

ENHANCED TRANSDERMAL PERMEATION OF DICLOFENAC SODIUM USING MANGO SEED KERNEL STARCH NANOPARTICLES

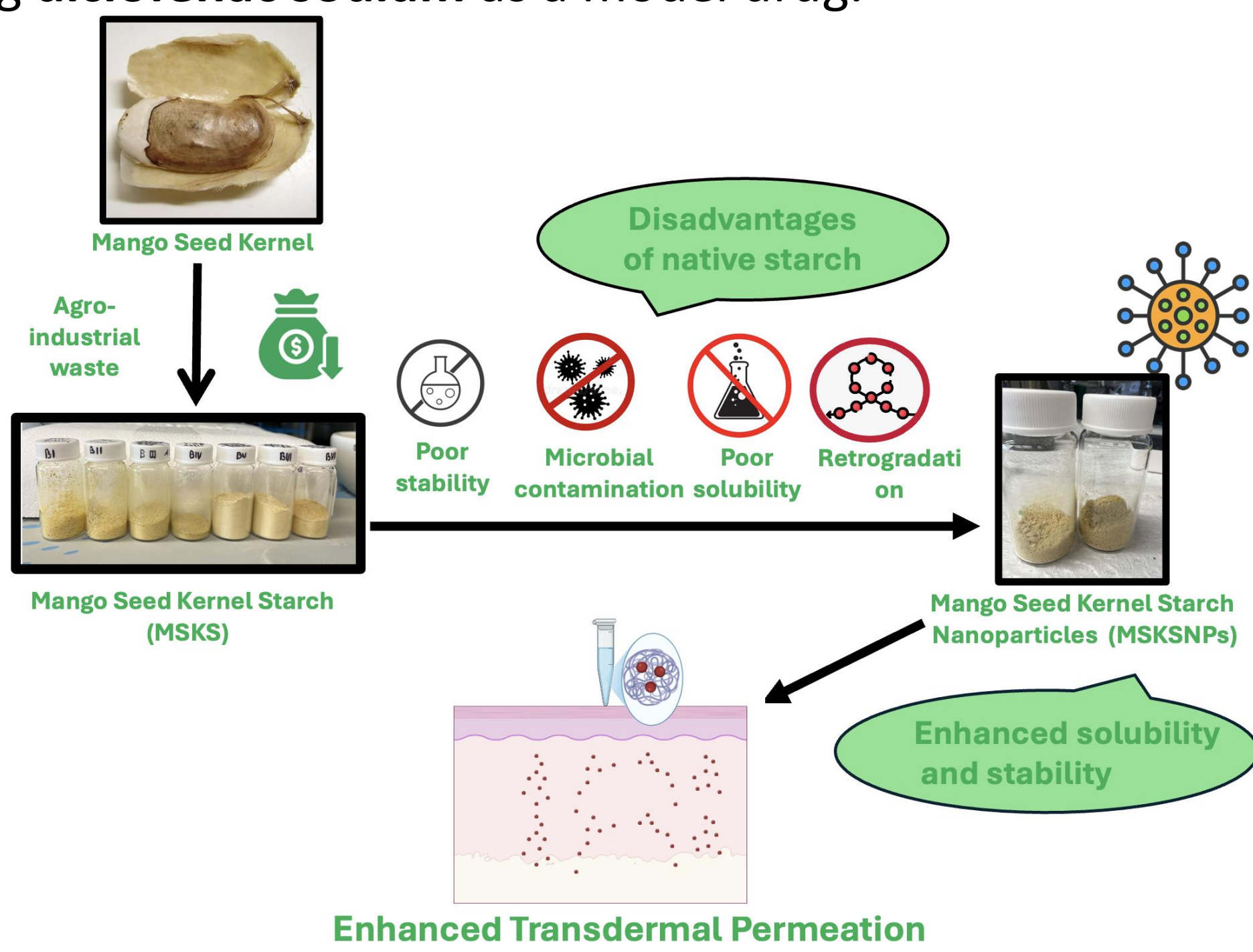
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Purpose

- The purpose of this study was to prepare and characterize diclofenac-loaded nanoparticles obtained from **starch isolated from mango seed kernels (MSKS)** and evaluate the **transdermal permeation**.
- MSKS **extraction** was performed using an **alkaline method** followed by **freeze-drying**, and its physicochemical properties were compared with commercial corn starch. MSKS **nanoparticles (MSKSNPs)** were synthesized using **mild alkali hydrolysis** and **ultrasonication**, then characterized for physicochemical properties, *in- vitro* **transdermal permeation** using **diclofenac sodium** as a model drug.



