WORLD JOURNAL OF PHARMACEUTICAL RESEARCH
SJIF Impact Factor 8.084
Volume 11, Issue 3, 1590-1602. Research Article ISSN 2277– 7105

DEVELOPMENT AND VALIDATION OF STABILITY INDICATING ASSAY METHOD FOR THE DETERMINATION OF ASPIRIN AND DOMPERIDONE IN BULK AND COMBINED DOSAGE FORM

[1]*Afiya Khan, [2]Dr. Mohd. Jalaluddin, [3]Mrs. Sheenam Mansuri, [4]Dr. Brajesh Kumar Arjariya and [5]Mr. Rahim Khan

Malhotra College of Pharmacy, Bhopal India.

Article Received on
02 Jan. 2022,

Revised on 23 Jan. 2022,
Accepted on 13 Feb. 2022

DOI: 10.20959/wjpr20223-23298

***Corresponding Author**
Afiya Khan
Malhotra College of Pharmacy, Bhopal India.

ABSTRACT

The aim of present study was to investigate the development and validation of Stability Indicating Assay Method for the determination of aspirin and domperidone. Method Development and Validation for Estimation of Domperidone and Aspirin in bulk or formulation by using RP-HPLC. The RP-HPLC method was developed for estimation of Aspirin and Domperidone in synthetic mixture by isocratically using 10 mM KH2PO4:Acetonitrile (20:80) as mobile phase, Prontosil C-18 column (4.6 x 250 mm, 5µparticle size) column as stationary phase and chromatogram was recorded at 231 nm. Then developed method was validated by using various parameters such as, linearity, Range accuracy, precision repeatability, intermediate precision, robustness, limit of detection, limit of quantification. The proposed methods were found to be linear with correlation coefficient close to one. Precision was determined by repeatability, Intermediate precision and reproducibility of the drugs. The robustness of developed method was checked by changing in the deliberate variation in solvent. The result obtained shows the developed methods to be Cost effective, Rapid (Short retention time), Simple, Accurate (the value of SD and % RSD less than 2), Precise and can be successfully employed in the routine analysis of these drugs in bulk drug as well as in tablet dosage form. The Simplicity, Rapidly and Reproducibility of the proposed method completely fulfil the objective of this research work.

KEYWORDS: Aspirin; Domperidone; HPLC; Ultra Violet; Validation.

INTRODUCTION

Day by day numbers of new drugs are introduced into market. Frequently, there is bigger time

period between date of presentation of medication into business sector and date of its incorporation in pharmacopeias. This happens as result of conceivable vulnerabilities in nonstop and more extensive utilization of these medications, report of new toxicities and improvement of patient resistance and presentation of better medications by contenders. In such cases standard scientific strategies for these medications may not be accessible in Pharmacopeia's. It gets to be fundamental, in this way, to grow new explanatory system for such medications. Additionally, quality is vital in every item or administration in pharmaceuticals as it includes life. Investigation which helps in discovering spatial plan of ion as in particle and vicinity or position of certain natural useful gathering in given compound. What's more surface examination assumes imperative part in material studies to get surface related physical properties, for example, geography, profundity profiling, introduction of atom and so forth. Concoction examination has some fundamental strides like, decision of system, testing, preparatory specimen treatment, partitions, last estimation and appraisal of results. It is with first step viz. decision of system, consideration ought to be practiced to choose best possible instrument to do productive examination. Wrong choice as of right now will prompt good for nothing examination. Analytical methods are broadly classified as Physical, Chemical and Instrumental analysis. Physical observation includes description of compound, measurements of its dimension (shape, size), color, odor etc. Chemical analysis includes titrimetric analysis of compound such as potentiometric, audiometry, argentometry, permagnometry etc. Instrumental methods of chemical analysis have become backbone of experimental chemistry. Method development is done for new products and for existing products. More difficulty is short of imminent in equilibrium among wanted and previous understanding to carry out sufficient optimization lesson and enhancement in acquaintance gained through such lesson.[1]

