

An Application of Nano Multilayers Found in the Wires of a Pacemaker

Maria Masood

Scientist, Germany

*Corresponding author: Maria Masood, Scientist, Germany.

Submitted: 19 September 2025 Accepted: 30 September 2025 Published: 07 October 2025

doi <https://doi.org/10.63620/MKJESER.2025.1019>

Citation: Masood, M. (2025). An Application of Nano Multilayers Found in the Wires of a Pacemaker. J of Electron Sci and Electrical Res 2(4), 01-05.

Abstract

This paper explores the application of nano multilayers-developed in our previous research – as conductive wires in cardiac pacemakers. Specifically, it proposes a novel design approach where pacemaker wires are embedded directly into the ventricular walls of the heart to optimize interaction with cardiomyocytes, the cells responsible for initiating and regulating heart beats. The study emphasizes the importance of precise placement and miniaturization of wires, advocating for diameters below 1mm using nanotechnology, and transfer matrix method (TMM) simulations. To meet biocompatibility and functionality requirements, the paper suggests using Titanium Nitride (TiN) as the conductive layer and silicon as the flexible, insulating substrate forming an Insulator-Metal-Insulator (IMI) structure. These materials are selected for their electrical conductivity, biocompatibility, flexibility and corrosion resistance, making them suitable for long-term implantation in cardiac tissues. This work aims to advance pacemaker design by improving integration with the heart's natural electrical system while ensuring safety and material compatibility.

Keywords: Pacemaker, Insulator-Metal-Insulator, Titanium Nitride, Pacemaker Wires, Diameter of Leads.

Introduction

The human heart stands as one of the most extraordinary and vital organs within the human body, entrusted with the indispensable role of continuously circulating blood. This circulation is critical for delivering oxygen and essential nutrients to every tissue and organ, thereby sustaining life and enabling the myriad physiological processes necessary for health and survival. The heart's ceaseless pumping action facilitates this intricate supply chain, ensuring that the body's cells receive the sustenance they require for metabolism, repair, and optimal function [1].

At the microscopic level, the heart's function depends on a specialized group of cells known as cardiomyocytes, or cardiac muscle cells. These cells are uniquely structured and highly specialized to meet the demanding mechanical requirements of the heart's pumping action. Unlike typical skeletal muscle cells, cardiomyocytes possess distinct structural and electrophysiological characteristics that en-

able them to contract in a highly coordinated and rhythmic manner. This contractile ability is precisely regulated by electrical impulses that propagate through cardiac tissue, orchestrating the heart's rhythmic beating.

Cardiomyocytes inherently respond to electrical stimulation by producing forceful contractions—a fundamental aspect of the heart's role as a pump that transforms electrical signals into mechanical work. The sarcomeric architecture of these cells, composed of interdigitating filaments of actin and myosin, facilitates contraction. Meanwhile, their rich mitochondrial content provides the energy necessary to sustain relentless activity over an individual's lifetime. Collectively, cardiomyocytes generate the pressure gradients required to move blood sequentially through the heart's chambers and into the arterial circulation.

A critical aspect of cardiac function is the synchronous contraction of cardiomyocytes, enabled by their electrical

coupling through specialized gap junctions. These junctions allow rapid propagation of action potentials across the myocardium, ensuring that the contraction wave efficiently spreads from the atria, the heart's upper chambers, to the ventricles, the more muscular lower chambers. Atrial contraction fills the ventricles with blood, which the ventricles then propel into systemic circulation through the aorta and into pulmonary circulation via the pulmonary artery. This seamless coordination is vital for maintaining optimal cardiac output and ensuring timely delivery of oxygenated blood to all body tissue [2,3].

The rhythmic beating of cardiomyocytes persists throughout an individual's life, often amounting to billions of contractions over several decades. This remarkable endurance reflects the specialized biology and finely tuned electrophysiological properties of these cells. The heartbeat is initiated by pacemaker cells located in the sinoatrial (SA) node, which generate spontaneous electrical impulses to set the pace for the myocardium, coordinating the precise temporal sequence of atrial and ventricular contractions and maintaining a regular heart rhythm.

In summary, the heart's critical role as a life-sustaining pump fundamentally relies on the unique properties and coordinated activity of cardiomyocytes. Their ability to convert electrical impulses into mechanical force, combined with synchronous contraction across the heart chambers, underpins the efficient, continuous blood flow necessary to meet metabolic demands. Understanding these cellular and physiological mechanisms provides essential insights into cardiac health and disease, guiding therapeutic strategies to manage conditions that disrupt this delicate balance.

