
Development of Fast Atom Bombardment Mass Spectral Methods for the Identification of Carcinogen-Nucleoside Adducts

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An analytical strategy using fast atom bombardment (FAB) ionization and tandem mass spectrometry has been developed to determine the molecular weight and major fragment ions, and to provide limited structural characterization of low picomole levels of carcinogen-nucleoside adducts. This strategy consists of three main components: (1) the sensitivity for analysis by FAB combined with mass spectrometry is increased via chemical derivatization; (2) the nucleoside adducts are selectively detected by using constant neutral loss scans; and (3) structurally characteristic fragments are obtained by using daughter ion scans. Trimethylsilyl derivatized arylamine-nucleoside adducts have been detected at levels as low as a few picomoles by using this approach. After experimental determination of the mass of the BH_2^+ fragment ion, daughter ion spectra have been used to probe the structure specificity associated with collision-activated decomposition of this fragment. With model C-8 substituted arylamine adducts [N-(deoxyguanosin-8-yl)-4-aminobiphenyl, N-(deoxyadenosin-8-yl)-4-aminobiphenyl, and N-(deoxyguanosin-8-yl)-2-aminofluorene], nucleoside-specific and carcinogen-specific fragmentation have been observed in daughter ion spectra. (*J Am Soc Mass Spectrom* 1992, 3, 360-371)

A central tenet of cancer research is that covalent carcinogen-DNA adducts are associated with the risk of developing tumors [1]. Not surprisingly, it has been suggested that these adducts could be used as biological dosimeters of exposure to chemical carcinogens. This approach offers significant advantages over traditional methods of risk assessment, such as environmental monitoring [2-7]. This use of carcinogen-DNA adducts, however, is critically dependent upon the development of analytical methods that allow their identification and quantification at very low (picomole) levels [1].

Methods to measure DNA adducts in human tissues have employed enzyme immunoassays [8-13], fluorescence assays [11, 14], and ^{32}P -postlabeling [8, 15-17]. The principal advantage of these techniques is their great sensitivity. For example, the ^{32}P -postlabeling method has the capability to detect minute quantities of adducts, perhaps one adduct in 10^{10} nu-

cleotides [18]. Nevertheless, identifications using this method are generally based on cochromatography with known standards, and at best, only tentative identifications can be made for unknown adducts. The other widely used technique, immunoassay, involves the use of antibodies raised against either the specific carcinogen-modified DNA or nucleoside adducts coupled to a protein carrier [19]. Methods based on immunoassay suffer from some limitations when applied to the identification of unknowns. False-positive confirmations may arise due to cross reactivity with structurally similar adducts. Also, unknown adducts will likely not elicit a response to a specific antibody derived from a carcinogen of unrelated structure, and thus remain undetected. This combination of exquisite sensitivity and poor structure specificity has resulted in the isolation of carcinogen-DNA adducts that can be detected but not identified (see ref 8 for example).

Electron impact mass spectrometry (EI/MS) has the sensitivity to detect comparably low levels of volatile organic compounds. This technique has not been successfully applied to the analysis of carcinogen-modified deoxynucleosides isolated from tissue DNA, even

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when the compounds are derivatized to increase volatility. Chemical ionization mass spectral methods for the detection of specific DNA adducts have been reported, especially with electrophore labels to enhance negative ion formation [20-22]. Typically, the sugar is removed to enhance volatility prior to electrophore derivatization. Giese and co-workers [20, 21] accomplished this using sequential oxidation and reduction prior to derivatization. Similarly, Swenberg and co-workers [22] have used gas chromatography/mass spectrometry (GC/MS) with negative ion chemical ionization (NICI) to analyze trace levels of N²-3-ethenoguanine following electrophore derivatization.

Fast atom bombardment (FAB) ionization has been shown amenable to the identification of microgram levels of purified nucleosides [23-25] and carcinogen-nucleoside/nucleotide adducts [26]. We have reported the identification of deoxynucleoside/nucleotide adducts based, in part, on analysis by FAB combined with mass spectrometry (FAB/MS) [27-30]. Unfortunately, for applications involving the analysis of tissue samples, microgram quantities of adducts cannot easily be obtained. Thus, we have become interested in methods that allow the identification of unknown carcinogen-deoxynucleoside adducts at levels in the 1-10 ng range.

Trimethylsilylated derivatives of deoxynucleoside-carcinogen adducts show surprising sensitivity by FAB/MS, possibly even greater than EI/MS [27, 31]. For example, the adduct reported in ref 27, N-(deoxyadenosin-8-yl)-4-amino-3-chlorobenzyl alcohol, was correctly identified after FAB/MS analysis of the silylated derivative but not by using EI/MS. Recent results from Schram's laboratory [32, 33] show surprisingly low level detection for silylated derivatives of related compounds, including ribonucleosides and ribonucleotide monophosphate derivatives by FAB/MS. This approach is analogous to the electrophore derivatization methods used to enhance the detection of alkylated nucleosides and bases using NICI mass spectrometry [34], or electron capture GC [35].

In this article we report the development of FAB combined with tandem mass spectrometry methodologies for the analysis of derivatized arylamine-deoxynucleoside adducts at levels as low as 1 ng. The adducts are converted to trimethylsilyl (TMS) derivatives and examined by using FAB ionization and constant neutral loss (CNL) scans. FAB ionization provides a characteristic mass spectral fragment-marker ion (sugar loss) that can be used in CNL scans to search for potential nucleoside adducts. Subsequent daughter ion scans, using the experimentally determined BH₂⁺ fragment ion rather than the protonated (or deprotonated) molecule ion, are then used to provide evidence for adduct structures. This method is applicable even when the mass selected parent ion cannot be identified/recognized in the normal FAB mass spectrum. Results obtained by using underiva-

tized adducts and soft ionization methods other than FAB/MS are reported for comparison. Preliminary accounts of our work have been presented [36-38].

Experimental

Adduct synthesis. N-(2'-deoxyguanosin-8-yl)-4-aminobiphenyl was synthesized by a modification of a published procedure [39] and then purified by preparative high-performance liquid chromatography (HPLC). N-(guan-8-yl)-4-aminobiphenyl and N-(guan-8-yl)-4-aminobiphenyl-d₉ were synthesized by reaction of 4-aminobiphenyl or 4-aminobiphenyl-2,2',3,3',4,5,5',6,6'-d₉ with guanine-N-oxide in 10:1 dimethyl sulfoxide/dimethylformamide at room temperature overnight. The other adducts were obtained from Frederick A. Beland and Fred F. Kadlubar of the National Center for Toxicological Research.