An official analytical method is used to assess characterized normal for medication substance or medication item. Option logical technique is proposed by candidate for utilization rather than administrative systematic method. Security testing structures imperative piece of procedure of medication item advancement. reason for soundness testing is to give prove on how nature of medication substance or medication item differs with time affected by mixed bag of ecological components, for example, temperature, moistness, light and empowers suggestion of capacity conditions, retest periods and timeframes of realistic usability to be set up. Two primary parts of medication item that assume vital part in timeframe of realistic usability determination are test of dynamic medication and degrades created, amid soundness

study. Cutting edge techniques for decision for quantitative examination are UV, HPLC, GC, GCMS, LCMS and HPTLC which are very advanced. Chromatographic routines are normally utilized as part of administrative labs for subjective and quantitative examination of medication substances, drug items, crude materials and natural examples all through all periods of medication advancement from exploration to quality control.[2] Superior fluid chromatography (HPLC) is quickest developing diagnostic strategy for investigation of medications. Its effortlessness, high specificity and extensive variety of affectability make it perfect for examination of numerous medications in both measurement shapes and organic liquids. High performance thin-layer chromatography (HPTLC) is classical separative technique that has enjoyed wide spread popularity particularly in analysis of complex mixtures of natural origin. Now-a-days HPTLC is turning into routine investigative method because of its preferences of low working expense, high specimen throughput and requirement for least example clean-up.

Significant point of preference of HPTLC is that few specimens can be run at same time utilizing little amount of versatile stage not at all like HPLC, in this way bringing down examination time and expense per investigation.[3]

Aspirin [Chemically 2-acetoxybenzoic acid] is also known as Acetyl Salicylic acid. It is a medication used to treat pain, fever, or inflammation. Different inflammatory conditions like Kawasaki disease, pericarditis, and rheumatic fever are treated by Aspirin. Aspirin is also used to prevent further heart attacks, ischaemic strokes, and blood clots in people at high risk. It may also decrease the risk of certain types of Carcinomas, particularly colorectal cancer. Aspirin is a non-steroidal anti-inflammatory drug (NSAID) and works similarly to other NSAIDs but also suppresses the normal functioning of platelets. It can be given by oral and rectal route. Lysine acetylsalicylate is given by IV and IM.[4] Domperidone is a peripherally selective dopamine D2 receptor antagonist, it was developed by Janssen Pharmaceutica and is used as an antiemetic, gastroprokinetic agent, and galactagogue. It may be administered orally or rectally, and is available in the form of tablets, orally disintegrating tablets, suspension, and suppositories.[5]

MATERIALS AND METHODS

Chemicals and reagents: Aspirin, Domperidone and other chemicals obtained from pharmaceutical companies (J.B chemicals and other).

Identification and Characterization of drugs

IR spectrum of Aspirin and Domperidone

The concentration of the sample in KBr should be in the range of 0.2% to 1 %. The pellet is much thicker than a liquid film, hence a lower concentration in the sample is required (Beer's Law). For the die set that you will be using, about 80 mg of the mixture is needed. Too high of a concentration causes usually difficulties to obtain clear pellets. This pellet keeps into the sample cell and scanned between 4000-400 cm -1 and IR spectra is obtained.

Determination of λ max of Aspirin and Domperidone

The λmax of Aspirin and Domperidone were determined by running the spectrum of drug solution in double beam ultraviolet spectrophotometer.

Method development of Aspirin and Domperidone

Method development and validation for estimation of Aspirin and Domperidone using RP-HPLC.[9-11]

Selection of Mobile Phase

Initially to estimate Aspirin and Domperidone in fix dosage form number of mobile phase in different ratio were tried.

Taking into consideration the system suitability parameter like RT, tailing factor, No. of theoretical plates and HETP, the mobile phase found to be most suitable for analysis was 10 mM KH2PO4: acetonitrile (pH 3.5 with OPA) in the ratio of 20:80v/v. The mobile phase was filtered through 0.45 µ filter paper to remove particulate matter and then degassed by sonication. Flow rate employed for analysis was 1.0 ml/min 10.[6]

Procedure for preparation of mobile phase

10 mM KH2PO4: acetonitrile (pH 3.5) in the ratio of 20:80v/v, pH 3.0 with Ortho phosphoric acid. Filtered through 0.45 µ filter paper.

Selection of Diluent

Diluent used for preparation of sample were compatible with mobile phase and no any significant affect retention and resolution of analyte. After various trials Acetonitrile was used as diluents.

Selection of separation variable

1. Preparation of standard Stock solution

Accurately weighed 10 mg of Aspirin and Domperidone was transferred into 10 ml volumetric flasks separately and dissolved in 5 ml of acetonitrile and sonicate for 10 min., then volume was made up to 10 ml with acetonitrile. Concentration of Aspirin and Domperidone in acetonitrile was 1000µg/ml. (stock- A).

