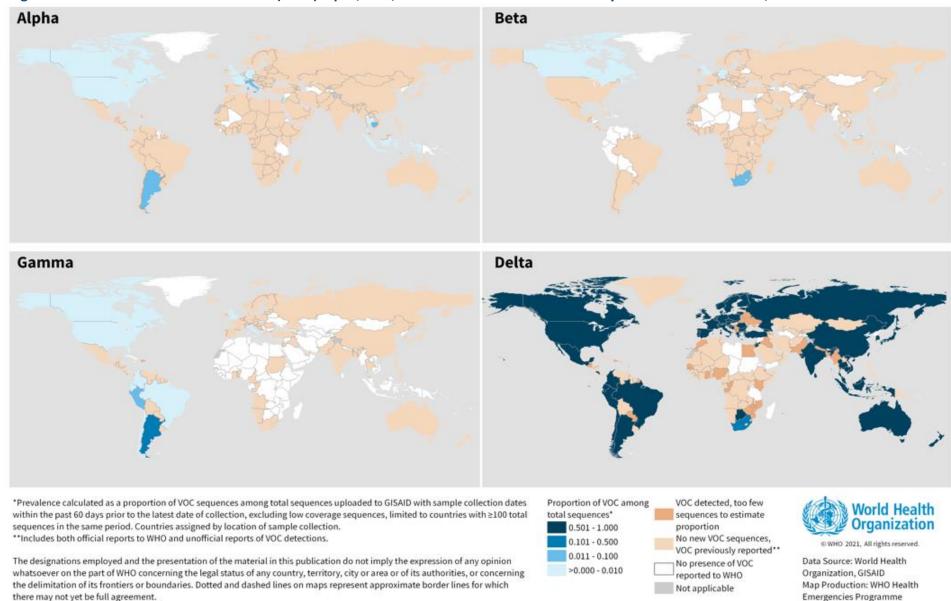
The current global epidemiology of SARS-CoV-2 is characterized by a predominance of the Delta variant, declining trend in the proportion of Alpha, Beta and Gamma, and the emergence of Omicron variant; however, regional and country-level variations continue to be observed (Figure 4 and 5; Annex 2). While most of the Omicron cases identified in November 2021 were travel-related, community transmission with associated clusters has now been reported in several countries. Of 879 779 sequences uploaded to GISAID with specimens collected in the last 60 days, 872 876 (99.2%) were Delta, 3 755 (0.4%) were Omicron, 206 (<0.1%) Alpha, 179 (<0.1%) Gamma, 16 (<0.1%) Beta, and <0.1% comprised other circulating variants (including VOIs Mu and Lambda). This week, for the first time since Delta was classified as a VOC in April 2021, the percentage of Delta sequences has declined in respect to other VOCs. However, this observation needs to be interpreted with caution as countries may perform targeted sequencing for Omicron and therefore upload fewer sequences on all other variants, including Delta.

To note, global VOCs distribution should be interpreted with due consideration of surveillance limitations, including differences in sequencing capacities and sampling strategies between countries, as well as delays in reporting.

#### **Additional resources**

- Tracking SARS-CoV-2 Variants
- COVID-19 new variants: Knowledge gaps and research
- Genomic sequencing of SARS-CoV-2: a guide to implementation for maximum impact on public health
- Considerations for implementing and adjusting PHSM in the context of COVID-19

Figure 4: Prevalence of Variants of Concern (VOCs) Alpha, Beta, Gamma and Delta in the last 60 days and historic detections, data as of 14 December 2021



See also Annex 2 for reported VOC detections by country/territory/area

Presence of Omicron variant reported Verified (70) //// Under verification (6) Presence not reported Not applicable

Figure 5. Presence of Variant of Concern (VOC) Omicron, data as of 14 December 2021 (4 pm CET)

The designations employed and the presentation of the material in this publication do not imply the expression of any opinion whatsoever on the part of WHO concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted and dashed lines on maps represent approximate border lines for which there may not yet be full agreement.

Data Source: World Health Organization, GISAID Map Production: WHO Health Emergencies Programme



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Presence of the Omicron variant is based on information reported to WHO. It includes countries/territories/areas reporting the detection of VOCs among travellers (e.g., imported cases detected at points of entry), or local cases (detected in the community). See also Annex 2 for reported VOC detections by country/territory/area.

#### **Update on Omicron VOC**

Since the last <u>update published on 7 December</u>, additional countries across all six WHO Regions have reported confirmed cases of the Omicron variant. As of 14 December 2021 (2 pm CET), the Omicron variant has been confirmed in 76 countries.

Based on current limited evidence Omicron appears to have a growth advantage over Delta. It is spreading faster than the Delta variant in South Africa where Delta circulation was low, but also appears to be spreading more quickly than the Delta variant in countries where the incidence of Delta is high, such as in the United Kingdom.

The data on the clinical severity of Omicron remains limited. More information on case severity associated with Omicron is expected in the coming weeks due to the time lag between an increase in the incidence of cases and an increase in the incidence of severe cases, and deaths.

Preliminary evidence suggests that there may be a reduction in vaccine efficacy and effectiveness against infection and transmission associated with Omicron, as well as an increased risk of reinfection. More data are needed to better under the extent to which Omicron may evade vaccine and/or infection derived immunity and the extent to which current vaccines continue to protect against severe disease and death associated with Omicron.

Diagnostic accuracy of routinely used PCR and antigen-based rapid diagnostic test (Ag-RDT) assays does not appear to be impacted by Omicron, and therapeutic interventions for the management of patients with severe or critical COVID-19 associated with the Omicron variant are expected to remain effective

As a result of this, the overall risk related to the new variant of concern Omicron remains very high. More information on Omicron variant can be found in the updated <u>Technical Brief and Priority Actions for Member States</u> that was published on 10 December 2021 by WHO.

#### Phenotypic characteristics

Available evidence on the phenotypic impacts of VOCs is summarized in Table 2, as well as in <u>previous editions</u> of the COVID-19 Weekly Epidemiological Update. Since the <u>last detailed update on 30 November</u>, there are several new publications on the phenotypic characteristics of VOCs, including recent literature on Omicron. Reported studies might have not been formally peer-reviewed and findings must be interpreted in the light of this limitation.

A cohort analysis reported by UK Health Security Agency<sup>1</sup>, estimated the odds of household transmission for Omicron variant index cases as compared with Delta variant index cases. The analysis included 72,761 index cases of Delta and 121 of Omicron in residential households with a specimen collection date between 15 and 28 November 2021. Household transmission was defined as an index (first) case followed by one or more laboratory confirmed SARS-CoV-2 cases at the same private dwelling within a 14-day period (minimum 7 days follow-up). Multivariable logistic regression model found the adjusted odds ratio for household transmission from an Omicron index case was 3.2 (95%Cl 2.0-5.0, p <0.001) compared to Delta index cases.

Reports describing Omicron cases among partially and fully vaccinated individuals have been recently released:

• The US CDC<sup>2</sup> reported the characteristics of the 43 investigated cases attributed to the Omicron variant. Twenty-five of them (58%) were in persons aged 18–39 years, and 14 (33%) persons reported international travel during

the 14 days preceding symptom onset or receipt of a positive test result. Thirty-four cases (79%) occurred in persons who completed the primary series of an FDA-authorized or approved COVID-19 vaccine ≥14 days before symptom onset or receipt of a positive SARS-CoV-2 test result, including 14 who had received an additional or booster dose; five of the 14 persons had received the additional dose <14 days before symptom onset. Six (14%) persons also had a documented previous SARS-CoV-2 infection. The most commonly reported symptoms were cough, fatigue, and congestion or runny nose. One vaccinated patient was hospitalized for 2 days, and no deaths among the 43 cases reported by US CDC have been reported to date. Case investigations have identified exposures associated with international and domestic travel, large public events, and household transmission.

- Preliminary findings published by the Norwegian Institute of Public Health (NIPH)<sup>3</sup>, described the result of the investigation of an Omicron outbreak that occurred during a Christmas party. Of 111 participants at the Christmas party, 73% (80 people) were subsequently diagnosed with SARS-CoV-2. Of these, 17 were confirmed with the Omicron variant by sequencing as of 8 December 2021. Analysis of additional samples is ongoing. Over 70% of cases reported cough, lethargy, headache, sore throat and over half of them reported fever. No hospital admissions have been reported. According to NIPH, most of the cases (the report does not specify the number) were aged between 30 and 50 years and were vaccinated with two doses of an mRNA vaccine between May and November 2021.
- A report from South Africa<sup>4</sup> described seven cases of breakthrough infection with the Omicron variant among visitors who received three doses of SARS-CoV-2 vaccines. On arrival in South Africa during the first half of November, all cases provided a negative SARS-CoV-2 PCR test and a record of complete vaccination including the third dose. Six cases were fully vaccinated with Pfizer BioNTech-Comirnaty and five of them also received a third dose of the same vaccine in October or early November 2021. One person received a third dose of ModernamRNA-1273 at the beginning of October. The seventh subject received an initial dose of AstraZeneca-Vaxzevria, followed by two doses of Pfizer BioNTech-Comirnaty for completion of primary immunization and as a third dose. None of them had a previous history of a SARS-CoV-2 infection. The cases developed onset of mild respiratory symptoms from 30 November to 2 December 2021 while in Cape Town and samples were collected between 2 to 4 days after onset of symptoms. Genome sequencing confirmed 5 of the cases to be due to Omicron variant; in two cases sequencing failed but they are inferred to be Omicron, too, based on their very close epidemiological links to the others.

Preliminary evidence shows that sera obtained from vaccinated and previously infected individuals has lower neutralization activity on Omicron VOC than with any other circulating VOCs of SARS-CoV-2 and the ancestral strain:

- A study (not yet peer reviewed) conducted in South Africa<sup>5</sup>, investigated whether the Omicron variant escapes antibody neutralization elicited by the Pfizer BioNTech-Comirnaty vaccine. Plasma samples from 12 participants fully vaccinated were tested. Six participants had a record of previous infection from the first SARS-CoV-2 wave in South Africa that was due to the ancestral SARS-Cov-2 strain. The other six participants had no previous record of SARS-CoV-2 infection nor detectable nucleocapsid antibodies indicative of previous infection. Overall, the geometric mean titer 50% focus reduction neutralization test (GMT FRNT50) was 1321 for the ancestral strain, indicating a very strong neutralization. However, the GMT FRNT50 for the Omicron variant was 32, a 41-fold reduction compared to the ancestral strain. Those participants who had a previous infection in addition to vaccination, had a higher GMT FRNT50, both versus Omicron variant and the ancestral strain.
- Another preprint study<sup>6</sup> analysed titers of neutralizing antibodies of sera from convalescent or vaccinated individuals against Omicron and compared them with titers against other VOCs (Alpha, Beta and Delta). Sera were selected from patients after an infection with Alpha (n=10), Beta (n=8) and Delta variants (n=7); from individuals

fully vaccinated with no previous SARS-CoV-2 infection (n=60) and fully vaccinated with previous SARS-CoV-2 infection (n=10). Sera from fully vaccinated individuals without previous SARS-CoV-2 infection neutralized the Omicron variant to a much lesser extent (not specified by authors) than any other VOCs analysed. Sera from fully vaccinated persons with previous SARS-CoV-2 infection, were able to neutralize Omicron variant, although to a lesser degree (not specified by authors) than the other VOCs.

• A study from Karolinska Institute<sup>7</sup> (not yet published) assessed neutralization activity against the Omicron variant. Two cohorts with laboratory confirmed SARS-CoV-2 in May 2020 and serum samples collected in November 2021, were included. Cohort 1 comprised serum samples with detectable neutralization against the ancestral SARS-CoV-2 strain from 17 anonymized blood donors. Cohort 2 comprised 17 serum samples from hospital workers with previous SARS-Cov-2 infection and that were fully vaccinated (the report does not specify with which vaccine). Almost all serum samples evaluated, retained some neutralization activity against the Omicron variant. Fold-reduction in the neutralization of Omicron relative to the ancestral strain, ranged from 1 to 23, with quartiles of 2.5, 5.5, and 11, measured by lentiviral pseudotype neutralization assay. As compared with Delta, Omicron showed a further reduction in neutralization activity, but that was not significant.

If specific antibodies neutralizing activity seems to be reduced versus the Omicron VOC, preliminary studies suggest that CD8+ T-cell responses may still maintain capability to recognize the Omicron VOC. A study from United States<sup>8</sup> not yet published, examined SARS-CoV-2 CD8+ T-cell responses from 30 recovered COVID-19 convalescent patients, evaluating if the previously identified viral epitopes targeted by CD8+ T-cells in these individuals (n=52 distinct epitopes) are mutated in the newly described Omicron variant. Within this population, only one low-prevalence (found in 2/30 (7%) of participants) epitope restricted to two HLA alleles from the Spike protein was found to be mutated in Omicron and contains only a single amino acid change. These data suggest that individuals with existing anti-SARS-CoV-2 CD8+ T-cell responses should recognize the Omicron variant, and that SARS-CoV-2 has not evolved extensive T-cell escape mutations.

A retrospective cohort study<sup>9</sup> was conducted in the United States to evaluate the duration of immunity to the Delta variant following infection. Of the samples tested prior from 9 March-31 December 2020, 15.5% (50 327/ 325 157) individuals were positive for SARS-CoV-2. During the Delta dominant period, protection of prior SARS-CoV-2 infection (defined as 1 minus the ratio of the infection rate for those initially positive to those initially negative) against reinfection was 85.4% (95%CI 80.0-89.3%) however, was lower for asymptomatic compared to symptomatic infection (66.6% (95%CI 40.6-81.2%). From 30 August 2020 to 9 September 2021, prior infection provided an overall 85.7% (95%CI 82.2-88.5%) protection against reinfection and again, protection against asymptomatic infection was lower than for symptomatic infection (52.2% (95%CI 35.3-64.7%) and 92.0% (95%CI 89.1-94.2%, respectively). Additionally, long-term protection among those aged 65 years and over was lower than for those aged under 65 years (76.3% compared to 88.9%, p<0.001).

A study conducted in the Republic of Korea<sup>10</sup> evaluated the transmissibility of the Delta variant among household contacts. A total of 405 cases with a median age of 19 years (1-71 years) who were infected with the Delta variant between 22 June to 31 July from Daejeon metropolitan city were included the study, of whom 325 (80.2%) were symptomatic. From these cases, six local clusters (two or more confirmed infections) were identified, all of which were associated with indoor facilities with the largest related to a sports academy (n=249 cases) and the second largest, a karaoke centre (n=47 cases). It was also estimated that 80% of all local transmission was caused by 15% (95%Cl 13-18%) of cases and from 258 infector-infectee transmission pairs, an estimated mean serial interval (the time between which the infector and infectee show symptoms) of 3.26 days (95% credible interval of 2.92-3.60 days) was calculated. A secondary attack rate of 63.4% (52/82 cases) was calculated based on data from 23 household

contacts in 32 homes. The study demonstrates the high transmissibility of the Delta variant in indoor settings and households. Contact tracing and isolation and the use of personal preventive measures during indoor activity remain imperative particularly given the potential for pre-symptomatic transmission.

Table 2: Summary of phenotypic impacts\* of variants of concern

WHO label	Alpha	Beta	Gamma	Delta	Omicron
Transmissibility	Increased transmissibility <sup>11</sup>	Increased transmissibility <sup>12,13</sup>	Increased transmissibility <sup>13,14</sup>	Increased transmissibility 13,15,16	No direct evidence for increased transmissibility.
Disease severity	Possible increased risk of hospitalization <sup>17,18</sup> , possible increased risk of severe disease and death <sup>19,20</sup>	Possible increased risk of hospitalization <sup>18</sup> , possible increased in-hospital mortality <sup>21</sup>	Possible increased risk of hospitalization <sup>18</sup> , possible increased risk of severe disease <sup>22</sup>	Possible increased risk of hospitalization <sup>23,24</sup>	Not yet known. Clinical outcome data are under review.
Risk of reinfection	Neutralizing activity retained <sup>25</sup> , risk of reinfection remains similar <sup>26</sup>	Reduction in neutralizing activity reported; T cell response elicited by D614G virus remains effective <sup>27</sup>	Moderate reduction in neutralizing activity reported <sup>28</sup>	Reduction in neutralizing activity reported <sup>29–31</sup>	Preliminary evidence suggests a possible increased risk of reinfection <sup>32</sup>
Impacts on diagnostics	Limited impact – S gene target failure (SGTF), no impact on overall result from multiple target RT-PCR; No impact on Ag RDTs observed <sup>33</sup>	PCR or Ag RDTs	None reported to date	No impact on RT- PCR or Ag RDTs observed <sup>34</sup>	PCR continues to detect Omicron. Impact on Ag-RDTs is under investigation.

