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MEDICAL TOURISM

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ABSTRACT

Medical tourism has become a major component of cross-border health services, characterized by patients travelling internationally to obtain medical care that is more affordable, faster, or perceived as higher in quality than what is available in their home jurisdictions. Cost differentials, lengthy waiting periods in certain public health systems, expanding private-sector healthcare capacity, and technological advances have collectively accelerated this market. India has emerged as a leading destination due to comparatively lower treatment costs, a large pool of specialized clinicians, the growth of accredited tertiary-care hospitals, and the co-existence of biomedical and traditional wellness systems. At the same time, the sector generates complex legal and ethical concerns, particularly in relation to patient safety, informed consent, professional accountability, dispute resolution, continuity of care, confidentiality, and distributive justice in access to health resources. This article analyses the concept and expansion of medical tourism, outlines its perceived benefits and structural risks, and argues for clearer governance mechanisms that safeguard patient rights while supporting sustainable sectoral growth.

Keywords: Medical Tourism, Cross-Border Healthcare, Medical Negligence, Informed Consent, Global Health Governance, etc.

CHAPTER 1 – INTRODUCTION

Globalization, improved connectivity, and the increasing commercialization of healthcare have contributed to the rapid expansion of medical tourism. The term generally refers to the cross-border movement of patients who seek medical or health-related services outside their home country, frequently motivated by lower costs, shorter waiting times, specialized expertise, or access to technologies not readily available domestically. Over time, what began as relatively sporadic travel for treatment has evolved into a structured industry integrating healthcare delivery with hospitality, travel facilitation, and international marketing.

Developing economies have increasingly positioned themselves as competitive destinations by investing in clinical infrastructure, specialist capacity, and hospital

branding. India, in particular, has drawn sustained international interest because it combines comparatively low procedure costs with advanced clinical services and an expanding network of hospitals that pursue national and international accreditation credentials. Alongside these economic and infrastructural benefits, medical tourism also produces regulatory stress points, including questions of legal liability for adverse outcomes, enforceability of patient rights, standards for informed consent in cross-cultural settings, and the distributional impact on domestic patients when providers prioritize higher-paying international demand.

Critical questions about patient safety standards, the validity of informed consent in cross-cultural settings, medical negligence liability, and the security of sensitive health data are brought up by the unchecked expansion of

this international sector.⁴ Furthermore, concerns about local citizens' fair access to healthcare are raised by the diversion of resources to serve foreign patients. Such an inquiry must not only evaluate the economic and soft-power benefits of the trade but also underscore the urgent necessity for a cohesive regulatory framework capable of ensuring ethical medical practice and the rigorous enforcement of patient rights in a cross-border setting.¹⁸⁵⁰

CHAPTER 2 – CONCEPTUAL FRAMEWORK: MEANING & SCOPE OF MEDICAL TOURISM

Medical tourism may be understood as the organized movement of individuals across national or regional boundaries for medical, surgical, therapeutic, rehabilitative, cosmetic, or wellness services. In many cases, medical travel is not limited to curative interventions; it can include preventive screenings, elective procedures, fertility services, rehabilitation programs, and wellness therapies. In jurisdictions such as India, the sector's scope is often described as including both conventional clinical care and structured traditional or alternative systems that are offered to international patients as part of broader "integrative" health packages.¹⁸⁵¹

Conceptually, the sector reflects the globalization of health services, where treatment becomes tradable across borders and patients behave, in part, as consumers selecting among competing destinations. This market orientation increases the importance of quality assurance, transparent pricing, ethical clinical practice, and clear accountability when treatment outcomes are disputed. Medical tourism has greatly broadened its reach beyond its original emphasis on life-saving or curative treatments. Modern research acknowledges a wider range of services, from elective interventions like cosmetic surgery, dental care, and reproductive treatments to

necessary surgeries like cardiac and orthopedic procedures. Additionally, the field is expanding to include wellness tourism, rehabilitation, and preventive care. This description is particularly thorough in the particular setting of India, fusing traditional indigenous medical systems with biological interventions.

Medical tourism is essentially a symbol of the globalization and commercialization of healthcare. It turns medical care from a regional public good into a transnational commodity that may be traded. This change redefines the patient-physician interaction within a framework driven by the market, posing crucial problems about consumer rights protection in an increasingly globalized medical market, regulatory control, and cross-border ethical norms.¹⁸⁵²

CHAPTER 3 – EVOLUTION & GROWTH TRENDS

Historically, international patient flows often moved from lower-income to higher-income countries for specialized treatment perceived to be unavailable locally. Over the late twentieth and early twenty-first centuries, that direction partially reversed as multiple developing states expanded private tertiary-care capacity, strengthened specialist training, and adopted internationally recognizable accreditation practices. These structural shifts, coupled with increased information access and improved air connectivity, supported the growth of medical tourism as a global industry.