2. Preparation of Sub Stock Solution

1 ml of solution was taken from stock-A of Aspirin and Domperidone and transferred into 10 ml volumetric flask separately and diluted up to 10 ml with diluent (Acetonitrile) to give concentration of 100µg/ml (Stock-B).

3. Preparation of Different Solution

1ml, 2ml, 3ml, 4ml and 5ml of stock-B was taken separately in 10 ml volumetric flask and volume was made up to 10ml with (Acetonitrile). This gives the solutions of 10 µg/ml, 20 µg/ml, 30 µg/ml, 40µg/ml, 50µg/ml for Aspirin. In same manner 1µg/ml, 2µg/ml, 3µg/ml, 4µg/ml, 5µg/ml of Domperidone also prepared.[7-8]

4. Linearity and Calibration Graph

To establish the linearity of analytical method, a series of dilution ranging from 10-50 µg/ml was prepared for aspirin and 1-5µg/ml for Domperidone. All the solutions were filtered through 0.2 µm membrane filter and injected, chromatograms were recorded at 231 nm and it was repeat for three times. A calibration graph was plotted between the mean peak area and respective concentration and regression equation was derived.

System Suitability Parameters

Separation variables were set and mobile phase was allowed to saturate the column at 1.00 ml/min. After complete saturation of column, three replicates of working standard of Domperidone 5 µg/ml and 50 µg/ml Aspirin was injected separately. Peak report and column performance report were recorded for all chromatogram.[12]

RESULT AND DISCUSSION

Identification and Characterization of drugs

IR spectrum of Aspirin and Domperidone

The IR spectrum of sample drug shows the peak values which are characteristics of the drug

and the graph were shown in figure 1-2.

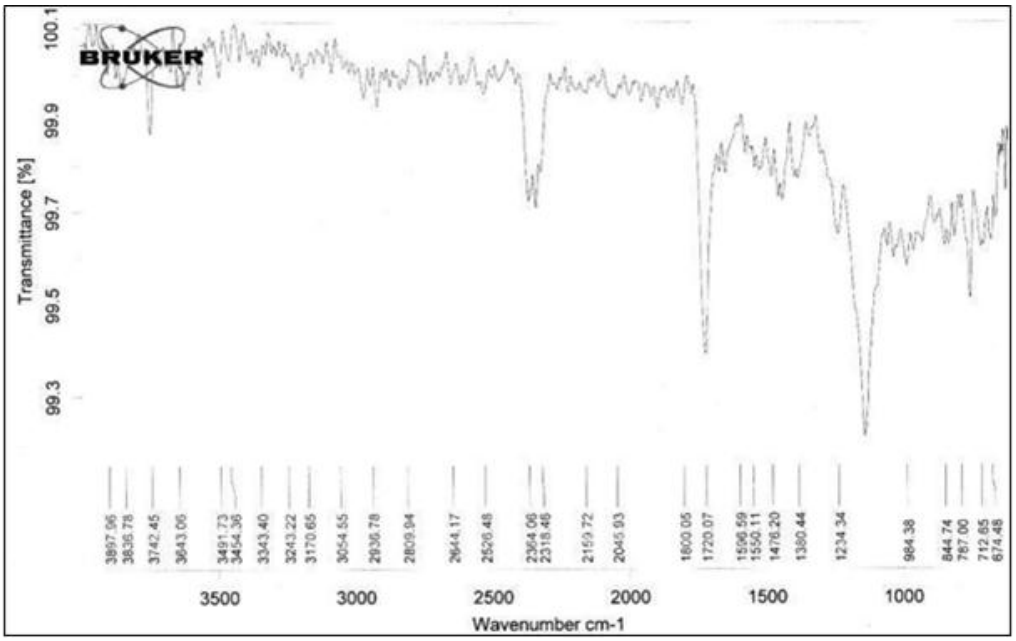

Figure 1: FT-IR Spectrum of Pure Drug (Aspirin).