<sup>\*</sup>Generalized findings as compared to previously/co-circulating variants. Based on emerging evidence, including non-peer-reviewed preprint articles and reports, all subject to ongoing investigation and revision.

Table 3 summarizes the impact of variants on product specific vaccine efficacy/effectiveness (VE) and quantifies the reduction in VE in the setting of variants compared to non-VOC settings. Since the <u>30 November update</u>, a total of 10 notable new studies have provided evidence of COVID-19 vaccine performance against the variants of concern.

As of December 12, seven recent studies have provided evidence of vaccine performance against the Omicron variant: 1 VE study and 6 neutralization studies. Note all studies are preliminary and more data are needed to confirm findings.

The first study of real-world VE against the Omicron variant (not yet peer-reviewed) provides preliminary evidence of reduced effectiveness of AstraZeneca-Vaxzevria and Pfizer BioNTech-Comirnaty against symptomatic disease due to Omicron.<sup>35</sup> No data on VE against severe disease was reported. <sup>35</sup>. This test-negative case-control study conducted in the United Kingdom found evidence that two doses of AstraZeneca-Vaxzevria was not effective at preventing symptomatic disease due to Omicron, at ≥15 weeks after the second dose. However, the authors note that early data for AstraZeneca-Vaxzevria are likely biased due to small numbers and because persons receiving two doses of the vaccine likely reflect an older population and those with more co-morbidities. VE of Pfizer BioNTech-Comirnaty against symptomatic disease was similar to Delta (88.0%, 95% CI: 65.9-95.8%) 2-9 weeks post complete vaccination, but then fell to 48.5% (24.3-65.0%) at 10-14 weeks post second dose (compared to VE of 77.7%, 95% CI: 76.3-79%, against Delta 10-14 weeks post vaccination). VE against symptomatic disease due to Omicron remained 34-37% from

15-to-25+ weeks post second dose, without evidence of further decrease. Two weeks after receiving a third dose of Pfizer BioNTech-Comirnaty, VE against symptomatic disease due to Omicron increased to 71.4% (41.8-86.0%) among those who had received a primary series of AstraZeneca-Vaxzevria and to 75.5% (95%CI: 56.1 to 86.3%) among those who had received a primary series of Pfizer BioNTech-Comirnaty; in contrast, VE of a third dose against symptomatic disease due to Delta was 93-94% when added to either primary series. While these results indicate reduced VE of AstraZeneca-Vaxzevria and Pfizer BioNTech-Comirnaty against Omicron, significant bias cannot be ruled out; differences in age and risk among early cases of Omicron among vaccinated versus unvaccinated persons, as well as the predominance of early Omicron cases among travelers and their close contacts, could explain some of the results. Moreover, due to the small number of Omicron cases detected to date, these early VE estimates are subject to significant uncertainty with wide confidence intervals.

Six studies (not yet peer reviewed) have assessed the ability of blood collected from vaccinated persons to neutralize the Omicron variant.

- One study found an average 41.3-fold reduction in neutralization capacity relative to the ancestral SARS-CoV-2 strain in 12 samples collected 10-39 days after complete vaccination with the Pfizer BioNTech-Comirnaty primary (two-dose) series.<sup>5</sup>
- A second study conducted by Pfizer, found a 25.8-fold reduction relative to the ancestral strain among approximately 20 samples collected 3 weeks after completion of the Pfizer BioNTech-Comirnaty primary series.
   The reduction was only 2.6-fold among samples collected from persons who had received a third dose of Pfizer BioNTech-Comirnaty one month prior to sample collection.<sup>36</sup>
- A third study found that neutralization capacity against Omicron was reduced by 33.5-fold relative to the ancestral strain in persons receiving 2 doses of Pfizer BioNTech-Comirnaty, while the majority of samples from persons receiving two doses of AstraZeneca-Vaxzevria, failed to neutralize the variant.<sup>37</sup>
- A fourth study found that blood collected from individuals who were previously vaccinated with 2 doses of mRNA vaccine 1.3 months prior showed a 127-fold reduction in capacity to neutralize Omicron relative to the ancestral strain; samples collected from persons vaccinated with Janssen-Ad26.COV 2.S 1 month prior failed to neutralize Omicron.<sup>38</sup>
- A fifth study found 20-fold, 11.4- fold, and 10-fold reductions in neutralization capacity compared to Delta for two doses of Moderna-mRNA-1273, for two doses of Pfizer BioNTech-Comirnaty, and for a single dose of AstraZeneca-Vaxzevria followed by second dose of Pfizer BioNTech-Comirnaty, respectively, among 14-19 samples collected from persons who had received their second dose 6-7 months prior.<sup>39</sup> Previous studies have found a median 3-fold reduction (IQR 2-4) of these vaccines against Delta relative to the ancestral strain. Adding a third dose of Pfizer BioNTech-Comirnaty to each of three primary vaccination series evaluated, an increase in neutralizing antibodies was observed relative to two doses, however, neutralization capacity against Omicron relative to Delta was still reduced by 23- to 37-fold.
- Finally, a sixth study found that blood collected from persons vaccinated with 2 doses of AstraZeneca-Vaxzevria, 2 doses of Moderna-mRNA-1273, 2 doses of Pfizer BioNTech-Comirnaty, or 1 dose of AstraZeneca-Vaxzevria followed by a second dose of Pfizer BioNTech-Comirnaty had reduced capacity to neutralize Omicron as *compared to Alpha, Beta, and Delta variants*; a much smaller reduction in neutralization capacity against Omicron was observed for blood collected from persons who had been previously infected and then vaccinated or previously vaccinated and then infected. Of note, these neutralization studies used different assays, sera at variable times after vaccination, and most included sera from a small number of persons.

While methods vary across the studies, and neutralization is only one marker of vaccine performance, these preliminary laboratory results suggest that the effectiveness of COVID-19 vaccines against infection with the Omicron variant may be reduced.

Three studies assessed COVID-19 vaccine effectiveness in settings where Delta was the predominant circulating variant.

- The first test-negative case-control peer-reviewed study conducted at two medical centers in India found AstraZeneca-Vaxzevria to be 63.1% (51.5-72.1) effective at preventing SARS-CoV-2 infection 14 or more days post second dose, with a maximum follow-up time up to 10 weeks following the second dose. 40 Authors also report that persons infected with SARS-CoV-2, two doses of AstraZeneca-Vaxzevria was 81.5% (9.9-99.0%) effective at preventing progression to moderate-to-severe disease.
- A second peer-reviewed retrospective cohort study from Israel evaluated the effectiveness of a third dose of Pfizer BioNTech-Comirnaty in preventing death among persons 50 years and older who had completed the primary vaccination series at least 5 months prior.<sup>41</sup> The authors found that a third dose<sup>41</sup> had a relative VE of 90% (86-93%) effective at preventing death due to COVID-19, compared to those who had received only 2 doses; the rate of death in the third dose group was 0.16 per 100,000 person-years compared to 2.98 per 100,000 person years in persons with 2 doses only.
- A third retrospective cohort study from Israel (not yet peer reviewed), found the rate of SARS-CoV-2 infection to be 2.6 (2.4-2.7) time lower in persons having received a third dose of Pfizer BioNTech-Comirnaty in the previous two months relative to persons who had received their second dose in the prior two months.<sup>42</sup> The study also found evidence of decreasing VE of two doses of Pfizer BioNTech-Comirnaty over time, with a 4-fold increase in the rate of infection among those receiving a second dose 6-8 months prior compared to those who had received their second dose only 0-2 months earlier.

Table 3. Summary of primary series vaccine performance against variants of concern (data as of 12 December 2021)

Table 3. Summary of pri						data as of 12 D	ecember 202	21)			
	WHO Emergency Use Listing (EUL) Qualified Vaccines*								Vaccines without WHO EUL <sup>+</sup>		
	AstraZeneca- Vaxzevria/SII - Covishield	Beijing CNBG- BBIBP CorV	Bharat-Covaxi	Janssen-Ad26.COV 2.S	Moderna-mRNA- 1273	Moderna-mRNA- 1273/ Pfizer BioN BioNTech-Comirnaty	Pfizer BioNTech- Comirnaty	Sinovac-CoronaVac	Anhui ZL- Recombinant	Gamaleya-Sputnik V	Novavax-Covavax
Alpha <sup>43,44</sup>											
Summary of VE*				Protecti	on retained aga	inst all outcomes	·				
- Severe disease	$\leftrightarrow_2$	-	-	-	$\leftrightarrow_2$	$\leftrightarrow_1$	$\leftrightarrow_6$	-	-	-	-
- Symptomatic disease	↔to↓₅	-	-	-	$\longleftrightarrow_1$	$\leftrightarrow_1$	$\longleftrightarrow_4$	-	-	-	$\downarrow_1$
- Infection	$\leftrightarrow$ to $\downarrow_4$	-	-		$\longleftrightarrow_3$	-	$\longleftrightarrow_3$	-	-	-	-
Neutralization	<>to↓8	$\longleftrightarrow_1$	$\leftrightarrow_2$	$\longleftrightarrow_4$	↔to↓₃	↔to↓₃	to↓₄₃	$\leftrightarrow$ to $\downarrow\downarrow_7$	$\longleftrightarrow_2$	$\leftrightarrow$ to $\downarrow_4$	<b>↓</b> 1
Beta <sup>45–48</sup>											
Summary of VE*			Protection	retained against s	severe disease; r	educed protection ag	gainst symptomat	ic disease; limite	ed evidence		
- Severe disease	-	-	-	$\leftrightarrow_1$	$\leftrightarrow_1$	-	$\leftrightarrow_3$	-	-	-	-
- Symptomatic disease	$\leftrightarrow$ to $\downarrow\downarrow\downarrow_2$	-	-	$\leftrightarrow_1$	$\leftrightarrow_1$	-	$\leftrightarrow_2$	-	-	-	$\downarrow\downarrow\downarrow\downarrow_1$
- Infection	-	-	-	-	$\leftrightarrow_1$	-	$\downarrow_1$	-	-	-	-
Neutralization	↓to↓↓ <sub>8</sub>	⇔to↓ <sub>2</sub>	<b>√</b> 2	↓to↓↓ <sub>8</sub>	↓to↓↓ <sub>17</sub>	<b>↓</b> ↓₃	↓to↓↓₄₃	$\sqrt{to}\sqrt{\sqrt{16}}$	⇔to↓₃	$\downarrow\downarrow\downarrow$ to $\downarrow\downarrow\downarrow\downarrow$ 5	$\downarrow\downarrow\downarrow\downarrow_1$
Gamma											
Summary of VE*	Unclear impact; very limited evidence										
- Severe disease	$\leftrightarrow_1$	-	-	-	$\leftrightarrow_1$	-	$\leftrightarrow_2$	-	-	-	-
- Symptomatic disease	$\leftrightarrow_1$	-	-	-	$\leftrightarrow_1$	-	$\leftrightarrow_1$	-	-	-	-
- Infection	$\leftrightarrow_1$	-	-	-	$\leftrightarrow_1$	-	$\leftrightarrow_1$	$\longleftrightarrow_1$	-	-	-
Neutralization	↔to↓₃	-	-	<b>⇔</b> to↓₄	<b>√</b> 8	$\leftrightarrow_1$	↔to↓₂₅	<b>↓</b> 5	$\leftrightarrow_1$	↓to↓↓3	-
Delta <sup>49</sup>								1		• •	
Summary of VE*		Protect	tion retained ag	gainst severe disea	ise; possible red	luced protection agai	nst symptomatic (	disease and inte	ction; ilmited	evidence	
- Severe disease	$\leftrightarrow_3$	-	-	-	$\leftrightarrow_3$	-	$\leftrightarrow_6$	-	-	-	-
- Symptomatic disease	$\sqrt{to}\sqrt{\sqrt{s}}$	-	$\downarrow_1$	-	$\leftrightarrow_1$	-	$\leftrightarrow$ to $\downarrow_4$	-	-	-	-
- Infection	$\leftrightarrow$ to $\downarrow_4$	-	-	$\downarrow\downarrow\downarrow\downarrow_1$	$\leftrightarrow_3$	-	$\leftrightarrow$ to $\downarrow_3$	-	-	-	-
Neutralization	$\downarrow_{\mathfrak{D}}$	-	$\leftrightarrow$ to $\downarrow_3$	$\leftrightarrow$ to $\downarrow\downarrow_8$	<b>√</b> 9	$\sqrt{10}\sqrt{13}$	↔to↓ <sub>23</sub>	$\downarrow$ to $\downarrow\downarrow\downarrow\downarrow_5$	$\leftrightarrow$ to $\downarrow_2$	$\sqrt{to}\sqrt{\sqrt{3}}$	-
Omicron											
Summary of VE*						No evidence					
- Severe disease	-	-	-	-	-	-	-	-	-	-	-
- Symptomatic disease	-	-	-	-	-	-	$\downarrow\downarrow\downarrow\downarrow_1$	-	-	-	-
- Infection	-	-	-	-	-	-	-	-	-	-	-
Neutralization	$\downarrow\downarrow\downarrow\downarrow_1$	-	-	$\downarrow\downarrow\downarrow\downarrow_1$	-	$\downarrow\downarrow\downarrow\downarrow_1$	$\downarrow\downarrow\downarrow\downarrow_3$	-	-	-	-

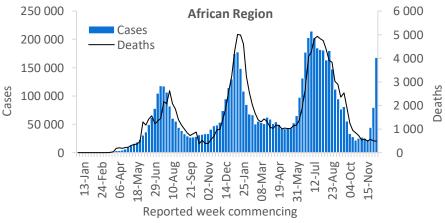
VE refers to vaccine effectiveness and vaccine efficacy. \*Summary of VE: indicates the general conclusions but only for the vaccines evaluated against the specific variant. Arrows generalize the magnitude of reduction in VE or neutralization: " $\leftrightarrow$ " <10% reduction in VE, or VE >90% with no comparator, or that there was a <2-fold reduction in neutralization; " $\downarrow$ " 10 to <20% reduction in VE, or 2 to <5-fold reduction in neutralization; " $\downarrow$ " 20% reduction in NE, or 5 to <10-fold reduction in neutralization; " $\downarrow$ " ≥30% reduction in VE, or ≥10-fold reduction in neutralization. When more than one neutralization study is available, the interquartile range (25th and 75th percentiles) of fold-reductions across all studies for specific vaccine/variant was used. "Moderna-mRNA-1273/Pfizer BioNTech-Comirnaty" indicates that both vaccines were evaluated together in study. The number of studies is shown as subscripts: vaccine effectiveness and neutralization studies informing this table can be found on the VIEW-hub Resources Library. References indicated by superscripts next to VOC name in column 1 are vaccine efficacy results from randomized controlled trials informing this table.

# WHO regional overviews Epidemiological week 6 – 12 December 2021

## **African Region**

The African Region reported over 167 000 new cases, an increase of 111% as compared to the previous week and the highest number of new weekly cases since early August 2021. Marked increases were observed in over two thirds (33/49; 67%) of countries in the Region with the majority (30/33; 91%) reporting increases of 25% or greater, as compared to the previous week. The highest numbers of new cases were reported from South Africa (109 053 new cases; 183.9 new cases per 100 000 population; a 76% increase), Zimbabwe (26 479 new cases; 178.2 new cases per 100 000; a 479% increase), and Mauritius (6415 new cases; 504.4 new cases per 100 000; a 775% increase).

