
Informed consent and ethical clearance

Tripartite informed consent was obtained from the patients, the patients' family, and the research ethics committee of the hospital. The patients were informed about the purpose of the study, the procedures, the risks, and the benefits of the study. The patients' family was informed about the purpose of the study, the procedures, the risks, and the benefits of the study. The research ethics committee of the hospital was informed about the purpose of the study, the procedures, the risks, and the benefits of the study.

Data collection

The data were collected from the patients' medical records, the patients' family, and the research ethics committee of the hospital. The data were collected from the patients' medical records, the patients' family, and the research ethics committee of the hospital.

