Artificial Intelligence Based Medical Device Technologies Implementation Strategies in Indian Health Care Industry

A. Narasimha Reddy[1], G. Naga praveen[2], D. Sanjana[3], SK. Sanjuda[4]
[1,2,3] *Department of Regulatory Affairs Hindu College of Pharmacy*
[4]*Co-Author, Hindu College of Pharmacy*

***Abstract*—Artificial intelligence (AI) is transforming the healthcare sector by enhancing disease diagnosis, improving treatment planning, accelerating drug discovery, and optimizing clinical decision-making. The integration of AI technologies, including machine learning, deep learning, natural language processing, and computer vision, has led to the development of intelligent medical devices and Software as a Medical Device (SaMD), enabling more accurate, efficient, and personalized healthcare services. AI-powered applications such as clinical decision support systems, medical imaging, remote patient monitoring, robotic-assisted surgery, electronic health records, and virtual health assistants have significantly improved healthcare delivery while reducing operational costs and human error.**

Despite these advancements, the widespread adoption of AI in healthcare presents several regulatory, ethical, legal, and technical challenges, including data privacy, algorithmic bias, cybersecurity, transparency, accountability, and clinical validation. Regulatory agencies worldwide, including the U.S. Food and Drug Administration (FDA), the Medicines and Healthcare products Regulatory Agency (MHRA), the International Medical Device Regulators Forum (IMDRF), and India's Central Drugs Standard Control Organization (CDSCO), are developing frameworks to ensure the safety, effectiveness, and quality of AI-enabled medical devices. In India, initiatives such as the Ayushman Bharat Digital Mission (ABDM), the Digital Personal Data Protection Act, 2023, and evolving medical device regulations are supporting the responsible integration of AI into healthcare.

This review summarizes the current applications of AI in healthcare, examines AI-enabled medical devices and their regulatory landscape, discusses major implementation challenges and ethical considerations, and highlights future directions for AI-driven healthcare systems. Continued collaboration among healthcare professionals, researchers, policymakers, regulatory authorities, and technology developers will be essential to establish trustworthy, transparent, and patient-centric AI solutions. With appropriate regulatory oversight, robust digital infrastructure, and responsible innovation, artificial intelligence has the potential to revolutionize global healthcare by improving patient outcomes, increasing healthcare accessibility, and promoting precision medicine.

***Index Terms*—Artificial Intelligence, Healthcare, Software as a Medical Device (SaMD), Machine Learning, Deep Learning, Medical Devices, Clinical Decision Support, Digital Health, Healthcare Regulation, Personalized Medicine, India.**

I. INTRODUCTION

AI-based medical device technologies, also known as AI-SaMD (Software as a Medical Device), use deep learning and machine learning to improve diagnostic precision, allow for real-time patient monitoring, and customize care. [1][2] Robotic surgery, wearable health monitors [1], and AI-powered imaging are important applications that greatly enhance patient outcomes while lowering administrative costs [1][2]. Artificial Intelligence (AI) has emerged as one of the most transformative technologies in modern healthcare, revolutionizing the way diseases are diagnosed, treated, and managed[3]. AI refers to the ability of computer systems to perform tasks that normally require human intelligence, including learning, reasoning, problem-solving, decision-making, speech recognition, and visual perception. The rapid advancement of computational power, machine learning (ML), deep learning (DL), natural

language processing (NLP), and computer vision has significantly expanded the scope of AI applications across the healthcare sector. [4] By analyzing large volumes of structured and unstructured healthcare data, AI systems can generate accurate predictions, identify disease patterns, assist clinicians in diagnosis, and support evidence-based clinical decision-making.

The healthcare industry is currently facing numerous challenges, including an increasing prevalence of chronic diseases, aging populations, shortages of healthcare professionals, rising healthcare expenditures, and unequal access to quality medical services. These challenges have created an urgent need for innovative technologies capable of improving healthcare efficiency while maintaining high standards of patient safety and quality of care.[5] AI has emerged as a promising solution by enabling automation of repetitive tasks, enhancing diagnostic accuracy, optimizing treatment planning, and supporting personalized medicine.

