

# The Pictet–Spengler Condensation: A New Direction for an Old Reaction

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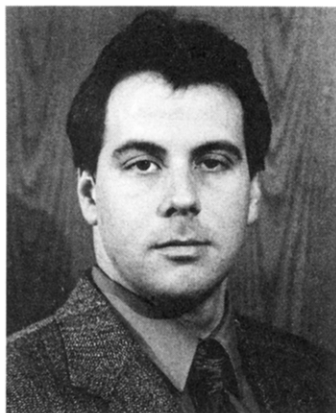
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## I. Introduction

The Pictet–Spengler reaction has long been an important reaction for the synthesis of both indole and isoquinoline alkaloids.<sup>1</sup> This condensation was discovered in 1911 by Amé Pictet and Theodor Spengler<sup>2</sup> when they condensed phenethylamine (**1**) with methylal to provide tetrahydroisoquinoline (**2**), as illustrated in Scheme 1.<sup>1,3,4</sup> The Pictet–Spengler

reaction was originally utilized exclusively to prepare tetrahydroisoquinolines. Conceived upon biogenetic grounds, it was felt that isoquinoline alkaloids were formed in plants by the condensation of  $\beta$ -arylethyl amines with carbonyl compounds.<sup>5</sup> Soon after the initial report, the Pictet–Spengler reaction became the standard method for the formation of tetrahydroisoquinolines. This process was first utilized with indole bases in 1928 by Tatsui during the preparation of 1-methyl-1,2,3,4-tetrahydro- $\beta$ -carboline (**4**).<sup>6</sup> The enzymatically catalyzed Pictet–Spengler condensation of tryptamine (**3**) with secologanin (**6**) to provide strictosidine<sup>7–11</sup> **7** is the key step in the biogenetic pathway to monoterpene indole alkaloids and is depicted in Figure 1. This condensation also occurs biogenetically with tryptophan (**5**).<sup>12,13</sup> The development of the enantiospecific Pictet–Spengler reaction in recent years has rendered this condensation an important synthetic method for the formation of macroline/sarpagine/ajmaline indole alkaloids.

The interest in the total synthesis of indole alkaloid natural products stems from their complex structures and diverse medicinal properties. For example, vincristine and vinblastine from *Catharanthus roseus*<sup>14,15</sup> have long been established as antitumor alkaloids<sup>16–18</sup> of clinical significance, while reserpine<sup>19,20</sup> as well as ajmaline<sup>21</sup> exhibit very important cardiovascular effects.<sup>7,22</sup> In regard to the present work, three bisindole alkaloids macralstonine acetate (**13**), macrocarpamine (**15**), and villalstonine (**16**), illustrated in Figure 2, have been found to exhibit significant activity *in vitro* against both *Entamoeba histolytica* and *Plasmodium falciparum*.<sup>23</sup> Macrocarpamine (**15**) was found to be the most active antiamebic alkaloid with an activity one-fourth that of the standard drug emetine. Villalstonine (**16**) was found to be the most potent of the three alkaloids tested against *P. falciparum* and was about 15 times less potent than the antimalarial drug chloroquine. Macralstonine acetate (**13**) was much more active against both types of protozoa when compared with the parent macralstonine (**14**). The use of *Alstonia angustifolia* in traditional medicine has been well documented and is, presumably, due in large part to the activity of these three alkaloids.<sup>24</sup> In addition, macralstonine (**14**) has also been shown to lower blood pressure in dogs.<sup>25,26</sup> The development of an enantiospecific route to these indole alkaloids, based upon a common intermediate, would therefore increase the amount of material available for study as well as provide the possibility for derivatization and greater potency. In addition, entry into the optical antipode of these bases would also be possible. Research on the stereospecific Pictet–Spengler condensation has led to the enantiospecific synthesis of such an intermedi-



Eric D. Cox was born in Milwaukee, WI, in 1965. He received his B.S. in Chemistry from the University of Wisconsin—Madison in 1988. From 1988 until 1991 he was employed at a biomedical company in St. Paul Minnesota where he worked in technical support. He joined Professor Cook's group at the University of Wisconsin—Milwaukee in 1991 and is currently working toward his Ph.D. in organic chemistry. His research interests include the design and synthesis of anxiolytic agents at the benzodiazepine receptor site. In addition he is currently working on the enantiospecific total synthesis of the bisindole alkaloid villalstonine and the indole alkaloid pleiocarpamine.

ate, the tetracyclic ketone. This intermediate will be discussed in detail later in this report. Work in this area was preceded with the discovery that the Pictet–Spengler reaction could be effected in nonacidic aprotic media as well as under the classical conditions of acid catalysis. Investigations to explore the scope of this reaction have led to an understanding of the factors which underlie the stereochemical control of this condensation. Detailed studies of the stereochemical and mechanistic factors which influence this process have been elucidated and are presented here along with the enantiospecific total synthesis of a number of indole alkaloid natural products made possible utilizing the key tetracyclic ketone mentioned above.

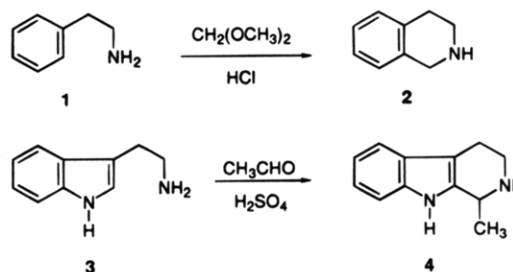
The Pictet–Spengler condensation has been of vital importance in the synthesis of numerous  $\beta$ -carbolines in addition to its use in the formation of indole alkaloids of more complex structure. Tetrahydro- $\beta$ -carbolines (THBC's) have been isolated from *Virola theiodora* and other plants of South American origin and employed by Indian tribes as a botanical source of intoxicating snuffs.<sup>27,28</sup> In addition, some THBC's derived from tryptamine and 5-methoxytryptamine have been shown to inhibit monoamine oxidase A and bind with nanomolar affinity in the central nervous system (CNS) to serotonin receptors.<sup>29–31</sup> Moreover, Langer et al.<sup>32</sup> have reported that 6-methoxy-THBC is present in high concentrations in the human pineal gland, and that it potently inhibited the high affinity binding of [<sup>3</sup>H]mipramine in human platelets. There is also a substantial amount of work which has suggested the involvement of THBC's in the etiology of alcoholism,<sup>33,34</sup> although, this has not yet been confirmed. The present work, however, represents the use of tryptophan alkyl esters in the Pictet–Spengler condensation and the tryptamine-related  $\beta$ -carbolines mentioned above will not be discussed in further detail.

The use of  $\beta$ -carbolines has been instrumental in the development of the inverse agonist/agonist phar-

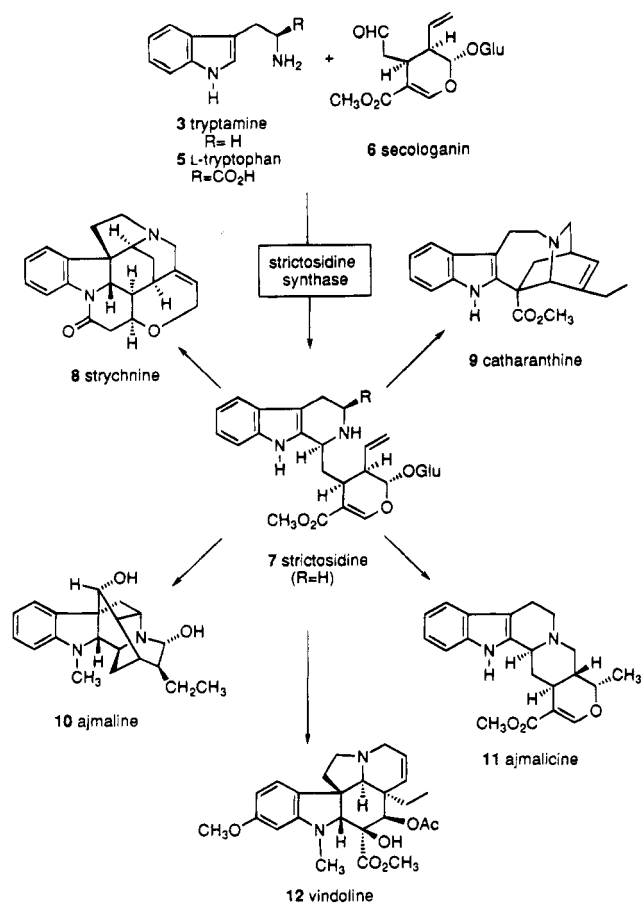


Professor James Cook received his B.S. degree in Chemistry, with honors, in 1967 from West Virginia University and his Ph.D. in 1971 from the University of Michigan. While at Michigan, he worked on the isolation and structure determination of monomeric and antihypertensive bisindole alkaloids from *Alstonia* species. During 1972–1973, he was a National Institutes of Health Postdoctoral Fellow at the University of British Columbia working on the synthesis of the antitumor alkaloids vincristine and vinblastine. He joined the faculty of the University of Wisconsin—Milwaukee in 1973 and has been Professor since 1986. An organic chemist by training, Professor Cook's interests include synthetic organic and natural products chemistry, as well as medicinal chemistry. His current research interests include the use of the chirally controlled Pictet–Spengler reaction for the total synthesis of antileukemic, antitumor, and antihypertensive indole alkaloids as well as the chemistry of reserpine, quinidine, and quinine in relation to their biological activity. This has culminated recently in an enantiospecific synthesis of the sarpagine- and ajmaline-related alkaloids, alstonerine, suaveoline, and raumacline as well as a partial synthesis of villalstonine. His group has carried out seminal studies on the use of the Weiss reaction for the synthesis of polyquinanes and is currently investigating this reaction as a source of strained polyquinenes to study bonding character in organic chemistry. The major thrust of his interest in medicinal chemistry is directed toward investigation of the structure, topology, and function of benzodiazepine (Valium) receptor subsites.  $\beta$ -Carbolines and diindoles which have been prepared in this study are important tools for studying anxiety, convulsions, sleep, and memory-learning, as well as reversal of the effect of Valium–alcohol or barbiturate–alcohol overdose. Many of these analogs are Bz<sub>1</sub> and Bz<sub>5</sub> receptor subsite specific agents. Students in his group have also recently designed inhibitors of the multidrug resistance pump (MDR) of drug-resistant strains of cancer cells as well as prepared inhibitors of the indolamine 2,3-dioxygenase enzyme. This latter enzyme is involved in the pathogenesis of many inflammatory diseases including the dementia experienced by AIDS patients.

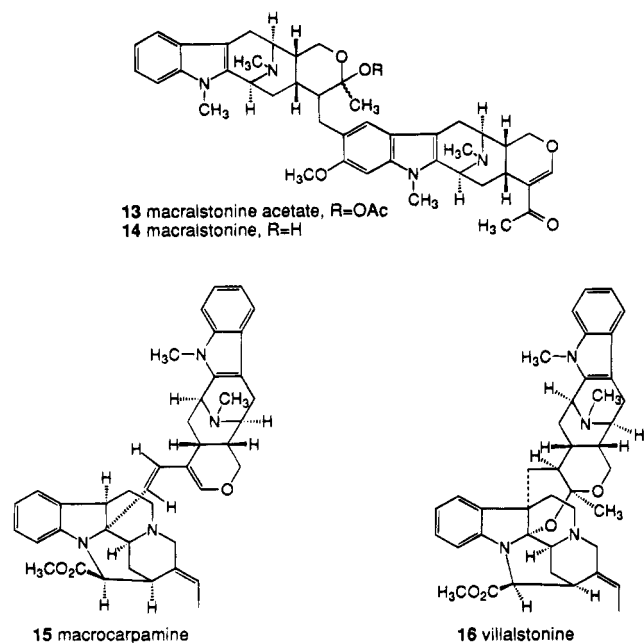
#### Scheme 1



macophore of the benzodiazepine receptor (BzR) site. The BzR is located on the GABA<sub>A</sub> receptor ion channel and plays a central role in the molecular mechanisms controlling anxiety,<sup>35,36</sup> convulsions, memory-learning,<sup>37</sup> and sleep.<sup>38</sup> Positive or negative modulation of the GABA<sub>A</sub> receptor system by various structural classes of ligands has resulted in markedly different pharmacological effects. Agonist ligands are widely used as anxiolytics, anticonvulsants, and myorelaxants. In contrast, there are ligands which elicit pharmacological actions opposite to those of



**Figure 1.** The strictosidine synthase-mediated Pictet–Spengler condensation of tryptamine with secologanin to provide the key biogenetic intermediate strictosidine.



**Figure 2.** Several important bisindole alkaloids isolated from *Alstonia* species.

agonist ligands. These are known as inverse agonists and are represented here by  $\beta$ -carboline; they exhibit anxiogenic, somnolytic, convulsant, and proconvulsant activity. At this time there is a need for selective inverse agonist and agonist ligands to be employed in the treatment of a variety of processes mediated in the CNS. More specifically, there is a

need for benzodiazepine (Valium) receptor partial agonists which elicit anxiolytic and anticonvulsant effects but are devoid of the muscle relaxant and sedative side effects of classical 1,4-benzodiazepines. Moreover, partial or selective inverse agonists which enhance neuronal firing in the CNS in the absence of proconvulsant or convulsant<sup>39–43</sup> effects represent agents with potential therapeutic use as drugs to enhance cognition, to reverse the effects of hepatic encephalopathy, or to reverse the effects of barbiturate–alcohol-induced CNS depression (overdose).<sup>44</sup> A thorough understanding of the factors affecting these site(s) will permit the development of highly selective psychoactive drugs. Extensive studies using computer graphics of the inverse agonist and agonist pharmacophores coupled with synthetic results has permitted the development of the partial inverse agonist 3-EBC and the partial agonist 6-PBC as well as the Bz<sub>1</sub> selective antagonist  $\beta$ Cct. The results of these studies will be presented in this review as well as future plans regarding medicinal chemistry in the benzodiazepine receptor area.

## II. Pictet–Spengler Condensations in Nonacidic Aprotic Media

In the course of work directed toward the construction of potential antihypertensive agents, the need arose for the preparation of *N*<sub>b</sub>-benzyltryptophan methyl ester (21). This ester can be prepared by stirring tryptophan methyl ester 17 and benzaldehyde 18 in benzene at room temperature, followed by reduction of the resulting imine 19 with sodium borohydride, similar to the work of Yoneda (Scheme 2).<sup>45</sup> To improve the conversion of ester 17 into imine 19, benzaldehyde and amine 17 were heated in benzene at reflux while a Dean–Stark trap was employed to remove water formed during the process. Although the imine 19 was initially observed, after prolonged heating the products of this reaction were the *cis* and *trans* diastereomers of 1-phenyl-3-(methoxycarbonyl)-1,2,3,4-tetrahydro- $\beta$ -carboline 20 in 95% yield. This result was surprising for generally the Pictet–Spengler reaction had been carried out in a protic solvent with acid catalysts.<sup>1,46–52</sup> Jackson et al. earlier reported that tryptamine (3) and benzaldehyde (18) yielded only imine when heated in benzene at reflux.<sup>53</sup> Presumably, the Pictet–Spengler reaction with tryptophan methyl ester 17 had occurred without the aid of acid catalysts, therefore, it was decided to make a detailed study of this observation.<sup>54</sup>

### Scheme 2

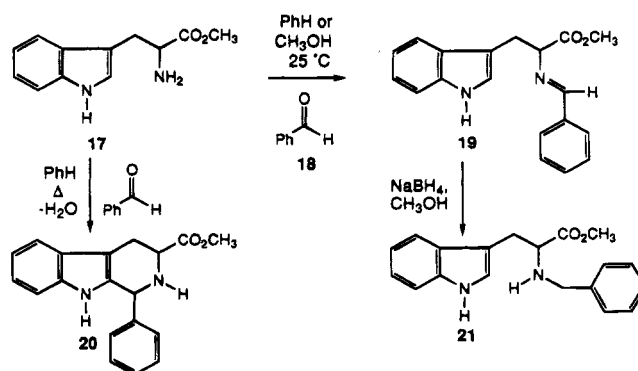
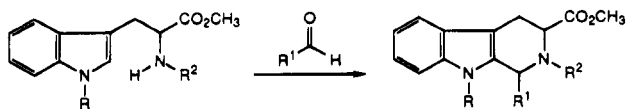
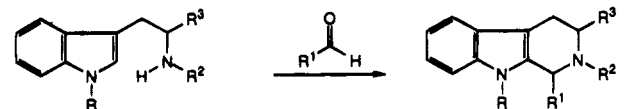


Table 1. Pictet–Spengler Cyclization of Tryptophan Methyl Ester Derivatives



amine	R	R <sup>2</sup>	aldehyde	product	R <sup>1</sup>	aprotic % yield	aqueous % yield
<b>17</b>	H	H	<b>18</b>	<b>20</b>	Ph	95	90
<b>17</b>	H	H	<b>23</b>	<b>25</b>	C <sub>6</sub> H <sub>11</sub>	85	73
<b>21</b>	H	Bn	<b>18</b>	<b>26</b>	Ph	95	
<b>21</b>	H	Bn	<b>23</b>	<b>27</b>	C <sub>6</sub> H <sub>11</sub>	87	
<b>22</b>	CH <sub>3</sub>	Bn	<b>23</b>	<b>28</b>	C <sub>6</sub> H <sub>11</sub>	87	68
<b>21</b>	H	Bn	<b>24</b>	<b>29</b>	CH <sub>2</sub> CH <sub>2</sub> CO <sub>2</sub> C <sub>2</sub> H <sub>5</sub>	80	50

Table 2. Reaction of Amines with Acid Labile Aldehydes



amine	R	R <sup>2</sup>	R <sup>3</sup>	aldehyde	R <sup>1</sup>	product	aprotic % yield	aqueous % yield
<b>22</b>	CH <sub>3</sub>	Bn	CO <sub>2</sub> CH <sub>3</sub>	<b>32</b>	HC(OC <sub>2</sub> H <sub>5</sub> ) <sub>2</sub>	<b>35</b>	90	<20
<b>30</b>	H	Bn	H	<b>32</b>	HC(OC <sub>2</sub> H <sub>5</sub> ) <sub>2</sub>	<b>36</b>	92	<25
<b>31</b>	CH <sub>3</sub>	H	CO <sub>2</sub> CH <sub>3</sub>	<b>32</b>	HC(OC <sub>2</sub> H <sub>5</sub> ) <sub>2</sub>	<b>37</b>	65	<15
<b>17</b>	H	H	CO <sub>2</sub> CH <sub>3</sub>	<b>32</b>	HC(OC <sub>2</sub> H <sub>5</sub> ) <sub>2</sub>	<b>38</b>	62	<15
<b>21</b>	H	Bn	CO <sub>2</sub> CH <sub>3</sub>	<b>32</b>	HC(OC <sub>2</sub> H <sub>5</sub> ) <sub>2</sub>	<b>39</b>	75	<25
<b>22</b>	CH <sub>3</sub>	Bn	CO <sub>2</sub> CH <sub>3</sub>	<b>33</b>	C <sub>2</sub> H <sub>4</sub> COCH <sub>2</sub> CH <sub>3</sub>	<b>40</b>	94	25
<b>22</b>	CH <sub>3</sub>	Bn	CO <sub>2</sub> CH <sub>3</sub>	<b>34</b>	CH <sub>2</sub> CH <sub>2</sub> CO <sub>2</sub> H <sup>a</sup>	<b>41</b>	97	50
<b>22</b>	CH <sub>3</sub>	Bn	CO <sub>2</sub> CH <sub>3</sub>	<b>24</b>	CH <sub>2</sub> CH <sub>2</sub> CO <sub>2</sub> C <sub>2</sub> H <sub>5</sub>	<b>42</b>	75	59

<sup>a</sup> Carbonyl substrate was  $\alpha$ -ketoglutaric acid.

A variety of tryptophan methyl ester derivatives have been employed in this condensation and some of these are outlined in Table 1. Excellent yields of tetrahydro- $\beta$ -carboline were obtained with tryptophan methyl ester **17**, *N*<sub>b</sub>-benzyltryptophan methyl ester (**21**), and *N*<sub>a</sub>-methyl-*N*<sub>b</sub>-benzyltryptophan methyl ester **22**. It was apparent from examination of the data in Table 1, that yields of the Pictet–Spengler reaction could be improved in nonacidic aprotic media. This was even more obvious when acid-labile aldehydes were used as substrates, as illustrated in Table 2. For example, the reaction of tryptophan methyl ester derivatives **17**, **21**, **22**, **30**, and **31** with glyoxal diethyl acetal **32** in benzene at reflux resulted in good to excellent yields of the corresponding tetrahydro- $\beta$ -carboline.<sup>55</sup> These same cyclizations carried out under aqueous acidic conditions resulted in considerably poorer yields of the tetrahydro- $\beta$ -carboline. The yields were generally two to three times better under nonacidic aprotic conditions than those obtained in an aqueous acidic medium. From the data in Table 2 it was obvious the Pictet–Spengler reaction could be extended to include aldehydes containing functionality such as acetals, esters, amides, and acetonides, heretofore too labile to be practical for this condensation.

Jackson and co-workers have extensively studied two possible pathways for the Pictet–Spengler reaction between tryptamine derivatives and carbonyl compounds.<sup>53,56,57</sup> Regardless of which path occurs, it is the electrophilic nature of the imine double bond which is the driving force of the cyclization.<sup>54</sup> Performing the reaction in nonacidic, aprotic media permitted a study of the correlation between the electron density on the aliphatic nitrogen atom with

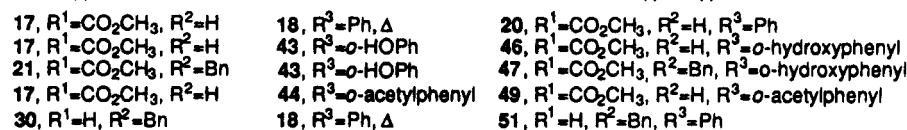
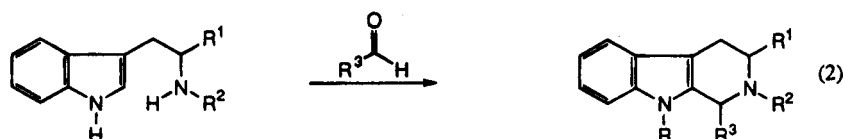
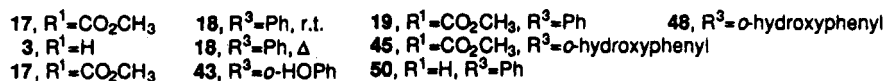
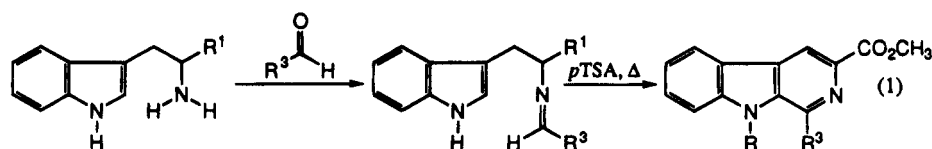
the ease of cyclization since protonation of the nitrogen atom by solvent was no longer a complicating factor.

The reaction of tryptophan methyl ester (**17**) with benzaldehyde (**18**) in benzene at reflux resulted in the tetrahydro- $\beta$ -carboline **20** in 90% yield. Conversely, the reaction of tryptamine (**3**) under the analogous conditions resulted only in the formation of the Schiff's base **50** in quantitative yield, as illustrated in Table 3. These results can be rationalized by examination of the  $pK_a$  values of the two bases: tryptamine,  $pK_a = 10.2$ ,<sup>58</sup> tryptophan methyl ester,  $pK_a = 7.29$ .<sup>59</sup> The tryptophan methyl ester imine intermediate **19** is clearly more electrophilic than the imine **50**, formed from tryptamine (**3**). In fact, even in the presence of small amounts of tryptophan or tryptophan methyl ester hydrochloride the tryptamine intermediate **50** failed to cyclize.

It was reported in 1974 by Hamaguchi et al.<sup>46</sup> that the condensation of salicylaldehyde (**43**) with tryptophan methyl ester (**17**) in acidic media provided the Pictet–Spengler product **46** (Table 3, eq 2) as a mixture of the *cis* and *trans* isomers **46a,b**, albeit in 3.5% overall yield. This same condensation carried out in benzene or toluene at reflux (Table 3, eq 1) provided only the imine **45**. This imine was so unreactive in fact, that cyclization was only achieved by heating in toluene at reflux for 48 h in the presence of *p*-toluenesulfonic acid. These harsh reaction conditions failed to provide the tetrahydro- $\beta$ -carboline, but rather led to the fully aromatic  $\beta$ -carboline **48** (Table 3, eq 1). These results can be rationalized by the mesomeric effect of the phenolic oxygen atom. The electron release by this atom rendered the imine double bond of tryptamine **45** less



Table 3. Influence of Electrophilic Character of Imine on Cyclization

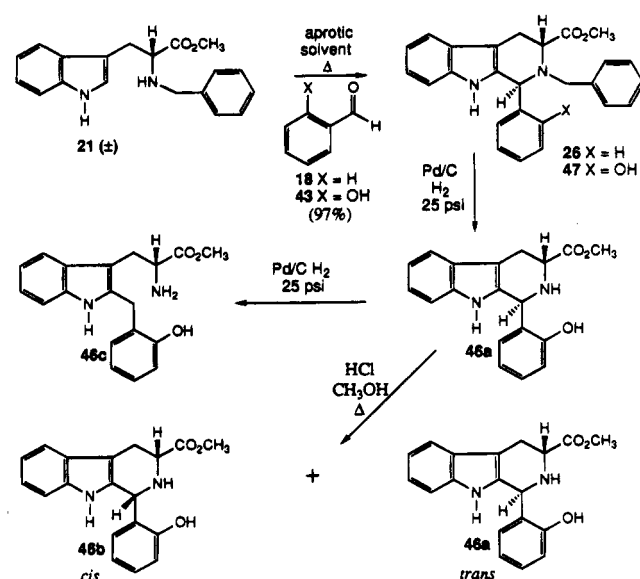


amine	R <sup>1</sup>	R <sup>2</sup>	aldehyde (Δ)	R <sup>3</sup>	product	medium	% yield	amine	R <sup>1</sup>	R <sup>2</sup>	aldehyde (Δ)	R <sup>3</sup>	product	medium	% yield
17	CO <sub>2</sub> CH <sub>3</sub>	H	18	Ph	20	benzene	90	17	CO <sub>2</sub> CH <sub>3</sub>	H	43	<i>o</i> -hydroxyphenyl	48	toluene/ pTSA	5
17	CO <sub>2</sub> CH <sub>3</sub>	H	43	<i>o</i> -hydroxyphenyl	45	toluene	100	17	CO <sub>2</sub> CH <sub>3</sub>	H	44	<i>o</i> -acetylphenyl	49	benzene	40
17	CO <sub>2</sub> CH <sub>3</sub>	H	43	<i>o</i> -hydroxyphenyl	46	methanol/ water	3.5	3	H	H	18	Ph	50	benzene	100
21	CO <sub>2</sub> CH <sub>3</sub>	Bn	43	<i>o</i> -hydroxyphenyl	47	benzene	62	30	H	Bn	18	Ph	51	benzene	98.5
21	CO <sub>2</sub> CH <sub>3</sub>	Bn	43	<i>o</i> -hydroxyphenyl	47	toluene	97	17	CO <sub>2</sub> CH <sub>3</sub>	H	18 (rt)	Ph	19	benzene	100

electrophilic in comparison to the same bond in imine 19. Further support for this hypothesis was obtained by the formation of *cis* and *trans* isomers of *o*-acetoxyphenyl-1,2,3,4-tetrahydro- $\beta$ -carboline 49 prepared in moderate yield (Table 3) from the condensation of acetylsalicylaldehyde (44) and tryptophan methyl ester (17) in benzene at reflux. In this case the electron-withdrawing effect of the acetyl group retarded the electron-donating capability of the phenolic oxygen atom and permitted cyclization to occur in the desired manner to provide the tetrahydro- $\beta$ -carboline represented by 49.

As alluded to earlier, the electrophilicity of the imine double bond was the limiting factor which determined whether cyclization would be achieved in nonacidic aprotic medium. With this in mind, the use of the *N*<sub>b</sub>-benzyl group would provide an electrophilic iminium ion intermediate which should readily undergo cyclization. In addition, the benzyl moiety could be easily removed later by catalytic hydrogenation. The results of this investigation are shown in Scheme 3 and the yields are listed in Table 3. The reaction of *N*<sub>b</sub>-benzyltryptophan methyl ester (21) with salicylaldehyde (43) in toluene at reflux provided tetrahydro- $\beta$ -carboline 47 in 97% yield. Catalytic debenzylation of 47 with Pd/C at 25 psi (H<sub>2</sub>) gave the *trans*-tetrahydro- $\beta$ -carboline 46a in 55–70% yield accompanied by the 2-benzyl derivative 46c, the product of a second hydrogenolysis. In every case in which Sandrin et al.<sup>55</sup> examined the substitution of a benzyl group on the aliphatic nitrogen atom of either tryptamine (3) or tryptophan methyl ester (21), the effect has been to speed the rate of the cyclization and to improve the yield. Furthermore, cyclization of an imine which contained an *N*<sub>b</sub>-isopropyl group

Scheme 3



(Table 1) resulted in lower yields of product which may be due to steric effects, electronic effects, or presumably, to a mixture of both effects.<sup>54,60</sup>

The utilization of high boiling solvents such as cumene or diglyme to facilitate the Pictet–Spengler cyclization without aldehyde decomposition is an advantage of the reaction in aprotic nonacidic media.<sup>46,52</sup> The synthetic potential of the Pictet–Spengler cyclization in aprotic media with either tryptophan methyl ester derivatives or *N*<sub>b</sub>-benzyltryptamines is quite general and can be employed with a variety of aldehydes which may contain acid-labile functional groups. The optimum conditions for

this modification appear to be the use of *N*<sub>1</sub>-benzyl derivatives in benzene or toluene at reflux. Care must be taken to assure that the boiling point of the aldehyde is higher than that of the solvent; aldehydes such as acetaldehyde readily distill off into the Dean–Stark trap and give poor yields of tetrahydro- $\beta$ -carbolines. In these cases the use of a sealed tube provides improved yields.<sup>61,62</sup> For substances that are not soluble in benzene or toluene, dioxane can be added to the reaction medium.

### A. Stereochemical Assignment of 1,3-Disubstituted 1,2,3,4-Tetrahydro- $\beta$ -carbolines

The use of nonacidic, aprotic media in the Pictet–Spengler reaction was a significant improvement upon standard reaction conditions for this condensation. Because this process resulted in the formation of *cis* and *trans* isomers it became necessary to establish a general method for the assignment of stereochemistry to differentiate between these diastereomeric 1,3-disubstituted 1,2,3,4-tetrahydro- $\beta$ -carbolines.

The use of ORD/CD had been successfully employed by Brossi et al.<sup>63</sup> in the 1-methyl-3-carboxyl series; however, accurate assignments using this technique required pure samples and such diastereomers were often difficult to separate. Inconsistent results often occurred with proton NMR due to the overlap of signals from the protons located at C-1. In addition, the assignment of stereochemistry by <sup>1</sup>H NMR for the *cis* and *trans* isomers in the 1-phenyl-3-methoxycarbonyl series (Scheme 4) has led to conflicting assignments.<sup>46,64</sup> Chemical correlation (preferential cyclization of only the *cis* isomer) of stereochemistry carried out by Smith<sup>13</sup> required the synthesis of specific compounds with the proper functionality at carbon atoms 1 and 3 to permit cyclization. This chemical approach was considered too laborious to be employed in a general sense, consequently attention turned to <sup>13</sup>C NMR spectroscopy.

Assignments in the 1-phenyl-3-methoxycarbonyl series using carbon-13 magnetic resonance spectroscopy had been reported in preliminary fashion by Sandrin et al. in 1976.<sup>65</sup> Moreover the use of <sup>13</sup>C NMR for the stereochemical assignment and structure proof of a number of yohimbinoid and ajmalcinoid alkaloid systems was also published in that same year by Wenkert et al.<sup>66</sup> It became clear that carbon NMR spectroscopy would be devoid of the complications observed in proton NMR; therefore, studies were initiated to determine the limits of the method described earlier.<sup>67</sup> The <sup>13</sup>C NMR method developed by Sandrin et al.<sup>65</sup> employed the well-documented compression effect<sup>68</sup> observed in <sup>13</sup>C NMR spectroscopy, and will be illustrated only briefly for the *cis*- and *trans*-tetrahydro- $\beta$ -carbolines **20a** and **20b**.

