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# **ProductInformation**

## p-Aminohippuric acid sodium salt

Product Number **A3759**Store at Room Temperature

## **Product Description**

Molecular Formula:  $C_9H_9N_2NaO_3$ Molecular Weight: 216.2 CAS Number: 94-16-6 pK<sub>a</sub> of conjugate acid: 3.6<sup>1</sup>

Melting point: 123-125 °C

Synonyms: 4-Aminohippuric acid sodium salt, N-(4-Aminobenzoyl)glycine sodium salt, PAH.

p-Aminohippuric acid (PAH) is a liver metabolite of p-aminobenzoic acid (PABA). PAH is formed by conjugation of PABA and glycine. The ability of the liver to convert PABA to PAH is a way of measuring liver function.<sup>2</sup> An HPLC method for the determination of PAH and PABA levels in urine has been published.<sup>3</sup>

This product has been used to measure renal function by two different methods. One method which uses PAH and inulin to measure renal flow is an enzymatic assay for inulin. The other is a direct assay for this compound which measures renal flow. Plasma clearance of aminohippurate is considered to be equal to the effective renal plasma flow. The mechanism by which this compound is secreted by the kidneys involves an organic anion transporter. This compound causes natriuresis (excessive loss of sodium in the urine). The mechanism for the natriuretic effect has been determined.

The effect of this compound on the central nervous system (CNS) distribution of the anti-AIDS drugs ddC and AZT has been studied.<sup>8</sup>

#### **Precautions and Disclaimer**

For Laboratory Use Only. Not for drug, household or other uses.

## **Preparation Instructions**

This product is soluble in water (50 mg/ml), yielding a clear, colorless solution. A USP formulation of this compound consists of an aqueous solution (200 mg/ml) of PAH free acid prepared with the addition of sodium hydroxide; it has a pH of 6.7-7.6. This compound is also soluble in dilute hydrochloric acid (200 mg/ml, with decomposition). It is freely soluble in alkaline solutions (with decomposition).

## Storage/Stability

A 1% solution of this compound at pH 7 has been found to stable after storage for one week at 80 °C.9

#### References

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- 8. Gibbs, J. E., and Thomas, S. A., The distribution of the anti-HIV drug, 2'3'-dideoxycytidine (ddC), across the blood-brain and blood-cerebrospinal fluid barriers and the influence of organic anion transport inhibitors. J. Neurochem., 80(3), 392-404 (2002).
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IRB/RXR 11/03