One of the most significant developments in this field is the emergence of Artificial Intelligence-enabled Medical Devices and Software as a Medical Device (SaMD). Unlike conventional medical devices that rely on fixed algorithms, AI-enabled medical devices continuously analyze clinical data, learn from new information, and improve their performance over time. These technologies are increasingly being used in diagnostic imaging, pathology, cardiology, ophthalmology, oncology, intensive care monitoring, wearable health devices, robotic surgery, and remote patient monitoring.[6] AI-powered Clinical Decision Support Systems (CDSS) assist healthcare professionals by providing real-time diagnostic suggestions, risk prediction, and treatment recommendations, thereby reducing medical errors and improving patient outcomes.

Artificial intelligence has also transformed pharmaceutical research and drug development. AI algorithms accelerate drug discovery by identifying novel drug targets, predicting molecular interactions, optimizing clinical trial recruitment, and improving pharmacovigilance. In precision medicine, AI integrates genomic, clinical, lifestyle, and imaging data to develop individualized treatment strategies that improve therapeutic efficacy while minimizing adverse drug reactions. Furthermore, AI-powered virtual health assistants and chatbots enhance patient engagement by providing appointment scheduling, medication reminders, symptom assessment, and health education through continuous digital interaction.

Despite its tremendous potential, the integration of AI into healthcare presents several ethical, legal, technical, and regulatory challenges. [7]AI systems rely heavily on high-quality clinical data, making data privacy, confidentiality, cybersecurity, and informed consent major concerns. Algorithmic bias arising from unrepresentative datasets may lead to unequal healthcare outcomes across different patient populations. Additionally, the "black-box" nature of many deep learning models raises concerns regarding transparency, explainability, accountability, and clinician trust. Questions regarding liability for AI-related medical errors and the extent of human oversight remain important issues requiring comprehensive regulatory guidance.

Recognizing these challenges, regulatory agencies worldwide are developing frameworks to ensure the safe and effective use of AI-enabled medical technologies. Organizations such as the U.S. Food and Drug Administration (FDA), the International Medical Device Regulators Forum (IMDRF), the Medicines and Healthcare products Regulatory Agency (MHRA) in the United Kingdom, and the World Health Organization (WHO) have introduced guidelines for the development, validation, and post-market surveillance of AI-based medical devices. In India, regulatory oversight is evolving through the[8] Central Drugs Standard Control Organization (CDSCO), the Medical Devices Rules, 2017, the Digital Personal Data Protection Act, 2023, the Ayushman Bharat Digital Mission (ABDM), and ethical guidelines issued by the Indian Council of Medical Research (ICMR). These initiatives aim to promote responsible AI adoption while ensuring patient safety, data security, transparency, and regulatory compliance.

The future of AI in healthcare is expected to be driven by advances in explainable AI, federated learning, digital therapeutics, precision medicine, genomics, robotics, Internet of Medical Things (IoMT), and interoperable digital health ecosystems [9]. Continued collaboration among healthcare professionals, researchers, policymakers, regulatory authorities, and technology developers will be essential to maximize the benefits of AI while addressing its associated risks. With appropriate governance, ethical implementation, and continuous technological innovation, AI has the

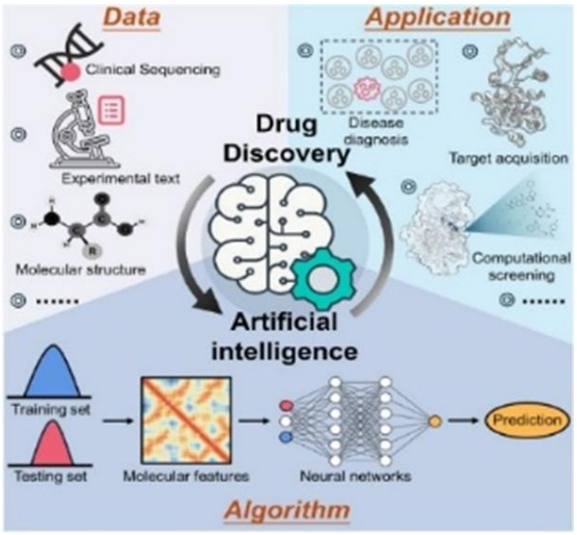
potential to reshape global healthcare by improving diagnostic accuracy, enhancing healthcare accessibility, reducing costs, and ultimately delivering safer, more efficient, and patient-centered care.