Results

Sl. No	Drying method	Solid: solvent ratio	Sedimentation time (Hour)	% yield
1	Tray Dryer (40°C) for 6 hours	1:10	24	17.23±5.96
2	Tray Dryer (40°C) for 6 hours	1:10	48	20.2±3.24
3	Tray Dryer (40°C) for 6 hours	1:12	24	15.4±6.41
4	Air dry (RT 20°C) for 24 hours	1:12	24	18.6±2.32
5	Air dry (RT 20°C) for 24 hours	1:12	48	36.2±3.23
6	Freeze drying for 24 hours	1:10	48	20.2±2.15
7	Freeze drying for 24 hours	1:12	48	30.4±3.15
8	Freeze drying for 24 hours	1:14	48	67.7±5.20
9	Freeze drying for 24 hours	1:15	48	60.5±3.21

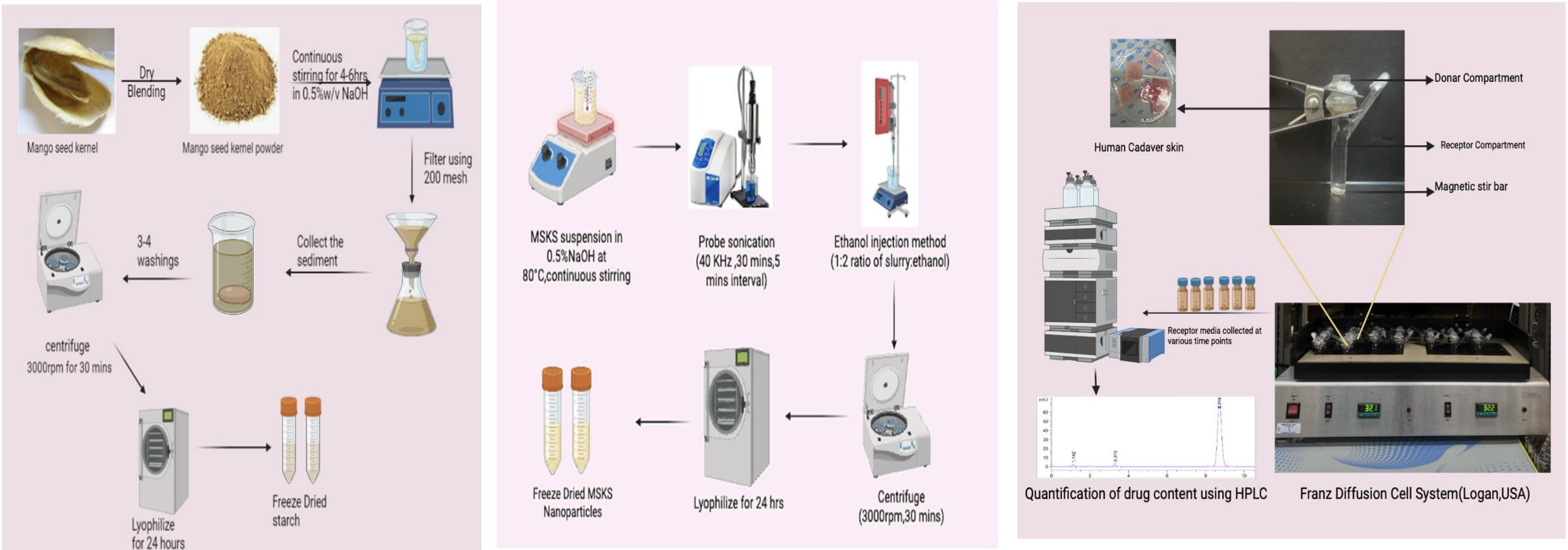
Table 1:Effect of process parameters on percentage yield.

Sl. No.	Parameter	MSKS (Mean±SD)	Corn starch (Mean±SD)	PSD
1	Solubility (%)	17±2.8	14±3.2	• 140±3.6 nm with a polydispersity index of 0.42±0.03.
2	pH	7.0±1.2	7.0±0.6	
3	Moisture content (%)	7.4±0.8	11.8±1.2	• Confirmed the chemical integrity of Starch.
4	Water Holding Capacity (%)	79.35±0.8	72.93±0.6	
5	Swelling Power(g/g)	3.2±0.16	2.3±0.52	• Confirmed the globular structure of MSKSNPs with particle sizes lower than 100nm.
6	Gelatinization temperature(°C)	60±2.5	66.7	
7.	Amylose/ Amylopectin content	0.35	0.33	• The degree of crystal size of diclofenac is reduced to 14 nm when compared with the pure drug, 33 nm.
				• 82.34±5.2.

