Consent of the patient (guardian) to participate in medical research

Research Title	Clinical study to compare the Efficacy and Safety of casirivimab & imdevimab, Remdesivir, and Favipravir in hospitalized COVID-19 patients
Search Steps	1- Dividing cases into 3 groups according to antiviral treatment 2- Follow-up cases before, during and after taking medication doses, while they are in the hospital and after their discharge 3- Collecting clinical data on cases in order to compare them statistically and obtain results
Search duration	6 months after taking the approval of the scientific committee
Where to search	Isolation Hospital - Mansoura University
Benefits of Research	Benefiting patients with the Corona virus, by determining the most appropriate antiviral treatment according to the case

	Informing the community of the results of this research, which has changed in the treatment protocol for Corona virus patients
Possible side effects	Some of the rare allergy symptoms that may result from intravenous injections of drugs Therefore, cases are followed up while taking the medicine

I, the undersigned, acknowledge that the researcher has informed me that:

1- The research does not contradict with the values and ethics of society.

2- With emphasis on the confidentiality of the research and my right to leave it without being held accountable and without affecting medical care

Participant Patient's Name.....#### #### ###..... Signature of the participating patient (guardian)..... .....#### #### ###.....

Signature of the Principal Investigator Ahmed Hosny Hassan Moussa.
Date..... 11/8/2022