IMMUNOSUPPRESSIVE TREATMENT MANAGEMENT IN A COHORT OF HOSPITALIZED SOLID ORGAN RECIPIENTS AFFECTED BY COVID-19

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BACKGROUND AND OBJECTIVES

Solid organ transplantation (SOT) recipients' immunosuppression management is challenging. Drugs used in COVID-19 involve drug-drug interactions (DDI) with immunosuppressants. The aim of this study was:

- Describe DDIs in hospitalized SOT recipients (SOTr).
- Analyze DDIs management and their clinical impact.

MATERIALS AND METHODS

- Retrospective unicentric study including SOTr with COVID-19 hospitalized during March11th- April25th.
- Clinical data and pharmacotherapy was were recorded from admission up to 28 days(d) or discharge.
- Lexicomp[®] was used to detect and categorize DDI according to: risk level, reliability rating and severity.

RESULTS

46 patients were included:

- 33 (71.7%) men

 -62.7 ± 12.6 (mean \pm SD) years.

Kidney Liver Lung 30 (56.2%) 13 (28.3%) 3 (6.5%)

Immunosuppression at admission - tacrolimus: 41 (89.1%) - mycophenolate mofetil/ mycophenolate sodium: 28 (60.9%) - prednisone: 39 (84.8%) - everolimus: 7 (15.2%) - sirolimus: 7 (15.2%) - cyclosporine: 1 (2.2%)

106 DDIs were detected and affected 42 (91.3%) patients.

Immunosuppressant was withheld in 33 (71.7%) patients due to DDI.





Severity: major

50

100

- 36 (87.7%) out of 41 patients receiving tacrolimus suffered 65 DDIs.

- Treatment was withdrew in 22 patients (61.1%), dose was reduced in 18(50%) patients and increased in 4 (11.1%).
- Through tacrolimus levels were supratherapeutic in 8 (25%) patients at admission, 13 (43.3%) at 48h, 10 (31.3%) at 7d and 2 at 14d (17.7%, n= 28).

7 patients receiving everolimus had 12 DDIs. Treatment was stopped in all of them.

- No graft rejection was detected.
- 2cases of kidney acute failure were attributable to tacrolimus. —

4 patients receiving **sirolimus** had 6 DDIs. Treatment was stopped in all cases.

DDIs were highly prevalent in hospitalized SOTr with COVID-19.

Pharmaceutical care is critical to promptly detect and manage DDIs in SOTr.

