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Separation of a L-aromatic amino acid decarboxylase from proline-rich protein (P5) by chromatofocusing and affinity chromatography. Proline-rich protein (P5), a major extracellular matrix protein found in culture media of mouse fibroblasts, has been shown to contain several different polypeptides by SDS-polyacrylamide gel electrophoresis. In this paper, we report on the separation of a L-aromatic amino acid decarboxylase from mouse skin, as well as the partial purification of the enzyme which is associated with P5 using ion-exchange and affinity chromatographies. The decarboxylase was selectively purified 18,500-fold using chromatofocusing on Mono S and affinity chromatography on poly-L-proline-Sepharose. Purified decarboxylase (also termed P5C decarboxylase) had a native molecular weight of 67,000 and a subunit size of 29,000. The enzyme catalyzed the decarboxylation of L-proline, L-tyrosine and L-phenylalanine with the highest activity toward L-proline. The decarboxylase had a pH optimum of 6.6 and was stable over a broad pH range.

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