The Region reported just under 500 new deaths, a number similar to the number reported in the previous week. The highest numbers of new deaths were reported from South Africa (151 new deaths; <1 new death per 100 000 population; a 13% decrease), Mauritius (92 new deaths; 7.2 new deaths per 100 000; a 27% decrease), and Algeria (41 new deaths; <1 new death per 100 000; a 7% decrease).

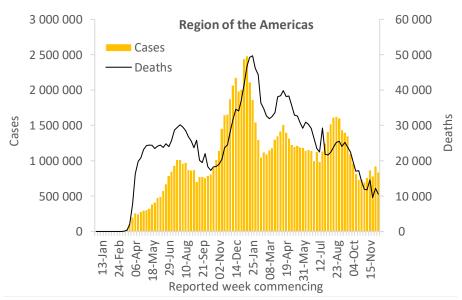


#### Updates from the African Region

## **Region of the Americas**

The Region of the Americas reported over 837 000 new cases and over 10 000 new deaths, decreases of 10% and 14% respectively as compared to the previous week. Nevertheless, 28% (15/56) of countries in the Region reported over 10% increases in cases, with the greatest observed in the Caribbean islands of Saint Barthélemy (350%; from 2 cases to 9 cases), Turks and Caicos Islands (285%) and Saint Martin (111%). The highest numbers of new cases were reported from the United States of America (674 019 new cases; 203.6 new cases per 100 000; a 9% decrease), Brazil (38 372 new cases; 18.1 new cases per 100 000; a 38% decrease), and Canada (25 332 new cases; 67.1 new cases per 100 000; a 25% increase).

The highest numbers of new deaths were reported from the United States of America (6909 new deaths; 2.1 new deaths per 100 000; a 16% decrease), Mexico (1122 new deaths; <1 new death per 100 000; an 85% increase), and Brazil (851 new deaths; <1 new death per 100 000; a 41% decrease).

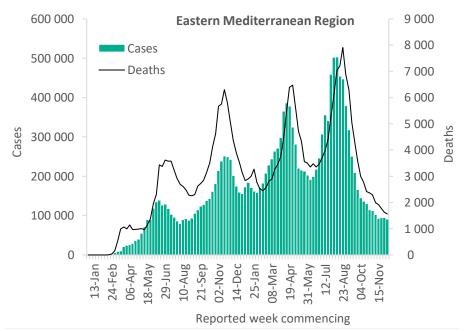


Updates from the Region of the Americas

## **Eastern Mediterranean Region**

The weekly incidence of cases and deaths in the Eastern Mediterranean Region remained stable this week, with over 90 000 new cases and over 1500 new deaths reported. However, three countries (3/22, 13%) in the Region reported an increase of over 10% in weekly incidence of cases. The highest numbers of new cases were reported from Jordan (34 735 new cases; 340.4 new cases per 100 000; an 8% increase), the Islamic Republic of Iran (21 168 new cases; 25.2 new cases per 100 000; a 19% decrease), and Lebanon (11 341 new cases; 166.2 new cases per 100 000; a 9% increase).

The highest numbers of new deaths continued to be reported from the Islamic Republic of Iran (537 new deaths; <1 new death per 100 000; a 7% decrease), Egypt (333 new deaths; <1 new death per 100 000; a 12% decrease), and Jordan (226 new deaths; 2.2 new deaths per 100 000; a 13% increase).

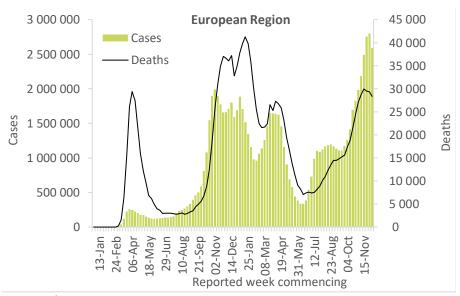


Updates from the Eastern Mediterranean Region

### **European Region**

The European Region reported just under 2.6 million new cases, a 7% decrease as compared to the previous week and a decline since early September. The number of new deaths reported this week was just over 28 000, remaining similar to the number reported in the previous week. Despite the declining trend, a small proportion (10/61; 6%) of countries still reported over a 10% increase in cases as compared to the previous week. The highest numbers of new cases were reported from Germany (351 738 new cases; 422.9 new cases per 100 000; an 11% decrease), the United Kingdom (350 340 new cases; 516.1 new cases per 100 000; a 13% increase), and France (335 972 new cases; 516.6 new cases per 100 000; a 19% increase).

The highest numbers of new deaths were reported from the Russian Federation (8205 new deaths; 5.6 new deaths per 100 000; a similar number to that of the previous week), Poland (2804 new deaths; 7.4 new deaths per 100 000; a 6% increase), and Ukraine (2747 new deaths; 6.3 new deaths per 100 000; a 13% decrease).

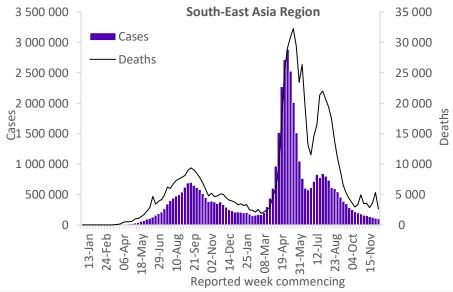


Updates from the European Region

## **South-East Asia Region**

The declining trend in reported new weekly cases and deaths continued this week in the South-East Asia Region. Over 98 000 new cases and over 2600 new deaths were reported, amounting to a 10% and 50% decrease respectively as compared to the previous week. Only two countries reported an increase in weekly cases, Bangladesh (from 1659 to 1882, a 13% increase) and Bhutan (from 1 to 8; a 700% increase). The highest numbers of new cases were reported from India (57 255 new cases; 4.1 new cases per 100 000; a 6% decrease), Thailand (27 405 new cases; 39.3 new cases per 100 000; a 20% decrease), and Sri Lanka (5220 new cases; 24.4 new cases per 100 000; similar to the number reported in the previous week).

The highest numbers of new deaths were reported from India (2108 new deaths; <1 new death per 100 000; a 56% decrease), Thailand (227 new deaths; <1 new death per 100 000; similar to the number reported in the previous week), and Sri Lanka (153 new deaths; <1 new death per 100 000; similar to the number reported in the previous week).

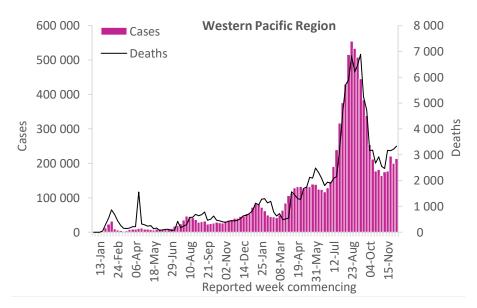


Updates from the South-East Asia Region

## **Western Pacific Region**

The Western Pacific Region reported 214 000 new cases, a 7% increase as compared to the previous week. Four of the 27 countries in the region, reported an increase in case incidence of over 10%, Northern Mariana Islands (62%), Republic of Korea (37%), Lao People's Democratic Republic (17%) and Japan (12%). The highest numbers of new cases were reported from Viet Nam (103 635 new cases; 106.5 new cases per 100 000; a 6% increase), Republic of Korea (44 238 new cases; 86.3 new cases per 100 000; a 38% increase), and Malaysia (33 675 new cases; 104.0 new cases per 100 000; similar to the number reported in the previous week).

The Region reported over 3300 new deaths, a number similar to that of the previous week. The highest numbers of new deaths were reported from Viet Nam (1550 new deaths; 1.6 new deaths per 100 000; a 13% increase), the Philippines (866 new deaths; <1 new death per 100 000; a 16% decrease), and Republic of Korea (401 new deaths; <1 new death per 100 000; a 32% increase).



Updates from the Western Pacific Region

## **Summary of the COVID-19 Weekly Operational Update**

The <u>Weekly Operational Update</u> is a report provided by the COVID-19 Strategic Preparedness and Response Plan (SPRP) Monitoring and Evaluation team, which aims to update on the ongoing global progress against the <u>COVID-19 SPRP 2021</u> framework, and to highlight country-level actions and WHO support to countries. In this week's edition published on 13 December, highlights include the following:

- WHO provides testing kits to Uganda for screening the COVID-19 Omicron variant of concern
- COVID-19 Contact Tracing Communication in Honduras
- WHO/Europe laboratory system strengthening mission to Kazakhstan
- UN agencies support intensive COVID-19 vaccination drive in the Philippines
- Emergency Medical Teams (EMT) in the Pacific: Strengthening national capacity for health emergency response
- Updates on WHO's financing to support countries on COVID-19 response implementation to suppress transmission, reduce exposure, and protect the vulnerable and save lives
- Progress on a subset of global indicators that demonstrate country and global progress to end the acute phase of the pandemic

## **Technical guidance and other resources**

- WHO technical guidance
- WHO COVID-19 Dashboard
- WHO Weekly Operational Updates on COVID-19
- WHO COVID-19 case definitions
- COVID-19 Supply Chain Inter-Agency Coordination Cell Weekly Situational Update
- Research and Development
- Open WHO courses on COVID-19 in official UN languages and in additional national languages
- WHO Academy COVID-19 mobile learning app
- <u>The Strategic Preparedness and Response Plan</u> (SPRP) outlining the support the international community can provide to all countries to prepare and respond to the virus
- EPI-WIN: tailored information for individuals, organizations, and communities
- Recommendations and advice for the public:
  - Protect yourself
  - Questions and answers
  - Travel advice

#### **Annexes**

#### Annex 1. Additional notes on VOC impacts on vaccines

- Reductions in VE do not necessarily mean loss of protection, as indicated by the absolute VE estimate. For example, a 10-percentage point reduction in VE against symptomatic disease for mRNA vaccines would still mean high vaccine effectiveness of ~85%. Likewise, vaccines have shown higher VE against severe disease; thus, small reductions in VE against severe disease due to VOCs may still mean substantial protection.
- Table 3 summarizes the impact of VOCs on COVID-19 vaccine performance in the absence of waning, and, therefore, does not include studies that only assess VE greater than 4 months post final dose.
- Studies reporting VOC-specific VE estimates for full vaccination (≥7 days post final dose) are assessed against
  a comparator VE estimate for that vaccine product to determine level of reduction in VE. For symptomatic
  disease, VOC VE is compared against phase 3 RCT results from non-VOC settings. For severe disease and
  infection, due to instability or lack of phase 3 RCT estimates, VOC VE is compared to non-VOC VE estimates
  from the same study when available (or to Alpha VE from same study when assessing Beta, Gamma, or Delta);
  with an exception for AstraZeneca-Vaxzevria for infection (when a phase 3 estimate of VE against infection
  due to non-VOC is available and used as comparator). In some instances, a study may be included for severe
  disease or infection outcome even without a comparator if a very high VE estimate is reported against a VOC
  (i.e., >90%).
- It is also important to note that studies vary in population, outcome definitions, study design and other
  methodological considerations, which may in part explain differences when comparing VE estimates for a
  product between different studies. In addition, the reductions summarized in the table represent VE point
  estimates and do not represent the uncertainty intervals around these estimates which vary substantially
  across studies. The reductions in VE noted should be interpreted with these limitations in mind.
- Neutralization studies that use samples collected >7 days and < 6 months after complete vaccination and that use an ancestral strain as the reference are included in Table 3.

Annex 2. List of countries/territories/areas reporting variants of concern as of 14 December 2021

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Afghanistan	•	-	•	-	-
Albania	•	-	0	-	-
Algeria	•	-	•	-	-
Andorra	0	0	0	-	-
Angola	•	•	•	•	-
Anguilla	•	-	•	-	-
Antigua and Barbuda	•	•	•	•	-
Argentina	•	•	•	•	•
Armenia	•	-	•	-	-
Aruba	•	•	•	•	-
Australia	•	•	•	•	•
Austria	•	•	•	•	•
Azerbaijan	•	-	0	-	-
Bahamas	•	-	•	•	-
Bahrain	•	•	•	•	•*
Bangladesh	•	•	•	0	•*
Barbados	•	-	•	•	-
Belarus	•	-	0	-	-
Belgium	•	•	•	•	•
Belize	•	-	•	•	-
Benin	•	•	•	•	-
Bermuda	•	•	•	-	•*
Bhutan	•	•	•	-	-
Bolivia (Plurinational State of)	•	-	•	•	-
Bonaire	•	-	•	•	-
Bosnia and Herzegovina	•	•	0	•	-
Botswana	0	•	•	-	•
Brazil	•	•	•	•	•
British Virgin Islands	•	-	•	•	-
Brunei Darussalam	•	•	•	-	-

Bulgaria  Burkina Faso  Burundi  Cabo Verde  Cambodia	- - -	-
Burundi • • • • Cabo Verde • - •		-
Cabo Verde • - •	-	_
	-	-
Cambodia • • •		-
	-	-
Cameroon • • •	•	-
Canada ● ● ●	•	•
Cayman Islands • • •	•	-
Central African Republic • • •	-	-
Chad •	-	-
Chile • • •	•	•
China • • •	•	•
Colombia • - •	•	-
Comoros - • •	-	-
Congo • • •	•	-
Costa Rica • • •	•	-
Croatia • • o	•	•
Cuba • • •	-	•*
Curação • • •	•	-
Cyprus • • o	-	•*
Czechia • • •	•	•
Côte d'Ivoire   ● • ○	-	-
Democratic Republic of the Congo	-	-
Denmark ● ● ●	•	•
Djibouti	-	-
Dominica • - •	-	-
Dominican Republic • - •	•	-
Ecuador • - •	•	-
Egypt • - •	-	-
El Salvador • - •	•	-

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Equatorial Guinea	•	•	•	-	-
Estonia	•	•	0	0	•
Eswatini	0	•	•	-	-
Ethiopia	•	•	•	-	-
Falkland Islands (Malvinas)	•	•	-	-	-
Faroe Islands	•	-	-	•	-
Fiji	0	-	•	-	•
Finland	•	•	•	•	•
France	•	•	•	•	•
French Guiana	•	•	•	•	-
French Polynesia	•	•	•	•	-
Gabon	•	•	•	-	-
Gambia	•	-	•	-	-
Georgia	•	0	•	-	-
Germany	•	•	•	•	•
Ghana	•	•	•	•	•
Gibraltar	•	-	0	-	•*
Greece	•	•	•	•	•
Greenland	-	-	•	-	-
Grenada	•	-	•	•	-
Guadeloupe	•	•	•	•	-
Guam	•	•	•	•	-
Guatemala	•	•	•	•	-
Guinea	•	•	•	-	-
Guinea-Bissau	•	•	•	-	-
Guyana	-	-	•	•	-
Haiti	•	-	•	•	-
Honduras	•	-	•	•	-
Hungary	•	0	0	•	•*
Iceland	•	•	•	•	•

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
India	•	•	•	•	•
Indonesia	•	•	•	-	-
Iran (Islamic Republic of)	•	•	•	-	-
Iraq	•	•	•	•	-
Ireland	•	•	•	•	•
Israel	•	•	•	•	•
Italy	•	•	•	•	•
Jamaica	•	-	•	-	-
Japan	•	•	•	•	•
Jordan	•	•	•	•	•*
Kazakhstan	•	0	•	-	-
Kenya	•	•	•	-	-
Kosovo[1]	•	0	0	-	-
Kuwait	•	•	•	-	•*
Kyrgyzstan	•	•	•	-	-
Lao People's Democratic Republic	•	-	•	-	-
Latvia	•	•	0	•	•
Lebanon	•	-	•	-	•*
Lesotho	-	•	0	-	-
Liberia	•	•	•	-	-
Libya	•	•	-	-	-
Liechtenstein	•	-	0	0	0*
Lithuania	•	•	0	•	-
Luxembourg	•	•	•	•	•
Madagascar	•	•	-	-	-
Malawi	•	•	•	-	•*
Malaysia	•	•	•	-	•
Maldives	•	-	•	-	•
Mali	-	-	•	-	-
Malta	•	0	0	•	-
Martinique	•	•	•	•	-

