**REMAP-CAP PRESS RELEASE (LAY PUBLIC)**

**Antiplatelet therapy not effective in treating critically ill COVID-19 patients**

For critically ill patients with COVID-19, antiplatelet therapy, either with aspirin or a P2Y12 inhibitor, was ineffective at improving outcome when compared with no antiplatelet therapy.

These findings, based on an analysis of 1467 critically ill patients, which are yet to be published, come from the REMAP-CAP trial, led in the UK by Imperial College and ICNARC. REMAP-CAP began investigating treatments for COVID-19 in March 2020, enrolling hospitalized non-critically ill and critically ill patients. The study design randomises patients to multiple combinations of treatments, enabling researchers to evaluate different treatments for COVID-19, including drugs which modulate the immune response, drugs affecting the coagulation system, and therapies that modulate or support other vital aspects of the body's response to the virus.

Since the start of the pandemic, over 12,500 randomisations of patients with COVID-19 have occurred in the trial, at over 320 hospitals worldwide. REMAP-CAP investigates 48 current or completed interventions in 14 different treatment domains. The study has released and published several platform conclusions (see <https://www.remapcap.eu/publications/> ) that have contributed to finding the best treatment for these patients.

The investigators are working on more detailed analyses, that will be submitted to a peer-reviewed journal shortly. These valuable research findings add significantly to the accumulating evidence of antithrombotic use in patients with COVID-19 and represent the hard work of many individuals. We would like to sincerely thank the investigators, research coordinators, participants and their families, who continue to support REMAP-CAP for their contribution.

**REMAP-CAP PRESS RELEASE (SCIENTIFIC COMMUNITY)**

**Antiplatelet therapy not effective in treating critically ill COVID-19 patients**

For patients with COVID-19 who are receiving organ support in an ICU, antiplatelet therapy, either with aspirin or P2Y12 inhibitor (which were found to be equivalent), was ineffective at improving the composite endpoint of death and organ support free days when compared to no antiplatelet therapy.

These findings, based on an analysis of 1467 critically ill patients, which are yet to be published, come from the REMAP-CAP trial, led in Europe by the University Medical Center in Utrecht, the Netherlands [link]. At a planned adaptive analysis, the probability of futility of antiplatelet therapy (defined as an odds ratio of < 1.2) was 98%, well above the platform threshold of 95%. Previously, aspirin and P2Y12 inhibition had been proven to be equivalent in the trial. The odds ratio for improving the primary outcome of death and organ support free days was 0.99 (95% Credible Interval 0.82 – 1.19) for these drugs, compared to control.

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We look forward to receiving more detailed analysis that includes thrombotic and bleeding outcomes to properly understand the context of these results. These valuable research findings add significantly to the accumulating evidence of antithrombotic use in patients with COVID-19 and represent the hard work of many individuals. We would like to sincerely thank the investigators, research coordinators, participants and their families, who continue to support REMAP-CAP for their contribution.

**REMAP-CAP PRESS RELEASE (TWITTER)**

For patients with COVID-19 receiving organ support in ICU, antiplatelet therapy with aspirin or P2Y12 inhibitor (which were found to be equivalent), was ineffective at improving death and organ support free days when compared to no antiplatelet therapy 1/8

These are initial results from 1467 patients in REMAP-CAP, which are yet to be published. At a planned adaptive analysis, the probability of futility of antiplatelet therapy (defined as an odds ratio of < 1.2) was 98%, well above the platform threshold of 95% 2/8

Previously, aspirin and P2Y12 inhibition had been proven to be equivalent in the trial. The odds ratio for improving the primary outcome of death and organ support free days was 0.99 (95% Credible Interval 0.82 – 1.19) for these drugs, compared to control 3/8

In REMAP-CAP, patients are randomized to multiple combinations of treatments, enabling researchers to evaluate different treatments for COVID-19, including drugs affecting the coagulation system, drugs which modulate the immune response and others 4/8

Since the start of the pandemic, over 12.500 randomizations of patients with COVID-19 have occurred in the trial, at over 320 hospitals worldwide. REMAP-CAP investigates 48 current or completed interventions in 14 different treatment domains 5/8

The study has released and published several platform conclusions (see https://www.remapcap.org) that have contributed to finding the best treatment for these patients.6/8

We look forward to receiving more detailed analysis that includes thrombotic and bleeding outcomes to properly understand the context of these results 7/8

These findings add significantly to accumulating evidence of antithrombotic use in patients with COVID-19 and represent the hard work of many individuals. We sincerely thank investigators, research coordinators, participants and their families for their contribution 8/8