



## Sun Screen Potential of *Murraya koenigii* (L.)

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### ABSTRACT

*Murraya koenigii* (L.) Spreng, colloquially known as meethi neem, kari patta or curry leaves belongs to the Rutaceae. The plant is renowned in various traditional system of medicine for its peculiar aroma and therapeutic significance. The plants prosper well in tropical and subtropical climates *M. koenigii* is significantly used in Indian culinary and complementary system of medicine to mitigate various disorder. UV-radiation perceived as major cause that vitiate the intuitive nature and function of skin. UV-radiation may cause detrimental effect to the usual characteristics of human skin. UV-filters or sunscreens are the agent that could help to assuage the deleterious effect of UV radiation by absorbing and tempering the harmfulness to a major extent. The present study supports the sun protective efficacy of *Murraya koenigii* in concentration dependent manner that could broad cosmeceutical horizon of *Murraya koenigii* (L.).

**Keywords:** Sun screen, *Murraya koenigii* , UV radiations, SPF

### INTRODUCTION

Plants are vital for human existence in the earth. Plants recognized as a preferential source of medicine and used to alleviate, recuperate and treat variety of ailments from ancient times. *Murraya koenigii* (L.) Spreng, colloquially known as meethi neem, kari patta or curry leaves belongs to the Rutaceae. The plant is renowned in various traditional system of medicine for its peculiar aroma and therapeutic significance. The plants prosper well in tropical and subtropical climates *M. koenigii* is significantly used in Indian culinary and complementary system of medicine to mitigate various disorder. Leaves, bark and roots are used in skin eruptions, stimulant, tonic, stomachic, dysentery, diarrhea, anthelmintic, inflammation, analgesic, curing piles, itching, leukoderma, vomiting and blood disorders. The vital constituents of plant are O-phellandrene, P-caryophyllene, P-elemene and P-gurjunene,  $\alpha$ -pinene,  $\beta$ -phellandrene,  $\beta$ -caryophyllene carbazole alkaloids, coumarins, acridine, carbazole alkaloids, essential oil, carotenoids, saponins, proteins, steroids, tannins, flavonoids, phenylpropanoids, glycosides, phenolics, nicotinic acid, vitamin C, sesquiterpenes etc. The leaves of plant are green, lanceolate, exstipulate, bipinnately compound with reticulate venation. The plant has been asserted to possess array of therapeutic properties includes anti-oxidant, antimicrobial, antibacterial, immunomodulation, hepatoprotective, antidiabetic, wound healing, antipyretic, cytotoxic, antiulcer, cholesterol lowering, anti-obesity, neuroprotection and antitrichomonal effects. Due to presence of vital corresponding component, sunscreen activity has been chosen for the study. Sun screen agent are compounds which principally used to protect the skin from deleterious UV radiations and oxidative stress. [1-6]

Minimal erythema dose in sunscreen protected skin

$$\text{SPF} = \frac{\text{Minimal erythema dose in sunscreen protected skin}}{\text{Minimal erythema dose in non-sunscreen protected skin}}$$

MED is the minimum time interval or dosage of ultraviolet irradiation engender noticeable erythema on protected or unprotected skin. Higher SPF value would be beacon of higher protection against UV radiation. The sample absorbance was recorded at 5 nm interval in the range of 290-320 nm. SPF is determined by spectrophotometer. The SPF value was reckoned by using the formula. [7-14]

$$\text{SPF}_{\text{Spectrophotometer}} = \frac{320}{290} \times \sum \frac{EE(\lambda) \times I(\lambda) \times \text{Abs}(\lambda)}{CF}$$

Where CF is denoting Correction factor (10),  $EE(\lambda)$  indicates erythmogenic effect of bring about by radiation at wavelength ( $\lambda$ )  $\text{Abs}(\lambda)$  corresponds to spectrophotometric absorbance values at discrete wavelength ( $\lambda$ ). The value of  $EE(\lambda) \times I(\lambda)$  taken as a constant and displayed in Table 1.



## MATERIAL AND METHODS-

Analytical grade chemical and glassware of ASGI mark had been used to perform study. The analysis of sample was done in UV-VIS Spectrophotometer model UV-1700 Pharmaspec Shimadzu.

### Collection and Processing of Plant material

The leaves of *Murraya koenigii* have been collected in the month of March from Madhav university campus, abu road, Rajasthan, India. The collected plant leaves were thoroughly washed with tap water then shade dried till crumpled. The dried leaves were used to made coarse powder. The powder is shifted to obtain uniform size, The powdered was then subjected to extraction with selected solvents.

