

## Tradition of “Traditional Medicine” - Bangladesh Perspective

Type: Review Article

Received: September 21, 2025

Published: October 02, 2025

**Citation:**

Mamun Al Mahtab., et al. “Tradition of “Traditional Medicine” - Bangladesh Perspective”. PriMera Scientific Surgical Research and Practice 6.3 (2025): 13-16.

**Copyright:**

© 2025 Mamun Al Mahtab., et al. This is an open-access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

**Sheikh Mohammad Fazle Akbar<sup>1</sup>, Chowdhury Faiz Hossain<sup>2</sup>, Gazi Nurun Nahar Sultana<sup>3</sup>, Sitesh Chandra Bachar<sup>4</sup>, Shahabuddin K Choudhuri<sup>5</sup>, Rezaul Karim<sup>6</sup>, Sakirul Khan<sup>7</sup>, Md. Zakir Sultan<sup>8</sup>, Md. Enayet Ali Pramanik<sup>9</sup>, Rakibul Hasan<sup>10</sup>, Debabrata Karmakar<sup>11</sup>, Rejina Afrin<sup>12</sup>, Md. Abdur Rahim<sup>13</sup>, Sheikh Mohammad Noor E Alam<sup>14</sup>, Rokshana Begum<sup>15</sup>, Ahmed Lutful Moben<sup>16</sup>, Manas Saha<sup>17</sup>, Md. Rezwanur Rahman<sup>18</sup>, Musarrat Mahtab<sup>19</sup> and Mamun Al Mahtab<sup>20\*</sup>**

<sup>1</sup>*Ehime University, Ehime, Japan, Oita University, Oita, Japan and Miyakawa Memorial Research Foundation, Tokyo, Japan*

<sup>2</sup>*Department of Pharmacy, East West University, Dhaka, Bangladesh*

<sup>3</sup>*Centre for Advanced Research in Sciences (CARS), University of Dhaka, Dhaka, Bangladesh*

<sup>4</sup>*Department of Pharmacy, University of Dhaka, Dhaka, Bangladesh*

<sup>5</sup>*Department of Pharmacy, Jahangirnagar University, Dhaka, Bangladesh*

<sup>6</sup>*Institute of Technology Transfer and Innovation, Bangladesh Council of Scientific and Industrial Research, Dhaka, Bangladesh*

<sup>7</sup>*Department of Microbiology, Faculty of Medicine, Oita University, Oita, Japan*

<sup>8</sup>*Centre for Advanced Research in Sciences (CARS), University of Dhaka, Dhaka, Bangladesh*

<sup>9</sup>*On-Farm Research Division, Bangladesh Agricultural Research Institute, Rajshahi, Bangladesh*

<sup>10</sup>*Institute of Technology Transfer and Innovation, Bangladesh Council of Scientific and Industrial Research, Dhaka, Bangladesh*

<sup>11</sup>*Institute of Technology Transfer and Innovation, Bangladesh Council of Scientific and Industrial Research, Dhaka, Bangladesh*

<sup>12</sup>*Department of Pharmacy, East West University, Dhaka, Bangladesh*

<sup>13</sup>*Department of Hepatology, International Medical College, Gazipur, Bangladesh*

<sup>14</sup>*Department of Hepatology, Bangladesh Medical University, Dhaka, Bangladesh*

<sup>15</sup>*Department of Hepatology, Shaheed Suhrawardy Medical College, Dhaka, Bangladesh*

<sup>16</sup>*Department of Hepatology, Kurmitola General Hospital, Dhaka, Bangladesh*

<sup>17</sup>*Department of Hepatology, Khulna Medical College, Khulna, Bangladesh*

<sup>18</sup>*National Gastroenterology Institute and Hospital, Dhaka, Bangladesh*

<sup>19</sup>*Department of Biochemistry, North South University, Dhaka, Bangladesh*

<sup>20</sup>*Department of Hepatology, Bangladesh Medical University, Dhaka, Bangladesh*

**\*Corresponding Author:** Professor Mamun Al Mahtab, Department of Hepatology, Bangladesh Medical University, Dhaka, Bangladesh.

## Abstract

History of medicine has its roots in ancient times dating before the birth of Christ. Medicine has since then undergone massive evolution, more so in the 20<sup>th</sup> and 21<sup>st</sup> centuries. Traditional medicine has now mostly been replaced by evidence-based Allopathic medicine. However, the need to align traditional medicine with modern day medicine is being recognized. Countries like Bangladesh, with rich history of traditional systems of medicine, can be in the forefront. For this further work including research, appropriate policy for pharmaceutical industries etc. is now time demanding.

**Keywords:** Traditional medicine; history; Bangladesh; policy for pharmaceutical industry

## History of development of medicine

The history of medicine is a long and fascinating journey, evolving from ancient practices to modern science. Early medicine often involved herbal remedies, animal parts and minerals, with the earliest known medical texts dating back to 2500 BC in ancient Ebla. Key figures like Hippocrates, the ‘father of medicine’, emphasized on rational approach based on observation and logical reasoning, shifting the focus from superstition. Later, scholars like Galen, Leonardo da Vinci and William Harvey made significant contributions. The 20<sup>th</sup> century saw major advancements in understanding infectious diseases and developing remedies like antibiotics and vaccines [1]. The entire 20<sup>th</sup> and 21<sup>st</sup> century experienced unique developments of innovative therapies. However, most of these concepts and modalities have not served humanity adequately.

