JOC *Additions and Corrections*

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Sreekanth A. Ramachandran, Rajendra K. Kharul, Sylvain Marque, Pierre Soucy, Frédéric Jacques, Robert Chênevert, and Pierre Deslongchamps*. Synthetic Studies toward Highly Functionalized 5β-Lanosterol Derivatives: A Versatile Approach Utilizing Anionic Cycloaddition.

Page 6149. Olivier Lepage should be added to the paper as a coauthor. The author listing should be listed as follows: Sreekanth A. Ramachandran, Rajendra K. Kharul, Olivier Lepage, Sylvain Marque, Pierre Soucy, Frédéric Jacques, Robert Chênevert, and Pierre Deslongchamps.*

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Hermann A. Wegner, Helge Reisch, Karsten Rauch, Attila Demeter, Klaas A. Zachariasse, Armin de Meijere, and Lawrence T. Scott.*
Oligoindenopyrenes: A New Class of Polycyclic Aromatics.

Pages 9080−9087. All of the log ϵ values reported for the UV−vis absorption spectra in this paper were recorded incorrectly.

Page 9081. The last paragraph on this page should read:

Absorption Spectra of the Indenopyrenes. In their absorption spectra, the diindenopyrenes **3a** and **3b** exhibit two main absorption maxima (Figure 2), appearing at 267 nm (log ϵ = 4.55) and 307 nm (log ϵ = 4.65) for **3a** and at 264 nm (log ϵ = 4.56) and 316 nm (log ϵ = 4.49) for **3b**. These correlate with the absorption maxima for pyrene itself at 242 nm (log ϵ = 4.90) and 273 nm (log ϵ = 4.67) but are shifted to longer wavelengths.[11] The red colors of the diindenopyrenes result from their extended π-systems, which give rise to additional lower intensity maxima at 388 nm (log ϵ = 4.11) and 411 nm (log ϵ = 4.17) for **3b** and at even longer wavelength for **3a** at 410 nm (log ϵ = 4.15), 429 nm (log ϵ = 4.15), and 455 nm (log ϵ = 4.25). Monoindenopyrene **1** shows long wavelength absorptions also in this region at 376 nm (log ϵ = 4.14) and 386 nm (log ϵ = 4.06), which emphasizes the trend: the more indeno groups attached, the further the maxima shift to longer wavelength. This trend continues with the higher homologues of the indenopyrenes. The triindenopyrene **3c** exhibits strong maxima in its UV−vis spectrum at 276 nm (log ϵ = 4.71), 312 nm (log ϵ = 4.84) up to 325 nm (log ϵ = 4.55), and also in the region from 400 nm to beyond 500 nm. The tetraindenopyrene **3d** shows distinct absorption maxima well beyond 500 nm (545 nm (log ϵ = 3.79)) and an absorption tail that extends beyond 600 nm. These long wavelength absorption bands make the indenopyrenes appealing as candidates for new dyes and photoelectronic materials.[12]

Page 9086. Near the bottom of the first column, the absorption spectral data for diindeno[1,2,3-*cd*:1,2,3-*jk*]pyrene (**3a**) should read: UV λ_{max} ($CHCl_3$) nm (log ϵ) 455 (4.25), 429 (4.15), 410 (4.15), 389 (3.82), 334 (3.92), 307 (4.65), 298 (4.60), 267 (4.55), 260 (4.52), 248 (4.51).

Page 9087. Near the bottom of the first column, the absorption spectral data for 2,7,11-tri-*tert*-butyltriindeno[1,2,3-*cd*:1′,2′,3′-*fg*:1″,2″,3″-*jk*]pyrene (**3c**) should read: UV λ_{max} ($CHCl_3$) nm (log ϵ) 259 (4.62), 276 (4.71), 298 (4.70), 312 (4.84), 325 (4.55), 354 (4.33), 371 (4.18), 413 (4.08), 438 (4.23), 451 (4.05), 487 (3.30).

Page 9087. In the last paragraph of the Experimental Section, the absorption spectral data for 2,7,11,16-tetra-*tert*-butyltetraindeno[1,2,3-*cd*:1′,2′,3′-*fg*:1″,2″,3″-*jk*:1‴,2‴,3‴-*mn*]pyrene (**3d**) should read: UV λ_{max} ($CHCl_3$) nm (log ϵ) 278 (4.89), 288 (4.84), 298 (4.86), 312 (4.93), 333 (4.76), 363 (4.61), 397 (4.25), 419 (4.33), 437 (4.08), 467 (4.10), 545 (3.79).

Corrections also need to be made in the Supporting Information as follows:

Page S3. Near the bottom, the absorption spectral data for diindeno[1,2,3-*cd*:1′,2′,3′-*fg*]pyrene (**3b**) should read: UV λ_{max} ($CHCl_3$) nm (log ϵ) 551 (2.86), 512 (3.26), 454 (3.65), 411 (4.17), 388 (4.11), 366 (3.87), 332 (4.45), 327 (4.43), 316 (4.49), 282 (4.38), 264 (4.56).

Page S6. On lines 5 and 6, the absorption spectral data for 2,7,11-Tri-*tert*-butyltriindeno[1,2,3-*cd*:1′,2′,3′-*fg*:1″,2″,3″-*jk*-]pyrene (**3c**) should read: UV λ_{max} ($CHCl_3$) nm (log ϵ) 259 (4.62), 276 (4.71), 298 (4.70), 312 (4.84), 325 (4.55), 354 (4.33), 371 (4.18), 413 (4.08), 438 (4.23), 451 (4.05), 487 (3.30).

Page S8. In the last paragraph, the absorption spectral data for 2,7,11,16-tetra-*tert*-butyltetraindeno[1,2,3-*cd*:1′,2′,3′-*fg*:1″,2″,3″-*jk*:1‴,2‴,3‴-*mn*]pyrene (**3d**) should read: UV λ_{max} ($CHCl_3$) nm (log ϵ) 278 (4.89), 288 (4.84), 298 (4.86), 312 (4.93), 333 (4.76), 363 (4.61), 397 (4.25), 419 (4.33), 437 (4.08), 467 (4.10), 545 (3.79).

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