The *cis* and *trans* isomers were prepared using a Pictet–Spengler condensation between tryptophan methyl ester (**17**) and benzaldehyde (**18**) in nonacidic aprotic media<sup>54</sup> or in an aqueous acidic media (Scheme 4). The *cis* and *trans* diastereomers were isolated, separated, and the <sup>13</sup>C NMR spectrum of each was recorded. The signal assignments were made on the basis of nuclear Overhauser effects (nOe), correlations with known compounds, and off-resonance decoupled spectra. The carbon signals for carbon atoms C-1 and C-3 for the compounds in Scheme 4 are illustrated in Table 4. The carbon signals assigned to the *cis* isomer **20a** [C-1 (58.7 ppm) and C-3 (56.9 ppm)] were downfield relative to those of isomer **20b** [C-1 (54.9 ppm) and C-3 (52.3 ppm)] which was then assigned as the *trans* diastereomer. Conformational analysis and examination of molecular models indicated that of the two possible twist chair conformations (A or B, Scheme 5) for the *trans* isomer **20b**, conformer A should represent the structure of the more stable species due to the 1,4-gauche interaction between the hydrogen atom at C-1 and the substituent located at C-3 in conformer B. More importantly,

Scheme 4

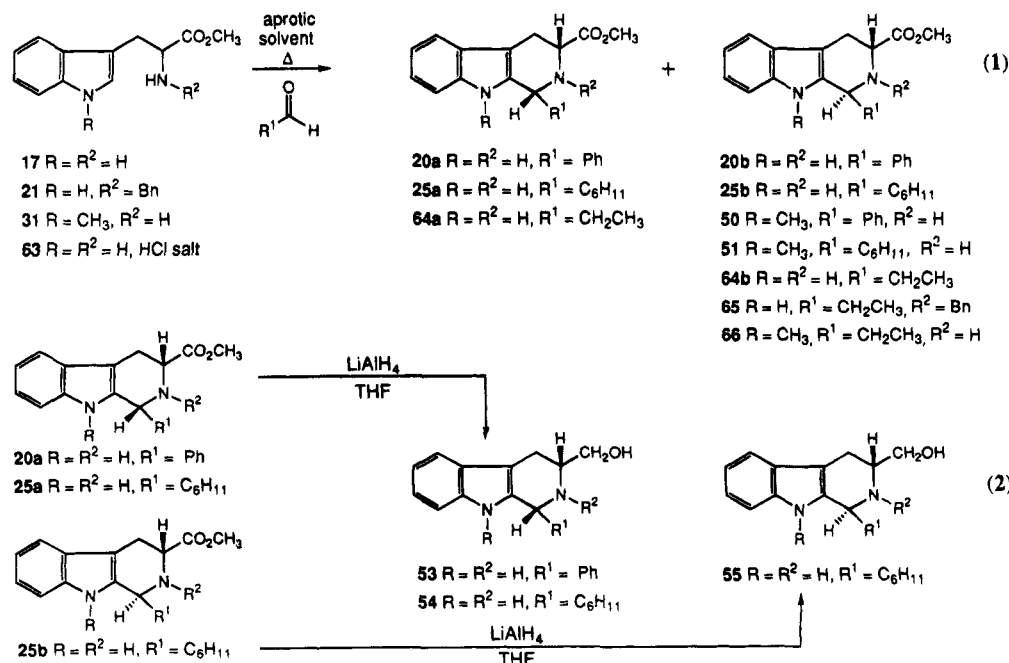
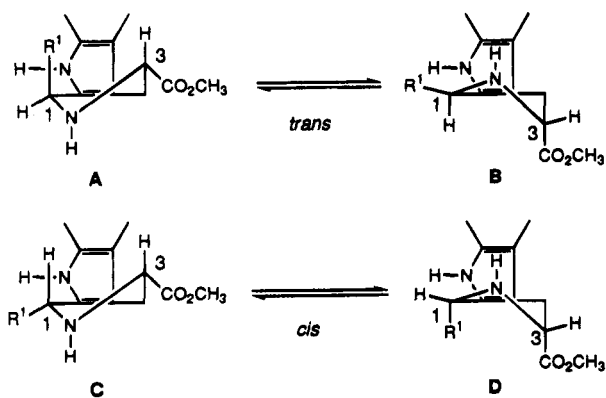


Table 4.  $^{13}\text{C}$  NMR Signals for C-1 and C-3

compound	stereochemical configuration	$^{13}\text{C}$ signal C-1 (ppm)	$^{13}\text{C}$ signal C-3 (ppm)
<b>20a</b>	<i>cis</i>	58.7	56.9
<b>20b</b>	<i>trans</i>	54.9	52.3
<b>25a</b>	<i>cis</i>	57.7	56.6
<b>25b</b>	<i>trans</i>	55.4	53.4
<b>50</b>	<i>trans</i>	54.8	51.1
<b>51</b>	<i>trans</i>	54.5	52.8
<b>52</b>	<i>cis</i> C(1) = $^2\text{H}$		57.0
<b>53</b>	<i>cis</i>	58.8	56.4
<b>54</b>	<i>cis</i>	57.9	55.8
<b>55</b>	<i>trans</i>	55.9	51.3
<b>56</b>	<i>cis</i>	53.1	56.5
<b>58</b>	<i>trans</i>	52.8	52.7
<b>60</b>	<i>trans</i>	49.9	52.9
<b>64a</b>	<i>cis</i>	53.8	56.5
<b>64b</b>	<i>trans</i>	51.7	52.8
<b>66</b>	<i>trans</i>	51.1	52.2

Scheme 5



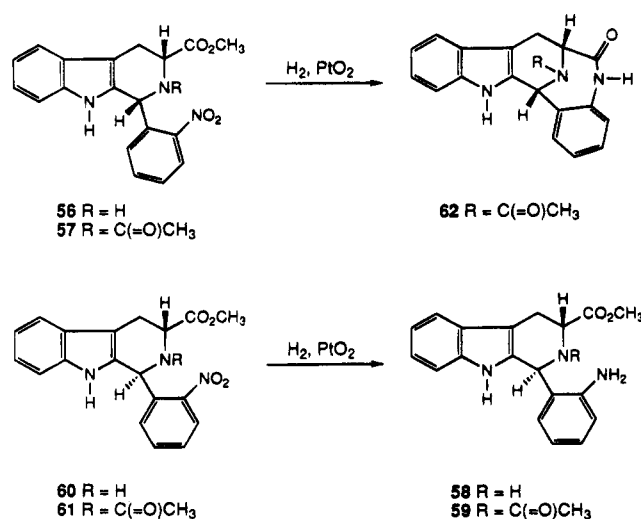
conformer B undergoes an unfavorable interaction between the indole  $N_a$ -H and the equatorial phenyl group located at C-1 [A<sup>(1,2)</sup> strain].<sup>69</sup> The methoxycarbonyl group at C-3 is located in the equatorial position in the more stable conformer A while the substituent at C-1, although now in the axial position, is devoid of the interaction between the substituent at C-1 and the  $N_a$ -indole hydrogen atom. A similar analysis of the *cis* diastereomer **20a** led to the conclusion that conformer C (with no  $^{13}\text{C}$ - $\gamma$  gauche effect between atoms bonded to carbons 1 and 3) should be much more stable than conformer D. Consequently, it would be expected that the signals for carbon atoms 1 and 3 in *trans* isomer **20b** (with a  $^{13}\text{C}$ - $\gamma$  effect) would appear at a higher field in the carbon NMR spectrum, respectively, than those of the corresponding *cis* isomer. This was found to be the case in 1-cyclohexyl-3-methoxycarbonyl bases **25a** (*cis*) and **25b** (*trans*) (Scheme 4) as well as the 1-phenyl compounds discussed above. Additional data were required, moreover, to support the assumption that the assignment of *cis* and *trans* configurations to the above indoles was correct and that the 1,3-disubstituted 1,2,3,4-tetrahydro- $\beta$ -carboline were not subject to other subtle influences which might lead to erroneous conclusions.

In this regard  $N_a$ -methyltryptophan methyl ester **31** was heated at reflux in an aprotic solvent with either benzaldehyde (**18**) or cyclohexanecarboxaldehyde (**23**) which provided the corresponding 1-substituted derivatives **50** and **51**, respectively, in excellent yields. Examination of molecular models had

indicated that the A<sup>(1,2)</sup> strain between the substituent at C-1 and the  $N_a$ -methyl group would permit only formation of the *trans* diastereomer under these conditions. In agreement with this, only one diastereomer was isolated from each synthesis (>85% yield); furthermore, the carbon signals for *trans* diastereomer **50** [C-1 (54.8 ppm) and C-3 (51.1 ppm)] were virtually identical to those of the *trans* isomer **20b** [C-1 (54.9 ppm) and C-3 (52.3 ppm)]. This same phenomenon was observed in the case of *trans* diastereomer **51**. While the latter two experiments demonstrated indirectly the validity of this approach, additional experiments were necessary to accurately determine which signals in Table 4 were due to the resonance at C-1.

For this purpose tryptophan methyl ester **17** was condensed with deuterobenzaldehyde<sup>70</sup> in the presence of *p*-toluenesulfonic acid (*p*TSA) to provide a mixture of two components, the  $R_f$  values of which were identical to those of phenyl analogs **20a** and **20b**. These diastereomers were separated by chromatography and the  $^{13}\text{C}$  NMR spectrum (except at C-1) and melting point of the more accessible *cis* compound **52** (Table 4) were found to be in complete agreement with a structure such as **20a** (**52** 1<sub>d</sub>). The carbon spectrum of **52** 1<sub>d</sub> was devoid<sup>71,72</sup> of the signal at 58.7 ppm due to C-1 in **20a**; therefore, it was clear that the signal at 58.7 ppm in **20a** resulted from the carbon atom at position 1 while that at 56.9 ppm corresponded to carbon atom C-3. To further corroborate this assignment and to extend this to the cyclohexyl bases **25a** and **25b**, *cis*-1-phenyl-**20a**, *cis*-1-cyclohexyl-**25a**, and *trans*-1-cyclohexyl-3-(methoxycarbonyl)-1,2,3,4-tetrahydro- $\beta$ -carboline (**25b**) were treated with  $\text{LiAlH}_4$  in THF. The 3-hydroxymethyl-substituted bases were isolated and subjected to  $^{13}\text{C}$  NMR spectroscopy (Table 4). In all cases [**20a**  $\rightarrow$  **53** ( $\delta$  -0.5 ppm), **25a**  $\rightarrow$  **54** ( $\delta$  -0.8 ppm), **25b**  $\rightarrow$  **55** ( $\delta$  -2.1 ppm) the signal for the carbon atom at C-3 was shifted upfield by 0.5 ppm or more; the same phenomenon had been observed on going from methyl acetate to ethanol ( $\delta$  -1.7 ppm).<sup>73</sup> Having now determined the chemical shifts for carbon atoms 1 and 3 in the 1-phenyl and 1-cyclohexyl series, direct chemical proof was required to establish that the  $^{13}\text{C}$  NMR spectroscopic method was correct in the most unequivocal sense. Smith and co-workers<sup>13</sup> had shown that *cis*-1-(hydroxymethyl)-3-(methoxycarbonyl)-1,2,3,4-tetrahydro- $\beta$ -carboline (as the  $N_b$ -amide) would cyclize to a lactone, while the *trans* isomer would not. Spenser<sup>74</sup> had demonstrated, however, that these 1-hydroxymethyl derivatives were quite labile and consequently the *cis*- and *trans*-1-(*o*-nitrophenyl) bases depicted in Scheme 6 were chosen for study. A mixture of *cis* and *trans* diastereomers were obtained from the Pictet–Spengler condensation of **17** with *o*-nitrobenzaldehyde, and the two compounds (**56** and **60**) were carefully separated by chromatography. The assignment of stereochemistry was initially based upon the carbon spectrum of these two molecules. When the *trans* isomer was subjected to catalytic hydrogenation, only 2-amino derivative **58** was isolated; however, the *cis* diastereomer **56**, under analogous conditions gave a complex mixture of products. According to Hobson<sup>75</sup> the formation of

## Scheme 6



an amide would flatten the twist chair conformation of ring C of a tetrahydro- $\beta$ -carboline and would bring the two groups *cis* 1,3-disposed into closer proximity to permit more facile cyclization. In view of this, both *o*-nitro bases **56** (*cis*) and **60** (*trans*) were converted into the corresponding acetamide derivatives **57** and **61**, respectively, by treatment with acetic anhydride and pyridine. The *trans* amide **61** gave only amide **59** when treated with hydrogen and PtO<sub>2</sub> while the *cis* isomer under the same reaction conditions was converted into the pentacyclic lactam **62** in 30% yield. This latter transformation clearly represented chemical proof of the validity of the <sup>13</sup>C NMR method for stereochemical assignments when position 1 was substituted with an aryl group.

Attention now turned to tetrahydro  $\beta$ -carbolines which contained smaller substituents at C-1. In this vein, it was decided to synthesize the *cis* and *trans* diastereomers of 1-ethyl-substituted tetrahydro- $\beta$ -carboline (**64a,b**) from amino ester **63** and propionaldehyde **68**, as illustrated in Scheme 4. Brossi<sup>63</sup> had shown that 1,3-disubstituted 1,2,3,4-tetrahydro- $\beta$ -carbolines which have a methyl group at C-1 are found predominantly as the *cis* isomer. It should be noted, however, that extrapolation to larger groups can lead to erroneous conclusions.<sup>64</sup> The condensation of **63** with propionaldehyde (**68**) in acidic solution provided the 1,3-disubstituted 1,2,3,4-tetrahydro- $\beta$ -carbolines as a mixture of *cis* and *trans* isomers represented by **64a** and **64b**, respectively. The diastereomeric ratio depicted in Table 5 suggested that A<sup>(1,2)</sup> strain was significant enough to favor the *trans*-1-ethyl diastereomer **64b**. Separation of the two diastereomers followed by examination of their <sup>13</sup>C NMR spectra led to the initial assignment of stereochemistry. Following the initial assignment of stereochemistry attempts were made to corroborate the findings by another method. Chemical correlation of the *cis* isomer (via cyclization) was not possible, however, Ungemach et al.<sup>76</sup> had reported earlier that *N*<sub>b</sub>-benzyltryptophan methyl ester (**21**) reacted with either salicylaldehyde (**43**), glyoxal diethyl acetal (**32**), or propionaldehyde (**68**) to provide only the *trans*-1,3-disubstituted  $\beta$ -carboline (e.g. **65**, illustrated in Scheme 4). When the *trans* base **65** was subjected to catalytic hydrogenation this resulted

Table 5. Ratio of *Cis/Trans* Diastereomers

compound	R	R <sup>1</sup>	diastereomeric ratio <sup>a</sup> ( <i>cis:trans</i> )	ref
<b>20a, 20b</b>	H	Ph	40:60	46
<b>25a, 25b</b>	H	C <sub>6</sub> H <sub>11</sub>	40:60	46
<b>50</b>	CH <sub>3</sub>	Ph	0:100	46
<b>51</b>	CH <sub>3</sub>	C <sub>6</sub> H <sub>11</sub>	0:100	46
<b>56, 60</b>	H	2-NO <sub>2</sub> C <sub>6</sub> H <sub>5</sub>	25:75	71
<b>64a, 64b</b>	H	CH <sub>2</sub> CH <sub>3</sub>	43:57	56
<b>66</b>	CH <sub>3</sub>	CH <sub>2</sub> CH <sub>3</sub>	0:100	56

<sup>a</sup> In a pair of diastereomers the second number represents the *trans* isomer in this table.

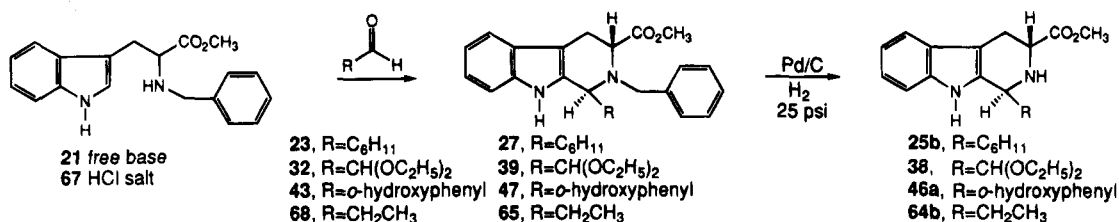
in the formation of 1,3-disubstituted  $\beta$ -carboline **64b** as the only isolable material. In addition, excellent agreement was observed when comparison of the chemical shifts for carbon atoms 1 and 3 in the spectrum of **64b** (51.7, 52.8 ppm) was made with the signals in the <sup>13</sup>C NMR spectrum of *trans* **66** [51.1, 52.2 ppm (*N*<sub>a</sub>-methyl)]. More importantly, the *trans* stereochemistry of the 1,3-disubstituted 1,2,3,4-tetrahydro- $\beta$ -carboline **64b** was confirmed by X-ray crystallography.<sup>77</sup>

In all of the examples examined here, the signals for C-1 and C-3 of the *cis* diastereomers are clearly distinct from the analogous resonance lines for the *trans* diastereomers. The enantiospecific synthesis of indole alkaloids in the macroline/sarpagine/ajmaline series rests on the accurate assignment of the stereogenic centers at C-1 and C-3 in the corresponding 1,3-disubstituted tetrahydro- $\beta$ -carbolines. Although Bailey et al. have reported a <sup>13</sup>C NMR method to differentiate between *cis* and *trans* diastereomers in the *N*<sub>a</sub>-hydrogen-*N*<sub>b</sub>-benzyl series, this method is not 100% effective as noted by the authors.<sup>78</sup> Moreover, Tóth et al. have examined this method for stereochemical assignments and also found exceptions.<sup>79</sup> To date, accurate stereochemical assignments for the *cis*- and *trans*-1,3-disubstituted *N*<sub>b</sub>-benzyltetrahydro- $\beta$ -carbolines in the above series can only be made in 100% of these cases by removal of the *N*<sub>b</sub>-benzyl group [catalytic debenzylation (CTH): Pd/C, NH<sub>4</sub>CO<sub>2</sub>H, CH<sub>3</sub>CH<sub>2</sub>OH] followed by identification of the diastereomers by the <sup>13</sup>C NMR method developed by Sandrin and Ungemach.<sup>65,77</sup> In addition, this <sup>13</sup>C NMR technique<sup>77</sup> provided for the first time a facile method to study the effects of A<sup>(1,2)</sup> strain in the determination of configurational preference of 1,3-disubstituted 1,2,3,4-tetrahydro- $\beta$ -carbolines in the Pictet–Spengler cyclization. With these results in hand, attention now turned to the stereospecific synthesis of *trans*-1,3-disubstituted-1,2,3,4-tetrahydro- $\beta$ -carbolines. Many of these tetrahydro- $\beta$ -carbolines have demonstrated interesting biological activity or are important intermediates in the total synthesis of indole alkaloids.

## B. Stereospecific Synthesis of *trans*-1,3-Disubstituted 1,2,3,4-Tetrahydro- $\beta$ -carbolines

Several groups<sup>13,46,52,54,64</sup> had earlier investigated the ratio of *cis/trans* isomers in the Pictet–Spengler

## Scheme 7



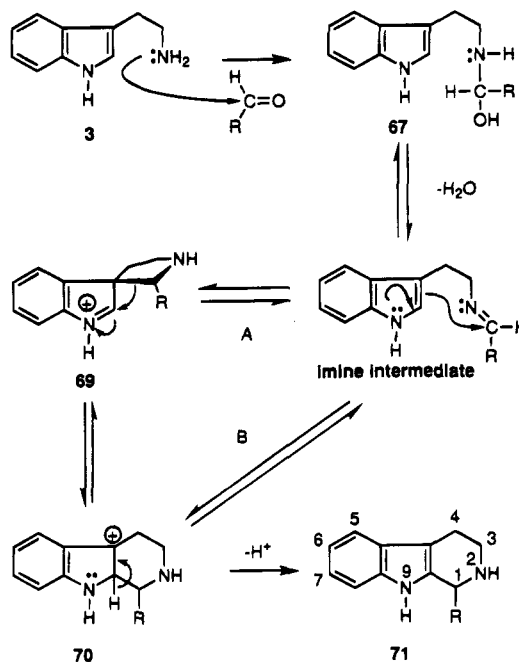
reaction. In all of the reactions examined previously, mixtures of *cis* and *trans* isomers had been reported. During the preparation of 1,3-disubstituted  $\beta$ -carbolines for <sup>13</sup>C NMR studies<sup>77</sup> Ungemach had discovered that the reaction of *N*<sub>b</sub>-benzyltryptophan methyl ester (**21**) with salicylaldehyde (**43**) in toluene at reflux provided a single diastereomer, the Pictet–Spengler product **47** in 97% yield (Table 3). Examination of the <sup>13</sup>C NMR spectrum confirmed that only one diastereomer had been formed in this process; however, steric interactions in this 1,2,3-trisubstituted  $\beta$ -carboline were too complex to permit an unequivocal assignment using <sup>13</sup>C NMR spectroscopy.<sup>77</sup>

Removal of the *N*<sub>b</sub>-benzyl function of this indole, however, permitted comparison of the properties of this base with those of authentic *cis*- or *trans*-1,3-disubstituted  $\beta$ -carbolines previously prepared by an independent route.<sup>54</sup> In this vein, the 2-benzyl derivative was subjected to catalytic hydrogenation (10% Pd/C, 25 psi) which resulted in a 75% yield of *trans*-tetrahydro- $\beta$ -carboline **46a** (Scheme 7). Earlier it had been reported<sup>46,54</sup> that treatment of tryptophan methyl ester with salicylaldehyde under the same conditions resulted in a mixture of *cis* and *trans* isomers (**46b,a**). The *N*<sub>b</sub>-benzyl group clearly directed this condensation in a *trans* stereospecific manner.<sup>60,76</sup> This represented the first completely stereoselective result in the *N*<sub>a</sub>-H tetrahydro- $\beta$ -carboline methyl ester series. The possibility that hydrogen bonding (hydroxyl group of the salicyl moiety **43**) might play a role in the stereoselectivity of this reaction led to the use of a second aldehyde, devoid of the hydroxyl group.

Cyclohexanecarboxaldehyde (**23**) was heated with *N*<sub>b</sub>-benzyltryptophan methyl ester (**21**) in benzene at reflux to provide the tetrahydro- $\beta$ -carboline (**27**) in 87% yield (Scheme 7). Again, examination of the <sup>13</sup>C NMR spectrum of this base indicated the presence of only one isomer in the reaction mixture, therefore, it was clear that the hydroxyl group played no role in directing the stereochemical outcome of the condensation. Catalytic debenzylation of **27** gave a 70% yield of *trans*-1-cyclohexyl-3-(methoxycarbonyl)-1,2,3,4-tetrahydro- $\beta$ -carboline (**25b**). As in the previous example this reaction carried out in the absence of the *N*<sub>b</sub>-benzyl function resulted in the formation of *cis* and *trans* isomers.<sup>54</sup> The *N*<sub>b</sub>-benzyl sequence had again occurred with 100% *trans* diastereoselectivity.

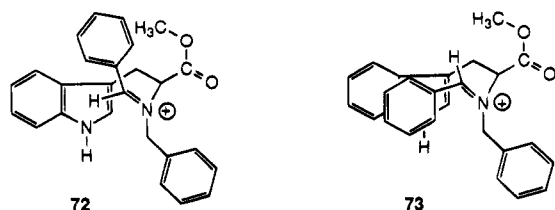
Previous studies had shown that A<sup>(1,2)</sup> strain between position 1 of the tetrahydro- $\beta$ -carboline and the *N*<sub>a</sub>-substituent was the dominant factor in determining the *cis/trans* ratio in the Pictet–Spengler reaction in the *N*<sub>b</sub>-H series.<sup>67,77</sup> With this in mind it was decided to condense smaller aldehydes with amine **21** and examine the *cis/trans* ratio. Glyoxal diethyl

## Scheme 8



acetal (**32**) was condensed with methyl ester **21** in refluxing benzene to furnish the desired tetrahydro- $\beta$ -carboline **39** in 75% yield (Scheme 7). Catalytic debenzylation of the base provided the 1,3-disubstituted  $\beta$ -carboline **38** in excellent yield. Careful examination of the <sup>13</sup>C NMR spectral data showed no evidence for the *cis* isomer. Even more important was the result when propionaldehyde (**68**) was treated with the hydrochloride salt of *N*<sub>b</sub>-benzyltryptophan methyl ester (**67**) in methanol/water at reflux (cold finger condenser) for 48 h. Three compounds were isolated from this reaction, one of which was the desired *trans*-*N*<sub>b</sub>-benzyl derivative **65** (46%) and the other two products appeared to arise from attack of the indole nitrogen atom on the aldehyde. Treatment of the base **65** with hydrogen over palladium on carbon yielded the *trans*-1-ethyl-3-(methoxycarbonyl)-1,2,3,4-tetrahydro- $\beta$ -carboline (**64b**) in 97% yield. These results clearly indicated the stereospecificity of this sequence was due largely to the effect of the *N*<sub>b</sub>-benzyl substituent and not solely the result of A<sup>(1,2)</sup> strain on the system.<sup>69</sup> This result was later confirmed when ester **21** was condensed with butyraldehyde in a sealed tube to provide the *trans* diastereomer with 77:23 (*trans/cis*) diastereoselectivity.<sup>61,62</sup>

An examination of the mechanism of this condensation with regard to diastereoselectivity is not a simple matter. The Pictet–Spengler reaction (see **67**–**71**) has generally been thought to proceed via a spiroindolenine intermediate<sup>5,56,78</sup> as shown in Scheme 8 (path A), although Casnati<sup>80</sup> has shown that



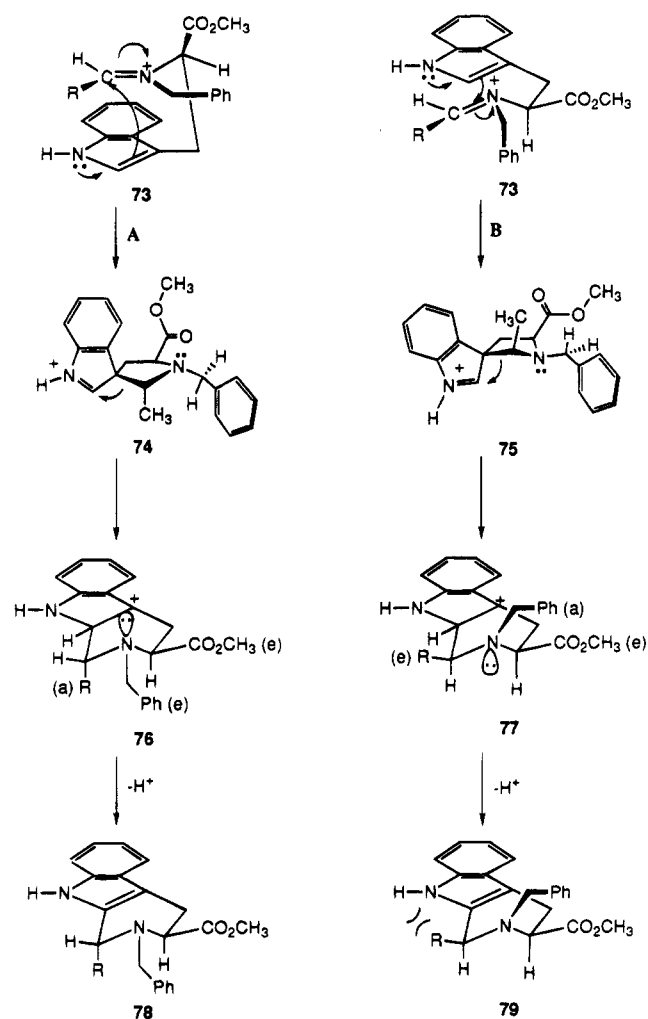
**Figure 3.** Steric interactions in the two possible iminium ion intermediates.

cyclization can occur by direct attack at position 2 of the indole (path B) when very reactive electrophiles are employed. The benzyl imine intermediate (when using *N*<sub>b</sub>-benzyltryptophan) can certainly be considered a reactive electrophile; therefore, both pathways must be considered when discussing the stereochemical outcome of this cyclization. Molecular models have been used to examine the steric interactions on the benzyl iminium ion intermediates **72** and **73** (Figure 3). These steric interactions which would develop in the transition state between the indole moiety and the phenyl group (or R group) in stereoisomer **72** are much higher in energy than the interactions between the indole group and the hydrogen atom of intermediate **73**. In addition, attack of the indole double bond of **72** (*Z* isomer) would result in a 1,3-interaction between the group at position -1 and the methyl ester in the transition state as the electronic character of the iminium ion approaches sp<sup>3</sup> hybridization. Conversely, this same unfavorable interaction would not develop in the *E* isomer **73**; therefore, stereoisomer **73** is believed to be the favored intermediate in this cyclization.

The second factor which introduces stereospecificity into this sequence is the stereoelectronic control in attack of the indole double bond on the benzyl imine **73**. Attack can occur from either C-2, or C-3 (see Scheme 9), but is expected to occur antiperiplanar to the methyl ester, similar to the attack of hydroxide ion on imidate salts reported by Deslongchamps.<sup>81</sup>

The structures which result from attack of the indole double bond at C-3 on the benzyl imine **73** from the bottom face (A) and the top face (B) of the carbon-nitrogen double bond are illustrated in Scheme 9. Intermediate **75** is clearly more crowded due to the *syn* alignment of the substituents at C-1, -2, and -3 than the *anti* spiroindolenine intermediate depicted in conformer **74**. Attack of the indole 2,3-double bond on imine **73** is believed to occur from the face opposite the ester function which minimizes interaction with this group during cyclization. Examination of Scheme 9 shows the stereospecificity of this sequence is even more obvious when intermediates **76** and **77** are examined. Spiroindolenine intermediate **74** results from the intramolecular attack (in a stereoelectronic sense) on imine **73** from the bottom face of the iminium ion, anti to the ester function. The result of this attack is occupation of the pseudo-equatorial position with the benzyl group while the phenyl group at C-1 would be pseudoaxial. Rearrangement of this intermediate leads to carbocation **76** which contains the two equatorial groups accompanied by the 1-axial substituent. Comparatively, attack on imine **73** from the top face of the iminium ion double bond would result in the more crowded spiroindolenine intermediate **75** which con-

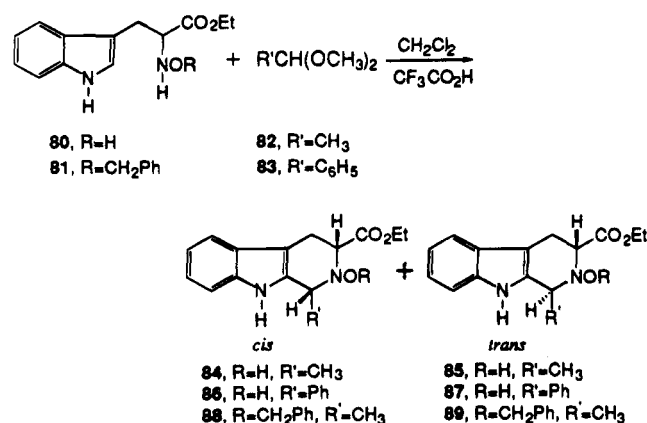
### Scheme 9



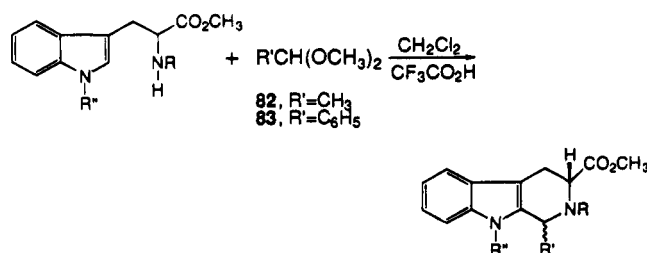
tains (after rearrangement) an axial benzyl function, an equatorial methoxycarbonyl group, and the equatorial 1-phenyl moiety. The *cis* diastereomer **77** is clearly the less stable diastereomer since the 2-benzyl function occupies an axial position in the transition state. In addition, the *cis* diastereomer **77** suffers from the additional unfavorable interaction between the equatorial substituent at position 1 and the indole *N*<sub>a</sub>-H function [A<sup>(1,2)</sup> strain]. It is therefore the combined influence of stereoelectronic control and conformational interactions which leads to complete stereospecificity in this Pictet-Spengler reaction to provide the *trans* isomer **78**.

Evidence suggested<sup>60</sup> that the *N*<sub>b</sub>-benzyl group when used in conjunction with large aldehydes resulted in the stereospecific formation of *trans*-1,3-disubstituted 1,2,3,4-tetrahydro- $\beta$ -carbolines. It was discovered, however, during the work of Ottenheim<sup>82</sup> on the synthesis of eudistomin alkaloids,<sup>83,84</sup> that the reaction of *N*<sub>b</sub>-(benzyloxy)tryptophan ethyl ester (**81**) with acetals **82** and **83** (Table 6) did not proceed with the same stereospecificity that had been earlier observed with *N*<sub>b</sub>-benzyltryptophan methyl ester (**21**).<sup>60</sup> Investigations in our laboratory have explored the reasons for variations in the stereospecificity of the Pictet-Spengler condensation. It was therefore of interest to determine if the oxygen substituent in the *N*<sub>b</sub>-hydroxyl- and *N*<sub>b</sub>-(benzyloxy)tryptophan ethyl esters (**80** and **81**, respectively) was responsible for



**Table 6. Condensation of *N*<sub>b</sub>-Oxytryptophan Ethyl Esters with Acetals**

reactants	R	R'	products		time (h)	yield (%)
			<i>cis</i>	<i>trans</i>		
80 + 82	H	CH <sub>3</sub>	84 (67)	85 (33)	72	95
80 + 83	H	C <sub>6</sub> H <sub>5</sub>	86 (40)	87 (60)	6	77
81 + 82	CH <sub>2</sub> Ph	CH <sub>3</sub>	88 (50)	89 (50)	3	96

**Table 7. Condensation of *N*<sub>b</sub>-Alkyltryptophan Methyl Esters with Acetals**

ester	R''	R	R'	product ratios <sup>a</sup>		time (h)
				<i>cis</i> (%)	<i>trans</i> (%)	
17	H	H	CH <sub>3</sub>	93 (75)	94 (25)	48
90	H	CH <sub>3</sub>	CH <sub>3</sub>	95 (66)	96 (34)	24
21	H	CH <sub>2</sub> Ph	CH <sub>3</sub>	97 (16)	98 (84)	72
91	H	CH <sub>2</sub> CH <sub>2</sub> Ph	CH <sub>3</sub>	99 (16)	100 (84)	168
21	H	CH <sub>2</sub> Ph	C <sub>6</sub> H <sub>5</sub>	101 (0)	102 (100)	48
92	CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>	103 (16)	104 (84)	288

<sup>a</sup> The stereochemistry and ratios of the *cis* and *trans* diastereomers were determined by <sup>1</sup>H and <sup>13</sup>C NMR spectroscopy.

the decreased stereospecificity when compared to the *N*<sub>b</sub>-methyl- and *N*<sub>b</sub>-phenethyltryptophan methyl esters (**90** and **91**, respectively), for a comparison see Tables 6 and 7.

The reactions which had been carried out in refluxing benzene were carried out under the conditions of Ottenheijm<sup>82</sup> in dichloromethane/trifluoroacetic acid (CH<sub>2</sub>Cl<sub>2</sub>/TFA). The *N*<sub>b</sub>-methyl- and *N*<sub>b</sub>-phenethyltryptophan methyl esters **90** and **91** were prepared to be employed as the carbon analogs of the corresponding *N*<sub>b</sub>-hydroxyl- and *N*<sub>b</sub>-(benzyloxy)tryptophan derivatives **80** and **81**. These derivatives permitted a comparison between the effect of size vs electronegativity (O vs C) on the stereoselectivity of the Pictet–Spengler condensation.

The reaction of *N*<sub>b</sub>-hydroxytryptophan **80** with acetaldehyde dimethyl acetal (**82**, Table 6) resulted in substantial amounts of the *cis* diastereomer, (67:33, *cis:trans*) as reported.<sup>82</sup> This same diastereose-

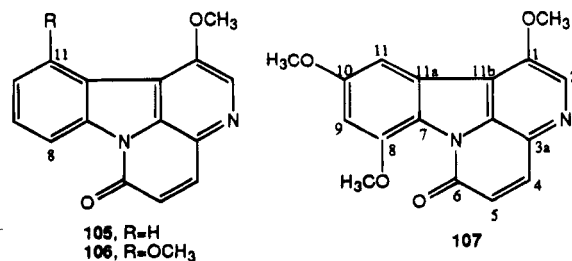
lectivity was realized when *N*<sub>b</sub>-methyltryptophan methyl ester (**90**) was condensed with acetal **82**. As expected, the size of the substituent on the *N*<sub>b</sub>-nitrogen atom led to higher *trans* diastereoselectivity in the reaction as illustrated in Tables 6 and 7. The reaction of *N*<sub>b</sub>-benzyloxytryptophan methyl ester (**81**) with **82** gave a 50:50 ratio of *cis:trans* diastereomers, while the corresponding reaction of the *N*<sub>b</sub>-phenethyl base **91** led to a high degree of *trans* selectivity (16:84). Comparison of the data in Tables 6 and 7 clearly shows that the electronic contributions from the oxygen atom decrease the stereoselectivity for the isosteric tryptophan methyl esters.<sup>85</sup>

The studies described in the preceding section have helped to shed some light on the factors which affect the diastereoselectivity of this widely used reaction. Moreover, a short two-step sequence has been developed to prepare *trans*-1,3-disubstituted 1,2,3,4-tetrahydro- $\beta$ -carbolines in 100% stereoselective fashion. Easy removal of the 2-benzyl function employed to direct the *trans* diastereoselectivity via catalytic debenylation (CTH) renders this method useful for the diastereoselective synthesis of many different 1,3-disubstituted tetrahydro- $\beta$ -carbolines and indole alkaloids of complex structure.