Cardiomyocytes also possess intrinsic automaticity, enabling them to generate spontaneous electrical impulses. However, this pace-making ability is mainly localized to a specialized subset of cells in the SA node. These pacemaker cells initiate the heartbeat by producing regular electrical impulses, which travel through the heart's conduction system, ensuring coordinated contraction of atria and ventricles in a precise temporal sequence. Disruptions in this electrical system can lead to arrhythmias, abnormal heart rhythms that may impair cardiac function and pose serious health risks, including heart failure and sudden cardiac death.

The Role of Pacemakers in Cardiac Therapy

In cases where the heart's natural pacemaker function is

impaired—whether due to aging, disease, or injury—artificial pacemakers play a critical therapeutic role. These devices are designed to restore and maintain appropriate heart rhythms by delivering timed electrical impulses to the cardiac tissue. The typical artificial pacemaker consists of a pulse generator implanted subcutaneously near the shoulder or chest wall and insulated wires, known as leads, which extend into the heart chambers to stimulate the myocardium.

The placement and design of these leads are crucial for effective pacing. Leads are commonly inserted through the venous system into the right atrium or right ventricle, depending on the pacing requirements. They deliver electrical stimuli to cardiomyocytes to initiate contraction when the heart's intrinsic rhythm is too slow (bradycardia) or irregular. Moreover, pacemakers can provide anti-tachycardia pacing (ATP), which involves delivering rapid electrical pulses to interrupt episodes of tachycardia—abnormally fast heart rates that exceed 100 beats per minute. ATP acts as a physiological “brake,” resetting the heart's electrical activity and restoring normal rhythm.

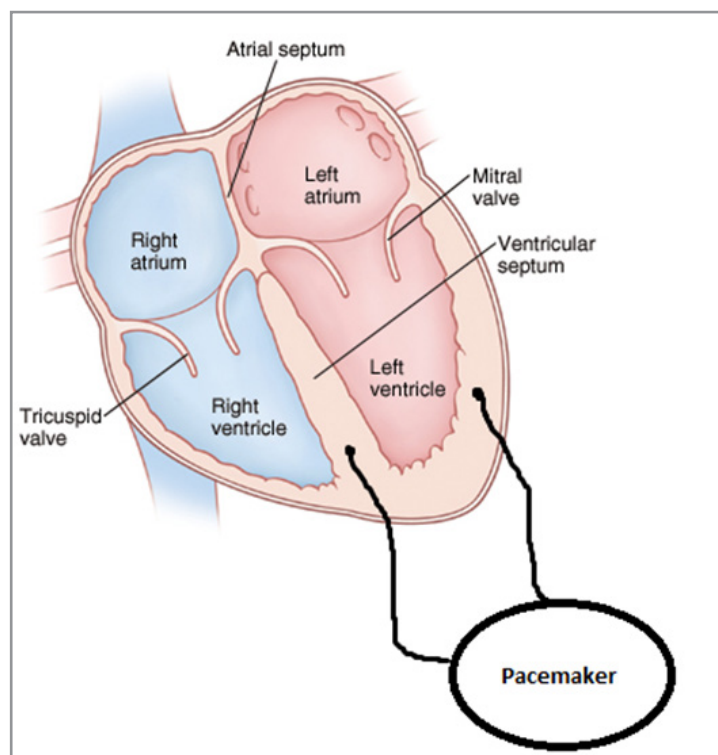
Challenges in Pacemaker Lead Placement

Despite decades of clinical success, conventional pacemaker lead systems have inherent limitations. Leads must be mechanically robust yet flexible enough to accommodate heart movements without causing tissue damage. Their diameter, often several millimeters, can cause discomfort, fibrosis, or even lead dislodgement over time. Additionally, leads may fail due to fracture, insulation breaches, or corrosion, necessitating replacement surgeries that pose further risk to patients.

The choice of lead placement site is also constrained by anatomical and functional factors. The right ventricular wall is often targeted for lead placement because it is accessible and provides effective pacing. However, its thickness is relatively small—approximately 3 to 4 millimeters—posing challenges for lead integration without causing injury or inflammation.

Proposed Advances in Lead Design and Placement

This paper introduces an innovative approach to pacemaker lead design that leverages recent advancements in nanotechnology and materials science. The core concept is to fabricate ultra-thin leads with diameters below 1 millimeter (~1000 nanometers), enabling more precise and less invasive insertion into the ventricular walls, particularly targeting the thinner right ventricular tissue.