Medical tourism's historical course has experienced a significant structural change. In previous decades, the predominant pattern of patient movement was unidirectional, with elites from poor countries flying to the Global North (mostly Western Europe and the United States) to obtain cutting-edge medical treatments that were not available in their home countries.¹ Disparities in clinical knowledge and technology capabilities were the driving forces behind this "South-to-North" shift. However, this trend has reversed due to

¹⁸⁵⁰ Valorie A. Crooks et al., Ethical and Legal Implications of the Risks of Medical Tourism for Patients: A Qualitative Study of Canadian Health and Safety Representatives' Perspectives, 3 *BMJ OPEN* e002302 (2013).

¹⁸⁵¹ I. Glenn Cohen, Patients with Passports: Medical Tourism, Law, and Ethics 5 (2014).

¹⁸⁵² Nathan Cortez, Recalibrating the Legal Risks of Cross-Border Health Care, 10 *YALE J. HEALTH POL'Y L. & ETHICS* 1, 6 (2010).

the emergence of globalization and the quick spread of technology. The contemporary market is driven by healthcare price disparities, capacity constraints in certain national systems, the availability of high-volume specialist centers, and patient preferences for faster access to elective and complex procedures. While the industry can generate foreign-exchange inflows and spur health infrastructure investment, it also raises transnational governance problems, including variability in standards, fragmented dispute pathways, and inconsistent post-operative follow-up once patients return home¹⁸⁵³.

As a result, modern medical tourism has developed into a multibillion dollar global industry. Demographic changes, particularly the aging of populations in developed nations and the rising prevalence of chronic illnesses, support the sector's growth. Additionally, the obstacles of entrance for potential medical tourists have been reduced by the standardization of care through international accreditation (like JCI) and better air connections. Medical tourism is now a vital part of the global healthcare scene, spurring cross-border competition and innovation, rather than a niche business.

CHAPTER 4 - GLOBAL LANDSCAPE

A number of states have actively branded themselves as medical tourism hubs by combining specialized clinical services with tourism infrastructure and targeted facilitation measures. Commonly discussed destinations include India, Thailand, Malaysia, Singapore, Turkey, and South Korea. The global market is shaped by uneven access to specialized care and by substantial differences in treatment costs and waiting times. Patients from systems with high out-of-pocket expenses or long queues may seek alternatives abroad, while provider states view medical tourism as both an

economic opportunity and a catalyst for private-sector expansion.

At the same time, cross-border healthcare markets intensify concerns about patient safety and continuity of care, especially where clinical standards, malpractice norms, and complaint mechanisms differ widely. These concerns underscore the importance of minimum standards, enforceable disclosure and consent processes, and credible oversight mechanisms capable of protecting patients even when disputes have cross-jurisdictional dimensions. Macro-economically speaking, systemic inequalities are the main forces behind this international commerce. Due to high treatment costs, insufficient insurance coverage, or long waiting lists that are a feature of public health systems (such as the NHS in the UK or Canadian Medicare), patients in industrialized countries are often forced to look for offshore alternatives. On the other hand, developing countries take advantage of these inequalities by providing a "value proposition" that offers significant cost advantages together with equivalent medical expertise. A consistent flow of foreign patients to emerging nations has been fueled by this arbitrage potential.¹⁸⁵⁴

Patient safety, the standardization of clinical norms, the enforceability of legal remedies across jurisdictions, and the moral ramifications of commodifying treatment are the most important of these.¹⁸⁵⁵ Therefore, improved international coordination and the creation of strong regulatory oversight systems to guarantee fair and safe practices globally are essential to the industry's sustainability.

CHAPTER 5 - MEDICAL TOURISM IN INDIA: KEY FEATURES

India is widely recognized as a major medical travel destination due to the combination of (i) comparatively lower costs for many procedures, (ii) large-scale tertiary-care

¹⁸⁵³ Valorie A. Crooks et al., What is Known About the Patient's Experience of Medical Tourism? A Scoping Review, 10 BMC HEALTH SERVS. RES. 266 (2010).

¹⁸⁵⁴ Meghann Orma, Medical Tourism in India: A Legal Perspective, 4 INTL J. L. MGMT. & HUMAN. 1205 (2021).

