



Corrigendum

Corrigendum to “A validated stability indicating UPLC method for desloratadine and its impurities in pharmaceutical dosage forms”

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The authors regret the following error:

3.2.5 Accuracy

The percentage recovery of desloratadine from tablets was ranged from 98.9 to 100.5%. The percentage recovery of impurities in desloratadine samples varied from 97.3 to 101.3%. The LC chromatogram of spiked sample at 0.20% level of all five impurities in desloratadine tablets sample is shown in Fig. 3. The % recovery values for desloratadine and impurities are presented in Table 3.

The authors would like to apologise for any inconvenience this may have caused to the readers of the journal.

Table 3

Evaluation of accuracy.

Amount spiked ^a	% Recovery ^b					
	Desloratadine	Imp-A	Imp-B	Imp-C	Imp-D	Imp-E
LOQ	98.9 ± 0.89	98.4 ± 0.83	98.6 ± 0.52	99.5 ± 0.61	98.2 ± 0.71	99.6 ± 0.51
50%	100.5 ± 0.25	99.7 ± 0.72	98.1 ± 0.66	98.1 ± 1.12	98.0 ± 0.77	101.3 ± 0.43
100%	99.3 ± 0.31	97.6 ± 0.56	98.8 ± 0.44	99.8 ± 0.45	98.9 ± 0.75	98.1 ± 0.69
150%	100.1 ± 0.15	99.0 ± 0.48	98.1 ± 0.27	99.5 ± 0.72	97.3 ± 1.66	97.3 ± 0.63

^a Amount of five impurities spiked with respect to 0.20% specification level individually to 0.5 mg/ml of desloratadine.^b Mean ± % RSD for three determinations.DOI of original article: [10.1016/j.jpba.2009.09.016](https://doi.org/10.1016/j.jpba.2009.09.016).

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