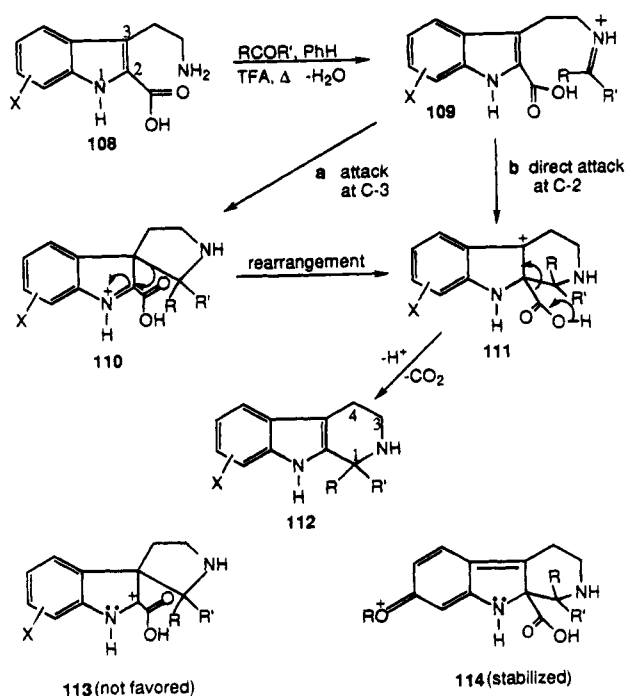
### C. Carboxyl-Mediated Pictet–Spengler Condensation

In the last several years an increasing number of  $\beta$ -carboline alkaloids that contain an oxygen substituent at position 4 have been isolated.<sup>86–90</sup> The 4-methoxy- $\beta$ -carbolines<sup>86–88</sup> and canthin-6-ones,<sup>86–89</sup> as well as several bisindoles<sup>91</sup> serve as representative examples. The alkaloids 1-methoxycanthin-6-one (**105**), and 1,11-dimethoxycanthin-6-one (**106**) and their congeners shown in Figure 4 have been reported to exhibit cytotoxic antileukemic activity via their inhibitory effects on DNA synthesis in GPK epithelial cells.<sup>92,93</sup> Oxygenation of the canthin-6-one skeleton either at position 1 (C-4 in the  $\beta$ -carboline numbering system) and/or ring A greatly enhanced the cytotoxic antileukemic activity of these bases.<sup>92,93</sup>

The synthesis of the parent 1-methoxycanthin-6-one (**105**) has been carried out and is described below. Efforts are currently focused on the synthesis of “unnatural products” such as 1,8,10-trimethoxycanthin-6-one (**107**) to investigate the mode of action of the cytotoxic activity of these unique oxygenated alkaloids. The approach to **106** or **107** requires a simple route to oxy-substituted tryptamines, the most straightforward of which was reported earlier by Abramovitch and Shapiro.<sup>94,95</sup> This process suffered, however, because the decarboxylation of the tryptamine-2-carboxylic acid to provide the substituted tryptamine often occurred in low yield, depending on

**Figure 4.** Methoxy-substituted canthin-6-ones.

## Scheme 10



the nature of the substituents on the indole ring.<sup>94,95</sup> It was this synthetic problem which led to the development of the carboxyl-mediated Pictet-Spengler reaction.<sup>96,97</sup> With this in mind, the mechanism of the Pictet-Spengler reaction<sup>5,56,80</sup> was reviewed. As outlined in Scheme 10, if the tryptamine-2-carboxylic acid (**108**) could be encouraged to form the Schiff's base **109**, and was then heated, this might provide the spiroindolenine intermediate **110** (C-3) or the carbocation **111** [path b (C-2) or from **110**]. Loss of both the proton and the elements of CO<sub>2</sub> from **111** to satisfy the positive charge would provide the desired 1,2,3,4-tetrahydro- $\beta$ -carboline **112**.

When a substituted tryptamine-2-carboxylic acid (**108**) was simply heated with the carbonyl compound in a solution of benzene/dioxane/trifluoroacetic acid at reflux with water removal (Dean-Stark trap) the desired tetrahydro- $\beta$ -carboline was produced. The results of the condensation between various tryptamine-2-carboxylic acids and a carbonyl component are summarized in Table 8. The process appears to be quite general for simple aldehydes such as benzaldehyde **18** as well as more reactive electrophiles including  $\alpha$ -keto acids and  $\alpha$ -keto esters undergo the cyclization with ease. The carboxyl-mediated Pictet-Spengler cyclization employed herein represented a considerable improvement over the reported syntheses of indoles **125**–**133**, and can be extended to include many ring A-oxygen substituted tryptamine-2-acids which fail to undergo decarboxylation during attempts to convert the tryptamine-2-carboxylic acids to tryptamines.<sup>94,95,97</sup> This is also important for acids **119** and **120**, the decarboxylation of which is very difficult under normal reaction conditions.<sup>97,98</sup>

Mechanistically, the iminium ion **109** can undergo attack at C-3 to provide the spiroindolenine intermediate<sup>5,56</sup> **110** or the ion **109** can undergo direct attack at C-2<sup>80</sup> to provide the carbocation **111**. The former intermediate is unlikely due to the localization of positive charge adjacent to the carbonyl group (see

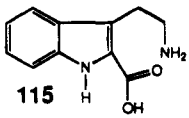
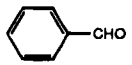
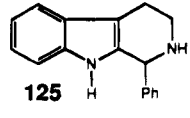
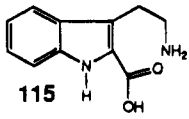
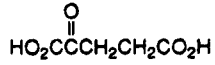
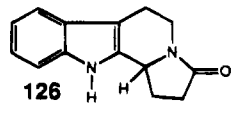
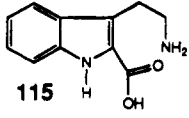
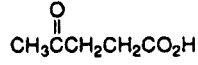
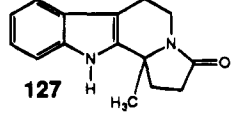
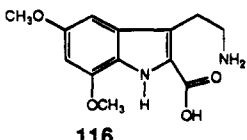
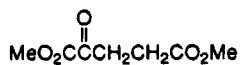
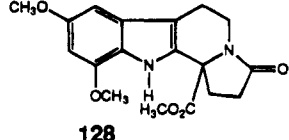
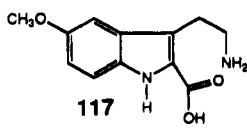
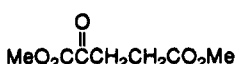
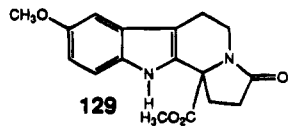
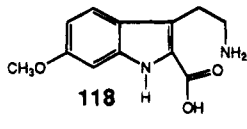
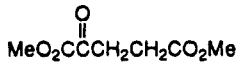
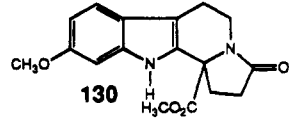
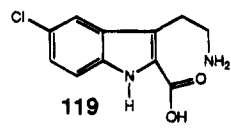
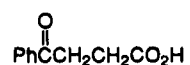
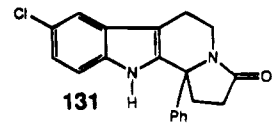
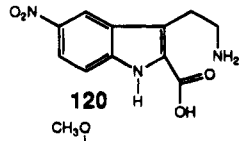
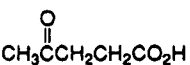
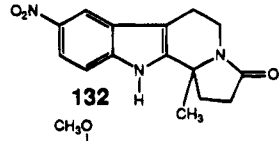
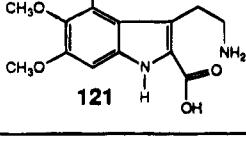
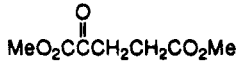
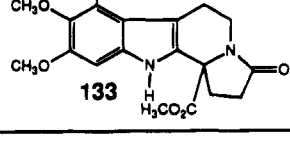
resonance structure **113**, Scheme 10). It is known that 6-alkoxy substituents facilitate attack at C-2<sup>80</sup> in the Pictet-Spengler reaction. If this were the case then resonance structure **114** might be expected to play a role in the stabilization of intermediate **111** in the condensation reaction described herein. From Table 8 it is clear that the treatment of 6-methoxytryptamine-2-carboxylic acid (**118**) with  $\alpha$ -keto ester **123** provided a higher yield of tetrahydro- $\beta$ -carboline than the 5-methoxy analog **117** in agreement with direct attack at C-2 for this process.

The synthesis of the cytotoxic antileukemic alkaloid 1-methoxycanthin-6-one **105** was completed by Hagen et al.<sup>99</sup> and is illustrated in Scheme 11. This synthetic study also resulted in a general method for the synthesis of 4-alkoxy- $\beta$ -carbolines from 4-oxo-1,2,3,4-tetrahydro- $\beta$ -carbolines. The treatment of tryptamine hydrochloride **3** with the dimethyl ester of 2-ketoglutaric acid (**123**) in methanol at reflux provided the desired indolizino[8,7-*b*]indole lactam **134** in 92% yield; this same condensation occurs readily between **115** and **123** as well. During this process a Pictet-Spengler cyclization had occurred and the  $\gamma$ -lactam **134** had formed in a one-pot reaction. The  $\gamma$ -lactam contained the necessary carbon atoms for the synthesis of **105**; moreover, both the N<sub>b</sub>-nitrogen atom and C-1 carbon atom were protected from interaction with 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ). The desired 3-acylindole **135** was obtained in good yield when the  $\gamma$ -lactam **134** was stirred with DDQ (1:2) in aqueous THF at room temperature. The 4-oxo-1,2,3,4-tetrahydro- $\beta$ -carboline **135** was heated in HCl/HOAc, according to the procedure of Hobson,<sup>75</sup> to remove the ester protecting group which resulted in the formation of 3-acylindole **136** in 88% yield. The  $\gamma$ -lactam of **136** proved to be resistant to hydrolysis under a variety of conditions.<sup>99,100</sup>

In order to facilitate cleavage of the  $\gamma$ -lactam to provide the  $\delta$ -lactam, it was decided to form the enol ether of the 3-acyl indole **136**. This would provide a 1,2-dihydro- $\beta$ -carboline, congeners of which are known to readily undergo oxidation (O<sub>2</sub>, air) or disproportionation to provide the fully aromatic  $\beta$ -carbolines.<sup>54,101</sup> With this in mind, the keto lactam **136** was heated with trimethyl orthoformate in methanol in the presence of *p*TSA to provide a 51% yield of 4-methoxy-1-(3-carbomethoxypropyl)- $\beta$ -carboline **137**. The ester group of **137** was hydrolyzed in quantitative yield with aqueous ammonia, followed by heating of the residual solid in the presence of *p*TSA to furnish 1-methoxy-4,5-dihydrocanthine-6-one (**138**) in 82% yield. Dehydrogenation of the  $\delta$ -lactam **138** was accomplished using DDQ in dioxane at reflux to provide 1-methoxycanthin-6-one (**105**) in 70% yield. This seven-step synthesis of **105** proceeded in an overall yield of 20% starting from tryptamine **3** and dimethyl 2-ketoglutarate (**123**). This represented the first synthesis of any of the 1-methoxycanthin-6-one alkaloids and provides a route to other alkaloids in this series.

The carboxyl-mediated Pictet-Spengler reaction is an effective method for the direct synthesis of 1,2,3,4-tetrahydro- $\beta$ -carbolines and hexahydro-3-oxo-indolizino[8,7-*b*]indoles from tryptamine-2-carboxylic acids. It is no longer necessary to remove the 2-carboxylic acid function to provide tryptamines prior to the

Table 8. Synthesis of Indolizino[8,7-*b*]indoles from Tryptamine-2-carboxylic Acids

tryptamine-2-acid	carbonyl component	product	(% yield)
			(75)
			(58)
			(85)
			(65)
			(60)
			(80)
			(60)
			(52)
			(66)

execution of the Pictet–Spengler reaction for the elements of CO<sub>2</sub> are lost during the process of cyclization. This greatly enhances the use of the Abramovitch–Shapiro method<sup>94,95</sup> for the synthesis of substituted- $\beta$ -carbolines especially in the area of highly oxygenated ring A-substituted heterocycles. This method can be potentially extended to other indoles which contain a carboxyl function located at position-2. Heterocycles such as indole-2-esters are readily available through the Fischer,<sup>102</sup> Reissert,<sup>103</sup> and Moody<sup>104,105</sup> routes.

#### D. Kinetic vs Thermodynamic Control

Previous reports highlighted the variance in the diastereochemical product ratios obtained when employing the Pictet–Spengler condensation.<sup>54,63,77</sup> Three such examples are shown in Table 9. The ratios of

diastereomers when acetaldehyde (**157**), propionaldehyde (**68**), and cyclohexanecarboxaldehyde (**23**) were heated with *N*<sub>5</sub>-H tryptophan methyl ester under acidic conditions reportedly varied from 75:25 *cis/trans* (**139/140**) to a more equal ratio of 43:57 *cis/trans* (**64a/64b**), to 41:59 *cis/trans* (**25a/25b**). The only significant difference between these reactions was the size of the substituent at C-1 of the tetrahydro- $\beta$ -carboline. We have noted previously that as the size of the substituent at C-1 increases, so too does the degree of *trans* stereoselectivity.<sup>54,77</sup> To examine the effect the steric bulk of the incoming aldehyde had upon the diastereomeric ratio realized in the Pictet–Spengler condensation in the presence of a benzyl group at the amino nitrogen, *N*<sub>5</sub>-benzyltryptophan methyl ester was prepared.<sup>52,106</sup> This base was treated individually with acetaldehyde,

## Scheme 11

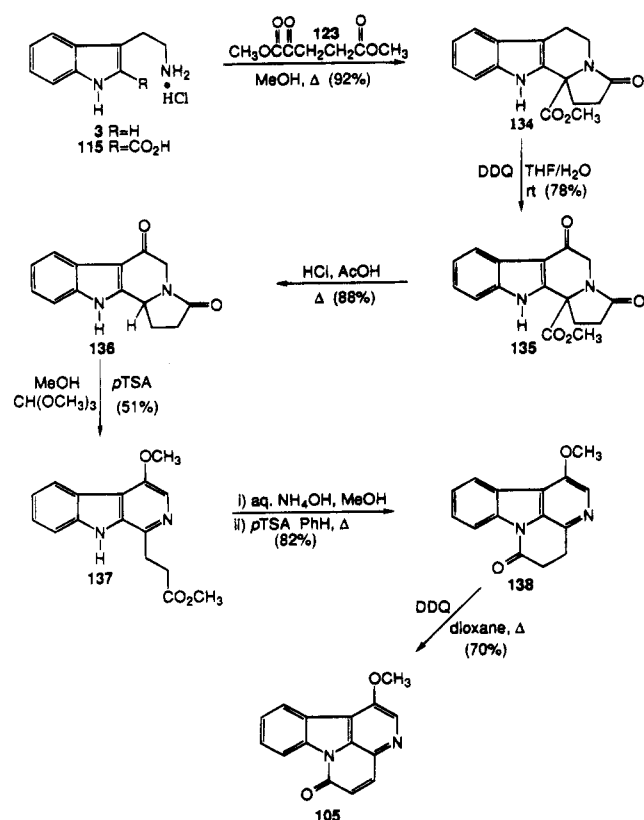


Table 9. *Cis:Trans* Ratios<sup>a</sup> of ( $\pm$ )-1,2-Disubstituted and 1,2,3-Trisubstituted Tetrahydro- $\beta$ -carbolines

compound	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	aprotic <i>cis:trans</i>	acidic <i>cis:trans</i>
139, 140 <sup>b</sup>	CH <sub>3</sub>	H	CH <sub>3</sub>		75:25
64a, 64b	CH <sub>2</sub> CH <sub>3</sub>	H	CH <sub>3</sub>		43:57 <sup>c</sup>
25a, 25b	C <sub>6</sub> H <sub>11</sub>	H	CH <sub>3</sub>	40:60 <sup>c</sup>	41:59 <sup>c</sup>
141, 142	CH <sub>3</sub>	Bn	CH <sub>3</sub>	26:74	12:88
143, 144 <sup>d</sup>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>	Bn	CH <sub>3</sub>	23:77	11:89
27	C <sub>6</sub> H <sub>11</sub>	Bn	CH <sub>3</sub>	0:100	0:100
145, 146	CH <sub>3</sub>	Bn	CH(CH <sub>3</sub> ) <sub>2</sub>	23:77	13:87
147, 148	CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>	Bn	CH(CH <sub>3</sub> ) <sub>2</sub>	13:87	12:88
149	C <sub>6</sub> H <sub>11</sub>	Bn	CH(CH <sub>3</sub> ) <sub>2</sub>	0:100	0:100
150, 151	CH <sub>3</sub>	CH(Ph) <sub>2</sub>	CH <sub>3</sub>	10:90	0:100
152	CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>	CH(Ph) <sub>2</sub>	CH <sub>3</sub>	0:100	0:100
153	C <sub>6</sub> H <sub>11</sub>	CH(Ph) <sub>2</sub>	CH <sub>3</sub>	0:0 <sup>e</sup>	0:100
154	CH <sub>3</sub>	CH(Ph) <sub>2</sub>	CH(CH <sub>3</sub> ) <sub>2</sub>	0:100	0:100
155	CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>	CH(Ph) <sub>2</sub>	CH(CH <sub>3</sub> ) <sub>2</sub>	0:0 <sup>e</sup>	0:100
156	C <sub>6</sub> H <sub>11</sub>	CH(Ph) <sub>2</sub>	CH(CH <sub>3</sub> ) <sub>2</sub>	0:0 <sup>e</sup>	0:100

<sup>a</sup> As determined by integration of the <sup>1</sup>H NMR spectrum ( $\pm 3\%$ ). <sup>b</sup> When a pair of numbers is present, the first is *cis*, the second *trans*. When only one number is present the reaction was 100% *trans* stereoselective. <sup>c</sup> Reference 77. <sup>d</sup> Carried out in the optically active series. <sup>e</sup> These transformations yielded only starting materials under nonacidic aprotic conditions but were converted into the 1-substituted tetrahydro- $\beta$ -carbolines upon addition of TFA to the reaction medium.

butyraldehyde, and cyclohexanecarboxyaldehyde under the nonacidic aprotic conditions of Sandrin et al.<sup>46,47,85</sup> This afforded a series of 1-alkyl-2-benzyl-3-(methoxycarbonyl)tetrahydro- $\beta$ -carbolines with the substituent at position 1 varying in steric bulk from

the relatively small methyl group, to propyl, to the larger cyclohexyl moiety.

The reactants were heated in benzene at reflux under nonacidic aprotic conditions for 36 h. All reactions were stopped after 36 h to provide uniform experimental conditions, thus enabling an unbiased comparison of the stereochemical effects of the various substituents. Chemical yields are thus unoptimized. The reactions utilizing acetaldehyde were carried out in a sealed glass tube to avoid loss of the aldehyde. Each aldehyde was purified by vacuum distillation and the IR spectrum obtained to verify that the reactants were indeed free of acid. In all cases, after 36 h a small aliquot was removed and the proton NMR spectrum recorded. It was difficult to accurately measure the diastereomeric ratio due to impurities which masked the signals necessary to determine the ratio of diastereomers which were formed. In these cases impurities were removed by running the reaction mixture through a short wash column of silica gel and measuring the NMR spectrum once again. Comparison of the spectra before and after chromatography assured that the diastereomeric ratio had not been altered. Flash chromatography on silica gel permitted separation of the *cis* and *trans* diastereomers and a close examination of the NMR spectra of the individual diastereomers clearly indicated the proton signals that differed significantly. These signals were then used to determine the diastereomeric ratio of *cis* to *trans* isomers in the spectra of the mixtures. Although CDCl<sub>3</sub> was initially used as an NMR solvent it was found that the acidic impurities present were in high enough concentration to effect epimerization of the *cis* diastereomers to their *trans* isomers. The cause of this epimerization will be expanded upon in a later section. These impurities could be removed but it was found easier to employ C<sub>6</sub>D<sub>6</sub> as the NMR solvent. Catalytic transfer hydrogenation of the 1,2,3-trisubstituted species afforded 1,3-disubstituted tetrahydro- $\beta$ -carbolines that are well-known compounds.<sup>54,77</sup> Consequently, the stereochemistry of the individual diastereomers could be unequivocally established by comparison of their <sup>13</sup>C NMR spectra to previously published work. Determination of the stereochemistry of these individual diastereomers had been carried out according to the carbon NMR method described earlier, a method which has been validated by independent sources.<sup>76-79</sup>

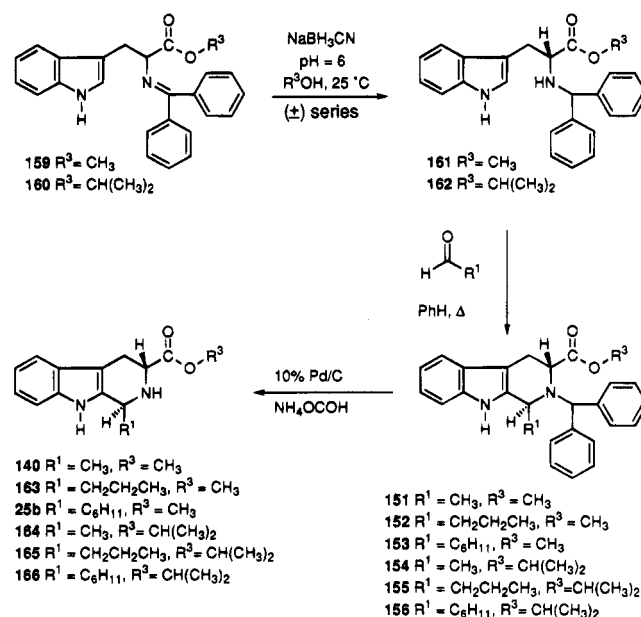
The condensation of *N*<sub>b</sub>-benzyltryptophan methyl ester (21) with acetaldehyde (157) or butyraldehyde (158) resulted in the formation of mixtures of *cis* and *trans* diastereomers 141–144, respectively. This same condensation employing cyclohexanecarboxaldehyde 23 provided only the *trans* diastereomer 27 as was expected.<sup>60</sup> Examination of the product ratios (Table 9) for the nonacidic aprotic condensations with acetaldehyde (26:74 *cis*-141/*trans*-142) and butyraldehyde (23:77 *cis*-143/*trans*-144) revealed that the ratio of *cis* to *trans* diastereomers for each was essentially the same. As the steric bulk of the substituents increased to cyclohexyl, however, the *cis* isomer was excluded from formation. Ungemach had proposed that this phenomenon was the result of a steric interaction between the ester function and the alkyl group in addition to the effect of A<sup>(1,2)</sup> strain.<sup>60</sup>

Since aldehydes similar in size to an *n*-butyl group provided a mixture of diastereomers the synthetic potential of the process was limited. If Ungemach's hypothesis were correct, however, any increase in the steric interactions between the ester and alkyl groups should shift the diastereomeric ratios toward increased *trans* stereoselectivity. To this end, replacement of the methyl ester function with the larger isopropyl group was undertaken. After the *N*<sub>b</sub>-benzyltryptophan isopropyl ester was synthesized in a fashion analogous to the methyl ester, the previously mentioned series of condensations were repeated and the diastereomeric ratios examined by NMR spectroscopy. In agreement with this hypothesis one of the condensations yielded an increased amount of *trans* isomer, the condensation of *N*<sub>b</sub>-benzyltryptophan isopropyl ester with butyraldehyde (158). In the 1-propyl methyl ester series (143 and 144) the ratio of *cis* to *trans* was 23:77 for this condensation while in the isopropyl ester series (147 and 148) the amount of *trans* isomer was increased to 13:87 (*cis:trans*).

The previous work in this area had clearly demonstrated that introduction of a benzyl group at the *N*<sub>b</sub>-amino nitrogen function favored formation of the *trans* diastereomer. Substituents at the amino *N*<sub>b</sub>-nitrogen function have come under scrutiny previously since it was shown by Ottenheim et al.<sup>82</sup> and Sandrin et al.<sup>85</sup> that electronic effects played a significant role in the stereochemical outcome of the reaction. The effect of the steric bulk of the *N*<sub>b</sub>-alkyl substituent upon the diastereochemical ratio had not been fully studied. In order to increase the size but not alter the electronic effects on the stereochemical preference of this condensation, the substituent at the *N*<sub>b</sub>-nitrogen function, a diphenylmethyl group, was employed. Introduction of this group (Scheme 12) was accomplished by transimination of the appropriate ester hydrobromide or hydrochloride salt with benzophenone imine, according to the procedure of Polt and O'Donnell.<sup>107</sup> Reduction of the resultant imine with sodium cyanoborohydride in methanol or 2-propanol under acidic conditions provided the desired *N*<sub>b</sub>-diphenylmethyl-substituted tryptophan derivative 161 or 162, respectively. This facile reduction could be carried out in as little as 5 min (depending upon the scale of the reaction) with complete conversion of the imine. This protocol provided excellent yields, could easily be scaled up to the 10 g level, and was not subject to disubstitution or the formation of quaternary ammonium salts. With the methyl and isopropyl esters of *N*<sub>b</sub>-diphenylmethyltryptophan in hand, the series of condensations were repeated (benzene at reflux) and the diastereomeric ratios measured by NMR spectroscopy.

In the methyl ester series the condensation with acetaldehyde reacted to form the desired 1,2,3,4-tetrahydro- $\beta$ -carboline (150 and 151) as evidenced by <sup>1</sup>H NMR spectroscopy (10:90) after initial removal of the nonindolic byproducts. However, attempts to isolate the small amount of *cis* diastereomer which was present were unsuccessful. Only the *trans* diastereomer 151 was found upon examination of the chromatographic fractions. In the reaction of *N*<sub>b</sub>-(diphenylmethyl)tryptophan isopropyl ester (162) with acetaldehyde, only the *trans* diastereomer 154

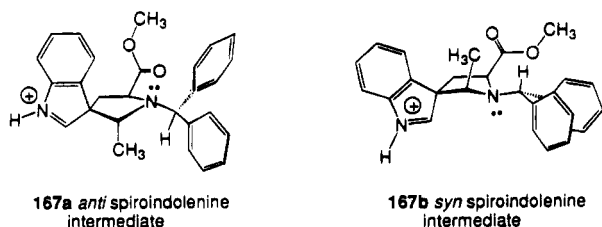
Scheme 12



was formed as determined by <sup>1</sup>H and <sup>13</sup>C NMR spectroscopy. In the methyl ester series the reaction of butyraldehyde also provided only the *trans* isomer 152. In three of the cases (153, 155, and 156) no reaction was observed under nonacidic aprotic conditions after 36 h presumably due to increased crowding in the transition states. Prolonged heating only increased the amount of byproducts which were formed. In order to effect the Pictet–Spengler cyclization it was necessary to add trifluoroacetic acid (TFA) to the reaction medium. The addition of TFA to catalyze the formation of 153, 155, and 156 resulted in the isolation of only the *trans* diastereomer in each case. Cleavage occasionally of diphenylmethyl groups is known to occur with TFA at low concentrations,<sup>108–110</sup> although it failed to do so here as evidenced by control reactions devoid of aldehyde. In all the reactions employed with the *N*<sub>b</sub>-diphenylmethyl group, the chemical yields are somewhat lower than in the *N*<sub>b</sub>-benzyl cases. This is not unreasonable since the presence of such bulky moieties hinders the formation of 1,3-disubstituted 1,2,3,4-tetrahydro- $\beta$ -carboline.<sup>82,108–110</sup>

A comparison of the reactions which employed acetaldehyde showed a marked trend. In the *N*<sub>b</sub>-H methyl ester case (139 and 140, 75:25 *cis/trans*), the Pictet–Spengler reaction yielded the *cis* isomer as the major product. Introduction of a benzyl function at the amino nitrogen atom inverted the stereochemical outcome to provide the *trans* diastereomer under aprotic conditions (141 and 142, 26:74 *cis/trans*) as the major product. Furthermore, when the *N*<sub>b</sub> substituent was increased to the larger diphenylmethyl group under similar conditions, a 9:1 stereoselective formation of the *trans* diastereomer 151 was observed (Table 9). When the methyl ester substituent was replaced with an isopropyl group a 100% stereoselective formation of the *trans* diastereomer 154 was realized.

As previously shown in Scheme 9, two distinct spiroindolenine intermediates may result from attack on the imine intermediate. In the *N*<sub>b</sub>-diphenylmeth-



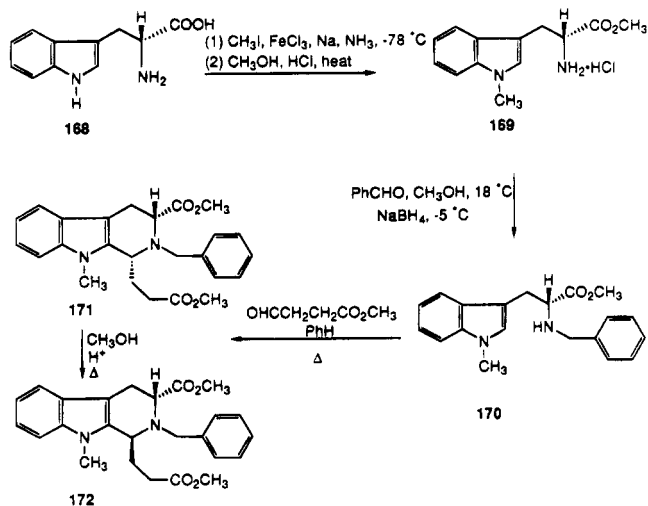
**Figure 5.** *Syn*- and *anti*-spiroindolenine intermediates. In the *anti*-spiroindolenine case, attack of the imine from the face opposite the ester function would provide the *anti* spiroindolenine **167a**. Attack from the same face as that occupied by the ester would result in a spiroindolenine of *syn* configuration such as intermediate **167b**, as illustrated in Figure 5. Molecular mechanics calculations combined with conformational searching (MacroModel 2.5–MM2 force field) revealed that the *anti* configuration is 2.1 kcal/mol lower in energy than the all eclipsed *syn* configuration.<sup>62</sup> On rearrangement to C-2 this *anti* spiroindolenine **167a** can only provide the *trans* diastereomer.

The data from these experiments under nonacidic aprotic conditions (benzene) reflect ratios which are the result of kinetic trapping experiments. Examination of the data in Table 9 clearly indicated that if mixtures of  $N_b$ -alkyl diastereomers realized in the nonacidic aprotic Pictet–Spengler reaction were exposed to acid, the diastereomeric ratios in the  $N_b$ -alkyl cases shifted to further increase the amount of *trans* diastereomer formed. This generality was confirmed by either the addition of TFA to a small aliquot of the reaction mixture or as in cases with tetrahydro- $\beta$ -carbolines **153**, **155**, and **156**, by reaction in the presence of TFA to catalyze cyclization. Examination of ratios **141/142**, **143/144**, **145/146**, and **150/151**, clearly indicated that the ratio of *cis* product to *trans* product formed under nonacidic aprotic conditions was higher than that realized under acidic conditions. The method by which the conversion of the *cis* diastereomers into the more thermodynamically stable *trans* diastereomers took place can now be rationalized.

