CS Bio Company

C S Bio was founded in 1993. Our initial product line was a complete line of automated synthesizers for manufacturing and process development, with a focus on cGMP production. Our instruments are the only synthesizers on the market that are fully validated for GMP production by an independent auditor (Weinberg, Spelton and Sax). In 1994, we opened our custom peptide facility. Initially opened to support R&D for our instrument line, we soon began producing high quality compounds for research organizations and pharmaceutical companies worldwide. Our current customers include Abbott Labs, Amgen, Merck, Pfizer, Roche, Sanofi-Aventis and many more. While we received our initial Drug Manufacturing License for cGMP production in 2001 for our San Carlos, CA facility, in 2004 we opened our state of the art peptide production facility in Menlo Park, CA with a primary focus on cGMP peptide production. Our goal is to provide our customers with the highest quality compounds available, in a time frame that matches our customer's requirements. Our cGMP peptide production facility has been issued a Drug Manufacturing License by the State of California Department of Health Services, Food and Drug Branch. This license is only issued after a thorough audit by the Food and Drug Branch, and the license must be renewed on a yearly basis. We have also undergone an FDA inspection from which we received a positive report and only several observations. The FDA report as well as a copy of our Drug Manufacturing License is available on request. Our approach to GMP manufacturing includes the dedication of one of our cGMP production lines and a team of chemists to each individual product. We have found this eliminates any chance of cross contamination, it keeps the chemists focused on the specific project at hand and this also goes a long way in regards to easing concerns of various regulatory agencies. We also supply our customers with all GMP related documentation including Batch Production Records, raw material C of A's and final QC data. It is imperative for us to work with our customers as closely as possible. We have undergone audits by many of our customers and we would certainly welcome any members of your staff to visit our facility for an audit/inspection. Should you wish to receive more information regarding our cGMP production capabilities such as overviews of recent projects, quality assurance information or contact information for references, we would be more than happy to send you the information ASAP. Should you have any questions regarding our cGMP production capabilities or our operations in general, please do not hesitate to contact us.

C S Bio Co.
People Who Know Peptides

C S Bio Company, Inc.

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