

be approximately 0° if the Mo atom were bonded solely to the nitrogen lone pair or 90° if it were bonded to the imine π bond.

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Novel Chiral Triorganotin Halides. Stabilization of Optically Active Tin Centers by Intramolecular Coordination

Sir:

Contrary to the high optical stability of tetraorganotin compounds¹ triorganotin halides are configurationally unstable.^{2,3} In the presence of nucleophiles the $RR'R''SnX$ enantiomers isomerize either via formation of penta- or hexacoordinate tin intermediates which are stereochemically nonrigid or via an SN_2 type mechanism which leads to direct inversion of configuration.⁴

We report here that chiral triorganotin halides with high optical stability may be obtained by rendering the tin atom pentacoordinate by intramolecular coordination so that stereoisomerization pathways are blocked.

Recently, we have reported⁵ a facile one-step synthesis for $(2-Me_2NCH_2C_6H_4)_2R_2SnBr$ compounds (**1**, $R = Me$; **2**, $R = Ph$). An x-ray investigation has shown that the tin atom in **2** is pentacoordinate as a result of intramolecular Sn-N coordination.⁶

We have now synthesized⁷ $(2-Me_2NCH_2C_6H_4)MePhSnBr$, **3**, in which the tin atom is a chiral center. The Me_2NCH_2 ligand, present in this structure, allows the study by NMR spectroscopy of the configurational stability of the chiral tin center as well as of the occurrence of intramolecular Sn-N coordination.⁸

Figure 1 shows the NMR spectrum of **3** in toluene- d_8 at 10, 30, and 123 °C. The benzylic protons (25 °C;⁹ AB, δ 3.02 and 2.81 ppm) are anisochronous up to 123 °C¹⁰ ($\Delta\delta$ decreases from 21 Hz (0 °C) to 15.5 Hz (123 °C)) and thus diastereotopic. Consequently, up to 123 °C the rate at which the absolute configuration of the chiral tin atom inverts ($Sn \rightleftharpoons n\bar{2}$) is slow on the NMR timescale, because it is only by this inversion process that the benzylic protons become enantiotopic and isochronous.

The observation below 30 °C of two singlets for the NMe protons at δ 1.50 and 1.28 ppm which coalesce above 30 ° to one singlet at 1.39 ppm is interpreted in terms of rate determining intramolecular Sn-N coordination. Pyramidal inversion at nitrogen in N,N -dialkylbenzylamines is a low energy process which has an estimated barrier lower than 6 kcal/mol.¹¹ Consequently, inversion at nitrogen (rate constant k_i) takes place in the tetracoordinate tin conformer. In the pentacoordinate tin conformer Sn-N coordination (rate constant k_c) makes a stable prochiral assembly of the NMe_2 group and renders the NMe groups diastereotopic and thus anisochronous because of the dissymmetry at tin. Two processes can be envisaged by which the NMe groups can become homotopic (enantiotopic): (i) by inversion of configuration at tin in the pentacoordinate conformer (rate constant k_r'), i.e., without prior Sn-N bond dissociation (vide infra), and (ii) by Sn-N bond dissociation (rate constant k_d) followed by inversion at N with concomitant 180 ° rotation about the CH_2-N bond and