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Mauritania	•	•	•	-	-
Mauritius	•	•	•	-	0*
Mayotte	•	•	0	-	-
Mexico	•	•	•	•	•
Monaco	•	•	•	-	-
Mongolia	•	-	•	-	-
Montenegro	•	-	0	0	0
Montserrat	•	-	•	•	-
Morocco	•	•	•	-	-
Mozambique	•	•	•	-	-
Myanmar	•	-	•	-	-
Namibia	•	•	•	•	•
Nepal	•	-	•	-	•
Netherlands	•	•	•	•	•
New Caledonia	•	-	•	-	-
New Zealand	•	•	•	•	-
Nicaragua	•	•	•	•	-
Niger	•	-	•	-	-
Nigeria	•	•	•	-	•
North Macedonia	•	•	0	-	-
Northern Mariana Islands (Commonwealth of the)	0	-	•	-	-
Norway	•	•	•	•	•
Occupied Palestinian Territory	•	•	•	-	_
Oman	•	•	•	-	-
Pakistan	•	•	•	•	•*
Panama	•	•	•	•	-
Papua New Guinea	-	-	•	-	-
Paraguay	•	-	•	•	-
Peru	•	-	•	•	-
Philippines	•	•	•	•	-
Poland	•	0	•	•	-

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Portugal	•	•	•	•	•
Puerto Rico	•	•	•	•	0*
Qatar	•	•	•	-	-
Republic of Korea	•	•	•	•	•
Republic of Moldova	•	-	•	-	-
Romania	•	•	•	•	•
Russian Federation	•	•	•	0	•
Rwanda	•	•	•	-	-
Réunion	•	•	0	•	•
Saba	-	-	•	-	-
Saint Barthélemy	•	-	•	-	-
Saint Kitts and Nevis	-	-	•	-	-
Saint Lucia	•	-	•	-	-
Saint Martin	•	•	•	-	-
Saint Pierre and Miquelon	-	-	•	-	-
Saint Vincent and the Grenadines	-	-	•	•	-
Sao Tome and Principe	•	-	0	-	-
Saudi Arabia	•	•	•	-	•
Senegal	•	•	•	-	•
Serbia	•	-	•	-	-
Seychelles	•	•	•	-	-
Sierra Leone	-	•	•	-	•*
Singapore	•	•	•	•	•
Sint Maarten	•	•	•	•	-
Slovakia	•	•	•	-	0*
Slovenia	•	•	•	•	-
Somalia	•	•	•	-	-
South Africa	•	•	•	0	•
South Sudan	•	•	•	-	-
Spain	•	•	•	•	•
Sri Lanka	•	•	•	-	•

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Sudan	•	•	-	•	-
Suriname	•	•	•	•	-
Sweden	•	•	•	•	•
Switzerland	•	•	•	•	•
Thailand	•	•	•	•	•
Timor-Leste	•	-	•	-	-
Togo	•	•	•	•	-
Trinidad and Tobago	•	-	•	•	•*
Tunisia	•	•	•	-	•
Turkey	•	•	•	•	0*

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Turks and Caicos Islands	•	-	•	•	-
Uganda	•	•	•	-	•
Ukraine	•	0	0	-	-
United Arab Emirates	•	•	•	•	•
United Kingdom	•	•	•	•	•
United Republic of Tanzania	-	•	-	-	-
United States Virgin Islands	•	•	•	•	-
United States of America	•	•	•	•	•
Uruguay	•	•	•	•	-
Uzbekistan	•	•	0	-	-

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Vanuatu	-	-	•	-	-
Venezuela (Bolivarian Republic of)	•	-	•	•	-
Viet Nam	•	•	•	-	-
Wallis and Futuna	•	-	-	-	-
Yemen	•	•	-	-	-
Zambia	•	•	•	-	•
Zimbabwe	•	•	•	-	•

<sup>\*</sup>Newly reported in this update. "•" indicates that information for this variant was received by WHO from official sources. "o" indicates that information for this variant was received by WHO from unofficial sources and will be reviewed as more information become available. \*\*Includes countries/territories/areas reporting the detection of VOCs among travellers (e.g., imported cases detected at points of entry), or local cases (detected in the community). Excludes countries, territories, and areas that have never reported the detection of a variant of concern. See also Annex 2: Data, table, and figure notes

#### Annex 3. Data, table, and figure notes

Data presented are based on official laboratory-confirmed COVID-19 case and deaths reported to WHO by country/territories/areas, largely based upon WHO <u>case definitions</u> and <u>surveillance guidance</u>. While steps are taken to ensure accuracy and reliability, all data are subject to continuous verification and change, and caution must be taken when interpreting these data as several factors influence the counts presented, with variable underestimation of true case and death incidences, and variable delays to reflecting these data at the global level. Case detection, inclusion criteria, testing strategies, reporting practices, and data cut-off and lag times differ between countries/territories/areas. A small number of countries/territories/areas report combined probable and laboratory-confirmed cases. Differences are to be expected between information products published by WHO, national public health authorities, and other sources.

Due to public health authorities conducting data reconciliation exercises that remove large numbers of cases or deaths from their total counts, negative numbers may be displayed in the new cases/deaths columns as appropriate. When additional details become available that allow the subtractions to be suitably apportioned to previous days, graphics will be updated accordingly. A record of historic data adjustment made is available upon request by emailing <a href="mailto:epi-data-support@who.int">epi-data-support@who.int</a>. Please specify the countries of interest, time period, and purpose of the request/intended usage. Prior situation reports will not be edited; see <a href="mailto:covid19.who.int">covid19.who.int</a> for the most up-to-date data. COVID-19 confirmed cases and deaths reported in the last seven days by countries, territories, and areas, and WHO Region (reported in previous issues) are now available at: <a href="https://covid19.who.int/table">https://covid19.who.int/table</a>.

'Countries' may refer to countries, territories, areas or other jurisdictions of similar status. The designations employed, and the presentation of these materials do not imply the expression of any opinion whatsoever on the part of WHO concerning the legal status of any country, territory, or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted and dashed lines on maps represent approximate border lines for which there may not yet be full agreement. Countries, territories, and areas are arranged under the administering WHO region. The mention of specific companies or of certain manufacturers' products does not imply that they are endorsed or recommended by WHO in preference to others of a similar nature that are not mentioned. Errors and omissions except, the names of proprietary products are distinguished by initial capital letters.

[1] All references to Kosovo should be understood to be in the context of the United Nations Security Council resolution 1244 (1999). In the map, the number of cases of Serbia and Kosovo (UNSCR 1244, 1999) have been aggregated for visualization purposes.

#### References

- 1. SARS-CoV-2 variants of concern and variants under investigation https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment\_data/file/1040076/Technical\_Briefing\_3 1.pdf. :42.
- 2. CDCMMWR. SARS-CoV-2 B.1.1.529 (Omicron) Variant United States, December 1–8, 2021. MMWR Morb Mortal Wkly Rep. 2021;70. doi:10.15585/mmwr.mm7050e1
- 3. Published. Preliminary findings from study after Christmas party in Oslo. Norwegian Institute of Public Health. Accessed December 14, 2021. https://www.fhi.no/en/news/2021/preliminary-findings-from-outbreak-investigation-after-christmas-party-in-o/
- 4. Kuhlmann C, Mayer CK, Claassen M, et al. *Breakthrough Infections with SARS-CoV-2 Omicron Variant Despite Booster Dose of MRNA Vaccine*. Social Science Research Network; 2021. doi:10.2139/ssrn.3981711
- 5. Cele S, Jackson L, Khan K, et al. SARS-CoV-2 Omicron Has Extensive but Incomplete Escape of Pfizer BNT162b2 Elicited Neutralization and Requires ACE2 for Infection.; 2021:2021.12.08.21267417. doi:10.1101/2021.12.08.21267417
- 6. Roessler A, Riepler L, Bante D, Laer D von, Kimpel J. SARS-CoV-2 B.1.1.529 Variant (Omicron) Evades Neutralization by Sera from Vaccinated and Convalescent Individuals.; 2021:2021.12.08.21267491. doi:10.1101/2021.12.08.21267491
- 7. Sheward et al. Preliminary Report Early release, subject to modification Quantification of the neutralization resistance of the Omicron Variant of Concern. Google Docs. Accessed December 14, 2021.
- https://drive.google.com/file/d/1CuxmNYj5cpluxWXhjjVmuDqntxXwlfXQ/view?usp=embed\_facebook
- 8. Redd AD, Nardin A, Kared H, et al. *Minimal Cross-over between Mutations Associated with Omicron Variant of SARS-CoV-2 and CD8+ T Cell Epitopes Identified in COVID-19 Convalescent Individuals*. Immunology; 2021. doi:10.1101/2021.12.06.471446
- 9. Kim P, Gordon SM, Sheehan MM, Rothberg MB. Duration of SARS-CoV-2 Natural Immunity and Protection against the Delta Variant: A Retrospective Cohort Study. *Clinical Infectious Diseases*. Published online December 3, 2021:ciab999. doi:10.1093/cid/ciab999
- 10. Hwang H, Lim JS, Song SA, et al. Transmission dynamics of the Delta variant of SARS-CoV-2 infections in South Korea. *The Journal of Infectious Diseases*. Published online December 2, 2021:jiab586. doi:10.1093/infdis/jiab586
- Buchan SA, Tibebu S, Daneman N, et al. Increased household secondary attacks rates with Variant of Concern SARS-CoV-2 index cases. *Clinical Infectious Diseases*. 2021;(ciab496). doi:10.1093/cid/ciab496
- 12. Tegally H, Wilkinson E, Giovanetti M, et al. Emergence of a SARS-CoV-2 variant of concern with mutations in spike glycoprotein. *Nature*. Published online 2021. https://doi.org/10.1038/s41586-021-03402-9
- 13. Sinha S, Tam B, Wang SM. Altered interaction between RBD and ACE2 receptor contributes towards the increased transmissibility of SARS CoV-2 delta, kappa, beta, and gamma strains with RBD double mutations. *bioRxiv*. Published online January 1, 2021:2021.08.30.458303. doi:10.1101/2021.08.30.458303
- 14. Curran J, Dol J, Boulos L, et al. Transmission characteristics of SARS-CoV-2 variants of concern Rapid Scoping Review. *medRxiv*. Published online January 1, 2021:2021.04.23.21255515. doi:10.1101/2021.04.23.21255515
- 15. Campbell F, Archer B, Laurenson-Schafer H, et al. Increased transmissibility and global spread of SARS-CoV-2 variants of concern as at June 2021. *Eurosurveillance*. 2021;26(24):2100509.
- 16. Dhar MS, Marwal R, Vs R, et al. Genomic characterization and epidemiology of an emerging SARS-CoV-2 variant in Delhi, India. *Science*. Published online October 14, 2021:eabj9932. doi:10.1126/science.abj9932
- 17. Bager P, Wohlfahrt J, Fonager J, Albertsen. Increased Risk of Hospitalisation Associated with Infection with SARS-CoV-2 Lineage B.1.1.7 in Denmark. doi:Bager, Peter and Wohlfahrt, Jan and Fonager, Jannik and Albertsen, Mads and Yssing Michaelsen, Thomas and Holten Møller, Camilla and Ethelberg, Steen and Legarth, Rebecca and Fischer Button, Mia Sara and Gubbels, Sophie Madeleine and Voldstedlund, Marianne and Mølbak, Kåre and Skov, Robert Leo and Fomsgaard, Anders and Grove Krause, Tyra, Increased Risk of Hospitalisation Associated with Infection with SARS-CoV-2 Lineage B.1.1.7 in Denmark. Available at SSRN: https://ssrn.com/abstract=3792894 or http://dx.doi.org/10.2139/ssrn.3792894
- 18. Paredes MI, Lunn SM, Famulare M, et al. Associations between SARS-CoV-2 variants and risk of COVID-19 hospitalization among confirmed cases in Washington State: a retrospective cohort study. *medRxiv*. Published online January 1, 2021:2021.09.29.21264272. doi:10.1101/2021.09.29.21264272
- 19. NERVTAG paper on COVID-19 variant of concern B.1.1.7. *GOVUK*. Published online 2021. https://www.gov.uk/government/publications/nervtag-paper-on-covid-19-variant-of-concern-b117, http://files/64/nervtag-paper-on-covid-19-variant-of-concern-b117.html %[2021/02/08/18:37:19]
- 20. Pascall DJ, Mollett G, Blacow R, Bulteel N, et al. The SARS-CoV-2 Alpha variant causes increased clinical severity of disease. https://www.medrxiv.org/content/10.1101/2021.08.17.21260128v1
- 21. Pearson CA, Eggo. Estimates of severity and transmissibility of novel South Africa SARS-CoV-2 variant 501Y.V2. https://cmmid.github.io/topics/covid19/reports/sa-novel-variant/2021 01 11 Transmissibility and severity of 501Y V2 in SA.pdf