Derivatization procedure A. Measured portions of the adduct were added to tapered glass vials and the ethanol solvent was removed with dry argon gas. The silylating reagent (N-methyl-N-trimethylsilyltri-fluoroacetamide [MSTFA], N,O-bis(trimethylsilyl)trifluoroacetamide [BSTFA], or N-methyl-N-(tert-butyl)dimethylsilyl-trifluoroacetamide [MTBSTFA]) in acetonitrile or dimethylformamide (1:1 v/v) was then added under argon and allowed to react 1-2 h at room temperature or at 60-100 °C for 1 h.

Derivatization procedure B. Adducts were silylated by adding the reagent directly to the dry sample and heating at 60-100 °C for 1-2 h. Mass spectral analyses were performed directly on the resulting solutions containing the derivatized sample dissolved in the derivatizing reagent.

Thermospray mass spectrometry (TS/MS). TS mass spectra were obtained by using a Finnigan MAT (San Jose, CA) TSQ-70 triple stage quadrupole mass spectrometer equipped with a Finnigan TS controller and source. An ISCO (Lincoln, NE) LC-5000 syringe pump connected to a Rheodyne 7125 injector (20 μL loop) provided a flow rate of ~1.5 mL/min. The mobile phase typically was 80-90% aqueous buffer (0.1 M ammonium acetate and methanol). The vaporizer temperature was adjusted to a cool setting (~90 °C) to maximize abundance of the protonated molecule.

Desorption chemical ionization (DCI) mass spectra. DCI mass spectra were obtained with a Finnigan MAT 4023 operating in the CI mode with methane or 10% ammonia in nitrogen as the reagent gas. Samples were introduced via a Vacumetrics (Ventura, CA) or a custom-made DCI probe incorporating a platinum filament heated at a rate of 50 mA/s for 60 s. The reagent gas was set to an uncorrected nominal source pressure of 0.25 torr. The source temperature was 200-220 °C.

FAB mass spectrometry. Samples were applied directly to a copper target via a syringe (0.5–1.0 μL) after application of 1–2 μL of thioglycerol FAB matrix liquid. Spectra were obtained by using a Finnigan MAT TSQ-70 with an Ion Tech (Middlesex, UK) gun. FAB ionization was effected by using xenon atoms accelerated to 8–10 keV. Scans were obtained from about 900 u down to about 70 u under control of the TSQ-70 data system.

FAB/MS/MS. (a) Constant neutral loss scans were acquired by using the Finnigan MAT TSQ 70 as described above for FAB/MS. To obtain a constant neutral loss scan the first and last quadrupoles, Q1 and Q3, were scanned at the same rate with a constant mass offset corresponding either to the mass of a deoxyribose moiety (116 u) or a *bis*-TMS derivatized deoxyribose (260 u). (b) Daughter ion scans were obtained by fixing the mass passed by Q1 to correspond to the protonated molecule, the BH_2^+ ion, or a TMS derivative of the former, while the mass of Q3 was scanned to detect the masses of all fragments formed by decomposition of the mass selected parent ion. Both types of scans were acquired by using argon as the collision gas to induce decomposition of the parent ion. Collision energies of 30–90 eV were utilized and collision cell pressures were varied to maximize the yield of useful daughter ions.

Results and Discussion

Sensitivity Studies

The initial objectives of the sensitivity studies were: (a) to demonstrate that FAB/MS is a sensitive soft ionization methodology for the analysis of carcinogen-modified deoxynucleosides; and (b) to optimize experimental conditions so that full-scan spectra could be obtained by using minimum amounts of sample. The analysis of carcinogen-DNA adducts from *in vivo* rather than *in vitro* studies is currently hindered by the inability to routinely obtain mass spectra from low nanogram to picogram levels of samples.

Comparison of ionization methods. To address the first objective, mass spectra of the model adduct N-(deoxyguanosin-8-yl)-4-aminobiphenyl (dG-C8-ABP, Figure 1) were obtained by using DCI, TS, and FAB ionization. [This adduct, the principal modified nucleoside isolated from urinary bladder DNA upon exposure to the arylamine carcinogen 4-aminobiphenyl (ABP), is attributed to a reaction between DNA and a reactive metabolite, namely N-hydroxy-ABP.] Analyses were attempted by using 1 μg , 500 ng, 200 ng, 100 ng, and 10 ng of adduct with each of the ionization techniques. Typically, spectra were obtained by scanning from 600 u down to 100 m. The nominal sample sizes giving reproducible full-scan spectra under the conditions described above are enumerated in Table

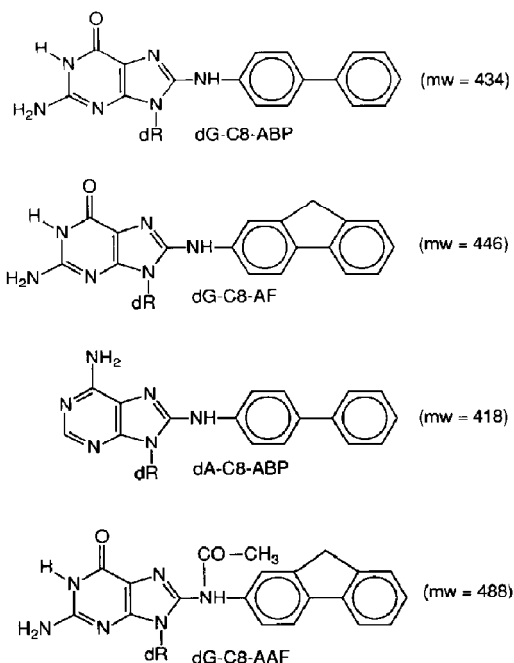


Figure 1. The structures of model nucleoside-carcinogen adducts used in this study: N-(deoxyguanosin-8-yl)-4-aminobiphenyl, dG-C8-ABP; N-(deoxyguanosin-8-yl)-2-aminofluorene, dG-C8-AF; N-(deoxyadenosin-8-yl)-4-aminobiphenyl, dA-C8-ABP, and N-(deoxyguanosin-8-yl)-2-acetylaminofluorene, dG-C8-AAF.

1. For each ionization technique, the lowest limit of detection was 100 or 200 ng. Figure 2 shows a typical spectrum obtained from the analysis of 100 ng of dG-C8-ABP by using FAB ionization. The protonated molecule and BH_2^+ ion are detectable above the near continuous background signal attributed to the FAB matrix liquid (thioglycerol). Moser and Wood [24] and Weng et al. [32] have reported comparable detection limits (250–500 ng) for unmodified nucleotides by FAB/MS using *p*-toluenesulfonic acid/glycerol or glycerol as matrix compounds.