Diagnostics and Clinical Imaging

AI in diagnostic and clinical imaging changes healthcare by using machine learning and deep learning to make image analysis more accurate and improve the flow of work [4][3]

Personalized Medicine :AI in personalized medicine uses genomic, clinical, lifestyle, and imaging data to find the best way to diagnose and treat each person [10][12]. It allows for accurate, proactive care, like predicting how drugs will work, finding cancers earlier, and managing long-term conditions [13]1[4].

Drug Discovery and Development

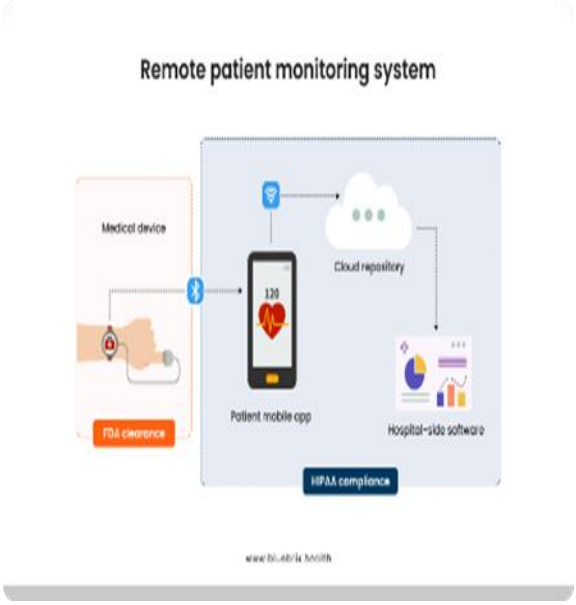


Artificial intelligence is revolutionizing drug discovery and development by significantly reducing research time from years to months, decreasing costs, and identifying new therapeutic candidates [12]. Utilizing machine learning and deep learning, AI enhances the processes of target identification, drug design, and clinical trials, leading to quicker, more accurate, and successful outcomes compared to traditional methodologies [16].

Clinical Decision Support

By analyzing patient data using machine learning and, frequently, LLMs, AI-driven clinical decision support systems (CDSS) improve healthcare by offering real-time diagnostics, individualized treatment recommendations, and better, more affordable care [17].

Remote Patient Monitoring



AI-powered remote patient monitoring (RPM) makes healthcare better by using AI to analyze data from wearables and biosensors in real time [18].

Virtual Health Assistants and Chatbots

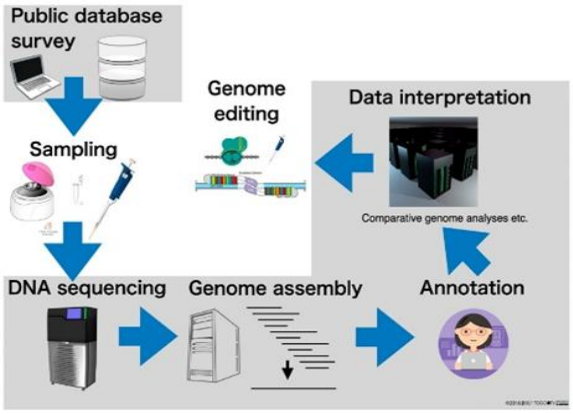
AI-driven virtual assistants using NLP and machine learning enhance patient engagement through real-time services such as 24/7 support, appointment scheduling, and basic health inquiries [19].

Electronic Health Records

AI enables the analysis of large datasets from EHRs containing both structured data (laboratory tests, procedures) and unstructured data (radiology reports, discharge summaries) [20].

Mental Healthcare

AI-powered tools assist in early detection and diagnosis of mental health conditions, generate customized treatment plans, and provide continuous support — bridging resource gaps and promoting stigma-free environments [21].



Research & Development, Bioinformatics, and Genome Editing

The biotechnology sector utilizes AI, especially Machine Learning (ML) and Deep Learning, to accelerate the analysis of vast datasets

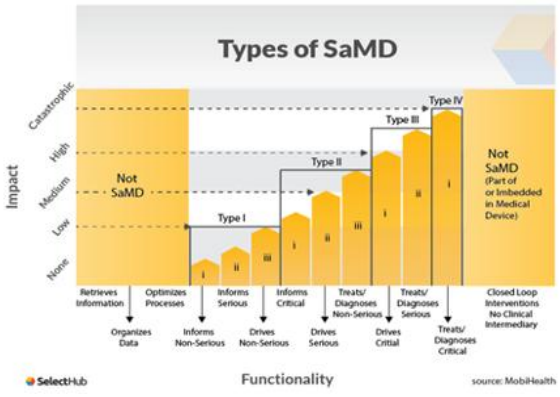
Clinical Trial Recruitment

AI streamlines clinical trial processes by identifying suitable candidates based on patient data, improving recruitment efficiency and diversity. AI also monitors trial data in real-time, enabling rapid detection of problems and protocol adjustments [22].