- 22. Freitas ARR, Beckedorff OA, Cavalcanti LP de G, et al. The emergence of novel SARS-CoV-2 variant P.1 in Amazonas (Brazil) was temporally associated with a change in the age and sex profile of COVID-19 mortality: A population based ecological study. *The Lancet Regional Health Americas*. 2021;1:100021. doi:10.1016/j.lana.2021.100021
- 23. Fisman DN, Tuite AR. Progressive Increase in Virulence of Novel SARS-CoV-2 Variants in Ontario, Canada. *medRxiv*. Published online July 12, 2021:2021.07.05.21260050. doi:10.1101/2021.07.05.21260050
- 24. McAlister FA, Nabipoor M, Chu A, Lee DS, Saxinger L, Bakal JA. *Lessons from the COVID-19 Third Wave in Canada: The Impact of Variants of Concern and Shifting Demographics*. Infectious Diseases (except HIV/AIDS); 2021. doi:10.1101/2021.08.27.21261857
- 25. Muik A, Wallisch AK, Sänger B, et al. Neutralization of SARS-CoV-2 lineage B.1.1.7 pseudovirus by BNT162b2 vaccine—elicited human sera. *Science*. Published online 2021:eabg6105.
- 26. Gallais F, Gantner P, Bruel T, et al. Anti-SARS-CoV-2 Antibodies Persist for up to 13 Months and Reduce Risk of Reinfection. *medRxiv*. Published online January 1, 2021:2021.05.07.21256823. doi:10.1101/2021.05.07.21256823
- 27. Wibmer CK, Ayres F, Hermanus T, et al. SARS-CoV-2 501Y.V2 escapes neutralization by South African COVID-19 donor plasma. *Nat Med*. Published online March 2021. https://www.ncbi.nlm.nih.gov/pubmed/33654292
- 28. Sabino EC, Buss LF, Carvalho MPS, et al. Resurgence of COVID-19 in Manaus, Brazil, despite high seroprevalence. *The Lancet*. 2021;397(10273):452-455.
- 29. Public Health England (PHE). SARS-CoV-2 Variants of Concern and Variants under Investigation in England. Technical Briefing 20. Public Health England; 2021.
- $https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment\_data/file/1009243/Technical\_Briefing\_2~0.pdf$
- 30. Planas D, Veyer D, Baidaliuk A, et al. *Reduced Sensitivity of Infectious SARS-CoV-2 Variant B.1.617.2 to Monoclonal Antibodies and Sera from Convalescent and Vaccinated Individuals*. Microbiology; 2021. doi:10.1101/2021.05.26.445838
- 31. Public Health England (PHE). SARS-CoV-2 Variants of Concern and Variants under Investigation..Technical Briefing 18.; 2021. https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment\_data/file/1001358/Variants\_of\_Concern\_VOC\_Technical\_Briefing\_18.pdf
- 32. Classification of Omicron (B.1.1.529): SARS-CoV-2 Variant of Concern. Accessed November 30, 2021. https://www.who.int/news/item/26-11-2021-classification-of-omicron-(b.1.1.529)-sars-cov-2-variant-of-concern
- 33. Public Health England. SARS-CoV-2 lateral flow antigen tests: evaluation of VOC1 (Kent, UK) and VOC2 (South Africa). GOV.UK. Accessed June 21, 2021. https://www.gov.uk/government/publications/sars-cov-2-lateral-flow-antigen-tests-evaluation-of-voc1-and-voc2/sars-cov-2-lateral-flow-antigen-tests-evaluation-of-voc1-kent-uk-and-voc2-south-africa
- 34. Bekliz M, Adea K, Essaidi-Laziosi M, et al. *Analytical Performance of Eleven SARS-CoV-2 Antigen-Detecting Rapid Tests for Delta Variant*. Infectious Diseases (except HIV/AIDS); 2021. doi:10.1101/2021.10.06.21264535
- 35. Andrews N, Stowe J, Kirsebom F, et al. Effectiveness of COVID-19 vaccines against the Omicron (B.1.1.529) variant of concern. :16.
- 36. Pfizer and BioNTech Provide Update on Omicron Variant. Published December 8, 2021. Accessed December 11, 2021. https://www.businesswire.com/news/home/20211208005542/en/Pfizer-and-BioNTech-Provide-Update-on-Omicron-Variant
- 37. Dejnirattisai W, Shaw RH, Supasa P, et al. *Reduced Neutralisation of SARS-COV-2 Omicron-B.1.1.529 Variant by Post-Immunisation Serum.*; 2021:2021.12.10.21267534. doi:10.1101/2021.12.10.21267534
- 38. Schmidt F, Muecksch F, Weisblum Y. Plasma neutralization properties of the SARS-CoV-2 Omicron variant. Google Docs. Accessed December 13, 2021. https://drive.google.com/file/d/1zjJWsybGaa3egiyn5nQqTzBtl0kmvMUu/view?usp=embed\_facebook
- 39. Wilhelm A, Widera M, Grikscheit K, et al. *Reduced Neutralization of SARS-CoV-2 Omicron Variant by Vaccine Sera and Monoclonal Antibodies*. Infectious Diseases (except HIV/AIDS); 2021. doi:10.1101/2021.12.07.21267432
- 40. Thiruvengadam R, Awasthi A, Medigeshi G, et al. Effectiveness of ChAdOx1 nCoV-19 vaccine against SARS-CoV-2 infection during the delta (B.1.617.2) variant surge in India: a test-negative, case-control study and a mechanistic study of post-vaccination immune responses. *The Lancet Infectious Diseases*. Published online November 2021:S1473309921006800. doi:10.1016/S1473-3099(21)00680-0
- 41. Arbel R, Hammerman A, Sergienko R, et al. BNT162b2 Vaccine Booster and Mortality Due to Covid-19. *New England Journal of Medicine*. Published online December 8, 2021. doi:10.1056/NEJMoa2115624
- 42. Goldberg Y, Mandel M, Bar-On YM, et al. *Protection and Waning of Natural and Hybrid COVID-19 Immunity*. Epidemiology; 2021. doi:10.1101/2021.12.04.21267114
- 43. Emary KRW, Golubchik T, Aley PK, et al. Efficacy of ChAdOx1 nCoV-19 (AZD1222) vaccine against SARS-CoV-2 variant of concern 202012/01 (B.1.1.7): an exploratory analysis of a randomised controlled trial. *The Lancet*. 2021;397(10282):1351-1362. doi:10.1016/S0140-6736(21)00628-0
- 44. Heath PT, Eva Galiza FP, David Neil Baxter M, et al. Efficacy of the NVX-CoV2373 Covid-19 Vaccine Against the B.1.1.7 Variant. *medRxiv*. Published online May 2021:2021.05.13.21256639-2021.05.13.21256639. doi:10.1101/2021.05.13.21256639
- 45. Madhi SA, Baillie V, Cutland CL, et al. Efficacy of the ChAdOx1 nCoV-19 Covid-19 Vaccine against the B.1.351 Variant. *New England Journal of Medicine*. Published online March 2021:NEJMoa2102214-NEJMoa2102214. doi:10.1056/NEJMoa2102214

- 46. Sadoff J, Gray G, Vandebosch A, et al. Safety and Efficacy of Single-Dose Ad26.COV2.S Vaccine against Covid-19. *New England Journal of Medicine*. Published online April 2021:NEJMoa2101544-NEJMoa2101544. doi:10.1056/NEJMoa2101544
- 47. Shinde V, Bhikha S, Hoosain MZ, et al. Preliminary Efficacy of the NVX-CoV2373 Covid-19 Vaccine Against the B.1.351 Variant [Authors, highest degree, and affiliation/institution]. *medRxiv*. Published online March 2021:2021.02.25.21252477-2021.02.25.21252477. doi:10.1101/2021.02.25.21252477
- 48. Thomas SJ, Moreira ED, Kitchin N, et al. Six Month Safety and Efficacy of the BNT162b2 mRNA COVID-19 Vaccine. *medRxiv*. Published online July 28, 2021:2021.07.28.21261159. doi:10.1101/2021.07.28.21261159
- 49. Ella R, Reddy S, Blackwelder W, et al. Efficacy, safety, and lot to lot immunogenicity of an inactivated SARS-CoV-2 vaccine (BBV152): a double-blind, randomised, controlled phase 3 trial. *medRxiv*. Published online July 2, 2021:2021.06.30.21259439. doi:10.1101/2021.06.30.21259439



## **COVID-19 Weekly Epidemiological Update**

#### Edition 71, published 21 December 2021

#### In this edition:

- Global overview
- Special focus: Update on SARS-CoV-2 Variants of Interest and Variants of Concern
- WHO regional overviews
- Summary of the Weekly Operational Update

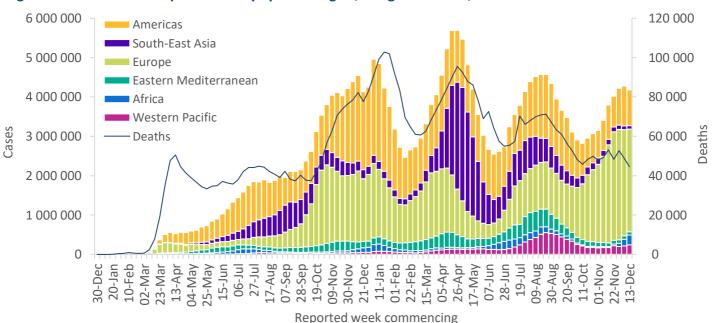
#### **Global overview**

#### Data as of 19 December 2021

During the week 13-19 December, the global number of new cases remained similar to those reported during the previous week (Table 1); however, the weekly incidence of deaths decreased by 9%. Nonetheless, this corresponds to over 4.1 million new cases and just under 45 000 new deaths. As of 19 December, over 273 million cases and over 5.3 million deaths have been reported globally (Figure 1).

The African Region continued to report the largest increase in new cases in the last week (53%), followed by the Western Pacific Region, which reported an increase of 12%. The South-East Asia and the Eastern Mediterranean regions both reported decreases of 12% and the Region of the Americas reported a 10% decrease. The number of new weekly cases reported by the European Region was similar to the numbers reported in the previous week. The African Region was the only region to report an increase in the number of new weekly deaths (15%). The Region of the Americas reported the largest decrease (15%), followed by the Eastern Mediterranean Region (12%), the European Region (7%) and the Western Pacific and South-East Asia Regions (both 6%).

Figure 1. COVID-19 cases reported weekly by WHO Region, and global deaths, as of 19 December 2021\*\*



<sup>\*\*</sup>See Annex 2: Data, table, and figure notes

The European Region continued to report the highest incidence of weekly cases (279.9 new cases per 100 000 population), followed by the Region of the Americas (88.5 new cases per 100 000 population). Both regions also reported the highest weekly incidence in deaths of 2.9 and 1.0 per 100 000 population, respectively, while all other regions reported <1 new death per 100 000.

The highest numbers of new cases were reported from the United States of America (725 750 new cases; 12% decrease), the United Kingdom (507 984 new cases; 45% increase), France (358 175 new cases; 7% increase) and Germany (283 673 new cases; 19% decrease).

Please note, the next two issues of the Weekly Epidemiological Report (to be published on 28 December and 6 January) will be condensed versions covering only the global and regional epidemiology.

Table 1. Newly reported and cumulative COVID-19 confirmed cases and deaths, by WHO Region, as of 19 December 2021\*\*

WHO Region	New cases in last 7 days (%)	Change in new cases in last 7 days *	Cumulative cases (%)	New deaths in last 7 days (%)	Change in new deaths in last 7 days *	Cumulative deaths (%)
Europe	2 611 478 (63%)	-3%	94 345 936 (35%)	26 802 (60%)	-7%	1 626 294 (30%)
Americas	904 789 (22%)	-10%	99 606 828 (36%)	10 255 (23%)	-15%	2 384 550 (45%)
Africa	256 031 (6%)	53%	6 778 548 (2%)	564 (1%)	15%	154 330 (3%)
Western Pacific	239 159 (6%)	12%	10 823 510 (4%)	3 144 (7%)	-6%	150 683 (3%)
South-East Asia	86 545 (2%)	-12%	44 823 551 (16%)	2 475 (6%)	-6%	716 778 (13%)
Eastern Mediterranean	79 620 (2%)	-12%	17 016 594 (6%)	1 376 (3%)	-12%	313 674 (6%)
Global	4 177 622 (100%)	-2%	273 395 731 (100%)	44 616 (100%)	-9%	5 346 322 (100%)

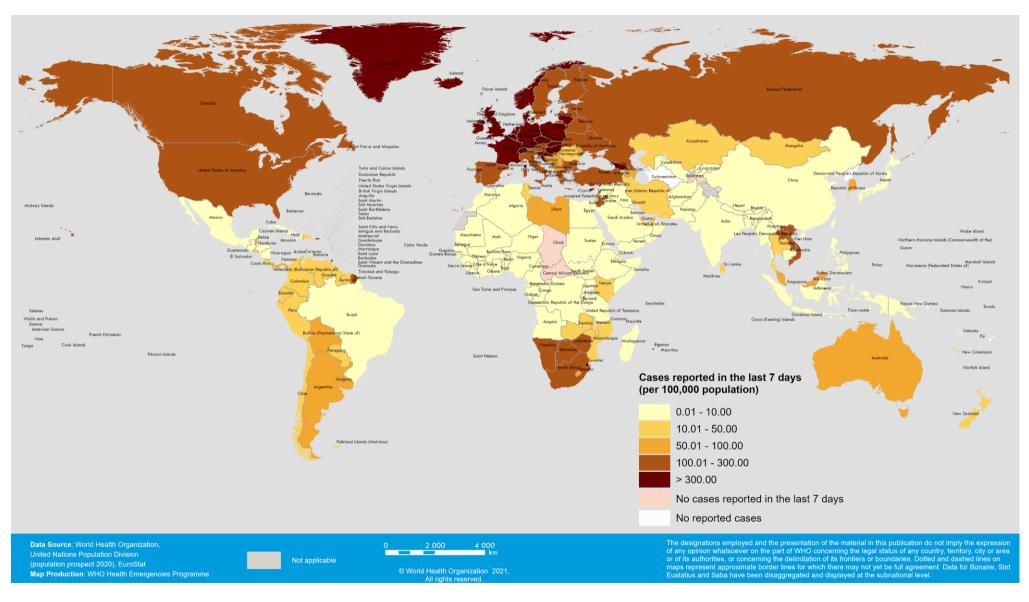
<sup>\*</sup>Percent change in the number of newly confirmed cases/deaths in the past seven days, compared to seven days prior

For the latest data and other updates on COVID-19, please see:

- WHO COVID-19 Dashboard
- WHO COVID-19 Weekly Operational Update and previous editions of the Weekly Epidemiological Update

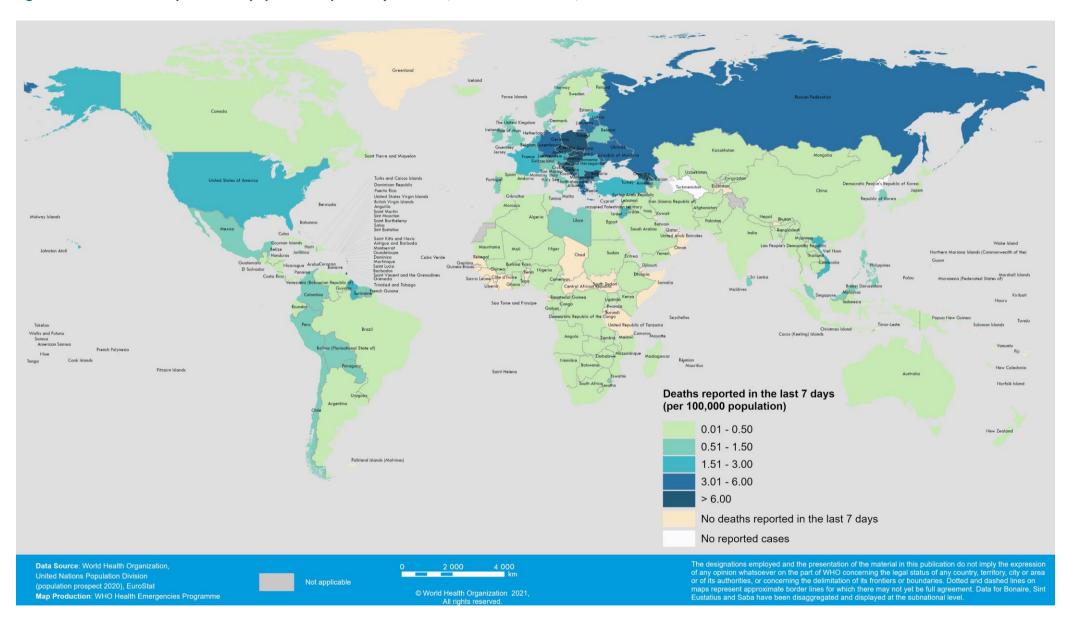
<sup>\*\*</sup>See Annex 2: Data, table, and figure notes

Figure 2. COVID-19 cases per 100 000 population reported by countries, territories and areas, 13 December - 19 December 2021\*\*



<sup>\*\*</sup>See Annex 2: Data, table, and figure notes

Figure 3. COVID-19 deaths per 100 000 population reported by countries, territories and areas, 13 December - 19 December 2021\*\*



<sup>\*\*</sup>See Annex 2: Data, table, and figure notes

#### Special Focus: Update on SARS-CoV-2 variants of interest and variants of concern

WHO, in collaboration with national authorities, institutions and researchers, routinely assesses if variants of SARS-CoV-2 alter transmission or disease characteristics, or impact effectiveness of vaccines, therapeutics, diagnostics or public health and social measures (PHSM) applied by national authorities to control disease spread. Potential variants of concern (VOCs), variants of interest (VOIs) or variants under monitoring (VUMs) are regularly assessed based on the risk posed to global public health. As evidence becomes available, classifications of variants will be revised to reflect the continuous evolution of circulating variants and their changing epidemiology. Criteria for variant classification, and the current lists of VOCs, VOIs and VUMs, are available on the WHO Tracking SARS-CoV-2 variants website. National authorities may choose to designate other variants of local interest/concern and are encouraged to investigate and report on the impacts of these variants.

#### Geographic spread and prevalence of VOCs

The current global epidemiology of SARS-CoV-2 is characterized by a predominance of the Delta variant, a declining trend in the proportion of Alpha, Beta and Gamma variants, which have been circulating at a very low prevalence for several weeks, and the emergence of Omicron variant (Figure 4; Annex 1). Following the classification of Omicron as a VOC, many countries have adopted targeted sequencing strategies to detect the variant. The change in sampling strategy, away from community-based surveillance sequencing, may result in biases in the proportions of variants being reported. Thus, the recent declines in the proportion of Delta variant reported by some countries may reflect changes in sampling strategy, rather than declines in the proportion of Delta variant cases among all COVID-19 cases.

While most of the Omicron cases identified in November 2021 were travel-related, community transmission with associated clusters has now been reported in several countries. Of 1 051 598 sequences<sup>i</sup>, 1 009 253 (96%) were Delta, 16 988 (1.6%) were Omicron, 176 (<0.1%) were Gamma, 53 (<0.1%) were Alpha, 16 (<0.1%) were Beta and 188 (<0.1%) comprised other circulating variants (including VOIs Mu and Lambda). To note, global VOCs distribution should be interpreted with due consideration of surveillance limitations, including differences in sequencing capacities and sampling strategies between countries, as well as delays in reporting.

