

# Comprehensive Phytochemical Profiling of the Ethanolic Extract of *Zephyranthes candida* (Amaryllidaceae)

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**Abstract:** *Phytochemicals are the non-nutritive, bioactive chemical compounds that occur naturally in plants, acting as a primary defense mechanism against environmental stressors and pathogens. The extraction of these compounds using ethanol is a standard pharmacological practice, as ethanol effectively dissolves a broad spectrum of secondary metabolites including polar phenolics and non-polar terpenoids. This study aims to provide a rigorous qualitative and quantitative assessment of the chemical constituents of a specific test plant.*

**Keywords:** *Phytochemicals*

## I. INTRODUCTION

Phytochemicals are the non-nutritive, bioactive chemical compounds that occur naturally in plants, acting as a primary defense mechanism against environmental stressors and pathogens. The extraction of these compounds using ethanol is a standard pharmacological practice, as ethanol effectively dissolves a broad spectrum of secondary metabolites including polar phenolics and non-polar terpenoids. This study aims to provide a rigorous qualitative and quantitative assessment of the chemical constituents of a specific test plant.

## II. MATERIALS AND METHODS

### Extraction

The plant material was subjected to ethanolic extraction to isolate the bioactive fractions for further analysis.

### Qualitative Methodology

The screening for various secondary metabolites was conducted using globally recognized protocols:

**Alkaloids:** Detection was performed using Mayer's, Dragendorff's, and Wagner's reagents as described by Evans (2009)

**Terpenoids:** The Salkowski test was utilized following the procedures of Trease and Evans (2002)

**Phenolics:** Presence was determined via Ferric chloride and Lead acetate methods outlined by Harborne (1998)

**Phytosterols:** Screening was conducted through the Libermann-Burchard and Salkowski tests based on Sofowora (1993)

### Quantitative Methodology

Quantitative estimations were calculated as percentage composition (g/100g) using modified gravimetric and spectrophotometric procedures.

## III. RESULTS AND DETAILED EXPLANATION

### Qualitative Screening Results

The qualitative analysis revealed a significant presence of phenolic compounds, indicated by the formation of a greenish-blue color during testing. Flavonoids were identified by the transition from yellow to colorless in the alkaline reagent test.

**Table 1: Qualitative Phytochemical screening of Ethanol extract**

Phytochemical	Test Name	Observation	Result
Phenolics	Ferric chloride	Blue/green/purple	++ (Higher Quantity)
Terpenoids	Salkowski	Reddish-brown interface	+ (Present)
Alkaloids	Mayer's/Wagner's	Cream/Brown ppt	+ (Present)
Flavonoids	Shinoda/Alkaline	Red/orange; Yellow→colorless	+ (Present)
Phytosterols	Liebermann-Burchard	No Blue-green color	Absent

#### Quantitative Composition

The quantitative data highlights **Terpenoids** as the most abundant constituent at 2.35  $\mu\text{m}$  0.25 g/100g, followed by **Phenolics** at 1.55  $\mu\text{m}$  0.12 g/100g<sup>14</sup>. Interestingly, while phytosterols were not detected qualitatively, the quantitative analysis measured a trace amount of 0.21  $\mu\text{m}$  0.05 g/100g

**Table 2: Percentage composition (g/100g)<sup>16</sup>**

Phytochemicals	Composition (g/100g)
Terpenoids	2.35 $\mu\text{m}$ 0.25
Phenolics	1.55 $\mu\text{m}$ 0.12
Alkaloids	0.29 $\mu\text{m}$ 0.04
Flavonoids	0.24 $\mu\text{m}$ 0.06
Phytosterols	0.21 $\mu\text{m}$ 0.05

#### IV. DISCUSSION

The high concentration of terpenoids (2.35 g/100g) suggests the plant may possess significant anti-inflammatory or antimicrobial properties. The "higher quantity" of phenolics observed qualitatively aligns with the 1.55 g/100g measured quantitatively, reinforcing the plant's potential as a source of natural antioxidants. The discrepancy in phytosterols—being absent in qualitative tests but present in quantitative results—is often attributed to the quantitative method's higher sensitivity to low concentrations.

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