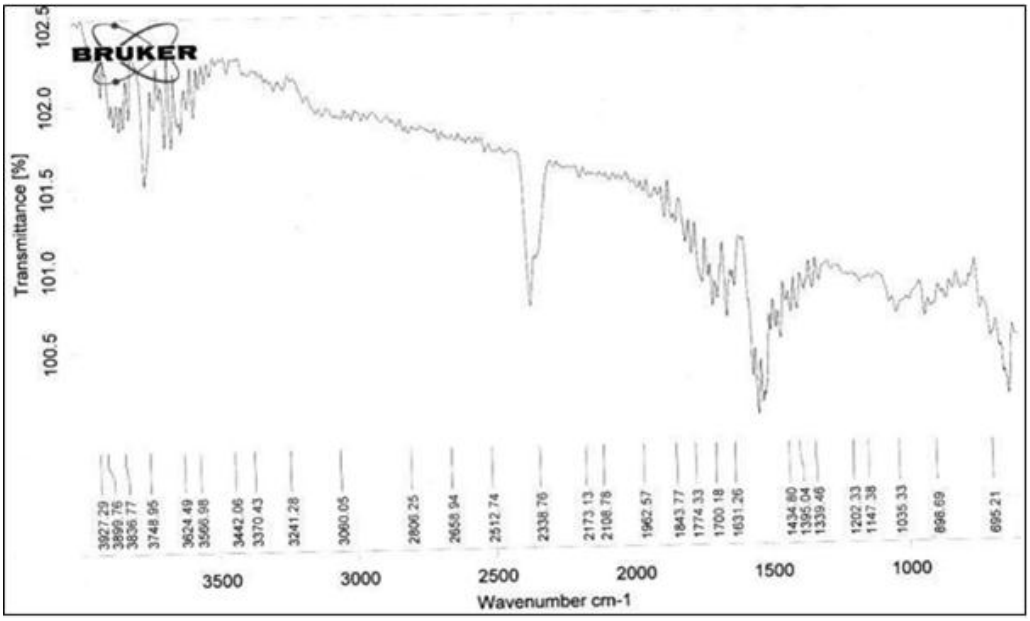

Figure 2: FT-IR Spectrum of Pure Drug (Domperidone).

Determination of λ max of Aspirin and Domperidone

Standard solution (10µg/ml) of Aspirin and Domperidone was prepared. The pure drug solution was scanned on UV spectrophotometer, and λmax was determined.

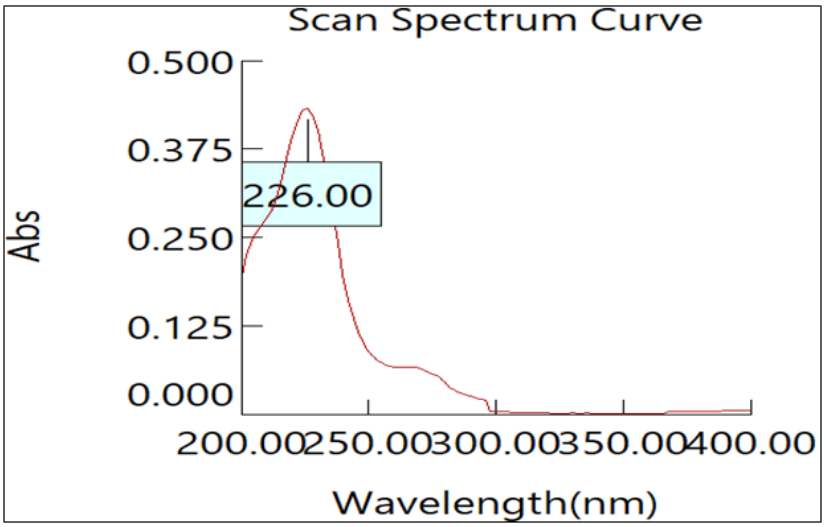

Figure 3: Determination of λmax of Aspirin.

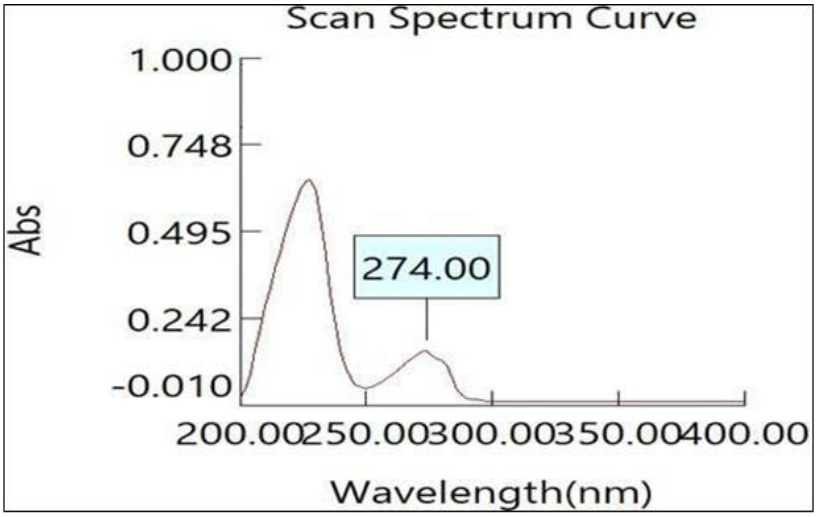

Figure 4: Determination of λmax of Domperidone.