### E. Epimerization at C-1 by Scission of the Carbon–Nitrogen Bond (Thermodynamic Control)

For these studies optically active  $N_a$ -methyl- $N_b$ -benzyl-D-(+)-tryptophan methyl ester was synthesized from D-(+)-tryptophan (**168**, Scheme 13). Alkylation of D-(+)-tryptophan with methyl iodide in sodium–liquid ammonia at  $-78^\circ\text{C}$  and esterification of the resultant solid by heating in methanolic hydrogen chloride at reflux provided ester **169**. This indole was converted into  $N_a$ -methyl- $N_b$ -benzyltryptophan methyl ester (**170**) on stirring the free base with benzaldehyde at  $18^\circ\text{C}$  for 2 h, followed by reduction of the imine with sodium borohydride at  $-5^\circ\text{C}$ . Racemization was observed if the benzylation process took too long or the temperature rose too high,<sup>111</sup> although in the racemic series (17 h) it made no difference. The optical purity of the above compounds was verified by  $^1\text{H}$  NMR spectroscopy in the presence of the chiral shift reagent tris[3-(trifluoromethyl)hydroxymethylene]-(+)-camphorato]europium(III) and later by HPLC analysis of diastereomeric ureas.<sup>24,106,112</sup>

### Scheme 13



**Table 10.**  $^1\text{H}$  NMR Data for **171** and **172**

proton	172 ( <i>trans</i> )		171 ( <i>cis</i> )	
	chemical shift $\delta$ (ppm)	$J$ (Hz), multiplicity	chemical shift $\delta$ (ppm)	$J$ (Hz), multiplicity
H (1)	3.51	15.7/5.2, dd	3.75	9.5, d
H (3)	4.10	11.0/5.5, dd	3.92	6.3/2.1, dd
H (4 <sub>a</sub> )	3.05	15.8/5.5, dd	3.05	18.6/6.3, dd
H (4 <sub>b</sub> )	3.12	15.8/11.0, dd	3.37	18.6/2.1, dd
H (5)	7.60	8.0, d	7.58	9.5, d
CH <sub>3</sub> ( $N_b$ )	3.84	s	3.69	s
CH <sub>3</sub> (ester)	3.65	s	3.65	s
CH <sub>3</sub> (ester)	3.50	s	3.56	s
H <sub>8</sub> (aromatic)	7.12–7.40	m	7.10–7.48	m
H (14 <sub>a</sub> )	3.39	13.1, d	3.89	s (14 <sub>a</sub> /14 <sub>b</sub> )
H (14 <sub>b</sub> )	3.81	13.1, d		
H (1' <sub>a</sub> )	1.85	m	1.50	m
H (1' <sub>b</sub> )	2.02	m	1.95	m
H (2' <sub>a</sub> )	2.38	17.5/5.6, dt	2.51	18.0/6.0, dt
H (2' <sub>b</sub> )	2.61	17.5,9.6/5.6, d, dd	2.81	18.0,9.8/6.0, d, dd

Reaction of optically active  $N_a$ -methyl- $N_b$ -benzyltryptophan methyl ester **170** with methyl 3-formylpropionate in benzene at reflux provided a mixture of the optically active diastereomers **171** [ $[\alpha]_D^{25} = -36.8^\circ$  ( $c = 0.95$ ,  $\text{CHCl}_3$ ), and **172** [ $[\alpha]_D^{25} = +20.0^\circ$  ( $c = 0.96$ ,  $\text{CHCl}_3$ ) in a *trans/cis* ratio of 72:28 (81%). The *trans* isomer **172** (mp  $119$ – $120^\circ\text{C}$ ) was identical spectrometrically to the ( $\pm$ )-*trans* isomer (mp  $145$ – $146^\circ\text{C}$ ) whose structure had been confirmed by Yoneda by X-ray crystallographic analysis.<sup>113</sup> It was interesting to note that both the C-1 substituent and the  $N_b$ -benzyl function of the *trans* isomer occurred in the axial position in the solid state as confirmed by the X-ray analysis.<sup>113</sup> The coupling constants observed between H-3 and H-4<sub>a</sub>, H-4<sub>b</sub> ( $J = 5.5$  and  $11.0$  Hz) indicate that H-3 was located on the  $\beta$ -axial position of the chair-like C-ring with a dihedral angle of  $135^\circ$  with respect to H-4<sub>a</sub> and  $10^\circ$  in regard to H-4<sub>b</sub> (Table 10). Hence the *trans* isomer **172** contained an axial substituent at C-1, an equatorial substituent at C-3 and an axial benzyl group in the preferred



Table 11.  $^{13}\text{C}$  NMR Data for **171** and **172**

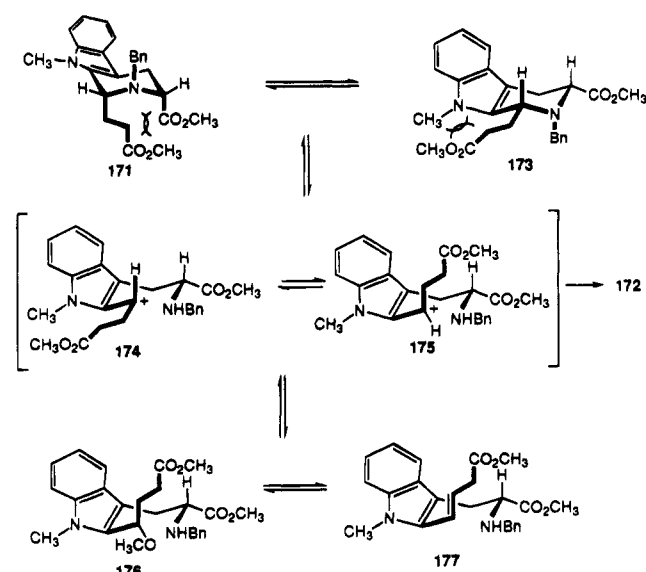
carbon atom	<b>172</b>	<b>171</b>	racemic <sup>a</sup>
C (1)	53.32	54.09	53.5
C (3)	56.12	57.44	56.2
C (4)	20.25	17.95	20.4
C (5)	118.12	118.25	118.2
C (6)	119.11	118.93	119.2
C (7)	121.32	121.28	121.4
C (8)	108.90	108.79	108.9
C (10)	135.67	134.65	135.7
C (11)	106.29	104.70	106.4
C (12)	126.52	126.58	126.7
C (13)	137.46	137.49	137.6
C (14)	52.79	61.16	52.9
C (14a)	139.25	138.93	139.2
C (14b)	129.29	129.07	129.3
C (14c)	128.14	128.36	128.2
C (14d)	126.96	127.31	127.0
C (1')	27.90	29.10	28.0
C (2')	29.60	29.66	29.68
N-CH <sub>3</sub>	29.71	29.66	29.74
COOCH <sub>3</sub>	51.26	51.30	51.3
COOCH <sub>3</sub>	52.00	51.83	52.0
C=O	173.37	174.07	173.3
C=O	173.87	174.25	173.9

<sup>a</sup> Reference 118.

conformation in solution. This was further confirmed by  $^{13}\text{C}$  NMR spectroscopy. In the  $^{13}\text{C}$  NMR spectrum of the two diastereomeric 1,2,3,4-tetrahydro- $\beta$ -carbolines, it was well documented that the exocyclic benzylic methylene carbon atom (C-14) of the  $N_b$ -benzyl-*cis*-1,3-disubstituted-1,2,3,4-tetrahydro- $\beta$ -carboline resonated downfield from that of the corresponding *trans* isomer.<sup>68,114</sup> As illustrated in Table 11, the chemical shift of carbon atom 14 in the *cis* isomer **171** ( $\delta$  61.16) appeared 8.37 ppm downfield with respect to the corresponding carbon atom in the *trans* diastereomer **172** ( $\delta$  52.79). This phenomenon was due to the  $\gamma$ -gauche effect.<sup>68,77,114</sup> The preferred position of the  $N_b$ -benzyl group in the *cis* isomer **171** must be *trans* to the two 1,3-substituents, while in the corresponding *trans* isomer, any position occupied by the benzyl group will necessarily be *cis* either to the substituent at C-1 or C-3. Consequently, this carbon atom (C-14) is compressed in the  $^{13}\text{C}$  NMR spectrum of *trans* isomer **172**, and resonates up field with respect to the corresponding carbon atom in the *cis* isomer **171**. The coupling constants between H-3 and H-4a, H-4b ( $J = 6.3$  and  $2.1$  Hz) indicate that hydrogen atom H-3 of the *cis* isomer **171** was located in the  $\alpha$ , equatorial position with a dihedral angle of  $35^\circ$  with respect to H-4a and  $85^\circ$  with regard to H-4b (Table 10). The benzyl group in the *cis* diastereomer should be in the  $\beta$ , axial position in order to reduce the steric repulsion with the *cis* 1,3-substituents. This was confirmed by the  $^{13}\text{C}$  NMR spectrum as described previously on the  $N_b$ -H analogs.<sup>5</sup> The acids corresponding to the *trans* **172** and *cis* **171** diastereomers were also obtained (58:42) in optically active form when the Pictet–Spengler reaction of **170** was executed with  $\alpha$ -ketoglutaric acid (84%) in benzene-dioxane.<sup>111</sup> The mechanism of diastereoselectivity in the Pictet–Spengler condensation of  $\alpha$ -keto acids is different from the mechanism which has been discussed thus far.<sup>1,111</sup>

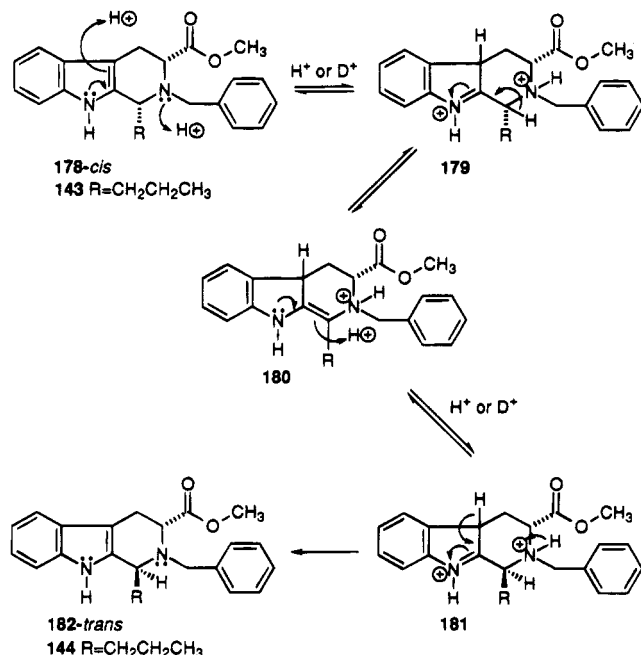
There were a number of reports in the literature regarding the synthesis of analogs of the *cis* isomer

Scheme 14



**171**.<sup>45,113,115–120</sup> However, little was known about the effect of the substituents located 1,3 on the tetrahydro- $\beta$ -carboline ring with regard to inversion of configuration in these molecules. It was observed that the *cis* isomer (+)-**171**, could be converted into the thermodynamically more stable *trans* diastereomer (–)-**172** (85%),  $[\alpha]_D^{25} = -36.6^\circ$  ( $c = 1.0$ ,  $\text{CH}_3\text{Cl}$ ), by heating *cis* isomer **171** in a 1% anhydrous HCl–methanol solution at reflux. The optically active *trans* isomer **172** remained unaffected when treated under the analogous conditions. The epimerization must have occurred at C-1 of the *cis* diastereomer since only optically pure *trans* ester **172** was isolated from the reaction. If epimerization of the *cis* diastereomer had occurred at the C-3 carbon atom (ester) of **171**, it would have resulted in the formation of the enantiomer of **172** with an expected specific rotation ( $[\alpha]_D^{25} = +36^\circ$ ) equal and opposite to that of **172**. This was contrary to the observed results. Further evidence for the epimerization of *cis* **171** at C-1 to provide *trans* **172** was obtained on isolation of intermediates **176** and **177** (Scheme 14). Both the methyl ether **176** and alkene **177** were minor products (7%) obtained from the acid promoted conversion of *cis* **171** into *trans* **172**. The structures of **176** and **177** were determined on the mixture by COSY NMR and mass spectroscopy.<sup>111</sup> When a mixture of intermediates **176** and **177** was heated in methanolic HCl (1%), the *trans* diastereomer **172** was isolated as the sole product (70%),  $[\alpha]_D^{25} = -36.6^\circ$  ( $c = 1.0$ ,  $\text{CHCl}_3$ ). None of the *cis* isomer **171** was observed. On the basis of the above experiments, the epimerization of (1*R*,3*R*)-(+)–*cis*-**171** to provide (1*S*,3*R*)-(–)-*trans*-**172** had occurred regioselectively at C-1 and could be rationalized, as illustrated in Scheme 14. Under the conditions of heat and acid the  $N_b$ -nitrogen atom of *cis* isomer **171** can be protonated, followed by ring cleavage across the 1,2(C–N) bond to furnish the carbocations **174** or **175**. The carbocation **174** (or **175**) may either react with methanol to give 1-substituted ether **176** or lose a proton to furnish alkene **177**. These carbocations, **174** or **175**, which occupy a central position in the equilibrium, may also cyclize via a sterically favored conformation **175** to furnish

**Scheme 15. Alternate Mechanism for the *Cis* to *Trans* Epimerization at C-1 Does Not Operate Here**

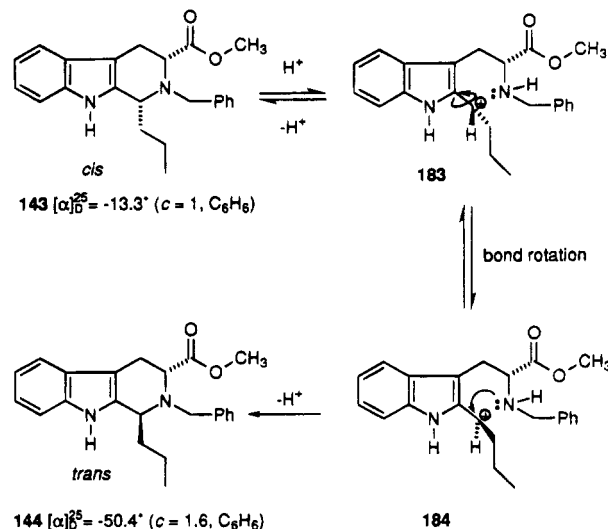


the optically active *trans* diastereomer **172**. Presumably, an equilibrium between the *cis* 1,3-diaxial **171** and *cis* 1,3-diequatorial **173** conformers existed at high temperature. The driving force for the ring scission between C-1 and N-2 was presumed to result from relief of A<sup>(1,2)</sup> strain<sup>69,121</sup> between substituents located at C-1 and N-9 in the diequatorial conformer **173** or the 1,3-diaxial interactions between substituents at C-1 and C-3 in the corresponding diaxial conformer **171**.

Another possible mechanism for this *cis*–*trans* epimerization involved the initial protonation of the indole 3-position and subsequent formation of an olefinic species (Scheme 15). This olefin could then be protonated from either face of the double bond to yield the more thermodynamically stable *trans* diastereomer **182**. Although Zhang provided strong evidence with  $\alpha$ -ketoglutaric acid that this olefinic equilibration was not involved,<sup>111</sup> recent results have further confirmed this earlier work.

In order to test the model for carbon–nitrogen bond cleavage versus the olefinic protonation model in the absence of an *N*<sub>a</sub>-methyl function, two experiments were carried out. In the first, optically active *N*<sub>b</sub>-benzyltryptophan methyl ester was synthesized from D-(+)-tryptophan according to the method employed for the related *N*<sub>a</sub>-methyl compound, as described above. This material was stirred under the nonacidic aprotic conditions of benzene at reflux under a nitrogen atmosphere with freshly distilled butyraldehyde **158** to kinetically trap the optically active diastereomers as a 23:77 *cis/trans* mixture. Separation of the diastereomers on silica gel with CH<sub>2</sub>Cl<sub>2</sub> by gravity chromatography provided the pure *cis* and *trans* diastereomers **143** and **144**. The specific rotation  $[\alpha]^{25}_D$  of the *trans* diastereomer **144** was measured as  $-50.4^\circ$  ( $c = 1$ , benzene) and the *cis* **143** as  $[\alpha]^{25}_D = -13.3^\circ$  ( $c = 1$ , benzene). The (–)-*cis* diastereomer **143** was then taken up in CH<sub>2</sub>Cl<sub>2</sub> and stirred

**Scheme 16. Proposed Mechanism for the Acid-Catalyzed Epimerization of the *Cis* to *Trans* Tetrahydro- $\beta$ -carboline**



with 2 equiv of TFA. Examination of the mixture by TLC after 12 h revealed that the *cis* isomer had been completely converted into the *trans* diastereomer **144** (Scheme 16). Workup with mild base (NaHCO<sub>3</sub>) and chromatography on a short wash column of silica gel (CH<sub>2</sub>Cl<sub>2</sub>) yielded the (–)-*trans* enantiomer **144**. The specific rotation  $[\alpha]^{25}_D$  of this sample was  $-50.4^\circ$  ( $c = 1.6$ , benzene). Since the specific rotations of the two (–)-*trans* compounds were identical (as well as the NMR and mass spectra) the epimerization that Zhang observed across the C(1)–N(2) bond had occurred here as well.

To test the alternate mechanism of *cis*–*trans* epimerization, the same experiment was repeated with 2 molar equiv of pure CF<sub>3</sub>COOD (Scheme 15). If the olefinic protonation mechanism had operated, then deuterium would have been incorporated into the tetrahydro- $\beta$ -carboline at C-1 upon protonation of the olefin **180** to re-form the *trans* tetrahydro- $\beta$ -carboline **144**. Repke et al. in an ibogaine derivative, have shown that incorporation of deuterium at position 1 was found to occur when an olefinic mechanism of isomerization was in operation.<sup>122</sup> We found no evidence for deuterium incorporation at C-1 of **144** by integration of the <sup>1</sup>H NMR spectrum nor by mass spectroscopy.

It was clear that the isomerization of the *cis* diastereomers to the more thermodynamically stable *trans* diastereomers (in both cases—Scheme 14 and Scheme 16) did not take place by the olefinic intermediate **180** (Scheme 15), but presumably by cleavage across the C(1)–N(2) bond (Schemes 14 and 16). In the experiments which involve *N*<sub>a</sub>-H derivatives **143** and **144**, it was observed that upon treatment of the *cis* isomer **143** with acid, **143** was completely converted into the *trans* diastereomer **144**, as illustrated in Scheme 16. After examination of the reaction mixture by TLC, *cis* isomer **143** was not observed and the reaction mixture was then heated to reflux and followed hourly by TLC. After prolonged heating no *cis* isomer (**143**) was observed in the mixture. Furthermore, there was no evidence of the original starting material (*N*<sub>a</sub>-hydro-*N*<sub>b</sub>-benzyl-

tryptophan methyl ester) which would be indicative of a retro-Pictet–Spengler-type process. From these experiments it has been concluded that the *trans* diastereomers are thermodynamically more stable than their *cis* counterparts in the  $N_a$ -hydro- $N_b$ -alkyl series as well. Hence, exclusive formation of the *trans* isomer would occur if the *cis* diastereomer were allowed to react under acidic conditions. Massiot and Mulamba<sup>123</sup> reported that a *trans* tetrahydro- $\beta$ -carboline was formed exclusively when methyl 4-formyl-2,2-bis(phenylthio)butyrate and tryptophanamide were heated to form an imine after which cyclization was effected with TFA at room temperature in  $\text{CH}_2\text{Cl}_2$ . On the basis of the results under aprotic conditions which yielded diastereomeric mixtures (Table 9), it seems more reasonable that a mixture of the *cis* and *trans* diastereomers had been formed by Massiot et al. and that upon introduction of TFA the *cis* diastereomer had epimerized to the more thermodynamically stable *trans* diastereomer. When optically active  $N_b$ -benzyl-D-(+)-tryptophan methyl ester and butyraldehyde were again heated in benzene at reflux until TLC indicated the consumption of all the indolic starting material, it was found that the *trans* diastereomer **144** could exclusively be formed upon addition of 10 equiv of TFA at room temperature to the reaction mixture. The epimerization of the *cis* diastereomer **143** to the *trans* diastereomer **144** was monitored by TLC (74% yield). This has important implications for the enantiospecific synthesis of indole alkaloids. The use of  $N_b$ -benzyl or  $N_b$ -diphenylmethyl substituents in the tryptophan methyl ester series can provide the *trans* diastereomer kinetically with 100% diastereoselectivity. In cases where a mixture of the *cis* and *trans* diastereomers are formed, the thermodynamic preference for the *trans* isomer in the  $N_b$ -alkyl series can be employed to convert the entire mixture into the desired *trans* isomer with 100% diastereoselectivity. Moreover, if the Pictet–Spengler reaction is carried out in benzene at reflux followed by the addition of TFA afterward, this will provide the *trans* isomer stereospecifically (substituent at C-1 greater than  $\text{CH}_3$ ). Since both D-(+) and L-(-)-tryptophan are commercially available, both optical antipodes of the alkaloids are available enantiospecifically. Furthermore, the realization of the recent enantiospecific synthesis of both 5-methoxy- and 6-methoxy-D-(+)- or L-(-)-tryptophans provides enantiospecific entry into over 70 different alkaloids via this method.

### III. Modeling the Pharmacophore of Benzodiazepine Receptor Sites

The study of benzodiazepine receptors has been one of the most rapidly increasing areas of research in molecular neuropsychopharmacology in recent years.<sup>124–127</sup> Benzodiazepines exhibit a wide range of clinical uses including the treatment of anxiety and related emotional disorders as well as sleep disorders and as anticonvulsants. They behave as centrally acting muscle relaxants. In 1975 examination of evidence from behavioral, electrophysiological, and biochemical experiments, indicated that benzodiazepines act at synapses in which  $\gamma$ -aminobutyric acid (GABA) is the acting neurotransmitter.<sup>128</sup> In 1977

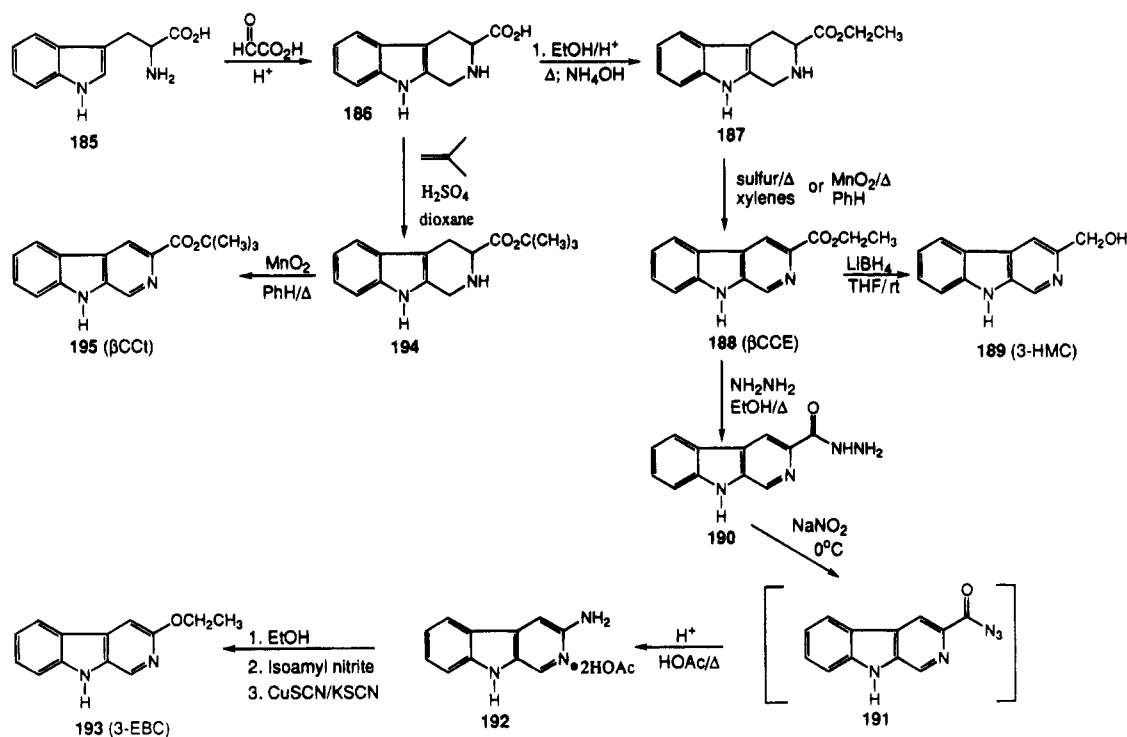
the identification of specific, saturable, high-affinity binding sites for tritiated diazepam by two independent research groups, Möhler and Okada,<sup>129</sup> as well as Squires and Braestrup,<sup>130</sup> generated a tremendous interest in benzodiazepine receptor (BzR) sites. It was later determined through various biochemical techniques that the BzR comprises a portion of the GABA<sub>A</sub> receptor chloride ion complex. In addition to the benzodiazepine binding site, there exists at least three other modulatory binding domains on the complex corresponding to sites for barbiturates, picrotoxin, and anesthetic steroids.<sup>126</sup>

More recently, the GABA<sub>A</sub> receptor has been shown to be a heterooligomeric family of ligand-gated ion channels which constitutes the major inhibitory neurotransmitter system in the central nervous system (CNS).<sup>125,126,131–134</sup> This membrane-bound protein complex plays a central role in the molecular mechanisms which underlie anxiety, sleep, convulsions, memory-learning, and consequently it represents an important target for the design of selective agents to treat specific disease states in the CNS.<sup>131–139</sup> The inhibitory GABA neurotransmitter or ligand, acts by binding to its receptor site which is followed by the opening of an intrinsic chloride ion channel. The increase in chloride ion flux results in hyperpolarization of the neuronal cell membrane with a concomitant decrease in neuronal transmission. The biological activity of modulatory BzR ligands of interest here encompasses a wide range of physiological responses ranging from full agonist (anxiolytic, muscle relaxant/ataxic, amnesic, sedative/hypnotic, and anticonvulsant) to full inverse agonist (anxiogenic, proconvulsant, and convulsant). Clinically, there is a need for partial agonists which exhibit anxiolytic/anticonvulsant activity in the absence of myorelaxant, ataxic, and sedative-hypnotic activity. Moreover, there is a need for partial (or selective) inverse agonists which enhance neuronal firing in the CNS but are devoid of the proconvulsant/convulsant activity of full inverse agonists. These latter ligands would be useful for treatment of barbiturate–alcohol induced CNS depression (overdose), hepatic encephalopathy, and cognition enhancement among others.<sup>137–139</sup> Recent studies have shown that many  $\beta$ -carbolines do elicit these pharmacological properties.<sup>44</sup>

### A. Molecular Biology

In 1987 Seeburg, Schofield, and co-workers reported the cloning and functional expression of a GABA/Bz receptor complex from bovine brain.<sup>140</sup> It was initially proposed the complex was composed of  $\alpha$ - and  $\beta$ -subunits. Subsequent to this report, several other subunits of the GABA<sub>A</sub> receptor have been characterized including  $\gamma$ -,  $\delta$ -, and  $\rho$ -subunits.<sup>127,141–143</sup> The identification of multiple  $\alpha$ -,  $\beta$ -, and  $\gamma$ -subunits is consistent with the pharmacological evidence of multiple GABA<sub>A</sub> receptor isoforms in the CNS.<sup>144</sup> Recent molecular biological studies have established that expression of at least two ( $\alpha$ ,  $\gamma$ ) and preferably three ( $\alpha$ ,  $\beta$ ,  $\gamma$ ) subunits is necessary to constitute a functional receptor which mimics many of the pharmacological, biochemical, and electrophysiological properties of native receptors.<sup>145–147</sup> Previous studies of site-directed mutagenesis have shown that changes

Scheme 17



in the  $\alpha$ -subunit altered the activity of the benzodiazepine receptor-mediated response.<sup>37,138,148,149</sup> It is now known that changes in the  $\gamma$ -subunit also alter this response.<sup>147,150–152</sup> However, neither the stoichiometry, nor the composition of native  $\text{GABA}_A$  receptors in the CNS has been unequivocally established. It appears now, on the basis of preliminary evidence,<sup>132,141,143,147,151,152</sup> that the BzR binding site lies between the  $\alpha$ - and  $\gamma$ -subunits.

## B. Receptor Subtypes

Initially it was proposed that the type-I BzR was responsible for the anxiolytic and anticonvulsant effects<sup>153</sup> of the benzodiazepines while the type-II BzR mediated the muscle relaxant, sedative-hypnotic properties of these ligands. Structure–activity relationship (SAR) data from a variety of recently synthesized BzR ligands,<sup>44,143,154–156</sup> however, demonstrated that the actual pharmacological situation is much more complicated. Bz type-I selective ligands (i.e. CL 217872) have now been shown to exhibit some sedative effects,<sup>154,157</sup> in contrast to previous reports.<sup>153</sup> In fact zolpidem is now marketed as a sedative-hypnotic. Receptor isoforms whose pharmacology resembles that of previously reported Bz-I and Bz-II receptors have recently been expressed: the type-I BzR was constructed from an  $\alpha 1\beta 2\gamma 2$  [ $\alpha 1\beta 2\gamma 2 = \omega 1 = \alpha 1$ ] combination of subunits while the type-II BzR was comprised of  $\alpha 2\beta 2\gamma 2$  [ $\alpha 2\beta 2\gamma 2 = \omega 2 = \alpha 2$ ],  $\alpha 3\beta 2\gamma 2$  [ $\alpha 3\beta 2\gamma 2 = \omega 3 = \alpha 3$ ], and  $\alpha 5\beta 2\gamma 2$  [ $\alpha 5\beta 2\gamma 2 = \omega 5 = \alpha 5$ ] (zolpidem insensitive)<sup>156</sup> receptor isoforms. Moreover, photolabeling experiments<sup>158</sup> with [ $^3\text{H}$ ]Ro 15-4513 have identified a site termed the “diazepam-insensitive” (DI) receptor which is made up of the subunits designated  $\alpha 4\beta 2\gamma 2$  [ $\alpha 4\beta 2\gamma 2 = \omega 4 = \alpha 4$ ] and  $\alpha 6\beta 2\gamma 2$  [ $\alpha 6\beta 2\gamma 2 = \omega 6 = \alpha 6$ ],<sup>158,159</sup> the latter of which has been studied extensively.<sup>44</sup> The physiological activity of agonists and inverse

agonists at benzodiazepine receptors in the CNS is therefore a consequence of action at one or more of the receptor subtypes. Studies are currently underway to correlate receptor subtype selectivity with a biological response.

## C. Biology of Selected 3-Substituted $\beta$ -Carbolines

It was originally proposed by Braestrup that  $\beta$ -carboline-3-carboxylic acid ethyl ester ( $\beta\text{CCE}$ , 188) shown in Scheme 17 was the endogenous ligand associated with the BzR.<sup>160,161</sup> Subsequent studies, however, have shown this ligand was formed during the isolation procedure.<sup>124,162</sup> Nonetheless, the demonstration that certain  $\beta$ -carbolines potently inhibit [ $^3\text{H}$ ]diazepam binding with high affinity (5 nM) has led to numerous studies of ligands which bind to benzodiazepine receptors as well as the development of potential new therapeutic agents.

It has been shown that the full inverse agonist  $\beta\text{CCE}$  (188) antagonized the anticonvulsant actions of diazepam, lowered the seizure threshold of the convulsant pentylenetetrazole, and antagonized the sedative actions of flurazepam.<sup>163–165</sup> More importantly, when  $\beta\text{CCE}$  (188) was administered to primates<sup>36</sup> it elicited a profound behavioral and physiological syndrome reminiscent of “fear” or “anxiety” related to the same effects experienced in man. For example, in *Rhesus* monkeys this substance produced dramatic elevations in heart rate, blood pressure, plasma cortisol, and catecholamines.<sup>36</sup> These effects were blocked by benzodiazepines and the specific benzodiazepine receptor antagonist Ro 15-1788 (flumazenil). The results of this study demonstrated that the administration of  $\beta\text{CCE}$  (188) to animals may represent a reliable and reproducible model of human anxiety and as such, could be valuable in studying the postulated role of anxiety and stress in a variety of human diseases, including cardiovascu-

lar, ulcerative, and neoplastic disorders. In a beautiful series of experiments with  $\beta$ CCE (**188**) Mueller has shown that a BzR mechanism participates in the physiological regulation of  $\beta$ -endorphin-like immunoreactivity ( $\beta$ -END-LI) secretion from the AL of the rat pituitary gland. Accordingly, the ability of the anxiogenic  $\beta$ -carbolines  $\beta$ CCE (**188**) and  $\beta$ CCM (methyl ester) to stimulate rapid and pronounced increases in plasma  $\beta$ -END-LI indicates that anxiety, *per se*, is a stimulus for  $\beta$ -END-LI release. Furthermore, the findings raise the possibility that pituitary  $\beta$ -END peptides may have functions related specifically to anxiety and separate from pain.<sup>166</sup>

Although  $\beta$ CCE (**188**) is a potent inverse agonist, it has several serious drawbacks when used for *in vivo* studies.  $\beta$ CCE (**188**) is readily susceptible to esterase hydrolysis<sup>43</sup> to the inactive acid (especially in rodents), and thus large quantities of this compound are needed for pharmacological studies. In addition,  $\beta$ CCE (**188**) is only sparingly soluble in water, making administration of this compound difficult. Therefore, the search for water-soluble, long-lived partial inverse agonists led to the synthesis of 3-ethoxy- $\beta$ -carboline (3-EBC, **193**), 3-(hydroxymethyl)- $\beta$ -carboline (3-HMC, **189**) and  $\beta$ -carboline-3-carboxylic acid *tert*-butyl ester ( $\beta$ CCt, **195**), as shown in Scheme 17. The biology of these 3-substituted  $\beta$ -carbolines is discussed below.

Pharmacological studies on 3-HMC<sup>167</sup> (**189**) have shown this  $\beta$ -carboline inhibited [<sup>3</sup>H]diazepam binding *in vitro* ( $K_i = 1470$  nM), and antagonized both the anticonvulsant and anxiolytic actions of diazepam at doses which did not elicit overt behavioral effects. This  $\beta$ -carboline also inhibited the sleep-inducing effects of flurazepam<sup>38</sup> and was oxidized to the active aldehyde ( $IC_{50} = 62$  nM) *in vivo*. Thus the hypnotic actions of flurazepam were strongly felt to originate through interaction with the benzodiazepine receptor. At slightly higher doses, 3-HMC (**189**) increased wakefulness in rodents by significantly increasing sleep latency and reducing non-REM (but not REM) sleep. Consequently 3-HMC (**189**) was not merely a benzodiazepine antagonist but exerted a pharmacological action opposite to that effected by 1,4-benzodiazepines. Although often drugs such as amphetamines and methylxanthines can reduce sleep, they also invariably cause profound alterations in behavior and motor activity. Compounds that reduce sleep without eliciting major changes in motor activity may, therefore, be more properly termed "somnolytics".<sup>38</sup> The suggestion (with 3-HMC, **189**) that the BzR was involved in both physiological and pharmacologically induced sleep<sup>38</sup> could lead to the development of  $\beta$ -carbolines or related compounds for treating human sleep disorders, especially those characterized by excessive somnolence.

3-Ethoxy- $\beta$ -carboline (3-EBC, **193**) bound with high affinity to the BzR ( $IC_{50} = 24$  nM). Trullas et al.<sup>43</sup> have shown that 3-EBC (**193**) potentiated the convulsant actions of pentylenetetrazole in mice consequently, it is proconvulsant. Furthermore, this compound reduced both the time spent and the total entries in the open arms of an elevated plus maze. Moreover 3-EBC (**193**) inhibited stress-induced ulcer formation in mice. These effects are common to

benzodiazepine receptor inverse agonists, suggesting this compound is an inverse agonist at benzodiazepine receptors. Although 3-EBC (**193**) is proconvulsant, even at doses as high as 40 mg/kg it does not exhibit convulsant effects, consequently, it is termed a partial inverse agonist. In addition, this ligand has a higher affinity for benzodiazepine receptors and better water solubility (13 mg/mL of H<sub>2</sub>O)<sup>168</sup> as a hydrochloride salt than the commonly employed inverse agonist FG 7142 (<1 mg/mL of H<sub>2</sub>O), making this  $\beta$ -carboline an attractive ligand for *in vivo* studies in models of anxiety.

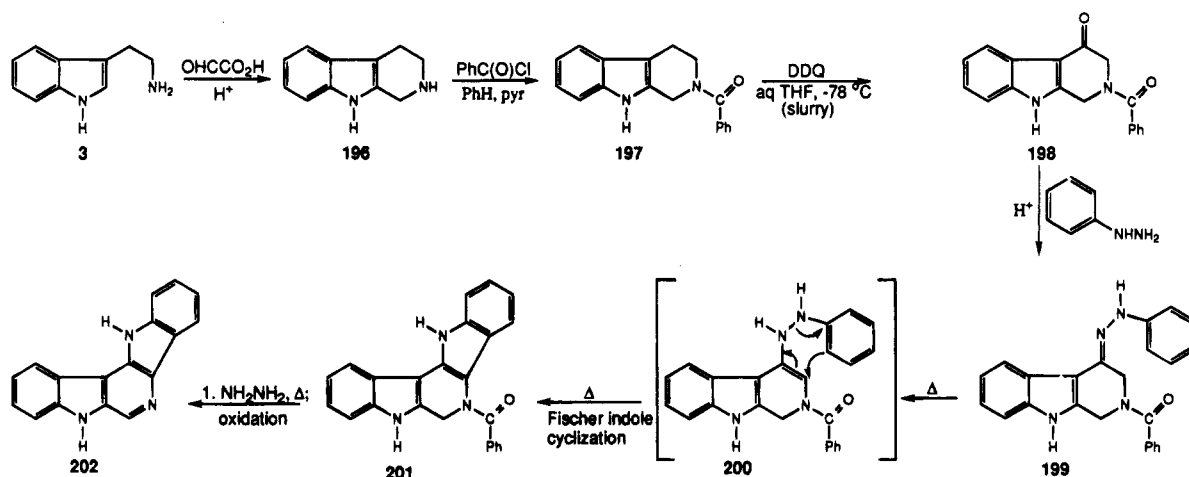
The  $\beta$ -carboline  $\beta$ CCt (**195**) was synthesized to provide a BzR ligand which would possess a longer duration of action than either ethyl ester  $\beta$ CCE (**188**) or methyl ester  $\beta$ CCM. This  $\beta$ -carboline **195** did bind tightly to BzR ( $IC_{50} = 10$  nM) obtained from synaptosomal membranes and was much longer lived *in vivo* than the related ethyl or methyl esters.<sup>39</sup> The ability of  $\beta$ CCt (**195**) to antagonize the anticonvulsant, anxiolytic, ataxic, and muscle relaxant effects of diazepam (Valium) was evaluated by Shannon.<sup>39</sup> In mice  $\beta$ CCt (**195**) at doses of 3 and 10 mg/kg produced a dose-related antagonism of the anticonvulsant effects of diazepam against pentylenetetrazole (80 mg/kg) induced seizures. A dose of 30 mg/kg of  $\beta$ CCt (**195**) did not produce a further shift in the diazepam dose–effect curve, apparently because  $\beta$ CCt (**195**) failed to block the muscle relaxant effects of diazepam. Furthermore, **195** (30 mg/kg) failed to antagonize the ataxic effects of diazepam in the inverted wire screen test.