Bangladesh is a country of 170 million population. In this country there are thousands of plants endowed with medicinal properties. These plants are used in traditional medicine for centuries in Bangladesh and its neighboring countries [2]. Mostly, these herbs and plants are regarded as efficacious in various pathological conditions. However, these drugs, in most cases have not been properly evaluated for their safety, efficacy and proper indications.

## A brief sketch of traditional medicine

Traditional medicine has a long history. It is the sum of knowledge, skills and practices based on the theories, beliefs and experiences indigenous to different cultures, used for maintaining health and prevention, diagnosis and treatment of physical and mental illnesses.

For centuries, traditional and complementary medicine has been an integral resource for health in households and communities. One hundred seventy countries of the world use traditional medicine, with acupuncture being the most common form of practice in 113 countries. Many developed countries have also begun recognizing and integrating traditional medicine into their healthcare systems [3]. According to 2012 data, almost half the population in many industrialized countries uses some form of traditional and complementary medicine (e.g. USA 42%, Australia, 48%, France, 49%, Canada 70%).

The World Health Organization (WHO) traditional medicine strategy 2014-2023 was developed and launched in response to the World Health Assembly resolution on traditional medicine (WHA62.13). The strategy aims to support member states in developing proactive policies and implementing action plans that will strengthen the role traditional medicine plays in keeping populations healthy. Addressing the challenges, responding to the needs identified by member states and building on the work done under the WHO traditional medicine strategy 2002-2005, the updated strategy for the period 2014-2023 devotes more attention to prioritizing health services and systems, including traditional and complementary medicine products, practices and practitioners.

With the support of the Indian government, WHO founded the Traditional Medicine Centre (GTMC) in 2022 to meet this demand worldwide. The Division of Universal Health Coverage and Life Course, the GTMC is a WHO Headquarters department that is stationed in Jamnagar, Gujarat, India. In order to promote traditional wisdom and contemporary science for the health and well-being of people and the earth, the Centre works to advance research, facilitate knowledge exchange, conserve biodiversity, and build collaborations.

### ***Traditional medicine in Bangladesh***

Bangladesh possesses rich flora of medicinal plants. Continuous use of these plants as ingredients of traditional medicine in the treatment and management of various health problems for generation after generation has made the system an integral part of the culture of this country. Both the Unani and Ayurvedic systems of traditional medicine have firm roots in Bangladesh and are widely practiced all over the country [2-7]. The Unani-Tibb or Graeco-Arab system was developed by the Arab and Muslim scholars from the ancient Greek system, while the Ayurvedic system, which is the old Indian system is based on the Vedas, scriptures from the Aryan age.

Out of the estimated 5000 species of different plants growing in the country, more than 1000 are regarded as having medicinal properties. Use of these plants for therapeutic purposes has been in practice in this country since time immortal [8]. Now, in this age of highly advanced Allopathic medicine, vast majority of the population of this country still prefer traditional medicine in the treatment of most of their diseases even though modern medical facilities may be available in the neighborhood.

### ***Importance of safety assessment in establishing a medicine***

The formal monitoring of medicine safety, known as pharmacovigilance, began in the early 1950s, with the US Food and Drug Administration and academic medical centers taking the lead. This was largely in response to concerns about the safety of chloramphenicol, which was linked to aplastic anemia and other blood disorders. The 1962 Kefauver-Harris Amendments to the Federal Food, Drug, and Cosmetic Act further solidified this by requiring adverse event evaluation and reporting [9].

Prior to this, while there was some awareness of medication-related harms, it wasn't systematically monitored or regulated. The development of pharmacovigilance, including activities like clinical trial monitoring, post-market surveillance and adverse event reporting, has been a gradual process, with ongoing improvements and expansions in scope. The WHO Programme for International Drug Monitoring (WHO PIDM), established in 1968, also played a crucial role in coordinating international efforts to monitor medication safety.

The concept of drug safety, also called 'Medication Safety', is not new, especially in the developed countries in the field of health. For instance, USA has an experience for more than a century in the field of the safety of medications. In October 1937, the use of the antibiotic sulfanilamide caused the death of more than 100 people in the USA. These deaths were not due to the active ingredient itself, rather, they were caused by addition of diethylene glycol (DEG), the excipient used as a solvent for the active drug [10, 11]. DEG was supposed to be inert with no therapeutic benefits. However, it was this toxic substance that led to those fatal side effects. The company claimed they did not foresee these side effects, which was true, as they did not commit animal studies before they marketed the drug. Because of this incident, the US Food and Drug Administration (FDA) approved an act to ensure the safety of any drug by conducting non-clinical and clinical studies before the drugs are marketed for public use [12-21].

