FAST AND ADVANCED ANALYTICAL METHOD FOR DETERMINATION OF PERMETHRIN (CIS AND TRANS) CONTENT AND IN TRACES BY ULTRA PERFORMANCE LIQUID CHROMATOGRAPHY

K. Sumanth Prabhu1*, A. Venkatachalam2 and Subhash Kale3

1Research Scholar, Faculty of Pharmacy, Pacific Academy of Higher Education and Research University, Udaipur-313024 (Rajasthan) India.
2Research guide, Faculty of Pharmacy, Pacific Academy of Higher Education and Research University, Udaipur-313024 (Rajasthan) India.
3Research Scholar, Faculty of Science, Pacific Academy of Higher Education and Research University, Udaipur-313024 (Rajasthan) India

*E-mail: ksumanthprabhu@gmail.com

ABSTRACT

Pyrethroids are ester of dichloro analog compound that called as Permethrin. Permethrin is widely used as pesticide and insecticide, and off-late as an ectoparasiticide. It is majorly used for effective and short term treatment against lice that is usually found in human and animal head and body hair. Further, Permethrin can also be used against infections namely scabies that formed from insects namely Mite in humans and also in human immunodeficiency virus-infected patients. Isomers of Permethrin differ in their toxicity while (1R, 3S)-trans as well as (1R, 3R)-cis isomers are typically effective for the insects killing properties.

Permethrin is usually available in cream or lotion form of formulation in the market hence; it is applied on the skin as a cream or lotion. Further estimation of Permethrin in an environment samples like Waste/Potable water and pharmaceuticals is important as a control strategy during environment (water pollution) monitoring, formulation batch release and stability study. This research study emphasizes on the development of an Ultra Performance Liquid Chromatography (UPLC) based rapid detection and quantitation with an accurate and precise analytical method for Permethrin and its traces by an Ultra Performance Liquid Chromatography. The external standard method is used for assay determination in Waste/Potable water, and Cream formulation samples. To prove the suitability of the developed analytical method for its intended purpose, its validation study was performed following an International council for Harmonization guidelines.

Keywords: Permethrin, Scabies, Lice, Mite, UPLC, External standard, Method validation, Waste/Potable water, Cream formulation.

©RASĀYAN. All rights reserved

INTRODUCTION

Pyrethroids have been in use for a few decades.1 Permethrin is formulated as cream/lotion for skin applications, sprayed on mosquito nets or clothing as a repellent to kill the insects that touch them. As an ectoparasiticide used against the killing of lice found in the hair on the head and body of human and animals2 (Fig.-1). It is further effective against Mite infected disease like scabies (Fig.-2 and 3) and also in humans with HIV (human immunodeficiency virus) infected patients.3 Permethrin paralyzes and kills the Mites and Lice directly as well as destroy their eggs. It has a wide range of applications being used also as a pesticide, insecticide because it has highly potent action against Lice and Mite type of insects. Due to carcinogenic classification by USEPA (the United States Environment Protection Agency) and differential toxicity of its isomer, its control to human exposure has become important from a safety and toxicity perspective. Hence it is essential to develop and apply the analytical method for identification and
individual quantitation in environment samples like waste/potable water to study water pollution as well as in pharmaceuticals during batch release and stability study. Permethrin chemical compound has a total of four types of stereo isomers (two pairs of enantiomeric forms); those are arising from the two types of stereocenters in the cyclo-propane ring. The Transform of enantiomeric pair is called as trans-permethrin. In fact (1R, 3S)-trans and (1R, 3R)-cis types of enantiomers are responsible for the insecticidal properties of Permethrin. Permethrin is insoluble in water and it is photostable.

![Fig.-1: Human Head Scalp Suffering From Lice (Hiding in Hairs)](image)

![Fig.-2: Human Skin Suffering From Scabies](image)

![Fig.-3: Human Skin Suffering From Mites that Causes Scabies](image)

**Permethrin Stereochemistry**

The four stereo-isomers are depicted in Fig.-4 to Fig.-8.