Table 2:Physico-chemical properties of MSKS compared with corn starch.

Table 3:Physico-chemical Characterization Of Mango Seed Kernel Starch Nanoparticles

Methods and Materials



Extraction of starch from mango seed kernels by modified Alkaline method

Preparation of MSKSNP'S nanoparticles by mild alkaline hydrolysis and ultrasonication method.

Invitro permeation testing using Franz Diffusion Cell System

Figure 1:Schematic Illustration of Isolation of Mango Seed Kernel Starch, Preparation of Mango Seed Kernel Starch Nanoparticles, and In-vitro Permeation Testing

Results

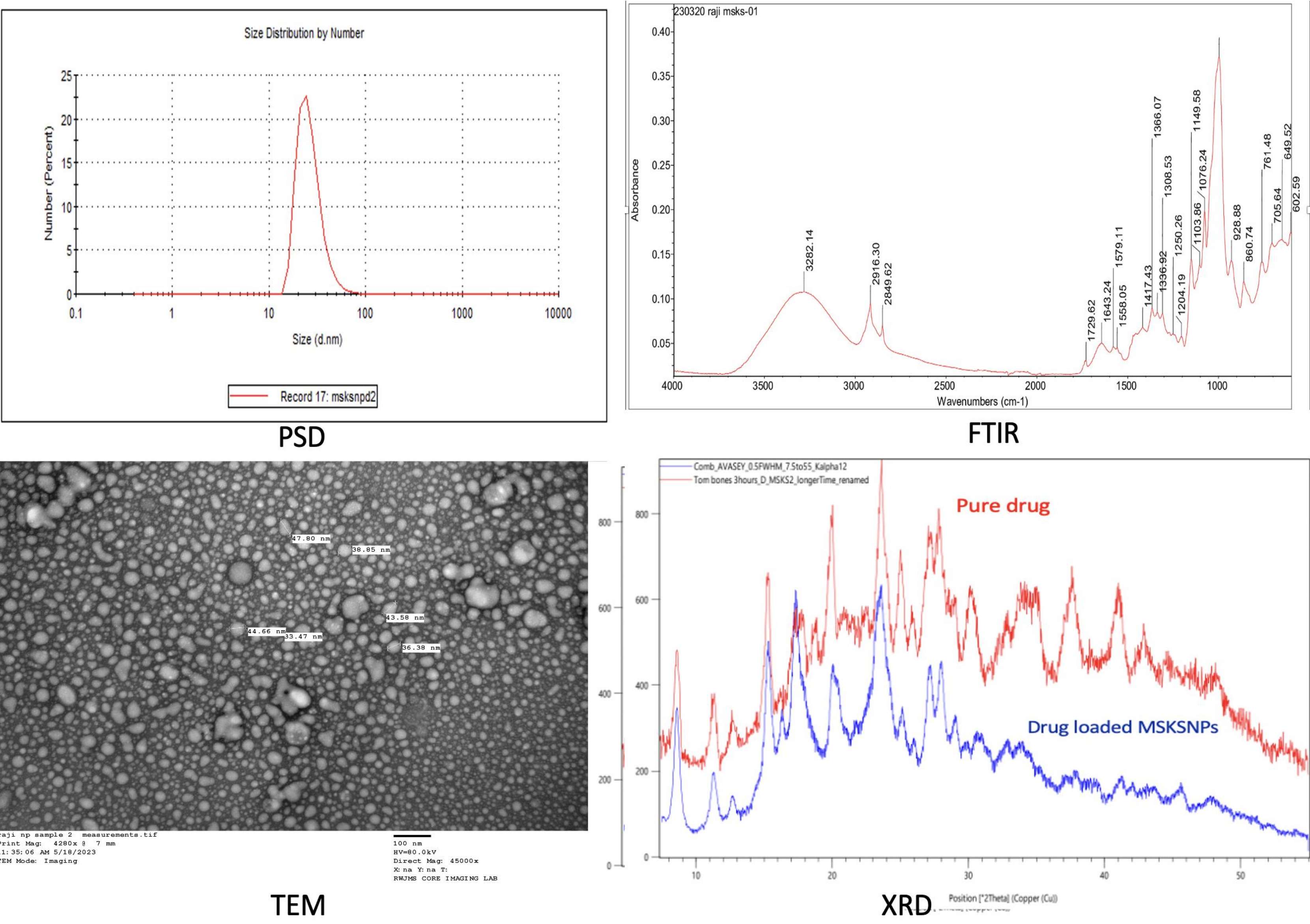


