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Review Paper

Microneedle-Based Drug Delivery System: A Painless and Patient-Friendly Approach

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ABSTRACT

Transdermal drug delivery systems offer a non-invasive alternative to oral and injectable routes; however, their clinical application is significantly limited by the barrier function of the stratum corneum. Microneedle-based drug delivery systems have emerged as an innovative strategy to overcome this limitation by creating transient microchannels in the skin, thereby enhancing drug permeation without stimulating deep dermal pain receptors. Microneedles are micron-scale projections, typically 150–1500 μm in length, designed to penetrate the outer skin layers safely and effectively. This review aims to summarize the design principles, classification (solid, coated, dissolving, hollow, and hydrogel-forming), fabrication materials, and mechanisms of drug release associated with microneedle systems. Recent advancements in biodegradable polymers, microfabrication techniques, and sustained-release formulations have expanded their applications in vaccination, chronic pain management, insulin delivery, dermatological therapy, and cosmetic treatments. Compared to conventional hypodermic injections, microneedles offer minimal pain, improved patient compliance, reduced systemic side effects, and potential for self-administration. Despite promising clinical outcomes, challenges related to large-scale manufacturing, regulatory approval, and long-term safety evaluation remain. Continued research and technological innovation are expected to further enhance the translational potential of microneedle-based therapeutics in modern healthcare.

INTRODUCTION

The skin acts as a highly efficient protective barrier against environmental insults and chemical exposure. While this function is physiologically

essential, it limits the transdermal absorption of most drug molecules. Conventional hypodermic injections provide effective systemic delivery but are associated with discomfort, needle anxiety, and reduced long-term adherence. Conversely, topical

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formulations often fail to achieve therapeutic drug concentrations due to limited permeability across the stratum corneum.

Microneedle-based systems were developed to overcome these limitations. These devices consist of arrays of microscopic projections typically ranging from 150–1500 μm in length. Their dimensions permit penetration through the outermost skin layer while avoiding deeper vascular and neural structures, thereby minimizing pain and tissue trauma. This approach enhances dermal drug transport and improves patient acceptance.

The skin is composed of three principal layers: epidermis, dermis, and hypodermis. The epidermis contains multiple strata, with the stratum corneum serving as the primary permeability barrier. The dermis is a connective tissue layer containing blood vessels, nerve endings, and appendages that support thermoregulation and sensation. The hypodermis consists largely of adipose tissue and provides insulation and structural support. Knowledge of this layered structure is fundamental in designing effective microneedle systems.

Skin Anatomy ⁽⁵⁾

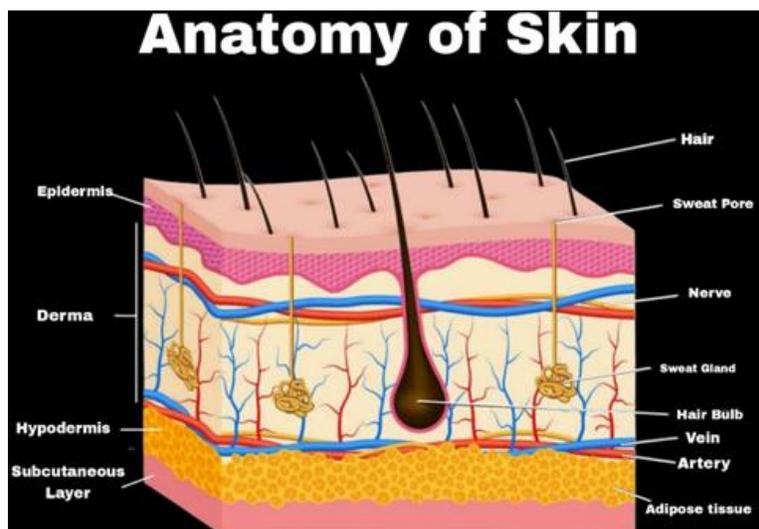


Figure 1- Anatomy of skin

Conventional Drug Delivery for Pain Management ⁽⁶⁻⁸⁾

Pain is commonly managed using oral, parenteral, topical, and transdermal drug delivery approaches. Each route offers certain advantages; however, significant limitations remain.

