MAHARAJAH'S INSTITUTE OF MEDICAL SCIENCES

D.No. 31-15, NELLIMARLA - 535 217,
VIZIANAGARAM (DIST.,) Andhra Pradesh, INDIA
(AFFILIATED TO Dr. NTR UNIVERSITY OF HEALTH SCIENCES, VIJAYAWADA)
(Recognized by Medical Council of India / Govt. of India)
(Sponsored by SRI RAMA EDUCATIONAL TRUST)

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Subject: Formation of Medical Device-Related Adverse Event Committee – Reg.

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In compliance with the guidelines of the Ministry of Health & Family Welfare and Medical Device Rules (MDR), 2017, and to ensure timely identification, reporting, and review of any medical device-related adverse events in the hospital, the following **Medical Device-Related Adverse Event Committee (MDAEC)** is hereby constituted with immediate effect:

Medical Device-Related Adverse Event Committee (MDAEC)

S. No	Name of the Member	Designation	Role in Committee	Contact Number
1	DR KETAVATH CHANGA THAVARYA NAIK.	Medical Superintendent	Chairperson	7981540734
2	DR. B. KRISHNA CHAITNAYA	Professor of Anaesthesia	Member	9700561544
3	DR. CHANDRADEVE VARNA BSK	Professor of Paediatrics	Member	7013220392
4	DR TUMMIDI V V VINAY KUMAR	Associate Professor of ENT	Coordinator	9652277741
5	DR. MOULEESWARA KUMAR TAMMA	Associate Professor of General Medicine	Member	9989482403
6	DR ANIL KUMAR KALLEPALLY	Associate Professor of Radiology	Member	9985366699
7	DR PAVANI JERRY	Assistant Professor of Ophthalmology	Member	8985039904
8	DR PRADEEP PALAVALASA	Assistant Professor of Orthopedics	Member	9177353282
9	MS. K.V. PADMA	I/C Nursing Superintendent	Member	9346631938
10	MR. U. JITHENDRA RAJU	OT in-charge	Member	9949329055
11	MR.J VIJAY KUMAR	Manager - Quality	Member	9666343044
12	MR. B. SEETHA RAMA RAJU	Biomedical Supervisor	Member	9492746135
13	MR. I.NAGARAJU	Biomedical Engineer	Member	9330713664

Responsibilities of the Committee:

1. To monitor and assess all medical device-related adverse events reported in the hospital.
2. To ensure timely submission of reports to the Indian Pharmacopoeia Commission (IPC) under the Materiovigilance Programme of India (MvPI).
3. To recommend corrective and preventive actions (CAPA) in cases of recurring events.
4. To educate and sensitize hospital staff on medical device vigilance and reporting protocol.
5. To maintain documentation of all adverse events and investigation reports.

All departments are hereby directed to promptly report any suspected adverse event or malfunction related to medical devices to this committee.

This committee shall remain functional until further orders.

DEAN
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