Figure 2:Physico: chemical characterization of MSKSNPs

References

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Results

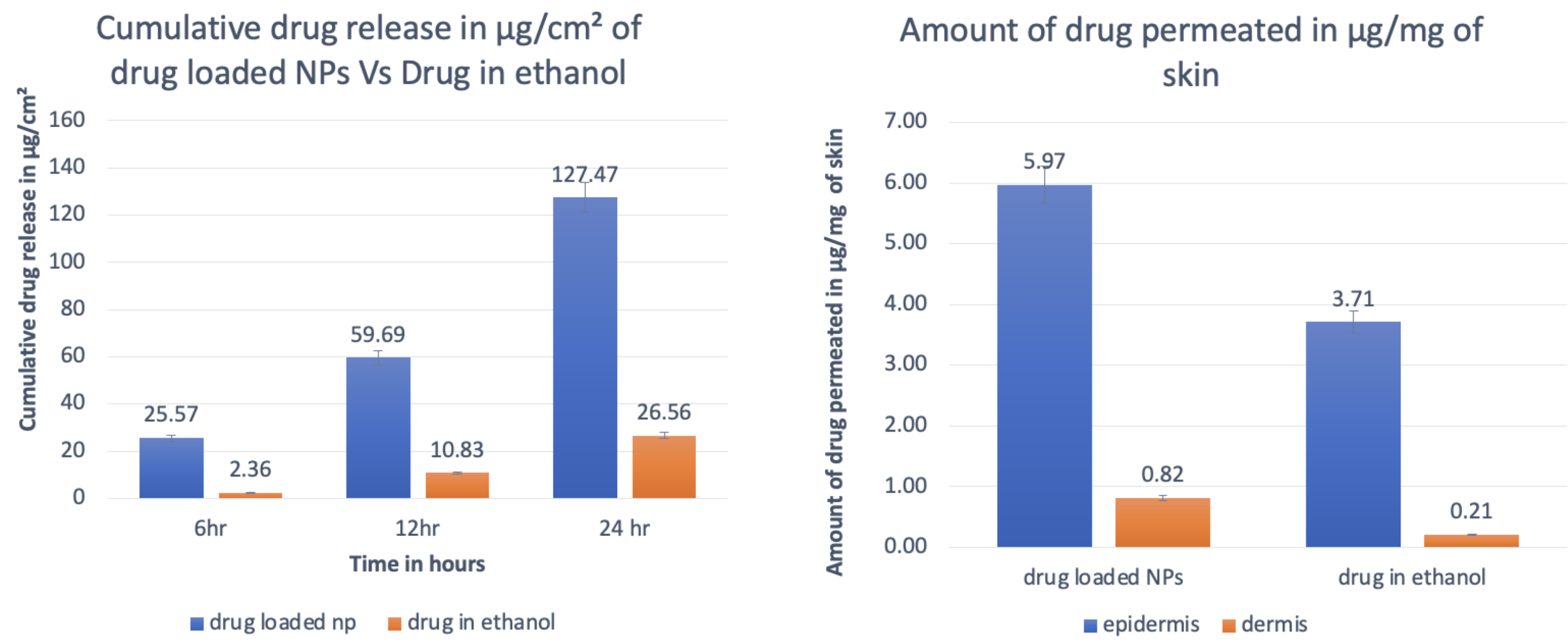


Figure 3: IVPT data of diclofenac Sodium loaded Mango Seed Kernel Starch Nanoparticles (MSKSNPs) as compared with control ethanolic diclofenac solution. Concentrations of the drug were 10mg/ml in both test and control samples (n=3). A) Cumulative drug release of drug-loaded nanoparticles vs ethanolic diclofenac solution for 6,12, and 24-hour time points. B) Amount of drug permeated in µg/mg of epidermis and dermis.

