Enzymatic synthesis of radiolabeled phosphonoacetaldehyde

Guofeng Zhang, a Karen N. Allen, b and Debra Dunaway-Mariano a,*

a Department of Chemistry, University of New Mexico, Albuquerque, NM 87131-0001, USA
b Department of Physiology and Biophysics, Boston University School of Medicine, Boston, MA 02118-2394, USA

Received 11 June 2003

Abstract

Phosphonoacetaldehyde (Pald) is formed in a variety of biosynthetic pathways leading to natural phosphonates and is an intermediate in the degradation pathway of the natural product 2-aminoethylphosphonate. To facilitate the investigation of the enzymes catalyzing these pathways, a method for the synthesis of radiolabeled Pald was developed. The enzyme pyruvate phosphate dikinase was used to prepare phosphoenolpyruvate (PEP) from pyruvate, adenosine triphosphate (ATP), and orthophosphate. Then PEP was converted to phosphonopyruvate (Ppyr) with PEP mutase and then to Pald with Ppyr decarboxylase. By using [\(\beta\)-32P]ATP or [2-14C]pyruvate as precursor, [32P]Pald or [1-14C]Pald was obtained, respectively. The utilization of the synthetic, radiolabeled Pald as a probe of enzyme mechanism was demonstrated with the enzyme phosphonoacetaldehyde hydrolase (trivial name phosphonatase). The single turnover time course for the formation and consumption of radiolabeled covalent enzyme species evidenced a kinetically competent covalent intermediate.

© 2003 Elsevier Inc. All rights reserved.

Keywords: Phosphonate; Phosphonoacetaldehyde; [32P]Pald; [14C]Pald; Radiolabel; Enzymatic synthesis

Radiolabeled probes have applications in biological studies, such as tracing the distribution and usage of a metabolite in an organism, monitoring the absorption and excretion kinetics of drugs in the body, and identifying the chemical steps of metabolic pathways. At the enzyme level, radiolabeled substrates are used in activity assays and in determination of the mechanism of catalysis. Mechanistic studies that focus on the characterization of reaction intermediates often rely on the use of radiolabeled reactants in combination with rapid quench technology [1].

Phosphonoacetaldehyde (Pald) \(^1\) is an intermediate in a variety of biosynthetic pathways leading to natural phosphonates and an intermediate in the degradation pathway of the most ubiquitous phosphonate, 2-aminoethylphosphonate (Fig. 1) [2,3]. Phosphonates are synthesized in microbes and lower animals to carry out specialized functions which include microbe eradication, cell–cell signaling, host infection and/or persistence, or phosphorus storage (reviewed in [4] and references therein). To facilitate the study of the biosynthetic pathways leading to phosphonate natural products and the enzymes catalyzing these pathways, radiolabeled Pald was prepared as a probe.

Here, we report an enzymatic synthetic pathway leading to [32P]Pald and [1-14C]Pald. To demonstrate their application in enzyme mechanism studies these two probes were used to identify a covalent enzyme intermediate formed during Pald hydrolysis catalyzed by the enzyme phosphonoacetaldehyde hydrolase (hereafter called phosphonatase; EC 3.11.1.1).

Materials and methods

Materials

[2-14C]pyruvic acid sodium salt (specific activity (SA) = 23 mCi/mmol) and [\(\beta\)-32P]ATP (SA = 7000 mCi/mmol) were purchased from Perkin–Elmer Life sciences.

---

\(^1\) Abbreviations used: Pald, phosphonoacetaldehyde; SA, specific activity; PPDK, pyruvate phosphate dikinase; Ppyr, phosphonopyruvate; PEP, phosphoenolpyruvate; P, orthophosphate; PP, pyrophosphate; cpm, counts per minute.
Recombinant *Clostridium symbiosum* pyruvate phosphate dikinase (PPDK) was prepared according to the published procedure [5]. *Bacteroides fragilis* phosphono-pyruvate (Ppyr) decarboxylase was prepared as described [6]. Recombinant *Mytilus edulis* phospho-enolpyruvate (PEP) mutase, prepared according to published procedure, was kindly provided by Mr. Zhi-bing Lu in this laboratory [7]. Yeast inorganic pyrophosphatase was purchased from Sigma. Recombinant wild-type and mutant phosphonatases were prepared according to [8]. Thiamine pyrophosphate, unlabeled pyruvate and ATP, and all buffer components were the highest grade purchased from Sigma–Aldrich and were used without further purification.

