

MICROFLUIDIC INTEGRATION: REVOLUTIONIZING LAB-ON-A-CHIP TECHNOLOGY FOR COMPACT AND EFFICIENT BIOMEDICAL ANALYSIS

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Abstract: A lab-on-a-chip (LOC) device is a miniaturized platform that integrates various laboratory functions onto a single chip typically only a few square centimeters in size. These devices are designed to perform a wide range of biochemical and biomedical analyses, often with small sample volumes and rapid turnaround times. Key components of a lab-on-a-chip device may include microfluidic channels, where fluids such as blood or chemical reagents are manipulated and processed; sensors for detecting analytes or biological molecules; and actuators for controlling fluid flow or other operations. Lab-on-a-chip devices offer several advantages over traditional laboratory techniques, including reduced sample and reagent consumption, faster analysis times, portability, and potential for automation. They have applications in fields such as medical diagnostics, environmental monitoring, drug discovery, and biotechnology.

IndexTerms - Biosensor, Lab-on-Chip, MEMS, Microfluidics, PCB

I.INTRODUCTION

Lab-on-a-Chip technology refers to methods that carry out different laboratory tasks on a smaller scale, such as chemical emulsion and analysis on a single chip, which results in a portable and handheld device. Stated differently, LoC is a tool that can measure one or more laboratory functions at the chip level. This chip is between a few millimetres and several square centimetres in size [1]. The integration of fluidics, electronics, optics, and biosensors is the main element of LoC [2]. The main motive of LoC is the need for the state- of- art pathological analysis on the- go. LoCs prove to be useful for chancing the styles for the early stage opinion of deadly and habitual conditions. Due to the appearance of advanced technologies similar as MEMS,NEMS, the integration of large number of interdisciplinary modules on a single chip is possible [3].

fig 1 represents the various steps involved in Lab-on-Chip processing. The first step in the LoC procedure is to gather a physiological sample, from which a specific analyte or biomarker is created. The transducer will operate on the analyte electrically, electromechanically, optically, or mechanically, depending on the biological function. The coming step involves counting, sorting and modification of the transducer affair is performed according to the operation. Eventually, the amplified sample is reused using microelectronics ways. Current trend shows the growth of research in this area .In numerous universities across the world, nu merous groups are formed that are earmarking their exploration in this area .For illustration, memoirs in University of Twente, Mina Med in Germany, and Nanobe in Finland [3] are some of the groups. Their main motive is to understand naonofluidics and nanosensing, to connect micro/ nanoeng. with biomedical and life wisdom fields, to develop new micro and nano technologies for LOC, and to demonstrate new LOC operations.



fig 1. schematic representation of lab-on-chip process

II.LITERATURE SURVEY

The study [4] (Duffy, David C., et al) presents a process that allows microfluidic devices to be designed, fabricated, and se aledin less than 24 hours using the elastomeric material poly(dimethylsiloxane) (PDMS). Using a CAD programme, a network of microfluidic channels (wider than 20 µm) is created. A high-resolution printer turns this design into a transparency, which is then used as a mask in photolithography to make a positive relief photoresist master. When PDMS is cast against the master, a polymeric replica with a channel network is produced. This replica's surface, together with the surface of a flat PDMS slab, is oxidised in an oxygen plasma. When these oxidised surfaces come into conformal contact, they seal firmly and permanently. An further benefit of oxidising the PDMS is that it creates channels with negatively charged walls that allow electroosmotic pumping and are easily filled with high surface energy liquids, particularly water. Using this fast prototyping method, a miniature capillary electrophoresis device has been built to assess the performance of microfluidic systems. Using this method, aqueous solutions containing amino acids, charge ladders of positively and negatively charged proteins, and DNA fragments were separated with a resolution similar to that achieved through the use of fused silica capillaries.

The paper [5] (C. D. Chin, V. Linder et al and S. K. Sia et al) 2012 discusses the advancements and applications of lab- on- a- chip (LOC) technology. It provides insights into the miniaturization and integration of laboratory functions onto a single chip, enabling efficient and high-throughput analysis. The authors likely explore microfabrication techniques utilized in creating these devices, along with their potential applications in various fields such as biology, chemistry, and medicine. The paper may also discuss the challenges and future directions in the development and commercialization of lab-on-a-chip devices. Overall, it serves as a valuable resource for researchers and professionals interested in the rapidly evolving field of microfluidics and miniaturized analytical systems.

The article [6] (Abill Robert, Elisha Blessing et al and Hubert Klaus et al) 2024 likely explores innovative methods and technologies for swiftly detecting and identifying plant pathogenic microorganisms. Lab-on-a-chip systems, which miniaturize laboratory processes onto a single chip, are probably a focal point. The authors likely discuss the advantages of these systems, such as rapidity and portability, in contrast to conventional diagnostic methods. Additionally, the article may delve into the specific techniques, sensors, and detection mechanisms employed within these lab-on-a-chip systems to achieve accurate and efficient diagnosis of plant diseases. Overall, the aim is likely to highlight advancements in plant pathology diagnostics and the potential of lab-on-a-chip technologies in agricultural settings.

