

The first part of the text discusses the importance of data monitoring and follow-up in clinical trials. It highlights that these processes are essential for ensuring the safety and efficacy of the interventions being tested. The text mentions that data monitoring committees (DMCs) are often established to oversee the trial and make recommendations based on the data collected.

The second part of the text discusses the challenges of data monitoring and follow-up. It notes that these processes can be time-consuming and costly, and that they require a high level of expertise and resources. The text also mentions that there are often difficulties in obtaining and maintaining accurate data, and that this can lead to delays and inefficiencies in the trial.

The third part of the text discusses the benefits of data monitoring and follow-up. It notes that these processes can help to identify safety issues early on, which can prevent serious harm to participants. It also mentions that data monitoring and follow-up can help to ensure that the trial is conducted in a transparent and ethical manner, and that the results are reliable and valid.

The fourth part of the text discusses the role of data monitoring and follow-up in the overall clinical trial process. It notes that these processes are integral to the trial, and that they are essential for ensuring that the trial is conducted in a safe and effective manner. The text also mentions that data monitoring and follow-up can help to build trust between researchers and participants, and that this is essential for the success of the trial.

The fifth part of the text discusses the future of data monitoring and follow-up. It notes that there are many challenges ahead, and that there is a need for continued research and innovation in this field. The text also mentions that there are many opportunities for improvement, and that there is a need for greater collaboration and communication between researchers, regulators, and participants.

In conclusion, data monitoring and follow-up are essential components of clinical trials. They are essential for ensuring the safety and efficacy of the interventions being tested, and for ensuring that the trial is conducted in a transparent and ethical manner.

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Calls on the UN system, in consultation with the international financial institutions, to develop transparent measurements of progress on sustainable development that go beyond per capita income; these should recognize poverty in all of its forms and dimensions, and the social, economic and environmental dimensions of domestic output and structural gaps at all levels (129, MoI 17.19)

Commits to seek to develop and implement tools to mainstream sustainable development, as well as to monitor sustainable development impacts for different economic activities, including for sustainable tourism (129, MoI 12.b)

Committed to work with the international community to develop and implement tools to mainstream sustainable development, as well as to monitor sustainable development impacts for different economic activities, including for sustainable tourism (129, MoI 12.b)