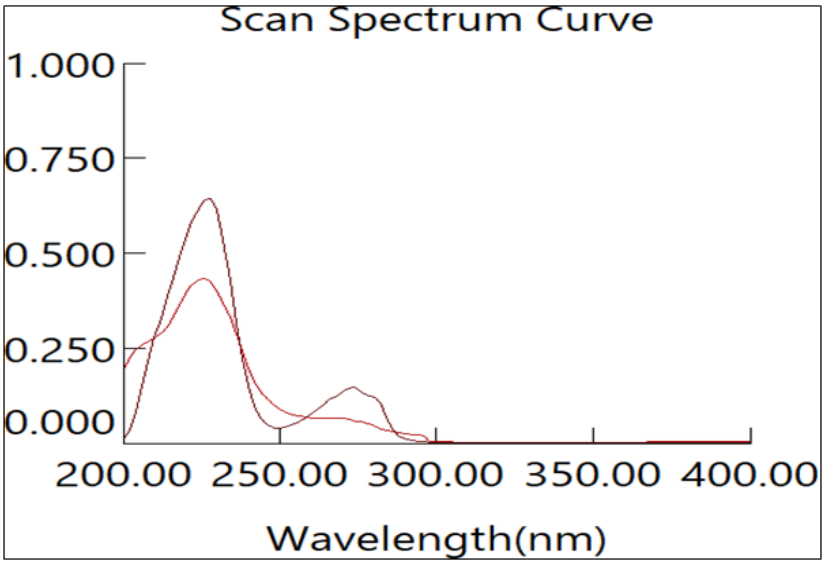

Figure 5: Overlain spectra of Aspirin and Domperidone.

Method development of Aspirin and Domperidone

Method development and validation for estimation of Aspirin and Domperidone using RP- HPLC

Selection of Mobile Phase

Initially to estimate Aspirin and Domperidone in fix dosage form number of mobile phase in different ratio were tried. A result was shown in Table 1.

Taking into consideration the system suitability parameter like RT, Tailing factor, No. of theoretical plates and HETP, the mobile phase found to be most suitable for analysis was 10 mM KH2PO4: acetonitrile (pH 3.5 with OPA) in the ratio of 20:80v/v. The mobile phase was filtered through 0.45 µ filter paper to remove particulate matter and then degassed by sonication. Flow rate employed for analysis was 1.0 ml/min.

Procedure for preparation of mobile phase

10mM KH2PO4: acetonitrile (pH 3.5) in the ratio of 20:80v/v, pH 3.0 with Ortho phosphoric acid. Filtered through 0.45 µ filter paper.

Table 1: Mobile Phase Selection.

Solvent	Ratio	Observation
10 mM KH2PO4: acetonitrile (pH 3.5)	20:80 v/v	Both sharp peak was observed (Most suitable)

Selection of Diluent

Diluent used for preparation of sample were compatible with mobile phase and no any significant affect retention and resolution of analyte. After various trials Acetonitrile was used as diluents.

Selection of separation variable:Table 2: Separation Variable

Variable	Condition
Particle Size	5 µ
Bonded Phase	Octadecylsilane (C18)
Mobile Phase	10 mM KH2PO4: acetonitrile (pH 3.5)
10mM KH2PO4	20
Acetonitrile	80
Diluent	Acetonitrile
Flow rate	1.0 ml/min
Temperature	Ambient
Sample Size	20 µ
Detection wavelength	231 mm
Aspirin	2.112 ± 0.3 min
Domperidone	4.316 ± 0.3 min

Preparation of standard Stock solution

Accurately weighed 10 mg of Aspirin and Domperidone was transferred into 10 ml volumetric flasks separately and dissolved in 5 ml of acetonitrile and sonicate for 10 min., then volume was made up to 10 ml with acetonitrile. Concentration of Aspirin and Domperidone in acetonitrile was 1000µg/ml. (stock- A).

Preparation of Sub Stock Solution

1 ml of solution was taken from stock-A of Aspirin and Domperidone and transferred into 10 ml volumetric flask separately and diluted up to 10 ml with diluent (Acetonitrile) to give concentration of 100µg/ml (Stock-B).

