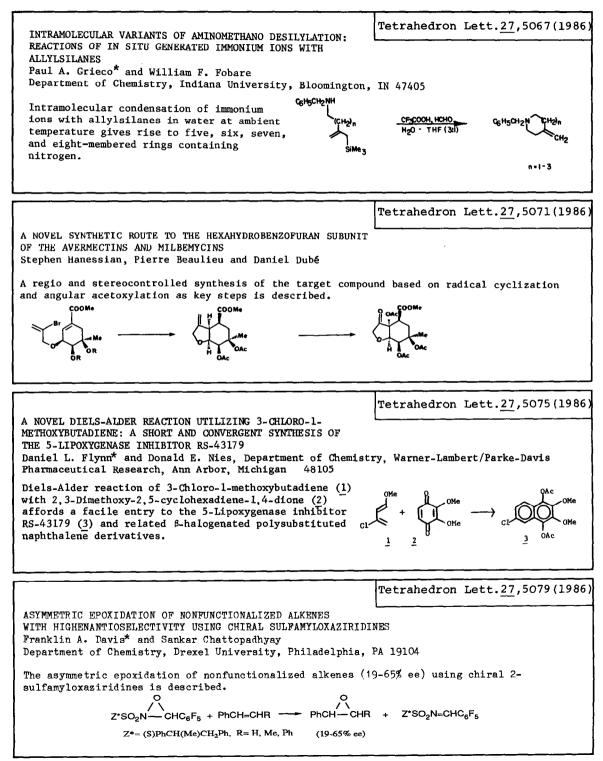
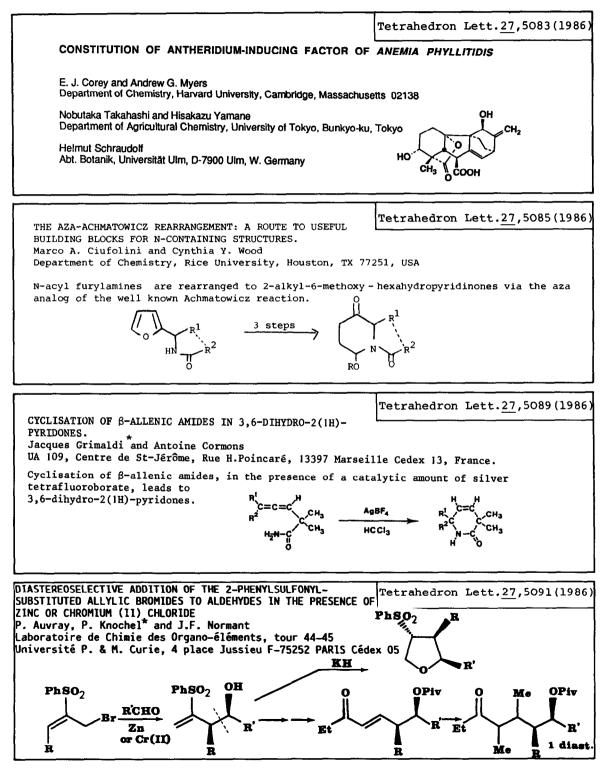
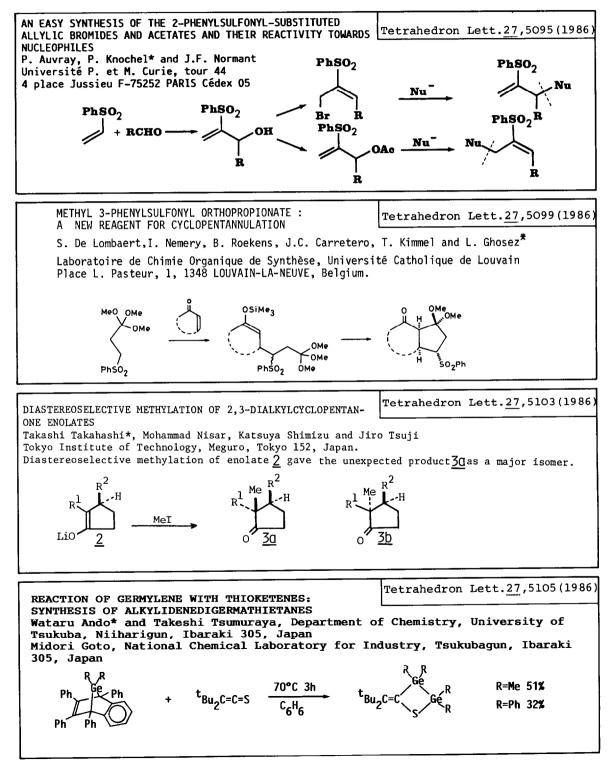
GRAPHICAL ABSTRACTS