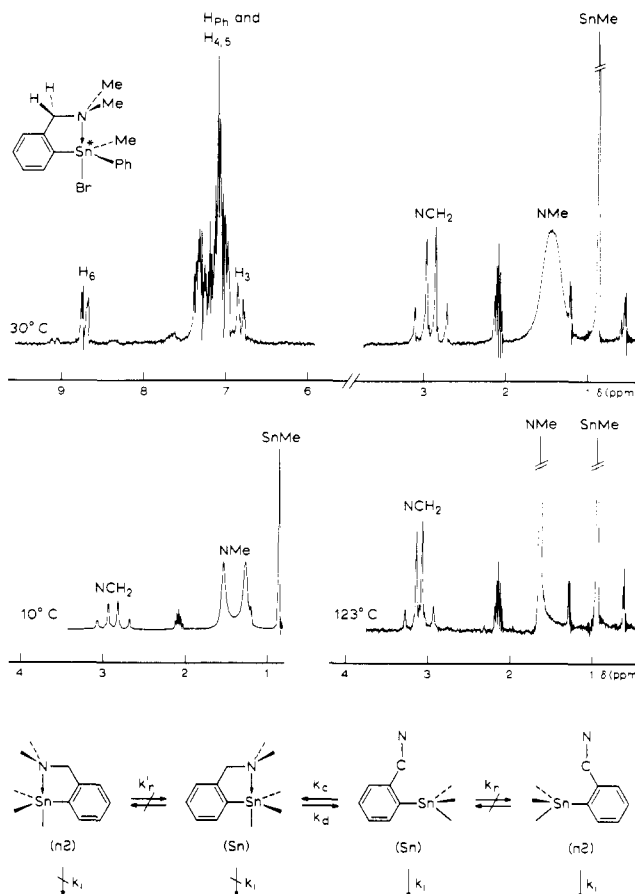


Figure 1. 1H NMR spectrum (100 MHz) of $(2-Me_2NCH_2C_6H_4)(C_6H_5)MeSnBr$ in toluene- d_8 (resonances at δ 2.1 and 7.1 ppm are due to hydrogen containing sites of the solvent). Only the high field part of the spectra recorded at 10° and 123 °C is shown.

re-formation of the Sn-N bond. The latter mechanism accounts for the observed dynamic NMR pattern whereas the observation that the NCH_2 and NMe proton resonance patterns coalesce at different rates excludes the former mechanism.

External ligands such as PPh_3 , NPh_3 , DABCO, and pyridine, which readily cause inversion of configuration in other chiral triorganotin halides,^{1,2} do not interact with the tin atom in **3**¹² as can be concluded from the absence of any effect on the resonance patterns or on the T_c value of the NMe resonances. However, it has been shown that in $(2-Me_2NCH_2C_6H_4)_2MeSnBr$, **4**,¹³ which contains two internal ligands, intramolecular exchange does take place. The NMR of **4** at -50 °C shows two sets of diastereotopic NCH_2 protons (two AB patterns: $\Delta\delta$, 1.48 ppm, irradiation at 2.7 ppm resulted in a singlet at 4.27 ppm; $\Delta\delta_2$ 0.58 ppm), two multiplets for H_6 at δ 8.85 and 7.20 ppm, and two sets of NMe protons (two singlets at δ 1.34 and 1.52 ppm for diastereotopic NMe groups and one singlet at 1.62 ppm for homotopic NMe groups) (Figure 2). This pattern is compatible with a pentacoordinate structure at low temperature containing one C,N- and one C-bonded $2-Me_2NCH_2C_6H_4$ group. In the temperature range -50 ° to 20 °C the two sets of resonance patterns for the $2-Me_2NCH_2C_6H_4$ groups coalesce to give an averaged multiplet for H_6 at δ 8.12 ppm, one broadened singlet for the NMe protons at the position of homotopic NMe groups while the two sets of diastereotopic NCH_2 protons coalesce to an averaged AB pattern with $\Delta\delta$ 0.58 ppm. The two types of $Me_2NCH_2C_6H_4$ groups observable below -50 °C have now become equivalent on the NMR timescale by an intramolecular exchange process (pentacoordinate $Sn \rightleftharpoons$ pentacoordinate

