1.810 and 0.649 - 1.503 and is well within the acceptance criteria of less than 2% at all the wavelength. The low value of % RSD indicates that the proposed method was precise and accurate [Figure 14 & 15, Table 5, 6].

Recovery

The percentage recovery of the drug from the synthetic mixture was found to be in the range of 98.74 - 100.26 % w/w, which was within the acceptance limit of 97 - 103 % w/w as per the ICH guideline as shown in Table 7 and Figure 16.

CONCLUSION

The suggested UV spectrophotometric method employing multivariate calibration technique was novel, uncomplicated, accurate, precise, profitable and sensitive for the quantification of Zaleplon in its pharmaceutical formulations. Hence, this method is very useful with very simple mathematical contents, is more reliable than the other spectrophotometric methods and strongly recommends the developed method for a routine analysis of Zaleplon in pharmaceutical formulations.

ACKNOWLEDGEMENT

Authors are thankful to Vice Chancellor, SRM University and management of SRM College of Pharmacy, SRM University, Kattankulathur for providing various reprographic sources for carrying out this work.

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