Since the last <u>update published on 14 December</u>, additional countries across all six WHO Regions have reported confirmed cases of the Omicron variant. As of 21 December 2021 (2 pm CET), the Omicron variant has been confirmed in 106 countries.

Recent evidence indicates that Omicron variant has a growth advantage over the Delta variant and is spreading rapidly, even in countries with documented community transmission and high levels of population immunity. It remains uncertain to what extent the observed rapid growth rate can be attributed to immune evasion, intrinsic increased transmissibility, or a combination of both. There are still limited data on the clinical severity of Omicron. Hospitalizations in the UK and South Africa continue to rise, and given rapidly increasing case numbers, it is possible that healthcare systems may become overwhelmed. Preliminary data suggest that there is a reduction in neutralization of Omicron in those who have received a primary vaccination series or in those who have had prior SARS-CoV-2 infection, which may suggest a level of humoral immune evasion.

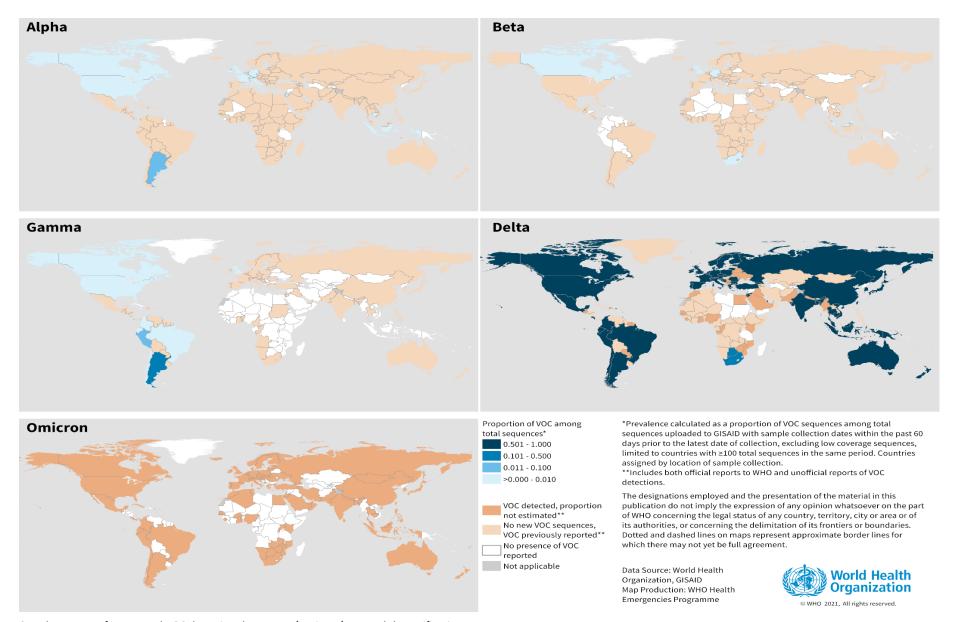
As a result of this, the overall risk related to the new variant of concern Omicron remains very high. More information on this variant can be found in the updated <u>Technical Brief and Priority Actions for Member States</u> that was published on 17 December 2021 by WHO.

<sup>&</sup>lt;sup>1</sup> Includes sequences submitted to GISAID with sample collected dates from 20 October to 19 December 2021 (last reported sample at the time of data extraction), excluding low coverage sequences.

#### **Additional resources**

- Tracking SARS-CoV-2 Variants
- COVID-19 new variants: Knowledge gaps and research
- Genomic sequencing of SARS-CoV-2: a guide to implementation for maximum impact on public health
- Considerations for implementing and adjusting PHSM in the context of COVID-19

Figure 4: Presence and prevalence of variants of concern (VOCs) in the last 60 days and historic detections, data as of 21 December 2021



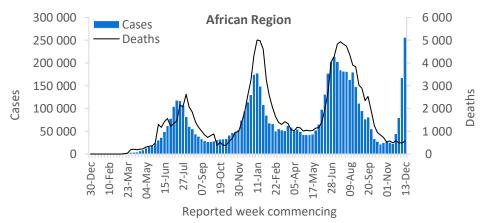
See also Annex 1 for reported VOC detections by country/territory/area and the verification status.

# WHO regional overviews Epidemiological week 13 – 19 December 2021

## **African Region**

The African Region continued to see an increase in the reported case incidence in the last month. Over 256 000 new cases were reported (an increase of 53%) as compared to the previous week, the highest number of weekly cases reported since the start of the pandemic. Increases in incidence of over 50% were observed in nearly half (23/49; 47%) of countries in the Region. The highest numbers of new cases continued to be reported from South Africa (162 987 new cases; 274.8 new cases per 100 000 population; a 50% increase) and Zimbabwe (26 671 new cases; 179.4 new cases per 100 000 population; similar to the previous week's number), with Eswatini now reporting the third highest incidence of cases (7540 new cases; 649.9 new cases per 100 000 population; a 57% increase).

The Region reported just under 500 new weekly deaths, similar to the number in the previous week. The highest numbers of new deaths were reported from South Africa (229 new deaths; <1 new death per 100 000 population; a 52% increase), Mauritius (60 new deaths; 4.7 new deaths per 100 000; a 35% decrease), and Zimbabwe (47 new deaths; <1 new death per 100 000; an 81% increase).

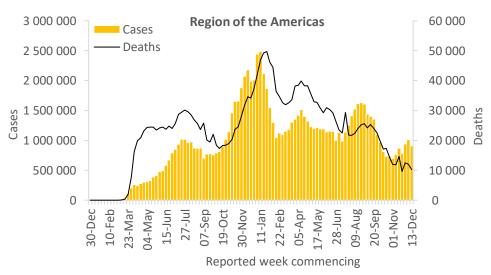


Updates from the African Region

## **Region of the Americas**

The Region of the Americas reported over 904 000 new cases and over 10 000 new deaths, decreases of 10% and 15% respectively as compared to the previous week. However, a quarter of the countries (14/56) reported increases of over 10%, with Puerto Rico reporting the highest increase (3736 new cases, a 425% increase). The highest numbers of new cases were reported from the United States of America (725 750 new cases; 219.3 new cases per 100 000; a 12% decrease), Canada (39 216 new cases; 103.9 new cases per 100 000; a 55% increase), and Argentina (32 013 new cases; 70.8 new cases per 100 000; a 91% increase).

The highest numbers of new deaths were reported from the United States of America (6723 new deaths; 2.0 new deaths per 100 000; a 20% decrease), Mexico (892 new deaths; <1 new death per 100 000; a 49% increase), and Brazil (704 new deaths; <1 new death per 100 000; a 45% decrease).

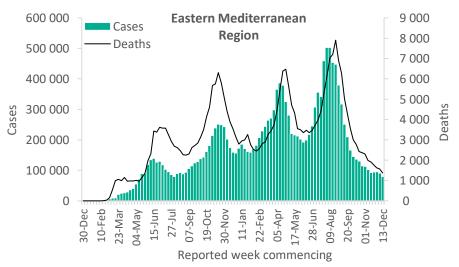


Updates from the Region of the Americas

## **Eastern Mediterranean Region**

The weekly incidence of cases and deaths in the Eastern Mediterranean Region both decreased by 12%, with over 79 000 new cases and over 1300 new deaths reported. However, the percentage of countries reporting increases of over 10% rose from 13% to 45% (10/22 countries), with the highest increases reported from Somalia (95 vs 23 new cases, a 313% increase); the United Arab Emirates (1133 vs 441 new cases, a 157% increase) and Djibouti, although case numbers remain low (14 vs 6 new cases; a 133% increase). The highest numbers of new cases were reported from Jordan (27 333 new cases; 267.9 new cases per 100 000; an 21% decrease), the Islamic Republic of Iran (16 487 new cases; 19.6 new cases per 100 000; a 22% decrease), and Lebanon (10 949 new cases; 160.4 new cases per 100 000; a 3% decrease).

The highest numbers of new deaths continued to be reported from the Islamic Republic of Iran (372 new deaths; <1 new death per 100 000; a 31% decrease), Egypt (300 new deaths; <1 new death per 100 000; a 10% decrease), and Jordan (250 new deaths; 2.5 new deaths per 100 000; an 11% increase).

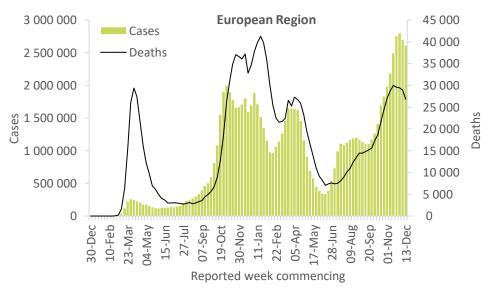


Updates from the Eastern Mediterranean Region

## **European Region**

The European Region reported over 2.6 million new cases, similar to the previous week's number. However, the weekly incidence of deaths decreased by 7%, with over 26 000 new deaths reported. Despite the stable trend, 20% of countries (12/61) still reported an increase of over 10% in cases as compared to the previous week, with the highest increases reported from Malta (1103 vs 603 new cases, an 83% increase) and Gibraltar (270 vs 148 new cases, an 82% increase). The highest numbers of new cases were reported from the United Kingdom (507 984 new cases; 748.3 new cases per 100 000; a 45% increase), France (358 175 new cases; 550.7 new cases per 100 000; a 7% increase) and Germany (283 673 new cases; 341.1 new cases per 100 000; a 19% decrease).

The highest numbers of new deaths were reported from the Russian Federation (7720 new deaths; 5.3 new deaths per 100 000; a 6% decrease), Poland (3006 new deaths; 7.9 new deaths per 100 000; a 7% increase), and Germany (2595 new deaths; 3.1 new deaths per 100 000; similar to the previous week).

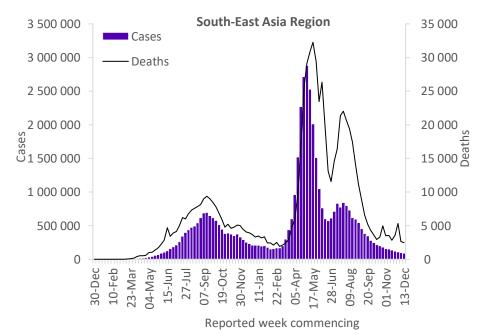


Updates from the European Region

## **South-East Asia Region**

The declining trend in the number of weekly cases and deaths reported has continued in the South-East Asia Region. Over 86 000 new cases and over 2400 new deaths were reported, decreases of 12% and 6%, respectively as compared to the previous week. Only two countries reported an increase in weekly cases, Timor Leste (4 vs 1 new case, a 300% increase) and Sri Lanka. The highest numbers of new cases continued to be reported from India (49 765 new cases; 3.6 new cases per 100 000; a 13% decrease), Thailand (22 882 new cases; 32.8 new cases per 100 000; a 17% decrease), and Sri Lanka (6783 new cases; 31.7 new cases per 100 000; a 30% increase).

The highest numbers of new deaths also continued to be reported from India (1988 new deaths; <1 new death per 100 000; a 6% decrease), Thailand (206 new deaths; <1 new death per 100 000; a 9% decrease), and Sri Lanka (138 new deaths; <1 new death per 100 000; a 10% decrease).

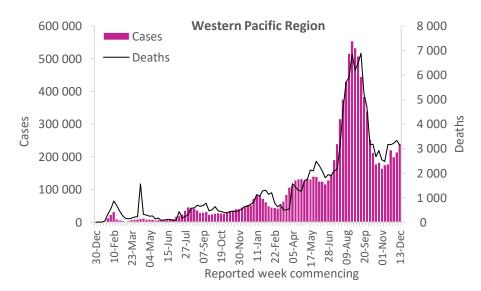


Updates from the **South-East Asia Region** 

## **Western Pacific Region**

The Western Pacific Region continues to see an increase in the incidence of cases with 239 000 new cases reported during the previous week (a 12% increase). Three of the 27 countries in the region, reported an increase in case incidence of over 10% including: Australia (19 415 vs 10 805 new cases, a 80% increase), Japan (1088 vs 861 new cases, a 26% increase) and Viet Nam. The highest numbers of new cases were reported from Viet Nam (125 955 new cases; 129.4 new cases per 100 000; a 22% increase), the Republic of Korea (47 825 new cases; 93.3 new cases per 100 000; an 8% increase), and Malaysia (27 698 new cases; 85.6 new cases per 100 000; an 18% decrease).

The incidence of deaths decreased by 6% as compared to the previous week, with over 3100 new deaths reported. The highest numbers of new deaths continued to be reported from Viet Nam (1740 new deaths; 1.8 new deaths per 100 000; a 12% increase), the Philippines (579 new deaths; <1 new death per 100 000; a 33% decrease), and the Republic of Korea (469 new deaths; <1 new death per 100 000; a 17% increase).



Updates from the Western Pacific Region

## **Summary of the COVID-19 Weekly Operational Update**

The <u>Weekly Operational Update</u> is a report provided by the COVID-19 Strategic Preparedness and Response Plan (SPRP) Monitoring and Evaluation team, which aims to update on the ongoing global progress against the <u>COVID-19 SPRP 2021</u> framework, and to highlight country-level actions and WHO support to countries. In this week's edition published on 21 December, highlights include the following:

- Implementing national studies on the real-world effectiveness of COVID-19 vaccines in Armenia
- Delivering critical equipment and supplies in the Western Pacific to Vanuatu and Fiji
- Boosting Sierra Leone's COVID-19 response and disease surveillance with laboratory commodities
- Bringing COVID-19 vaccination to the most vulnerable via the Global Health Cluster
- Supporting the global scale-up of infodemic management
- Strengthening genomic surveillance: WHO in collaboration with GISAID organizes training workshops for laboratory experts
- Conducting the first Universal Health and Preparedness Review (UHPR) Pilot: Bangui, Central African Republic
- Updates on WHO's financing to support countries on COVID-19 response implementation to suppress transmission, reduce exposure, and protect the vulnerable and save lives
- Progress on a subset of global indicators that demonstrate country and global progress to end the acute phase of the pandemic

## **Technical guidance and other resources**

- WHO technical guidance
- WHO COVID-19 Dashboard
- WHO Weekly Operational Updates on COVID-19
- WHO COVID-19 case definitions
- COVID-19 Supply Chain Inter-Agency Coordination Cell Weekly Situational Update
- Research and Development
- Open WHO courses on COVID-19 in official UN languages and in additional national languages
- WHO Academy COVID-19 mobile learning app
- <u>The Strategic Preparedness and Response Plan</u> (SPRP) outlining the support the international community can provide to all countries to prepare and respond to the virus
- EPI-WIN: tailored information for individuals, organizations, and communities
- Recommendations and advice for the public:
  - Protect yourself
  - Questions and answers
  - <u>Travel advice</u>

## **Annexes**

Annex 1. List of countries/territories/areas reporting variants of concern as of 21 December 2021

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Afghanistan	•	-	•	-	-
Albania	•	-	0	-	-
Algeria	•	-	•	-	•*
Andorra	0	0	0	-	-
Angola	•	•	•	•	-
Anguilla	•	-	•	-	-
Antigua and Barbuda	•	•	•	•	-
Argentina	•	•	•	•	•
Armenia	•	-	•	-	-
Aruba	•	•	•	•	-
Australia	•	•	•	•	•
Austria	•	•	•	•	•
Azerbaijan	•	-	0	-	-
Bahamas	•	-	•	•	-
Bahrain	•	•	•	•	•
Bangladesh	•	•	•	0	•
Barbados	•	-	•	•	-
Belarus	•	-	0	-	-
Belgium	•	•	•	•	•
Belize	•	-	•	•	-
Benin	•	•	•	•	-
Bermuda	•	•	•	-	•
Bhutan	•	•	•	-	-
Bolivia (Plurinational State of)	•	-	•	•	-
Bonaire	•	-	•	•	-
Bosnia and Herzegovina	•	•	0	•	-
Botswana	0	•	•	-	•
Brazil	•	•	•	•	•

