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In chemistry, stereoselectivity is the property of a chemical reaction in which a single reactant forms an unequal mixture of stereoisomers during the non-stereospecific creation of a new stereocenter or during the non-stereospecific transformation of a pre-existing one.

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A stereoselective synthesis of (+)-herboxidiene is described. The convergent synthesis utilized a Suzuki cross-coupling reaction to assemble the key segments. The synthesis of the functionalized tetrahydropyran ring utilized an Achmatowicz reaction as the key step. The synthesis of the C10-C19 segment was accomplished using Brown's

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The synthesis of optically active secondary and tertiary propargylic alcohols was accomplished by addition of lithium acetylide to chiral α -sulfinyl enones.

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A highly stereoselective method for the synthesis of functionalized 3-oxabicyclo[3.3.1]nonan-2-one derivatives with four contiguous stereogenic centers, including one tetrasubstituted stereogenic

Topics in current chemistry, 216, Stereoselective ...

Stereoselective Heterocyclic Synthesis III Volume Editor: Peter Metz With contributions by A. Hassner, I. N. N. Namboothiri, U. Nubbemeyer, S. D. Rychnovsky, C. J. Sinz, P. ten Holte, B. Zwanenburg Springer The series Topics in Current Chemistry presents critical reviews of the present and future trends in modern chemical research. The scope of coverage includes all areas of chemical science.

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A stereoselective synthesis of a bioactive triarylmethane is described. Key to the synthesis is a nickel-catalyzed Suzuki-Miyaura coupling which proceeds with retention at the benzylic center.

Synthesis of tetrahydropyrans from sugar lactones - [PDF ...

Pergamon Tetrahedron 54 (1998) 13591-13620

TETRAHEDRON Synthesis of tetrahydropyrans from sugar lactones Juan C. Estevez a, Antony J. Fairbanks b^{*} and George W.J. Fleet b^{*}

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