For comparison the experiment was repeated by using dG-C8-ABP derivatized to incorporate TMS groups. Solutions of the derivatizing reagent, a solvent, and the adduct were prepared so that the appropriate amounts of adduct (1 μg to 10 ng) could be analyzed directly from the derivatization solution. TS/MS was not used in this comparison as the derivatized adducts were not expected to be stable in an aqueous solution. Procedure A was used for derivatization with MSTFA. Mass spectra were acquired over a wide mass range (900 u) to simulate the experimental conditions appropriate for the examination of unknown adducts. For example, the molecular weight of the *tetrakis*-TMS derivative of a deoxyguanosine adduct of a carcinogen having a molecular weight above 350 u would exceed 900 u.

Table 1. Compilation of the minimum amount of carcinogen-nucleoside adduct needed to obtain full-scan mass spectra by using various soft ionization mass spectrometry methods

Method	Adduct	Derivative ^a	Amount ^b	Figure
DCI/MS ^c	dG-C8-ABP	—	200 ng	—
DCI/MS ^d	dG-C8-ABP	—	200 ng	—
TS/MS	dG-C8-ABP	—	200 ng	—
FAB/MS	dG-C8-ABP	—	100 ng	2
DCI/MS ^c	dG-C8-ABP	MSTFA (A)	200 ng	3
DCI/MS ^d	dG-C8-ABP	MSTFA (A)	200 ng	—
FAB/MS	dG-C8-ABP	MSTFA (A)	10 ng	4
FAB/MS	dG-C8-AF	MSTFA (A)	5 ng	5
FAB/MS	dG-C8-AF	MSTFA (B)	1 ng	—

Abbreviations: DCI = desorption chemical ionization;
 FAB = fast atom bombardment ionization;
 MSTFA = N-methyl-N-trimethylsilyl-trifluoroacetamide;
 ng = nanograms;
 TS = thermospray ionization.

^aA and B refer to the derivatization procedures (see Experimental).

^bApplied to the probe tip.

^cMethane reagent gas.

^dAmmonia reagent gas.

Incorporation of TMS groups did not significantly reduce the amount of adduct needed to produce spectra using DCI/MS with either methane or ammonia as the reagent gas. Typical of these results is the methane DCI mass spectrum shown in Figure 3. About 200 ng of dG-C8-ABP derivatized with MSTFA/DMF was required to obtain reproducible spectra. While the protonated molecules for the *tris*- and *tetrakis*-TMS containing derivatives (m/z 651 and 723) were observed, the fragment ions corresponding to TMS containing BH_2^+ ions (m/z 391 or 463) were not. The ions at m/z 487, 413, and 307 have not been identified.

In contrast to the DCI work, FAB mass spectra were obtained by using smaller amounts of silylated adduct. Figure 4 shows the FAB mass spectrum obtained when a 100-ng sample of dG-C8-ABP was derivatized (Procedure A) and 10% (10 ng) applied to the probe tip for analysis. The ions at m/z 651 and

579 correspond to the protonated molecules for the *tris*- and *bis*-TMS derivatives, whereas the ion at m/z 391 corresponds to the mono-TMS derivative of the BH_2^+ fragment ion. This fragment could occur via loss of a *bis*-silylated deoxyribose from m/z 651 or via loss of a monosilylated sugar from m/z 579. Based on previous reports of complete silylation of the deoxyribose moiety, even with incomplete derivatization of the carcinogen/base moiety, the former pathway appears to be more likely [40]. The unsilylated fragment ion, BH_2^+ , was also observed at m/z 319 but is not displayed in Figure 4. This ion is attributed to loss of the disilylated deoxyribose moiety from those *bis*-silylated adduct protonated molecules having both TMS groups attached to the sugar.

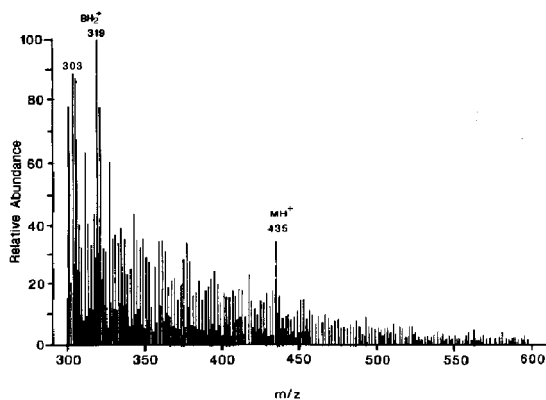


Figure 2. The FAB mass spectrum obtained by using 100 ng of dG-C8-ABP showing the protonated molecule (m/z 435) and BH_2^+ ion (m/z 319).

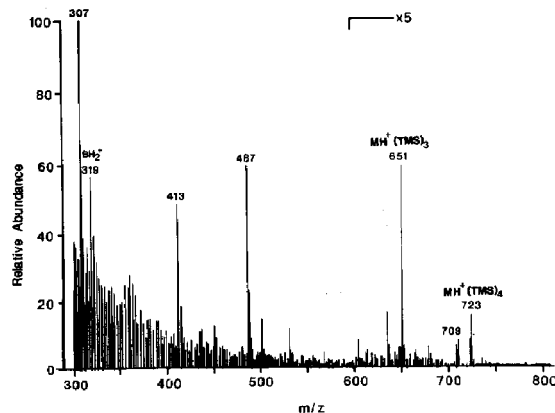


Figure 3. The methane DCI mass spectrum obtained by using 200 ng of dG-C8-ABP derivatized with MSTFA/dimethylformamide showing protonated molecules for the *tetrakis*-TMS derivative (m/z 723) and the *tris*-TMS derivative (m/z 651), as well as a BH_2^+ ion with no TMS groups (m/z 319).

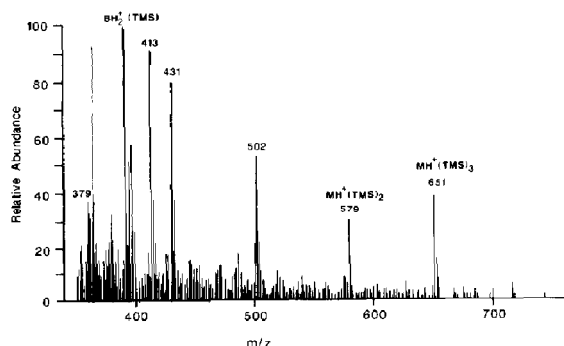


Figure 4. The FAB mass spectrum obtained by using 10 ng of dG-C8-ABP derivatized with MSTFA/dimethylformamide showing protonated molecules for the *tris*-TMS (m/z 651) and *bis*-TMS species (m/z 579) and a BH_2^+ fragment ion containing a single TMS group (m/z 391).