¹⁸⁵⁵ I. Glenn Cohen, Protecting Patients with Passports: Medical Tourism and the Patient-Protective Argument, 95 IOWA L. REV. 1467 (2010).

capacity in private and corporate hospitals, (iii) specialization in areas such as cardiac care, orthopedics, oncology, transplantation, reproductive care, and cosmetic procedures, and (iv) the availability of traditional systems marketed internationally as wellness complements. Many hospitals also seek to demonstrate quality through accreditation systems, including those administered by the National Accreditation Board for Hospitals & Healthcare Providers and international accrediting entities.¹⁸⁵⁶

India's public policy has also supported the sector through visa facilitation and coordinated promotion of health-related travel. However, rapid market growth increases the need for clearer regulatory allocation of duties—among clinicians, hospitals, facilitators, and insurers—and more predictable dispute resolution for international patients. India has firmly established itself as a premier destination within the global medical tourism landscape. The country attracts a substantial volume of international patients, drawn by a distinctive convergence of cost-effectiveness, highly qualified medical personnel, and state-of-the-art clinical infrastructure. To ensure alignment with global safety benchmarks, a growing number of Indian hospitals have secured accreditation from the Joint Commission International (JCI), alongside rigorous national certification from the National Accreditation Board for Hospitals & Healthcare Providers (NABH). This dual-layer accreditation framework serves as a critical signal of quality assurance to prospective international patients.

Even if the sector creates a large number of jobs and foreign exchange gains, its quick growth demands that the regulatory framework be strengthened concurrently to handle new ethical, legal, and accessibility issues.

DRIVERS IN MEDICAL TOURISM

Multiple factors explain why patients choose cross-border treatment. First, significant cost differentials encourage patients to seek care abroad, particularly when domestic insurance does not adequately cover elective or specialist interventions. Second, long waiting times in some public systems drive patients toward destinations where scheduling is faster. Third, patients may perceive destination states as offering specialized expertise, high-volume centers, or advanced technologies. Fourth, facilitative policies—such as medical visas, patient concierge services, and bundled treatment-and-travel arrangements—reduce transaction costs for foreign patients.¹⁸⁵⁷

Beyond these structural drivers, cultural and therapeutic preferences (including interest in wellness or traditional therapies) also influence patient decisions, particularly where destinations actively market such services as part of recovery or preventive care programs.

ROLE OF GOVERNMENT & POLICY MEASURES

Governments influence medical tourism by shaping visa policy, quality regulation, professional licensing, and consumer protections. In India, professional standards and ethics for medical practice are regulated through the statutory framework governing medical education and licensure, including the National Medical Commission regime. In addition, healthcare marketing and delivery interact with broader consumer-protection norms, which can become relevant when patients allege service deficiencies.¹⁸⁵⁸

From a governance standpoint, medical tourism benefits from coordinated policy across health, tourism, immigration, and external affairs institutions. Yet coordination alone is insufficient without enforceable patient-safety requirements, standardized disclosure practices, and credible monitoring that

¹⁸⁵⁶ Accredited Hospitals, NATL ACCREDITATION BD. FOR HOSPS. & HEALTHCARE PROVIDERS, <https://www.nabh.co/> (last visited Dec. 18, 2025).

¹⁸⁵⁷ Milad Haghdoost et al., Medical Tourism in India: A Systematic Review, 12 J. EDUC. HEALTH PROMOTION 1, 3 (2023).

¹⁸⁵⁸ Rami J. Al-Azri, Medical Tourism in India: A Legal Perspective, 4 INT'L J. L. MGMT. & HUMAN. 1205, 1210 (2021).

prevents unsafe or misleading conduct in an industry that often involves intermediaries.

CHAPTER 7 – LEGAL AND REGULATORY FRAMEWORK: ACCOUNTABILITY AND REMEDIES

India's medical tourism activities are regulated through a combination of professional standards, hospital accreditation practices, and general legal frameworks applicable to healthcare services. Professional conduct expectations flow from medical regulatory norms administered through the National Medical Commission system. Remedies for deficient services may also arise under consumer-protection law, including the Consumer Protection Act, 2019, which provides institutional pathways for addressing consumer grievances regarding service deficiencies.

However, significant challenges persist because medical tourism disputes are frequently cross-border in nature. International patients may face barriers relating to jurisdiction, applicable law, evidence collection, and enforcement of awards. Moreover, where care involves facilitators or package providers, questions arise regarding the allocation of responsibility among the hospital, clinician, intermediary, and any insurance entity.

ETHICAL ISSUES & PATIENT RIGHTS

The ethical risks of medical tourism are closely tied to asymmetries of information, differences in language and culture, and the commercialization of clinical decision-making. Informed consent is central: patients must receive understandable, accurate disclosures regarding risks, benefits, alternatives, expected outcomes, and post-treatment needs. Confidentiality and responsible handling of patient data also become more complex in cross-border settings where information may be transferred across jurisdictions with different privacy protections.

Medical tourism can also pose distributive justice concerns. Where private capacity is diverted to higher-paying international patients,

local populations may experience reduced access, longer waits, or increased prices. Ethical governance therefore requires safeguards that ensure international market participation does not undermine fair access for domestic patients, particularly for essential and urgent care. These concerns align with widely accepted principles for research and clinical ethics emphasizing respect for persons, beneficence, and justice.

