

RESEARCH MONOGRAPH . COMPOUND PROFILE

F-Phenibut (Fluorophenibut)

A 4-fluoro-substituted phenibut analog with higher GABA-B affinity and lower dose requirement; sold as a research chemical with elevated dependence concerns.

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ABSTRACT

F-Phenibut (4-amino-3-(4-fluorophenyl)butanoic acid) is a fluorinated analog of phenibut with the addition of a para-fluoro substituent on the phenyl ring. The structural modification increases GABA-B receptor affinity and produces effective doses approximately 5- to 10-fold lower than phenibut. The compound is not approved in any jurisdiction and is sold as a research chemical. Mechanism parallels phenibut (GABA-B agonist) with the affinity increase. Dependence liability is at least as substantial as phenibut and may be greater on a dose-equivalent basis. Pharmacokinetics are not formally characterized in humans; rodent half-life is similar to phenibut. Research-grade vendor doses are typically 50 to 250 mg per administration. F-phenibut should be regarded as a higher-potency phenibut analog with all the dependence concerns of phenibut amplified by potency.

1. Compound Identification

F-Phenibut (Fluorophenibut) is described as gaba-b agonist / fluoro-phenibut. A 4-fluoro-substituted phenibut analog with higher GABA-B affinity and lower dose requirement; sold as a research chemical with elevated dependence concerns.

1.1 Class summary

GABA-B agonist / fluoro-phenibut

1.2 Subtitle

A 4-fluoro-substituted phenibut analog with higher GABA-B affinity and lower dose requirement; sold as a research chemical with elevated dependence concerns.

2. Abstract

F-Phenibut (4-amino-3-(4-fluorophenyl)butanoic acid) is a fluorinated analog of phenibut with the addition of a para-fluoro substituent on the phenyl ring. The structural modification increases GABA-B receptor affinity and produces effective doses approximately 5- to 10-fold lower than phenibut. The compound is not approved in any jurisdiction and is sold as a research chemical. Mechanism parallels phenibut (GABA-B agonist) with the affinity increase. Dependence liability is at least as substantial as phenibut and may be greater on a dose-equivalent basis. Pharmacokinetics are not formally characterized in humans; rodent half-life is similar to phenibut. Research-grade vendor doses are typically 50 to 250 mg per administration. F-phenibut should be regarded as a higher-potency phenibut analog with all the dependence concerns of phenibut amplified by potency.

3. Mechanism of Action

Higher-affinity GABA-B agonist than phenibut; same general pharmacology.

The mechanistic profile described above represents the consensus position from peer-reviewed literature where available; for compounds with sparse formal characterization, this section reflects vendor-grade reports and structurally-related class pharmacology and should be interpreted with appropriate caution.

4. Pharmacokinetics and Routes of Administration

Pharmacokinetic data are summarized within the abstract for compounds with formal Phase 1 records. Compounds without published human pharmacokinetic studies are noted as such; investigators should treat the absence of data as a research limitation.

5. Reported Research Dose Ranges

50 to 250 mg per administration (research-grade).

Doses are scaled from clinical use where the compound is approved as a medicine, from Phase 2/3 trials where development advanced, or from rodent dose-response data scaled by allometric assumptions where the compound has no human clinical record. Investigators should treat dose ranges from research-chemical-grade vendors as starting points for their own pharmacokinetic and tolerability characterization rather than validated dosing recommendations.

6. Sourcing and Quality Verification

Investigators should obtain analytical confirmation of identity (mass spectrometry; for peptides, additionally amino acid analysis or N-terminal sequencing) and purity (HPLC area-percent, ideally greater than 95 percent) on every lot. Vendor certificates of analysis vary in quality; third-party analytical confirmation through independent laboratories (e.g., Janoshik Analytical, Anabolic Lab) is the gold standard for research-grade peptides and small molecules.

For compounds with established pharmaceutical-grade marketing in any jurisdiction, sourcing from the marketed product (where legal and accessible) provides the highest quality assurance.

7. Reconstitution and Handling

Peptides should be reconstituted in bacteriostatic water or sterile water for injection in the volume appropriate to the desired concentration; lyophilized peptides should be allowed to come to room temperature before reconstitution to minimize precipitation. Reconstituted peptides should be stored refrigerated and used within the timeframe specified by the manufacturer (typically 14 to 30 days for bacteriostatic water reconstitutions, shorter for sterile water).

Small molecules: storage conditions vary by compound; consult the certificate of analysis for the specific product. Most are stable at room temperature in dry, dark conditions; oxidation-sensitive compounds (e.g., catecholamine precursors) require cool storage and protection from light.

8. Stack Interaction Considerations

Investigators combining this compound with other research substances should consider receptor-level interactions (additive effects at shared targets), enzyme-level interactions (CYP-mediated drug-drug interactions where applicable), and pharmacodynamic interactions (additive sedation, additive serotonergic activity, additive hemodynamic effects).

The stack interaction section in published monographs of well-characterized compounds is necessarily incomplete for any specific combination; investigators should treat novel combinations with elevated caution and titrate gradually.

9. References

1. Sparse formal literature; phenibut analog pharmacology papers.

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