Tetrahedron Lett.27,5051(1986) ISOTOPE EFFECTS AND MECHANISM IN THE BROMINATION OF ALPHA- AND BETA-CARBON-14 LABELED 4-NITRO-4'-METHYLSTILBENES Pandurang B. Kokil and Arthur Fry* Department of Chemistry and Biochemistry, University of Arkansas, Fayetteville, AR 72701, USA Carbon-14 istotpe effects are reported, for the first time, in a simple bromination reaction, $k/^{\alpha}k = 1.013 \pm 0.003$; $k/^{B}k = 1.0035 \pm 0.003$. Br₂ in CH₂Cl₂ 8_ NO 26° C Br 8r Tetrahedron Lett.27,5055(1986) ISOTOPE EFFECTS AND MECHANISM IN THE BASE-PROMOTED DEHYDROCHLORINATION OF 1,1-DICHLORO-2,2-DIARYLETHANES-1- $^{14}\mathrm{C}$ Agnes Pulay and Arthur Fry* Department of Chemistry and Biochemistry, University of Arkansas, Fayetteville, AR 72701, USA Carbon-14 isotope effects, k/ lpha k, are 1.045 and 1.024 for dehydrochlorinations of Z = H and $Z = NO_2$ DDD-type compounds. Both compounds react by the E2 mechanism. →)₂ch-^achci₂ → base (z-))-),с=^аснс≀ Tetrahedron Lett.27,5059(1986) CARBON-14 KINETIC ISOTOPE EFFECTS AND MECHANISM IN THE SOLVOLYSIS OF 1,1,1-TRIFLUORO-2-PHENYL-2-PROPYL-3-¹⁴C p-TOLUENESULFONATE Zili Guo and Arthur Fry Department of Chemistry and Biochemistry, University of Arkansas, Fayetteville, AR 72701, USA Solvolysis of 1,1,1-trifouoro-2-phenyl-2-propyl-3- 14 C p-toluenesulfonate takes place with only a very small $^{\beta}$ C isotope effect, k/^Bk = 1.008 ± 0.002. The mechanism is S_N1/E1. SH TsOH + Ph- $\alpha C - \beta CH_3$ + Ph- $\alpha C = \beta CH_2$ heat ĊFą CF3 ĊFa Tetrahedron Lett.27,5063(1986) CARBON-14 KINETIC ISOTOPE EFFECTS AND MECHANISMS OF ADDITION OF 2,4-DINITROBENZENESULFENYL CHLORIDE TO SUBSTITUTED STYRENES IN THE PRESENCE OF LITHIUM PERCHLORATE Marianna Kanska and Arthur Fry* Department of Chemistry and Biochemistry, University of Arkansas, Fayetteville, AR 72701, USA Different ${}^{\alpha}$ C and ${}^{\beta}$ C isotope effect behavior in the absence and presence of LiClO₄ demonstrate different mechanisms for Z = Me (open carbenium ion) than for Z = H or Cl (cyclic intermediate ion pair, less associated with added salt). LICIO $-^{\alpha}CH-^{\beta}CH_{2}-Ar + Z-O-^{\alpha}CH^{\beta}CH_{2}-Ar$ -^αCH=^βCH₂ ArSCI . Å