III.OBJECTIVE

Lab-on-a-chip (LOC) devices represent a transformative approach to laboratory analysis by consolidating multiple laboratory functions onto a single chip. The primary objective of these devices is to miniaturize and integrate complex laboratory processes, typically performed in conventional benchtop settings, onto a compact and portable platform.

By achieving miniaturization, LOC devices enable the analysis of samples using significantly reduced volumes of reagents and samples, thereby conserving resources and minimizing waste. Moreover, the integration of various functions, suchas sample preparation, mixing, reaction, separation, and detection, onto a single chip streamlines workflows and reduces processing time. Automation further enhances the efficiency of LOC devices by minimizing human intervention and improving reproducibility. These features make LOC devices particularly well-suited for point-of-care testing (POCT) applications, where rapid analysis at the site of need is essential for timely diagnosis and treatment decisions. Additionally, the versatility of LOC devices allows them to be adapted for a wide range of applications, including biomedical diagnostics, environmental monitoring, drug discovery, and food safety testing. Overall, the overarching goal of lab-on-a-chip technology is to revolutionize laboratory practices by providing compact, integrated, and user- friendly platforms for efficient and cost-effective analysis across diverse fields and settings.

IV.CHIP MATERIALS AND FABRICATION TECHNOLOGIES

Photolithography forms the foundation of most LOC fabrication methods. Since these highly developed technologies were directly descended from the manufacture of semiconductors, the majority of procedures were initially in silicon. In response to demands for, among other things, specific optical characteristics, bio- or chemical compatibility, lower production costs, and faster

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prototyping, new processes have been developed, including glass, ceramics, and metal etching, deposition and bonding, polydimethylsiloxane (PDMS) processing (e.g., soft lithography), Off-stoichiometry thiol-ene polymers (OSTEmer) processing ,thick-film- and stereolithography-based 3D printing,[7] as well as fast replication methods via electroplating, injection moulding , and embossing. A straightforward technique for creating PDMS microfluidic devices was developed in response to the need for low-cost, straightforward LOC prototyping: ESCARGOT (Embedded SCAffold RemovinG Open Technology) [8]. This method uses a dissolvable scaffold (produced, for example, via 3D printing) [9] to create microfluidic channels within a single block of PDMS. Moreover, the boundaries between nanotechnology, precision engineering, and lithography-based microsystem technology are increasingly being crossed by the LOC field. Printing is regarded as a mature yet well-established technique for chip manufacture quick prototyping [10].

These distinguishing features—commercially available substrates with integrated electronics, sensors, and actuators; lowcost, disposable devices with a high potential for commercialization—make the development of LOC devices using printed circuit board(PCB) substrates an intriguing alternative [11]. A few benefits of PCB technology are as follows:

- PCB-based circuit design is very flexible and maybe customised to meet certain needs [12].
- The integration of electronic and sensor modules on the same platform is made possible by PCB technology, which reduces device size without sacrificing detection accuracy.
- The large-scale, economically viable production of PCB-based detecting devices is made possible by the well- established and standardised PCB manufacturing process.
- The development of wearable detecting devices has been fueled by the advancement of flexible PCB technology. Consequently, a plethora of publications on the application of Lab-on-PCB to many biomedical domains have been published during the past ten years.
- PCBs can be used with wet deposition techniques, enabling the creation of sensors with cutting-edge nanomaterials like graphene [13].

V.METHODOLOGY

Sample Preparation and Introduction:-

Sample Handling:- The device facilitates the handling of raw samples, such as blood, saliva, or environmental fluids, by integrating mechanisms for filtration, concentration, and extraction.

Preparation:- It prepares the sample for analysis by separating target analytes from interfering substances, removing particulates, and adjusting the sample's chemical composition if necessary.

Introduction to Microfluidic System: The prepared sample is introduced into the microfluidic system, where it undergoes further processing and analysis.

• Analysis and Detection:-

Microfluidic Manipulation:- Within the microfluidic channels, the sample is manipulated using precise fluid control mechanisms, such as pumps, valves, and mixers, to perform various analytical functions.

Analyte Detection: - The device incorporates sensors, detectors, or assay platforms to detect and quantify specific analytes of interest. This may involve optical, electrochemical, or biological detection methods, depending on the application.

Chemical and Biological Reactions: Chemical reactions, such as PCR amplification or enzymatic assays, as well as biological reactions, such as antigen-antibody interactions, can be conducted within the microfluidic system to generate measurable signals indicative of the analyte's presence or concentration.