Where to Place Pacemaker Leads in Heart

To achieve this level of miniaturization and performance, we propose the use of nano multilayer structures composed of bio-compatible materials. Inspired by prior research in selective emitters for thermophotovoltaic systems, multilayer films constructed via the transfer matrix method (TMM) allow for the precise tuning of optical, electrical, and mechanical properties at the nanometer scale. Our group has developed a Python-based simulation framework capable of modeling and optimizing such multilayer architectures [4-7].

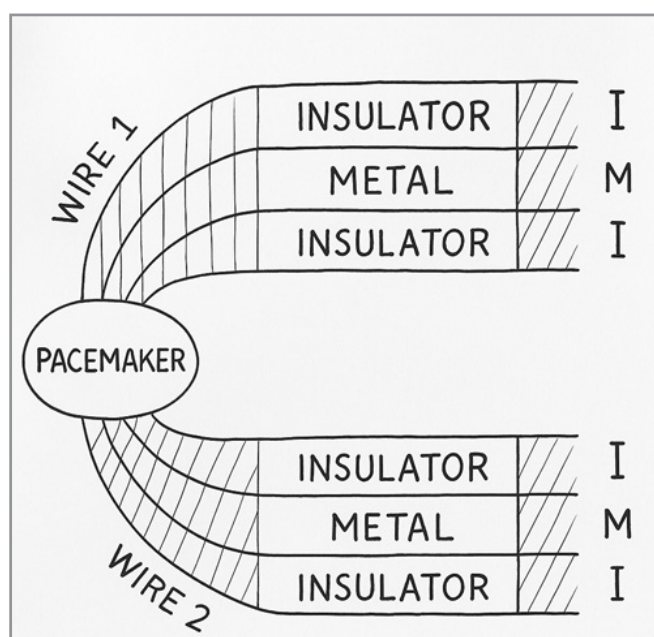
Materials and Biocompatibility Considerations

Biocompatibility is a critical requirement for all implantable cardiac devices. Materials must be non-toxic, corrosion-resistant, and mechanically compatible with the dynamic cardiac environ-

ment to minimize immune response and mechanical wear.

Titanium Nitride (TiN) is selected as the conductive layer in our multilayer design due to its excellent electrical conductivity, chemical stability, and well-documented biocompatibility. TiN coatings have been widely used in various medical implants and have shown to reduce allergic reactions while enhancing the durability of the device [8].

For the insulating layers, silicon is proposed owing to its established use in biomedical applications and its mechanical flexibility when fabricated in thin-film or flexible substrate form. Silicon-based insulating films provide the necessary electrical isolation while allowing the overall lead to maintain flexibility and conform to the cardiac tissue movements, reducing mechanical stress.



Idea Introduced in this Research

By combining TiN and silicon in an insulator–metal–insulator (IMI) multilayer configuration, the resulting structure can be engineered for optimal electrical performance, flexibility, and biocompatibility. The multilayer design facilitates miniaturization while maintaining mechanical integrity and functionality.

Significance and Future Directions

The proposed ultra-thin, flexible multilayer pacemaker leads have the potential to significantly improve patient outcomes by reducing complications related to lead implantation and longevity. Smaller lead diameters may allow for less invasive implantation techniques, decreased risk of lead displacement, and improved integration with cardiac tissue. Furthermore, the ability to precisely tune electrical properties at the nanoscale opens new avenues for responsive pacing therapies and adaptive electrical stimulation protocols.

Future research will focus on experimental validation of the proposed multilayer structures, including fabrication processes, mechanical testing, and in vitro biocompatibility studies. Integration with existing pacemaker systems and in vivo functional assessments will be crucial steps toward clinical translation.

Conclusion

This research article has explored an innovative approach to improving artificial pacemaker technology by addressing critical challenges associated with pacemaker lead design and placement. The human heart's intrinsic ability to rhythmically contract is governed by cardiomyocytes, driven by pacemaker cells within the sinoatrial node that generate electrical impulses. When this natural system falters due to disease or injury, artificial pacemakers become indispensable, restoring proper cardiac rhythms through electrical stimulation. Despite their lifesaving role, conventional pacemaker leads face several limitations, including bulkiness, mechanical rigidity, risk of tissue damage, and eventual device failure requiring risky replacement surgeries.

In response to these challenges, the study proposed and theoretically developed ultra-thin pacemaker leads using advanced nanotechnology, particularly multilayer thin-film structures, to enable safer and more effective lead placement, especially in the delicate right ventricular walls. By harnessing the transfer matrix method (TMM) for precise multilayer design and optimization, the research demonstrated how biocompatible materials such as Titanium Nitride (TiN) and silicon could be engineered in an insulator-metal-insulator (IMI) configuration. This approach allows for sub-millimeter lead diameters that maintain excellent electrical conductivity, mechanical flexibility, and biocompatibility essential for long-term implantation.