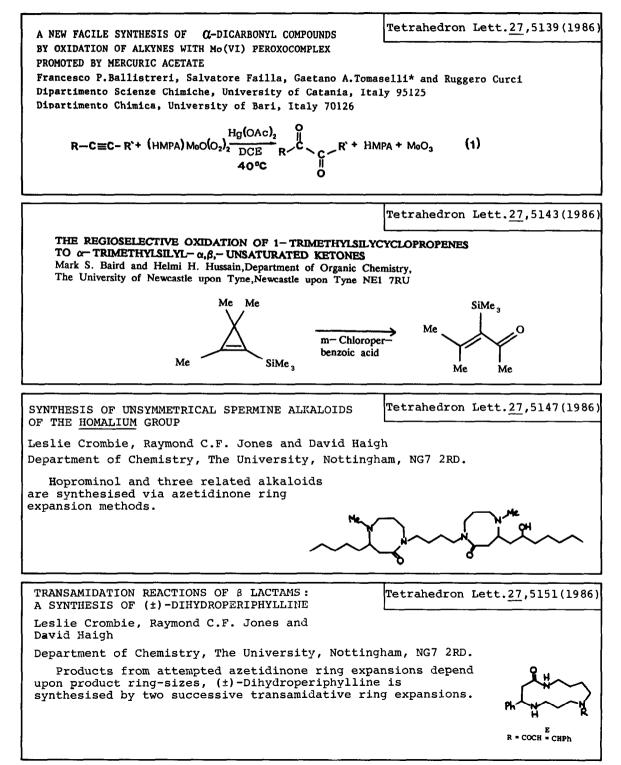






$$\begin{array}{c} \hline \\ \text{RMMESIS OF OPTICALLY ACTIVE 4-HYDROWALK-2-ENENTFILIES:}\\ \hline \\ \text{Retaining the optically active 2-o-chargements supervised by the set of the s$$

Tetrahedron Lett.27,5125(1986) A NOVEL DEGRADATIVE STRATEGY FOR THE SYNTHESIS OF p-QUINONES José M. Saá, Jerónimo Morey and Antonio Costa Departamento de Química. Universitat de les Illes Balears. E-07071 Palma de Mallorca. Spain. E: D,Me,-Fremy's salt promoted oxidative degradation of p-hydroxy benzylalcohols Tetrahedron Lett. 27, 5129 (1986) REGIOCONTROLLED ADDITION IN THE REACTION OF N-(α -METHOXYPHENACYL)ANILINES WITH METHYL LITHIOISOBUTYRATE B. Alcaide^{*}, C. López-Mardomingo, R. Pérez-Ossorio, J. Plumet, and J. Rodriguez-López Dpto. Química Orgánica, Facultad de Química, Universidad Complutense,28040-Madrid, Spain Reaction of the title compounds gives compounds 2-6 through a regiocontrolled process. Ar N Ph OH O 3 Ph H OME Tetrahedron Lett.27,5133(1986) A NOVEL CYCLOPENTANE ANNULATION REACTION : NEW SYNTHESIS OF ESTRONE Nurani S. Narasimhan* and Prakash A. Patil Garware Research Centre, Department of Chemistry, University of Poona, Pune - 411 007, INDIA. Tetrahedron Lett.27,5135(1986) SYNTHESIS WITH ORGANOBORANES. 2. SYNTHESIS OF ~- AND /-DAMASCONE Marek Zaidlewicz Institute of Chemistry, Nicolas Copernicus University, 87-100 Torun, Poland Syntheses of ∞ -damascone /1/ and f-damascone /2/ via allylic organoboranes. ++



Tetrahedron Lett.27,5155(1986) Heterocyclic Silyl Enol Ether Chemistry : Synthesis and Reactivity of 3-Trimethylsiloxyfuran and 3-Trimethylsiloxythiophene. Lucia Camici, Alfredo Rícci and Maurizio Taddei Centro CNR Composti Eterociclicici, Dipartimento Chimica Organica, Firenze, Italy. Treatment of 3-Lithiofuran and-thiophene with $(Me_3Si)_2 o_2$ gave the corresponding trimethyl-siloxy-heterocycles which were further functionalised. $\int_{0}^{0 \text{SiMe}_{3}} R' \xrightarrow{\text{R}} \frac{X=0}{2}$ (Me₃Si)₂O₂ Tetrahedron Lett.27,5159(1986) BENZOCALICENES AND BENZOTRIAHEPTAFULVALENES FROM CYCLOPROPARENES B. Halton, S.J. Buckland, Q. Mei, and P.J. Stang, Chemistry Departments, Victoria University of Wallington, Private Bag, Wellington, New Zealand, and The University of Utah, Salt Lake City, Utah 84112, U.S.A. Peterson olefination of cyclopropabenzene and cyclopropa[b]naphthalene gives the fulvalenes (8)/(10)/(12) and (9)/(11)/(13) respectively. The dipole moments of 8 R = H (8) and (12) have been measured. 10 0 - 4 9 BB = benze 11 RR = F