Robotic Surgery

AI-assisted robotic surgical systems enhance surgeon precision, reduce human error, and shorten recovery times. Real-time AI feedback makes surgeries more efficient and safer, with applications across minimally invasive procedures in neurosurgery, cardiovascular surgery, and beyond [23].

II. AI-ENABLED MEDICAL DEVICES: KEY CONSIDERATIONS



The SaMD Framework

Most national regulatory authorities have converged on classifying AI-based healthcare applications as Software as a Medical Device (SaMD) — software applications not being part of a hardware medical device but intended for medical purposes such as diagnosis, treatment, or monitoring. SaMD can function independently or interact with other devices. Its regulatory framework typically involves risk assessment, clinical validation, and quality, safety, and performance standards [11][13].

Personalized Healthcare Solutions

Machine learning and predictive analytics are used in AI-powered personalized healthcare to customize medical interventions, diagnosis, and prevention based on a patient's genetic, lifestyle, and medical information [13][14].

Patient Safety and Adverse Event Monitoring

By enabling real-time, automated safety risk detection through machine learning and natural language processing, artificial intelligence (AI) improves patient safety and adverse event (AE) monitoring.

III. REGULATORY LANDSCAPE: GLOBAL PERSPECTIVES

Regulators worldwide are taking strides toward regulating AI-based medical technologies to ensure patient safety, address ethical concerns, and protect data privacy. These efforts aim to minimize risks, ensure fairness, promote transparency, and establish clear accountability [1][2].

Table - 1

Dimension	India	United States	United Kingdom
Framework status	No AI-specific legislation; existing healthcare and IT laws apply	FDA-led guidance with multiple published frameworks	Multi-agency approach with dedicated AI programmes in motion
Lead regulatory body	CDSCO (medical devices); MeitY	FDA — Center for Devices and	MHRA — Medicines and Healthcare

	(digital/AI); NHA (digital health)	Radiological Health (CDRH)	products Regulatory Agency
AI / SaMD classification	Under Medical Device Rules, 2017 — typically Class B or C risk	Classified as SaMD under existing device regulations	Classified as SaMD; addressed under the AI as a Medical Device Change Programme
Key guidance documents	Medical Device Rules, 2017 Telemedicine Practice Guidelines, 2020 DPDPA, 2023 (data protection)	Good Machine Learning Practice (GMLP) AI/ML SaMD Action Plan 2024 Report on AI in Medical Products Jan 2025 draft on AI in regulatory decisions	Evidence Standards Framework (NICE, 2019) Software and AI as a Medical Device Change Programme (MHRA, 2021) Regulatory Horizons Council Report (2022)
Post-market & lifecycle monitoring	Static MDR framework — does not account for AI's continuous learning; post-approval change rules create operational burden	Predetermined Change Control Plans (PCCPs) allow pre-approved algorithm updates; ongoing post-market surveillance required	Explicit lifecycle approach — covers cybersecurity, algorithmic changes, bias, and post-market monitoring throughout device life
Liability & accountability	Consumer Protection Act, 2019; liability for design, manufacturing, and marketing defects; AI cannot hold medical licences	Liability shared among developers, healthcare providers, and institutions; FDA enforces product accountability	Patient and public involvement emphasised; MHRA addresses interpretation of AI-driven decisions and accountability

IV. REGULATION OF AI FOR HEALTHCARE IN INDIA

India's regulatory framework for AI-based medical devices and healthcare technologies is evolving, combining existing laws applicable to healthcare and AI technology independently [12].

Key Regulatory Bodies

Multiple regulatory bodies oversee AI in healthcare: CDSCO regulates AI-based medical devices; NHA manages the Ayushman Bharat Digital Health Mission; ICMR ensures clinical validation and ethics; MoHFW drives AI adoption in public health; MeitY governs broader AI and digital health technologies; CCPA protects consumers from unsafe products; and CERT-In ensures cybersecurity for AI healthcare applications [14].