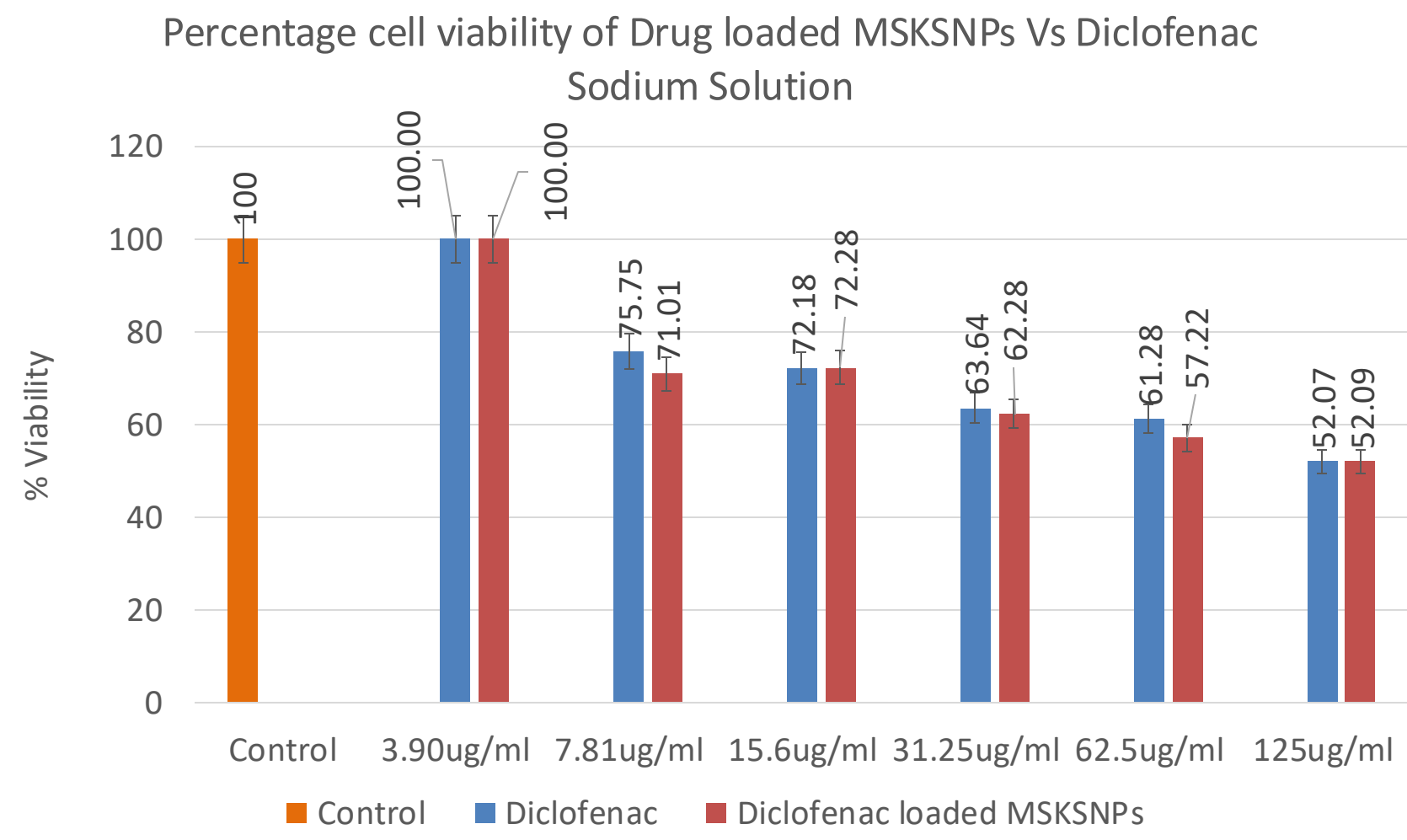


Figure 4: Percentage cell viability using Alamar Blue Assay of drug-loaded Mango Seed Kernel Starch Nanoparticles (MSKSNPs) and ethanolic solution of diclofenac sodium. The study was conducted in three replicates with three individual groups.

Conclusions

- Mango seed starch nanoparticles were successfully synthesized using mild alkali hydrolysis and ultrasonication.
- The method was **relatively simple** and took **less time** than acid hydrolysis. The MSKSNPs were found to be **more amorphous** compared to the native starch.
- These nanoparticles showed **enhanced transdermal permeation** compared with the ethanolic drug solution.

Future Directions

- The study will be expanded to include other BCS Class II drugs.
- Antioxidant and antimicrobial studies will be conducted on the nanoparticle formulations.

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