Optimization of derivatization conditions and comparison with other adducts. Similar experiments were also performed by using other adducts and other derivatization reagents. Figure 5 shows the spectrum obtained from 5 ng of N-(deoxyguanosin-8-yl)-2-aminofluorene, dG-C8-AF, derivatized with MSTFA (Procedure B). The ions at m/z 663 and 591 correspond to the *tris*- and *bis*-TMS containing protonated molecules, whereas m/z 403 and 331 are the BH_2^+ fragments having one and no TMS groups, respectively. For comparison, the m/z 217 ion corresponds to the protonated thioglycerol matrix dimer. It should be noted that this matrix background ion is about 200 times smaller relative to the analyte protonated molecule ions using 5 ng of derivatized adduct (Figure 5) than when using 1 μ g of underivatized adduct (Figure 6). This shows the striking improvement in

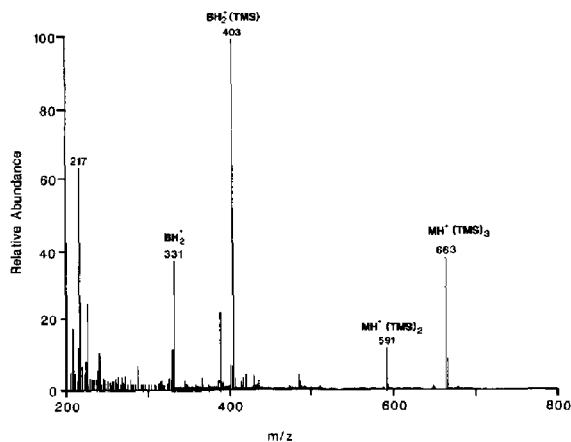


Figure 5. The FAB mass spectrum obtained by using 5 ng of MSTFA derivatized dG-C8-AF showing protonated molecules for the *tris*- and *bis*-TMS containing ions (m/z 663 and 591) as well as the BH_2^+ TMS ion (m/z 403) and the BH_2^+ ion (m/z 331).

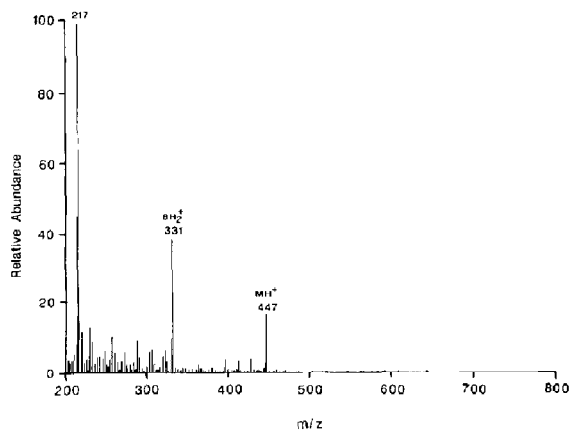


Figure 6. The FAB mass spectrum obtained by using 1 μ g of underivatized dG-C8-AF showing the protonated molecule (m/z 447), the BH_2^+ (m/z 331), and the relatively large contribution of the matrix ion at m/z 217.

the analyte to matrix signal levels observed with derivatized adducts as compared to their underivatized forms. The relative abundance of the analyte protonated molecules with respect to the thioglycerol matrix ion (m/z 217) is even more surprising when the partitioning of the molecular signal among the two TMS derivatives (m/z 663 and 591) is considered.

In a related study, Schram and co-workers [32] reported a method for the analysis of low nanogram levels of clinically important nucleotide-monophosphates by FAB/MS. Derivatization was accomplished in capillary tubes by using small volumes of BSTFA in pyridine. Analysis by EI and CI/MS indicated complete derivatization of the ribose monophosphate moiety, but not the base. They observed the loss of TMS groups, which they attributed to hydrolysis, by using either glycerol or 3-nitrobenzyl alcohol (NBA) as a FAB matrix liquid. One TMS group was lost with NBA, whereas more extensive hydrolysis and low signal levels were reported for glycerol. Hydrolysis was not observed with tetraglyme, but other experimental difficulties, including lack of sensitivity, limited its use as a FAB matrix. No results for thioglycerol were reported.

We also see evidence for hydrolysis with FAB/MS using thioglycerol. Differences in the number of TMS groups are observed when TMS derivatized adducts are analyzed by DCI/MS and FAB/MS (Figures 3 and 4, for example). However, the number of TMS groups appears constant from scan to scan via either method, suggesting that hydrolysis occurs rapidly and to completion prior to introduction of the sample into the ion source. In other words, matrix induced hydrolysis likely occurs almost immediately upon introduction of the sample into the FAB matrix liquid.

If the DCI mass spectrum is assumed to accurately reflect the number of TMS groups added prior to introduction of the analyte into the FAB matrix liquid,

the number of groups expected by FAB/MS without hydrolysis would be 3-4. The addition of this number of TMS groups, although it represents incomplete derivatization, is in good agreement with a previous study of carcinogen-nucleoside adducts using BSTFA and EI/MS [41]. The TMS derivatization of a similar C8-substituted deoxyguanosine adduct produced *tris*- and *tetrakis*-TMS derivatives with a ratio of about 1:2. In our work we have typically observed *bis*- and *tris*-TMS derivatives in about the same ratio (i.e., Figures 4 and 5). These data suggest that only one TMS group is lost via hydrolysis in thioglycerol. This is a lesser extent of hydrolysis than was attributed to the use of glycerol in ref 32, and is more consistent with the results reported for NBA. In other words, we attribute the observation of partially derivatized DNA-carcinogen adducts in thioglycerol to two causes. First, typical arylamine nucleoside adducts are not likely to be completely derivatized prior to their introduction into the matrix. Subsequently the thioglycerol matrix also induces loss of approximately one additional TMS group. The derivatization procedure may slightly affect the extent of this hydrolysis. When pure MSTFA is used for derivatization, the subsequent FAB spectra sometimes show a small increase in the *tris*-TMS derivative relative to the *bis*-TMS derivatized adduct. For example, the FAB/MS spectrum shown in Figure 4 was obtained by using 1:1 MSTFA/dimethylformamide, whereas Figure 5 represents derivatization in pure MSTFA. By either method, FAB/MS analysis of the TMS derivatives gives about the same detection limit in thioglycerol previously reported for NBA.