AI as a Medical Device

By increasing diagnostic precision, customizing treatment regimens, and enhancing patient monitoring, artificial intelligence (AI) and machine learning (ML) in software as a medical device (SaMD) are revolutionizing healthcare. [13]

Restrictions on AI in Medical Practice

Only Registered Medical Practitioners (RMPs) licensed by the National Medical Commission (NMC) may practice medicine in India. AI's role in patient evaluation or management necessitates NMC licensing, as it cannot hold medical licenses.

Data Protection and Privacy

The current data protection framework under the IT Act, 2000 and IT (SPDI) Rules, 2011 treats patient health information as Sensitive Personal Data or Information (SPDI). [24]

Intellectual Property Considerations

AI healthcare innovations can be protected by various intellectual property (IP) mechanisms, including patents for novel applications, copyrights for source code and data, trademarks for brand names, and trade secrets for proprietary models.

Consumer Protection and Product Liability

AI healthcare companies may face liability for defects in AI systems causing harm. Liability can arise from: design defects (AI systems with inherently flawed algorithms causing misdiagnosis); manufacturing defects (incorrectly programmed software causing errors); and marketing defects (failure to warn about known risks).

V. CHALLENGES TO REGULATING AI IN INDIA

The regulation of AI as a medical device in India faces challenges due to the dynamic nature of AI technologies. Unlike traditional medical devices, which are static, the current regulatory framework does not adequately account for AI's ability to learn, adapt, and evolve.

Licensing Requirements

While CDSCO has recognised AI-driven SaMD, specific relaxations or guidance remain absent. AI-driven SaMDs are often freely downloadable from websites or app stores channels that cannot realistically obtain medical device sales licences under the MDR [23].

Clinical Investigations

The MDR requires clinical investigations in human participants before novel medical devices can be marketed. CDSCO has not yet issued guidance on how such investigations should be conducted for SaMD — creating regulatory uncertainty for AI product developers. [24]

Post-Approval Changes

MDR's post-approval change framework (major changes requiring prior approval; minor changes requiring notification was designed for static, predictable devices. Given AI systems' dynamic nature — continuously learning and updating — frequent notifications or approvals could result, creating significant operational burden.

Labelling

Mandatory labelling requirements under the MDR and Legal Metrology (Packaged Commodities) Rules, 2011 present unique challenges for AI-driven SaMDs: there is no physical label for downloadable software. Adapting these requirements to software products remains an unresolved regulatory challenge.

Pricing and Advertising

All medical devices are subject to Drugs (Price Control) Order, 2013 pricing restrictions, limiting price increases to 10% per twelve-month period.

Healthcare Practitioner Promotion

The Uniform Code for Marketing Practices in Medical Devices (UCMPMD) governs promotional activities and interactions with healthcare practitioners. AI-driven SaMDs require ongoing training for healthcare professionals — but repeated educational interactions could attract scrutiny under the UCMPMD.

VI. RISKS AND ETHICAL CHALLENGES

Bias and Discrimination

AI models may carry biases from training data, resulting in discriminatory outcomes like unequal healthcare for certain demographic groups. [25]

Data Quality

AI system performance is fundamentally dependent on the quality and representativeness of training data. Without comprehensive data across the full spectrum of patient populations, AI models may produce skewed results, creating risks in diagnosis and treatment. Defining relevant and actionable clinical metrics is itself a complex challenge.[24]

Doctor-Patient Relationship

Increasing AI reliance risks undermining the human dimensions of healthcare. Patients may feel alienated if AI systems assume larger decision-making roles, potentially eroding the therapeutic value of the doctor-

patient relationship and reducing human empathy and judgment in clinical settings.[25]

Privacy and Confidentiality
The integration of AI with big data brings significant ethical, legal, and operational risks regarding patient data privacy. [12]

Physician Preparedness
Widespread AI adoption requires significant training and awareness across the healthcare workforce. In India, where AI adoption may be more challenging in rural or smaller regions, barriers include limited technology access, language differences, lack of training, and varying levels of digital expertise.

Cybersecurity
Growing reliance on AI and big data makes healthcare systems increasingly vulnerable to cyberattacks that could compromise sensitive patient data, disrupt clinical operations, and jeopardize patient safety. [15]

Environmental Impact
The computational power required to train complex AI models — particularly deep neural networks — results in substantial energy consumption and carbon footprint.