• Data Analysis and Interpretation: -

Signal Processing:- The device processes the raw data obtained from the detection step, which may involve amplification, filtering, or signal conversion to enhance the signal-to-noise ratio and improve the accuracy of the analysis.

Data Interpretation:- Algorithms or analytical models are applied to interpret the processed data and extract relevant information, such as concentration levels, molecular profiles, or diagnostic outcomes.

Result Presentation:- The final results are presented in a user-friendly format, such as numerical values, graphical plots, or diagnostic reports, allowing for easy interpretation and decision-making by end-users, such as researchers, clinicians, or environmental scientists.

VI.FUTURE SCOPE

Microfluidic LoCs are commonly believed to have the potential to completely transform the biological profession and to significantly improve the healthcare industry. However, this LoC technology still seems like a pipe dream, particularly in underdeveloped nations. Low-resource regions, including developing nations, still need to work on improving the economic model that drives the production of LoC devices and raising public awareness of how effectively they may be used. Many infectious diseases are there such as malaria, HIV and AIDS, measles, TB, lower respiratory conditions and so on that require timely diagnosis and treatment so as to reduce the mortality rate. Now-a-days, Swine Flu, Zika Virus and Ebola Virus are killing thousands of people and causing the risk of epidemic.

Because LoCs diagnostics can provide a diagnosis in real time, they are vital for all of these disorders and can significantly enhance the landscape of disease management. Moreover, this technology can proven to be useful for finding a novel way to treat central

nervous system disorders such as Parkinson's disease and spinal cord injury with the help of extracting sufficient cerebrospinal fluid needed to perform conventional assays. The potential of this technology can further be explored for autoimmune joint diseases like rheumatoid arthritis.

VII.CONCLUSION

The survey results indicate that there is a broad range of uses for LoC devices. The primary areas of research in this field include human diagnostics, chemical synthesis, and DNA analysis. They can be used in different areas such as in diagnostics, bioanalysis, and biosensing, for environmental monitoring including testing of water and food quality, for testing of different drugs, in pharmaceutics and so on. Their advantages include portability, modularity, reconfigurability, and low power consumption. Due to their various capabilities, it becomes now possible to detect the chronic and life threatening diseases like cancer, TB etc. in the early stages leading to the reduction in mortality rate. They automate the laboratory processes like sample transport, dispensing and mixing and have the capability of reducing the time of laboratory tests to a great extent. Due to their various capabilities, it becomes now possible to detect. TB etc. in the early stages leading to the reduction and life threatening diseases like cancer, TB etc. in the reduction in mortality rate. They automate the laboratory processes like sample transport, dispensing and mixing and have the capability of reducing the time of laboratory tests to a great extent. Due to their various capabilities, it becomes now possible to detect the chronic and life threatening diseases like cancer, TB etc. in the early stages leading to the reduction in mortalityrate. Because LoCs devices offer quick and accurate diagnosis results, they are appropriate for point-of-care diagnostics. Till today, Lab-on-Chip for monitoring the lithium level in maniac patients is successfully tested.

REFERENCES

[1] Lisa R. Volpatti and Ali K. Yetisen, "Commercialization of microfluidic devices", Trends inbiotechnology, vol. 32, no. 7, (2014), pp. 347-350.

[2] A. T. Giannitsis and M. Min, "Usage of microfluidic lab-on-chips in biomedicine", Proceedings of 12th Biennial Baltic Electronics Conference, Tallinn, (2010), pp. 249-252

[3] "Microfluidics, Lab-On-Chip", http://lab-on-chip.gene quantification.info, accessed 29 June 2016.

[4] Duffy, David C., et al. "Rapid prototyping of microfluidic systems in poly(dimethylsiloxane)." Analytical chemistry 70.23 (1998): 4974-4984.

[5] C. D. Chin, V. Linder and S. K. Sia, Lab Chip, 2012, 12, 2118-2134

[6] Abill Robert; Elisha Blessing; Hubert Klaus; Advanced Diagnostic Techniques for Rapid Detection and Identification of Plant Pathogenic Microorganisms Using Lab-on-a Chip Systems, 2024.

[7] Gonzalez, Gustavo; Chiappone, Annalisa; Dietlikee, Kurt; Pirri, Fabrizio; Roppolo, Ignazio (2020). "Fabrication and Functionalization of 3D Printed Polydimethylsiloxane-Based Microfluidic Devices Obtained through Digital Light Processing". Advanced Materials Technologies. 5 (9): 2000374. doi:10.1002/admt.202000374. S2CID 225360332.

[8] Saggiomo, V.; Velders, H. A. (Jul 2015). "Simple 3D Printed Scaffold-Removal Method for the Fabrication of Intricate Microfluidic Devices". Advanced Science. 2 (8): X. doi:10.1002/advs.201500125. PMC 5115388. PMID 27709002.