In another recent report, Vourous and co-workers [31] showed FAB/MS spectra of TMS derivatives of deoxynucleoside-carcinogen adducts. They reported extensive hydrolysis using thioglycerol, based on the presumption that the adducts were initially completely derivatized. However, they showed no evidence regarding the extent of derivatization prior to FAB/MS analysis. Based on results already cited from ref 41, as well as a review of the EI/MS data for 16 silylated deoxynucleoside-arylamine adducts [40], the assumption of complete derivatization prior to analysis by FAB/MS may have been incorrect. Nevertheless, the FAB mass spectra in ref 31 are very similar to those we report. For example, a spectrum obtained by using 67 ng of TMS derivatized dG-C8-AF was very similar to Figure 5, obtained by using 5 ng. One notable difference is that the extent of hydrolysis is slightly greater in all of the FAB mass spectra reported in ref 31. Vourous's group reported that the *bis*-TMS protonated molecule was always larger than the *tris*-TMS species under the conditions they described, whereas we often see the opposite. More interesting is the report of better results by using thioglycerol rather than NBA as a FAB matrix.

Derivatization with MSTFA, BSTFA, and MTBSTFA produced similar improvements in sensitivity in

our experiments. FAB/MS spectra were readily obtained by using 5 ng of dG-C8-AF derivatized with MTBSTFA (spectrum not shown); however, the abundance of the protonated molecule was relatively small. The *tris*-(dimethyl, *tert*-butylsilyl) (DMTBS) derivatized protonated molecule was observed at m/z 789 as was a mono-DMTBS BH_2^+ ion (m/z 445) ion attributed to loss of the *bis*-DMTBS derivatized sugar. The *bis*-DMTBS protonated molecule (m/z 675) was not observed and the BH_2^+ ion was less intense than either m/z 789 or 445. Numerous other ions in the spectrum were not identified. A particular problem with MTBSTFA derivatization was that for each DMTBS group added, the adduct increased in mass by 114 u rather than the 72 u for TMS. Hence, a DMTBS derivatized nucleoside-carcinogen adduct including a large carcinogen could exceed 1000 u in mass.

A major limitation upon reduction of the sample sizes for mass spectral analysis was dilution of the adducts in the derivatization reaction mixture. The derivatization procedure was changed by omitting the cosolvent to reduce the overall sample volume (Procedure B, Experimental). In subsequent experiments the derivatizing reagent was added directly to the dry nucleoside-adduct residue in a small conical vial. Using this procedure it was possible to derivatize as little as 2 ng of the modified nucleoside with 2 μ L of MSTFA and to then remove 1 μ L for analysis. The use of pure MSTFA may also have resulted in a minor reduction in the extent of hydrolysis, as indicated above. This same approach has been proposed in ref 31. In that article the authors recommended derivatization of 200 ng of adduct using 10-20 μ L BSTFA [31]. However, they also apparently observed more hydrolysis. We have observed similar results by using other arylamine adducts, namely, N-(deoxyadenosin-8-yl)-4-aminobiphenyl (dA-C8-ABP), and N-(deoxyguanosin-8-yl)-2-acetylaminofluorene (dG-C8-AAF) (spectra not shown).

Constant Neutral Loss Scans

The use of CNL scans for the analysis of modified nucleosides was also evaluated. By using this approach all of the ions that arise via a mass loss corresponding to the deoxyribose moiety can then be identified. The characteristic cleavage to produce the $[BH_2]^+$ ion has been observed in mass spectra obtained from protonated nucleosides/nucleotides [23-25], carcinogen-modified nucleosides [26, 30, 31], and silylated carcinogen-modified nucleosides [27, 31]. We have recently reported the successful application of CNL scans (116 u) to the identification of an unknown carcinogen-nucleoside adduct. When HPLC separated fractions from the *in vitro* reaction of DNA with a nitro-PAH metabolite, 2-nitropyrene, were examined by using CNL scans, two adducts were identified. One of these was an unusual C8-modified

deoxyadenosine adduct [30]. By using the same approach, Claeys and co-workers [42] have demonstrated the utility of CNL scans for the analysis of nucleosides of phenyl glycidyl ethers. They reported detection limits of about 1-10 ng. Because daughter ion scans of the protonated molecule were not particularly characteristic of parent ion structures, they used daughters of the $[M-H]^-$ ion, in the negative ion mode, for structure characterization, whereas we employ BH_2^+ ions for the same purpose.

Of particular importance to our study was whether the CNL scan method would allow additional reduction in the minimum amount of sample needed for the analysis of nucleoside-carcinogen adducts by mass spectrometry. As the signals for the protonated molecule and the BH_2^+ ion become weaker at lower concentrations, confirmation of the relationship between these two ions, in unknown samples, becomes increasingly difficult and important. Moreover, any improvement in sensitivity relative to that obtained by using full-scan FAB/MS of MSTFA derivatives would allow detection of adducts at levels approaching 1 pmol.

Underivatized adducts. Underivatized adducts were examined by using CNL scans (116 u) to determine if the observation of parent ions showing loss of the deoxyribose moiety was possible at levels too low to obtain full-scan FAB mass spectra. Whereas full-scan spectra required the analysis of 100 ng of underivatized adducts, the use of these CNL scans allowed adducts to be detected at levels as low as 5 ng (Table 2). Figure 7 shows a FAB CNL mass spectrum obtained by using 5 ng of dG-C8-AF. The protonated molecule at m/z 447 is the largest peak in the spectrum. The other signals are attributed to the high noise level at this lower limit of detection. Similar results have been obtained by using 8 ng of N-(deoxyadenosin-8-yl)-4-aminobiphenyl, dA-C8-ABP (spectrum not shown).

Derivatized adducts. Significant reduction in the noise level was observed in CNL scans (260 u) of MSTFA derivatized adducts. With FAB ionization, CNL scans were readily obtained by using 1-5 ng of MSTFA

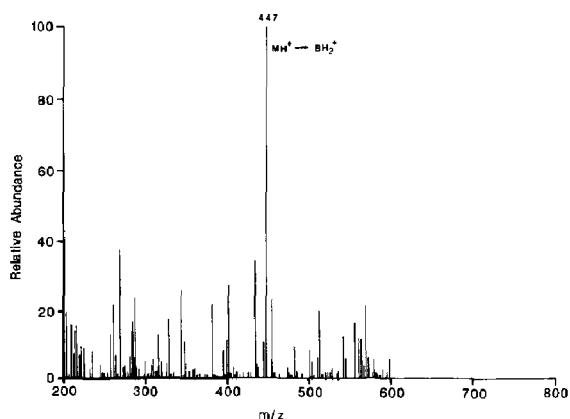


Figure 7. The FAB/CNL spectrum (116 u) obtained by using 5 ng of dG-C8-AF showing the loss of the deoxyribose moiety from the parent ion (protonated molecule) at m/z 447.

derivatized dG-C8-AF. Because a derivatized nucleoside was examined, the CNL was 260 u rather than 116 u, corresponding to the mass of a *bis*-TMS containing deoxyribose moiety. Typical of these results is the spectrum obtained by using 1 ng dG-C8-AF (Figure 8). The ions at m/z 591 and 663 correspond to the protonated molecules of the *bis*- and *tris*-TMS derivatives, whereas the ions at m/z 484 and 412 are attributed to loss of the AF moiety during derivatization. (The facile cleavage of the carcinogen from TMS derivatives of arylamine-nucleoside adducts, especially C-8 substituted deoxyguanosine derivatives, has been reported elsewhere [40].) These results should be contrasted to those shown in Figure 5, obtained by using 5 ng of MSTFA derivatized adduct via FAB/MS. Satisfactory constant neutral loss scans were consistently obtained by using smaller amounts of sample when the adducts were TMS derivatized. However, the extent of enhancement observed upon TMS derivatization was less with CNL scans (Table 2) than expected based on the results obtained by using FAB/MS (Table 1).

