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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. [Mesh] type = GeneratedMesh dim = 2 nx = 10 ny = 10 [Variables] [./u] [./] [Kernels] [./diff] type = Coefficient variable = u coefficient = 0.1 [./] [Materials] [./elasticity] type = ComputeSubsystemElasticity pos = '0 0 0' scale_along_x = 1 scale_along_y = 1 [./] [Executioner] type = Transient dt = 0.1 solve_type = 'PJFNK' petsc_options_iname = '-pc_type -pc_hypre_type -pc_hypre_boomeramg_max_3da54e8ca3

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