$\beta$ CCt (**195**) did selectively antagonize the effects (anxiolytic) of diazepam on punished behavior as well as the anticonvulsant effects of Valium; however, **195** failed to antagonize the rate-decreasing and ataxic effects of diazepam. These results suggested that  $\beta$ CCt (**195**) was a selective Bz<sub>I</sub> benzodiazepine receptor antagonist employing the earlier terminology of Beer and Lippa.<sup>153,169</sup>

The response–rate-decreasing effects of diazepam under both the punished and unpunished components of the multiple schedule occurred at doses which produced ataxia as measured on the rotarod in rats and are likely a direct result of the sedative, muscle relaxant, and ataxic effects of diazepam. Consequently, the ineffectiveness of  $\beta$ CCt **195** in blocking the rate-decreasing effects of diazepam is further evidence that **195** failed to block the ataxic and sedative effects of diazepam. This selectivity of the antagonist actions of  $\beta$ CCt (**195**) distinguished it from  $\beta$ CCE (**188**) or  $\beta$ CCM (**218**). The latter two  $\beta$ -carbolines antagonized the diazepam- or phenobarbital-induced impairment of motor performance in the horizontal wire test in mice.

In summary,  $\beta$ CCt (**195**) antagonized the anticonvulsant and antipunishment (anxiolytic) effects of diazepam but not the ataxic, sedative, and muscle-relaxant effects. It was proposed in the early 1980s<sup>153,169</sup> that Bz<sub>I</sub> receptors mediated the anxiolytic and anticonvulsant properties of benzodiazepines, whereas Bz<sub>II</sub> (type II) receptors mediated the depressant and ataxic effects of the benzodiazepines. Within the framework of this early constraint,  $\beta$ CCt (**195**) appeared to be a Bz<sub>I</sub> receptor antagonist. Although

Scheme 18



the antagonist properties of **195** are as efficacious as those of the clinically employed flumazenil (Ro 15-1788)  $\beta$ CCt (**195**) appeared to be more Bz<sub>1</sub> selective than the Roche antagonist.

The long half-life *in vivo* of  $\beta$ CCt (**195**) when compared to  $\beta$ CCE (**188**) rendered **195** an important pharmacological probe of sleep<sup>170</sup> and other processes mediated by the CNS.<sup>171</sup> Because physostigmine had been reported to reverse the sedation and paradoxical delirium induced by the 1,4-benzodiazepines, Hoffman employed  $\beta$ CCt (**195**) to study this phenomenon. Hoffman<sup>171</sup> found that physostigmine increased cerebral blood flow (CBF) and cerebral oxygen consumption (CMR<sub>O<sub>2</sub></sub>) in the CNS as reported previously. The increase in CBF was linked primarily to increases in CMR<sub>O<sub>2</sub></sub> and a normal coupling between blood flow and metabolism. The benzodiazepine agonist midazolam, decreased CBF and CMR<sub>O<sub>2</sub></sub> in a dose-related manner, and this decrease was antagonized by physostigmine.  $\beta$ CCt (**195**) increased CBF and CMR<sub>O<sub>2</sub></sub> but this change was further potentiated by physostigmine.<sup>171</sup> The additive effects of  $\beta$ CCt (**195**) and physostigmine suggested that physostigmine antagonized the central action of benzodiazepines by a central stimulatory action, and not by a direct competitive effect at the BzR as previously suggested by others. If physostigmine produced stimulation by an action at the central BzR/GABA<sub>A</sub> complex, this action was probably independent of the BzR and in keeping with the cholinergic properties of physostigmine.

#### D. Pharmacophore Models of the Benzodiazepine Receptor Site

The practice of incorporating biofunctionality into a rigid framework to enhance activity or selectivity of action is often employed when trying to define a pharmacophore for a specific receptor site. The early work of Bentley et al.<sup>172,173</sup> in the morphine area is an excellent example of this. Previous structure-activity relationship (SAR) studies<sup>174-177</sup> suggested that one necessary criteria for high-affinity binding of ligands to benzodiazepine receptors was the ability of these molecules to assume a planar or pseudoplanar topography.<sup>174,175</sup> With these goals in mind Trudell et al.<sup>178</sup> in 1987 first reported the biological

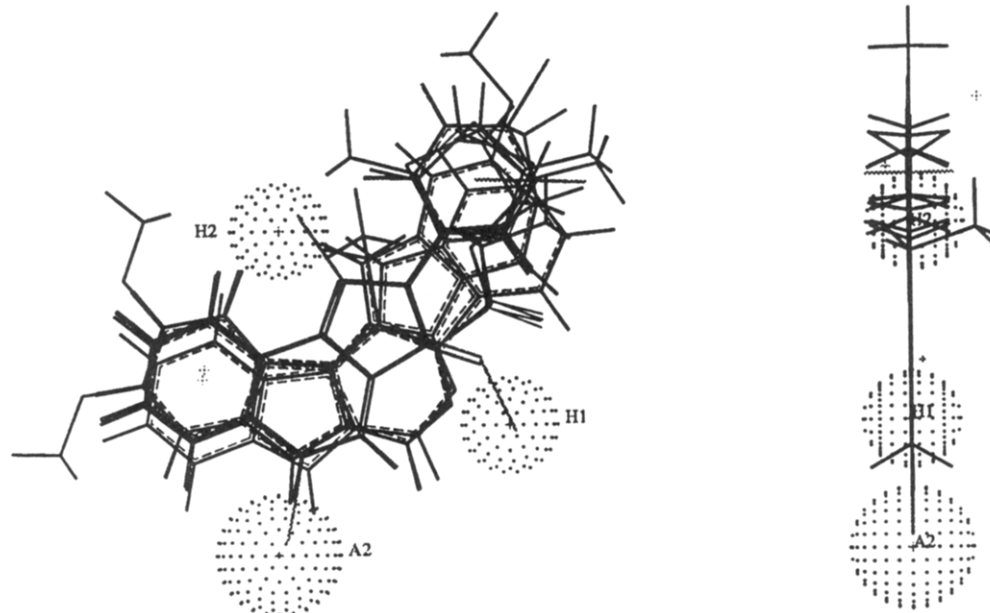
activity of the rigid planar 7,12-dihydropyridof[3,2-*b*:5,4-*b'*]diindole (**202**) the synthesis of which is illustrated in Scheme 18.

The tetrahydro- $\beta$ -carboline **196** required for this synthesis was prepared in 84% yield from tryptamine **3** via a Pictet-Spengler reaction with glyoxylic acid monohydrate (Scheme 18). Protection of the amine function at N-2 by treatment of the tetrahydro- $\beta$ -carboline **196** with benzoyl chloride furnished the two rotameric benzoyl-1,2,3,4-tetrahydro- $\beta$ -carbolines represented by **197** in 79% yield. The benzamide **197** which resulted was then oxidized<sup>179</sup> regioselectively at C-4 with dichlorodicyanoquinone (DDQ) in aqueous THF at  $-78$  °C to furnish the 2-benzoyl-4-oxo-1,2,3,4-tetrahydro- $\beta$ -carboline **198**, in 56% yield.<sup>99</sup> The execution of this procedure at low temperature provided the regioselectivity of this process; however, it required reaction with DDQ as a slurry. Treatment of the 4-oxo ketobenzamide with phenylhydrazine resulted in the formation of hydrazone **199** in excellent yield. When the hydrazone **199** was heated, the obligatory [3,3] sigmatropic rearrangement of the Fischer indole reaction took place to provide the indole analog **201**. Removal of the amide functionality was accomplished by heating indolobenzamide **201** in the presence of hydrazine to furnish the fully aromatic dihydropyridodiindole (**202**) in excellent yield.

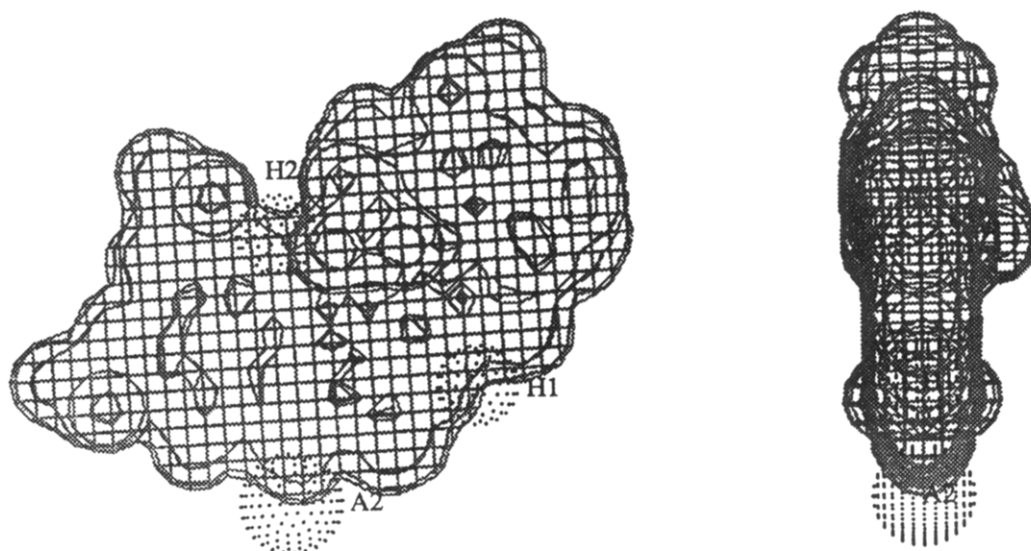
The dihydropyridodiindole (**202**) was found to bind tightly to the BzR (4 nM) and exhibited inverse agonist activity reminiscent of the activity of  $\beta$ CCE (**188**) and FG 7142. The rigid nature of this ligand permitted the effective superposition (using a template approach) of 12 other inverse agonists via the active analog approach of Marshall,<sup>180,181</sup> to arrive at the topography of the inverse agonist/antagonist site in the binding cleft at the BzR (Figure 6). An included volume analysis of these ligands resulted in the inverse agonist/antagonist pharmacophore illustrated in Figure 7. In addition to the pyridodiindoles, >100 3-substituted  $\beta$ -carbolines have been synthesized and modeled<sup>40-42,44</sup> in our laboratory to define the topography of the inverse agonist/antagonist BzR pharmacophore receptor model illustrated in Figure 7.

The results of this study suggested for potent inverse agonist activity the involvement of an indole





**Figure 6.** The superposition of 12 inverse agonist ligands at the benzodiazepine binding site ( $\beta$ CCE, DMCM, 3-EBC, 3-propionyl- $\beta$ -carboline, CGS-8216, 7,12-dihydropyrido[3,2-*b*:5,4-*b'*]diindole, 2-methoxy-7,12-dihydroxyprido[3,2-*b*:5,4-*b'*]diindole, 2-thienylpyrazolo[3,4-*c*]quinolin-3-one, 2-(4'-methylthienyl)pyrazolo[3,4-*c*]quinolin-3-one, 3-thienylpyrazolo[3,4-*c*]quinolin-3-one, 3-(5'-methylthienyl)pyrazolo[3,4-*c*]quinolin-3-one, and 3-(4'-methylthienyl)pyrazolo[3,4-*c*]quinolin-3-one). The inverse agonist pharmacophoric descriptors H<sub>1</sub> and H<sub>2</sub> represent hydrogen-bond donor sites on the protein; A<sub>2</sub> represents a hydrogen-bond acceptor site on the protein. The illustration on the right side of the picture (orthographic stereoview) originates from rotation of the pharmacophore 90° to the right.



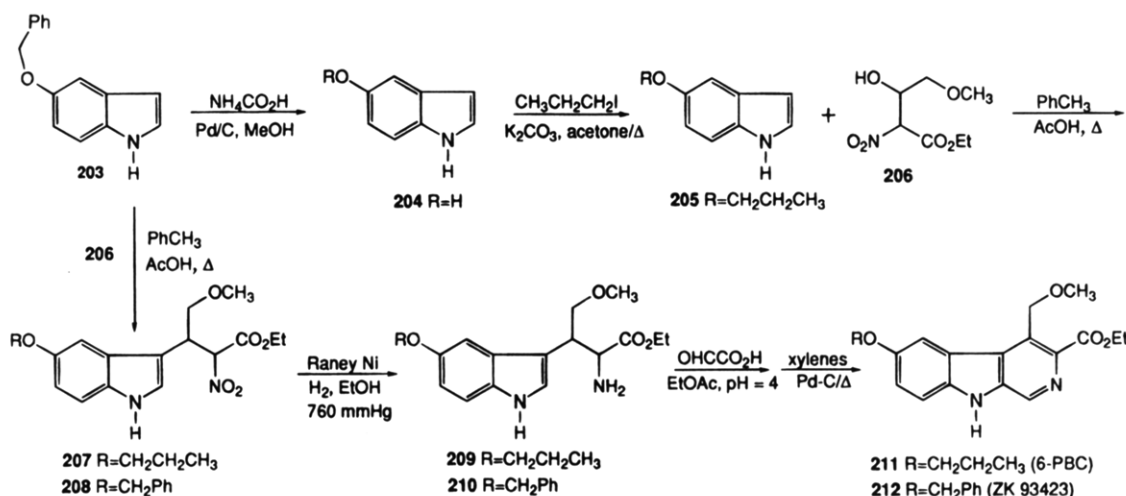
**Figure 7.** The included volume analysis of 12 inverse agonist ligands at the benzodiazepine binding site.

N(9)–H in a hydrogen-bond interaction of the ligand with a hydrogen-bond acceptor site (A<sub>2</sub>) on the receptor was necessary, in contrast to several other pharmacophore models.<sup>44,182–185</sup> In addition, the proposed model required the pyridine N-2 nitrogen atom of a  $\beta$ -carboline or N-5 of a diindole to exhibit hydrogen bond acceptor (N:) properties and interact with a receptor hydrogen bond donating site (H<sub>1</sub>) on the protein.<sup>40,186</sup> Lastly, it was proposed there existed a hydrophobic pocket (L<sub>1</sub>) in the receptor protein near position 3 of the  $\beta$ -carboline framework. The affinity of various 3-substituted  $\beta$ -carbolines suggested this pocket has a definite length and width.<sup>41,42</sup> It was also noted by Allen et al.<sup>42</sup> that substituents at the 3-position of  $\beta$ -carbolines have a strong influence on both the affinity and type of activity. The synthesis of the long-lived water-soluble partial inverse agonist

3-ethoxy- $\beta$ -carboline **193** (whose biology was described above) resulted from this modeling.<sup>40–42</sup> With the development of the inverse agonist/antagonist pharmacophore, attention then focused on modeling the agonist pharmacophore of the BzR. It should be noted that controversy existed in the literature as to whether inverse agonists and agonists bound to the same receptor or to different receptor sites.<sup>187,188</sup> Inverse agonists and agonists were treated as separate entities in our work although the common sites of overlap between the two pharmacophores were established from molecular modeling in agreement with the previously reported domain model of Skolnick.<sup>134</sup>

There are a wide variety of compounds which exhibit full agonist activity at the BzR but are structurally unrelated to 1,4-benzodiazepines (e.g.

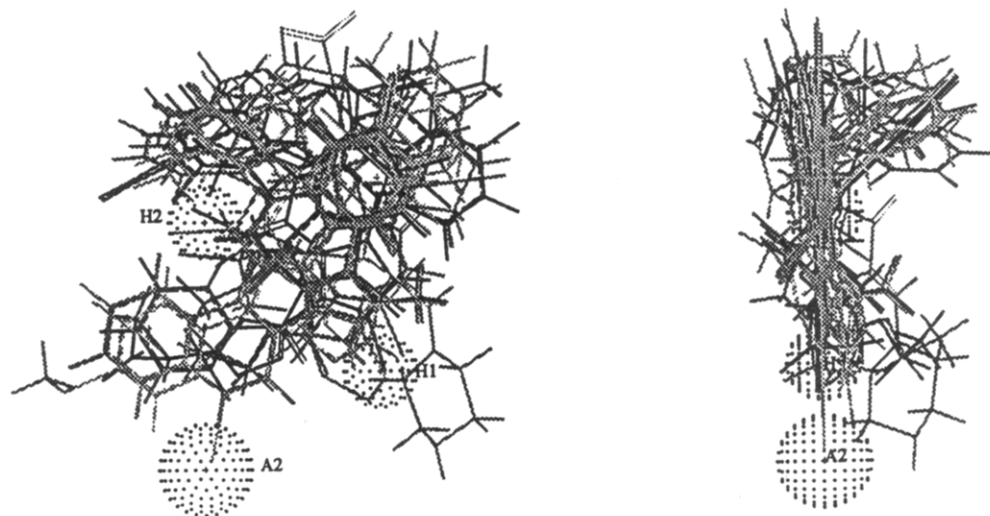
## Scheme 19



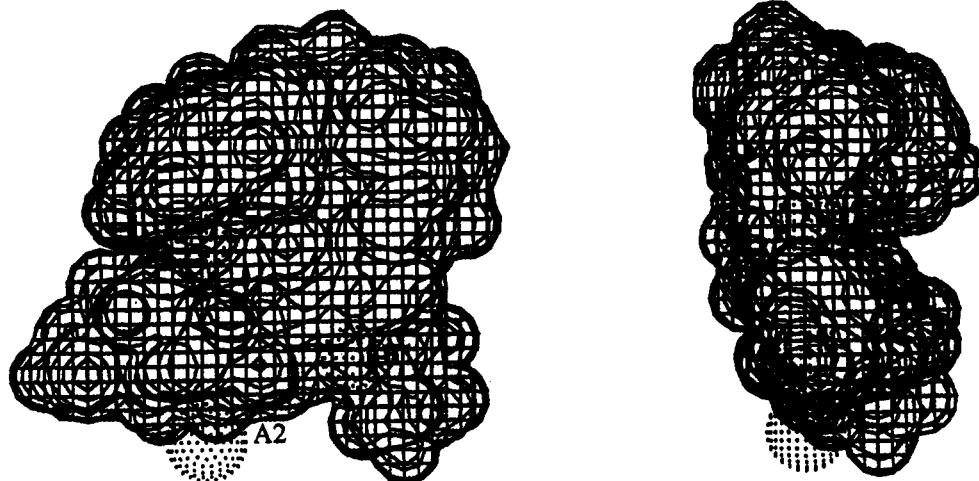
Valium). There had only been one  $\beta$ -carboline reported to date<sup>189</sup> which exhibited *full* agonist activity at the BzR, i.e. the  $\beta$ -carboline derivative 6-(benzyloxy)-4-(methoxymethyl)- $\beta$ -carboline-3-carboxylic acid ethyl ester (ZK 93423, **212**) (Scheme 19). The  $\beta$ -carboline ligands are some of the most difficult to model<sup>190,191</sup> due to the conformational flexibility of substituents located at positions 3, 4, 5, and 6, therefore, the use of ZK 93423 (**212**) was important in establishing the points of electronic interaction necessary for agonist activity. The development of the agonist pharmacophore was carried out in our laboratory by Diaz-Arauzo et al.<sup>191</sup> and then overlapped with the inverse agonist/antagonist model. The strategy employed was analogous to that reported for the inverse agonist/antagonist receptor model and employed 36 different ligands which

belonged to 10 different structural families (now 13) and represented 136 different ligands.<sup>44,190,191</sup> The alignment of the agonist ligands and the included volume analysis of these ligands is illustrated in Figures 8 and 9.

The data from the synthetic and computer-assisted analysis of the agonist pharmacophore at the BzR<sup>191</sup> suggested that this site contained two hydrogen bond donating sites (H<sub>1</sub> and H<sub>2</sub>), as illustrated in Figure 8. These two sites are located about 6.7 Å from each other. The binding site H<sub>1</sub> is common to both the agonist and inverse agonist pharmacophores, but region H<sub>2</sub> (agonist model, Figure 8) corresponded to the 4-position of  $\beta$ -carbolines or interaction at N-4 of the 1,4-benzodiazepines.<sup>44</sup> With respect to  $\beta$ -carbolines it was believed the oxygen atom at this position was of critical importance for agonists in the  $\beta$ -car-



**Figure 8.** The superposition of 31 agonist ligands to define the agonist pharmacophore at the benzodiazepine binding site (diazepam, flunitrazepam, brotizolam, midazolam, triazolam, norflunitrazepam, 7-aminoflurazepam, 7,2'-dichloro-thieno[2,3-*e*][1,4]benzodiazepine, 1-methyl-8-chloro-2'-fluoro-*s*-triazolo[4,3-*a*][1,4]benzodiazepine, 2,9-dichloropyrimido[5,4-*d*][2]benzazepine, 4-methoxy-6-(alkylamino)-3-aryl-1,2,4-triazolo[3,4-*a*]phthalazine, 4-chloro-6-(alkylamino)-3-aryl-1,2,4-triazolo[3,4-*a*]phthalazine, 4-methoxy-6-(alkylamino)-3-aryl-1,2,4-triazolo[3,4-*a*]phthalazine, 4-fluoro-6-(alkylamino)-3-aryl-1,2,4-triazolo[3,4-*a*]phthalazine, 2-benzoyl-5-methoxy-7-ethylimidazo[1,2-*a*]quinoline, 2-benzoyl-5-(methylthio)-6-ethyl-7-methylimidazo[1,2-*c*]pyrimidine, 2-(4'-chlorophenyl)pyrazolo[3,4-*c*]quinolin-3-one, 2-(4'-methoxyphenyl)pyrazolo[3,4-*c*]quinolin-3-one, 2-(2'-methylthienyl)pyrazolo[3,4-*c*]quinolin-3-one, 2-(5'-ethylthienyl)pyrazolo[3,4-*c*]quinolin-3-one, 2-(5'-butylthienyl)pyrazolo[3,4-*c*]quinolin-3-one, 2-(4',5'-dimethylthienyl)pyrazolo[3,4-*c*]quinolin-3-one, 2-(5'-butylthienyl)pyrazolo[3,4-*c*]quinolin-3-one, ZK 93423, ZK 93426, 6-(benzyloxy)-4-(methoxymethyl)- $\beta$ -carboline-3-carboxylic acid isopropyl ester, 6-PBC, loprazolam, delorazepam). The agonist pharmacophoric descriptors H<sub>1</sub> and H<sub>2</sub> represent hydrogen-bond donor sites on the protein. The illustration on the right side of the picture (orthographic stereoview) originates from rotation of the pharmacophore 90° to the right.

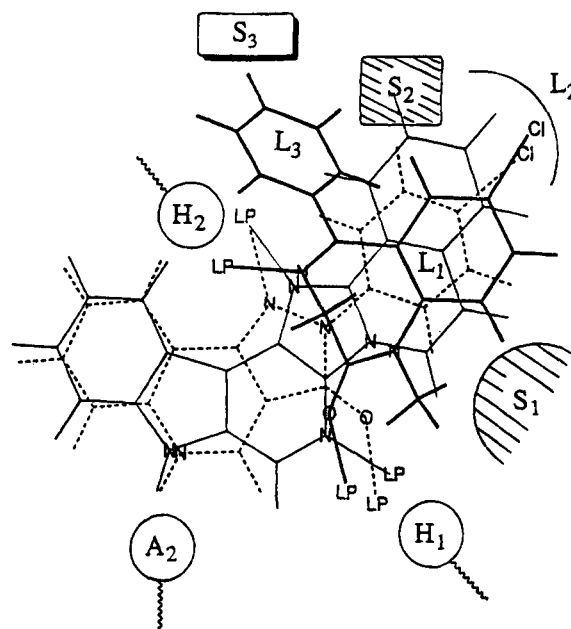


**Figure 9.** The included volume analysis of 31 agonist ligands at the benzodiazepine binding site.

boline series and directed the ligand into the active site of the agonist receptor region through the formation of a hydrogen bond between the ether oxygen atom at C-4 and H<sub>2</sub> of the receptor protein. In addition there are three areas of lipophilic interaction (L<sub>1</sub>, L<sub>2</sub>, and L<sub>3</sub>). Occupation of the areas L<sub>2</sub>/L<sub>3</sub> and interaction at H<sub>1</sub>, H<sub>2</sub>, and L<sub>1</sub> are important for full agonist activity; full agonist activity appears to require complete occupation of L<sub>2</sub> and L<sub>3</sub>. This is in agreement with previous work.<sup>40,41,176,177,182,184-186,192-194</sup>

Substituents which occupy L<sub>3</sub> cannot lie in the same plane as H<sub>1</sub>, H<sub>2</sub>, and L<sub>1</sub> for they would interfere with the hydrogen bonding protein-ligand interaction at H<sub>2</sub> which would eliminate agonist activity. Areas of negative steric interaction (S<sub>1</sub>, S<sub>2</sub>, and S<sub>3</sub>) between the ligand and receptor-binding protein have also been defined and are illustrated in Figure 10.

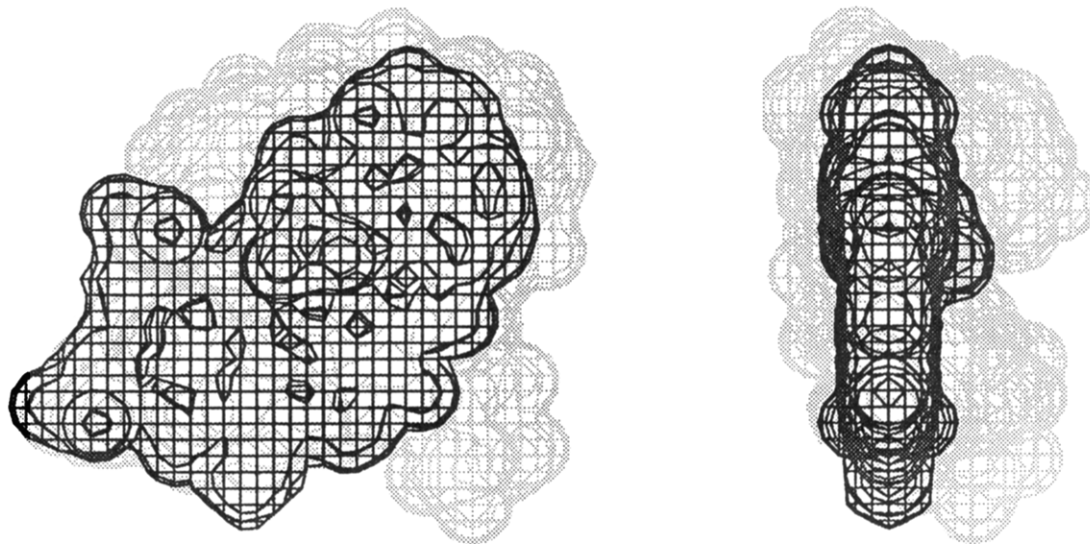
With regard to this pharmacophore, it is felt the alignment rule for agonist  $\beta$ -carbolines is different from that which elicit inverse agonist activity. Examination of Figure 11 shows the overlap of the inverse agonist (light-colored area) and agonist (dark-colored area) pharmacophores. It is now clear from molecular graphics that inverse agonists bind to the same domain as agonists.<sup>44,168,195</sup> As illustrated, the inverse agonist pharmacophore is planar while the agonist pharmacophore contains a lipophilic pocket (L<sub>3</sub>) out of the plane. It is also apparent that the inverse agonist pharmacophore is considerably smaller in size relative to the agonist receptor/pharmacophore model. As alluded to earlier, it has been proposed that these pharmacophores bind to the same area at the receptor site, but contain different pharmacophoric descriptors in order to elicit the opposite biological response. As evident from the included volumes the interaction at the pharmacophoric descriptors required for agonist activity (H<sub>1</sub>, H<sub>2</sub>, L<sub>1</sub>, L<sub>2</sub>, and/or L<sub>3</sub>) and inverse agonist activity (H<sub>1</sub>, A<sub>2</sub>, and L<sub>1</sub>) are clearly different although H<sub>1</sub> and L<sub>1</sub> are common descriptors for both types of ligands. Evidence to date suggests that ligands which exhibit full agonist activity (diazepam, ZK 93423, etc.) fully occupy lipophilic regions L<sub>2</sub> and L<sub>3</sub> as well as interacting at L<sub>1</sub>, H<sub>1</sub>, and H<sub>2</sub>. Analysis of limited evidence suggests that the target partial agonists do not fully occupy both L<sub>2</sub> and L<sub>3</sub>, but that one or more interactions is diminished in this class of ligands with respect to the



**Figure 10.** The pyrazolo[3,4-c]quinolin-3-one ligand CGS-9896 (dotted line), diazepam (thick line), and pyridodiindole (thin line) fitted to a schematic representation of the inclusive pharmacophore model for the BzR. The sites H<sub>1</sub> and H<sub>2</sub> designate hydrogen-bond donor sites on the receptor protein while A<sub>2</sub> represents a hydrogen-bond acceptor site. Interaction with the lipophilic pocket L<sub>1</sub>, as well as with H<sub>1</sub> and A<sub>2</sub> is required for potent inverse agonist activity. Agonist activity requires interaction with H<sub>1</sub>, H<sub>2</sub>, L<sub>1</sub>, L<sub>2</sub>, and/or L<sub>3</sub>. Receptor descriptors S<sub>1</sub>, S<sub>2</sub>, and S<sub>3</sub> are regions of negative steric repulsion.

binding of full agonists. Receptor site selectivity of course plays a critical role in this behavior.

Clinically, there is a need for partial agonists which exhibit anxiolytic/anticonvulsant activity but are devoid of the ataxic/muscle relaxant effects associated with full agonist ligands. As stated earlier, there are only a few  $\beta$ -carboline ligands reported which elicit full agonist activity, and these compounds are the most challenging to model since the multiple rotamers produced by the substituents at positions 3, 4, 5, and 6 present difficulties.<sup>44</sup> In addition, the modeling studies suggested that  $\beta$ -carboline full agonists completely occupied lipophilic pockets L<sub>2</sub> and L<sub>3</sub>.<sup>44</sup> On the basis of modeling and the CGS 9895/9896 series it was postulated by Diaz-Araujo,<sup>191</sup> therefore, that partial occupation of L<sub>3</sub> may result



**Figure 11.** Superposition of the inclusive volumes of inverse agonists (Figure 7) and agonists (Figure 9). The agonist pharmacophore is depicted in gray while that of inverse agonists is the darker region.

in a partial agonist response. As a result of this, the synthesis<sup>190</sup> of a new anxiolytic/anticonvulsant, 6-*n*-propoxy-4-(methoxymethyl)- $\beta$ -carboline-3-carboxylic acid ethyl ester (6-PBC, **211**), which bound to the BzR with an IC<sub>50</sub> value of 8.1 nM, was designed. This compound was found to be devoid of the undesired myorelaxant and ataxic effects normally found with full agonist ligands. The synthesis of this compound via the method of Neef et al.<sup>196</sup> is illustrated in Scheme 19.

The 5-(benzyloxy)indole (**203**) was treated with ammonium formate in the presence of a palladium catalyst to provide the 5-hydroxyindole (**204**) in 92% yield. The indole **204** was alkylated with propyl iodide and potassium carbonate in acetone at reflux to furnish 3-(*n*-propoxy)indole (**205**) in excellent yield. A Michael reaction between nitro ester **206** and indole **205** in toluene at reflux yielded the 3,5-disubstituted indole **207**. Reduction of this compound using Raney nickel in ethanol under hydrogen at atmospheric pressure yielded the desired amino compound **209**. The Pictet–Spengler condensation of glyoxylic acid and **209** in ethyl acetate at pH = 4, followed by decarboxylation and subsequent oxidation furnished 6-PBC (**211**). This same procedure had been employed earlier by Neef et al.<sup>196</sup> to prepare the full agonist ZK 93423 (**212**).

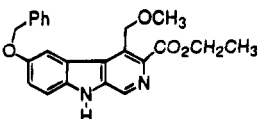
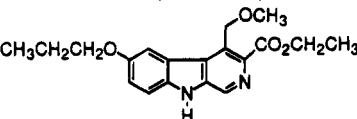
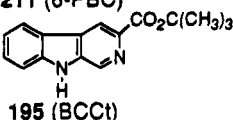
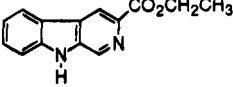
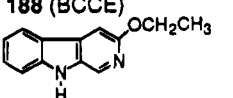
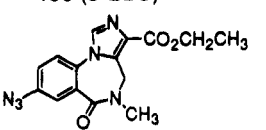
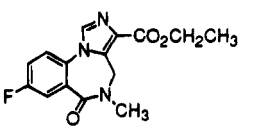
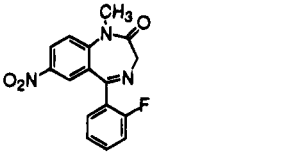
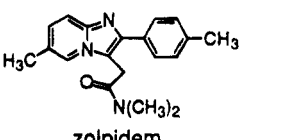
The synthesis and biological activity of **211** has important implications for the development of  $\beta$ -carbolines as partial agonists and anxiolytic agents. The proper lipophilic substituent at position 4 of  $\beta$ -carbolines directs the ligand into the agonist pharmacophore, the alignment of which is clearly different from that for inverse agonists. This new anxiolytic agent exhibited anticonvulsant/anxiolytic activity, but was devoid of the muscle relaxant/ataxic effects often associated with the classical 1,4-benzodiazepines.<sup>190</sup> More importantly, **211** actually antagonized the myorelaxant actions of diazepam, one of the most widely prescribed benzodiazepines currently in use. The design, synthesis, and pharmacological actions of **211**, based on computer-assisted analysis of the agonist pharmacophore, support the validity of this model.

### E. Benzodiazepine Receptor Subtype Selectivity

Recently, Zhang et al.<sup>44</sup> have shown with pyrazoloquinolines and imidazobenzodiazepines that the pharmacophoric descriptors H<sub>1</sub>, H<sub>2</sub>, and L<sub>1</sub> appear to be common to all six recently cloned [ $\omega$ 1,  $\omega$ 1,  $\omega$ 3,  $\omega$ 4,  $\omega$ 5,  $\omega$ 6 (DI)] benzodiazepine receptor sites in agreement with Doble and Martin.<sup>154</sup> The difference between these six major subsites is felt to originate from the size of (or interactions with) the lipophilic pockets, two of which have been designated L<sub>2</sub> and L<sub>3</sub> (see Figures 6–11). Molecular modeling studies using  $\omega$ 1 and  $\omega$ 5 selective ligands suggest that the lipophilic pocket (L<sub>2</sub>) in the  $\omega$ 5 site is larger than the analogous pocket in the Bz<sub>1</sub>( $\omega$ 1) site. Moreover, Zhang et al. have shown via chemical synthesis and molecular modeling that the lipophilic pocket designated L<sub>3</sub> in the DI ( $\omega$ 6) site is much smaller or nonexistent when compared to the same pocket in the  $\omega$ 1,  $\omega$ 2,  $\omega$ 3, and  $\omega$ 5 sites.<sup>44,197</sup> With these six cloned subtypes (McKernan et al.<sup>145</sup>) it is now possible to pursue the synthesis of ligands selective for one subsite over the other and to correlate the pharmacology with an interaction at that specific subsite. For the purposes of this review, illustrated in Table 12 are some selective  $\beta$ -carbolines, as well as imidazobenzodiazepines (Ro 15-1788 and Ro 15-4513) and their corresponding subsite selectivities.