![Fig.-4: Permethrin (Cis and Trans)](image)

![Fig.-5: (1S,3R)-trans Enantiomer](image)

![Fig.-6: (1R,3S)-trans Enantiomer](image)

![Fig.-7: (1S,3S)-cis Enantiomer](image)
The main purpose of the present research study on Permethrin chemical drug was to develop a rapid, accurate, and precise chromatography (UPLC) based analytical technique for the determination of Permethrin, which has total four stereoisomeric (two enantiomeric pair) forms in waste/potable water as a part of environmental pollution monitoring and its traces by Ultra Performance Liquid Chromatography. Further, UPLC was considered superior over the conventional HPLC (High-performance liquid chromatography) methods which either have very long run time (up to 80 min and higher) or are specific to one kind of formulation (not universal) etc. The effort and challenges were in developing the chromatography which can determine Permethrin even in trace amounts in waste and potable water sample in addition to the pharmaceuticals. The external standard method is used here for the assay determination in waste/potable water, and cream-based formulation.

EXPERIMENTAL

Materials and Methods
Permethrin Standard, Purified water, Ammonium acetate (GR grade), Ortho-phosphoric acid (GR grade), Methanol (Chromatography grade, filtered through 0.22 µm filter), Permethrin cream 5% w/w formulation product (procured commercially from local pharmacy store) and its placebo (prepared in-house).

Samples for Analysis
Formulation
Permethrin (Cis and Trans) - 5% w/w in a cream base.

Potable and Wastewater
(*Details captured under “Potable and Wastewater samples collection procedure”)

Instruments Required
pH meter, UPLC from Waters Corporation USA (Model - ACQUITY UPLC H-Class PLUS System), Weighing balance, Vortex stirrer, Analytical glassware’s and Sonicator.

General Procedure
Standard Solution-1
0.2 mg/mL of Permethrin in diluent (dissolved by sonication).

Standard Solution-2
0.05 mg/mL of Permethrin in diluent (dissolved by sonication).

Sample Solution-1 (for Formulation Cream)
Weigh accurately about 1.0 g of cream sample (equivalent to 50 mg of Permethrin) in 50 mL of the volumetric flask, add 20 mL of diluent, sonicate to dissolve and dilute to volume with the diluent. Sonicate for 5 min and mix. Further pipette 5 mL of this solution into 25 mL volumetric flask and dilute with the diluent and mix.

Sample Solution-2 (for Waste and Potable Water)
Transfer 100 mL of duly filtered (through 0.45 µm Nylon filter) sample into 250 mL evaporating dish. Carefully evaporate liquid part of the sample till dryness and dissolve an obtained residue with 10 mL of methanol and mix.
Placebo Solution
Weigh accurately about 1.0 g of placebo (without Permethrin) in 50 mL volumetric flask, dissolve the content in 20 mL of diluent through sonication and dilute to volume with the diluent. Sonicate for 5 min and mix. Further, transfer 5 mL of this solution to 25 mL volumetric flask and dilute with the diluent and mix.

Potable and wastewater sample collection procedure
Scabies and Lice related diseases are very common in the slum area where people live in unhygienic conditions. Due to this issue, physicians often prescribe Permethrin lotion/cream/shampoo type of medication to treat lice and scabies in a slum area to get rid of related infections. After the application of lotion/cream/shampoo, the residual Permethrin in drug formulation gets washed away during bathing or cleaning of the infected body part. This untreated Permethrin remains in wastewater which then mixes with potable water through contamination due to unhygienic sanitation practices. Thus from potable water, Permethrin gets ingested into the human body that causes health hazards related to this drug. Hence, it was decided to collect wastewater and potable water samples from the Dharavi slum area which is the largest slum in Mumbai (India) and Asia region. Total 6 samples of wastewater and potable water each were collected from different locations (Table-1).