Analysis of unknown adducts. We have already reported the application of CNL scans to the identifi-

Table 2. Typical sample sizes needed to obtain FAB CNL spectra for loss of 116 u (the deoxyribose moiety) or 260 u (the *bis*-TMS derivatized deoxyribose) from the protonated molecule of carcinogen-nucleoside adducts

Adduct	Derivative	CNL mass loss	Amount ^a	Figure
dG-C8-AF	—	116	5 ng	7
dA-C8-ABP	—	116	8 ng	—
dG-C8-AF	MSTFA	260	1 ng	8

Abbreviations: CNL = constant neutral loss;
 FAB = fast atom bombardment ionization;
 MSTFA = N-methyl-N-trimethylsilyl-trifluoroacetamide;
 ng = nanograms.

^a Applied to the probe tip.

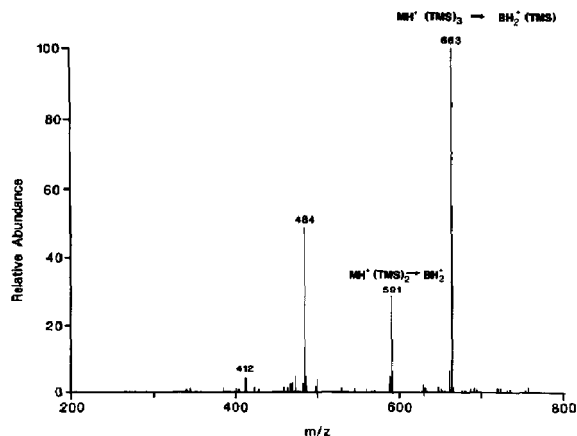


Figure 8. The FAB/CNL spectrum (260 u) obtained by using 1 ng of dG-C8-AF derivatized with MSTFA.

cation of unknown carcinogen-DNA adducts from nitropyrene metabolites [30]. Another recent experiment also demonstrates this application. Several isolated HPLC fractions, some containing carcinogen-modified DNA, were submitted to the laboratory for identification. Prior to mass spectral analysis the DNA was reacted with the N-hydroxy derivative of the heterocyclic aromatic amine 2-amino-1-methyl-6-phenylimidazo[4,5-b]pyridine (PHiP), the modified DNA was hydrolyzed to the nucleosides, and HPLC fractions containing UV adsorbing peaks with retention times consistent with modified nucleosides were collected. The individual fractions were dried, TMS derivatized, and submitted to the laboratory as unknowns. The CNL scan (260 u) obtained from one of the fractions is shown in Figure 9. The intense ion at m/z 706 corresponds to the *tris*-TMS derivative of dG-C8-PHiP. Additional evidence for this assignment was obtained by comparison of daughter ions from the BH_2^+ ion (m/z 374) and synthetic G-C8-PHiP.

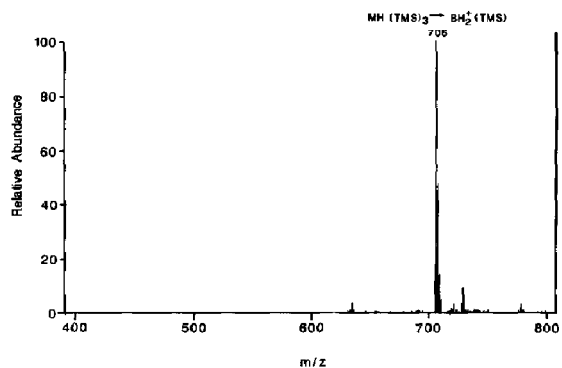


Figure 9. The FAB CNL spectrum (260 u) of an unknown HPLC separated component from the hydrolysate of N-hydroxy-PHiP treated DNA.

Daughter Ion Scans

The use of daughter ion scans to provide evidence regarding nucleoside-adduct structures was also investigated. For unknowns, even modest details regarding adduct structure could be extremely useful. For example, identification of a specific base would allow the weight of the carcinogen to be deduced. The MS/MS methodology is widely accepted as a means of detecting trace components in complex mixtures [43]. Vouros and co-workers [31] have developed a method for the detection of specific carcinogen-DNA adducts at the 100 ng level based on TMS derivatization, FAB/MS analysis, and, finally, B/E linked scans. Our method differs primarily based on the use of a CNL scan rather than FAB/MS in the first step of mass spectral analysis. These CNL scans simplify the identification of the parent ions that are used in the second step of identification based on daughter ion scans. We also note some differences in daughter ions obtained by using a triple quadrupole mass spectrometer and low energy collisional activation (CA), described below.

The typical procedure for characterizing analyte structures via soft ionization and MS/MS is CA of the analyte protonated molecule. For underivatized arylamine-nucleoside adducts this produces an intense BH_2^+ ion. For example, the daughter ion spectrum obtained when the protonated molecule (m/z 447) of dG-C8-AF was mass selected and collisionally activated showed a single major fragment ion at m/z 331 (spectrum not shown). Similar results were observed with the other model arylamine-nucleoside adducts. This fragment ion provides no additional evidence regarding the adduct's structure beyond that also observed in the FAB/MS and FAB CNL scans.