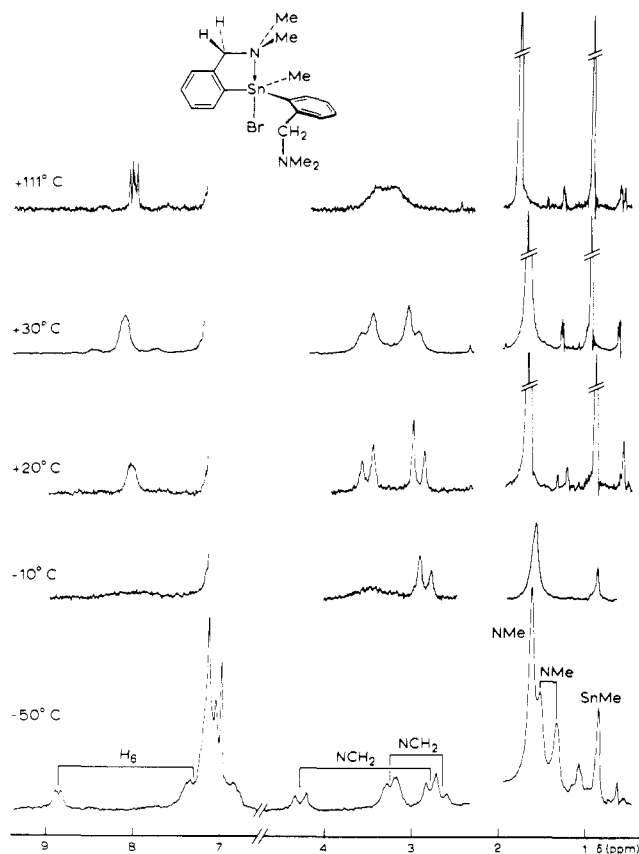
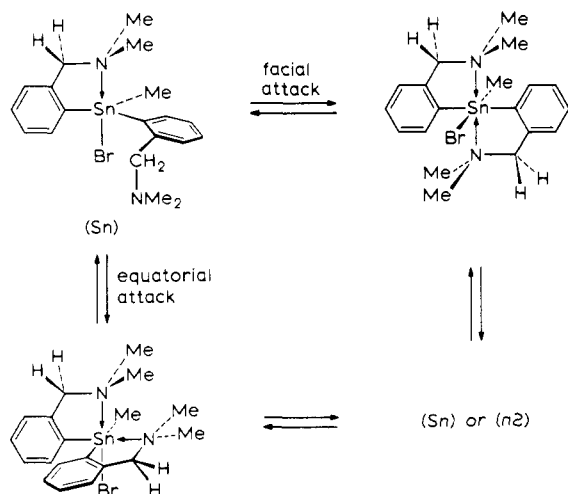


Figure 2. ^1H NMR spectrum (100 MHz) of $(2\text{-Me}_2\text{NCH}_2\text{C}_6\text{H}_4)_2\text{MeSnBr}$ in toluene- d_8 .

Scheme I



Sn') which presumably involves a hexacoordinate transition state or intermediate, e.g., Scheme I. Above 20°C the rate constants for both the dissociation process (pentacoordinate $\text{Sn} \rightleftharpoons$ tetracoordinate Sn) and the inversion process at nitrogen increase as is reflected in a sharpening of the H_6 multiplet as well as in a decrease of the $\Delta\delta$ value of the anisochronous CH_2 protons.¹⁴

We ascribe the high configurational stability of these triorganotin halides to a combination of factors. Complex formation with external Lewis bases, believed to be the first step in the stereoisomerization process of triorganotin halides, cannot compete with intramolecular coordination.¹⁵ Furthermore, stereoisomerization routes in the pentacoordinate conformer,¹⁶ e.g., by a Berry pseudorotation mechanism, are energetically unfavorable in view of (i) the strong site prefer-

ence of the electronegative bromine and nitrogen atoms for axial positions¹⁷ and (ii) the geometric constraints of the five-membered chelate ring (N-Sn-C , 75.3°).⁶ Study of the corresponding $(2\text{-Me}_2\text{NCH}_2\text{C}_6\text{H}_4\text{CH}_2)\text{MePhSnBr}$, **5**,¹⁸ for which intramolecular Sn-N coordination leads to a six-membered chelate ring¹⁹ supports this idea.

The NMR spectrum of **5** in toluene- d_8 shows at room temperature isochronous resonances for the $-\text{CH}_2\text{Sn}$ (δ 4.07 ppm, averaged $^2J_{\text{Sn-H}} = 86$ Hz), $-\text{CH}_2\text{N}$ (δ 3.91 ppm), and NMe (δ 1.41 ppm) protons. On cooling the NMe groups ($T_c \approx -37^\circ\text{C}$; $\Delta\delta$ 38 Hz) and CH_2Sn and CH_2N protons ($T_c \approx 0^\circ\text{C}$; $\Delta\delta$ 15 and 9 Hz, respectively) become diastereotopic and thus anisochronous at about the same rate. It is only in the pentacoordinate conformer that inversion of the absolute configuration at the tin atom is reflected simultaneously in the resonance patterns of these three prochiral groupings. Ligand site exchange is now a lower energy process because of the lower constraints of a six-membered chelate ring containing two sp^3 carbon atoms as compared with the five-membered chelate rings of **3** and **4**.