Preparation of Different Solution 1 ml, 2 ml, 3 ml, 4 ml and 5 ml of stock-B was taken separately in 10 ml volumetric flask and volume was made up to 10ml with (Acetonitrile). This gives the solutions of 10µg/ml, 20µg/ml, 30µg/ml, 40µg/ml, 50µg/ml for Aspirin. In same manner 1µg/ml, 2µg/ml, 3µg/ml, 4µg/ml, 5µg/ml of Domperidone also prepared.

Linearity and Calibration Graph

To establish the linearity of analytical method, a series of dilution ranging from 10-50µg/ml was prepared for aspirin and 1-5µg/ml for Domperidone. All the solution were filtered through 0.2µm membrane filter and injected, chromatograms were recorded at 231 nm and it was repeat for three times. A calibration graph was plotted between the mean peak area and respective concentration and regression equation was derived.

Table: Linearity of Aspirin.

Standard Concentration g/ml	Area under Curve (AUC)						Mean
	Rep-1	Rep-2	Rep-3	Rep-4	Rep-5	Rep-6	
0	**0**	0	0	0	0	0	0
10	235.654	240.565	229.896	230.478	245.587	231.658	235.640
20	465.658	475.658	460.587	455.698	472.125	479.985	468.285
30	699.854	705.658	713.458	445.569	685.458	679.985	654.997
40	935.471	925.698	930.145	940.587	928.741	920.325	930.161
50	1170.658	1165.254	1169.987	1176.654	1175.654	1180.325	1173.089
Correl Coeff (r^2)							0.998
Slope (m)							23.24
Intercept (c)							-4.094

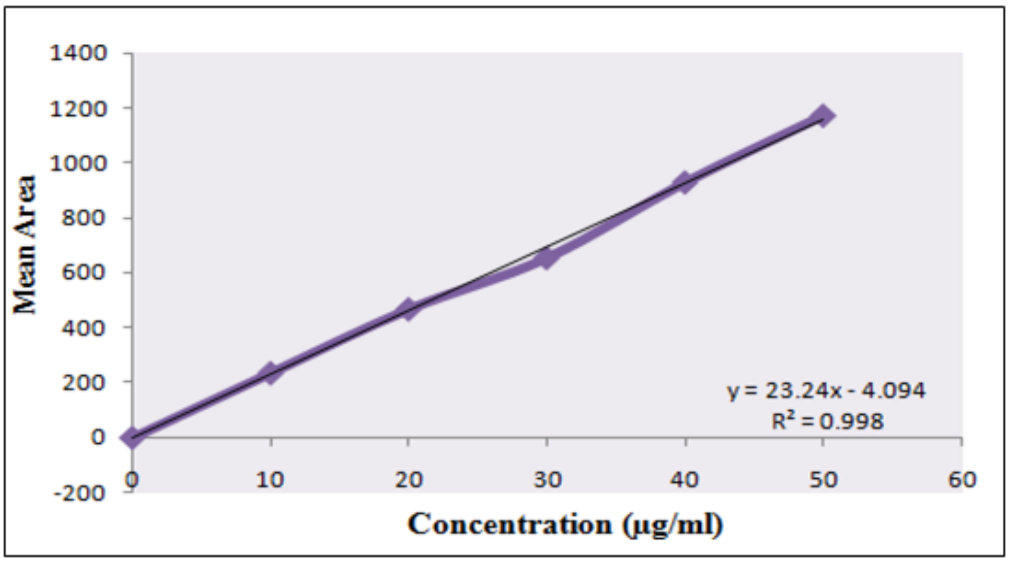

Figure: Calibration Curve of Aspirin.

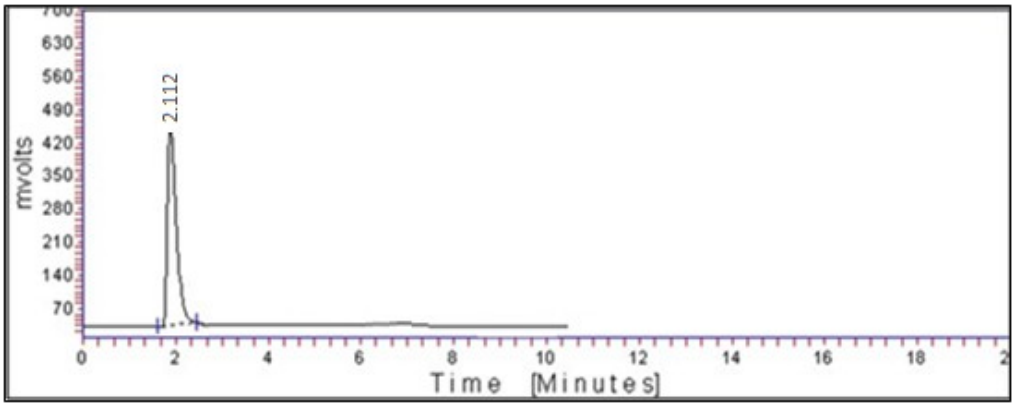

Figure: Chromatogram of Aspirin.