With derivatized adducts the results were similar. The daughter ions of the mono-TMS derivative of dG-C8-AF included a BH_2^+ ion at m/z 331 and a TMS- BH_2^+ ion 72 u higher in mass at m/z 403. This spectrum (Figure 10) illustrates two important points. First, as with underivatized nucleoside-adducts,

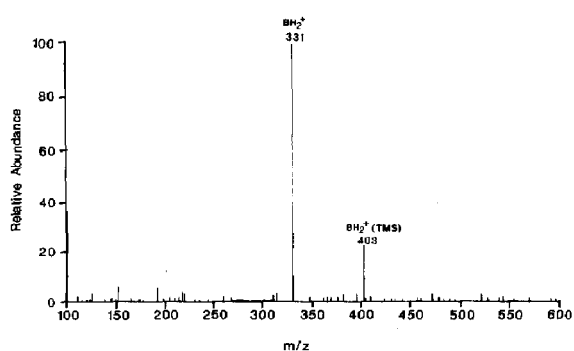


Figure 10. The FAB daughter ion spectrum from the protonated molecule (m/z 519) of the mono-TMS derivative of dG-C8-AF.

daughter ions from the protonated molecule do not provide additional data regarding adduct substructure. Second, derivatization complicates interpretation when an adduct containing TMS groups may include more than one isomer. The ion at m/z 403 in Figure 10 must have arisen via the loss of an underivatized sugar from a protonated molecule having the TMS group attached to the base or the AF moiety. On the other hand, the ion at m/z 331 must have arisen from a protonated molecule having a TMS group on the sugar but not the base or carcinogen moiety.

A simple strategy was developed to avoid problems associated with the formation of isomers accompanying the TMS derivatization step. Derivatized adducts were used to obtain CNL (260 u) and daughter ion spectra; but the specific daughter ions were underivatized BH_2^+ fragments rather than TMS-containing protonated molecules. The incomplete derivatization of the carcinogen/base moiety and the loss of one TMS in the FAB matrix liquid have resulted in the observation of non-TMS containing BH_2^+ ions from all of the MSTFA derivatized carcinogen-nucleoside adducts examined in our laboratory. Fortunately, for carcinogen-deoxynucleoside adducts, the BH_2^+ ions show considerably more structurally characteristic fragmentation than protonated molecules [31]. Be-

cause the carcinogen is invariably bound to the base rather than the deoxyribose moiety in carcinogen-DNA adducts, this strategy should not result in loss of structure specificity. The observation that daughter ion spectra from several protonated G-C8-adducts are indistinguishable from the daughters of the corresponding BH_2^+ fragments supports this hypothesis. For example, the daughters of the BH_2^+ fragment ion of dG-C8-ABP and the protonated molecule of the corresponding carcinogen base adduct (N-(guan-8-yl)-4-aminobiphenyl) are compared in Table 3.

Table 3 shows the masses of significant daughter ions observed from the BH_2^+ of dG-C8-ABP. The frequency of occurrence of daughters using various collision energies is also reported. Subjective review of these data, and the individual daughter ion spectra used to prepare Table 3, suggested that although there was significant variability between spectra at all collision energies, 50-eV collisions generally produced more abundant and reproducible fragmentation. To provide consistency this value was subsequently used in the remaining daughter ion experiments.

A daughter ion spectrum obtained by using the BH_2^+ ion from 20 ng of dG-C8-AF derivatized by using MSTFA is shown in Figure 11. These results are compared in Table 4 with daughters obtained by using dG-C8-ABP and are shown in Table 3. Peaks

Table 3. The frequency of observation of selected daughter ions from the BH_2^+ ion of dG-C8-ABP and a comparison with daughter ions from the protonated molecule of synthetic G-C8-ABP^a

Ion mass	Mass loss	Experiment Number				
		1	2	3	2 ^b	4 ^c
302	17	+	+	+	+	+
277	42	3/4	+	2/3	+	+
276	43	2/4	2/6	—	—	—
275	44	—	2/6	—	—	+
249	70	2/4	+	1/3	+	+
247	72	3/4	5/6	—	+	+
208	111	+	+	+	+	+
207	112	+	5/6	+	+	+
195	124	+	+	2/3	+	+
180	139	+	+	+	+	+
168	151	—	5/6	+	+	+
165	154	+	+	1/3	+	+
153	166	2/4	+	+	+	+
140	179	+	+	+	+	+
123	196	+	+	+	+	+
112	207	+	+	+	+	+
105	214	—	3/6	—	—	—
95	224	1/4	+	2/3	+	+
Collision energy (eV) ^d		30	50	90	50	50

^aA "+" symbol indicates that daughter ion was detected in all experiments. A fraction indicates the frequency with which a daughter ion was observed.

^bThe daughter ions expected more than 50% of the time from a single analysis at 50 eV.

^cThe daughter ions observed from the protonated molecule of G-8-ABP, for comparison.

^deV = electron volts, the laboratory value for collision energy.

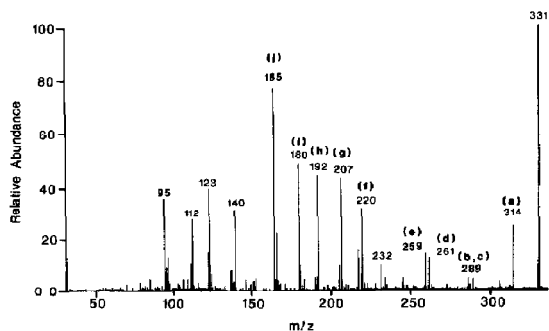


Figure 11. The FAB daughter ion spectrum from the BH_2^+ fragment ion obtained by using 20 ng of dG-C8-AF derivatized with MSTFA.

labeled a to j occurred via the same neutral losses for both of these C-8 substituted deoxy-guanosine adducts. This is attributed to fragmentation in which the carcinogen is retained within the part of the molecule having the positive charge. Experiments using a deuterated adduct (dG-C8-ABP-d₉) confirmed retention of the carcinogen moiety in these fragments. Daughter ions including the carcinogen moiety retained eight or nine of the nine deuterium atoms originally bound to the biphenyl group in the ABP moiety of the deuterated adduct.

Other characteristic ions showed the same masses rather than identical neutral losses. The ions at m/z 95, 112, 123, and 140, observed in the spectra of dG-C8-AF and dG-C8-ABP, are attributed to fragments from which the carcinogen moiety has been lost. Because the ions a to j resulted from retention of the carcinogen moiety, ions resulting from these specific neutral losses should be observed from the BH_2^+ ions of all C8-dG adducts. Similarly, the ions showing loss of the carcinogen should have common masses (m/z 95, 112, 123, and 140) for all C8-dG adducts. An exception is that because fragment j occurs via cleavage within the carcinogen, it may be specific for C8 adducts linked to the carcinogen via an amine moiety (i.e., arylamines).