Acknowledgment. We thank Miss N. Natan and Mr. J. T. B. H. Jastrzebski for preparative assistance and Mr. J. W. Marsman for helpful discussions.

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- (7) Obtained in 90% yield from the 1/1 reaction of $\text{Me}_2\text{NCH}_2\text{C}_6\text{H}_4\text{-Cu}$, **6**, or $-\text{Li}$, **7**, with MePhSnBr_2 according to the procedures described in ref 5. Mp $121\text{--}125^\circ\text{C}$, monomeric in benzene (osmometry) concentration independent, 432; $\text{C}_{16}\text{H}_{20}\text{BrNSn}$ calcd, 425.
- (8) We have used the Me_2NCH_2 ligand as a probe for the detection of dissymmetry in the metal core of arylmetal 1b cluster compounds.
- (9) Analysis of the AB part of the ABX spectrum due to ^{117}Sn and ^{119}Sn isotopes which allows the determination of a $^4J_{\text{Sn-H}}$ is underway; see also Holloway et al.²
- (10) I. e., the racemization process has an activation energy barrier much higher than 100 kJ mol^{-1} which should allow separation of the enantiomers.
- (11) E.g., H. Kessler, *Angew. Chem.*, **82**, 237 (1970).
- (12) This conclusion is also supported by the observation that $^2J_{\text{Sn-CH}_3} = 68.2/65.5$ Hz and $^3J_{\text{Sn-H}_6} = 70$ Hz are solventdependent whereas the chemical shift values of various protons in **3** are very solvent dependent, i.e., the series toluene- d_6 /pyridine- d_5 / CCl_4 (δ ppm): NMe, 1.45/1.87/2.00; CH_2N , 3.02 and 2.81/3.39 and 3.38/3.50 and 3.37 ($J_{\text{gem}} = 14$ Hz); CH_2Sn , 0.87/1.16/1.00 and H_6 , 8.72/8.76/8.30. The latter data show that preferential solvent-solute interactions occur at the site of the intramolecular ligand CH_2NMe_2 .⁵
- (13) Obtained in 60% yield from the addition in 1/1 molar ratio of Me_2SnBr_2 to **7** as well as from the 1/1 reaction of $(2\text{-Me}_2\text{NCH}_2\text{C}_6\text{H}_4)_2\text{Me}_2\text{Sn}$ with Me_2SnBr_2 . Mp $142\text{--}144^\circ\text{C}$, monomeric in benzene (osmometry) found, 484; $\text{C}_{10}\text{H}_{27}\text{Br}_2\text{N}_2\text{Sn}$ calcd, 481.
- (14) The benzylic carbon atoms in the tetracoordinate tin conformer are not C_2 centers and thus the benzylic protons remain diastereotopic.
- (15) The occurrence of intramolecular Sn-N coordination in the tetracoordinate tin conformer of **3** can be viewed as the first step of an $\text{S}_{\text{N}}2$ type substitution process at tin. Continuation of this process beyond the TBP configuration resulting in Sn-Br bond dissociation, would require opening of the N-Sn-C angle from about 75.3° to formally 109° . This is unlikely in view of the geometric constraints of the chelate ring. However, in $[2,6\text{-}(\text{Me}_2\text{NCH}_2)\text{C}_6\text{H}_3]\text{MePhSnBr}$ the Sn-Br bond is dissociated because the C_3Sn cationic state is stabilized by Sn-N coordination of the two Me_2NCH_2 ligands resulting in a TBP configuration around tin.
- (16) E. L. Muettterties and R. A. Schunn, *Q. Rev., Chem. Soc.*, **20**, 245 (1966); G. M. Whitesides and H. L. Mitchell, *J. Am. Chem. Soc.*, **91**, 5384 (1969); F. H. Westheimer, *Acc. Chem. Res.*, **1**, 70 (1968); R. R. Holmes, *ibid.*, **5**, 296 (1972); J. R. Shapley and J. A. Osborn, *ibid.*, **6**, 305 (1973).
- (17) B. Y. K. Ho and J. J. Zuckerman, *J. Organomet. Chem.*, **49**, 1 (1973).
- (18) Isolated in 30% yield from 1/1 reaction of $2\text{-Me}_2\text{NCH}_2\text{C}_6\text{H}_4\text{CH}_2\text{Li}$ with MePhSnBr_2 , see ref 5. Mp $152\text{--}160^\circ$; monomeric in benzene (ebulliometry)

found, 439; C₁₇H₂₂BrNSn calcd, 438.