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
British Virgin Islands	•	-	•	•	-
Brunei Darussalam	•	•	•	-	-
Bulgaria	•	•	•	-	-
Burkina Faso	•	-	•	-	•*
Burundi	•	•	•	-	-
Cabo Verde	•	-	•	-	-
Cambodia	•	•	•	-	•*
Cameroon	•	•	•	•	-
Canada	•	•	•	•	•
Cayman Islands	•	•	•	•	•*
Central African Republic	•	•	•	-	-
Chad	•	-	-	-	-
Chile	•	•	•	•	•
China	•	•	•	•	•
Colombia	•	-	•	•	0*
Comoros	-	•	•	-	-
Congo	•	•	•	•	-
Costa Rica	•	•	•	•	-
Croatia	•	•	0	•	•
Cuba	•	•	•	-	•
Curaçao	•	•	•	•	-
Cyprus	•	•	0	-	•
Czechia	•	•	•	•	•
Côte d'Ivoire	•	•	0	-	-
Democratic Republic of the Congo	•	•	•	-	-
Denmark	•	•	•	•	•
Djibouti	•	•	•	-	-

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Dominica	•	-	•	-	-
Dominican Republic	•	-	•	•	-
Ecuador	•	-	•	•	•*
Egypt	•	-	•	-	0*
El Salvador	•	-	•	•	-
Equatorial Guinea	•	•	•	-	-
Estonia	•	•	0	0	•
Eswatini	0	•	•	-	-
Ethiopia	•	•	•	-	-
Falkland Islands (Malvinas)	•	•	-	-	-
Faroe Islands	•	-	-	•	-
Fiji	0	-	•	-	•
Finland	•	•	•	•	•
France	•	•	•	•	•
French Guiana	•	•	•	•	-
French Polynesia	•	•	•	•	•*
Gabon	•	•	•	-	-
Gambia	•	-	•	-	-
Georgia	•	0	•	-	•*
Germany	•	•	•	•	•
Ghana	•	•	•	•	•
Gibraltar	•	-	0	-	•
Greece	•	•	•	•	•
Greenland	-	-	•	-	-
Grenada	•	-	•	•	-
Guadeloupe	•	•	•	•	•*
Guam	•	•	•	•	-
Guatemala	•	•	•	•	-

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Guernsey	-	-	-	-	•*
Guinea	•	•	•	-	-
Guinea-Bissau	•	•	•	-	-
Guyana	-	-	•	•	-
Haiti	•	-	•	•	-
Honduras	•	-	•	•	-
Hungary	•	0	0	•	•
Iceland	•	•	•	•	•
India	•	•	•	•	•
Indonesia	•	•	•	-	•*
Iran (Islamic Republic of)	•	•	•	-	•*
Iraq	•	•	•	•	-
Ireland	•	•	•	•	•
Israel	•	•	•	•	•
Italy	•	•	•	•	•
Jamaica	•	-	•	-	-
Japan	•	•	•	•	•
Jordan	•	•	•	•	•
Kazakhstan	•	0	•	-	-
Kenya	•	•	•	-	0*
Kosovo[1]	•	0	0	-	-
Kuwait	•	•	•	-	•
Kyrgyzstan	•	•	•	-	-
Lao People's Democratic Republic	•	-	•	-	-
Latvia	•	•	0	•	•
Lebanon	•	-	•	-	•
Lesotho	-	•	0	-	-
Liberia	•	•	•	-	-
Libya	•	•	-	-	-
Liechtenstein	•	-	0	0	0
Lithuania	•	•	0	•	•*

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron	Country/T
Luxembourg	•	•	•	•	•	Oman
Madagascar	•	•	-	-	-	Pakistan
Malawi	•	•	•	-	•	Panama
Malaysia	•	•	•	-	•	Papua Nev
Maldives	•	-	•	-	•	Paraguay
Mali	•*	-	•	-	-	Peru
Malta	•	0	0	•	-	Philippines
Martinique	•	•	•	•	-	Poland
Mauritania	•	•	•	-	-	Portugal
Mauritius	•	•	•	-	0	Puerto Ric
Mayotte	•	•	0	-	-	Qatar
Mexico	•	•	•	•	•	Republic o
Monaco	•	•	•	-	-	Republic o
Mongolia	•	-	•	-	0*	Romania
Montenegro	•	-	0	0	0	Russian Fe
Montserrat	•	-	•	•	-	Rwanda
Morocco	•	•	•	-	•*	Réunion
Mozambique	•	•	•	-	•*	Saba
Myanmar	•	-	•	-	-	Saint Barth
Namibia	•	•	•	•	•	Saint Kitts
Nepal	•	-	•	-	•	Saint Lucia
Netherlands	•	•	•	•	•	Saint Mart
New Caledonia	•	-	•	-	-	Saint Pierr
New Zealand	•	•	•	•	•*	Saint Vince
Nicaragua	•	•	•	•	-	Grenadine
Niger	•	-	•	-	-	Sao Tome
Nigeria	•	•	•	-	•	Saudi Arab
North Macedonia	•	•	0	-	-	Senegal
Northern Mariana Islands	0	_	•	_	_	Serbia
(Commonwealth of the)						Seychelles
Norway	•	•	•	•	•	Sierra Leor
Occupied Palestinian Territory	•	•	•	-	-	Singapore

Gamma	Omicron	Count	Country/Territory/Area				Delta	Gamma	Omicron
•	•	Omar	1	•	•	•	-	•*	
-	-	Pakist	an		•	•	•	•	•
-	•	Panar	ma		•	•	•	•	•*
-	•	Papua	New Guinea		-	-	•	-	-
-	•	Parag	uay		•	-	•	•	-
-	-	Peru			•	-	•	•	•*
•	-	Philip	pines		•	•	•	•	•*
•	-	Polan	d		•	0	•	•	•*
-	-	Portu	gal		•	•	•	•	•
-	0	Puert	o Rico		•	•	•	•	•
-	-	Qatar	Qatar				•	-	-
•	•	Repul	Republic of Korea			•	•	•	•
-	-	Repul	Republic of Moldova		•	-	•	-	-
	0*	Roma	Romania		•	•	•	•	•
0	0	Russia	Russian Federation		•	•	•	0	•
•	-	Rwan	Rwanda		•	•	•	-	0*
-	•*	Réuni	Réunion		•	•	0	•	•
-	•*	Saba			-	-	•	-	-
-	-	Saint	Barthélemy		•	-	•	-	-
•	•	Saint	Kitts and Nevis		-	-	•	-	-
	•	Saint	Lucia		•	-	•	-	-
•	•	Saint	Martin		•	•	•	-	•*
-	-	Saint	Pierre and Mique	lon	-	-	•	-	-
•	•*		Vincent and the adines		-	-	•	•	-
-	_	Sao To	ome and Principe	!	•	-	0	-	-
	•		Arabia		•	•	•	-	•
_		Seneg	gal		•	•	•	-	•
		Serbia	 Э		•	-	•	-	-
-	-	Seych	elles		•	•	•	-	-
•	•		Leone		-	•	•	-	•
-	-	Singa	pore		•	•	•	•	•

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Sint Maarten	•	•	•	•	-
Slovakia	•	•	•	-	•
Slovenia	•	•	•	•	•*
Somalia	•	•	•	-	-
South Africa	•	•	•	0	•
South Sudan	•	•	•	-	-
Spain	•	•	•	•	•
Sri Lanka	•	•	•	-	•
Sudan	•	•	-	•	-
Suriname	•	•	•	•	-
Sweden	•	•	•	•	•
Switzerland	•	•	•	•	•
Thailand	•	•	•	•	•

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Timor-Leste	•	-	•	-	-
Togo	•	•	•	•	•*
Trinidad and Tobago	•	-	•	•	•
Tunisia	•	•	•	-	•
Turkey	•	•	•	•	0
Turks and Caicos Islands	•	-	•	•	-
Uganda	•	•	•	-	•
Ukraine	•	0	0	-	•*
United Arab Emirates	•	•	•	•	•
United Kingdom	•	•	•	•	•
United Republic of Tanzania	-	•	-	-	-
United States Virgin Islands	•	•	•	•	-
United States of America	•	•	•	•	•

Country/Territory/Area	Alpha	Beta	Delta	Gamma	Omicron
Uruguay	•	•	•	•	-
Uzbekistan	•	•	0	-	-
Vanuatu	-	-	•	-	-
Venezuela (Bolivarian Republic of)	•	-	•	•	-
Viet Nam	•	•	•	-	-
Wallis and Futuna	•	-	-	-	-
Yemen	•	•	-	-	-
Zambia	•	•	•	-	•
Zimbabwe	•	•	•	-	•

<sup>\*</sup>Newly reported in this update. "•" indicates that information for this variant was received by WHO from official sources. "o" indicates that information for this variant was received by WHO from unofficial sources and will be reviewed as more information become available. \*\*Includes countries/territories/areas reporting the detection of VOCs among travellers (e.g., imported cases detected at points of entry), or local cases (detected in the community). Excludes countries, territories, and areas that have never reported the detection of a variant of concern. See also Annex 2: Data, table, and figure notes

#### Annex 2. Data, table, and figure notes

Data presented are based on official laboratory-confirmed COVID-19 case and deaths reported to WHO by country/territories/areas, largely based upon WHO <u>case definitions</u> and <u>surveillance guidance</u>. While steps are taken to ensure accuracy and reliability, all data are subject to continuous verification and change, and caution must be taken when interpreting these data as several factors influence the counts presented, with variable underestimation of true case and death incidences, and variable delays to reflecting these data at the global level. Case detection, inclusion criteria, testing strategies, reporting practices, and data cut-off and lag times differ between countries/territories/areas. A small number of countries/territories/areas report combined probable and laboratory-confirmed cases. Differences are to be expected between information products published by WHO, national public health authorities, and other sources.

Due to public health authorities conducting data reconciliation exercises that remove large numbers of cases or deaths from their total counts, negative numbers may be displayed in the new cases/deaths columns as appropriate. When additional details become available that allow the subtractions to be suitably apportioned to previous days, graphics will be updated accordingly. A record of historic data adjustment made is available upon request by emailing <a href="mailto:epi-data-support@who.int">epi-data-support@who.int</a>. Please specify the countries of interest, time period, and purpose of the request/intended usage. Prior situation reports will not be edited; see <a href="mailto:covid19.who.int">covid19.who.int</a> for the most up-to-date data. COVID-19 confirmed cases and deaths reported in the last seven days by countries, territories, and areas, and WHO Region (reported in previous issues) are now available at: <a href="https://covid19.who.int/table">https://covid19.who.int/table</a>.

'Countries' may refer to countries, territories, areas or other jurisdictions of similar status. The designations employed, and the presentation of these materials do not imply the expression of any opinion whatsoever on the part of WHO concerning the legal status of any country, territory, or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted and dashed lines on maps represent approximate border lines for which there may not yet be full agreement. Countries, territories, and areas are arranged under the administering WHO region. The mention of specific companies or of certain manufacturers' products does not imply that they are endorsed or recommended by WHO in preference to others of a similar nature that are not mentioned. Errors and omissions except, the names of proprietary products are distinguished by initial capital letters.

[1] All references to Kosovo should be understood to be in the context of the United Nations Security Council resolution 1244 (1999). In the map, the number of cases of Serbia and Kosovo (UNSCR 1244, 1999) have been aggregated for visualization purposes.



## **COVID-19 Weekly Epidemiological Update**

#### Edition 72, published 28 December 2021

#### In this edition:

- Global overview
- Brief Update on SARS-CoV-2 Omicron variant
- WHO regional overviews

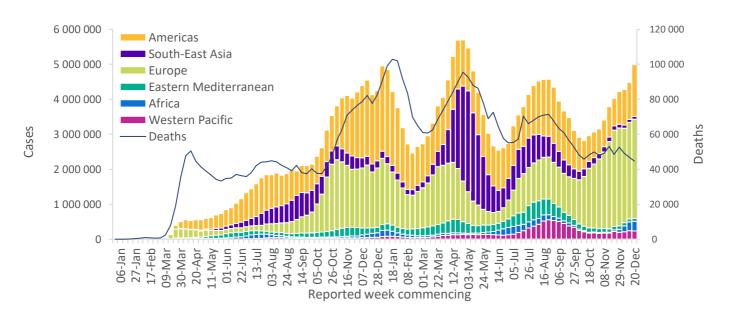
#### Global overview

#### Data as of 26 December 2021

During the week 20-26 December, following a gradual increase since October, the global number of new cases increased by 11% as compared to the previous week (Table 1); while the number of new deaths remained similar to the number reported during the previous week. This corresponds to just under 5 million new cases and over 44 000 new deaths. As of 26 December, over 278 million cases and just under 5.4 million deaths have been reported globally (Figure 1).

The Region of the Americas reported the largest increase in new cases in the last week (39%), followed by the African Region, which reported an increase of 7%. The South-East Asia Region continued to report a decrease in new cases as compared to the previous week (12%) while in the European, Eastern Mediterranean, and Western Pacific Regions, the number of new cases was similar to those reported during the previous week. The African Region reported the highest increase in the number of new deaths (72%), followed by the South-East Asia Region (9%) and the Region of the Americas (7%). The European and Eastern Mediterranean Region reported decreases of 12% and 7% respectively, in the incidence of deaths, while in the Western Pacific Region, the incidence was similar to the previous weeks.

Figure 1. COVID-19 cases reported weekly by WHO Region, and global deaths, as of 26 December 2021\*\*



<sup>\*\*</sup>See Annex 1: Data, table, and figure notes

The European Region continued to report the highest incidence of weekly cases (304.6 new cases per 100 000 population), followed by the Region of the Americas (144.4 new cases per 100 000 population). Both regions also reported the highest weekly incidence in deaths of 2.6 and 1.2 per 100 000 population, respectively, while all other regions reported <1 new death per 100 000.

The highest numbers of new cases were reported from the United States of America (1 185 653 new cases; 34% increase), the United Kingdom (611 864 new cases; 20% increase), France (504 642 new cases; 41% increase); Italy (257 579 new cases; 62% increase) and Germany (197 845 new cases; 30% decrease).

Please note, the next issue of the Weekly Epidemiological Update (to be published on 6 January) will be condensed versions covering only the global and regional epidemiology of COVID-19.

