This is a promotional email intended for healthcare professionals only.



WHO EXPANDS RECOMMENDATION FOR VEKLURY TO PATIENTS WITH SEVERE COVID-19¹

I'm writing with important news for the treatment of patients with COVID-19.

Dear [Doctor],

The WHO now conditionally recommends VEKLURY for patients

Non-severe COVID-19 Severe COVID-19

VEKLURY is the only antiviral treatment recommended

severe or critical COVID-19

Absence of any criteria for

with severe COVID-19.1

by the WHO for patients with:1

Signs of pneumonia

Signs of severe respiratory distress

Oxygen saturation <90% on room air

The panel did not find evidence of benefit in patients with critical COVID-19

and recommends against the use of VEKLURY in critical patients.*1

of patients with severe COVID-19 is mainly driven by the final results of the WHO-sponsored SOLIDARITY study.²

Data supporting the new recommendation

The WHO SOLIDARITY primary outcome assessed in-hospital mortality, both overall and subdivided by disease severity at entry.2

The WHO conditional recommendation for VEKLURY's use in the treatment

For patients who required supplemental oxygen at baseline,

13% 0/0

lower relative risk of death or progression to IMV vs SoC (RR: 0.83; 95% CI: 0.75–0.93)

who were already being ventilated²

standard of care in hospitals. 1,4-10

mortality for COVID-19 patients.^{1,3}

subgroups in either ACTT-1 or SOLIDARITY.^{1,3}

SOLIDARITY showed:2

of mortality vs SoC (RR: 0.87; 95% CI: 0.76–0.99) VEKLURY had no significant effect on patients with COVID-19

lower relative risk

These findings complement results from NIAID ACTT-1, a double-blind placebo-controlled trial in which a post-hoc subgroup analysis showed a mortality reduction for VEKLURY vs placebo in patients on low-flow oxygen

a mortality benefit in the overall population or other baseline oxygen

View the new WHO guideline here In addition to the WHO, VEKLURY is recommended by several major guidelines worldwide, including the NIH, NICE, ESCMID, IDSA, MHLW,

AWMF and COVRIIN, supporting VEKLURY's role as the COVID-19 antiviral

This data reflects the robust body of evidence for VEKLURY, highlighting its

ability to reduce disease progression, fast forward to recovery and reduce

at baseline (HR: 0.30; 95% CI: 0.14–0.64). VEKLURY did not demonstrate

To help your patients benefit, initiate VEKLURY right away in appropriate patients upon hospitalisation.¹¹ If you would like any additional information, I can follow up with you directly.

VEKLURY is indicated for the treatment of coronavirus disease 2019 (COVID19) in adults and paediatric

(low- or high-flow oxygen or other non-invasive ventilation at start of treatment), and adults and paediatric

patients (weighing at least 40 kg) who do not require supplemental oxygen and who are at increased risk

patients (at least 4 weeks of age and weighing at least 3 kg) with pneumonia requiring supplemental oxygen

[Insert contact details here]

of progressing to severe COVID-19. Please click here to access the Prescribing Information.

Accessed September 2022.

Best wishes,

ratio; IDSA, Infectious Diseases Society of America; MHLW, Ministry of Health, Labour and Welfare; NIAID, National Institute of Allergy and Infectious Diseases; NICE, National Institute of Health and Care Excellence; NIH, National Institutes of Health; RR, relative risk; SoC, standard of care; WHO, World Health Organisation. * Critical disease is defined as requiring life-sustaining treatment, ARDS, sepsis or septic shock.1 References:

ACTT, Adaptive COVID-19 Treatment Trial; ARDS, acute respiratory distress syndrome; AWMF, Association

of Scientific Medical Societies in Germany; CI, confidence interval; COVID-19, coronavirus disease 2019;

COVRIIN, Department of Intensive Care Medicine, Infectiology and Emergency Medicine; HR, hazard

3. Beigel JH, et al. N Engl J Med. 2020; 383: 1813–1826. **4.** NIH. Therapeutic management of adults with COVID-19. Available at: https://www.covid19treatmentguidelines.nih.gov/management/clinical-management/

hospitalizedadults--therapeutic-management/. Accessed September 2022. **5.** NICE. Guideline [NG191]. COVID-19 rapid guideline: managing COVID-19. Available at: https://app.magicapp.org/#/guideline/L4Qb5n/section/ERYAXn. Accessed September 2022. **6.** Bartoletti M, et al. Clinical Microbiology and Infection. DOI: 10.1016/j.cmi.2021.11.007. 7. IDSA. IDSA Guidelines on the Treatment and Management of Patients with COVID-19. Available at:

Available at https://www.awmf.org/uploads/tx_szleitlinien/113-001LGI S3 Empfehlungen-zurstationaeren-Therapie-von-Patienten-mit-COVID-19 2022-03.pdf. Accessed September 2022. 10. Infographic: COVID-19: Drug and non-drug therapy recommendations by disease phase.

8. Eleven things you need to know NOW about COVID-19. Available at https://www.mhlw.go.jp/content/

11. Insert reference to local VEKLURY Product Information. This medicinal product is subject to additional monitoring. This will allow quick identification of new safety information. Healthcare professionals are asked to report any suspected adverse reactions.

Adverse events should be reported to Gilead at safety FC@gilead.com or +44 (0) 1223 897500.

This information is intended for healthcare professionals. September 2022 GLB-VKY-0046

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1. Agarwal A, et al. BMJ. 2020; 370: m3379. 2. WHO Solidarity Trial Consortium. Lancet. 2022. DOI: 10.1016/S0140-6736(22)00519-0.

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https://www.idsociety.org/practice-guideline/covid-19-guideline-treatmentand-management/.

