



Spectrophotometric Studies on Determination of Tenoxicam in Pharmaceutical Formulations via Complexation with Thorium (IV) Ion

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Introduction









Tenoxicam is a novel NSAID often used for treating musculoskeletal, and joint disorders, analgesic and relief of post-surgical inflammation.

The most common tenoxicam determination techniques are costly, time-consuming, and chemically intensive. Therefore, a **simple, direct, inexpensive,** and **accurate** method to assay tenoxicam in dosage forms.



3D Chemical Structure of Tenoxicam

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• **Bathochromic shifting** of Tenoxicam by complexation to maximum UV absorbance wavelength (λ max).

 The main objective of this study is the development of a new and simple analytical method for the determination of Tenoxicam in selected Pharmaceutical formulations.





04 Experimental Design



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Results and Discussion



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That means the TXM is present in the solution in a Zwitterionic state (ZWC). The chelating reaction of

Tenoxicam with metal ions can be expected through three coordination sites (-OH, —CONH, and N pyridine Ring). (Mamdouh S Masoud et al., 2020).



Fig 2: Structure For Dissociation Forms of TNX In Solution





Fig 3: The mechanism of the formed complex (illustrates phase I and phase II of the formed complex. (Mamdouh S Masoud *et al.*, 2020).





Effect of pH.

This complex is studied within a pH range of 4.5 to 12.3.



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Effect of temperature

UV absorbance of a TNX-TH4+ complex generally decreases with increasing

temperature, (According to the study (Ito, 1960), higher temperatures cause changes in the **electronic transitions** and **molecular structure** leading to dissociation and precipitation of the complex.







Assay of the content of TNX in selected Pharmaceutical Preparations for marketed brands.

The proposed analytical method for TNX determination in Tablets and suppositories.

Table 1: Assay of the content of TNX in(Tablet and suppositories)

Dosage form	Content (mg)	Absorbance	Founded concentration (mg)	Recovery (%)
Tablet	20	0.301	20.12	100.65%
Suppositories	20	0.285	19.77	98.85%







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Limitation and Future Perspective



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Limitation:

• There are limited options for tenoxicam-containing formulations only tablets,

and suppositories are available.

> Future prospectives:

• Applying the assay method to test the contents of different generic

Pharmaceutical products of tenoxicam.

• Extensibility of the assay method to other oxicam derivatives.

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• Based on the experimental results, could be concluded:

The development analytical method is considered a novel approach for the determination of tenoxicam in tablets and suppositories.

The proposed spectrophotometric method is accurate (average recovery range 98.85-100.96%)

The proposed method was applied successfully for the assay of the Tenoxicam in selected pharmaceutical products















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