Oral administration of analgesics and anti-inflammatory agents is widely preferred because it is convenient and non-invasive. Despite this practicality, oral therapy often requires repeated dosing due to short duration of action. Additionally, delayed onset of relief,

gastrointestinal irritation, variable absorption, and reduced bioavailability may compromise therapeutic effectiveness. Long-term use of certain oral analgesics is also associated with systemic adverse effects.

Parenteral administration provides rapid onset of action and precise dosing, making it useful in acute pain conditions. However, it is not ideal for routine or long-term management. Repeated injections increase the risk of infection, tissue damage, and local inflammation at the injection site. Moreover, needle-related anxiety, discomfort, and fear

significantly reduce patient acceptance and adherence to therapy.

Topical drug delivery systems were introduced to minimize systemic side effects and improve safety. These formulations allow localized treatment and are generally well tolerated. Nevertheless, their clinical effectiveness is frequently limited by poor drug penetration across the skin barrier, necessitating frequent reapplication. Prolonged use may also cause local irritation or sensitivity.

Transdermal drug delivery systems represent a more advanced alternative. By facilitating controlled drug release across the skin, they provide sustained therapeutic levels while avoiding first-pass hepatic metabolism. Compared with oral and injectable routes, transdermal systems can reduce systemic toxicity, improve bioavailability, and enhance patient compliance. However, their performance remains constrained by the barrier properties of the stratum corneum, which restrict the delivery of many therapeutic agents.

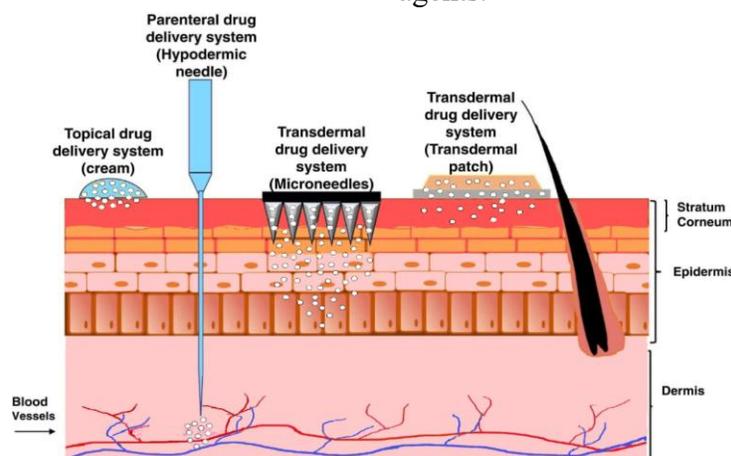


figure 2 – conventional drug delivery

Types of Microneedles ^(9,10)

1. Solid Microneedles: Used to create microchannels followed by topical drug application.
2. Coated Microneedles: Drug-coated surfaces allow rapid release upon insertion.

3. Dissolving Microneedles: Fabricated from biodegradable polymers that dissolve after insertion.
4. Hollow Microneedles: Contain a lumen for controlled liquid drug infusion.
5. Hydrogel-Forming Microneedles: Swell after insertion and enable sustained drug diffusion.

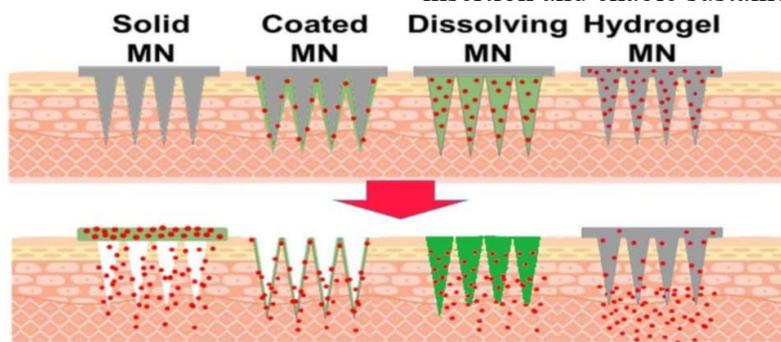


Figure 3 – Types of microneedles

Mechanism of Drug Delivery ⁽¹¹⁾

Microneedles function by mechanically breaching the stratum corneum to create temporary aqueous

pathways. Drug molecules subsequently diffuse into the epidermis or dermis, depending on formulation design. The microchannels typically

reseal within hours, reducing infection risk and preserving skin integrity.