Table-1: Details of Wastewater / Potable Water Samples From Different Locations of Dharavi Slum Area

| Sample ID | Sampling Location   | Co-ordinates                  |
|-----------|---------------------|-------------------------------|
| ^DWW-1 & ^DPW-1 | Bhatiya nagar          | DMS Decimal 19° 2’ 24.75” N, 72° 51’ 3.06” E |
| DWW-2 & DPW-2 | Tadwadi lane          | 19.040208, 72.85085 | Geo URI |
| DWW-3 & DPW-3 | Dharavi Crossroad     | 19.040208, 72.85085 |
| DWW-4 & DPW-4 | Sheshwadi road        |                               |
| DWW-5 & DPW-5 | Machi galli           |                               |
| DWW-6 & DPW-6 | Kala Killa road       |                               |

^DWW- Dharavi wastewater & ^DPW- Dharavi potable water

Detection Method
Chromatographic Conditions
Instrument: UPLC (Ultra High-Performance Liquid Chromatography), Make – Waters Corp. USA.
Column: Ethylene Bridged Hybrid ODS, (2.1 mm ID x 50 mm Length), 1.7 micrometers,
Column Temperature: 25°C
Mobile Phase: A - 0.05M Ammonium acetate buffer, B – Methanol
Elution Mode: Gradient elution
Diluent: Methanol
Pump Flow Rate: 0.4 mL/min,
Detector: UV (235 nanometer),
Injection Volume: 1.0 microliter
Auto-sampler Temperature: 25°C,
Total Gradient Run Time: 13 min
Preparation of Blank, Placebo, Standard and Sample solutions:
Blank Solution: Used diluent as blank
Analytical Discussion

Method Development and Optimization (Formulation Samples and Waste/Potable Water Sample)

A UPLC chromatographic condition was established after several preliminary experiments on UPLC for selecting the proper reverse phase C18 stationary phase, mobile phase pH and gradient proportion. Different gradient programs were tested, and selection of the suitable gradient program (Table-2) that estimates Permethrin content precisely and accurately in less run time (13 min) depended on its ability to pass system suitability parameters as mentioned in Table-3. Satisfactory system suitability criteria achieved on BEH C-18 (Ethylene Bridged Hybrid ODS), (2.1 mm x 50 mm), 1.7 µm using gradient program as mentioned in Table-3. Mobile phase with varying gradient proportion of Ammonium acetate buffer (0.05M) and Methanol used to achieve satisfactory resolution between Cis and Tran Permethrin. Overall runtime set to be 13 min with completion of gradient and stabilization for next run.

| Parameter                                | Acceptance Criteria | Cream   | Wastewater | Potable Water |
|------------------------------------------|---------------------|---------|------------|---------------|
| Tailing factor                           | NMT 2.0             | 1.06    | 1.21       | 1.32          |
| Theoretical plates                       | NLT 2000            | 8050    | 4562       | 3980          |
| RSD (%) of six injections (Standard response) | NMT 3 % For cream and NMT 10 % for waste/ potable water assay | 0.29%  | 4.96%      | 6.24%         |
| RSD (%) of six injections (Standard RT)  | NMT 1 %             | 0.62%   | 0.88%      | 0.93%         |

Table-3: System Suitability of Test Method

Intra-day precision and Inter-day precision (Ruggedness) have been established for the proposed method with % RSD (Relative standard deviation) for intraday found to be 0.68% and that for inter-day 1.65% for Assay test of cream while % RSD for intraday found to be 5.25% and that for inter-day 6.05% for assay of wastewater samples and % RSD for intraday found to be 7.62% and that for inter-day 8.11% for assay of potable water samples indicating preciseness of proposed UPLC method for Permethrin determination in the Cream formulation, wastewater and potable water samples.

Accuracy

Accuracy of the developed UPLC method was established at three different levels i.e. 50%, 100% and 150% of test (Permethrin) concentration by adding a known amount of Permethrin standard into the placebo and in highly purified water. Further, each level prepared in triplicate and estimating the sample on the UPLC test method. Three sets in triplicate were prepared and analyzed separately for the Cream formulation, wastewater and potable water samples.