- (19) This six-membered ring is puckered both at the CH₂Sn as well as at the SnN(Me₂)CH₂ part. In the pentacoordinate {5-MeO-8-Me₂NCM₂naphthyl}MeRSnBr (R = i-Me or Ph) compounds puckering is restricted to the SnN(Me₂)CH₂ part which results in two different conformations for the six-membered chelate ring. Both conformations can be frozen out on the NMR timescale below -30 °C.

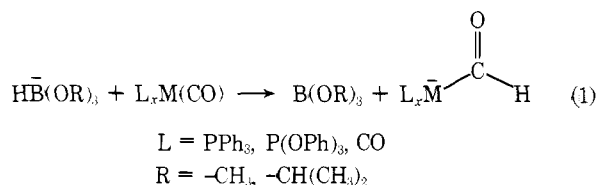
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Synthesis of Transition Metal Formyl Compounds by Nucleophilic Attack of Hydride on Coordinated CO

Sir:

There is renewed interest in the conversion of coal to gaseous and liquid fuels. These conversions can be accomplished by the reaction of coal with water to give a 1:1 mixture of CO:H₂, which can subsequently be passed over various heterogeneous transition metal catalysts to give methane or methanol or (via the Fischer-Tropsch reaction) gasoline.¹⁻³ Little is known about the mechanism of these reductions, but transition metal formyl complexes may well be important intermediates in the initiation step for the catalytic reduction of CO by H₂. However, attempts to obtain evidence for the intermediacy of metal formyl complexes in the reactions of metal carbonyl hydrides have uniformly met with failure.⁴ Our interest in the mechanism of the Fischer-Tropsch reaction and in the development of a homogeneous catalyst for the reduction of CO has led us to undertake a study of the synthesis and reactions of metal formyl complexes. The only previously reported metal formyl complex, [(Ph₃P)₂N]⁺(CO)₄FeCHO⁻, **1**, was prepared by Collman and Winter in 1973 by the reaction of Na₂Fe(CO)₄ with acetic formic anhydride.⁷ Here we report (1) a new synthesis of metal formyl complexes from the reactions of metal carbonyl compounds with trialkoxyborohydrides, (2) the isolation of a second stable metal formyl complex, [Et₄N]⁺*trans*-[(PhO)₃P](CO)₃FeCHO⁻, **2**, and (3) the observation of a number of unstable metal formyl complexes.

The reactions of borohydrides with transition metal carbonyl compounds have been used to facilitate the substitution of one or more ligands for CO,⁸⁻¹⁰ and to synthesize certain group 6 dinuclear metal carbonyl hydrides.^{11,12} We have found that the reaction of Na⁺HB(OCH₃)₃⁻ or K⁺HB(O-*i*-Pr)₃⁻ with various metal carbonyl compounds leads to transition metal formyl compounds according to eq 1. Reaction of Na⁺H-



B(OCH₃)₃⁻ with Fe(CO)₅, followed by cation exchange with [(Ph₃P)₂N]⁺Cl⁻, gives [(Ph₃P)₂N]⁺(CO)₄FeCHO⁻, **1**, in 80-90% yield by NMR analysis¹⁴ and in 39% isolated yield, mp 122-132 °C (sealed tube) dec.¹⁵ The infrared spectrum and ¹H and ¹³C NMR spectrum of **1** prepared from Fe(NO)₅ and Na⁺HB(OCH₃)₃⁻¹⁶ were identical with the spectra of a sample of **1** prepared from Na₂Fe(CO)₄ and acetic formic anhydride.⁷ The formyl proton characteristically appears at very low field, δ 14.95, in the ¹H NMR; similarly, the formyl carbon appears at characteristically low field, δ 270.5, in the ¹³C NMR spectrum. Reaction of **1** (prepared from Na⁺HB(OCH₃)₃⁻ and Fe(CO)₅) with HCl in THF gives formal-

dehyde as previously reported.^{7,17} A THF solution of **1** reacts with HCl in the presence of Ph₃P to give (CO)₄FePPh₃ (57%) and (CO)₃Fe(PPh₃)₂ (20%); similar results were reported in the NaBH₄ "facilitated" substitution of Fe(CO)₅ by PPh₃, although iron formyl species were not proposed for these reactions.⁸

