

Experimental Procedures Solid Phase Peptide Synthesis Spps

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Peptide SynthesisRunning a solid phase organic synthesis
Experimental Procedures Solid Phase Peptide
Experimental procedures Solid phase peptide synthesis (SPPS) Solid phase peptide synthesis (SPPS) was performed using a microwave-assisted peptide synthesizer (CEM) or in a standard manual reaction vessel under argon. Rink-amide MBHA resin and Wang resin were purchased from Sigma-Aldrich. DMF, DMSO, NMP, DCM, MeOH, ACN and DIEA were dried and distilled using standard protocols.

Experimental procedures Solid phase peptide synthesis (SPPS)
An effective synthetic procedure where the DOTA-linked peptide amide is synthesized on a single solid-phase support offers several advantages: the ease of coupling and time to synthesize one DOTA-peptide amide is dramatically reduced because DOTA is introduced to the peptide backbone [3, 5, 9, 10] while the peptide still attached to resin support, resulting in the formation of relatively pure DOTA-peptide product. Also, a large molar excess of DOTA will not be required as in solution-phase ...

A convenient and efficient total solid-phase synthesis of ...
General Procedures for the Fmoc Solid-Phase Peptide Synthesis. Peptides were synthesized with Fmoc solid-phase peptide synthesis (SPPS) on rink amide resin (initial loading = 0.15 mmol) manually. Fmoc deprotection was realized using 20% piperidine in DMF for 10 min at room temperature.

Frontiers | Design, Synthesis, and Antitumor Activities ...
Then, a solid-phase extraction (SPE) vacuum manifold (Figure S3d) (Phenomenex, Torrance, USA) was used for filtering and washing procedures of the resin by means of a PTFE universal stopcock (Figure S3a) that allowed the solvent passing through the resin and collected in a waste container.

Boosting Fmoc Solid-Phase Peptide Synthesis by Ultrasonication
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The solid-phase peptide synthesis starts with a resin which is insoluble under the conditions of the synthesis, usually a copolymer of polystyrene with 1 % divinylben-zene sometimes grafted with polyethylene glycol (Zalipsky et al. 1994), and which must also have an anchor for the synthesis to be carried out in the solid support (Merrield

Some Mechanistic Aspects on Fmoc Solid Phase Peptide Synthesis
Solid Phase Peptide Synthesis (SPPS) can be defined as a process in which a peptide anchored by its C-terminus to an insoluble polymer is assembled by the successive ad- dition of the protected amino acids consti- tuting its sequence.

Solid Phase Peptide Synthesis - Bachem
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Experimental Procedures Solid Phase Peptide Synthesis Spps
Diseases of immunity, including autoimmune diseases such as multiple sclerosis, transplantation graft rejection, allergy, and asthma, are prevalent and increasing in prevalence. They contribute to significant morbidity and mortality; however, few if any curative therapies exist, and those that are available lack either potency or specificity. Dendritic cells (DCs) are sentinels of the immune ...

Steroid|Peptide Immunoconjugates for Attenuating T Cell ...
experimental procedures solid phase peptide Experimental procedures Solid phase peptide synthesis (SPPS) Solid phase peptide synthesis (SPPS) was performed using a microwave-assisted peptide synthesizer (CEM) or in a standard manual reaction vessel under argon. Rink-amide MBHA resin and Wang resin were purchased from Sigma-Aldrich. DMF, DMSO ...

Experimental Procedures Solid Phase Peptide Synthesis Spps ...
Experimental Procedures Solid Phase Peptide Experimental procedures Solid phase peptide synthesis (SPPS) Solid phase peptide synthesis (SPPS) was performed using a microwave-assisted peptide synthesizer (CEM) or in a standard manual reaction vessel under argon. Rink-amide MBHA resin and Wang resin were purchased from Sigma-Aldrich.

Experimental Procedures Solid Phase Peptide Synthesis Spps
We demonstrate a solid-phase capture strategy suitable for the proteolysis, purification, and subsequent chemical modification of peptides. We use this resin on an HEK293T cell lysate and perform one-pot proteolysis, capture, and derivatization to generate a cellular proteome that identified over 40,000 bead-bound peptides.

Solid-Phase Peptide Capture and Release for Bulk and ...
Experimental Procedure. The procedure is composed of six steps: 1 1. A few milligrams of resin is placed in a small glass tube and washed with DMF. 2. DMF (3 ml) is added followed by NMM (1 ml) and TCT (5 mg). 3. The tube is heated at 70° for 20 min. 4. The solution is pipetted off and the resin is washed thoroughly with DMF. 5.

Qualitative Colorimetric Tests for Solid Phase Synthesis ...
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Experimental Procedures Solid Phase Peptide Synthesis Spps ...
Solid-state 15 N NMR experiments were performed on samples of the M6(252)269, C252A) peptide fragment of the Ste2p receptor in lipid bilayers. The peptides contain a single specific 15 N-labeled amino acid at the position Leu4, Leu5, Val6, Ile9, Ile10, Phe11, Ile12, Leu13, Ala14, or Leu17.

Structure and Topology of a Peptide Segment of the 6th ...
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Abstract We report a general and operationally simple method for the solid phase synthesis of α -ketoamide peptides using standard Fmoc solid phase peptide synthesis. The method delivers deprotected peptide α -ketoamides directly upon resin cleavage without any additional steps, and tolerates all side chain functional groups.

Facile Synthesis of Internal and C-Terminal Peptide α ...
Typically, the peptides are cleaved from the support together with the side-chain protecting groups, but there are several experimental procedures, which require solid-phase-bound peptides. The SPOT technique uses membranes as the solid support and enables the parallel synthesis and testing of hundreds to thousands of peptides at different locations on one membrane (Frank 1992).

Chemistry of Fmoc Peptide Synthesis on Membranes ...
The most commonly N-terminal protecting group used in Solid Phase Peptide Synthesis (SPPS) is the Fmoc group (9-fluorenylmethoxycarbony-) (Scheme 1, Table 1) [1] [3]. Besides the coupling procedure, the Fmoc deprotection step is another most crucial stage in peptide synthesis. Above all, the property which makes it a valuable tool is its stability in acidic conditions.

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