The receptor subsite binding data for the full agonist  $\beta$ -carboline ZK 93423 (**212**) is given in Table 12. Clearly, there is no selectivity demonstrated between the  $\alpha$ 1,  $\alpha$ 2,  $\alpha$ 3, and  $\alpha$ 5 BzR subsites for this full agonist. The  $\alpha$ 4 site is felt to be very similar to the  $\alpha$ 6 site and is not discussed here.<sup>145</sup> The binding affinity of ZK 93423 at  $\alpha$ 6, however, is very poor ( $K_i > 1000$  nM). As discussed above, modeling studies suggest this is due to the lack of the lipophilic pocket L<sub>3</sub> at the  $\alpha$ 6 (DI) site, it is this pocket which would interact with the 6-benzyloxy substituent.<sup>191</sup> This same trend is observed with respect to the affinity of 6-PBC (**211**) at the  $\alpha$ 6 site ( $K_i = 1343$  nM). Presumably, the 6-propyloxy substituent cannot bind at the diazepam insensitive site [ $\alpha$ 6 $\beta$ 2 $\gamma$ 2, ( $\omega$ 6)] for this group would need to occupy L<sub>3</sub>. More interest-

Table 12. Receptor Subsite Selectivity Data for Selected BzR Ligands (Nanomolar)

Compound	$\alpha_1$	$\alpha_2$	$\alpha_3$	$\alpha_5$	$\alpha_6$
 212 (ZK 93423)	4.1	4.2	6	4.5	> 1000
 211 (6-PBC)	0.49	1.21	2.2	2.39	1343
 195 (BCct)	0.72	15	18.9	110.8	>5000
 188 (BCCE)	1.2	4.9	5.7	26.8	2700
 193 (3-EBC)	6.43	25.1	ND	826	>1000
 Ro15-4513	3.3	2.6	2.5	0.26	3.8
 Ro15-1788	0.8	0.9	1.05	0.6	148
 flunitrazepam	2.2	2.5	4.5	2.1	>2000
 zolpidem	26.7	156	383	>10,000	>10,000

ingly, 6-PBC (**211**) which is a partial agonist devoid of muscle relaxant activity, binds with a greater selectivity for Bz<sub>I</sub> ( $\omega_1 = \alpha_1$ ) receptor sites. This is in agreement with the earlier work of Beer and Lippa<sup>153,169</sup> which supports the hypothesis that muscle relaxant activity may arise from interactions at Bz<sub>II</sub> (Bz<sub>2</sub>, Bz<sub>3</sub>, and Bz<sub>5</sub>) sites.

The three  $\beta$ -carbolines  $\beta$ CCt (**195**),  $\beta$ CCE (**188**), and 3-EBC (**193**) demonstrated  $\alpha_1$  selectivity, in fact  $\beta$ CCt (**195**) is the most Bz<sub>I</sub> ( $\alpha_1$ ) selective antagonist (20 fold) reported to date. This now confirms why  $\beta$ CCt **195** antagonized the anxiolytic/anticonvulsant activity of diazepam but not the muscle relaxant or ataxic properties of this agent. Although more data is needed before an accurate correlation can be made, these results coupled with the biological profiles of these ligands, strongly suggest that receptor subsite

selectivity and pharmacological action are related. In agreement with Lippa and Beer it appears that Bz<sub>II</sub> receptors (Bz<sub>2</sub>, Bz<sub>3</sub>, and perhaps Bz<sub>5</sub>) are responsible for the muscle relaxant effects of diazepam rather than the Bz<sub>I</sub> sites.<sup>44</sup>

In conclusion, BzR pharmacology is complicated by the existence of multiple receptor isoforms. The inclusive pharmacophore model (Figures 6–11) for the "diazepam sensitive" (DS) BzR is based upon the weighted average of these receptor subtypes. Numerous  $\beta$ -carbolines, diindoles, 1,4-benzodiazepines, 1,4-benzazepines, triazolobenzodiazepines, imidazobenzodiazepines, and pyrazoloquinolines have been employed to model the inclusive pharmacophore at the benzodiazepine receptor site. In addition, receptor isoforms whose pharmacology resembles that of previously reported BzR-I (Bz<sub>1</sub>) and BzR-II (Bz<sub>2</sub>, Bz<sub>3</sub>,



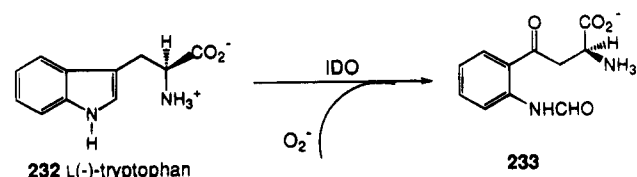
and Bz<sub>5</sub>) receptors have been expressed: the type-I BzR was constructed from an  $\alpha 1\beta 2\gamma 2$  combination of subunits as described previously while the type-II BzR were composed of  $\alpha 2\beta 2\gamma 2$ ,  $\alpha 3\beta 2\gamma 2$ , and  $\alpha 5\beta 2\gamma 2$  (zolpidem insensitive)<sup>156</sup> receptor isoforms.<sup>198,199</sup> As receptor-selective SAR data is accumulated for each of the cloned receptor isoforms, separate pharmacophore models will need to be developed for each. While these individual models may differ in detail from the inclusive model (inverse agonist/antagonist, and agonist) presented here, these models will nevertheless share many features in common with the inclusive model. The SAR and pharmacophore models which will result from future studies will permit the design of more selective ligands for each receptor subtype and may allow one to decouple the broad spectrum of effects exhibited by the current class of non-selective ligands. An outgrowth of this approach will be new therapeutic opportunities. A direct result of the BzR pharmacophore/receptor model has been the design and synthesis of a new partial agonist,<sup>190</sup> as well as partial inverse agonists.<sup>41</sup> The data obtained from these new ligands will be useful in refining the model and may ultimately lead to better drugs for treatment of anxiety disorders and of other maladies associated with neurotransmission in the CNS.

#### IV. Indolamine 2,3-Dioxygenase Inhibition in Inflammatory Diseases

Although 3-substituted  $\beta$ -carbolines have been shown to interact at benzodiazepine receptors, a number of related  $\beta$ -carbolines have proven to be effective noncompetitive inhibitors of the indolamine-2,3-dioxygenase (IDO) enzyme system.<sup>200</sup> Furthermore, the most interesting two agents studied to date (3-*n*-butyl- $\beta$ -carboline and 3-nitro- $\beta$ -carboline)<sup>40</sup> bind with poor affinity to BzR sites which decreases possible side effects in regard to IDO inhibition.

The metabolism of L-tryptophan (**232**) in mammals proceeds by the serotonin pathway,<sup>201,202</sup> the kynurenine pathway,<sup>201,202</sup> and a third proposed pathway<sup>203–205</sup> which involves the reversible conversion of L-tryptophan (**232**) into indole-3-pyruvic acid and subsequent transformation into kynurenic acid. The initial process in the kynurenine pathway involves the conversion of L-tryptophan **232** and other indoleamines into formylkynurenines by indolamine 2,3-dioxygenase (IDO) or tryptophan 2,3-dioxygenase (TDO). The conversion of tryptophan (**232**) into formylkynurenine (**233**) by IDO is depicted in Scheme 20. The affinity ( $K_m$ ) of L-tryptophan for IDO is 13  $\mu$ M.<sup>206</sup> Eventually the kynurenine pathway produces a variety of bioactive metabolites; the most significant of these are kynurenic acid, quinolinic acid (QA) and nicotinamide adenine dinucleotide (NAD). Quinolinic acid has been shown to be present in the mammalian brain, to be an agonist for the excitatory amino acid receptors of the *N*-methyl-D-aspartate (NMDA) receptor ion channel complex and to cause excitotoxic brain lesions when present in high concentrations.<sup>207–209</sup> This overstimulation of NMDA receptors can lead to a variety of neuropsychologic disorders.<sup>207,210–212</sup> Recently, Heyes and co-workers have shown that only macrophage-derived cells and certain liver cells synthesize labeled QA from labeled

Scheme 20



L-tryptophan following immune stimulation and that macrophages may provide a means for the production of large amounts of quinolinic acid in the brain following inflammation.<sup>213</sup> The activity of later stage enzymes which directly produce QA have been identified in brain tissue. Thus, metabolites from the kynurenine-pathway are manufactured by cells other than the brain and may enter the brain to be later metabolized to quinolinic acid *in vivo*. Kynurenic acid, however, is a noncompetitive agonist which interacts with the same receptor system that is affected by QA but can also cause seizures at high levels.<sup>213,214</sup>

Indolamine 2,3-dioxygenase is a 41 kD, monomeric heme-containing enzyme,<sup>215–220</sup> that utilizes superoxide to cleave the 2,3-double bond of indoleamines.<sup>221–224</sup> A mechanistic model of the conversion of indolamines to formylkynurenines by IDO, analogous to the mechanism of photolytically derived O<sub>2</sub> with tryptophan, has been proposed.<sup>225</sup> Unlike tryptophan 2,3-dioxygenase, an enzyme that is isofunctional to IDO but found only in the liver,<sup>226</sup> IDO is found in a variety of tissues, such as the brain, lung, and small intestine.<sup>227</sup> Furthermore, IDO is induced by interferon- $\gamma$ ,<sup>228–230</sup> consequently, IDO assumes a major role in immunological responses, as well as being a key factor for the *in vivo* production of bioactive metabolites which can be toxic at high levels.

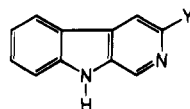
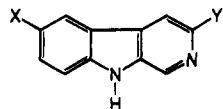
Upon severe infection, large quantities of interferon- $\gamma$  are produced which subsequently activate IDO to high levels.<sup>231,232</sup> This activation of IDO can lead to aberrant tryptophan metabolism and has been implicated in immunological diseases which affect the central nervous system. It has been demonstrated by Heyes and co-workers that quinolinic acid is produced from L-tryptophan by human macrophages and that the central nervous system could be affected by macrophages which have entered the CNS.<sup>233</sup> Aberrant tryptophan metabolism is characterized by the removal of tryptophan from the amino acid pool and the production of high levels of kynurenine, kynurenic acid and quinolinic acid.<sup>228–230</sup> This condition has been implicated in many inflammatory diseases, including acquired immune deficiency syndrome (AIDS),<sup>234–236</sup> hepatic encephalopathy,<sup>237</sup> polio virus,<sup>238</sup> and others.<sup>239–242</sup>

Norharman (**213**, 91% inhibition of rabbit small intestine IDO,  $K_i = 120 \mu$ M) was shown to function as an uncompetitive inhibitor for IDO,<sup>243</sup> but a noncompetitive mechanism of inhibition for this compound was later established on the basis of kinetic and spectroscopic studies.<sup>244</sup> Moreover, this later work demonstrated that norharman competed with oxygen for the heme-iron site. Several other examples of noncompetitive inhibitors have been discovered which are not  $\beta$ -carbolines.<sup>206,245</sup>

The results of this present study were intended to illustrate some of the structural requirements of the



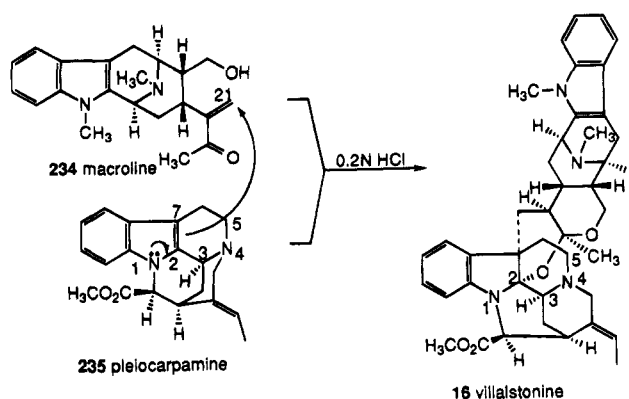
Table 13.  $\beta$ -Carbolines Evaluated for IDO Inhibition

		
213 Y=H	220 Y=CO <sub>2</sub> t-Bu	228 X=F; Y=CO <sub>2</sub> Me
214 Y=OEt	221 Y=C(O) <i>n</i> -Pr	229 X=Br; Y=CO <sub>2</sub> Me
215 Y=O <i>n</i> -Pr	222 Y=CH <sub>2</sub> OH	230 X=F; Y=CO <sub>2</sub> Et
216 Y=OCH <sub>2</sub> CH <sub>2</sub> OH	223 Y= <i>n</i> -Bu	231 X=NCS; Y=CO <sub>2</sub> Me
217 Y=CO <sub>2</sub> H	224 Y=NH <sub>2</sub>	
218 Y=CO <sub>2</sub> Me	225 Y=NCS	
219 Y=CO <sub>2</sub> <i>n</i> -Pr	226 Y=NO <sub>2</sub>	
	227 Y=OH	

noncompetitive binding site of human monocyte/macrophage IDO for the purpose of developing a potent inhibitor of this enzyme and are presented in Table 13. A series of 3-substituted  $\beta$ -carbolines **213**–**227** and four 3,6-disubstituted  $\beta$ -carbolines **228**–**231** were prepared and evaluated for *in vitro* human monocyte/macrophage indolamine 2,3-dioxygenase (IDO) inhibition.<sup>200</sup> Of these, 3-*n*-butyl- $\beta$ -carboline (**223**,  $K_i = 3.3 \mu\text{M}$ ) was found to be the most potent inhibitor of IDO reported from any source to date to these authors' knowledge. Substitution of position 6 of 3-(methoxycarbonyl)- $\beta$ -carboline (**218**,  $\beta\text{CCM}$ ) with fluoro or isocyanate substituents ( $K_i = 7.4$  and  $8.5 \mu\text{M}$ ) furnished derivatives **228** and **231** which were more potent inhibitors of IDO than the parent  $\beta\text{CCM}$  (**218**,  $K_i = 259 \mu\text{M}$ ). The 6-fluoro ethyl ester derivative **230** ( $K_i = 21.0 \mu\text{M}$ ) was found to be less potent than the methyl ester analog **228**. The inhibition was negated with substitution of position-6 of  $\beta\text{CCM}$  (**218**) with bromine. Moderate inhibition was produced by  $\beta$ -carboline-3-carboxylate (**217**,  $K_i = 40.6 \mu\text{M}$ ) and 3-nitro- $\beta$ -carboline (**226**,  $K_i = 37.5 \mu\text{M}$ ). When the 3-substituent of norharman (**213**) was replaced with a polar hydroxyl, amino, hydroxymethyl, or hydroxyethoxy group, only weak inhibition was observed. All of the  $\beta$ -carbolines which were investigated by kinetic methods were found to be noncompetitive or uncompetitive [in the cases of  $\beta$ -carboline-3-carboxylate (**217**) and 3-(hydroxymethyl)- $\beta$ -carboline (**222**)] inhibitors with respect to the substrate L-tryptophan (**232**). The natural products camalexin and brassilexin ( $K_i = 5.4 \mu\text{M}$ ) were also found to be noncompetitive inhibitors of IDO and provide important structural leads for further investigations. Both 3-*n*-butyl- $\beta$ -carboline (**223**) and 3-nitro- $\beta$ -carboline (**226**) are being evaluated by Heyes et al.<sup>238–241</sup> in *in vivo* models of IDO at the present time.

### V. Enantiospecific Total Synthesis of Indole Alkaloids

Bisindole alkaloids comprise a major portion of the macroline/sarpagine class of indole alkaloids and these natural products as a class have been the subject of several reviews, most notably those of Kutney, Lounasmaa, and Cordell.<sup>9,246,247</sup> Bisindoles in general, whether macroline related or not, are a class of alkaloids that present a significant synthetic challenge for the natural products chemist. The biomimetic synthesis of the macroline-related *Alstonia* bisindoles was pioneered by LeQuesne<sup>248</sup> and

Scheme 21<sup>248</sup>

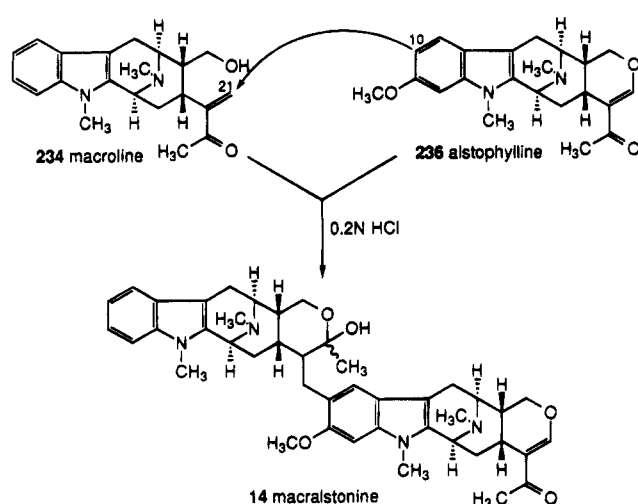
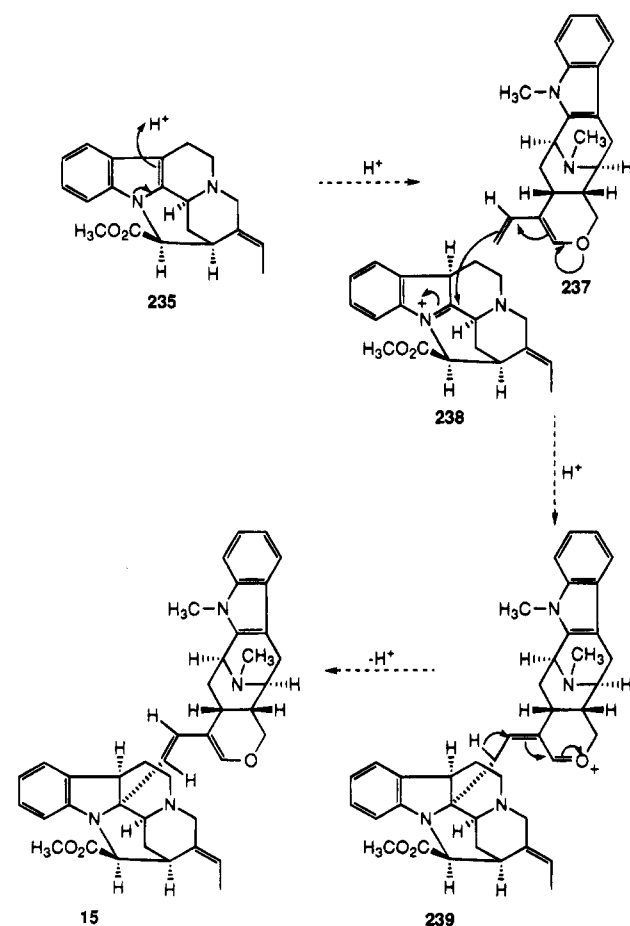
their isolation and structure determination have been reviewed.<sup>24</sup> As alluded to earlier in this review, three of these alkaloids, macralstonine (**14**), macrocarpamine (**15**), and villalstonine (**16**) possess interesting biological activity. As such, these three alkaloids are obvious synthetic targets and progress toward their total synthesis will be presented.

Many of the bisindole alkaloids from *Alstonia* species are comprised of two units directly related to the monomeric derivative macroline, while others originate from the condensation of macroline with another alkaloid. The biomimetic synthesis of *Alstonia* alkaloids involves the Michael addition of a monomeric alkaloid to C-21 of the  $\alpha, \beta$ -unsaturated enone moiety of (+)-macroline (**234**).<sup>248</sup> The biomimetic coupling of macroline (**234**) and pleiocarpamine (**235**) to form villalstonine (**16**) is represented in Scheme 21. The C-7 carbon atom of pleiocarpamine was activated by the lone pair of electrons on the indole nitrogen atom. This activation facilitated the Michael addition to the C-21 enone of macroline (**234**). The iminium ion which formed in this process was then attacked nucleophilically by the oxygen atom of the developing hemiacetal to provide villalstonine (**16**) in a stereospecific coupling process.<sup>248</sup> Since the total synthesis of (+)-macroline (**234**) has been completed and coupled to natural pleiocarpamine (**235**), a partial synthesis of villalstonine (**16**) has recently been completed.<sup>24,249</sup> Macralstonine (**14**), a ring A-oxygenated bisindole, was formed biomimetically by a similar process; however, the Michael addition took place between the C-10 carbon atom of alstophylline (**236**) and C-21 of macroline. Again, hemiketal formation followed the Michael addition and macralstonine **14** resulted (Scheme 22).

The structure of macrocarpamine (**15**), composed of the subunits of pleiocarpamine (**235**) and an alstonerine derivative **237**, was reported in 1978 by Hesse et al.<sup>250,251</sup> A biomimetic coupling process between pleiocarpamine (**235**) and olefin **237** has been proposed by Hesse to account for the formation of **15** in *Alstonia* species. This process, recently confirmed by Gan in our laboratory, serves as the foundation for the total synthesis of **15** and is depicted in Scheme 23.

The total synthesis of these bisindoles in optically active form therefore requires the enantiospecific preparation of the following monomeric indole alkaloids: macroline, alstophylline, deoxydehydroalston-

## Scheme 22

Scheme 23<sup>250,251</sup>

erine, and pleiocarpamine. The first three of these alkaloids have very similar carbon skeletons; consequently, an ideal approach to these target bases might rest upon the multigram synthesis of a common, optically active intermediate which could be employed for the synthesis of many related natural products. This common intermediate would at the very least contain the requisite tetracyclic ring system which could be readily functionalized for further transformations. The (–)-tetracyclic ketone **240** (Figure 12) was synthesized in 1988 with these goals in mind<sup>45,54,106,252,253</sup> while the racemic compound had been prepared on kilogram scale in the late 1970s.<sup>254</sup>

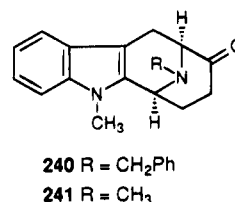
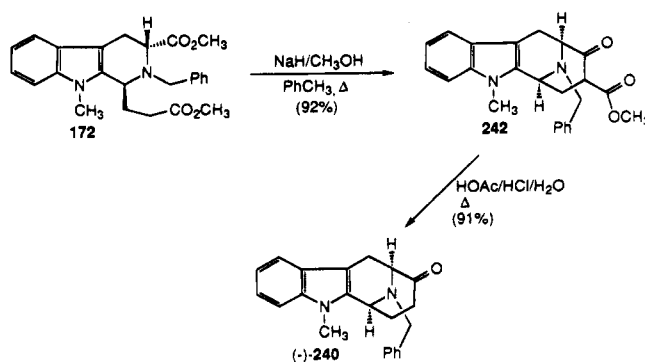


Figure 12. The optically active (–)-tetracyclic ketone.

### A. Enantiospecific Synthesis of the (–)-Tetracyclic Ketone

The synthesis of (±)-5-methyl-9-oxo-12-benzyl-6,7,8,9,10,11-hexahydro-6,10-imino-5*H*-cyclooct[*b*]indole (**240**) was first reported by Yoneda<sup>45</sup> and was improved by Soerens.<sup>254</sup> The enantiospecific preparation of the tetracyclic ketone **240** in optically active form was developed by Zhang<sup>106,252</sup> and is illustrated in Scheme 24. The synthesis of **240** began with D-(+)-tryptophan since it had been found earlier that the Pictet–Spengler reaction of aldehydes with *N*<sub>b</sub>-benzyl-substituted tryptophan methyl esters exhibited a strong preference for the enantiomerically pure *trans* diester. The 1,3-transfer of chirality from position 3 to position 1 of **172** would impart the correct configuration at C-1 for the synthesis of all the macroline, sarpagine, and ajmaline alkaloids. Hence, as depicted earlier in Scheme 13, methylation of D-(+)-tryptophan **168** was accomplished with sodium in liquid ammonia and methyl iodide in 92% yield. Fischer esterification of the methylated D-(+)-tryptophan gave *N*<sub>a</sub>-methyltryptophan methyl ester (**169**) in 87% yield. The benzylation of the *N*<sub>b</sub>-nitrogen function was carried out without racemization if care was taken to keep the imine intermediate cold during the reduction and to limit the time of reaction (3 h). The tryptophan methyl ester **169** was treated with benzaldehyde at 22 °C, and the imine which resulted was reduced with sodium borohydride (at –5 °C) to provide *N*<sub>a</sub>-methyl-*N*<sub>b</sub>-benzyltryptophan methyl ester **170** (greater than 98% ee) in 88% yield.<sup>252</sup> The Pictet–Spengler condensation of **170** with α-ketoglutaric acid in benzene/dioxane, accompanied by removal of water via a Dean–Stark trap, was followed by esterification in 1% methanolic HCl to afford the required *trans* diester **172** enantiospecifically. In the synthesis of the optically active tetracyclic ketone **240**, the Pictet–Spengler reaction was employed to set the stereochemistry at C-1 of the tetrahydro β-carboline ring system in stereospecific fashion. Yoneda<sup>45</sup> had earlier reported the

## Scheme 24



synthesis of a mixture of the racemic diesters **171** and **172** via the Pictet–Spengler reaction. The *cis* isomer **171** was originally reported to be the main constituent<sup>45</sup> but this was later corrected<sup>255</sup> to consist of a mixture of *trans*-**172** and *cis*-**171** diastereomers in a ratio of 5:4, respectively (89.1% yield). In the optically active series, Sakai<sup>255</sup> and co-workers extended the study of the Pictet–Spengler reaction to include the synthesis of (–)-tryptargine. Although the synthesis by Sakai et al. was in the *N<sub>a</sub>*-H series, use of the method developed by Ungemach<sup>60</sup> with an *N<sub>b</sub>*-benzyl group provided a remarkable *trans* to *cis* preference.<sup>255</sup> In the *N<sub>a</sub>*-methyl series, however, Zhang<sup>252</sup> observed a 72:28 diastereomeric ratio of *trans*-**172** to *cis*-**171** isomers when *N<sub>a</sub>*-methyl-*N<sub>b</sub>*-benzyltryptophan was treated with methyl 3-formylpropionate under nonacidic aprotic conditions (90% yield). More importantly, there was no racemization at C-3. Under the protic conditions involving  $\alpha$ -ketoglutaric acid, Zhang observed almost complete *trans* stereospecificity after esterification (1% methanolic HCl at reflux). The remaining small amount of *cis* isomer had been converted, with no loss of optical activity, into the *trans* diastereomer upon heating in 1% methanolic HCl. Hence, a sequence had been developed to provide the *trans* isomer in high enantiomeric purity even in the *N<sub>a</sub>*-methyl series in the absence of time-consuming separations.

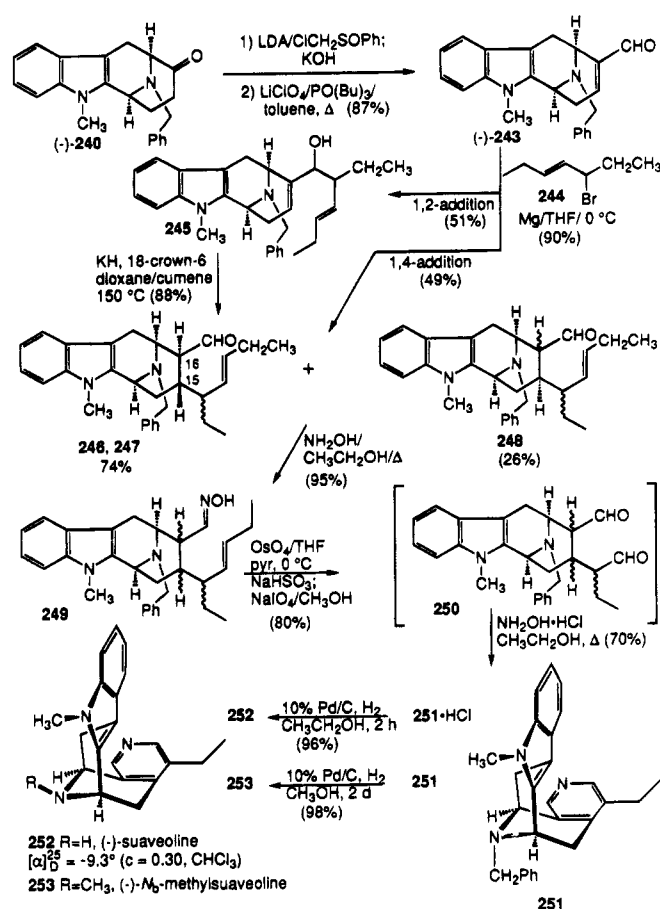
Dieckmann cyclization (Scheme 24) of the *trans* diester **172** afforded the  $\beta$ -ketoester **242** (92%).<sup>252,256</sup> After acid-mediated decarboxylation of **242**, the (–)-tetracyclic ketone **240** was obtained in 91% yield. The enantiomeric purity of this ketone (–)-**240** was shown to be greater than 98% ee by use of both <sup>1</sup>H NMR spectroscopy with the chiral shift reagent<sup>112</sup> tris[3-[(heptafluoropropyl)hydroxymethylene]-(+)-camphorato]europium(III) and by HPLC on a diastereomeric urea derivative of **240**.<sup>256</sup> The utility of this enantiospecific sequence rests on the fact that these reactions can be run on multigram scale to provide the (–)-tetracyclic ketone **240**, which can now be considered a readily available starting material for the synthesis of the macroline, sarpagine, and ajmaline related alkaloids. In addition, D-(+)-tryptophan and L-(–)-tryptophan are both readily available from commercial sources permitting entry into either antipode of the natural products for screening.

## B. (–)-Suaveoline

In 1972 Potier et al. isolated suaveoline **252** from the trunk bark of *Rauwolfia suaveolens* S. and reported the specific rotation  $[\alpha]_D^{25} = 0^\circ \pm 2^\circ$  for this base. The structure of **252** was elucidated on the basis of mass and proton spectroscopy, as well as a partial synthesis from ajmaline.<sup>257,258</sup> In 1989 Trudell reported the total synthesis of ( $\pm$ )-suaveoline (**252**)<sup>253</sup> and in 1992 Fu completed the first enantiospecific total synthesis of (–)-suaveoline.<sup>259,260</sup> A specific rotation of  $[\alpha]_D^{25} = -9.3^\circ$  ( $c = 0.30$ ,  $\text{CHCl}_3$ ) was determined for pure **252** in contrast to earlier reports.<sup>257,258</sup>

The total synthesis of (–)-suaveoline (Scheme 25) will be described beginning from (–)-*N<sub>a</sub>*-methyl-*N<sub>b</sub>*-benzyltetracyclic ketone (**240**), the synthesis of which was illustrated in Schemes 13 and 24. Conversion of the carbonyl function of (–)-**240** into the  $\alpha$ ,

## Scheme 25



$\beta$ -unsaturated aldehyde **243** via the spirooxirano-phenyl sulfoxide was accomplished in 87% yield by the method of Trudell<sup>253,261</sup> in the racemic series and of Zhang<sup>106</sup> in the (–)-*N<sub>b</sub>*-methyl series. The pseudosymmetric Grignard reagent, available from 5-bromo-3-heptene (**244**), was then added to the  $\alpha$ ,  $\beta$ -unsaturated aldehyde **243** at low temperature to provide the products of 1,2- (**245**) and 1,4-addition (**246–248**) in a combined yield of 90%. When this sequence was repeated at room temperature, only the product of 1,2-addition **245** was isolated and in high yield (88%). The alcohol **245** was purified and subjected to conditions that promote an oxyanion-Cope rearrangement (150 °C) to furnish the same C-15 functionalized tetracyclic systems **246**, **247**, and **248** obtained from the 1,4-addition in a ratio of 3:2. Although the stereoselectivity in the oxyanion-Cope process was only 3:2 with the preferred attack from the desired bottom face of the C(15)–C(16) olefinic bond, the mixture of aldehydes **246–248** could be employed in the synthesis of (–)-suaveoline (**252**). The 1,4-addition of **244** to **243** was unprecedented in these systems and provided the diastereomeric aldehydes **246** and **247** with the ajmaline configuration at C-15 and C-16 in a ratio of 3:1 **246,247/248**. Previous attempts<sup>115,254,262</sup> to effect 1,4-addition to  $\alpha$ ,  $\beta$ -unsaturated aldehyde **243** had proven unsuccessful; therefore, this example serves as the first case of such an addition in this hindered *N<sub>b</sub>*-benzylazabicyclo[3.3.1]nonane system. Since the configurations of the newly formed stereocenters in aldehydes **246–248** will eventually be destroyed, the aldehyde functions of the mixture of **246–248** were protected by treatment with hydroxylamine hydrochloride in ethanol at reflux. A diaster-

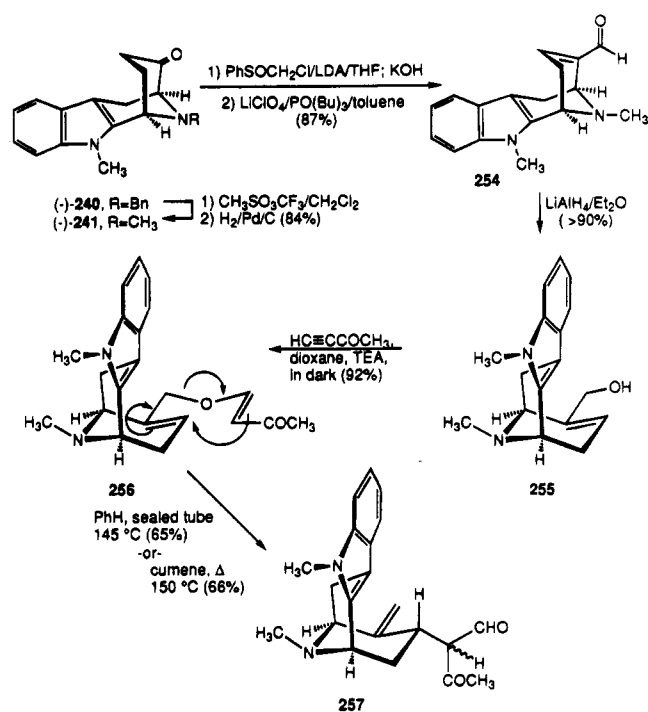
eomeric mixture of oximes represented by **249** was obtained in 95% yield. The mixture of oximes was osmolyated and subsequently hydrolyzed reductively with  $\text{NaHSO}_3$  to provide the desired diol which was subjected directly to the oxidative cleavage sequence ( $\text{NaIO}_4$ ). The desired dialdehyde **250** was obtained in 80% overall yield based on recovered starting oxime **249**. The mixture of dialdehydes **250** was cyclized *in situ* with hydroxylamine hydrochloride to provide  $N_b$ -benzylsuaveoline (**251**) in 70% yield. When **251** was subjected to the conditions of catalytic debenzylation with excess 10% Pd/C (1.5:1 w/w) and hydrogen in methanol, a 98% yield of  $(-)$ - $N_b$ -methylsuaveoline (**253**,  $[\alpha]_D^{25} = -89.5^\circ$ ,  $c = 0.35$ ,  $\text{CHCl}_3$ ) was realized in greater than 98% ee. Although the exact mechanism of the benzyl/methyl transformation is still not clear, it provided a simple manner in which to execute a benzyl/methyl transfer in the latter stages of the synthesis. This process can be employed in the preparation of a number of macroline/sarpagine/ajmaline alkaloids.<sup>24,263</sup> Catalytic debenzylation of the hydrochloride salt of  $(-)$ - $N_b$ -benzylsuaveoline (**251**) with 10% Pd/C (0.7:1.0 w/w) and hydrogen in ethanol provided a 96% yield of  $(-)$ -suaveoline (**252**).<sup>259,260</sup>

This sequence represents the first enantiospecific total synthesis of  $(-)$ -suaveoline and provides material upon which an accurate optical rotation could be obtained. Since the intermediates in this route are closely related to those previously reported in the synthesis of  $(\pm)$ -ajmaline, the strategy employed in the macroline series can be extended to alkaloids of the ajmaline family. Later in 1993, Bailey<sup>264</sup> described a formal synthesis of  $(-)$ -suaveoline (from L-tryptophan) which rested on the preparation of the optically active  $N_a$ -methyl- $N_b$ -benzyl tetracyclic ketone **240**. This  $(-)$ -ketone is identical with that reported earlier by Zhang<sup>252</sup> in 1988 and Fu<sup>259</sup> in 1992.