The specific cleavages observed in the daughter ion spectrum (see Figure 11) are shown in Scheme I. These cleavages were chosen based on the hypothesis that for low energy collisions, the even electron BH_2^+ fragment ion would not fragment via loss of radicals. A previous report of radical ions in the FAB mass spectra of TMS derivatives of related compounds was attributed to the presence of an Si atom in the TMS moiety [33]. Because no TMS groups are present on the BH_2^+ ions used as parent ions, radical ions would not be expected. The correspondence between the spectra for the two dG-C8 adducts, as well as the

Table 4. Comparison of the major daughter ions in the tandem mass spectra of dG-C8-ABP, dG-C8-AF, and dA-C8-ABP showing common neutral losses or common fragment masses^a

Mass loss	dG-C8-ABP ^b	dG-C8-AF	Fragment ^c	dA-C8-ABP
17	+	+	a	+
42	+	+	b	
44	(2/6)	+	c	+
70	+	+	d	
72	(5/6)	+	e	
111	+	+	f	
112	+	+		
124	+	+ ^d	g	
139	+	+	h	
151	+	+ ^e	i	
166	+	+	j	
Mass				
207	+	+ ^d		+
195	+		g	+
180	+	+ ^e	h	+
168	+		i	+
153	+		j	+
140	+	+		
123	+	+		
112	+	+		
105	+			
95	+	+		

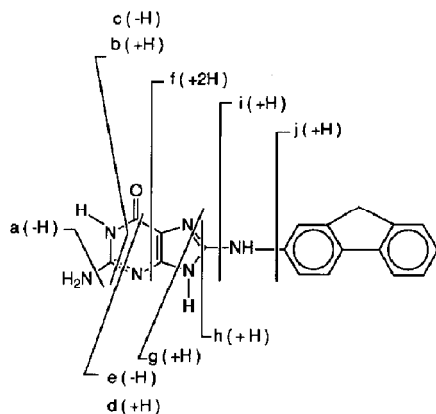
^a The collision energy (laboratory value) was 50 eV. A "+" indicates the daughter ion or mass loss is observed.

^b A "2/6" indicates that this daughter ion was observed in only two of six replicate analyses.

^c This term refers to proposed structures for the daughter ions labeled in Figure 11 as shown in Scheme I.

^d This fragment at m/z 207 is attributed to loss of 124 u via cleavage g in Scheme I.

^e This fragment at m/z 180 is attributed to loss of 151 u via cleavage i in Scheme I.



Scheme 1. Proposed fragmentation of the BH_2^+ ion from dG-C8-AF corresponding to peaks labeled a to j in Figure 11 and Table 4.

ready assignment of the carcinogen containing fragments from the adduct having deuterium isotopes on the aromatic hydrogens of the carcinogen moiety, provide strong evidence for the assignments in Scheme I. However, even if these include errors, the daughter ion methodology can be used for structure characterization based on the empirically observed correlation between neutral losses and fragment ion masses with the analyte structures.

A comparison was made with a model adduct of a different nucleoside having the same carcinogen and similar carcinogen-base bonding (dA-C8-ABP). The daughter ion spectrum (not shown) was obtained by using 42 ng of adduct. Most of the dG characteristic losses associated with retention of the carcinogen moiety in the neutral fragment were not observed with the dA adduct (see Table 4). Exceptions are ions labeled a and c. These losses, NH_3 and $\text{C}_2\text{N}_2\text{H}_2$, are relatively small molecules and might be expected from either dA or dG adducts. However, some ions are common to both the ABP adducts. For example, ions labeled g, h, i, and j are attributed to loss of most of the base moiety. Any remaining portion of the base substructure is identical for dG-C8-ABP and dA-C8-ABP, as shown in Scheme I.

Some confusion may arise because of the coincidence of daughter ion masses attributed to different fragmentation pathways in two of the adducts. The ions at m/z 207 and 180 for dG-C8-AF and dG-C8-ABP are not attributed to the same structures (see Table 4). The ions at m/z 207 and 180 in dG-C8-AF correspond to loss of 124 and 151 u, respectively, from the BH_2^+ . Hence, the m/z 207 and 180 ions from dG-C8-AF and the m/z 168 and 195 ions from dG-C8-ABP would be attributed to the same cleavage within the base moiety. The m/z 180 from dG-C8-AF arises from loss of the dG moiety to give $[\text{C}_{13}\text{H}_{10}\text{N}]^+$.

These results should be contrasted to those reported in ref 31. Daughter ions from the BH_2^+ ion

from dG-C8-AF were examined by using high energy collisional activation and B/E scans. Significant daughter ions were reported at m/z 314, 284, 259, 232, 206, 191, 180, and 165. The ions at m/z 314, 259, 232, 180, and 165 are prominent in Figure 11 as well. These ions, common to both studies, were attributed in ref 31 to the same structures we propose in Scheme I. The ions at m/z 284, 206, and 191 were observed in high energy, but not low energy, collisions. The ions at m/z 206 and 191 could be radical fragments resulting from high energy collisions, or they could be misassigned in the B/E scan. The authors of ref 31 pointed out that an error in their daughter ion assignments of 1 u was possible because of the use of linked scans. Because we have observed m/z 207 and 192 in numerous daughter ion scans from dG-C8-AF, and this same mass loss (of 124 and 151 u) from dG-C8-ABP, and dG-C8-AF, under a variety of experimental conditions, we attribute this difference to an error in the linked scan derived daughter ion mass assignments. The other ion observed in ref 31, at m/z 284, was not observed in our experiments, but was also attributed to an even electron fragment. These results suggest daughter ion spectra obtained by using low or high energy collisions provide sufficient structure specificity for use in characterizing adduct structures, even when the experiments are performed in different laboratories. Furthermore, with the exception of the two daughter ions in the B/E scans that can be attributed to an error in mass assignment, all of the daughter ions from the BH_2^+ ions of several adducts, in both studies, can be attributed to even electron fragmentation.

Conclusion

Our methodology combines three experimental approaches to the characterization of low levels of carcinogen-deoxynucleoside adducts. The utility of each component approach for the analysis of these or related compounds by mass spectrometry has been previously reported. The method incorporates the use of CNL scans for screening [30, 42], TMS derivatization for sensitivity enhancement [31-33], and daughters of the BH_2^+ ion for structure characterization [31].

FAB ionization with CNL scans of MSTFA derivatized samples allowed identification of carcinogen-deoxynucleoside adducts at levels as low as 1 ng. For a carcinogen-nucleoside adduct with a molecular weight of 500 u, this corresponds to about 2 pmol. With this detection limit the analysis of in vivo samples from animals dosed with carcinogens should be possible based on one adduct per 10^6 nucleotides (manuscript in preparation).

Daughter ion scans were used to differentiate C-8 substituted arylamine adducts of deoxyadenosine and deoxyguanosine by using the BH_2^+ ion as the parent. For dG-C8-ABP this fragment ion showed the same daughter ion spectrum as the protonated molecule of

the carcinogen-base, G-C8-ABP. Fragmentation of the BH_2^+ parent ions was largely in agreement with previously reported results obtained by using high energy collisions.

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