Table: Linearity of Domperidone.

Standard Concentration g/ml	Area under Curve (AUC)						Mean
	Rep-1	Rep-2	Rep-3	Rep-4	Rep-5	Rep-6	
0	**0**	0	0	0	0	0	0
1	95.569	98.856	92.325	98.789	96.658	93.321	95.920
2	186.658	180.125	189.658	182.325	175.658	193.321	184.624
3	276.458	270.325	275.658	265.589	273.325	269.987	271.890
4	365.458	360.254	369.987	345.658	372.325	360.458	362.357
5	465.581	470.586	475.658	460.325	470.325	478.954	470.238
Correl Coeff (r^2)							0.998
Slope (m)							92.50
Intercept (c)							-0.430

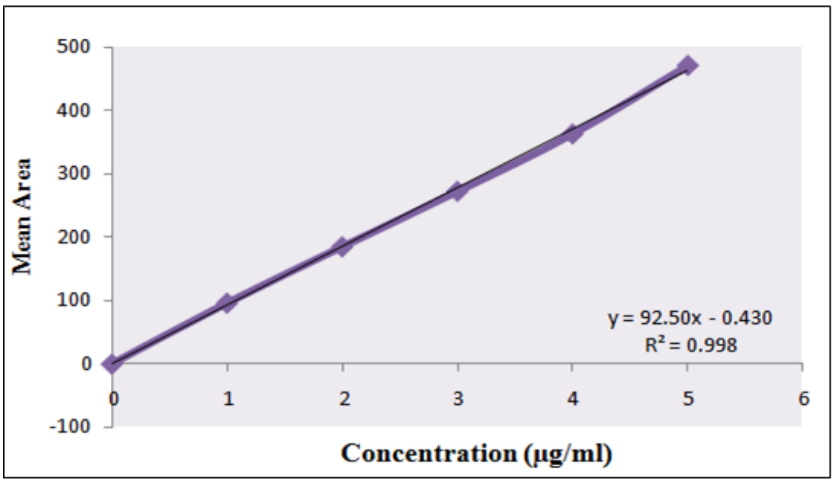

Figure: Calibration Curve of Domperidone.

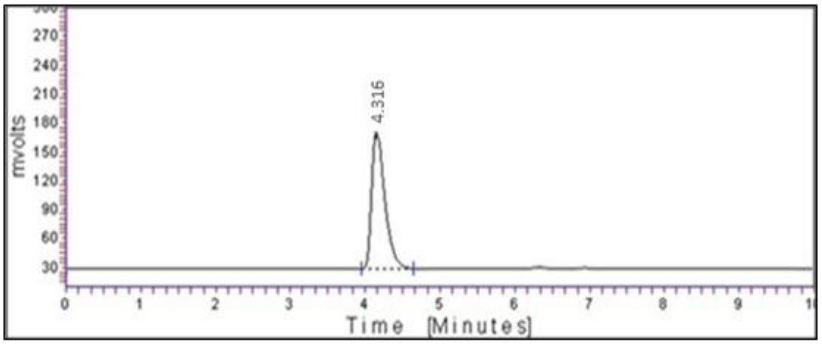

Figure: Chromatogram of Domperidone.

System Suitability Parameters

Separation variables were set and mobile phase was allowed to saturate the column at 1.00ml/min. After complete saturation of column, three replicates of working standard of Domperidone 5mg/ml and 50mg/ml Aspirin was injected separately. Peak report and column performance report were recorded for all chromatogram.

Table: System Suitability Parameters of Aspirin.

System suitability Parameter →	RT	AUC	No. of theoretical plates	Tailing factor
Rep-1	2.112	1170.658	3256	1.25
Rep-2	2.113	1165.254	3156	1.26
Rep-3	2.114	1169.987	3265	1.32
Rep-4	2.114	1176.654	3156	1.25
Rep-5	2.113	1175.654	3255	1.32
Rep-6	2.116	1180.325	3265	1.45
Mean	2.114	1173.089	3225.500	1.308
S.D.	0.001	5.451	54.003	0.077
% R.S.D.	0.065	0.465	1.674	5.869

Table: System Suitability Parameters of Domperidone.