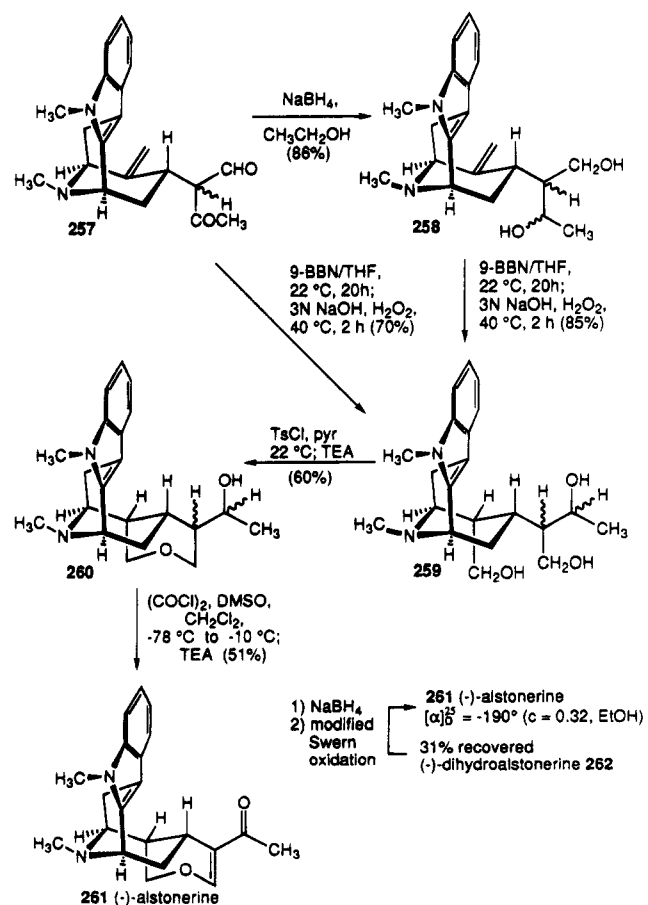
### C. $(-)$ -Alstonerine

Alstonerine (**261**) was first isolated from *Alstonia muelleriana* Domin by Elderfield and Gilman<sup>265,266</sup> and its structure was elucidated by LeQuesne et al.<sup>267</sup> The indole alkaloid alstonerine **261** is closely related to the oxindole alkaloid alstonisine (**294**).<sup>268</sup> Zhang<sup>106</sup> reported the synthesis of  $(-)$ -alstonerine in 1990 starting from the  $(-)$ -tetracyclic ketone **240** which was prepared earlier in enantiospecific fashion.<sup>252,256</sup> The  $N_b$ -benzyltetracyclic ketone **240** was methylated with methyl trifluoromethanesulfonate followed by catalytic debenzylation with Pd/C and hydrogen to afford the  $N_b$ -methyltetracyclic ketone **241** in high yield (Scheme 26). The ketone **241** was converted into the  $\alpha,\beta$ -unsaturated aldehyde **254** in 80% overall yield using conditions analogous to those reported by Trudell.<sup>253</sup> The  $\alpha,\beta$ -unsaturated aldehyde **254** was then transformed into the allylic alcohol **255** with lithium aluminum hydride in ether at  $-20^\circ\text{C}$ . Michael addition to 3-buten-2-one in the absence of light gave the desired enone **256** in excellent yield. The Claisen rearrangement (**256** to **257**) proceeded via the preferred chair transition state primarily from the bottom face of the double bond to afford the desired  $\beta$ -dicarbonyl compound **257** with a diaste-

Scheme 26



Scheme 27



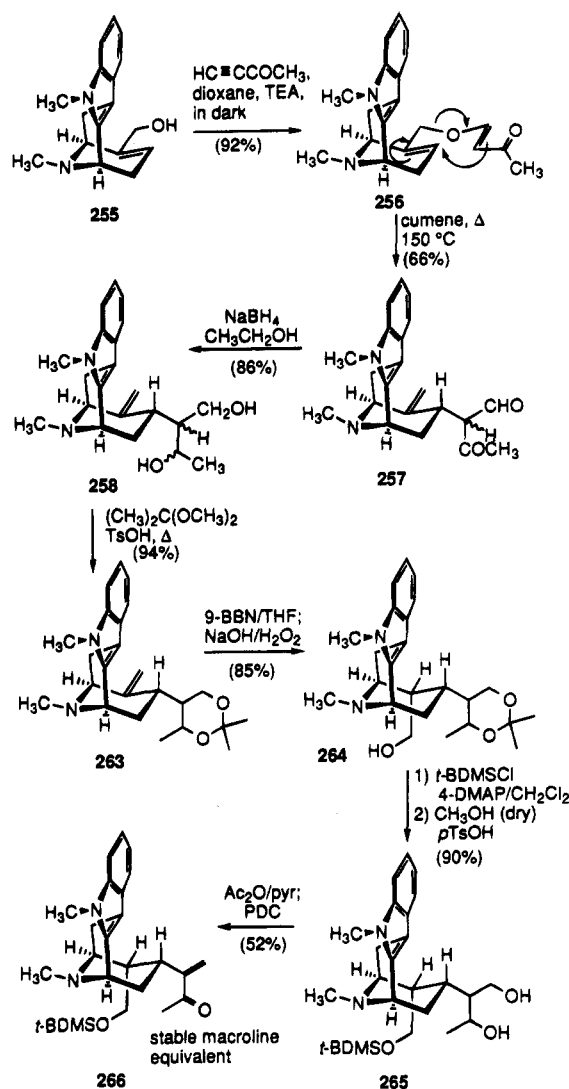
reoselectivity in cumene ( $150^\circ\text{C}$ ) of greater than 4:1 in 82% yield.<sup>106</sup> The  $\beta$ -dicarbonyl compound **257** was then reduced with sodium borohydride to the diol **258** as illustrated in Scheme 27. Hydroboration of the exocyclic methylene function of **258**, with an excess of 9-BBN occurred stereospecifically from the  $\beta$ -face of the double bond and after oxidative workup

provided the triol **259**. One equivalent of 9-BBN complexed to the  $N_b$ -nitrogen function which hindered attack from the bottom face of the double bond and resulted in exclusive hydroboration from the  $\beta$ -face of the exocyclic methylene function. Upon stirring with tosyl chloride (1 equiv) in pyridine followed by treatment with triethylamine, the triol **259** was regioselectively cyclized to the desired monol **260** in 60% yield, accompanied by recovered starting triol **259** (33%). Additional quantities of **260** could be obtained by subjecting the recovered triol **259** to the same tosylation process. The alcohol **260** underwent a modified Swern oxidation to provide (–)-alstonerine (**261**) in 51% yield, accompanied by dihydroalstonerine (**262**, 31%). A proposed mechanism for this transformation has been reported by Bi et al.<sup>24</sup> The dihydroalstonerine intermediate **262** could be recycled to provide additional quantities of **261** by sodium borohydride reduction, the monol of which was subjected to the conditions of the modified Swern oxidation. This procedure may provide a general method for the conversion of hydroxy-substituted tetrahydropyrans into enones which are commonly found in other *Alstonia* alkaloids such as alstophylline (**236**) and alstonisine (**294**). The enantiospecific synthesis of the tetracyclic ketone (–)-**240** coupled with the Claisen rearrangement (C-15) and the hydroboration process (C-16) provided a route of high diastereoselectivity for the enantiospecific synthesis of the macroline/sarpagine alkaloid, (–)-alstonerine **261**. Substitution of the 6-methoxy derivative of tryptophan alkyl ester (**172**) for the parent ester will provide a route to alstophylline (**236**), the monomeric unit required for the synthesis of macralstonine.

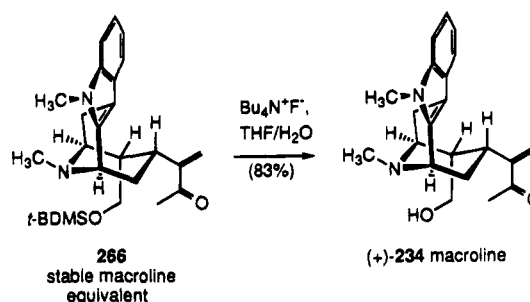
#### D. (+)-Macroline

The synthesis of (+)-macroline (**234**) has recently been completed (Schemes 28 and 29) in enantiospecific fashion starting with D-(+)-tryptophan.<sup>23,249</sup> The significance of this synthesis becomes apparent when one considers that >70 macroline related alkaloids have been isolated.<sup>24</sup> Macroline is not stable over long periods of time; therefore, the synthesis of the stable macroline equivalent **266** which can be employed for the synthesis of *Alstonia* bisindoles is presented in Scheme 28. The conversion of **266** into (+)-macroline (**234**) follows in Scheme 29. The tetracyclic ketone (–)-**240** was employed for the synthesis of (–)-alstonerine<sup>106</sup> and the required allylic alcohol **255** had been prepared by the route shown in Scheme 26. The synthesis of macroline **234**, as illustrated in Scheme 28, began with the intermediate allylic alcohol **255**. Michael addition of **255** to 3-butyne-2-one in the absence of light provided the enone **256** in excellent yield. The Claisen rearrangement of **256** took place stereoselectively from the desired  $\alpha$ -face (4:1) in cumene at 150 °C to afford the same dicarbonyl compound **257** employed for the synthesis of (–)-alstonerine (**261**). Although the stereoselectivity was reported to be 4:1, it may be much higher because the three byproducts formed in this pericyclic event were inseparable, rendering their structure determination difficult at this juncture. Reduction of dicarbonyl compound **257** produced the diol **258**. The diol **258** was converted into

Scheme 28

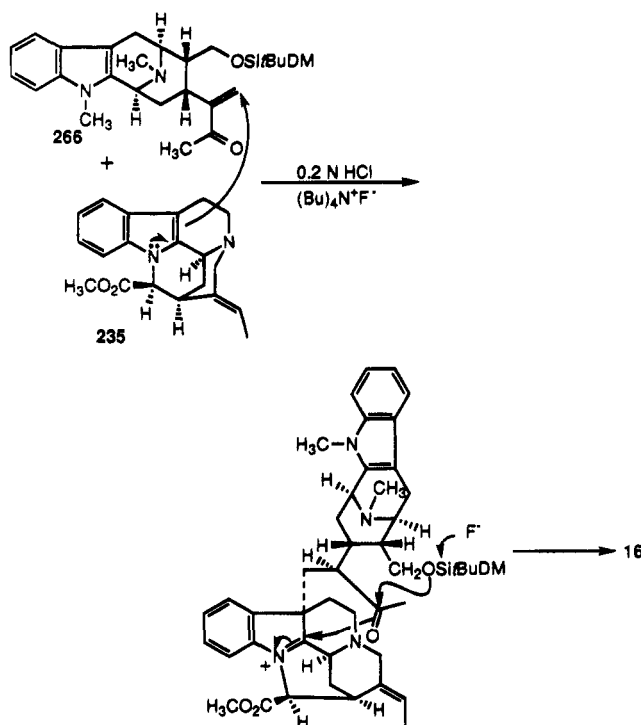


Scheme 29



the triol previously described in Scheme 27; however, attempts to utilize the triol **259** for the synthesis of macroline proved impractical. Consequently, **258** was protected as the acetonide **263** before the hydroboration/oxidation process with 9-BBN/OH<sup>−</sup>/H<sub>2</sub>O<sub>2</sub> was carried out. Hydroboration of **263** occurred exclusively from the  $\beta$ -face of the C(16–17) olefinic bond, as planned, to provide the desired primary alcohol **264**. The primary hydroxyl moiety of **264** was converted into the *tert*-butyldimethylsilyl ether, after which the acetonide was selectively removed upon stirring this compound with *p*-toluenesulfonic acid in dry methanol under argon. Acetic anhydride was then used to protect the primary alcohol in diol **265**, and the acetate which resulted served as the desired

Scheme 30



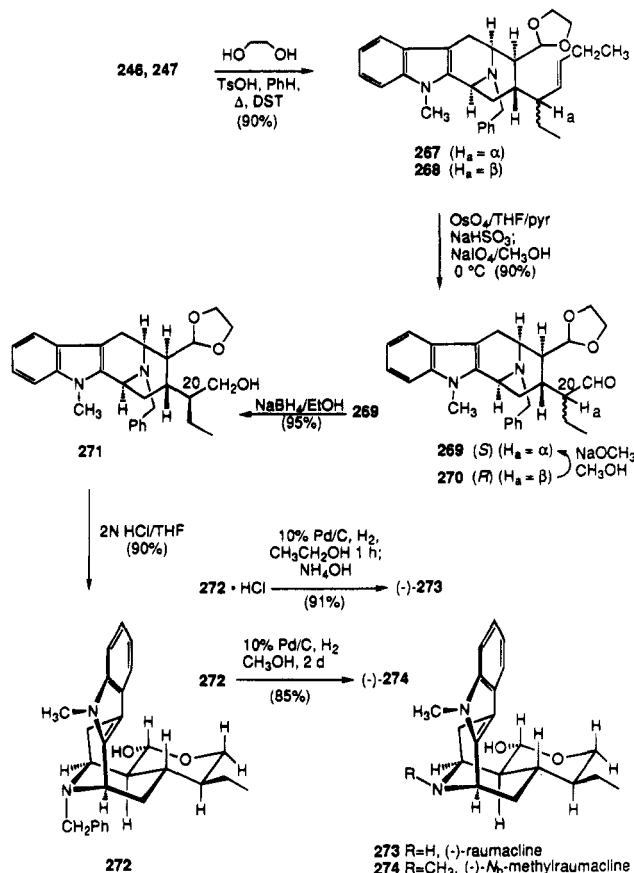
leaving group. This protection of the primary hydroxyl group of **265** was followed by oxidation with pyridinium dichromate (PDC) to provide the stable macroline derivative **266** in a one-pot process. After the oxidation of the secondary alcohol of **265** to the corresponding ketone had occurred, the pyridine present in solution promoted the loss of the  $\beta$ -ketoacetate function to provide the stable macroline enone **266**. When **266** was stirred in THF with tetrabutylammonium fluoride, (+)-macrolone (**234**) was obtained (Scheme 29). Macrolone is known to cyclize to dihydroalstonerine **262** when exposed to base; therefore, the synthesis of the macroline equivalent **266** was designed to facilitate its use in the synthesis of bisindole alkaloids.

The (-)-tetracyclic ketone **240** has been converted into the macroline equivalent **266** by a series of stereocontrolled steps as described above. When the synthetic macroline equivalent **266** was stirred with plant-derived pleiocarpamine in 0.2 N aqueous hydrochloric acid in the presence of fluoride ion, vilalstonine (**16**) was the only observable product, as illustrated in Scheme 30.<sup>250</sup>

## E. (-)-Raumacline

Since the complete structure of (+)-ajmaline has been well documented<sup>269–271</sup> and confirmed by X-ray crystallography<sup>272</sup> this commercially available alkaloid has been widely used as a starting material for the preparation of other alkaloids. Stöckigt and Sakai, et al.<sup>273–275</sup> isolated several new alkaloids, termed the raumaclines, from cell cultures of *Rauwolfia serpentina* Benth after feeding experiments with ajmaline. Raumacline (**273**) and  $N_b$ -methylraumacline (**274**) were first detected as products of these feeding experiments in 1990. The structures of these alkaloids were elucidated by spectroscopic methods and partial synthesis from ajmaline.<sup>275</sup> Later, Stöck-

Scheme 31



igt et al. isolated<sup>273,274</sup> four more raumacline alkaloids from *Rauwolfia serpentina* Benth cells cultivated in the presence of (+)-ajmaline.

Soon after the isolation of this new class of alkaloids, Fu<sup>259</sup> completed the total synthesis of (-)-raumacline (**273**) and (-)- $N_b$ -methylraumacline (**274**). The enantiospecific nature of the total synthesis of **273** and **274** is important for several reasons. The relationship between raumacline **273** and ajmaline has now been established chemically by Sakai *et al.*; therefore, entry into other ajmaline alkaloids including the total synthesis of the unnatural antipode (-)-ajmaline can be envisaged. The synthetic route executed by Fu employed the same oxyanion-Cope rearrangement developed for the total synthesis of (-)-suaveoline (**252**)<sup>259</sup> and  $N_b$ -methylsuaveoline (**253**).

The formyl group of the mixture of aldehydes **246** and **247** was protected as the ethylene acetal, followed by oxidative cleavage of the double bond to provide two epimeric aldehydes **269** and **270** in excellent yield (Scheme 31) which were separated by flash chromatography. Aldehyde **269** possessed the desired chirality (*S*) at C-20 for the synthesis of (-)-raumacline (**273**) and  $N_b$ -methylraumacline (**274**). For this reason the epimer **270** was treated with base and converted into an equilibrium mixture of **269** and **270** (1:1), which was easily separated by flash chromatography on silica gel. The combined (-)-(*S*)-aldehyde **269** was then reduced to the alcohol **271**. This was followed by deprotection of the aldehyde function and cyclization under acidic conditions to provide (-)- $N_b$ -benzyltraumacline (**272**). It is worth noting in this last sequence that the formation of **272** from aldehyde **269** was stereospecific. Catalytic



debenzylation of the hydrochloride salt of (–)-**272** in ethanol furnished (–)-raumacline (**273**) in 91% yield. When the base (–)-**273** was subjected to catalytic debenzylation with excess Pd/C and hydrogen in methanol, an 85% yield of (–)-*N*<sub>b</sub>-methylraumacline (**274**) was realized. These two syntheses of (–)-**273** and (–)-**274** and that of (–)-suaveoline (**252**) represent the first enantiospecific synthesis of members of the ajmaline family of indole alkaloids and demonstrate that the strategy employed for the preparation of the macroline related sarpagine alkaloids can be extended to other families of indole alkaloids.

## VI. Enantiospecific Synthesis of 5-Methoxy-D-(+)- or L-(–)-tryptophan

Over the past several years the isolation of a number of C-10 ring A-oxygenated indole alkaloids in the macroline/sarpagine series have been reported (Figure 13)<sup>24,276</sup> including 18-hydroxylochnerine (**275**), spogatrine (**276**), lochneram (**277**), 10-methoxyvellosimine (**278**), sarpagine (**279**), lochnerine (**280**), *N*<sub>a</sub>-methylsarpagine (**281**), neosarpagine (**283**), verticillatine (**284**), and 19,20-dehydro-10-methoxytalcarpine (**285**). Interest in the synthesis of *N*<sub>a</sub>-methylsarpagine (**281**) as well as 19,20-dehydro-10-methoxytalcarpine (**285**) has prompted the need for a preparative synthesis of 5-alkoxy-D-(+)-tryptophans via a route which would also provide the L-(–)-enantiomers, if desired. Since, the enantiospecific synthesis of a number of *Alstonia* macroline/sarpagine alkaloids has been demonstrated<sup>24,276</sup> by employing the *trans* 1,3-transfer of chirality during the Pictet–Spengler condensation, the use of D-(+)-tryptophan would provide the natural indole alkaloid while the L-(–) enantiomer would furnish the unnatural antipode for biological screening. Recently, a general method for the enantiospecific synthesis of 5-methoxy-D-(+)-tryptophan (**293**) or the L-(–) optical anti-

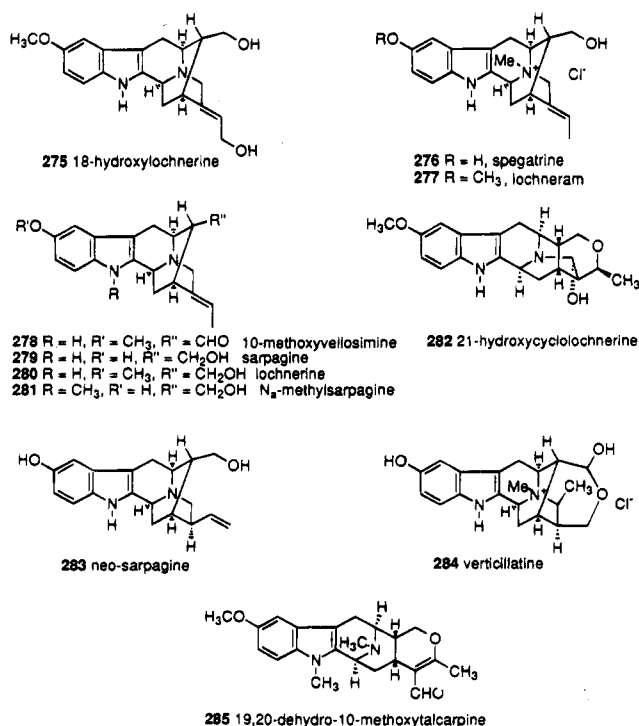
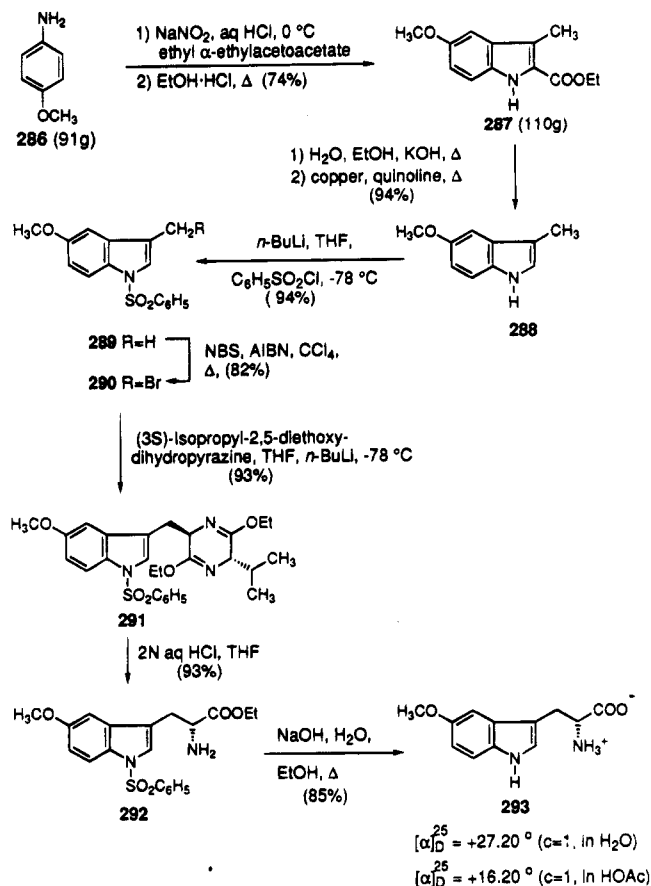


Figure 13. Ring A-oxygenated indole alkaloids.

## Scheme 32



pode was completed by Zhang<sup>277</sup> and is illustrated in Scheme 32.

In regard to the total synthesis of alstophylline and macralstonine, the preparation of 1-(phenylsulfonyl)-6-methoxy-D-(+)-tryptophan ethyl ester had already been carried out in our laboratories<sup>278</sup> by employing the Schöllkopf chiral auxiliary.<sup>279</sup> The Schöllkopf chiral auxiliary had been chosen for the desired D-(+)-tryptophan would be available from L-valine while the L-(–) enantiomer would originate from D-valine. The success of this sequence rested upon the ability to scale up the first few steps to multihundred gram scale. For this reason, the well-known Fischer indole cyclization,<sup>102</sup> via the thermally mediated [3,3]sigmatropic rearrangement, was chosen as the method by which to generate large quantities of 5-methoxy-3-methylindole (**288**).

Ethyl 5-methoxy-3-methylindole-2-carboxylate (**287**) was prepared on a large scale from *p*-anisidine and ethyl α-ethylacetoacetate by the Fischer indole cyclization via a Japp–Klingman azo-ester intermediate (Scheme 32).<sup>280</sup> This process has been fully explored by Abramovitch and Shapiro as well as reviewed.<sup>95,281</sup> Alkaline hydrolysis of ester **287** and subsequent copper/quinoline-mediated decarboxylation of the carboxylic acid furnished the 5-methoxy-3-methylindole (**288**) in excellent yield. Care must be exercised on decarboxylation of the corresponding acid on a large scale. The best yields were obtained when the carboxylic acid was carefully dried and the decarboxylation was executed at reflux in a well-stirred minimum amount of distilled quinoline (1.5–2 equiv of quinoline with respect to the carboxylic acid). Only a catalytic amount of copper powder was

required to ensure yields of **288** in excess of 90%.

In order to employ the Schöllkopf chiral auxiliary in the indole series, protection/deactivation of the indole N(H) group was required. This was accomplished by treatment of indole **288** with *n*-BuLi and benzenesulfonyl chloride in THF at  $-78\text{ }^{\circ}\text{C}$  to provide sulfonamide **289** in 94% yield.<sup>279,282</sup> The protected indole **289** was then treated with NBS<sup>283</sup> under free radical conditions (AIBN) to afford the protected 3-(bromomethyl)indole **290** in excellent yield. The alkylation of the anion of the Schöllkopf chiral auxiliary derived from L-valine was performed under conditions analogous to those described in the literature<sup>278,282</sup> and afforded the protected tryptophan derivative **291** stereospecifically. The substituted pyrazine group was hydrolyzed under acidic conditions (aqueous 2 N HCl, THF) to provide the desired 1-(phenylsulfonyl)-5-methoxytryptophan ethyl ester (**292**) as the hydrochloride salt. Alkaline hydrolysis of both the 1-phenylsulfonyl protecting group and the ethyl ester moiety furnished 5-methoxy-D-(+)-tryptophan (**293**). Racemization of the chiral center of the amino acid **293** was not observed under the conditions of hydrolysis (8 h) for even prolonged heating under equivalent reaction conditions ( $\text{OH}^-$ ) returned the same amino acid **293** with the identical optical rotation observed on hydrolysis of **293** for only 8 h.

The applicability of this route rests on the ease of execution of each step,<sup>277</sup> moreover the optically active 5-methoxytryptophan was obtained in only five steps from 5-methoxy-3-methylindole (**288**). Recently we reported the enantiospecific synthesis of 6-methoxy-D-(+)-tryptophan<sup>278</sup> which was being employed for the total synthesis of macralstonine as stated. The recent work of Zhang<sup>277</sup> however can be employed to provide 6-methoxy-D-(+)-tryptophan via a much shorter route than the previous sequence.<sup>267,268</sup> In addition, the route by Zhang has been applied to *o*- and *m*-anisidine to provide 7-methoxy-3-methylindole and 6-methoxy-3-methylindole, respectively, in good yields. Attempts to convert these indoles into optically pure 7-methoxy- and 6-methoxytryptophans are currently underway. The sequence by Zhang represents the first synthetic entry into either 5-methoxy-D-(+)- or L(-)-tryptophan<sup>284,285</sup> and makes these materials available on multigram scale for total synthesis.

## VII. Oxindole Alkaloids

The first macroline-related oxindole alkaloid alstonisine (**294**), was isolated from *Alstonia muelleriana* Domin and reported by Elderfield and Gilman (Figure 14).<sup>265,266</sup> The structure of this alkaloid was reported by Nordman,<sup>286</sup> unfortunately an error in transposition of this to paper resulted in an incorrect representation of the structure of alstonisine.<sup>286</sup> The absolute configuration of this base at C-3, C-5, C-15, and C-16 was later determined by Le Quesne et al.<sup>287</sup> when alstonisine (**294**) was biomimetically transformed into talpinine; however, direct confirmation of the stereochemistry at the spirocenter (C-7) has not been reported to date. The establishment of the stereochemistry at the spirocenter C-7 therefore constitutes one of the principal reasons for interest in the enantiospecific total synthesis of alstonisine

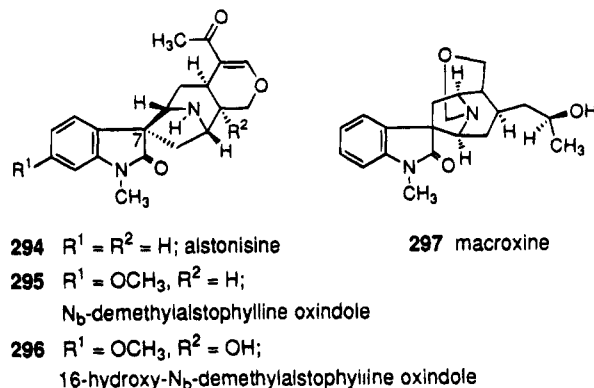


Figure 14. Macroline-related oxindoles.

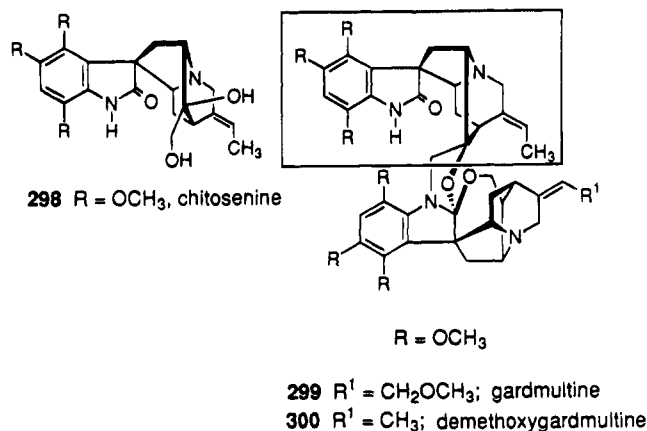


Figure 15. Oxindoles isolated from *Gardneria multiflora* Makino.

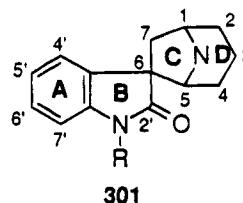
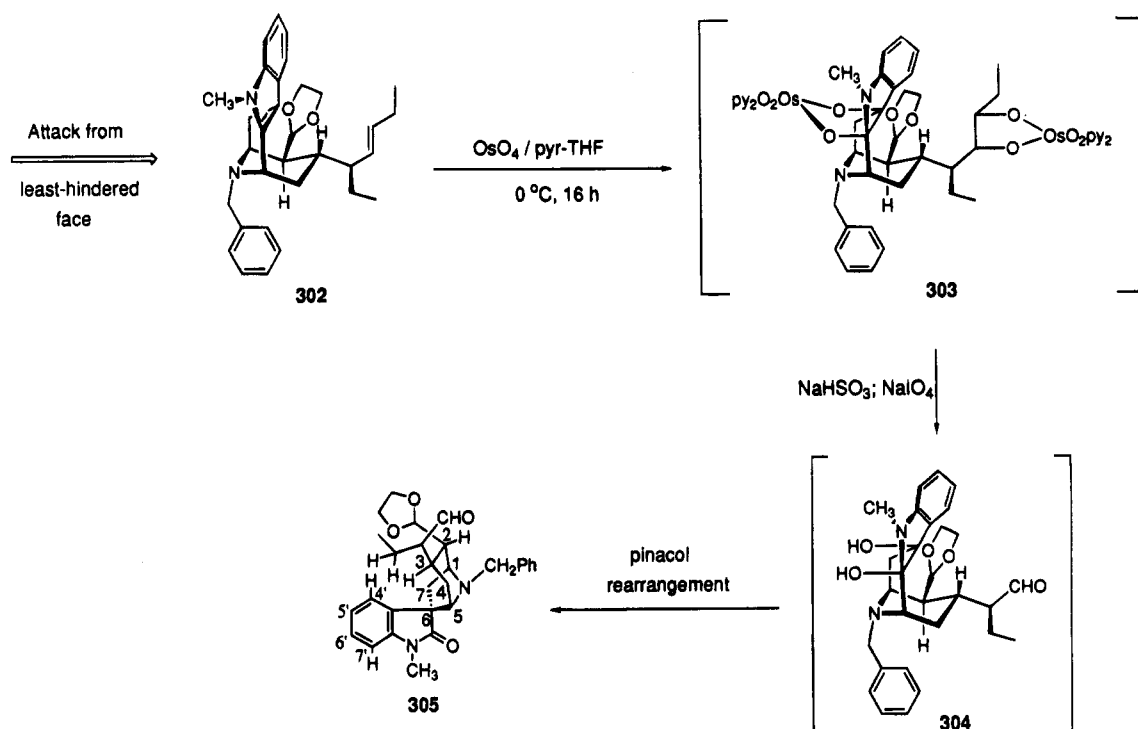


Figure 16. The tetracyclic oxindole ring system.

(**294**). Several other macroline-related oxindole alkaloids have recently been isolated from *Alstonia macrophylla* Wall including *N<sub>b</sub>*-demethylalstophylline oxindole (**295**),<sup>288</sup> 16-hydroxy-*N<sub>b</sub>*-demethylalstophylline oxindole (**296**),<sup>289</sup> and macroxine (**297**, Figure 14).<sup>290</sup> The configuration of oxindole alkaloids **295** and **296** at C-7 has been determined by NOE spectroscopic experiments.<sup>288,289</sup> The biological importance of these bases is unclear at this time due to limited amounts of material available for study.

The monomeric base chitosenine (**298**)<sup>291</sup> as well as the bisindoles gardmultine (**299**) and desmethoxygardmultine (**300**)<sup>292</sup> isolated from *Gardneria multiflora* Makino have been shown to exhibit short-lived inhibitory activity *in vivo* of ganglionic transmission in both rats and rabbits (Figure 15).<sup>293</sup> The configuration of the spirocyclic carbon (C-7 in alstonisine **294**) of these oxindoles, however, is opposite to that found in the *Alstonia* oxindoles **294–297**. The isolation of alkaloids **298–300** in addition to oxindoles **294–297** suggests that alkaloids which contain the substructure **301** (Figure 16) may be more prevalent in plants than previously realized. The development of a stereochemically complementary method into

Scheme 33



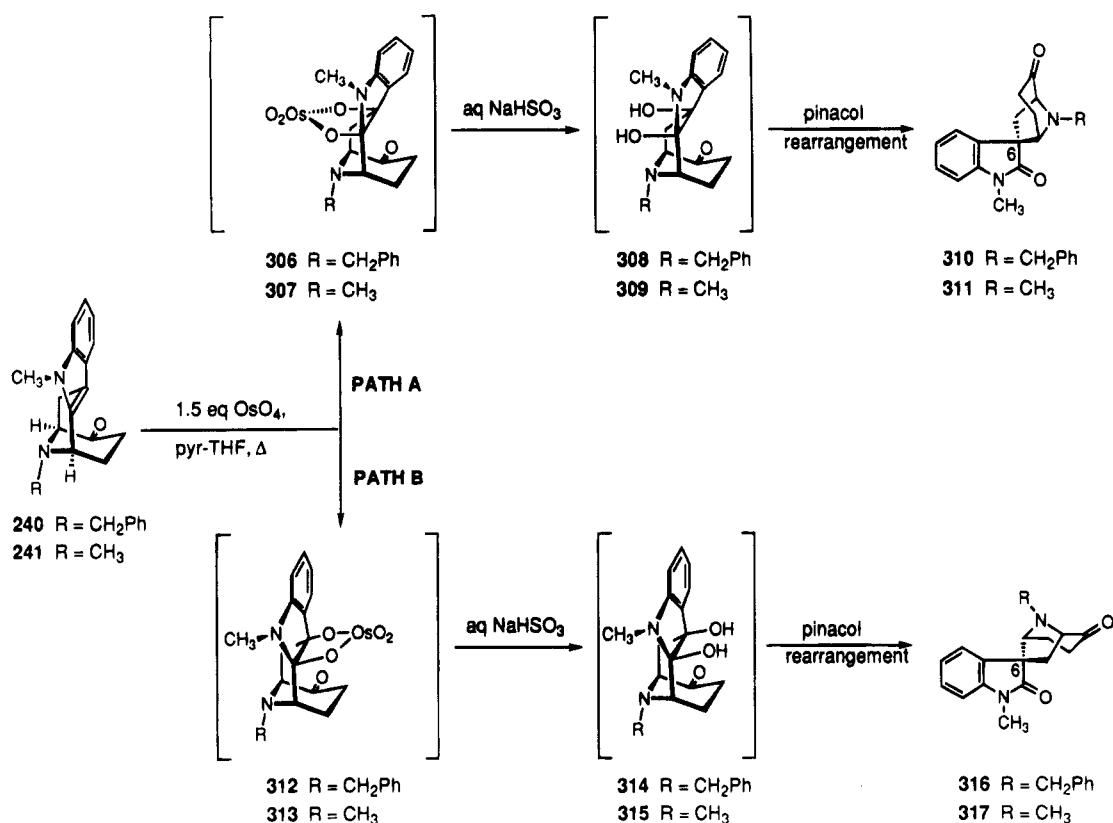
oxindoles related to **301** of either chirality at the spirocyclic carbon would provide material for further the study of these unique oxygenated bases.