CHAPTER 8 – ECONOMIC IMPLICATIONS

Medical tourism can contribute to foreign-exchange inflows, private investment in high-end clinical infrastructure, employment generation across healthcare and hospitality sectors, and reputational gains that attract further international collaborations. These effects can indirectly benefit domestic patients where infrastructure and skill development spill over into broader health services.

At the same time, unregulated expansion risks entrenching inequities, encouraging aggressive marketing that over-promises outcomes, and creating incentives for volume-driven care that may conflict with patient-centered clinical judgment. Sustainable economic benefit therefore depends on credible oversight, transparent pricing, and accountable clinical governance.¹⁸⁵⁹

KEY CHALLENGES

Medical tourism faces recurring challenges that affect safety and legitimacy. Regulatory gaps remain where there is no single, dedicated framework specifically tailored to cross-border patient pathways and the intermediary structures that often facilitate them. Continuity of care after discharge is another problem; patients returning home may struggle to secure follow-up care, manage complications, or obtain access to the treating team for post-operative guidance. Dispute resolution is also difficult, as cross-border claims can be expensive and procedurally complex.

¹⁸⁵⁹ Rupa Chanda, Trade in Health Services, 80 BULL. WORLD HEALTH ORG. 158, 159 (2002).

Communication barriers—language differences, culturally distinct expectations of disclosure, and varying health literacy—can undermine informed consent and treatment compliance. Quality assurance is uneven where accreditation is voluntary or poorly monitored. Ethical concerns persist where commercial incentives may skew provider priorities or where foreign patient demand influences resource allocation.¹⁸⁶⁰

COMPARATIVE PERSPECTIVES

Differences in cost structures, regulatory rigor, and specialization shape the competitive positioning of medical tourism destinations. Some states emphasize high-end tertiary care, others focus on elective procedures, cosmetic services, or wellness tourism. These differences highlight the importance of quality assurance and legal predictability: destinations with credible accreditation and effective complaint mechanisms can build higher trust and reduce disputes.

For India, comparative analysis reinforces the need to align rapid growth with enforceable minimum standards, transparent information practices, and clearer accountability norms across hospitals, clinicians, and intermediaries.

NEED FOR REFORMS & REGULATORY STRENGTHENING

To sustain growth while safeguarding patient rights, reforms should focus on standard-setting, monitoring, and dispute resolution design. Effective governance would include clearer rules on disclosure and consent; transparent pricing obligations; minimum service standards for facilities that cater to international patients; and defined responsibilities for intermediaries and package providers. Strengthening post-treatment follow-up protocols, including cross-border care coordination and telemedicine-based continuity pathways, can mitigate clinical risk.¹⁸⁶¹

There is a critical need to enforce uniform quality protocols across all clinical establishments catering to foreign nationals to mitigate disparities in treatment outcomes. To decide medical negligence claims involving international plaintiffs, a specific legal framework is needed to handle complicated conflict-of-laws issues and guarantee easily accessible corrective procedures. Finally, credible enforcement—through licensing oversight, consumer protection institutions, and hospital accountability mechanisms—can improve trust and reduce preventable harm.

CHAPTER 9 - CONCLUSION

Medical tourism has become a prominent feature of the global healthcare economy, driven by price differentials, access constraints, and expanding private clinical capacity. India's growth in this sector reflects skilled clinical labor markets, expanding infrastructure, and the strategic packaging of modern and traditional health services. Yet the same market dynamics that generate economic benefits also create heightened legal and ethical exposure, particularly regarding informed consent, negligence and liability, data confidentiality, and equitable access for domestic populations. A more coherent regulatory approach—one that clarifies duties, strengthens quality assurance, and improves remedy mechanisms for international patients—can support sustainable development while protecting patient rights and ethical medical practice.

However, there are significant structural inconsistencies with the quick commercialization of cross-border care. Complex legal and ethical vulnerabilities have resulted from the industry's growth, especially with regard to medical malpractice liability, the legitimacy of informed consent in cross-cultural contexts, and the possible displacement of domestic health priorities. Therefore, market forces alone cannot ensure the survival of medical tourism. It depends on the development of a strict regulatory framework that strikes a balance between financial needs

¹⁸⁶⁰ Jeremy Snyder et al., Medical Tourism Facilitators: Patterns of Service Differentiation, 15 J. MED. INTERNET RES. e281 (2013).

¹⁸⁶¹ Nicolas P. Terry, Under-Regulated Healthcare Phenomena: An American Perspective, 9 YALE J. HEALTH POLY L. & ETHICS 379, 412 (2009).



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