### Extraction of Plant Material

The hydro alcoholic extract has been prepared by steeping the plant drug in the selected solvent for seven days consecutive with occasional stirring. 200g of powdered plant material was accurately weighed, each 50 g of drug was extracted with 60%,70%,80% and 90% of alcohol respectively, the extract is then filtered thrice through whatman filter, the filtrate was collected, evaporated to dryness. The residual solvent was expunged in desiccator. The yield of individual extract was reckoned and recorded. [8,9]

### Sample Preparation

10 mg of plant extract mixed with 100 mL of hydroalcoholic solution to get 100µg/mL. The mixture is then filtered through Whatman filter paper, three dilution 40µg/mL, 50µg/mL and 60µg/mL were made with stock solution, each sample had been scanned thrice for selected wavelength at 5 nm intervals through UV spectrophotometer. The base line correction was made with similar solvent used for extraction. The absorbance of selected dilutions of *Lantana camara* extract was recorded. [8,9]

### In vitro SPF Determination

The UV absorption efficacy of *Murraya koenigii* extracts were evaluated by spectrophotometric method. The 40µg/mL, 50µg/mL and 60µg/mL dilutions of disparate extract were made from initial stock solution, the devised dilutions were scanned thrice in the range of 290 nm to 320 nm at 5 nm interval. The mean of absorbance was taken for each discrete concentration, the absorbance values has been multiplied with the constant. the summation of those multiplied with correction factor constant10. [7-14]

Table 1: Product Function Used in Calculation of SPF

Sr. No	Wavelength in nm	EE(λ) X I (normalized)
1	290	0.015
2	295	0.0817
3	300	0.2874
4	305	0.3278
5	310	0.1864
6	315	0.0839
7	320	0.018
Total		1

## RESULT AND DISCUSSION

The percentage yield of plant extract with different solvents was found as *Murraya koenigii* 5.8%,6.3%,6.8%,7.1% The result corroborated that 90% hydroalcoholic solvent had more extractable efficiency in terms of maximum yield compared to other extraction solvent used in the study. The spectrophotometric SPF evaluation method could be useful in development of sun protective preparation. It would be the better alternative for preliminary evaluation before going to *In-vivo* study. In this study plant extracts were evaluated by UV-spectrophotometry. The SPF was reckoned by Mansur equation. The observation and outcome revealed that hydro alcoholic extracts of *Murraya koenigii* had potential of sun screen and could be used as favorable sunscreen ingredients in cosmeceutical development. 90% hydroalcoholic extract had shown greater potential compared to other used in the study, although extract of 60% hydroalcoholic had observed as lowest UV screening potential.



Table no.2- In vitro SPF value at concentration 40µg/mL

Sr. No	Wave length in nm	EE(λ) X I (normalized)	MK 60% (absorbance) 40µg/ml	MK 70% (absorbance) 40µg/ml	MK 80% (absorbance) 40µg/ml	MK 90% (absorbance) 40µg/ml
1	290	0.015	3.1155±0.019	3.9554±0.012	4.7425±0.018	5.8577±0.016
2	295	0.0817	2.8254±0.017	3.5249±0.011	4.4531±0.016	5.6141±0.020
3	300	0.2874	2.6224±0.021	3.2124±0.018	4.1982±0.014	5.2912±0.015
4	305	0.3278	2.2552±0.015	2.9629±0.015	3.8625±0.015	4.9674±0.016
5	310	0.1864	1.7973±0.011	2.5154±0.018	3.5501±0.019	3.7658±0.018
6	315	0.0837	1.5752±0.014	2.1204±0.013	3.2217±0.018	3.5515±0.017
7	320	0.018	1.1106±0.019	1.9565±0.011	2.9859±0.018	3.1480±0.012

Value=Mean± SD, MK- *Murraya koenigii*

Table no.3- In vitro SPF value at concentration 50µg/mL

Sr. No	Wave length in nm	EE(λ)XI (normalized)	MK 60% (absorbance) 50µg/ml	MK 70% (absorbance) 50µg/ml	MK 80% (absorbance) 50µg/ml	MK 90% (absorbance) 50µg/ml
1	290	0.015	3.9684±0.011	4.8911±0.012	5.7582±0.012	6.4471±0.011
2	295	0.0817	3.6274±0.015	4.6954±0.015	5.5528±0.017	6.1934±0.015
3	300	0.2874	3.3081±0.019	4.3514±0.014	5.2625±0.018	5.8187±0.017
4	305	0.3278	3.1224±0.018	4.1952±0.011	4.9162±0.019	5.5264±0.018
5	310	0.1864	2.8117±0.017	3.8181±0.021	4.5014±0.021	5.2781±0.021
6	315	0.0837	2.5442±0.012	3.5859±0.016	4.2512±0.015	4.9254±0.012
7	320	0.018	1.9241±0.014	2.9114±0.019	3.8914±0.012	4.3715±0.015