Recently Peterson<sup>268</sup> has developed a method to convert  $N_a$ -methyltetracyclic ketones into their corresponding oxindoles with a high degree of diastereoselectivity. During the synthesis of (–)-raumacline (**273**),<sup>259</sup> Fu discovered that the treatment of the synthetic  $N_b$ -benzyltetracyclic monoketal **302** with osmium tetroxide in pyridine followed by periodate oxidation provided the oxindole **305**, as illustrated in Scheme 33. This conversion occurred with complete diastereoselectivity with a configuration identical to that proposed for alstonisine **294**. Esmond and Le Quesne had also observed a similar formation of an oxindole during dihydroxylation of a key intermediate during their biomimetic synthesis of macroline.<sup>294</sup> Attack of the osmium tetroxide was proposed to occur from the less hindered convex face of the indole 2,3-double bond to furnish an intermediate bisosmate ester **303**, as illustrated in Scheme 33. It was believed that the apically positioned acetal group effectively blocked the concave face of the double bond to attack by the  $\text{OsO}_4$ /pyridine reagent. Conversion of intermediate **303** into the diol-aldehyde **304** and subsequent pinacol rearrangement provided oxindole **305**. The recently developed method of Peterson employing the Sharpless osmylation sequence converts  $N_a$ -methyltetracyclic ketone<sup>252</sup> analogs into their corresponding oxindoles with a high degree of diastereoselectivity. These substrates are devoid of a group other than hydrogen situated at either the equatorial position at C-16 or the axial position at C-15 to direct the stereoselectivity. More importantly, this approach provides entry into either spirocyclic oxindole, diastereomeric at C-6 (see Figure 16), from the same (–)-antipode of the tetracyclic ketone **240**.

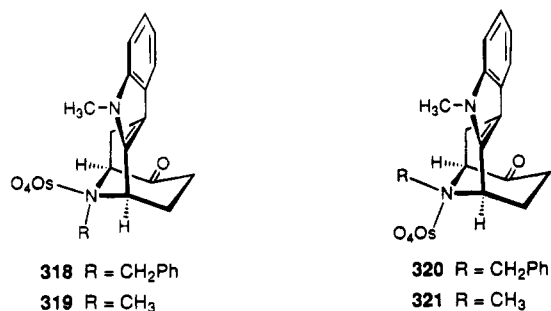
Optically active (–)- $N_b$ -benzyltetracyclic ketone **240** was treated with osmium tetroxide in THF at room temperature, followed by reductive workup with aqueous  $\text{NaHSO}_3$  and flash chromatography, the oxindoles **310** and **316** were produced in a 91:9 ratio in 42% yield (Scheme 34; Table 14). It is believed that the osmium tetroxide first complexed with the piperidine nitrogen atom of ketone **240** to furnish complex **318** (Figure 17). This complexation was presumably favored due to the axial preference (with respect to the D ring) of the benzyl group. Single-crystal X-ray analysis of an  $N_b$ -benzyl tetracyclic derivative indicated that the benzyl group rested in the axial position of the D ring in the crystal.<sup>295</sup> The concomitant complexation of osmium at the equatorial position (with respect to ring D) facilitated intramolecular attack of the osmium reagent to furnish osmate ester **322** (Figure 18) upon heating at reflux. If complexation occurred at the axial position (with respect to the D ring) to give complex **320**, intramolecular delivery of the osmium reagent would be unlikely. The osmate ester **322** was then reduced by sodium bisulfite, and the *cis*-diol **308** which resulted underwent a pinacol rearrangement to furnish oxindole **310** with a 10:1 overall diastereoselectivity. The configuration about the spirocyclic C-6 in oxindole **310** was found to be the same as in alstonisine **294** (C-7) by NMR spectroscopy.<sup>268</sup>

Further evidence for the advent of the complexation/intramolecular delivery of osmium tetroxide in the previous example was obtained by attempted conversion of  $N_b$ -benzoyl ketone **323** into  $N_b$ -benzoyloxindole **324** or its diastereomer (Scheme 35). Only the starting ketone **323** was recovered (95% recovery) from this sequence. Clearly, the benzoyl group of substrate **323** was approximately the same size as the benzyl group in ketone **240**. However, the lone pair of electrons of the piperidine nitrogen are

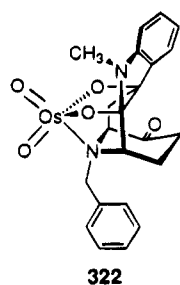
## Scheme 34



delocalized into the carbonyl group of the amide function and not readily available to coordinate with osmium tetroxide. This example demonstrated that neither the complexation of OsO<sub>4</sub> (and subsequent intramolecular oxidation of the indole 2,3-double bond) occurred nor uncomplexed OsO<sub>4</sub> reacted with substrate **323** even from the concave face of the indole double bond at room temperature. Evidently in these systems the OsO<sub>4</sub> was not reactive enough at room temperature to oxidize the indole double bond without previous ligation to a nitrogen atom.

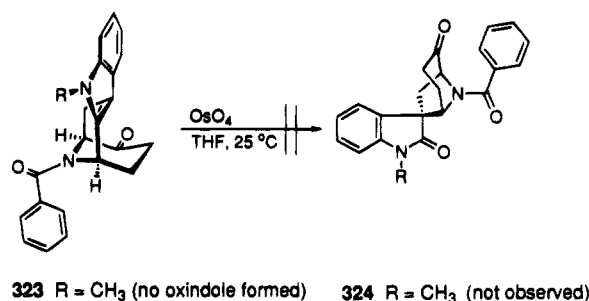


**Figure 17.** The complexation of osmium tetroxide to the piperidine nitrogen atom.



**Figure 18.** Intermediate osmate ester.

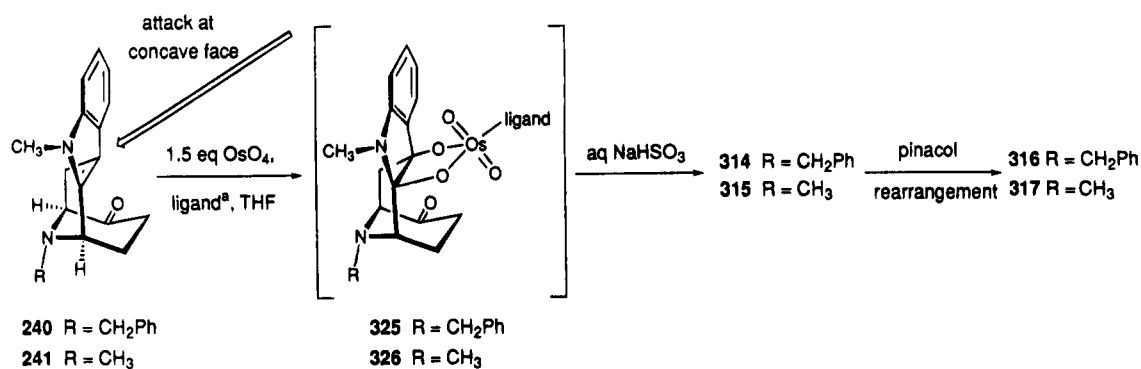
## Scheme 35



When a solution of the same (-)-ketone **240** was treated with dihydroquinine 4-chlorobenzoate in THF (Scheme 36, Table 14, entry 2), ketone (-)-**240** was converted into oxindole **316** with 30:1 diastereoselectivity in 91% isolated yield. Employment of the bulky Sharpless phthalazine ligands,<sup>296</sup> (DHQ)<sub>2</sub>PHAL and (DHQD)<sub>2</sub>PHAL (Table 14; entries 4 and 5), resulted in reduced facial discrimination. Some matching of configurations between the *Cinchona* derivative and the substrate were necessary to optimize the diastereoselectivity. In the cases immediately above (OsO<sub>4</sub>/*Cinchona* derivative) attack of the osmium reagent occurred preferentially from the concave face of the indole 2,3-double bond of substrate **240** to provide osmate ester **325** (Scheme 36). Hydrolysis of ester **325** (aqueous NaHSO<sub>3</sub>) and subsequent pinacol rearrangement provided oxindole **316**. More importantly, attack of the osmium reagents which contain bulky amino ligands on the indole 2,3-double bond occurred preferentially from the concave face without regard to asymmetry in the pendent ligand.<sup>268</sup>

When *N*<sub>1</sub>-methyl ketone **241** was treated with osmium tetroxide, osmium tetroxide/pyridine, or

## Scheme 36



<sup>a</sup>ligand = DHQ-CLB, DHQD-CLB, (DHQ)<sub>2</sub>PHAL, (DHQD)<sub>2</sub>PHAL

Table 14. Results from the Treatment of Ketones 323, 240, and 241 with Osmium Reagents<sup>a</sup>

entry	ketone	ligand <sup>b</sup>	OsO <sub>4</sub> (equiv)	T (°C)	t (days)	oxindole	yield (%)	310:316 <sup>c</sup>
1	(-)-240		1.0	reflux	3	310/316	42	91:9
2	(-)-240	DHQ-CLB	1.5	rt	3	310/316	91	3:97
3	(-)-240	DHQD-CLB	1.5	rt	3	310/316	77	20:80
4	(-)-240	(DHQ) <sub>2</sub> PHAL	1.5	rt	3	310/316	81	25:75
5	(-)-240	(DHQD) <sub>2</sub> PHAL	1.5	rt	3	310/316	82	20:80
6	(±)-323		1.0	rt	3	(±)-324	0	
7	(±)-241		1.0	reflux	3	(±)-317	36	0:100
8	(±)-241	DHQ-CLB	1.5	rt	3	(±)-317	66	0:100

<sup>a</sup> Reactions conducted in THF under a nitrogen atmosphere. <sup>b</sup> Ligands: DHQ-CLB, dihydroquinine 4-chlorobenzoate; DHQD-CLB, dihydroquinidine 4-chlorobenzoate; (DHQ)<sub>2</sub>PHAL, dihydroquinine 1,4-phthalazinediyl diether; (DHQD)<sub>2</sub>PHAL, dihydroquinidine 1,4-phthalazinediyl diether. <sup>c</sup> Ratios of diastereomers were obtained by <sup>1</sup>H NMR spectroscopy using a pulse delay of 15 s. The diastereomeric ratios of the mixtures of *N*<sub>b</sub>-benzyl oxindoles 310 to 316 were determined by <sup>1</sup>H NMR spectroscopy (500 MHz, CDCl<sub>3</sub>) on the purified mixture (flash chromatography) by integration of the *N*<sub>a</sub>-methyl singlets ( $\delta$  3.23 for 316 and  $\delta$  3.19 for 310) and confirmed by integration of the H-7 $\alpha$  protons ( $\delta$  2.54 for 316 and  $\delta$  2.91 for 310). For most reactions the diastereomeric ratios were also determined on the crude product mixtures. No significant difference in the 310:316 ratio was observed between crude and purified mixtures.

osmium tetroxide/dihydroquinine 4-chlorobenzoate, only one diastereomer, *N*<sub>b</sub>-methyloxindole 317 was produced and in 36–66% yields (Scheme 36 Table 14, entries 7 and 8). The smaller *N*<sub>b</sub>-methyl substituent in 241 and other macroline-related indoles is believed to preferentially occupy the equatorial position of the D ring.<sup>297</sup> As a result, the ligation and subsequent attack of the osmium reagent on the double bond would be hindered by the *N*<sub>b</sub>-methyl substituent and only osmate ester 313 (or 326 in the case of OsO<sub>4</sub>/DHQ-CLB or 321 in the case of OsO<sub>4</sub> alone) and subsequently diol 315 were formed regardless of the osmium reagent. Diol 315 underwent a pinacol rearrangement to furnish spirocyclic oxindole 317 with complete diastereoselectivity. The chirality at C-6 of *N*<sub>b</sub>-methyloxindole 317 is identical to that of the spirocyclic carbons present in chitosenine 298. To date, no previous examples are known in which stereoselective osmylation of the 2,3-double bond of indole alkaloids has been reported to take place in an intramolecular fashion. Stereoselective intramolecular osmylations have, however, been demonstrated for acyclic olefins containing an allylic function that is capable of coordination to osmium.<sup>298,299</sup>

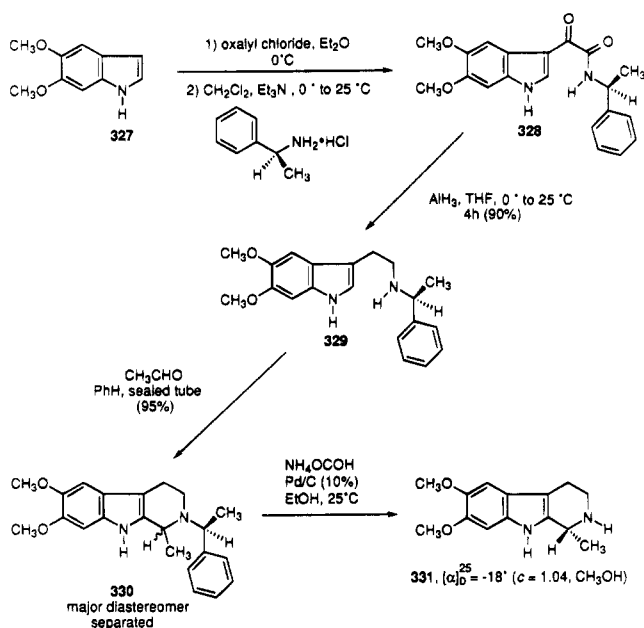
Examination of the results of this study demonstrated that oxindole 310 which is related to alstonisine (294) can be prepared with a 10:1 diastereoselectivity from the (-)-*N*<sub>b</sub>-benzyltetracyclic ketone 240 by an intramolecular OsO<sub>4</sub> complexation–control mechanism. An *N*<sub>b</sub>-benzyloxindole related to chitosenine (298), which exhibited the opposite configuration to that of alstonisine (294) about the spiro-

juncture (C-7), was also prepared by treatment of (-)-*N*<sub>b</sub>-benzyltetracyclic ketone 240 with osmium tetroxide reagents that contain bulky amino ligands. From the same optical antipode of tetracyclic ketone (-)-240 the synthesis of either the *Alstonia* oxindoles or the *Gardneria* and *Voacanga* oxindole alkaloids (diastereomeric at C-7) can be pursued. Furthermore, this approach via the inter- vs intramolecular complexation of osmium reagents may be applicable to the diastereoselective conversion of other classes of indole alkaloids into their respective oxindoles.

### VIII. Epimerization in Natural Products by Cleavage Across the Carbon–Nitrogen Bond

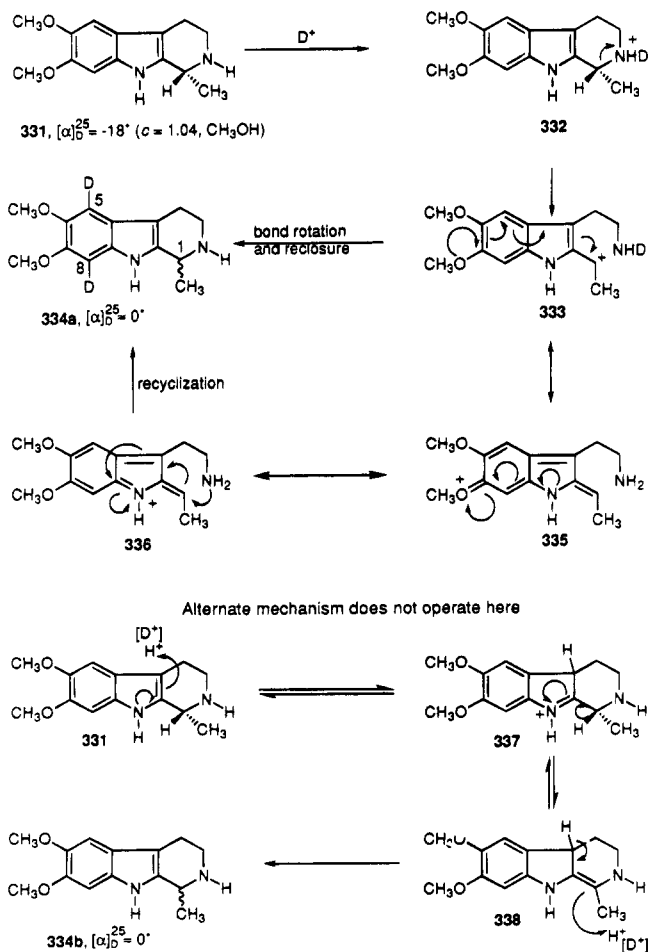
The epimerization of the stereocenter at C-1 of tetrahydro  $\beta$ -carboline by acid catalyzed cleavage across the C(1)–N(2) bond may be more general in scope than was first thought. Reddy has shown that the natural product (-)-1,2,3,4-tetrahydroroeharminine (331) can be epimerized to racemic material by stirring with acid. The synthesis of 331 (Scheme 37) began with the treatment of indole 327 with oxalyl chloride in ether which furnished the (5,6-dimethoxyindole-3-glyoxalyl chloride as an insoluble solid. On reaction with (*S*)- $\alpha$ -methylbenzylamine hydrochloride in the presence of excess triethylamine in dichloromethane, glyoxamide 328 was formed in 75% yield.<sup>300</sup> Reduction of both oxygen functions of glyoxamide 328 with AlH<sub>3</sub>, generated *in situ* in THF, provided the desired tryptamine 329 in yields ranging from 85 to 90% without racemization of the chiral

## Scheme 37



auxiliary. Pictet–Spengler condensation of **329** with acetaldehyde under the nonacidic aprotic conditions furnished 1-methyltetrahydro-β-carboline **330** as a mixture of diastereomers in a ratio of 2:1. Chromatographic separation of the major diastereomer of **330** followed by catalytic transfer hydrogenation provided (Pd/C, NH<sub>4</sub>CO<sub>2</sub>H, EtOH) (-)-1,2,3,4-tet-

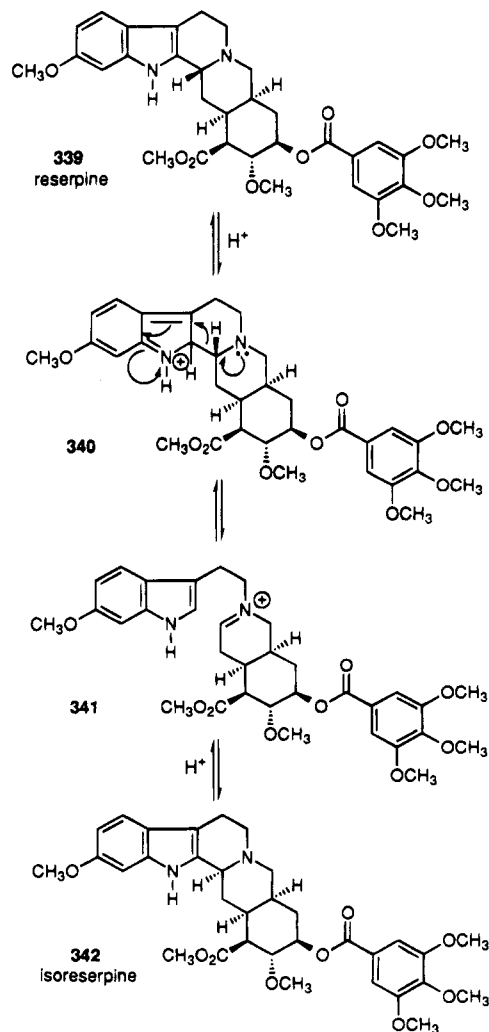
## Scheme 38. Cleavage across the C(1)–N(2) Bond



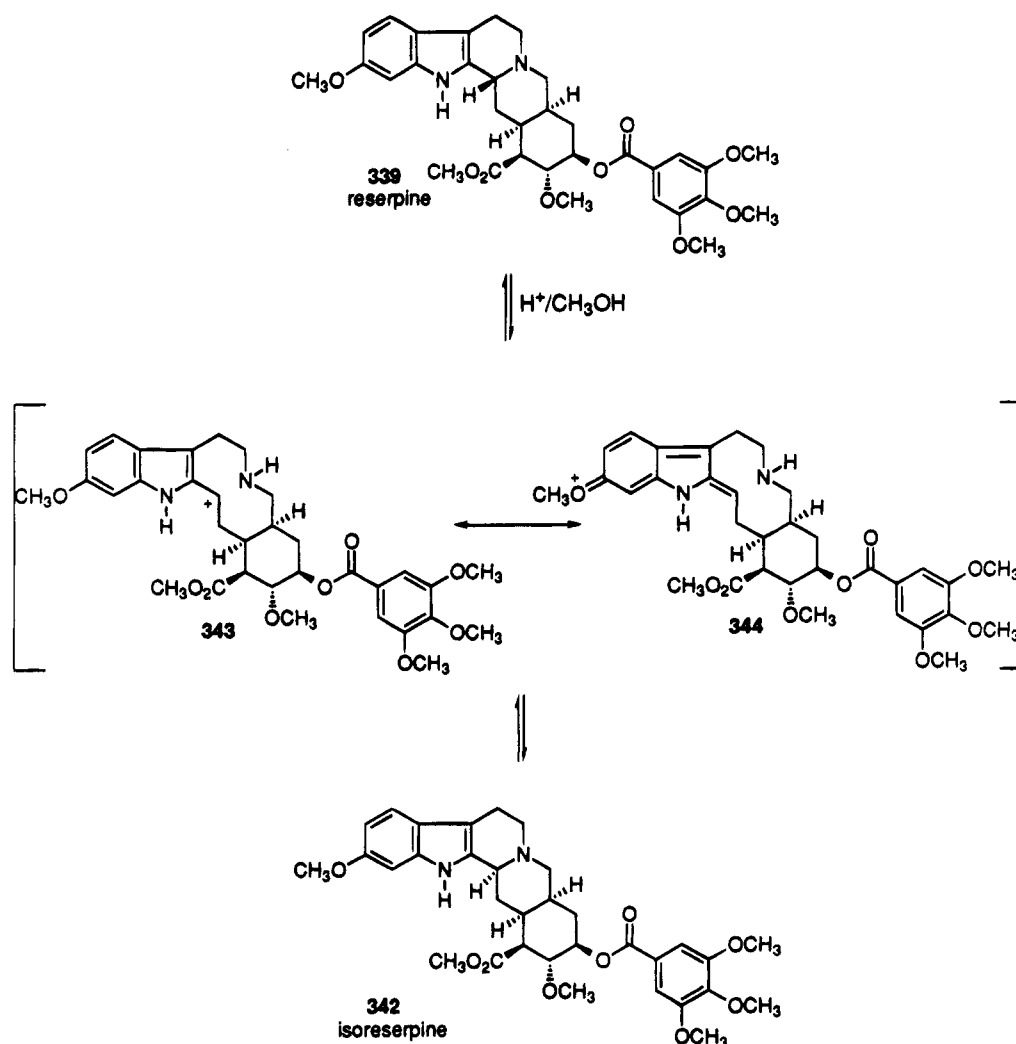
rahydroreserpine (**331**), the spectral properties of which were identical to the natural product except for the specific rotation. The specific rotation  $[\alpha]_D^{25}$  of synthetic **331** was found to be  $-18^\circ$  ( $c = 1.04$ , CH<sub>3</sub>-OH), while that reported for the natural product was  $-4^\circ$  ( $c = 0.12$ , CH<sub>3</sub>OH).<sup>301</sup>

It was believed that (-)-1,2,3,4-tetrahydroreserpine (**331**) had undergone partial racemization during the acid/base mediated isolation procedure of Gozler et al.<sup>301</sup> To test this hypothesis, optically pure **331** was exposed to deuterated trifluoroacetic acid in dichloromethane at room temperature, as illustrated in Scheme 38. The proton NMR spectrum and the *R<sub>f</sub>* value of the alkaloid which resulted were unchanged; however, the optical rotation of this material was now  $-0.8^\circ$ . On the basis of this experiment, it is believed that the mechanism of racemization of **331** occurred as illustrated at the top of Scheme 38. Racemization, we believe has occurred by cleavage across the C(1)–N(2) bond in agreement with previous work from this laboratory. Although deuterium incorporation occurred at C-5 and C-8, no deuterium was found at C-1 of **331**, consequently epimerization could not have occurred by the alternate olefinic protonation mechanism illustrated at the bottom of Scheme 38.

## Scheme 39. Previously Proposed Mechanism of Isomerization of Reserpine into Isoreserpine





**Scheme 40. Carbocation-Mediated Epimerization of Reserpine into Isoreserpine via C–N Cleavage**

The results observed here are important for all chemists who employ acidic conditions during isolation of *ring A-methoxylated* indole alkaloids. Care must be taken to avoid exposure of these natural products to acid to prevent the opportunity for racemization. In addition, it is strongly believed the racemization of (–)-tetrahydroharmine reported by Brossi<sup>302</sup> occurred in a similar fashion to that depicted in Scheme 38.<sup>300</sup>

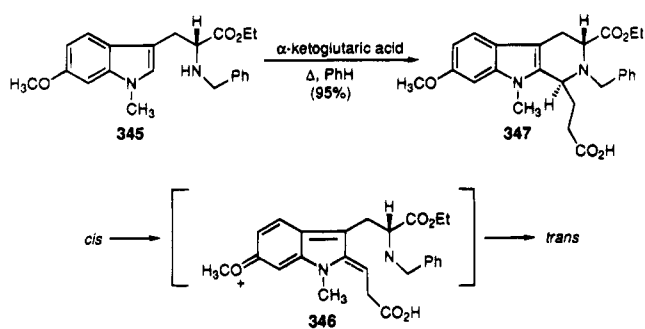
The natural product, reserpine (**339**), which was originally isolated from the Indian snake root *Rauwolfia serpentina* Benth,<sup>121,303,304</sup> is the preminent member of the yohimboid class of indole alkaloids because of its structural complexity coupled with its clinical importance as a hypotensive agent.<sup>305–310</sup> This base also exhibits significant activity as a sedative and tranquilizer.<sup>311</sup> Over the years, reserpine has been the subject of extensive chemical and synthetic investigations.<sup>312–317</sup> It has been reported that the epimerization of reserpine (**339**) to isoreserpine (**342**) can be effected under either acidic or basic conditions, moreover under acidic conditions **342** has been found to predominate over **339** in a ratio of 3.5:1.<sup>318–321</sup> Woodward suggested three possible mechanisms to account for this equilibration<sup>312,313</sup> one of which is similar to that reported by Wenkert and Liu for the epimerization of alloyohimbine to epialloyohimbine.<sup>322</sup> Joule, however, reported that the mechanism involving initial protonation at C-2 followed by

reverse Mannich fission of the C(2)–C(3) bond through iminium ion intermediate **341** was responsible for the isomerization between **339** and **342**, as illustrated in Scheme 39.<sup>321</sup>

Observations by Martin<sup>317</sup> during the total synthesis of **339** prompted a reinvestigation of the equilibration in our laboratory. In brief, Martin formed iminium ion **341** by  $\text{Hg}(\text{OAc})_2$  oxidation of its corresponding dihydro derivative. Cyclization of the iminium ion **341**, proceeded via a half-chair/chair conformation to provide reserpine (**339**) as the major product accompanied by isoreserpine in a ratio of 4.3:1. If iminium ion **341** was indeed an intermediate in the conversion of **339** into **342**, as proposed earlier by Joule<sup>321</sup> then isoreserpine should have been isolated as the major product<sup>321</sup> contrary to the experimental findings of Martin.<sup>317</sup> The Joule intermediate, iminium ion **341**, therefore cannot be the predominant species in the acid-catalyzed isomerization.

The reader will recall that *cis*-1,3-disubstituted 1,2,3,4-tetrahydro- $\beta$ -carboline **171** could be converted into the thermodynamically more stable *trans* isomer **172** by bond cleavage across the C(1)–N(2) bond followed by intramolecular recyclization, as discussed above (Scheme 13).<sup>252</sup> Encouraged by these results, reserpine **339** was heated in methanolic hydrogen chloride (1% HCl), under conditions analogous to those employed to convert **171** into **172**. This process

## Scheme 41



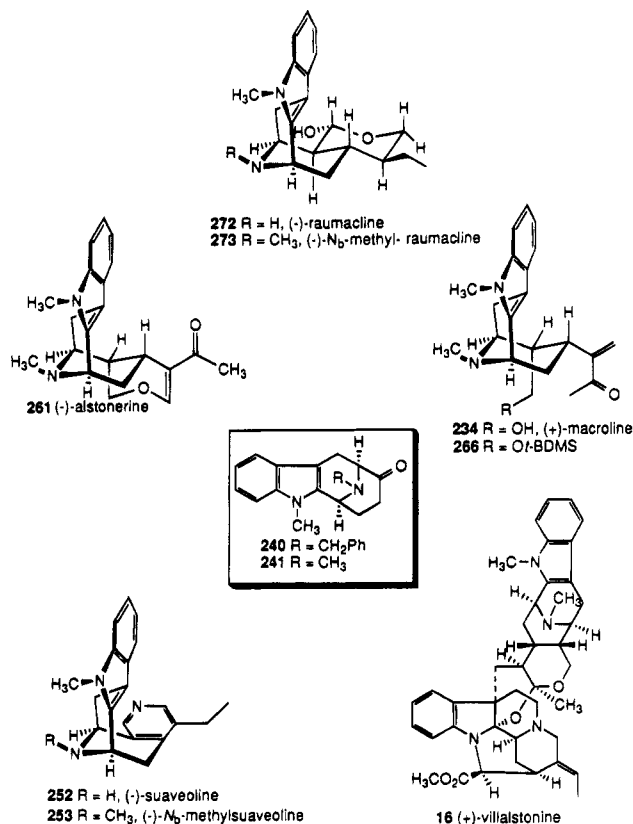
gave a mixture of isoreserpine (**342**) and reserpine (**339**) with **342** predominating in the product mixture (3:1). Moreover, when isoreserpine (**342**) was heated in methanolic hydrogen chloride it remained the major alkaloid isolated from this equilibration in agreement with its thermodynamic stability.

Upon the basis of the cyclization of **341** reported by Martin,<sup>317</sup> numerous experiments,<sup>111,323</sup> and stereochemical considerations, it was felt that the epimerization of reserpine at C-3 occurred via the pathway outlined in Scheme 40. Protonation of **339** at N-4 followed by ring scission of the C(3)–N(4) bond would afford the carbocation intermediate **343**. This carbocation can then cyclize to furnish isoreserpine (**342**), a thermodynamically more stable molecule with the indole group in an equatorial position relative to ring C.<sup>323</sup> In agreement with the mechanism of cleavage across the C(1)–N(2) bond for the epimerization of *cis*-1,3-disubstituted diastereomers into the *trans* isomers, Hamaker has recently synthesized the key 6-methoxy analog **347** required for the enantiospecific synthesis of alstoneryline and macralstonine. When 6-methoxy-*N*<sub>b</sub>-benzyl-D-(+)-tryptophan ethyl ester (**345**) was treated with  $\alpha$ -ketoglutaric acid in benzene at reflux only the required *trans* diastereomer **347** was isolated in 95% yield (>98% ee). Presumably the ring A alkoxy group stabilized the intermediate carbocation at C-1 (Scheme 41) permitting the conversion of any *cis* diastereomer so generated into the desired *trans* diastereomer.<sup>324</sup>

## IX. Conclusion

The detailed study of the Pictet–Spengler reaction has progressed from the discovery of improved non-acidic aprotic reaction conditions to probing the mechanistic phenomena involved. We have been able to explore the scope of the condensation including the underlying causes for stereospecificity, the mechanism of epimerization of the C-1 carbon atom, and the optimal conditions for effecting highly stereoselective reactions. Although these studies are important, their purpose was to provide a gateway through which to access pharmacologically interesting indole alkaloids.

It is clear that enantiomerically pure *N*<sub>b</sub>-benzyl-tryptophan alkyl esters can be condensed with aldehydes (acid labile or otherwise) to provide *trans*-1,3-disubstituted 1,2,3,4-tetrahydro- $\beta$ -carbolines in stereospecific fashion. In those cases in which a small amount of the undesired *cis* diastereomer is formed, addition of excess TFA converts the mixture into the enantiomerically pure *trans* diastereomer. Pictet–



**Figure 19.** Indole alkaloids prepared from the (-)-tetracyclic ketone.

Spengler reaction of these same aldehydes and esters under acidic conditions should also provide the *trans* isomer with 100% diastereoselectivity, albeit with slightly lower yields.

As such, the enantiospecific synthesis of (-)-alstoneryline (**261**) and (+)-macroline **234** have been completed in greater than 98% ee starting from the optically active tetracyclic ketone **240** which had been prepared by an asymmetric Pictet–Spengler reaction. The syntheses of **234** and **261** are the first enantiospecific syntheses of members of the *Alstonia* class of indole alkaloids. Since the three intramolecular reactions (the Pictet–Spengler reaction, the stereocontrolled Dieckmann cyclization, and the Claisen rearrangement) employed in the syntheses provide an intermediate **240** which possessed the same stereochemical configuration at C-3, C-5, and C-15 as those in the macroline, sarpagine, and ajmaline alkaloids with high stereoselectivity, a general approach for the preparation of these alkaloids has been developed (Figure 19). The significance of the synthesis of (+)-macroline (**234**) becomes apparent when one considers that >70 macroline related alkaloids have been isolated and that macroline is known to serve as a biogenetic precursor for many of the bisindole alkaloids as well. Macroline (**234**) is known to cyclize to dihydroalstoneryline when exposed to base and is not stable in a vial for long periods of time; therefore, the synthesis of the stable macroline equivalent **266** described herein will obviously facilitate the synthesis of bisindole alkaloids which exhibit greater biological activity than the monomers that constitute them.<sup>24,276</sup> Synthetic (+)-macroline (**234**) has now been coupled stereospecifically with natural pleiocarpamine (**235**) to furnish the antiprotozoal

alkaloid villalstonine (**16**) which contains 11 chiral centers and 11 carbocyclic rings. The macroline intermediate **234** will further provide a means to achieve the total synthesis of other dimeric indole alkaloids in this series.

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