**Synthesis of [1-14C]Pald**

A 1-ml reaction solution contained 50 mM K⁺-Hepes (pH 7.0), 20 mM ATP, 5 mM [2-14C]pyruvic acid (SA = 23 mCi/mmol), 20 mM Na₃PO₄, 5 mM MgCl₂, 40 mM NH₄Cl, 4.5 μM PPDK, and 5 units of inorganic pyrophosphatase (excess ATP and pyrophosphate hydrolysis, catalyzed by inorganic pyrophosphatase, were used to drive the reaction to completion). The reaction was allowed to stand at 25°C for 1 h and then applied to a 1.5 x 40-cm DEAE Sepharose column equilibrated with 0.1 M triethylammonium bicarbonate (pH 7.2). The column was eluted at 25°C with a 1.6-L linear gradient (0.1 to 0.4 M) of triethylammonium bicarbonate (pH 7.2) at a flow rate of 200 ml/h. The column fractions (20 ml/fraction) were assayed for radioactivity by dissolving an aliquot in Ready Safe liquid scintillation cocktail (Beckman Coulter) and carrying out liquid scintillation counting with a Beckman LS 6500 multipurpose Scintillation Counter. The PEP-containing fractions (eluted at ~0.28–0.30 M triethylamine bicarbonate) were combined and concentrated in vacuo to 1 ml by rotatory evaporation at 25°C. The sample was twice diluted with 10 ml of deionized water and concentrated to remove triethylamine bicarbonate. The resulting clear oil (~0.5 ml; 50% yield) was diluted to 1 ml with deionized water and stored at ~20°C. The purity of the [2-14C]PEP sample was confirmed by HPLC analysis; retention time 8.35 min (same as PEP standard) (Beckman Coulter System Gold instrument equipped with Phenomenex phenosphere SAX 00 G-3151-E0 ion-exchange column; mobile phase 0.4 M KCl, 0.1 M KH₂PO₄, pH 5.0; 0.7 ml/min flow rate).

[1-14C]Pald was synthesized immediately before use according to the following procedure. A 0.5-ml reaction solution containing 50 mM K⁺-Hepes (pH 7.5), 200 μM [2-14C]PEP (SA = 23 mCi/mmol), 10 mM MgCl₂, 4 mM thiamine diphosphate, 5 μM recombinant *M. edulis* PEP mutase, and 5 μM Ppyr decarboxylase was incubated at 25°C for 1 h. The enzymes were removed by centrifugation of the solution through a 5-kDa cutoff filter (twice) and the [1-14C]Pald was used without further purification. The [1-14C]Pald sample was confirmed by HPLC analysis; retention time 6.23 min (same as chemical standard) (Beckman Coulter System Gold instrument equipped with Phenomenex phenosphere SAX 00 G-3151-E0 ion-exchange column; mobile phase 0.4 M KCl, 0.1 M KH₂PO₄, pH 5.0; 0.7 ml/min flow rate).

The yield (>95%) of prepared Pald (0.2 mM) was determined using unlabeled PEP treated under the same conditions employed with the radiolabeled PEP. The concentration of Pald in the product sample was measured with a spectrophotometric assay based on the phosphonatase reaction. Accordingly, a 0.5-ml aliquot from the product solution was added to a 1-ml assay...
solution containing 0.01 μM phosphonatase, 5 units of alcohol dehydrogenase, 200 μM β-NADH, 10 mM MgCl2, and 50 mM K+ Hepes (pH 7.5, 25 °C) in a 1-cm-path-length cell. The ratio of the ΔAbs (340 nm) = 0.6 ODU and the molar extinction coefficient = 6200 M⁻¹ cm⁻¹ were used to calculate the Pald concentration.

**Synthesis of [³²P]Pald**

A 1-ml reaction solution contained 50 mM K⁺ Hepes (pH 7.0), 1 mM [β-³²P]ATP (SA = 7000 mCi/mmol), 20 mM pyruvic acid, 20 mM Na₃PO₄, 5 mM MgCl₂, 40 mM NH₄Cl, 4.5 μM PPK, and 5 units of inorganic pyrophosphatase. Excess pyruvic acid and inorganic pyrophosphatase were used to drive the reaction to completion. The [³²P]PEP was purified and concentrated using the same procedure described for [2-¹⁴C]PEP. The resulting clear oil (~0.2 ml) was diluted to 0.5 ml with deionized water and stored at −20°C. The purity of the stock was checked by HPLC (the same conditions as described above).

[³²P]Pald was synthesized just before use employing the following procedure. A 0.5-ml reaction solution containing 50 mM K⁺ Hepes (pH 7.5), 200 μM [³²P]PEP (SA = 5000 mCi/mmol), 10 mM MgCl₂, 4 mM thiamine diphosphate, 5 μM M. edulis PEP mutase, and 5 μM Ppyr decarboxylase was incubated at 25 °C for 1 h. The enzymes were removed from the solution and the identity and concentration of the Pald product (0.2 mM; ~100% yield) determined as previously described for the [1-¹⁴C]Pald.