Table 1. Newly reported and cumulative COVID-19 confirmed cases and deaths, by WHO Region, as of 26 December 2021\*\*

WHO Region	New cases in last 7 days (%)	Change in new cases in last 7 days *	Cumulative cases (%)	New deaths in last 7 days (%)	Change in new deaths in last 7 days *	Cumulative deaths (%)
Europe	2 842 375 (57%)	3%	97 359 631 (35%)	23 900 (53%)	-12%	1 650 729 (31%)
Americas	1 476 724 (30%)	39%	101 243 155 (36%)	12 782 (29%)	7%	2 399 735 (44%)
Africa	274 342 (6%)	7%	7 055 628 (3%)	952 (2%)	72%	155 292 (3%)
Western Pacific	238 654 (5%)	0%	11 062 163 (4%)	3 063 (7%)	-3%	153 746 (3%)
Eastern Mediterranean	76 875 (2%)	-3%	17 093 469 (6%)	1 275 (3%)	-7%	314 949 (6%)
South-East Asia	76 123 (2%)	-12%	44 899 674 (16%)	2 708 (6%)	9%	719 486 (13%)
Global	4 985 093 (100%)	11%	278 714 484 (100%)	44 680 (100%)	-4%	5 393 950 (100%)

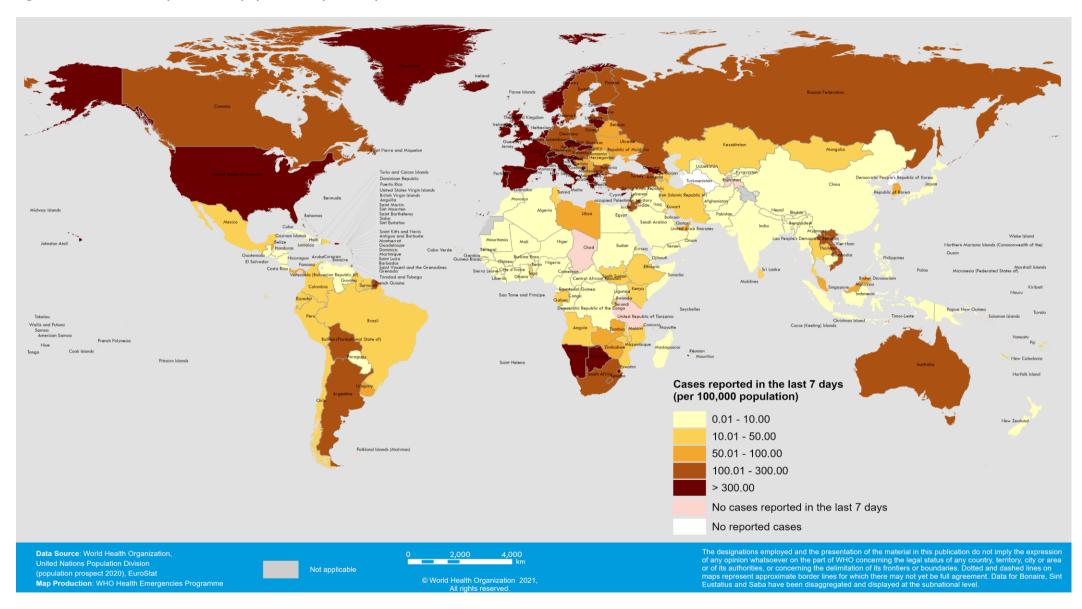
<sup>\*</sup>Percent change in the number of newly confirmed cases/deaths in the past seven days, compared to seven days prior

For the latest data and other updates on COVID-19, please see:

- WHO COVID-19 Dashboard
- WHO COVID-19 Weekly Operational Update and previous editions of the Weekly Epidemiological Update

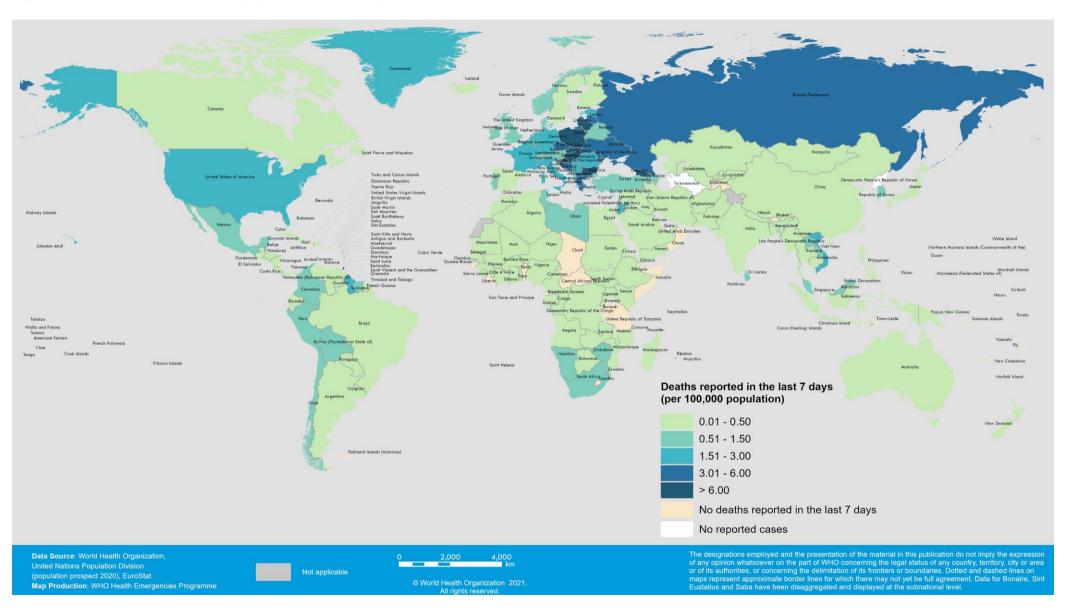
<sup>\*\*</sup>See Annex 1: Data, table, and figure notes

Figure 2. COVID-19 cases per 100 000 population reported by countries, territories and areas, 20 - 26 December 2021\*\*



<sup>\*\*</sup>See Annex 1: Data, table, and figure notes

Figure 3. COVID-19 deaths per 100 000 population reported by countries, territories and areas, 20 - 26 December 2021\*\*



<sup>\*\*</sup>See Annex 1: Data, table, and figure notes

## **Brief Update on SARS-CoV-2 Omicron variant**

The overall risk related to the new variant of concern Omicron remains very high. Consistent evidence shows that the Omicron variant has a growth advantage over the Delta variant with a doubling time of 2-3 days and rapid increases in the incidence of cases is seen in a number of countries, including those where the variant has become the dominant SARS-CoV-2 variant, such as the United Kingdom and the United States of America. However, a decline in the incidence of cases has now been observed in South Africa. The rapid growth rate is likely to be a combination of both immune evasion and intrinsic increased transmissibility of the Omicron variant. Early data from the United Kingdom, South Africa and Denmark suggests there is a reduced risk of hospitalization for the Omicron compared to the Delta variant, however, further data are needed to understand the clinical markers of severity including the use of oxygen, mechanical ventilation and death, and how severity may be impacted by vaccination and/or prior SARS-CoV-2 infection. It is also expected that corticosteroids and interleukin 6 receptor blockers will remain effective in the management of patients with severe disease, however, preliminary data suggest that monoclonal antibodies may be less able to neutralize the Omicron variant. Reassuringly, preliminary data suggests testing using either PCR or antigen-based rapid diagnostic tests (Ag-RDT) assays does not appear to be impacted by the Omicron variant. More information on this variant can be found in the updated Technical Brief and Priority Actions for Member States that was published on 23 December 2021 by WHO.

#### **Additional resources**

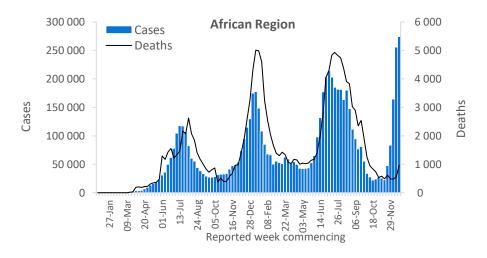
- Tracking SARS-CoV-2 Variants
- COVID-19 new variants: Knowledge gaps and research
- Genomic sequencing of SARS-CoV-2: a guide to implementation for maximum impact on public health
- Considerations for implementing and adjusting PHSM in the context of COVID-19

# WHO regional overviews Epidemiological week 20 – 26 December 2021

## **African Region**

The African Region reported over 274 000 new cases, however, the weekly increase in incidence was smaller (7%) as compared to the incidence of the previous week (53%). Increases in case incidence of over 50% were observed in nearly two-thirds (32/49; 65%) of countries in the Region. The highest numbers of new cases were reported from South Africa (115 328 new cases; 194.5 new cases per 100 000 population; a 29% decrease), Ethiopia (19 940 new cases; 17.3 new cases per 100 000 population; a 610% increase) and Kenya (19 023 new cases; 35.4 new cases per 100 000; a 207% increase).

The Region reported over 900 new weekly deaths, a 72% increase as compared to the number reported during the previous week. The highest numbers of new deaths were reported from South Africa (428 new deaths; <1 new death per 100 000 population; an 87% increase), Zimbabwe (103 new deaths; <1 new death per 100 000; a 119% increase) and the Democratic Republic of the Congo (79 new deaths; <1 new death per 100 000; an 888% increase).

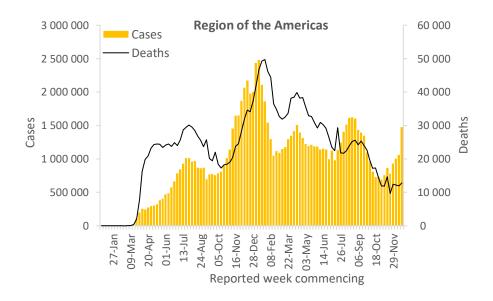


Updates from the African Region

## **Region of the Americas**

The Region of the Americas reported over 1.4 million new cases and over 12 000 new deaths, increases of 39% and 7% respectively, as compared to the previous week. Half of the countries (28/56) reported increases of over 10%, with the highest increases reported from Saint Kitts and Nevis (38 vs 3 new cases, an 1167% increase albeit small numbers); the United States Virgin Islands (367 vs 42 new cases, a 774% increase) and Puerto Rico (32 162 vs 3736 new cases, a 761% increase). However, the highest numbers of new cases continued to be reported from the United States of America (1 185 653 new cases; 358.2 new cases per 100 000; a 34% increase), Canada (78 847 new cases; 208.9 new cases per 100 000; a 101% increase), and Argentina (65 966 new cases; 146.0 new cases per 100 000; a 106% increase).

The highest numbers of new deaths were reported from the United States of America (9355 new deaths; 2.8 new deaths per 100 000; a 7% increase), Brazil (997 new deaths; <1 new death per 100 000; a 42% increase), and Mexico (797 new deaths; <1 new death per 100 000; a 41% increase).

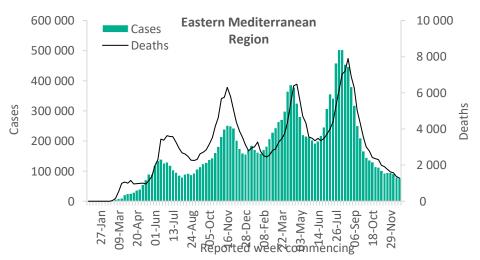


Updates from the Region of the Americas

## **Eastern Mediterranean Region**

The weekly incidence of cases in the Eastern Mediterranean Region remained similar to the incidence reported during the previous week, with over 76 000 new cases reported. Half of the countries (11/22) reported increases in cases of over 10%, with the highest increases reported from the United Arab Emirates (5678 vs 1133 new cases, a 401% increase); Somalia (363 vs 95 new cases, a 282% increase) and Saudi Arabia (1668 vs 549 new cases, 204% increase). The highest numbers of new cases continued to be reported from Jordan (17 952 new cases; 267.9 new cases per 100 000; a 34% decrease), the Islamic Republic of Iran (13 894 new cases; 16.5 new cases per 100 000; a 16% decrease), and Lebanon (11 795 new cases; 172.8 new cases per 100 000; an 8% increase).

The weekly incidence of deaths in the Region decreased by 7%, while the highest numbers of new deaths continued to be reported from the Islamic Republic of Iran (315 new deaths; <1 new death per 100 000; a 15% decrease), Egypt (256 new deaths; <1 new death per 100 000; a 15% decrease), and Jordan (253 new deaths; 2.5 new deaths per 100 000; similar to the previous week).

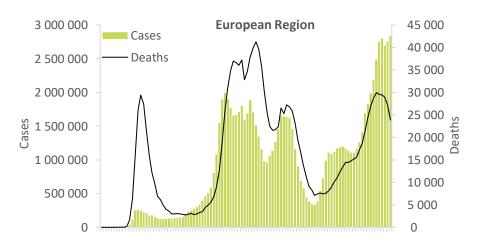


Updates from the Eastern Mediterranean Region

## **European Region**

The European Region reported over 2.8 million new cases, similar to the number reported during the previous week. However, the weekly incidence of deaths decreased by 12%, with over 24 000 new deaths reported. Despite the stable trend, one third of countries (20/61) reported a weekly increase in cases of over 10%, with the highest increases reported from Malta (4107 vs 1103 new cases, an 272% increase), Israel (9076 vs 4886 new cases, an 86% increase) and Portugal (55 217 vs 30427 new cases, an 81% increase). The highest numbers of new cases were reported from the United Kingdom (611 864 new cases; 901.3 new cases per 100 000; a 20% increase), France (504 642 new cases; 775.9 new cases per 100 000; a 41% increase) and Italy (257 579 new cases; 431.9 new cases per 100 000; a 62% increase).

The highest numbers of new deaths continued to be reported from the Russian Federation (7015 new deaths; 4.8 new deaths per 100 000; a 9% decrease), Poland (2842 new deaths; 7.5 new deaths per 100 000; a 5% decrease), and Germany (2131 new deaths; 2.6 new deaths per 100 000; an 18% decrease).



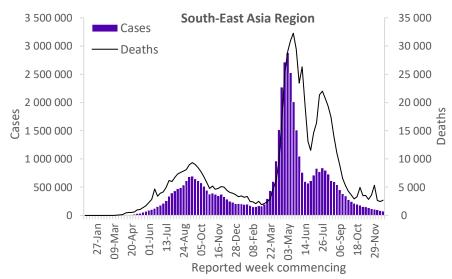
Reported week commencing

Updates from the European Region

## **South-East Asia Region**

The declining trend in the number of weekly cases reported has continued in the South-East Asia Region, with over 76 000 new cases reported, a 12% decrease as compared to the number reported during the previous week. However, the number of new weekly deaths increased by 9%, with 2700 new deaths reported. Only two countries in the Region reported an increase in weekly cases: Bangladesh (2170 vs 310 new cases, a 23% increase) and the Maldives (907 vs 838 new cases, an 8% increase). However, the highest numbers of new cases continued to be reported from India (46 527 new cases; 3.4 new cases per 100 000; a 7% decrease), Thailand (18 442 new cases; 26.4 new cases per 100 000; a 19% decrease), and Sri Lanka (3964 new cases; 18.5 new cases per 100 000; a 42% decrease).

The highest numbers of new deaths also continued to be reported from India (2260 new deaths; <1 new death per 100 000; a 14% increase), Thailand (203 new deaths; <1 new death per 100 000; similar to the previous week), and Sri Lanka (132 new deaths; <1 new death per 100 000; similar to the previous week).

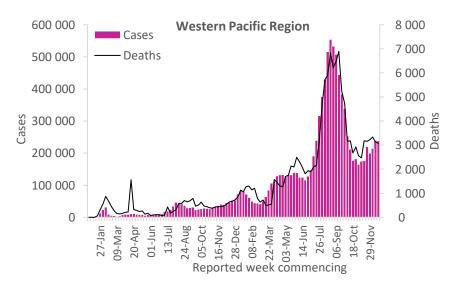


#### Updates from the South-East Asia Region

## **Western Pacific Region**

The incidence of cases and deaths reported in the Western Pacific Region was similar to those reported during the previous week, with over 238 000 new cases and over 3000 new deaths reported. However, seven of the 27 countries in the region, reported an increase in case incidence of over 10%, with the highest increases reported from French Polynesia (40 vs 8 new cases, a 400% increase), Fiji (192 vs 39 new cases, a 392% increase) and Australia. The highest numbers of new cases were reported from Viet Nam (112 087 new cases; 115.2 new cases per 100 000; a 11% decrease), Australia (45 560 new cases; 178.7 new cases per 100 000, a 135% increase), and the Republic of Korea (42 367 new cases; 82.6 new cases per 100 000; an 11% decrease).

The highest numbers of new deaths were reported from Viet Nam (1656 new deaths; 1.7 new deaths per 100 000; similar to the previous week), the Republic of Korea (523 new deaths; 1.0 new death per 100 000; a 12% increase), and the Philippines (512 new deaths; <1 new death per 100 000; a 12% decrease).



Updates from the Western Pacific Region

## **Technical guidance and other resources**

- WHO technical guidance
- WHO COVID-19 Dashboard
- WHO Weekly Operational Updates on COVID-19
- WHO COVID-19 case definitions
- COVID-19 Supply Chain Inter-Agency Coordination Cell Weekly Situational Update
- Research and Development
- Open WHO courses on COVID-19 in official UN languages and in additional national languages
- WHO Academy COVID-19 mobile learning app
- <u>The Strategic Preparedness and Response Plan</u> (SPRP) outlining the support the international community can provide to all countries to prepare and respond to the virus
- EPI-WIN: tailored information for individuals, organizations, and communities
- Recommendations and advice for the public:
  - Protect yourself
  - Questions and answers
  - Travel advice

#### Annex 1. Data, table, and figure notes

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[1] All references to Kosovo should be understood to be in the context of the United Nations Security Council resolution 1244 (1999). In the map, the number of cases of Serbia and Kosovo (UNSCR 1244, 1999) have been aggregated for visualization purposes.