[9] Vittorio Saggiomo (17 July 2015). "Simple fabrication of complex microfluidic devices(ESCARGOT)". Archived from the original on 2021-12-22 – via YouTube.

[10] Loo J, Ho A, Turner A, Mak WC (2019). "Integrated Printed Microfluidic Biosensors". Trends inBiotechnology. 37 (10): 1104–1120. doi:10.1016/j.tibtech.2019.03.009. hdl:1826/15985. PMID 30992149.S2CID 119536401.

[11] Perdigones, Francisco (2021). "Lab-on-PCB and Flow Driving: A Critical Review". Micromachines.12(2):175. doi:10.3390/mi12020175. PMC 7916810. PMID 33578984.

[12] Zhao, Wenhao; Tian, Shulin; Huang, Lei; Liu, Ke; Dong, Lijuan (2020). "The review of Lab-on-PCB forbiomedical application". Electrophoresis. 41 (16–17): 1433–1445. doi:10.1002/elps.201900444. PMID 31945803. S2CID 210699552.

[13] Fenech-Salerno, Benji; Holicky, Martin; Yao, Chengning; Cass, Anthony E. G.; Torrisi, Felice (2023). "Asprayed graphene transistor platform for rapid and low-cost chemical sensing". Nanoscale. 15 (7): 3243– 3254.doi:10.1039/d2nr05838c. hdl:10044/1/102808. PMID 36723120. S2CID 2562617.

[14] C.D. Chin, Vincent Linder and S.K. Samuel. "Commercialization of microfluidic point-of-care diagnostic devices", Lab on a Chip Journal, vol. 12, no. 12, (2012), pp. 2118-2134.

[15] Krishna, Katla Sai, Yuehao Li, Shuning Li, and Challa SSR Kumar, "Lab-on-a-chip synthesis of inorganic nanomaterials and quantum dots for biomedical applications", Advanced Drug Delivery Reviews Journal, vol. 65, no. 11, (2013), pp. 1470-1495.

[16] "Lab-on-a-Chip' Technology Cuts Costs of Lab Tests for HIV, Lyme Disease, Other Diseases", http://news.rutgers.edu/research-news/%E2%80%98lab-chip%E2%80%99-technology-cuts-costs-labtests- hiv-lyme-disease-other-diseases/20150910#.VtWeUPI97IU, accessed 15 June 2016.

[17] A.M. Cabibbe, P. Miotto, R. Moure, F. Alcaide, S. Feuerriegel, G. Pozzi, V. Nikolayevskyy, F. Drobniewski, S. Niemann, K. Reither and D.M. Cirillo, "Lab-on-chip-based platform for fast molecular diagnosis of multidrug-resistant tuberculosis", Journal of clinical microbiology, vol. 53, no. 12, (2015), pp. 3876-3880.

[18] E Primiceri, MS Chiriaco, R Rinaldi and G Maruccio, "Cell chips as new tools for cell biology – results, perspectives and opportunities", Lab on a Chip Journal, vol. 13, no. 19, (2013), pp. 3789-3802.

[19] E. T. Carlen and A. van den Berg, "Labs-on-a-chip and nanosensors for medical applications and life sciences", Proceedings of IEEE International Electron Devices Meeting, Washington, DC, (2013), December 9-11, pp. 8.7.1-8.7.4.

[20] K. Grenier, D. Dubuc, T. Chen, T. Chrétiennot, M. Poupot and J. J. Fournié, "Microfluidic on-chip for biomedical applications", Proceedings of IEEE Bipolar/BiCMOS Circuits and Technology Meeting, Atlanta, GA, (2011), October 9-11, pp. 129-132.

[21] Foudeh, Amir M., Tohid Fatanat Didar, Teodor Veres, and Maryam Tabrizian, "Microfluidic designs and techniques using labon-a-chip devices for pathogen detection for point-of-care diagnostics", Lab on a Chip Journal, vol. 12, no. 18, (2012), pp. 3249-3266.

[22] J. Auerswald, S. Berchtold, J.M. Diserens, M.A. Gijs, Y.H. Jin, H.F. Knapp, Y. Leterrier, J.A.E Manson, G. Suarez, and G. Voirin, "Lab-on-a-chip for Analysis and Diagnostics: Application to Multiplexed Detection of Antibiotics in Milk", In Nanosystems Design and Technology, Springer US, (2009), pp. 117-142.

[23] BL Ziober, MG Mauk, EM Falls, Z Chen, AF Ziober and Haim H. Bau, "Lab on a chip for oral cancer screening and diagnosis", Head & Neck Journal, vol. 30, no. 1, (2008), pp. 111-121.