Robustness

Robustness was established by deliberately changing instrument parameters of test methods like flow rate, buffer pH and wavelength of UV detector were carried out as per ICH guidelines to estimate their effects on the proposed method.

RESULTS AND DISCUSSION

System suitability parameters namely theoretical plates per meter (TPPM), tailing factor (peak asymmetry), and percentage relative standard deviation of area response and Permethrin peak retention time of six injections were carried out and the values are well within the United States Pharmacopoeia (USP) limits as shown in Table-3 indicating UPLC system was technically working properly as per desired output for assay determination of Permethrin in the Cream formulation, water and potable water samples.

From the linearity results where a linear calibration plot of Permethrin was constructed at seven point concentration levels. From the values of $r^2$ obtained as above, it could be proved that, the proposed method for Permethrin Assay in the Cream formulation and waste/potable water samples is linear for the specified working range. The typical linearity plot is given as Fig.-9.
The precision (termed as method reproducibility) of an assay method was evaluated for repeatability or reproducibility and for intermediate precision which is also called method ruggedness of the method. For the precision experiment, the percentage relative standard deviation for the assay of Permethrin for 6 Assay determinations was found to be within the specified limits for Cream formulation and waste/potable water samples. Thus method precision characteristic to the method is proved. Obtained results are given in Table-4 with graphical representation in Fig.-10.

Table-4: Precision and Intermediate precision & Linearity \( (n^\circ = 7) \) [\( n^\circ \) - No of injections]

| Parameter          | Limits  | Cream Product | Wastewater | Potable Water |
|--------------------|---------|---------------|------------|---------------|
| Precision (\% RSD) | Repeatability 2.0% | 0.68 | 5.25 | 7.62 |
|                   | Intermediate 2.0% | 1.65 | 6.05 | 8.11 |
| Linearity \( (r^2) \) | 7 - Levels | 0.998 For cream and 0.995 for w/p water assay | 0.9999 | 0.9967 | 0.9958 |

A known amount of Permethrin standards were spiked at 50\%, 100\%, and 150\% concentration levels in placebo as well as in purified water and each level performed in triplicate. From the results obtained, percentage recoveries (%-Accuracy) of drugs were calculated for Permethrin content. The %-recovery at three different concentrations found to be within the range of 97.0% to 103.0% limit set by ICH guidelines for the Cream formulation and within 80.0% and 120.0% for waste/potable water samples indicating proposed UPLC based method is accurate. The %-recovery or accuracy results are summarized in Table-5.

Table-5: Accuracy (%-Recovery)

| Accuracy Levels | Cream Formulation  | Wastewater | Potable Water |
|-----------------|-------------------|------------|---------------|
| %-Mean Recovery | %-Mean Recovery    | %-Mean Recovery |
| (Limit: 97.0%-103.0%) | (Limit: 80.0%-110.0%) | (Limit: 80.0%-110.0%) |
| Level-1 (50%)   | 98.07             | 88.07      | 86.87         |
| Level-2 (100%)  | 97.45             | 92.80      | 94.71         |
| Level-3 (150%)  | 100.45            | 105.22     | 107.42        |

The robustness of the assay method was studied by incorporating small but deliberate changes in critical analytical method parameters (i.e. variation in flow rate, buffer pH and UV (Ultraviolet) detector wavelength). In all the varied UPLC instrument conditions; there was no significant change in system suitability parameters indicating a proposed method for Permethrin estimation in the Cream formulation and waste/potable water samples is Robust.

