

RESEARCH MONOGRAPH . COMPOUND PROFILE

Rolziracetam

A bicyclic piracetam analog with sparse published literature; primarily of historical and research-chemical interest.

DOCUMENT	KDC-MN-040
ISSUED	2026-05-09
REVISION	1.0
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COMPOUND CLASS	Bicyclic pyrrolidinone racetam

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ABSTRACT

Rolziracetam is a bicyclic racetam analog with sparse published literature and no clinical development record. The compound features a fused bicyclic scaffold built on the pyrrolidinone core. Mechanism is presumed to overlap with piracetam pharmacology but has not been formally characterized. The compound is occasionally available from research chemical vendors; identity confirmation by mass spectrometry is essential before any investigational use. There is no published human pharmacokinetic data, no Phase 1 safety record, and no peer-reviewed dose-finding studies. This entry is included for completeness of the racetam class survey and as a marker that "racetam" as a class label spans well-characterized compounds (piracetam, levetiracetam) and obscure analogs with thin evidence bases.

1. Compound Identification

Rolziracetam is described as bicyclic pyrrolidinone racetam. A bicyclic piracetam analog with sparse published literature; primarily of historical and research-chemical interest.

1.1 Class summary

Bicyclic pyrrolidinone racetam

1.2 Subtitle

A bicyclic piracetam analog with sparse published literature; primarily of historical and research-chemical interest.

2. Abstract

Rolziracetam is a bicyclic racetam analog with sparse published literature and no clinical development record. The compound features a fused bicyclic scaffold built on the pyrrolidinone core. Mechanism is presumed to overlap with piracetam pharmacology but has not been formally characterized. The compound is occasionally available from research chemical vendors; identity confirmation by mass spectrometry is essential before any investigational use. There is no published human pharmacokinetic data, no Phase 1 safety record, and no peer-reviewed dose-finding studies. This entry is included for completeness of the racetam class survey and as a marker that "racetam" as a class label spans well-characterized compounds (piracetam, levetiracetam) and obscure analogs with thin evidence bases.

3. Mechanism of Action

Presumed overlap with piracetam; no formal characterization.

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

No reliable data.

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. No substantial peer-reviewed published references for this specific analog.

This monograph is provided for research use only. Not for medical, diagnostic, or therapeutic purposes. Not for human consumption. Compound class generalizations are noted where compound-specific data are sparse; investigators should weight the evidence accordingly.