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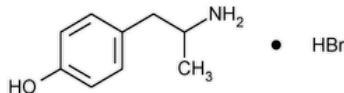
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## Hydroxyamphetamine Hydrobromide

 $C_9H_{13}NO \cdot HBr$  232.12

Phenol, 4-(2-aminopropyl)-, hydrobromide.

 $(\pm)$ -*p*-(2-Aminopropyl)phenol hydrobromide CAS RN®: 306-21-8; UNII: 59IG47SZ0E.» Hydroxyamphetamine Hydrobromide contains not less than 98.0 percent and not more than 101.5 percent of  $C_9H_{13}NO \cdot HBr$ , calculated on the dried basis.**Packaging and storage**—Preserve in well-closed, light-resistant containers.**USP REFERENCE STANDARDS (11)**—[USP Hydroxyamphetamine Hydrobromide RS](#)**Identification**—**Change to read:****A:** [▲ Spectroscopic Identification Tests \(197\), Infrared Spectroscopy: 197K▲](#) (CN 1-May-2020) .**B:** Dissolve about 500 mg of ammonium molybdate in 10 mL of sulfuric acid, and add to this solution about 2 mg of Hydroxyamphetamine Hydrobromide: an intense blue color is produced (*distinction from similar amino compounds such as amphetamine and methamphetamine, which, lacking a phenolic hydroxyl, do not undergo this reaction*).**C:** Dissolve about 200 mg in 2 mL of water, and add a solution of 500 mg of potassium carbonate in 2 mL of water. Extract with two 10-mL portions of ether, allow the clear ether solution to evaporate to dryness, and dry at about 80°: the hydroxyamphetamine so obtained melts between 124° and 127° (see *Class I* under [Melting Range or Temperature \(741\)](#)).**D:** To a solution of about 10 mg of it in 10 mL of water add 1 mL of 2 N nitric acid, then add silver nitrate TS: a pale yellow precipitate is formed, and it is slightly soluble in 6 N ammonium hydroxide.**MELTING RANGE (741):** between 189° and 192°.**LOSS ON DRYING (731):**—Dry it at 105° for 2 hours: it loses not more than 0.5% of its weight.**RESIDUE ON IGNITION (281):** not more than 0.1%.**Bromide content**—Accurately weigh about 400 mg, and dissolve in 50 mL of water. Add 50 mL of methanol and 10 mL of glacial acetic acid, then add eosin Y TS, and titrate with 0.1 N silver nitrate VS. Each mL of 0.1 N silver nitrate is equivalent to 7.990 mg of Br: the content of Br, calculated on the dried basis, is between 33.6% and 35.2%.**ORDINARY IMPURITIES (466)**—**Test solution:** methanol.**Standard solution:** methanol.**Eluant:** a mixture of toluene, methanol, and ammonium hydroxide (10:4:0.25).**Visualization:** 1.**Assay**—Dissolve about 400 mg of Hydroxyamphetamine Hydrobromide, accurately weighed, in a mixture of 10 mL of glacial acetic acid and 10 mL of mercuric acetate TS, warming slightly, if necessary, to effect solution. Add crystal violet TS, and titrate with 0.1 N perchloric acid VS. Perform a blank determination, and make any necessary correction. Each mL of 0.1 N perchloric acid is equivalent to 23.21 mg of  $C_9H_{13}NO \cdot HBr$ .**Auxiliary Information** - Please [check for your question in the FAQs](#) before contacting USP.

Topic/Question	Contact	Expert Committee
HYDROXYAMPHETAMINE HYDROBROMIDE	<a href="#">Documentary Standards Support</a>	SM32020 Small Molecules 3
REFERENCE STANDARD SUPPORT	RS Technical Services <a href="mailto:RSTECH@usp.org">RSTECH@usp.org</a>	SM32020 Small Molecules 3

**Chromatographic Database Information:** [Chromatographic Database](#)

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