The reaction of trialkoxyborohydrides as a hydride source with metal carbonyl compounds is an excellent organometallic synthetic method. The rapidity and ease of the procedure, along with the ability to obtain adequate yields, make the method quite useful. Moreover, this procedure can be extended to the synthesis of metal formyl compounds from systems other than Fe(CO)₅, whereas in our hands the reaction of acetic formic anhydride with metal carbonyl anions could not. Reaction of CH₃C(O)OC(O)H with C₅H₅Fe(CO)₂⁻, Cr(CO)₅²⁻, (Ph₃P)Mn(CO)₄⁻, and (Ph₃P)₂Mn(CO)₃⁻ produced no products with ¹H NMR resonances below δ 10 expected for metal formyl compounds. However, reaction of Na⁺HB(OCH₃)₃⁻ with the appropriate metal carbonyl compound in THF in a sealed NMR tube led, in many cases, to the appearance of characteristic formyl ¹H NMR resonances below δ 10. We now have NMR evidence for the formation of metal formyl compounds from Cr(CO)₆ (δ 15.2, s),¹⁸ W(CO)₆ (δ 15.9, s), (CO)₅CrPPh₃ (δ 15.3, s), (CO)₅WPPh₃ (δ 15.8, s), and (CO)₄FePPh₃ (δ 15.5, d, J = 24 Hz).¹⁹ Quantitative NMR studies on several of the above systems indicated that the extent of the reaction producing the desired metal formyl compound was generally quite low (<10%),¹⁴ and as a result we tried analogous reactions of K⁺HB(O-*i*-Pr)₃⁻ with several metal carbonyl compounds in an attempt to increase the extent of reaction by using a more reactive hydride.^{13b} We found that Cr(CO)₆ reacts with K⁺HB(O-*i*-Pr)₃⁻ to give a maximum 76% conversion to the formyl compound in 25 min at ambient temperature;¹⁴ the product formyl complex, K⁺(CO)₅CrCHO⁻, is half decomposed in an additional 40 min. In contrast, the reaction of Cr(CO)₆ with Na⁺HB(OCH₃)₃⁻ gives a maximum of 4% of the metal formyl after 25 min at ambient temperature.¹⁴ The reaction of K⁺HB(O-*i*-Pr)₃⁻ with W(CO)₆ and (CO)₄FePPh₃ gives qualitatively similar results.

We have been able to isolate and characterize the second known compound of the metal formyl class by the reaction of a sixfold excess of K⁺HB(O-*i*-Pr)₃⁻ with (CO)₄FeP(OPh)₃ in THF.¹⁵ Cation exchange of the resulting potassium salt with [Et₄N]⁺Br⁻ and 1 N NaOH gave [Et₄N]⁺*trans*-[(PhO)₃P](CO)₃FeCHO⁻, **2**, as a light tan powder in 64% yield, mp 40-55 °C (sealed tube) dec.²¹ Solutions of **2** are rapidly oxidized by air and undergo thermal decomposition at 65° over several hours. **2** shows spectral properties fully consistent with its proposed structure: ¹H NMR (acetone-*d*₆) δ 1.30 (tt, J = 7.7 Hz, J_{NH} = 1.5 Hz, 12 H, NCH₂CH₃), 3.40 (q, J = 7.7 Hz, 8 H, NCH₂CH₃), 6.8-7.8 (m, 15 H, C₆H₅), 14.82 (d, J_{PH} = 44.4 Hz, 1 H, FeCHO); ir (THF) 2515 cm⁻¹ (w, aldehyde C-H²²), 1960 (m, MC≡O), 1872 (vs, MC≡O²³), 1584 (m, formyl C=O).

This new synthesis of metal formyl complexes will enable us to study the kinetic stability of metal formyl complexes, the equilibrium between metal hydrides and metal formyl complexes, and the reaction of metal formyl compounds with hydrogen and other reducing agents.

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Supplementary Material Available: Experimental procedures (2 pages). Ordering information is given on any current masthead page.