Accountability and Explainability
As AI systems gain autonomy in clinical decision-making, the complexity of determining responsibility for errors or harm increases. Transparency, auditability, and the ability to explain their reasoning are crucial to maintain patient trust and facilitate human oversight. Urgent guidelines and disclaimers from regulatory authorities are necessary.[25]

VII. KEY ENABLERS: INDIA'S DIGITAL HEALTH ECOSYSTEM

Ayushman Bharat Digital Mission (ABDM)
The ABDM is India's pivotal initiative for creating an integrated digital health ecosystem. Its primary components — the Ayushman Bharat Health Account, Health Facility Registry, Healthcare Professionals Registry, and Unified Health Interface — aim to facilitate interoperability and data-driven healthcare delivery across the country.[21]

Digital Public Infrastructure
India's robust digital public infrastructure — accessible, secure, and interoperable — has played a transformative role in the digital economy. Combined with mobile technology proliferation and improving connectivity, this infrastructure provides the foundational layer for AI-enabled healthcare delivery at scale, including telemedicine expansion.[25]

Telemedicine — e-Sanjeevani
Telemedicine platforms are effectively overcoming geographical barriers, particularly in rural and underserved regions. e-Sanjeevani exemplifies a telemedicine service delivering healthcare to millions of individuals [12]

Government Initiatives
The Indian government is promoting digital health through the National Health Policy and National Digital Health Mission, fostering the adoption of digital technologies in healthcare. Initiatives like the Pradhan Mantri Jan Arogya Yojana are establishing an ecosystem for AI-driven healthcare innovations.[12]

VIII. FUTURE DIRECTIONS

Accelerated AI Integration
AI is projected to drive significant advancements across all facets of healthcare delivery. This technology will enhance diagnostic capabilities, improve operational efficiency through automation of administrative tasks, and contribute to improved patient outcomes. Healthcare AI investment globally is expected to reach $11.78 billion by 2025.[25]

Expansion of Telemedicine Services

Telemedicine services are expected to continue their substantial expansion, maintaining a vital role in broadening healthcare access, particularly in underserved regions. As regulatory frameworks mature, AI's role in telemedicine will likely evolve from purely assistive to more substantive clinical functions, under appropriate oversight mechanisms.[12]

Development of Digital Health Ecosystems

The ABDM is spearheading development of more interconnected digital health infrastructure, aiming to enhance interoperability among healthcare providers and deliver seamless patient experiences. This includes integration of AI tools across diagnostic, therapeutic, and administrative workflows to create cohesive digital health systems. [15]

Data-Driven and Federated Healthcare

AI-based analytics will become a key driver of personalized medicine and predictive diagnostics. Innovations like federated learning — training predictive models across decentralized data sources without centralizing patient data — offer promising solutions that enable collaboration while maintaining privacy. [21]

Emerging Technologies Beyond AI

Several adjacent technologies will significantly impact healthcare delivery, including: Virtual and Augmented Reality (VR/AR) for medical training and procedures; Robotics enhancing surgical efficiency and access; Nanotechnology for targeted drug delivery in treatments like cancer; 3D Bioprinting addressing organ shortages by creating biomimetic structures; and Pharmacogenomics improving drug prescription through genetic testing to enhance efficacy and reduce adverse reactions.

Global Collaboration and Standardization

The WHO is advocating for increased global collaboration in digital health policy development, focusing on interoperability, security, and data-driven policy applications. Standardized data exchange protocols (e.g., HL7 FHIR) are crucial for enabling seamless integration across heterogeneous healthcare information systems. International alignment on AI standards will be essential for ensuring that digital health solutions are safe, effective, and mutually transferable.[25]

IX. CONCLUSION

• AI integration in healthcare can transform patient care, efficiency, and medical research, particularly in India.
• India faces challenges like healthcare workforce shortages and chronic disease burdens, making AI a significant solution.
• The regulatory landscape is adapting, addressing issues such as liability, accountability, and algorithmic transparency as AI systems gain autonomy.
• Multi-stakeholder collaboration is essential: policymakers need adaptive frameworks, healthcare providers must enhance AI literacy, and technology developers must ensure safety and fairness.
• Patient empowerment as informed participants in AI-enabled care is crucial.
• Investment in tech infrastructure, skill development, and regulatory standards is necessary for effective AI deployment.
• India's ethical guidelines and emerging AI standards position it as a model for global AI health transformation, aiming to improve health outcomes and frameworks.

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