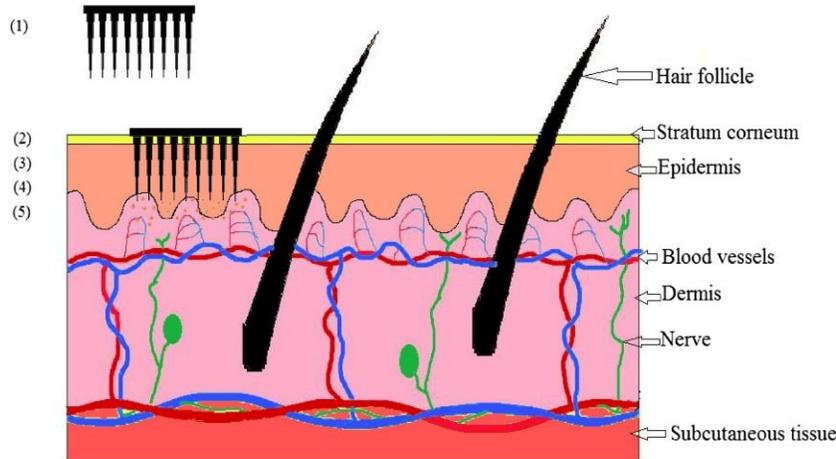


Figure 4 – Mechanism of medicine delivery

- (1) microneedle device with drug solution,
- (2) device inserted into the skin,
- (3) temporary mechanical disruption of the skin,
- (4) releasing the drug in epidermis,
- (5) transport of drugs to site of action

Microneedle systems have demonstrated potential in osteoarthritis, rheumatoid arthritis, neuropathic pain, atopic dermatitis, and psoriatic arthritis. Localized dermal delivery reduces systemic exposure and enhances therapeutic precision. Sustained-release designs further improve long-term symptom control.

Applications in Chronic Pain-Associated Disorders (12-15)

Schematic representation of mechanism of action of microneedles

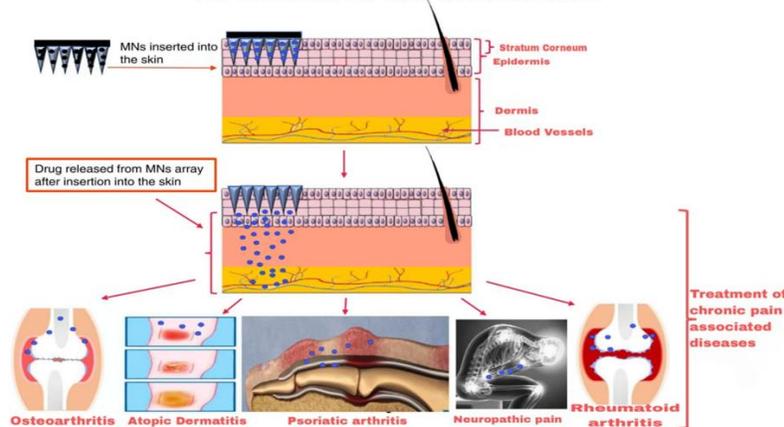


Figure 5 – microneedles in chronic pain associated disease

Materials Used in Fabrication (16,17)

Common fabrication materials include biodegradable polymers (PVA, PVP, PLA, PLGA), carbohydrates, metals, silicon, ceramics,

and glass. Material selection influences mechanical strength, biocompatibility, drug loading capacity, and release kinetics.

Approved microneedle products



The first microneedle product was derma comber. request and are approved for medical and numerous microneedle products are coming in the ornamental use.

Sr. No.	PRODUCT NAME	COMPANY	DESCRIPTION	USES
1.	Dermaroller®	Dermaroller Germany, White Lotus	A small roller with solid or essence-coated microneedles, ranging from 0.2 to 2.5 mm in length.	Improves skin texture and appearance.
2.	Soluvia®	Sanofi Pasteur Europe	Hollow microneedle attached to a syringe-like device for precise delivery.	Administers influenza vaccination.
3.	MicroHyal®	CosMed	Dissolving microneedle patch containing hyaluronic acid.	Reduces wrinkles and promotes skin hydration.
4.	LiteClear®	Nanomed Skincare	Solid silicon microneedles used for pre-treatment before applying topical medicines.	Helps in acne management and enhances topical treatment efficacy.