### ***Compilation of the policy for pharmaceutical industries in Bangladesh and research directions***

The pharmaceutical industry in Bangladesh is a well-developed sector, contributing significantly to the nation's economy and health-care. It fulfills 98% of the country's medicine requirements and has also emerged as a prominent exporter, reaching over 160 countries.

The policy of the pharmaceutical industries of Bangladesh is to buy active ingredients and packing them as consumable drugs. There is lack of research and development (R&D) concept. Universities and medical institutions rarely undertake studies directed to drug development. In fact, the first step of drug development that requires animal study has neither been initiated nor formulated in this country in medical field. Thus, there is a need to have a strong collaboration among different institutions so that R&D may move to its target of drug development. Universities also lack proper manpower, instinct and direction for drug development.

## Conclusion

Our pharmaceutical industries now enjoy patent exemption, which will disappear after 2033, by the time when Bangladesh completes its graduation from least developed to developing country status. This will likely result in a surge in drug prices. It is therefore time demanding that collaboration between universities to conduct research with available herbal drug of Bangladesh. The following features are important facts that were considered for fixing aims and objectives.

## References

1. Cook HJ. “The history of medicine and scientific revolution”. The University of Chicago Press Journal 102.1 (2011).
2. Saha D. “Potentials and Practice of Traditional Medicine in Bangladesh”. Al-Kufa University Journal for Biology 15.1 (2023).
3. Qureshi Mahmud Shah. “Tribal Culture in Bangladesh”. Institute of Bangladesh Studies, Rajshahi University, Bangladesh (1984).
4. Ghani Abdul. “Traditional Medicine”. Jahangirnagar University, Savar, Dhaka, Bangladesh (1990).
5. Rashid KM, Khabiruddin and Md. And Hyder S. “Textbook of Community Medicine and Public Health”. 1st edn., RKH Publishers, Dhaka, Bangladesh (1992).
6. Hatip Al Khatib I, et al. “Determination of the effectiveness of components of the herbal medicine Toki Shakuyaku San and fractions of Angelica acutiloba in improving the scopolamine induced impairment of rat’s spatial cognition in eight armed radial maze test”. JPharmacol Sci 96 (2004): 33-41.
7. Ghani Abdul. “Medicinal Plants of Bangladesh: Chemical Constituents and Uses”. Asiatic Society of Bangladesh, Dhaka (1998).
8. Paterson GR, Neilson JB and Roland CG. “History of medicine”. Canadian Medical Association Journal 127.10 (1982): 948.
9. Alshammari TM. “Drug safety: The concept, inception and its importance in patients’ health”. Saudi Pharm J 24.4 (2016): 405-12.
10. Hanif M, et al. “Fatal renal failure caused by diethylene glycol in paracetamol elixir: the Bangladesh epidemic”. BMJ 311.6997 (1995): 88-91.
11. Baranwal M, Kaur A and Kumar R. “Challenges in utilizing diethylene glycol and ethylene glycol as excipient: A thorough overview”. Pharmaspire 15.1 (2023): 8-15.
12. Niazi SK. “Bridging the Regulatory Divide: A Dual-Pathway Framework Using SRA Approvals and AI Evaluation to Ensure Drug Quality in Developing Countries”. Pharmaceuticals (Basel) 18.7 (2025): 1024.
13. Cavaleri M, et al. “A roadmap for fostering timely regulatory and ethics approvals of international clinical trials in support of global health research systems”. Lancet Glob Health 13.4 (2025): e769-e777.
14. Lai Y, et al. “Recent advances in the translation of drug metabolism and pharmacokinetics science for drug discovery and development”. Acta Pharm Sin B 12.6 (2022): 2751-2777.
15. Hutt JA, et al. “Scientific and Regulatory Policy Committee Points to Consider: Nonclinical Research and Development of In Vivo Gene Therapy Products, Emphasizing Adeno-Associated Virus Vectors”. Toxicol Pathol 50.1 (2022): 118-146.
16. Kang HN, et al. “Regulatory challenges with biosimilars: an update from 20 countries”. Ann N Y Acad Sci 1491.1 (2021): 42-59.
17. Lexchin J. “Cost recovery by Health Canada and drug safety: a time-series analysis”. CMAJ Open 6.4 (2018): E471-E477.
18. Hjorth R, van Hove L and Wickson F. What can nanosafety learn from drug development? The feasibility of “safety by design”. Nanotoxicology 11.3 (2017): 305-312.
19. Tyner K and Sadrieh N. “Considerations when submitting nanotherapeutics to FDA/CDER for regulatory review”. Methods Mol Biol 697 (2011): 17-31.
20. Alban S. “The ‘precautionary principle’ as a guide for future drug development”. Eur J Clin Invest 35 Suppl 1.Suppl 1 (2005): 33-44.
21. Borgert CJ, Fuentes C and Burgoon LD. “Principles of dose-setting in toxicology studies: the importance of kinetics for ensuring human safety”. Arch Toxicol 95.12 (2021): 3651-3664.