System suitability Parameter→	RT	AUC	No. of theoretical plates	Tailing factor
Rep-1	4.316	465.581	3250	1.45
Rep-2	4.325	470.586	3150	1.46
Rep-3	4.321	475.658	3250	1.45
Rep-4	4.326	460.325	3050	1.44
Rep-5	4.329	470.325	3150	1.46
Rep-6	4.322	478.954	3250	1.52
Mean	4.323	470.238	3183.333	1.463
S.D.	0.005	6.715	81.650	0.029
% R.S.D.	0.105	1.428	2.565	1.965

CONCLUSION

In the present research work, a successful attempt was made for "Method Development and Validation for the Estimation of Domperidone and Aspirin in Bulk or Formulation Using HPLC" which was developed by experimentation based on thorough literature survey and ascertained by statistical parameters of sampling. The simplicity, rapidity, accurate and reproducibility of the proposed methods completely fulfil the objective of the research work of estimation of the drugs. Liquid chromatographic system from waters comprising of manual injector, waters 515 pumps for constant flow and constant pressure delivery and UV Visible Detector connected to data ace software for controlling the instrumentation as well as processing the data generated were used. Drug sample was extracted by precipitating method using 5ml of methanol for each ml of plasma sample. The proposed methods were found to be linear with correlation coefficient close to one. Precision was determined by repeatability, Intermediate precision and reproducibility of the drugs. The robustness of developed method was checked by changing in the deliberate variation in solvent. The result obtained shows the developed methods to be Cost effective, Rapid (Short retention time), Simple, Accurate (the value of SD and % RSD less than 2), Precise and can be successfully employed in the routine analysis of these drugs in bulk drug as well as in tablet dosage form. The Simplicity, Rapidly and Reproducibility of the proposed method completely fulfil the objective of this research work.

REFERENCES

1. Sharma BK. Instrumental methods of chemical analysis, Introduction to Analytical chemistry: Goel Publishing House Meerut, 23th edition, 2004.
2. Saeed AM, Hamzah MJ, Ahmed NQ. Quantitative assay of aspirin and (salicylic acid and heavy metals as impuraties) in Iraqi's market aspirin tablets using different analytical

methods. International Journal of Applied Pharmaceutics, 2018; 10(5): 167-172.

3. Willard HH, Merritt LL, Dean JJA, Frank AS. Instrumental method of analysis: CBS Publishers and Distributors, New Delhi, 7th Edition, 1986.
4. R Vani, M Sunitha Analytical Method Development and Validation for theDetermination of Omeprazole And Aspirin using Reverse Phase HPLC Method in Bulk and Dosage Form. Universal Journal of Pharmaceutical Research, 2017; 2(4): 25-29.
5. S. Vidyadhara, RLC Sasidhar, B.Praveen Kumar, NT Ramarao and N.Sriharita, Method Development and Validation for Simultaneous Estimation of Ranitidine andDomperidone in Pharmaceutical Dosage Forms by RP-HPLC. Oriental journal of chemistry, 2012; 28(4): 1691-1696.
6. Willard HH, Merritt LL, Dean JA, Settle FA. HPLC Theory and Instrumentation; In Instrumental Methods of Analysis, CBS Publishers and Distributors, New Delhi, 8th edn., 2002; 1-12.
7. Snyder LR, Kirkland JJ, Glajch LJ. Getting Started; In Practical HPLC Method Development, John Willey and Sons, Inc, New York, 2nd edn., 1997; 5-17.
8. Snyder LR, Kirkland JJ, Glajch LJ. Non-ionic Samples; Reversed- and Normal-Phase HPLC, In Practical HPLC Method Development, John Wiley and Sons, Inc, New York, 2nd edn., 1997; 233-291.
9. Billiet HAH, Rippel G. Method Development and Selectivity Optimization in High-Performance Liquid Chromatography; In Advances in Chromatography, Marcel Dekker, Inc, New York, 1998; 39: 263-310.
10. Sharaf MA. Assessment of Chromatographic Peak Purity; In Advances in hromatography, Market Dekker, Inc, New York, 1997; 37: 1-6.
11. Cindy Green. J Vali Teach, In; Analytical method validation. RAC, 2000; 6: 625- 31.
12. Elhance DN. Foundation of statistics. Kitab mahal, 47th edn, 2003.