**Results and discussion**

The goal of this work was to develop a method to synthesize [1-¹⁴C]Pald and [³²P]Pald. The reaction sequence used to this end is outlined in Fig. 2. In the first step of this sequence, the enzyme pyruvate phosphate dikinase catalyzes the interconversion of ATP, Pi, and pyruvate with AMP, PPi, and PEP. The reaction is driven to completion (in the PEP-forming direction) by inorganic pyrophosphatase-catalyzed hydrolysis of PPi. For the synthesis of [2-¹⁴C]PEP, commercial [2-¹⁴C]pyruvate is reacted with excess ATP and Pi. For the synthesis of [³²P]PEP, commercial [β-³²P]ATP is reacted with excess pyruvate and Pi. The (>99%) pure radiolabeled PEP is obtained in 50% yield by anion-exchange column chromatography of the reaction mixture. The specific activity of the [2-¹⁴C]PEP prepared in this manner was 23 mCi/mmol while that of the [³²P]PEP was 5000 mCi/mmol (calculated from the SA of the [β-³²P]ATP and the known decay rate).

Fig. 2. Enzymatic synthetic pathway of the radiolabeled Pald. PPDK, pyruvate phosphate dikinase; PEP mutase, phosphoenolpyruvate/phosphono-pyruvate phosphomutase; Ppyr, phosphono-pyruvate.
Based on our own experience, radiolabeled PEP is chemically stable when stored in deionized water at −20°C, while radiolabeled Pald is not. Thus, when desired for use, the radiolabeled Pald is generated from the radiolabeled PEP precursor in >95% yield and with the same specific activity as that of the starting PEP. As shown in Fig. 2, this is accomplished via the actions of two enzymes: PEP mutase and Ppyr decarboxylase. PEP mutase catalyzes the intramolecular rearrangement of PEP to Ppyr. This reaction is thermodynamically unfavorable ($K_{eq} \sim 1 \times 10^{-2}$) [9]. The ensuing decarboxylation step catalyzed by Ppyr decarboxylase is required to drive the Ppyr-forming reaction forward. Thus, acting together, the mutase and decarboxylase convert PEP to Pald. The enzymes are removed from the sample by size filtration. If needed, the Mg(II) can be removed with Chelex resin. Remaining is the thiamine pyrophosphate, which can be employed in the reaction at a much reduced concentration (10 μM). Likewise, the Hepes buffer can be reduced in concentration or a different buffer substituted. Attempts to purify the Pald using anion-exchange column chromatography (such as that used in the purification of the radiolabeled PEP) resulted in degradation of the Pald during the dilution–evaporation cycles.

The utilization of the synthetic, radiolabeled Pald as a probe of enzyme mechanism was demonstrated with the enzyme phosphonatase. As illustrated in Fig. 3, phosphonatase catalysis of Pald hydrolysis to acetaldehyde and orthophosphate proceeds via several covalent enzyme intermediates. Of these covalent intermediate species, only the Lys53 N-ethylenamine(III) has been trapped by borohydride reduction during steady state turnover [10].

In this experiment [1-14C]Pald and [32P]Pald were used in a single turnover reaction (i.e., concentration of phosphonatase ≫ concentration of radiolabeled Pald), terminated at varying conversion levels by rapid acid-quench, to identify one or more of the covalent adducts represented in Fig. 3. Following the quench, the enzyme was precipitated from the reaction and analyzed for radioactivity. The enzyme fractions isolated from control reactions (viz. reaction between the inactive phosphonatase mutant Lys53Arg and the radiolabeled Pald, or acid-quench added to wild-type phosphonatase before reaction with radiolabeled Pald) contained only ~2% of the total counts measured for the 14C-labeled enzyme (~60 to 80 compared to ~2800 to 3000 cpm) and only ~6% (~300 to 400 cpm compared to ~4500 to 5000 cpm) of the total counts measured for the [32P]-labeled enzyme.

Fig. 4 shows the time courses measured for formation and consumption of 14C-labeled or 32P-labeled enzyme in the single turnover reaction. In both cases, the radiolabeled enzyme reached a maximum level of ~15% of the limiting substrate at 50–60 ms and then decreased to near zero within 400 ms. Phosphonatase has a $k_{cat}$ of 15 s$^{-1}$; hence, the single turnover reaction should be completed.
in \( \sim 325 \text{ ms} \) (estimated by using the equation \( t_{1/2} = \ln 2/k_{\text{cat}} \) and assuming a greater than 99% conversion at seven half-lives). Therefore, the time courses for the reactions of the \([1-^{14}\text{C}]\text{Pald}\) and \([^{32}\text{P}]\text{Pald}\) demonstrate the formation and consumption of the kinetically competent covalent intermediates. Future experiments, to be carried out with double-labeled Pald and with different chemical quenches, will determine the time courses for the individual intermediates.

This study demonstrates that the \([1-^{14}\text{C}]\text{Pald}\) and \([^{32}\text{P}]\text{Pald}\) are useful probes in the study of the catalytic mechanism of phosphonatase. We anticipate their application in the investigation of Pald-dependent biochemical processes. Investigators may obtain the enzymes used in the synthesis of the radiolabeled Pald by writing to the corresponding author at the address provided.

References