The selection of TiN as the conductive layer addresses multiple requirements simultaneously—its high electrical conductivity ensures efficient pacing signals, its chemical stability and corrosion resistance minimize degradation in the dynamic cardiac environment, and its proven biocompatibility reduces adverse immune reactions. Silicon, chosen for the insulating layers, complements TiN by providing flexibility and electrical isolation while adapting well to the mechanical strains imposed by cardiac motion. Together, the IMI multilayer structure enables

a novel pacemaker lead design that potentially overcomes the mechanical and biological shortcomings of traditional leads.

The implications of this research are significant. Miniaturized leads reduce the invasiveness of implantation procedures, thereby lowering patient trauma and the risk of lead dislodgement or fibrosis that often complicate long-term pacemaker use. Moreover, the ability to finely tune electrical and mechanical properties at the nanoscale opens pathways for developing adaptive pacing therapies. These could offer more physiological pacing strategies that respond dynamically to patient needs, improving overall cardiac function and quality of life.

While this study primarily focused on the theoretical design and simulation of multilayer structures, it lays a robust foundation for subsequent experimental work. Future research must emphasize the fabrication of these nanoscale leads using state-of-the-art thin-film deposition techniques. Mechanical testing under simulated cardiac cycles will be essential to validate durability and flexibility claims. Additionally, thorough in vitro biocompatibility studies will ensure that the materials and designs do not elicit harmful immune responses or cytotoxicity.

Integration with existing pacemaker hardware and electronic control systems represents another critical step. This includes evaluating the electrical interface between the miniaturized leads and the pulse generator, ensuring that signal transmission remains reliable. Eventually, preclinical in vivo testing in appropriate animal models will be needed to assess functional pacing efficacy, tissue integration, and long-term safety before clinical trials can be considered.

In summary, this research advances the field of cardiac therapy by proposing a cutting-edge nanomaterials-based solution to longstanding issues in pacemaker lead technology. By blending the disciplines of materials science, nanotechnology, and biomedical engineering, it opens new horizons for creating safer, more effective, and more patient-friendly cardiac pacing devices. The use of TiN and silicon multilayers in an IMI configuration provides a versatile platform that could revolutionize pacemaker design, reduce complications and enhance the therapeutic outcomes for millions of patients worldwide.

Ultimately, the successful realization of these ultra-thin, flexible leads could not only improve the durability and comfort of pacemakers but also catalyze further innovations in cardiac electrophysiology devices. The potential for integrating smart, responsive electronics within these nanoscale architectures suggests a future where cardiac pacing is more personalized, adaptive, and closely aligned with the heart's natural physiology. This paradigm shift promises to improve survival rates and quality of life for patients suffering from cardiac arrhythmias and conduction system disorders, marking a transformative milestone in cardiovascular medicine.

Funding Declaration

There was no funding provided or needed for this manuscript.

Ethics, Consent to Participate, And Consent to Publish Declarations

Not Applicable.

Reference

1. Hall, J. E. (2016). Textbook of medical physiology (13th ed.). Elsevier.
2. Makita, Y., Nakano, Y., Oda, N., Suenari, K., Sairaku, A., Kajihara, K., ... & Kihara, Y. (2012). Use of preprocedural multidetector computed tomography to decrease atrial fibrillation recurrence following extensive encircling circumferential pulmonary vein isolation. *Journal of cardiology*, 60(3), 236-241.
3. Katz, A. M. (2010). *Physiology of the heart* (5th ed.). Lippincott Williams & Wilkins. (Assumed 5th ed. and full name "A. M. Katz"; please correct if different.)
4. Masood, M. (2025). Trends in metal-insulator-metal-insulator-metal multilayers built using Fabry–Perot cavities. *Journal of Electronics Science and Electrical Research*, 12(1), xx–xx. (Volume, issue, and page numbers should be confirmed.)
5. Masood, M. (2025). Extracting optical parameters of plasmonic titanium nitride using Ref FIT. *Journal of Electronics Science and Electrical Research*, 12(2), xx–xx.
6. Masood, M. (2025). Fabry–Perot cavity-based selective emitters intended for medium. *International Journal of Quantum Technologies*, 6(1), xx–xx.
7. Masood, M. (2025). Validation of transfer matrix method simulations with experiment. *Journal of Electronics Science and Electrical Research*, 12(3), xx–xx.
8. Gobby. (2019, March 6). Orthopedic implants: Coating with TiN. *Biomedical Journal of Scientific & Technical Research*. <https://biomedres.us/fulltexts/BJSTR.MS.ID.002786.php>