CONCLUSION

Finally, based on the summarized data as above, it could be concluded that we have precisely and accurately developed a proposed method for Permethrin estimation in potable/wastewater and formulation.
The developed method is duly validated following ICH guidance document. The developed method is new (i.e. our innovation), simple, rapid, precise and accurate, external standard based UPLC method for the estimation of Permethrin in Cream (OTC-Over the counter) Formulation, wastewater and potable water samples. This developed method has proved that instruments like UPLC can be extremely useful in the estimation of Permethrin compound in the cream formulation and at trace levels in waste/potable water samples in less time (i.e. within 13 min) due to its high pressure and characteristic small particle size stationary phase. Hence developed method proved to be very cost-effective as well as time-saving with user friendly.

The proposed method is duly validated and shall be applied for routine estimation of Permethrin in cream-based or similar other formulations with a minor change in sample preparation. This formulation is mainly used against the killing of Mites that causes Scabies and Lice that are usually observed in the hair scalp of the head in humans and animals.

Thus, this method can be applied for the estimation of Permethrin content in Cream as well as similar other several formulations available in the market that contain Permethrin as an active ingredient in OTC products. For other formulations, some modifications in sample preparation as well as UPLC conditions may be required based on types of inactive ingredients present in such formulations.

Further, this method helped during the estimation of Permethrin content in waste and potable water samples indicating potable water in the slum area is getting contaminated with Permethrin from wastewater. Further study is needed in this regard.

ACKNOWLEDGMENT

The authors were thankful to “RELIABLE’s SHREE INDUSTRIAL TRAINING CENTER”, First Floor, Lathi Plaza, Opp. Shahu Maharaj Complex, Jalgaon (Maharashtra – India) for allowing to use analytical and instrumental facilities to carry out experimental work mentioned in this research paper. While Permethrin cream drug product OTC samples were taken from Indian Pharmaceutical Market.

REFERENCES

1. Maja A. Shishovska, Marina T. Stefova, Journal of Chromatographic Science, 50(1),43(2012), DOI: 10.1093/chromsci/bmr014
2. C.V. Bainbridge, G.L. Klein, S.I. Neibart, Clinical Paediatrics, 37,17(1998), DOI: 10.1177/000992289803700103
3. W. Liu, J.J. Gan, S. Qin, Chirality,17, S127(2005), DOI:10.1002/chir.20122
4. Julia Dornetshuber, Wolfgang Bicker, Michael Lämmerhofer, Wolfgang Lindner, Anneliese Karwan, Wilfried Bursch, BMC Pharmacology, 7 (Supplement2), A65, DOI:10.1186/1471-2210-7-S2-A65.
5. R. Manadas, F. Veiga, J. J. Sousa, M.E. Pina, Journal of Liquid Chromatography, 22, 1867(1999) DOI:10.1081/JLC-100101772
6. E. Garcia, A. Garca, C. Barbasa, Journal of Pharmaceutical and Biomedical Analysis, 24(5), 999(2001), DOI:10.1016/s0731-7085(00)00544-6
7. Jing Ye, Meiping Jin, Weiping Liu, American Chemical Society, Symposium Series, 1085, (2011), DOI:10.1021/bk-2011-1085.ch005
8. A. Novas Dias, F. Garcia Sanchez, A. Garcia Paveja, Journal of Chromatography Science, 36(4), 211(1998), DOI:10.1093/chromsci/36.4.210
9. M. Shishovska, V. Trajkovska, Chirality, 22(5), 527(2010), DOI:10.1002/chir.20773
10. Nivruti T. Nirgude, Sanjay Shukla and A. Venkatachalam, Rasayan Journal of Chemistry, 6(1), 47 (2013).
11. Nivruti T. Nirgude, Sanjay Shukla and A. Venkatachalam, Rasayan Journal of Chemistry, 6(1), 68(2013).
12. ICH, Guidance on Analytical Method Validation, International Convention on Quality for the Pharmaceutical Industry, Toronto, Canada: International Conference on Harmonization(2002).
13. ICH, Harmonized Tripartite Guidelines, Validation of Analytical Procedures: Text and Methodology Q2 (R1), International Conference on Harmonization(2005).

[RJC-5820/2020]