Advantages ^(18,19)

Microneedles offer minimal pain, improved patient compliance, avoidance of first-pass metabolism, reduced systemic side effects, and suitability for self-administration.

Future Prospects

Future research focuses on stimuli-responsive microneedles, wearable integration, and personalized drug delivery systems. Advances in microfabrication and biomaterials are expected to enhance scalability and clinical translation.

CONCLUSION

Microneedle-based drug delivery represents a progressive advancement in transdermal therapeutics. By transiently disrupting the skin barrier without causing significant discomfort, microneedles provide a patient-centered and minimally invasive alternative to conventional injections. Continued innovation in material science and device engineering is likely to expand their clinical applications in the coming years.

REFERENCES

1. Ita K: Transdermal delivery of drugs with microneedles—potential and challenges. *Pharmaceutics* 2015; 7: 90-105.
2. Bora P, Kumar L and Bansal AK: Microneedle technology for advanced drug delivery: evolving vistas. *Critical Reviews in Therapeutic Drug Carrier Systems* 2008; 25: 427-465.
3. Amarnani R and Shende P: Microneedles in diagnostic, treatment and theranostics: an advancement in minimally invasive delivery system. *Biomedicine and Pharmacotherapy* 2018; 102: 123-133.
4. Waghule T, Singhvi G, Dubey SK, Pandey MM, Gupta G, Singh M and Dua K: Microneedles: a smart approach and increasing potential for transdermal drug delivery system. *Biomedicine and Pharmacotherapy* 2019; 109: 1249-1258.
5. Norlén LPO: The skin barrier—structure and physical function. Ph.D. Dissertation, Karolinska Institute, Stockholm, Sweden 1999.



6. Puntillo F, Giglio M and Varrassi G: The routes of administration for acute postoperative pain medication. *Pain Therapy* 2021; 10: 909-925.
7. Marty FM, Lowry CM, Rodríguez M, Milner DA Jr, Pieciak WS and Sinha A: Parenteral administration of ivermectin in a patient with disseminated strongyloidiasis. *Clinical Infectious Diseases* 2005; 41: e5-e8. (*Students must verify correct article if different one intended.*)
8. Rizwan M, Aqil M, Talegaonkar S, Azeem A, Sultana Y and Ali A: Enhanced transdermal drug delivery techniques: an extensive review of patents. *Recent Patents on Drug Delivery and Formulation* 2009; 3: 105-124.
9. Yang D, Chen M, Sun Y, Jin Y, Lu C, Pan X, Quan G and Wu C: Microneedle-mediated transdermal drug delivery for treating diverse skin diseases. *Acta Biomaterialia* 2021; 121: 119-133.
10. Zhang Y, Brown K, Siebenaler K, Determan A, Dohmeier D and Hansen K: Development of lidocaine-coated microneedle product for rapid, safe and prolonged local analgesic action. *Pharmaceutical Research* 2012; 29: 170-177.
11. Prausnitz MR and Langer R: Transdermal drug delivery. *Advanced Drug Delivery Reviews* 2008; 60: 1-2.
12. Ita K: Current status of microneedles for transdermal drug delivery. *Current Medical Delivery* 2015; 12: 138-150.
13. Gupta J, Gill HS, Andrews SN and Prausnitz MR: Kinetics of skin resealing after insertion of microneedles in human subjects. *European Journal of Pharmaceutics and Biopharmaceutics* 2011; 79: 618-625.
14. Nguyen HX and Banga AK: Delivery of biologics using microneedles. *Journal of Controlled Release* 2019; 301: 1-17.
15. Kim YC, Park JH and Prausnitz MR: Microneedles for drug and vaccine delivery. *Advanced Drug Delivery Reviews* 2012; 64: 1547-1568.
16. Ita K: Recent advances and future prospects of topical and transdermal drug delivery systems. *Frontiers in Drug Delivery* 2022; 2: 1-15.
17. Sharma D: Microneedles: an approach in transdermal drug delivery—a review. *International Journal of Pharmaceutical Sciences and Research* 2017; 8: 456-465.
18. Singh A and Yadav S: Microneedling: advances and widening horizons. *Indian Dermatology Online Journal* 2016; 7: 244-254.
19. Donnelly RF, Singh TRR and Woolfson AD: Microneedle-based drug delivery systems: microfabrication, drug delivery and safety. *Drug Delivery* 2012; 19: 1-14.

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