Value=Mean± SD, MK - *Murraya koenigii*

Table no.4- In vitro SPF value at concentration 60µg/mL

Sr. No	Wave length in nm	EE(λ)XI (normalized)	MK 60% (absorbance) 60µg/ml	MK 70% (absorbance) 60µg/ml	MK 80% (absorbance) 60µg/ml	MK 90% (absorbance) 60µg/ml
1	290	0.015	4.9361±0.017	5.4194±0.022	6.7451±0.016	7.6935±0.018
2	295	0.0817	4.6224±0.019	5.2584±0.017	6.5074±0.012	7.3812±0.014
3	300	0.2874	4.2101±0.024	4.9552±0.014	6.1403±0.021	7.1511±0.015
4	305	0.3278	3.9581±0.015	4.5621±0.012	5.7741±0.017	6.8204±0.011
5	310	0.1864	3.7538±0.012	4.1851±0.011	5.4714±0.019	6.6851±0.013
6	315	0.0837	3.4714±0.011	3.9887±0.017	4.9887±0.013	6.3175±0.022
7	320	0.018	3.1141±0.015	3.2115±0.011	4.5281±0.015	5.9134±0.019

Value=Mean± SD, MK - *Murraya koenigii*



Table no.5- Spectrophotometric values of SPF at different concentration

Sr.No.	Extract	SPF 40µg/ml	SPF 50µg/ml	SPF 60µg/ml
1	MK 60%	3.224	4.431	5.722
2	MK 70%	4.176	5.924	6.574
3	MK 80%	5.561	7.041	8.298
4	MK 90%	6.787	7.944	9.843

MK - *Murraya koenigii*

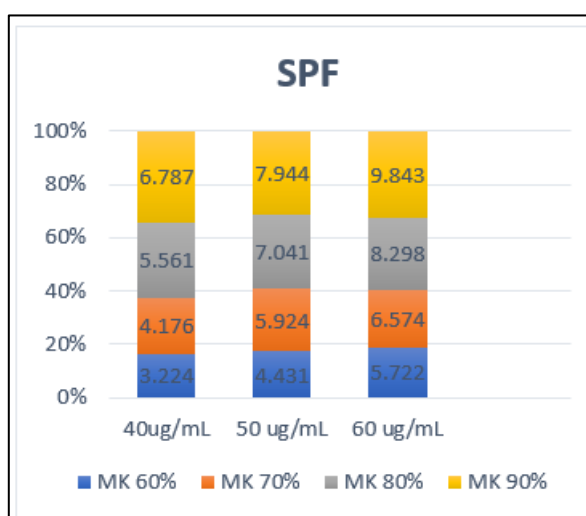


Figure 1-Graphical Presentation of SPF value

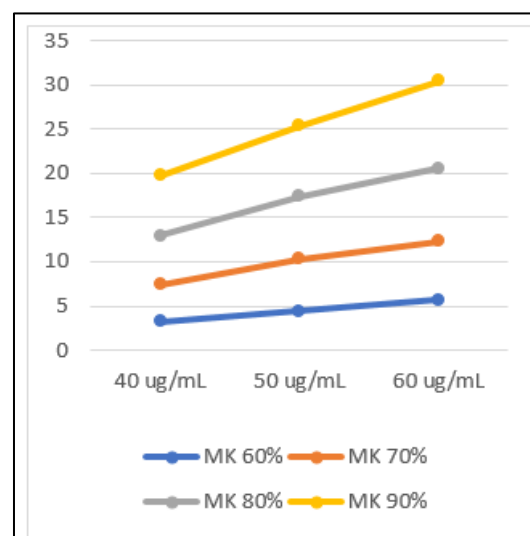


Figure 2 – Line diagram for SPF value

The result substantiates the photo protective efficiency is contingent to concentration of solute, As the ratio of extract increased the SPF has been increased that might be due to more promising solute at higher concentration. The plant selected for the study already had multitudinous medicinal benefits, this additional property would extend therapeutic vein of *Murraya koenigii*.

## CONCLUSION

*Murraya koenigii* prospers well in tropical and subtropical climates. The plant is renowned for its therapeutic significance. The present study unfolds new horizon and scope for investigation of drug. The efficacy of hydroalcoholic extract of leaves made plant more apposite to use as sun screen agent. The nature could redress suffering and ailments of mankind. In the recent years many novel compounds have been successfully discovered from the natural sources and many are there in the vault of nature. Natural therapy is always convivial and biocompatible than synthetic counterparts. *Murraya koenigii* is traditionally acclaimed for its medicinal properties. The study paves the way for further research and investigation in relevant field. The present study unfolded the sun protective efficacy of *Murraya koenigii* in concentration dependent manner that could heightened cosmeceutical horizon of the drug.

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